Review Paper: A Review of Signet Ring Cells in Salivary Gland Tumors

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Introduction: Signet Ring Cells (SRCs) are often present in gastrointestinal tract, breast, bile duct, and lung invasive carcinomas. Observing these cells in benign and malignant lesions of the salivary glands is usual. Signet-ring cell adenocarcinoma is rare and is commonly seen in secondary salivary glands. The presence of SRCs in other neoplasms is usually associated with poor prognosis; however, the prognosis and the survival of malignant tumors of the salivary glands containing SRCs are not well known and few studies have been conducted on this topic.

Materials and Methods: In this review article, 18 articles were reviewed from 1984 to 2017 in the PubMed, Google Scholar, and MEDLINE databases with the keywords of “salivary glands”, “parotid”, “submandibular”, “sublingual”, and “signet ring cells”.

Results: Viewing SRCs in the salivary glands is not common. Although the SRC adenocarcinoma has a low incidence, some of them may have been reported as Not Otherwise Specified (NOS) and in fact, have a higher incidence.

Conclusion: SRC adenocarcinoma of the salivary glands must be differentially diagnosed with more common tumors of these glands. Further studies are required to determine the exact prevalence of these salivary tumors that contain SRCs. Also, it seems necessary to predict their biological behavior.

ABSTRACT

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1. Introduction

Signet Ring Cells (SRCs) are often present in gastrointestinal invasive carcinomas, the gallbladder, breast, and lungs. These cells have been seen in many lesions and other organs with a much smaller outbreak. SRCs are not always indicative of malignancy, as SRCs have been already seen in the microscopic samples obtained from the colon of patients suffering from colitis and in the bile ducts removed because of the stones [1].

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Signet ring cell adenocarcinoma of the minor salivary glands was first introduced by Ghannoum and Freedman in 2004 [2]. Salivary gland tumors are an important group of head and neck tumors and 9% to 23% of primary salivary epithelial tumors are formed in the secondary salivary glands, 50% of which are malignant. Although SRC adenocarcinoma is a very rare subcutaneous salivary gland tumor, it includes 2% of the total malignancy of the secondary salivary glands.

The age range of the patients is 32-72 years and the female to male ratio is 2.5:1. Of course, the incidence of these tumors may be slightly higher, as some of them have been reported as Not Otherwise Specified (NOS) [3]. The most common site of minor salivary gland tumors is the hard palate and where the hard and soft palate meet [4]. They are also reported to be seen in different areas such as lips and buccal mucousa [5, 6]. These tumors are generally or typically seen as slowly growing asymptomatic masses, without invading the lower bone 4.

2. Materials and Methods

In this review study, articles from 1984 to 2017 in the PubMed, Google Scholar were searched with the keywords of “Salivary gland carcinoma”, “parotid”, “submandibular”, “sublingual”, and “signet ring cells”. Eventually, 18 articles were found.

3. Results

Viewing SRCs in the salivary glands is not common. Although the signet ring cell adenocarcinoma has a low incidence, some of them may have been reported as NOS and in fact, it might have a higher incidence.

4. Discussion

The observation of SRCs in neoplasms and tumor-like lesions of the salivary glands is rare1. SRCs have been reported in benign tumors such as Warthin and malignant tumors, such as minor salivary gland adenocarcinomas and a mucin-rich subgroup of salivary duct carcinoma and in the carcinoma ex-pleomorphic adenoma [9-11]. In the case of the Warthin tumor, the presence of oncocytes along with SRCs is helpful in the final diagnosis [9].

The mucin-rich variant of the Salivary Duct Carcinoma (mSDC) is very rare, and so far only 8 cases have been reported, of which 5 have SRCs. In 3 of these 5 samples containing SRCs, mSDC was a malignant component in pleomorphic adenoma and as a carcinoma ex-pleomorphic adenoma. In all of these cases, the survival rate has been reported to be low and about 10 to 28 months [11]. Signet ring cell adenocarcinoma is a kind of mucin-secreting adenocarcinoma, which often appears in minor salivary glands as an exophytic mass attached to the underlying tissue. The mean age of the affected patients is 56.4 years and is more common in women. However, this tumor has also been seen in the major salivary glands, such as parotid gland [12, 13].

In some cases, reports have indicated the presence of SRCs in the absence of mucin pools or lakes containing mucin [2, 12, 14]. The tumor is non-capsular and infiltrative. In cases reported by Singh et al. and Ghannoum et al., SRCs have been seen as cords and sheets of malignant cells with minor pleomorphism in salivary gland tumors [2, 12]. One case of mucus-producing adenopapillary carcinoma has also been reported in the minor salivary gland cells of the palate, whose microscopic view indicated a papillary pattern along with a large number of SRCs and mucin-contents [5]. This tumor in micro-
scopic view may be mistaken with mucoepidermoid carcinoma, where one should attend to the absence of epidermoid, intermediate, and clear cells and their differences with SRCs for differentiation [12].

