~~COMPOUND COMPOSITION FOR INHALATION USED FOR TREATING~~ ASTHMA

[0001] This application claims the priority of Chinese Patent Application 201010502339.3, filed on September 28, 2010, which is incorporated by reference as if fully set forth herein in its entirety.

[0002] The present invention relates to a pharmaceutical composition of compounds, in particular to the composition which contains an inhaled corticosteroid combined with a short (rapid) acting beta-2 agonist.

[0003] Inhaled medicines for treating Asthma or Chronic Obstructive Pulmonary Disease, COPD, is preferred than oral dosage form product because they delivers medication directly to the target acting site, hence the given dose is minimized and the systemic side effects are reduced. Most commonly used inhaled medicines include inhaled corticosteroids, beta-2 agonists, anticholinergics, or combinations of the above medicine.

[0004] Commercially available inhaled corticosteroid includes Fluticasone propionate, Budesonide, Ciclesonide, Momethasone furoate, Beclomethasone dipropionate, Triamcinolone acetonide, and Tipredane, etc. Due to the anti-inflammatory action, they were used as controller in asthma or COPD treatment either administer alone or combined with long acting beta-2 agonist (LABA).

[0005] Commercially available beta-2 agonists include Albuterol sulfate (also called Salbutamol sulphate), Procaterol hydrochloride, Fenoterol hydrocromide, Reproterol hydrochloride, Formoterol, Terbutaline sulphate, and Salmeterol xinafoate. Both Formoterol and Salmeterol are long acting beta-2 agonists (LABA) with duration of action up to twelve hours. In practice, LABA is suggested to be combined with inhaled corticosteroid and given twice daily to act as controller in patients with moderate to severe Asthma. The reason why LABA is not suggested to be used alone in asthma is that it will develop tolerance through beta-2 receptors down regulation which will results reduced efficacy or even higher

exacerbation rate or deaths under routine treatment if used alone. While inhaled corticosteroid could reduce beta-2 receptors' down regulation, consequently reduce the extent of beta-2 agonists' tolerance, hence was suggested to combine with LABA as controller for asthma's long term treatment.

[0006] Fenterol, Aalbuterol, Terbutaline, or Procaterol are short acting beta-2 agonist (SABA) with the duration of action between 3 to 6 hours or 4 to 8 hours. In clinical application, SABA is mainly used alone as needed to relieve the status of acute bronchocontriction, if necessarily, it can be used thrice to four times a day.

[0007] In the prior art, there are several related patent allocation, such as: 1). the Taiwan application No. 200303767 discloses that the superfine formula of Formoterol contains 0.003-0.192% w/v of (R,R)-(±)-Formoterol fumarate; 2). the Taiwan patent No. 329837 discloses a pharmaceutical composition containing Mometasone furoate for treating airway disease and lung disease; 3). the China application No. CN1305380 discloses a composition including Formoterol and Budesonide for applying on preventing and treating acute asthma; 4). the U.S. patent No. 5,972,919 disclose a formulation containing an effective amount Formoterol and Budesonide and the molar ratio of the content is 1:4 to 1:70, 5). the U.S. patents No. 6,932,962, No.6,799,572, No. 6,638,495, No. 6,962,151 and No. 7,321,059 have disclosed a combination about inhaled corticosteroids, which is one selected from Budesonide, Fluticasone, Mometasone, Beclomethasone, or Ciclesonide, etc. and Beta-2 agonist which is one selected from Fenoterol, Albuterol, Procaterol, Salmeterol, or Formoterol, etc. and 6). the U.S. patents No. 7,244,742, No. 7,481,995, and No. 6,596,261 further discloses that the compound of inhaled corticosteroids collocating with beta-2 agonists is added to Anticholinergics of Ipratropium. However, what recited in the above-mentioned references are all about techniques of Pharmaceutical formulation, and it is not what disclosed by the present invention that using combinations of acting time, dosage, and usage to improve clinical performances.

