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PATHO-PHYSIOLOGIC RESPONSE TO SINGLE AND MULTIPLE AIR POLLUTANTS IN HUMANS AND ANIMALS

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Pittsburgh, University Pittsburgh, Pennsylvania

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FINAL REPORT

PATHO-PHYSIOLOGIC RESPONSE TO SINGLE AND MULTIPLE AIR POLLUTANTS IN HUMANS AND ANIMALS

Covering the Period

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U. S. Public Health Service

By

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with the assistance of
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July 1, 1970

N O T I C E

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Abstracts (Cont'd)

concentration of sulfur dioxide finality, either alone or in the presence of NaCl aerosol (10 mg/m3). When the pollutants were administered via endotracheal catheter and face mask, an increased frequency of significant changes in pulmonary flow resistance in these animals were used in the former studies. All alterations in parameters of response were reversible shortly after exposure ceased. Morphological examination of lung tissue sections after rapid freezing with Freon indicated that measurement of alterations in airway size is not possible in the range of changes of pulmonary flow resistance reported here (< 100%). Methods and data for all experiments are presented in detail.

b

SUMMARY

Twenty healthy, adult male cats were lightly anesthetized (Nembutal), tracheotomized and were then breathed by a Harvard pump at a fixed frequency and tidal volume. Purified Medical Grade breathing air with or without sulfur dioxide in air or sulfur dioxide in combination with sodium chloride aerosol in air, was delivered to the animals in predetermined exposure sequences and for fixed durations of time. Parameters of response used to judge adaptation of cats to the inhaled challenges of pollutants were pulmonary flow resistance and lung compliance. Measurement methods were standard and included continuous trace recordings of air flow, tidal volume, transpulmonary pressure and blood pressure. Each animal acted as his own control. In addition, pollutant mixtures were delivered to animals via endotracheal catheter and/or face mask to evaluate the possible influence of receptors which may be present in the nasopharyngeal chamber and in the trachea above the tracheal cannula. After selected exposures, the pleural cavity was opened and liquid Freon was used to freeze the lungs. Procedures were developed to obtain histological sections in order to measure changes in airway size.

The major finding of the study was the variability of the responses of the test animals. Certain subjects showed increased pulmonary flow resistance at low SO₂ concentration, and were the

analogues of the "reactors" in human populations. Approximately 20 ppm SO₂ in air were required to evoke a significant change in pulmonary flow resistance in "reactors". The majority of animals showed no response at this concentration of sulfur dioxide in air, either alone or in the presence of NaCl aerosol (10 mg/m³). When the pollutants were administered via endotracheal catheter and face mask, an increased frequency of significant changes in pulmonary flow resistance in these animals was suggested, as compared to animals receiving challenges by tracheal cannula. However, fewer test animals were used in the former studies.

All alterations in parameters of response were reversible shortly after exposure ceased. This finding in cats is similar to reports of early reversal of these parameters in spontaneously breathing human subjects exposed to the same pollutants. In guinea pigs, pulmonary flow resistance, which was elevated by exposure to pollutants, returned to normal after a prolonged period (at least one hour) following cessation of exposure.

Morphological examination of lung tissue sections after rapid freezing with Freon indicated that measurement of alterations in airway size is not possible in the range of changes of pulmonary flow resistance reported here (< 100%).

Methods and data for all experiments are presented in detail.

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I. INTRODUCTION

A. Review of Contract Aims and Efforts

This Contract was initiated on January 3, 1967. Its aim was to perform a series of experiments in animals and in man to determine the extent of and the mechanisms associated with, synergism between inert aerosols and irritant gas in combination after inhalation. The parameters used to judge synergistic response were to be changes in pulmonary mechanics. Initially, it was our aim to study humans in both the resting state and during exercise. The departure from this laboratory of Dr. George Burton for Loma Linda University in 1968 resulted in curtailment of human studies because a physician was not associated with the study, thus precluding adherence to University guidelines for experiments involving human subjects. However, studies involving exposures of human subjects at rest were completed and a publication entitled "Response of Healthy Men to Inhaled Low Concentrations of Gas-Aerosol Mixtures" by Burton, G., Corn, M., Gee, B. L., Vasallo, C. and Thomas, A. P. resulted and appeared in the AMA Archives of Environmental Health 18, 681 (1969). (Appendix I.) Efforts on this Contract then shifted to studies of synergism in cats exposed to the same aerosol-irritant gas mixture as was used in the human studies. In addition to the presence or absence of response, we were interested in the mechanisms associated with the response.

Quarterly Progress Reports have been submitted since the inception of this Contract. The publication referenced above summarized the work accomplished during the initial 18 months of the Contract. The purpose of this report is to summarize efforts during the latter 18 months of the investigation. Progress during this period was exclusively associated with synergistic studies utilizing cats as test animals. The work reported here is scheduled for presentation at the 10th Air Pollution Medical Research Conference to be held in New Orleans, October 5-7, 1970. The paper is entitled "Response of Cats to Inhaled SO₂ and SO₂-NaCl Aerosol Mixtures in Air" by M. Corn, N. Kotsko, D. Stanton, and W. Bell. The presentation will be a summary version of data and discussion presented here.

Background for Inhalation Studies Using Cats as Subjects В. The biological assay procedure for air pollutants, as originally developed by Amdur and Mead (1), utilizes guinea pigs as experimental animals. The results of work by Amdur and her coworkers is reviewed in the document Air Quality Criteria for Sulfur Oxides (2). The more than a decade of work by Amdur, in which guinea pigs have been exposed to a variety of gaseous and particulate pollutants, provides the most extensive body of information available on the response of an animal species to air pollutants. The data relative to human response to mixtures of pollutants, as judged by alterations in parameters of pulmonary mechanics, have not substantiated the findings in guinea pigs. The studies on humans are few in number and are contradictory, as discussed by Amdur in a recent appraisal of sulfur dioxideaerosol mixtures and their effects on animals and man (3). Because of heavy reliance on the guinea pig assay system and the difficulties inherent in extrapolating these data to man, or even in drawing conclusions from these data relative to effects of these systems on man, it was considered appropriate to study the effects of these mixtures on another species.

A reasonable and appropriate question is "why select the cat as the species of choice in these studies?" A series of investigations by Widdicombe (4,5,6) and Nadel (7,8) delineated the mechanisms of bronchoconstriction and peripheral airway constriction in cats. Thus, studies of response to pollutant gas and aerosol mixtures in cats promised to answer whether the responses reported in guinea pigs occurred in other animal species. Also, if they occurred, the mechanisms of action could be investigated by isolating previously studied mechanisms and pathways of response by means of well developed experimental techniques. The investigation reported herewas a follow-up to this reasoning. We report on the effects of sulfur dioxide alone, and in combination with sodium chloride aerosol, on the pulmonary flow resistance and lung compliance of cats. In addition, results will be reported for pathological examination of lungs initially rapidly frozen with liquid Freon following exposure to pollutant free or to pollutant laden air. Experimental methods will be described and this will be followed by results and discussion.

II. EXPERIMENTAL METHODS

A. Pollutant Aerosol and Gas Concentration

Pollutant mixtures for the exposures were produced using a portable aerosol and pollutant gas supply apparatus designed and constructed for this study (Fig. 1).

Medical-grade compressed air was passed through activated carbon and silica gel. The stream of air was metered by the use of calibrated orifices, before entering the Dautrebande D_{30} l generator or the Venturi tube for mixing with aerosol and gas. Sulfur dioxide gas was supplied to the Venturi throat by a syringe driven by an infusion pump. The Dautrebande D_{30} l aerosol generator was filled with isotonic NaCl and placed in an opening at the base of the Venturi tube. The salt solution was replaced every 15 minutes to prevent a significant increase in NaCl concentration due to the evaporation of water.

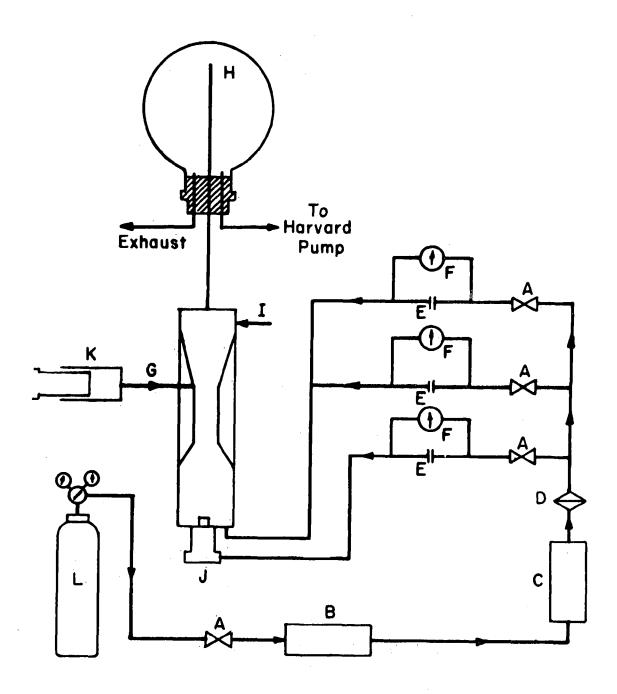
The mixture exited from the Venturi mixing tube into a reservoir balloon, where it was either exhausted, or withdrawn by the Harvard pump.

All components of the system, with the exception of the balloon, were of stainless steel, rigid plastic, or Teflon.

A three way valve at the entry to the Harvard pump could be set for pollutant mixture or room air entry to the pump. The Harvard pump was equipped to permit the setting of tidal volume and breathing frequency. The most commonly used settings for these parameters were 75 ml and 20 cps, respectively. The inspiratory stroke of the pump was under positive pressure. The animal expired under the driving force of lung elasticity.

FIGURE 1

SCHEMATIC OF POLLUTANT GENERATION APPARATUS



Schematic of pollutant generation apparatus. A, Valves; B, "catch-all" air cleaner; C, silica gel; D, Millipore HA filter; E, Critical orifice; F, pressure gauge; G, SO₂ inlet; H, mixing balloon; I, Herschel-Type Venturi tube; J, Dautrebande D₃₀l; K, Motor driven syringe; L, Medical grade compressed air; triangle, flowdirection.

B. Pollutant Gas and Aerosol Sampling and Analysis

Samples for measurements of aerosol and pollutant gas concentrations were withdrawn from a stainless steel port in the exhaust line following the balloon reservoir. It was ascertained that concentrations measured above the tracheal cannula did not differ from those withdrawn at the former site.

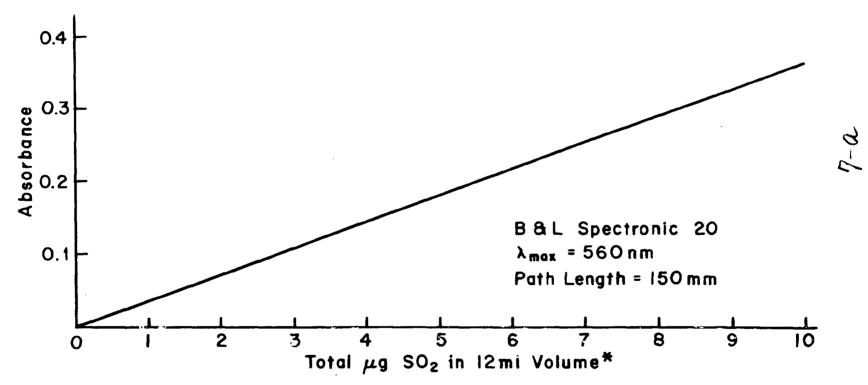
1. Sulfur Dioxide

In order to determine SO₂ concentration, 0.76 liters/min. of the pollutant mixture was drawn for three minutes into a midget impinger containing 10 ml of West-Gaeke reagent. The samples were then analyzed spectrophotometrically using the West-Gaeke method ⁽⁹⁾ with the Pararosaniline modification specified by Pate ⁽¹⁰⁾. The high sulfur dioxide concentrations in the experimental protocol required additional modifications in the analytical method, as follows.

- a) A 1 ml aliquot was removed from the original 10 ml sample and was diluted with 10 ml of unexposed absorbing reagent. The 9 ml and 1 ml aliquots were then prepared according to the original West-Gaeke procedure (9).
- b) A calibration curve was prepared using standardized solutions of sodium bisulfite, which ranged in concentration from 0.1 to 5.0 µg of SO₂ per ml. The solutions were standardized by the iodometric titration method described by the Intersociety Committee for Ambient Air Sampling and Analysis (11). A new calibration curve was prepared whenever a new stock dye solution was used. A typical calibration curve is shown in Figure 2. All calibration

FIGURE 2

CALIBRATION CURVE FOR SO₂ USING MODIFIED WEST-GAEKE PROCEDURE



* 12 ml includes IOml. absorbing reagent +1 ml dye + 1 ml formaldehyde

curves adhered to Beer's Law in the absorbance range less than 0.700. If a 9 ml sample yielded an absorbance value greater than 0.700, it was discarded and the 1 ml sample was analyzed.

The concentration of SO_2 in μg SO_2 per ml was calculated as follows:

$$\mu g SO_2 per ml = \frac{V_T \times Absorbance}{Slope \times V_F}$$

where $V_m = total$ volume of solution, ml

 V_{F} = fraction of original sample analyzed

The total volume of the 9 ml sample was 11 ml, which consisted of 9 ml of the original sample, 1 ml of dye and 1 ml of formaldehyde. The 1 ml sample contained a total volume of 13 ml, which consisted of 1 ml of the original sample, 10 ml of unexposed absorbing reagent, 1 ml of dye and 1 ml of formaldehyde.

The concentration of sulfur dioxide was converted to Parts Per Million (ppm) in air as follows:

$$PPM SO_2 = \frac{382 \times C}{R \times T}$$

where 382 = Conversion factor for ug/ml to ppm at 25°C, 760 mm.

C = Concentration of SO₂ in µg/ml

R = Sample flow rate in cc per minute

T = Sample time in minutes

It should be noted that whenever sodium chloride was present in the system, sampling for sulfur dioxide was performed by first drawing a sample through HA Millipore filter paper to eliminate sodium chloride interference. Several calibration runs indicated that the loss of sulfur dioxide on the filter paper was negligible.

2. Sodium Chloride Aerosol

The concentration of sodium chloride aerosol was determined by withdrawing the exposure mixture at 21 liters/min. for 20 minutes through HA Millipore paper, leaching the salt by filter immersion in distilled water, and analyzing electrical conductivity. A calibration curve was prepared using reagent-grade sodium chloride. Because of the length of time required to sample for the sodium chloride aerosol, this was done at the end of the exposure periods. Several nonexposure and postexposure checks found the concentrations to be very consistent over a period of several hours.

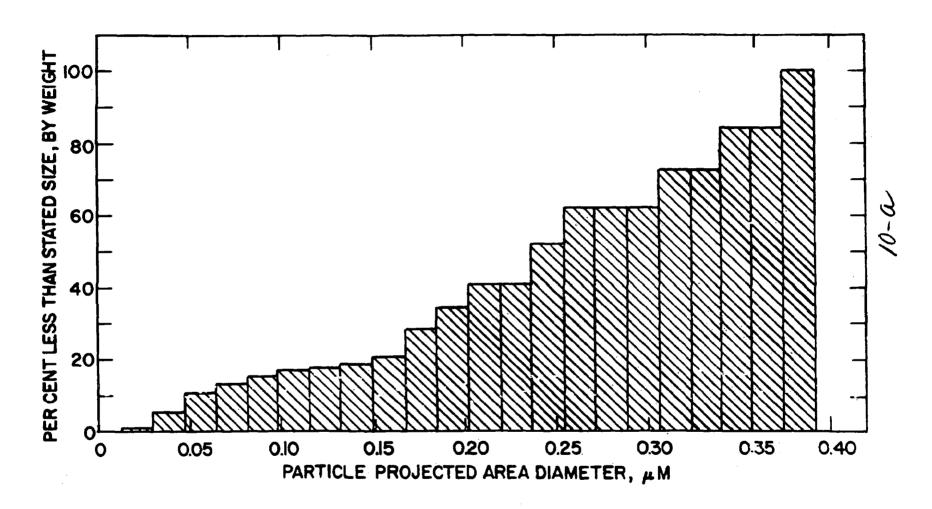
The particle size distribution of the aerosol was determined by first sampling with an oscillating thermal precipitator onto a carbon-coated glass coverslip. The carbon film was transferred to a 200 mesh electron microscope grid prior to obtaining photographs with an electron microscope. The particles were sized using a Zeiss TGZ3 particle sizing unit. The particles were all smaller than 0.40 µm by weight; the mean size and standard deviation were 0.25 µm and + 0.01 µm, respectively. (Figure 3.)

The concentrations of SO₂ gas and NaCl aerosols used in these studies will be cited in the Results and Discussion section of this report. However, it is appropriate here to indicate the variations in concentrations used in these studies. LOW sulfur dioxide concentrations were 15-25 ppm. HIGH sulfur dioxide concentrations were 30-40 ppm. Sodium chloride aerosol concentrations were 9-10 mg/m³.

FIGURE 3

CUMULATIVE PARTICLE SIZE DISTRIBUTION CURVE FOR TEST

AEROSOL SIZED AT 32,000x



CUMULATIVE PARTICLE-SIZE DISTRIBUTION CURVE FOR TEST AEROSOLS (NaCl), SIZED AT 32,000x WITH ZEISS TGZ-3 AND ELECTRON MICROSCOPE

A representative time-concentration diagram for each pollutant during a single experiment is shown in Figure 4 (Cat No. 1610).

C. Animal Handling and Preparation

Upon receipt of animals a routine examination was made to determine the presence of gross abnormalities. Animals with abnormalities were returned to the supplier; all other animals were weighed and were then injected with feline pneumonitis vaccine. (Veterinary care or consultation was available when needed.) An initial isolation period of 2 weeks was observed.

All animals were caged, fed, and attended in accordance with the rules and regulations of the United States Department of Agriculture and Federal Act of August 24, 1966 (P.L. 89-544).

A trained technician observed, weighed and kept a daily record of any changes that occurred. At the conclusion of the isolation period animals were permitted to exercise daily.

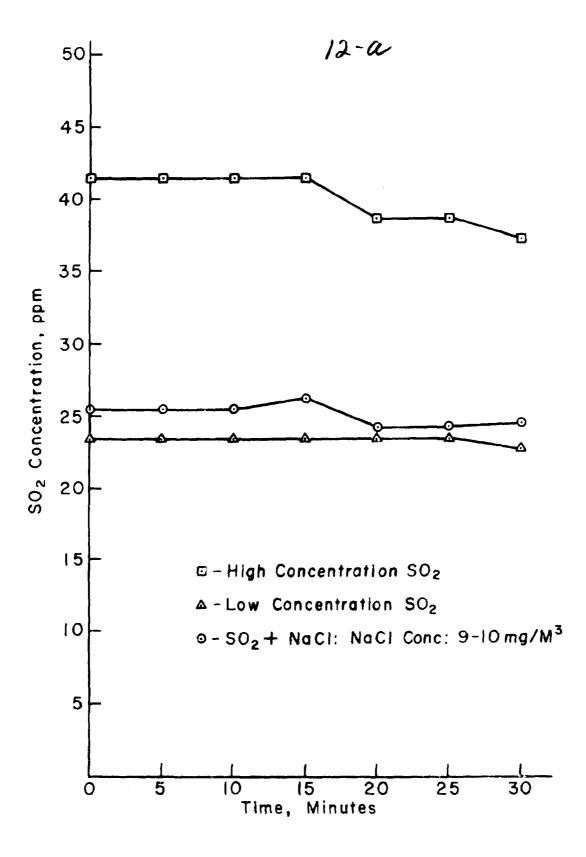
Food and water were removed from an animal cage 18 hours prior to inhalation exposure. An examination was performed for symptoms of diarrhea, diuresis, conjunctivitis, etc. The entire ventral and cervical region of the cat was then shaved and a vacuum was used to remove excess hair.

Nembutal (30 mg/Kg, intraperitoneally) was used to anesthetize adult cats weighing 2-5 Kg. Occasionally additional nembutal was required. We preferred to keep the cats in a light surgical anesthesia (Stage III). A check of the pedal and corneal reflexes were made to determine if additional nembutal was required. The booster case, if required, was given intravenously in diluted

FIGURE 4

TYPICAL VARIATIONS IN POLLUTANT CONCENTRATIONS

DURING AN EXPERIMENT



TYPICAL SYSTEM PERFOR-MANCE: SO₂ EXPOSURE CONCENTRATION VS. TIME

ethiodide, a muscle relaxant, was administered. It produces a nondepolarization block at the neuromuscular junction. The dosage was 0.05 mg/Kg, intravenously.

Two surgical procedures were used. In the first method, a tracheotomy was performed by using a Bard Parker #3 holder with #15 rib back blade. A 1 inch incision was made approximately 20 mm below the larynx in the center of the ventral region of the throat. The muscle was separated by using two small hemostats which were spread in opposite directions until the trachea was visible. A slightly larger hemostat was inserted under the trachea and was then lifted on the opposite side, where at this time a 6" length of suture was attached. This hemostat was left in position. A small horizontal cut was made between the cartilage rings of the trachea and a suitable size cannula was inserted. The cannulas which were used were 6.35 mm i.d. and 7.98 mm i.d. with 7/16" diameter side air tube. The hemostat was withdrawn after bringing 3" of the suture to the opposite side of the trachea and tying the cannula to hold it intact. In vagotomized cats we tied off the vagus nerve before inserting the cannula. The side air tube of the cannula was connected to a Fleisch Pneumotachograph (0), which in turn was connected to a Model #607 Harvard Animal Breathing Pump.

The second method utilized similar procedures, but in this case an endotracheal tube was placed at the entrance of the larynx. The epiglottis was sighted with the aid of a laryngscope and was then held open by a small hemostat. A #16 Foregger endotracheal tube

(i.d. 2.5 mm, o.d. 5.3 mm), previously interlined with Teflon and having a built-in cuff, was then placed into the mouth of the larynx and the hemostat was released from the epiglottis. The cuff was inflated, clamped with a hemostat, and the tube was connected to the air pump in the usual manner. Whenever excessive mucus was found, suction was used to remove it.

A third and final preparation was used where a tracheotomy was initially performed, as described in the first method. A second cannula (same size) was inserted to provide a pathway to the nasopharyngeal chamber of the animal. The side air tube of the cannula was connected to an exhaust outlet. A face mask made of Lucite and anatomically sculptured for perfect fit was placed on the animal's face and was held in place with adhesive tape. The Teflon tubing nipple of the mask was connected to a small pump for flushing.

Preparation for intrapleural catheter insertion was initiated by a 5 mm incision in the right lateral thoracic region between the fourth and fifth rib. We used two small hemostats to probe and spread in opposite directions until we entered the intrapleural space. The hemostat was held in position immediately upon entry into the thoracic cavity. A #10 Malecot intrapleural catheter (.089" i.d.) was clamped onto the second hemostat and in a concerted movement one hemostat was withdrawn and the other, with the attached catheter, was inserted. The incision was sealed with wound clips. The catheter was connected to the side arm of a Statham differential strain gauge (Transducer Model No. PM 5 ± 0.2-350). The other arm of the strain gauge was connected to an opening on the side of the tracheal cannula.

In order to record blood pressure, intramedic polyethylene tubing (.045" i.d.) was inserted into the femoral artery and was then connected to a Statham strain gauge (Model No. P23Db series pressure transducer). The pressure dome of the transducer was filled with 1/200 solution of heparin and sodium chloride. The same size catheter was inserted into the femoral vein for administering drugs. Blood loss from surgery was minimal.

At the conclusion of inhalation exposures, the cat was sacrificed by administering an additional dose of nembutal. The thorax was opened by making a midline incision in the ventral thoracic region. The muscle was separated, costal cartilage dissected, sternum and eight pairs of ribs were removed. Portions of all lung lobes were exposed. We applied a modified Staub freeze technique (12) using Freon 12 (Dichlorodifluoromethane). Freon 12 was chosen because it was not hazardous, it required no extensive preparations for use, and good frozen sections were obtained.

D. Determination of Pulmonary Flow Resistance and Lung Compliance
The surgical preparations described above were followed by the
hook-up of Statham transducers to the Electronics for Medicine
Model DR-8 amplifiers for oscilloscope readout. The signals monitored and their respective transducers are shown in Table 1.

The signals from the DR-8 amplifiers or integrators could be visually read on an integral oscilloscope or recorded on LW-27, 18 cm photographic paper. A second oscilloscope provided visual display of R_L and C_L loops. These could also be permanently recorded on the photographic paper. Dry records were obtained in four seconds by means of a Rapid Writer attachment.

PARAMETERS RECORDED ON THE ELECTRONICS FOR MEDICINE UNIT AND ASSOCIATED TRANSDUCERS

TABLE 1

Parameter	Parameter Transducer	
Intrapleural Pressure (TPP)	Statham Model No. PM 5 + 0.2-350 Serial No. 12394	+ 0.2 psid
Air Flow (V)	Fleisch Pneumotachograph with Stat- ± 0.15 psid ham Model No. PM 283TC ± 0.15-350	
Volume of Air (V)	Signal from V is electronically in- tegrated	
Blood Pressure	Statham Model No. P23Db. Serial No. 11680	

Representative sweep and loop tracings from the Rapid Writer are shown in Figure 5.

Figure 6 is a photograph which shows the test cat connected to the Harvard Pump and the pneumotachograph in position.

In these studies animals were ventilated at 50-80 cc per stroke at a frequency of 18-21 strokes per minute. Thus, on the basis of an average tidal volume of 26 cc for the cat, these animals were hyperventilated.

When using oscilloscope loops, the total lung resistance was measured by subtracting a voltage proportional to lung volume change (and this proportional to compliance pressure) from the pressure axis of a pressure flow trace recorded on the oscilloscope screen (13). The voltage subtracted was sufficient to close the loop. The slope of the line resulting from closing the loop represented total lung resistance, which includes airway resistance and the viscous resistance of the lung tissue.

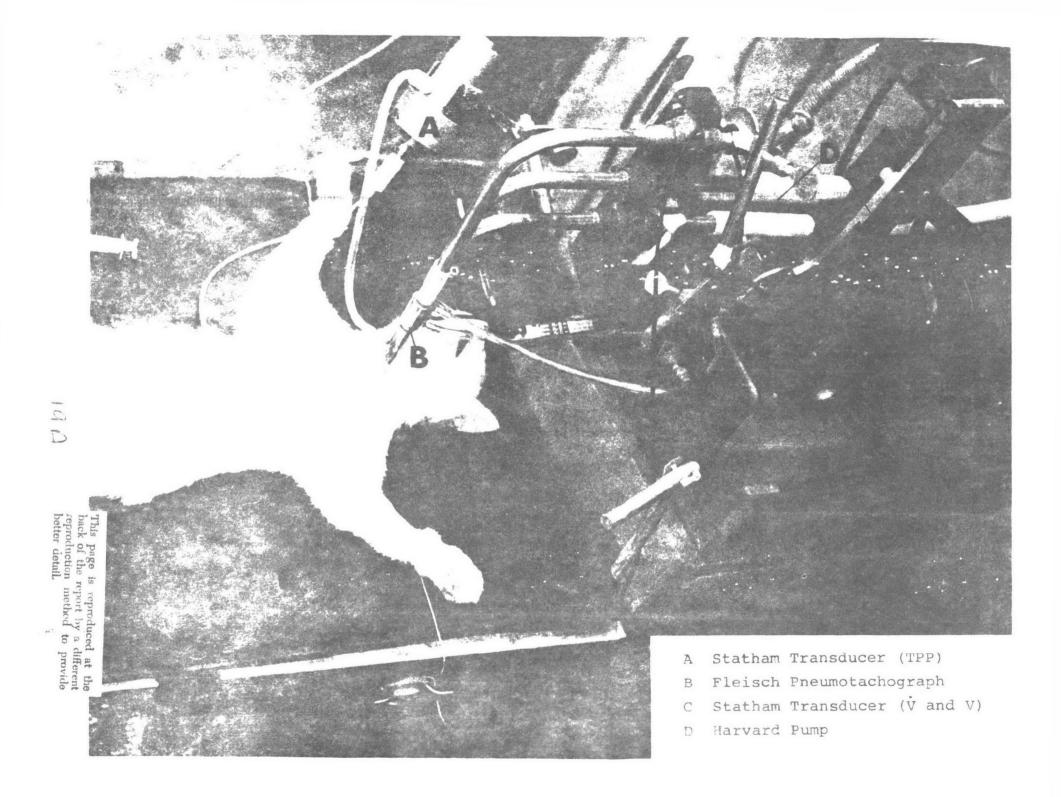
Calibration of all transducers was performed at the completion of each experiment. The correspondence between water or mercury manometers as primary standards and the LW-27 photographic paper readout of BP and TPP was recorded. Air Flow (v) was calibrated by utilizing compressed air and a rotameter calibrated against a wet test meter. Volume was calibrated by alternately depressing or withdrawing the barrels of two opposed 100 ml syringes in a closed air loop with the Fleisch Pneumotachograph; 20 ml increments of volume yielded a step function on the record chart. Loop signals were calibrated in a similar manner, except that signal deflections were measured directly on the oscilloscope face.

