Sequential treatment from mandibulectomy to reconstruction on mandibular oral cancer – Case review I: mandibular ramus and angle lesion of primary intraosseous squamous cell carcinoma

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Abstract (J Korean Assoc Oral Maxillofac Surg 2021;47:120-127)

Primary intraosseous squamous cell carcinoma (PIOSCC) is very rare type of squamous cell carcinoma (SCC) that occurs within the jaw and arises from remnants of odontogenic epithelium with no connection to the oral mucosa. This study reports two cases of PIOSCC of the mandible. Reported in this article are two cases of PIOSCC of the mandible that were treated with resection and reconstruction using a fibular free flap. The first case was a 36-year-old male patient who complained of right mandibular pain. Computed tomography (CT) and panoramic radiograph revealed a large radiolucency in the mandibular ramus area. At first, an odontogenic keratocyst was tentatively diagnosed, and an excision procedure was carried out at another clinic. A final biopsy after cyst enucleation revealed well-differentiated SCC, so we proceeded with segmental mandibulectomy and reconstruction using a fibular free flap. The second case was a 48-year-old male patient with left mandibular pain. CT and panoramic radiograph revealed irregular radiolucency in the mandibular angle area near tooth #38. At first, osteomyelitis was tentatively diagnosed, and a curettage was carried out. A later biopsy revealed well-differentiated SCC, so segmental mandibulectomy and reconstruction with a fibular free flap were secondarily performed. Our two cases have had no recurrence. The facial appearance of both patients is satisfactory, and the neo-mandibular body created using a fibular bone transfer displays adequate bony volume.

Key words: Oral squamous cell carcinoma, Mandibular reconstruction

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I. Introduction

Primary intraosseous squamous cell carcinoma (PIOSCC) is a type of squamous cell carcinoma (SCC) that arises within the jaw, has no initial connection with the oral mucosa, and presumably develops from residues of the odontogenic epithelium or from an odontogenic cyst or tumor. According to the World Health Organization (WHO) classification published in 2005, the term “PIOSCC” is intended to replace old terms, and its subtypes include solid type and that derived from odontogenic cyst or tumor. There are three subcategories of PIOSCC:

1. A solid neoplasm that invades the bone marrow space and induces osseous resorption
2. SCC that arises from the lining of an odontogenic cystic lesion, a category that is subdivided into carcinomas arising in odontogenic keratocysts and carcinomas arising in other odontogenic cysts
3. SCC associated with other epithelial odontogenic tumors.

This study presents the features of two cases of PIOSCC and a review of the literature.

II. Case Presentation

1. Case 1

A 36-year-old male patient visited Pusan National University Dental Hospital in July 2013, for pain in the right mandibular region from the mandibular body to the mandibular ramus. Computed tomography (CT) and panoramic radiography revealed a large mandibular radiolucency.
Odontogenic keratocyst was tentatively diagnosed, and excision with extraction of teeth #47 and #48 was carried out at another clinic. The specimen biopsy from the first surgery indicated the presence of well-differentiated SCC (cT4N0M0).

The patient underwent additional magnetic resonance imaging (MRI), enhanced CT, and positron emission tomography-CT (PET-CT). The MRI and CT revealed a 3.9-cm restricted diffusion in the right mandibular body. No enhancing lymph nodes with diffusion restriction were found. The maximum standardized uptake value (SUVmax) was 7.8 on PET-CT, which showed no metastatic lymph nodes. (Fig. 2)

Supraomohyoid neck dissection (SOHND) was performed at our hospital on the right side along with wide excision and segmental mandibulectomy. This patient had an intraoral...
defect and ipsilateral neck vessels, so a contralateral fibular free flap was chosen to cover the intraoral defect and resected mandible areas. Mandibular ramus reconstruction with a two-piece fibular free flap was carried out after mandibular resection. (Fig. 3) The biopsy indicated the presence of a solid tumor of the PIOSCC type (pT4aN0M0). (Fig. 4) At first, the multilocular radiolucency seen on radiographs was considered to be SCC arising from an odontogenic cyst. However, pathologic findings revealed no cystic lining in the lesion. The mandibular resection defect was successfully anastomosed with a fibular free flap. (Fig. 5) The patient was not given postoperative radiotherapy or chemotherapy due to the presence of a clear surgical margin and the absence of neck node metastasis. Seven years after mandibular resection and reconstruction, no recurrence has been observed.

2. Case 2

A 48-year-old male patient visited Pusan National University Dental Hospital in April 2020, with pain in the left mandibular region and paresthesia of the left lower lip. A radiolucent lesion was visible near the root of his wisdom tooth (#38) at the mandibular angle. Osteomyelitis was tentatively diagnosed, and curettage along with extraction of tooth #38 was carried out. (Fig. 6) The biopsy suggested a PIOSCC (cT4aN1M0) lesion (SCC arising from the lining of an odontogenic cyst).

The patient underwent additional MRI, enhanced CT, and PET-CT. MRI and CT revealed a 1.8-cm restricted diffusion on the left mandibular body that extended to the lateral aspect of the left medial pterygoid muscle and the medial aspect of the left masseter muscle. An enhancing lymph node with diffusion restriction in the left neck at the IIa level also was identified. The SUVmax was 6.3 on PET-CT, which revealed a level-IIa metastatic lymph node. (Fig. 7)

SOHND was performed on the patient’s left side. Wide excision and segmental mandibulectomy were carried out. A Mandibular Reconstruction Angled Plate (Stryker, Kalamazoo, MI, USA) was prepared preoperatively using a rapid prototype model and was used as the reconstruction plate after being cut according to the bone resection site. (Fig. 8) This patient also had an intraoral defect and ipsilateral neck vessels, so a contralateral fibular free flap was chosen. After resection, mandibular reconstruction was carried out using a fibular free flap. (Fig. 9) The biopsy result indicated the pres-
ence of well-differentiated SCC arising from the lining of an odontogenic cyst (pT4aN1M0). (Fig. 10) The mandibular resection defect was successfully anastomosed with the fibular free flap. (Fig. 11) The patient received 30 cycles of postoperative radiotherapy (60 Gy) and chemotherapy using cisplatin to prevent tumor recurrence. Four months after mandibular resection and reconstruction, there was no inflammation or other complications, and no recurrence has occurred.
III. Discussion

PIOSCC is a rare malignant tumor that develops in the jaw. Diagnosis of PIOSCC can be difficult as it must be distinguished from other cancers, including malignant tumor metastases to the jaw and oral mucosal epithelium tumors that have invaded the bone. The etiology, prevalence, and incidence of PIOSCC have not been concluded.

