Assessment of intralesional injection of botulinum toxin type A injection for hypertrophic scars

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ABSTRACT

Background: Hypertrophic scars are dermal fibroproliferative disorders that typically develop after a skin injury heals. They can cause physical, psychological, and cosmetic problems. The management of such scars remains a matter of debate due to lack of effective treatment methods and the inability to prevent recurrences. Recent reports have demonstrated that botulinum toxin type A improves wound healing so it may play a role in treating hypertrophic scars. Aims: We assessed the effectiveness of intralesional botulinum toxin type A injection for treating hypertrophic scars. Methods: This prospective clinical study included twenty patients with hypertrophic scars. Intralesional injection of botulinum toxin type A was given once a month for three months with a follow-up period of six months. Each lesion was injected until slight blanching occurred. Therapeutic satisfaction of the patient and physician were recorded. Lesions were assessed for erythema, itching and pliability. Each item was assessed on a 5-point scale. Results: Therapeutic satisfaction was recorded as ‘good’ in 14 patients and ‘excellent’ in the remaining six. The mean erythema score decreased from 3.2 to 1.0, the mean pliability score from 3.3 to 0.8 and the mean itching score from 2.7 to 0.7. All of these were statistically significant. Limitations: A larger sample size and longer follow-up period would have given a better evaluation but was not feasible due to the high expenses involved. Conclusion: Botulinum toxin type A is a novel and promising therapy for hypertrophic scars with few side effects.

Key words: Botulinum toxin type A, hypertrophic scar, scars

INTRODUCTION

Hypertrophic scars are dermal fibroproliferative disorders that usually develop after a skin injury heals.[1] Patients often experience physical deformities, restricted range of motion, pain and pruritus.[2] These scars are seen in about 50% of post surgical wounds and in more than 50% of healed deep burns.[3] Certain sites are predisposed because of increased skin tension, such as the chest and back.[4] These scars represent an aberration in the fundamental process of wound healing which includes hemostasis, inflammation, proliferation and the remodeling process.[5] Only hypertrophic scars have myofibroblasts with α smooth muscle actin expression which is believed to be an important element in the pathogenesis of contractures.[6] The exact etiopathogenesis of hypertrophic scars is still unknown.[7] The numerous treatments available include surgical excision, intralesional steroid injection, radiation therapy, lasers and pressure therapy, none of which offer satisfactory therapeutic results.

Botulinum toxin type A is an exotoxin of the anaerobic spore forming bacterium, Clostridium botulinum. It has become a useful tool for treating the hyperactive muscles of facial expression. It acts by inhibiting the exocytosis of acetylcholine which indirectly blocks neuromuscular transmission resulting in muscle...
flaccidity.\[^8\] Although initial studies focussed on conditions related to skeletal muscle hyperactivity, it has become clear that botulinum toxin is effective in other disorders as well. It is being investigated for the treatment of smooth muscle conditions such as achalasia, anal fissure and overactive bladder. It is an accepted treatment for a few autonomic disorders such as primary hyperhidrosis. This led to the exploration of additional mechanisms of action, apart from the inhibition of acetylcholine release.\[^9\] Several papers have reported that botulinum toxin type A can minimize facial scarring by reducing the muscle tension that acts on the healing wound. It may produce changes in the muscle spindles, that could lead to altered sensory input and changes in the cell cycle distribution of fibroblasts derived from the hypertrophic scar.\[^10,11\]

**METHODS**

This study included twenty patients with hypertrophic scars who attended our dermatology outpatient clinic between May 2013 and November 2013. The study was assessed and approved by the institutional ethics committee. The procedure was explained to the patients and informed consent was obtained. All patients underwent history taking and clinical examination. Lesions that were present for more than one year and patients who were above the age of eighteen were included in the study. Exclusion criteria were pregnancy, lactation, local infection, patients on retinoids or anabolic steroids, severe systemic disease, pre-existing neuromuscular or cardiovascular disease, patients with unrealistic expectations or those who were treated for hypertrophic scars in the past.

