European Federation of Societies for Ultrasound in Medicine and Biology (EFSUMB) Position Statement on Dermatologic Ultrasound

Stellungnahme der European Federation of Societies for Ultrasound in Medicine and Biology (EFSUMB) zu Dermatologischem Ultraschall

Authors
Fernando Alfageme1, Ximena Wortsman2, Orlando Catalano3, Gaston Roustan1, Maria Crisan4, Diana Crisan5, Diana E. Gaitini6, Eugenio Cerezo7, Radu Badea8

Affiliations
1 Dermatology, Hospital Universitario Puerta De Hierro Majadahonda, Madrid, Spain
2 Department of Dermatology, Universidad de Chile, Santiago de Chile
3 Radiology, Institut Pascale, Naples, Italy
4 Dermatology, University of Medicine and Pharmacy Iuliu Hatieganu, Cluj-Napoca, Romania
5 Dermatology, Universitätsklinikum Ulm Klinik für Dermatologie und Allergologie, Ulm, Germany
6 Radiology, Rambam Medical Center, Haifa, Israel
7 Ultrasound, Clinica DKV, Madrid, Spain
8 Regional Institute of Gastroenterology and Hepatology, University of Medicine and Pharmacy, "Iuliu Hatieganu", Cluj-Napoca, Romania

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Correspondence
Dr. Fernando Alfageme
Dermatology, Hospital Universitario Puerta De Hierro Majadahonda, Manuel de Falla 2, 28022 Madrid, Spain
Tel.: ++34/6 95 54 72 62
dermalfgame@gmail.com

ABSTRACT
Dermatologic ultrasound is a recent application of ultrasound for the evaluation of healthy skin and appendages and their diseases. Although the scientific literature regarding this application is still not sufficient for evidence-based guidelines, general recommendations issued by scientific societies are necessary. The EFSUMB (European Federation of Societies for Ultrasound in Medicine and Biology) steering committee for dermatologic ultrasound has developed a series of consensus position statements regarding the main fields of dermatologic ultrasound (technical requirement, normal skin and appendages, inflammatory skin diseases, tumoral skin diseases, aesthetic dermatology and practice-training requirements). This document is the foundation for future evidence-based recommendations and guidelines for dermatologic ultrasound practice.

ZUSAMMENFASSUNG
Introduction

Dermatologic ultrasound (DERMUS) is an application of ultrasonography (US) in the study of the normal and diseased state of the skin and appendages (nails and hair) [1]. As it is a growing, recently developed application, guidelines and recommendations based on scientific evidence are not methodologically possible. However, a position statement from a scientific society with regards to this application of US is useful for physicians involved in dermatologic US, allowing for the foundations of present clinical practice and evidence generation to be developed [2, 3].

Methodological structure and classification of the consensus levels

The executive board of the European Federation of Societies for Ultrasound in Medicine and Biology (EFSUMB) designated a dermatologic ultrasound steering committee based on qualifications including relevant publications, clinical experience and absence of conflict of interest. The Policy Document Development Strategy for Clinical Practice Guidelines, Position Statements and Technological Reviews of the EFSUMB was adhered to throughout the process for this position statement [4].

The main topics regarding dermatologic US were selected by the steering group and a comprehensive scientific literature search was performed to identify relevant studies.

Recommendations were elaborated by steering committee members and a consensus meeting for expert evaluation of these recommendations was convened at the EUROSON 2019 Congress (Granada, Spain).

A position statement was approved if > 75 % of voting members were in agreement (broad agreement: > 75–95 % of votes, strong consensus > 95 % of the votes). For discussion a nominal group technique was applied [4]. In the case of disagreement (≤ 50 % of the votes or less in favor) or if the rephrased or alternative position statement again failed to gain > 75 % of votes, the position statement was removed. A lack of consensus on this particular issue would be recorded in the text as recommended by the EFSUMB policy document [4].

