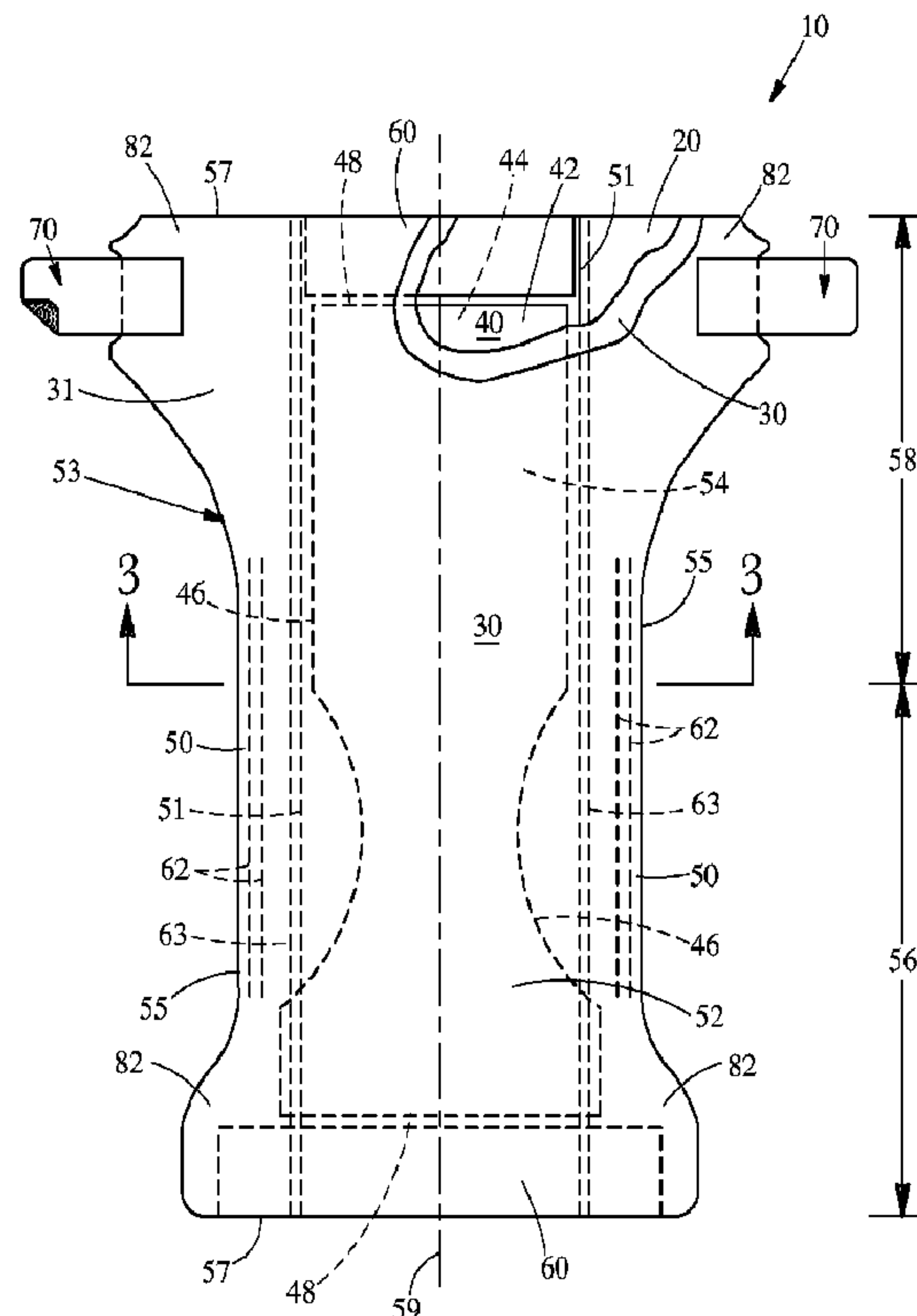




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(54) **Title: WEB MATERIAL(S) FOR ABSORBENT ARTICLES**



(57) **Abrégé/Abstract:**

The present disclosure, in part, relates generally to an absorbent article to be worn about the lower torso. The absorbent article comprises a chassis comprising a topsheet, a backsheet, an absorbent core disposed between the topsheet and the backsheet,



(57) **Abrégé(suite)/Abstract(continued):**

and a pair of longitudinal barrier cuffs attached to the chassis. Each of the longitudinal barrier cuffs is formed of a web of material. The web of material comprises a first nonwoven component layer comprising fibers having an average diameter in the range of about 8 microns to about 30 microns, a second nonwoven component layer comprising fibers having an average diameter of less than about 1 micron, and a third nonwoven component layer comprising fibers having an average diameter in the range of about 8 microns to about 30 microns. The second nonwoven component layer is disposed intermediate the first nonwoven component layer and the third nonwoven component layer.

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WEB MATERIAL(S) FOR ABSORBENT ARTICLES

FIELD OF THE INVENTION

The present disclosure generally relates to consumer products such as absorbent articles and methods for manufacturing the same, and more particularly relates to web material configurations for absorbent articles and methods of manufacturing the same.

BACKGROUND OF THE INVENTION

Nonwoven fabric webs may be useful in a wide variety of applications. Various nonwoven fabric webs may comprise spunbond, meltblown, spunbond (“SMS”) webs comprising outer layers of spunbond thermoplastics (e.g., polyolefins) and an interior layer of meltblown thermoplastics. Such SMS nonwoven fabric webs may comprise spunbond layers which are durable and an internal meltblown layer which is porous but which may inhibit fast strikethrough of fluids, such as bodily fluids, for example, or the penetration of bacteria through the fabric webs. In order for such a nonwoven fabric web to perform to particular characteristics, it may be desirable for the meltblown layer to have a fiber size and a porosity that assures breathability of the nonwoven fabric web while at the same time inhibiting the strikethrough of fluids.

Absorbent articles such as diapers, training pants, incontinent wear and feminine hygiene products, for example, may also utilize nonwoven fabric webs for many purposes such as liners, transfer layers, absorbent media, barrier layers and cuffs, backings, and other components. For many such applications, the barrier properties of the nonwoven fabric web play an important role in the performance of the fabric webs, such as the performance as a barrier to fluid penetration, for example. Absorbent articles may comprise multiple elements such as a liquid permeable topsheet intended to be placed next to the wearer’s skin, a liquid impermeable backsheet which forms, in use, the outer surface of the absorbent article, various barrier cuffs, and an absorbent core disposed between the topsheet and the backsheet.

When absorbent articles are produced, webs of materials, such as nonwoven materials, are bonded to each other. The bonding of these materials can be done for example via a mechanical bonding process. Reducing the manufacturing cost of absorbent articles by reducing the basis weight of the webs while preserving, if not improving, their functionality remains a challenge. For example, it is believed that when the combined basis weight of the webs to be

bonded is less than 30 gsm, a reduction in basis weight of currently available spunbond, or SMS nonwoven webs can result in a significantly higher rate of bond defects. Those defects can lead to increased leakage of the absorbent article. There remains a need to provide an absorbent article comprising low basis weight webs that have a high quality of bonds with a low rate of defect when webs are bonded together.

There is also a need for low basis weight nonwoven webs that may be used in the manufacture of absorbent articles at high production rates and packaged under significant compaction for extended periods of time while achieving and maintaining soft, air permeable (i.e. breathable) and liquid barrier materials with good tactile properties and good barrier properties to low surface tension fluid. Structural, mechanical and fluid-handling properties of available nonwoven webs are believed not to be sufficient. Therefore, there is also a need for improved nonwoven web structures.

Absorbent articles that incorporate nonwoven webs in elements that act as barriers to liquids should be able to contain low surface tension liquids. Currently available nonwoven webs often require expensive hydrophobic coatings or melt-additives that are added to the webs in order to achieve satisfactory low surface tension fluid strike-through times while remaining air permeable. It is believed that in addition to their cost, such coated/treated nonwoven webs may still not be sufficient to contain low surface tension body exudates with a surface tension of 45 Mn/m or less. As a result, there is a need for absorbent articles comprising breathable elements made of lower cost nonwoven webs having superior barrier properties. Such new nonwoven materials can enable the simplification of the absorbent article design, such as, for example, replacing a multiple layer barrier cuff construction with a single layer cuff construction.

SUMMARY OF THE INVENTION

In one embodiment, the present disclosure, in part, relates generally to an absorbent article to be worn about the lower torso. The absorbent article comprises a chassis comprising a topsheet, a backsheet, an absorbent core disposed between the topsheet and the backsheet, and a pair of longitudinal barrier cuffs attached to the chassis. Each of the longitudinal barrier cuffs is formed of a web of material. The web of material comprises a first nonwoven component layer comprising fibers having an average diameter in the range of about 8 microns to about 30 microns, a second nonwoven component layer comprising fibers having a number-average

diameter of less than about 1 micron, a mass-average diameter of less than about 1.5 microns, and a ratio of the mass-average diameter to the number-average diameter less than about 2, and a third nonwoven component layer comprising fibers having an average diameter in the range of about 8 microns to about 30 microns. The second nonwoven component layer is disposed intermediate the first nonwoven component layer and the third nonwoven component layer.

In one embodiment, the present disclosure, in part, relates generally to an absorbent article to be worn about the lower torso. The absorbent article comprises a chassis comprising a topsheet, a backsheet, an absorbent core disposed between the topsheet and the backsheet, and a pair of longitudinal barrier cuffs attached to the chassis. Each of the longitudinal barrier cuffs is formed of a web of material. The web of material comprises a first nonwoven component layer comprising fibers having an average diameter in the range of about 8 microns to about 30 microns, a second nonwoven component layer comprising fibers having a number-average diameter of less than about 1 micron, a mass-average diameter of less than about 1.5 microns, and a ratio of the mass-average diameter to the number-average diameter less than about 2, a third nonwoven component layer comprising fibers having an average diameter in the range of about 8 microns to about 30 microns, and a fourth nonwoven component layer comprising fibers having an average diameter in the range of about 1 micron to about 8 microns. The second and fourth nonwoven component layers are disposed intermediate the first nonwoven component layer and the third nonwoven component layer.

In one embodiment, the present disclosure, in part, relates generally to an absorbent article to be worn about the lower torso. The absorbent article comprises a chassis comprising a topsheet, a backsheet, an absorbent core disposed between the topsheet and the backsheet, and a pair of longitudinal barrier cuffs attached to the chassis. Each of the longitudinal barrier cuffs is formed of a web of material. The web of material comprises a first nonwoven component layer comprising fibers having an average denier in the range of about 0.4 to about 6, a second nonwoven component layer comprising fibers having an average denier in the range of about 0.00006 to about 0.006, and a third nonwoven component layer comprising fibers having an average denier in the range of about 0.4 to about 6. The second nonwoven component layer is disposed intermediate the first nonwoven component layer and the third nonwoven component layer.

BRIEF DESCRIPTION OF THE DRAWINGS

The above-mentioned and other features and advantages of the present disclosure, and the manner of attaining them, will become more apparent and the disclosure itself will be better understood by reference to the following description of non-limiting embodiments of the disclosure taken in conjunction with the accompanying drawings, wherein:

FIG. 1 is a plan view of an absorbent article in accordance with one non-limiting embodiment of the present disclosure.

FIG. 2 is a perspective view of the absorbent article of FIG. 1.

FIGS. 3 A-B are cross-sectional views of the absorbent article of FIG. 1 taken along line 3--3.

FIG. 4 is a schematic diagram of a forming machine used to make a nonwoven web of material in accordance with one non-limiting embodiment of the present disclosure.

FIG. 5 is a cross-sectional view of a nonwoven web of material in a three layer configuration in accordance with one non-limiting embodiment of the present disclosure.

FIG. 6 is a perspective view of the web of material of FIG. 5 with various portions of nonwoven component layers cut away to show the composition of each nonwoven component layer in accordance with one non-limiting embodiment of the present disclosure.

FIG. 7 is a top view photograph of a web of material.

FIG. 8 is a cross-sectional photograph of the web of material of FIG. 7 taken through a calendering bond.

FIG. 9 is a top view photograph of a web of material in accordance with one non-limiting embodiment of the present disclosure.

FIG. 10 is a cross-sectional photograph of the web of material of FIG. 9 taken through a calendering bond in accordance with one non-limiting embodiment of the present disclosure.

FIG. 11 is a cross-sectional view of a web of material in a four layer configuration in accordance with one non-limiting embodiment of the present disclosure.

FIG. 12 is a perspective view of the web of material of FIG. 11 with various portions of nonwoven component layers cut away to show the composition of each nonwoven component layer in accordance with one non-limiting embodiment of the present disclosure.

FIG. 13 is a top view photograph of a web of material in accordance with one non-limiting embodiment of the present disclosure.

FIG. 14 is a cross-sectional photograph of the web of material of FIG. 13 in accordance with one non-limiting embodiment of the present disclosure.

FIG. 15 illustrates a simplified dynamic mechanical bonding apparatus in accordance with one non-limiting embodiment of the present disclosure.

FIG. 16 illustrates a patterned cylinder in accordance with one non-limiting embodiment of the present disclosure.

FIG. 17 is a plan view of a fragmentary portion of a bonded web of material in accordance with one non-limiting embodiment of the present disclosure.

FIG. 18A-D illustrate patterns of bond sites in accordance with various non-limiting embodiments of the present disclosure.

FIG. 19 is a cross-sectional view taken along line 19--19 of FIG. 17, which illustratively shows a bond site in accordance with one non-limiting embodiment of the present disclosure.

FIG. 20 is a cross-sectional perspective view of the bond site of FIG. 19.

FIG. 21A illustrates mechanical bond quality and the templates for determining defects.

FIG. 21B illustrates the use of defect area templates for defects of holes, skips and tears.

FIGS. 22-25 graphically illustrate data from Tables 1A and 1B of Example 1.

FIG. 26 graphically illustrates the low surface tension fluid strikethrough times of various samples of Table 2A of Example 2A.

FIG. 27 graphically illustrates the low surface tension fluid strikethrough times of various samples of Table 2B of Example 2B compared the number-average diameter of the samples.

FIG. 28 graphically illustrates the sidedness of an SMNS web of the present disclosure having the properties specified in Table 2C.

FIGS. 29 and 30 graphically illustrate the low surface tension fluid strikethrough times of various SMS webs compared with the low surface tension fluid strikethrough times of the SMNS webs of the present disclosure.

FIG. 31 graphically illustrates the pore size distribution of Samples G, B, and A with respect to Example 3.

FIG. 32 graphical illustrates the bond defects of various samples of Table 32 as a function of basis weight COV.

FIGS 33A-33G illustrate examples of various mechanical bonds.

DETAILED DESCRIPTION OF THE INVENTION

Various non-limiting embodiments of the present disclosure will now be described to provide an overall understanding of the principles of the structure, function, manufacture, and use of the apparatuses and methods disclosed herein. One or more examples of these non-limiting embodiments are illustrated in the accompanying drawings. Those of ordinary skill in the art will understand that the apparatuses and methods specifically described herein and illustrated in the accompanying drawings are non-limiting example embodiments and that the scope of the various non-limiting embodiments of the present disclosure are defined solely by the claims. The features illustrated or described in connection with one non-limiting embodiment may be combined with the features of other non-limiting embodiments. Such modifications and variations are intended to be included within the scope of the present disclosure.

Definitions:

In this description, the following terms have the following meanings:

The term “absorbent article” refers to a device that is placed against or in proximity to a body of a wearer to absorb and contain various exudates discharged from the body. Example absorbent articles comprise diapers, training pants, pull-on pant-type diapers (i.e., a diaper having a pre-formed waist opening and leg openings, such as illustrated in U.S. Pat. No. 6,120,487, issued to Ashton, on September 19, 2000), refastenable diapers, incontinence briefs and undergarments, diaper holders and liners, feminine hygiene garments, panty liners, and absorbent inserts, for example.

The term “air permeability” is defined by the Air Permeability Test set forth below. Air permeability is set forth in $\text{m}^3/\text{m}^2/\text{minute}$ (m/min).

The term “basis weight” is defined by the Basis Weight Test set forth below. Basis weight is set forth in grams/m^2 (gsm).

The term “bond area” refers to the area of an individual bond site. Bond area is set forth in mm^2 .

The term “bond density” is the number of bonds in an area. Bond density is set forth in bonds per cm^2 . A relative bond area is the bond density multiplied by the bond area (all converted to same unit area), and given in a percentage.

The term “cross direction” refers to a direction that is generally perpendicular to the machine direction.

The term “defect occurrence rate” is defined by the Defect Occurrence Rate Test set forth below.

The term “denier” refers to a unit of fineness of a fiber that is equal to the weight (in grams) per 9000 m of fiber.

The term “diameter” when referring to fibers is defined by the Fiber Diameter and Denier Test set forth below. Diameter of fibers is set forth in microns.

The term “elongatable material,” “extensible material,” or “stretchable material” are used interchangeably and refer to a material that, upon application of a biasing force, can stretch to an elongated length of at least 150% of its relaxed, original length (i.e. can stretch to 50% more than its original length), without complete rupture or breakage as measured by EDANA method 20.2-89. In the event such an elongatable material recovers at least 40% of its elongation upon release of the applied force, the elongatable material will be considered to be “elastic” or “elastomeric.” For example, an elastic material that has an initial length of 100mm can extend to 150mm, and upon removal of the force retracts to a length of at least 130mm (i.e., exhibiting a 40% recovery). In the event the material recovers less than 40% of its elongation upon release of the applied force, the elongatable material will be considered to be “substantially non-elastic” or “substantially non-elastomeric”. For example, a material that has an initial length of 100mm can extend at least to 150mm, and upon removal of the force retracts to a length of 145mm (i.e., exhibiting a 10% recovery).

The term “elastic strand” or “elastic member” refers to a ribbon or strand (i.e. great length compared to either width and height or diameter of its cross-section) as may be part of the inner or outer cuff gathering component of an article.

The term “fiber” refers to any type of artificial fiber, filament, or fibril, whether continuous or discontinuous, produced through a spinning process, a meltblowing process, a melt fibrillation or film fibrillation process, or an electrospinning production process, or any other suitable process.

The term “film” refers to a polymeric material, having a skin-like structure, and it does not comprise individually distinguishable fibers. Thus, “film” does not include a nonwoven material. For purposes herein, a skin-like material may be perforated, apertured, or micro-porous and still be deemed a “film.”

The term “grommet ring”, or “grommet”, refers to a ring (not necessarily circular or oval) that is formed around the periphery of a mechanical bond site. FIG. 19 shows a bond site 351b with a bottom surface 351bb and a grommet ring 376.

The term “hydrophobic” refers to a material or composition having a contact angle greater than or equal to 90° according to The American Chemical Society Publication "Contact Angle, Wettability, and Adhesion," edited by Robert F. Gould and copyrighted in 1964. In certain embodiments, hydrophobic surfaces may exhibit contact angles greater than 120°, greater than 140°, or even greater than 150°. Hydrophobic liquid compositions are generally immiscible with water. The term “hydrophobic melt additive” refers to a hydrophobic composition that has been included as an additive to a hot melt composition (i.e., blended into a thermoplastic melt), which is then formed into fibers and/or a substrate (e.g., by spunbonding, meltblowing, or extruding).

The term “hydrophobic surface coating” refers to a composition that has been applied to a surface in order to render the surface hydrophobic or more hydrophobic. “Hydrophobic surface coating composition” means a composition that is to be applied to a surface in order to provide a hydrophobic surface coating.

The term “local basis weight variation” is defined by the Local Basis Weight Variation Test set forth below. Local basis weight variation is set forth in percentage.

The term “low surface tension fluid” refers to a fluid having a surface tension of less than 45 mN/m.

The term “low surface tension fluid strikethrough time” is defined by the Low Surface Tension Fluid Strikethrough Time Test set forth below. Low Surface Tension Fluid Strikethrough Time is set forth in seconds.

The term “machine direction” (MD) refers to the direction of material flow through a process.

The term “mass-average diameter” refers to a mass-weighted arithmetic mean diameter of fibers calculated from the fiber diameter, which is measured by the Fiber Diameter and Denier Test set forth below. Mass-average diameter of fibers is calculated by the Fiber Diameter Calculations set forth below. The mass-average diameter of fibers is set forth in microns.

The term “mean-flow pore diameter” in a nonwoven sample refers to a pore diameter corresponding to pressure at which the flow through pores in a “wet sample” is 50% of the flow

through pores in a “dry sample”. The mean flow pore diameter is measured by the Pore Size Distribution Test set forth below. The mean-flow pore diameter is such that the 50% of flow is through pores larger than the mean-flow pore diameter, and the rest of the flow is through the pores smaller than the mean-flow pore diameter. The mean-flow pore diameter is set forth in microns.

The term “calender bond” or “thermal bond” refers to a bond formed between fibers of a nonwoven by pressure and temperature such that the polymers within the bond melt together to form a continuous film-like material. The term “calendar bond” does not comprise a bond formed using an adhesive nor through the use of pressure only as defined by mechanical bond below. The term “thermal bonding” or “calender bonding” refers to the process used to create the thermal bond.

The term “mechanical bond” refers to a bond formed between two materials by pressure, ultrasonic attachment, and/or other mechanical bonding process without the intentional application of heat. The term mechanical bond does not comprise a bond formed using an adhesive.

The term “mechanical bonding” refers to the process used to create a mechanical bond.

As used herein, the term “nonwoven” means a porous, fibrous material made from continuous (long) filaments (fibers) and/or discontinuous (short) filaments (fibers) by processes such as, for example, spunbonding, meltblowing, carding, and the like. “Nonwoven” does not include a film, woven cloth, or knitted cloth..

The term “nonwoven component layer” refers to one sheet, ply or layer of a web of material.

The term “number-average diameter,” alternatively “average diameter”, refers to an arithmetic mean diameter of fibers calculated from the fiber diameter, which is measured by the Fiber Diameter and Denier Test set forth below. Number-average diameter of fibers is calculated by the Fiber Diameter Calculations set forth below. The number-average diameter of fibers is set forth in microns.

The term “polydispersity” refers to a measure of the width of a distribution calculated by a ratio of the mass-average diameter to the number-average diameter.

The term “porosity” refers to a measure of void volume of the nonwoven layer with the fibers composed of a material, and is calculated as $(1 - [\text{basis weight}]/[\text{thickness} \times \text{material density}])$ with the units adjusted so that they cancel out.

The term “relative standard deviation” (RSD) refers to a measure of precision calculated by dividing the statistic standard deviation for a series of measurements by the statistic average measurement of the series of measurements. This is often also called coefficient of variation or COV.

The terms “web” or “web of material” refer to a sheet-like structure such as a nonwoven or a film.

Nonwoven webs of material, such as nonwoven fabric webs, may comprise sheets of individual nonwoven component layers bonded together using mechanical, thermal, or chemical bonding processes. Nonwoven webs may be formed as flat, porous sheets made directly from individual fibers, from molten plastic, and/or plastic film. Some nonwoven structures may be strengthened or reinforced by a backing sheet, for example. Nonwoven structures may be engineered fabrics that may be a limited life, single-use fabric, or a very durable fabric. In various embodiments, nonwoven webs provide specific functions, such as absorbency, liquid repellency, resilience, stretch, softness, strength. These properties are often combined to create fabrics suited for specific applications, while achieving a good balance between product useful life and cost.

Continuous and discontinuous fiber spinning technologies of molten materials and typically of thermoplastics are commonly referred to as spinnmelt technologies. Spinnmelt technologies may comprise both the meltblowing process and spunbonding processes. A spunbonding process comprises supplying a molten polymer, which is then extruded under pressure through a large number of orifices in a plate known as a spinneret or die. The resulting continuous fibers are quenched and drawn by any of a number of methods, such as slot draw systems, attenuator guns, or Godet rolls, for example. In the spunlaying or spunbonding process, the continuous fibers are collected as a loose web upon a moving foraminous surface, such as a wire mesh conveyor belt, for example. When more than one spinneret is used in line for forming a multi-layered web, the subsequent nonwoven component layers are collected upon the uppermost surface of the previously formed nonwoven component layer.

The meltblowing process is related to the spunbonding process for forming a layer of a nonwoven material, wherein, a molten polymer is extruded under pressure through orifices in a spinneret or a die. High velocity gas impinges upon and attenuates the fibers as they exit the die. The energy of this step is such that the formed fibers are greatly reduced in diameter and are fractured so that micro-fibers of indeterminate length are produced. This differs from the

spunbonding process where the continuity of the fibers are generally preserved. Often meltblown nonwoven structures are added to spunbond nonwoven structures to form spunbond, meltblown (“SM”) webs or spunbond, meltblown, spunbond (“SMS”) webs, which are strong webs with some barrier properties.

Other methods to produce fine fibers comprise melt fibrillation and electrospinning. Melt fibrillation is a general class of making fibers defined in that one or more polymers are molten and are extruded into many possible configurations (e.g., co-extrusion, homogeneous or bicomponent films or filaments) and then fibrillated or fiberized into filaments. Meltblowing is one such specific method (as described herein). Melt film fibrillation is another method that may be used to produce submicron fibers. A melt film is produced from the melt and then a fluid is used to form fibers from the melt film. Examples of this method comprise U.S. Patent Nos. 6,315,806, 5,183,670, and 4,536,361, to Torobin et al., and U.S. Patent Nos. 6,382,526, 6,520,425, and 6,695,992, to Reneker et al. and assigned to the University of Akron. The process according to Torobin uses one or an array of co-annular nozzles to form a tube of film which is fibrillated by high velocity air flowing inside this annular film. Other melt film fibrillation methods and systems are described in the U.S. Pat. Publ. No. 2008/0093778, to Johnson, et al., published on April 24, 2008, U.S. Pat. No. 7,628,941, to Krause et al., and U.S. Pat. Publ. No. 2009/0295020, to Krause, et al., published on December 3, 2009 and provide uniform and narrow fiber distribution, reduced or minimal fiber defects such as unfiberized polymer melt (generally called “shots”), fly, and dust, for example. These methods and systems further provide uniform nonwoven webs for absorbent hygiene articles.

Electrospinning is a commonly used method of producing sub-micron fibers. In this method, typically, a polymer is dissolved in a solvent and placed in a chamber sealed at one end with a small opening in a necked down portion at the other end. A high voltage potential is then applied between the polymer solution and a collector near the open end of the chamber. The production rates of this process are very slow and fibers are typically produced in small quantities. Another spinning technique for producing sub-micron fibers is solution or flash spinning which utilizes a solvent.

There is a distinct difference between submicron diameter fibers made with electrospinning versus those made with melt-fibrillation, namely the chemical composition. Electrospun submicron fibers are made of generally soluble polymers of lower molecular weight than the fibers made by melt-fibrillation. Commercially-viable electrospinning methods have been

described in U.S. Pat. No. 7,585,437, to Jirsak et al., U.S. Pat. No. 6,713,011 to Chu et al., and U.S. Pat. Publ. No. 2009/0148547, to Petras et al. Electro-spinning is recently explored in combination with a molten polymer rather than a polymer solution, as described in a reference by Lyons et al., "Melt-electrospinning Part I: Processing Parameters and Geometric Properties", published in the journal POLYMER 45 (2004) pp. 7597-7603; and by Zhou et al., "The Thermal Effects on Electrospinning of Polylactic Acid Melts", published in the journal POLYMER 47 (2006) pp. 7497-7505. The researchers in these studies have observed that electrospun fibers have average diameters generally greater than 1 micron as compared to solution electrospun fibers that are submicron (i.e., less than 1 micron). With motivation to reduce the fiber diameter, researchers have more recently started optimizing process and polymer parameters. Generally, the goal of the researchers has been to reduce the number-average diameter, but not reduce the mass-average diameter, and narrow the fiber diameter distribution. Improvements in melt electrospinning show that fiber diameter may be decreased, though to a limited extent but still above 1 micron (generally, in the range of 2 micron to 40 micron for polypropylene with molecular weights in the range of 12,000 to 200,000 Daltons) by the research works of Kong et al., "Effects of the Spin Line temperature Profile and Melt Index of Poly(propylene) on Melt-electrospinning", published in the journal POLYMER ENGINEERING AND SCIENCE 49 (2009) pp. 391-396 (average fiber diameter of 20 micron using polypropylene of melt flow index of 1500); by Kadomae et al., "Relation Between Tacticity and Fiber Diameter in Melt-electrospinning of Polypropylene", published in the journal FIBERS AND POLYMERS 10 (2009) pp. 275-279 (fiber diameters in the range of 5-20 microns using polypropylene with 12,000 and 205,000 molecular weight), and by Yang et al., "Exploration of Melt-electrospinning Based on the Novel Device", published in the Proceedings of the IEEE International Conference on Properties and Applications of Dielectric Materials, 2009, pp. 1223-1226 (finest fiber diameter of 5 micron). Most recently, the melt electrospinning has been modeled by Zhmayev et al., "Modeling of Melt Electrospinning for Semi-crystalline Polymers", published in the journal POLYMER 51 (2010) pp. 274-290. Even their models show that the fiber diameter of melt electrospun Nylon 6 (with a melt flow index of 3) is 2 microns, similar to that obtained by experiments. A prior work by Dalton et al., "Electrospinning of Polymer Melts: Phenomenological Observations", showed that fiber diameter of melt electrospun high molecular weight polypropylene fibers (with MFI in the range of 15 cm³/10 min to 44 cm³/10 min) may be significantly reduced to submicron by adding 1.5% of

viscosity reducing additive, such as Irgatec CR 76 (from Ciba Specialty Chemicals, Switzerland). However, viscosity reducing additives, such as Irgatec CR 76, for example, significantly reduce the molecular weight of the polymer, as described in U.S. Pat. No. 6,949,594 to Roth et al., and by Gande et al., "Peroxide-free Vis-breaking Additive for Improved Qualities in Meltblown Fabrics", in the conference proceedings of the International Nonwovens Technical Conference, 2005, St. Louis, Missouri, USA. Therefore, melt electrospun fibers have fiber diameters generally above 1 micron, or a high standard deviation leading to a broad fiber diameter distribution using commercial-grade high molecular weight polymers. Also, the polymer used in successful electrospinning of polymer melts uses a polymer of low molecular weight, e.g., in the case of PLA starting from 186,000 Dalton and degrading to actually 40,000 Dalton in the spun fibers used by Zhou et al., and use of viscosity reducing additive Irgatec CR 76 by Dalton et al. to reduce the melt viscosity by reducing the molecular weight. This compares to PLA used in melt-fibrillation processes of where e.g. the Natureworks 6202D resin starts at a molecular weight Mw of 140,000 Dalton and 'degrades' only to a molecular weight of 130,000 to 135,000 Dalton compared to the 40,000 of the melt-electrospun fibers. Also other grades of PLA (e.g. with Mw of 95,000 or 128,000) drop in molecular weight from neat resin to fiber form by less 10,000 or even less than 1,000 Dalton (less than 10% or less than 1%). Therefore, not only is the electrospinning process including the melt-electrospinning process at present still low in throughput, but it is structurally and chemically distinct from the fine fibers (i.e., the second nonwoven component layer) of the present disclosure. However, it is desirable to develop the electrospinning method towards making fine fibers at higher throughput and a narrow submicron diameter distribution as described herein.

In various embodiments, the fibers of the nonwoven structure may be made of polyesters, including PET and PBT, polylactic acid (PLA), and alkyds, polyolefins, including polypropylene (PP), polyethylene (PE), and polybutylene (PB), olefinic copolymers from ethylene and propylene, elastomeric polymers including thermoplastic polyurethanes (TPU) and styrenic block-copolymers (linear and radial di- and tri-block copolymers such as various types of Kraton), polystyrenes, polyamides, PHA (polyhydroxyalkanoates) and e.g. PHB (polyhydroxybutyrate), and starch-based compositions including thermoplastic starch, for example. The above polymers may be used as homopolymers, copolymers, e.g., copolymers of ethylene and propylene, blends, and alloys thereof.

A variety of mass-produced consumer products such as diapers, paper towels, feminine care products, incontinence products and similar materials, employ nonwoven webs, such as SMS webs, in their manufacture. One of the largest users of SM and SMS webs is the disposable diaper and feminine care products industry. When the nonwoven webs are incorporated in an absorbent article, however, achieving a barrier against fluids that have a surface tension on a similar level of the surface energy of the SMS structure is sometimes difficult. For example, some SMS webs may have a surface energy level of approximately 30 mN/m, e.g., when made of PP, while the fluids sought to be blocked (i.e., infant urine or runny feces) may have surface tensions of 40-50 mN/m, or in some cases as low as 32 to 35 mN/m. For various components of absorbent articles, such as barrier leg cuffs, for example, in order to achieve a desired fluid barrier, hydrophobic surface coatings may be applied to the webs or hydrophobic melt-additives may be used in the production of the nonwoven webs. Such techniques, however, may add to the production costs associated with the absorbent product and generally increase the production complexity. If hydrophilic surfactants or materials are used on other portions of the absorbent article (such as for example the topsheet), they may migrate or wash off toward other absorbent article components during wet and/or dry conditions. During dry conditions, for example, the hydrophilic surfactants or materials may migrate after absorbent articles are manufactured and packaged and while being stored over the course of weeks and attach to the barrier cuff, thereby, possibly leading to an increased leakage rate. In addition, during wet conditions, the hydrophilic surfactants or materials may also wash off of a diaper topsheet, for example, and then attach to the barrier cuffs, thereby, again possibly leading to an increased leakage rate. One advantage of the additional hydrophobic materials in the web is that they resist and repel the hydrophilic surfactants. Therefore, it would be desirable to combine that advantage without the additional complexities and costs.

Further to the above, a number of undesirable holes extending through the nonwoven webs, such as SMS webs, for example, may be created during the mechanical bonding process of various structures. Current equipment and processes are not sufficient to bond combinations of SMS and spunbond (S, SS, SSS) materials at total basis weights below 25 gsm using a pressure/shear bonding without an increase in the number of holes created by the process. Holes are created from the bonding nub punching through thin areas of the SMS or SS web. Increased holes through the bonded materials result in higher product failure rates (i.e., leakage). When an

absorbent article that incorporates such a nonwoven web is subsequently worn by a user, the presence of the holes may result in undesirable leaks.

In view of the above, low cost nonwoven webs having low basis weights, adequate air permeability, (i.e., breathable), adequate tactile characteristics, and low surface tension fluid strikethrough times exceeding certain parameters are desired. It is also desirable for the nonwoven materials to have more structural uniformity (i.e., less local basis weight variation), especially at lower basis weights (e.g., less than 25 gsm, alternatively, less than 15 gsm, alternatively, less than 13 gsm, and, alternatively, less than 10 gsm). An increased structural uniformity in nonwoven webs of 25 gsm or less reduces the amount of defects (e.g., holes) created during mechanical bonding processes. With specific regard to barrier cuff materials, in one embodiment, it is desired to have soft low basis weight webs with an improved barrier against low surface tension body exudates to give the absorbent core more time to absorb the fluid, especially with recent and future trend of more “body-fitting” diaper designs and thinner absorbent cores.

As described in more detail below, a nonwoven component layer having fine fibers (“N-fibers”) with an average diameter of less than 1 micron (an “N-fiber layer”) may be added to, or otherwise incorporated with, other nonwoven component layers to form a nonwoven web of material. In some embodiments, the N-fiber layer may be used to produce a SNS nonwoven web or SMNS nonwoven web, for example. The N-fibers may be comprised of a polymer, e.g., selected from polyesters, including PET and PBT, polylactic acid (PLA), alkyds, polyolefins, including polypropylene (PP), polyethylene (PE), and polybutylene (PB), olefinic copolymers from ethylene and propylene, elastomeric polymers including thermoplastic polyurethanes (TPU) and styrenic block-copolymers (linear and radial di- and tri-block copolymers such as various types of Kraton), polystyrenes, polyamides, PHA (polyhydroxyalkanoates) and e.g. PHB (polyhydroxybutyrate), and starch-based compositions including thermoplastic starch, for example. The above polymers may be used as homopolymers, copolymers, e.g., copolymers of ethylene and propylene, blends, and alloys thereof. The N-fiber layer may be bonded to the other nonwoven component layers by any suitable bonding technique, such as the calender bond process, for example, also called thermal point bonding.

In some embodiments, the use of an N-fiber layer in a nonwoven web may provide a low surface tension barrier that is as high as other nonwoven webs that have been treated with a hydrophobic coating or a hydrophobic melt-additive, and still maintain a low basis weight (e.g.,

less than 15 gsm or, alternatively, less than 13 gsm). The use of the N-fiber layer may also provide a soft and breathable (i.e., air permeable) nonwoven material that, at least in some embodiments, may be used in single web layer configurations in applications which previously used double web layer configurations. Furthermore, in some embodiments, the use of the N-fiber layer may at least reduce the undesirable migration of hydrophilic surfactants toward the web and, therefore, may ultimately result in better leak protection for an associated absorbent article. Also, when compared to an SMS web having a similar basis weight, the use of a nonwoven web comprising the N-fiber layer may decrease the number of defects (i.e., holes or pinholes through the mechanical bond site) created during the mechanical bonding process.

Without intending to be bound by any particular theory, with regard to fluid barrier characteristics of the webs disclosed herein, it is believed that the small size of the pores created in the web by the use of the N-fiber layer along with the tightness or proximity of the fibers may increase the hydrostatic pressure required to penetrate through the pores for low surface tension fluids and potentially increase capillary drag forces. The fine pores may increase the capillary drag forces applied to a low surface tension fluid passing through the fine pores of the web to slow down low surface tension fluid strikethrough. Further, it is found that multiple aspects of the pore structure are relevant, more than the average pore size, such as, for example, the narrowness of the pore size distribution, mean-flow pore size, and modes of pore size distribution.

As discussed in more detail below, the webs of materials incorporating the N-fiber layer may be used in the construction of various absorbent articles. In one embodiment, the absorbent articles of the present disclosure may comprise a liquid pervious topsheet, a backsheet attached or joined to the topsheet, and an absorbent core disposed between the topsheet and the backsheet. Absorbent articles and components thereof, including the topsheet, backsheet, absorbent core, and any individual layers of these components, generally have an interior surface (or wearer-facing surface) and an exterior surface (or garment-facing surface).

The following description generally discusses a suitable absorbent core, a topsheet, and a backsheet that may be used in absorbent articles, such as disposable diapers, for example. It is to be understood that this general description applies to the components of the specific absorbent article shown in FIGS. 1, 2, and 3A-3B, which are further described below, and to other absorbent articles which are described herein.

FIG. 1 is a plan view of an absorbent article 10 in accordance with one non-limiting embodiment of the present disclosure. The absorbent article 10 is illustrated in its flat, uncontracted state (i.e., with its elastic induced contraction removed for illustration and with portions of the absorbent article 10 being cut-away to more clearly show the construction of the absorbent article 10. A portion of the absorbent article 10 which faces away from the wearer is oriented towards the viewer. FIG. 2 is a perspective view of the absorbent article 10 of FIG.1 in a partially contracted state. As shown in FIG. 1, the absorbent article 10 may comprise a liquid pervious first topsheet 20, a liquid impervious backsheet 30 joined with the topsheet 20, and an absorbent core 40 positioned between the topsheet 20 and the backsheet 30. The absorbent core 40 has an exterior surface (or garment-facing surface) 42, an interior surface (or a wearer-facing surface) 44, side edges 46, and waist edges 48. In one embodiment, the absorbent article 10 may comprise gasketing barrier cuffs 50 and longitudinal barrier cuffs 51. The longitudinal barrier cuffs 51, in some embodiments, may extend generally parallel to a central longitudinal axis 59. For example, the longitudinal barrier cuffs 51 may extend substantially between the two end edges 57. The absorbent article 10 may comprise an elastic waist feature multiply designated as 60 (also referred to herein as a waistband or a belt) and a fastening system generally multiply designated as 70.

