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DIABETE

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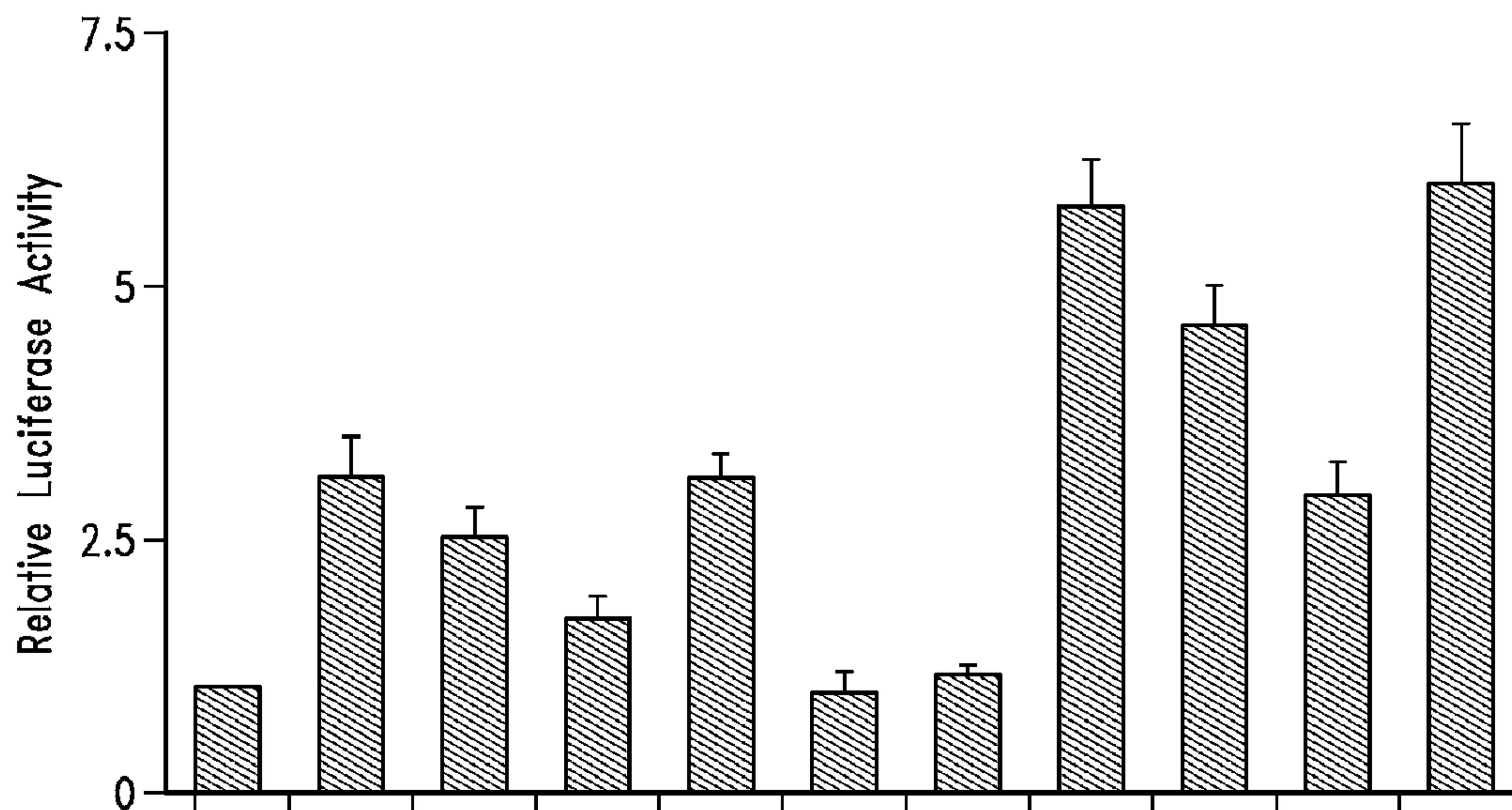


Fig. 124C

(57) Abrégé/Abstract:

Provided are electrokinetically-altered fluids (gas-enriched electrokinetic fluids) comprising an ionic aqueous solution of charge-stabilized oxygen-containing nanostructures in an amount sufficient to provide modulation of at least one of cellular membrane potential and cellular membrane conductivity, and therapeutic compositions and methods for use in treating diabetes and diabetes-associated conditions or disorders (e.g., insulin resistance), or symptoms thereof. Provided are electrokinetically-altered ionic aqueous fluids optionally in combination with other therapeutic agents. Particular aspects provide for regulating or modulating intracellular signal transduction associated with said inflammatory responses by modulation of at least one of cellular membranes, membrane potential, membrane proteins such as membrane receptors, including but not limited to G-Protein Coupled Receptors (GPCR), and intercellular junctions (e.g., tight junctions, gap junctions, zona adherens and desmosomes). Other embodiments include particular routes of administration or formulations for the electrokinetically-altered fluids (e.g., electrokinetically-altered gas-enriched fluids and solutions) and therapeutic compositions.

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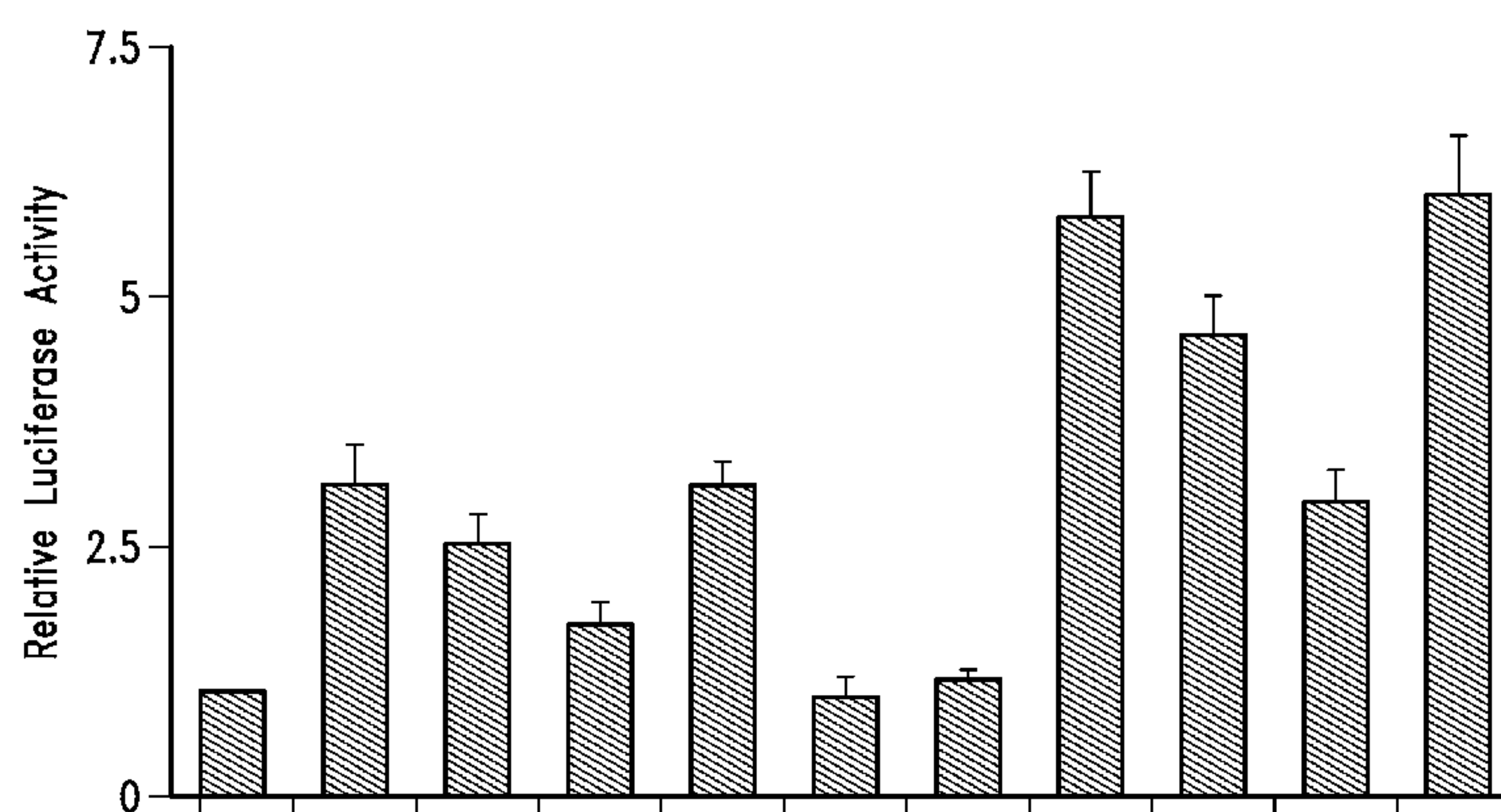


Fig. 124C

(57) **Abstract:** Provided are electrokinetically-altered fluids (gas-enriched electrokinetic fluids) comprising an ionic aqueous solution of charge-stabilized oxygen-containing nanostructures in an amount sufficient to provide modulation of at least one of cellular membrane potential and cellular membrane conductivity, and therapeutic compositions and methods for use in treating diabetes and diabetes-associated conditions or disorders (e.g., insulin resistance), or symptoms thereof. Provided are electrokinetically-altered ionic aqueous fluids optionally in combination with other therapeutic agents. Particular aspects provide for regulating or modulating intracellular signal transduction associated with said inflammatory responses by modulation of at least one of cellular membranes, membrane potential, membrane proteins such as membrane receptors, including but not limited to G-Protein Coupled Receptors (GPCR), and intercellular junctions (e.g., tight junctions, gap junctions, zona adherens and desmosomes). Other embodiments include particular routes of administration or formulations for the electrokinetically-altered fluids (e.g., electrokinetically-altered gas-enriched fluids and solutions) and therapeutic compositions.



WO 2010/126908 A1

COMPOSITIONS AND METHODS FOR TREATING INSULIN RESISTANCE AND DIABETES MELLITUS

5 FIELD OF THE INVENTION

Certain embodiments disclosed herein relate to treating insulin resistance and/or a diabetic conditions or disorders, or at least one symptom thereof, in a subject by administering a therapeutic composition comprising at least one electrokinetically-altered fluid as disclosed herein. Particular embodiments disclosed herein relate to
10 regulating or modulating intracellular signal transduction by modulation of at least one of cellular membranes, membrane potential, membrane proteins such as membrane receptors, including but not limited to G-Protein Coupled Receptors, and intercellular junctions (e.g., tight junctions, gap junctions, zona adherins and desmosomes). Certain aspects relate to electrokinetically-altered fluids (gas-enriched electrokinetic fluids)
15 comprising an ionic aqueous solution of charge-stabilized oxygen-containing nanostructures in an amount sufficient to provide modulation of at least one of cellular membrane potential and cellular membrane conductivity.

CROSS-REFERENCE TO RELATED APPLICATIONS

20 This application claims the benefit of priority to United States Provisional Patent Application Serial No. 61/173,134 filed 27 April 2009.

BACKGROUND OF THE INVENTION

Diabetes mellitus is a serious lifelong metabolic disease that is defined by the presence of chronically elevated levels of blood glucose (hyperglycemia). This disease
25 often leads to blindness, heart and blood vessel disease, strokes, kidney failure, amputations, and nerve damage. Uncontrolled diabetes can complicate pregnancy, and birth defects are more common in babies born to women with diabetes. Diabetes is widely recognized as one of the leading causes of death and disability in the United States. Diabetes can lead to serious and premature complications like diabetic
30 retinopathy, for example.

Recent scientific discoveries have suggested a link between chronic inflammation and insulin resistance and/or diabetes. Inflammation is the immune and/or vascular response to trauma or infection by microbes, such as bacterial or

viruses, and it may be an acute or chronic and/or localized or systemic. Inflammatory reactions typically destroy, dilute, or confine the injurious agent and the injured tissue in the subject. Inflammation is characterized, particularly in the acute form, by the classic signs of pain, heat, redness, swelling, and possibly loss of function. At a histological level, inflammation involves a complex series of events, including dilation of arterioles, capillaries, and venules, and an increased permeability and blood flow, exudation of fluids, including plasma proteins, and leukocyte migration into the area of inflammation, particularly with a localized reaction.

Therapeutic treatments for diabetic disorders include a wide array of pharmaceutical drugs. However, most of the treatments available today have considerable side effects, such as abdominal pain, diarrhea, nausea, gas, bloating, loss of appetite, weight gain, hypoglycemia and fluid retention. Thus, there is a need for better diabetic therapeutics and treatment methods. With the link between inflammation and insulin resistance and diabetes, a combinational treatment whereby both anti-inflammatory and diabetes drugs were used in concert would likely result in a superior treatment.

SUMMARY OF THE INVENTION

Particular aspects provide a method for treating diabetes or a diabetes-associated condition or disorder, or symptoms thereof, comprising administering to a subject in need thereof a therapeutically effective amount of an electrokinetically altered aqueous fluid comprising an ionic aqueous solution of charge-stabilized oxygen-containing nanostructures substantially having an average diameter of less than about 100 nanometers and stably configured in the ionic aqueous fluid in an amount sufficient for treating diabetes or a diabetes-associated condition or disorder, or at least one symptom thereof. In certain embodiments, the charge-stabilized oxygen-containing nanostructures are the major charge-stabilized gas-containing nanostructure species in the fluid. In particular aspects, the percentage of dissolved oxygen molecules present in the fluid as the charge-stabilized oxygen-containing nanostructures is a percentage selected from the group consisting of greater than: 0.01%, 0.1%, 1%, 5%; 10%; 15%; 20%; 25%; 30%; 35%; 40%; 45%; 50%; 55%; 60%; 65%; 70%; 75%; 80%; 85%; 90%; and 95%. In certain embodiments, the total dissolved oxygen is substantially present in

the charge-stabilized oxygen-containing nanostructures. In particular aspects, the charge-stabilized oxygen-containing nanostructures substantially have an average diameter of less than a size selected from the group consisting of: 90 nm; 80 nm; 70 nm; 60 nm; 50 nm; 40 nm; 30 nm; 20 nm; 10 nm; and less than 5 nm. In preferred
5 aspects, the ionic aqueous solution comprises a saline solution. In certain embodiments, the fluid is superoxygenated. In particular embodiments, the fluid comprises a form of solvated electrons.

In particular aspects, alteration of the electrokinetically altered aqueous fluid comprises exposure of the fluid to hydrodynamically-induced, localized electrokinetic
10 effects. In particular embodiments, exposure to the localized electrokinetic effects comprises exposure to at least one of voltage pulses and current pulses. In certain aspects, the exposure of the fluid to hydrodynamically-induced, localized electrokinetic effects, comprises exposure of the fluid to electrokinetic effect-inducing structural features of a device used to generate the fluid.

In certain embodiments, the diabetes-associated condition or disorder comprises at least one selected from the group consisting of: diabetes; insulin-dependent diabetes mellitus or IDDM (Type 1); non-insulin dependent diabetes mellitus or NIDDM (Type 2); insulin resistance; and diabetic retinopathy. In preferred aspects, the diabetes-associated condition or disorder comprises at least one of diabetes and insulin
15 resistance. Preferably, the diabetes-associated condition or disorder comprises insulin resistance. In certain aspects, the at least one symptom of the diabetes-associated condition or disorder is related to at least one condition selected from the group consisting of: chronic inflammation, acute inflammation, insulin resistance.

In certain aspects, the electrokinetically altered aqueous fluid modulates
25 localized or cellular levels of nitric oxide.

In particular embodiments, the electrokinetically altered aqueous fluid promotes a localized decrease at the site of administration of at least one cytokine selected from the group consisting of: IL-1beta, IL-8, TNF-alpha, and TNF-beta.

In certain aspects, the methods further comprise a synergistic or non-synergistic
30 inhibition or reduction in inflammation by simultaneously or adjunctively treating the subject with another anti-inflammatory agent. Preferably, said other anti-inflammatory

agent comprises a steroid or glucocorticoid steroid. In particular embodiments, the glucocorticoid steroid comprises Budesonide or an active derivative thereof.

In certain embodiments, the methods further comprise combination therapy, wherein at least one additional therapeutic agent is administered to the patient. In particular aspects, the at least one additional therapeutic agent is selected from the group consisting of: Biguanides including metformin, buformin, and phenformin, insulin, alpha-glucosidase inhibitors, biguanides, DPP-4 inhibitors, meglitinides, sulfonylureas, thiazolidinediones, alpha-glucosidase inhibitors including, acarbose and miglitol, DPP-4 inhibitors including vildagliptin, sitagliptin, saxagliptin, linagliptin, and alogliptin, sulfonylureas including acetohexamide, chlorpropamide, tolbutamide, tolazamide, glipizide, gliclazide, glibenclamide (glyburide), gliquidone, glycopyramide, and glimepiride, meglitinides including nateglinide, mitiglinide, and repaglinide. thiazolidinediones including troglitazone, pioglitazone, and rosiglitazone, inhibitors of MMPs including inhibitor of MMP-9 and MMP-2, short-acting β_2 -agonists, long-acting β_2 -agonists, anticholinergics, corticosteroids, systemic corticosteroids, mast cell stabilizers, leukotriene modifiers, methylxanthines, β_2 -agonists, albuterol, levalbuterol, pirbuterol, arformoterol, formoterol, salmeterol, anticholinergics including ipratropium and tiotropium; corticosteroids including beclomethasone, budesonide, flunisolide, fluticasone, mometasone, triamcinolone, methyprednisolone, prednisolone, prednisone; leukotriene modifiers including montelukast, zafirlukast, and zileuton; mast cell stabilizers including cromolyn and nedocromil; methylxanthines including theophylline; combination drugs including ipratropium and albuterol, fluticasone and salmeterol, budesonide and formoterol; antihistamines including hydroxyzine, diphenhydramine, loratadine, cetirizine, and hydrocortisone; immune system modulating drugs including tacrolimus and pimecrolimus; cyclosporine; azathioprine; mycophenolatemofetil; and combinations thereof.

In certain aspects, the at least one additional therapeutic agent is a TSLP and/or TSLPR antagonist. Preferably, the TSLP and/or TSLPR antagonist is selected from the group consisting of neutralizing antibodies specific for TSLP and the TSLP receptor, soluble TSLP receptor molecules, and TSLP receptor fusion proteins, including TSLPR-immunoglobulin Fc molecules or polypeptides that encode components of more than

one receptor chain. In certain aspects, treating comprises altering at least one of cellular membrane structure or function comprising altering at least one of a conformation, ligand binding activity, or a catalytic activity of a membrane associated protein. In particular embodiments, the membrane associated protein comprises at least one selected from the group consisting of receptors, transmembrane receptors, ion channel proteins, intracellular attachment proteins, cellular adhesion proteins, and integrins. In certain aspects, the transmembrane receptor comprises a G-Protein Coupled Receptor (GPCR). In particular aspects, the G-Protein Coupled Receptor (GPCR) interacts with a G protein α subunit. Preferably, the G protein α subunit comprises at least one selected from the group consisting of G_{α_s} , G_{α_i} , G_{α_q} , and $G_{\alpha_{12}}$. Preferably, the at least one G protein α subunit is G_{α_q} .

In particular aspects, the charge-stabilized oxygen-containing nanostructures are stably configured in the ionic aqueous fluid in an amount sufficient to provide, upon contact of a living cell by the fluid, modulation of at least one of cellular membrane potential and cellular membrane conductivity. In certain aspects, modulating cellular membrane conductivity, comprises modulating whole-cell conductance. In particular aspects, modulating whole-cell conductance, comprises modulating at least one voltage-dependent contribution of the whole-cell conductance. Particular aspects comprise modulation of a calcium dependant cellular messaging pathway or system (e.g., modulation of phospholipase C activity; modulation of adenylate cyclase (AC) activity). Particular aspects comprise modulation of intracellular signal transduction associated with at least one condition or symptom selected from the group consisting of: chronic inflammation, acute inflammation, and insulin resistance.

Particular aspects of the methods comprise administration to a cell network or layer, and further comprise modulation of an intercellular junction therein (e.g., at least one selected from the group consisting of tight junctions, gap junctions, zona adherens and desmosomes). In certain aspects, the cell network or layers comprises at least one selected from the group consisting of endothelial cells, endothelial-astrocyte tight junctions in CNS vessels, blood-cerebrospinal fluid tight junctions or barrier, pulmonary epithelium-type junctions, bronchial epithelium-type junctions, and intestinal epithelium-type junctions.

In particular embodiments of the methods, the electrokinetically altered aqueous fluid is oxygenated, and wherein the oxygen in the fluid is present in an amount of at least 8 ppm, at least 15, ppm, at least 25 ppm, at least 30 ppm, at least 40 ppm, at least 50 ppm, or at least 60 ppm oxygen at atmospheric pressure. In certain aspects, the electrokinetically altered aqueous fluid comprises at least one of a form of solvated electrons, and electrokinetically modified or charged oxygen species. In certain aspects, the solvated electrons or electrokinetically modified or charged oxygen species are present in an amount of at least 0.01 ppm, at least 0.1 ppm, at least 0.5 ppm, at least 1 ppm, at least 3 ppm, at least 5 ppm, at least 7 ppm, at least 10 ppm, at least 15 ppm, or at least 20 ppm. In particular embodiments of the methods, electrokinetically altered oxygenated aqueous fluid comprises solvated electrons stabilized, at least in part, by molecular oxygen.

Certain aspects comprise the ability to alter cellular membrane structure or function sufficient to provide for modulation of intracellular signal transduction persisting for at least two, at least three, at least four, at least five, at least 6, at least 12 months, or longer periods, in a closed gas-tight container.

In particular embodiments of the methods, the amount of oxygen present in charge-stabilized oxygen-containing nanostructures of the electrokinetically-altered fluid is at least 8 ppm, at least 15, ppm, at least 20 ppm, at least 25 ppm, at least 30 ppm, at least 40 ppm, at least 50 ppm, or at least 60 ppm oxygen at atmospheric pressure.

Additional aspects provide a method of formulating a therapeutic agent suitable for use in treating diabetes, or a diabetes-associated condition or disorder, or symptoms thereof, comprising: obtaining a therapeutic agent suitable for use in treating diabetes, or a diabetes-associated condition or disorder, or symptoms thereof, of a subject; and combining the therapeutic agent with an amount of an electrokinetically altered aqueous fluid comprising an ionic aqueous solution of charge-stabilized oxygen-containing nanostructures substantially having an average diameter of less than about 100 nanometers and stably configured in the ionic aqueous fluid in an amount sufficient for treating inflammation or at least one symptom thereof, wherein formulating a therapeutic agent suitable for use in treating diabetes, or a diabetes-associated condition or disorder, or symptoms thereof is afforded. In particular embodiments, the

charge-stabilized oxygen-containing nanostructures are stably configured in the ionic aqueous fluid in an amount sufficient to provide, upon contact of a living cell by the fluid, modulation of at least one of cellular membrane potential and cellular membrane conductivity.

5 Yet further aspects provide a pharmaceutical composition, comprising: a therapeutic agent suitable for use treating diabetes, or a diabetes-associated condition or disorder, or symptoms thereof, of a subject; and an amount of an electrokinetically altered aqueous fluid comprising an ionic aqueous solution of charge-stabilized oxygen-containing nanostructures substantially having an average diameter of less than about
10 100 nanometers and stably configured in the ionic aqueous fluid in an amount sufficient for treating inflammation or at least one symptom thereof. Particular aspects provide a pharmaceutical composition, prepared by the method of claim 48. Certain aspects comprise administration by at least one of topical, inhalation, intranasal, and intravenous routes.

15 In particular aspects, the membrane associated protein comprises CCR3. In certain aspects, treating comprises modulation of NF- κ B expression and/or activity.

BRIEF DESCRIPTION OF THE DRAWINGS

Figure 1 is a partial cross-section, partial block diagram of a prior art mixing
20 device.

Figure 2 is block diagram of an exemplary embodiment of a mixing device.

Figure 3 is an illustration of an exemplary system for delivering a first material to the mixing device of Figure 2.

Figure 4 is a fragmentary partial cross-sectional view of a top portion of the
25 mixing device of Figure 2.

Figure 5 is a fragmentary cross-sectional view of a first side portion of the mixing device of Figure 2.

Figure 6 is a fragmentary cross-sectional view of a second side portion of the mixing device of Figure 2.

30 Figure 7 is a fragmentary cross-sectional view of a side portion of the mixing device of Figure 2 located between the first side portion of Figure 5 and the second side portion of Figure 6.

Figure 8 is a perspective view of a rotor and a stator of the mixing device of Figure 2.

Figure 9 is a perspective view of an inside of a first chamber of the mixing device of Figure 2.

5 Figure 10 is a fragmentary cross-sectional view of the inside of a first chamber of the mixing device of Figure 2 including an alternate embodiment of the pump 410.

Figure 11 is a perspective view of an inside of a second chamber of the mixing device of Figure 2.

10 Figure 12 is a fragmentary cross-sectional view of a side portion of an alternate embodiment of the mixing device.

Figure 13 is a perspective view of an alternate embodiment of a central section of the housing for use with an alternate embodiment of the mixing device.

Figure 14 is a fragmentary cross-sectional view of an alternate embodiment of a bearing housing for use with an alternate embodiment of the mixing device.

15 Figure 15 is a cross-sectional view of the mixing chamber of the mixing device of Figure 2 taken through a plane orthogonal to the axis of rotation depicting a rotary flow pattern caused by cavitation bubbles when a through-hole of the rotor approaches (but is not aligned with) an aperture of the stator.

20 Figure 16 is a cross-sectional view of the mixing chamber of the mixing device of Figure 2 taken through a plane orthogonal to the axis of rotation depicting a rotary flow pattern caused by cavitation bubbles when the through-hole of the rotor is aligned with the aperture of the stator.

25 Figure 17 is a cross-sectional view of the mixing chamber of the mixing device of Figure 2 taken through a plane orthogonal to the axis of rotation depicting a rotary flow pattern caused by cavitation bubbles when a through-hole of the rotor that was previously aligned with the aperture of the stator is no longer aligned therewith.

Figure 18 is a side view of an alternate embodiment of a rotor.

30 Figure 19 is an enlarged fragmentary cross-sectional view taken through a plane orthogonal to an axis of rotation of the rotor depicting an alternate configuration of through-holes formed in the rotor and through-holes formed in the stator.

Figure 20 is an enlarged fragmentary cross-sectional view taken through a plane passing through and extending along the axis of rotation of the rotor depicting a

configuration of through-holes formed in the rotor and through-holes formed in the stator.

Figure 21 is an enlarged fragmentary cross-sectional view taken through a plane passing through and extending along the axis of rotation of the rotor depicting an alternate offset configuration of through-holes formed in the rotor and through-holes formed in the stator.

Figure 22 is an illustration of a shape that may be used to construct the through-holes of the rotor and/or the apertures of the stator.

Figure 23 is an illustration of a shape that may be used to construct the through-holes of the rotor and/or the apertures of the stator.

Figure 24 is an illustration of a shape that may be used to construct the through-holes of the rotor and/or the apertures of the stator.

Figure 25 is an illustration of a shape that may be used to construct the through-holes of the rotor and/or the apertures of the stator.

Figure 26 is an illustration of an electrical double layer ("EDL") formed near a surface.

Figure 27 is a perspective view of a model of the inside of the mixing chamber.

Figure 28 is a cross-sectional view of the model of Figure 27.

Figure 29 is an illustration of an experimental setup.

Figure 30 illustrates dissolved oxygen levels in water processed with oxygen in the mixing device of Figure 2 and stored in a 500 ml thin walled plastic bottle and a 1,000 ml glass bottle each capped at 65° Fahrenheit.

Figure 31 illustrates dissolved oxygen levels in water processed with oxygen in the mixing device of Figure 2 and stored in a 500 ml plastic thin walled bottle and a 1,000 ml glass bottle both refrigerated at 39° Fahrenheit.

Figure 32 illustrates the dissolved oxygen retention of a 500 ml beverage fluid processed with oxygen in the mixing device of Figure 2.

Figure 33 illustrates the dissolved oxygen retention of a 500 ml braun balanced salt solution processed with oxygen in the mixing device of Figure 2.

Figure 34 illustrates a further experiment wherein the mixing device of Figure 2 is used to sparge oxygen from water by processing the water with nitrogen in the mixing device of Figure 2.

Figure 35 illustrates the sparging of oxygen from water by the mixing device of Figure 2 at standard temperature and pressure.

Figure 36 is an illustration of an exemplary nanocage.

Figure 37A and B illustrate Rayleigh scattering effects of an oxygen-enriched
5 fluid;

Figure 38 illustrates the cytokine profile of a mitogenic assay in the presence of a gas-enriched fluid and deionized control fluid; and

Figure 39 illustrates the difference in the growth rates of *Pseudomonas* bacteria at various dissolved oxygen saturation ratios.

10 Figures 40A and 40B illustrate *in vitro* healing of wounds using an oxygen-enriched cell culture media and a non-gas-enriched media.

Figures 41A through 41F show histological cross-sections of dermal and epidermal *in vivo* wound healing.

Figure 42 illustrates the expression of Hale's stain in treated and control healing
15 wounds, used to detect acid mucopolysaccharides, such as hyaluronic acid.

Figure 43 illustrates the expression of von Willebrand's Factor stain used to detect angiogenesis in treated and control healing wounds.

Figure 44 illustrates the detection of Luna's stain used to detect elastin in treated and control healing wounds.

20 Figure 45 illustrates the number of mast cells per visual field for treated and control healing wounds.

Figure 46 illustrates the percentage of dead cells at separate time points in a corneal fibroblast assay using inventive gas-enriched culture media and control culture media.

25 Figure 47 illustrates the shelf life of the inventive gas-enriched fluid in a polymer pouch.

Figure 48 illustrates the results of contacting splenocytes with MOG in the presence of pressurized pot oxygenated fluid (1), inventive gas-enriched fluid (2), or control deionized fluid (3).

30 Figures 49-58 show the results of whole blood sample evaluations of cytokines.

Figures 59-68 show the corresponding cytokine results of bronchoalveolar lavage fluid (BAL) sample evaluations.

Figures 69-75 show studies where the Bradykinin B2 membrane receptor was immobilized onto aminopropylsilane (APS) biosensor. The Sample plate setup was as designated in Figure 69 and the binding of Bradykinin to the immobilized receptor was assessed according to the sample setup as designated in Figure 71. Results of
5 Bradykinin binding are shown in Figure 72. Bradykinin binding to the receptor was further titrated according to the setup as designated in Figure 73. As indicated in Figure 74, Bradykinin binding to the B2 receptor was concentration dependent, and binding affinity was increased in the proprietary gas-enriched saline fluid of the instant disclosure compared to normal saline. Stabilization of Bradykinin binding to the B2
10 receptor is shown in Figure 75.

Figures 76-83 show data showing the ability of particular embodiments disclosed herein to affect regulatory T cells. The study involved irradiating antigen presenting cells, and introducing antigen and T cells.

Figure 84 shows that the inventive electrokinetically-generated fluids decreased
15 serum uptake of salmon calcitonin and an animal model. The results are consistent with enhancement of tight junctions.

Figures 85-89 show the expression levels of tight junction-related proteins in lung tissue from the animal model used to generate the data of Figure 84.

Figures 90-94 show data obtained from human foreskin keratinocytes exposed
20 to RDC1676-01 (sterile saline processed through the instant proprietary device with additional oxygen added; gas-enriched electrokinetically-generated fluid (Rev) of the instant disclosure) showing up-regulation of NOS1 and 3, and Nostrin, NOS3.

Figures 95 and 96 show data supporting localized electrokinetic effects (voltage/current) occurring in a mixing device comprising insulated rotor and stator
25 features to allow for detection of voltage/current effects during electrokinetic fluid generation.

Figures 97A-C show results of nuclear magnetic resonance (NMR) studies conducted to further characterize the fundamental nature of the inventive electrokinetically-generated fluids. The electrokinetically-generated fluids increased the
30 ^{13}C -NMR line-widths of the reporter Trehalose solute.

Figures 98 and 99 show results of voltametric studies (i.e., square wave voltametry (Fig. 98) and stripping polarography (Fig. 99)) conducted to further

characterize the fundamental nature of the inventive electrokinetically-generated fluids. Square wave voltametry peak differences (with respect to control) unique to the electrokinetically-generated fluids were observed at -0.14V, -0.47V, -1.02V and -1.36V. Pronounced polarographic peaks were seen at -0.9 volts for the electrokinetically-generated Revera and Solas fluids, and the spectra of the non-electrokinetically-generated blank and saline control fluids show characteristic peaks at -0.19 and -0.3 volts that are absent in the spectra for the electrokinetically-generated fluids.

Figures 100-106 show results of patch clamping techniques that assessed the effects of the electrokinetically-generated fluid test on epithelial cell membrane polarity and ion channel activity. The results indicate that the inventive electrokinetically-generated fluids affect a voltage-dependent contribution of the whole-cell conductance.

Figures 107A-D and 108A-D show data indicating that the inventive electrokinetically-generated fluids (e.g., RDC1676-00, RDC1676-01, RDC1676-02 and RDC1676-03) protected against methacholine-induced bronchoconstriction when administered alone or as diluents for albuterol sulfate in male guinea pigs.

Figures 109-114 show results of budesonide experiments performed to assess the airway anti-inflammatory properties of the inventive electrokinetically-generated fluids in a Brown Norway rat ovalbumin sensitization model. The inventive electrokinetically-generated fluids decreased eosinophil count, showed strong synergy with Budesonide in decreasing eosinophil count, decreased Penh values, increased Tidal Volume, decreased blood levels of Eotaxin, significantly enhanced the Blood levels of two major key anti-inflammatory cytokines, IL10 and Interferon gamma at 6 hours after challenge as a result of treatment with the inventive electrokinetically-generated fluid (e.g., Rev 60) alone or in combination with Budesonide, and decreased systemic levels of Rantes. The data show that there is a substantial synergistic effect of Budesonide 750 ug/kg and the inventive electrokinetically-generated fluids (e.g., Rev 60).

Figure 115 shows that the inventive electrokinetically-generated fluid (e.g., Revera 60 and Solas) reduced DEP-induced TSLP receptor expression in bronchial epithelial cells (BEC) by approximately 90% and 50%, respectively, whereas normal saline (NS) had only a marginal effect.

Figure 116 shows the inventive electrokinetically-generated fluid (e.g., Revera 60 and Solas) inhibited the DEP-induced cell surface-bound MMP9 levels in bronchial epithelial cells by approximately 80%, and 70%, respectively, whereas normal saline (NS) had only a marginal effect.

5 Figures 117 A-C demonstrate the results of a series of patch clamping experiments that assessed the effects of the electrokinetically generated fluid (e.g., RNS-60 and Solas) on epithelial cell membrane polarity and ion channel activity at two time-points (15 min (left panels) and 2 hours (right panels)) and at different voltage protocols.

10 Figures 118 A-C show, in relation to the experiments relating to Figures 117 A-C, the graphs resulting from the subtraction of the Solas current data from the RNS-60 current data at three voltage protocols (A. stepping from zero mV; B. stepping from -60 mV; C. stepping from -120 mV) and the two time-points (15 mins (open circles) and 2 hours (closed circles)).

15 Figures 119 A-D demonstrate the results of a series of patch clamping experiments that assessed the effects of the electrokinetically generated fluid (e.g., Solas (panels A. and B.) and RNS-60 (panels C. and D.)) on epithelial cell membrane polarity and ion channel activity using different external salt solutions and at different voltage protocols (panels A. and C. show stepping from zero mV; panels B. and D. show stepping from -120 mV).

20 Figures 120 A-D show, in relation to the experiments relating to Figures 119 A-D, the graphs resulting from the subtraction of the CsCl current data (shown in Figure 119) from the 20 mM CaCl_2 (diamonds) and 40 mM CaCl_2 (filled squares) current data at two voltage protocols (panels A. and C. stepping from zero mV; B. and D. stepping from -120 mV) for Solas (panels A. and B.) and Revera 60 (panels C. and D.).

Figure 121 A shows 1 mm² AFM scan for RNS60-1 (rns60-1 1um 3D.jpg). The small peaks ("1") represent hydrophobic nanobubbles which are ~20 nm wide and ~1.5 nm tall or smaller.

30 Figure 121 B shows 1 mm² scan for PNS60-1 (pp60-1 1um 3d.jpg). This scan reveals peaks ("2") (hydrophobic nanobubbles) that are substantially larger (~60 nm wide and ~5 nm tall) than those visible with RNS60-1.

Figures 122 A-B demonstrate the results of Fluorescence-Activated Cell Sorting (FACS) analysis wherein the levels of expression of the cell surface receptor, CD193 (CCR3), on white blood cells was compared using either normal saline or RNS-60. The X-axis represents the log fluorescence of the sample and the Y-axis represents the events of fluorescence that occur in the sample.

Figures 123 A-C demonstrate the results of Fluorescence-Activated Cell Sorting (FACS) analysis wherein the levels of expression of cell surface receptors, CD154 (CD40L) (panel A); CD11B (panel B); and CD3 (panel C), on white blood cells was compared using either normal saline or RNS-60. The X-axis represents the log fluorescence of the sample and the Y-axis represents the events of fluorescence that occur in the sample.

Figures 124 A-C show the results from two gel shift experiments (panels A and B) and a luciferase activity (reporter gene) assay (panel C) that examined the effects of RNS60 on the activation of NF κ B in MBP-primed T cells.

DETAILED DESCRIPTION OF THE INVENTION

Certain embodiments disclosed herein relate to providing compositions and methods of treatment of insulin resistance and/or a diabetic condition or disorder, or at least one symptom thereof, by administering to a subject, a therapeutic composition comprising a gas-enriched fluid. In certain specific embodiments, the gas-enriched fluid comprises oxygen-enriched water.

Diabetic disorders and conditions

Certain embodiments herein relate to therapeutic compositions and methods of treatment for a subject by preventing or alleviating at least one symptom of insulin resistance and/or a diabetic associated condition or disease.

In further embodiments herein relating to the therapeutic compositions and methods of treatment for preventing or alleviating complications related to insulin resistance and/or diabetic-associated condition, including alleviating the symptoms of diabetic retinopathy, for example.

Electrokinetically-generated fluids:

“Electrokinetically-generated fluid,” as used herein, refers to Applicants’ inventive electrokinetically-altered fluids generated, for purposes of the working Examples herein, by the exemplary Mixing Device described in detail herein (see also US200802190088 and WO2008/052143, both incorporated herein by reference in their entirety). The electrokinetic fluids, as demonstrated by the data disclosed and presented herein, represent novel and fundamentally distinct fluids relative to prior art non-electrokinetic fluids, including relative to prior art oxygenated non-electrokinetic fluids (e.g., pressure pot oxygenated fluids and the like). As disclosed in various aspects herein, the electrokinetically-generated fluids have unique and novel physical and biological properties including, but not limited to the following:

In particular aspects, the electrokinetically altered aqueous fluid comprise an ionic aqueous solution of charge-stabilized oxygen-containing nanostructures substantially having an average diameter of less than about 100 nanometers and stably configured in the ionic aqueous fluid in an amount sufficient to provide, upon contact of a living cell by the fluid, modulation of at least one of cellular membrane potential and cellular membrane conductivity.

In particular aspects, electrokinetically-generated fluids refers to fluids generated in the presence of hydrodynamically-induced, localized (e.g., non-uniform with respect to the overall fluid volume) electrokinetic effects (e.g., voltage/current pulses), such as device feature-localized effects as described herein. In particular aspects, said hydrodynamically -induced, localized electrokinetic effects are in combination with surface-related double layer and/or streaming current effects as disclosed and discussed herein.

In particular aspects the administered inventive electrokinetically-altered fluids comprise charge-stabilized oxygen-containing nanostructures in an amount sufficient to provide modulation of at least one of cellular membrane potential and cellular membrane conductivity. In certain embodiments, the electrokinetically-altered fluids are superoxygenated (e.g., RNS-20, RNS-40 and RNS-60, comprising 20 ppm, 40 ppm and 60 ppm dissolved oxygen, respectively, in standard saline). In particular embodiments, the electrokinetically-altered fluids are not-superoxygenated (e.g., RNS-10 or Solas, comprising 10 ppm (e.g., approx. ambient levels of dissolved oxygen in standard

saline). In certain aspects, the salinity, sterility, pH, etc., of the inventive electrokinetically-altered fluids is established at the time of electrokinetic production of the fluid, and the sterile fluids are administered by an appropriate route. Alternatively, at least one of the salinity, sterility, pH, etc., of the fluids is appropriately adjusted (e.g.,
5 using sterile saline or appropriate diluents) to be physiologically compatible with the route of administration prior to administration of the fluid. Preferably, and diluents and/or saline solutions and/or buffer compositions used to adjust at least one of the salinity, sterility, pH, etc., of the fluids are also electrokinetic fluids, or are otherwise compatible.

10 In particular aspects, the inventive electrokinetically-altered fluids comprise saline (e.g., one or more dissolved salt(s); e.g., alkali metal based salts (Li^+ , Na^+ , K^+ , Rb^+ , Cs^+ , etc.), alkaline earth based salts (e.g., Mg^{++} , Ca^{++}), etc., or transition metal-based positive ions (e.g., Cr, Fe, Co, Ni, Cu, Zn, etc.), in each case along with any suitable anion components, including, but not limited to F^- , Cl^- , Br^- , I^- , PO_4^- , SO_4^- , and
15 nitrogen-based anions. . Particular aspects comprise mixed salt based electrokinetic fluids (e.g., Na^+ , K^+ , Ca^{++} , Mg^{++} , transition metal ion(s), etc.) in various combinations and concentrations, and optionally with mixtures of counterions. In particular aspects, the inventive electrokinetically-altered fluids comprise standard saline (e.g., approx. 0.9% NaCl, or about 0.15 M NaCl). In particular aspects, the inventive
20 electrokinetically-altered fluids comprise saline at a concentration of at least 0.0002 M, at least 0.0003 M, at least 0.001 M, at least 0.005 M, at least 0.01 M, at least 0.015 M, at least 0.1 M, at least 0.15 M, or at least 0.2 M. In particular aspects, the conductivity of the inventive electrokinetically-altered fluids is at least 10 $\mu\text{S}/\text{cm}$, at least 40 $\mu\text{S}/\text{cm}$, at least 80 $\mu\text{S}/\text{cm}$, at least 100 $\mu\text{S}/\text{cm}$, at least 150 $\mu\text{S}/\text{cm}$, at least 200 $\mu\text{S}/\text{cm}$, at least
25 300 $\mu\text{S}/\text{cm}$, or at least 500 $\mu\text{S}/\text{cm}$, at least 1 mS/cm, at least 5 mS/cm, 10 mS/cm, at least 40 mS/cm, at least 80 mS/cm, at least 100 mS/cm, at least 150 mS/cm, at least 200 mS/cm, at least 300 mS/cm, or at least 500 mS/cm. In particular aspects, any salt may be used in preparing the inventive electrokinetically-altered fluids, provided that they allow for formation of biologically active salt-stabilized nanostructures (e.g., salt-stabilized oxygen-containing nanostructures) as disclosed herein.
30

According to particular aspects, the biological effects of the inventive fluid compositions comprising charge-stabilized gas-containing nanostructures can be

modulated (e.g., increased, decreased, tuned, etc.) by altering the ionic components of the fluids, and/or by altering the gas component of the fluid.

According to particular aspects, the biological effects of the inventive fluid compositions comprising charge-stabilized gas-containing nanostructures can be modulated (e.g., increased, decreased, tuned, etc.) by altering the gas component of the fluid. In preferred aspects, oxygen is used in preparing the inventive electrokinetic fluids. In additional aspects mixtures of oxygen along with at least one other gas selected from Nitrogen, Oxygen, Argon, Carbon dioxide, Neon, Helium, krypton, hydrogen and Xenon. As described above, the ions may also be varied, including along with varying the gas constituent(s).

Given the teachings and assay systems disclosed herein (e.g., cell-based cytokine assays, patch-clamp assays, etc.) one of skill in the art will readily be able to select appropriate salts and concentrations thereof to achieve the biological activities disclosed herein.

TABLE 1. Exemplary cations and anions.

Common Cations:

<u>Name</u>	<u>Formula</u>	<u>Other name(s)</u>
Aluminum	Al^{+3}	
Ammonium	NH_4^+	
Barium	Ba^{+2}	
Calcium	Ca^{+2}	
Chromium(II)	Cr^{+2}	Chromous
Chromium(III)	Cr^{+3}	Chromic
Copper(I)	Cu^+	Cuprous
Copper(II)	Cu^{+2}	Cupric
Iron(II)	Fe^{+2}	Ferrous
Iron(III)	Fe^{+3}	Ferric
Hydrogen	H^+	
Hydronium	H_3O^+	
Lead(II)	Pb^{+2}	
Lithium	Li^+	
Magnesium	Mg^{+2}	
Manganese(II)	Mn^{+2}	Manganous
Manganese(III)	Mn^{+3}	Manganic

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Mercury(I)	Hg_2^{+2}	Mercurous
Mercury(II)	Hg^{+2}	Mercuric
Nitronium	NO_2^+	
Potassium	K^+	
Silver	Ag^+	
Sodium	Na^+	
Strontium	Sr^{+2}	
Tin(II)	Sn^{+2}	Stannous
Tin(IV)	Sn^{+4}	Stannic
Zinc	Zn^{+2}	

Common Anions:

Simple ions:

Hydride	H^-	Oxide	O^{2-}
Fluoride	F^-	Sulfide	S^{2-}
Chloride	Cl^-	Nitride	N^{3-}
Bromide	Br^-		
Iodide	I^-		

Oxoanions:

Arsenate	AsO_4^{3-}	Phosphate	PO_4^{3-}
Arsenite	AsO_3^{3-}	Hydrogen phosphate	HPO_4^{2-}
		Dihydrogen phosphate	H_2PO_4^-
Sulfate	SO_4^{2-}	Nitrate	NO_3^-
Hydrogen sulfate	HSO_4^-	Nitrite	NO_2^-
Thiosulfate	$\text{S}_2\text{O}_3^{2-}$		
Sulfite	SO_3^{2-}		
Perchlorate	ClO_4^-	Iodate	IO_3^-
Chlorate	ClO_3^-	Bromate	BrO_3^-
Chlorite	ClO_2^-		
Hypochlorite	OCl^-	Hypobromite	OBr^-
Carbonate	CO_3^{2-}	Chromate	CrO_4^{2-}
Hydrogen carbonate or Bicarbonate	HCO_3^-	Dichromate	$\text{Cr}_2\text{O}_7^{2-}$

Multivalent Cations

Formula	Charge	Name
Cr^{2+}	2	chromium(II) or chromous ion
Cr^{3+}	3	chromium(III) or chromic ion
Mn^{2+}	2	manganese(II) or manganous ion
Mn^{4+}	4	manganese(IV) ion
Fe^{2+}	2	iron(II) or ferrous ion
Fe^{3+}	3	iron(III) or ferric ion
Co^{2+}	2	cobalt(II) or cobaltous ion
Co^{3+}	3	cobalt(III) or cobaltic ion
Ni^{2+}	2	nickel(II) or nickelous ion
Ni^{3+}	3	nickel(III) or nickelic ion
Cu^{+}	1	copper(I) or cuprous ion
Cu^{2+}	2	copper(II) or cupric ion
Sn^{2+}	2	tin(II) or stannous ion
Sn^{4+}	4	tin(IV) or stannic ion
Pb^{2+}	2	lead(II) or plumbous ion
Pb^{4+}	4	lead(IV) or plumbic ion

Monoatomic Anions

Formula	Charge	Name
H^{-}	1-	hydride ion
F^{-}	1-	fluoride ion
Cl^{-}	1-	chloride ion
Br^{-}	1-	bromide ion
I^{-}	1-	iodide ion
O^{2-}	2-	oxide ion
S^{2-}	2-	sulfide ion
N^{3-}	3-	nitride ion

Polyatomic Anions

Formula	Charge	Name
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OH^-	1-	hydroxide ion
CN^-	1-	cyanide ion
SCN^-	1-	thiocyanate ion
$\text{C}_2\text{H}_3\text{O}_2^-$	1-	acetate ion
ClO^-	1-	hypochlorite ion
ClO_2^-	1-	chlorite ion
ClO_3^-	1-	chlorate ion
ClO_4^-	1-	perchlorate ion
NO_2^-	1-	nitrite ion
NO_3^-	1-	nitrate ion
MnO_4^{2-}	2-	permanganate ion
CO_3^{2-}	2-	carbonate ion
$\text{C}_2\text{O}_4^{2-}$	2-	oxalate ion
CrO_4^{2-}	2-	chromate ion
$\text{Cr}_2\text{O}_7^{2-}$	2-	dichromate ion
SO_3^{2-}	2-	sulfite ion
SO_4^{2-}	2-	sulfate ion
PO_3^{3-}	3-	phosphite ion
PO_4^{3-}	3-	phosphate ion

The present disclosure sets forth novel gas-enriched fluids, including, but not limited to gas-enriched ionic aqueous solutions, aqueous saline solutions (e.g., standard aqueous saline solutions, and other saline solutions as discussed herein and as would be recognized in the art, including any physiological compatible saline solutions), cell culture media (e.g., minimal medium, and other culture media) useful in the treatment of diabetes or diabetes related disorders. A medium, or media, is termed "minimal" if it only contains the nutrients essential for growth. For prokaryotic host cells, a minimal media typically includes a source of carbon, nitrogen, phosphorus, magnesium, and trace amounts of iron and calcium. (Gunsalus and Stanter, The Bacteria, V. 1, Ch. 1 Acad. Press Inc., N.Y. (1960)). Most minimal media use glucose as a carbon source, ammonia as a nitrogen source, and orthophosphate (e.g., PO_4) as the phosphorus source. The media components can be varied or supplemented according to the specific prokaryotic or eukaryotic organism(s) grown, in order to

encourage optimal growth without inhibiting target protein production. (Thompson et al., *Biotech. and Bioeng.* 27: 818-824 (1985)).

In particular aspects, the electrokinetically-altered aqueous fluids are suitable to modulate ^{13}C -NMR line-widths of reporter solutes (e.g., Trehelose) dissolved therein.

5 NMR line-width effects are an indirect method of measuring, for example, solute 'tumbling' in a test fluid as described herein in particular working Examples.

In particular aspects, the electrokinetically-altered aqueous fluids are characterized by at least one of: distinctive square wave voltametry peak differences at any one of -0.14V, -0.47V, -1.02V and -1.36V; polarographic peaks at -0.9 volts; and an
10 absence of polarographic peaks at -0.19 and -0.3 volts, which are unique to the electrokinetically-generated fluids as disclosed herein in particular working Examples.

In particular aspects, the electrokinetically-altered aqueous fluids are suitable to alter cellular membrane conductivity (e.g., a voltage-dependent contribution of the whole-cell conductance as a measure in patch clamp studies disclosed herein).

15 In particular aspects, the electrokinetically-altered aqueous fluids are oxygenated, wherein the oxygen in the fluid is present in an amount of at least 15, ppm, at least 25 ppm, at least 30 ppm, at least 40 ppm, at least 50 ppm, or at least 60 ppm dissolved oxygen at atmospheric pressure. In particular aspects, the electrokinetically-altered aqueous fluids have less than 15 ppm, less than 10 ppm of dissolved oxygen at
20 atmospheric pressure, or approximately ambient oxygen levels.

In particular aspects, the electrokinetically-altered aqueous fluids are oxygenated, wherein the oxygen in the fluid is present in an amount between approximately 8 ppm and approximately 15 ppm, and in this case is sometimes referred to herein as "Solas."

25 In particular aspects, the electrokinetically-altered aqueous fluid comprises at least one of solvated electrons (e.g., stabilized by molecular oxygen), and electrokinetically-modified and/or charged oxygen species, and wherein in certain embodiments the solvated electrons and/or electrokinetically-modified or charged oxygen species are present in an amount of at least 0.01 ppm, at least 0.1 ppm, at least
30 0.5 ppm, at least 1 ppm, at least 3 ppm, at least 5 ppm, at least 7 ppm, at least 10 ppm, at least 15 ppm, or at least 20 ppm.

In particular aspects, the electrokinetically-altered aqueous fluids are suitable to alter cellular membrane structure or function (e.g., altering of a conformation, ligand binding activity, or a catalytic activity of a membrane-associated protein) sufficient to provide for modulation of intracellular signal transduction, wherein in particular aspects, 5 the membrane-associated protein comprises at least one selected from the group consisting of receptors, transmembrane receptors (e.g., G-Protein Coupled Receptor (GPCR), TSLP receptor, beta 2 adrenergic receptor, bradykinin receptor, etc.), ion channel proteins, intracellular attachment proteins, cellular adhesion proteins, and integrins. In certain aspects, the effected G-Protein Coupled Receptor (GPCR) 10 interacts with a G protein a subunit (e.g., G_{α_s} , G_{α_i} , G_{α_q} , and $G_{\alpha_{12}}$).

In particular aspects, the electrokinetically-altered aqueous fluids are suitable to modulate intracellular signal transduction, comprising modulation of a calcium dependant cellular messaging pathway or system (e.g., modulation of phospholipase C activity, or modulation of adenylate cyclase (AC) activity).

15 In particular aspects, the electrokinetically-altered aqueous fluids are characterized by various biological activities (e.g., regulation of cytokines, receptors, enzymes and other proteins, and intracellular signaling pathways) described in the working Examples and elsewhere herein.

In particular aspects, the electrokinetically-altered aqueous fluids display synergy 20 and/or additive activity with metformin.

In particular aspects, the electrokinetically-altered aqueous fluids reduce DEP-induced TSLP receptor expression in bronchial epithelial cells (BEC) as shown in working Examples herein.

In particular aspects, the electrokinetically-altered aqueous fluids inhibit the 25 DEP-induced cell surface-bound MMP9 levels in bronchial epithelial cells (BEC) as shown in working Examples herein.

In particular aspects, the biological effects of the electrokinetically-altered aqueous fluids are inhibited by diphtheria toxin, indicating that beta blockade, GPCR blockade and Ca channel blockade affects the activity of the electrokinetically-altered 30 aqueous fluids (e.g., on regulatory T cell function) as shown in working Examples herein.

In particular aspects, the physical and biological effects (e.g., the ability to alter cellular membrane structure or function sufficient to provide for modulation of intracellular signal transduction) of the electrokinetically-altered aqueous fluids persists for at least two, at least three, at least four, at least five, at least six months, or longer periods, in a closed container (e.g., closed gas-tight container).

Therefore, further aspects provide said electrokinetically-generated solutions and methods of producing an electrokinetically-altered oxygenated aqueous fluid or solution, comprising: providing a flow of a fluid material between two spaced surfaces in relative motion and defining a mixing volume therebetween, wherein the dwell time of a single pass of the flowing fluid material within and through the mixing volume is greater than 0.06 seconds or greater than 0.1 seconds; and introducing oxygen (O₂) into the flowing fluid material within the mixing volume under conditions suitable to dissolve at least 20 ppm, at least 25 ppm, at least 30, at least 40, at least 50, or at least 60 ppm oxygen into the material, and electrokinetically alter the fluid or solution. In certain aspects, the oxygen is infused into the material in less than 100 milliseconds, less than 200 milliseconds, less than 300 milliseconds, or less than 400 milliseconds. In particular embodiments, the ratio of surface area to the volume is at least 12, at least 20, at least 30, at least 40, or at least 50.

Yet further aspects provide a method of producing an electrokinetically-altered oxygenated aqueous fluid or solution, comprising: providing a flow of a fluid material between two spaced surfaces defining a mixing volume therebetween; and introducing oxygen into the flowing material within the mixing volume under conditions suitable to infuse at least 20 ppm, at least 25 ppm, at least 30 ppm, at least 40 ppm, at least 50 ppm, or at least 60 ppm oxygen into the material in less than 100 milliseconds, less than 200 milliseconds, less than 300 milliseconds, or less than 400 milliseconds. In certain aspects, the dwell time of the flowing material within the mixing volume is greater than 0.06 seconds or greater than 0.1 seconds. In particular embodiments, the ratio of surface area to the volume is at least 12, at least 20, at least 30, at least 40, or at least 50.

Additional embodiments provide a method of producing an electrokinetically-altered oxygenated aqueous fluid or solution, comprising use of a mixing device for creating an output mixture by mixing a first material and a second material, the device

comprising: a first chamber configured to receive the first material from a source of the first material; a stator; a rotor having an axis of rotation, the rotor being disposed inside the stator and configured to rotate about the axis of rotation therein, at least one of the rotor and stator having a plurality of through-holes; a mixing chamber defined between
5 the rotor and the stator, the mixing chamber being in fluid communication with the first chamber and configured to receive the first material therefrom, and the second material being provided to the mixing chamber via the plurality of through-holes formed in the one of the rotor and stator; a second chamber in fluid communication with the mixing chamber and configured to receive the output material therefrom; and a first internal
10 pump housed inside the first chamber, the first internal pump being configured to pump the first material from the first chamber into the mixing chamber. In certain aspects, the first internal pump is configured to impart a circumferential velocity into the first material before it enters the mixing chamber.

Further embodiments provide a method of producing an electrokinetically-altered
15 oxygenated aqueous fluid or solution, comprising use of a mixing device for creating an output mixture by mixing a first material and a second material, the device comprising: a stator; a rotor having an axis of rotation, the rotor being disposed inside the stator and configured to rotate about the axis of rotation therein; a mixing chamber defined
20 between the rotor and the stator, the mixing chamber having an open first end through which the first material enters the mixing chamber and an open second end through which the output material exits the mixing chamber, the second material entering the mixing chamber through at least one of the rotor and the stator; a first chamber in communication with at least a majority portion of the open first end of the mixing
25 chamber; and a second chamber in communication with the open second end of the mixing chamber.

Additional aspects provide an electrokinetically-altered oxygenated aqueous fluid or solution made according to any of the above methods.

Diabetes and Insulin Resistance

30 Diabetes mellitus is a serious lifelong metabolic disease that is defined by the presence of chronically elevated levels of blood glucose (hyperglycemia). This state of hyperglycemia is the result of a relative or absolute lack of activity of the peptide

hormone, insulin. Insulin is produced and secreted by the β cells of the pancreas. Insulin promotes glucose utilization, protein synthesis, and the formation and storage of carbohydrate energy as glycogen. Glucose is stored in the body as glycogen, a form of polymerized glucose, which may be converted back into glucose to meet metabolism requirements. Under normal conditions, insulin is secreted at both a basal rate and at enhanced rates following glucose stimulation, all to maintain metabolic homeostasis by the conversion of glucose into glycogen.

The term diabetes mellitus encompasses several different hyperglycemic states. These states include Type 1 (insulin-dependent diabetes mellitus or IDDM) and Type 2 (non-insulin dependent diabetes mellitus or NIDDM) diabetes. Type 1 diabetes is associated with deficient, reduced, or nonexistent levels of insulin that are insufficient to maintain blood glucose levels within the physiological range.

Type 2 diabetes is an increasingly prevalent disease of aging. It is initially characterized by decreased sensitivity to insulin or insulin resistance and a compensatory elevation in circulating insulin concentrations, the latter of which is required to maintain normal blood glucose levels. Increased insulin levels are caused by increased secretion from the pancreatic β cells, and the resulting hyperinsulinemia is associated with cardiovascular complications of diabetes. As insulin resistance worsens, the demand on the pancreatic β cells steadily increases until the pancreas can no longer provide adequate levels of insulin, resulting in elevated levels of glucose in the blood. Ultimately, overt hyperglycemia and hyperlipidemia occur, leading to the devastating long-term complications associated with diabetes, including cardiovascular disease, renal failure, and blindness.

One specific complication resulting from impaired cardiovascular circulation caused by Type II diabetes is diabetic retinopathy, where the eye and, more specifically, the retina undergoes damage due reduced cardiovascular functioning. In the retina the damage from lack of circulation initially manifests itself by weakening of the arteries which causes them to leak. This leakage results in small, dot-like hemorrhages and swelling. As the disease progresses the circulation problems cause parts of the retina to become ischemic or oxygen-deprived, resulting in new, fragile blood vessels developing as the circulatory system tries to maintain appropriate oxygen levels within the retina. This process, called neovascularization, leads to frequent leaks

of blood into the retina, because the new vessels are delicate and hemorrhage easily. In the later phases of diabetes, abnormal vessel growth continues and scar tissue develops. The result from the weak vessels and scar tissue can be quite serious, often leading to retinal detachment, glaucoma, and blindness.

5 The exact mechanism(s) causing type 2 diabetes are unknown, but result in impaired glucose transport into skeletal muscle and increased hepatic glucose production, in addition to inadequate insulin response. Dietary modifications are often ineffective, therefore the majority of patients ultimately require pharmaceutical intervention in an effort to prevent and/or slow the progression of the complications of
10 the disease. Many patients can be treated with one or more of the many oral anti-diabetic agents available, including insulin, alpha-glucosidase inhibitors, biguanides, DPP-4 inhibitors, meglitinides, sulfonylureas, and thiazolidinediones.

Alpha-glucosidase inhibitors (including, acarbose and miglitol) are saccharides that competitively inhibit certain enzymes that digest carbohydrates, specifically alpha-
15 glucosidase enzymes in the small intestines, where the membrane-bound alpha-glucosidases hydrolyze oligosaccharides, trisaccharides, and disaccharides into glucose and other monosaccharides.

Biguanides (such as metformin, buformin, and phenformin) reduce the level of serum glucose by inhibiting liver gluconeogenesis and by increasing the absorption of
20 glucose already present in the blood stream. DPP-4 inhibitors (including vildagliptin, sitagliptin, saxagliptin, linagliptin, and alogliptin) are thought to increase incretin levels (GLP-1 and GIP). Incretins inhibit the release of glucagon which limits glucose levels, increases insulin secretion, and decreases gastric emptying.

Sulfonylureas (including acetohexamide, chlorpropamide, tolbutamide, tolazamide, glipizide, gliclazide, glibenclamide (glyburide), gliquidone, glycopyramide, and glimepiride) bind to ATP-dependent K^+ (K_{ATP}) channel on the cell membrane of
25 pancreatic β cells, thereby increasing the secretion of (pro)insulin. Sulfonylureas also are believed to 1) sensitize β -cells to glucose, 2) limit glucose production in the liver, 3) decrease lipolysis (breakdown and release of fatty acids by adipose tissue) and 4)
30 decrease clearance of insulin by the liver.

Meglitinides (including nateglinide, mitiglinide, and repaglinide) mechanism of action is similar to that of sulfonylureas. Meglitinides bind to ATP-dependent K^+ (K_{ATP})

channel on the cell membrane of pancreatic β cells, which in turn increases the secretion of (pro)insulin.

Thiazolidinediones (including troglitazone, pioglitazone, and rosiglitazone) bind to peroxisome proliferator-activated receptors, a group of receptor molecules inside the cell nucleus. The normal ligands for these receptors are free fatty acids (FFAs) and eicosanoids. When activated, the receptor migrates to the DNA, activating transcription of a number of specific genes. The activation of these different genes results in 1) decreasing insulin resistance, 2) modifying adipocyte differentiation, 3) inhibiting VEGF-induced angiogenesis, 4) decreasing leptin levels (leading to an increased appetite), 5) decreasing certain interleukins (e.g. IL-6) levels, and 6) increasing adiponectin levels.

Recent scientific discoveries have suggested a link between chronic inflammation and insulin resistance, which is a precursor to diabetes. Solinas et al., *Cell Metabolism* 6, 386-397 (2007) (hereby incorporated by reference in its entirety); Duncan et al., *Diabetes* 52, 1799-1805 (2003) (hereby incorporated by reference in its entirety); Kathryn E. Wellen and Gökhan S. Hotamisligil, *J. Clin. Invest.* 115, 1111-1119 (2005) (hereby incorporated by reference in its entirety). More specifically, inflammation induces insulin resistance by promoting serine-phosphorylation of insulin receptor substrate-1 (IRS-1), which impairs insulin signaling. By impairing insulin signaling, individual cells become more resistant to serum insulin and instead signal for the release of glucose (from glycogen) into the blood stream. This suggests that pharmacological targeting of inflammation, in particular, might lower insulin resistance thereby preventing diabetes. In addition, using a dual approach, combining anti-inflammation therapy with diabetes drug treatment might result in enhanced relief of insulin resistance and diabetes symptoms.

Inflammation

Inflammation may occur as a defensive response to invasion of the subject by foreign material, particularly of microbial origin. Additionally, mechanical trauma, toxins, and neoplasia may induce inflammatory responses. The accumulation and subsequent activation of leukocytes are central events in the pathogenesis of most forms of inflammation. Inflammation deficiencies can compromise the host, leaving it susceptible to worsening infection or trauma. Excessive inflammation, such as

prolonged inflammatory responses, may lead to inflammatory diseases including but not limited to insulin resistance, diabetes, arteriosclerosis, cataracts, chronic skin disorders, reperfusion injury, and cancer, to post-infectious syndromes such as in infectious meningitis, rheumatic fever, and to rheumatic diseases such as systemic lupus erythematosus and rheumatoid arthritis. These diseases affect millions of people worldwide every year, and lead to increased mortality and morbidity. The commonality of the inflammatory response in these varied disease processes makes its regulation a major element in the prevention, or treatment, of human disease.

Overproduction of pro-inflammatory cytokines has been implicated in the pathogenesis of numerous inflammatory and autoimmune diseases. Secretion of TNFa is a primary event in the initiation of the inflammatory cascade (Brennan F. M., et. al. *Lancet*, 1989, 2:244-7; Haworth C, et. al. *Eur. J. Immunol.* 1991, 21:2575-2579) and directly contributes to the initiation and maintenance of these diseases, particularly insulin resistance and diabetes. Other cytokines also play a role, including interleukin 1 β (IL-1 β), IL-6, IL-8, IL-12, nitric oxide (NO), IFN- γ , granulocyte colony stimulating factor (G-CSF), granulocyte macrophage-colony stimulating factor (GM-CSF), and IL-10. Certain of these cytokines (e.g., IL-8) may increase or exacerbate an inflammatory response, while others (e.g., IL-10) may decrease or alleviate the inflammatory response.

Cells of the immune system, macrophages in particular, secrete many of these cytokines in response to activating stimuli. Target cells of the cytokines may be localized in any body compartment and may act via long-distance mechanisms, or may act on neighboring cells. Thus, cytokines may regulate inflammation in a localized or systemic manner.

Thymic stromal lymphopoietin (TSLP). Thymic stromal lymphopoietin (TSLP) is an IL-7-like cytokine that triggers dendritic cell-mediated Th2-type inflammatory responses and is considered as a master switch for allergic inflammation. TSLP is an integral growth factor to both B and T cell development and maturation. Particularly, murine TSLP supports B lymphopoieses and is required for B cell proliferation. Murine TSLP plays a crucial role in controlling the rearrangement of the T cell receptor-gamma (TCR.gamma.) locus and has a substantial stimulatory effect on thymocytes and mature T cells. See, for example, Friend et al., *Exp. Hematol.*, 22:321-328, 1994; Ray

et al., *Eur. J. Immunol.*, 26:10-16, 1996; Candeias et al., *Immunology Letters*, 57:9-14, 1997.

TSLP possesses cytokine activity similar to IL-7. For instance, TSLP can replace IL-7 in stimulating B cell proliferation responses (Friend et al., *supra*). Although
5 TSLP and IL-7 mediate similar effects on target cells, they appear to have distinct signaling pathways and likely vary in their biologic response. For Example, although TSLP modulates the activity of STAT5, it fails to activate any Janus family tyrosine kinase members (Levin et. al., *J. Immunol.*, 162:677-683, 1999).

TSLP effects on dendritic cells and TNF production. After human TSLP and the
10 human TSLP receptor were cloned in 2001, it was discovered that human TSLP potently activated immature CD11c⁺ myeloid dendritic cells (mDCs) (see, e.g., Reche et al., *J. Immunol.*, 167:336-343, 2001 and Soumelis et al., *Nat. Immunol.*, 3:673-680, 2002). Th2 cells are generally defined in immunology textbooks and literature as CD4⁺ T cells that produce IL-4, IL-5, IL-13, and IL-10, and Th1 cells such as CD4⁺ T cells
15 produce IFN- γ and sometimes TNF. When TSLP-DCs are used to stimulate naive allogeneic CD4⁺ T cells in vitro, a unique type of Th2 cell is induced which produces the classical Th2 cytokines IL-4, IL-5, and IL-13, and large amounts of TNF, but little or no IL-10 or interferon γ (Reche et al., *supra*) (see also, e.g., Soumelis et al., *Nat. Immunol.*, 3:673-680, 2002). TNF is not typically considered a Th2 cytokine. However,
20 TNF is prominent in asthmatic airways and genotypes that correlate with increased TNF secretion are associated with an increased asthma risk. See Shah et al., *Clin. Exp. Allergy.*, 25:1038-1044, 1995 and Moffatt, M.F. and Cookson, W.O., *Hum. Mol. Genet.*, 6:551-554, 1997.

TSLP induces human mDCs to express the TNF superfamily protein OX40L at
25 both the mRNA and protein level (Ito et al., *J. Exp. Med.*, 202:1213-1223). The expression of OX40L by TSLP-DCs is important for the elaboration of inflammatory Th2 cells. Thus, TSLP-activated DCs create a Th2-permissive microenvironment by up-regulating OX40L without inducing the production of Th1-polarizing cytokines. *Id.*

TSLP expression, allergen-specific responses, and asthma. In Early studies
30 have shown that TSLP mRNA was highly expressed by human primary skin keratinocytes, bronchial epithelial cells, smooth muscle cells, and lung fibroblasts (Soumelis et al., *Nat. Immunol.*, 3:673-680, 2002). Because TSLP is expressed mainly

in keratinocytes of the apical layers of the epidermis, this suggests that TSLP production is a feature of fully differentiated keratinocytes. TSLP expression in patients with atopic dermatitis was associated with Langerhans cell migration and activation in situ which suggests that TSLP may contribute directly to the activation of these cells which could subsequently migrate into the draining lymph nodes and prime allergen-specific responses. *Id.* In a more recent study, it was shown by in situ hybridization that TSLP expression was increased in asthmatic airways and correlated with both the expression of Th2-attracting chemokines and with disease severity which provided a link between TSLP and asthma (Ying et al., *J. Immunol.*, 174:8183-8190, 2005).

TSLP receptor (TSLPR) and allergy, asthma. The TSLP receptor (TSLPR) is approximately 50 kDa protein and has significant similarity to the common γ -chain. TSLPR is a novel type 1 cytokine receptor, which, combined with IL-7Ra (CD127), constitutes a TSLP receptor complex as described, for example, in Pandey et al., *Nat. Immunol.*, 1:59-64, 2000. TSLPR has a tyrosine residue near its carboxyl terminus, which can associate with phosphorylated STAT5 and mediate multiple biological functions when engaged with TSLP (Isaksen et al., *J. Immunol.*, 168:3288-3294, 2002).

Human TSLPR is expressed by monocytes and CD11c+ dendritic cells, and TSLP binding induces the expression of the Th2 cell-attracting chemokines CCL17 and CCL22. Furthermore, as stated above, the TSLPR-induced activation of dendritic cells indirectly results in the increased secretion of Th2 cytokines IL-4, -5 and -13, which may be necessary for the regulation of CD4+ T cell homeostasis. In mice, deficiency of TSLPR has no effect on lymphocyte numbers. However, a deficiency of TSLPR and common γ -chain results in fewer lymphocytes as compared to mice deficient in the common γ -chain alone. See Reche et al., *J. Immunol.*, 167:336-343, 2001 and Soumelis et al., *Nat. Immunol.*, 3:673-680, 2002.

Studies have found that TSLP and the TSLPR play a critical role in the initiation of allergic diseases in mice. In one study, it was demonstrated that mice engineered to overexpress TSLP in the skin developed atopic dermatitis which is characterized by eczematous skin lesions containing inflammatory infiltrates, a dramatic increase in circulating Th2 cells and elevated serum IgE (Yoo et al., *J. Exp. Med.*, 202:541-549, 2005). The study suggested that TSLP may directly activate DCs in mice. In another study, conducted by Li et al., the group confirmed that transgenic mice overexpressing

TSLP in the skin developed atopic dermatitis which solidifies the link between TSLP and the development of atopic dermatitis.

Another set of studies demonstrated that TSLP is required for the initiation of allergic airway inflammation in mice in vivo. In one study, Zhou et al. demonstrated that
5 lunch specific expression of a TSLP transgene induced allergic airway inflammation (asthma) which is characterized by massive infiltration of leukocytes (including Th2 cells), goblet cell hyperplasia, and subepithelial fibrosis, and increased serum IgE levels (Zhou et al., *Nat. Immunol.*, 6:1047-1053, 2005). However, in contrast, mice lacking the TSLPR failed to develop asthma in response to inhaled antigens (Zhou et al., *supra* and
10 Al-Shami et al., *J. Exp. Med.*, 202:829-839, 2005). Thus, these studies together demonstrate that TSLP is required for the initiation of allergic airway inflammation in mice.

Further, in a study conducted by Yong-Jun et al., it was demonstrated that epithelial cell-derived TSLP triggers DC-mediated inflammatory Th2 responses in
15 humans which suggest that TSLP represents a master switch of allergic inflammation at the epithelial cell-DC interface (Yong-Jun et al., *J. Exp. Med.*, 203:269-273, 2006).

In a recent study, it was shown that modulation of DCs function by inhibiting the TSLPR lessened the severity in mice (Liyun Shi et al., *Clin. Immunol.*, 129:202-210, 2008). In another set of studies, it was demonstrated that the TSLPR was not only
20 expressed in DCs, but also on macrophages, mast cells, and CD4+ T cells (Rochman et al., *J. Immunol.*, 178:6720-6724, 2007 and Omori M. and Ziegler S., *J. Immunol.*, 178:1396-1404, 2007). In order to rule out the direct effects of the TSLPR neutralization on CD4+ T cells or other effector cells in allergic inflammation, Liyun Shi et al. performed experiments wherein OVA-loaded DCs were in vitro treated with anti-
25 TSLPR before adoptive transfer to the airways of naive mice. It has previously been found that OVA-DCs triggered strong eosinophilic airway inflammation and accompanied with massive production of Th2 cytokines such as IL-4 and IL-5 (Sung et al., *J. Immunol.*, 166:1261-1271 and Lambrecht et al., *J. Clin. Invest.*, 106:551-559, 2000). However, pretreating OVA-DCs with anti-TSLPR resulted in a significant
30 reduction of eosinophils and lymphocyte infiltration as well as IL-4 and IL-5 levels, further illuminating the role that TSLPR plays in DC-primed allergic disease. This result

also supports that blocking of TSLPR on DCs will aid in controlling airway inflammation (Liyun Shi et al., *supra*).

There has been a growing body of experiments implicating the role of TSLP/TSLPR in various physiological and pathological processes. Physiological roles of TSLP include modulating the immune system, particularly in stimulating B and T cell proliferation, development, and maturation. TSLP plays a vital role in the pathobiology of allergic asthma and local antibody mediated blockade of TSLP receptor function to alleviate allergic diseases. Thus, interplay between TSLP and TSLP receptor is believed to be important in many physiological disease processes such as: allergic inflammation, skin lesions of patients with atopic dermatitis or atopic eczema, insulin resistance or diabetes, and asthma.

Inflammation-Associated Conditions or Diseases

Certain embodiments herein relate to therapeutic compositions and methods of treatment for a subject by preventing or alleviating at least one symptom of inflammation associated with certain conditions or diseases. For example, the therapeutic compositions and/or methods disclosed herein may be useful for treating or preventing insulin resistance and diabetes.

Methods of Treatment

The term “treating” refers to, and includes, reversing, alleviating, inhibiting the progress of, or preventing a disease, disorder or condition, or one or more symptoms thereof; and “treatment” and “therapeutically” refer to the act of treating, as defined herein.

A “therapeutically effective amount” is any amount of any of the compounds utilized in the course of practicing the invention provided herein that is sufficient to reverse, alleviate, inhibit the progress of, or prevent a disease, disorder or condition, or one or more symptoms thereof.

Certain embodiments herein relate to therapeutic compositions and methods of treatment of a diabetes-associated condition or disorder, or symptoms thereof. For

example, the therapeutic compositions and/or methods disclosed herein may be useful for treating or preventing one or more condition or disease selected from the group consisting of: diabetes; insulin-dependent diabetes mellitus or IDDM (Type 1); non-insulin dependent diabetes mellitus or NIDDM (Type 2); insulin resistance; and diabetic
5 retinopathy.

Many conditions or diseases associated with inflammation have been treated with steroids, methotrexate, immunosuppressive drugs including cyclophosphamide, cyclosporine, azathioprine and leflunomide, nonsteroidal anti-inflammatory agents such as aspirin, acetaminophen and COX-2 inhibitors, gold agents and anti-malarial
10 treatments. These drugs have a variety of disadvantages, and adverse reactions including injection site reactions, rash, upper respiratory infections, autoimmune disorders and increased susceptibility to infections. In addition, many anti-inflammatory pharmaceutical drugs require intravenous (IV) or subcutaneous (SC) administration, as opposed to more convenient and compliant oral or topical dermal routes. Accordingly,
15 a need still exists for the development of novel medicaments and treatment methods for conditions and diseases relating to inflammation.

Combination therapy

Additional aspects provide the herein disclosed inventive methods, further comprising combination therapy, wherein at least one additional therapeutic agent is administered to the patient. In certain aspects, the at least one additional therapeutic agent is selected from the group consisting of insulin, alpha-glucosidase inhibitors, biguanides, DPP-4 inhibitors, meglitinides, sulfonylureas, and thiazolidinediones.
20

In particular embodiments, the at least one additional therapeutic agent is selected from the group consisting of: alpha-glucosidase inhibitors consisting of acarbose and miglitol; biguanides consisting of metformin, buformin, and phenformin; DPP-4 inhibitors, consisting of vildagliptin, sitagliptin, saxagliptin, linagliptin, and alogliptin; sulfonylureas, consisting of acetohexamide, chlorpropamide, tolbutamide, tolazamide, glipizide, gliclazide, glibenclamide (glyburide), gliquidone, glycopyramide,
25 and glimepiride; meglitinides consisting of nateglinide, mitiglinide, and repaglinide;
30

thiazolidinediones, consisting of troglitazone, pioglitazone, and rosiglitazone; and combinations thereof.

Additional aspects provide the herein-disclosed inventive methods, further comprising combination therapy with a TSLP and/or TSLPR antagonist (e.g., neutralizing antibodies specific for TSLP and the TSLP receptor, soluble TSLP receptor molecules, and TSLP receptor fusion proteins, such as TSLPR-immunoglobulin Fc molecules or polypeptides that encode components of more than one receptor chain, that thereby mimic a physiological receptor heterodimer or higher order oligomer. If the receptor includes more than one polypeptide chain, a single chain fusion can be utilized).

Therapeutic treatments for inflammation and/or diabetes include a wide array of pharmaceutical drugs administered intravenously, subcutaneously, topically, or orally depending on the outcome sought. However, most of the anti-inflammatory treatments available today have considerable drawbacks, including severe reactions at injection site, increased susceptibility to infection, rash, or other side effects. Thus, there is a need for better anti-inflammatory therapeutics and treatment methods.

Anti-inflammatory Activity of the Electrokinetically-generated Gas-Enriched Fluids and Solutions:

According to certain aspects of the present invention, the gas-enriched fluids and/or solutions disclosed herein have anti-inflammatory properties and effects, and can be used as anti-inflammatory agents for the treatment of subjects afflicted by diseases or disorders relating to inflammation. Figure 38 shows the experimental results of cytokine profiles in stimulated lymphocytes from a healthy blood donor. As can be seen in Figure 38, the inventive oxygen-enriched fluid (water) affected a down regulation of particular cytokines, especially IL-6, IL-8, and IL-1 β .

Increased production of pro-inflammatory cytokines has been implicated in the pathogenesis of numerous inflammatory and autoimmune diseases. Secretion of TNFa is a primary event in the initiation of the inflammatory cascade (Brennan F. M., et. al. *Lancet*, 1989, 2:244-7; Haworth C, et. al. *Eur. J. Immunol.* 1991, 21:2575-2579) and directly contributes to the initiation and maintenance of inflammatory and autoimmune diseases. Other pro-inflammatory cytokines also play a role, including interleukin 1 β

(IL-1 β), IL-6, IL-8, IL-12 nitric oxide, IFN- γ and GM-CSF, while anti-inflammatory cytokines such as IL-10 may reduce disease. Cells of the immune system, macrophages in particular, secrete many of these cytokines in response to activating stimuli.

5 A variety of cell types are involved in the inflammatory process. Overproduction of TNFa by monocytes, macrophages, and other immune cells is a key element in the pathogenesis of a multitude of diseases. Macrophages and T cells in particular play a central role in the initiation and maintenance of the immune response. Once activated by pathological or immunogenic stimuli, macrophages respond by releasing a host of
10 cytokines, including TNF- α , IL-1 β , IL-8, IL-12, nitric oxide (NO), IL-6, GM-CSF, G-CSF, M-CSF and others. T cells release IL-2, IL-4, INF- γ , and other inflammatory cytokines. These cytokines activate other immune cells and some can also act as independent cytotoxic agents. Excessive release of macrophage and T cell-derived inflammatory mediators can particularly lead to damage of normal cells and surrounding tissues.

15 Pro-inflammatory cytokines have been implicated to contribute to many diseases and disease symptoms including diabetes. The induction of NO from smooth muscle cells mediates decreased mean arterial pressure and systemic vascular resistance during septic shock, suggesting a fundamental role for NO. Thus, therapies that target downregulatory effects on IL-8, IL-1 β , and NO could be beneficial in the treatment of
20 inflammatory diseases or disorders, including sepsis, septic shock, endotoxic shock, and diabetes.

Overproduction of TNFa contributes to the clinical features of numerous autoimmune diseases such as diabetes and rheumatoid arthritis. Systemic lupus erythematosus (SLE) is also precipitated by increased IL-1 β and TNFa levels. Within
25 lupus patients, serum C-reactive protein, IL-1 β , and TNFa levels were higher than in controls, suggesting that an increased inflammatory response plays a role in the disease (Liou L. B. *Clin. Exp. Rheumatol.* 2001, 19:515-523). A study of patients with one form of SLE, neuropsychiatric lupus erythematosus (NPLE), showed that the number of peripheral blood mononuclear cells expressing mRNA for TNFa as well as
30 the cerebrospinal fluid level of NO metabolites correlated with NPLE disease severity (Svenungsson E., et al. *Ann. Rheum. Dis.* 2001, 60:372-9).

IL-1 and TNFa play a central role in various acute as well as chronic responses in animal models. Additionally, IL-11, IFNa, and IFN β may also up-regulate inflammatory reactions. Conversely, several cytokines may be involved in down-regulation of inflammatory responses (i.e. IL-4, IL-10, IL-13, among others). As set forth in Example 1, cells contacted with the inventive gas-enriched fluid showed an increase in IFN- γ levels with T3 antigen than in the control culture media with T3 antigen, while IL-8 was lower in the inventive gas-enriched culture media with T3 antigen than in the control culture media with T3 antigen. Additionally, IL-6, IL-8, and TNF-a levels were lower in the inventive gas-enriched media with PHA, than in the control media with PHA, while IL-1 β levels were lower in the inventive gas-enriched fluid with PHA when compared with control media with PHA. In the inventive gas-enriched media alone, IFN- γ levels were higher than in control media. These results are consistent with an anti-inflammatory microenvironment.

NO is recognized as a mediator and regulator of inflammatory responses. It possesses cytotoxic properties toward pathogens, but can also have deleterious effects on the subject's own tissues. (Korhonen et al., *Curr Drug Targets Inflamm Allergy* 4(4): 471-9, 2005). NO reacts with soluble guanylate cyclase to form cyclic guanosine monophosphate (cGMP), which mediates many of the effects of NO. NO can also interact with molecular oxygen and superoxide anion to produce reactive oxygen species that can modify various cellular functions. These indirect effects of NO have a significant role in inflammation, where NO is produced in high amounts by inducible NO synthase (iNOS) and reactive oxygen species are synthesized by activated inflammatory cells. In fact, overproduction of NO contributes to impairment of both muscle cell insulin action and β cell function in obesity, which can lead to insulin resistance and diabetes.

NO can be produced by keratinocytes, fibroblasts, endothelial cells, and possibly others. Some of the vascular actions of NO include vasodilation, inhibiting platelet adhesion to the vascular endothelium, inhibiting leukocyte adhesion to the vascular endothelium, and scavenging superoxides. (Shah et al., *Env. Health Persp.* v. 106 (5): 1139-1143.)

Furthermore, inhibition of NO synthesis has been shown to delay wound contraction, alter collagen organization, and alter neoepidermis thickness. (Amadeu

and Costa, *J. Cutan. Pathol.* 33: 465-473, 2006.) Mast cell migration and angiogenesis in wounds is also affected by inhibition of NO. (*Id.*) Without being bound to any particular theory of mechanism, in certain embodiments the inventive gas-enriched fluids may be modulating localized and/or cellular NO production, or degradation, consistent with the spectrum of wound healing effects illustrated in the Examples section disclosed herein. Due to variable pathways of regulation, in certain embodiments, the inventive gas-enriched fluid may increase NO production and/or retard NO degradation, whereas in other certain embodiments, the inventive gas-enriched fluid may decrease NO production and/or hasten NO degradation.

Specifically, wounds treated with oxygen-enriched saline solution showed an increase in wound healing at days 4 through 11, and between days 3 and 11, the new epidermis in wounds treated with the oxygen-enriched saline solution migrated at two to four times as fast as the epidermis of the wounds treated with the normal saline solution, as set forth in Example 8 herein. The study also showed that between 15 and 22 days, wounds treated by the oxygen-enriched saline solution differentiated at a more rapid rate as evidenced by the earlier formation of more mature epidermal layers. At all stages, the thickening that occurs in the epidermis associated with normal healing did not occur within the wounds treated by the oxygen-enriched saline solution.

Thus, in accordance with this spectrum of wound-healing effects, but without wishing to be bound by any particular theory, it is believed that the oxygen-enriched saline solution may modulate the localized and/or cellular level of NO within the wounds. NO modulates growth factors, collagen deposition, inflammation, mast cell migration, epidermal thickening, and neovascularization in wound healing. Furthermore, nitric oxide is produced by an inducible enzyme that is regulated by oxygen.

In the case of mast cell migration, differences also occurred in early and late migration for the oxygen-enriched solution. This is consistent with what is known in the art regarding inhibition of NO synthesis (Amadeu and Costa, *J. Cutan Pathol* 33: 465-473, 2006).

Referring now to Figure 41A through 41F, various illustrations compare the wound healing results of the porcine epidermal tissues with or without oxygen-enriched

saline solution. As can be seen, the healing of the control wound and of the wound using the oxygen-enriched saline solution was followed for days 1, 4, and 16.

Figure 41A illustrates the wound healing for the control wound on day 1. As can be seen, the wound shows epidermal/dermal thickening and a loss of contour. Figure 41B illustrates the wound healing on day 1 for the wound treated using the oxygen-enriched saline solution. The wound shows normal epidermal/dermal thickness and normal contouring is typical on a new wound.

Referring now to Figures 41C and 41D, there are illustrated the wound healing for the control wound on day 4 and the wound healing for the wound treated with the oxygen-enriched saline solution on day 4. For the control wound illustrated in Figure 41C, the wound shows a 600 micron epidermal spur. In the wound treated with the oxygen-enriched saline solution in Figure 41D, there is illustrated a 1200 micron epidermal spur. Thus, in the first 4 days of the experiment, the epidermal spur created in the wound treated using the oxygen-enriched saline solution shows an epidermal growth rate of twice of that of the wound that was not treated with the oxygen-enriched saline solution.

Referring now to Figure 41E, there is illustrated the control wound at day 16. The wound shows less differentiated epidermis with loss of epidermal/dermal contour than that illustrated by the wound treated with the oxygen-enriched saline solution, illustrated in Figure 41F. Figure 41F shows more differentiated epidermis and more normal epidermal/dermal contouring in the wound.

In the first two phases of the inflammatory process, the foreign body is either destroyed, for example, if the foreign body is an organism, or the tissue around it is loosened, for example, if it is a splinter. In the healing phase, the inflammation begins to subside, individual blood vessels and vascular patterns become normal once again, and repair of the wound commences. The three main events in the repair process are: (1) formation of new connective tissue by proliferating fibroblasts, (2) regeneration of epithelium, and (3) outgrowth of new capillaries.

Even before the inflammation subsides, fibroblasts begin moving into the injured area from the surrounding normal tissue, where they usually exist in a dormant state. They migrate by an amoeboid movement along strands of fibrin and distribute themselves throughout the healing area. Once fixed into position in the injured tissue,

they begin to synthesize collagen and secrete this protein, which arranges itself into fibers. The fibers orient themselves with their longitudinal axes in the direction of the greatest stress. As the collagen bundles grow in firmness, the fibroblasts gradually degenerate and attach closely to the bundles, and the injured area transforms into scar
5 tissue.

Simultaneously with scar tissue formation, the intact epidermal cells on the edge of the wound begin to proliferate and move, as one sheet, toward the center of the injured area. As the inflammation subsides, a need for a direct supply of blood arises, and angiogenesis occurs at the wound site.

10 Inflammation is a complex process that involves multiple cell types. For example, mast cells release mediators that trigger an early phase of vasodilation, accompanied by the separation of endothelial cells and exposure of collagen fibers in the subendothelial layer. Fibers in the intercellular gaps that form in blood vessels trap platelets and trigger the release of mediators from these cells.

15 In addition to platelets, the exposed collagen fibers also interact with proteins of the plasma that filter through the pores of the dilated vessel wall, including the triggering factor of the blood-clotting cascade, increased vasodilation, increased blood vessel permeability, and chemotaxis.

20 Additionally, the complement cascade can be activated by several stimuli: the injured blood vessels, the proteolytic enzymes released by the damaged cells, the membrane components of any participating bacteria, and antigen-antibody complexes. Some of the activated complement components act as chemotactic factors, responsible for the influx of leukocytes into the inflamed area, while others facilitate phagocytosis and participate in cell lysis.

25 In addition, it is believed that the inventive gas-enriched fluids or solutions may also regulate at least one cytokine involved in at least one aspect of inflammation, the cytokine(s) including, but not limited to MAF (macrophage activating factor), MMIF (macrophage migration inhibition factor), MCF (macrophage chemotactic factor), LMIF (leukocyte migration inhibition factor), HRFs (histamine releasing factors), TF (transfer
30 factors), interleukins (IL-1, IL-2, IL-3, IL-4, IL-5, IL-6, IL-7, IL-8, IL-9, IL-10, IL-11, IL-12, IL-13, IL-14, IL-15, etc.), TNF- α , TNF- β , interferons (IFN- α , IFN- β , IFN- γ , IFN- ζ , IFN-d, etc.), G-CSF (granulocyte colony stimulating factor), GM-CSF (granulocyte-

macrophage CSF), M-CSF (macrophage CSF), multi-CSF (IL-3), fibroblast growth factor (aFGF, bFGF), EGF (epidermal growth factor), NGF (nerve growth factor), PDGF (platelet-derived growth factor), VEGF (vascular endothelial growth factor), transforming growth factors (TGF- α , TGF- β , etc.), NAP-2 (neutrophil-activating protein 2), PF-4
 5 (platelet factor 4), thromboglobulin, MCP-1 (monocyte chemoattractant protein 1), MCP-3, MIP-1 α , MIP-1 β -+ (macrophage inflammatory proteins), RANTES (regulated upon activation normal T expressed and presumably secreted chemokine), HSPs (heat shock proteins), GRPs (glucose-regulated proteins), ubiquitin, and others.

Thus, in certain embodiments, the gas-enriched fluids and/or therapeutic
 10 compositions may increase production and/or secretion of anti-inflammatory molecules or cytokines or decrease the degradation of anti-inflammatory molecules or cytokines, thereby alleviating or preventing at least one symptom of inflammation. In other embodiments, the gas-enriched fluids and/or therapeutic compositions of the present invention may decrease production and/or secretion of pro-inflammatory molecules or
 15 cytokines or increase the degradation of pro-inflammatory molecules or cytokines, thereby alleviating or preventing at least one symptom of inflammation.

Previous studies had shown a critical role of anti-MOG antibodies in augmentation of demyelination and worsening of EAE (experimental autoimmune encephalomyelitis), an animal model system for the human autoimmune disorder of
 20 rheumatoid arthritis. (Linington, et al. 1992. *J. Neuroimmunol.* 40:219-224). Additionally, antibodies against MOG have been implicated in the pathogenesis of multiple sclerosis. (Berger et al. *N. Engl. J. Med.* 2003 Jul 10;349(2):139-45).

As set forth in Figure 48 and Example 12, the inventive gas-enriched fluid of the present invention amplifies the lymphocyte response to an antigen for which an animal
 25 was previously primed. As indicated in Figure 48, lymphocyte proliferation was greater for response to MOG challenge when cultured in fluid reconstituted with the inventive gas-enriched fluid comprising solvated electrons, when compared with pressurized, oxygenated fluid (pressure pot) or control deionized fluid.

30 Inventive Gas-Enriched Fluids and Solutions

Diffusing or enriching a fluid with another fluid may result in a solution or suspension of the two fluids. In particular, enriching a liquid with a gas (e.g. oxygen)

may be beneficial for certain applications, including therapeutic treatments. As utilized herein, "fluid," may generally refer to a liquid, a gas, a vapor, a mixture of liquids and/or gases, or any combination thereof, for any particular disclosed embodiment. Furthermore, in certain embodiments a "liquid" may generally refer to a pure liquid or
5 may refer to a gel, sol, emulsion, fluid, colloid, dispersion, or mixture, as well as any combination thereof, any of which may vary in viscosity.

In particular embodiments disclosed herein, the dissolved gas comprises ambient air. In a preferred embodiment, the dissolved gas comprises oxygen. In another embodiment, the dissolved gas comprises nitric oxide.

10 There are several art-recognized methods of gas-enriching liquids (such as oxygen-enriching water). For example, a turbine aeration system can release air near a set of rotating blades of an impeller, which mixes the air or oxygen with the water, or water can be sprayed into the air to increase its oxygen content. Additionally, other systems on the market inject air or oxygen into the water and subject the water/gas to a
15 large-scale vortex. Naturally occurring levels of oxygen in water are typically no more than 10 ppm (parts per million), which is considered to be a level of 100% dissolved oxygen. Tests on certain devices have shown that under ideal conditions, the device can attain upwards of approximately 20 ppm, or twice the natural oxygen levels of water. In certain embodiments, the oxygen level may be even higher.

20 In certain embodiments disclosed herein, a gas-enriched fluid of the present invention provides an anti-inflammatory benefit. Certain embodiments disclosed herein relate to a therapeutic composition comprising a gas-enriched fluid of the present invention, and optionally at least one additional therapeutic agent, such as a pharmaceutical drug, a metal, a peptide, a polypeptide, a protein, a nucleotide, a
25 carbohydrate or glycosylated protein, a fat (including oils or waxes), or other agent that prevents or alleviates at least one symptom of a condition or disease associated with inflammation.

Furthermore, certain embodiments disclosed herein include therapeutic compositions and methods related to inflammation of wounds. Wound care is desirable
30 to improve health and appearance of underlying dermal tissues. Wounds, either injury induced, such as cuts, abrasions or blisters, or surgically induced, such as surgical incisions or ostomies, require localized treatment to remedy the affected area and to

prevent further dermal damage. If wounds are not properly treated, further dermal irritation can result, such as inflammation, and may result in secondary infections and further discomfort to the subject.

Particular embodiments provided herein relate to a diffuser-processed therapeutic fluid as defined herein, comprising: a fluid host material; an infusion material diffused into the host material; and optionally, at least one therapeutic agent dispersed in the host material, wherein the infusion material comprises oxygen micro-bubbles in the host fluid, wherein the majority of the micro-bubbles are less than 0.2 microns, or preferably less than 0.1 microns in size. In certain embodiments, the dissolved oxygen level in the infused fluid host material may be maintained at greater than about 30 ppm at atmospheric pressure for at least 13 hours. In other particular embodiments, the dissolved oxygen level in the infused fluid host material may be maintained at greater than 40 ppm at atmospheric pressure for at least 3 hours.

In additional embodiments, the infused fluid host material further comprises a saline solution. In further embodiments, the infused fluid host material maintains a dissolved oxygen level of at least about 20 ppm to about 40 ppm for a period of at least 100 days, preferably at least 365 days within a sealed container at atmospheric pressure. In certain embodiments, the infused fluid host material may have a dissolved oxygen level of at least 50 ppm at atmospheric pressure.

In certain embodiments, the infused fluid host material exhibits Rayleigh scattering for a laser beam shining therethrough for a selected period of time after the oxygen has been diffused into therein.

Table 3 illustrates various partial pressure measurements taken in a healing wound treated with an oxygen-enriched saline solution and in samples of the gas-enriched oxygen-enriched saline solution of the present invention.

TABLE 3

TISSUE OXYGEN MEASUREMENTS	
Probe Z082BO	
In air: 171 mmHg	23° C
Column	Partial Pressure (mmHg)
B1	32-36
B2	169-200

B3	20-180*
B4	40-60
*wound depth minimal, majority >150, occasional 20 s	

BUBBLE SIZE MEASUREMENTS

Experimentation was performed to determine a size of the bubbles of gas diffused within the fluid by the mixing device 100. While experiments were not performed to measure directly the size of the bubbles, experiments were performed that established that the bubble size of the majority of the gas bubbles within the fluid was smaller than 0.1 microns. In other words, the experiments determined a size threshold value below which the sizes of the majority of bubbles fall.

This size threshold value or size limit was established by passing the output material 102 formed by processing a fluid and a gas in the mixing device 100 through a 0.22 filter and a 0.1 micron filter. In performing these tests, a volume of the first material 110, in this case, a fluid, and a volume of the second material 120, in this case, a gas, were passed through the mixing device 100 to generate a volume of the output material 102 (i.e., a fluid having a gas diffused therein). Sixty milliliters of the output material 102 was drained into a 60 ml syringe. The DO level of the fluid within the syringe was then measured using an Orion 862a. The Orion 862a is capable of measuring DO levels within a fluid. The fluid within the syringe was injected through a 0.22 micron filter into a 50 ml beaker. The filter comprised the Milipor Millex GP50 filter. The DO level of the material in the 50 ml beaker was then measured. The experiment was performed three times to achieve the results illustrated in Table 4 below.

TABLE 4

DO IN SYRINGE	DO AFTER 0.22 MICRON FILTER
42.1 ppm	39.7 ppm
43.4 ppm	42.0 ppm
43.5 ppm	39.5 ppm

As can be seen, the DO levels measured within the syringe and the DO levels measured within the 50 ml beaker were not changed drastically by passing the output

material 102 through the 0.22 micron filter. The implication of this experiment is that the bubbles of dissolved gas within the output material 102 are not larger than 0.22 microns otherwise there would be a significantly greater reduction in the DO levels in the output material 102 passed through the 0.22 micron filter.

5 A second test was performed in which the 0.1 micron filter was substituted for the 0.22 micron filter. In this experiment, saline solution was processed with oxygen in the mixing device 100 and a sample of the output material 102 was collected in an unfiltered state. The DO level of the unfiltered sample was 44.7 ppm. The output material 102 was filtered using the 0.1 micron filter and two additional samples were
10 collected. The DO level of the first sample was 43.4 ppm. The DO level of the second sample was 41.4 ppm. Then, the filter was removed and a final sample was taken from the unfiltered output material 102. The final sample had a DO level of 45.4 ppm. These results were consistent with those seen using the Millipore 0.2 micron filter. These results lead to the conclusion that there is a trivial reduction in the DO levels of
15 the output material 102 passed through the 0.1 micron filter providing an indication that the majority of the bubbles in the processed saline solution are no greater than 0.1 micron in size. The DO level test results described above were achieved using Winkler Titration.

As appreciated in the art, the double-layer (interfacial) (DL) appears on the
20 surface of an object when it is placed into a liquid. This object, for example, might be that of a solid surface (e.g., rotor and stator surfaces), solid particles, gas bubbles, liquid droplets, or porous body. In the mixing device 100, bubble surfaces represent a significant portion of the total surface area present within the mixing chamber that may be available for electrokinetic double-layer effects. Therefore, in addition to the surface
25 area and retention time aspects discussed elsewhere herein, the relatively small bubble sizes generated within the mixer 100 compared to prior art devices 10, may also contribute, at least to some extent, to the overall electrokinetic effects and output fluid properties disclosed herein. Specifically, in preferred embodiments, as illustrated by the mixer 100, all of the gas is being introduced via apertures on the rotor (no gas is
30 being introduced through stator apertures). Because the rotor is rotating at a high rate (e.g., 3,400 rpm) generating substantial shear forces at and near the rotor surface, the bubble size of bubbles introduced via and adjacent to the spinning rotor surface

apertures would be expected to be substantially (e.g., 2- to 3-times smaller) smaller than those introduced via and near the stationary stator. The average bubble size of the prior art device 10 may, therefore, be substantially larger because at least half of the gas is introduced into the mixing chamber from the stationary stator apertures.

- 5 Because the surface area of a sphere surface varies with r^2 , any such bubble component of the electrokinetic surface area of the mixing device 100 may be substantially greater than that of the prior art diffusion device 10.

Therefore, without being bound by theory, not only does the mixing chamber of the mixing device 100 have (i) a substantially higher surface to volume ratio than that of
10 the prior art device 10 (the prior art device 10 has a ratio of surface to volume of 10.9, whereas the present mixer 100 has a surface to volume ratio of 39.4), along with (ii) a 7-fold greater dwell-time, but (iii) the unique properties of the current output solutions may additionally reflect a contribution from the substantially larger bubble surface area in the mixing device 100. These distinguishing aspects reflect distinguishing features of
15 the present mixer 100, and likely each contribute to the unique electrokinetic properties of the inventive output materials/fluids.

Referring now to Figure 30, there is illustrated the DO levels in water enriched with oxygen in the mixing device 100 and stored in a 500 ml thin-walled plastic bottle and a 1000 ml glass bottle out to at least 365 days. Each of the bottles was capped
20 and stored at 65° Fahrenheit. As can be seen in the figure, the DO levels of the oxygen-enriched fluid remained fairly constant out to at least 365 days.

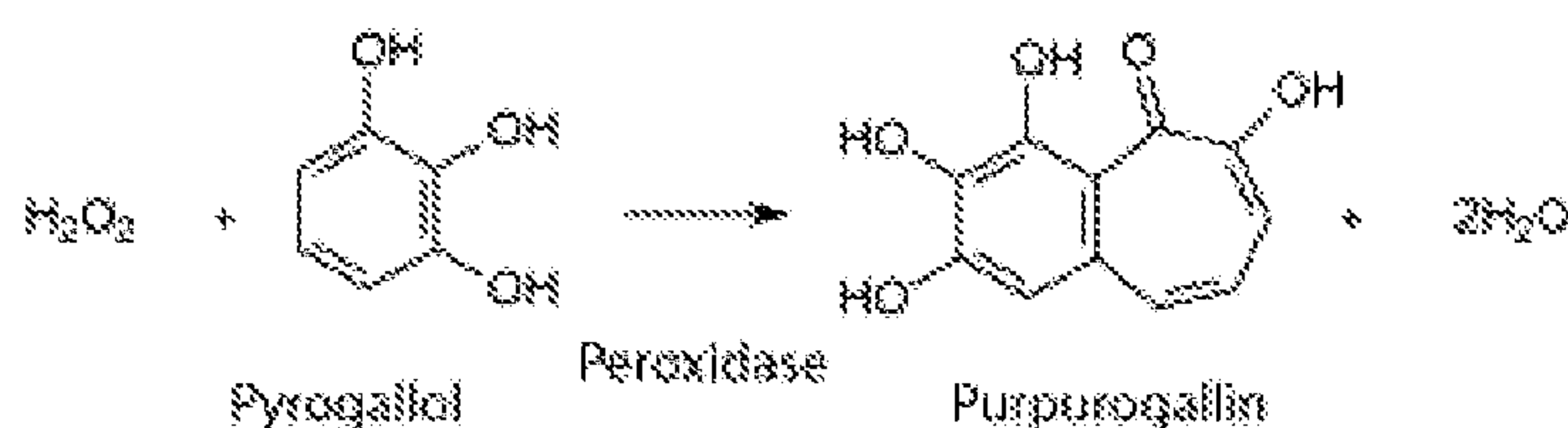
Referring to Figure 31, there is illustrated the DO levels in water enriched with oxygen in the mixing device 100 and stored in a 500 ml plastic thin-walled bottle and a 1000 ml glass bottle. Both bottles were refrigerated at 39° Fahrenheit. Again, DO
25 levels of the oxygen-enriched fluid remained steady and decreased only slightly out to at least 365 days.

Compositions comprising hydrated (solvated) electrons imparted to the inventive compositions by the inventive processes

30 In certain embodiments as described herein (see under “double-layer”), the gas-enriched fluid is generated by the disclosed electromechanical processes in which molecular oxygen is diffused or mixed into the fluid and may operate to stabilize

charges (e.g., hydrated (solvated) electrons) imparted to the fluid. Without being bound by theory or mechanism, certain embodiments of the present invention relate to a oxygen-enriched fluid (output material) comprising charges (e.g., hydrated (solvated) electrons) that are added to the materials as the first material is mixed with oxygen in the inventive mixer device to provide the combined output material. According to particular aspects, these hydrated (solvated) electrons (alternately referred to herein as 'solvated electrons') are stabilized in the inventive solutions as evidenced by the persistence of assayable effects mediated by these hydrated (solvated) electrons. Certain embodiments may relate to hydrated (solvated) electrons and/or water-electron structures, clusters, etc., (See, for example, Lee and Lee, *Bull. Kor. Chem. Soc.* 2003, v. 24, 6; 802-804; 2003).

Horseradish peroxidase (HRP) effects. Horseradish peroxidase (HRP) is isolated from horseradish roots (*Amoracia rusticana*) and belongs to the ferroporphyrin group (Heme group) of peroxidases. HRP readily combines with hydrogen peroxide or other hydrogen donors to oxidize the pyrogallol substrate. Additionally, as recognized in the art, HRP facilitates auto-oxidative degradation of indole-3-acetic acid in the absence of hydrogen peroxide (see, e.g., Heme Peroxidases, H. Brian Dunford, Wiley-VCH, 1999, Chapter 6, pages 112-123, describing that auto-oxidation involves a highly efficient branched-chain mechanism; incorporated herein by reference in its entirety). The HRP reaction can be measured in enzymatic activity units, in which Specific activity is expressed in terms of pyrogallol units. One pyrogallol unit will form 1.0 mg purpurogallin from pyrogallol in 20 sec at pH 6.0 at 20°C. This purpurogallin (20 sec) unit is equivalent to approx. 18 µM units per min at 25°C.



It is known that Horseradish peroxidase enzyme catalyzes the auto-oxidation of pyrogallol by way of facilitating reaction with the molecular oxygen in a fluid. (Khajepour et al., *PROTEINS: Struct, Funct, Genet.* 53: 656-666 (2003)). It is also

known that oxygen binds the *heme* pocket of horseradish peroxidase enzyme through a hydrophobic pore region of the enzyme (between Phe68 and Phe142), whose conformation likely determines the accessibility of oxygen to the interior. According to particular aspects, and without being bound by mechanism, because surface charges
5 on proteins are known in the protein art to influence protein structure, the solvated electrons present in the inventive gas-enriched fluid may act to alter the conformation of the Horseradish peroxidase such that greater oxygen accessibility may result. The greater accessibility of oxygen to the prosthetic heme pocket of the Horseradish peroxidase enzyme may in turn allow for increased HRP reactivity, when compared
10 with prior art oxygenated fluids (pressure-pot, fine-bubbled).

In any event, according to particular aspects, production of output material using the inventive methods and devices comprises a process involving: an interfacial double layer that provides a charge gradient and movement of the materials relative to surfaces pulling charge (e.g., electrons) away from the surface by virtue of a
15 triboelectric effect, wherein the flow of material produces a flow of solvated electrons. Moreover, according to additional aspects, and without being bound by mechanism, the orbital structure of diatomic oxygen creates charge imbalances (e.g., the two unpaired electrons affecting the hydrogen bonding of the water) in the hydrogen bonding arrangement within the fluid material (water), wherein electrons are solvated
20 and stabilized within the imbalances.

Several chemical tests of the inventive oxygen-enriched fluid for the presence of hydrogen peroxide were conducted as described below, and none of these tests were positive (sensitivity of 0.1 ppm hydrogen peroxide). Thus, the inventive oxygen-enriched fluid of the instant application contain no, or less than 0.1 ppm, hydrogen
25 peroxide.

According to particular aspects, despite the absence of hydrogen peroxide, the inventive combination of oxygen-enrichment and solvated electrons imparted by the double-layer effects and configuration of the presently claimed devices may act to alter the conformation and/or heme group accessibility of the Horseradish peroxidase.

30

Glutathione Peroxidase Study

The inventive oxygen-enriched output fluid material was tested for the presence of hydrogen peroxide by testing the reactivity with glutathione peroxidase using a standard assay (Sigma). Briefly, glutathione peroxidase enzyme cocktail was constituted in deionized water and the appropriate buffers. Water samples were tested by adding the enzyme cocktail and inverting. Continuous spectrophotometric rate determination was made at A_{340} nm, and room temperature (25 degrees Celsius). Samples tested were: 1. deionized water (negative control), 2. inventive oxygen-enriched fluid at low concentration, 3. inventive oxygen-enriched fluid at high concentration, 4. hydrogen peroxide (positive control). The hydrogen peroxide positive control showed a strong reactivity, while none of the other fluids tested reacted with the glutathione.

Device for Generating Gas-Enriched Fluids or SolutionsDescription of the Related Art

Figure 1 provides a partial block diagram, partial cross-sectional view of a prior art device 10 for diffusing or emulsifying one or two gaseous or liquid materials ("infusion materials") into another gaseous or liquid material ("host material") reproduced from U.S. Patent No. 6,386,751, incorporated herein by reference in its entirety. The device 10 includes a housing configured to house a stator 30 and a rotor 12. The stator 30 encompasses the rotor 12. A tubular channel 32 is defined between the rotor 12 and the stator 30. The generally cylindrically-shaped rotor 12 has a diameter of about 7.500 inches and a length of about 6.000 inches providing a length-to-diameter ratio of about 0.8.

The rotor 12 includes a hollow cylinder, generally closed at both ends. A gap exists between each of the first and second ends of the rotor 12 and a portion of the housing 34. A rotating shaft 14 driven by a motor 18 is coupled to the second end of the rotor 12. The first end of the rotor 12 is coupled to an inlet 16. A first infusion material passes through the inlet 16 and into the interior of the rotor 12. The first infusion material passes from the interior of the rotor 12 and into the channel 32 through a plurality of openings 22 formed in the rotor 12.

The stator 30 also has openings 22 formed about its circumference. An inlet 36 passes a second infusion material to an area 35 between the stator 30 and the housing 34. The second infusion material passes out of the area 35 and into the channel 32 through openings 22.

5 An external pump (not shown) is used to pump the host material into a single inlet port 37. The host material passes through a single inlet port 37 and into the channel 32 where it encounters the first and second infusion materials, which enter the channel 32 through openings 22. The infusion materials may be pressurized at their source to prevent the host material from passing through openings 22.

10 The inlet port 37, is configured and positioned such that it is located along only a relatively small portion (< about 5%) of the annular inlet channel 32, and is substantially parallel to the axis of rotation of the rotor 12 to impart an axial flow toward a portion of the channel 32 into the host material.

Unfortunately, before entering the tubular channel 32, the host material must
15 travel in tortuous directions other than that of the axial flow (e.g., including in directions substantially orthogonal thereto) and down into and between the gap formed between the first end of the rotor 12 and the housing 34 (i.e., down a portion of the first end of the rotor adjacent to the inlet 16 between the end of the rotor 12 and the housing 34). The non-axial and orthogonal flow, and the presence of the host material in the gap
20 between the first end of the rotor 12 and the housing 34, causes undesirable and unnecessary friction. Further, it is possible for a portion of the host material to become trapped in eddy currents swirling between the first end of the rotor and the housing. Additionally, in the device 10, the host material must negotiate at least two right angles to enter any aspect of the annular of the annular inlet of the tubular channel 32.

25 A single outlet port 40 is formed in the housing 34. The combined host material and infusion material(s) exit the channel 32 via the outlet 40. The outlet port 40, which is also located along only a limited portion (< about 5%) of the annular outlet of tubular channel 32, is substantially parallel to the axis of rotation of the rotor 12 to impart or allow for an axial flow of the combined materials away from the limited portion of the
30 annular outlet of tubular channel 32 into the outlet port 40. An external pump 42 is used to pump the exiting fluid through the outlet port 40.

Unfortunately, before exiting the channel 32, a substantial portion of the exiting material must travel in a tortuous direction other than that of the axial flow (e.g., including in directions substantially orthogonal thereto) and down into and between the gap formed between the second end of the rotor 12 and the housing 34 (i.e., down a portion of the second end of the rotor adjacent to the shaft 14 between the end of the rotor 12 and the housing 34). As mentioned above, the non-axial and orthogonal flow, and the presence of the host material in the other gap between the end (in this case, the second end) of the rotor 12 and the housing 34, causes additional undesirable and unnecessary friction. Further, it is possible for a portion of the host material to become trapped in eddy currents swirling between the second end of the rotor and the housing. Additionally, in the device 10, a substantial portion of the exiting combined material must negotiate at least two right angles as it exits from the annular exit of the tubular channel 32 into the outlet port 40.

As is apparent to those of ordinary skill in the art, the inlet port 37 imparts only an axial flow to the host material. Only the rotor 21 imparts a circumferential flow into the host material. Further, the outlet port 40 imparts or provides for only an axial flow into the exiting material. Additionally, the circumferential flow velocity vector is imparted to the material only after it enters the annular inlet 37 of the tubular channel 32, and subsequently the circumferential flow vector must be degraded or eliminated as the material enters the exit port 40. There is, therefore, a need for a progressive circumferential acceleration of the material as it passes in the axial direction through the channel 32, and a circumferential deceleration upon exit of the material from the channel 32. These aspects, in combination with the tortuous path that the material takes from the inlet port 37 to the outlet port 40, create a substantial friction and flow resistance over the path that is accompanied by a substantial pressure differential (26 psi, at 60 gallons/min flow rate) between the inlet 37 and outlet 40 ports, and these factors, *inter alia*, combine to reduce the overall efficiency of the system.

Electrokinetically Oxygen-Enriched Aqueous Fluids and Solutions

Figure 2 provides a block diagram illustrating some of the components of a mixing device 100 and the flow of material into, within, and out of the device. The mixing device 100 combines two or more input materials to form an output material 102,

which may be received therefrom into a storage vessel 104. The mixing device 100 agitates the two or more input materials in a novel manner to produce an output material 102 having novel characteristics. The output material 102 may include not only a suspension of at least one of the input materials in at least one of the other input materials (e.g., emulsions) but also a novel combination (e.g., electrostatic combinations) of the input materials, a chemical compound resulting from chemical reactions between the input materials, combinations having novel electrostatic characteristics, and combinations thereof.

The input materials may include a first material 110 provided by a source 112 of the first material, a second material 120 provided by a source 122 of the second material, and optionally a third material 130 provided by a source 132 of the third material. The first material 110 may include a liquid, such as water, saline solution, chemical suspensions, polar liquids, non-polar liquids, colloidal suspensions, cell growing media, and the like. In some embodiments, the first material 110 may include the output material 102 cycled back into the mixing device 100. The second material 120 may consist of or include a gas, such as oxygen, nitrogen, carbon dioxide, carbon monoxide, ozone, sulfur gas, nitrous oxide, nitric oxide, argon, helium, bromine, and combinations thereof, and the like. In preferred embodiments, the gas is or comprises oxygen. The optional third material 130 may include either a liquid or a gas. In some embodiments, the third material 130 may be or include the output material 102 cycled back into the mixing device 100 (e.g., to one or more of the pumps 210, 220, or 230, and/or into the chamber 310 and/or 330).

Optionally, the first material 110, the second material 120, and the optional third material 130 may be pumped into the mixing device 100 by an external pump 210, an external pump 220, and an external pump 230, respectively. Alternatively, one or more of the first material 110, the second material 120, and the optional third material 130 may be stored under pressure in the source 112, the source 122, and the source 132, respectively, and may be forced into the mixing device 100 by the pressure. The invention is not limited by the method used to transfer the first material 110, the second material 120, and optionally, the third material 130 into the mixing device 100 from the source 112, the source 122, and the source 132, respectively.

The mixing device 100 includes a first chamber 310 and a second chamber 320 flanking a mixing chamber 330. The three chambers 310, 320, and 330 are interconnected and form a continuous volume.

The first material 110 is transferred into the first chamber 310 and flows therefrom into the mixing chamber 330. The first material 110 in the first chamber 310 may be pumped into the first chamber 310 by an internal pump 410. The second material 120 is transferred into the mixing chamber 330. Optionally, the third material 130 may be transferred into the mixing chamber 330. The materials in the mixing chamber 330 are mixed therein to form the output material 102. Then, the output material 102 flows into the second chamber 320 from which the output material 102 exits the mixing device 100. The output material 102 in the mixing chamber 330 may be pumped into the second chamber 320 by an internal pump 420. Optionally, the output material 102 in the second chamber 320 may be pumped therefrom into the storage vessel 104 by an external pump 430 (e.g., alone or in combination with the internal pump 410 and/or 420).

In particular aspects, a common drive shaft 500 powers both the internal pump 410 and the internal pump 420. The drive shaft 500 passes through the mixing chamber 330 and provides rotational force therein that is used to mix the first material 110, the second material 120, and optionally, the third material 130, together. The drive shaft 500 is powered by a motor 510 coupled thereto.

Figure 3 provides a system 512 for supplying the first material 110 to the mixing device 100 and removing the output material 102 from the mixing device 100. In the system 512, the storage vessel 104 of the output material 102 and the source 112 of the first material 110 are combined. The external pump 210 is coupled to the combined storage vessel 104 and source 112 by a fluid conduit 514 such as hose, pipe, and the like. The external pump 210 pumps the combined first material 110 and output material 102 from the combined storage vessel 104 and source 112 through the fluid conduit 514 and into a fluid conduit 516 connecting the external pump 210 to the mixing device 100. The output material 102 exits the mixing device 100 through a fluid conduit 518. The fluid conduit 518 is coupled to the combined storage vessel 104 and source 112 and transports the output material 102 exiting the mixing device 100 to the combined storage vessel 104 and source 112. The fluid conduit 518 includes a

valve 519 that establishes an operating pressure or back pressure within the mixing device 100.

Referring to Figures 2, 4-9, and 11, a more detailed description of various components of an embodiment of the mixing device 100 will be provided. The mixing device 100 is scalable. Therefore, dimensions provided with respect to various components may be used to construct an embodiment of the device or may be scaled to construct a mixing device of a selected size.

Turning to Figure 4, the mixing device 100 includes a housing 520 that houses each of the first chamber 310, the mixing chamber 330, and the second chamber 320. As mentioned above, the mixing device 100 includes the drive shaft 500, which rotates during operation of the device. Therefore, the mixing device 100 may vibrate or otherwise move. Optionally, the mixing device 100 may be coupled to a base 106, which may be affixed to a surface such as the floor to maintain the mixing device 100 in a substantially stationary position.