1. Technical requirements for dermatologic ultrasound

The main technological advance that has made dermatologic US possible is the introduction of high-frequency and very high-frequency transducers with enough spatial resolution to study the superficial structures of the skin and appendages [5, 6].

According to the DERMUS group (an international group of experts in dermatologic ultrasound), dermatologic US for the skin and appendages should be performed using a linear multiple frequency transducer (30–70 MHz) as the minimum standard [2]. Newer very high-frequency (> 20 MHz) and ultra-high-frequency transducers (30–70 MHz) allow exploration of small adnexal structures, such as sebaceous glands and apocrine and eccrine glands [7].

POSITION STATEMENT 1

Operators performing a dermatology ultrasound study should be aware of patient history and clinical findings. A detailed request from the referring clinician should be available (strong agreement 9/9, 100 %).

POSITION STATEMENT 2

The minimum transducer frequency for dermatologic ultrasound should be 15 MHz. Higher transducer frequencies may provide further information that may be relevant (broad agreement 8/9, 88.9 %).

Both gel hips and gel pads, which make it possible to separate the epidermis from the transducer, should be used for accurate epidermal evaluation and avoidance of superficial vascular plexus compression [8, 9].

Also according to the DERMUS group, all US examinations should include color, power or spectral Doppler US to ascertain the presence of a vascular anomaly, providing fundamental information regarding inflammation and neovascularization [2, 10].

In order to detect superficial dermal, subdermal, small vessels, the pulse repetition frequency (PRF) should be adjusted accordingly and the gain should be adjusted to reduce flare artifacts [11]. Proper training in color and spectral Doppler US should be included in dermatologic US training programs [3] (see training section).

Trapezoid field-of-view (FOV) and extended FOV facilities are useful in the evaluation of large or deep lesions. Three-dimensional reconstruction software and new non-Doppler facilities for microvascularity assessment are useful and should be employed, if available on the ultrasound machine [12].

Current experience regarding dermatology application of elastography is limited, and contrast-enhanced ultrasound (CEUS) has not been deployed in dermatology US practice [13, 14].

POSITION STATEMENT 3

Color Doppler/power Doppler and pulsed spectral Doppler (in the case of vascular anomalies) are recommended to establish inflammatory state of skin and appendages and the presence or neovascularization (strong agreement 9/9, 100 %).

2. Ultrasound of normal skin and appendage

Anatomical and histological structures of skin and appendages present differential echogenicity that must be known as changes may indicate pathology [11, 15].
2.1 Ultrasound of normal skin
Ultrasound of the skin is basically a correlation of the different layers (epidermis, dermis and subcutaneous tissue) with the acoustic interphases they produce [16].
- The epidermis is a hyperechoic line due to its keratin content,
- The dermis is a band inferior to the epidermal line which is less hyperechoic than the dermis, due to the rich collagen content
- Subcutaneous fat is a complex structure composed of hyperechoic collagen septa and hypoechoic fatty lobules

2.2 Ultrasound of hair tracts and follicles
A hair tract (the visible part of hair) is the final keratinized product of hair follicles, which are oblique cellular structures in the dermis [17, 18].
- A hair tract is a bilaminar or trilaminar lineal hyperechoic structure depending on the presence of medulla (terminal follicles) [19]
- Hair follicles are oblique hypoechoic structures that may vary in depth from subcutis to upper dermis depending on the phase of growth (early anagen follicles are deeper than superficial catagen follicles)

2.3 Ultrasound of the normal nail apparatus
The nail apparatus has a very close relationship with the distal interphalangeal joint.
Four elements must be assessed [20, 21]
- Nail plate: Bilaminar hyperechoic structure due to the keratin content
- Nail bed: Between nail plate and cortex of the distal phalanx
- Distal phalanx: Hyperechoic lineal interphase under the nail-bed
- Nail Matrix: Ill-defined hypoechoic area that surrounds the proximal nail plate area.