[0008] Austria et al. disclose that a control therapy by adding low or high dosage of inhaled Budesonides to oral Procaterol for treating patients between 7 to 18 year-old and found under regular asthma treated by Procaterol and Budesonide (Chest, 2005) can improve the asthma condition. However, the numbers of patients are too few, and there is no difference between the low dosage and the high dosage. In addition, on April 4, 2010, Japan disclosed that using 15 to 30 mcg inhaled solution of Procaterol and 250 mcg inhaled suspension of Budesonide on a nebulizer twice a week, after one week, switch to use Budesonide alone for treating regular asthma of teenagers. The above two respiratory therapies are using oral beta-2 agonists to combined with inhaled corticosteroids or using a nebulizer to continuously spray these two medicines, and the above-mentioned respiratory therapies are different from the metered dose inhaled combination product or dry-powder inhalation of the present invention. Furthermore, when the drug delivered through nebulizing solution, patients need to inhale the medicine for a period of 10 to 15 minutes, hence, the drug amount given to the patients are much higher than administered from the metered dose inhaler or the Dry-powder inhaler, and in addition they were all given twice daily as control therapy. Therefore, they are different from the present invention in partial polarity control therapy (eccentric therapy) in order to provide a drug blood concentration variation within a day to meet the need of daily lung function circadian rhythm changes.

[0009] Respiratory diseases such as asthma and chronic obstructive pulmonary disease (COPD) are caused by urbanization with environmental pollution problems. Such respiratory diseases have become one of the major diseases of globalization. Most research findings show that the high asthma mortality is related to low diagnosis rates and treatment inadequate. In 1993 World Health Organization and the National Institutes of Health invited experts to discuss control solutions of asthma and organized the Global Initiative for Asthma, GINA, which is an ongoing council organization. Every few years, based on medical evidences, the

hydrofluoroalkanes and alkyl saccharides, HFA; U.S. patent No. 7,244,742 related to adding anti-cholinergic drugs; U.S. patent No. 7,267,813 containing crystalline spherical inhalation particles; U.S. patent No. 7,459,146 related to using for modified polyethylene glycol, PEG, nanoparticles HFA inhalation aerosol propellants. And what mentioned above is not same as the present invention.

[0017] References have also showed that the adding of the inhaled corticosteroid only alleviates part of acute drug tolerance phenomenon of the beta2-agonist. In our research, we also found that when the beta2-agonist was given too high single dose, it will produce an acute drug tolerance. This reveals that there are reasons other than beta2 receptor downregulationto cause the acute tolerance.

[0018] According to the relevant references, the mechanism of beta2-agonist's tachyphylaxis may be related to the exhaust of some endogenous trachea-relaxing substances which mediate trachea relaxation. Such as substantially increased intracellular cyclic adenosine monophosphate (cAMP) concentrations after administration of beta2-agonist. The released cAMP is catalyzed by proteases and generates serial reactions to relax the trachea. Therefore, when given a large amount or high frequency of beta2-agonist, the endogenous trachea-relaxing substances will be exhausted simultaneously. When the internal recovery rate of these substances is less than the consumption, it might induce the acute drug tolerance phenomenon.

[0019] It is also well know that asthma patient's lung functions will change during a day. Due to the neurohormonal circadian change, the lung function is the worst at 04:00 am and is the best at 16:00 pm. Based on the circadian rythm of the lung function and the above-mentioned mediating substance exhausted mechanism, the inventor proposed a new way of therapy to asthma or chronic pulmonary obstruction. After carefully testing and research with a spirit of, and a spirit of perseverance, the inventor eventually has proposed an invention, "Inhaled

~~Combination Product for Asthma." The summary of the present invention is~~ described as follows.

[0020] The present invention provides an inhaled composition combination which contains an effective amount of beta2-agonist and an effective amount of inhaled corticosteroid. When it is necessary, it can include a pharmaceutically acceptable carrier.

[0021] The present invention further provides an inhaled metered dose aerosol or powder inhalator combination which contains an effective amount of beta2-agnoist and an effective amount of inhaled corticosteroid. When it is necessary, it can include a pharmaceutically acceptable carrier.

[0022] According to the above-mentioned aspect of the present invention, the so called beta2-agonist is selected from short or fast acting beta2-agonists such as Albuterol, Fenoterol, Procaterol, terbutaline, Albuterol sulfate, Procaterol HCl, Fenoterol hydrobromide and terbutaline sulphate.

