INTRODUCTION

Multiple symmetric lipomatosis (MSL) is a very rare lipid metabolism disorder characterized by the accumulation of multiple, symmetric, nonencapsulated lipomatous masses on the face, neck, shoulders, and upper arms [1]. The disorder was first described by Brodie in 1846. Madelung then reported the cases of 33 patients in 1888, and Launois and Bensaude provided the standard description of MSL in 1898. Thus, MSL is also known as Madelung’s disease or Launois-Bensaude syndrome [2]. The cause of the condition is unknown, but a number of theories have been proposed. The incidence rate is higher in men than in women, and the average age of incidence is approximately 60 years; the condition is known to be linked to alcoholism [1,2].

Herein, we report a case of MSL in a 66-year-old man and review the current literature on the condition.

CASE REPORT

A 66-year-old man presented with bilateral breast bulging. His breasts had slowly become enlarged over a number of years and this had led to huge breast bulging; the patient reported that he was ashamed of his breast disfigurement when in public baths and swimming pools. He had undergone an excisional biopsy of a mass in the central upper back region 1 year earlier, which revealed a lipoma. He reported more than 25 years of alcohol abuse, and confirmed that he consumed alcohol 5 times a week. Furthermore, he had a history of cigarette use, but had stopped smoking 4 years previously. He also had a 10-year history of hypertension and diabetes mellitus. The patient’s father had a history of multiple large disfiguring masses in the neck and upper body.

Physical examination of the patient revealed multiple large, disfiguring masses in the occipital region as well as on the breasts, upper arms, shoulders, neck, and upper trunk. These appeared to be symmetrical on the breasts, upper arms, and shoulders. Palpation revealed that the masses had an elastic consistency, could be moved without causing the patient any pain, and had no definite margins (Figure 1A).

Laboratory examinations revealed normal liver and thyroid function, and a hormone assay confirmed that the results were within the normal ranges. A mammography was per-
formed which revealed that the patient’s breasts were comprised almost entirely of fat. Breast ultrasonography revealed excessively fatty breasts without any definite glandular tissue (Figure 1B). Computed tomography and magnetic resonance imaging showed diffuse and marked fatty deposits in the superficial soft tissue layers of the neck, chest, abdomen, and shoulder wall with marked bulging contours of both breasts (Figure 2).

The patient underwent bilateral nipple-sparing mastectomy with nipple interposition. Pathology revealed soft tissue lipomas measuring 30 × 17 × 5 cm and 27 × 16 × 5 cm, and both lipomas were homogenous without hemorrhage and necrosis (Figures 3, 4). No hidden malignancy was found. The patient did well postoperatively and was satisfied with the cosmetic result.

Figure 1. Physical findings and imaging studies of the patient. (A) Breast masses had elastic consistency and could be moved without causing any pain. Mammography (B) and ultrasonography (C) shows bilateral excessive fatty breast without definite glandular tissue (pseudogynecomastia).

Figure 2. Computed tomography (CT) and magnetic resonance imaging (MRI) findings of the patient. (A) Marked bulging contours of both breasts with fat depositions are observed on CT scan. (B) MRI shows fat deposition in occipital lesion (white arrow). (C) CT scan shows bilateral fat deposition in shoulder (white arrow). (D) Excessive focal fat deposition as another focus of lipoma is also observed in superficial soft tissue layer of upper central abdomen on CT scan (white arrow).
DISCUSSION

MSL is a rare disease with an incidence of approximately 1 in 25,000 men. The condition is more common in men, especially those aged 30 to 60 years; the male to female ratio is approximately 15:1 to 30:1. Its incidence is known to be associated with a history of alcoholic abuse. The etiology of MSL remains unknown, however, several theories have been suggested, such as mitochondrial disorder related to genetic dysfunction, defective embryogenic brown adipose tissue, and a lack of adrenergic-induced lipolysis [1-5]. MSL presents as large painless growing adipose masses with little circumscription that are usually located in the neck region, on the shoulders and upper extremities, and in the upper trunk region. Pathologically, MSL shows characteristics of normal adipose tissue and accumulation of fat in the brown adipose tissue [1-3]. MSL may occur sporadically or be inherited (autosomal dominant transmission). Sporadic cases are more frequently observed and are strongly related to alcoholic abuse and tobacco smoking. In the case reported here, the family history supports a hereditary origin of the disease [1,2].

Enzi [6] classified MSL into two types according to its distribution site. Type I is horse collar lipoma, which presents as circumscribed protruding masses in the neck area as well as the clavicular and deltoid regions. In severe cases, this type can cause mediastinal extension and tracheal obstruction. Type II is characterized by a pseudoathletic appearance with diffuse involvements of the trunk and the proximal end of the extremities, resembling obesity [1,3]. Donhauser et al. [4] added gynecoid type to the categories of the disease and classified MSL as follows: type I, horse collar lipoma; type II, pseudoathletic appearance; and type III, gynecoid type.

There was one interesting difference in the case presented here compared with previous reports. The presenting symptom of MSL is somewhat different according to gender: men who presented with condition usually present with a submental mass [2,5,7,8]. However, the patient in our case presented mainly with large bilateral breast masses. Due to this extraordinary clinical presentation, we first had to exclude simple gynecomastia by taking a through medical history and per-

Figure 3. Postoperative physical finding and breast specimen. (A) The patient underwent both nipple-sparing mastectomy with nipple interposition. (B) Breast specimen shows multiple lipomas, which were measured 30 × 17 × 5 cm (right) and 27 × 16 × 5 cm (left), respectively.

