

# Prevalence of Head and Neck Sarcomas in the Main Health Centers in Yazd from 1994 to 2014

## Original Article

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## Abstract

### Introduction:

Head and neck sarcomas involve a group of rare malignant diseases with a high histological variability involving various anatomical sites that can lead to under-reporting of the true incidence of these neoplasms. This study aimed to epidemiologically investigate the occurrence of sarcomas of the head and neck within the past 20 years in Yazd, Iran (1994–2014).

### Materials and methods:

In this descriptive, cross-sectional and retrospective study, 16114 patient's records with malignant tumors were examined via the census method, which were available in the archives of 8 main treatment centers in Yazd, Iran within a 20-year period. Age, sex, occupation, habitat, type of sarcoma, tumor location and grade, metastasis, recurrence, and history of head and neck irradiation were recorded. Data were analyzed in SPSS software version 17.

### Results:

Among 586 cases of sarcomas, 59 cases (10.06%) were identified with head and neck sarcomas. The mean age of the patients was  $32.22 \pm 8.31$  years, of which 26 (44.01%) patients were males and 33 (55.9%) were females. Soft tissue sarcomas were noted in 41 cases (69.5%); rhabdomyosarcoma was the most common (27.1%). Eighteen (30.5%) patients had hard tissue sarcomas; osteosarcoma (15.3%) was the most common. Soft tissues of the head and neck were the most (49.20%) involved sites. Most sarcomas were low grade. In 5 patients (8.5%), metastases occurred to the head and neck, and the tumor relapsed in 16 patients (27.1%).

### Conclusion:

The findings of the current study were in agreement with those of other reports referred to in different studies. This suggests that the epidemiology of head and neck sarcomas in Yazd, Iran is similar to other geographical regions.

### Key words:

•Head and Neck Neoplasms •Sarcoma •Yazd

## Introduction

Sarcomas involve a rare and heterogeneous variety of malignant tumors of mesenchymal origin with a specific and distinct histopathology<sup>(1, 2)</sup>. The mesenchymal cells can develop into tumors affecting the soft tissues of muscle, fat, and fibrous tissue. Bone and nerves can also be involved.<sup>(2)</sup> Occasionally, these tumors are associated with trauma, genetic syndromes as well as exposure to previous radiation, though there is mostly no apparent cause. Pathological classification is most valuable in the treatment and prognosis of head and neck sarcomas.<sup>(3)</sup>

The incidence of sarcoma is more prevalent in children than adults. Approximately 1% of all adult cancers<sup>(2-4)</sup> and 10%–20% of pediatric cancers are sarcomas<sup>(2)</sup>. About 5%–15%<sup>(2, 3)</sup> of adult sarcomas are in the head and neck region; 35%<sup>(3, 5)</sup> of children are diagnosed with head and neck sarcomas. The findings of different studies have revealed that the incidence of sarcoma is more in men than in women, which is about 50% to 60%.<sup>(6-11)</sup>

Common soft tissue sarcomas in order of frequency include liposarcoma, malignant fibrous histiocytoma, fibrosarcoma, rhabdomyosarcoma, leiomyosarcoma, synovial sarcoma, malignant peripheral nerve sheath tumors, angio-sarcoma and kaposi sarcoma, whereas hard tissue sarcomas in order of frequency entail osteosarcoma, chondrosarcoma, as well as Ewing's sarcoma<sup>(12)</sup>. The results of some studies conducted in Iran demonstrated that 60% of sarcomas occurred in males with a mean age of 36 years, that the most common sarcomas in adults were malignant fibrous histiocytoma and synovial sarcoma and osteosarcoma and in children, osteosarcoma, Ewing sarcoma, and rhabdomyosarcoma in the were the most common sarcomas.<sup>(11, 13)</sup>

The ratio of bone sarcoma to sarcoma of soft tissue was 3:1 in patients aged under 16 years and 1:3 among adults<sup>(13)</sup>. In a study conducted on the epidemiology of soft tissue sarcomas in Shahid Sadoughi hospital of Yazd in 2005, the most common sarcomas were synovial sarcoma and malignant fibrous histiocytoma respectively among males and females. The study findings indicated that the sarcoma incidence in Yazd was similar to that of Western countries.<sup>(11)</sup>

Epidemiological studies provide vital informa-

tion that forms the basis of future research. Because investigating the prevalence of sarcomas in the head and neck has received scant attention in Iran, the present study is intended to provide the epidemiology of sarcomas of the head and neck over the past 20 years in Yazd, Iran (1994–2014) based on the histopathologic examinations.