2.4 Ultrasound of skin tumors
Ultrasound examination of the skin is widely used for the assessment of cutaneous tumors [22]. Even though histological examination remains the “gold standard” for the diagnosis of oncologic skin pathology [23], US can help guide preoperative diagnosis and improve both oncological and aesthetic outcomes. Furthermore, US plays an important role not only in the therapeutic outcome after surgical management of skin tumors, but also in the follow-up process [24].

Modern US technology can be successfully and effectively used in the field of skin oncology, offering particular descriptive information which should be integrated in the overall examination to achieve a complex evaluation of the nature of the investigated tumor pathology. Conventional B-mode provides significant morphological data related to the tumoral lateral and depth extension, relationship to neighboring tissues (involvement of muscle, cartilage, bone), contour, echogenicity and echo-structure [14, 22]. The tumor thickness (Breslow histological index) is one of the most important prognosis factors and establishes the therapeutic strategy especially in cases of melanoma. The evidence indicates that there is a significant correlation between the sonographic “depth index” and the histological Breslow index [22, 23].

3. Ultrasound of skin tumors
Ultrasound examination of the skin is widely used for the assessment of cutaneous tumors [22]. Even though histological examination remains the “gold standard” for the diagnosis of oncologic skin pathology [23], US can help guide preoperative diagnosis and improve both oncological and aesthetic outcomes. Furthermore, US plays an important role not only in the therapeutic outcome after surgical management of skin tumors, but also in the follow-up process [24].

Doppler, contrast-enhanced ultrasound and elastography in skin tumors
A color Doppler US examination combined with spectral Doppler US can reveal macro-circulation at the tumor bed level. The vascular features that may suggest a malignant lesion include hypervascularization, disorganized blood flow model with peripheral or mixed distribution, increased blood velocity and multiple vascular pedicles [23]. The lack of evidence of supply vessels in certain cases, however, reveals the limitations of Doppler US examination, which can be overcome by CEUS, which allows for assessment of the microcirculation. Analysis of the time-intensity curves during CEUS examination suggests that the vascular dynamics of the tumor depend on vascular resistance, shunts, histological type and location. The disposition of the tumor vessels, the uptake pattern of the contrast agent, the blood flow velocity are parameters that may suggest the malignant or benign nature of tumors. According to published data, malignant tumors display an inhomogeneous uptake pattern on a CEUS examination with a significantly higher value for the washout time. Compared to benign tu-
mors, CEUS allows analysis of tumoral vascularization from the early arterial to the late venous phase, emphasizing maximal uptake, wash-up time, duration of passage and distribution pattern in the area of interest [14, 23, 24].

Strain elastography is also a useful examination in the diagnosis of skin cancer. The qualitative elastography appearance is significantly associated with semi-quantitative elastography (strain ratio) measurements. Any type of elastography provides information about tissue stiffness. According to the literature, malignant tumors display a high or medium increase in stiffness. The stiffness positively correlates with the thickness of the tumors. The role of elastography in skin tumors has not been subject to active research. As malignant tumors are stiffer than benign ones, elastography added to B-mode and color Doppler US examination has the potential to improve the accuracy of traditional clinical diagnosis [25, 26].

**POSITION STATEMENT 8**
The use of dermatologic ultrasound as an examination for palpable skin tumors is recommended, since it offers valuable information regarding tumoral extension, vascularization, delimitation and degree of compressibility for optimized treatment (broad agreement 7/9, 77.8%).

