

Treatment of Idiopathic Granulomatous Mastitis: Combined Intralesional Steroid Injection with Topical Steroid Administration vs. Systemic (Oral) Steroid

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Abstract

Background

Idiopathic granulomatous mastitis (IGM) is characterized by non-caseating granuloma and microabscess formation limited to mammary gland lobules. It is a form of chronic mastitis of unknown pathogenesis. In this study, we aimed to compare the effectiveness of intralesional steroid injection with topical steroids versus systemic steroid therapy in the treatment of IGM.

Methods

Between June 2017 and December 2020, the patients were collected and assessed. IGM was diagnosed histopathologically by tru-cut biopsy in patients with breast mass, pain, and erythema with suspicion of IGM. Included in the study were one hundred and eleven patients who were diagnosed with IGM and who were treated with local or oral administration of corticosteroids, with at least 6 months of follow-up. The patients were divided into 2 groups: a local corticosteroid treatment group (LC, group 1, n=57); and a peroral corticosteroid treatment group (OC, group 2, n=54). Demographic characteristics, treatment responses, recurrence rates, side effects of the steroid, and the need for surgery were compared.

Results

While the rate of smoking was 12.3% in the LC group, was 20.4% in the OC group. There was no history of oral contraceptive use in either group of patients. Previous steroid use was significantly lower in the LC group (10.5%) compared to the OC group (55.6%) ($p = 0.001$). Previous antibiotic use was significantly lower in the LC group (75.4%) compared to the OC group (100%) ($p = 0.001$). When complete and partial responders were grouped as "responders," 96.5% in the LC group and 75.9% in the OC group responded after the first course of treatment ($p=0.001$). 98.2% in the LC group and 87.0% in the OC group were complete responders after the third course of treatment ($p=0.003$). While 7% of the patients in the LC group had recurrence, 37% in the OC group had recurrence ($p=0.001$). Steroid-related side effects were lower in the LC group (no) compared to the OC group (11.1%) ($p=0.010$). Surgery was performed in 3.5% of the LC group, while surgery was performed in 57.3% of the OC group ($p=0.001$).

Conclusion

Steroid injection is an IGM treatment with proven efficacy, short response time, low recurrence rate, and lower need surgery. The treatment is easy to administer and minimizes the risk of systemic side effects. Even in clinically severe cases, we believe that steroid injection might be used as monotherapy, or in combination with other therapies, to improve treatment efficacy and create a transition to less aggressive treatments.

Introduction

Idiopathic granulomatous mastitis (IGM) is characterized by non-caseating granuloma and microabscess formation limited to mammary gland lobules. It is a form of benign chronic mastitis of unknown pathogenesis. Kessler and Wolloch first described this disease in 1972 [1, 2]. IGM is usually observed as a persistent or recurrent tender and inflammatory mass in the breast in premenopausal women with a history of breastfeeding. Although the etiology of IGM is still not well known, many different developmental mechanisms have been suggested in the literature. While the autoimmune response is the most accepted theory, infection and hormonal disruption are considered the three leading hypotheses [3]. Because IGM is clinically similar to breast cancer and inflammatory breast diseases, IGM needs to be differentiated from breast cancer and granulomatous inflammation or autoimmune diseases such as tuberculosis,

sarcoidosis, fungal/parasitic infection, and Wegener's granulomatosis [4]. Epidemiologically, IGM is rare and accounts for approximately 0.44-1.6% of breast biopsy specimens. IGM is more common in Asian and Spanish women, and it is mostly seen in childbearing age [5]. Misdiagnosis of IGM is common, especially in the beginning, and it can turn into a frustrating long-term illness for the patient and doctor. There is still no consensus on the treatment of IGM. Treatment includes many options, such as surgical resection, steroids, methotrexate, bromocriptine, colchicine, immunosuppressive agents, antibiotics and even follow-up observation [6]. Currently, treatment mostly depends on the experience of the clinic or surgeon and the severity of the patient's disease [7].

In this study, we compared the efficacy of local steroid therapy that is combined intralesional steroid injection with topical steroid administration versus systemic steroid therapy in the treatment of IGM.

