Case report

Intraosseous lipoma of the jaws: Review of the literature and rare case report

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\textbf{A R T I C L E  I N F O}

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\textbf{A B S T R A C T}

\textbf{Background:} Lipoma is a benign tumor that arises at the expense of mature adipose tissue, it can occur anywhere in the body that contains adipose tissue. However, the intraosseous lipoma is considered a rare bone tumor, that affects the long and flat bones, but it is uncommon in the jaws. Usually, diagnosis based on clinical and radiographic features is insufficient and histopathology analysis is of utmost importance for the final diagnosis.

\textbf{Case presentation:} In this article, we present a case report of a 28-year-old female patient who visited the dentist with the intention of extracting an impacted tooth. She had no medical history with medications or genetic diseases. The tooth was extracted with a small soft mass that was subjected to histological analysis and it was found to be an intraosseous lipoma.

\textbf{Discussion:} Intraosseous lipoma is a rare tumor of the oral cavity, and it can be difficult to diagnose when only clinical and radiographic features are based, and the definitive diagnosis is based on histological examination. The treatment is complete surgical excision.

\textbf{Conclusions and literature review:} 30 cases were discovered and archived since 1948 until now, most of them were in the posterior region of mandibular, and the infection of females is higher than males with different final sub-diagnosis of types of intraosseous lipoma.

1. Introduction

\textbf{Lipoma} is a benign tumor of mature adipose tissue without cellular atypia, well-circumscribed, painless and slow-growing soft mass located in subcutaneous tissue, muscles, retroperitoneal space and bones, it is considered to be the rarest benign primary tumors of the bones [1].

Some hypotheses suggest that osseous lesions are primary benign tumor [2], while some of these hypotheses are likely to be a reactive bone lesion following bone infarct or trauma. Another hypothesis suggests that the cause of the tumor is conglomeration of fatty marrow, as commonly seen in calcaneus or vertebral bodies [3]. Although this tumor has been reported in the long bones: (humerus, radius, femur, tibia, fibula) as well as the calcaneus and cervical spine, also in the flat bones (ribs, pelvis, and skull), but in 20 % of all cases they are found in the head and neck region. However, only 1–4 % of lipomas are located in the oral cavity and the most common places of infection are the parotid gland, buccal mucosa, lip and tongue [4].

The literature reports that intraosseous lipoma usually occurs at age between (20–65) years, and males are reported to be more frequently affected than females [2].

In general, intraosseous lipoma is asymptomatic [2] can be clinically silent, and then the lesion may be found incidentally during radiological investigation following an injury in the same region, but in some cases patients may suffer from swelling, pain and paresthesia [1].

Radiologically, some cases of intraosseous lipoma appear as well-defined osteolytic lesions with sclerotic borders [5] [6], But in other cases, some intraosseous lipomas are difficult to diagnose radiologically, just like the case in this article. Consequently, diagnosis of this lesion is based on clinical, radiological and histopathological features [5] [6], while emphasizing that the gold standard for accurate diagnosis remains the histopathological evaluation [7].

According to Cakarer et al. [5]. The etiology and characteristics of the jaw lesions are not clear, due to the small number of cases so far, stating the importance of documentation of each new case of intraosseous mandibular lipoma. Therefore, we present this rare case of mandibular lipoma, which was written according to SCARE criteria [8].
2. Presentation of case

Patient, female, aged 28 years was referred to her dentist with the aim of extraction the right impacted third molar. The patient's general health was good and there was neither a history of medical problems nor trauma, she did not suffer from any psychiatric or genetic diseases, and had no history with any specific medications. Extraoral examination did not reveal any abnormal results. Intraoral examination showed no dilatation or signs of inflammation.

The radiographic examination did not show clear signs of the presence of the lesion and there was no resorption of the root of the third molar (Fig. 1).

The surgery was performed by a general dentist with 20 years clinical experience, under local anesthesia and surgical suturing with 4/0 silk sutures, and the suture removed after a week. After the surgery, the patient was prescribed routine medications (analgesics and anti-edema).

On clinical examination, and after the extraction, the doctor found a yellowish, non-cystic mass, soft and not attached to the crown of the tooth or the root, measured in biggest dimensions: 12 \times 10 \times 06 \text{mm} (Fig. 2) with no traces of pus in the area. Dentigerous cyst was excluded as a differential diagnosis of the lesion because it is not associated with the crown of an impacted molar. Odontogenic Keratocyst was also excluded as a differential diagnosis due to the non-cystic lesion shape.

