Measurement of nasal patency in anesthetized and conscious dogs

MICHAEL C. KOSS,1 YONGXIN YU,1 JOHN A. HEY,2 AND ROBBIE L. McLEOD2
1Department of Cell Biology, University of Oklahoma, College of Medicine, Oklahoma City, Oklahoma 73190; and 2Allergy, Schering-Plough Research Institute, Kenilworth, New Jersey 07033

Received 27 August 2001; accepted in final form 23 September 2001

Koss, Michael C., Yongxin Yu, John A. Hey, and Robbie L. McLeod. Measurement of nasal patency in anesthetized and conscious dogs. J Appl Physiol 92: 617–621, 2002.—Experiments were undertaken to characterize a noninvasive chronic, model of nasal congestion in which nasal patency is measured using acoustic rhinometry. Compound 48/80 was administered intranasally to elicit nasal congestion in five beagle dogs either by syringe (0.5 ml) in thiopental sodium-anesthetized animals or as a mist (0.25 ml) in the same animals in the conscious state. Effects of mast cell degranulator Compound 48/80, which has been shown to produce nasal congestion in cats (4, 22, 23), methods; nasal congestion; Compound 48/80; nasal airway volume; minimal cross-sectional distance to minimal cross-sectional area

MATERIALS AND METHODS

General procedures. Two series of experiments were undertaken using five adult male, purpose-bred beagle dogs (C and C Kennels, Wewoka, OK) weighing 9–11 kg. All studies were performed in an Association for Assessment and Accreditation of Laboratory Animal Care-accredited facility and were undertaken in accordance with the National Institutes of Health (NIH) Guide for the Care and Use of Laboratory Animals (DHHS Publication No. (NIH) 85-23, revised 1985, Office of Science and Health Reports, Bethesda, MD 20892).

In one series of experiments, the animals were anesthetized with intravenous thiopental sodium (e.g., 25 mg/kg bolus plus 50-mg supplements at 15- to 30-min intervals as needed). After tracheal intubation with a cuffed endotracheal tube, blood pressure and heart rate were monitored using a V6004 monitor (Surgi Vet, Waukeha, WI). Body temperature was maintained at ~37°C by using a recirculating hot-water system. An 8,500-V pulse oximeter (Nonin Medical, Plymouth, MN) was used to continuously monitor arterial PO2.

For studies without anesthesia, dogs were trained (daily over a period of ~1 mo) to remain quiet during the measurement period of ~10–15 s required for three determinations. Animals were gradually acclimated to the procedure with positive reinforcement (dog treats) offered in response to the desired behavior. This initially included training the animals to sit quietly during presentation of the clicking sound produced by the acoustic rhinometer and gradually working up to acceptance of having the probe placed into the nasal cavity. The soft nosepiece used, together with the intranasal application of the probe, allowed for an effective seal without changes in nasal cavity geometry in a large-animal model using dogs. Because the technique is noninvasive, chronic, repeated measurements can be made in the same experimental subjects. In addition, it is possible to train dogs to remain quiet during simple experimental manipulations. To test this model, sequential experiments were undertaken in five beagle dogs in both anesthetized and nonanesthetized states. Nasal cavity geometry was assessed after topical application of the mast cell degranulator Compound 48/80, which has been shown to produce nasal congestion in cats (4, 22, 23).

Address for reprint requests and other correspondence: M. C. Koss, Dept. of Cell Biology, Univ. of Oklahoma College of Medicine, PO Box 26901, Oklahoma City, OK 73190 (E-mail: michael-koss@ouhsc.edu).

The costs of publication of this article were defrayed in part by the payment of page charges. The article must therefore be hereby marked “advertisement” in accordance with 18 U.S.C. Section 1734 solely to indicate this fact.

http://www.jap.org 8750-7587/02 $5.00 Copyright © 2002 the American Physiological Society
need of sealant material (as is needed for similar measurements in humans).

For consistency, all determinations were made at the end of expiration. The average of three to five acoustic rhinometry readings was taken for each time period represented (30-min intervals). In the majority of cases, three sequential readings were consistent and averaged. In a few trials, an obviously out-of-line measurement was obtained. When this occurred, two additional determinations were obtained with all, except for the questionable measurement, averaged and reported for that time point.