[0023] According to the above-mentioned aspect of the present invention, the corticosteroid is selected from Budesonide, Fluticasone, Mometasone, Ciclesonide, Beclomethasone, Triamcinolone, Tipredane, Fluticasone propionate, Beclomethasone dipropionate, Triamcinolone Acetonide and so on.

[0024] According to the above-mentioned aspect of the present invention, one purpose of the present invention, "Inhaled Combination Product For Asthma," is using for eccentric way control therapy of asthma or chronic pulmonary obstruction diseases. Based on the above-mention circadian change of pulmonary function of asthma

patients, the so called eccentric therapy by giving medicines before go to bed and after wake up, might be a better treatment option for patients' long term treatment. The eccentric treatment gives higher drug concentration during the worse stage of lung function and giving a lower concentration or drug free period during the better lung function stage during a day. Therefore, patients' body will have time to recover and accumulate the endogenous trachea-relaxing substances. In addition, such eccentric therapy could also use the same combination medicine as reliever to a patient with asthma or chronic pulmonary obstruction diseases when it is necessary. The advantage is that the additional inhaled corticosteroid will help to reduced and recover the beta2-agonist acute tolerance and helping the patients to control their disease earlier. Therefore, compared with the current combination control therapy which contains a 12 hours acting drugs (LABA) used twice daily which having the potential to exhaust tracheal relaxing mediator, the eccentric therapy using inhaled corticosteroids plus SABA (ICS+SABA) could be a better treatment choice to preserve patient's lung function.

[0025] In addition, such combination could also be used as needed to improve the current short acting beta2-agonist alone reliever treatment. The reason is that when there are asthma symptom developed and need a bronchodilator, it also means that the tracheal are under inflammation condition. Hence, the addition of inhaled corticosteroids could help reduce the tracheal inflammation and help the patients to recover earlier.

[0026] Other objects, advantages and efficacy of the present invention have been studies through the following experiment with reference to the accompanying drawings, in which:

In order to understand the alleviating effects of the present invention, pharmaceutical composition, compare with that the sensitized mice are given long-active or short-acting beta2-agonist alone (Salmeterol or Procaterol), and compare with the reaction of the normal mice and the control group.

[0034] The results are shown as in Figure 2 to 4, no matter administrating alone or administering in combination of the long-acting Salmeterol or medium-short-acting Procaterol, when giving a single super high dose can both make the sensitized mice to show an acute drug tolerance phenomenon of reducing the ability of trachea relaxing. The result shows that there is a suitable dose range for displaying medical effects. Administering the beta2-agonist twice daily cannot avoid the phenomenon of the acute tolerance, giving the inhaled corticosteroid, Budesonide, under a suitable dose range can help to relieve the acute drug tolerance. However, when using too high dose of the beta2-agnist, Budesonide cannot relieve the acute drug tolerance, and it shows that the dosage of the beta2-agonist has exceeded body affordability. These results also indirectly proved that the present invention proposes an idea and assumption that "an acute drug resistance results from exhaustion of endogenously trachea-relaxing substances.

[0035] Relative to the listed suggestions of GINA guidelines, it is suggested to use short-acting beta2-agonist alone as reliever for treating emergent exacerbation of all kind of patients. If swapping the administering way of the short-acting beta2-agonist to another administering way by using a combination inhaler of the short-acting beta2-agonist with the inhaled corticosteroid, i.e., using the present

invention which is a combination of inhaled corticosteroid with medium-short acting beta2-agonist for all level patients with emergent exacerbations as reliever, it might help the patient's recover from their asthma exacerbations.

[0036] From the fact that, when the step 5 patients under severe exacerbation, they need to use additional oral corticosteroid to control their acute asthma exacerbations, we can see that the patients may require additional corticosteroids to help patients to restore the relaxing functions of trachea by control the inflammation, counteract the beta2-receptor down regulation and help to restore the level of tracheal relaxing mediator.

