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CASE REPORTS

MICROBIOLOGY

Fabrizio Cialente, et al.: Running title

Laryngeal tuberculosis in renal transplant recipients: A case report and review of the literature

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ABSTRACT

Renal allograft recipients are at greater risk of developing tuberculosis than the general population. A woman with a kidney transplant was admitted to the emergency department with high temperature, dysphonia, odynophagia, and asthenia. The final diagnosis was laryngeal tuberculosis. Multidisciplinary collaboration enabled accurate diagnosis and successful treatment. Laryngeal tuberculosis should be considered in renal allograft recipients with hoarseness. A more rapid diagnosis of tuberculosis in renal transplant recipients is desirable when the site involved, such as the larynx, exhibits specific manifestations and the patient exhibits specific symptoms. In these cases, prognosis is excellent, and with adequate treatment, a complete recovery is often achieved.

KEYWORDS: Laryngeal tuberculosis; kidney transplant; laryngeal leukoplakia

INTRODUCTION

Tuberculosis (TB) is an infectious disease caused by Mycobacterium tuberculosis [1]. Geographically, most TB cases in 2018 were in Southeast Asia (44%), Africa (24%), and the Western Pacific nations (18%); smaller numbers of cases have occurred in eastern Mediterranean nations (8%), the Americas (3%), and Europe (3%) [1]. In 2018, the number of new cases of rifampicin-resistant TB was approximately half a million, of which 78% were multidrug-resistant TB [1]. Although the prevalence of TB is low in Italy, its epidemiology is changing. Since 1955, more than 160,000 people in Italy have died from this potentially preventable and curable disease [10]. In 2018, the estimated incidence of TB in Italy was 4300 new cases (7 cases per 100,000 population), and the estimated number of deaths from TB in Italy was 370. The number of TB case notifications in 2018 in Italy was 3912 (71% with pulmonary involvement) [1].

Although the lung is the most common site of involvement, a number of extrapulmonary organs, including the larynx, can be involved as well. Laryngeal TB is an uncommon condition and has been rarely reported in the literature [1,2]. Nevertheless, it represents a public health concern [3] and may appear similar to malignancy on imaging and laryngoscopy [4,5].

Renal allograft recipients are at greater risk of developing tuberculosis (TB), commonly atypical and extrapulmonary, than is the general population. The prevalence of TB in renal transplant recipients ranges from 1% to 4% in developed Western countries [6-9].

We present the case of a kidney transplant recipient who developed laryngeal TB. This case represents a rare condition.

CASE REPORT

A 38-year-old woman with chronic renal failure caused by chronic diabetes mellitus from the age of 9 was admitted to the emergency department of the Sapienza University Hospital, Rome, Italy, in December 2018 with high temperature, dysphonia, odynophagia, general discomfort, and severe asthenia. She reported having undergone a thoracotomy with right-sided pneumectomy at the age of 16 for previous pulmonary aspergillosis and renal transplantation in 2015 at the age of 35 and

subsequent treatment with steroids, mycophenolate mofetil, and tacrolimus. Laryngeal and pulmonary mycosis were initially suspected, and she was transferred to the Department of Infectious Diseases and treated with antimycotic drugs.

A neck and pulmonary computed tomographic (CT) scan showed asymmetry of the glottic space with marked contrast enhancement of the epiglottis up to the false right vocal fold; submucosal imbibition up to the thyroid cartilage, which appeared to be inhomogeneous with pseudo-focality; and right-sided level II lymphadenopathy with a maximum lymph node diameter of 12 mm. We also found right-sided lung atelectasis with ipsilateral mediastinal displacement (as a result of the earlier thoracotomy with right pneumectomy) and hyperplasia of the left lung with interstitial thickening. Otolaryngologic examination with high-definition fiber optic laryngoscopy (Figure 1) showed swelling of the right false vocal fold with a whitish lesion; motility and respiratory space were normal. Endoscopy of the larynx with narrow-band imaging confirmed that the lesion was not malignant.

The patient underwent microlaryngoscopy under general anesthesia, and a biopsy sample of the whitish lesion was obtained. Histologic examination (Figure 2) revealed that the mucosa was lined by respiratory epithelium free from atypia. Lymphocyte-rich inflammatory infiltrates, along with multiple small, nonnecrotizing granulomatous aggregates of epithelioid histiocytes (CD68R+), were detected in the lamina propria. Ziehl-Neelsen and periodic acid–Schiff staining did not yield conclusive findings. Subsequent microbiologic bronchoalveolar lavage and sputum tests confirmed the diagnosis of pulmonary TB with extrapulmonary laryngeal extension. The patient was then isolated for 30 days and treated daily with 200 mg of isoniazid, 400 mg of ethambutol, 500 mg of pyrazinamide, 400 mg of moxifloxacin, and 300 mg of vitamin B_6 (Benadon). As of this writing, the patient is not isolated and undergoes regular follow-up visits with clinical examinations and CT scans every 2 months.

