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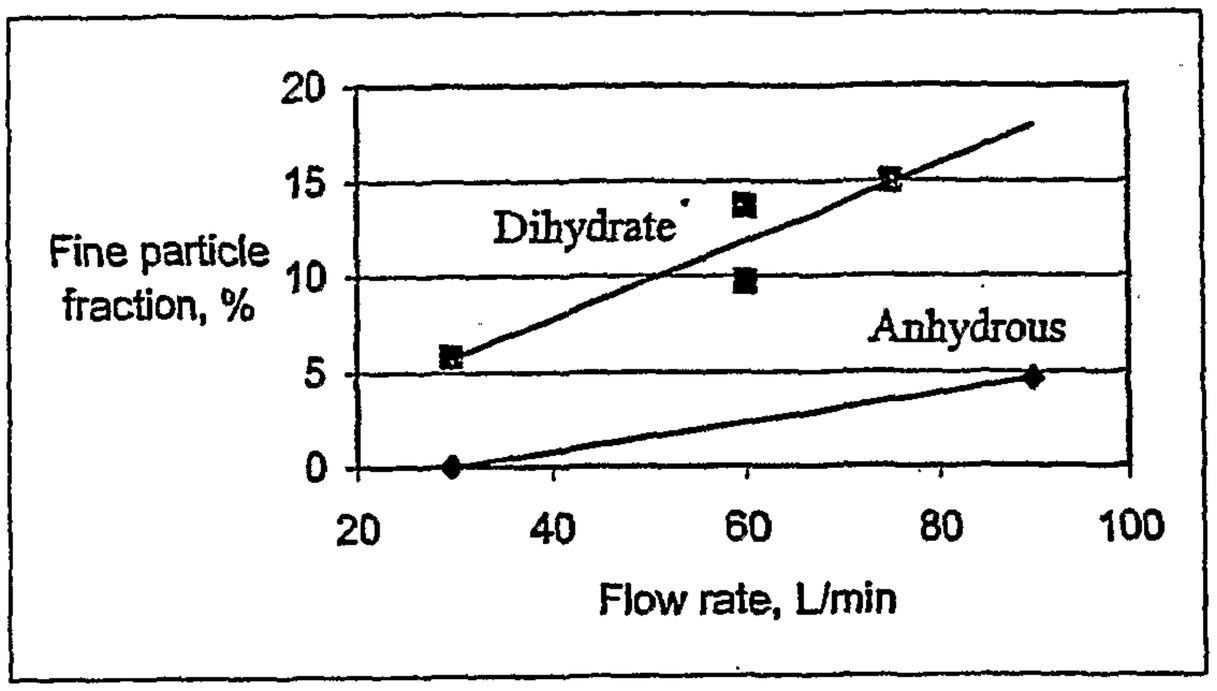
(54) Titre : COMBINAISON CONSTITUEE DE DESHYDROEPIANDROSTERONE OU DE DESHYDROEPIANDROSTERONE-SULFATE ET D'UN BRONCHODILATATEUR BETA-AGONISTE DESTINEE AU TRAITEMENT DE L'ASTHME OU DE LA BRONCHOPNEUMOPATHIE CHRONIQUE OBSTRUCTIVE

(54) Title: COMBINATION OF DEHYDROEPIANDROSTERONE OR DEHYDROEPIANDROSTERONE-SULFATE WITH A BETA-AGONIST BRONCHODILATOR FOR TREATMENT OF ASTHMA OR CHRONIC OBSTRUCTIVE PULMONARY DISEASE



(57) Abrégé/Abstract:
A pharmaceutical or veterinary composition, comprises a first active agent selected from a dehydroepiandrosterone and/or dehydroepiandrosterone-sulfate, or a salt thereof, and a second active agent comprising a β2-agonist bronchodilator for the treatment of asthma, chronic obstructive pulmonary disease, or other respiratory diseases. The composition is provided in various formulations and in the form of a kit. The products of this patent are applied to the prophylaxis and treatment of asthma, chronic obstructive pulmonary disease, or other respiratory diseases.

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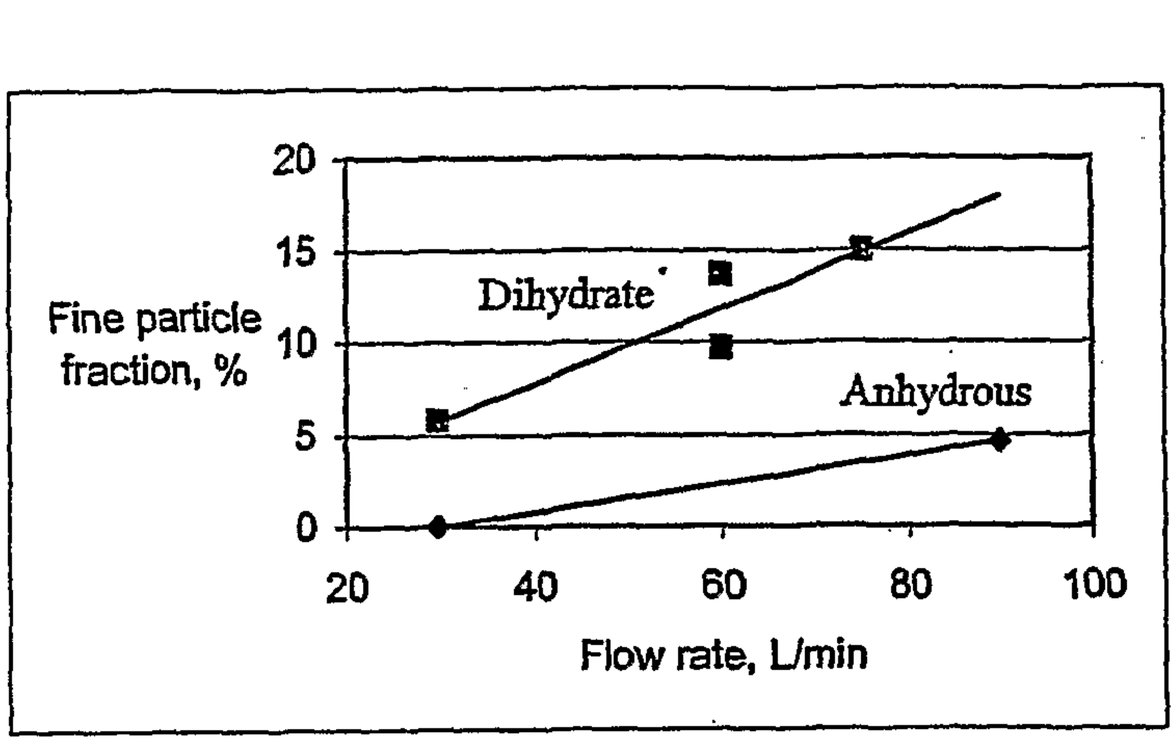

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COMBINATION OF DEHYDROEPIANDROSTERONE OR DEHYDROEPIANDROSTERONE-SULFATE WITH A BETA-AGONIST BRONCHODILATOR FOR TREATMENT OF ASTHMA OR CHRONIC OBSTRUCTIVE PULMONARY DISEASE

REFERENCE TO RELATED APPLICATIONS

This application is a continuation-in-part of U.S. Application Serial No. 10/698,078, filed on October 29, 2003, which claims priority to the U.S. Provisional Patent Application Serial No. 60/492,232, filed on July 31, 2003, which are herein incorporated by reference in their entirety.

BACKGROUND OF THE INVENTION

Field of the Invention

This invention relates to a composition comprising a non-glucocorticoid steroid including dehydroepiandrosterone (DHEA), DHEA-Sulfate, or a salt thereof, and a β2-agonist bronchodilator. These compositions are useful in the treatment of asthma, chronic obstructive pulmonary disease (COPD), or other respiratory diseases.

Description of the Background

Respiratory ailments, associated with a variety of conditions, are extremely common in the general population. In some cases they are accompanied by inflammation, which aggravates the condition of the lungs. Respiratory ailments include asthma, chronic obstructive pulmonary disease (COPD), and other upper and lower airway respiratory diseases, such as, allergic rhinitis, Acute Respiratory Distress Syndrome (ARDS), and pulmonary fibrosis.

Asthma, for example, is one of the most common diseases in industrialized countries. In the United States it accounts for about 1% of all health care costs. An alarming increase in both the prevalence and mortality of asthma over the past decade has been reported, and asthma is predicted to be the preeminent occupational lung disease in the next decade. Asthma is a condition characterized by variable, in many instances reversible obstruction of the airways. This process is associated with lung inflammation and in some cases lung allergies. Many patients have acute episodes

referred to as "asthma attacks," while others are afflicted with a chronic condition. The asthmatic process is believed to be triggered in some cases by inhalation of antigens by hypersensitive subjects. This condition is generally referred to as "extrinsic asthma." Other asthmatics have an intrinsic predisposition to the condition, which is thus referred to as "intrinsic asthma," and may be comprised of conditions of different origin, including those mediated by the adenosine receptor(s), allergic conditions mediated by an immune IgE-mediated response, and others. All asthmatics have a group of symptoms, which are characteristic of this condition: episodic bronchoconstriction, lung inflammation and decreased lung surfactant. Existing bronchodilators and anti-inflammatories are currently commercially available and are prescribed for the treatment of asthma. The most common anti-inflammatories, corticosteroids, have considerable side effects but are commonly prescribed nevertheless. Most of the drugs available for the treatment of asthma are, more importantly, barely effective in a number of patients.

COPD is characterized by airflow obstruction that is generally caused by chronic bronchitis, emphysema, or both. Commonly, the airway obstruction is incompletely reversible but 10-20% of patients do show some improvement in airway obstruction with treatment. In chronic bronchitis, airway obstruction results from chronic and excessive secretion of abnormal airway mucus, inflammation, bronchospasm, and infection. Chronic bronchitis is also characterized by chronic cough, mucus production, or both, for at least three months in at least two successive years where other causes of chronic cough have been excluded. In emphysema, a structural element (elastin) in the terminal bronchioles is destroyed leading to the collapse of the airway walls and inability to exhale "stale" air. In emphysema there is permanent destruction of the alveoli. Emphysema is characterized by abnormal permanent enlargement of the air spaces distal to the terminal bronchioles, accompanied by destruction of their walls and without obvious fibrosis. COPD can also give rise to secondary pulmonary hypertension. Secondary pulmonary hypertension itself is a disorder in which blood pressure in the pulmonary arteries is abnormally high. In severe cases, the right side of the heart must work harder than usual to pump blood against the high pressure. If this continues for a long period, the right heart enlarges and functions poorly, and fluid collects in the ankles (edema) and

belly. Eventually the left heart begins to fail. Heart failure caused by pulmonary disease is called *cor pulmonale*.

COPD characteristically affects middle aged and elderly people, and is one of the leading causes of morbidity and mortality worldwide. In the United States it affects about 14 million people and is the fourth leading cause of death, and the third leading cause for disability in the United States. Both morbidity and mortality, however, are rising. The estimated prevalence of this disease in the United States has risen by 41% since 1982, and age adjusted death rates rose by 71% between 1966 and 1985. This contrasts with the decline over the same period in age-adjusted mortality from all causes (which fell by 22%), and from cardiovascular diseases (which fell by 45%). In 1998 COPD accounted for 112,584 deaths in the United States.

COPD, however, is preventable, since it is believed that its main cause is exposure to cigarette smoke. Long-term smoking is the most frequent cause of COPD. It accounts for 80 to 90% of all cases. A smoker is 10 times more likely than a non-smoker to die of COPD. The disease is rare in lifetime non-smokers, in whom exposure to environmental tobacco smoke will explain at least some of the airways obstruction. Other proposed etiological factors include airway hyper responsiveness or hypersensitivity, ambient air pollution, and allergy. The airflow obstruction in COPD is usually progressive in people who continue to smoke. This results in early disability and shortened survival time. Smoking cessation shows the rate of decline to that of a non-smoker but the damage caused by smoking is irreversible. Other risk factors include: heredity, second-hand smoke, exposure to air pollution at work and in the environment, and a history of childhood respiratory infections. The symptoms of COPD include: chronic coughing, chest tightness, shortness of breath at rest and during exertion, an increased effort to breathe, increased mucus production, and frequent clearing of the throat.

There is very little currently available to alleviate symptoms of COPD, prevent exacerbations, preserve optimal lung function, and improve daily living activities and quality of life. Many patients will use medication chronically for the rest of their lives, with the need for increased doses and additional drugs during exacerbations. Medications that are currently prescribed for COPD patients include: fast-acting β2-agonists, anticholinergic bronchodilators, long-acting bronchodilators, antibiotics, and expectorants. Amongst the currently available treatments for COPD,

short term benefits, but not long term effects, were found on its progression, from administration of anti-cholinergic drugs, β2 adrenergic agonists, and oral steroids. Oral steroids are only recommended for acute exacerbations with long term use contributing to excess mortality and morbidity.

Short and long acting inhaled β2 adrenergic agonists achieve short-term bronchodilation and provide some symptomatic relief in COPD patients, but show no meaningful maintenance effect on the progression of the disease. Short acting β2 adrenergic agonists improve symptoms in subjects with COPD, such as increasing exercise capacity and produce some degree of bronchodilation, and even an increase in lung function in some severe cases. The maximum effectiveness of the newer long acting inhaled, β2 adrenergic agonists was found to be comparable to that of short acting β2 adrenergic agonists. Salmeterol was found to improve symptoms and quality of life, although only producing modest or no change in lung function. The use of β2-agonists can produce cardiovascular effects, such as altered pulse rate, blood pressure and electrocardiogram results. In rare cases, the use of β2-agonists can produce hypersensitivity reactions, such as urticaria, angioedema, rash and oropharyngeal edema. In these cases, the use of the β2-agonist should be discontinued. Continuous treatment of asthmatic and COPD patients with the bronchodilators ipratropium bromide or fenoterol was not superior to treatment on an as-needed basis, therefore indicating that they are not suitable for maintenance treatment. The most common immediate adverse effect of β2 adrenergic agonists, on the other hand, is tremors, which at high doses may cause a fall in plasma potassium, dysrhythmias, and reduced arterial oxygen tension. The combination of a β2 adrenergic agonist with an anti-cholinergic drug provides little additional bronchodilation compared with either drug alone. The addition of ipratropium to a standard dose of inhaled β2 adrenergic agonists for about 90 days, however, produces some improvement in stable COPD patients over either drug alone. Overall, the occurrence of adverse effects with β2 adrenergic agonists, such as tremor and dysrhythmias, is more frequent than with anti-cholinergics. Thus, neither anti-cholinergic drugs nor β2 adrenergic agonists have an effect on all people with COPD; nor do the two agents combined.

Anti-cholinergic drugs achieve short-term bronchodilation and produce some symptom relief in people with COPD, but no improved long-term prognosis. Most

COPD patients have at least some measure of airways obstruction that is somewhat alleviated by ipratropium bromide. "The Lung Health Study" found spirometric signs of early COPD in men and women smokers and followed them for five years. Three treatments were compared over a five year period and results show that ipratropium bromide had no significant effect on the decline in the functional effective volume of the patient's lungs whereas smoking cessation produced a slowing of the decline in the functional effective volume of the lungs. Ipratropium bromide, however, produced adverse effects, such as cardiac symptoms, hypertension, skin rashes, and urinary retention.

Theophyllines produce modest bronchodilation in COPD patients whereas they have frequent adverse effects, and a small therapeutic range. Serum concentrations of 15-20 mg/l are required for optimal effects and serum levels must be carefully monitored. Adverse effects include nausea, diarrhea, headache, irritability, seizures, and cardiac arrhythmias, occurring at highly variable blood concentrations and, in many people, even within the therapeutic range. The theophyllines' doses must be adjusted individually according to smoking habits, infection, and other treatments, which is cumbersome. Although theophyllines have been claimed to have an anti-inflammatory effect in asthma, especially at lower doses, none has been reported in COPD. The adverse effects of theophyllines and the need for frequent monitoring limit their usefulness.

Oral corticosteroids have been shown to improve the short term outcome in acute exacerbations of COPD but long term administration of oral steroid has been associated with serious side effects including osteoporosis and inducing overt diabetes. Inhaled corticosteroids have been found to have no real short-term effect on airway hyper-responsiveness to histamine. In two studies of 3 year treatment with inhaled fluticasone, moderate and severe exacerbations were significantly reduced as well as a modest improvement in the quality of life without affecting pulmonary function. COPD patients with more reversible disease seem to benefit more from treatment with inhaled fluticasone.

Mucolytics have a modest beneficial effect on the frequency and duration of exacerbations but an adverse effect on lung function. Neither N-acetylcysteine nor other mucolytics, however, have a significant effect in people with severe COPD (functional effective volume<50%) in spite of evidencing greater reductions in

frequency of exacerbation. N-acetylcysteine produced gastrointestinal side effects. Long-term oxygen therapy administered to hypoxaemic COPD and congestive cardiac failure patients, had little effect on their rates of death for the first 500 days or so, but survival rates in men increased afterwards and remained constant over the next five years. In women, however, oxygen decreased the rates of death throughout the study. Continuous oxygen treatment of hypoxemic COPD patients for 19.3 years decreased overall risk of death. To date, however, only life style changes, smoking cessation and long term treatment with oxygen (in hypoxaemics), have been found to alter the long-term course of COPD.

Antibiotics are also often given at the first sign of a respiratory infection to prevent further damage and infection in diseased lungs. Expectorants help loosen and expel mucus secretions from the airways, and may help make breathing easier. In addition, other medications may be prescribed to manage conditions associated with COPD. These may include: diuretics (which are given as therapy to avoid excess water retention associated with right-heart failure), digitalis (which strengthens the force of the heartbeat), and cough suppressants. This latter list of medications help alleviate symptoms associated with COPD but do not treat COPD. Thus, there is very little currently available to alleviate symptoms of COPD, prevent exacerbations, preserve optimal lung function, and improve daily living activities and quality of life.

Acute Respiratory Distress Syndrome (ARDS), or stiff lung, shock lung, pump lung and congestive atelectasis, is believed to be caused by fluid accumulation within the lung which, in turn, causes the lung to stiffen. The condition is triggered within 48 hours by a variety of processes that injure the lungs such as trauma, head injury, shock, sepsis, multiple blood transfusions, medications, pulmonary embolism, severe pneumonia, smoke inhalation, radiation, high altitude, near drowning, and others. In general, ARDS occurs as a medical emergency and may be caused by other conditions that directly or indirectly cause the blood vessels to "leak" fluid into the lungs. In ARDS, the ability of the lungs to expand is severely decreased and produces extensive damage to the air sacs and lining or endothelium of the lung. ARDS' most common symptoms are labored, rapid breathing, nasal flaring, cyanosis blue skin, lips and nails caused by lack of oxygen to the tissues, anxiety, and temporarily absent breathing. A preliminary diagnosis of ARDS may be confirmed with chest X-rays and the measurement of arterial blood gas. In some cases ARDS appears to be associated

with other diseases, such as acute myelogenous leukemia, with acute tumor lysis syndrome (ATLS) developed after treatment with, e.g. cytosine arabinoside. In general, however, ARDS appears to be associated with traumatic injury, severe blood infections such as sepsis, or other systemic illness, high dose radiation therapy and chemotherapy, and inflammatory responses which lead to multiple organ failure, and in many cases death. In premature babies ("preemies"), neither the lung tissue nor the surfactant is fully developed. When Respiratory Distress Syndrome (RDS) occurs in preemies, it is an extremely serious problem. Preterm infants exhibiting RDS are currently treated by ventilation and administration of oxygen and surfactant preparations. When preemies survive RDS, they frequently develop bronchopulmonary dysplasia (BPD), also called chronic lung disease of early infancy, which is often fatal.

Allergic rhinitis afflicts one in five Americans, accounting for an estimated $4 to 10 billion in health care costs each year, and occurs at all ages. Because many people mislabel their symptoms as persistent colds or sinus problems, allergic rhinitis is probably underdiagnosed. Typically, IgE combines with allergens in the nose to produce chemical mediators, induction of cellular processes, and neurogenic stimulation, causing an underlying inflammation. Symptoms include ocular and nasal congestion, discharge, sneezing, and itching. Over time, allergic rhinitis sufferers often develop sinusitis, otitis media with effusion, and nasal polyposis. Approximately 60% of patients with allergic rhinitis also have asthma and flares of allergic rhinitis aggravate asthma. Degranulation of mast cells results in the release of preformed mediators that interact with various cells, blood vessels, and mucous glands to produce the typical rhinitis symptoms. Most early- and late-phase reactions occur in the nose after allergen exposure. The late-phase reaction is seen in chronic allergic rhinitis, with hypersecretion and congestion as the most prominent symptoms. Repeated exposure causes a hypersensitivity reaction to one or many allergens. Sufferers may also become hyperreactive to nonspecific triggers such as cold air or strong odors. Nonallergic rhinitis may be induced by infections, such as viruses, or associated with nasal polyps, as occurs in patients with aspirin idiosyncrasy.

Medical conditions such as pregnancy or hypothyroidism and exposure to occupational factors or medications may cause rhinitis. The so-called NARES syndrome (Nonallergic Rhinitis with Eosinophilia Syndrome) is a non-allergic type of

rhinitis associated with eosinophils in the nasal secretions, which typically occurs in middle-age and is accompanied by some loss of sense of smell. Treatment of allergic and non-allergic rhinitis is unsatisfactory. Self-administered saline improves nasal stuffiness, sneezing, and congestion and usually causes no side effects and it is, thus, the first treatment tried in pregnant patients. Saline sprays are generally used to relieve mucosal irritation or dryness associated with various nasal conditions, minimize mucosal atrophy, and dislodge encrusted or thickened mucus. If used immediately before intranasal corticosteroid dosing, saline sprays may help prevent drug-induced local irritation. Anti-histamines such as terfenadine and astemizole are also employed to treat allergic rhinitis; however, use of antihistamines have been associated with a ventricular arrhythmia known as Torsades de Points, usually in interaction with other medications such as ketoconazole and erythromycin, or secondary to an underlying cardiac problem. Loratadine, another non-sedating anti-histamine, and cetirizine have not been associated with an adverse impact on the QT interval, or with serious adverse cardiovascular events. Cetirizine, however, produces extreme drowsiness and has not been widely prescribed. Non-sedating anti-histamines, e.g. Claritin, may produce some relieving of sneezing, runny nose, and nasal, ocular and palatal itching, but have not been tested for asthma or other more specific conditions. Terfenadine, loratadine and astemizole, on the other hand, exhibit extremely modest bronchodilating effects, reduction of bronchial hyper-reactivity to histamine, and protection against exercise- and antigen-induced bronchospasm. Some of these benefits, however, require higher-than-currently-recommended doses. The sedating-type anti-histamines help induce night sleep, but they cause sleepiness and compromise performance if taken during the day. When employed, anti-histamines are typically combined with a decongestant to help relieve nasal congestion. Sympathomimetic medications are used as vasoconstrictors and decongestants. The three commonly prescribed systemic decongestants, pseudoephedrine, phenylpropanolamine and phenylephrine cause hypertension, palpitations, tachycardia, restlessness, insomnia and headache. The interaction of phenylpropanolamine with caffeine, in doses of two to three cups of coffee, may significantly raise blood pressure. In addition, medications such as pseudoephedrine may cause hyperactivity in children. Topical decongestants, nevertheless, are only indicated for a limited period of time, as they are associated with a rebound nasal dilatation with overuse.

Anti-cholinergic agents are given to patients with significant rhinorrhea or for specific conditions such as "gustatory rhinitis", usually caused by ingestion of spicy foods, and may have some beneficial effects on the common cold. Cromolyn, for example, if used prophylactically as a nasal spray, reduces sneezing, rhinorrhea, and nasal pruritus, and blocks both early- and late-phase hypersensitivity responses, but produces sneezing, transient headache, and even nasal burning. Topical corticosteroids such as Vancenase are effective in the treatment of rhinitis, especially for symptoms of itching, sneezing, and runny nose but are less effective against nasal stuffiness. Depending on the preparation, however, corticosteroid nose sprays may cause irritation, stinging, burning, or sneezing, as well. Local bleeding and septal perforation can also occur sometimes, especially if the aerosol is not aimed properly. Topical steroids generally are more effective than cromolyn sodium in the treatment of allergic rhinitis. Immunotherapy, while expensive and inconvenient, often provides benefits, especially for inpatients who experience side effects from other medications. So-called blocking antibodies, and agents that alter cellular histamine release, eventually result in decreased IgE, along with many other favorable physiologic changes. This effect is useful in IgE-mediated diseases, e.g., hypersensitivity in atopic patients with recurrent middle ear infections.

Pulmonary fibrosis, interstitial lung disease (ILD), or interstitial pulmonary fibrosis, include more than 130 chronic lung disorders that affect the lung by damaging lung tissue, and produce inflammation in the walls of the air sacs in the lung, scarring or fibrosis in the interstitium (or tissue between the air sacs), and stiffening of the lung. Breathlessness during exercise may be one of the first symptoms of these diseases, and a dry cough may be present. Neither the symptoms nor X-rays are often sufficient to differentiate various types of pulmonary fibrosis. Some pulmonary fibrosis patients have known causes and some have unknown or idiopathic causes. The course of this disease is generally unpredictable and the disease is inevitably fatal. Its progression includes thickening and stiffening of the lung tissue, inflammation and difficult breathing. Most people may need oxygen therapy and the only treatment is lung transplantation.

Lung cancer is the most common cancer in the world. During 2003, there will be about 171,900 new cases of lung cancer (91,800 among men and 80,100 among women) in the US alone and approximately 375,000 cases in Europe. Lung cancer is

the leading cause of cancer death among both men and women. There will be an estimated 157,200 deaths from lung cancer (88,400 among men and 68,800 among women) in 2003, accounting for 28% of all cancer deaths in the US alone. More people die of lung cancer than of colon, breast, and prostate cancers combined (American Cancer Society Web site, 2003, Detailed Guide: Lung Cancer: What are the Key Statistics?). Tobacco smoking is well established as the main cause of lung cancer and about 90% of cases are thought to be tobacco related. There is a clear dose-response relation between lung-cancer risk and the number of cigarettes smoked per day, degree of inhalation, and age at initiation of smoking. Lifelong smokers have a lung-cancer risk 20-30 times greater than a non-smoker. However, risk of lung cancer decreases with time since smoking cessation. The relative risk of male ex-smokers decreases strongly with time since end of exposure, but does not reach the risk of non-smokers, and does not decrease as much as for female ex-smokers (Tyczynski et al., *Lancet Oncol.* 4(1):45-55 (2003).

Frequently, COPD and lung cancer are co-morbid diseases and the degree of underlying COPD may dictate whether a particular patient is a surgical candidate. For NSCLC (non small cell lung cancer), only surgery (with or without radiation therapy or adjuvant chemotherapy) is curative.

- The 1-year survival rate (the number of people who live at least 1 year after their cancer is diagnosed) for lung cancer was 42% in 1998, largely due to improvements in surgical techniques.
- The 5-year survival rate for all stages of non-small cell lung cancer combined is only 15%. For small cell lung cancer the 5-year relative survival rate is about 6%.
- For people whose NSCLC is found and treated early with surgery, before it has spread to lymph nodes or other organs, the average 5-year survival rate is about 50%. However, only 15% of people with lung cancer are diagnosed at this early, localized stage.

Clearly, there is much room for improvement in chemoprophylaxis of lung cancer as well as treatment of lung cancer.

Dehydroepiandrosterone (DHEA) (3β-hydroxyandrost-5-en-17-one) is a naturally occurring steroid secreted by the adrenal cortex with apparent chemoprotective properties. Epidemiological studies have shown that low

endogenous levels of DHEA correlate with increased risk of developing some forms of cancer, such as pre-menopausal breast cancer in women and bladder cancer in both sexes. The ability of DHEA and DHEA analogues, such as DHEA-S (DHEA-sulfate), to inhibit carcinogenesis is believed to result from their uncompetitive inhibition of the activity of the enzyme glucose-6-phosphate dehydrogenase (G6PDH). G6PDH is the rate limiting enzyme of the hexose monophosphate pathway, a major source of intracellular ribose-5-phosphate and NADPH. Ribose-5-phosphate is a necessary substrate for the synthesis of both ribo- and deoxyribonucleotides. NADPH is a cofactor also involved in nucleic acid biosynthesis and the synthesis of hydroxmethylglutaryl Coenzyme A reductase (HMG CoA reductase). HMG CoA reductase is an unusual enzyme that requires two moles of NADPH for each mole of product, mevalonate, produced. Thus, it appears that HMG CoA reductase would be ultrasensitive to DHEA-mediated NADPH depletion, and that DHEA-treated cells would rapidly show the depletion of intracellular pools of mevalonate. Mevalonate is required for DNA synthesis, and DHEA arrests human cells in the G1 phase of the cell cycle in a manner closely resembling that of the direct HMG CoA. Because G6PDH is required to produces mevalonic acid used in cellular processes such as protein isoprenylation and the synthesis of dolichol, a precursor for glycoprotein biosynthesis, DHEA inhibits carcinogenesis by depleting mevalonic acid and thereby inhibiting protein isoprenylation and glycoprotein synthesis. Mevalonate is the central precursor for the synthesis of cholesterol, as well as for the synthesis of a variety of non-sterol compounds involved in post-translational modification of proteins such as farnesyl pyrophosphate and geranyl pyrophosphate; and for dolichol, which is required for the synthesis of glycoproteins involved in cell-to-cell communication and cell structure.

It has long been known that patients receiving steroid hormones of adrenocortical origin at pharmacologically appropriate doses show increased incidence of infectious disease. U.S. Patent No. 5,527,789 discloses a method of combating cancer by administering to a patient DHEA and ubiquinone, where the cancer is one that is sensitive to DHEA.

DHEA is a 17-ketosteroid which is quantitatively one of the major adrenocortical steroid hormones found in mammals. Although DHEA appears to serve as an intermediary in gonadal steroid synthesis, the primary physiological function of DHEA has not been fully understood. It has been known, however, that

levels of this hormone begin to decline in the second decade of life (reaching 5% of the original level in the elderly.) Clinically, DHEA has been used systemically and/or topically for treating patients suffering from psoriasis, gout, hyperlipemia, and it has been administered to post-coronary patients. In mammals, DHEA has been shown to have weight optimizing and anti-carcinogenic effects, and it has been used clinically in Europe in conjunction with estrogen as an agent to reverse menopausal symptoms and also has been used in the treatment of manic depression, schizophrenia, and Alzheimer's disease. DHEA has been used clinically at 40 mg/kg/day in the treatment of advanced cancer and multiple sclerosis. Mild androgenic effects, hirsutism, and increased libido were the side effects observed. These side effects can be overcome by monitoring the dose and/or by using analogues. The subcutaneous or oral administration of DHEA to improve the host's response to infections is known, as is the use of a patch to deliver DHEA. DHEA is also known as a precursor in a metabolic pathway which ultimately leads to more powerful agents that increase immune response in mammals. That is, DHEA acts as a prodrug: it acts as an immuno-modulator when converted to androstenediol or androst-5-ene-3β,17β-diol (βAED), or androstenetriol or androst-5-ene-3β,7β,17β-triol (βAET). However, in vitro DHEA has certain lymphotoxic and suppressive effects on cell proliferation prior to its conversion to βAED and/or βAET. It is, therefore, believed that the superior immunity enhancing properties obtained by administration of DHEA result from its conversion to more active metabolites.

Adenosine is a purine involved in intermediary metabolism, and may constitute an important mediator in the lung for various diseases, including bronchial asthma, COPD, CF, RDS, rhinitis, pulmonary fibrosis, and others. The potential role of its receptor was suggested by the finding that asthmatics respond to aerosolized adenosine with marked bronchoconstriction whereas normal individuals do not. An asthmatic rabbit animal model, the dust mite allergic rabbit model for human asthma, responded in a similar fashion to aerosolized adenosine with marked bronchoconstriction whereas non-asthmatic rabbits showed no response. More recent work with this animal model suggested that adenosine-induced bronchoconstriction and bronchial hyperresponsiveness in asthma may be mediated primarily through the stimulation of adenosine receptors. Adenosine has also been shown to cause adverse effects, including death, when administered therapeutically for other diseases and

conditions in subjects with previously undiagnosed hyper-reactive airways.

Adenosine plays a unique role in the body as a regulator of cellular metabolism. It can raise the cellular level of AMP, ADP and ATP that are the energy intermediates of the cell. Adenosine can stimulate or down regulate the activity of adenylate cyclase and hence regulate cAMP levels. cAMP, in turn, plays a role in neurotransmitter release, cellular division and hormone release. Adenosine's major role appears to be to act as a protective injury autocoid. In conditions in which ischemia, low oxygen tension or trauma occurs adenosine appears to play a role. Defects in synthesis, release, action and/or degradation of adenosine have been postulated to contribute to the over activity of the brain excitatory amino acid neurotransmitters, and hence various pathological states. Adenosine has also been implicated as a primary determinant underlying the symptoms of bronchial asthma and other respiratory diseases, the induction of bronchoconstriction and the contraction of airway smooth muscle. Moreover, adenosine causes bronchoconstriction in asthmatics but not in non-asthmatics. Other data suggest the possibility that adenosine receptors may also be involved in allergic and inflammatory responses by reducing the hyperactivity of the central dopaminergic system. It has been postulated that the modulation of signal transduction at the surface of inflammatory cells influences acute inflammation. Adenosine is said to inhibit the production of super-oxide by stimulated neutrophils. Recent evidence suggests that adenosine may also play a protective role in stroke, CNS trauma, epilepsy, ischemic heart disease, coronary by-pass, radiation exposure and inflammation. Overall, adenosine appears to regulate cellular metabolism through ATP, to act as a carrier for methionine, to decrease cellular oxygen demand and to protect cells from ischemic injury. Adenosine is a tissue hormone or inter-cellular messenger that is released when cells are subject to ischemia, hypoxia, cellular stress, and increased workload, and or when the demand for ATP exceeds its supply. Adenosine is a purine and its formation is directly linked to ATP catabolism. It appears to modulate an array of physiological processes including vascular tone, hormone action, neural function, platelet aggregation and lymphocyte differentiation. It also may play a role in DNA formation, ATP biosynthesis and general intermediary metabolism. It is suggested that it regulates the formation of cAMP in the brain and in a variety of peripheral tissues. Adenosine regulates cAMP formation through two receptors A_1 and A_2. Via A_1 receptors, adenosine reduces adenylate cyclase activity,

while it stimulates adenylate cyclase at A_2 receptors. The adenosine A_1 receptors are more sensitive to adenosine than the A_2 receptors. The CNS effects of adenosine are generally believed to be A_1-receptor mediated, where as the peripheral effects such as hypotension, bradycardia, are said to be A_2 receptor mediated.

A handful of medicaments have been used for the treatment of respiratory diseases and conditions, although in general they all have limitations. Amongst them are glucocorticoid steroids, leukotriene inhibitors, anti-cholinergic agents, anti-histamines, oxygen therapy, theophyllines, and mucolytics. Glucocorticoid steroids are the ones with the most widespread use in spite of their well documented side effects. Most of the available drugs are nevertheless effective in a small number of cases, and not at all when it comes to the treatment of asthma. No treatments are currently available for many of the other respiratory diseases. Theophylline, an important drug in the treatment of asthma, is a known adenosine receptor antagonist which was reported to eliminate adenosine-mediated bronchoconstriction in asthmatic rabbits. A selective adenosine A1 receptor antagonist, 8-cyclopentyl-1, 3-dipropylxanthine (DPCPX) was also reported to inhibit adenosine-mediated bronchoconstriction and bronchial hyperresponsiveness in allergic rabbits. The therapeutic and preventative applications of currently available adenosine A1 receptor-specific antagonists are, nevertheless, limited by their toxicity. Theophylline, for example, has been widely used in the treatment of asthma, but is associated with frequent, significant toxicity (gastrointestinal, cardiovascular, neurological and biological disturbances) resulting from its narrow therapeutic dose range. DPCPX is far too toxic to be useful clinically. The fact that, despite decades of extensive research, no specific adenosine receptor antagonist is available for clinical use attests to the general toxicity of these agents.

Currently, the β2-agonist bronchodilator salmeterol is available commercially for treating wheezing, shortness of breath, and troubled breathing caused by asthma and COPD, which includes chronic bronchitis, emphysema, and other lung diseases. Salmeterol also is used to prevent breathing difficulties (bronchospasm) during exercise. Salmeterol comes as an aerosol to use by oral inhalation. It is marketed as Serevent™ Accuhaler™, Serevent™ Diskhaler™ and Serevent™ Inhaler in the United Kingdom (Glaxo Wellcome, UK).

Currently, the β2-agonist bronchodilator formoterol is available commercially for treating bronchoconstriction inpatients with COPD, including chronic bronchitis and emphysema. It is marketed in capsule form containing a dry powder formulation (Foradil®), containing 12 μg of formoterol fumarate and 25 mg of lactose, to be used for oral inhalation only with the Aerolizer® Inhaler (Schering Corp., Kenilworth, NJ).

U.S. Patent No. 5,660,835 (and corresponding PCT publication WO 96/25935) discloses a novel method of treating asthma or adenosine depletion in a subject by administering to the subject a dehydroepiandrosterone (DHEA) or DHEA-related compound. The patent also discloses a novel pharmaceutical composition in regards to an inhalable or respirable formulation comprising DHEA or DHEA-related compounds that is in a respirable particle size.

U.S. Patent No. 5,527,789 discloses a method of combating cancer in a subject by administering to the subject a DHEA or DHEA-related compound, and ubiquinone to combat heart failure induced by the DHEA or DHEA-related compound.

U.S. Patent No. 6,087,351 discloses an in vivo method of reducing or depleting adenosine in a subject's tissue by administering to the subject a DHEA or DHEA-related compound.

U.S. Patent Application Ser. No. 10/454,061, filed June 3, 2003, discloses a method for treating COPD in a subject by administering to the subject a DHEA or DHEA-related compound.

U.S. Patent Application Ser. No. 10/462,901, filed June 17, 2003, discloses a stable dry powder formulation of DHEA in a nebulizable form sealed in a container.