FIGURE 5

REPRESENTATIVE SWEEP AND LOOP TRACINGS USING E FOR M RAPID WRITER AND LW-27, 18 cm PHOTOGRAPHIC PAPER

FIGURE 6

PHOTOGRAPH SHOWING CAT ARTIFICIALLY VENTILATED BY
HARVARD PUMP



The dead space of the tracheal cannula and Fleisch Pneumotachograph, i.e. volume from bifurcation above Pneumotachograph leading to Harvard pump inlet or discharge, to the trachea of the test animal, was 4.7 cc. A correction was subtracted from each value of total lung resistance in order to correct for the resistance of the tracheal cannula.

E. Conduct of Experiments

Each animal acted as his own control with respect to resting state values of TPP, V, V, R_L, and C_L. The criterion for significant changes in these parameters following inhalation of polluted air was that the values of these parameters after exposure should be different from the values of these parameters in the same animal before exposure. This approach requires that care be taken to ensure that the animal is in a stable resting state prior to being challenged by a polluted atmosphere. Another criterion of these studies was that following exposure an animal had to return to his initial, preexposure values of these parameters before he was challenged with another polluted atmosphere. These are stringent experimental criteria which resulted in long experiments.

Immediately following surgical preparations the animal was maintained in a resting state for approximately one half hour. This was followed by a 15 minute control period, with recorded records at five minute intervals and continuous sweep visual display. The animal was then tested by mechanical stimulus to ensure the intactness of the vagally mediated reflexes for control of airway constriction (14). Initially, the vagii were surgically

isolated and an electrical stimulus was applied, but the mechanical stimulus was judged to be equally effective and less traumatic. A further control period of 15 to 60 minutes followed the mechanical stimulus test. If the vagii were not responsive, as judged by an immediate increase in R_L, then anesthesia was too deep and the animal remained in a resting state until a positive test was obtained for intactness of vagal pathways. This procedure of testing was repeated prior to the first, and after each succeeding pollutant challenge.

The order of exposures in these studies was varied in order to rule out the possibility of a long-term effect due to initial exposure to High SO₂, for instance. Thus, certain animals were initially exposed to High SO₂ (following control period), and then to Low SO₂ or the SO₂-aerosol mixture. All orders of exposure were used. After initial exposure to High SO₂, a 1-2 hour resting period was required to permit the animal to meet the criterion established for return to initial resting values of parameters.

Another series of experiments was performed to determine if receptors which could affect airway size were being bypassed by introduction of pollutant gases (in breathing air) by means of the tracheal cannula. These tests were made with an endotracheal catheter. In this case, the animal was ventilated by the Harvard Pump and the tracheal cannula, but room air (for control) or pollutant laden air was simultaneously flushed through the endotracheal catheter by means of a small pump delivering 2.4 cc/stroke.

Only Low SO, concentrations were delivered in these studies.

In a similar manner, a series of tests was performed to determine if receptors in the nose could cause airway constriction. The usual procedures were followed, but another cannula was inserted back to back to the cannula flushing upwards into the mouth. A mask was placed over the nose and mouth. A small pump ventilated the mask while the second cannula served as the exhaust. Thus, the airway above the tracheal cannula was being flushed via the nose and mouth. The lower airways received breathing air or pollutant gas via the original tracheal cannula. Low and High SO₂ in air were used in this series of experiments (Figure 7).

F. Treatment of Data

Values of R_L and C_L were calculated from sweep tracings of \dot{V} , V and TPP as described in the original method $^{(1)}$, or the values were obtained from reading recorded loops. R_L and C_L values were then graphed as a function of time. Although continuous records were made, it was necessary to establish the intervals for transcribing data in order to show experimental trends. Figure 8 is a representative graphical summary showing values of $:R_L$ and C_L at five minute intervals. In this case each point represents a series of breathing cycles from the sweep record and clearly shows the stability of these parameters during control periods.

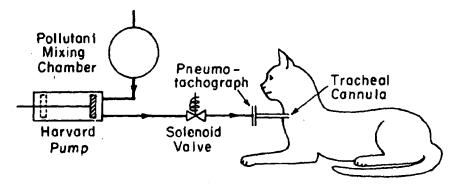
Table 2 contains the numerical results obtained from the experiment on Cat #1610.

FIGURE 7

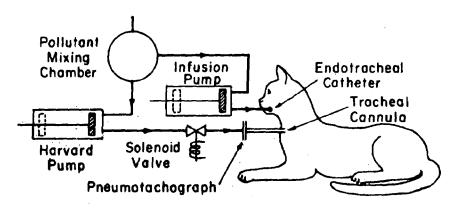
CANNULA ARRANGEMENTS FOR EXPOSURE VIA TRACHEAL CANNULA AND FOR ABOVE AND BELOW TRACHEAL CANNULA

FIGURE 8

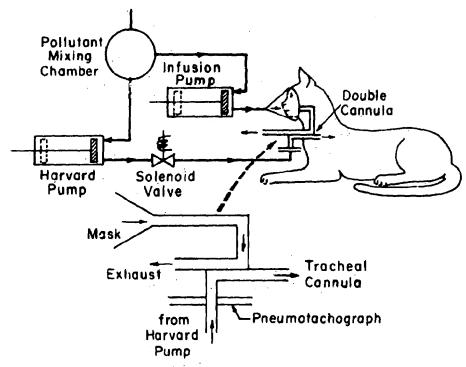
GRAPHICAL SUMMARY OF VARIATIONS IN R_L AND C_L DURING AN EXPERIMENT



(a) Via Tracheal Cannula

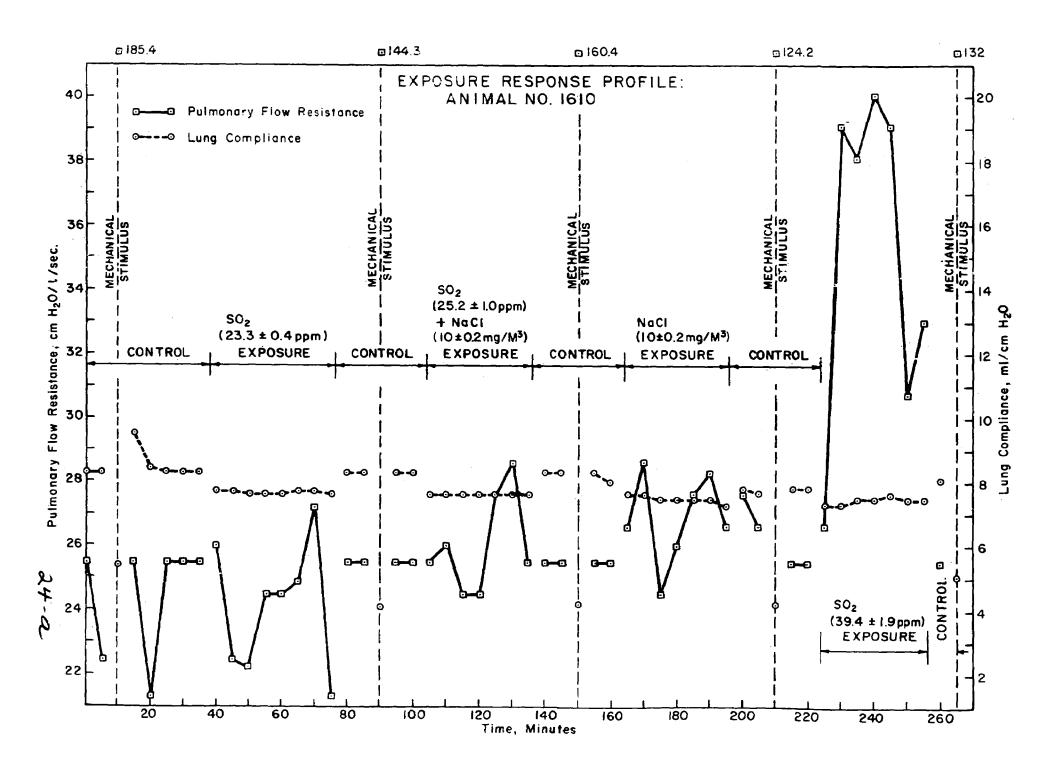


(b) Via Tracheal Cannula and Endotracheal Catheter



(c) Via Naso-pharyngeal Chamber

23-a



Final calculations are based on data from the final fifteen minutes of the control and exposure periods. Other time periods were evaluated, but the last fifteen minute period proved to be the best indication of response. The table shows the mean and the standard arithmetic deviations of parameters during this time period. The percentage change in parameters noted in Table 2 was calculated from control period values as a baseline. Thus,

Tabular and graphical summaries of individual experiments conducted during this study are contained in the Results and Discussion section of this report.

Before an animal's response was termed "significant" it was necessary to compare R_L and C_L values obtained during the last fifteen minutes of the control and exposure periods. The comparison was based on a modified Student t-test for the difference of means. The degrees of freedom (d.f.) for the test was calculated from the variances and sample sizes. Thus,

$$t = \frac{\bar{x}_2 - \bar{x}_1}{\sqrt{s_1^2/n_1 + s_2^2/n_2}}$$

$$d.f. = \frac{[(s_1^2/n_1) + (s_2^2/n_2)]^2}{(s_1^2/n_1)^2/(n_1 + 1) + (s_2^2/n_2)^2/(n_2 + 1)} - 2$$

where mean values are designated x_1 and x_2 , the standard deviations are S_1 and S_2 , and sample sizes are n_1 and n_2 .

The level of significance accepted as meaningful with the ttest was P < 0.01 (two tailed test).

TABLE 2 Cat No. 1610 RESPONSE TO VARIOUS STIMULI EXPRESSED AS PERCENT CHANGE RELATIVE TO CONTROL VALUES*

	Pu	1monary Flow Resistance	(6)	
Medh. % Control Stim. Change	Low 본 Control SO ₂ Change	Mech. ½ Control Stim. Change	SO2+ ½ Control NaCl Change	Mech. ½ Control Stim. Change
25.5 185.4 674.1 22.4	25.5 24.9 -2.5 25.5 27.2 5.7 25.5 21.3 -16.5	25.5 14/4.3 -65.9 25.5	25.5 27.6 8.2 25.5 28.6 12.2 25.5 0	25.5 160.4 529.0 25.5
Mean± 24.0 185.4 S.D. 2.2	25.5 24.5 0 3. 0	25.5 144.3 0	25.5 27.2 0 1.6	25.5 150.4 0
REFLEX INTACT	ΔN.S.	REFLEX INTACT	ΔN.S.	REFLEX INTACT
		Lung Compliance (c)		
Mech. % Control Stim. Change	Low 1/ Control SO ₂ Shange	lech. ½ Control Stim.Change	SO ₂ + % Control NaCl Change	Mech. % Control Stim. Change
9.3 5.4 -34.9 9.3	8.3 7.7 -7.2 8.3 7.7 -7.2 8.3 7.6 -9.4	E.3 4.1 -50.6 B.3	8.3 7.6 -8.4 8.3 7.6 -8.4 7.6 -8.4	8.3 4.2 -49.4 8.3
reari 9.3 5.4 5.0. 0	8.3 7.6 0 0.1	8.3 4.1	3.3 7.6 0 0	8.3 4.2 0
	P<0.01		P< 0.01	
Sequence of (1)	(2)	(3)	(4)	(5)

*Control for 15 minutes preceding challenge.

(a) See Figure.

(b) Pulmonary Flow Resistance, Cm H₂0/1/sec. (c) Lung Compliance, ml/cm H₂0.

 \triangle N.S. = Difference between means not sigificant (P>0.05)

TABLE 2 (Continued) Cat ilo. 1610 RESPONSE TO VARIOUS STIMULI EXPRESSED AS PERCENT CHANGE RELATIVE TO CONTROL VALUES*

		Pu	lmonary Flow Resistance	(b)	
Control NaCl	्रं Change	Mech. % Control Stim. Change	High % Control \$0 ₂ Change	Mech. ½ Control Stim. Change	
25.5 27.6 25.5 23.3 26.6	9.2 11.0 4.3	27.6 124.7 353.3 26.6	25.5 39.1 53.3 25.5 30.7 20.4 33 29.4	25.5 132 417.6	
Meari 25.5 27.5 S.D. 0 0.8		27.1 124.7 0.7	25.5 34.3 0 4.3	25.5 132	
<u> </u>		REFLEX INTACT	Δ N.S.	REFLEX INTACT	
			Lung Compliance (c)		
Control NaCl	% Change	Mech. Control Stim. Change	High % Contro! \$02 Change	Mech. 火 Control Stim. Change	
8.3 7.5 8.0 7.5 7.3	-8.0 -9.0 -10.4	7.8 4.2 -45.8 7.7	7.8 7.6 -2.6 7.8 7.5 -3.8 7.5 -3.8	9.1 5.1 -37.0	
Mean± 9.2 7.4 S.D. 0.2 0.1		7.7 4.2 0.1	7.8 7.5 0 0.1	8.1 5.1	
P< 0.05			P< 0.05		
Sequence of (a)	(6)	(7)	(8)	(9)	

*Control for 15 minutes preceding challenge.

(a) See Figure.
(b) Pulmonary Flow Resistance, Cm H₂0/1/sec.
(c) Lung Compliance, m1/cm H₂0.

A N.S. = Difference between means not sigificant (P>0.05)

G. Pathological Procedures

Six weeks after fixation tissues were removed from freezer and Carnoy's fluid poured off. The Storey and Staub (15) method was used to process and stain most of the lung tissues except for a modification as follows:

1. Nitrocellulose embedded tissues were sectioned 75 µm thick on a Reitchert sliding microtome. Using Lundy's method, serial sections were stapled in sequence, approximately 5 mm apart (by means of an office stapling machine), on a strip of 500 gauge polyethylene film, cut slightly wider than the width of the sections. The length of the strips was determined by the size of our staining dishes (Pyrex Baking Dish 5" x 9"). After the tissue was stained, sections were detached from the film with scissors and mounted in sequence on numbered slides. Strips of polyethylene film holding sections not immediately required for staining were rolled up and stored in 70% alcohol.

In order to obtain thin sections, a small piece of each lung tissue was put into Bouin's fluid for 24 hours for further fixation. A routine manual process was used to obtain paraffin blocks. Blocks were sectioned 5 µm on an American Optical rotary microtome. Serial sections were stained with Harris Hematoxylin and Eosin and mounted on glass slides with permount.

We photographed serial sections 5 µm thick and 75 µm thick with a Zeiss photomicroscope. To obtain a total estimated magnification of 130x, a planachromate 2.5x objective with an N.A. = 0.08, optavar = 2.0x, intermediate camera magnification

of 3.2x and photographic enlargement estimated at 8.0x was used.

A calibrated field finder was used in returning to the same
photographic field of each slide in the series.

Measurements were made of the alveoli on the enlarged prints of thick tissue sections. We noted in our findings that there were no significant changes in the sizes of the small conducting airways of cats which had shown small $R_{\rm L}$ and small $C_{\rm L}$ changes.

In an attempt to measure the same segment of lung lobe of each cat we measured 2 cm from the tip of the third lobe of the right lung and removed that portion of the lobe for frozen sectioning and for comparison. Biological factors such as weight and size of the lungs presented a problem. We questioned whether we could be sure that we were in the same segment of the lobe, when some of the lungs were larger than others. Despite this obvious flaw, we cut the same distance from the tip of the same lobe of each animal to obtain comparable sections. We realized, however, that we had encountered a source of error in using a method of measurement on a histologic section.

Thin lung sections were studied, of cats that had been challenged with ${\rm SO}_2$ in the air and ${\rm SO}_2$ + NaCl aerosol in medical grade breathing air. The absence of comparable changes in cell structures suggested that small R_L changes and small C_L changes could not be detected histologically.

III. RESULTS AND DISCUSSION

All calculated results for individual experiments are contained in tabular and graphical summaries in this Section. The results presented were obtained by extracting data from continuous records as described in Section II F., above. Results are presented for a total of twenty-nine complete experimental protocols in which twenty-nine different cats were used. A discussion of this vast amount of data requires that a simple summary be used to present findings. The reader can examine calculated results of individual animal experiments to substantiate generalizations made in this discussion.

A. Changes in Pulmonary Flow Resistance and Lung Compliance in Cats Following Pollutant Challenges

Table 3 summarizes the challenge atmospheres which produced significant changes in pulmonary flow resistance or lung compliance when the pollutant mixture was administered via tracheal cannula. Table 4 is a similar summary for pollutants administered via endotracheal catheter and via double cannula to produce pollutant flushing of the nasopharyngeal chamber and any receptors above the trachea cannula. First, the data suggest that with the cat, approximately 20 ppm $\rm SO_2$ in air are required before any animals show significant alterations in $\rm R_L$. Prior to the studies reported here, it had been shown (7) that pure $\rm SO_2$ delivered into the lower airways and lungs during a single respiratory cycle increased $\rm R_L$ (mean, + .246 percent; P < 0.05). The animal returned to control levels within one minute (7). The work reported here

indicates that with the same physiological preparation approximately 20 ppm $\rm SO_2$ in air are required to trigger this response—and at this concentration only two animals in twenty responded. One animal in twenty showed a significant decrease in $\rm R_L$ at this concentration.

It is interesting to compare concentrations of SO, in air required to cause bronchoconstriction in different species. Frank, et al. (16) demonstrated a significant increase in resistance to air flow in volunteers exposed to 5 ppm. This finding is consistent with that of Burton, et al. (Appendix I) of perhaps one out of ten "human reactors" to approximately 3 ppm SO, in air. Balchum, Dybicki and Meneely (17) exposed ten anesthetized dogs to concentrations of SO, in air ranging from 1.8 to 148 ppm for periods from 30 to 40 minutes. Increases in the nonelastic resistance to breathing ranged from 150 to over 300% and these increases occurred within nine seconds after the onset of breathing SO2; increases disappeared as quickly following the end of exposure. Exposures of guinea pigs to SO2 in air for one hour results in a 10% increase in preliminary flow resistance at 0.16 ppm (18). The guinea pig, as used by Amdur, "may be the accidental analog of the sensitive segment of the population" (3). The results reported here suggest that the cat, as used in our preparation, is an analog of the more resistant segment of the population.

An aspect of our studies which was stressed by Frank, et al. (15) and Burton, et al. (Appendix I) in studies involving human exposures but was not stressed in studies employing dogs or guinea pigs,

is the great variability of response of individuals. In our studies, "reactors" were characterized by large variability of response in R_L and C_L and hypersensitivity during preparation. Examination of graphical summaries of results of exposures will demonstrate the great variability of response for animals with cervical vagosympathetic nerve conduction intact.

Two other characteristics of response should be noted for their contrast with guinea pig studies. Our animals returned to control levels shortly after exposure, although we often waited for an extended period to reduce the range of variation about the mean R_L or C_L following exposure, i.e. variability about the mean was greater following exposure, which we did not desire. We chose to permit the animal to settle down. The guinea pig, on the other hand, returns slowly to control levels after exposure. As noted above, following exposure human subjects and dogs also return rapidly to control values of resistance to air flow.

In animals showing changes in pulmonary flow resistance, we confirmed the effects of cooling the vagii, injection of atropine or severing the vagii, on blocking these changes, as reported by Nadel, et al. (7)

Table 4 summarizes results collected to focus on the sites of receptors responsible for bronchoconstriction or peripheral airway constriction. These results relate to findings of nearly total uptake of SO_2 in air in the nose and upper airways in animals $^{(19,20)}$ and in human subjects $^{(21)}$. Less than 1% of the inhaled concentration of SO_2 in air is estimated to

reach the larynx and more distal airways in man (20). The endotracheal catheter and double cannula were used in these studies to determine if the number of animals responding increased, or the degree of response increased in respondents when the pollutant mixtures were administered via these pathways in cats. Three animals were exposed with the endotracheal catheter and two with the double cannula. Unfortunately, these experiments must be contrasted to findings in twenty cats where a tracheal cannula was used for delivery of pollutant mixtures. However, comparison of Tables 3 and 4 suggest that delivery of pollutant via tracheal cannula did evoke fewer significant responses in R, or C, thus suggesting that certain receptors were bypassed when pollutant was delivered by tracheal cannula. Thus, our findings suggest that receptors in the nasal pharyngeal chamber and proximal to the tracheal cannula used here can increase total lung resistance distal to the tracheal cannula. This finding contrasts with that by Nadel and Widdicombe (22) that mechanical irritation of the nasal mucosa did not change total lung resistance in cats anesthetized with chloralose and urethan. Additional cats should be studied to substantiate our findings.

Small but insignificant changes in lung compliance were found in a few animals following each of the challenges presented, including one animal who showed a significant decrease in $C_{\rm L}$ following inhalation of NaCl aerosol. These results suggest that the physiological mechanisms generally delineated for upper airway

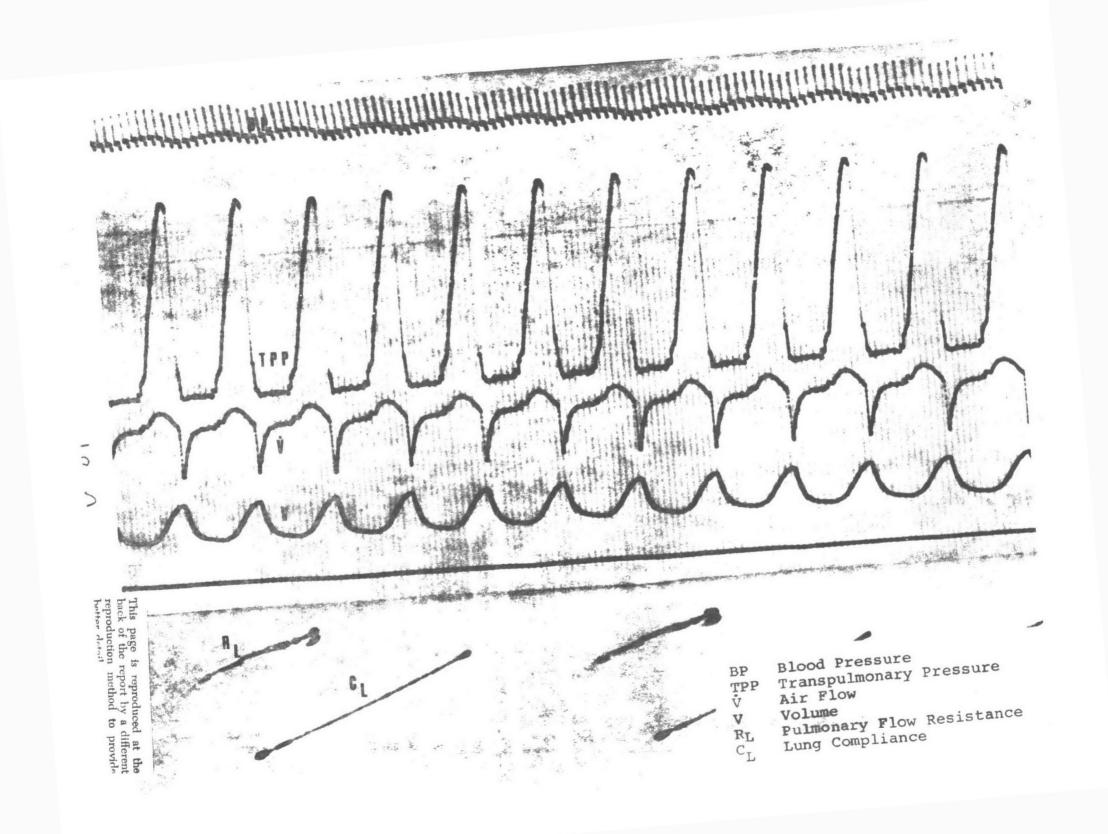
and for peripheral airway constriction by acute challenges of physical or chemical agents may not be completely independent. Other investigators have suggested that a reflex mediated by vagal efferent fibers could partially contribute to peripheral airway changes (8). Alternately, the small quantities of pollutants which penetrated the upper airways may be sufficient to cause peripheral airway constriction, but this possibility, although suggested elsewhere (21), seems remote to us as a mechanism for peripheral airway constriction.

Finally, it should be noted that while upper airway changes after inhalation of pollutants could be due to mucous secretion, cooling of vagii, intravenous atropine or deep anesthesia blocked these changes, thus suggesting that they were due to changes in airway calibre.

B. Changes in Pulmonary Flow Resistance and Lung Compliance in Guinea Pigs Following Pollutant Challenges

The rapid return of C_L and R_I, values in cats, dogs and man following exposures to pollutants used in this study differs from the slow recovery of guinea pig pulmonary flow resistance following similar exposures. Therefore, we used the exact experimental preparations described above to study the responses of guinea pigs. Animals weighing approximately one kilogram were used to permit insertion of a tracheal cannula; a difficult procedure with small animals.

It was not possible to obtain valid data for guinea pigs with our preparation. The animals secreted abundant mucous



SUPPLARY OF R_L^a AND C_L^b RESPONSES (P > 0.01) OF CATS EXPOSED TO POLLUTANT MIXTURES IN THIS STUDY (TRACHEAL CANNULA POLLUTANT DELIVERY)

hallenge	Low	Concentr	ation of	f \$02 i	n Airc	so ₂ d	and	NaCl	Aero	sole ;	n Air	NaC.) Aer	osol	in	ire	High	Co	ncent	rati	on S	02 i	n Air	T
Cat		RL		CL			RL			c_{L}			RL		C				RL				L	_
Number	+			<u> </u>		+			+			+			<u>+</u>		+					+		
14629	į					1						l					i				x			
1117	1																•							
1112	×											1					×							
1113	l				:	1						1]							
1564	1											i					•				×			
1566	}				×							l					1							
1144	×					l						l					ł							
1593	ł					l						1					ł						×	ž.
1606	i											1					ł							
1611	l					1						1					(
1610	İ				×	l					×	ļ					ł							
1633	1					ł						1					(
1612	1				×)			×		1							
1609	1					Ì						1)							
1651	ł			×		İ						ł												
5354g,i																								_
2135g,i	1											i					1							
1486g,i	ļ		x		x :						×	1					ì							
1487g,i	1				×	l					×	I					l							
1676g,i	1				x) x					×							
1781h	!											1					1						×	_
1801h	i				İ							1				ж	1							
1988h	1				×	ŀ					×	1					1				x			
1807h	ł				x						- "	1				ж	i					х		

 $^{^{}a}R_{L}$ = Pulmonary Flow Resistance, Cm H₂0/1/sec (+ denotes increase, - denotes decrease).

 b_{C_L} = Lung Compliance, m1/Cm H₂0 (+ denotes increase, - denotes decrease).

 c_{SO_2} in air concentration for all exposures, expressed as Mean \pm S.D. = 19.0 \pm 5.9 ppm.

