Lipedema Can Be Treated Non-Surgically: A Report of 5 Cases

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Case series

Patients: Femal, 39-year-old • Femal, 64-year-old • Femal, 52-year-old • Femal, 34-year-old • Femal, 90-year-old

Final Diagnosis: Lipedema

Symptoms: Edema • pain

Medication: —

Clinical Procedure: Medications

Specialty: Endocrinology and Metabolic

Objective: Unusual clinical course

Background: Lipedema is a chronic progressive disease characterized by the abnormal accumulation of fat in the subcutaneous region. Both medical and surgical treatments have been described in international guidelines; however, the current literature is biased toward promoting liposuction as the primary treatment of lipedema, and this can lead to the misapprehension that liposuction is the only form of definitive treatment.

Case Reports: In the present study, we report 5 cases at various stages of the evolution of lipedema, all with different therapeutic objectives. Case 1 reported having persistent bruising and pain, case 2 reported pain and fat deposition, case 3 reported night cramps and discomfort, case 4 reported leg thickening, and case 5 reported redness in the legs. All of were diagnosed with lipedema in different evolution stages. Our purpose was to demonstrate the possibility of non-surgical therapy, as well as to improve signs and symptoms of lipedema, using the QuASiL questionnaire and measuring changes in volumes and proportions. Good aesthetic outcomes improve both social and psychological status.

Conclusions: Currently, there are many described therapies available for lipedema. Liposuction surgery for lipedema should be considered one possible tool. Treatment objectives can be different for each patient. It is imperative to understand each patient’s needs in order to offer the best therapy attainable that meets patient requirements and induces a better quality of life. Non-surgical treatment of lipedema is feasible in selected cases, and it can meet the criteria for achieving selected clinical objectives.

Keywords: Lipedema • Lymphedema • Obesity • Obesity Management

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Background

Lipedema is an inherited, chronic, progressive disease characterized by the abnormal accumulation of fat in subcutaneous tissue, mainly in the lower and upper limbs [1]. It is a potentially aggressive disease because it limits mobility and damages the lymphatic vascular system [2], leading to deformities and loss of quality of life. Its prevalence in females is believed to be 11-39% [3-6]. The intensity of presentation is greatest in the inflammatory phase, manifesting as pain [7], sensitivity to touch, edema [8], chronic fatigue, and unprovoked ecchymosis [5,9].

Lipedema is often misdiagnosed as obesity, lymphedema, or chronic venous disease, although these diagnoses can often occur concomitantly [1,10,11]. Lipedema is not influenced substantially by diet or exercise [12]. In the absence of targeted treatment, there is no significant improvement in body disproportions or in signs and symptoms. Signs of lipedema are disproportionately increased limb volume, adipose tissue on the limbs, symmetric tissue, palpable tissue nodules, painful tissue (not always), and limb swelling (pitting or non-pitting), and the hands and feet not affected [13].

Clinical and surgical treatments for lipedema have been described extensively in international guidelines. The aim of all of these is to improve the signs and symptoms, to reduce volumes and disproportions of the affected limbs, and to prevent progression [14-18]. Manual lymphatic drainage and elastic compression have been reported to alleviate symptoms [5]. Diuretics are not indicated, as there is no accumulation of liquid. However, there is thickening of adipose tissue and there is no standard pharmacological protocol to treat this [19]. Diet has been presented as an adjuvant therapy [3], however, like physical exercise, it has been little studied [19]. Most often the most commonly used diet strategies are anti-inflammatory, low-carbohydrate, and ketogenic diets [20]. The current literature is biased in the sense that it presents liposuction as the main treatment for lipedema [3], and this can lead to the misapprehension that it is its only definitive treatment.

The treatment of lipedema must be individualized and must consider the technical limitations and clinical aspects of each patient. In the present study, we describe a series of 5 cases of lipedema in various stages (Figure 1 [21]), all of which were successfully treated clinically.

Stage I lipedema is characterized by normal skin with enlarged hypodermis. Stage II is defined as uneven skin texture with pleats of fat and large piles of tissue growing like unencapsulated masses. Stage III is characterized by hardening and thickening of the subcutaneous with large nodules and protrusion of cushions/accumulations of fat, especially in the thighs and around the knees. Finally, stage IV is associated with lymphedema, and is also called lipolymphedema.