U.S. Patent Application Ser. No. 10/462,927, filed June 17, 2003, discloses a stable dry powder formulation of dihydrate crystal form of DHEA-S suitable for treating asthma and COPD.

The above patents and patent applications are herein incorporated by reference in their entirety.

There exists a well defined need for novel and effective therapies for treating respiratory, lung and cancer ailments that cannot presently be treated, or at least for which no therapies are available that are effective and devoid of significant detrimental side effects. This is the case of ailments afflicting the respiratory tract, and more particularly the lung and the lung airways, including respiratory difficulties,

asthma, bronchoconstriction, lung inflammation and allergies, depletion or hyposecretion of surfactant, etc. Moreover, there is a definite need for treatments that have prophylctic and therapeutic applications, and require low amounts of active agents, which makes them both less costly and less prone to detrimental side effects.

Further, there is a need to better ensure patient compliance in the taking of medication, and a need to facilitate the taking of the plurality of compounds necessary for prevention or treatment of asthma, COPD, or other respiratory diseases.

SUMMARY OF THE INVENTION

The present invention provides for a composition comprising at least two active agents. A first active agent comprises a non-glucocorticoid steroid, such as an epiandrosterone (EA) or a salt thereof. A second active agent comprises a β2-agonist bronchodilator. The composition comprises a combination of the first active agent and the second active agent. The amount of the first active agent and the amount of the second active agent in the composition is of an amount sufficient to effectively prophylactically or therapeutically treat a subject in danger of suffering or suffering from asthma, COPD, or other respiratory diseases when the composition is administered to the subject. The composition can further comprise other bioactive agents and formulation ingredients. The composition is a pharmaceutical or veterinary composition suitable for administration to a subject or patient, such as a human or a non-human animal (such as a non-human mammal).

The composition is useful for treating asthma, COPD, or other respiratory diseases for which inflammation and its sequelae plays a role including conditions associated with bronchoconstriction, surfactant depletion and/or allergies.

The present invention also provides for methods for treating asthma, COPD, lung cancer, or other respiratory diseases comprising administering the composition to a subject in need of such treatment.

The present invention also provides for a use of the first active agent and the second active agent in the manufacture of a medicament for the prophylactic or therapeutic treatment of asthma, COPD, or other respiratory diseases described above.

The present invention also provides for a kit comprising the composition and a delivery device. The delivery device is capable of delivering the composition to the subject. Preferably, the delivery device comprises an inhaler provided with an aerosol or spray generating means that delivers particles. Preferably, the delivery is

to the airway of the subject. More preferably, the delivery is to the lung or lungs of the subject. Preferably, the delivery is directly to the desired location.

The main advantage of using the compositions is the compliance by the patients in need of such prophylaxis or treatment. Respiratory diseases such as asthma or COPD are multifactorial with different manifestations of signs and symptoms for individual patients. As such, most patients are treated with multiple medications to alleviate different aspects of the disease. A fixed combination of the first active agent, such as DHEA or DHEA-S, and the second active agent, such as salmeterol or formoterol, permits more convenient yet targeted therapy for a defined patient subpopulation. Patient compliances should be improved by simplifying therapy and by focusing on each patient's unique disease attributes so that their specific symptoms are addressed in the most expeditious fashion. Further, there is the added advantage of convenience or savings in time in the administering of both the first and second active agents in one administration. This is especially true when the composition is administered to a region of the body of the subject that has the potential of discomfort, such as the composition administered to the airways of the subject. This is also especially true when the administration of the compositions to the subject is invasive.

In addition, the first active agent, such as DHEA or DHEA-S, is an anti-inflammatory agent that is most effective when it is delivered or deposited in the distal peripheral airways rather than the conducting airways, in the alveolar membranes and fine airways. Asthma and some COPD patients have conducting airways that are constricted, which limit the delivery (due to earlier deposition caused by lower particle velocity) of the first active agent, such as DHEA, acting on these distal peripheral airways. Therefore, the combination of a bronchodilator drug (β2 agonist, antimuscarinic which reverses elevated tone) facilitates the delivery of an anti-inflammatory to the distal peripheral airways. Use of the combination provides an improved sustained pharmacologic effect that translates an improved disease management. The antileukotrienes reduce interstitial edema in the very small peripheral airways. This too would have the effect of increasing peripheral airway diameter and facilitate delivery of the first active agent. This is also true for antihistamines, which also reduce peripheral airways edema and facilitate distal airway delivery of the first active agent.

The drawings accompanying this patent form part of the disclosure of the invention, and further illustrate some aspects of the present invention as discussed below.

BRIEF DESCRIPTION OF THE DRAWINGS

Figure 1 depicts fine particle fraction of neat micronized DHEA-S·$2H_2O$ delivered from the single-dose Acu-Breathe inhaler as a function of flow rate. Results are expressed as DHEA-S. IDL data on virtually anhydrous micronized DHEA-S are also shown in this figure where the 30 L/min result was set to zero since no detectable mass entered the impactor.

Figure 2 depicts HPLC chromatograms of virtually anhydrous DHEA-S bulk after storage as neat and lactose blend for 1 week at 50°C. The control was neat DHEA-S stored at room temperature (RT)

Figure 3 depicts HPLC chromatograms for DHEA-S·$2H_2O$ bulk after storage as neat and lactose blend for 1 week at 50°C. The control was neat DHEA-S·$2H_2O$ stored at RT.

Figure 4 depicts solubility of DHEA-S as a function of NaCl concentration at two temperatures.

Figure 5 depicts DHEA-S solubility as a function of the reciprocal sodium cation concentration at 24-25 °C.

Figure 6 depicts DHEA-S solubility as a function of the reciprocal sodium cation concentration at 7-8 °C.

Figure 7 depicts solubility of DHEA-S as a function of NaCl concentration with and without buffer at RT.

Figure 8 depicts DHEA-S solubility as a function of the reciprocal of sodium cation concentration at 24-25 °C with and without buffer.

Figure 9 depicts solution concentration of DHEA-S versus time at two storage conditions.

Figure 10 depicts solution concentration of DHEA versus time at two storage conditions.

Figure 11 depicts the schematic for nebulization experiments.

Figure 12 depicts mass of DHEA-S deposited in by-pass collector as a function of initial solution concentration placed in the nebulizer.

Figure 13 depicts particle size by cascade impaction for DHEA-S nebulizer solutions. The data presented are the average of all 7 nebulization experiments.

Figure 14 depicts the inhibition of HT-29 SF cells by DHEA.

Figure 15 depicts the effects of DHEA on cell cycle distribution in HT-29 SF cells.

Figures 16a and 16b depict the reversal of DHEA-induced growth inhibition in HT-29 cells.

Figure 17 depicts the reversal of DHEA-induced G_1 arrest in HT-29 SF cells.

Figure 18 depicts the effect of EGF, IBMX, and DHEA on 3H-thymidine incorporation in human smooth muscle cultures.

Figure 19 depicts the ability of DHEA alone and in combination with IBMX to inhibit EGF stimulated proliferation.

Figure 20 depicts certain suitable analogs of DHEA.

Figure 21 depicts certain suitable analogs of DHEA.

Figure 22 depicts certain suitable analogs of DHEA.

Figure 23 depicts suitable modifications of the C-17 ketone of DHEA.

DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS

Definitions

In the present context, the terms “adenosine” and “surfactant” depletion are intended to encompass levels that are lowered or depleted in the subject as compared to previous levels in that subject, and levels that are essentially the same as previous levels in that subject but, because of some other reason, a therapeutic benefit would be achieved in the patient by modification of the levels of these agents as compared to previous levels.

The term ”airway”, as used herein, means part of or the whole respiratory system of a subject that is exposed to air. The airway includes, but not exclusively, throat, tracheobronchial tree, nasal passages, sinuses, among others. The airway also includes trachea, bronchi, bronchioles, terminal bronchioles, respiratory bronchioles, alveolar ducts, and alveolar sacs.

The term ”airway inflammation”, as used herein, means a disease or condition related to inflammation on airway of subject. The airway inflammation may be caused or accompanied by allergy(ies), asthma, impeded respiration, cystic fibrosis (CF), Chronic Obstructive Pulmonary Diseases (COPD), allergic rhinitis (AR), Acute

Respiratory Distress Syndrome (ARDS), microbial or viral infections, pulmonary hypertension, lung inflammation, bronchitis, cancer, airway obstruction, and bronchoconstriction.

The term “carrier”, as used herein, means a biologically acceptable carrier in the form of a gaseous, liquid, solid carriers, and mixtures thereof, which are suitable for the different routes of administration intended. Preferably, the carrier is pharmaceutically or veterinarily acceptable.

"An effective amount" as used herein, means an amount which provides a therapeutic or prophylactic benefit.

“Other therapeutic agents” refers to any therapeutic agent is not the first or second active agent of the composition.

The terms “prophylaxis”, as used herein, mean a prophylactic treatment made before a subject experiences a disease or a worsening of a previously diagnosed condition such that it can have a subject avoid, prevent or reduce the probability of having a disease symptom or condition related thereto. The subject can be one of increased risk of obtaining the disease or a worsening of a previously diagnosed condition.

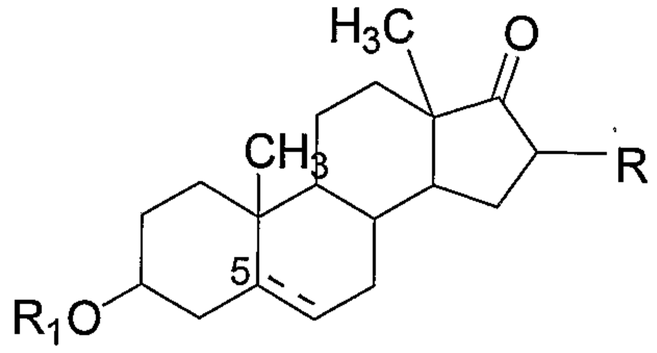
The term ”respiratory diseases”, as used herein, means diseases or conditions related to the respiratory system. Examples include, but not limited to, airway inflammation, allergy(ies), impeded respiration, cystic fibrosis (CF), allergic rhinitis (AR), Acute Respiratory Distress Syndrome (ARDS), cancer, pulmonary hypertension, lung inflammation, bronchitis, airway obstruction, bronchoconstriction, microbial infection, and viral infection, such as SARS.

The terms "treat", "treating" or “therapeutic”, as used herein, mean a treatment which decreases the likelihood that the subject administered such treatment will manifest symptoms of disease or other conditions.

The present invention provides for a composition comprising a first active agent comprising a non-glucocorticoid steroid, such as an epiandrosterone (EA), analogue thereof, or a salt thereof, in combination with a second active agent comprising a β2-agonist bronchodilator. The composition can further comprise a pharmaceutical or veterinarily acceptable carrier, diluent, excipient, bioactive agent or ingredient. The compositions are useful for treating asthma, COPD, or other respiratory diseases. Other respiratory diseases that the compositions are also useful

for treating are lung and respiratory diseases and conditions associated with bronchoconstriction, lung inflammation and/or allergies, and lung cancer.

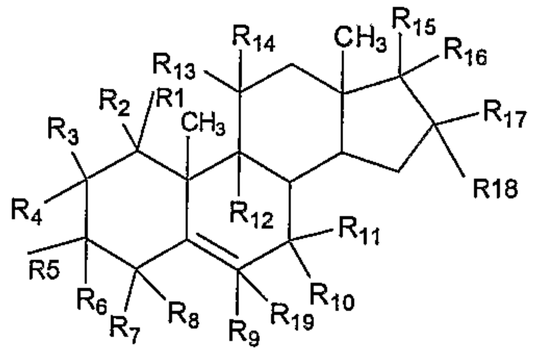
The first active agent is an epiandrosterone, an analogue or a pharmaceutically or veterinarily acceptable salt thereof. The epiandrosterone, an analogue or a pharmaceutically or veterinarily acceptable salt thereof is selected from a non-glucocorticoid steroid having the chemical formula



(I)

wherein the broken line represents a single or a double bond; R is hydrogen or a halogen; the H at position 5 is present in the alpha or beta configuration or the compound of chemical formula I comprises a racemic mixture of both configurations; and R^1 is hydrogen or a multivalent inorganic or organic dicarboxylic acid covalently bound to the compound;

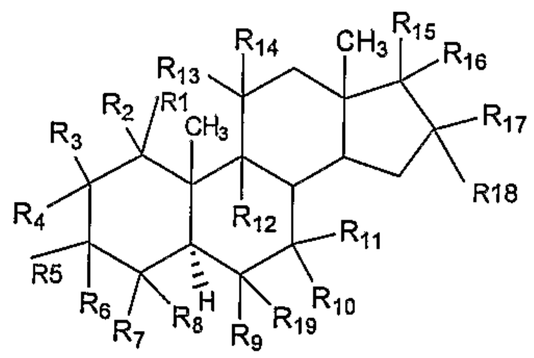
a non-glucocorticoid steroid of the chemical formula



(III)

; or

a non-glucocorticoid steroid of the chemical formula



(IV)

; or a combination thereof

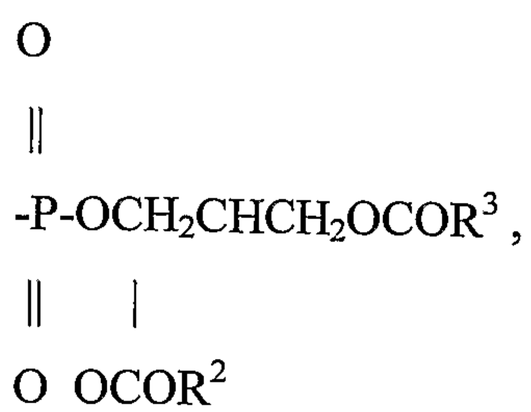
wherein R1, R2, R3, R4. R5, R7, R8, R9, R10, R12, R13, R14 and R19 are independently H, OR, halogen, (C1-C10) alkyl or (C1-C10) alkoxy, R5 and R11 are independently OH, SH, H, halogen, pharmaceutically acceptable ester, pharmaceutically acceptable thioester, pharmaceutically acceptable ether, pharmaceutically acceptable thioether, pharmaceutically acceptable inorganic esters, pharmaceutically acceptable monosaccharide, disaccharide or oligosaccharide,

spirooxirane, spirothirane, -OSO2R20, -OPOR20R21 or (C1-C10) alky, R5 and R6 taken together are =O, R10 and R11 taken together are =O; R15 is (1) H, halogen, (C1-C10) alkyl, or (C1-C10) alkoxy when R16 is -C(O)OR22, (2) H, halogen, OH or (C1-C10) alkyl when R16 is halogen, OH or (C1-C10) alkyl, (3) H, halogen, (C1-C10) alkyl, (C1-C10) alkenyl, (C1-C10) alkynyl, formyl, (C1-C10) alkanoyl or epoxy when R16 is OH, (4) OR, SH, H, halogen, pharmaceutically acceptable ester, pharmaceutically acceptable thioester, pharmaceutically acceptable ether, pharmaceutically acceptable thioether, pharmaceutically acceptable inorganic esters, pharmaceutically acceptable monosaccharide, disaccharide or oligosaccharide, spirooxirane, spirothirane, -OSO2R20 or -OPOR20R21 when R16 is H, or R15 and R16 taken together are =O; R17 and R18 are independently (1) H, -OH, halogen, (C1-C10) alkyl or -(C1-C10) alkoxy when R6 is H OR, halogen. (C1-C10) alkyl or -C(O)OR22, (2) H, (C1-C10 alkyl).amino, ((C1-C10) alkyl)n amino-(C1-C10) alkyl, (C1-C10) alkoxy, hydroxy - (C1-C10) alkyl, (C1-C10) alkoxy - (C1-C10) alkyl, (halogen)m (C1-C10) alkyl, (C1-C10) alkanoyl, formyl, (C1-C10) carbalkoxy or (C1-C10) alkanoyloxy when R15 and R16 taken together are =O, (3) R17 and R18 taken together are =O; (4) R17 or R18 taken together with the carbon to which they are attached form a 3-6 member ring containing 0 or 1 oxygen atom; or (5) R15 and R17 taken together with the carbons to which they are attached form an epoxide ring; R20 and R21 are independently OH, pharmaceutically acceptable ester or pharmaceutically acceptable ether; R22 is H, (halogen)m (C1-C10) alkyl or (C1-C10) alkyl; n is 0, 1 or 2; and m is 1, 2 or 3; or pharmaceutically or veterinarily acceptable salts thereof.

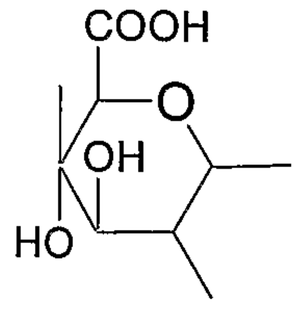
Preferably, for chemical formula (I), the multivalent organic dicarboxylic acid is SO_2OM, phosphate or carbonate, wherein M comprises a counterion. Examples of a counterion are H, sodium, potassium, magnesium, aluminum, zinc, calcium, lithium, ammonium, amine, arginine, lysine, histidine, triethylamine, ethanolamine, choline, triethanoamine, procaine, benzathine, tromethanine, pyrrolidine, piperazine, diethylamine, sulfatide

$$-SO_2O\text{-}CH_2CHCH_2OCOR^3;$$
$$\quad\quad\quad

and phosphatide



wherein R^2 and R^3, which may be the same or different, are straight or branched (C_1-C_{14}) alkyl or glucuronide



The hydrogen atom at position 5 of the chemical formula I may be present in the alpha or beta configuration, or the DHEA compound may be provided as a mixture of compounds of both configurations. Compounds illustrative of chemical formula I above are included, although not exclusively, are DHEA, wherein R and R^1 are each hydrogen, containing a double bond; 16- alpha bromoepiandrosterone, wherein R is Br, R^1 is H, containing a double bond; 16-alpha-fluoro epiandrosterone, wherein R is F, R^1 is H, containing a double bond; Etiocholanolone, wherein R and R^1 are each hydrogen lacking a double bond; and dehydroepiandrosterone sulphate, wherein R is H, R^1 is SO_2OM and M is a sulphatide group as defined above, lacking a double bond. Others, however, are also included. Also preferred compounds of formula I are those where R is halogen, e.g. bromo, chloro, or fluoro, where R1 is hydrogen, and where the double bond is present. A most preferred compound of formula I is 16-alpha-fluoro epiandrosterone. Other preferred compounds are DHEA and DHEA salts, such as the sulfate salt (DHEA-S).

In general, the non-glucocorticoid steroid, such as those of formulas (I), (III) and (IV), their derivatives and their salts are administered in a dosage of about 0.05, about 0.1, about 1, about 5, about 20 to about 100, about 500, about 1000, about 1500 about 1800, about 2500, about 3000, about 3600 mg/kg body weight. Other dosages, however, are also suitable and are contemplated within this patent. The first active agent of formula (I), (III) and (IV) may be made in accordance with known

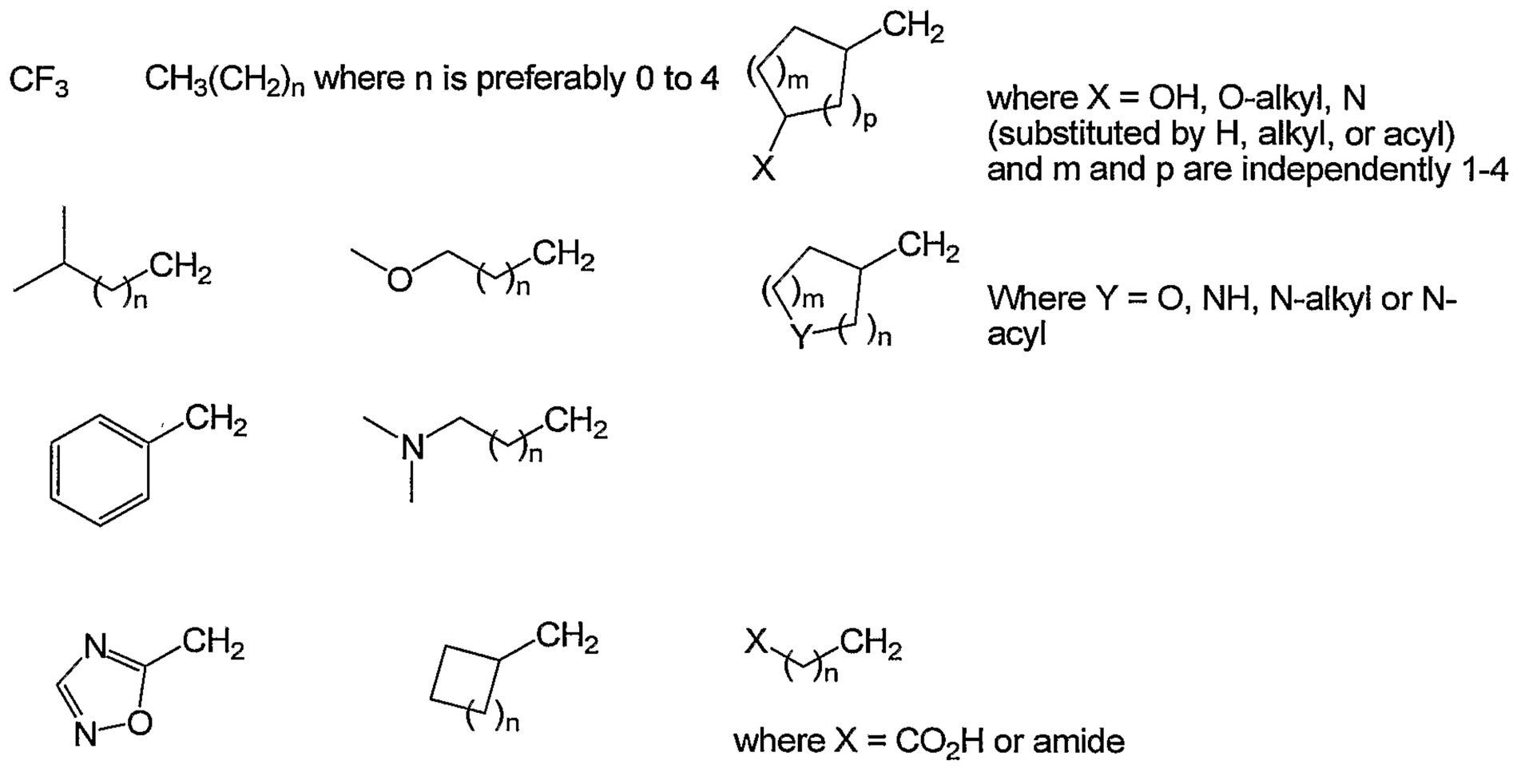
procedures, or variations thereof that will be apparent to those skilled in the art. See, for example, U.S. Patent No. 4,956,355; UK Patent No. 2,240,472; EPO Patent Application No. 429; 187, PCT Patent Publication No. WO 91/04030; U.S. Patent No. 5,859,000; Abou-Gharbia et al., J. Pharm. Sci. 70: 1154-1157 (1981); Merck Index Monograph No. 7710 (11th Ed. 1989), among others.

In some embodiments of the invention, the first active agent can be an epiandrosterone analog or derivative thereof. Also, prodrugs and active metabolites of epiandrosterone are encompassed by the present invention. Those of skill in the art will recognize that the compounds described herein may exhibit the phenomena of tautomerism, conformational isomerism, geometric isomerism and/or optical isomerism. It should be understood that the invention encompasses any tautomeric, conformational isomeric, optical isomeric, and/or geometric isomeric forms of the compounds having one or more of the utilities described herein, as well as mixtures of these various different forms.

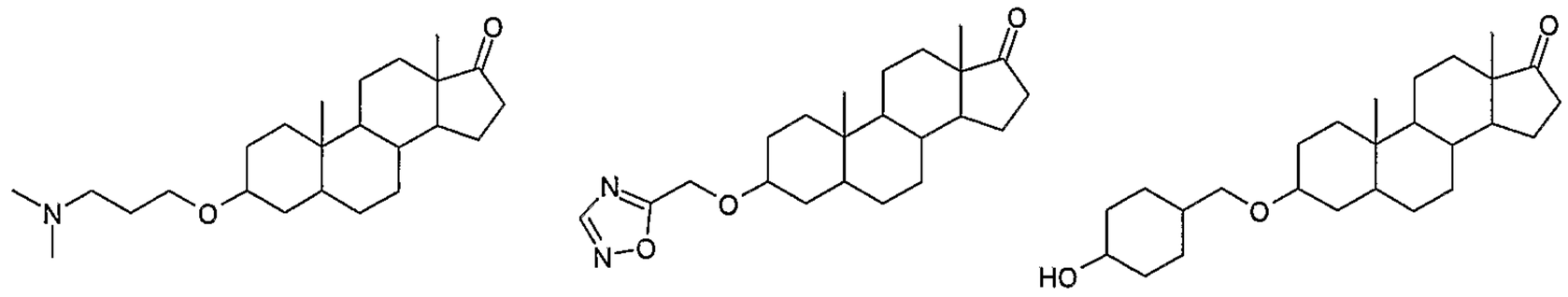
Metabolites of epiandrosterones, such as those described in the following references maybe used as the first active agent – Capillary gas chromatography of urinary steroids of terbutaline-treated asthmatic children, Chromatographia (1998), 48(1/2), 163-165; Androstenedione metabolism in human lung fibroblasts; Journal of Steroid Biochemistry (1986), 24(4), 893-7; Metabolism of androsterone and 5α-androstane-3α,17β-diol in human lung tissue and in pulmonary endothelial cells in culture, Journal of Clinical Endocrinology and Metabolism (1985), 60(2), 244-50; Testosterone metabolism by human lung tissue, Journal of Steroid Biochemistry (1978), 9(1), 29-32; Metabolism of androsterone and 5 alpha-androstane-3 alpha,17 beta-diol in human lung tissue and in pulmonary endothelial cells in culture, Journal of clinical endocrinology and metabolism (1985 Feb), 60(2), 44-50; Metabolism of dehydroisoandrosterone and androstenedione in human pulmonary endothelial cells in culture, Journal of clinical endocrinology and metabolism (1983 May), 56(5), 930-5; Metabolism of dehydroisoandrosterone and androstenedione by the human lung in vitro, Journal of steroid biochemistry (1977 Apr), 8(4), 277-84; and Testosterone metabolism in dog lung in vitro, Steroids and lipids research (1973), 4(1), 17-23, which references are herein incorporated by reference in their entirety.

Other suitable analogs of DHEA that can be used as the first active agent are described herein. Figure 20 depicts certain suitable analogs of DHEA, including

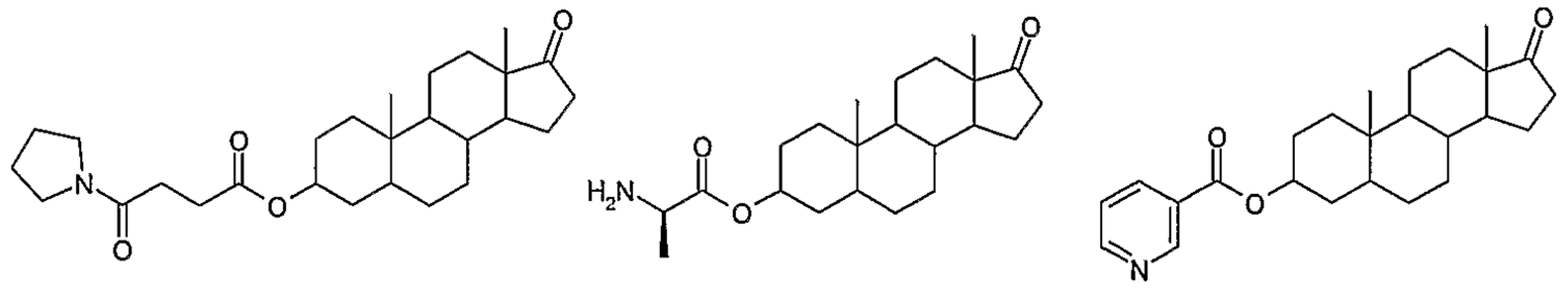
compounds of the Formulas IA, IB, IC, and ID. In the depiction of suitable R groups herein, the attachment point is indicated by a CH2 group or by an atom marked with an asterisk. R1 and R3 can be linear or branched alkyls including benzyl and optionally substituted alkyls, such as aminoalkyls, hydroxyalkyls, ethers, and carboxylic acids, and optionally substituted aryl and heteroaryls. R1 and R3 can be, for example,



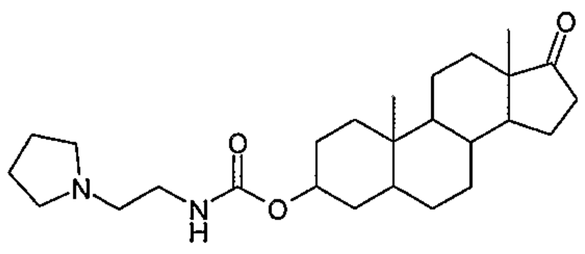
Examples of compounds of formula IA include,

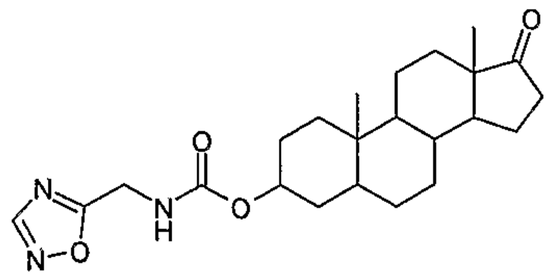


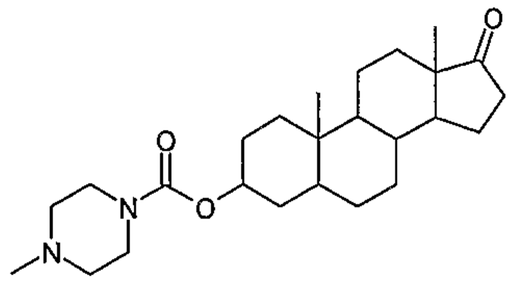
R2 is preferably a diacid-derived or amino acid derived substituent, potentially including chloracetyl derivatives and acrylate derivatives, or optionally substituted aryls such as benzyl and heterobenzoyl. Examples of compounds of formula IB include,



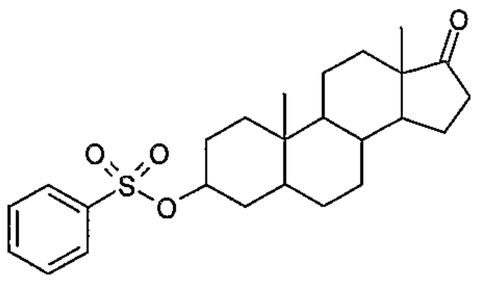
Examples of compounds of formula IC include,

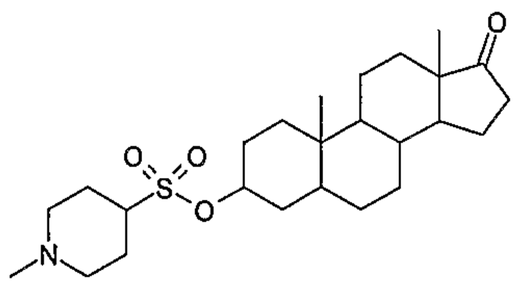






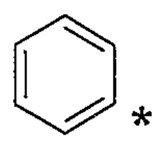
R4 can be aromatic in nature and examples of suitable compounds of formula ID include,

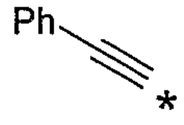




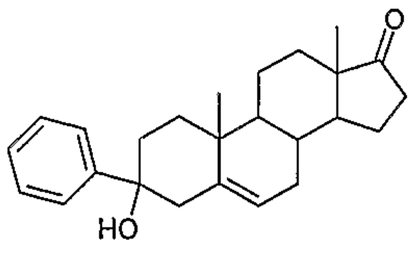
Other suitable analogs include the analogs wherein modifications have been performed at C-3 position by retaining OH or replacing OH with NH. These analogs are typically made by starting from C-17 acetal protected andrst-4-ene3, 17-dione, such as depicted in Figure 21. Compounds of Formula IE can be derived from Grignard reagents and possibly aryl-lithium reagents, such as aromatic in nature, and can also be alkynyl, alkenyl, and alkyl. Examples of R5 are

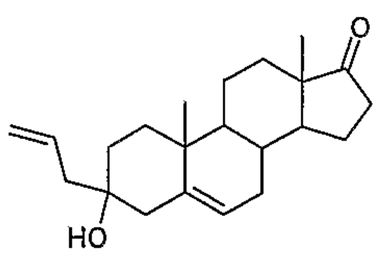
$CH_3(CH_2)_n$ where n=0 to 4

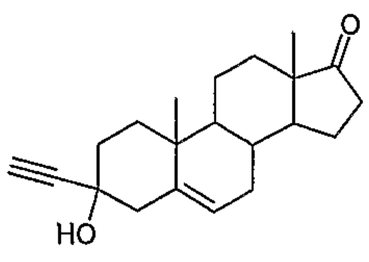




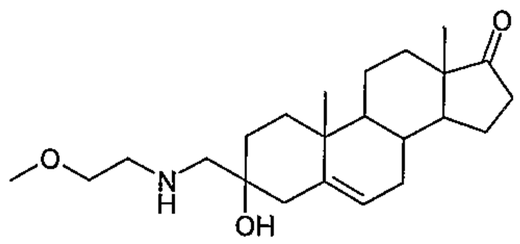
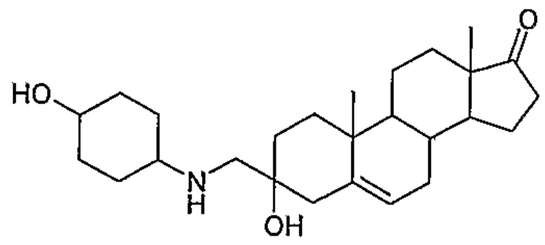
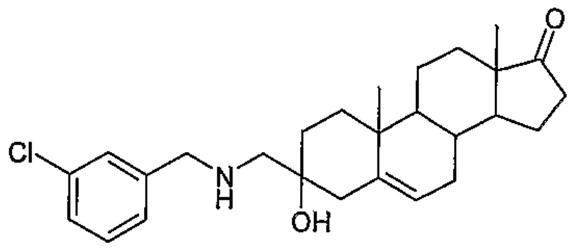
Examples of compounds of Formula IE include



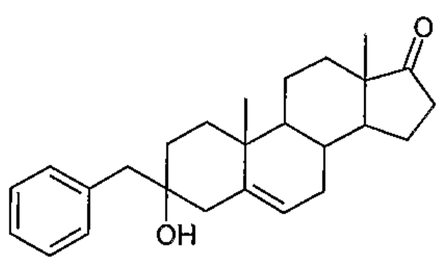
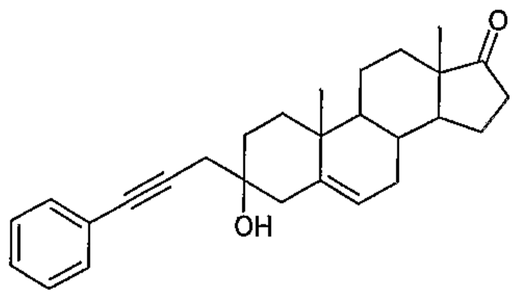
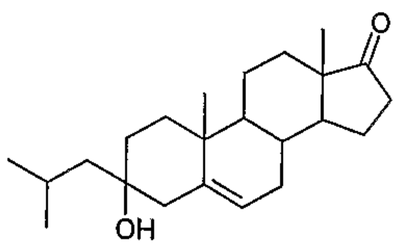




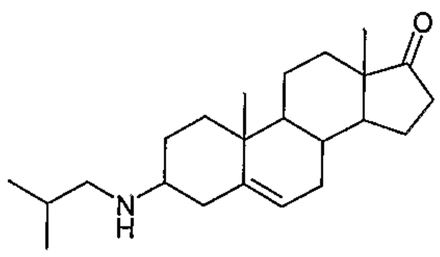
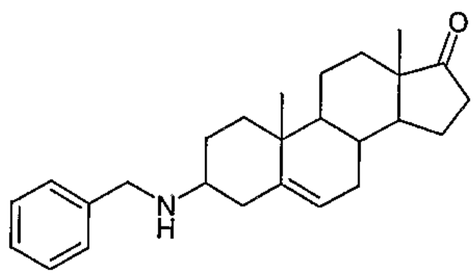
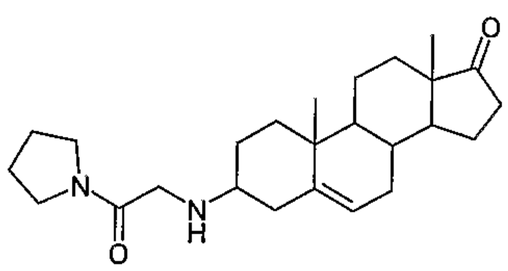
R6 and R8 can independently be a diverse set of amines and can include amines possessing the functionalities as described for the R1 group. Examples of suitable compounds of Formula IF include,

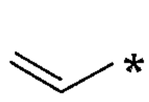
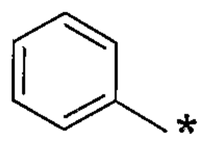
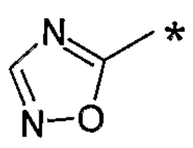
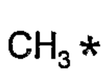
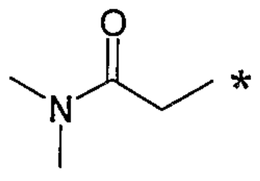
Suitable R7 groups can be derived from Grignard/organolithium reagents and so could encompass the functionalities described for R5. Examples of compounds of Formula IG include

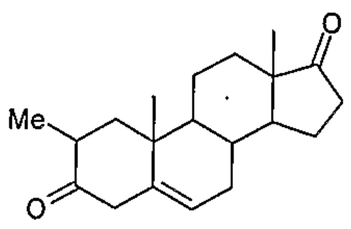
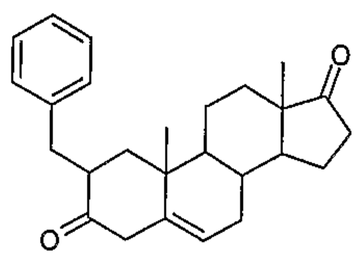
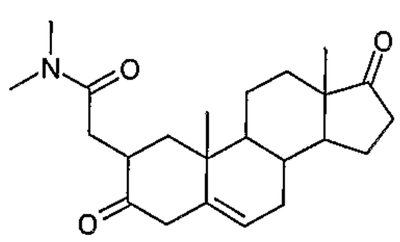
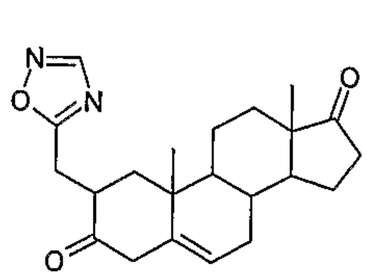
Examples of compounds of Formula IH include

Other suitable analogs include those compounds wherein the C-2 position of DHEA was modified. Suitable modifications are depicted in Figure 22. R9 can be derived from alkylating agents, such as alkyl, benzyl, heterobenzyl, and derivatives of other activated halides. Examples of R9 include,

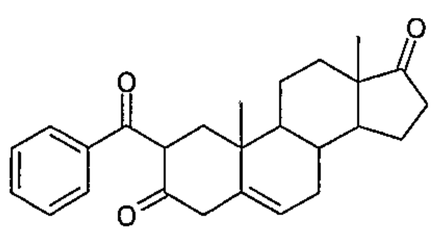
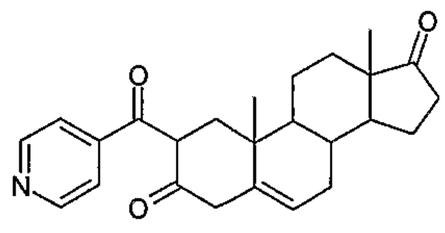
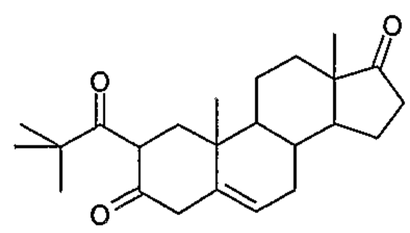






Example of Formula IJ compounds include,

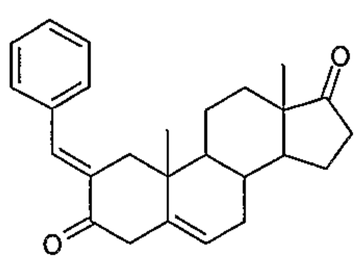
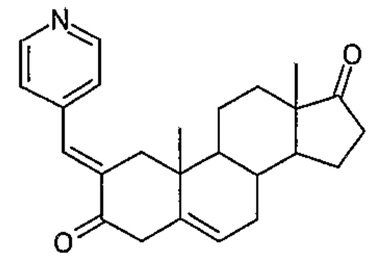
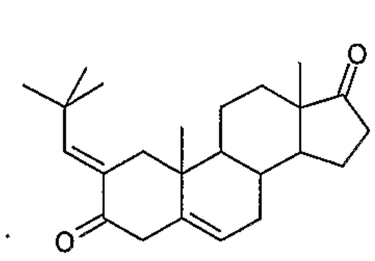





R10 can be aromatic esters such as with aryl or heteroaryl ring or enolisable alkyl

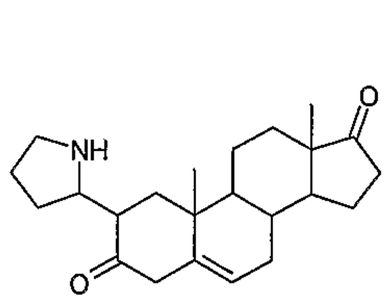
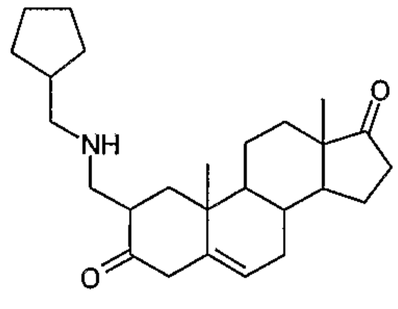
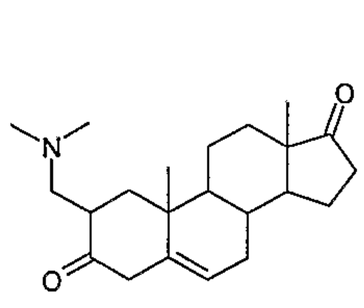
esters. Examples of compounds of formula IK include,

R11 can be a series of aromatic and heteroaromatic aldehydes such as benzene carboxaldehyde and substituted variants thereof, pyridine carboxaldehyde or non-enolisable aldehydes such as (CH3)3CCH=O. Examples of compounds of Formula IL include,

R12 can be a subset of an amine such as for R6. Examples of compounds of formula IN include,

Suitable modifications of the C-17 ketone of DHEA are depicted in Figure 23. The compounds depicted in Figure 23 can also be used as the first active agent.