## Materials and Methods

In this descriptive, cross-sectional, and retrospective study, 16114 patients records (case notes), diagnosed as malignant tumors were examined from the archive of the Shahid Sadooghi Dental School and several other hospitals (Shahid Sadoughi, Shahid Rahneemooon, Mojibiyani, Mortaz, ShohadayeKargar, Seyedoshohada and SavanehSookhtegi) over a 20-year period (1994–2014).

The records were obtained by proposing a research study from the Shahid Sadoughi medical university of Yazd (Ethical code: p.17.1.77710; date: 1393.4.22).

It should be noted that case records that were incomplete or cases in which the patients reported a written dissatisfaction were excluded from the study. In order to glean the study data, a checklist was devised consisting of the following variables: case record number, pathology department identification number, age, sex, occupation, place of residence, type of sarcoma, tumor location, tumor grade, the occurrence of metastasis, recurrence, history of head and neck irradiation. The patients' medical records as well as their pathology reports available in the mentioned health centers were collected and analyzed utilizing the SPSS software (Ver. 17) through descriptive statistics.

## Results

Out of 16,114 cases examined in this study, 586 cases (3.65%) of patients were diagnosed with sarcomas, among which 59 patients (10.06%) suffered from sarcomas of the head and neck, the study cohort. The mean age of the patients was  $32.22 \pm 8.31$  years with an age range of 1.5–83 years, of which 17 patients (28.8%) were <16 years of age, whereas 42 patients (71.2%) were > 16 years. A total of 26 patients (44.05%) were males and 33 (55.9%) were females. More than half of patients (59.3 %) lived in Yazd (Table 1). As demonstrated in Table 2, 41 cases (69.5%) of

sarcomas belonged to soft tissue sarcomas. The most prevalent type of soft tissue sarcomas was rhabdomyosarcoma (27.1%).

Among the bone tissue sarcomas, affecting 18 patients (30.5%), osteosarcoma (15.3%) and chondrosarcoma (11.9%) were the most prevalent (Table 2).

Rhabdomyosarcoma was reported to be the most common sarcoma (13.5 %) in the both the age groups, as well as in men (10.2%) and women (16.9%). The lowest incidence was found to be Kaposi sarcoma, which was only observed in one man >16 years old (1.7 %) (Table 2).

The most areas commonly affected were soft tissues of the head and neck (49.2 %), jaw bones (35.6 %) as well as the head and skull bones (15.3%). With respect to tumor grading, most of the sarcomas were of the low grade (40.4%) followed by moderate (30.5%) and high (28.8%) grades, respectively. Metastases from other parts of the body to the head and neck occurred in 5 patients (8.5%), whereas, in 2 patients (3.4%) the sarcoma metastasized to other parts of the body from the head and neck. Out of the 59 examined patients, 16 cases (27.1%) were observed to have suffered from recurrence.

Table 1. Distribution of the head and neck sarcomas according to the demographic characteristics

Variable		Number	Percent
Age	≤16	17	28.8
	>16	42	71.2
Sex	Male	26	44.05
	Female	33	55.9
Habitat	Yazd	35	59.3
	Out of Yazd	24	40.7

Table 2: Distribution of the head and neck sarcomas according to demographic variables and frequency

Type of sarcoma	Name of sarcoma	N	%	Total	Sex				Age			
					Female		Male		16<		16≥	
					%	n	%	n	%	n	%	n
Soft tissue sarcomas	Rhabdomyosarcoma	16	27.1	(69.5%) 41	16.9	10	10.2	6	13.5	8	13.5	8
	Neurofibrosarcoma	5	8.5		6.8	4	1.7	1	6.8	4	1.7	1
	Fibrosarcoma	5	8.5		6.8	4	1.7	1	6.8	4	1.7	1
	Liposarcoma	3	5.1		1.7	1	3.4	2	5.1	3	0	0
	Malignant fibrous histiocytoma	3	5.1		1.7	1	3.4	2	3.4	2	1.7	1
	Synovial Sarcoma	3	5.1		1.7	1	3.4	2	0	0	5.1	3
	Angiosarcoma	3	5.1		3.4	2	1.7	1	5.1	3	0	0
	Leimyosarcoma	2	3.4		0	0	3.4	2	3.4	2	0	0
Kaposi's Sarcoma	1	1.7	0	0	1.7	1	1.7	1	0	0		
Hard tissue sarcomas	Osteosarcoma	9	15.3	(30.5%) 18	10.2	6	5.1	3	15.3	9	0	0
	Chondrosarcoma	7	11.9		6.8	4	5.1	3	8.5	5	3.4	2
	Ewing's sarcoma	2	3.4		0	0	3.4	2	1.7	1	1.7	1