DISCUSSION

Renal transplant recipients at high risk for different infections because they must receive immunosuppressive therapy [11]. Using the keywords "laryngeal TB" and "kidney transplant," we searched PubMed for articles written in English from 1990 and on about the occurrence of laryngeal TB in kidney transplant recipients. Inclusion criteria were kidney transplantation and laryngeal TB, and we included only articles in which laryngeal involvement occurred before or at the same time as pulmonary involvement. We reviewed five reports about a total of eight patients [9-10,12-13]. In these cases, laryngeal symptoms were prominent (dysphonia, hoarseness) and directed the diagnosis. We excluded one article in which the authors diagnosed laryngeal TB after the occurrence of pulmonary TB (in a patient with cavitary pulmonary TB who was found to have laryngeal TB 1 year after discontinuing antituberculous drugs) [6]. In our review, we evaluated the following parameters: clinical manifestations, results of diagnostic tests, and treatment. In these patients (Table 1), the main symptoms were hoarseness, dysphonia, odynophagia, cough, and fever, and biopsy samples of tissue revealed granulomatous inflammation. The treatment of TB may be complex, and drug resistance often develops [14,15]. Standard therapeutic agents include isoniazid, rifampicin, and pyrazinamide, although modifications of therapy because of cost have been reported in the literature [10]; neither of the patients were given rifampicin because it would have necessitated an increase in the cyclosporine dose and, consequently, the cost of therapy. Since laryngeal TB in renal transplant recipients is an atypical presentation of TB, the diagnosis may be delayed, which can lead to increased morbidity and exposure of the patients' contacts, particularly through nosocomial transmission. The mechanism of laryngeal involvement is postulated to be through bronchogenic spread, with infected sputum from the lungs directly contaminating the larynx. However, this theory has been challenged by other authors [9], who suggested that hematogenous spread is responsible for laryngeal seeding. Immunosuppressive drugs promote reactivation of latent TB and acquisition of new disease. In many studies [10-12], the main symptoms that forced patients to seek medical attention were severe

fever and severe odynophagia, although patients had already had other symptoms, such as dysphonia and asthenia, for weeks. The differential diagnosis must include laryngeal TB. Larynx assessment with advanced endoscopy and narrow-band imaging enables diagnosis [16]. We used endoscopes capable of narrow-band imaging to rule out suspected malignant lesions of the larynx because our institution is a referral center for the treatment of laryngeal cancer in Italy. Whitish lesions of larynx could occur in many types of diseases, ranging from benign hyperkeratosis to squamous cell carcinoma. Other clinical conditions included in the differential diagnosis are sarcoidosis, Wegener's granulomatosis, syphilis, histoplasmosis, blastomycosis, coccidioidomycosis, cryptococcosis, amyloidosis, traumatic polypoid granulation, and laryngeal papilloma [16-18]. Thus, biopsy may be performed for differential diagnostic purpose. In two cases in the literature, standard treatment was isoniazid, rifampicin, and pyrazinamide, although both patients required threefold increases in the daily cyclosporine dosage to maintain therapeutic levels. Because of the drug interaction between rifampicin and cyclosporine, the dosage of cyclosporine must be increased substantially when it is coadministered with rifampicin [19, 20]. The results of both randomized and nonrandomized studies support the value of isoniazid as TB prophylaxis in renal transplant recipients at risk for active infection [21]. Clinicians should consider prophylaxis in renal transplant recipients in endemic areas or in recipients in nonendemic countries who are at risk. We treated our patient with isoniazid because she was at risk for active infection, according to the literature [22-25]. However, the evidence of the benefit of isoniazid prophylaxis in renal transplantation is not robust, and a large multicenter trial is needed to investigate isoniazid prophylaxis in kidney transplantation in an endemic area [26].

Laryngeal TB is rare in kidney transplant recipients who developed TB. The few cases reported in the literature indicate that laryngeal symptoms (dysphonia, dysphagia) in such patients must be assessed adequately, even in industrialized countries in which TB is not widespread. In these cases, prognosis is excellent and recovery is often achieved. When whitish lesions are found in the larynx, diagnostic tests for TB must be performed.

CONCLUSION

Laryngeal TB should be considered in renal allograft recipients with hoarseness. A more rapid diagnosis of TB in renal transplant recipients is desirable when the site involved, such as the larynx, exhibits specific manifestations and the patient exhibits specific symptoms. In these cases, prognosis is excellent, and with adequate treatment, a complete recovery is often achieved.

