Paradoxical effect of salbutamol in a model of acute organophosphates intoxication in guinea pigs: role of substance P release

Jaime Chávez,1 Patricia Segura,1 Mario H. Vargas,1* José Luis Arreola,1 Edgar Flores-Soto,2 and Luis M. Montaño2*

1Departamento de Investigación en Hiperreactividad Bronquial, Instituto Nacional de Enfermedades Respiratorias, and 2Departamento de Farmacología, Facultad de Medicina, Universidad Nacional Autónoma de México, México DF, México

Submitted 9 June 2005; accepted in final form 5 December 2006

Chávez J, Segura P, Vargas MH, Arreola JL, Flores-Soto E, Montaño LM. Paradoxical effect of salbutamol in a model of acute organophosphates intoxication in guinea pigs: role of substance P release. Am J Physiol Lung Cell Mol Physiol 292: L915–L923, 2007. First published December 8, 2006; doi:10.1152/ajplung.00253.2005.—Organophosphates induce bronchoconstriction in guinea pigs, and salbutamol only transiently reverses this effect, suggesting that it triggers additional obstructive mechanisms. To further explore this phenomenon, in vivo (barometric plethysmography) and in vitro (organ baths, including ACh and substance P concentration measurement by HPLC and immunooassay, respectively; intracellular Ca2+ measurement in single myocytes) experiments were performed. In vivo experiments, parathion caused a progressive bronchoconstriction until a plateau was reached. Administration of salbutamol during this plateau decreased bronchoconstriction up to 22% in the first 5 min, but thereafter airway obstruction rose again as to reach the same intensity as before salbutamol. Aminophylline caused a sustained decrement (71%) of the parathion-induced bronchoconstriction. In in vitro studies, paraoxon produced a sustained contraction of tracheal rings, which was fully blocked by atropine but not by TTX, ω-conotoxin (CTX), or epithelium removal. During the paraoxon-induced contraction, salbutamol caused a temporary relaxation of ~50%, followed by a partial reaction. This paradoxical reaction was avoided by the M2- or neurokinin-1 (NK1)-receptor antagonists (methoctramine or AF-DX 116, and L-732138, respectively), accompanied by a long-lasting relaxation. Forskolin caused full relaxation of the paraoxon response. Substance P and, to a lesser extent, ACh released from intracellular Ca2+ basal levels. Our results suggested that: 1) organophosphates caused smooth muscle contraction by accumulation of ACh released through a TTX- and CTX-resistant mechanism; 2) during such contraction, salbutamol relaxation is functionally antagonized by the stimulation of M2 receptors; and 3) after this transient salbutamol-induced relaxation, a paradoxical contraction ensues due to the subsequent release of substance P.

albuterol; β2-adrenoceptor agonist; parathion; paraoxon; tachykinins; physostigmine; airway smooth muscle

PARATHION IS ONE OF THE MAIN REPRESENTATIVES OF ORGANOPHOSPHATES, A FAMILY OF COMPOUNDS SYNTHESIZED SINCE THE 1940s AND STILL WIDELY USED ALL AROUND THE WORLD IN AGRICULTURE AND VETERINARY MEDICINE AS INSECTICIDES AND ANTIHELMENTICS (12). TO ACHIEVE BIOLOGICAL ACTIVITY, PARATHION MUST BE BIOTRANSFORMED IN THE LIVER AND OTHER TISSUES INTO PARAOXON. THE LATTER METABOLITE HAS A VERY STRONG INHIBITORY CAPACITY ON ACETYLCHOLINESTERASE (AChE) ACTIVITY (27, 36), LEADING TO AN INCREASE IN ACh EFFECTS. THE USE OF ORGANOPHOSPHATES AND CARbamates ARE RESPONSIBLE FOR OVER 1,000,000 POISONING VICTIMS PER YEAR WORLDWIDE (17). THIS FIGURE, HOWEVER, MIGHT WELL UNDERESTIMATE THE PROBLEM DUE TO THE AMOUNT OF UNREPORTED CASES, MAINLY IN DEVELOPING COUNTRIES (37).

