Intratracheal Sensitization/Challenge-Induced Biphasic Asthmatic Response and Airway Hyperresponsiveness in Guinea Pigs

Nobuaki Mizutani,*a Shouichi Inui,b Shin Yoshino,a and Takeshi Nabe,c

a Department of Pharmacology, Kobe Pharmaceutical University; 4–19–1 Motoyamakita, Higashinada, Kobe 658–8558, Japan; b Pharmacological Research Laboratories, Drug Safety Testing Center Co., Ltd.; 88–75 Shingo, Higashimatsuyama, Saitama 353–0071, Japan; and c Department of Pharmacology, Kyoto Pharmaceutical University; 5 Nakauchi, Misasagi, Yamashina, Kyoto 607–8414, Japan.

Received June 16, 2010; accepted September 22, 2010; published online September 27, 2010

In most experimental model of asthma using guinea pigs, the animals are made to inhale an aerosolized antigen which passes through the nasal cavity. In the present study, we attempted to create an animal model of asthma showing a biphasic asthmatic response and airway hyperresponsiveness, in which the allergic responses are restricted to the lung. Guinea pigs were sensitized by the intratracheal instillation of ovalbumin (OVA)+Al(OH)3, once a day for 7 d, and then intratracheally challenged with OVA 12 d after the last sensitization. The change in specific airway resistance (sRaw) and airway responsiveness to histamine were measured. Pranukast (100 mg/kg), theophylline (50 mg/kg), and dexamethasone (10 mg/kg) were orally administered 18 and 2 h before the antigen challenge. The challenge caused a marked biphasic elevation of sRaw with peaks at 5 min and 4 h. At 24 h, airway hyperresponsiveness to histamine was observed. Pranukast, theophylline, and dexamethasone suppressed the late asthmatic response and airway hyperresponsiveness. The early asthmatic response was inhibited by theophylline and dexamethasone. In conclusion, the intratracheal sensitization and challenge caused a biphasic asthmatic response and airway hyperresponsiveness in guinea pigs. This model may be useful for the evaluation of anti-asthma drugs.

Key words allergic asthma; animal model; intratracheal administration; biphasic asthmatic response; hyperresponsiveness

The guinea pig has long been employed as a model of allergic asthma, because the pulmonary responses following an antigen challenge in the animal is similar to those of asthmatic patients. The airway smooth muscle of guinea pigs well reacts with various chemical mediators such as histamine, cysteinyl leukotrienes (CysLTs), and thromboxane A2, which is very similar to that of human beings. In addition, when sensitized guinea pigs are exposed to an aerosolized antigen, a characteristic feature of chronic asthma known as the late asthmatic response (LAR) is evoked several hours after an early asthmatic response (EAR) that occurs approximately 10 min after the challenge.

We too have developed an animal model of asthma showing a LAR, in which guinea pigs were sensitized and challenged through exposure to aerosolized ovalbumin (OVA)+Al(OH)3 and OVA alone, respectively. The mists were generated by a glassware pressure nebulizer, as their mean diameters (approximately 2 μm) was smaller than that (approximately 5 μm) generated by an ultrasonic nebulizer. Thus, the asthmatic obstruction in the model should occur in the lower airway. However, approximately 20% of the mist trapped in the airway tissues was found to be deposited in the nasal mucosa despite using the glassware pressure nebulizer. In addition, it took more than 4 months to induce the LAR through repeated the aerosols, not particularly convenient for pharmacological examinations of anti-asthma drugs.

Direct administration into the trachea without a surgical operation is difficult in guinea pigs. However, we recently developed a method for intratracheal administration in conscious guinea pigs. In the present study, we attempted to develop a new model showing a biphasic asthmatic response, in which OVA+Al(OH)3 and OVA are delivered into the lung by intratracheal administrations over several weeks. In addition, we evaluated in the model whether airway hyperresponsiveness, and infiltration by leukocytes such as eosinophils were induced. Furthermore, effects of several anti-asthma drugs on changes in the parameters of asthmatic responses were assessed.

MATERIALS AND METHODS

Animals Male, 6-week-old Hartley guinea pigs (Japan SLC, Hamamatsu, Japan) were used. The animals were housed in an air-conditioned room at 23±1°C and 60±10% humidity, with lights on from 8:00 a.m. to 8:00 p.m.; they were fed a standard laboratory diet and given water ad libitum. The first sensitization was started 1 week after purchase. This animal study was approved by the Experimental Animal Research Committee at Drug Safety Testing Center Co., Ltd. (Saitama, Japan).