Figure 1. Classification of lipedema by evolutionary stages [21]. Stage 1: normal skin with an enlarged hypodermis; Stage 2: unevenness of the skin texture with pleats of fat and large piles of tissue growing like unencapsulated masses; Stage 3: hardening and thickening of the subcutaneous with large nodules and protrusion of cushions/accumulations of fat, especially in the thighs and around the knees; Stage 4: lipedema with lymphedema, so-called lipolymphedema.
Case Reports

Case 1, a Patient in Stage 1

The patient was a 39-year-old woman with persistent ‘bruising’ and pain that worsened with standing. Edema of her lower limbs worsened in the afternoon. She also reported ‘cellulite’ and fat on her legs that started 5 years prior to presentation, when she started taking contraceptives, in addition to finding her legs ‘disproportionately thick’. On the Lipedema Symptom Assessment Questionnaire (QuASiL: minimum 0, maximum 150 [22]) she presented 89 points. Superficial and deep-color Doppler ultrasound examinations of the lower limbs (EDCV-MMII) revealed only sparse reticular structures. Lymphoscintigraphy was within normal limits.

On past surgical history, she reported having undergone ‘hydrolipo’ 3 years prior and varicose vein surgery 5 years prior.

Physical examination revealed sparse reticular veins, absence of Godet and Stemmer signs, and pain on palpation of areas with greater fat deposition. The bioimpedance test (Tanita®, BC-601, Illinois, United States) showed a body mass index (BMI) of 21.2 kg/cm² with 29.8% body fat and lower-limb volume of 19 668.55 mL.

A diagnosis of grade I lipedema was made and clinical treatment was suggested based on the aforementioned guidelines. These included anti-inflammatory dietary measures, regular aquatic physical exercise, manual lymphatic drainage, and antioxidant phytotherapeutics in regular-usage doses. The patient returned with symptom improvement, with a score of 81 in QuASiL (improvement of 8.9%) and volumetric decrease of 491.62 mL. After 8 months of clinical treatment, she achieved 58 points on the QuASiL (improvement of 34.83%) and total volume loss of 1230.54 mL. There was significant improvement in her aesthetic concerns of gynecoid lipodystrophy (Figure 2). Her final BMI was 21.4 kg/cm².

Case 2, a Patient in Stage 1

The patient was a 64-year-old woman who presented with pain and fat in her legs since adolescence. On physical examination, there were fatty nodules in both lateral malleoli that were painful to palpation; her QuASiL score was 44. Stage 1 lipedema was diagnosed, and clinical treatment was provided with anti-inflammatory dietary measures, aquatic physical exercise, manual lymphatic drainage, and antioxidant herbal medicines. After 1 month of treatment, she reported significant improvement in the nodules (Figure 3) and symptoms, scoring 11 points on the QuASiL.

Figure 2. Patient with stage 1 lipedema before (A) and after (B) treatment, showing significant aesthetic improvement of gynecoid lipodystrophy (cellulitis), although it was not the main objective of the treatment.
Case 3, a Patient in Stage 2

A 52-year-old woman presented with a 7-month history of ankle edema. She reported discomfort in her calf, with a sensation that she could feel her veins, as well as sporadic night cramps and the appearance of spider veins in her right thigh. She had been taking chlortalidone and phlebotonics. On physical examination, she had telangiectasias, severe pain on palpation of the medial pre-tibial and genicular fat, absence of Godet or Stemmer signs, and 103 points on the QuASIL. EDCV-MMII revealed bilateral varicose tributaries, without reflux in saphenous veins or the deep venous system. The bioimpedance exam showed a BMI of 29.5 kg/cm$^2$, 38.6% body fat, and lower-limb volume of 23,737.71 mL. Stage 2 lipedema was diagnosed.

Testing for IgG hypersensitivity to food was requested, which identified yogurt, cow’s milk, cashews, poppy seeds, and mushrooms as the strongest triggers of hypersensitivity.

