Clinical Study

Potential Applications of ⁶⁸Ga-PSMA-11 PET/CT in the Evaluation of Salivary Gland Uptake Function: Preliminary Observations and Comparison with ^{99m}TcO₄ ⁻ Salivary Gland Scintigraphy

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Purpose. To preliminarily evaluate the feasibility and potential of using ⁶⁸Ga-PSMA-11 PET/CT in evaluating the function of salivary glands and lacrimal glands in comparison with ^{99m}Tc-pertechnetate (^{99m}TcO₄⁻) salivary gland scintigraphy (SGS). *Methods*. A retrospective study was performed in 15 patients with different degrees of xerostomia and suspected salivary gland dysfunction. Each patient underwent ⁶⁸Ga-PSMA-11 PET/CT first and SGS the next day, and the findings of both scans were compared. *Results*. The results of ⁶⁸Ga-PSMA-11 PET/CT and SGS were consistent in 12/15 patients (80%) and were inconsistent in the remaining patients (20%). For 5 (33.3%) of 15 patients, ⁶⁸Ga-PSMA-11 PET/CT provided more information than did SGS. Additionally, ⁶⁸Ga-PSMA-11 PET/CT corrected the misdiagnosis by SGS for 1 patient. *Conclusions*. ⁶⁸Ga-PSMA-11 PET/CT is a potentially useful imaging tool for evaluating the function of salivary glands and lacrimal glands. ⁶⁸Ga-PSMA-11 PET/CT can be a promising supplement to SGS, and its clinical value deserves further study.

1. Introduction

The salivary glands consist of three pairs of major salivary glands (parotid glands, submandibular glands, and sublingual glands) and multiple minor salivary glands that are widely distributed in the oral mucosa. Glandular function impairment occurs in approximately 60% of patients with xerostomia [1]. Dry mouth is frequently observed in salivary gland dysplasia, in obstructive diseases, in autoimmune syndrome, and in space-occupying lesions, with the help of radiotherapy and chemotherapy for head and neck tumors and with large-dose ¹³¹I treatment, and is most commonly observed in Sjogren's syndrome. According to the American-European Consensus Group (AECG) criteria, the diagnosis of salivary and lacrimal gland dysfunction is mainly based on the objective and subjective evaluations of patients [2]. However, most of the current studies use objective indicators as a more convincing means of evaluation because the subjective symptoms of patients do not necessarily reflect the dysfunction of salivary glands [3, 4].

The objective evaluation methods of clinical salivary gland function mainly include saliva analysis, radiography of the parotids, and ^{99m}Tc-pertechnetate (^{99m}TcO₄⁻) salivary gland scintigraphy (SGS). The first two methods are affected by physiological factors, and their application is complicated; therefore, they are not ideal for clinical use. SGS allows a noninvasive inspection of the condition and has the advantages of simplicity, safety, and relatively low cost [5]. SGS is a commonly used method for the clinical evaluation of salivary gland function and has high specificity and

sensitivity [6]. SGS has been generally proposed to assess salivary gland function in different indications [7, 8]. SGS can be used to directly observe the morphology and function of the bilateral parotid and submandibular glands. Image analysis for SGS depends on the observer and lacks uniform standards [9, 10].

Over the past few years, prostate-specific membrane antigen- (PSMA-) targeted radiotracers have been widely used in the diagnosis (⁶⁸Ga-PSMA) and treatment (¹⁷⁷Lu-PSMA) of prostate cancer. Among them, PSMA-targeted positron emission tomography (PET), especially ⁶⁸Ga-PSMA-11, has been widely used for the detection, staging, posttreatment efficacy evaluation, and recurrence assessment of prostate cancer [11-13]. However, the binding of PSMA ligands is not limited to prostate cancer cells, and their physiological distribution includes the lacrimal glands, salivary glands, kidneys, duodenum, and small intestine [14]. Salivary glands and lacrimal glands have high uptake levels [14-17]. ⁶⁸Ga-PSMA PET/CT can enable the visualization of head and neck lacrimal glands, primary and secondary salivary glands, and seromucous glands. All normal glands have increased tracer uptake [16]. A recent study by Rupp found that the normal submandibular gland had high ⁶⁸Ga-PSMA-11 aggregation, but uptake of ⁶⁸Ga-PSMA-11 in chronic inflammatory atrophy of the submandibular gland was significantly lower [18].

The purpose of this study was to explore the value of ⁶⁸Ga-PSMA-11 PET/CT in evaluating salivary and lacrimal gland function and whether ⁶⁸Ga-PSMA-11 PET/CT has a complementary or alternative role to SGS.