Patients were treated with intralesional botulinum toxin type A as monotherapy. An intralesional injection of botulinum toxin type A (Botox Allergan®, Irvine, CA, USA. 100 U vacuum-dried powder in a single-use vial for reconstitution diluted in 2 mL of sterile, preservative-free 0.9% saline to constitute a solution at a concentration of 4 U/0.1 mL) was administered once a month for a total period of three months. It was injected into the body of the scar with the help of a 9-gauge needle (0.3 mm × 0.8 mm) until slight blanching was visible. The dose was adjusted to 2.5 U/cm\(^3\) of the lesion, not exceeding 100 units per session. Photographs of the scars were taken before and immediately after each treatment session and at the 3\(^{rd}\) and 6\(^{th}\) month after the last treatment session. At the 6\(^{th}\) month follow-up period, the overall assessment information was obtained and graded subjectively on a 5-point scale as follows, 0: no improvement; 1: poor (up to 25% improvement); 2: fair (26–50% improvement); 3: good (51–75% improvement); or 4: excellent (76–100% improvement). The physician as well as the patient recorded this information. Erythema was graded on a 5-point scale as follows, 0: no erythema, 1: mild, 2: moderate, 3: severe, or 4: very severe. Lesion lightening was defined as the percentage of erythema reduction compared to the erythema before injection. Pliability (induration) also was graded on a 5-point scale as follows, 0: no induration, 1: mild, 2: moderate, 3: severe, or 4: very severe. Lesion softening was defined as the percentage of pliability reduction compared to the pliability before injection. The severity of itching was graded on a 5-point scale as follows, 0: no itching, 1: mild, 2: moderate, 3: severe, or 4: very severe. Improvement in itching was defined as the percentage of itching reduction compared to the itching before injection.

During the six month follow-up period, the patients were monitored for the development of any complications such as pain, bleeding, edema, erythema, hypopigmentation, hyperpigmentation, secondary infection and recurrence.

**Statistical analysis**

The data was analyzed using Statistical Package for Social Science version 20 (SPSS Inc., Chicago, IL, USA). Parametric data were expressed as mean ± standard deviation and nonparametric data were expressed as numbers and percentages of the total. Comparisons of the pre-treatment and post-treatment mean ± standard deviation were done using the paired \(t\)-test. A \(P\) value of \(P < 0.05\) was considered statistically significant.

**RESULTS**

The study enrolled 14 men and 6 women with a mean age of 23.9 ± 5.4 years (range 19–35 years). The mean duration of the hypertrophic scars was 1.9 ± 0.84 years (range 1–3 years), and the mean lesion size was 2.85 ± 1.4 cm\(^3\) (range 1.25–6.0 cm\(^3\)). There were six cases of hypertrophic scars on the face and neck, four on the chest and shoulders and the remaining ten were on the extremities [Table 1].

According to the overall assessment of clinical response by the physician, 14 lesions had “good” improvement, and 6 lesions had “excellent” improvement. Patient
assessment showed “good” improvement in 12 lesions and “excellent” improvement in 8 lesions. There was a significant difference in the mean erythema scores from the pre-treatment (3.2 ± 0.78) to post-treatment stage (1.0 ± 0.66) (P = 0.000) [Table 2, Figures 1 and 2].

There was no recurrence or complications in any of the lesions during the 6-month follow-up period.

**DISCUSSION**

Many researchers have tried to treat hypertrophic scars by reducing excess collagen with the least amount of complications or recurrence rates. Our study demonstrates the ability of botulinum toxin type A injection to significantly decrease erythema, induration and itching with an improvement in the overall appearance.

Botulinum toxin type A acts by blocking cholinergic neuromuscular transmission or cholinergic autonomic innervation of exocrine glands and smooth muscles. It inhibits the action of soluble N-ethylmaleimide-sensitive factor attachment protein receptor protein, which is involved in the fusion of acetylcholine-containing synaptic vesicles with the plasma membrane.[12] The action of botulinum toxin type A on hypertrophic scars is believed to be a temporary denervation of smooth muscle fibers decreasing the tension in the scar tissue. This is one of the main factors that determine the degree of scar formation.[13] Hypertrophic scars result from fibroblast proliferation which is responsible for the enhanced collagen production that leads to excessive scarring.[14] By decreasing the tension in the scar, botulinum toxin causes local fibroblasts to gradually change their functional status to proliferate slower, secrete less biologically active mediators and synthesize less extracellular matrix and collagen resulting in improvement of hypertrophic scars.[15]

Recent studies have elucidated the molecular connections between the biological mechanisms of botulinum toxin type A and hypertrophic scars. An *in vitro* study by Zhibo and Miaobo showed

