Botulinum toxin treatment of synkinesia and hyperlacrimation after facial palsy

B Boroojerdi, A Ferbert, M Schwarz, H Herath, J Noth

Abstract

Objectives—To investigate the effects of injection of botulinum toxin type A (BTX A) into the orbicularis oculi muscle and lacrimal gland in patients with aberrant regeneration after facial palsy (facial synkinesias and hyperlacrimation).

Methods—The effect of the toxin injection (on average 75 mouse units of BTX A) into the orbicularis oculi muscle on facial synkinesias was assessed on a five point (0 to 4) scale in 10 patients with aberrant regeneration of facial nerve fibres after a peripheral facial nerve palsy. Six patients underwent a videographic control, which was assessed by a blinded independent investigator. In two patients with hyperlacrimation an extra dose of botulinum toxin (on average 20 mouse units BTX A) was injected into the lacrimal gland and the effect was assessed using the Schirmer test and on a three point scale.

Results—Botulinum toxin type A had a good to excellent (grades 3 and 4) effect over an average of six months after 91% of injections. In 9% the injections had a moderate (grade 2) effect. Patients with hyperlacrimation showed a nearly complete recovery. There were no systemic side effects but focal side effects due to a temporary weakness of the orbicularis oculi muscle were not uncommon.

Conclusions—Botulinum toxin type A is the treatment of choice in motor and autonomic effects of aberrant regeneration of facial nerve after a peripheral palsy. The required dose is similar to or slightly lower than the dose usually recommended for hemifacial spasm.

Keywords: facial nerve palsy; aberrant regeneration; botulinum toxin

Patients and methods

We treated eight women and two men with facial palsy with a total of 32 injections. The average age of the patients was 59 (range 26 to 85) years. Six patients were treated in the Department of Neurology in Kassel and four in Aachen. An informed written consent was obtained from all patients. The table contains a list of the patients including their clinical and treatment features.

BTX A ( Dysport ) was used; 500 mouse units were dissolved in 2.5 ml normal saline and an average dose of 75 (range 40–120)
List of the patients, clinical, and treatment data

<table>
<thead>
<tr>
<th>Patient</th>
<th>Age (y)</th>
<th>Sex</th>
<th>Clinical symptoms</th>
<th>Aetiology of facial palsy</th>
<th>Duration before treatment (y)</th>
<th>Injection sessions (n)</th>
<th>Dose of injected toxin in different injection sessions (range (mu))</th>
<th>Range of clinical effect on synkinesia (S) and hyperlacrimation (H)</th>
<th>Mean difference (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>67</td>
<td>F</td>
<td>A: Yes</td>
<td>Idiopathic</td>
<td>?</td>
<td>2</td>
<td>60–120</td>
<td>S: 4</td>
<td>—</td>
</tr>
<tr>
<td>2</td>
<td>85</td>
<td>F</td>
<td>A: Yes</td>
<td>Idiopathic</td>
<td>2</td>
<td>8</td>
<td>40–120</td>
<td>S: 2–4</td>
<td>+57</td>
</tr>
<tr>
<td>3</td>
<td>58</td>
<td>F</td>
<td>A: Yes</td>
<td>Idiopathic</td>
<td>30</td>
<td>5</td>
<td>60–120</td>
<td>S: 4 H: 1–2</td>
<td>—</td>
</tr>
<tr>
<td>4</td>
<td>74</td>
<td>F</td>
<td>A: Yes</td>
<td>Idiopathic</td>
<td>1</td>
<td>1</td>
<td>70</td>
<td>S: 3 H: 2</td>
<td>—</td>
</tr>
<tr>
<td>5</td>
<td>80</td>
<td>F</td>
<td>A: Yes</td>
<td>Idiopathic</td>
<td>1</td>
<td>5</td>
<td>60–100</td>
<td>S: 2–4</td>
<td>+47</td>
</tr>
<tr>
<td>6</td>
<td>70</td>
<td>M</td>
<td>A: Yes</td>
<td>Zoster oticus</td>
<td>5</td>
<td></td>
<td>70–80</td>
<td>S: 2–4 H: 2</td>
<td>+45</td>
</tr>
<tr>
<td>7</td>
<td>68</td>
<td>M</td>
<td>A: Yes</td>
<td>Resection of acusticus neurinoma</td>
<td>7</td>
<td>1</td>
<td>80</td>
<td>S: 3 H: 1</td>
<td>+67</td>
</tr>
<tr>
<td>8</td>
<td>31</td>
<td>F</td>
<td>A: Yes</td>
<td>Idiopathic</td>
<td>5</td>
<td>2</td>
<td>80–100</td>
<td>S: 4 H: 2</td>
<td>+26</td>
</tr>
<tr>
<td>9</td>
<td>34</td>
<td>F</td>
<td>A: Yes</td>
<td>Idiopathic</td>
<td>4</td>
<td>1</td>
<td>60</td>
<td>S: 4 H: 2</td>
<td>—</td>
</tr>
<tr>
<td>10</td>
<td>26</td>
<td>F</td>
<td>A: Yes</td>
<td>Hypolacrimation</td>
<td>4</td>
<td>2</td>
<td>60</td>
<td>S: 4 H: No clinical effect on hyperlacrimation</td>
<td>+40</td>
</tr>
</tbody>
</table>