IN A PREVIOUS STUDY USING AN IN VIVO MODEL OF ORGANOPHOSPHATE POISONING, WE (35) FOUND THAT GUINEA PIGS WITH AN ONGOING PARATHION-INDUCED BRONCHOOBSTRUCTION RESPONDED TO SALBUTAMOL ADMINISTRATION WITH A TRANSIENT BRONCHODILATION, WHICH WAS FOLLOWED BY A RETURNING OF THE BRONCHOOBSTRUCTION. THOSE RESULTS ALLOWED US TO SPECULATE THAT SALBUTAMOL TRIGGERED SOME OBSTRUCTIVE MECHANISM THAT RAPIDLY COUNTERACTED ITS RELAXING ACTION. IN THIS REGARD, SOME STUDIES HAVE DEMONSTRATED THAT SALBUTAMOL CAN FACILITATE NEUROTRANSMITTER RELEASE, SUCH AS ACh RELEASE FROM PARASYMPATHETIC NERVES, THROUGH PREJUNCTIONAL β2-ADRENOCEPTOR STIMULATION (11, 44–46). BECAUSE SUCH AN EFFECT MIGHT WELL EXPLAIN THE AIRWAY OBSTRUCTION INDUCED BY SALBUTAMOL, IN THE PRESENT WORK THIS HYPOTHESIS AND SOME OTHER POSSIBLE MECHANISMS EXPLAINING THE PARADOXICAL EFFECT OF SALBUTAMOL WERE EXPLORED.

MATERIALS AND METHODS

Animals and experimental design. Male Hartley guinea pigs (500–600 g) bred in conventional conditions in our institutional animal facilities (filtered conditioned air, 21 ± 1°C, 50–70% humidity, sterilized bed) and fed with Harlan pellets and sterilized water were used. The protocol was approved by the Scientific and Bioethics Committees of the Instituto Nacional de Enfermedades Respiratorias. The experiments were conducted in accordance with the published Guiding Principles in the Care and Use of Animals approved by the American Physiological Society.

Most experiments were performed using either barometric plethysmography for freely moving guinea pigs or the organ bath technique for tracheal tissue. The sequence of drugs administered in these experiments is shown in Table 1. In addition, intracellular Ca2+ concentration was measured in enzymatically dispersed tracheal myocytes.

In vivo experiments. Guinea pigs were studied in a whole body, barometric plethysmography chamber for freely moving animals (Buxco, Troy, NY), and responses were recorded by a specialized software (BiosystemXVA 1.5.4, Buxco). The underlying principles of this technique have been already described by others (13, 15, 21). The main variable in this method is the “enhanced pause” (Penh), which has been shown to be correlated with total lung resistance in BALB/c...

* M. H. Vargas and L. M. Montaño contributed equally to this work.

Address for reprint requests and other correspondence: L. M. Montaño, Instituto Nacional de Enfermedades Respiratorias, Talpan 4502, CP 14080, México DF, México (e-mail: lmmr@servidor.unam.mx).
Table 1. Experimental design

<table>
<thead>
<tr>
<th>Plhysmography Experiments</th>
<th>Organ Bath Experiments</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sequence</td>
<td>n</td>
</tr>
<tr>
<td>Pth</td>
<td>7</td>
</tr>
<tr>
<td>Pth-Sal</td>
<td>7</td>
</tr>
<tr>
<td>Pth-Ami</td>
<td>7</td>
</tr>
<tr>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
</tr>
</tbody>
</table>

AF, AF-DX 116; Ami, aminophylline; Atr, atropine; CCh, carbachol; For, forskolin; L-732, L-732138; Phy, physostigmine; Pox, paraoxon; Pro, propranolol; Pth, parathion; Sal, salbutamol; TTX, tetrodotoxin; ω-CTX, ω-conotoxin GVIA; Met, methoctramine. *Separate experiments without epithelium (n = 5 for each group) were also performed.

mice (2, 21) and with specific airway resistance in guinea pigs (4, 9). The software was adjusted to only include breaths with a tidal volume of 1 ml or more, with a minimal inspiratory time of 0.15 s, a maximal inspiratory time of 3 s, and a maximal difference between inspiratory and expiratory volumes of 10%. A constant air flow (10 ml/s) was continuously supplied to the plethysmographic chamber without affecting the signal.

