

An Uncommon Co-Occurrence: A Case Report of Laryngeal Actinomycosis and Vocal Cord Carcinoma with Literature Review

Dr O. Qassab^{1*}, Pr. M. Hmidi², Pr A. Kessab², Pr. K. Nadour²

¹Otolaryngology - Head and Neck Surgery Department, Hassan II University Hospital of Fez, Morocco

²Otolaryngology - Head and Neck Surgery Department, Military Hospital of Moulay Ismail, Meknes, Morocco

DOI: <https://doi.org/10.36347/sjmcr.2026.v14i04.064>

| Received: 07.06.2025 | Accepted: 31.07.2025 | Published: 29.04.2026

*Corresponding author: Dr O. Qassab

Otolaryngology - Head and Neck Surgery Department, Hassan II University Hospital of Fez, Morocco

Abstract

Case Report

Background: Cervicofacial actinomycosis (CFA) is a rare but treatable infection typically associated with poor dental hygiene, immunosuppression, or prior mucosal injury in the upper aerodigestive tract. Although rare, when cervicofacial actinomycosis involves the larynx, it may be misinterpreted as a mucosal mass. Moreover, reports of association with laryngeal carcinoma are exceptionally rare in the literature. **Case presentation:** A male patient, in his fifth decade, presented with laryngeal carcinoma accompanied by actinomycosis, which was confirmed through biopsy. The patient received antibiotics to treat the infection and was subsequently referred for radiotherapy, as he declined surgical intervention for his carcinoma. His condition showed a favorable clinical outcome during follow-up. **Conclusion:** The simultaneous occurrence of laryngeal carcinoma and actinomycosis is exceedingly rare, making it a particularly unusual clinical scenario. Clinicians should be vigilant in evaluating patients with laryngeal carcinoma for signs of actinomycosis, as timely recognition and appropriate management of this rare co-occurrence can significantly impact patient outcomes and prevent misdiagnosis or delayed treatment.

Keywords: Actinomycosis; granulomatous, laryngeal; vocal cord; carcinoma; radiotherapy; antibiotics.

Copyright © 2026 The Author(s): This is an open-access article distributed under the terms of the Creative Commons Attribution 4.0 International License (CC BY-NC 4.0) which permits unrestricted use, distribution, and reproduction in any medium for non-commercial use provided the original author and source are credited.

BACKGROUND

Cervicofacial Actinomycosis is a chronic, suppurative granulomatous inflammation caused by *Actinomyces israelii* [1], a filamentous gram-positive organism belonging to the genus Actinobacteria. It is a normal commensal in humans in the oral cavity, lower gastrointestinal tract and female genital tract [1]. Based on the infection site, Actinomycosis can be classified into several categories: Cervicofacial actinomycosis, Thoracic or respiratory tract actinomycosis - usually due to aspiration of infected material-, Abdominal actinomycosis -commonly affecting the appendix, cecum, and liver- and Pelvic actinomycosis, often linked to complications from intrauterine contraceptive devices [2]. In rare instances, the infection may also involve the joints, skin, or central nervous system [2]. The incidence of actinomycosis in humans is relatively rare, with most major medical centers diagnosing approximately one case per year, leading to an overall incidence of about one in 300,000 [3].

CASE PRESENTATION

We report the case of a 49-year-old male with a smoking history who presented to our ENT department with hoarseness and dysphonia evolving for the last 10 months, along with intermittent dyspnea on exertion. Physical examination concluded poor dental hygiene with several carious teeth but no suspect lesion on the oral mucosa, no cervical lymphadenopathy or goiter. Fiber optic endoscopy revealed a mass on the posterior half of the right vocal cord, extending into the subglottic space, while both vocal cords had normal mobility. For further characterization of the mass, a computerized tomography (CT scan) with contrast injection was ordered [Figure 1A-1B].