Videographic assessment of the width of the affected lid was performed before and 2–4 weeks after the BTX A injection in patients 2, 5, 6, 7, 8, and 10. A: synkinesia; B: hyperlacrimation; C: gustatory sweating.

Mean difference = mean difference between relative width of affected lid before and 2–4 weeks after treatment v unaffected side.

Results

Injection of botulinum toxin into the orbicularis oculi muscle had a good (grade 3 and 4) clinical effect on the facial synkinesias after 29 injections (91%) and a moderate (grade 2) effect after three injections (9%). The average duration of the effect was 23.9 (range 13–40) weeks. The relative width of the affected lid compared with the unaffected side was measured both before and a few weeks after the treatment. A highly significant increase (47 (SD 18%); p<0.01) of the lid diameter during innervation of the orbicularis oris muscle was found (figure A, B).

The potential correlation between the dose and the subjective rating of the effect of the botulinum toxin was assessed in two ways: Firstly, all individual recordings of the dose and effect rating (k=32) were considered irrespective of whether several of them were obtained from the same patient. Secondly, the median dose level and the median effect rating were computed for each patient and the correlation was determined for these averaged values (n=10). The Spearman rank correlation coefficient “r” was computed to assess monotone relations between dose level and effect rating. For the individual values the correlation coefficient was r=−0.052 (p=0.788, k=32), indicating no systematic correlation between these two indices. For the median values the correlation coefficient was r=−0.431 (p=0.213, n=10), which was still not significantly different from zero.

Patients with hyperlacrimation who were treated with botulinum toxin injection into the lacrimal gland (patients 6 and 8) had a nearly complete recovery of this symptom. In patients 3, 4, 7, and 9 botulinum toxin was only injected...
into the orbicularis oculi muscle. All these patients showed a moderate to complete recovery of hyperlacrimation. The clinical effect of the injection was assessed using a Schirmer test before and after the toxin injection into the lacrimal gland, which showed a nearly equal lacrimation on both sides after the treatment (figure C, D).

Six patients (after 14 injections) had side effects, which lasted from a few days to two weeks. There were 12 episodes of incomplete lid closure, two episodes of ptosis, one episode of double vision, and two episodes of conjunctivitis. No patient showed dry eyes. There were neither serious nor systemic side effects. In three patients (patients 1, 3, and 6) only the highest doses of BTX A (100–120 mouse units) led to side effects, whereas with doses of 60–80 units no side effects could be found. In two patients (patients 2 and 3) no apparent relation between the dose and the incidence of side effects could be obtained, although these patients reported a subjective decline in the severity of their unwanted symptoms with decreasing doses of injected BTX A.

Discussion
With botulinum toxin type A we now have a powerful method to treat the synkinesias due to the aberrant regeneration of the facial nerve after a peripheral lesion. Moreover, our results suggest that it may be very effective in treating hyperlacrimation.