The housing 520 may be assembled from two or more housing sections. By way of example, the housing 520 may include a central section 522 flanked by a first mechanical seal housing 524 and a second mechanical seal housing 526. A bearing housing 530 may be coupled to the first mechanical seal housing 524 opposite the central section 522. A bearing housing 532 may be coupled to the second mechanical seal housing 526 opposite the central section 522. Optionally, a housing section 550 may be coupled to the bearing housings 530.

Each of the bearing housings 530 and 532 may house a bearing assembly 540 (see Figures 5 and 6). The bearing assembly 540 may include any suitable bearing assembly known in the art including a model number "202SZZST" manufactured by SKF USA Inc, of Kulpville, Pennsylvania, operating a website at www.skf.com.

Seals may be provided between adjacent housing sections. For example, o-ring 560 (see Figure 5) may be disposed between the housing section 550 and the bearing housing 530, o-ring 562 (see Figure 5) may be disposed between the first mechanical seal housing 524 and the central section 522, and o-ring 564 (see Figure 6) may be disposed between the second mechanical seal housing 526 and the central section 522.

MIXING CHAMBER 330

Turning now to Figure 7, the mixing chamber 330 is disposed inside the central section 522 of the housing 520 between the first mechanical seal housing 524 and the second mechanical seal housing 526. The mixing chamber 330 is formed between two components of the mixing device 100, a rotor 600 and a stator 700. The rotor 600 may have a sidewall 604 with an inside surface 605 defining a generally hollow inside portion 610 and an outside surface 606. The sidewall 604 may be about 0.20 inches to about 0.75 inches thick. In some embodiments, the sidewall 604 is about 0.25 inches thick. However, because the mixing device 100 may be scaled to suit a particular application, embodiments of the device having a sidewall 604 that is thicker or thinner than the values provided are within the scope of the present teachings. The sidewall 604 includes a first end portion 612 and a second end portion 614 and a plurality of through-holes 608 formed between the first end portion 612 and the second end portion 614. Optionally, the outside surface 606 of the sidewall 604 may include other features such as apertures, projections, textures, and the like. The first end portion 612 has a relieved portion 616 configured to receive a collar 618 and the second end portion 614 has a relieved portion 620 configured to receive a collar 622.

The rotor 600 is disposed inside the stator 700. The stator 700 has a sidewall 704 with an inside surface 705 defining a generally hollow inside portion 710 into which the rotor 600 is disposed. The sidewall 704 may be about 0.1 inches to about 0.3 inches thick. In some embodiments, the sidewall 604 is about 1.5 inches thick. The stator 700 may be non-rotatably coupled to the housing 520 in a substantially stationary position. Alternatively, the stator 700 may integrally formed with the housing 520. The sidewall 704 has a first end portion 712 and a second end portion 714. Optionally, a plurality of apertures 708 are formed in the sidewall 704 of the stator 700 between the first end portion 712 and the second end portion 714. Optionally, the inside surface 705 of the sidewall 704 may include other features such as through-holes, projections, textures, and the like.

The rotor 600 rotates with respect to the stationary stator 700 about an axis of rotation "a" in a direction indicated by arrow "C3" in Figure 9. Each of the rotor 600 and the stator 700 may be generally cylindrical in shape and have a longitudinal axis. The rotor 600 has an outer diameter "D1" and the stator 700 may have an inner

diameter "D2." The diameter "D1" may range, for example, from about 0.5 inches to about 24 inches. In some embodiments, the diameter "D1" is about 3.04 inches. In some embodiments, the diameter "D1" is about 1.7 inches. The diameter "D2," which is larger than the diameter "D1," may range from about 0.56 inches to about 24.25 inches.

5 In some embodiments, the diameter "D2" is about 4 inches. Therefore, the mixing chamber 330 may have a ring-shaped cross-sectional shape that is about 0.02 inches to about 0.125 inches thick (i.e., the difference between the diameter "D2" and the diameter "D1"). In particular embodiments, the mixing chamber 330 is about 0.025 inches thick. The channel 32 between the rotor 12 and the stator 34 of prior art
10 device 10 (see Figure 1) has a ring-shaped cross-sectional shape that is about 0.09 inches thick. Therefore, in particular embodiments, the thickness of the mixing chamber 330 is less than about one third of the channel 32 of the prior art device 10.

The longitudinal axis of the rotor 600 may be aligned with its axis of rotation "a." The longitudinal axis of the rotor 600 may be aligned with the longitudinal axis of the
15 stator 700. The rotor 600 may have a length of about 3 inches to about 6 inches along the axis of rotation "a." In some embodiments, the rotor 600 may have a length of about 5 inches along the axis of rotation "a." The stator 700 may have a length of about 3 inches to about 6 inches along the axis of rotation "a." In some embodiments, the stator 700 may have a length of about 5 inches along the axis of rotation "a."

20 While the rotor 600 and the stator 700 have been depicted as having a generally cylindrical shape, those of ordinary skill in the art appreciate that alternate shapes may be used. For example, the rotor 600 and the stator 700 may be conically, spherically, arbitrarily shaped, and the like. Further, the rotor 600 and the stator 700 need not be identically shaped. For example, the rotor 600 may be cylindrically shaped and the
25 stator 700 rectangular shaped or vice versa.

The apertures 708 of the stator 700 and the through-holes 608 depicted in Figures 4—7 are generally cylindrically shaped. The diameter of the through-holes 608 may range from about 0.1 inches to about 0.625 inches. The diameter of the apertures 708 may range from about 0.1 inches to about 0.625 inches. One or more of
30 apertures 708 of the stator 700 may have a diameter that differs from the diameters of the other apertures 708. For example, the apertures 708 may increase in diameter from the first end portion 712 of the stator 700 to the second end portion 714 of the

stator 700, the apertures 708 may decrease in diameter from the first end portion 712 of the stator 700 to the second end portion 714 of the stator 700, or the diameters of the apertures 708 may vary in another manner along the stator 700. One or more of through-holes 608 of the rotor 600 may have a diameter that differs from the diameters of the other through-holes 608. For example, the through-holes 608 may increase in diameter from the first end portion 612 of the rotor 600 to the second end portion 614 of the rotor 600, the through-holes 608 may decrease in diameter from the first end portion 612 of the rotor 600 to the second end portion 614 of the rotor 600, or the diameters of the through-holes 608 may vary in another manner along the rotor 600.

As described below with reference to alternate embodiments, the apertures 708 and the through-holes 608 may have shapes other than generally cylindrical and such embodiments are within the scope of the present invention. For example, the through-holes 608 may include a narrower portion, an arcuate portion, a tapered portion, and the like. Referring to Figures 7, each of the through-holes 608 includes an outer portion 608A, a narrow portion 608B, and a tapered portion 608C providing a transition between the outer portion 608A and the narrow portion 608B. Similarly, the apertures 708 may include a narrower portion, an arcuate portion, a tapered portion, and the like.

Figure 8 provides a non-limiting example of a suitable arrangement of the apertures 708 of the stator 700 and the through-holes 608 of the rotor 600. The apertures 708 of the stator 700 may be arranged in substantially parallel lateral rows "SLAT-1" through "SLAT-6" substantially orthogonal to the axis of rotation "a." The apertures 708 of the stator 700 may also be arranged in substantially parallel longitudinal rows "SLONG-1" through "SLONG-7" substantially parallel with the axis of rotation "a." In other words, the apertures 708 of the stator 700 may be arranged in a grid-like pattern of orthogonal rows (i.e., the lateral rows are orthogonal to the longitudinal rows) having the longitudinal rows "SLONG-1" through "SLONG-7" substantially parallel with the axis of rotation "a."

Like the apertures 708 of the stator 700, the through-holes 608 of the rotor 600 may be arranged in substantially parallel lateral rows "RLAT-1" through "RLAT-6" substantially orthogonal to the axis of rotation "a." However, instead of being arranged in a grid-like pattern of orthogonal rows, the through-holes 608 of the rotor 600 may

also be arranged in substantially parallel rows "RLONG-1" through "RLONG-7" that extend longitudinally along a helical path. Alternatively, the through-holes 608 of the rotor 600 may also be arranged in substantially parallel rows "RLONG-1" through "RLONG-7" that extend longitudinally at an angle other than parallel with the axis of rotation "a."

The apertures 708 of the stator 700 and the through-holes 608 of the rotor 600 may be configured so that when the rotor 600 is disposed inside the stator 700 the lateral rows "SLAT-1" to "SLAT-6" at least partially align with the lateral rows "RLAT-1" to "RLAT-6," respectively. In this manner, as the rotor 600 rotates inside the stator 700, the through-holes 608 pass by the apertures 708.

The through-holes 608 in each of the lateral rows "RLAT-1" to "RLAT-6" may be spaced apart laterally such that all of the through-holes 608 in the lateral row align, at least partially, with the apertures 708 in a corresponding one of the lateral rows "SLAT-1" to "SLAT-6" of the stator 700 at the same time. The longitudinally extending rows "RLONG-1" through "RLONG-6" may be configured such that the through-holes 608 in the first lateral row "RLAT-1" in each of the longitudinally extending rows passes completely by the apertures 708 of the corresponding lateral row "SLAT-1" before the through-holes 608 in the last lateral row "RLAT-6" begin to partially align with the apertures 708 of the corresponding last lateral row "SLAT-6" of the stator 700.

While, in Figure 8, six lateral rows and six longitudinally extending rows have been illustrated with respect to the rotor 600 and six lateral rows and seven longitudinally extending rows have been illustrated with respect stator 700, it is apparent to those of ordinary skill in the art that alternate numbers of lateral rows and/or longitudinal rows may be used with respect to the rotor 600 and/or stator 700 without departing from the present teachings.

To ensure that only one pair of openings between corresponding lateral rows will be coincident at any one time, the number of apertures 708 in each of the lateral rows "SLAT-1" to "SLAT-6" on the stator 700 may differ by a predetermined number (e.g., one, two, and the like) the number of through-holes 608 in each of the corresponding lateral rows "RLAT-1" to "RLAT-6" on the rotor 600. Thus, for example, if lateral row "RLAT-1" has twenty through-holes 608 evenly spaced around the

circumference of rotor 600, the lateral row "SLAT-1" may have twenty apertures 708 evenly spaced around the circumference of stator 700.

Returning to Figure 7, the mixing chamber 330 has an open first end portion 332 and an open second end portion 334. The through-holes 608 formed in the sidewall 604 of the rotor 600 connect the inside portion 610 of the rotor 600 with the mixing chamber 330.

The rotor 600 is rotated inside the stator 700 by the drive shaft 500 aligned with the axis of rotation "a" of the rotor 600. The drive shaft 500 may be coupled to the first end portion 612 and the second end portion 614 of the rotor 600 and extend through its hollow inside portion 610. In other words, a portion 720 of the drive shaft 500 is disposed in the hollow inside portion 610 of the rotor 600.

The collar 618 is configured to receive a portion 721 of the drive shaft 500 disposed in the hollow inside portion 610 and the collar 622 is configured to receive a portion 722 of the drive shaft 500 disposed in the hollow inside portion 610.

The portion 721 has an outer diameter "D3" that may range from about 0.5 inches to about 2.5 inches. In some embodiments, the diameter "D3" is about 0.625 inches. The portion 722 has an outer diameter "D4" that may be substantially similar to the diameter "D3," although, this is not required. The diameter "D4" may range from about 0.375 inches to about 2.5 inches.

The rotor 600 may be non-rotationally affixed to the portion 721 and the portion 722 of the drive shaft 500 by the collar 618 and the collar 622, respectively. By way of example, each of the collars 618 and 622 may be installed inside relieved portions 616 and 620, respectively. Then, the combined rotor 600 and collars 618 and 622 may be heated to expand them. Next, the drive shaft 500 is inserted through the collars 618 and 622 and the assembly is allowed to cool. As the collars 618 and 622 shrink during cooling, they tighten around the portions 722A and 722B of the drive shaft 500, respectively, gripping it sufficiently tightly to prevent the drive shaft 500 from rotating relative to the rotor 600. The collar 618, which does not rotate with respect to either the portion 721 or the relieved portion 616, translates the rotation of the drive shaft 500 to the first end portion 612 of the rotor 600. The collar 622, which does not rotate with respect to either the portion 722 or the relieved portion 620,

translates the rotation of the drive shaft 500 to the second end portion 614 of the rotor 600. The drive shaft 500 and the rotor 600 rotate together as a single unit.

5 The drive shaft 500 may have a first end portion 724 (see Figure 5) and a second end portion 726 (see Figure 6). The first end portion 724 may have a diameter "D5" of about 0.5 inches to about 1.75 inches. In particular embodiments, the diameter "D5" may be about 1.25 inches. The second end portion 726 may have a diameter "D6" that may be substantially similar to diameter "D5."

10 The second material 120 may be transported into the mixing chamber 330 through one of the first end portion 724 and the second end portion 726 of the rotating drive shaft 500. The other of the first end portion 724 and the second end portion 726 of the drive shaft 500 may be coupled to the motor 510. In the embodiment depicted in Figures 5 and 6, the second material 120 is transported into the mixing chamber 330 through the first end portion 724 and the second end portion 726 of the drive shaft 500 is coupled to the motor 510.

15 Turning to Figure 5, the drive shaft 500 may have a channel 728 formed therein that extends from the first end portion 724 into the portion 720 disposed in the inside portion 610 of the rotor 600. The channel 728 has an opening 730 formed in the first end portion 724. When the mixing device 100 is operating, the second material 120 is introduced into the channel 728 through the opening 730.

20 A valve 732 may be disposed inside a portion of the channel 728 located in the first end portion 724 of the drive shaft 500. The valve 732 may restrict or otherwise control the backward flow of the second material 120 from inside the hollow inside portion 610 through the channel 728 and/or the forward flow of the second material 120 into the channel 728. The valve 732 may include any valve known in the art including a
25 check valve. A suitable check valve includes a part number "CKFA1876205A," free flow forward check valve, manufactured by The Lee Company USA having an office in Bothell, WA, and operating a website at www.theleeco.com.

30 The drive shaft 500 may include an aperture 740 located in the inside portion 610 of the rotor 600 that connects the channel 728 with the inside portion 610 of the rotor 600. While only a single aperture 740 is illustrated in Figure 5, it is apparent to those of ordinary skill in the art that multiple apertures may be used to connect the channel 728 with the inside portion 610 of the rotor 600.

Referring to Figure 2, optionally, the external pump 220 may pump the second material 120 into the mixing device 100. By way of non-limiting example, the pump 220 may include any suitable pump known in the art including a diaphragm pump, a chemical pump, a peristaltic pump, a gravity fed pump, a piston pump, a gear pump, a combination of any of the aforementioned pumps, and the like. If the second material 120 is a gas, the gas may be pressurized and forced into the opening 730 formed in the first end portion 724 of the drive shaft 500 by releasing the gas from the source 122.

The pump 220 or the source 122 is coupled to the channel 728 by the valve 732. The second material 120 transported inside the channel 728 exits the channel 728 into the inside portion 610 of the rotor 600 through the aperture 740. The second material 120 subsequently exits the inside portion 610 of the rotor 600 through the through-holes 608 formed in the sidewall 608 of the rotor 600.

Referring to Figure 5, the mixing device 100 may include a seal assembly 750 coupled to the first end portion 724 of the drive shaft 500. The seal assembly 750 is maintained within a chamber 752 defined in the housing 520. The chamber 752 has a first end portion 754 spaced across the chamber from a second end portion 756. The chamber 752 also includes an input port 758 and an output port 759 that provide access into the chamber 752. The chamber 752 may be defined by housing section 550 and the bearing housing 530. The first end portion 754 may be formed in the housing section 550 and the second end portion 756 may be adjacent to the bearing housing 530. The input port 758 may be formed in the bearing housing 530 and the output port 759 may be formed in the housing section 550.

The seal assembly 750 includes a first stationary seal 760 installed in the first end portion 754 of the chamber 752 in the housing section 550 and the bearing housing 530. The first stationary seal 760 extends around a portion 762 of the first end portion 724 of the drive shaft 500. The seal assembly 750 also includes a second stationary seal 766 installed in the second end portion 756 of the chamber 752 in the bearing housing 530. The second stationary seal 766 extends around a portion 768 of the first end portion 724 of the drive shaft 500.

The seal assembly 750 includes a rotating assembly 770 that is non-rotatably coupled to the first end portion 724 of the drive shaft 500 between the portion 762 and

the portion 768. The rotating assembly 770 rotates therewith as a unit. The rotating assembly 770 includes a first seal 772 opposite a second seal 774. A biasing member 776 (e.g., a spring) is located between the first seal 772 and the second seal 774. The biasing member 776 biases the first seal 772 against the first stationary seal 760 and biases the second seal 774 against the second stationary seal 766.

A cooling lubricant is supplied to the chamber 752 and around rotating assembly 770. The lubricant enters the chamber 752 through the input port 758 and exits the chamber 752 through output port 759. The lubricant may lubricate the bearing assembly 540 housed by the bearing housing 530. A chamber 570 may be disposed between the bearing housing 530 and the mechanical seal housing 524. The bearing housing 530 may also include a second input port 759 connected to the chamber 570 into which lubricant may be pumped. Lubricant pumped into the chamber 570 may lubricate the bearing assembly 540. The seal assembly 750 may significantly, if not greatly, reduce frictional forces within this portion of the device caused by the rotation of the rotor 600 and may increase the active life of the seals 770. The seals may include surfaces constructed using silicon carbide.

Referring to Figure 9, as the rotor 600 rotates about the axis of rotation "a" in the direction indicated by arrow "C1," the rotor expels the second material 120 into the mixing chamber 330. The expelled bubbles, droplets, particles, and the like of the second material 120 exit the rotor 600 and are imparted with a circumferential velocity (in a direction indicated by arrow "C3") by the rotor 600. The second material 120 may be forced from the mixing chamber 330 by the pump 220 (see Figure 2), the centrifugal force of the rotating rotor 600, buoyancy of the second material 120 relative to the first material 110, and a combination thereof.

MOTOR 510

Returning to Figure 6, the second end portion 726 of the drive shaft 500 may be coupled to a rotating spindle 780 of a motor 510 by a coupler 900. The spindle 780 may have a generally circular cross-sectional shape with a diameter "D7" of about 0.25 inches to about 2.5 inches. In particular embodiments, the diameter "D7" may be about 0.25 inches to about 1.5 inches. While in the embodiment depicted in Figure 6, the diameter "D5" of the first end portion 724 of the drive shaft 500 is substantially equal to

the diameter "D7" and the spindle 780, embodiments in which one of the diameter "D5" and the diameter "D7" is larger than the other are within the scope of the present invention.

Referring also to Figure 4, it may be desirable to cover or shield the coupler 900.

5 In the embodiment illustrated in Figures 4 and 6, a drive guard 910 covers the coupler 900. The drive guard 910 may be generally U-shaped having a curved portion 914 flanked by a pair of substantially linear portions 915 and 916. The distal end of each of the substantially linear portions 915 and 916 of the drive guard 910 may have a flange 918 and 919, respectively. The drive guard 910 may be fastened by
10 each of its flanges 918 and 919 to the base 106.

The motor 510 may be supported on the base 106 by a support member 920. The support member 920 may be coupled to the motor 510 near the spindle 780. In the embodiment depicted, the support member 920 includes a through-hole through which the spindle 780 passes. The support member 920 may be coupled to the motor 510
15 using any method known in the art, including bolting the support member 920 to the motor 510 with one or more bolts 940.

The coupler 900 may include any coupler suitable for transmitting a sufficient amount of torque from the spindle 780 to the drive shaft 500 to rotate the rotor 600 inside to the stator 700. In the embodiment illustrated in Figures 4 and 6, the
20 coupler 900 is a bellows coupler. A bellows coupler may be beneficial if the spindle 780 and the drive shaft 500 are misaligned. Further, the bellows coupler may help absorb axial forces exerted on the drive shaft 500 that would otherwise be translated to the spindle 780. A suitable bellows coupler includes a model "BC32-8-8-A," manufactured by Ruland Manufacturing Company, Inc. of Marlborough, MA, which operates a website
25 at www.ruland.com.

The motor 510 may rotate the rotor 600 at about 0.1 revolutions per minute ("rpm") to about 7200 rpm. The motor 510 may include any motor suitable for rotating the rotor 600 inside to the stator 700 in accordance with the present teachings. By way of non-limiting example, a suitable motor may include a one-half horsepower electric
30 motor, operating at 230/460 volts and 3450 rpm. A suitable motor includes a model "C4T34NC4C" manufactured by LEESON Electric Corporation of Grafton, WI, which operates a website at www.leeson.com.

FIRST CHAMBER 310

Turning to Figures 4 and 7, the first chamber 310 is disposed inside the central section 522 of the housing 520 between the first mechanical seal housing 524 and the first end portions 612 and 712 of the rotor 600 and the stator 700, respectively. The first chamber 310 may be annular and have a substantially circular cross-sectional shape. The first chamber 310 and the mixing chamber 330 form a continuous volume. A portion 1020 of the drive shaft 500 extends through the first chamber 310.

As may best be viewed in Figure 4, the first chamber 310 has an input port 1010 through which the first material 110 enters the mixing device 100. The first material 110 may be pumped inside the first chamber 310 by the external pump 210 (see Figure 2). The external pump 210 may include any pump known in the art for pumping the first material 110 at a sufficient rate to supply the first chamber 310.

The input port 1010 is oriented substantially orthogonally to the axis of rotation "a." Therefore, the first material 110 enters the first chamber 310 with a velocity tangential to the portion 1020 of the drive shaft 500 extending through the first chamber 310. The tangential direction of the flow of the first material 110 entering the first chamber 310 is identified by arrow "T1." In the embodiment depicted in Figures 4 and 7, the input port 1010 may be offset from the axis of rotation "a." As is apparent to those of ordinary skill in the art, the direction of the rotation of the drive shaft 500 (identified by arrow "C1" in Figure 9) has a tangential component. The input port 1010 is positioned so that the first material 110 enters the first chamber 310 traveling in substantially the same direction as the tangential component of the direction of rotation of the drive shaft 500.

The first material 110 enters the first chamber 310 and is deflected by the inside of the first chamber 310 about the portion 1020 of the drive shaft 500. In embodiments wherein the first chamber 310 has a substantially circular cross-sectional shape, the inside of the first chamber 310 may deflect the first material 110 in a substantially circular path (identified by arrow "C2" in Figure 9) about the portion 1020 of the drive shaft 500. In such an embodiment, the tangential velocity of the first material 110 may cause it to travel about the axis of rotation "a" at a circumferential velocity, determined at least in part by the tangential velocity.

Once inside the first chamber 310, the first material 110 may be pumped from the first chamber 310 into the mixing chamber 330 by the pump 410 residing inside the first chamber 310. In embodiments that include the external pump 210 (see Figure 2), the external pump 210 may be configured to pump the first material 110 into the first chamber 310 at a rate at least as high as a rate at which the pump 410 pumps the first material 110 from the first chamber 310.

The first chamber 310 is in communication with the open first end portion 332 of the mixing chamber 330 and the first material 110 inside the first chamber 310 may flow freely into the open first end portion 332 of the mixing chamber 330. In this manner, the first material 110 does not negotiate any corners or bends between the mixing chamber 330 and the first chamber 310. In the embodiment depicted, the first chamber 310 is in communication with the entire open first end portion 332 of the mixing chamber 330. The first chamber 310 may be filled completely with the first material 110.

The pump 410 is powered by the portion 1020 of the drive shaft 500 extending through the first chamber 310. The pump 410 may include any pump known in the art having a rotating pump member 2022 housed inside a chamber (i.e., the first chamber 310) defined by a stationary housing (i.e., the housing 520). Non-limiting examples of suitable pumps include rotary positive displacement pumps such as progressive cavity pumps, single screw pumps (e.g., Archimedes screw pump), and the like.

The pump 410 depicted in Figures 7 and 9, is generally referred to as a single screw pump. In this embodiment, the pump member 2022 includes a collar portion 2030 disposed around the portion 1020 of the drive shaft 500. The collar portion 2030 rotates with the portion 1020 of the drive shaft 500 as a unit. The collar portion 2030 includes one or more fluid displacement members 2040. In the embodiment depicted in Figures 7 and 9, the collar portion 2030 includes a single fluid displacement member 2040 having a helical shape that circumscribes the collar portion 2030 along a helical path.

Referring to Figure 9, the inside of the first chamber 310 is illustrated. The pump 410 imparts an axial flow (identified by arrow "A1" and arrow "A2") in the first material 110 inside the first chamber 310 toward the open first end portion 332 of the

mixing chamber 330. The axial flow of the first material 110 imparted by the pump 410 has a pressure that may exceed the pressure obtainable by the external pump of the prior art device 10 (see Figure 1).

5 The pump 410 may also be configured to impart a circumferential flow (identified by arrow "C2") in the first material 110 as it travels toward the open first end portion 332 of the mixing chamber 330. The circumferential flow imparted in the first material 110 before it enters the mixing chamber 330 causes the first material 110 to enter the mixing chamber 330 already traveling in the desired direction at an initial circumferential velocity. In the prior art device 10 depicted in Figure 1, the first material 110 entered
10 the channel 32 of the prior art device 10 without a circumferential velocity. Therefore, the rotor 12 of the prior art device 10 alone had to impart a circumferential flow into the first material 110. Because the first material 110 is moving axially, in the prior art device 10, the first material 110 traversed at least a portion of the channel 32 formed between the rotor 12 and the stator 30 at a slower circumferential velocity than the first
15 material 110 traverses the mixing chamber 330 of the mixing device 100. In other words, if the axial velocity of the first material 110 is the same in both the prior art device 10 and the mixing device 100, the first material 110 may complete more revolutions around the rotational axis "a" before traversing the axial length of the mixing chamber 330, than it would complete before traversing the axial length of the
20 channel 32. The additional revolutions expose the first material 110 (and combined first material 110 and second material 120) to a substantially larger portion of the effective inside surface 706 (see Figure 7) of the stator 700.

In embodiments including the external pump 210 (see Figure 2), the circumferential velocity imparted by the external pump 210 combined with the input
25 port 1010 being oriented according to the present teachings, may alone sufficiently increase the revolutions of the first material 110 (and combined first material 110 and second material 120) about the rotational axis "a." Further, in some embodiments, the circumferential velocity imparted by the pump 210 and the circumferential velocity imparted by the pump 410 combine to achieve a sufficient number of revolutions of the
30 first material 110 (and combined first material 110 and second material 120) about the rotational axis "a." As is appreciated by those of ordinary skill in the art, other structural elements such as the cross-sectional shape of the first chamber 310 may contribute to

the circumferential velocity imparted by the pump 210, the pump 410, and a combination thereof.

In an alternate embodiment depicted in Figure 10, the pump 410 may include one or more vanes 2042 configured to impart a circumferential flow in the first material 110 as it travels toward the open first end portion 332 of the mixing chamber 330.

SECOND CHAMBER 320

Turning now to Figures 4 and 7, the second chamber 320 is disposed inside the central section 522 of the housing 520 between the second mechanical seal housing 526 and the second end portions 614 and 714 of the rotor 600 and the stator 700, respectively. The second chamber 320 may be substantially similar to the first chamber 310. However, instead of the input port 1010, the second chamber 320 may include an output port 3010. A portion 3020 of the drive shaft 500 extends through the second chamber 320.

The second chamber 320 and the mixing chamber 330 form a continuous volume. Further, the first chamber 310, the mixing chamber 330, and the second chamber 320 form a continuous volume. The first material 110 flows through the mixing device 100 from the first chamber 310 to the mixing chamber 330 and finally to the second chamber 320. While in the mixing chamber 330, the first material 110 is mixed with the second material 120 to form the output material 102. The output material 102 exits the mixing device 100 through the output port 3010. Optionally, the output material 102 may be returned to the input port 1010 and mixed with an additional quantity of the second material 120, the third material 130, or a combination thereof.

The output port 3010 is oriented substantially orthogonally to the axis of rotation "a" and may be located opposite the input port 1010 formed in the first chamber 310. The output material 102 enters the second chamber 320 from the mixing chamber 330 having a circumferential velocity (in the direction indicated by arrow "C3" in Figure 9) imparted thereto by the rotor 600. The circumferential velocity is tangential to the portion 3020 of the drive shaft 500 extending through the second chamber 320. In the embodiment depicted in Figures 4, 6, and 7, the output port 3010 may be offset from the axis of rotation "a." The output port 3010 is positioned so that the output

material 102, which enters the second chamber 320 traveling in substantially the same direction in which the drive shaft 500 is rotating (identified in Figure 9 by arrow "C1"), is traveling toward the output port 3010.

The output material 102 enters the second chamber 320 and is deflected by the inside of the second chamber 320 about the portion 3020 of the drive shaft 500. In embodiments wherein the second chamber 320 has a substantially circular cross-sectional shape, the inside of the second chamber 320 may deflect the output material 102 in a substantially circular path about the portion 3020 of the drive shaft 500.

Referring to Figure 2, optionally, the output material 102 may be pumped from inside the second chamber 320 by the external pump 430. The external pump 430 may include any pump known in the art for pumping the output material 102 at a sufficient rate to avoid limiting throughput of the mixing device 100. In such an embodiment, the external pump 430 may introduce a tangential velocity (in a direction indicated by arrow "T2" in Figures 4 and 11) to at least a portion of the output material 102 as the external pump 430 pumps the output material 102 from the second chamber 320. The tangential velocity of the portion of the output material 102 may cause it to travel about the axis of rotation "a" at a circumferential velocity, determined in part by the tangential velocity.

PUMP 420

Turning to Figures 6 and 7, the pump 420 residing inside the second chamber 320 may pump the output material 102 from the second chamber 320 into the output port 3010 and/or from the mixing chamber 330 into the second chamber 320. In embodiments that include the external pump 430, the external pump 430 may be configured to pump the output material 102 from the second chamber 320 at a rate at least as high as a rate at which the pump 420 pumps the output material 102 into the output port 3010.

The second chamber 320 is in communication with the open second end portion 334 of the mixing chamber 330 and the output material 102 inside the mixing chamber 330 may flow freely from the open second end portion 334 into the second chamber 320. In this manner, the output material 102 does not negotiate any corners

or bends between the mixing chamber 330 and the second chamber 320. In the embodiment depicted, the second chamber 320 is in communication with the entire open second end portion 334 of the mixing chamber 330. The second chamber 320 may be filled completely with the output material 102.

5 The pump 420 is powered by the portion 3020 of the drive shaft 500 extending through the second chamber 320. The pump 420 may be substantially identical to the pump 410. Any pump described above as suitable for use as the pump 410 may be used for the pump 420. While the pump 410 pumps the first material 110 into the mixing chamber 330, the pump 420 pumps the output material 102 from the mixing
10 chamber 330. Therefore, both the pump 410 and the pump 420 may be oriented to pump in the same direction.

As is appreciated by those of ordinary skill in the art, the first material 110 may differ from the output material 102. For example, one of the first material 110 and the output material 102 may be more viscous than the other. Therefore, the pump 410 may
15 differ from the pump 420. The pump 410 may be configured to accommodate the properties of the first material 110 and the pump 420 may be configured to accommodate the properties of the output material 102.

The pump 420 depicted in Figures 6 and 7, is generally referred to as a single screw pump. In this embodiment, the pump member 4022 includes a collar
20 portion 4030 disposed around the portion 3020 of the drive shaft 500. The collar portion 4030 rotates with the portion 3020 of the drive shaft 500 as a unit. The collar portion 4030 includes one or more fluid displacement members 4040. The collar portion 4030 includes a single fluid displacement member 4040 having a helical shape that circumscribes the collar portion 4030 along a helical path.

25 Referring to Figure 11, the inside of the second chamber 320 is illustrated. The pump 420 imparts an axial flow (identified by arrow "A3" and arrow "A4") in the output material 102 inside the second chamber 320 away from the open second end portion 334 of the mixing chamber 330.

The pump 420 may be configured to impart a circumferential flow (identified by
30 arrow "C4") in the output material 102 as it travels away from the open second end portion 334 of the mixing chamber 330. The circumferential flow imparted in the output

material 102 may help reduce an amount of work required by the rotor 600. The circumferential flow also directs the output material 102 toward the output port 3010.

In an alternate embodiment, the pump 420 may have substantially the same configuration of the pump 410 depicted in Figure 10. In such an embodiment, the one or more vanes 2042 are configured to impart a circumferential flow in the output material 102 as it travels away from the open second end portion 334 of the mixing chamber 330.

As is apparent to those of ordinary skill, various parameters of the mixing device 100 may be modified to obtain different mixing characteristics. Exemplary parameters that may be modified include the size of the through-holes 608, the shape of the through-holes 608, the arrangement of the through-holes 608, the number of through-holes 608, the size of the apertures 708, the shape of the apertures 708, the arrangement of the apertures 708, the number of apertures 708, the shape of the rotor 600, the shape of the stator 700, the width of the mixing chamber 330, the length of the mixing chamber 330, rotational speed of the drive shaft 500, the axial velocity imparted by the internal pump 410, the circumferential velocity imparted by the internal pump 410, the axial velocity imparted by the internal pump 420, the circumferential velocity imparted by the internal pump 420, the configuration of disturbances (e.g., texture, projections, recesses, apertures, and the like) formed on the outside surface 606 of the rotor 600, the configuration of disturbances (e.g., texture, projections, recesses, apertures, and the like) formed on the inside surface 706 of the stator 700, and the like.

ALTERNATE EMBODIMENT

Referring to Figure 12, a mixing device 5000 is depicted. The mixing device 5000 is an alternate embodiment of the mixing device 100. Identical reference numerals have been used herein to identify components of the mixing device 5000 that are substantially similar corresponding components of the mixing device 100. Only components of the mixing device 5000 that differ from the components of the mixing device 100 will be described.

The mixing device 5000 includes a housing 5500 for housing the rotor 600 and the stator 5700. The stator 5700 may be non-rotatably coupled by its first end

portion 5712 and its second end portion 5714 to the housing 5500. A chamber 5800 is defined between the housing 5500 and a portion 5820 of the stator 5700 flanked by the first end portion 5712 and the second end portion 5714. The housing 5500 includes an input port 5830 which provides access into the chamber 5800. The input port 5830
5 may be oriented substantially orthogonally to the axis of rotation "a;" however, this is not a requirement.

The stator 5700 includes a plurality of through-holes 5708 that connect the chamber 5800 and the mixing chamber 330 (defined between the rotor 600 and the stator 5700). An external pump 230 may be used to pump the third material 130 (which
10 may be identical to the second material 120) into the chamber 5800 via the input port 5830. The third material 130 pumped into the chamber 5800 may enter the mixing chamber 330 via the through-holes 5708 formed in the stator 5700. The third material 130 may be forced from the chamber 5800 by the pump 230, buoyancy of the third material 130 relative to the first material 110, and a combination thereof. As the
15 rotor 600 rotates, it may also draw the third material 130 from the chamber 5800 into the mixing chamber 330. The third material 130 may enter the mixing chamber 330 as bubbles, droplets, particles, and the like, which are imparted with a circumferential velocity by the rotor 600.

20 **ALTERNATE EMBODIMENT**

An alternate embodiment of the mixing device 100 may be constructed using a central section 5900 depicted in Figure 13 and a bearing housing 5920 depicted in Figure 14. Figure 13 depicts the central section 5900 having in its interior the stator 700 (see Figure 7). Identical reference numerals have been used herein to
25 identify components associated with the central section 5900 that are substantially similar corresponding components of the mixing device 100. Only components of the central section 5900 that differ from the components of the central section 522 will be described. The central section 5900 and the stator 700 are both constructed from a conductive material such as a metal (e.g., stainless steel). The input port 1010 and the
30 output port 3010 are both constructed from a nonconductive material such as plastic (e.g., PET, Teflon, nylon, PVC, polycarbonate, ABS, Delrin, polysulfone, etc.).

An electrical contact 5910 is coupled to the central section 5900 and configured to deliver a charge thereto. The central section 5900 conducts an electrical charge applied to the electrical contact 5910 to the stator 700. In further embodiments, the central section 5900 may be constructed from a nonconductive material. In such
5 embodiments, the electrical contact 5910 may pass through the central section 5900 and coupled to the stator 700. The electric charge applied by the electrical contact 5910 to the stator 700 may help facilitate redox or other chemical reactions inside the mixing chamber 330.

Optionally, insulation (not shown) may be disposed around the central
10 section 5900 to electrically isolate it from the environment. Further, insulation may be used between the central section 5900 and the first and second mechanical seals 524 and 526 that flank it to isolate it electrically from the other components of the mixing device.

Turning now to Figure 14, the bearing housing 5920 will be described. The
15 bearing housing 5920 is disposed circumferentially around the portion 726 of the drive shaft 500. An electrical contact 5922 is coupled to the bearing housing 5920. A rotating brush contact 5924 provides an electrical connection between the drive shaft 500 and the electrical contact 5922.

In this embodiment, the drive shaft 500 and the rotor 600 are both constructed
20 from a conductive material such as a metal (e.g., stainless steel). The bearing housing 5920 may be constructed from either a conductive or a nonconductive material. An electrical charge is applied to the drive shaft 500 by the electrical contact 5922 and the rotating brush contact 5924. The electrical charge is conducted by the drive shaft 500 to the rotor 600.

The alternate embodiment of the mixing device 100 constructed using the central
25 section 5900 depicted in Figure 13 and the bearing housing 5920 depicted in Figure 14 may be operated in at least two ways. First, the electrical contacts 5910 and 5922 may be configured not to provide an electrical charge to the stator 700 and the rotor 600, respectively. In other words, neither of the electrical contacts 5910 and 5922 are
30 connected to a current source, a voltage source, and the like.

Alternatively, the electrical contacts 5910 and 5922 may be configured to provide an electrical charge to the stator 700 and the rotor 600, respectively. For example, the

electrical contacts 5910 and 5922 may be coupled to a DC voltage source (not shown) supplying a steady or constant voltage across the electrical contacts 5910 and 5922. The negative terminal of the DC voltage source may be coupled to either of the electrical contacts 5910 and 5922 and the positive terminal of the DC voltage source
5 may be coupled to the other of the electrical contacts 5910 and 5922. The voltage supplied across the electrical contacts 5910 and 5922 may range from about 0.0001 volts to about 1000 volts. In particular embodiments, the voltage may range from about 1.8 volts to about 2.7 volts. By way of another example, a pulsed DC voltage having a duty cycle of between about 1% to about 99% may be used.

10 While the above examples of methods of operating the mixing device apply a DC voltage across the electrical contacts 5910 and 5922, as is apparent to those of ordinary skill in the art, a symmetrical AC voltage or non-symmetrical AC voltage having various shapes and magnitudes may be applied across the electrical contacts 5910 and 5922 and such embodiments are within the scope of the present invention.

15 MIXING INSIDE THE MIXING CHAMBER 330

As mentioned above, in the prior art device 10 (shown in Figure 1), the first material 110 entered the channel 32 between the rotor 12 and the stator 30 via a single limited input port 37 located along only a portion of the open second end of the
20 channel 32. Likewise, the output material 102 exited the channel 32 via a single limited output port 40 located along only a portion of the open first end of the channel 32. This arrangement caused undesirable and unnecessary friction. By replacing the single limited inlet port 37 and the single limited outlet port 40 with the chambers 310 and 320, respectively, friction has been reduced. Moreover, the first material 110 does not
25 negotiate a corner before entering the mixing chamber 330 and the output material 102 does not negotiate a corner before exiting the mixing chamber 330. Further, the chambers 310 and 320 provide for circumferential velocity of the material prior to entering, and after exiting the channel 32.

Accordingly, pressure drop across the mixing device 100 has been substantially
30 reduced. In the embodiments depicted in Figures 2, 4-9, and 11, the pressure drop between the input port 1010 and the output port 3010 is only approximately 12 psi when the mixing device 100 is configured to produce about 60 gallons of the output

material 102 per minute. This is an improvement over the prior art device 10 depicted in Figure 1, which when producing about 60 gallons of output material per minute was at least 26 psi. In other words, the pressure drop across the mixing device 100 is less than half that experienced by the prior art device 10.

5 According to additional aspects, the inclusion of pumps 410 and 420, which are powered by the drive shaft 500, provides a configuration that is substantially more efficient in mixing materials and that requires less energy than the external pumps used in the prior art.

10 MICRO-CAVITATION

During operation of the mixing device 100, the input materials may include the first material 110 (e.g., a fluid) and the second material 120 (e.g., a gas). The first material 110 and the second material 120 are mixed inside the mixing chamber 330 formed between the rotor 600 and the stator 700. Rotation of the rotor 600 inside the
15 stator 700 agitates the first material 110 and the second material 120 inside the mixing chamber 330. The through-holes 608 formed in the rotor 600 and/or the apertures 708 formed in the stator 700 impart turbulence in the flow of the first material 110 and the second material 120 inside the mixing chamber 330.

Without being limited by theory, the efficiency and persistence of the diffusion of
20 the second material 120 into the first material 110 is believed to be caused in part by micro-cavitation, which is described in connection with Figures 15-17. Whenever a material flows over a smooth surface, a rather laminar flow is established with a thin boundary layer that is stationary or moving very slowly because of the surface tension between the moving fluid and the stationary surface. The through-holes 608 and
25 optionally, the apertures 708, disrupt the laminar flow and can cause localized compression and decompression of the first material 110. If the pressure during the decompression cycle is low enough, voids (cavitation bubbles) will form in the material. The cavitation bubbles generate a rotary flow pattern 5990, like a tornado, because the localized area of low pressure draws the host material and the infusion material, as
30 shown in Figure 15. When the cavitation bubbles implode, extremely high pressures result. As two aligned openings (e.g., one of the apertures 708 and one of the through-holes 608) pass one another, a succussion (shock wave) occurs, generating significant

energy. The energy associated with cavitation and succussion mixes the first material 110 and the second material 120 together to an extremely high degree, perhaps at the molecular level.

The tangential velocity of the rotor 600 and the number of openings that pass each other per rotation may dictate the frequency at which the mixing device 100. It has been determined that operating the mixing device 100 within in the ultrasonic frequency range can be beneficial in many applications. It is believed that operating the mixing device 100 in the ultrasonic region of frequencies provides the maximum succession shock energy to shift the bonding angle of the fluid molecule, which enables it to transport an additional quantity of the second material 120 which it would not normally be able to retain. When the mixing device 100 is used as a diffuser, the frequency at which the mixing device 100 operates appears to affect the degree of diffusion, leading to much longer persistence of the second material 120 (infusion material) in the first material 110 (host material).

Referring now to Figure 15, an alternate embodiment of the rotor 600, rotor 6000 is provided. The cavitations created within the first material 110 in the mixing chamber 330 may be configured to occur at different frequencies along the length of the mixing chamber 330. The frequencies of the cavitations may be altered by altering the number and/or the placement of the through-holes 6608 along the length of the rotor 600. Each of the through-holes 6608 may be substantially similar to the through-holes 608 (discussed above).

By way of non-limiting example, the rotor 6000 may be subdivided into three separate exemplary sections 6100, 6200, and 6300. The through-holes 6608 increase in density from the section 6100 to the section 6200, the number of holes in the section 6100 being greater than the number of holes in the section 6200. The through-holes 6608 also increase in density from the section 6200 to the section 6300, the number of holes in the section 6200 being greater than the number of holes in the section 6300. Each of the sections 6100, 6200, and 6300 create succussions within their particular area at a different frequency due to the differing numbers of through-holes 6608 formed therein.

By manufacturing the rotor 6000 with a desired number of through-holes 6608 appropriately arranged in a particular area, the desired frequency of the succussions

within the mixing chamber 330 may be determined. Similarly, the desired frequency of the cavitations may be determined by a desired number of apertures 708 appropriately arranged in a particular area upon the stator 700 within which the rotor 600 rotates. Further, the desired frequency (or frequencies) of the succussions within the mixing chamber 330 may be achieved by selecting both a particular number and arrangement of the apertures 708 formed in the stator 700 and a particular number and arrangement of the through-holes 608 formed in the rotor 600.

Figures 19-21, depict various alternative arrangements of the apertures 708 formed in the stator 700 and the through-holes 608 formed in the rotor 600 configured to achieve different results with respect to the cavitations created. Figure 19 illustrates a configuration in which the apertures 708 and the through-holes 608 are aligned along an axis 7000 that is not parallel with any line (e.g., line 7010) drawn through the axis of rotation "a" of the rotor 600. In other words, if the rotor 600 has a cylindrical shape, the axis 7000 does not pass through the center of the rotor 600. Thus, the first material 110 within the mixing chamber 330 will not be oriented perpendicularly to the compressions and decompressions created by the apertures 708 and the through-holes 608. The compressions and decompressions will instead have a force vector that has at least a component parallel to the circumferential flow (in the direction of arrow "C3" of Figure 9) of first material 110 within the mixing chamber 330.

Relative alignment of the apertures 708 and the through-holes 608 may also affect the creation of cavitations in the mixing chamber 330. Figure 20 illustrates an embodiment in which the apertures 708 are in registration across the mixing chamber 330 with the through-holes 608. In this embodiment, rotation of the rotor 600 brings the through-holes 608 of the rotor into direct alignment with the apertures 708 of the stator 700. When in direct alignment with each other, the compressive and decompressive forces created by the apertures 708 and the through-holes 608 are directly aligned with one another.

In the embodiment depicted in Figure 21, the apertures 708 and the through-holes 608 are offset by an offset amount "X" along the axis of rotation "a." By way of non-limiting example, the offset amount "X" may be determined as a function of the size of the apertures 708. For example, the offset amount "X" may be approximately equal to one-half of the diameter of the apertures 708. Alternatively, the offset amount "X"

may be determined as a function of the size of the through-holes 608. For example, the offset amount "X" may be approximately equal to one-half of the diameter of the through-holes 608. If features (e.g., recesses, projections, etc.) other than or in addition to the through-holes 608 and the apertures 708 are included in either the rotor 600 or the stator 700, the offset amount "X" may be determined as a function of the size of such features. In this manner, the compressive and decompressive forces caused by the apertures 708 of the stator 700 and the through-holes 608 of the rotor 600 collide at a slight offset causing additional rotational and torsional forces within the mixing chamber 330. These additional forces increase the mixing (e.g., diffusive action) of the second material 120 into the first material 110 within the mixing chamber 330.

Referring now to Figures 22-25, non-limiting examples of suitable cross-sectional shapes for the apertures 708 and the through-holes 608 are provided. The cross-sectional shape of the apertures 708 and/or the through-holes 608 may be square as illustrated in Figure 22, circular as illustrated in Figure 23, and the like.

Various cross-sectional shapes of apertures 708 and/or the through-holes 608 may be used to alter flow of the first material 110 as the rotor 600 rotates within the stator 700. For example, Figure 24 depicts a teardrop cross-sectional shape having a narrow portion 7020 opposite a wide portion 7022. If the through-holes 608 have this teardrop shape, when the rotor 600 is rotated (in the direction generally indicated by the arrow "F"), the forces exerted on the first material 110, the second material 120, and optionally the third material 130 within the mixing chamber 330 increase as the materials pass from the wide portion 7022 of the teardrop to the narrow portion 7020.

Additional rotational forces can be introduced into the mixing chamber 330 by forming the apertures 708 and/or the through-holes 608 with a spiral configuration as illustrated in Figure 25. Material that flows into and out of the apertures 708 and/or the through-holes 608 having the spiral configuration experience a rotational force induced by the spiral configuration. The examples illustrated in Figures 22-25 are provided as non-limiting illustrations of alternate embodiments that may be employed within the mixing device 100. By application of ordinary skill in the art, the apertures 708 and/or the through-holes 608 may be configured in numerous ways to achieve various

successive and agitative forces appropriate for mixing materials within the mixing chamber 330.

DOUBLE LAYER EFFECT

5 The mixing device 100 may be configured to create the output material 102 by complex and non-linear fluid dynamic interaction of the first material 110 and the second material 120 with complex, dynamic turbulence providing complex mixing that further favors electrokinetic effects (described below). The result of these electrokinetic effects may be observed within the output material 102 as charge redistributions and
10 redox reactions, including in the form of solublized electrons that are stabilized within the output material.

 Ionization or dissociation of surface groups and/or adsorption of ions from a liquid cause most solid surfaces in contact with the liquid to become charged. Referring to Figure 26, an electrical double layer ("EDL") 7100 forms around exemplary
15 surface 7110 in contact with a liquid 7120. In the EDL 7100, ions 7122 of one charge (in this case, negatively charged ions) adsorb to the surface 7120 and form a surface layer 7124 typically referred to as a Stern layer. The surface layer 7124 attracts counterions 7126 (in this case, positively charged ions) of the opposite charge and equal magnitude, which form a counterion layer 7128 below the surface layer 7124
20 typically referred to as a diffuse layer. The counterion layer 7128 is more diffusely distributed than the surface layer 7124 and sits upon a uniform and equal distribution of both ions in the bulk material 7130 below. For OH⁻ and H⁺ ions in neutral water, the Gouy-Chapman model would suggest that the diffuse counterion layer extends about one micron into the water.

25 According to particular aspects, the electrokinetic effects mentioned above are caused by the movement of the liquid 7120 next to the charged surface 7110. Within the liquid 7120 (e.g., water, saline solution, and the like), the adsorbed ions 7122 forming the surface layer 7124 are fixed to the surface 7120 even when the liquid 7120 is in motion (for example, flowing in the direction indicated by arrow "G"); however, a
30 shearing plane 7132 exists within the diffuse counterion layer 7128 spaced from the surface 7120. Thus, as the liquid 7120 moves, some of the diffuse counterions 7126

are transported away from the surface 7120, while the absorbed ions 7122 remain at the surface 7120. This produces a so-called "streaming current."

Within the mixing chamber 330, the first material 110, the second material 120, and optionally, the third material 130 are subject to an electromagnetic field created by the inside surface 705 of the stator 700 and/or the outside surface 606 of the rotor 600, a voltage between the inside surface 705 and the outside surface 606, and/or an electrokinetic effect (e.g., streaming current) caused by at least one EDL formed in the first material 110. The at least one EDL may be introduced into the first material 110 by at least one of the inside surface 705 of the stator 700 and the outside surface 606 of the rotor 600.

Movement of the first material 110 through the mixing chamber 330 relative to surface disturbances (e.g., the through-holes 608 and apertures 708) creates cavitations in the first material 110 within the mixing chamber 330, which may diffuse the second material 120 into the first material 110. These cavitations may enhance contact between of the first material 110 and/or the second material 120 with the electric double layer formed on the inside surface 705 of the stator 700 and/or the electric double layer formed on the outside surface 606 of the rotor 600. Larger surface to volume ratios of the mixing chamber, an increased dwell time of the combined materials within the mixing chamber, and further in combination with a small average bubble size (and hence substantially greater bubble surface area) provide for effectively imparting EDL-mediated effects to the inventive output materials.

In embodiments in which the inside surface 705 and the outside surface 606 are constructed from a metallic material, such as stainless steel, the motion of the liquid 7120 and/or the streaming current(s) facilitate redox reactions involving H_2O , OH^- , H^+ , and O_2 at the inside surface 705 and the outside surface 606.

Referring to Figure 27, without being limited by theory, it is believed a section 7140 of the mixing chamber 330 between the inside surface 705 and the outside surface 606 may be modeled as a pair of parallel plates 7142 and 7144. If the first material 110 is a liquid, the first material 110 enters the section 7140 through an inlet "IN" and exits the section 7140 through an outlet "OUT." The inlet "IN" and the outlet "OUT" restrict the flow into and out of the section 7140.

Referring to Figure 28, the area between the parallel plates 7142 and 7144 has a high surface-area-to-volume ratio. Hence, a substantial portion of the counterion layer 7128 (and counterions 7126) may be in motion as the first material 110 moves between the plates 7142 and 7144. The number of counterions 7126 in motion may exceed the number allowed to enter the section 7140 by the inlet "IN" and the number allowed to exit the section 7140 by the outlet "OUT." The inlet "IN" and the outlet "OUT" feeding and removing the first material 110 from the section 7140, respectively, have far less surface area (and a lower surface-area-to-volume ratio) than the parallel plates 7142 and 7144 and thereby reduce the portion of the counterions 7126 in motion in the first material 110 entering and leaving the section 7140. Therefore, entry and exit from the section 7140 increases the streaming current locally. While a background streaming current (identified by arrow "BSC") caused by the flowing first material 110 over any surface is always present inside the mixing device 100, the plates 7142 and 7144 introduce an increased "excess" streaming current (identified by arrow "ESC") within the section 7140.

Without a conductive return current (identified by arrow "RC") in the plates 7142 and 7144 in the opposite direction of the flow of the first material 110, an excess charge 7146 having the same sign as the adsorbing ions 7122 would accumulate near the inlet "IN," and an excess charge 7148 having the same sign as the counterion 7126 would accumulate near the at outlet "OUT." Because such accumulated charges 7146 and 7148, being opposite and therefore attracted to one another, cannot build up indefinitely, the accumulated charges seek to join together by conductive means. If the plates 7142 and 7144 are perfectly electrically insulating, the accumulated charges 7146 and 7148 can relocate only through the first material 110 itself. When the conductive return current (identified by arrow "RC") is substantially equivalent to the excess streaming current (identified by arrow "ESC") in the section 7140, a steady state is achieved having zero net excess streaming current, and an electrostatic potential difference between the excess charge 7146 near the inlet "IN," and the excess charge 7148 near the outlet "OUT," creating a steady-state charge separation therebetween.

The amount of charge separation, and hence the electrostatic potential difference between the excess charge 7146 near the inlet "IN," and the excess

charge 7148 near the outlet "OUT," depends on additional energy per unit charge supplied by a pump (e.g., the rotor 600, the internal pump 410, and/or the external pump 210) to "push" the charge against the opposing electric field (created by the charge separation) to produce a liquid flow rate approximating a flow rate obtainable by
 5 a liquid without ions (i.e., ions 7122 and 7126). If the plates 7142 and 7144 are insulators, the electrostatic potential difference is a direct measure of the EMF the pump (e.g., the rotor 600, the internal pump 410, and/or the external pump 210) can generate. In this case, one could measure the electrostatic potential difference using a voltmeter having a pair of leads by placing one of the leads in the first material 110 near
 10 the inlet "IN," and the other lead in the first material 110 near the outlet "OUT."

With insulating plates 7142 and 7144, any return current is purely an ion current (or flow of ions), in that the return current involves only the conduction of ions through the first material 110. If other conductive mechanisms through more conductive pathways are present between the excess charge 7146 near the inlet "IN," and the
 15 excess charge 7148 near the outlet "OUT," the return current may use those more conductive pathways. For example, conducting metal plates 7142 and 7144 may provide more conductive pathways; however, these more conductive pathways transmit only an electron current and not the ion current.

As is appreciated by those of ordinary skill, to transfer the charge carried by an
 20 ion to one or more electrons in the metal, and vice versa, one or more oxidation-reduction reactions must occur at the surface of the metal, producing reaction products. Assuming the first material 110 is water (H₂O) and the second material 120 is oxygen (O₂), a non-limiting example of a redox reaction, which would inject a negative charge into the conducting plates 7142 and 7144 includes the following known half-cell
 25 reaction:



Again, assuming the first material 110 is water (H₂O) and the second material 120 is oxygen (O₂), a non-limiting example of a redox reaction includes the following known half-cell reaction, which would remove the negative charge from the conducting
 30 plates 7142 and 7144:



With conducting metal plates 7142 and 7144, most of the return current is believed to be an electron current, because the conducting plates 7142 and 7144 are more conductive than the first material 110 (provided the redox reactions are fast enough not to be a limiting factor). For the conducting metal plates 7142 and 7144, a smaller charge separation accumulates between the inlet "IN" and the outlet "OUT," and a much smaller electrostatic potential exists therebetween. However, this does not mean that the EMF is smaller.

As described above, the EMF is related to the energy per unit charge the pump provides to facilitate the flow of the first material 110 against the opposing electric field created by the charge separation. Because the electrostatic potential is smaller, the pump may supply a less energy per unit charge to cause the first material 110 to flow. However, the above example redox reactions do not necessarily occur spontaneously, and thus may require a work input, which may be provided by the pump. Therefore, a portion of the EMF (that is not reflected in the smaller electrostatic potential difference) may be used to provide the energy necessary to drive the redox reactions.

In other words, the same pressure differentials provided by the pump to push against the opposing electric field created by the charge separation for the insulating plates 7142 and 7144, may be used both to "push" the charge through the conducting plates 7142 and 7144 and drive the redox reactions.

Referring to Figure 29, an experimental setup for an experiment conducted by the inventors is provided. The experiment included a pair of substantially identical spaced apart 500 ml standard Erlenmeyer flasks 7150 and 7152, each containing a volume of deionized water 7153. A rubber stopper 7154 was inserted in the open end of each of the flasks 7150 and 7152. The stopper 7154 included three pathways, one each for a hollow tube 7156, a positive electrode 7158, and a negative electrode 7160. With respect to each of the flasks 7150 and 7152, each of the hollow tube 7156, the positive electrode 7158, and the negative electrode 7160 all extended from outside the flask, through the stopper 7154, and into the deionized water 7153 inside the flask. The positive electrode 7158 and the negative electrode 7160 were constructed from stainless steel. The hollow tubes 7156 in both of the flasks 7150 and 7152 had an open end portion 7162 coupled to a common oxygen supply 7164. The positive electrode 7158 and the negative electrode 7160 inserted into the flask 7152 where

coupled to a positive terminal and a negative terminal, respectively, of a DC power supply 7168. Exactly the same sparger was used in each flask.

Oxygen flowed through the hollow tubes 7156 into both of the flasks 7150 and 7152 at a flow rate (Feed) of about 1 SCFH to about 1.3 SCFH (combined flow rate). The voltage applied across the positive electrode 7158 and the negative electrode 7160 inserted into the flask 7152 was about 2.55 volts. This value was chosen because it is believed to be an electrochemical voltage value sufficient to affect all oxygen species. This voltage was applied continuously over three to four hours during which oxygen from the supply 7164 was bubbled into the deionized water 7153 in each of the flasks 7150 and 7152.

Testing of the deionized water 7153 in the flask 7150 with HRP and pyrogallol gave an HRP-mediated pyrogallol reaction activity consistent with the properties of fluids produced with the alternate rotor/stator embodiments described herein. The HRP optical density was about 20% higher relative to pressure-pot or fine-bubbled solutions of equivalent oxygen content. The results of this experiment indicate that mixing inside the mixing chamber 330 involves a redox reaction. According to particular aspects, the inventive mixing chambers provide for output materials comprising added electrons that are stabilized by either oxygen-rich water structure within the inventive output solutions, or by some form of oxygen species present due to the electrical effects within the process.

Additionally, the deionized water 7153 in both of the flasks 7150 and 7152 was tested for both ozone and hydrogen peroxide employing industry-standard colorimetric test ampoules with a sensitivity of 0.1 ppm for hydrogen peroxide and 0.6 ppm for ozone. There was no positive indication of either species up to the detection limits of those ampoules.

DWELL TIME

Dwell time is an amount of time the first material 110, the second material 120, and, optionally, the third material 130 spend in the mixing chamber 330. The ratio of the length of the mixing chamber 330 to the diameter of the mixing chamber 330 may significantly affect dwell time. The greater the ratio, the longer the dwell time. As mentioned in the Background Section, the rotor 12 of the prior art device 10 (see

Figure 1) had a diameter of about 7.500 inches and a length of about 6.000 inches providing a length to diameter ratio of about 0.8. In contrast, in particular embodiments, the length of the mixing chamber 330 of the mixing device 100 is about 5 inches and the diameter "D1" of the rotor 600 is about 1.69 inches yielding a length to diameter ratio of about 2.95.

Dwell time represents the amount of time that the first material 110, the second material 120, and optionally the third material 130 are able to interact with the electrokinetic phenomena described herein. The prior art device 10 is configured to produce about 60 gallons of the output material 102 per minute and the mixing device 100 is configured to produce about 0.5 gallons of the output material 102 per minute; the prior art device 10 (see Figure 1) had a fluid dwell time of about 0.05 seconds, whereas embodiments of the mixing device 100 have a substantially greater (about 7-times greater) dwell time of about 0.35 seconds. This longer dwell time allows the first material 110, the second material 120, and optionally the third material 130 to interact with each other and the surfaces 606 and 705 (see Figure 7) inside the mixing chamber 330 for about 7-times longer than was possible in the prior art device 10. In additional embodiments, the dwell time is at least 1.5-times, at least 2-times, at least 3-times, at least 4-times, at least 5-times, at least 6-times, at least 7-times or greater, than was possible in the prior art device 10.

With reference to Table 5 below, the above dwell times were calculated by first determining the flow rate for each device in gallons per second. In the case of the prior art device 10 was configured to operate at about 60 gallons of output material per minute, while the mixing device 100 is configured to operate over a broader range of flow rate, including at an optimal range of about 0.5 gallons of output material per minute. The flow rate was then converted to cubic inches per second by multiplying the flow rate in gallons per second by the number of cubic inches in a gallon (i.e., 231 cubic inches). Then, the volume (12.876 cubic inches) of the channel 32 of the prior art device 10 was divided by the flow rate of the device (231 cubic inches/second) to obtain the dwell time (in seconds) and the volume (0.673 cubic inches) of the mixing chamber 330 of the mixing device 100 was divided by the flow rate (1.925 cubic inches/second) of the device (in cubic inches per second) to obtain the dwell time (in seconds).

TABLE 5. Inventive device can accommodate a range of dwell times, including a substantially increased (e.g., 7-times) dwell time relative to prior art devices.

Device	Flow Rate Gallons/ Minute	Flow Rate Gallons/ Second	Flow Rate Cubic Inches/ Second	Volume Mixing Chamber (Cubic Inches)	Dwell Time (Seconds)
Prior art device 10	60	1.000	231.000	12.876	0.056
Mixing device 100	2	0.033	7.700	0.673	0.087
Mixing device 100	0.5	0.008	1.925	0.673	0.350

5

RATE OF INFUSION

Particular aspects of the mixing device 100 provide an improved oxygen infusion rate over the prior art, including over prior art device 10 (see Figure 1). When the first material 110 is water and the second material 120 is oxygen, both of which are processed by the mixing device 100 in a single pass (i.e., the return block of Figure 2 is set to "NO") at or near 20° Celsius, the output material 102 has a dissolved oxygen level of about 43.8 ppm. In certain aspects, an output material having about 43.8 ppm dissolved oxygen is created in about 350 milliseconds via the inventive flow through the inventive non-pressurized (non-pressure pot) methods. In contrast, when the first material 110 (water) and the second material 120 (oxygen) are both processed in a single pass at or near 20° Celsius by the prior art device 10, the output material had dissolved oxygen level of only 35 ppm in a single pass of 56 milliseconds.

OUTPUT MATERIAL 102

When the first material 110 is a liquid (e.g., freshwater, saline, GATORADE®, and the like) and the second material 120 is a gas (e.g., oxygen, nitrogen, and the like), the mixing device 100 may diffuse the second material 120 into the first material 110. The following discusses results of analyses performed on the output material 102 to characterize one or more properties of the output material 102 derived from having been processed by the mixing device 100.

When the first material 110 is saline solution and the second material 120 is oxygen gas, experiments have indicated that a vast majority of oxygen bubbles produced within the saline solution are no greater than 0.1 micron in size.

5 DECAF OF DISSOLVED OXYGEN LEVELS

Referring now to Figure 30, there is illustrated the DO levels in water processed with oxygen in the mixing device 100 and stored in a 500 ml thin-walled plastic bottle and a 1000 ml glass bottle. Each of the bottles was capped and stored at 65 degrees Fahrenheit. Point 7900 is the DO level at bottling. Line 7902 illustrates the Henry's
10 Law equilibrium state (i.e., the amount of dissolved oxygen that should be within the water at 65 degrees Fahrenheit), which is a DO level of slightly less than 10 ppm. Points 7904 and 7906 represent the DO levels within the water in the plastic bottle at 65 days and 95 days, respectively. As can be seen at point 7904, when the plastic bottle is opened approximately 65 days after bottling, the DO level within the water is
15 approximately 27.5 ppm. When the bottle is opened approximately 95 days after bottling, as indicated at point 7906, the DO level is approximately 25 ppm. Likewise, for the glass bottle, the DO level is approximately 40 ppm at 65 days as indicated at point 7908 and is approximately 41 ppm at 95 days as illustrated at point 7910. Thus, Figure 30 indicates the DO levels within both the plastic bottle and the glass bottle
20 remain relatively high at 65 degrees Fahrenheit.

Referring now to Figure 30, there is illustrated the DO levels in water enriched with oxygen in the mixing device 100 and stored in a 500 ml thin-walled plastic bottle and a 1000 ml glass bottle out to at least 365 days. Each of the bottles was capped and stored at 65 degrees Fahrenheit. As can be seen in the Figure, the DO levels of
25 the oxygen-enriched fluid remained fairly constant out to at least 365 days.

Referring to Figure 31, there is illustrated the DO levels in water enriched with oxygen in the mixing device 100 and stored in a 500 ml plastic thin-walled bottle and a 1000 ml glass bottle. Both bottles were refrigerated at 39 degrees Fahrenheit. Again, DO levels of the oxygen-enriched fluid remained steady and decreased only slightly out
30 to at least 365 days.

MOLECULAR INTERACTIONS

Conventionally, quantum properties are thought to belong to elementary particles of less than 10^{-10} meters, while the macroscopic world of our everyday life is referred to as classical, in that it behaves according to Newton's laws of motion.

5 Recently, molecules have been described as forming clusters that increase in size with dilution. These clusters measure several micrometers in diameter, and have been reported to increase in size non-linearly with dilution. Quantum coherent domains measuring 100 nanometers in diameter have been postulated to arise in pure water, and collective vibrations of water molecules in the coherent domain may eventually
10 become phase locked to electromagnetic field fluctuations, providing for stable oscillations in water, providing a form of 'memory' in the form of excitation of long lasting coherent oscillations specific to dissolved substances in the water that change the collective structure of the water, which may in turn determine the specific coherent oscillations that develop. Where these oscillations become stabilized by magnetic field
15 phase coupling, the water, upon dilution may still carry 'seed' coherent oscillations. As a cluster of molecules increases in size, its electromagnetic signature is correspondingly amplified, reinforcing the coherent oscillations carried by the water.

Despite variations in the cluster size of dissolved molecules and detailed microscopic structure of the water, a specificity of coherent oscillations may
20 nonetheless exist. One model for considering changes in properties of water is based on considerations involved in crystallization.

With reference to Figure 36, a simplified protonated water cluster forming a nanoscale cage 8700 is shown. A protonated water cluster typically takes the form of $H^+(H_2O)_n$. Some protonated water clusters occur naturally, such as in the ionosphere.
25 Without being bound by any particular theory, and according to particular aspects, other types of water clusters or structures (clusters, nanocages, etc) are possible, including structures comprising oxygen and stabilized electrons imparted to the inventive output materials. Oxygen atoms 8704 may be caught in the resulting structures 8700. The chemistry of the semi-bound nanocage allows the oxygen 8704 and/or stabilized
30 electrons to remain dissolved for extended periods of time. Other atoms or molecules, such as medicinal compounds, can be caged for sustained delivery purposes. The

specific chemistry of the solution material and dissolved compounds depend on the interactions of those materials.

Fluids processed by the mixing device 100 have been shown via experiments to exhibit different structural characteristics that are consistent with an analysis of the fluid in the context of a cluster structure.

Water processed through the mixing device 100 has been demonstrated to have detectable structural differences when compared with normal unprocessed water. For example, processed water has been shown to have more Rayleigh scattering than is observed in unprocessed water. In the experiments that were conducted, samples of processed and unprocessed water were prepared (by sealing each in a separate bottle), coded (for later identification of the processed sample and unprocessed sample), and sent to an independent testing laboratory for analysis. Only after the tests were completed were the codes interpreted to reveal which sample had been processed by the mixing device 100.

At the laboratory, the two samples were placed in a laser beam having a wavelength of 633 nanometers. The fluid had been sealed in glass bottles for approximately one week before testing. With respect to the processed sample, Sample B scattered light regardless of its position relative to the laser source. However, Sample A did not. After two to three hours following the opening of the bottle, the scattering effect of Sample B disappeared. These results imply the water exhibited a memory causing the water to retain its properties and dissipate over time. These results also imply the structure of the processed water is optically different from the structure of the unprocessed fluid. Finally, these results imply the optical effect is not directly related to DO levels because the DO level at the start was 45 ppm and at the end of the experiment was estimated to be approximately 32 ppm.

Charge-stabilized nanostructures (e.g., charge stabilized oxygen-containing nanostructures):

As described herein above under "Double Layer Effect," "Dwell Time," "Rate of Infusion," and "Bubble size Measurements," the mixing device 100 creates, in a matter of milliseconds, a unique non-linear fluid dynamic interaction of the first material 110 and the second material 120 with complex, dynamic turbulence providing complex mixing in contact with an effectively enormous surface area (including those of the

device and of the exceptionally small gas bubbles of less than 100 nm) that provides for the novel electrokinetic effects described herein. Additionally, feature-localized electrokinetic effects (voltage/current) were demonstrated herein (see working Example 20) using a specially designed mixing device comprising insulated rotor and stator features.

As well-recognized in the art, charge redistributions and/or solvated electrons are known to be highly unstable in aqueous solution. According to particular aspects, Applicants' electrokinetic effects (e.g., charge redistributions, including, in particular aspects, solvated electrons) are surprisingly stabilized within the output material (e.g., saline solutions, ionic solutions). In fact, as described herein, the stability of the properties and biological activity of the inventive electrokinetic fluids (e.g., RNS-60 or Solas) can be maintained for months in a gas-tight container, indicating involvement of dissolved gas (e.g., oxygen) in helping to generate and/or maintain, and/or mediate the properties and activities of the inventive solutions. Significantly, as described in the working Examples herein, the charge redistributions and/or solvated electrons are stably configured in the inventive electrokinetic ionic aqueous fluids in an amount sufficient to provide, upon contact with a living cell (e.g., mammalian cell) by the fluid, modulation of at least one of cellular membrane potential and cellular membrane conductivity (see, e.g., cellular patch clamp working Examples 23 and 24).

As described herein under "Molecular Interactions," to account for the stability and biological compatibility of the inventive electrokinetic fluids (e.g., electrokinetic saline solutions), Applicants have proposed that interactions between the water molecules and the molecules of the substances (e.g., oxygen) dissolved in the water change the collective structure of the water and provide for nanoscale cage clusters, including nanostructures comprising oxygen and/or stabilized electrons imparted to the inventive output materials. Without being bound by mechanism, and according to the properties and activities described herein, the configuration of the nanostructures in particular aspects is such that they: comprise (at least for formation and/or stability and/or biological activity) dissolved gas (e.g., oxygen); enable the electrokinetic fluids (e.g., RNS-60 or Solas saline fluids) to modulate (e.g., impart or receive) charges and/or charge effects upon contact with a cell membrane or related constituent thereof;

and in particular aspects provide for stabilization (e.g., carrying, harboring, trapping) solvated electrons in a biologically-relevant form.

According to particular aspects, and as supported by the present disclosure, in ionic or saline (e.g., standard saline, NaCl) solutions, the inventive nanostructures
5 comprise charge stabilized nanostructures (e.g., average diameter less than 100 nm) that may comprise at least one dissolved gas molecule (e.g., oxygen) within a charge-stabilized hydration shell. According to additional aspects, and as described elsewhere herein, the charge-stabilized hydration shell may comprise a cage or void harboring the at least one dissolved gas molecule (e.g., oxygen). According to further aspects, by
10 virtue of the provision of suitable charge-stabilized hydration shells, the charge-stabilized nanostructure and/or charge-stabilized oxygen containing nano-structures may additionally comprise a solvated electron (e.g., stabilized solvated electron).

Without being bound by mechanism or particular theory, after the present priority date, charge-stabilized microbubbles stabilized by ions in aqueous liquid in equilibrium
15 with ambient (atmospheric) gas have been proposed (Bunkin et al., *Journal of Experimental and Theoretical Physics*, 104:486-498, 2007; incorporated herein by reference in its entirety). According to particular aspects of the present invention, Applicants' novel electrokinetic fluids comprise a novel, biologically active form of charge-stabilized oxygen-containing nanostructures, and may further comprise novel
20 arrays, clusters or associations of such structures.

According to the charge-stabilized microbubble model, the short-range molecular order of the water structure is destroyed by the presence of a gas molecule (e.g., a dissolved gas molecule initially complexed with a nonadsorptive ion provides a short-range order defect), providing for condensation of ionic droplets, wherein the defect is
25 surrounded by first and second coordination spheres of water molecules, which are alternately filled by adsorptive ions (e.g., acquisition of a 'screening shell of Na⁺ ions to form an electrical double layer) and nonadsorptive ions (e.g., Cl⁻ ions occupying the second coordination sphere) occupying six and 12 vacancies, respectively, in the coordination spheres. In under-saturated ionic solutions (e.g., undersaturated saline
30 solutions), this hydrated 'nucleus' remains stable until the first and second spheres are filled by six adsorptive and five nonadsorptive ions, respectively, and then undergoes

Coulomb explosion creating an internal void containing the gas molecule, wherein the adsorptive ions (e.g., Na^+ ions) are adsorbed to the surface of the resulting void, while the nonadsorptive ions (or some portion thereof) diffuse into the solution (Bunkin et al., *supra*). In this model, the void in the nanostructure is prevented from collapsing by
5 Coulombic repulsion between the ions (e.g., Na^+ ions) adsorbed to its surface. The stability of the void-containing nanostructures is postulated to be due to the selective adsorption of dissolved ions with like charges onto the void/bubble surface and diffusive equilibrium between the dissolved gas and the gas inside the bubble, where the negative (outward electrostatic pressure exerted by the resulting electrical double layer
10 provides stable compensation for surface tension, and the gas pressure inside the bubble is balanced by the ambient pressure. According to the model, formation of such microbubbles requires an ionic component, and in certain aspects collision-mediated associations between particles may provide for formation of larger order clusters (arrays) (*Id*).

15 The charge-stabilized microbubble model suggests that the particles can be gas microbubbles, but contemplates only spontaneous formation of such structures in ionic solution in equilibrium with ambient air, is uncharacterized and silent as to whether oxygen is capable of forming such structures, and is likewise silent as to whether solvated electrons might be associated and/or stabilized by such structures.

20 According to particular aspects, the inventive electrokinetic fluids comprising charge-stabilized nanostructures and/or charge-stabilized oxygen-containing nanostructures are novel and fundamentally distinct from the postulated non-electrokinetic, atmospheric charge-stabilized microbubble structures according to the microbubble model. Significantly, this conclusion is in unavavoidable, deriving, at least in
25 part, from the fact that control saline solutions do not have the biological properties disclosed herein, whereas Applicants' charge-stabilized nanostructures provide a novel, biologically active form of charge-stabilized oxygen-containing nanostructures.

According to particular aspects of the present invention, Applicants' novel electrokinetic device and methods provide for novel electrokinetically-altered fluids
30 comprising significant quantities of charge-stabilized nanostructures in excess of any amount that may or may not spontaneously occur in ionic fluids in equilibrium with air,

or in any non-electrokinetically generated fluids. In particular aspects, the charge-stabilized nanostructures comprise charge-stabilized oxygen-containing nanostructures. In additional aspects, the charge-stabilized nanostructures are all, or substantially all charge-stabilized oxygen-containing nanostructures, or the charge-stabilized oxygen-
5 containing nanostructures the major charge-stabilized gas-containing nanostructure species in the electrokinetic fluid.

According to yet further aspects, the charge-stabilized nanostructures and/or the charge-stabilized oxygen-containing nanostructures may comprise or harbor a solvated electron, and thereby provide a novel stabilized solvated electron carrier. In particular
10 aspects, the charge-stabilized nanostructures and/or the charge-stabilized oxygen-containing nanostructures provide a novel type of electride (or inverted electride), which in contrast to conventional solute electrides having a single organically coordinated cation, rather have a plurality of cations stably arrayed about a void or a void containing an oxygen atom, wherein the arrayed sodium ions are coordinated by water hydration
15 shells, rather than by organic molecules. According to particular aspects, a solvated electron may be accommodated by the hydration shell of water molecules, or preferably accommodated within the nanostructure void distributed over all the cations. In certain aspects, the inventive nanostructures provide a novel 'super electride' structure in solution by not only providing for distribution/stabilization of the solvated electron over
20 multiple arrayed sodium cations, but also providing for association or partial association of the solvated electron with the caged oxygen molecule(s) in the void—the solvated electron distributing over an array of sodium atoms and at least one oxygen atom. According to particular aspects, therefore, 'solvated electrons' as presently disclosed in association with the inventive electrokinetic fluids, may not be solvated in the traditional
25 model comprising direct hydration by water molecules. Alternatively, in limited analogy with dried electride salts, solvated electrons in the inventive electrokinetic fluids may be distributed over multiple charge-stabilized nanostructures to provide a 'lattice glue' to stabilize higher order arrays in aqueous solution.

In particular aspects, the inventive charge-stabilized nanostructures and/or the
30 charge-stabilized oxygen-containing nanostructures are capable of interacting with cellular membranes or constituents thereof, or proteins, etc., to mediate biological

activities. In particular aspects, the inventive charge-stabilized nanostructures and/or the charge-stabilized oxygen-containing nanostructures harboring a solvated electron are capable of interacting with cellular membranes or constituents thereof, or proteins, etc., to mediate biological activities.

5 In particular aspects, the inventive charge-stabilized nanostructures and/or the charge-stabilized oxygen-containing nanostructures interact with cellular membranes or constituents thereof, or proteins, etc., as a charge and/or charge effect donor (delivery) and/or as a charge and/or charge effect recipient to mediate biological activities. In particular aspects, the inventive charge-stabilized nanostructures and/or the charge-
10 stabilized oxygen-containing nanostructures harboring a solvated electron interact with cellular membranes as a charge and/or charge effect donor and/or as a charge and/or charge effect recipient to mediate biological activities.

In particular aspects, the inventive charge-stabilized nanostructures and/or the charge-stabilized oxygen-containing nanostructures are consistent with, and account
15 for the observed stability and biological properties of the inventive electrokinetic fluids, and further provide a novel electrone (or inverted electrone) that provides for stabilized solvated electrons in aqueous ionic solutions (e.g., saline solutions, NaCl, etc.).

In particular aspects, the charge-stabilized oxygen-containing nanostructures substantially comprise, take the form of, or can give rise to, charge-stabilized oxygen-
20 containing nanobubbles. In particular aspects, charge-stabilized oxygen-containing clusters provide for formation of relatively larger arrays of charge-stabilized oxygen-containing nanostructures, and/or charge-stabilized oxygen-containing nanobubbles or arrays thereof. In particular aspects, the charge-stabilized oxygen-containing nanostructures can provide for formation of hydrophobic nanobubbles upon contact
25 with a hydrophobic surface (see elsewhere herein under EXAMPLE 25).

In particular aspects, the charge-stabilized oxygen-containing nanostructures substantially comprise at least one oxygen molecule. In certain aspects, the charge-stabilized oxygen-containing nanostructures substantially comprise at least 1, at least 2, at least 3, at least 4, at least 5, at least 10 at least 15, at least 20, at least 50, at least
30 100, or greater oxygen molecules. In particular aspects, charge-stabilized oxygen-containing nanostructures comprise or give rise to nanobubbles (e.g., hydrophobic

nanobubbles) of about 20 nm x 1.5 nm, comprise about 12 oxygen molecules (e.g., based on the size of an oxygen molecule (approx 0.3 nm by 0.4 nm), assumption of an ideal gas and application of $n=PV/RT$, where $P=1$ atm, $R=0.082057$ l.atm/mol.K; $T=295$ K; $V=\pi r^2 h=4.7 \times 10^{-22}$ L, where $r=10 \times 10^{-9}$ m, $h=1.5 \times 10^{-9}$ m, and $n=1.95 \times 10^{-22}$ moles).

In certain aspects, the percentage of oxygen molecules present in the fluid that are in such nanostructures, or arrays thereof, having a charge-stabilized configuration in the ionic aqueous fluid is a percentage amount selected from the group consisting of greater than: 0.1%, 1%; 2%; 5%; 10%; 15%; 20%; 25%; 30%; 35%; 40%; 45%; 50%; 55%; 60%; 65%; 70%; 75%; 80%; 85%; 90%; and greater than 95%. Preferably, this percentage is greater than about 5%, greater than about 10%, greater than about 15%, or greater than about 20%. In additional aspects, the substantial size of the charge-stabilized oxygen-containing nanostructures, or arrays thereof, having a charge-stabilized configuration in the ionic aqueous fluid is a size selected from the group consisting of less than: 100 nm; 90 nm; 80 nm; 70 nm; 60 nm; 50 nm; 40 nm; 30 nm; 20 nm; 10 nm; 5 nm; 4 nm; 3 nm; 2 nm; and 1 nm. Preferably, this size is less than about 50 nm, less than about 40 nm, less than about 30 nm, less than about 20 nm, or less than about 10 nm.

In certain aspects, the inventive electrokinetic fluids comprise solvated electrons. In further aspects, the inventive electrokinetic fluids comprises charge-stabilized nanostructures and/or charge-stabilized oxygen-containing nanostructures, and/or arrays thereof, which comprise at least one of: solvated electron(s); and unique charge distributions (polar, symmetric, asymmetric charge distribution). In certain aspects, the charge-stabilized nanostructures and/or charge-stabilized oxygen-containing nanostructures, and/or arrays thereof, have paramagnetic properties.

By contrast, relative to the inventive electrokinetic fluids, control pressure pot oxygenated fluids (non-electrokinetic fluids) and the like do *not* comprise such charge-stabilized biologically-active nanostructures and/or biologically-active charge-stabilized oxygen-containing nanostructures and/or arrays thereof, capable of modulation of at least one of cellular membrane potential and cellular membrane conductivity.

Systems for Making Gas-Enriched Fluids

The presently disclosed system and methods allow gas (e.g. oxygen) to be enriched stably at a high concentration with minimal passive loss. This system and methods can be effectively used to enrich a wide variety of gases at heightened percentages into a wide variety of fluids. By way of example only, deionized water at room temperature that typically has levels of about 2-3 ppm (ppm) of dissolved oxygen can achieve levels of dissolved oxygen ranging from at least about 5 ppm, at least about 10 ppm, at least about 15 ppm, at least about 20 ppm, at least about 25 ppm, at least about 30 ppm, at least about 35 ppm, at least about 40 ppm, at least about 45 ppm, at least about 50 ppm, at least about 55 ppm, at least about 60 ppm, at least about 65 ppm, at least about 70 ppm, at least about 75 ppm, at least about 80 ppm, at least about 85 ppm, at least about 90 ppm, at least about 95 ppm, at least about 100 ppm, or any value greater or therebetween using the disclosed systems and/or methods. In accordance with a particular exemplary embodiment, oxygen-enriched water may be generated with levels of about 30-60 ppm of dissolved oxygen.

Routes and Forms of Administration

As used herein, "subject," may refer to any living creature, preferably an animal, more preferably a mammal, and even more preferably a human.

In particular exemplary embodiments, the gas-enriched fluid of the present invention may function as a therapeutic composition alone or in combination with another therapeutic agent such that the therapeutic composition prevents or alleviates at least one symptom of a disorder associated with insulin resistance and/or diabetes. The therapeutic compositions of the present invention include compositions that are able to be administered to a subject in need thereof. In certain embodiments, the therapeutic composition formulation may also comprise at least one additional agent selected from the group consisting of: carriers, adjuvants, emulsifying agents, suspending agents, sweeteners, flavorings, perfumes, and binding agents.

As used herein, "pharmaceutically acceptable carrier" and "carrier" generally refer to a non-toxic, inert solid, semi-solid or liquid filler, diluent, encapsulating material, or formulation auxiliary of any type. Some non-limiting examples of materials which can serve as pharmaceutically-acceptable carriers are sugars such as lactose, glucose, and

sucrose; starches such as corn starch and potato starch; cellulose and its derivatives such as sodium carboxymethyl cellulose, ethyl cellulose, and cellulose acetate; powdered tragacanth; malt; gelatin; talc; excipients such as cocoa butter and suppository waxes; oils such as peanut oil, cottonseed oil, safflower oil, sesame oil, olive oil, corn oil, and soybean oil; glycols, such as propylene glycol; esters such as ethyl oleate and ethyl laurate; agar; buffering agents such as magnesium hydroxide and aluminum hydroxide; alginic acid; pyrogen-free water, isotonic saline, Ringer's solution, ethyl alcohol and phosphate buffer solutions; as well as other non-toxic compatible lubricants such as sodium lauryl sulfate and magnesium stearate; as well as coloring agents; releasing agents; coating agents; sweetening, flavoring and perfuming agents; and preservatives and antioxidants can also be present in the composition, according to the judgment of the formulator. In particular aspects, such carriers and excipients may be gas-enriched fluids or solutions of the present invention.

The pharmaceutically acceptable carriers described herein, for example, vehicles, adjuvants, excipients, or diluents, are well known to those who are skilled in the art. Typically, the pharmaceutically acceptable carrier is chemically inert to the therapeutic agents and has no detrimental side effects or toxicity under the conditions of use. The pharmaceutically acceptable carriers can include polymers and polymer matrices, nanoparticles, microbubbles, and the like.

In addition to the therapeutic gas-enriched fluid of the present invention, the therapeutic composition may further comprise inert diluents such as additional non-gas-enriched water or other solvents, solubilizing agents and emulsifiers such as ethyl alcohol, isopropyl alcohol, ethyl carbonate, ethyl acetate, benzyl alcohol, benzyl benzoate, propylene glycol, 1,3-butylene glycol, dimethylformamide, oils (in particular, cottonseed, groundnut, corn, germ, olive, castor, and sesame oils), glycerol, tetrahydrofurfuryl alcohol, polyethylene glycols and fatty acid esters of sorbitan, and mixtures thereof. As is appreciated by those of ordinary skill, a novel and improved formulation of a particular therapeutic composition, a novel gas-enriched therapeutic fluid, and a novel method of delivering the novel gas-enriched therapeutic fluid may be obtained by replacing one or more inert diluents with a gas-enriched fluid of identical, similar, or different composition. For example, conventional water may be replaced or

supplemented by a gas-enriched fluid produced by mixing oxygen into water or deionized water to provide gas-enriched fluid.

In certain embodiments, the inventive gas-enriched fluid may be combined with one or more therapeutic agents and/or used alone. In particular embodiments, incorporating the gas-enriched fluid may include replacing one or more solutions known in the art, such as deionized water, saline solution, and the like with one or more gas-enriched fluid, thereby providing an improved therapeutic composition for delivery to the subject.

Certain embodiments provide for therapeutic compositions comprising a gas-enriched fluid of the present invention, a pharmaceutical composition or other therapeutic agent or a pharmaceutically-acceptable salt or solvate thereof, and at least one pharmaceutical carrier or diluent. These pharmaceutical compositions may be used in the prophylaxis and treatment of the foregoing diseases or conditions and in therapies as mentioned above. Preferably, the carrier must be pharmaceutically acceptable and must be compatible with, i.e. not have a deleterious effect upon, the other ingredients in the composition. The carrier may be a solid or liquid and is preferably formulated as a unit dose formulation, for example, a tablet that may contain from 0.05 to 95% by weight of the active ingredient.

Possible administration routes include oral, sublingual, buccal, parenteral (for example subcutaneous, intramuscular, intra-arterial, intraperitoneally, intracisternally, intravesically, intrathecally, or intravenous), rectal, topical including transdermal, intravaginal, intraocular, intraotical, intranasal, inhalation, and injection or insertion of implantable devices or materials.

Administration Routes

Most suitable means of administration for a particular subject will depend on the nature and severity of the disease or condition being treated or the nature of the therapy being used, as well as the nature of the therapeutic composition or additional therapeutic agent. In certain embodiments, oral or topical administration is preferred.

Formulations suitable for oral administration may be provided as discrete units, such as tablets, capsules, cachets, syrups, elixirs, chewing gum, "lollipop" formulations, microemulsions, solutions, suspensions, lozenges, or gel-coated ampules, each

containing a predetermined amount of the active compound; as powders or granules; as solutions or suspensions in aqueous or non-aqueous liquids; or as oil-in-water or water-in-oil emulsions.

Formulations suitable for transmucosal methods, such as by sublingual or buccal administration include lozenges patches, tablets, and the like comprising the active compound and typically, a flavored base such as sugar and acacia or tragacanth, and pastilles comprising the active compound in an inert base, such as gelatin and glycerine or sucrose acacia.

Formulations suitable for parenteral administration typically comprise sterile aqueous solutions containing a predetermined concentration of the active gas-enriched fluid and possibly another therapeutic agent; the solution is preferably isotonic with the blood of the intended recipient. Additional formulations suitable for parenteral administration include formulations containing physiologically suitable co-solvents and/or complexing agents such as surfactants and cyclodextrins. Oil-in-water emulsions may also be suitable for formulations for parenteral administration of the gas-enriched fluid. Although such solutions are preferably administered intravenously, they may also be administered by subcutaneous or intramuscular injection.

Formulations suitable for urethral, rectal, or vaginal administration include gels, creams, lotions, aqueous or oily suspensions, dispersible powders or granules, emulsions, dissolvable solid materials, douches, and the like. The formulations are preferably provided as unit-dose suppositories comprising the active ingredient in one or more solid carriers forming the suppository base, for example, cocoa butter. Alternatively, colonic washes with the gas-enriched fluids of the present invention may be formulated for colonic or rectal administration.

Formulations suitable for topical, intraocular, intraotic, or intranasal application include ointments, creams, pastes, lotions, gels (such as hydrogels), sprays, dispersible powders and granules, emulsions, sprays or aerosols using flowing propellants (such as liposomal sprays, nasal drops, nasal sprays, and the like) and oils. Suitable carriers for such formulations include petroleum jelly, lanolin, polyethyleneglycols, alcohols, and combinations thereof. Nasal or intranasal delivery may include metered doses of any of these formulations or others. Likewise, intraotic or intraocular may include drops, ointments, irritation fluids, and the like.

Formulations of the invention may be prepared by any suitable method, typically by uniformly and intimately admixing the gas-enriched fluid optionally with an active compound with liquids or finely-divided solid carriers or both, in the required proportions and then, if necessary, shaping the resulting mixture into the desired shape.

5 For example a tablet may be prepared by compressing an intimate mixture comprising a powder or granules of the active ingredient and one or more optional ingredients, such as a binder, lubricant, inert diluent, or surface active dispersing agent, or by molding an intimate mixture of powdered active ingredient and a gas-enriched fluid of the present invention.

10 Suitable formulations for administration by inhalation include fine particle dusts or mists which may be generated by means of various types of metered dose pressurized aerosols, nebulisers, or insufflators. In particular, powders or other compounds of therapeutic agents may be dissolved or suspended in a gas-enriched fluid of the present invention.

15 For pulmonary administration via the mouth, the particle size of the powder or droplets is typically in the range 0.5-10 μM , preferably 1-5 μM , to ensure delivery into the bronchial tree. For nasal administration, a particle size in the range 10-500 μM is preferred to ensure retention in the nasal cavity.

Metered dose inhalers are pressurized aerosol dispensers, typically containing a
20 suspension or solution formulation of a therapeutic agent in a liquefied propellant. In certain embodiments, as disclosed herein, the gas-enriched fluids of the present invention may be used in addition to or instead of the standard liquefied propellant. During use, these devices discharge the formulation through a valve adapted to deliver a metered volume, typically from 10 to 150 μL , to produce a fine particle spray
25 containing the therapeutic agent and the gas-enriched fluid. Suitable propellants include certain chlorofluorocarbon compounds, for example, dichlorodifluoromethane, trichlorofluoromethane, dichlorotetrafluoroethane, and mixtures thereof.

The formulation may additionally contain one or more co-solvents, for example, ethanol surfactants, such as oleic acid or sorbitan trioleate, anti-oxidants and suitable
30 flavoring agents. Nebulisers are commercially available devices that transform solutions or suspensions of the active ingredient into a therapeutic aerosol mist either by means of acceleration of a compressed gas (typically air or oxygen) through a

narrow venturi orifice, or by means of ultrasonic agitation. Suitable formulations for use in nebulisers consist of another therapeutic agent in a gas-enriched fluid and comprising up to 40% w/w of the formulation, preferably less than 20% w/w. In addition, other carriers may be utilized, such as distilled water, sterile water, or a dilute aqueous alcohol solution, preferably made isotonic with body fluids by the addition of salts, such as sodium chloride. Optional additives include preservatives, especially if the formulation is not prepared sterile, and may include methyl hydroxy-benzoate, antioxidants, flavoring agents, volatile oils, buffering agents, and surfactants.

Suitable formulations for administration by insufflation include finely comminuted powders that may be delivered by means of an insufflator or taken into the nasal cavity in the manner of a snuff. In the insufflator, the powder is contained in capsules or cartridges, typically made of gelatin or plastic, which are either pierced or opened in situ and the powder delivered by air drawn through the device upon inhalation or by means of a manually-operated pump. The powder employed in the insufflator consists either solely of the active ingredient or of a powder blend comprising the active ingredient, a suitable powder diluent, such as lactose, and an optional surfactant. The active ingredient typically comprises from 0.1 to 100 w/w of the formulation.

In addition to the ingredients specifically mentioned above, the formulations of the present invention may include other agents known to those skilled in the art, having regard for the type of formulation in issue. For example, formulations suitable for oral administration may include flavoring agents and formulations suitable for intranasal administration may include perfumes.

The therapeutic compositions of the invention can be administered by any conventional method available for use in conjunction with pharmaceutical drugs, either as individual therapeutic agents or in a combination of therapeutic agents.

The dosage administered will, of course, vary depending upon known factors, such as the pharmacodynamic characteristics of the particular agent and its mode and route of administration; the age, health and weight of the recipient; the nature and extent of the symptoms; the kind of concurrent treatment; the frequency of treatment; and the effect desired. A daily dosage of active ingredient can be expected to be about 0.001 to 1000 milligrams (mg) per kilogram (kg) of body weight, with the preferred dose being 0.1 to about 30 mg/kg.

Dosage forms (compositions suitable for administration) contain from about 1 mg to about 500 mg of active ingredient per unit. In these pharmaceutical compositions, the active ingredient will ordinarily be present in an amount of about 0.5-95% weight based on the total weight of the composition.

5 Ointments, pastes, foams, occlusions, creams and gels also can contain excipients, such as starch, tragacanth, cellulose derivatives, silicones, bentonites, silica acid, and talc, or mixtures thereof. Powders and sprays also can contain excipients such as lactose, talc, silica acid, aluminum hydroxide, and calcium silicates, or mixtures of these substances. Solutions of nanocrystalline antimicrobial metals can be
10 converted into aerosols or sprays by any of the known means routinely used for making aerosol pharmaceuticals. In general, such methods comprise pressurizing or providing a means for pressurizing a container of the solution, usually with an inert carrier gas, and passing the pressurized gas through a small orifice. Sprays can additionally contain customary propellants, such as nitrogen, carbon dioxide, and other inert gases.
15 In addition, microspheres or nanoparticles may be employed with the gas-enriched therapeutic compositions or fluids of the present invention in any of the routes required to administer the therapeutic compounds to a subject.

The injection-use formulations can be presented in unit-dose or multi-dose sealed containers, such as ampules and vials, and can be stored in a freeze-dried
20 (lyophilized) condition requiring only the addition of the sterile liquid excipient, or gas-enriched fluid, immediately prior to use. Extemporaneous injection solutions and suspensions can be prepared from sterile powders, granules, and tablets. The requirements for effective pharmaceutical carriers for injectable compositions are well known to those of ordinary skill in the art. See, for example, *Pharmaceutics and*
25 *Pharmacy Practice*, J. B. Lippincott Co., Philadelphia, Pa., Banker and Chalmers, Eds., 238-250 (1982) and *ASHP Handbook on Injectable Drugs*, Toissel, 4th ed., 622-630 (1986).

Formulations suitable for topical administration include lozenges comprising a gas-enriched fluid of the invention and optionally, an additional therapeutic and a flavor,
30 usually sucrose and acacia or tragacanth; pastilles comprising a gas-enriched fluid and optional additional therapeutic agent in an inert base, such as gelatin and glycerin, or sucrose and acacia; and mouth washes or oral rinses comprising a gas-enriched fluid

and optional additional therapeutic agent in a suitable liquid carrier; as well as creams, emulsions, gels, and the like.

Additionally, formulations suitable for rectal administration may be presented as suppositories by mixing with a variety of bases such as emulsifying bases or water-
5 soluble bases. Formulations suitable for vaginal administration may be presented as pessaries, tampons, creams, gels, pastes, foams, or spray formulas containing, in addition to the active ingredient, such carriers as are known in the art to be appropriate.

Suitable pharmaceutical carriers are described in Remington's Pharmaceutical Sciences, Mack Publishing Company, a standard reference text in this field.

10 The dose administered to a subject, especially an animal, particularly a human, in the context of the present invention should be sufficient to effect a therapeutic response in the animal over a reasonable time frame. One skilled in the art will recognize that dosage will depend upon a variety of factors including the condition of the animal, the body weight of the animal, as well as the condition being treated. A
15 suitable dose is that which will result in a concentration of the therapeutic composition in a subject that is known to affect the desired response.

The size of the dose also will be determined by the route, timing and frequency of administration as well as the existence, nature, and extent of any adverse side effects that might accompany the administration of the therapeutic composition and the
20 desired physiological effect.

Most suitable means of administration for a particular subject will depend on the nature and severity of the disease or condition being treated or the nature of the therapy being used, as well as the nature of the therapeutic composition or additional therapeutic agent. In certain embodiments, oral or topical administration is preferred.

25 Formulations suitable for oral administration may be provided as discrete units, such as tablets, capsules, cachets, syrups, elixirs, chewing gum, "lollipop" formulations, microemulsions, solutions, suspensions, lozenges, or gel-coated ampules, each containing a predetermined amount of the active compound; as powders or granules; as solutions or suspensions in aqueous or non-aqueous liquids; or as oil-in-water or
30 water-in-oil emulsions.

Additional formulations suitable for oral administration may be provided to include fine particle dusts or mists which may be generated by means of various types

of metered dose pressurized aerosols, atomizers, nebulisers, or insufflators. In particular, powders or other compounds of therapeutic agents may be dissolved or suspended in a gas-enriched fluid of the present invention.

Formulations suitable for transmucosal methods, such as by sublingual or buccal
5 administration include lozenges patches, tablets, and the like comprising the active compound and, typically a flavored base, such as sugar and acacia or tragacanth and pastilles comprising the active compound in an inert base, such as gelatin and glycerine or sucrose acacia.

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10 aqueous solutions containing a predetermined concentration of the active gas-enriched fluid and possibly another therapeutic agent; the solution is preferably isotonic with the blood of the intended recipient. Additional formulations suitable for parenteral administration include formulations containing physiologically suitable co-solvents and/or complexing agents such as surfactants and cyclodextrins. Oil-in-water
15 emulsions may also be suitable for formulations for parenteral administration of the gas-enriched fluid. Although such solutions are preferably administered intravenously, they may also be administered by subcutaneous or intramuscular injection.

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20 emulsions, dissolvable solid materials, douches, and the like. The formulations are preferably provided as unit-dose suppositories comprising the active ingredient in one or more solid carriers forming the suppository base, for example, cocoa butter. Alternatively, colonic washes with the gas-enriched fluids of the present invention may be formulated for colonic or rectal administration.

Formulations suitable for topical, intraocular, intraotic, or intranasal application
25 include ointments, creams, pastes, lotions, pastes, gels (such as hydrogels), sprays, dispersible powders and granules, emulsions, sprays or aerosols using flowing propellants (such as liposomal sprays, nasal drops, nasal sprays, and the like) and oils. Suitable carriers for such formulations include petroleum jelly, lanolin,
30 polyethyleneglycols, alcohols, and combinations thereof. Nasal or intranasal delivery may include metered doses of any of these formulations or others. Likewise, intraotic

or intraocular may include drops, ointments, irritation fluids and the like.

Formulations of the invention may be prepared by any suitable method, typically by uniformly and intimately admixing the gas-enriched fluid optionally with an active compound with liquids or finely divided solid carriers or both, in the required proportions
5 and then, if necessary, shaping the resulting mixture into the desired shape.

For example a tablet may be prepared by compressing an intimate mixture comprising a powder or granules of the active ingredient and one or more optional ingredients, such as a binder, lubricant, inert diluent, or surface active dispersing agent, or by molding an intimate mixture of powdered active ingredient and a gas-enriched
10 fluid of the present invention.

Suitable formulations for administration by inhalation include fine particle dusts or mists which may be generated by means of various types of metered dose pressurized aerosols, atomizers, nebulisers, or insufflators. In particular, powders or other compounds of therapeutic agents may be dissolved or suspended in a gas-
15 enriched fluid of the present invention.

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Metered dose inhalers are pressurized aerosol dispensers, typically containing a suspension or solution formulation of a therapeutic agent in a liquefied propellant. In certain embodiments, as disclosed herein, the gas-enriched fluids of the present invention may be used in addition to or instead of the standard liquefied propellant. During use, these devices discharge the formulation through a valve adapted to deliver
20 a metered volume, typically from 10 to 150 μL , to produce a fine particle spray containing the therapeutic agent and the gas-enriched fluid. Suitable propellants include certain chlorofluorocarbon compounds, for example, dichlorodifluoromethane, trichlorofluoromethane, dichlorotetrafluoro-ethane and mixtures thereof.

The formulation may additionally contain one or more co-solvents, for example, ethanol surfactants, such as oleic acid or sorbitan trioleate, anti-oxidants and suitable
30 flavoring agents. Nebulisers are commercially available devices that transform

solutions or suspensions of the active ingredient into a therapeutic aerosol mist either by means of acceleration of a compressed gas (typically air or oxygen) through a narrow venturi orifice, or by means of ultrasonic agitation. Suitable formulations for use in nebulisers consist of another therapeutic agent in a gas-enriched fluid and comprising up to 40% w/w of the formulation, preferably less than 20% w/w. In addition, other carriers may be utilized, such as distilled water, sterile water, or a dilute aqueous alcohol solution, preferably made isotonic with body fluids by the addition of salts, such as sodium chloride. Optional additives include preservatives, especially if the formulation is not prepared sterile, and may include methyl hydroxy-benzoate, anti-oxidants, flavoring agents, volatile oils, buffering agents and surfactants.

Suitable formulations for administration by insufflation include finely comminuted powders that may be delivered by means of an insufflator or taken into the nasal cavity in the manner of a snuff. In the insufflator, the powder is contained in capsules or cartridges, typically made of gelatin or plastic, which are either pierced or opened in situ and the powder delivered by air drawn through the device upon inhalation or by means of a manually-operated pump. The powder employed in the insufflator consists either solely of the active ingredient or of a powder blend comprising the active ingredient, a suitable powder diluent, such as lactose, and an optional surfactant. The active ingredient typically comprises from 0.1 to 100 w/w of the formulation.

In addition to the ingredients specifically mentioned above, the formulations of the present invention may include other agents known to those skilled in the art, having regard for the type of formulation in issue. For example, formulations suitable for oral administration may include flavoring agents and formulations suitable for intranasal administration may include perfumes.

The therapeutic compositions of the invention can be administered by any conventional method available for use in conjunction with pharmaceutical drugs, either as individual therapeutic agents or in a combination of therapeutic agents.

The dosage administered will, of course, vary depending upon known factors, such as the pharmacodynamic characteristics of the particular agent and its mode and route of administration; the age, health and weight of the recipient; the nature and extent of the symptoms; the kind of concurrent treatment; the frequency of treatment;

and the effect desired. A daily dosage of active ingredient can be expected to be about 0.001 to 1000 milligrams (mg) per kilogram (kg) of body weight, with the preferred dose being 0.1 to about 30 mg/kg. According to certain aspects daily dosage of active ingredient may be .001 liters to 10 liters, with the preferred dose being from about .01
5 liters to 1 liter.

Dosage forms (compositions suitable for administration) contain from about 1 mg to about 500 mg of active ingredient per unit. In these pharmaceutical compositions, the active ingredient will ordinarily be present in an amount of about 0.5-95% weight based on the total weight of the composition.

10 Ointments, pastes, foams, occlusions, creams and gels also can contain excipients, such as starch, tragacanth, cellulose derivatives, silicones, bentonites, silica acid, and talc, or mixtures thereof. Powders and sprays also can contain excipients such as lactose, talc, silica acid, aluminum hydroxide, and calcium silicates, or mixtures of these substances. Solutions of nanocrystalline antimicrobial metals can be
15 converted into aerosols or sprays by any of the known means routinely used for making aerosol pharmaceuticals. In general, such methods comprise pressurizing or providing a means for pressurizing a container of the solution, usually with an inert carrier gas, and passing the pressurized gas through a small orifice. Sprays can additionally contain customary propellants, such as nitrogen, carbon dioxide, and other inert gases.
20 In addition, microspheres or nanoparticles may be employed with the gas-enriched therapeutic compositions or fluids of the present invention in any of the routes required to administer the therapeutic compounds to a subject.

The injection-use formulations can be presented in unit-dose or multi-dose sealed containers, such as ampules and vials, and can be stored in a freeze-dried
25 (lyophilized) condition requiring only the addition of the sterile liquid excipient, or gas-enriched fluid, immediately prior to use. Extemporaneous injection solutions and suspensions can be prepared from sterile powders, granules, and tablets. The requirements for effective pharmaceutical carriers for injectable compositions are well known to those of ordinary skill in the art. See, for example, *Pharmaceutics and
30 Pharmacy Practice*, J. B. Lippincott Co., Philadelphia, Pa., Banker and Chalmers, Eds., 238-250 (1982) and *ASHP Handbook on Injectable Drugs*, Toissel, 4th ed., 622-630

(1986).

Formulations suitable for topical administration include lozenges comprising a gas-enriched fluid of the invention and optionally, an additional therapeutic and a flavor, usually sucrose and acacia or tragacanth; pastilles comprising a gas-enriched fluid and optional additional therapeutic agent in an inert base, such as gelatin and glycerin, or sucrose and acacia; and mouth washes or oral rinses comprising a gas-enriched fluid and optional additional therapeutic agent in a suitable liquid carrier; as well as creams, emulsions, gels and the like.

Additionally, formulations suitable for rectal administration may be presented as suppositories by mixing with a variety of bases such as emulsifying bases or water-soluble bases. Formulations suitable for vaginal administration may be presented as pessaries, tampons, creams, gels, pastes, foams, or spray formulas containing, in addition to the active ingredient, such carriers as are known in the art to be appropriate.

Suitable pharmaceutical carriers are described in Remington's Pharmaceutical Sciences, Mack Publishing Company, a standard reference text in this field.

The dose administered to a subject, especially an animal, particularly a human, in the context of the present invention should be sufficient to effect a therapeutic response in the animal over a reasonable time frame. One skilled in the art will recognize that dosage will depend upon a variety of factors including the condition of the animal, the body weight of the animal, as well as the condition being treated. A suitable dose is that which will result in a concentration of the therapeutic composition in a subject that is known to affect the desired response.

The size of the dose also will be determined by the route, timing and frequency of administration as well as the existence, nature, and extent of any adverse side effects that might accompany the administration of the therapeutic composition and the desired physiological effect.

It will be appreciated that the compounds of the combination may be administered: (1) simultaneously by combination of the compounds in a co-formulation or (2) by alternation, i.e. delivering the compounds serially, sequentially, in parallel or simultaneously in separate pharmaceutical formulations. In alternation therapy, the delay in administering the second, and optionally a third active ingredient, should not be

such as to lose the benefit of a synergistic therapeutic effect of the combination of the active ingredients. According to certain embodiments by either method of administration (1) or (2), ideally the combination should be administered to achieve the most efficacious results. In certain embodiments by either method of administration (1) or (2),
5 ideally the combination should be administered to achieve peak plasma concentrations of each of the active ingredients. A one pill once-per-day regimen by administration of a combination co-formulation may be feasible for some patients suffering from inflammatory neurodegenerative diseases. According to certain embodiments effective peak plasma concentrations of the active ingredients of the combination will be in the
10 range of approximately 0.001 to 100 μ M. Optimal peak plasma concentrations may be achieved by a formulation and dosing regimen prescribed for a particular patient. It will also be understood that the inventive fluids and glatiramer acetate, interferon-beta, mitoxantrone, and/or natalizumab or the physiologically functional derivatives of any thereof, whether presented simultaneously or sequentially, may be administered
15 individually, in multiples, or in any combination thereof. In general, during alternation therapy (2), an effective dosage of each compound is administered serially, where in co-formulation therapy (1), effective dosages of two or more compounds are administered together.

The combinations of the invention may conveniently be presented as a
20 pharmaceutical formulation in a unitary dosage form. A convenient unitary dosage formulation contains the active ingredients in any amount from 1 mg to 1 g each, for example but not limited to, 10 mg to 300 mg. The synergistic effects of the inventive fluid in combination with glatiramer acetate, interferon-beta, mitoxantrone, and/or natalizumab may be realized over a wide ratio, for example 1:50 to 50:1 (inventive fluid:
25 glatiramer acetate, interferon-beta, mitoxantrone, and/or natalizumab). In one embodiment the ratio may range from about 1:10 to 10:1. In another embodiment, the weight/weight ratio of inventive fluid to glatiramer acetate, interferon-beta, mitoxantrone, and/or natalizumab in a co-formulated combination dosage form, such as a pill, tablet, caplet or capsule will be about 1, i.e. an approximately equal amount of inventive fluid
30 and glatiramer acetate, interferon-beta, mitoxantrone, and/or natalizumab. In other exemplary co-formulations, there may be more or less inventive fluid and glatiramer

acetate, interferon-beta, mitoxantrone, and/or natalizumab. In one embodiment, each compound will be employed in the combination in an amount at which it exhibits anti-inflammatory activity when used alone. Other ratios and amounts of the compounds of said combinations are contemplated within the scope of the invention.

5 A unitary dosage form may further comprise inventive fluid and glatiramer acetate, interferon-beta, mitoxantrone, and/or natalizumab, or physiologically functional derivatives of either thereof, and a pharmaceutically acceptable carrier.

It will be appreciated by those skilled in the art that the amount of active ingredients in the combinations of the invention required for use in treatment will vary
10 according to a variety of factors, including the nature of the condition being treated and the age and condition of the patient, and will ultimately be at the discretion of the attending physician or health care practitioner. The factors to be considered include the route of administration and nature of the formulation, the animal's body weight, age and general condition and the nature and severity of the disease to be treated.

15 It is also possible to combine any two of the active ingredients in a unitary dosage form for simultaneous or sequential administration with a third active ingredient. The three-part combination may be administered simultaneously or sequentially. When administered sequentially, the combination may be administered in two or three administrations. According to certain embodiments the three-part combination of
20 inventive fluid and glatiramer acetate, interferon-beta, mitoxantrone, and/or natalizumab may be administered in any order.

The following examples are meant to be illustrative only and not limiting in any way.

25 EXAMPLES

EXAMPLE 1

Dissolved Oxygen Stability

As indicated in Figure 30, there is illustrated the dissolved oxygen levels in a 500
30 ml thin-walled plastic bottle and a 1000 ml glass bottle which were each capped and stored at 65 degrees Fahrenheit.

As can be seen, when the plastic bottle is opened approximately 65 days after bottling, the dissolved oxygen level within the water is approximately 27.5 ppm. When a second bottle is opened at approximately 95 days after bottling, the dissolved oxygen level is approximately 25 ppm. Likewise, for the glass bottle, the dissolved oxygen level is approximately 40 ppm at 65 days and is approximately 41 ppm at 95 days. Thus, this chart indicates that the dissolved oxygen levels within both plastic and glass bottles are maintained at relatively high rates at 65 degrees Fahrenheit when the oxygen is diffused within the fluid using the described system and method.

EXAMPLE 2

Decayed Oxygen Content in Balanced Salt Solution

Figure 33 illustrates the dissolved oxygen retention of a 500 ml balanced salt solution that originally had a dissolved oxygen level of 5 ppm. Following enrichment of the solution at standard temperature and pressure with the diffuser of the present invention, the dissolved oxygen level was approximately 41 ppm. The solution was kept in an amber glass bottle. After an hour, the dissolved oxygen level was 40 ppm; 36 ppm after two hours; 34 ppm after three hours; and slightly more than 30 ppm after approximately four and a half hours. The final measurement was taken shortly before six hours, at which point the dissolved oxygen level was approximately 28 ppm.

EXAMPLE 3

Microbubble Size

Experiments were performed with a gas-enriched fluid by using the diffuser of the present invention in order to determine a gas microbubble size limit. The microbubble size limit was established by passing the gas-enriched fluid through 0.22 and 0.1 micron filters. In performing these tests, a volume of fluid passed through the diffuser of the present invention and generated a gas-enriched fluid. Sixty ml of this fluid was drained into a 60 ml syringe. The dissolved oxygen level of the fluid within the syringe was then measured by Winkler Titration. The fluid within the syringe was injected through a 0.22 micron Millipore Millex GP50 filter and into a 50 ml beaker. The dissolved oxygen rate of the material in the 50 ml beaker was then measured. The

experiment was performed three times to achieve the results illustrated in Table 6 below.

TABLE 6. Dissolved oxygen.

5

DO IN SYRINGE	DO AFTER 0.22 MICRON FILTER
42.1 ppm	39.7 ppm
43.4 ppm	42.0 ppm
43.5 ppm	39.5 ppm

As can be seen, the dissolved oxygen levels that were measured within the syringe and the dissolved oxygen levels within the 50 ml beaker were not significantly changed by passing the diffused material through a 0.22 micron filter, which implies that the microbubbles of dissolved gas within the fluid are not larger than 0.22 microns.

A second test was performed in which a batch of saline solution was enriched with the diffuser of the present invention and a sample of the output solution was collected in an unfiltered state. The dissolved oxygen level of the unfiltered sample was 44.7 ppm. A 0.1 micron filter was used to filter the oxygen-enriched solution from the diffuser of the present invention and two additional samples were taken. For the first sample, the dissolved oxygen level was 43.4 ppm. For the second sample, the dissolved oxygen level was 41.4 ppm. Finally, the filter was removed and a final sample was taken from the unfiltered solution. In this case, the final sample had a dissolved oxygen level of 45.4 ppm. These results were consistent with those in which the Millipore 0.22 micron filter was used. Thus, the majority of the gas bubbles or microbubbles within the saline solution are approximately less than 0.1 microns in size.

EXAMPLE 4

Sparging Effects

Figures 34 and 35 illustrate the sparging affects of the diffuser of the present invention on a fluid passing therethrough. The sparging of oxygen-enriched water

occurred in an 8 gallon tank at standard temperature and pressure. As indicated, initially the oxygen-enriched water had a dissolved oxygen level of approximately 42 ppm. After 2 minutes of running through the diffuser, the nitrogen had sparged the oxygen-enriched water such that the dissolved oxygen level was then slightly more than 20 ppm. At 6 minutes, the dissolved oxygen level was approximately 6 ppm. The dissolved oxygen level of the oxygen-enriched water reached a minimum value slightly greater than zero (0) at approximately 14 minutes after the beginning of the process. These figures illustrate the manner in which nitrogen may be diffused into water to sparge the oxygen from the water. However, any gas could be used within any fluid to sparge one gas from the other and diffuse the other gas into the fluid. The same experiment could utilize any host fluid material, and any fluid infusion material.

EXAMPLE 5

Rayleigh Effects

Fluids processed through the diffuser device described herein exhibit differences within the structure of the water when compared with normal unprocessed water. Gas-enriched water made by embodiments disclosed herein has been shown to have more Rayleigh scattering compared to unprocessed water.

In experiments conducted, samples of gas-enriched and non-enriched water were prepared and sent for optical analysis. The purpose of these tests was to determine whether there are any gross optical differences between normal (unprocessed) deionized water and water enriched by the diffuser device of the present invention.

The two samples, were coded to maintain their identities in secrecy, and only after the tests were completed were the samples identified. The two samples were placed in a laser beam of 633 nanometers according to the diagram illustrated in Figure 37A. Sample B, which was gas-enriched fluid according to certain embodiments disclosed herein, exhibited scattered light regardless of its position relative to the laser source. The Sample B fluid had been sealed in glass bottles for approximately one week. After two to three hours of opening the bottle, the scattering effect disappeared. Thus, the structure of the gas-enriched fluid is optically different from the structure of the unprocessed fluid. The optical effect is not directly related to dissolved oxygen

levels since the dissolved oxygen level at the start was approximately 45 ppm and at the end of the experiment was estimated to be approximately 32 ppm. Results are shown in Figure 37B.

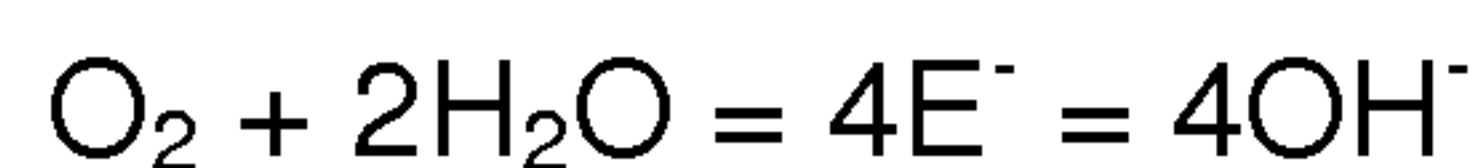
5

EXAMPLE 6

Generation of Solvated Electrons

Additional evidence has also indicated that the enriching process generated by the diffuser device of the present invention results in solvated electrons within the gas-enriched fluid. Due to the results of the polarographic dissolved oxygen probes, it is
10 believed that the diffused fluid exhibits an electron capture effect and thus the fluid included solvated electrons within the gas-enriched material.

There are two fundamental techniques for measuring dissolved oxygen levels electrically: galvanic measuring techniques and polarographic measurements. Each process uses an electrode system wherein the dissolved oxygen levels within the
15 solution being tested react with a cathode of the probe to produce a current. Dissolved oxygen-level sensors consist of two electrodes, an anode and a cathode, which are both immersed in electrolyte within the sensor body. An oxygen permeable membrane separates the anode and cathode from the solution being tested. Oxygen diffuses across the membrane and interacts with the internal components of the probe to
20 produce an electrical current. The cathode is a hydrogen electrode and carries negative potential with respect to the anode. The electrolyte solution surrounds the electrode pair and is contained by the membrane. When no oxygen is present, the cathode is polarized by hydrogen and resists the flow of current. When oxygen passes through the membrane, the cathode is depolarized and electrons are consumed. The
25 cathode electrochemically reduces the oxygen to hydroxyl ions according to the following equation:



When performing dissolved oxygen level measurements of a gas-enriched solution according to the systems of the present invention, an overflow condition has
30 been repeatedly experienced wherein the dissolved oxygen meter displays a reading that is higher than the meter is capable of reading. However, evaluation of the gas-enriched solution by Winkler Titration indicates lower dissolved oxygen (DO) level for

the solution than indicated by the probe. Typically, a DO probe (such as the Orion 862 used in these experiments) has a maximum reading of 60 ppm. However, when the meter is left in gas-enriched water of the present invention, it overflows.

Without wishing to be bound by any particular mechanism of action, the mechanism of the meter responds to electrons where the oxygen reacts. However, according to electron spin resonance, no free ions are present in the fluid. Thus, the fluid presumably contains solvated electrons stabilized by the oxygen species that is also present in the fluid.

EXAMPLE 7

In vitro Wound Healing

The effects of a gas-enriched fluid (enriched with oxygen) were tested for the ability of cultured human epidermal keratinocytes to seal a wound.

