Introduction

Sjogren’s syndrome (SS) is a systemic autoimmune disease characterized by the development of a functional impairment of exocrine salivary and lacrimal glands. The pathogenesis is related to the presence of a chronic lymphocytic infiltrate in the glandular parenchyma. Clinical presentation may be various, although the typical form of presentation includes xerostomia and xerophthalmia (sicca symptoms), which are present in the majority of patients (1). In younger patients, recurrent swelling of the parotid glands is also commonly reported (1). SS recognizes two forms of presentation. Primary SS (pSS) is characterized by the presence of sicca symptoms in absence of other systemic involvement, while secondary SS is characterized by the development of sicca symptoms in association with a concomitant systemic autoimmune disease (1,2). Considering pSS, both sexes at all ages can be affected,
although the disease is more often encountered in women between 30 and 50 years of age, with a prevalence range of 0.02–0.1% (2).

At the state of the art, the diagnosis of pSS is performed by a combination of clinical, serological, and histologic investigations, according to the American College of Rheumatology/European League Against Rheumatism (ACR/EULAR) criteria (1,3). Clinical examination aims at identifying the presence of sicca symptoms. Laboratory tests support the assessment of autoimmune activation, with the presence of positive autoantibodies Ro/SSA and/or La/SSB. Finally, minor salivary gland (SG) biopsy provides histologic confirmation and evaluation of the severity of lymphocytic infiltrate in glandular parenchyma through focus score estimation.

Recent technological advances in head and neck imaging, including facial and oral imaging, and in particular SGs imaging, allowed to improve the multimodality study of this challenging anatomical region (4–12). The following step is to integrate patients’ clinical and histopathology data with imaging biomarkers with the ultimate aim of using imaging biomarkers as decision-support tools to improve pSS diagnostic process and patients’ stratification in clinical practice.

From this perspective, the role of salivary glands ultrasound (SGUS) has progressively increased, due to the several advantages of this easy-to-perform technique that is non-invasive, widely available and consistently repeatable. At the present, the dominant pSS classification systems do not include ultrasound (US) findings in their criteria (13), although their inclusion in the ACR/EULAR criteria has been suggested by different authors in place of sialography and salivary scintigraphy (14–17).

Noteworthy, in the very recent past, the use of ultra-high frequency ultrasound (UHFUS) has emerged in various clinical settings, including the study of skin, peripheral blood vessels and nerves, and oral mucosa, showing a significant correlation with the information deriving from tissue histology and opening new avenues for the integration of tissue and imaging biomarkers (18,19).

Herewith, we provide a brief review of the literature about the role of imaging biomarkers in pSS focusing on conventional US of the major SGs and the technical background of UHFUS in order to discuss the current imaging concepts as well as the potential future applications of SGs UHFUS in the pSS clinical setting and research. We present the following article in accordance with the NARRATIVE REVIEW reporting checklist (available at http://dx.doi.org/10.21037/gs-20-529).

Methods

A literature search was performed in the PubMed record to write this narrative review. In particular, for conventional SGUS, only relevant review articles published from 1990 to 2020 were included. Given the limited number of publications regarding UHFUS, the search was extended to other databases (Web of Science and Google Scholar) to maximize the results and no relevant research work was excluded on the basis of year of publication, article type or study design. Only articles published in English language were included. Reference lists were also examined to find additional relevant articles.

Discussion

SGUS in SS: a brief review

Conventional SGUS is usually performed with linear-array probes with frequencies up to 22 MHz (13,20–23). The main indications for SGUS as stated by American Institute of Ultrasound in Medicine (AIUM) include the presence of diffuse enlargement and tenderness consistent with inflammatory sialadenitis and recurrent swelling suggesting pSS are included (24). A recent review stated that SGUS is an accurate method to identify the pSS echostructural anomalies within the SGs with specificity and sensitivity of 0.91% and 0.83%, respectively. Despite the clinical and methodological heterogeneity of the studies included in the review, these results are clinically interesting and highlight the significant contribution of SGUS in pSS diagnosis and follow-up (25–27). The variability of the interpretation of US images in patients with pSS still represents a limitation not yet overcome. However, imaging features like echogenicity and homogeneity were identified as reliable for pSS diagnosis, if assessed by a trained expert and evaluated in association with other classification criteria (28). While normal echostructure and homogeneity of SGs resembles the normal thyroid gland (hyperechoic compared to muscles), in affected glands echogenicity appears reduced by the presence of hypo/anechoic areas and/or increased by fibrosis (linear-shaped) or fatty infiltration (focal/diffuse), with a progressive evolution from hypo- to hyperechoic alterations. Parenchymal inhomogeneity with heterogeneous echostructure typically occurs in all the major SGs in...
patients with pSS, with demonstration of hypo/anechoic areas and hyperechoic foci/bands (28). Additionally, other US findings, such as cysts, calcifications, overall glands size variations, irregularities in glandular profiles and the presence of intra- or periglandular lymph nodes, may support pSS diagnosis (25).