2. Materials and Methods

2.1. Patient Selection. A total of 15 patients (10 men, 5 women; aged 19-75 years $[56.9 \pm 15.4 \text{ years}]$) with xerostomia and suspected salivary gland dysfunction were recruited in this study. The study included 4 patients with Sjogren's syndrome, 10 patients with head and neck tumors after radiotherapy and chemotherapy, and 1 patient with surgery for parotid space-occupying lesions. All patients underwent ⁶⁸Ga-PSMA PET/CT first, followed by SGS the next day. The inclusion criteria were as follows: clinical diagnosis of Sjogren's syndrome; radiotherapy and chemotherapy for head and neck tumors; high-dose ¹³¹I therapy; salivary gland surgery; salivary gland space-occupying lesions; and salivary gland dysplasia. The exclusion criteria were as follows: impairment of liver and kidney function, low white blood cell count, or pregnancy or lactation. The study followed the 1964 Helsinki Declaration and its later amendments or comparable ethical standards.

2.2. Image Acquisition

2.2.1. ⁶⁸Ga-PSMA-11 PET/CT. Images were acquired from the skull vertex to the root of neck using a PHILIPS Gemini TF PET/CT 16-tier system approximately 60 minutes after intravenous injection of 1.85MBq/kg ⁶⁸Ga-PSMA-11. Patients did not need specific preparation before PET/CT imaging. A low-dose CT scan (voltage: 120 kV, current: 100 mAs, 5 mm layer, 512×512 matrix, and 60 cm FOV) was performed first. Then, PET was performed (9-10 beds, 3 minutes/bed). After reconstruction, image analysis was performed using PHILIPS postprocessing fusion software.

2.2.2. ^{99m}TcO₄⁻ Salivary Gland Scintigraphy. On the day after ⁶⁸Ga-PSMA-11 PET/CT, the patients underwent SGS. The patients were injected with 185 MBq of ^{99m}TcO₄⁻, and dynamic images were acquired using a dual-head gamma camera (GE Healthcare, USA) equipped with a low-energy, high-resolution collimator (128 × 128 matrix, 140 KeV energy peak, 20% window width, 2 times amplification). The anterior image was collected with a probe, and the field of vision covered the entire thyroid and salivary glands. The blood perfusion phase image was collected immediately after intravenous injection of ^{99m}TcO₄⁻ (2 s/frame, 30 frames in total). Then, functional imaging was performed for 30 minutes at 40 s/frame. Vitamin C was given to stimulate salivary secretion 5 minutes before completion of the imaging. Images were analyzed on a Xeleris Workstation (GE Healthcare).

2.3. Image Analysis

2.3.1. ⁶⁸Ga-PSMA-11 PET/CT

Visual analysis. Three experienced nuclear physicians independently read the PET/CT images. When opinions differed, the majority opinion prevailed. Homogeneous, symmetrical, and strong uptake of ⁶⁸Ga-PSMA-11 in the bilateral parotid and submandibular glands was defined as normal. Abnormal uptake of ⁶⁸Ga-PSMA-11 in salivary glands included one of the following conditions. (a) The level of ⁶⁸Ga-PSMA-11 uptake in salivary glands was similar to that in background tissues (e.g., subcutaneous soft tissue of the head and neck); and (b) ⁶⁸Ga-PSMA-11 uptake into the glands presents a visible decrease compared to the contralateral side. Quantitative analysis. To quantify ⁶⁸Ga-PSMA-11 uptake, the volumes of interest (VOIs) of the bilateral lacrimal glands, parotid glands, submandibular glands, and thyroid glands were mapped on serial images. The maximum standardized uptake value (SUVmax) in the VOI of each gland was obtained by the software.

2.3.2. $^{99m}TcO_4^-$ Salivary Gland Scintigraphy. Visual evaluation: the images were independently and visually assessed by three experienced nuclear physicians. When the opinions differed, the majority opinion prevailed. The physicians were unaware of the results of other clinical studies. Classification of salivary gland function was based on the criteria used in the study by Kim et al. [19]. Salivary gland uptake is considered normal when it is similar to thyroid uptake. Normal excretion is defined as salivary gland activity similar to background activity after stimulation. Salivary gland function was graded on a scale from 1 to 3: 1, normal; 2, mild-tomoderate; and 3, severe. Grade 1 indicated that the uptake and excretion of salivary glands were normal; grade 2 indicated low uptake and/or delayed excretion; and grade 3 was defined as severe dysfunction with a complete absence of radioactivity in the salivary glands. Semiquantitative analysis: semiquantitative analysis was performed using the region of interest (ROI) technique. The ROIs of the bilateral parotid gland and submandibular gland were depicted, and the bilateral temporal region was used as a background reference. The computer automatically plotted the timeactivity curves. From the time-activity curves, the maximum value before vitamin C administration, the minimum value after vitamin C administration, and the background uptake value were obtained. The uptake ratio (UR) and excretion fraction (EF) of each salivary gland were calculated according to the following equation: UR = (maximumbackground uptake)/background uptake; EF = (maximumminimum)/(maximum-background uptake)×100% [20]. Decreased uptake function for the respective glands was defined as follows: UR (parotid gland) <2.28 and UR (submandibular gland) <1.60 [19]. Abnormality was indicated if the criteria were met in either visual analysis or semiquantitative analysis.