After each guinea pig was put inside the plethysmographic chamber, recording was initiated 5 min later, and a basal measurement of Penh was done. Approximately 20 min later, parathion (10 mg/kg) or vehicle (propylenglycol-ethanol, 10:1 vol/vol) was administered by the intraperitoneal route, and from this point onward respiratory parameters were recorded at minutes 5, 10, and every 10 min thereafter. The parathion dose was selected based on previous (35) and new experiments, to obtain the desired effect on airways without significant neurological effects. Because respiratory parameters were calculated by the computer in each breath, adjustments were made to the software to average values from all breaths occurring within 15 s and then to average those values during the last 5 min of each period. To evaluate whether a contraction of the airway smooth muscle was a component of the pulmonary toxicity of parathion, the bronchodilator effect of salbutamol (10 μg/kg ip) or aminophylline (40 mg/kg ip) was assessed during the plateau of the parathion-induced airway obstruction, while control animals received intraperitoneal saline solution. These salbutamol and aminophylline doses corresponded to the therapeutic dosage used in humans, and both of them were applied 90 min after parathion administration.

In vitro experiments: organ baths. Guinea pigs were euthanized by a pentobarbital overdose, and four rings (submitted to different experimental conditions) were obtained from the middle of the trachea and studied in a 5-ml organ bath system, as previously described (7). Tissues were stimulated three times with KCl (60 mM), and then paraoxon (10 μM) or physostigmine (10 μM) was added to the organ bath. Contractile responses to these drugs were expressed as percentage of the third KCl response. We corroborated that, at these concentrations, both paraoxon and physostigmine produced comparable contractile responses (91.9 ± 9.4% vs. 89.3 ± 3.3% of KCl contraction, respectively; P = 0.83). In some experiments, tissues were preincubated with 10 μM tetrodotoxin (TTX), 0.32 μM ω-conotoxin GVIA (ω-CTX; an N-type Ca2+ channel blocker), or 1 μM atropine during the 15 min before addition of paraoxon. In a separate set of tissues, during the plateau of the paraoxon- or physostigmine-induced maximum contraction, salbutamol (0.1 μM) or forskolin (1 μM; an adenylyl cyclase activator) were added with the aim to evaluate their relaxing effect.

To explore the role of β2- and M3-receptors in the effect of salbutamol, the former were blocked with propranolol (10 μM) and the latter with methoctramine (0.31 μM) or AF-DX 116 (0.1 μM), added 15 min before salbutamol. We corroborated that these concentrations of propranolol, methoctramine, and AF-DX 116 do not modify the plateau of tracheal contraction induced by paraoxon. Likewise, a possible participation of substance P on the effect of salbutamol was evaluated by using L-732138 (10 nM), a neurokinin-1 (NK1) receptor antagonist (28), added to the organ baths 15 min before paraoxon. We corroborated that these concentrations of L-732138 do not modify the contractile response to paraoxon. With the aim to assess if the salbutamol effect was similar during a paraoxon-induced contraction as during a typical cholinergic contractile response, additional tissues were precontracted with 0.32 μM carbachol against paraoxon. Likewise, paraoxon was added 15 min before salbutamol. We corroborated that these concentrations of paraoxon-induced contraction as during a typical cholinergic contractile response, additional tissues were precontracted with 0.32 μM carbachol against paraoxon. Likewise, paraoxon was added 15 min before salbutamol. We corroborated that these concentrations of carbachol and paraoxon did not modify the contractile response to paraoxon. Likewise, a possible participation of substance P on the effect of salbutamol was evaluated by using L-732138 (10 nM), a neurokinin-1 (NK1) receptor antagonist (28), added to the organ baths 15 min before paraoxon. We corroborated that these concentrations of L-732138 do not modify the contractile response to paraoxon. With the aim to assess if the salbutamol effect was similar during a paraoxon-induced contraction as during a typical cholinergic contractile response, additional tissues were precontracted with 0.32 μM carbachol against paraoxon. Likewise, paraoxon was added 15 min before salbutamol. We corroborated that these concentrations of carbachol and paraoxon did not modify the contractile response to paraoxon.

To evaluate the role of airway epithelium in the paraoxon-induced responses, guinea pig tracheal smooth muscle strips (in which epithelial layer and connective tissue surrounding smooth muscle were removed by dissection under microscopic microscopy) were used instead of tracheal rings. These smooth muscle strips were submitted to 10 μM paraoxon stimulation with or without 10 μM TTX preincubation.