### 3.1 Ultrasonographic aspects of non-melanoma and melanoma skin cancer

**Basal cell carcinoma**
Basal cell carcinoma (BCC) displays US features such as the presence of hyperechoic ‘dots’ or punctiform hyperechoic areas within the lesion (which correspond to calcium deposits, keratinized cells or prominent basal cell aggregates), anechoic or hyperechoic areas (suggestive for cysts, mucin deposits), the presence of two or more vascular pedicles [23]. The blood flow is more prominent at the inferior aspect of the lesion. Tumor margins are difficult to assess in cases of morpheaform and infiltrative BCC, although in most cases BCCs tend to have well-defined borders [28, 29]. The presence of cutaneous elastosis can occur when the measurement of the lesion difficult [24]. Furthermore, the hyperechoic ‘dots’ within the tumor, determined by the presence of keratin or basal cell nests, can be of use in differentiating a basal cell carcinoma from melanoma and even different histologic varieties of BCC [29, 30].

**Squamous cell carcinoma**
Squamous cell carcinoma (SCC) can present perpendicular shadows determined by superficial scales or crusts. Hyperkeratosis and the presence of abundant inflammatory infiltrate may lead to an overestimation of tumor size [22]. SCCs frequently invade deeper structures. Due to the risk of the development of metastasis, loco-regional lymph nodes should also be evaluated [31]. Regarding the microcirculation pattern, SCCs usually present with two or more vascular pedicles and have a mixed or peripheral intratumoral circulatory pattern [23, 24].

**Melanoma**
Malignant melanoma (MM) usually presents as homogeneous or inhomogeneous, hypoechogenic, with an irregular contour, intense chaotic vascularization (mostly arterial vessels) [32], two or more vascular pedicles, increased echogenicity of the underlying subcutaneous tissue, increased or moderate stiffness on elastography [33]. When a melanoma is suspected, it is important to check for satellite lesions, in-transit lesions and lymph node metastasis [33, 34].

**Other skin malignancies**
Skin lymphoma, angiosarcoma, Kaposi sarcoma, adnexal carcinoma and other non-melanoma skin cancer (NMSC) can also be assessed using US [35–37]. Cutaneous lymphoma may appear either as a nodular or diffuse mass. The nodular mass appears solid hypoechogenic and is poorly defined. The diffuse form appears as a hyperechogenic, poorly defined area with increased thickness of the subepithelial layers. The thickness or degree of infiltration of cutaneous lymphoma can also be assessed by US [37]. Moreover, the changes in skin infiltration following therapy can be evaluated by B-mode US examination and elastography. An angiosarcoma can also have a characteristic malignant appearance on US examination, appearing as a hypoechogenic, poorly defined lesion with intense vascularization that infiltrates deeper structures [38]. Kaposi sarcoma is seen as a hypoechoic, intensely vascularized plaque or nodule [35]. As the biology, growth rate and metastasis risk of malignant skin tumors are diverse, sonographic follow-up should be done according to the normally accepted clinical practice for skin cancer.

**POSITION STATEMENT 9**
Ultrasound is useful for dermatological tumor follow-up. The frequency and length of follow-up are based on each tumor type (broad agreement 8/9, 88.9%).

### 4. Inflammatory skin diseases
Although inflammation of the skin and appendages can usually be assessed by clinical visual inspection or palpation, some deep processes are difficult to evaluate as they involve deeper cutaneous structures such as deep dermis and subcutaneous tissue [1, 8, 39]. Moreover, skin sclerosing diseases (i.e., morphea, scleroderma, chronic graft versus host disease) have episodes of inflammatory change, which are difficult to evaluate, and will require a different therapeutic approach [40]. Therefore, the role of US in these clinical scenarios is to add useful information to the clinical visual exploration for more accurate patient assessment and staging [41].
4.1 General inflammatory ultrasound signs in skin diseases

Although there is an immense variety of inflammatory diseases of the skin and appendages, common shared US findings in inflammatory skin disorders include [39]:

- Hypoechogenic areas in the subepidermal portion of the dermis
- Increased local blood flow demonstrated by color Doppler US
- Hypoechogenic septa and hyperechoic fatty lobules when the subcutaneous tissue is affected

These features help to determine the level and extent of the changes in inflammatory disorders of the skin, hair, and nails [41].