 $^{^{}d}$ SO₂ in air concentration for all exposures, expressed as Mean \pm S.D. = 17.9 \pm 8.3 ppm.

eNaCl aerosol in air concentration, expressed as Mean \pm S.D. = $10 \pm 0.2 \text{ mg/m}^3$.

fso, in air concentration for all exposures, expressed as Mean + S.D. = 34.6 + 12.3 ppm.

 g_{In} the following experiments, the Low Concentration of SO₂ in air was the first challenge followed by Control Period SO₂ NaCl Aerosol, etc.

hIn the following experiments, the High Concentration of SC₂ in air was the first challenge, followed by Control Period NaCl Aerosol, etc.

¹These animals were characterized by extreme values of SO₂ concentration.

TABLE 4 SUMMARY OF R_L^a AND C_L^b RESPONSES (P 0.01) OF CATS EXPOSED TO POLLUTANT MIXTURES IN THIS STUDY (ENDOTRACHEAL CATHETER POLLUTANT DELIVERY)

Challenge	so ₂	in	Air	Via	Tracheal	Cannula	so ₂	in	Air	Via	Endotracheal	Catheterd	so ₂	in	Air	Via	Both	Site	se
Cat Number	+	RL		+	C _L		+	RL		+	C _L	_	+	R _L	_	+	CL		
2984 2206 53859			×			x x						×			x			:	x x

SUMMARY OF R_T AND C_T (NASO-PHARYNGEAL FLUSH WITH DOUBLE CANNULA)

Challenge	Low Concentration of SO ₂ in Air ^f	High Concentration SO ₂ in Air ^g
Cat Number	+ C _L -	+ R _L C _L -
2393 5674		

 $^{^{}a}$ R $_{L}$ = Pulmonary Flow Resistance, Cm H $_{2}$ 0/1/sec (+ denotes increase, - denotes decrease). b C $_{L}$ = Lung Compliance, m1/Cm H $_{2}$ 0 (+ denotes increase, - denotes decrease).

 $^{^{\}text{C}}\text{SO}_2$ in air concentration for all exposures, expressed as Mean \pm S.D. = 18.2 \pm 1.9 ppm.

 $^{^{}d}$ SO₂ in air concentration for all exposures, expressed as Mean \pm S.D. = 17.2 \pm 0.9 ppm.

 $^{^{}e}_{50}^{2}_{2}$ in air concentration for all exposures, expressed as Mean \pm S.D. = 17.1 \pm 0.9 ppm

 $^{^{}f}$ SO₂ in air concentration for all exposures, expressed as Mean \pm S.D. = 14.4 \pm 1.8 ppm.

 $^{^{9}}SO_{2}$ in air concentration for all exposures, expressed as Mean \pm S.D. = 22.5 \pm 2.9 ppm.

which required frequent withdrawal by catheter connected to a suction source. The changes in pulmonary mechanics which stemmed from mucous secretion in control animals would have overwhelmed any changes associated with airway calibre alterations due to pollutants. Mucous secretion under these conditions cannot be compared to that which may occur during spontaneous breathing during exposure to pollutants, the conditions for Amdur's studies $^{(3)}$. However, the slow return to control values of R_L by guinea pigs strongly suggests mucous secretion as a contributor to R_L , a hypothesis which could be easily tested experimentally.

C. Pathological Changes in Cats Following Pollutant Exposures
Rapid lung freezing procedures and preparation of samples
were described above. While alterations in airway calibre
could be detected by this method following severe acute pollutant challenges (8), it was not possible to detect differences
in airway calibre in thick and thin lung sections of control
and exposed animals in the studies reported here. A control
animal was one previously exposed to SO₂ or SO₂-aerosol mixture
in air, but whose pulmonary flow resistance subsequently returned
to the preexposure value. Figures 9 and 10 show the close correspondence between photomicrographs of thick lung sections from a
control and an exposed animal, respectively. Figure 11 is a
photomicrograph of a typical thin lung section.

The work of Macklem and Mead (23) stimulated these attempts to detect changes in peripheral airway size when alterations in pulmonary flow resistance were present. Macklem and Mead

demonstrated that at high lung volumes, R_L increased and this resistance was almost entirely due to that between airways 1.5-2.5 mm and the trachea in dogs. Thus, large changes in peripheral airways resistance could go undetected by measurement of R_L , which is insensitive to alterations in peripheral airway resistance. Our studies show that careful morphological examination of airway calibre are also insensitive to any changes that may occur in the ranges of R_L increases cited here.

FIGURE 9

PHOTOMICROGRAPH OF A THICK SECTION OF THE RIGHT LUNG LOBE OF A MALE CAT (NO. 1112). FROZEN IMMEDIATELY AFTER CAT HAD RETURNED TO CONTROL STATE FOLLOWING 15 MINUTES PREVIOUS EXPOSURE TO 27 PPM SO₂.

39 A

FIGURE 10

PHOTOMICROGRAPH OF A THICK SECTION OF THE RIGHT LUNG LOBE OF A FEMALE CAT (NO. 1144). FROZEN IMMEDIATELY FOLLOWING EXPOSURE TO 40 PPM SO₂ FOR 30 MINUTES.

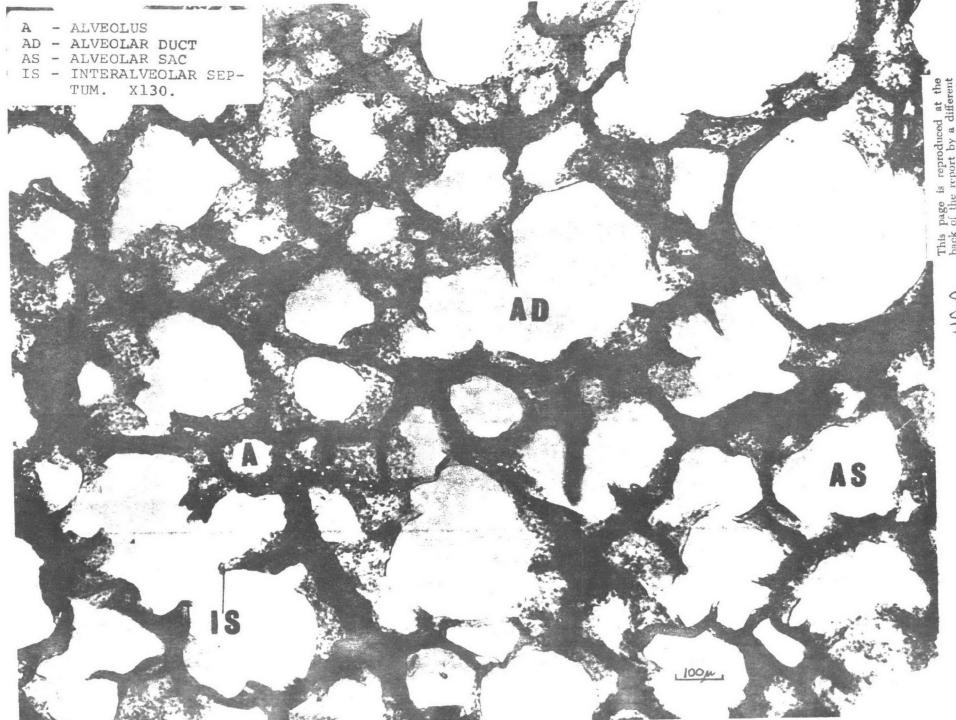


FIGURE 11

PHOTOMICROGRAPH OF A THIN SECTION OF THE RIGHT LUNG LOBE OF A MALE CAT (NO. 1610). FROZEN AFTER CAT HAD RETURNED TO CONTROL STATE FOR 15 MINUTES FOLLOWING PREVIOUS EXPOSURES TO 40 PPM SO₂.

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REFERENCES

- 1. Amdur, M. O. and Mead, J.: "Mechanics of Respiration in Unanesthetized Guinea Pigs". Am. J. Physiol. 192, 364 (1958).
- 2. Air Quality Criteria for Sulfur Oxides. U. S. Dept. Health, Education, and Welfare, Public Health Service, Consumer Protection and Environmental Health Service, National Air Pollution Control Administration, Washington, D. C. January, 1969. NAPCA Pub. No. AP-50.
- 3. Amdur, M. O.: "Toxicological Appraisal of Particulate Matter, Oxides of Sulfur, and Sulfuric Acid". J. Air Pollution Control Assoc. 19, 639 (1969).
- 4. Widdicombe, J. G.: "Respiratory Reflexes from the Trachea and Bronchi of the Cat". J. Physiol. 123, 55 (1954).
- 5. Widdicombe, J. G.: "Receptors in the Trachea and Bronchi of the Cat". J. Physiol. 123, 71 (1954).
- 6. Widdicombe, J. G., Kent, D. C. and Nadel, J. A.: "Mechanism of Bronchoconstriction during Inhalation of Dust". J. Appl. Physiol. 17, 613 (1962).
- 7. Nadel, J. A., Salem, H., Tamplin, B. and Yokiwa, Y.: "Mechanism of Bronchoconstriction". Arch. Environ. Health 10, 175 (1965).
- 8. Nadel, J. A., Corn, M., Zwi, S., Flesch, J. and Graf, P.:
 "Location and Mechanism of Airway Constriction after Inhalation of Histamine Aerosol and Inorganic Sulfate Aerosol".
 In Inhaled Particles and Vapors, II, Davies, C. N., Editor,
 Pergamon Press, New York, 1966. p. 55.
- 9. West, P. W. and Gaeke, G. C.: "Fixation of Sulfur Dioxide as Disulfitomercurate (II) and Subsequent Colorimetric Estimation". Anal. Chem. 12, 1816 (1956).
- 10. Pate, J. B., Ammons, B. E., Swanson, G. A. and Lodge, J. P.: "Nitrite Interference in Spectrophotometric Determination of Atmospheric Sulfur Dioxide. Anal. Chem. 37, 942 (1965).
- 11. Tentative Method of Analysis for Sulfur Dioxide Content of the Atmosphere (Colorimetric). Intersociety Committee Methods for Ambient Air Sampling and Analysis. Health Laboratory Science 6. 228 (October 1969).

- 12. Staub, N. C. and Storey, W. F.: "Relation Between Morphological and Physiological Events in Lung Studied by Rapid Freezing". J. Appl. Physiol. 17, 381 (1962).
- Mead, J. and Whittenberger, J. L.: "Physical Properties of Human Lungs Measured During Spontaneous Respiration".
 J. Appl. Physiol. 5, 779 (1953).
- 14. Nadel, J. A.: "Mechanisms of Airway Response to Inhaled Substances". Arch. Environ. Health 16, 171 (1968).
- 15. Storey, N. C. and Staub, W. F.: "Ventilation of Terminal Air Units".). Appl. Physiol. 17, 391 (1962).
- 16. Frank, N. R., Amdur, M. O., Worchester, J. and Whitten-berger, J. L.: "Effects of Acute Controlled Exposure to SO₂ on Respiratory Mechanics in Healthy Male Adults". J. Appl. Physiol. 17, 252 (1962).
- 17. Balchum, O. J., Dybicki, J. and Meneely, G. R.: "Pulmonary Resistance and Compliance with Concurrent Radioactive Sulfur Distribution in Dogs Breathing S³⁵⁰2". J. Appl. Physiol. 15, 62 (1960).
- 18. Amdur, M. O.: "Respiratory Absorption Data and SO₂ Dose-Response Curves". Arch. Environ. Health 12, 729 (1966).
- 19. Dalhamn, T. and Strandberg, L.: "Acute Effects of Sulfur Dioxide on Rate of Ciliary Beat in Trachea of Rabbit in vivo and in vitro, with Studies on Absorptional Capacity of Nasal Cavity". Int. J. Air Water Poll. 4, 154 (1961).
- 20. Strandberg, L. G.: "SO₂ Absorption in the Respiratory Tract". Arch. Environ. Health 9, 160 (1964).
- 21. Speizer, F. E. and Frank, N. R.: "The Uptake and Release of SO₂ by the Human Nose". Arch. Environ. Health 12, 725 (1966).
- 22. Nadel, J. A. and Widdicombe, J. G.: "Reflex Effects of Upper Airway Irritation on Total Lung Resistance and Blood Pressure". J. Appl. Physiol. 17, 861 (1962).
- 23. Macklem, P. T. and Mead, J.: "Resistance of Central and Peripheral Airways Measured by a Retrograde Catheter". J. Appl. Physiol. 22, 395 (1967).

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Gene Bingham, University of Pittsburgh Veterinarian, demonstrated
and helped us to perfect the endotracheal tube methodology.

APPENDICES

- A. Reprint of "Response of Healthy Men to Inhaled Low Concentrations of Gas-Aerosol Mixtures" by Burton, G. G., Corn, M., Gee, J. B. L., Vasallo, C., and Thomas, A. P. AMA Arch. Env. Hlth. 18, 681 (1969).
- B. Tabular and Graphical Records of Individual Experiments.

Response of Healthy Men to Inhaled Low Concentrations of Gas-Aerosol Mixtures

George G. Burton, M.D.: Morton Corn. PhD; J. Bernard L. Goe: M.D.: Charles Wasalio, M.D.; and Armand P. Thomas, BA. Pittsburgh EXISTING laboratory studies dealing with the acute effects of inhaled pollutants in humans and experimental animals have recently been reviewed. These studies had failed to demonstrate changes in lung mechanics of healthy adults exposed to single pollutants at concentrations representative of those in ambient urban air. LaBelle et aland later Goetza had suggested that gasacrosol synergism might explain the hypothesized adverse effect on health, and the animal studies of Amdur et alaba gave the weight of considerable sound experimental data to the concept.

Several studies have been done, to date, to determine the presence or absence of gasacrosol synergism in man, and the results are conflicting. Frank et al⁹ using mixtures which consisted of SO₂ and a submicron NaCl acrosol, could demonstrate no synergistic effect in healthy adults over the range of 1 to 17 ppm SO₂. Also, no significant changes in pulmonary flow resistance (R₁) occurred during exposure to 1 to 2 ppm SO₂ with or without the added acrosol.

Later, Toyama' studied the effects of a wide range of concentrations of SO₂ alone and in combination with a monodisperse submicronic aerosol of NaCl (concentration, 7.4 mg/cu m). He concluded that a synergistic response producing increased airwing resistance was present, even at law concentrations of SO₂, though the number of

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From the Graduate School of Public Health and School of Medicine, University of Pittsburgh, Pa. Dr. Burton is now at the Lonn Linda University School of Medicine, Lonn Linda, Calif.

Read before the ninth AMA Air Pollution Medical Research Conference, Derven July 24, 1988.

Raprint requests to Department of Medicine, Section on Medical Chest Diseases, Long. Linda University School of Medicine, Long. Linda, Calif 92354 (Dr. Durfon).

Experimental Subjects													
Pollutants	6	7	8	10	9	1	2	3	5	4	Subjects		
SO ₂ (10 ft)	1.9	3.0 0.08	2.4 ± 0.15	3.2 .÷0.13	2.1 ± 0.00	2.2 ±0.00	1.4	1.2	1.9 ± 0.56	1.9 +0.00	2.1		
SO ₂ (20 ft)	1.9 ±0.13	3.1	2.8 ± 0.28	3.6 ±0.00	1.9 +0.06	2.1 ± 0.03	1? ±0.03	1.2	1.8 +0.22	1.7	2.1 +0.3		
Aerosol-Mix (10 ft)	1.9 + 0.09	3.0 · 0.00	2.3 ± 0.10	3.0 - 0.00	1.9 + 0.18	2.3 ± 0.10	1.4	1.2	1.4 + 0.07	1.1 -0.14	.1.0 +0.22		
Aerosol-Mix (2D It)	1.9 ± 0.63	3.0 ± 0.00	2.4 ± 0.22	3.3 +0.16	1.7 ± 0.11	1.9 ± 0.03	1.2	1.1 ± 0.08	1.8	1.8 + 0.08	2.0		
[NaCl]	2.4 mg/	2.0 mg/ cu m	2.0 mg/	2.7 mg/	2.1 mg/ cu m	2.5 mg/ cu m	2.3 mg/	2.1 mg/	2.0 mg/ cu m	2.2 mg/	0.08		

Table 1.- Exposure Concentrations of SO2 and NaC1 Aerosol*

subjects exposed to < 5.0 ppm SO₂ and aerosol was small.

At the sixth annual Air Pollution Medical Research Conference in 1963, Toyamas presented studies of eight healthy young men whom he exposed to SO₂ concentrations (3 to 40 ppm) with and without inhalations of Kawasaki industrial area dusts (concentration, 10 to 50 mg/cu m). Again, he concluded that a synergistic response could be demonstrated, though there were "fairly wide individual differences."

We decided to extend these studies by measuring airway resistance (\mathbf{R}_{Λ}) —a more easily performed test of irritant responsement \mathbf{R}_{to} during precisely controlled and characterized pollutant exposures. Furthermore, our studies were designed to detect possible changes following pollutant exposure at concentrations which resembled those found in urban air.

Materials and Methods

Subject Exposure Procedure and Pulmonary Mechanics Measurement.—Studies were performed using ten healthy men volunteers ranging in age from 25 to 34 years as subjects. All subjects had no previous history of, or physical findings auggesting significant cardiopulmonary disease. Five were eigerette smokers; five were not.

Pulmonary flow resistance (R_L) was measured with an esophageal balloon and a low resistance spirometer using the technique of Mead and Whittenberger. Recordings of flow, volume, and esophageal (intrapleural) pressure were made on a multichancl galvanometric recorder. Airway resistance (R_A) was measured using the body plethysmograph airway-interruption technique of the same authors. Those

racic gas volume (TGV) was determined by a technique modified after Dubois et all!: Apparatus resistance across the R_L, C_L apparatus was 0.38 cm H₂O at 1 liter/sec; across the tubing of the plethysmograph it was 0.51 cm H₂O at 1 liter/sec. Since Widdicombe and Nadel!² had suggested that work of breathing should increase with increasing respiratory frequency (f) and airflow velocities, particularly if total airways dead space (V_D) is increased, we measured compliance (C_L) and R_L during normal resting and forced ventilation at airflows which did not exceed 2.5 liters/sec.

The subjects were nose clips and mouthbreathed, warmed, humidified filtered medicalgrade air from the dilution board schematically described below. Air breathing measurements were made after five minutes, first on the low resistance spirometer and then in the body plethysmograph. Measurements of lung resistance and compliance were complete within one minute following exposure—the plethysmograph data were obtained within the following two minutes. We felt that earlier measurements of these parameters were unnecessary, and did not attempt to make the exposures in the plethysmograph itself. Subject comfort during the hour-long total exposure was a factor in our decision to proceed in this fashion.

After control measurements were made, the subject was then exposed to SO₂ or an SO₂-sodium chloride aerosol mixture. The order of exposure to gas or gas-aerosol mixture was randomized. The exposures lasted 30 minutes each, with measurements being made at 10 minutes and 30 minutes. Sufficient time was allowed between exposures to allow airway and total lung resistance to return to control levels, if any change had occurred. Of 19 studies performed, ten successfully fulfilled the complete criteria of the experimental protocol.

Pollutant Aerosol and Gas Generation and Characterization.—Pollutant mixtures for the

^{* \$02,} parts per million; NaCl, mg/cu m. Values are mean ± 1 SE.

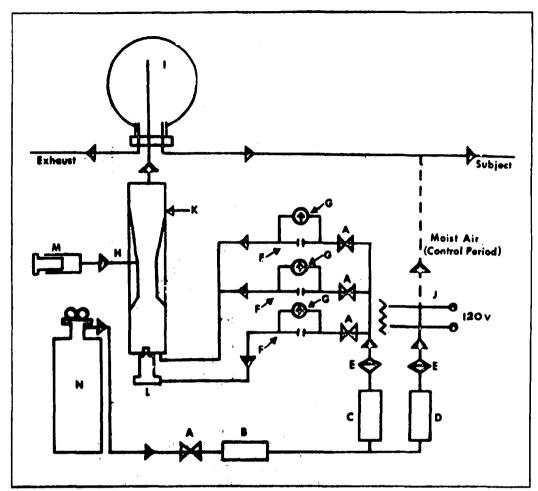
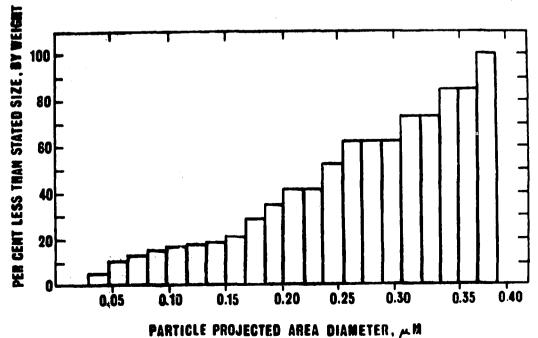


Fig 1.—Schematic of pollutant generation apparatus. A, Valve; B, "catch-ail" air cleaner; C, silica gel; D, watting tube, E, Millipore HA filters; F, critical orifice; G, pressure gage, H, SO_c inlet, I, mixing balloon; J, heating coil with rheostat control, K, Herschel-Type Vent, in Tube, L, Dautreband D_o,1, M, motor driven syringe; N, medical grade compressed air; triangle, flow direction.

Fig 2.—Cumulative particle—size distribution curve for test aerosols (NaC1), sized at $32,000 \times \text{with Zeiss TGZ-3}$ electron microscope.



exposures were produced using a portable aerosol-gas supply apparatus designed and constructed for this study (Fig 1).

Medical-grade compressed air was passed through activated carbon and silica gell, moistened and warmed, then metered by the use of calibrated orifices, before entering the Dautrebande $D_{30}1$ generator or the Venturi tube for mixing with aerosol and gas. Sulfur dioxide gas was supplied to the Venturi throat by a syringe driven by an infusion pump. The Dautrebande $D_{30}1$ aerosol generator was filled with 0.225% by weight solution of NaCl and placed in an opening at the base of the Venturi tube. The salt solution was replaced every 15 minutes to prevent a significant increase in NaCl concentration due to the evaporation of water.

The mixture exited from the Venturi mixing tube into a reservoir balloon, where it was either exhausted, or withdrawn by the subject under test.

All components of the system, with the exception of the balloon were of stainless steel, rigid plastic, or Teflon. The aerosol mixture was delivered to the subject through a 1-inch stainless steel Teflon-lined tube. At the end of the tube was a three-way valve. One port was connected to the "control" air source; the other port was attached to a two-way "J" valve. An exhaust line was connected to the outlet side of the "J" valve. A moulded contour rubber mouthpiece which fitted inside the mouth of the subject was used. Care was taken during the exposures to keep saliva from collecting inside of the "J" valve.

Measurement of pollutant concentration during exposures were made through a stainless steel tap which was injected into the inspiratory side of the breathing valve as close to the mouth as possible.

In order to determine SO₂ concentration, 2.8 liters/min of the pollutant mixture was drawn for two minutes into a midget impinger containing 50 ml of West-Gaeke reagent. The samples were then analyzed spectrophotometrically using the modified West-Gaeke method.¹³

During the first ten-minute exposure, two samples were taken, four minutes apart. During the 20-minute exposure three samples were taken at six-minute intervals. Whenever sodium chloride aerosol was present in the system, sampling was performed by first drawing a sample through HA Millipore filter paper to eliminate sodium chloride interferences. Soveral nonexposure calibration runs indicated that the loss of sulfur dioxide on the filter paper was negligible.

The concentration of sodium chloride aerosol was determined by withdrawing the exposure

mixture at 21 liters/min for 20 minutes through HA Millipore paper, leaching the salt by filter immersion in distilled water, and analyzing by electrical conductivity. A calibration curve was prepared using reagent-grade sodium chloride. Because of the length of time required to sample for the sodium chloride aerosol, this was done at the end of the exposure periods. Several nonexposure checks and postexposure checks found the concentrations to be very consistent over a period of several hours, ie, 2.2 ± 0.08 mg/cu m (mean ± SD).

The particle size distribution of the aerosol was determined by first sampling with an oscillating thermal precipitator onto a carbon-coated glass coverslip. The carbon film was transferred to a 200-mesh electron microscope grid prior to obtaining photographs with an electron microscope. The particles were sized using a particle sizing unit. The particles were all smaller than 0.40μ by weight, the mean size and standard deviation were 0.25μ and $\pm 0.001\mu$, respectively (Fig 2).

Table 1 is a summary of pollutant concentrations to which subjects were exposed in this study. Variations are due to operating characteristics of the generation apparatus. As experience with the unit increased, outward leaks, and other problems were eliminated and reproducibility of concentrations improved. These mixtures of aerosol and gaseous pollutant were generated to specifically simulate the urban milieu. The concentrations of SO₂ are slightly higher than those ever recorded in an acute air pollution episode (London, 1952). During that catastrophe, the concentration of particulate matter was 0.9 mg/cu m.

Results

The data (Table 2) shown represent the mean of six determinations of R_{λ} and TGV in the body plethysmograph. Other parameters of pulmonary mechanics are derived. Pulmonary flow resistance data is presented as an average of inspiratory and expiratory flow resistances based on six to ten breaths.