Figure 4. Microscopic pictures of specimen. Diffuse proliferation of normal adipose tissue is observed in microscopic pictures of specimen (H&E stain, × 100).
forming a physical examination.

MSL is a benign disease and its malignant transformation is very rare. Malignant degeneration of fatty tissue into liposarcoma has only been reported in two cases [9,10]. Despite the benign nature of MSL, it can present a serious clinical scenario. In the early period of the disease, it grows rapidly and thereafter, shows slow progression or remains stable for a long period of time. The long-term deposition of lipomatous tissue often results in a cosmetic problem, and more seriously it can cause obstructive symptoms such as dysphagia, dysphonia, or dyspnea [2,3,5]. In some cases, neurological symptoms, which are related to mitochondrial defect, maybe present [11].

Differential diagnoses of MSL are simple obesity, Cushing syndrome, angiolipoma, neurofibromatosis, salivary gland diseases, conditions with a neck mass such as thyroid goiter, Frohlich’s syndrome, Dercum’s disease, and lipomatosis in human immunodeficiency virus [1,2]. For the accurate diagnosis of MSL, taking the clinical history of patients and performing physical examinations are very important. In the case presented here, the patient presented with large bilateral breast masses, which had slowly become enlarged over a number of years. He also had a history of a lipomatous mass on his central upper back and 25 years of alcohol abuse. Physical examination revealed multiple large, disfiguring masses in the occipital region, upper arms, shoulders, and upper trunk. Of note, the masses on the arms and shoulders were located symmetrically. We were able to exclude the possibility of simple gynecomastia and diagnose MSL through careful history taking and physical examination.

Surgery is known to be the most effective treatment for MSL [7]. Alcohol withdrawal and weight reduction are recommended, however, these cannot reverse or stop the course of the disease [12]. Ultrasound-guided liposuction can also achieve good effects. However, liposuction is of limited effectiveness with large masses [13]. The overall recurrence rate following surgery is approximately 63%, open surgery results in a recurrence rate of 51% compared to a recurrence rate of 95% with liposuction [7].

In conclusion, MSL can present as gynecomastia. Increasing awareness of the clinical characteristics of MSL, especially among breast surgeons, could facilitate its recognition. Surgery is known to be the most effective treatment for MSL. In addition, because MSL is a genetic disorder, family members should be followed up to allow for early detection and appropriate surgical treatment, thus avoiding a delayed diagnosis and subsequent unsatisfactory treatment outcome accompanying a series of further operations.

CONFLICT OF INTEREST

The authors declare that they have no competing interests.

REFERENCES

1. Ardeleanu V, Chicos S, Georgescu C, Tutunaru D. Multiple benign symmetric lipomatosis: a differential diagnosis of obesity. Chirurgia (Bucur) 2013;108:580-3.
2. Meningaud JP, Pitak-Arnnop P, Bertrand JC. Multiple symmetric lipomatosis: case report and review of the literature. J Oral Maxillofac Surg 2007;65:1365-9.
3. Lopez-Ceres A, Aguilar-Lizarralde Y, Villalobos Sánchez A, Prieto Sánchez E, Valiente Álvarez A. Benign symmetric lipomatosis of the tongue in Madelung’s disease. J Craniofac Surg 2006;17:489-93.
4. Donhauser G, Vieuf D, Ruzicka T, Braun-Falco O. Benign symmetric Launois-Bensaude type III lipomatosis and Bureau-Barrière syndrome. Hautarzt 1991;42:311-4.
5. Mevrio E, Sbrocca M, Mullace M, Viggione S, Mevio N. Multiple symmetric lipomatosis: a review of 3 cases. Case Rep Otolaryngol 2012;2012:910526.
6. Enzi G. Multiple symmetric lipomatosis: a revised clinical report. Medicine (Baltimore) 1984;63:56-64.
7. Brea-Garcia B, Cameselle-Teijeiro J, Couto-González I, Taboada-Suárez A, González-Alvarez E. Madelung’s disease: comorbidities, fatty mass distribution, and response to treatment of 22 patients. Aesthetic Plast Surg 2013;37:409-16.
8. Busetto L, Sträter D, Enzi G, Coin A, Sergi G, Inelmen EM, et al. Differential clinical expression of multiple symmetric lipomatosis in men and women. Int J Obes Relat Metab Disord 2003;27:1419-22.
9. Borriello M, Lucidi A, Carbone A, Iannone V, Ferrandina G. Malignant transformation of Madelung’s disease in a patient with a coincidental diagnosis of breast cancer: a case report. Diagn Pathol 2012;7:116.
10. Tizian C, Berger A, Vykoupil KE. Malignant degeneration in Madelung’s disease (benign lipomatosis of the neck): case report. Br J Plast Surg 1983;36:187-9.
11. Plummer C, Spring PJ, Marotta R, Chin J, Taylor G, Sharpe D, et al. Multiple symmetrical lipomatosis: a mitochondrial disorder of brown fat. Mitochondrion 2013;13:269-76.
12. Heike Z, Gudrun UM, Frank RD, Vetter H, Walger P. Multiple benign symmetric lipomatosis: a differential diagnosis of obesity: is there a rationale for fibrate treatment? Obes Surg 2008;18:240-2.
13. Verhelle NA, Nizet JL, Van den Hof B, Guelinckx P,Heymans O. Liposuction in benign symmetric lipomatosis: sense or senseless? Aesthetic Plast Surg 2003;27:319-21.