Short-term safety of facial rejuvenation using an absorbable polydioxanone monofilament thread in patients with mild-to-moderate facial skin sagging

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INTRODUCTION

Over the past decade, facial rejuvenation procedures using surgical lifting through excision of redundant skin have been considered the gold standard [1,2]. However, surgical facial rejuvenation procedures may still produce postoperative complications such as infection, skin necrosis, hematoma, seroma, and damage to the branches of the facial nerve. Disadvantages also include visible scars and a prolonged recovery time [3,4]. Patients are therefore in need of minimally invasive facial rejuvenation techniques with lower morbidity and relatively rapid wound healing [3]. This has led to the introduction of novel nonsurgical facial rejuvenation procedures with volumizing effects. However, the disadvantages of these procedures include increased facial volume, unnatural contour, and a visible shift of the center of gravity to the lower third of the face [2,4-6]. Ablative or non-ablative resurfacing techniques are effective in improving the skin surface, but their disadvantages include a lack of sufficient lifting of the underlying ptotic tissues [7,8].
Facial rejuvenation techniques involving thread lifts are performed by passing sutures under the facial skin to treat sagging and facial flaccidity, and their advantages include a relatively short recovery time and relatively small incisions [6,9]. As such, facial rejuvenation techniques using threads are considered an alternative to conventional rhytidectomy procedures [10,11]. Their efficacy, safety, and durability therefore deserve special attention from four standpoints: (1) treatment effects, (2) sustainability, (3) in vivo safety, and (4) the risk of serious adverse events, such as intraoperative damage to the facial nerve [6].

Mint Lift (HansBiomed Co., Ltd., Seoul, Korea) is a violet-colored, absorbable polydioxanone monofilament thread with a wire length of 43 cm and a United States Pharmacopeia size of 0. Classified as a class IV medical device, its properties include transparency visible during the first postoperative month, bidirectional helical barbs that providing strong initial skin anchorage, and a lack of attachment of the yarn to the needle. It is initially inserted using a curved needle (5/8) and then later using an 18-gauge blunt cannula, for which a disposable external trocar is concomitantly used [9,12].

Given this background, we describe the short-term safety of a facial rejuvenation technique using the Mint Lift in patients with mild-to-moderate facial skin sagging.

METHODS

Study patients and setting
The current multicenter, retrospective, case-series study was conducted in a total of 196 patients who visited two local clinics in Seoul, Korea between 2016 and 2018. The inclusion criteria for the current study were as follows: (1) apparent good health; (2) mild-to-moderate skin sagging; (3) facial rejuvenation surgery performed using the Mint Lift 17 or 43; and (4) available follow-up data. The exclusion criteria for the current study were as follows: (1) a past history of incisional facelift within the 12 months prior to undergoing facial rejuvenation at a study center; (2) loss to follow-up (n = 17); and (3) otherwise ineligible for the current analysis according to our judgment (n = 0).

As such, we evaluated a total of 179 patients in the current study; informed consent was waived due to the study’s retrospective nature. The current study was conducted in compliance with the relevant ethical guidelines.

Surgical design and technique
The surgical procedure was performed under local anesthesia using 1% to 2% lidocaine hydrochloride mixed with epinephrine hemitartrate at a ratio of 1:200,000 at the needle insertion point and the exit point along the passage line for thread insertion.

For the surgical design and technique using the Mint Lift 43 and 17 threads, both the entry and exit points were designed on one side of the face.

The entry points for the Mint Lift 43 were located in the temporal region; two entry points were chosen 2 cm medial to the temporal hair line, superior to the superior helix of the ear, with a gap distance of approximately 2–3 cm. After a slit incision was made, the Mint Lift 43 was anchored to the entry and exit points using a specialized needle. This was followed by insertion of a two-piece cannula into the entry and exit points and removal of the inner guide. Then, two or three Mint Lift 43 threads were inserted into the outer cannula in the deep temporal fascia using a specialized needle, such as a temporal needle, and were then pulled through the exit point. The outer cannula was then removed. During this process, special attention should be paid to the passage of the outer cannula through the deep subcutaneous layer.

For the insertion of the Mint Lift 17, two or three entry points were chosen along the hair line, and the exit point was at the target site 1 cm proximal to the nasolabial folds or the apex of the jowl. Unlike in the insertion of the Mint Lift 43, no passage of the Mint Lift 17 through the exit point was required. For the insertion of the Mint Lift 17, a small hole was made at the entry point using an 18-gauge needle, and a two-piece cannula was inserted. Upon advancement through the deep subcutaneous layer to the target site, the inner guide was removed and three or four Mint Lift 17 threads were inserted. This was followed by the removal of the outer cannula.

Patient evaluation and criteria
All data from our clinical series of patients were expressed as mean ± standard deviation or as the number of cases and corresponding percentage, where appropriate. Moreover, the safety profile of this technique was assessed in terms of the incidence of postoperative complications.