The patient underwent direct laryngoscopy under general anesthesia. The procedure identified a tumor on the right vocal cord extending into the subglottic region [Figure 2]. The anterior and posterior commissures, both ventricles and ventricular bands, and the left vocal cord were unaffected. Multiple biopsies were taken and histopathology confirmed squamous cell carcinoma (SCC) with concurrent actinomycosis [Figure 3-4].

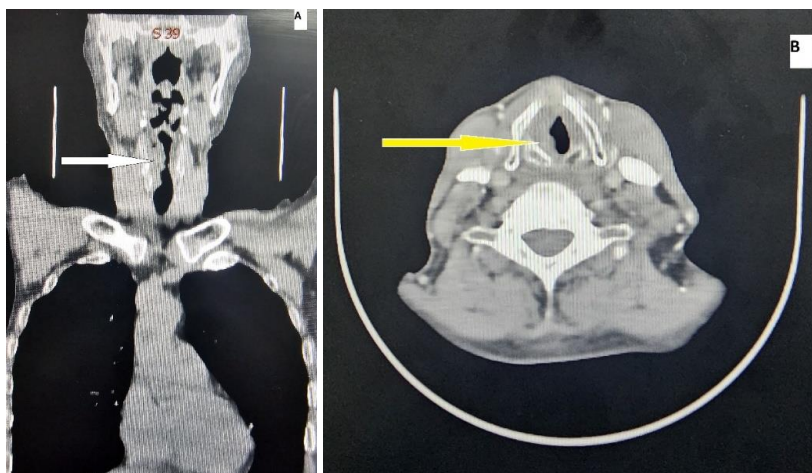


Figure 1: Cervical CT scan on coronal (A) and axial (B) view showed a tumor on the right vocal cord extending into the subglottic region

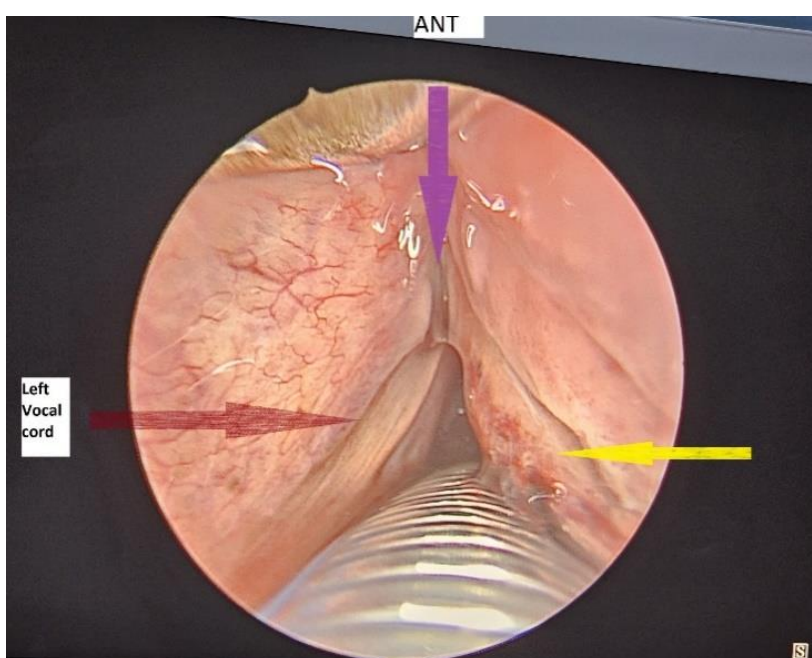


Figure 2: an endoscopic image showing a mass in the posterior half of the right vocal cord (yellow arrow)