REFERENCES

- 1. Global tuberculosis report 2019. https://www.who.int/tb/publications/global_report/en/
- 2. Kurokawa M, Nibu K, Ichimura K, Nishino H. Laryngeal tuberculosis: A report of 17 cases. Auris Nasus Larynx. 2015;42(4):305–310.
- 3. Zavod MB. Laryngeal tuberculosis: A public health concern. Ear Nose Throat J. 2015;94(10-11):E51–E52.
- 4. El Kettani NE, El Hassani M, Chakir N, et al. Primary laryngeal tuberculosis mimicking laryngeal carcinoma: CT scan features. Indian J Radiol Imaging 2010;20(1):11–12.
- 5. Suhail A, Ahmed MS, Sobani ZU, Ghaffar S. Laryngeal tuberculosis presenting as laryngeal carcinoma. J Pak Med Assoc. 2012;62(2):167–168.
- 6. Yildiz A, Sever MS, Türkmen A, Ecder T, Beşişik F, Tabak L, Ece T, Kiliçarslan I, Ark E. Tuberculosis after renal transplantation: experience of one Turkish centre. Nephrol Dial Transplant. 1998;13(7):1872–1875.
- 7. Conversano M. Update on the epidemiology of tuberculosis in Italy. J Rheumatol Suppl. 2014;91:4–10. doi: 10.3899/jrheum.140096
- 8. Rosado Rubio, C., et al., Atypical localization of tuberculosis in kidney transplants. Nefrologia. 2011;31(5): 622–624.
- 9. Lui, S.L., et al., Tuberculous infection in southern Chinese renal transplant recipients. Clin Transplant. 2004;18(6):666–671.
- 10. Jha, V., et al., Laryngeal tuberculosis in renal transplant recipients. Transplantation, 1999. **68**(1): p. 153-5.
- 11. Lim MA, Kohli J, Bloom RD. Immunosuppression for kidney transplantation: Where are we now and where are we going? Transplant Rev (Orlando). 2017;31(1):10–17.
- 12. Tato, A.M., et al., Laryngeal tuberculosis in renal allograft patients. Am J Kidney Dis. 1998;31(4):701–705.

- 13. Poniachik, J., et al., Tuberculosis in kidney transplant patients. Rev Med Chil. 1990;118(10):1105–1110.
- 14. Hunter, A.M., et al., The changing pattern of laryngeal tuberculosis. J Laryngol Otol. 1981;95(4):393–398.
- 15. Wang M, Wei H, Zhao Y, et al. Analysis of multidrug-resistant bacteria in 3223 patients with hospital-acquired infections (HAI) from a tertiary general hospital in China. Bosn J Basic Med Sci. 2019;19(1):86–93.
- 16. Kimza, H., Jackowska, J., and Wierzbicka, M. The usefulness of the NBI narrow band imaging for the larynx assessment. Otolaryngol Pol. 2018;72(3):1–3.
- 17. Parker, N.P., Vocal fold leukoplakia: incidence, management, and prevention. Curr Opin Otolaryngol Head Neck Surg. 2017;25(6):464–468.
- 18. Fang, T.J., et al., Classification of vocal fold leukoplakia by clinical scoring. Head Neck. 2016;38 Suppl 1:E1998–E2003.
- 19. Vachharajani, T., et al., Diagnosis and treatment of tuberculosis in hemodialysis and renal transplant patients. Am J Nephrol. 2000;20(4):273–277.
- 20. Campana, C., et al., Clinically significant drug interactions with cyclosporin. An update. Clin Pharmacokinet. 1996;30(2):141–179.
- 21. Naqvi A, Akhtar S, Ahmed E, Noor H, Saeed T, Akhtar F, Rizvi A. Use of isoniazid chemoprophylaxis in renal transplant recipients. Nephrol Dial Transplant. 2010;25(2):634–637.
- 22. Higgins RM, Cahn AP, Porter D, Richardson AJ, Mitchel RJ, Hopkin JM, Morris PJ. Mycobacterial infections after renal tranplantation. Q J Med 1991;78:145–153.
- 23. EBPG. Expert Group on Renal Transplantation. European best practice guidelines for renal transplantation. Section IV: Long-term management of the transplant recipient. IV.7.2. Late infections. Tuberculosis. Nephrol Dial Transplant. 2002;17(Suppl 4):39–43.

- 24. John GT, Thomas PP, Thomas M, Jeyaseelan L, Jacob CK, Shastry JC. A double-blind randomized controlled trial of primary isoniazid prophylaxis in dialysis and transplant patients. Transplantation 1994;57:1683.
- 25. Vachharajani T, Abreo K, Phadke A, Oza U, Kirpalani A. Diagnosis and treatment of tuberculosis in hemodialysis and renal transplant patients. Am J Nephrol. 2000;20:273.
- 26. Currie, A.C., Knight, S.R., and Morris, P.J. Tuberculosis in renal transplant recipients: the evidence for prophylaxis. Transplantation. 2010;90(7):695–704.

