Effects of sodium metabisulphite on guinea pig contractile airway smooth muscle responses in vitro

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Abstract

Background — Sodium metabisulphite (MBS) is known to induce bronchoconstriction in asthmatic patients. The effects of MBS on guinea pig airway smooth muscle and on neurally mediated contraction in vitro have been examined.

Methods — Tracheal and bronchial airway segments were placed in oxygenated buffer solution and electrical field stimulation was performed in the presence of indo-methacin (10^-3 M) and propranolol (10^-4 M) for the measurement of isometric tension. Atropine (10^-6 M) was added to bronchial tissues.

Results — Concentrations of MBS up to 10^-3 M had no direct effect on airway smooth muscle contraction and did not alter either tracheal smooth muscle contraction induced by electrical field stimulation at all frequencies or acetylcholine-induced tracheal smooth muscle contraction. There was a similar response in the absence of epithelium, except for potentiation of the response induced by electrical field stimulation at 0.5 Hz (24 (10)% increase). However, MBS (10^-5, 10^-4 and 10^-3 M) augmented neurally-mediated non-adrenergic non-cholinergic contractile responses in the bronchi (13-3 (3-2)%), 23-8 (9-6)% and 6-4 (1-6)% respectively). MBS had no effect on the contractile response induced by substance P, but at higher concentrations (10^-3 M and 10^-2 M) it caused a time-dependent attenuation of responses induced by either electrical field stimulation or exogenously applied acetylcholine or substance P.

Conclusions — MBS had no direct contractile responses but enhanced bronchoconstriction induced by activation of non-cholinergic neural pathways in the bronchus, probably through increased release of neuropeptides. At high concentrations MBS inhibited contractile responses initiated by receptor or neural stimulation.

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Keywords: sodium metabisulphite, cholinergic nerves, non-adrenergic non-cholinergic, sulphur dioxide, airway smooth muscle.

Sodium metabisulphite (MBS) is widely used as a food preservative and can induce bronchoconstriction in patients with asthma but not in normal individuals.1,4 MBS-induced bronchoconstriction is mediated through cholinergic pathways to a small extent,2,3 and it is possible that the non-cholinergic component of this response involves the release of tachykinins. Studies in the anaesthetised guinea pig have shown that, while a cholinergic reflex is partly involved in MBS-induced bronchoconstriction,1 tachykinins acting partially through both neurokinin NK₁ and NK₂ receptors may also be released.5 In addition, MBS induces plasma extravasation in the airways of guinea pigs, an effect which is also partly mediated through tachykinin release.6 It is not known, however, whether MBS could have direct or indirect effects on airway smooth muscle responses.

In the present study we have examined the responses of the guinea pig airway smooth muscle to MBS added directly to airway preparations in vitro under isometric conditions. In addition, we determined whether MBS could modulate airway smooth muscle responses induced by the release of acetylcholine and tachykinins from airway nerves. Because guinea pig tracheal preparations demonstrate a predominantly cholinergic response and bronchial preparations both cholinergic and non-cholinergic components following electrical field stimulation,7 we examined the effects of MBS on the contractile responses of both tracheal and bronchial preparations to electrical field stimulation.

Methods

Tissue Preparation

Male Dunkin-Hartley guinea pigs weighing 350–500 g were sacrificed by cervical dislocation. The airways (from larynx to whole lung) were rapidly removed and placed in oxygenated modified Krebs-Henseleit solution of the following composition (mM): NaCl 118, KCl 5-9, MgSO₄ 1-2, CaCl₂ 2-5, NaHCO₃ 25-5, and glucose 5-05. After careful dissection of connective tissue and lung parenchyma from the airway (from larynx to second generation bronchi), the trachea was opened longitudinally by cutting through the cartilage in its anterior aspect (opposite the smooth muscle layer) and then cut transversely with each segment containing 3–4 cartilaginous rings. In some experiments removal of the airway epithelial layer was achieved by gently rubbing the luminal surface to the trachea with a cotton wool applicator, which consistently achieves epithelial demodulation as assessed by light microscopy.8 For the bronchial preparations bronchi were separated into main and hilar sections respectively.
Tracheal and bronchial strips were suspended between platinum plate electrodes approximately 15 cm apart in 15 ml organ baths containing Krebs solution bubbled continuously with a mixture of 95% O₂ and 5% CO₂ at 37°C with pH at 7.4. Indomethacin (10⁻³ M) and propranolol (10⁻⁶ M) were present throughout the experiments, and atropine (10⁻⁶ M) was added for studies involving bronchial strips only. Atropine was not added to tracheal strips because it completely abolishes responses to electrical field stimulation. The segments were left to equilibrate for at least one hour with frequent washing and the resting tensions were adjusted to 1.5 g for the trachea and 0.5 g for the bronchi. These resting tensions were previously found to be optimal for measuring changes in isometric tensions. Isometric contractile responses were measured with Grass FT.03 force displacement transducers and recorded on a Graphic Linear recorder (Mark VII) polygraph (Graphtec Ltd, Nantwich, Cheshire, UK).

