



REVIEW

Characteristics of Laryngeal Osteosarcoma: A Critical Review

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ABSTRACT

Laryngeal sarcomas constitute an extremely rare entity among head and neck malignancies. Furthermore, most of them are chondrosarcomas, and the osteogenic form remains a true rarity. In general, there is a lack of information on the characteristics of laryngeal osteosarcoma. Thus, we sought to critically review the existing world literature on laryngeal osteosarcoma in order to develop a more accurate clinicopathological profile of this malignancy.

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Laryngeal osteosarcoma has a predilection for elderly male patients, as 87% were male in the present series and the mean age was 62 years (range 32–80), and without a direct association with tobacco exposure. Osteosarcoma of the larynx is typically a highly malignant neoplasm that metastasizes early, has a propensity for hematogenous spread and also has a marked tendency to recur. Twelve (41%) out of the 29 cases in the present review with follow-up data had metastatic disease. The aspects that distinguish osteosarcoma from its differential diagnostic alternatives are discussed in this review.

Keywords: Larynx; Osteosarcoma; Radiotherapy; Sarcoma; Surgery

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Key Summary Points

Laryngeal sarcomas constitute an extremely rare entity among head and neck malignancies.

Osteosarcoma of the larynx is typically a highly malignant neoplasm that metastasizes early, has a propensity of hematogenous spread and also has a marked tendency to recur.

A full photomicrographic documentation is essential for the diagnosis and registration of osteosarcoma cases.

The diagnosis of this highly malignant tumor would preferably warrant an international consultation network including a registry of rare head and neck neoplasms.

INTRODUCTION

Despite being the most common primary malignant neoplasm of the bony skeleton, osteosarcoma is one of the rarest malignant mesenchymal neoplasms of the larynx. Sarcomas encountered in the larynx are usually chondrosarcomas. Therefore, even physicians

in referral-based academic centers of pathology or otorhinolaryngology–head and neck surgery may encounter only a single patient with a true laryngeal osteosarcoma in their entire career. Of note, we searched the REFCOR database (réseau d’Expertise Français sur les Cancers ORL Rares [French Network of Rare Head and Neck Tumors]) in order to find cases to report in the present review. The REFCOR has been registering rare head and neck cancers, i.e., rare cancers of the aerodigestive tract excluding squamous cell carcinoma and undifferentiated nasopharyngeal carcinoma, and cancers of the salivary glands, ear, nasal cavities, and sinus, since 2008. Out of the 7530 patients registered up to August 2019, only 54 osteosarcoma cases were diagnosed, and none of these was located in the larynx.

Laryngeal osteosarcoma was first described by Jackson and Jackson in 1942 without histological documentation [1]. In 1946, Clerf et al. reported eight cases of sarcoma of the larynx, and one of these was classified as fibro-osteosarcoma, but the photomicrograph of the tumor was not convincing [2]. The first well-documented case of laryngeal osteosarcoma was described by Sprinkle et al. in 1966 [3]. Of note, in 1974, the textbook *Tumors of the Head and Neck: Clinical and Pathological Considerations* by Batsakis concluded that no authentic case of osteosarcoma in the upper respiratory tract, including the larynx, had been reported at that time [4]. In 2017, Bennion et al. reported 33 laryngeal osteosarcomas having been reported in the English literature since 1942, but many of the reports listed in their review, including their own cases, did not present pathological documentation [5].

It is noteworthy that morphological features of sarcomas typically represent a heterogeneous variation in morphology. This will lead to challenging differential diagnostic considerations, especially when the obtained biopsies remain small. In contrast to a low-grade chondrosarcoma—in which a limited biopsy specimen may still leave open the question of the malignant nature of that lesion—it is usually quite apparent for the pathologist that a laryngeal osteosarcoma specimen represents a high-grade malignancy. More importantly, the

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precise subclassification of the malignancy will need to be determined.

The existing reports of laryngeal osteosarcoma cases do not consistently include pathological documentation that would allow critical evaluation of the final pathological diagnosis. Therefore, our primary aim was to create a current series of all the published laryngeal osteosarcoma cases by critically evaluating the published reports of this rare malignancy and to include only those cases with adequate pathological description of the findings. The secondary aim was to obtain comprehensive information regarding the characteristics of these neoplasms to enable a discussion of possibilities for enhancing their diagnostic accuracy.