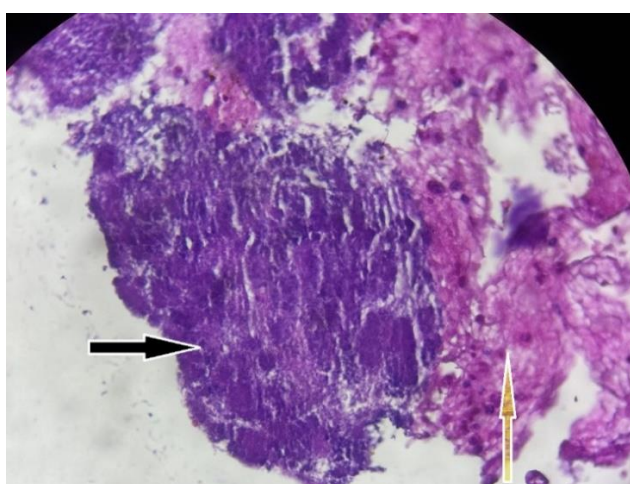


Figure 3: characteristic sulfur granules (black arrow) with a “sunburst” appearance are seen in the purulent discharge (yellow arrow) (Splendore–Hoeppli reaction)

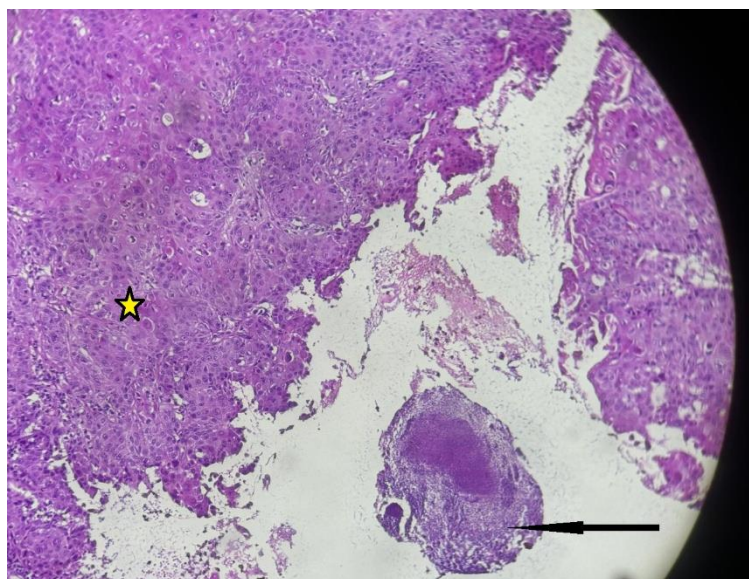


Figure 4: Sulfur granules (arrow) surrounded by a dense inflammatory cell infiltrate (star)

Regional and distant metastases were excluded through CT scan of the neck, thorax, and abdomen. The tumor was staged as T2N0M0. The patient refused surgery for his carcinoma and was referred for radiation therapy along with antibiotic treatment (amoxicillin with a dosage of 500 mg three times a day during 8 weeks). During the follow-up period, he remains alive and free of recurrence of both laryngeal carcinoma and actinomycosis. The patient's episodes of care timeline are reported in figure 5.

DISCUSSION

Actinomycosis is an exceptionally rare condition, with laryngeal involvement being even more uncommon [3-4], as fewer than 30 cases have been reported in the literature [5]. Adding to its rarity, the association of actinomycosis with an initial diagnosis of laryngeal carcinoma has been reported in only three other cases in the literature [2], making our case the fourth documented instance worldwide. This highlights the exceptional nature of such occurrences and the uncommon intersection of these conditions.

Actinomycosis was first identified by Israel in 1878, and the causative germ, *Actinomyces israelii*, was isolated by Wolfe in 1891 [1]. This gram-positive anaerobe has been linked to CFA [1]. CFA is often referred to as a "great masquerader" in head and neck diseases [6].