Clinical treatment was initiated, including changes in lifestyle habits, combined with an anti-inflammatory diet and progression to a low-carbohydrate diet associated with aquactic exercise, manual lymphatic drainage, and antioxidant herbal medicines. After 1 month, she returned with 69 points on the QuASIL (33% improvement) and volume loss of 968.96 mL. After 5 months of clinical treatment, she scored 22 points on the QuASIL (78.64% improvement), with total limb volume of 21,817.51 mL (a difference of 1920.2 mL) (Figure 4).

Case 4, a Patient in Stage 3

The patient was a 34-year-old woman with longstanding ‘stubborn constitution’ and weight gain after menarche. She reported that, during adolescence, her legs thickened despite her being very active. Dieting caused her to lose truncal fat, without changes in the lower limbs.

She reported a maximum weight of 95 kg, which reduced with exercise to a minimum of 65 kg, but without decreasing the volume of the legs, which remained the same.
Physical examination revealed absence of Godet and Stemmer signs, deposition of fat that was tender to palpation in the lower limbs, and 115 points on the QuASiL. EDCV-MMII revealed small varicosities in the thighs and legs, without significant reflux, with a measured dermal thickness of 22.4 mm and 21.2 mm in the right and left pre-tibial regions, respectively.

The bioimpedance exam showed a BMI of 34.2 kg/cm$^2$, 41.6% body fat, and lower-limb volume calculated by bioimpedance of 32,572.21 mL. She was diagnosed with grade III lipedema. Although surgical treatment with tumescent liposuction was indicated, the patient initially opted for clinical treatment that included an anti-inflammatory diet followed by a ketogenic diet associated with aquatic physical exercise, manual lymphatic drainage, and antioxidant herbal medicines. After 1 month of treatment, she returned with 107 points on the QuASiL (6.9% improvement), and a volume loss of 3972.44 mL. After 3 months, she scored 86 points on the QuASiL (25.21% improvement), with a volumetric loss of 6472.47 mL. After 6 months, she achieved 76 points on the QuASiL (33.91% improvement), with volumetric loss of 6,346.34 mL. After 11 months, she achieved 59 points in QuASiL (48.6% improvement) with final volume of 21,832.31 mL (volumetric loss 10,739.90 mL) (Figures 5, 6).

Case 5, a patient in stage 4

A 90-year-old woman presented with ‘redness in the leg’ of 1-year duration, with no improvement following unspecified treatments. She reported that, since adolescence, her legs had been slowly growing in volume and she had never managed to prevent progression. Two years prior to presentation, orthopedic surgical treatment of the knees was indicated after significant weight loss. She reported loss of mobility, which progressed to inability to walk. She was unable to lose weight despite changes in eating habits. She scored 75 points on the QuASiL. On physical exam she presented with redness, lymphangitis, erysipelas, severe columnar form, and painful symmetric fat tissue. Pitting edema over her feet suggested associated lymphedema, and the bilateral edema also suggested that the right lymphangitis was not the only cause of lymphedema (Figure 7). EDCV-MMII revealed only small superficial varicosities, while pre-tibial skin and subcutaneous thickness of 57.27 mm left and 55.9 mm right confirmed lipedema [23]. Lymphoscintigraphy was not requested because of infection. After starting clinical measures to treat lipedema (anti-inflammatory diet, moderate exercise, antioxidants and lymphatic drainage) and lymphedema (lymphatic drainage) associated with the treatment of erysipelas (amoxicillin-clavulanate...
and enoxaparin), there was quick resolution of erysipelas and stasis dermatitis.

Discussion

There is a need to define the objective of the treatment for lipedema in cooperation with the patient. These objectives can vary widely: improvement of signs and symptoms, weight loss, fat, reduction of limb volume, and improvement of mobility and aesthetics. Often, the patient’s aesthetic desire unduly overrides the objective need to improve mobility and symptoms, and to prevent progression. Using objective clinical tools for analysis and monitoring of lipedema, such as the QuASiL [22], symptom monitoring questionnaires that are validated and culturally adapted, and volumetric measurements of limbs and bioimpedance, it is possible to quantify clinical progression. BMI does not differentiate obesity from lipedema. Nevertheless, we have no tool to measure the aesthetic aspects of the disease.

