A CASE REPORT: LUDWIG’S ANGINA IN A PATIENT WITH SYSTEMIC LUPUS ERYTHEMATOSUS (SLE) AND FAMILIAL MEDITERRANEAN FEVER (FMF)

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Abstract

Ludwig’s angina is potentially life-threatening, rapidly spreading involving bilaterally submandibular, sublingual and sub mental spaces. It is characterized by induration of mouth, swelling, elevation of tongue and airway obstruction. There is three main points with early diagnosis which are airway management, intravenous antibiotic therapy and surgical intervention. We present a case of Ludwig’s angina in a 14-year-old boy. The etiology, management and potential complications of Ludwig’s angina are discussed.


Keywords: Ludwig’s angina, systemic diseases and, maxillofacial infection.

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Introduction

Ludwig’s angina is potentially life-threatening, rapidly spreading involving bilaterally submandibular, sublingual and sub mental spaces.¹ Ludwig’s angina was described by Wilhelm Frederick von Ludwig in 1836 as a rapidly progressive, gangrenous cellulitis and edema of the soft tissues of the neck and floor of the mouth.² It is characterized by brawny induration and elevation of the tongue with the airway obstruction.³ Mortality rates for Ludwig’s angina exceeded 50% in the pre-antibiotic era.⁴ As a result of antibiotic therapy, mortality recently averages approximately 8 %.⁵-⁶ Mortality is a result of upper airway obstruction.⁷-⁸

Odontogenic infection is the major etiologic factor of the Ludwig’s angina and primarily resulting from infection of the second and third molars.⁶-⁹ The roots of these teeth penetrate the mylohyoid ridge and because of that any dental infection or abscess has gateway to the submaxillary space.¹⁰ Infection spreads contiguously to the sublingual and submental space, posteriorly to the parapharyngeal and retropharyngeal space.¹¹-¹² Other etiologic factors are mandibular fractures, oral lacerations, infection of an oral malignancy, peritonsillar or parapharyngeal abscesses and submandibular sialadenitis.¹⁰ Predisposing factors include dental caries, recent dental treatment, systemic illness such as malnutrition, diabetes mellitus, compromised immune system such as immunodeficiency syndrome (AIDS), organ transplantation and trauma.¹³-¹⁴

Case Report

A 14-year-old boy was presented to our clinic complaining of severe painful tissues to the touch, diffuse swelling, difficulty swallowing. In his medical history, systemic lupus erythematosus (SLE) and Familial Mediterranean Fever (FMF) were present. The patient was febrile (temperature 38.4 °C), blood pressure was 123 / 72 mm Hg, and he was seemed uncomfortable, anxious, toxic appearance (Figure 1a). The patient’s erythrocyte sedimentation rate (ESR) was 27 mm/h and
white blood cells (WBCs) were 15,3 x10³/µL. The patient weighted 55 kg, heighed 170 cm and had a bodily mass index of 19.

In the radiographic examination, bilateral first molar teeth were detected as etiologic factor of Ludwig’s angina (Figure 3a).

During the intraoral examination, floor of mouth induration caused elevation of the tongue and floor of mouth was palpated that filling with the purulent matter (Figure 2a).

submandibular and sublingual space was widespread swelling, fluctuating and erythema. The patient was diagnosed as a Ludwig’s angina.

Corticosteroids are used to due to SLE. Therefore antibiotic prophylaxis was recommended for 3 days before the surgery. The patient had received twice a day with crystallized penicillin 800,000 units intravenously.

The patient was prepared for surgical intervention, provided asepsis and antisepsis. Extraoral and intraoral incision was made to drain of pus. Extraoral drainage was made the most fluctuant point of left mandible and intraoral, first lingual aspect of incisor teeth and bilateral buccal sulcus of premolar teeth. Blunt scissors was introduced to open up the tissue spaces and pus was drained. The wound was irrigated with normal saline and drain was placed and fixed to the skin with silk sutures. Postoperatively, crystallized penicillin intravenously, non-steroid anti-inflammatory were prescribed for 3 days. Postoperative irrigation was done within the drain and likewise intraoral three drainage point. Subsequently, drain was taken three days later and also infected right, left first mandibular molar teeth were extracted in the fifth day after mouth opening was provided and curettage was done. During this time, the patient was recommended that applied hot application over swelling surface. Every day for one week, 15th day, 1st month and 2nd month, the patient were followed and observed that swelling, erythema, pus, tongue elevation disappeared, sufficient mouth opening (Figure 2b).

Discussion

Ludwig’s angina generally has a dental origin and developing from mandibular molar teeth infection, periapical abscess (70-90 %). Although one report has shown that second or third molars caused infection and continued down the mylohyoid line of the mandible into the submaxillary space, our patient’s infection originated at the bilateral first mandibular molar teeth.

One third of cases associated with systemic diseases like diabetes mellitus, compromised immune system illnesses (AIDS, HIV+), lupus erythematosus, neutropenia, glomerulonephritis and aplastic anemia. Maxillofacial spatial infections are complicated in patients with compromised immune system.
such as SLE. Our patient had FMF and treatment of lupus erythematosus so that used corticosteroids and ferrous glycine sulphate complex and B vitamins complex.

Submandibular space is main infection region of the Ludwig’s angina. This space subdivided by the mylohyoid muscle into the sublingual space superiorly and the submaxillary space inferiorly. This space bounded superiorly by the floor of mouth mucosa and inferiorly by the superficial layer of deep cervical facia. Submandibular space infection can spread posteriorly to the retropharyngeal space and then mediastinum and causing mediastinitis.

Treatment of the Ludwig’s angina should include airway control, intravenous antibiotics and surgical intervention. A patient with Ludwig’s angina primarily should be secured airway management. In a retrospective review, 10% of children with Ludwig’s angina needed airway control, whereas 52 % of patients > 15 years of age underwent tracheostomy. In our patient, there is no need to endotracheal intubations, tracheostomy or monitoring for airway control. Intravenous antibiotics provide to treat and limit the spreading of infection. Antibiotics used before the antibiogram results, have been penicillin G intravenous, aminoglycosides, metronidazole, clindamycin, gentamicin. Steroid therapy has been suggested as a mean of reducing soft tissue swelling and edema and diminishing the potentiality for the need of a surgical airway in Ludwig’s angina. In the surgical intervention, all patients underwent incision and drainage. Surgical drainage requires large incisions exposing infected spaces and drains are inserted postoperatively.

Conclusion

Ludwig’s angina is a potentially lethal cellulitis of the submandibular, sublingual and submental spaces which was largely fatal during pre-antibiotic era. Airway control, aggressive intravenous antibiotic therapy and surgical decompresion and debridement are the prior treatment approach. As a medical therapy, our patient took ferrous glycine sulphate complex and corticosteroids due to his systemic disease. In Ludwig’s angina, patients who have systemic diseases like our patient, need antibiotic prophylaxis and consultation.

Declaration of Interest

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References