Human epidermal keratinocytes were isolated from neonatal foreskins that were obtained from routine circumcision and de-identified. Foreskins were washed twice in PBS and incubated in 2.4 U/mL Dispase II in order to separate the dermis from the epidermis. The epidermis was incubated with 0.25% trypsin/1 mM EDTA, neutralized with soy bean trypsin inhibitor, agitated, and passed through a 70 um sieve to separate the cells. Next, the cell suspension was centrifuged and resuspended in cell culture medium (M154) supplemented with 0.07 mM CaCl_2 , and human keratinocyte growth supplements (0.2% hydrocortisone, 0.2 ng/mL human epidermal growth factor) and penicillin/streptomycin, amphotericin antibiotic cocktail. The keratinocyte cell suspensions were plated onto uncoated 12-well culture dishes and the medium replaced after 24 hours, and every 48 hours after the initial seeding.

Upon reaching cellular confluence, linear scratches were made with a sterile p1000 pipette tip, which resulted in a uniform cell-free wound. The monolayers were washed several times with Dulbecco's PBS in order to remove any cellular debris. The wound monolayers were then incubated in the following media: i) the complete growth media (as described above in this Example); ii) the complete growth media diluted 1:1 with a sheared version of saline without oxygen (control fluid that was processed using the disclosed diffuser device but without adding a gas); and iii) the complete growth media diluted 1:1 with oxygen-enriched saline. Each study was done in triplicate.

Prior to incubation, the wells were filled with the respective media and sealed by placing a 25 x 25 mm glass coverslip on top of each well. At 6, 12, 24, and 48 hours post-wounding, oxygen measurements were made, and cultures were imaged.

Six hours post-wounding, the edges of the wounds in the saline and gas-enriched media were more ruffled than those in the media control that was processed with the diffuser device disclosed herein, but without the addition of a gas. Twelve hours post-wounding the edges of the wounds in all three media appeared uneven, with keratinocytes along the borders migrating toward the center of the wounds. Quantification of migrating keratinocytes revealed approximately the same level of keratinocyte migration in the saline and gas-enriched media. Results of the experiment are shown in Figures 40A and 44B.

EXAMPLE 8

Improved Wound Healing

A study was performed to determine the improved healing characteristics of wounds that were exposed to an oxygen-enriched saline solution that was processed according to embodiments disclosed herein. In this experiment, bandages were placed on porcine dermal excision biopsy wounds. The bandages soaked in oxygen-enriched saline solution or a control group of bandages soaked in a saline solution that was not oxygen-enriched. Microscopically, several factors were evaluated by the study including: 1) epidermalization; 2) neovascularization; 3) epidermal differentiation; 4) mast cell migration; and 5) mitosis.

Externally, the wounds appeared to heal at varying rates. The wounds treated with the oxygen-enriched saline solution showed an increase in wound healing at days 4 through 11. However, both wounds seemed to complete healing at approximately the same time. The study showed that between days 3 and 11, the new epidermis in wounds treated with the oxygen-enriched saline solution migrated at two to four times as fast as the epidermis of the wounds treated with the normal saline solution. The study also showed that between 15 and 22 days, the wound treated by the oxygen-enriched saline solution differentiated at a more rapid rate as evidenced by the earlier formation of more mature epidermal layers. At all stages, the thickening that occurs in

the epidermis associated with normal healing did not occur within the wounds treated by the oxygen-enriched saline solution.

Without wishing to be bound by any particular theory, it is believed that the oxygen-enriched saline solution may increase the localized level of NO within the wounds. NO modulates growth factors, collagen deposition, inflammation, mast cell migration, epidermal thickening, and neovascularization in wound healing. Furthermore, NO is produced by an inducible enzyme that is regulated by oxygen.

Thus, while not wishing to be bound to any particular theory, the inventive gas-enriched fluid may stimulate NO production, which is in accordance with the spectrum of wound-healing effects seen in these experiments.

The epidermis of the healing pigs experienced earlier differentiation in the oxygen-enriched saline group at days 15 through 22. In the case of mast cell migration, differences also occurred in early and late migration for the oxygen-enriched solution. A conclusive result for the level of mitosis was unascertainable due to the difficulty in staining.

Referring now to Figure 41A through 41F, various illustrations compare the wound-healing results of the porcine epidermal tissues with or without oxygen-enriched saline solution. Thus, the healing of the control wound and of the wound using the oxygen-enriched saline solution was followed for days 1, 4 and 16. Figure 41A illustrates the wound healing for the control wound on day 1. As can be seen, the wound shows epidermal/dermal thickening and a loss of contour. Figure 41B illustrates the wound healing on day 1 for the wound treated using the oxygen-enriched saline solution. The wound shows normal epidermal/dermal thickness and normal contouring is typical on a new wound.

Referring now to Figures 41C and 41D, there are illustrated the wound healing for the control wound on day 4 and the wound healing for the wound treated with the oxygen-enriched saline solution on day 4. For the control wound illustrated in Figure 41C, the wound shows a 600 micron epidermal spur. In the wound treated with the oxygen-enriched saline solution in Figure 41D, there is illustrated a 1200 micron epidermal spur. Thus, in the first 4 days of the experiment, the epidermal spur created in the wound treated using the oxygen-enriched saline solution shows an epidermal

growth rate of twice of that of the wound that was not treated with the oxygen-enriched saline solution.

Referring now to Figure 41E, there is illustrated the control wound at day 16. The wound shows less differentiated epidermis with loss of epidermal/dermal contour than that illustrated by the wound treated with the oxygen-enriched saline solution illustrated in Figure 41F. Figure 41F shows more differentiated epidermis and more normal epidermal/dermal contouring in the wound.

Thus, as illustrated with respect to Figures 41A through 41F, the wound treated with the oxygen-enriched saline solution shows much greater healing characteristics than the untreated wound and shows a greater differentiated epidermis with more normal epidermal/dermal contour.

EXAMPLE 9

Glutathione Peroxidase Study

The inventive oxygen-enriched fluid was tested for the presence of hydrogen peroxide by testing the reactivity with glutathione peroxidase using a standard assay (Sigma). Water samples were tested by adding the enzyme cocktail and inverting. Continuous spectrophotometric rate determination was made at A_{340} nm, and room temperature (25 degrees Celsius). Samples tested were: 1) deionized water (negative control), 2) inventive oxygen-enriched fluid at low concentration, 3) inventive oxygen-enriched fluid at high concentration, and 4) hydrogen peroxide (positive control). The hydrogen peroxide positive control showed a strong reactivity, while none of the other fluids tested reacted with the glutathione peroxidase.

EXAMPLE 10

(Electrokinetically-generated superoxygenated fluids and Solas were shown to provide for synergistic prolongation effects (e.g., suppression of bronchoconstriction) with Albuterol in vivo in an art-recognized animal model of human bronchoconstriction (human asthma model))

Experiment 1:

In an initial experiment, sixteen guinea pigs were evaluated for the effects of bronchodilators on airway function in conjunction with methacholine-induced bronchoconstriction. Following determination of optimal dosing, each animal was

dosed with 50 µg/mL to deliver the target dose of 12.5 µg of albuterol sulfate in 250 µL per animal.

The study was a randomized blocked design for weight and baseline PenH values. Two groups (A and B) received an intratracheal instillation of 250 µL of 50 µg/mL albuterol sulfate in one or two diluents: Group A was deionized water that had passed through the inventive device, without the addition of oxygen, while Group B was inventive gas-enriched water. Each group was dosed intratracheally with solutions using a Penn Century Microsprayer. In addition, the animals were stratified across BUXCO plethysmograph units so that each treatment group is represented equally within nebulizers feeding the plethysmographs and the recording units.

Animals that displayed at least 75% of their baseline PenH value at 2 hours following albuterol administration were not included in the data analyses. This exclusion criteria is based on past studies where the failure to observe bronchoprotection with bronchodilators can be associated with dosing errors. As a result, one animal from the control group was dismissed from the data analyses.

Once an animal had greater than 50% bronchoconstriction, the animal was considered to be not protected. As set forth in Table 7 below, 50% of the Group B animals (shaded) were protected from bronchoconstriction out to 10 hours (at which time the test was terminated).

TABLE 7. Bronchoconstriction Protection as Measured with Methacholine Challenge.

Group A

Percent protection from bronchoconstriction by animal number

Time (hours)	1	2	3	4	5	6	7
0	100.00	100.00	100.00	100.00	100.00	100.00	100.00
2	20.81	23.82	32.89	11.56	7.91	24.95	20.15
6	15.67	9.96	8.53	8.40	81.66	75.60	91.97
10	173.92	130.34	95.45	68.14	57.85	103.95	69.03

Group B

Percent protection from bronchoconstriction by animal number

Time (hours)	1	2	3	4	5	6	7	8
0	100.00	100.00	100.00	100.00	100.00	100.00	100.00	100.00
2	15.85	18.03	17.88	24.09	18.59	15.18	21.33	13.33
6	211.57	10.96	68.79	23.72	11.09	99.00	118.26	6.95
10	174.54	12.87	88.16	20.40	21.45	31.60	123.47	8.46

Experiment 2: A Bronchoconstriction Evaluation of RDC1676 With Albuterol Sulfate in Male Hartley Guinea Pigs.

5 An additional set of experiments was conducted using a larger number of animals to evaluate the protective effects of the inventive electrokinetically-generated fluids (e.g, RDC1676-00, RDC1676-01, RDC1676-02 and RDC1676-03) against methacholine-induced bronchoconstriction when administered alone or as diluents for albuterol sulfate in male guinea pigs.

10

Materials:

Guinea Pigs (*Cavia porcellus*) were Hartley albino, CrI:(HA)BR from Charles River Canada Inc. (St. Constant, Quebec, Canada). Weight: Approximately 325 ± 50 g at the onset of treatment. Number of groups was 32, with 7 male animals per group (plus 24 spares form same batch of animals). Diet: All animals had free access to a standard certified pelleted commercial laboratory diet (PMI Certified Guinea Pig 5026; PMI Nutrition International Inc.) except during designated procedures.

15

Methods:

20 Route of administration was intratracheal instillation via a Penn Century Microsprayer and methacholine challenge via whole body inhalation. The intratracheal route was selected to maximize lung exposure to the test article/control solution. Whole body inhalation challenge has been selected for methacholine challenge in order to provoke an upper airway hypersensitivity response (i.e. bronchoconstriction).

25

Duration of treatment was one day.

Table 8 shows the experimental design. All animals were subjected to inhalation exposure of methacholine (500µg/ml), 2 hours following TA/Control administration. All animals received a dose volume of 250µl. Therefore, albuterol sulfate was diluted (in the control article and the four test articles) to concentrations of 0, 25, 50, and 100µg/ml.

5 Thirty minutes prior to dosing, solutions of albuterol sulfate of 4 different concentrations (0, 25, 50, and 100µg/ml) was made up in a 10x stock (500µg/mL) in each of these four test article solutions (RDC1676-00, RDC1676-01, RDC1676-02, and RDC1676-03). These concentrations of albuterol sulfate were also made up in non-electrokinetically-generated control fluid (control 1). The dosing solutions were
10 prepared by making the appropriate dilution of each stock solution. All stock and dosing solutions were maintained on ice once prepared. The dosing was completed within one hour after the test/control articles are made. A solution of methacholine (500µg/ml) was prepared on the day of dosing.

Each animal received an intratracheal instillation of test or control article using a
15 Penn Century microsyringe. Animals were food deprived overnight and were anesthetized using isoflurane, the larynx was visualized with the aid of a laryngoscope (or suitable alternative), and the tip of the microsyringe was inserted into the trachea. A dose volume of 250 µl/animal of test article or control was administered.

The methacholine aerosol was generated into the air inlet of a mixing chamber
20 using aeroneb ultrasonic nebulizers supplied with air from a Buxco bias flow pump. This mixing chamber in turn fed four individual whole body unrestrained plethysmographs, each operated under a slight negative pressure maintained by means of a gate valve located in the exhaust line. A vacuum pump was used to exhaust the inhalation chamber at the required flow rate.

25 Prior to the commencement of the main phase of the study, 12 spare animals were assigned to 3 groups (n=4/group) to determine the maximum exposure period at which animals may be exposed to methacholine to induce a severe but non-fatal acute bronchoconstriction. Four animals were exposed to methacholine (500µg/mL) for 30 seconds and respiratory parameters were measured for up to 10 minutes following
30 commencement of aerosol. Methacholine nebulizer concentration and/or exposure time of aerosolization was adjusted appropriately to induce a severe but non-fatal acute/reversible bronchoconstriction, as characterized by a transient increase in penes.

Once prior to test article administration (Day -1) and again at 2, 6, 10, 14, 18, 22, and 26 hours postdose, animals were placed in the chamber and ventilatory parameters (tidal volume, respiratory rate, derived minute volume) and the enhanced pause Penh were measured for a period of 10 minutes using the Buxco Electronics BioSystem XA system, following commencement of aerosol challenge to methacholine. Once animals were within chambers baseline, values were recorded for 1-minute, following which methacholine, nebulizer concentration of 500ug/mL were aerosolized for 30 seconds, animals were exposed to the aerosol for further 10 minutes during which time ventilatory parameters were continuously assessed. Penh was used as the indicator of bronchoconstriction; Penh is a derived value obtained from peak inspiratory flow, peak expiratory flow and time of expiration. $\text{Penh} = (\text{Peak expiratory flow} / \text{Peak inspiratory flow}) * (\text{Expiratory time} / \text{time to expire 65\% of expiratory volume} - 1)$.

Animals that did not display a severe acute bronchoconstriction during the predose methacholine challenge were replaced. Any animal displaying at least 75% of their baseline Penh value at 2 hours post dose were not included in the data analysis. The respiratory parameters were recorded as 20 second means.

Data considered unphysiological was excluded from further analysis.

Changes in Penh were plotted over a 15 minute period and Penh value was expressed as area under the curve. Numerical data was subjected to calculation of group mean values and standard deviations (as applicable).

TABLE 8. Experimental design; 7 male guinea pigs per group.

Group ID	Albuterol (0 µg/animal)	Albuterol (6/25 µg/animal)	Albuterol (12.5 µg/animal)	Albuterol (25 µg/animal)
1 (control 1) (ambient oxygen)	7 males	7 males	7 males	7 males
5 (RDC1676-00 (Solas)	7 males	7 males	7 males	7 males
6 (RDC1676-01 (20 ppm oxygen)	7 males	7 males	7 males	7 males
7 (RDC1676-02 (40 ppm oxygen)	7 males	7 males	7 males	7 males
8 (RDC1676-03	7 males	7 males	7 males	7 males

Group ID	Albuterol (0 µg/animal)	Albuterol (6/25 µg/animal)	Albuterol (12.5 µg/animal)	Albuterol (25 µg/animal)
(60 ppm oxygen)				

Results:

As shown in Figure 107A-D, in the absence of albuterol, administration of the inventive electrokinetically-generated fluids had no apparent effect on mean percent
5 baseline PenH values, when measured over a 26 hour period.

Surprisingly, however, as shown in Figure 108A-D, administration of albuterol (representative data for the 25 µg albuterol/animal groups are shown) formulated in the inventive electrokinetically-generated fluids (at all oxygen level values tested; ambient (Figure 108-A), 20 ppm (Figure 108-B), 40 ppm (Figure 108-C) and 60 ppm (Figure
10 108-D)) resulted in a striking prolongation of anti-bronchoconstrictive effects of albuterol, compared to control fluid. That is, the methacholine results showed a prolongation of the bronchodilation of albuterol out to at least 26 hours. Figures 108 A-D shows that there were consistent differences at all oxygen levels between RDC1676 and the normal saline control. Combining all 4 RDC1676 fluids, the p value for the overall
15 treatment difference from normal saline was 0.03.

According to particular aspects of the present invention, therefore, the inventive electrokinetically-generated solutions provide for synergistic prolongation effects with albuterol, thus providing for a decrease in a patient's albuterol usage, enabling more efficient cost-effective drug use, fewer side effects, and increasing the period over
20 which a patient may be treated and responsive to treatment with albuterol.

EXAMPLE 11Cytokine Profile

Mixed lymphocytes were obtained from a single healthy human volunteer donor.
25 Buffy coat samples were washed according to standard procedures to remove platelets. Lymphocytes were plated at a concentration of 2×10^6 per plate in RPMI media (+ 50 mm HEPES) diluted with either inventive gas-enriched fluid or distilled water (control). Cells were stimulated with 1 microgram/mL T3 antigen, or 1 microgram/mL phytohemagglutinin (PHA) lectin (pan-T cell activator), or unstimulated (negative

control). Following 24 hour incubation, cells were checked for viability and the supernatants were extracted and frozen.

The supernatants were thawed, centrifuged, and tested for cytokine expression using a XMAP® (Luminex) bead lite protocol and platform. Notably, IFN-gamma level was higher in the inventive gas-enriched culture media with T3 antigen than in the control culture media with T3 antigen, while IL-8 was lower in the inventive gas-enriched culture media with T3 antigen than in the control culture media with T3 antigen. Additionally, IL-6, IL-8, and TNF-alpha levels were lower in the inventive gas-enriched media with PHA, than in the control media with PHA, while IL-1b levels were lower in the inventive gas-enriched fluid with PHA when compared with control media with PHA. In gas-inventive media alone, IFN-gamma levels were higher than in control media.

Two million cells were plated into 6 wells of a 24-well plate in full RPMI + 50 mM Hepes with either inventive oxygen-enriched fluid (water) (wells 1, 3, and 5) or distilled water (2, 4 and 6) (10X RPMI diluted into water to make 1 x). Cells were stimulated with 1 ug/ml T3 antigen (wells 1 and 2) or PHA (wells 3 and 4). Control wells 5 and 6 were not stimulated. After 24 hours, cells were checked for viability and supernatants were collected and frozen. Next, the supernatants were thawed and spun at 8,000 g to pellet. The clarified supernatants were assayed for the cytokines listed using a LUMINEX BEAD LITE protocol and platform. The numerical data is tabulated in Table 9.

TABLE 9.

Sample	IFN	IL-10	IL-12p40	IL-12p70	IL-2	IL-4	IL-5	IL-6	IL-8	IL-1 β	IL-10	TNF α
1	0	0	0	2.85	0	0	7.98	20.3	1350	7.56	11500	15.5
2	0	0	0	3.08	0	0	8	15.2	8940	3.68	4280	7.94
3	0	581	168	3.15	0	0	8	16400	2200	3280	862	13700
4	0	377	56.3	4.22	0	0	8.08	23800	22100	33600	558	16200
5	0	0	0	2.51	0	0	7.99	24	1330	7.33	5900	8.55
6	0	0	0	2.77	0	0	8	5.98	3210	4.68	3330	0

EXAMPLE 12Myelin Oligodendrocyte Glycoprotein (MOG)

5 As set forth in Figure 48, lymphocyte proliferation in response to MOG antigenic peptide was increased when cultured in the presence of the inventive gas-enriched fluid when compared to pressurized, oxygenated fluid (pressure pot) or deionized control fluid. Thus, the inventive gas-enriched fluid amplifies the lymphocyte proliferative response to an antigen to which the cells were previously primed.

10 Myelin oligodendrocyte glycoprotein peptide 35-55 (MOG 35-55) (M-E-V-G-W-Y-R-S-P-F-S-R-O-V-H-L-Y-R-N-G-K) (SEQ ID NO:1; see publication US20080139674, incorporated by reference herein, including for purposes of this SEQ ID NO:1) corresponding to the known mouse sequence was synthesized. Next, 5×10^5 spleen cells were removed from MOG T cell receptor transgenic mice previously immunized
 15 with MOG, and were cultured in 0.2 ml TCM fluid reconstituted with inventive gas-enriched fluid, pressurized oxygenated water (pressure pot water) or with control deionized water. Splenocytes were cultured with MOG p35-55 for 48 or 72 hours, respectively. Cultures were pulsed with 1Ci [3H]-thymidine and harvested 16 hours later. Mean cpm of [3H] thymidine incorporation was calculated for triplicate cultures.

20 Results are shown in Figure 48.

EXAMPLE 13Cytokine Expression

In particular aspects, human mixed lymphocytes were stimulated with T3 antigen
 25 or PHA in electrokinetically-generated oxygen-enriched fluid, or control fluid, and changes in IL-1 β , IL-2, IL-4, IL-5, IL-6, IL-7, IL-8, IL-10, IL-12 (p40), IL-12 (p70), IL-13, IL-17, Eotaxin, IFN- γ , GM-CSF, MIP-1 β , MCP-1, G-CSF, FGFb, VEGF, TNF- α , RANTES, Leptin, TNF- β , TFG- β , and NGF were evaluated. As can be seen from Figure 38, pro-inflammatory cytokines (IL-1 β , TNF- α , IL-6, and GM-CSF), chemokines
 30 (IL-8, MIP-1a, RANTES, and Eotaxin), inflammatory enzymes (iNOS, COX-2, and MMP-9), allergen responses (MHC class II, CD23, B7-1, and B7-2), and Th2 cytokines (IL-4, IL-13, and IL-5) tested were reduced in test fluid versus control fluid. By contrast,

anti-inflammatory cytokines (e.g., IL1R-a, TIMPs) tested were increased in test fluid versus control fluid.

To expand on these data, Applicants used an art-recognized model system involving ovalbumin sensitization, for assessing allergic hypersensitivity reactions. The end points studied were particular cytologic and cellular components of the reaction as well as serologic measurements of protein and LDH. Cytokine analysis was performed, including analysis of Eotaxin, IL-1A, IL-1B, KC, MCP-1, MCP-3, MIP-1A, RANTES, TNF-A, and VCAM.

Briefly, male Brown Norway rats were injected intraperitoneally with 0.5 mL Ovalbumin (OVA) Grade V (A5503-1G, Sigma) in solution (2.0 mg/mL) containing aluminum hydroxide (Al (OH)₃) (200 mg/mL) once each on days 1, 2, and 3. The study was a randomized 2 x 2 factorial arrangement of treatments (4 groups). After a two-week waiting period to allow for an immune reaction to occur, the rats were either exposed or were treated for a week with either RDC1676-00 (sterile saline processed through the Mixing Device), and RDC1676-01 (sterile saline processed through the Mixing Device with additional oxygen added). At the end of the 1 week of treatment for once a day, the 2 groups were broken in half and 50% of the rats in each group received either Saline or OVA challenge by inhalation.

Specifically, 14 days following the initial serialization, 12 rats were exposed to RDC 1676-00 by inhalation for 30 minutes each day for 7 consecutive days. The air flow rate through the system was set at 10 liters/minute. A total of 12 rats were aligned in the pie chamber, with a single port for nebulized material to enter and evenly distribute to the 12 sub-chambers of the AERONEB.

Fifteen days following initial sensitization, 12 rats were exposed to RDC 1676-01 by ultrasonic nebulization for 30 minutes each day for 7 consecutive days. The air flow was also set for 10 liters/minute, using the same nebulizer and chamber. The RDC 1676-00 was nebulized first and the AERONEB chamber thoroughly dried before RDC 1676-01 was nebulized.

Approximately 2 hours after the last nebulization treatment, 6 rats from the RDC 1676-00 group were re-challenged with OVA (1% in saline) delivered by intratracheal instillation using a Penn Century Microsprayer (Model 1A-1B). The other 6 rats from the RDC 1676-00 group were challenged with saline as the control group delivered by

way of intratracheal instillation. The following day, the procedure was repeated with the RDC 1676-01 group.

Twenty-four hours after re-challenge, all rats in each group were euthanized by overdose with sodium pentobarbital. Whole blood samples were collected from the inferior vena-cava and placed into two disparate blood collection tubes: Qiagen PAXgene™ Blood RNA Tube and Qiagen PAXgene™ Blood DNA Tube. Lung organs were processed to obtain bronchoalveolar lavage (BAL) fluid and lung tissue for RT-PCR to assess changes in markers of cytokine expression known to be associated with lung inflammation in this model. A unilateral lavage technique was to be employed in order to preserve the integrity of the 4 lobes on the right side of the lung. The left “large” lobe was lavaged, while the 4 right lobes were tied off and immediately placed into TRIzol™, homogenized, and sent to the lab for further processing.

BAL analysis. Lung lavage was collected and centrifuged for 10 minutes at 4° C at 600-800 g to pellet the cells. The supernatants were transferred to fresh tubes and frozen at -80° C. BAL fluid was separated into two aliquots. The first aliquot was spun down, and the supernatant was snap frozen on crushed dry ice, placed in -80° C, and shipped to the laboratory for further processing. The amount of protein and LDH present indicates the level of blood serum protein (the protein is a serum component that leaks through the membranes when it's challenged as in this experiment) and cell death, respectively. The proprietary test side showed slightly less protein than the control.

The second aliquot of BAL fluid was evaluated for total protein and LDH content, as well as subjected to cytological examination. The treated group showed total cells to be greater than the saline control group. Further, there was an increase in eosinophils in the treated group versus the control group. There were also slightly different polymorphonuclear cells for the treated versus the control side.

Blood analysis. Whole blood was analyzed by transfer of 1.2-2.0 mL blood into a tube, and allowing it to clot for at least 30 minutes. The remaining blood sample (approximately 3.5-5.0 mL) was saved for RNA extraction using TRIzol™ or PAXgene™. Next, the clotted blood sample was centrifuged for 10 minutes at 1200 g at room temperature. The serum (supernatant) was removed and placed into two fresh tubes, and the serum was stored at -80° C.

For RNA extraction utilizing TRI Reagent (TB-126, Molecular Research Center, Inc.), 0.2 mL of whole blood or plasma was added to 0.75 mL of TRI Reagent BD supplemented with 20 μ L of 5N acetic acid per 0.2 mL of whole blood or plasma. Tubes were shaken and stored at -80° C. Utilizing PAXgene™, tubes were incubated for approximately 2 hours at room temperature. Tubes were then placed on their side and stored in the -20° C freezer for 24 hours, and then transferred to -80° C for long term storage.

Luminex analysis. By Luminex platform, a microbead analysis was utilized as a substrate for an antibody-related binding reaction which is read out in luminosity units and can be compared with quantified standards. Each blood sample was run as 2 samples concurrently. The units of measurement are luminosity units and the groups are divided up into OVA-challenged controls, OVA-challenged treatment, and saline challenged treatment with proprietary fluid.

For Agilent gene array data generation, lung tissue was isolated and submerged in TRI Reagent (TR118, Molecular Research Center, Inc.). Briefly, approximately 1 mL of TRI Reagent was added to 50-100 mg of tissue in each tube. The samples were homogenized in TRI Reagent, using glass-Teflon™ or Polytron™ homogenizer. Samples were stored at -80° C.

Blood Samples:

Figures 49-58 show the results of whole blood sample evaluations.

Exemplary Figure 49 shows the basic luminosity data presentation format for the blood sample data. Letters designating the identity of the measured cytokine (in this case KC) are at the top right of each data figure. The data is presented both as data points (upper graph) and bar graphs (lower graph) of the individual samples. In either case, the graphs are divided, from left to right, in four groups. The first 2 groups (RDC1676-00 OVA and RDC1676-01 OVA, respectively) were those that were re-challenged with OVA by inhalation, whereas the last two groups (RDC1676-00 OVA and RDC1676-01 OVA, respectively) were those that were re-challenged with saline control only. Again, the suffix 00 represents saline treatment and suffix 01 represents electrokinetically-generated fluid-treated groups.

Each blood sample was split into two samples and the samples were run concurrently. The units of measure are units of luminosity and the groups, going from

left to right are: OVA-challenged controls, OVA-challenged electrokinetically-generated fluid treatment, followed by saline-challenged saline treatment, and saline-challenged electrokinetically-generated fluid treatment. To facilitate review, both the RDC1676-01 groups are highlighted with gray-shaded backdrops, whereas the control saline treatment groups have unshaded backdrops.

Generally, in comparing the two left groups, while the spread of the RDC1676-01 group data is somewhat greater, particular cytokine levels in the RDC1676-01 group as a whole are less than the samples in the control treated group, typically with about a 30% numerical difference between the two groups. Generally, in comparing the right-most two groups, the RDC1676-01 group has a slightly higher numerical number compared to the RDC1676-00 group.

Figure 50 shows analysis of RANTES (IL-8 super family) in blood sample data according to particular exemplary aspects. Luminosity units for the left-most two groups (the OVA-challenged groups) indicate that generally values in the RDC1676-01 treated group were less than the RDC1676-00 control group as shown by the dot plot in the upper graph portion which again shows a 30-35% differential between the two groups, whereas in the saline-only exposed groups the cytokine level values were roughly the same, or perhaps slightly increased in the RDC1676-01 treated group.

Figure 51 shows analysis of MCP-1 in blood sample data according to particular exemplary aspects. Luminosity units for the left-most two groups (the OVA-challenged groups) indicate that generally values in the RDC1676-01 treated group were less than the RDC1676-00 control group as shown by the dot plot in the upper graph portion, whereas in the saline-only exposed groups the cytokine level values were roughly the same, or perhaps slightly increased in the RDC1676-01 treated group.

Figure 52 shows analysis of TNF alpha in blood sample data according to particular exemplary aspects. Luminosity units for the leftmost two groups (the OVA-challenged groups) indicate that generally values in the RDC1676-01 treated group were less than the RDC1676-00 control group as shown by the dot plot in the upper graph portion, whereas in the saline-only exposed groups the cytokine level values were roughly the same, or perhaps slightly increased in the RDC1676-01 treated group.

Figure 53 shows analysis of MIP-1 alpha in blood sample data according to particular exemplary aspects. Luminosity units for the left-most two groups (the OVA-challenged groups) indicate that generally values in the RDC1676-01 treated group were less than the RDC1676-00 control group as shown by the dot plot in the upper graph portion, whereas in the saline only exposed groups the cytokine level values where roughly the same, or perhaps slightly increased in the RDC1676-01 treated group.

Figure 54 shows analysis of IL-1 alpha in blood sample data according to particular exemplary aspects. Luminosity units for the left-most two groups (the OVA-challenged groups) indicate that generally values in the RDC1676-01 treated group were less than the RDC1676-00 control group as shown by the dot plot in the upper graph portion, whereas in the saline-only exposed groups the cytokine level values where roughly the same, or perhaps slightly increased in the RDC1676-01 treated group.

Figure 55 shows analysis of Vcam in blood sample data according to particular exemplary aspects. Luminosity units for the left-most two groups (the OVA-challenged groups) indicate that generally values in the RDC1676-01 treated group were less than the RDC1676-00 control group as shown by the dot plot in the upper graph portion, whereas in the saline-only exposed groups the cytokine level values where roughly the same, or perhaps slightly increased in the RDC1676-01 treated group.

Figure 56 shows analysis of IL-1 beta in blood sample data according to particular exemplary aspects. Luminosity units for the left-most two groups (the OVA-challenged groups) indicate that generally values in the RDC1676-01 treated group were less than the RDC1676-00 control group as shown by the dot plot in the upper graph portion, whereas in the saline-only exposed groups the cytokine level values where roughly the same, or perhaps slightly increased in the RDC1676-01 treated group.

Figures 57 and 58 show analysis of Eotaxin and MCP-3, respectively, in blood sample data according to particular exemplary aspects. In each case, luminosity units for the left-most two groups (the OVA challenged groups) indicate that generally values in the RDC1676-01 treated group were less than the RDC1676-00 control group as shown by the dot plot in the upper graph portion, whereas in the saline-only exposed

groups the cytokine level values were roughly the same, or perhaps slightly increased in the RDC1676-01 treated group.

Bronchial Lavage Samples:

Figures 59-68 show the corresponding results of BAL fluid sample evaluations.

5 Figure 59 shows analysis of KC in BAL data according to particular exemplary aspects. In this instance the response level, coupled with sampling variability, was inconclusive with respect to a difference between the RDC1676-01 and RDC1676-00-treated groups; that is, KC showed relatively little difference between the two groups, but the units of luminosity were very small.

10 Likewise, Figure 60 shows analysis of RANTES in BAL data according to particular exemplary aspects, and showed marked variability in the RDC1676-01 group with one reading being markedly higher than the others, skewing the results.

Likewise, Figure 61 shows analysis of TNF alpha in BAL data according to particular exemplary aspects, and showing relatively little significance in the way of
15 difference between the RDC1676-01 and RDC1676-00 treated groups.

Figure 62 shows analysis of MCP-1 in BAL data according to particular exemplary aspects, and showing relatively little significance in the way of difference between the RDC1676-01 and RDC1676-00 treated groups.

Figures 63 through 68 show analysis of MIP1-A, IL-1 alpha, Vcam, IL-1 beta,
20 MCP-3, and Eotaxin, respectively, in BAL data according to particular exemplary aspects, and showing relatively little significance in the way of difference between the RDC1676-01 and RDC1676-00 treated groups.

In summary, this standard assay of inflammatory reaction to a known sensitization produced, at least in the blood samples, a marked clinical and serologic
25 affect. Additionally, while significant numbers of control animals were physiologically stressed and nearly dying in the process, none of the RDC1676-01 treated group showed such clinical stress effects. This was reflected then in the circulating levels of cytokines, with approximately 30% differences between the RDC1676-01 treated and the RDC1676-01 treated groups in the OVA-challenged groups. By contrast, there
30 were small and fairly insignificant changes in cytokine, cellular, and serologic profiles between the RDC1676 01-treated and the RDC1676-01 treated groups in the non-OVA

challenged groups, which likely merely represent minimal baseline changes of the fluid itself.

EXAMPLE 14

5 Bradykinin B2 Receptor Affinity Binding

A Bio-Layer Interferometry biosensor, Octet Rapid Extended Detection (RED) (forteBio™) was utilized in order to examine membrane receptor affinity binding of Bradykinin ligand with the Bradykinin B2 receptor. The biosensor system consists of a polished fiber optic embedded into a polypropylene hub with a sensor-specific
10 chemistry at the tip. The biosensor set-up has a layer of molecules attached to the tip of an optic fiber that creates an interference pattern at the detector. Any change in the number of molecules bound causes a measured shift in the pattern of light.

As shown in Figure 69 the Bradykinin B2 membrane receptor was immobilized onto aminopropylsilane (APS) biosensor. The sample plate set up was as designated
15 in Figure 69 and analyzed in Figure 70. Next, the binding of Bradykinin to the immobilized receptor was assessed according to the sample set up as designated in Figure 71. Results of Bradykinin binding are shown in Figure 72. Bradykinin binding to the receptor was further titrated according to the set-up as designated in Figure 73.

As indicated in Figure 74, Bradykinin binding to the B2 receptor was
20 concentration dependent, and binding affinity was increased in the proprietary gas-enriched saline fluid of the instant disclosure compared to normal saline. Stabilization of Bradykinin binding to the B2 receptor is shown in Figure 75.

EXAMPLE 15

25 *(A regulatory T cell assay was used to show effects of the inventive electrokinetically generated fluids in modulation of T cell proliferation and elaboration of cytokines (Il-10) and other proteins (e.g., GITR, Granzyme A, XCL1, pStat, and Foxp3)) in regulatory T cell assays, and of, for example, tryptase in PBMC)*

The ability of particular embodiments disclosed herein to regulate T cells was
30 studied by irradiating antigen-presenting cells, and introducing antigen and T cells. Typically, these stimulated T cells proliferate. However, upon the introduction of regulatory T cells, the usual T cell proliferation is suppressed.

Methods:

Briefly, FITC-conjugated anti-CD25 (ACT-1) antibody used in sorting was purchased from DakoCytomation (Chicago, IL). The other antibodies used were as follows: CD3 (HIT3a for soluble conditions), GITR (PE conjugated), CD4 (Cy-5 and
5 FITC-conjugated), CD25 (APC-conjugated), CD28 (CD28.2 clone), CD127-APC, Granzyme A (PE-conjugated), FoxP3 (BioLegend), Mouse IgG1 (isotype control), and XCL1 antibodies. All antibodies were used according to manufacturer's instructions.

CD4⁺ T cells were isolated from peripheral whole blood with CD4⁺ Rosette Kit (Stemcell Technologies). CD4⁺ T cells were incubated with anti-CD127-APC, anti-
10 CD25-PE and anti-CD4-FITC antibodies. Cells were sorted by flow cytometry using a FACS Aria into CD4⁺CD25^{hi}CD127^{lo}/nTreg and CD4⁺CD25⁻ responder T cells.

Suppression assays were performed in round-bottom 96 well microtiter plates. 3.75 x 10³ CD4⁺CD25^{neg} responder T cells, 3.75 x 10³ autologous T reg, 3.75 x 10⁴ allogeneic-irradiated CD3-depleted PBMC were added as indicated. All wells were
15 supplemented with anti-CD3 (clone HIT3a at 5.0 ug/ml). T cells were cultured for 7 days at 37° C in RPMI 1640 medium supplemented with 10% fetal bovine serum. Sixteen hours before the end of the incubation, 1.0 mCi of ³H-thymidine was added to each well. Plates were harvested using a Tomtec cell harvester and ³H-thymidine incorporation determined using a Perkin Elmer scintillation counter. Antigen-presenting
20 cells (APC) consisted of peripheral blood mononuclear cells (PBMC) depleted of T cells using StemSep human CD3⁺ T cell depletion (StemCell Technologies) followed by 40 Gy of irradiation.

Regulatory T cells were stimulated with anti-CD3 and anti-CD28 conditions and then stained with live/dead red viability dye (Invitrogen), and surface markers CD4,
25 CD25, and CD127. Cells were fixed in the Lyze/Fix PhosFlowTM buffer and permeabilized in denaturing Permbuffer III[®]. Cells were then stained with antibodies against each particular selected molecule.

Statistical analysis was performed using the GraphPad Prism software. Comparisons between two groups were made by using the two-tailed, unpaired
30 Student's t-test. Comparisons between three groups were made by using 1-way ANOVA. P values less than 0.05 were considered significant (two-tailed). Correlation

between two groups were determined to be statistically significant via the Spearman coefficient if the r value was greater than 0.7 or less than -0.7 (two-tailed).

Results:

As indicated in Figure 76, regulatory T cell proliferation was studied by stimulating cells with diesel exhaust particulate matter (PM, from EPA). The x-axis of Figure 76 shows activated autologous CD4⁺ effector T cells (responder cells) as a solid black bar, and regulatory T cells alone in the gray bar (shown for confirmation of anergy) which were mixed at a 1:1 ratio as shown in the white bar. The y-axis shows proliferation as measured by uptake of ³H-thymidine. As shown from left to right along the x-axis, "PM" indicates diesel exhaust derived Particulate Matter, "PM + Rev" indicates PM plus a gas-enriched electrokinetically-generated fluid (Rev) of the instant disclosure, "Solis" indicates an electrokinetically-generated fluid of the instant disclosure and device that is not gas-enriched beyond ambient atmosphere, only (no PM added), "Rev" indicates Rev alone (no PM added) as defined above, "Media" indicates the cell growth media alone control (minus PM; no Rev, no Solis), and "Saline Con" indicates the saline control (minus PM; no Rev, no Solis), "V" indicates verapamil, and "P" indicates propranolol, and "DT" is DT390 at 1:50.

As shown in Figure 77, cells stimulated with PM (no Rev, no Solis) resulted in a decrease in secreted IL-10, while cells exposed to PM in the presence of the fluids of the instant disclosure ("PM + Rev") resulted in a maintained or only slightly decreased production of IL-10 relative to the Saline and Media controls (no PM). Furthermore, Diphtheria toxin (DT390, a truncated diphtheria toxin molecule; 1:50 dilution of std. commercial concentration) was titrated into inventive fluid samples, and blocked the Rev-mediated effect of increase in IL-10 in Figure 77. Note that treatment with Rev alone resulted in higher IL-10 levels relative to Saline and Media controls.

Likewise, similar results, shown in Figures 78-82, were obtained with GITR, Granzyme A, XCL1, pStat, and Foxp3, respectively. In Figures, "NSC" is the same as "Solis" (no PM).

Figure 83 shows AA PBMC data, obtained from an allergic asthma (AA) profile of peripheral blood mononuclear cells (PBMC) evaluating tryptase. The AA PBMC data was consistent with the above T-regulatory cell data, as cells stimulated with PM showed high levels of tryptase, while cells treated with PM in the presence of the fluids

of the instant disclosure ("PM + Rev") resulted in significantly lower tryptase levels similar to those of the Saline and Media controls. Consistent with the data from T-regulatory cells, exposure to DT390 blocked the Rev-mediated effect on tryptase levels, resulting in an elevated level of tryptase in the cells as was seen for PM alone (minus
5 Rev, no Rev, no Solis). Note that treatment with Rev alone resulted in lower tryptase levels relative to Saline and Media controls.

In summary, the data of Figure 76, showing a decreased proliferation in the presence of PM and Rev relative to PM in control fluid (no Rev, no Solis), indicates that the inventive electrokinetically-generated fluid Rev improved regulatory T-cell function
10 as shown by relatively decreased proliferation in the assay. Moreover, the evidence of this Example and Figures 76-83, indicate that beta blockade, GPCR blockade and Ca channel blockade affects the activity of Revera on Treg function.

EXAMPLE 16

15 *(Treatment of primary bronchial epithelial cells (BEC) with the inventive electrokinetically-generated fluids resulted in reduced expression and/or activity of two key proteins of the airway inflammatory pathways, MMP9 and TSLP)*

Overview. As shown in Example 14 above (e.g., Figure 75, showing Stabilization of Bradykinin binding to the B2 receptor using Bio-Layer Interferometry
20 biosensor, Octet Rapid Extended Detection (RED) (forteBio™)), Bradykinin binding to the B2 receptor was concentration dependent, and binding affinity was increased in the electrokinetically-generated fluid (e.g., Rev; gas-enriched electrokinetically-generated fluid) of the instant disclosure compared to normal saline. Additionally, as shown in Example 15 in the context of T-regulatory cells stimulated with diesel exhaust PM, the
25 data showed a decreased proliferation of T-regulatory cells in the presence of PM and Rev relative to PM in control fluid (no Rev, no Solis) (Figure 76), indicating that the inventive electrokinetically-generated fluid Rev improved regulatory T cell function (e.g., as shown by relatively decreased proliferation in the assay). Moreover, exposure to the inventive fluids resulted in a maintained or only slightly decreased production of IL-10
30 relative to the Saline and Media controls (no PM). Likewise, in the context of the AA profiles of PBMC stimulated with PM, the data showed that exposure to the fluids of the instant disclosure ("PM + Rev") resulted in significantly lower tryptase levels similar to

those of the Saline and Media controls. Additionally, the Diphtheria toxin (DT390, a truncated diphtheria toxin molecule; 1:50 dilution of std. commercial concentration) effects shown in Example 15 and Figures 76-83, indicate that beta blockade, GPCR blockade and Ca channel blockade affects the activity of the electrokinetically-generated fluids on Treg and PBMC function. Furthermore, the data of Example 18 shows that, according to additional aspects, upon exposure to the inventive fluids, tight junction-related proteins were upregulated in lung tissue. Figures 85-89 show upregulation of the junction adhesion molecules JAM 2 and 3, GJA1, 3, 4, and 5 (junctional adherins), OCLN (occludin), claudins (e.g., CLDN 3, 5, 7, 8, 9, 10), TJP1 (tight junction protein 1), respectively. Furthermore, as shown in the patch clamp studies of Example 23, the inventive electrokinetically-generated fluids (e.g., RNS-60) affect modulation of whole cell conductance (e.g., under hyperpolarizing conditions) in Bronchial Epithelial Cells (BEC; e. g., Calu-3), and according to additional aspects, modulation of whole cell conductance reflects modulation of ion channels.

In this Example, Applicants have extended these discoveries by conducting additional experiments to measure the effects of production of two key proteins of the airway inflammatory pathways. Specifically, MMP9 and TSLP were assayed in primary bronchial epithelial cells (BEC).

Materials and Methods:

Commercially available primary human BEC (HBEpC-c from Promocell, Germany) were used for these studies. Approximately 50,000 cells were plated in each well of a 12-well plate until they reached ~80% confluence. The cells were then treated for 6 hours with normal saline, control fluid Solas or the test fluid Revera 60 at a 1:10 dilution (100ul in 1ml of airway epithelial growth medium) along with the diesel exhaust particulate matter (DEP or PM) before being lifted for FACS analysis, as described in Example 8 herein. Both MMP9 and TSLP receptor antibodies were obtained from BD Biosciences and used as per manufacturer's specifications.

Results:

In Figures 115 and 116, DEP represents cells exposed to diesel exhaust PM alone, "NS" represents cells exposed to normal saline alone, "DEP+NS" represent cells treated with PM in the presence of normal saline, "Revera 60" refers to cells exposed only to the test material, "DEP + Revera 60" refer to cells treated with PM in the

presence of the test material Revera 60. In addition, "Solas" and "DEP + Solas" represents cells exposed to the control fluid Solas alone or in combination with the PM, respectively.

Figure 115 shows that the test material Revera 60 reduces DEP-induced TSLP receptor expression in BEC by approximately 90%. Solas resulted in a 55% reduction in TSLP receptor expression, while normal saline failed to produce similar level of reduction in TSLP receptor expression (approximately 20% reduction). The effect of the inventive solution in reducing TSLP receptor expression is a significant discovery in view of recent findings showing that TSLP plays a pivotal role in the pathobiology of allergic asthma and local antibody mediated blockade of TSLP receptor function alleviated allergic disease (Liu, YJ, Thymic stromal lymphopoietin: Master switch for allergic inflammation, *J Exp Med* 203:269-273, 2006; Al-Shami et al., A role for TSLP in the development of inflammation in an asthma model, *J Exp Med* 202:829-839, 2005; and Shi et al., Local blockade of TSLP receptor alleviated allergic disease by regulating airway dendritic cells, *Clin Immunol.* 2008, Aug 29. (Epub ahead of print)).

Likewise, Figure 116 shows the effect of Revera 60, Solas and normal saline on the DEP-mediated increase in MMP 9. Specifically, Revera 60 inhibited the DEP-induced cell surface bound MMP9 levels in bronchial epithelial cells by approximately 80%, and Solas had an inhibitory effect of approximately 70%, whereas normal saline (NS) had a marginal effect of about 20% reduction. MMP-9 is one of the major proteinases involved in airway inflammation and bronchial remodeling in asthma. Recently, it has been demonstrated that the levels of MMP-9 are significantly increased in patients with stable asthma and even higher in acute asthmatic patients compared with healthy control subjects. MMP-9 plays a crucial role in the infiltration of airway inflammatory cells and the induction of airway hyperresponsiveness indicating that MMP-9 may have an important role in inducing and maintaining asthma (Vignola et al., Sputum metalloproteinase-9/tissue inhibitor of metalloproteinase-1 ratio correlates with airflow obstruction in asthma and chronic bronchitis, *Am J Respir Crit Care Med* 158:1945-1950, 1998; Hoshino et al., Inhaled corticosteroids decrease subepithelial collagen deposition by modulation of the balance between matrix metalloproteinase-9 and tissue inhibitor of metalloproteinase-1 expression in asthma, *J Allergy Clin Immunol* 104:356-363, 1999; Simpson et al., Differential proteolytic enzyme activity in

eosinophilic and neutrophilic asthma, *Am J Respir Crit Care Med* 172:559–565,2005; Lee et al., A murine model of toluene diisocyanate-induced asthma can be treated with matrix metalloproteinase inhibitor, *J Allergy Clin Immunol* 108:1021–1026, 2001; and Lee et al., Matrix metalloproteinase inhibitor regulates inflammatory cell migration by
5 reducing ICAM-1 and VCAM-1 expression in a murine model of toluene diisocyanate-induced asthma, *J Allergy Clin Immunol* 2003;111:1278–1284).

According to additional aspects, therefore, the inventive electrokinetically-generated fluids have substantial therapeutic utility for modulating (e.g., reducing) TSLP receptor expression and/or for inhibiting expression and/or activity of MMP-9, including,
10 for example, for treatment of inflammation and asthma.

EXAMPLE 17

(The inventive electrokinetically-generated fluids were shown to have a synergistic anti-inflammatory effect with Budesonide in an art-recognized animal model for allergic
15 asthma)

This working Example describes experiments performed to assess the airway anti-inflammatory properties of the inventive electrokinetically-generated fluids (e.g., RDC-1676-03) in a Brown Norway rat ovalbumin sensitization model. The Brown
20 Norway rat is an art-recognized model for determining the effects of a test material on airway function and this strain has been widely used, for example, as a model of allergic asthma. Airway pathology and biochemical changes induced by ovalbumin sensitization in this model resemble those observed in man (Elwood et al., *J Allergy Clin Immuno* 88:951–60, 1991; Sirois & Bissonnette, *Clin Exp Immunol* 126:9–15,
25 2001). The inhaled route was selected to maximize lung exposure to the test material or the control solution. The ovalbumin-sensitized animals were treated with budesonide alone or in combination with the test material RDC 1676-03 for 7 days prior to ovalbumin challenge. At time points of 6 and 24 hours following the challenge, total blood count and levels of several pro and anti-inflammatory cytokines as well as various
30 respiratory parameters were measured to estimate any beneficial effect of administering the test material on various inflammatory parameters.

Materials and Methods:

Brown Norway rats of strain Bn/Crl were obtained from Charles River Kingston, weighing approximately 275 ± 50 g at the onset of the experiment. All animal studies

were conducted with the approval by PCS-MTL Institutional Animal Care and Use Committee. During the study, the use and care of animals were conducted according to guidelines of the USA National Research Council as well as Canadian Council of Animal Care.

5 *Sensitization.* On day 1 of the experiment, animals (14 animals in each treatment group) were sensitized by administration of a 1 ml intraperitoneal injection of a freshly-prepared solution of 2 mg ovalbumin/100mg Aluminum Hydroxide per 1 ml of 0.9% Sodium Chloride, followed by repeat injection on day 3.

10 *Treatment.* Fifteen days following the initial sensitization, animals were subjected to nebulized exposure to control (normal saline) or test solutions (electrokinetically-generated fluids RDC1676-00, RDC1676-02 and RDC-1676-03), either administered alone or in combination with Budesonide, once daily for 15 minutes for 7 consecutive days. Animals were dosed in a whole body chamber of approximately 20L, and test atmosphere was generated into the chamber air inlet using aeroneb
15 ultrasonic nebulizers supplied with air from a Buxco bias flow pump. The airflow rate was set at 10 liters/min.

Ovalbumin challenge. On day 21, 2 hours following treatment with the test solutions, all animals were challenged with 1% ovalbumin nebulized solution for 15 minutes (in a whole body chamber at airflow 2 L/min).

20 *Sample collection.* At time points of 6 and 24 hours after the ovalbumin challenge, blood samples were collected for total and differential blood cell counts as well as for measuring levels of various pro and anti-inflammatory cytokines. In addition, immediately after and at 6 and 24 hours following ovalbumin challenge, the enhanced
25 pause Penh and tidal volume were measured for a period of 10 minutes using the Buxco Electronics BioSystem XA system.

Results:

 Eosinophil Count: As expected, and as shown in Figure 109, treatment with Budesonide ("NS + Budesonide 750 µg/Kg"; densely crosshatched bar graph) reduced the total eosinophil count in the challenged animals relative to treatment with the NS
30 alone control (open bar graph). Additionally, while treatment with the inventive fluid "RDC1676-03" alone (lightly crosshatched bar graph) did not significantly reduce the eosinophil count, it nonetheless displayed a substantial synergy with Budesonide in

reducing the eosinophil count ("RDC1676-03 + Budesonide 750 µg/Kg", solid dark bar graph). Similarly, in Figure 110, the Eosinophil percentage also reflected a similar trend. While RDC1676-03 (lightly crosshatched graph bar) or Budesonide 750 ug/kg (densely crosshatched bar graph) alone did not have a significant effect on Eosinophil percentage count in the challenged animals, the two in combination reduced the Eosinophil percentage significantly (solid dark bar graph).

Therefore, Figure 109 and 110 show, according to particular aspects of the present invention that the inventive electrokinetically-generated fluids (e.g., RDC1676-03) were demonstrated to have a substantial synergistic utility in combination with Budesonide to significantly reduce eosinophil count ("Eosinophil %" and total count) in an art-recognized rat model for human allergic asthma.

Respiratory parameters:

Figures 111 A-C and 112 A-C demonstrate the observed effect of the test fluids on Penh and tidal volume as measured immediately, and at 6 and 24 hours after the ovalbumin challenge. Penh is a derived value obtained from peak inspiratory flow, peak expiratory flow and time of expiration and lowering of penh value reflects a favorable outcome for lung function.

$$\text{Penh} = (\text{Peak expiratory flow/Peak inspiratory flow}) * (\text{Expiratory time/time to expire 65\% of expiratory volume} - 1).$$

As evident from figures 111 A-C, treatment with Budesonide (at both 500 and 750 ug/kg) alone or in combination with any of the test fluids failed to significantly affect the Penh values immediately after the challenge. However, 6 hours after the challenge, animals treated with RDC1676-03 alone or in combination with Budesonide 500 or 750 ug/kg demonstrated a significant drop in Penh values. Although the extent of this drop was diminished by 24 hours post challenge, the trend of a synergistic effect of Budesonide and RDC fluid was still observed at this time point.

Tidal volume is the volume of air drawn into the lungs during inspiration from the end-expiratory position, which leaves the lungs passively during expiration in the course of quiet breathing. As shown in figure 112 A-C, animals treated with Budesonide alone showed no change in tidal volumes immediately after the challenge. However, RDC1676-03 alone had a significant stimulatory effect on tidal volume even at this early time point. And again, RDC1676-03 in combination with Budesonide (both 500 and 750

ug/kg) had an even more-pronounced effect on Tidal volume measurements at this time point. Six hours after the challenge, RDC1676-03 alone was sufficient to cause a significant increase in tidal volume and addition of Budesonide to the treatment regimen either alone or in combination had no added effect on tidal volume. Any effect
5 observed at these earlier time points were, however, lost by the 24 hours time point.

Taken together, these data demonstrate that RDC1676-03 alone or in combination with Budesonide provided significant relief to airway inflammation as evidenced by increase in tidal volume and decrease in Penh values at 6 hours post challenge.

10 Cytokine Analysis:

To analyze the mechanism of the effects seen on the above-discussed physiological parameters, a number of pro- as well as anti-inflammatory cytokines were measured in blood samples collected at 6 and 24 hours after the challenge, immediately following the physiological measurements.

15 Figures 113A and 113B clearly demonstrate that Rev 60 (or RDC1676-03) alone lowered the blood level of eotaxin significantly at both 6 and 24 hours post challenge. Budesonide 750ug/kg also reduced the blood eotaxin levels at both of these time points, while Budesonide 250 ug/kg only had a notable effect at the later time point. However, the test solution Rev 60 alone showed effects that are significantly more
20 potent (in reducing blood eotaxin levels) than both concentrations of Budesonide, at both time points. Eotaxin is a small C-C chemokine known to accumulate in and attract eosinophils to asthmatic lungs and other tissues in allergic reactions (e.g., gut in Crohn's disease). Eotaxin binds to a G protein-coupled receptor CCR3. CCR3 is expressed by a number of cell types such as Th2 lymphocytes, basophils, and mast
25 cells but expression of this receptor by Th2 lymphocyte is of particular interest as these cells regulate eosinophil recruitment. Several studies have demonstrated increased production of eotaxin and CCR3 in asthmatic lung as well as establishing a link between these molecules and airway hyper-responsiveness (reviewed in Eotaxin and the attraction of eosinophils to the asthmatic lung, Dolores M Conroy and Timothy J
30 Williams *Respiratory Research* 2001, 2:150-156). It is of particular interest to note that these studies completely agree with the results in Figures 109 and 110 on eosinophil counts.

Taken together these results strongly indicate that treatment with RDC1676-03 alone or in combination with Budesonide can significantly reduce eosinophil total count and percentage in blood 24 hours after the ovalbumin challenge. This correlates with a significant drop in eotaxin levels in blood observed as early as 6 hours post challenge.

5 Blood levels of two major key anti-inflammatory cytokines, IL10 and Interferon gamma are also significantly enhanced at 6 hours after challenge as a result of treatment with Rev 60 alone or in combination with Budesonide. Figures 113C and 113D show such effects on Interferon gamma and IL 10, respectively. It is evident from these figures that Rev 60 alone or Rev 60 in combination with Budesonide 250 ug/kg
10 significantly increased the blood level of IL10 in the challenged animals up to 6 hrs post challenge. Similarly, Rev 60 alone or in combination with Budesonide 250 or 750 ug/kg significantly increased the blood level of IFN gamma at 6 hours post challenge. Increase in these anti-inflammatory cytokines may well explain, at least in part, the beneficial effects seen on physiological respiratory parameters seen 6 hours post
15 challenge. The effect on these cytokines was no longer observed at 24 hour post challenge (data not shown).

Rantes or CCL5 is a cytokine expressed by circulating T cells and is chemotactic for T cells, eosinophils and basophils and has an active role in recruiting leukocytes into inflammatory sites. Rantes also activates eosinophils to release, for example,
20 eosinophilic cationic protein. It changes the density of eosinophils and makes them hypodense, which is thought to represent a state of generalized cell activation. It also is a potent activator of oxidative metabolism specific for eosinophils.

As shown in Figure 114, systemic levels of Rantes were reduced significantly at 6 hours, but not at 24 hours post challenge in animals treated with Rev 60 alone or in
25 combination of Budesonide 250 or 750 ug/kg. Once again, there is a clear synergistic effect of Budesonide 750 ug/kg and Rev 60 that is noted in this set of data. A similar downward trend was observed for a number of other pro-inflammatory cytokines, such as KC or IL8, MCP3, IL1b, GCSF, TGFb as well as NGF, observed either at 6 or at 24 hours post challenge, in animals treated with Rev 60 alone or in combination with
30 Budesonide.

EXAMPLE 18

(The inventive therapeutic fluids have substantial utility for modulating intercellular tight junctions)

According to particular aspects, the inventive diffuser-processed therapeutic fluids have substantial utility for modulating intercellular tight junctions, including those relating with pulmonary and systemic delivery and bioavailability of polypeptides, including the exemplary polypeptide salmon calcitonin (sCT).

Example Overview. Salmon calcitonin (sCT) is a 32 amino acid peptide with a molecular weight of 3,432 Daltons. Pulmonary delivery of calcitonin has been extensively studied in model systems (*e.g.*, rodent model systems, rat model systems, etc) to investigate methods to enhance pulmonary drug delivery (*e.g.*, intratracheal drug delivery). According to particular exemplary aspects, the inventive diffuser-processed therapeutic fluid has substantial utility for modulating (*e.g.*, enhancing) intercellular tight junctions, for example those associated with pulmonary and systemic delivery and bioavailability of sCT in a rat model system.

Methods:

Intratracheal drug delivery. According to particular embodiments, sCT is formulated in the inventive therapeutic fluid and administered to rats using an intratracheal drug delivery device. In certain aspects, a Penn Century Micro-Sprayer device designed for rodent intratracheal drug delivery is used, allowing for good lung delivery, but as appreciated in the art, with relatively low alveolar deposition resulting in poor systemic bioavailability of peptides. According to particular aspects, this art-recognized model system was used to confirm that the inventive diffuser-processed therapeutic fluid has substantial utility for modulating (*e.g.*, enhancing) intercellular tight junctions, including those associated with pulmonary and systemic delivery and bioavailability of polypeptides.

Animal groups and dosing. In certain aspects, rats are assigned to 1 of 3 groups (*n*=6 per group): a) sterile saline; b) base solution without O₂ enrichment ('base solution'); or c) inventive diffuser processed therapeutic fluid ('inventive enriched based solution'). The inventive enriched-based solution is formed, for example by infusing oxygen in 0.9% saline. Preferably, the base solution comprises about 0.9% saline to minimize the potential for hypo-osmotic disruption of epithelial cells. In certain embodiments, sCT is separately reconstituted in the base solution and the inventive

enriched based solution and the respective solutions are delivered to respective animal groups by intratracheal instillation within 60 minutes (10 µg sCT in 200 µL per animal).

Assays. In particular aspects, blood samples (e.g., 200 µl) are collected and placed into EDTA coated tubes prior to dosing and at 5, 10, 20, 30, 60, 120, and 240 minutes following dosing. Plasma is harvested and stored at = -70° C until assayed for sCT using an ELISA.

For Agilent gene array data generation, lung tissue was isolated and submerged in TRI Reagent (TR118, Molecular Research Center, Inc.). Briefly, approximately 1 mL of TRI Reagent was added to 50-100 mg of tissue in each tube. The samples were homogenized in TRI Reagent, using glass-Teflon™ or Polytron™ homogenizer. Samples were stored at -80° C.

Results:

Enhancement of tight junctions. Figure 84 shows that RDC1676-01 (sterile saline processed through the instant proprietary device with additional oxygen added; gas-enriched electrokinetically-generated fluid (Rev) of the instant disclosure) decreased systemic delivery and bioavailability of sCT. According to particular aspects, the decreased systemic delivery results from decreased adsorption of sCT, most likely resulting from enhancement of pulmonary tight junctions. RDC1676-00 signifies sterile saline processed according to the presently-disclosed methods, but without oxygenation.

Additionally, according to particular aspects, tight junction-related proteins were upregulated in lung tissue. Figures 85-89 show upregulation of the junction adhesion molecules JAM 2 and 3, GJA 1,3,4 and 5 (junctional adherins), OCLN (occludin), claudins (e.g., CLDN 3, 5, 7, 8, 9, 10), TJP1 (tight junction protein 1), respectively.

EXAMPLE 19

(The inventive therapeutic fluids have substantial utility for modulating Nitric Oxide levels)

According to particular aspects, the inventive diffuser-processed therapeutic fluids have substantial utility for modulating nitric oxide levels, and/or related enzymes. Figures 90-94 show data obtained from human foreskin keratinocytes exposed to RDC1676-01 (sterile saline processed through the instant proprietary device with

additional oxygen added; gas-enriched electrokinetically-generated fluid (Rev) of the instant disclosure) showing up-regulation of NOS1 and 3, and Nostrin, NOS3. By contrast, data obtained from rat lung tissue (tissue of above Example entitled "Cytokine Expression") shows down regulation of NOS2 and 3, Nostrin and NOS1AP with Rev (Figures 93, 94).

EXAMPLE 20

(Localized electrokinetic effects (voltage/current) were demonstrated using a specially-designed mixing device comprising insulated rotor and stator features)

In this Example, feature-localized electrokinetic effects (voltage/current) were demonstrated using a specially-designed mixing device comprising insulated rotor and stator features.

Overview. As discussed in detail herein above under "Double Layer Effect" (see also Figures 26 and 28) The mixing device 100 may be configured to create the output material 102 by complex and non-linear fluid dynamic interaction of the first material 110 and the second material 120 with complex, dynamic turbulence providing complex mixing that further favors electrokinetic effects. According to particular aspects, the result of these electrokinetic effects may be present within the output material 102 as charge redistributions and redox reactions, including in the form of solublized electrons that are stabilized within the output material.

In addition to general surface-related double layer effects in the mixing chamber, Applicants additionally reasoned that localized electrokinetic effects may be imparted by virtue of the feature-induced microcavitation and fluid acceleration and deceleration in the vicinity of the features. The studies of this Example were thus performed to further investigate and confirm said additional electrokinetic aspects.

Materials:

A test device similar to the inventive mixing devices described herein was constructed, comprising a stainless steel rotor 12 having two features 18 (disposed at 180 degrees), and a stator 14 with a single feature 16 positioned to be rotationally opposable to the rotor features 18 and stator features 16. Significantly, the rotor and stator features, in each case, are insulated from the respective rotor and stator bodies (Figure 95). The device was machined to provide for a consistent rotor:stator gap 20 of

0.020 inches to conform with the devices disclosed elsewhere herein. There is a rotating contact (not shown) at the end of the rotor shaft (not shown) that provides an electrical path for the rotor surface and for the insulated rotor features. Likewise the stator has a similar insulated feature 16 (Figure 95), wherein the stator inner surface and the insulated stainless steel feature are connected to respective contacts on the stator exterior.

A operational amplifier (OpAmp) circuit (M) 22 is connected between the contacts. The operational amplifier (OpAmp) circuit was constructed to provide for collection of very low voltage measurements by taking advantage of the high input impedance of such amplifiers. The outputs of the OpAmp are fed to the inputs of an oscilloscope (e.g., a battery-powered laptop running an oscilloscope application with a Pico Scope 3000TM).

To eliminate the introduction of any ambient noise (e.g., RF radiation from wireless network signals and from the 60 Hz power line) during testing of the device, a fine copper mesh, RF-shielded compartment (approx. three by four by four feet) was constructed to provide a Faraday cage. This configuration provided for excellent signal to noise ratios during experimental testing, as interfering signals from 60 Hz AC noise (e.g., of approximately 2 volts) and high-frequency RF was reduced well below the signals of interest. Using a battery-powered laptop running an oscilloscope application with a Pico Scope 3000 enabled detection of the 30 mV signals (as in Figure 96) created by the features of the test device. In addition, a variable-speed DC motor was positioned outside the Faraday cage and coupled to the rotatable test device via a non-metallic shaft to effectively isolate the motor noise away from the test device.

Methods:

The OpAmp circuit was used to measure voltage potential between the contacts connecting the stator inner surface 12 and the insulated stator feature 16. With the particular circuit arrangement, only a potential was measured. The rotational speed of the device could be varied between about 700 to about 2800 rpm (with the data of Figure 96 being measured with the device running at about 1800 rpm).

To avoid any extraneous voltage generation due to a pump or peristaltic pump, fluid flow through the device was accomplished using inert nitrogen or air or argon acting on fluid in tanks connected to the device. There was no perceptible voltage

contribution from the flow mechanism, and typically air was used as the pumping force to provide for fluid flow through the device.

Fluid flow rate through the device was about 1 L/min.

5 An initial set of non-rotational experiments was conducted by directing fluid flow through the device chamber but without rotation of the rotor in order to assess the presence of any voltage between the stator body 12 and the isolated feature 16. Separate experiments were conducted for both flow directions.

10 An additional set of rotational experiments was then conducted with the same fluid flow rate, and with the device rotor rotating at various speeds from about 300 to about 1800 rpm. For any given experiment, the flow rate and rotational speed were held constant.

Results:

15 With respect to the non-rotational experiments, with fluid flowing through the device in either direction without any rotor rotation, there was only a barely perceptible voltage (e.g., 1 to 2 mV) between the body of the stator and the insulated feature.

20 With respect to the rotational experiments, and with reference to Figure 96, it can be seen that voltage pulses (potential pulses), temporally correlating (in this case at about 1800 rpm) with rotational alignment of opposing rotor stator features, were measurable with the OpAmp in the operating test device. Moreover, such periodic voltage pulses, correlating with feature alignments, could be observed over a range from about 250 or 300 rpm to about 1800 rpm. Additionally, with or without fluid flow, such voltage pulses were observed in the rotational experiments as long as the cavity/fluid chamber of the device was filled with fluid. According to particular aspects, and without being bound by mechanism, rapid, violent compression (e.g., cavitation),
25 acceleration, and deceleration of fluid flow in the vicinity of the repetitive rotationally-aligned features created the respective local voltage pulses that correlate exactly with the rotational period, providing, at least in part, for electrokinetically-generated fluid according to the present invention. Additional experiments revealed that the amplitude (peak shape and height) of the voltage pulses increased with increasing rotational
30 velocity, being initially observable at about 250 to 300 rpm in this particular test device, and increasing up to at least about 2800 rpm. The magnitude of the violent acceleration and deceleration, etc., of fluid flow in the vicinity of the rotationally-aligned

features would be expected to generally increase with increasing rotational velocity, at least until a maximum was reached reflecting physical limits imposed by the geometry, configuration, and/or flow rate of the device. According to additional aspects, because localized voltage spikes are present, localized current flow (e.g., current pulses) is generated in the vicinity of the features, providing, at least in part, for electrokinetically-generated fluid according to the present invention (e.g., without being bound by mechanism, providing for electrochemical reactions as discussed elsewhere herein).

According to additional aspects, and without being bound by mechanism, such feature-localized effects (e.g., voltage pulses and current and/or current pulses) contribute to generation of the electrokinetically-generated fluids in combination with more general surface-related double layer and streaming current effects discussed elsewhere herein above under "Double Layer Effect" (see also Figures 26 and 28).

EXAMPLE 21

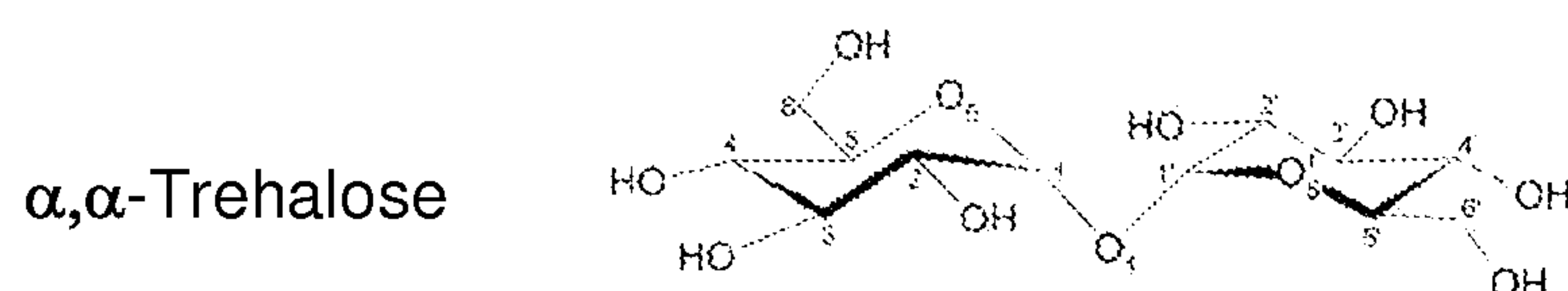
(Relative to non-electrokinetically-generated control fluids, the inventive electrokinetically-generated fluids were shown to differentially affect line widths in ^{13}C NMR analysis of the dissolved solute α,α -Trehalose)

Overview. Applicants' data disclosed elsewhere herein support utility and mechanism wherein the inventive electrokinetically-generated fluids mediate regulation or modulation of intracellular signal transduction by modulation of at least one of cellular membranes, membrane potential/conductance, membrane proteins (e.g., membrane receptors such as G protein coupled receptors), calcium-dependant cellular signaling systems, and intercellular junctions (e.g., tight junctions, gap junctions, zona adherins, and desmosomes). Specifically, using a variety of art-recognized biological test systems and assays, Applicants' data shows, relative to control fluids, differential effects of the inventive fluid on, for example: regulatory T cell proliferation; cytokine and protein levels (e.g, IL-10, GITR, Granzyme A, XCL1, pStat5, and Foxp3, typtase, tight junction-related proteins, TSLP receptor, MMP9, etc.); binding of Bradykinin ligand with the Bradykinin B2 receptor; expression of TSLP receptor, whole cell conductance; etc. Moreover, the Diphtheria toxin (DT390) effects shown herein indicate that beta blockade (beta 2 adrenergic receptor), and/or GPCR blockade and/or Ca channel blockade affects the activity of the electrokinetically-generated fluids on, for example, Treg and PBMC function.

Taken together these effects indicate that the inventive electrokinetically-generated fluids are not only fundamentally distinguished from prior art fluids, but also that they provide for novel compositions and substantial utilities such as those presently disclosed and claimed herein.

5 *In this Example.* Applicants have in this Example performed nuclear magnetic resonance (NMR) studies to further characterize the fundamental nature of the inventive electrokinetically-generated fluids. Specifically, Applicants have analyzed the ^{13}C NMR spectra of α,α -Trehalose dissolved in the electrokinetically-generated fluid, compared to dissolution in non-electrokinetically-generated fluid. Trehalose (shown
10 below with carbons numbered for reference) is a cosmotrophic solute and is known, for example, to protect against protein denaturation, membrane desiccation, organism viability upon freezing, etc. Applicants, given the data summarized above, reasoned that α,α -Trehalose might provide an effective tool to further probe the properties/structure of the inventive electrokinetically-generated fluids. Applicants
15 reasoned that NMR-related ‘chemical shifts’ and effects on ‘line widths’ could be used to assess properties of the inventive fluids. For these studies, a non-superoxygenated inventive electrokinetically-generated fluid (referred to herein as “Solas”) was employed to minimize the possibility that paramagnetic impurities, such as dissolved oxygen, might act to counter or otherwise mask the effects being analyzed.

20



Materials and Methods:

Solution Preparation. The Phosphate (sodium salt) and D-(+)-Trehalose dihydrate (T9531-10G, reduced metal content) and 99.9% D₂O containing 1% DSS
25 were purchased from Sigma. The “Normal Saline” is 0.9% Sodium Chloride, pH 5.6 (4.5-7.0), from Hospira. The 0.25 M α,α -Trehalose solutions were prepared by dissolving 0.949 g Trehalose into 965 μL NS and 35 mL Phosphate Buffered Saline (100mM Phosphate Buffer in 0.9% NaCl prepared in such a way that when 35 μL of this buffer are added to 1.0 mL Trehalose solution the pH becomes 6.93).

Nuclear Magnetic Resonance Spectra Collection. Spectra were collected at the University of Washington NMR facility using either an 500 MHz or 300 MHz Bruker Avance series instrument fitted with a Bruker BBO : X {1H} probe and running XWINNMR 3.5. ¹³C NMR spectra were collected at 125.7 MHz or 75.46 MHz using a 14000 Hz or 7900 Hz sweep width using 64K or 128K data points and 128 or 256 scans. The resulting FIDs were zero-filled twice and processed with a 1.0 Hz line broadening factor. Temperature was controlled using the Bruker Biospin Variable Temperature unit. External deuterium locking was employed by placing 99.9% D₂O + 1% DSS + a trace of acetone in a coaxial NMR insert tube, purchased from Wilmad. The NMR data was processed using the iNMR software v. 2.6.4 from Mestrelab Research.

Results:

Sample Spectra. FIGURE 97A-C shows expansions of six ¹³C-NMR spectra overlaid on top of each other such that the DSS signals line up at -2.04 ppm. The DSS signals are shown at the far right of the figure, and the acetone methyl signal is shown near 30.9 ppm. The remaining signals correspond to the 6 carbons of Trehalose as shown in the α,α -Trehalose structure above. As can be seen, the carbon signals in the Solas solutions show small chemical shifts (generally upfield) compared to the control solutions.

Line Width Measurements. TABLE 10 below shows the measured ¹³C NMR line widths for the six carbons of trehalose and the methyl carbon of acetone at 3 different temperatures for Solas Saline (an inventive electrokinetically-generated fluid). The corresponding NS samples represent non-electrokinetic control solutions at each temperature. In the Solas solutions, the line widths are significantly different from the line widths in the control solution for each carbon atom. The smaller linewidths in the Solas solutions at lower temperatures likely result from a faster tumbling rate of the trehalose molecule as a whole (including any solvated water molecules) compared to the control solutions.

TABLE 10. ¹³C NMR Line Widths for α,α -Trehalose in Solas & Normal Saline^{a,b}

Test Fluid (Temp. degrees K)	C-1	C-2	C-3	C-4	C-5	C-6	Acetone
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PCT/US2010/032620

Solas (277)	8.4	8.22	8.3	8.15	8.3	11.1	5.1
Normal (269.9)	15.4	16.1	15.8	14.9	15.4	21.7	5.1
Solas (293)	9.52	8.7	9.28	9	8.9	11.25	5.63
Normal (292.9)	10.33	10.23	10.23	9.93	10.23	13.13	5.63
Solas (310)	2.28	2.03	2.18	2.19	2	2.55	0.67
Normal (309.9)	1.17	0.99	1.1	1.02	0.97	1.42	0.67

^a1.0 Hz was subtracted from all line width values due to the 1.0 Hz line broadening used during processing. In addition, line width values were normalized relative to the acetone signal in the external reference tube in order to compensate for magnetic field inhomogeneities. This was done by subtracting from the Normal Saline line widths the amount by which the acetone peak was broadened in the corresponding Solas Saline spectra.

^bError in line width measurements estimated to be within +/-0.30 Hz

The ¹³C NMR line widths for α,α -Trehalose in Solas and normal saline, in each case normalized with respect to the Acetone line, are shown graphically in Figure 97A. In conclusion, the NMR data for ¹³C NMR line widths for α,α -Trehalose in Solas and normal saline indicate that there is a property of the inventive solution which alters solute tumbling.

Taken together with the biological activities summarize above and elsewhere herein, these ¹³C NMR line width effects indicate that the inventive electrokinetically-generated fluids are not only fundamentally distinguished from prior art fluids in terms of solute interactions, but also that they provide for novel compositions and substantial utilities such as those presently disclosed and claimed herein.

EXAMPLE 22

(Relative to non-electrokinetically-generated control fluids, the inventive electrokinetically-generated fluids produced differential square wave voltammetry profiles and displayed unique electrochemical properties under stripping polarography)

Overview. Applicants' data disclosed elsewhere herein support utility and mechanism wherein the inventive electrokinetically-generated fluids mediate regulation or modulation of intracellular signal transduction by modulation of at least one of cellular membranes, membrane potential/conductance, membrane proteins (e.g., membrane receptors such as G-Protein-Coupled receptors), calcium-dependant cellular signaling systems, and intercellular junctions (e.g., tight junctions, gap junctions, zona adherins, and desmosomes). Specifically, using a variety of art-recognized biological test systems and assays. Applicants' data shows, relative to control fluids, differential

effects of the inventive fluid on, for example: regulatory T cell proliferation, cytokine and protein levels (e.g, IL-10, GITR, Granzyme A, XCL1, pStat5, and Foxp3, typtase, tight junction-related proteins, TSLP receptor, MMP9, etc.), binding of Bradykinin ligand with the Bradykinin B2 receptor, expression of TSLP receptor and whole cell conductance; etc. Moreover, the Diphtheria toxin (DT390) effects shown herein indicate that beta blockade (beta 2 adrenergic receptor), and/or GPCR blockade and/or Ca channel blockade affects the activity of the electrokinetically-generated fluids on, for example, Treg and PBMC function.

Taken together these effects indicate that the inventive electrokinetically-generated fluids are not only fundamentally distinguished from prior art fluids, but also that they provide for novel compositions and substantial utilities such as those presently disclosed and claimed herein.

In this Example. Applicants have, in this Example, performed voltametry studies to further characterize the fundamental nature of the inventive electrokinetically-generated fluids. Voltametry is frequently used to determine the redox potential or measure kinetic rates and constants of fluids. The common characteristic of all voltametric methods is that they involve the application of a potential to an electrode and the resultant current flowing is monitored through an electrochemical cell. The applied potential produces a change in the concentration of an electroactive species at the electrode surface by electrochemically reducing or oxidizing the species.

Specifically, Applicants have utilized voltametric methods (i.e., square wave voltametry and stripping polarography) to further characterize fundamental differences between control saline fluid and the inventive electrokinetically-generated test fluids (e.g., Solas and Revera). Applicants, given the biological and membrane effects data summarized above, reasoned that square wave voltametry and stripping polarography would provide an effective means to further characterize the unique properties of the inventive electrokinetically-generated fluids.

Applicants further reasoned that differences in current at specific voltages, production of different concentrations of an electroactive redox compound, creation of new redox compounds, and possession of unique electrochemical properties could be used to assess and characterize properties of the inventive fluids. For these studies,

both a superoxygenated electrokinetically-generated fluid (Revera), and a non-super-oxygenated inventive electrokinetically-generated fluid (Solas) were used.

Materials and Methods:

Materials and Solution Preparation. The experiments were conducted on an EG & G SMDE 303A polarographer (Princeton Applied Research). The electrolyte, NaOH, used in the square wave voltammetry experiment, was purchased from Sigma. A 10 mL sample of the inventive fluid solution was prepared by adding 100 μ L of NaOH to 9.9 mL of Revera Saline to make a 0.18 molar solution. With regards to the stripping polarography experiment, no extra electrolyte was utilized.

Square Wave Voltammetry. As stated above, voltammetry is used to determine the redox potential or measure kinetic rates and constants in fluids. In the square wave voltammetry experiment, a potential of 0.0 to approximately -1.75 V was applied to an electrode and the resultant current flowing through the electrochemical cell was monitored.

Stripping Polarography. The stripping polarography method is similar to the square wave voltammetry method. However, no electrolyte was utilized as stated above and also involved a pre-step. In the pre-step, the static mercury drop electrode was held for 30 seconds at -1.1 V to amalgamate any compounds whose reduced form was soluble in mercury. Then, the potentials between -1.1 V and 0.0 V were scanned and the resultant current flowing through the electrochemical cell was monitored. A linear scan into the negative potentials on this amalgam provided a sensitive measurement of these compounds.

Results:

Square Wave Voltammetry. As evident from figure 98, the current profiles at -0.14V, -.47V, -1.02V and -1.36V differ between the various tested agents. According to particular aspects, the differences in current generated at the various specific voltages indicate at least one of a different concentration of an electroactive redox compound and/or a new or unique electroactive redox compound, and/or a change in the diffusion-limiting electrical double layer surrounding the mercury drop.

Stripping Polarography. Figure 99 shows that the inventive electrokinetically-generated fluids, Revera and Solas, show unique spectra with pronounced peaks at -0.9 volts that are not present in the non-electrokinetically-generated blank and saline

control fluids. Additionally, the spectra of the non-electrokinetically-generated blank and saline control fluids show characteristic peaks at -0.19 and -0.3 volts that are absent in the spectra for the electrokinetically-generated Solas and Revera fluids.

According to particular aspects, therefore, these results show unique
5 electrochemical properties of the inventive electrokinetically-generated Solas and Revera fluids compared to non-electrokinetically-generated Saline control fluid. According to additional aspects, the results indicate the presence or generation of at least one of a different concentration of an electroactive redox compound and a new and/or unique electroactive redox compound in electrokinetically-generated versus non-
10 electrokinetically-generated fluids.

On top of the various biological data presented elsewhere herein, this differential voltametry data, particularly when considered along with the differential effects on whole cell conductance, ^{13}C NMR line-width analysis, and the mixing device feature-localized effects (e.g., voltage pulses and current and/or current pulses) indicate that
15 the inventive electrokinetically-generated fluids are not only fundamentally distinguished from prior art fluids, but also provide for novel compositions and substantial utilities such as those presently disclosed and claimed herein.

EXAMPLE 23

20 *(Patch clamp analysis conducted on bronchial epithelial cells (BEC) perfused with inventive electrokinetically-generated fluid (RNS-60) revealed that exposure to RNS-60 resulted in a decrease in whole cell conductance, and stimulation with a cAMP stimulating "cocktail", which dramatically increased the whole-cell conductance. also increased the drug-sensitive portion of the whole-cell conductance, which was ten-*
25 *times higher than that observed under basal conditions)*

In this Example, patch clamp studies were performed to further confirm the utility of the inventive electrokinetically-generated fluids to modulate intracellular signal transduction by modulation of at least one of membrane structure, membrane potential
30 or membrane conductivity, membrane proteins or receptors, ion channels, and calcium dependant cellular messaging systems.

Overview. As shown in Example 14 above (e.g., Figure 75, showing Stabilization of Bradykinin binding to the B2 receptor using Bio-Layer Interferometry biosensor, Octet Rapid Extended Detection (RED) (forteBioTM)), Bradykinin binding to
35 the B2 receptor was concentration-dependent, and binding affinity was increased in the

electrokinetically-generated fluid (e.g., Rev; gas-enriched electrokinetically-generated fluid) of the instant disclosure compared to normal saline. Additionally, as shown in Example 15 in the context of T-regulatory cells stimulated with PM, the data showed a decreased proliferation of T-regulatory cells in the presence of PM and Rev relative to PM in control fluid (no Rev, no Solis) (Figure 76), indicating that the inventive electrokinetically-generated fluid Rev improved regulatory T cell function; e.g., as shown by relatively decreased proliferation in the assay. Moreover, exposure to the inventive fluids resulted in a maintained or only slightly decreased production of IL-10 relative to the Saline and Media controls (no PM). Likewise, in the context of the AA profiles of PBMC stimulated with PM, the data showed that exposure to the fluids of the instant disclosure ("PM + Rev") resulted in significantly lower tryptase levels similar to those of the Saline and Media controls. Additionally, the Diphtheria toxin (DT390) effects shown in Example 15 and figures 76-83, indicate that beta blockade, GPCR blockade and Ca channel blockade affects the activity of the electrokinetically-generated fluids on Treg and PBMC function. Furthermore, the data of Example 18 shows that, according to additional aspects, upon expose to the inventive fluids, tight junction-related proteins were upregulated in lung tissue. Figures 85-89 show upregulation of the junction adhesion molecules JAM 2 and 3, GJA1, 3, 4 and 5 (junctional adherins), OCLN (occludin), claudins (e.g., CLDN 3, 5, 7, 8, 9, 10), TJP1 (tight junction protein 1), respectively.

Patch clamp studies were performed to further investigate and confirm said utilities.

Materials and Methods:

The Bronchial Epithelial line Calu-3 was used in Patch clamp studies. Calu-3 Bronchial Epithelial cells (ATCC #HTB-55) were grown in a 1:1 mixture of Ham's F12 and DMEM medium that was supplemented with 10% FBS onto glass coverslips until the time of the experiments. In brief, a whole cell voltage clamp device was used to measure effects on Calu-3 cells exposed to the inventive electrokinetically-generated fluids (e.g., RNS-60; electrokinetically-treated normal saline comprising 60 ppm dissolved oxygen; sometimes referred to as "drug" in this Example).

Patch clamping techniques were utilized to assess the effects of the test material (RNS-60) on epithelial cell membrane polarity and ion channel activity. Specifically,

whole cell voltage clamp was performed upon the Bronchial Epithelial line Calu-3 in a bathing solution consisting of: 135mM NaCl, 5mM KCl, 1.2mM CaCl₂, 0.8mM MgCl₂, and 10mM HEPES (pH adjusted to 7.4 with N-methyl D-Glucamine). Basal currents were measured after which RNS-60 was perfused onto the cells.

5 More specifically, patch pipettes were pulled from borosilicate glass (Garner Glass Co, Claremont, CA) with a two-stage Narishige PB-7 vertical puller and then fire-polished to a resistance between 6-12 Mohms with a Narishige MF-9 microforge (Narishige International USA, East Meadow, NY). The pipettes were filled with an intracellular solution containing (in mM): 135 KCl, 10 NaCl, 5 EGTA, 10 Hepes, pH was
10 adjusted to 7.4 with NMDG (N-Methyl-D-Glucamine).

The cultured Calu-3 cells were placed in a chamber containing the following extracellular solution (in mM): 135 NaCl, 5 KCl, 1.2 CaCl₂, 0.5 MgCl₂ and 10 Hepes (free acid), pH was adjusted to 7.4 with NMDG.

Cells were viewed using the 40X DIC objective of an Olympus IX71 microscope
15 (Olympus Inc., Tokyo, Japan). After a cell-attached gigaseal was established, a gentle suction was applied to break in, and to attain the whole-cell configuration. Immediately upon breaking in, the cell was voltage clamped at -120, -60, -40 and 0 mV, and was stimulated with voltage steps between ± 100 mV (500 ms/step). After collecting the whole-cell currents at the control condition, the same cell was perfused through bath
20 with the test fluid comprising same extracellular solutes and pH as for the above control fluid, and whole-cell currents at different holding potentials were recorded with the same protocols.

Electrophysiological data were acquired with an Axon Patch 200B amplifier, low-pass filtered at 10 kHz, and digitized with 1400A Digidata (Axon Instruments, Union
25 City, CA). The pCLAMP 10.0 software (Axon Instruments) was used to acquire and to analyze the data. Current (I)-to-voltage (V) relationships (whole cell conductance) were obtained by plotting the actual current value at approximately 400 msec into the step, versus the holding potential (V). The slope of the I/V relationship is the whole cell conductance.

30 *Drugs and Chemicals.* Whenever indicated, cells were stimulated with a cAMP stimulatory cocktail containing 8-Br-cAMP (500 mM), IBMX (isobutyl-1-methylxanthine, 200 mM) and forskolin (10 mM). The cAMP analog 8-Br-cAMP (Sigma Chem. Co.) was

used from a 25 mM stock in H₂O solution. Forskolin (Sigma) and IBMX (Sigma) were used from a DMSO solution containing both 10 mM Forskolin and 200 mM IBMX stock solution.

Patch Clamp Results:

5 Figure 100 shows whole-cell currents under basal (no cAMP) conditions, with a protocol stepping from zero mV holding potential to ± 100 mV. Representative tracings are the average of $n=12$ cells. The tracings on the left are the control, followed by the whole-cell tracings while perfusing the test solution (middle). The tracings on the right are the composite delta obtained by subtraction of the test average values, from those
10 under control conditions. The whole-cell conductance, obtained from the current-to-voltage relationships is highly linear under both conditions, and reflects a modest, albeit significant, change in conductance due to the test conditions. The contribution to the whole-cell conductance, i.e., the component inhibited by the drug (inventive electrokinetically-generated fluid) is also linear, and the reversal potential is near zero
15 mV. There is a decrease in the whole cell conductance under hyperpolarizing conditions.

 Figure 101 shows whole-cell currents under basal conditions, with a protocol stepping from -40 mV holding potential to ± 100 mV. Representative tracings are the average of $n=12$ cells. The tracings on the left are the control, followed by the whole-
20 cell tracings while perfusing the test solution (middle). The tracings on the right are the composite delta obtained by subtraction of the test average values, from those under control conditions. The whole-cell conductance obtained from the current-to-voltage relationships is highly linear under both conditions, and reflects a modest, albeit significant change in conductance due to the test conditions. The contribution to the
25 whole-cell conductance, i.e., the component inhibited by the drug (inventive electrokinetically-generated fluid) is also linear, and the reversal potential is near zero mV. Values are comparatively similar to those obtained with the zero mV protocol.

 Figure 102 shows whole-cell currents under basal conditions, with a protocol stepping from -60 mV holding potential to ± 100 mV. Representative tracings are the
30 average of $n=12$ cells. The tracings on the left are the control, followed by the whole-cell tracings while perfusing the test solution (middle). The tracings on the right are the composite delta obtained by subtraction of the test average values, from those under

control conditions. The whole-cell conductance obtained from the current-to-voltage relationships is highly linear under both conditions, and reflects a minor, albeit significant, change in conductance due to the test conditions. The contribution to the whole-cell conductance, i.e., the component inhibited by the drug is also linear, and the reversal potential is near zero mV. Values are comparatively similar to those obtained with the zero mV protocol.

Figure 103 shows whole-cell currents under basal conditions, with a protocol stepping from -120 mV holding potential to ± 100 mV. Representative tracings are the average of $n=12$ cells. The tracings on the left are the control, followed by the whole-cell tracings while perfusing the test solution (middle). The tracings on the right are the composite delta obtained by subtraction of the test average values, from those under control conditions. The whole-cell conductance obtained from the current -to-voltage relationships is highly linear under both conditions, and reflects a minor, albeit significant change in conductance due to the test conditions. The contribution to the whole-cell conductance, i.e., the component inhibited by the drug is also linear, and the reversal potential is near zero mV. Values are comparatively similar to those obtained with the zero mV protocol.

Figure 104 shows whole-cell currents under cAMP-stimulated conditions, obtained with protocols stepping from various holding potentials to ± 100 mV. Representative tracings are the average of $n=5$ cells. The tracings on the left are the control, followed by the whole-cell tracings after cAMP stimulation, followed by perfusion with the drug-containing solution. The tracings on the right are the composite delta obtained by subtraction of the test average values in drug + cAMP, from those under control conditions (cAMP alone). The tracings on the Top are those obtained from voltage protocol at zero mV, and the ones below, at -40 mV. The whole-cell conductance obtained from the current-to-voltage relationships is highly linear under all conditions, and reflects a change in conductance due to the test conditions.

Figure 105 shows whole-cell currents under cAMP-stimulated conditions, obtained with protocols stepping from various holding potentials to ± 100 mV. Representative tracings are the average of $n=5$ cells. The tracings on the left are the control, followed by the whole-cell tracings after cAMP stimulation, followed by perfusion with the drug-containing solution. The tracings on the right are the composite

delta obtained by subtraction of the test average values in drug + cAMP, from those under control conditions (cAMP alone). The tracings on the Top are those obtained from voltage protocol at -60 mV, and the ones below, at -120 mV. The whole-cell conductance, obtained from the current-to-voltage relationships, is highly linear under all conditions, and reflects a change in conductance due to the test conditions.

Figure 106 shows the effect of holding potential on cAMP-activated currents. The effect of the drug (the inventive electrokinetically-generated fluids; RNS-60; electrokinetically-treated normal saline comprising 60 ppm dissolved oxygen) on the whole-cell conductance was observed under different voltage protocols (0, -40, -60, -120 mV holding potentials). Under basal conditions, the drug-sensitive whole-cell current was identical at all holding potentials (voltage-insensitive contribution, top left panel). In the cAMP-activated conditions, however, the drug-sensitive currents were much higher, and sensitive to the applied voltage protocol. The current-to-voltage relationships are highly nonlinear. This is further observed in the subtracted currents (bottom panel), where the contribution of the whole cell conductance at zero mV was further subtracted for each protocol (n=5).

Summary of Example. According to particular aspects, therefore, the data indicate that there is a modest but consistent effect of the drug (the inventive electrokinetically-generated fluids; RNS-60; electrokinetically-treated normal saline comprising 60 ppm dissolved oxygen) under basal conditions. To enhance the effect of the drug on the whole-cell conductance, experiments were also conducted by perfusing the drug after stimulation with a cAMP stimulating "cocktail," which dramatically increased the whole-cell conductance. Interestingly, this protocol also increased the drug-sensitive portion of the whole-cell conductance, which was *ten-times* higher than that observed under basal conditions. Additionally, in the presence of cAMP stimulation, the drug showed different effects with respect to the various voltage protocols, indicating that *the electrokinetically-generated fluids affect a voltage-dependent contribution of the whole-cell conductance*. There was also a decrease in a linear component of the conductance, further suggesting at least a contribution of the drug to the inhibition of another pathway (e.g., ion channel, voltage gated cation channels, etc.).

In particular aspects, and without being bound by mechanism, Applicants' data are consistent with the inventive electrokinetically-generated fluids (e.g., RNS-60; electrokinetically-treated normal saline comprising 60 ppm dissolved oxygen) producing a change either on a channel(s), being blocked or retrieved from the plasma
5 membrane.

Taken together with Applicants' other data (e.g., the data of working Examples) particular aspects of the present invention provide compositions and methods for modulating intracellular signal transduction, including modulation of at least one of membrane structure, membrane potential or membrane conductivity, membrane
10 proteins or receptors, ion channels, and calcium dependant cellular signaling systems, comprising use of the inventive electrokinetically-generated solutions to impart electrochemical and/or conformational changes in membranous structures (e.g., membrane and/or membrane proteins, receptors or other components) including but not limited to GPCRs and/or g-proteins. According to additional aspects, these effects
15 modulate gene expression, and may persist, dependant, for example, on the half lives of the individual messaging components, etc.

EXAMPLE 24

*(Patch clamp analysis conducted on Calu-3 cells perfused with inventive electrokinetically generated fluids (RNS-60 and Solas) revealed that (i) exposure to
20 RNS-60 and Solas resulted in increases in whole cell conductance, (ii) that exposure of cells to the RNS-60 produced an increase in a non-linear conductance, evident at 15 min incubation times, and (iii) that exposure of cells to the RNS-60 produced an effect of RNS-60 saline on calcium permeable channels)*

Overview. In this Example, patch clamp studies were performed to further
25 confirm the utilities, as described herein, of the inventive electrokinetically generated saline fluids (RNS-60 and Solas), including the utility to modulate whole-cell currents. Two sets of experiments were conducted.

The summary of the data of the *first* set of experiments indicates that the whole
30 cell conductance (current-to-voltage relationship) obtained with Solas saline is highly linear for both incubation times (15 min, 2 hours), and for all voltage protocols. It is however evident, that longer incubation (2 hours) with Solas increased the whole cell conductance. Exposure of cells to the RNS-60 produced an increase in a non-linear conductance, as shown in the delta currents (Rev-Sol subtraction), which is only

evident at 15 min incubation time. The effect of the RNS-60 on this non-linear current disappears, and is instead highly linear at the two-hour incubation time. The contribution of the non-linear whole cell conductance, as previously observed, was voltage sensitive, although present at all voltage protocols.

5 The summary of data of the *second* set of experiments indicates that there is an effect of the RNS-60 saline on a non-linear current, which was made evident in high calcium in the external solution. The contribution of the non-linear whole cell conductance, although voltage sensitive, was present in both voltage protocols, and indicates an effect of RNS-60 saline on calcium permeable channels.

10 First set of experiments (increase of conductance; and activation of a non-linear voltage regulated conductance)

Methods for first set of experiments:

See EXAMPLE 23 for general patch clamp methods. In the following first set of experiments, patch clamp studies were performed to further confirm the utility of the inventive electrokinetically generated saline fluids (RNS-60 and Solas) to modulate whole-cell currents, using Calu-3 cells under basal conditions, with protocols stepping from either zero mV holding potential, -120 mV, or -60 mV.

The whole-cell conductance in each case was obtained from the current-to-voltage relationships obtained from cells incubated for either 15 min or two hours, to further confirm the results of EXAMPLE 23. In this study, groups were obtained at a given time, for either Solas or RNS-60 saline solutions. The data obtained are expressed as the mean $\pm$ SEM whole cell current for 5-9 cells.

Results:

Figures 117 A-C show the results of a series of patch clamping experiments that assessed the effects of the electrokinetically generated fluid (e.g., RNS-60 and Solas) on epithelial cell membrane polarity and ion channel activity at two time-points (15 min (left panels) and 2 hours (right panels)) and at different voltage protocols (A, stepping from zero mV; B, stepping from -60 mV; and C, stepping from -120 mV). The results indicate that the RNS-60 (filled circles) has a larger effect on whole-cell conductance than Solas (open circles). In the experiment similar results were seen in the three voltage protocols and at both the 15 minute and two-hour incubation time points.

Figures 118 A-C show graphs resulting from the subtraction of the Solas current data from the RNS-60 current data at three voltage protocols ("Delta currents") (A, stepping from zero mV; B, stepping from -60 mV; and C, stepping from -120 mV) and the two time-points (15 mins (open circles) and 2 hours (filled circles)). These data indicated that at the 15 minute time-point with RNS-60, there is a non-linear voltage-dependent component that is absent at the 2 hour time point.

As in previous experiments, data with "Normal" saline gave a very consistent and time-independent conductance used as a reference. The present results were obtained by matching groups with either Solas or RNS-60 saline, and indicate that exposure of Calu-3 cells to the RNS-60 saline under basal conditions (without cAMP, or any other stimulation), produces time-dependent effect(s), consistent with the activation of a voltage-regulated conductance at shorter incubation times (15 min). This phenomenon was not as apparent at the two-hour incubation point. As described elsewhere herein, the linear component is more evident when the conductance is increased by stimulation with the cAMP "cocktail". Nonetheless, the two-hour incubation time showed higher linear conductance for both the RNS-60 and the Solas saline, and in this case, the RNS-60 saline doubled the whole cell conductance as compared to Solas alone. This evidence indicates that at least two contributions to the whole cell conductance are affected by the RNS-60 saline, namely the activation of a non-linear voltage regulated conductance, and a linear conductance, which is more evident at longer incubation times.

Second set of experiments (effect on calcium permeable channels)

Methods for second set of experiments:

See EXAMPLE 23 for general patch clamp methods. In the following second set of experiments, yet additional patch clamp studies were performed to further confirm the utility of the inventive electrokinetically generated saline fluids (RNS-60 and Solas) to modulate whole-cell currents, using Calu-3 cells under basal conditions, with protocols stepping from either zero mV or -120 mV holding potentials.

The whole-cell conductance in each case was obtained from the current-to-voltage relationships obtained from cells incubated for 15 min with either saline. To determine whether there is a contribution of calcium permeable channels to the whole

cell conductance, and whether this part of the whole cell conductance is affected by incubation with RNS-60 saline, cells were patched in normal saline after the incubation period (entails a high NaCl external solution, while the internal solution contains high KCl). The external saline was then replaced with a solution where NaCl was replaced
5 by CsCl to determine whether there is a change in conductance by replacing the main external cation. Under these conditions, the same cell was then exposed to increasing concentrations of calcium, such that a calcium entry step is made more evident.

Results:

Figures 119 A-D show the results of a series of patch clamping experiments that
10 assessed the effects of the electrokinetically generated fluid (e.g., Solas (panels A and B) and RNS-60 (panels C and D)) on epithelial cell membrane polarity and ion channel activity using different external salt solutions and at different voltage protocols (panels A and C show stepping from zero mV, whereas panels B and D show stepping from -120 mV). In these experiments one time-point of 15 minutes was used. For Solas (panels
15 A and B) the results indicate that: 1) using CsCl (square symbols) instead of NaCl as the external solution, increased whole cell conductance with a linear behavior when compared to the control (diamond symbols); and 2) CaCl_2 at both 20 mM CaCl_2 (circle symbols) and 40 mM CaCl_2 (triangle symbols) increased whole cell conductance in a non-linear manner. For RNS-60 (panels C and D), the results indicate that: 1) using
20 CsCl (square symbols) instead of NaCl as the external solution had little effect on whole cell conductance when compared to the control (diamond symbols); and 2) CaCl_2 at 40 mM (triangle symbols) increased whole cell conductance in a non-linear manner.

Figures 120 A-D show the graphs resulting from the subtraction of the CsCl current data (shown in Figure 119) from the 20 mM CaCl_2 (diamond symbols) and 40
25 mM CaCl_2 (square symbols) current data at two voltage protocols (panels A and C, stepping from zero mV; and B and D, stepping from -120 mV) for Solas (panels A and B) and RNS-60 (panels C and D). The results indicate that both Solas and RNS-60 solutions activated a calcium-induced non-linear whole cell conductance. The effect was greater with RNS-60 (indicating a dosage responsiveness), and with RNS-60 was
30 only increased at higher calcium concentrations. Moreover, The non-linear calcium

dependent conductance at higher calcium concentration was also increased by the voltage protocol.

The data of this second set of experiments further indicates an effect of RNS-60 saline and Solas saline for whole cell conductance data obtained in Calu-3 cells. The data indicate that 15-min incubation with either saline produces a distinct effect on the whole cell conductance, which is most evident with RNS-60, and when external calcium is increased, and further indicates that the RNS-60 saline increases a calcium-dependent non-linear component of the whole cell conductance.

The accumulated evidence suggests activation by Revalesio saline of ion channels, which make different contributions to the basal cell conductance.

Taken together with Applicants' other data (e.g., the data of Applicants other working Examples) particular aspects of the present invention provide compositions and methods for modulating intracellular signal transduction, including modulation of at least one of membrane structure, membrane potential or membrane conductivity, membrane proteins or receptors, ion channels, lipid components, or intracellular components which are exchangeable by the cell (e.g., signaling pathways, such as calcium dependant cellular signaling systems, comprising use of the inventive electrokinetically generated solutions to impart electrochemical and/or conformational changes in membranous structures (e.g., membrane and/or membrane proteins, receptors or other membrane components) including but not limited to GPCRs and/or g-proteins. According to additional aspects, these effects modulate gene expression, and may persist, dependant, for example, on the half lives of the individual messaging components, etc.

EXAMPLE 25

(Atomic Force Microscopy (AFM) Measurements of the inventive electrokinetic fluid (RNS-60) indicated the presence and/or formation of hydrophobic surface nanobubbles that were substantially smaller than those present in control 'pressure pot' (PNS-60) fluid)

Overview. Applicants used Atomic Force Microscopy (AFM) measurements to characterize hydrophobic nanobubbles in the inventive electrokinetic fluid (RNS-60).

Materials and Methods:

AFM studies. AFM studies were preformed at an art-recognized Nanotech User Facility (NTUF). For AFM studies, a very small and sensitive needle is dipped into a droplet of water placed onto a hydrophobic surface. The needle then scans over the water/surface interface at rates such as 1 mm² in ~15 minutes. The needle records any imperfections in the surface geometry, and is sensitive enough to record the presence of small bubbles.

The Silicon substrate upon which the water droplets were placed was prepared using Trichloro(1H,1H,2H,2H-perfluorooctyl)silane), and the resulting hydrophobic surface causes water to bead up with contact angles of approximately 95 degrees. This coating is used in many AFM studies, in part, because it is particularly durable.

Solution Preparation. Two test solutions were studied: RNS-60 and PNS-60. RNS-60 is an inventive electrokinetic fluid comprising 60 ppm oxygen, whereas PNS-60 is a non-electrokinetic control fluid comprising 60 ppm oxygen prepared by conventional exposure to a pressurized oxygen head (i.e., pressure pot oxygenated fluid). Each test solution was initially buffered by addition of a small amount of neutral phosphate buffer (pH 7) solution, and approximately 60-70 uL of each buffered test solution (approximately 22°C) was placed onto a previously prepared silica plate.

Results:

Under AFM, the RNS-60 droplet displayed a distribution of about 20 hydrophobic nanobubbles in a 1 mm² area, having dimensions of ~20 nm wide and ~1.5 nm tall or smaller (Figure 121 A). By contrast, under AFM, the PNS-60 droplet displayed approx 5 hydrophobic nanobubbles in a 1 mm² area, having dimensions of ~60 nm wide and ~5 nm tall (Figure 121 B). The PNS-60 droplet, therefore, had much fewer and much larger hydrophobic nanobubbles compared to the RNS60 droplet.

According to particular aspects, therefore, there is a substantial difference in the size and distribution of hydrophobic surface nanobubbles between the RNS-60 and PNS-60 test solutions, where the nanobubbles are either initially present in, and/or formed within the test fluids during AFM measurement.

As discussed elsewhere herein, according to particular aspects of the present invention, the inventive electrokinetically altered fluids comprise an ionic aqueous solution of charge-stabilized oxygen-containing nanostructures substantially having an

average diameter of less than about 100 nanometers and stably configured in the ionic aqueous fluid in an amount sufficient to provide, upon contact of a living cell by the fluid, modulation of at least one of cellular membrane potential and cellular membrane conductivity.

5 Applicants point out, however, that the *hydrophobic* bubbles (forming on a hydrophobic surface), such as those observed in AFM experiments are likely fundamentally different from inventive biologically-active charge-stabilized nanostructure disclosed herein. According to particular aspects therefore, while the AFM experiments in this working Example support, based on the size and distribution
10 hydrophobic bubble formation, that the inventive electrokinetic fluids (e.g., RNS-60) are fundamentally distinct from non-electrokinetic control fluids, the hydrophobic bubbles are likely distinct from and/or derived from the inventive charge-stabilized oxygen-containing nanostructures described in detail elsewhere herein. In any event, relative to the inventive electrokinetic fluids, control pressure pot oxygenated fluids do not
15 comprise charge-stabilized oxygen-containing nanostructures capable of modulation of at least one of cellular membrane potential and cellular membrane conductivity.

EXAMPLE 26

20 *(RNS-60 was shown by Fluorescence-Activated Cell Sorting (FACS) analysis to have a pronounced effect on Expression of Cell Surface Receptors: CD193 (CCR3); CD154 (CD40L); CD11B; and CD3)*

Overview. Applicants used Fluorescence-Activated Cell Sorting (FACS) analysis to compare the levels of expression of cell surface receptors, CD193 (CCR3); CD154
25 (CD40L); CD11B; and CD3, on white blood cells incubated with either the inventive electrokinetic fluid (RNS-60) or normal saline control fluid.

Methods:

Ficoll-hypaque separated PBMC (apheresis – All Cells) preincubated approximately 1 hour in 30% solutions of RNS60 or Normal Saline (NS);

30 PBMC activated with 2 µg/ml of PHA-L for 24 or 40 hours;

Cells collected and washed into blocking/staining buffer, stained and fixed; and

Cells were analyzed by flow cytometry.

Results:

With respect to CD193 (CCR3), as shown in Figure 122 B, the receptor is substantially down-regulated in the presence of RNS-60 when compared to the level of the receptor expression in the normal saline control. This down regulation affects the phosphorylation of MAPK p38 (data not shown) which in turn down-regulates eotaxin (e.g., see Example 13 and Figure 57) which in turn down regulates IL 5 (data not shown) and as well alters eosinophil counts (e.g., see Example 13), which is one of the factors that, that example, alters the bronchoconstrictive response.

As discussed above in Example 13 in the context of the ovalbumin challenge model and shown in Figure 57, RNS-60 decreased the serum eotaxin levels in the OVA challenged groups when compared to the effect of normal saline. Therefore, according to particular aspects, RNS-60 has the potential to decrease both the ligand eotaxin and its receptor CCR3.

With respect to CD154 (CD40L), as shown in Figure 123 A, the receptor is down-regulated in the presence of RNS-60 when compared to the level of the receptor expression in normal saline.

With respect to CD11B, as shown in Figure 123 B, the receptor is down-regulated in the presence of RNS-60 when compared to the level of the receptor expression in normal saline.

With respect to CD3, as shown in Figure 123 C, the receptor is down-regulated in the presence of RNS-60 when compared to the level of the receptor expression in normal saline.

EXAMPLE 27

(RNS60, but not normal saline (NS), attenuated the activation of NFκB in MBP-primed T cells)

This Example shows that RNS60, but not normal saline (NS), attenuated the activation of NFκB in MBP-primed T cells. According to particular aspects, therefore, the present electrokinetically-generated fluids have substantial utility for treating inflammation and inflammation-mediated conditions and diseases, including but not limited to, diabetes and related metabolic disorders, insulin resistance,

neurodegenerative diseases (e.g., M.S., Parkinson's, Alzheimer's, etc), asthma, cystic fibrosis, vascular/coronary disease, retinal and/or macular degeneration, digestive disorders (e.g., inflammatory bowel disease, ulcerative colitis, Crohn's, etc.).

5 Overview:

It is increasingly clear that inhibition of insulin receptor signaling pathways is a central mechanism through which inflammatory and stress responses mediate insulin resistance (see, e.g., review by Wellen & Hotamisligil, *The Journal of Clinical Investigation*, 115:1111-1119, 2005).

10 *Overlap of metabolic and immune pathways.* Several serine/threonine kinases are activated by inflammatory or stressful stimuli and contribute to inhibition of insulin signaling, including JNK, inhibitor of NF- κ B kinase (IKK), and PKC- θ (Zick, Y. 2003. Role of Ser/Thr kinases in the uncoupling of insulin signaling. *Int. J. Obes. Relat. Metab. Disord.* 27(Suppl. 3):S56–S60). Again, the activation of these kinases in obesity
15 highlights the overlap of metabolic and immune pathways; these are the same kinases, particularly IKK and JNK, that are activated in the innate immune response by Toll-like receptor (TLR) signaling in response to LPS, peptidoglycan, double-stranded RNA, and other microbial products (Medzhitov, R. 2001. Toll-like receptors and innate immunity. *Nat. Rev. Immunol.* 1:135–145). Hence it is likely that components of TLR signaling
20 pathways will also exhibit strong metabolic activities.

PKC and IKK are activated by cellular lipid metabolites. Two other inflammatory kinases that play a large role in counteracting insulin action, particularly in response to lipid metabolites, are IKK and PKC- θ . Lipid infusion has been demonstrated to lead to a rise in levels of intracellular fatty acid metabolites, such as diacylglycerol (DAG) and
25 fatty acyl CoAs. This rise is correlated with activation of PKC- θ and increased Ser307 phosphorylation of IRS-1 (Yu, C., et al. 2002. Mechanism by which fatty acids inhibit insulin activation of insulin receptor substrate-1 (IRS-1)-associated phosphatidylinositol 3-kinase activity in muscle. *J. Biol. Chem.* 277:50230–50236). PKC- θ may impair insulin action by activation of another serine/threonine kinase, IKK β , or JNK (Perseghin,
30 G., Petersen, K., and Shulman, G.I. 2003. Cellular mechanism of insulin resistance: potential links with inflammation. *Int. J. Obes. Relat. Metab. Disord.* 27(Suppl. 3):S6–S11). Other PKC isoforms have also been reported to be activated by lipids and may

also participate in inhibition of insulin signaling (Schmitz-Peiffer, C. 2002. Protein kinase C and lipid-induced insulin resistance in skeletal muscle. *Ann. N. Y. Acad. Sci.* 967:146–157).

IKK β can impact insulin signaling by activating NF- κ B. IKK β can impact on insulin signaling through at least 2 pathways. First, it can directly phosphorylate IRS-1 on serine residues (Yin, M.J., Yamamoto, Y., and Gaynor, R.B. 1998. The anti-inflammatory agents aspirin and salicylate inhibit the activity of I κ B kinase- β . *Nature*. 396:77–80, Gao, Z., et al. 2002. Serine phosphorylation of insulin receptor substrate 1 by inhibitor kappa B kinase complex. *J. Biol. Chem.* 277:48115–48121).

Second, it can phosphorylate inhibitor of NF- κ B (I κ B), thus activating NF- κ B, a transcription factor that, among other targets, stimulates production of multiple inflammatory mediators, including TNF- α and IL-6 (Shoelson, S.E., Lee, J., and Yuan, M. 2003. Inflammation and the IKK β /I κ B/NF- κ B axis in obesity- and diet-induced insulin resistance. *Int. J. Obes. Relat. Metab. Disord.* 27(Suppl. 3):S49–S52). Mice heterozygous for IKK β are partially protected against insulin resistance due to lipid infusion, high-fat diet, or genetic obesity (Yuan, M., et al. 2001. Reversal of obesity- and diet induced insulin resistance with salicylates or targeted disruption of IKK β *Science*. 293:1673–1677; Kim, J.K., et al. 2001. Prevention of fat-induced insulin resistance by salicylate. *J. Clin. Invest.* 108:437–446; doi:10.1172/JCI200111559).

Moreover, inhibition of IKK β in human diabetics by high-dose aspirin treatment also improves insulin signaling, although at this dose, it is not clear whether other kinases are also affected (Hundal, R.S., et al. 2002. Mechanism by which high-dose aspirin improves glucose metabolism in type 2 diabetes. *J. Clin. Invest.* 109:1321–1326. doi:10.1172/JCI200214955). Recent studies have also begun to tease out the importance of IKK in individual tissues or cell types to the development of insulin resistance. Activation of IKK in liver and myeloid cells appears to contribute to obesity-induced insulin resistance, though this pathway may not be as important in muscle (Cai, D., et al. 2005. Local and systemic insulin resistance resulting from hepatic activation of IKK β and NF- κ B. *Nat. Med.* 11:183–190; Arkan, M.C., et al. 2005. IKK β links inflammation to obesity-induced insulin resistance. *Nat. Med.* 11:191–198; and Rohl, M., et al. 2004. Conditional disruption of I κ B kinase 2 fails to prevent obesity-induced insulin resistance. *J. Clin. Invest.* 113:474–481; doi:10.1172/JCI200418712).

Methods. For the experiments shown in Figures 124 A and 124 B, T cells isolated from MBP-immunized mice were re-primed with MBP and after 24 h, cells received different concentrations of RNS60 and NS. After 2 h of treatment, DNA-binding activity of NF- κ B was monitored in nuclear extracts by electrophoretic mobility shift assay (EMSA).

For experiments shown in Figure 124 C, T cells isolated from MBP-immunized mice were transfected with PBIIIX-Luc, an NF- κ B dependent reporter construct, followed by repriming with MBP. After 24 h of MBP priming, cells were treated with different concentrations of RNS60 and NS for 2 h followed by assay of luciferase activity in total cell extracts by a luciferase assay kit (Promega). In other cases, MBP-primed T cells were also stimulated with 30 nM PMA for 1 h. In these cases, PMA was added after 1 h of pretreatment with RNS60 and NS. Results are mean + SD of three different experiments.

Results. Figures 124 A-C show that RNS60, but not normal saline (NS), attenuated the activation of NF- κ B in MBP-primed T cells. Specifically, Figures 124 A and 124 B show that RNS60 (see middle three lanes of Figures 124 A and 124 B), but not NS (see right-most lane of Figures 124 A and 124 B), attenuated the activation of NF- κ B in MBP-primed T cells in a dose-responsive manner.

Likewise, the bar graph of Figure 124 C shows that that RNS60 (see second, third and fourth bars of Figures 124 A and 124 B), but not NS (see fifth bar of Figures 124 A and 124 B), attenuated the activation of NF- κ B in MBP-primed T cells, and hence also attenuated luciferase activity from the transfected NF- κ B-dependent reporter construct (PBIIIX-Luc) in total cell extracts, in a dose-responsive manner.

According to particular aspects, therefore, the disclosed electrokinetically-generated fluids have substantial utility for treating inflammation and inflammation-mediated conditions and diseases, including but not limited to, diabetes and related metabolic disorders, insulin resistance, neurodegenerative diseases (e.g., M.S., Parkinson's, Alzheimer's, etc), asthma, cystic fibrosis, vascular/coronary disease, retinal and/or macular degeneration, digestive disorders (e.g., inflammatory bowel disease, ulcerative colitis, Crohn's, etc.).

Incorporation by Reference

All of the above U.S. patents, U.S. patent application publications, U.S. patent applications, foreign patents, foreign patent applications, and non-patent publications referred to in this specification and/or listed in the Application Data Sheet, are
5 incorporated herein by reference, in their entirety.

It should be understood that the drawings and detailed description, herein are to be regarded in an illustrative rather than a restrictive manner, and are not intended to limit the invention to the particular forms and examples disclosed. On the contrary, the invention includes any further modifications, changes, rearrangements, substitutions,
10 alternatives, design choices, and embodiments apparent to those of ordinary skill in the art, without departing from the spirit and scope of this invention, as defined by the following claims. Thus, it is intended that the following claims be interpreted to embrace all such further modifications, changes, rearrangements, substitutions, alternatives, design choices, and embodiments.

The foregoing described embodiments depict different components contained within, or connected with, different other components. It is to be understood that such depicted architectures are merely exemplary, and that in fact many other architectures can be implemented which achieve the same functionality. In a conceptual sense, any arrangement of components to achieve the same functionality is effectively “associated”
15 such that the desired functionality is achieved. Hence, any two components herein combined to achieve a particular functionality can be seen as “associated with” each other such that the desired functionality is achieved, irrespective of architectures or intermedial components. Likewise, any two components so associated can also be viewed as being “operably connected,” or “operably coupled,” to each other to achieve
20 the desired functionality.

While particular embodiments of the present invention have been shown and described, it will be obvious to those skilled in the art that, based upon the teachings herein, changes and modifications may be made without departing from this invention and its broader aspects and, therefore, the appended claims are to encompass within
30 their scope all such changes and modifications as are within the true spirit and scope of this invention. Furthermore, it is to be understood that the invention is solely defined by the appended claims. It will be understood by those within the art that, in general,

terms used herein, and especially in the appended claims (e.g., bodies of the appended claims) are generally intended as “open” terms (e.g., the term “including” should be interpreted as “including but not limited to,” the term “having” should be interpreted as “having at least,” the term “includes” should be interpreted as “includes but is not limited to,” etc.). It will be further understood by those within the art that if a specific number of an introduced claim recitation is intended, such an intent will be explicitly recited in the claim, and in the absence of such recitation no such intent is present. For example, as an aid to understanding, the following appended claims may contain usage of the introductory phrases “at least one” and “one or more” to introduce claim recitations.

5 However, the use of such phrases should not be construed to imply that the introduction of a claim recitation by the indefinite articles “a” or “an” limits any particular claim containing such introduced claim recitation to inventions containing only one such recitation, even when the same claim includes the introductory phrases “one or more” or “at least one” and indefinite articles such as “a” or “an” (e.g., “a” and/or “an” should

10 typically be interpreted to mean “at least one” or “one or more”); the same holds true for the use of definite articles used to introduce claim recitations. In addition, even if a specific number of an introduced claim recitation *is* explicitly recited, those skilled in the art will recognize that such recitation should typically be interpreted to mean *at least* the recited number (e.g., the bare recitation of “two recitations,” without other modifiers,

15 typically means *at least* two recitations, or *two or more* recitations). Accordingly, the invention is not limited except as by the appended claims.