Materials And Methods

The data collected between June 2017 and December 2020, was analyzed retrospectively. Patients who presented with breast mass, breast pain, mastitis, and suspected idiopathic granulomatous mastitis underwent core needle abscess drainage and incisional biopsy, or excisional biopsy, according to their clinical presentation. The diagnosis of IGM was made histopathologically. Infectious granulomatous mastitis agents were excluded with Gram, Periodic Acid-Schiff (PAS), Ziehl-Neelsen (EZN) and Giemsa Stains in the histopathological diagnosis of each patient. Included in the study were patients diagnosed with IGM who were treated with oral or local administration of corticosteroids, with at least 6 months of follow-up. Patients with recurrent disease or a contraindication for the use of corticosteroids, patient has breast abscess, and patients who were pregnant or lactating were excluded from the study. Extracted from patients' files were their age, number of pregnancies, history of smoking, complaint at disease onset, radiologic method preferred for the diagnosis and follow-up [such as breast ultrasound (USG), mammography (MMG) and breast magnetic resonance imaging (MRI) depending on the patients age and clinical status], and the biopsy method used for diagnosis. No specific patient selection was made for the two groups, the patients were selected consecutively. We didn't calculate sample size. The patients were selected consecutively according to the order they applied to the clinic. In our clinic, we have tried to treat almost all patients with IGM with LC for the last 2 years. Most of the patients in the OC group were patients who were treated before 2 years. No choice was made regarding the extent of the disease between the two groups.

Treatment protocol

There were 2 patient groups: a local corticosteroid treatment group (LC, group 1, n=57) and a peroral corticosteroid treatment group (OC, group 2, n=54). LC protocol was an intralesional triamcinolone acetonide which a medium-to-strong potency corticosteroid (Kenacort-A amp. 40mg/ml, Deva, Istanbul, Turkey) injection in the lesions once a month, under the USG guided (Figure 1) and topical administration of triamcinolone acetonide 0.1% pomade (Kenacort-A %0.1, 20gr pomad, Deva, Istanbul, Turkey) on the skin of the affected breast area twice a day on alternate days for one month. We diluted 1 amp of Keacort-A with 3-5 cc saline to the part of the lesion, depending on the size of the lesion. Injection was made at different points at 5 cm interval in patients with diffuse disease. Injections were made separately. In patients with multiple lesions if groups of lesions were more than 5 cm apart and if the individual lesions were less than 5 apart, the larger group was injected. OC protocol was administered as methylprednisolone 32mg/day, with each tablet containing 16 mg methylprednisolone. (Prednol 16 mg tb, Deva, Istanbul, Turkey). Before the treatment, the patient was informed about the treatments. however, 4 patients were excluded from treatment because they could not follow up due to treatment non-compliance. While About 10% of the LC patients and about half of the OC patients were treated in other centers. After this treatment patients applied to our clinic for biopsy or surgery. LC treatment lasted for 1 month. We diluted 1 amp of the drug into 3-5 cc and injected it into the area where the

disease was. The patients felt pain in the injection, but this pain was not intolerable. It was done just like a normal injection. Also, we have rescue medications as a methotrexate, but we don't need to use it.

One month later, the response of the patients was evaluated with clinical and radiological ultrasonography in cases who completed their treatment. Patients were classified depending on resolution of the disease clinically and radiologically as "complete response," "partial response," "no response," or "worsening disease." Patients with "complete response" were consigned to follow-up, while patients with "partial response," "no response," and "worsening disease" received a second and if necessary, a third course of treatment with monthly check-ups, until the complete resolution of the disease. If there was no significant improvement in the disease despite the treatments in one patient, we considered this as a persistent disease. If a patient had complete resolution after treatment, after a while mastitis seen again, we defined that the recurrence of the disease. The reason for the long follow-up period in the OC group was these patients were treated in the first years. We started the follow-up period from the time the patients' treatment ended. Information about the course of the disease available treatment options, and their rate of success were given all patients at their initial and follow-up exams. Surgical treatment was performed for patients who no response or worsening disease, and preferred surgery during follow-up. In our clinic, we do not perform surgical treatment on almost any of the patients with IGM. We have been trying to treat it completely with LC in the last 2 years.

The various treatment modalities were low or single box OC, various antibiotics, surgery such as lumpectomy and abscess drainage, and follow-up. Each "course" of OC treatment lasted for 1 month and was taper-off the steroid regimen by reducing the steroid dose. LC treatment was not applied to the group in which OC was used, and vice versa. There is only one patient who underwent lumpectomy in the LC group, and that patient was also a patient who did not respond to treatment. In fact, it had a medical indication. Although we wanted to continue LC treatment, the patient used the choice from surgery. He had used antibiotherapy for a long time and did not provide any benefit. We considered the patient's choice reasonable. No surgery was performed without medical indication. All patients who underwent surgery had medical indications.