The presence of bilateral changes in all the major SGs is the most relevant detectable US finding suggestive for pSS with the highest agreement with scintigraphy of the SGs, sialography, and minor labial salivary gland biopsy (LSGB). Finally, it has been suggested that in case of normal SGUS findings in patients with clinically suspected pSS with negative serum antibodies, can rule out pSS avoiding the LSGB (29). Indeed, SGUS findings have been evaluated as a predictor of the biopsy score. However, since the diagnosis of pSS is mainly based on the biopsy of minor SGs whereas conventional ultrasonography explores the echostucture of the major SGs the correlation between imaging biomarkers and minor SG histopathological grading remains moderate (30,31).

Different SGUS-based semiquantitative scores have been proposed but their wide variability, including heterogeneous cut-offs and inhomogeneous parameters for image interpretation, led to inconsistent specificity and sensitivity (20,25). However, the scoring system proposed by Jousse-Joulin et al. (21) seems to be effective in assessing SGs involvement with good diagnostic accuracy (25). Despite the promising results, further studies are needed to consolidate the role of SGUS as non-invasive tool for pSS diagnosis.

A role for prognosis estimation in pSS has also been claimed for SGUS due to the presence of a correlation between pathological US findings and increase in disease activity, risk of lymphoma, and development of systemic complications (32).

In addition to the SGUS morphologic evaluation, colour/power Doppler US (C/PDUS) can provide important information regarding glandular vascularization. Pathological C/PDUS findings in pSS include glandular hypervascularization, mainly with diffuse pattern: the small vessels within the glands are dynamically visible as punctiform areas of colour doppler signal. Indeed, hypervascularization showed a positive correlation with the parenchymal heterogeneity and the number of cyst-like alterations (25,33). Interestingly, quantitative estimation of pulsatility and resistivity indexes showed a significant reduction in patients with the early phase of SS, presumably due to the vasodilatation caused by the active glandular inflammation (25). However, a global reduction of C/PDUS signals was observed in patients with advanced pSS in association with an overall reduction of glandular volumes due to the limited active inflammation and to the increased fibrosis (25,33-35). Additionally, Martinoli et al. described the dynamic C/PDUS evaluation after lemon juice stimulation that caused increased vascular signal and decreased pulsatility and resistivity indexes in pSS pathological glandular parenchyma with subsequent normalization after about 20 seconds (33). Conversely, the facial artery in patients with pSS showed significantly lower changes in pulsatility and resistivity indexes compared to healthy controls (34). Among the C/PDUS parameters evaluated, Carotti et al. found that peak systolic velocity was more sensitive compared to the resistive index (36).

Recently, a new step towards the inclusion of SGUS in the 2016 ACR/EULAR pSS classification criteria has been made. Jousse-Joulin et al. (37) reported that SGUS had similar weight to other minor items already included in the 2016 ACR/EULAR pSS classification criteria, such as, namely, Schirmer test and unstimulated whole salivary flow rate. Moreover, including SGUS in the 2016 ACR/EULAR pSS classification criteria provided a higher sensitivity with no relevant change in specificity.

Nowadays, a consensus for a SGUS scoring system still represents an unmet need and the inclusion in the 2016 ACR/EULAR pSS classification criteria is not yet approved. However, SGUS examination with C/PDUS evaluation seems useful for the diagnosis and follow-up in patients with pSS and it is considered the first-time examination as widely available, repeatable, non-invasive, safe and low-cost investigation.

As in other clinical fields (18,38-43), UHFUS is being explored in the pSS scenario in order to evaluate its potential contribution to improve the evaluation of the SGs, particularly regarding minor SGs.

**UHFUS: technical notes**

UHFUS is a novel diagnostic technique which is finding an increasingly important role in several medical fields (18,19,38-45), including ophthalmology (38), dermatology (39,40), oral medicine (18,19,44,45), musculoskeletal (41,42) and vascular imaging (43). Since the axial resolution achievable increases with the frequency, the use of frequencies between 30–100 MHz allows to investigate superficial structures at a higher axial resolution compared to conventional US (46). The outstanding improvements in
image quality are provided by the shorter US wavelength (inversely proportional to frequency). Nevertheless, the wavelength reduces the US penetration, limiting the thickness of the superficial tissues that can be evaluated with UHFUS (47). Thus, the most appropriate trade-off between resolution and penetration should be found for each different clinical application (48).