20

CLAIMS

1. A method for treating diabetes or a diabetes-associated condition or disorder, or symptoms thereof, comprising administering to a subject in need thereof a therapeutically effective amount of an electrokinetically altered aqueous fluid comprising an ionic aqueous solution of charge-stabilized oxygen-containing nanostructures substantially having an average diameter of less than about 100 nanometers and stably configured in the ionic aqueous fluid in an amount sufficient for treating diabetes or a diabetes-associated condition or disorder, or at least one symptom thereof.

2. The electrokinetic fluid of claim 1, wherein the charge-stabilized oxygen-containing nanostructures are the major charge-stabilized gas-containing nanostructure species in the fluid.

3. The electrokinetic fluid of claim 1, wherein the percentage of dissolved oxygen molecules present in the fluid as the charge-stabilized oxygen-containing nanostructures is a percentage selected from the group consisting of greater than: 0.01%, 0.1%, 1%, 5%; 10%; 15%; 20%; 25%; 30%; 35%; 40%; 45%; 50%; 55%; 60%; 65%; 70%; 75%; 80%; 85%; 90%; and 95%.

4. The electrokinetic fluid of claim 1, wherein the total dissolved oxygen is substantially present in the charge-stabilized oxygen-containing nanostructures.

5. The electrokinetic fluid of claim 1, wherein the charge-stabilized oxygen-containing nanostructures substantially have an average diameter of less than a size selected from the group consisting of: 90 nm; 80 nm; 70 nm; 60 nm; 50 nm; 40 nm; 30 nm; 20 nm; 10 nm; and less than 5 nm.

6. The electrokinetic fluid of claim 1, wherein the ionic aqueous solution comprises a saline solution.

7. The electrokinetic fluid of claim 1, wherein the fluid is superoxygenated.

8. The electrokinetic fluid of claim 1, wherein the fluid comprises a form of solvated electrons.

9. The method of claim 1, wherein alteration of the electrokinetically altered aqueous fluid comprises exposure of the fluid to hydrodynamically-induced, localized electrokinetic effects.

10. The method of claim 9, wherein, exposure to the localized electrokinetic effects comprises exposure to at least one of voltage pulses and current pulses.

11. The method of claim 9, wherein the exposure of the fluid to hydrodynamically-induced, localized electrokinetic effects, comprises exposure of the fluid to electrokinetic effect-inducing structural features of a device used to generate the fluid.

12. The method of claim 1, wherein the diabetes-associated condition or disorder comprises at least one selected from the group consisting of: diabetes; insulin-dependent diabetes mellitus or IDDM (Type 1); non-insulin dependent diabetes mellitus or NIDDM (Type 2); insulin resistance; and diabetic retinopathy.

13. The method of claim 12, wherein the diabetes-associated condition or disorder comprises at least one of diabetes and insulin resistance.

14. The method of claim 13, wherein the diabetes-associated condition or disorder comprises diabetes.

15. The method of claim 1, wherein the at least one symptom of the diabetes-associated condition or disorder is related to at least one condition selected from the group consisting of: chronic inflammation, acute inflammation, insulin resistance.

16. The method of claim 1, wherein the electrokinetically altered aqueous fluid modulates localized or cellular levels of nitric oxide.

17. The method of claim 1 wherein the electrokinetically altered aqueous fluid promotes a localized decrease at the site of administration of at least one cytokine selected from the group consisting of: IL-1beta, IL-8, TNF-alpha, and TNF-beta.

18. The method of claim 1, further comprising a synergistic or non-synergistic inhibition or reduction in inflammation by simultaneously or adjunctively treating the subject with another anti-inflammatory agent.

19. The method of claim 18, wherein said other anti-inflammatory agent comprises a steroid or glucocorticoid steroid.

20. The method of claim 19, wherein the glucocorticoid steroid comprises Budesonide or an active derivative thereof.

21. The method of claim 1, further comprising combination therapy, wherein at least one additional therapeutic agent is administered to the patient.

22. The method of claim 21, wherein the at least one additional therapeutic agent is selected from the group consisting of: Biguanides including metformin, buformin, and phenformin, insulin, alpha-glucosidase inhibitors, biguanides, DPP-4 inhibitors, meglitinides, sulfonylureas, thiazolidinediones, alpha-glucosidase inhibitors including, acarbose and miglitol, DPP-4 inhibitors including vildagliptin, sitagliptin, saxagliptin, linagliptin, and alogliptin, sulfonylureas including acetohexamide, chlorpropamide, tolbutamide, tolazamide, glipizide, gliclazide, glibenclamide (glyburide), gliquidone, glycopyramide, and glimepiride, meglitinides including nateglinide, mitiglinide, and repaglinide. thiazolidinediones including troglitazone, pioglitazone, and rosiglitazone, inhibitors of MMPs including inhibitor of MMP-9 and MMP-2, short-acting β_2 -agonists, long-acting β_2 -agonists, anticholinergics, corticosteroids, systemic corticosteroids, mast cell stabilizers, leukotriene modifiers, methylxanthines, β_2 -agonists, albuterol, levalbuterol, pirbuterol, arformoterol, formoterol, salmeterol, anticholinergics including ipratropium and tiotropium; corticosteroids including beclomethasone, budesonide, flunisolide, fluticasone, mometasone, triamcinolone, methyprednisolone, prednisolone, prednisone; leukotriene modifiers including montelukast, zafirlukast, and zileuton; mast cell stabilizers including cromolyn and nedocromil; methylxanthines including theophylline; combination drugs including ipratropium and albuterol, fluticasone and salmeterol, budesonide and formoterol; antihistamines including hydroxyzine, diphenhydramine, loratadine, cetirizine, and hydrocortisone; immune system modulating drugs including tacrolimus and pimecrolimus; cyclosporine; azathioprine; mycophenolatemofetil; and combinations thereof.

23. The method of claim 21, wherein the at least one additional therapeutic agent is a TSLP and/or TSLPR antagonist.

24. The method of claim 23, wherein the TSLP and/or TSLPR antagonist is selected from the group consisting of neutralizing antibodies specific for TSLP and the TSLP receptor, soluble TSLP receptor molecules, and TSLP receptor fusion proteins, including TSLPR-immunoglobulin Fc molecules or polypeptides that encode components of more than one receptor chain.

25. The method of claim 1, wherein treating comprises altering at least one of cellular membrane structure or function comprising altering at least one of a conformation, ligand binding activity, or a catalytic activity of a membrane associated protein.

5 26. The method of claim 25, wherein the membrane associated protein comprises at least one selected from the group consisting of receptors, transmembrane receptors, ion channel proteins, intracellular attachment proteins, cellular adhesion proteins, and integrins.

10 27. The method of claim 26, wherein the transmembrane receptor comprises a G-Protein Coupled Receptor (GPCR).

28. The method of claim 27, wherein the G-Protein Coupled Receptor (GPCR) interacts with a G protein a subunit.

29. The method of claim 28, wherein the G protein a subunit comprises at least one selected from the group consisting of G_{α_s} , G_{α_i} , G_{α_q} , and $G_{\alpha_{12}}$.

15 30. The method of claim 29, wherein the at least one G protein a subunit is G_{α_q} .

20 31. The method of claim 1, wherein the charge-stabilized oxygen-containing nanostructures are stably configured in the ionic aqueous fluid in an amount sufficient to provide, upon contact of a living cell by the fluid, modulation of at least one of cellular membrane potential and cellular membrane conductivity.

32. The method of claim 31, wherein modulating cellular membrane conductivity, comprises modulating whole-cell conductance.

25 33. The method of claim 33, wherein modulating whole-cell conductance, comprises modulating at least one voltage-dependent contribution of the whole-cell conductance.

34. The method of claim 31, comprising modulation of a calcium dependant cellular messaging pathway or system.

35. The method of claim 31, comprising modulation of phospholipase C activity.

30 36. The method of claim 31, comprising modulation of adenylate cyclase (AC) activity.

37. The method of claim 31, comprising modulation of intracellular signal transduction associated with at least one condition or symptom selected from the group consisting of: chronic inflammation, acute inflammation, insulin resistance.

38. The method of claim 1, comprising administration to a cell network or
5 layer, and further comprising modulation of an intercellular junction therein.

39. The method of claim 38, wherein the intracellular junction comprises at least one selected from the group consisting of tight junctions, gap junctions, zona adherins and desmosomes.

40. The method of claim 38, wherein the cell network or layers comprises at
10 least one selected from the group consisting of endothelial cells, endothelial-astrocyte tight junctions in CNS vessels, blood-cerebrospinal fluid tight junctions or barrier, pulmonary epithelium-type junctions, bronchial epithelium-type junctions, and intestinal epithelium-type junctions.

41. The method of claim 1, wherein the electrokinetically altered aqueous fluid
15 is oxygenated, and wherein the oxygen in the fluid is present in an amount of at least 8 ppm, at least 15, ppm, at least 25 ppm, at least 30 ppm, at least 40 ppm, at least 50 ppm, or at least 60 ppm oxygen at atmospheric pressure.

42. The method of any one of claims 1 through 41, wherein the electrokinetically altered aqueous fluid comprises at least one of a form of solvated
20 electrons, and electrokinetically modified or charged oxygen species.

43. The method of claim 42, wherein the solvated electrons or electrokinetically modified or charged oxygen species are present in an amount of at least 0.01 ppm, at least 0.1 ppm, at least 0.5 ppm, at least 1 ppm, at least 3 ppm, at least 5 ppm, at least 7 ppm, at least 10 ppm, at least 15 ppm, or at least 20 ppm.

25 44. The method of claim 43, wherein the electrokinetically altered oxygenated aqueous fluid comprises solvated electrons stabilized, at least in part, by molecular oxygen.

45. The method of claim 1, wherein the ability to alter cellular membrane structure or function sufficient to provide for modulation of intracellular signal
30 transduction persists for at least two, at least three, at least four, at least five, at least 6, at least 12 months, or longer periods, in a closed gas-tight container.

46. The method of claim 1, wherein the amount of oxygen present in charge-stabilized oxygen-containing nanostructures of the electrokinetically-altered fluid is at least 8 ppm, at least 15 ppm, at least 20 ppm, at least 25 ppm, at least 30 ppm, at least 40 ppm, at least 50 ppm, or at least 60 ppm oxygen at atmospheric pressure.

5 47. A method of formulating a therapeutic agent suitable for use in treating diabetes, or a diabetes-associated condition or disorder, or symptoms thereof, comprising:

obtaining a therapeutic agent suitable for use in treating diabetes, or a diabetes-associated condition or disorder, or symptoms thereof, of a subject; and

10 combining the therapeutic agent with an amount of an electrokinetically altered aqueous fluid comprising an ionic aqueous solution of charge-stabilized oxygen-containing nanostructures substantially having an average diameter of less than about 100 nanometers and stably configured in the ionic aqueous fluid in an amount sufficient for treating inflammation or at least one symptom thereof, wherein formulating a
15 therapeutic agent suitable for use in treating diabetes, or a diabetes-associated condition or disorder, or symptoms thereof is afforded.

48. The method of claim 48, wherein the charge-stabilized oxygen-containing nanostructures are stably configured in the ionic aqueous fluid in an amount sufficient to provide, upon contact of a living cell by the fluid, modulation of at least one of cellular
20 membrane potential and cellular membrane conductivity.

49. A pharmaceutical composition, comprising: a therapeutic agent suitable for use treating diabetes, or a diabetes-associated condition or disorder, or symptoms thereof, of a subject; and an amount of an electrokinetically altered aqueous fluid comprising an ionic aqueous solution of charge-stabilized oxygen-containing
25 nanostructures substantially having an average diameter of less than about 100 nanometers and stably configured in the ionic aqueous fluid in an amount sufficient for treating inflammation or at least one symptom thereof.

50. A pharmaceutical composition, prepared by the method of claim 48.

51. The method of claim 1, wherein treating comprises administration by at
30 least one of topical, inhalation, intranasal, and intravenous.

52. The method of claim 25, wherein the membrane associated protein comprises CCR3.

53. The method of claim 1, wherein treating comprises modulation of intracellular NF- κ B expression and/or activity.

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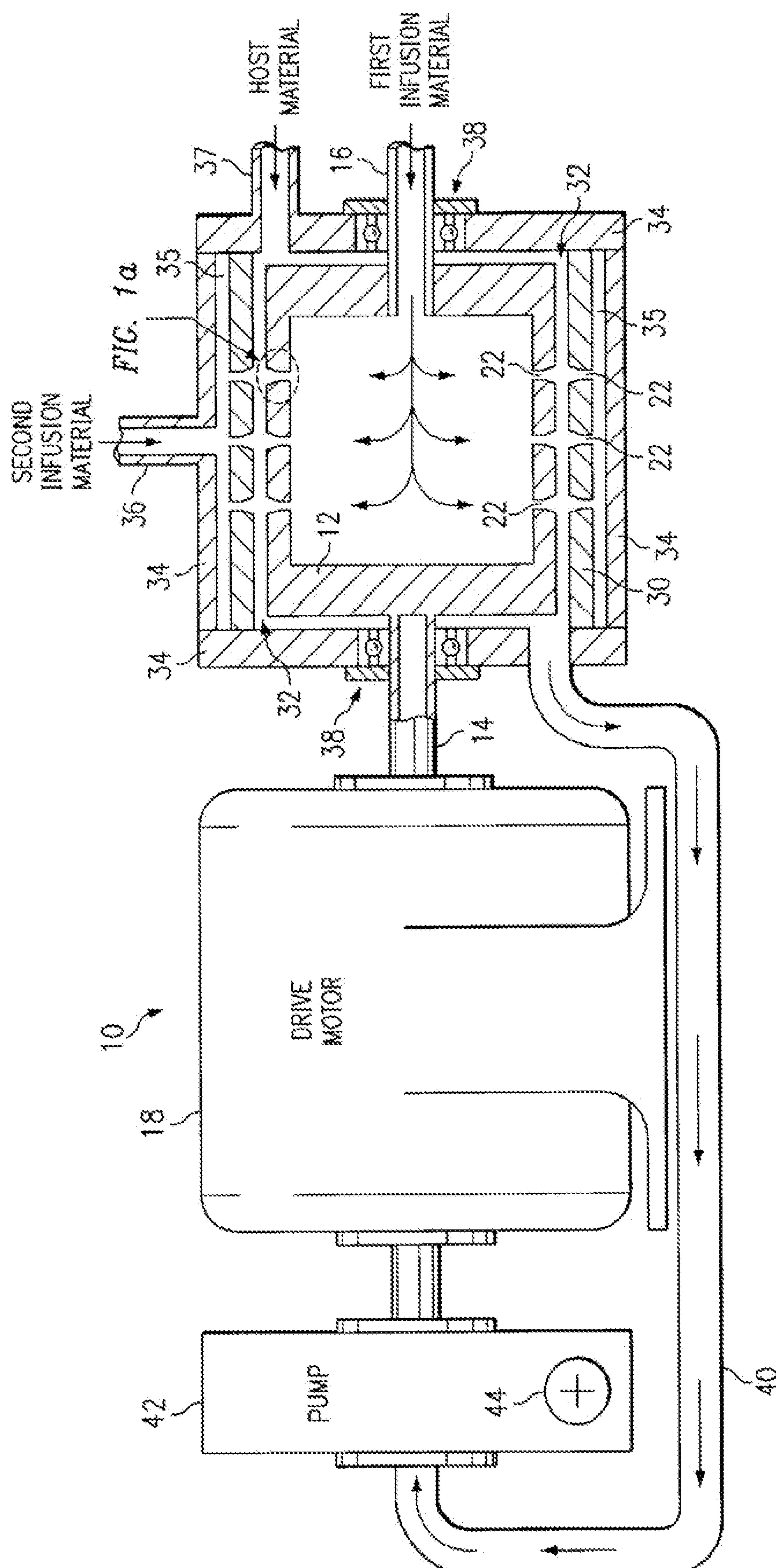


Fig. 1
(Prior Art)

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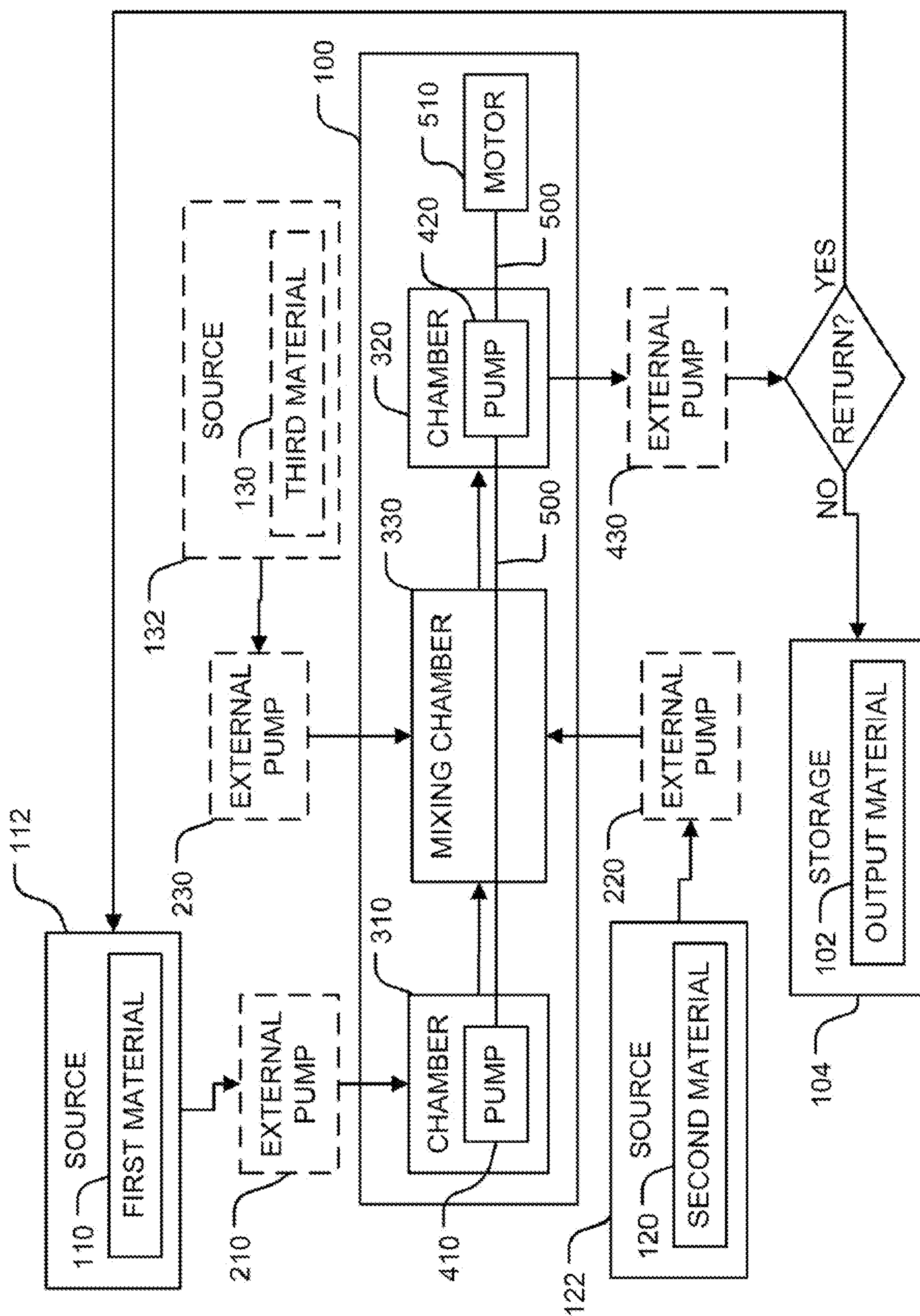


Fig. 2

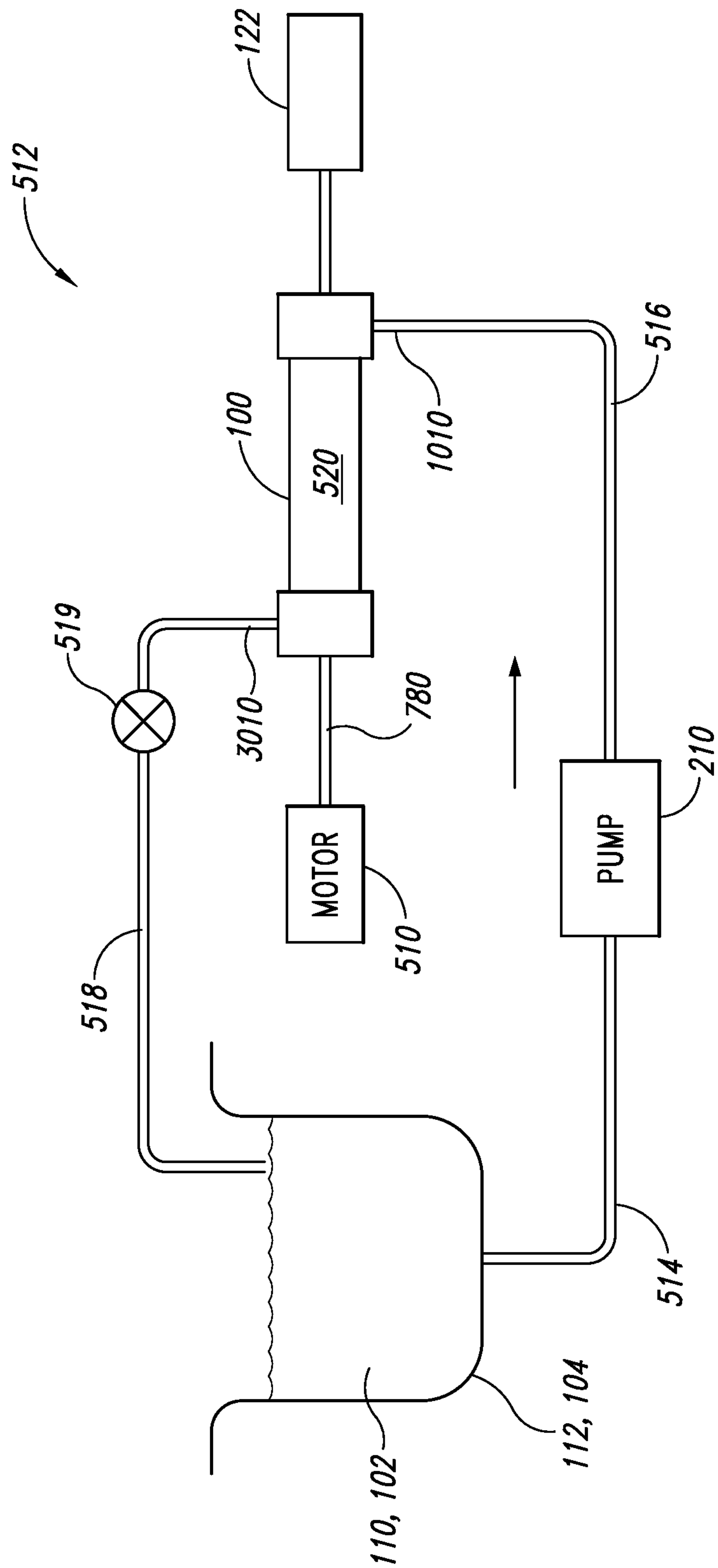


Fig. 3

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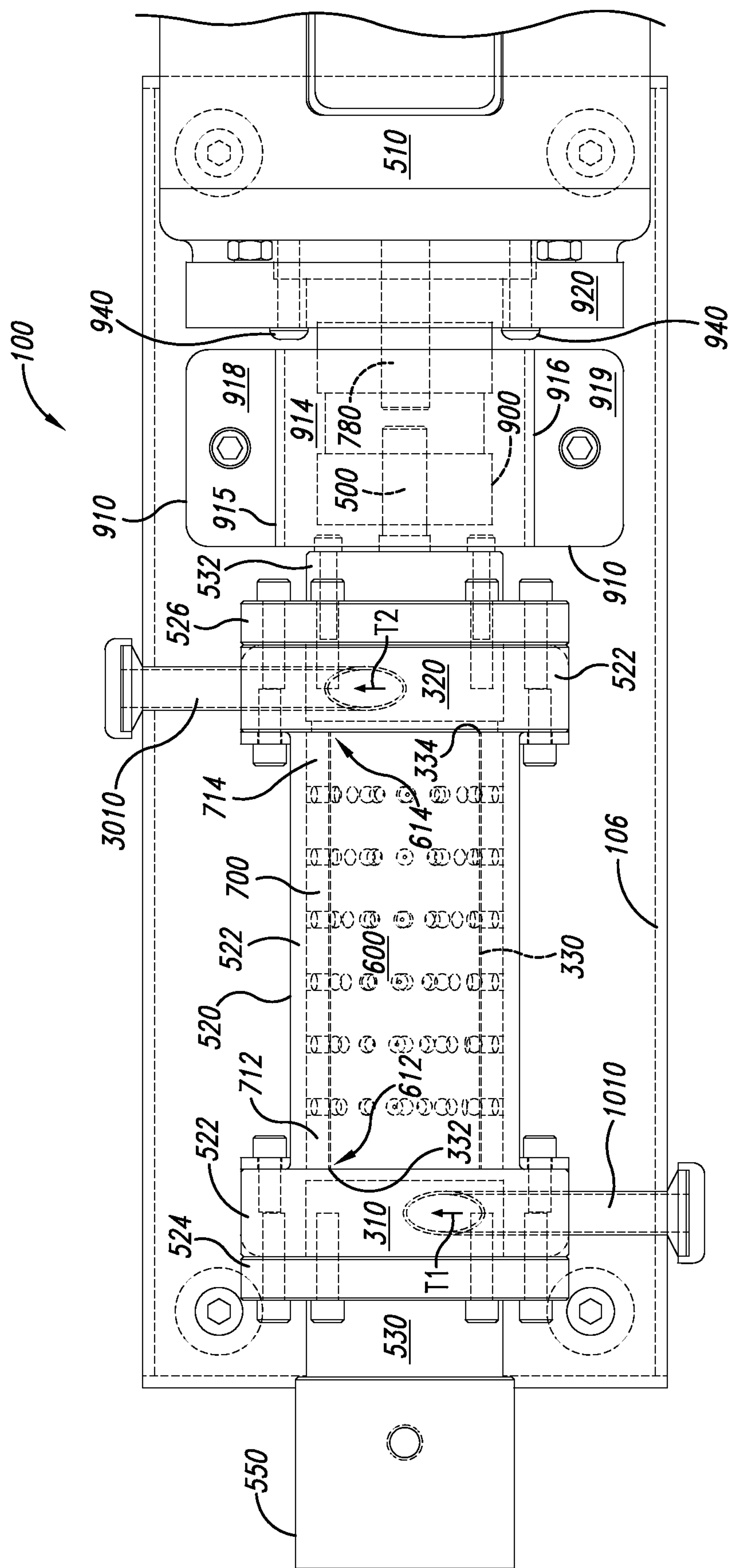


Fig. 4

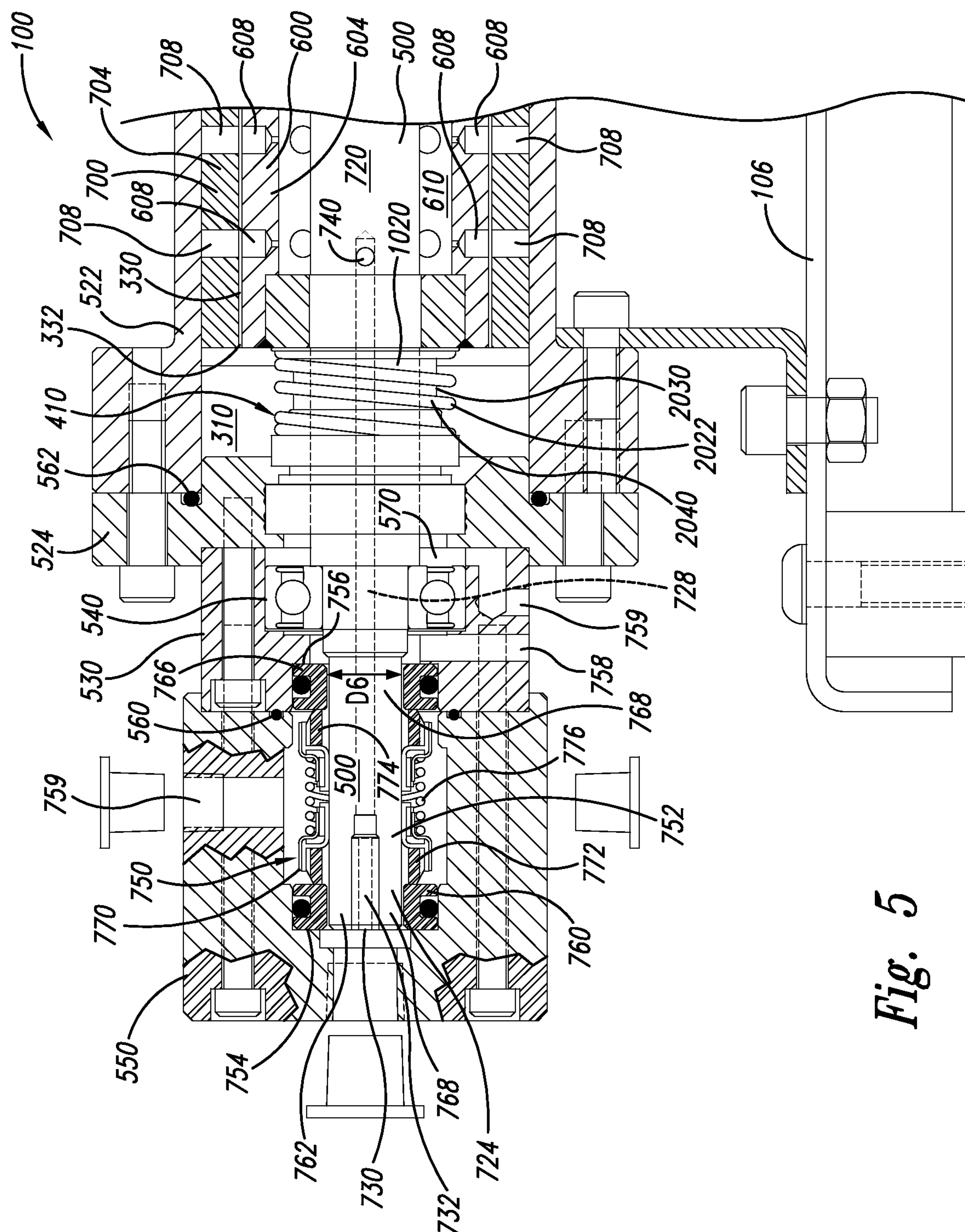


Fig. 5

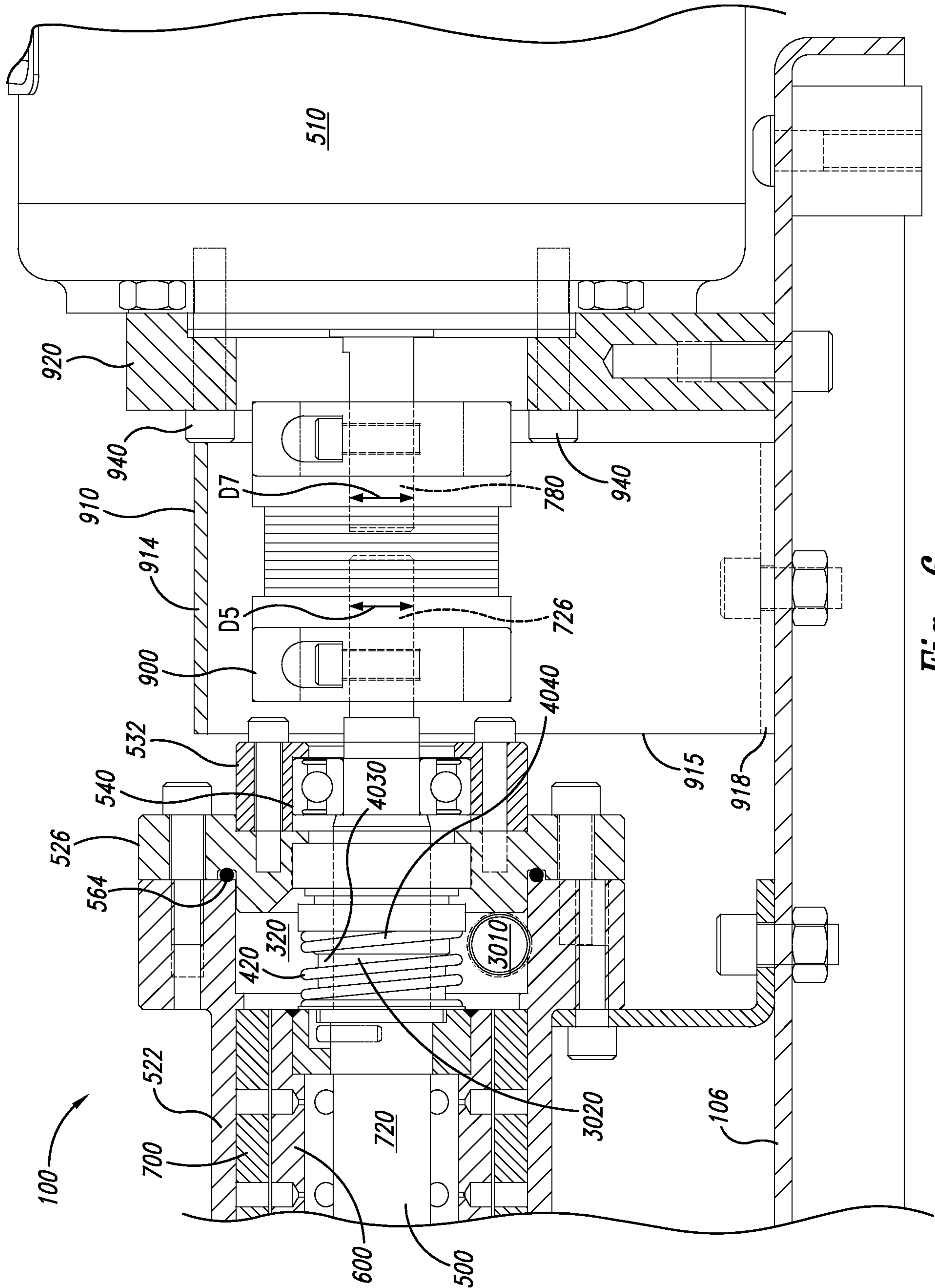


Fig. 6

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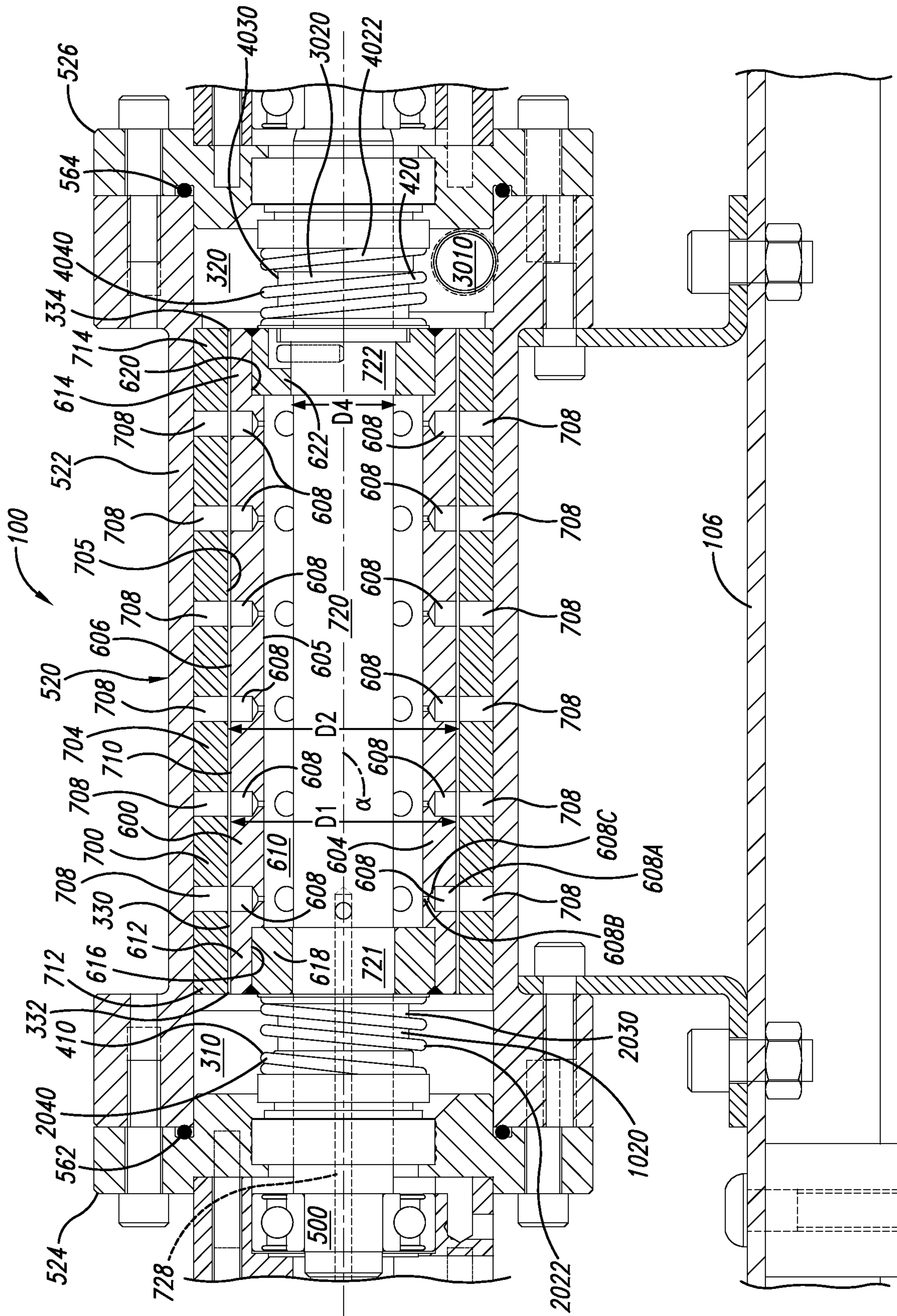


Fig. 7

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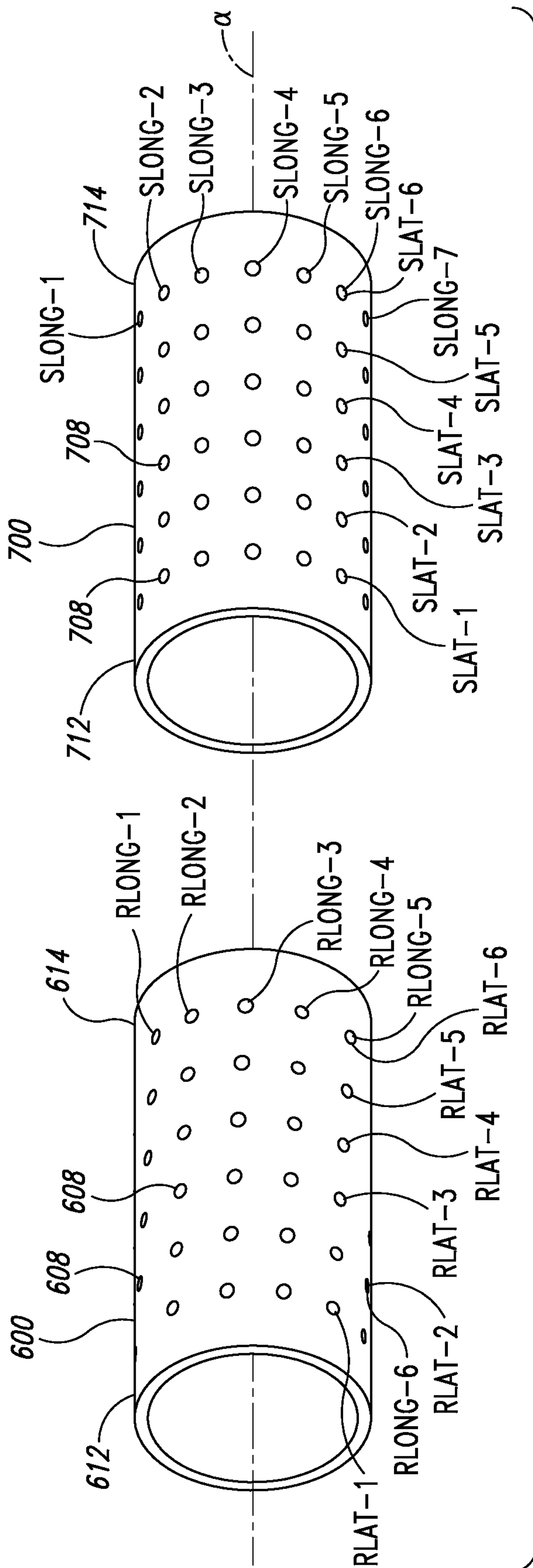


Fig. 8

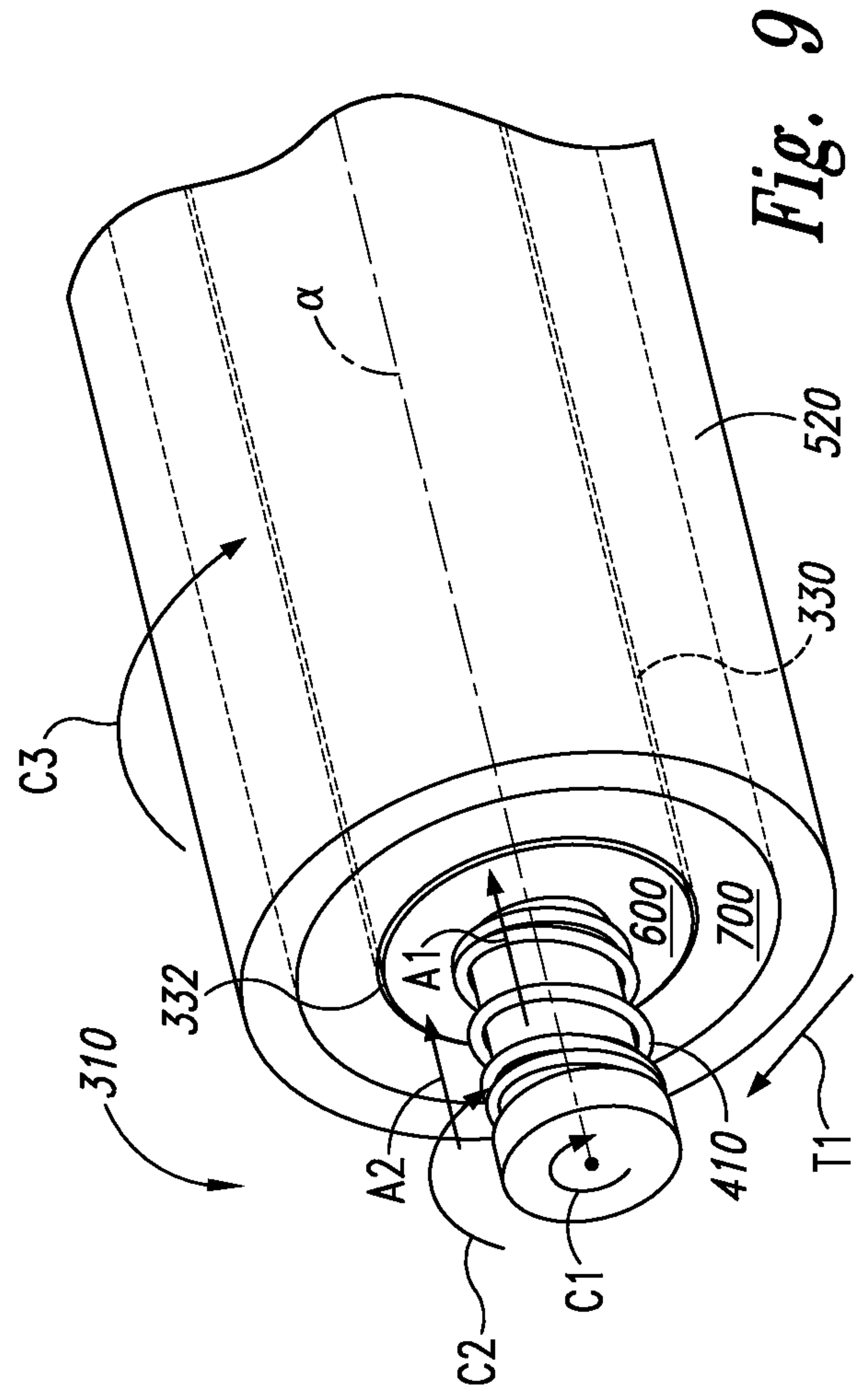
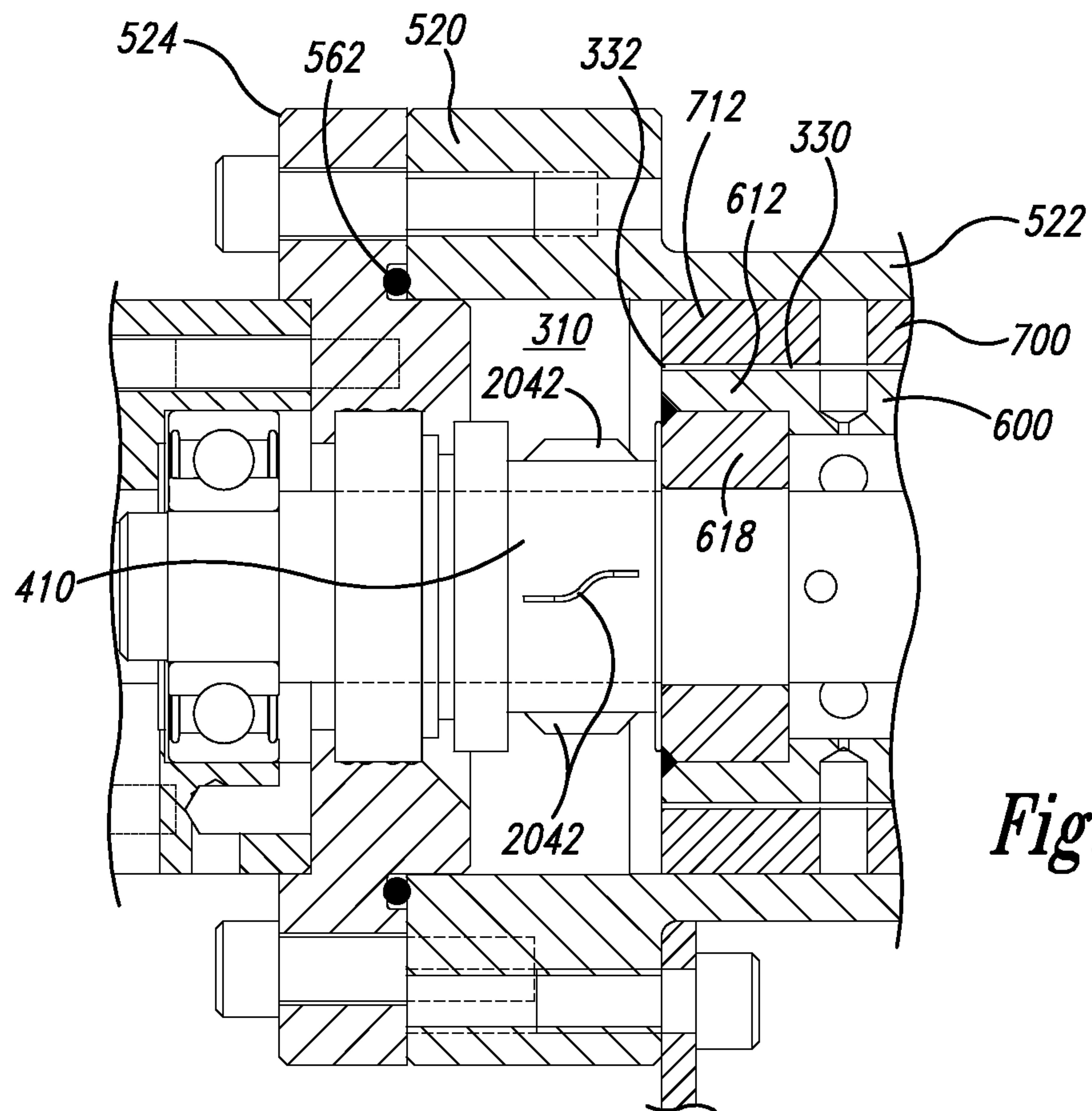
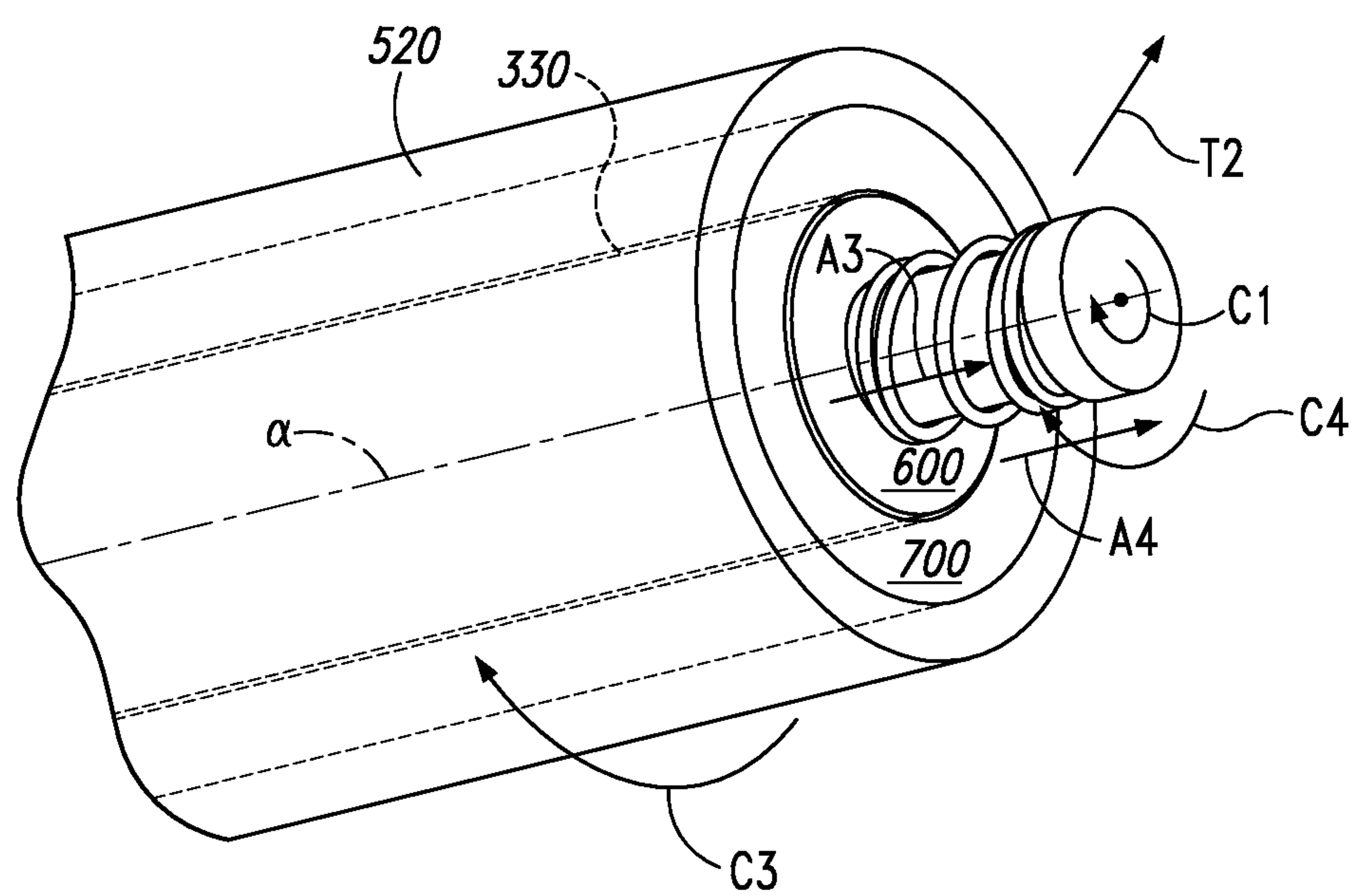
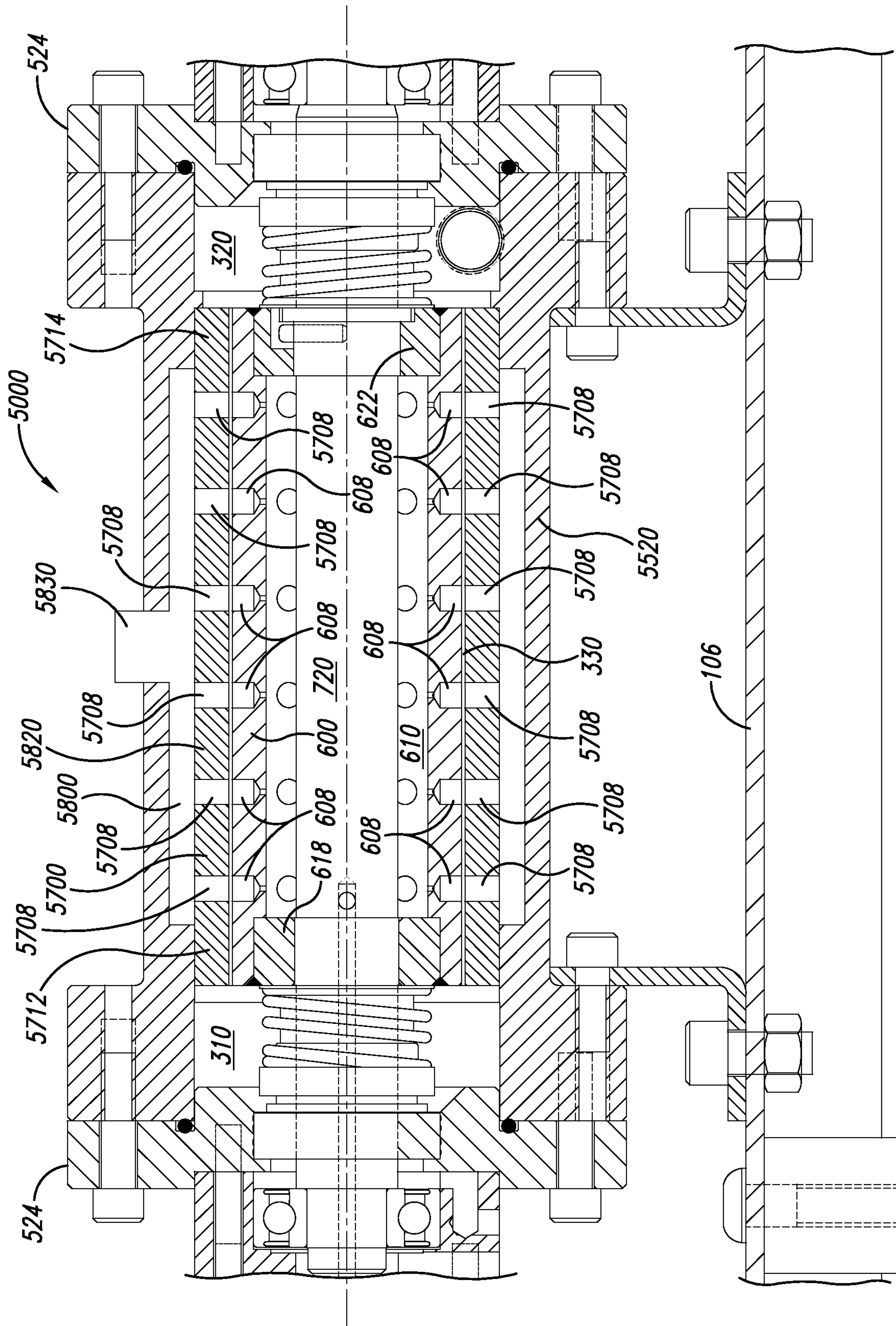


Fig. 9

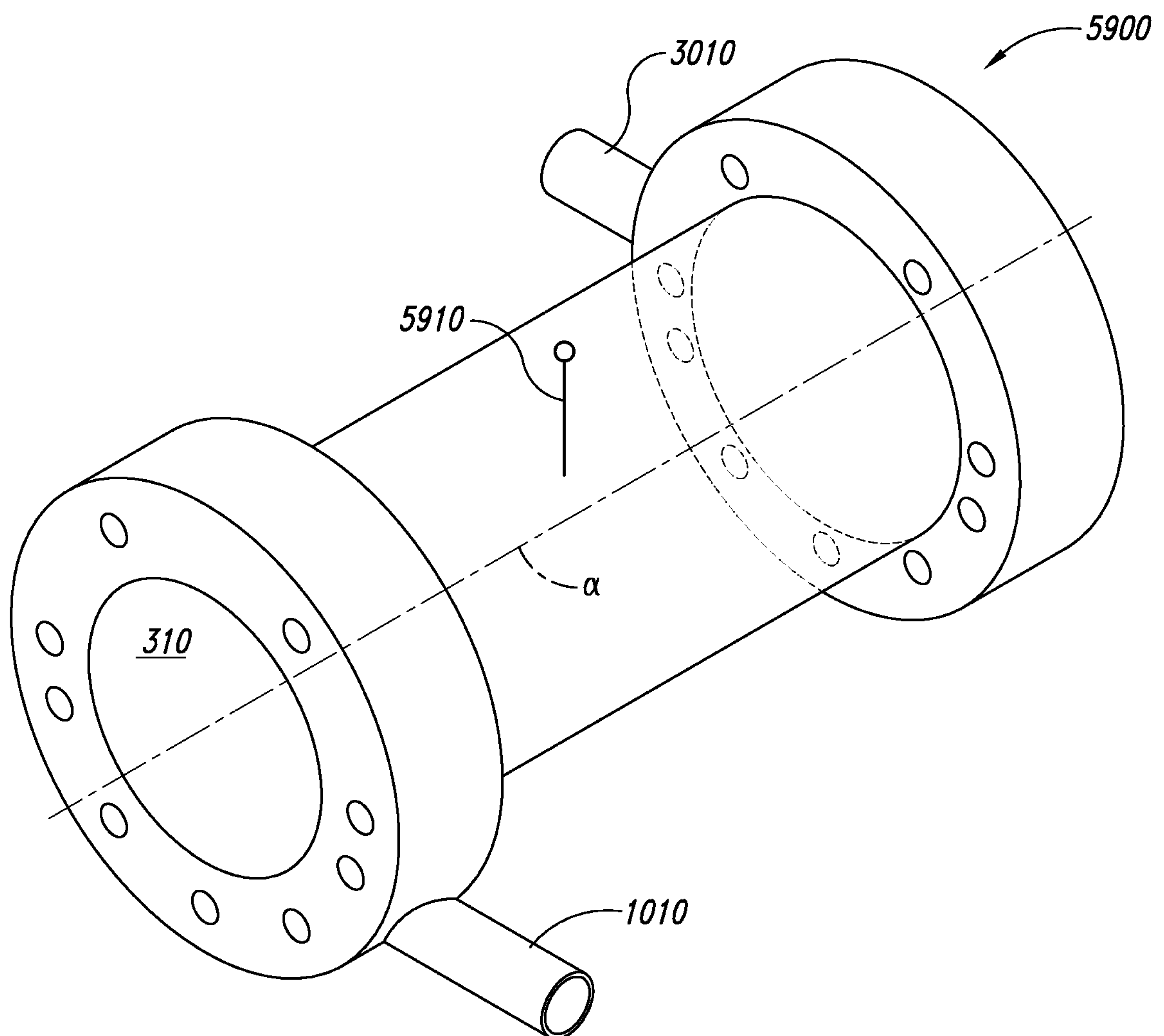
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*Fig. 10**Fig. 11*

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*Fig. 12*

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*Fig. 13*

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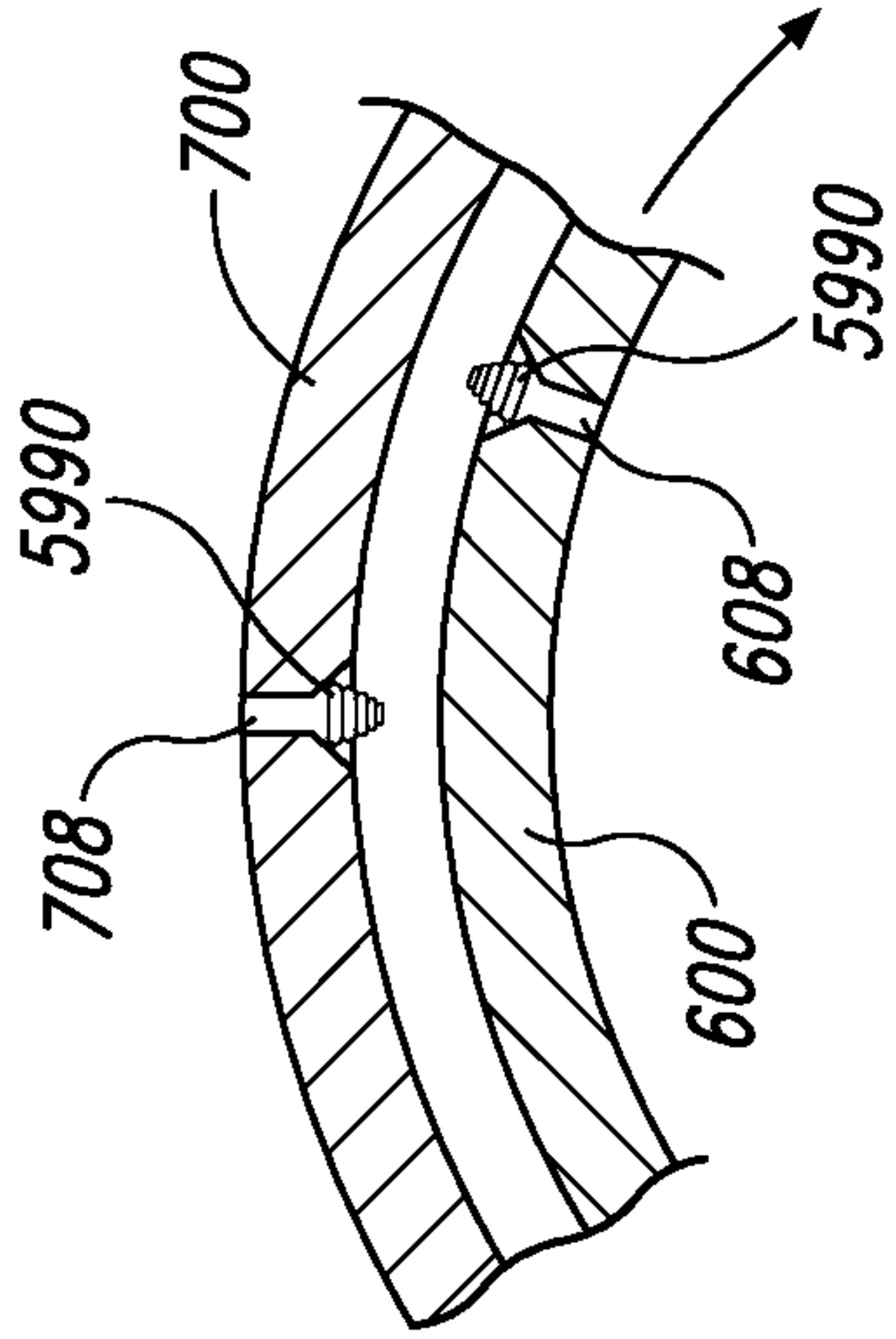


Fig. 15

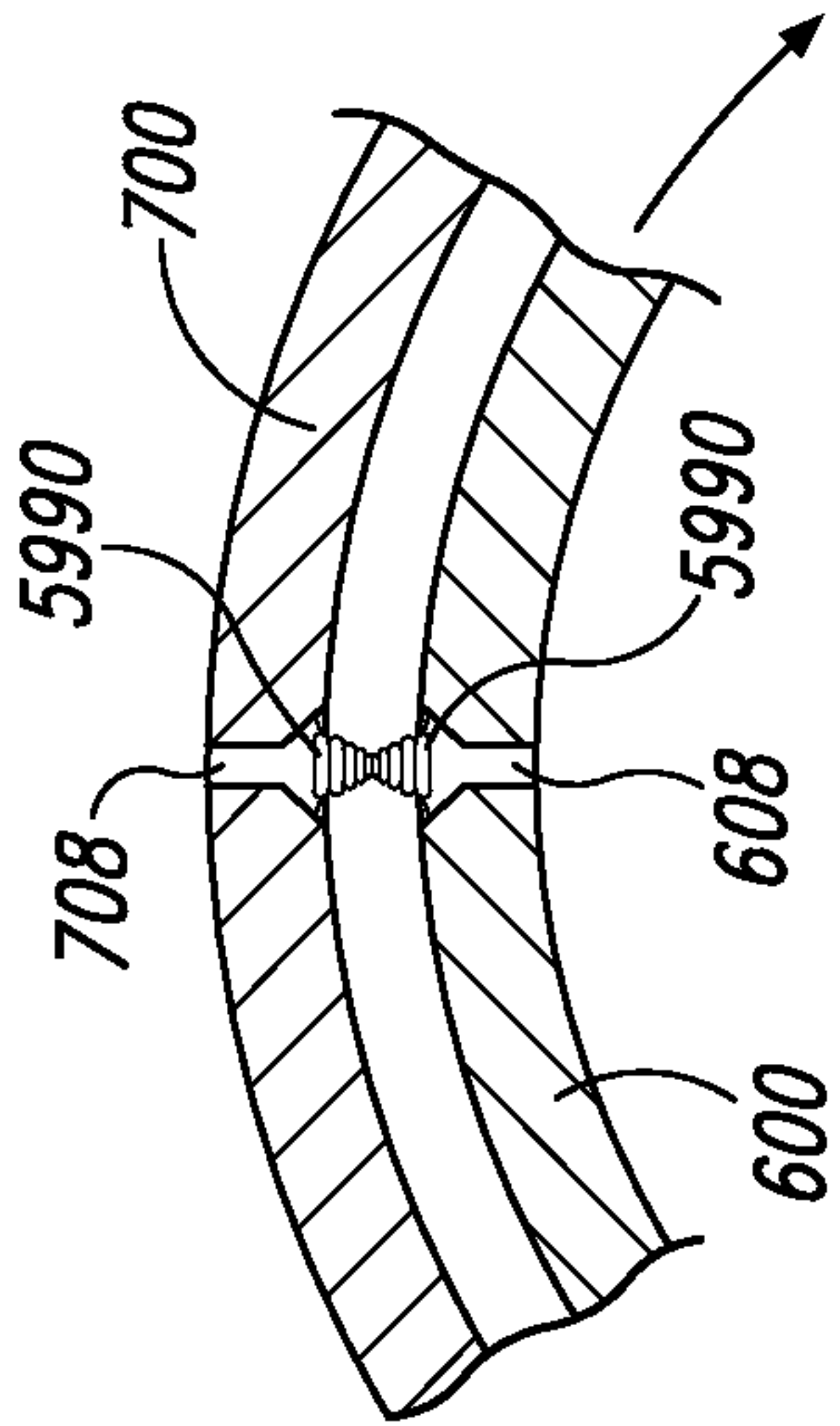


Fig. 16

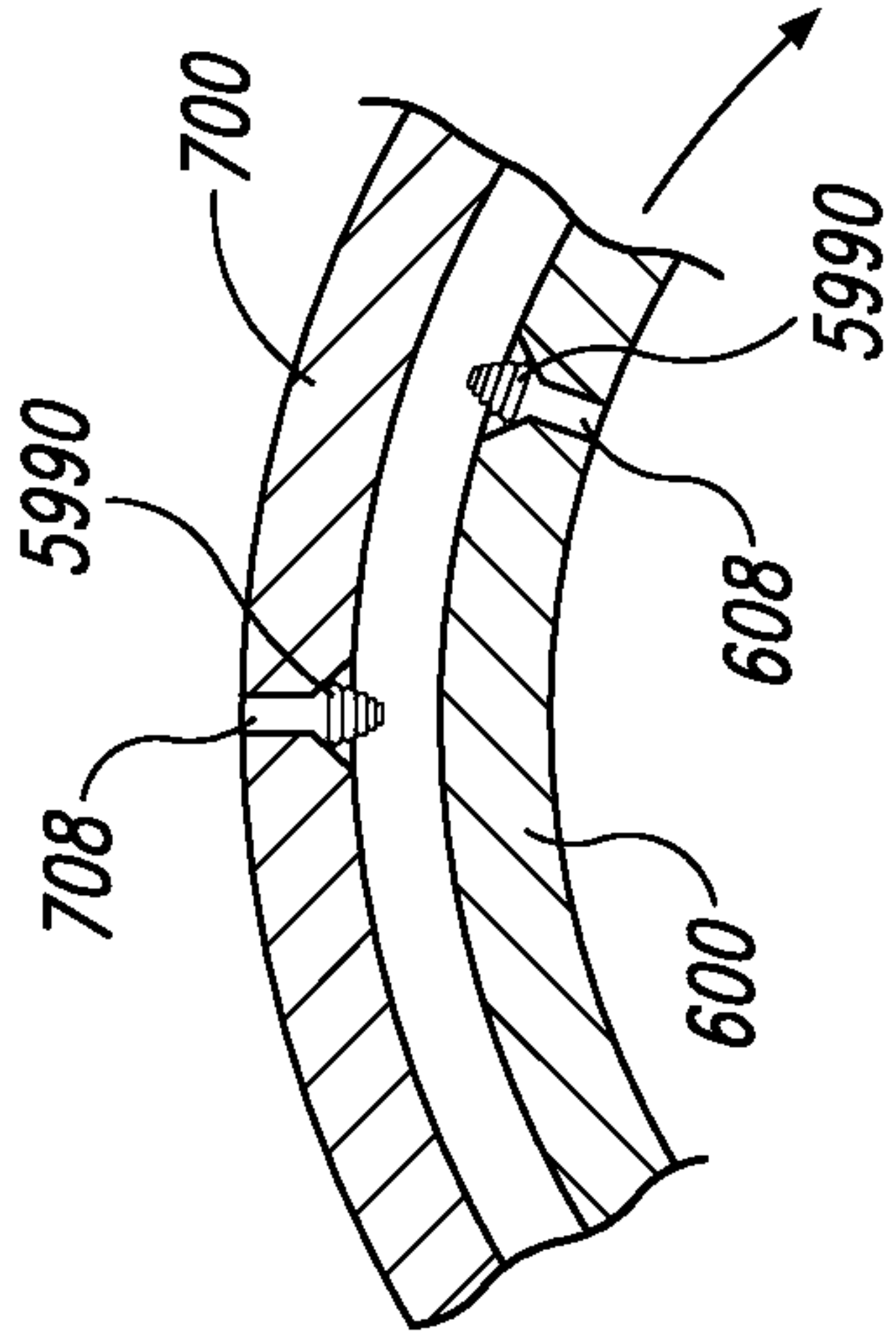


Fig. 17

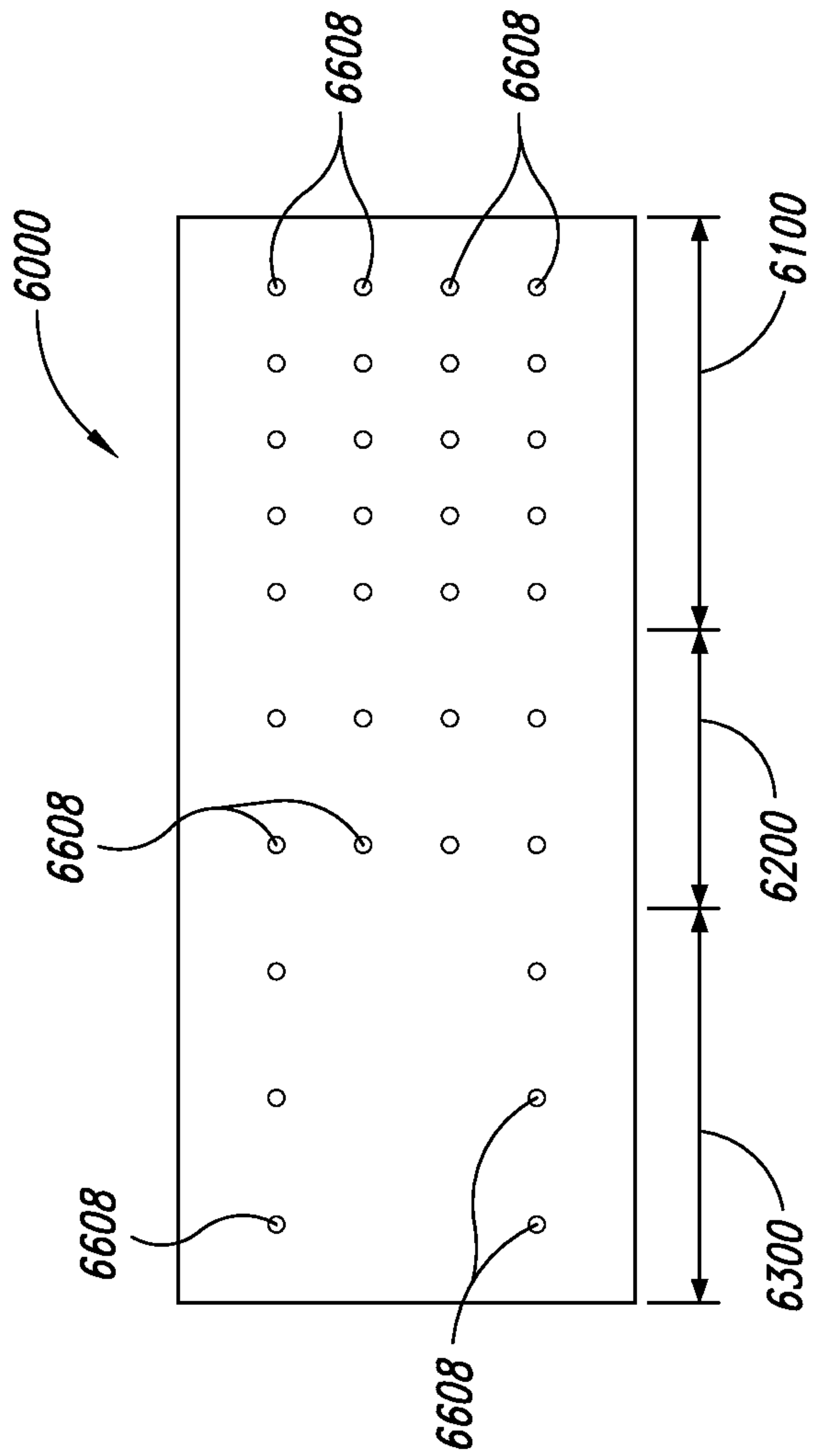
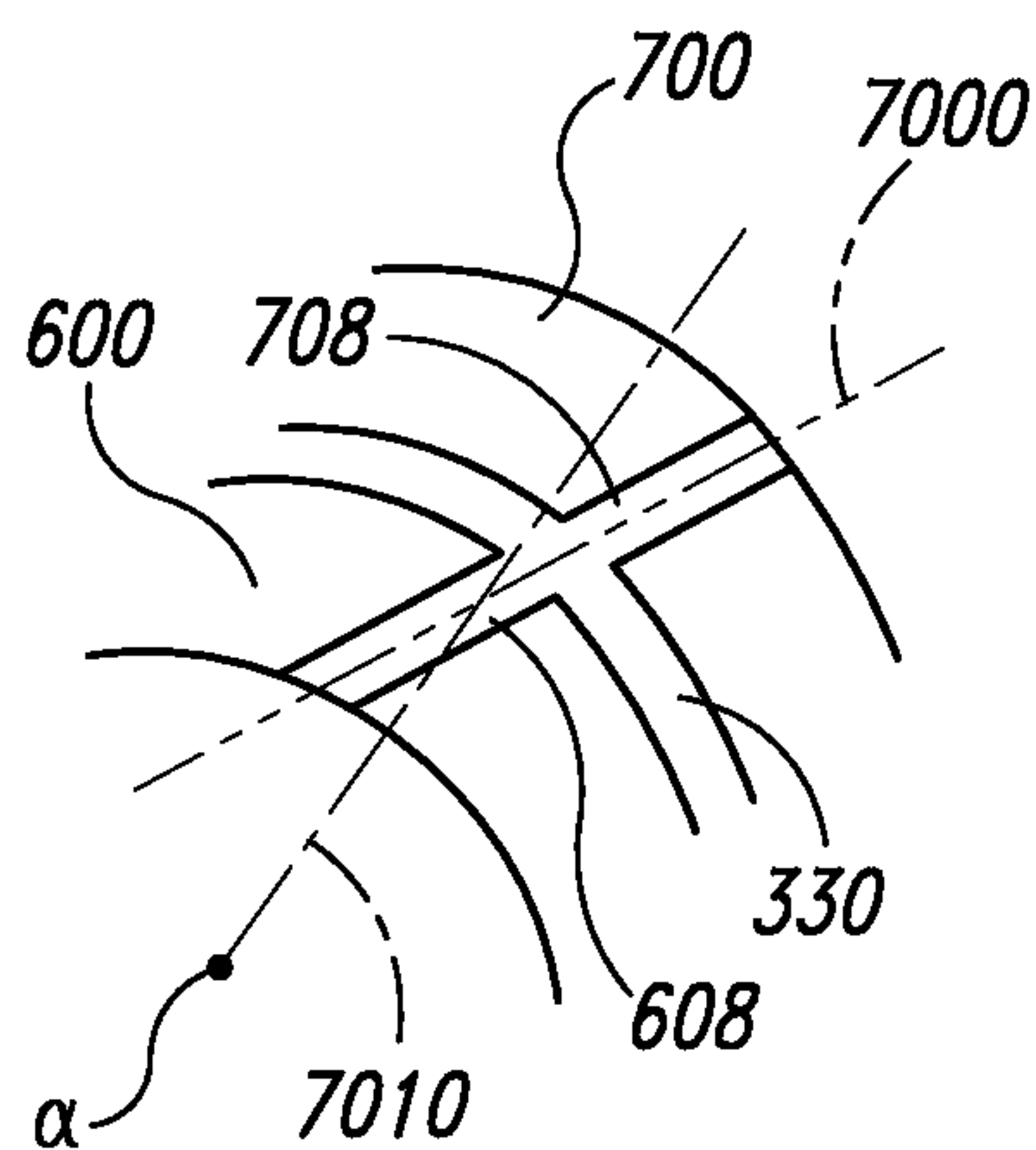
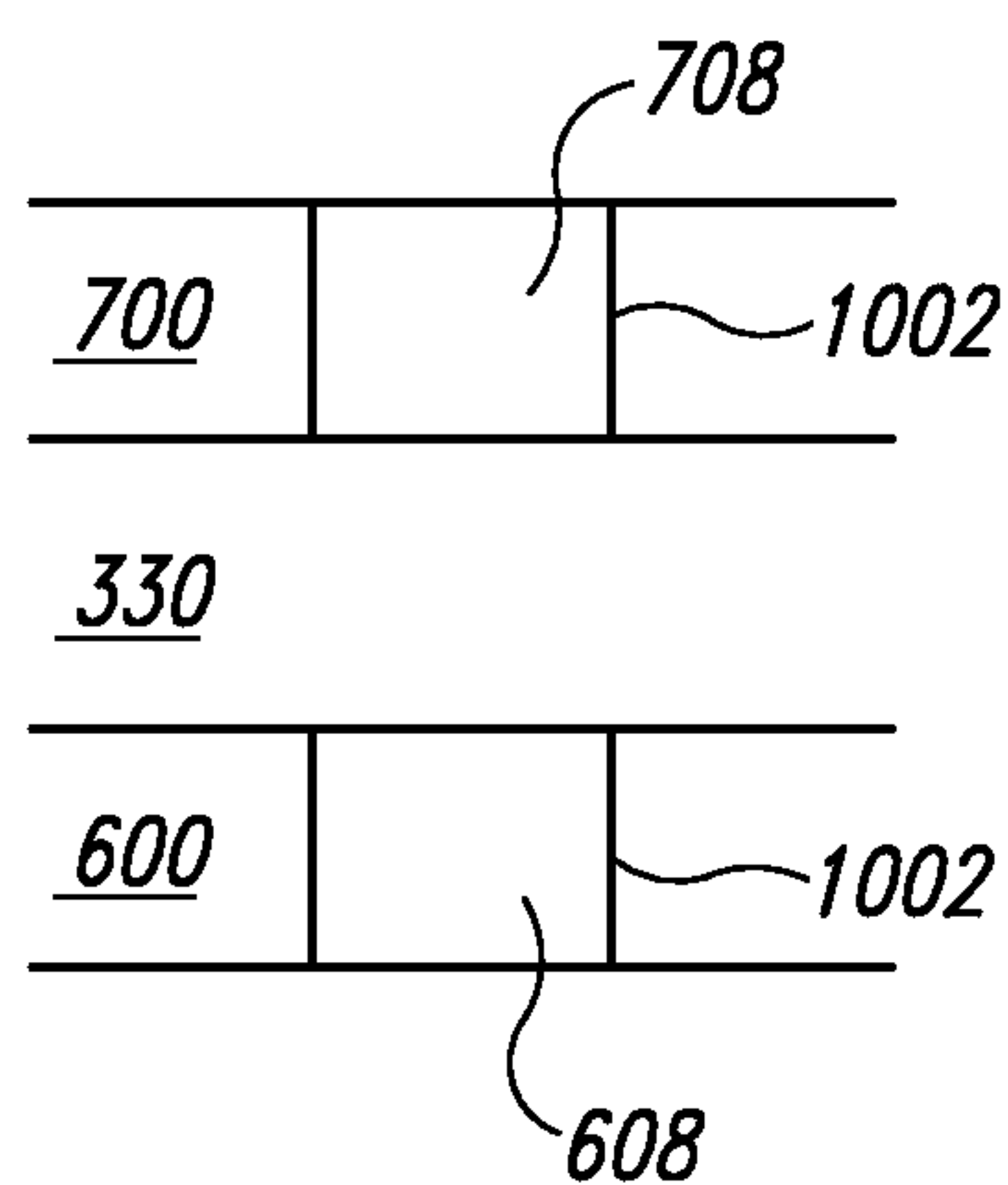
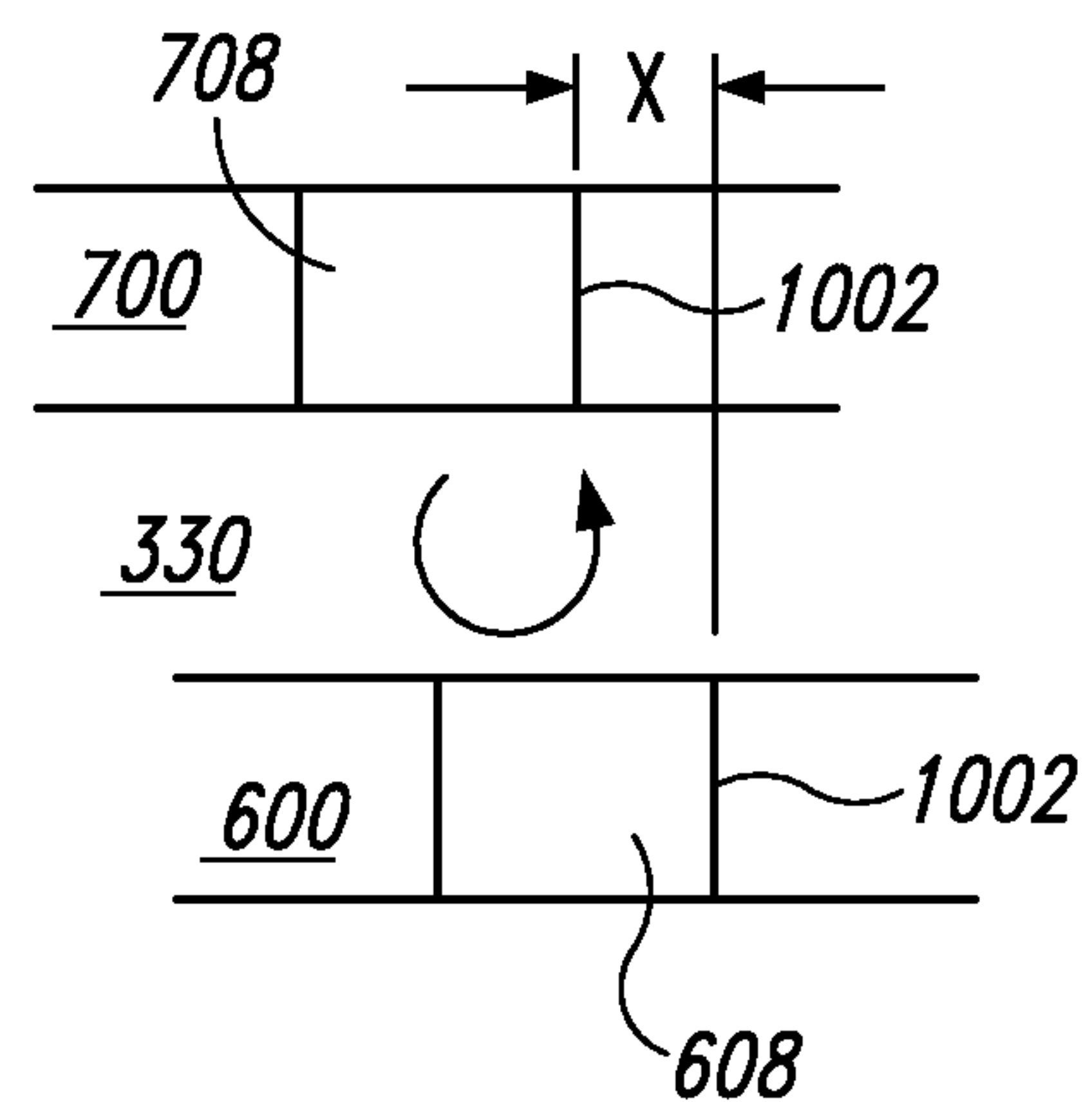
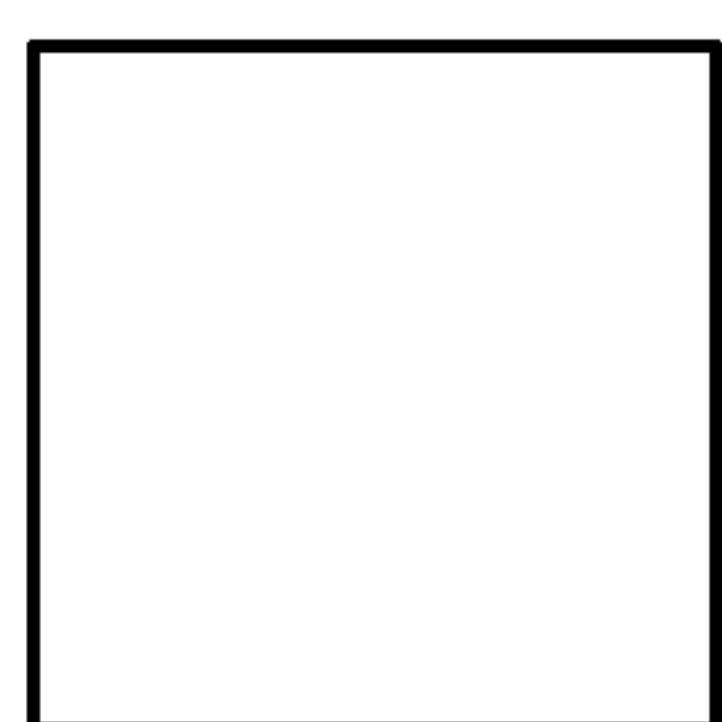
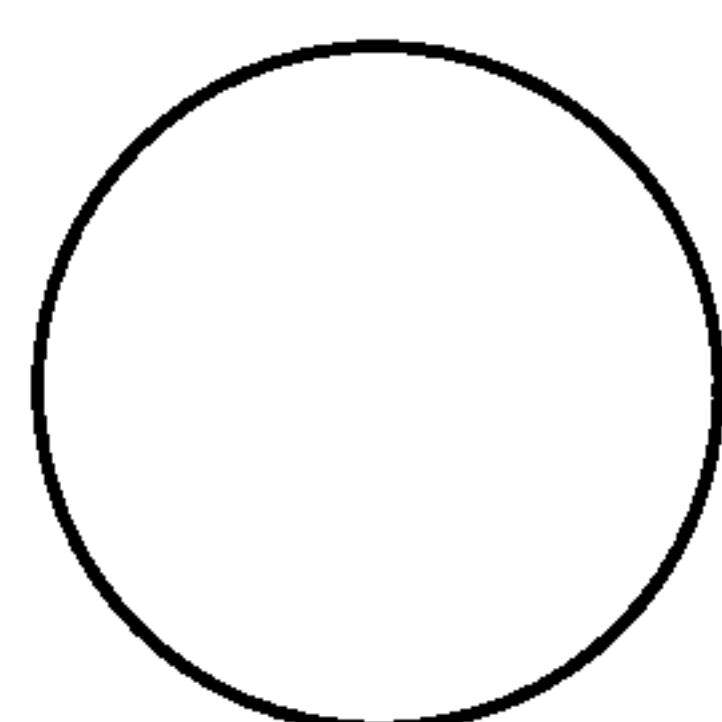
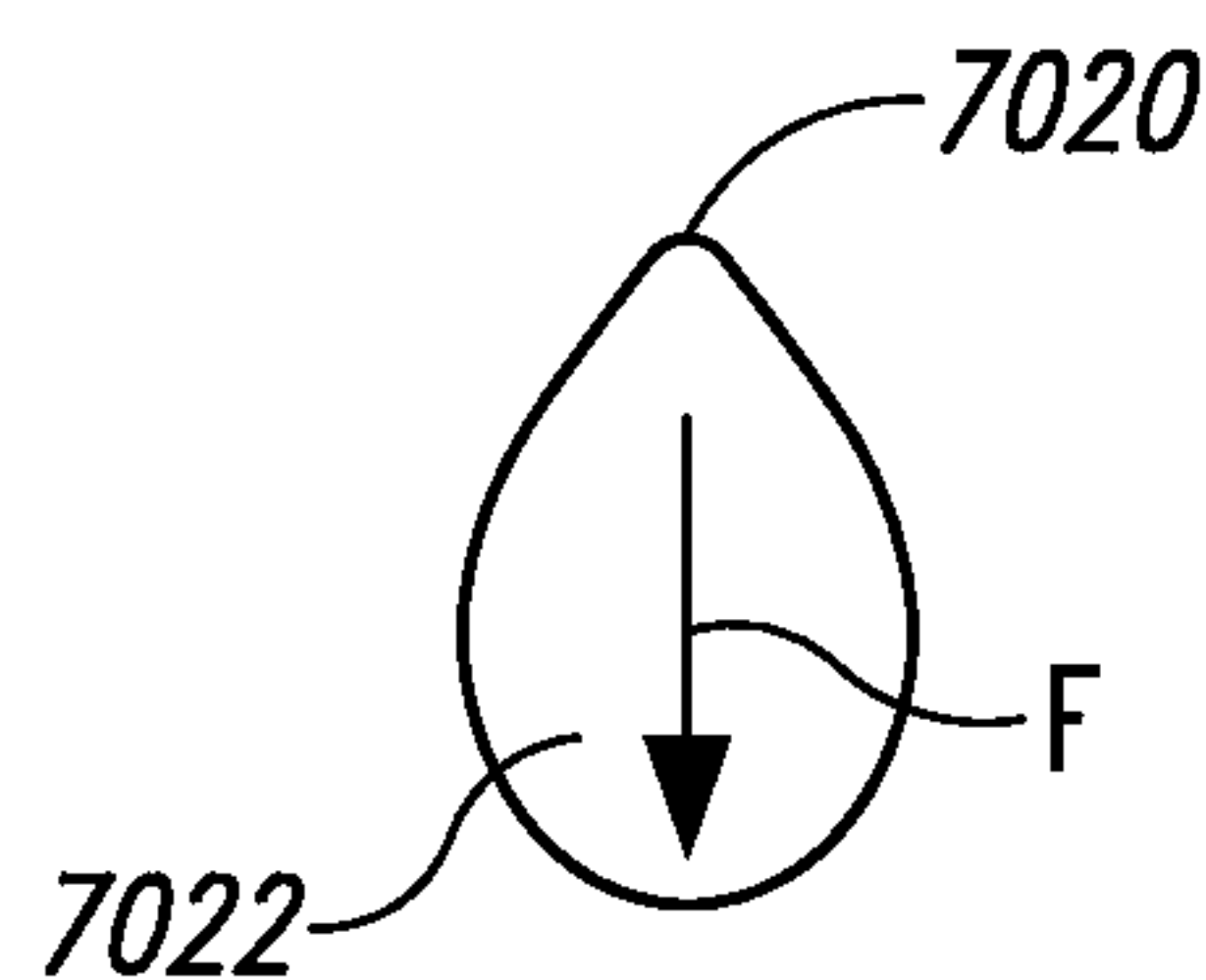
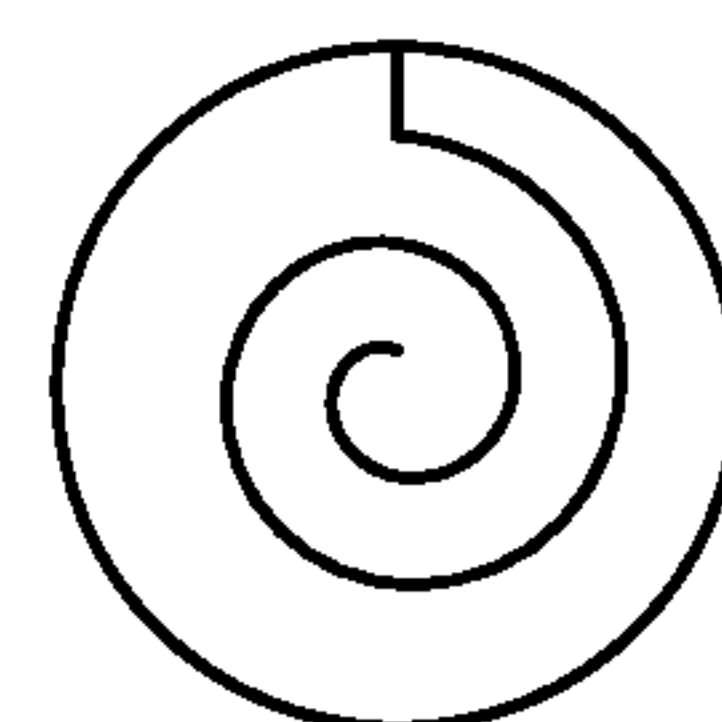
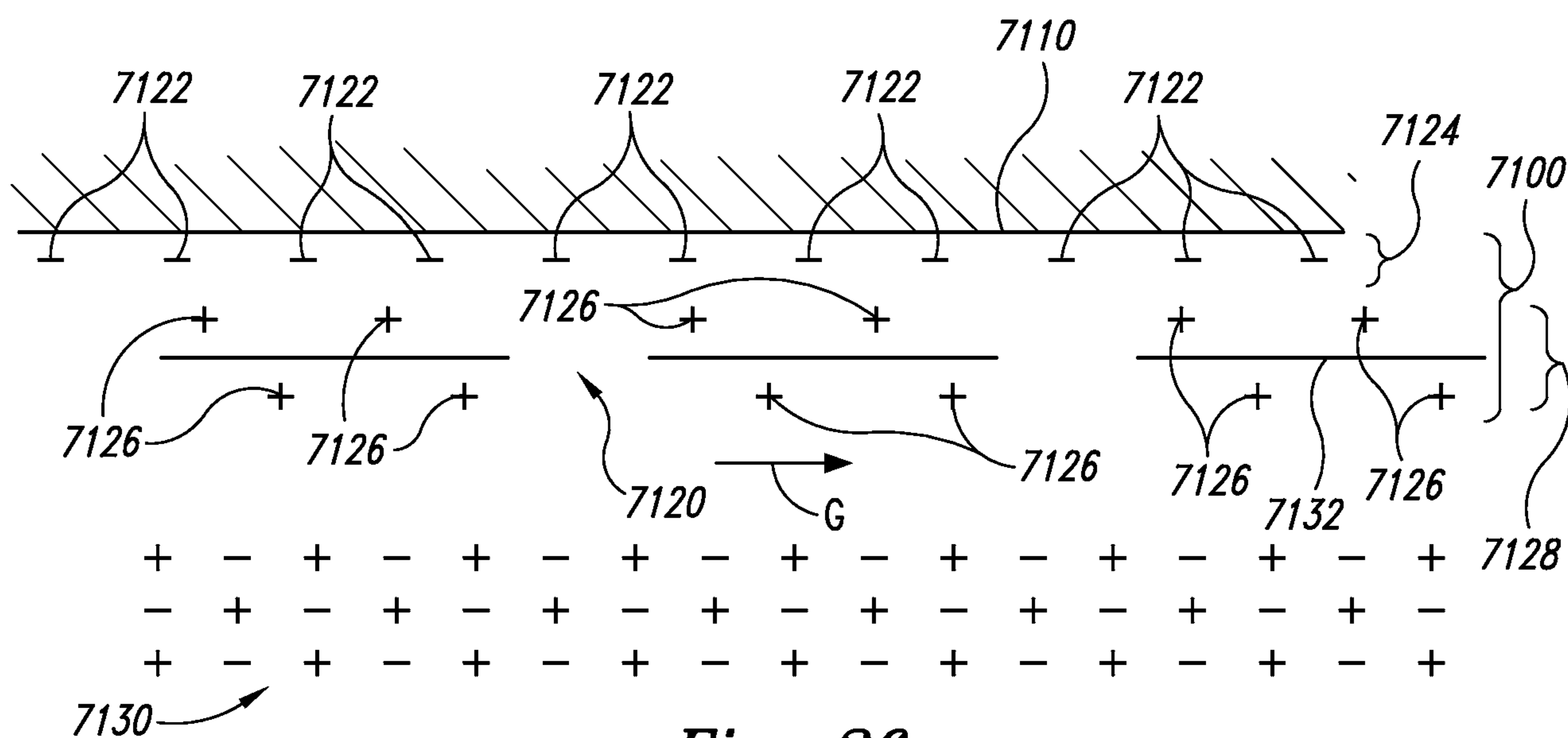
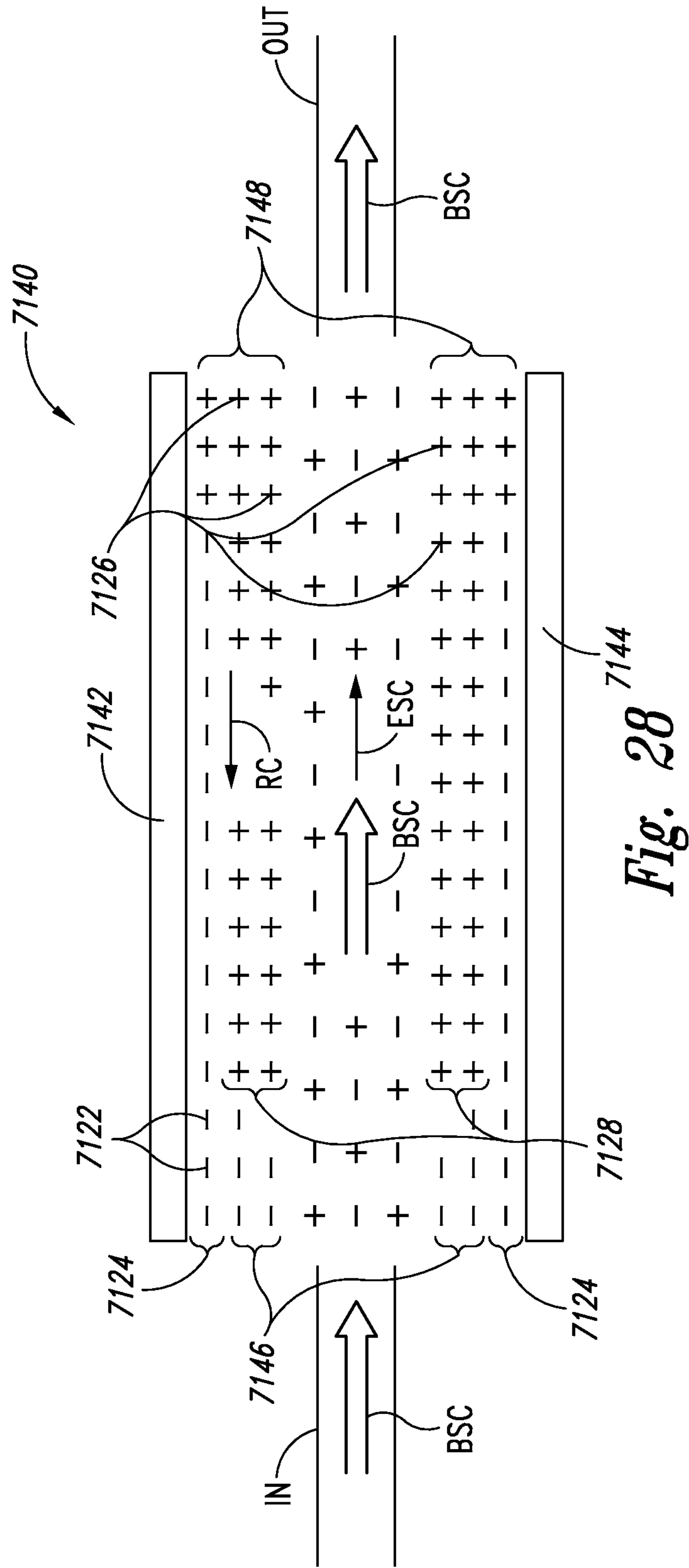
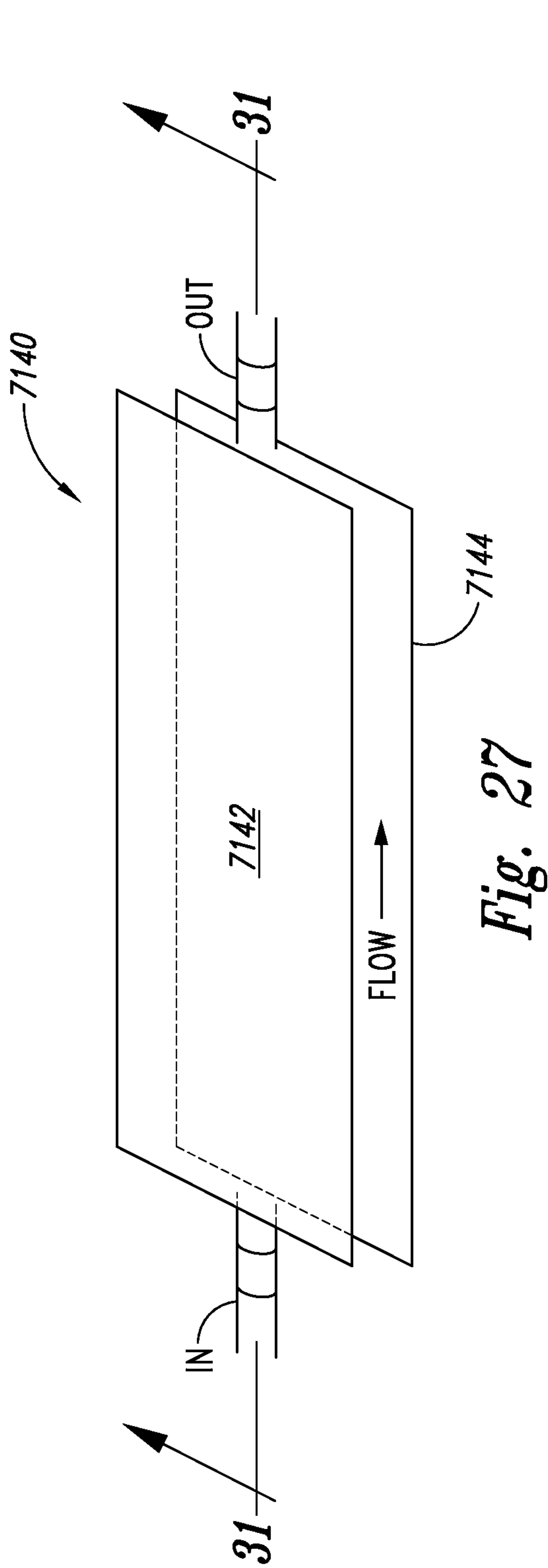


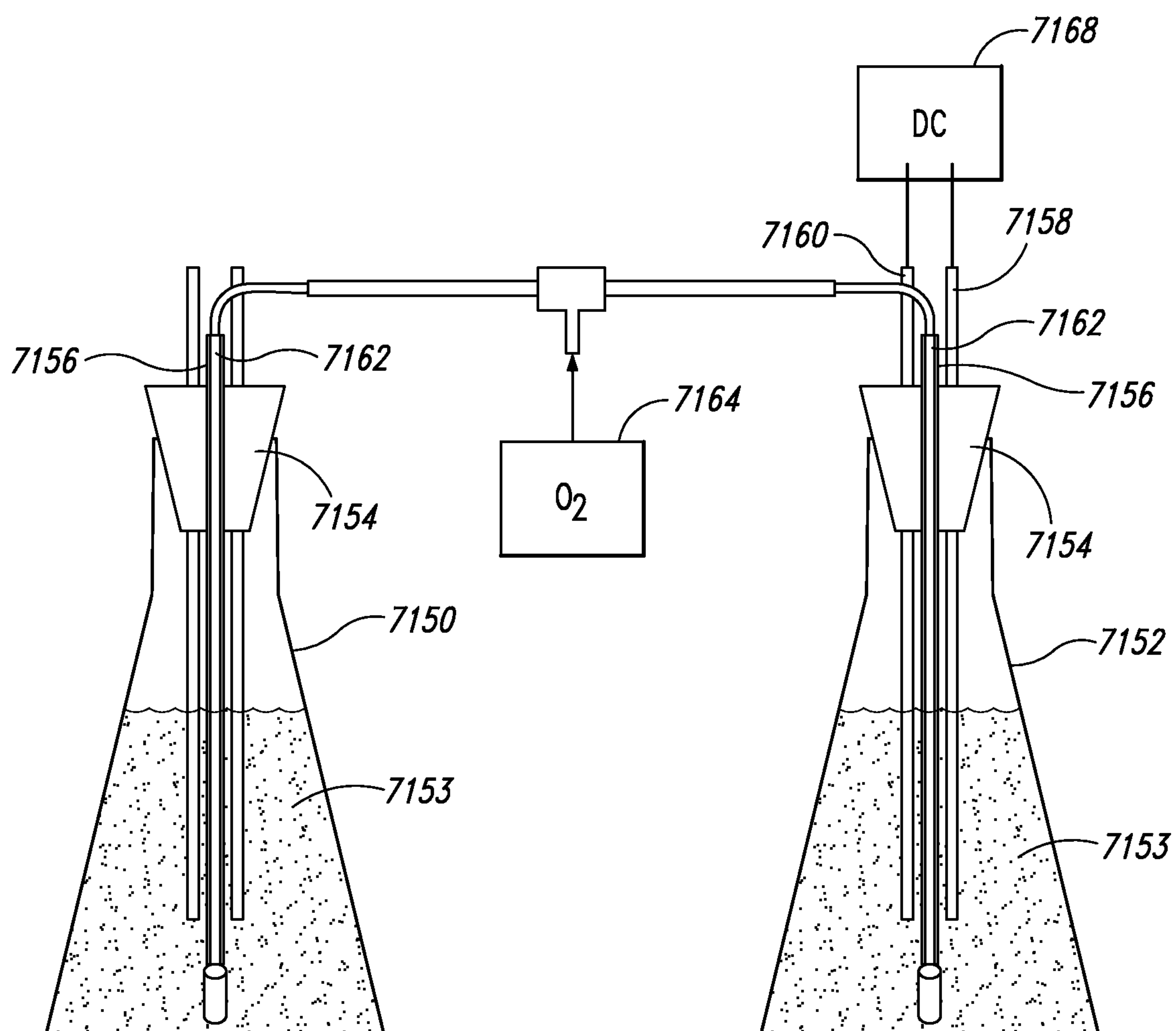
Fig. 18

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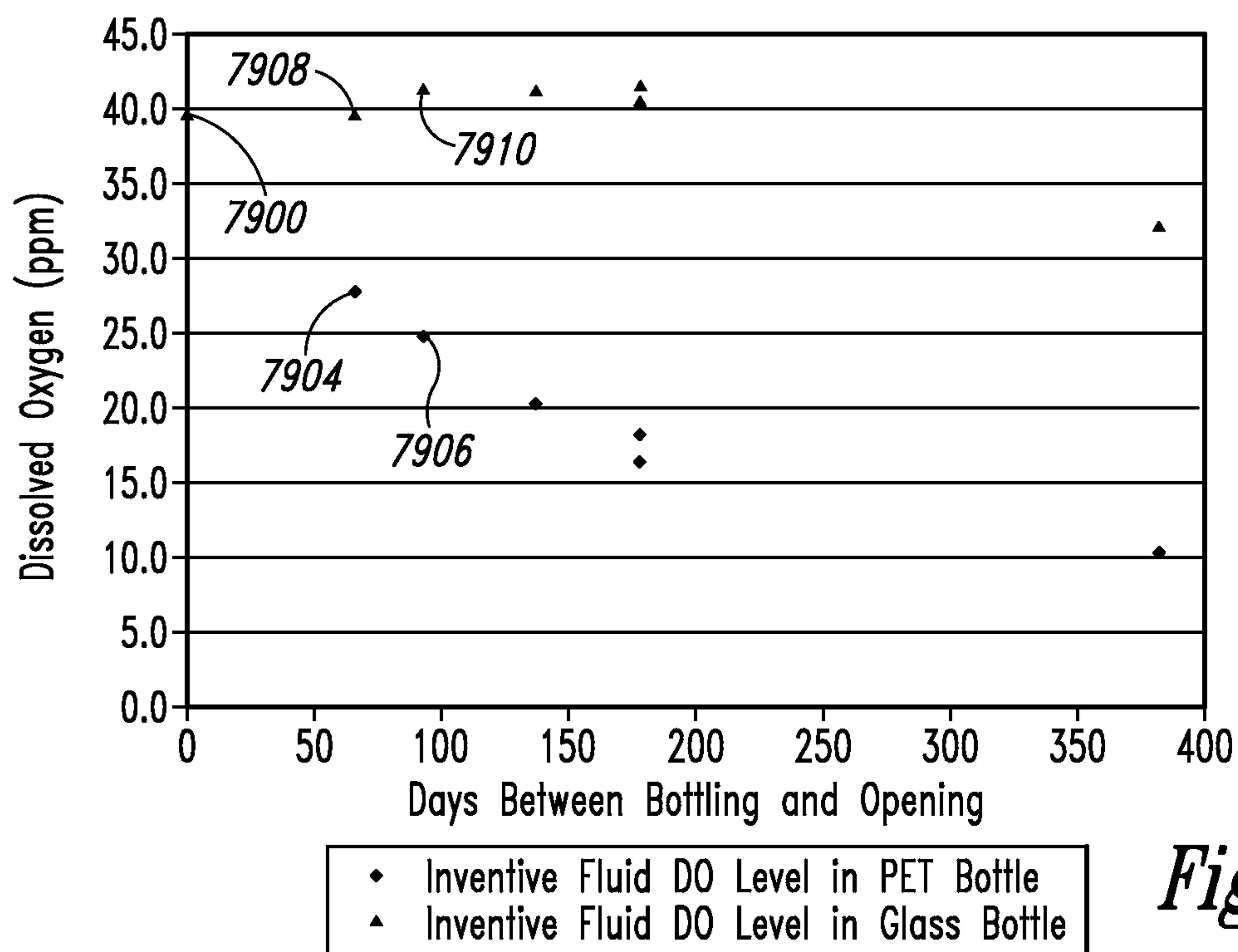
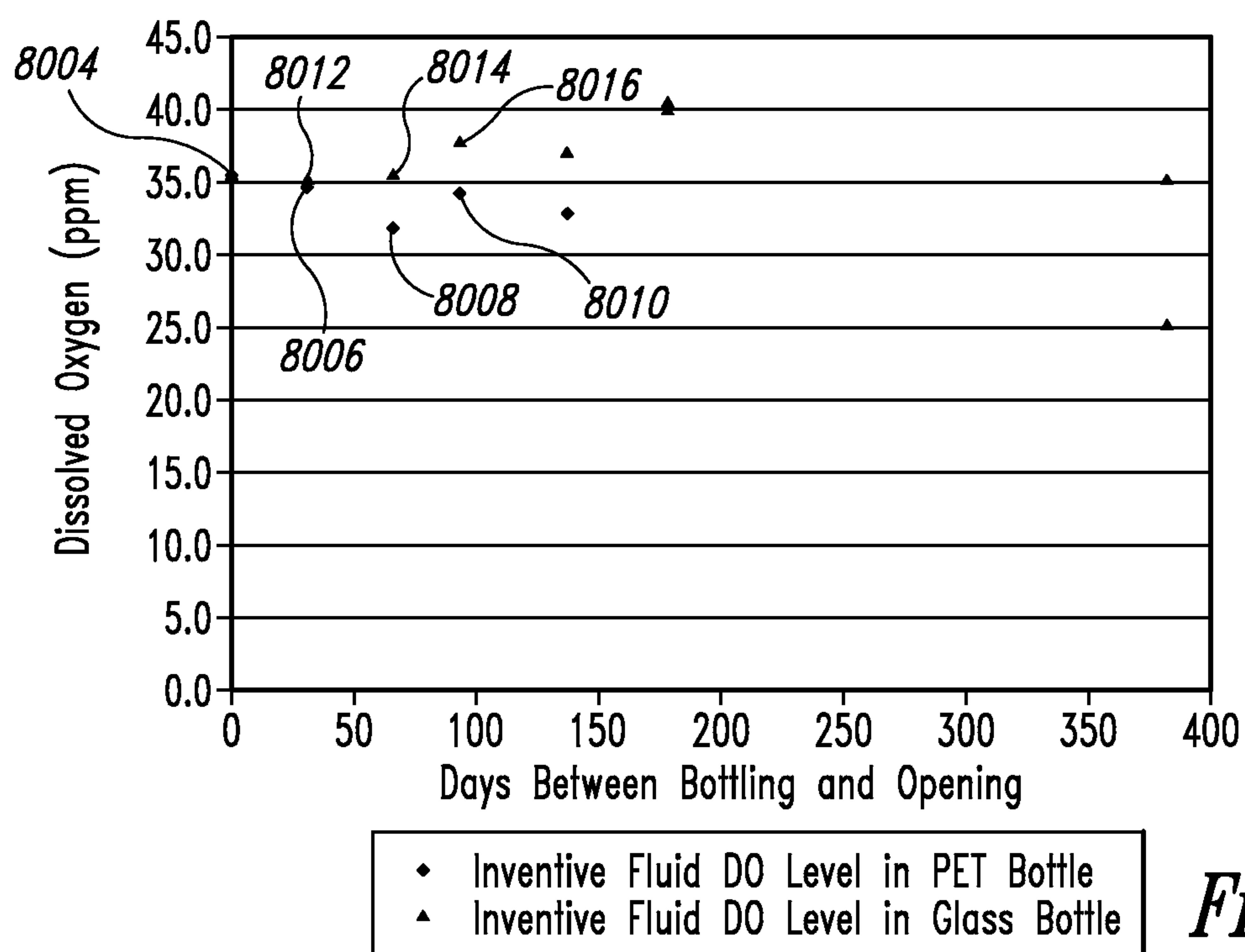
*Fig. 19**Fig. 20**Fig. 21**Fig. 22**Fig. 23**Fig. 24**Fig. 25**Fig. 26*



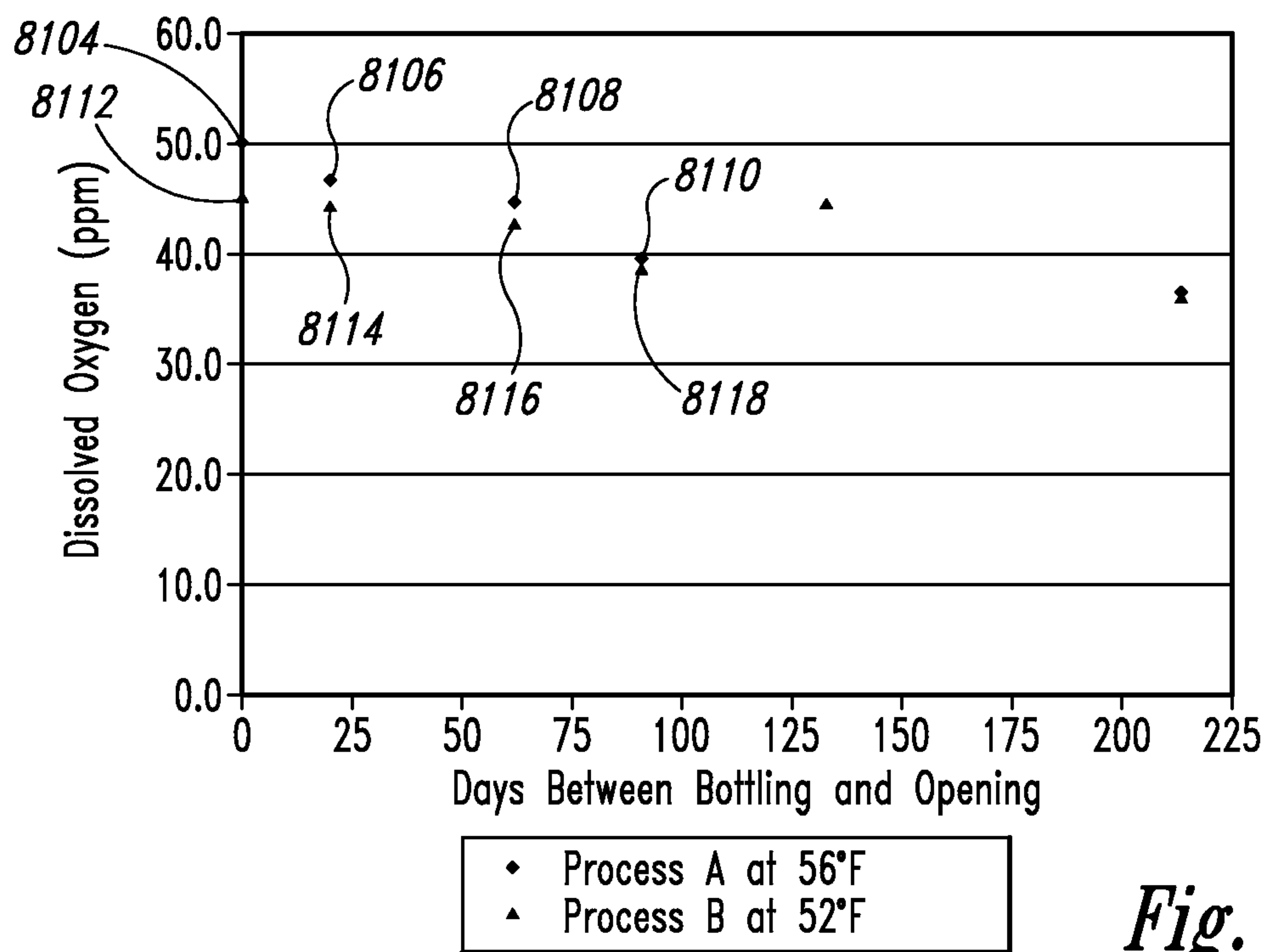
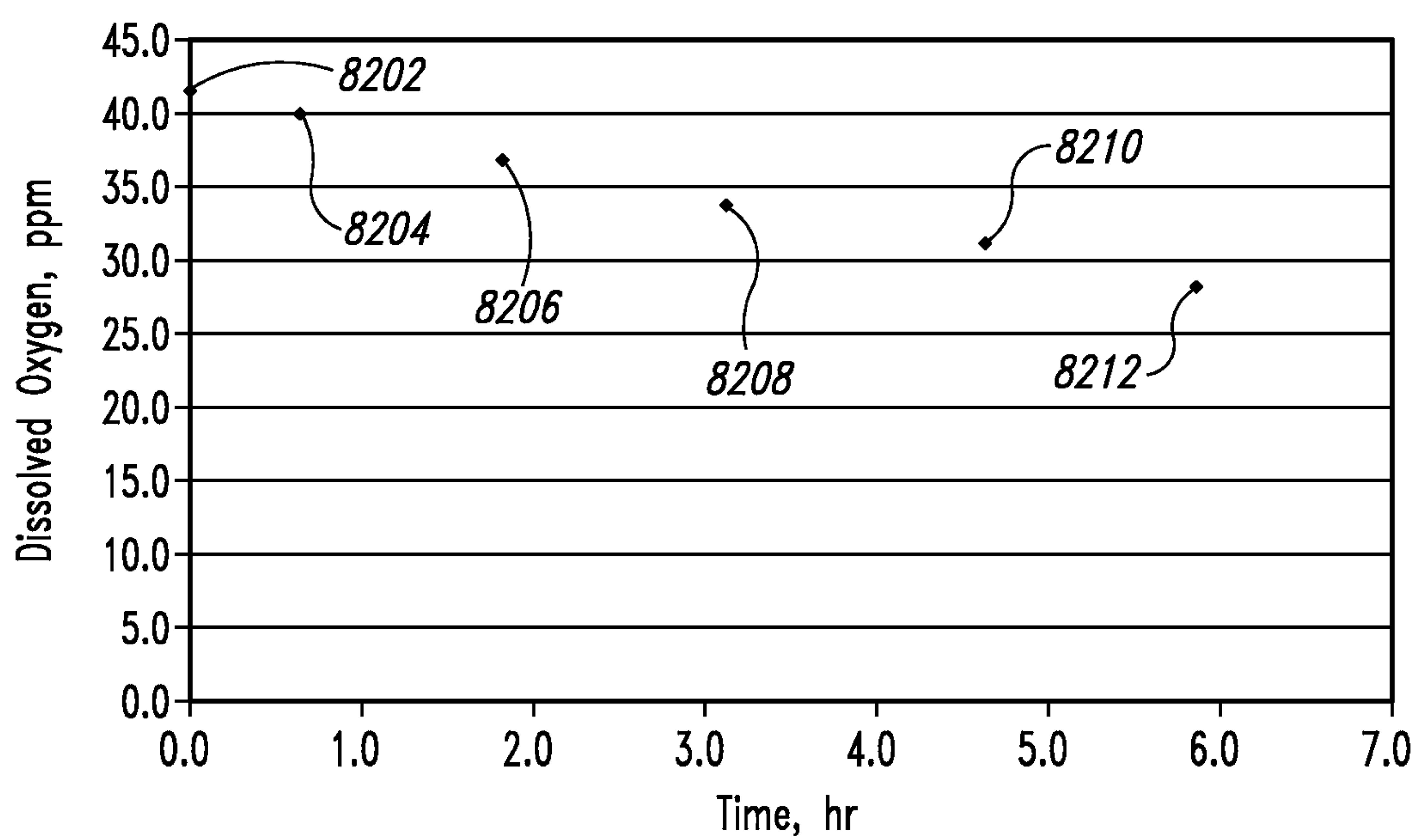
16/94