In one embodiment, the absorbent article 10 may have an outer surface 52, an inner surface 54 opposed to the outer surface 52, a first waist region 56, a second waist region 58, and a periphery 53 which is defined by longitudinal edges 55 and the end edges 57. (While the skilled artisan will recognize that an absorbent article, such as a diaper, is usually described in terms of having a pair of waist regions and a crotch region between the waist regions, in this application, for simplicity of terminology, the absorbent article 10 is described as having only waist regions comprising a portion of the absorbent article which would typically be designated as part of the crotch region). The inner surface 54 of the absorbent article 10 comprises that portion of the absorbent article 10 which is positioned adjacent to the wearer's body during use (i.e., the inner surface 54 is generally formed by at least a portion of the first topsheet 20 and other components that may be joined to the topsheet 20). The outer surface 52 comprises that portion of the absorbent article 10 which is positioned away from the wearer's body (i.e., the outer surface 52 is generally formed by at least a portion of the backsheet 30 and other components that may be joined to the backsheet 30). The first waist region 56 and the second

waist region 58 extend, respectively, from the end edges 57 of the periphery 53 to the lateral centerline (cross-sectional line 3-3) of the absorbent article 10.

FIG. 2 shows a perspective view of the absorbent article 10 which comprises a pair of longitudinal barrier cuffs 51 in accordance with one non-limiting embodiment of the present disclosure. FIG. 3 depicts a cross-sectional view taken along line 3--3 of FIG. 1.

In one embodiment, the absorbent core 40 may take on any size or shape that is compatible with the absorbent article 10. In one embodiment, the absorbent article 10 may have an asymmetric, modified T-shaped absorbent core 40 having a narrowing of the side edge 46 in the first waist region 56, but remaining generally rectangular-shaped in the second waist region 58. Absorbent core construction is generally known in the art. Various absorbent structures for use as the absorbent core 40 are described in U.S. Pat. Nos. 4,610,678, issued to Weisman et al., on September 9, 1986, 4,673,402, issued to Weisman, et al., on June 16, 1987, 4,888,231, issued to Angstadt, on December 19, 1989, and 4,834,735, issued to Alemany et al., on May 30, 1989. In one embodiment, the absorbent core 40 may comprise a dual core system containing an acquisition/distribution core of chemically stiffened fibers positioned over an absorbent storage core as described in U.S. Pat. Nos. 5,234,423, issued to Alemany, et al., on August 10, 1993, and 5,147,345, issued to Young et al., on September 15, 1992. The absorbent core 40 may also comprise a core cover 41 (as shown in FIGS. 3A-B and as described in detail below) and a nonwoven dusting layer that is disposed between the absorbent core 40 and the backsheet 30.

In one embodiment, the topsheet 20 of the absorbent article 10 may comprise a hydrophilic material that promotes rapid transfer of fluids (e.g., urine, menses, and/or runny feces) through the topsheet 20. The topsheet 20 may be pliant, soft feeling, and non-irritating to the wearer's skin. Further, the topsheet may be fluid pervious, permitting fluids (e.g., menses, urine, and/or runny feces) to readily penetrate through its thickness. In one embodiment, the topsheet 20 may be made of a hydrophilic material or at least the upper surface of the topsheet may be treated to be hydrophilic so that fluids will transfer through the topsheet more rapidly and enter the absorbent core 40. This diminishes the likelihood that body exudates will flow off of the topsheet 20 rather than being drawn through the topsheet 20 and being absorbed by the absorbent core 40. The topsheet 20 may be rendered hydrophilic by treating it with a surfactant, for example. Suitable methods for treating the topsheet 20 with a surfactant comprise spraying the topsheet 20 with the surfactant and immersing the topsheet 20 into the surfactant. A more

detailed discussion of such a treatment is contained in U.S. Pat. Nos. 4,988,344, issued to Reising, on Jan. 29, 1991, and 4,988,345, issued to Reising, on Jan. 29, 1991.

In one embodiment, the backsheet 30 may be impervious, or at least partially impervious, to low surface tension fluids (e.g., menses, urine, and/or runny feces). The backsheet 30 may be manufactured from a thin plastic film, although other flexible fluid impervious materials may also be used. The backsheet 30 may prevent, or at least inhibit, the exudates absorbed and contained in the absorbent core 40 from wetting articles which contact the absorbent article 10, such as bedsheets, clothing, pajamas, and undergarments, for example. The backsheet 30 may comprise a woven or a nonwoven web, polymeric films such as thermoplastic films of polyethylene or polypropylene, and/or composite materials such as a film-coated nonwoven material or a film-nonwoven laminate. In one embodiment, a suitable backsheet 30 may be a polyethylene film having a thickness of from 0.012 mm (0.5 mils) to 0.051 mm (2.0 mils). Exemplary polyethylene films are manufactured by Clopay Corporation of Cincinnati, Ohio, under the designation P18-1401 and by Tredegar Film Products of Terre Haute, Ind., under the designation XP-39385. The backsheet 30 may be embossed and/or matte finished to provide a more cloth-like appearance. Further, the backsheet 30 may permit vapors to escape from the absorbent core 40 (i.e., the backsheet 30 is breathable and has an adequate air permeability), while still preventing exudates from passing through the backsheet 30. The size of the backsheet 30 may be dictated by the size of the absorbent core 40 and the exact absorbent article design selected. In one embodiment, the backsheet 30 may comprise an SNS and/or an SMNS web, as described in greater detail below.

Other optional elements of the absorbent article 10 may comprise a fastening system 70, elasticized side panels 82, and a waist feature 60. The fastening system 70 allows for the joining of the first waist region 56 and the second waist region 58 in an overlapping configuration such that lateral tensions are maintained around the circumference of the absorbent article 10 to maintain the absorbent article 10 on the wearer. Exemplary fastening systems 70 are disclosed in U.S. Pat. Nos. 4,846,815, issued to Scripps, on July 11, 1989, 4,894,060, issued to Nestegard, on January 16, 1990, 4,946,527, issued to Battrell, on August 7, 1990, 3,848,594, issued to Buell, on November 19, 1974, 4,662,875, issued to Hirotsu et al., on May 5, 1987, and 5,151,092, issued to Buell et al., on September 29, 1992. In certain embodiments, the fastening system 70 may be omitted. In such embodiments, the waist regions 56 and 58 may be joined by the absorbent article manufacturer to form a pant-type diaper having a preformed waist opening

and leg openings (i.e., no end-user manipulation of the diaper is needed to form the waist opening and leg openings). Pant-type diapers are also commonly referred to as “closed diapers,” “prefastened diapers,” “pull-on diapers,” “training pants,” and “diaper-pants”. Suitable pants are disclosed in U.S. Pat. Nos. 5,246,433, issued to Hasse et al., on September 21, 1993, 5,569,234, issued to Buell et al., on October 29, 1996, 6,120,487, issued to Ashton, on September 19, 2000, 6,120,489, issued to Johnson et al., on September 19, 2000, 4,940,464, issued to Van Gompel et al., on July 10, 1990, and 5,092,861, issued to Nomura et al., on March 3, 1992. Generally, the waist regions 56 and 58 may be joined by a permanent or refastenable bonding method.

In certain embodiments, the absorbent article 10 may comprise at least one barrier member. In one embodiment, barrier members are physical structures joined to, applied to, and/or formed with the absorbent article 10 to improve the barrier properties of the absorbent article 10. In one embodiment, barrier members may comprise structures such as a core cover, an outer cover, a longitudinal barrier cuff, a gasketing cuff, an elasticized topsheet, and combinations thereof. It may be desirable that a barrier member comprise the SNS web and/or the SMNS web, as described in further detail below.

In one embodiment, the absorbent article 10 may comprise one or more longitudinal barrier cuffs 51 which may provide improved containment of fluids and other body exudates. The longitudinal barrier cuffs 51 may also be referred to as leg cuffs, barrier leg cuffs, longitudinal leg cuffs, leg bands, side flaps, elastic cuffs, or “stand-up” elasticized flaps. Elasticity may be imparted to the longitudinal barrier cuffs 51 by one or more elastic members 63. Elastic members 63 may provide elasticity to the longitudinal barrier cuff 51 and may aid in keeping longitudinal barrier cuff 51 in a “stand-up” position. U.S. Pat. No. 3,860,003, issued to Buell, on July 14, 1975, describes a disposable diaper that provides a contractible leg opening having a side flap and one or more elastic members to provide an elasticized leg cuff. U.S. Pat. Nos. 4,808,178 and 4,909,803 issued to Aziz et al. on Feb. 28, 1989 and Mar. 20, 1990, respectively, describe absorbent articles comprising “stand-up” elasticized flaps that improve the containment at the leg regions of the absorbent article 10. Additionally, in some embodiments, the one or more longitudinal barrier cuffs 51 may be integral with one or more gasketing cuffs 50. For example, the longitudinal barrier cuffs 51 and the gasketing cuffs 50 may be formed from a single web of material, as illustrated in FIGS. 3A-3B. As with the longitudinal barrier cuffs 51, the gasketing cuffs 50 may comprise one or more elastic members 62.

FIGS. 3A-B shows a cross-sectional view of the absorbent article 10 of FIG. 1 taken along line 3--3. FIGS. 3A-B depict various cuff constructions; however, modifications may be made to the cuff construction without departing from the spirit and scope of the present disclosure. A gasketing cuff 50 and a longitudinal barrier cuff 51 are both shown in FIGS. 3A-B, but a single cuff design is equally feasible. FIG. 3A illustrates a gasketing cuff 50 and a longitudinal barrier cuff 51 construction in accordance with one non-limiting embodiment. Both cuffs 50, 51 may share a common web 65, such as an SNS web or an SMNS web, for example. The longitudinal barrier cuff 51 is shown in a single layer configuration where over a substantial portion of the lateral width of the longitudinal barrier cuff 51 comprises a single ply of the web 65. FIG. 3B illustrates a gasketing cuff 50 and longitudinal barrier cuff 51 construction with the longitudinal barrier cuff 51 in a multiple layer configuration in accordance with another non-limiting embodiment. In the multiple layer construction, at least two plies of the web (such as an SNS web or an SMNS web, for example) exist over a substantial portion of the lateral width of the longitudinal barrier cuff 51. Those of skill in the art will recognize that the exact configuration of the web 65 may be altered in various embodiments.

A variety of suitable materials may be used as the web 65 in the cuffs described above. Suitable embodiments may have the web 65 comprising a plurality of layers, such as two spunbond layers and at least one N-fiber layer disposed between the two spunbond layers, for example, as described in greater detail below. Some embodiments of the web 65 may comprise a hydrophobic material, as described in greater detail below.

As shown in FIGS. 3A-B, a core cover 41 may be included in certain embodiments of the absorbent article 10 to provide structural integrity to the absorbent core 40. The core cover 41 may contain the absorbent core 40 components such as cellulosic material and absorbent gelling material, which both may tend to migrate, move, or become airborne without a physical barrier. The core cover 41 may entirely envelop the core 40, as shown in FIGS. 3A-B, or may partially cover the absorbent core 40. The core cover 41 may generally comprise a nonwoven web. In certain embodiments, the core cover 41, or other components of the absorbent article 10, may comprise an SNS web and/or an SMNS web.

In certain embodiments, the absorbent article 10 may comprise an outer cover 31. The outer cover 31 may cover all of, or substantially all of, the exterior surface of the absorbent article 10. In some embodiments, the outer cover 31 may be coterminous with the backsheet 30. The outer cover 31 may be bonded to a portion of the backsheet 30 to form a laminate structure.

Bonding may be performed by any conventional methods, such as adhesive bonding, mechanical bonding, and thermal bonding, for example. The outer cover 31 may be utilized to provide extra strength or bulk to the absorbent article 10. Outer covers 31 are often used to improve the aesthetic quality of the exterior surface of the absorbent article 10. It is also desirable that the exterior surface of the absorbent article 10 exhibit a cloth-like look and feel, as such features are pleasing to consumers. Various materials are suitable for use as the outer cover 31. Such materials comprise woven webs, foams, scrims, films, and loose fibers. However, in certain embodiments, the outer cover 31 may be constructed to provide increased barrier protection. In certain embodiments, the outer cover 31 may comprise an SNS web and/or an SMNS web.

FIG. 4 shows a schematic diagram of a forming machine 110 used to make a nonwoven web 112, such as an SNS web or an SMNS web, for example, in accordance with one embodiment. To make an SMNS web, the forming machine 110 is shown as having a first beam 120 for producing first coarse fibers 135, an optional second beam 121 for producing intermediate fibers 127 (e.g., meltblown fibers), a third beam 122 for producing fine fibers 131 (e.g., N-fibers), and a fourth beam 123 for producing second coarse fibers 124. The forming machine 110 may comprise an endless forming belt 114 which travels around rollers 116, 118 so the forming belt 114 is driven in the direction as shown by the arrows 114. In various embodiments, if the optional second beam 121 is utilized, it may be positioned intermediate the first beam 120 and the third beam 122 (as illustrated), or may be positioned intermediate the third beam 122 and the fourth beam 124, for example.

In one embodiment, the first beam 120 may produce first coarse fibers 135, such as by use of a conventional spunbond extruder with one or more spinnerets which form continuous fibers of polymer. Forming spunbond fibers and the design of such a spunbond forming first beam 120 is within the ability of those of skill in the art. Spunbond machines may be acquired from Reicofil GmbH in Troisdorf, Germany, for example. Suitable thermoplastic polymers comprise any polymer suitable for spunbonding such as polyesters, including PET and PBT, polylactic acid (PLA), and alkyds, polyolefins, including polypropylene (PP), polyethylene (PE), and polybutylene (PB), olefinic copolymers from ethylene and propylene, elastomeric polymers including thermoplastic polyurethanes (TPU) and styrenic block-copolymers (linear and radial di- and tri-block copolymers such as various types of Kraton), polystyrenes, polyamides, PHA (polyhydroxyalkanoates) and e.g. PHB (polyhydroxybutyrate), and starch-based compositions including thermoplastic starch, for example. The above polymers may be

used as homopolymers, copolymers, e.g., copolymers of ethylene and propylene, blends, and alloys thereof. The polymer is heated to become fluid, typically at a temperature of 100-350°C, and is extruded through orifices in the spinneret. The extruded polymer fibers are rapidly cooled and attenuated by air streams to form the desired denier fibers. The first coarse fibers 135 resulting from the first beam 120 may be dispensed or laid onto the forming belt 114 to create a first nonwoven component layer 136. The first nonwoven component layer 136 may be produced from multiple beams or spinnerets of the type of the first beam 120, but still creates one nonwoven component layer when the fibers produced from the multiple beams or spinnerets are of the same diameter, shape, and composition. The first beam 120 may comprise one or more spinnerets depending upon the speed of the process or the particular polymer being used. The spinnerets of the first beam 120 may have orifices with a distinct shape that imparts a cross-sectional shape to the first coarse fibers 135. In one embodiment, the spinnerets may be selected to yield fibers with cross-sectional shapes including, but not limited to, circular, oval, rectangular, square, triangular, hollow, multi-lobal, irregular (i.e., nonsymmetrical), and combinations thereof.

In one embodiment, the second beam 121, if used, may produce intermediate diameter fibers 127, such as meltblown fibers, for example. The meltblown process results in the extrusion of a thermoplastic polymer through a die 119 containing a plurality of orifices. In some embodiments, the die 119 may contain from 20 to 100, or even more, orifices per inch of die width. As the thermoplastic polymer exits the die 119, high pressure fluid, usually hot air may attenuate and spread the polymer stream to form the intermediate fibers 127. The intermediate fibers 127 resulting from the second beam 121 may be dispensed or laid onto the first nonwoven component layer 136 carried by the forming belt 114, to create a fourth nonwoven component layer 128. The fourth nonwoven component layer 128 may be produced from multiple, adjacent beams of the type like the second beam 121.

In one embodiment, the third beam 122 may produce the fine fibers 131 (i.e., N-fibers). In some embodiments, the N-fibers may be produced using systems and melt film fibrillation methods described in U.S. Patent Nos. 6,315,806, 5,183,670, and 4,536,361, to Torobin et al., and U.S. Patent Nos. 6,382,526, 6,520,425, and 6,695,992, to Reneker et al. and assigned to the University of Akron. Other melt film fibrillation methods and systems are described in the U.S. Pat. Publ. No. 2008/0093778, to Johnson, et al., published on April 24, 2008, U.S. Pat. No. 7,628,941, to Krause et al., and U.S. Pat. Publ. No. 2009/0295020, to Krause, et al., published

on December 3, 2009 and provide uniform and narrow fiber distribution, reduced or minimal fiber defects such as unfiberized polymer melt (generally called “shots”), fly, and dust, and further provide uniform N-fibers layer 132 for absorbent articles, such as those described by the present disclosure. The improvements in the melt film fibrillation method, specifically the design of converging-diverging gas passage specifications and the fluid curtain, described by the Johnson et al. and Krause et al., respectively, may provide the N-fibers of desired structural attributes such as number-average fiber diameter distribution, mass-average fiber diameter distribution, pore-size distribution, and structural uniformity (i.e., less local basis weight variation) for the embodiments of the present disclosure as described herein. Generally, in one embodiment, a pressurized gas stream flows within a gas passage confined between first and second opposing walls, which define respective upstream converging and downstream diverging wall surfaces. A polymer melt is introduced into the gas passage to provide an extruded polymer film on the heated wall surfaces that is impinged by the gas stream flowing within the gas passage, effective to fibrillate the polymer film into sub-micron diameter fibers or fibers. The fine fibers 131 may then be dispensed or laid onto the first nonwoven component layer 136 to create the second nonwoven component layer 132. In some embodiments, such as during the production of an SMNS web, for example, the fine fibers 131 may be dispensed or laid onto the fourth nonwoven component layer 128, which is carried on the forming belt 114. Alternatively, in some embodiments, the fine fibers 131 may be laid onto the first nonwoven component layer 136 and subsequently the intermediate fibers 127, such as meltblown fibers, may be laid onto the layer of fine fibers 131. The fine fiber layer 132 may be produced from more than one beam of the type of the third beam 122.

In one embodiment, the fourth beam 123 (or multiple beams like 120) may produce the second coarse diameter fibers 124 that are similar to the first coarse fibers 135. The second coarse fibers 124 may be dispensed or laid onto the second nonwoven component layer 132 of the web 112, such as during the production of an SNS web, for example. The resulting web 112 may be fed through thermal bonding rolls 138, 140. The bonding rolls 138, 140 are commonly referred to as a calender. The surfaces of one or both of the bonding rolls 138, 140 may be provided with a raised pattern or portions such as spots, grids, pins, or nubs, for example. In one embodiment, the bonding rolls 138, 140 may be heated to the softening temperature of the polymer used to form the nonwoven component layers of the web 112. As the web 112 passes between the heated bonding rolls 138, 140, the nonwoven component layers may be embossed

by the bonding rolls 138, 140 in accordance with the pattern on the bonding rolls 138, 140 to create a pattern of discrete areas, such as calender bond 168 shown in FIG. 5. The discrete areas are bonded from nonwoven component layer to nonwoven component layer with respect to the particular fibers within each layer. Such discrete area, or calender bond site, may be carried out by heated rolls or by other suitable techniques. Another thermal fiber bonding technique comprises blowing hot air through the web 112. Air-through bonding techniques may generally be used with low melting point matrix fibers, biocomponent fibers, and powders. While a nonwoven web is described herein as comprising three to four nonwoven component layers, any suitable number of nonwoven component layers may be used and are within the scope of the present disclosure.

FIG. 5 illustrates a cross-sectional view of an SNS web at a calender bond site 168 in accordance with one non-limiting embodiment. A three layer nonwoven web 112 is illustrated that was produced by the forming machine 110 described above without the optional second beam 121 (e.g., the meltblown layer). The nonwoven web 112 may comprise a first nonwoven component layer 125 which itself may be comprised of coarse fibers, such as spunbond fibers, for example. In one embodiment, the first nonwoven component layer 125 may comprise fibers having an average diameter, alternatively, number-average diameter, in the range of 8 microns to 30 microns and, alternatively, in the range of 10 microns to 20 microns, with a relative standard deviation in the range of 4% to 10%. Stated another way, the first nonwoven component layer 125 may comprise fibers having an average denier in the range of 0.4 to 6.0, with a relative standard deviation in the range of 8% to 15%. The mass-average fiber diameter in the same embodiment may be in the range of 8 microns to 30 microns and, alternatively, in the range of 10 microns to 20 microns, with a relative standard deviation in the range of 4% to 10%. In one embodiment, the first nonwoven component layer 125 may have a basis weight in the range of 1 gsm to 10 gsm and, alternatively, in the range of 2 gsm to 7 gsm, e.g., 5.5 gsm. In certain embodiments, the fibers in the first nonwoven component layer 125 may have non-circular cross-sections, such as trilobal cross-sections, for example, or may be bicomponent fibers, such as sheath-core or side by side, for example.

In one embodiment, the nonwoven web 112 may comprise a second nonwoven component layer 132 which itself may be comprised of fine fibers, such as N-fibers. In one embodiment, the second nonwoven component layer 132 may comprise fine fibers having a number-average diameter (alternatively “average diameter”) less than 1 micron, alternatively, in

the range of 0.1 microns to 1 micron, alternatively in the range of 0.2 microns to 0.9 microns, alternatively in the range of 0.3 microns to 0.8 microns and, alternatively, in the range of 0.5 microns to 0.7 microns, with a relative standard deviation of less than 100%, alternatively less than 80%, alternatively less than 60%, alternatively less than 50%, such as in the range of 10% to 50%, for example; and with over 80%, such as over 90%, or 95 to 100%, for example, of the fibers having less than 1 micron diameter, i.e. submicron. The mass-average diameter of fibers in the second nonwoven component layer 132 may be less than 2 microns, alternatively, in the range of 0.1 micron to 2 microns, alternatively, in the range of 0.1 microns to 1.5 microns, alternatively, in the range of 0.1 microns to 1 micron, alternatively, in the range of 0.2 microns to 0.9 microns, alternatively, in the range of 0.3 microns to 0.8 microns and, alternatively, in the range of 0.5 microns to 0.7 microns, with a relative standard deviation of less than 100%, alternatively less than 80%, alternatively less than 60%, alternatively less than 50%, such as in the range of 10% to 50%, for example. Stated another way, the second nonwoven component layer 132 may comprise fine fibers having an average denier in the range of 0.00006 to 0.006, alternatively, in the range of 0.0002 to 0.005, alternatively, in the range of 0.0016 to 0.005, and alternatively, in the range of 0.002 to 0.004, with a relative standard deviation in the range of less than 200%, alternatively, less than 150%, and alternatively, less than 120%; and with over 80%, alternatively, over 90%, and alternatively, 95 to 100% of the fibers less than 0.006 denier.

In an embodiment with the mass-average fiber distribution of less than 1 micron, almost all the fibers must have a diameter less than 1 micron. Even with very few fibers above 1 micron, it would make the mass-average fiber diameter greater than 1 micron. Thicker fibers have larger mass; thus, the presence of thicker fibers with larger mass increases the mass-average fiber diameter more than the number-average fiber diameter as described in the Fiber Diameter Calculations set forth below. For example, a fiber with a diameter of 3 microns (a typical meltblown fiber) has 36 times more mass than a submicron N-fiber of the same length and with a typical diameter of 0.5 microns because the 3 micron fiber has a cross-sectional area 36 times larger than that of a 0.5 micron diameter fiber. Alternatively, a single 3 micron fiber diameter fiber may take the place of 36 fibers of 0.5 micron diameter, and increase the mass-average fiber diameter of the second component layer. Conversely, to reduce the mass-average fiber diameter, it is critical to reduce the number of fibers with diameter greater than 1 micron. In one embodiment, the second nonwoven component layer may comprise fibers having a number-average diameter of less than 1 micron, a mass-average diameter of less than 1.5

microns, and a ratio of the mass-average diameter to the number-average diameter less than 2. In some embodiments, the second nonwoven component layer may comprise fibers having a number-average diameter of less than 1 micron, a mass-average diameter of less than 1 micron, and a ratio of the mass-average diameter to the number-average diameter less than 1.5, for example.

Without intending to be bound by any particular theory, it is believed that the finer fibers make finer pores in the nonwoven web. As set forth herein, the finer pores provide greater fluid strikethrough performance of the nonwoven web. Therefore, it is desirable to have as many fine fibers as possible in the nonwoven web to improve low surface tension fluid strikethrough times. By reducing the number of thicker fibers and increasing the number of fine fibers less than 1 micron in the N-layer, the embodiments of the present disclosure achieve finer pore sizes and higher low surface tension fluid strikethrough times than conventional webs. In one embodiment, the mean-flow pore diameter in the second component layer 132 may be less than 20 micron, alternatively less than 15 micron, alternatively less than 10 micron, and alternatively less than 5 micron. The mean-flow pore diameter corresponds to the pressure (called mean-flow pressure) below which half the flow happens, while the rest half of the flow happens above that pressure. Since pore diameter and pressure are inversely related, smaller mean-flow pore diameter suggests higher mean-flow pressure or flow resistance that slows down the flow, and increases the fluid strikethrough time. Because the mean-flow pore diameter is a flow attribute of a structure it is distinct from the average pore diameter that is just a statistical number average of pore diameter distribution, and the average pore diameter may not correlate to any fixed flow attribute. Alternatively, the average pore diameter may not necessarily become smaller as the mean-flow pore diameter becomes smaller, e.g, as the fiber diameter is reduced. It is believed that it is critical for an embodiment of the present disclosure to have the mean-flow pore diameter in the second component layer 132 less than 20 micron, alternatively less than 15 micron, alternatively less than 10 micron, and alternatively less than 5 micron.

The pore size distribution of the nonwoven web of the present disclosure may have one or more peaks or modes (where the mode of a pore size distribution is defined as the pore size value with highest frequency) corresponding to the multiple component layers. In one embodiment, the pore size corresponding to the lowest or the first mode of the pore size distribution corresponds to the second component layer 132 comprising N-fibers. In such embodiment, the lowest or the first mode of the pore size distribution may be less than 15

micron, alternatively less than 10 micron, and alternatively 5 micron or less. As described above, smaller pore diameter suggests higher resistance to the flow, and accordingly greater fluid strikethrough time. In some embodiments, the diameter corresponding to the lowest mode (corresponding to the smallest fibers) blocks the last 20% or more of the fluid flow (that is the pore diameters larger than the lowest mode allow the 80% or less of the fluid flow). Therefore, it is believed that the smallest pores, the higher their number the better, provide the highest resistance to the flow, and increase fluid strikethrough time.

The porosity of the second component layer 132 may be greater than 50%, alternatively greater than 70%, and alternatively greater than 80%. Since porosity corresponds to the void volume through which flow may happen, lower porosity resists the flow, and accordingly increases the liquid strikethrough time. The second component layer 132 may have at least 50% fibers with the number-average diameter less than 1 micron, alternatively at least 70% fibers with the number-average diameter less than 1 micron, alternatively at least 80% fibers with the number-average diameter less than 1 micron, and alternatively at least 90% fibers with the number-average diameter less than 1 micron. Nonwoven structures with a significant number of fibers of diameter less than 1 micron have been described by Isele et al. in U.S. Pat. Publ. Nos. 2006/0014460 published on January 1, 2006, and 2005/0070866 published March 31, 2005, both assigned to The Procter and Gamble Company, using the methods described by Torobin et al. and Reneker et al. However, having even more than 90% fibers with diameter less than 1 micron in the second nonwoven component layer 132 is not sufficient (but necessary) to have the mass-average diameter less than 1 micron, even though the number-average diameter may be less than 1 micron as described herein. In one embodiment, the second nonwoven component layer 132 may have at least 99% of fibers with the number-average diameter less than 1 micron. Therefore, in an embodiment of the present disclosure with the second nonwoven component layer 132 comprising fibers with the mass-average diameter less than 1 micron and the number-average fiber diameter less than 1 micron, almost all the fibers may have a diameter less than 1 micron, alternatively all the fibers of the second nonwoven component layer 132 in such an embodiment are submicron.

The polydispersity of fiber diameter distribution, defined as the ratio of the mass-average diameter to the number-average diameter, of the fibers comprising the second nonwoven component layer 132 may be less than 2, alternatively less than 1.8, alternatively less than 1.5, alternatively less than 1.25, alternatively less than 1.1, and alternatively 1.0. The polydispersity

of fiber diameter distribution measures the width of fiber distribution. The higher the value of the polydispersity of the distribution, the wider is the distribution. In one embodiment, as the polydispersity approaches 1, that is, the mass-average and number-average fiber diameters are the same, the second nonwoven component layer 132 may have an extremely uniform and narrow fiber distribution. The arithmetic difference between the mass-average diameter and the number-average diameter may be less than one standard deviation of the number-average diameter, alternatively, the difference may be less than three-fourths of one standard deviation of the number-average diameter, alternatively, the difference may be less than one-half of one standard deviation of the number-average diameter. Because of the above-mentioned fiber diameter averages and polydispersity of fiber diameter distribution, the N-fibers in the second nonwoven component layer 132 of the present disclosure differ from typical ultra-fine meltblown fibers that may also have the number-average diameter less than 1 micron, but typically have the mass-average diameter greater than 1 micron, and even greater than 2 microns or higher due to presence of a finite number of fibers with the diameter greater than 1 micron. As mentioned above, even with significantly large percentage of fibers, alternatively greater than 90% of fibers, having a diameter less than 1 micron, the ultra-fine meltblown fibers may not have the mass-average diameter near or less than 1 micron. The difference between the mass-average and the number-average diameters of the ultra-fine fibers may be greater than one-half of one standard deviation of the number-average diameter, more typically, the difference may be greater than one standard deviation of the number-average diameter, alternatively, the difference may be greater than two standard deviations of the number-average diameter of the ultra-fine meltblown fibers. In one embodiment, the second nonwoven component layer 132 may have a basis weight in the range of 0.1 gsm to 10 gsm, alternatively, in the range of 0.2 gsm to 5 gsm, alternatively, in the range of 0.5 to 3 gsm, and, alternatively 1 to 1.5 gsm.

In one embodiment, the nonwoven web 112 may comprise a third nonwoven component layer 136 which itself is comprised of coarse fibers, such as spunbond fibers, and may be similar to the first nonwoven component layer 125.

If the fourth nonwoven component layer 128 is used, such as a meltblown layer, these intermediate diameter fibers may comprise fibers having an average diameter, alternatively number-average diameter, in the range of 0.7 microns to 8 microns, alternatively in the range of 1 micron to 8 microns, and, alternatively, in the range of 1 micron to 5 microns, with a relative standard deviation in the range of 20% to over 100%. The mass-average diameter of the fourth

nonwoven component layer 128, such as a meltblown layer, may be in range of 0.7 microns to 8 microns, alternatively in the range of 1 micron to 8 microns, and, alternatively, in the range of 1 micron to 5 microns, and alternatively in the range of 2 to 5 micron, with a relative standard deviation in the range of 20% to over 100%. In addition, the polydispersity of the fiber diameters in the intermediate fiber layer is in the range from 1 to 10, alternatively from 2 to 8, alternatively from 2 to 6, alternatively from 1.5 to 5. Stated another way, the fourth nonwoven component layer 128 may comprise fibers having an average denier in the range of 0.003 to 0.4, alternatively, in the range of 0.006 to 0.3, with a relative standard deviation of in the range of 50% to 600%, alternatively in the range of 150% to 300%. In one embodiment, the meltblower layer may have a basis weight in the range of 0.1 gsm to 10 gsm, alternatively, in the range of 0.2 gsm to 5 gsm, and, alternatively, in the range of 0.5 gsm to 3 gsm and, alternatively, in the range of 1 to 1.5 gsm.

Also, the intermediate and fine diameter fibers may be of a bicomponent or polymer blend type, for example.

In one embodiment, referring to FIGS. 1-3, the absorbent article 10 may be configured to be worn about a lower torso of a wearer. In various embodiments, the absorbent article 10 may comprise a chassis 47 comprising a topsheet 20, a backsheet 30, and an absorbent core 40 disposed between, or at least partially between, the topsheet 20 and the backsheet 30. A pair of longitudinal barrier cuffs 51 may be attached to and/or formed with a portion of the chassis 47, such as the topsheet 20, for example. Each longitudinal barrier cuff 51 may be formed of a web of material, such as an SNS web or an SMNS web. In one embodiment, the web of material may be formed of a plurality of nonwoven component layers arranged in various combinations and permutations of a plurality of spunbond, meltblown, and N-fiber layers, including but not limited to SMN, SMNMS, SMMNMS, SSMMNS, SSNNSS, SSSNSSS, SSMMNNSS, SSMMNNMS, and the like. The webs of material disclosed herein exhibit exceptional, unexpected properties when compared to related webs of material as described in further detail below.

In one embodiment, referring to FIGS. 5 and 6, a web of material 112 may comprise a first nonwoven component layer 125 comprising fibers having an average diameter in the range of 8 microns to 30 microns, a second nonwoven component layer 132 comprising fibers having a number-average diameter of less than 1 micron, a mass-average diameter of less than 1.5 micron, and a polydispersity ratio less than 2, and a third nonwoven component layer 136

comprising fibers having an average diameter in the range of from 8 microns to 30 microns. Stated another way, the web of material 112 may comprise the first nonwoven component layer 125 comprising fibers having an average denier in the range of 0.4 to 6, the second nonwoven component layer 132 comprising fibers having an average denier in the range of 0.00006 to 0.006, and a third nonwoven component layer 136 comprising fibers having an average denier in the range of 0.4 to 6. In such an embodiment, the second nonwoven component layer 132 may be disposed intermediate the first nonwoven component layer 125 and the third nonwoven component layer 136. Also, the first nonwoven component layer 125, the second nonwoven component layer 132, and the third nonwoven component layer 136 may be intermittently bonded to each other using any suitable bonding process, such as a calendering bonding process, for example. In one embodiment, the web of material 112 does not comprise a film. In various embodiments, the web of material 112 may comprise a spunbond layer, which may correspond to the first nonwoven component layer 125, an N-fiber layer, which may correspond to the second nonwoven component layer 132, and a second spunbond layer, which may correspond to the third nonwoven component layer 136, together referred to herein as an “SNS web.”

SMS (spunbond-meltblown-spunbond) webs may have pore sizes which sometimes allow low surface tension fluids to penetrate therethrough after a particular increment of time. Some photographs of such SMS webs are illustrated in FIGS. 7 and 8. FIG. 7 is a top view of an 13 gsm SMS web 215 at 500 times magnification. FIG. 8 is a cross-sectional view of the SMS web 215 of FIG. 7 taken through a calendering bond site in the SMS web at 500 times magnification. Non-limiting example photographs, which are taken using a scanning electron microscope (SEM), of an 15 gsm SNS web 212 are illustrated in FIGS. 9 and 10. FIG. 9 is a top view of the SNS web 212 at 200 times magnification. FIG. 10 is a cross-sectional view of the SNS web 212 of FIG. 9 taken through a calendering bond site in the SNS web 212 at 500 times magnification. In one embodiment, other configurations (i.e., layering patterns) of the web of material 212 are envisioned and are within the scope of the present disclosure, such as a web of material comprising a spunbond layer, an N-fiber layer, a second spunbond layer, and a third spunbond layer of different composition or fiber cross-section, for example.

In one embodiment, a web of material, such as the SNS web 212, for example, may have a total basis weight of less than 30 gsm, alternatively, less than 15 gsm, alternatively, e.g., 13 gsm, alternatively, less than 10 gsm, and alternatively, in the range of 7 gsm to 15 gsm. In such an embodiment, the web of material may not comprise a film and has an air permeability of at

least $1 \text{ m}^3/\text{m}^2/\text{min}$, alternatively, at least $10 \text{ m}^3/\text{m}^2/\text{min}$, alternatively, at least $20 \text{ m}^3/\text{m}^2/\text{min}$, and alternatively, at least $40 \text{ m}^3/\text{m}^2/\text{min}$ but less than $100 \text{ m}^3/\text{m}^2/\text{min}$. In one embodiment, the web of material may have a local basis weight variation of less than 10%, alternatively, less than 8%, and alternatively, less than 6%, and a 32 mN/m low surface tension fluid strikethrough time of at least 19 seconds, alternatively, at least 23 seconds, alternatively, at least 30 seconds, alternatively, at least 35 seconds, alternatively, at least 40 seconds, alternatively, at least 45 seconds, and alternatively, at least 50 seconds.

In one embodiment, referring to FIGS. 11 and 12, a web of material 212' may comprise a first nonwoven component layer 225' comprising fibers having an average diameter in the range of 8 microns to 30 microns, a second nonwoven component layer 232' comprising fibers having a number average diameter of less than 1 micron, a mass-average diameter of less than 1.5 micron, and a polydispersity ratio less than 2, a third nonwoven component layer 236' comprising fibers having an average diameter in the range of 8 microns to 30 microns, and a fourth nonwoven component layer 228' comprising fibers having an average diameter in the range of 1 micron to 8 microns. Stated another way, the web of material 212' may comprise the first nonwoven component layer 225' comprising fibers having an average denier in the range of 0.4 to 6, the second nonwoven component layer 232' comprising fibers having an average denier in the range of 0.00006 to 0.006, a third nonwoven component layer 236' comprising fibers having an average denier in the range of 0.4 to 6, and a fourth nonwoven component layer 228' comprising fibers having an average denier in the range of 0.006 to 0.4. In such an embodiment, the second nonwoven component layer 232' and the fourth nonwoven component layer 228' may be disposed intermediate the first nonwoven component layer 225' and the third nonwoven component layer 236'. Also, the first nonwoven component layer 225', the second nonwoven component layer 232', the third nonwoven component layer 236', and the fourth nonwoven component layer 228' may be intermittently bonded to each other using any bonding process, such as a calendering bonding process, for example. In one embodiment, the web of material 212' does not comprise a film. In various embodiments, the web of material 212' may comprise a spunbond layer, which may correspond to the first nonwoven component layer 225', a meltblown layer, which may correspond to the fourth nonwoven component layer 228', an N-fiber layer, which may correspond to the second nonwoven component layer 232' and a second spunbond layer, which may correspond to the third nonwoven component layer 236', together referred to herein as an "SMNS web." Non-limiting example photographs, which are taken

using a scanning electron microscope, of an SMNS web 212'' are illustrated in FIGS. 13 and 14. FIG. 13 is a top view of the SMNS web 212'' at 1000 times magnification. FIG. 14 is a cross-sectional view of the SMNS web 212'' of FIG. 13 at 500 times magnification. In one embodiment, other configurations of webs of material are envisioned and are within the scope of the present disclosure, such as a web of material comprising a spunbond layer, a meltblown layer, an N-fiber layer, a second spunbond layer, and a third spunbond layer of different structure or composition, for example.

In one embodiment, referring to FIG. 1, the chassis 47 may define the two end edges 57, and the central longitudinal axis 59 may be defined in the chassis 47 and extend from one midpoint of an end edge 57 to the midpoint of the other end edge 57. In various embodiments, referring to FIGS. 1, 3A, 11 and 12, the third nonwoven component layer 236' may be positioned most proximal to the central longitudinal axis 59, the first nonwoven component layer 225' may be positioned most distal from the central longitudinal axis 59, and the second nonwoven component layer 232' may be disposed intermediate the third nonwoven component layer 236' and the fourth nonwoven component layer 228'. FIG. 3A comprises an exploded portion of the web 212' which illustrates this configuration. In certain other embodiments, the fourth nonwoven component layer 228' may be disposed intermediate the third nonwoven component layer 236' and the second nonwoven component layer 232', for example. It is possible to determine where the second nonwoven component layer 232' and/or the fourth nonwoven component layer 228' are positioned within a web using an SEM. In general, low surface tension fluid strikethrough times appear to improve by 10% to 15%, for example, when the second nonwoven component layer 232' is positioned closer to the skin of the wearer (i.e., closer to the central longitudinal axis 59 of the absorbent article 10). This is referred to as "sidedness."

