Facial Augmentation by Intra-Oral Delivery of Autologous Fat

**INTRODUCTION**

Changes in the deep subcutaneous tissues of the face are intrinsic to aging and most often associated with infrastructure collapse [1]. Indeed, atrophy of the facial form is a hallmark of the aging face [2]. As fat harvesting techniques have improved, public confidence in synthetic materials (e.g., silicone, Teflon® [Dupont, Wilmington, DE, USA]), and foreign tissues (collagen) has waned, and problems with absorption and long-term survival of translocated fat have diminished. Autologous fat transplants—injectons of extracted fatty tissue from the abdomen, hips, buttocks, or thighs—are widely used for facial augmentation. Conventional techniques involve placing harvested fat deep in subcutaneous tissue through skin punctures. Trans-oral fat injections at the buccal and labial sulci and through the mucosa of the oral commissures obviate external scars. We report experience with this technique for augmentation of the cheeks, zygomatic arch, nasolabial folds, upper and lower lips, and chin in 130 patients over a 14-year period.

**Background** As harvesting techniques have improved the viability of transposed adipose tissue, autologous fat transfer is increasingly utilized for facial augmentation. Conventional techniques involve placing harvested fat deep in subcutaneous tissue through skin punctures. Trans-oral fat injections at the buccal and labial sulci and through the mucosa of the oral commissures obviate external scars. We report experience with this technique for augmentation of the cheeks, zygomatic arch, nasolabial folds, upper and lower lips, and chin in 130 patients over a 14-year period.

**Methods** We report an observational case series of 147 procedures performed in 130 patients (118 women and 12 men, 19–69 years of age) in which 10 to 60 mL autologous fat supernatant harvested from the hips, buttocks, or abdomen were injected through the oral mucosa for augmentation. Single preoperative doses of antibiotic and methylprednisolone were administered in all cases. Fat frozen and banked during initial treatments was used for additional augmentation 3 to 6 months after initial treatments in 46% of cases. Outcomes were assessed clinically with physical exams and photographs.

**Results** Mucosal puncture wounds healed uneventfully. Most patients tolerated the procedures well and recovered rapidly compared to historical experience with conventional percutaneous techniques. One patient developed a sterile seroma 3 weeks after facial augmentation that responded to needle aspiration.

**Conclusions** Transmucosal, intra-oral autologous fat grafting was associated with minimal visible scarring, swelling, or infection at the recipient site. This technique may be considered an alternative to percutaneous transfer for proximate peri-oral facial augmentation.

**Keywords** Cosmetic techniques, Reconstructive surgical procedures, Rejuvenation, Subcutaneous fat, Tissue transplantation
scribe our experience with 147 intra-oral procedures in 130 patients for augmentation of the cheeks, zygomatic arch (ZA), nasolabial folds, chin, and upper and lower lips.

METHODS

The technique involved injection of fat through intra-oral punctures at the buccal and labial sulci and through the mucosa of the oral commissures. Between 10 and 60 milliliters of supernatant fat harvested from the hips, buttocks, or abdomen, were injected using 10-gauge needles under local anesthesia. A single dose of an antibiotic, generally cephalexin, was given preoperatively, and a single dose of methylprednisolone was administered intra-operatively in all cases.

The donor sites were selected for cosmetic benefit of fat removal, usually the outer thighs, buttocks, or abdomen, since fat from these regions has high lipogenic activity [2,9]. Patients were positioned supine, prepped with benzalkonium chloride solution, and draped. Mild preoperative sedation with oral diazepam was supplemented with nitrous oxide administered via nasal cannula throughout the procedure. Xylocaine® (AstraZeneca LP, Wilmington, DE, USA) (1%) and epinephrine (1:100,000) were injected at the donor site in 400 mL lactated Ringer’s solution with lidocaine (Marcaine®; Hospira, Lake Forest, IL, USA). A 1-mm incision was made with a #15 blade. Fat was then aspirated through the incision with a 60-mL syringe using a size-3 Becker Toomey cannula. A total of 100 mL of fat was harvested, mixed, cleansed with saline without centrifugation, and transferred to 10-mL syringes. The donor site incision was closed with 5-0 Vicryl® sutures. Cleaned fat was stored for later use in a standard freezer compartment in 10-mL sterile syringes and sterile plastic bags.

The recipient sites—generally the nasolabial folds, cheeks, lips, and chin—were anesthetized intra-orally with 1% Xylocaine® (AstraZeneca) with epinephrine (1:100,000) using 25-gauge needles, and fat was injected using 10-gauge needles. Intra-oral incisions 1 mm in length were made on the right and left corners of the commissures of the lip, and the needle was advanced through the subcutaneous tissue while fat was gently injected. Injections in the cheeks and chin were made through a 1-mm buccal or labial incision above the periosteum, delivering enough fat to achieve approximately 30% over-correction (Fig. 1). Approximately 30 to 40 mL was injected to the cheeks, nasolabial folds, lips, and chin during the initial procedures. Typically, an additional 20 to 30 mL were injected during subsequent sessions 3 to 6 months later. Postoperatively, durability of fat grafting, infection, and scarring were assessed clinically with physical exams and photographs.