The pathogenesis of laryngeal actinomycosis remains unclear. It is not well understood why *Actinomyces* species, normally found in the mouth as a commensal saprophyte of the oral flora, become pathogenic. Some factors such as poor dental hygiene, oropharyngeal damage and trauma, diabetes, immunodeficiency, steroid therapy and malnutrition [7] are believed to predispose individuals to this condition. Consequently, we hypothesize that in our patient, dental

caries along with his laryngeal carcinoma may have compromised the normal mucosal barrier, facilitating the onset of the infection. In fact, a review of the literature indicates that laryngeal actinomycosis is frequently associated with a history of squamous cell carcinoma of the larynx and subsequent radiation therapy [8]. Additionally, cases have been reported in patients with systemic lupus erythematosus (SLE) and those who are immunosuppressed following renal transplantation [9]. However, actinomycosis can arise without any identifiable risk factors, with around 25% of patients having no known predisposing conditions [5].

Clinically, patients with actinomycosis may experience a range of symptoms including dysphonia, dysphagia, dyspnea, stridor, cough, odynophagia, pharyngitis or a sensation of a foreign body or heaviness in the pharynx. They may also present with general symptoms such as weight loss and fever, depending on the infection's location. In our case, the infection involved the vocal cord, which is the most frequently affected site, occurring in 58.3% of reported cases [5]. This is followed by the anterior and posterior commissures and the vestibular folds, each involved in 20.8% of cases. The aryepiglottic fold is affected in 16.7% of cases, while the epiglottis and pyriform sinus each appear in 4 cases. There are also isolated reports of invasion into the thyroid cartilage and subglottic region [5].

The clinical presentation of actinomycosis is nonspecific and may mimic conditions such as laryngeal carcinoma [10], papilloma [11], radionecrosis or recurrent carcinoma [12]. Thus, it is crucial to differentiate actinomycosis from these other conditions through biopsy. Differential diagnoses might also include abscesses, congenital anomalies, tuberculosis, or fungal infections [13]. While imaging findings are not specific and blood cultures and serologic tests are not

particularly useful for diagnosis [14], a biopsy and microscopic examination of the specimen remain the most effective and sensitive methods for confirming actinomycosis [15]. Microscopically, typical sulfur granules can be observed amidst the suppurative inflammation. These granules consist of conglomeration of filamentous gram-positive bacilli that are ensconced in a biofilm-like substance; they appear within the center of suppurative lesions. The bacilli are surrounded by eosinophilic amorphous material with a club-shaped configuration, a phenomenon known as the Splendore–Hoeppli reaction [2,5]. This reaction features clusters of bacteria surrounded by radiating, intensely eosinophilic material, which is believed to represent deposits of antigen-antibody complexes and debris from host inflammatory cells [2,5].

The treatment of actinomycosis infection depends on the extent of the lesions. Laryngeal actinomycosis generally responds well to prolonged penicillin or tetracycline therapy, with complications being rare but sometimes necessitating surgical intervention [13,15,16]. Excisional biopsy combined with penicillin-based therapy is often considered the definitive treatment approach [13]. However, surgery remains a critical option for advanced cases, involving thorough incision and drainage of abscesses, and wide excision of necrotic tissues, fistulous tracts, and affected bone [13,15,16]. For our patient, medical therapy alone was deemed sufficient, as the excision performed during the biopsy effectively addressed most of the lesion.

Table 1: Summarizes All Laryngeal Actinomycosis Cases Reported in the Literature to Date

Authors	Year	Country	N° Of Cases	Age/ Sex	Location	Risk Factors/ Symptoms	Treatment	Outcome
<i>J.H. Brandenburg and al</i>	1977	USA	1	67/M	Subglottic +/- tonsillar	None / seven-month history of progressive dyspnea, hoarseness, and nocturnal stridor with weight-loss	Excision+ long term antibiotics (2 million units of penicillin IU every four hours for two weeks and continued on 800,000 units daily for a total of six months)	Full recovery
<i>S.O. Shateen and al</i>	1983	United Kingdom	1	45/M	right cricothyroid area and vocal cord	Removal of molar tooth / recurrent sore throats, swelling	IV penicillin was given 1 megaunit 6 hourly for 23 days, and then oral penicillin 2 megaunits daily for 10 days	complete resolution and a normal repeat tomogram
<i>R.A. Hughes and al</i>	1984	USA	1	66/M	Pyrimform sinus, aryepiglottic fold, hypopharyngeal wall	Diabetes/ Pharyngitis, stridor dysphagia over 5 days and weight loss	Cephalexin for 4 months	Full recovery
<i>D.H. Tsuji</i>	1991	Japan	1	68/M	Left glottic area	radiation therapy for laryngeal cancer / sore throat, odynophagia, and hoarseness with fatigue and slight dyspnea.	Penicillin G 10 million units per day for 40 days f	full recovery