In experiments aimed to obtain samples of bath fluid for substance P measurement, tracheal rings were incubated with paraoxon (10 μM) for 1 h with or without addition of salbutamol (100 nM) in the second half of this period. Afterwards, the bath liquid was recovered and stored in vials at −70°C until its study. These experiments were performed with 3.2 μM phosphoramidon to avoid the enzymatic degradation of substance P by neutral endopeptidases. The same procedure was used to obtain samples for ACh measurement, excepting that physostigmine (10 μM) was used instead of paraoxon (because the organophosphate spoils the analytic column for ACh and choline measurement used in the HPLC system), and three concentrations of salbutamol were separately tested (1, 10, or 100 μM). In addition, these last samples were filtered through 0.2-μm nylon membrane Acrodics (Pall Gelman Sciences, Ann Arbor, MI) before its storage at −70°C.

AChE activity measurement. The AChE activity was determined in guinea pig lung lobe and plasma samples using a colorimetric method based on the Ellman reaction (14) and expressed as percentage of inhibition. Briefly, the lung tissue was homogenized in phosphate buffer (1 ml of phosphate buffer per 100 mg of tissue) with an homogenizer (Polytron Kinematica PT3100; Brinkmann, Westbury, NY). The homogenate was centrifuged for 15 min at 3,200 g. The supernatant was filtered (Whatman 1 filter paper), and 300 μl of supernatant was added to a cuvette containing 2.5 ml of 5,5’-dithiobis(2-nitrobenzoic acid) (DTNB; 0.32 mM) and 300 μl of phosphate buffer (64 mM). The background absorbance per minute was measured at 405 nm at 25°C with a spectrophotometer (DU-640; Beckman, Fullerton, CA). Afterwards, 100 μl of AChE substrate (42 mM acetylthiocholine) was added to the cuvette, and the change in absorbance per minute was measured. Once the background absorbance was subtracted, the AChE activity was calculated as international units (IU) by means of the following equation

\[
IU = \frac{A \times 3,200}{0.36 \times 10^4 \times 300 \times Co}
\]

where A is the change in absorbance per minute, 0.36 × 104 is the extinction coefficient of DTNB, Co is the amount of tissue in the
 supernatant (milligram of tissue per milligram of buffer), and 3,200 and 300 are the total volume (μL) of the cuvette and the volume (μL) of the supernatant, respectively.

Regarding plasma samples, after 1:25 dilution in phosphate buffer, 100 μL was added to a cuvette containing 2.5 mL of DTNB and 500 μL of phosphate buffer. The remaining steps were the same as those described for lung homogenates. The formula was changed (100 μL of sample instead of 300 μL), and IU corresponded to substrate hydrolyzed mol·min⁻¹·ml⁻¹ of plasma.

\[ \text{ACh measurement by HPLC.} \]
ACh and choline concentrations in the physostigmine-containing samples were measured by cation exchange HPLC electrochemical detection method as described by Potter et al. (30). In this technique, an analytic column for ACh and choline (MF-6150; Bioanalytical Systems, West Lafayette, IN), an immobilized enzyme reactor (Bioanalytical Systems), and an electrochemical detector (Coulochem II; ESA, Chelmsford, MA) were coupled to the HPLC (model 9012; Varian, Walnut Creek, CA). Mobile phase (50 mM Tris/NaClO₄ plus 1% ProClin reagent, pH 8.5) was pumped at a rate of 1 mL/min. Standard curves of ACh and choline (1–100 mM) were used for calibration. The detection limit of our HPLC system was ~0.1 nA in a 15-μL sample. Data were stored in a microcomputer and analyzed using a data acquisition and analysis software (Star Chromatography Workstation v4.01, Varian). ACh production in the samples was expressed as the sum of ACh plus choline detected by the HPLC system and expressed as nanomoles per liter.

\[ \text{Substance P measurement.} \]
Substance P concentration was measured through a competitive enzyme immunoassay kit (Cayman, Ann Arbor, MI). Samples were read at 405 nm using a Multiskan MS photometer (Labsystems Oy, Helsinki, Finland). Substance P concentration was expressed as the sum of ACh plus choline detected by the HPLC system and expressed as nanomoles per liter.