Table 1. Baseline characteristics of the patients

<table>
<thead>
<tr>
<th>Variable</th>
<th>Total</th>
<th>Grace Plastic Surgery Clinic (n=107)</th>
<th>Songdo IB Clinic (n=72)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (yr)</td>
<td>44.4±14.4</td>
<td>40.7±12.1</td>
<td>49.9±15.7</td>
</tr>
<tr>
<td>Sex</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Male</td>
<td>6 (3.4)</td>
<td>4 (3.7)</td>
<td>2 (2.8)</td>
</tr>
<tr>
<td>Female</td>
<td>173 (96.6)</td>
<td>103 (96.3)</td>
<td>70 (97.2)</td>
</tr>
<tr>
<td>Nationality</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Korean</td>
<td>131 (73.2)</td>
<td>75 (70.1)</td>
<td>56 (77.8)</td>
</tr>
<tr>
<td>Chinese</td>
<td>26 (14.6)</td>
<td>24 (22.4)</td>
<td>2 (2.8)</td>
</tr>
<tr>
<td>Japanese</td>
<td>22 (12.2)</td>
<td>8 (7.5)</td>
<td>14 (19.4)</td>
</tr>
<tr>
<td>Follow-up period (mo)</td>
<td>22.3±9.8</td>
<td>22.5±9.6</td>
<td>21.9±10.1</td>
</tr>
</tbody>
</table>

Values are presented as means±SD or number (%).
should easily dissolve at the same rate as that at which the tissue causes the material used for thread lifts should be strong, but also tensile strength are particularly essential features of thread lifts, be absorbability [15-17]. High biodegradability in body fluids and high lack of an allergic reaction, lack of carcinogenicity, and high ab-
lack of bacterial growth, high tensile strength, ease of sterilization, inertness, lack of tissue reaction, ease of handling, ease of knotting, lack of bacterial growth, high tensile strength, ease of sterilization, lack of an allergic reaction, lack of carcinogenicity, and high absorbability [15-17]. High biodegradability in body fluids and high tensile strength are particularly essential features of thread lifts, because the material used for thread lifts should be strong, but also should easily dissolve at the same rate as that at which the tissue 

RESULTS

Baseline characteristics of patients
Our clinical series included 179 patients (six men and 173 women; mean age, 44.4 ± 14.4 years). The mean follow-up was 22.3 ± 9.8 months. In total, 107 and 72 patients were treated at the Grace Plastic Surgery Clinic and the IB Plastic Surgery Clinic, respectively. The baseline characteristics of the patients are presented in Table 1.

Illustrative cases
Treatment outcomes were described using photographs taken preoperatively and at 3 or 6 months, for which two representative cases are illustrated (Figs. 1, 2).

Safety outcomes
In our series, the postoperative complications included ecchymosis (1.7%), infection (0.4%), thread extrusion (1.1%), and skin dimpling (1.5%). No cases of iatrogenic nerve injury were observed, and all postoperative complications spontaneously resolved.

DISCUSSION

Over the past several years, there have been marked changes in facial rejuvenation techniques using absorbable thread lifts with polydioxanone threads [13-16].

Thread lifts are used for nonsurgical facial tightening. Ideal requirements for threads used in these procedures include biological inertness, lack of tissue reaction, ease of handling, ease of knotting, lack of bacterial growth, high tensile strength, ease of sterilization, lack of an allergic reaction, lack of carcinogenicity, and high absorbability [15-17]. High biodegradability in body fluids and high tensile strength are particularly essential features of thread lifts, because the material used for thread lifts should be strong, but also should easily dissolve at the same rate as that at which the tissue

**Fig. 1.** A 44-year-old woman receiving the Mint Lift 43 at 3 months. (A) A 44-year-old woman visited us with a chief concern of skin sagging on the right side of the face, for which she underwent facial rejuvenation using the Mint Lift 43. Two threads were inserted in the deep temporal fascia, a process in which special attention should be paid to the passage of the outer cannula through the deep subcutaneous layer. (B) At 3 months, the patient was satisfied with the treatment outcomes and reported no postoperative complications.

**Fig. 2.** A 31-year-old woman receiving the Mint Lift 17 at 6 months. (A) A 31-year-old woman with skin sagging on the left side of the face underwent insertion of four Mint Lift 17 threads in the deep subcutaneous layer. (B) At 6 months, the patient was satisfied with the treatment outcomes and reported no postoperative complications.

It is apparent that no single thread lift fulfills all of the above criteria. Plastic surgeons should therefore select the optimal thread lift based on the type of aesthetic procedure they plan to perform. This issue is also closely associated with a lack of consistent requirements for thread lifts when these tools are used to lift the skin [10,19]. Finally, the use of optimal thread lifts for aesthetic facial treatments is an essential factor for both obtaining good cosmetic results and avoiding complications [6,14,15,20-22].

In our series, only nonserious complications were encountered, such as ecchymosis, infection, thread extrusion, and skin dimpling, all of which spontaneously resolved. These results indicate that Mint Lift is a safer modality than other facial lifting procedures or surgical methods; this may be due to the simplicity of the procedure, the low risk of iatrogenic tissue injuries, and differential features of our techniques, such as the use of a specialized cannula and absorbable sutures, as well as variability in the depth of cannula insertion. The short-term safety of the Mint Lift has been documented previously. According to Baek et al. [12], only a few minor complications arose with no notable problems in a 24-week, prospective, single-center, single-arm, pre-post test design, open-label clinical study of 61 patients with deep nasolabial folds and lower facial drooping (three men and 58 women). Those complications included pustule formation, pain, swelling, a subjective feeling of tightness, and skin dimpling, although they were mostly self-limited. Thus, Baek et al. concluded that facial rejuvenation using the Mint Lift may be a favorable modality. Moreover, in a separate study, Bae et al. [23] reported no major complications such as thread exposure, alopecia, or parotid gland injury. Although five patients complained of ecchymosis for up to 3 weeks, they did not require treatment and returned to their daily lives within 1 week postoperatively.

Herein, we describe the safety of a facial rejuvenation technique using the Mint Lift with a review of previous published studies on this topic. However, further long-term follow-up studies are warranted to corroborate our results.
NOTES

Conflict of interest
This study was supported by HansBiomed Co., Ltd. No other potential conflict of interest relevant to this article was reported.

Ethical approval
The study was performed in accordance with the principles of the Declaration of Helsinki.

Patient consent
The patients provided written informed consent for the publication and the use of their images.

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REFERENCES