In one embodiment, by positioning the second nonwoven component layer 232' closer to the central longitudinal axis 59 than the fourth nonwoven component layer 228', the second nonwoven component layer 232' is positioned closer to the skin of the wearer when the absorbent article 10 is positioned about the lower torso of the wearer. Without intending to be bound by any particular theory, applicants believe that the SMNS web exhibits more desirable properties and/or characteristics (e.g., low surface tension fluid strikethrough time) when the second nonwoven component layer 232' is positioned closer to the skin of the wearer and the source of the fluid insult into the absorbent article (and prior to use, closer to the central

longitudinal axis 59) than the fourth nonwoven component layer 228'. The arrow 213 of FIG. 3A illustrates the direction of flow of a body exudates or fluid relative to the positioning of the various nonwoven component layers.

In one embodiment, a web of material, such as the SMNS web 212', may have the same or similar properties as the properties as that described above with regard to an SNS web 212. For example, the SMNS web 212' may have a total basis weight of less than 30 gsm, alternatively, less than 15 gsm, alternatively, e.g., 13 gsm, alternatively, less than 10 gsm, and alternatively, in the range of 7 gsm to 15 gsm. In such an embodiment, the web of material may not comprise a film and may have an air permeability of at least $1 \text{ m}^3/\text{m}^2/\text{min}$, alternatively, at least $10 \text{ m}^3/\text{m}^2/\text{min}$, alternatively, at least $20 \text{ m}^3/\text{m}^2/\text{min}$, and alternatively, at least $40 \text{ m}^3/\text{m}^2/\text{min}$ but less than $100 \text{ m}^3/\text{m}^2/\text{min}$. In one embodiment, the web of material may have a local basis weight variation of less than 10%, alternatively, less than 8%, and alternatively, less than 6% and a 32 mN/m low surface tension fluid strikethrough time of at least 19 seconds, alternatively, at least 23 seconds, alternatively, at least 30 seconds, alternatively, at least 35 seconds, alternatively, at least 40 seconds, alternatively, at least 45 seconds, and alternatively, at least 50 seconds.

In one embodiment, the webs described herein, such as the SNS web and/or the SMNS web, for example, may exhibit the specified properties even without comprising a hydrophobic material, such as a hydrophobic melt additive or a hydrophobic surface coating, for example. Such features provide the webs of the present disclosure significant cost-saving advantages over related webs as adding hydrophobic materials leads to additional manufacturing cost and complexity. The inclusion of the N-fiber layer within the webs allows the webs to maintain a desirable low surface tension fluid strikethrough time and air permeability without any hydrophobic materials or films. Without intending to be bound by any particular theory, applicants believe that the N-fiber layer reduces the pore size of the webs by filling in voids within the spunbond and meltblown layers. By creating webs with smaller pore sizes when compared to the pore sizes of related webs, the webs of the present disclosure may have higher capillary drag forces to fluid penetration and, thereby, a longer low surface tension fluid strikethrough time, even without comprising a hydrophobic material or a film. Still, when looking at the structure of the SNS or the SMNS webs, the efficacy of the N-fiber layer in boosting the barrier performance of the web was not expected.

As referenced above, some absorbent articles comprise hydrophilic surfactants or materials on topsheets and/or central portions thereof, for example, and also may comprise

hydrophobic materials on barrier cuffs thereof. The hydrophilic surfactants or materials may be used to draw bodily fluids toward an absorbent core of an absorbent article, while the hydrophobic materials restrict the flow of bodily fluids through the barrier cuffs. In some instances, the hydrophilic surfactants or materials may naturally migrate toward other materials prior to use of the absorbent articles. When the hydrophilic surfactants or materials come into contact with the barrier cuffs formed of webs of materials, they reduce the web's ability to hinder low surface tension bodily fluid flow through the barrier cuffs. However, the applicants have found that the webs provided herein, such as the SNS web and/or the SMNS web, for example, may reduce the degradation of barrier properties of the web after hydrophilic surfactant's or material's migration from the topsheet or other central portion of an absorbent article to the barrier cuffs, owing perhaps to the fact that the webs of the present disclosure have higher surface areas and dilute the migrating hydrophilic surfactants when used as the barrier cuffs, or used as a portion of the barrier cuffs. In that, in one embodiment, no hydrophobic material may be present on the barrier cuffs, the hydrophilic surfactants or materials may not spread out fully on the barrier cuffs and, therefore, may not reduce the barrier cuff's ability to restrict the flow of low surface tension bodily fluids therethrough.

In other embodiments, it may be desirable for the webs to comprise a hydrophobic melt additive and/or a hydrophobic surface coating. The hydrophobic melt additive and/or the hydrophobic surface coating may increase the low surface tension fluid strikethrough time of the SNS web and/or the SMNS web, while not significantly decreasing the air permeability.

Hydrophobic additive formulations and methods for incorporating them in nonwoven webs are described by Catalan in US Publication Nos. 2006/0189956 filed on February 18, 2005 and 2005/0177123 filed on February 10, 2005, and in US Publication Nos. 2010-0222757 filed on January 22, 2010, and 2010-0221407 filed on January 22, 2010 both to JJ Tee et al. that are all assigned to The Procter and Gamble Company. Some suitable, but not limiting, hydrophobic materials used as hydrophobic surface coatings and/or hydrophobic melt additives may comprise one or more silicone polymers that are also substantially free of aminosilicones. Suitable silicone polymers are selected from the group of silicone MQ resins, polydimethylsiloxanes, crosslinked silicones, silicone liquid elastomers, and combinations thereof. Typically, the molecular weight of such silicone polymers should be at least 4000 MW. However, the molecular weight of such silicone polymers may be at least 10,000 MW, at least 15,000 MW, at least 20,000 MW, or at least 25,000 MW. Suitable

polydimethylsiloxanes are selected from the group consisting of vinyl-terminated polydimethylsiloxanes, methyl hydrogen dimethylsiloxanes, hydroxyl-terminated polydimethylsiloxanes, organo-modified polydimethylsiloxanes, and combinations thereof.

Alternatively, fluorinated polymers may also be used as the hydrophobic surface coatings and/or the hydrophobic melt additives. Suitable fluorinated polymers are selected from the group of telomers and polymers containing tetrafluoroethylene and/or perfluorinated alkyl chains. For instance, fluorinated surfactants, which are commercially available from Dupont under the tradename Zonyl® are suitable for use herein.

In one embodiment, these hydrophobic materials may be deposited onto the surface of the SNS web and/or the SMNS web in amounts of from at least 1 µg of coating per 1 g of a web. A suitable amount of silicone polymer present on the surface may be at least 100 µg/g. In certain embodiments, the amount of silicone polymer present on the surface may be at least 200 µg/g. In other embodiments, the amount of silicone polymer present on the surface may be at least 300 µg/g, alternatively, at least 400 µg/g or, alternatively, in the range of 1000 µg/g to 10,000 µg/g, for example.

The hydrophobic surface coating may be delivered to a substrate and/or fiber surface by any conventional methods. Without intending to be bound by any particular theory, it is believed that the hydrophobic surface coatings disclosed herein, when topically applied to the surface of a fibrous substrate (e.g., nonwoven surface), tend to envelop or at least partially coat one or more fibers and/or fibrous structures of the web in such a way that a cohesive, uniform film-like network is formed around the fiber and/or fibrous structures, and partially also fills the pore network of the web. In certain embodiments, hydrophobic materials may be included as an additive to a hot melt composition (e.g., blended into a thermoplastic melt), which is then formed into fibers and/or a substrate (e.g., by spunbonding, meltblowing, or extruding) (referred to herein as “hydrophobic melt additives”). Those minute additions of hydrophobic materials (chemical components) increase the contact angle of the fibers with liquid to some degree; namely for 1000 µg/g the contact angle for water increases from 100 to 110 degrees.

In one embodiment, a web of material comprising a hydrophobic surface coating and/or a hydrophobic melt additive, such as an SNS web or an SMNS web comprising these materials, for example, may have a total basis weight of less than 30 gsm, alternatively, less than 15 gsm, e.g., 13 gsm, alternatively, less than 10 gsm, and alternatively, in the range of 7 gsm to 15 gsm. In such an embodiment, the web of material may not comprise a film and may have an air

permeability of at least $1 \text{ m}^3/\text{m}^2/\text{min}$, alternatively, at least $10 \text{ m}^3/\text{m}^2/\text{min}$, alternatively, at least $20 \text{ m}^3/\text{m}^2/\text{min}$, and alternatively, at least $40 \text{ m}^3/\text{m}^2/\text{min}$ but less than $100 \text{ m}^3/\text{m}^2/\text{min}$. In one embodiment, the web of material may have a local basis weight variation of less than 10%, alternatively, less than 8%, and alternatively, less than 6% and a 32 mN/m low surface tension fluid strikethrough time of at least 30 seconds, alternatively, at least 35 seconds, alternatively, at least 40 seconds, alternatively, at least 47 seconds, alternatively, at least 50 seconds, alternatively, at least 55 seconds, alternatively, at least 60 seconds, alternatively, at least 65 seconds, and alternatively, at least 70 seconds.

In one embodiment, the webs of the present disclosure, for example, the SNS or the SMNS webs, and in the relevant comparisons, e.g., with SMS, all have a porosity (% void fraction) of over 80% (e.g., 85%). The porosity of 85% arises since the M and N fiber layers have 80% to 85% porosity and the first nonwoven component layers 132 have 85% to 92% porosity. A lower porosity may be achieved by flat calendering and reducing the breathability or by referring to a film, e.g., a microporous film, however the desired air permeabilities listed above then may become unachievable.

MECHANICAL BONDING

During construction of an absorbent article, such as absorbent article 10, for example, a web, such as an SNS web and/or an SMNS web, for example, may need to be attached to another component of the absorbent article 10. In some embodiments, as described in more detail below, a first portion of the web may be mechanically bonded to a second portion of the web, thereby creating a hem, for example. The components of the absorbent article sought to be mechanically bonded may be passed through a mechanical bonding apparatus.

FIG. 15 illustrates a simplified dynamic mechanical bonding apparatus 320 in accordance with one non-limiting embodiment of the present disclosure. The mechanically bonding apparatus 320 may comprise a patterned cylinder 322, an anvil cylinder 324, an actuating system 326 for adjustably biasing cylinders 322 and 324 towards each other with a predetermined pressure within a predetermined range of pressures, and drivers 328 and 329 for rotating the cylinders 322 and 324, respectively, at independently controlled velocities to provide an optional predetermined surface velocity differential therebetween. In one embodiment, the cylinders 322 and 324 may be biased towards each other at approximately 10,000 psi, for example.

A web 341, a web 342, and a laminate 345 are also shown in FIG. 15. In various embodiments, the web 341 may be various webs of nonwoven material, such as a 13 gsm polypropylene SNS web and/or SMNS web, for example, and the web 342 may be, for example, a 12gsm, 1.5 denier polypropylene spunbond topsheet, or other component of an absorbent article. Additionally, the apparatus 320 may comprise a frame, not shown, and drivers, not shown, for driving rolls 331 through 338 for controllably forwarding the web 341 and the web 342 through the nip 343 defined between the patterned cylinder 322 and the anvil cylinder 324, and for enabling forwarding the resulting laminate (laminate 345) to a downstream apparatus, such as a roll winder or web converting apparatus: for example, a disposable diaper converter. As used herein, “laminate” refers to at least two components of an absorbent article sharing at least one mechanical bond. Generally, the driving rolls 331 through 338, inclusive, may be provided for guiding and advancing the webs or the web 341 and the web 342, and the laminate 345 through and away from nip the 343. These rolls 331 through 338 may be driven at surface velocities which maintain predetermined levels of tension or stretch so that neither slack web conditions nor excessively tensioned/stretched webs and/or laminate precipitate undesirable deleterious consequences.

For the purposes of clarity, neither the upstream ends or sources of the web 341 and the web 342, nor the downstream destination or user of the laminate 345 are shown. In some embodiments, the mechanically bonding apparatus 320 may received more than two laminates for bonding, and the laminates to be mechanically bonded may comprise, for example, thermoplastic films, nonwoven materials, woven materials, and other webs in roll form; and to provide upstream unwinding and splicing devices to enable forwarding continuous lengths of such laminate through the mechanical bonding apparatus 320 and/or other converters to make products comprising laminated and/or other web elements at controlled velocities and under controlled tension. Furthermore, for simplicity and clarity, the mechanical bonding apparatus 320 is described herein as comprising the cylinders 322 and 324. However, the cylinders 322 and 324 are but one embodiment of nip defining members as stated. Accordingly, it is not intended to thereby limit the present disclosure to an apparatuses comprising cylinders. Similarly, the use of the term “pattern element” is not intended to limit the present disclosure to bonding patterns comprising only discrete, spaced pattern elements to the exclusion of other patterns: e.g., reticulated patterns or patterns comprising continuous or elongate lines of bonding.

In one embodiment, the actuating system 326 for biasing the patterned cylinder 322 towards the anvil cylinder 324 may comprise a pressure regulator 355, and a pneumatic actuator 356, for example. The pressure regulator 355 may be adapted to have its inlet connected to a supply source "P" of pressurized air, and to have its outlet connected to the pneumatic actuator 356 in order to adjust and control the pneumatic actuator means loading of the cylinders 322 and 324 towards each other. Although only one pneumatic actuator 356 is illustrated in FIG. 15, additional actuators may be connected to each end journal of the patterned cylinder 322; and each end journal may be supported by frame members and ancillary hardware (not shown) to be vertically moveable so that, in fact, the pressure biasing mechanism may be effective.

In one embodiment, the drivers 328 and 329, are provided to independently drive the cylinders 322 and 324, respectively. Thus, they may rotate the cylinders 322 and 324 so that there is a predetermined but adjustable relationship between the surface velocities of the cylinders 322 and 324. In various embodiments, the rotation may be synchronous, or asynchronous; equal surface velocities; or with a predetermined surface velocity differential with either of the cylinders 322 and 324 being driven faster than the other. In one embodiment that is integrated into a disposable diaper converter, the patterned cylinder 322 is driven by a converter line drive through a gear train so that its surface velocity is essentially matched to the line velocity of the converter; and, the anvil cylinder 324 is powered by an independently speed controlled DC (direct current) drive. This implementation may enable adjustment of the surface velocity of the anvil cylinder 324 to be equal to, or less than, or greater than the surface velocity of the patterned cylinder 322 by predetermined amounts or percentages.

Referring now to FIG. 16, the patterned cylinder 322 may be configured to have a cylindrical surface 352, and a plurality of pins, nubs, or other protuberances, collectively referred to as pattern of elements 351, which extend outwardly from the surface 352. As shown in FIG. 16, the patterned cylinder 322 may have a saw-tooth shape pattern of elements 351, which may extend circumferentially about each end of the patterned cylinder 322. Such a patterned cylinder 322 may be configured, for example, to laminate, lap-seam, or otherwise mechanically bond together the laminate 341 and the laminate 342. In one embodiment, the patterned cylinder 322 may be comprised of steel and may have a diameter of 11.4 inches (about 29 cm.), for example. While the illustrated embodiment shows two sets of pattern of elements 351 extending circumferentially around the patterned cylinder 322, in other embodiments, the patterned cylinder 322 may have more or less patterns of elements 351 and the overall width of

the patterned cylinder 322 may vary accordingly. The anvil cylinder 324 (FIG. 15) may be smooth surfaced, right circular cylinder of steel. In one embodiment, the anvil cylinder 324 may have a 4.5 inch (about 11.4 cm.) diameter and may be independently power rotated by a speed controlled direct current motor, for example, although the embodiments are not limited to such configurations.

FIG. 17 is a plan view of a fragmentary portion of the laminate 345 of FIG. 16 comprising overlapping edge portions of the laminate 341 and the laminate 342 which have been mechanically bonded together by a pattern of bond sites 351b: the pattern being the pattern of pattern elements which extends circumferentially about one end of patterned cylinder 322 (FIG. 16). The bond sites 351b (e.g., bond points, bond areas, dimples, nubs, land areas, cells, or elements) on the laminate 345 may have any suitable geometric shape (e.g., triangle, square, rectangle, diamond, other polygonal shapes, circle, ellipse, oval, oblong, and/or any combinations thereof). The shape and size of the bond pattern may be selected to yield bond sites 351b having predetermined strength and elasticity characteristics in the MD and CD directions, generally referred to in the art as tensile and elongation physical properties. The bond sites' 351b arrangement may be hexagonal, rectangular, square, or any other suitable polygonal shapes, for example. Generally, compressed fibers at the bond sites 351b give strength and reinforcement to the laminate 345, such a barrier cuff nonwoven web comprising an SNS web and/or SMNS web bonded to a spunbond topsheet of an absorbent article, for example. For clarity, the MD oriented edges of the laminate 341 and the laminate 342 are designated as 341e and 342e, respectively, in FIG. 17.

As is to be appreciated, the pattern of elements 351 on the patterned cylinder 322 may be configured to generate a variety of bond site patterns. FIGS. 18A-D illustrate patterns of bond sites in accordance with various non-limiting embodiments. In certain embodiments, the arrangement of the bond sites 351b may be staggered to reduce or eliminate the stress concentration of a "straight" line in the MD. The width (illustrated as "W") of the pattern may vary. For example, in certain embodiments the width may be less than 10 mm, alternatively, less than 5 mm, alternatively, less than 4 mm, and, alternatively, less than 3 mm. Some patterns, for example, may comprise bond sites 351b having different shapes and/or cross sectional areas. In one embodiment, individual bond sites 351b may be 2 mm long and 1 mm wide, and, in one embodiment, individual bond sites 351b may be 4 mm long and 1 mm wide, although other bond site sizing may be used in other embodiments. Furthermore, the area of individual bond

sites 351b may vary. In one embodiment the bond area may be 4 mm^2 , alternatively, alternatively, 2 mm^2 , and, alternatively, 1.5 mm^2 or less. The bond density per square cm may vary based on the particular application. For example, in one embodiment, there may be 15 bonds per cm^2 , alternatively, 10 bonds per cm^2 , and, alternatively, less than 10 bonds per cm^2 . Based on the bond density, the relative bond area (which is the bond density multiplied by the bond area per pin) may be 50% or less in some embodiments and, alternatively, may be 30% or less in other embodiments.

As the nonwoven web, such as the SNS web and the SMNS web, for example, is compressed during the mechanical bonding process, without intending to be bound by any particular theory, it is believed that the rapid compression of the materials beneath the protuberances 351 causes the respective materials to be rapidly deformed and at least partially expressed from beneath the pattern of elements 351. As a result, structures of entangled or otherwise combined material are formed beneath and/or around the protuberances to create mechanical bonds in the nonwoven web. In various embodiments, the mechanical bonds may be created without the use of adhesives, heat sources for a thermal welding process, or an ultrasonic wave source.

FIG. 19 is a sectional view taken along line 19--19 of FIG. 17, which illustratively shows a bond site 351b which mechanically bonds the web 341 and the web 342 together to form the laminate 345. In the illustrated embodiment, the web 341 may be an SNS web material, with an N-fiber layer 432 positioned intermediate a first nonwoven component layer 425 and a second nonwoven component layer 436. The web 342 may comprise any suitable materials, such as a topsheet of an absorbent article, a spunbond or another SNS web, or a second portion of the web 341, for example. In some embodiments, one or both of the web 341 and the web 342 may comprise an SMNS web, comprising both a meltblown layer and a N-fiber layer, in addition to two spunbond layers. In some embodiments, at least one of the webs 341, 342 may comprise a polypropylene component. In one embodiment, if an SMNS web is passing through the mechanical bonding apparatus 320 (FIG. 15), the material may be oriented such that the nubs (or pins) exert force on the N-fiber layer before exerting force on the meltblown layer. This configuration may lead to a displacement and more uniform expression of the N fibers into the underlying and surrounding fibrous structure, with a resulting higher bond strength than when the M layer (or generally coarser fiber layer) are more proximate to the nubs.

As shown in FIG. 19, the bond site 351b may have a bottom surface 351bb and a ring 376 formed substantially around the periphery of the bond site 351b, defined as the grommet ring. The grommet ring 376 may extend above the first nonwoven component layer 425 to form a ridge-like structure generally surrounding each bond site 351b. Without intending to be bound by any particular theory, it is believed that that compression forces applied to the laminate 341 and the laminate 342 during the mechanical bonding process cause material flow (e.g., fiber flow) from a bond center 378 toward the bond's periphery thereby forming the grommet ring 376. In some embodiments, the thickness of the bond site 351b at the bond center 378 may be less than 50 micrometers and, alternatively, less than 15 micrometers. Despite the formation of a robust bond using the aforementioned techniques, the bond site 351b may and should still maintain a material barrier 380 across the entire bottom surface 351bb. If the material barrier 380 across the bottom surface 351bb is breached, the laminate 345 may undesirably leak through the breach when a fluid is introduced to the bond site 351b.

Compared to the bond site 351b, in a thermal bond or a calender bond most of the adhesive force comes from fusion of materials in the bond center, and formation of a grommet ring is may not occur. In fact, the average mass of material per unit area (i.e., basis weight) inside of a thermal bond point is generally the same as in the unbonded surrounding area. In contrast, the grommet ring 376, for example, is postulated to provide most of the bond strength for the mechanical bond, and the bond center 378 has a significantly reduced basis weight compared to the surrounding area. Furthermore, the use of the N-fiber layer(s) in the nonwoven webs helps to provide a significant increase in the uniformity. In some embodiments, the local basis weight variation may be less than 15% , alternatively, less than 10%, and, alternatively, in the range of 5% to 10%.

Without intending to be bound by any particular theory, with regard to performance during the mechanical bonding process, applicants believe that the N-fibers (with diameters less than 1 micron) in the nonwoven web significantly increase the surface area of the web by 4 to 5 times (inversely proportional to the diameters of the fibers that are produced) compared to SMS or spunbond nonwoven webs of same basis weight. The increase in surface area may serve to increase the number of fibers underneath the pattern of elements during the mechanical bonding process to better distribute the energy from the pattern of elements and distribute it throughout the web. Additionally, the use of the N-fibers may allow the web to be covered more densely to create a more uniform web having a relatively low basis weight variation (e.g., less than 10%

local basis weight variation). As a result, the materials incorporating the N-fibers display less defects within the bond sites. In some embodiments, mechanically bonded webs comprising at least one N-fiber layer may have a defect occurrence rate of less than 0.9%, alternatively less than 0.54% and, alternatively, less than 0.25%. with the bonded nonwoven web having a basis weight (combined basis weight of two webs or more) of less than 25gsm. Furthermore, in accordance with the embodiments of the present disclosure, webs incorporating the N-fiber layer, such as SNS webs and SMNS webs, for example, may utilize generally small bond areas as compared to other webs, such as SMS webs. Moreover, the desired performance of the webs may be achieved with lower basis weights and/or lower stock heights when the N-fiber layer is used. In some embodiments, the bonded nonwoven material may have a low basis weight (e.g., less than 25 gsm or less than 15 gsm) and achieve mechanical bonds with suitable defect occurrence rates.

FIG. 20 is a sectional perspective view of the bond site 351b shown in FIG. 19. As illustrated, the grommet ring 376 extends generally around the periphery of the bond site 351b. Additionally, the material barrier 380, such as a membrane, extends across the bond site 351b in order to substantially “seal” the bond to maintain the bond’s fluid barrier characteristics.

Utilizing the aforementioned mechanical bonding techniques, a barrier cuff, for example, may be attached to, or otherwise integrated with, an absorbent article. Referring to FIGS. 1, 2, 3A-3B and 5, the absorbent article 10 may comprise a pair of longitudinal barrier cuffs 51, attached to a chassis 47. The chassis 47 may be any component or portion, or collection of components or portions, of the absorbent article 10, such as the topsheet 20, for example. Each longitudinal barrier cuff 51 may be comprised of a web 65, such as an SNS web or an SMNS web, having the characteristics described above. For example, the web 65 may comprise a first nonwoven component layer 125 comprising fibers having an average diameter in the range of 8 microns to 30 microns and a second nonwoven component layer 132 comprising fibers having an average diameter of less than 1 micron. The web of material 65 of the longitudinal barrier cuffs 51 may have a local basis weight variation less than 10%, alternatively, less than 8%, or alternatively, less than 6%. In fact, applicants estimated that a low defect rate of less than 10 bond defects per 5 m of a 25 gsm laminate (bond occurrence rate less than 0.35%) would require an SMS web to have an even lower local basis weight variation of 3% or less. In one embodiment, an SNS or SMNS web of 13gsm (comprising an N and M layer of 1 gsm each) or less, when combined with a spunbond layer of 12 gsm or less is sufficient to require a local basis

weight variation of 6% or less in order to achieve a defect rate of less than 10 bond defects per 5 m of laminate (bond occurrence rate less than 0.35%). The variation of 10% would suffice for an SNS or SMNS web of 13 to 15gsm with the N layer of 1.5 gsm to 3 gsm, or combining two layers of SNS or SMNS webs of 12 gsm to 13 gsm each. Each of the longitudinal barrier cuffs 51 may comprise a longitudinal zone of attachment 49 where the longitudinal barrier cuff 51 attaches to the chassis 47. In some embodiments, the longitudinal zone of attachment 49 may extend generally parallel to the central longitudinal axis 59 (FIG. 1). In some embodiments, the zone of attachment 49 may be generally linear or may be curved, or a combination.

Furthermore, the zone of attachment 49 may be substantially continuous along the absorbent article or, alternatively, discontinuous. Furthermore, each longitudinal barrier cuffs 51 may have a longitudinal free edge 64 and a plurality of mechanical bonds 68 disposed between the zone of attachment 49 and the free edge 64. In one embodiment, the plurality of mechanical bonds 68 forms a hem proximate to the longitudinal free edge 64. For example, the plurality of mechanical bonds 68 may attach, for example, a first portion of the web material 59 to a second portion 61 of the web 65, which may be referred to as a hem fold bond. In some embodiments, the mechanical bonds 68 may attach the web 65 to a portion of the absorbent article 10. The mechanical bonds 68 may be similar to the bond site 351b illustrated in FIGS. 19-20, for example. The mechanical bonds 68 may, for example, bond the topsheet 20 to the longitudinal barrier cuffs 51. Furthermore, the mechanical bonds 68 may be disposed in any suitable pattern or configuration, such as the patterns illustrated in FIGS. 18A-18D, for example.

In another embodiment, referring to FIG. 3B, longitudinal barrier cuffs 51 of the absorbent article 10 may each comprises a first layer of the web of material 65a and a second layer of the web of material 65b. The first and second layers of web material 65a and 65b, may each comprise an SNS web or an SMNS web, for example. Furthermore, as illustrated, the longitudinal barrier cuff 51 may be folded in order to form to two layers of web material 65a and 65b. In other embodiments, two separate webs of material 65a and 65b may be joined, bonded, or otherwise attached to form the longitudinal barrier cuff 51. The longitudinal barrier cuffs 51 may comprise a longitudinal zone of attachment 49 where the longitudinal barrier cuff attaches to the chassis 47 and a longitudinal free edge 64. A plurality of mechanical bonds 68 may attach the first and second layers of the web of material 65a and 65b. In some embodiments, the plurality of mechanical bonds 68 attach at least on of the first and second layers of the web of material 65a and 65b to the chassis 47. In one embodiment, the plurality of mechanical bonds

68 have a defect occurrence rate of less than 0.9%, alternatively, less than 0.5% and, alternatively, less than 0.25%. In some embodiments, the plurality of mechanical bonds 68 may be disposed along, or generally proximate to, the longitudinal zone of attachment 49.

In one embodiment, the SNS web and/or the SMNS web may comprise, or may comprise a portion of, a component of an absorbent article other than a longitudinal barrier cuff, such as a backsheet of a diaper, for example, owing to the webs' superior properties of air permeability, low surface tension fluid strikethrough time, basis weight, and local basis weight variation. Likewise, the SNS web and/or the SMNS web may also be used to comprise any other suitable portions of various consumer absorbent articles or other suitable non-absorbent articles or portions thereof. Some non-limiting examples of non-absorbent articles that may be formed of, or formed partially of, the SNS web and/or the SMNS web are consumer disposable water filtration components, air freshener components using perfume release for odor elimination, and surfactant release components in detergents and detergent capsules.

In other embodiments, the SNS web and/or the SMNS web may be formed with, attached to, and/or used with a film, such as microporous or micro-apertured films (or films with risk of pin holes), for example, to increase the low surface tension fluid strikethrough times of the webs for desired applications, such as when used as a backsheet of a diaper, for example. In still other embodiments, the SNS web and/or the SMNS web may comprise or be coated with a hydrophobic melt additive and/or a hydrophobic surface coating to again increase the low surface tension fluid strikethrough times of the webs for desired applications. In one embodiment, the SNS web and/or the SMNS web may comprise both a film and a hydrophobic melt additive and/or a hydrophobic surface coating, for example. Such web embodiments with the film, the hydrophobic melt additive, and/or the hydrophobic surface coating may comprise or may be used as components of any suitable absorbent or non-absorbent articles, such as diaper backsheets, catamenial pad topsheets or backsheets, for example.

TESTS

Air Permeability Test

The air permeability is determined by measuring the flow rate of standard conditioned air through a test specimen driven by a specified pressure drop. This test is particularly suited to materials having relatively high permeability to gases, such as nonwovens, apertured films and the like.

A TexTest FX3300 instrument or equivalent is used. (Available by Textest AG in Switzerland (www.textest.ch), or from Advanced Testing Instruments ATI in Spartanburg SC, USA.) The Test Method conforms to ASTM D737. The test is operated in a laboratory environment at $23 \pm 2^\circ\text{C}$ and $50 \pm 5\%$ relative humidity. In this test, the instrument creates a constant differential pressure across the specimen which forces air through the specimen. The rate of air flow through the specimen is measured in $\text{m}^3/\text{m}^2/\text{min}$, which is actually a velocity in m/min , and recorded to three significant digits. The test pressure drop is set to 125 Pascal and the 5 cm^2 area test head is used. After getting the system operational, the 1 cm^2 insert is installed (also available from Textest or from ATI). The sample of interest is prepared and specimens cut out to fit into the 1 cm^2 head insert. After making the measurement of a specimen according to operating procedure, the result is recorded to three significant digits accounting for the area difference between the 1 cm^2 test area insert and the 5 cm^2 head. If the FX3300 instrument is not accounting for this automatically, then each specimen's result is manually recalculated to reflect the actual air permeability by accounting for the area difference between the 1 cm^2 test area insert and the 5 cm^2 head. The average of 10 specimens' air permeability data of this sample is calculated and reported.

Surface Tension of a Liquid

The surface tension of a liquid is determined by measuring the force exerted on a platinum Wilhelmy plate at the air-liquid interface. A Kruss tensionmeter K11 or equivalent is used. (Available by Kruss USA (www.kruss.de)). The test is operated in a laboratory environment at $23 \pm 2^\circ\text{C}$ and $50 \pm 5\%$ relative humidity. The test liquid is placed into the container given by the manufacturer and the surface tension is recorded by the instrument and its software.

Surface tension of a Fiber

Basis Weight Test

A 9.00 cm^2 large piece of web, i.e. 1.0 cm wide by 9.0 cm long, is cut out of the product, and it needs to be dry and free from other materials like glue or dust. Samples are conditioned at 23°C ($\pm 2^\circ\text{C}$) and at a relative humidity of about 50% ($\pm 5\%$) for 2 hours to reach equilibrium. The weight of the cut web pieces is measured on a scale with accuracy to 0.0001 g . The resulting mass is divided by the specimen area to give a result in g/m^2 (gsm). Repeat for at least 20 specimens for a particular sample from 20 identical products, If the product and component is

large enough, more than one specimen can be obtained from each product. An example of a sample is the left diaper cuff in a bag of diapers, and 10 identical diapers are used to cut out two 9.00 cm^2 large specimens of cuff web from the left side of each diaper for a total of 20 specimens of “left-side cuff nonwoven.” If the local basis weight variation test is done, those same samples and data are used for calculating and reporting the average basis weight.

Mechanical Bond Defect Occurrence Rate Test

The defect occurrence rate of a mechanical bonding pattern is determined by determining the percentage of defective bonds in 5.0 meters of bonded material. Defects are holes or skips or tears. Holes are defined as an area of at least 0.39 mm^2 that is apertured or missing from the film-like membrane formed at the bond site material. Skips are defined as an area of at least 1.00 mm^2 where the intended mechanical bond site does not visually show a film-like membrane. The third type of defect, a tear, is the result of a broken perimeter of the membrane where at least 1.0 mm of the membrane’s perimeter is torn or broken. See FIG. 20 for illustration of an example material barrier 380 (or “membrane”) within a mechanical bond grommet. FIG. 21 illustrates what constitutes a good mechanical bond, a bad, but not defective mechanical bond, and a defective mechanical bond during a Mechanical Bond

Defect Occurrence Rate Test.

A visual procedure is used to measure the defect occurrence rate from a produced web of two or more webs, or from a web that is cut out of a product or product feature. First, take 5 m of the nonwoven web or equivalent number of products (e.g., 10 consecutive diapers of 0.5 m pad cuff length) and inspect one side (e.g., the left longitudinal side or the right longitudinal side of the diaper of the bond sites on the nonwoven webs for defects. Care is taken not to disrupt and damage the bonds and to select the section where the mechanical bonds have not been overbonded with a mechanical bond a second time or more.

If the component with the bonds of interest cannot be removed by simply cutting without disrupting and damaging the bonds, another method for disintegration may be used, such as use of a THF bath to dissolve the adhesives. After carefully cutting out the component with the bonds of interest, label the specimens for tracking and later analysis.

Each mechanical bond pattern has a certain repeat length. The total target number of bonds in the 5m laminate web is obtained by multiplying the 5m length (5000mm) with the

number of bonds per repeat length (#bonds/mm). If the mechanical bonds of the bond pattern of interest are so large as to extend the whole diaper length, the diaper length is defined as repeat length. Cut out an extra (per example 18th) section according to above from the sample of interest, tape its ends to a flat surface so the section is fully extended (manually extended to full length with reasonable force without damaging the web and to remove wrinkles and extend any elastomeric contraction) then slide a thin black piece of cardboard under the taped sample. Find a repeat length of the bond pattern over at least a 100mm section, which means for repeat lengths less than 100mm long, that multiple individual repeat lengths are selected. For example the bond pattern of Figure 18A, when measuring the length from the top to the bottom of the shown pattern and it gives 200mm, then the repeat length of the pattern in Figure 18A is from the top edge of the C-shaped bond on the very top, to the top edge of the third C-shaped bond from the top, and in this example would give 142mm. All the bonds, even if of multiple shapes, are counted and added up in this overall repeat length. In the example of 18A, the overall repeat length is 142 mm, from top of first C-shaped bond to top of third. The number of bonds in this 142 mm repeat length is 16 bonds., The total number of bonds within the 5000mm length is thus 5000 mm multiplied by 16 bonds divided by 142mm, which is 563 bonds.

Each bond site is examined under a microscope at 25x magnification. The lens is used in conjunction with a the respective defect determination templates; i.e. for holes template with a 0.39 mm^2 large circle ($0.705 \pm 0.005 \text{ mm}$ diameter), for skips the template with a 1.00 mm^2 large circle (1.0 mm diameter), and for tears the template with a 1.0 mm diameter circle, which can be seen on the specimen when viewed through the eyepiece. See illustration in Figure 21B, and outlined here further for a hole defect. If the circle can fit within the hole, then the hole is counted as a hole defect. (see Figure 21B) After one bond site is inspected, the next consecutive bond to be inspected is in the lengthwise direction of the diaper.

Holes are classified as H1, H2,...orH5, with the number reflecting the number of consecutive mechanical bonds with a hole. Consecutive defects in the same row in the diaper length direction are counted as a single defect, i.e., five consecutive holes are counted as one H5 defect. Record the results of the analysis in a data table like below, where for each specimen and each image the number of holes and skips is recorded.

Image	H1	H2	H3	H4	H5	S1	S2	S3	S4	S5	T1	T2	T3	T4	T5	Defects per specimen
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Spec 1 – image 1																
S1-i2																
Etc.																

If there are more bond shapes not yet analyzed for holes, repeat this step for those and determine the number of its defects like above using this bond shape's hole defect limit.

Skip failures are classified with the respective template and recorded.. as S1, S2,..., or S5, with the number reflecting the number of consecutive missing mechanical bonds. Consecutive defects in the same row in the diaper length direction are counted as a single defect, i.e., 5 consecutive skips is counted as one S5 defect. Tear failures are classified with the respective template and recorded. as T1, T2 . . or T5 with the number reflecting the number of consecutive missing mechanical bonds. Consecutive defects in the same row in the diaper length direction are counted as a single defect i.e. five consecutive tears are counted as one T5 defect. The total number of defects of all holes, skips and tears are added up to obtain the number of defects per 5.0 m of web. Dividing this by the theoretical number of mechanical bonds (mechanical bond density in number of mechanical bonds/cm times the length of the laminate (500cm)) and multiplied by 100% yields the defect occurrence rate in %. The theoretical number includes all mechanical bonds that would be on the 5 m laminate regardless of whether material is properly bonded or not.

See FIGURES 21A, 21B, and 33A to 33G for illustration of identifying the defects with this test.

Fiber Diameter and Denier Test

The diameter of fibers in a sample of a web is determined by using a Scanning Electron Microscope (SEM) and image analysis software. A magnification of 500 to 10,000 times is chosen such that the fibers are suitably enlarged for measurement. The samples are sputtered with gold or a palladium compound to avoid electric charging and vibrations of the fibers in the electron beam. A manual procedure for determining the fiber diameters is used. Using a mouse and a cursor tool, the edge of a randomly selected fiber is sought and then measured across its width (i.e., perpendicular to fiber direction at that point) to the other edge of the fiber. A scaled and calibrated image analysis tool provides the scaling to get actual reading in micrometers (μm).

Several fibers are thus randomly selected across the sample of the web using the SEM. At least two specimens from the web (or web inside a product) are cut and tested in this manner.

Altogether at least 100 such measurements are made and then all data are recorded for statistic analysis. The recorded data are used to calculate average (mean) of the fiber diameters, standard deviation of the fiber diameters, and median of the fiber diameters. Another useful statistic is the calculation of the amount of the population of fibers that is below a certain upper limit. To determine this statistic, the software is programmed to count how many results of the fiber diameters are below an upper limit and that count (divided by total number of data and multiplied by 100%) is reported in percent as percent below the upper limit, such as percent below 1 micrometer diameter or %-submicron, for example.

If the results are to be reported in denier, then the following calculations are made.

Fiber Diameter in denier = Cross-sectional area (in m²) * density (in kg/m³) * 9000 m * 1000 g/kg.

The cross-sectional area is $\pi \cdot \text{diameter}^2 / 4$. The density for polypropylene, for example, may be taken as 910 kg/m³.

Given the fiber diameter in denier, the physical circular fiber diameter in meters (or micrometers) is calculated from these relationships and vice versa. We denote the measured diameter (in microns) of an individual circular fiber as d_i .

In case the fibers have non-circular cross-sections, the measurement of the fiber diameter is determined as and set equal to the hydraulic diameter which is four times the cross-sectional area of the fiber divided by the perimeter of the cross of the fiber (outer perimeter in case of hollow fibers).