\[ \text{Ca}^{2+} \text{ measurement in guinea pig tracheal myocytes.} \]
Airway smooth muscle from guinea pig trachea was placed in 5 mL of Hank's solution containing 2 mg of cysteine and 0.05 U/ml papain and incubated for 10 min at 37°C. The tissue was washed with Leibovitz's solution to remove the enzyme excess and then placed in a Krebs solution containing 2 mg of cysteine and 0.05 U/ml papain and incubated for 10 min at 37°C. The tissue was gently dispersed by mechanical agitation until detached cells were observed. Enzymatic activity was stopped by adding Leibovitz's solution, the cells were centrifuged at 600 rpm for 5 min, and the supernatant was discarded. This last step was repeated twice again. Afterward, cells were loaded with fura-2/AM in low Ca²⁺-free Krebs solution containing 1 mg/ml collagenase type I and 4 mg/ml dispase II (neutral protease) for ~10 min at 37°C. The tissue was gently dispersed by mechanical agitation until detached cells were observed. Enzymatic activity was stopped by adding Leibovitz's solution, the cells were centrifuged at 600 rpm for 5 min, and the supernatant was discarded. This last step was repeated twice again. Afterward, cells were loaded with 0.5 μM fura-2/AM in low Ca²⁺ (0.1 mM) at room temperature (22–25°C). After 1 h, cells were allowed to settle down into a heated perfusion chamber with a glass cover in the bottom. This chamber was mounted on an inverted microscope (Diaphot 200; Nikon, Tokyo, Japan), and the cells adhered to the glass were continuously perfused at a rate of 2–2.5 mL/min with Krebs solution containing 10 mM HEPES, 4 mM CaCl₂, 2.0 mM MgSO₄, 1.2 mM NaHCO₃, 25 mM KH₂PO₄, 1.2 mM glucose, 11 mM CaCl₂, 37°C, equilibrated with 5% CO₂ in oxygen, pH 7.4.

Addition of paraoxon to organ baths produced a sustained decrease in ACh response, which was expressed as median and range.

\[ \text{RESULTS} \]
In vitro studies. We corroborated that the parathion dose used in our in vivo experiments (10 mg/kg) caused (90 min later) a 97.8 ± 0.8% and 67.3 ± 4.9% inhibition of AChE activity in guinea pig lung homogenates and plasma, respectively (n = 5 for each group; Fig. 1A).

Administration of parathion to control male guinea pigs caused a progressive increase of the basal Penh values in up to 250%, from 0.213 ± 0.010 to 0.566 ± 0.080, n = 7, and remained unchanged until the end of the recording period (Fig. 2). Salbutamol (10 μg/kg ip) caused a 31% decrease (P < 0.05) of the ongoing parathion-induced bronchoconstriction 5 min after its intraperitoneal administration, compared with the averaged Penh values in the previous 10-min period (Fig. 3). After this transient relaxation, Penh was raised again as to reach the same values as before salbutamol administration. A higher salbutamol dose (30 μg/kg ip) also produced this paradoxical reaction, though somewhat delayed in time (data not shown). By contrast, aminophylline produced a progressive decrement (P < 0.05 to P < 0.001) of the parathion-induced bronchoconstriction that was maximal at 20 min (77% decrement) and remained relatively unchanged thereafter.

In vitro studies. In guinea pig lung homogenates, we confirmed that paraoxon and physostigmine caused (3 min later) a concentration-dependent inhibition of AChE activity (Fig. 1B). Specifically, at the concentration used in the present study (10 μM) either paraoxon or physostigmine produced 99.9 ± 0.1% inhibition of AChE activity (n = 4 for each group).

Addition of paraoxon to organ baths produced a sustained smooth muscle contraction of guinea pig tracheal rings (Fig. 4A). This response was not modified by previous incubation with 0.3-CTX or TTX (Fig. 4B), but was fully prevented and reverted...
by atropine (Fig. 4, C and D). In some tissues, we corroborated
that these ω-CTX and TTX concentrations were enough to
abolish or significantly diminish, respectively, the contractile
response induced by electrical field stimulation (data not
shown). Elimination of epithelium and connective tissue from
the airway preparation not only did not interfere with the
contractile effect of paraoxon, but it caused an increased
response to this organophosphate, compared with whole tra-
cheal rings (190.3 ± 39.3% vs. 102.1 ± 11.4% of KCl
contraction, respectively; P < 0.02). This last response was not
modified by TTX pretreatment (188.0 ± 41.7% of KCl con-
traction).