When compared with individual or mean group controls, no significant increases in C_L and R_L were seen during quiet breathing or during hyperventilation, either after SO_2 alone, or after the SO_2 -aerosol mixture. Similarly, no significant changes in R_A , airway conductance (G_A) , or specific conductance could be demonstrated. Thoracia gas volume did not change significantly. These studies confirm those of Frank et al.¹⁴ Measure of the second state of the second sta

Table 2.—Effect of Exposures on Various Pulmonary Mechanics Measurements*

····	Control	p 10 ft SO2	p 30 ft SO ₂	Control	p 10 ft Mix	p 30 ft Mix	Order
ndividual data							
Subject 6							
RL	0.41, 1.23	0.91, 1.40	0.90, 1.23	1.63, 1.80	1.56, 1.90	1.30, 1.87	
Cr C	0.20, 0.21	0.28, 0.22	0.27, 0.25	0.27, 0.20	0.18, 0.18	0.22, 0.18	
RA Ga	0.70 — 1.42 —	0.97 — 1.03 —	0.95 1.05	1.00 — 1.00 —	0.96	0.95 —	Gas, Mi
TGV	1.42 — 3.36 —	3.37	3.50 -	3.40	1.04 — 3.24 —	1.05 3. 28	
. GA/TGV	0.42 —	0:31	0.30 -	0.29 —	0.32 —	0.32	
Subject 7				<u> </u>	0.52	0.52	
RL	2.02, 1.69	1.90, 1.80	1.33, 1.66	1.50 —	2.00, 1.90	1.80, 1.60	
C.	0.25, 0.21	0.29, 0.26	0.27, 0.24	0.24 —	0.28, 0.30	0.35, 0.23	
RA .	1.12 -	1.18 —	1.03 —	1.10 —	0.98 —	1.08 -	
GA	0.89	0.85 -	0.97 —	0.91 -	1.02 -	0.93	Gas, Mi
TGV	5.20 -	4.94 —	5.12 —	.5.10 —	5.27 -	5.05 -	•
GA/TGV	0.17 —	0.17 —	0.19 -	0.18 —	0.19 —	0.18 —	
Subject 8							
RL	1.70, 2.30	<u> </u>	1.05, 2.50	2.02 -	1.90, 2.30	2.00, 2.40	
CL	0.21, 0.19	<u> </u>	0.18, 0.13	0.20 -	0.16, 0.18	0.16, 0.15	
RA	1.04 —	1.11 —	1.25 —	1.02 —	1.23 —	1.13 —	
G _A	0.96 —	0.90	0.80	0.98 —	0.81 —	0.88 —	Gas, Mi
TGV	2.88 — 0.33 —	2.94 — 0.31 —	2.94 —	2.99 — 0.33 —	2.83 —	2:97 -	
G _A /TGV	0.33 —	0.31 —	0.27 —	0.33 —	0.29	0.30 —	
Subject 10	1 84 3 00	2.03, 2.90	1 70 2 00	1 90 200	170 250	1 60 2 10	
R _L .	1.84, 2.00 0.24, 0.21	0.31, 0.23	1.70, 2.00 0.31, 0.30	1.80, 2.00 0.32, 0.22	1.70, 2.50 0.29, 0.22	1.50, 2.10	
C _{I.} Ra	1.25 -	1.40 —	1.09 —	1.15 -	1.08 -	0.26, 0.14 1.12	Gos, Mi
G _A	0.80 -	0.71 —	0.92	0.87 —	0.93 -	0.89 -	G05, W1
TGV	5.28 —	5.08 —	5.38 -	4.87 —	5.25 —	5.16 —	
GA/TGV	0.15~ —	0.14 —	0.17 —	0.18 -	0.18 —	0.17 -	
Subject 9							
RL	1.60, 3.30	2.70, 2.80	1.40, 1.70	1.40, 1.80	1.90, 1.80	1.60, 1.50	
CL	0.27, 0.25	0.30, 0.30	0.36, 0.23	0.25, 0.31	0.27, 0.32	0.30, 0.25	
RA	1.04 —	1.05 -	1.04 —	1.03	0.97 —	1.02	Gas, Mi
GA	0.96 —	0.95 —	0.96 —	0.97 —	1.03 —	0.98 —	•
TGV	4.48 —	4.58 —	4.50 —	4.65 ~	4.82 —	4.84	
GA/TGV	0.21 -	0.21 —	0.21 —	0.21 -	0.21 -	0.20 —	
Subject 1			•				
RL	1.04, 1.25	1.20, 1.10	1.50, 1.70	1.20, 1.60	1.20, 1.70	0.96, 1.80	
CL	0.39, 0.63	0.36, 0.63		0.28, 0.30	0.30 —	0.23, 0.25	
.RA	0.69 —	0.74 —	0.64 -	0.66	0.85	0.70 —	.Gas, Mi
GA TOM	1.45 —	1.35	1.56	1.52 -	1.18 — 4.16 —	1.43	
TGV.	4.40 —	4.20 — 0.32 —	4:35 0:36	3.97 — 0.38 —		3.90 —	
G _A /TGV	0.33	0.32 -	0.36		0.28 —	0.37 -	
Subject 2	. 00	150 150	1 40 1 50	2.00 2.40	2 10 1 40	1.60 1.60	
RL C	1.80 — -0.25 —	1:60, 1.50 0.29, 0.28	1.40, 1.50	2.60, 2.40 0.36, 0.37	2.10. 1.40	1.50, 1.60 0.29, 0.40	
C _L R _A	·0.25 — 1. 59 —	1.65 -	0.37, 0.42 1.61	1.57	0.30, 0.52 1.62 —	1.55	Mix. Ga
GA	0.63 —	0.61	1.61 0.62	0.64	1.62 - 0.62 -	0.65 —	WIIX, CLO
TGV	5.68 —	5.74 -	5.54 —	6.26 —	5.54	5.64	
GA/TGV	0.11 —	0.11 -	0.11 -	0.10 -	0.11 -	0.12 -	
Subject 3							
	0.99, 2.00	1.40, 1.90	1.50, 1.90	0.83, 1.30	1.80, 2.10	1.30, 2.00	
CL	0.24, 0.29	0.27, 0.25	0.27, 0.26	0.22, 0.22	0.27, 0.23	0.28, 0.27	
RΛ	1.05 -	1.20	1.45	1.32	1.56	1.42 —	Mix, Ga
GA	0.95 —	0.83 -	0.69 —	0.76	0.64 —	0.70 —	•
TGV	3.78 -	3.58	3 .35	3.44	3.18	3.38 —	
GA/TGV	0.25 -	0.23 —	0.21 -	0.22 —	0.20 —	0.21	
Subject 5							
R _L	1.41, 1.55	1.10, 1.71	1.78, 1.71	2.36, 2.37	1.93, 1.62	1.41, 1.55	
CL	0.27, 0.25	0.25, 0.18	0.23, 0.26	0.15, 0.14	0.18, 0.23	0.27, 0.25	
ŔA	1,42 —	1.47 —	1.42	1.42	1.47 —	1.38 —	
GA TOM	0.70 —	0.68	0.70 —	0.70 —	0.68	0.72 —	Mlx, Ga
TGV	4.85 —	4.55	4.61 —	4.73 —	4.62 -	4.85	
GA/TGV	0.14. —	0.15	0.15 —	0.15 —	0.15 —	0.15 —	<u> </u>
Subject 4	10	1 00 5		1.00			
Rt.	1.97, 2.09	1.89, 2.12	1.89, 2.30	1.82, 1.61	1.47, 2.05	2.28, 2.01	
C _L	0.16, 0.15	0.15, 0.13	0.16, 0.15	0.15, 0.14	0.15, 0.13	0.15, 0.13	Miss of
RA	1.97 —	2.05 —	2.06	1.67 —	2.00	1.83 —	Mix, Ga
GA TGV	0.51 · 2.85	0.49 — 2.76 —	0.48 — .2.78 —	0.60 — 3.00 —	0.50 — 2.73 —	0.55 2.89	
	4.00			J.UU —		£.0.	

Table 2.—Effect of Exposures on Various Pulmonary Mechanics Measurements*—(Continued)

	Control	p 10 ft SO2	p 30 ft SO ₂	Control	p 10 ft Mix	p 30 ft Mix	Orde
Grouped data	A						
. Ru	1.48, 1.93	1.64, 1.92	1.44, 1.82	1.72, 1.86	1.76, 1.93	1.57, 1.84	
C _L	0.25, 0.27	0.28, 0.28	0.27, 0.25	0.24, 0.24	0.24, 0.26	0.25, 0.23	
R_{Λ}	1.19 —	1.28 -	1.25 -	1.19	1.27	1.22 -	
⊁.G∧	0.84 —	0.88 —	0.88 —	0.90	0.85	0.88	
, TGV	4.27 —	4.18 —	4.21 —	4.24	4.16	4.20	
GA/TGV	0.23 —	0.21 -	0.21 -	0.22 —	0.21	0.22	

^{*} RL, cm H2O/liter/sec; CL, liter per centimeter H2O; RA, cm H2O/liter/sec; GA, 1/RA; 1GV, liters; and GA/TGV (specific conductance), sec 1 cm H2O-1.

surements of R₁ and C₁, during rapid breathing were also unaffected by any exposure.

Figures 3 to 5 illustrate the absence of significant change in group mean values of R_L , C_L , or G_Λ per TGV following any exposure condition. The R_L and C_L data seemed to add little to the results, for they follow the same trends as the more simply obtained body plethysmograph data.

The wide scatter of individual values and the lack of significant trend can be seen in Table 2 and 3, and Fig 6 to 8. One or two possible "hyperreactors" can be identified here. Control val-

ues are all within reported normal ranges for these measurements.

Except for subject 10, who complained of some dryness of the throat, there were no subjective symptoms associated with any exposure.

Comment

This study confirms existing evidence^{0,14,15} that human exposures to low concentrations (< 3.0 ppm) of SO_n in air do not result in immediate physiologic effects on measures of pulmonary mechanics. Wide subject variability, and hour-to-hour variation in airway resistance and conductance¹⁶ made detection and interpretation of small transient changes difficult. Furthermore, time-series analysis studies in New York¹⁷ and Tennessee¹⁸ have demonstrated a 1 to 2 day lag between peak ambient levels of SO. and development of cough or worsening of

Table 3.—Effects of Exposures on Lung Resistance and Specific Airway Conductance

			% Change	vs Co	ntrol		
Sub-			_	Sub-			
ject		Rt*	G∧/TGV+	ject		Ru	GA/TGV
	p 10′ SO₂	+122.2	26.2		_p 10′ SO₂	+ 15.3	3.0
	<u>p</u> 30′ SO₂	+119.5	-28.6		β 30′ SO ₂	+ 44.3	+ 90
6	<u>₱</u> 10′ Mix	– 4.7	+10.3	1	<u>μ</u> 10′ Μίχ	0.0	-26.3
	p 30' Mix	- 20.2	+10.3		ੂੰ 30′ Mix	20.0	2.7_
_	p 10' SO2	- 6.0	0.0		ji 10′ SO₂	- 11.1	0.0
	p 30′ SO₂	- 34.2	+11.7		p 30′ SO ₂	- 22.2	0.
7	p 10' Mix	+ 33.3	+ 9.5	2	p 10' Mix	- 19.2	4-10.0
	p 30' Mix	+ 20.0	0.0		7 30' Mix	- 42.3	+20.0
	p 10' SO2		- 6.0		p 10' SO:	+ 41.4	- 8.0
	p 30, 205	— 38.2	- 8.2		₽ 30′ SO ₂	4- 51.5	16.0
8	p 10' Mix	 5.9	-12.1	3	p 10' Mix	+116.9	9.1
	p 30' Mix	- 1.0	- 9.0		ji 30' Mix	+.56.6	4.5
	p 10' SO2	+ 10.3	- 6.7		p 10' 80 ·	- ::::0	
	p 30′ SO₂	- 7.6	+13.3		₽ 30° 80°	4 26.2	4. 7.1
10	p 10' Mix	- 5.6	0.0	5	p 10' Mix	- 18.2	OO
	p 30' Mlx	- 16.7	- 5.5		ji ∃0′ M⇔	40.3	0.0
	p 10' SO2	+ 68.8	0.0		i 10′ SO₂	4.1	0,0
	p 30' SO₂	- 12.5	0.0		ர் 30′ ≒⊖∞	- 4.1	fs.fr
9	p 10' Mix	+ 35.7	0.0	4	ji IO′Mi≀	$\sim 19.2^{\circ}$	- 10 0
	₱ 30' Mix	+ 14.3	- 4.8		<u> 7</u> 30′ Mi≀	+ 25.3	ht

^{*} RL, cm H2O/liter/sec; GA/TGV, sec 11 cm H2O 1.

asthma. Spicer¹⁶ has confirmed this relationship in Baltimore, using a sophisticated statistical analysis of changing SO₂ concentrations and measurements of pulmonary airway conductance and resistance as a function of time (a so-called power spectrum analysis). Such work suggests that the expected effects of low concentration SO₂ exposures are delayed unless pulmonary defense mechanisms are in some other way altered.

Inhaled aerosols may yet be shown to act in this fashion, in some way altering the immediate adaptive mechanisms of the airways and lung, rendering them more susceptible to otherwise ineffective concentrations of gaseous pollutant. Stokinger¹⁹ and Anderson²⁰ have recently discussed the potential of gas-aerosol interaction, and the limitations of testing adequately for its presence or absence. It is unclear whether or not

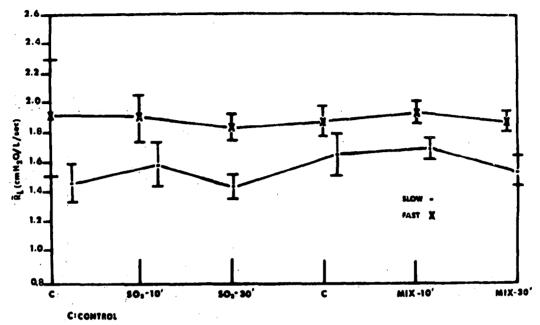
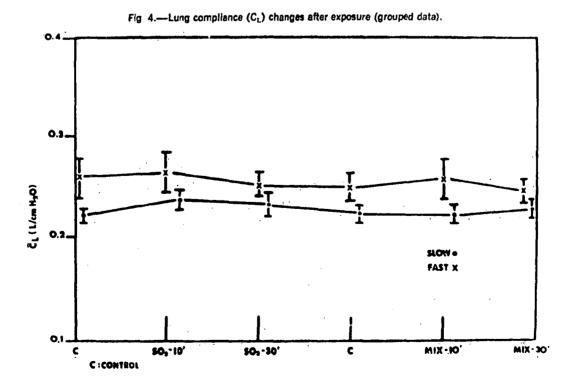


Fig 3:-Total lung resistance (R_I) changes after exposure (grouped data).



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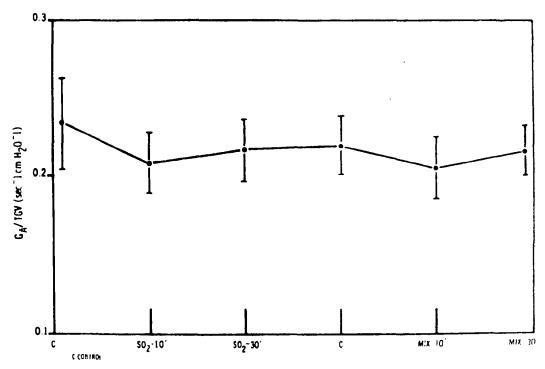


Fig 5.--Specific airway conductance changes after exposure (grouped data).

the aerosols in themselves need to be "irritant" to produce an effect in man, though they need not be in animals. Toyama's industrial dusts may have been more irritant than the NaCl acrosol of Frank et al^a and our own, and this may account in part for the evocation of response from his SO₂-dust exposed subjects.

We were able to identify at least one possible "hyperreactor" to SO₂ in the present study (subject 3, a 24-year-old nonsmoker). Our work suggests, as have other studies, 14,15 that there may be only one or two physiologic "reactors" for every ten exposed subjects. The implication for future exposure studies is that large numbers of "normal" subjects will need to be studied to locate persons who show effects of inhaled pollutants on pulmonary mechanics.

A study which probed immediate gasacrosol synergism in patients already affected by pulmonary disease might report positive results where ours have been negative. Support for this exists in the literature,²¹ though complete acrosol characterization data are not given. Medicolegal and ethical considerations make studies of this kind difficult.

Conclusions

In summary, like Frank et ale and the work recently reported by Snell and Luchsinger,22 we could not demonstrate gas-aero sol synergism for SO₂ and inert aerosols at concentrations which approximate those in urban atmospheres. These experiments suggest that grouped population data may not be as sensitive an indicator of effect in the experimental exposure situation as they are in the epidemiological setting. These findings in humans are in marked contrast to the study in animals of Amdur et al. 45 where grouped data, as well as single responses. gave evidence of a synergistic effect when guinea pigs were exposed to mixtures of aerosol and gas similar to those reported here. (She used the same indicator of response, namely pulmonary flow resistance. in her studies as we did.)

While gas-acrosol synergism may yet be proven an important toxicologic mechanism in man, we suspect that reactor, characteristics of reactant, and timing and sensitivity of measurement will have to be more carefully considered in future studies if such an effect is to be demonstrated.

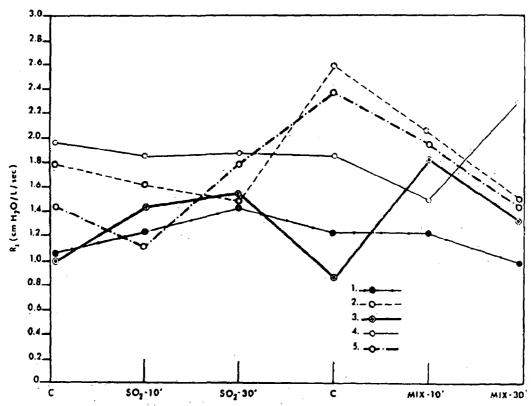
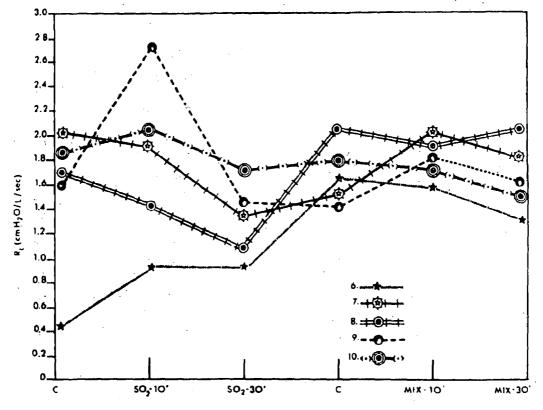


Fig 6:-Plot of lung resistance changes in individual subjects after exposure to SO2 gas and NaC1 acrosol.



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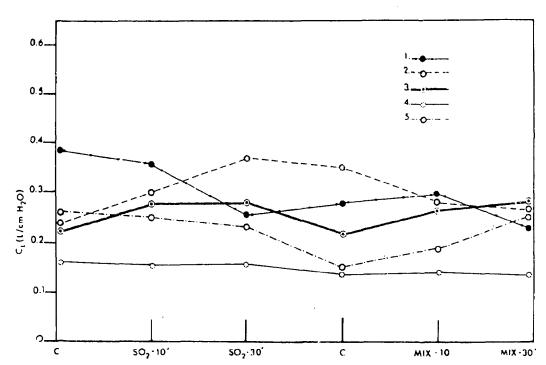
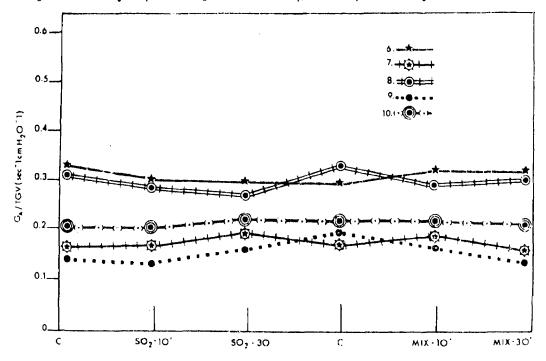


Fig 7.—Plot of leng compliance changes in individual subjects after exposure to SO, gas and NaC1 aerosol.



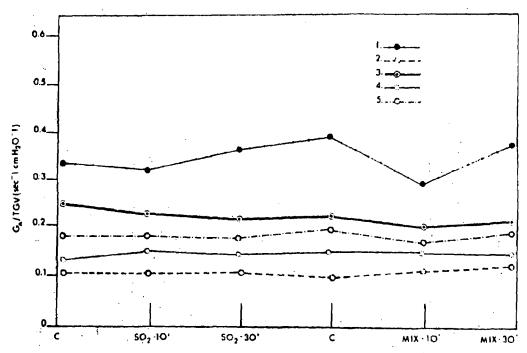
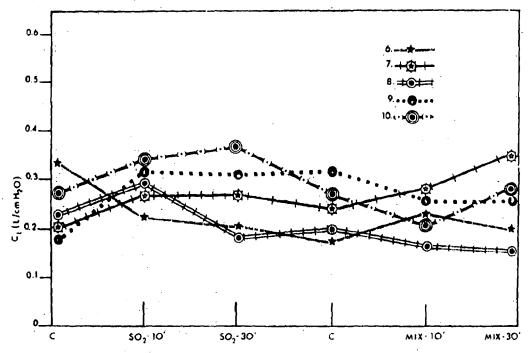


Fig 8.—Plot of specific airway conductance in individual subjects after exposure to SO₂ gas and NaC1 aerosol.



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Resting pulmonary mechanics studies may not represent the best approach to problems of acute-effect air pollution toxicology in man. Studies of distribution of ventilation or changes in pulmonary me-

chanics following exposure during exercise may possibly be more sensitive indicators of response.

This study was supported by Public Health Service grant PH86-67-73 from the National Center for Air Pollution.

References

- 1. Com, M., and Burton, G.: The Irritant Potential of Pollutants in the Atmosphere, Arch Environ Health 14:54-61 (Jan) 1967.
- 2. LaBelle, C.W.; Long, J.E.; and Christofano, E.E.: Synergistic Effects of Aerosols, Arch Industr Health 11:297-304 (April) 1955.
- 3. Goetz, A.: On the Nature of the Synergistic Action of Aerosols, Int J Air Water Pollut 4:168-184 (Sept) 1961.
- 4. Amdur, M.O.: The Influence of Acrosols Upon the Respiratory Response of Guinea Pigs to Sulphur Dioxide, Industr Hyg Assoc Quart 18:149-155 (June) 1957.
- 5. Amdur, M.O.. The Physiological Response of Guinea Pigs to Atmospheric Pollutants. Int J Air Pollut 1:170-183 (March) 1959.
- 6. Frank, N.R.; Amdur, M.O.; and Whittenberger, J.L.: A Comparison of the Acute Effects of SO₈ Administered Alone or in Combination With N.C.1 Particles on the Respiratory Mechanics of Healthy Adults, Air Water Pollut 8:125-133 (Feb.) 1964.
- 7. Toyama, T.: Studies on Aerosols: I. Synergistic Response of Pulmonary Airway Resistance on Inhaling Sodium Chloride and SO₂ in Man, Jap J Industr Med 4:18-27, 1962.
- 8. Toyama, T.: Air Pollution and Its Health Effects in Japan, Arch Environ Health 8:153-173 (Jan) 1964.
- 9. Mead, J., and Whittenberger, J.L.: Physical Properties of Human Lungs Measured During Spontaneous Respiration, J Appl Physiol 5:779-796 (June) 1953.
- 10. Mead, J., and Whittenberger, J.L.: Evaluation of Airway Interruption Technique as a Method for Measuring Pulmonary Airflow Resistance, J Appl Physiol 6:408-416 (Jan) 1954.
- 11. Dubois, A.B., et al: A Rapid Plethysomographic Method for Measuring Thoracic Gas Volume: A Comparison With a Nitrogen Washout Method for Measuring Functional Residual Capacity in Normal Subjects, J Clin Invest 35:322 (March) 1956.

- 12. Widdicombe, J.G., and Nadel, J.A.: Airway Volume, Airway Resistance, and Work and Force of Breathing: Theory, J. Appl. Physiol. 18:863-868 (Sept.) 1963.
- 13. West, P.W., and Gaeke, G.C.: Fixation of Sulfur Dioxide as Disulfitomercurate (II) and Subsequent Colorimetric Estimation, Anal Chem 12:1816-1819 (Dec.) 1956.
- 14. Frank, N.R., et al: Effects of Acute Controlled Exposure to SO₂ on Respiratory Mechanics in Healthy Male Adults, *J Appl Physiol* 17:252-258 (March) 1962.
- 15. Lawther, P.J.: Effects of Inhalation of Sulphur Dioxide on Respiration and Pulse Rate in Normal Subjects, Lancet 2:745-748 (Oct 8) 1955.
- 16. Spicer, W.S.: Air Pollution and Meteorologic Factors, Arch Environ Health 14:185-188 (Jun) 1967.
- 17. McCarroll, J., et al: Healthy and the Urban Environment: V. Air Pollution and Illness in a Normal Urban Population, Arch Environ Health 14:178-183 (Jan) 1967.
- 18. Zeidberg, L.D.; Prindle, R.A.; and Landau, E.: The Nashville Air Pollution Study: I. Sulphur Dioxide and Bronchial Asthma, A Preliminary Report, Amer Rev Resp Dis 84:489-503 (Oct.) 1961.
- 19. Stokinger, H.: Toxicological Interactions of Mixtures of Air Pollutants, Int J Air Pollut 2:313-326 (June) 1960.
- 20. Anderson, D.O.: The Effects of Air Contamination on Health: II. A Review, Canad Med Assoc J 97:585-593 (Sept) 1967.
- 21. Lovejoy, F.W., Jr., et al: Measurement of Gas Trapped in the Lungs During Acute Changes in Airway Resistance in Normal Subjects and in Patients With Chronic Pulmonary Disease, Amer J Med 30:884-892 (June) 1961.
- 22. Snell, R.E., and Luchsinger, P.C.: Effects of Sulfur Dioxide on Expiratory Flow Rates and Tabal Respiratory Resistance in Normal Human Subjects, Arch Environ Health 18:690-698 (April) 1969.

TABLE _5_ Cat No. 1676 RESPONSE TO VARIOUS STIMULI EXPRESSED AS PERCENT CHANGE RELATIVE TO CONTROL VALUES.

	Pu	lmonary Flow Resistance	(b)	
Mech. % Control Stim. Change	Low 2 Control SO ₂ Change	Mech. % Control Stim. Change	SO2 & % Control NaCl Change	Mech. % Control Stim. Change
9.5 68.2 617.9 °.5	8.4 9.5 0 9.5 10.6 11.6 9.5 12.9 35.8	9.5 59.7 528.4	14.0 15.2 60 10.6 16.3 71.6 9.5 15.2 60	10.6 72.8 586.8
Mean [±] 9.5 69.2 S.O. 0	9.1 11.0 0.6 1.7	9.5 59.7	11.4 15.6 2.3 0.6	10.6 72.8
REFLEX INTACT	£ N.S.	REFLEX INTACT	Δ N.S.	REFLEX INTACT
		Lung Compliance (c)		
Mech. % Control Stim. Change	Low % Control SO ₂ Change	Mech. % Control Stim. Change	SO ₂ & % Control NaCl Change	Mech. Control Stim. Change
14.5 3.8 -73.8 14.5	15.0 12.3 -13.7 15.0 12.4 -16.4 14.5 12.3 -13.0	13.2 3.4 -74.2	13.6 11.4 -13.6 13.2 11.4 -13.6 12.8 11.4 -13.6	12.1 8.3 -31.4
Mean [±] 14.5 · 3.? S.D. 0	14.9 12.7 0.3 0.2	13.2 3.4	13.2 11.4 0.4 0	12.1 8.3
	P<0.01		P<0.05	
Sequence of (a) (1)	(2)	(3)	(4)	(5)

*Control for .15 minutes preceding challenge.

(a) See Figure.
(b) Pulmonary Flow Resistance, Cm H₂0/1/sec.
(c) Lung Compliance, m1/cm H₂0.

TABLE 5 (Continued) Cat No. 1676

					Pu	lmonary	Flow R	esistance	(b)			
Control	NaC1	% Change	Control	Mech. Stim.	% Change	Control	High SO ₂	% Change	Control	Mech. Stim.	% Change	
9.5 9.5	12.9 14.0 12.9	35.3 47.4 35.8	9.5	98.6	937.9	9.5 9.5 9.5	10.6 12.9 12.9	11.6 35.08 35.8	9.5	96.3	808.4	
Mean± 9.5 S.D. 0	13.3		9.5	98.6		9.5 0	12.1		9.5	86.3		
	P<0.0	1	RE	FLEX IN	TACT		ΔN.S.		RI	EFLEX 1	NTACT	
						Lung C	omplia	nce (c)				
Control	NaC1	% Chan ge	Control	Mech. Stim.	?∤ Change	Contro	High 01 SO ₂	% Change	Contro	Mecb. 1 Stim.	. ¾ . Change	
12.4 12.1	11.4 11.1 11.4	-6.9 -9.4 -6.9	11.7	4.1	-65	11.4 11.4 11.4	16.2 16.2 16.2	42.1	16.2	4.7	-71	
Mean 12.3 S.D. 0.2	11.3		11.7	4.1		11.4	16.2		16.2	4.7		
	P< 0.	05					P 0.0	01				
Sequence of Challenge	^f a)	(6)	- 	(7)			(8)			(9)		

*Control for 15 minutes preceding challenge.

(a) See Figure.

(b) Pulmonary Flow Resistance, Cm H₂0/1/sec.(c) Lung Compliance, ml/cm H₂0.