Statistical Analysis

In calculating the sample width of the study, Power (the power of the test) of at least 80% for each variable and Type 1 error were determined by taking 5%. Whether the continuous measurements in the study were normally distributed or not was examined by Kolmogorov-Smirnov ($n > 50$) and Skewness-Kurtosis. Tests and parametric tests were applied because the measurements were normally distributed. Descriptive statistics for continuous variables in the study; mean and standard deviation; and categorical variables were expressed as numbers (n) and percentages (%). The "Independent T-test" was used to compare "continuous measures" according to Prednol Treatment Groups. Chi-square test was used to determine the relationship between Prednol Treatment Groups and categorical variables. Statistical significance level was taken as 5% in the calculations and SPSS (IBM SPSS for Windows, ver.25) statistical package program was used for analysis.

Results

One hundred and eleven female patients diagnosed with IGM were included in the study. 57 patients received local corticosteroid (LC) therapy, and 54 patients received oral corticosteroid (OC) therapy. Patients' mean age was 32.7 ± 7.08 in the LC and 36.04 ± 8.42 in the OC ($p = 0.014$). The localization of lesions was similar between two groups ($p = 0.224$). The main complaint at onset presented with all three signs of disease as a breast mass with pain and skin erythema in two groups ($p = 0.112$). Duration until treatment was 10.98 ± 21.08 weeks in LC and 4.17 ± 2.08 weeks in OC ($p = 0.020$). The reason for the high duration of complaints in the LC group was the receipt of unsuccessful treatment, such as antibiotics, steroids, or other medications, or they are having undergone surgeries such as abscess drainage

and lumpectomy in other hospitals. At the beginning, we performed tru-cut biopsy and started treatment early. The mean number of children was 2.62 ± 1.58 in the LC and 3.28 ± 2.10 in the OC ($p=0.066$). Total breast-feeding time was 41.52 ± 30.47 months in the LC and 48.41 ± 30.78 months in the OC ($p=0.245$). Smoking ratio was similar between the two groups ($p=0.248$). Patients and their clinical characteristics are shown in Table 1.

Table 1
Patients and their clinical characteristics

		Local Corticosteroid (LC, n=57)		Oral Corticosteroid (OC, n=54)		<i>p.</i>
		N	%	N	%	
Localization of lesions	Right	23	40.4%	28	51.9%	<i>.224</i>
	Left	34	59.6%	26	48.1%	
Main complaint at onset	Mass+Pain	19	33.3%	26	48.1%	<i>.112</i>
	Mass+pain+Erythema	38	66.7%	28	51.9%	
Smoking	Yes	7	12.3%	11	20.4%	<i>.248</i>
	No	50	87.7%	43	79.6%	
Previously received corticosteroid treatment	Yes	6	10.5%	30	55.6%	<i>.001</i>
	No	51	89.5%	24	44.4%	
Previously received antibiotic treatment	Yes	43	75.4%	54	100.0%	<i>.001</i>
	No	14	24.6%	0	0.0%	
Tissue sampling method	Core needle	53	93.0%	38	70.4%	<i>.002</i>
	Incisional	4	7.0%	10	18.5%	
	Excisional	0	0.0%	6	11.1%	
Recurrence	Yes	4	7.0%	20	37.0%	<i>.001</i>
	No	53	93.0%	34	63.0%	
Steroid-related side effects	Yes	0	0.0%	6	11.1%	<i>.010</i>
	No	57	100.0%	48	88.9%	
Surgery	Abscess drainage	1	1.75%	26	48.1%	<i>.001</i>
	Lumpectomy	1	1.75%	5	9.2%	
Age (Mean±SD)		32.7±7.08		36.04±8.42		<i>.014</i>
Duration until treatment (Mean±SD)		10.98±21.08		4.17±2.08		<i>.020</i>
Number of children (Mean±SD)		2.62±1.58		3.28±2.10		<i>.066</i>
Breast-feeding time (Mean±SD)		41.52±30.47		48.41±30.78		<i>.245</i>
Follow-up (Mean±SD)		21.68±9.99		41.81±18.11		

