Correction of post-traumatic anterior open bite
by injection of botulinum toxin type A into the anterior belly
of the digastric muscle: case report

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Abstract
Post-traumatic anterior open bite can occur as a result of broken balance among the masticatory muscles. The superior hyoid muscle group retracts the mandible downward and contributes to the anterior open bite. Denervation of the digastric muscle by injection of botulinum toxin type A (BTX-A) can reduce the power of the digastric muscle and help to resolve the post-traumatic anterior open bite. A patient with a bilateral angle fracture had an anterior open bite even after undergoing three operations under general anesthesia and rubber traction. Although the open bite showed some improvement by the repeated operation, the occlusion was still unstable six weeks after the initial treatment. To eliminate the residual anterior open bite, BTX-A was injected into the anterior belly of the digastric muscle. Following injection of BTX-A, the anterior open bite showed immediate improvement. Complication and relapse were not observed during follow-up. Long-standing post-traumatic open bite could be successfully corrected by injection of BTX-A into the anterior belly of the digastric muscle without complication.

Key words: Open bite, Neck muscles, Botulinum toxins

I. Introduction
Bilateral mandibular angle fractures separate two posterior condyle-bearing segments from the anterior tooth-bearing segment\(^1\). Treatment of bilateral mandibular angle fractures is difficult because of the high morbidity rate\(^2\). After the reduction of the angle fractures, anterior open bite can occur because two posterior condyle-bearing fragments are drawn superomedially by the action of masseter, temporalis, and medial pterygoid muscles. On the contrary, the mandibular anterior fragment is depressed by muscles such as digastric, geniobyoid, and mylohyoid\(^3\).

The position of the mandible is mainly determined by attached muscles. The muscle group for closing the mouth is mainly attached to the ramus, and the muscle group for opening the mouth is attached to the body. In case of bilateral mandibular angle fractures, the mandibular body is disconnected to the ramus area. Though open reduction restores the connection between the ramus area and the body area, the masseter muscle loses most of its attachment to the ramus because of the muscle being stripped during open reduction. Therefore, the balance of power between the muscle groups for closing and opening should be tipped toward the opening muscle group. The opening muscles will then track the mandible posteroinferiorly. This mechanism can induce the anterior open bite\(^1\). Since it is important for opening the mouth, the digastric muscle can be a treatment target for restoring the balance.

Intramuscular injection of botulinum toxin type A (BTX-A) can denervate muscle by preventing the release of acetylcholine from the terminal of the motor nerve; this leads to the paralysis of muscle\(^3,4\). BTX-A can cause muscle atrophy, thereby reducing the volume of muscle\(^5\). In this

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Thus, hardware removal and reduction and fixation were planned intra-orally. After hardware removal, bony malunion areas were separated using chisel and mallet. After the separation of proximal and distal segments, proper occlusions were established by manipulation. Internal fixation was done using reconstruction plates and screws. Genioplasty was performed simultaneously for esthetic reasons. One day after operation, the anterior open bite was improved compared to the preoperative condition, although it remained (Fig. 2, A). To resolve the residual anterior open bite, we applied additional study, the injection of BTX-A into the digastric muscle demonstrated the correction of post-operative anterior open bite. To the best of our knowledge, this is the first technical report using BTX-A for the correction of post-operative anterior open bite.

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rubber traction over a week. Rubber was located on the anterior tooth area and both premolar areas using 8 screws anchored on the alveolar bone. (Fig. 2. B) In spite of active rubber traction, there was no improvement in the anterior open bite. To eliminate residual anterior open bite, we decided to use BTX-A (Meditoxin; Medy-Tox, Seoul, Korea). Ten days after the third operation, 24-gauge intravenous catheters were inserted into the anterior belly of the digastric muscle at 4 sites. (Fig. 3) This muscle is attached to the digastric fossa as an internal aspect of the mandible and both sides of the midline. We could easily find the anterior belly of the digastric muscle using the digastric fossa. At each point, 5 units of BTX-A were injected into the muscle through the plastic catheter. A total of 20 units of BTX-A was used.

There was no early complication after the injection of BTX-A. The anterior open bite completely disappeared 3 days after BTX-A injection. (Fig. 4) Elastic traction was not used after achieving stable occlusion, with only screws kept in preparation against any unexpected accident. CBCT was taken 4 weeks after BTX-A injection, showing the resolution of anterior open-bite and good occlusion. (Fig. 5) During the follow-up, no event occurred. Occlusion was stable until 6 months after BTX-A injection. Thus, repeated BTX-A injection was not considered.