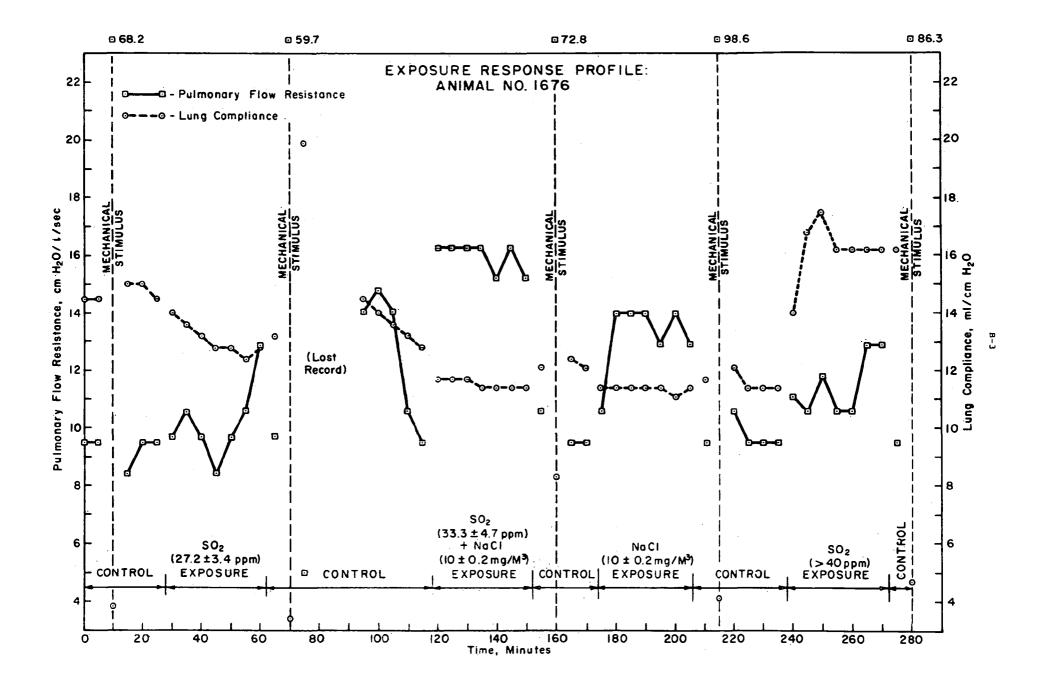


TABLE 6 Cat No. 1651 RESPONSE TO VARIOUS STIMULI EXPRESSED AS PERCENT CHANGE RELATIVE TO CONTROL VALUES*

	Pu	Imonary Flow Resistance	(b)	
Mech. % Control Stim. Change	Low ½ Control SO ₂ Change	Mech. % Control Stim. Change	SO ₂ +	Mech. % Control Stim. Change
14.5 54.4 275.2 14.5	14.5 11.8 -18.6 14.5 13.0 -10.3 14.5 11.8 -18.6	14.5 163.9 1030	14.5 13.0 -10.3 14.5 14.5 0 14.5 13.0 -10.3	10.5 84.1 572.8 14.5
1ean± 14.5 54.4 5.0. 0	14.5 12.2 0 0.7	14.5 163.9	14.5 13.5 0 0.9	12.5 84.1 2.8
REFLEX INTACT	P <0. 05	REFLEX INTACT	Δ N.S.	REFLEX INTACT
	_	Lung Compliance (c)		
Mech. % Control Stim. Change	Low % Control SO ₂ Change	Mech. ½ Control Stim. Change	SO ₂ + ½ Control NaCl Change	Mech. % Control Stim. Change
11.1 7.2 -35.1 11.1	10.5 14.1 30.6 10.8 13.6 25.9 11.1 13.6 25.9	14.1 5.3 -62.4	12.4 15.7 28.7 12.1 14.1 15.6 12.1 15.1 23.8	16.3 5.2 -67.5 15.7
"ean [±] 11.1 7.2 S.D. 3	10.8 13.8 0.3	14.1 5.3	12.2 15.0 0.2 0.8	16.0 5.2 0.4
	P<0.01		P<0.05	
Sequence of (1)	(2)	(3)	(4)	(5)

*Control for 15 minutes preceding challenge.

(a) See Figure.
(b) Pulmonary Flow Resistance, Cm H₂0/1/sec.
(c) Lung Compliance, m1/cm H₂0.

TABLE 6 (Continued) Cat No. 1651

	Pu	Imonary Flow Resistance	(b)	
<pre> Control NaCl Change </pre>	Mech. ⅓ Control Stim. Change	High % Control SO ₂ Change	Mech. % Control Stim. Change	
13.1 11.7 -19.3 14.5 11.7 -19.3 11.8 -18.6	11,8 151.1 1267.4 10.3	14.5 19.3 35.1 14.5 22.9 57.3 19.3 33.1	9.3 119 1179.6	
Mean [±] 13.8 11.7 S.D. 1.0 0.1	11.1 151.1 1.1	14.5 20.5 0 2.1	9.3 119	
N.S.	REFLEX INTACT	P< 0.05	REFLEX INTACT	
		Lung Compliance (c)		
% Control NaCl Change	Control Stim. Change	Control SO ₂ Change	Mech. % Contro ¹ Stim. Change	·
14.1 15.1 5.2 14.5 15.1 5.2 15.1 5.2	15.1 3.º -74.4 14.6	13.6 15.7 13.4 14.1 15.7 13.4 15.7 13.4	15.7 4.9 -68.8	
Mean± 14.4 15.1 S.D. 0.4 0	14.9 3.9 0.4	13.9 15.7 0.4 0	15.7 4.9	
N.S.		∆ %. S.		
Sequence of (6)	(7)	(8)	(9)	

*Control for 15 minutes preceding challenge.

(a) See Figure.

(b) Pulmonary Flow Resistance, Cm H₂0/1/sec.
 (c) Lung Compliance, ml/cm H₂0.

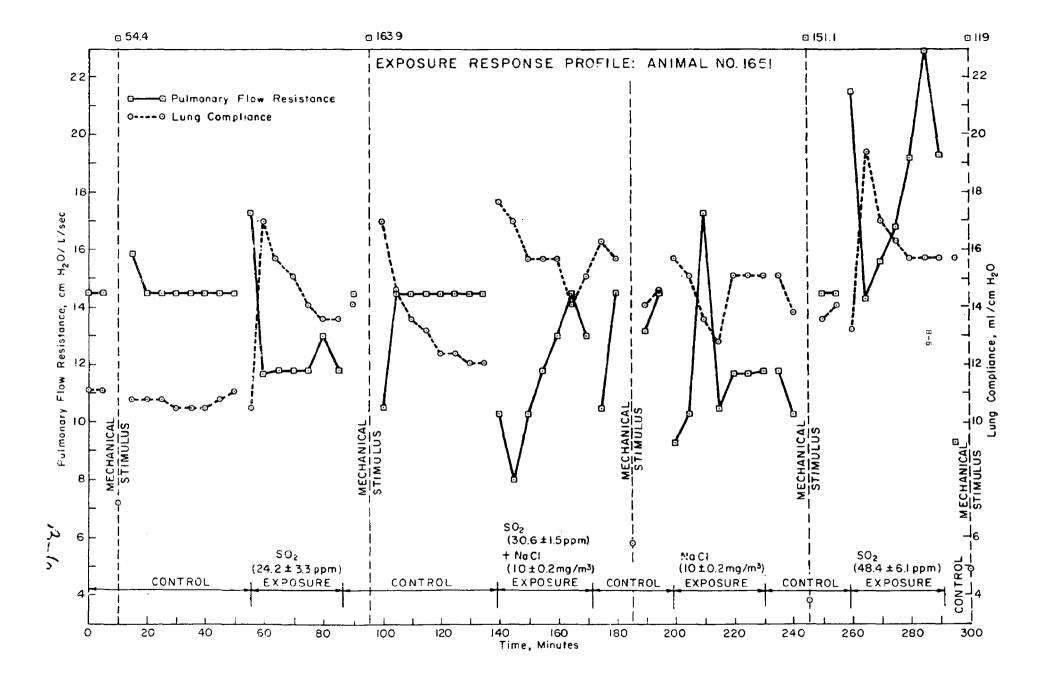


TABLE 7 Cat No. 1612 RESPONSE TO VARIOUS STIMULI EXPRESSED AS PERCENT CHANGE RELATIVE TO CONTROL VALUES*

	Pu1	lmonary Flow Resistance	(6)	
Mech. % Control Stim. Change	Low % Control SO ₂ Change	Mech. % Control Stim. Change	Control NaCl Change	Mech: % Control Stim. Change
20.9 156 646.4 20.9	20.9 27.1 29.7 20.9 28.1 34.4 20.9 30.2 44.5	28.1 103 329.2 23 20.9	20.9 29.2 39.7 20.9 25.6 22.5 27.1 29.7	20.9 119 469.4
Mean‡ 20.9 156 5.D. 0	20.9 29.5 0 1.6	24.0 103 3.7	20.9 27.3 0 1.8	20.9 119
REFLEX INTACT	+0.05	REFLEX INTACT	P<0.05	. REFLEX INTACT
		Lung Compliance (c)		
Mech. % Control Stim. Change	Low ½ Control SO ₂ Change	Mech. ½ Control Stim.Change	SO ₂ + % Control NaCl Change	Mech. ¾ Control Stim. Change
6.3 5.0 -21.3 6.4	7.0	6.9 6.9 0 7.0 6.9	7.4 7.1 -1.4 7.0 8.4 16.7 7.1 -1.4	7.7 7.7 0
Mean± 6.3 S.D. 0.1	7. 6.5 0.1 5.1	6.9 6.9 0.1	7.2 7.5 0.3 0.6	7.7 7.7
•	P<0.01		Δ N.S.	
Sequence of (1)	(2)	(3)	(4)	(5)

^{*}Control for 15 minutes preceding challenge.

(a) See Figure.
(b) Pulmonary Flow Resistance, Cm H₂0/1/sec.
(c) Lung Compliance, ml/cm H₂0.

TABLE 7 (Continued)

Cat ilo. 1612

	Pu	Imonary Flow Resistance	(6)	
₹ Control NaCl Change	Mech. ½ Control Stim. Change	High % Control SO ₂ Change	Mech. % Control Stim. Chang≘	
20.9 38.7 85.2 20.9 30 43.5 33.9 62.2	20.9 81.6 290.4	20.9 28.1 34.4 20.9 27.1 29.7 23 32.2 54.1	20.9 134 54.1	
lean± 20.9 34.2	20.9 81.6	21.6 29.1 1.2 2.7	20.9 134	
P< 0.05	REFLEX INTACT	P< 0.05	REFLEX INTACT	į
		Lung Compliance (c)		
% Control NaCl Change	Mech. ₹ Control Stim. Change	High ½ Control SO2 Change	Mech. % Control Stim. Change	
7.7 9.3 9.2 7.5 9.4 10.5 8.1 6.6	8.8 6.7 -23.9	7.7 6.4 -13.1 6.5 8.4 14.0 7.9 6.1 -17.2	7.1 5.0 -29.6	
P<0.01	R. R. F. 7	7.4 7.0 0.8 1.3 Δ N.S.	7.1 5.0	
Sequence of (6)	(7)	(9)	(9)	

^{*}Control for 15 minutes preceding challenge.

(a) See Figure.

(b) Pulmonary Flow Resistance, Cm H₂0/1/sec.

(c) Lung Compilance, m1/cm H₂0.

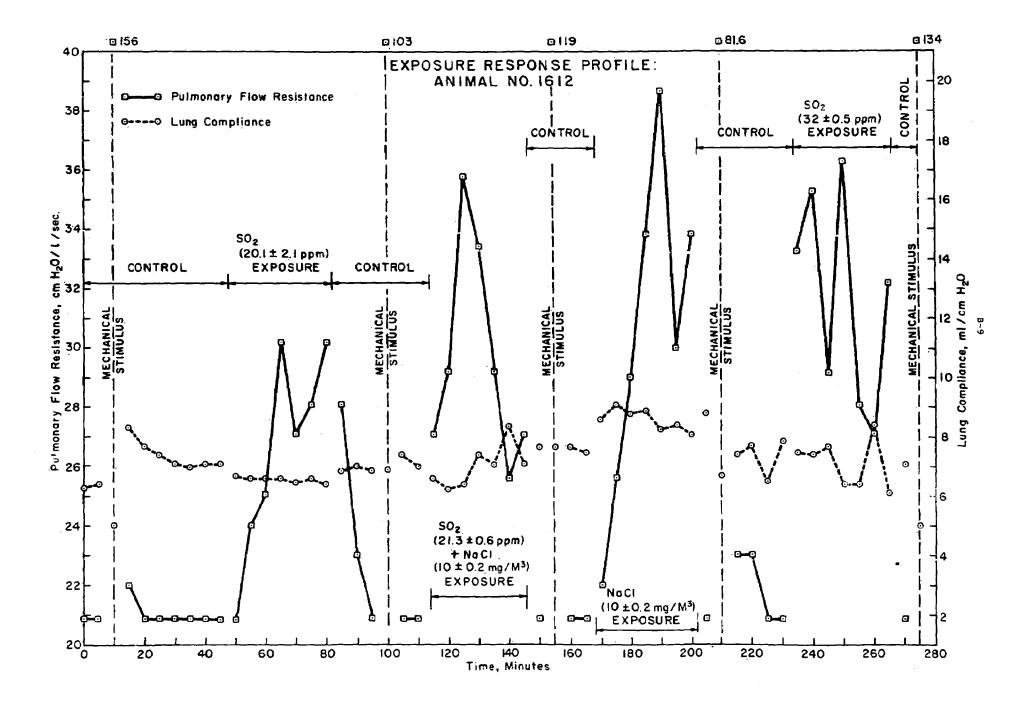


TABLE 8 Cat No. 1633 RESPONSE TO VARIOUS STIMULI EXPRESSED AS PERCENT CHANGE RELATIVE TO CONTROL VALUES*

					Pu	lmonary F	low Re	sistance	(P)					
Control	Mech. I Stim.	% Change	Control	Low SO ₂	½ Change	Control	Mech. Stim.	% Change	Contro	S0 ₂ + NaC1	% Change	Control	Mech. Stim.	% Change
16.9 17.9	122	601	15.9 15.9 15.9	14.0 15.6 14.2	-11.9 -1.9 -10.7	16,8 15.9	80.1	389.9	15.9 15.9 15.9	19.1 17.9 20.8	20.1 12.6 30.8	17.9 19.5	82	338.5
Mean 17.4 S.D. 0.7	122		15.9 0	14.6 0.9		16.4 0.6	80.1		15.9 0	19.3		18.7 1.1	82	
RE	FLEX INT	ГАСТ		N.S.		RE	FLEX	INTACT		ΔN.S.		RE	FLEX IN	TACT
						Lung Co	mplian	ce (c)						
Control	Mech. Stin.	% Change	Control	Low SO ₂	:/ Change	Control	Mech. Stim.	% Change	Contro!	S0 ₂ + NaC1	% Change	Control	Mech. Stim.	% Change
3.0 7.7	3.6	-54.1	7.7 7.7 7.7	7.5 7.5 7.7	-2.6 -2.6 0	8.1 7.5	8.1	3.8	7.7 7.5 7.5	7.0 7.1 7.3	-7.5 -6.2 -3.5	7.7 7.3	4.2	-44.0
Mean± 7.9 S.D. 0.2	3.6		7.7 0	7.6 0.1		7.8 0.4	8.1	ı	7.6 0.1	7.1 0.2		7.5 0.3	4.2	
				N.S.					P<	-0.05				
Sequence q Challenge	(a)	(1)	<u></u>	(2)			(3)			(4)			(5)	

^{*}Control for 15 minutes preceding challenge.

⁽a) See Figure.

⁽b) Pulmonary Flow Resistance, Cm H₂0/1/sec.(c) Lung Compliance, ml/cm H₂0.

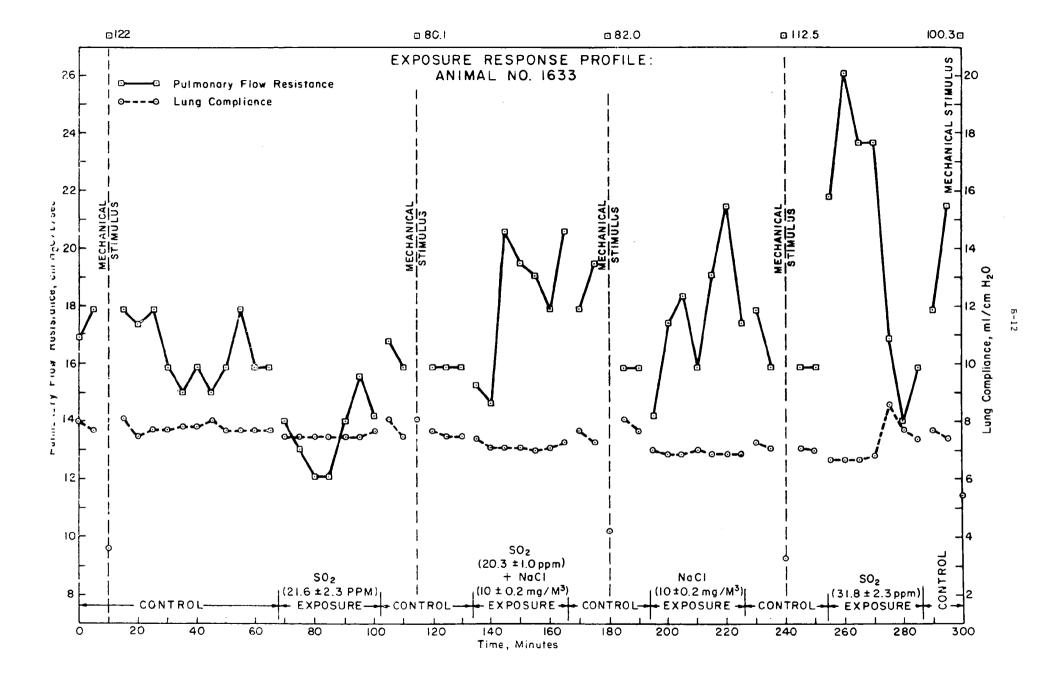
Cat No. 1633 RESPONSE TO VARIOUS STIMULI EXPRESSED AS PERCENT CHANGE RELATIVE TO CONTROL VALUES*

TABLE > 8 (Continued)

				•		Pu	imonary Flow	Res	sistance	(Б)			
Co	ontro1	NaC1 ,	% Change	Control	Mech. Stim.	% Change	H Control S	igh 02		Control	Mech. Stim.	% Change	
	15.9 15.9	19.1 21.5 17.4	20.1 35.2 9.4	17.9 15.9	112.5	565.7		6.9 4.0 5.9	-11.9	17.9 21.5	100.3	409.1	
Meant 1 S.D.	15.9	19.3		16.9 1.4	112.5		15.9 1 0	5.6 1.5		19.7 2.5	100.3		
	•	N.S.		REF	LEX INT	TACT	Δ	N.S.		RE	FLEX I	NTACT	
							Lung Comp	liand	ce (c)				
Co	ontro1	NàC1	% Change	Control	Mech. Stim.	(nange	H Control S		光 Change	Contr o 1	Mech. Stim.	½ Change	
	8.1 7.7	6.9 6.9 6.9	-12.7 -12.7 -12.7	7.3 7.1	3.3	-54.2	7.0 7	3.6 7.7 7.4	22 9.2 5	7.7 7.4	5.4	2.3	
Mean [‡] 7 S.D. (6.9 0	-	7.2 0.1	3.3			7.9).6		7.6 0.2	5.4		
		N.S.					۵ ۸	ı.s.					
Sequei Challe	nce of enge)	(6)		(7)		. ((8)	·		(9)		

*Control for 15 minutes preceding challenge.

⁽a) See Figure.
(b) Pulmonary Flow Resistance, Cm H₂0/1/sec.
(c) Lung Compliance, m1/cm H₂0.



(5)

TABLE 9 Cat No. 1606

	_			Pu	lmonary F	ow Res	istance	(b)	- -				
Mech. Control Stim.	% Change	Control	Low SO ₂	% . Change	-Control	Mech. Stim.	% Change	Contro	50 ₂ + NaC1	% Change	Control	Mech. Stim.	% Change
19.9 127 19.9	538	19.9 18.5 18.5	15.7 15.7 21.9	-17.2 -17.2 15.5	18.5	87.4	372.4	17.1 14.1 18.5	16.6 16.6 16.6	0.2 0.2 0.2	14.1	112	694.3
Mean± 19.9 127 S.D. 0		19.0 0.8	17.9 3.6		18.5	87.4		16.6 2.2	16.6 0		14.1	112	
REFLEX INT	ACT		N.S.		RE	FLEX II	NTACT		∆ N.S.		RE F	LEX INT	ACT
					Lung Con	pliano	:e (c)						
Mech. Control Stim.	% Change	Control	Low Sn ₂	% Change	Control	Mech. Stim.	% Change	Contro	SO ₂ + NaC1	% Change	Control	Mech. Stim.	% Change
9.7 6.9 9.3	-27.4	9.1 9.3 9.5	9.1 9.3 9.5	-2.2 0.0 2.2	9.7	4.7	-51.5	9.9 10.2 8.7	7.6 7.3 7.5	-20.8 -24 -21.9	8.4	4.1	-51.2
Mean± 9.5 6.9 S.D. 0.3		9.3 0.2	9.3 0.2		9.7	4.7		9.6 0.8	7.5 0.2		8.4	4.1	

(3)

*Control for 15 minutes preceding challenge.

Sequence of a) Challenge

(a) See Figure.
(b) Pulmonary Flow Resistance, Cm H₂0/1/sec.
(c) Lung Compliance, m1/cm H₂0.

Δ N.S.

(2)

(1)

A N.S. = Difference between means not sigificant (P>0.05)

P <0.05

(4)

TABLE 9 (Continued) Cat No. 1606

			·	Pu	1monary F	low Re	s istance	(b)					
Control NaCl)/. Change	Control	Mech. Stim.	% Change	Control	High SO _{2.}	% Change	Contro _. l	Mech. Stim.	% Change	-		
14.1 11.7 18.5 11.6 18.5 8.4	-31.3 -31.9 -50.7	19.2	96.3	401.6	19.9 18.5 18.5	22.3 25.3 19.9	17.6 23.4 4.9	18.5	61.1	230.3			
Mean± 17.0 10.6 S.D. 2.5 1.9		19.2	95.3		19.0 0.8	22.5		18.5	61.1				
P<0.05		REF	LEX INT	АСТ	Δ	N.S.		R€	FLEX I	NTA CT	•		
				\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\	Lung Co	mplian	ce (c)			-	·		
Control NaCl	7. Change	Control	Mech. Stim.	% Change	Contro!	High SO ₂	% Change	Control	Mech. Stim.	% Change			
9.3 7.6 8.9 7.6 8.5 7.7	-14.6 -14.6 -13.5	8.5	4.1	-51.8	6.7 7.2 7.2	9.2 7.2 7.1	16.6 2.4 0.9	7.9	8.5	7.6			
Mean [±] 8.9 7.6 S.D. 0.4 0.1		8.5	4.1		7.0 0.3	7.5 0.6		7.9	8.5				
P< 0.05					·	ΔN.S.						 	
Sequence of (a) Challenge	(6)		(7)			(8)			(9)			· ·	

^{*}Control for 15 minutes preceding challenge.

⁽a) See Figure.
(b) Pulmonary Flow Resistance, Cm H₂0/1/sec.
(c) Lung Compliance, m1/cm H₂0.

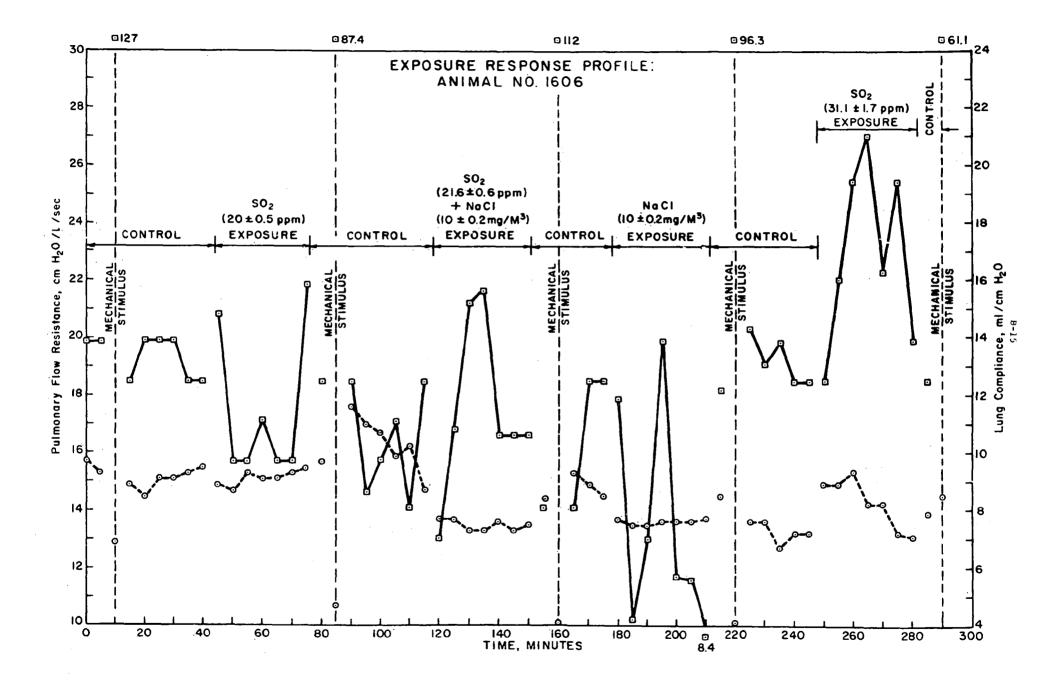


TABLE 10 Cat No. 1112 RESPONSE TO VARIOUS STIMULI EXPRESSED AS PERCENT CHANGE RELATIVE TO CONTROL VALUES*

	Pu	Imonary Flow Resistance	(b)	
Low %	Mech. %	SO ₂ + %	Mech. %	%
Control SO ₂ Change	Control Stim. Change	Control NaCl Change	Control Stim. Change	Control NaCl Change
18.1 27.6 57.1	19.7 65.7 233.5	18.1 25.9 43.1	18.9 49.8 156.2	18.9 14.9 -23.3
18.1 29.2 66.2	19.7	18.1 34.4 90.1	19.7	19.7 20.3 4.5
16.5 27.6 57.1	19.7	18.1 28.8 59.1	19.7	19.7 23.1 18.9
Mean± 17.6 28.1	19.7 65.7	18.1 29.7	19.4 49.8	19.4 19.4
S.D. 0.9 0.9	0	0 4.3	0.5	0.5 4.2
t-test P< 0.(1	REFLEX INTACT	p< 0.05	REFLEX INTACT	Δ N.S.
		Lung Compliance (c)		
Low %	Mech. %	SO ₂ + %	%	%
Control SO ₂ Change	Control Stim. Change	Control NaCl Change	Control Stim. Change	Control- NaCl Change
15.5 12.8 -4.2	11.5 11.5 +0.0	11.3 12.8 13.6	11.5 16.0 47.7	11.5 9.2 -15.1
12.8 13.2 -1.2	11.3	11.0 14.5 28.7	10.5	10.5 10.0 -7.7
11.8 12.8 -4.2	11.8	11.5 12.5 10.9	10.5	10.5 9.2 -15.1
Mean [‡] 13.4 12.9	11.5 11.5	11.3 13.3	10.8 16.0	10.8 9:5
S.D. 1.9 0.2	0.3	0.3 1.1	0.6	0.6 0.5
t-test Δ N.S.		Δ N.S.		-0.05
Sequence of (1) Challenge	(2)	(3)	(4)	(5)

*Control for 15 minutes preceding challenge.

(a) See Figure.

(b) Pulmonary Flow Resistance, Cm H₂0/1/sec.
 (c) Lung Compliance, ml/cm H₂0.

△ N.S. = Difference between means not sigificant (P>0.05)

TABLE 10 (Continued)

Cat No. 1112

RESPONSE TO VARIOUS STIMULI EXPRESSED AS PERCENT CHANGE RELATIVE TO CONTROL VALUES*

	Pulmonary Flow Resistance (b)								
High Control SO ₂	½ Change	Control	Mech. Stim.	꽃 Change					
16.5 37.3	149.1 126.1 172.1	20.3 20.3 20.9	37-3.	82					
Mean± 16.5 41.1 S.D. 0 3.8		20.5 0.3	37.3						
t-test P<0.01		. RE	FLEX IN	TACT					
					Lung Compliance (c)				
High Control SO ₂	¾ Change	Contro1	Stim.	¥ Change					
10.5 15.0	63.8 42.9 77.1	11.5 11.0 10.0	14.0	29.6					
Mean± 10.5 16.9 S.D. 0 1.8	,	10.8 0.8	14.0						
t-test P< 0.05		 							
Sequence of Challenge	(6)		(7)	· · · · · · · · · · · · · · · · · · ·					

*Control for 15 minutes preceding challenge.

(a) See Figure.
(b) Pulmonary Flow Resistance, Cm H₂0/1/sec.
(c) Lung Compliance, ml/cm H₂0.