*Fig. 29*

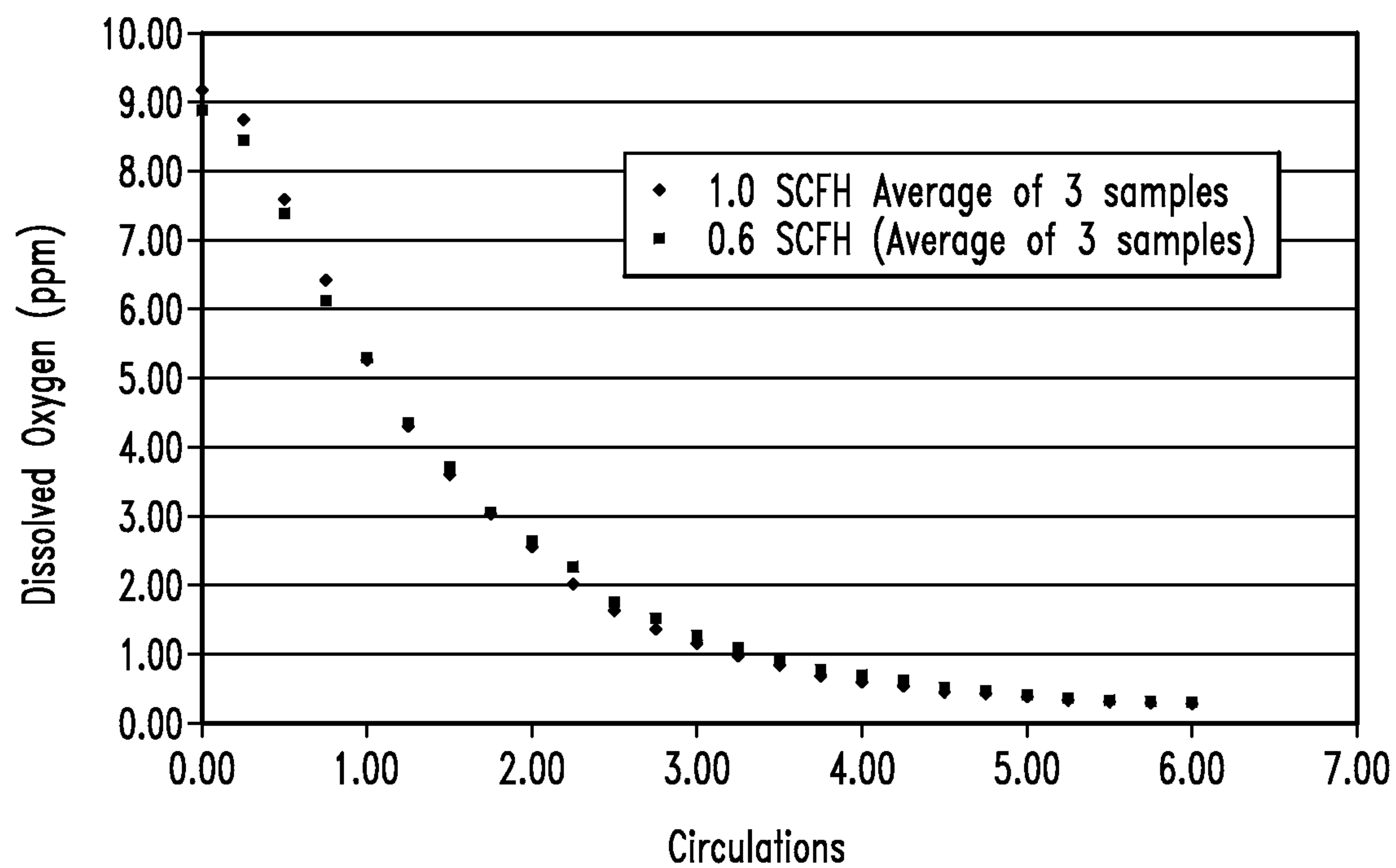
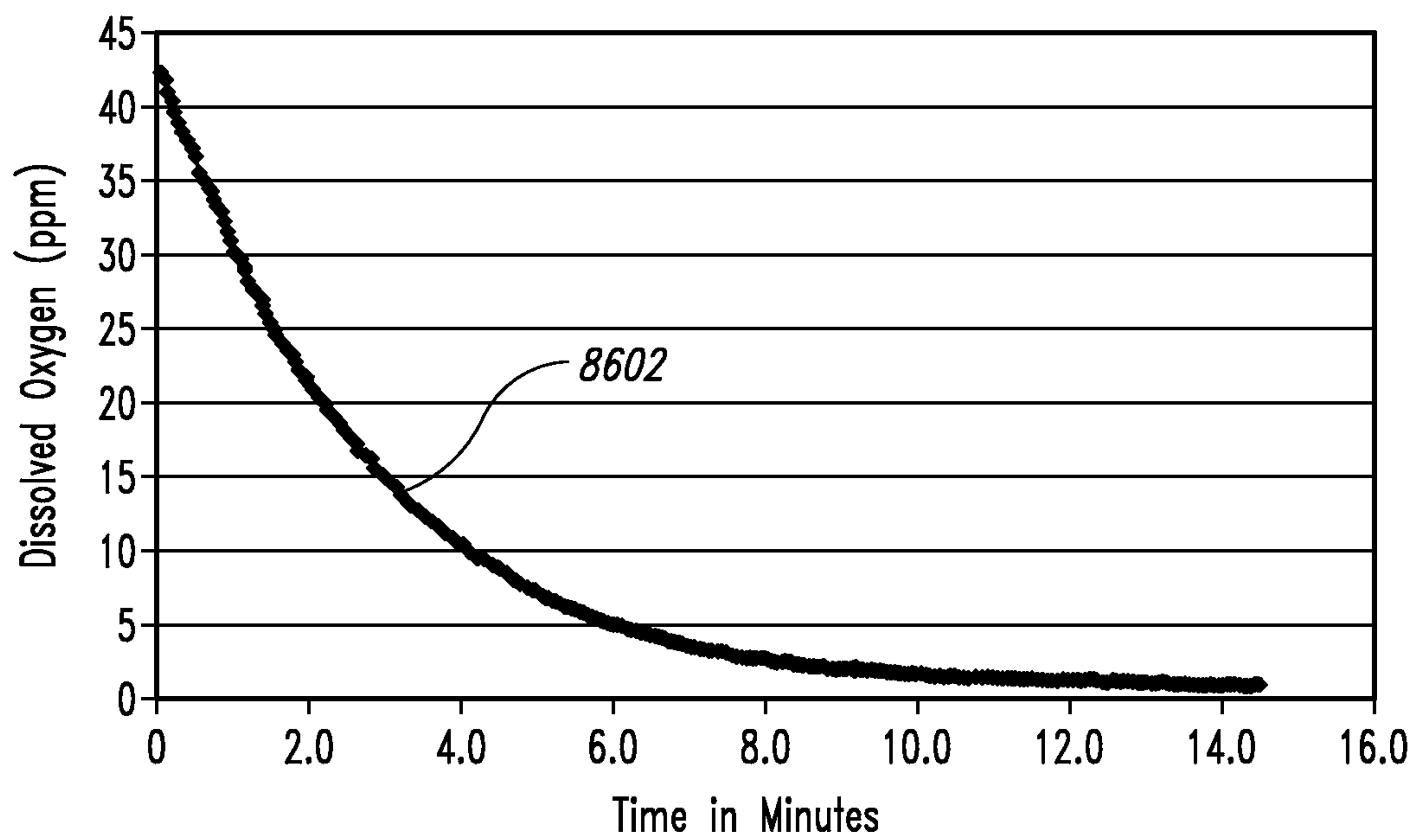
17/94

*Fig. 30**Fig. 31*

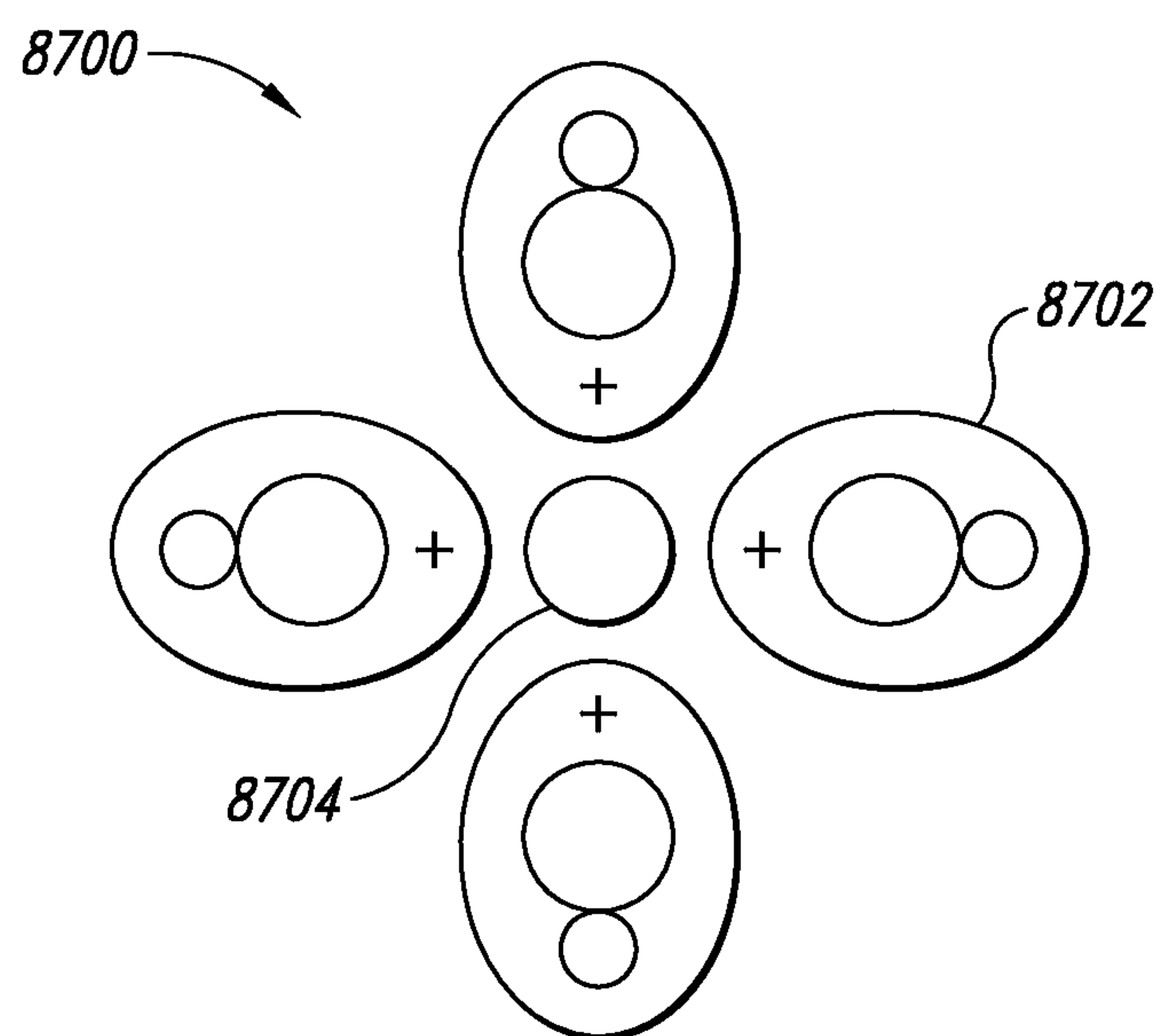
18/94

*Fig. 32**Fig. 33*

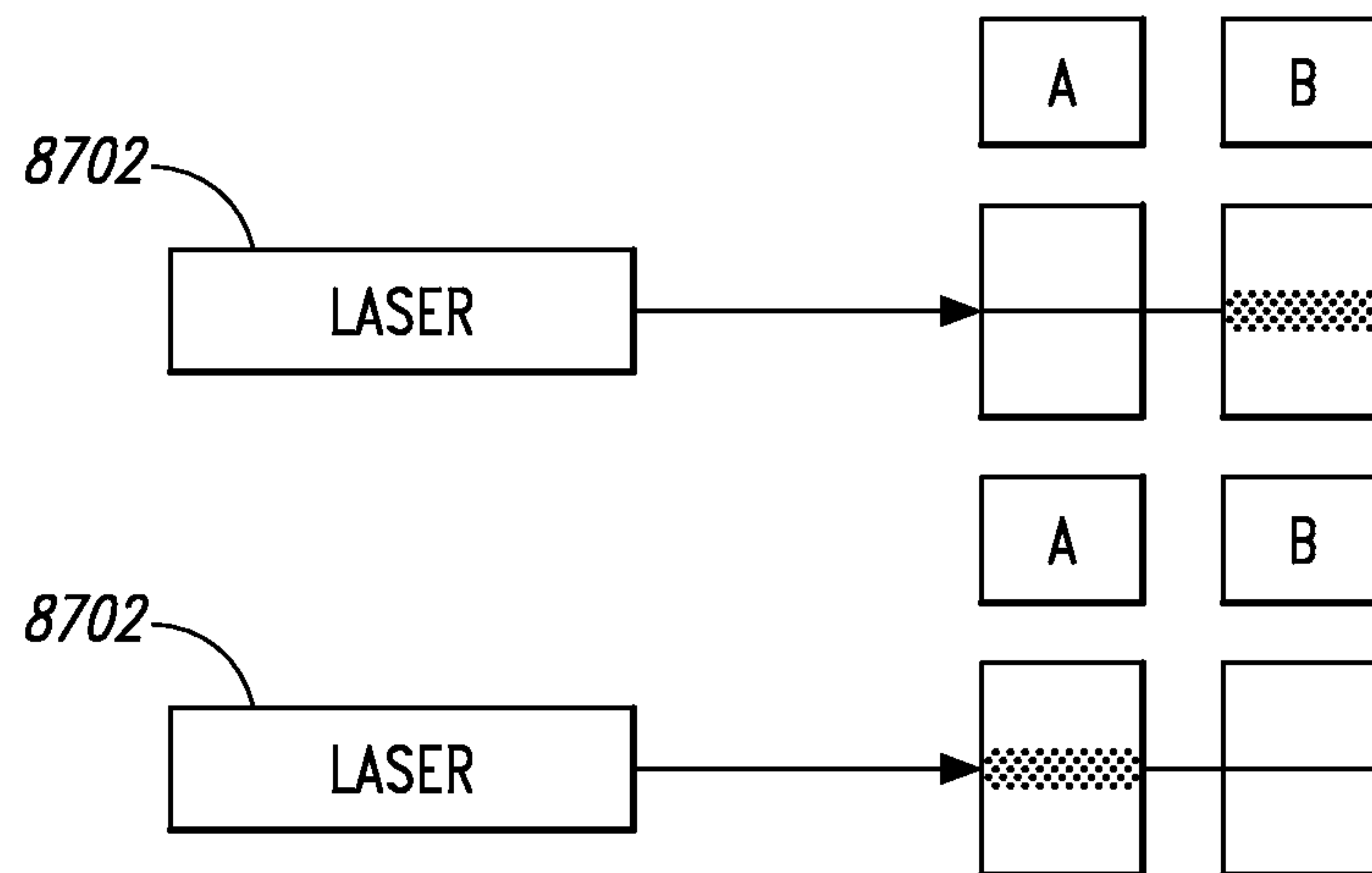
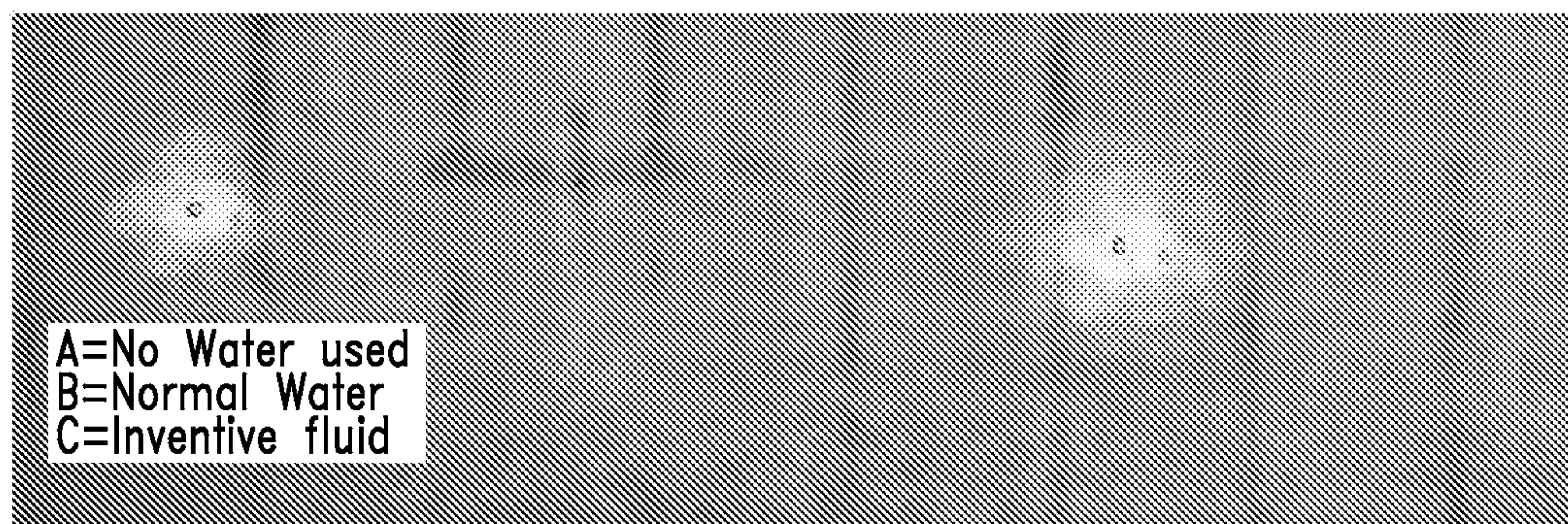
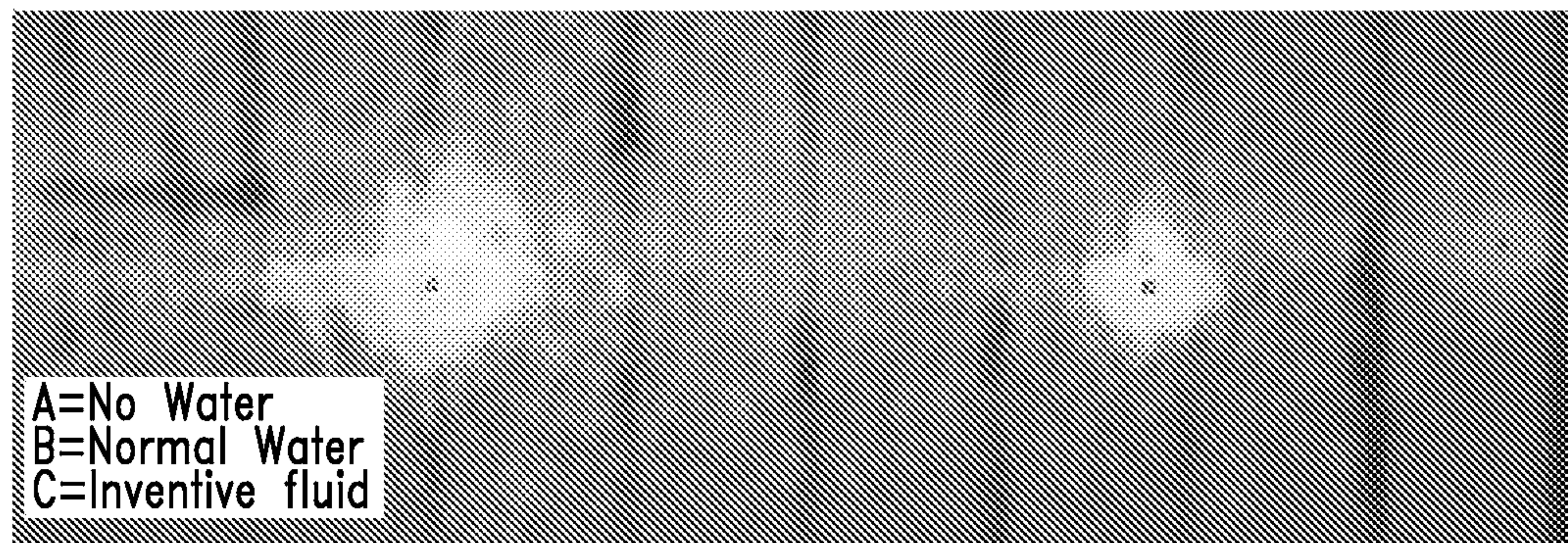
19/94

*Fig. 34**Fig. 35*

20/94

*Fig. 36*

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*Fig. 37A**Fig. 37B*

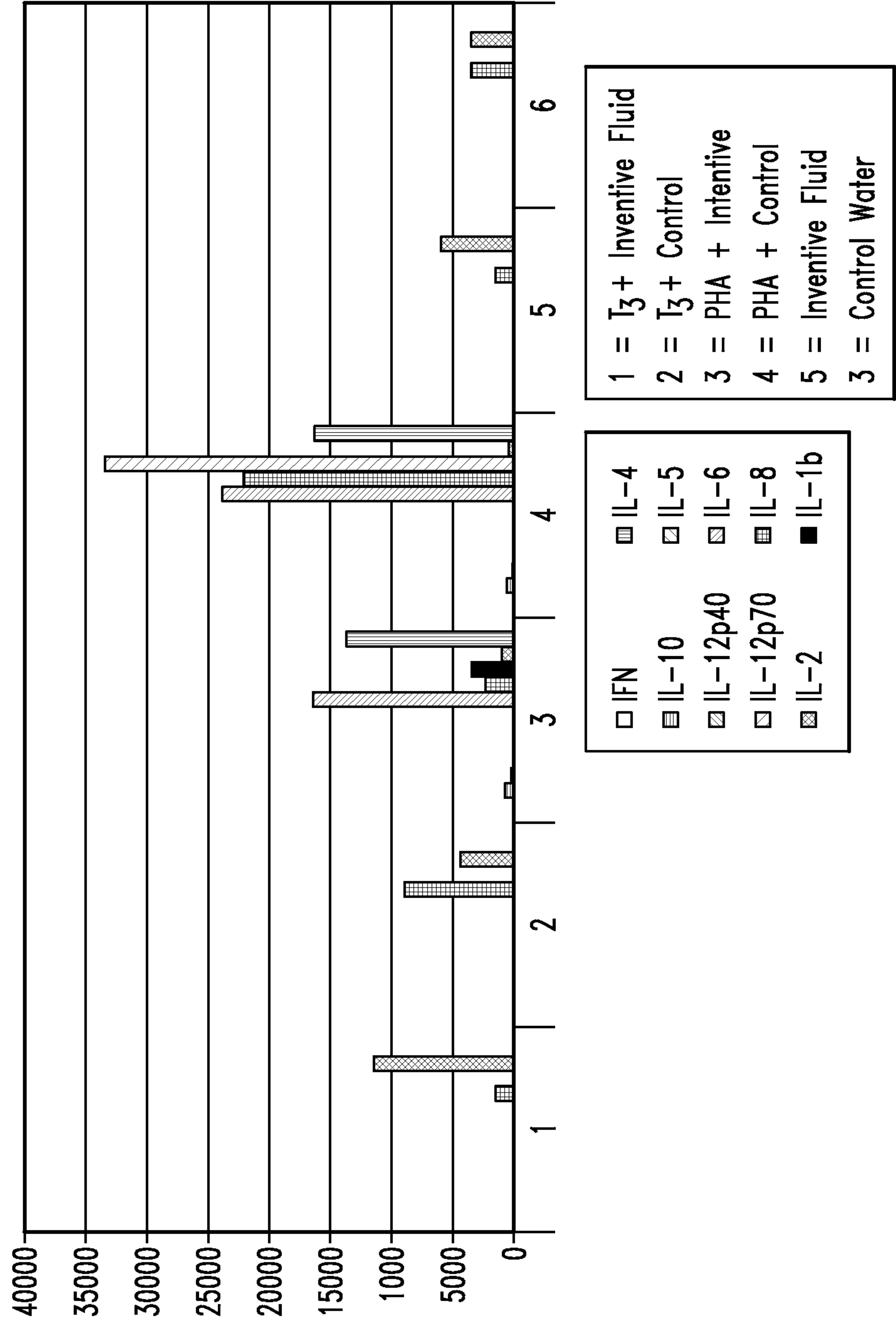
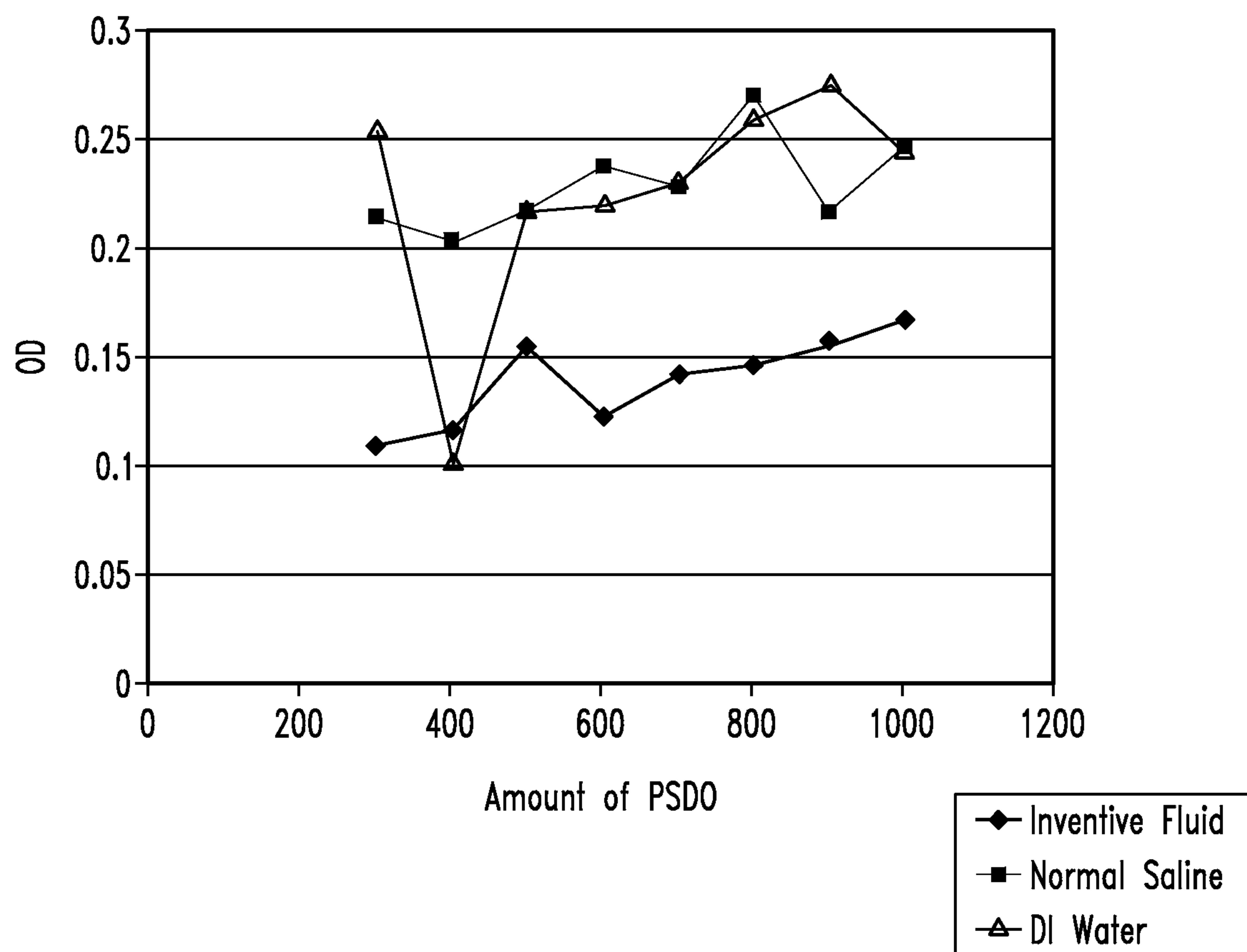
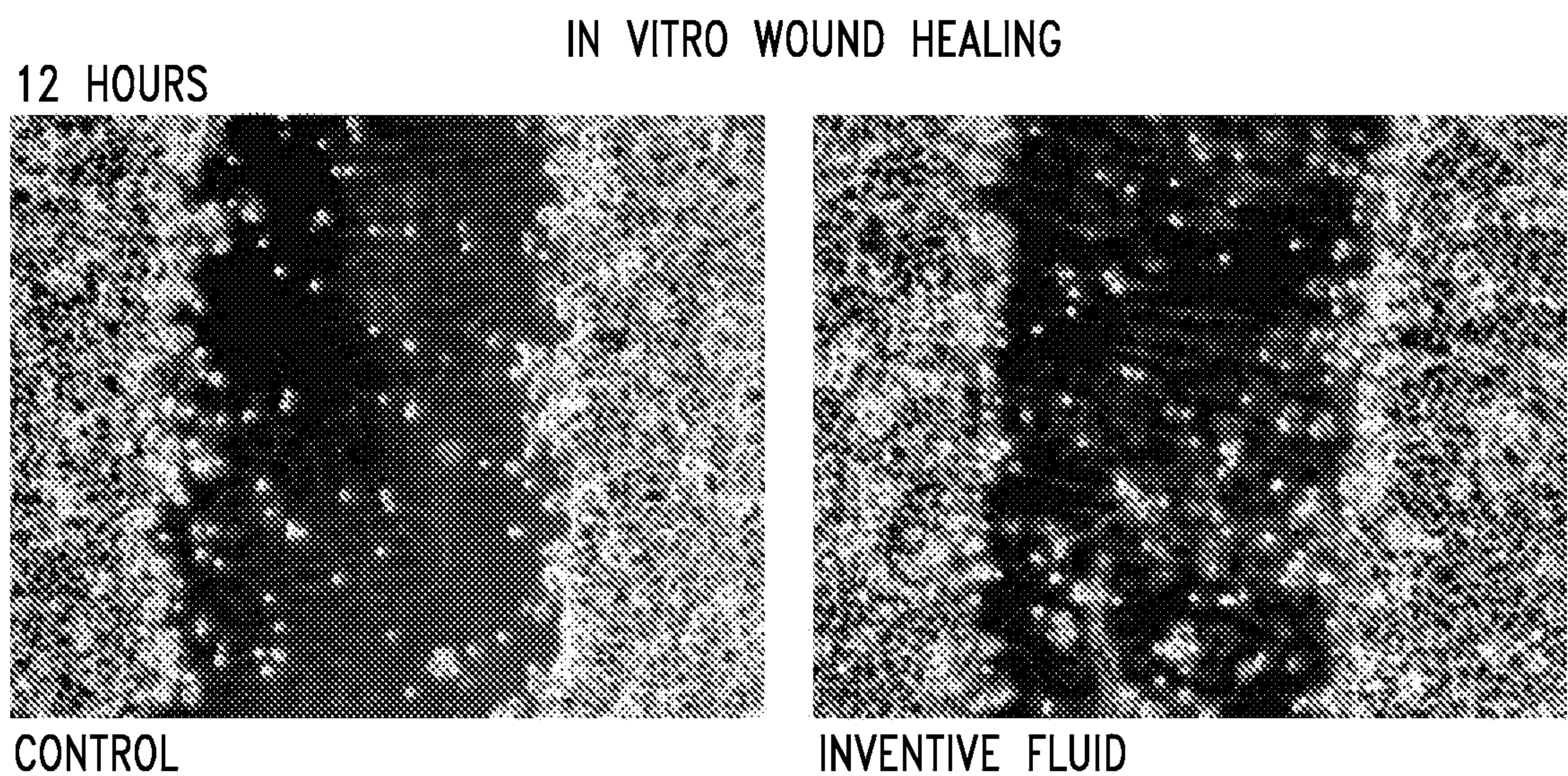
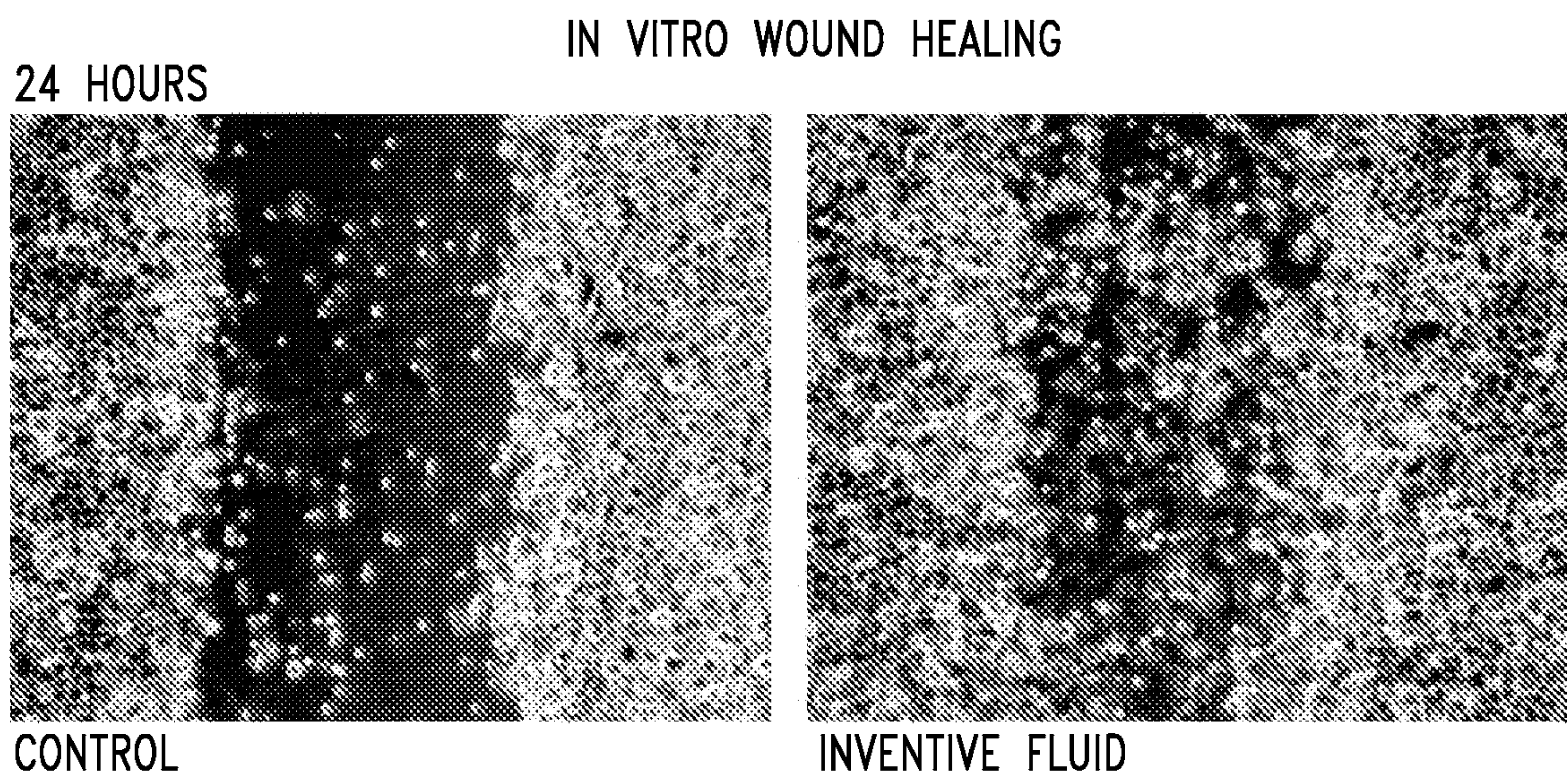


Fig. 38

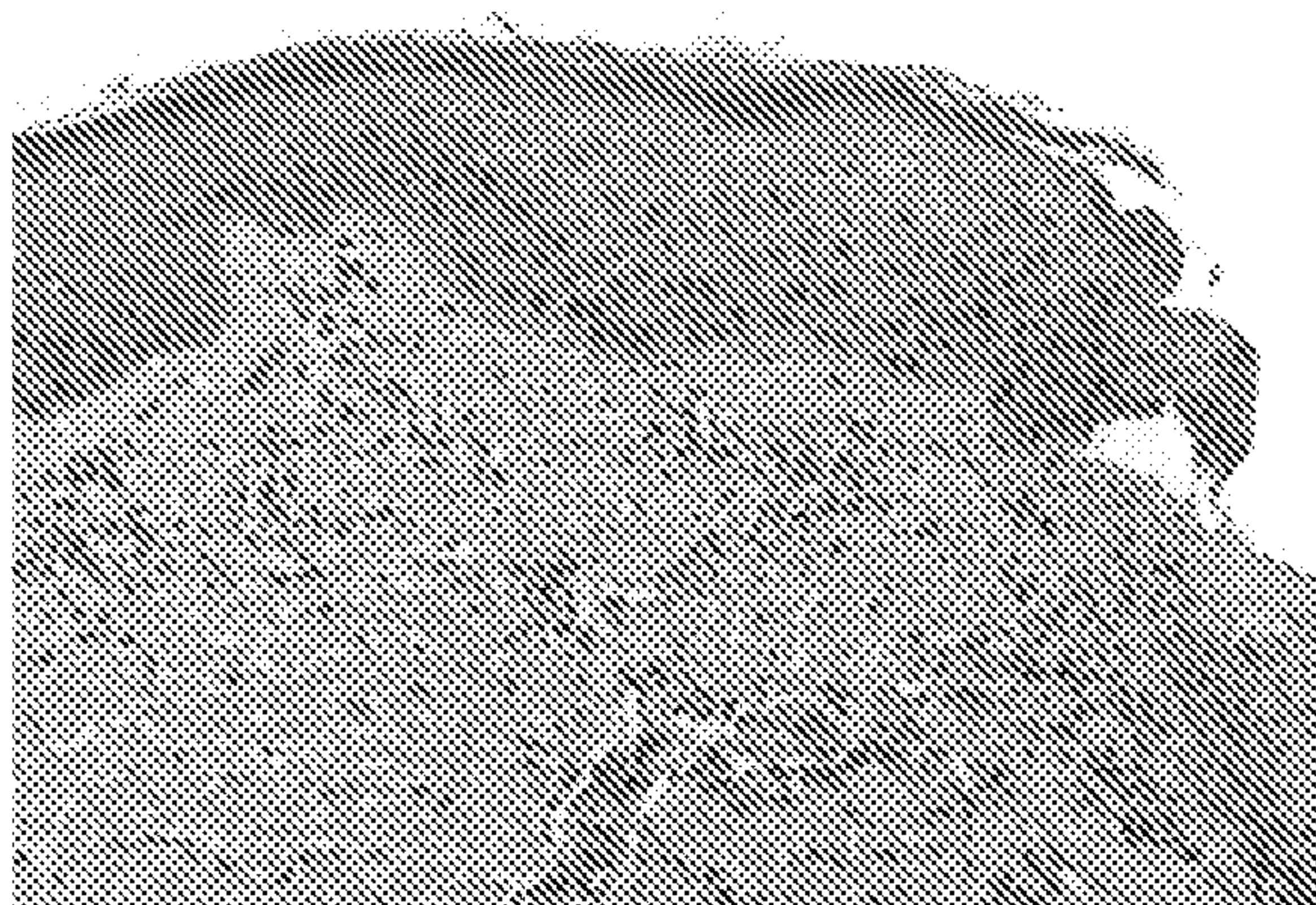
23/94

*Fig. 39*

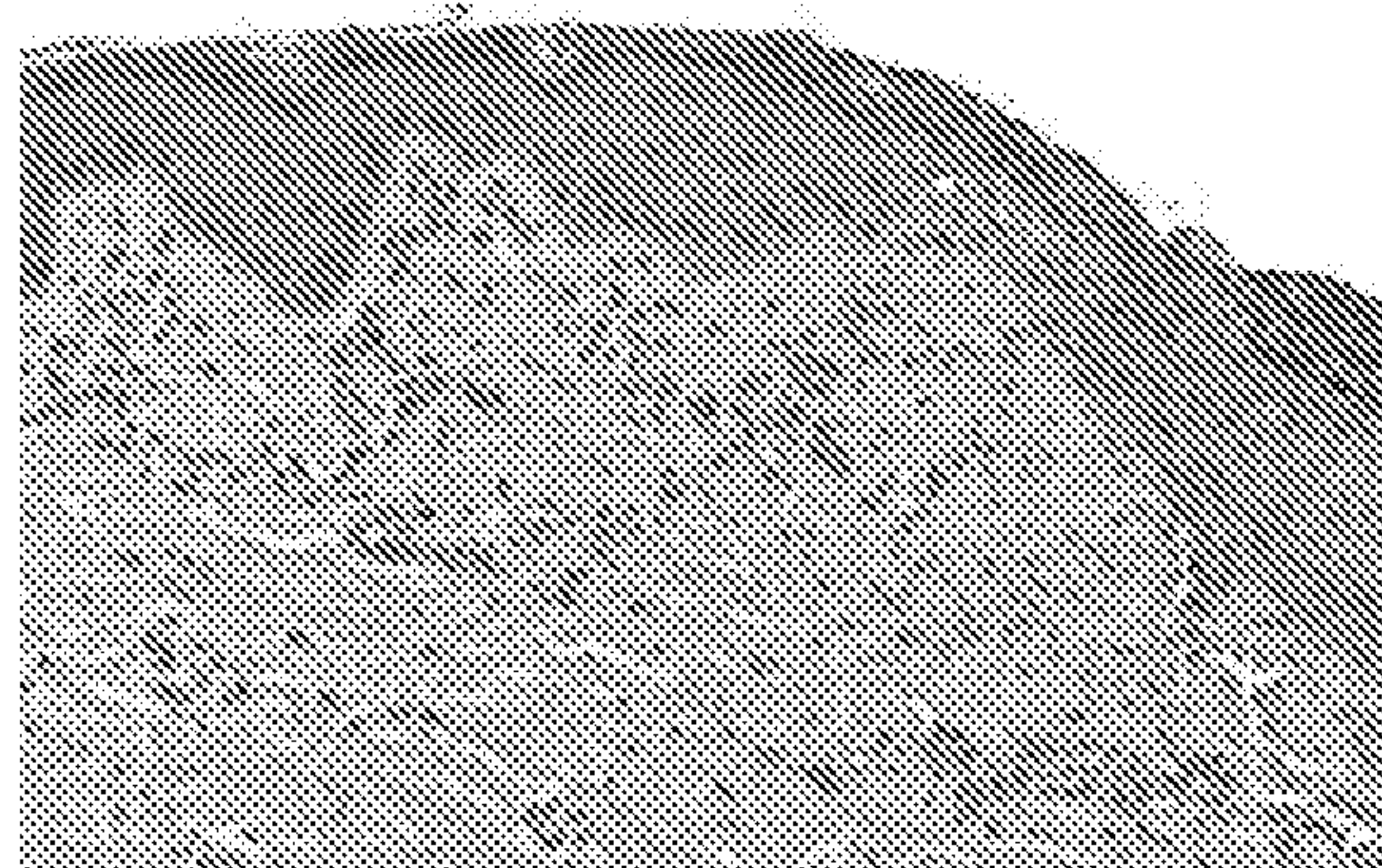
24/94

*Fig. 40A**Fig. 40B*

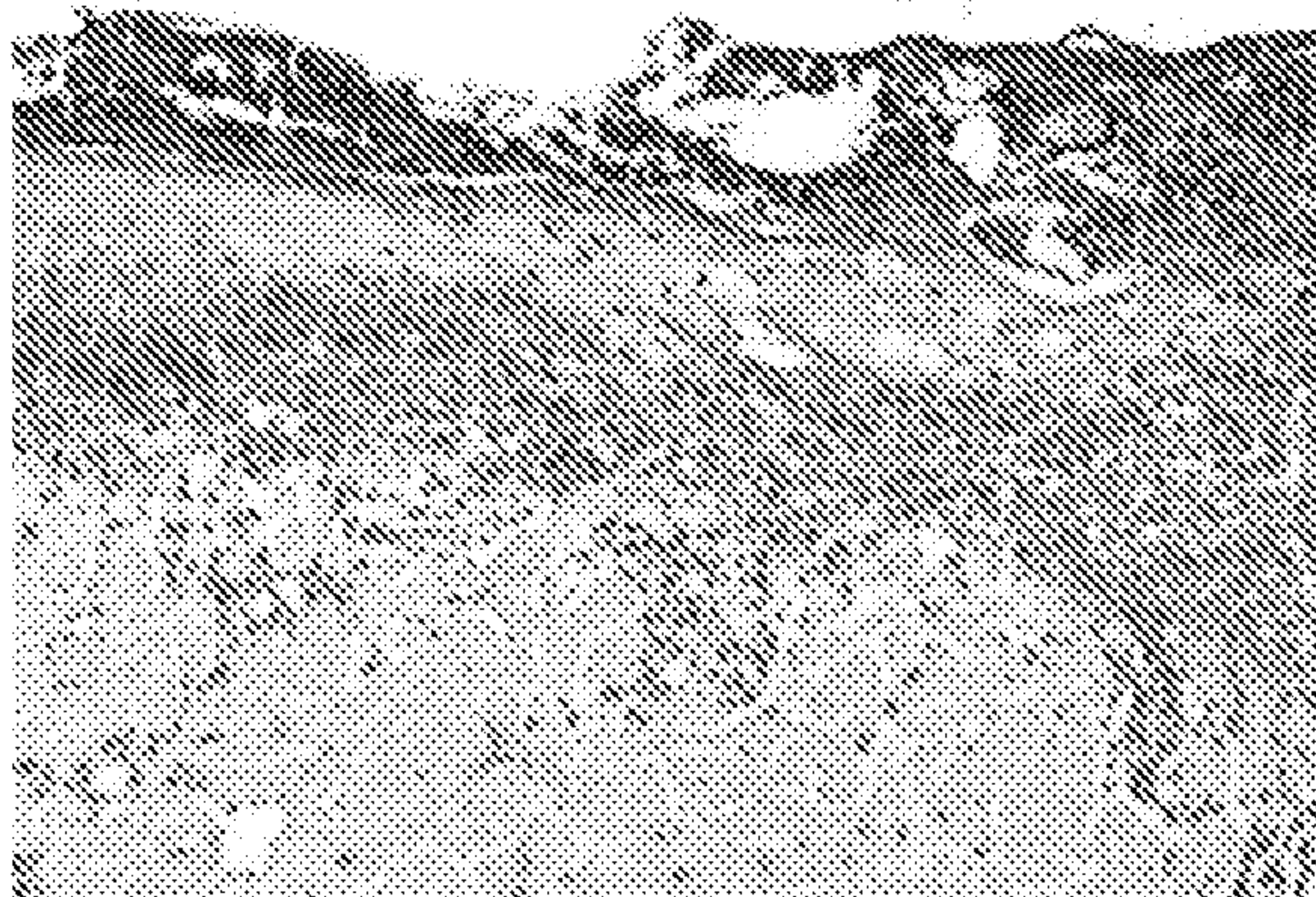
25/94



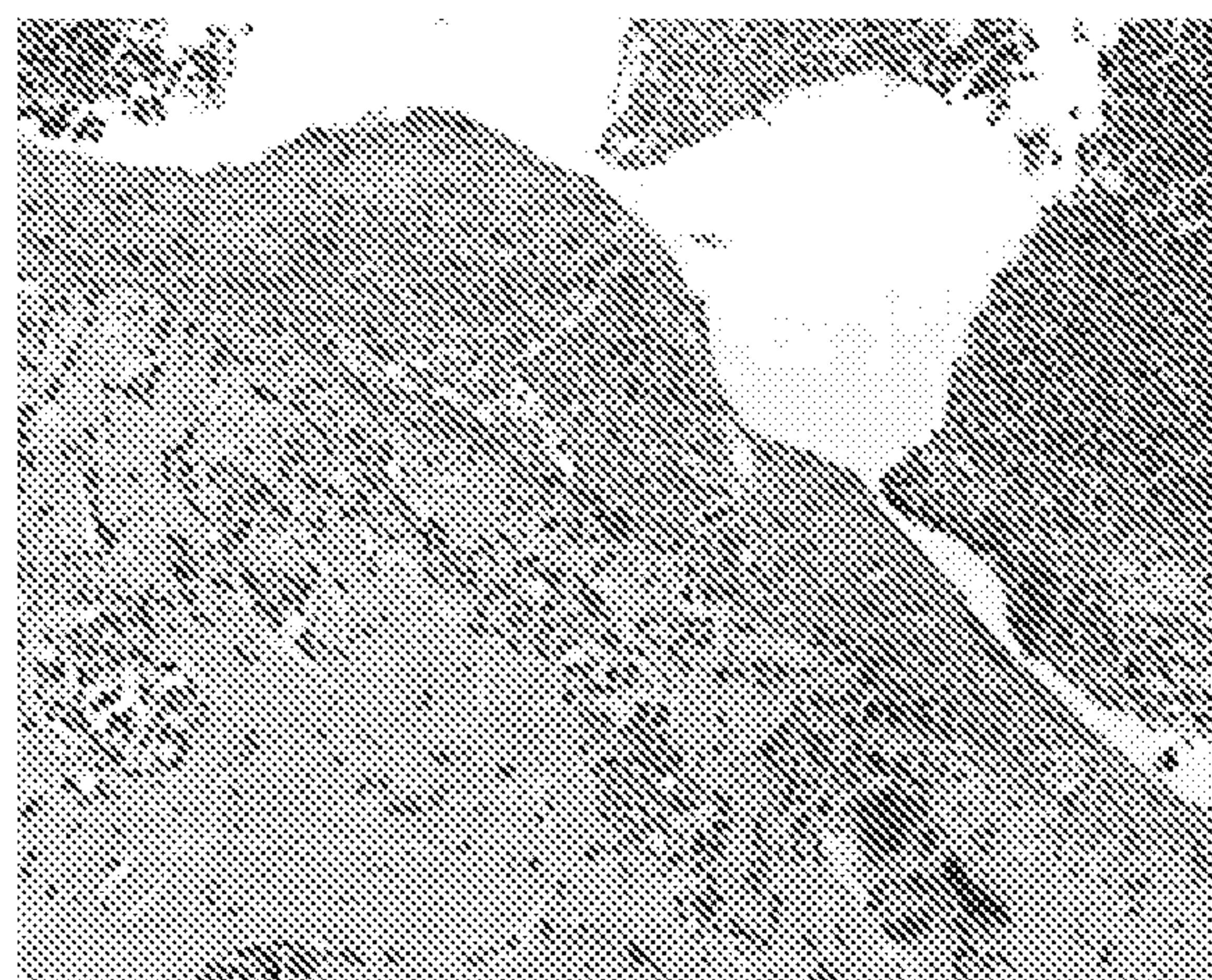
Control one day post wounding
showing epidermal/dermal thickening
and loss of contour

Fig. 41A

Experimental with inventive fluid one day post
wounding showing normal epidermal/dermal
thickness and normal contour

Fig. 41B

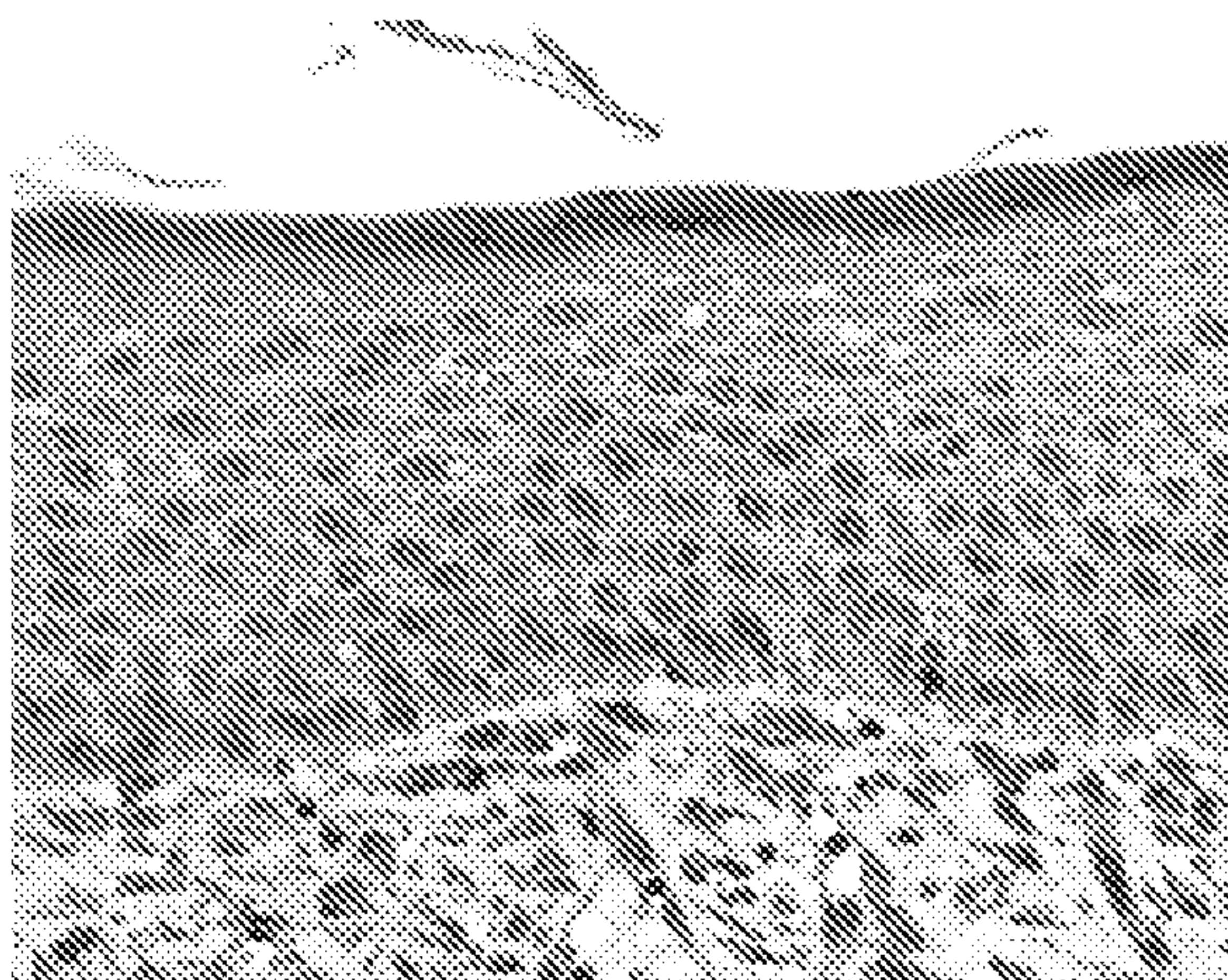
Control day four showing 600 micron
epidermal spur

Fig. 41C

Experimental day four with inventive fluid
showing 1200 micron epidermal spur

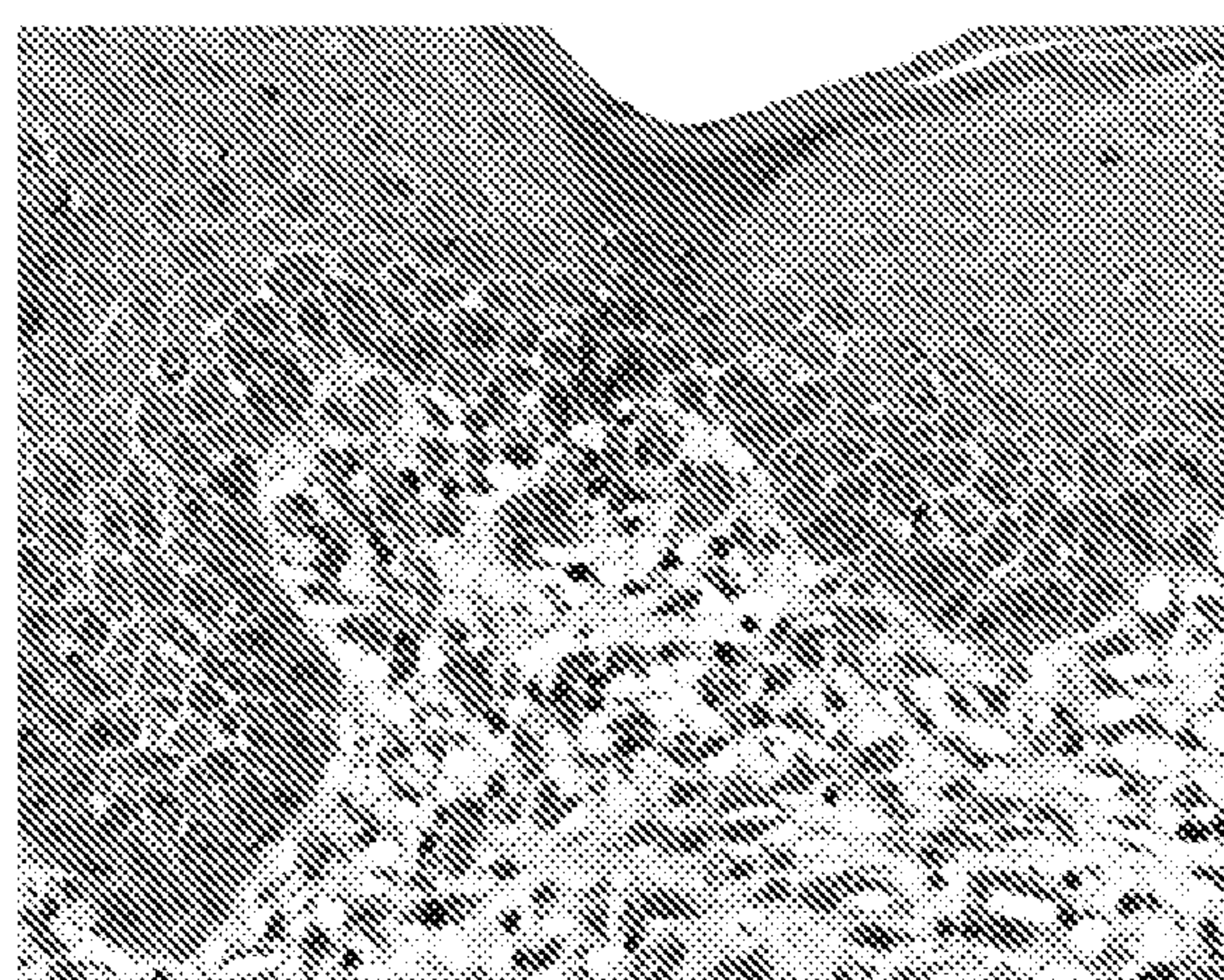
Fig. 41D

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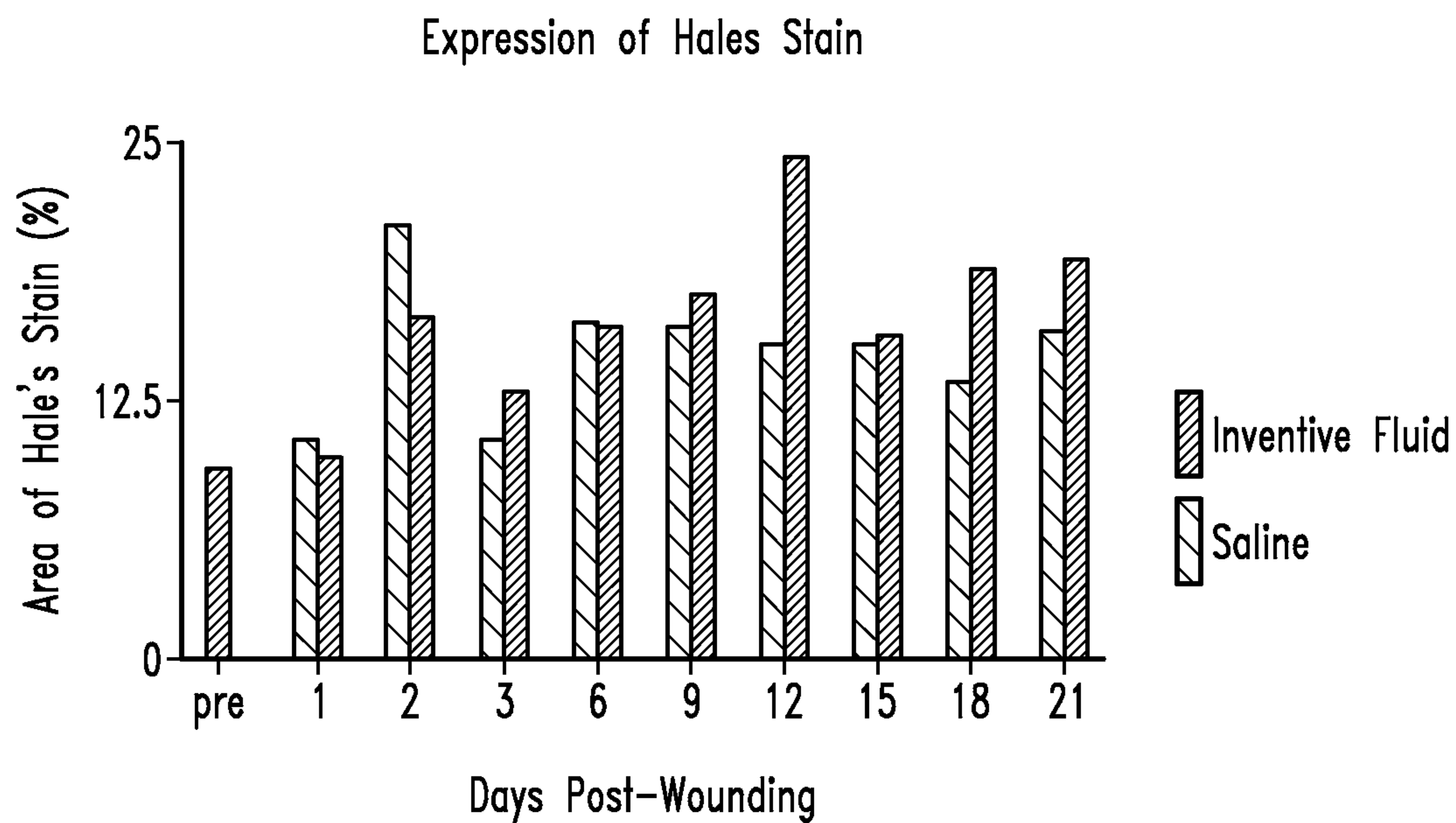
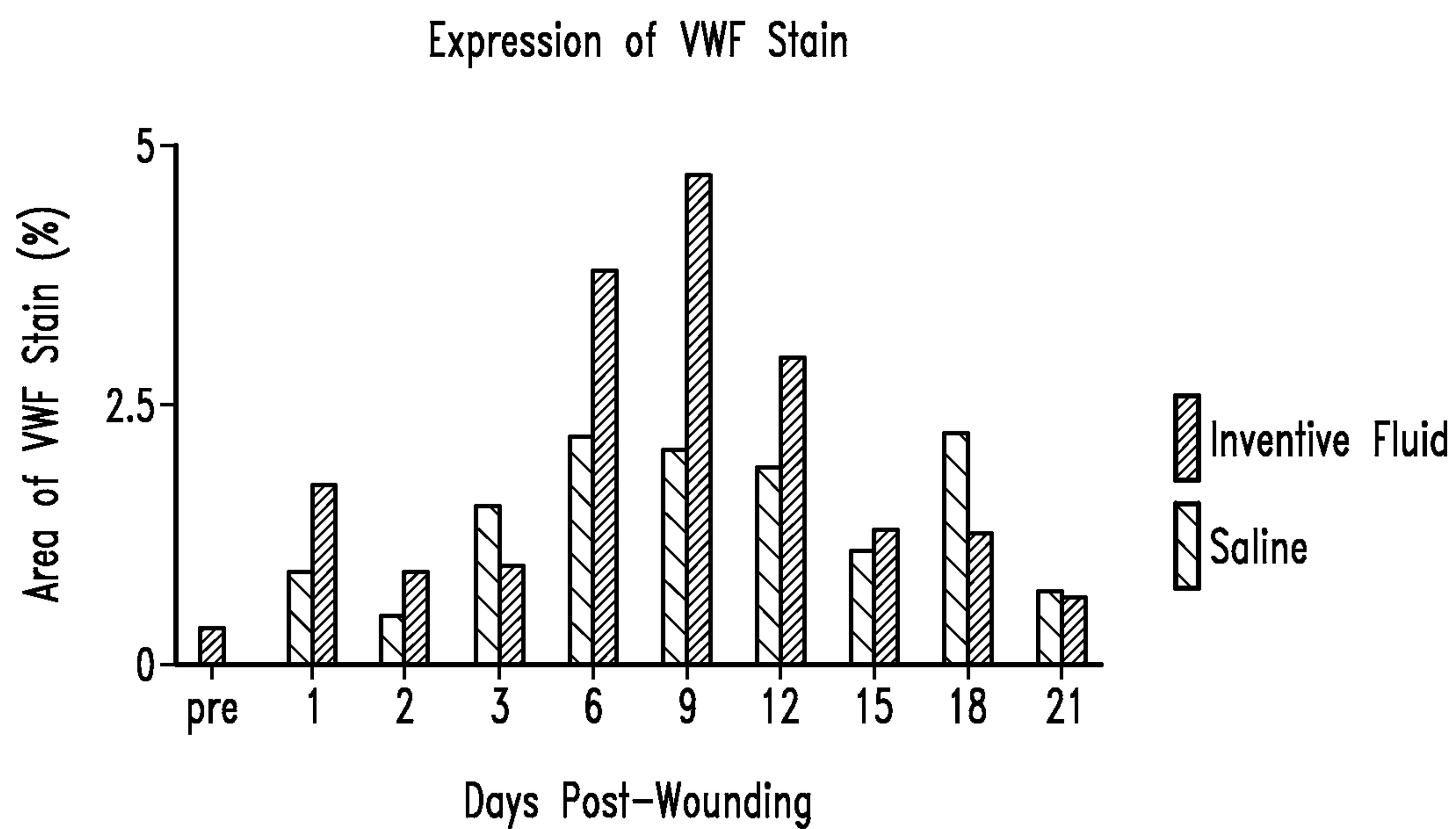
Control day 16 showing less differentiated epidermis with loss of epidermal/dermal contour

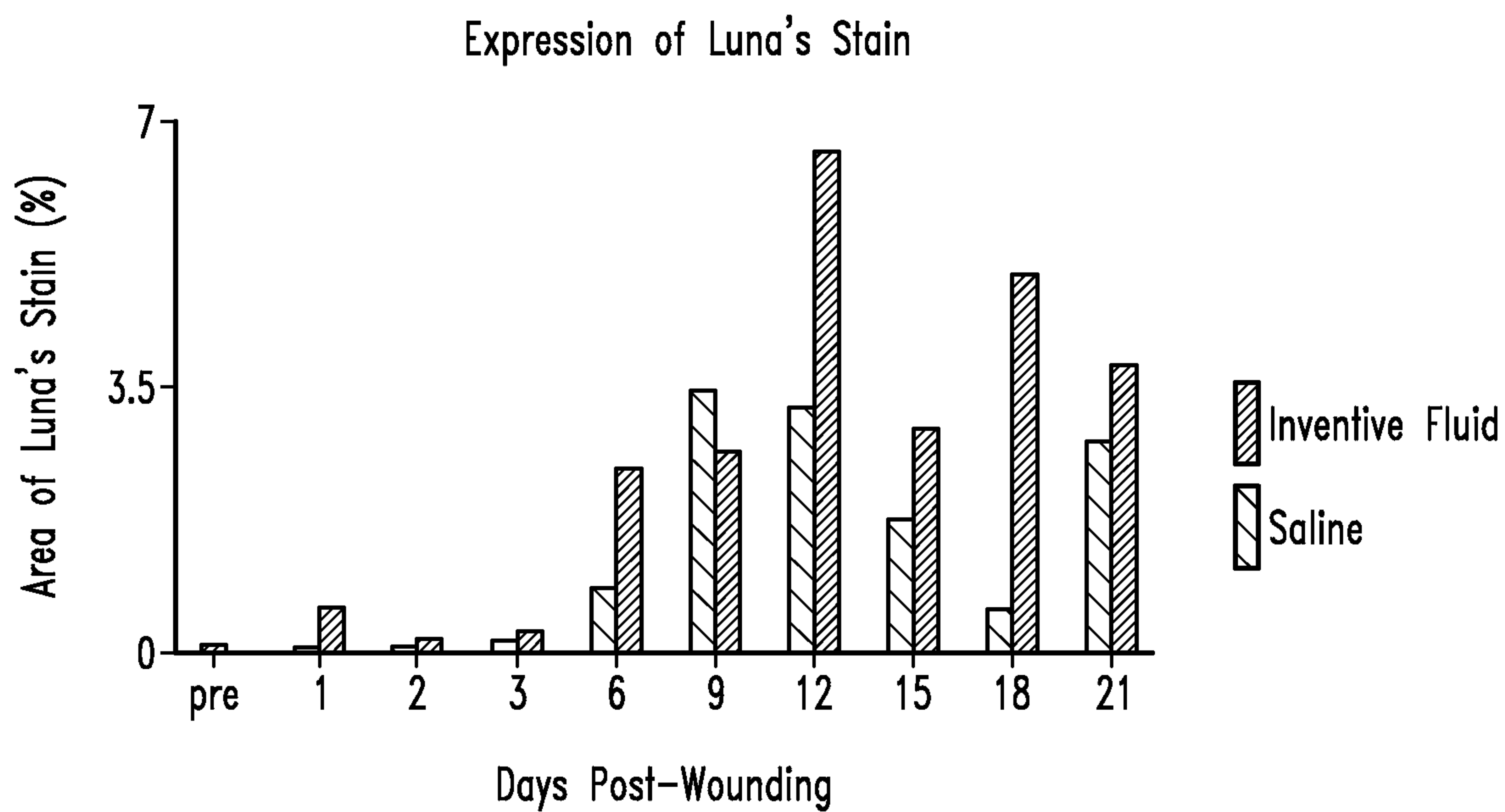
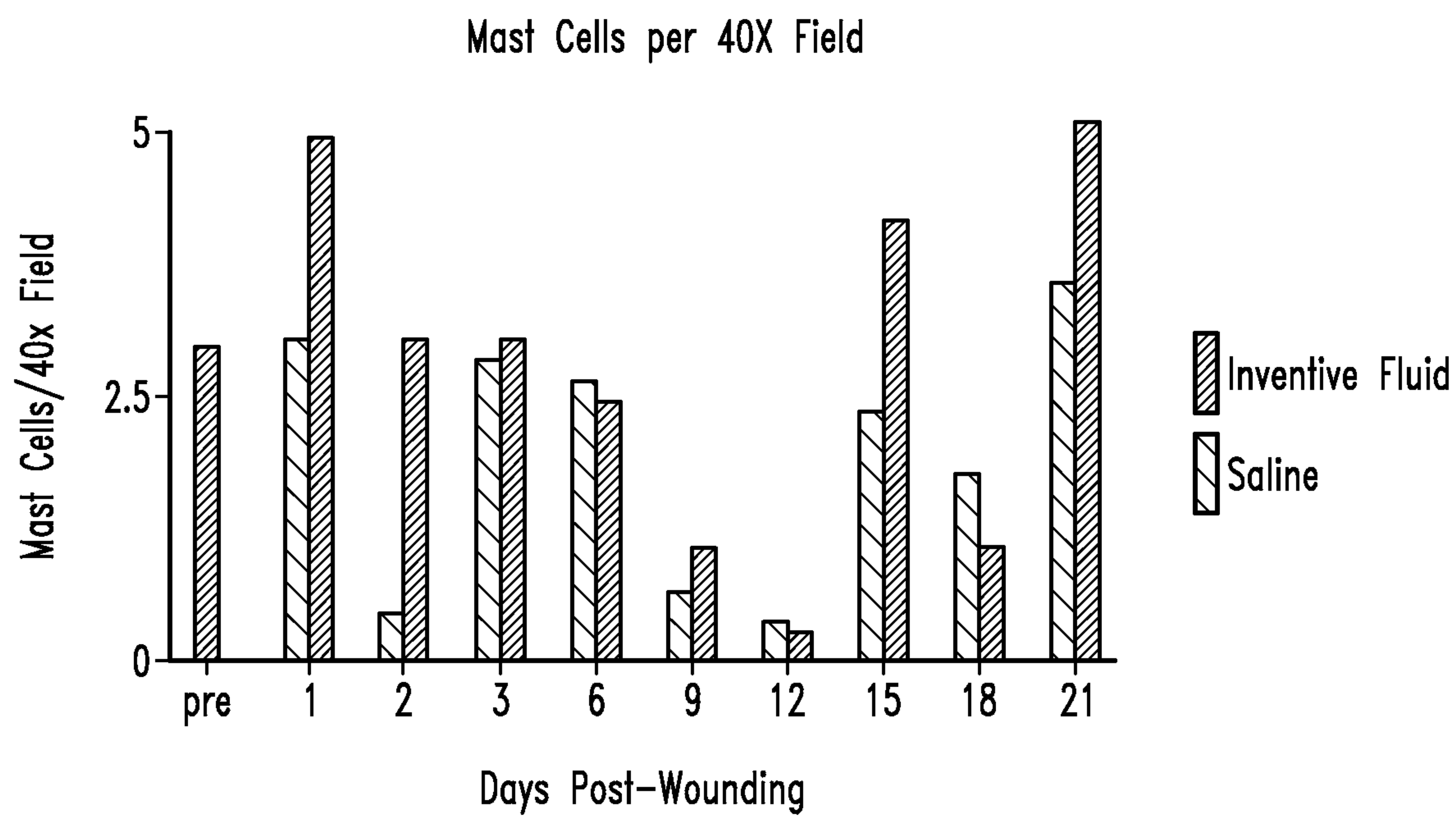
Fig. 41E



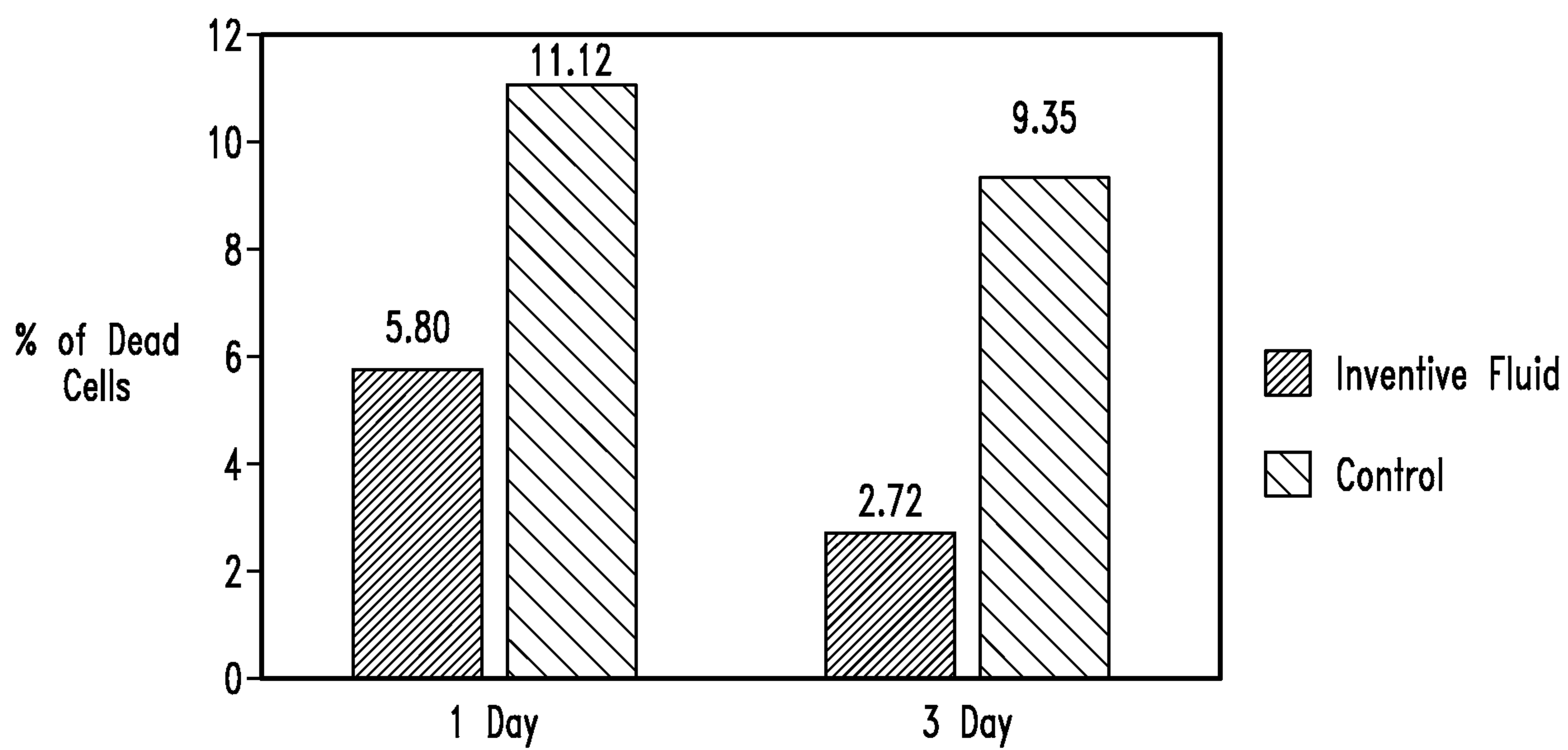
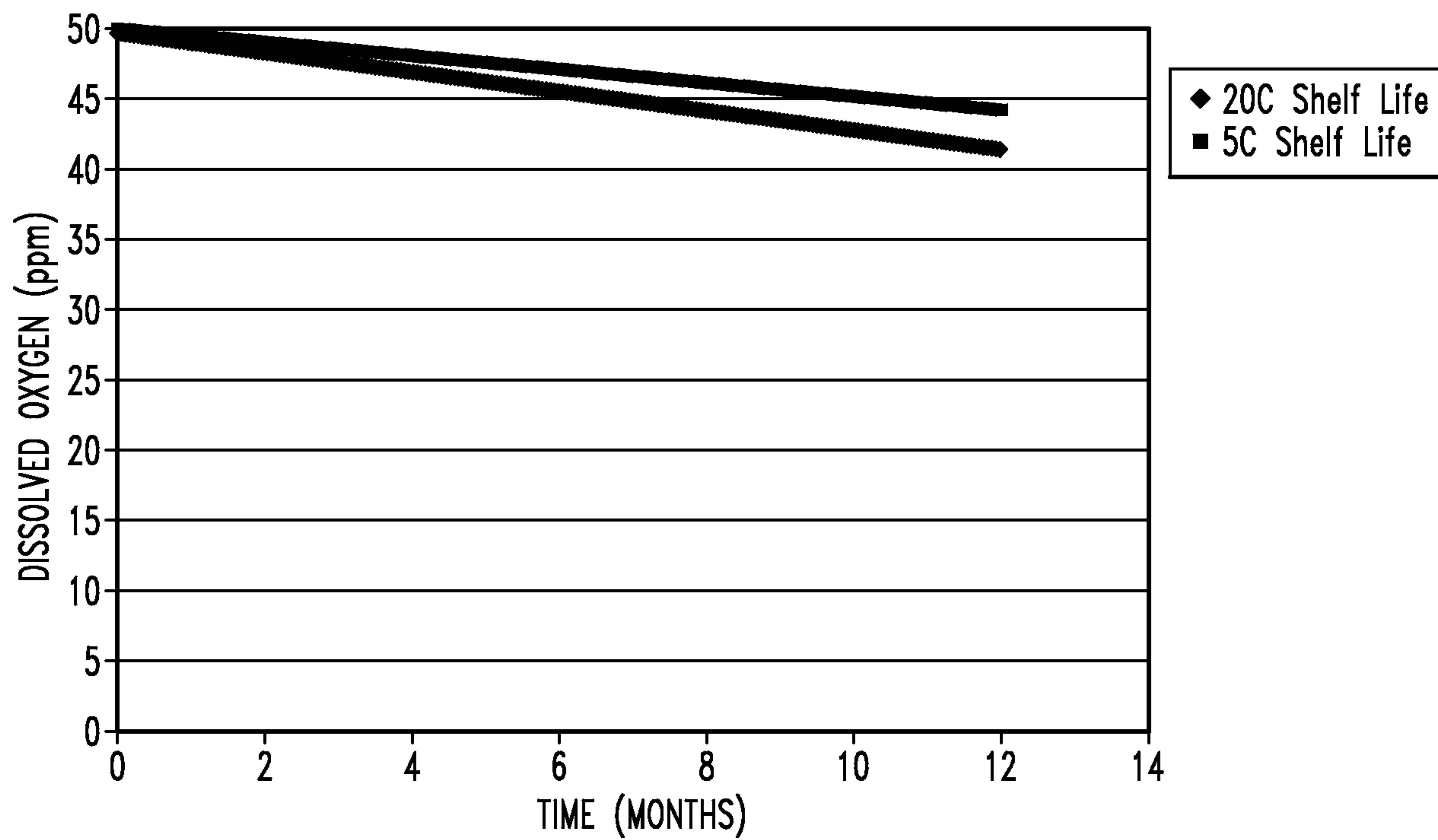
Experimental day 16 with inventive fluid showing more differentiated epidermis with more normal epidermal/dermal contour

Fig. 41F

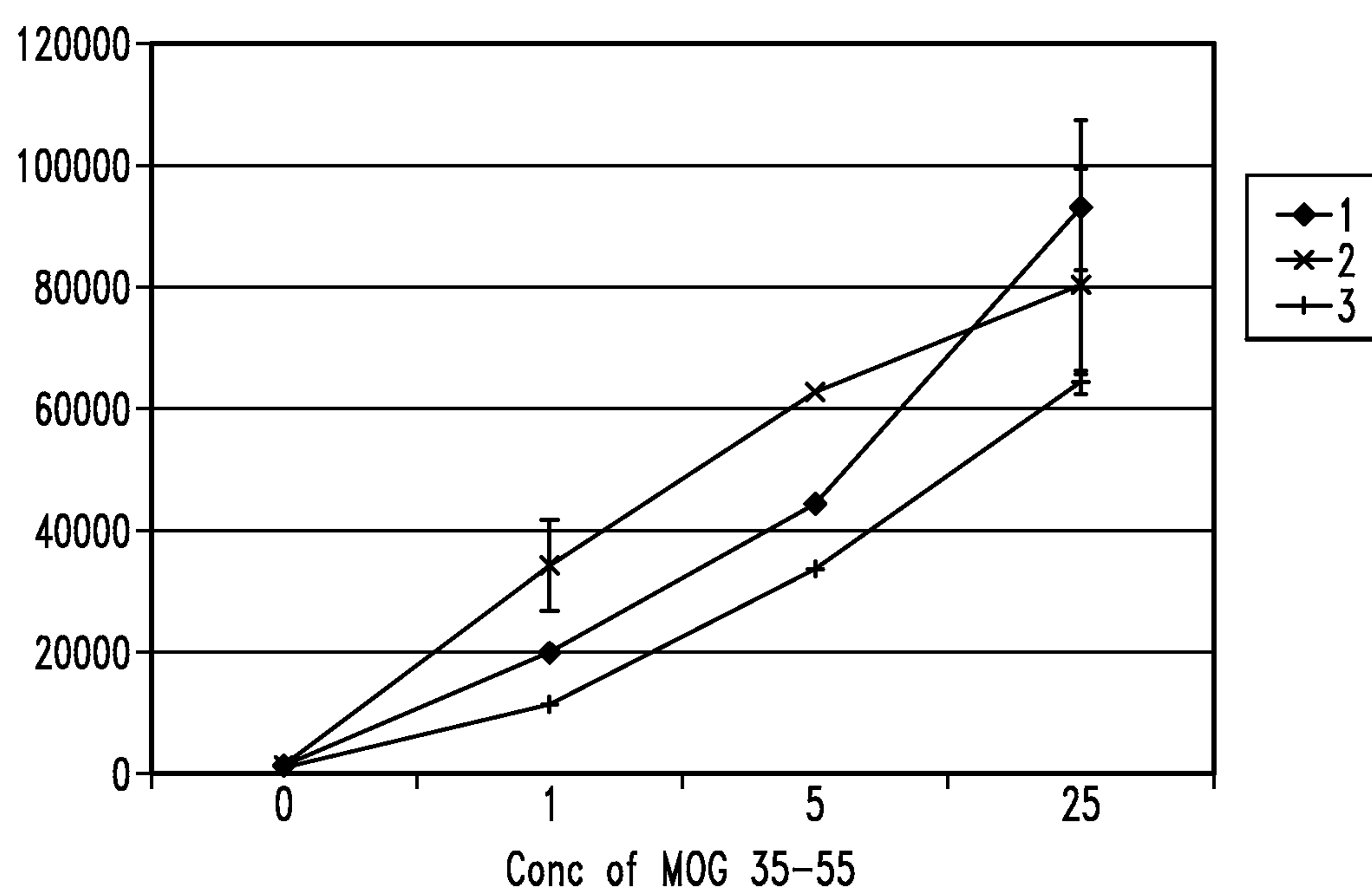
27/94**Fig. 42****Fig. 43**

28/94*Fig. 44**Fig. 45*

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*Fig. 46**Fig. 47*

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*Fig. 48*

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● RDC1676-00 OVA ⊗ RDC1676-01 OVA ○ RDC1676-00 SALINE ⊕ RDC1676-01 SALINE

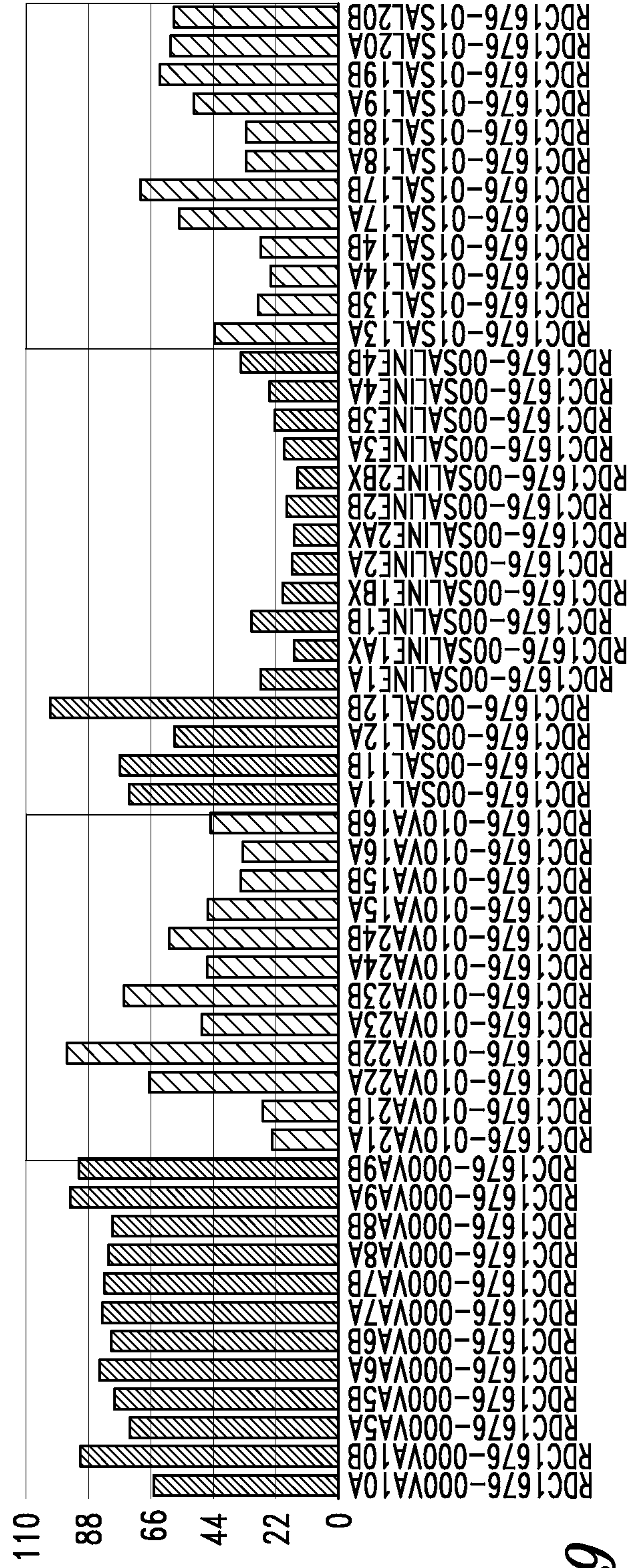
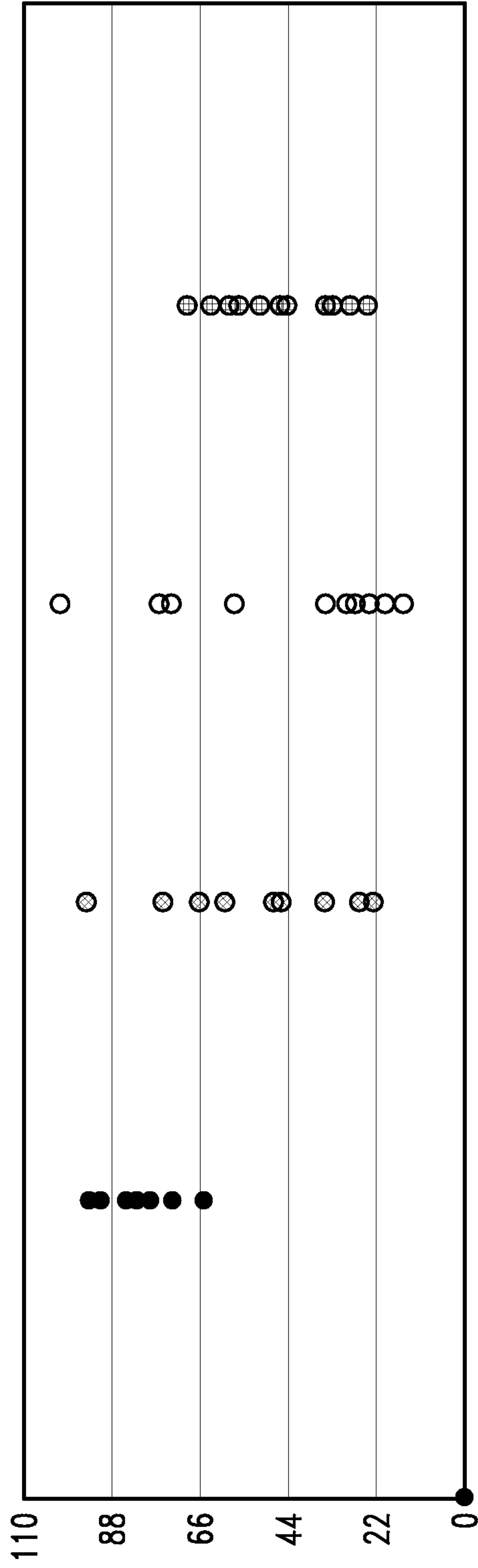


Fig. 49

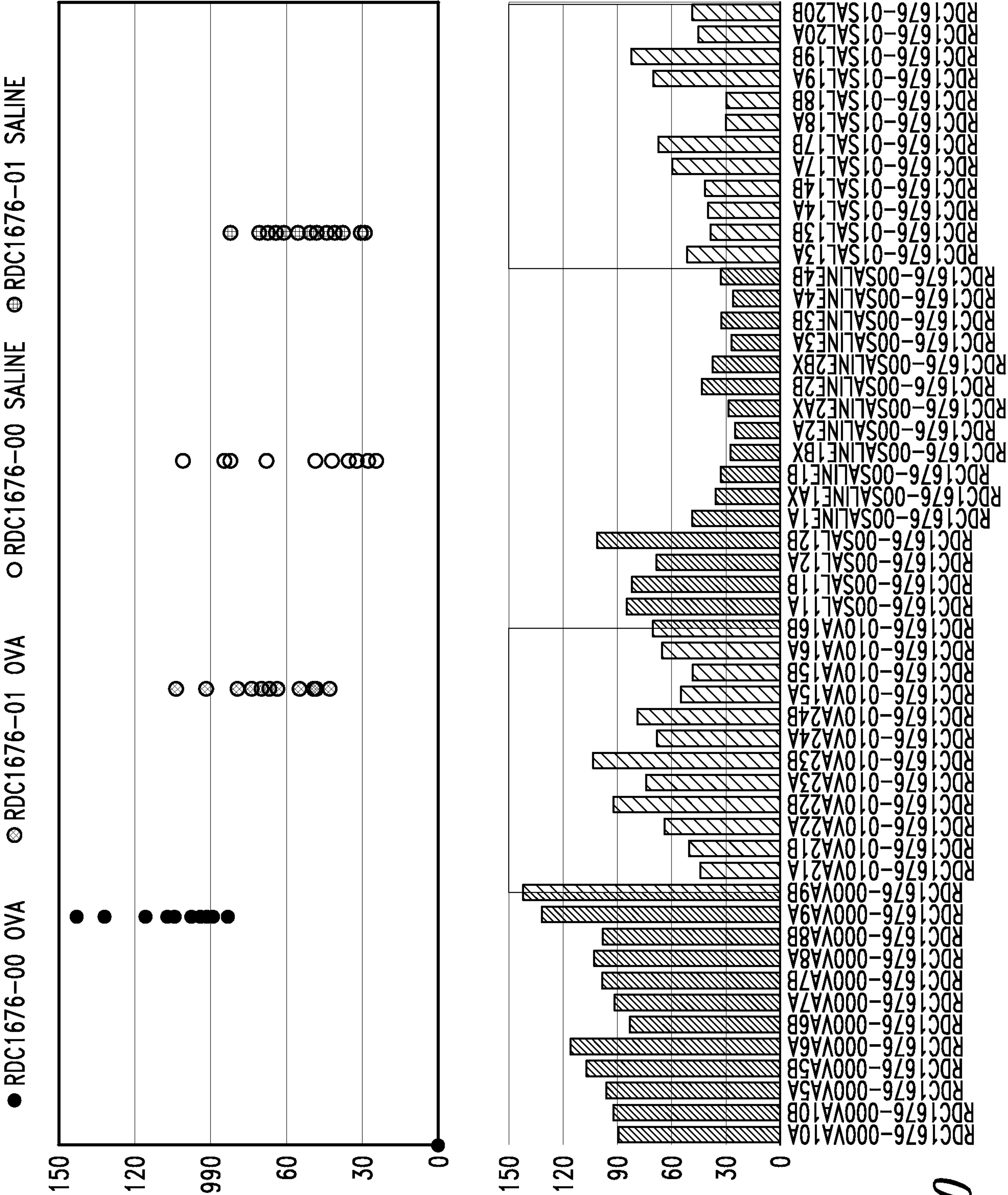


Fig. 50

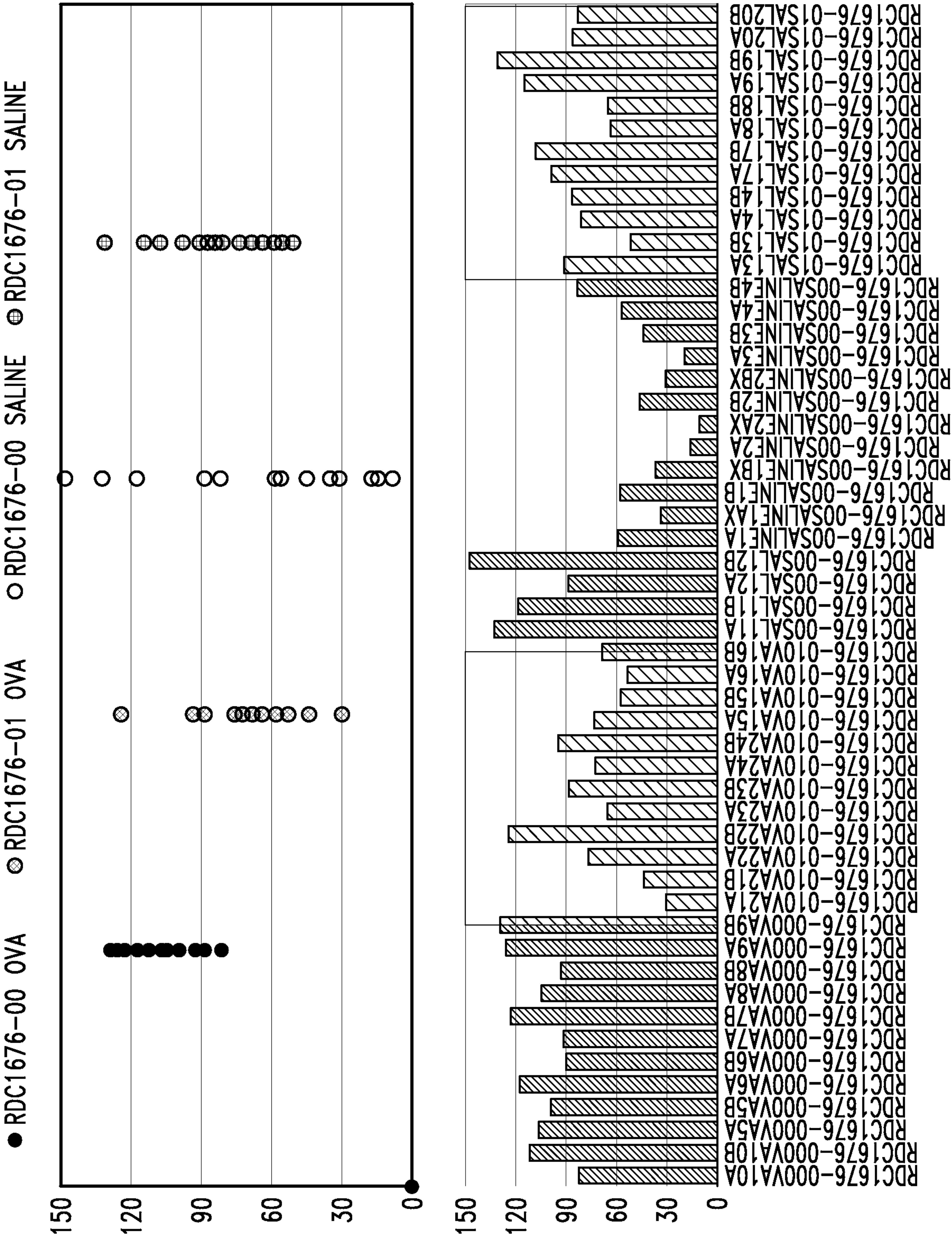


Fig. 51

Fig. 52

Fig. 53

● RDC1676-00 OVA ● RDC1676-01 OVA ○ RDC1676-00 SALINE ⊕ RDC1676-01 SALINE

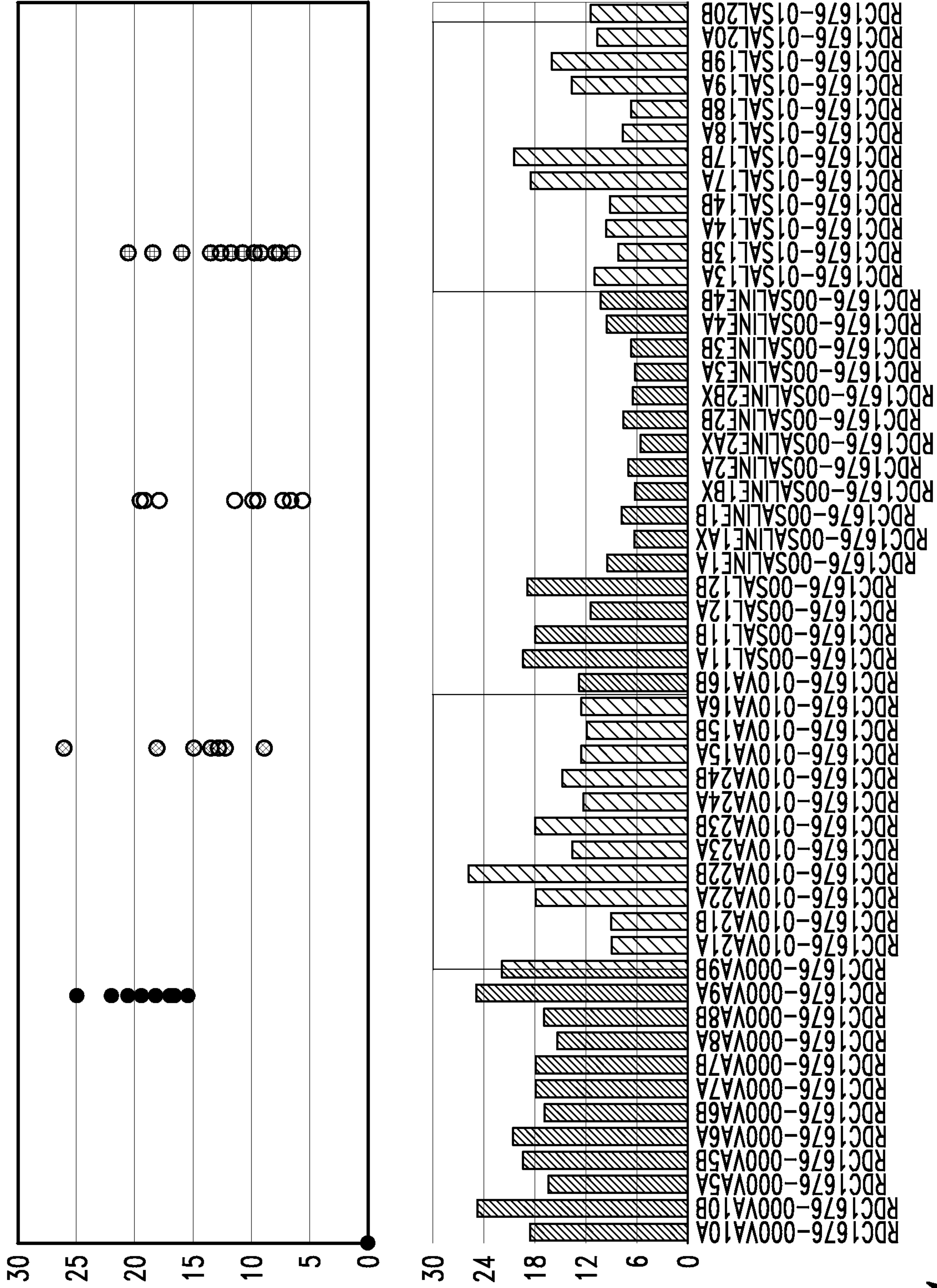


Fig. 54

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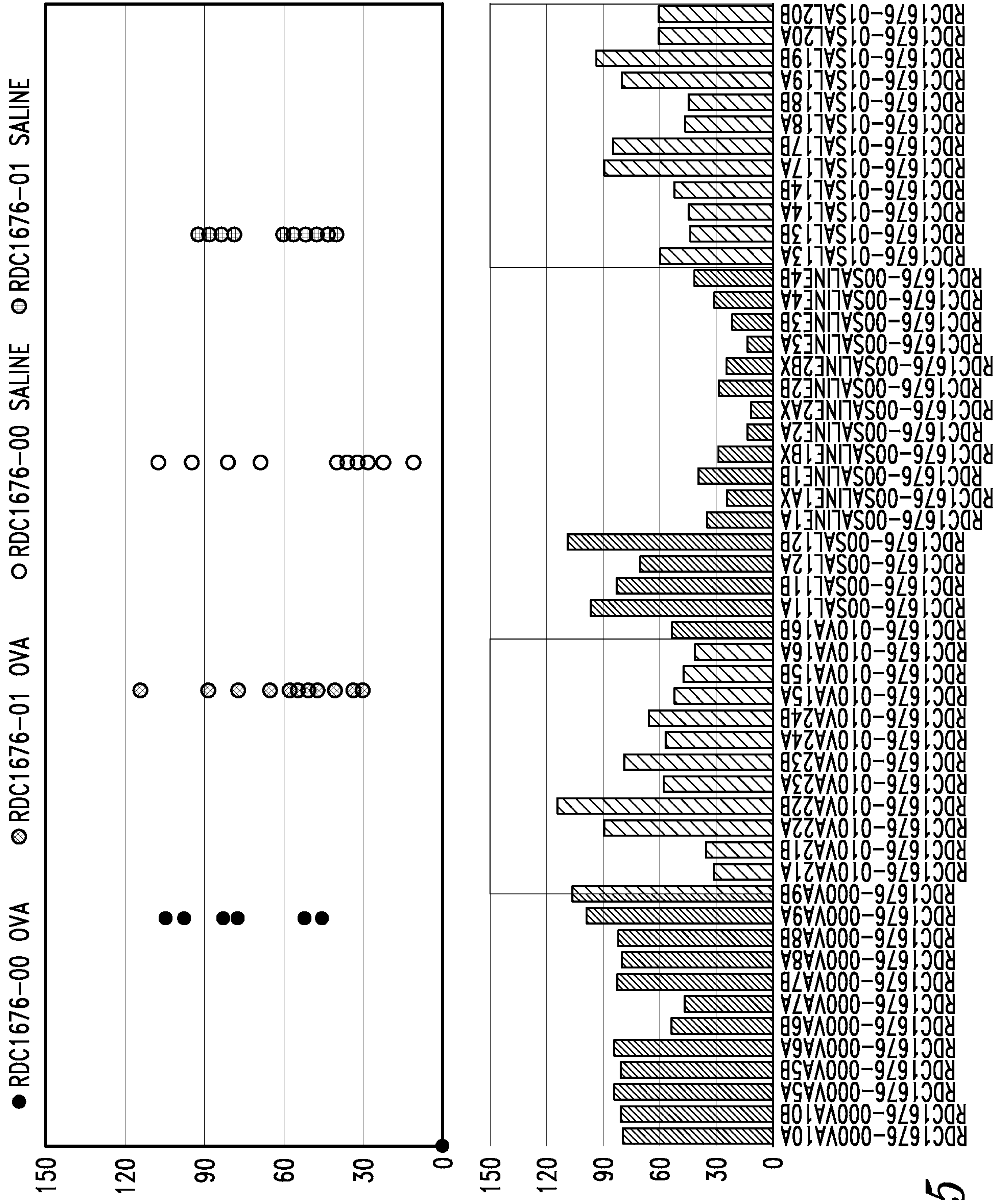


Fig. 55

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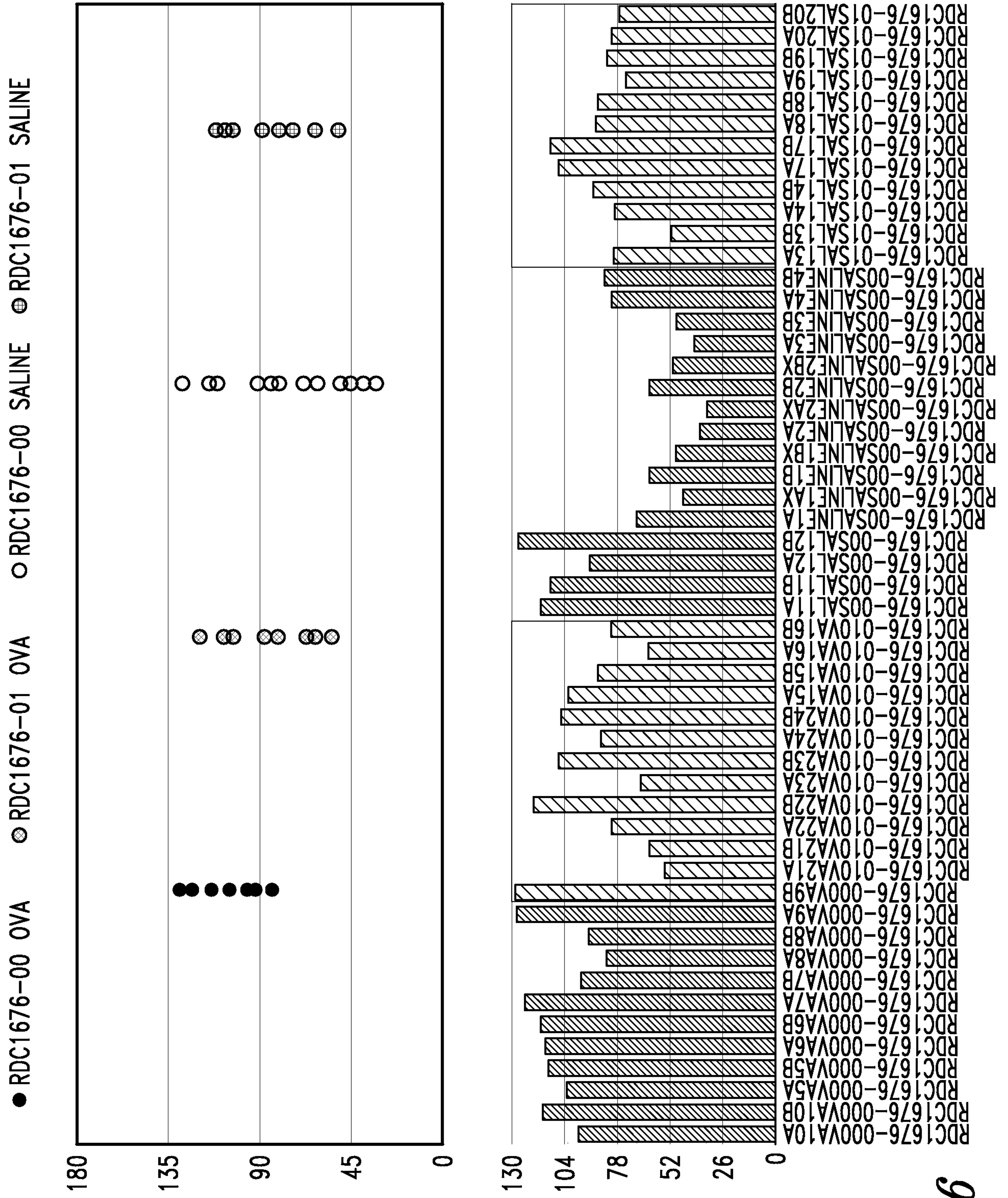