3. Results

The general information and PET and SGS imaging results for all 15 patients are shown in Table 1. This study included 4 patients with Sjogren's syndrome (patients 1–4), 10 patients with head and neck tumors after radiotherapy and chemotherapy (patients 5–14), and 1 patient with surgery for parotid space-occupying lesions (patient 15).

3.1. Image Findings by ${}^{68}Ga$ -PSMA-11 PET/CT and ${}^{99m}TcO_4^-$ Salivary Gland Scintigraphy. The results of ${}^{68}Ga$ -PSMA-11 PET/CT and SGS were consistent in 12/15 patients (80%; patients 1–12). Four patients with Sjogren's syndrome and 4 patients with head and neck cancer after radiotherapy and chemotherapy showed positive results, whereas four patients with head and neck cancer after radiotherapy and chemotherapy showed negative results.

The results of the two examinations were not consistent with those in the other 3 patients (patients 13–15). One patient who underwent surgery for parotid space-occupying lesions was positive only on ⁶⁸Ga-PSMA-11 PET/CT, which corrected the misdiagnosis by SGS. Two patients with head and neck cancer after radiotherapy and chemotherapy were positive only on SGS.

In 2 (13.3%; patients 5 and 15) of 15 patients, ⁶⁸Ga-PSMA-11 PET/CT provided more accurate information about the salivary glands than did SGS.

Additionally, in 3 (20%; patients 1–3) of 15 patients, ⁶⁸Ga-PSMA-11 PET/CT provided information about the lacrimal gland; decreased uptake of ⁶⁸Ga-PSMA-11 was observed.

Among all 15 patients, only one had an incidental finding of diffuse uptake of ⁶⁸Ga-PSMA-11 in the thyroid (patient 1: Figure 1, SUVmax 4.6). No significant uptake of ⁶⁸Ga-PSMA-11 was observed in the thyroids of the other patients. This patient was diagnosed with Hashimoto's thyroiditis.

4. Discussion

Klein et al. assessed the physiological distribution of ⁶⁸Ga-PSMA-11 in the salivary glands and seromucous glands of the head and neck and observed increased tracer uptake in all gland locations. The mean SUVmax ± standard deviation varied as follows: parotid, 12.3 ± 3.9 (range 5.2–22.9); sub-mandibular gland, 11.7 ± 3.5 (range 6.0–22.2); sublingual gland, 4.5 ± 1.9 (range 1.2–8.5); and lacrimal gland, 6.2 ± 2.2 (range 2.5–13.6) [16]. Our study also confirmed the high uptake of ⁶⁸Ga-PSMA-11 in the salivary and lacrimal glands, which is consistent with previous studies [14–17].

In our study, the results of ⁶⁸Ga-PSMA-11 PET/CT and SGS were consistent in 12 patients and inconsistent in the other 3 patients. Moreover, ⁶⁸Ga-PSMA-11 PET/CT corrected the misdiagnosis of patient 15 (Figure 2) by SGS and provided more information for 5 patients (33%) than SGS. ⁶⁸Ga-PSMA-11 PET/CT showed promising potential in evaluating the function of the lacrimal gland and salivary gland.

SGS is a planar SPECT imaging technique, while ⁶⁸Ga-PSMA-11 PET/CT is a positron emission computed tomography fusion imaging technique. ⁶⁸Ga-PSMA-11 PET/ CT can provide functional metabolic and anatomical images with higher resolution than SGS. ⁶⁸Ga-PSMA-11 PET/CT can accurately assess the morphology of the gland and the degree of atrophy to evaluate gland function more accurately. For example, in patient 1 (Figure 1) and patient 5 (Figure 3), ⁶⁸Ga-PSMA-11 PET/CT clearly shows the degree of atrophy and the density changes in the parotid and submandibular glands. In addition to the inherent advantages of PET/CT, ⁶⁸Ga-PSMA-11 PET/CT has the following unique advantages in the evaluation of lacrimal and salivary gland function.