[0037] Regarding to the related researches, only the reference which sponsored by Chiesi published on New England Journal of Medicine 2007; 356: 2040-2052 involves a research of inhaled corticosteroids with rapidly short-acting beta2-agonist. The result indicates that after six months treatment in mild asthma patients, the as needed(which means does not give drugs until exacerbations of asthma) administration of combination of Beclomethasobne and Albuterol, shows that, compared with using Albuterol alone, the disease status is controlled better, acute exacerbation and hospitalizations were also reduced. Although this study only discussed the situation in patients with mild asthma, and not in moderately to severe patients under the conventional control medicine, it revealed that when the patients have exacerbations, beside the beta2-agonist, giving more corticosteroid is helpful for controlling the condition of disease.

[0038] Therefore, the present invention which is a compound of corticosteroid with rapidly short-acting beta2-agonist which can offer a better choice for using as an reliever for treating acute bronchial constriction of asthma in all levels patents. Such extra corticosteroids given can provide extra anti-inflammatory effect to assist patients to relieve from disease condition earlier.

[0039] In accordance with composition of the formula of the present embodiment, medium-short acting beta2-agonist are used, such as Procaterol, Fenoterol, Terbutaline, Albuterol and its base drugs, which has bronchodilating actions of 6 to 8 hours. If the above medicine was administrated 3 to 4 times a day, the phenomenon of acute tolerance will be produced rapidly, and the duration of broncodilating effect will be reduced to 3 to 4 hours. Use the medium-short acting beta2-agonist together with an inhaled corticosteroid, such as budesonide, and adapting an eccentric way of before sleeping and after awake can restore duration of action back to 6 to 8 hours. Therefore, the present invention can help the long-term control of patients' disease and reducing the acute exacerbation.

[0040] The beta2-agonist with 3 to 8 hours duration, such as Procaterol HCl, Procaterol, Albuterol, Albuterol sulfate, Fenoterol, and Fenoterol hydrobromide etc. was use to combined with inhaled corticosteroid, such as Budesonide, Fluticasone, Beclomethasone, Mometasone, Ciclesonide, Triamcinolone and their base morphology, generate a quantitative combination metered dose inhaler or a dry powder inhaler. The weight ratio of the beta2-agonist and the inhaled

corticosteroid is about 1:2 w/w% to 1:70 w/w%, and the better range is about 1:5 w/w% to 1:60 w/w%.

[0041] The clinical usage is giving the combination drug to asthma patients as controller before sleeping and after waking up. The advantage is to provide patients' body a low dose period to recover trachea relaxing mediators for the next day treatment during their better lung function period in the afternoon,. Hence, it reduces the chance of acute tolerance of the beta2-agonist and help patients to control their conditions stably.

[0042] Compared with using short-acting beta2-agonist (Procaterol, Albuterol, Terbutaline, Fenoterol and its base drugs) alone, the combination has additional anti-inflammatory inhaled corticosteroid. The inhaled corticosteroid provided the anti-inflammatory and mitigation of the acute tolerance resulting from overuse of beta2-agonist, it can help to relieve conditions and reduce numbers of severe exacerbations.

[0043] The present invention which is an eccentric treatment of the combination of inhaled beta2-agonist with corticosteroid and can be administered via HFA MDI or DPI (dry powder inhaler). The above-mentioned pharmaceutically acceptable vectors are provided as an excipient system for need of preparing the combination product of the beta2-agonist with the corticosteroid, so it makes the administered animals or humans not to have adverse reactions, allergies or other non-appropriate responses. Carriers or an excipient system can also include the proper amount of surface active agents, solvents, suspending agents and propellants for stabilizing prescriptions. The formulation of HFA

MDI used in the present invention is usually adopted 1,1,1,2-tetrafluoroethane (Tetrafluoroethane, HFA 134a, HFC 134a) or 1,1,1,2, 3,3,3 - Heptafluoropropane (Heptafluoro-n-propane, HFC 227ea, HFC 227, HFA 227), and depending on necessity, it can also be a mixture formulation of HFA 134a and HFA 227. Dry powder inhalation could be a single dose inhaler or multiple dose inhaler composed of carrier-free active pharmaceutical ingredients or use lactose as a carrier.