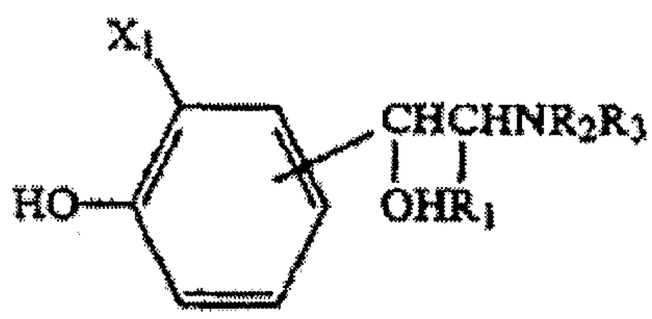
Other suitable DHEA analogs are described in U.S. Patent 6,635,629; European Patent 934745; Dehydroepiandrosterone and analogs inhibit DNA binding of AP-1 and airway smooth muscle proliferation, Journal of Pharmacology and Experimental Therapeutics (1998), 285(2), 876-883; and Dehydroepiandrosterone and related steroids inhibit mitochondrial respiration in vitro, International Journal of Biochemistry (1989), 21(10), 1103-7, all of which are herein incorporated by reference in their entirety.

The second active agent is a bronchodilator comprising a long-acting β2-agonist. Bronchodilators relax the muscle bands that tighten around the airways, thereby allowing more air in and out of the lungs and improve breathing. Bronchodilators also help clear mucus from the lungs, and as the airways open, the mucus moves more freely and can be coughed or cleared out more easily. There are short-acting and long-acting forms of bronchodilators: the short-acting form relieves or stops asthma symptoms while the long-acting form provide control of asthma

symptoms and prevent asthma attacks.

The range of β2 agonists encompassed in this invention encompasses the compounds defined in U.S. Patent Nos. 3,994,974; 4,600,710; 4,894,219; 4,992,474; 5,108,363; 5,126,375; 5,225,445; 5,234,404; 5,258,385; 5,286,252; 5,460,605; 5,684,199; 6,156,503; 6,297,382; and 6,441,181 (herein incorporated by reference in their entirety). Examples of β2 agonists are ephedrine, isoproterenol, isoetharine, epinephrine, metaproterenol (short-acting form(s): Alupent®, Boehringer Ingelheim (Canada) Ltd, Metaprel), terbutaline (short-acting form(s): Brethaire, Riker Laboratories, Inc., Brethine, Novartis), fenoterol, procaterol, albuterol (short-acting form(s): albuterol (generic, IVAX Corp.; Ventolin, Allen & Hanburys, Proventil, Schering Corp., Proventil HFA, Key pharmaceuticals, Inc; long-acting forms: Proventil Repetabs, Schering Corp., Volmax, Muro Pharmaceutical), salmeterol (long-acting forms: Serevent, GlaxoSmithKline, Severent Diskus, Glaxo Wellcome), pirbuterol (short-acting form(s): Maxair, 3M Pharmaceuticals), formoterol (long-acting forms: Foradil Aerolizer, Schering Corporation), bitolterol (short-acting form(s): Tornalate, Elan Pharmaceuticals and Sanofi), levalbuterol (short-acting form(s): Xoponex, Sepracor), bambuterol, salbutamol, and seretide, among others. Preferably, the long-acting β2-agonist is salmeterol or formoterol.

Salmeterol comprises compounds defined by chemical formulae (V) and (VI). The compounds defined by chemical formulae (V) (British Patent Specification No. 1200886) are:

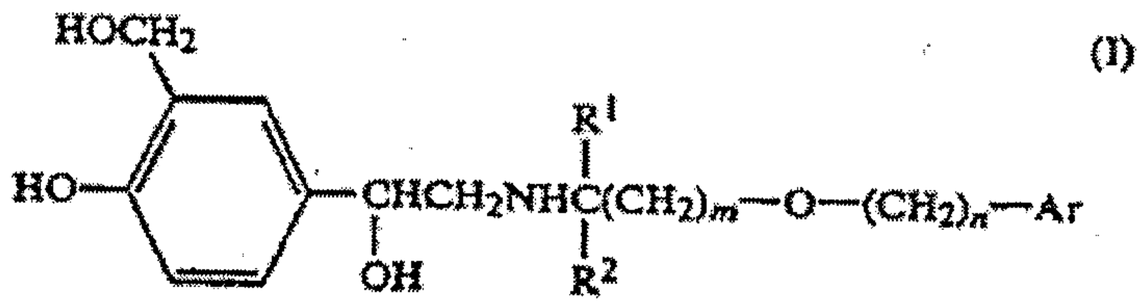


where, inter alia, X 1 is hydroxyalkyl, R1 and R is each a hydrogen atom, and R3 is straight or branched C_{1-6} alkyl, aralkyl or aryloxyalkyl. One compound from within this particular group has been developed for clinical use. This is salbutamol [(α1 -tert-butylaminomothyl)-4-hydroxy-m-xylene-α1,α3 -diol; X1 =CH_2 OH, R1 =--H; R2 =--H; R3 =t-butyl, above] which at the present time is widely prescribed for the treatment of conditions such as bronchial asthma and chronic bronchitis. The success

of salbutamol devolves from its profile of action, in particular its potency, coupled with a selective stimulant action at β2-adrenoreceptors.

All β2-stimulants currently used in clinical practice suffer from the disadvantage that they have a relatively short duration of action when administered by inhalation. A β2-stimulant with a relatively long duration of action would therefore offer a significant advance in the treatment of bronchial asthma and related disorders.

The long-acting β2-agonist also includes compounds of the chemical formula (VI):



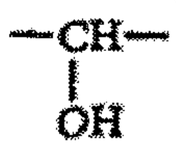
(I)

wherein m is an integer from 2 to 8 and n is an integer from 1 to 7 with the proviso that the sum total of m+n is 4 to 12; Ar represents a phenyl group optionally substituted by one or two substituents selected from halogen atoms, C_{1-3} alkyl or C_{1-3} alkoxy groups, or by an alkylenedioxy group of formula --$O(CH_2)p$ O-- where p is 1 or 2; and R1 and R2 each represents a hydrogen atom or a C_{1-3} alkyl group with the proviso that the sum total of carbon atoms in R1 and R2 is not more than 4; and physiologically acceptable salts and solvates (e.g. hydrates) thereof.

The compounds of general formula (VI) possess one or two asymmetric carbon atoms, namely the carbon atom of the

group and, when R1 and R2 are different groups, the carbon atom to which these groups are attached.

The compounds according to the invention thus include all enantiomers, diastereoisomers and mixtures thereof, including racemates. Compounds in which the carbon atom in the



group is in the R configuration are preferred.

In the general formula (VI), the chain --$(CH_2)_m$ -- may be, e.g., --(CH_2) 3 --, --(CH_2) 4 --, --(CH_2) 5 --, --(CH_2) 6 -- or --(CH_2) 7 --, and the chain --(CH_2)n -- may be, e.g., --(CH_2) 2 --, --(CH_2) 3 --, --(CH_2) 4 --, --(CH_2) 5 -- or (CH_2) 6 --.

Preferably the sum total of the number of carbon atoms in the chains --(CH_2)m -- and --(CH_2)n is 6 to 12 inclusive and may be, e.g., 7, 8, 9 or 10. Compounds wherein the sum total of m+n is 7, 8 or 9 are particularly preferred.

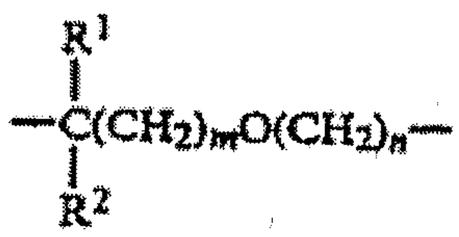
Preferred compounds of general formula (VI) are those wherein m is 3 and n is 6, or m is 4 and n is 3, 4 or 5, or m is 5 and n is 2, 3, 4 or 5, or m is 6 and n is 2 or 3.

R1 and R2, e.g., may each be methyl, ethyl, propyl, or isopropyl groups except that if one of R1 and R2 is a propyl or isopropyl group, the other is a hydrogen atom or a methyl group. Thus, e.g., R1 may be a hydrogen atom or a methyl, ethyl or propyl group. R2, e.g., may be a hydrogen atom or a methyl group.

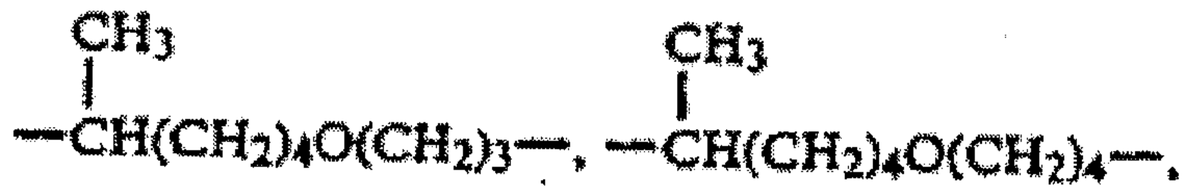
R1 and R2 are each preferably a hydrogen atom or a methyl group.

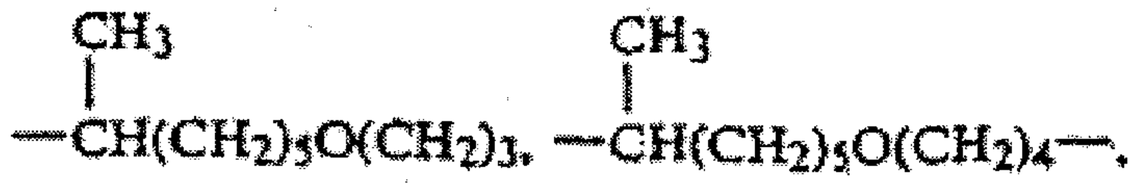
A preferred group of compounds is that wherein R1 and R2 are both hydrogen atoms. In another preferred group of compounds R1 is a hydrogen atom and R2 is a C_{1-3} alkyl group, particularly a methyl group. In yet another preferred group of compounds R1 and R2 are both methyl groups.

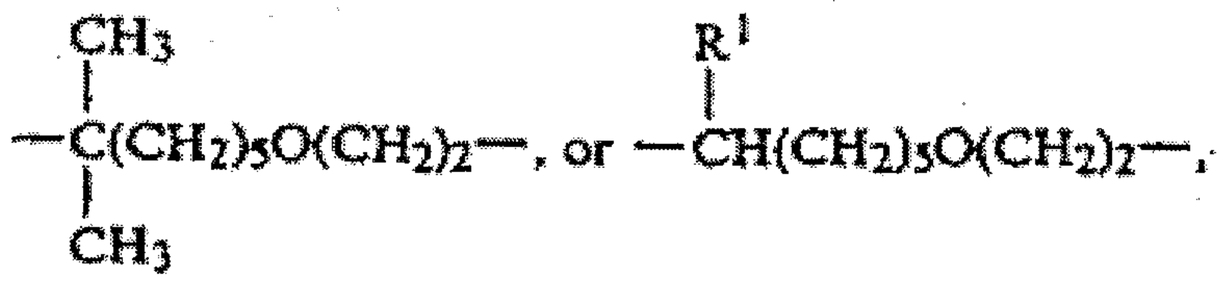
The chain



in general formula (VI) may be, e.g. --(CH_2)$_4$ O(CH_2)$_4$ --, (CH_2)$_5$ O(CH_2)$_2$ -- --(CH_2)$_5$ O(CH_2)$_3$, --(CH_2)$_5$ O(CH_2)$_4$ --,







where R1 is methyl, ethyl or propyl.

Examples of the optional substituents which may be present on the phenyl group represented by Ar include bromine, iodine or, in particular, chlorine or fluorine atoms, or methyl, ethyl, methoxy or ethoxy groups. In general, Ar is preferably an unsubstituted phenyl group. According to another preference, Ar is a phenyl group substituted by one substituent, particularly a fluorine or chlorine atom or a methoxy or methyl group.

Suitable physiologically acceptable salts of the compounds of general formula (V) include acid addition salts derived from inorganic and organic acids, such as hydrochlorides, hydrobromides, sulphates, phosphates, maleates, tartrates, citrates, benzoates, 4-methoxybenzoates, 2- or 4-hydroxybenzoates, 4-chlorobenzoates, p-toluenesulphonates, methanesulphonates, ascorbates, salicylates, acetates, fumarates, succinates, lactates, glutarates, gluconates, tricarballylates, hydroxynaphthalenecarboxylates e.g. 1-hydroxy- or 3-hydroxy-2-naphthalenecarboxylates, or oleates. The compounds may also form salts with suitable bases. Examples of such salts are alkali metal (e.g. sodium and potassium), and alkaline earth metal (e.g. calcium and magnesium) salts.

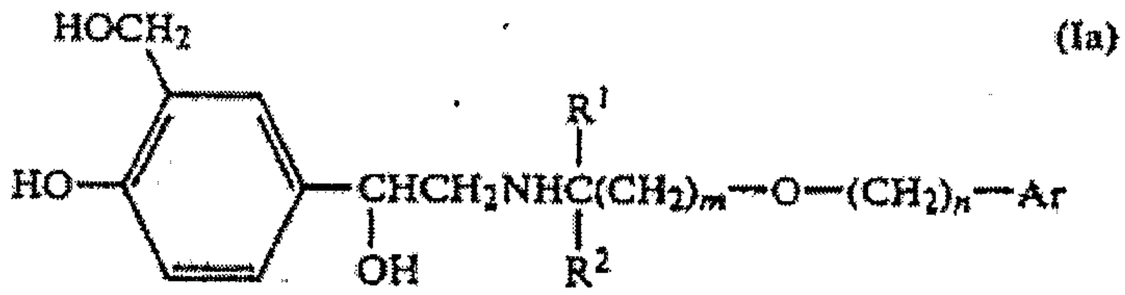
The compounds of the second active agent have a selective stimulant action at β2 -adrenoreceptors, which furthermore is of a particularly advantageous profile. The stimulant action have been demonstrated in the guinea-pig, where compounds have been shown to cause relaxation of PGF2α-contracted isolated trachea. In another test, compounds of the invention have been shown to afford protection against histamine-induced broncho-constriction when administered by inhalation or by an oral route in conscious guinea-pigs. In both tests, compounds according the invention have shown a particularly long duration of action. These compounds have been demonstrated in the rat or guinea pig. Where compounds were shown to have little or no effect on

isolated rat or guinea pig left atria (β1-adrenoreceptor tissues) at concentrations where they cause relaxation of PGF2α -contracted isolated trachea. Compounds according to the invention have also been shown to inhibit the anaphylactic release of spasmagens and inflammagens from sensitised human tissues e.g. lung fragments (U.S. Patent Nos. 4,992,474; 5,126,375; and, 5,225,445; herein incorporated by reference in their entirety.)

The compounds according to the invention may be used in the treatment of diseases associated with reversible airways obstruction such as asthma, COPD, and chronic bronchitis.

The compounds according to the invention may also be used for the treatment of premature labour, depression and congestive heart failure, and are also indicated as useful for the treatment of inflammatory and allergic skin diseases, psoriasis, proliferative skin diseases, glaucoma, and in the treatment of conditions in which there is an advantage in lowering gastric acidity, particularly in gastric and peptic ulceration.

A particularly important group of compounds, which has been shown to have an advantageously long duration of action (U.S. Patent Nos. 4,992,474; 5,126,375; and, 5,225,445; herein incorporated by reference in their entirety), has the chemical formula (VII):



(Ia)

in which R1 and R2 are as defined for general formula (VI); m is an integer from 3 to 6, n is an integer from 2 to 6, and Ar is phenyl or phenyl substituted by a methoxy or methyl group, or more preferably a fluorine or chlorine atom, and the physiologically acceptable salts and solvates thereof, in each instance the sum total of carbon atoms in the chains --$(CH_2)_m$ -- and --$(CH_2)_n$ -- being an integer from 7 to 10 inclusive, in particular 7, 8 or 9.

A preferred group of compounds of chemical formula (VII) is that wherein R1 and R2 is each a hydrogen atom.

In another preferred group of compounds of chemical formula (VII) R1 is a hydrogen atom or a methyl group and R2 is a methyl group.

In a further group of compounds of chemical formula (VII) R1 and R2 each is a hydrogen atom and Ar is phenyl or phenyl substituted by a methoxy group, or more preferably a fluorine or chlorine atom.

A particularly preferred group of compounds has the chemical formula (VII) in which R1 and R2 each is a hydrogen atom or a methyl group, m is 4 or 5, n is 2, 3 or 4, and Ar is phenyl or phenyl substituted by a chlorine or fluorine atom or a methoxy or methyl group and the physiologically acceptable salts and solvates thereof.

Particularly important compounds are:

4-hydroxy-α1 [[[6-(4-phenylbutoxy)hexyl]amino]methyl]-1,3-benzenedimethanol; and the physiologically acceptable salts thereof;

4-hydroxy-α1 [[[6-(3-phenylpropoxy)hexyl]amino]methyl]-1,3-benzenedimethanol; and the physiologically acceptable salts thereof;

4-hydroxy-α1 -[[[6-(2-phenylethoxy)hexyl]amino]methyl]-1,3-benzenedimethanol; and the physiologically acceptable salts thereof;

4-hydroxy-α1 -[[[5-(4-phenylbutoxy)pentyl]amino]methyl]-1,3-benzenedimethanol; and the physiologically acceptable salts thereof;

4-hydroxy-α1 -[[[1-methyl-6-(2-phenylethoxy)hexyl]amino]methyl]-1,3-benzenedimethanol; and the physiologically acceptable salts thereof;

4-hydroxy-α1 -[[[1-methyl-5-(3-phenylpropoxy)pentyl]amino]methyl]-1,3-benzenedimethanol ; and the physiologically acceptable salts thereof;

4-hydroxy-α1 -[[[1-methyl-5-(4-phenylbutoxy)pentyl]amino]methyl]-1,3-benzenedimethanol; and the physiologically acceptable salts thereof;

4-hydroxy-α1 -[[[1-ethyl-6-(2-phenylethoxy)hexyl]amino]methyl]-1,3-benzenedimethanol; and the physiologically acceptable salts thereof;

α1 -[[[1,1-dimethyl-6-(2-phenylethoxy)hexyl]amino]methyl-4-hydroxy-1,3-benzen edimethanol; and the physiologically acceptable salts thereof;

α1 -[[[6-[2-(4-fluorophenyl)ethoxy]-1-methylhexyl]amino]methyl]-4-hydroxy-1,3 - benzenedimethanol; and the physiologically acceptable salts thereof;

4-hydroxy-α1 -[[[6-[3-(4-methoxyphenyl)propoxy]-1-methylhexyl]amino]methyl]-1,3-benzene dimethanol; and the physiologically acceptable salts thereof;

4-hydroxy-α1 -[[[1-methyl-6-(4-phenylbutoxy)hexyl]amino]methyl]-1,3-

benzenedimethanol; and the physiologically acceptable salts thereof;

4-hydroxy-α1 -[[[6-[2-(4-methylphenyl)ethoxy]hexyl]amino]methyl]-1,3-benzenedimethanol; and the physiologically acceptable salts thereof;

α1 -[[[6-[2-(3-chlorophenyl)ethoxy]hexyl]amino]methyl]-4-hydroxy-1,3-benzened imethanol; and the physiologically acceptable salts thereof;

4-hydroxy-α1 -[[[6-[2-(4-methoxyphenyl)ethoxy]hexyl]-amino]-methyl]-1,3-benzenedimethan ol; and the physiologically acceptable salts thereof;

α1 -[[[6-[3-(4-fluorophenyl)propoxy]hexyl]amino]methyl]-4-hydroxy-1,3-benzene dimethanol; and the physiologically acceptable salts thereof.

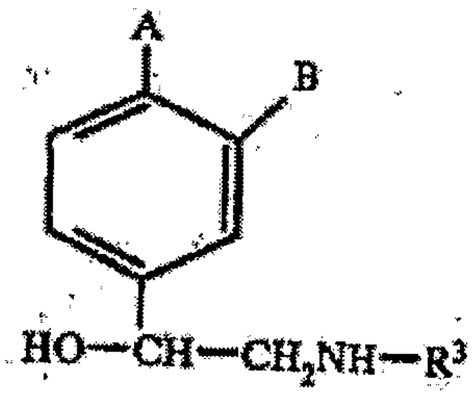
The invention accordingly further provides compounds of formula (V) and their physiologically acceptable salts and solvates for use in the therapy or prophylaxis of diseases associated with reversible airways obstruction in human or animal subjects. The invention also provides compounds of chemical formula (V) and their physiologically acceptable salts and solvates and compositions containing them in association with instructions for their use in the therapy or prophylaxis of diseases associated with reversible airways obstruction in human or animal subjects.

The compounds according to the invention may be formulated for administration in a convenient way. The invention therefore includes within its scope pharmaceutical compositions comprising at least one compound of chemical formula (V) or a physiologically acceptable salt or solvate thereof formulated for use in human or veterinary medicine. Such compositions may be presented for use with physiologically acceptable carriers or excipients, optionally with supplementary medicinal agents.

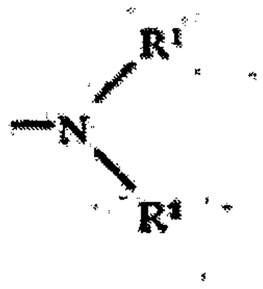
A proposed daily dosage of the second active agent is 0.0005 mg to 100 mg, which may be conveniently administered in one or two doses. The precise dose employed will of course depend on the age and condition of the patient and on the route of administration. Thus a suitable dose for administration by inhalation is 0.0005 mg to 10 mg, for oral administration is 0.02 mg to 100 mg, and for parenteral administration is 0.001 mg to 2 mg.

The salmeterol compounds are prepared and isolated by the methods described in U.S. Patent Nos. 4,992,474; 5,126,375; and, 5,225,445; herein incorporated by reference in their entirety.

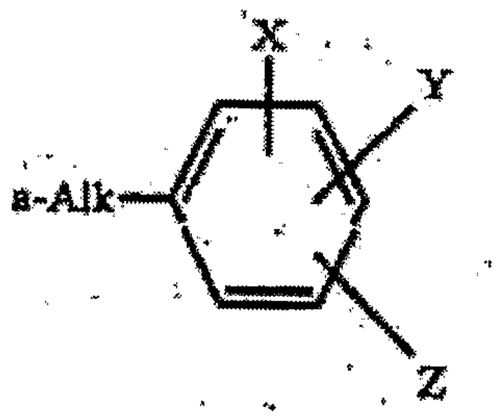
Formoterol comprises α-aminomethylbenzyl alcohol derivatives represented by the chemical formula (VIII):



wherein one of A and B represents a hydrogen atom or a hydroxyl group while the other of them represents a



group (in which R1, which is different from R2, represents a hydrogen atom or a C_{1-7} alkyl and R2 represents a hydrogen atom or a --CO--R4 group in which R4 represents a hydrogen atom, a C_{1-7} hydroxyalkyl or a C_{2-10} alkanoylaminoalkyl) and R3 represents a C_{3-7} alkyl, a C_{6-12} cycloalkylalkyl or



Group (wherein Alk represents a straight or branched C_{1-7} alkylene, and X, Y and Z are the same of different from each other and each represents a hydrogen atom, a hydroxy group, a C_{1-7} alkanoylamino, a C_{1-7} alkyl or a C_{1-7} alkoxy).

Formoterols have utility as β-adrenergic stimulants and thus have great activity on respiratory smooth muscle and are suitable as bronchodilating agents.

As a compound having bronchodilating effects, there are known hitherto various compounds and especially, Isoproterenol and Trimetoquinol, these compounds being well known among the bronchodilating drugs and being widely sold since they have strong effects. Further a bronchlodilating agent should not have ill-

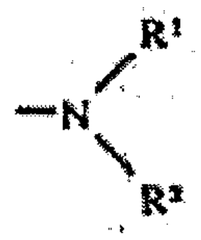
effects on the heart that is, it should have high selectivity, and Salbutamol satisfies this requirement and is also widely sold.

Further, compounds having a similar structure to the present compounds are the known 3-amino-4-hydroxy-α-isopropylaminomethylbenzyl alcohol (Dutch Patent No. 85197: "Chemical Abstract", 52 11121d (1958)), 3-ethoxycarbonylamino-4-hydroxy-α-isopropylaminomethylbenzyl alcohol (Belgian Patent No. 765,986), α-(isopropylaminomethyl)-4-hydroxy-3-ureido benzyl alcohol (Japanese Patent Application Public Open No. 2676/1971). These patents and patent applications are herein incorporated by reference in their entirety.

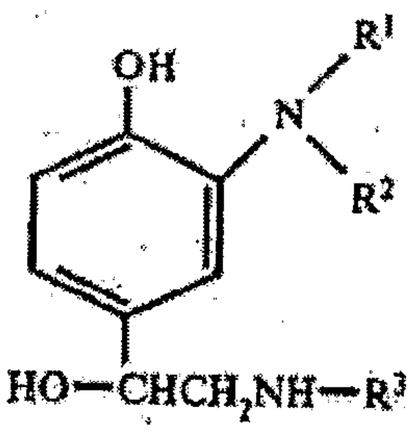
In chemical formula (VIII) representing the compounds of this invention, examples of R1 are a hydrogen atom, an alkyl group such as a methyl group, an ethyl group, a propyl group, an isopropyl group, a n-butyl group, a tert butyl group, a 1, 3-dimethylbutyl group, a 1, 3-dimethylpentyl group, a 2, 3-dimethylbutyl group, a 2, 3, 3-tributyl group, etc.; examples of R4 are a hydrogen atom, a hydroxyalkyl group such as a hydroxymethyl group, a hydroxyethyl group, a hydroxypropyl group, a hydroxybutyl group, etc., an alkanoylaminoalkyl group such as a formamidemethyl group, an acetylaminomethyl group, an acetylaminoethyl group, an acetylaminopropyl group, a butyrylaminoethyl group, etc. Examples of an alkyl group of R3 are a propyl group, an isopropyl group, a n-butyl group, a tert-butyl group, a 1, 3-dimethylbutyl group, a 1, 3-dimethylpentyl group, a 2, 3-dimethylbutyl group, a 2, 3, 3-trimethylbutyl group, etc.; examples of the cycloalkyl group of R3 are a cyclopentylmethyl group, a 2-cycloethyl group, a cyclohexylmethyl group, a 2-cyclohexylethyl group, a 3-cyclohexyl-1-methylpropyl group, etc. Examples of Alk are an alkylene group such as methylene group, an ethylene group, a propylene group, a butylene group, a 1-methylethylene group, a 1-ethylethylene group, a 1-methylpropylene group, a 1-ethylpropylene group, a 2-methylpropylene group, a 3-methylbutylene group, a 2-ethylbutylene group, etc. The examples of X, Y and Z are a hydrogen atom, a hydroxy group, al alkanoylamino group such as a formamide group, an acetylamino group, a propionylamino group, a butyrylamino group, etc., an alkyl group such as a methyl group, an ethyl group, a propionyl group, an isopropyl group, an isobutyl group, a tert-butyl group, etc., or an alkoxy group such as a methoxy group, an ethoxy group, a propoxy group, an isopropoxy group, a butoxy group, etc.

The particularly useful compounds of this invention are 3-formamido-4-hydroxy-α-[N-(1-methyl-2-p-hydroxyphenylethyl) amino-methyl] benzyl alcohol, 3-formamido-4-hydroxy-α-[N-(1-methyl-2-p-methoxyphenylethyl) aminomethyl]benzyl alcohol, 4-hydroxy-3-methylamino-α-(N-tert-butylaminomethyl) benzyl alcohol, etc.

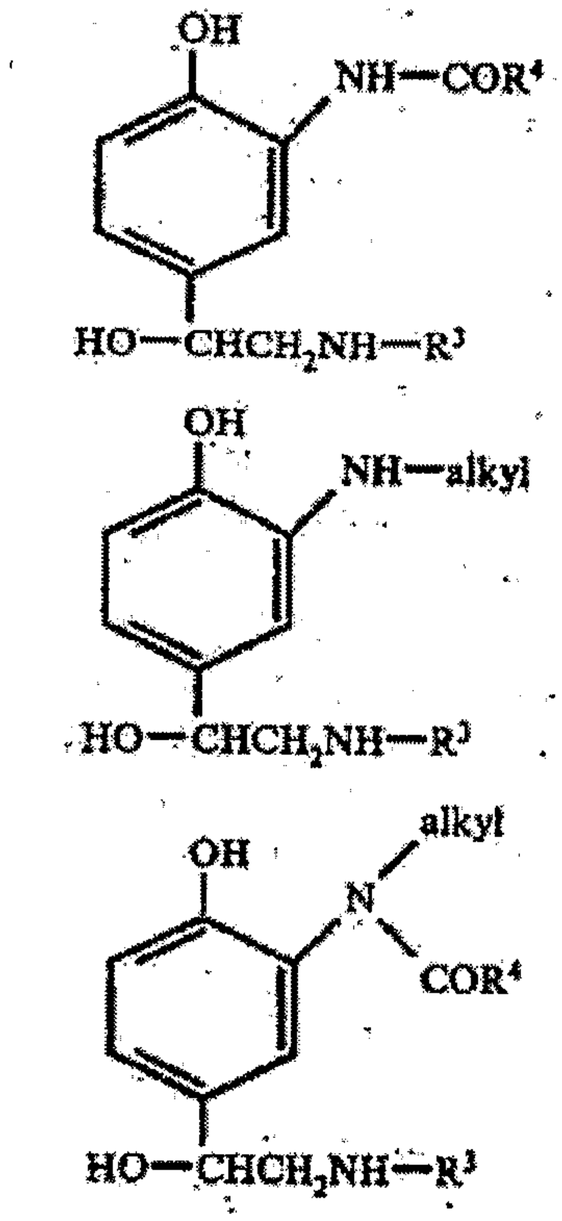
When A of the chemical formula (VIII) representing the compounds of this invention is a hydroxyl group and B is the



group, the compounds of this invention are shown by the chemical formula (VIII) are:



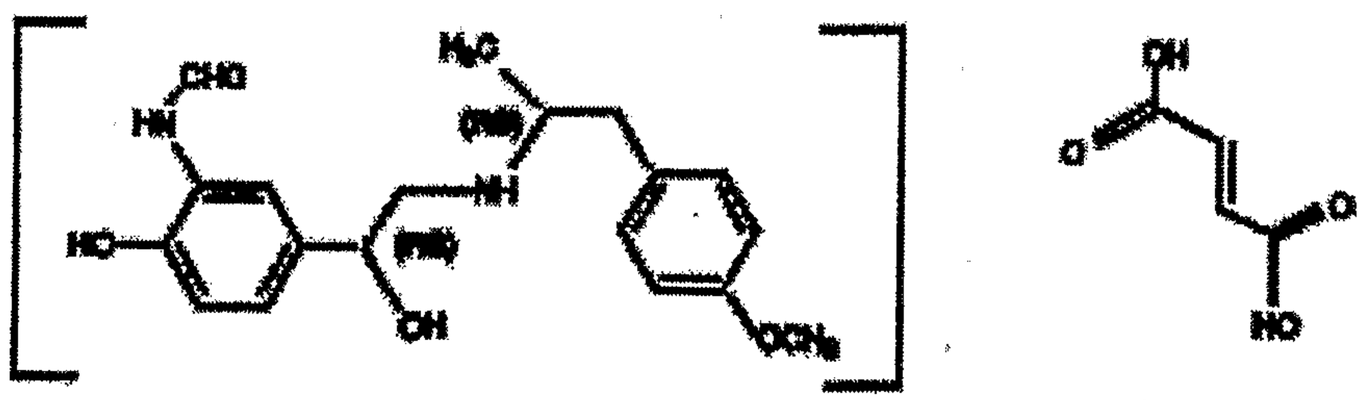
wherein R1, R2, and R3 have the same significance as in the formula(VII) and more specifically the formula (VIII) includes the following three formulae;



in the above formulae, R3 and R4 have the same significance as in formula (VII).

Preferably, formoterol is N-[2-hydroxy-5-(1-hydroxy-2-((2-(4-methoxyphenyl)-1-methylethyl) amino)-ethyl)phenyl] formamide, or a pharmaceutically acceptable salt thereof or a solvate of formoterol or said salt. Preferably, it is in the form of its fumarate salt. Formoterol is a bronchodilator used in the treatment of inflammatory or obstructive airways diseases.

A particularly preferred formoterol is formoterol fumarate with the following chemical formula:



Pharmaceutically acceptable salts of formoterol include, e.g., salts of inorganic acids such as hydrochloric, hydrobromic, sulfuric and phosphoric acids, and organic acids such as fumaric, maleic, acetic, lactic, citric, tartaric, ascorbic, succinic, glutaric,

gluconic, tricarballylic, oleic, benzoic, p-methoxybenzoic, salicylic, o- and p-hydroxybenzoic, p-chlorobenzoic, methanesulfonic, p-toluenesulfonic and 3-hydroxy-2-naphthalene carboxylic acids.

Formoterol may be in any isomeric form or mixture of isomeric forms, e.g. a pure enantiomer, a mixture of enantiomers, a racemate or a mixture thereof. It may be in the form of a solvate, e.g. a hydrate, thereof, e.g. as described in U.S. Patent Nos. 3,994,974 or 5,684,199, and may be present in a particular crystalline form, e.g. as described in WO95/05805. These patens and patent applications are herein incorporated by reference in their entirety. Preferably, formoterol is formoterol fumarate, especially in the form of the dihydrate.