TABLE 11 Cat No. 1609 RESPONSE TO VARIOUS STIMULI EXPRESSED AS PERCENT CHANGE RELATIVE TO CONTROL VALUES*

	•					Pu	lmonary F1	ow Res	istance	(b)					
·	Control	Mech. Stim.	% Change	Control	Low SO ₂	% Change	Control	Mech. Stim.	% Change	Cóntro	SO ₂ + NaC1	% Change	Control	Mech. Stim.	½ Change
	19.3 20.5 18.8	114	483.6	20. I 18. 1 18. 1	19.3 20.5 20.5	6.6 13.3 13.3	18.1	162	795	18.1 18.1 17.0	18.1 20.5 15.8	0 13.3 -12.7	15.8 13.3	72.5	398.3
Mean: S.D.	19.5	114.		19.0 1.0	20.1 0.7		18.1	162		17.7 0.6	18.1 2.4		14.6 1.8	72.5	
	RE	FLEX IN	TACT		N.S.		RE	FLEX II	NTACT		∆N.S.		REF	LEX INT	ACT
							Lung Con	pliano	e (c)						
	Control	Mech. Stim.	% Change	Control	Low SO ₂	½ Change	Control	Mech. Stim.	% Change	Contro	SO ₂ + NaC1	% Change	Control	Mech. Stim.	½ Change
	8.9 9.1 9.2	5.0	-44.9	9.4 9.2 9.2	8.5 9.4 8.9	-9.6 1.1 -4.3	9.7	4.5	-53.6	9.7 10.1 9.9	9.4 9.4 9.4	-5.1 -5.1 -5.1	10.6 10.3	7.9	-24.4
Mean S.D.	0.2	5.0		9.3 0.1	8.° 0.5		9.7	4.5		9.9 0.2	9.4 0		10.5	7.9	
				1	N.S.						Δ N.S.				
Seq	uence of	a)	(1)		(2)			(3)			(4)			(5)	

^{*}Control for 15 minutes preceding challenge.

⁽a) See Figure.

 ⁽b) Pulmonary Flow Resistance, Cm H₂0/1/sec.
 (c) Lung Compliance, ml/cm H₂0.

TABLE 11 (Continued)

Cat No. 1609

RESPONSE TO VARIOUS STIMULI EXPRESSED AS PERCENT CHANGE RELATIVE TO CONTROL VALUES*

	Pulmonary Flow Resistance (b)								
High Control SO ₂	% Change	Control	Mech. Stim.	% Change					
17.0 !7.6 18.1 22.8 18.1 21.7	-2.8 26.0 19.9	15.8	122.7	67÷.6 .					
Mean 17.7 20.7 S.D. 0.6 2.7		15.9	122.7						
t-test ∆ N.S.		REF	LEX INT	ACT					
					Lung Compliance (c)				
High Control SO ₂	% Change	Control	Mech. Stim.	% Chan ge					
9.9 9.2 9.9 9.4 10.1 8.2	-7.7 -5.7 -7.7	9.9	5.3	-46.5					
Mean± 10.0 8.9 S.D. 0.1 0.6		9.9	5 .3						
t-test ∆ N.S.		<u></u>							
Sequence of Challenge	(6)		(7)						

^{*}Control for 15 minutes preceding challenge.

- (a) See Figure.
- (b) Pulmonary Flow Resistance, Cm H₂0/1/sec.
 (c) Lung Compliance, m1/cm H₂0.



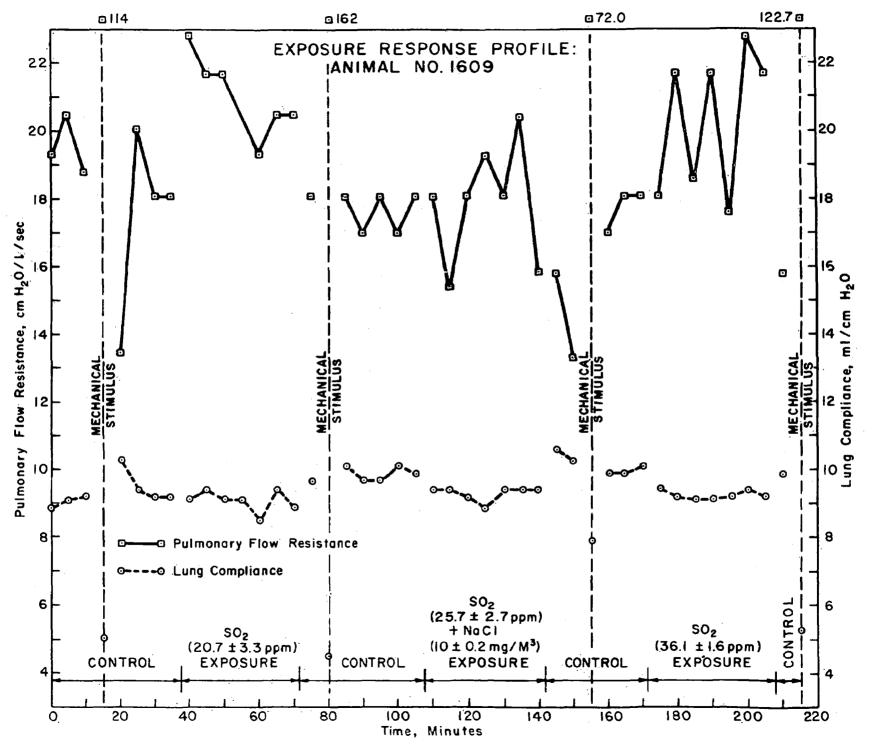


TABLE 12 Cat No. 1564 RESPONSE TO VARIOUS STIMULI EXPRESSED AS PERCENT CHANGE RELATIVE TO CONTROL VALUES*

	Pu	llmonary Flow Resistance	(b)	
Mech. % Contro! Stim. Chan	Low ½ Control SO ₂ Change	Mech. % Control Stim. Change	\$0 ₂ + % Control NaCl Change	Mech. % Control Stim. Change
11.1 76.8 645. 9.9 9.9	9.7 7.1 -29.9 9.7 8.4 -17.1 11.0 9.5 -6.2	8.4 121.7 1337.4 7.1 9.9	11.0 9.5 8.8 7.6 8.8 0.8 7.6 8.8 0.8	11.0 29.5 237.8 7.6 7.6
lean± 13.3 76.9	10.1 8.3 0.8 1.2	8.5 121.7 1.4	8.7 9.0 2.0 0.4	8.7 29.5 2.0
REFLEX INTACT	Δ N.S.	REFLEX INTACT	Δ N.S.	REFLEX INTACT
		Lung Compliance (c)		
Mech. % Control Stim. Shan	Low % ge Gontrol SO ₂ Change	Mech. % Contro¹ Stim. Change	\$0 ₂ + % Contro! NaC1 Change	Mech. % Control Stim. Change
9.9 6.3 -36. 9.9 9.9	14.6 14.2 1.2 14.2 12.9 -8.1 13.3 13.2 -5.2	20.1 9.9 -49.5 17.6 21.1	11.2 12.9 19.1 11.2 12.2 12.2 10.1 23.4 116.0	11.2 8.1 -25.2 11.2 10.1
Mean <u>†</u> 9.9 6.3 3.0. 0	14.0 13.4 0.7 0.7	19.6 9.9 1.8	10.8 16.2 0.6 6.3	10.8 0.6
	∆ N.S.		∆ N.S.	
Sequence of (1)	(2)	(3)	(4)	(5)

^{*}Control for 15 minutes preceding challenge.

(a) See Figure.

(b) Pulmonary Flow Resistance, Cm H₂0/1/sec.

(c) Lung Compliance, m1/cm H₂0.

TABLE 12 (Continued)

Cat No. 1564

RESPONSE TO VARIOUS STIMULI EXPRESSED AS PERCENT CHANGE RELATIVE TO CONTROL VALUES*

	Pulmonary Flow Resistance (b)	
High % Control SO2 Change		
9.7 7.1 -23.4 8.4 5.2 -43.9 9.7 5.8 -37.4		
ean± 9.3 6.0 .D. 0.9 1.0		
P< 0.01		
	Lung Compliance (c)	
High ½ Control 302 Change		
19.2 16.3 7.9 10.9 13.7 -9.3 15.2 14.6 -3.3		
lean		
Δ N.S.		

*Control for 15 minutes preceding challenge.

(a) See Figure.
(b) Pulmonary Flow Resistance, Cm H₂0/1/sec.
(c) Lung Compliance, m1/cm H₂0.

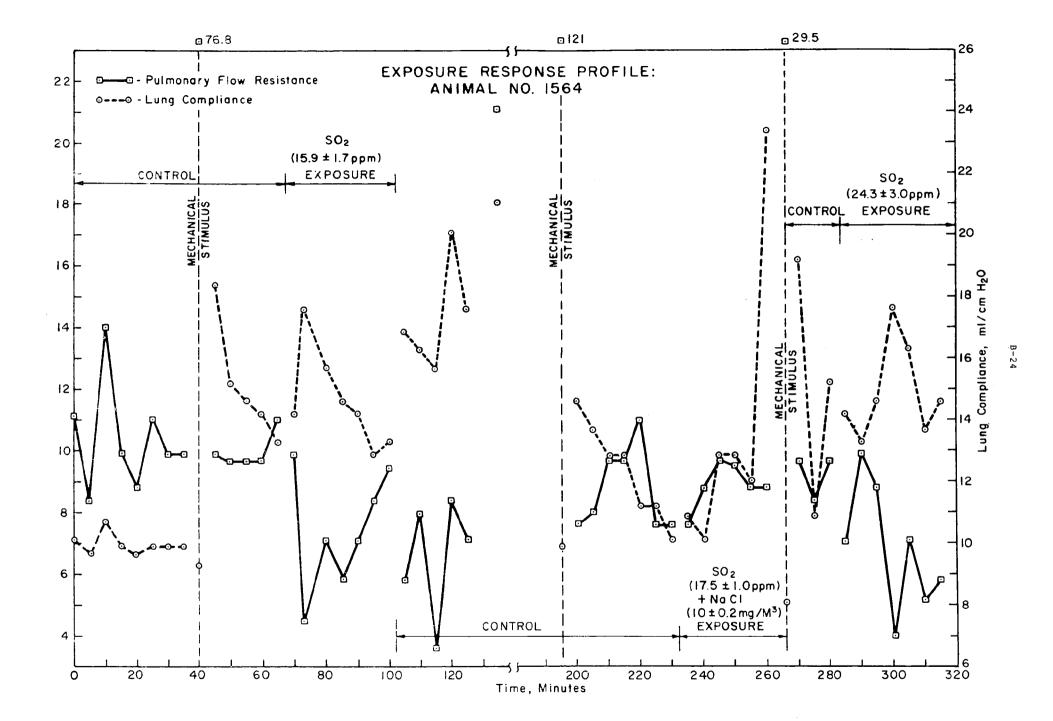


TABLE 13 Cat No. 1593 RESPONSE TO VARIOUS STIMULI EXPRESSED AS PERCENT CHANGE RELATIVE TO CONTROL VALUES*

	Pu	Imonary Flow Resistance	(b)	
Mech. %	Low ½	Mech. %	SO ₂ +	Mech. %
Contro! Stim. Change	Control SO ₂ Change	Control Stim. Change		Control Stim. Change
8.2 110 1298	11.1 6.4 -33.1	11.1 80.1 731.5	9.0 10.1 3.8	10.1 82.8 719.8
9.0	8.8 11.2 17.1	9.0	10.1 12.4 27.4	
6.4	8.8 5.8 -39.4	9.8	10.1 8.8 -9.6	
Mean ⁺ 7.9 110	9.6 7.8	9.6 80.1	9.7 10.4	10.1 82.8
S.D. 1.3	1.3 3.0	1.3	0.6 1.8	
t-test REFLEX INTACT	Δ N. S.	REFLEX INTACT	∆ N.S.	REFLEX INTACT
		Lung Compliance (c)		
Mech. ₹	Low ⅔	Mech. %	SO2+ %	Mech. %
Control Stim. Change	Contro! SO ₂ Change	Control Stim. Change	Control NaCl Change	Control Stim. Change
10.4 6.0 -55.4	15.2 17.3 11.4	17.3 4.9 -73.8	14.7 18 22.2	20.2 6.4 -68.3
10.4	15.2 16.7 7.5	19.4	15.2 15.2 3.2	
19.4	16.2 17.3 11.4	19.4	14.3 15.7 6.6	
Mean [±] 13.5 6.0	15.5 17.1	18.7 4.9	14.7 16.3	20.2 6.4
S.D. 5.3	0.6 0.3	1.2	0.5 1.5	
t-test	P< 0.05		ΔN.S.	
Sequence of (i) Challenge	(2)	(3)	(4)	(5)

*Control for 15 minutes preceding challenge.

⁽a) See Figure.
(b) Pulmonary Flow Resistance, Cm H₂0/1/sec.
(c) Lung Compliance, m1/cm H₂0.

TABLE 13 (Continued) Cat No. 1593

				-	Pu	Imonary Flow Resistance (b)	
Control	High SO ₂	% Change	Control	Mech. Stim.	½ Change		
11.1 8.9 15.9	13.5 14.3 14.7	19.8 23.2 13.1	11.1 8.8 15.9	67.4	464.9		
Mean <u>†</u> 11.9 S.D. 3.6	14.2		11.9 3.6	67.4			
t-test ∆	N.S.		RE	FLEX IN	TACT		
						Lung Compliance (c)	
Control	High SO ₂	½ Change	Control	Mech. Stim.	⅓ Change		
18 18 17.3	15.2 15.2 15.2	-14.4 -14.4 -17.3	18 18 17.3	4.6	-74.1		
Mean [±] 17.8 S.D. 0.4	15.2		17.8	4.6			
t-test P	< 0.01						
Sequence of Challenge	(a)	(6)		(7)			

*Control for 15 minutes preceding challenge.

(a) See Figure.

(b) Pulmonary Flow Resistance, Cm H₂0/1/sec.
 (c) Lung Compliance, m1/cm H₂0.

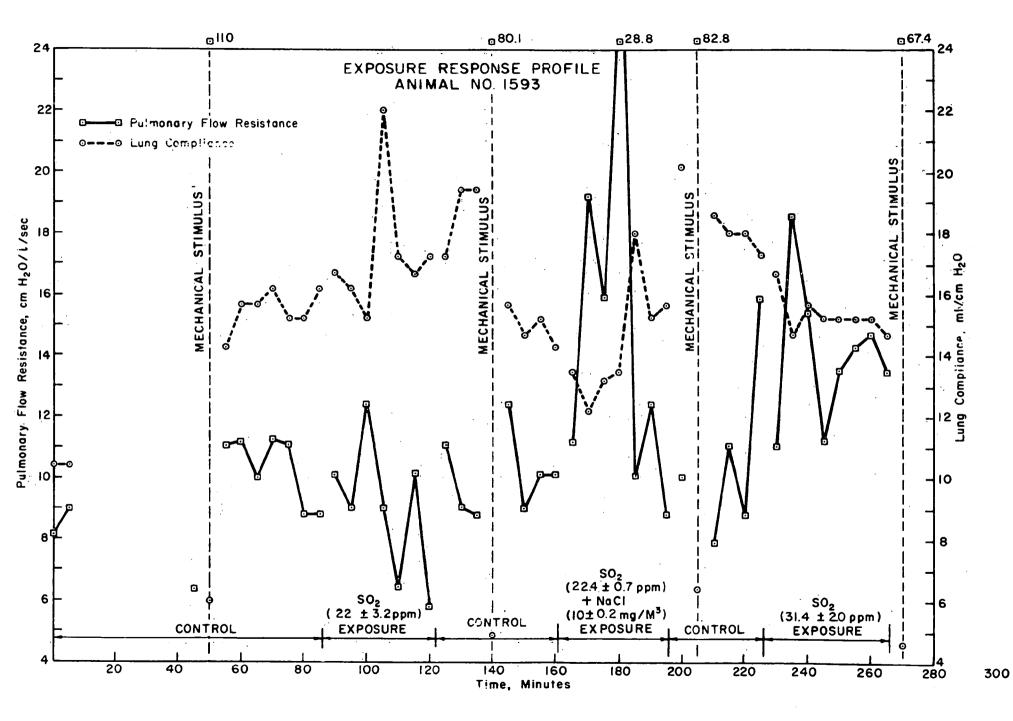


TABLE 14 (Continued)

Cat No. 1566

					Pu	lmonary Flow Resistance	(b)	
Contro1	Mech. Stim.	% Change	Control	Mech. Stim.	% Change	High % Control SO ₂ Change	Control	
9.6 9.6 9.6	175.1	1724	10.7 9.2 15.3	201	1613	15.0 18.5 32.1 13.5 11.3 -19.3 13.5 12.2 -12.9	14.8 16.9	
Mean± 9.6 S.D. 0	175.1		11.7	201		14.0 14.0 0.9 3.8	15.9 1.5	
t-test REFL	EX INTA	CT	RE	FLEX IN	TACT	Δ N.S.		
						Lung Compliance (c)	_	
Control	Mech. Stim.	/ Change	Control	Mech. Stim.	½ Change	High % Control SO ₂ Change	Control	
7.5 6.7 6.8	4.5	-35.7	8.5 5.5 11.1	4.5	-46.2	8.1 7.9 0.9 7.9 7.5 -4.3 7.5 7.5 -4.3	8.1 7.7	
Mean± 7.0 S.D. 0.4	4.5		8.4 2.8	4.5		7.8 7.6 0.3 0.2	7.9 0.3	
t-test						Δ N.S.		
Sequence of Challenge	fa)	(6)	<u> </u>	(7)		(8)	(9)	

*Control for 15 minutes preceding challenge.

(a) See Figure.
(b) Pulmonary Flow Resistance, Cm H₂0/1/sec.
(c) Lung Compliance, m1/cm H₂0.

TABLE 14 Cat No. 1566 RESPONSE TO VARIOUS STIMULI EXPRESSED AS PERCENT CHANGE RELATIVE TO CONTROL VALUES*

	Pu	Imonary Flow Resistance	(6)	
Mech. % Control Stim. Change	Mech. % Control Stim. Change	Low % Control SO ₂ Change	Mech. % Control Stim. Change	SO ₂ + % Control NaCl Change
11.9 93.5 646 13.6 12.1	8.8 232.2 2318.8 10.4	9.2 10.0 8.7 9.2 10.2 10.9 9.2 10.4 13.0	11.9 112.2 984.1 8.8	9.6 13.5 40.6 9.6 11.1 15.6 9.6 15.0 56.3
ean [±] 12.5 93.5 .D. 0.9	9.6 232.2 1.1	9.2 10,2 0 0.2	10.4 112.2 2.2	9.6 13.2 0 2.0
-test REFLEX INTACT	REFLEX INTACT	P<0.05	REFLEX INTACT	Δ N.S.
		Lung Compliance (c)		
Mech. % Control Stim. Change	Mech. % Control Stim. Change	Low % Control SO ₂ Change	Mech. % Control Stim. Change	\$0 ₂ + % Control NaCl Change
5.9 6.6 11.9 5.9 5.9	9.1 3.3 -62.1 8.3	9.4 6.9 -23.9 8.9 6.8 -25 8.9 6.8 -25	7.2 4.1 -43.8 7.4	7.5 5.7 -18.6 6.7 6.8 -2.9 6.8 5.8 -17.1
Hean [±] 5.9 6.6 0	8.7 3.3 0.6	9.1 6.8 0.3 0.1	7.3 4.1 0.1	7.0 6.1 0.4 0.6
t-test		P<0.01		Δ N.S
Sequence of (1):	(2)	(3)	(4)	(5)

^{*}Control for 15 minutes preceding challenge.

 ⁽a) See Figure.
 (b) Pulmonary Flow Resistance, Cm H₂O/1/sec.
 (c) Lung Compliance, m1/cm H₂O.

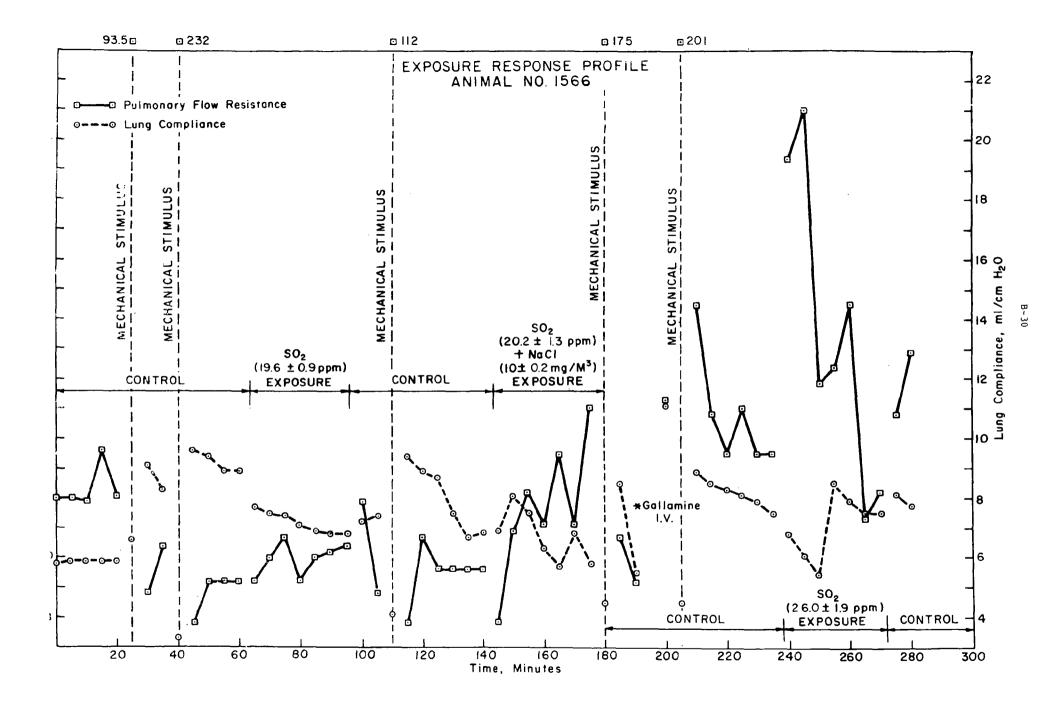


TABLE 15 Cat No. 1113 RESPONSE TO VARIOUS STIMULI EXPRESSED AS PERCENT CHANGE RELATIVE TO CONTROL VALUES*

	Pu	lmonary Flow Resistance	(b)	
Mech. %	Low %	Mech. %	SO ₂ + %	High %
Contro! Stim. Change	Control SO ₂ Change	Control Stim. Change	Control NaCl Change	Control SO ₂ Change
16.1 233.2 1349	16.1 19.2 19.3	15.2 129.3 718	16.1 10.8 -32.9	16.1 21.2 31.7
16.1	16.1 17.8 10.6	16.1	16.1 18.8 16.8	16.1 19.5 31.1
16.1	16.1 24.6 52.8	16.1	8.9 -45.3	16.1 17.8 10.6
Mean± 16.1 233.2	16.1 20.5	15.8 129.3	16.1 12.8	16.1 19.5
S.D. 0	0 3.6	0.5	0 5.3	0 1.7
REFLEX INTACT	Δ N.S.	REFLEX INTACT	Δ N.S.	ΔN.S.
		Lung Compliance (c)		
Mech. ½	Low %	Mech. %	SO ₂ + %	High %
Contro¹ 3tim. Change	Control SO ₂ Change	Control Stim. Change	Contro! NaCl Change	Control SO ₂ Change
12.4 4.7 -60.3	11.1 12.0 6.2	11.4 5.0 -55.5	10.8 9.1 -20.2	11.1 10.8 -0.9
12	11.1 11.4 0.9	11.1	12.0 10.2 -10.5	11.1 10.8 -0.9
11.1	11.7 12.4 9.7	11.1	9.1 -20.2	10.5 10.5 -3.7
Mean± 11.8 4.7	11.3 11.9	1.1.2 5.0	11.4 9.5	10.9 10.7
S.D. 0.7	0.3 0.5	0.2	0.8 0.6	0.3 0.2
	ΔN.S.	·	ΔN.S.	Δ N.S.
Sequence of (1)	(2)	(3)	(4)	(5)

^{*}Control for 15 minutes preceding challenge.

△ N.S. = Difference between means not sigificant (P>0.05)

⁽a) See Figure.
(b) Pulmonary Flow Resistance, Cm H₂0/1/sec.
(c) Lung Compliance, m1/cm H₂0.



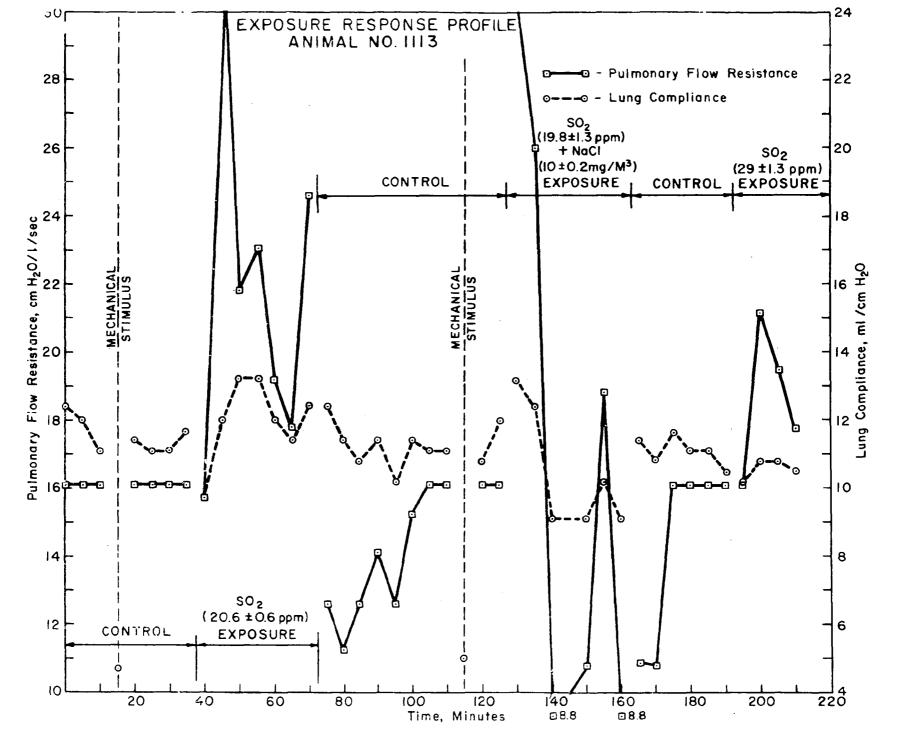


TABLE 16 Cat No. 1482 RESPONSE TO VARIOUS STIMULI EXPRESSED AS PERCENT CHANGE RELATIVE TO CONTROL VALUES*

	Pu	Imonary Flow Resistance	(ь)	
Mech. % Control Stim. Change	Low % Control SO ₂ Change	SO ₂ + % Control NaCl Change	High % Control SO ₂ Change	Control
49.0 70 88.7 35.9 26.5	20.9 20.1 -11.1 23.9 21.6 -4.4 23.1 19.1 -15.5	24.6 22.1 7.1 19.1 23.3 12.9 18.2 26.7 29.4	20.5 11.8 -42.7 23.3 10 -51.5 18 13.9 -32.5	18.4 11.7
Mean	22.6 20.3 1.6 1.3 Δ N.S.	20.6 24.0 3.5 2.4 Δ N.S.	20.6 11.9 2.7 2.0 P< 0.01	15.1 4.7
	<u>' </u>	Lung Compliance (c)		
Mech. % Control Stim. Change	Low % Control SO ₂ Change	SO ₂ + % Control NaCl Change	High % Control SO ₂ Change	Control
9.4 17.6 74.3 9.3 12.5	9.9 10.1 0.7 9.3 11.8 17.6 10.9 11.2 11.6	14.1 14.6 11.5 11.5 14.6 11.5 13.7 12.9 -1.5	13.3 10.4 -11.6 10.5 8.4 -28.6 11.5 8.4 -28.6	8.4 10.1
Mean±10.1 17.6 S.D. 2.2	10.0 11.0 0.8 0.9	13.1 14.0 1.4 1.0	11.8 9.1 1.4 1.2	9.3 1.2
•	≙ N.S.	Δ N.S.	P< 0.05	
Sequence of (1) Challenge	(2)	(3)	(4)	(5)

(a) See Figure.
 (b) Pulmonary Flow Resistance, Cm H₂0/1/sec.
 (c) Lung Compliance, ml/cm H₂0.