Fig. 56

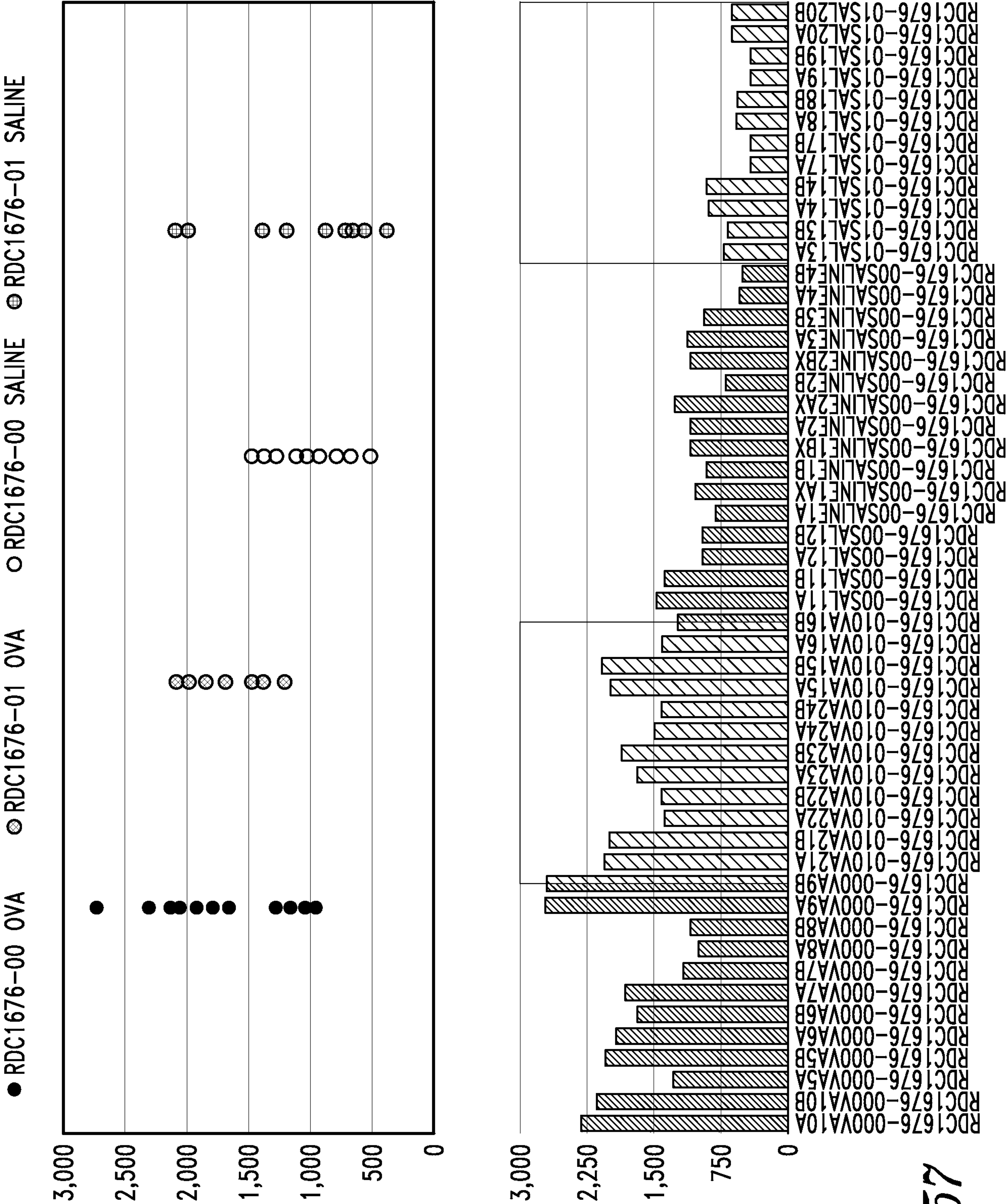


Fig. 57

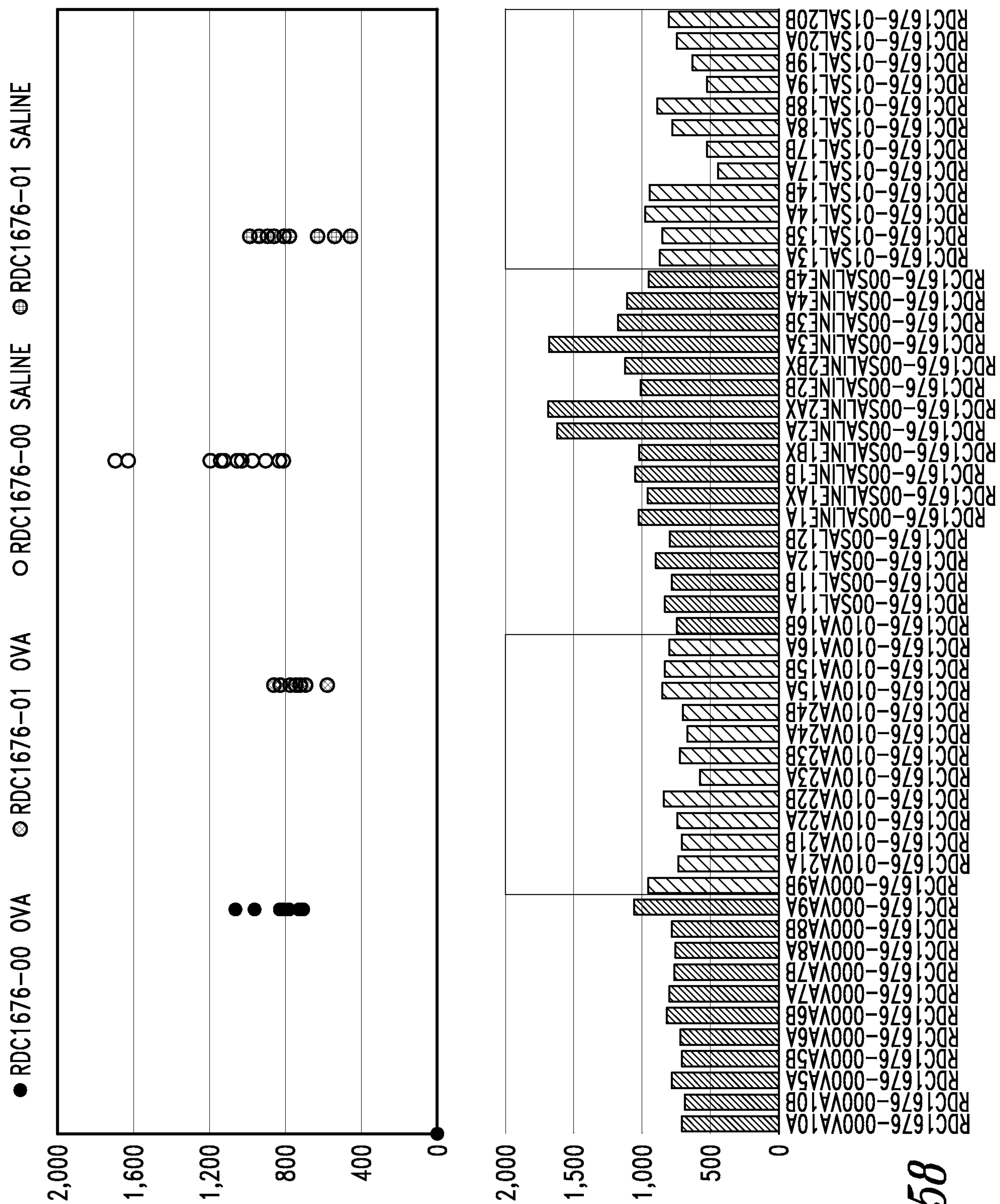


Fig. 58

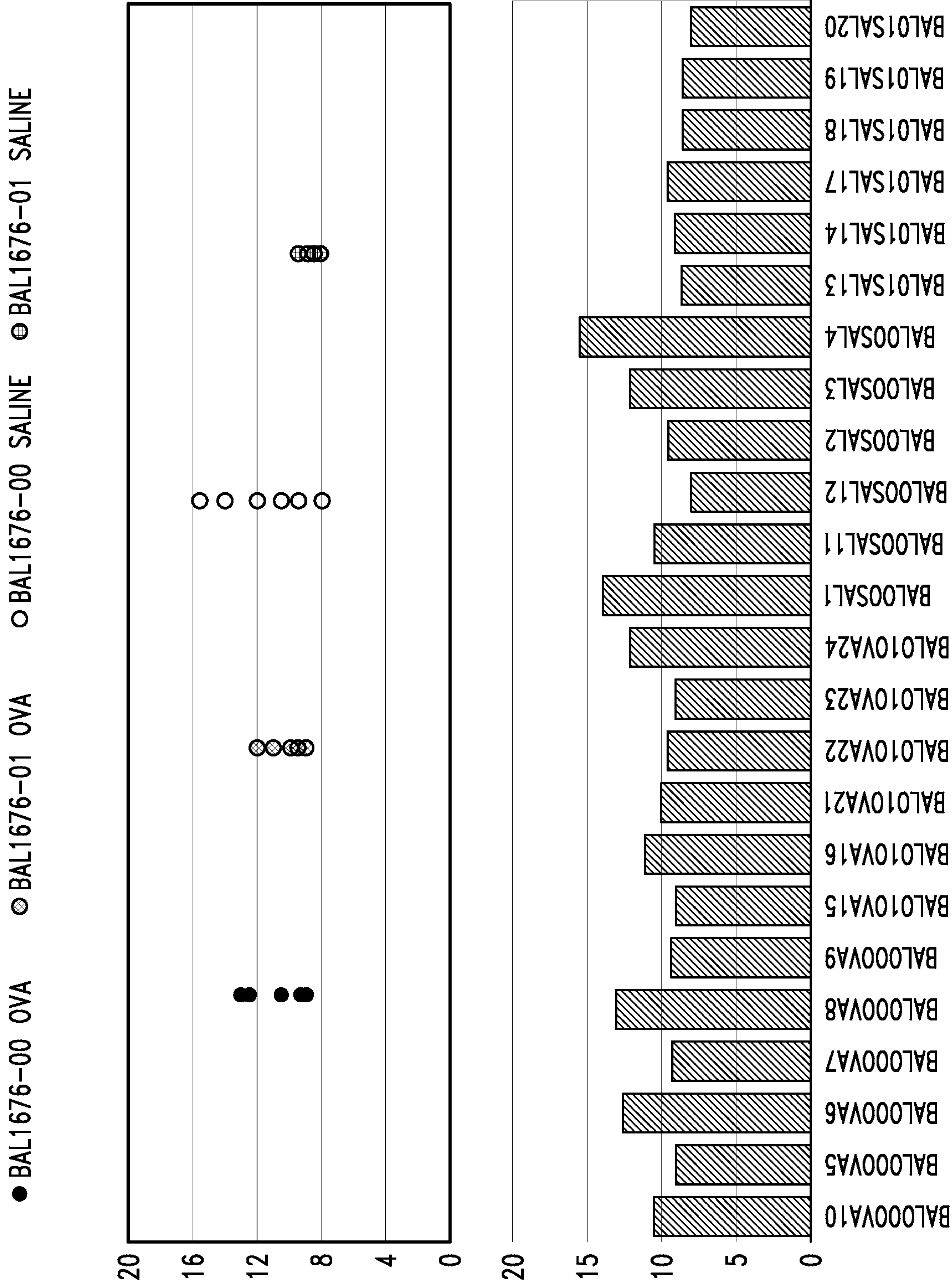


Fig. 59

● BAL1676-00 OVA ⊗ BAL1676-01 OVA ○ BAL1676-00 SALINE ⊕ BAL1676-01 SALINE

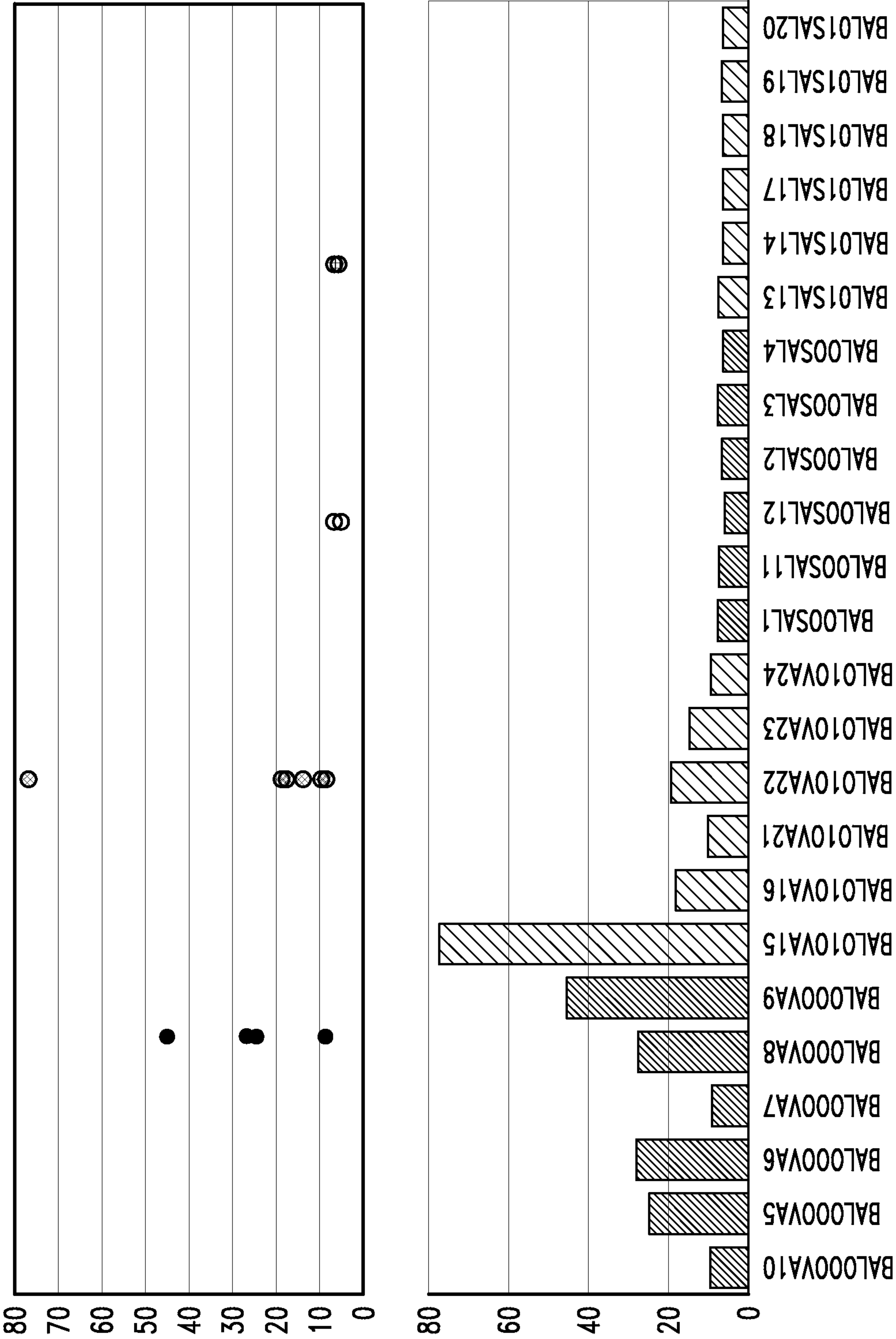


Fig. 60

● BAL1676-00 OVA ⊗ BAL1676-01 OVA ○ BAL1676-00 SALINE ⊕ BAL1676-01 SALINE

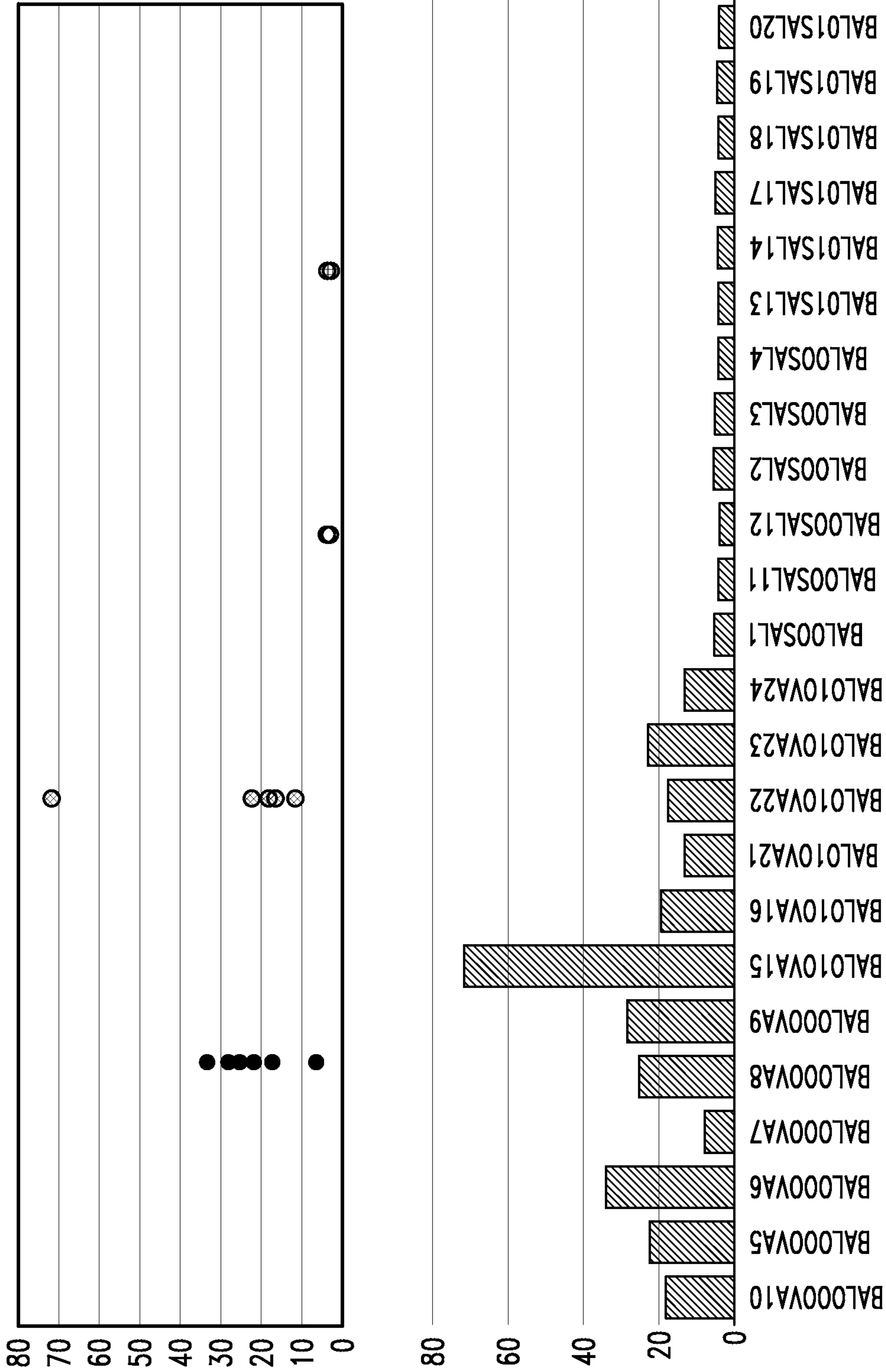


Fig. 61

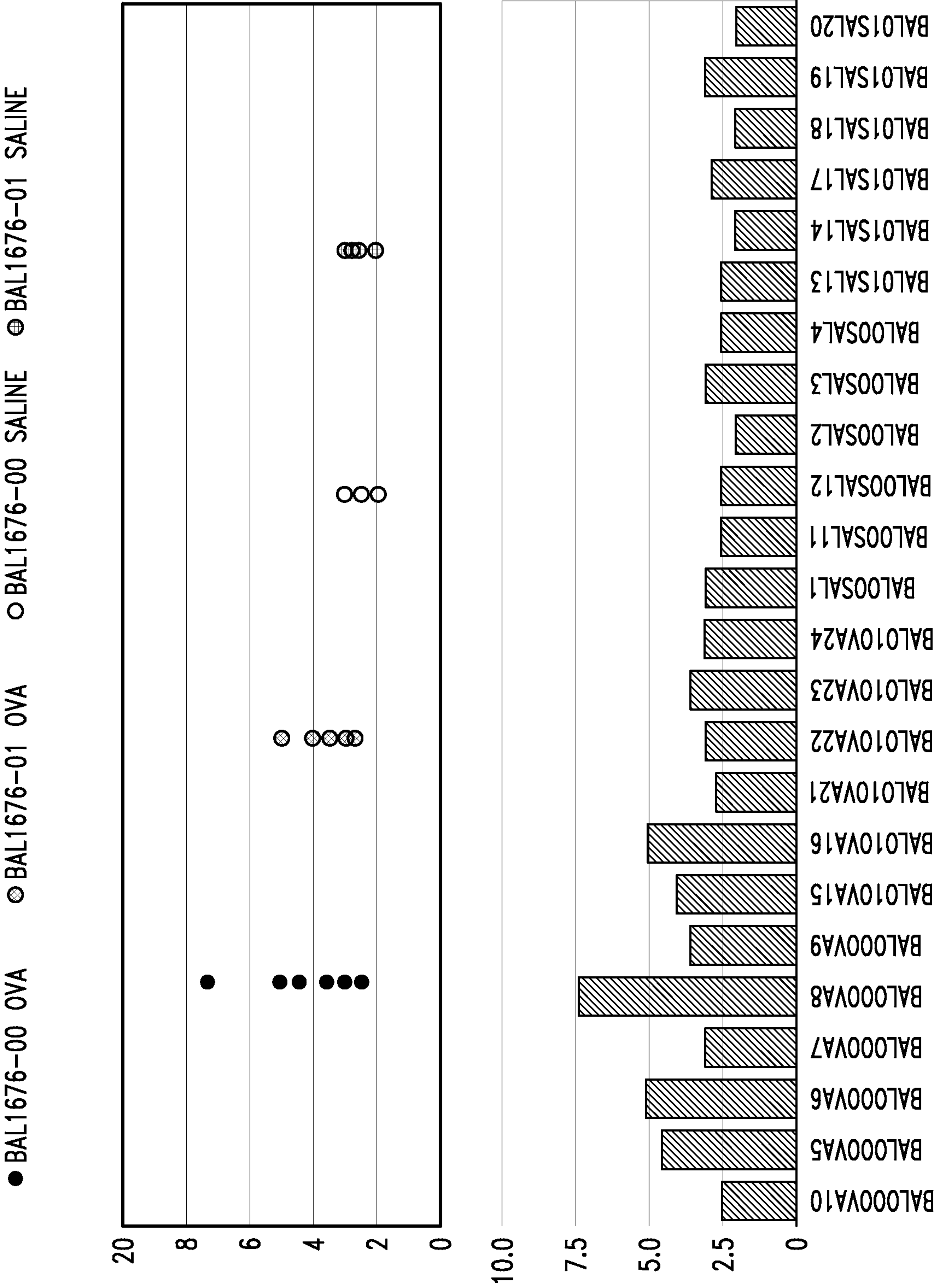


Fig. 62

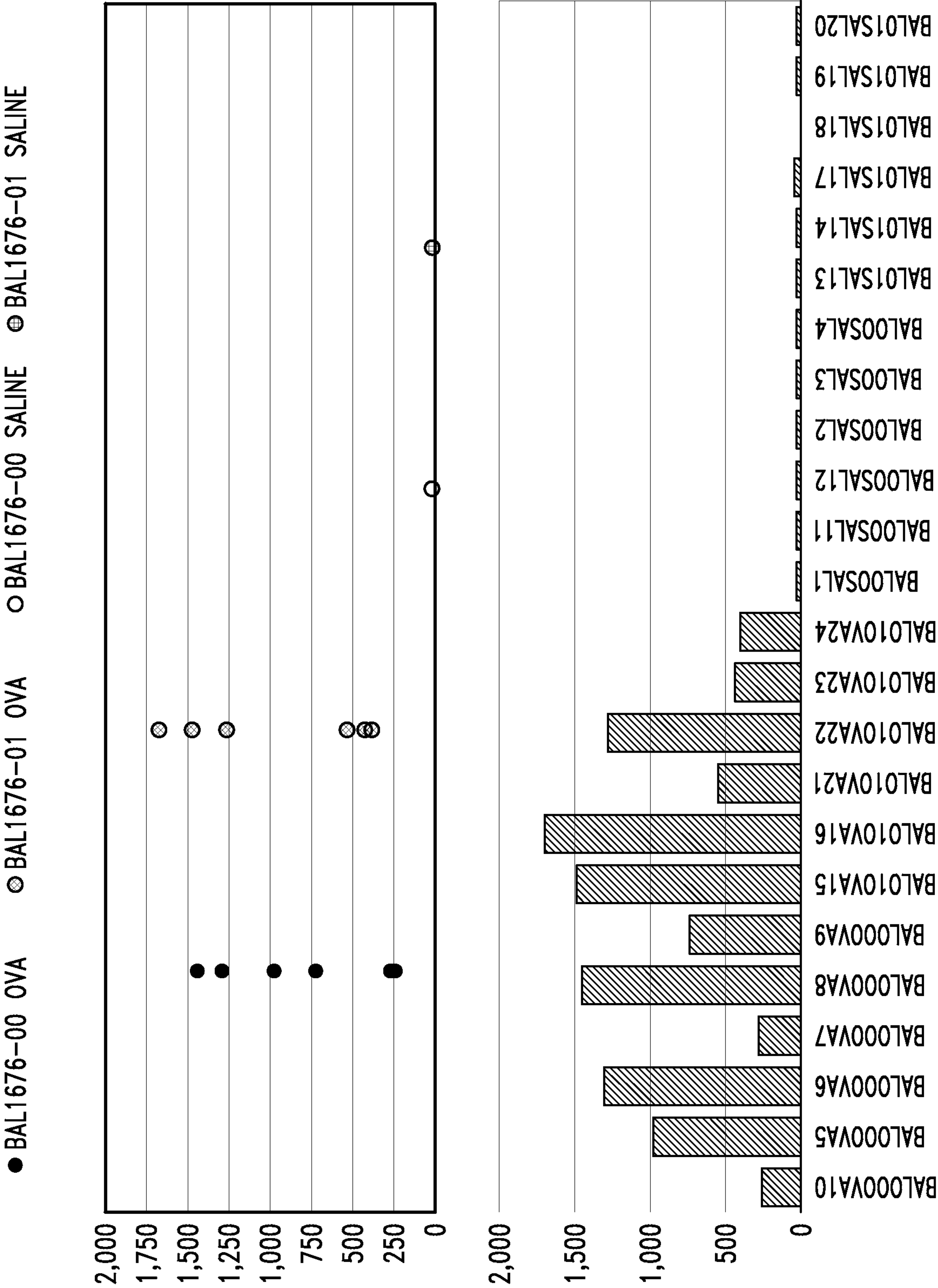


Fig. 63

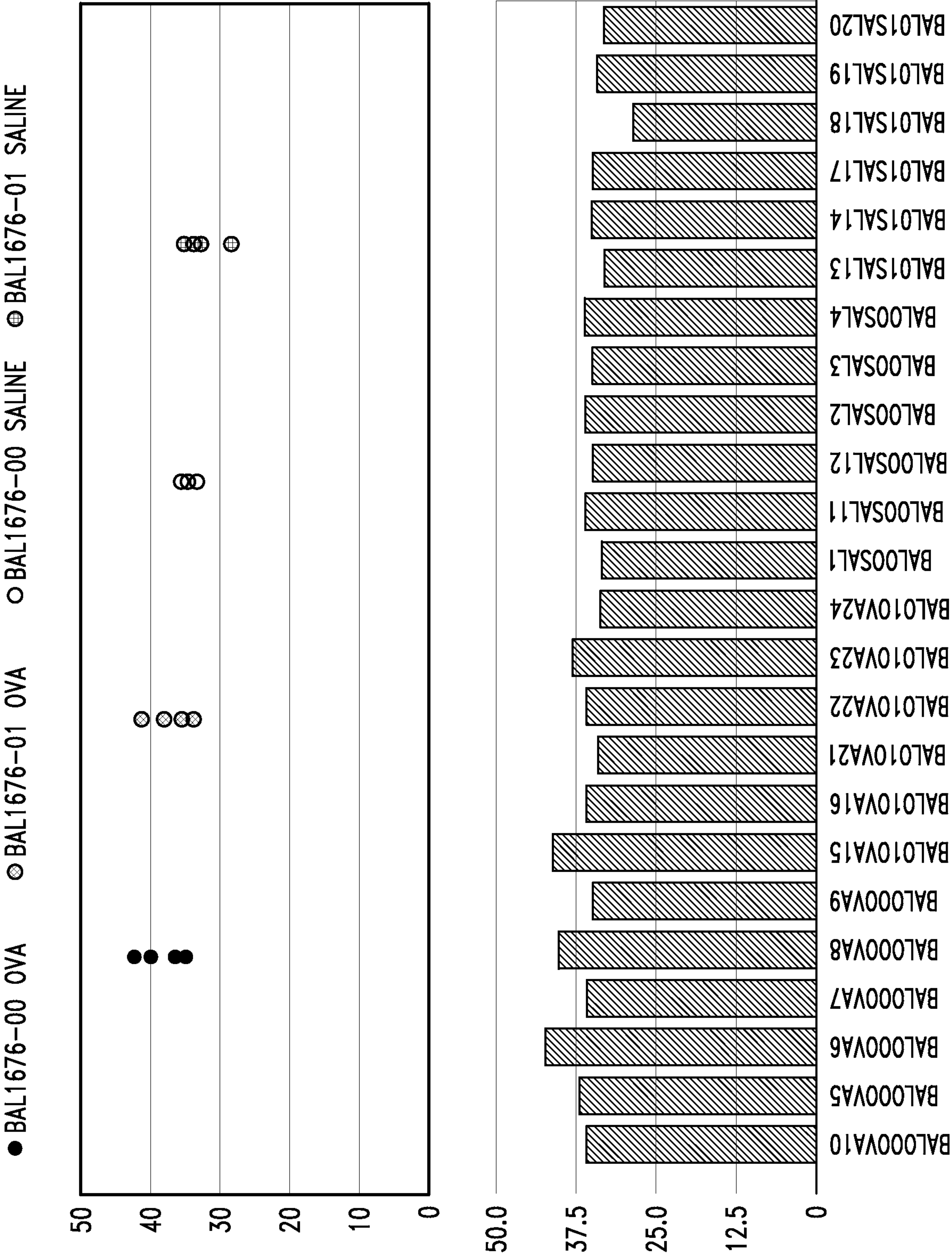


Fig. 64

● BAL1676-00 OVA ⊗ BAL1676-01 OVA ○ BAL1676-00 SALINE ⊕ BAL1676-01 SALINE

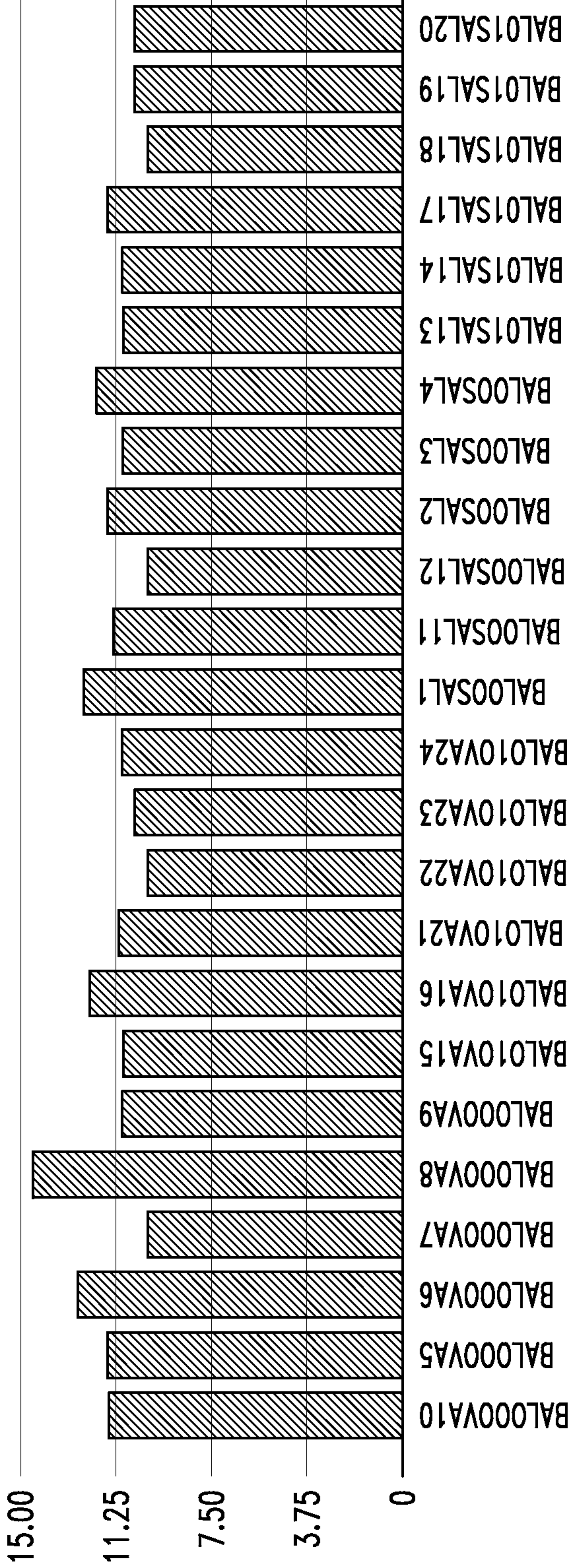
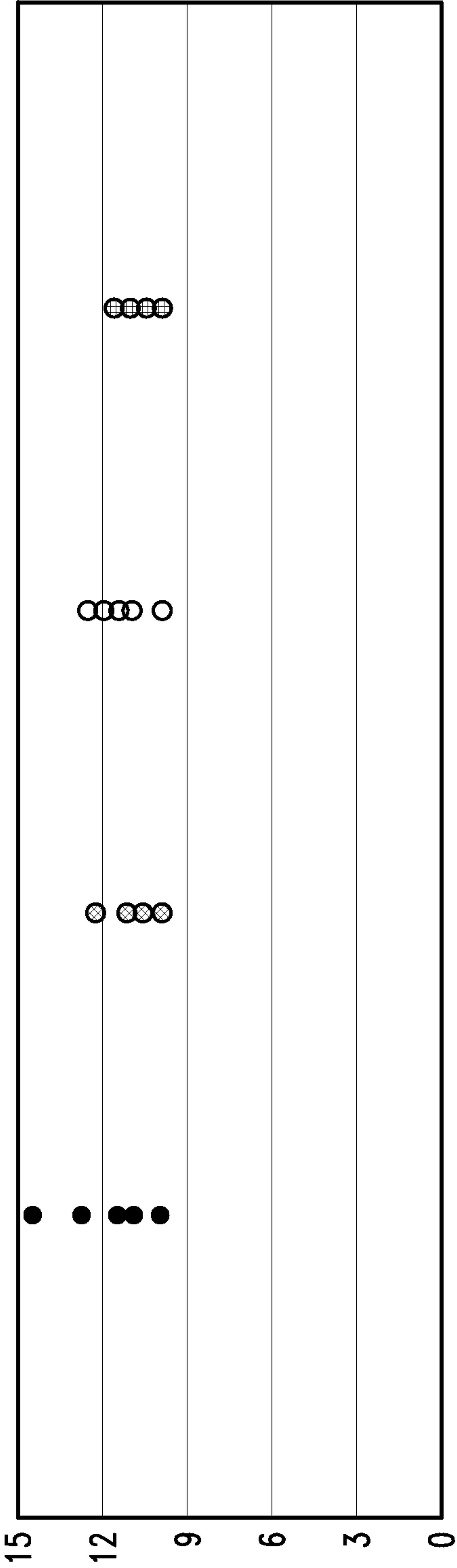


Fig. 65

● BAL1676-00 OVA ⊗ BAL1676-01 OVA ○ BAL1676-00 SALINE ⊕ BAL1676-01 SALINE

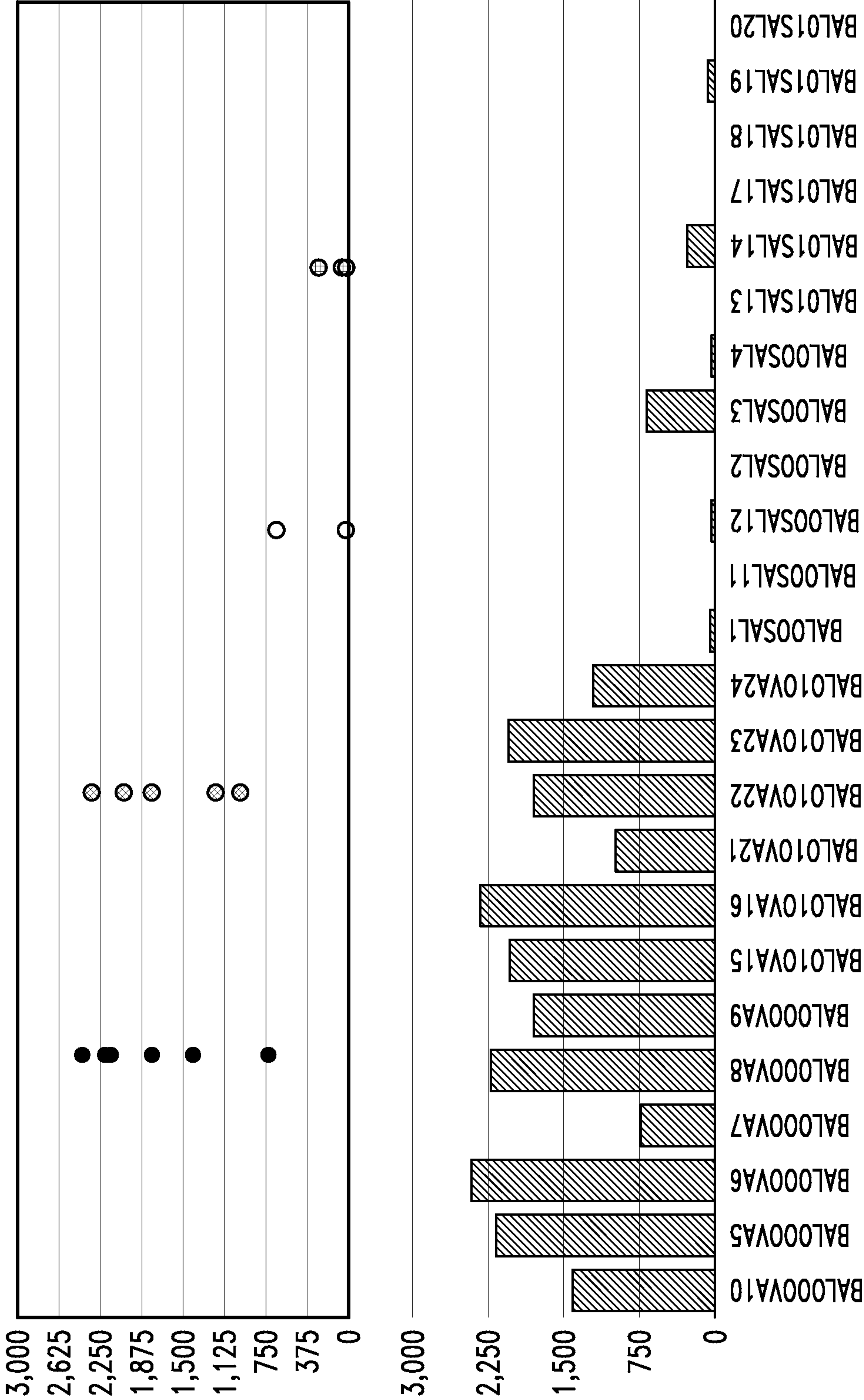


Fig. 67

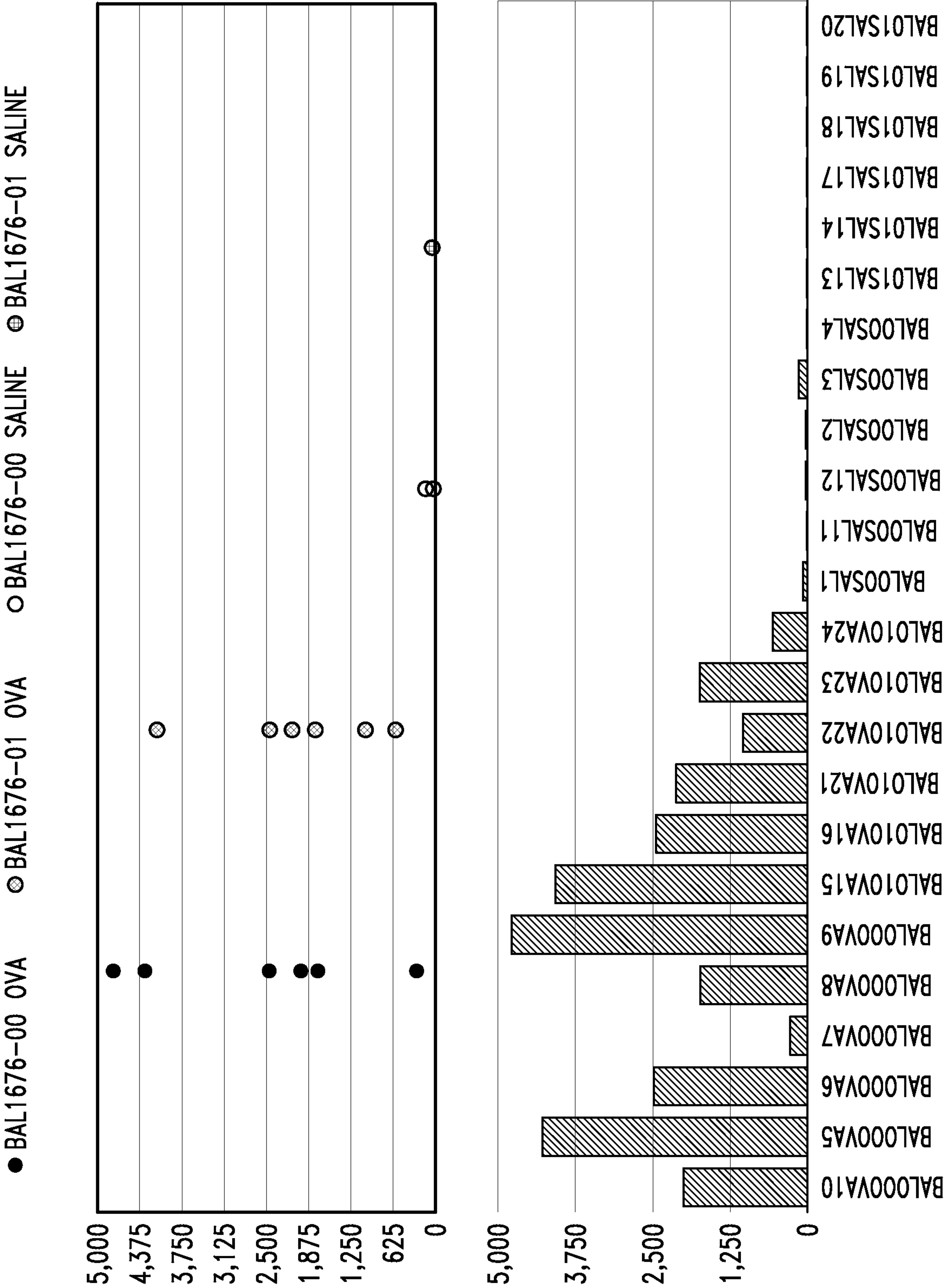


Fig. 68

Step Data Setup-

Data Name	Assay Time	Flow Rate	Type
Def Assoc.	900	1000	Assoc.
Def Dissoc.	900	1000	Dissoc.
Def Baseline No Fl...	600	0	Baseline
Def Loading No Flow	900	0	Loading
Def Activation	600	1000	Activation
Def Quenching	600	1000	Quenching
Def Regeneration	30	1000	Regeneration
baseline1	30	1000	Baseline
loading	400	1000	Loading
baseline2	60	1000	Baseline
bradykinin	600	1000	Assoc.
dissociation	900	1000	Dissoc.

☐ + ☐ Add	☐ - ☐ Remove	Time in (sec), flow rate in (rpm)
---	--	-----------------------------------

Assay Step Lists

[illegible]

	Sample plate set-up
	Column 1: saline
	Column 2: receptor B2 (50ug/mL) in saline
	Column 3: A-D revsaline; E-H saline
	Column 4: A-D Bradykinin at 1.25mM, 125uM 12.5uM in revsaline
	E-H Bradykinin at 1.25mM, 125uM, 12.5uM in saline
	 Move

Fig. 69

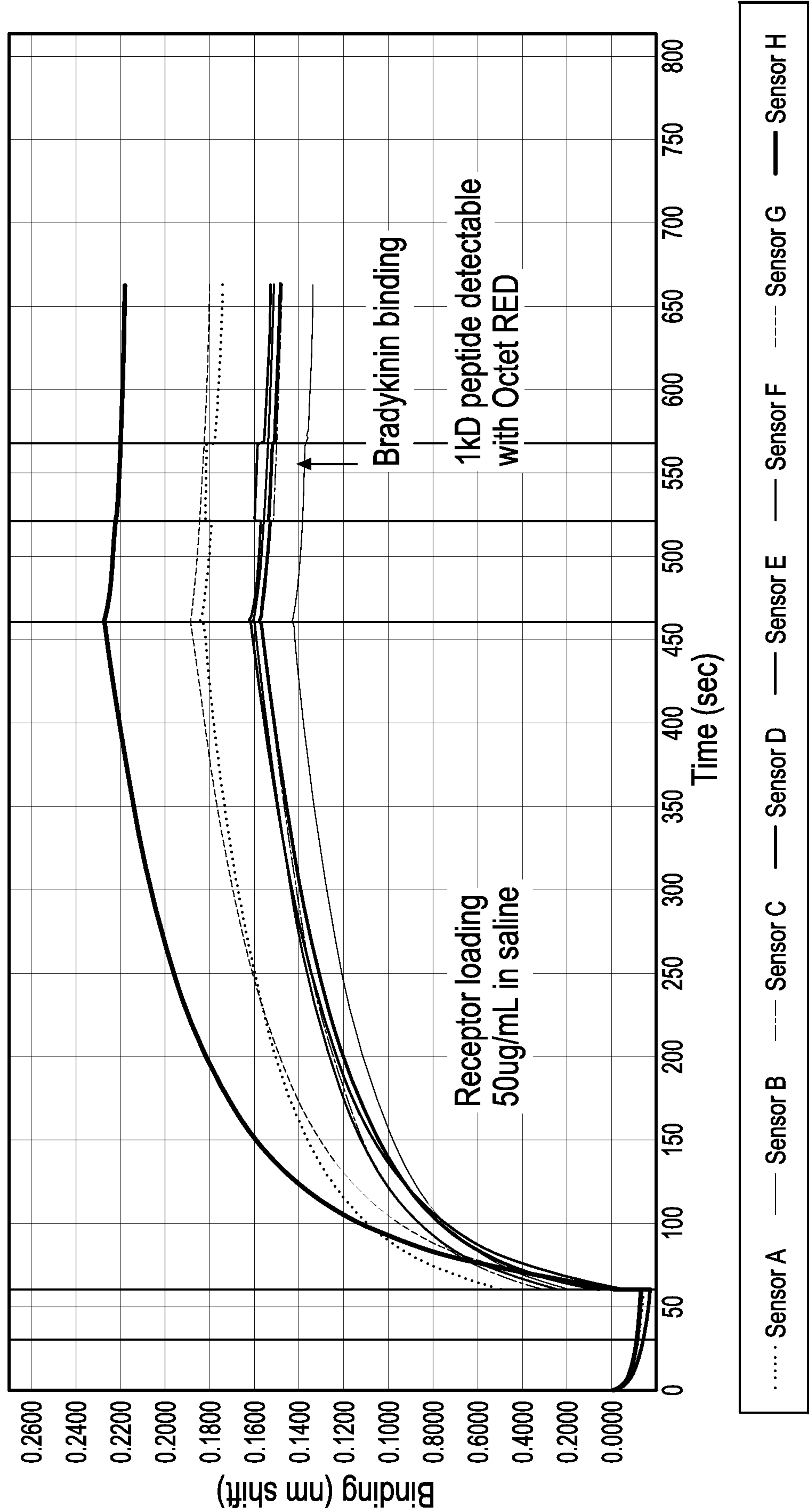


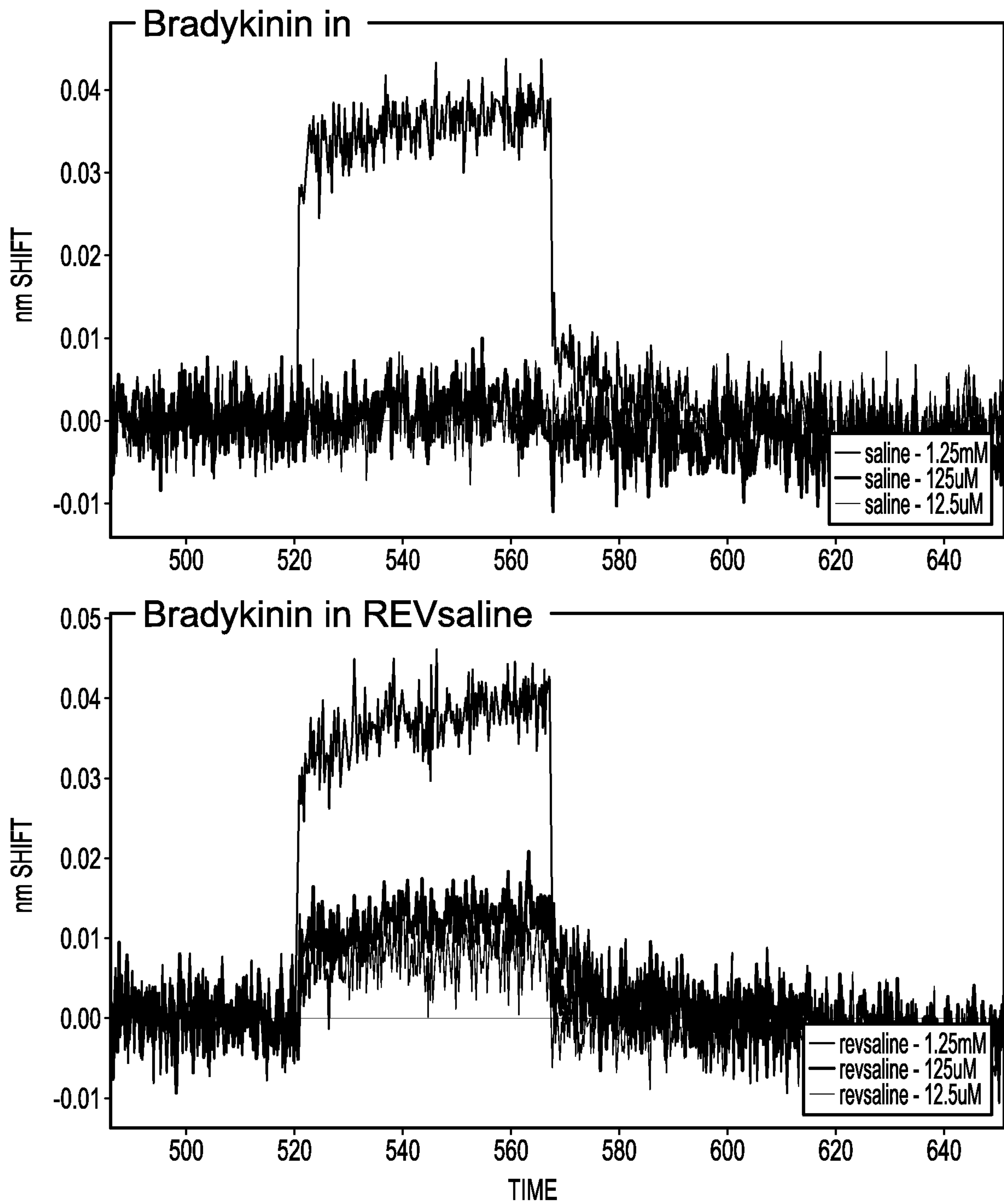
Fig. 70

Figure 1 displays two plates used for the experiment: a Sensor Plate and a Sample Plate. The Sensor Plate is an 8x12 grid of squares. The first column (A) contains 'X' marks, while the other columns (B-H) are empty. The Sample Plate is an 8x12 grid of circles. The first column (A) contains circles with diagonal lines. The other columns (B-H) contain circles with horizontal lines, vertical lines, cross-hatch patterns, and diagonal lines.

[illegible]

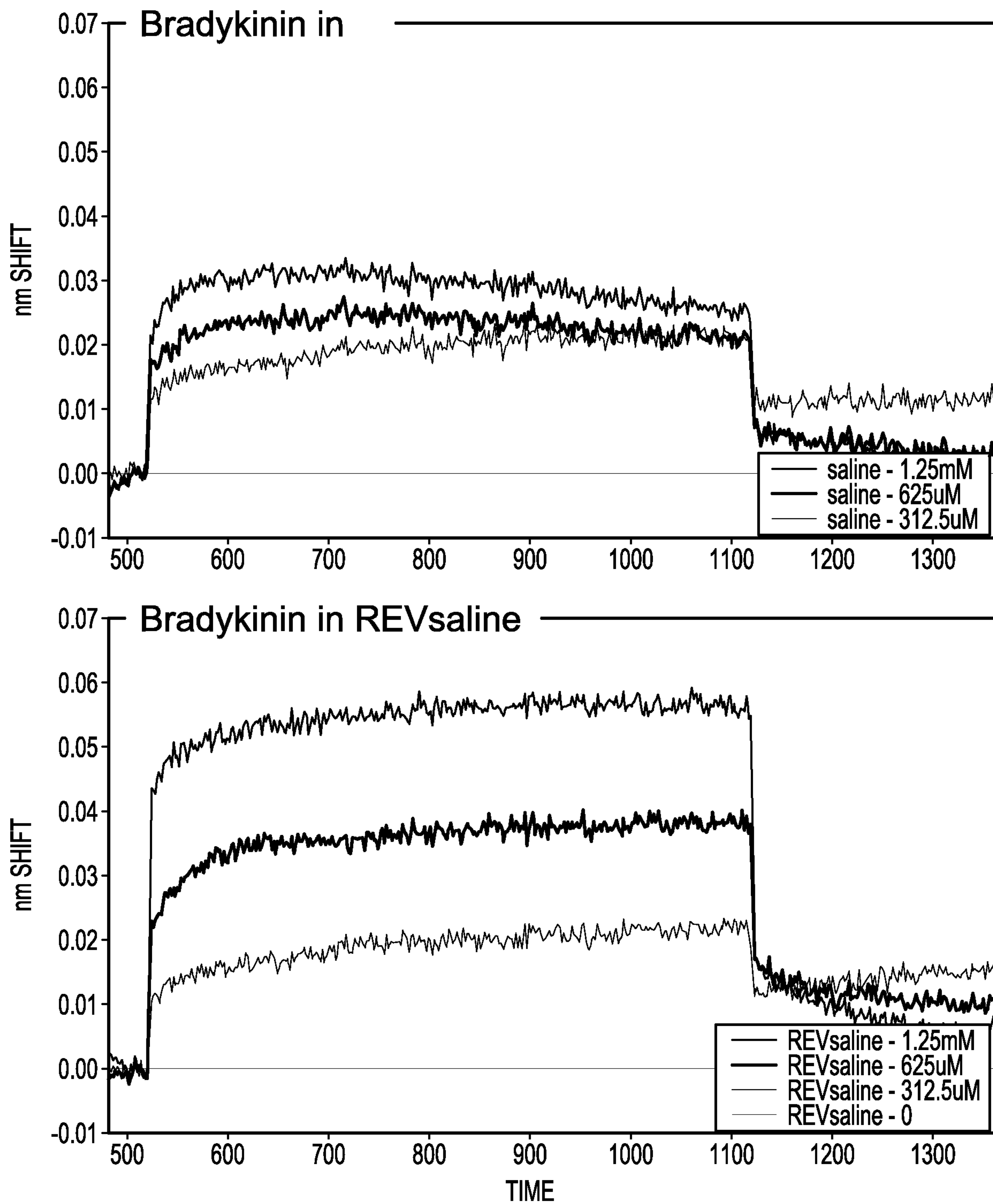
Fig. 71

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*Fig. 72*

[illegible]

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*Fig. 74*

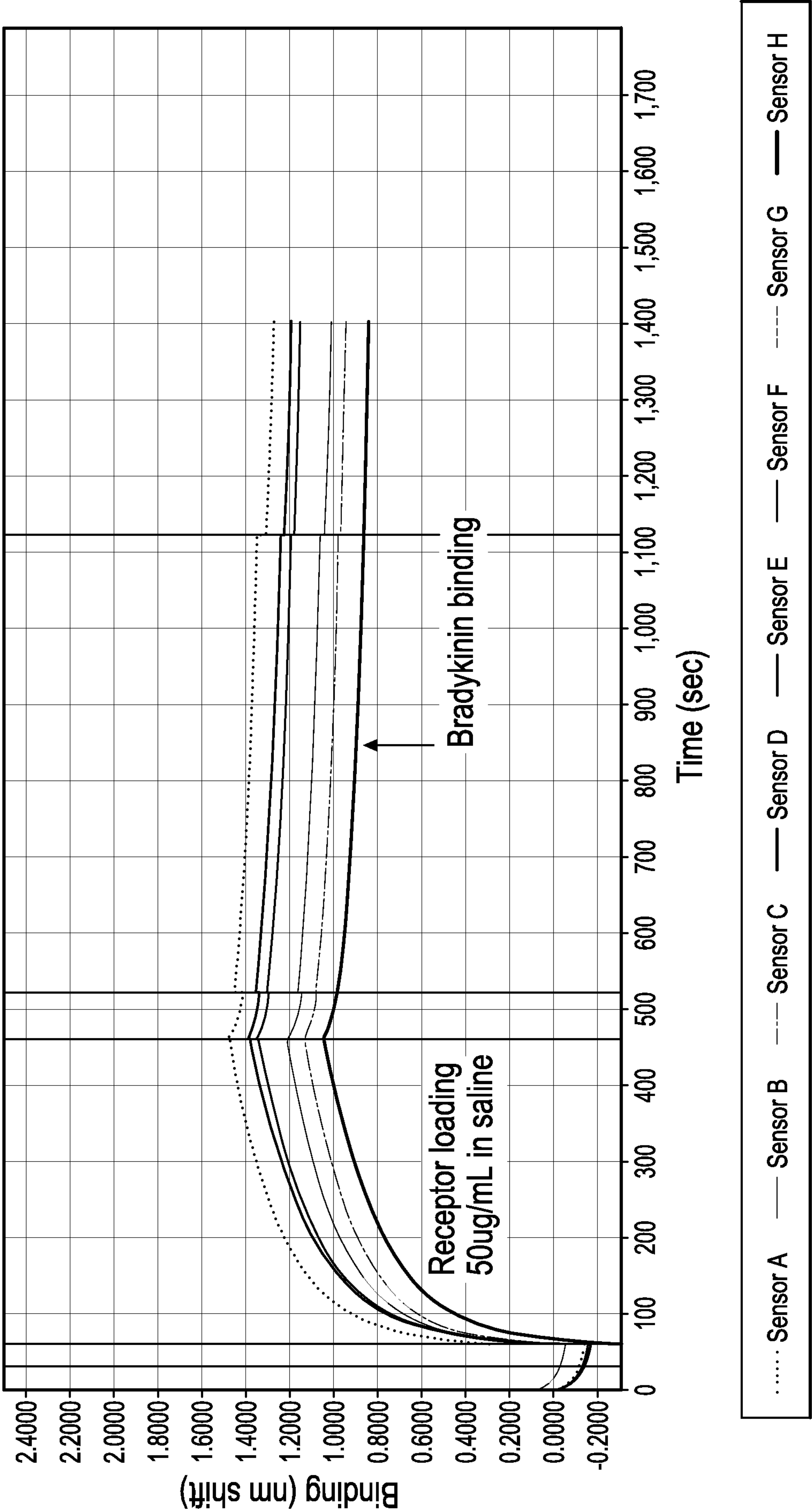


Fig. 75

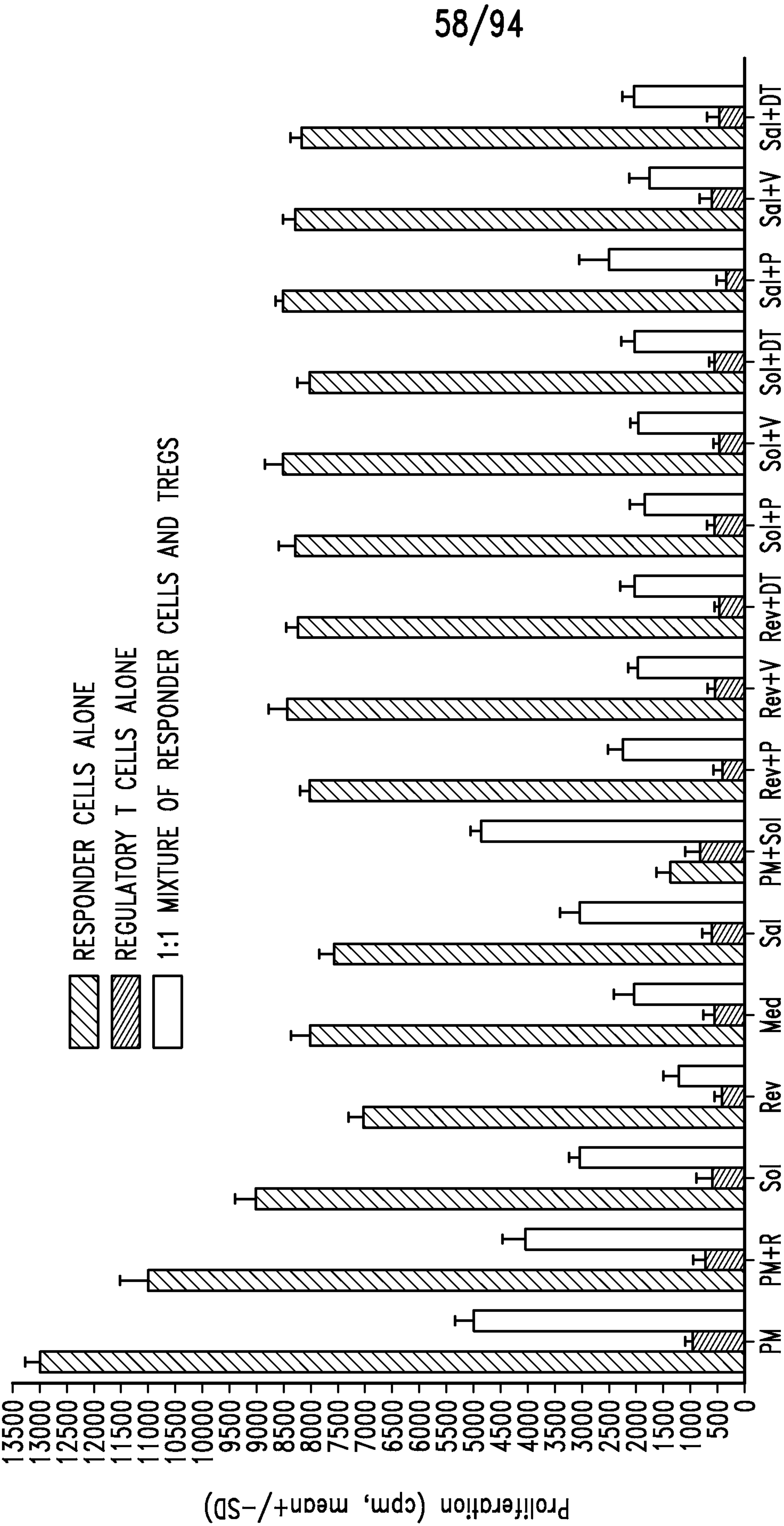
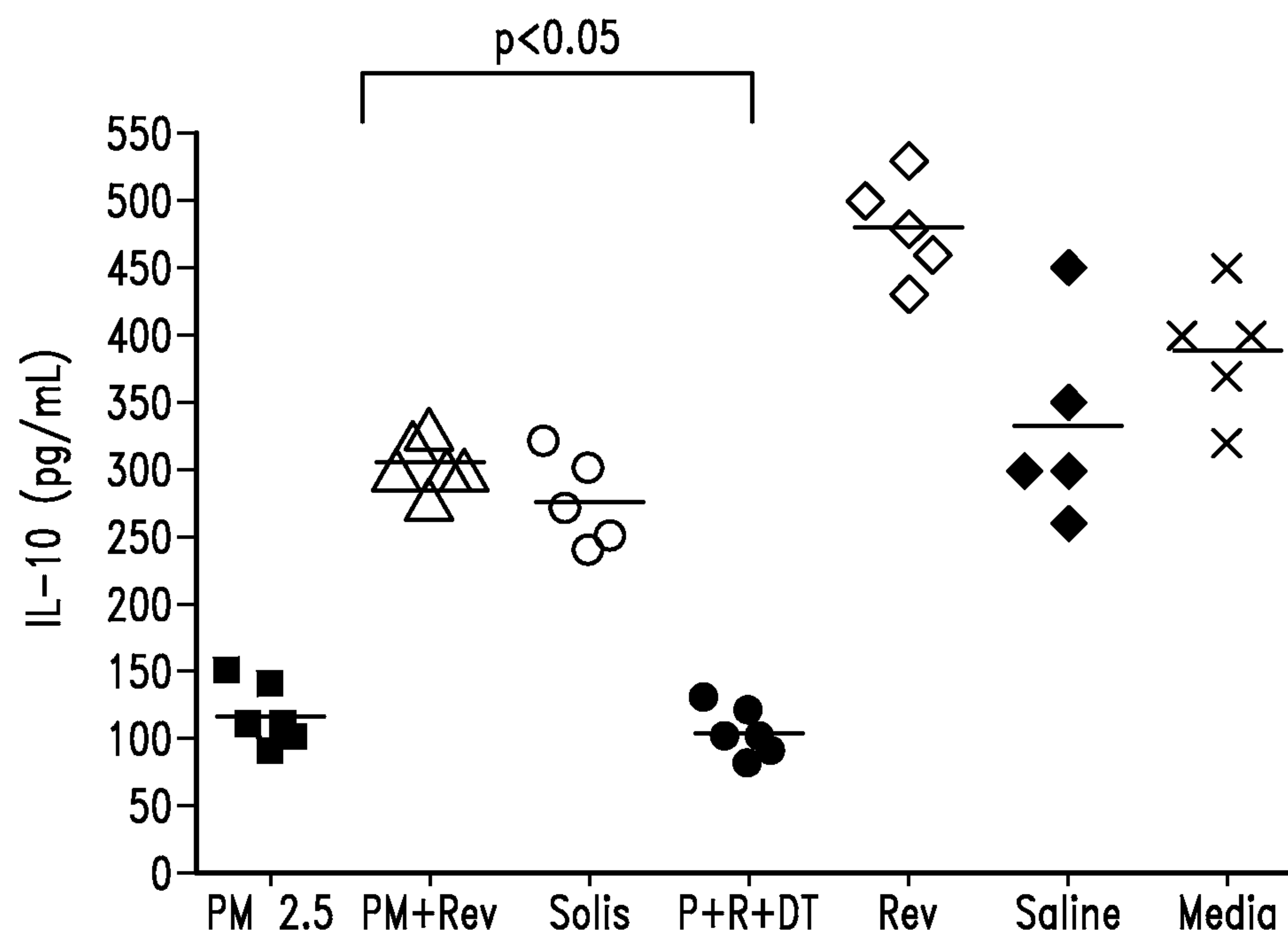
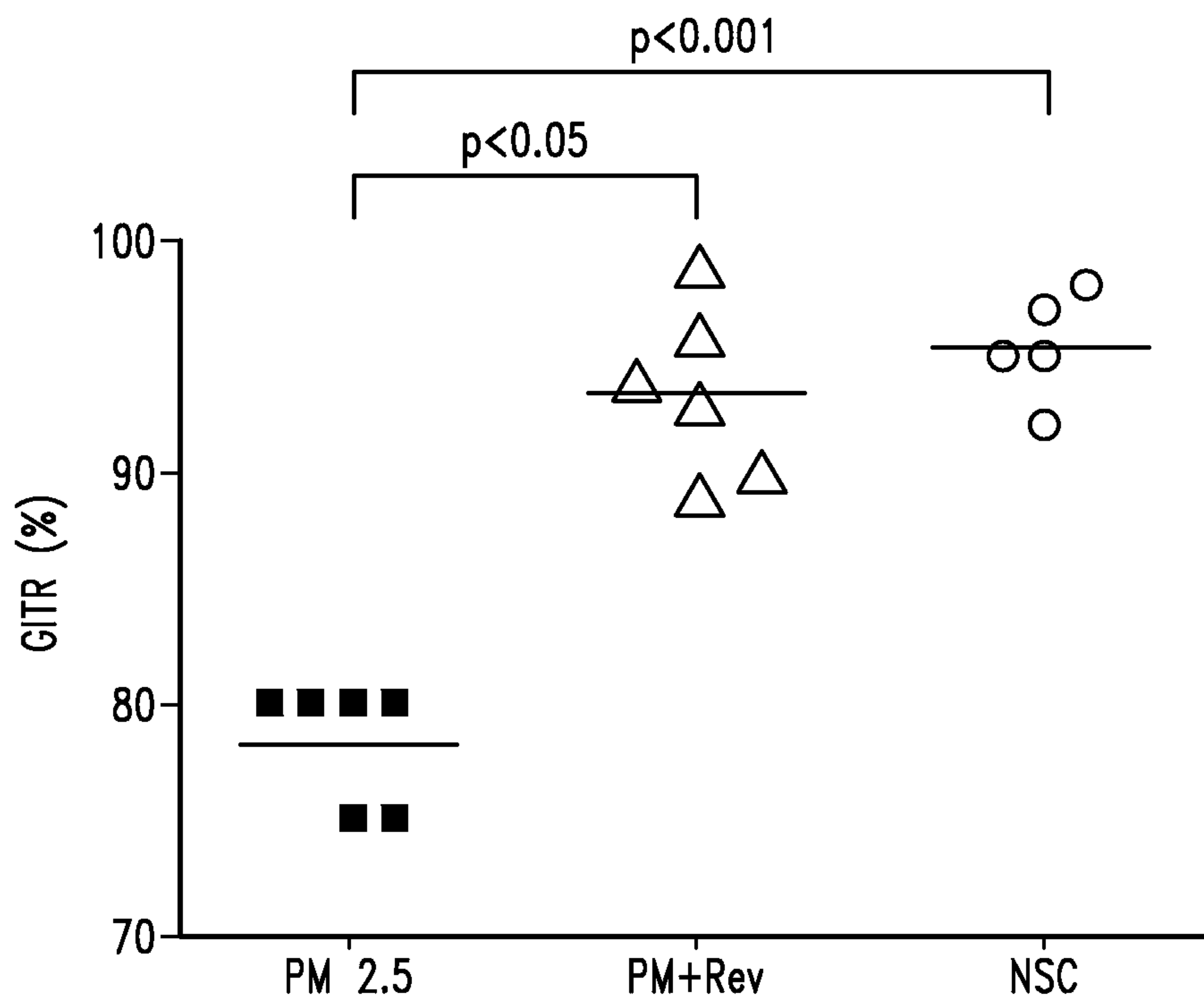
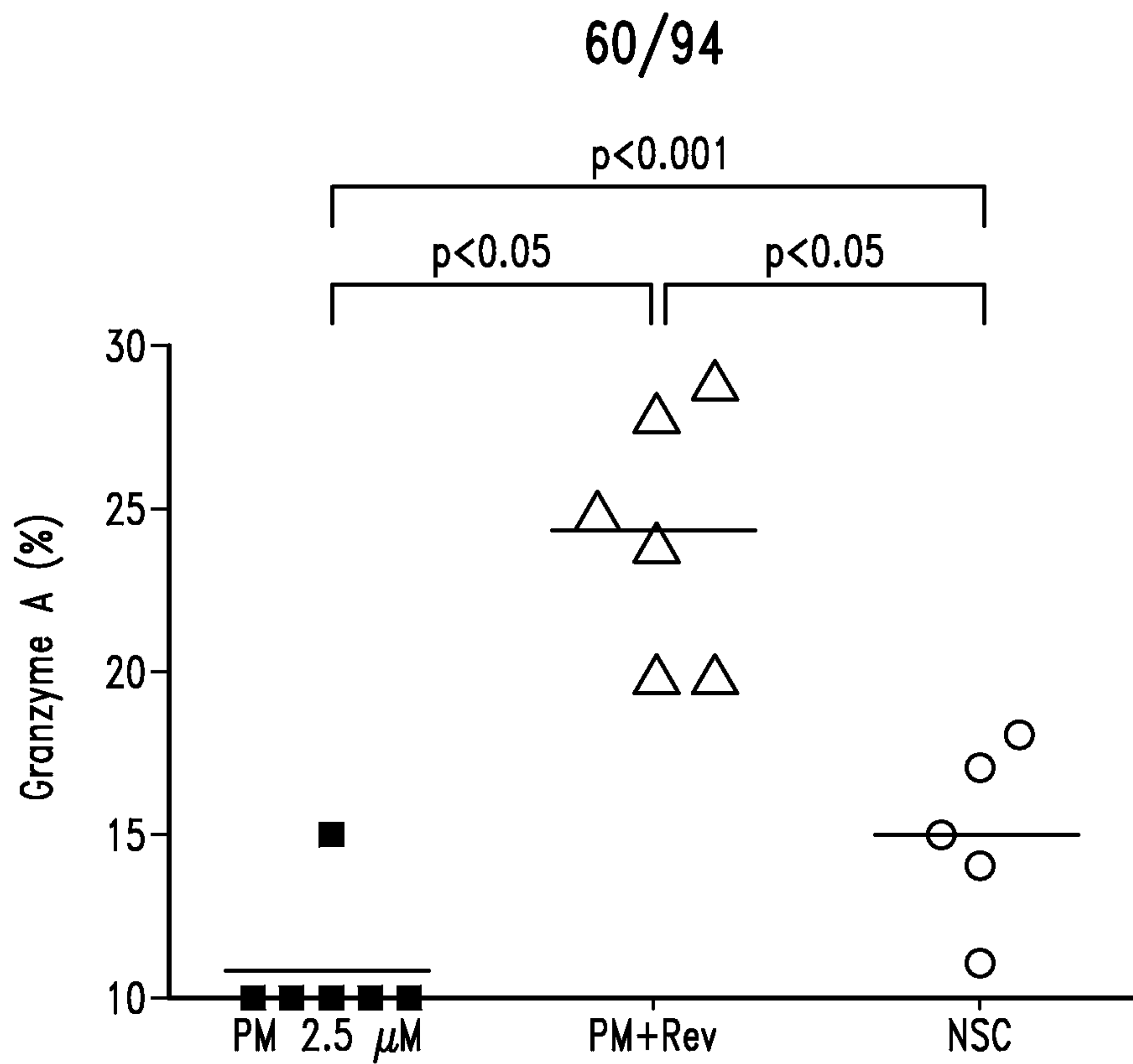
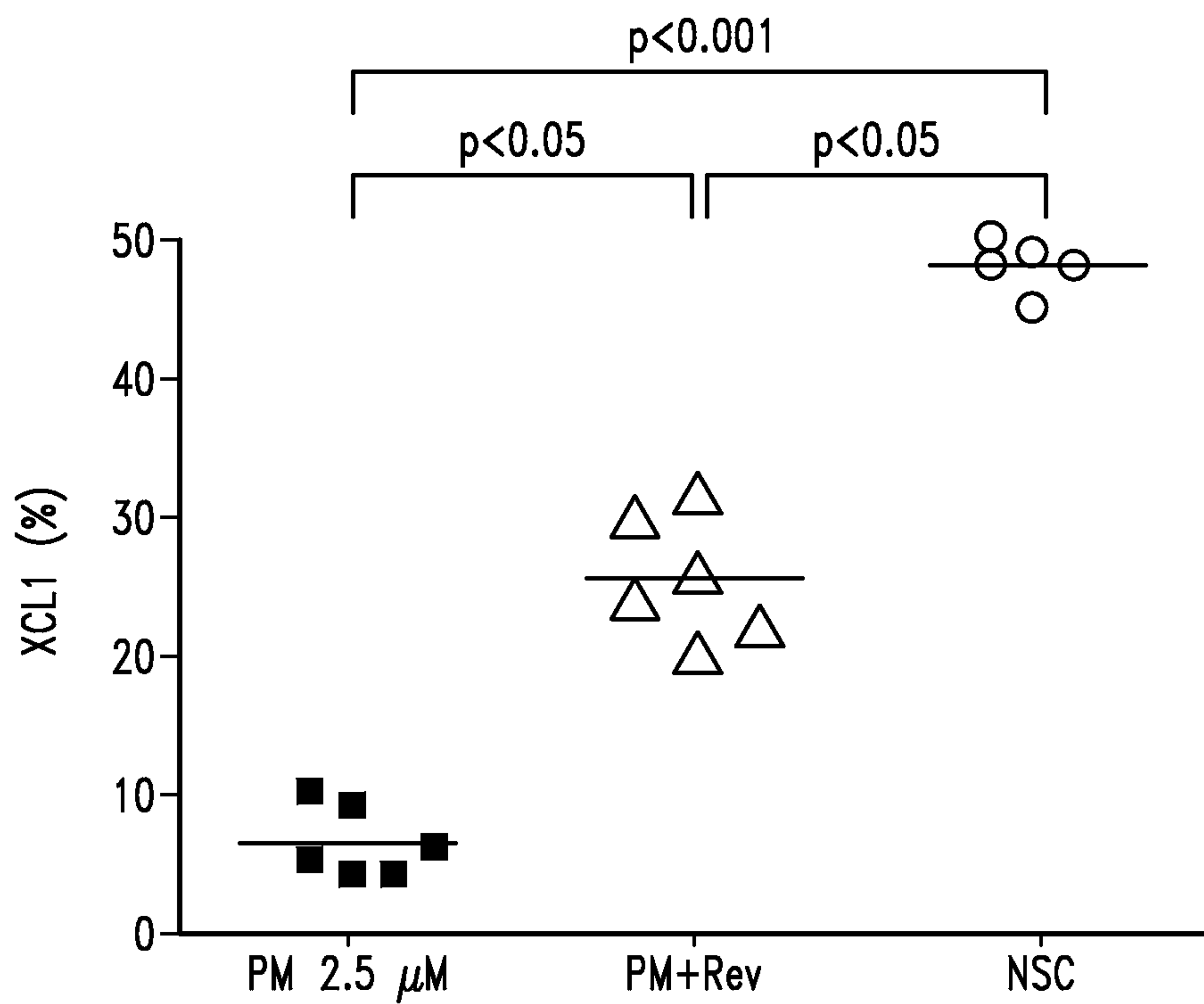
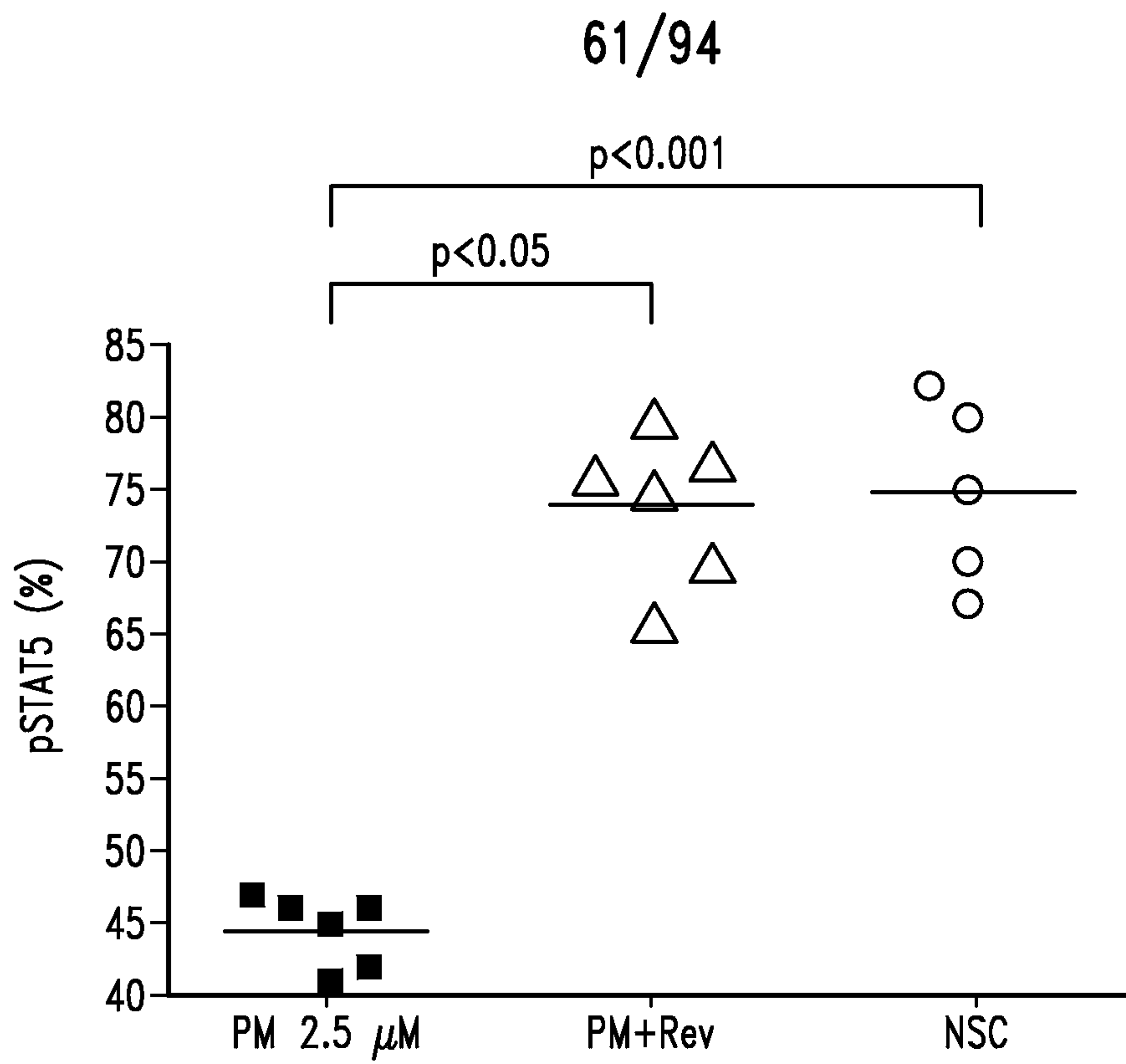
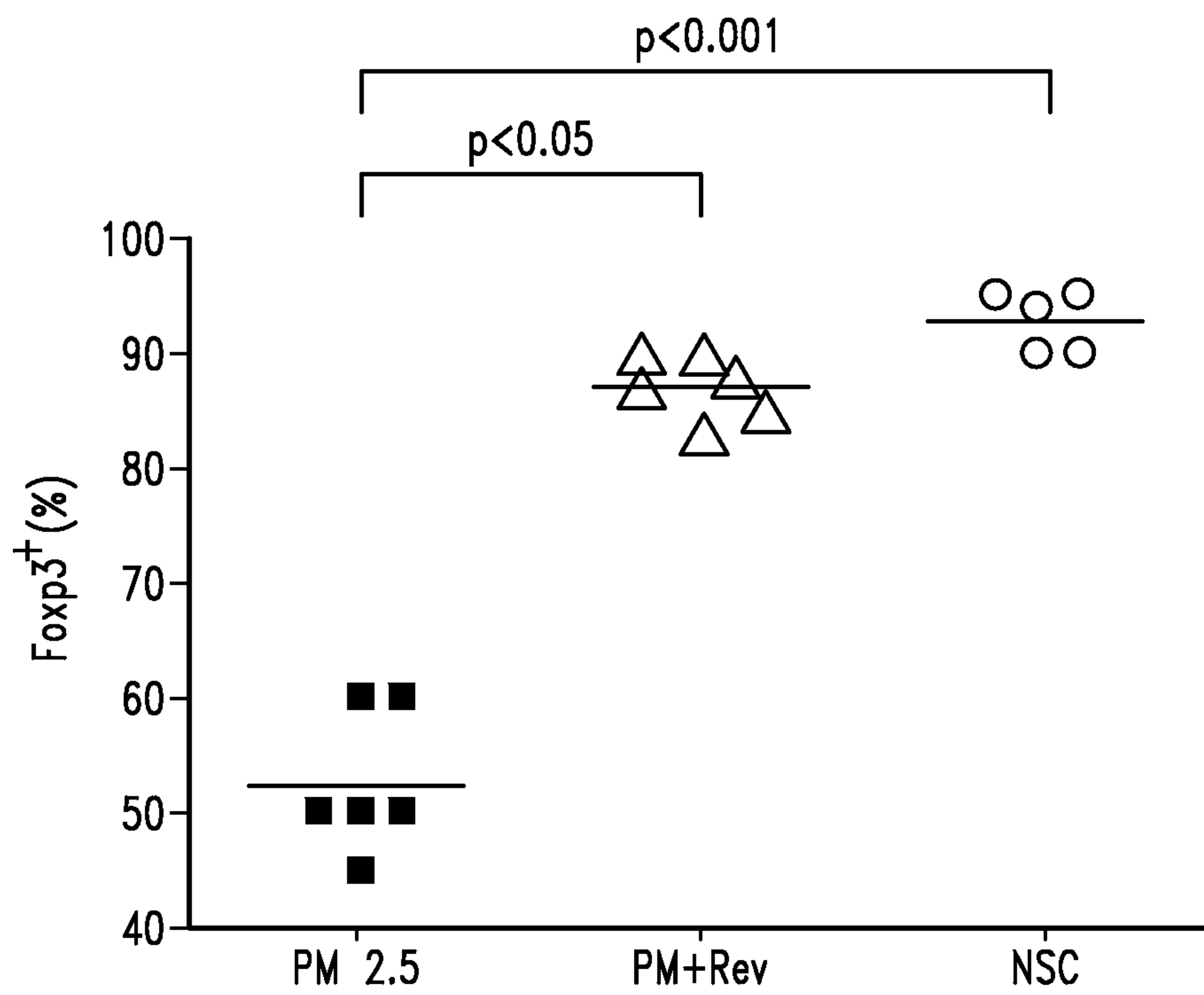


Fig. 76

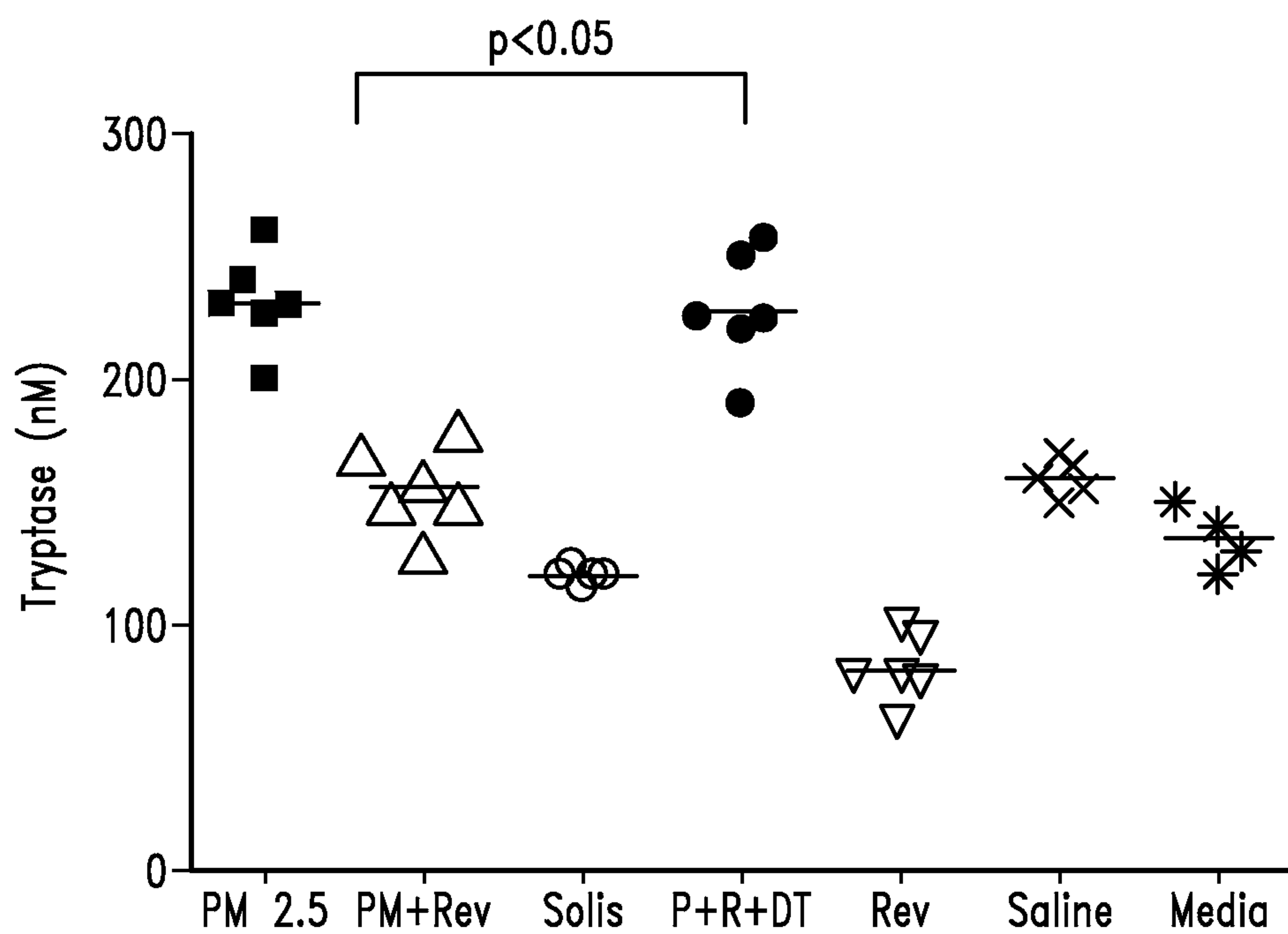
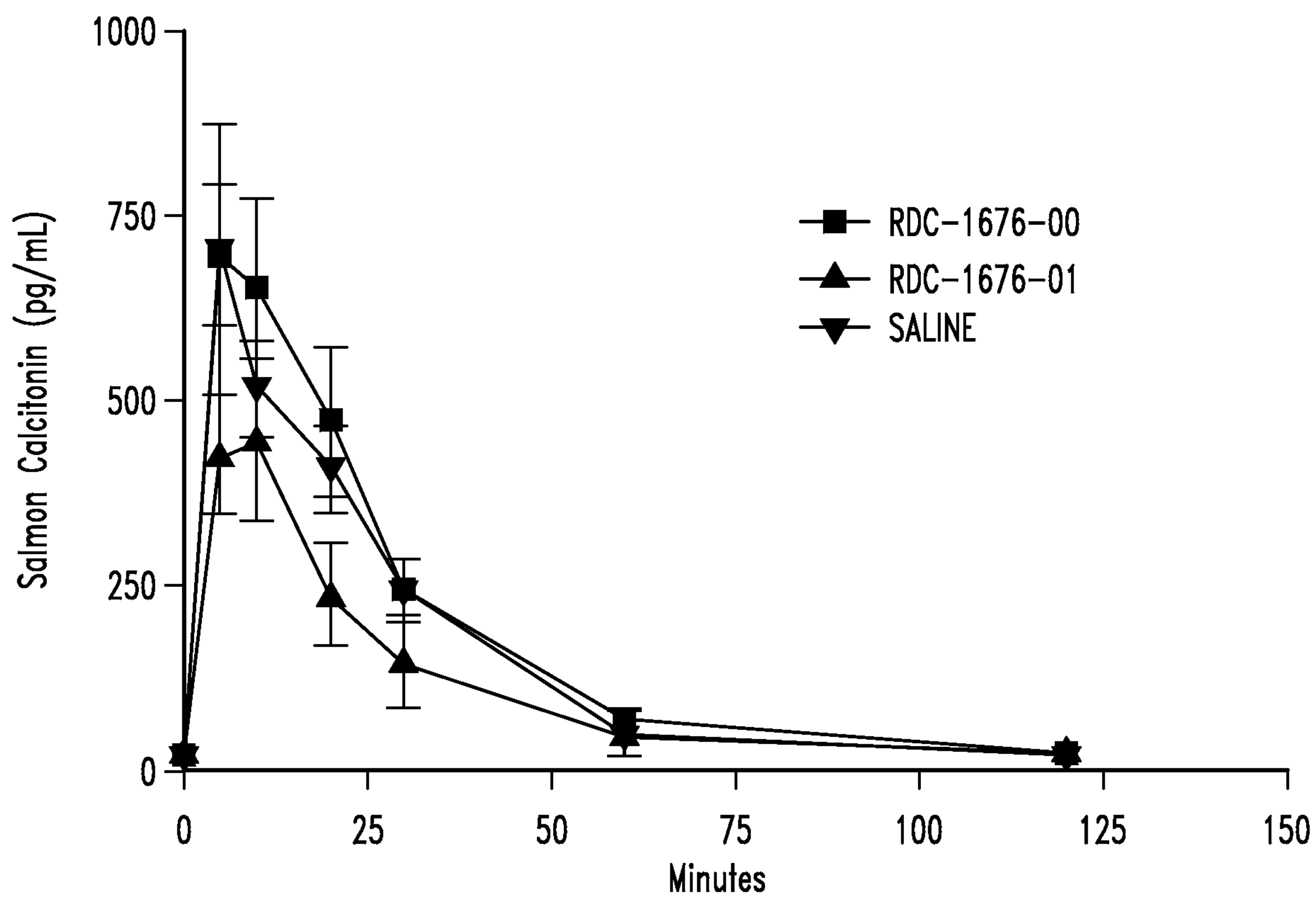
59/94

*Fig. 77**Fig. 78*

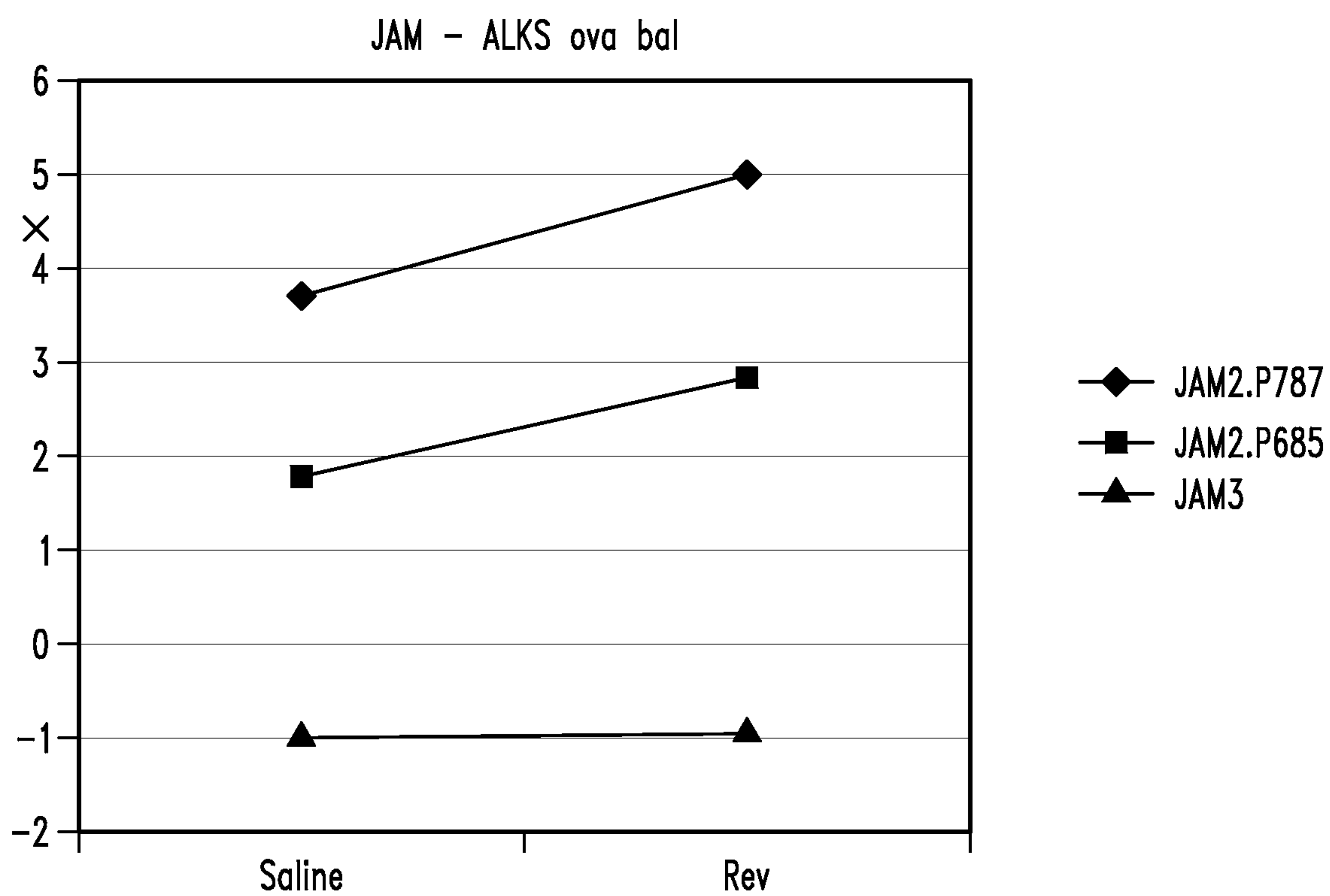
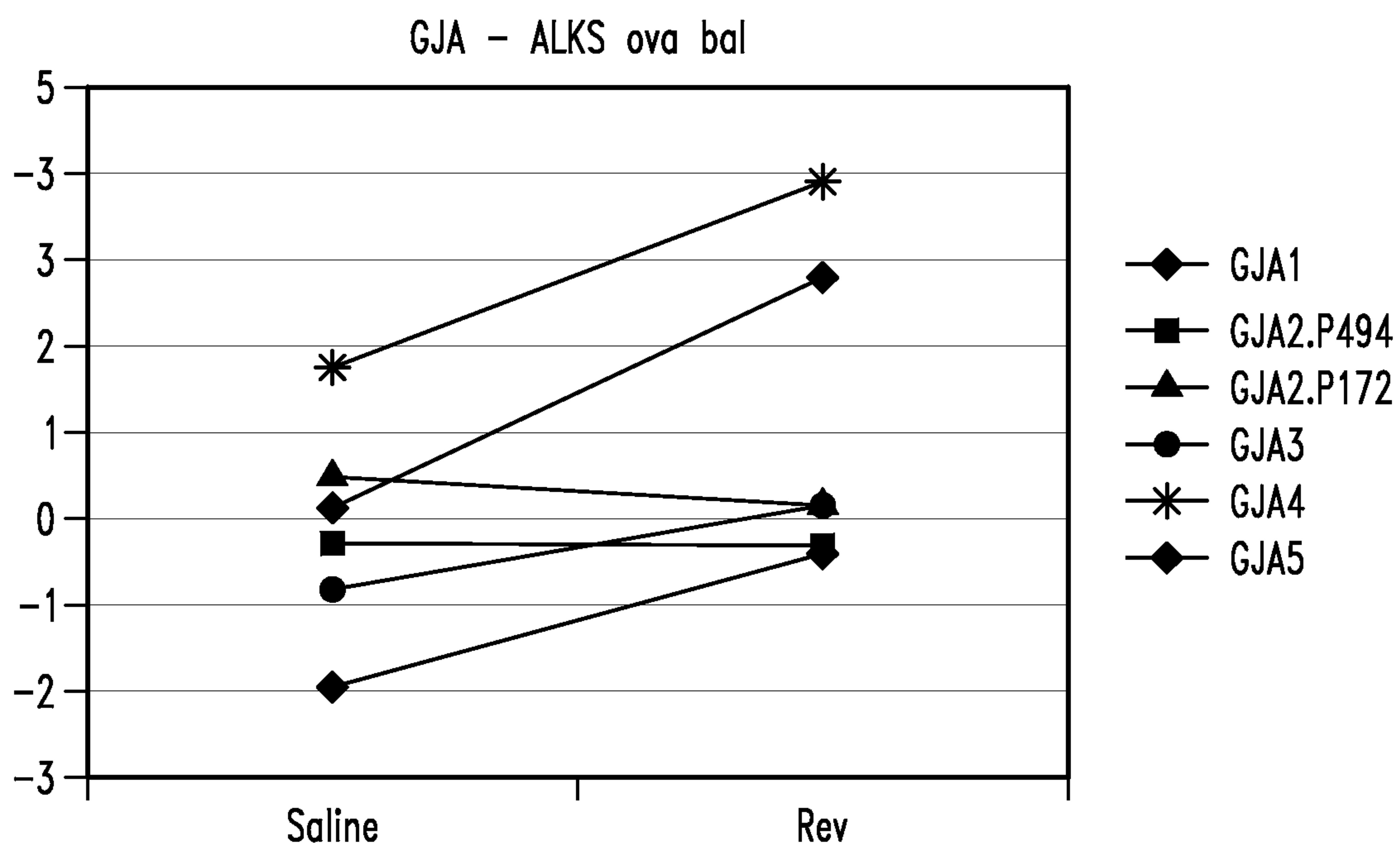
*Fig. 79**Fig. 80*

*Fig. 81**Fig. 82*

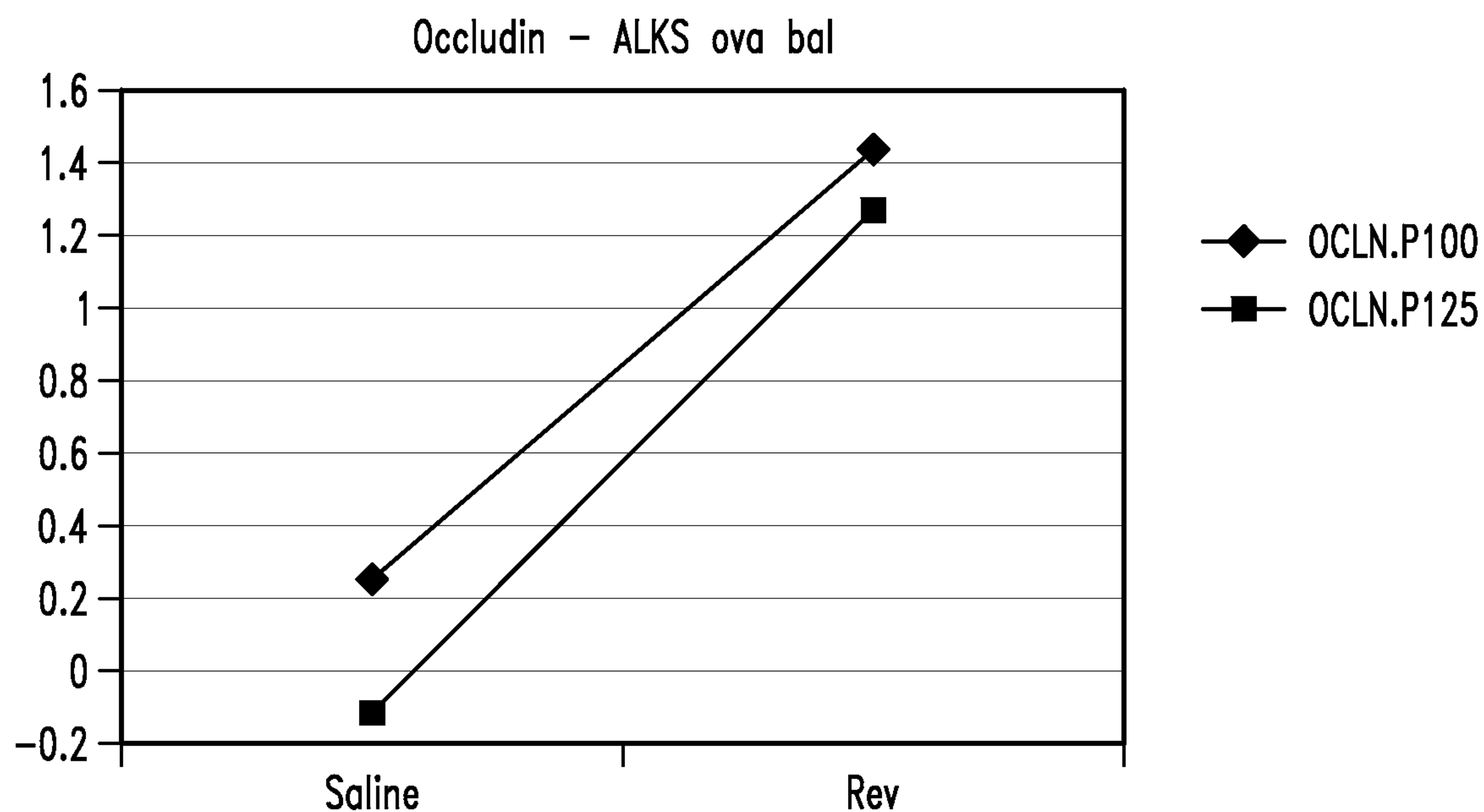
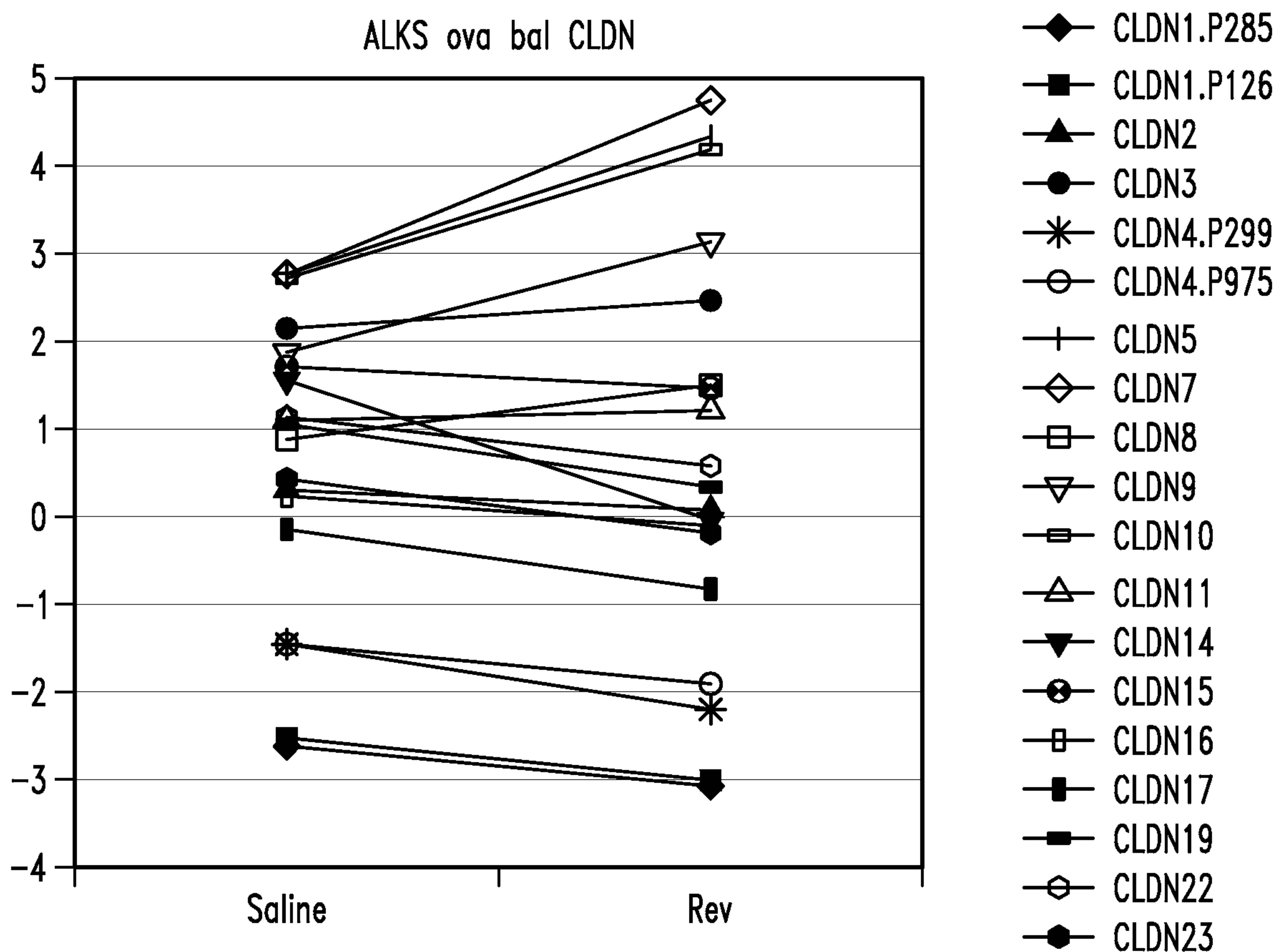
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*Fig. 83**Fig. 84*

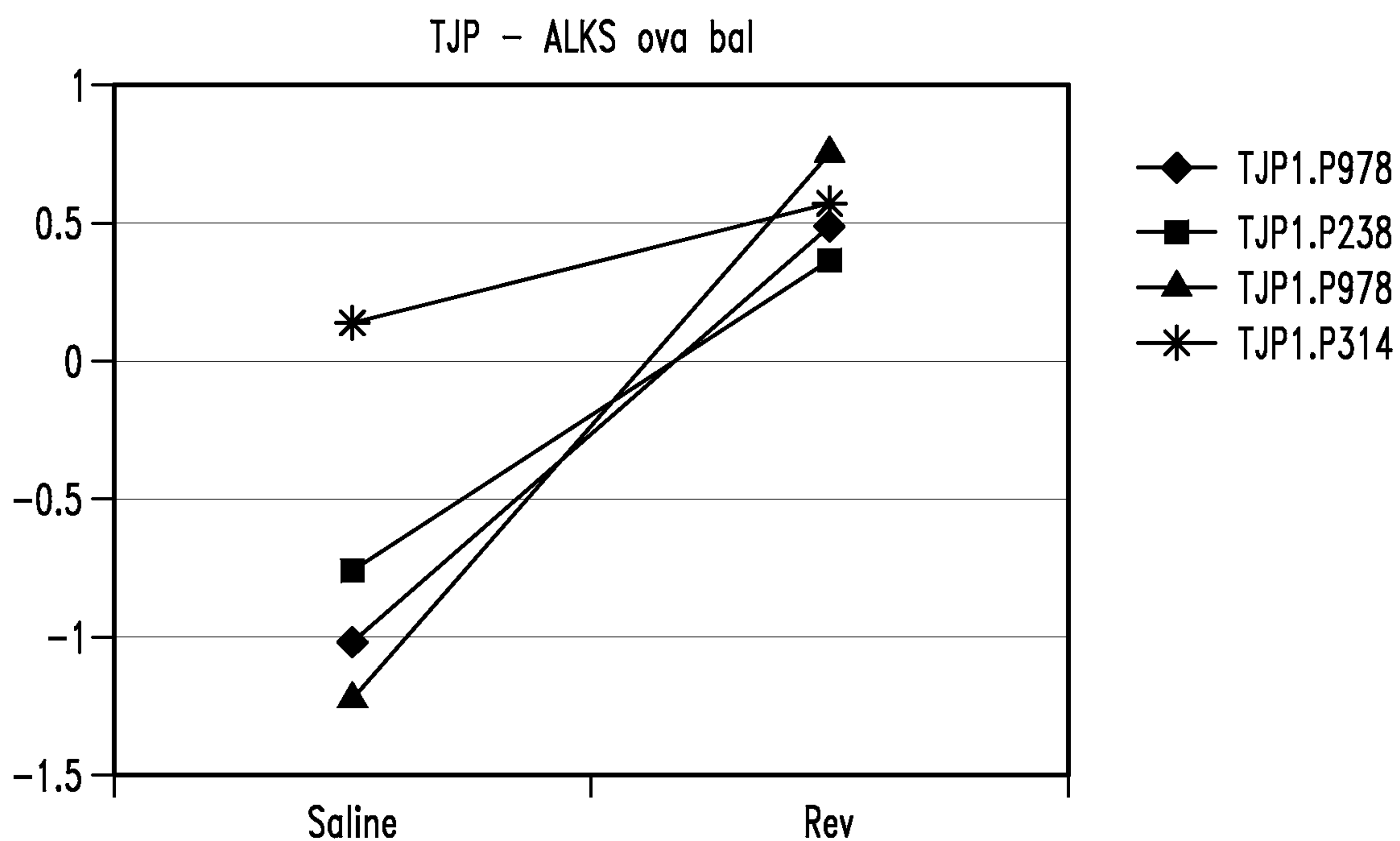
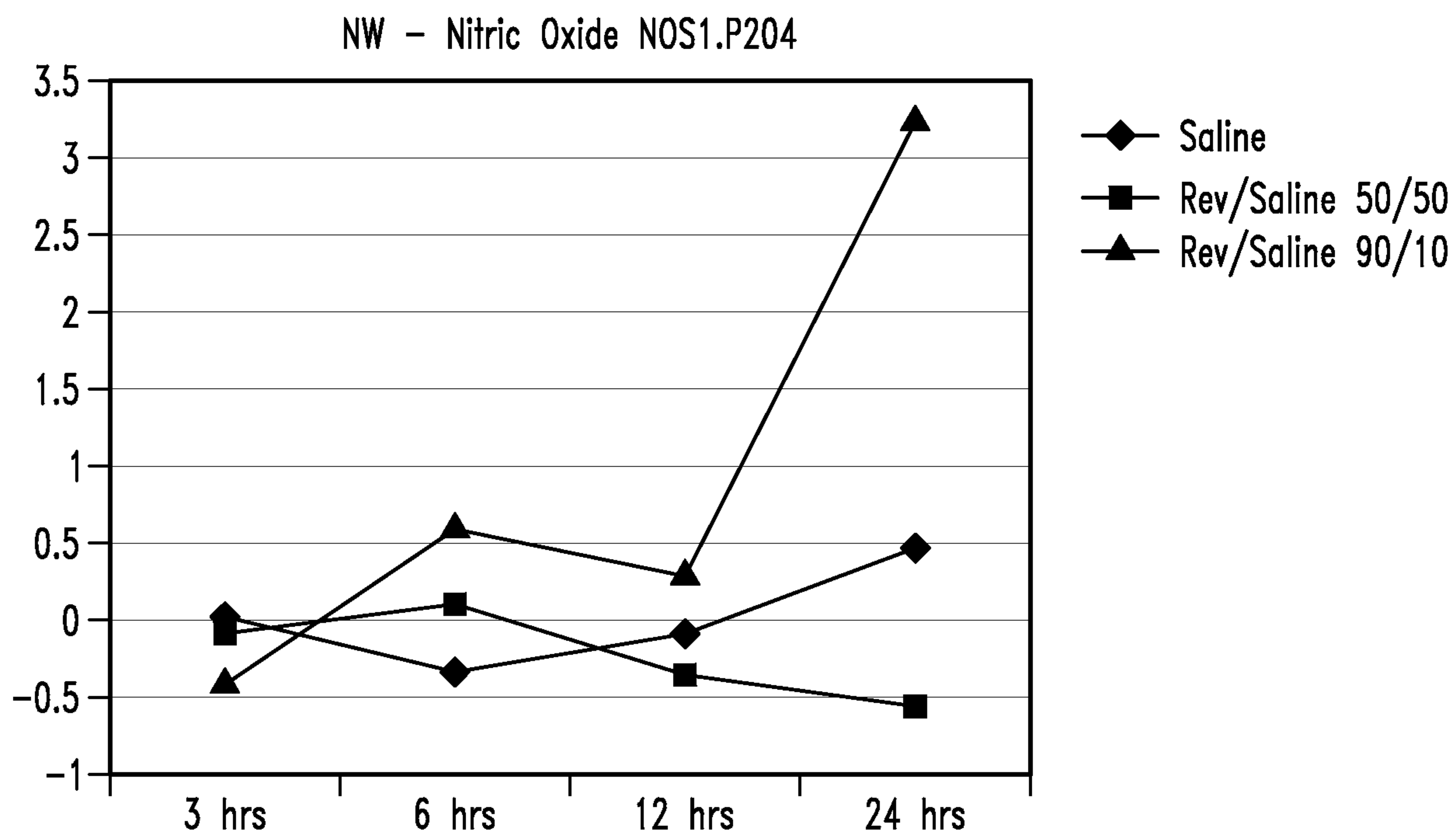
63/94

*Fig. 85**Fig. 86*

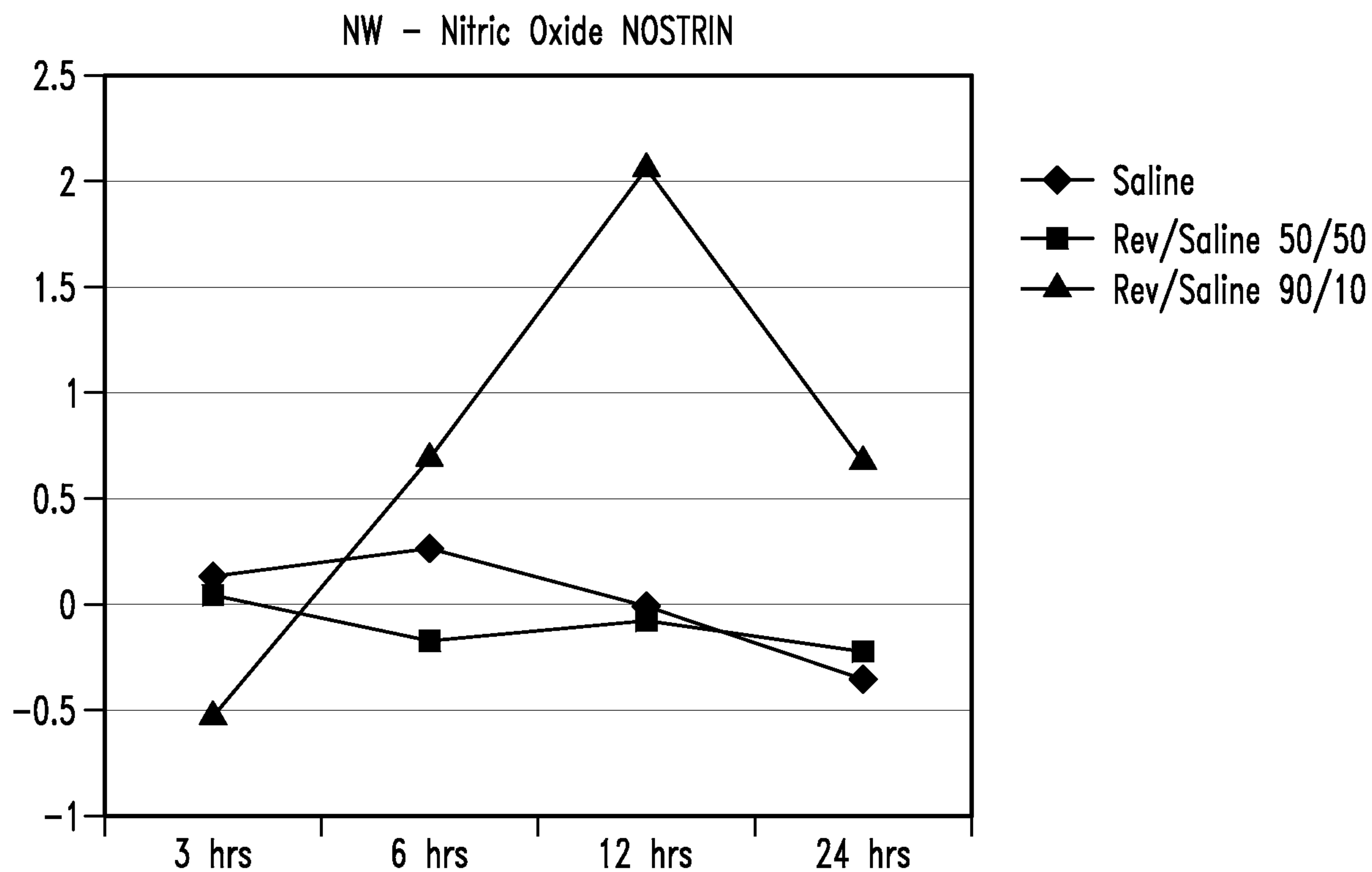
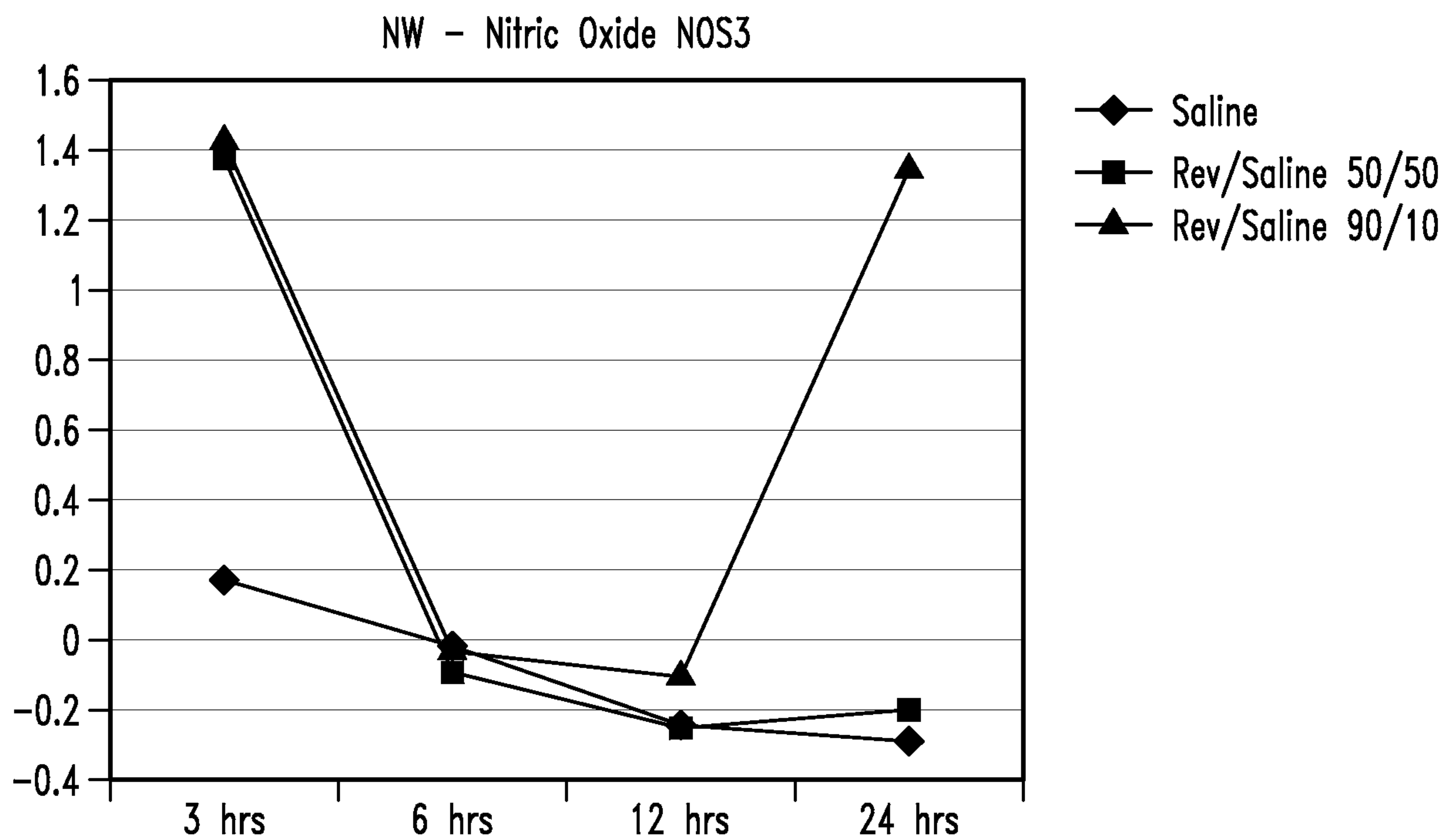
64/94