A suitable daily dose of formoterol, or salt or solvate thereof, particularly as formoterol fumarate dihydrate, for inhalation may be from 1 to 72 μg, e.g. from 1 to 60 μg, generally from 3 to 50 μg, preferably from 6 to 48 μg, for instance from 6 to 24 μg. The precise dose used will depend on the condition to be treated, the patient and the efficiency of the inhalation device. The unit doses of formoterol and its frequency of administration may be chosen accordingly. A suitable unit dose of formoterol, particularly as formoterol fumarate dihydrate, may be from 1 to 72 μg, e.g. from 1 to 60 μg, generally from 3 to 48 μg, preferably from 6 to 36 μg, especially from 12 to 24 μg. These unit doses may suitably be administered once or twice daily in accordance with the suitable daily dose mentioned hereinbefore. For on demand usage, unit doses of 6 μg to 12 μg of formoterol are preferred.

The formoterol compounds are prepared and isolated by the methods described in U.S. Patent Nos. 3,994,974 and 5,684,199 (herein incorporated by reference in their entirety). Formoterol fumarate is available commercially. It is marketed in capsule form containing a dry powder formulation (Foradil®), containing 12 μg of formoterol fumarate and 25 mg of lactose, to be used for oral inhalation only with the Aerolizer® Inhaler (Schering Corp., Kenilworth, NJ).

Foradil® Aerolizer® is indicated for long-term, twice daily (morning and evening) administration in the treatment of asthma and the prevention of bronchospasm in adults and children (5 years or older) with reversible obstructive airways disease, including patients with symptoms of nocturnal asthma, who require regular treatment with inhaled short-acting, β2-agonist. It is also indicated for acute prevention of exercise-induced bronchospasm in adults and children (12 years or

older). It is also indicated to treat asthma concomitantly with short-acting β2-agonist, inhaled or systemic corticosteroids, and theophylline therapy. It is also indicated to for the long-term, twice daily (morning and evening) administration in the maintenance treatment of bronchoconstriction inpatients with COPD including chronic bronchitis and emphysema.

The first and second active agents are used to treat respiratory and lung diseases, and any of the additional agents listed below, may be administered per se or in the form of pharmaceutically acceptable salts, as discussed above, all being referred to as "active compounds or agents". The first and second active agents may also be administered in combination with one another, in the form of separate, or jointly in, pharmaceutically or veterinarily acceptable formulation(s). The active compounds or their salts may be administered either systemically or topically, as discussed below.

The present invention also provides for methods for treating asthma, COPD, or other respiratory diseases comprising administering the composition to a subject in need of such treatment. The method is for prophylactic or therapeutic purposes. The method comprises an in vivo method. The method is effective for treating a plurality of diseases, whatever their cause, including steroid administration, abnormalities in adenosine or adenosine receptor metabolism or synthesis, or other causes. The method comprises treating respiratory and lung diseases, whether by reducing adenosine or adenosine receptor levels, reducing hypersensitivity to adenosine, or other mechanisms, particularly in the lung, liver, heart and brain, or any organ that is need of such treatment. Other respiratory diseases referred to herein include cystic fibrosis (CF), dyspnea, emphysema, wheezing, pulmonary hypertension, pulmonary fibrosis, lung cancer, hyper-responsive airways, increased adenosine or adenosine receptor levels, particularly those associated with infectious diseases, pulmonary bronchoconstriction, lung inflammation, lung allergies, surfactant depletion, chronic bronchitis, bronchoconstriction, difficult breathing, impeded and obstructed lung airways, adenosine test for cardiac function, pulmonary vasoconstriction, impeded respiration, Acute Respiratory Distress Syndrome (ARDS), administration of certain drugs, such as adenosine and adenosine level increasing drugs, and other drugs for, e.g. treating SupraVentricular Tachycardia (SVT), and the administration of adenosine stress tests, infantile Respiratory Distress Syndrome (infantile RDS), pain, allergic rhinitis, decreased lung surfactant, severe acute respiratory syndrome (SARS),

among others.

In one embodiment, the invention is a method for the prophylaxis or treatment of asthma comprising administering the composition to a subject in need of such treatment an amount of the composition sufficient for the prophylaxis or treatment of asthma in the subject.

In one embodiment, the invention is a method for the prophylaxis or treatment of COPD comprising administering the composition to a subject in need of such treatment an amount of the composition sufficient for the prophylaxis or treatment of COPD in the subject.

In one embodiment, the invention is a method for the prophylaxis or treatment of bronchoconstriction, lung inflammation or lung allergy comprising administering the composition to a subject in need of such treatment an amount of the composition sufficient for the prophylaxis or treatment of bronchoconstriction, lung inflammation or lung allergy in the subject.

In one embodiment, the invention is a method for the reducing or depleting adenosine in a subject's tissue comprising administering the composition to a subject in need of such treatment an amount of the composition sufficient to reduce or deplete adenosine in the subject's tissue.

The present invention also provides for a use of the first active agent and the second active agent in the manufacture of a medicament for the treatment of asthma, COPD, or other respiratory diseases, including lung cancer. The medicament comprises the composition described throughout this disclosure.

The daily dosage of the first active agent and the second active agent to be administered to a subject will vary with the overall treatment programmed, the first active agent and the second active agent to be employed, the type of formulation, the route of administration and the state of the patient. Examples 11 to 18 show aerosolized preparations in accordance with the invention for delivery with a device for respiratory or nasal administration, or administration by inhalation. For intrapulmonary administration, liquid preparations are preferred. In the case of other bioactive agents, there exist FDA recommended amounts for supplementing a person's dietary intake with additional bioactive agents, such as in the case of vitamins and minerals. However, where employed for the treatment of specific conditions or for improving the immune response of a subject they may be utilized in

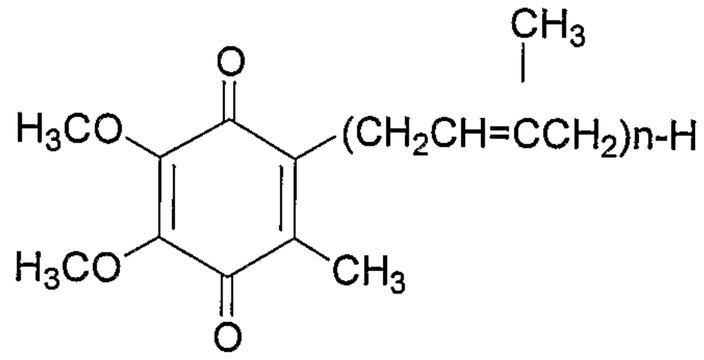
dosages hundreds and thousands of times higher. Mostly, the pharmacopeia's recommendations cover a very broad range of dosages, from which the medical artisan may draw guidance. Amounts for the exemplary agents described in this patent may be in the range of those currently being recommended for daily consumption, below or above those levels. The treatment may typically begin with a low dose of a bronchodilator in combination with a non-glucocorticoid steroid, or other bioactive agents as appropriate, and then a titration up of the dosage for each patient. Higher and smaller amounts, including initial amounts, however, may be administered within the confines of this invention as well.

Preferable ranges for the first and second active agents, or any other therapeutic agent, employed here will vary depending on the route of administration and type of formulation employed, as an artisan will appreciate and manufacture in accordance with known procedures and components. The active compounds may be administered as one dose (once a day) or in several doses (several times a day). The compositions and method of preventing and treating respiratory, cardiac, and cardiovascular diseases may be used to treat adults and infants, as well as non-human animals afflicted with the described conditions. Although the present invention is concerned primarily with the treatment of human subjects, it may also be employed, for veterinary purposes in the treatment of non-human mammalian subjects, such as dogs and cats as well as for large domestic and wild animals. The terms "high" and "low" levels of "adenosine" and "adenosine receptors" as well as "adenosine depletion" are intended to encompass both, conditions where adenosine levels are higher than, or lower (even depleted) when compared to previous adenosine levels in the same subject, and conditions where adenosine levels are within the normal range but, because of some other condition or alteration in that patient, a therapeutic benefit would be achieved in the patient by decreasing or increasing adenosine or adenosine receptor levels or hypersensitivity. Thus, this treatment helps regulate (titrate) the patient in a custom tailored manner. Whereas the administration of the first active agent may decrease or even deplete adenosine levels in a subject having either normal or high levels prior to treatment, the further administration of the second active agent will improve the subject's respiration in a short period of time. The further addition of other therapeutic agents will help titrate undesirably low levels of adenosine, which may be observed upon the administration of the present treatment, particularly until an

optimal titration of the appropriate dosages is attained.

Other therapeutic agents that may be incorporated into the present composition are one or more of a variety of therapeutic agents that are administered to humans and animals.

The composition can further comprise, in addition to the first and second active agents, a ubiquinone and/or folinic acid. A ubiquinone is a compound represented by the formula:



(II)

or pharmaceutically acceptable salt thereof.

Preferably, the ubiquinone is a compound according to the chemical formula given above, wherein n=1-10 (Coenzymes Q_{1-10}), more preferably n=6-10, (Coenzymes Q_{6-10}) and most preferably n=10 (Coenzyme Q_{10}). The ubiquinone is administered in a therapeutic amount for treating the targeted disease or condition, and the dosage will vary depending upon the condition of the subject, other agents being administered, the type of formulation employed, and the route of administration. The ubiquinone is preferably administered in a total amount per day of about 0.1, about 1, about 3, about 5, about 10, about 15, about 30 to about 50, about 100, about 150, about 300, about 600, about 900, about 1200 mg/kg body weight. More preferred the total amount per day is about 1 to about 150 mg/kg, about 30 to about 100 mg/kg, and

most preferred about 5 to about 50 mg/kg. Ubiquinone is a naturally occurring substance and is available commercially.

The active agents of this invention are provided within broad amounts of the composition. For example, the active agents may be contained in the composition in amounts of about 0.001%, about 1%, about 2%, about 5%, about 10%, about 20%, about 40%, about 90%, about 98%, about 99.999% of the composition. The amount of each active agent may be adjusted when, and if, additional agents with overlapping activities are included as discussed in this patent. The dosage of the active compounds, however, may vary depending on age, weight, and condition of the subject. Treatment may be initiated with a small dosage, e.g. less than the optimal dose, of the first active agent of the invention. This may be similarly done with the second active agent, until a desirable level is attained. Or vice versa, for example in the case of multivitamins and/or minerals, the subject may be stabilized at a desired level of these products and then administered the first active compound. The dose may be increased until a desired and/or optimal effect under the circumstances is reached. In general, the active agent is preferably administered at a concentration that will afford effective results without causing unduly harmful or deleterious side effects, and may be administered either as a single unit dose, or if desired in convenient subunits administered at suitable times throughout the day. The second therapeutic or diagnostic agent(s) is (are) administered in amounts which are known in the art to be effective for the intended application. In cases where the second agent has an overlapping activity with the principal agent, the dose of one of the other or of both agents may be adjusted to attain a desirable effect without exceeding a dose range that avoids untoward side effects. Thus, for example, when other analgesic and anti-inflammatory agents are added to the composition, they may be added in amounts known in the art for their intended application or in doses somewhat lower that when administered by themselves.

Pharmaceutically acceptable salts should be pharmacologically and pharmaceutically or veterinarily acceptable, and may be prepared as alkaline metal or alkaline earth salts, such as sodium, potassium or calcium salts. Organic salts and esters are also suitable for use with this invention. The active compounds are preferably administered to the subject as a pharmaceutical or veterinary composition, which includes systemic and topical formulations. Among these, preferred are

formulations suitable for inhalation, or for respirable, buccal, oral, rectal, vaginal, nasal, intrapulmonary, ophthalmic, optical, intracavitary, intratraccheal, intraorgan, topical (including buccal, sublingual, dermal and intraocular), parenteral (including subcutaneous, intradermal, intramuscular, intravenous and intraarticular) and transdermal administration, among others.

The present invention also provides for a kit comprising the composition and a delivery device. The compositions may conveniently be presented in single or multiple unit dosage forms as well as in bulk, and may be prepared by any of the methods which are well known in the art of pharmacy. The composition, found in the kit, whether already formulated together or where the first and second active agents are separately provided along with other ingredients, and instructions for its formulation and administration regime. The kit may also contain other agents, such as those described in this patent and, for example, when for parenteral administration, they may be provided with a carrier in a separate container, where the carrier may be sterile. The present composition may also be provided in lyophilized form, and in a separate container, which may be sterile, for addition of a liquid carrier prior to administration. See, e.g. U.S. Patent No. 4,956,355; UK Patent No. 2,240,472; EPO Patent Application Serial No. 429,187; PCT Patent Publication WO 91/04030; Mortensen, S. A., et al., *Int. J. Tiss. Reac.* XII(3): 155-162 (1990); Greenberg, S. et al., *J. Clin. Pharm.* 30: 596-608 (1990); Folkers, K., et al., *Proc. Natl. Acad. Sci. USA* 87: 8931-8934 (1990), the relevant preparatory and compound portions of which are incorporated by reference above.

The present composition is provided in a variety of systemic and topical formulations. The systemic or topical formulations of the invention are selected from the group consisting of oral, intrabuccal, intrapulmonary, rectal, intrauterine, intradermal, topical, dermal, parenteral, intratumor, intracranial, intrapulmonary, buccal, sublingual, nasal, subcutaneous, intravascular, intrathecal, inhalable, respirable, intraarticular, intracavitary, implantable, transdermal, iontophoretic, intraocular, ophthalmic, vaginal, optical, intravenous, intramuscular, intraglandular, intraorgan, intralymphatic, slow release and enteric coating formulations. The actual preparation and compounding of these different formulations is known in the art and need not be detailed here. The composition may be administered once or several times a day.

Formulations suitable for respiratory, nasal, intrapulmonary, and inhalation administration are preferred, as are topical, oral and parenteral formulations. All methods of preparation include the step of bringing the active compound into association with a carrier which constitutes one or more accessory ingredients. In general, the formulations are prepared by uniformly and intimately bringing the active compound into association with a liquid carrier, a finely divided solid carrier, or both, and then, if necessary, shaping the product into desired formulations.

Compositions suitable for oral administration may be presented in discrete units, such as capsules, cachets, lozenges, or tablets, each containing a predetermined amount of the active compound; as a powder or granules; as a solution or a suspension in an aqueous or non-aqueous liquid; or as an oil-in-water or water-in-oil emulsion.

Compositions suitable for parenteral administration comprise sterile aqueous and non-aqueous injection solutions of the active compound, which preparations are preferably isotonic with the blood of the intended recipient. These preparations may contain anti-oxidants, buffers, bacteriostats and solutes which render the compositions isotonic with the blood of the intended recipient. Aqueous and non-aqueous sterile suspensions may include suspending agents and thickening agents. The compositions may be presented in unit-dose or multi-dose containers, for example sealed ampoules and vials, and may be stored in a freeze-dried or lyophilized condition requiring only the addition of the sterile liquid carrier, for example, saline or water-for-injection immediately prior to use.

Nasal and instillable formulations comprise purified aqueous solutions of the active compound with preservative agents and isotonic agents. Such formulations are preferably adjusted to a pH and isotonic state compatible with the nasal mucous membranes.

Formulations for rectal or vaginal administration may be presented as a suppository with a suitable carrier such as cocoa butter, or hydrogenated fats or hydrogenated fatty carboxylic acids.

Ophthalmic formulations are prepared by a similar method to the nasal spray, except that the pH and isotonic factors are preferably adjusted to match that of the eye. Otical formulations are generally prepared in viscous carriers, such as oils and the like, as is known in the art, so that they may be easily administered into the ear without spilling.

Compositions suitable for topical application to the skin preferably take the form of an ointment, cream, lotion, paste, gel, spray, aerosol, or oil. Carriers which may be used include Vaseline, lanolin, polyethylene glycols, alcohols, transdermal enhancers, and combinations of two or more thereof. Compositions suitable for transdermal administration may be presented as discrete patches adapted to remain in intimate contact with the epidermis of the recipient for a prolonged period of time.

The first and second active agents disclosed herein may be administered into the respiratory system either by inhalation, respiration, nasal administration or intrapulmonary instillation (into the lungs) of a subject by any suitable means, and are preferably administered by generating an aerosol or spray comprised of powdered or liquid nasal, intrapulmonary, respirable or inhalable particles. The respirable or inhalable particles comprising the active compound are inhaled by the subject, i.e, by inhalation or by nasal administration or by instillation into the respiratory tract or the lung itself. The formulation may comprise respirable or inhalable liquid or solid particles of the active compound that, in accordance with the present invention, include respirable or inhalable particles of a size sufficiently small to pass through the mouth and larynx upon inhalation and continue into the bronchi and alveoli of the lungs. In general, particles ranging from about 0.05, about 0.1, about 0.5, about 1, about 2 to about 4, about 6, about 8, about 10 microns in diameter. More particularly, about 0.5 to less than about 5 μm in diameter, are respirable or inhalable. Particles of non-respirable size which are included in an aerosol or spray tend to deposit in the throat and be swallowed. The quantity of non-respirable particles in the aerosol is, thus, preferably minimized. For nasal administration or intrapulmonary instillation, a particle size in the range of about 8, about 10, about 20, about 25 to about 35, about 50, about 100, about 150, about 250, about 500 μm (diameter) is preferred to ensure retention in the nasal cavity or for instillation and direct deposition into the lung. Liquid formulations may be squirted into the respiratory tract (nose) and the lung, particularly when administered to newborns and infants.

Liquid pharmaceutical compositions of active compound for producing an aerosol may be prepared by combining the active compound with a stable vehicle, such as sterile pyrogen free water. Solid particulate compositions containing respirable dry particles of micronized active compound may be prepared by grinding dry active compound with a mortar and pestle, and then passing the micronized

composition through a 400 mesh screen to break up or separate out large agglomerates. A solid particulate composition comprised of the active compound may optionally contain a dispersant that serves to facilitate the formation of an aerosol. A suitable dispersant is lactose, which may be blended with the active compound in any suitable ratio, e.g., a 1 to 1 ratio by weight. The U.S. Patent Application Ser. Nos. 10/462,901 and 10/462,927 disclose a stable dry powder formulation of DHEA in a nebulizable form and a stable dry powder formulation of dihydrate crystal form of DHEA-S, respectively (these patent applications are herein incorporated by reference in their entirety).

Aerosols of liquid particles comprising the active compound may be produced by any suitable means, such as with a nebulizer. See, e.g. U.S. Patent No. 4,501,729 (the disclosure of which is incorporated by reference). Nebulizers are commercially available devices which transform solutions or suspensions of the active ingredient into a therapeutic aerosol mist either by means of acceleration of a compressed gas, typically air or oxygen, through a narrow venturi orifice or by means of ultrasonic agitation. Suitable compositions for use in nebulizer consist of the active ingredient in liquid carrier, the active ingredient comprising up to 40% w/w composition, but preferably less than 20% w/w carrier being typically water or a dilute aqueous alcoholic solution, preferably made isotonic with body fluids by the addition of, for example sodium chloride. Optional additives include preservatives if the composition is not prepared sterile, for example, methyl hydroxybenzoate, anti-oxidants, flavoring agents, volatile oils, buffering agents and surfactants. Aerosols of solid particles comprising the active compound may likewise be produced with any sold particulate medicament aerosol generator. Aerosol generators for administering solid particulate medicaments to a subject product particles which are respirable, as explained above, and generate a volume of aerosol containing a predetermined metered dose of a medicament at a rate suitable for human administration. Examples of such aerosol generators include metered dose inhalers and insufflators.

The composition may be delivered with any delivery device that generates liquid or solid particulate aerosols, such as aerosol or spray generators. These devices produce respirable particles, as explained above, and generate a volume of aerosol or spray containing a predetermined metered dose of a medicament at a rate suitable for human or animal administration. One illustrative type of solid particulate aerosol or

spray generator is an insufflator, which are suitable for administration of finely comminuted powders. In the insufflator, the powder, e.g. a metered dose of the composition effective to carry out the treatments described herein, is contained in a capsule or a cartridge. These capsules or cartridges are typically made of gelatin, foil or plastic, and may be pierced or opened in situ, and the powder delivered by air drawn through the device upon inhalation or by means of a manually-operated pump. The composition employed in the insufflator may consist either solely of the first and second agents or of a powder blend comprising the first and second agents, typically comprising from 0.01 to 100 % w/w of the composition. The composition generally contains the first and second agents in an amount of about 0.01% w/w, about 1% w/w/, about 5% w/w, to about 20%, w/w, about 40% w/w, about 99.99% w/w. Other ingredients, and other amounts of the agent, however, are also suitable within the confines of this invention.

In one embodiment, the composition is delivered by a nebulizer. This means is especially useful for patients or subjects who are unable to inhale or respire the composition under their own efforts. In serious cases, the patients or subjects are kept alive through artificial respirator. The nebulizer can use any pharmaceutically or veterinarily acceptable carrier, such as a weak saline solution. The nebulizer is the means by which the powder pharmaceutical composition is delivered to the target of the patients or subjects in the airways.

The composition is also provided in various forms that are tailored for different methods of administration and routes of delivery. In one embodiment, the composition comprises a respirable formulation, such as an aerosol or spray. The composition of the invention is provided in bulk, and in unit form, as well as in the form of an implant, a capsule, blister or cartridge, which may be openable or piercable as is known in the art. A kit is also provided, that comprises a delivery device, and in separate containers, the composition of the invention, and optionally other excipient and therapeutic agents, and instructions for the use of the kit components.

In one embodiment, the composition is delivered using suspension metered dose inhalation (MDI) formulation. Such a MDI formulation can be delivered using a delivery device using a propellant such as hydrofluroalkane (HFA). Preferably, the HFA propellants contain 100 parts per million (PPM) or less of water.

In one embodiment, the delivery device comprises a dry powder inhalator (DPI) that delivers single or multiple doses of the composition. The single dose inhalator may be provided as a disposable kit which is sterilely preloaded with enough formulation for one application. The inhalator may be provided as a pressurized inhalator, and the formulation in a piercable or openable capsule or cartridge. The kit may optionally also comprise in a separate container an agent such as other therapeutic compounds, excipients, surfactants (intended as therapeutic agents as well as formulation ingredients), antioxidants, flavoring and coloring agents, fillers, volatile oils, buffering agents, dispersants, surfactants, antioxidants, flavoring agents, bulking agents, propellants and preservatives, among other suitable additives for the different formulations.

Having now generally described this invention, the same will be better understood by reference to certain specific examples, which are included herein for purposes of illustration only and are not intended to be limiting of the invention or any embodiment thereof, unless so specified.

EXAMPLES

Examples 1 and 2: In vivo Effects of Folinic Acid and DHEA on Adenosine Levels

Young adult male Fischer 344 rats (120 grams) were administered dehydroepiandrosterone (DHEA) (300 mg/kg) or methyltestosterone (40 mg/kg) in carboxymethylcellulose by gavage once daily for fourteen days. Folinic acid (50 mg/kg) was administered intraperitoneally once daily for fourteen days. On the fifteenth day, the animals were sacrificed by microwave pulse (1.33 kilowatts, 2450 megahertz, 6.5 seconds (s)) to the cranium, which instantly denatures all brain protein and prevents further metabolism of adenosine. Hearts were removed from animals and flash frozen in liquid nitrogen with 10 s of death. Liver and lungs were removed en bloc and flash frozen with 30 s of death. Brain tissue was subsequently dissected. Tissue adenosine was extracted, derivatized to 1, N6-ethenoadenosine and analyzed by high performance liquid chromatography (HPLC) using spectrofluorometric detection according to the method of Clark and Dar (J. of Neuroscience Methods 25:243 (1988)). Results of these experiments are summarized in Table 1 below. Results are expressed as the mean ±SEM, with κ $p<0.05$ compared to control group

and ψ $p<0.05$ compared to DHEA or methyltestosterone-treated groups.

Table 1: In vivo Effect of DHEA, δ -1-methyltestosterone and Folinic Acid on Adenosine Levels in various Rat Tissues

Treatment	Intracellular adenosine (nmols)/mg protein			
	Heart	Liver	Lung	Brain
Control	10.6 ± 0.6 (n=12) κ	14.5 ± 1.0 (n=12) κ	3.1 ± 0.2 (n=6) κ	0.5 ± 0.04 (n=12) κ
DHEA (300 mg/kg)	6.7 ± 0.5 (n=12) κ	16.4 ± 1.4 (n=12) κ	2.3 ± 0.3 (n=6) κ	0.19 ± 0.01 (n=12) κ
Methyltestosterone (40 mg/kg)	8.3 ± 1.0 (n=6) κ	16.5 ± 0.9 (n=6) κ	N.D.	0.42 ± 0.06 (n=6) κ
Methyltestosterone (120 mg/kg)	6.0 ± 0.4 (n=6) κ	5.1 ± 0.5 (n=6) κ	N.D.	0.32 ± 0.03 (n=6) κ
Folinic Acid (50 mg/kg)	12.4 ± 2.1 (n=5) κ	16.4 ±2.4 (n=5) κ	N.D.	0.72 ± 0.09 (n=5) κ
DHEA (300 mg/kg) + Folinic Acid (50 mg/kg)	11.1 ± 0.6 (n=5) ψ	18.8 ± 1.5 (n=5) ψ	N.D.	0.55 ± 0.09 (n=5) ψ
Methyltestosterone (120 mg/kg) + Folinic Acid (50 mg/kg)	9.1 ± 0.4 (n=6) ψ	N.D.	N.D.	0.60 ± 0.06 (n=6) ψ

N.D.=Not determined

The results of these experiments indicate that rats administered DHEA or methyltestosterone daily for two weeks showed multi-organ depletion of adenosine. Depletion was dramatic in brain (60% depletion for DHEA, 34% for high dose methyltestosterone) and heart (37% depletion for DHEA, 22% depletion for high dose methyltestosterone). Coadministration of folinic acid completely abrogated steroid-mediated adenosine depletion. Folinic acid administered alone induce increase in adenosine levels for all organs studied.

Example 3: Airjet Milling of Anhydrous DHEA-S and Determination of Respirable Dose

DHEA-S is evaluated as an asthma therapy. The solid-state stability of sodium dehydroepiandrostenone sulfate (NaDHEA-S) has been studied for both bulk and milled material (Nakagawa, H., Yoshiteru, T., and Fujimoto, Y. (1981) *Chem. Pharm. Bull.* 29(5) 1466-1469; Nakagawa, H., Yoshiteru, T., and Sugimoto, I. (1982) *Chem. Pharm. Bull.* 30(1) 242-248). DHEA-S is most stable and crystalline as the dihydrate form. The DHEA-S anhydrous form has low crystallinity and is very hygroscopic. The DHEA-S anhydrous form is stable as long as it picks up no water on storage. Keeping a partially crystalline material free of moisture requires specialized manufacturing and packing technology. For a robust product, minimizing sensitivity to moisture is essential during the development process.

(1) Micronization of DHEA-S

Anhydrous DHEA-S was micronized using a jet milling (Jet-O-Mizer Series #00, 100-120 PSI nitrogen). Approximately 1 g sample was passed through the jet mill, once, and approximately 2 g sample were passed through the jet mill twice. The particles from each milling run were suspended in hexane, in which DHEA-S was insoluble and Spa85 surfactant added to prevent agglomeration. The resulting solution was sonicated for 3 minutes and appeared fully dispersed. The dispersed solutions were tested on a Malvern Mastersizer X with a small volume sampler (SVS) attachment. One sample of dispersed material was tested 5 times. The median particle size or D (v, 0.5) of unmilled material was 52.56 μm and the %RSD (relative standard deviation) was 7.61 for the 5 values. The D (v, 0.5) for a single pass through

the jet mill was 3.90 μm and the %RSD was 1.27, and the D (v, 0.5) from a double pass through the jet mill 3.25 μm and the %RSD was 3.10. This demonstrates that DHEA-S can be jet milled to particles of size suitable for inhalation.

(2) HPLC Analysis

Two vials (A; single-pass; 150 mg) and (B double-pass; 600 mg) of the micronized drug were available for determining drug degradation during jet milling micronization. Weighed aliquots of DHEA-S from vials A and B were compared to a standard solution of unmilled DHEA-S (10 mg/ml) in an acetonitrile-water solution (1:1). The chromatographic peak area for the HPLC assay of the unmilled drug standard solution (10 mg/ml) gave a value of 23,427. Weighed aliquots of micronized DHEA-S form vials A and B, (5 mg/ml) was prepared in an acetonitrile-water solution (1:1). The chromatographic peak areas for vials A and B were 11,979 and 11,677, respectively. Clearly, there was no detectable degradation of the drug during the jet milling micronization process.

(3) Emitted Dose Studies

DHEA-S powder was collected in Nephele tubes and assayed by HPLC. Triplicate experiments were performed at each airflow rate for each of the three dry powder inhalers tested (Rotahaler, Diskhaler and IDL's DPI devices). A Nephele tube was fitted at one end with a glass filter (Gelman Sciences, Type A/E, 25 μm), which in turn was connected to the airflow line to collect the emitted dose of the drug from the respective dry powder inhaler being tested. A silicone adapter, with an opening to receive the mouthpiece of the respective dry powder inhaler being tested at the other end of the Nephele tube was secured. A desired airflow, of 30, 60, or 90 L/min, was achieved through the Nephele tube. Each dry powder inhaler's mouthpiece was inserted then into the silicone rubber adapter, and the airflow was continued for about four s after which the tube was removed and an end-cap screwed onto the end of each tube. The end-cap of the tube not containing the filter was removed and 10 ml of an HPLC grade water- acetonitrile solution (1:1) added to the tube, the end-cap reattached, and the tube shaken for 1-2 minutes. The end-cap then was removed from the tube and the solution was transferred to a 10 ml plastic syringe fitted with a filter (Cameo 13N Syringe Filter, Nylon, 0.22μm). An aliquot of the solution was directly filtered into an HPLC vial for later drug assay via HPLC. The

emitted dose experiments were performed with micronized DHEA-S (about 12.5 or 25 mg) being placed in either a gelatin capsule (Rotahaler) or a Ventodisk blister (Diskhaler and single-dose DPI (IDL)). When the micronized DHEA-S (only vial B used), was weighed for placement into the gelatin capsule or blister, there appeared to be a few aggregates of the micronized powder. The results of the emitted dose tests conducted at an airflow rate of 30, 60 and 87.8 L/min are displayed in Tables 2. Table 2 summarizes the results for the Rotahaler experiments at 3 different flow rates, for the Diskhaler experiments at 3 different flow rates, and of the multi-dose experiments at 3 different flow rates.

Table 2. Emitted Dose Comparison of Three Different Dry Powder Inhaler Devices

Inhaler Device	Airflow Rate (L/min)	Emitted Dose (%)
Rotahaler	87.8	73.2, 67.1, 68.7
Average		69.7
Rotahaler (2nd study)	87.8	16.0, 24.5, 53.9
Average		31.5
Diskhaler	87.8	65.7, 41.6, 46.5
Average		51.3
Diskhaler (2nd study)	87.8	57.9, 59.9, 59.5
Average		59.1
IDL Multi-Dose	87.8	71.3, 79.0, 67.4
Average		72.6
IDL Multi-Dose (2nd study)	87.8	85.7, 84.6, 84.0
Average		84.8
Rotahaler	60	58.1, 68.2, 45.7
Average		57.3
Diskhaler	60	63.4, 38.9, 58.0
Average		68.2
IDL Multi-Dose	60	78.8, 83.7, 89.6
Average		84.0
Rotahaler	30	34.5, 21.2, 48.5
Average		34.7
Diskhaler	30	53.8, 53.4, 68.7
		58.6
IDL Multi-Dose	30	78.9, 88.2, 89.2
Average		85.4

(4) Respirable Dose Studies

The respirable dose (respirable fraction) studies were performed using a standard sampler cascade impactor (Andersen), consisting of an inlet cone (an impactor pre-separator was substituted here), 9 stages, 8 collection plates, and a backup filter within 8 aluminum stages held together by 3 spring clamps and gasket O-ring seals, where each impactor stage contains multiple precision drilled orifices. When air is drawn through the sampler, multiple jets of air in each stage direct any airborne particles toward the surface of the collection plate for that stage. The size of the jets is constant for each stage, but is smaller in each succeeding stage. Whether a particle is impacted on any given stage depends upon its aerodynamic diameter. The range of particle sizes collected on each stage depends upon on the jet velocity of the stage, and the cut-off point of the previous stage. Any particle not collected on the first stage follows the air stream around the edge of the plate to the next stage, where it is either impacted or passed on to the succeeding stage, and so on, until the velocity of the jet is sufficient for impaction. To prevent particle bounce during the cascade impactor test, the individual impactor plates were coated with a hexane-grease (high vacuum) solution (100:1 ratio). As noted above, the particle size cut-off points on the impactor plates changed at different airflow rates. For example, Stage 2 corresponds to a cut-off value greater than 6.2 μm particles at 60 L/min, and greater than 5.8 μm particles at 30 L/min, and stage 3 had a particle size cut-off value at 90 L/min greater than 5.6 μm. Thus, similar cut-off particle values are preferentially employed at comparable airflow rates, i.e. ranging from 5.6 to 6.2 μm. The set-up recommended by the United States Phamacopeia for testing dry powder inhalers consists of a mouthpiece adapter (silicone in this case) attached to a glass throat (modified 50 ml round-bottom flask) and a glass distal pharynx (induction port) leading top the pre-separator and Andersen sampler. The pre-separator sample includes washings from the mouthpiece adaptor, glass throat, distal glass pharynx and pre-separator. 5 ml acetonitrile:water (1:1 ratio) solvent was placed in the pre-separator before performing the cascade impactor experiment, that were performed in duplicate with 3 different dry powder inhaler devices and at 3 airflow rates, 30, 60 and 90 L/min. The drug collected on the cascade impactor plates were assayed by the HPLC, and a drug mass balance was performed for each Diskhaler and multi-dose cascade impactor experiment consisting of determining the amount of drug left in the blister, the

amount of drug remaining in the device (Diskhaler only), the non-respirable amount of the dose retained on the silicone rubber mouth piece adaptor, glass throat, glass distal pharynx and pre-separator, all combined into one sample, and the respirable dose, i.e. Stage 2 through filter impactor plates for airflow rates of 30 and 60 L/min and Stages 1 through filter impactor plates for 90 L/min experiments.

Table 3. Cascade Impactor Experiments (90L/min)

Inhaler Device	Preseparator (%)	Blister (%)	Respirable Dose (%)	Device (%)	Mass Balance (%)
Diskhaler	72.7	6.6	2.9	22.1	104.3
Diskhaler	60.2	10.1	2.4	13.3	86.0
Multi-dose	65.8	3.9	3.8	26.5 *a	100.0
Multi-dose	73.3	3.8	3.6	19.3 *a	100.0
Multi-dose *b	78.7	2.8	4.6	13.9 *a	100.0
Multi-dose *c	55.9	5.0	1.2	37.9 *a	100.0

*a: Multi-dose device was not washed; as solvents would attack SLA components. Multi-dose device retention percentage is obtained by difference.

*b: oven dried drug for 80 minutes

*c: oven dried drug for 20 hours

Based on the results of the emitted dose and cascade impactor experiments, the low respirable dose values achieved in the cascade impactor experiments were due to agglomerated drug particles, which could not be separated, even at the highest airflow rate tested. Agglomeration of the drug particles is a consequence of static charge build up during the mechanical milling process used for particles size reduction and that this situation is further compounded by subsequent moisture

absorption of the particles. A micronization method that produces less static charge or a less hygroscopic, fully hydrated crystalline form of DHEA-S (i.e. dihydrate form) should provide a freer flowing powder with diminished potential for agglomeration.

Example 4: Spray Drying of Anhydrous DHEA-S and Determination of Respirable Dose

(1) Micronization of the Drug

1.5 g of anhydrous DHEA-S were dissolved to 100 ml of 50% ethanol:water to produce a 1.5 % solution. The solution was spray-dried with a B-191 Mini Spray-Drier (Buchi, Flawil, Switzerland) with an inlet temperature of 55°C, outlet temperature of 40°C, at 100% aspirator, at 10% pump, nitrogen flow at 40 mbar and spray flow at 600 units. The spray-dried product was suspended in hexane and Span85 surfactant added to reduce agglomeration. The dispersions were sonicated with cooling for 3-5 minutes for complete dispersion and the dispersed solutions tested on a Malvern Mastersizer X with a Small Volume Sampler (SVS) attachment. The two batches of spray dried material were found to have mean particle sizes of 5.07 ±0.70 μm and 6.66 ±0.91 μm. Visual examination by light microscope of the dispersions of each batch confirmed that spray drying produced small respirable size particles. The mean particle size was 2.4 μm and 2.0 μm for each batch, respectively. This demonstrates that DHEA-S can be spray dried to a particle size suitable for inhalation.

(2) Respirable Dose Studies

The cascade impactor experiments were conducted as described in Example 3. Four cascade impactor experiments were done, three with a IDL multi-dose device and one with a Diskhaler, all at 90 L/min. The results of the cascade impactor experiments are presented in Table 4 below. The spray-dried anhydrous material in these experiments produced a two-fold increase in the respirable dose compared to micronized anhydrous DHEA-S. It appears that spray drying obtained higher respirable doses as compared to jet-milling. However, the % respirable dose was still low. This was likely the result of moisture absorption of the anhydrous form.

Table 4: Cascade Impactor Results with Spray-Dried Drug Product

Device	Diskhaler	Multi-dose	Multi-dose	Multi-dose
Number of Blisters	3	3	4	4
Drug per Blister (mg)	38.2	36.7	49.4	50.7
Preseparator (%)	56.8	71.9	78.3	85.8
Device (%)	11.2	7.9	8.9	7.6
Blisters (%)	29.0	6.4	8.2	4.8
Respirable Dose (%)	5.6	7.8	5.3	2.6
Mass Balance Recovery (%)	102.7	94.0	103.3	98.1

Example 5: Air Jet Milling of DHEA-S Dihydrate (DHEA-S ·2H_2O) and Determination of Respirable Dose

(1) Recrystallization of DHEA-S dihydrate.