- (a) ⁶⁸Ga-PSMA-11 PET/CT is significantly less affected by surgery than SGS. For example, in patient 15 (Figure 2), the left parotid gland region showed significant uptake of ^{99m}TcO₄⁻ and slight uptake of ⁶⁸Ga-PSMA-11. The ⁶⁸Ga-PSMA-11 PET/CT axial images clearly showed the absence of the left parotid gland and the structural disorder of the soft tissue. The $^{99m}TcO_{4}^{-}$ uptake in the left parotid gland region was considered to be caused by the operation according to the patient's history of total parotid gland resection one month previously. Without an accurate medical history, it may be a mistake to presume that the uptake function of the left parotid gland is normal. However, ⁶⁸Ga-PSMA-11 PET/CT can accurately show the postoperative changes in the left parotid gland and is significantly less affected by surgery.
- (b) ⁶⁸Ga-PSMA-11 PET/CT has higher sensitivity and image contrast for the uptake of imaging agents in glands than SGS. For example, on the ⁶⁸Ga-PSMA-11 PET/CT images, patient 5 showed slight uptake of ⁶⁸Ga-PSMA-11 in the right submandibular gland (Figure 3(a, c-e) small arrows, SUVmax, 2.6) and differences in the bilateral submandibular glands (SUVmax, right: left = 2.6:1.9). It could not be observed on the SGS image (Figure 3(b)). The bilateral

| | | | | | <u>(0</u> | |
|----------------|-----|------------|--|--|---|---|
| Patient no. | Sex | Age (Y) | History | Abnormal uptake of ^{99m} TcO ₄ ⁻ in PG or SM | Abnormal uptake of ⁶⁸ Ga- PSMA-11 in PG or SM | Additional findings from ⁶⁸ Ga-PSMA-11 PET/CT |
| 1 | F | 55 | Siggren's syndrome | Lt PG | Lt PG | Rt LG T |
| 2 | F | 67 | Siggren's syndrome | Rt SM Lt SM | Rt SM Lt SM | Rt LG Lt LG |
| 3 | M | 71 | Siggren's syndrome | Rt PG It PG | Rt PG It PG | Rt IG It IG |
| 4 | M | 70 | Siggren's syndrome | Rt SM | Rt. FG, Et. FG | |
| 5 | М | 64 | NPC after radiotherapy and chemotherapy | Rt. SM, Lt. SM | Rt. SM, Lt. SM | Rt. SM |
| | | | Vocal cord cancer after | | | |
| 6 | М | 64 | radiotherapy and chemotherapy | Lt. PG, Lt. SM | Lt. PG, Lt. SM | _ |
| 7 | F | 41 | NPC after radiotherapy and chemotherapy | Lt. PG | Lt. PG | _ |
| 8 | F | 66 | NPC after radiotherapy and chemotherapy | Rt. SM | Rt. SM | _ |
| | | | Papillary carcinoma of the | | | |
| 9 | М | 75 | jaw after radiotherapy and chemotherapy | Ν | Ν | _ |
| | | | Vocal cord cancer after | | | |
| 10 | М | 72 | radiotherapy and chemotherapy | Ν | Ν | — |
| 11 | М | 46 | NPC after radiotherapy and chemotherapy | Ν | Ν | — |
| | | | EMP of nasal cavity after | | | |
| 12 | М | 19 | radiotherapy and chemotherapy | Ν | Ν | — |
| 13 | М | 50 | NPC after radiotherapy and chemotherapy | Rt. SM | Ν | — |
| 14 | F | 43 | NPC after radiotherapy and chemotherapy | Lt. PG | Ν | — |
| | | | Postoperative treatment of | | | |
| 15 | М | 50 | left parotid acinar cell carcinoma | Ν | Lt. PG | Lt. PG |

TABLE 1: Patient data and analysis of the uptake characteristics of $^{99m}TcO_4^-$ and $^{68}Ga-PSMA-11$ in all 15 patients.

NPC = nasopharyngeal carcinoma; EMP = extramedullary plasmacytoma; N = negative; LG = lacrimal gland; PG = parotid gland; SM = submandibular gland; T = Thyroid. "—" means no relevant data.

parotid glands of patient 1 (Figure 1) showed better contrast on ⁶⁸Ga-PSMA-11 PET/CT (SUVmax_{right}: SUVmax_{left} = 9.2:1.7 = 5.4:1) than that on SGS (UR_{right}: UR_{left} = 10.0:2.5 = 4:1). ⁶⁸Ga-PSMA-11 PET/CT can clearly show the uptake of the imaging agent in the salivary glands, even in cases of low levels of imaging agents that are not normally visible on SGS. ⁶⁸Ga-PSMA-11 PET/CT can more clearly show and provide better image contrast for visible and invisible differences in gland pairs on SGS.