[0044] Experimental methods:

[0045] Balb/c mice are divided into normal control group, asthma control group (Ova control group), and each group which is contained 2 to 28 mice (in the pre-exploration stage, the number of the mice is fewer, and in the post stage after confirming dose, the number of the mice for repeatedly confirming experiments is more) is supplied with the same water and food. In the asthma control group, at first, the mice are received i.p. with ovalbumin (OVA) with the interval of 10 days as shown in Figure 1, and the mice are made to have allergies, on the 24th days, the blood are collected from the eye sockets of the mice to be examined by IgE Elisa quantification for confirming that the mice have allergies. In the normal control group, the mice are not made to have allergies and are given salt water. In the asthma control group, the mice are also given salt water but not medicine for treating, and 3 days before administration, the mice are continuously given OVA in the nasal to enhance allergies.

[0046] The mice of each group are divided into subgroups by different medicine and different dose. Each subgroup is divided into single administration within 7 days (one dose), once administration a day for 7 days (qd) or twice administrations a day (bid) for 7 days. The salt water or mixture drug of Procaterol mixing with Budesonide in the salt water is directly administered in the throats of the mice.

COMPOUND COMPOSITION FOR INHALATION USED FOR TREATING ASTHMA

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[0005] Commercially available beta-2 agonists include Albuterol sulfate (also called Salbutamol sulphate), Procaterol hydrochloride, Fenoterol hydrocromide, Reproterol hydrochloride, Formoterol, Terbutaline sulphate, and Salmeterol xinafoate. Both Formoterol and Salmeterol are long acting beta-2 agonists (LABA) with duration of action up to twelve hours. In practice, LABA is suggested to be combined with inhaled corticosteroid and given twice daily to act as controller in patients with moderate to severe Asthma. The reason why LABA is not suggested to be used alone in asthma is that it will develop tolerance through beta-2 receptors down regulation which will results reduced efficacy or even higher

exacerbation rate or deaths under routine treatment if used alone. While inhaled corticosteroid could reduce beta-2 receptors' down regulation, consequently reduce the extent of beta-2 agonists' tolerance, hence was suggested to combine with LABA as controller for asthma's long term treatment.

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[0009] Respiratory diseases such as asthma and chronic obstructive pulmonary disease (COPD) are caused by urbanization with environmental pollution problems. Such respiratory diseases have become one of the major diseases of globalization. Most research findings show that the high asthma mortality is related to low diagnosis rates and treatment inadequate. In 1993 World Health Organization and the National Institutes of Health invited experts to discuss control solutions of asthma and organized the Global Initiative for Asthma, GINA, which is an ongoing council organization. Every few years, based on medical evidences, the

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[0018] According to the relevant references, the mechanism of beta2-agonist's tachyphylaxis may be related to the exhaust of some endogenous trachea-relaxing substances which mediate trachea relaxation. Such as substantially increased intracellular cyclic adenosine monophosphate (cAMP) concentrations after administration of beta2-agonist. The released cAMP is catalyzed by proteases and generates serial reactions to relax the trachea. Therefore, when given a large amount or high frequency of beta2-agonist, the endogenous trachea-relaxing substances will be exhausted simultaneously. When the internal recovery rate of these substances is less than the consumption, it might induce the acute drug tolerance phenomenon.

[0019] It is also well know that asthma patient's lung functions will change during a day. Due to the neurohormonal circadian change, the lung function is the worst at 04:00 am and is the best at 16:00 pm. Based on the circadian rythm of the lung function and the above-mentioned mediating substance exhausted mechanism, the inventor proposed a new way of therapy to asthma or chronic pulmonary obstruction. After carefully testing and research with a spirit of, and a spirit of perseverance, the inventor eventually has proposed

the current techniques." The invention is defined in the independent claim. Some optional features of the invention are defined in the dependent claims.

[0020] There is disclosed an inhaled composition combination which contains an effective amount of beta2-agonist and an effective amount of inhaled corticosteroid. When it is necessary, it can include a pharmaceutically acceptable carrier.

