Effects of botulinum toxin A on the sphincter of Oddi: an in vivo and in vitro study

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Abstract

Background—Botulinum toxin A is a potent inhibitor of the release of acetylcholine from nerve endings. Local injection of botulinum toxin has recently been suggested to be helpful in sphincter of Oddi dyskinesia by decreasing sphincter of Oddi pressure.

Aims—To explore the mechanism of action of botulinum toxin A on sphincter of Oddi (SO) muscle.

Methods—Four piglets underwent duodenoscopy and SO manometry was performed. After obtaining a baseline pressure, the SO was injected with normal saline and the experiment repeated after one week. The SO was then injected endoscopically with botulinum toxin (40 U) with follow up manometry one week later. The sphincter of Oddi was removed from 10 pigs, cut into three rings, and placed in an organ bath. The force of contraction was measured and registered on a polygraph. Rings were stimulated by 70 V (10 Hz, 0.5 ms) electrical field stimulation for 20 seconds, exogenous acetylcholine (100 µM), and KCl (125 mM). Botulinum toxin (0.1 U/ml) or atropine (1 µM) was added to the incubation medium and the stimulation was repeated.

Results—Mean basal SO pressure in the pigs remained unchanged after saline injection but decreased to about 50% of baseline value following botulinum toxin injection (p=0.04). The contractions induced by direct stimulation of SO smooth muscle with KCl were not significantly affected by either atropine or botulinum toxin. In all rings exogenous acetylcholine induced contractions, which were totally blocked by atropine, but not by botulinum toxin. Electrical field stimulation induced contractions that were inhibited by both atropine and botulinum toxin.

Conclusion—Botulinum toxin inhibits pig sphincter of Oddi smooth muscle contractions by a presynaptic cholinergic mechanism, similar to that described in skeletal muscle.

Keywords: sphincter of Oddi; botulinum toxin; pig; ex vivo

EX VIVO STUDIES

Ten pigs (25–28 kg) were used for this study. The animals were anaesthetised with controlled 1–2% efrane inhalation via tracheostomy, performed under ketamine (10 mg/kg, intra-
and thiopental (10 mg/kg intravenously) induction. After laparotomy and duodenotomy, the entire SO was harvested and placed in an oxygenated physiological salt solution (pH 7.4, NaCl 119.0 mM, NaHCO₃ 25.0 mM, glucose 11.1 mM, CaCl₂ 1.6 mM, KCl 4.7 mM, KH₂PO₄ 1.2 mM, and MgSO₄ 1.2 mM). The duodenal mucosa of the papilla was stripped away under a microscope, but the inner ampullary epithelium was left intact. The section beyond the SO was cut into three transverse sections (rings), each 1–2 mm thick. The rings were placed between two hooks in an organ bath chamber containing continuously oxygenated physiological salt solution at 37°C. The force of the SO contraction was measured with an isometric force displacement transducer and registered on a polygraph (FT03 transducer, Model 7E Polygraph; Grass Instrument Co., Quincy, Massachusetts).

After a stabilisation period of 30 minutes, the SO rings were stimulated by either 70 V (10 Hz, 0.5 ms) electrical field stimulation for 20 seconds, or 100 µM exogenous acetylcholine (acetylcholine chloride; Sigma Chemical Co., St Louis, Missouri). Potassium chloride (125 mM), a potent direct smooth muscle stimulant, was used as positive control to test the viability of the preparation. After each stimulant the rings were rinsed twice with fresh incubation medium. Then 0.1 U/ml of botulinum toxin (Oculinum, Allergan Inc., Irvine, California) or 1 µM of atropine (atropine sulphate; Sigma) was added to the medium, and the stimulation was repeated after 20 minutes of incubation. All results are expressed as mean (SEM), unless otherwise stated. Paired Student’s t test (with logistic transformation in skewed distributions) was used to calculate the significance of differences. The protocol was approved by the Animal Care Committee of the University of Tampere.

Results

In vivo effects

No evidence of adverse effects to botulinum toxin injection were apparent. Basal SO pressure (day 0) in the four pigs was 13.3 (4) mm Hg (table 1). While no significant change occurred on day 7 after saline injection, basal SO pressure decreased by approximately 50% to 6.9 (2.8) mmHg after botulinum toxin injection (p=0.04).

Ex vivo effects

Baseline stimulation

KCl, exogenous acetylcholine, and electrical field stimulation induced remarkable contractions in SO rings (table 2). Our preliminary experiments had shown that each of these three stimuli elicited reproducible contractions in pig SO rings. Thus, there was no significant shift in contractile force generation when the preparation was repeatedly (up to three times) challenged with the given stimuli (data not shown).

Effects of atropine and botulinum toxin on SO contractile response

Neither atropine nor botulinum toxin had any effect on the KCl induced contractions (table 2).