| **Table 1: Patient characteristics** |
|-----------------------------------|
| Patient number | Site of the lesion | Age of the patient (in years) | Size of the lesion (cm²) | Sex | Duration of the lesion (in years) |
|----------------|--------------------|-------------------------------|--------------------------|-----|----------------------------------|
| 1              | Left arm           | 19                           | 1.25                     | Male | 1                                |
| 2              | Forehead           | 18                           | 1.75                     | Male | 3                                |
| 3              | Right wrist        | 23                           | 4                        | Female | 3                              |
| 4              | Neck               | 19                           | 2.5                      | Male | 2                                |
| 5              | Right thigh        | 25                           | 2.5                      | Female | 3                              |
| 6              | Right forearm      | 19                           | 2                        | Male | 1.5                              |
| 7              | Chest              | 23                           | 2.5                      | Male | 1                                |
| 8              | Right shoulder     | 33                           | 6                        | Female | 2                              |
| 9              | Left elbow         | 23                           | 2                        | Male | 1.5                              |
| 10             | Neck               | 35                           | 3                        | Male | 1                                |
| 11             | Left cheek         | 26                           | 4                        | Male | 3                                |
| 12             | Left cheek         | 21                           | 2                        | Male | 1                                |
| 13             | Chest              | 25                           | 2.5                      | Female | 2                              |
| 14             | Right ankle        | 35                           | 4                        | Male | 3                                |
| 15             | Left leg           | 18                           | 3.5                      | Male | 1.5                              |
| 16             | Right knee         | 20                           | 2.25                     | Female | 2.5                            |
| 17             | Right arm          | 30                           | 1.75                     | Male | 2                                |
| 18             | Left leg           | 19                           | 6                        | Female | 1                              |
| 19             | Forehead           | 22                           | 2                        | Male | 2                                |
| 20             | Left shoulder      | 23                           | 1.5                      | Male | 1                                |

| **Table 2: Improvement of hypertrophic scars after intralesional injection of botulinum toxin type A** |
|------------------------------------------------------------------------------------------------|
| Evaluation factors | Pre-treatment X±standard deviation | Post-treatment X±standard deviation | Total X±standard deviation | T-test | P value |
|---------------------|------------------------------------|-------------------------------------|---------------------------|--------|---------|
| Pliability          | 3.3±0.48                           | 0.80±0.42                           | 2.5±0.16                  | 15.00  | 0.000*  |
| Itching             | 2.7±0.67                           | 0.70±0.48                           | 2.0±0.81                  | 7.74   | 0.000*  |
| Erythema            | 3.2±0.78                           | 1.0±0.66                            | 2.2±0.63                  | 11.00  | 0.000*  |

*Significance level P<0.05
that a higher percentage of fibroblasts were in the non-proliferative phases (G0 and G1) when treated with botulinum toxin type A (64%) as compared to the control group (36%).[11] They also confirmed the ability of botulinum toxin type A to reduce the expression of transforming growth factor \( \beta_1 \) in keloid fibroblasts.[16] Transforming growth factor \( \beta_1 \) is considered to be the main regulator in the pathogenesis of hypertrophic scars and is associated with excessive deposition of scar tissue and fibrosis.[17]

Another in vitro study demonstrated that botulinum toxin type A could promote apoptosis, inhibit transforming growth factor \( \beta_1 \) protein expression, affect the cell cycle and cell dynamics to inhibit the proliferation of fibroblasts derived from hypertrophic scars.[11] Botulinum toxin type A also inhibits the release of substance \( P \) which requires the soluble N-ethylmaleimide-sensitive factor attachment protein receptor protein activity. This contributes to the improvement of erythema, pain, and itching.[18,19]

However, another study showed that the expression of cytokines and growth factors in human keloid tissue was unaffected by botulinum toxin type A incubation suggesting that there was no significant therapeutic role for botulinum toxin type A injections in keloid treatment.[20]

Clinically, several studies have confirmed the positive effect of intradermal botulinum toxin type A in hypertrophic scars.[21] Xiao et al. treated 19 patients with hypertrophic scars once monthly with intradermal botulinum toxin type A for 3 months, with a follow-up for at least 6 months.[22] Twelve patients noted that the improvement of their lesions was “good,” and the other seven said that the improvement was “excellent.” According to the overall assessment by the plastic surgeons, the improvement in fifteen lesions and “excellent” improvement in the remaining four. The mean erythema score decreased from 3.41 to 1.23, the pliability score decreased from 3.85 to 0.78, and the itching score decreased from 3.50 to 0.83. Zhibo and Miaobo have studied intradermal injection with botulinum toxin type A at a concentration of 35 U/mL (range 70–140 U/session) with 3-month intervals for a maximum of 9 months in twelve patients.[16] At 1-year of follow-up, three patients demonstrated excellent improvement, five had good improvement and the remaining four had fair improvement. There were no signs of recurrence during the follow-up period. In contrast, Gauglitz et al. determined that repeated injections with botulinum toxin type A for up to six months did not change the macroscopic appearance, morphology or size of scars in four patients. They
had used optical profilometry to confirm the visual impression.[23]

Limitations
A larger sample size and longer follow-up period would have given a better evaluation, but was not feasible due to the high cost of the study.

CONCLUSION
This study concludes that monthly injections of intraliesional botulinum toxin type A for a period of three months is associated with an improvement of erythema, pliability, and itching in hypertrophic scars. It is a novel and promising therapeutic agent with no side effects noted so far.

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Conflicts of interest
There are no conflicts of interest.

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