SRCs in parotid other than mucin-secreting adenocarcinoma, have also been reported by Michal et al. in a rare case of oncocytic cystadenoma. In this case, numerous SRCs with bland morphology and oncocytic cells have been reported. In the microscopic view of this tumor, the sheets of the oncocytic cells are surrounded by myoepithelial cells, and most of the oncocytic cells contain large intracytoplasmic vacuoles (4-50 μm) and nuclei is located environmental and have appeared in the form of a signet ring cell. In this case, staining for musicarmin and cytokeratin was negative but staining for anti-mitochondrial antibody have been reported positive [13]. Rarely, signet ring lymphoma has also been seen in salivary glands, which is different from salivary gland tumors in terms of morphology and immunohistochemical profiles [12].

One recurrence of centroblastic and centrocytic nodular lymphoma has been reported in the salivary glands, which likens a signet ring cell lymphoma, and in ultrastructural studies, the cells contain empty vacuoles or residual membrane and myelinoid [15].

In salivary gland malignancies where SRCs are seen, certain metastatic lesions from other organs should be checked by accurate clinical and radiological examinations and laboratory evaluations such as thyroglobulin, Cytokeratin (CK), Carcino Embryonic Antigen (CEA), Alpha Feto Protein (AFP) , Cancer Antigen (CA-125), and Carbohydrate Antigen (CA)19-9 tests [12].

In cases where SRCs are seen in the tumor and Smooth Muscle Actin (SMA) marker reported as positive, clear cell epithelial-myoepithelial carcinoma is considered in differential diagnosis. This diagnosis could be ruled out thanks to the specific morphology of the SRCs and the negativity of S100 [12]. Because of the morphology of the tumor and its mucin content, it may be mistaken with mucoepidermoid carcinoma, particularly its clear cell variant, mucinous adenocarcinoma, and Polymorphous Low-Grade Adenocarcinoma (PLGA), which can be distinguished from mucoepidermoid carcinoma and mucinous adenocarcinoma with due attention to the general cytomorphology and growth pattern. Mucinous adenocarcinoma is a rare malignant neoplasm composed of floating epithelial cells in basophilic mucin lakes, separated by fibrous septa.

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The cells are single or form duct-like structures, dense cords, or cell clusters. These cells are polyhedral or cubic and have eosinophilic and occasionally clear cytoplasm [3]. In PLGA, evident cellular eosinophilic cytoplasm, oval/round, and bland nuclei, pagetoid pattern around the nerve and infiltrative pattern of Indian file (Single file) around the tumor can help to diagnose and differentiate properly [3].

The mucinous variant of myoepithelioma is also a myoepithelial tumor with clear prominent cytoplasmic areas that often contain mucin content with the morphology of signet ring [16]. Unlike signet ring carcinomas in other organs, this tumor in the salivary glands is often referred to as low-grade neoplasm, where its recurrence and metastasis are rare after complete excision [12]. The subtype of the "signet ring" in gastric and esophageal adenocarcinoma is invasive and metastatic with poor prognosis [17]. However, in rare cases where metastases to
the lymph node exist, besides complete excision, neck dissection and radiotherapy are also needed [18].

In the sample reported by Horn et al. a signet ring cell adenocarcinoma was found in the parotid gland that unlike the minor salivary glands, its metastasis was seen to the lymph node and brain [14]. The Ki67 index is the most useful immunohistochemical marker for determining the prognosis of patients with salivary gland tumors. In the case of high Ki67, the prognosis is poor and the survival rate is often low [19].

5. Conclusion

Signet ring cell adenocarcinoma of the salivary gland is rare and must be differentially diagnosed with more common tumors of the salivary glands such as mucoepidermoid carcinoma and PLGA. Although the features and prognosis of signet ring cell adenocarcinoma are known in other parts of the body such as the gastrointestinal tract, further studies and investigations are required to determine the exact prevalence of salivary gland signet ring adenocarcinoma, its features, prognosis, and treatment methods.

Paraffin blocks of the samples labeled as NOS adenocarcinoma should also be investigated in this regard because some of the signet ring cell adenocarcinomas of the salivary glands have been inappropriately reported as adenocarcinomas NOS.

Ethical Considerations

Compliance with ethical guidelines

There is no ethical principle to be considered doing this research.

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Authors contributions

Laleh Maleki: Study concept, study design, literature review, manuscript preparation; literature review, manuscript preparation: Elha Arzhang and Leili Alizadeh; and editing and review: Leili Alizadeh.

Conflict of interest

The authors declared no conflict of interest.

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References


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