ELECTRICAL FIELD STIMULATION

Biphasic square wave pulses were delivered for 20 second periods from an electrical stimulator with a voltage of 20 V at source and pulse duration of 0.5 ms. Tracheal strips were first stimulated with the maximum frequency of 50 Hz four successive times, at intervals of three minutes each. A control frequency-response curve of the strips was performed after the resting tension had returned to a stable baseline. For each frequency-response curve, pulses of increasing frequency (from 0.5 to 50 Hz) were delivered every three minutes. Two successive reproducible control frequency-response curves were obtained before MBS or diluent was added to the bath. After incubating for 20 minutes, electrical field stimulation was repeated in the same tissue with increasing frequencies of stimulation.

For bronchial strips, stimuli at a frequency of 8 Hz were delivered every 30 minutes after the resting tension of the smooth muscle had returned to its baseline. A frequency-response curve of the bronchial preparation was not possible to perform because repeated stimulations at different frequencies gave inconsistent responses. However, repeated stimulations at a single frequency were reproducible and at least two successive reproducible control responses to 8 Hz were obtained before MBS or diluent was added to the bath. Electrical field stimulation was repeated after incubation for 20 minutes with the drugs. Only one concentration of MBS was tested for each tracheal or bronchial preparation (10⁻⁷ to 10⁻³ M).

ACETYLCHOLINE AND SUBSTANCE P RESPONSES

The effects of MBS (10⁻³ M and 10⁻⁶ M) on cumulative concentration responses to acetylcholine (10⁻⁶ to 10⁻³ M) were studied in tracheal strips, and to substance P (10⁻⁸ to 10⁻⁵ M) were studied in both tracheal (with epithelium intact or removed) and bronchial strips. The results were expressed as a percentage of the maximum contraction induced by the agonists themselves, but for substance P the results were expressed as a percentage of the maximum contraction induced by carbachol (10⁻⁷ M).

DRUGS AND SOLUTIONS

Drugs and chemicals were obtained from the following sources: atropine sulphate (Phoenix Pharmaceuticals Ltd, Gloucester, UK); propranolol hydrochloride (Zeneca plc, Macclesfield, UK); substance P, acetylcholine chloride, indomethacin, and MBS (Sigma Chemical Co, Poole, UK). Stock solutions of MBS were made up in distilled water 24 hours

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**Figure 1** Effect of sodium metabisulphite (MBS 10⁻⁴ M) or control on contractile responses of guinea pig tracheal smooth muscle strips induced by increasing frequencies of electrical field stimulation (0.5–50 Hz, 20 V, 0.5 ms, 20 s) in the presence of propranolol (10⁻⁴ M) and indomethacin (10⁻³ M). The epithelium has been left intact. Data are shown as mean (SE) of 5–6 guinea pigs.

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Figure 2 Enhancing effect of sodium metabisulphite (MBS 10⁻⁷ to 10⁻³ M) expressed as percentage increase above control responses on non-adrenergic, non-cholinergic contraction of guinea pig bronchial smooth muscle strips induced by electrical field stimulation at 8 Hz (20 V, 0.5 ms, 20 s) in the presence of propranolol (10⁻⁴ M), atropine (10⁻⁶ M), and indomethacin (10⁻³ M). Data are shown as mean (SE) of 6–8 guinea pigs. *p<0.05; **p<0.01 compared with vehicle-treated tissues.
before the experiment and then further diluted also in distilled water to the appropriate concentration on the experimental day. Fresh solutions of drugs were made up daily. The control solutions for MBS were made by adjusting the pH of distilled water to the same pH as MBS solution (10⁻³ M; pH 3-75) using sulphuric acid.