METHODS

We conducted a literature search in the PubMed database using the following keywords: “Laryngeal” OR “Larynx” AND “Osteosarcoma”. The search covered all years from the origin of the database until 31 December 2019. The Preferred Reporting Items for Systematic Review and Meta-Analysis (<https://www.equator-network.org/>) methodology was applied. A

total of 61 hits were retrieved (Fig. 1). Of these, 30 publications were relevant as having reported a new case/s of laryngeal osteosarcoma (N = 33) that met the inclusion criteria of providing proper pathological evaluation and photomicrographic documentation of the finding. Cases that did not meet these inclusion criteria were excluded. This article is based on previously conducted studies and does not contain any studies with human participants or animals performed by any of the authors.

RESULTS AND DISCUSSION

Clinical Features of the Reported Cases in the Literature

Many reports indicate that this neoplasm occurs in men in their sixth and eighth decades of life, and that the lesion is not directly associated with alcohol or tobacco exposure [6, 7]. Laskin et al. and Sheen et al. reported two patients with a history of radiotherapy prior to the development of laryngeal osteosarcoma [8, 9]. These tumors usually arise from the vocal cord and cricoid cartilage. Therefore, the most common symptoms are hoarseness, dyspnea, and obstruction.

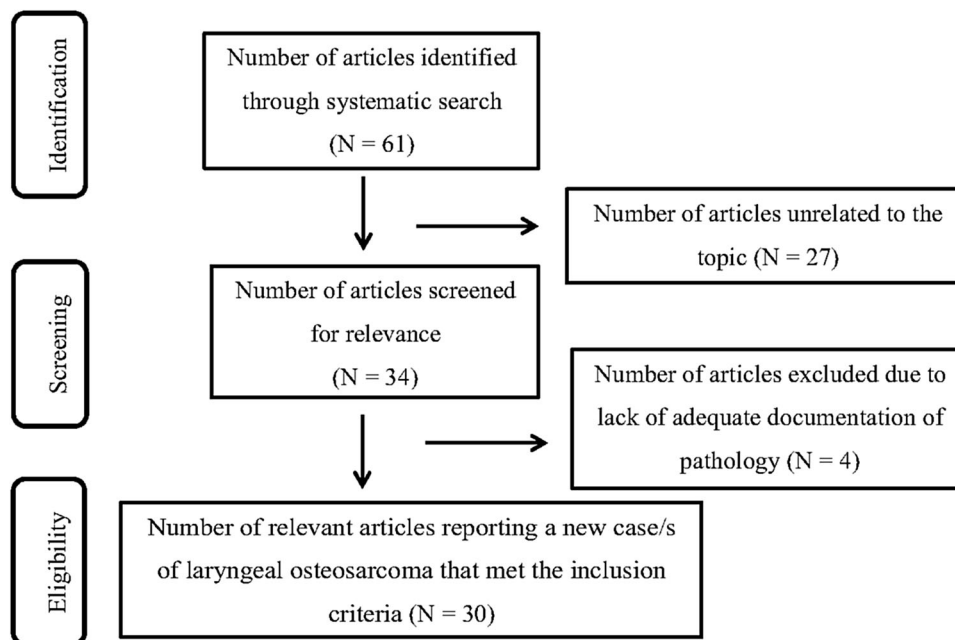


Fig. 1 Flow chart describing the PRISMA search

The tumors included in this review and with information on the exact location ($N = 22$) involved either subglottic [2], supraglottic [3] or transglottic [2] structures, vocal cords [10], or thyroid and/or cricoid cartilages [6].

Osteosarcoma is typically a highly malignant neoplasm that metastasizes early, with a propensity for hematogenous spread and also a marked tendency to recur. There are reports of metastases to regional lymph nodes, but these remain rare, and instead, hematogenous spread to the lungs is the main route. In the present review, 12 out of the 33 patients were diagnosed with distant metastases, most frequently in the lungs. Seventeen out of the 25 patients with follow-up data available in the published case reports had experienced persistent or early recurrent disease, typically within 1–2 years.

Pathology of Laryngeal Osteosarcoma

Macroscopically, the neoplasm is mainly polypoid or exophytic in appearance but may also mimic other sarcomas. The gross specimen is typically grainy and, in some areas, difficult or

impossible to cut with a scalpel because of its ossification. The tissue may have a variable color ranging from yellow to brown or red.

