Lipodissolve for Body Sculpting  
Safety, Effectiveness, and Patient Satisfaction

Efficacy and Safety of a Low-Molecular Weight Hyaluronic Acid Topical Gel in the Treatment of Facial Seborrheic Dermatitis

Osteopathic Manipulative Treatment  
Novel Application to Dermatological Disease

Invasive Ductal Breast Carcinoma  
Underneath a Lipoma in a Male Patient

Lymphocytic Thrombophilic Arteritis  
Induced by Minocycline

Sister Mary Joseph Nodule as a Presenting Sign of Pancreatobiliary Adenocarcinoma

Acute Painful Nodules in a Young Healthy Adult
Lipodissolve for Body Sculpting
Safety, Effectiveness, and Patient Satisfaction

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ABSTRACT
Lipodissolve, to reduce superficial deposits of fat, has gained popularity in recent years. A simple solution of phosphatidylcholine in deoxycholate evolved around 2004 and has been used by two collaborating physicians in Minnesota. Their experience encompassing 1,616 patients receiving a total of 15,122 treatments is described. Relatively modest volumes of injections produced satisfactory and smooth results in 74.5 to 86.5 percent of the patients in the two practices. No serious complications developed. Minor and rare side effects included pain, lightheadedness, tender nodules, pigmentation, and ulceration in two patients. The authors offer useful tips to enhance safety, effectiveness, and patient satisfaction with the procedure. (J Clin Aesthet Dermatol. 2012;5(10):16–19.)

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with 3 to 6cc. Ultrasound, for a few minutes, was added before or after injections for the possibility of improved adipocyte lysis.11,12

RESULTS

All patients had some stinging and pain during, and for 30 minutes after, the procedure, which was reduced among the patients who received 4 to 5 minutes of icing before the procedure. A variable amount of swelling and bruising developed in almost all patients, but mostly subsided within 10 days. The overall results differed somewhat depending upon the volume of solution with each treatment (Table 1).

Practice A, using higher injection volume per treatment needed fewer treatments, 3 versus 4 per area, than practice B. The satisfaction rate was slightly better among those receiving higher injection volume, 86.6 percent in practice A versus 74.4 percent in practice B. When separated for body sculpting and cellulite, in practice B, the satisfaction rates were 81 and 44 percent, respectively. When response could be easily measured, such as abdominal circumference reduction in 95 patients, it varied from 0.125 to 3 inches per treatment, with an average of 1.1 inch. The fat reduction was generally associated with some degree of skin tightening, resulting in a smooth appearance. No patient developed lumpy, bumpy, or uneven appearance, as seen in some patients after liposuction. Complications were minor and extremely rare and developed in 1 to 2 percent of the patients (Table 1). Lightheadedness from hyperventilation, pain, and prolonged tender nodules were slightly more common in patients receiving a higher volume of injections (practice A). Complications of liposuction, such as pulmonary embolism, hemorrhage, perforation, lidocaine/epinephrine toxicity, third space fluid shifts, and fat embolism,13,14 did not occur with lipodissolve.

DISCUSSION

These results, in conformity to others reported in the literature (cited above), reveal that lipodissolve is a highly effective procedure in reducing unwanted deposits of localized fat and that it can tighten the skin to some extent, providing smooth cosmetic results (Figures 1–4).

This is an observational report, not a prospective study, so institutional review board approval was not indicated. Because different areas of the body would show different amounts of shrinkage (e.g., abdomen vs. arm) and because the amount of fat being treated is highly variable in different individuals, the authors felt that they could not attach significance to circumferential reduction in centimeters to the entire group, but they have listed the measured reduction in 100 treated patients. In evaluating abdominal circumference reduction, the authors did see a statistically significant diminution (see Table 2). The most important parameter to the authors for this report was patient satisfaction with the procedure. The procedure should be preferred over liposuction for small-to-moderate areas of superficial fat to avoid possible complications of liposuction. Fat embolism, one of the complications of liposuction, has never been reported with lipodissolve. To the contrary, phosphatidylcholine has been used to treat fat embolism, such as with bone fractures.15 There has been some conjecture and/or fear that lipodissolve may increase cholesterol. However, phosphatidylcholine has been shown to improve lipids and reduce cardiovascular risk factors.16–18

Large volumes of PC/DC injections, 100cc per area as
suggested a few years ago, do not seem to be necessary. A high degree of patient satisfaction can be achieved with as little as 40 to 60cc per treatment. More treatments may be needed with lower volumes, but as the authors’ experience reveals, these treatments are better tolerated and with a lesser chance of complications.

Based on personal experience and review of the literature, the authors offer the following tips to enhance safety, effectiveness, and patient satisfaction with the procedure.

SAFETY

- Avoid pregnant patients and patients with any systemic disease, such as heart disease, kidney disease, uncontrolled diabetes or hypothyroidism, infections, active or previous autoimmune disease, or active skin disorders.
- Avoid patients on aspirin or other anti-inflammatory drugs and those with a known bleeding tendency.
- Avoid patients with soy allergy (the usual PC/DC mixture is soy based).
- Avoid injecting breast or axillary tail of the breast. Prolonged inflammation could possibly stimulate malignant cells.
- Avoid injecting around the knees (too close to ligaments) and below the knee, as these areas may be more prone to skin breakdown.
- Inject the minimum volume necessary depending upon the area and amount of fat, usually 40 to 60cc.
- Injections should be 1.5cm apart and in the middle of the fat pad. Avoid injecting too close to the skin or underlying muscle and fascia.
- Avoid rubbing the area or wearing garments that are too tight.

![Figure 1. Abdominal treatments, before and after (4 treatments)](image)

![Figure 2. Bra/back fat treatments, before and after (4 treatments)](image)

![Figure 3. Flank treatments, before and after (4 treatments)](image)

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<th>TABLE 2. Reduction in inches</th>
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<td><strong>MEANS WITH THE SAME LETTER ARE NOT SIGNIFICANTLY DIFFERENT</strong></td>
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<td><strong>DUNCAN GROUPING</strong></td>
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• Patient should avoid putting on lotions immediately after treatment to avoid infection. Wait for a few days until puncture sites have healed.

EFFECTIVENESS
• Select patients with relatively localized areas of soft fat. Firm and fibrous fat tends not to respond as well.
• In patients with cellulite or skin laxity, inject somewhat more superficially, but still at a depth of 5mm, at least. Avoid patients with excessive skin laxity with minimal underlying fat.
• Palpate and mark the areas to be treated exactly. For example, some patients have a contiguous fat pad around the umbilicus, others have a large area below the umbilicus or both below and above the umbilicus, and others have multiple areas over the abdomen.
• Ultrasound immediately before or after the procedure may help fat cell lysis.
• Wait for 6 to 8 weeks before re-injecting the same area to have full effect of the first treatment.

PATIENT SATISFACTION
• Avoid patients with undue expectations, such as obese or overweight patients who feel this may lead to weight loss and really thin patients who imagine they have extra fat.
• Go over informed consent and the possibility of even rare complications. Inform them that there is a possibility of prolonged palpable nodules and pigmentation in a small number of patients, and pain that may last for three or more days.
• Inform the patient that it may take 3 to 4 treatments per area to have the desired effect. However, if there is absolutely no effect six weeks after the first treatment, it is helpful to have a frank discussion with the patient and better to avoid further treatment if the patient is skeptical.

REFERENCES