(c) The visual analysis of SGS imaging is affected by the extent of thyroid imaging agent uptake, but ⁶⁸Ga-PSMA-11 PET/CT image analysis is not affected. In addition, ⁶⁸Ga-PSMA-11 PET/CT can also unexpectedly identify thyroid lesions. Visual analysis of SGS images is usually based on the level of imaging agent uptake in the thyroid gland. The thyroid uptake of ^{99m}TcO₄⁻ increased in patient 15 (Figure 2). At this time, image analysis based on the thyroid uptake level underestimates the uptake function of salivary glands. No significant ^{99m}TcO₄⁻ uptake was observed in the thyroid of patient 1 (Figure 1(b)), and visual analysis of salivary gland function relied

on the experience of the observer. ⁶⁸Ga-PSMA-11 PET/CT images can be directly visualized and quantitatively analyzed without the thyroid as a reference. In patient 1 (Figure 1(f)-1(h)), diffuse uptake of ⁶⁸Ga-PSMA-11 in the thyroid was unexpectedly found but without 99m TcO₄⁻ uptake on the corresponding SGS imaging. The patient was diagnosed with Hashimoto's thyroiditis. A study by Kirchner, J et al. found that diffuse uptake in the thyroid (SUVmax \pm SD: 4.5 \pm 1.2) of ⁶⁸Ga-PSMA-11 was found in 22% (12/55) of patients. However, there were no indications about thyroid malignancies or other serious diseases in their data [14]. Some reports described the uptake of ⁶⁸Ga-PSMA in follicular adenoma, differentiated thyroid cancer, and medullary thyroid cancer [21–23]. A study by Silver et al. showed no expression of PSMA in the thyroid gland [24]. A previous study has shown that neovascular PSMA expression is common in thyroid cancer but may also rarely be found in benign thyroid diseases, such as follicular adenoma [25].

(d) ⁶⁸Ga-PSMA-11 PET/CT can provide accurate localization and quantitative analysis for lacrimal

Contrast Media & Molecular Imaging



(g)

(h)

FIGURE 1: A 55-year-old woman (patient 1) with dry mouth and a dry right eye for half a year was initially diagnosed with sjogren's syndrome. ⁶⁸Ga-PSMA-11 PET/CT MIP (a) and tomographic images (c-e) showed that the left parotid gland (SUVmax, 1.7) was smaller in volume and lower in density without ⁶⁸Ga-PSMA-11 uptake than the right parotid gland (SUVmax, 9.2). SGS (b) confirmed that the uptake function of the left parotid gland was severely decreased (UR_{left parotid}, 2.5; UR_{right parotid}, 10.0). In addition, it was found that the ⁶⁸Ga-PSMA-11 uptake of the right lacrimal gland was lower than that of the left side (SUVmax_{right lacrimal gland}, 2.0; SUVmax_{left lacrimal gland}, 4.2). Combined with ⁶⁸Ga-PSMA-11 PET/CT and the patient's medical history, the uptake function of the right eye was considered to be reduced. It was also found that the thyroid (f–h) density decreased with ⁶⁸Ga-PSMA-11 diffuse uptake but without ^{99m}TcO₄⁻ uptake. The patient with a history of taking supplements had a 7-year history of Hashimoto's thyroiditis.

glands. This is not possible on SGS. Three patients showed decreased uptake of 68 Ga-PSMA-11 in their lacrimal glands. For example, in patient 1 (Figure 1), the uptake of lacrimal gland 68 Ga-PSMA-11 on the right side was lower than that on the left side (SUVmax, right:left = 2.0:4.2). Combined with the patient's symptom of right eye dryness and the diagnosis of Sjogren's syndrome, the uptake

function of the right lacrimal gland was considered to be reduced. SGS can only evaluate the morphology and function of the parotid and submandibular glands, but it is useful for imaging and observing small glands, such as the lacrimal gland and sublingual gland. Our study shows that ⁶⁸Ga-PSMA-11 PET/CT can be used to visualize the lacrimal gland and evaluate its function.



FIGURE 2: A 50-year-old male (patient 15) underwent left parotid gland resection for a left parotid mass one month previously. Postoperative pathology revealed left parotid acinar cell carcinoma. ⁶⁸Ga-PSMA-11 PET/CT maximum density projection (MIP) (a) and corresponding tomography (c–e) show the absence of the left parotid gland and structural disorder of the soft tissue with slight uptake of ⁶⁸Ga-PSMA-11 (SUVmax, 2.7; right parotid gland SUVmax, 20.8). SGS (b) shows that ^{99m}TcO₄⁻ uptake in the left parotid gland region (UR 4.3) was slightly lower than that in the right parotid gland (UR 5.6). In addition, the thyroid uptake of ^{99m}TcO₄⁻ was significantly increased; this patient had a history of hyperthyroidism and took propylthiouracil for half a year.

However, ⁶⁸Ga-PSMA-11 PET/CT imaging also has the following limitations in assessing salivary gland function. (a) ⁶⁸Ga-PSMA-11 PET/CT cannot evaluate the excretion function of salivary glands. There is no effective method to reduce PSMA ligand uptake in salivary glands [26]. Further research is warranted. (b) Visual evaluation of ⁶⁸Ga-PSMA-11 PET/CT may have limitations in patients with mild to moderate reduction of salivary gland uptake on SGS. In patient 14 (Figure 4), SGS showed a mild-to-moderate decrease in bilateral parotid and submandibular gland uptake. However, there was no significant abnormality on ⁶⁸Ga-PSMA PET/CT imaging. This may be a defect in the visual assessment of the ⁶⁸Ga-PSMA-11 PET/CT, which may be compensated by more accurate quantitative analysis criteria.