TABLE 17 Cat No. 1611 RESPONSE TO VARIOUS STIMULI EXPRESSED AS PERCENT CHANGE RELATIVE TO CONTROL VALUES*

	Pu	lmonary Flow Resistance	(6)			
Mech. % Control Stim. Change	Low % Control SO ₂ Change	Mech. % Control Stim. Change	SO ₂ + % Control NaCl Change	Mech. % Control Stim. Change		
17.7 135.6 666.1 17.7	17.7 24.6 35.9 18.9 21.2 17.1 17.7 22.3 23.2	17.7 167.4 845.8	17.7 25.7 45.2 17.7 22.2 25.4 17.7 22.3 26.0	17.7 97.0 448		
Mean [±] 17.7 135.6 S.D. 0	18.1 22.7 0.7 1.7	17.7 167.4	17.7 23.4 0 2.0	17.7 97.0		
REFLEX INTACT	2€0.05	REFLEX INTACT	P< 0.05	REFLEX INTACT		
		Lung Compliance (c)				
Mech. % Control Stim. Chang e	Low ½ Control SO ₂ Change	Mech. % Control Stim. Change	\$02+ % Control NaCl Change	Mech. % Control Stim. Change		
9.5 4.5 - 52.6 9.5	8.9 9.1 -8.6 8.8 8.6 -3.0 8.9 8.4 -5.3	9.1 4.8 -47.3	10.5 8.6 -11.9 10.0 8.2 -16.0 8.8 8.6 -11.9	9.8 4.6 -53.1		
Mean [±] 9.5 4.5 \$. D . 0	8.7 8.4 0.1 0.3	9.1 4.8	9.8 8.5 0.9 0.2	9.8 4.6		
	ΔN.S.		ΔN.S.			
Sequence of (a) (1)	(2)	(3)	(4)	(5)		

^{*}Control for 15 minutes preceding challenge.

△ N.S. = Difference between means not sigificant (P>0.05)

⁽a) See Figure.
(b) Pulmonary Flow Resistance, Cm H₂0/1/sec.
(c) Lung Compliance, m1/cm H₂0.

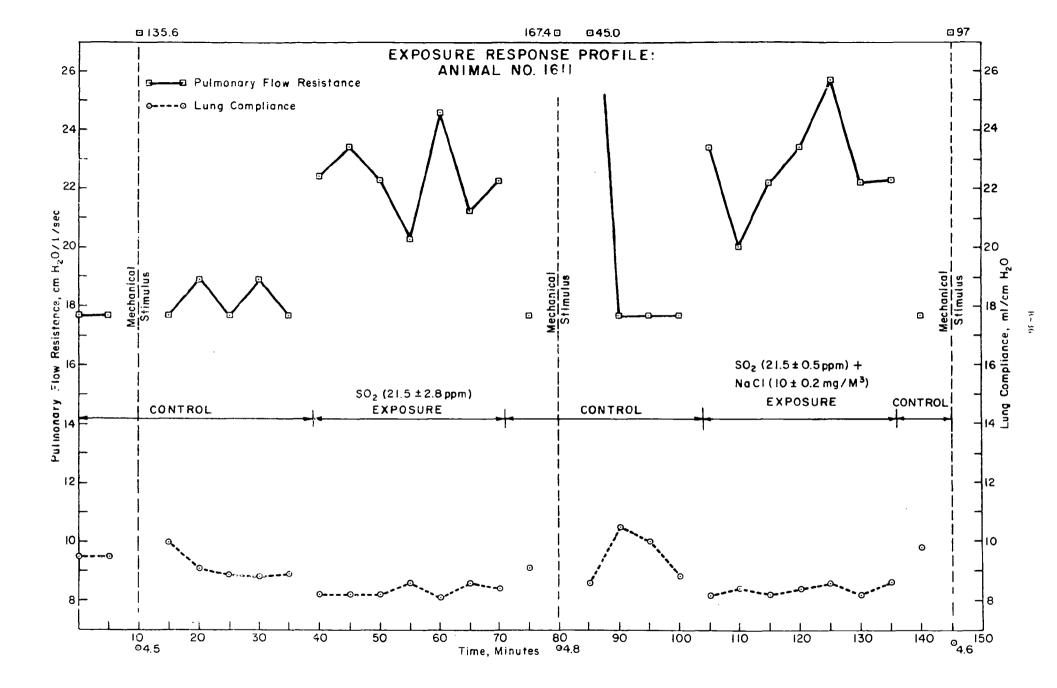


TABLE 18 Cat No. 1478 RESPONSE TO VARIOUS STIMULI EXPRESSED AS PERCENT CHANGE RELATIVE TO CONTROL VALUES*

					Pu	lmonary F	low Re	sistance	(6)					
Me Control S	ech. tim.	¾ Change	Control	Low SO ₂	% Change	Control	Mech. Stim.	% Change	Control	S0 ₂ + NaC1	% Change	Control	Mech. Stim.	% Change
22.4 1	15	413	17.5 16.9 18.7	21.2 17.3 21.2	19.8 -2.3 19.8	17.3	104		19 19 19	19 21.2 17.3	0 11.6 -8.9	17.7	89	403
Mean [±] 22.4 1 S.G.	15		17.7	19.9		17.3	104		19.0 0	19.2		17.7	89	
-test REFLEX INTACT				REFLEX INTACT		Δ N.S.			REFLEX INTACT					
						Lung Co	mplian	ce (c)						
M Control S	lech.	½ Change	Control	Low SO ₂	% Change	Control	Mech. Stim.	% Change	Contro 1	S0 ₂ + NaC1	% Change	Contro1	Mech. Stim.	% Change
22.6 3	33.5	49.2	22.6 22.6 22.6	20.3 20.3 20.3	-10.2 -10.2 -10.2	19.4	12.8		22.6 22.6 22.6	17.7 17.7 17.7	-21.7 -21.7 -21.7	20.3	12.2	-39.9
Mean ± 22.6 3 S.D.	33.5		22.6	20.3		19.4	12.8		22.6 0	17.7 0		20.3	12.2	
t-test			1	P < 0.01	+					P<0.01				
Sequence of (a)	,	(1)		(2)			(3)			(4)			(5)	

(a) See Figure.
(b) Pulmonary Flow Resistance, Cm H₂0/1/sec.
(c) Lung Compliance, m1/cm H₂0.

TABLE 18 (Continued) Cat No. 1478 RESPONSE TO VARIOUS STIMULI EXPRESSED AS PERCENT CHANGE RELATIVE TO CONTROL VALUES*

	Pu	ulmonary Flow Resistance (b)
Cig. % Control Smoke Change	Mech. % Control Stim. Change	High % Control SO ₂ Change
17.4 13.1 -25.8 17.9 13.6 -22.9 12.3 -30.3	15.2 77.9 321 20.1 20.1	23.4 21.3 -6.0 21.2 18.3 -19.3 23.4 14.3 -36.9
Mean± 17.7 13.0 S.D. 0.4 0.7	19.5 77.9 2.8	22.7 18.0 1.3 3.5
t-test	REFLEX INTACT	Δ N.S.
		Lung Compliance (c)
Cig. ½ Control Smoke Change	Mech. ½ Control Stim. Change	High % Control SO ₂ Change
21.4 21.4 0 21.4 18.5 -13.6 18.5 -13.6	20.3 17.7 -7.3 18.5 18.5	16.3 :4.6 -14.1 17.7 14.5 -14.1 14.1 -17.1
Mean± 21.4 19.5 S.D. 0 t-test	19.1 17.7 1.0	17.0 1.0 14.4 0.3 \$\triangle N.S.
Sequence of (6)	(7)	(8)

(a) See Figure.

(b) Pulmonary Flow Resistance, Cm H₂0/1/sec.
 (c) Lung Compliance, m1/cm H₂0.



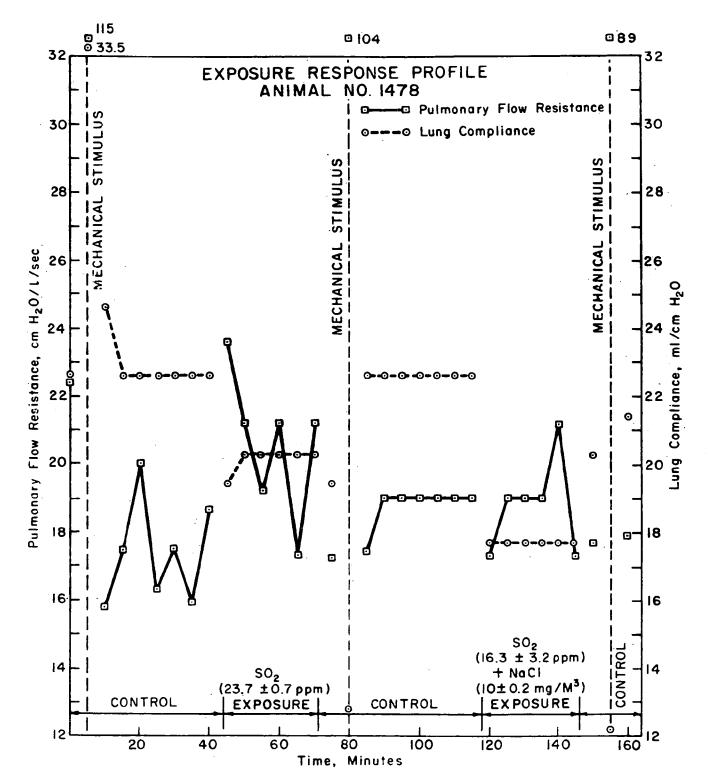


TABLE 19 Cat No. 32135 RESPONSE TO VARIOUS STIMULI EXPRESSED AS PERCENT CHANGE RELATIVE TO CONTROL VALUES*

	Pu	Imonary Flow Resistance	(5)			
Elec. % Control Stim. Change	. Low ₹ Control SO ₂ Change	Elec. % Control Stim. Change	Elec. % Control Stim. Change	SO2+ ⅓ Control Na€1 Change		
19 30.7 61.6	11.6 11.8 7.6 9.7 12.5 14.0 9.2 12.5 14.0	11.8 9.9 -8.3 9.8	11.2 10.4 -2.8 9.8 11.1	9.7 13.2 36.1 11.8 21.6 12.2 25.3		
Meani 19 30.7 S.D.	10.2 12.3 1.3 0.4	10.8 9.9 1.4 REFLEX	10.7 10.4 0.8 REFLEX	9.7 12.4 0.7		
REFLEX INTACT	A N.S.	QUESTIONABLE	QUESTIONABLE	P<.0.05		
		Lung Compliance (c)				
Elec. % Control Stim. Change	Low 劣 Control SO ₂ Change	Elec. % Control Stim. Change	Elec. % Control Stim. Change	SO ₂ + % Control NaCl Change		
13.1 10.2 -22.1	13.6 14.5 18.2 11.6 14.0 14.1 11.6 12.7 13.5	12.7 14.5 2.5 15.6	12 13.1 9.2 12 12 0 12 12 0	12 10.5 -12.5 9.7 -19.2 10 -16.7		
rean± 13.1 !0.2 S.D.	12.3 13.7 1.2 0.8	14.2 14.5 2.1	12.0 12.4 0 0.6	12.0 10.1 0 0.4		
t-test	Δ N.S.			P< 0.05		
Sequence of (1)	(2)	(3)	(4)	(5)		

^{*}Control for 15 minutes preceding challenge.

⁽a) See Figure.
(b) Pulmonary Flow Resistance, Cm H₂0/1/sec.
(c) Lung Compliance, ml/cm H₂0.

TABLE 20 (Continued) Cat No. 32135

RESPONSE TO VARIOUS STIMULI EXPRESSED AS PERCENT CHANGE RELATIVE TO CONTROL VALUES.

	Pi	ulmonary Flow Resistance (b)
Elec. % Control Stim. Change	% Control NaCl Change	
11.7 14.8 31 11.7 10.5	11.7 11.3 0 11.7 14.4 27.4 10.5 12.1 7.1	
Mean± 11.3 14.8 S.D. 0.7 REFLEX t-test QUESTIONABLE	11.3 12.6 0.7 1.6 Δ N.S.	The state of the s
		Lung Compliance (c)
Elec. ½ Control Stim. Change	Control NaCl Change	
12.7 11.6 -7.2 12.7 12.0	12.7 10.7 -14.2 12.7 10.7 -14.2 12.0 10.7 -14.2	
Mean <u>†</u> 12,5 11.6 S.D. 0.4	12.5 10.7 0.4 0	
t-test	P<0.05	
Sequence of (6)	(7)	

^{*}Control for 15 minutes preceding challenge.

⁽a) See Figure.
(b) Pulmonary Flow Resistance, Cm H₂0/1/sec.
(c) Lung Compliance, m1/cm H₂0.

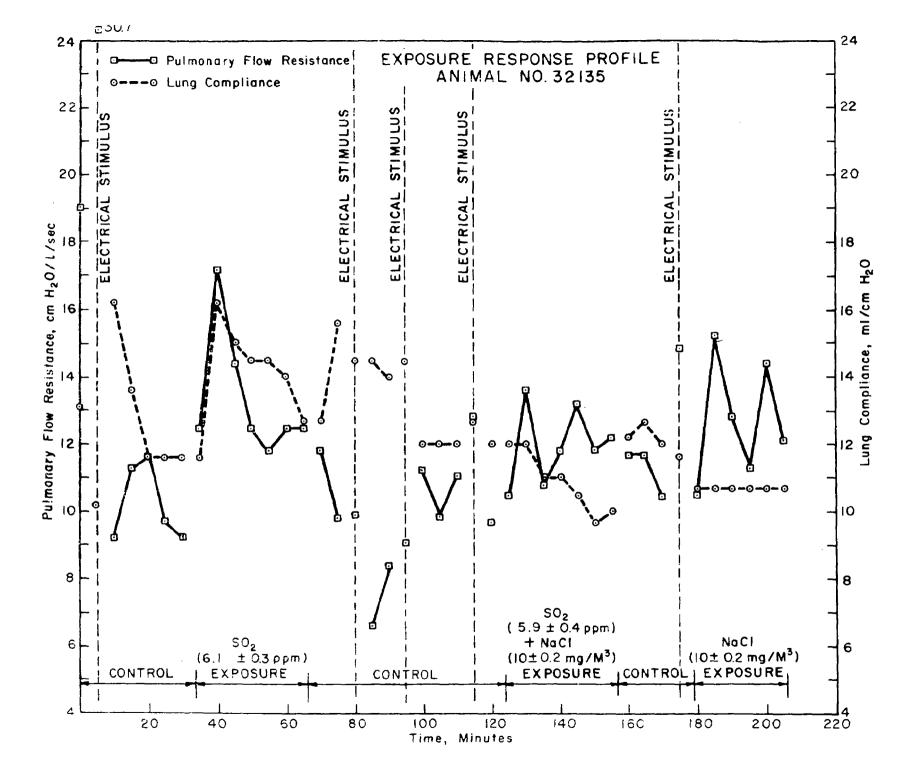


TABLE 21 Cat No. 35354 RESPONSE TO VARIOUS STIMULI EXPRESSED AS PERCENT CHANGE RELATIVE TO CONTROL VALUES*

	Pu	Imonary Flow Resistance	(b)	
Elec. ½ Control Stim. Change	% . Contro! SO ₂ Change	SO ₂ + % Control laC1 Change	Elec. % Control Stim. Change	
20.3 51.6 154	21.8 27.9 18.7 25.5 27.9 18.7 23.2 27.9 18.7	19.2 26.4 21.7 21.6 20.5 -5.5 24.4 20.8 -4.1	18.4 19.6 -10.1 23.0 24.0	
Mean± 20.3 51.6 S.D.	23.5 .27.9 1.9 0	21.7 22.6 2.6 3.3	21.8 19.6 3.0	
t-test REFLEX INTACT	Δ N.S.	Δ N.S.	REFLEX QUESTIONABLE	
		Lung Compliance (c)		
Elec. % Control Stim. Change	Control SO ₂ Change	\$0 ₂ + % Control NaCl Change	% Control Stim. Change	
10.9 10.5 -3.7	9.5 9.5 1.1 9.3 9.5 1.1 9.3 9.9 5.3	10.7 10.0 -6.5 10.7 10.0 -6.5 10.7 10.4 -2.8	10.5 10.5 1.0 10.1 10.7	
Mean± 10.3 10.5 S.D.	9.4 9.6 0.1 0.2	10.7 10.1 0 0.2	10:4 10.5 0.3	
t-test	ΔN.S.	Δ N.S.		
Sequence of (1)	(2)	(3)	(4)	

^{*}Control for 15 minutes preceding challenge.
(a) See Figure.
(b) Pulmonary Flow Resistance, Cm H₂0/1/sec.
(c) Lung Compliance, m1/cm H₂0.

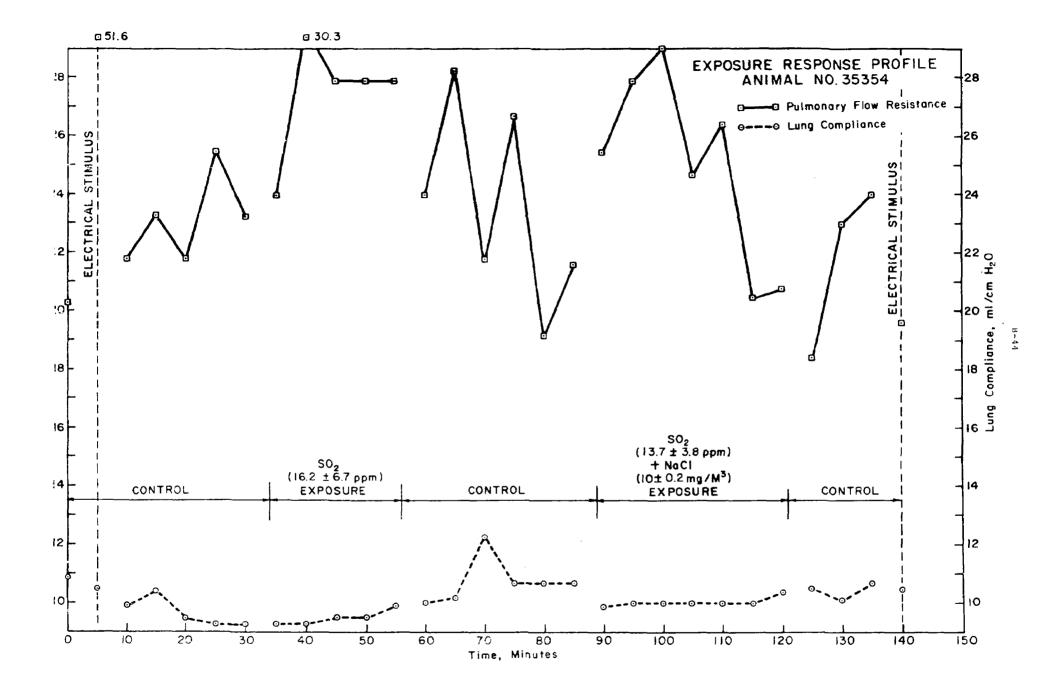


TABLE 22 Cat No. 1807 RESPONSE TO VARIOUS STIMULI EXPRESSED AS PERCENT CHANGE RELATIVE TO CONTROL VALUES*

				Pu	Imonary Flow Resistance	(b)	
Me Control St	ch. % im. Change	Control	High SO ₂	% Change	Mech. % Control Stim. Change	% Control NaCl Change	Mech. % Control Stim. Change
15.4 55 16.6 16.6	.1 +240,1	16.6 16.6 16.6	9.7 12.3 7.1	-41.6 -25.9 -57.2	12.2 17.0 38.8 12.3	10.8 13.7 7.6 13.7 12.2 -4.2 13.7 13.7 7.6	12.3 31.8 158
Mean [±] 16.2 55 S.D. 0.7	.1	16.6 0	9.7 2.6		12.5 17.0 0.1	12.7 13.2 1.7 0.9	12.3 31.8
t-test REFL	LEX INTACT		P<-0.	05	REFLEX Question ab ee	Δ N.S.	REFLEX INTACT
					Lung Compliance (c)		
Me Contro¹ St	ech. % im. Change	Control	High SO ₂	浅 Change	Mech. % Control Stim. Change	% Control NaCl Change	Mech. % Control Stim. Change
9.9 5. 9.9 9.9	6 -43.4	9.9 9.9 10.1	14.2 14.6 15.1	42.5 46.5 51.5	9.5 7.7 -42.8 17.4	14.2 11.7 -14.4 13.4 11.4 -16.6 13.4 11.4 -16.6	12.0 10.6 -11.7
Mean	6	10.0	14.6 0.5	·	13.5 7.7 5.6	13.7 11.5 0.5 0.2	12.0 10.6
		F	P<0.01			P<-0.01	
Sequence of (a)	(1)		(2)		(3)	(4)	(5)

*Control for 15 minutes preceding challenge.

(a) See Figure.

(b) Pulmonary Flow Resistance, Cm H₂0/1/sec.

(c) Lung Compliance, m1/cm H₂0.

TABLE 22 (Continued) Cat No. 1807

RESPONSE TO VARIOUS STIMULI EXPRESSED AS PERCENT CHANGE RELATIVE TO CONTROL VALUES*

	Pu	Imonary Flow Resistance	(b)		
Control					
13.6 16.3 13.7					
Mean± 14.5 S.D. 1.5					
		Lung Compliance (c)			
Control					
11.4 11.1 11.4					
Mean± 1!.3 S.D. 0.2					
Sequence of (6) Challenge				<u> </u>	1

*Control for 15 minutes preceding challenge.

- (a) See Figure.
- (b) Pulmonary Flow Resistance, Cm H₂0/1/sec.
 (c) Lung Compliance, m1/cm H₂0.

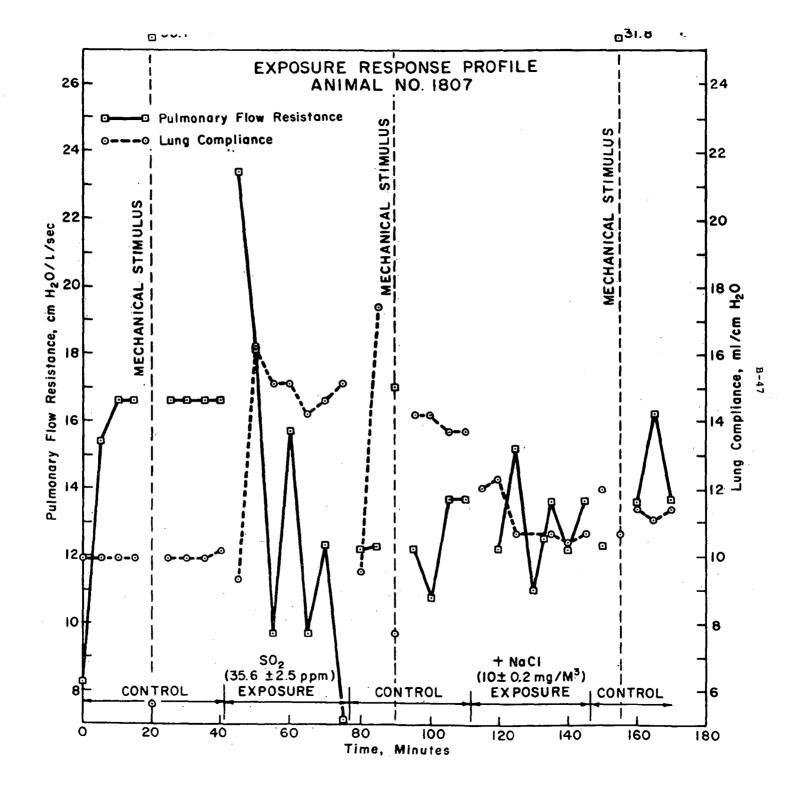


TABLE 23 Cat No. 1486 RESPONSE TO VARIOUS STIMULI EXPRESSED AS PERCENT CHANGE RELATIVE TO CONTROL VALUES

	Pu	Imonary Flow Resistance	(6)	
Mech. % Control Stim. Change	SO ₂ + % Control NaCl Change	Mech. % Control Stim. Change	Low % Control SO ₂ Change	Control
25.6 128 376 27.0 27.9	29.2 32.2 7.3 31.0 33.8 12.7 29.8 30.2 0.7	28.5 117 302 29.2	35.9 24.7 -31.2 26.9 -25.1 26.4 -26.5	26.9 24.2
Mean [±] 26.9 S.D. 1.2	30.0 32.1 0.9 1.8	29.1 117 0.2	35.9 26.0 1.2	25.5
t-test REFLEX INTACT	Δ N.S.	REFLEX INTACT	P< 0.01	
		Lung Compliance (c)		
Mech. % Contro! Stim. Change	SO ₂ + % Control NaC1 Change	Mech. ½ Control Stim. Change	Low % Control SO ₂ Change	Control
5.3 3.8 -24.5 4.9 4.9	5.0 4.5 -10.0 5.0 4.6 -8.0 5.0 4.6 -8.0	5.3 3.6 -40 6.5	8.1 6.2 -23.5 6.0 -25.9 5.6 -30.9	6.3 6.3
Mean [†] 5.0 5.0. 0.2	5.0 4.5 0.0 0.1	5.9 3.6 0.8	5.9 0.3	6.3
t-test	P≪0.01		P< 0.01	
Sequence of (1) Challenge (1)	(2)	(3)	(4)	(5)

(a) See Figure.

A N.S. = Difference between means not sigificant (P>0.05)

⁽b) Pulmonary Flow Resistance, Cm H₂0/1/sec. (c) Lung Compliance, m1/cm H₂0.

