Extraparotid Warthin Tumors Imitating Metastasis of Oral Cancers

Michał Gontarz1, Krzysztof Gąsiorowski1, Jakub Bargiel1, Tomasz Marecik1, Paweł Szczurowski1, Jan Zapala1, Grażyna Wyszęńska-Pawelec1

1 Department of Cranio-Maxillofacial Surgery, Jagiellonian University Medical College, Cracow, Poland

Address for correspondence Michał Gontarz, DDS, PhD, Assistant Prof., Department of Cranio-Maxillofacial Surgery, Jagiellonian University Medical College, University Hospital, 2 Jakubowskiego Street, 30-688 Cracow, Poland (e-mail: mgontarz@op.pl).

Abstract

Introduction Extraparotid Warthin tumor (WT) is a very rare entity, especially when synchronous with oral cancer (OC).

Objective The present study presents a case series of extraparotid WTs detected in the surgical specimen of patients treated for OC.

Methods From 2007 to 2016, 336 patients were operated for OC in our institution. Neck dissection was performed in 306 patients.

Results In the 306 patients operated for OC whose necks were dissected, unexpected WTs were observed in 4 surgical neck specimens. In 3 cases, extraparotid WTs were responsible for tumor, node, metastasis (TNM) overstaging before surgery.

Conclusion Extraparotid WTs may be discovered during neck dissection in ~ 1% of OC patients, and they may mimic neck metastasis, especially in positron-emission tomography/computed tomography (PET/CT) imaging.

Keywords ► Warthin tumor ► adenolymphoma ► squamous cell carcinoma ► mouth neoplasms ► head and neck squamous cell carcinoma

Introduction

Warthin tumor (WT) is the second most common salivary gland tumor, most commonly found in the tail of the parotid gland. The extraparotid localization of this tumor is uncommon and is observed up to 8% of all WTs. Warthin tumor can be observed in periparotid lymph nodes, especially at the first and second neck levels and may mimic metastasis in patients suffering from oral cancer (OC). Only a few case reports of synchronous WT and OC have been published in the English-language literature. This study presents a case series of extraparotid WTs detected in surgical specimen of patients treated for OC.

Methods

Between January 2007 and December 2016, 336 patients were operated due to OC in our institution. Neck dissection was performed in 306 patients, and, in 187 cases, neck dissection was bilateral. Unexpected WTs were observed in 4 surgical specimens. Two cases of synchronous parotid WT and squamous cell carcinoma of retroauricular region and two cases of metachronous parotid WT and OC were excluded from the study. Medical charts of the patients were evaluated retrospectively according to histopathological aspects and treatment. This study was approved by the institutional review board (No: 122.6120.287.2016).

Result

Four hundred and ninety-three neck dissection were performed in 306 patients, 184 of which were modified-radical/radical neck dissections (MRND/RND) (37.5%) and 309 of which were selective neck dissections (SND) (62.5%). Extraparotid WTs were observed in the neck specimen of 4
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Table 1 Characteristics of patients with extraparotid Warthin tumor

<table>
<thead>
<tr>
<th>No</th>
<th>Age</th>
<th>Sex</th>
<th>Site of OC</th>
<th>cTNM</th>
<th>Neck dissection</th>
<th>Site of WT</th>
<th>pN</th>
<th>Additional treatment</th>
<th>Follow-up (months)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>86</td>
<td>F</td>
<td>buccal mucosa</td>
<td>T3N1M0</td>
<td>SOHND</td>
<td>Ila</td>
<td>N2b</td>
<td>PORT</td>
<td>18</td>
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<tr>
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<td>67</td>
<td>F</td>
<td>tongue</td>
<td>T2N1M0</td>
<td>MRND</td>
<td>Ila</td>
<td>N0</td>
<td>No</td>
<td>35</td>
</tr>
<tr>
<td>3</td>
<td>65</td>
<td>F</td>
<td>lower gingiva</td>
<td>T2N1M0</td>
<td>SOHND</td>
<td>Ila</td>
<td>N0</td>
<td>No</td>
<td>28</td>
</tr>
<tr>
<td>4</td>
<td>54</td>
<td>M</td>
<td>FOM</td>
<td>T3N1M0</td>
<td>SOHND</td>
<td>lb</td>
<td>N0</td>
<td>No</td>
<td>46</td>
</tr>
</tbody>
</table>

Abbreviations: cTNM, clinical tumor, node, metastasis stage; F, female; FOM, floor of mouth; M, male; MRND, modified radical neck dissection; OC, oral cancer; pN, pathologic node stage; PORT, postoperative radiotherapy; SOHND, supraomohyoid neck dissection; WT, Warthin tumor.