In our institution, Vevo MD (VisualSonics, Toronto, Canada) a UHFUS equipment is available since 2016; The UHFUS system is equipped with two probes: UHF48 operating up to 46 MHz (center frequency 30 MHz), providing an image depth of 23.5 mm and an axial resolution of 50 µm, and UHF70 operating up to 71 MHz (center frequency 50 MHz), with image depth of 10.0 mm and axial resolution of 30 µm. UHFUS applications in different clinical fields have been explored, including rheumatology.

The current concepts of UHFUS application in major and minor SGs evaluation are reported and discussed in order to analyse the potential contribution of this technique in patients with pSS. Further considerations based on our preliminary clinical experience are also reported and discussed.

**SGs UHFUS in SS: current concepts**

**Major SGs UHFUS**

UHFUS of the major SGs is performed with the 48 MHz probe, similarly to conventional SGUS scan with both axial and longitudinal acquisitions of all the major SGs. Similarly to SGUS, normal UHFUS appearance of salivary glandular tissues is characterized by homogeneous echogenicity with defined glandular margins (Figure 1). In pathologic glands, diffuse parenchymal inhomogeneity, hypoechoic areas, and hyperechoic foci/bands can be detected (Figures 2 and 3). In patients with suspected pSS, the typical alterations in glandular echostructure, in particular inhomogeneity, can be better depicted with UHFUS due to the increased axial resolution and improved contrast-to-noise ratio (49). Thus, UHFUS of major SGs might improve diagnosis performance even at an early stage, allowing the identification of initial signs of inhomogeneity in the gland echostructure and of hypoechoic areas of small dimensions, which cannot be detected with conventional SGUS. Given the potential bimodal comparative evaluation that can be performed in major SGs, further comparison studies between SGUS and SGs UHFUS will permit to assess the role of SGs UHFUS in the reduction of false negative US examinations. It may be expected that the application of UHFUS to major SGs investigation may lead to an increase of the overall diagnostic accuracy in patient with pSS, particularly in the early phases. Indeed, the role of SGUS in the early diagnosis of pSS is still controversial (50). Although some authors stated that normal US findings in SGs have been observed in patients suffering from pSS for about 4.2 years (51), a recent study described the role of SGUS in the early diagnosis of pSS considering patients with symptoms duration inferior to 5 years (52). According to these considerations, the potential role of UHFUS in the early diagnosis of pSS should be further explored.

*Figure 1* Normal parotid (A) and submandibular (B) glands depicted by using UHFUS with the 48 MHz probe. Although the whole glandular structure cannot be imaged due to a limited depth of penetration of the ultrasound beam, high resolution provides a clear depiction of regular glandular margins (dotted line) and homogeneous parenchymal echostructure. UHFUS, ultra-high frequency ultrasound.
**Figure 2** Mild alteration of the parenchymal echostructure of parotid (A) and submandibular (B) glands depicted by using UHFUS with the 48 MHz probe. Unlike normal glands, initial phases of glandular inflammation secondary to pSS correspond to the presence of a fine inhomogeneity of the glandular parenchyma with occasional slightly hypoechoic areas with blurred edges (arrows). UHFUS, ultra-high frequency ultrasound; pSS, primary Sjogren’s syndrome.

**Figure 3** Severe alterations in the glandular parenchyma of parotid (A) and submandibular (B) glands in course of pSS depicted by using UHFUS with the 48 MHz probe. Multiple single (arrows) or confluent (arrowheads) hypoechoic areas can be detected, along with disruption of normal glandular structure. pSS, primary Sjogren’s syndrome; UHFUS, ultra-high frequency ultrasound.

**Minor SGs UFHUS**

Minor SGs are localized in the submucosal layer of the buccal mucosa, labial mucosa, lingual mucosa, soft/hard palate, and floor of the mouth (53). The study of minor SGs is of particular interest when considering labial localization. In fact, labial minor SGs are an important part of the diagnostic algorithm of pSS, as bioptic sampling is performed at this level to obtain histologic confirmation of diagnosis. In this sense, inferior labial mucosa represents an accessible site for surgical biopsy, due to a relatively superficial localization of the gland and a low risk of surgical complications. While minor SGs cannot be normally imaged with conventional US frequencies (53), UFHUS allows the evaluation of minor SGs thanks to the higher spatial resolution (up to 30 μm) provided by the 70 MHz probe (19). Among all the minor SGs localizations, the inferior labial mucosa is the most accessible for UFHUS scanning. Given the superficial localization of the labial glands and the relatively small thickness of the labial mucosa, the US penetration of this probe—up to 10 mm—results adequate to investigate these structures.