Microscopically, the tumor is composed of frankly malignant, spindle-shaped mesenchymal cells, associated with osteoid and immature neoplastic bone formation, as was shown in the reviewed reports (Fig. 2). The nuclei show considerable hyperchromasia or pleomorphism, and mitoses may be numerous. Atypical and bizarre giant cells and multinucleated osteoclast-like cells have been described. Venous invasions may be identified.

Immunohistochemical staining has shown that the neoplasm is positive for vimentin and negative for desmin, S-100 protein, cytokeratin (AE1/AE3, MNF-116, CK5/6, CK7, CK8, CK19, CK20), and epithelial membrane antigen. Both osteonectin and osteocalcin are expressed in osteogenic sarcoma but are not sufficiently specific. However, osteonectin and osteocalcin will highlight osteoid matrix and distinguish it from collagen matrix.

Electron microscopy studies [10–12] have shown cells resembling osteoblasts with

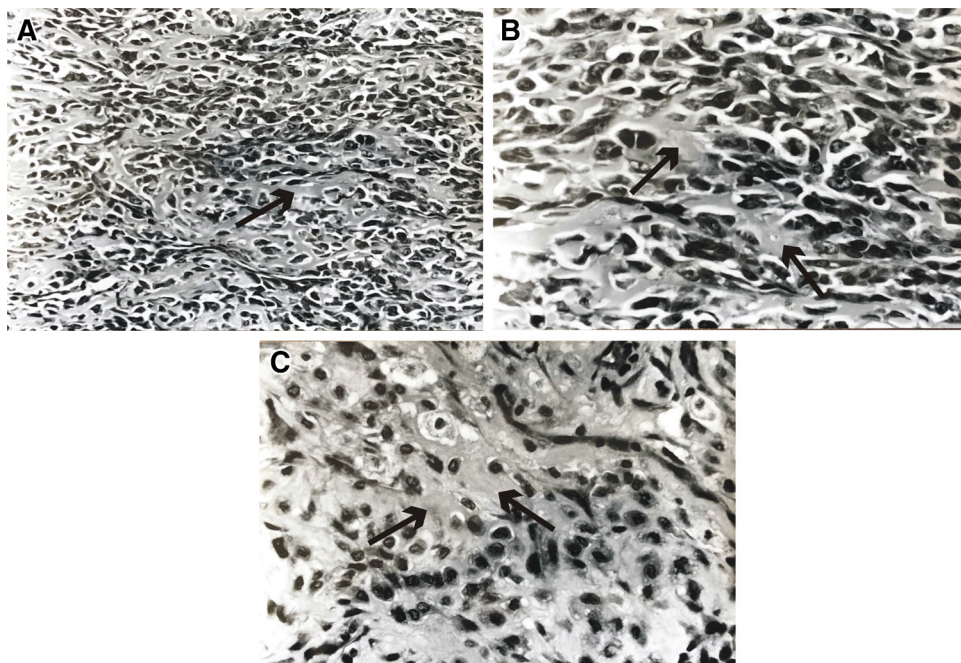


Fig. 2 **a** Laryngeal osteosarcoma with osteoid formations (arrow) produced by malignant cells (hematoxylin–eosin). **b** Osteoid formation (arrows) by the tumor cells is readily

apparent and confirms the diagnosis of osteosarcoma (hematoxylin–eosin). **c** Another area of the tumor-producing osteoid (arrows) (hematoxylin–eosin)

pleomorphic nuclei and nucleoli of variable size within the abundant cytoplasm containing numerous dilated cisternae of rough endoplasmic reticulum. The cells were enveloped by interlacing collagen fibers. Osteoblastic elements predominated but cartilaginous and fibrous tissue could also be found. No desmosomes or other evidence of epithelial differentiation was found in the tumor.

Diagnosis of Laryngeal Osteosarcoma

The definite diagnosis of osteosarcoma depends on the identification of osteoid production by malignant cells in the biopsy sample, as was shown in the reviewed reports. The differential diagnosis includes spindle-cell squamous carcinoma with osteoid differentiation, chondrosarcoma, malignant fibrous histiocytoma, fibrosarcoma, and myositis ossificans affecting the larynx. Spindle-cell squamous carcinoma may contain benign and rarely malignant osteoid areas [13]. Therefore, older reports of laryngeal osteosarcoma may represent spindle-cell squamous carcinoma with osteoid formation. Any bone-producing malignant mesenchymal lesion should at first be suspected as a

spindle-cell carcinoma. The presence of islands of carcinoma in situ or microinfiltrating carcinoma excludes the diagnosis of osteosarcoma.