*Fig. 87**Fig. 88*

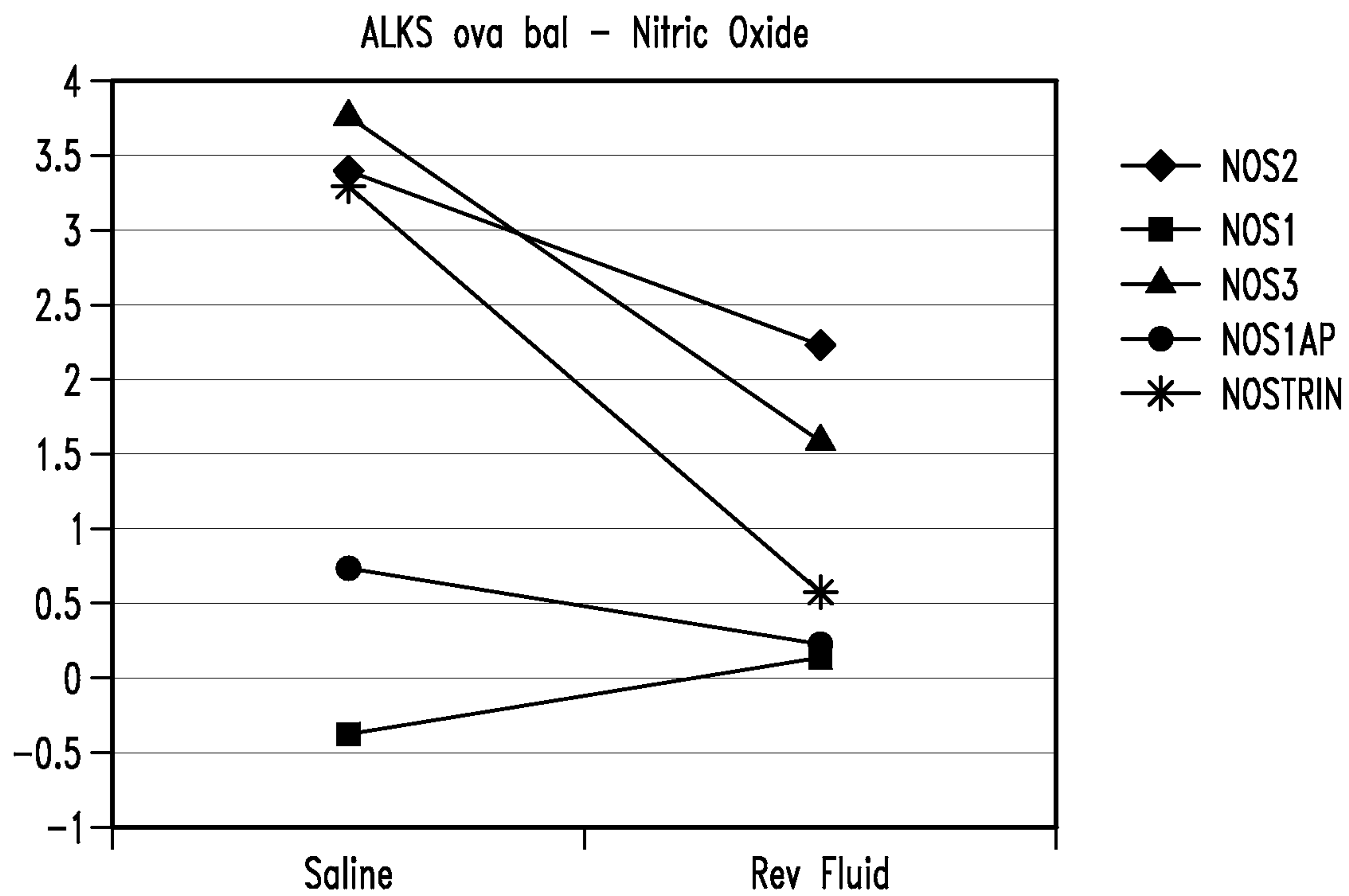
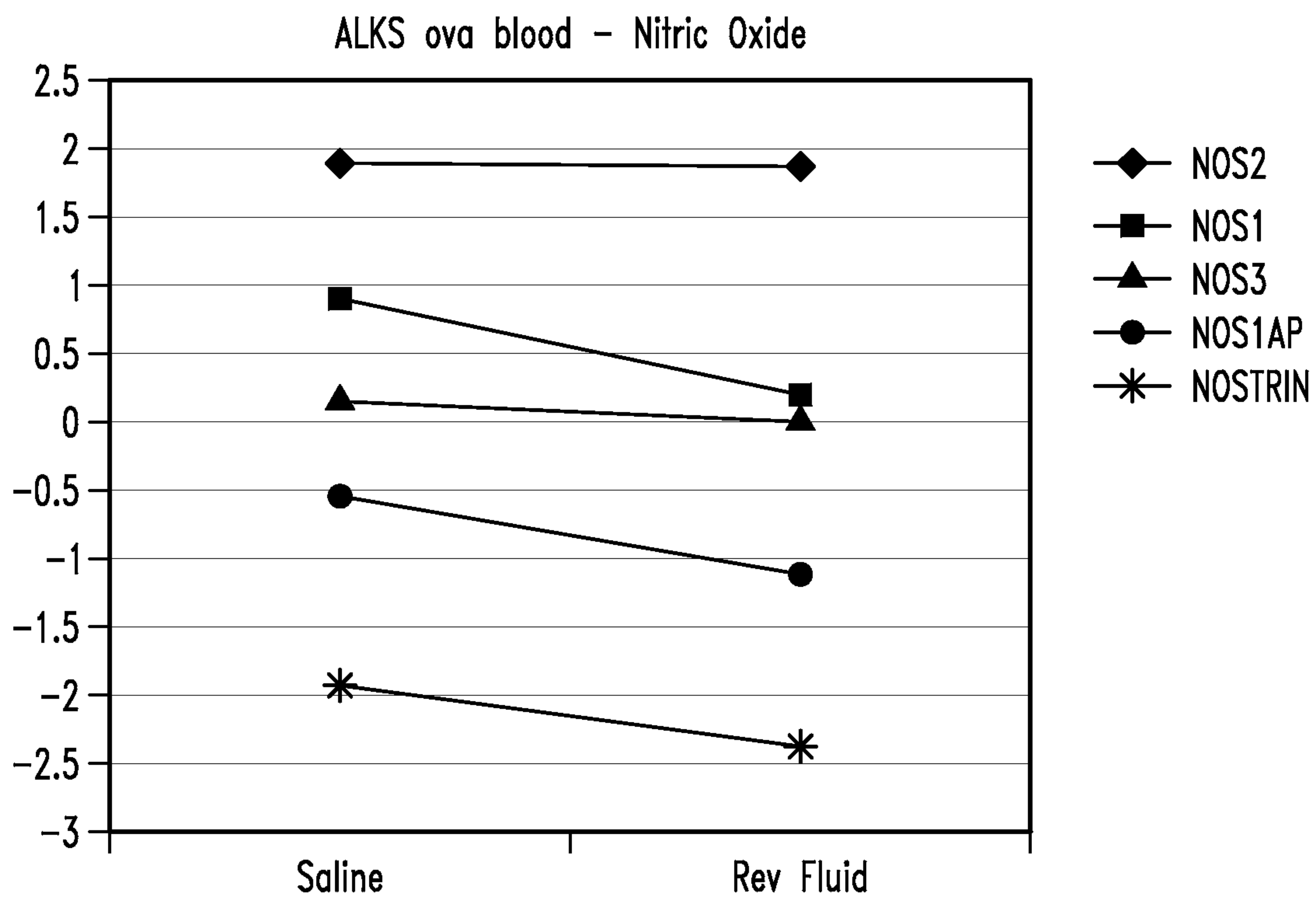
65/94

*Fig. 89**Fig. 90*

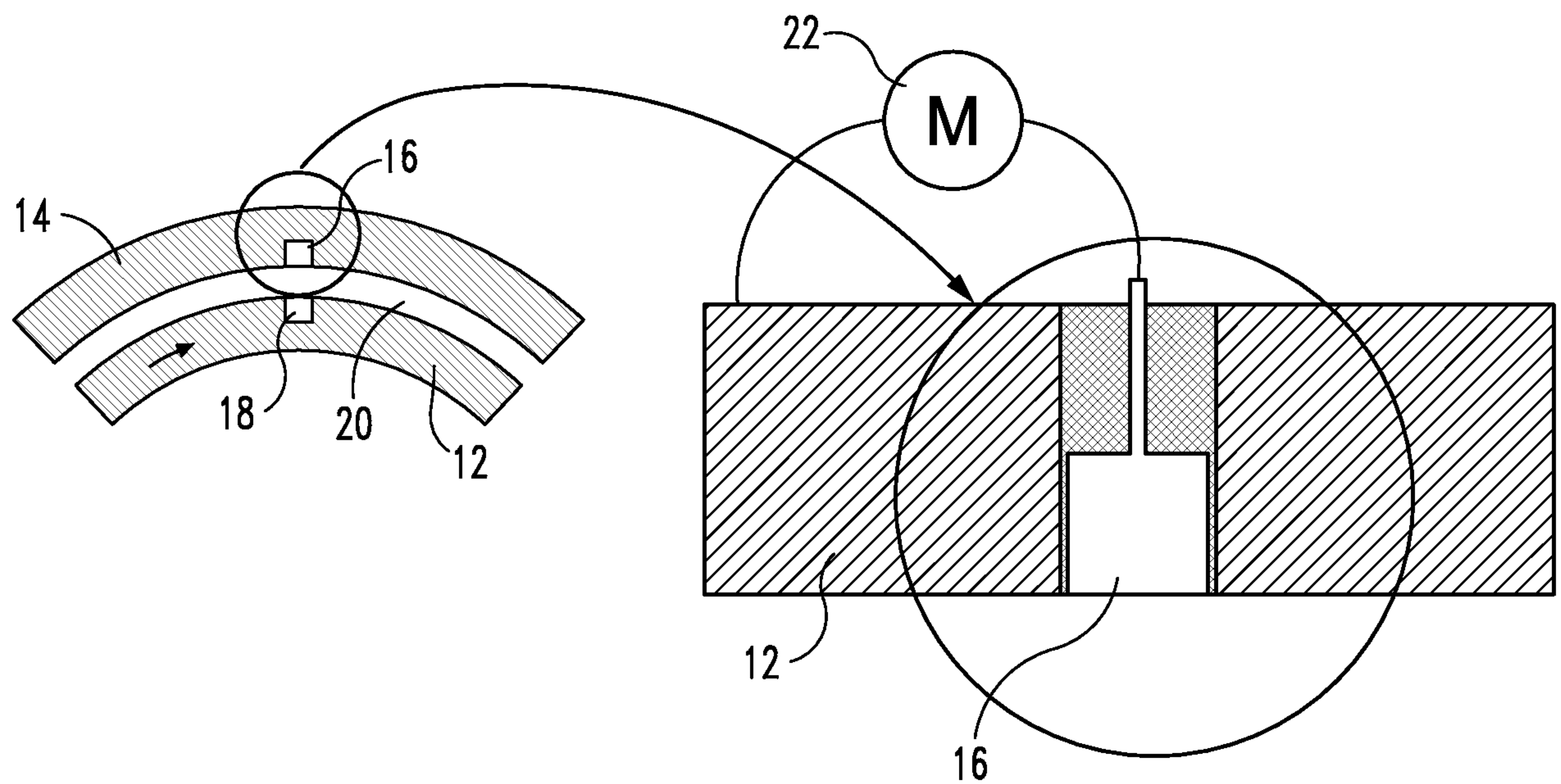
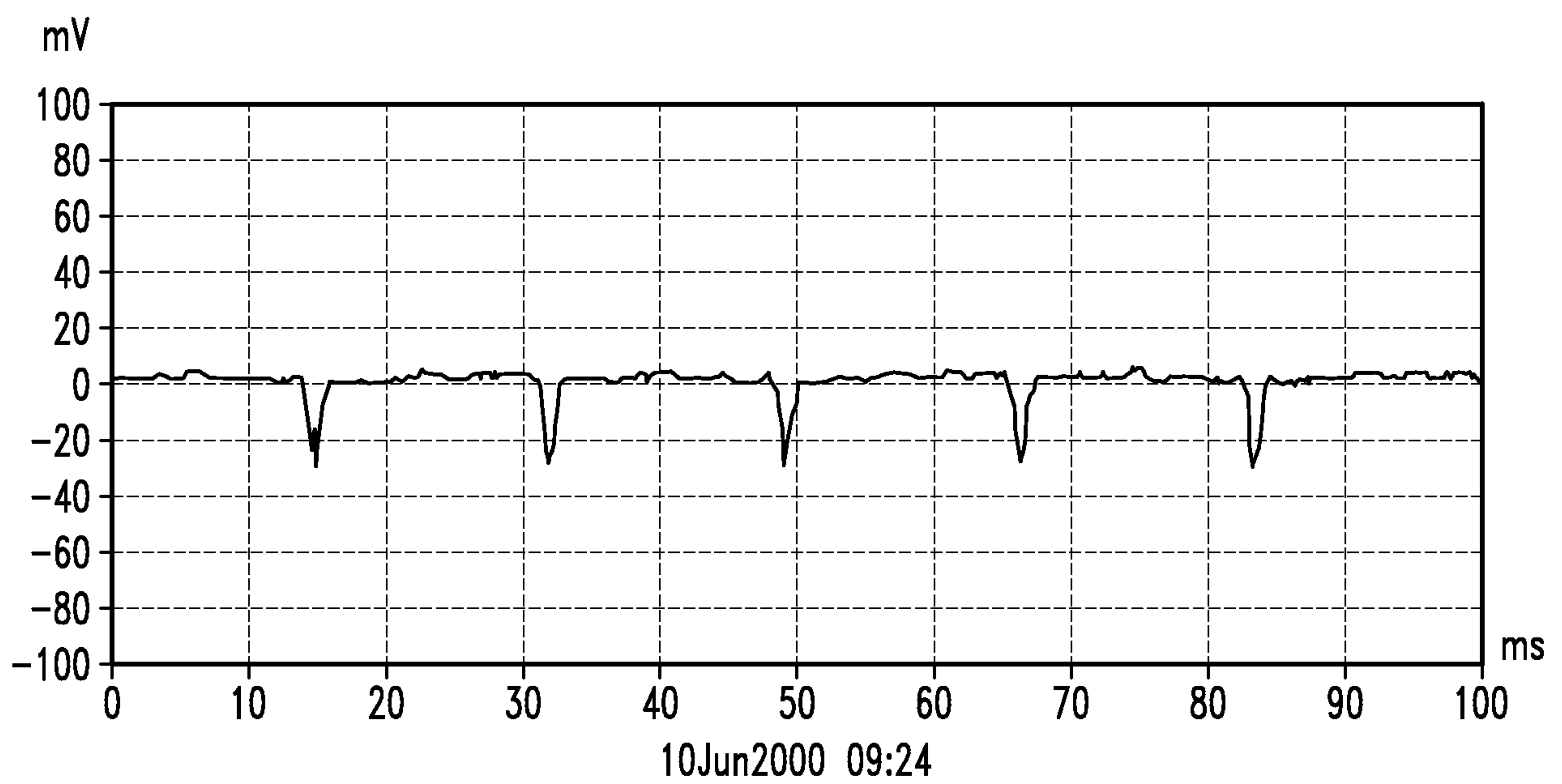
66/94

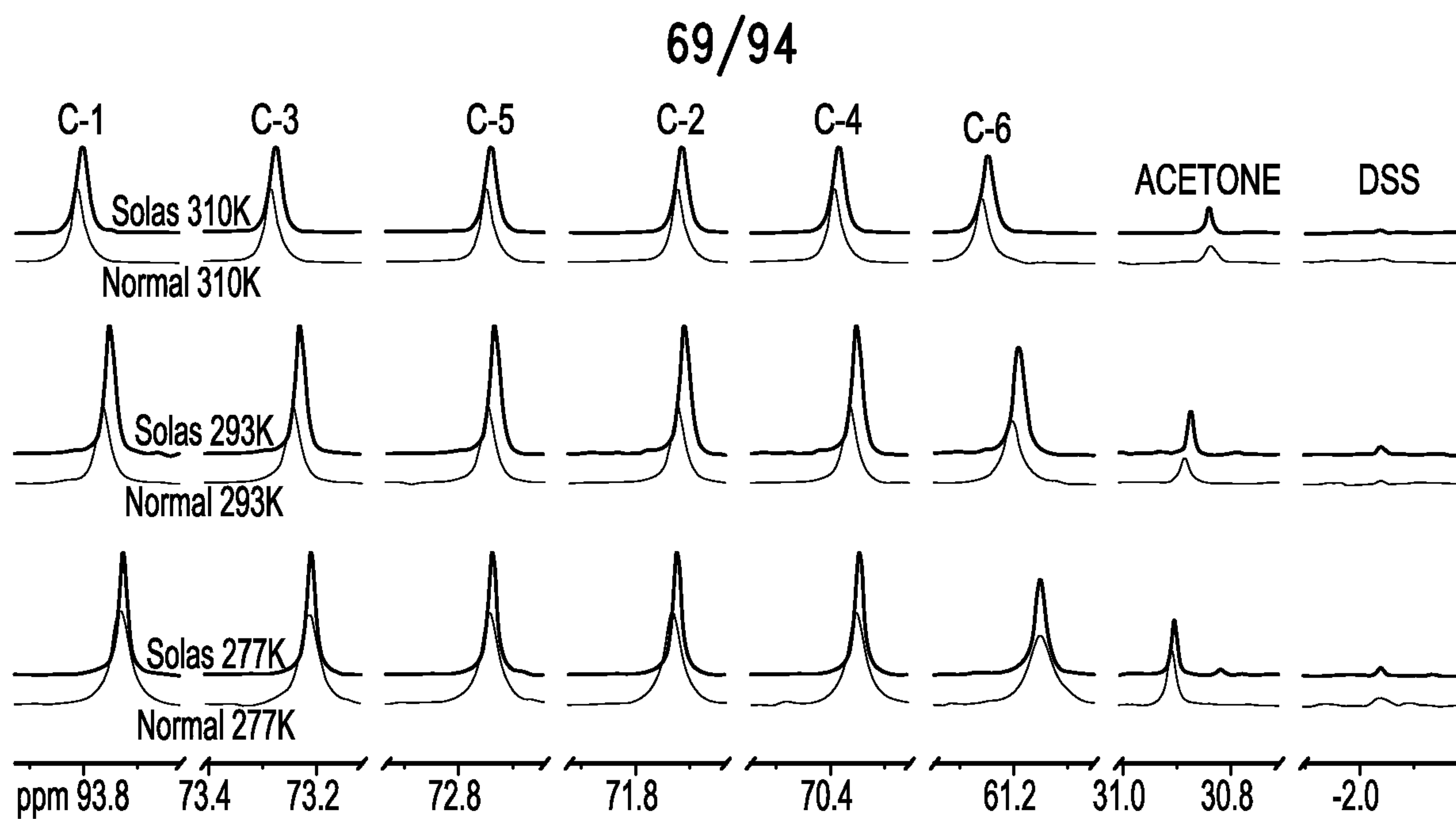
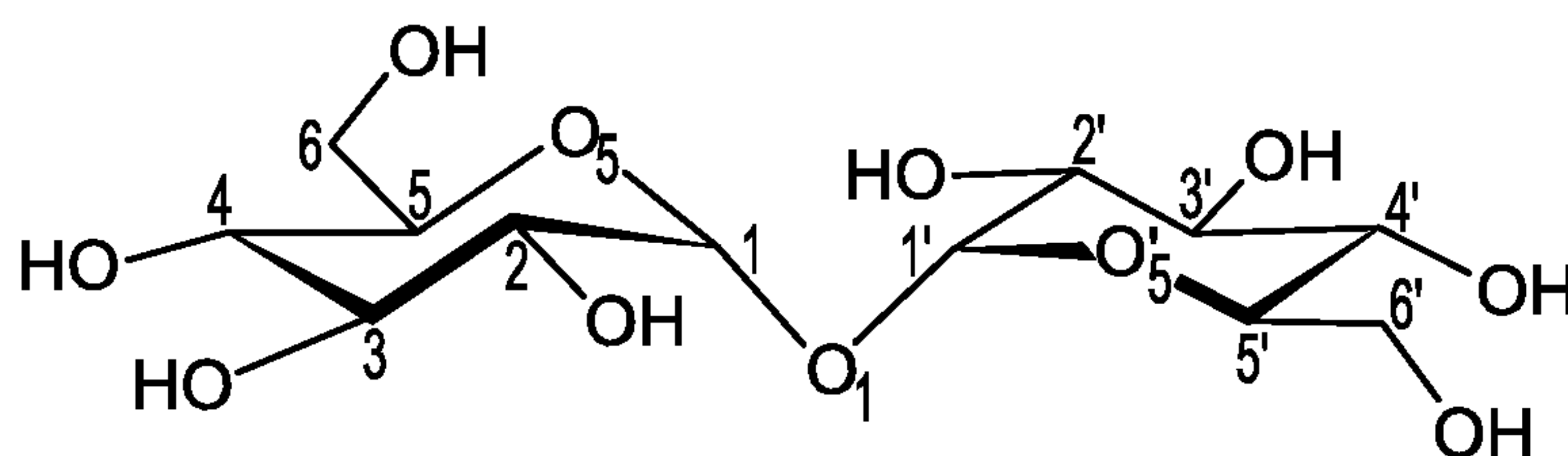
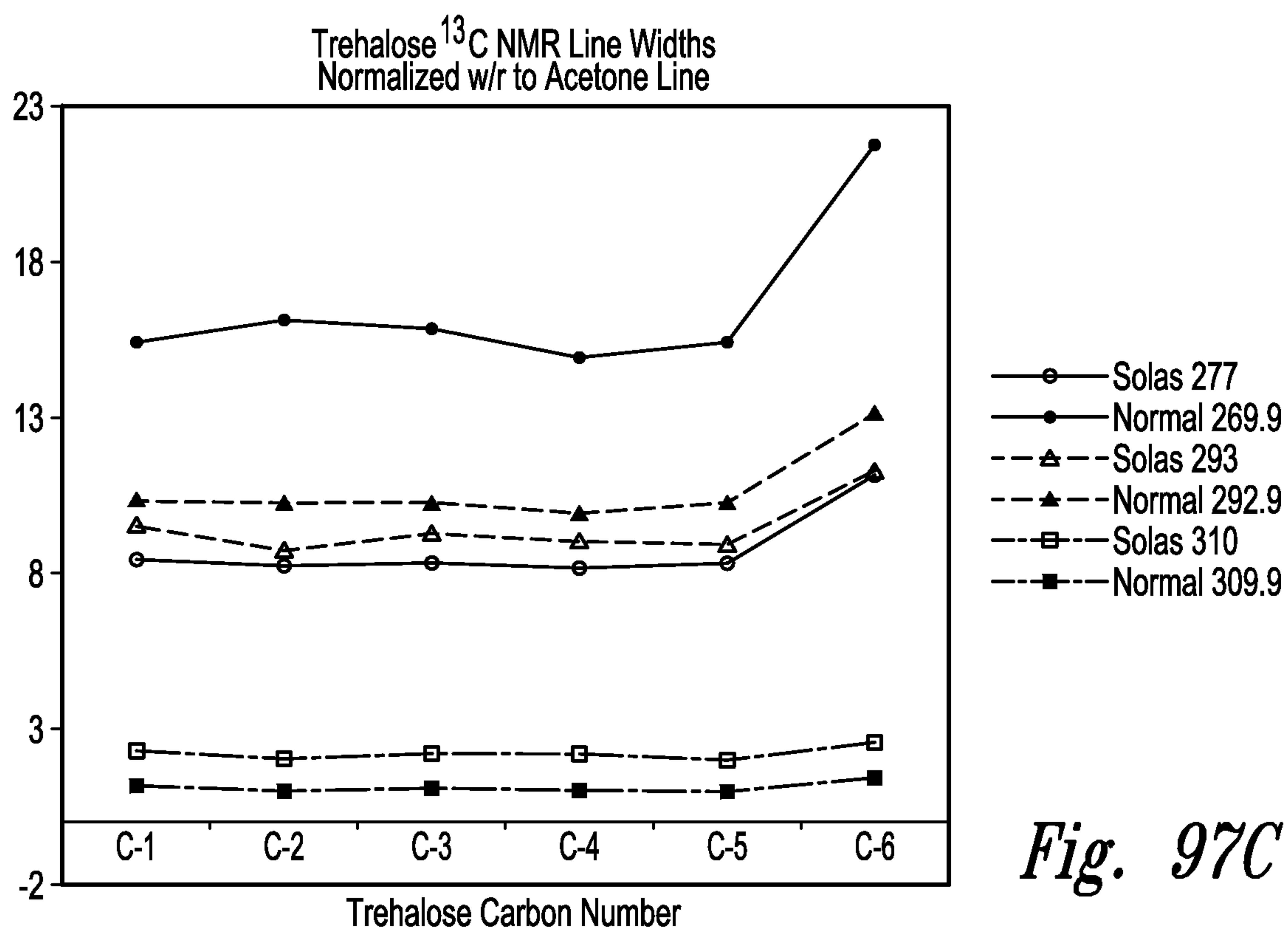
*Fig. 91**Fig. 92*

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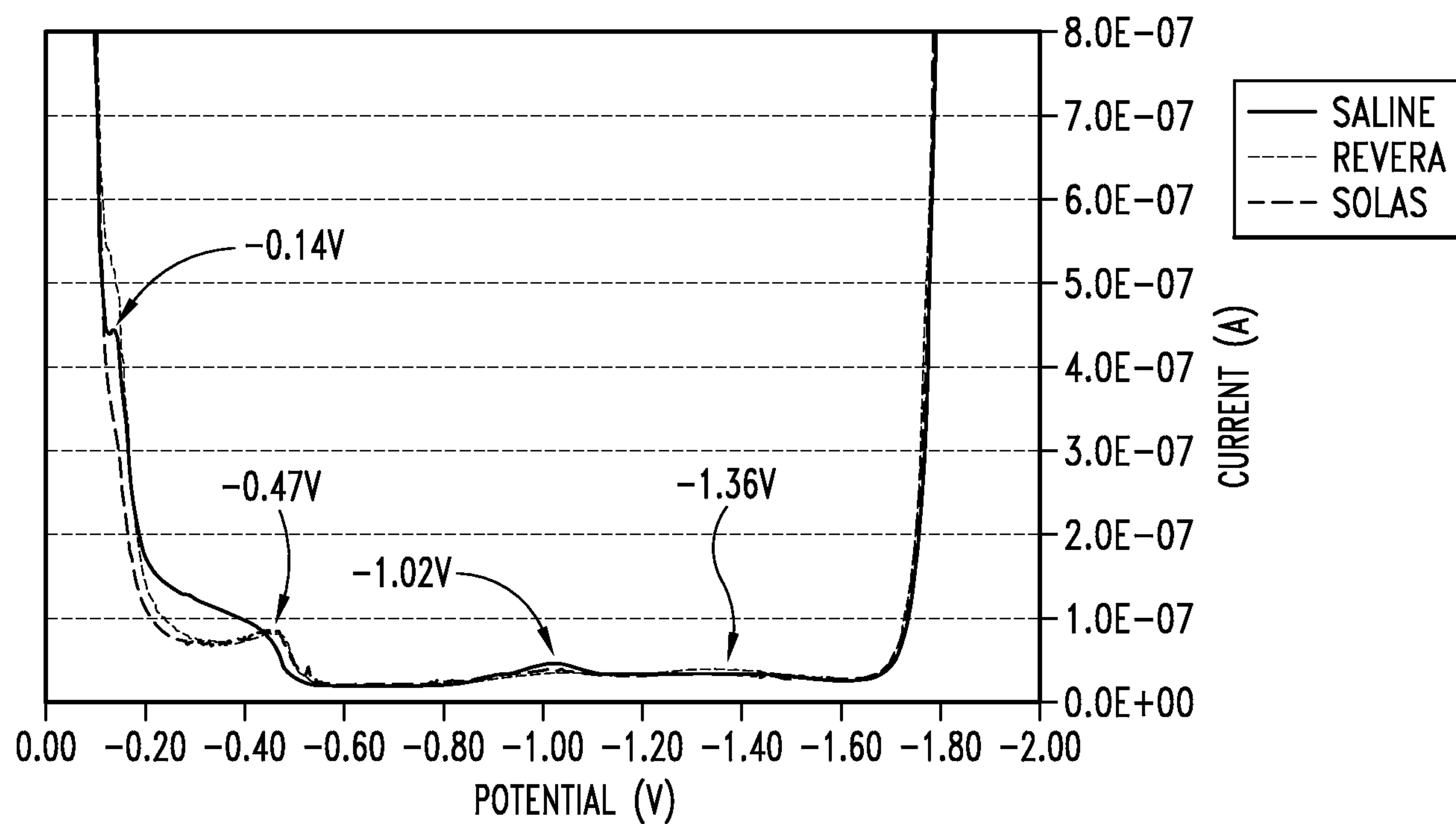
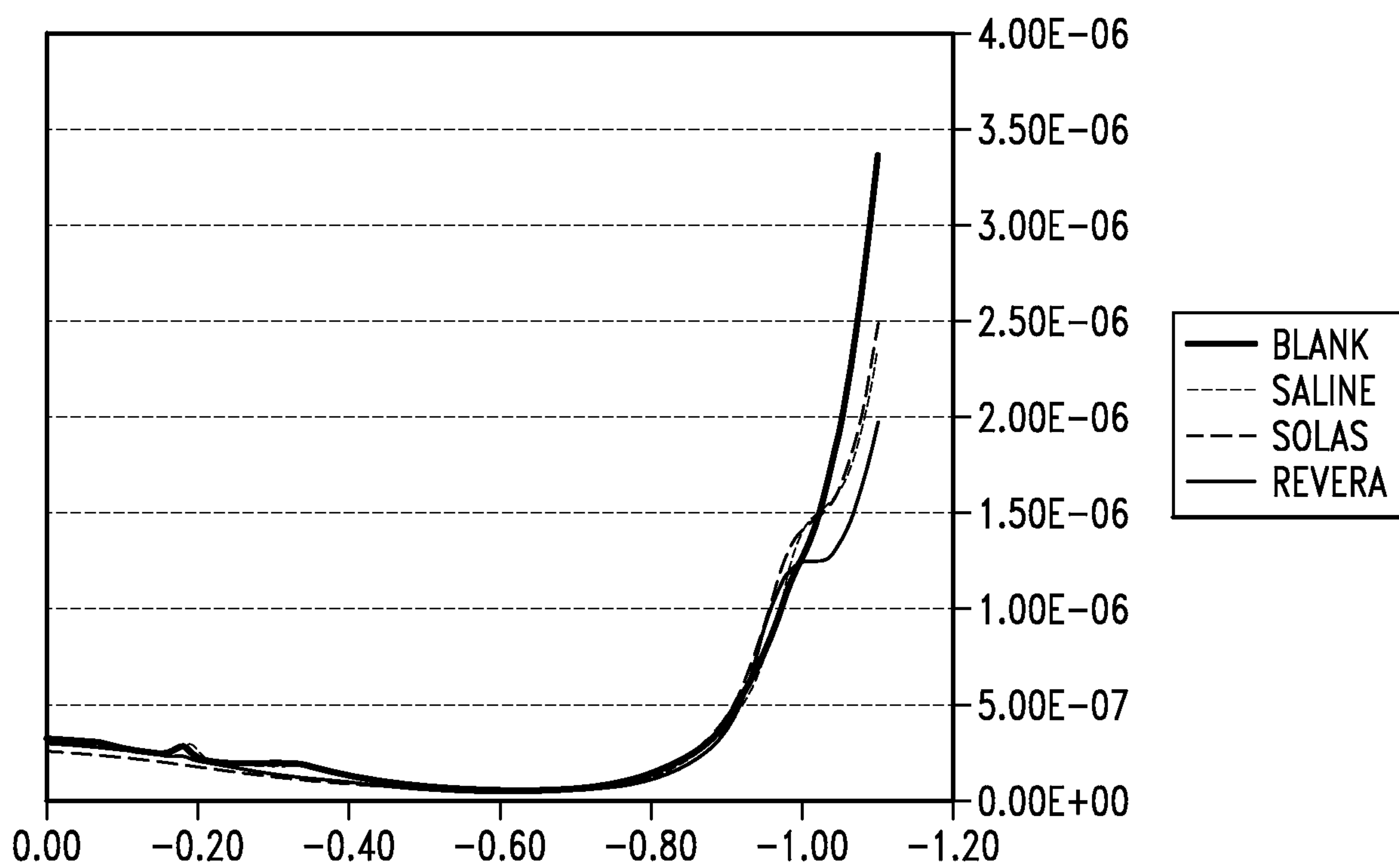
*Fig. 93**Fig. 94*

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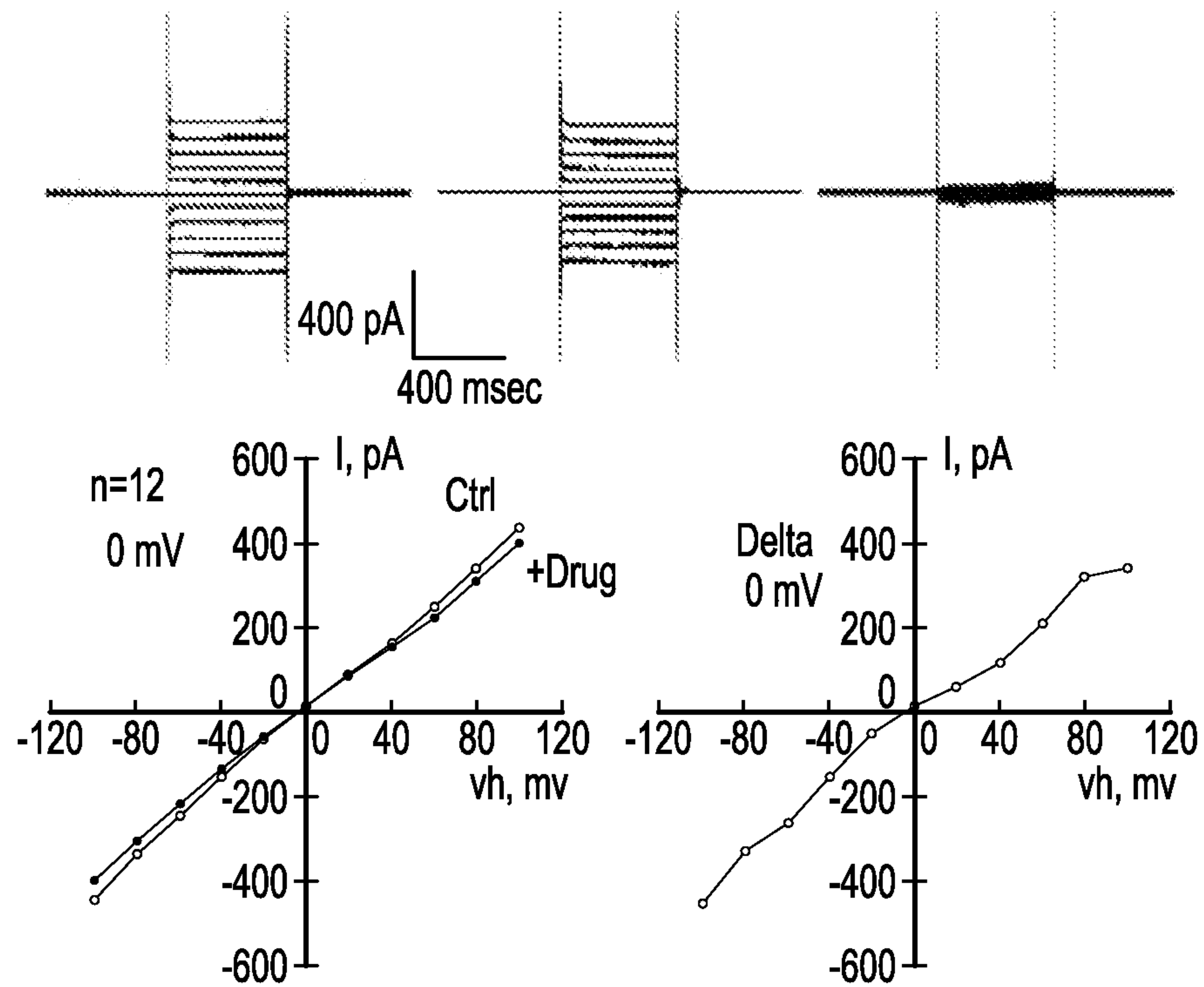
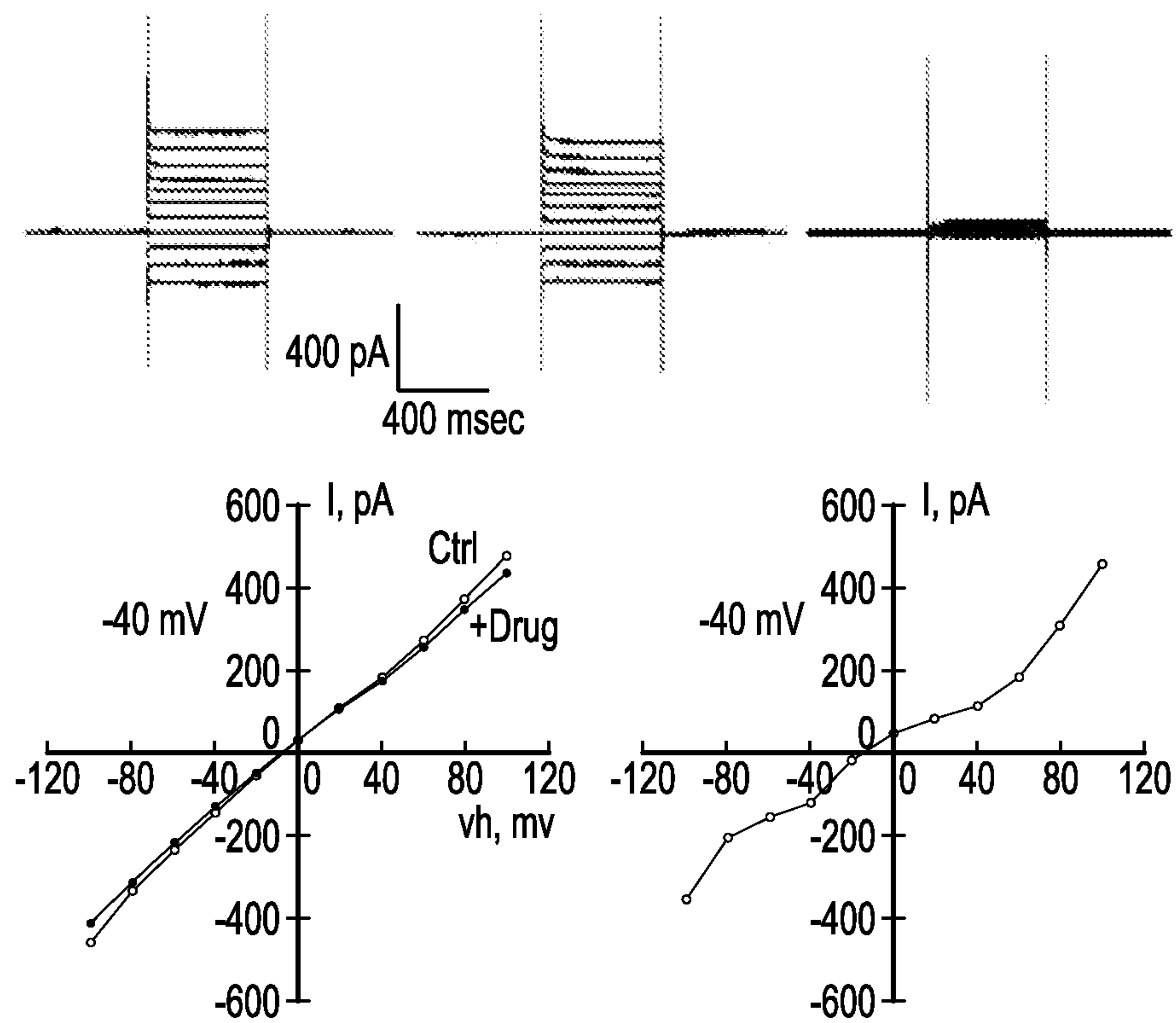
*Fig. 95**Fig. 96*

*Fig. 97A**Fig. 97B**Fig. 97C*

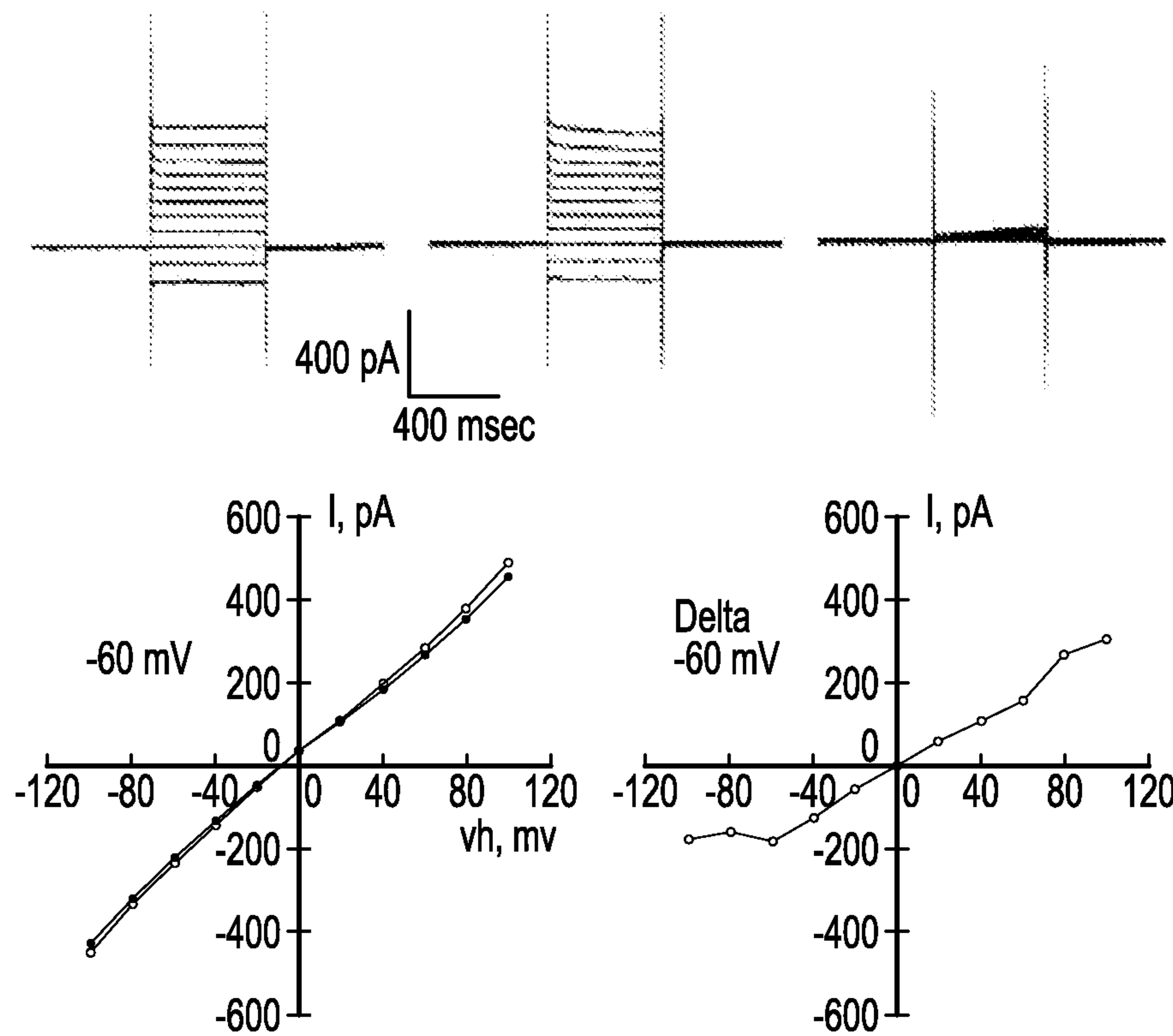
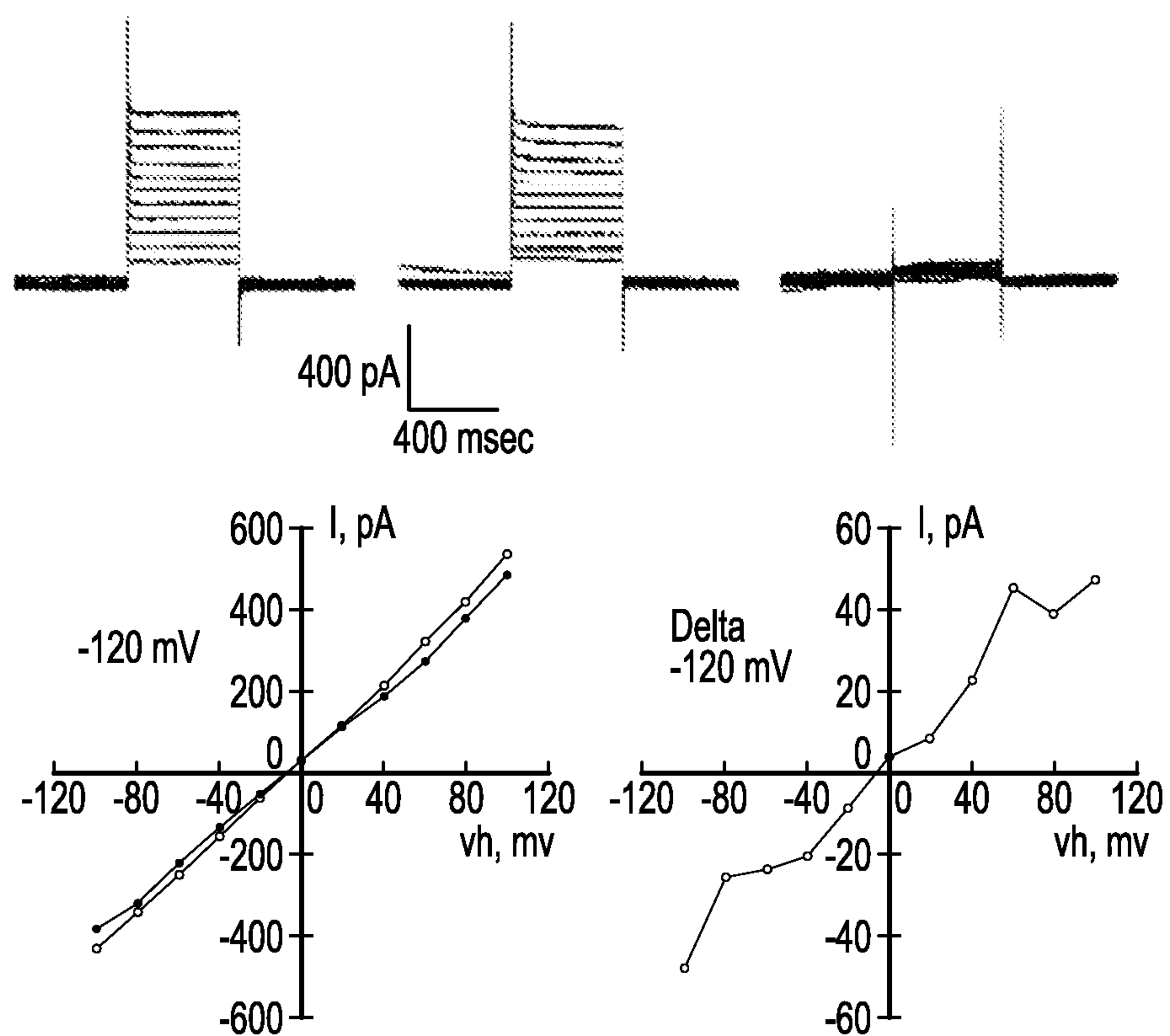
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*Fig. 98**Fig. 99*

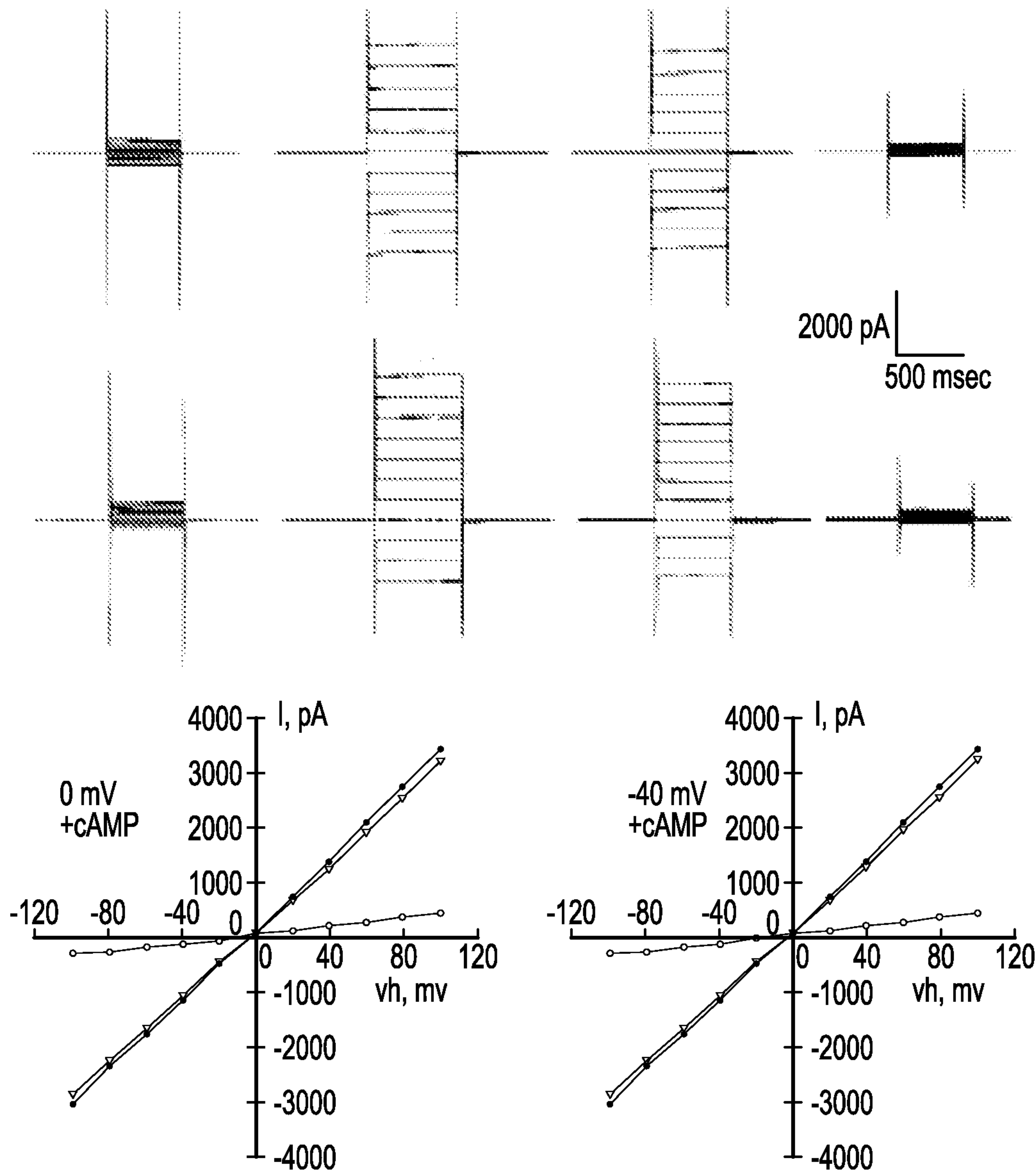
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*Fig. 100**Fig. 101*

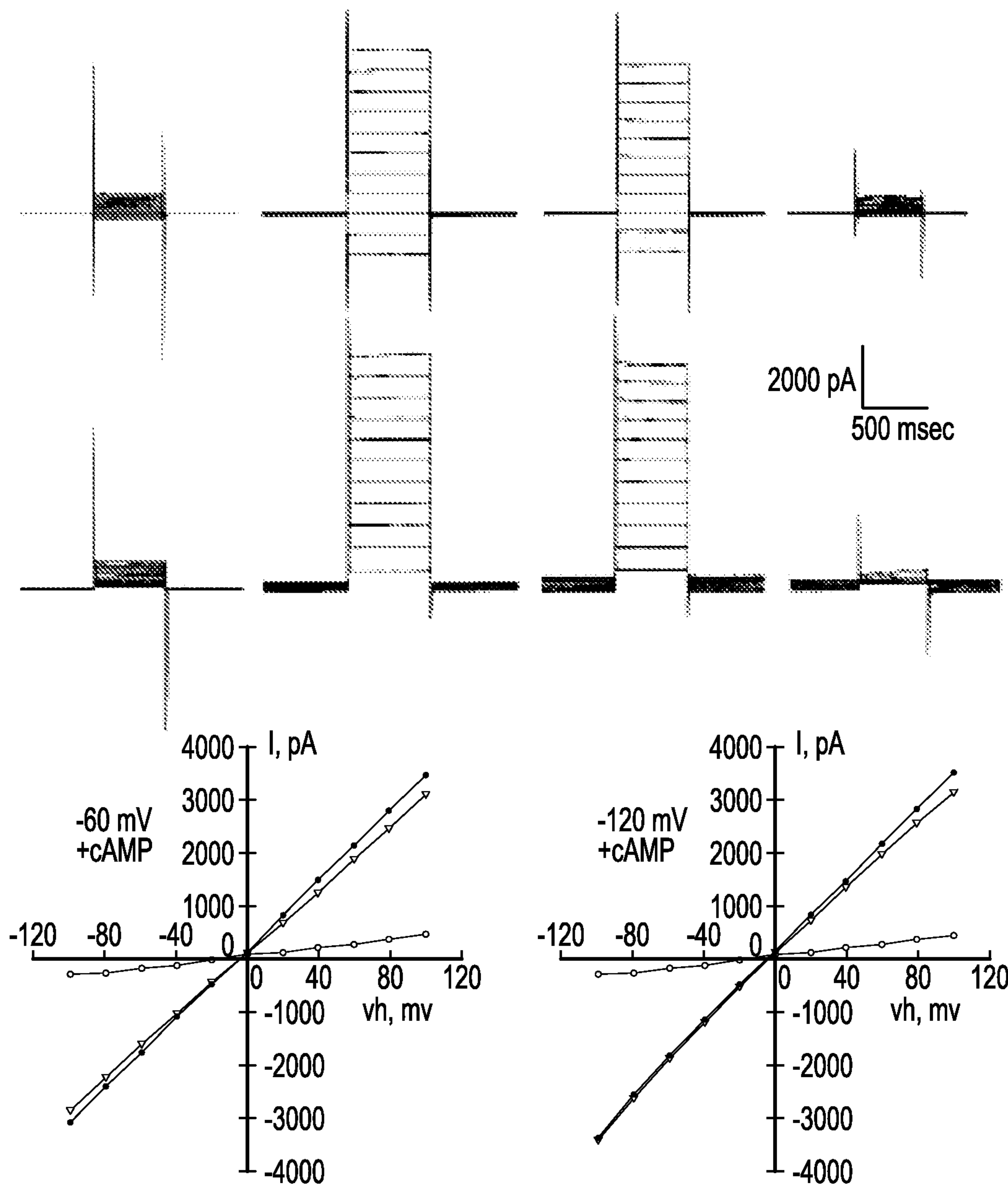
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*Fig. 102**Fig. 103*

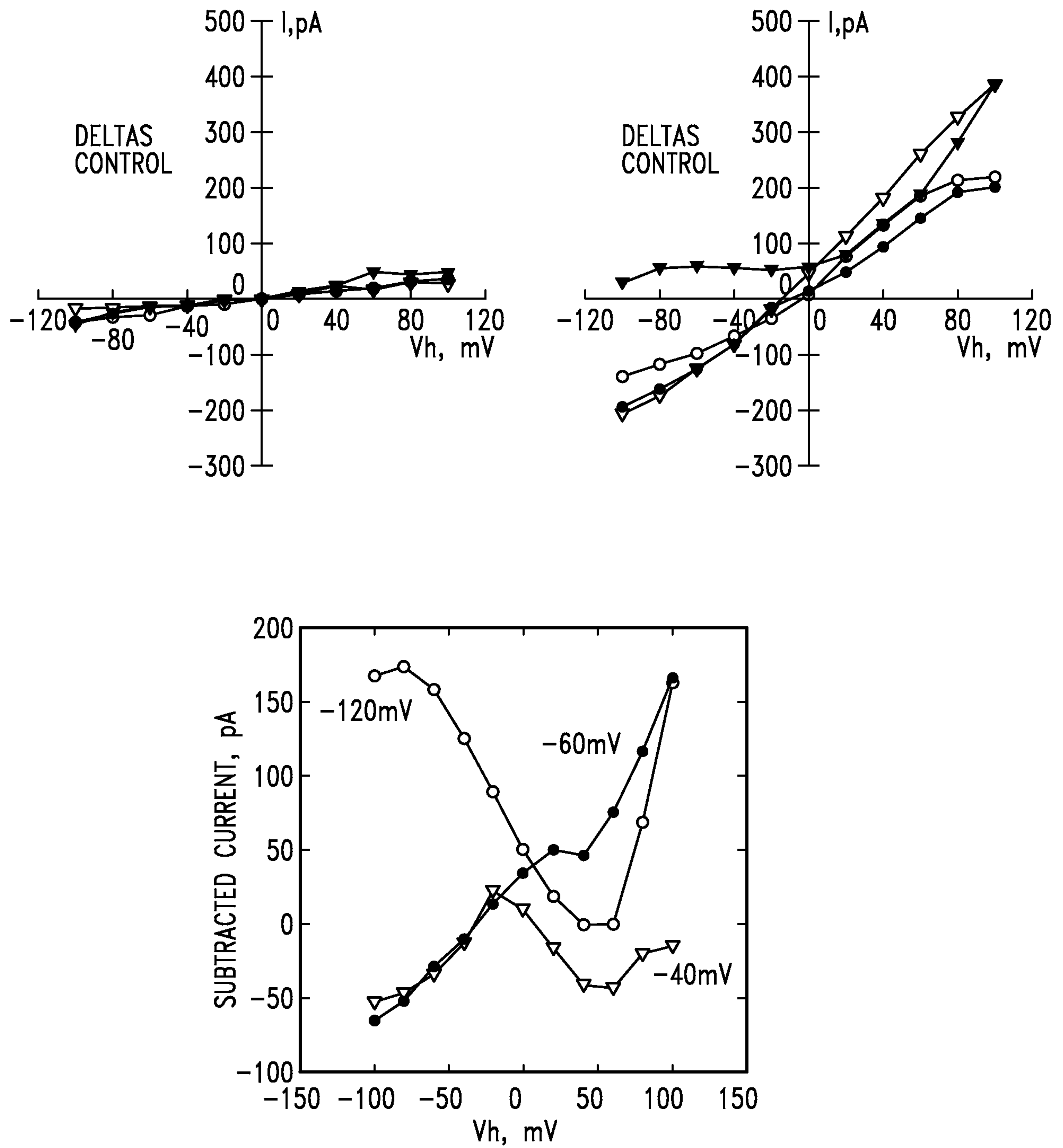
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*Fig. 104*

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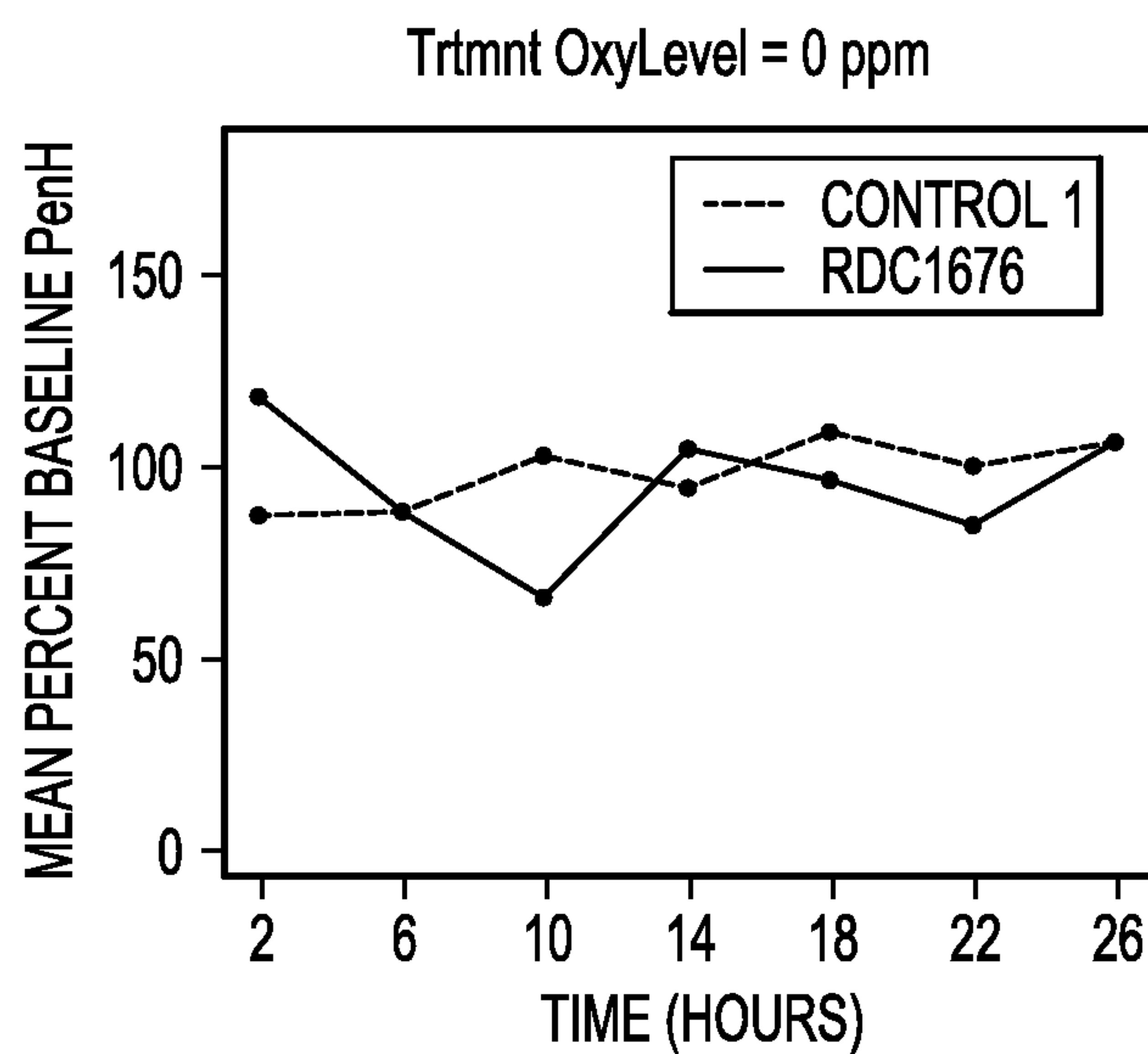
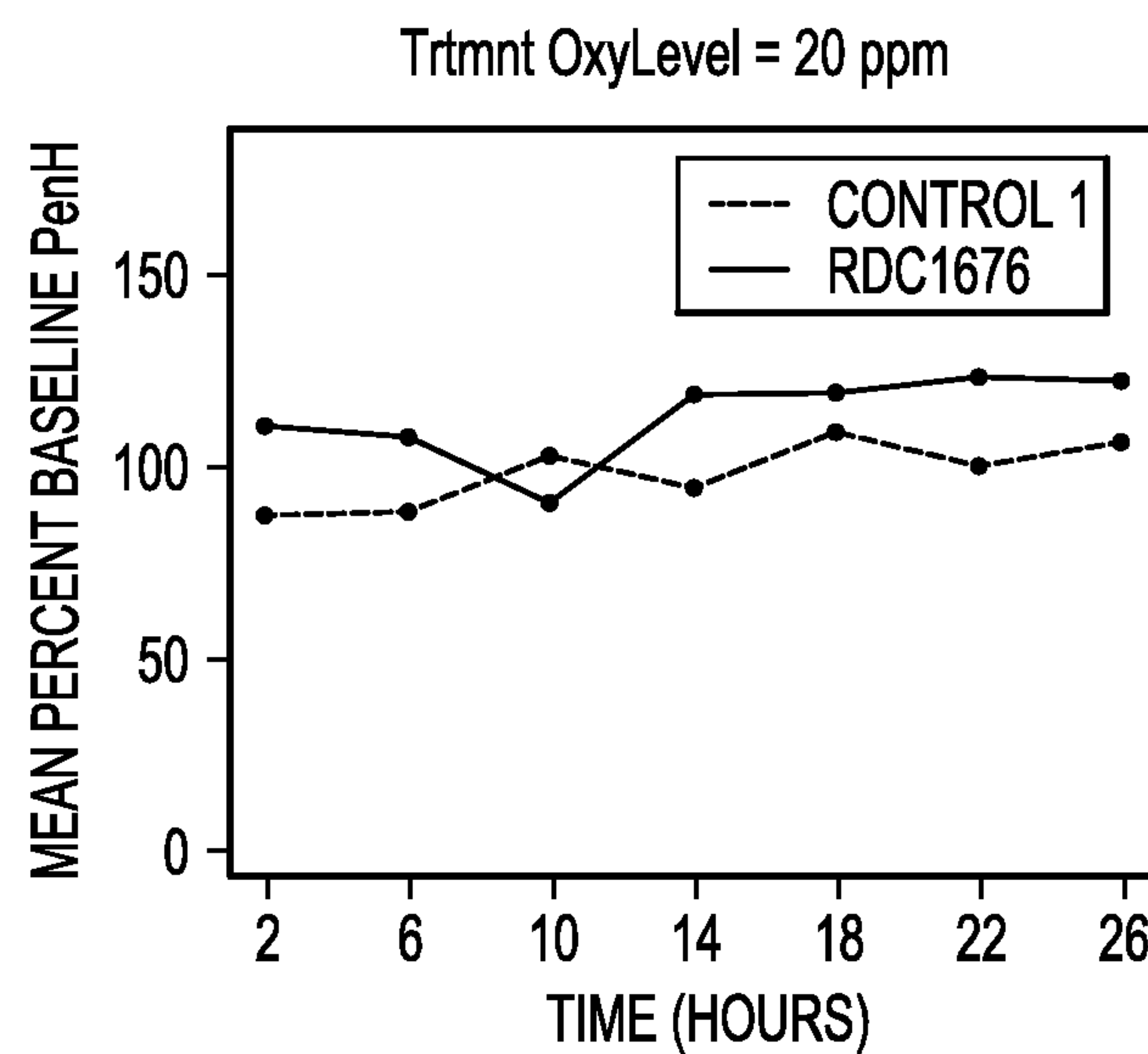
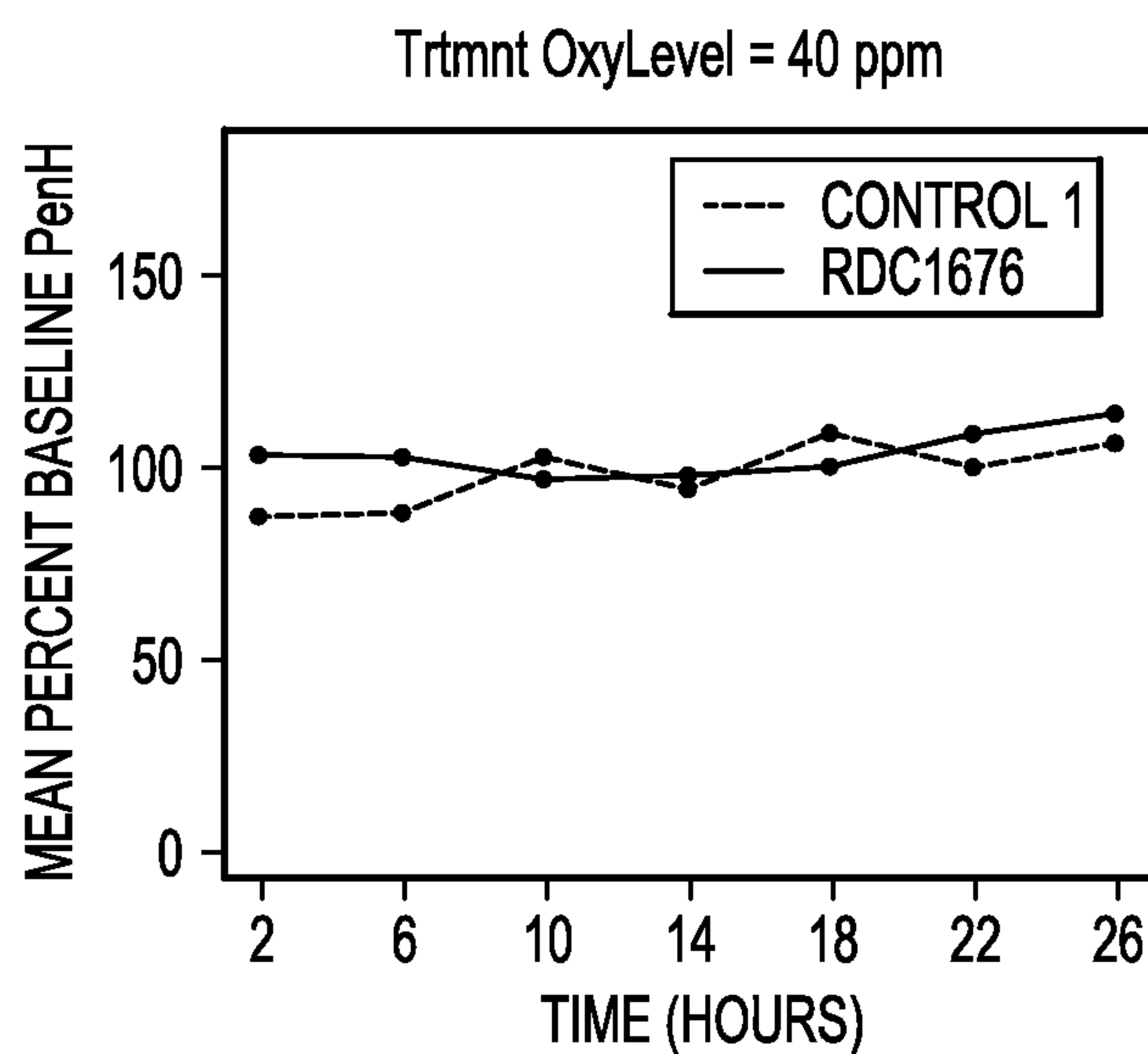
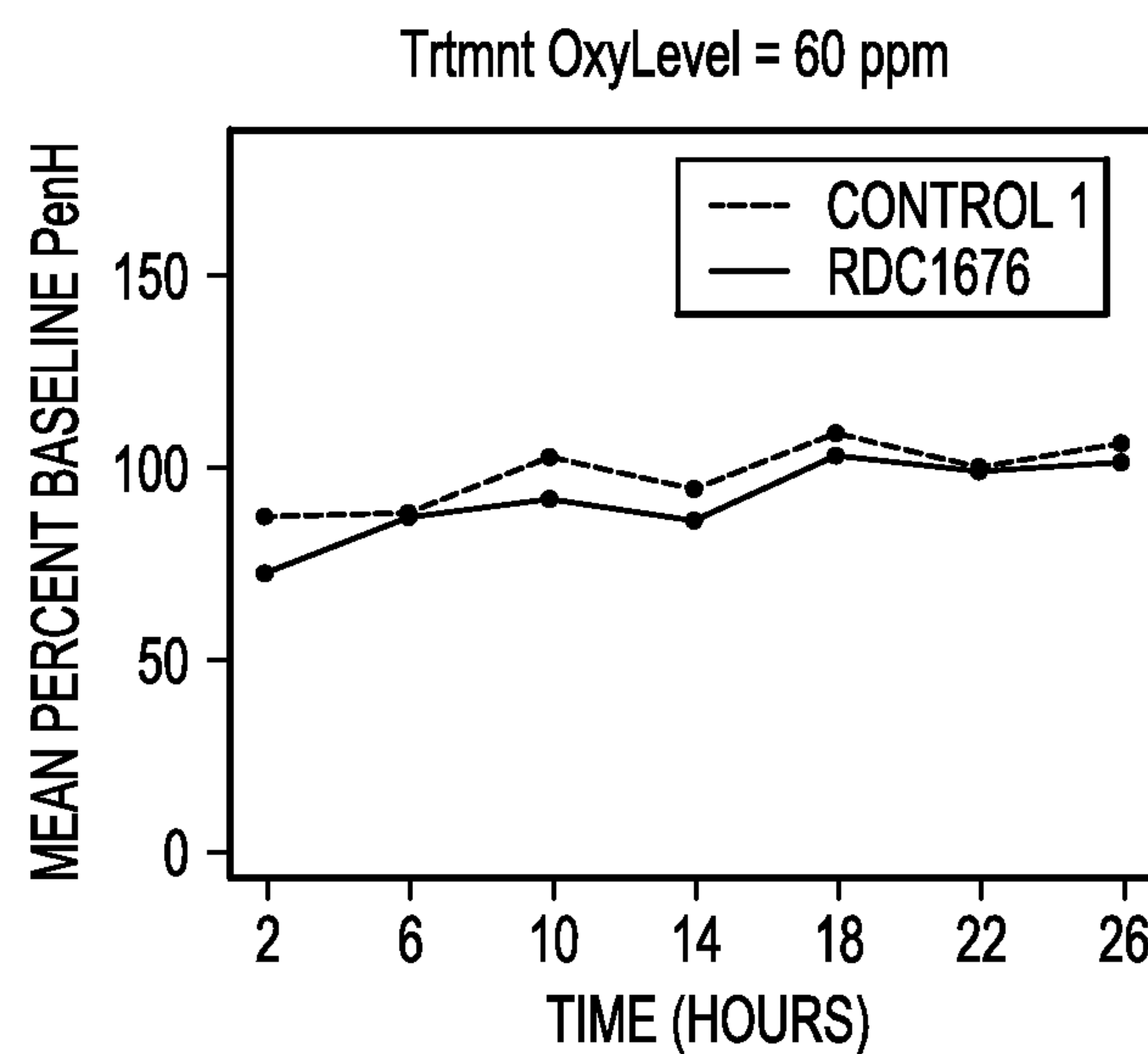
*Fig. 105*

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*Fig. 106*

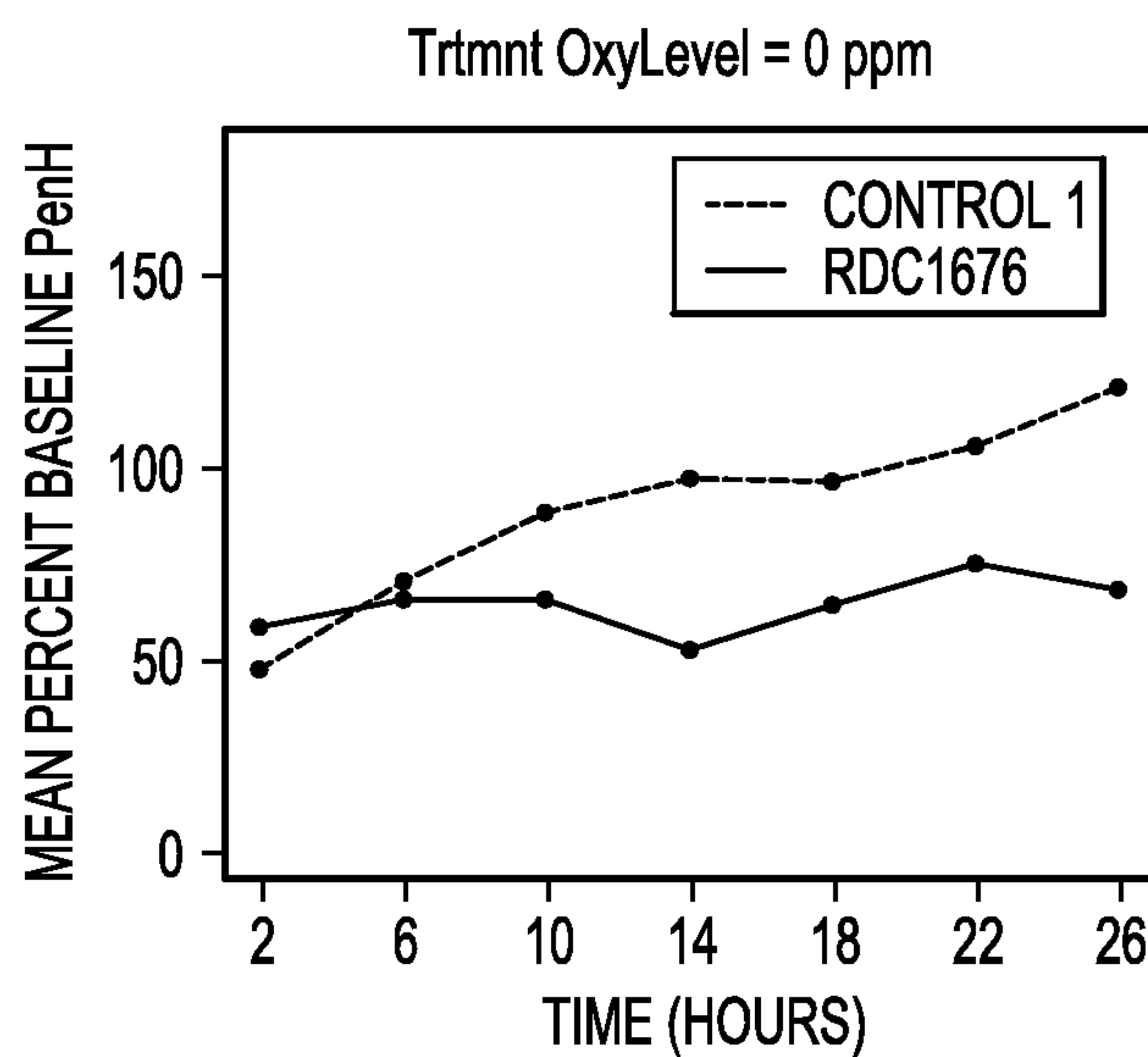
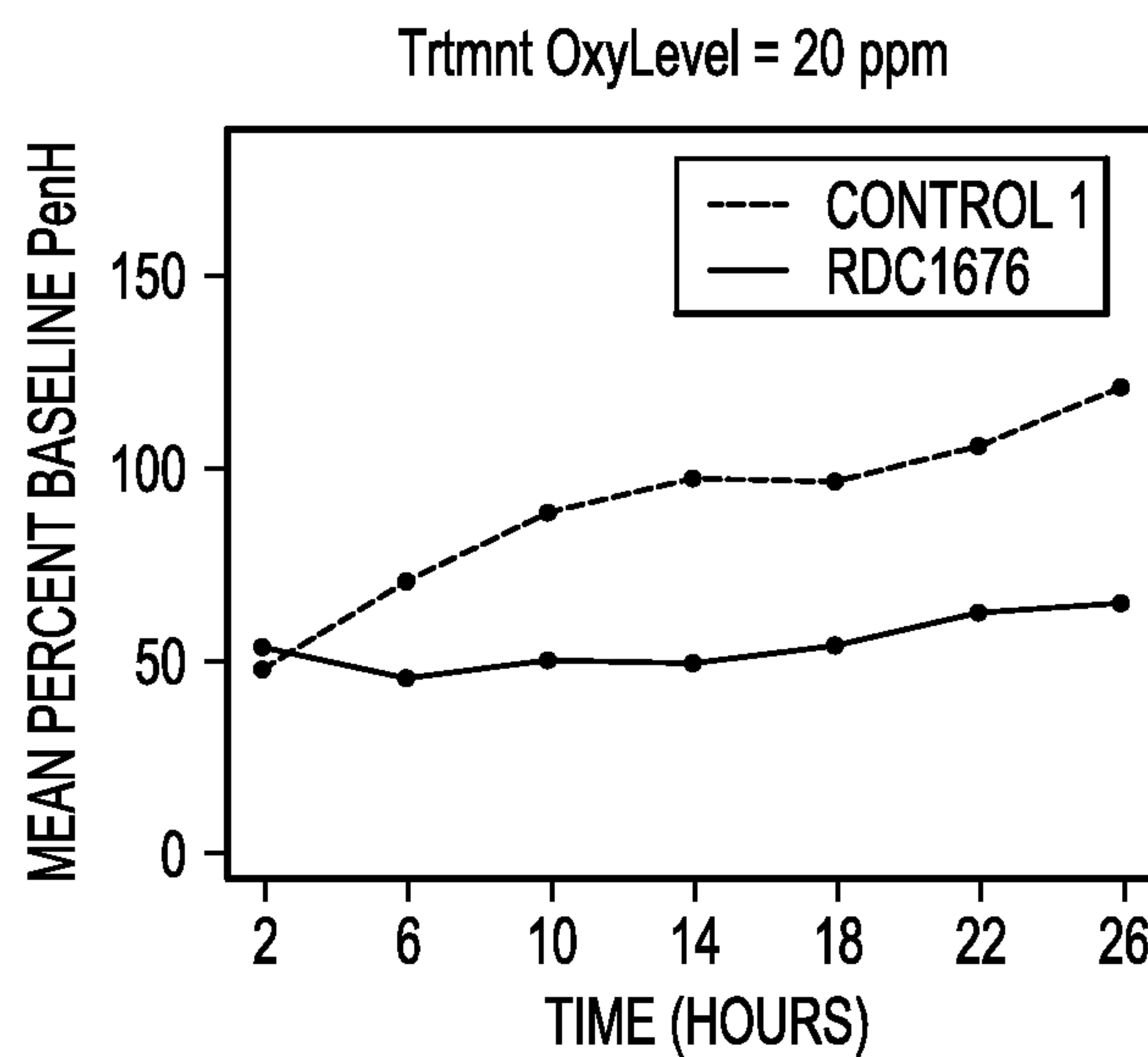
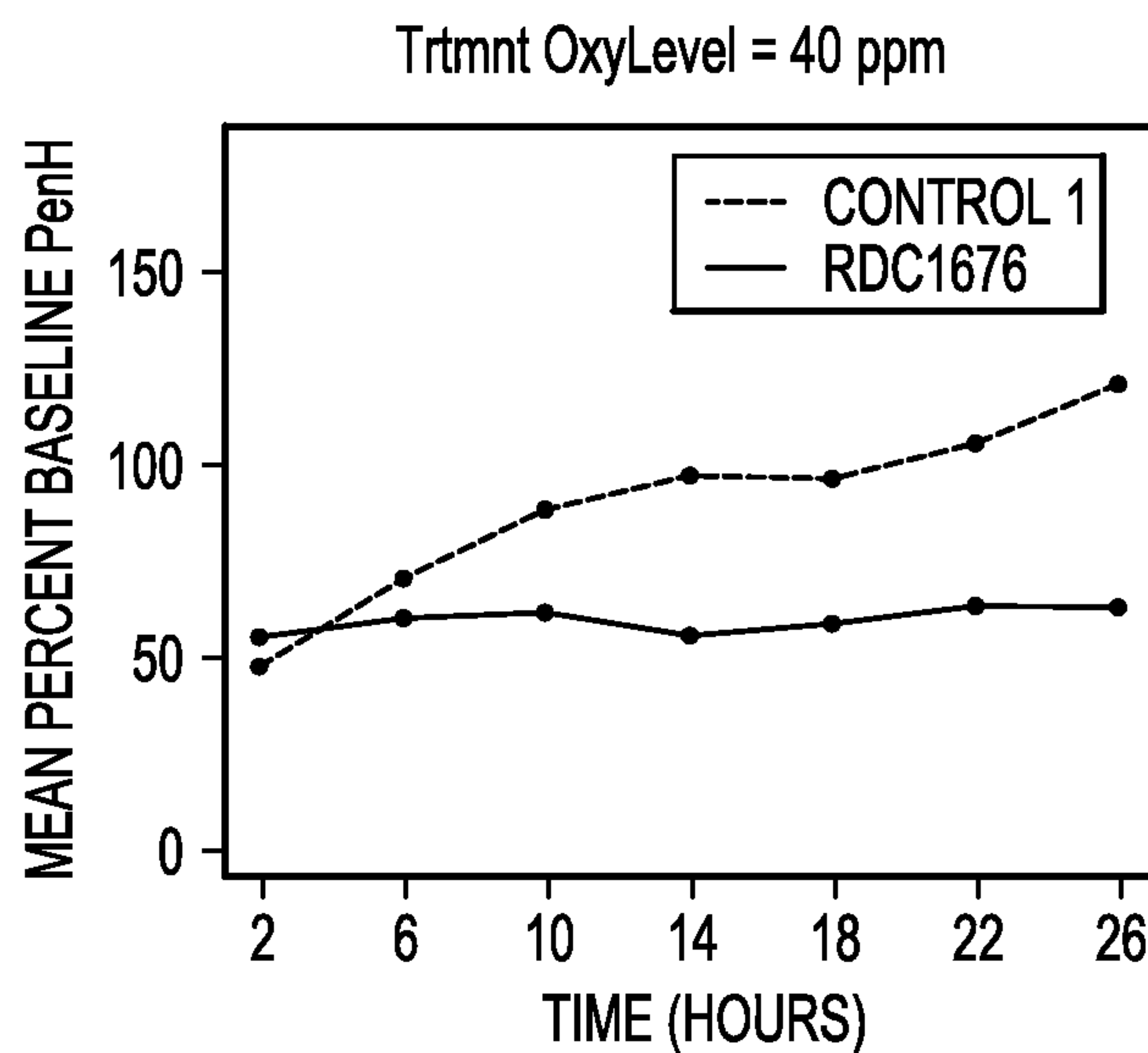
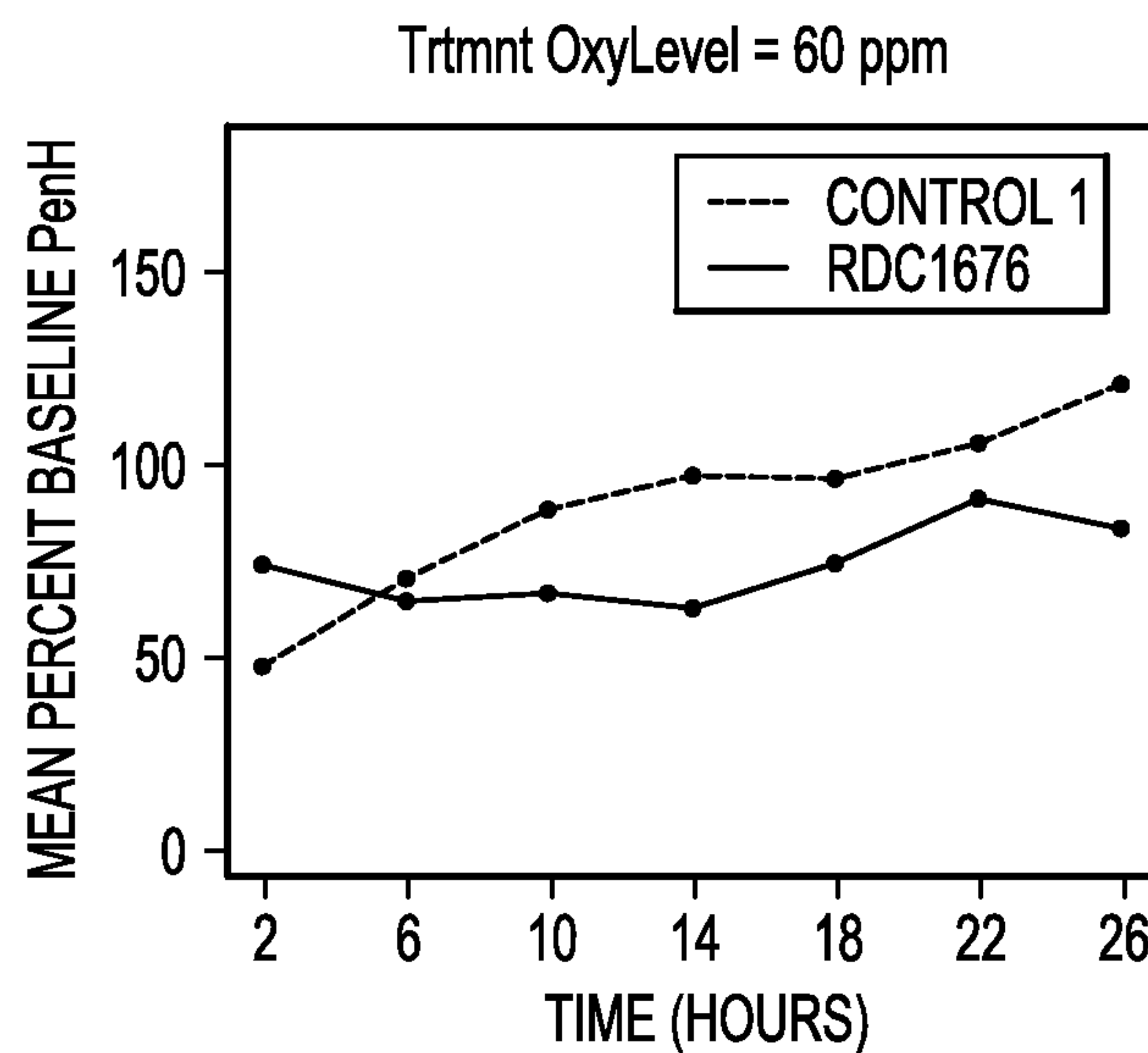
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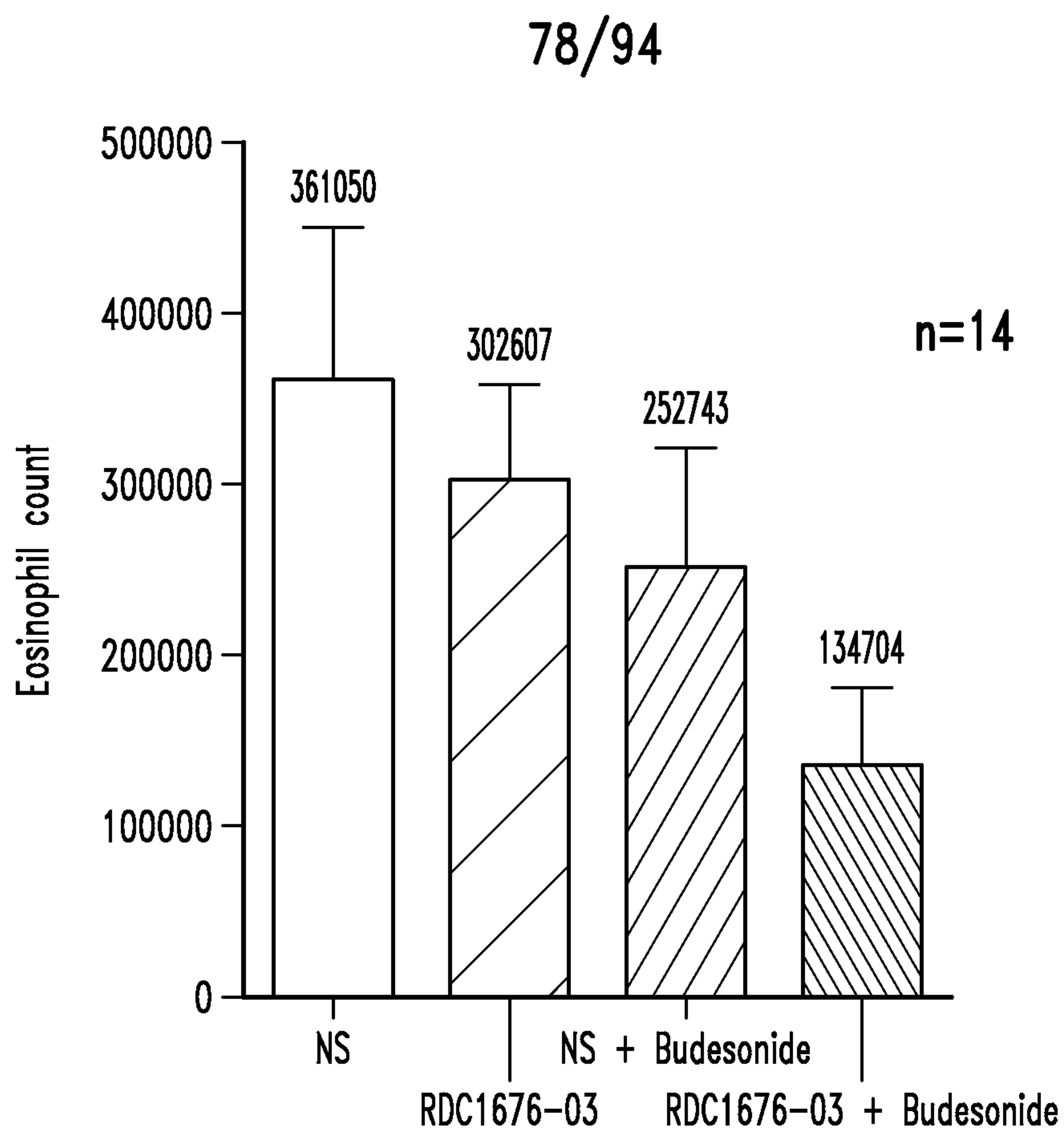
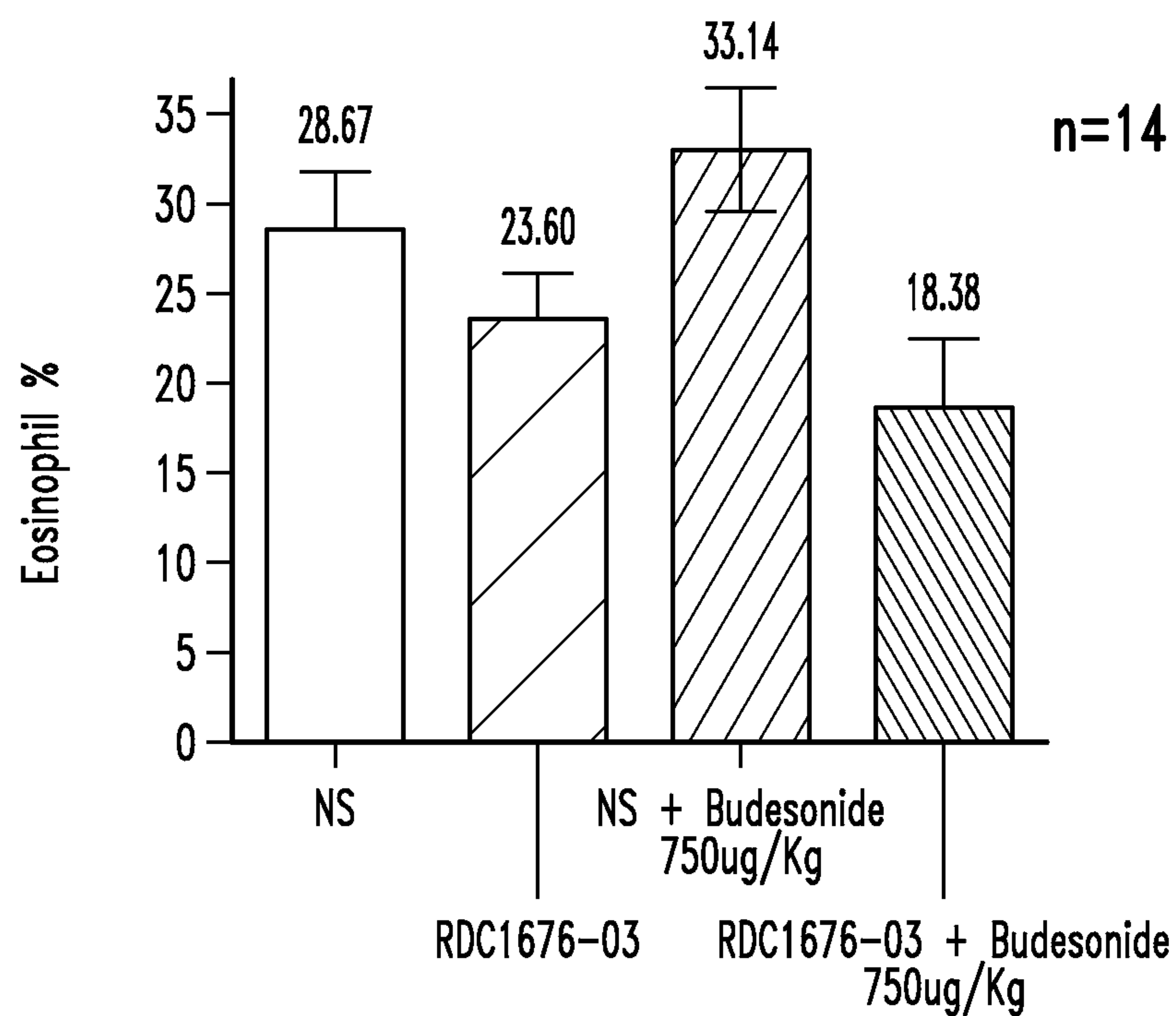
Albuterol = 0

*Fig. 107A**Fig. 107B**Fig. 107C**Fig. 107D*

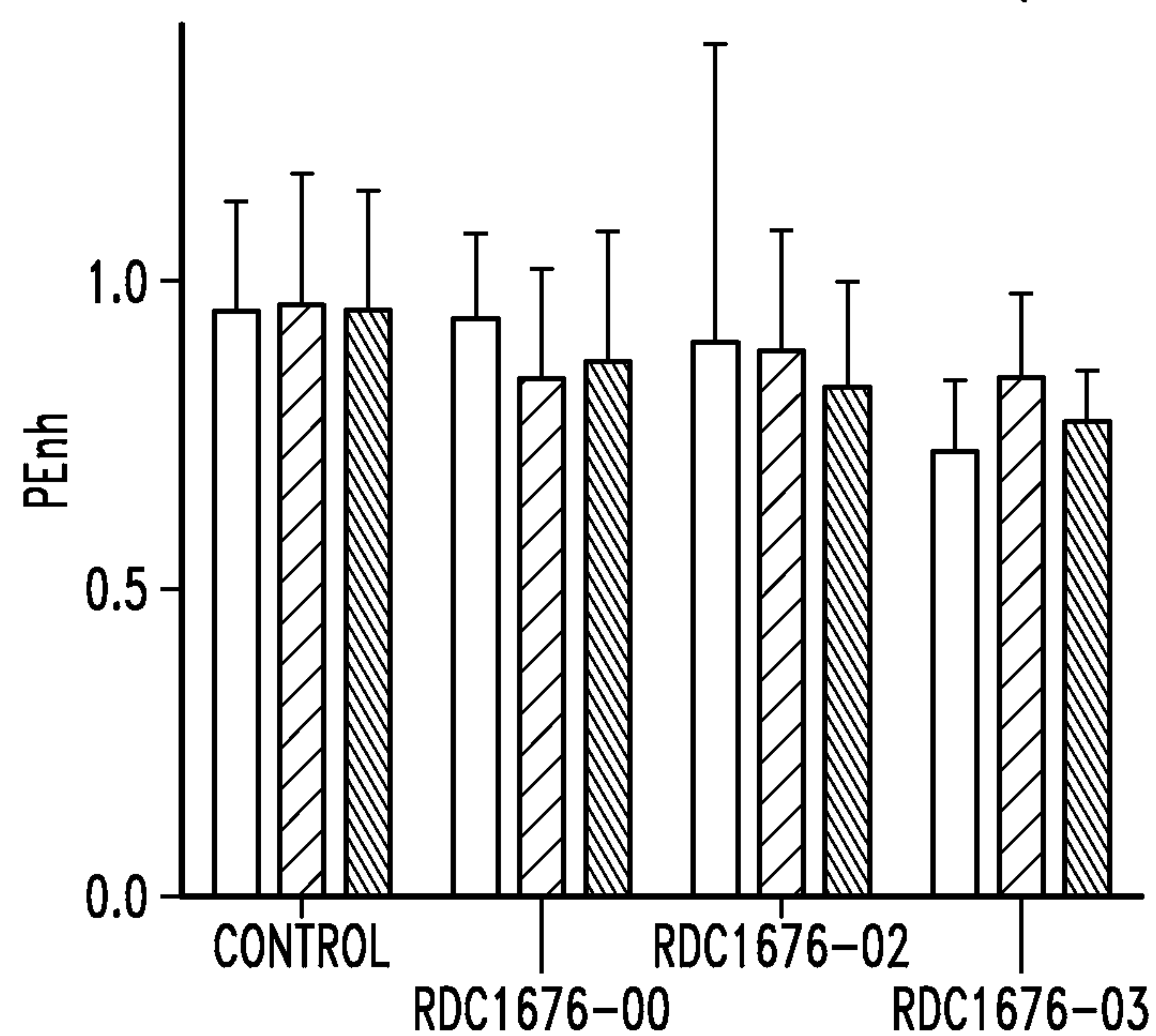
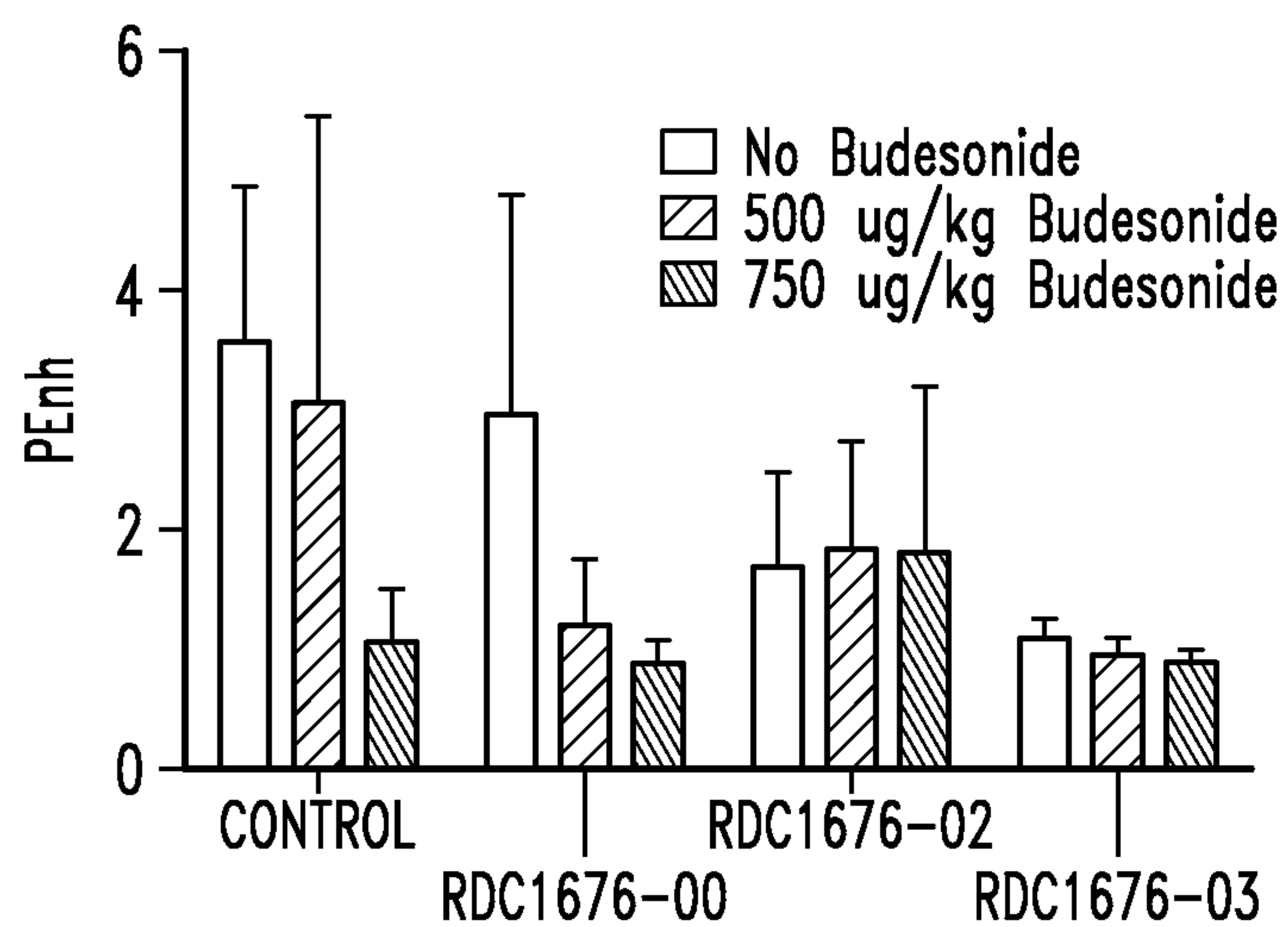
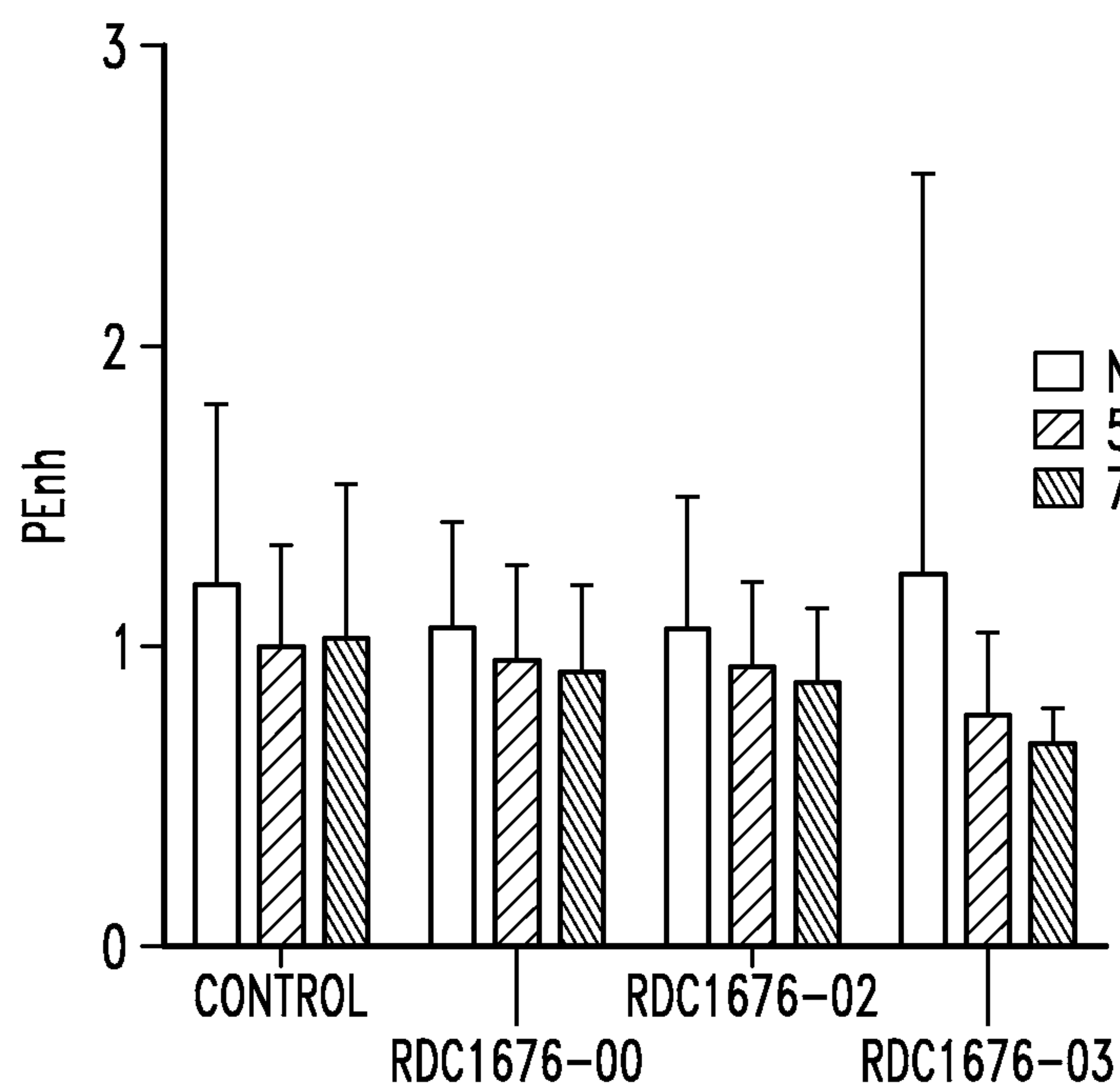
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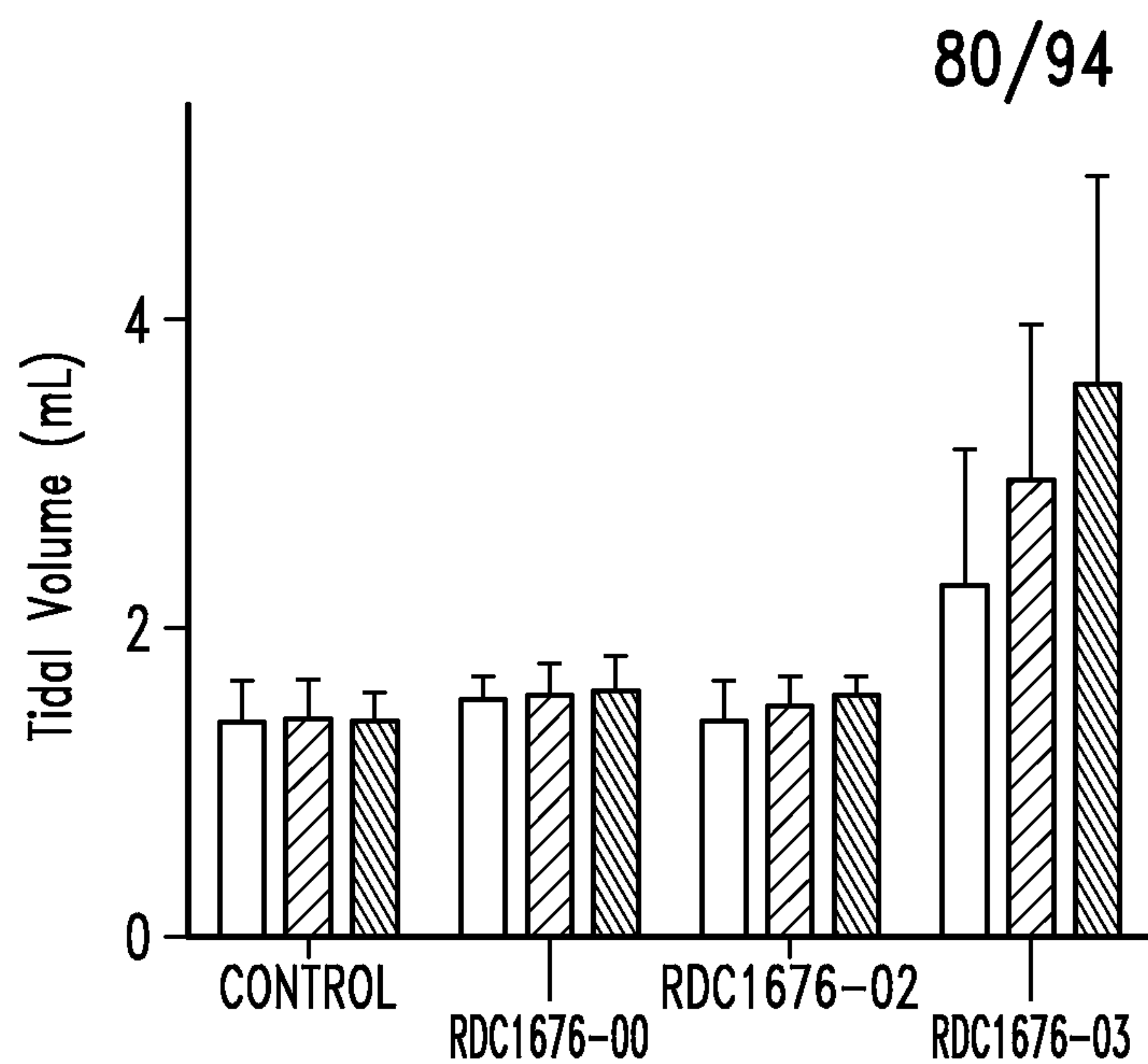
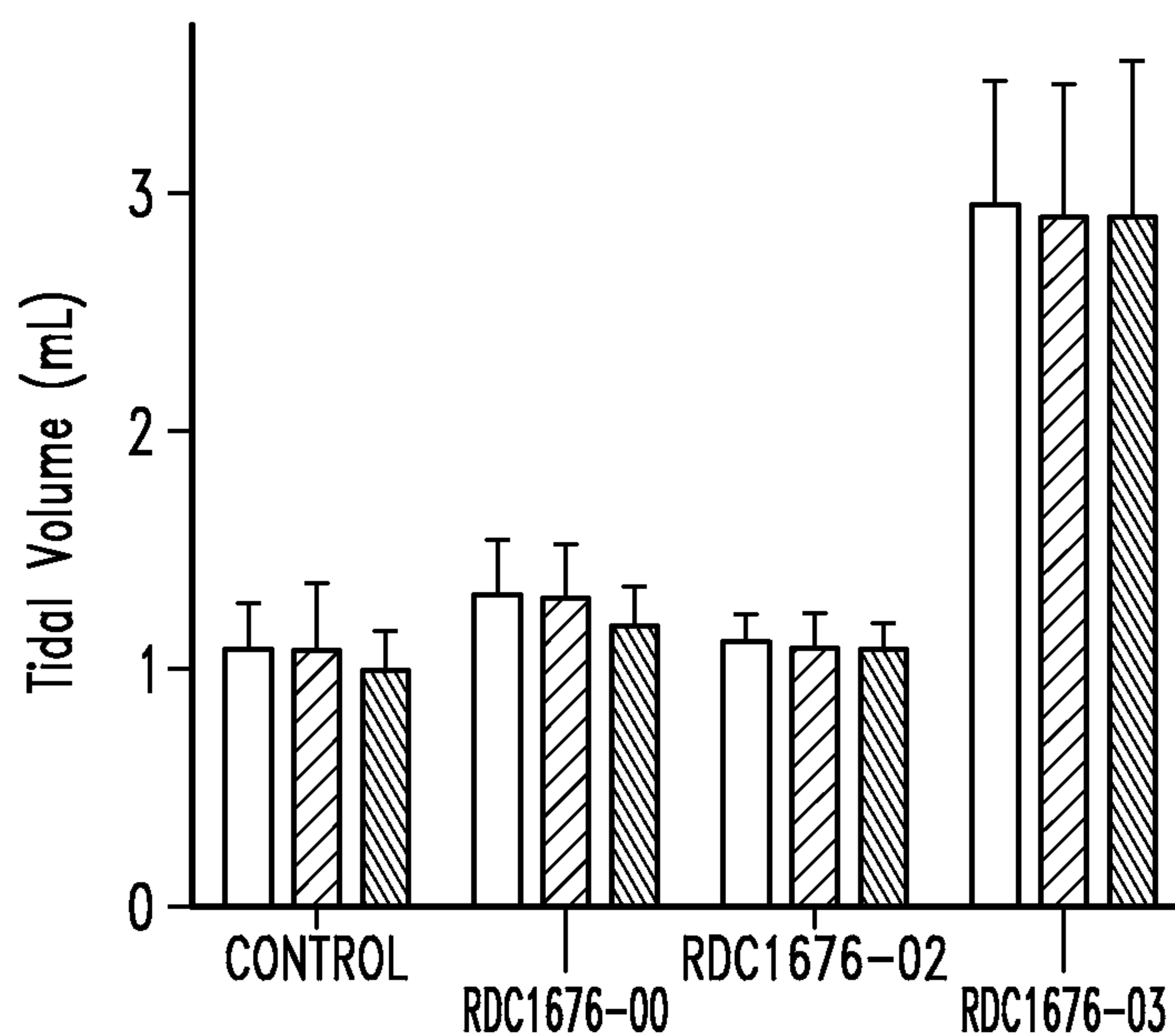
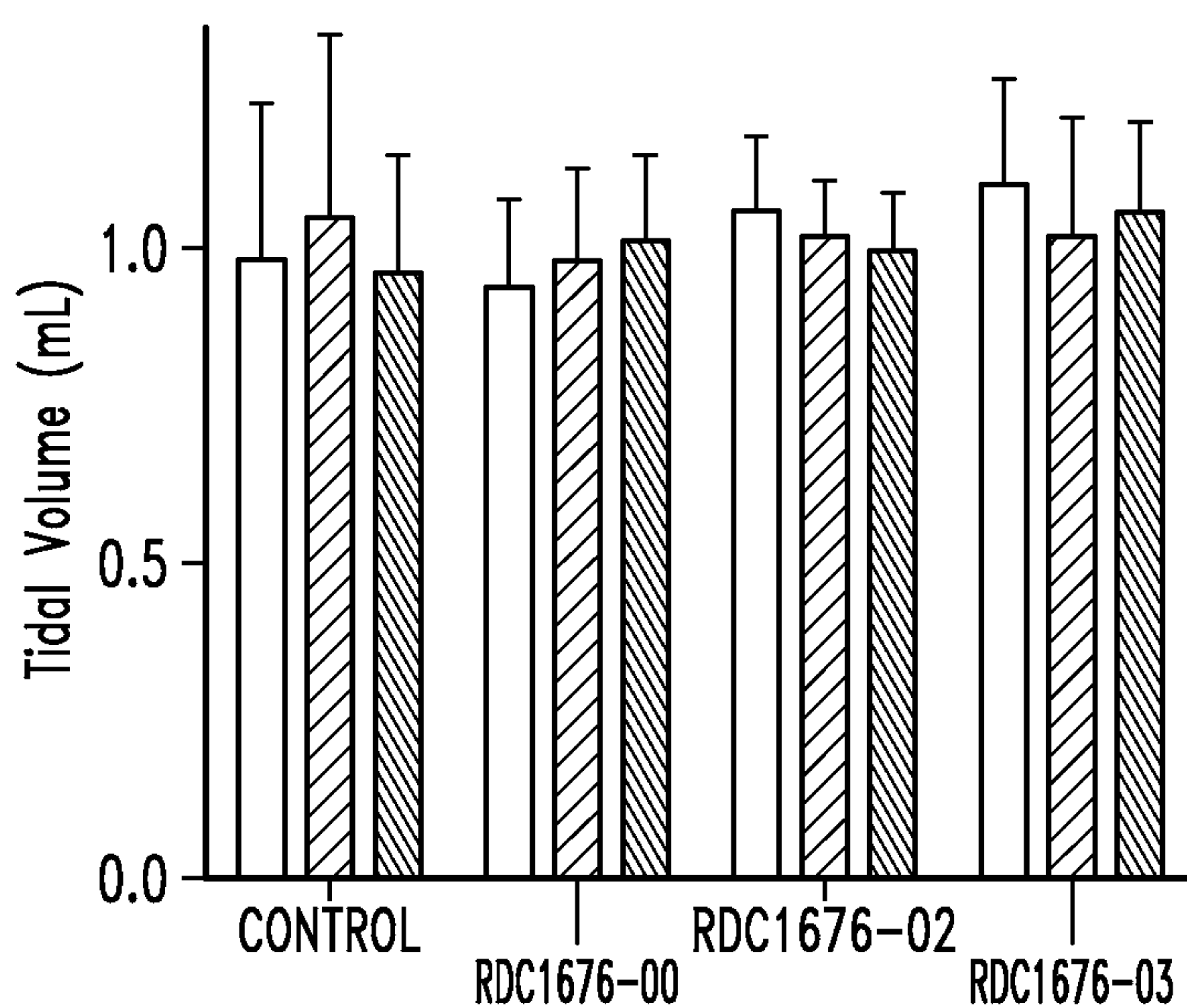
Albuterol = 25