RESULTS

The mean age of the 118 women and 12 men was 50 years (range 19–69 years). One hundred percent of patients had augmentation of the nasolabial folds and cheeks, 50% also had lip augmentation, and 30% also had augmentation of the chin. Fat frozen and banked at the time of initial treatments was used for additional augmentation 3 to 6 months after initial treatment in 46% of cases. We observed no visible scars, and regional swelling due to the volume of injected liquid was rapidly absorbed, so the appearance of over-correction subsided after 2 weeks (Fig. 2). Mucosal puncture wounds healed uneventfully, and most patients tolerated the procedures well and recovered rapidly compared to historical experience with conventional percutaneous techniques. One patient developed a sterile seroma 3 weeks after facial augmentation that responded to needle aspiration. The overall infection rate was the same for men and women and across each injection site. Long-term durability depended on initial over-correction, similar to the conventional approach. This technique was employed to augment the cheeks, ZA, nasolabial folds, upper and lower lips, and chin (Fig. 2 and 3).

DISCUSSION

Fat grafting for facial augmentation was first described in 1893 when...
Fig. 2. Attenuated lips, prominent nasolabial folds, and malar flattening. A 39-year-old female with attenuated lips, prominent nasolabial folds, and malar flattening received autologous trans-oral fat injection. (A) A preoperative photograph. (B) Postoperative week one. (C) Postoperative week two.

Fig. 3. Malar flattening, visible nasolabial folds, and attenuation of lips. A 50-year-old female with malar flattening, visible nasolabial folds, and attenuation of the lips received autologous trans-oral fat injection. (A) A preoperative photograph. (B) Postoperative month 6.
Neuber [10] reconstructed facial defects by transferring 1-cm pieces of fat from the upper arm. Injection of autologous fat to correct post-rhinoplasty deformities was first reported by Brunnings in 1911, who injected small pieces of fat using a hypodermic needle and syringe [11]. Fat transplantation was revived after the introduction of liposuction in 1977 [12], and Fournier coined the term "liposculpture" to describe a procedure by which adipose tissue was removed with liposuction and selectively injected to correct irregularities in other areas [12].

Conventionally, percutaneous autologous fat transfer relocates harvested fat deep into subcutaneous tissue using a syringe and large-core needle, or cannula attached to a suction device. Advantages include ease of performance and supplementation, minimal induction of immune-mediated inflammation, relatively low cost, avoidance of secondary defects, rapid recovery, and minimal risk. Major drawbacks are reabsorption, which can be lessened by new techniques and overcome by repeated treatments, and bruising and redness at the recipient site, due mainly to puncture wounds [8,10].

In the 1990s, Coleman [3] further refined the harvesting, purification, and injection techniques. The use of small-caliber cannulas (3–4 mm), so threads of transplanted adipose tissue do not exceed 3 mm in diameter, reduces fat necrosis and improves fat survival. Recent reports describe transfer of injected fat particles as small as 3 mm in diameter, reduces fat necrosis and improves fat survival. These advantages have been associated with greater viability of transposed autologous fat.

Previous studies described the use of autologous fat injection grafting for cosmetic facial augmentation [14,15], but these studies did not involve an intra-oral approach. We identified one report of intra-oral fat injection for correction of lip hemangiomas [16]. A common criticism of intra-oral fat transfer is the belief that oral injection is associated with a greater risk of infection than use of the percutaneous approach. However, the precise incidence of infection using a trans-oral or transmucosal route of injection has not been reported in the literature to support this claim. We did not observe an increased rate of infection using this technique, mucosal puncture wounds healed uneventfully, recovery was rapid, and swelling, bruising, redness, and scarring at the recipient site were negligible inside the mouth. These findings are consistent with use of the transmucosal route in many other surgeries, such as synthetic malar and chin implants, and maxillary and mandibular osteotomies with plate and wire fixation.

The majority of the patients we encountered tolerated the procedure well, and 46% chose additional augmentation 3 to 6 months later, using fat banked and frozen during the first session. This duration of fat stability is consistent with reports of extra-oral fat grafting [9,17]. The decision to receive additional reinjections after the initial fat augmentation was based on the patient’s own subjective assessment and preferences as to the amount of augmentation desired and was independent of the initial injection site. As with the conventional technique, long-term persistence of augmentation depended on the initial over-correction. Clinically applicable objective methods for assessing the longevity of fat injections are lacking. While some studies use magnetic resonance imaging or other formal imaging techniques, this is often not practical or cost-effective in clinical practice. We judged the duration of fat stability based on physical exam and photographs taken at each visit, similar to previous studies [2,14].

Although this is a small case series and more formal controlled studies are necessary, our study demonstrates that transmucosal, intra-oral delivery of autologous fat for proximate peri-oral facial augmentation can reduce visible scarring and swelling at the recipient site. Furthermore, this technique was associated with minimal risk of infection. Transmucosal intra-oral fat transfer should be considered an alternative to percutaneous transfer for peri-oral augmentation.

**PATIENT CONSENT**

Patients provided written consent for the use of their images.

**REFERENCES**

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