Anhydrous DHEA-S is dissolved in a boiling mixture of 90% ethanol/water. This solution is rapidly chilled in a dry ice/methanol bath to recrystallize the DHEA-S. The crystals are filtered, washed twice with cold ethanol, than dried in a vacuum desiccator at RT for 36 h. During the drying process, the material is periodically mixed with a spatula to break large agglomerates. After drying, the material is passed through a 500 μm sieve.

(2) Micronization and physiochecmical testing.

DHEA-S dihydrate is micronized with nitrogen gas in a jet mill at a venturi pressure of 40 PSI, a mill pressure of 80 PSI, feed setting of 25 and a product feed rate of about 120 to 175 g/hour. Surface area is determined using five point BET analyses are performed with nitrogen as the adsorbing gas (P/P_o = 0.05 to 0.30) using a Micromeritics TriStar surface area analyzer. Particle size distributions are measured by laser diffraction using a Micromeritics Saturn Digisizer where the particles are suspended in mineral oil with sodium dioctyl sodium sulfosuccinate as a dispersing agent. Drug substance water content is measured by Karl Fischer titration (Schott Titroline KF). Pure water is used as the standard and all relative standard deviations for triplicates are less than 1%. Powder is added directly to the titration media. The

physicochemical properties of DHEA-S·dihydrate before and after micronization are summarized in Table 5.

Table 5. Physicochemical properties of DHEA-S·dihydrate before and after micronization.

Property	Bulk	Micronized
Particle size ($D_{50\%}$)	31 microns	3.7 microns
Surface area (m^2/g)	Not measured	4.9
Water (% w/w)	8.5	8.4
Impurities	No significant peaks	No significant peaks

The only significant change measured is in the particle size. There is no significant loss of water or increase in impurities. The surface area of the micronized material is in agreement with an irregularly shaped particle having a median size of 3 to 4 microns. The micronization successfully reduces the particle size to a range suitable for inhalation with no measured changes in the solid-state chemistry.

(3) Aerosolization of DHEA-S·dihydrate.

The single-dose Acu-Breathe device is used for evaluating DHEA-S·dihydrate. Approximately 10 mg of neat DHEA-S·dihydrate powder is filled and sealed into foil blisters. These blisters are actuated into the Andersen 8-stage cascade impactor at flow rates ranging from 30 to 75 L/min with a glass twin-impinger throat. Stages 1-5 of the Andersen impactor are rinsed together to obtain an estimate of the fine particle fraction. Pooling the drug collected from multiple stages into one assay make the method much more sensitive. The results for this series of experiments is shown in Figure 1. At all flow rates, the dihydrate yields a higher fine particle fraction than the virtually anhydrous material. Since the dihydrate powder is aerosolized using the single-dose inhaler, it is very reasonable to conclude that its aerosol properties are significantly better than the virtually anhydrous material. Higher crystallinity and stable moisture content are the most likely factors contributing the dihydrate's superior aerosol properties. This unique feature of DHEA-S·dihydrate has not been reported in any previous literature. While the improvement in DHEA-S's aerosol performance with the dihydrate form is significant, neat drug substance may not be the

optimal formulation. Using a carrier with a larger particle size typically improves the aerosol properties of micronized drug substances.

Example 6: Anhydrous DHEA-S and DHEA-S Dihydrate Stability with and without Lactose

The initial purity (Time=0) was determined for anhydrous DHEA and for DHEA-S dihydrate by high pressure liquid chromatography (HPLC). Both forms of DHEA-S were then either blended with lactose at a ratio of 50:50, or used as a neat powder and placed in open glass vials, and held at 50°C for up to 4 weeks. These conditions were used to stress the formulation in order to predict its long-term stability results. Control vials containing only DHEA-S (anhydrous or dihydrate) were sealed and held 25°C for up to 4 weeks. Samples were taken and analyzed by HPLC also at 0, 1, 2, and 4 weeks to determine the amount of degradation, as determined by formation of DHEA. After one week, virtually anhydrous DHEA-S blended with lactose (50% w/w, nominally) stored at 50°C in sealed glass vials acquires a brown tinge that is darker for the lactose blend. This color change is accompanied by a significant change in the chromatogram as shown in Figure 1. The primary degradant is DHEA. Qualitatively from Figure 2, the amount of DHEA in the blend is higher than the other two samples. To quantitatively estimate the % DHEA in the samples, the area for the DHEA peak is divided by the total area for the DHEA-S and DHEA peaks (see Table 6). The higher rate of decomposition for the blend indicates a specific interaction between lactose and the virtually anhydrous DHEA-S. In parallel with the increase in DHEA, the brown color of the powders on accelerated storage increased over time. The materials on accelerated storage become more cohesive with time as evidenced by clumping during sample weighing for chemical analysis. Based on these results, it is not possible to formulate virtually anhydrous DHEA-S with lactose. This is a considerable disadvantage since lactose is the most commonly used inhalation excipient for dry powder formulations. Continuing with the virtually anhydrous form would mean limiting formulations to neat powder or undertaking more comprehensive safety studies to use a novel excipient.

Table 6: DHEA % formed from Anhydrous DHEA-S at 50°C

Formulation	Time (Weeks)	1	2	4
Control	2.774	2.694	2.370	2.666
DHEA-S. Alone		9.817	14.954	20.171
DHEA-S+Lactose (50:50)		24.085	30.026	38.201

In contrast to Figure 2, there is virtually no DHEA generated after storage for 1 week at 50 °C (see Figure 3). Furthermore, the materials show no change in color. The moisture content of DHEA-S·dihydrate remains virtually unchanged after one week at 50 °C. The water content after accelerated storage is 8.66% versus a starting value of 8.8%. The %DHEA measured during the course of this stability program is shown in Table 7.

Table 7: Percent DHEA formed from DHEA-S Dihydrate at 50°C

Formulation	Time (Weeks)	1	3	4
Control	0.213			0.218
DHEA-S alone		0.216	0.317	0.374
DHEA-S:Lactose (50:50)		0.191	0.222	0.323

By comparing Figures 1 and 2 and Tables 6 and 7, one can see that the dihydrate form of DHEA-S is the more stable form for progression into further studies. The superior compatibility of DHEA-S·dihydrate with lactose over that of the virtually anhydrous material has not been reported in the patent or research literature. The solubility of this substance is reported in the next section as a portion of the development work for a nebulizer solution.

Example 7: DHEA-S Dihydrate/Lacotse blends, Determination of Respirable Dose and Stability

(1) DHEA-S dihydrate/Lactose blend.

Equal weights of DHEA-S and inhalation grade lactose (Foremost Aero Flo 95) are mixed by hand then passed through a 500 μm screen to prepare a pre-blend. The pre-blend is then placed in a BelArt Micro-Mill with the remaining lactose to yield a 10% w/w blend of DHEA-S. The blender is wired to a variable voltage source to regulate the impeller speed. The blender voltage is cycled through 30%, 40%, 45% and 30% of full voltage for 1, 3, 1.5, and 1.5 minutes, respectively. The content uniformity of the blend was determined by HPLC analysis. Table 8 shows the result of content uniformity samples for this blend. The target value is 10% w/w DHEA-S. The blend content is satisfactory for proximity to the target value and content uniformity.

Table 8: Content uniformity for a blend of DHEA-S·dihydrate with lactose.

Sample	% DHEA-S, w/w
1	10.2
2	9.7
3	9.9
4	9.3
5	9.4
Mean	9.7
RSD	3.6%

(2) Aerosolization of DHEA-S·dihydrate/Lactose blend.

Approximately 25 mg of this powder is filled and sealed in foil blisters and aerosolized using the single-dose device at 60 L/min. Two blisters are used for each test and the results for fine particle fraction (material on stages 1-5) are shown in Table 9. The aerosol results for this preliminary powder blend are satisfactory for a respiratory drug delivery system. Higher fine particle fractions are possible with optimization of the powder blend and blister/device configuration. The entire particle size distribution of Test 2 is shown in Table 10. This median diameter for DHEA-S for this aerosol is ~2.5 μm. This diameter is smaller than the median diameter

measured for micronized DHEA-S·dihydrate by laser diffraction. Irregularly shaped particles can behave aerodynamically as smaller particles since their longest dimension tends to align with the air flow field. Therefore, it is common to see a difference between the two methods. Diffraction measurements are a quality control test for the input material while cascade impaction is a quality control test for the finished product.

Table 9: Fine particle fraction for lactose blend in two different experiments

Test	Total powder weight in two blisters (mg)	DHEA-S collected Stages 1-5 (mg)	Fine particle fraction, %
1	52.78	1.60	31
2	57.09	1.62	29

Table 10: Particle size distribution of aerosolized DHEA-S dihydrate/Lactose Blend

Size (μm)	6.18	9.98	3.23	2.27	1.44	0.76	0.48	0.27
% Particles Under	100	87.55	67.79	29.87	10.70	2.57	1.82	0.90

(3) Stability of DHEA-S Dihydrate/Lactose Blend.

This lactose formulation is also placed on an accelerated stability program at 50°C. The results for DHEA-S content are in Table 11. The control is the blend stored at RT. There is no trend in the DHEA-S content over time for either condition and all the results are within the range of samples collected for content uniformity testing (see Table 11). Furthermore, there are no color changes or irregularities observed in the chromatograms. The blend appears to be chemically stable.

Table 11: Stressed stability data on DHEA-S·dihydrate/lactose blend at 50°C.

Time (weeks)	% DHEA-S w/w for control condition	% DHEA-S w/w for stressed condition
0	9.7	9.7
1	9.6	9.6
1.86	9.5	9.7
3	10	9.9

Example 8: Nebulizer Formulation of DHEA-S

Solubility of DHEA-S.

An excess of DHEA-S dihydrate, prepared according to "Recrystallization of DHEA-S·Dihydrate (Example 5)", is added to the solvent medium and allowed to equilibrate for at least 14 hours with some periodic shaking. The suspensions are then filtered through a 0.2 micron syringe filter and immediately diluted for HPLC analysis. To prepare refrigerated samples, the syringes and filters are stored in the refrigerator for at least one hour before use. Inhalation of pure water can produce a cough stimulus. Therefore, it is important to add halide ions to a nebulizer formulation with NaCl being the most commonly used salt. Since DHEA-S is a sodium salt, NaCl could decrease solubility due to the common ion effect. The solubility of DHEA-S at RT (24-26 °C) and refrigerated (7-8 °C) as a function of NaCl concentration is shown in Figure 4. DHEA-S's solubility decrease with NaCl concentration. Lowering the storage temperature decrease the solubility at all NaCl concentrations. The temperature effect is weaker at high NaCl concentrations. For triplicates, the solubility at ~25 °C and 0% NaCl range from 16.5-17.4 mg/mL with a relative standard deviation of 2.7%. At 0.9% NaCl refrigerated, the range for triplicates is 1.1-1.3 mg/mL with a relative standard deviation of 8.3%.

The equilibrium between DHEA-S in the solid and solution states is:

$$NaDHEA\text{-}S_{solid} \leftrightarrow DHEA\text{-}S^{-} + Na^{+}$$

$$K = [DHEA\text{-}S^{-}][Na^{+}]/[NaDHEA\text{-}S]_{solid}$$

Since the concentration of DHEA-S in the solid is constant (i.e., physically stable dihydrate), the equilibrium expression is simplified:

$$Ksp = [DHEA\text{-}S^{-}][Na^{+}]$$

Based on this presumption, a plot of DHEA-S solubility versus the reciprocal of the total sodium cation concentration is linear with a slope equal to Ksp. This is shown in Figures 5 and 6 for equilibrium at RT and refrigerated, respectively. Based on the correlation coefficients, the model is a reasonable fit to the data at both room and refrigerated temperatures where the equilibrium constants were 2236 and 665 mM^2, respectively. To maximize solubility, the NaCl level needs to be as low as possible. The minimum halide ion content for a nebulizer solution should be 20 mM or 0.12% NaCl.

To estimate a DHEA-S concentration for the solution, a 10 °C temperature drop in the nebulizer during use is assumed (i.e., 15 °C). Interpolating between the equilibrium constants versus the reciprocal of absolute temperature, the Ksp at 15 °C would be ~ 1316 mM^2. Each mole of DHEA-S contributes a mole of sodium cation to the solution, therefore:

$$Ksp = [DHEA\text{-}S^-][Na^+]=[DHEA\text{-}S^-][Na^+ + DHEA\text{-}S^-]$$
$$= [DHEA\text{-}S^-]^2 + [Na^+][DHEA\text{-}S^-]$$

which is solve for [DHEA-S^-] using the quadratic formula. The solution for 20 mM Na^+ with a Ksp of 1316 mM^2 is 27.5 mM DHEA-S^- or 10.7 mg/mL. Therefore a 10 mg/mL DHEA-S solution in 0.12% NaCl is selected as a good candidate formulation to progress into additional testing. The estimate for this formula does not account for any concentration effects due to water evaporation from the nebulizer. The pH of a 10 mg/mL DHEA-S solution with 0.12% NaCl range from 4.7 to 5.6. While this would be an acceptable pH level for an inhalation formulation, the effect of using a 20 mM phosphate buffer is evaluated. The solubility results at RT for buffered and unbuffered solutions are shown in Figure 7. The presence of buffer in the formulation suppress the solubility, especially at low NaCl levels. As shown in Figure 8, the solublity data for the buffered solution falls on the same equilibrium line as for the unbuffered solution. The decrease in solubility with the buffer is due to the additional sodium cation content. Maximizing solubility is an important goal and buffering the formulation reduces solubility. Furthermore, Ishihora and Sugimoto ((1979) *Drug Dev. Indust. Pharm.* 5(3) 263-275) did not show a significant improvement in NaDHEA-S stability at neutral pH.

Stability Studies.

A 10 mg/mL DHEA-S formulation is prepared in 0.12% NaCl for a short-term solution stability program. Aliquots of this solution are filled into clear glass vials and stored at RT (24-26 °C) and at 40 °C. The samples are checked daily for DHEA-S content, DHEA content, and appearance. For each time point, duplicate samples are withdrawn and diluted from each vial. The DHEA-S content over the length of this study is shown in Figures 9 and 10. At the accelerated condition, the solution show a faster decomposition rate and became cloudy after two days of storage. The solution stored at RT is more stable and a slight precipitate is observed on the third day. The

study is stopped on day three. DHEA-S decomposition is accompanied by an increase in DHEA content as shown in Figure 10. Since DHEA is insoluble in water, it only takes a small quantity in the formulation to create a cloudy solution (accelerated storage) or a crystalline precipitate (room storage). This explains why earlier visual evaluations of DHEA-S solubility severely underestimate the compound's solubility: small quantities of DHEA would lead the experimenter to conclude the solubility limit of DHEA-S had been exceeded. The solution should easily be stable for the day of reconstitution in a clinical trial. The following section describes the aerosol properties of this formulation.

Nebulizer Studies.

DHEA-S solutions are nebulized using a Pari ProNeb Ultra compressor and LC Plus nebulizer. The schematic for the experiment set-up is shown in Figure 11. The nebulizer is filled with 5 mL of solution and nebulization is continued until the output became visually insignificant (4½ to 5 min.). Nebulizer solutions are tested using a California Instruments AS-6 6-stage impactor with a USP throat. The impactor is run at 30 L/min for 8 s to collect a sample following one minute of nebulization time. At all other times during the experiment, the aerosol is drawn through the by-pass collector at approximately 33 L/min. The collection apparatus, nebulizer, and impactor are rinsed with mobile phase and assayed by HPLC. 5 mL of DHEA-S in 0.12% NaCl is used in the nebulizer. This volume is selected as the practical upper limit for use in a clinical study. The results for the first 5 nebulization experiments are shown below:

Table 12. Results for nebulization studies with DHEA-S

Solution-Nebulizer #	Left in Nebulizer, mg	Deposited in Collector, mg	Deposited in Impactor, mg	Total, mg
10 mg/mL-1	17.9*	16.3	0.38	34.6
10 mg/mL-2	31.2	17.2	0.48	49.0
7.5 mg/mL-1	19.3	16.3	0.35	36.0
7.5 mg/mL-1	21.7	15.4	0.30	37.4
5.0 mg/mL-1	14.4	10.6	0.21	25.2

* Only assayed liquid poured from nebulizer; did not weigh before and after aerosolization or rinse entire unit

Nebulizer #1 runs to dryness in about 5 minutes while Nebulizer #2 takes slightly less than 4.5 minutes. In each case, the liquid volume remaining in the nebulizer is approximately 2 mL. This liquid is cloudy initially after removal from the nebulizer then clears within 3-5 minutes. Even after this time, the 10 mg/mL solutions appear to have a small amount of coarse precipitate in them. Fine air bubbles in the liquid appear to cause the initial cloudiness. DHEA-S appears to be surface active (i.e., promoting foam) and this stabilizes air bubbles within the liquid. The precipitate in 10 mg/mL solutions indicates that the drug substance's solubility is exceeded in the nebulizer environment. Therefore, the additional nebulization experiments in Table 13 are run at lower concentrations. able 13 presents additional data of "dose" linearity versus solution concentration.

Table 13. Results from additional nebulizer experiments with DHEA-S.

Solution-Nebulizer #	Left in Nebulizer, mg	Deposited in Collector, mg	Deposited in Impactor, mg	Total, mg
6.25 mg/mL-2	17.8	12.1	0.24	30.1
7.5 mg/mL-3	21.2	13.8	0.33	35.3

Nebulizer #3 takes slightly less than 4.5 minutes to reach dryness. The mass in the by-pass collector is plotted versus the initial solution concentration in Figure 12. There is good linearity from 0 to 7.5 mg/mL then the amount collected appears to start leveling-off. While the solubility reduction by cooling is included in the calculation of the 10 mg/mL solution, any concentration effects on drug and NaCl content were neglected. Therefore, it is possible for a precipitate to form via supersaturation of the nebulizer liquid. The data in Figure 12 and the observation of some particulates in the 10 mg/mL solution following nebulization indicate that the highest solution concentration for a proof of concept clinical trial formulation is approximately 7.5 mg/mL. An aerosol sample is drawn into a cascade impactor for particle size analysis. There is no detectable trend in particle size distribution with solution concentration or nebulizer number. The average particle size distribution for all nebulization experiments is shown in Figure 13. The aerosol particle size measurements are in agreement with published/advertised results for this nebulizer (i.e., median diameter ~2 μm). While the *in vitro* experiments demonstrate that a nebulizer formulation can

deliver respirable DHEA-S aerosols, the formulation is unstable and takes 4-5 minutes of continuous nebulization. Therefore, a stable DPI formulation has significant advantages. DHEA-S·dihydrate is identified as the most stable solid state for a DPI formulation. An optimal nebulizer formulation is 7.5 mg/mL of DHEA-S in 0.12% NaCl for clinical trials for DHEA-S. The pH of the formulation is acceptable without a buffer system. The aqueous solubility of DHEA-S is maximized by minimizing the sodium cation concentration. Minimal sodium chloride levels without buffer achieve this goal. This is the highest drug concentration with 20 mM of Cl^- that will not precipitate during nebulization. This formulation is stable for at least one day at RT.

Example 9: Preparation of the Experimental Model

Cell cultures, HT-29 SF cells, which represent a subline of HY-29 cells (ATCC, Rockville, Md.) and are adapted for growth in completely defined serum-free PC-1 medium (Ventrex, Portland, ME), were obtained. Stock cultures were maintained in this medium at 37°C (in a humidified atmosphere containing 5% CO_2). At confluence cultures were replated after dissociation using trypsin/EDTA (Gibco, Grand Island, NY) and re-fed every 24 hours. Under these conditions, the doubling time for HT-29 SF cells during logarithmic growth was 24 hours.

Flow Cytometry

Cells were plated at 10^5/60-mm dish in duplicate. For analysis of cell cycle distribution, cultures were exposed to 0, 25, 50, or 200 μM DHEA. For analysis of reversal of cell cycle effects of DHEA, cultures were exposed to either 0 or 25 μM DHEA, and the media were supplemented with MVA, CH, RN, MVA plus CH, or MVA plus CH plus RN or were not supplemented. Cultures were trypsinized following 0, 24, 48, or 74 hours and fixed and stained using a modification of a procedure of Bauer et al., *Cancer Res.* 46, 3173-3178 (1986). Briefly, cells were collected by centrifugation and resuspended in cold phosphate-buffered saline. Cells were fixed in 70% ethanol, washed, and resuspended in phosphate-buffered saline. One ml hypotonic stain solution (50 μg/ml propidium iodide (Sigma Chemical Co.), 20 μg/ml Rnase A (Boehringer Mannheim, Indianapolis, IN), 30 mg/ml polyethylene glycol, 0.1% Triton X-100 in 5 mM citrate buffer) was then added, and after 10 min at room temperature, 1 ml of isotonic stain solution (propidium iodide, polyethylene

glycol, Triton X-100 in 0.4M NaCl) was added and the cells were analyzed using a flow cytometer, equipped with pulse width/pulse area doublet discrimination (Becton Dickinson Immunocytometry Systems, San Jose, CA) After calibration with fluorescent beads, a minimum of $2x10^4$ cells/sample were analyzed, data were displayed s total number of cells in each of 1024 channels of increasing fluorescence intensity, and the resulting histogram was analyzed using the Cellfit analysis program (Becton Dickinson).

DHEA Effect on Cell Growth

Cells were plated 25,000 cells/30 mm dish in quadruplicate, and after 2 days received 0, 12.5, 25, 50, or 200 μM DHEA. Cell number was determined 0, 24, 48, and 72 hours later using a Coulter counter (model Z; Coulter Electronics, Inc. Hialeah, FL). DHEA (AKZO, Basel, Switzerland) was dissolved in dimethyl sulfoxide, filter sterilized, and stored at -20°C until use.

Figure 14 illustrates the inhibition of growth for HT-29 cells by DHEA. Points refer to numbers of cells, and bars refer to SEM. Each data point was performed in quadruplicate, and the experiment was repeated three times. Where SEM bars are not apparent, SEM was smaller than symbol. Exposure to DHEA resulted in a reduced cell number compared to controls after 72 hours in 12.5 μM, 48 hours in 25 or 50 μM, and 24 hours in 200 μM DHEA, indicating that DHEA produced a time- and dose-dependent inhibition of growth.

DHEA Effect on Cell Cycle

To examine the effects of DHEA on cell cycle distribution, HT-29 SF cells were plated (10^5 cells/60 mm dish), and 48 hours later treated with 0,25, 50, or 200 μM DHEA. Fig. 15 illustrates the effects of DHEA on cell cycle distribution in HT-29 SF cells. After 24, 48, and 72 hours, cells were harvested, fixed in ethanol, and stained with propidium iodide, and the DNA content/cell was determined by flow cytometric analysis. The percentage of cells in G_1, S, and G_2M phases was calculated using the Cellfit cell cycle analysis program. S phase is marked by a quadrangle for clarity. Representative histograms from duplicate determinations are shown. The experiment was repeated three times.

The cell cycle distribution in cultures treated with 25 or 50 μM DHEA was unchanged after the initial 24 hours. However, as the time of exposure to DHEA

increased, the proportion of cells in S phase progressively decreased, and the percentage of cells in G_1, S and G_2M phases was calculated using the Cellfit cell cycle analysis program. S phase is marked by a quadrangle for clarity. Representative histograms from duplicate determinations are shown. The experiment was repeated three times.

The cell cycle distribution in cultures treated with 25 or 50 μM DHEA was unchanged after the initial 24 hours. However, as the time of exposure to DHEA increased, the proportion of cells in S phase progressively decreased and the percentage of cells in G_1 phase was increased after 72 hours. A transient increase in G_2M phase cells was apparent after 48 hours. Exposure to 200μM DHEA produced a similar but more rapid increase in the percentage of cells in G_1 and a decreased proportion of cells in S phase after 24 hours, which continued through the treatment. This indicates that DHEA produced a G_1 block in HT-29 SF cells in a time-and dose-dependent manner.

Example 10: Reversal of DHEA-mediated Effect on Growth and Cell Cycle

Reversal of DHEA-mediated Growth Inhibition.

Cells were plated as above, and after 2 days received either 0 or 25 μM DHEA-containing medium supplemented with mevalonic acid ("MVA"; mM) squalene (SQ; 80 μM), cholesterol (CH; 15 μg/ml), MVA plus CH, ribonucleosides (RN; uridine, cytidine, adenosine, and guanosine at final concentrations of 30 μM each), deoxyribonucleosides (DN; thymidine, deoxycytidine, deoxyadenosine and deoxyguanosine at final concentrations of 20 μM each). RN plus DN, or MVA plus CH plus RN, or medium that was not supplemented. All compounds were obtained from Sigma Chemical Co. (St. Louis, MO) Cholesterol was solubilized in ethanol immediately before use. RN and DN were used in maximal concentrations shown to have no effects on growth in the absence of DHEA.

Figure 16 illustrates the reversal of DHEA-induced growth inhibition in HT-29 SF cells. In A, the medium was supplemented with 2 μM MVA, 80 μM SQ, 15 μg/ml CH, or MVA plus CH (MVA+CH) or was not supplemented (CON). In B, the medium was supplemented with a mixture of RN containing uridine, cytidine, adenosine, and guanosine in final concentrations of 30 μM each; a mixture of DN containing thymidine, deoxycytidine, deoxyadenosine and deoxyguanosine in final

concentrations of 20 μM each; RN plus DN (RN+DN); or MVA plus CH plus RN (MVA+CH+RN). Cell numbers were assessed before and after 48 hours of treatment, and culture growth was calculated as the increase in cell number during the 48 hour treatment period. Columns represent cell growth percentage of untreated controls; bars represent SEM. Increase in cell number in untreated controls was 173,370”6518. Each data point represents quadruplicate dishes from four independent experiments. Statistical analysis was performed using Student’s t test κ $p<0.01$; ψ p<, 0.001; compared to treated controls. Note that supplements had little effect on culture growth in absence of DHEA.

Under these conditions, the DHEA-induced growth inhibition was partially overcome by addition of MVA as well as by addition of MVA plus CH. Addition of SQ or CH alone had no such effect. This suggest that the cytostatic activity of DHEA was in part mediated by depletion of endogenous mevalonate and subsequent inhibition of the biosynthesis of an early intermediate in the cholesterol pathway that is essential for cell growth. Furthermore, partial reconstitution of growth was found after addition of RN as well as after addition of RN plus DN but not after addition of DN, indicating that depletion of both mevalonate and nucleotide pools is involved in the growth-inhibitory action of DHEA. However, none of the reconstitution conditions including the combined addition of MVA, CH, and RN completely overcame the inhibitory action of DHEA, suggesting either cytotoxic effects or possibly that additional biochemical pathways are involved.

Reversal of DHEA Effect on Cell Cycle

HT-29 SF cells were treated with 25 FM DHEA in combination with a number of compounds, including MVA, CH, or RN, to test their ability to prevent the cell cycle-specific effects of DHEA. Cell cycle distribution was determined after 48 and 72 hours using flow cytometry.

Figure 17 illustrates reversal of DHEA-induced arrest in HT-29 SF cells. Cells were plated (10^5 cells/60 mm dish) and 48 hours later treated with either 0 or 25 FM DHEA. The medium was supplemented with 2 FM MVA; 15 Fg/ml CH; a mixture of RN containing uridine, cytidine, adenosine, and guanosine in final concentrations of 30 FM; MVA plus CH (MVA+CH); or MVA plus CH plus RN (MVA+CH+RN) or was not supplemented. Cells were harvested after 48 or 72 hours, fixed in ethanol,

and stained with propidium iodine, and the DNA content per cell was determined by flow cytometric analysis. The percentage of cells in G_1, S, and G_2M phases were calculated using the Cellfit cell cycle profile analysis program. S phase is marked by a quadrangle for clarity. Representative histograms from duplicative determinations are shown. The experiment was repeated two times. Note that supplements had little effect on cell cycle progression in the absence of DHEA.

With increasing exposure time, DHEA progressively reduced the proportion of cells in S phase. While inclusion of MVA partially prevented this effect in the initial 48 hours but not after 72 hours, the addition of MVA plus CH was also able to partially prevent S phase depletion at 72 hours, suggesting a requirement of both MVA and CH for cell progression during prolonged exposure. The addition of MVA, CH, and RN was apparently most effective at reconstitution but still did not restore the percentage of S phase cells to the value seen in untreated control cultures. CH or RN alone had very little effect at 48 hours and no effect at 72 hours. Morphologically, cells responded to DHEA by acquiring a rounded shape, which was prevented only by the addition of MVA to the culture medium. Some of the DNA histograms after 72 hours DHEA exposure in FIG.4 also show the presence of a subpopulation of cells possessing apparently reduced DNA content. Since the HT-29 cell line is known to carry populations of cells containing varying numbers of chromosomes (68-72; ATCC), this may represent a subset of cells that have segregated carrying fewer chromosomes.

Conclusions

The Examples 9-10 above provide evidence that in vitro exposure of HT-29 SF human colonic adenocarcinoma cells to concentrations of DHEA known to deplete endogenous mevalonate results in growth inhibition and G_1 arrest and that addition of MVA to the culture medium in part prevents these effects. DHEA produced effects upon protein isoprenylation which were in many respects similar to those observed for specific 3-hydroxy-3-methyl-glutaryl-CoA reductase inhibitors such as lovastatin and compactin. Unlike direct inhibitors of mevalonate biosynthesis, however, DHEA mediates its effects upon cell cycle progression and cell growth in a pleiotropic manner involving ribo-and deoxyribonucleotide biosynthesis and possibly other factors as well.

Example 11: Metered Dose Inhaler

Active Ingredient	Target per Actuation
Salmeterol (as hydroxynaphthoate)	25.0 µg
DHEA	400 mg
Stabilizer	5.0 µg
Trichlorofluoromethane	23.70 mg
Dichlorodifluoromethane	61.25 mg

Example 12: Metered Dose Inhaler

Active Ingredient	Target per Actuation
Salmeterol (as hydroxynaphthoate)	25.0 µg
DHEA-S	400 mg
Stabilizer	7.5 µg
Trichlorofluoromethane	23.67 mg
Dichlorodifluoromethane	61.25 mg

Example 13: Metered Dose Inhaler

Active Ingredient	Target per Actuation
formoterol fumarate	12.0 µg
DHEA	400.0 mg
Stabilizer	15.0 µg
Trichlorofluoromethane	23.56 mg
Dichlorodifluoromethane	61.25 mg

Example 14: Metered Dose Inhaler

Active Ingredient	Target per Actuation
formoterol fumarate	12.0 µg
DHEA-S	400.0 mg
Stabilizer	15.0 µg
Trichlorofluoromethane	23.56 mg
Dichlorodifluoromethane	61.25 mg

In the following Examples 15-18, the first and second active agents are micronized and bulk blended with lactose in the proportions given above. The blend is filled into hard gelatin capsules or cartridges or into specifically constructed double foil blister packs (Rotadisks blister packs, Glaxo® to be administered by an inhaler such as the Rotahaler inhaler (Glaxo®) or in the case of the blister packs with the Diskhaler inhaler (Glaxo®).

Example 15: Metered Dose Dry Powder Formulation

Active Ingredient	/cartridge or blister
Salmeterol (hydroxynaphthoate)	72.5 µg
DHEA	1 mg
Lactose Ph. Eur.	To 12.5 or 25.0 mg

Example 16: Metered Dose Dry Powder Formulation

Active Ingredient	/cartridge or blister
Salmeterol (hydroxynaphthoate)	12.5 µg
DHEA-S	1 mg
Lactose Ph. Eur.	To 12.5 or 25.0 mg

Example 17: Metered Dose Dry Powder Formulation

Active Ingredient	/cartridge or blister
Salmeterol (hydroxynaphthoate)	72.5 µg
DHEA	1 mg
Lactose Ph. Eur.	To 12.5 or 25.0 mg

Example 18: Metered Dose Dry Powder Formulation

Active Ingredient	/cartridge or blister
Salmeterol (hydroxynaphthoate)	72.5 µg
DHEA-S	1 mg
Lactose Ph. Eur.	To 12.5 or 25.0 mg

Example 19: Effects of cAMP on Human Airway Smooth Muscle Cell Function in Asthma

In the following study, how DHEA-S can augment the effects of drugs acting via cAMP in inhibiting human ASM cell growth was evaluated. The mitogen chosen for this study (EGF) is increased in the bronchoalveolar lavage (BAL) of patients with asthma or those asthmatics who underwent allergen challenge. The concentrations used approximate to those found in the BAL fluid from such studies.

Human ASM cells were obtained from the distal trachea of transplant donors in compliance with the University of Pennsylvania's Institutional Review Board. Distal trachealis was dissected free of connective tissue and enzymatically digested to yield approximately 1 x 10^4 human ASM cells. These cells were then grown to confluence in 10% fetal bovine serum and quiesced in serum-free media containing 1% bovine serum albumin for 24 hours. Subsequently, cells were incubated either with diluent or EGF (1 ng/ml) for 24 hours. After the initial incubation, cells were then exposed to [^{3}H]-thymidine and DNA synthesis was measured over 24 hours using thymidine incorporation. The cells were scraped and placed on filters which were then counted using a beta counter. All experiments were repeated in a minimum of three human ASM cell lines. All conditions per experiment were run with six replicates. Data is represented as mean ± standard error of the mean and statistical analysis was performed using Bonferroni-Dunn correction and ANOVA with a significance demonstrated with $P < 0.05$.

In some experiments, cells were incubated with the specified growth factor and 10µM DHEA-S. DNA synthesis, as measured by thymidine incorporation, was performed as described above. The concentration to inhibit 50% of the growth factor-induced DNA synthesis was calculated and described as the IC_{50}.

The role of DHEA-S in modulating cAMP concentration on ASM mitogenesis

was also examined. cAMP is known to be important as a secondary signaling pathway for smooth muscle proliferation. There are three classes of commonly used anti-asthma medications that alter cAMP concentrations and/or adenosine concentrations. These classes of drugs are: methylxanthines, beta-agonists and phosphodiesterase inhibitors. For each of these drug classes, isobutylmethylxanthine (IBMX) was used as the tool compound to inhibit smooth muscle proliferation. In these experiments, DHEA-S at an IC_{50} concentration was added simultaneously to cells treated with IBMX and the effects on growth factor-induced DNA synthesis were examined.

Figure 18 depicts the effect of EGF, IBMX, and DHEA-S on ^{3}H-thymidine incorporation in human smooth muscle cultures. These data show that neither IBMX nor DHEA-S alone or in combinations inhibits smooth muscle proliferation. EGF at 1 ng/mL concentration stimulated ^{3}H thymidine incorporation approximately 10 fold.

In this series of experiments, the ability of DHEA-S alone and in combination with IBMX was studied for the ability to inhibit EGF stimulated proliferation. As can be seen from Figure 19, DHEA-S at concentrations of 10 μM reduced ASM proliferation by approximately 11% (comparison of second and third columns). IBMX (0.1- 100 μM) had variable effects on thymidine incorporation (columns 8-11). However, when this was repeated in the presence of 10 μM DHEA-S, IBMX caused a concentration-dependent (columns 4-7) inhibition of smooth muscle proliferation with an IC_{50} around 10 μM.

These data show that the combination of DHEA-S and IBMX had a greater effect on inhibiting smooth muscle proliferation than either agent alone. IBMX was used as a tool compound to evaluate the actions of cAMP. Thus, these experiments indicate that agents that modulate cAMP, such as beta-agnists, can be used in combination with DHEA-S in the treatment of respiratory diseases. Not intending to limit to one mechanism of action, the inhibition of smooth muscle proliferation would be expected to prevent ongoing airway remodeling noted in patients with moderate to severe asthma. These results demonstrate that combining DHEA-S with a cAMP modulator, such as a beta-agonist, may lead to a reduction in airway smooth muscle remodeling.

Example 20: Effects of DHEA-S and Salbutamol on Histamine Release

Model 1: Rat Peritoneal Mast Cells

Freshly isolated rat peritoneal mast cells ($2x10^5$ cells) were pre-incubated for 5 min at 37°C in a Balanced Salt Solution containing 150 mM NaCl (pH 7.4), 2.7 mM KCl, 0.9 mM CaCl2, 4 mM Na2HPO4, 2.7 mM KH2PO4, 1.75 mg/ml BSA and 0.1 µg/ml compound 48/80. Thereafter, DHEA-S, combination of DHEA-S and salbutamol, or water (control) are added and the mixture is incubated for 2 min at 37°C.

Following incubation, the mixture is cooled to 4°C then centrifuged at 4500 rpm for 5 min. The supernatant is collected and mixed for 30 min at 4°C with 5% TCA (trichloroacetic acid) to cause protein precipitation. After centrifugation at 9500 x g for 15 min, the supernatant is collected and mixed with 0.25 M HCl. Thereafter, the samples are mixed with 2M NaOH and 0.2% OPT (ortho-phtalaldehyde) and incubated for 30 min at 4°C in darkness, after which time the reaction is stopped by addition of 0.5 M H_2SO_4.

The amount of secreted histamine is quantified by measuring the fluorescence intensity at $\lambda ex = 360$ nm and $\lambda em = 450$ nm using a spectrofluorimeter (GeminiXS, Molecular Devices). The results are expressed as a percent inhibition of control histamine secretion.

Salbutamol was studied at a concentration 1, 10 and 100 pM, a range known to encompass the EC50 for binding to the β_2 adrenoceptor. There was no or trivial inhibition of compound 48/80-induced histamine release at these concentrations (see table below). DHEA-S at 30 µM caused a 46% inhibition of mast cell release. When 30 µM DHEAS was combined with 10 or 100 pM salbutamol a synergistic effect was observed with the inhibition of 93% being greater than either drug alone.

% inhibition of histamine release
DHEA-S and Salbutamol
Rat mast cell 48/80 histamine release

		DHEA-S		
		0	30 uM	100 uM
Control	0		46	8
Salbutamol	1pM	-1	41	32
	10pM	7	93	36
	100pM	2	93	2

Since beta-agonists are established to be effective in respiratory disease, it follows from the above findings that the combination of DHEA-sulfate together with a beta-agonist will act in a multiplicative or synergistic fashion to reduce the deleterious role of histamine release in an allergic patient. Therefore, the patient will derive clinical benefit from the combination of these two drugs.