Salbutamol (100 nM) administered during the paraoxon-
induced contraction produced a temporary relaxation of
~50% (Figs. 5 and 6) 10–20 min after its administration. From this
point onwards, tissues began to contract again but without
reaching the initial maximal paraoxon-induced contraction. A
similar pattern was observed when tissues were precontracted
with carbachol instead of paraoxon (Fig. 6). We corroborated
that the relaxation induced by salbutamol was due to β-adre-
noceptor stimulation, since preincubation with propranolol
completely abolished this response (Fig. 5).

The M3 antagonist methoctramine did not modify the on-
going paraoxon-induced contraction (Fig. 4E), but fully pre-
vented the paradoxical contraction occurring after the transient
salbutamol-induced relaxation, favoring a sustained relaxing
response of ~90% (Fig. 5). In this same context, a lower
concentration of salbutamol (32 nM) induced a partial relax-
ation of the carbachol-induced contraction, and methoctramine
(administered during the plateau of such relaxing response)
was able to completely relax the tissue (Fig. 4F). Likewise,
AF-DX 116 did not affect the paraoxon-induced contraction in
tracheal rings, but permitted a complete relaxing effect of
salbutamol (140.4 ± 27.1% relaxation, not shown). Addition-
ally, we found that tissue incubation with the NK1 receptor
antagonist L-732138 allowed salbutamol to fully relax the
tracheal rings and almost abolished the paradoxical recontrac-
tion (Fig. 7). By contrast, forskolin completely relaxed the
paraoxon response from 10 min ahead (Fig. 6). Some of these
phenomena were essentially reproduced when phystostigmine
was used instead of paraoxon (Fig. 8).

ACh and substance P measurements. As can be seen in Fig.
9, control levels of ACh released from tracheal rings during a
60-min incubation with phystostigmine had a tendency to in-
crease when 1, 10, or 100 nM salbutamol was administered in
the second half of this period, reaching statistically significant
difference at the intermediate concentration. On the other hand, addition of 100 nM salbutamol significantly increased the concentration of substance P recovered in the organ bath fluid (median 34.9 pg/ml, range 32.0–79.4), compared with control tissues without salbutamol (median <3.9 pg/ml, range <3.9–9.4; \( P < 0.0001 \); Fig. 10).

**Single cell studies.** Baseline concentration of \([Ca^{2+}]_i\) in airway myocytes was 49 ± 7 nM, \( n = 4 \). Carbachol addition produced a sustained smooth muscle contraction of the guinea pig tracheal rings (A). This response was not modified by previous incubation with TTX (B) but was completely blocked (C) and reverted (D) by atropine. Methoctramine did not modify the contraction induced by paraoxon but potentiated the relaxation induced by 0.1 μM salbutamol (Sal) (E) and removed the functional antagonism between carbachol (CCh) (0.32 μM) and salbutamol (0.032 μM) (F). Pox, paraoxon (10 μM); Atr, atropine (1 μM); KCl, potassium chloride (60 mM); Met, methoctramine (0.31 μM); TTX, tetrodotoxin (10 μM).

**Fig. 5.** Effect of \( M_2 \) receptor antagonism on the salbutamol response during the paraoxon-induced contraction in tracheal rings. Paraoxon (○, 10 μM, \( n = 11 \)) induced a contractile response, which was partially reversed by salbutamol (●, 0.1 μM, \( n = 11 \)) with a tendency to contract again from 10 min onward. The paradoxical contraction occurring after the transient salbutamol-induced relaxation was fully abolished by the \( M_2 \) antagonist methoctramine favoring a relaxing response of \( >90\% \) (●, 0.31 μM, \( n = 8 \)). Propranolol (▲, 1 μM, \( n = 6 \)) completely abolished the salbutamol relaxant effect. Symbols represent the means ± SE. *\( P < 0.05 \) and **\( P < 0.01 \) compared with the paraoxon group and †\( P < 0.01 \) compared with the paraoxon-salbutamol group (one-way ANOVA with Bonferroni multiple comparisons test).**