III. Discussion

Simple elastics traction can treat most cases of post-traumatic anterior open bite. The traction period may differ according to the trauma severity, but approximately 2-4 weeks has been reported to be adequate. If the post-traumatic open bite does not improve in spite of extensive elastic traction, the patient must feel discomfort and complain. Though the open bite is corrected through 3 weeks of rubber traction, the patient will suffer from 3 weeks' treatment, and the overall expense will also increase. BTX-A injection is a relatively safe, non-invasive method for reducing muscle volume and power. In the presented case, BTX-A injection showed rapid correction of post-traumatic open bite. Occlusal stability was maintained until 6 months' follow-up, and repeated injections were not required. Therefore, BTX-A injection into the anterior belly of the digastric muscle was an effective method for the correction of post-traumatic open bite.

BTX-A has been injected periodically for the correction of masseter muscle hypertrophy. Though BTX-A injection denervates the muscle, motor nerves regenerate and renervate again. Note, however, that BTX-A injection was done 1 time for the correction of post-traumatic open bite. Unlike masseter muscle hypertrophy, the post-traumatic open bite was due to the broken balance among muscles. The muscle group for closing the mouth was detached from the mandibular body in the bilateral angle fracture. Even after proper reduction, these muscles were still in a state of being
stripped from the bone. After the correction of open bite, the reattachment of these muscles could occur. Though the power of the digastric muscle would be recovered 3-4 months later, the muscle group for closing the mouth also exerted power after reattachment. In this case, the balance among muscles would not be impaired. For similar reasons, once the closure of the anterior open bite is achieved by elastic internmaxillary traction, anterior open bite seldom recurs after the removal of the elastics.

In single mandibular angle fracture, only one plate is usually applied on the external oblique ridge, which is a tension zone of the mandible. To secure proper stabilization, one more plate can be applied on the inferior border of the mandibular angle area. Note, however, that we applied only metal plate on the mid-portion of the fracture line. Because of previous surgery, severe bone resorption was observed on the external oblique ridge and inferior border of the mandibular angle, so we could not fix the plate and screw on this area. Thus, we had to apply only one reconstruction plate on the mid-portion of the mandible angle fracture line. Wound dehiscence, swelling, and hotness were observed prior to surgery; thus, we tried to operate with minimal dissection and plate screwing.

In this study, the anterior belly of the digastric muscle was selected as target of BTX-A injection. The main role of the digastric muscle is to depress the mandible, so it is primarily related to opening the mouth. The lateral pterygoid muscle is also related to opening the mouth. Note, however, that the lateral pterygoid muscle is seated deeply, and the complication after BTX injection will be higher. Suprahyoid myotomy can reduce muscle power. Radiofrequency therapy to the digastric muscle has been considered an additional therapy for preventing open bite or correcting the reestablished open bite. As an aggressive procedure, suprahyoid myotomy can cause severe morbidity. Radiofrequency therapy is a relatively less invasive method but requires an expensive device such as radiofrequency generator. In contrast, the BTX-A injection method is relatively cheap, requiring no special device.

BTX-A has been used for the treatment of dystonia. Previous studies showed that injection of BTX-A into the digastic muscle can treat jaw opening and oromandibular dystonia. The suggested doses of BTX-A ranged from 3.8 to 30 units. In this study, we used 20 units of BTX-A. We injected BTX-A into the muscle through plastic intravenous catheter instead of using metal needle injection syringe. Plastic intravenous catheter is a very useful drug delivery tool to the muscle. The diameter of the 24-gauge plastic intravenous catheter is very small, and the plastic catheter is not rigid. If BTX-A is injected into the blood vessel, there was no effect on the muscle, possibly causing rare but severe complication such as botulism accompanied by dysphagia, muscle weakness, and respiratory distress. When the plastic catheter is placed in a vessel, the aspirated blood can be observed in the plastic catheter. Therefore, unwanted intravascular injection can be avoided.

We believe the presented technique can be applied to correct open bite in orthodontic practice or orthognathic surgery. Simple BTX-A injection may speed up overall treatment and minimize the post-treatment relapse. To provide evidence of this hypothesis, a large-scale, case control study will be required. In this stage, BTX-A injection may be applied to selective open bite patients who do not respond to extensive rubber traction.

In conclusion, long-standing post-traumatic open bite could be corrected successfully by BTX-A injection into the anterior belly of the digastic muscle without any complication.

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