Case Report

Metastatic squamous cell carcinoma of the gingiva appearing as a solitary branchial cyst carcinoma: diagnostic role of PET/CT

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Abstract: We herein present a case of a left cervical cystic mass, for which the initial pathological diagnosis was branchial cleft cyst carcinoma (following complete mass excision). Thorough postoperative examinations, including with FDG positron emission tomography/computed tomography (PET/CT), revealed a primary tumor in the retromolar region of the left mandible. A 52-year-old female presented with a 2-month history of a painless, progressively enlarged left-sided neck mass. Fine-needle aspiration biopsy suggested a branchial cleft cyst. Physical examination revealed a 3 × 3-cm smooth, tender mass in the upper-left neck and anterior border of the sternocleidomastoid muscle. Examination using nasendoscopy and a strobolaryngoscope revealed no abnormalities of the nasal cavity, nasopharynx, oropharynx, hypopharynx or larynx. MRI of the neck revealed a solitary, round, cystic mass under the left parotid gland. The mass was excised completely. Pathologic results indicated a branchial cleft cyst carcinoma. According to the diagnostic criteria for a branchial cleft cystic carcinoma, PET/CT was performed to detect the occult primary site. PET/CT revealed high FDG uptake in the tooth root of the left mandible. Frozen sections of the mass were indicative of moderate, differentiated squamous cell carcinoma. The carcinoma in the retromolar region of the left mandible was locally excised under general anesthesia. A partial left maxillectomy, partial mandibulectomy, and left radical neck dissection were performed. The patient received postoperative concurrent chemoradiotherapy, and was disease-free at the 8-month follow-up. True branchial cleft cyst carcinoma is rare; once diagnosed, it should be distinguished from metastatic cystic cervical lymph and occult primary carcinoma. FDG PET/CT is useful in the identification of occult primary tumor.

Keywords: Branchial cleft cyst carcinoma, metastatic cystic cervical lymph node, carcinoma of unknown primary site, positron emission tomography/computed tomography

Introduction

The majority of metastatic cervical lymph nodes in head-and-neck squamous cell carcinomas (HNSCCs) are solid masses. However, between 33% and 62% of cases are cystic metastatic squamous cell carcinomas (SCC) [1-4]. Diagnosis and management for both clinicians and pathologists is problematic if lesions are characterized by a solitary cystic appearance [5].

The most-common variety of solitary cervical cystic lesion is the branchial cleft cyst (BCC), which can become malignant. According to the strict diagnostic criteria for branchial cleft cystic carcinoma (BCCC), proposed by Martin et al. [6] and modified by Khafif et al. [7], fewer than 40 cases have been reported [1]. The majority of previously suspected BCCCs were actually metastatic cystic lymph nodes; misdiagnosis can occur due to the absence of long-term follow-up [6, 8]. Thus, solitary metastatic cystic lymph nodes, from carcinomas of the upper aerodigestive tract, are difficult to differentiate from BCCCs, especially in patients more than 40 years of age [9]. The proportion of metastatic SCCs initially diagnosed as BCCs, or BCCCs, ranges between 11% and 21% [2, 10, 11]. Metastatic cystic SCCs in the upper lateral neck are easily mistaken for BCCCs if the primary site of the carcinoma is not detected [5].
Diagnostic role of PET/CT in metastatic squamous cell carcinoma

The challenge, therefore, is to distinguish BCCC from occult metastatic cystic SCC, and to identify the primary site of metastatic cystic SCC in the head and neck. Routine work-ups, with ultrasound (US), computed tomography (CT), magnetic resonance image (MRI), and panendoscopy, can aid in diagnosis and differentiation. Several studies have emphasized the role of fine-needle aspiration cytology (FNAC), or fine needle aspiration biopsy (FNAB), under US guidance. [1, 5] **[18]F** 2-Fluoro-2-deoxy-D-glucose (**[18]F-FDG**) positron emission tomography/CT (PET/CT) imaging has been widely used to detect unknown primary tumors [12]. However, the utility of FDG PET/CT for the diagnosis and differentiation of cervical cystic masses is limited, and its role disputed [13-15], especially in adults in whom the primary tumor is very small [15]. Therefore, the utility of PET/CT requires further validation.