Fiber Diameter Calculations

The number-average diameter, alternatively average diameter, $d_{num} = \frac{\sum_{i=1}^n d_i}{n}$

The mass-average diameter is calculated as follows:

$$\text{mass average diameter, } d_{mass} = \frac{\sum_{i=1}^n (m_i \cdot d_i)}{\sum_{i=1}^n m_i} = \frac{\sum_{i=1}^n (\rho \cdot V_i \cdot d_i)}{\sum_{i=1}^n (\rho \cdot V_i)} = \frac{\sum_{i=1}^n \left(\rho \cdot \frac{\pi d_i^2 \cdot \Delta x}{4} \cdot d_i \right)}{\sum_{i=1}^n \left(\rho \cdot \frac{\pi d_i^2 \cdot \Delta x}{4} \right)} = \frac{\sum_{i=1}^n d_i^3}{\sum_{i=1}^n d_i^2}$$

where

fibers in the sample are assumed to be circular/cylindrical,

d_i = measured diameter of the i^{th} fiber in the sample,

∂x = infinitesimal longitudinal section of fiber where its diameter is measured, same for all the fibers in the sample,

m_i = mass of the i^{th} fiber in the sample,

n = number of fibers whose diameter is measured in the sample

ρ = density of fibers in the sample, same for all the fibers in the sample

V_i = volume of the i^{th} fiber in the sample.

The polydispersity of fiber diameter distribution = $\frac{(\text{mass average fiber diameter})}{(\text{number average fiber diameter})}$

Low Surface Tension Fluid Strikethrough Time Test

The low surface tension fluid strikethrough time test is used to determine the amount of time it takes a specified quantity of a low surface tension fluid, discharged at a prescribed rate, to fully penetrate a sample of a web (and other comparable barrier materials) which is placed on a reference absorbent pad. As a default, this is also called the 32 mN/m Low Surface Tension Fluid Strikethrough Test because of the surface tension of the test fluid and each test is done on two layers of the nonwoven sample simply laid on top of each other.

For this test, the reference absorbent pad is 5 plies of Ahlstrom grade 989 filter paper (10cm x 10cm) and the test fluid is a 32 mN/m low surface tension fluid.

Scope

This test is designed to characterize the low surface tension fluid strikethrough performance (in seconds) of webs intended to provide a barrier to low surface tension fluids, such as runny BM, for example.

Equipment

Lister Strikethrough Tester: The instrumentation is like described in EDANA ERT 153.0-02 section 6 with the following exception: the strike-through plate has a star-shaped orifice of 3 slots angled at 60 degrees with the narrow slots having a 10.0 mm length and a 1.2mm slot width. This equipment is available from Lenzing Instruments (Austria) and from W.

Fritz Metzger Corp (USA). The unit needs to be set up such that it does not time out after 100 seconds.

Reference Absorbent Pad: Ahlstrom Grade 989 filter paper, in 10 cm x 10 cm areas, is used. The average strikethrough time is 3.3 ± 0.5 seconds for 5 plies of filter paper using the 32 mN/m test fluid and without the web sample. The filter paper may be purchased from Empirical Manufacturing Company, Inc. (EMC) 7616 Reinhold Drive Cincinnati, OH 45237.

Test Fluid: The 32 mN/m surface tension fluid is prepared with distilled water and 0.42 ± 0.001 g/liter Triton-X 100. All fluids are kept at ambient conditions.

Electrode-Rinsing Liquid: 0.9% sodium chloride (CAS 7647-14-5) aqueous solution (9g NaCl per 1L of distilled water) is used.

Test Procedure

- Ensure that the surface tension is 32 mN/m $\pm$ 1 mN/m. Otherwise remake the test fluid.
- Prepare the 0.9% NaCl aqueous electrode rinsing liquid.
- Ensure that the strikethrough target (3.3 ± 0.5 seconds) for the Reference Absorbent Pad is met by testing 5 plies with the 32 mN/m test fluid as follows:
 - Neatly stack 5 plies of the Reference Absorbent Pad onto the base plate of the strikethrough tester.
 - Place the strikethrough plate over the 5 plies and ensure that the center of the plate is over the center of the paper. Center this assembly under the dispensing funnel.
 - Ensure that the upper assembly of the strikethrough tester is lowered to the pre-set stop point.
 - Ensure that the electrodes are connected to the timer.
 - Turn the strikethrough tester “on” and zero the timer.
 - Using the 5 mL fixed volume pipette and tip, dispense 5 mL of the 32 mN/m test fluid into the funnel.
 - Open the magnetic valve of the funnel (by depressing a button on the unit, for example) to discharge the 5 mL of test fluid. The initial flow of the fluid will complete the electrical circuit and start the timer. The timer will stop when the fluid has penetrated into the Reference Absorbent Pad and fallen below the level of the electrodes in the strikethrough plate.
- Record the time indicated on the electronic timer.

- Remove the test assembly and discard the used Reference Absorbent Pad. Rinse the electrodes with the 0.9% NaCl aqueous solution to “prime” them for the next test. Dry the depression above the electrodes and the back of the strikethrough plate, as well as wipe off the dispenser exit orifice and the bottom plate or table surface upon which the filter paper is laid.
- Repeat this test procedure for a minimum of 3 replicates to ensure the strikethrough target of the Reference Absorbent Pad is met. If the target is not met, the Reference Absorbent Pad may be out of spec and should not be used.
- After the Reference Absorbent Pad performance has been verified, nonwoven web samples may be tested.
- Cut the required number of nonwoven web specimens. For web sampled off a roll, cut the samples into 10 cm by 10 cm sized square specimens. For web sampled off of a product, cut the samples into 15 by 15 mm square specimens. The fluid flows onto the nonwoven web specimen from the strike through plate. Touch the nonwoven web specimen only at the edge.
- Neatly stack 5 plies of the Reference Absorbent Pad onto the base plate of the strikethrough tester.
- Place the nonwoven web specimen on top of the 5 plies of filter paper. Two plies of the nonwoven web specimen are used in this test method. If the nonwoven web sample is sided (i.e., has a different layer configuration based on which side is facing in a particular direction), the side facing the wearer (for an absorbent product) faces upwards in the test.
- Place the strikethrough plate over the nonwoven web specimen and ensure that the center of the strikethrough plate is over the center of the nonwoven web specimen. Center this assembly under the dispensing funnel.
- Ensure that the upper assembly of the strikethrough tester is lowered to the pre-set stop point.
- Ensure that the electrodes are connected to the timer. Turn the strikethrough tester “on” and zero the timer.
- Run as described above.
- Repeat this procedure for the required number of nonwoven web specimens. A minimum of 5 specimens of each different nonwoven web sample is required. The average value is the 32 mN/m low surface tension strikethrough time in seconds.

35 mN/m Low Surface Tension Fluid Strikethrough Test

This test is done as described above with two exceptions. First, the testing is done with one layer of the nonwoven web sample. Second, the test fluid has a surface tension of 35 mN/m. The test fluid is created by mixing 2 parts of the 32 mN/m fluid and 5 parts of deionized water. Before testing, the actual surface tension of the fluid needs to be checked to ensure that it is 35 $\pm$ 1 mN/m. If this fluid is not 35 $\pm$ 1 mN/m, it should be discarded and another fluid should be prepared.

*Local Basis Weight Variation Test**Purpose*

The local basis weight variation test is intended to measure variability of mass distribution of 9 cm² areas throughout a lot of a nonwoven web. The local basis weight variation parameter describes a lack of desirable uniformity through a nonwoven web. Lower local basis weight variation is desirable since it helps in consistency of most other qualities, such as barrier properties, strength, and bonding, for example.

Principle

The mass of 1 cm by 9 cm area nonwoven web samples are measured and analyzed to determine the local basis weight variation (i.e., mass distribution) throughout a lot of a web production. All individual data of the lot, or of a portion of the lot, of interest is analyzed as standard deviation and average and then the quotient is taken to provide the local basis weight variation. Stated another way, this gives a relative standard deviation (RSD) or coefficient of variation (COV) of the small area basis weight distribution.

- The size of 1 cm by 9 cm for each replicate was selected such that the mass of each replicate may be measured with sufficient digits and accuracy on the specified scale.
- Mass is measured in grams.
- Grammage and basis weight are synonymous and are measured in g/m² (also written gsm) units.
- Samples of the nonwoven web are taken in the machine direction (the web needs to be at least 1 cm wide such that it may be cut into specimens).

Equipment

- Scale with a 0.0001g sensitivity (alternatively, a scale with 0.00001g sensitivity or with accuracy to within 0.1% of a target basis weight) (e.g., 13 gsm in 1 cm by 9 cm area weighs 0.0117g; 0.1% of this mass is 0.00001g)
- Die with 1.0 cm by 9.0 cm or 9 cm² area rectangular cut area optionally with soft foam for easier sample removal. The die areas need to be within about 0.05 mm side length.
- Hydraulic press: The hydraulic press is used to stamp out the nonwoven web samples with the die.

Test Procedure

Sampling:

- At least 40 data points are needed to assess the local basis weight variation of a defined nonwoven web sample. These data points are to be sampled evenly throughout the nonwoven web sample.
- Test specimens should be free of wrinkles and free from contaminants such as dust or glue.

Conditioning:

-Use only clean and dry nonwoven web samples, at normal lab conditions (50+/-5% relative humidity and 23+/-2 degrees C).

Procedure:

- Cut out the replicate with the prepared die 9 cm² and the hydraulic press. One layer is cut out. Paper may be put between the cutting board and the sample for easier removal after cutting.
- Be sure that the scale reads exactly zero (0.0000g), or tare the scale to 0.0000g.
- Measure the cut out replicate on the scale to the nearest 0.0001g (alternatively, to the nearest 0.00001g).
- Record the lot, nonwoven web sample, replicate and result.
- Continue the above steps for all selected replicates.

When the analysis is done for absorbent articles (e.g., diapers) then identical products are used, preferably consecutive diapers are tested within one bag, package, or case. Either the right

of the left leg barrier cuff may be selected for the samples. For purposes of this description, we assume that the right leg barrier cuff has been selected.

- Carefully cut the leg barrier cuffs out of the absorbent articles and number the cuffs sequentially (e.g., right leg barrier cuff of absorbent article 1). Proceed with doing the same for the remaining absorbent articles in the bag, package or case.
- Beginning with the cut out leg barrier cuff from absorbent article 1, fasten (e.g., tape) the leg barrier cuff to a piece of cardboard or plastic sheeting and put the die with the cut area (1 cm by 9 cm) onto the barrier cuff and cut out a specimen. If there is still enough sample length left, repeat this procedure one or two more times for two or three more specimens out of the barrier cuff.
- Weigh the cut out parts to the nearest 0.0001g and record the result.
- Proceed with the other cut out right side leg barrier cuffs from the other absorbent articles and measure the mass of the die cut 1cm by 9cm sized pieces and record the data.
- Repeat this procedure for as many absorbent articles as needed and, if necessary, several bags of absorbent articles, until the right side barrier cuff of the absorbent articles is characterized with 40 data points. Since a package of absorbent articles typically holds over twenty absorbent articles, it is possible to cut out and measure 40 or more replicates per side (in this case the right side) for each sample package of absorbent articles.

Repeat the whole procedure for the other side of the product (in this case the left side). The local basis weight variation should be calculated for each side.

Calculations

- Calculate the average weight of the nonwoven web sample (40 individual replicates)
- Calculate the standard deviation of the nonwoven web sample
- Calculate the Local Basis Weight Variation (standard deviation/average weight).

Reporting

Report the local basis weight variability to the nearest first decimal point 0.1%, e.g., 7.329% becomes 7.3%.

Surface Tension Measurement of Fluid

The measurement is done with a video-based optical contact angle measuring device, OCA 20, by DataPhysics Instrument GmbH, or equivalent. Choose a clean glass syringe and dosing needle (with 1.65~3.05 mm size) before filling the syringe with liquid to test; and then remove the bubble from the syringe/needle; adjust the position of the syringe, dosing needle and stage; a drop of the test liquid with known volume will be formed at the lower end of the dosing needle. The detection of the drop shape is done by the software SCA20 and the surface tension is calculated according to the Young-Laplace equation. The measurement is carried out on an anti-vibration table in a closed hood.

The surface energy of fibers is also determined with this instrument following the Sessile Drop Technique.

Thickness or Caliper Test

The thickness test is done according to EDANA 30.5-99 normal procedure with a foot of 15 mm diameter pushing down at 500 Pascal (i.e., a force of 0.0884N). Start the test, wait for 5 seconds so the result stabilizes, and record the result in millimeters to the nearest 0.01mm. The sample analysis should include at least 20 measurements from different locations spread throughout the available sample.

Pore Size Distribution Test

The pore size distribution of nonwoven web samples is measured with the Capillary Flow Porometer, the APP 1500 AEXi from Porous Materials, Inc. or equivalent. The available pressure of the clean and dry air supply should be at least 100 psi so that pores down to 0.08 microns may be detected. A nonwoven web sample is first cut and fully soaked in a low surface tension fluid, namely Galwick with a surface tension of 15.9 mN/m. The nonwoven web sample size is 7 mm diameter. The soaked nonwoven web sample is placed into the sample chamber of the instrument and the chamber is then sealed. Upon starting the automatic measurement cycle, gas flows into the sample chamber behind the nonwoven web sample and then the gas pressure is slowly increased via the computer to a value sufficient to overcome the capillary action of the fluid in the pore having the largest diameter in the nonwoven web sample. This is the bubble point. The pressure inside the chamber is further increased in small increments resulting in a flow of gas that is measured until all of the pores in the nonwoven web sample are empty of the

low surface tension fluid. The gas flow versus pressure data represents the “wet curve.” When the curve continues to rise linearly, the sample is considered to be dry (i.e., the pores are emptied of the low surface tension fluid). The pressure is then decreased in steps producing the “dry curve.” From the relationships of the “wet” and “dry” curves, the computer calculates the pore parameters including the mean-flow pore diameter and a histogram of pore diameters across the tested range (e.g., from the bubble point down to about 0.08 microns or even less with higher gas pressure) as is known to those of skill in the porous media field.

Some key parameters for the test procedure with the capillary flow porometer are the following: the test fluid is Galwick with 15.9 mN/m surface tension; the test area opening size is 7 mm; and the tortuosity parameter is set to 1. Other parameters of the instrument are set to max flow: 100,000 cc/min, bubble flow 3 cc/min, F/PT parameter 1000, zero time 2s, v2incr 25 cts*3, preginc 25 cts*50, pulse delay 0s, maxpres 1 bar, pulsewidth 0.2s, mineqtime 10s, presslew 10 cts*3, flowslew 30 cts*3, equiter 10*0.1s, aveiter 10*0.1s, max press diff 0.01 bar, max flow diff 40 cc/min, starting press 0.1 bar, and starting flow 500 cc/min.

Nonwoven Tensile Strength (in CD)

The nonwoven tensile strength (in CD) is measured using an Instron MTS 3300 tensile tester, or equivalent according to WSP 110.4(05)B. A nonwoven web sample of 15 mm x 50 mm, where the 50 mm length is along the length of the diaper product. The sample width is 50 mm, The gauge length is 5 mm, allowing for 5 mm to be placed in each sample clamp. The test speed is 100 mm/min. A stress-strain curve is measured until the sample breaks. The nonwoven tensile strength is defined as the maximum stress value observed of the curve.

Bond Peel Strength

The bond peel strength is defined as the force required to separate the two bonded layers of barrier leg cuff and the topsheet in the longitudinal direction. The test is measured using an MTS 3300 tensile tester or equivalent. A nonwoven laminate specimen of 15 mm x 170 mm is removed from the product. A free end is created in the last 20mm by manually peeling apart the topsheet from the barrier leg cuff layer, thus obtaining a free end with a cuff face and a topsheet face. The test speed is 305 mm/min. The specimens are obtained from the product as described in the Mechanical Bond defect occurrence rate test.

Procedure

- Insert the free end of the barrier leg cuff layer of the specimen into the lower jaw with the length axis of the sample perpendicular to the upper edge of the jaw, and close the jaw. Align the specimen between the lower and upper jaws. Insert the free end of the topsheet layer of the specimen into the upper jaw with the length axis of the sample perpendicular to the lower edge of the jaw and close the jaw with enough tension to eliminate any slack, but less than 5 grams of force on the load cell. Do NOT zero the instrument after the specimen has been loaded.
- Start the tensile tester and data collection device simultaneously as described by the manufacturer's instructions.
- Remove the specimen from the clamps and return the crosshead to the starting position in preparation for the next specimen.

If tearing has occurred during testing, cut another specimen from the same general area of the sample. If tearing occurs during testing of this second specimen also, record the bond strength for the specimen as "total bond".

- Disregard results for the first 2.5 cm of peel. If the tensile tester is computer interfaced, set the program to calculate the average peel force in grams for the specimen.

EXAMPLES

EXAMPLE 1

In this example, the second nonwoven component layer 132 comprises N-fibers having fiber diameters (measured per the Fiber Diameter and Denier Test set forth herein), polydispersity, fiber diameter ranges (minimum – maximum measured), and amounts of submicron diameter fibers (less than 1 micron) illustrated in Table 1A below:

Table 1A

Sample No.	Number Average Diameter (microns)	Mass Average Diameter (microns)	Polydispersity Ratio	Standard Deviation (microns)	Fiber Diameter Range (microns)	Amount of Submicron Fibers (%)
N1	0.34	0.39	1.14	0.09	0.15 – 0.55	>99%
N2	0.33	0.45	1.36	0.09	0.08 – 0.78	>99%
N3	0.38	0.48	1.27	0.14	0.17 – 0.77	>99%
N4	0.68	0.73	1.08	0.14	0.40 – 0.98	>99%
N5	0.57	0.95	1.66	0.31	0.11 – 2.23	92%
N6	0.84	0.96	1.13	0.22	0.25 – 1.55	74%
N7	0.85	1.02	1.19	0.27	0.26 – 1.60	79%

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N8	0.69	1.12	1.63	0.37	0.23 – 1.84	85%
N9	1.03	1.21	1.18	0.33	0.28 – 1.98	43%
N10	0.78	1.23	1.59	0.39	0.29 – 2.31	80%

Comparative Example 1: A nonwoven component layer comprises meltblown fibers having fiber diameters (measured per the Fiber Diameter and Denier Test set forth herein), polydispersity, fiber diameter ranges (minimum – maximum measured), and amounts of submicron diameter fibers (less than 1 micron) illustrated in Table 1B below.

Table 1B

Sample No.	Number Average Diameter (microns)	Mass Average Diameter (microns)	Polydispersity Ratio	Standard Deviation (microns)	Fiber Diameter Range (microns)	Amount of Submicron Fibers (%)
M1	0.69	1.64	2.39	0.58	0.15 – 2.68	80%
M2	0.45	1.97	4.35	0.44	0.10 – 5.55	93%
M3	0.61	2.99	4.91	0.65	0.07 – 8.44	86%
M4	1.36	1.86	1.37	0.56	0.41 – 3.32	21%
M5	1.78	2.15	1.21	0.55	0.84 – 3.99	4%
M6	1.44	2.25	1.56	0.71	0.46 – 4.40	26%
M7	1.71	2.62	1.54	0.82	0.70 – 4.76	10%
M8	3.16	4.16	1.32	1.23	1.80 – 6.80	0%
M9	1.85	4.10	2.22	1.39	0.67 – 6.44	23%
M10	1.54	4.60	2.99	1.47	0.20 – 8.18	38%
M11	2.27	6.17	2.72	1.85	0.55 – 12.17	10%

In Table 1B, the samples identified by the numbers M1 through M3 represent ultra-fine meltblown fibers, the samples identified by the numbers M4 through M7 represent fine meltblown fibers, and the samples identified by the numbers M8 through M11 represent intermediate meltblown fibers.

The data set forth in Table 1A and Table 1B is illustrated in FIGS. 22 through 25. The number average diameter and the mass average diameter values, shown in the Tables 1A and 1B, are depicted on the statistically fitted curves to the fiber diameter distributions in FIGS. 22 through 25. FIG. 22 compares the fiber diameter distribution of the N-Fibers sample N1 with the fiber diameter distribution of the ultra-fine meltblown fibers sample M1. Similarly, FIG. 23 compares the fiber diameter distribution of the N-Fibers samples N1 through N4 with the fiber diameter distribution of the ultra-fine meltblown fibers samples M1 through M3. The comparison of N-Fibers and ultra-fine meltblown fibers shows that even though ultra-fine

meltblown fibers samples comprise significant number of fibers (at least 80%) with diameters less than 1 micron, they also comprise finite number of fibers (about 6% to 20%) with diameters greater than 1 micron (to up to 8.4 microns), making the fiber distributions with long tails on the large diameter end. The long large diameter end tails of fiber distributions are well-described by the mass average diameters, which range between 1.64 and 2.99 along with a polydispersity ratio ranging between 2.39 and 4.91. FIGS. 24 and 25 compare the fiber diameter distributions of N-Fibers samples N1 through N4 with the fine and intermediate size meltblown fiber samples, respectively. The meltblown fiber samples are labeled in FIGS. 24 and 25. The fiber diameter distributions of the meltblown samples in FIGS. 24 and 25, and Table 1B depict that fiber diameters range from submicron (< 1 micron) to as large as 12 microns, making the fiber distribution significantly wide with long tails on the large fiber diameter end. Owing to the presence of large diameter fibers (illustrated by the long tails of the fiber distributions on the large fiber diameter end) in the measured samples, listed in Table 1B, both the mass average and the number average diameters for all the measured meltblown samples lie on the distribution tails, and the mass average diameters are more than about 1 standard deviation greater than the number average diameters. In comparison, the N-fibers have a very small number of large diameter fibers in the measured samples. Therefore, the fiber diameter distributions of N-Fibers have short tails, and both the number average and the mass average diameters are tended towards the center of the fiber distributions, and are within about 1 standard deviation of the number average diameters.

EXAMPLE 2A

In this example, various samples of nonwoven web materials A-i are tested. Their various properties are displayed in Table 2A. Samples G-i are embodiments of nonwoven web materials of the present disclosure, while SMS samples A-F are provided merely for comparison purposes. The low surface tension fluid strikethrough times of the various samples are illustrated graphically in FIG. 26 (with the exception of sample J to provide a graph with a better scale). As can be seen from FIG. 26, the low surface tension fluid strikethrough times of samples G-I of the present disclosure are significantly higher than SMS samples A-F, even when the SMS webs are coated with a hydrophobic coating (see SMS samples D-F). The low surface tension fluid strikethrough values are determined using two plies of each sample and a 32 mN/m low surface tension fluid.

Table 2A

Sample No.	Material Type	Total Basis Weight (g/m ²)	Fine Fiber Basis Weight (g/m ²)	Low Surface Tension Fluid Strikethrough (s)	Air Permeability (m/min)
A	SMS	15.7	1	13	91
B	SMS	16.9	3	16	72
C	SMS	13.3	1.5	13	80
D	SMS + Hydrophobic Coating 1	15.2	1	19	96
E	SMS + Hydrophobic Coating 1	17.1	3	20	84
F	SMS + Hydrophobic Coating 2	15.1	1	23	70
G	SNS	15.5	1.5	32	52
H	SNS + Hydrophobic Coating 1	15.6	1.5	47	50
I	SMNS	13.3	1 (M) + 1 (N)	33	59

EXAMPLE 2B

In this example, various samples of nonwoven web materials A-I (same as Example 2A) are tested. Their various properties are displayed in Table 2B. Samples G-I are embodiments of nonwoven web materials of the present disclosure, while SMS samples A-F are provided merely for comparison purposes. The low surface tension fluid strikethrough times of the various samples are plotted against their number average diameter (microns) in FIG. 27. As is illustrated in FIG. 27, the low surface tension fluid strikethrough time increases based on the smaller number average diameter of the fibers in the sample. The low surface tension fluid strikethrough values are determined using two plies of each sample and a 32 mN/m low surface tension fluid.

Table 2B

Sample ID Material Type	A SMS	B SMS	D SMS	F SMS	G SNS	H SNS	I SMNS
Hydrophobic Coating	-	-	Type 1	Type 2	-	Type 1	-
Total Basis Weight (g/m ²)	15.7	16.9	15.2	15.1	15.5	15.6	13.3
MeltBlown Fiber Basis Weight (g/m ²)	1	3	1	1	-	-	1
N-Fiber Basis Weight (g/m ²)	-	-	-	-	1.5	1.5	1
Spunbond Number Average Diameter (micron)	14.85	15.57	15.95	18.40	16.98	16.98	15.61
Spunbond Mass Average Diameter (micron)	15.03	15.71	16.10	18.47	16.99	16.99	15.67
MeltBlown Number Average Diameter (micron)	1.96	1.85	2.20	2.69	-	-	2.04
MeltBlown Mass Average Diameter (micron)	2.46	4.10	2.93	3.10	-	-	3.72
Submicron M Fibers (%)	8%	23%	2%	0%	-	-	11%
N-Fibers Number Average Diameter (micron)	-	-	-	-	0.49	0.49	0.35
N-Fibers Mass Average Diameter (micron)	-	-	-	-	0.54	0.54	0.43
Submicron N Fibers (%)	-	-	-	-	>99%	>99%	>99%
Low Surface Tension Fluid Strikethrough (s)	13	16	19	23	32	47	33
Air Permeability (m/min)	91	72	96	70	52	50	59

EXAMPLE 2C

In this example, the sidedness (i.e., which layer, the meltblown layer or the N-fiber layer, is positioned more proximal to the source of the low surface tension fluid) of the SMNS nonwoven webs of the present disclosure is illustrated against the low surface tension fluid strikethrough times of the SMNS webs. Referring to FIG. 28, in the data set on the left, the meltblown layer (i.e., the fourth nonwoven component layer) was positioned more proximal to the low surface tension fluid than the N-fiber layer in an SMNS web sample. In the data set on the right, the N-fiber layer (i.e., the second nonwoven component layer) was positioned more proximal to the low surface tension fluid than the meltblown layer of the SMNS sample. As illustrated in FIG. 28, when the N-fiber layer is positioned closer to the source of the fluid, the SMNS web provides a higher low surface tension fluid strikethrough time.

Turning to Table 2C below, a single layer of the SMNS web is tested using the 35 mN/m Low Surface Tension Fluid Strikethrough Test.

Table 2C

	M facing liquid	N facing liquid
LSTST at 35mN/m, 1 layer	Liquid → SMNS	Liquid → SNMS
Average	202	230
StDev	69.1	76.8

The single layer SMNS web has a basis weight of 13 gsm (for more specifics, see sample I in Example 2A and 2B). The variation in this Example 2C is which side of the SMNS material is facing the source of the fluid (i.e., is the material positioned fluid-SMNS or fluid-SNMS). In the set of data on the left side of FIG. 28, the sample is positioned fluid-SMNS and in the data set on the right side of Fig. 28 is positioned fluid-SNMS.

Statistical analysis shows that when the N-layer is positioned most proximal to the low surface tension fluid source, a statistically significant benefit of greater low surface tension fluid strikethrough times (with 89% certainty) is provided. Therefore, in one embodiment, an absorbent article of the present disclosure, using the SMNS web as a barrier to fluid penetration, may have the N-layer of the SMNS web facing inwards, towards the wearer of the absorbent article (i.e., wearer-SNMS). This concept is illustrated in FIG. 3A, where the N-layer of the longitudinal barrier cuff 51 is positioned more proximal to the central longitudinal axis 59 than the M-layer.

EXAMPLE 2D

In this example, a single layer of a nonwoven web is tested using the 35 mN/m Low Surface Tension Fluid Strikethrough Test. Table 2D shows the results of some comparative samples (SMS) and a sample of an SMNS web of the present disclosure.

Table 2D

Material	LSTST 1 layer at 35mN/m (Average, s)
15gsm SMS (1gsm M, 7+1+7gsm total) (= sample A)	81.0
15gsm SMS (3gsm M, 6+3+6gsm total) (= sample B)	133.9
Hydrophobic surface coated 15gsm SMS (1gsm M; additive is PDMS with surface energy 20mN/m) (= sample D)	311.7
SMNS* 13gsm (5.5+1+1+5.5gsm total) (= sample I)	229.6

The first sample in this table is equal to sample A of Example 2A and 2B. The second sample is similar to sample B of Example 2A and 2B, but has a lower overall basis weight (i.e., less spunbond basis weight) the fiber diameters of sample B's meltblown layer have a number

average diameter between 2 and 3 micrometers and a mass-average diameter of about 4 micrometers. The third sample in Table 2D is sample D from Example 2A and 2B and is coated with a hydrophobic surface additive according to Catalan in U.S. Pat. Publ. No. 2006/0189956 A1 in the following manner: a 3% solution of a vinyl terminated PDMS (commercially available from Momentive as SM3200) and a methyl hydrogen PDMS (commercially available from Momentive as SM3010) is prepared and mixed for 30 minutes. The SMS web is dipped into the solution and the excess liquid is squeezed out such that at least about 400 μ g/g of the aqueous silicone mixture is deposited on the SMS web. The SMS web is then dried in a convective oven at 120°C for 1 minute and then cooled and stored in a dry and clean location until the SMS web is ready for testing. The weight gain of the SMS web (i.e., the dry coating amount per square meter) needs to be less than 1%. The fourth sample in Table 2D is sample I from Example 2A and 2B.

Referring to Figs. 29 and 30, sample I shows a surprisingly large advantage in low surface tension fluid strikethrough times compared to the SMS samples (the first three samples of Table 2D) and is more than halfway to the performance of a hydrophobic-coated SMS in this single layer 35 mN/m Low Surface Tension Fluid Strikethrough Test. The SMNS sample (sample I) has a lower total basis weight than any of the other SMS samples (the first three samples of Table 2D), and does not have the advantage of the PDMS coating which has a low surface energy of 20mN/m to provide a higher contact angle. Sample I, even with having such a low basis weight and such a low fine fiber basis weight, and without hydrophobic chemical modification, still is capable of producing very high low surface tension fluid strikethrough times (e.g., above 150 seconds or even above 200 seconds).

EXAMPLE 3

In this example, pore size distribution of the SMS samples A and B from Example 2A are compared with the SNS sample G and the SMNS sample I from Example 2A. The pore size distribution of the embodiment of samples G and I comprising N-fibers as the finest fiber layer is significantly different and much narrower than the SMS samples A and B comprising meltblown fibers as the finest fiber layer, as illustrated in FIG. 31. The pore size distributions for all the samples have been statistically fitted with a mixture of constituent distributions (shown as dotted lines in the FIG. 31) corresponding to fine fiber and spunbond layers, with the largest pores corresponding to the spunbond layer because of larger fiber diameters than the fine fibers. While the largest mode corresponds to the largest frequency of the thick spunbond fibers,

the lowest mode corresponds to the largest frequency of the fine fibers, and the intermediate mode (for the samples A, B, and I) corresponds to the largest frequency of intermediate size fibers. The lowest mode value, mean flow, and bubble point pore diameters describing the pore size distribution are listed in Table 3 below for the samples A, B, G, and I along with their respective basis weights, fiber size distributions, low surface tension fluid strikethrough times, and air permeability values. The percent flow blocked by the lowest mode diameter is calculated from intersection of the “wet flow” and “dry flow” curves (set forth in the Pore Size Distribution Test) at the pressure corresponding to the lowest mode diameter. Table 3 also shows that the mean flow pore diameter correlates with the mass-average diameter. Additionally, the low surface tension fluid strikethrough time and air permeability correlate with the mean flow and the lowest mode pore diameters. Clearly, samples G and I of the present disclosure have significantly smaller pores and significantly longer low surface tension fluid strikethrough times when compared to SMS samples A and B.

Table 3

Sample ID	A	B	G	I
Material Type	SMS	SMS	SNS	SMNS
Total Basis Weight (g/n ²)	15.7	16.9	15.5	13.3
MeltBlown Fiber Basis Weight (g/n ²)	1	3	-	1
N-Fiber Basis Weight (g/n ²)	-	-	1.5	1
MeltBlown Number-Average Diameter (microns)	1.96	1.85	-	2.04
MeltBlown Mass-Average Diameter (microns)	2.46	4.10	-	3.72
Submicron M Fibers (%)	8%	23%	-	11%
N-Fibers Number-Average Diameter (microns)	-	-	0.49	0.35
N-Fibers Mass-Average Diameter (microns)	-	-	0.54	0.43
Submicron N Fibers (%)	-	-	>99%	>99%
Lowest Mode Pore Diameter (microns)	13.5	11.1	7.8	5.2
Flow Blocked by the Lowest Mode Pore Diameter (microns)	7%	1%	19%	9%
Mean Flow Pore Diameter (microns)	21.4	29.5	10.1	15.1
Bubble Point Pore Diameter (microns)	67.2	79	69.1	110.1
Low Surface Tension Fluid Strikethrough Time (secs)	13	16	32	33
Air Permeability (m/min)	91	72	52	59

Surprisingly, the mean flow pore diameter appears to be more important than the bubble point in order to obtain low surface tension fluid strikethrough times above 12 seconds with untreated (no hydrophobic additive) nonwoven webs having a basis weight of 15 gsm or less with 3 gsm or less fine fibers (i.e., less than 1 micron). Thus, in one embodiment, a mean flow pore diameter of 15 microns or less, alternatively of 12 microns or less, alternatively of 10

microns or less is provided. A mean flow pore diameter greater than 1 micron, alternatively greater than 3 microns, and alternatively greater than 5 microns, is provided for breathability.

EXAMPLE 4

In this example, the mechanical bonds of various nonwoven webs are evaluated using the basis weight coefficient of variation (COV) of 900 mm² samples. 5 m samples of the same materials are bonded to a 12 gsm topsheet in a docking station using a hem bond pattern at 3.5 bar and a linear speed of ~300 m/min. Various samples of web materials BLC1- BLC6 are tested. Their various properties are displayed in Table 4.

Table 4			
The bond defects characterized following	Sample No.	Material Type	mechanical are using the criteria:
	BLC1	13gsm SSMMMS (with 4gsm M)	
	BLC2	13gsm SMMMS (with 1gsm M)	
	BLC3	13gsm SMMMS #2 (with 1gsm M)	
	BLC4	13gsm SSMS (with 1gsm M)	
	BLC5	15gsm SNS (1.5gsm N, sample G)	
	BLC6	15gsm SMS (1gsm M)	

“Hole”: an aperture with a size of at least 0.39 mm² in the bond area (hole defect limit). Hole failures are classified as H1, H2,..., or H5, with the number reflecting the number of consecutive mechanical bonds with a hole. Consecutive defects are counted as a single defect, i.e., 5 holes are counted as one H5 defect.

“Skip”: a mechanical bond is missing at least an area of 1.00 mm² (skip defect limit). Skip failures are classified as S1, S2,..., or S5, with the number reflecting the number of consecutive missing mechanical bonds. Consecutive defects are counted as a single defect, i.e., 5 skips are counted as one S5 defect.

“Tear”: a tearing of the perimeter such that 1.0 mm or greater of the perimeter of the grommet ring has been torn (tear defect limit). Tear failures are classified as T1, T2,..., or T5, with the number reflecting the number of consecutive missing mechanical bonds. Consecutive defects are counted as a single defect, i.e., 5 tears are counted as one T5 defect.

The total number of defects was added up of each kind of defect.

It should be noted that a SSMMMS 13 gsm (sample BLC1) barrier leg cuff shows a significant increase in the number of mechanical bond defects. Extrapolation of a linear fit of BLC1, BLC2, BLC3, and BLC4 leads to an intersection with the horizontal line of BLC6 at a basis weight COV of 0.03 (3%). Therefore, a basis weight COV (local basis weight variation)

of 0.03 would be needed in order to attain the current levels of defects found for the 15 gsm barrier leg cuff when using a 13 gsm barrier leg cuff.

FIG. 32 is graphical illustration of the bond defects of samples BLC1- BLC6 of Table 32 as a function of basis weight COV. The line BLC6 represents the average number of defects observed over the range of basis weight COV values observed in current 15 gsm barrier leg cuffs. Previous manufacturer trials have shown that the basis weight uniformity may be increased through increasing the amount of the meltblown basis weight. The results suggests that if 13 gsm barrier leg cuff could achieve a basis weight COV value of 0.03, it would be theoretically possible to attain the current levels of bond defects and bond strength observed in the 15 gsm barrier leg cuff.

The dimensions and values disclosed herein are not to be understood as being strictly limited to the exact numerical values recited. Instead, unless otherwise specified, each such dimension is intended to mean both the recited value and a functionally equivalent range surrounding that value. For example, a dimension disclosed as "40 mm" is intended to mean "about 40 mm".

The citation of any document, including any cross referenced or related patent or application, is not an admission that it is prior art with respect to any invention disclosed or claimed herein or that it alone, or in any combination with any other reference or references, teaches, suggests, or discloses any such invention. Further, to the extent that any meaning or definition of a term in this document conflicts with any meaning or definition of the same term in a document cited herein, the meaning or definition assigned to that term in this document shall govern.

While particular embodiments of the present invention have been illustrated and described, it would be obvious to those skilled in the art that various other changes and modifications can be made without departing from the invention described herein.

Claims:

1. An absorbent article to be worn about the lower torso, the absorbent article comprising:

a chassis comprising a topsheet, a backsheet, and an absorbent core disposed between the topsheet and the backsheet; and

a pair of longitudinal barrier cuffs attached to the chassis, each longitudinal barrier cuff formed of a web of material, the web of material comprising:

a first nonwoven component layer comprising fibers having an average diameter in the range of about 8 microns to about 30 microns;

a second nonwoven component layer comprising fibers having a number-average diameter of less than about 1 micron, a mass-average diameter of less than about 1.5 microns, and a ratio of the mass-average diameter to the number-average diameter less than about 2; and

a third nonwoven component layer comprising fibers having an average diameter in the range of about 8 microns to about 30 microns;

wherein the second nonwoven component layer is disposed intermediate the first nonwoven component layer and the third nonwoven component layer.

2. The absorbent article of claim 1, wherein the first nonwoven component layer, the second nonwoven component layer, and the third nonwoven component layer are intermittently bonded.

3. The absorbent article of claim 1, comprising a fourth nonwoven component layer comprising fibers having an average diameter in the range of about 1 micron to about 8 microns.

4. The absorbent article of claim 3, wherein the chassis defines a first end and a second end, wherein a central longitudinal axis is defined in the chassis and extends from the first end to the second end, wherein the third nonwoven component layer is positioned most proximal to the central longitudinal axis, wherein the first nonwoven

component layer is positioned most distal from the central longitudinal axis, and wherein the second nonwoven component layer is disposed intermediate the third nonwoven component layer and the fourth nonwoven component layer.

5. The absorbent article of claim 3, wherein the chassis defines a first end and a second end, wherein a central longitudinal axis is defined in the chassis and extends from the first end to the second end, wherein the third nonwoven component layer is positioned most proximal to the central longitudinal axis, wherein the first nonwoven component layer is positioned most distal from the central longitudinal axis, and wherein the second nonwoven component layer is disposed intermediate the fourth nonwoven component layer and the first nonwoven component layer.

6. The absorbent article of claim 1, wherein the web of material comprises one of a hydrophobic melt additive and a hydrophobic surface coating.

7. The absorbent article of claim 1, wherein the first, second, and third nonwoven component layers together have a total basis weight less than about 15 gsm.

8. The absorbent article of claim 1, wherein the first, second, and third nonwoven component layers together have a total basis weight in the range of about 7 gsm to about 15 gsm.

9. The absorbent article of claim 1, wherein the web of material has a low surface tension fluid strikethrough time of at least about 19 seconds.

10. The absorbent article of claim 1, wherein the web of material has a low surface tension fluid strikethrough time in the range of about 19 seconds to about 40 seconds.

11. The absorbent article of claim 1, wherein the web of material has an air permeability of at least about 20 m.³/m²/min.