FIGURES AND TABLES

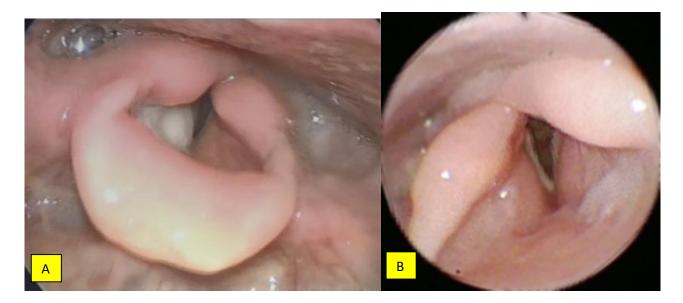


FIGURE 1. Laryngoscopic images. (A) Before antitubercular therapy. Swelling of the right false vocal fold with a whitish lesion, normal motility, and respiratory space. (B). After antitubercular therapy. Normal motility and structure of larynx, with normal respiratory space.

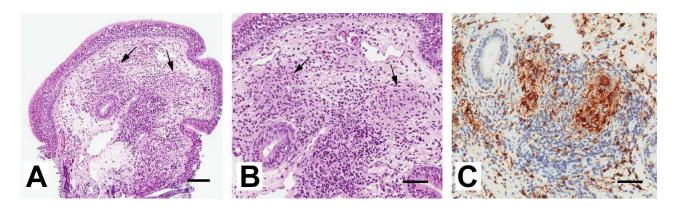


FIGURE 2. Eosin and hematoxylin staining of the sampled mucosa under low- (A) and high-power (B) magnification illustrates the lymphocyte-rich inflammatory infiltrates along with multiple small, nonnecrotizing granulomas (arrows in A and B). The immunohistochemical stain for CD68R (C) highlights the small granulomatous aggregates of histiocytes. Bars represent 100 μm in A and 50 μm in B and C.

TABLE 1. Cases of laryngeal tuberculosis reported in the literature

Subject of	No. of	Clinical	Diagnostic tests	Treatment
study	patients	manifestation		
Laryngeal	2	Prolonged	Histological study of	No different from
tuberculosis in		hoarseness and	laryngeal biopsy sample	tuberculosis elsewhere.
renal		painful	revealed large numbers of	Neither patient was
transplant		dysphagia	inflammatory cells, scattered	given rifampicin
recipients [6]			epithelioid cells, and	because it would have
			histiocytes along with a few	necessitated an increase
			multinucleated giant cells and	in the cyclosporine
			a few areas of necrosis. Ziehl-	dose and, consequently,
			Nielsen staining was strongly	the cost of therapy
			positive for acid-fast bacilli.	
Laryngeal	2	2-week febrile	Direct laryngoscopy and	Isoniazid, rifampicin,
tuberculosis in		illness with	sputum results were positive	and pyrazinamide in
renal allograft		hoarseness and	for acid-fast bacilli that	each case; both patients
recipients [7]		dysphagia	subsequently grew	required threefold
			Mycobacterium tuberculosis	increases in daily
				cyclosporine dosage to
				maintain therapeutic
				levels.
Atypical	1	Dysphonia with	Direct laryngoscopy with	Isoniazid, rifampicin,
localization of		whitish lesions	laryngeal biopsy revealed	and pyrazinamide for 2
tuberculosis in		on arytenoid	tuberculoid granulomatous	months and isoniazid
kidney		cartilage and	chronic laryngitis. Ziehl-	and rifampicin for 9
transplant		epiglottis	Neelsen staining and	months
recipients [4]			Löwenstein test yielded	
			positive results. Acid-fast	

			bacilli were demonstrated in	
			sputum	
Tuberculous	2	Hoarseness of	Direct laryngoscopy with	Both patients received
infection in		voice, fever,	laryngeal biopsy, test of	isoniazid, 300 mg/day;
southern		cough	sputum, bronchoalveolar	rifampicin, 450–600
Chinese renal			lavage, and chest radiography	mg/day; and
transplant			were performed. Laryngeal	pyrazinamide, 1.5–2
recipients [5]			biopsy revealed typical	g/day. Pyrazinamide
			caseating granulomata and	was stopped after 3
			acid-fast bacilli	months
Tuberculosis in	1	Fever, dysphonia	Direct laryngoscopy with	
kidney			laryngeal biopsy	
transplant			demonstrated acid-fast bacilli	
recipients [8]			in sputum	