All patients investigated had a moderate to good recovery of their motor symptoms (facial synkinesias) and hyperlacrimation. We treated facial synkinesias with an average dose of 75 mouse units BTX A and hyperlacrimation with 20 mouse units in two patients. The administered dose was comparable with that used in other studies.12 13 There are no reports in the literature on the treatment of hyperlacrimation after facial nerve palsy. Although injections of BTX A into the lacrimal gland had an excellent clinical effect on hyperlacrimation, a considerable recovery could be found after subcutaneous toxin injections into the lateral part of the orbicularis oculi muscle, possibly due to diffusion of BTX A into the lacrimal gland.

Duration of the effect of the toxin in our patients was somewhat longer than that reported and lasted about 24 weeks; in the literature there is a reported duration of 11 to 20 weeks in patients with facial synkinesias after facial nerve palsy.12 13

Statistical analysis disclosed no significant correlation between the injected dose and the clinical effect of the toxin. One of the most important criteria for choosing the appropriate dose of the toxin was the rate and the subjective severity of the side effects. Accordingly the injected dose was not always chosen on the basis of the effect, but also on that of the expected side effects. This could possibly explain the lack of correlation between the dose and the clinical effect of the toxin. In addition, the size of the treatment group may have been too small to allow statistical significance and the dose range used by us was...
probably not large enough to establish such a correlation.

As in some patients only highest doses of BTX A (100–120 units) led to side effects, a starting dose of 60–80 units may be recommended to reduce the incidence of undesirable effects.

On average the patients in the present study needed a lower toxin dose, which showed a longer duration of the clinical effects than was the case for patients with blepharospasm. The required dose was comparable with or slightly lower than the dose given to patients with hemifacial spasm. As in some patients only highest doses of BTX A (100–120 units) led to side effects, a starting dose of 60–80 units may be recommended to reduce the incidence of undesirable effects.

On average the patients in the present study needed a lower toxin dose, which showed a longer duration of the clinical effects than was the case for patients with blepharospasm. The required dose was comparable with or slightly lower than the dose given to patients with hemifacial spasm. As in some patients only highest doses of BTX A (100–120 units) led to side effects, a starting dose of 60–80 units may be recommended to reduce the incidence of undesirable effects.

Due to the few patients investigated, this study presents insufficient data to comment on the long-term effects of the therapy of facial synkinesias and hyperlacrimation with the toxin. The effect of the injection on hyperlacrimation seems to be more stable than the effect on synkinesias.

Local injection of BTX A into the orbicularis oculi muscle is a very effective treatment of the facial synkinesias occurring after a peripheral facial nerve palsy. With doses of slightly less than half of the required dosage for blepharospasm and similar to that required in hemifacial spasm a good clinical effect may be obtained over about six months. Subcutaneous toxin injections into the lateral part of the orbicularis oculi muscle may have a beneficial effect on hyperlacrimation in most cases. With a suboptimal response to this treatment, injection of BTX A into the lacrimal gland may be more effective.

We are grateful to Dr K Willmes-v Hinckeldey for the statistical analysis and to Dr S Fellows (Department of Neurology, RWTH Aachen) for reviewing the English version of the manuscript.

Botulinum toxin treatment of synkinesia and hyperlacrimation after facial palsy

B Boroojerdi, A Ferbert, M Schwarz, H Herath and J Noth

J Neurol Neurosurg Psychiatry 1998 65: 111-114
doi: 10.1136/jnnp.65.1.111

Updated information and services can be found at:
http://jnnp.bmj.com/content/65/1/111

These include:

References
This article cites 12 articles, 1 of which you can access for free at:
http://jnnp.bmj.com/content/65/1/111#BIBL

Email alerting service
Receive free email alerts when new articles cite this article. Sign up in the box at the top right corner of the online article.

Topic Collections
Articles on similar topics can be found in the following collections

Drugs: CNS (not psychiatric) (1945)
Neuromuscular disease (1311)
Ophthalmology (842)

Notes

To request permissions go to:
http://group.bmj.com/group/rights-licensing/permissions

To order reprints go to:
http://journals.bmj.com/cgi/reprintform

To subscribe to BMJ go to:
http://group.bmj.com/subscribe/