In addition to DHEA-sulfate, other suitable non-glucocorticoid steroids can be used as the first active agent, including, but not limited to, epiandrosterone and derivative, analogs, and pharmaceutically acceptable salts thereof. For example, such as the compounds depicted by Formulas I, III, and IV, herein.

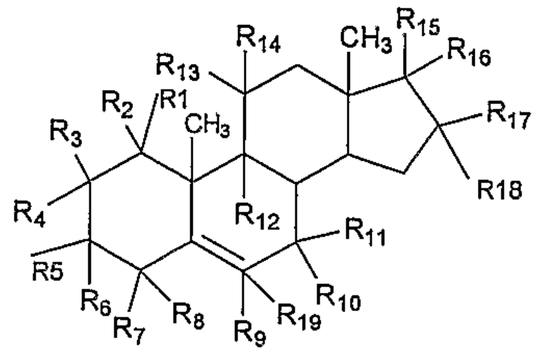
Although the invention has been described with reference to the presently preferred embodiments, it should be understood that various modifications can be made without departing from the spirit of the invention.

All publications, patents, and patent applications, and web sites are herein incorporated by reference in their entirety to the same extent as if each individual publication, patent, or patent application, was specifically and individually indicated to be incorporated by reference in its entirety.

WHAT IS CLAIMED IS:

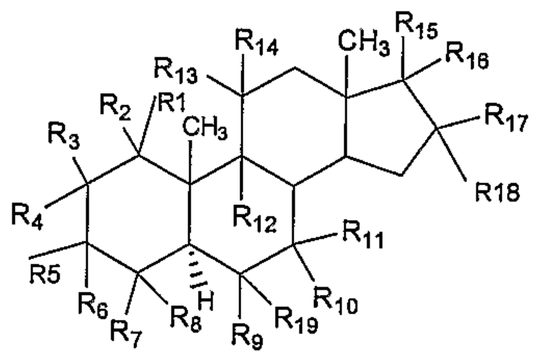
1. A pharmaceutical composition, comprising a pharmaceutically or veterinarily acceptable carrier, a first active agent and a second active agent effective to treat asthma, chronic obstructive pulmonary disease, or a respiratory or lung disease,

(a) the first active agent is at least one of a non-glucocorticoid steroid selected from a non-glucocorticoid steroid having the chemical formula



(III)

and a non-glucocorticoid steroid of the chemical formula



(IV)

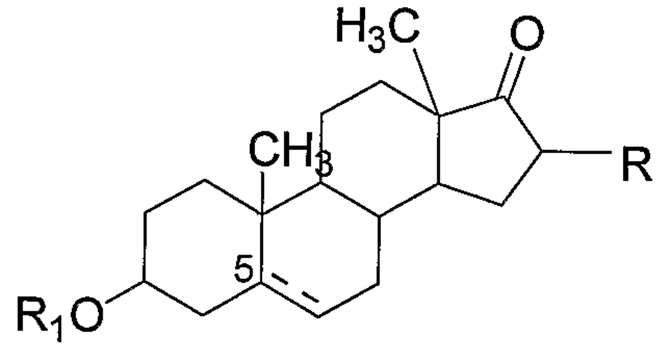
;

wherein R1, R2, R3, R4. R5, R7, R8, R9, R10, R12, R13, R14 and R19 are independently H, OR, halogen, (C1-C10) alkyl or (C1-C10) alkoxy, R5 and R11 are independently OH, SH, H, halogen, pharmaceutically acceptable ester, pharmaceutically acceptable thioester, pharmaceutically acceptable ether, pharmaceutically acceptable thioether, pharmaceutically acceptable inorganic esters, pharmaceutically acceptable monosaccharide, disaccharide or oligosaccharide, spirooxirane, spirothirane, -OSO2R20, -OPOR20R21 or (C1-C10) alky, R5 and R6 taken together are =O, R10 and R11 taken together are =O; R15 is (1) H, halogen, (C1-C10) alkyl, or (C1-C10) alkoxy when R16 is -C(O)OR22, (2) H, halogen, OH or (C1-C10) alkyl when R16 is halogen, OH or (C1-C10) alkyl, (3) H, halogen, (C1-C10) alkyl, (C1-C10) alkenyl, (C1-C10) alkynyl, formyl, (C1-C10) alkanoyl or epoxy when R16 is OH, (4) OR, SH, H, halogen, pharmaceutically acceptable ester, pharmaceutically acceptable thioester, pharmaceutically acceptable ether, pharmaceutically acceptable thioether, pharmaceutically acceptable inorganic esters, pharmaceutically acceptable monosaccharide, disaccharide or oligosaccharide, spirooxirane, spirothirane, -OSO2R20 or -OPOR20R21 when R16 is H, or R15 and

R16 taken together are =O; R17 and R18 are independently (1) H, -OH, halogen, (C1-C10) alkyl or -(C1-C10) alkoxy when R6 is H OR, halogen. (C1-C10) alkyl or -C(O)OR22, (2) H, (C1-C10 alkyl).amino, ((C1-C10) alkyl)n amino-(C1-C10) alkyl, (C1-C10) alkoxy, hydroxy - (C1-C10) alkyl, (C1-C10) alkoxy - (C1-C10) alkyl, (halogen)m (C1-C10) alkyl, (C1-C10) alkanoyl, formyl, (C1-C10) carbalkoxy or (C1-C10) alkanoyloxy when R15 and R16 taken together are =O, (3) R17 and R18 taken together are =O; (4) R17 or R18 taken together with the carbon to which they are attached form a 3-6 member ring containing 0 or 1 oxygen atom; or (5) R15 and R17 taken together with the carbons to which they are attached form an epoxide ring; R20 and R21 are independently OH, pharmaceutically acceptable ester or pharmaceutically acceptable ether; R22 is H, (halogen)m (C1-C10) alkyl or (C1-C10) alkyl; n is 0, 1 or 2; and m is 1, 2 or 3; or pharmaceutically or veterinarily acceptable salts thereof; and

(b) the second active agent is a β2-agonist bronchodilator.

2. The pharmaceutical composition of claim 1 wherein said first active agent is a non-glucocorticoid steroid having the chemical formula



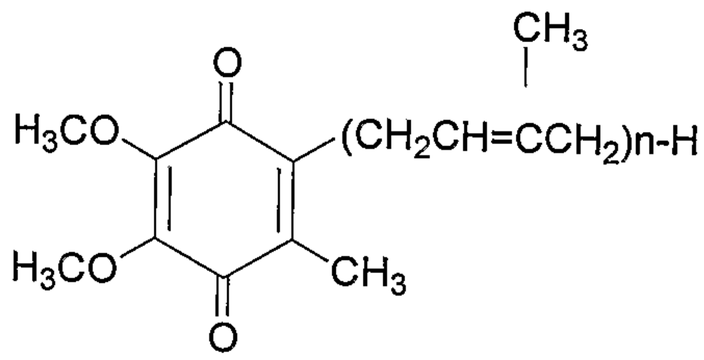
(I)

wherein the broken line represents a single or a double bond; R is hydrogen or a halogen; the H at position 5 is present in the alpha or beta configuration or the compound of chemical formula I comprises a racemic mixture of both configurations; and R^1 is hydrogen or a multivalent inorganic or organic dicarboxylic acid covalently bound to the compound.

3. The pharmaceutical composition of claim 1, wherein the first active agent is a non-glucocorticoid steroid having the chemical formula (I), wherein said multivalent organic dicarboxylic acid is SO_2OM, phosphate or carbonate, wherein M comprises a counterion, wherein said counterion is H, sodium, potassium, magnesium, aluminum, zinc, calcium, lithium, ammonium, amine, arginine, lysine, histidine, triethylamine, ethanolamine, choline, triethanoamine, procaine, benzathine,

tromethanine, pyrrolidine, piperazine, diethylamine, sulfatide

$$-SO_2O-CH_2CHCH_2OCOR^3;$$
$$\quad\quad\quad



(II)

wherein n is 1 to 12.

8. The pharmaceutical composition of claim 1, wherein the pharmaceutical composition comprises particles of inhalable or respirable size.

9. The pharmaceutical composition of claim 8, wherein the particles are about 0.01 μm to about 10 μm in size.

10. The pharmaceutical composition of claim 8, wherein the particles are about 10 μm to about 100 μm in size.

11. A kit comprising a delivery device and the pharmaceutical composition of claim 1.

12. The kit of claim 11, wherein the delivery device is an aerosol generator or spray generator.

13. The kit of claim 11, wherein the aerosol generator comprises an inhaler.

14. The kit of claim 13, wherein the inhaler delivers individual pre-metered doses of the formulation

15. The kit of claim 13, wherein the inhaler comprises a nebulizer or insufflator.

16. A method for reducing the probability of or treating asthma in a subject, comprising administering to a subject in need of such treatment a

prophylactically or therapeutically effective amount of the pharmaceutical composition of claim 1.

17. A method for reducing the probability of or treating of chronic obstructive pulmonary disease in a subject, comprising administering to a subject in need of such treatment a prophylactically or therapeutically effective amount of the pharmaceutical composition of claim 1.

18. A method for treatment of respiratory, lung or malignant disorder or condition, or for reducing levels of, or sensitivity to, adenosine or adenosine receptors in a subject, comprising administering to a subject in need of such treatment a prophylactically or therapeutically effective amount of the pharmaceutical composition of claim 1.

19. The method of claim 18, wherein the disorder or condition comprises asthma, chronic obstructive pulmonary disease (COPD), cystic fibrosis (CF), dyspnea, emphysema, wheezing, pulmonary hypertension, pulmonary fibrosis, hyper-responsive airways, increased adenosine or adenosine receptor levels, adenosine hyper-sensitivity, infectious diseases, pulmonary bronchoconstriction, respiratory tract inflammation or allergies, lung surfactant or ubiquinone depletion, chronic bronchitis, bronchoconstriction, difficult breathing, impeded or obstructed lung airways, adenosine test for cardiac function, pulmonary vasoconstriction, impeded respiration, Acute Respiratory Distress Syndrome (ARDS), administration of adenosine or adenosine level increasing drugs, infantile Respiratory Distress Syndrome (infantile RDS), pain, allergic rhinitis, cancer, or chronic bronchitis.

Fig. 1

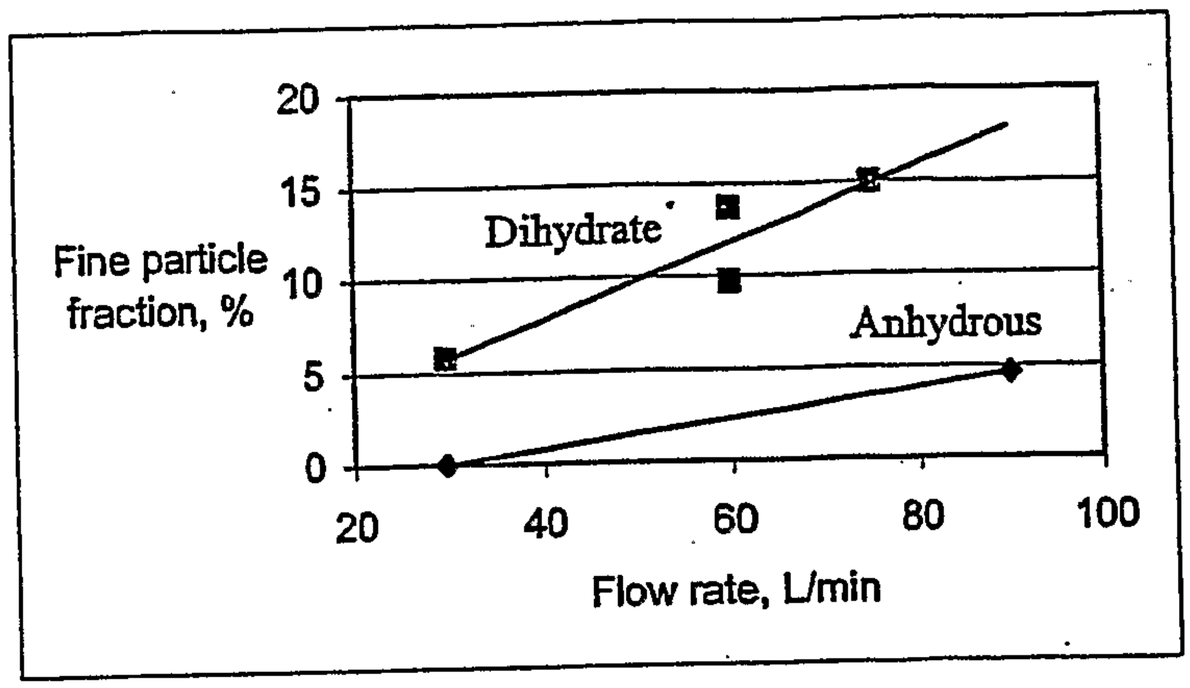

Fine particle fraction, %
20
15
10
5
0
Dihydrate
Anhydrous
20
40
60
80
100
Flow rate, L/min

Fig. 2

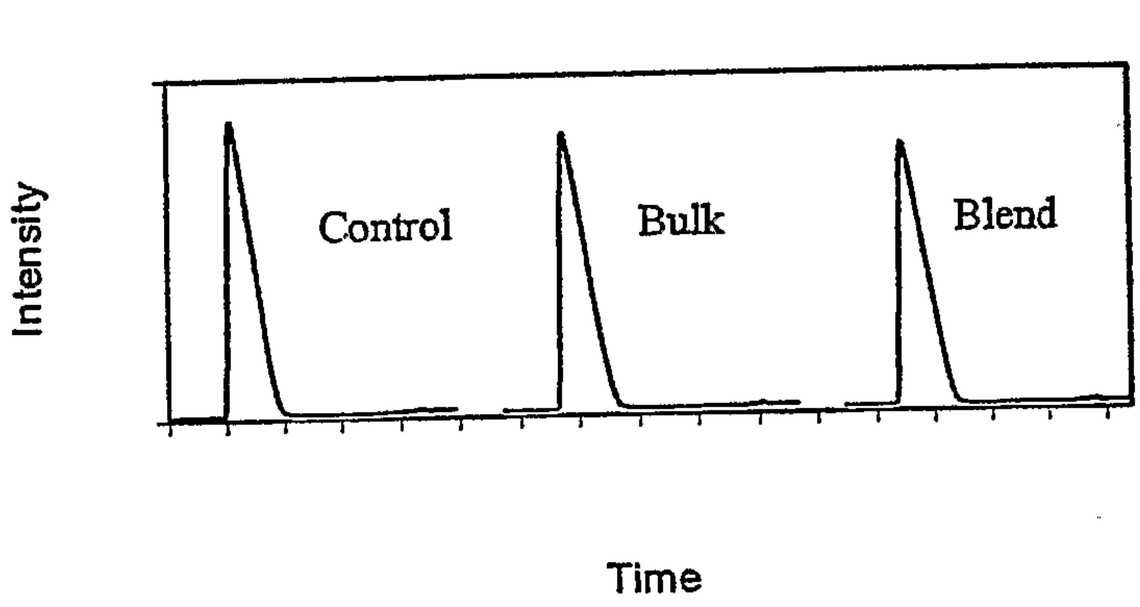

Intensity
Control
Bulk
Blend
Time

Fig. 3

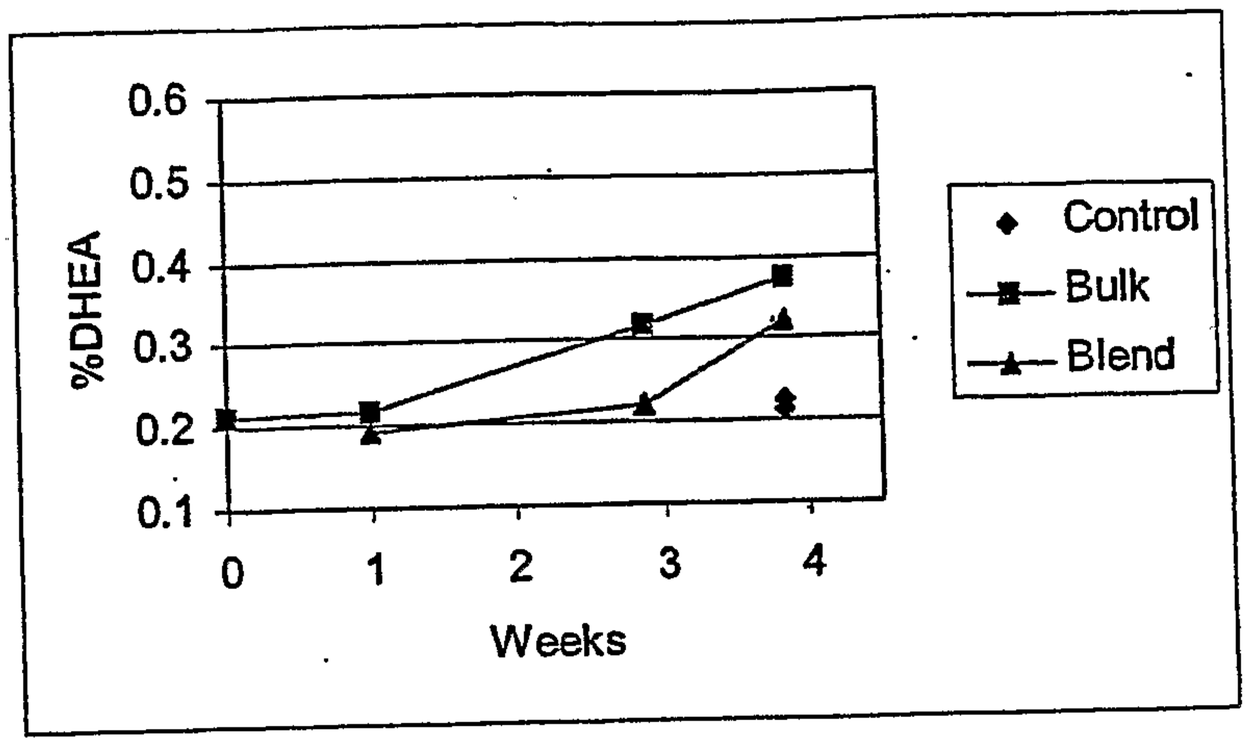

0.6
0.5
0.4
0.3
0.2
0.1
%DHEA
0
1
2
3
4
Weeks
Control
Bulk
Blend

Fig. 4

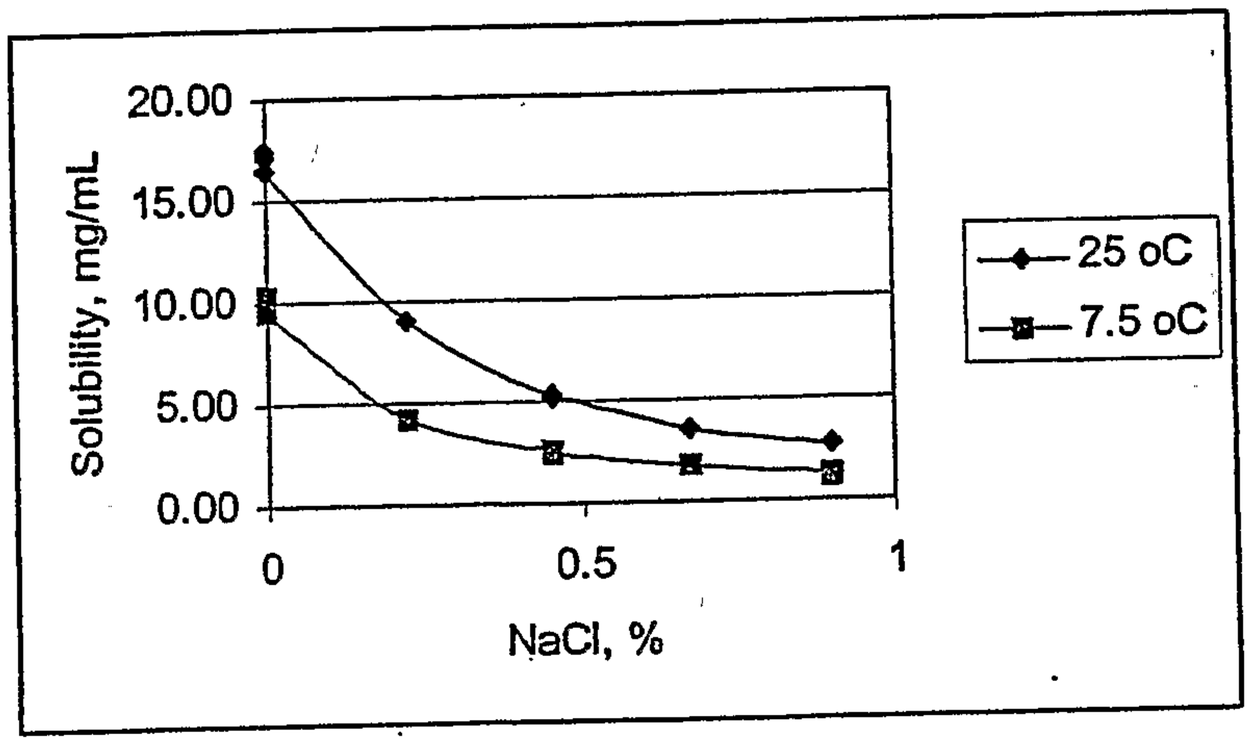

20.00
15.00
10.00
5.00
0.00
Solubility, mg/mL
0
0.5
1
NaCl, %
25 oC
7.5 oC

Fig. 5

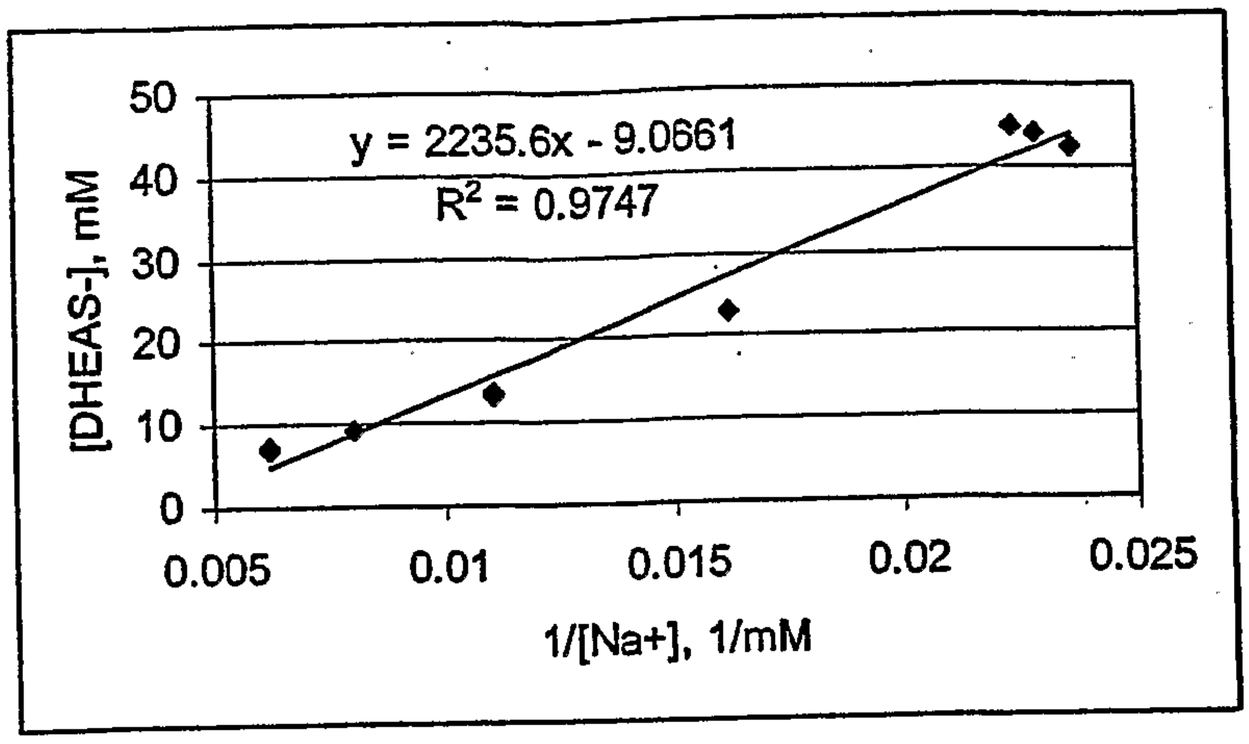

y = 2235.6x - 9.0661
R² = 0.9747
[DHEAS-], mM
50
40
30
20
10
0
0.005
0.01
0.015
0.02
0.025
1/[Na+], 1/mM

Fig. 6

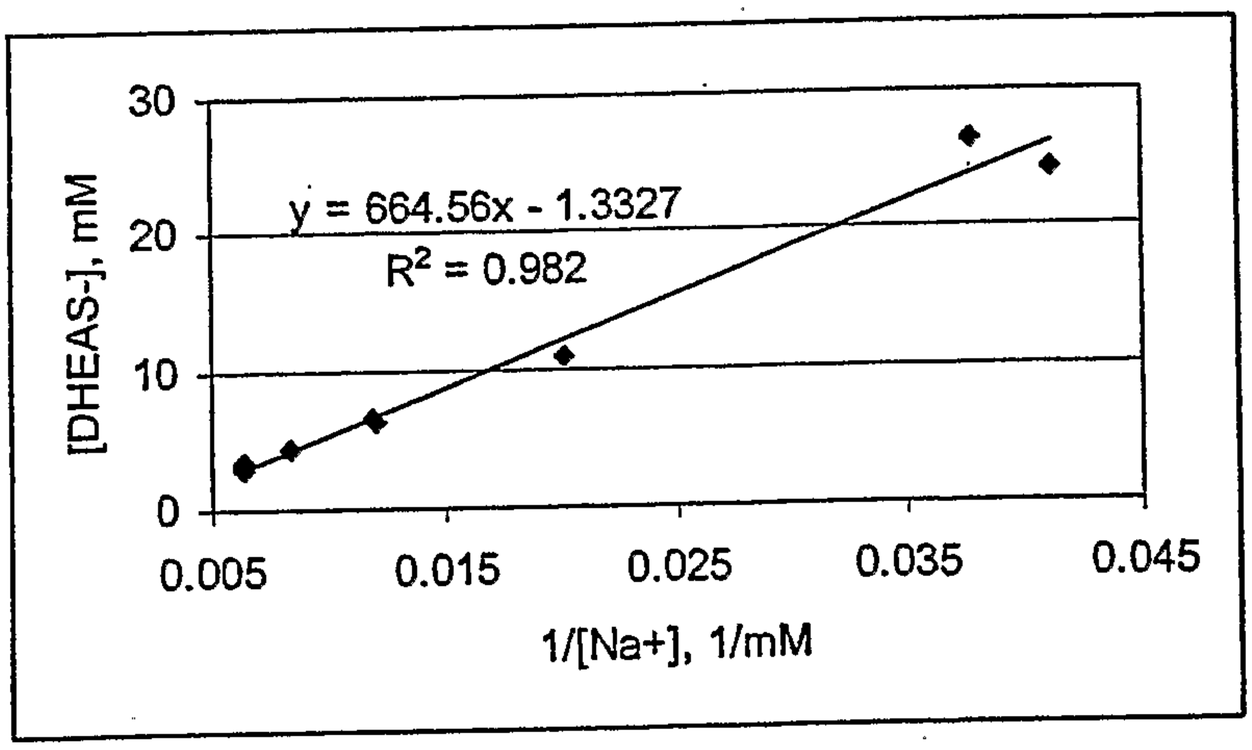

y = 664.56x - 1.3327
R² = 0.982
[DHEAS-], mM
30
20
10
0
0.005
0.015
0.025
0.035
0.045
1/[Na+], 1/mM

Fig. 7

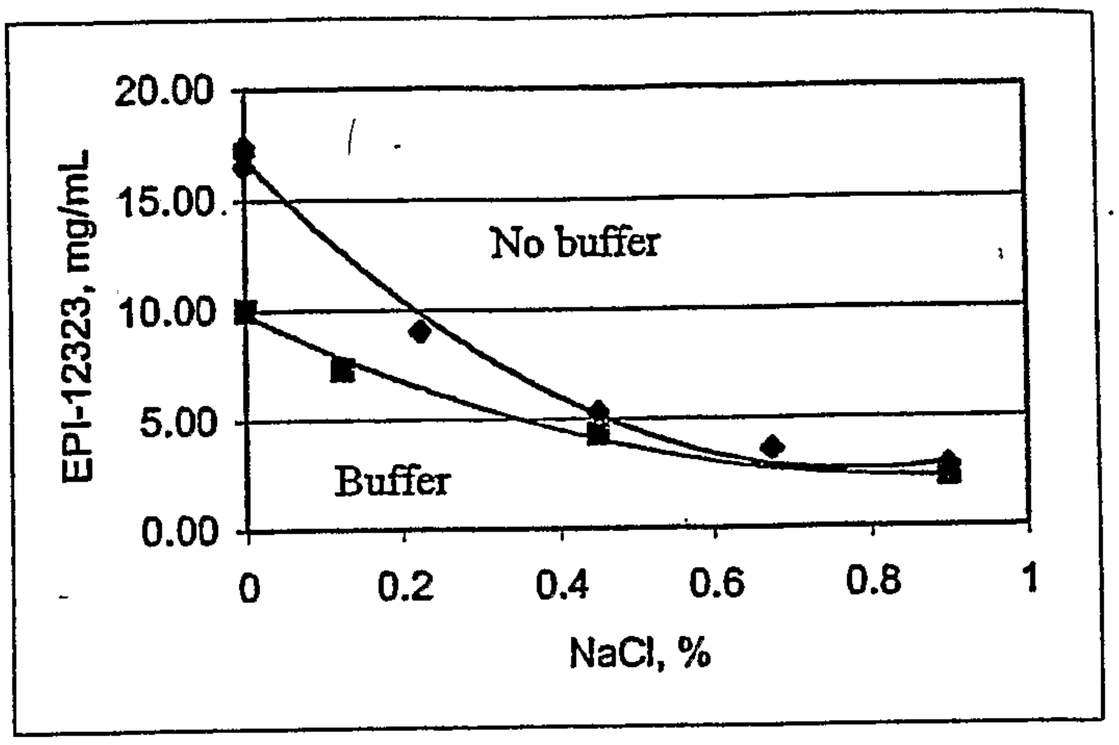

20.00
15.00
10.00
5.00
0.00
EPI-12323, mg/mL
No buffer
Buffer
0
0.2
0.4
0.6
0.8
1
NaCl, %

Fig. 8

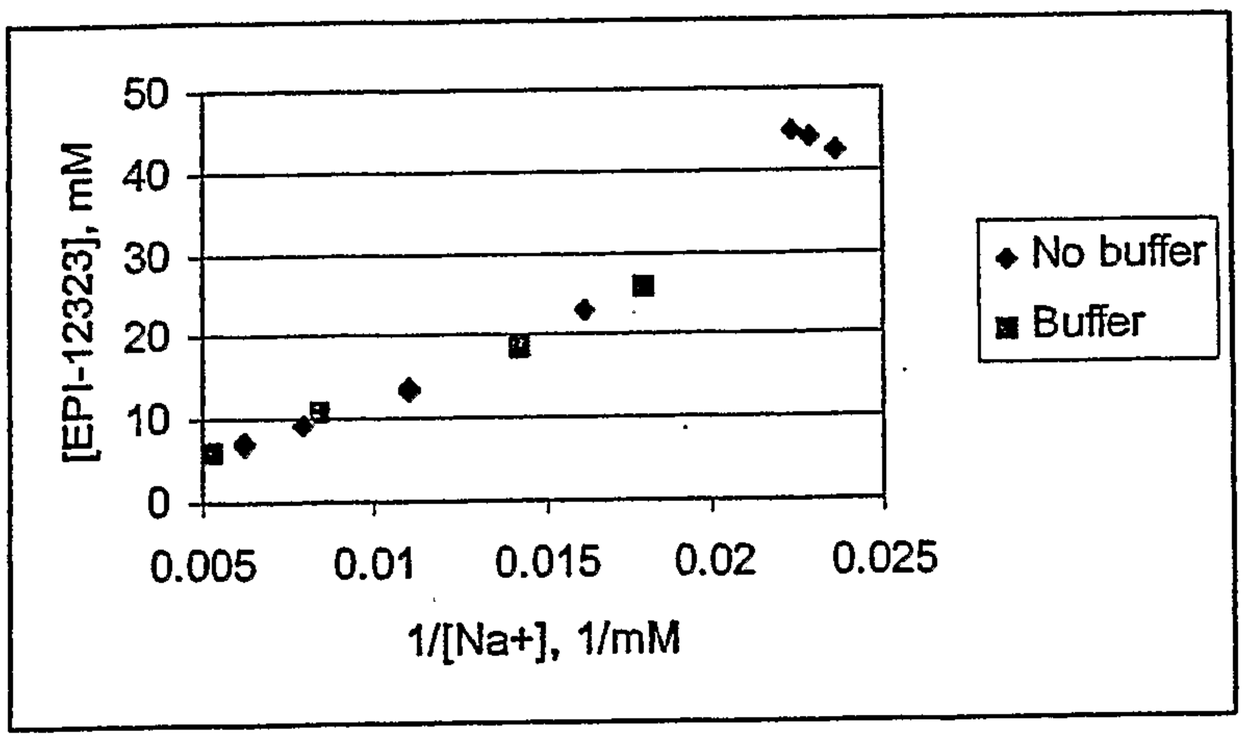

50
40
30
20
10
0
[EPI-12323], mM
0.005
0.01
0.015
0.02
0.025
1/[Na+], 1/mM
No buffer
Buffer

Fig. 9

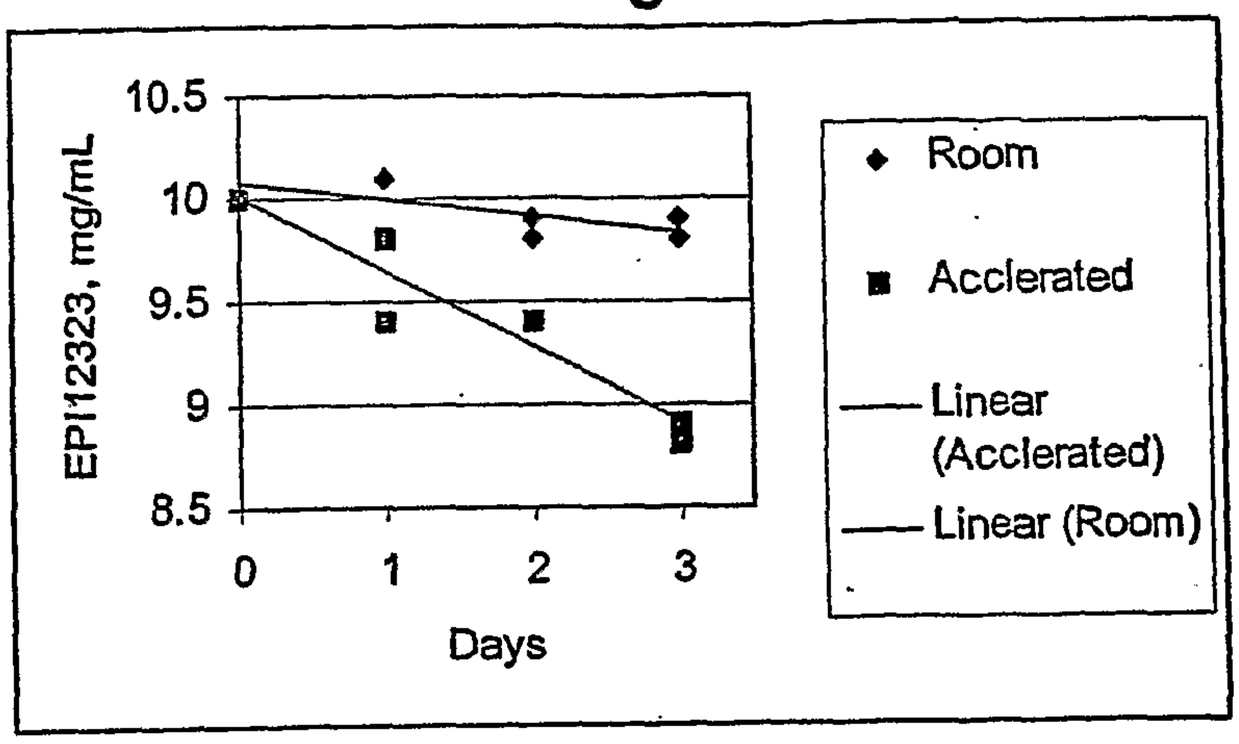

EPI12323, mg/mL
10.5
10
9.5
9
8.5
0
1
2
3
Days
Room
Acclerated
Linear (Acclerated)
Linear (Room)