**POSITION STATEMENT 10**

With inflammatory diseases of the skin and appendages, the level and extent of inflammation should be assessed and reported as it may influence treatment (broad agreement 8/9, 88.9%).

4.2 Skin ultrasound in infectious diseases

Ultrasound can be used to assess the extent of plantar warts and to monitor treatment response in human papillomavirus infections [42]. Ultrasound assessment of abscesses has also become widespread in emergency departments in the United States. Ultrasound evaluation in the pediatric emergency department can alter the treatment strategy (drainage vs. no drainage) in 15% of cases evaluated by physical examination alone [43]. An additional advantage of US in the diagnosis of abscesses is that it can be performed with minimal point-of-care training [44].

**POSITION STATEMENT 11**

Ultrasound is useful in evaluating and detecting subclinical subcutaneous abscesses in the emergency department. Ultrasound equipment and trained personnel are recommended for this kind of evaluation (broad agreement 7/9, 77.8%).

Psoriasis

Ultrasound characteristics of psoriasis include epidermal and dermal thickening and subepidermal hypoechogenic areas with increased blood flow on color Doppler US. These US findings, and in particular, dermal thickness, have been found to be correlated with disease severity measured using the Psoriasis Area Severity Index and other scales assessing the severity or extent of disease [45, 46]. In a multicenter study by the Spanish Rheumatology Society, high-frequency US evaluation showed a reduction in plaque thickness and Doppler signal intensity in the dermis of patients treated with infliximab [47]. Psoriatic nails appear thicker and irregular mainly on the ventral aspect of the nail plate in comparison with healthy nails or nails affected by other diseases (atopic dermatitis, mycoses) [48, 49]. Nail disease in psoriasis has been correlated with the presence of enthesopathy and psoriatic arthritis, even in the absence of clinical signs. Therefore, with clinically inconclusive signs of a psoriatic onychopathy, US may add useful information [50–52].

**POSITION STATEMENT 12**

In patients with suspected psoriatic arthritis, US findings of psoriatic onychopathy can support the diagnosis of psoriasis (strong agreement 9/9, 100%).

Hidradenitis suppurativa

Hidradenitis suppurativa is a predominantly dermal and subcutaneous inflammation that can be assessed in detail by US. Comparing epidermal and dermal thickness in patients with hidradenitis and healthy controls, Wortsman et al. found that areas of the body affected by hidradenitis exhibited an increased dermal-epidermal thickness, areas of lower echogenicity, formed by pseudo-cysts, fistulous tracts and fluid collections that allowed US diagnosis of hidradenitis suppurativa [53, 54].

In a multicenter study ultrasound better assessed hidradenitis suppurativa patients than clinical inspection alone, resulting in upstaging of most patients as a consequence of the identification of occult and deep fistulous tracts, previously considered to be part of an inflammatory nodule [55].

**POSITION STATEMENT 13**

Ultrasound is recommended for supporting diagnosis, staging and treatment monitoring in hidradenitis suppurativa treatment (strong agreement 9/9, 100%).

Collagenosis

Scleroderma is the collagenosis disease most widely studied by US, as treatment varies according to the stage of disease (inflammatory or sclerotic). In a study of 104 morphea plaques in 59 patients, Wortsman et al. [56] showed that US had a sensitivity of 100% and a specificity of 98.4% for differentiating between the inflammatory and the sclerotic phases. In the same study, US detected subclinical inflammation in five patients with Parry-Romberg syndrome. Attempts have been made to standardize these results to create semi-quantitative scales to assess the effectiveness of treatments [57]. With respect to scleroderma ulcers, inflammation and superinfection can also be assessed by US as increased vascularity is evidenced on color Doppler US [58]. Calcinoses associated with scleroderma and dermatomyositis can also be detected and evaluated by US [58, 59]. Recently shear wave elastography evaluation of scleroderma patients has been confirmed to be useful for the evaluation of generalized scleroderma patients [60].
5. Aesthetic dermatologic ultrasound