While 10.5% (6 patients) of the patients in the LC group had previously received corticosteroid treatment, 55.6% (30 patients) in the OC group had previously received corticosteroid treatment ($p=0.001$). While 75.4% (43 patients) of the patients in the LC group had received antibiotic treatment, 100% (54 patients) in the OC group had received antibiotic treatment initially ($p=0.001$). This is due to the low response rate to oral steroids and starting antibiotics without a definitive diagnosis, making second and third doses necessary. Histopathological diagnosis was made by tru-cut biopsy in 93% (53 patients) of the patients in the LC group, while the diagnosis was made by tru-cut biopsy in 70.4% (38 patients) in the OC group ($p=0.002$). In addition, 18.5% (10 patients) in the OC group were treated with incisional biopsy and 11.1% (6 patients) with excisional biopsy (Table 1). The reason for the high rate of tru-cut biopsy in the LC group was that the diagnosis and treatment of these patients were made in our center, which is a university hospital.

In the LC group, two patients with a complete response after the first course of treatment received a second course of treatment due to recurrence. Similarly, a third course of treatment was given to two patients in the LC group who developed recurrence after the second course of treatment. In the LC group, lumpectomy was performed in one patient who did not achieve a complete response after the second course of treatment. We performed oncoplastic surgery on our own patients and their breast appearance was not so bad. The patient in Fig. 2 had surgery with the bad cosmetic result of surgery in another center. In the OC group, lumpectomy was performed in five patients who did not achieve a complete response. While there were no steroid-related side effects in the LC group, 11.1% (6 patients) of the OC group had steroid-related side effects such as steroid-induced hirsutism, acne, weight gain, diffuse bone pain, and muscle weakness ($p=0.010$). These patients used steroids and these side effects were of course steroid related. These patients were also consulted to the endocrine clinic and followed up with them. The follow-up period for the patients was 21.68 ± 9.99 in the LC group and 41.81 ± 18.11 in the OC group. (Table 1).

While 7% (4 patients) of the patients in the LC group had recurrence, 37% (20 patients) in the OC group had recurrence ($p=0.001$). Surgery was performed in 3.5% (2 patients) in the LC group, while surgery was performed in 57.3% (31 patients) in the OC group ($p=0.001$). Abscess drainage was performed in 1.75% (1 patient), and lumpectomy was performed in 1.75% (1 patient) in the LC group, whereas abscess drainage was performed in 48.1% (26 patients) and lumpectomy in 9.2% (5 patients) in the OC group. The reason for the high number of surgical interventions in the OC group is the lower response rates to corticosteroid treatment and different treatment protocols (antibiotic, low dose oral corticosteroid) are used by other centers (Table 1).

In the first course of treatment, 28.1% in the LC group (14 partial response, 2 no response) did not achieve complete response, but 70.4% did not achieve complete response in the OC group (25 partial response, 12 no response and 1 worsening disease) ($p=0.001$). In the second course of treatment, 12.3% (7 partial response) in the LC group and 22.2% in the OC group (10 partial response, 2 no response) did not achieve complete response ($p=0.058$). In the third course of treatment, 1.75% (1 partial response) in the LC group and 13.0% in OC group (5 partial response, 2 no response) did not achieve complete response ($p=0.004$) (Table 2).

Table 2
Response rates of patients in treatment arms

	1st administration		2nd administration		3rd administration	
	Local Corticosteroid (n=57)	Oral Corticosteroid (n=54)	Local Corticosteroid (n=18)	Oral Corticosteroid (n=37)	Local Corticosteroid (n=7)	Oral Corticosteroid (n=13)
Complete response	41(71.9%)	16(29.6%)	11(19.2%)	25(46.3%)	6(10.5%)	6(11.1%)
Partial response	14(24.6%)	25(46.3%)	7(12.3%)	10(18.5%)	1(1.75%)	5(9.3%)
No response	2(3.5%)	12(22.2%)	0	2(3.7%)	0	2(3.7%)
Worsening disease	0	1(1.9%)	0	0	0	0

When patients with complete and partial response were pooled in a single group to determine the rate of “responders,” 96.5% (55 patients) in the LC group and 75.9% (41 patients) in the OC group were responders after the first course of treatment ($p=0.001$); 91.1% (52 patients) in the LC group and 75.9% (41 patients) in the OC group were complete responders after the second course of treatment; and 98.2% (56 patients) in the LC group and 87.0% (47 patients) in the OC group were complete responders after the third course of treatment ($p=0.003$), (Figure 3a,b and 4a,b and 5a,b). The fourth dose of treatment was administered to one patient who was stable and then continued with partial responses in the LC group (Table 2).