12. The absorbent article of claim 1, wherein the web of material has an air permeability of at least about $40 \text{ m}^3/\text{m}^2/\text{min}$.

13. An absorbent article to be worn about the lower torso, the absorbent article comprising:

a chassis comprising a topsheet, a backsheet, and an absorbent core disposed between the topsheet and the backsheet; and

a pair of longitudinal barrier cuffs attached to the chassis, each longitudinal barrier cuff formed of a web of material, the web of material comprising:

a first nonwoven component layer comprising fibers having an average denier in the range of about 0.4 to about 6;

a second nonwoven component layer comprising fibers having an average denier in the range of about 0.00006 to about 0.006 and fibers having a number-average diameter of less than about 1 micron, a mass-average diameter of less than about 1.5 microns, and a ratio of the mass-average diameter to the number-average diameter less than about 2; and

a third nonwoven component layer comprising fibers having an average denier in the range of about 0.4 to about 6;

wherein the second nonwoven component layer is disposed intermediate the first nonwoven component layer and the third nonwoven component layer.

14. The absorbent article of claim 13, comprising a fourth nonwoven component layer comprising fibers having an average denier in the range of about 0.003 to about 0.4.

15. The absorbent article of claim 13, wherein the web of material has a basis weight of less than about 15 gsm.

16. The absorbent article of claim 13, wherein the web of material has an air permeability of at least about $40 \text{ m}^3/\text{m}^2/\text{min}$.

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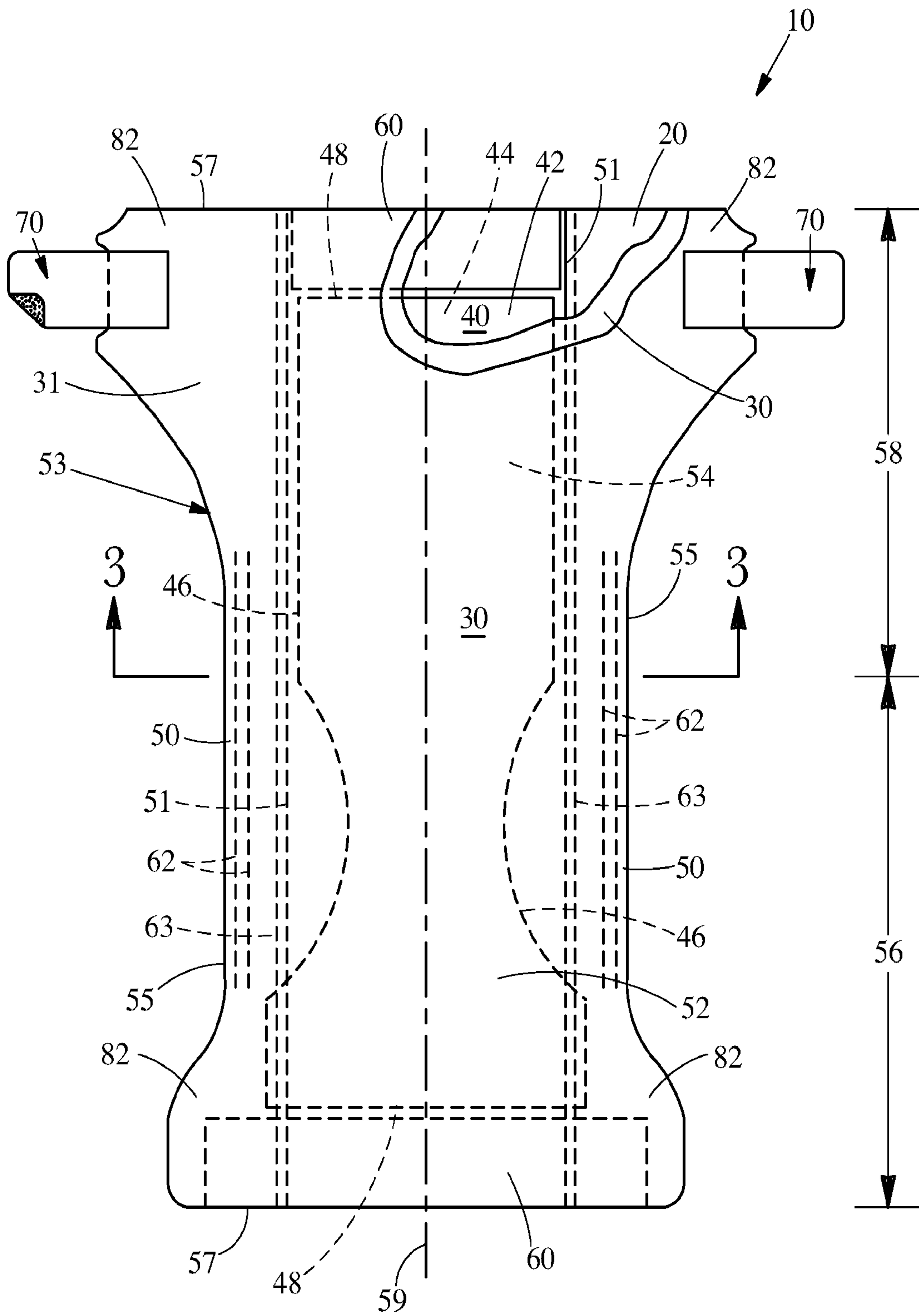


Fig. 1

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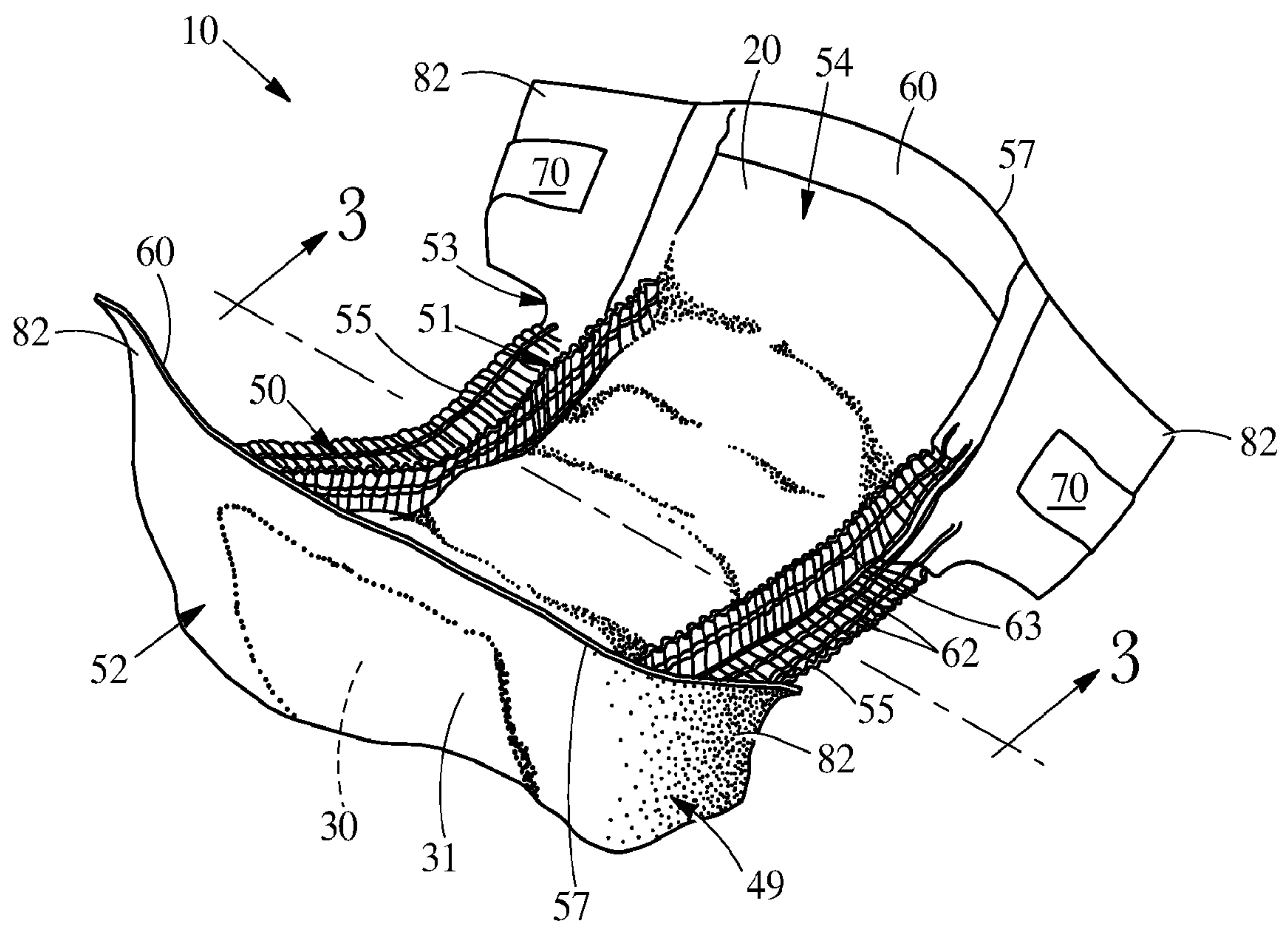


Fig. 2

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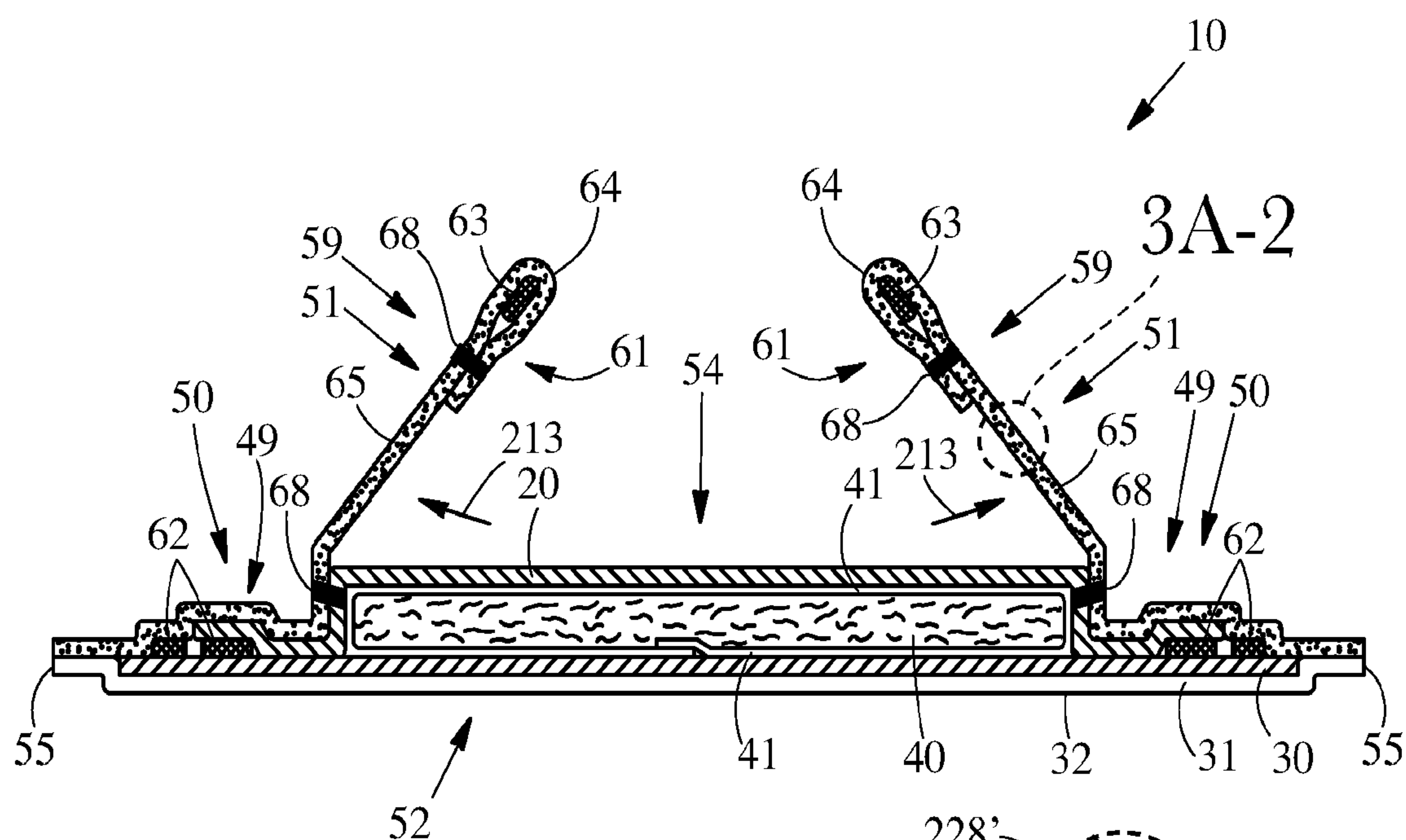


Fig. 3A-1

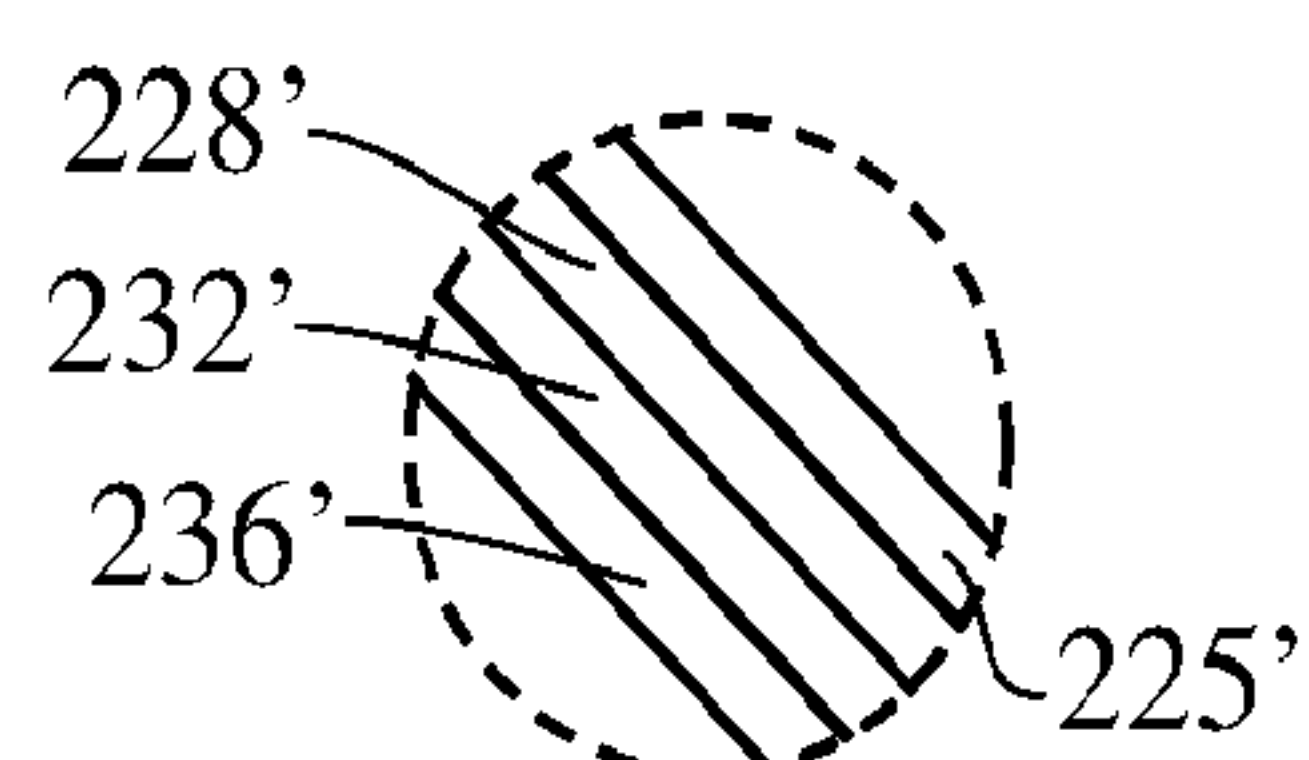


Fig. 3A-2

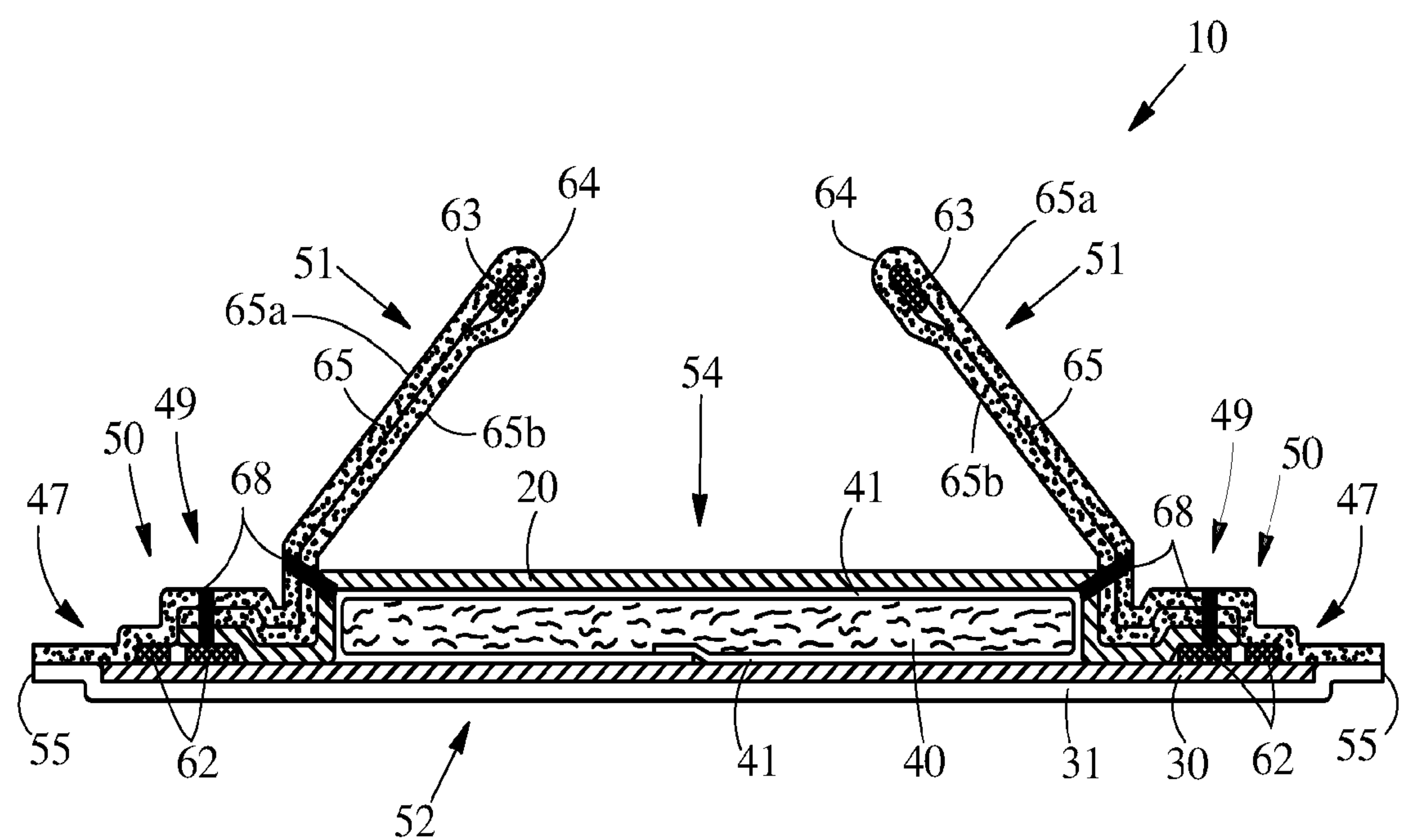


Fig. 3B

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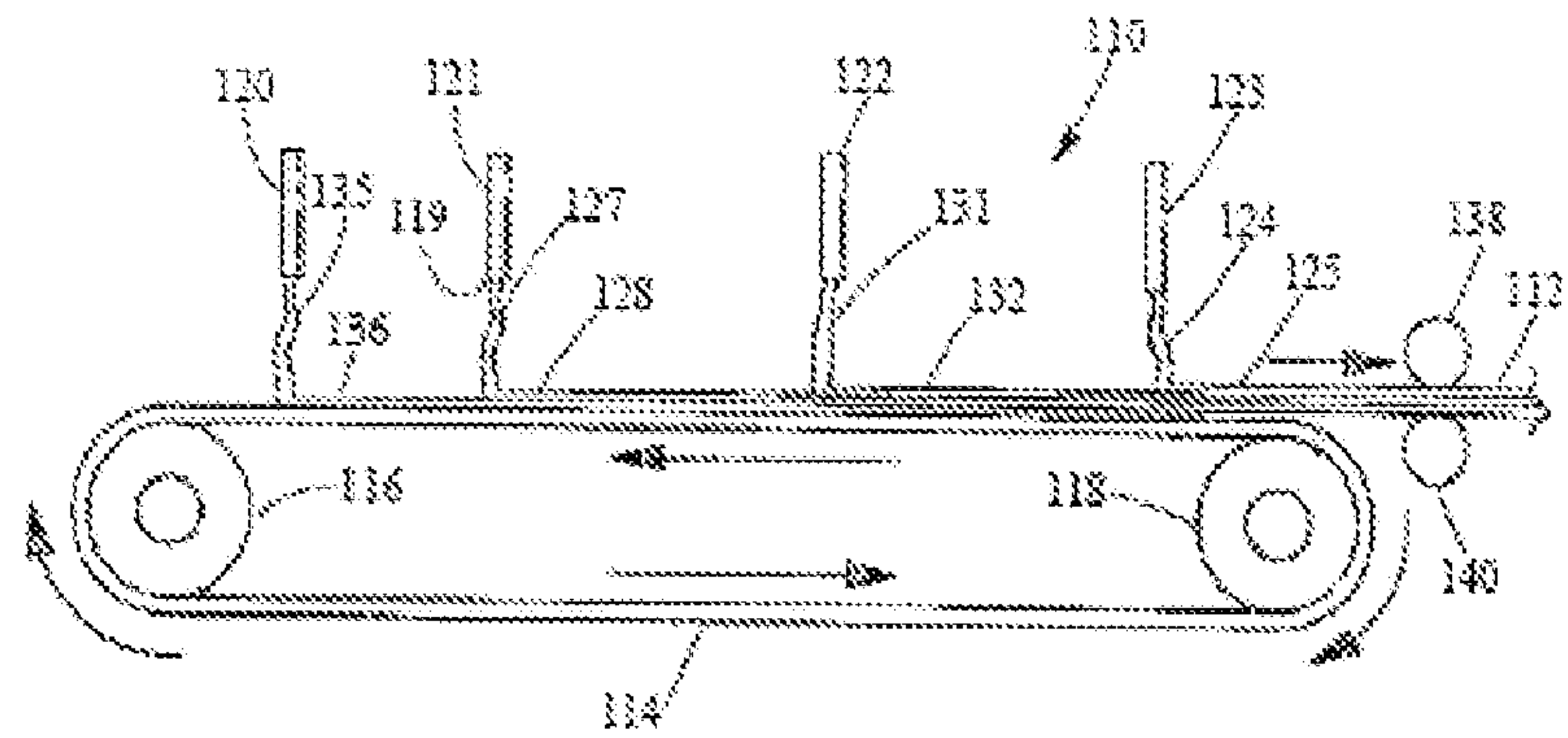


Fig. 4

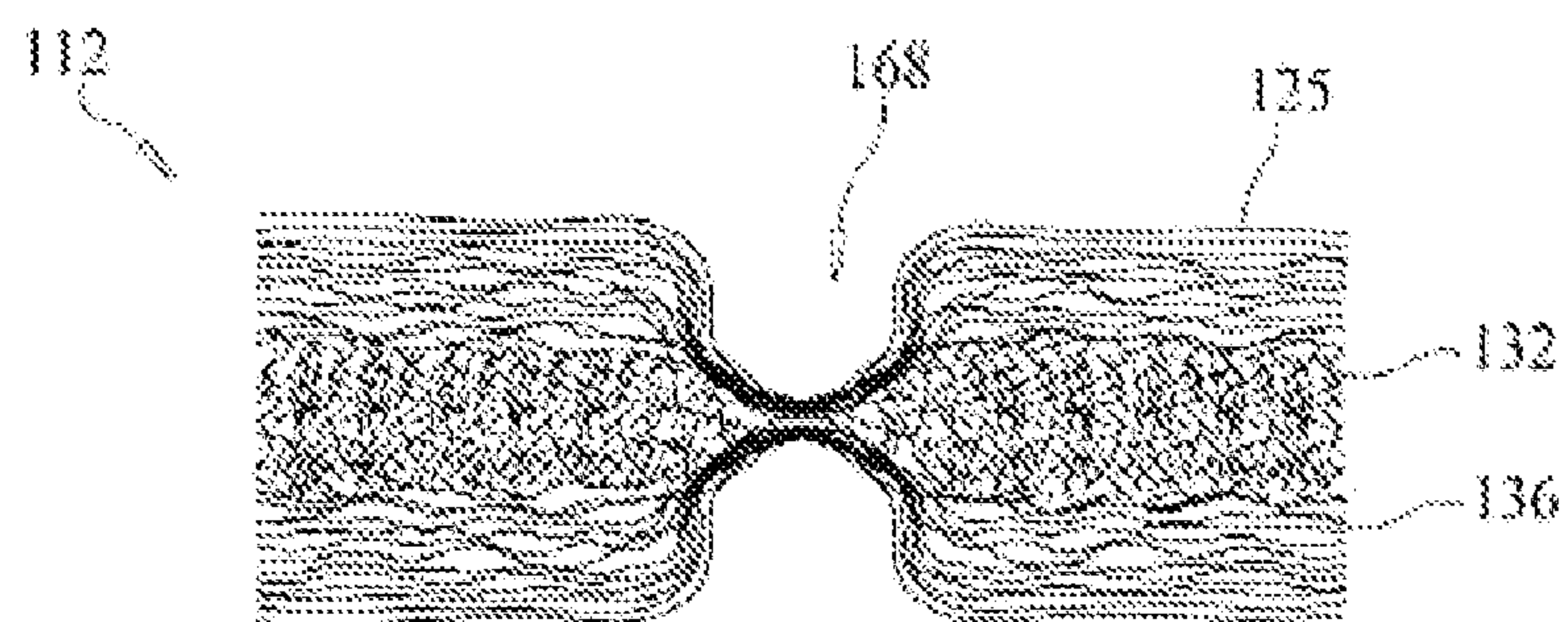


Fig. 5

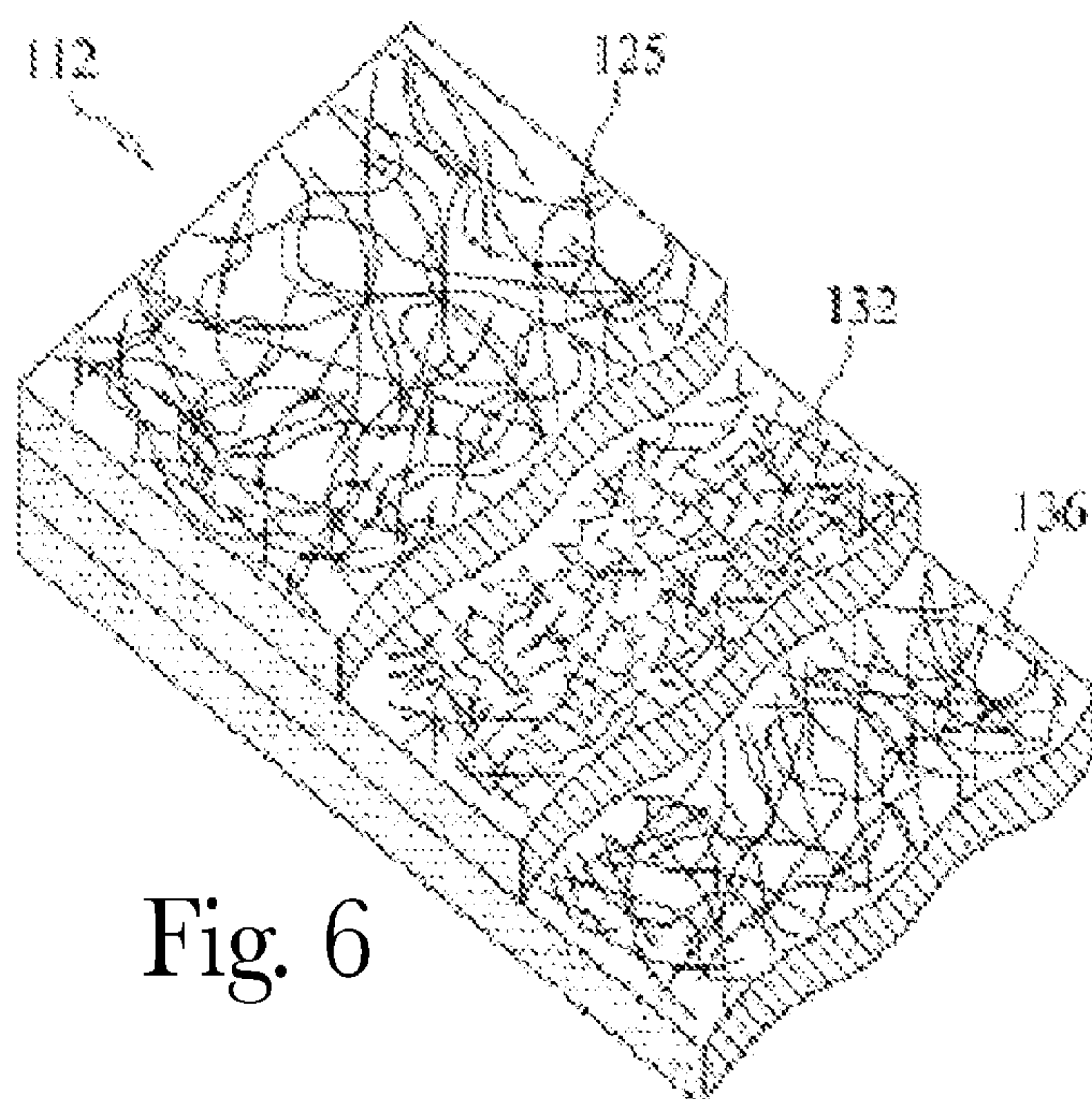


Fig. 6

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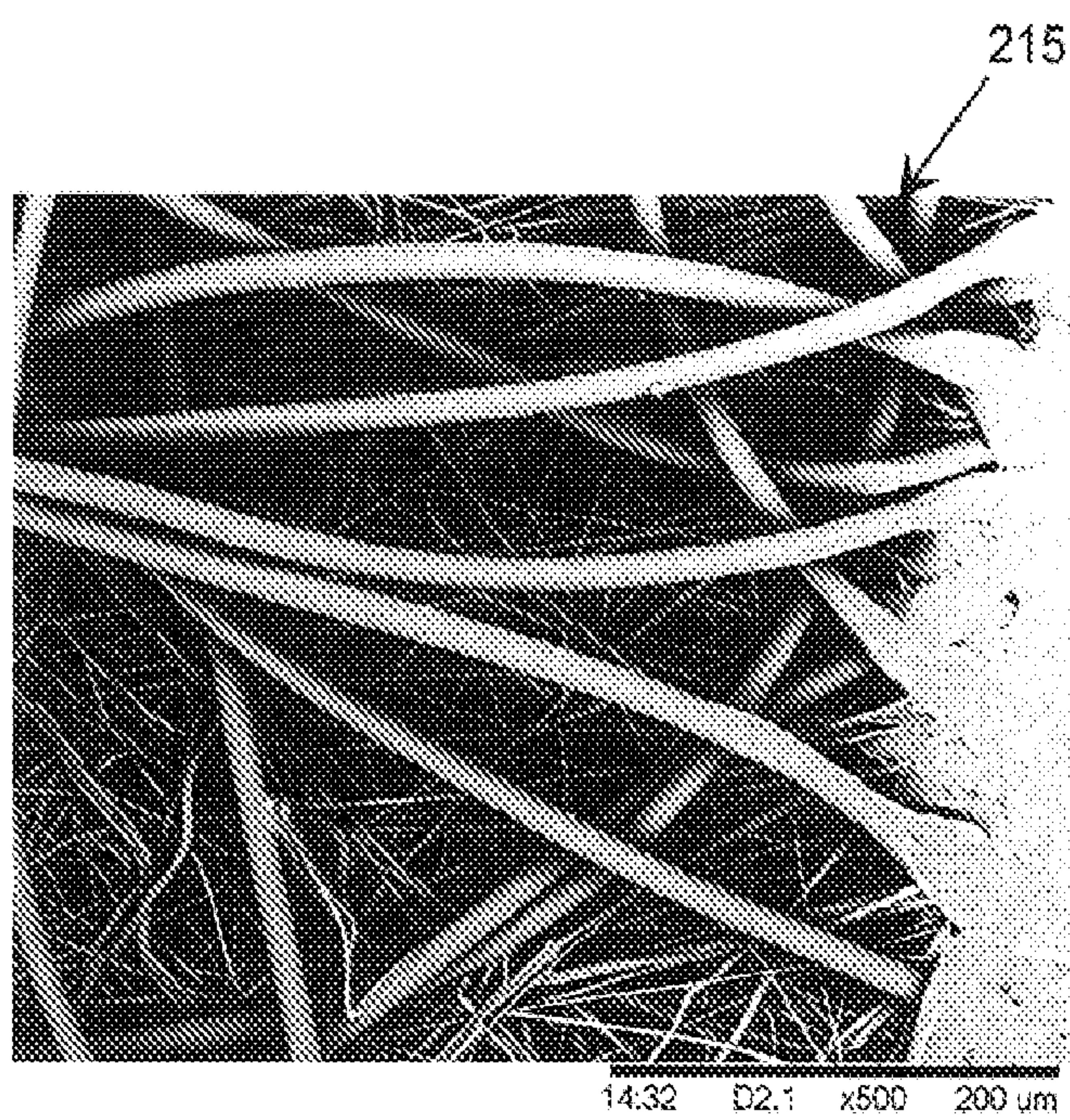


Fig. 7

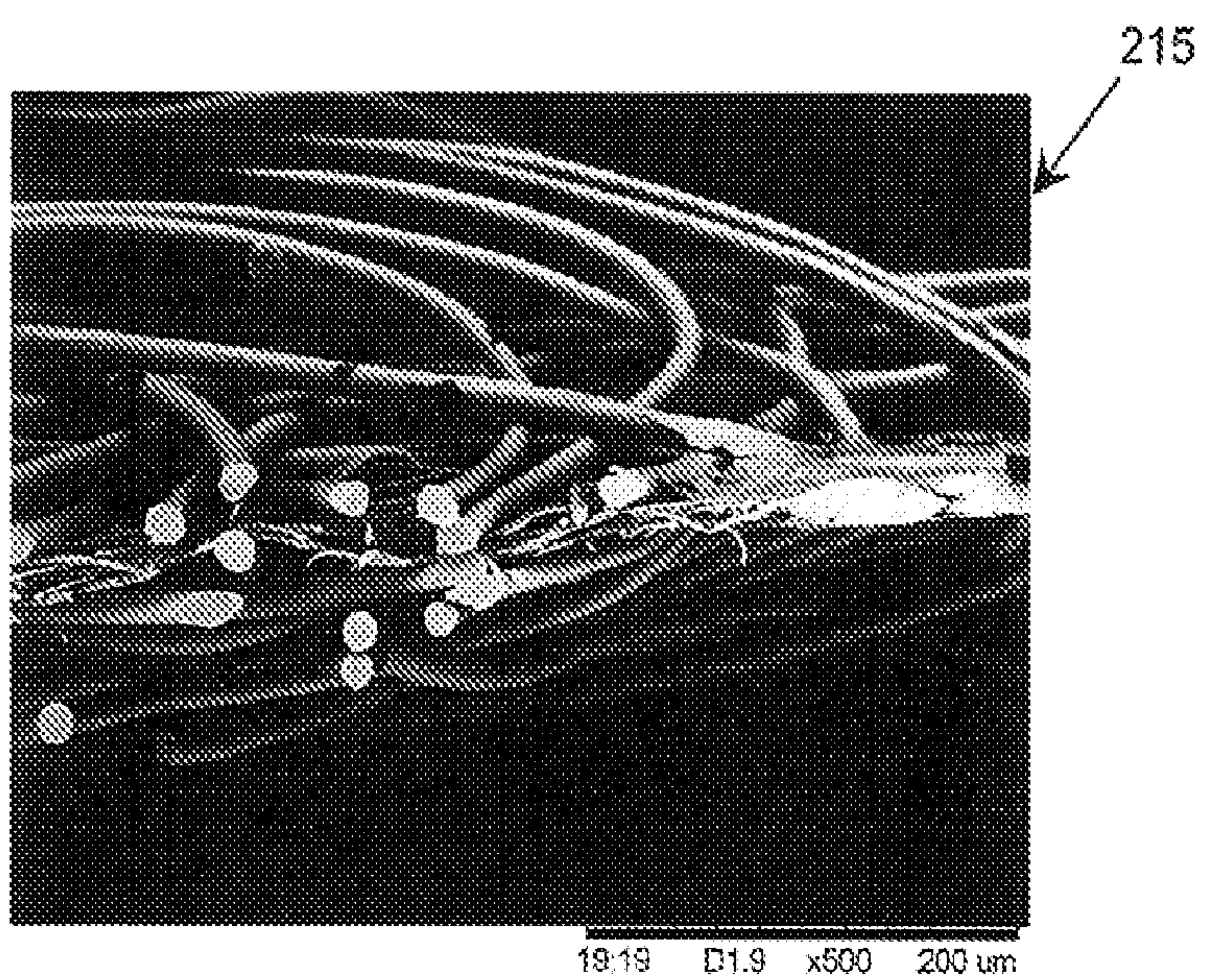


Fig. 8

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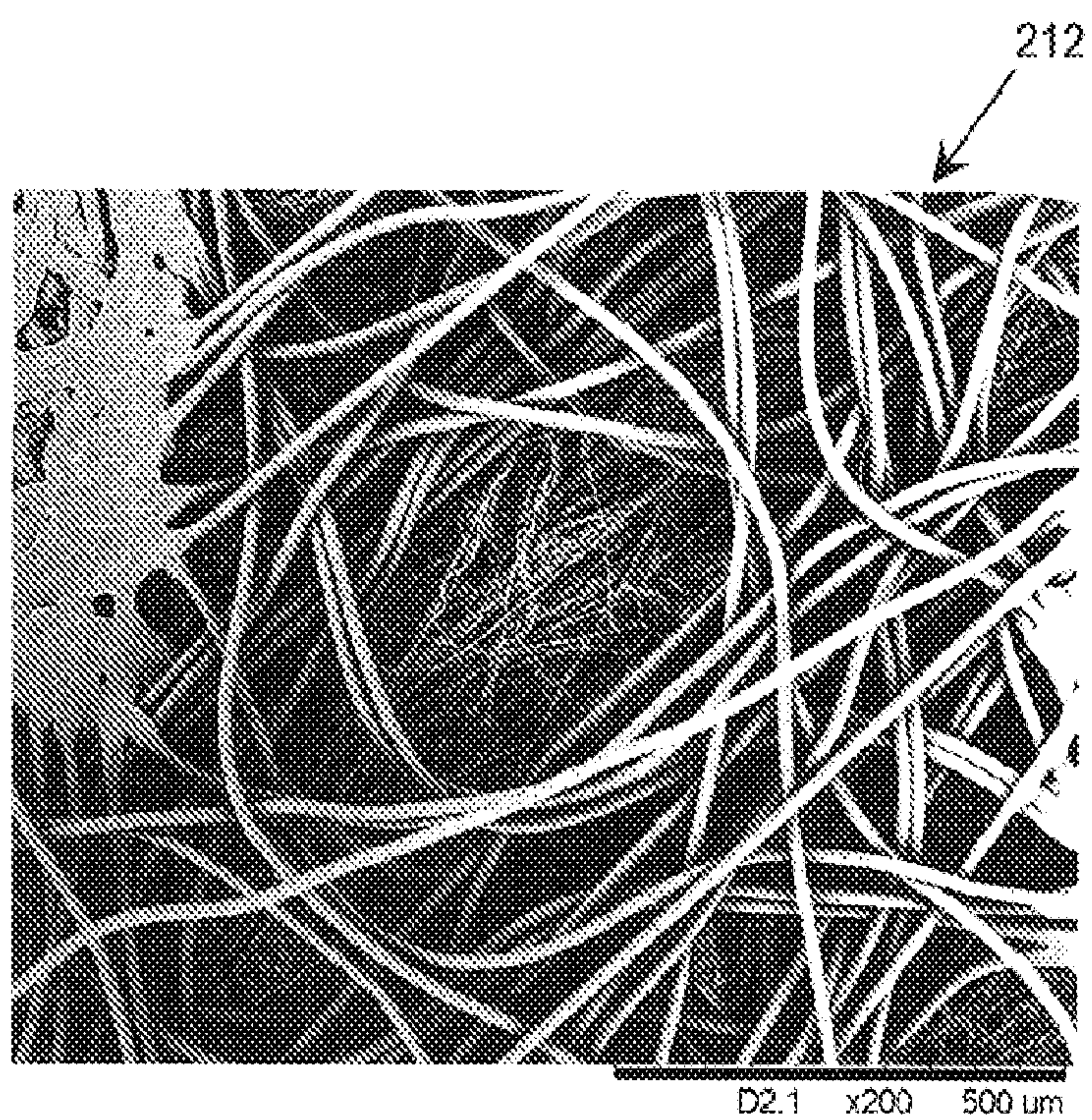