Intratracheal Instillation The intratracheal instillation was performed in conscious guinea pigs without any surgery, as described previously. Briefly, after the removal of any food struck to the inner surface of the oral cavity with cotton wool, the mucosal surface of the oral cavity and larynx was topically anesthetized by painting on a 1% lidocaine solution absorbed into cotton wool. Then, stainless steel tubing (external diameter: 3.40 mm; internal diameter: 2.84 mm; length: 70 mm), the tip of which was bent at an angle of 120°, was properly attached to the opening of the trachea at the larynx. Teflon tubing (30 mm) was attached to the other end of the oral cannula, and the inner surface of the Teflon tubing was monitored for moisture indicating expiration by the guinea pig through the oral cannula. After confirmation that the guinea pig was breathing through the stainless steel tubing, OVA+Al(OH)3 or OVA was instilled into the tube with a pipet at a volume of 100 μl/animal.

* To whom correspondence should be addressed. e-mail: mizutani@kobepharma-u.ac.jp © 2010 Pharmaceutical Society of Japan
Sensitization and Challenge  As shown in Fig. 1, guinea pigs were sensitized by the intratracheal instillation of a suspension of OVA adsorbed onto Al(OH)$_3$ at a dose of 100 µg OVA/mg Al(OH)$_3$/100 µl/animal once a day for 7 d. The sensitized animals were then intratracheally challenged with OVA at a dose of 1 mg/100 µl/animal 12 d after the last sensitization. In order to prevent anaphylactic death, pyrilamine (10 mg/kg) was intraperitoneally administered 30 min before the OVA challenge.

Pranlukast (100 mg/kg), theophylline (50 mg/kg), and dexamethasone (10 mg/kg) were administered orally twice at 18 and 2 h before the antigen challenge.

Measurement of the Biphasic Asthmatic Response  Specific airway resistance (sRaw) was measured 1 h before and 5 min to 8 h after the OVA challenge using a two-chambered, double-flow plethysmograph system (Pulmos-I; M.I.P.S., Osaka, Japan) according to the method of Pennock et al. Data were expressed as the change in sRaw (% increase). Effects of anti-asthmatic drugs on LAR were expressed as the area under the response curve (AUC) for changes in sRaw at 2—8 h after the challenge.

Measurement of Airway Hyperresponsiveness to Histamine  To assess airway hyperresponsiveness to histamine, changes in lung resistance (RLung) after the administration of histamine were measured by Pulmos-II (M.I.P.S., Osaka, Japan) according to the method of Pennock et al. Data were expressed as the change in Rlung (% increase). Effects of anti-asthmatic drugs on airway hyperresponsiveness were expressed as AUC calculated from dose–response curves for histamine.

Leukocytes in Bronchoalveolar Lavage Fluid (BALF)  Immediately after the assessment of airway hyperresponsiveness, mice were killed and the lung was lavaged with saline (5 ml/animal once a day for 7 d. The sensitized guinea pigs were anesthetized with an intraperitoneal injection of pentobarbital sodium at 50 mg/kg, a tracheotomy was performed, and an endotracheal cannula was inserted. The animals were artificially ventilated via the cannula using a small animal respirator adjusted to a tidal volume of 6 ml/kg at a rate of 60 beats/min. While being artificially ventilated, the animals were injected intravenously with a physiological saline solution containing various concentrations of histamine (5, 10, 20, 40, 80 µg/ml). Data were expressed as the change in Rlung (% increase) and then each dose was converted logarithmically. Effects of anti-asthmatic drugs on airway hyperresponsiveness were expressed as AUC.

RESULTS

Biphasic Asthmatic Response  Figure 2 shows the time-course of change in sRaw after the challenge with OVA in the sensitized guinea pigs. The challenge induced a biphasic elevation of sRaw peaking at 5 min (EAR) and 4 h later (LAR). In the nonsensitized-challenged animals, no change in sRaw was induced.

Before the challenge, the sRaw in the sensitized-challenged group (1.23±0.12) was not statistically different from that in the nonsensitized-challenged group (1.32±0.16).

Figure 3 shows the effects of pranlukast, theophylline, and dexamethasone on the biphasic asthmatic response. All three significantly suppressed the LAR. Furthermore, theophylline and dexamethasone, but not pranlukast, significantly suppressed the EAR.

Leukocytes in BALF were counted using a hemocytometer and based on their morphological criteria, classified as monocytes, neutrophils, or eosinophils.