Accordingly, previous steroid use was significantly lower in the LC group (10.5%) compared to OC group (55.6%) ($p=0.001$). Previous antibiotic use was significantly lower in the LC group (75.4%) compared to the OC group (100%) ($p=0.001$). Treatment response rate was significantly higher in the LC group than in the OC group ($p=0.001$). The recurrence rate was significantly lower in the LC group (7%) compared to the OC group (37%) ($p=0.001$). Steroid-related side effects were lower in the LC group (none) compared to the OC group (11.1%) ($p=0.010$). The need for surgical treatment was significantly lower in the LC group (3.5%) compared to the OC group (57.3%) ($p=0.001$).

Discussion

Since Kessler and Wolloch defined IGM in 1972, despite time and developing technology and treatment modalities, uncertainty in the physiopathology of IGM and debates on its treatment continue [2, 3]. Although autoimmunity is the cause most supported by the literature, various etiological factors such as various hormonal disorders (hyperprolactinemia, etc.), oral contraceptive use, autoimmune factors, microorganisms, smoking, and alpha 1-antitrypsin deficiency have been blamed [8–10]. The fact that the physiopathology of IGM is not well known has hampered effective treatment. [11]. In addition to surgical and non-surgical treatment modalities, the use of local steroids has recently been discussed in the literature. In this study, we aimed to discuss cases with IGM treated with local and systemic steroids in light of current literature.

The rate of smoking and oral contraceptive use, which are etiological factors, was 0-23.7% and 15.3%-52.6%, respectively [3, 12–15]. In our study, while smoking was 12.3% in the LC group, it was 20.4% in the OC group. There was no history of oral contraceptive use in either group of our patients. In this context, the effects of smoking and oral contraceptive use on the pathophysiology of the disease are questionable.

Although most patients receive antibiotic treatment before the definitive diagnosis because of diagnostic uncertainty, routine antibiotic use is not recommended given that IGM is a sterile condition [3, 16]. Oral steroid and antibiotic use rates were very high in the OC group. The lack of high treatment responses has led to the reuse of these medications. High doses and long-term use of these medications cause some complications and increase the cost of treatment. The fact that these were lower in the LC group (steroid use;10.5%, antibiotic use;75.4%) indicates the effectiveness of the local steroid treatment and a more effective professional approach to the disease. Similarly, the tru-cut biopsy rates are high in the LC group.

Although one of the most used methods in the treatment of IGM is steroid therapy and surgery, there are many options such as follow-up, antibiotics, non-steroidal anti-inflammatory drugs, colchicine, methotrexate, azathioprine, imuran, and mycophenolate mofetil. And there is a current study about local steroid use [3, 11–13, 15, 17, 18]. There are opinions that wide local excision is the most ideal treatment. However, surgical excision may have a high recurrence rate, poor wound healing, fistulization, and cosmetic problems [19]. In recent years, medical treatments have been more preferred in the treatment of IGM [15]. Corticosteroids have been defined as first-line therapy because of their positive results [12]. On the other hand, steroids are known to cause side effects that affect all systems such as blood sugar irregularity (steroid-induced diabetes mellitus), weight gain, Cushing's syndrome, and body weight gain, and as a result they require a change in treatment algorithm [11, 15, 20–22]. Topical corticosteroids are used in the treatment of many skin diseases, especially atopic dermatitis, due to their anti-inflammatory, vasoconstrictive, antiproliferative and immunosuppressive effects. While steroids can be easily absorbed through normal skin, their absorption at the site of inflammation is increased [18, 23].

Topical steroid use was first described in an IGM case by Altintoprak *et al.* in 2011. They saw a significant improvement in the patient's clinic after 5 weeks of treatment [18]. In their next studies, they treated 28 patients with only topical steroid for an average of 8.2 weeks (range 4–12 weeks) and noted that the long-term efficacy of treatment was more than 90% [11]. After that Munot *et al.* applied local steroids to the breast lesion in 4 patients and noted clinical and radiological improvement in all patients [24]. In our previous study, in which we treated patients with uncomplicated IGM with local steroids, we showed that there was 93.5% improvement in the lesion in the first month after the first dose. The rate of patients with complete response after the third dose was 93.4% [3].