Fig. 9

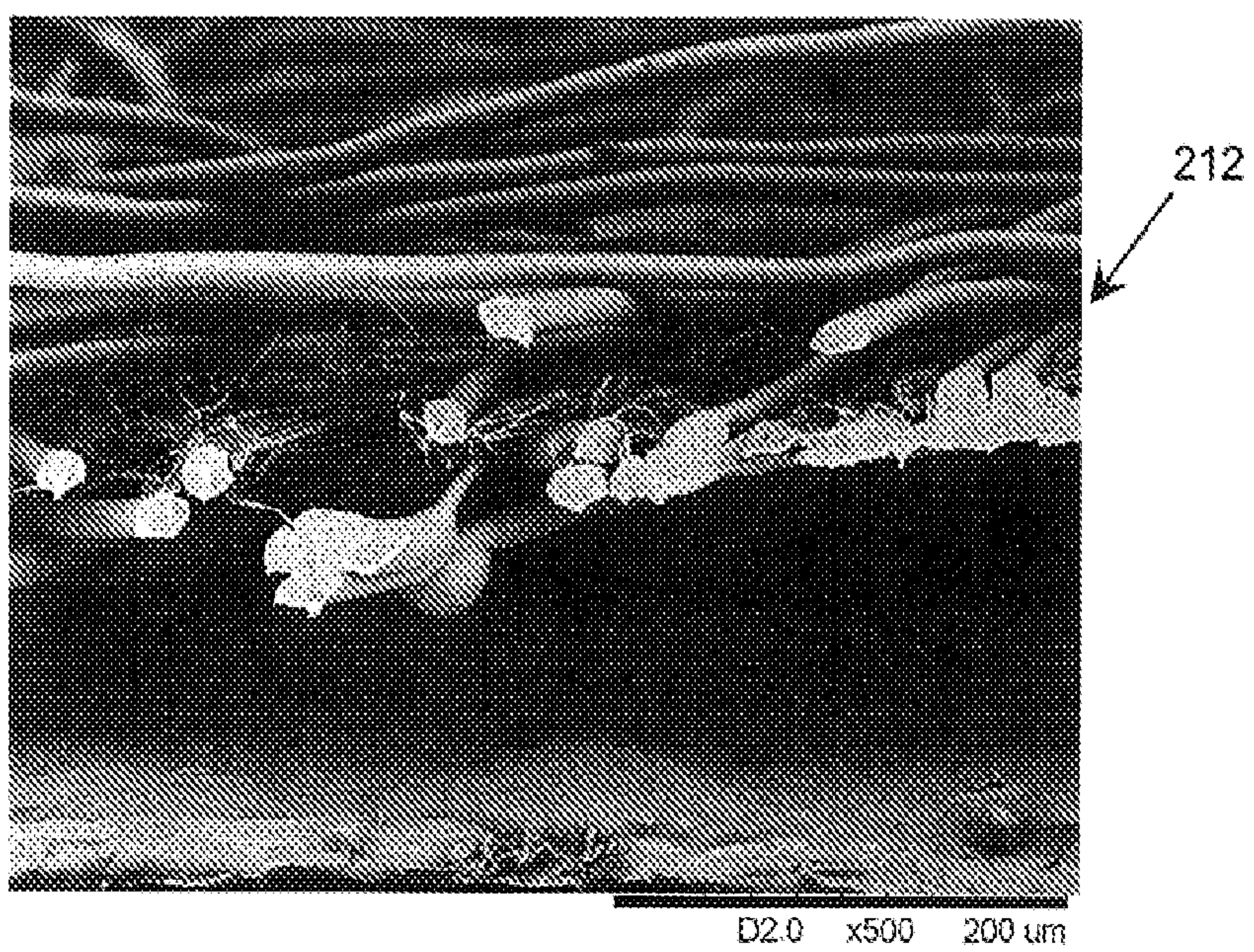


Fig. 10

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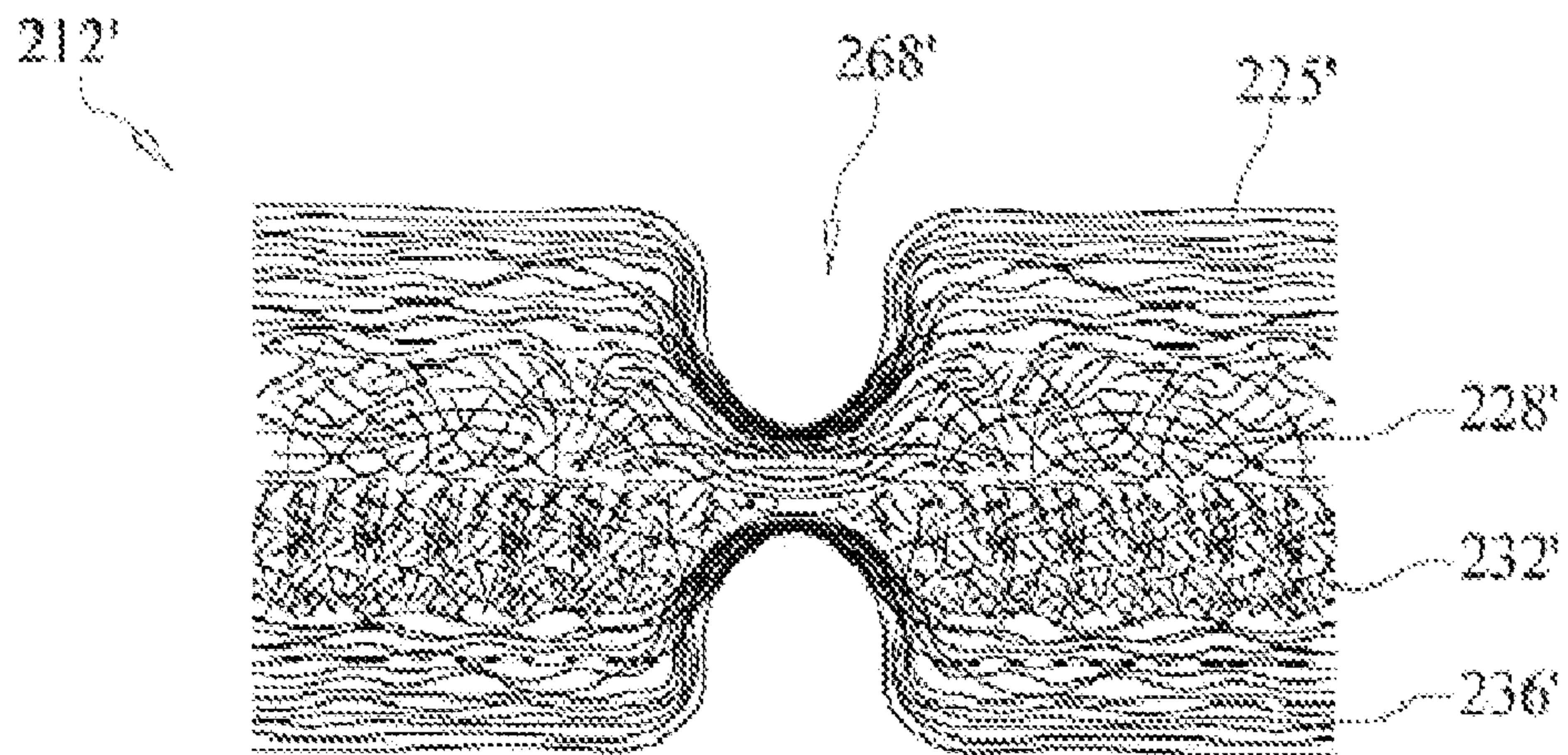


Fig. 11

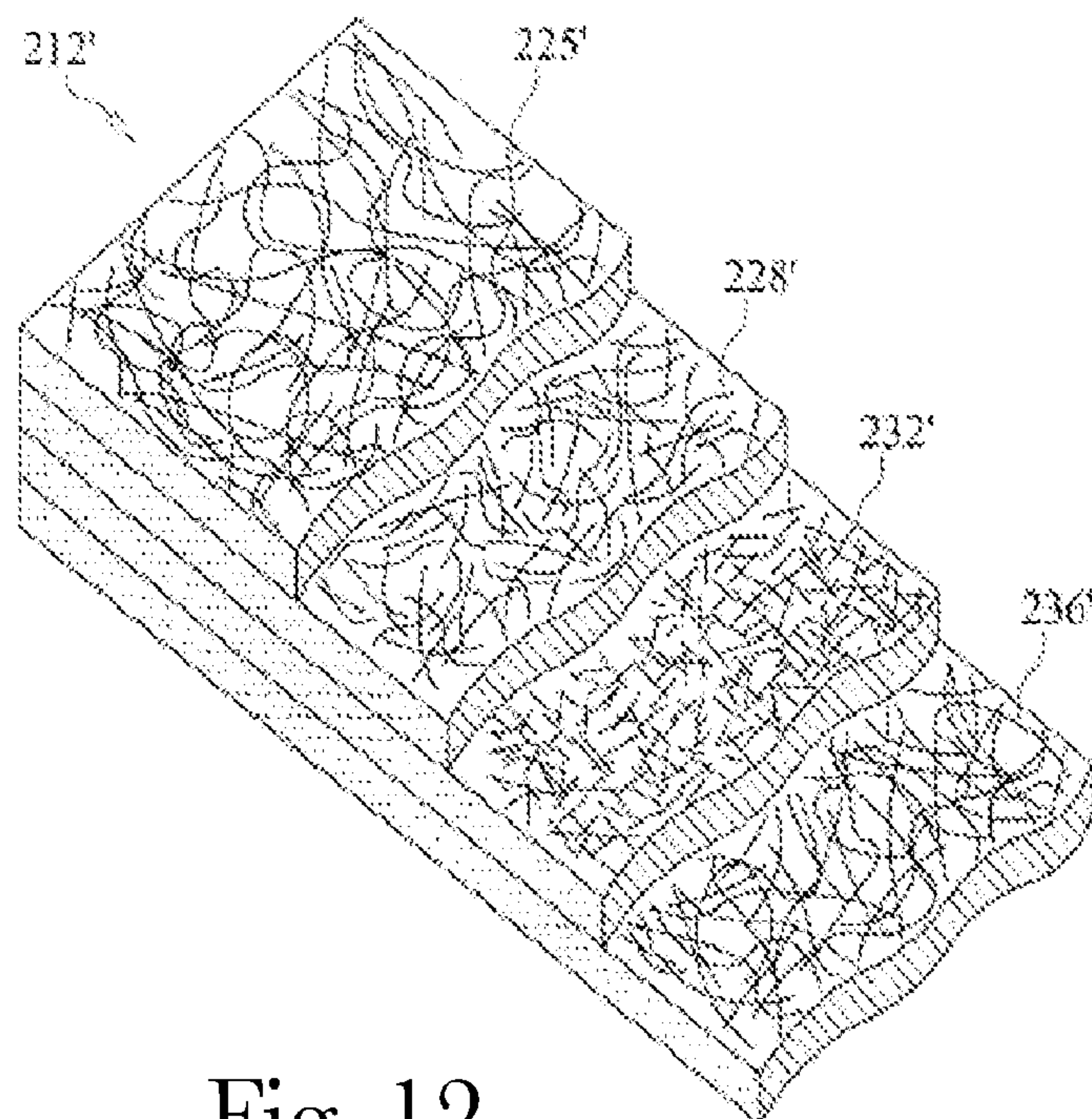


Fig. 12

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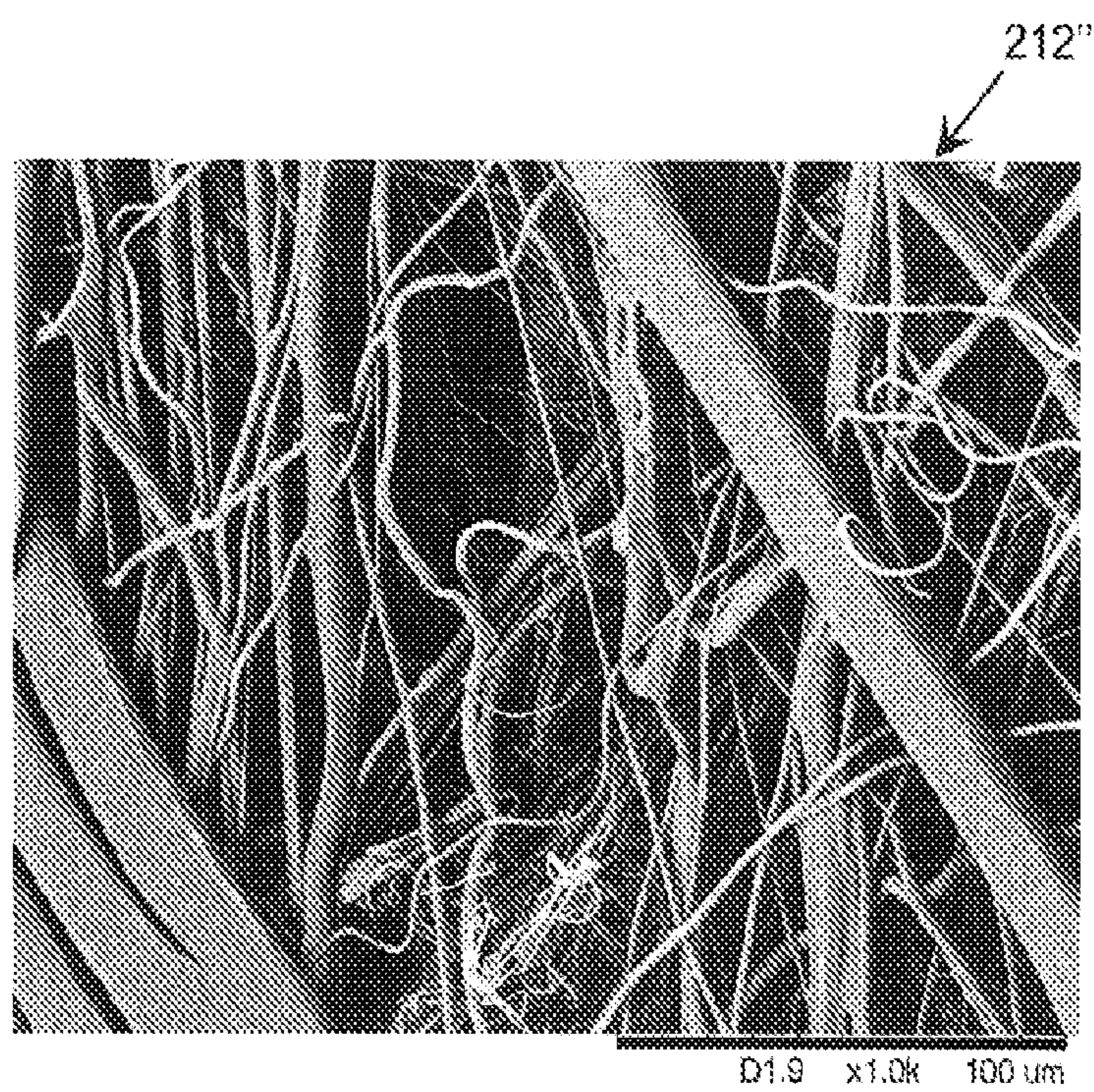


Fig. 13

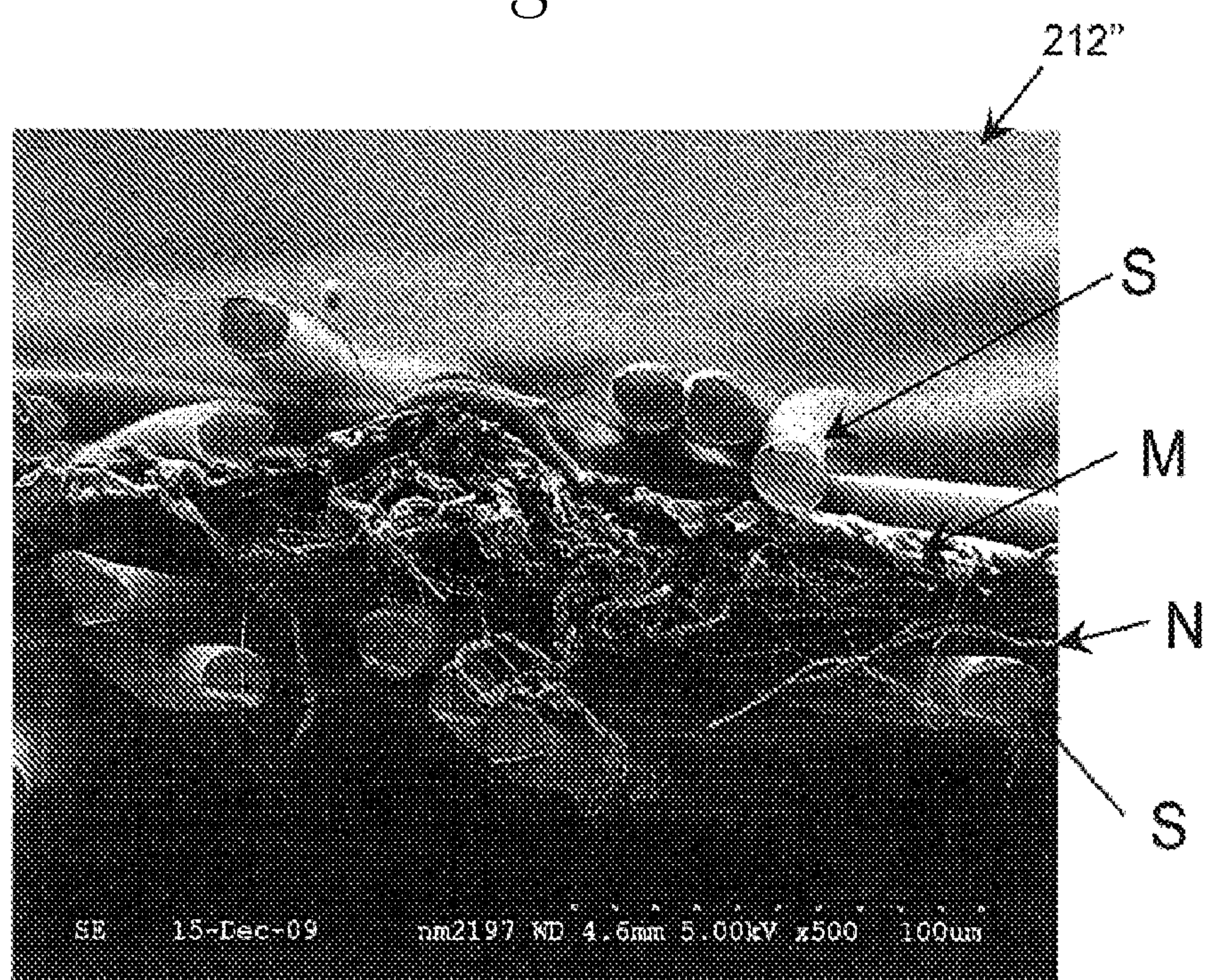


Fig. 14

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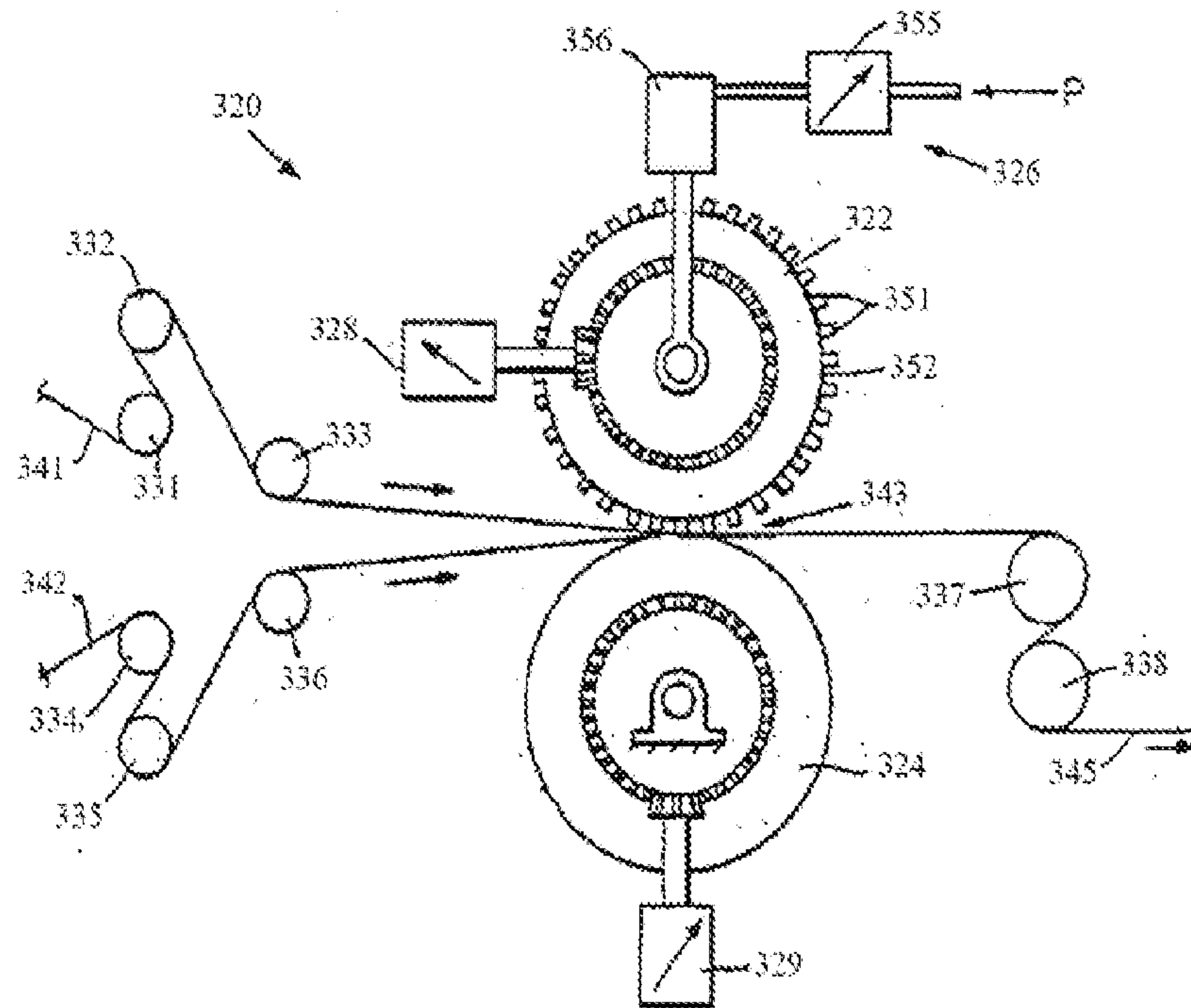


Fig. 15

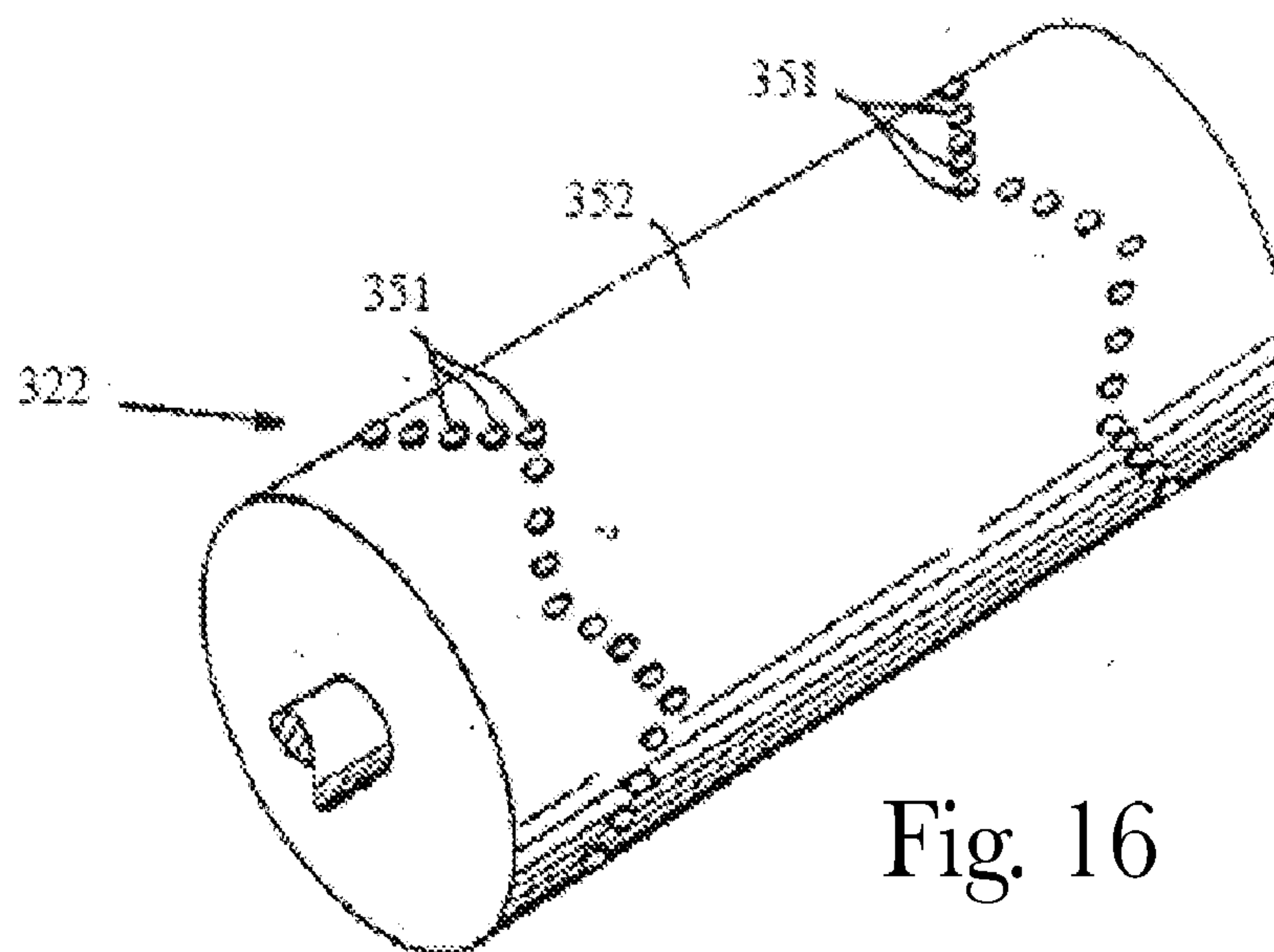


Fig. 16

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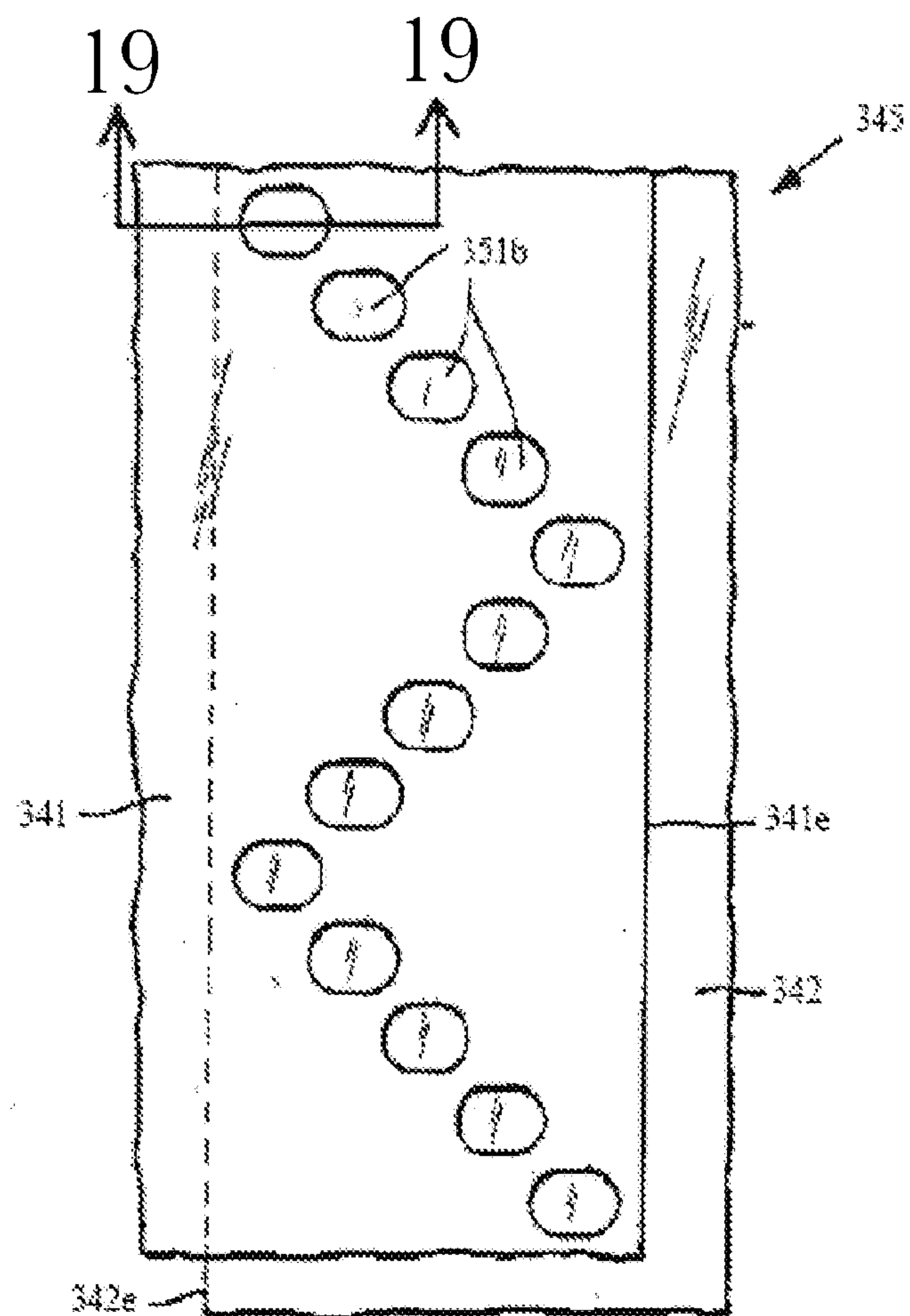
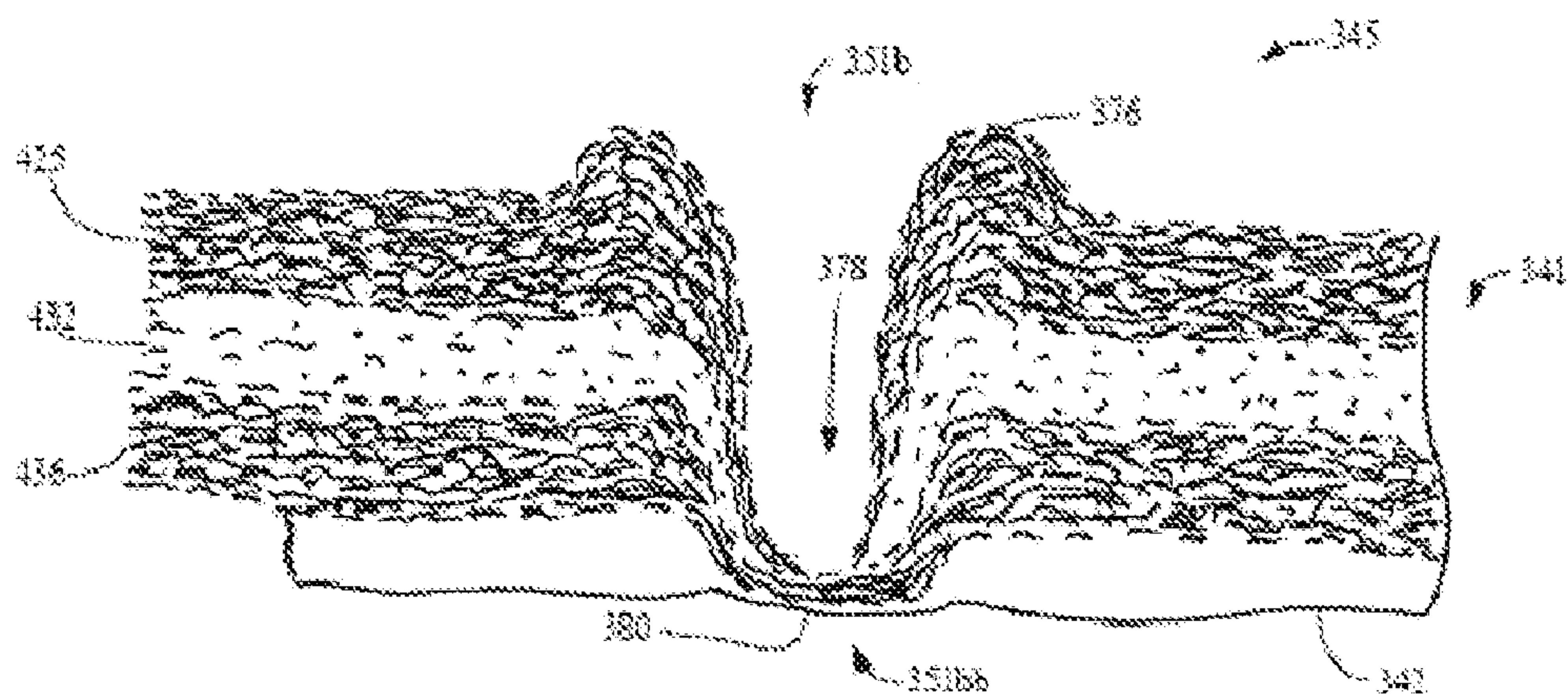
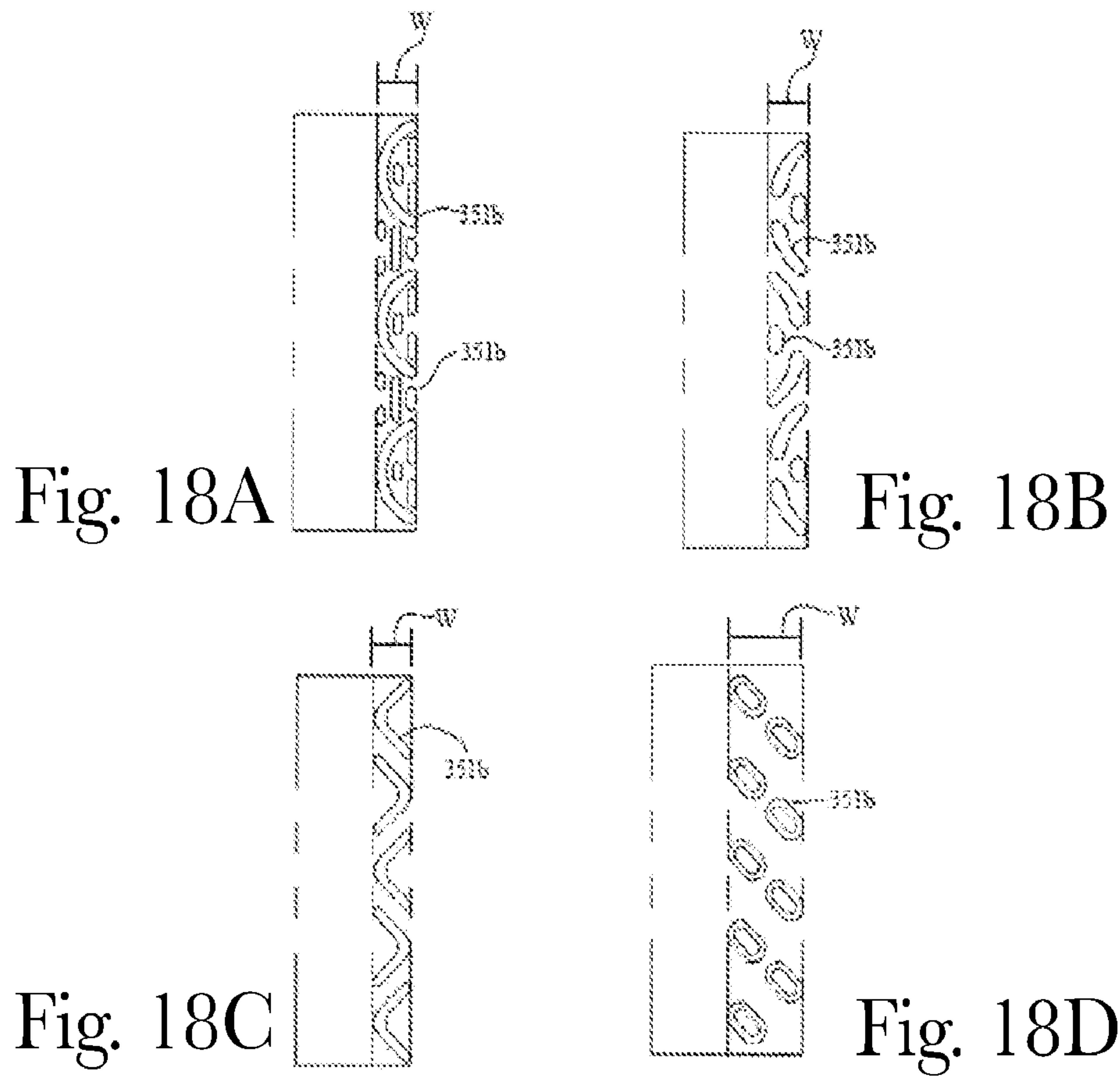