In osteogenic sarcoma, clearly sarcomatous areas are present, and the “zonal phenomenon”—which is characterized by a central cellular area, an intermediate zone of osteoid formation, and a peripheral shell of highly organized bone—is absent. The phenomenon of formation of mature bone is most prominent in the periphery of the lesion and is an important criterion for differentiating myositis ossificans from osteogenic sarcoma. In this latter lesion, mature osteoids are located in the center of neoplasm, and neoplastic mesenchymal cells—that form the osteoids—are in the periphery.

Osteogenic sarcoma metastatic to larynx has also been reported [14, 15]. This unlikely event can be ruled out with clinical assessment of the patient.

Treatment and Prognosis

Wide local resection with clear margins, which typically warrants total laryngectomy, remains the treatment of choice for laryngeal osteosarcoma to achieve local control and long-term

Table 1 Key features of laryngeal osteosarcoma

Rarity
Demographics
Large lesions arising predominantly in men in their sixth to eight decades
Histology
High-grade sarcoma with area of osteoid production by the tumor cells at least focally
Chondroid elements may be present, as may areas reminiscent of fibrosarcoma, malignant fibrous histiocytoma, or sarcoma (not further subclassified)
Immunohistochemical staining can be utilized and will add to the histological diagnosis
Differential diagnosis
Spindle-cell squamous carcinoma with osteoid differentiation, chondrosarcoma, malignant fibrous histiocytoma, fibrosarcoma, and myositis ossificans
Prognosis
Laryngeal osteosarcoma usually results in death in less than 2 years from either extensive local disease or pulmonary metastases

Table 2 Cases of laryngeal osteosarcoma with pathological documentation

Institute	Authors	Year	Characteristics and treatment	Follow-up
University of Washington, Seattle, WA, USA	Mantilla et al. [38]	2019	75/M. Former smoker. Progressive hoarseness. Anterior commissure High-grade. No invasion of TC. No Metastasis. LE	Lost to follow-up
Rashid Hospital, Dubai	AbdullGaffar and Keloth [16]	2018	32/F. Non-smoker. Hoarseness for 1.5 months Subglottic, 1.5 cm. Small-cell OS	Lost to follow-up
Chandigarh, India	Bahl et al. [6]	2015	50/M. Hoarseness for 2 months Transglottic. No metastases. LE, thyroidectomy, partial pharyngectomy. CT	NED at 9 months
Nicolaus Copernicus University, Torun, Poland	Sawicki et al. [17]	2015	58/F. Dyspnea for 6 months Subglottic. Immobile VC. No metastases. LE	NED > 22 months
Tygerberg Hospital, Cape Town, South Africa	Mosalleum et al. [7]	2015	55/M. Former smoker (2PY) Dysphonia, dyspnea for 4 months. LE	Bilateral lung metastases at 8 months
Saitama Medical University, International Medical Center, Hidaka, Saitama, Japan	Kuba et al. [18]	2015	55/M. Non-smoker. Odynophagia Neck mass High-grade osteoblastic OS No metastases. Partial LE	NED at 20 months
Baroda, Gujarat, India	Bhatt et al. [19]	2014	38/M. Smoker. Hoarseness for 2 months. No metastases Near-total LE	NED at 15 months
Istanbul University, Turkey	Ulusan et al. [20]	2012	59/M. Smoker. Hoarseness and dyspnea. Anterior commissure Radiation-induced OS (RT 5 years earlier). Normal chest, abdominal, and bone radiographs. No metastasis LE	NED at 8 years