When we look at the treatment durations of different studies in the literature; Lai *et al.* [25] found that the duration of treatment in the observation treatment group was 14.5 months. Cetinkaya *et al.* [15] in their study, found that the duration of improvement in the observation group was 5.6 months (1.3-13.8) and 3.9 months (0.8-9.8) in the systemic steroid treatment group. Azizi *et al.* [12] described the recovery period of the observation group as 9 months in their study. By 15 months of treatment with MTX, 75% achieved disease remission, and 13-15 (range 1-30) months were median duration of treatment [13]. In a study by Montazer *et al.* [26] improvement was seen after 12 months in 93.3% of patients in the group treated with high-dose steroids (50 mg daily), while 53.3% of patients in the low-dose steroid group were able to improve after 6 months [27]. In their study in which they performed steroid injection, Alper *et al.* showed that 89.3% of the patients had complete recovery in their mean follow-up of 11.8 (5-20) months. In our study, 98.2% (56 patients) in the LC group and 87.0% (47 patients) in the OC group were complete responders after three months ($p=0.001$). When comparing our study with the literature, we see that the treatment time in the LC group was significantly shorter. In other words, we see that we can reach a high cure rate in a shorter time.

Equally as difficult as treatment in patients with IGM is the recurrence of the disease. Although it varies according to the type of treatment in the literature, the recurrence rate in IGM is reported to be between 5-50%. The recurrence rate in this study was 24.8% [12]. Montazer *et al.* [26] found no significant difference in the recurrence rate between the conservative and surgical treatment groups. Cetinkaya *et al.* [15] found that recurrence rates were 14.3% in the

observation group, 42.9% in the systemic corticosteroid group, and 21.4% in the antibiotic group. Postolova *et al.* [13] found that recurrence rate was 50% after surgery. They found that 15.8% of patients had recurrence during their treatment with MTX. In our study, while 7% (4 patients) of the patients in the LC group had recurrence, 37% (20 patients) in the OC group had recurrence ($p=0.001$). The literature indicates that the recurrence rate was significantly lower according to the use of systemic.

The greatest restriction on the use of the steroid is undoubtedly its side effects. Cetinkaya *et al.* [15] reported in their study that the treatment was not successful in more than half of the patients they treated with steroids and that the patients had problems due to various side effects. In another study involving treatment with MTX, side effects were reported in 15.7% of patients [13]. Yin *et al.* [2] stated in their article that the side effects of topical, systemic, and combined steroid treatments were 2.4%, 38.2% and 30.3%, respectively. Munot *et al.* [24] reported they had no side effects in any of the 4 patients they treated with steroid injection [27]. In a study in which they used local steroids, Alper *et al.* found that none of the 28 patients with IGM observed steroid-related side effects. In our previous study, which was multicentric, the rate of steroid-related side effects was 2.2% in the local steroid group, while it was 9.4% in the systemic steroid group [3]. In our study, there were no steroid-related side effects in the LC group and 11.1% (6 patients) in the OC group.

Wide surgical excision may be required to prevent recurrence; however, it causes large defects in the breast, resulting in negative cosmetic outcomes and psychological burden [2]. Chirappapa *et al.* [28] reported in their cohort study that surgical resection offered no advantage over steroids in terms of healing time, and even 52% (13/25) of patients had wound complications. Shin *et al.* [29] reported that surgical resection may result in significant breast deformity, extensive scarring, and other complications. In our study, the patients' need for surgery was 3.5% in the LC group and 59.3% in the OC group ($p=0.001$).

Conclusion

Steroid injection is a treatment that is easy to administer and minimizes the risk of systemic side effects. It carries with it a proven efficacy, short response time, and low recurrence rate. Even in clinically severe cases, we believe that steroid injection might be used as monotherapy, or in combination with other therapies, to improve treatment efficacy and create a transition to less aggressive treatments. Prospective randomized studies investigating the effect of different doses of local steroid injections, long-term responses, and side effects should be carried out to aid in the development of future treatment algorithms for patients with IGM.

Each treatment modality has advantages and disadvantages. Considering the superiority of local steroid use to other treatments, it should be considered a primary treatment modality.

Declarations

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Statement of Ethics

This study was approved by the local ethics committee of Van Yüzüncü Yıl University, Turkey. (Reg. No. 2021/06-03).