[0021] There is also disclosed an inhaled metered dose aerosol or powder inhalator combination which contains an effective amount of beta2-agnoist and an effective amount of inhaled corticosteroid. When it is necessary, it can include a pharmaceutically acceptable carrier.

[0022] According to the above-mentioned aspect, the so called beta2-agonist is selected from short or fast acting beta2-agonists such as Albuterol, Fenoterol, Procaterol, terbutaline, Albuterol sulfate, Procaterol HCl, Fenoterol hydrobromide and terbutaline sulphate.

[0023] According to the above-mentioned aspect, the corticosteroid is selected from Budesonide, Fluticasone, Mometasone, Ciclesonide, Beclomethasone, Triamcinolone, Tipredane, Fluticasone propionate, Beclomethasone dipropionate, Triamcinolone Acetonide and so on.

[0024] According to the above-mentioned aspect, one purpose of the present techniques is using for eccentric way control therapy of asthma or chronic pulmonary obstruction diseases. Based on the above-mention circadian change of pulmonary function of asthma

patients, the so called eccentric therapy by giving medicines before go to bed and after wake up, might be a better treatment option for patients' long term treatment. The eccentric treatment gives higher drug concentration during the worse stage of lung function and giving a lower concentration or drug free period during the better lung function stage during a day. Therefore, patients' body will have time to recover and accumulate the endogenous trachea-relaxing substances. In addition, such eccentric therapy could also use the same combination medicine as reliever to a patient with asthma or chronic pulmonary obstruction diseases when it is necessary. The advantage is that the additional inhaled corticosteroid will help to reduced and recover the beta2-agonist acute tolerance and helping the patients to control their disease earlier. Therefore, compared with the current combination control therapy which contains a 12 hours acting drugs (LABA) used twice daily which having the potential to exhaust tracheal relaxing mediator, the eccentric therapy using inhaled corticosteroids plus SABA (ICS+SABA) could be a better treatment choice to preserve patient's lung function.

[0025] In addition, such combination could also be used as needed to improve the current short acting beta2-agonist alone reliever treatment. The reason is that when there are asthma symptom developed and need a bronchodilator, it also means that the tracheal are under inflammation condition. Hence, the addition of inhaled corticosteroids could help reduce the tracheal inflammation and help the patients to recover earlier.

[0026] Other objects, advantages and efficacy of the present techniques have been studies through the following experiment with reference to the accompanying drawings, in which:

In order to understand the alleviating effects of the present techniques, pharmaceutical composition, compare with that the sensitized mice are given long-active or short-acting beta2-agonist alone (Salmeterol or Procaterol), and compare with the reaction of the normal mice and the control group.

[0034] The results are shown as in Figure 2 to 4, no matter administrating alone or administering in combination of the long-acting Salmeterol or medium-short-acting Procaterol, when giving a single super high dose can both make the sensitized mice to show an acute drug tolerance phenomenon of reducing the ability of trachea relaxing. The result shows that there is a suitable dose range for displaying medical effects. Administering the beta2-agonist twice daily cannot avoid the phenomenon of the acute tolerance, giving the inhaled corticosteroid, Budesonide, under a suitable dose range can help to relieve the acute drug tolerance. However, when using too high dose of the beta2-agnist, Budesonide cannot relieve the acute drug tolerance, and it shows that the dosage of the beta2-agonist has exceeded body affordability. These results also indirectly proved that the present techniques propose an idea and assumption that "an acute drug resistance results from exhaustion of endogenously trachea-relaxing substances.

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technique which is a combination of inhaled corticosteroid with medium-short acting beta2-agonist for all level patients with emergent exacerbations as reliever, it might help the patient's recover from their asthma exacerbations.

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[0038] Therefore, the present technique which is a compound of corticosteroid with rapidly short-acting beta2-agonist which can offer a better choice for using as an reliever for treating acute bronchial constriction of asthma in all levels patents. Such extra corticosteroids given can provide extra anti-inflammatory effect to assist patients to relieve from disease condition earlier.