<table>
<thead>
<tr>
<th>Type of stimulation</th>
<th>No</th>
<th>Control</th>
<th>Treatment*</th>
<th>p Value</th>
</tr>
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<tbody>
<tr>
<td>Potassium chloride</td>
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<td>2160 (526)</td>
<td>1770 (393)</td>
<td>&gt;0.1</td>
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<tr>
<td>Atropine</td>
<td>8</td>
<td>832 (195)</td>
<td>785 (176)</td>
<td>&gt;0.1</td>
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<tr>
<td>Acetylcholine</td>
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<td>&lt;0.01</td>
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<tr>
<td>Atropine</td>
<td>9</td>
<td>571 (133)</td>
<td>533 (118)</td>
<td>&gt;0.1</td>
</tr>
<tr>
<td>Botulinum toxin</td>
<td></td>
<td>918 (830)</td>
<td>27 (27)</td>
<td>&lt;0.01</td>
</tr>
<tr>
<td>Botulinum toxin</td>
<td>5</td>
<td>422 (258)</td>
<td>31 (20)</td>
<td>&lt;0.01</td>
</tr>
</tbody>
</table>

* In the presence of atropine or botulinum toxin.

Figure 1  (A) Representative tracings of sphincter of Oddi smooth muscle contractions induced by electrical field stimulation (EFS), and the effects of botulinum toxin and atropine. Small arrows indicate washes with fresh physiological salt solution. (B) Representative tracings of the sphincter of Oddi smooth muscle contractions induced by exogenous acetylcholine (ACH), and the effects of botulinum toxin and atropine. Small arrows indicate washes with fresh physiological salt solution.
2). On the other hand, both atropine and botulinum toxin seemed to inhibit electrically induced contractions (table 2, fig 1A). Atropine seemed to inhibit totally contractions induced by acetylcholine. However, contractions induced by acetylcholine were unaffected by botulinum toxin (table 2, fig 1B).

Discussion
The effects of botulinum toxin have been extensively studied in skeletal muscle systems. However, its effects on gastrointestinal smooth muscle, first described many decades ago, have only recently received attention. Pasricha et al showed for the first time that local injection of botulinum toxin inhibits resting lower oesophageal sphincter tone in a live animal model. This property was subsequently shown to be useful in the treatment of achalasia, a condition characterised by the failure of the lower oesophageal sphincter to relax. Since then, local injection of botulinum toxin has been used in an attempt to lower SO pressures in patients with sphincter of Oddi dysfunction, characterised by a high basal SO pressure.

The rationale behind the use of botulinum toxin in this condition is based on the assumption that net smooth muscle tone in the sphincter is determined by a balance between two opposing sets of nerve influences: an excitatory limb (acetylcholine, substance P, and others) and an inhibitory limb (vasoactive intestinal polypeptide, nitric oxide, and other neurotransmitters). The muscle in the sphincter of Oddi is richly innervated by cholinergic fibres. Acetylcholine is released by exocytosis from vesicles of cholinergic nerve endings and binds to muscarinic receptors on the smooth muscle cells, resulting in muscle contractions. This muscarinic effect of acetylcholine is blocked by atropine at the receptor level. Thus as expected, atropine reduces smooth muscle tone in the sphincter of Oddi, while cholinergic stimulation increases it. In contrast to atropine, botulinum toxin exerts its “anticholinergic” effect on striated muscle not by receptor antagonism but by inhibiting the release of acetylcholine from nerve endings. The effects of botulinum toxin therefore are expected to differ in important ways from those of atropine. We attempted to study these in a porcine model. The pig was chosen for these studies because the physiological nature of pig SO seems to be similar to that in man. Furthermore, we have previously shown that the peptidergic innervation of pig SO is very similar to that of human SO.

Our studies in live animals showed a significant reduction in basal SO pressure after botulinum toxin injection directly into the SO. Our experiments have therefore shown that net cholinergic impulses play a dominant role in the maintenance of basal SO tone in vivo. This has been an area of controversy in previous reports with conflicting results reported after vagotomy, as well as in response to atropine and other antimuscarinic agents.

Exogenous acetylcholine induced marked contractions in all SO rings. The acetylcholine induced contractions were totally inhibited by atropine but were unaffected by botulinum toxin. This proves that botulinum toxin has no postsynaptic anticholinergic effects such as blockade of muscarinic receptors or binding of acetylcholine to its receptors. Acetylcholine induced contractions in SO rings that could be inhibited almost totally by atropine, suggesting that they were mediated by acetylcholine. The effects of botulinum toxin were similar to atropine in this preparation. In our model, the replacement of all NaCl in the medium with KCl induced contractions in all SO rings, and these contractions were not significantly affected by either atropine or botulinum toxin. This implies that the previously noted effect of botulinum toxin on stimulated smooth muscle contraction is not a result of a non-specific toxic inhibition. Since we have shown that botulinum toxin has no postsynaptic anticholinergic effect, the toxin must therefore act by interfering with the release of endogenous acetylcholine.

In conclusion, botulinum toxin inhibits pig SO smooth muscle contractions by a presynaptic cholinergic mechanism, most likely similar to that described in skeletal muscle. In addition to its therapeutic potential, botulinum toxin may represent a powerful physiological tool to study the intricate workings of the enteric nervous system.
