Actinomycotic Osteomyelitis of the Mandible: A Case Report

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Actinomycosis is rare, chronic disease caused by Actinomyces spp. and once mistaken for fungal infection [1]. Actinomyces spp. is Gram-positive anaerobic bacteria, including Actinomyces israelii, A. bovis, A. odontolyticus, and A. viscosus. They can be found in calculus, gingival crevices, infected dentin and commonly live in pharynx and tonsil [2]. Actinomyces has no pathogenic potential when exposed to normal skin or healthy mucosa. Once losing membrane continuity by trauma or dental surgery, Actinomyces can invade into mucous membrane and form localized abscess [3].

Actinomycosis can be divided in three categories: cervicofacial (55%), abdominopelvic (20%), thoracic (15%) depending on the region [4]. Cervicofacial actinomycosis is the most common type; it can spread in acute form to hypopharynx, larynx, lacrimal duct and gland, major salivary gland, oral mucous, mandible, paranasal sinus and scalp; it can be chronic condition forming granuloma or multiple sinus tract and fistula [5]. Periapical actinomycosis, less invasive subtype of cervicofacial actinomycosis, has been reported, too [6].

Pus culture and bacterial examination is needed for diagnosis. Culture of Actinomyces spp. require specific oxidation-reduction condition associated with microbiome. In clinic, diagnosis of actinomycosis can be difficult even in highly suspected. Several months of antibiotics are required of treating actinomycosis and it can recur if treatment is stopped early. Making early definite diagnosis is important for preventing recurrence of actinomycosis [7].

There are few case reports of actinomycosis and even much rare case reports of osteomyelitis by Actinomyces except for medication related osteonecrosis of jaw. This report is about a
A 80-year-old man presented with mandibular incisor pain and swelling. Pain has been for 20 days and there was no surgery, trauma history and no underlying systemic disease. In physical examination, right cervical lymph node enlargement and multiple fistula in mandibular lateral incisor, canine, first premolar and first molar were observed. Elective pulp test on those teeth were all positive and no suspected teeth involvement was observed in panorama (Fig. 1) and periapical view. Clinically, actinomycosis was suspected and pus culture and bacterial examination were held but failed to culture actinomyces. Computed tomography (CT) and biopsy were performed for diagnosis. Irregular destruction on mandibular incisor labial bone was observed in CT (Fig. 2) and histological examination confirmed actinomycotic osteomyelitis. Patient was admitted to the emergency room presenting with severe swelling on right lower face, pain and limited mouth opening during administration of prescribed amoxicillin under cooperating with division of infectious diseases. In enhanced neck computed tomography, there was abscess pocket showing irregular contrast enhancement in right submasseteric space and pterygomandibular space (Fig. 3). Drainage on pus pocket and curettage of bone on incisor area were performed under local anesthesia.

Meanwhile, chest computed tomography was performed as routine ward laboratory examination despite of no symptom, observing tuberculosis-like lesion in left lower lobe in chest X-ray (Fig. 4A). Considering patient’s history,
disseminated actinomycosis was suspected, left lower lobectomy was performed under general anesthesia in department of cardiothoracic surgery. It was finally confirmed actinomycosis by histopathologic examination. Patient was prescribed doxycycline over 10 months after discharge. After 2 years, patient visited for another reason with no sign of recurrence of actinomycosis. CT showed healing state of mandibular bone (Fig. 5).

DISCUSSION

After first reporting of actinomycosis, misunderstood to fungal infection in 1845 by Van Langenbeck, Bollinger et al. assumed that Actinomyces bovis were fungus breaking out ‘lumpy jaw’ in cow. Wolff et al. succeeded in isolation culture from human body in 1891 and Waksman concluded that Actinomyces are Gram-positive anerobic bactera, not fungus in 1960s [8].

Cervicofacial actinomycosis is usually associated with soft tissue of chin and neck area and make multiple fistula on skin with swelling. It is characterized by features that infection spreads ignoring anatomical plane. Infected skin has abscess with purplish red and indurated, wood-like area. Special sulfur granule is observed in fistula drainage. Since another fistula appears when the former fistula disappears, extensive scarring and facial deformity in prolonged duration of untreated patient with actinomycosis.