Fig. 10

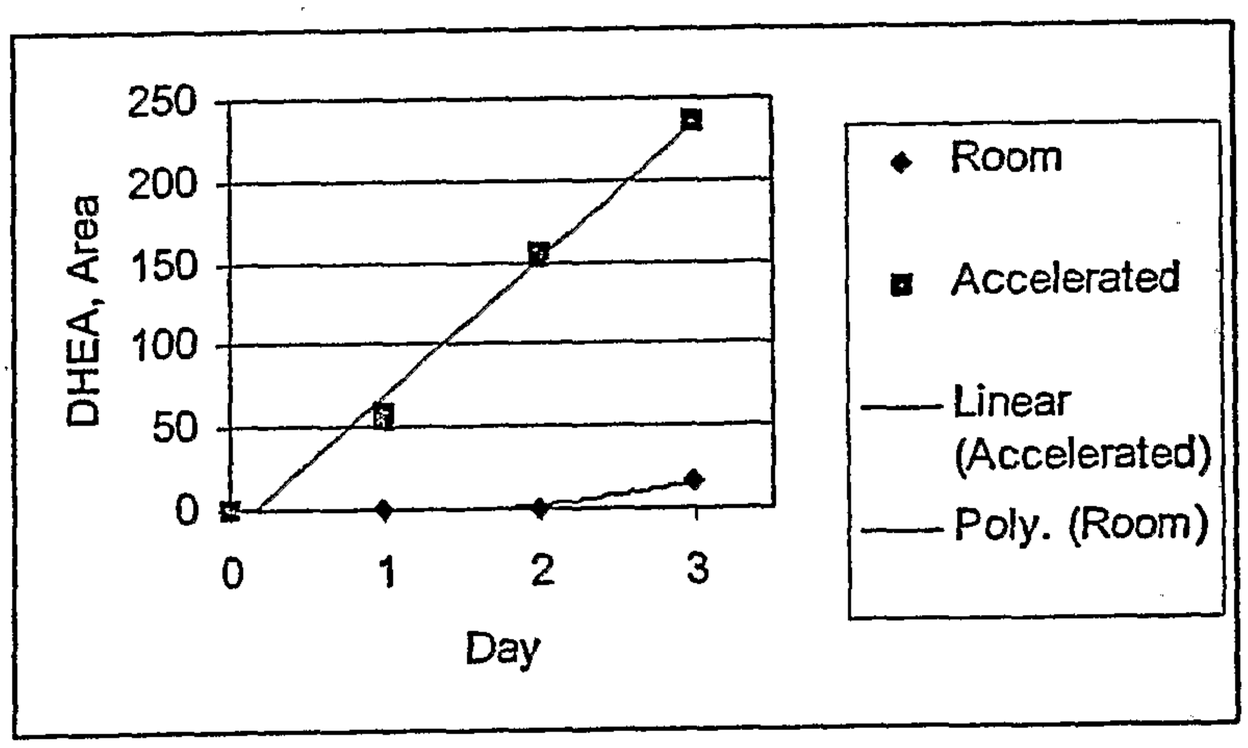

DHEA, Area
250
200
150
100
50
0
0
1
2
3
Day
Room
Accelerated
Linear (Accelerated)
Poly. (Room)

Fig. 11

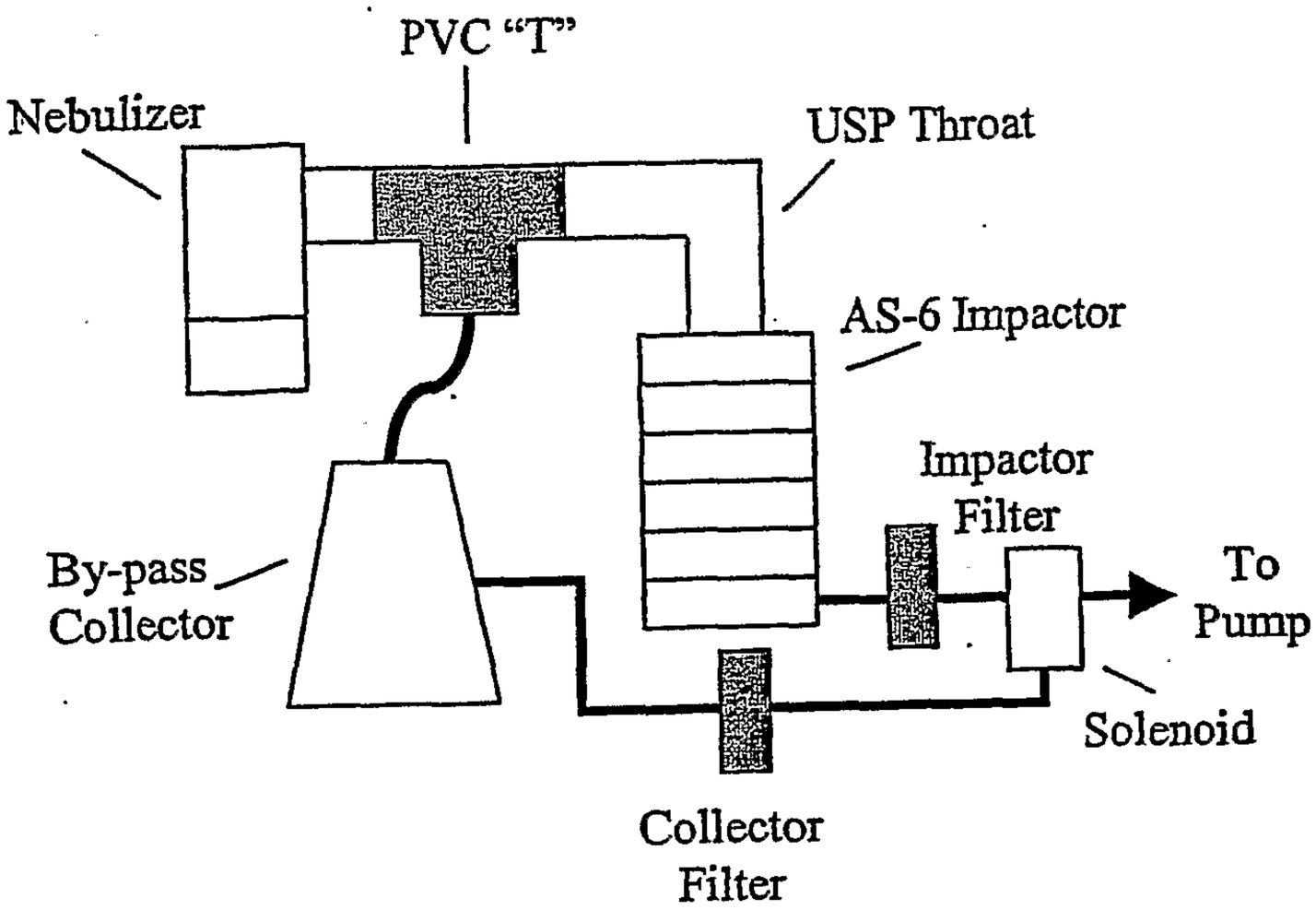

PVC "T"
Nebulizer
USP Throat
AS-6 Impactor
Impactor
Filter
By-pass
Collector
To
Pump
Solenoid
Collector
Filter

Fig. 12

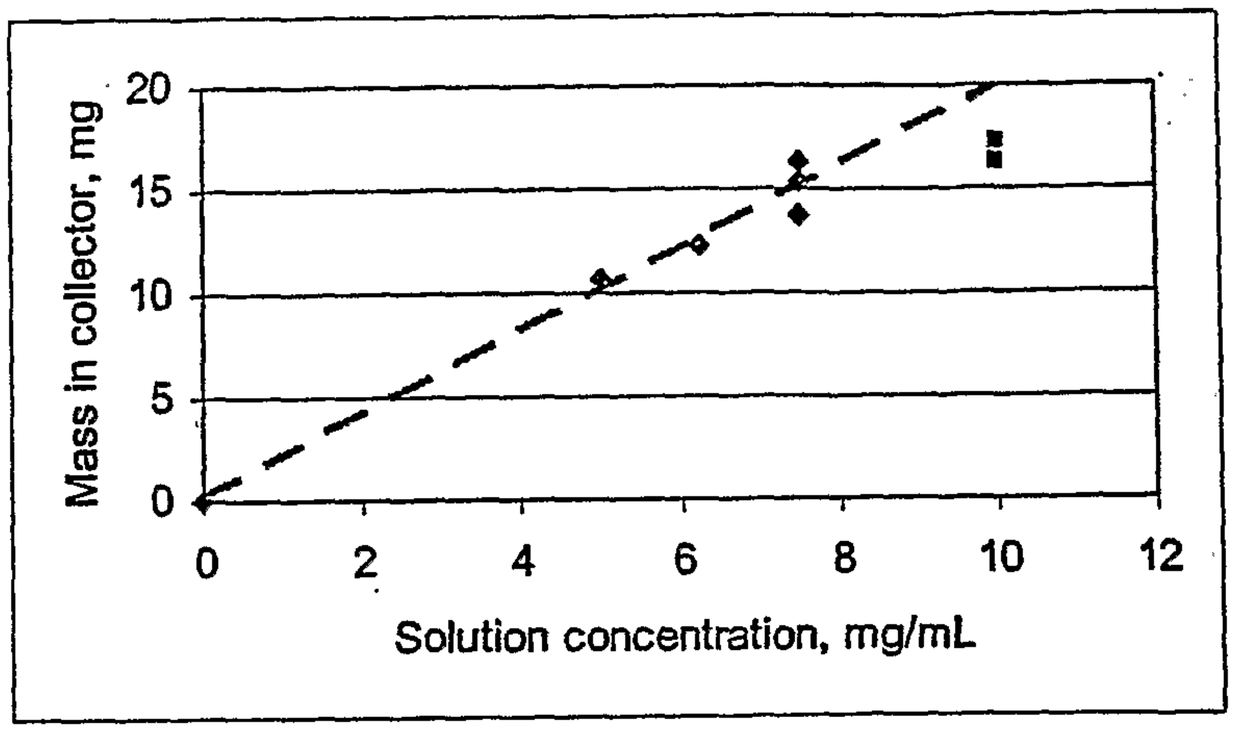

Mass in collector, mg
20
15
10
5
0
0
2
4
6
8
10
12
Solution concentration, mg/mL

Fig. 13

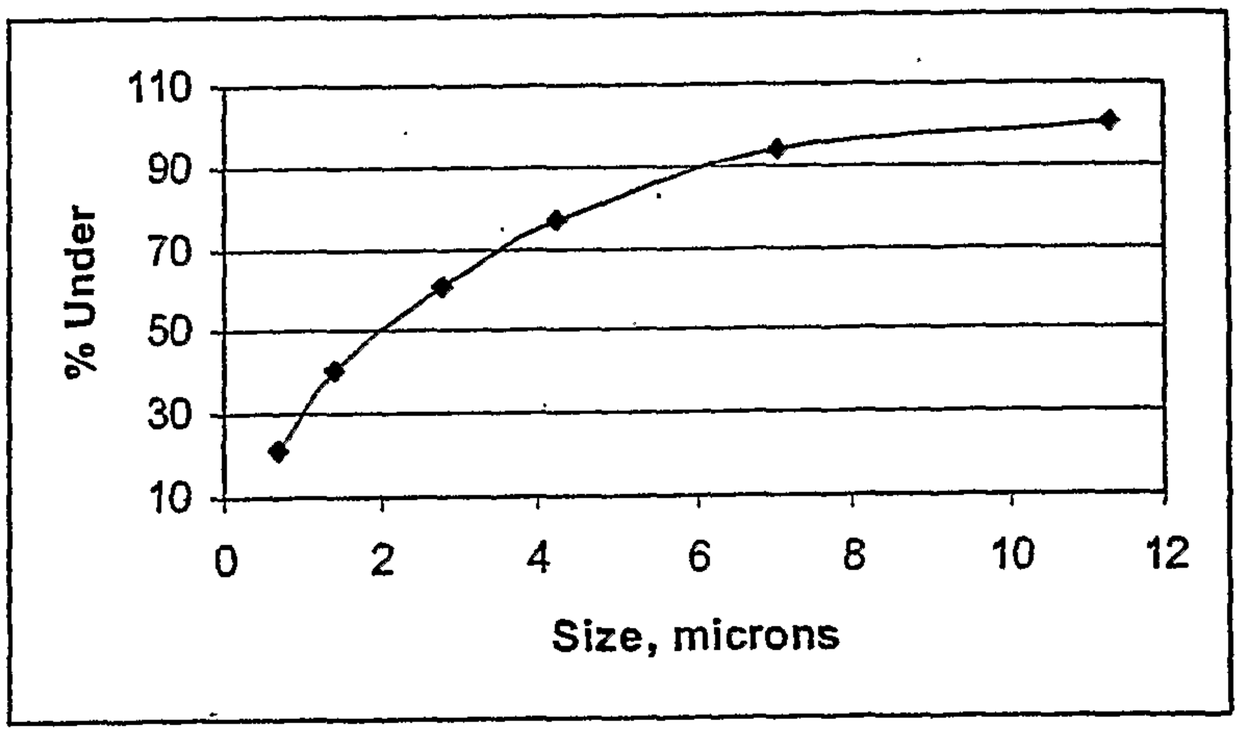

% Under
110
90
70
50
30
10
0
2
4
6
8
10
12
Size, microns

Fig. 14

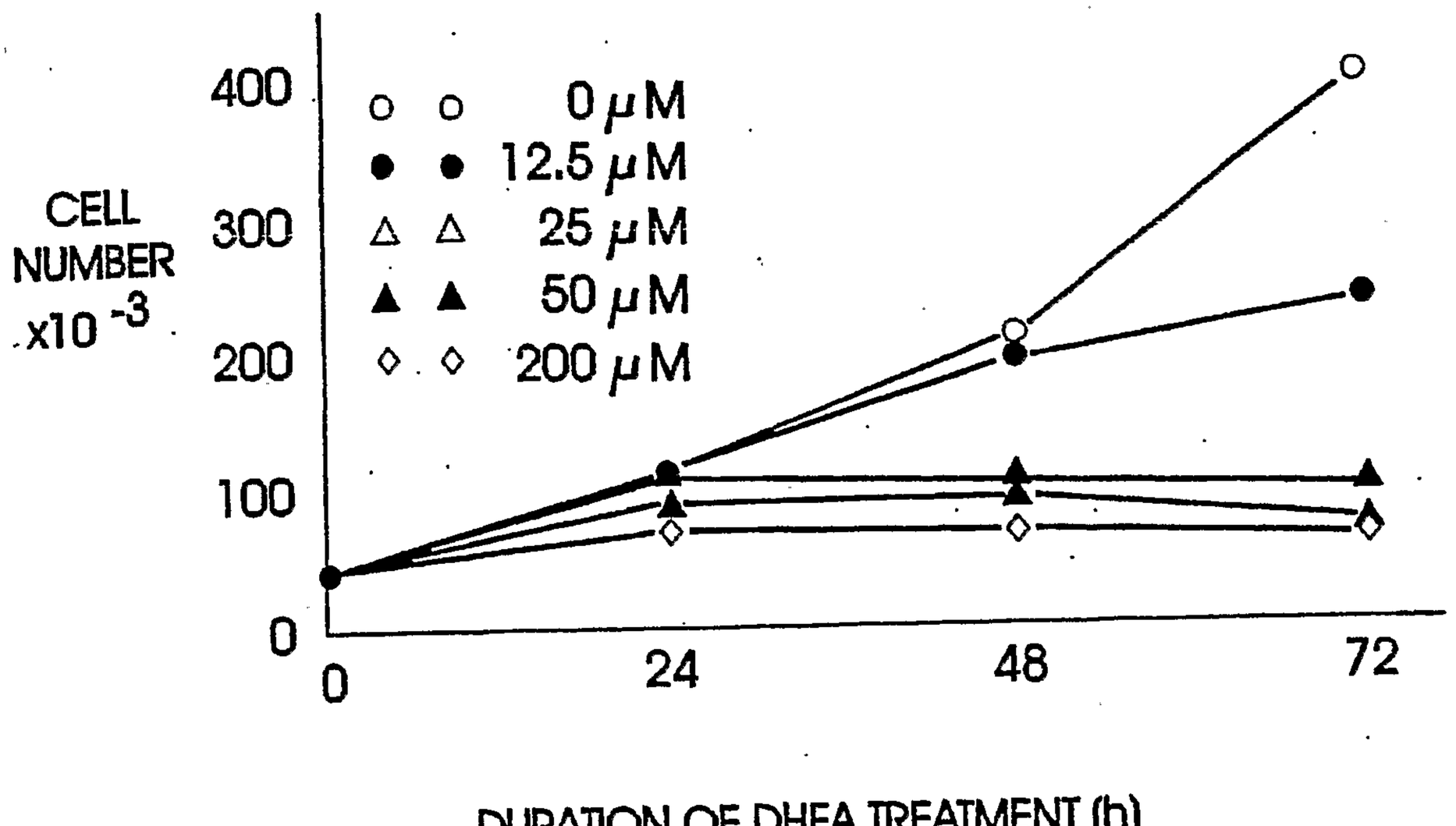

CELL NUMBER x10 -3
400
300
200
100
0
0 μM
12.5 μM
25 μM
50 μM
200 μM
0
24
48
72
DURATION OF DHEA TREATMENT (h)

Fig. 15A

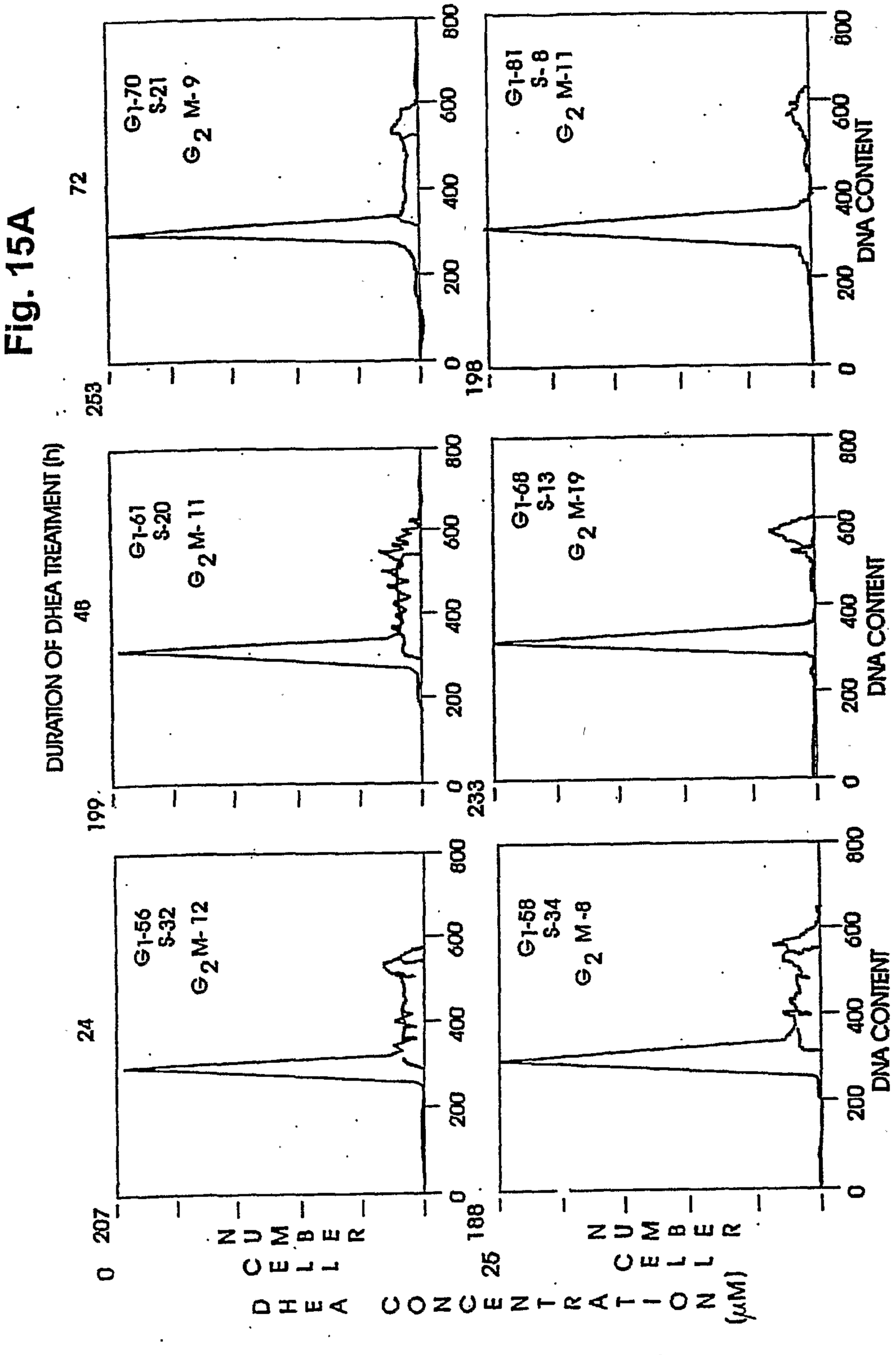

DURATION OF DHEA TREATMENT (h)
72
48
24
0
G1-70
S-21
G2 M-9
253
G1-81
S-8
G2 M-11
198
G1-61
S-20
G2 M-11
233
G1-68
S-13
G2 M-19
188
G1-56
S-32
G2 M-12
207
G1-58
S-34
G2 M-8
199
0
200
400
600
800
DNA CONTENT
NUMBER
CELL
25
DHEA CONCENTRATION (μM)

Fig. 15B

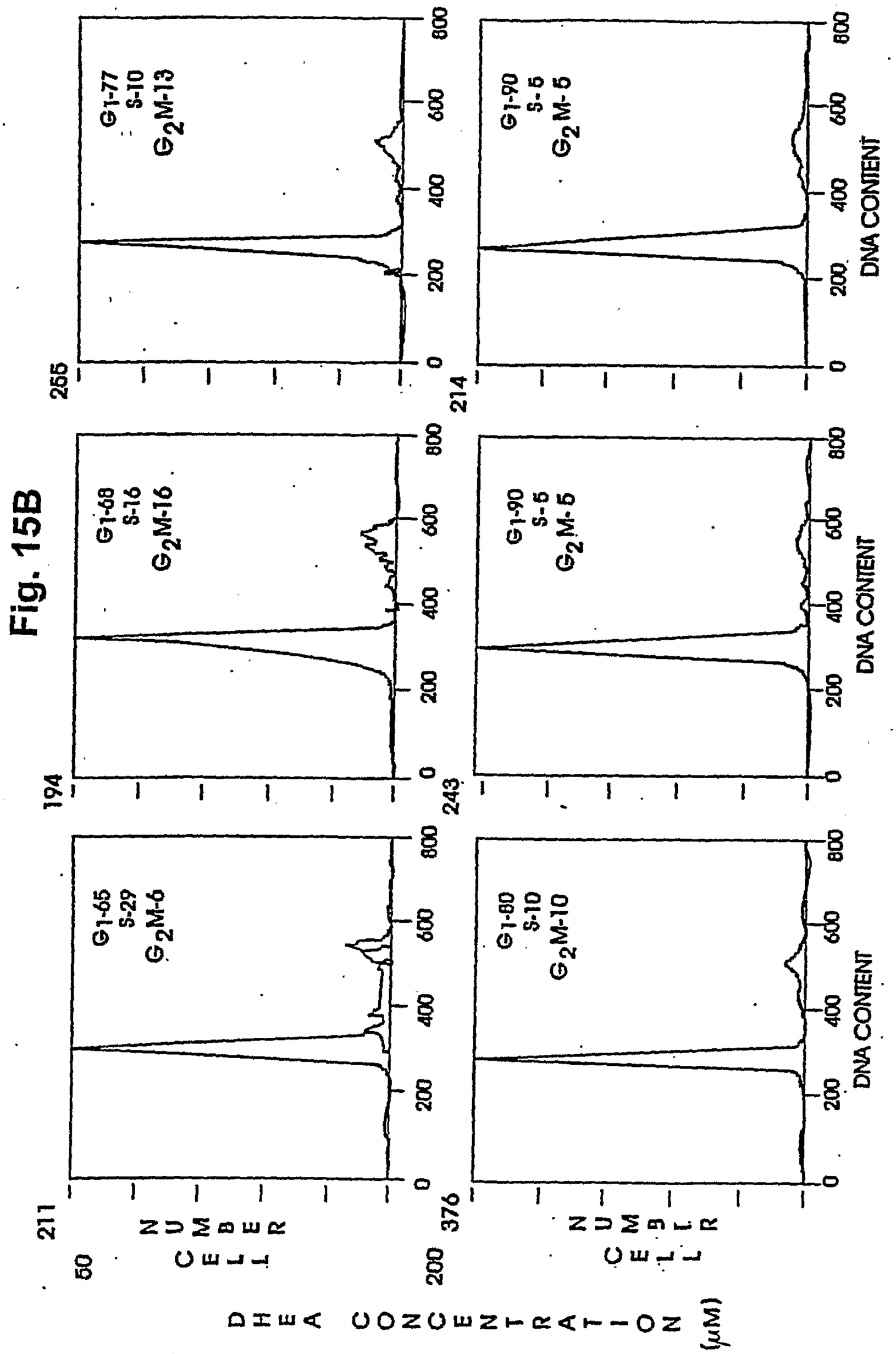

G1-77
S-10
G2M-13
255
G1-90
S-5
G2M-5
214
G1-68
S-16
G2M-16
194
G1-90
S-5
G2M-5
243
G1-65
S-29
G2M-6
211
G1-80
S-10
G2M-10
376
0
200
400
600
800
DNA CONTENT
NUMBER CELLS
DNA CONCENTRATION (μM)

Fig. 16A

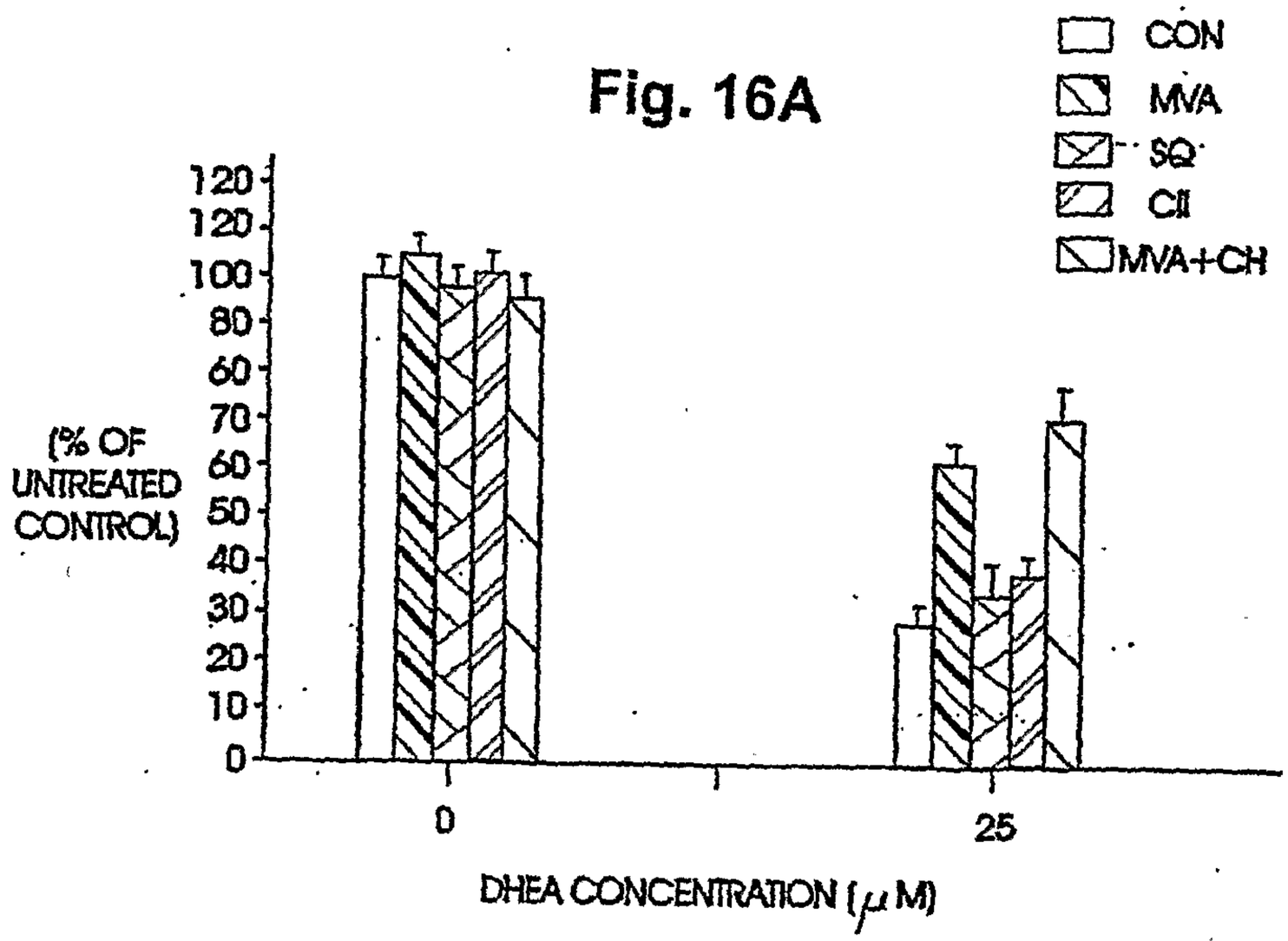


Fig. 16B

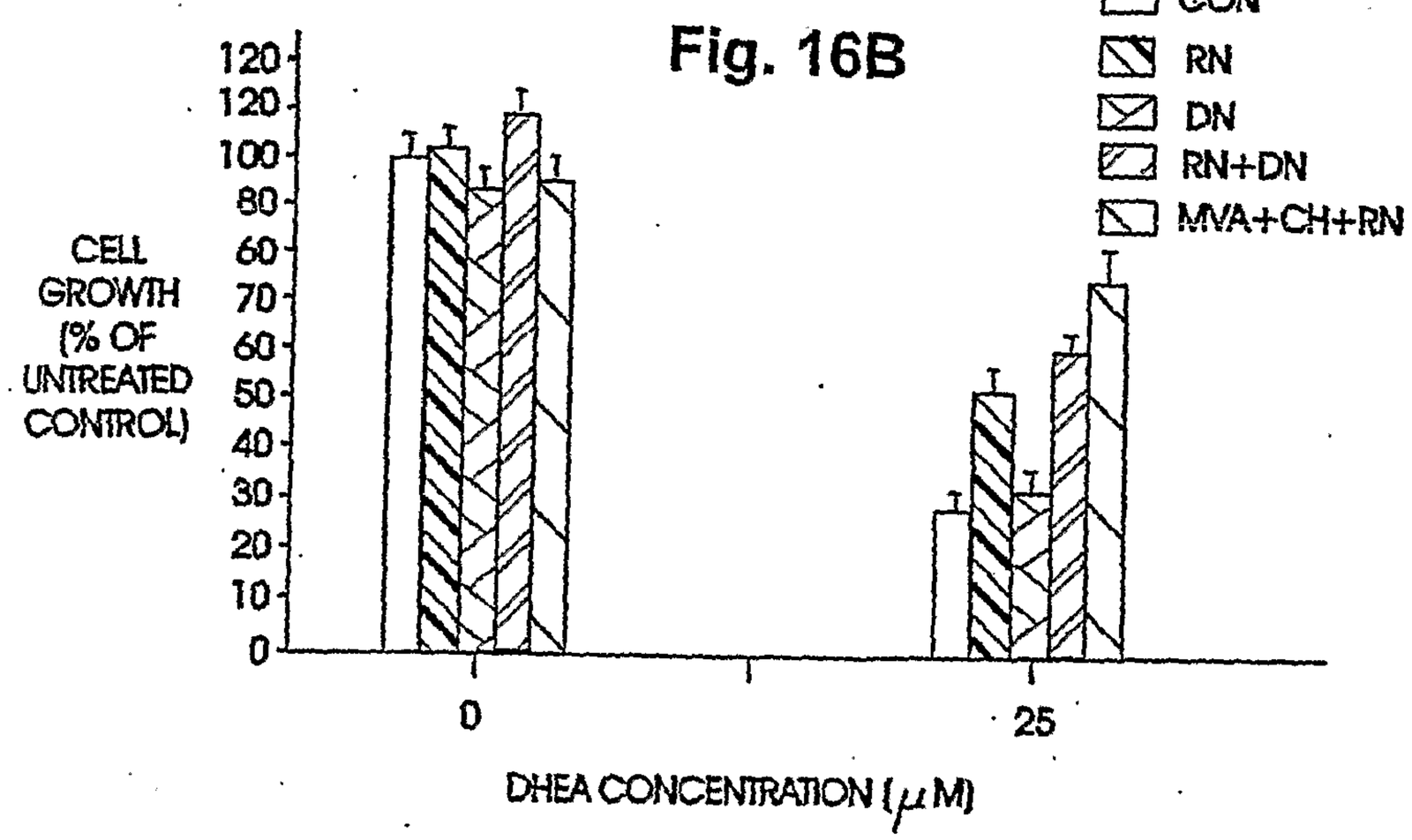


Fig. 17A

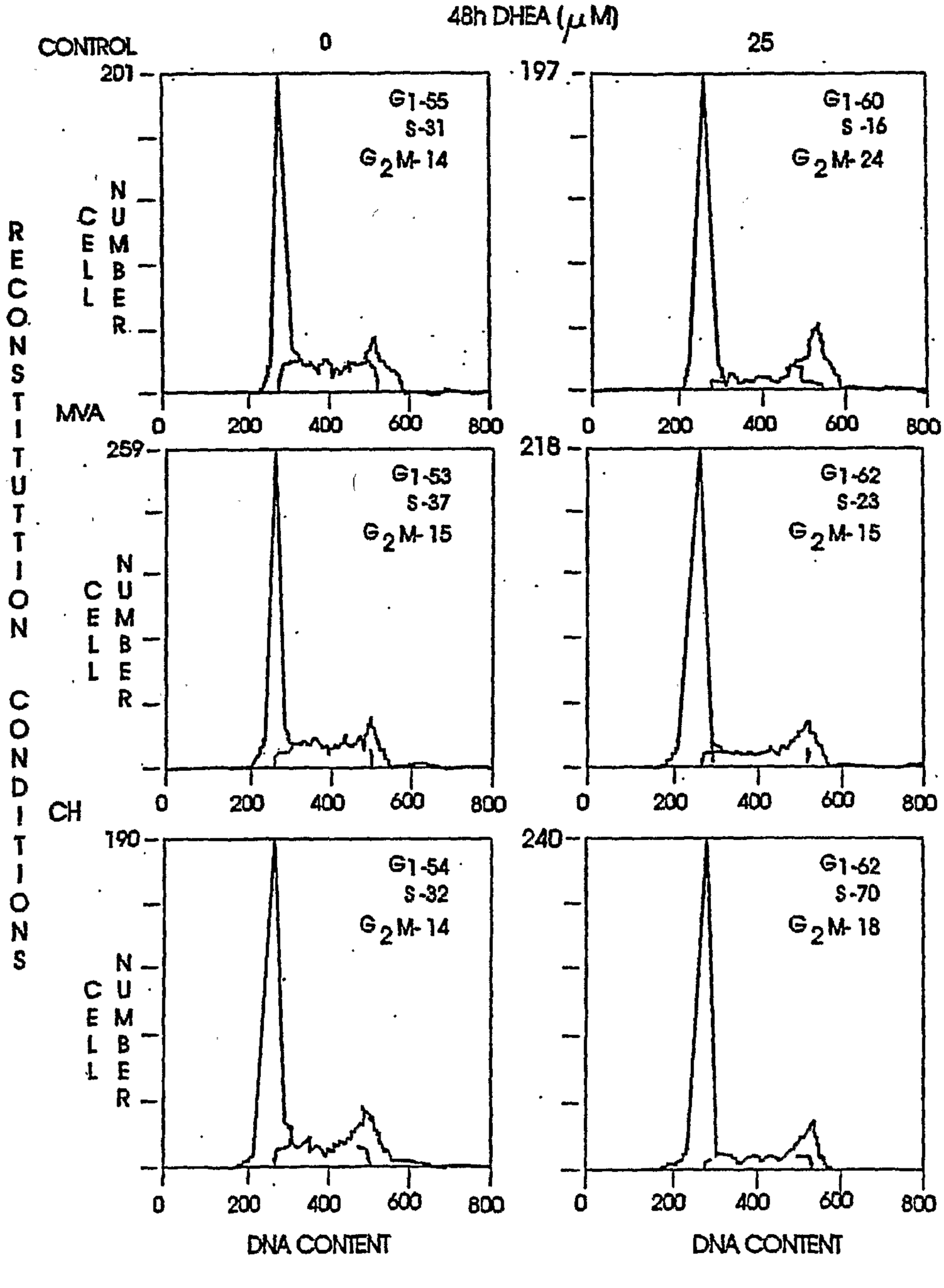

DURATION OF TREATMENT
48h DHEA (μM)
CONTROL
0
25
RECONSTITUTION CONDITIONS
CH
MVA
CELL NUMBER
201
G1-55
S-31
G2M-14
197
G1-60
S-16
G2M-24
259
G1-53
S-37
G2M-15
218
G1-62
S-23
G2M-15
190
G1-54
S-32
G2M-14
240
G1-62
S-70
G2M-18
0
200
400
600
800
DNA CONTENT

Fig. 17B

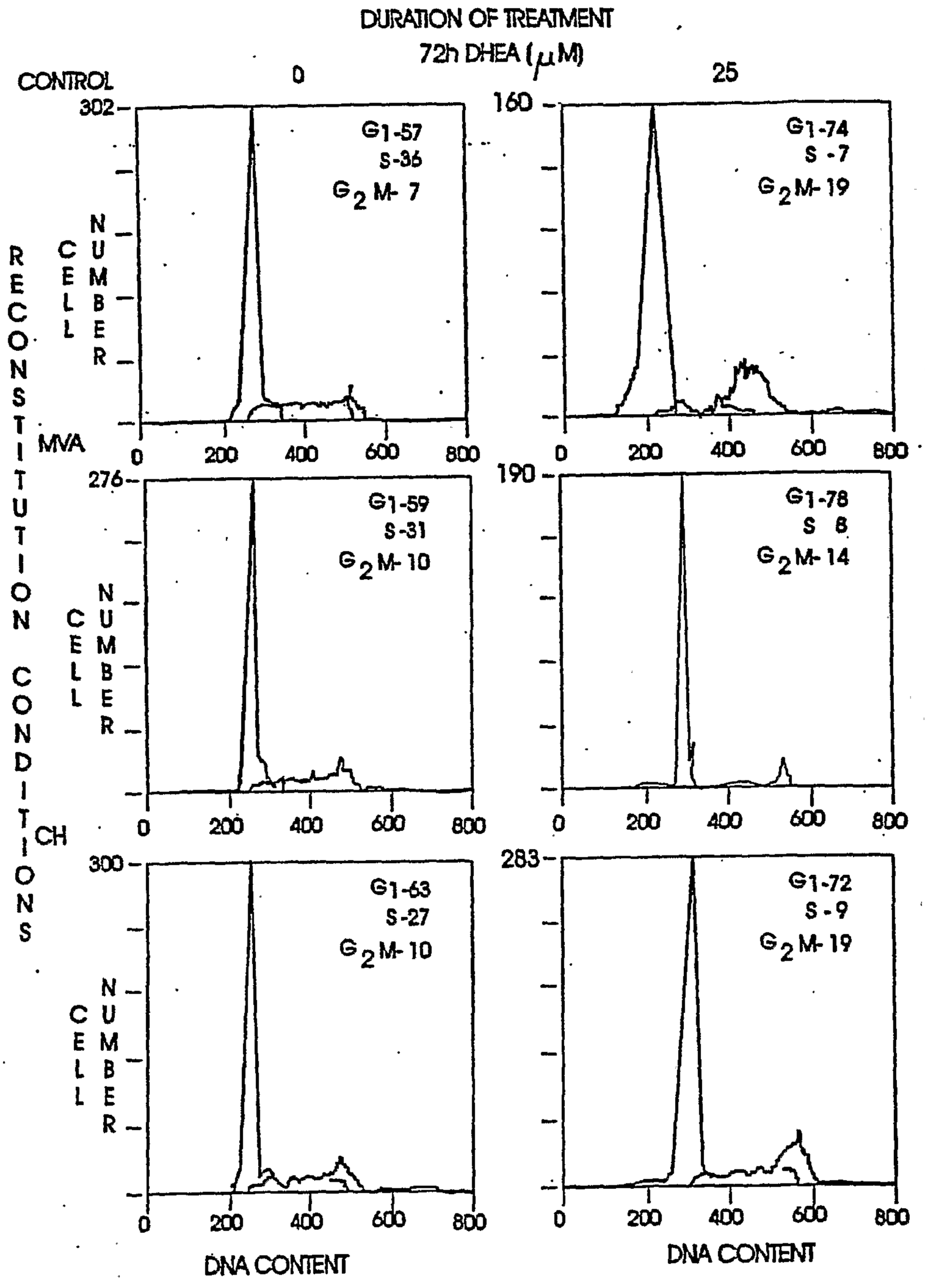

DURATION OF TREATMENT
72h DHEA (μM)
CONTROL
0
25
RECONSTITUTION CONDITIONS
CELL NUMBER
MVA
302
G1-57
S-36
G2M-7
160
G1-74
S-7
G2M-19
276
G1-59
S-31
G2M-10
190
G1-78
S 8
G2M-14
CH
300
G1-63
S-27
G2M-10
283
G1-72
S-9
G2M-19
0
200
400
600
800
DNA CONTENT