## Discussion

Sarcomas are rare and involve 1% of human cancers, among which currently 4%–10% occur in the head and neck.<sup>(14)</sup> In the present study, 59 cases (10.06%) of 586 sarcomas patients were observed to suffer from the head and neck sarcomas.

In this study, 12 types of sarcoma histopathology were observed, which in turn demonstrates the wide histological variety of these tumors. Pacheco et al.<sup>(16)</sup> in a study on 36 patients with the head and neck sarcomas, reported 12 types of sarcomas. Lajer et al. observed 15 histopathological types of sarcoma in a study consisting of 36 patients, which are in line with the findings of the present study.

Most of the head and neck sarcomas belong to the soft tissue category, and only 20% are bone sarcomas or have a cartilage source<sup>(17)</sup>. In this study, 59 sarcomas were investigated among which 41 (69.5 %) soft tissue sarcomas, and 18 cases (30.5%) were hard tissue sarcomas. In a similar study in the Canadian population, Aljabab et al. observed 80% of sarcomas in the hard tissue and 20% of sarcomas in the soft tissue.<sup>(18)</sup>

In the current study, the most prevalent type of sarcoma, rhabdomyosarcoma, involved 27.1% of the total tumors. Multiple reports<sup>(19-23)</sup> have noted that approximately half of rhabdomyosarcomas commonly occur in the head and neck areas. This is in concurrence with the findings of the current study. However, the incidence of head and neck rhabdomyosarcoma can vary. Few studies mention the incidence to be in the range of 8% to 16%<sup>(24-26)</sup>, or higher (33.3%)<sup>(11)</sup>. This could be because of geographical and racial differences as well as the different sample size of the different studies. Various studies mention that no geographical preference can be taken into consideration for the occurrence of soft tissue sarcomas<sup>(36, 37)</sup>. The majority of the patients in our study<sup>(59,35)</sup> lived in Yazd, and the rest were from other cities traveled to Yazd for treatment. However, this might affect follow-up of patients in the study centers.

In this study, more than half of the involved patients with the head and neck sarcomas (56%) were females, which is in line with the findings by Bree et al.<sup>(27)</sup>, although in several other studies<sup>(23, 28-31)</sup> men have been reported to be affect-

ed more than the women. Some of these studies have indicated the predilection of males for the disease is nearly twice as much than females<sup>(29, 31)</sup>. Due to the rarity of head and neck sarcomas, and limited sample size of most studies conducted in Iran and the world, drawing conclusions and comparisons is not very effective, yet; perhaps one of the reasons for the higher prevalence of females in this study was that women were referred to health centers more than men.

As was expected, sarcomas of the head and neck can occur in any age. In fact, studies have demonstrated that soft tissue sarcomas affects 80%–90% of adults and 10%–20% of children.<sup>(18, 32)</sup> The mean age of the patients involved in this study was  $32.22 \pm 8.31$  years which supports the findings of the studies conducted by Pacheco et al.<sup>(16)</sup> and Dudhat et al.<sup>(30)</sup> who noted a similar average mean age of  $39.7 \pm 25.1$ . However, Epstein and Gorsky<sup>(33)</sup> reported the mean age of 40.4 years for this disease. Moreover, Mendenhall et al.<sup>(34)</sup> reviewed the literature published between 1972 and 2000, and indicated the mean age of 50–55 years for the head and neck sarcomas, though findings of some studies propose that generally sarcomas involving the head and neck affect younger people including children and teenagers compared to squamous cell carcinoma.<sup>(35)</sup>

In the present study, the anatomical distribution of most of the head and neck sarcomas included the jaw bone as well as the soft tissue of head and neck. Kraus et al.<sup>(32)</sup> also indicated these two areas as the most prevalent involved locations with the head and neck sarcomas. Tajudeen et al.<sup>(9)</sup> found the nasal cavity and sinuses as the most commonly involved locations in 22% of his patients. Penel et al.<sup>(28)</sup> reported a 39.3% involvement in the neck tissue that is consistent with the findings of the current study.