Actinomycotic osteomyelitis is rare, which accounts for about 12% of total osteomyelitis [9]. Cervicofacial actinomycotic osteomyelitis usually occurs in mandibular body, following symphysis, ramus, mandibular angle. Maxilla and temporomandibular joint are comparatively resistant for the disease and prevalence is a ratio mandible:maxilla of 4:1. It is assumed that mandible is more commonly affected than maxilla due to the lesser vascularity and the denser cortical bone, similar to osteoradionecrosis [10]. In this case, co-infection of cervicofacial and thoracic type was shown by invasion of symphysis and lung.

Actinomycyes in oral flora in patient with poor oral hygiene could be inhaled to lung through the pharynx. It could be cause of thoracic actinomycosis. Thoracic actinomycosis forms micro-abscess and focal consolidation in parenchyma, fibrosis of local tissue. It was shown like tuberculosis, infiltration, malignancy in chest PA [11]. Thoracic
Actinomycosis cannot be diagnosed without surgical intervention, while cervicofacial actinomycosis can be treated by removal of fistula and long-term antibiotics. Biopsy is required to rule out pulmonary malignancy especially [12].

In this case, thoracic type actinomycosis was likely to occur because of long-time untreated condition and inhalation of abscess with poor oral hygiene due to mouth opening limitation because of abscess pocket of masticator space. To prevent thoracic actinomycosis, meticulous oral hygiene is essential in cervicofacial actinomycosis patient.

Pathophysiology of actinomycotic osteomyelitis has not clearly defined yet. Actinomyces can be found in tonsil, sulcus fluid, mucus membrane, infected dentin and tooth extrated area but cannot be infectious by itself. Actinomycotic osteomyelitis may be shown in the circumstance with broken balance of normal flora and pathologic change by chronic inflammation for some reason. Actinomyces itself is less invasive and toxic than other pathogens. Another bacteria help actinomyces to invade tissue with production of toxin and enzyme. These polymicrobial conditions make proper environment for growing anaerobic bacteria having low oxido-reduction potential [13]. Not forming sulfur granule of actinomyces in laboratory environment, it suggests that invasion of host immune system and colonization of associated bacteria are cornerstones to actinomycosis [14]. Once infection established on bone, the host makes inflammation response, suppuration, forming granuloma, fibrosis and making fistula.

Actinomycosis is shown clinically like tuberculosis and granuloma mimicking malignant lesion. Actinomycotic osteomyelitis usually proceeds slowly, but there were severe cases that infection invaded to mid-face, skull base, and cranium [15], and that proceed so rapidly that mandible was fractured, requiring reconstruction in immunosuppressive patient [16]. It is very important to diagnose actinomycosis, but it is reported that usual bacterial examination can confirm actinomycosis in less than 50% and first visit examination in less than 10% [17]. Four case series of actinomycotic osteomyelitis of Sezer et al. [18] reported to fail to make definite diagnosis of actinomycosis by usual microbiological examination. If the examination are failed repeatedly and delayed, it can get more inaccurate by effect of antibiotics. In this case, biopsy is highly recommended for diagnosis [19]. Actinomyces are strongly stained by H&E and Periodic acid-Schiff, Giemsa (GMS), especially filaments are well-shown in GMS [20].

In this case, there was no collection of actinomyces in first test. After several test, penicillin-resistant Streptococcus pneumonia, S. parascuinius, candida was found but not found actinomyces at all. Definite diagnosis is performed by biopsy of mandible and lung. When actinomycosis is highly suspected clinically, biopsy helps more accurate diagnosis, as well as scrutinizing microbiological test.

It can be treated long-term antibiotic application in case of cervicofacial actinomycosis localized in soft tissue, but when osteomyelitis is accompanied as this case, surgical intervention is essential. If acute abscess formation and necrotic tissue exists, surgical treatment is also required to distribute antibiotics more effectively. Antibiotic treatment for actinomycosis usually consists of 2-6 weeks of penicillin antibiotics by intravenous injection and 2-12 months of amoxicillin oral administration after IV [21]. Tetracyclin, minocycline, erythromycin, cephalosporin, sulfonamide, streptomycin can be used if patient has allergy for penicillin. In case of penicillin-resistant bacteria, clindamycin can be used by oral administration [22]. In this case, penicillin-resistant S. pneumonia, S. parascuinius is detected, ceftriaxone and vancomycin was prescribed intravenously. Doxycycline was also prescribed through oral administration over 10 months after discharge.

In conclusion, early and thorough diagnosis, appropriate surgery, long-term administration of antibiotics is required for actinomycotic osteomyelitis. When actinomycosis is clinically suspected, biopsy is recommended that cannot be diagnosed by typical microbiological culture test.

**CONFLICT OF INTEREST**

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

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