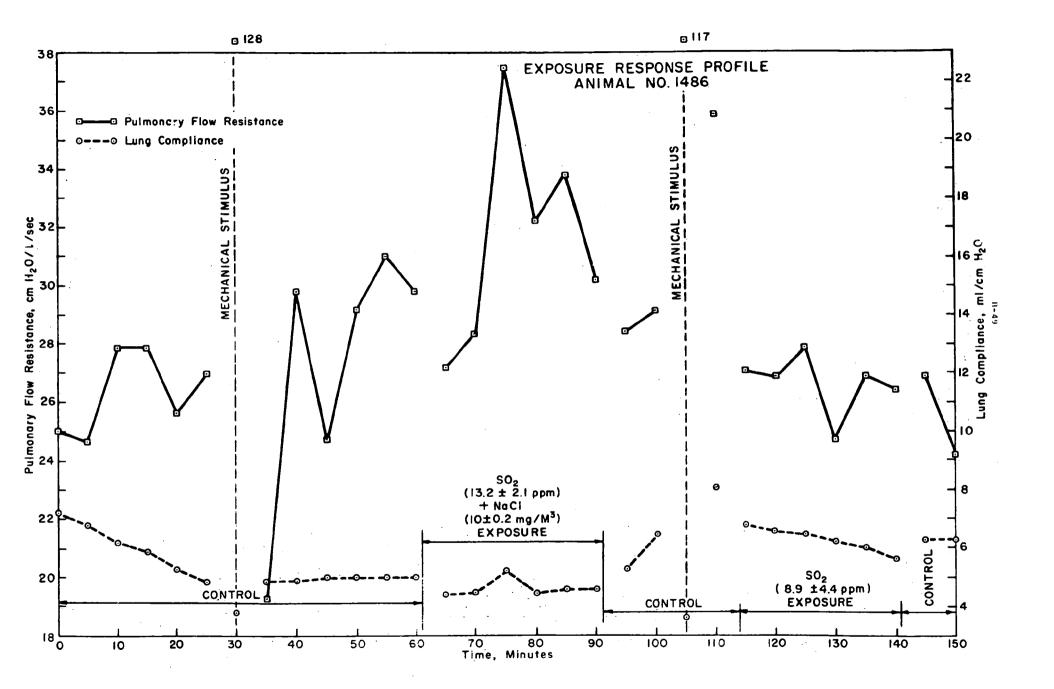


TABLE 24 Cat No. 1144 RESPONSE TO VARIOUS STIMULI EXPRESSED AS PERCENT CHANGE RELATIVE TO CONTROL VALUES*

						Pu	lmonary f	low Re	sistance	(b)					
. 1013	Control	-	Percent Change	Control	Low SO ₂	Percent Change	! Control	Mech. Stim.	Percent Change		S0 ₂ + NaC1	Percent Change	Control	Mech. Stim.	Percen t
	30.2 31.2	131	326.7	34.6 33.8 36.5	41.8 40.5 40.5	19 ,5 15.8 15.8	35.2	99.3	182.1	47.9 36.5 36.5	45 45 45	11.7 11.7 11.7	43.5	84.7	94.7
lean [±] .0,	30.7 0.7	131		35.0 1.4	40.9 0.8		35.2	99.3		40.3 6.6	45.0 0		43.5	84.7	
-test	est REFLEX INTACT P<0.01		REFLEX INTACT Δ N.S.				REFLEX INTACT								
							Lung Co	mpliane	ce (c)				*	· · · · · · · · · · · · · · · · · · ·	
	Control ⁻		Percent Change	Control	Low SO ₂	Percent Change	Control	Mech. Stim:	Percent Change	Control	S0 ₂ + NaC1	Percent Change	Control	Mech. Stim.	Per cen t Change
	5.8 6.0	4.2	-28. ⁹	6.4 6.6 6.6	6.7 6.4 6.4	3.1 -1.5 -1.5	7.8	4.0	-49.7	6.4 6.3 6.2	5.9 5.9 5.9	-6.3 -6.3	6.4	4.4	-31.3
iean‡	5.9 0.1	4.2	: !	6.5 0.1	6.5 0.2		7.8	4.0		6.3 0.1	5.9 0		6.4	4.4	
-test				,	N.S.				-	P	< 0.05		ĺ		

^{*}Control for 15 minutes preceding challenge.

- (a) See Figure.
- (b) Pulmonary Flow Resistance, Cm H₂0/1/sec.
 (c) Lung Compliance, ml/cm H₂0.

Cat No. 1144

RESPONSE TO VARIOUS STIMULI EXPRESSED AS PERCENT CHANGE RELATIVE TO CONTROL VALUES*

				14	Pu 1	monary Flo	w Resistance	(p)					
	Control	High SO2	Percent Change							·		;	
	40.5 36.5 36.5	55.3 47.1 45	46.2 24.5 18.9										
ean± .D.	37.8 2.3	49.1 5.4			1								
-test	. !	P< 0.05	.								}		
				·		Lung Com	liance (c)						
	Control	High SO ₂	Percent Change										
	5.8 5.9 5.9	6.0 5.9 5.9	2.3 0.6 0.6										
ean‡ .D.	5.8 0.1	5.9 0.1									·		
-test		N.S.		<u> </u>	 1			<u> </u>	·		<u> </u>		
Sequ	ence of lenge)											

*Control for 15 minutes preceding challenge.

(a) See Figure.
(b) Pulmonary Flow Resistance, Cm H₂0/1/sec.
(c) Lung Compliance, ml/cm H₂0.

TABLE 25 Cat No. 2393 RESPONSE TO VARIOUS STIMULI EXPRESSED AS PERCENT CHANGE RELATIVE TO CONTROL VALUES*

						Pu	ilmonary Flow Res	istance (b)	
	Control	Low SO ₂	Percent Change	Control		Percent Change	Control		_
	47.4 33.6 41.1	34.9	-25.5 -19.3 -14.8	45, 3 44, 2 41, 1	49.9	4.8 11.9 8.2	48.9		
Meant S.D.	42.7 4.1	34.4 2.3		43.7 2.4	47.3 1.6		48.9		
t-test	t	№ 0.05			14.S				
							Lung Complianc	_e (c)	
	Control	Low SO ₂	Percent Change	Control	Нј д ћ ЅО2	Percent Change	Control		
	7.9 9.2 8.8	8.2 8.7 8.4	-4.7 1.2 -2.3	8.6 8.7 9.0	9.7 8.6 9.5	-0.8 -1.9 8.4	9.5		
Mean‡ S.D.	ዓ.6 0.7	8.4 0.3		8.8 0.2	8.9 0.5		9.5		
t-test	_ Δ	٧.5.		<u></u>	. N.S.				

(a) See Figure.
 (b) Pulmonary Flow Resistance, Cm H₂0/1/sec.
 (c) Lung Compliance, m1/cm H₂0.

TABLE 26 Cat No. 1988 RESPONSE TO VARIOUS STIMULI EXPRESSED AS PERCENT CHANGE RELATIVE TO CONTROL VALUES*

						Pc	lmonary F	low Res	istance	(ъ)			-		
	:Control		Percent Change	Control		Percent Change	Control		Percent Change	B	NaC1	Percent Change	Control		Percent Change
	64.2 58.9	130.5	112.0	53.1 51.1 53.1	31.8 35.6 31.8	-39.4 -32.1 -39.4	27.7	73.7	166.1	22.1 29.8 29.8	18.2 18.2 24.0	-33.2 -33.2 -11.9	22.7	49.3	117.2
Mean <u>†</u> S.D.	61.6 3.7	930,5	-	52.4 1.2	33.1		27.7	7 3. 7		27.2 4.4	20.1 3.3	•	22.7	49.3	
t-tes	t REF	LEX INT	ACT.	. 1	P< 0.01		REF	LEX INT	ACT -	Δ	N.S.		REF	LEX IN	TACT
							Lung Co	mpliand	e (c)						
	Control	Mech. Stim.	Percent Change	Control	High SO ₂	Percent Change	Control	Mech. Stim.		Control	NaC1	Percent Change	Control		Percent Change
	3.9 3.9	3.3	-15.4	4.3 4.3 4.3	5.2 5.4 8.4	20.9 25.6 95.3	6,1	3.3	-45.9	7.6 7.4 7.2	7.6 7.2 7.2	2.7 -2.7 -2.7	7.4	4.3	-41.9
Mean <u>+</u>	3.9 0.0	3.3		4.3 0.0	6.3 1.8		6.1	3.3		7.4 0.2	7.3 0.2		7.4	4.3	
t-tes	t				∆n.s.		1			Δ	N.S.	1	l		

^{*}Control for 15 minutes preceding challenge.

⁽a) See Figure.

 ⁽b) Pulmonary Flow Resistance, Cm H₂0/1/sec.
 (c) Lung Compliance, m1/cm H₂0.

TABLE 26 (Continued) Cat No. 1988

RESPONSE TO VARIOUS STIMULI EXPRESSED AS PERCENT CHANGE RELATIVE TO CONTROL VALUES*

						Pt	llmonary F	low Res	sistance	(P)			
	Control	50 ₂ + NaC1	Percent Change	Contro1		Percent Change	Control	so ₂	Percent Change	Contro1		Percent Change	·
	27.9 29.5 27.8	27.8 27.8 31.2	-2.1 -2.1 9.9	24.7	33.1	34.0	26.2 27.7 26.2	26.2 30.7 31.8	-1.9 15.0 19.1	29.8	97.6	227.5	·
Mean± S.D.	29.4 1.0	28.9 2.0		24.7	33.1		26.7 0.9	29.6 3.0		29.8	97.6		
t-tes	t	∆N.S.			STIONA			Δ N.S.		REF	LEX IN	TACT	
							Lung Co	mp1ian	ce (c)				
	Control	50 ₂ + NaC1	Percent Change	Control		Percent Change	Control	so ₂	Percent Change	Control		Percent Change	
	7.4 7.0 6.6	5.4 5.4 5.2	-22.9 -22.9 -25.7	5.9	5.4	-8.5	5.4 5.6 5.6	4.9 4.9 5.0	-10.9 -10.9 - 9.1	5.7	4.3	-24.6	
Mean‡ S.D.	7.0 0.4	5.3 0.1		5.9	5.4		5.5 0.1	4.9 0.1		5.7	4.3		
	t!	P< 0.01					1	0.01		•			1

*Control for 15 minutes preceding challenge.

(a) See Figure.

(b) Pulmonary Flow Resistance, Cm H₂0/1/sec.(c) Lung Compliance, m1/cm H₂0.

TABLE 27 Cat No. 1801 RESPONSE TO VARIOUS STIMULI EXPRESSED AS PERCENT CHANGE RELATIVE TO CONTROL VALUES*

	Pu	Imonary Flow Resistance	(b)	
Mech. % Control Stim. Change	High % Control SO ₂ Change	Mech. % Control Stim. Change	火 Control NaCl Change	Mech. % Control Stim. Change
23.3 33.7 43.7 23.6	18.1 18.1 -16.8 23.6 25.3 16.2 23.6 22.0 1.1	23.6 45 90.7	22.0 45.8 93.8 23.6 56.4 133.6 25.3 61.4 159.8	46.6 89.6 92.3
Mean [±] 23.5 33.7 S.O. 0.2	21.8 21.8 3.2 3.6	23.6 45.0	23.6 54.5 1.7 8.0	46.6 89.6
REFLEX INTACT	· Δ N.S.	REFLEX INTACT	P<0.05	REFLEX INTACT
		Lung Compliance (c)		
Mech. Control Stim. Change	High ½ Control SO2 Change	Mech. % Control Stim.Change	% Control NaCl Change	Mech. % Control Stim. Change
6.3 4.9 -21.6 6.2	5.2 6.0 12.5 5.4 5.7 6.9 5.4 5.4 1.3	6.0 5.7 -5.0	5.7 4.7 -15.6 5.6 4.6 -17.4 5.4 4.5 -19.2	4.8 4.8 0
Mean [±] 6.2 4.9 S.D. 0.1	5.3 5.7 0.1 0.3	6.0 5.7	5.6 4.6 0.2 0.1	4.8 4.8
	Δ N.S.		P<0.01	
Sequence of (a) (1)	(2)	(3)	(4)	(5)

^{*}Control for 15 minutes preceding challenge.

A N.S. = Difference between means not sigificant (P>0.05)

⁽a) See Figure.
(b) Pulmonary Flow Resistance, Cm H₂0/1/sec.
(c) Lung Compliance, ml/cm H₂0.

Cat No. 1801 RESPONSE TO VARIOUS STIMULI EXPRESSED AS PERCENT CHANGE RELATIVE TO CONTROL VALUES*

TABLE 27 (Continued)

	Pu	lmonary Flow Resistance	(6)	
SO ₂ + ½ Contro! NaCi Change	Mech. ½ Control Stim. Change	% Control SO ₂ Change	Mech. % Control Stim. Change	
23.6 25.3 -2.7 23.6 23.9 -8.1 30.8 21.0 -19.2	25.3 57.0 125.3	22.0 25.3 15.0 22.0 28.6 30.0 22.0 32.6 48.2	21.4 64.5 201.4	
Mean± 26.0 23.4 S.D. 4.2 2.2	25.3 57.0	22.0 23.8 0.1 3.7	21.4 64.5	
Δ N.S.	REFLEX INTACT	· Δ N.S.	REFLEX INTACT	
		Lung Compliance (c)		
SO ₂ +	Mech. ½ Control Stim. Chang⊖	% Control SO ₂ Change	Mech. % Control Stim. Change	
5.9 4.3 -20.4 5.6 4.3 -20.4 4.7 4.3 -20.4	5.1 5.2 2.0	5.0 4.2 -12.5 4.7 4.0 -16.7 4.7 4.2 -12.5	4.9 4.9 0	
Mean [±] 5.4 4.3 0.6 0	5.1 5.2	4.9 4.1 0.2 0.1	4.9 4.9	
Δ N.S.		P<0.01		
Sequence of Challenge (6)	(7)	(8)	(9)	

- (a) See Figure.
- (b) Pulmonary Flow Resistance, Cm H₂0/1/sec.
 (c) Lung Compliance, m1/cm H₂0.

^{*}Control for 15 minutes preceding challenge.

TABLE 28 Cat No. 1781 RESPONSE TO VARIOUS STIMULI EXPRESSED AS PERCENT CHANGE RELATIVE TO CONTROL VALUES*

•				-	Pu	lmonary Flow	Res	istance	(b)			
Control	Mech. Stim.	% Change		iigh 502	关 Change	Me Control St	ech.					
26.1 31.9	80.6	177.9	16.6	9.6 3.7 3.0	-42.1 12.9 3.7	16.1 4	4.4	175.8				
Mean± 29.0 S.D. 4.1	80.6		16.6	15.4 5.1		16.1 4	4.4			•		• .
t-test REFLEX	X INTAC	Γ	Δ	N.S.		REFLEX	INT	АСТ			_	
						Lung Compi	ianc	e (c)		-		
Control	Mech.	% Change	Control S	High SO ₂	浅 Change	Control S		% Change				
6.7 7.4	6.0	-14.9	7.3	6.6 6.6 6.7	-8.8 -8.8 -7.4	6.7 5	.6	-16.4				
Mean± 7.1 0.5	6.0			6.6		6.7 5	.6			٠		
t-test			₽<	0,01					· · · · · · · · · · · · · · · · · · ·	· .	 	·
Sequence of Challenge	f _a }	(1)		(2)		()	3)					

^{*}Control for 15 minutes preceding challenge.

 ⁽a) See Figure.
 (b) Pulmonary Flow Resistance, Cm H₂0/1/sec.
 (c) Lung Compliance, ml/cm H₂0.

TABLE 29 Cat No. 2206 RESPONSE TO VARIOUS STIMULI EXPRESSED AS PERCENT CHANGE RELATIVE TO CONTROL VALUES*

					Pu	Imonary	Flow Res	istance	(b)	
Control	S02 Both Sites	Percent Change	Control	50 ₂ Mouth	Percent Change	Control	S0 ₂ Trachea	Percent Change	Control	
44.6 44.6 44.6	37.3 42.2 42.2	-16.4 -5.4 -5.4	44.6 44.6 44.6	34.9 37.3 34.1	-21.7 -16.4 -23.5	44.6 44.6 44.6	32.5 27.3 30.0	-27.1 -38.8 -32.7	42.2	
Mean± 44.6 S.D. 0.0	40.6 2.8		44.6 0. 0	35.4 1.7	!	44.6 0.0	29.9 2.6	·	42.2	
t-test	ΔN.S.		F	< 0.05		P	< 0.05			
						Lung C	omplianc	e (c)		
Control	SU2 Both Sites	Perc e nt Chang e	Control	SO ₂ Mouth	Percent Change	Control	SO ₂ Trachea	Percent Change	Control	
5.3 5.6 5.6	4.6 4.5 5.0	-16.4 -18.2 - 9.1	5.2 5.5 5.3	5.8 5.3 5.6	8.8 -0.6 5:0	5.6 5.6 5.6	5.0 4.8 5.0	-10.7 -14.3 -10.7	5.0	
Mean [±] 5.5 S.D. 0.2	4.7 0.3		5.3 0.2	5.6 0.3		5.6 0.0	5.0 0.1		5.0	·
t-test	P<0.01			۵N.S.	•	[P< -0.01			
Sequence of Challenge			 							

(a) See Figure.

⁽b) Pulmonary Flow Resistance, Cm H₂0/1/sec. (c) Lung Compliance, m1/cm H₂0.

TABLE 30 Cat No. 53869 RESPONSE TO VARIOUS STIMULI EXPRESSED AS PERCENT CHANGE RELATIVE TO CONTROL VALUES

S02 Both Perce Sites Change 13.5 44.6 14.2 52.1 13.5 44.6 13.7 0.4	9.6 13 9.6 13 9.6 12 9.6 12 0.0	02 Percent outh Change 3.6 41.7 1.8 22.9 2.0 25.0 2.5 1.0	Control Trachea 10.9 13.5 10.9 14.2 10.9 12.9 10.9 13.5 0.0 0.7 P< 0.05	Change Con 23.9 30.3 18.3		
14.2 52.1 13.5 44.6 13.7 0.4	9.6 12 9.6 12 9.6 12	1.8 22.9 2.0 25.0 2.5 1.0	10.9 14.2 10.9 12.9 10.9 13.5 0.0 0.7 P< 0.05	30.3 18.3		
0.4	0.0	1.0	0.0 0.7 P< 0.05	12.	.3	
< 0.01	P< (0.05		- 1		}
						- {
	•		Lung Compliance	(ċ)		
SO ₂ Both Perce Sites Chang	nt SC Control Mo	0 ₂ Percent outh Change	SO ₂ Pe Control Trachea Cl	ercent hange Cont	rol	
19.4 -14.5 19.4 -14.5 20.2 -11.0	22.7	8.1 -20.3 9.4 -14.5 8.7 -17.6	19.4 16.4 -	19.2 18. 16.6 19.2	7	
19.7 0.5			19.7 16.1 0.5 0.3	18.	7	
0.01	P<	ე. 01	P<0.01			· .
19	9.7 9.5	0.2 -11.0 22.7 1 0.7 22.7 1 0.5 0.0	22.7 18.7 -17.6 22.7 18.7 22.7 0.0 0.7	0.2 -11.0 22.7 18.7 -17.6 20.2 15.9 - 0.7 22.7 18.7 19.7 16.1 0.5 0.5 0.3	0.2 -11.0 22.7 18.7 -17.6 20.2 15.9 -19.2 0.7 22.7 18.7 19.7 16.1 18. 0.5 0.0 0.7 0.5 0.3	0.2 -11.0 22.7 18.7 -17.6 20.2 15.9 -19.2 0.7 22.7 18.7 19.7 16.1 18.7 0.5 0.0 0.7 0.5 0.3

*Control for 15 minutes preceding challenge.
(a) See Figure.
(b) Pulmonary Flow Resistance, Cm H₂0/1/sec.
(c) Lung Compliance, ml/cm H₂0.

TABLE 31 Cat No. 2984 RESPONSE TO VARIOUS STIMULI EXPRESSED AS PERCENT CHANGE RELATIVE TO CONTROL VALUES*

SO ₂ Control Trache	Percent ea Change	Control	SO ₂ Mouth	Percent Change	Control	SU2 Both Sites	Percent Change	Control	
42.2 25.7 42.2 23.0 42.2 23.0	-39.1 -45.5 -45.5	42.2 42.2 42.2	39.4 42.2 42.2	-6.6 0 0	42.2 42.2 42.2	23.0 21.9 25.3	-45.5 -48.1 -40.0	36.7	
Mean [±] 42.2 23.9 S.D. 0.0 1.6		42.2 0.0	41.3		42.2 0.0	23.4		36.7	
t-test P< 0.01			۵ N. S.		}	P< 0.0	1		
					Lung C	omplian	ce (c)		
S0 ₂ Control Trach	Percent ea Change	Control	SO ₂ Mouth	Percent Change	Control	SO ₂ Both Sites	Percent Change	Control	
5.3 5.6 5.4 5.6 6.0 5.6	0.6 0.6 0.6	5.3 6.4 5.8	6.7 6.9 6.7	14.9 1 8.3 14.9	6.0 6.4 6.7	5.8 5.6 5.8	-8.9 -12.0 -8.9	8.2	
Mean ± 5.6 5.6 0.4 0.0		5.8 0.6	6.8 0.1		6.4 0.4	5.7 0.1		8.2	
t-test △ N.S.			∆N.S.		l l	∆ N.S.			

^{*}Control for 15 minutes preceding challenge.

⁽a) See Figure.

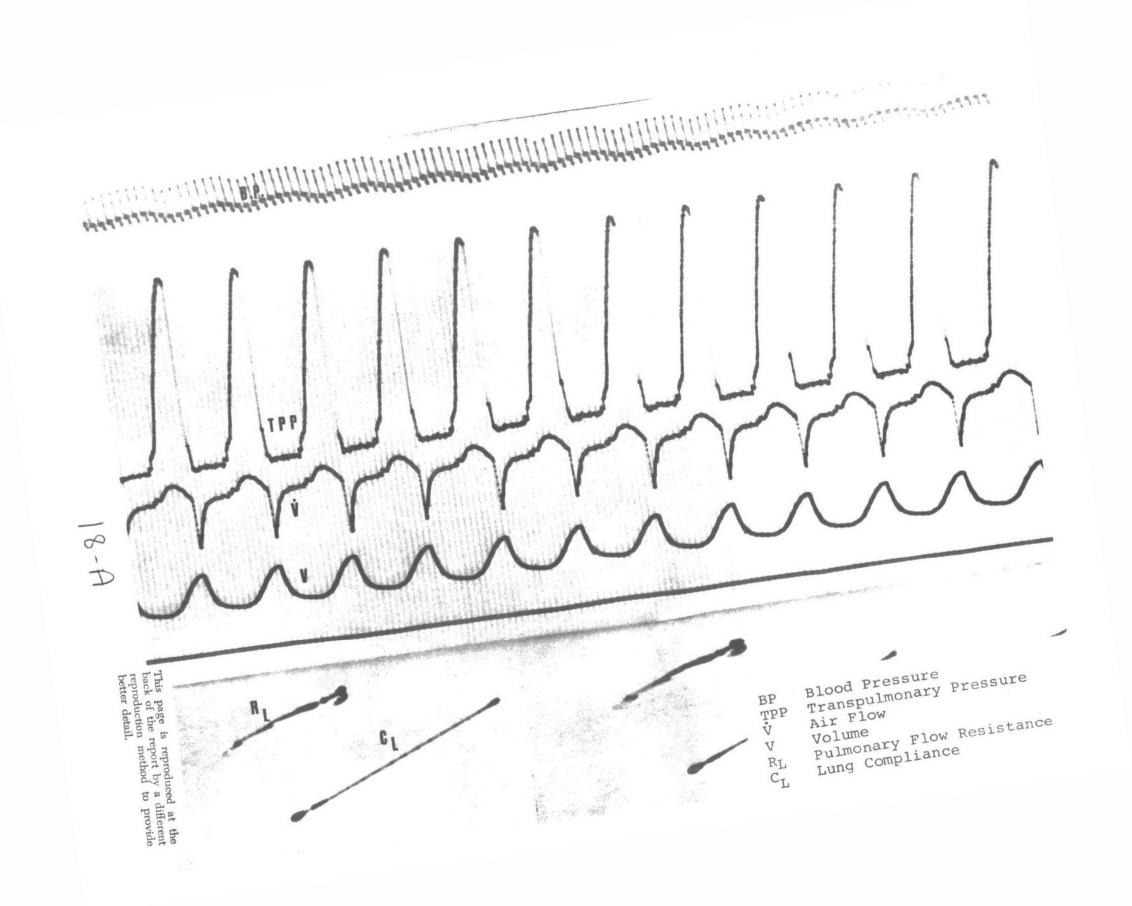
⁽b) Pulmonary Flow Resistance, Cm H₂0/1/sec. (c) Lung Compliance, m1/cm H₂0.

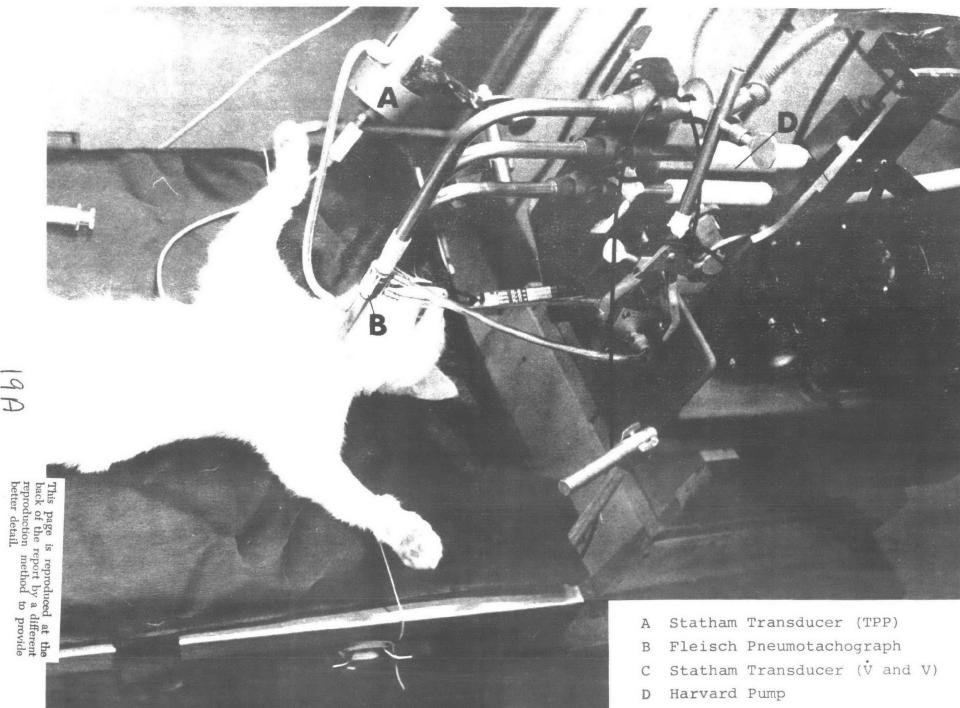
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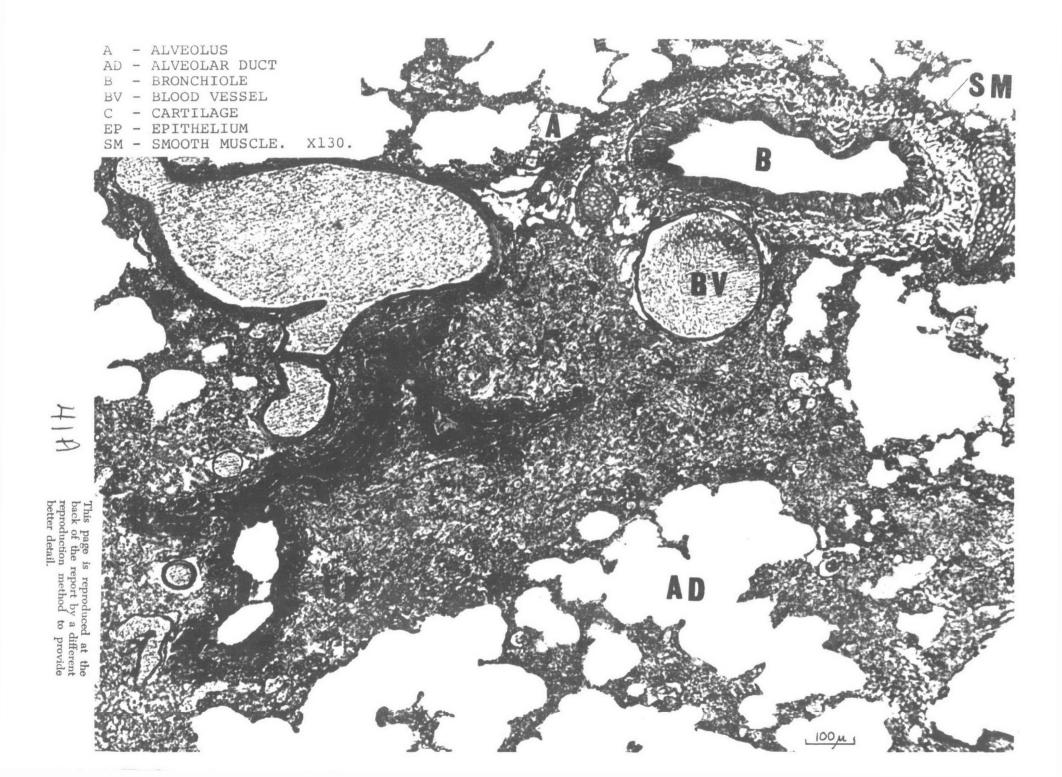
REPORT. THEY HAVE BEEN REPRODUCED HERE BY

A DIFFERENT METHOD TO PROVIDE BETTER DETAIL.





19A



Patho-Physiologic Response to Single and Multiple Air Pollutants in Humans and Animals