Batur çalis and al		L. Artesi and al	G. Lozano and al	M.A. Syed	M. Yasuda and al	S.H. Fernandez	M. Moreno and al
2006	2006	2004	2001	2000	1999	1997	
Turkey	Italy	Spain	United Kingdom	Japan	Asia	Spain	
2	1	1	1	1	1	1	
45/M	75/M	53/M	74/M	53/M	30/F	52/F	
right ventricle and both vocal cord	Epiglottis + left aryepiglottic fold	Vocal cord	vocal folds and arytenoids + post cricoid space	Anterior commissure	Vocal cord (secondary pulmonary localization)	Posterior commissure and vocal cord	
Smoking + Concurrent carcinoma with radiotherapy treatment / hoarseness, dyspnea and dysphagia	Smoking + alcohol / 2-month history of worsening dysphagia complicated by slight dyspnea	Smoking/ Dysphonia	course of radical radiotherapy for a TINOM0 squamous cell carcinoma of his left vocal fold / a two-month history of dysphonia and progressive dysphagia	T cell leukemia / Dysphonia	None/ Dysphonia for 02 months	Diabetes/ hoarseness and Cough	
Total laryngectomy (TL) + bilateral modified radical neck dissection (MRND) type 3 + primary voice restoration + palliative chemotherapy (carcinoma od esophagus)	IV penicillin (5 MIU/ die) for 15 days followed by oral treatment with clindamycin (600 mg) for 4 months	Penicillin 600,000 units orally every 6 hours for 3 weeks	amoxycillin 500 mg three times daily (malignancy have been excluded by biopsies and histological examination)	Penicillin 20M IU daily for 35 days	IU penicillin for 06 weeks	Cefuroxime- axetil 250 mg twice a day for 3 weeks	
Still alive during follow-up with oncology department	Full recovery confirmed after 7 months follow-up	Full recovery	Slow but satisfactory recovery	Good response to treatment	Full recovery	Resolution of symptoms	
	no evidence of recurrence of laryngeal carcinoma (LC) or actinomycosis.						

<i>M. Wierbicka and al</i>	2012	Poland	1	22/M	Epiglottitis, aryepiglottic folds and postcricoid region	None/ discreet dysphagia and weight-loss for several months	IV penicillin and clindamycin	Full recovery
<i>M. Meidani and al</i>	2011	Iran	1	77/M	Larynx+ lung	Prolonged intubation (heart surgery / Fever, Cough, Weight loss and an unexplained respiratory failure	High dose of penicillin	Full recovery
<i>B. Khademi and al</i>	2011	Iran	1	14/M	Vocal cord	Recent tooth extraction/ Dysphonia worsening over 2 months	IV penicillin for 2 wks. Then PO penicillin for 3 months	Full recovery
<i>R. Schumann and al</i>	2010	Germany	1	56/M	Aryepiglottic fold and pyriform sinus	accidental ingestion of corn / dysphagia	Surgical excision of the abscess and amoxicillin and clavulanic acid for 3 weeks	Good response to treatment
<i>M. Sari and al</i>	2007	Turkey	1	21/M	anterior one-third part of the left vocal cord	None/ 6-month history of hoarseness of voice	amoxicillin-clavulanate 625 mg orally three times a day for 8 weeks.	Full recovery
<i>H.S. Sims ans al</i>	2007	USA	1	47/M	posterior glottic tissue	Systemic lupus erythematosus (prednisone therapy) + renal transplantation / hoarseness and dysphagia complicated with increased dyspnea with exertion even under initial treatment	IV penicillin for two weeks followed by oral penicillin for three months	Full recovery
<i>Sara and al</i>	2006	USA	1	69/M	Thyroid cartilage +Trachea (Co-occurrence with a laryngeal carcinoma)	History of smoking + Diabetes/ 8-month history of hoarseness, a 30-pound weight loss, trouble speaking, and a persistent cough	Neoadjuvant chemotherapy+ total laryngectomy with extended tracheal resection down to the 8th tracheal ring + IV ampicillin 500 mg every 6 hours for 12 weeks, then PO for 6 to 12 months	Well response to antibiotics
<i>M.C. Menezes</i>	2006	Brazil	1	77/M	left aryepiglottic fold	Smoker + alcoholic / mild throat pain for about a year + weight-loss	antibiotics for 21 days	Full recovery