Table 2 continued

Institute	Authors	Year	Characteristics and treatment	Follow-up
Tabriz University Tabriz, Iran	Sanaat et al. [21]	2009	71/M. Smoker. True and false cords Dyspnea, dysphagia, and odynophagia Solid mass destroying both cords with supra- and subglottic extension Bone and lung metastases. LE	DOD at 12 months
Sisli Etfal Education & Research, Hospital, Department of Radiation Oncology, Istanbul, Turkey	Arslan et al. [22]	2008	69/M. Larynx. NOS. Persistent hoarseness. LE and postoperative RT. Neck recurrence 5 months later Lung and neck metastases	DOD
Hospital San Giovanni, di Dio e Ruggi d'Aragona, Via San Leonardo, Salerno, Italy	Mottola et al. [23]	2008	56/M. Non-smoker. TC, CC, and paratracheal space. Dyspnea, dysphagia, and dysphonia. Tumor mass affecting glottis and supraglottis. LE	DOD 3 months
Oddział Otolaryngologiczny Szpitala im. F. Raszei w Poznaniu	Rydzewki et al. [24]	2005	Bilateral neck and lung metastases LE, ND, and CT	DOD
University of Texas, Southwestern Medical Center, Dallas, TX, USA	Athré et al. [25]	2005	69/F. Progressive dyspnea. Larynx, strap muscles, and thyroid gland Dystrophic calcification outside laryngeal confines. Total LE, pharyngectomy, esophagectomy Lung metastasis	DOD at 10 months
Departments of ORL and Pathology Okmeydanı Training Hospital, Istanbul, Turkey	Topaloglu et al. [26]	2004	80/M. RVC, LVC, and ventricular bands. Hoarseness, progressive dyspnea. Soft tissue mass narrowing the airway. Extended LE	NED 15 months
M.D. Anderson Cancer Center, Houston, TX, US	Madrigal et al. [27]	2002	50/M. RVC. Hoarseness. Partial frontolateral LE 59/M. Smoker. TC. Hoarseness. Destructive lesion of the TC extending to parapharyngeal soft tissue. Total LE, CRT. Multiple lung metastases	NED at 24 months DOD 18 at months

Table 2 continued

Institute	Authors	Year	Characteristics and treatment	Follow-up
			60/M. Smoker. Polypoid mass of the RVC with anterior commissure extension. Dyspnea. LE and CRT	NED at 60 months
			Neck recurrence 10 months later	
			69/M. Smoker. True vocal cord	Lost to follow-up
			Dyspnea, dysphonia, and cough	
			Lung metastases	
Ohio State University, OH, USA	Rossi et al. [28]	1998	68/M. Former smoker	DOD at 3 months
			Wide-field LE. Lung metastases	
The Long Island Jewish Medical Center. The Long Island Campus of the Albert Einstein College of Medicine, New Hyde Park, NY, USA	Myssiorek et al. [29]	1998	47/M. Smoker. TC. Hoarseness, dyspnea, otalgia, hemoptysis	DOD at 12 months
			Destructive lesion of TC with extension into soft tissues.	
			LE. Bone metastasis	
University of Pittsburgh, Medical Center, Pittsburgh, PA, USA	Berge et al. [30]	1998	60/F. Non-smoker. CC. Dysphonia, dysphagia, and weight loss	NED at 44 months
			Expansion of the medullary portion of the cricoid cartilage with encroachment on the airway	
			LE and ND	
National Taiwan University Hospital, Taipei, Taiwan	Sheen et al. [9]	1997	56/M. RT for nasopharyngeal carcinoma 32 years earlier	DOD at 21 months
			Left neck and anterior TC	
			Hoarseness. Hyperdense mass in the left upper neck. LE	
Institute AFIP	Devaney and Ferlito [31]	1996	Pathology consultation case	Lost to follow-up
Hasharon Hospital, Golda Medical Center, Petah Tikva	Zohar et al. [32]	1996	Surgery	Lost to follow-up
Centre Hospitalier Universitaire, de Bordeaux, France	Pinsolle et al. [12]	1990	65/M. Former smoker. Dyspnea, dysphonia. 1.5 cm polypoid tumor involving the RVC and anterior commissure. RT	NED at 60 months