Fig. 17

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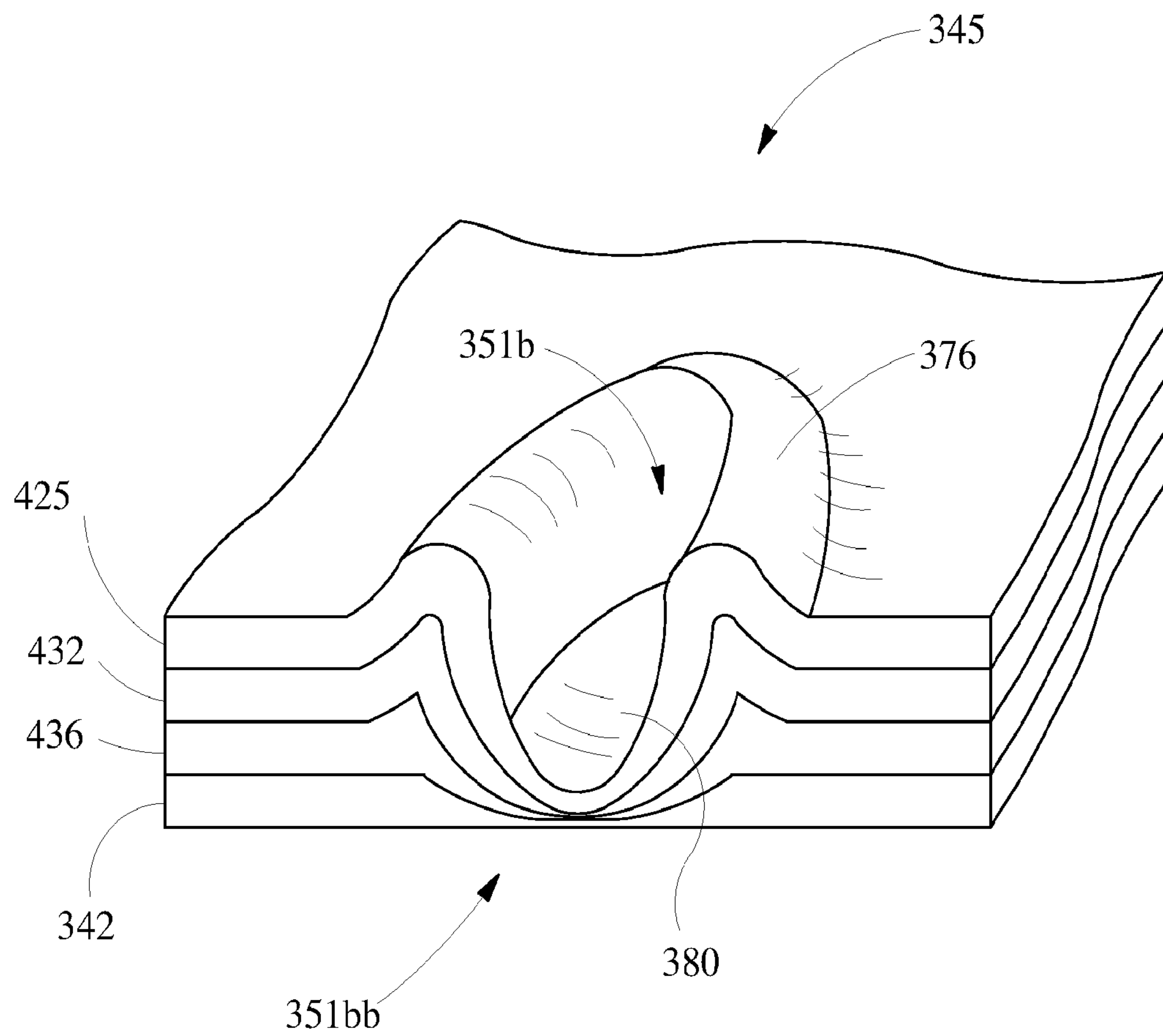


Fig. 20

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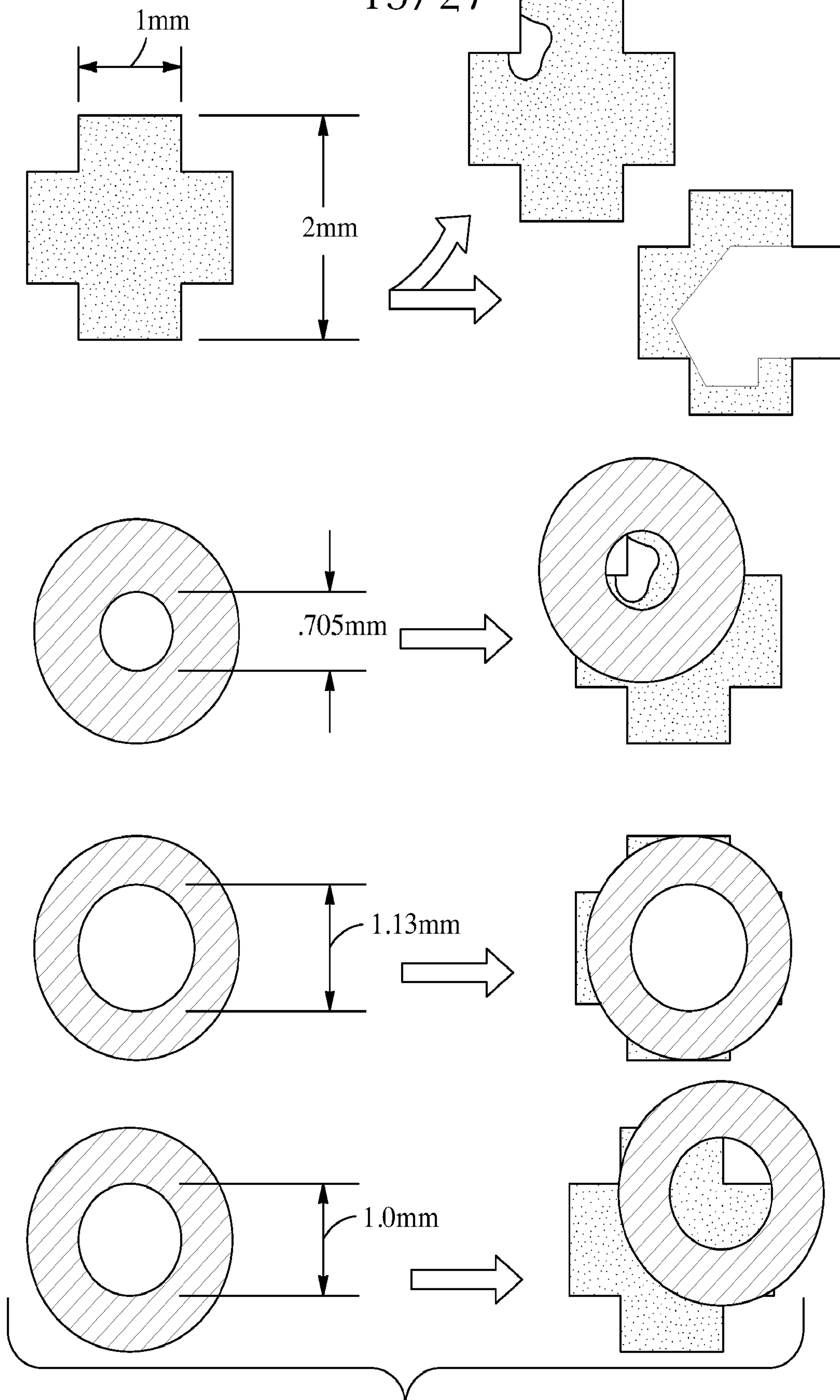


Fig. 21A

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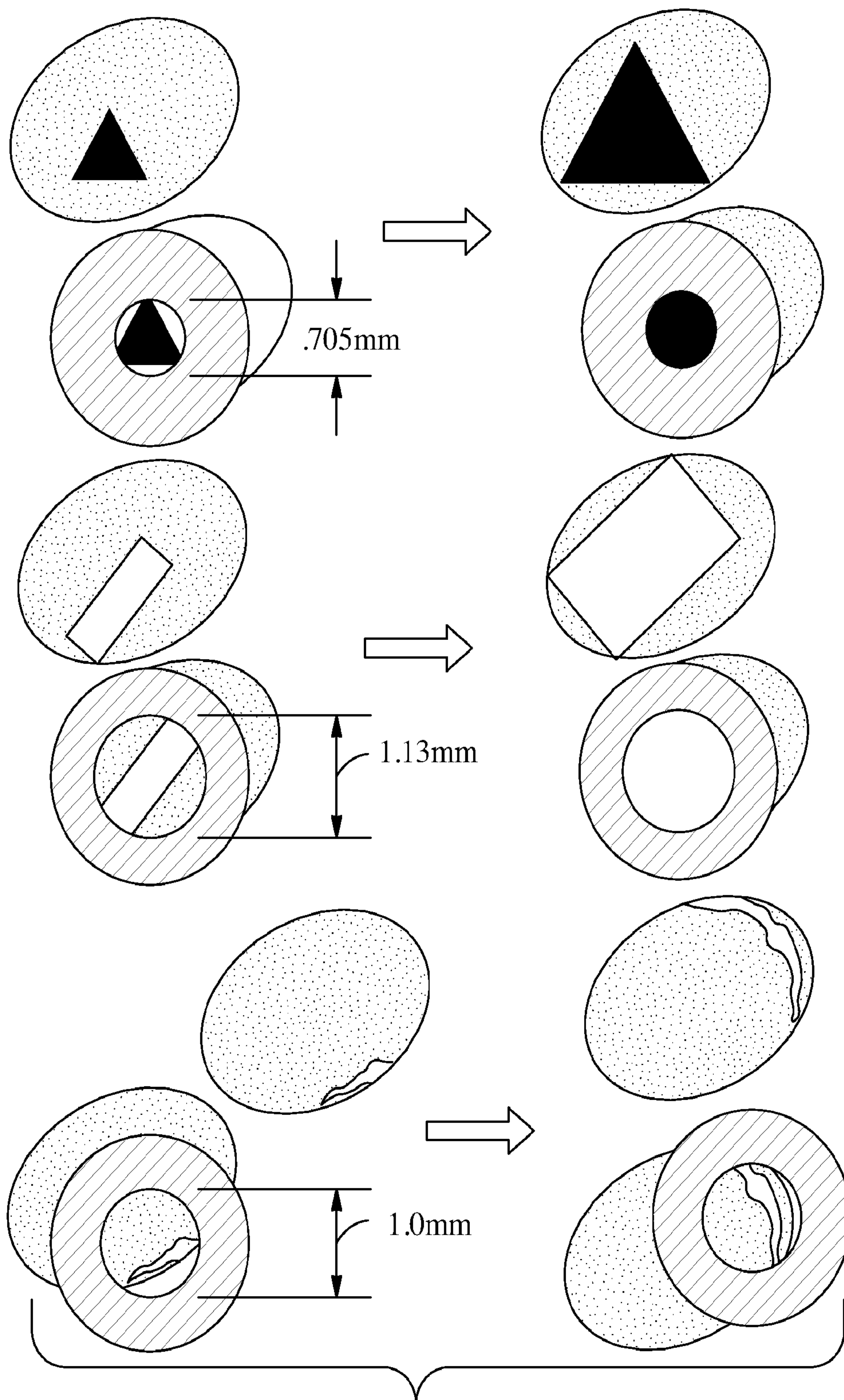


Fig. 21B

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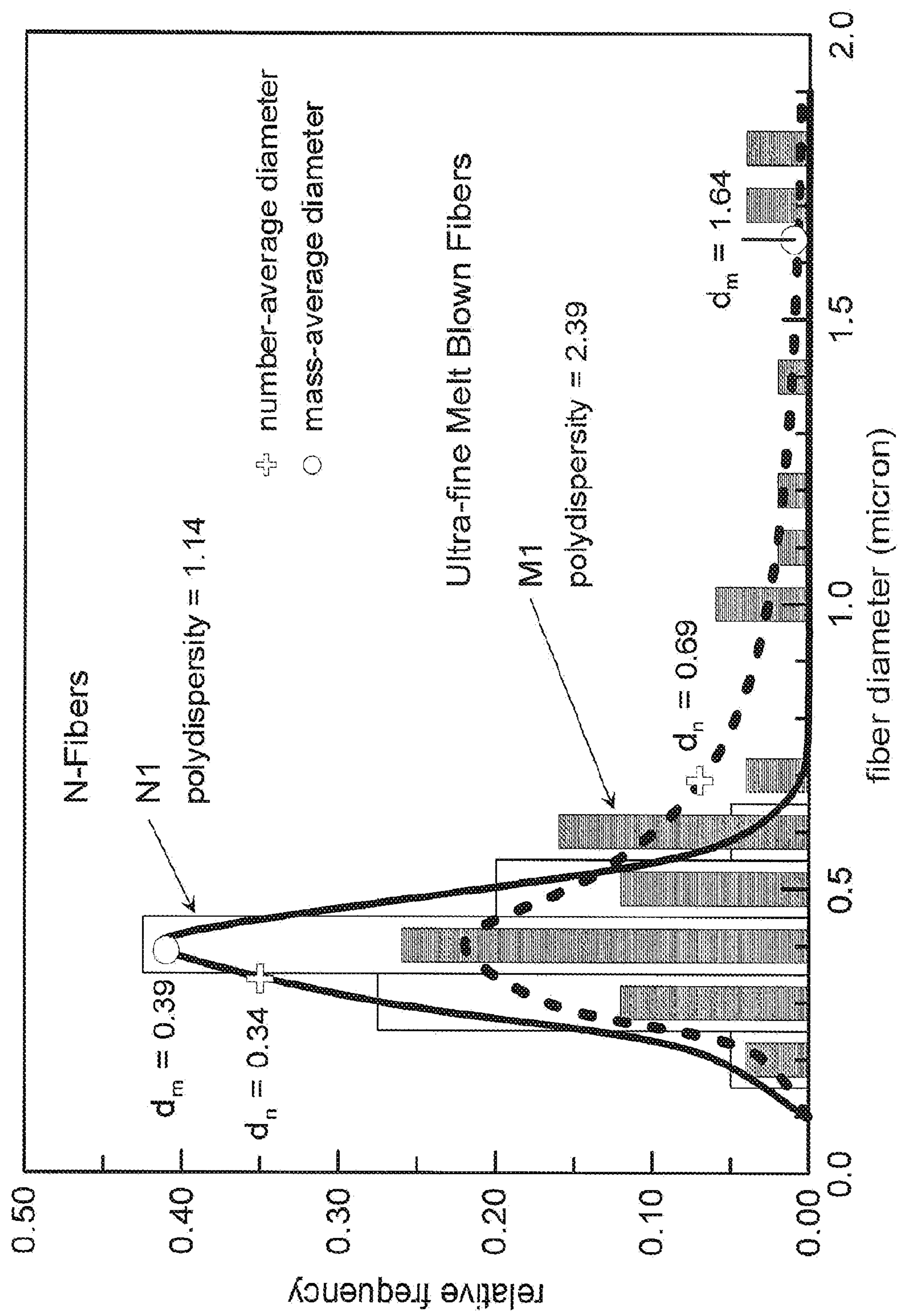


Fig. 22

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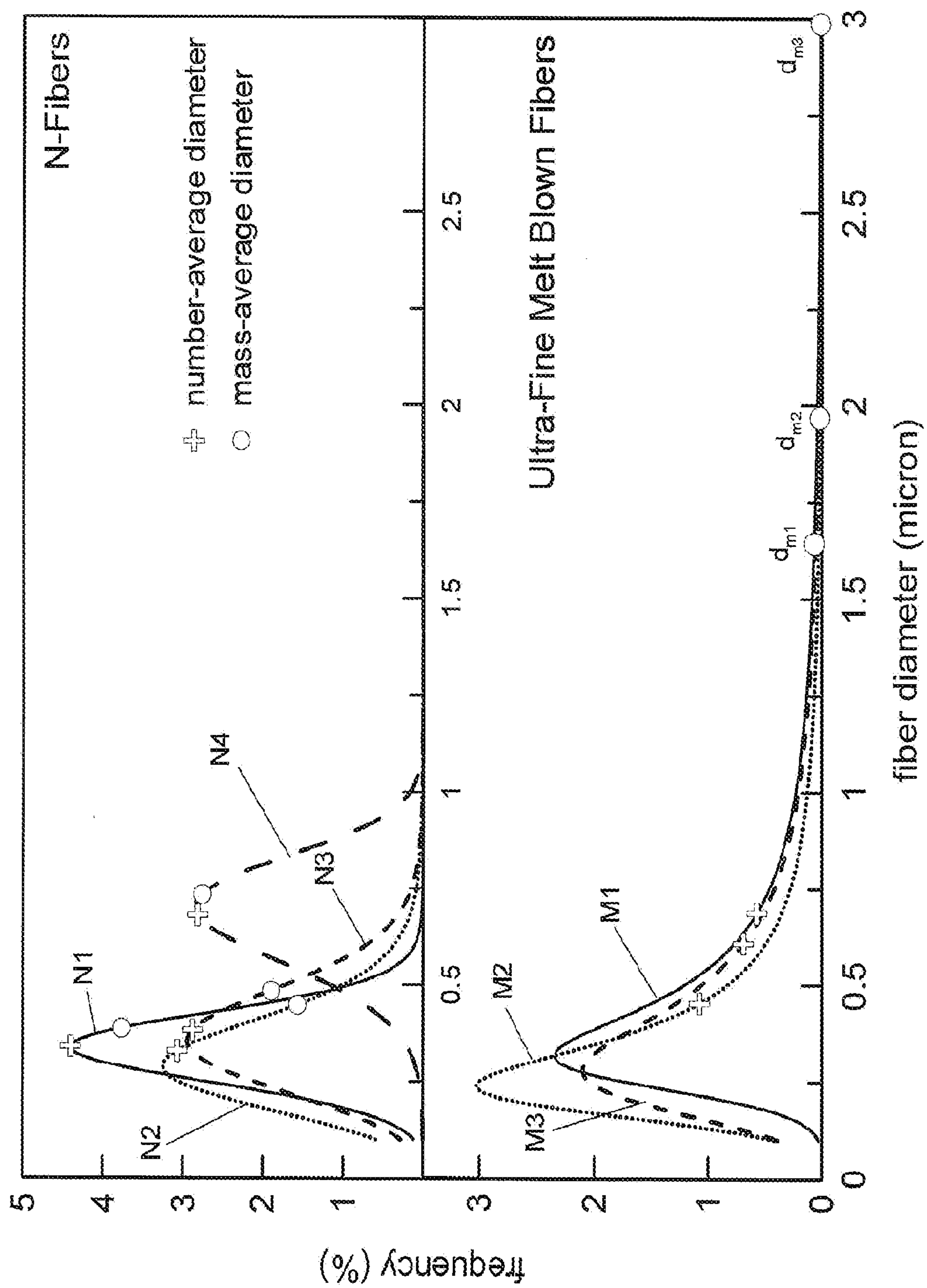


Fig. 23

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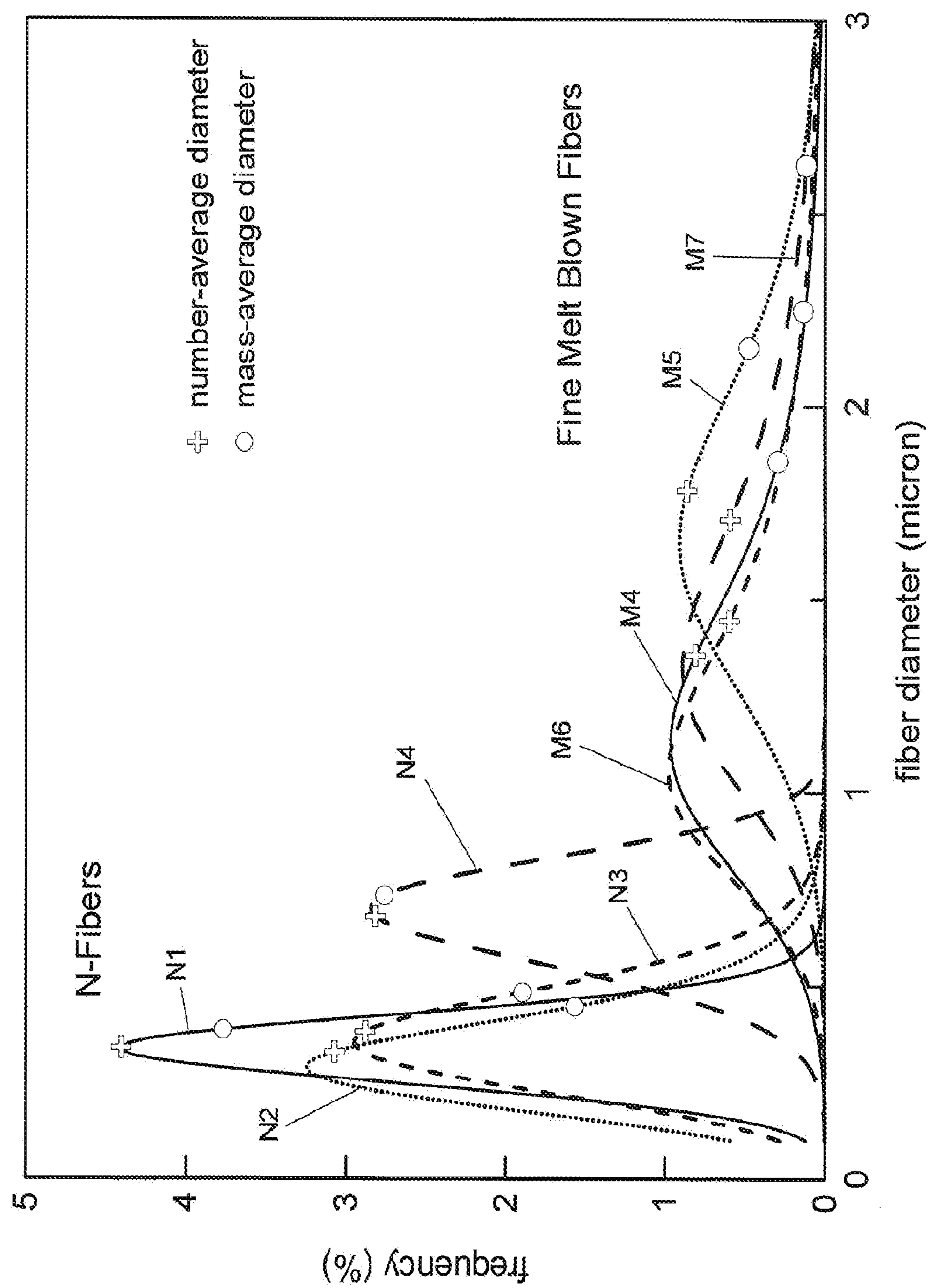


Fig. 24

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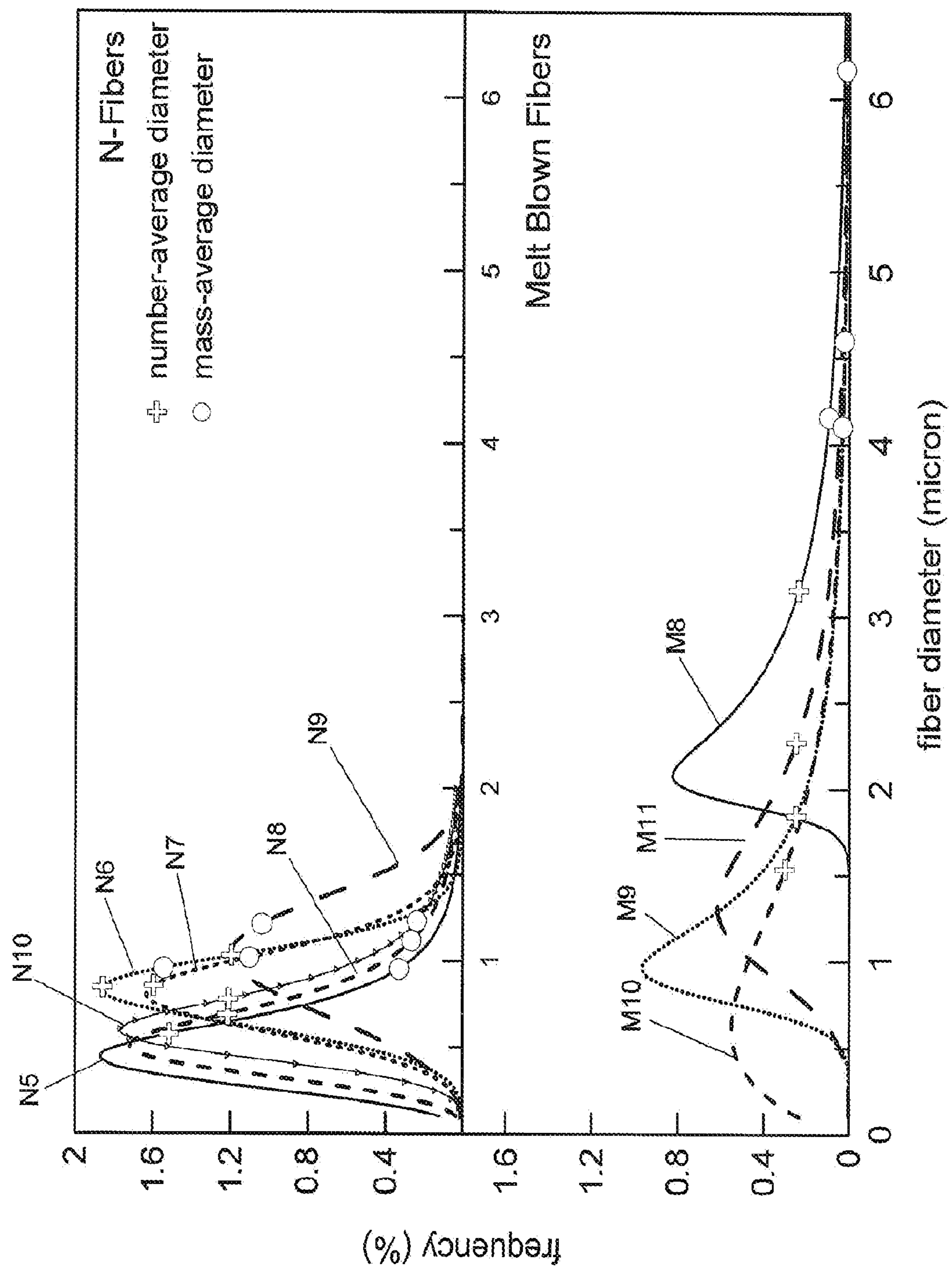


Fig. 25

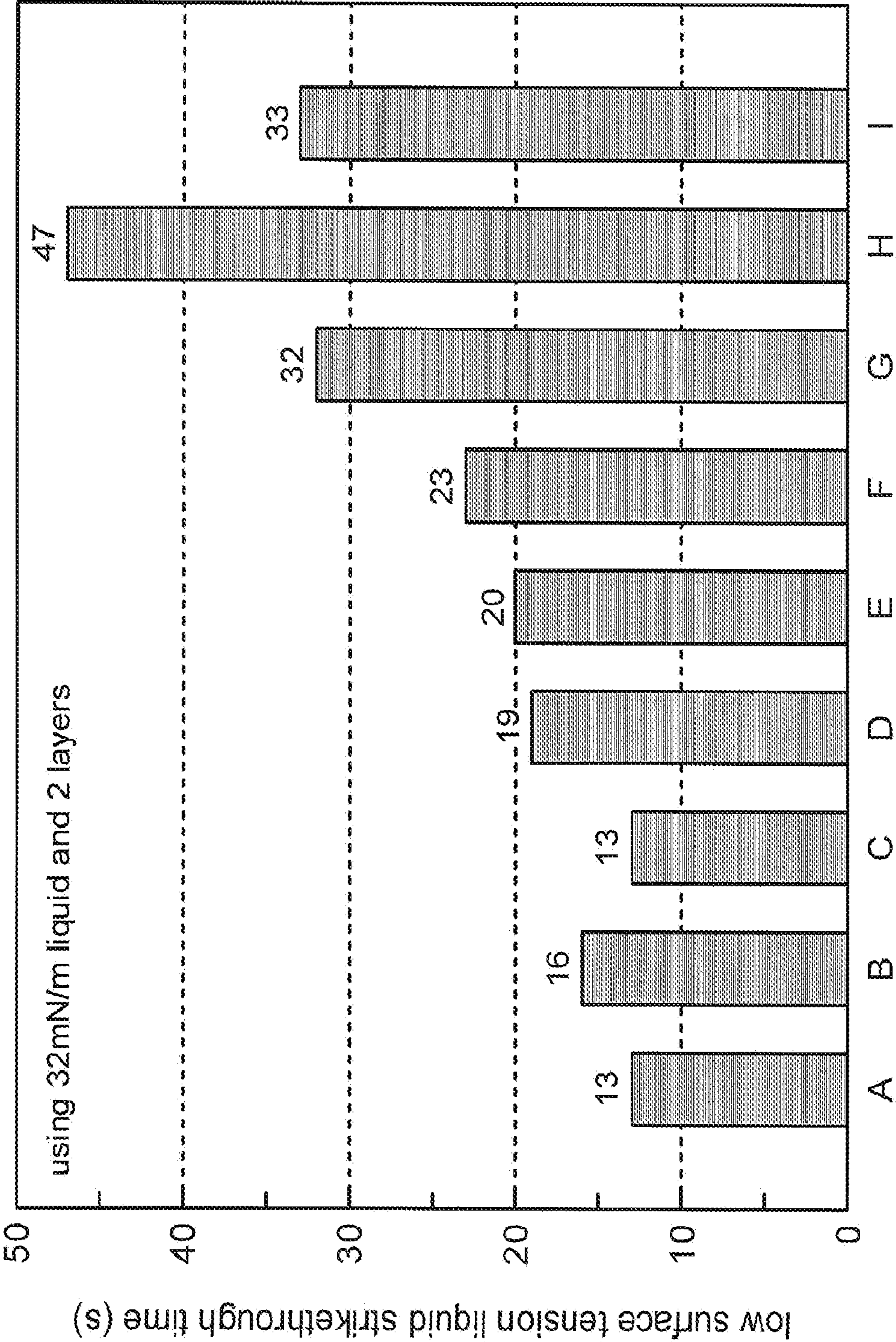


Fig. 26

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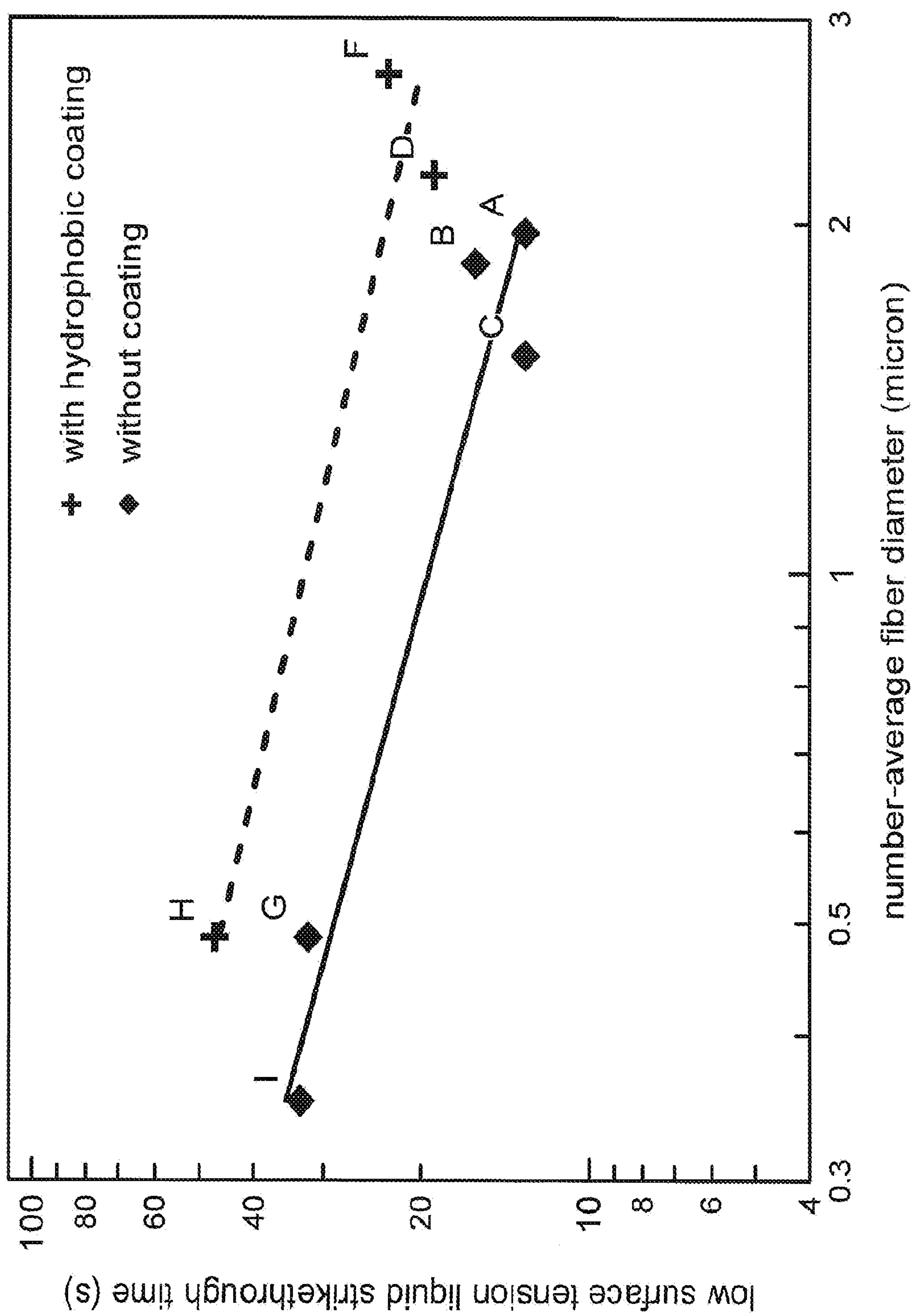


Fig. 27

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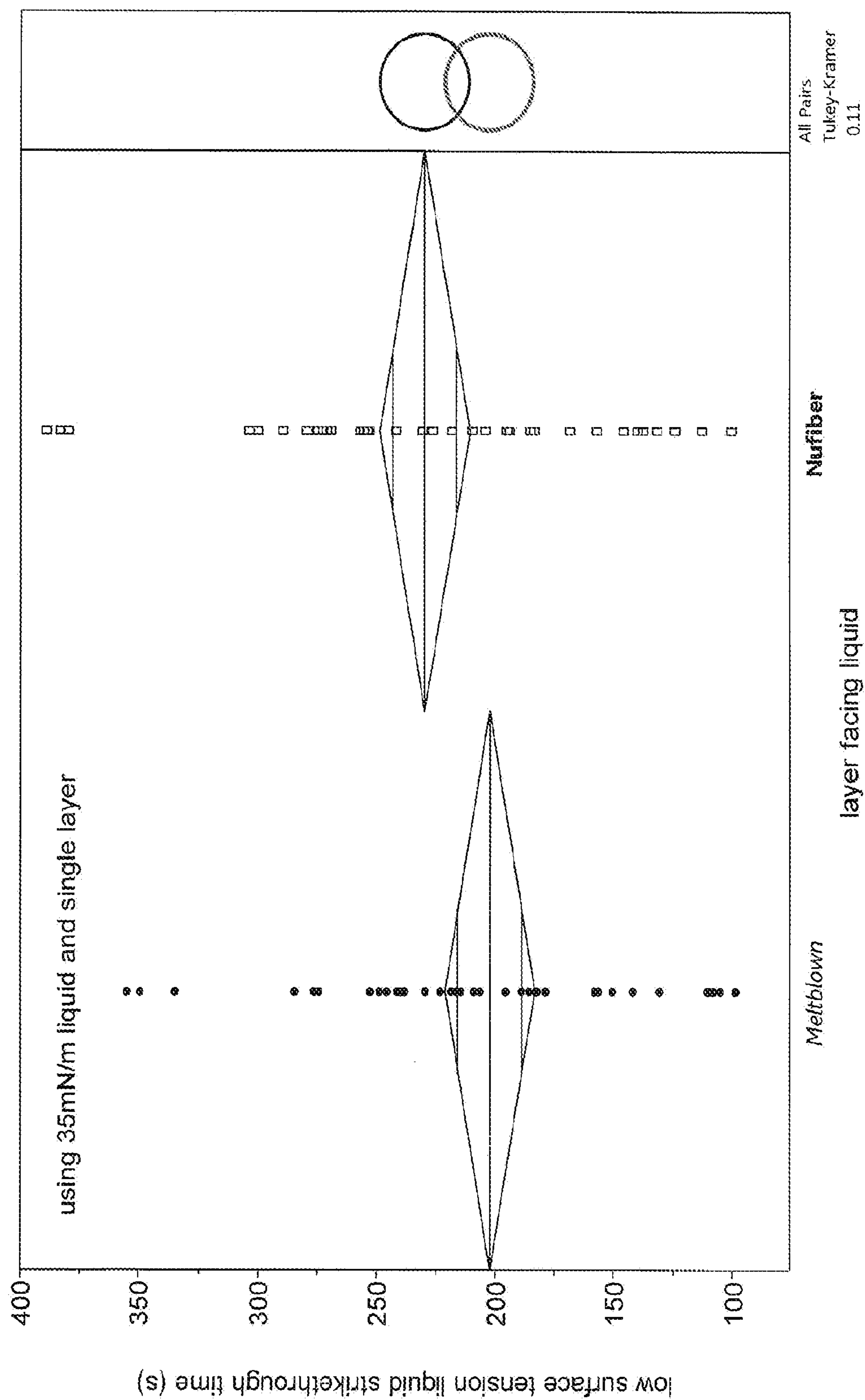


Fig. 28

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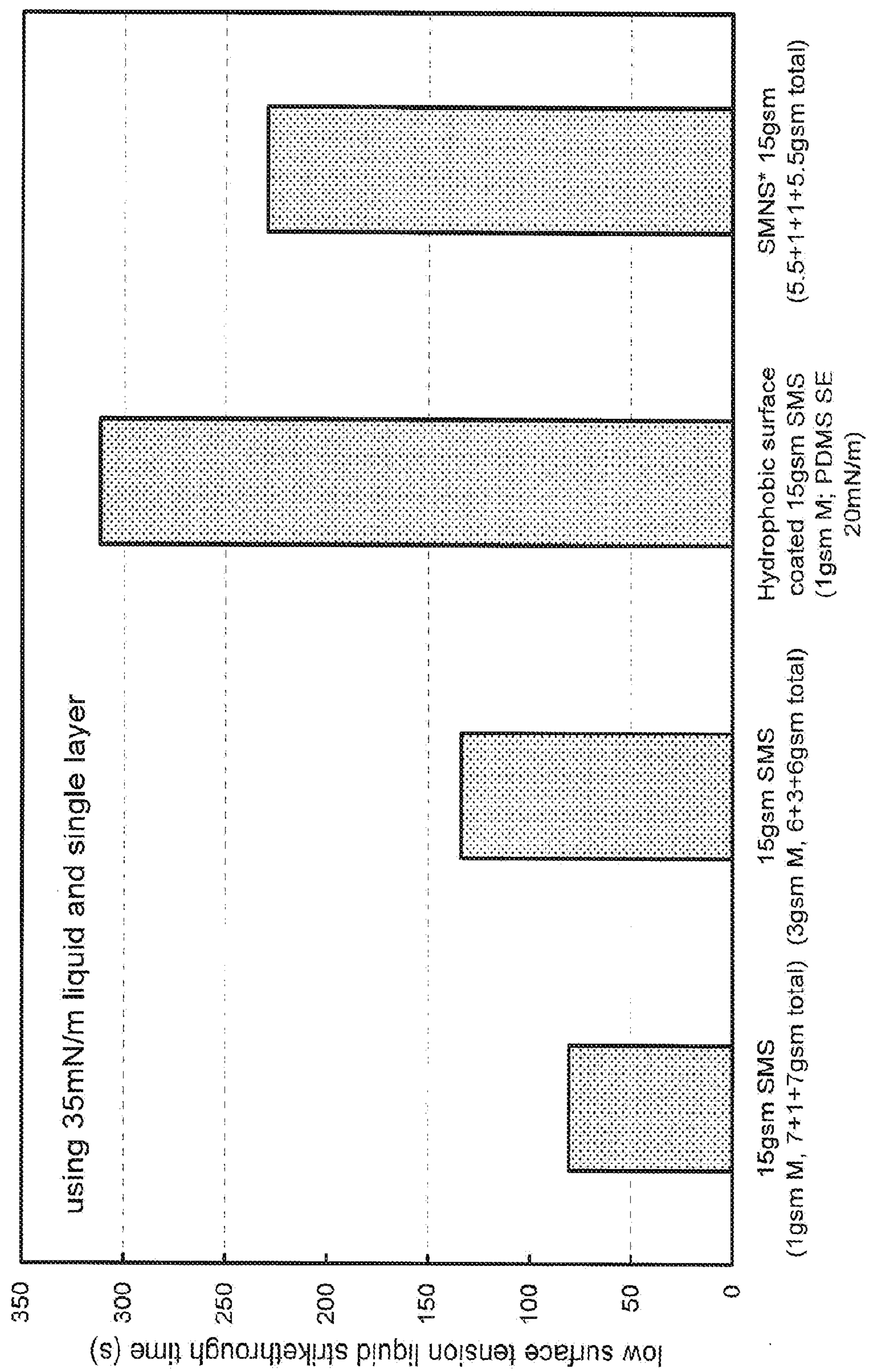