We present our preliminary experience regarding UFHUS of the minor labial salivary glands (MLSGs), including our methodological approach, findings and
Figure 4 Normal labial minor salivary glands evaluated with UHFUS operating at 70 MHz. Similarly to major salivary glands, each minor labial salivary gland shows homogeneous parenchymal echostructure and defined and regular margins (dotted line). UHFUS, ultra-high frequency ultrasound.

Figure 5 Mildly altered labial minor salivary glands depicted by using UHFUS with the 70 MHz probe. In the initial stage of pSS, salivary glands usually preserve normal dimensions, although slight inhomogeneity and rare hypoechoic areas (arrows) can be detected. UHFUS, ultra-high frequency ultrasound; pSS, primary Sjogren’s syndrome.

Figure 6 Severe alteration of labial minor salivary glands in course of pSS. Similarly to major salivary glands, minor salivary glands present irregular margins (dotted line) and extensive echostructural alterations, with hypoechoic areas (arrows), which in some cases appear confluent (arrowheads). Glands can be slightly decreased in terms of dimensions. pSS, primary Sjogren’s syndrome.

observations.

MLSGs UHFUS examination is carried out with the patient’s mouth slightly open and the lower lip gently stretched out with a sterile gauze. First, a small amount of US gel is placed on the scanning surface of the transducer, which is then enveloped with a disposable probe-cover to avoid cross-infection during intraoral UHFUS scan. Subsequently, a sufficient amount of isotonic physiological solution is placed on the covered probe and on the labial mucosa to optimize the US transmission and to reduce reflection. Finally, the labial mucosa is scanned with longitudinal and transverse positioning of the probe respectively in the central, lateral left, and lateral right compartment in order to provide a complete scan of the lip.

On UHFUS, MLSGs are depicted as roundish structures disposed in groups with characteristics of glandular parenchyma in terms of echostructure, as previously reported in the literature (19), appearing slightly hypoechoic compared to the hyperechoic submucosal connective tissue (Figure 4). Glandular dimensions and distribution vary through the different compartments, with numerous larger glands being localized in the lateral compartments and fewer smaller ones in the central area.

At UHFUS examination, normal MLSGs show a homogeneous echogenicity similar to major SGs. Despite the low amount of parenchyma in the MLSGs, alterations in the glandular echostructure can be detected, including parenchymal inhomogeneity, hypoechoic areas and hyperechoic foci/bands (Figures 5 and 6).

Based on a preliminary experience, Baldini et al. first demonstrated the feasibility of MLSGs UHFUS in 32 consecutive patients with clinically suspected pSS (54). All the patients were evaluated with a complete diagnostic work-up according to the ACR/EULAR criteria 2016, including LSGB (55). The pSS diagnosis was confirmed in 12/32 (37.5%) cases. According to their results (54), no differences between patients with pSS and controls with no-SS sicca syndrome were observed with UHFUS evaluation of MLSGs distribution and C/PDUS vascular parameters. On the contrary, a statistically significant increase of UHFUS inhomogeneity of glandular parenchyma (P=0.006) and a higher degree of fibrosis (P=0.01) was observed in the MLSGs of the lateral compartments (Figure 7) (54). This
Figure 7 Bar graphs showing the statistically significant increase of UHFUS inhomogeneity of glandular parenchyma (P=0.006) and the higher degree of fibrosis (P=0.01) observed in the MLSGs of the lateral compartments. Image and data were retrieved from ref (54). CTRL, controls; pSS, patients with primary Sjogren syndrome; G0-G3 and G0-G2 refer to the degree of inhomogeneity and fibrosis, respectively. UHFUS, ultra-high frequency ultrasound; MLSG, minor labial salivary gland.

work paved the way for a novel application of UHFUS technique in the assessment of pSS, allowing for the first time a non-invasive and highly detailed evaluation of the MLSGs. Further studies are needed to confirm the potential role of MLSGs UHFUS in the pSS clinical setting.