Salivary gland analysis in ⁶⁸Ga-PSMA-11 PET/CT still has some problems that need to be solved. (*a*) The uptake mechanism of ⁶⁸Ga-PSMA-11 in salivary glands is unclear. Tracer uptake in prostate cancer is based on the expression of PSMA receptors in cells. However, the previous literature shows that the mechanism of the intense accumulation of PSMA radioligands in salivary glands is unclear [18]. Normal salivary glands and lacrimal glands have high PSMA ligand uptake. It was previously reported that PSMA is

physiologically expressed in normal salivary glands but at much lower levels than those in prostate cancer tissue [18, 27]. There are also studies showing that there is no PSMA expression in salivary glands [24, 28]. The intense uptake of PSMA ligands in salivary glands is not correlated with the expression level of PSMA. Some studies have shown that the uptake of PSMA-targeted radioligands in salivary gland tissues may be caused by both nonspecific and PSMAspecific uptake. However, the exact proportion of each form of uptake is still uncertain, and the mechanism of the nonspecific uptake is still unclear [29-32]. Recently, it has been reported that the accumulation of PSMA-targeted radioligands in salivary gland tissue is mainly caused by nonspecific uptake [18, 33]. Further research will be necessary to investigate the exact mechanism of radioligand accumulation in salivary glands. (b) At present, there is no uniform classification standard for ⁶⁸Ga-PSMA-11 uptake in *lacrimal glands and major salivary glands.* The clinical experience of ⁶⁸Ga-PSMA-11 PET/CT image interpretation is still limited. The uptake patterns of ⁶⁸Ga-PSMA-11 in salivary and lacrimal glands may need to be analyzed in future large sample studies and to develop accurate and reliable classification criteria.



FIGURE 3: A 64-year-old male patient (patient 5) with nasopharyngeal carcinoma received radiotherapy and chemotherapy. The ⁶⁸Ga-PSMA-11 PET/CT MIP (a) and tomographic images (c–e) clearly show a slight uptake of ⁶⁸Ga-PSMA-11 in the right submandibular gland (small arrow; SUVmax, 2.6) but no significant uptake of imaging agents in the left submandibular gland (large arrow; SUVmax, 1.9). The slice images (d–e) show that the volume of the left submandibular gland was significantly reduced. SGS (b) showed no significant ^{99m}TcO₄⁻ uptake in the bilateral submandibular glands, confirming a severe decrease in its uptake function.



FIGURE 4: A 43-year-old female (patient 14) with nasopharyngeal carcinoma after radiotherapy and chemotherapy. ⁶⁸Ga-PSMA-11 PET/CT MIP (a) shows that the imaging agents were evenly and symmetrically distributed in bilateral lacrimal glands and major salivary glands (right parotid gland, left parotid gland, right submandibular gland, and left submandibular gland; SUVmax values of 8.1, 8.0, 10.2, and, 8.2, respectively). No obvious abnormalities were observed. However, 99m TcO $_{4}^{-}$ SGS (b) shows a mild-to-moderate decrease in the uptake function of the left parotid (right parotid gland, left parotid gland, right submandibular gland, and left submandibular gland; UR values of 3.1, 2.2, 3.4, and 2.8, respectively).

5. Conclusions

⁶⁸Ga-PSMA-11 PET/CT imaging shows promising results in terms of lacrimal gland and major salivary gland uptake function. In contrast to SGS, ⁶⁸Ga-PSMA-11 PET/CT can visualize small glands, such as the lacrimal gland and sublingual gland, and is less affected by surgery. ⁶⁸Ga-PSMA-11 PET/CT may be used as a supplementary examination tool for SGS in the evaluation of salivary gland function. This study is a preliminary exploratory study, and the results of this study still need to be verified in subsequent large-scale studies.

Data Availability

The data used to support the findings of this study are included within the article.

Conflicts of Interest

The authors declare that they have no conflicts of interest.