**Acoustic rhinometry measurement in dogs.** Anesthetized dogs were placed in a supine position (on a heated thermal blanket) throughout the experiment. Conscious animals were trained to sit quietly on an operating table. Nasal cavity volumes, minimal cross-sectional areas (A_{min}), and the distance to the A_{min} (D_{min}) were determined by using an Eccovision Acoustic Rhinometry System (Hood Laboratories, Pembroke, MA) according to established methods (8, 22). In brief, a wave tube containing a spark sound generator was connected with the nasal cavity by means of a flexible plastic nosepiece. The distance measured from the nostril opening into the nasal cavity was 10 cm. This distance was chosen on the basis of nasal cast impressions and X-ray determinations made of the dog nasal cavity. Acoustic reflections were recorded and amplified with a computer analysis made of the local acoustic impedance changes, which, in turn, are used to provide estimates of volume and cross-sectional area of the nares.

After the experimental procedure, the anesthetized animals were placed in a recovery cage and closely monitored throughout the recovery from anesthesia under the close supervision of a member of the University of Oklahoma College of Medicine Veterinary staff. When the animals had completely recovered from the anesthesia, they were then returned to their home cage. Conscious animals did not require close postexperimental supervision. All of the animals tolerated repeated procedures well with no signs of distress and with no residual side effects.

**Effects of topical application of Compound 48/80.** Acoustic rhinometry was used to assess the effects of mast cell degranulation by Compound 48/80 on nasal geometry in five anesthetized dogs. Each dog received three doses of Compound 48/80 in a crossover design. In anesthetized preparations, the histamine releaser, Compound 48/80 was administered ipsilaterally into the nasal cavity at three dosage levels (1.5, 5, and 15 mg) using a syringe. The volume was held constant at 0.5 ml. Conscious animals received Compound 48/80 as a nasal mist by using an atomizer (model IA-IB, Delong Distributors), also at three dosage levels (5, 15, and 45 mg), with a volume of 0.25 ml. In a preliminary study, with conscious animals, the mist and drops of Compound 48/80 had similar effects on the ipsilateral nasal cavity. However, drugs seemed to also enter into the contralateral side when applied in droplet form with the dogs in the conscious state. This is likely due to reflex responses in the conscious dogs leading to some mixing of fluid containing Compound 48/80 between the two sides of the nasal cavities. A comparable administration of PBS was used for control experiments for both groups. All measurements were taken before, and for 3 h after, administration of Compound 48/80 (at 30-min intervals). A 2-wk washout period was allotted between each experimental group receiving either PBS or Compound 48/80.

**Drugs and statistics.** Compound 48/80 was purchased from Sigma Chemical (St. Louis, MO) and was dissolved in PBS. Control experiments were undertaken by using PBS alone. Nasal cavity volumes, A_{min}, and D_{min} were derived directly from the computer calculations of the acoustic rhinometry apparatus. Statistical significance was determined, for values (means ± SE) taken at 30-min intervals, by using ANOVA followed with Dunnett’s two-tailed t-test. Cardiovascular parameters, before and after treatment, were evaluated by using a paired two-tailed Student’s t-test. Differences were considered statistically significant at P < 0.05 levels.

**RESULTS**

**Effect of Compound 48/80 on nasal airway patency in anesthetized dogs.** There were no differences between the baseline volumes obtained for the left and right nares or between the baseline values of the dogs before Compound 48/80 administrations. Basal nasal volumes were 7.0 ± 0.3, 6.7 ± 0.6, and 7.2 ± 0.5 cm³ for the 1.5-, 5-, and 15-mg Compound 48/80 trials, respectively. Similarly, no significant differences were seen with regard to A_{min} or to D_{min} baseline values (see Tables 1 and 2).

Typical examples of area-distance curves taken before and after challenge with Compound 48/80 are shown in Fig. 1. In these, control values for nasal volume and A_{min} were, respectively, 8.52 cm³ and 0.37 cm² in the anesthetized state and 11.49 cm³ and 0.45 cm² in the same dog without anesthesia. These parameters were reduced to 2.71 cm³ and 0.12 cm², and to 5.63 cm³ and 0.27 cm², 3 h after topical application of Compound 48/80 (15 and 45 mg), respectively. D_{min} increased from 0.66 cm to 3.06 cm and from 0.42 to 5.23 cm after administration of Compound 48/80.