Statistical Analyses  Data are shown as the mean±S.E. Statistical comparison between two groups were made using Student’s t-test. To compare more than two groups, Dunnett’s test was used after an one-way analysis of variance (ANOVA). A probability value of $p<0.05$ was considered to indicate statistical significance.

Fig. 2. Time Course of Changes in Airway Resistance (sRaw) after the Antigen Challenge in Sensitized Guinea Pigs

Each value represents the mean±S.E. for 8 animals. ++$p<0.05$ and +++$p<0.01$, compared with the NS-C group. sRaw, specific airway resistance; S-C, sensitized-challenged; NS-C, nonsensitized-challenged; EAR, early asthmatic response; LAR, late asthmatic response.

Fig. 3. Effects of Pranlukast, Theophylline, and Dexamethasone on the Biphasic Asthmatic Response Induced by the Antigen Challenge in Sensitized Guinea Pigs

Each value represents the mean±S.E. for 8 animals. ++$p<0.01$, compared with the NS-C group. sRaw, specific airway resistance; S-C, sensitized-challenged; NS-C, nonsensitized-challenged; EAR, early asthmatic response; LAR, late asthmatic response.
In the present study, so as to restrict allergic responses to the lung, we chose to intratracheally administer OVA + Al(OH)$_3$ and OVA, thereby avoiding passage through the nasal cavity. Guinea pigs were sensitized with intratracheal instillations of OVA + Al(OH)$_3$ for 7 d and challenged with a single intratracheal administration of OVA 12 d after the last sensitization. The challenge caused a marked biphasic asthmatic response and airway hyperresponsiveness.

Most studies using experimental models of asthma have not attempted to restrict allergic responses to the lower airway, even though sensitized guinea pigs are known to exhibit nasal blockage after the intranasal instillation of an antigen. In addition, we have reported that when guinea pigs inhaled an Evans blue mist generated by an ultrasonic nebulizer, 80% of the dye trapped in airway tissue was retained in the upper airway. Therefore, it is important to deliver an antigen specifically to the lung in experimental models of asthma. The intratracheal sensitization and challenge used here efficiently elicited allergic responses in the lungs, enhancing the value of this model.

Our previous animal model of asthma was developed by exposing guinea pigs to a fine mist of antigen solution. However, it took more than 4 months to induce an LAR. Another key finding of the current study is that the EAR, LAR, and airway hyperresponsiveness were induced by the first antigen challenge, and took only 19 d, although only the EAR was induced after the first challenge in our previous model. These results suggest the intratracheal method to be an efficient means of sensitization/challenge in a short period of time.

The findings suggest this experimental asthmatic model to be suitable for the development of anti-asthma drugs. Therefore, we examined effects of theophylline, pranlukast, and dexamethasone, which have been used clinically for the treatment of asthma. The effect of theophylline on the EAR may be due to the inhibition of bronchoconstriction and airway edema (plasma leakage) as previously reported. Dexamethasone also inhibited the EAR, though glucocorticoids do not prevent the immediate activation of mast cells resulting in the release of chemical mediators. The effectiveness of dexamethasone against the EAR may be associated with various inflammatory changes, such as a suppression of vascular permeability.

The LAR and AHR were also inhibited by theophylline. The effect may be dependent upon not only a decrease in bronchoconstriction and/or mucosal edema but also anti-inflammatory actions, such as the non-selective inhibition of phosphodiesterase, antagonism of adenosine receptor and/or activation of histone deacetylase, which can lead to reduced levels of cytokines that contribute to the development of LAR and AHR. Furthermore, pranlukast inhibited the LAR and AHR, suggesting that CysLTs are important mediators in the development of these responses. However, the suppression by pranlukast was weaker than that by theophylline or dexamethasone. This difference may be related to the recruitment of eosinophils, which theophylline and dexamethasone but not pranlukast inhibited, suggesting the recruitment of eosinophils to be related to the LAR and AHR, suggesting that CysLTs are important mediators in the development of these responses. However, the suppression by pranlukast was weaker than that by theophylline or dexamethasone. This difference may be related to the recruitment of eosinophils, which theophylline and dexamethasone but not pranlukast inhibited, suggesting the recruitment of eosinophils to be related to the LAR and AHR in this model.

In conclusion, by employing intratracheal sensitization and challenge, we have developed a guinea pig model of asthma showing a biphasic asthmatic response and airway hyper-
responsiveness. This model may be useful for elucidating the pathogenesis of asthma, which could possibly lead to the development of new therapeutic drugs.

REFERENCES