M. Pujani and al	2017	India	1	28/M	Anterior commissure	History of microlaryngoscopic excision of a polyp 7 months before / Hoarseness and change in voice for previous 12 months	Amoxicillin therapy	Good response to treatment
V. Manimaran and al	2017	India	1	33/M	Anterior commissure and anterior one third of both vocal cords	NONE / Hoarseness for 03 months	cotrimoxazole (800/160mg) tablets twice daily for 04 months	Full recovery
Gogge and al	2016	USA	1	55/F	Vocal cord	Smoking/ hoarseness and intermittent dysphonia+ pharyngitis	amoxicillin 500 mg orally three times per day for one month	Full recovery
S. Patel and al	2014	USA	1	74/M	Vocal cord and Vestibular fold	chronic lymphocytic leukemia and febrile neutropenia/odynophagia and dysphagia	Ciprofloxacin and amoxicillin/clavulanic acid for 1 ^o month	near-complete resolution of his symptoms
F. Lensing and al.	2014	USA	1	24/M	Cricoid cartilage and adjacent fat pad	Oropharyngeal trauma/ Acute dyspnea and laryngeal pain 2 days after recent airway dilation	IV penicillin 6 weeks	resolution of hypermetabolic activity in the larynx on imaging and resolution of the soft tissue mass.
T. Abed and al	2013	United Kingdom	1	35/F	Anterior commissure	Systemic lupus erythematosus (prednisone therapy) + renal transplantation / Hoarseness	Excision of the abscess and 3-month course of oral penicillin V and referred for speech therapy	Full recovery
K. Yoshihama and al	2013	Japan	1	49/M	anterior one-third part of the left vocal cord	None / hoarseness of voice for 2 years	amoxicillin-clavulanate 62.5 mg orally three times a day for 8 weeks	Full recovery
T. Ferry and al	2012	France	1	67/M	Vocal cord and vestibular fold	Smoking+ chemotherapy, lateral pharyngolaryngectomy was performed, followed by radiotherapy for his Laryngeal carcinoma / Mild dyspnea on rest	Oral amoxicillin (6 g/day) for 3 months (relapse excluded by biopsies and histological examination)	No relapse occurred with full recovery

<i>H. Sei and al</i>	2018	Japan	1	14/F	Right arytenoid of the larynx	Bone marrow transplantation for acute lymphocytic leukemia 4 months before/ sore throat for 02 weeks	Oral amoxicillin 750 mg/day for 70 days	Full recovery
<i>B. Raggio and al</i>	2019	USA	1	93M	Anterior vocal folds bilaterally with anterior commissure	Chronic kidney disease/ 6month history of progressively worsening dysphonia and odynophagia	Renally dose amoxicillin for 03 weeks	Withdraw of treatment after diagnosis of omental carcinomatosis with lung metastasis

CONCLUSION

By presenting an additional case of vocal cord actinomycosis in this paper, we aim to highlight that actinomycotic infections can accompany and/or complicate head and neck carcinomas. The rarity of this combination underscores the importance of considering actinomycosis in patients with laryngeal carcinoma who present with atypical symptoms or complications. Given that actinomycosis can mimic or complicate the clinical picture of laryngeal cancer, awareness of this possibility is crucial for accurate diagnosis and effective treatment. Therefore, we seek to alert clinicians and pathologists to consider the possibility of such rare concurrent actinomycotic infections.