To make a final diagnosis, the sample was fixed in formaldehyde solution and sent directly to oral maxillofacial pathology service for histological examination. Microscopically, using traditional staining (H&E) it was found that the lesion consisted of mature adipose tissue without cellular atypia. It was confirmed using immunohistochemical stains and the final diagnosis was Intraosseous lipoma (Figs. 3–5).

3. Discussion

Intraosseous lipoma(IOL) is one of the rarest benign tumors of the bone accounting for less than 0.1 % of primary bone tumor [5] [6], only in 20 % of all cases they are found in the head and neck region, while in the oral cavity it affects the soft tissues and the reason for the rareness of this tumor inside the jaws may be attributed to the lack of the lipid component inside these bones, in addition to the fact that the number of
Summary of site, radiographic features and histological diagnosis of the published cases of Intraosseous lipoma of the jaw since 1948.

Table 1

| Authors          | Year | Sex | Age | Site                                      | Radiographic features | Histological diagnosis  |
|------------------|------|-----|-----|-------------------------------------------|------------------------|-------------------------|
| Oringer          | 1948 | F   | 37  | Lower second molar                        | Well-defined radiolucency | Intraosseous Lipoma    |
| Newman           | 1957 | M   | 65  | Associated to impacted third molar         | Well-defined radiolucency | Fibrolipoma             |
| Johnson          | 1969 | M   | 21  | Associated to impacted second and third molars | Well-defined radiolucency | Intraosseous Lipoma    |
| Polte et al.     | 1976 | M   | 39  | Second premolar to second molar            | Moderate well-defined radiolucency | Angiolipoma             |
| Lewis et al.     | 1980 | F   | 56  | Mandibular body                           | Multilocular well-defined | Angiolipoma             |
| Steiner et al.   | 1981 | M   | 50  | Right third molar                         | Well-defined radiolucency | Parosteal lipoma        |
| Miller et al.    | 1982 | M   | 51  | Impacted left third molar                 | Well-defined radiolucency | Intraosseous Lipoma    |
| Heir and Geron   | 1983 | F   | 43  | Ramus                                    | Well-defined radiolucency | Intraosseous Lipoma    |
| Barker and Sloan | 1986 | F   | 53  | Impacted right third molar                | Well-defined radiolucency | Intraosseous Lipoma    |
| Manganaro et al. | 1994 | F   | 51  | Posterior left mandible and mandibular ramus | Well-defined radiolucency | Angiolipoma             |
| Koami et al.     | 1995 | M   | 59  | Mandibular symphysis                      | Well-defined radiolucency | Intraosseous Lipoma    |
| Sakashita et al. | 1998 | M   | 17  | Left side of Maxilla                      | Well-defined radiolucency | Intraosseous Lipoma    |
| Burić et al.     | 2001 | F   | 62  | Mandibular symphysis                      | Multilocular well-defined | Intraosseous Lipoma    |
| Keogh et al.     | 2004 | F   | 56  | Impacted right third molar                | Well-defined radiolucency | Intraosseous Lipoma    |
| Colella et al.   | 2005 | F   | 20  | Posterior left mandible                   | Well-defined radiolucency | Intraosseous Lipoma    |
| Darling and Daley| 2005 | F   | 22  | Anterior mandible (not associated with the apex of the anterior teeth) | Well-defined radiolucency | Intraosseous Lipoma    |
| Cakar et al.     | 2009 | F   | 45  | Anterior mandible (associated to the apex of the anterior teeth) | Well-defined radiolucency | Intraosseous Lipoma    |
| Gonzales et al.  | 2010 | F   | 61  | Left mandibular ramus                     | Radiolucency            | Intraosseous Lipoma    |
| Silva et al.     | 2010 | M   | 18  | Right mandibular body                     | Well-defined radiolucency | Intraosseous Lipoma    |
| Morais et al.    | 2011 | F   | 39  | Upper left third molar                    | Well-defined radiolucency | Intraosseous Lipoma    |
| Hemavathy et al. | 2012 | F   | 21  | From the right mandibular lateral incisor to the left ramus | Well-defined radiolucency | Intraosseous Lipoma    |
| Olgać et al.     | 2012 | F   | 48  | Incisor area of the mandible              | Radiolucency            | Intraosseous Lipoma    |
| Basheer et al.   | 2013 | M   | 15  | Right mandibular                          | Radiopaque              | Intraosseous Lipoma    |
| Castellaneti et al. | 2015 | F   | 25  | Right mandibular ramus                    | Well-defined radiolucency | Intraosseous Lipoma    |
| Waikowska et al. | 2017 | M   | 32  | Right body of the mandible                | Well-defined radiolucency | Intraosseous Lipoma    |
| Sanjuan et al.   | 2017 | F   | 50  | Left mandibular ramus and condyle          | Multilocular well-defined | Intraosseous Lipoma    |
| Cooper et al.    | 2017 | M   | 53  | Right mandible                            | Well-demarcated unilocular radiolucency | Spindle cell lipoma    |
| Tabakovic et al. | 2018 | F   | 43  | Left side of Maxilla                      | Radiopaque              | Intraosseous Lipoma    |
| Moghadam et al.  | 2021 | M   | 39  | Left lateral incisor and canine of mandibular | Well-defined radiolucency | Fibrolipoma             |
| Maksoud and Aoun | 2022 | F   | 37  | Anterior region of the mandible           | Well-defined unilocular radiolucency | Intraosseous Lipoma    |