Fig. 17C

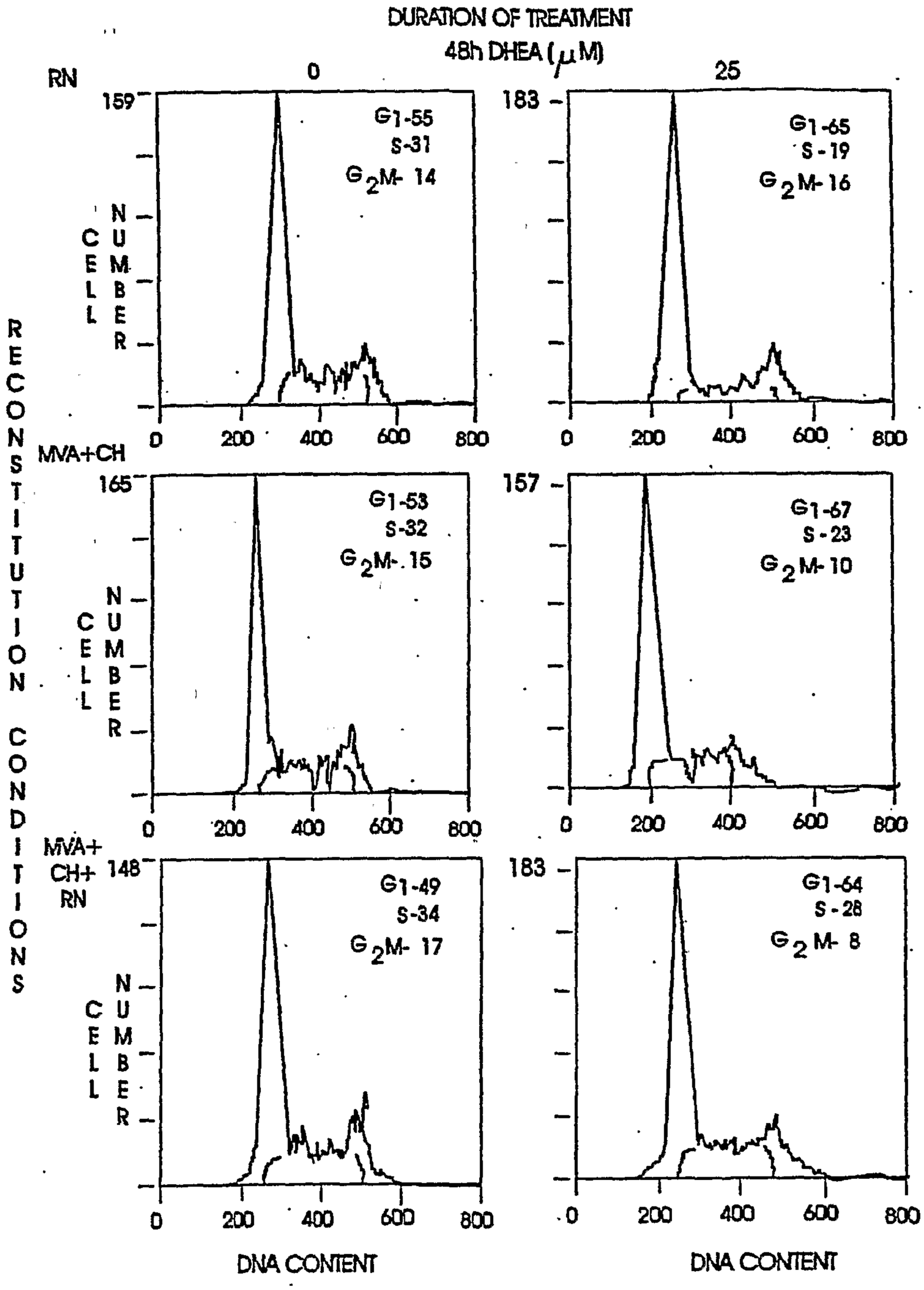

DURATION OF TREATMENT
48h DHEA (μM)
0
25
RN
159
183
G1-55
S-31
G2M- 14
G1-65
S-19
G2M-16
CELL NUMBER
RECONSTITUTION CONDITIONS
MVA+CH
165
157
G1-53
S-32
G2M-15
G1-67
S-23
G2M-10
MVA+
CH+
RN
148
183
G1-49
S-34
G2M- 17
G1-64
S-28
G2M- 8
0
200
400
600
800
DNA CONTENT

Fig. 17D

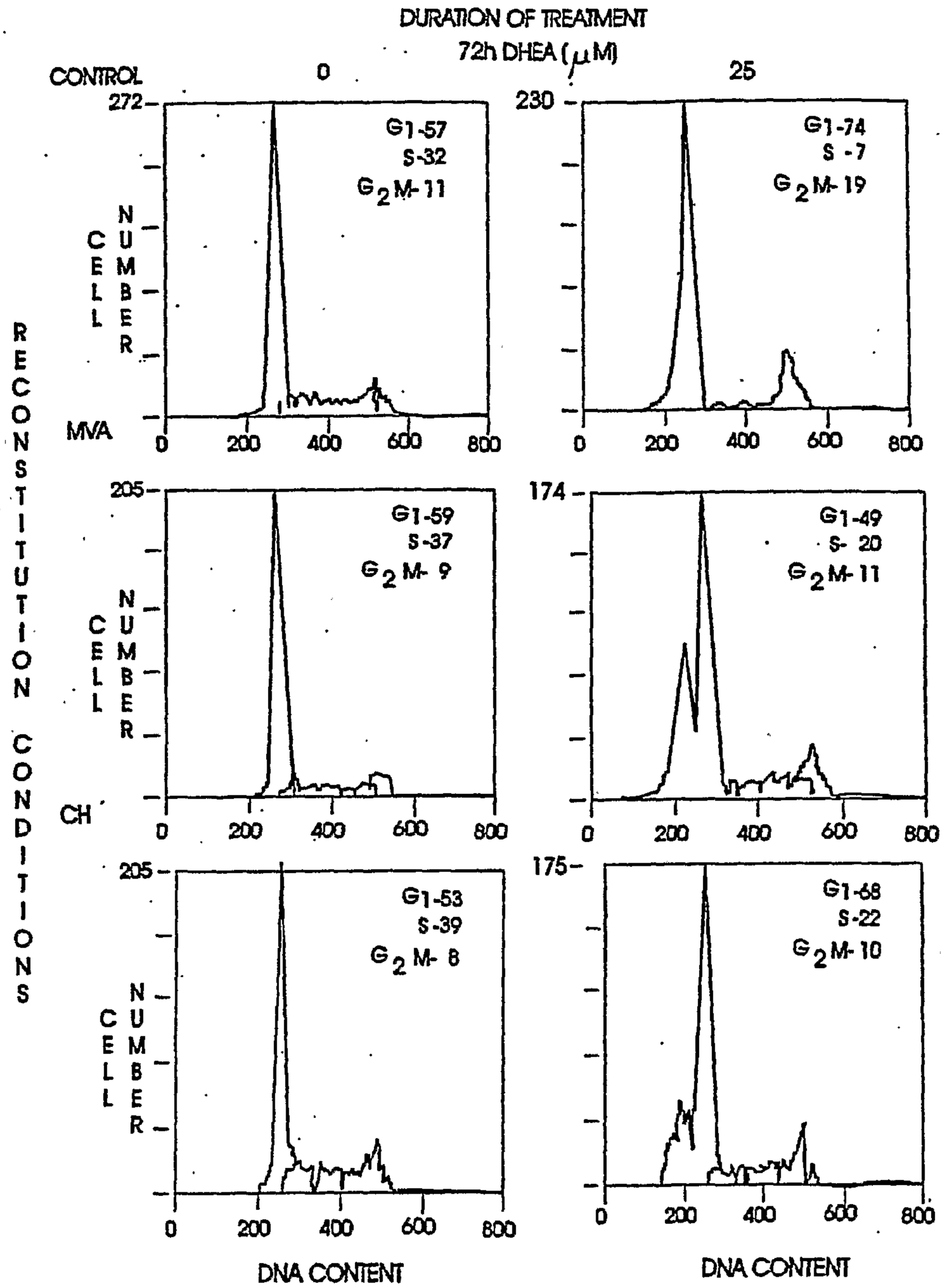

DURATION OF TREATMENT
72h DHEA (μM)
CONTROL
0
25
RECONSTITUTION CONDITIONS
CELL NUMBER
MVA
CH
272
230
205
174
205
175
G1-57
S-32
G2M-11
G1-74
S -7
G2M-19
G1-59
S-37
G2M- 9
G1-49
S- 20
G2M-11
G1-53
S-39
G2M- 8
G1-68
S-22
G2M-10
0
200
400
600
800
DNA CONTENT

Fig. 18

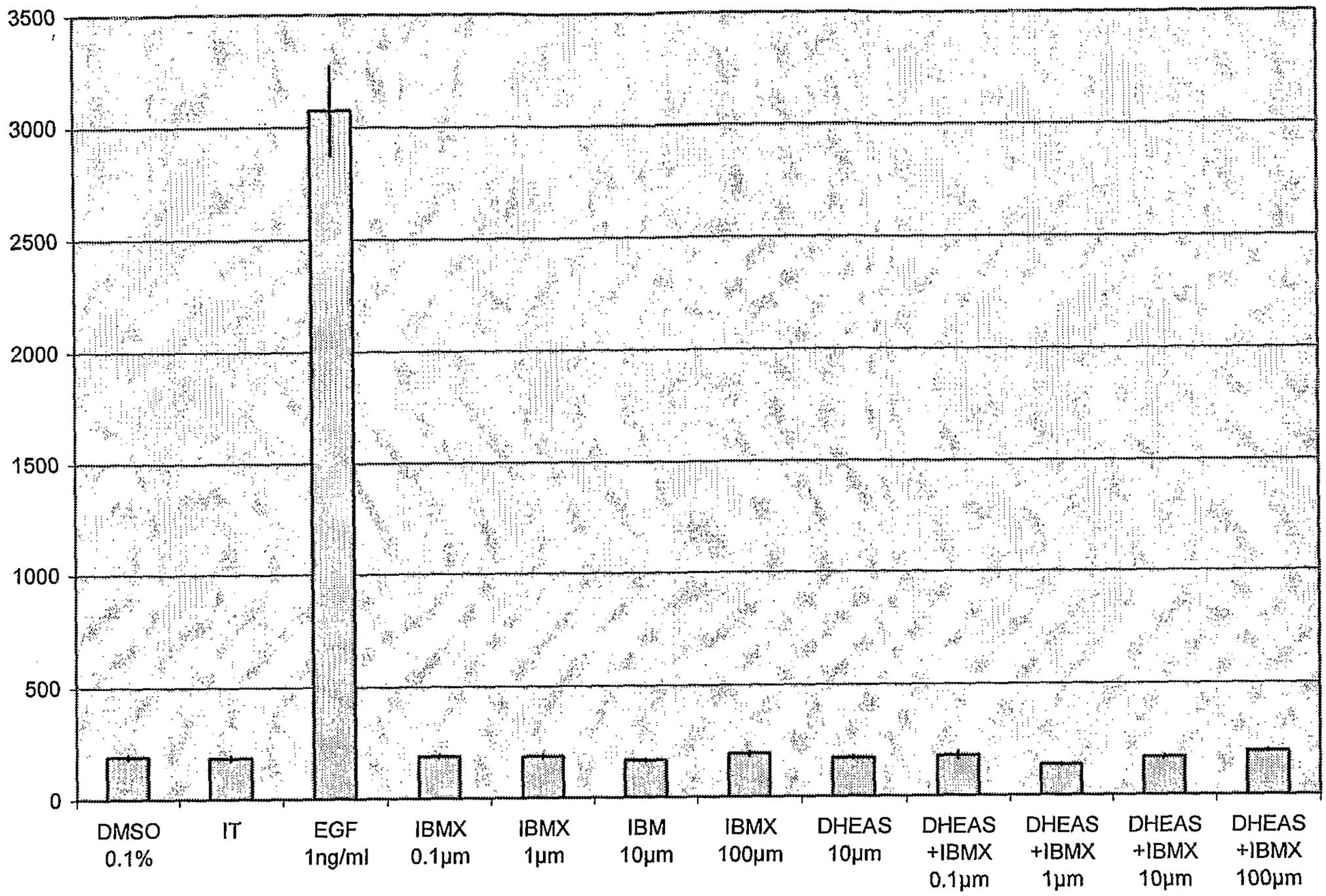

3500
3000
2500
2000
1500
1000
500
0
DMSO 0.1%
IT
EGF 1ng/ml
IBMX 0.1µm
IBMX 1µm
IBM 10µm
IBMX 100µm
DHEAS 10µm
DHEAS +IBMX 0.1µm
DHEAS +IBMX 1µm
DHEAS +IBMX 10µm
DHEAS +IBMX 100µm

Fig. 19

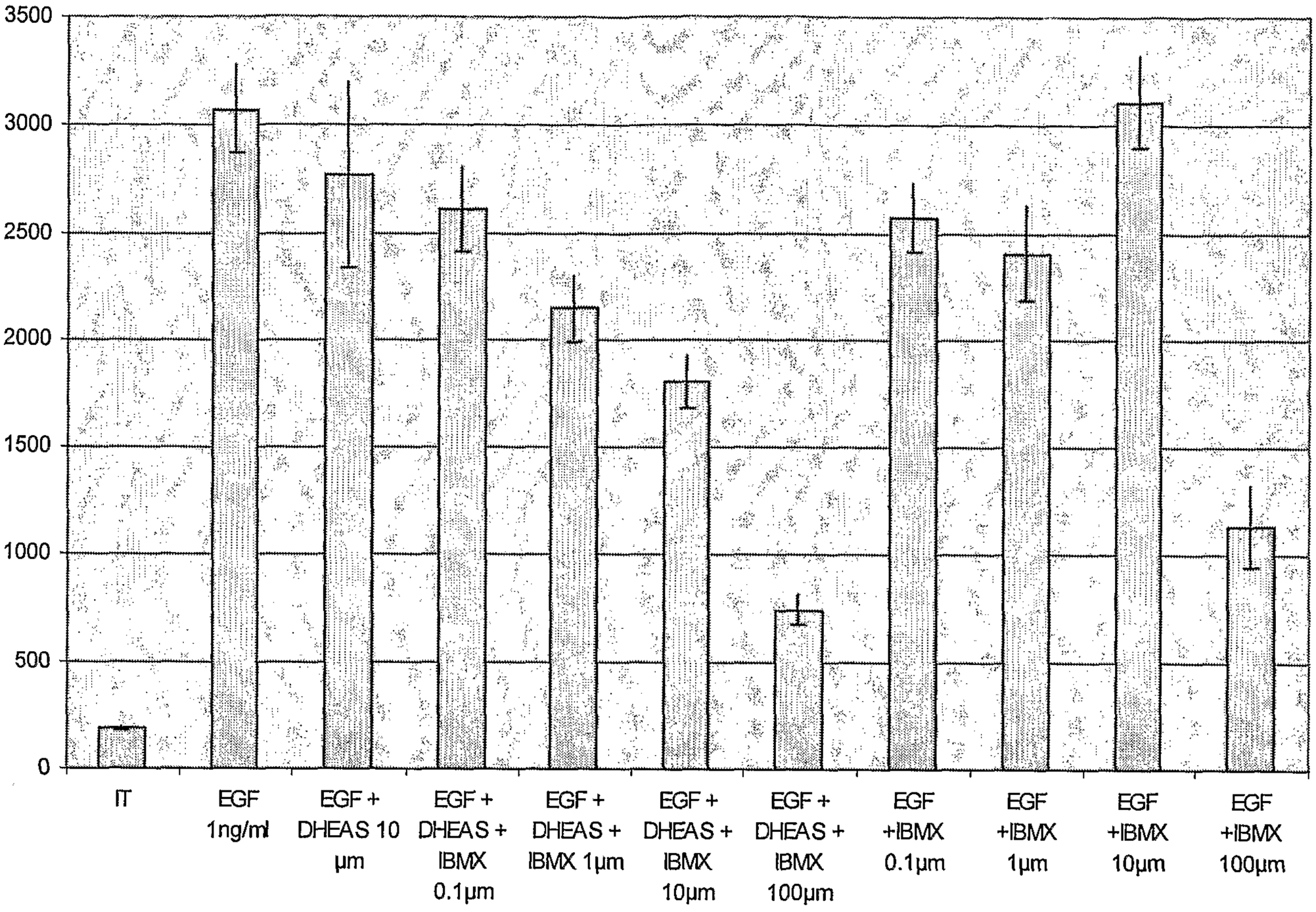

3500
3000
2500
2000
1500
1000
500
0
IT
EGF 1ng/ml
EGF + DHEAS 10 µm
EGF + DHEAS + IBMX 0.1µm
EGF + DHEAS + IBMX 1µm
EGF + DHEAS + IBMX 10µm
EGF + DHEAS + IBMX 100µm
EGF +IBMX 0.1µm
EGF +IBMX 1µm
EGF +IBMX 10µm
EGF +IBMX 100µm

Fig. 20

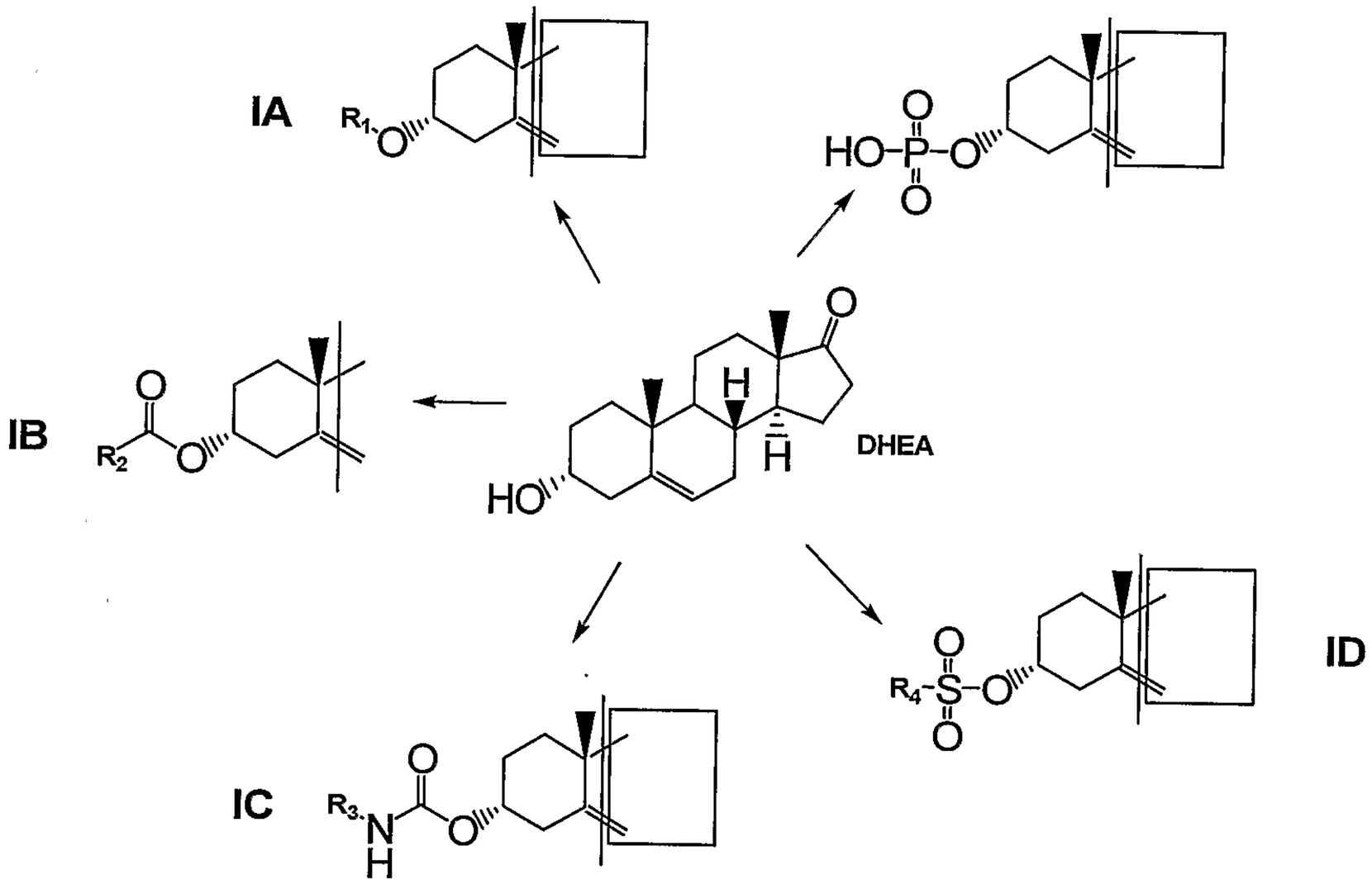


Fig. 21

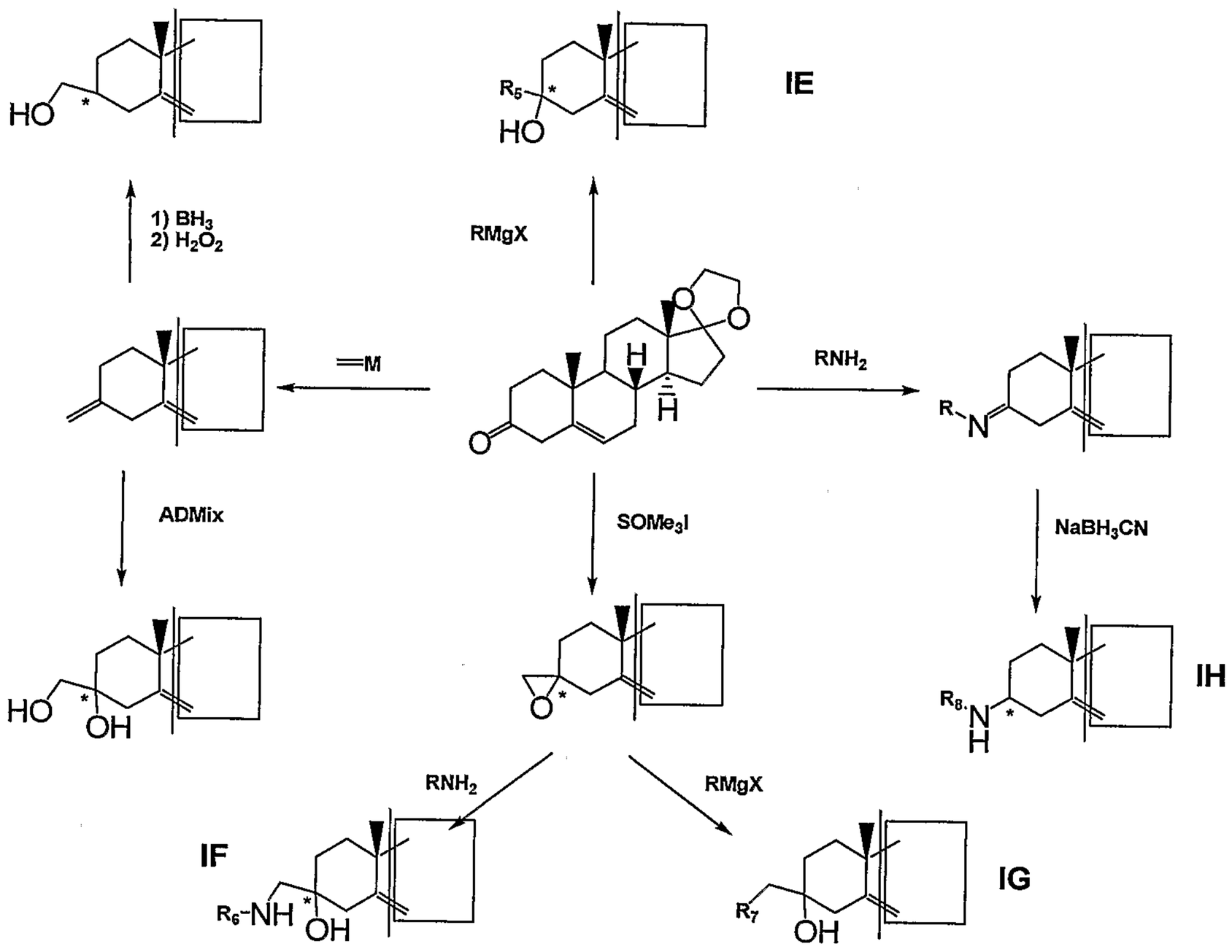

HO
*
R5
*
HO
IE
1) BH3
2) H2O2
RMgX
=M
RNH2
R
N
ADMix
SOMe3I
NaBH3CN
HO
*
OH
O
*
R8
N
H
*
IH
RNH2
RMgX
IF
R6-NH
*
OH
R7
OH
IG

Fig. 22

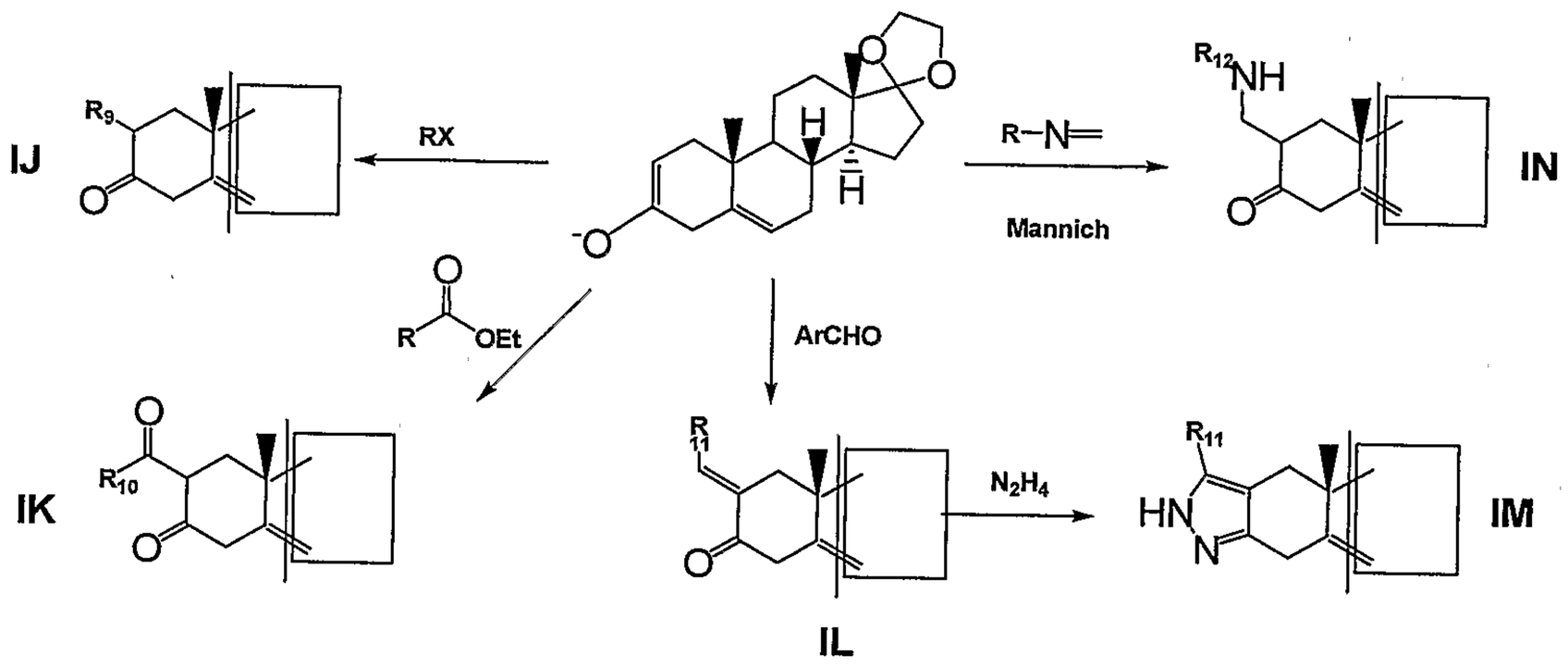
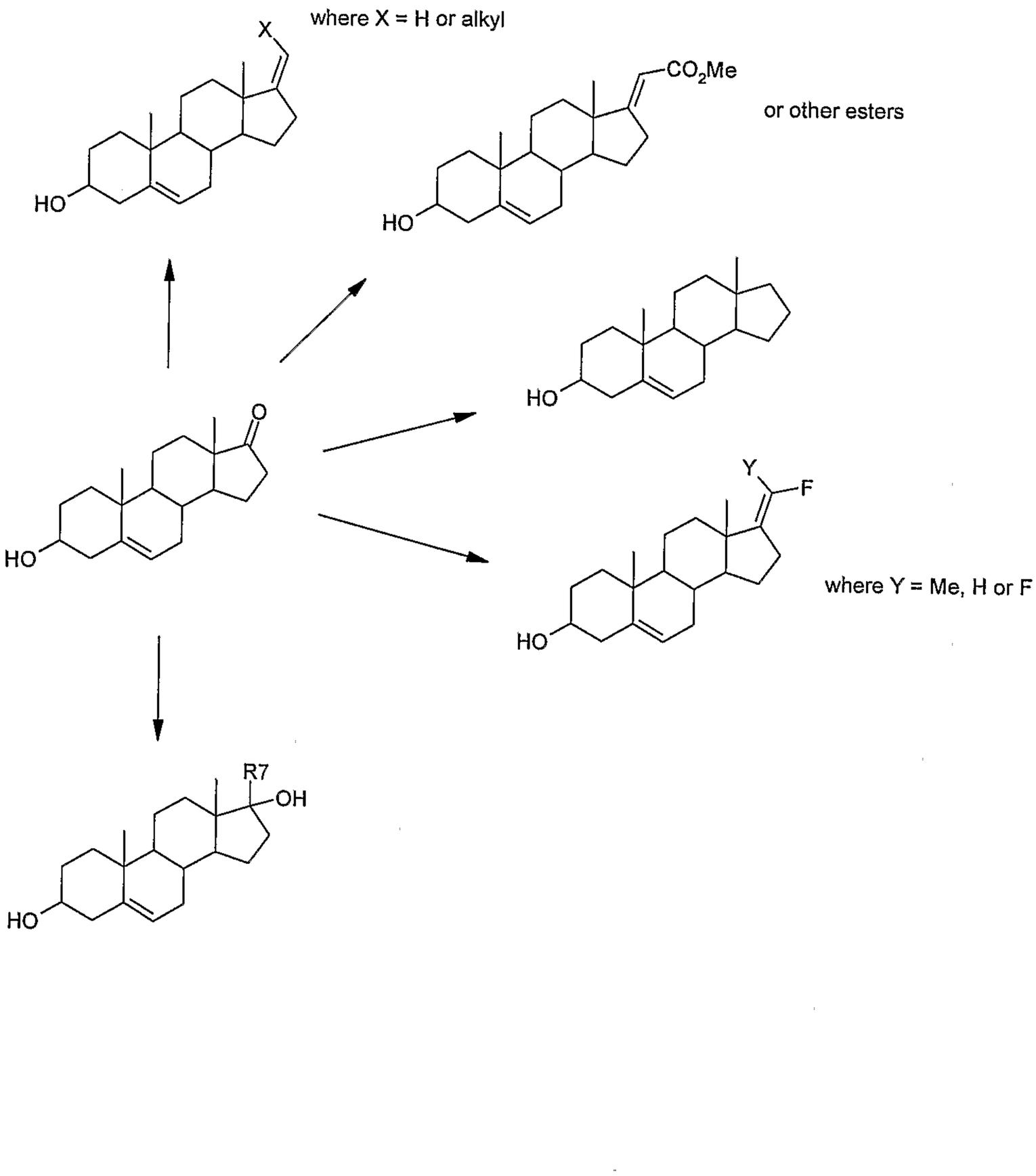
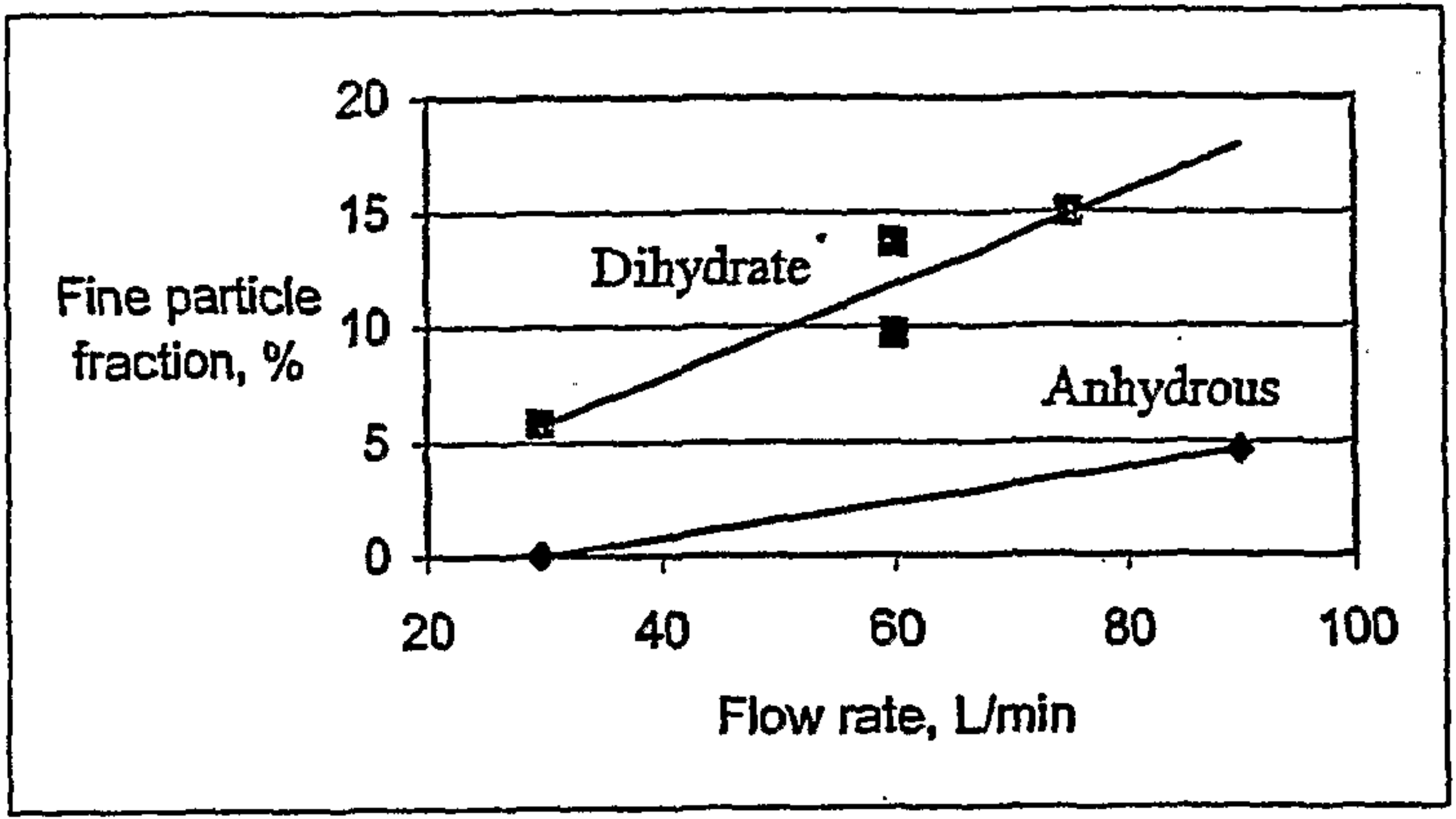

IJ
R9
RX
H
H
O
O
O
R12NH
R–N=
Mannich
IN
R
OEt
ArCHO
IK
R10
O
O
R11
N2H4
HN
N
R11
IM
IL

Fig. 23


where X = H or alkyl
X
HO
CO_2Me
or other esters
HO
HO
O
HO
Y
F
where Y = Me, H or F
HO
R7
OH
HO


20
15
10
5
0
Fine particle fraction, %
Dihydrate
Anhydrous
20
40
60
80
100
Flow rate, L/min