Patient’s perspective

I recently had hoarseness along with dysphonia so I went seeing an ENT specialist. In the ORL outpatient clinic, doctor performed a fibroendoscopy and discovered a tumor. I must say that I had intermittent breathing difficulty that I didn’t took seriously. I had a CT scan and the ORL told me they had to perform laryngoscopy under general anesthesia in order to take biopsies. My pathology results showed cancer with an infection for which my doctor prescribed antibiotics for 8 weeks. I refused surgery for my cancer, so my doctor referred me to an oncologist. I had radiation therapy and since then, my breathing and hoarseness had improved and my doctors confirmed that I am actually cancer free.

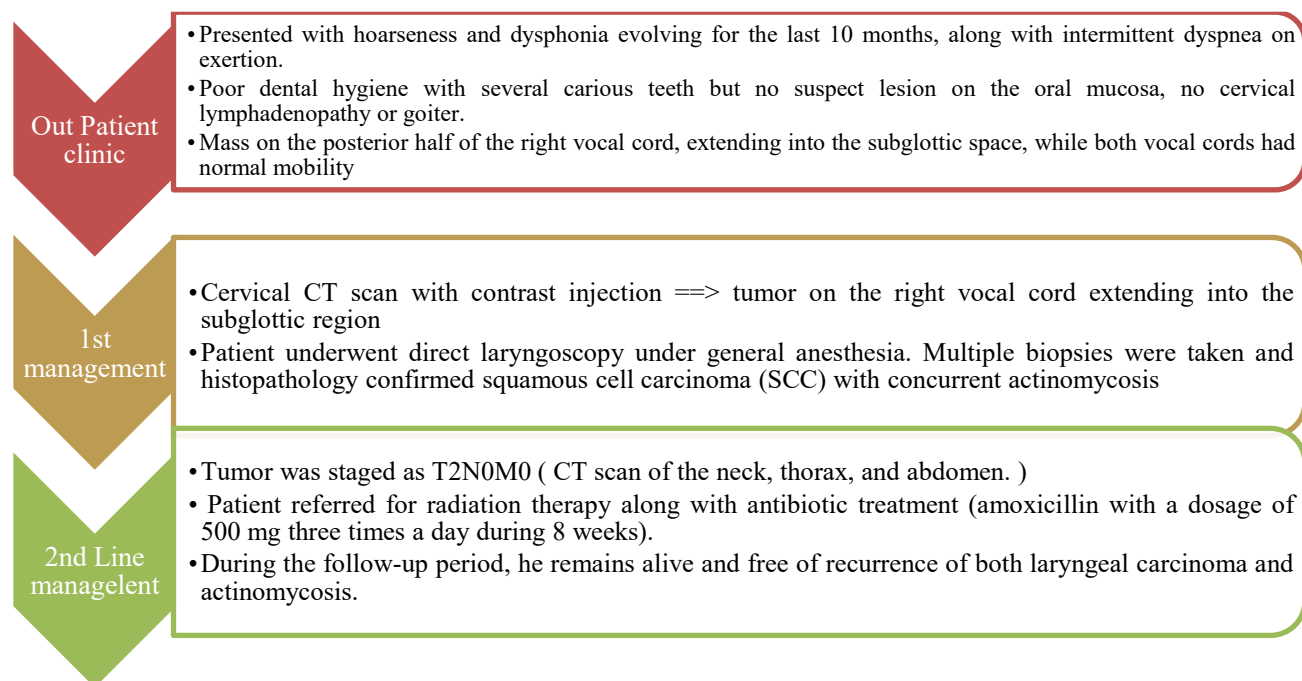


Figure 5: Patient’s case management timeline

Declarations:**Ethics approval and consent to participate:**

The IRB approved our study and consents to participate to the study were signed by the patients. Our IRB is CEHUF (comité d'éthique hospital-universitaire de Fès).