PIOSCC is defined as an SCC that arises within the jaw. The WHO has proposed the term “primary intraosseous odontogenic carcinoma” to describe this type of SCC. The WHO has subcategorized this lesion into three subtypes: a solid-type, a tumor arising from an odontogenic keratocyst, and a tumor associated with other epithelial odontogenic tumors.

PIOSCC has a predilection for males (2:1 male-to-female ratio) and occurs often in the mandibular body and angle. In this study, 36-year-old and 48-year-old males were affected, and the lesions involved the mandibular body, angle, and ramus areas.

Clinically, PIOSCC has some distinct symptoms according to type and size. These symptoms are pain, swelling, lip numbness, and teeth mobility. In this case study, the 36-year-old male patient reported pain in the lower right gin-
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The hallmark radiographic features of PIOSCCs are similar to those of odontogenic tumors. Early stage PIOSCC can mimic odontogenic tumor or dental disorder, which could lead to misdiagnosis\(^\text{11}\). In this study, the author misdiagnosed the early stage PIOSCC lesion in the 48-year-old male as osteomyelitis at the first visit. While the other patient did not have early stage PIOSCC, that patient’s first dentist diagnosed the lesion as an odontogenic keratocyst. After resection and neck dissection, adjuvant radiotherapy and chemotherapy were not performed due to lack of neck metastasis.

Radiographically, in most cases, osteolytic bony changes are characterized by diffuse, ill-defined, and irregular margins\(^{11}\). In this study, the lesions showed similar features, including irregular osteolytic bony changes in both the 36-year-old and 48-year-old patients. MRI and PET-CT scanning modalities have advantages over other imaging techniques for anatomical assessment of head and neck cancers, especially with regard to mapping neck metastasis and primary tumor extent\(^{12}\).

In our cases, the 48-year-old male patient had a left level-IIa neck metastasis visible on both MRI and PET-CT.

The prognosis of PIOSCC is generally not good, although well-differentiated SCC is reported to have a better prognosis\(^{6}\). Wenguang et al.\(^{13}\) reported estimated two-year and five-year overall survival rates of 68.9% and 38.8%, respectively. de Morais et al.\(^{14}\) reported that PIOSCC has a poor prognosis, with high rates of mortality. Neck lymph node metastasis and histological grading have been suggested as prognostic indi-

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Table 1. Summary of patient characteristics

| Variable          | Case 1          | Case 2          |
|-------------------|-----------------|-----------------|
| Age (yr)          | 36              | 48              |
| Sex               | Male            | Male            |
| Symptom           | Pain            | Pain            |
| Location          | Right mandible  | Left mandible   |
| Initial diagnosis | OKC             | Osteomyelitis   |
| Initial treatment | Cyst enucleation| Curettage       |
| Final treatment   | Wide excision   | Wide excision   |
| Subtype           | Solid type      | Arising from cyst |
| Stage             | pT4aN0M0        | pT4aN1M0        |
| Adjuvant treatment| Negative        | Positive        |
| Follow-up period  | 7 yr            | 4 mo            |

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Fig. 10. Histological findings of Case 2 (H&E staining). A. Hole view (×20) of cyst wall (square) and normal squamous epithelium (arrow). B. Magnification view (×100) of the part of primary intraosseous squamous cell carcinoma (PIOSCC) (square) and normal squamous epithelium (arrow). C. Magnification view (×200) of the part of PIOSCC. It shows a pattern of invasion to basal membrane and a small and thick nucleus which is different in size between other nuclei.

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Fig. 11. The patient’s intraoral view after 2 months postoperatively.

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cators of these tumors. Fortunately, the carcinomas were well-differentiated in both of our cases. In the 48-year-old male, a left level-IIa neck metastasis was confirmed. Concomitant chemotherapy and radiation therapy was performed, and the follow-up period was very short. Further follow-up will be needed to provide additional review of these cases.

IV. Conclusion

The author presented two cases of PIOSCC. (Table 1) Case 1 (a 36-year-old male) had no lymph node metastasis and has shown no recurrence after seven years. Case 2 (a 48-year-old male) demonstrated a level-IIa lymph node metastasis and has shown no recurrence after six months. Both cases were misdiagnosed as other odontogenic diseases due to the type of PIOSCC lesion. The present cases highlight the importance of careful interpretation of radiographs.

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Authors’ Contributions

W.B.L. participated in data collection and wrote the manuscript. D.S.H. contributed to conception and revision of the manuscript. U.K.K. provided with clinical data, participated in the study design and data analysis. All authors read and approved the final manuscript.

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Ethics Approval and Consent to Participate

The study was approved by the Institutional Review Board of Pusan National University Dental Hospital (IRB No. PNUDH-2020-034). Informed consent was waived by the IRB.

Consent for Publishing Photographs

Written informed consent was obtained from the patients for publication of this article and accompanying images.

Conflict of Interest

No potential conflict of interest relevant to this article was reported.

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