In the present study, we report a case of a left cervical cystic mass for which the initial pathological diagnosis was of branchial cleft cyst carcinoma (following complete excision of the mass). Thorough postoperative examination, including with FDG PET/CT, revealed a primary tumor in the retromolar region of the left mandible.

**Figure 1.** MRI of the neck revealed a solitary, round mass under the left parotid gland: (A) The T1- and (B) T2-weighted signals were hyperintense; (C) DWI suggested hyperintense lesions ($b = 1000 \text{ s/mm}^2$); (D) Contrast-enhanced T1-weighted images revealed peripheral enhancement, but no enhancement in the cystic region.

**Figure 2.** The mass was excised completely using a left-neck lateral approach under general anesthesia (A); a surgical sample (B).

**Case report**

A 52-year-old female presented with a 2-month history of a painless, progressively enlarged left-sided neck mass. There was no nasal obstruction, nasal bleeding, respiratory distress, hoarseness, cough, odynophagia, weight loss, or dysphagia, and no childhood history of neck swellings. In November, 2013, the patient visited a local hospital. US and CT revealed a cystic mass in the left upper neck. FNAB further demonstrated that the mass contained yellow liquid, and that squamous epithelial cells and a portion of nuclear cells were atypical. The pathological results suggested a branchial cleft cyst accompanied by infection. The patient was admitted to our department for surgery. Physical examinations revealed a 3 × 3-cm smooth, tender mass in the upper-left neck, and in the anterior border of the sternocleidomastoid muscle. Nasendoscopy, and a strobolaryngoscope, revealed no abnormalities in the nasal cavity, nasopharynx, oropharynx, hypopharynx or larynx. MRI of the neck revealed a solitary, round mass under the left parotid gland. The T1- and T2-weighted signals were hyperintense. Diffusion-weighted imaging (DWI) suggested hyperintense lesions ($b = 1000 \text{ s/mm}^2$): contrast-enhanced T1-weighted images revealed peripheral, but not cystic, enhancement (Figure 1). On November 13, 2013, the mass was excised completely using a left-neck lateral approach under general anesthesia (Figure 2). Pathologic results indicated a BCCC. According to the diagnostic criteria for BCCC, proposed by Martin et al. [6] and modified by Khaffif et al. [7], PET/CT was performed to detect the occult primary site. The scan revealed high FDG uptake in the surgical
regions previously delineated, and in the tooth root of the left mandible. The surrounding mandibular bone was partially degraded, with high FDG uptake (Figure 3). Oral examination revealed a 1 × 1-cm rough mass in the retromolar region of the left mandible. On December 27, 2013, frozen sections of the mass indicated moderately differentiated SCC. Carcinoma in the retromolar region of the left mandible was locally excised under general anesthesia. A partial left maxillectomy, partial mandibulectomy and left radical neck dissection were performed. The tongue flap was used to address the surgical defect. The postoperative period was uneventful: the patient received postoperative concurrent chemoradiotherapy (CCR; 5,000 cGy in 200-cGy fractions delivered over 25 days and chemotherapy using cisplatin, at 37 mg, on days 1-3 [one cycle per 4 weeks, four cycles in total]). The patient was disease-free at 8 months postoperatively.

**Discussion**

BCCC is currently regarded as an uncommon clinicopathological entity [1]. In 1950, Martin et al. reviewed 250 cases of BCCC, of which the majority represented metastatic head and neck primary cancers resulting from an absence of long-term follow-up [6]. Strict diagnostic criteria for BCCC were subsequently established, as follows: a) tumor located along the anterior border of the sternocleidomastoid muscle; b) histological findings consistent with tissue originating from a branchial cleft; c) histological evidence of carcinoma arising in the wall of an epithelial-lined cyst; and d) no evidence of a primary source during a minimum 5-year follow-up period [6]. In 1989, Khafif et al. modified the above criteria as follows: a) tumor located in the anatomic region of a branchial cleft cyst; b) histological appearance consistent with a branchial vestige origin; c) for a squamous cell carcinoma, presence of the carcinoma within the lining of an identifiable epithelial cyst; d) evidence of transition from a normal squamous epithelium of the cyst to carcinoma; and e) absence of any identifiable primary malignant tumor following exhaustive evaluation of the patient [7]. According to these criteria, fewer than 40 cases were reported [1].