Funding: not applicable

REREFENCES

- Smego RA, Foglia G. Actinomycosis. *Clin Infect Dis* 1998 ;26(6):1255– 1261.
- Valour F, Sénéchal A, Dupieux C, Karsenty J, Lustig S, *et al.*, (2014) Actinomycosis: etiology, clinical features, diagnosis, treatment, and management. *Infect Drug Resist* 7: 183-197.
- Belmont MJ, Behar PM, Wax MK (1999) Atypical presentations of actinomycosis. *Head Neck* 21: 264-268.
- B. Khademi, S. H. Dastgheib-Hosseini, and M. J. Ashraf, "Vocal cord actinomycosis: a case report," *Iranian Journal of Otorhinolaryngology*, vol. 23, no. 2, pp. 49–52, 2011.
- Benjamin G, David K, Kim G, John S (2016) Laryngeal Actinomycosis: A Case Report and Systematic Review of 32 Cases in the Literature. *J Otolaryng Head Neck Surg* 2: 006.
- Rankow RM, Abraham DM. Actinomycosis: masquerader in the head and neck. *Ann Otol Rhinol Laryngol* 1978;87(2 Pt 1):230–237.
- I. Zajc, Z. Orihovac, and M. Bagatin, "Temporal actinomycosis: report of a case," *Journal of Oral and Maxillofacial Surgery*, vol. 57, no. 11, pp. 1370–1372, 1999.
- A. Batur Calis, A. E. Ozbal, T. Basak, and S. Turgut, "Laryngeal actinomycosis accompanying laryngeal carcinoma: report of two cases," *European Archives of Oto-Rhino-Laryngology*, vol. 263, no. 8, pp. 783–785, 2006.
- H. S. Sims and B. B. Heywood, "Post-transplant actinomycosis of the posterior glottis involving both vocal processes," *Otolaryngology—Head and Neck Surgery*, vol. 137, no. 6, pp. 967– 968, 2007.
- Keisuke Yoshihama, Yasumasa Kato, and Yuh Baba, Vocal Cord Actinomycosis Mimicking a Laryngeal Tumor, *Case Reports in Otolaryngology Volume 2013*, Article ID 361986, 2 pages <http://dx.doi.org/10.1155/2013/361986>
- Mukta Pujani, Sabina Khan, Sujata Jetley and Seema Monga, Laryngeal actinomycosis with dysplasia in a young male with a recurrent laryngeal polyp, *Tropical doctor* 0(0) 1–3, 2017, DOI: 10.1177/0049475516688161
- Ferry T, Buiret G, Pignat J-C, Chidiac C. Laryngeal actinomycosis mimicking relapse of laryngeal carcinoma in a 67-year-old man. *BMJ Case Reports* 2012;10.1136/bcr-2012-007084
- Hughes RA, Paonessa DF, Conway WF (1984) Actinomycosis of the larynx. *Ann Otol Rhinol Laryngol* 93:520–524
- Carpenter JL, Artenstein MS (1976) Use of diagnostic microbiologic facilities in the diagnosis of head and neck infections. *Otolaryngol Clin North Am* 9(3):611–629
- Syed MA, Ayshford CA, Uppal HS, Cullen RJ (2001) Actinomycosis of the post-cricoid space: an unusual case of dysphagia. *J Laryngol Otol* 115:428–429
- Pillsbury HC, Sasaki CT (1982) Granulomatous diseases of the larynx. *Otolaryngol Clin North Am* 15(3):539–551