The number of aesthetic procedures has been increasing explosively worldwide over the last decade, and most of these techniques are performed blindly, sometimes in different institutions and by operators with variable levels of training from medical and non-medical backgrounds. Moreover, patients may forget or be unaware of the type of treatment that they received [61, 62]. Thus, anatomical information can be challenging to obtain, but is critical for management and outcome in aesthetic medicine, where good results are the goal, and scars and complications are unwanted. The use of US is expanding in aesthetic medicine since US can provide relevant information that includes data on facial anatomical variants, the type, location and extent of common cosmetic fillers, the identification of implants, the complications of lipolytic procedures, and the possibility of percutaneous US guidance for the procedure [62, 63].

5.1 Main indications for ultrasound in aesthetic medicine

1– Assessment of photoaging
Ultrasound can detect and measure signs of photoaging caused by prolonged exposure to the sun through the observation of the subepidermal low echogenic band (SLEB) which is produced by the deposit of glycosaminoglycans in the papillary dermis (upper dermis) [64].

2– Recognition of relevant anatomical data
Anatomical variants in vessels, muscles or glands, measurement of the thickness of skin layers and the assessment of blood flow in complications are relevant for planning or managing cosmetic techniques and complications. Moreover, the face, the most common corporal region for cosmetic procedures, presents a complex anatomical structure where the skin layers are thinner in comparison to other body regions, and any abnormality is highly visible [65, 66].

3– Management of cosmetic fillers
Ultrasound allows for the detection and identification of common cosmetic fillers as well as the assessment of their location and extent and potential complications. These fillers include deposits such as hyaluronic acid, polymethylmethacrylate, silicone (pure or oily forms), calcium hydroxyapatite, polyacrylamide, and polycaprolactone that are approved by the Food and Drug Administration (FDA) and European Union Medical Agencies as well as those that are not approved. Percutaneous US guidance of the injection of hyaluronidase has also been reported in the management of hyaluronic acid complications [67–69].

4– Detecting implants and their complications
Ultrasound can support the detection of organic and synthetic implants and their complications. Examples of organic implants are fat, cartilage and bone grafts. Examples of synthetic implants are pure silicone, polyethylene, and polydioxanone (e.g. tensor threads). Complications of implants include an excessively superficial location, extrusion, chronic inflammatory and fibrotic reactions as well as rupture [62, 65, 69–71].

5– Detection of complications of lipolytic procedures
The goal of these procedures is to decrease the amount of hypodermal fatty tissue using techniques that generate inflammation and liquefaction of the fat. These procedures include radiofrequency, mesotherapy or cryolipolysis. Ultrasound can provide anatomical information for planning these techniques, detect the extent and location of the inflammatory changes and may serve as a monitoring technique for the assessment of the outcomes or the management of potential complications [62, 72].
Ultrasound can detect complications of cosmetic or plastic surgery procedures, such as dermal and hypodermal edema, lymphedema, seromas, hematomas, abscesses, fistulous tracts, fat necrosis, granulomas and loose sutures [62, 65].

Ultrasound is recommended for the evaluation of plastic surgery procedures and their complications (broad agreement 7/9, 77.8 %).

Percutaneous ultrasound guidance of aesthetic or plastic surgery procedures can improve the precision of injections, drainage procedures and corrections and decrease potential adverse reactions or complications due to injury to neighboring structures [73, 74].

Percutaneous ultrasound guidance of non-surgical and surgical aesthetic procedures is recommended as it increases precision and decreases the potential for unwanted effects or neighboring injuries (strong agreement 9/9, 100 %).

A written report must be provided after a dermatologic US examination. High quality images should be recorded, stored and made available for follow-up examinations (broad agreement 8/9, 88.9 %)

Basic courses on dermatologic US should be both theoretical and practical with a minimum of two days of training. Clinical images should be available for training since clinical-sonographic correlation is key in this application (broad agreement 8/9, 88.9 %).
**References**