**Fig. 6.** Effect of salbutamol and forskolin on the paraoxon-induced contraction in tracheal rings. Paraoxon (○, 10 μM, \( n = 11 \)) induced a contractile response, which was partially reversed by salbutamol (●, 0.1 μM, \( n = 11 \)) with a tendency to contract again from 10 min onward. Forskolin (▲, 1 μM, \( n = 7 \)) completely abolished the paraoxon-induced contraction. The sustained contraction induced by carbachol (●, 0.32 μM, \( n = 8 \)) was moderately reversed by salbutamol following a similar pattern as was observed in the paraoxon group (●, 0.1 μM, \( n = 11 \)). Symbols represent the means ± SE. *\( P < 0.01 \) compared with their respective control group; †\( P < 0.01 \) compared with the paraoxon-salbutamol group (one-way ANOVA with Bonferroni multiple comparisons test).
caused a transient Ca\(^{2+}\) peak followed by a plateau (Fig. 11). Paraoxon did not modify the [Ca\(^{2+}\)] or the response to a superimposed stimulation with carbachol.

**DISCUSSION**

In the present study, we found that, under conditions of AChE inhibition, salbutamol induced the release of substance P, and to a lesser extent ACh, with a subsequent contraction large enough to overcome its relaxing effect. Parathion is an organophosphate pesticide with a well known and potent inhibitory effect on AChE activity (8, 35) mediated through its active metabolite, paraoxon. One toxicological consequence of the resulting ACh accumulation is airway obstruction, which in in vivo conditions has been postulated to be mediated by a combination of airway smooth muscle contraction, augmented mucous secretion, and edema (35). In the present work, we found that the main physiopathological mechanism involved in the organophosphate-induced airway obstruction was the smooth muscle contraction, since in in vivo conditions aminophylline rapidly reverted much of this obstruction (71%; Fig. 3), while forskolin caused a rapid and complete relaxation of the paraoxon-induced contraction in vitro (Fig. 6). The cholinergic nature of this response was confirmed by the total abolishment of the contraction by atropine in the in vitro preparation. This response, however, was TTX- and \(\alpha\)-CTX-resistant, suggesting that spontaneous ACh release involves mechanisms other than those classically occurring during neuronal depolarization. Such additional mechanisms have been already postulated by others (1). Because there is good evidence that minute amounts of ACh may also be produced by nonneuronal sources (27, 29), an alternative possibility is that ACh was not coming from nerves but from other airway cell types. In separate experiments, we ruled out a major role of airway epithelium as additional source of ACh in the organophosphate-induced contraction, but further experiments are needed to discard other sources such as airway smooth muscle (41).

Contrasting with forskolin and aminophylline, salbutamol was unable to induce a sustained relaxation (in vitro) or bronchodilation (in vivo) and, furthermore, its moderate and...
transient relaxing effect was rapidly overlapped by an additional contraction. At least in the in vivo experiments, the dose used in our study (10 µg/kg ip) was close to the intravenous dose recommended for the acute relief of severe asthmatic bronchospasm in children (5). Additionally, in a previous work (35), we found that aerosolized salbutamol at therapeutic doses used in humans (2 mg/ml, 2 min) also produced the paradoxical effect. Thus the paradoxical effect of salbutamol in guinea pigs occurred at doses equivalent to those used in humans.

In addition to their main pharmacological effect on airways (relaxation of smooth muscle), several studies have demonstrated that salbutamol and other β2-adrenoceptor agonists are capable of stimulating vagal nerve ends to induce ACh release (11, 44–46). This might explain the increment of ACh concentration observed in the organ bath fluids after salbutamol administration. Nevertheless, this increment was small enough as to raise doubts about its potential role in the paradoxical recontraction induced by salbutamol.

Contrasting with the ACh results, salbutamol was able to induce a significant increase of substance P concentration. This last tachykinin may explain the secondary contraction observed during the salbutamol response, which is in agreement with the almost full relaxation occurring during the antagonism of NK1 receptors. It is well known that an extensive network of tachykinin-containing nerve ends (C fibers) is located within and below the airway epithelium, including the trachea (38, 42). Thus one possible explanation of salbutamol-induced tachykinin release is that salbutamol directly stimulates C fibers. However, to our knowledge, the presence of prejunctional β2-adrenoceptors in C fibers has not been described so far, and thus this possibility requires further demonstration. The possible role of C fibers is in agreement with a recent work by Keir et al. (25). They demonstrated that a 10-day treatment with β2-adrenoceptor agonist (RS- and S-albuterol) induced bronchial hyperresponsiveness (BHR) in sensitized and non-sensitized guinea pigs. The development of BHR was prevented by capsaicin treatment, suggesting a role of sensory nerves (C fibers) in this phenomenon.