Table 2 continued

Institute	Authors	Year	Characteristics and treatment	Follow-up
			Recurrence after 1 month	
			Resection and pectoralis flap	
University of Liverpool, England	Van Laer et al. [33]	1989	75/M. LVC and left pyriform fossa	Recurrence at 13 months
			Dysphagia, discomfort in the throat	
			Amorphous mass of calcification. LE	
Pathologisches Institut Oldenburg, Germany	Remagen et al. [34]	1983	65/M. LVC. RT. Recurrent disease	DOD at 24 months
Mayo Clinic, Rochester, NY, USA	Gorenstein et al. [35]	1980	75/M. RVC and ventricle	DOD at 14 months
			Hoarseness 5 months. LE and ND	
			Local and distant metastases	
Mahidol University, Bangkok, Thailand,	Suchatlampong et al. [11]	1981	67/M. LVC, anterior and posterior commissure and supraglottis	DOD at 6 months
			Hoarseness and dyspnea. Mass involving the glottis with supraglottic extension. RT	
University of Texas, Southwestern Medical School, Dallas, TX, USA	Dahm et al. [10]	1978	79/M. RVC and anterior commissure	DOD
			Hoarseness and difficulty breathing	
			Polypoid mass occupying the airway	
			Multiple lung metastases. LE	
State Univ. of New York, Buffalo, NY, USA	Haar et al. [36]	1978	66/M. Smoker. Hoarseness. CC, posterior wall of the proximal trachea and displacing the esophagus anteriorly	DOD at 21 months
			Lung metastasis. LE	
Royal Victoria Infirmary, Newcastle, UK	Morley et al. [37]	1973	62/M. Hoarseness, dyspnea, acute respiratory obstruction. Occasions of loss of consciousness. Polypoid lesion of RVC	DOD at 3 months
			LVC with supraglottic extension	
			Subtotal LE and RT	

Table 2 continued

Institute	Authors	Year	Characteristics and treatment	Follow-up
Univ. of Virginia, Charlottesville, VA, USA	Sprinkle et al. [3]	1966	71/M. Hoarseness. Posterior surface of RVC. No metastasis. LE	NED > 24 months

F female, *M* male, *OS* osteosarcoma, *LE* laryngectomy, *ND* neck dissection, *Tx* treatment, *NED* no evidence of disease, *PY* pack years, *RT* radiotherapy, *CT* chemotherapy, *CRT* chemoradiotherapy, *DOD* dead of disease, *TC* thyroid cartilage, *CC* cricoid cartilage, *RVC* right vocal cord, *LVC* left vocal cord, *ND* neck dissection

survival. In the present cases, laryngectomy was performed for 26 out of the 29 cases with treatment-related data available. Adjuvant radiotherapy needs to be considered individually, as osteosarcomas are relatively radioresistant. Chemotherapy might improve the prognosis for patients with disseminated lesions, and it should be emphasized that in the results of the present review, the number of patients with distant metastasis was high (36%).

In the present review, five out of the 33 patients had no reported follow-up data. Eleven out of the 28 patients with follow-up data available were alive with no evidence of disease after a variable follow-up time (range 9 months to 8 years).

The important features of laryngeal osteosarcoma are summarized in Table 1. A summary of the reported laryngeal osteosarcoma cases is shown in Table 2, with the first case published in 1966 [3] and the last in 2019 [38]. Table 3 provides a description of the cases that have been reported without pathological documentation.

CONCLUSIONS

True laryngeal osteosarcoma remains an extremely rare malignancy. The true incidence of this entity is not known but based on the number of existing well-documented cases ($n = 33$) in the literature during 1966–2019, it is obviously not encountered at most of the institutions managing laryngeal tumors.

The diagnosis of this highly malignant tumor would preferably warrant an international consultation network including a registry of rare head and neck neoplasms. Electronic

Table 3 Reported cases of laryngeal osteosarcoma with no pathological documentation

Bennion et al. [5]	2017	48/F. Shortness of breath. Erythematous mass inferior to the level of the true vocal cords at the posterior commissure. The tumor appeared to arise from the cricoid cartilage in computed tomography imaging. No microphotography
Laskin et al. [8]	1988	56/M. Three years after radiotherapy. No follow-up data. No microphotography
Clerf et al. [2]	1946	51/M. Hoarseness and dyspnea. Later with local recurrence and lung metastasis. DOD 14 months. Pathological documentation not convincing
Jackson and Jackson [1]	1942	51/M. Hoarseness, dyspnea, and dysphagia. Cricoid cartilage. Local recurrence after 3 months. DOD after 6 months with mediastinal metastasis. No other data available

F female, *M* male, *DOD* dead of disease

platforms for such activities exist. Our review demonstrates that full photomicrographic documentation is essential for the diagnosis and registration of osteosarcoma cases. Furthermore, such a collaborative effort would enable multidisciplinary recommendations for individual management as well. Naturally, management of laryngeal osteosarcoma should be

centralized to a few experienced tertiary care academic centers.

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Compliance with Ethics Guidelines. This article is based on previously conducted studies and does not contain any studies with human participants or animals performed by any of the authors.

Data Availability. All data generated or analyzed during this study are included in this published article.

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