The top priority in the treatment of lipedema, regardless of type, is the recovery and preservation of mobility [3]. Medial contact of the thighs when walking, which already occurs in the early stages of the disease, is difficult to quantify; however, it can be considered the beginning of mobility limitation.

Secondarily, the objective is to improve symptoms, as assessed by the QuaSiL questionnaire, in addition to volumetric and proportional improvements. Finally, the aesthetic improvement that is eventually achieved improves psychological status.

In the first case, it can be argued that lipedema had been treated surgically by previous liposuction; however, the return of lipedema symptoms demonstrated the need for clinical treatment even after surgical treatment. The prior varicose vein surgery highlights comorbidities and overlap of symptoms. She
experienced volumetric, symptomatic, and aesthetic improvements. In the second case, symptoms and fat nodules showed significant improvement. In the third case, in addition to the symptomatic improvement, there was significant volume loss. In the fourth patient, who had grade I obesity on the basis of BMI, there was disproportion and substantial fat deposition in the lower limbs. In this case, lipedema mimicked obesity; nevertheless, there was significant volumetric decrease in addition to symptomatic improvement. In the fifth case, the initial objective was the treatment of refractory erysipelas; nevertheless, clinical treatment facilitated the resolution of the persistent infection.

The clinical treatment of lipedema involves, in most aspects, the search for a healthy lifestyle, including changes in attitudes. These measures are mostly harmless to those who do not have lipedema and can be suggested to all patients who do not have contraindications. Although compression bandages or garments are widely recognized treatment, they were not used for the present patients because of high pain patients reported before lowering inflammation. We opted for lymphatic drainage as a strategy the patients would find more acceptable. Hesperidin and diosmin are antioxidant treatments currently suggested for lipedema treatment [13], and use of quercetin, pycnogenol [24], flavonoids [25], rutosides [26], and butcher’s broom [27] was previously reported in treatment of lipedema and/or lymphedema. It was suggested that a combined sequence of natural complex antioxidants sets up a reductive cascade to match the progressive oxidative stresses of the free-radical cascade [28].

In patients with lipedema with comorbid diseases such as varicose veins of the lower limbs (53% of cases) or obesity (50% of cases) [29], failure in the clinical treatment of lipedema may cause failure in the treatment of the associated disease [30]. The high prevalence of the disease and co-presentation with other common diseases, as well as its numerous clinical variations, justify the consideration of clinical treatment as initial and supporting treatments for patients with lipedema.

Surgical treatment should not be performed in the inflammatory phase of the disease [10], and the best time for decision-making is when the patient has reached the best symptomatic state. Undoubtedly, the advent of liposuction that protects the lymphatic system brought new perspectives and hope in the treatment of this disease [31]; nevertheless, liposuction does not exclude non-surgical therapy.

Figure 7. Patient with lipolymphedema, stage 4 lipedema, with erysipelas that was difficult to treat before (A) and after (B) non-surgical treatment for lipedema.
This series of cases exemplifies and contextualizes lipedema, which until recently had few therapeutic options in the daily practice of vascular surgeons. Our study has several limitations inherent to a series of cases: not all aspects of lipedema were addressed, and patients were selected to exemplify chosen aspects.

Conclusions

Currently, there are many described therapies available for lipedema. Liposuction surgery for lipedema should be considered a possible tool to be used and not the only available treatment. Treatment objectives should be individualized for each patient. It is imperative to understand each patient’s needs to offer the best therapy attainable that meets patient requirements and induces a better quality of life. Some patients may demand better aesthetics, which in some cases is reachable without surgery. Most patients report pain and discomfort, symptoms that can improve without surgery. Other patients may be concerned about leg volume. Volume and disproportion reduction may seem challenging without surgery, but we have shown it is possible. Complications of lipedema, like ulcers, lymphangitis, and erysipelas, can be the main patient concern and can be treated without liposuction. Because of all available therapies, lipedema requires an integrated multispecialty and multidisciplinary teamwork patient-centered approach.

Non-surgical treatment of lipedema is feasible in selected cases, and it can meet the criteria for achieving selected clinical objectives.

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