Fig. 29

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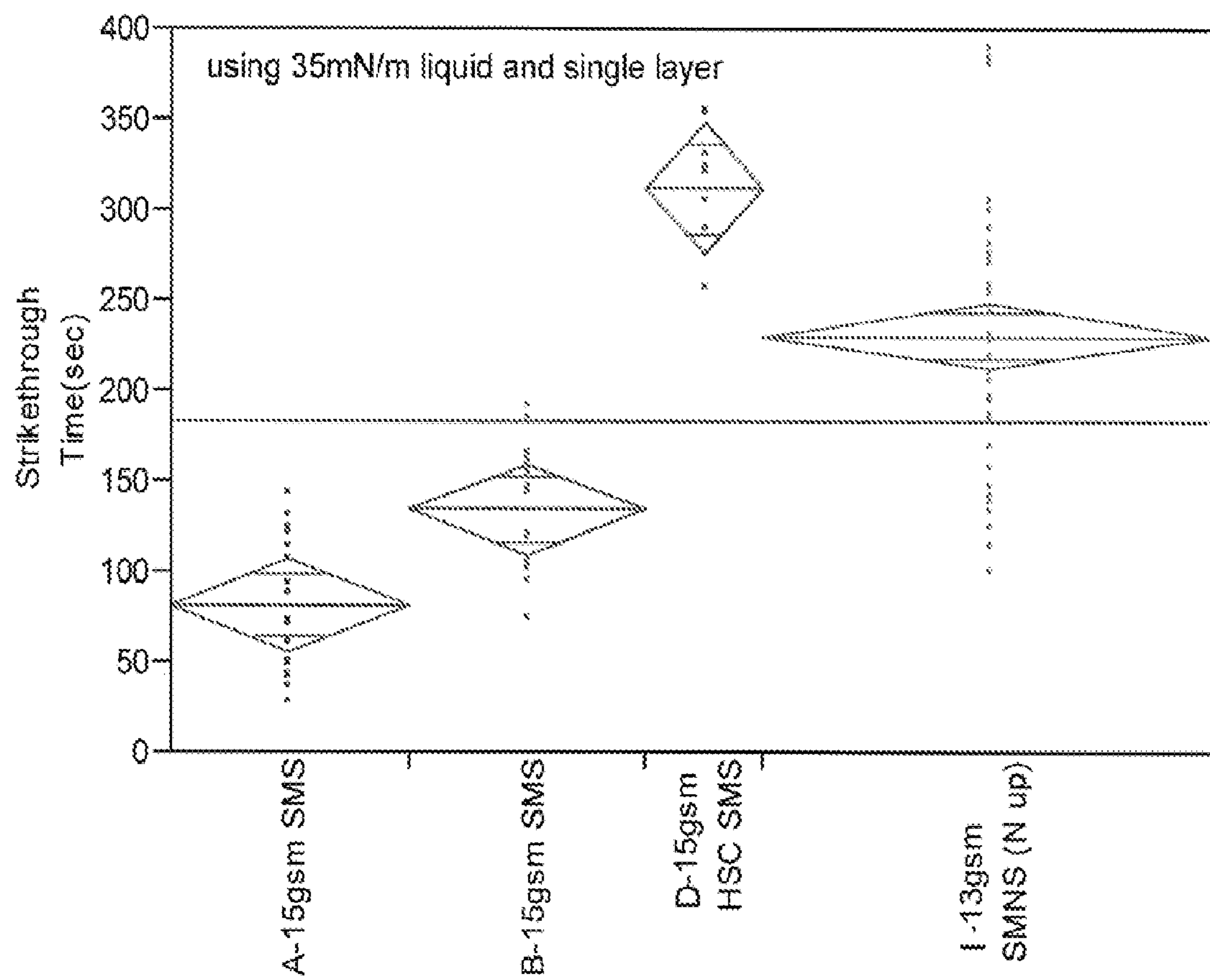


Fig. 30

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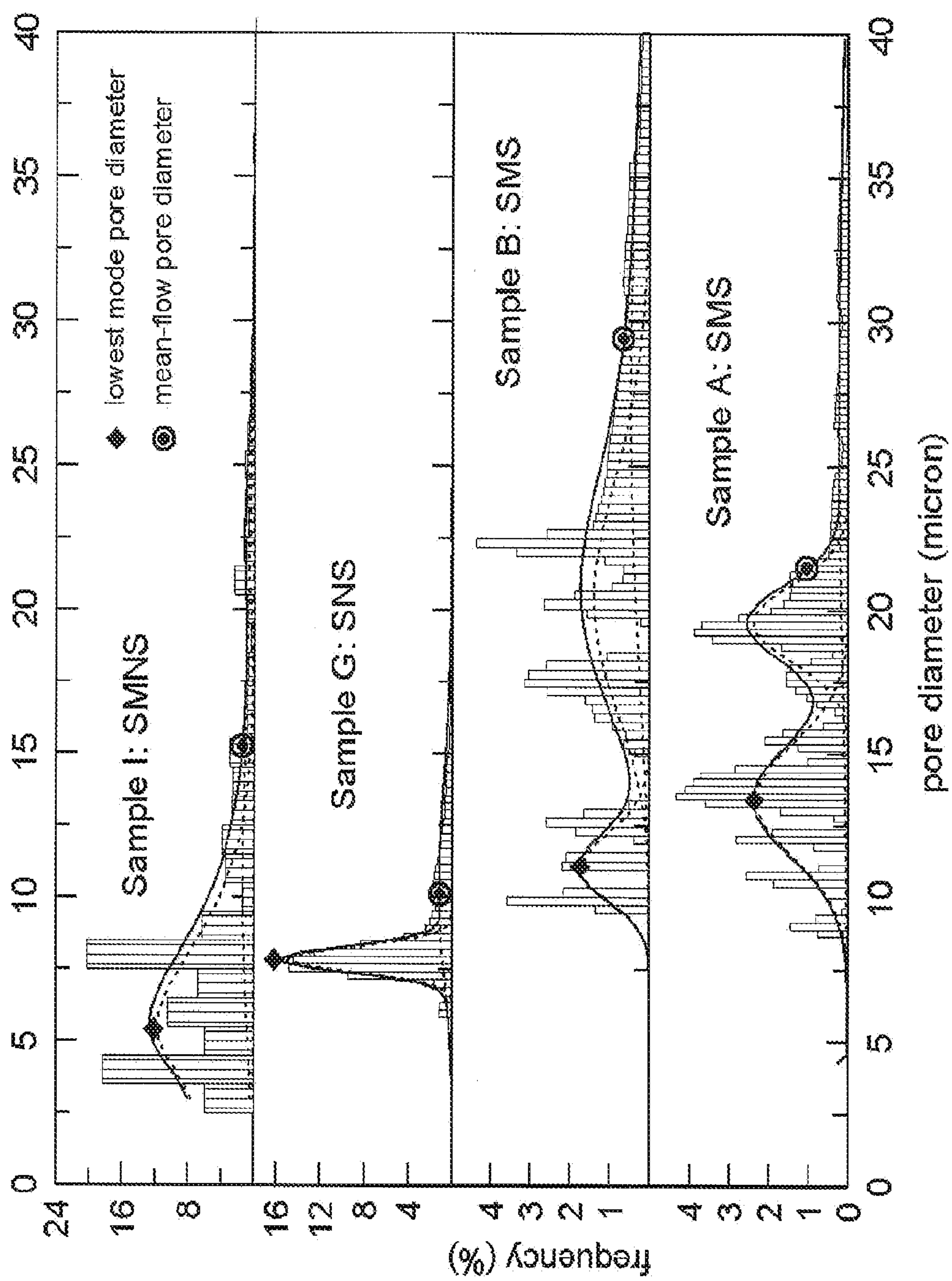


Fig. 31

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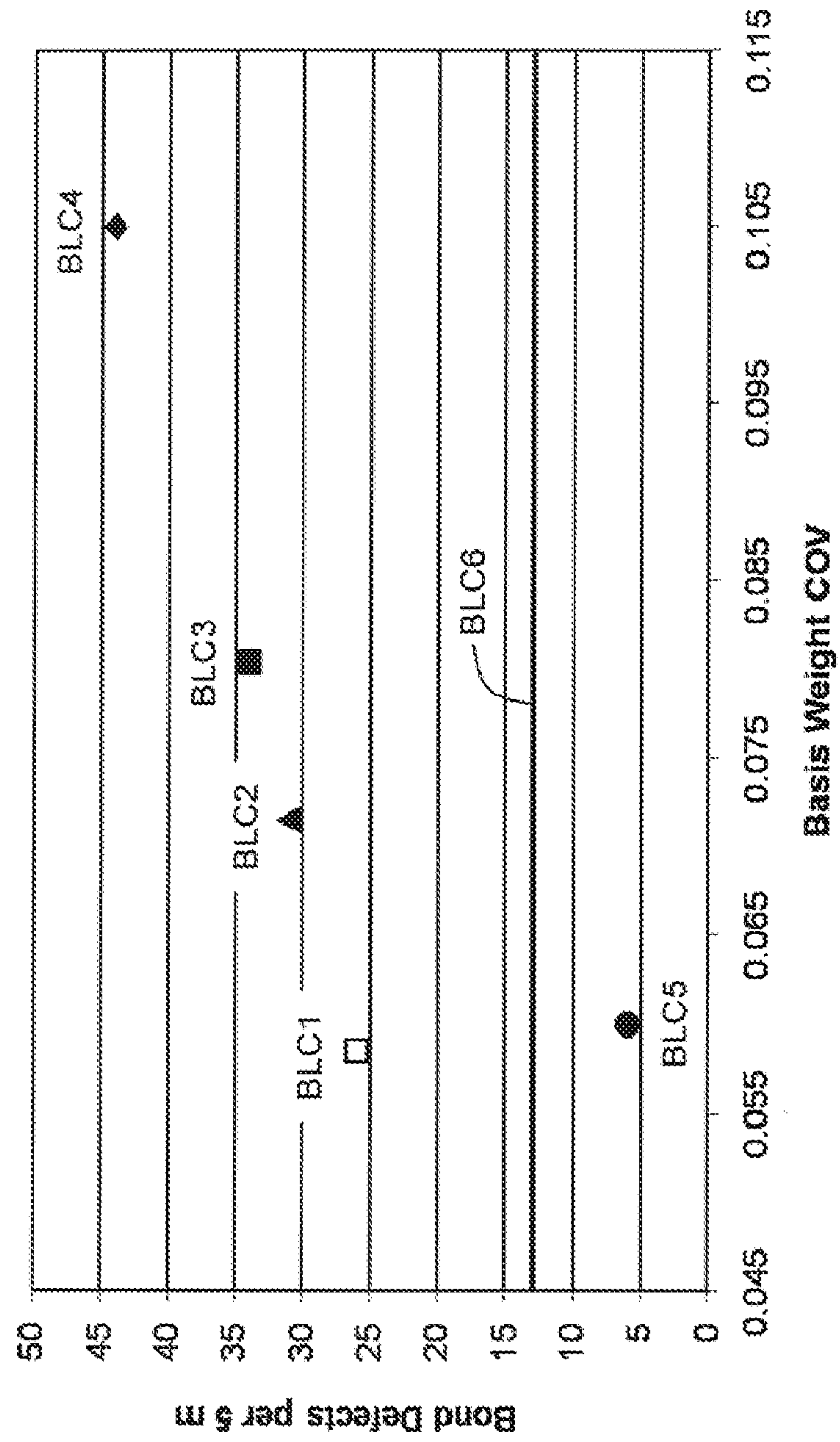


Fig. 32

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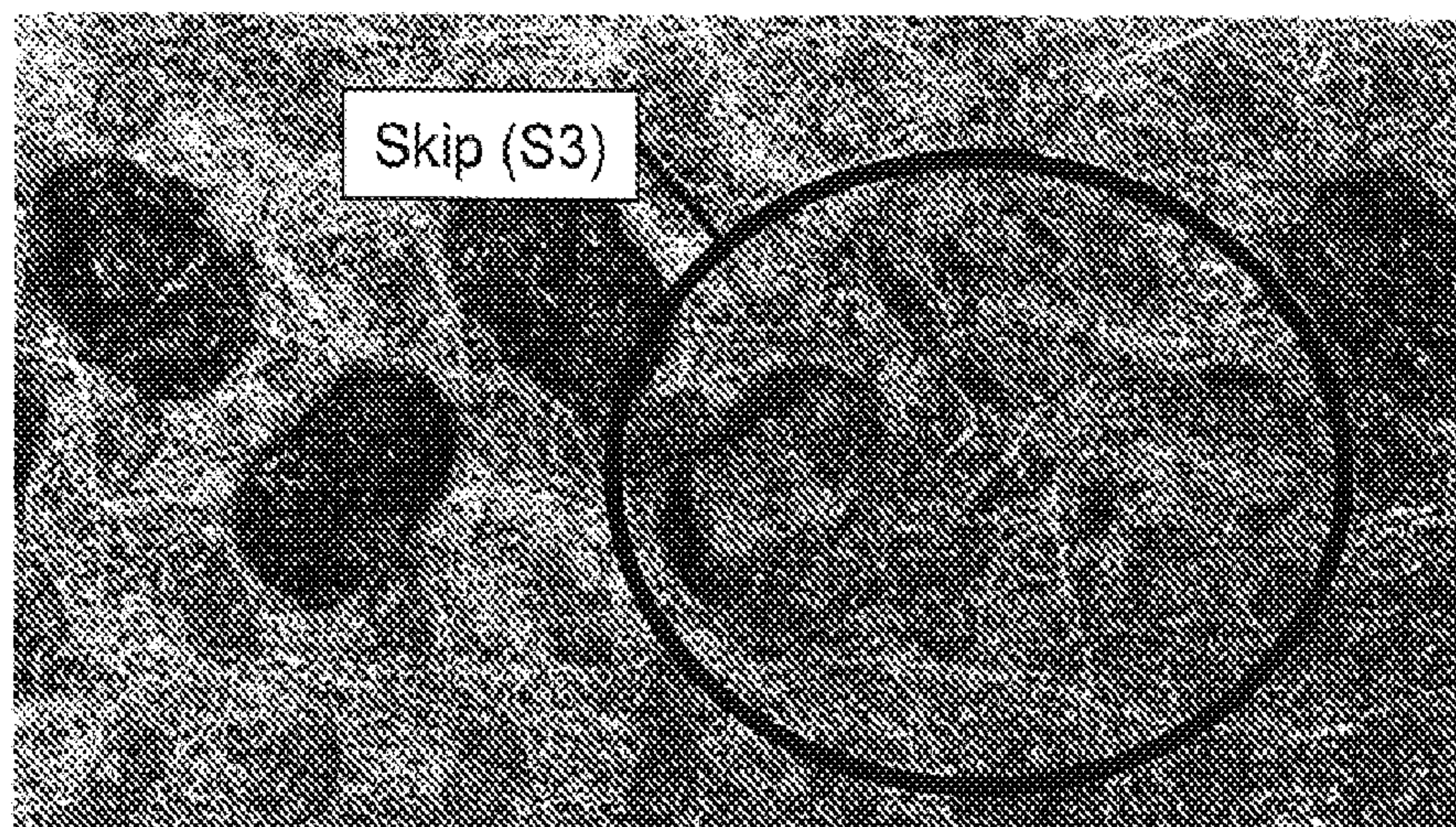


Fig. 33A

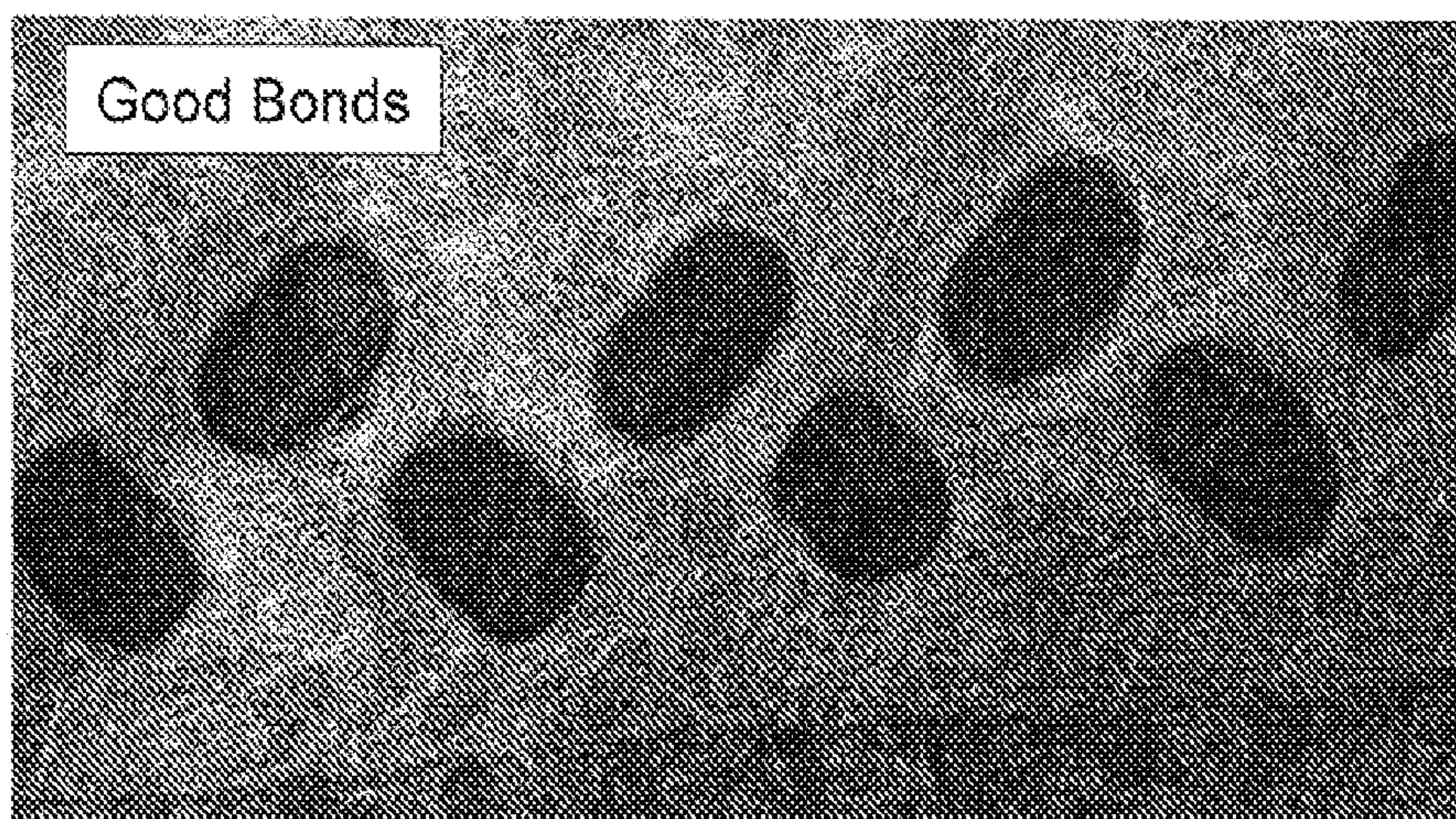


Fig. 33B

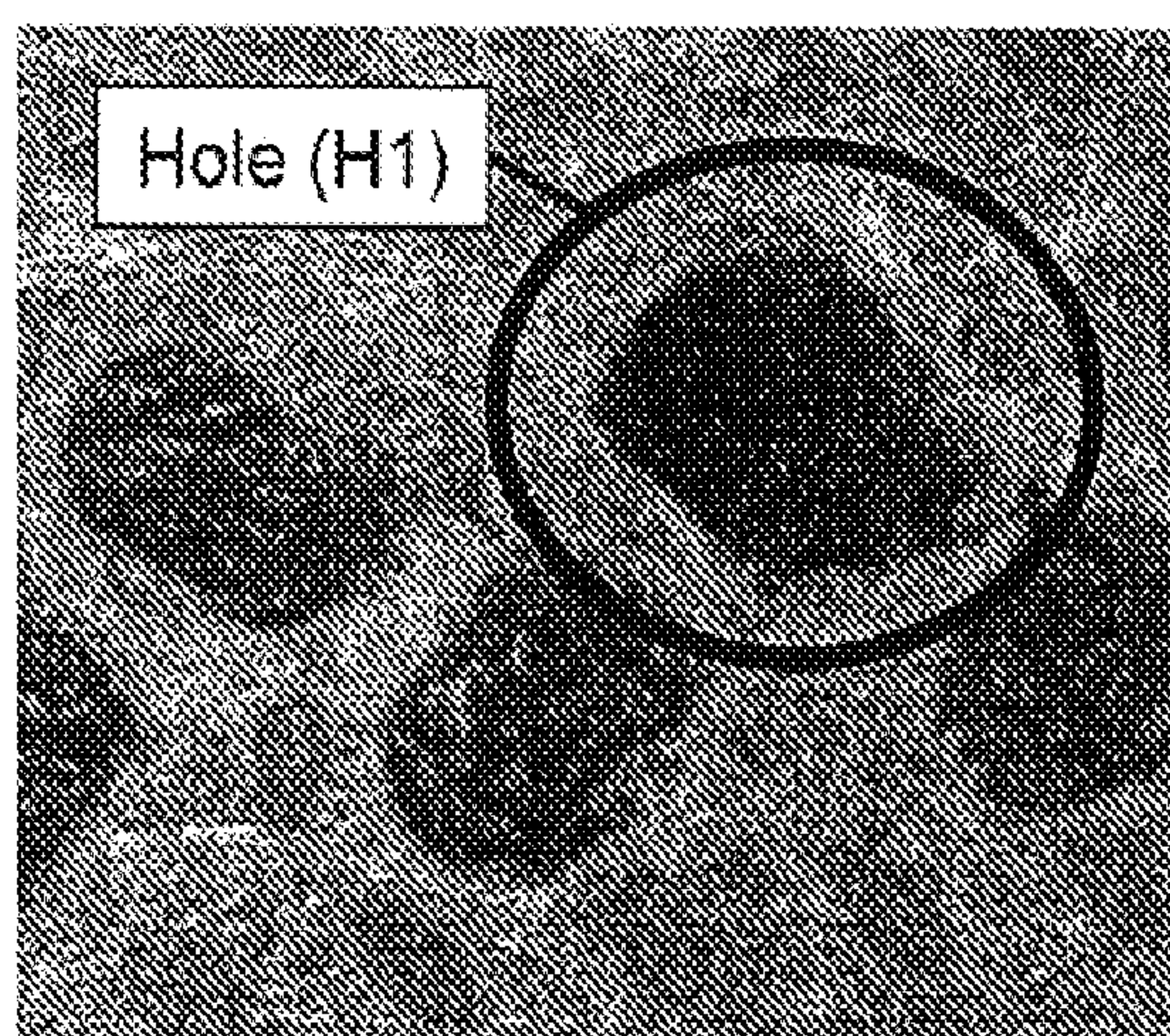


Fig. 33C

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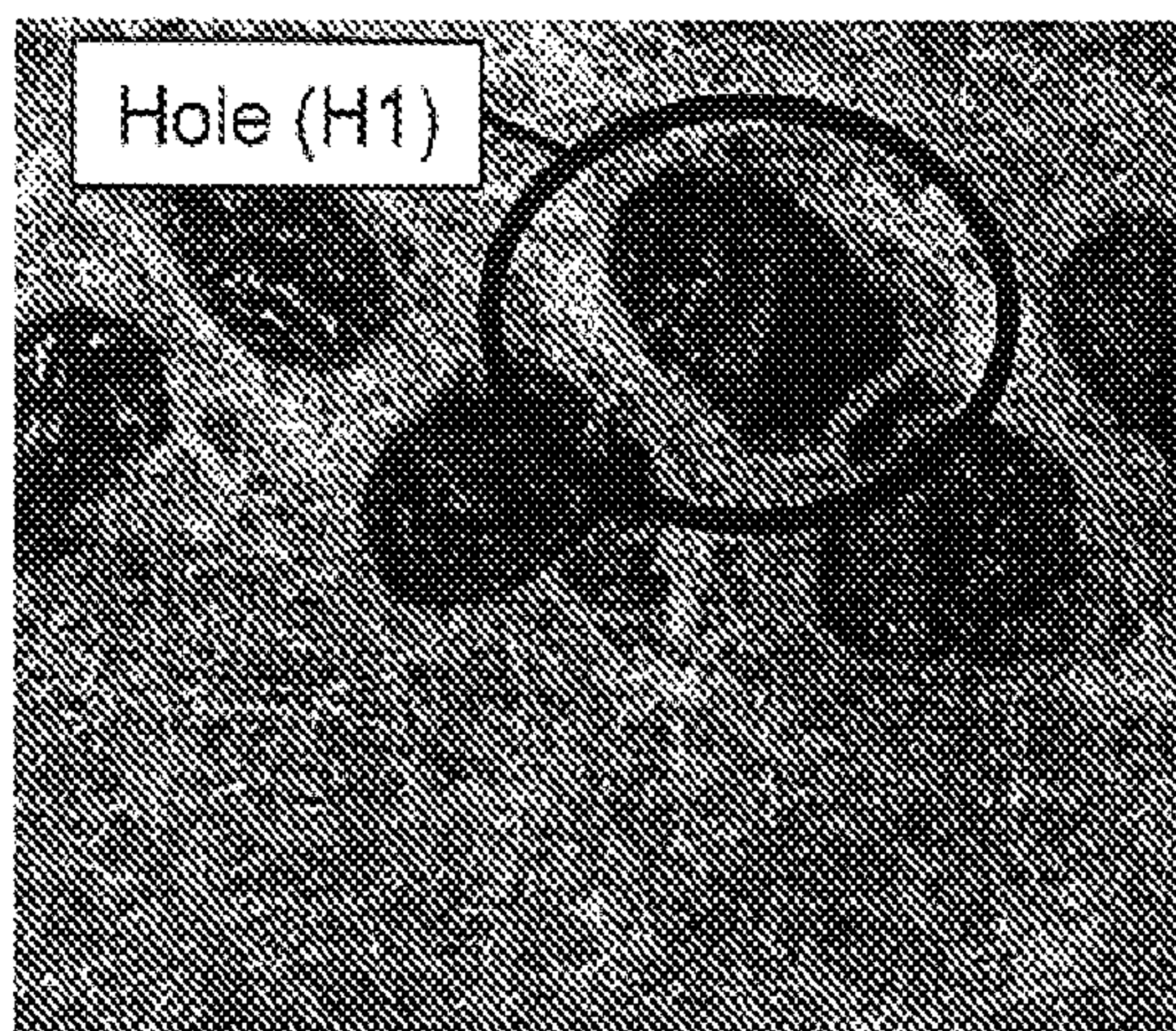


Fig. 33D

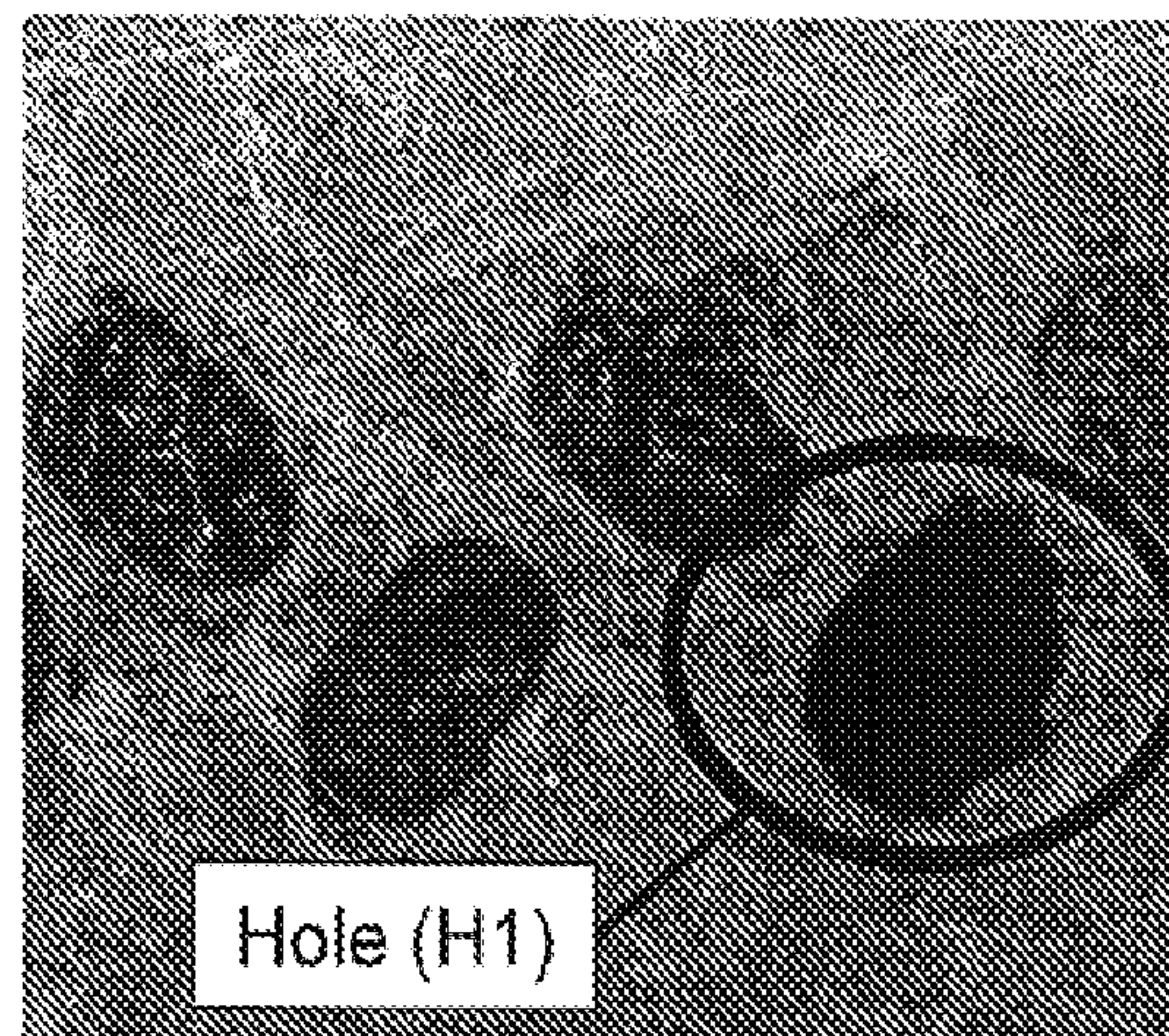


Fig. 33E

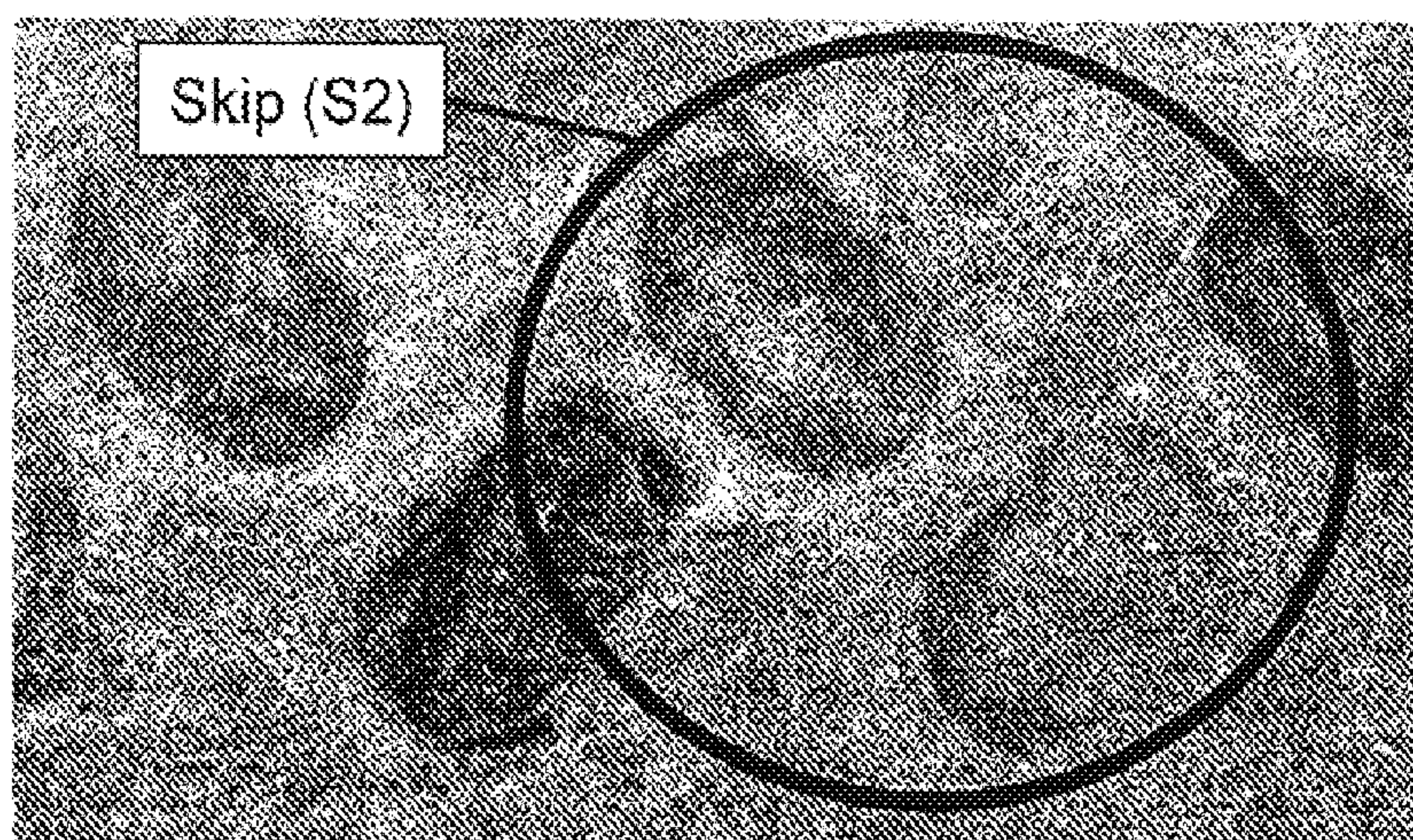


Fig. 33F

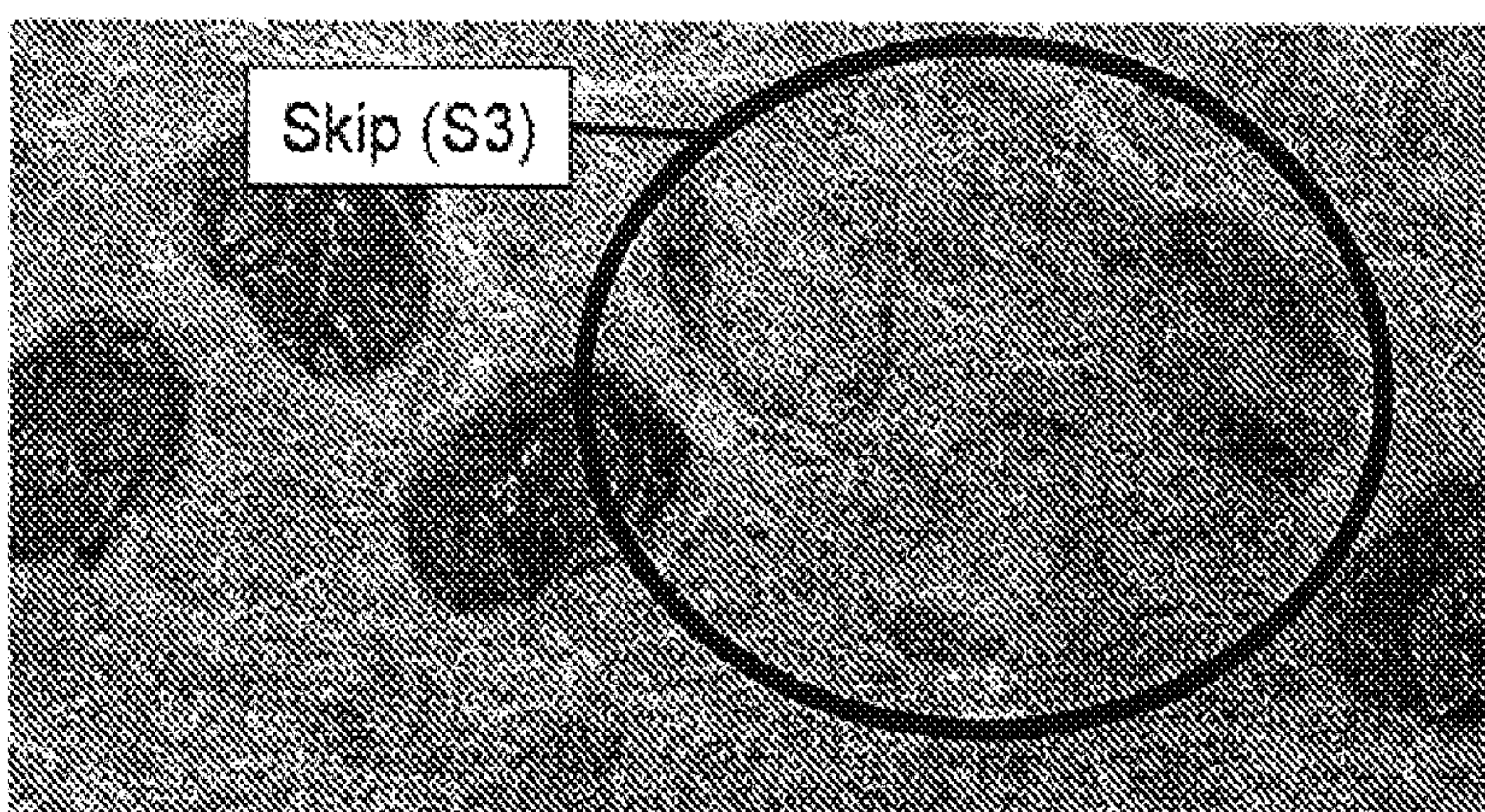


Fig. 33G