*Fig. 108A**Fig. 108B**Fig. 108C**Fig. 108D*

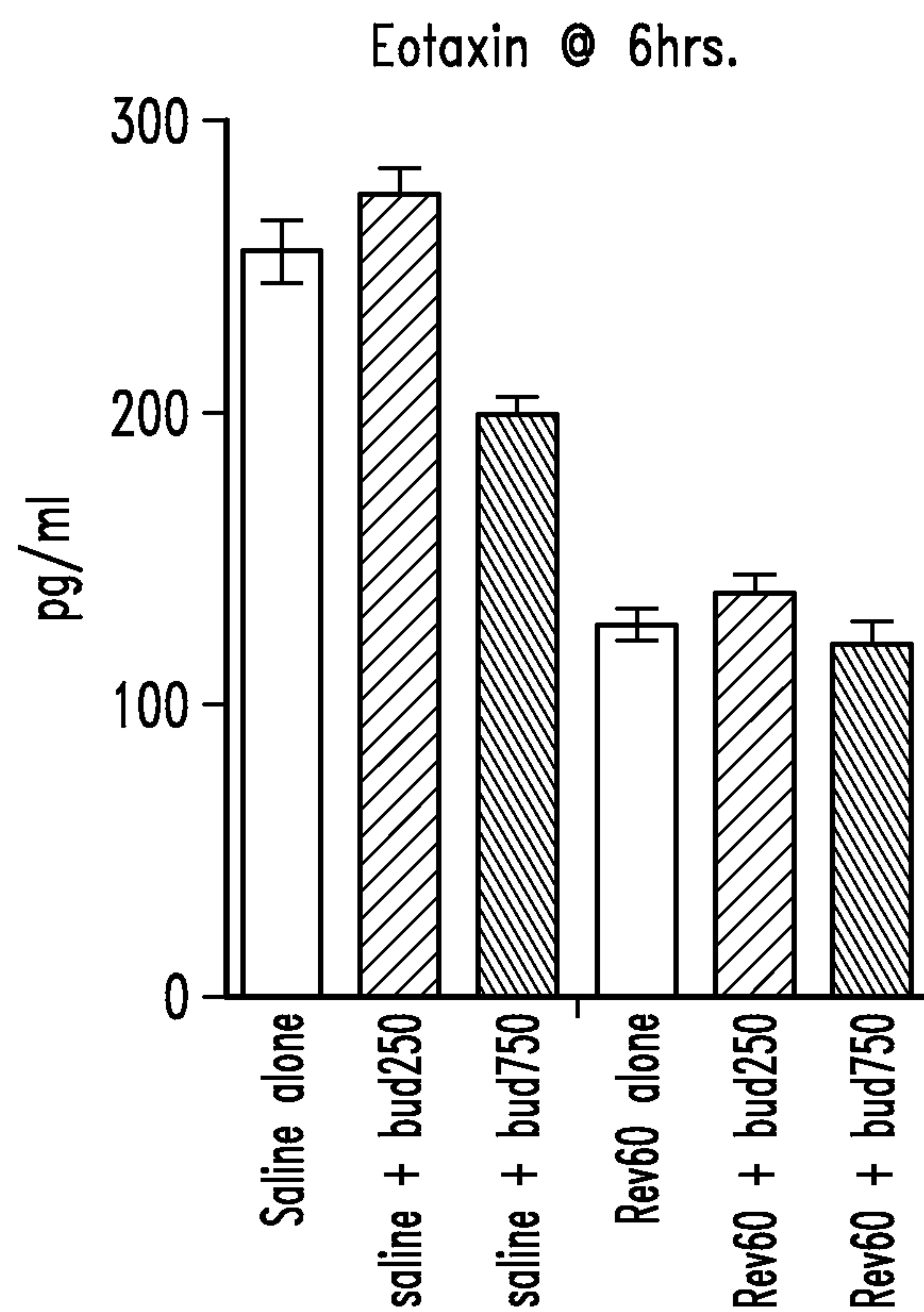
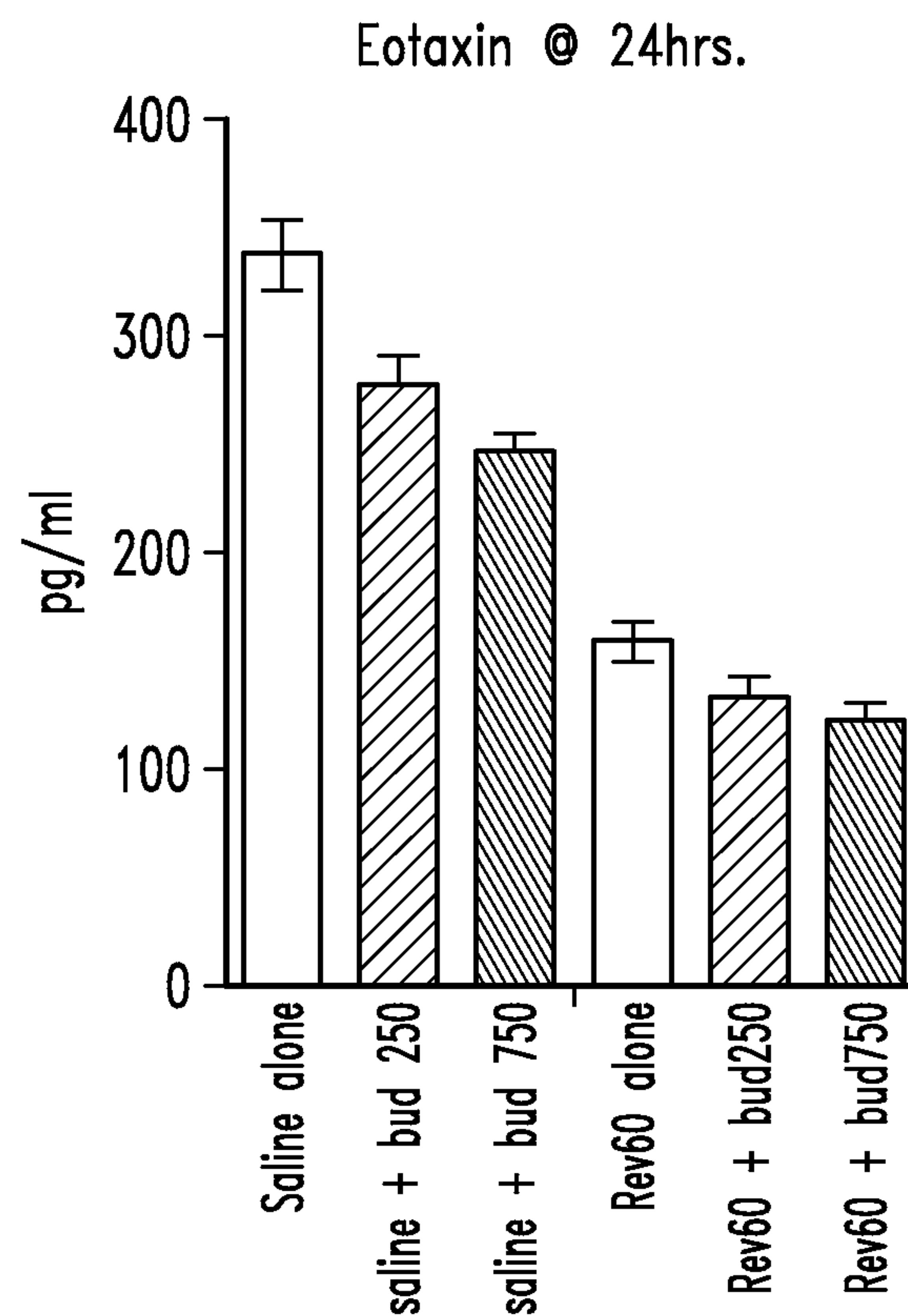
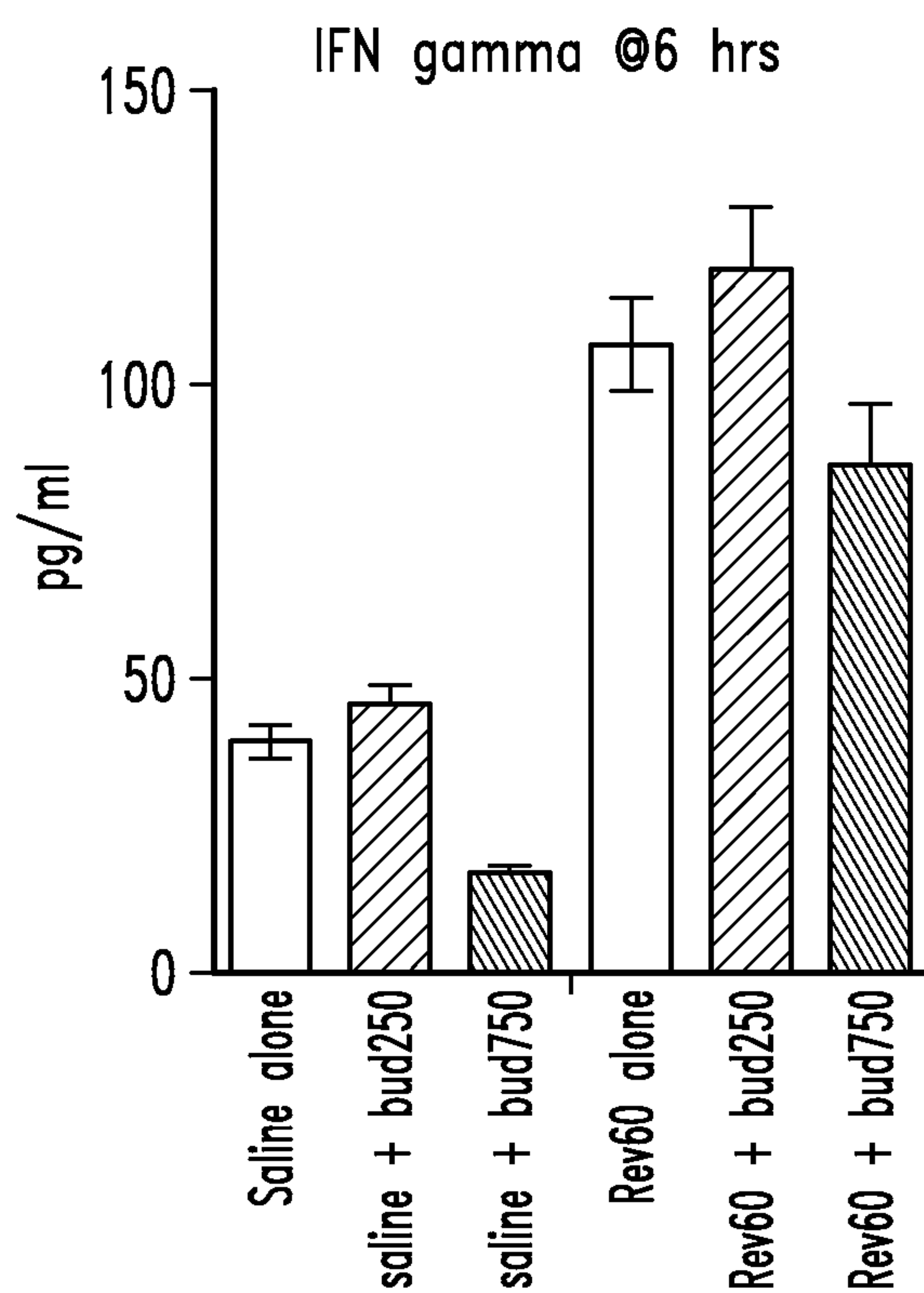
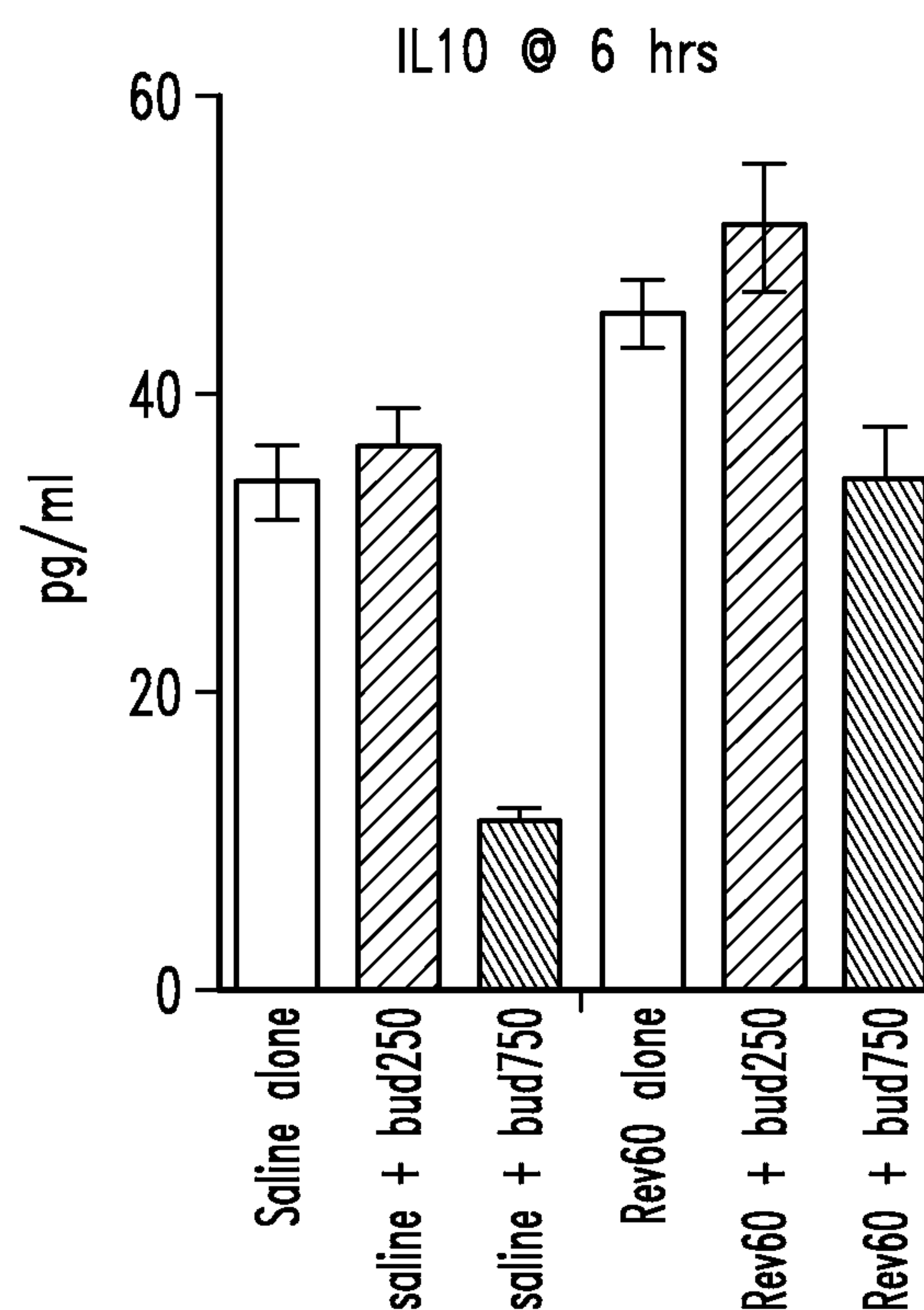
*Fig. 109**Fig. 110*

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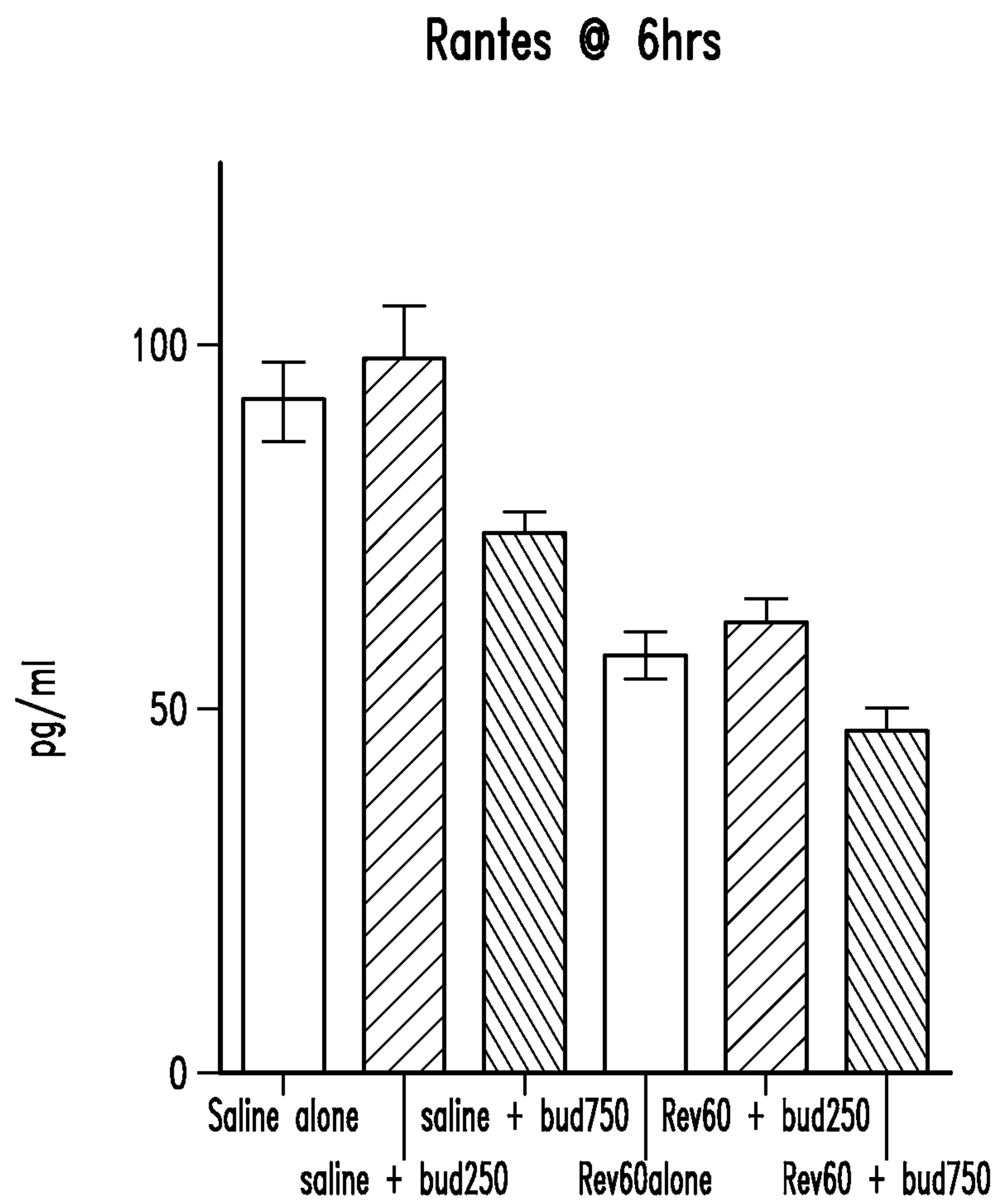
*Fig. 111A**Fig. 111B**Fig. 111C*

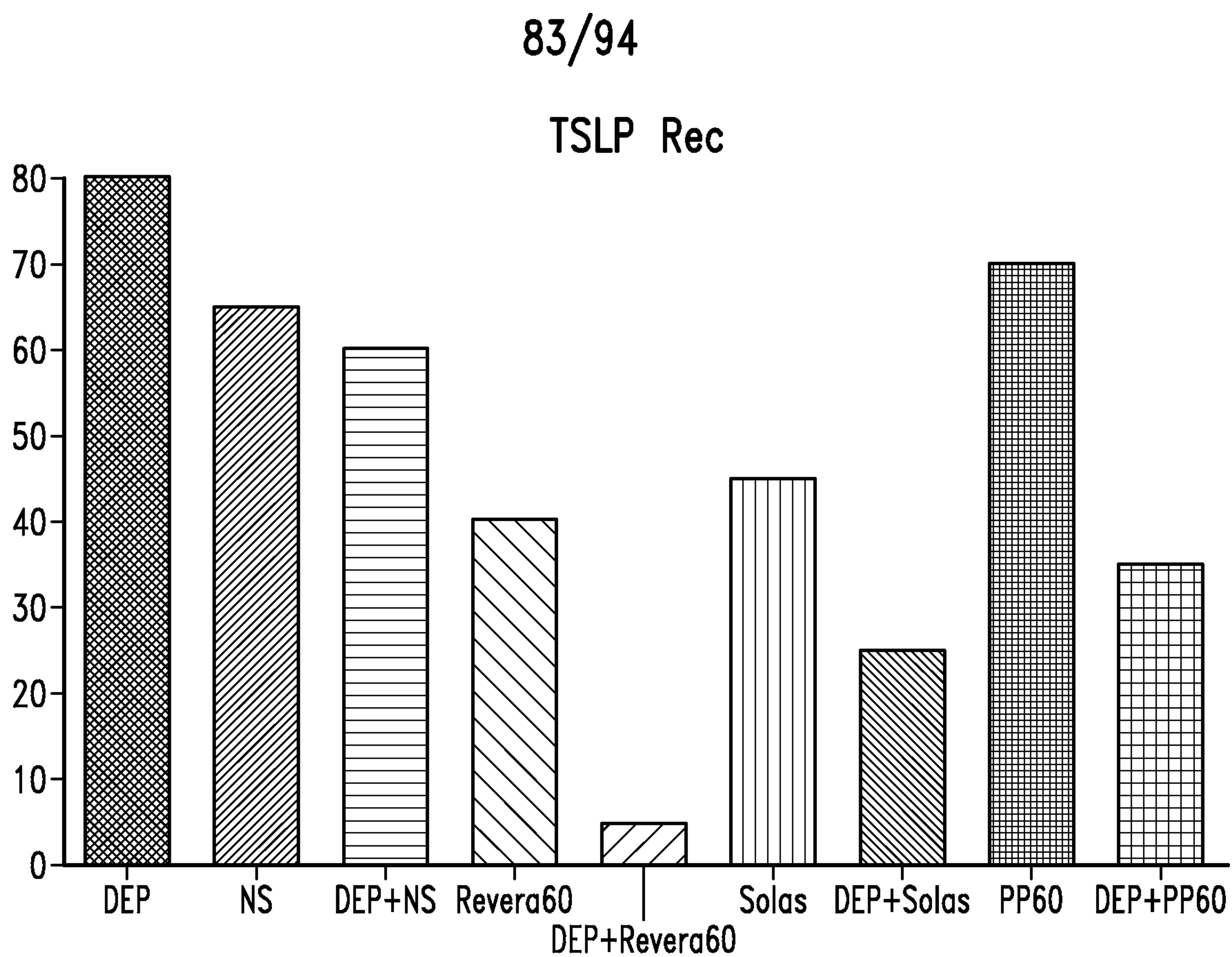
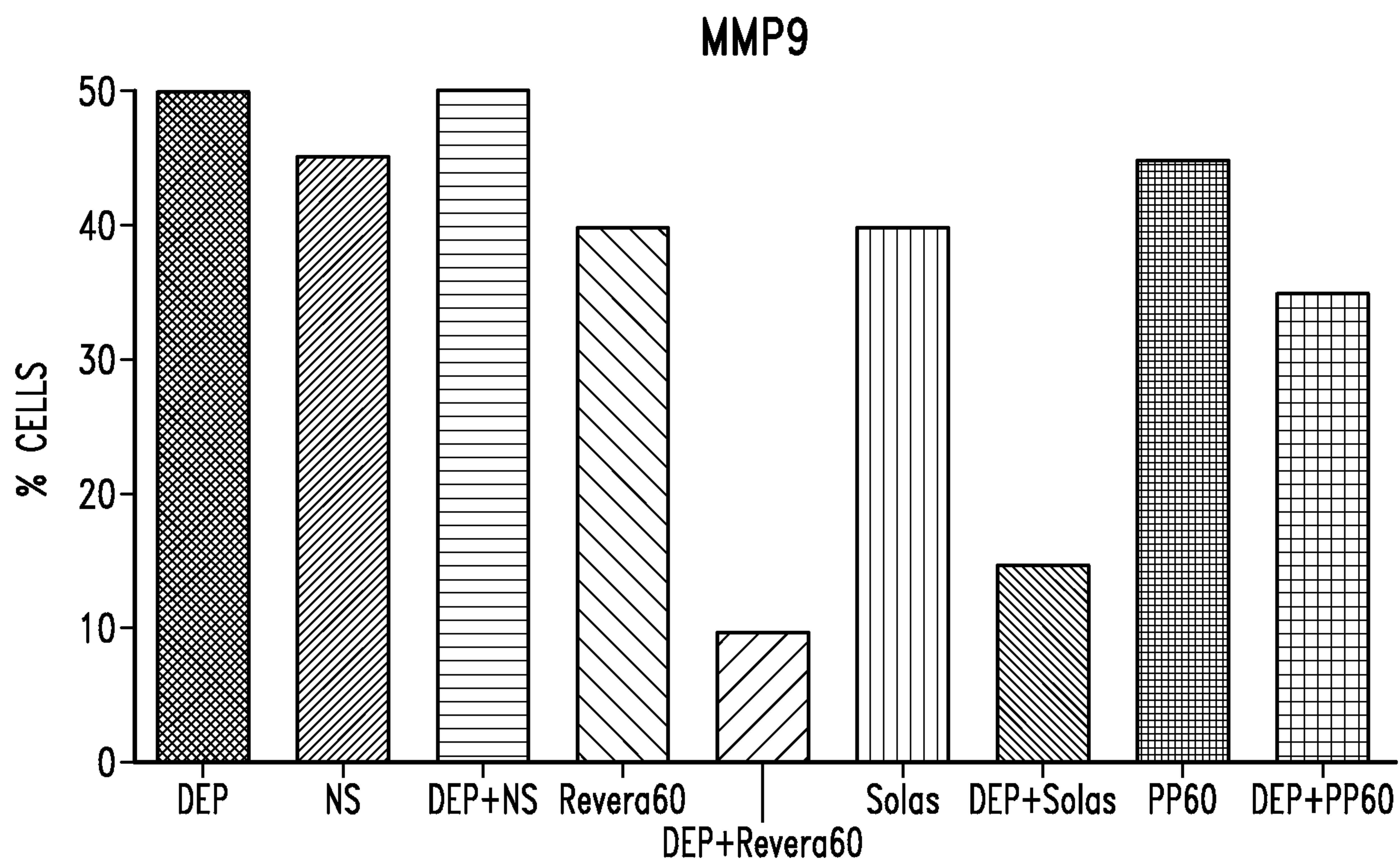
*Fig. 112A**Fig. 112B**Fig. 112C*

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*Fig. 113A**Fig. 113B**Fig. 113C**Fig. 113D*

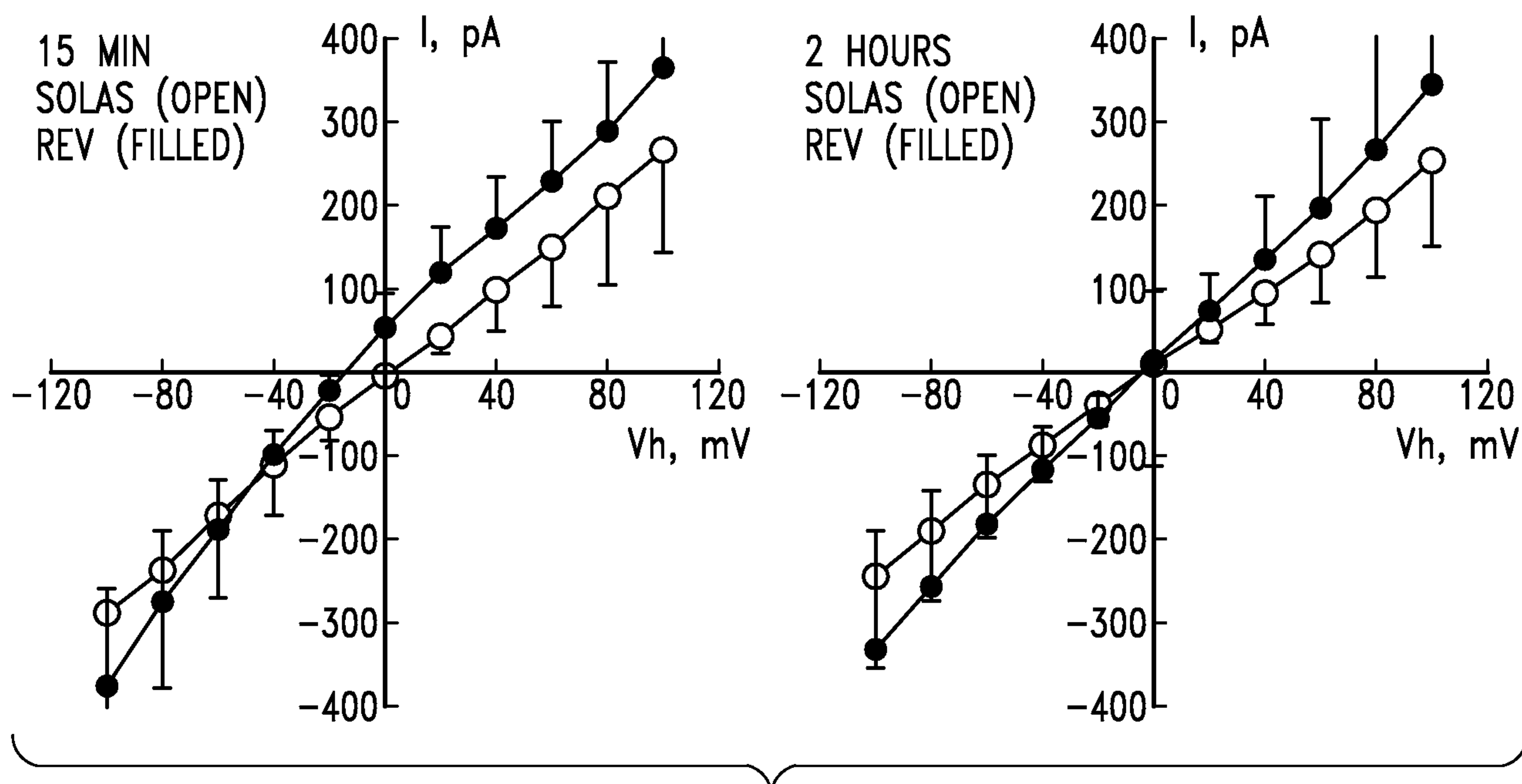
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*Fig. 114*

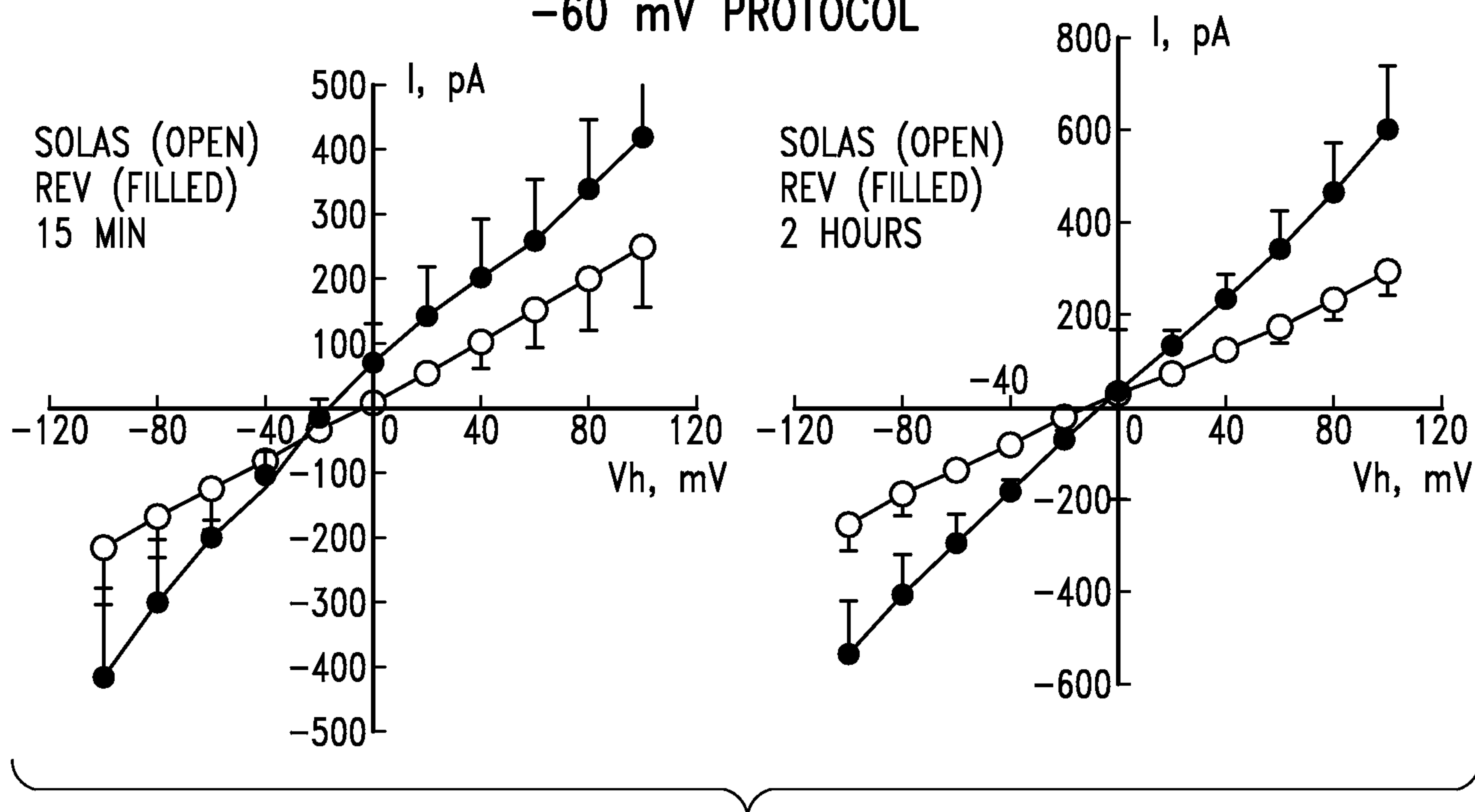
*Fig. 115**Fig. 116*

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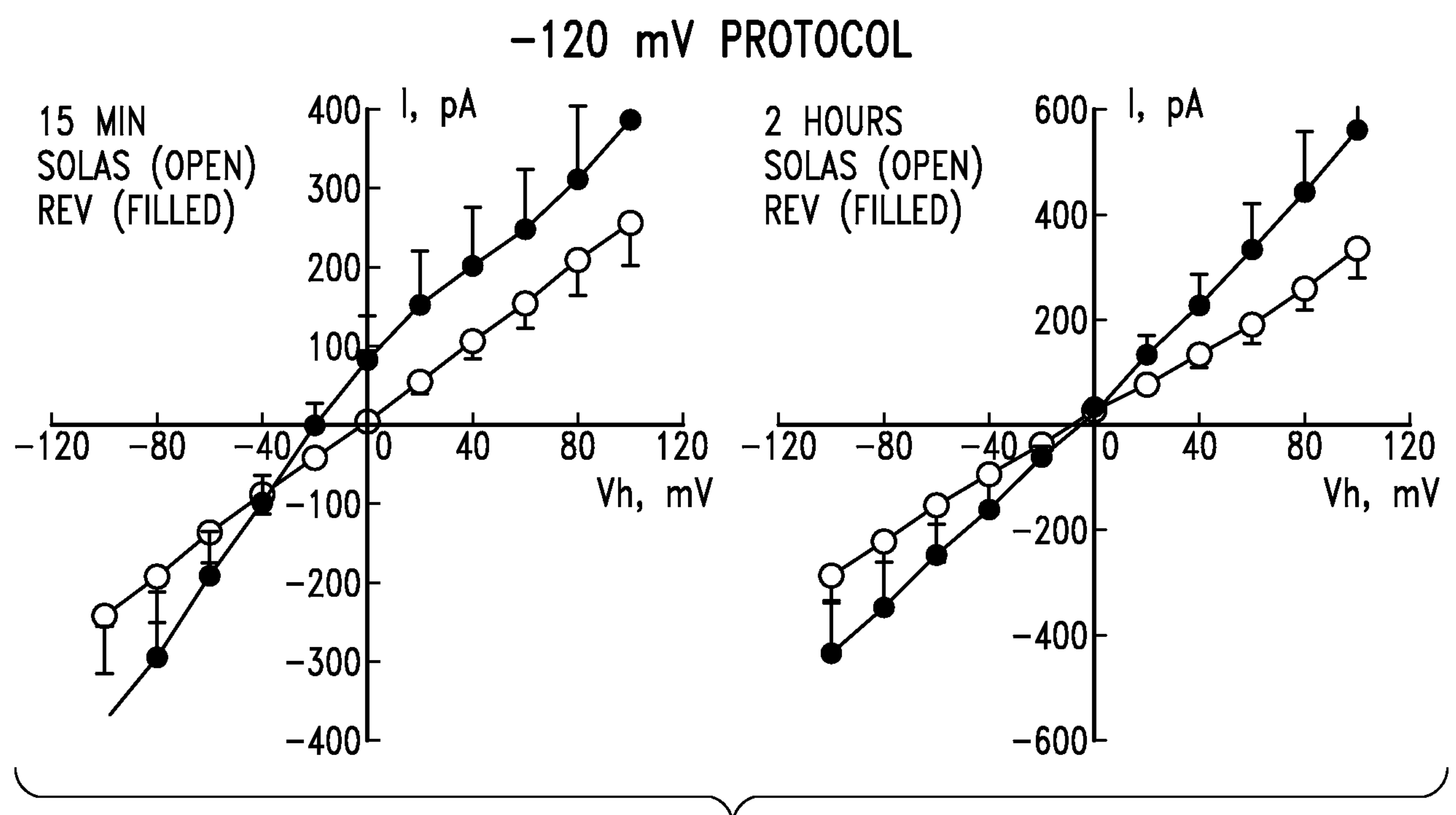
0 mV PROTOCOL

*Fig. 117A*

-60 mV PROTOCOL

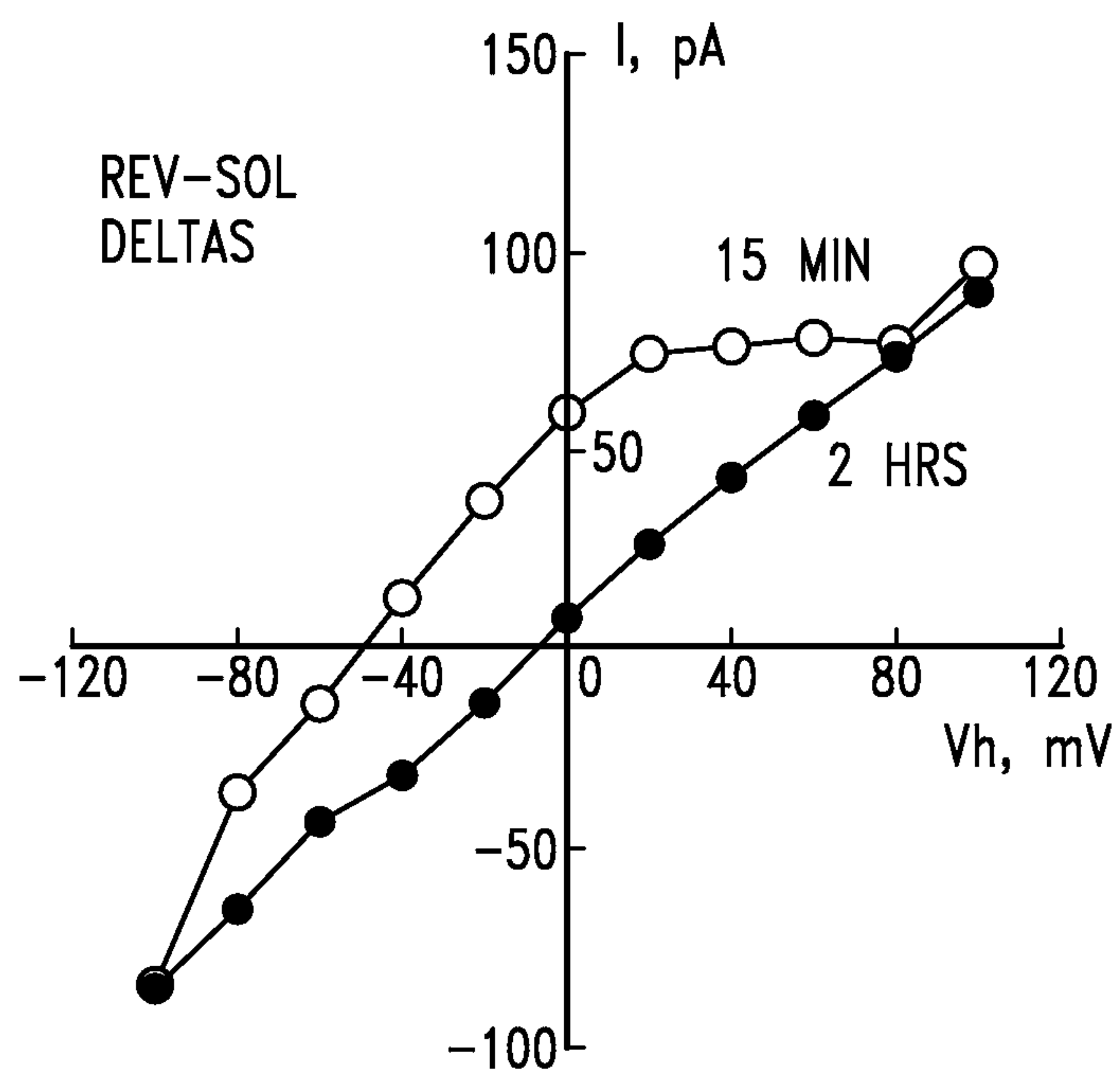
*Fig. 117B*

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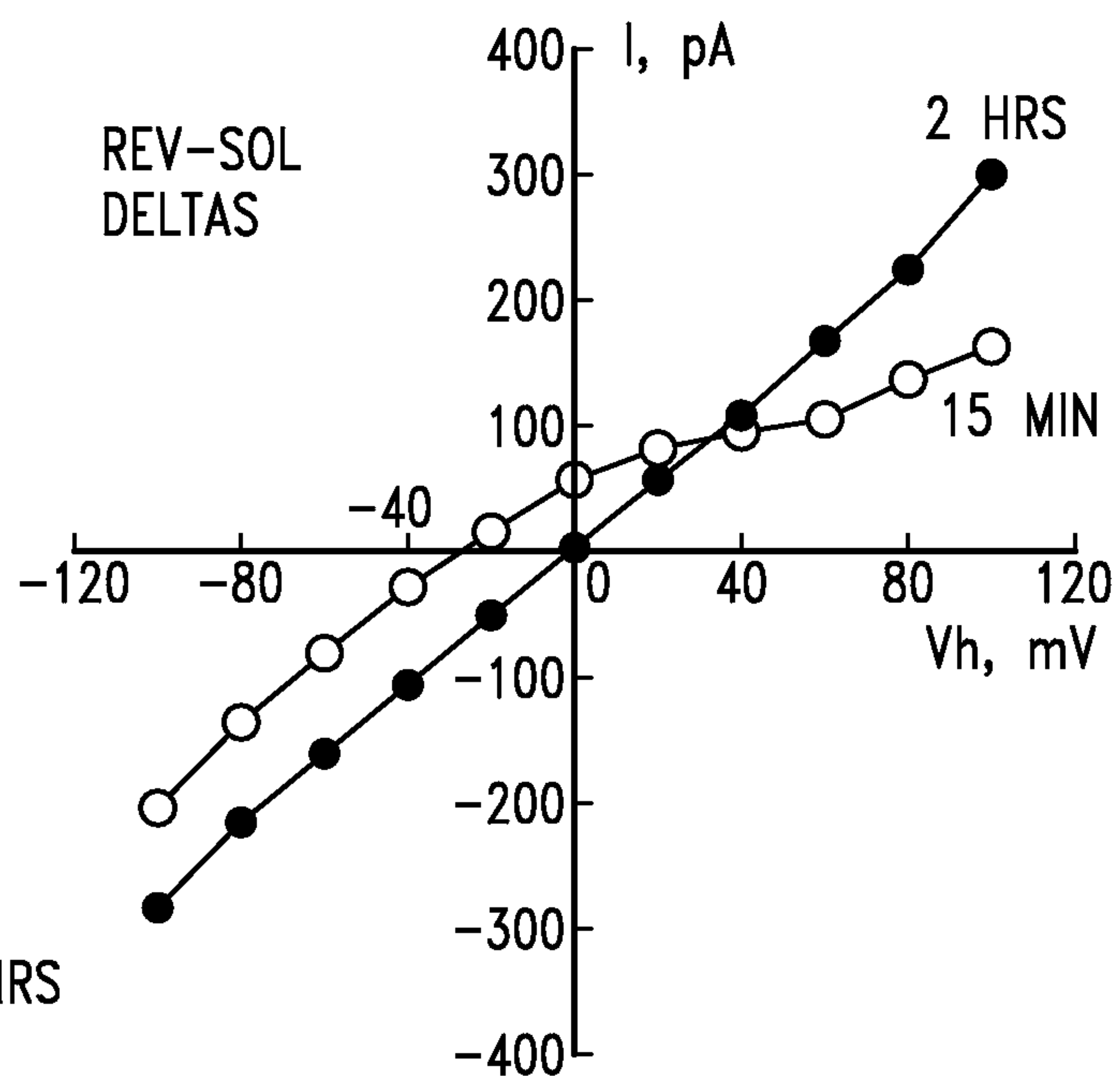
*Fig. 117C*

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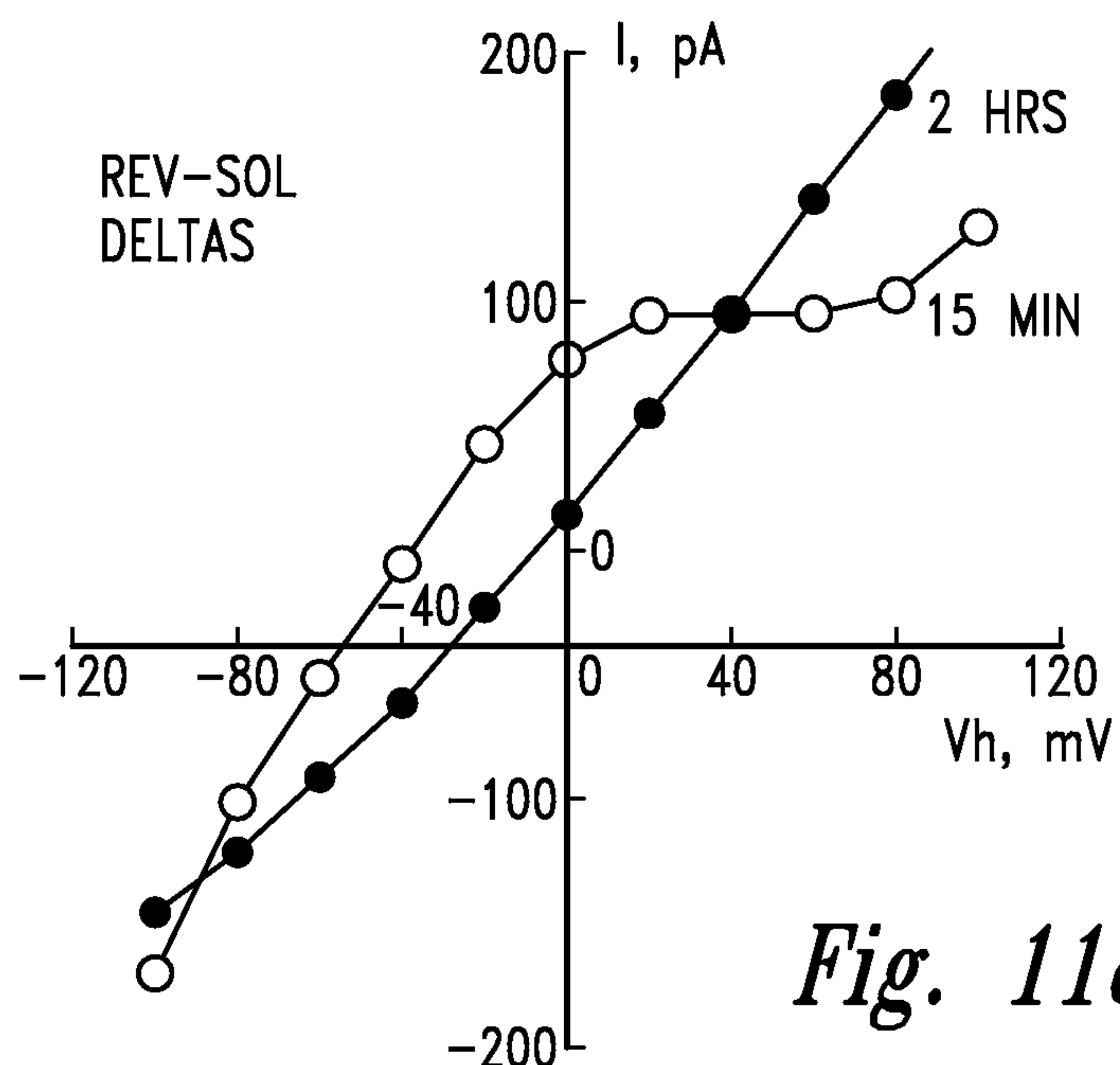
0 mV PROTOCOL

*Fig. 118A*

-60 mV PROTOCOL

*Fig. 118B*

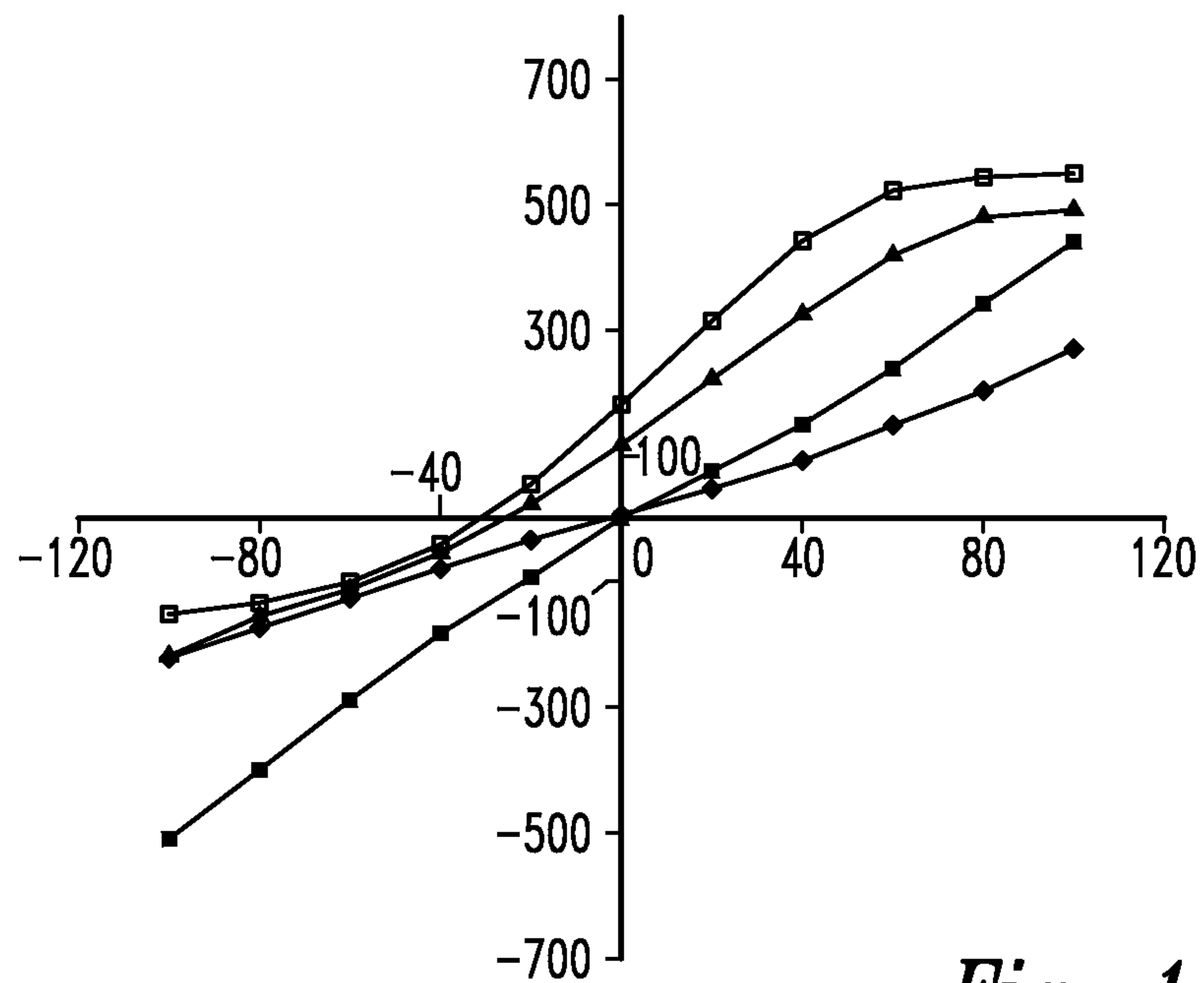
-120 mV PROTOCOL

*Fig. 118C*

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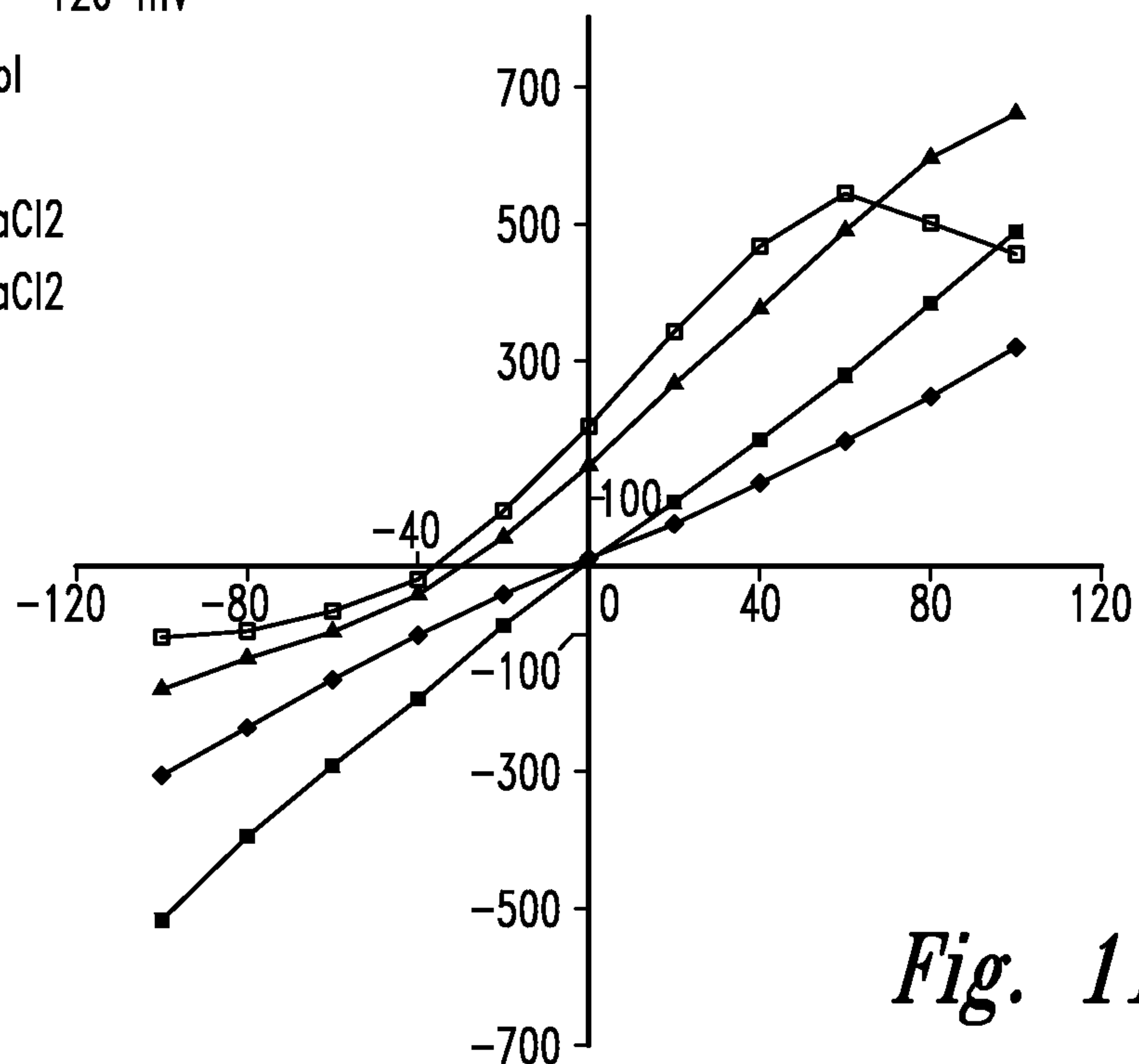
Solas 15 min, 0 mV

—◆— Control
—■— CsCl
—□— 20 CaCl₂
—▲— 40 CaCl₂

*Fig. 119A*

Solas 15 min, -120 mV

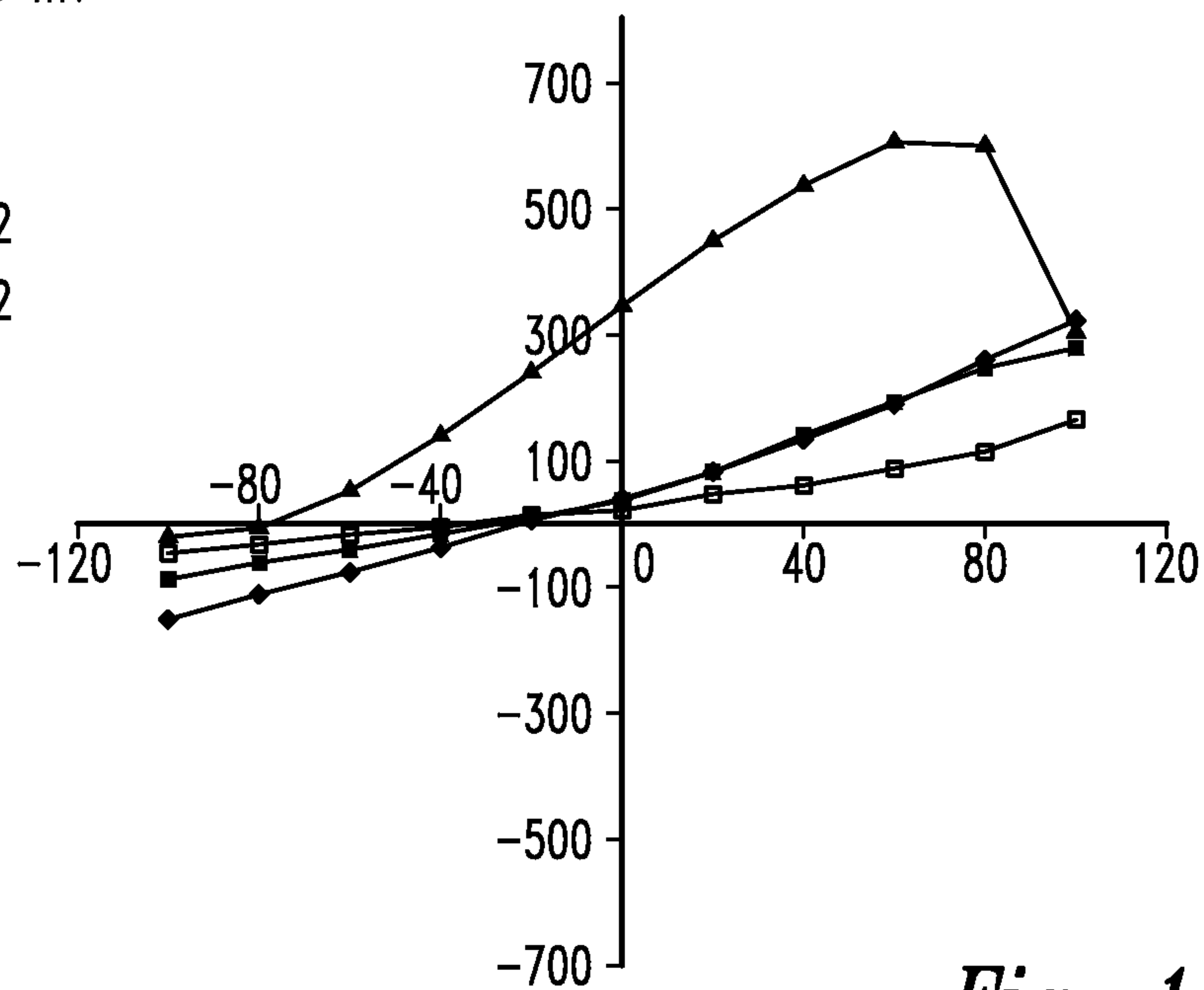
—◆— Control
—■— CsCl
—□— 20 CaCl₂
—▲— 40 CaCl₂

*Fig. 119B*

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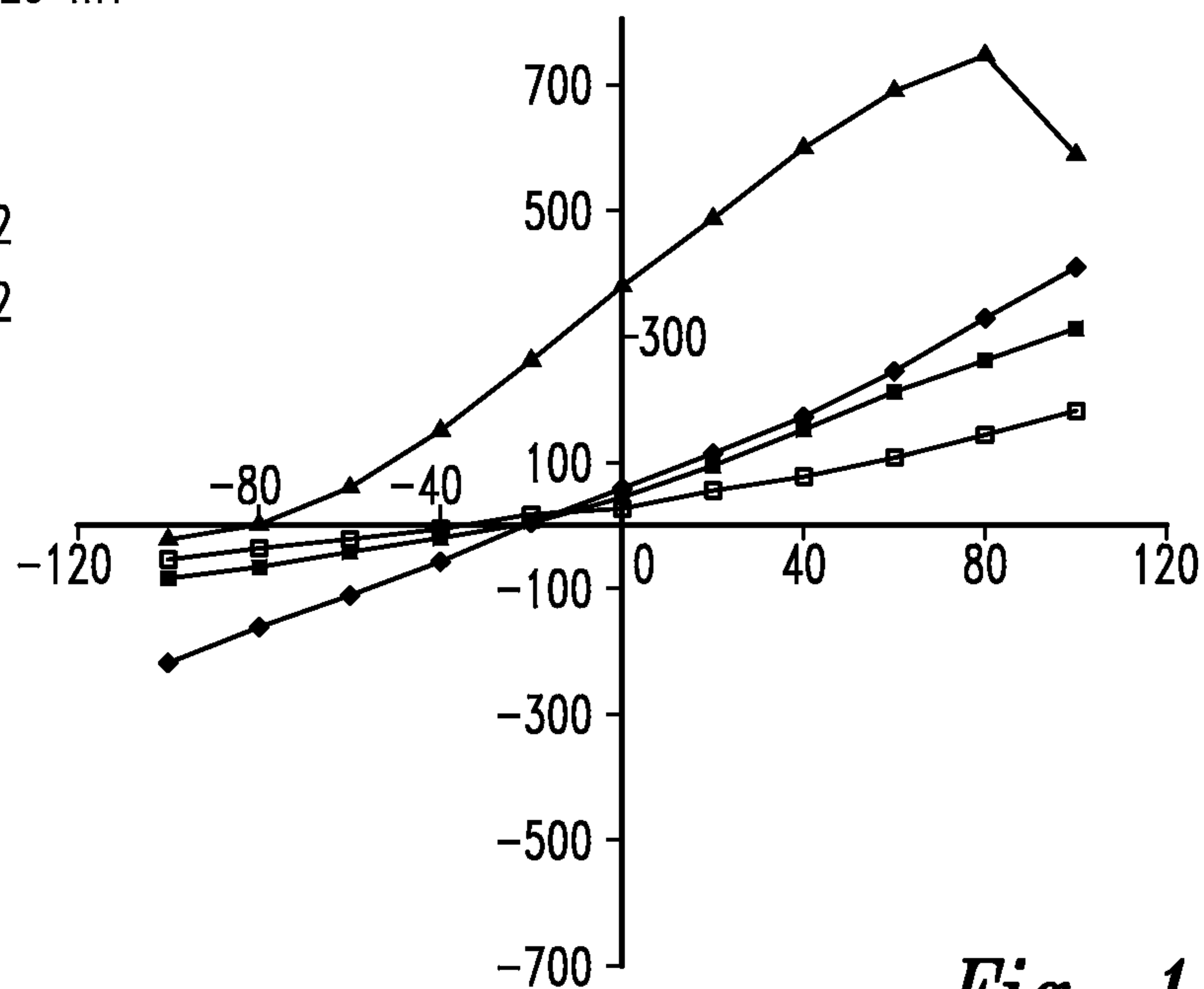
Rev 15 min, 0 mV

—◆— Control
—■— CsCl
—□— 20 CaCl₂
—▲— 40 CaCl₂

*Fig. 119C*

Rev 15 min, -120 mV

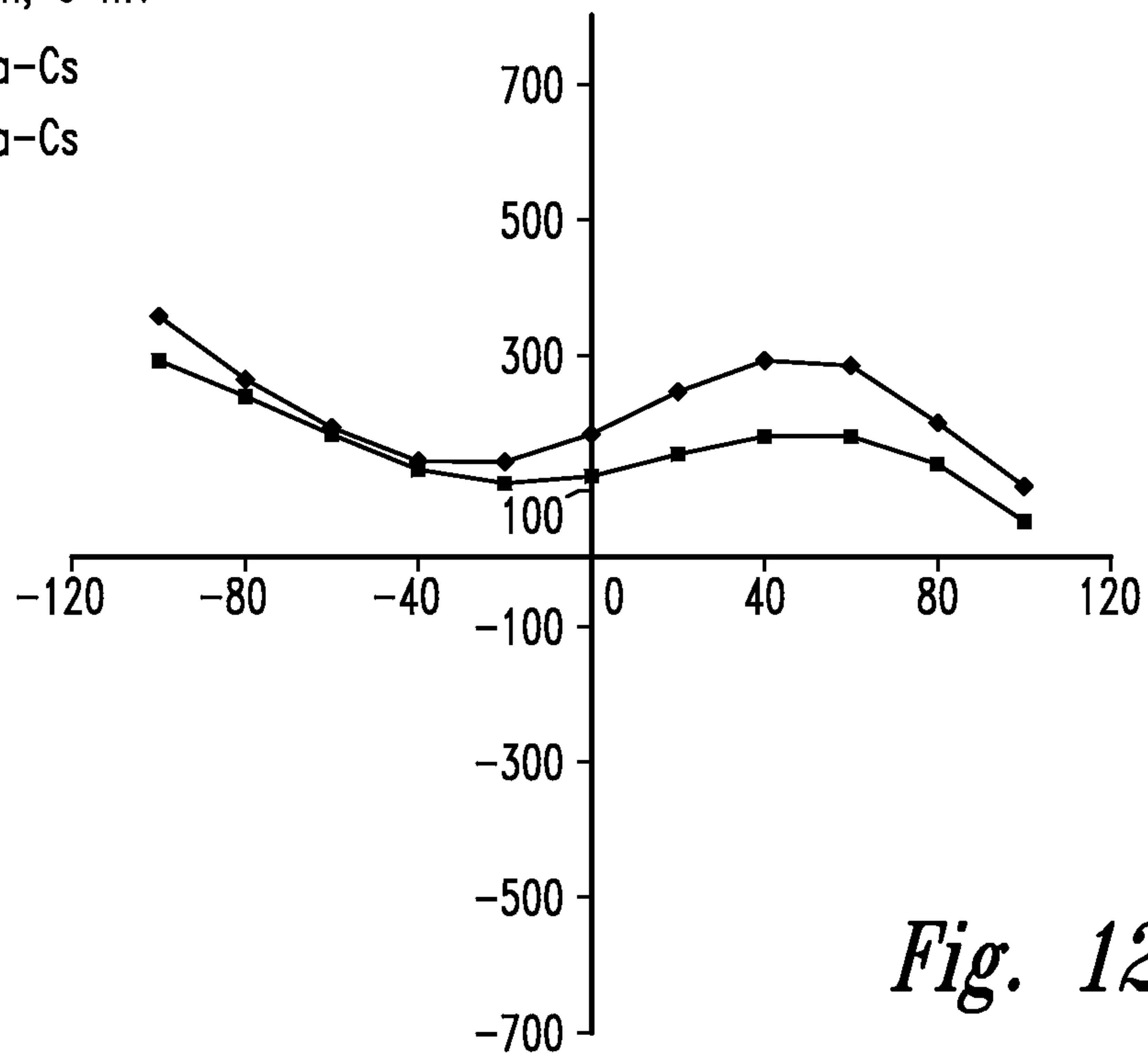
—◆— Control
—■— CsCl
—□— 20 CaCl₂
—▲— 40 CaCl₂

*Fig. 119D*

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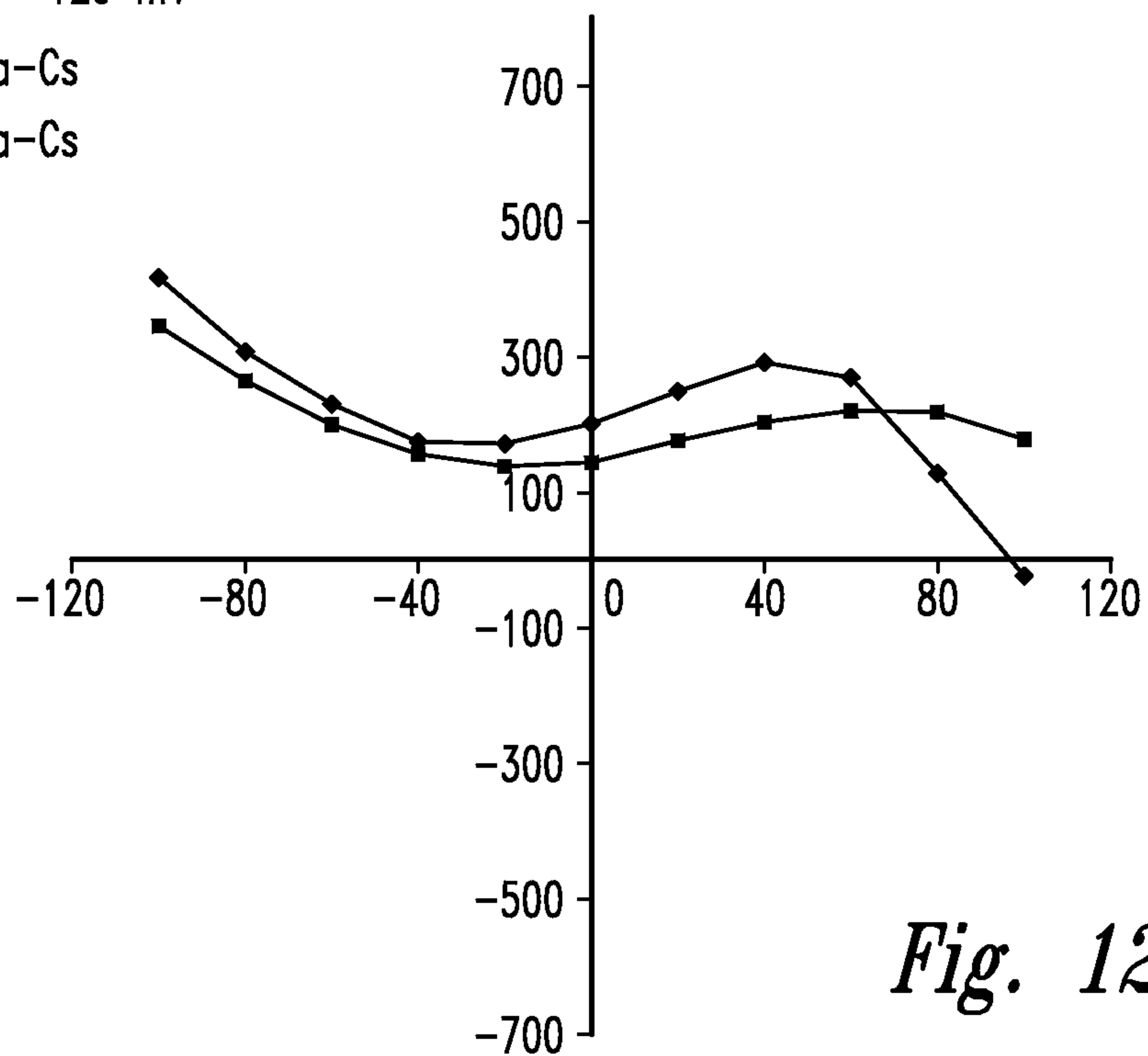
Solus 15 min, 0 mV

—◆— 20 Ca-Cs
—■— 40 Ca-Cs

*Fig. 120A*

Solus 15 min, -120 mV

—◆— 20 Ca-Cs
—■— 40 Ca-Cs

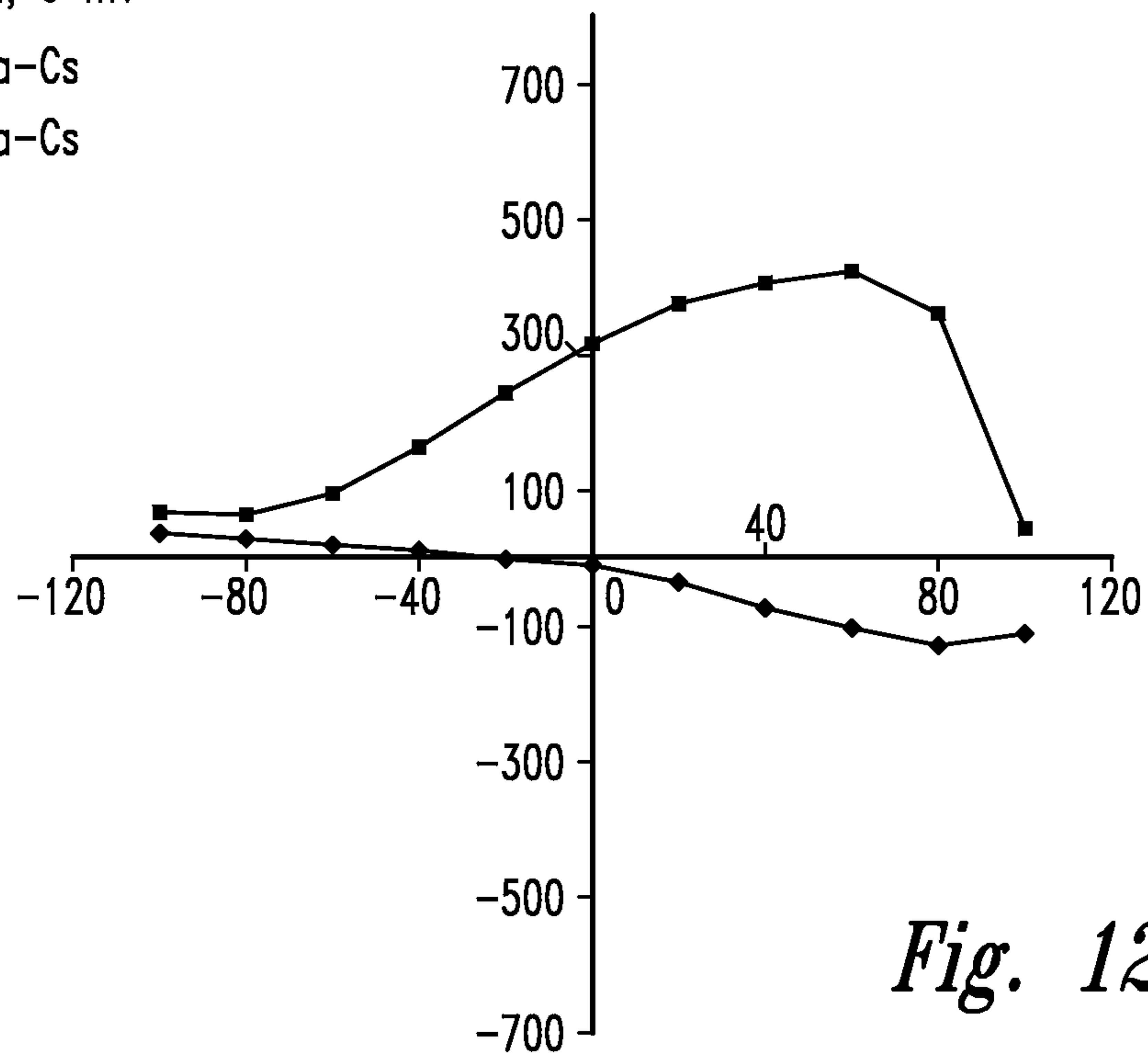
*Fig. 120B*

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Rev 15 min, 0 mV

—◆— 20 Ca-Cs

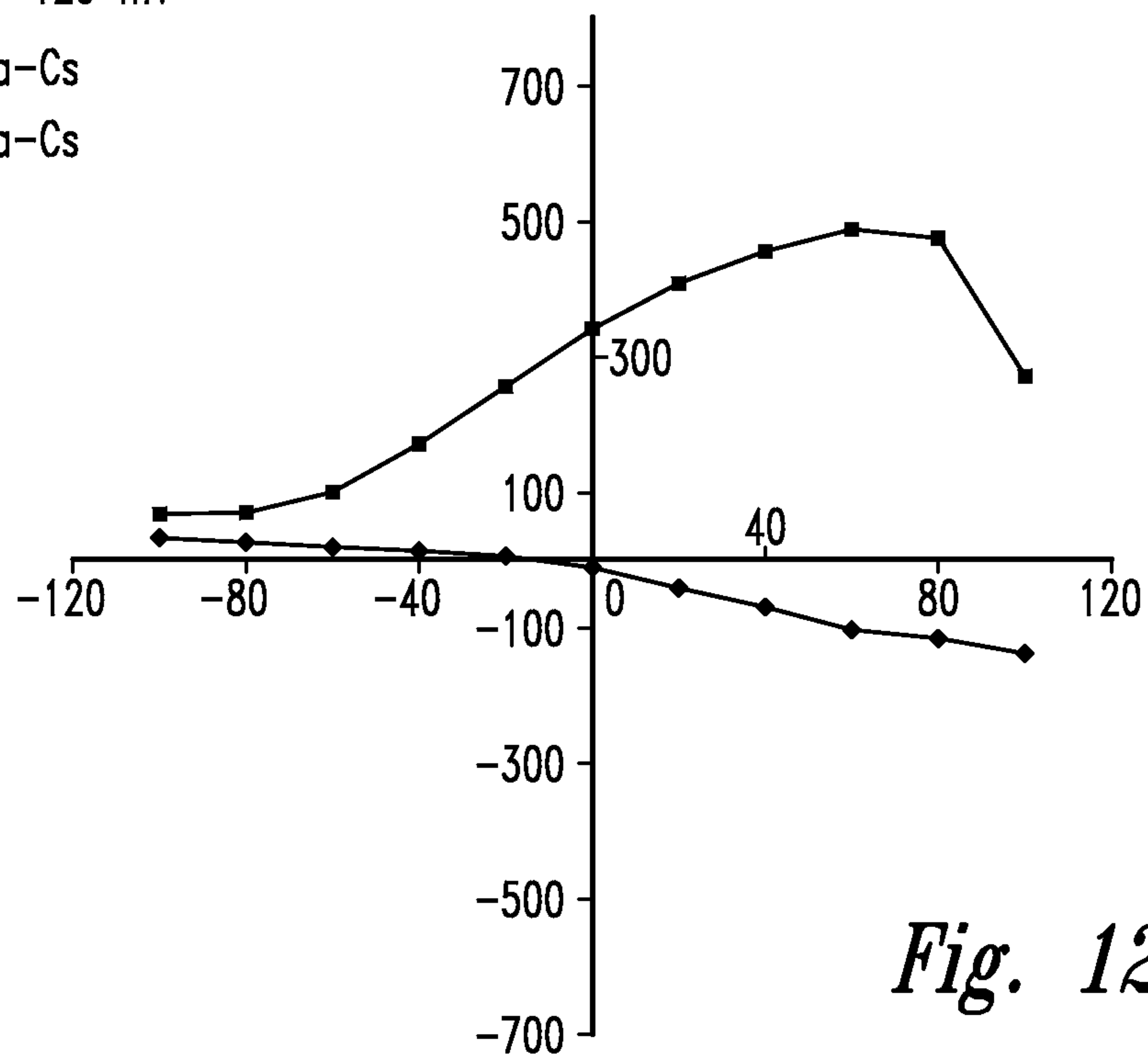
—■— 40 Ca-Cs

*Fig. 120C*

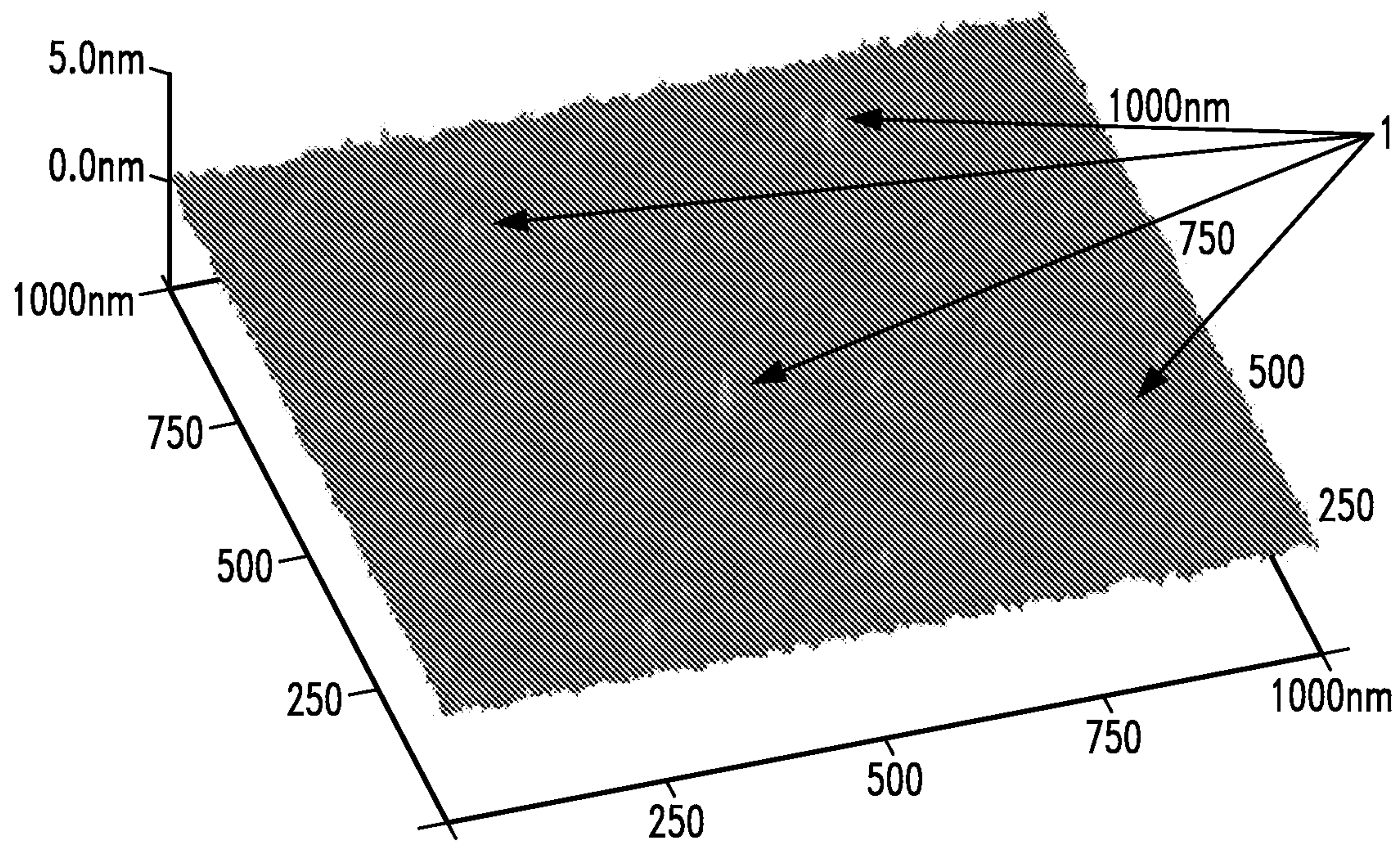
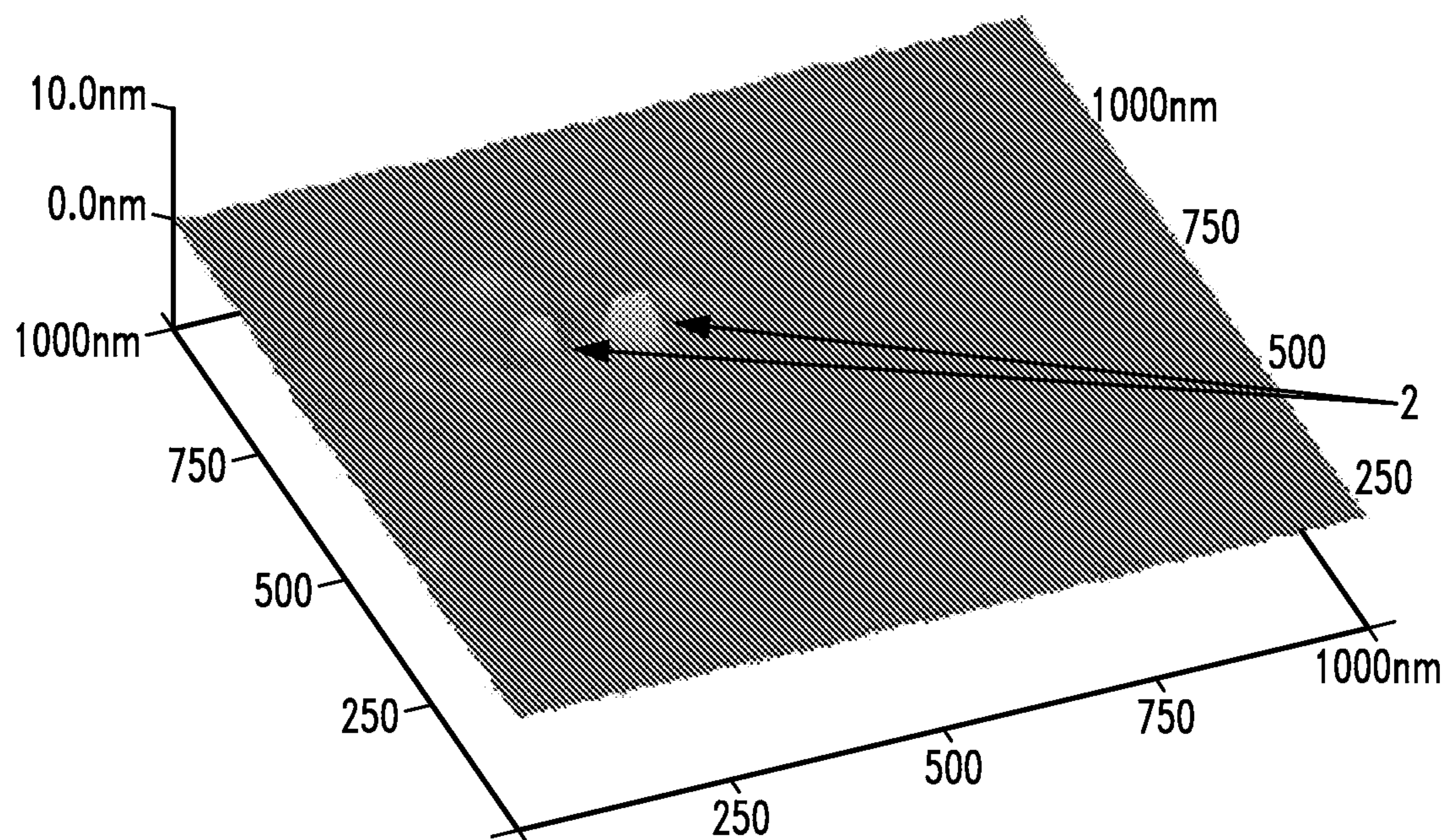
Rev 15 min, -120 mV

—◆— 20 Ca-Cs

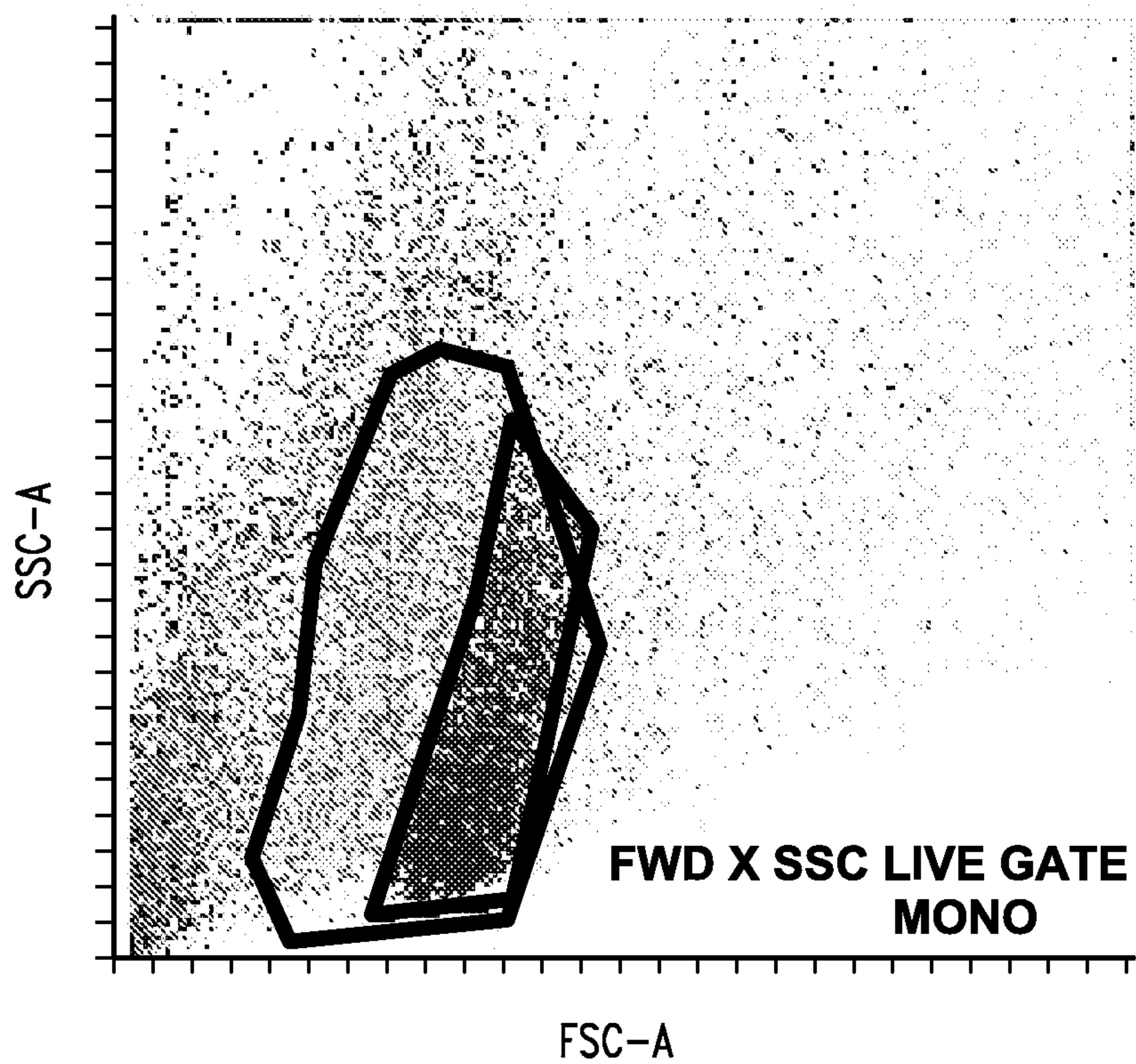
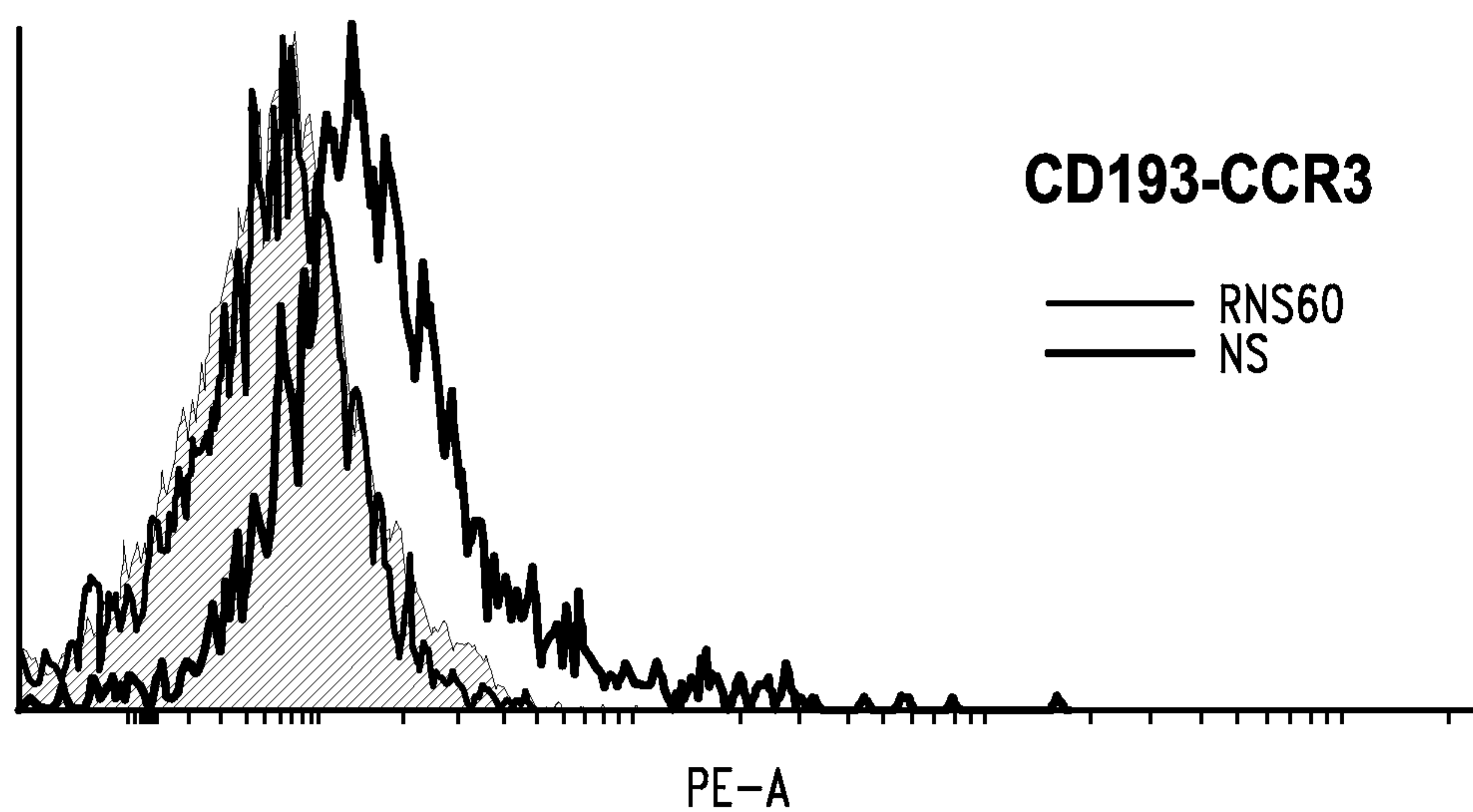
—■— 40 Ca-Cs

*Fig. 120D*

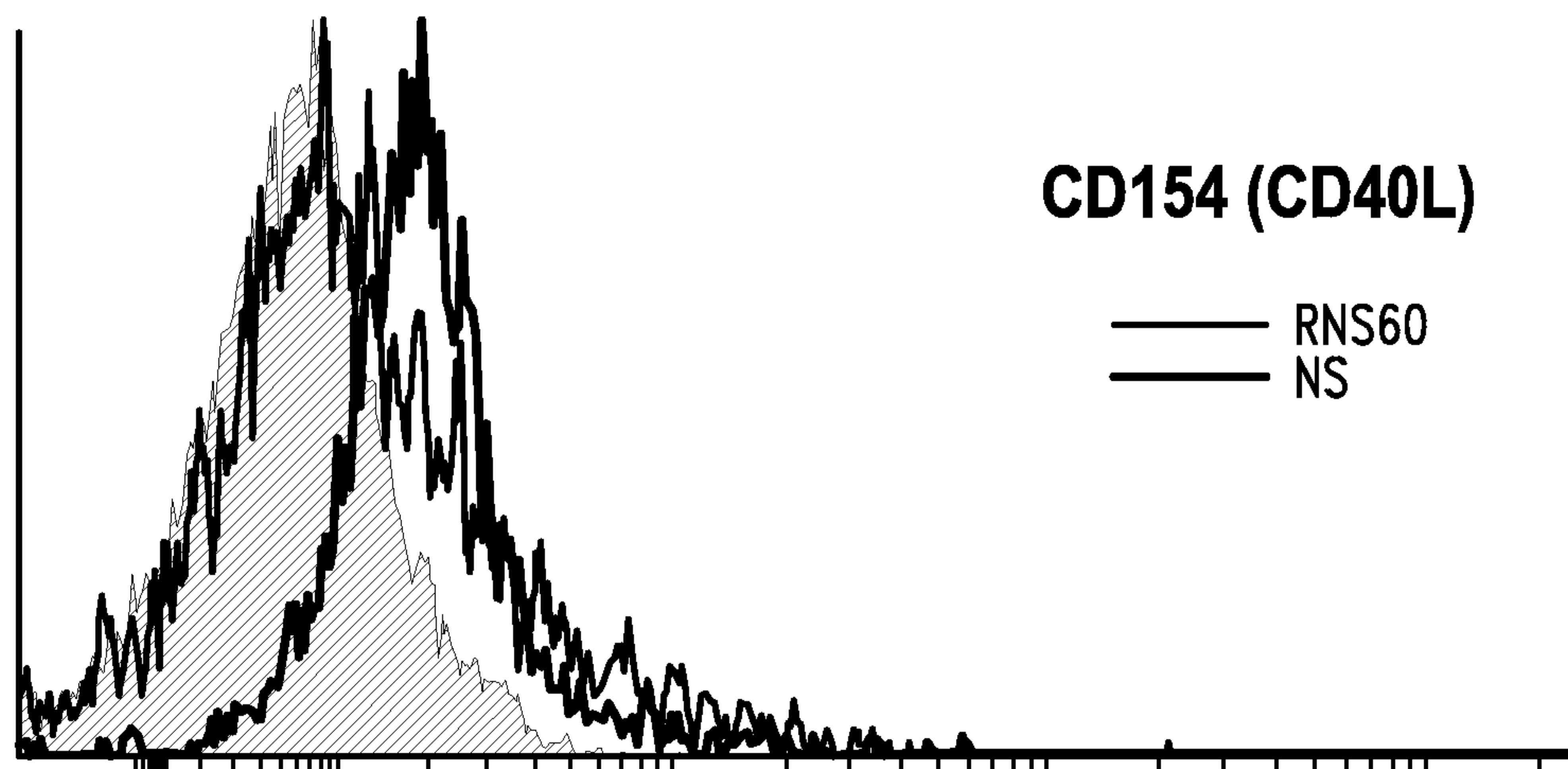
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*Fig. 121A**Fig. 121B*

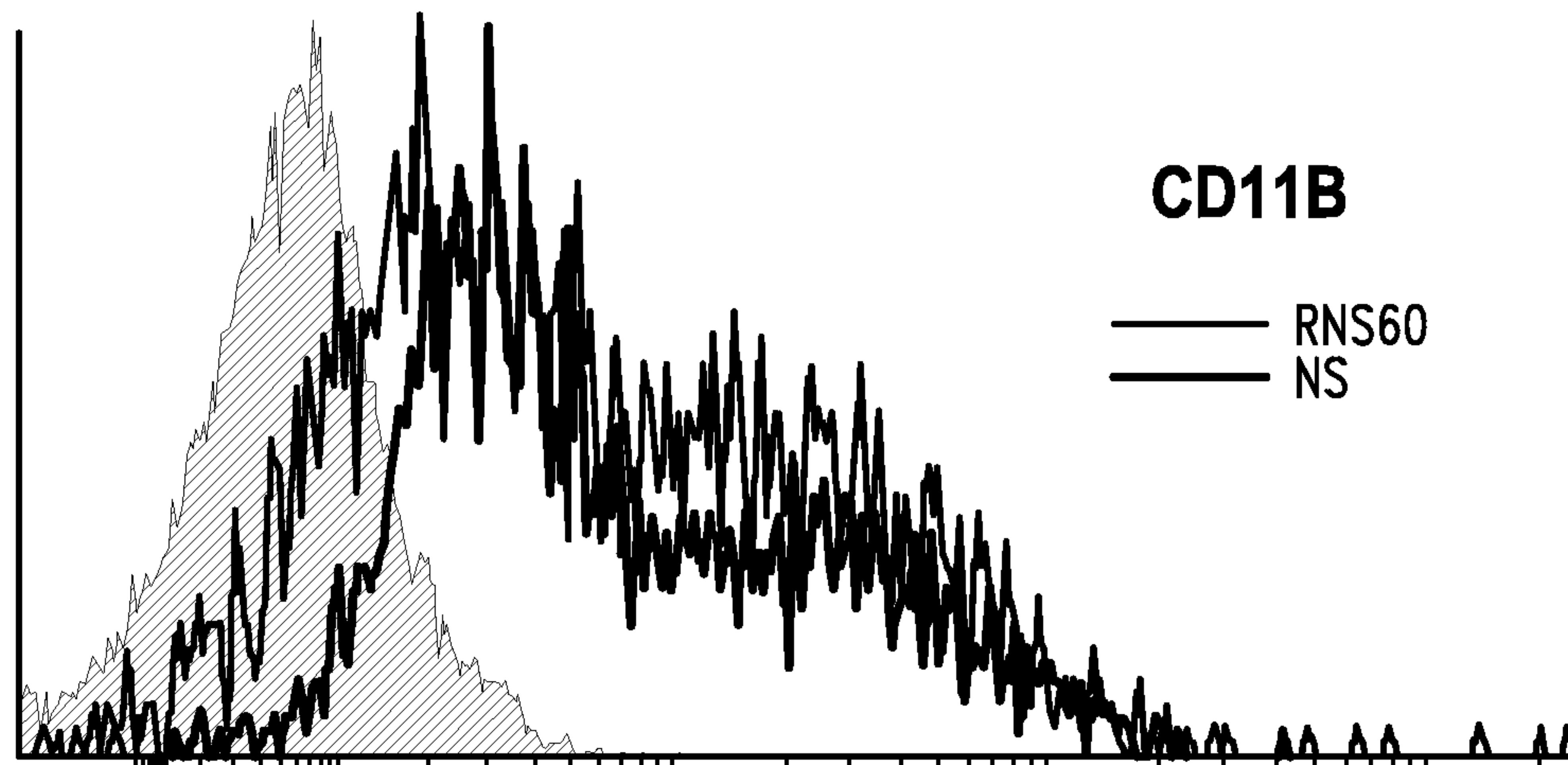
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*Fig. 122A**Fig. 122B*

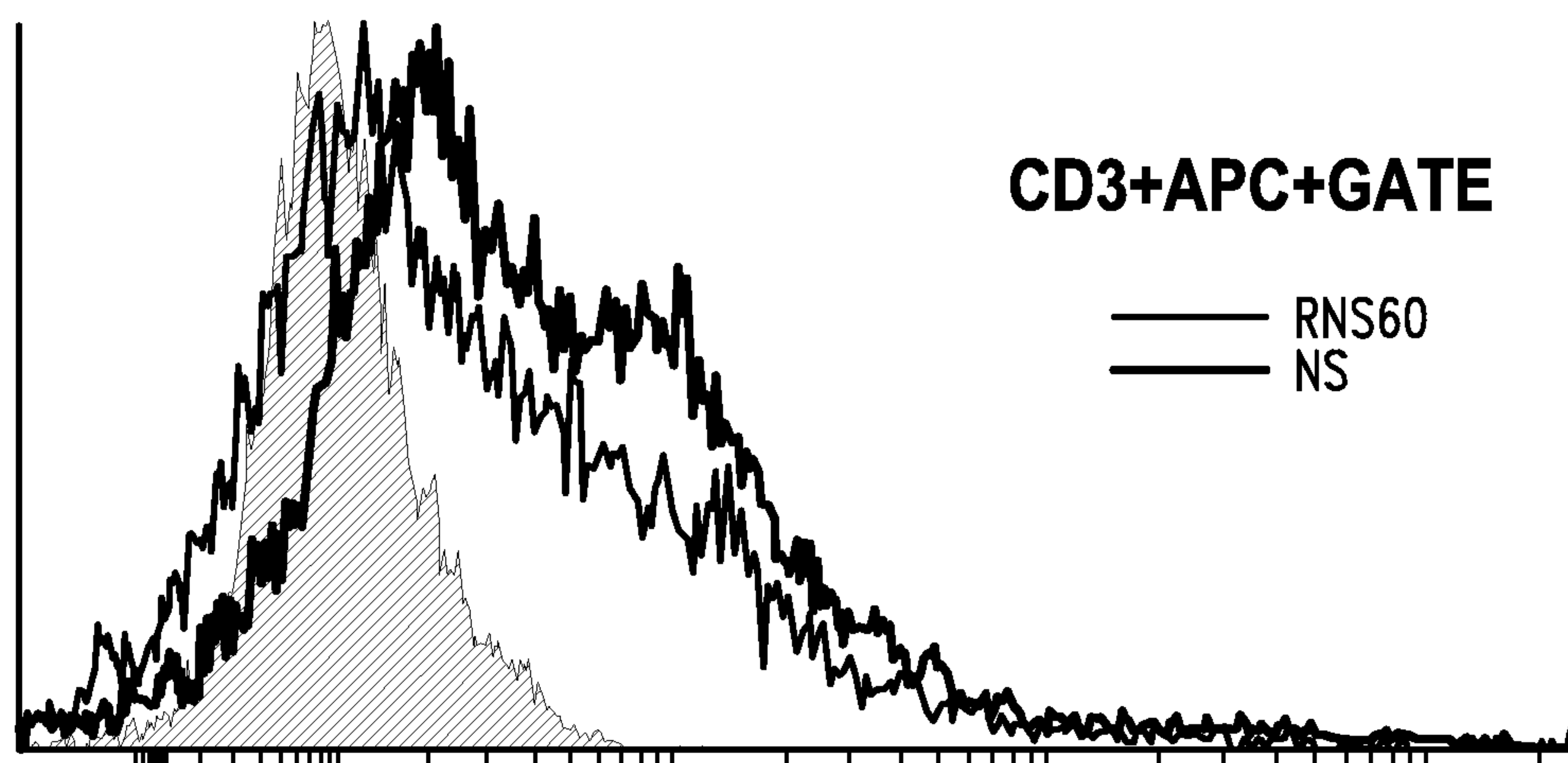
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*Fig. 123A*

PE-A

*Fig. 123B*

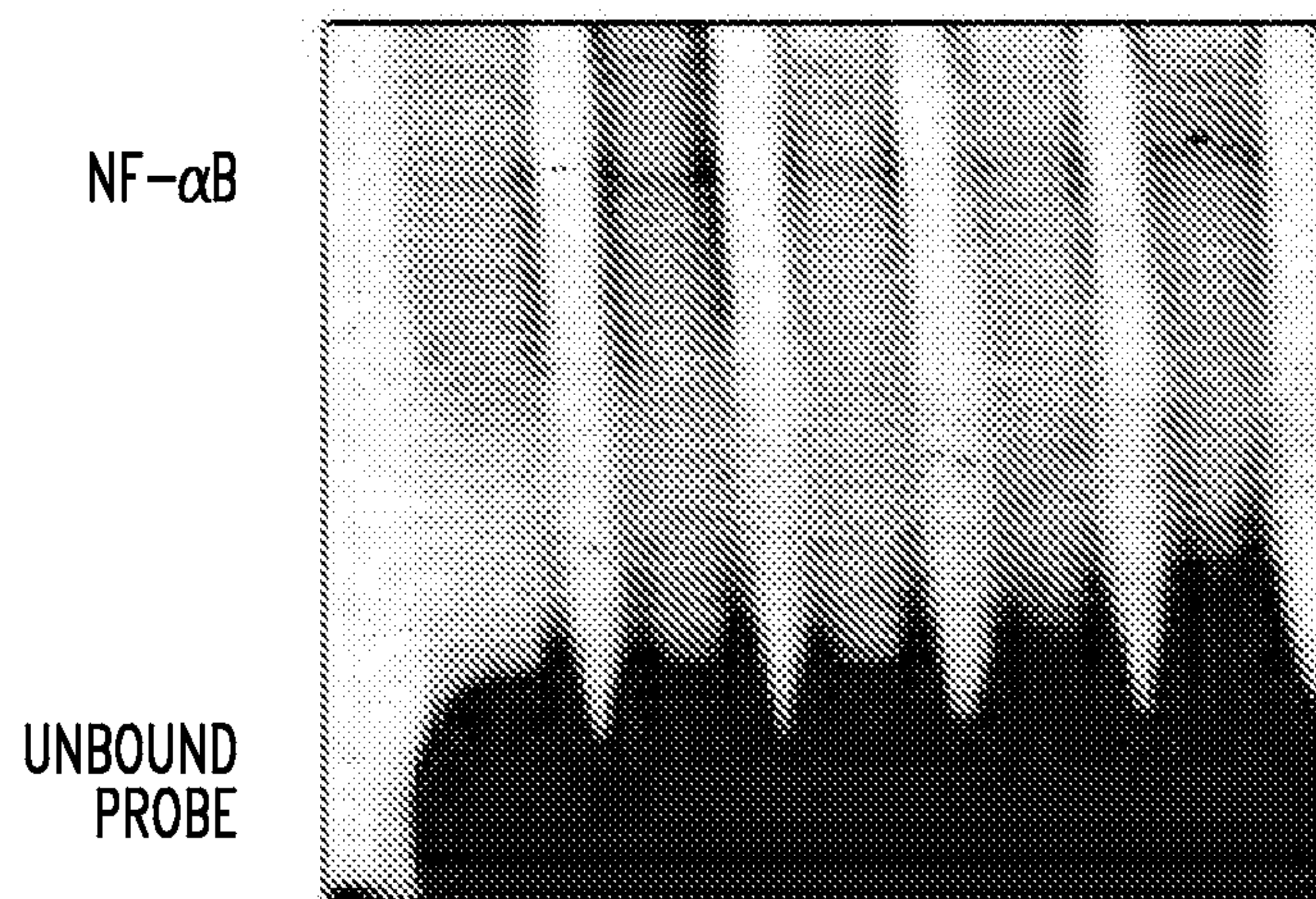
PE-A

*Fig. 123C*

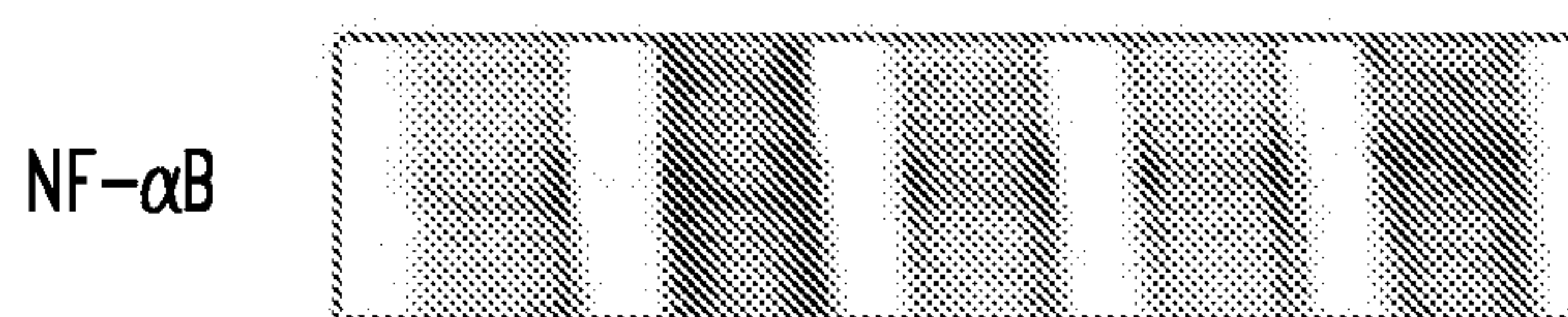
FITC-A

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NS (%)	0	0	0	0	10
RNS60 (%)	0	0	5	10	0
MBPT cells	-	+	+	+	+
Cont T cells	+	-	-	-	-



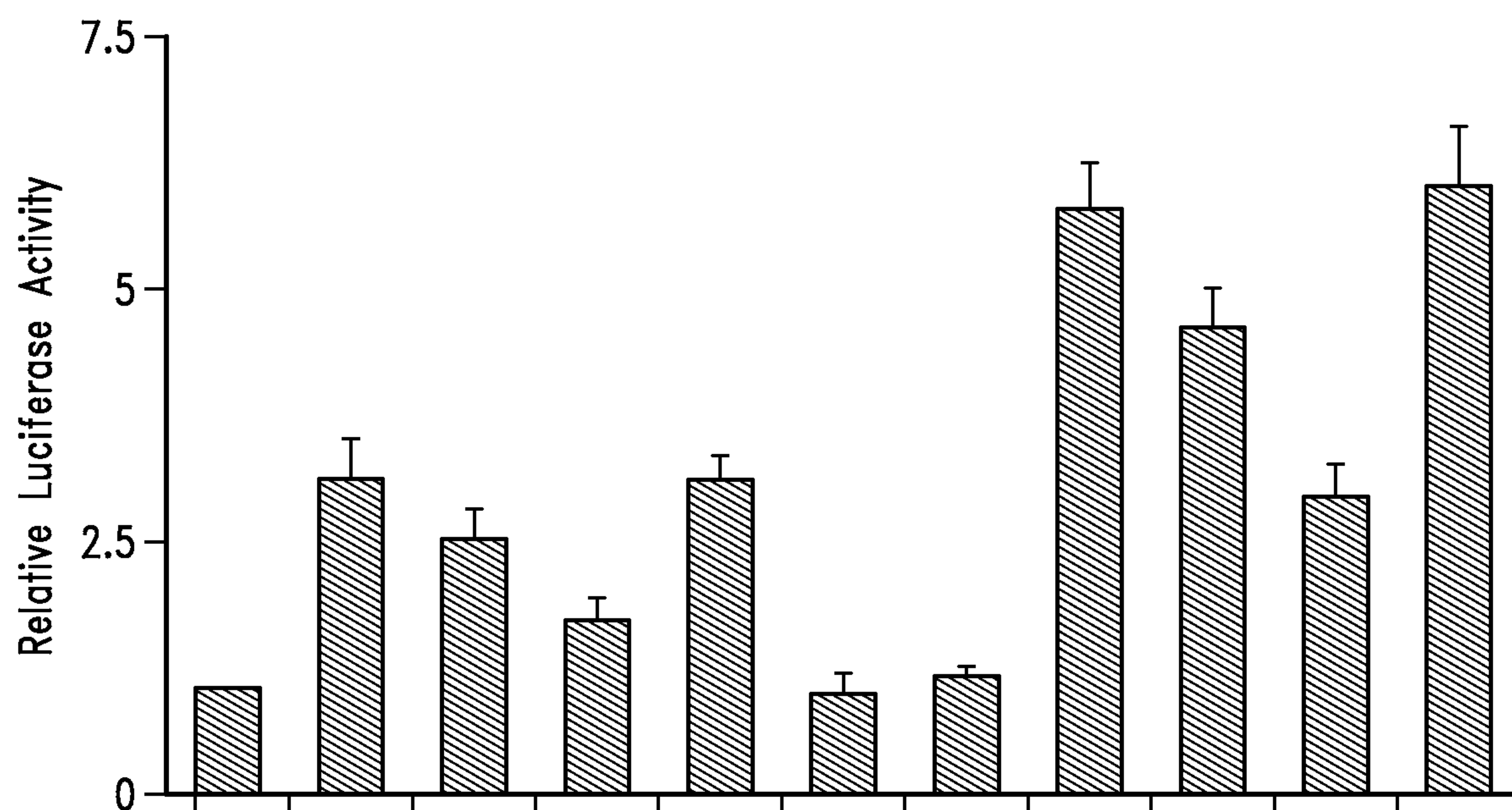
Expt-I

Fig. 124A

Expt-II

Fig. 124B

PMA	-	-	-	-	-	-	-	+	+	+	+
NS (%)	0	0	0	0	10	0	10	0	0	0	10
RNS60 (%)	0	0	5	10	0	10	0	0	5	10	0
MBPT cells	-	+	+	+	+	-	-	+	+	+	+
Cont T cells	+	-	-	-	-	+	+	-	-	-	-

*Fig. 124C*

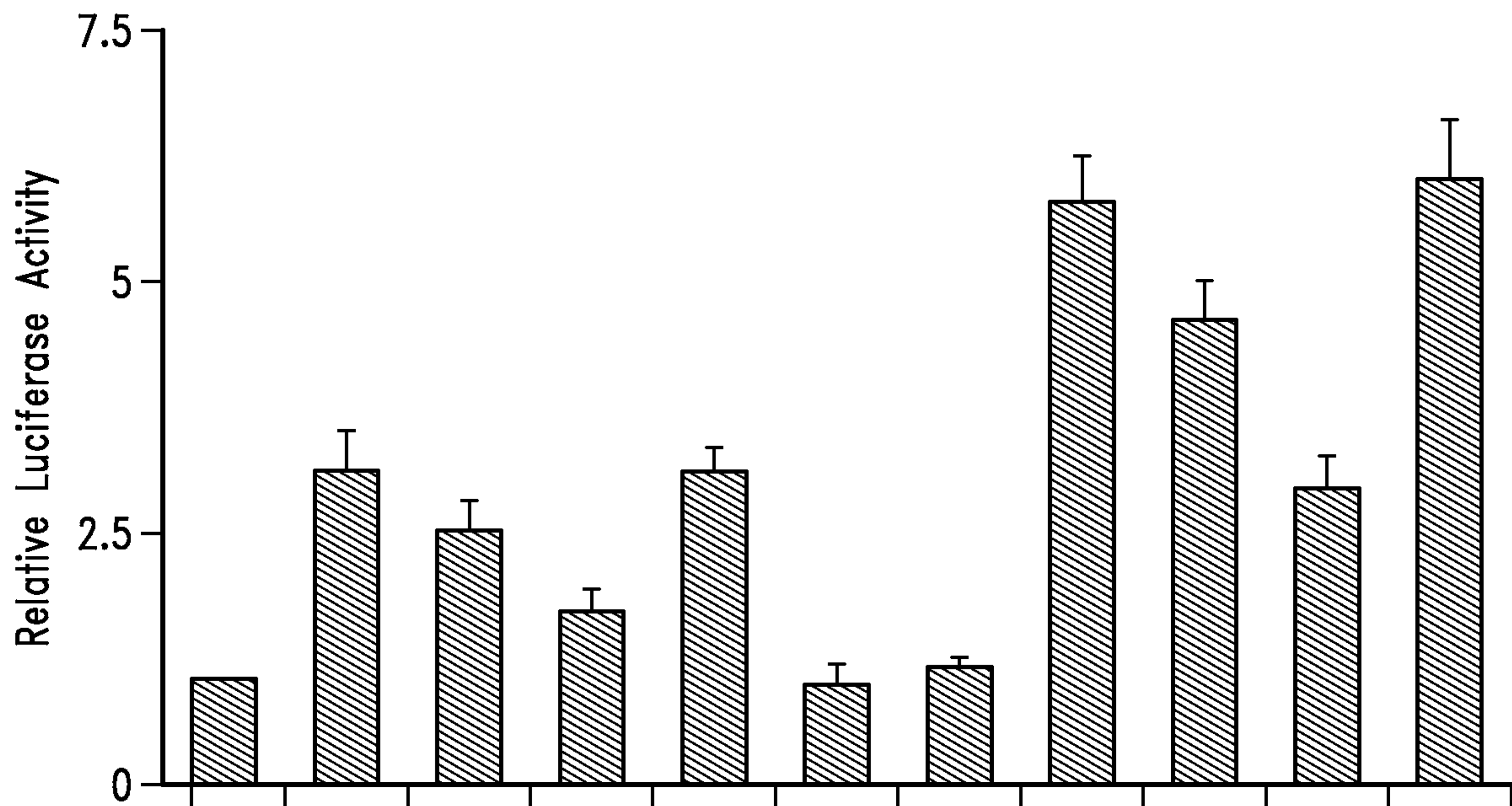


Fig. 124C