Informed consent was not received due to the retrospective registry nature of the study

Declaration of conflicting interests

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Author Contribution

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Figures

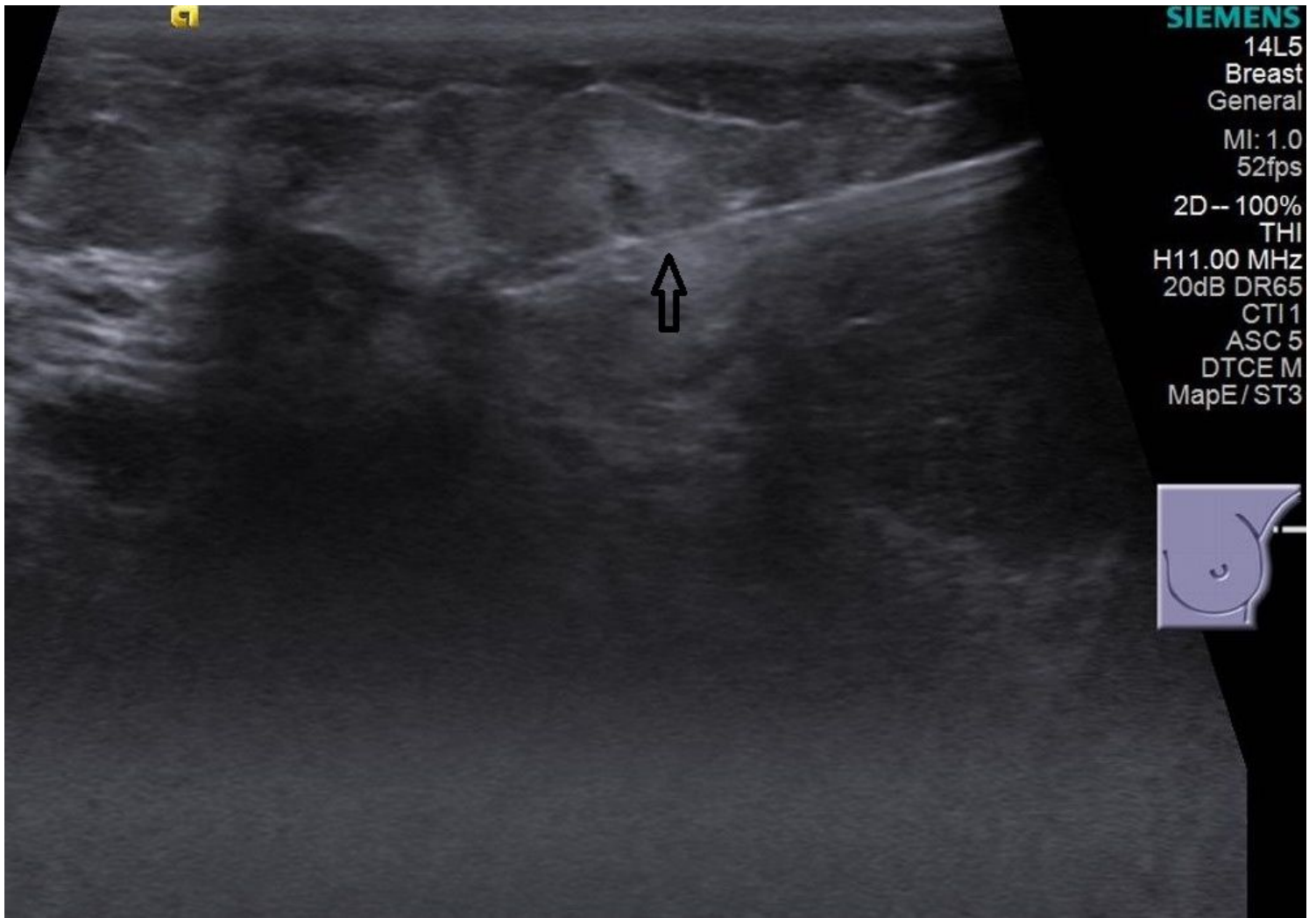


Figure 1

Ultrasound-guided steroid injection (Thin arrow is the needle in the center of IGM lesion).



Figure 2

Poor cosmetic result after surgery.



Figure 3

a/b: Follow-up of the patient who received steroid injection; a) Before treatment, b) After 3rd dose.



Figure 4

a/b: Follow-up of another patient who received steroid injection; a) Before treatment, b) After 2nd dose.



Figure 5

a/b: Follow-up of the patient who received oral steroid; a) Before treatment, b) After 3rd dose.