[0039] In accordance with composition of the formula of the present embodiment, medium-short acting beta2-agonist are used, such as Procaterol, Fenoterol, Terbutaline, Albuterol and its base drugs, which has bronchodilating actions of 6 to 8 hours. If the above medicine was administrated 3 to 4 times a day, the phenomenon of acute tolerance will be produced rapidly, and the duration of broncodilating effect will be reduced to 3 to 4 hours. Use the medium-short acting beta2-agonist together with an inhaled corticosteroid, such as budesonide, and adapting an eccentric way of before sleeping and after awake can restore duration of action back to 6 to 8 hours. Therefore, the present technique can help the long-term control of patients' disease and reducing the acute exacerbation.

[0040] The beta2-agonist with 3 to 8 hours duration, such as Procaterol HCl, Procaterol, Albuterol, Albuterol sulfate, Fenoterol, and Fenoterol hydrobromide etc. was use to combined with inhaled corticosteroid, such as Budesonide, Fluticasone, Beclomethasone, Mometasone, Ciclesonide, Triamcinolone and their base morphology, generate a quantitative combination metered dose inhaler or a dry powder inhaler. The weight ratio of the beta2-agonist and the inhaled

corticosteroid is about 1:2 w/w% to 1:70 w/w%, and the better range is about 1:5 w/w% to 1:60 w/w%.

[0041] The clinical usage is giving the combination drug to asthma patients as controller before sleeping and after waking up. The advantage is to provide patients' body a low dose period to recover trachea relaxing mediators for the next day treatment during their better lung function period in the afternoon,. Hence, it reduces the chance of acute tolerance of the beta2-agonist and help patients to control their conditions stably.

[0042] Compared with using short-acting beta2-agonist (Procaterol, Albuterol, Terbutaline, Fenoterol and its base drugs) alone, the combination has additional anti-inflammatory inhaled corticosteroid. The inhaled corticosteroid provided the anti-inflammatory and mitigation of the acute tolerance resulting from overuse of beta2-agonist, it can help to relieve conditions and reduce numbers of severe exacerbations.

[0043] The present technique which is an eccentric treatment of the combination of inhaled beta2-agonist with corticosteroid and can be administered via HFA MDI or DPI (dry powder inhaler). The above-mentioned pharmaceutically acceptable vectors are provided as an excipient system for need of preparing the combination product of the beta2-agonist with the corticosteroid, so it makes the administered animals or humans not to have adverse reactions, allergies or other non- appropriate responses. Carriers or an excipient system can also include the proper amount of surface active agents, solvents, suspending agents and propellants for stabilizing prescriptions. The formulation of HFA

MDI used in the present technique is usually adopted 1,1,1,2-tetrafluoroethane (Tetrafluoroethane, HFA 134a, HFC 134a) or 1,1,1,2, 3,3,3 - Heptafluoropropane (Heptafluoro-n-propane, HFC 227ea, HFC 227, HFA 227), and depending on necessity, it can also be a mixture formulation of HFA 134a and HFA 227. Dry powder inhalation could be a single dose inhaler or multiple dose inhaler composed of carrier-free active pharmaceutical ingredients or use lactose as a carrier.

[0044] Experimental methods:

[0045] Balb/c mice are divided into normal control group, asthma control group (Ova control group), and each group which is contained 2 to 28 mice (in the pre-exploration stage, the number of the mice is fewer, and in the post stage after confirming dose, the number of the mice for repeatedly confirming experiments is more) is supplied with the same water and food. In the asthma control group, at first, the mice are received i.p. with ovalbumin (OVA) with the interval of 10 days as shown in Figure 1, and the mice are made to have allergies, on the 24th days, the blood are collected from the eye sockets of the mice to be examined by IgE Elisa quantification for confirming that the mice have allergies. In the normal control group, the mice are not made to have allergies and are given salt water. In the asthma control group, the mice are also given salt water but not medicine for treating, and 3 days before administration, the mice are continuously given OVA in the nasal to enhance allergies.

[0046] The mice of each group are divided into subgroups by different medicine and different dose. Each subgroup is divided into single administration within 7 days (one dose), once administration a day for 7 days (qd) or twice administrations a day (bid) for 7 days. The salt water or mixture drug of Procaterol mixing with Budesonide in the salt water is directly administered in the throats of the mice.