In our in vitro studies, we found that the transitory character of the relaxation induced by salbutamol was similarly observed during the contraction induced either by paraoxon, physostigmine (a nonorganophosphate compound), or carbachol, suggesting that increased concentrations of ACh in the microenvironment surrounding the smooth muscle is necessary for the paradoxical effect of salbutamol to take place. Cholinergic contraction of smooth muscle involves activation of both M3 and M2 receptors. The latter is known to be coupled to Gi protein and hence to produce inhibition of adenylate cyclase (33). Thus mechanisms induced by M2 receptor activation are particularly opposed to those produced during β2-adrenoceptor stimulation. This functional antagonism between M2 and β2 receptors has been already reported (6, 16) and might be largely involved in our results, inasmuch as the blockade of M2 receptors by methoctramine or AF-DX 116 enhanced the salbutamol-induced relaxation and abolished the paradoxical effect. We were able to demonstrate the existence of this functional antagonism by using a slightly lower concentration of salbutamol (32 nM). In these experiments, salbutamol caused an ∼50% relaxation of the carbachol-induced contraction, and the subsequent blockade of M2 receptors by methoctramine produced a rapid and complete relaxation of the tissues (Fig. 4F). In summary, our results strongly suggest that at least two mechanisms are involved in the paradoxical effect of salbutamol during an acute organophosphate intoxication: 1) the functional antagonism between M2 and β2-receptors; and 2) the salbutamol-induced release of substance P.

However, additional mechanisms might also be involved. For example, van den Beukel et al. (39) studied the effect of parathion and paraoxon on Chinese hamster ovary cells transfected with muscarinic M3 receptors. They concluded that these organophosphates act as agonists on M3 receptors (which, in airway smooth muscle, mediate the ACh contractile effect). In this context, we did not find a direct effect of paraoxon in single airway smooth muscle cells. On the other hand, Fryer et al. (18) and Lein and Fryer (26) recently postulated that organophosphate insecticides induce BHR through a decrease in the neuronal M2 muscarinic receptor function. This prejunctional receptor limits the release of ACh from pulmonary vagal nerves (19) and thus its inhibition by organophosphates would favor the release of this neurotransmitter if a subsequent neural stimulation takes place. To what extent these mechanisms are participating in the paradoxical effect of salbutamol remains to be elucidated.

Exposure to organophosphate pesticides may worsen symptoms in asthmatic patients or induce respiratory complaints such as wheezing in nonasthmatic subjects (3, 22, 32, 43). In the case that an organophosphate poisoning is recognized, the appropriate treatment with oximes and atropine is likely to be started. Nevertheless, in many cases, organophosphate poisoning is low-grade in nature, and the exposure history to organophosphates would not always be evident (22, 37). In this last setting, it is expected that a β2-adrenoceptor agonist such as salbutamol is administered for the treatment of wheezing. In this context, our current results in these in vivo and in vitro models of organophosphate pesticide intoxication allow us to speculate that effectiveness of such treatment might be less than expected or may even worsen the bronchial obstruction. This last speculation warrants further investigation. Additionally, clinical studies in asthma have found that chronic treatment with β2-adrenoceptor agonists worsens asthma control (34) and induces BHR to a number of bronchoconstrictor stimuli, including methacholine, histamine, exercise, and allergen (10, 23, 34, 40). Therefore, one possible explanation of this
adverse effect of salbutamol could be related to its effect on tachykinin release.

According to our results, we concluded that: 1) organophosphates caused smooth muscle contraction by accumulation of ACh released through a TTX- and ω-CTX-resistant mechanism; 2) during such contraction, salbutamol relaxation is functionally antagonized by the stimulation of M₂ receptors; and 3) after this transient salbutamol-induced relaxation, a paradoxical contraction ensues due to substance P release. Thus further studies on the potential adverse effect of salbutamol during organophosphate-induced bronchoostriction in humans are warranted.

ACKNOWLEDGMENTS

We thank the Programa de Doctorado en Ciencias Biomédicas, Facultad de Medicina, Universidad Nacional Autónoma de México.

GRANTS

This study was partially supported by Dirección General de Apoyo al Personal Académico, Universidad Nacional Autónoma de México (DGAPA-UNAM) Grant IN203502.

REFERENCES