A solitary cystic metastatic cervical lymph node, from a small occult primary tumor, is easily misdiagnosed as BCCC, especially during initial presentation. Hardee et al. reported two cases of solitary cystic metastatic cervical lymph nodes initially presenting as branchial cysts [16], one of which was not located at the primary site. In the other case, primary carcinoma was discovered in the right tonsil [16]. Solitary cystic masses located in level II of the neck are problematic for clinicians. Briggs et al. suggested the following reasons for this diag-

![PET/CT revealed high FDG uptake in the above-mentioned surgical regions (A; SUVmax = 2.27) and at the site of the tooth root of the left mandible (B; SUVmax = 20.77); the surrounding, the mandibular bone was partially degraded.](image)
nostic difficulty: a) the two diseases always manifest in similar locations; b) cervical cystic metastases might represent the initial presentation of an occult primary carcinoma in the upper aerodigestive tract; and c) histological features usually confer difficulty with respect to differentiation of a branchial cleft carcinoma from solitary cystic degeneration of a metastatic cervical lymph node [9]. In the present case, the patient also presented with a painless solitary cystic mass in the left side of level II of the neck. FNAB suggested a branchial cleft cyst. A primary lesion was not discovered during thorough work-up investigations. Subsequently, the mass was completely excised. The postoperative histological results revealed that the lesion was a BCCC, and should therefore not be considered an occult primary carcinoma according to the criteria of Martin et al. [6] and Khafif et al. [7].

Routine work-ups usually employ US, CT, MRI and panendoscopy, and can aid in the detection of the primary site. Between 72% and 90% of primary tumors of cervical metastatic SCC are located within Waldeyer’s ring [1, 2, 4]. Directed biopsy of this area using panendoscopy or tonsillectomy is recommended if these routine work-ups do not detect any primary lesions. Several researchers have recommended that bilateral tonsillectomy represent the standard approach to detect the primary site of a carcinoma of unknown primary treatment (CUP) in the head and neck [17]. Metastatic cervical lymph nodes in thyroid carcinoma are also a common cause of cystic masses in the neck [18, 19]. Recently, PET/CT has been shown to confer advantages in the detection of CUPs. Researchers within our center have assessed the role of PET/CT in the diagnosis of CUPs [20-22]. In a previous study, we reported that the success rates of FDG PET and CT in the detection of CUPs were 73.3% and 52.4%, respectively [22]. We initially reported a case of tonsillar metastasis of small-cell neuroendocrine carcinoma of the lung using FDG PET/CT following tonsillectomy [20]. Wong et al. reported use of FDG PET/CT to detect 30 primary cancers in 75 patients in whom no primary tumor was detected using CT/MRI. Furthermore, PET/CT was still useful in patients subjected to tonsillectomy [23]. However, Ferris et al. reported that PET/CT might not confer diagnostic advantages in adults with suspicious cystic neck masses if the primary tumor is very small [24]. However, with technical improvements, contrast-enhanced FDG PET/CT now possesses advantages compared with non-enhanced FDG PET/CT in the detection of cystic lymph nodes [25]. In the present case, FDG PET/CT revealed that the primary site was the retromolar region of the left mandible. Findings during surgery and postoperative pathologic results confirmed the PET/CT findings. The patient received timely and appropriate treatment.

In conclusion, true branchial cleft cyst carcinoma is rare. Once diagnosed, it should be distinguished from metastatic cystic cervical lymph and occult primary carcinoma. FDG PET/CT is useful for the identification of an occult primary tumor.

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Disclosure of conflict of interest

None.

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