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References

- S. Dugonjic, D. Stefanovic, B. Ethurovic, V. Spasic-Jokic, and B. Ajdinovic, "Evaluation of diagnostic parameters from parotid and submandibular dynamic salivary glands scintigraphy and unstimulated sialometry in sjogren's syndrome," *Hellenic Journal of Nuclear Medicine*, vol. 17, no. 2, pp. 116–122, 2014.
- [2] C. Vitali, S. Bombardieri, R. Jonsson et al., "Classification criteria for sjogren's syndrome: a revised version of the European criteria proposed by the American-European Consensus Group," *Annals of the Rheumatic Diseases*, vol. 61, no. 6, pp. 554–558, 2002.
- [3] C. Vitali, H. M. Moutsopoulos, and S. Bombardieri, "The European Community Study Group on diagnostic criteria for sjogren's syndrome. Sensitivity and specificity of tests for ocular and oral involvement in sjogren's syndrome," *Annals of the Rheumatic Diseases*, vol. 53, no. 10, pp. 637–647, 1994.
- [4] A. Vissink, W. W. I. Kalk, K. Mansour et al., "Comparison of lacrimal and salivary gland involvement in sjogren's syndrome," Archives of Otolaryngology-Head & Neck Surgery, vol. 129, no. 9, pp. 966–971, 2003.
- [5] M. N. Mojsak and F. Rogowski, "Application scintigraphy in evaluation of salivary gland function," *Polski Merkuriusz Lekarski: Organ Polskiego Towarzystwa Lekarskiego*, vol. 28, no. 165, pp. 214–219, 2010.
- [6] J. Chen, X. Zhao, H. Liu et al., "A point-scoring system for the clinical diagnosis of sjogren's syndrome based on quantified SPECT imaging of salivary gland," *PLoS One*, vol. 11, no. 5, Article ID e0155666, 2016.

- [7] H. P. van den Akker and E. Busemann-Sokole, "Absolute indications for salivary gland scintigraphy with ^{99m}Tc-pertechnetate," *Oral Surgery, Oral Medicine, Oral Pathology*, vol. 60, no. 4, pp. 440–447, 1985.
- [8] I. H. Liem, R. A. Valdés Olmos, A. J. M. Balm et al., "Evidence for early and persistent impairment of salivary gland excretion after irradiation of head and neck tumours," *European Journal* of Nuclear Medicine, vol. 23, no. 11, pp. 1485–1490, 1996.
- [9] F. B. Vivino and G. A. Hermann, "Role of nuclear scintigraphy in the characterization and management of the salivary component of sjogren's syndrome," *Rheumatic Disease Clinics* of North America, vol. 34, no. 4, pp. 973–986, 2008.
- [10] F. Vinagre, M. J. Santos, A. Prata, J. C. da Silva, and A. I. Santos, "Assessment of salivary gland function in sjogren's syndrome: the role of salivary gland scintigraphy," *Autoimmunity Reviews*, vol. 8, no. 8, pp. 672–676, 2009.
- [11] A. Afshar-Oromieh, A. Malcher, M. Eder et al., "PET imaging with a [⁶⁸Ga]gallium-labelled PSMA ligand for the diagnosis of prostate cancer: biodistribution in humans and first evaluation of tumour lesions," *European Journal of Nuclear Medicine and Molecular Imaging*, vol. 40, no. 4, pp. 486–495, 2013.
- [12] A. Afshar-Oromieh, E. Avtzi, F. L. Giesel et al., "The diagnostic value of PET/CT imaging with the ⁶⁸Ga-labelled PSMA ligand HBED-CC in the diagnosis of recurrent prostate cancer," *European Journal of Nuclear Medicine and Molecular Imaging*, vol. 42, no. 2, pp. 197–209, 2015.
- [13] A. Afshar-Oromieh, T. Holland-Letz, F. L. Giesel et al., "Diagnostic performance of ⁶⁸Ga-PSMA-11 (HBED-CC) PET/CT in patients with recurrent prostate cancer: evaluation in 1007 patients," *European Journal of Nuclear Medicine and Molecular Imaging*, vol. 44, no. 8, pp. 1258–1268, 2017.
- [14] J. Kirchner, B. M. Schaarschmidt, L. M. Sawicki et al., "Evaluation of practical interpretation hurdles in ⁶⁸Ga-PSMA PET/CT in 55 patients: physiological tracer distribution and incidental tracer uptake," *Clinical Nuclear Medicine*, vol. 42, no. 7, pp. e322–e327, 2017.
- [15] M. S. Hofman, R. J. Hicks, T. Maurer, and M. Eiber, "Prostatespecific membrane antigen PET: clinical utility in prostate cancer, normal patterns, pearls, and pitfalls," *RadioGraphics*, vol. 38, no. 1, pp. 200–217, 2018.
- [16] T. J. W. Klein Nulent, M. H. Valstar, B. de Keizer et al., "Physiologic distribution of PSMA-ligand in salivary glands and seromucous glands of the head and neck on PET/CT," Oral Surgery, Oral Medicine, Oral Pathology and Oral Radiology, vol. 125, no. 5, pp. 478–486, 2018.
- [17] L. W. M. van Kalmthout, M. Lam, B. de Keizer et al., "Impact of external cooling with icepacks on ⁶⁸Ga-PSMA uptake in salivary glands," *EJNMMI Research*, vol. 8, no. 1, p. 56, 2018.
- [18] N. J. Rupp, C. A. Umbricht, D. A. Pizzuto et al., "First clinicopathological evidence of a non PSMA-related uptake mechanism for (68)Ga-PSMA-11 in salivary glands," *Journal of Nuclear Medicine*, vol. 60, no. 9, pp. 1270–1276, 2019.
- [19] H.-A. Kim, S.-H. Yoon, J.-K. Yoon et al., "Salivary gland scintigraphy in sjogren's syndrome. Comparison of the diagnostic performance of visual and semiquantitative analysis," *Nuklearmedizin*, vol. 53, no. 4, pp. 139–145, 2014.
- [20] S. Klutmann, K. H. Bohuslavizki, S. Kroger et al., "Quantitative salivary gland scintigraphy," *Journal of Nuclear Medicine Technology*, vol. 27, no. 1, pp. 20–26, 1999.
- [21] G. L. Kanthan, J. Drummond, G. P. Schembri, M. A. Izard, and E. Hsiao, "Follicular thyroid adenoma showing avid uptake on ⁶⁸Ga PSMA-HBED-CC PET/CT," *Clinical Nuclear Medicine*, vol. 41, no. 4, pp. 331-332, 2016.