The application of UHFUS technique for MLSGs evaluation presents some intrinsic limitations. Contrary to SG-UHFUS, a direct comparison between UHFUS and conventional US in MLSGs is not feasible, due to the limitation of the latter hindering minor SGs imaging. From a technical point of view, the US scan with the 70 MHz probe is more difficult to perform due to the impact on the images of even minimal movements of the probe. The lack of dedicated probes may represent a limit especially in terms of invasiveness of the intraoral scan. The management of cross-infection risk is obviously an important issue which must be addressed by adopting adequate precautions in probe covering when performing intraoral UHFUS. However, the disposable cover may cause artefacts in the superior part of the scan due to the presence of multiple interfaces for the US beam prior to reaching the target tissue.

For these reasons, a training period may be needed to achieve the manual skills required to correctly perform the examination.

Future perspectives

The application of UHFUS technique appears promising for several future perspectives in the SGs evaluation for pSS assessment, moving towards a more accurate integration of tissue and imaging biomarkers. On the one hand, the higher resolution of UHFUS in major SGs may have a role in the early diagnosis of pSS, thus improving management and follow-up of patients. On the other hand, the possibility of direct UHFUS evaluation of the MLSGs
Figure 8 Overall comparison of the UHFUS visualisation of the normal appearance and the different degrees of alterations related to pSS in parotid (D,E,F), submandibular (J,K,L) and minor labial salivary glands (M,N,O). Comparative conventional US images for parotid (A,B,C) and submandibular (G,H,I) glands are included. Images acquired in the university hospital of Pisa. pSS, primary Sjogren syndrome; US, ultrasound; UHFUS, ultra-high frequency ultrasound.
may complete the diagnostic algorithm of pSS, in terms of giving additional information on the echostructure of these glands and of providing direct comparison with histology. The direct visualization of the MLGSs may improve pre-surgical planning for biopsies and subsequent histological examination, potentially reducing the number of inadequate biopsies. Moreover, the performance of comparative studies between UHFUS and histology may highlight a relevance for UHFUS investigation and potentially re-define the criteria for biopsy according to UHFUS-based parameters. If further studies will reveal the presence of a positive correlation between histology and UHFUS of minor SGs, pSS diagnostic algorithm will potentially be oriented to a less invasive approach limiting the biopsy of minor SGs to few selected cases.

Conclusions

This work aimed to present the current concepts of UHFUS as a novel technique for the evaluation of SGs by placing them in the state-of-the-art context of US applications in pSS, focusing on the pros and cons of this novel technique and its future potential applications.

Conventional US of major SGs appears a valid support for pSS diagnosis, although its role is still controversial. As for conventional US, UHFUS has the advantages of being safe, repeatable, and non-invasive. The high-resolution provided by the UHFUS thanks to the 48MHz and 70MHz probes allows to reach a high-detailed evaluation of the major and minor SGs. In Figure 8, the UHFUS aspect of normal major and minor SGs is shown, along with the different degrees of alterations related to pSS. For major SGs, comparative conventional US images are also provided (Figure 8).

UHFUS investigation of major SGs in the diagnostic algorithm of pSS seems promising, as the high resolution provided by the technique can be extremely valuable in detecting glandular alterations even at an early stage.

UHFUS of minor SGs represents a promising tool in the pSS work-up, as it is the only technique which allows a non-invasive evaluation of these structures with high resolution and high diagnostic confidence. Moreover, it could provide direct correlation between histology and echostructure, due to the biopsy sampling being performed on labial minor SGs.

This new technique applied to the major and minor SGs evaluation might lead to several potential improvements, including earlier diagnosis, reduction of unnecessary and inadequate biopsies and better management and follow-up of patients with pSS. However, given the small amount of preliminary data available, further studies are needed to confirm the role of UHFUS in the pSS clinical setting and research.

Acknowledgments

Funding: None.

Footnote

Provenance and Peer Review: This article was commissioned by the Guest Editor (Antonio Barile) for the series “Multimodality Advanced Imaging and Intervention in Gland Diseases” published in Gland Surgery. The article has undergone external peer review.

Reporting Checklist: The authors have completed the Narrative Review reporting checklist. Available at http://dx.doi.org/10.21037/gs-20-529

Conflicts of Interest: All authors have completed the ICMJE uniform disclosure form (available at http://dx.doi.org/10.21037/gs-20-529). The series “Multimodality Advanced Imaging and Intervention in Gland Diseases” was commissioned by the editorial office without any funding or sponsorship. The authors have no other conflicts of interest to declare.

Ethical Statement: The authors are accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved.

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