Fig. 5. Photomicrography showing mature adipose tissue (H&E; ×40).

Returning to the medical literature, we found 30 cases of this tumor in the jaws since 1948 (Table 1) [9]. IOL of the jaws, like long bone IOL, can develop between (20–65) years old and the lesions were most commonly found in the fourth and fifth decades [10]. Usually, IOL is asymptomatic and some authors even have stated that the intraosseous lipoma belongs to the “leave me alone” group of bony lesions and that any invasive method is unnecessary, although swelling, pain, chin numbness, hypoesthesia, trismus, and periodontal signs are also reported, that depends on the location and size of the lesion [7]. Intraosseous mandibular lipomas mainly arise in the posterior region of the mandible [1] [10]. In radiological X-rays they appear as clearly defined osteolytic lesions often with an osteosclerotic border and usually a single, soft, well-defined and slowly growing radiolucent lesion [6], but diagnosis of IOLs of the jaws is very difficult when based only on clinical and radiological features [7]. Thus, the definitive test that confirms the nature of the lesion is the result of histopathological examination [6].

4. Conclusion and literature review

Since the first case reported in 1948 only 30 cases have been reported. Almost all of them were uniclar radiolucent lesion (83.33 %), most cases had clear boundaries, but 16.66 % of cases did not specify the nature of the lesion border. The main location was body and posterior region of mandibular with percentage (66.66 %) followed by: Anterior mandibular (23.33 %) and maxillary (10 %). Although patients ranged between 20 and 65 (most commonly in 4th, 5th and 6th decades with 20 % per decade). However, there were 3 cases (10 %) of patient’s ages below normal. Previously, the incidence of males was more than females, but after reviewing all archived cases in the medical literature, it was found that the incidence of females is higher (18 cases out of 30, with percentage 60 %). Pathological diagnosis: 21 cases were lipoma (70 %), 4 cases were angiolipoma (13.33 %), 3 cases were fibrolipoma...
According to Milgram: the transformation of Lipoma to Liposarcoma should not be excluded, but the rate of Liposarcoma is mainly very low (1 case per 2.5 million people) [11], and the percentage of IOL is also low (0.1 %) and less than all of that is IOL in the jaws (30 cases).

Thus, the ratio of transformation of IOL in the jaws to Liposarcoma may be excluded so far due to the paucity of archived cases.

We conclude from this that we need to record and archive each case of IOL in the jaws to have a comprehensive idea of the transformation of this type of tumor towards malignancy, and the manifestations of the transformation clinically, radiologically and histologically. In addition, we have a lot of questions that we don’t have an exact answer to yet. For example:

1. Was the long-term follow-up (more than 5 years) done for all cases of IOL and monitoring for any manifestations of transformation, recurrence and metastasis?
2. Were all archived lesions encapsulated?
3. Does the presence of the capsule affect the transformation to malignancy or not?
4. Does the sub-diagnosis of lipoma affect the rate of transformation or not?

These and other questions necessitate that we archive and write down everything we know about this type of tumor to increase medical knowledge about all these mysterious points and more.

Consent

Written informed consent was obtained from the patient for publication of this case report and accompanying photographs.

Sources of funding

We have no sponsors.

Ethical approval

Ethical approval has been exempted because this is a case report and no new studies or new techniques were carried out.

Registration of research studies

This paper is case report. The authors don’t need to register this work.

Guarantor

Corresponding author is the guarantor for this case report.

Provenance and peer review

Not commissioned, externally peer-reviewed.

Patient perspective

Patient was quite happy with the surgical procedure and overall treatment.

CRediT authorship contribution statement

Nabil Kochaji: diagnosis of case and reporting, and paper writing.
Suleiman Alhessani: literature review and paper writing.
Abdul Hadi Drbaa: surgery part of the work and following up patient.

Declaration of competing interest

Authors have no conflicts of interest.

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