Figure 2 shows composite nasal volume responses of all five dogs in response to topical application (0.5 ml)

---

**Table 1. Comparison of nasal cavity minimal cross-sectional areas in anesthetized and conscious dogs after topical Compound 48/80 administration**

<table>
<thead>
<tr>
<th>Time, min</th>
<th>1.5 mg</th>
<th>5 mg</th>
<th>15 mg</th>
<th>5 mg</th>
<th>15 mg</th>
<th>45 mg</th>
</tr>
</thead>
<tbody>
<tr>
<td>0</td>
<td>0.36 ± 0.02</td>
<td>0.36 ± 0.01</td>
<td>0.37 ± 0.01</td>
<td>0.40 ± 0.01</td>
<td>0.41 ± 0.01</td>
<td>0.42 ± 0.01</td>
</tr>
<tr>
<td>30</td>
<td>0.33 ± 0.03</td>
<td>0.32 ± 0.03</td>
<td>0.26 ± 0.03*</td>
<td>0.40 ± 0.04</td>
<td>0.41 ± 0.01</td>
<td>0.34 ± 0.03*</td>
</tr>
<tr>
<td>60</td>
<td>0.24 ± 0.04*</td>
<td>0.24 ± 0.05*</td>
<td>0.37 ± 0.02</td>
<td>0.37 ± 0.02</td>
<td>0.35 ± 0.03*</td>
<td></td>
</tr>
<tr>
<td>90</td>
<td>0.23 ± 0.02*</td>
<td>0.19 ± 0.03†</td>
<td>0.35 ± 0.02</td>
<td>0.38 ± 0.02</td>
<td>0.33 ± 0.03*</td>
<td></td>
</tr>
<tr>
<td>120</td>
<td>0.21 ± 0.03†</td>
<td>0.15 ± 0.02‡</td>
<td>0.36 ± 0.01*</td>
<td>0.36 ± 0.01*</td>
<td>0.31 ± 0.02*</td>
<td></td>
</tr>
<tr>
<td>180</td>
<td>0.23 ± 0.03*</td>
<td>0.14 ± 0.02‡</td>
<td>0.34 ± 0.02*</td>
<td>0.36 ± 0.02*</td>
<td>0.35 ± 0.03*</td>
<td></td>
</tr>
</tbody>
</table>

Values are means ± SE given in cm² for 5 dogs. Compound 48/80 was administered as droplets (0.5 ml) to the anesthetized animals and as a mist (0.25 ml) to the conscious dogs. *P < 0.05 compared with initial values. †P < 0.01 compared with initial values.
of 3 doses of Compound 48/80 (1.5, 5, and 15 mg) in the anesthetized state. Effects of Compound 48/80 on $A_{\text{min}}$ and $D_{\text{min}}$ values are shown in Tables 1 and 2.

Composite basal mean arterial blood pressure before Compound 48/80 administration, under anesthesia ($n = 15$), was 124.8 ± 4.6 mmHg and heart rate was 117.4 ± 7.2 beats/min. There were no significant alterations of these values after any of the doses of Compound 48/80 in these anesthetized dogs.

**Effect of Compound 48/80 on nasal airway patency in conscious dogs.** Acoustic rhinometry was used to assess the effects of mast cell degranulation by Compound 48/80 on nasal geometry in these same dogs, in this case, without anesthesia. As described in *Effects of topical application of Compound 48/80*, each dog received three doses of Compound 48/80 in a crossover design with no differences between the baseline volumes obtained for the left and right nares or between the baseline values (means ± SE) of the dogs before Compound 48/80 administration. Basal nasal volumes were $13.5 ± 1.0$, $12.1 ± 0.3$, and $12.6 ± 0.3$ cm$^3$ for the 5-, 15-, and 45-mg Compound 48/80 trials, respectively. Similarly, no significant differences were seen with regard to $A_{\text{min}}$ or to $D_{\text{min}}$ baseline values (Tables 1 and 2).

Figure 3 shows composite nasal volume responses of all five dogs in response to topical application (0.25-ml mist) of three doses of Compound 48/80 (5, 15, and 45 mg) in the nonanesthetized condition. Cardiovascular parameters were not measured in the freely moving conscious animals. Tables 1 and 2 document effects of Compound 48/80 on $A_{\text{min}}$ and $D_{\text{min}}$ in these animals.

**DISCUSSION**

Allergic rhinitis is among the most common medical conditions worldwide and presents with a decrease in nasal patency resulting from inflammation of the nasal mucosa, congestion, and rhinorrhea. In the United States, it is estimated that 10–20% of all adults are affected (3). Preclinical studies designed to elucidate the pathophysiological mechanisms as well as drug discovery research in this area have used a variety of experimental animal models.

The “ideal” model for assessment of nasal congestion in animals would be noninvasive, reproducible, easily performed, and focus on the nasal cavity compared with the other airway components. The ability to use conscious animals also would eliminate potential confounding influences of general anesthetic agents.