**DATA ANALYSIS**

Results are expressed as mean (SE). Electrical field stimulation frequency-response curves were first obtained in the absence of MBS and then repeated in the presence of one single concentration of MBS or diluent. A two-tailed paired Student's t test was used to determine the difference between sham-treated and MBS-treated groups. The effects of MBS were expressed as a percentage of pretreatment control response at each frequency of stimulation. For agonist-induced concentration response curves, either acetylcholine-induced or substance P-induced responses were performed in the presence of either MBS or diluent. For responses to acetylcholine the effective concentration needed to cause 50% of the maximal contractile response (EC₅₀) was calculated from concentration-response curves. EC₅₀ values are reported as geometric means and geometric standard error of the mean. The Mann-Whitney U test was also used to analyse the differences. p values of <0.05 were considered to be significant.

**Results**

MBS in concentrations ranging from 10⁻⁷ M to 10⁻³ M had no direct contractile effect on guinea pig trachea or bronchial preparations (n=5–8). Removal of the airway epithelium had no effect on the direct contractile effects.
EFFECT OF MBS ON RESPONSES TO ELECTRICAL FIELD STIMULATION

MBS (10^-7–10^-3 M) had no effect on electrical field stimulation responses at frequency stimulation of 0.5–50 Hz in tracheal strips, apart from an augmentation at 0.5 Hz. Figure 1 shows the effect of MBS at 10^-6 M in tracheal strips with an intact epithelium. By contrast, concentrations of MBS at 10^-7, 10^-6, and 10^-5 M significantly enhanced non-adrenergic non-cholinergic contractile responses in the guinea pig bronchial preparations (fig 2). The enhancement was maximal at a concentration of 10^-3 M (23.8 (9.6)%). Although the enhancement found at 10^-2 M was significant when compared with responses after vehicle alone, this effect was small. The addition of diluent alone had no significant effect. This enhancement was unaffected by removal of airway epithelium in both tracheal and bronchial preparations.

At higher concentrations of MBS (10^-3 and 10^-4 M) there was a time-dependent attenuation of contractile responses induced by electrical field stimulation in both tracheal and bronchial preparations. Maximal inhibition was observed with 10^-3 M MBS within 15 minutes of incubation with a gradual loss of inhibition with successive stimulations (fig 3). Maximal recovery to baseline responses occurred by 75 minutes of incubation (fig 3). This was not affected by removal of the epithelium.

EFFECT OF MBS ON RESPONSES TO RECEPTOR-MEDIATED STIMULI

MBS (10^-6 M) had no effect on acetylcholine-induced isometric contractile responses in the tracheal strips either with or without epithelial denudation. However, at the higher concentration of 10^-3 M there was a significant increase in the EC50 from 2.36 (geometric standard error of mean 1.42) x 10^-6 M to 19.1 (1.77) x 10^-6 M (n=5; p<0.05), but maximal responses to acetylcholine were not inhibited. In addition, similar responses were observed with MBS at 10^-3 M on substance P-induced contractile responses which were significantly inhibited at all concentrations for both tracheal and bronchial preparations; thus, at 10^-6 M, substance P-induced a contractile response of 28 (5.0)% of the maximal response compared with only 6.2 (2.0)% in the presence of MBS (10^-1 M) (p<0.005; fig 4).

Discussion

We have shown that MBS does not have direct contractile responses on guinea pig airway smooth muscle. Although MBS did not augment the cholinergic component of the tracheal response to electrical field stimulation, it increased non-adrenergic non-cholinergic responses in bronchi at concentrations that did not affect contractile responses to substance P. Thus, MBS may augment the release of tachykinins from airway nerves induced by electrical field stimulation, while having no effect in the unstimulated airway preparation. At higher concentrations MBS inhibited contractile responses to acetylcholine and substance P in tracheal and bronchial strips, suggesting a generalised effect on airway contractile responses, possibly by an interaction of MBS with airway smooth muscle receptors. Our study shows that MBS has a complex modulatory role on airway smooth muscle depending on the level of the airway and the concentration of MBS.