- [22] F. A. Verburg, T. Krohn, A. Heinzel, F. M. Mottaghy, and F. F. Behrendt, "First evidence of PSMA expression in differentiated thyroid cancer using [(68)Ga]PSMA-HBED-CC PET/CT," *European Journal of Nuclear Medicine and Molecular Imaging*, vol. 42, no. 10, pp. 1622-1623, 2015.
- [23] R. Ciappuccini, A. Edet-Sanson, V. Saguet-Rysanek, M. Gauthé, and S. Bardet, "Thyroid incidentaloma on 18Ffluorocholine PET/CT and ⁶⁸Ga-PSMA PET/CT revealing a medullary thyroid carcinoma," *Clinical Nuclear Medicine*, vol. 44, no. 8, pp. 663–665, 2019.
- [24] D. A. Silver, I. Pellicer, W. R. Fair, W. D. Heston, and C. Cordon-Cardo, "Prostate-specific membrane antigen expression in normal and malignant human tissues," *Clinical Cancer Research*, vol. 3, no. 1, pp. 81–85, 1997.
- [25] B. Heitkotter, K. Steinestel, M. Trautmann et al., "Neovascular PSMA expression is a common feature in malignant neoplasms of the thyroid," *Oncotarget*, vol. 9, no. 11, pp. 9867–9874, 2018.
- [26] T. Langbein, G. Chaussé, and R. P. Baum, "Salivary gland toxicity of PSMA radioligand therapy: relevance and preventive strategies," *Journal of Nuclear Medicine*, vol. 59, no. 8, pp. 1172-1173, 2018.
- [27] J. K. Troyer, M. L. Beckett, and G. L. Wright Jr., "Detection and characterization of the prostate-specific membrane antigen (PSMA) in tissue extracts and body fluids," *International Journal of Cancer*, vol. 62, no. 5, pp. 552–558, 1995.
- [28] P. Mhawech-Fauceglia, S. Zhang, L. Terracciano et al., "Prostate-specific membrane antigen (PSMA) protein expression in normal and neoplastic tissues and its sensitivity and specificity in prostate adenocarcinoma: an immunohistochemical study using mutiple tumour tissue microarray technique," *Histopathology*, vol. 50, no. 4, pp. 472–483, 2007.
- [29] D. Taïeb, J.-M. Foletti, M. Bardiès, P. Rocchi, R. J. Hicks, and U. Haberkorn, "PSMA-targeted radionuclide therapy and salivary gland toxicity: why does it matter?" *Journal of Nuclear Medicine*, vol. 59, no. 5, pp. 747-748, 2018.
- [30] R. P. Baum, T. Langbein, A. Singh et al., "Injection of botulinum toxin for preventing salivary gland toxicity after PSMA radioligand therapy: an empirical proof of a promising concept," *Nuclear Medicine and Molecular Imaging*, vol. 52, no. 1, pp. 80-81, 2018.
- [31] C. Kratochwil, F. L. Giesel, K. Leotta et al., "PMPA for nephroprotection in PSMA-targeted radionuclide therapy of prostate cancer," *Journal of Nuclear Medicine*, vol. 56, no. 2, pp. 293–298, 2015.
- [32] H. Rathke, C. Kratochwil, R. Hohenberger et al., "Initial clinical experience performing sialendoscopy for salivary gland protection in patients undergoing ²²⁵Ac-PSMA-617 RLT," *European Journal of Nuclear Medicine and Molecular Imaging*, vol. 46, no. 1, pp. 139–147, 2019.
- [33] R. Tonnesmann, P. T. Meyer, M. Eder, and A. C. Baranski, "(¹⁷⁷)Lu-PSMA-617 salivary gland uptake characterized by quantitative in vitro autoradiography," *Pharmaceuticals*, vol. 12, no. 1, 2019.