Although a number of different techniques have been utilized to assess nasal patency in animals, none of these fulfills all of the above criteria. For example, although plethysmographic techniques (9) are noninvasive and can be undertaken in conscious animals, plethysmographic airway resistance measurements...
Acoustic rhinometry is commonly used to study nasal patency in humans. Its use has been validated by using magnetic resonance imaging (8) as well as by comparison with cast impressions of nasal cavities of humans, guinea pigs, and cats (7, 11, 21, 22). Caveats concerning potential sources of artifact and possible misinterpretations of components of the acoustic rhinometry tracings have been addressed (31). Acoustic rhinometry is a noninvasive, reproducible technique, and correlates well with rhinometric (21, 27) and magnetic resonance techniques (8). Initial preclinical applications of acoustic rhinometry were limited to studies in the guinea pig (25, 26) in which correlations with nasal resistance changes and directly measured nasal cavity volume have been established (11, 24).

More recently, an experimental model using acoustic rhinometry has been applied to studies on the anesthetized cat (4, 22, 23). Nasal congestion is produced by topical administration of histamine or by nasal application of a liberator of mast cell histamine, Compound 48/80 (17). In these studies, both procedures significantly reduce the ipsilateral volume and \( A_{\min} \) while increasing \( D_{\min} \) (4, 22, 23). These findings indicate that this cat model may prove useful in investigations of the basic mechanisms of nasal congestion as well as for elucidating the mechanisms of action of antiallergic agents. Although this cat model fulfills many of the criteria listed above, it is unlikely that cats could be easily trained to allow the acoustic rhinometry procedure to be performed in the absence of anesthesia.

In comparison, dogs are more amenable to training for procedures using conscious animals. Use of dogs for studies of nasal congestion may provide several added benefits over other animal models. Not the least of these is the fact that dogs are widely used in the pharmaceutical industry as a model to profile pharmacokinetic and safety aspects of potential new drugs. More specifically, with regard to the upper airway, dogs have a total nasal cavity volume more comparable to that of humans and the nasal physiology and pharmacology is well characterized (19, 20, 28).

In the present study, Compound 48/80 produced a consistent dose-related decrease of nasal volume and \( A_{\min} \) in the anesthetized dog. A more variable increase of the \( D_{\min} \) was found, consistent with previous studies in human and other species (4, 12, 26). Compared with results reported in the anesthetized cat, the peak responses were somewhat delayed in time, in that they occurred at -2 h after topical application of Compound 48/80. Peak responses, in cats, were seen at -1 h posttreatment (4, 22). Dogs also appear to be somewhat less sensitive to the actions of Compound 48/80, because the maximal dose of 15 mg was somewhat greater than the 5-mg dose needed for maximal responsiveness in the anesthetized cat (22).

Compound 48/80 also produced alterations of the nasal geometry in conscious dogs. As in the anesthetized animals, there was a dose-related decrease in nasal volume and \( A_{\min} \), as well as a variable increase of the \( D_{\min} \) at the highest dose (45 mg). Overall, the same dogs appeared to be less sensitive to topical Compound 48/80...
ments for nasal congestion. This observation could be due to the anesthetic drug, thiopental sodium, or the fact that the nasal volume was much larger in the conscious state.

The pronounced difference in nasal volume between the anesthetized and nonanesthetized preparations is likely due to anesthesia-induced depression of sympathetic neural tone to the nasal vasculature. Nasal blood vessels appear to be highly innervated, because sympathetic nerve section, even in anesthetized animals, results in significantly increased nasal blood flow in rats (13), cats, and dogs (15, 16). In the anesthetized dog, sympathetic nerve section increases nasal blood flow by between 14 and 43% (15, 16).

In this study, we have characterized a chronic dog model of nasal congestion. Topical application of Compound 48/80 was utilized to decrease nasal patency (due to local histamine release from mast cells) as measured by acoustic rhinometry. Mast cell degranulation resulted in a dose-related decrease in nasal cavity volume and $A_{\text{min}}$, in both anesthetized and conscious dogs. Increased sympathetic nerve tone in the nonanesthetized preparations was reflected in a much larger basal nasal volume. Acoustic rhinometry in dogs may be a useful tool in investigating pathophysiological mechanisms of allergic rhinitis, as well as for drug discovery oriented toward novel pharmacological treatments for nasal congestion.

The authors thank Linda Hess for expert technical assistance in these studies.

REFERENCES