We performed this study in order to elucidate the mechanisms by which inhaled MBS induces bronchoconstriction in patients with asthma, although non-asthmatic subjects usually do not respond.12 MBS has been shown to increase bronchial blood flow in non-asthmatic subjects and to increase the release of airway microvascular leakage in the guinea pig.6 MBS may therefore cause luminal airway narrowing through bronchial vasodilation and the induction of bronchial wall oedema. In view of the limited effect of anticholinergic agents in blocking MBS-induced bronchoconstriction in both asthmatic patients and guinea pigs,24 it has been proposed that MBS induces airway narrowing in vivo through indirect mechanisms such as activation of non-cholinergic nerves. We could not show a direct effect of MBS on isometric contractile responses in guinea pig tracheal preparations; however, we did observe a non-specific inhibitory effect of MBS (10^-3 M) in guinea pig tracheal and bronchial preparations.

There has been some evidence to suggest that the effect of MBS is mediated through the activation of the cholinergic receptors. We have shown that MBS-induced effects on guinea pig airway smooth muscle are mediated through the activation of the cholinergic receptors. Thus, MBS-induced effects on guinea pig airway smooth muscle are mediated through the activation of the cholinergic receptors.

There is evidence to suggest that sulphur dioxide, which is generated from MBS solutions, may activate airway nerves. Thus, SO2 delivered to the lower airways of anaesthetised dogs stimulates bronchial and pulmonary C fibres with reflex increases in tracheal smooth muscle tone.11 Our studies suggest that, in
addition to inducing reflex effects, SO₂, generated from MBS may enhance the stimulation of local airway C fibres. This may represent a mechanism by which local axon reflexes, which have been postulated in the airways, 12 may be enhanced. An increase in the number of substance P-containing airway nerves has been reported in the airways of patients with asthma, 13 an observation which may account for a greater propensity for bronchoconstriction to MBS in asthmatic than in non-asthmatic subjects. 12

One mechanism by which sulphur dioxide and sulphites may induce bronchoconstriction in asthmatics is through their effect in increasing airway mucosal acidity. Sodium metabisulphite and sulphur dioxide convert to bisulphite 14 and bisulphite, in turn, enters into equilibrium with the sulphite ions. 15 These reactions are accompanied by the release of hydrogen ions, thereby acting as acids. Although the bronchoconstrictor effects of sodium sulphite, to which MBS is converted, are pH-dependent with the greatest effects occurring at low pH, acidity itself is unlikely to be the stimulus for bronchoconstriction. 16 It is unlikely that the in vitro airway preparations were exposed to an acid solution because of the buffering capacity of the Kreb’s solution. Addition of MBS to a maximum concentration of 10⁻³ M to Kreb’s solution did not alter its pH. In addition, a solution of sulphuric acid with a similar pH to MBS at 10⁻³ M did not induce airway smooth muscle contraction or augmentation of non-adrenergic, non-cholinergic responses.

Sodium sulphite is highly reactive with proteins 17 and can bind to rat airways. 18 Sulphite ions interact with disulphide bonds which can play a critical part in numerous cell membrane receptors. 19 Sulphonation by sulphites of specific disulphide bonds in the neuromuscular junction has been shown to increase spontaneous acetylcholine release. 20, 21 Thus, MBS may increase the release of acetylcholine or tachykinins induced by electrical field stimulation through sulphonation of disulphide bonds prejunctionally.

At high concentrations of MBS the direct contractile effects of acetylcholine and substance P were significantly inhibited with a rightward shift of the concentration-response curve without affecting the maximal response for the acetylcholine response. This suggests that MBS may be directly or indirectly acting as a competitive antagonist at the receptor sites of acetylcholine and substance P in a reversible fashion. Contractile responses to electrical field stimulation were inhibited at high concentrations of MBS, which may result from an effect of the contractile neural mediator on airway smooth muscle rather than an effect on their release, although the latter mechanism cannot be completely excluded. These inhibitory effects were reversible with repeated successive stimulations. The mechanism for this reversible inhibitory effect is unclear. One potential mechanism may involve the further interaction of MBS at high concentrations with sulphhydryl groups or disulphide bridges of various receptors on the smooth muscle membrane to alter receptor function or binding properties. 23-25

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