Discussion

Warthin tumors are the second most common benign salivary gland tumors after pleomorphic adenoma. The majority of WTs occur in the parotid gland. The multifocal and bilateral appearance of WTs might occur synchronously or metachronously. Extraparotid WTs are rare and sometimes multifocal. Most extraparotid WTs are located in the periparotid lymph nodes at the Ib, Ila, and III levels of the neck. The synchronous appearance of extraparotid WTs and OC is an extremely rare entity. In the English-language literature, there are only a few papers describing single-case reports. Only Shehan et al. presented one case of extraparotid WT (0.5%) as an unexpected finding in neck dissection in 202 patients operated for head and neck squamous cell carcinoma. In the current study, extraparotid WTs in neck specimen were observed in 4 patients (1.3%) with OC.

Microscopically, WTs comprise eosinophilic oncocytic double-layered epithelium lining the cyst lumen and hematoxyphilic lymphoid aggregates in the stroma. The epithelial and lymphatic components of WT are explained by two theories. The first one is that during embryogenesis, salivary elements might be trapped heterotopically within the extraparotid and parotid lymph nodes. This might explain why there is no evidence of WTs in the lower neck levels (IV and V). An alternative theory suggests that WT is not a neoplastic lesion but, most probably, it is caused by a delayed hypersensitivity reaction to degenerated oncocyttes. The pathogenesis of WTs is still unclear. Some studies suggest that epithelial components and lymphocytic infiltrations are polyclonal, which may be evidence of the non-neoplastic nature of WTs. On the other hand, some WTs are characterized by a presence of at (11;19) translocation, similar to that of mucoepidermoid carcinoma. Such WTs are monoclonal and truly neoplastic and may lead to malignant transformation. But it is still unclear which factors trigger the development of WTs.

Smoking cigarettes has a strong association with WT development. In their study, Lewis et al. support that smoking cigarettes cause an oxidative damage of the mitochondrial DNA of oncocytic cell, which lead to the occurrence of WTs. Smokers have eight times higher risk of tumor development than non-smokers. Tobacco abuse is also one of the most important risk factors for OC. This might explain the coincidence of synchronous and metachronous WTs and OC.

A useful diagnostic tool for the staging and treatment planning of patients suffering from OC is the 18-fluorodeoxyglucose positron emission tomography (FDG-PET/CT). However, false positive FDG uptake is observed in inflammations and benign tumors. Warthin tumors are known to have high FDG uptake and, as such, might mislead to diagnosis of metastatic lymph nodes. Also, PET/CT is used relatively often in the follow-up of oncological patients. In this case, WT as an incidentaloma may be found in the tail of the parotid gland. In those cases, fine-needle aspiration cytology (FNAC) is recommended to confirm the diagnosis of WT. The diagnostic accuracy of FNAC for identifying WTs is good, ranging from 74 to 100%. According to the study by Schwajle et al., the growth rate of WTs is slow, especially in patient over 75 years old. For this reason, conservative management of WTs confidently diagnosed with FNAC is an option mainly for patients suffering from head and neck cancers treated with non-surgical modality.

Conclusion

Unexpected WTs may be discovered during neck dissection in ~1% of OC patients. Abnormal FDG uptake in PET/CT is typical for WTs, and it may suggest metastatic lymph node. For these reasons, the possibility of synchronous or metachronous WTs and OC should always be kept in mind by surgeons and radiotherapists, especially in patients with a history of WT.

Conflict of Interests

The authors declare that there is no conflict of interests.

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