In the present study, 5 patients (8.5%) were found to have metastasis from other parts of the body to the head and neck. Breast is the most common site of tumor metastasis to the bone of the jaw, while the lungs involves the most frequent source for metastasis to the soft tissue of the mouth and teeth. In 30% of cases, metastasis in the mouth has been found to demonstrate the first sign of an undiscovered cancer in another part of the body.<sup>(38)</sup> Compared to other head and

neck neoplasms (e.g., squamous cell carcinoma), soft tissue sarcomas have a lower rate of regional metastasis.<sup>(39)</sup> In this study, only 2 cases (3.4%) had metastasis to other parts of the body, whereas in a study conducted by Tajudeen et al.<sup>(9)</sup> metastasis to lymph nodes was 6.5% and neural invasion was observed in 6.5% of the cases. Salcedo-Hernández et al.<sup>(5)</sup> reported 50% of soft tissue sarcomas of the head and neck involving metastasis.

Singh et al.<sup>(31)</sup> reported local recurrence in 42% of patients and 42% of metastatic disease development in the lungs. Probably in this study, one reason for the low figures of head and neck sarcoma metastasis to other parts of the body is the lack of follow-up of individual patients specifically from other provinces, who included a large portion of the study sample.

Recurrences of the sarcomas were observed in 27.1% of patients in this study.

Because the patients were not actively followed up in order to evaluate recurrence, in reality, the recurrence rate in this study might have been more than what is being reported here. It has been demonstrated that local recurrence in head and neck sarcomas is more than that of other organs<sup>(28, 40-43)</sup>, which is probably due to the fact that reaching negative margins of tumor is more complex during sarcoma surgery of the head and neck.<sup>(44)</sup>

Regardless of the location and size of the tumor, one of the main factors for prognosis with sarcomas is the tumor histologic grade.<sup>(45, 46)</sup>

## References

1. Cardona AF, Zuluaga J, Carranza H, et al. Stem Cells in Cancer: Should We Believe or Not?: Springer; 2014. P. 245-61.
2. Ryan C, Meyer J. Clinical presentation, histopathology, diagnostic evaluation, and staging of soft tissue sarcoma. Available from: <http://www.uptodate.com/contents/clinical-presentation-histopathology-diagnostic-evaluation-and-staging-of-soft-tissue-sarcoma> Sturgis
3. Sturgis EM, Potter BO. Sarcomas of the head and neck region. *Curr Opin Oncol* 2003;15(3):239-52.
4. Pellitteri PK, Ferlito A, Bradley PJ, et al. Management of sarcomas of the head and neck in adults. *Oral Oncol* 2003;39(1):2-12.
5. Salcedo-Hernández RA, Lino-Silva LS, Mosqueda-Taylor A, Luna-Ortiz K. Soft tissue sarcomas of the head and neck. Clinical and pathological evaluation of 108 cases in Mexico. *J Craniomaxillofac Surg* 2014; 42(8):1566-71.
6. De Bree R, van der Waal I, de Bree E, René Leemans C. Management of adult soft tissue sarcomas of the head and neck. *Oral Oncol* 2010;46(11):786-90.
7. Eeles R, Fisher C, A'Hern R, et al. Head and neck sarcomas: prognostic factors and implications for treatment. *Br J Cancer* 1993;68(1):201-7.
8. Moretti G, Guimarães R, Oliveira KMd, Sanjar F, Voegels RL. Rhabdomyosarcoma of the head and neck: 24 cases and literature review. *Braz J Otorhinolaryngol* 2010;76(4):533-7.
9. Tajudeen BA, Fuller J, Lai C, et al. Head and neck sarcomas: the UCLA experience. *Am J Otolaryngol* 2014; 35(4):476-

In this study, the majority (40.7%) of sarcomas were low-grade tumors, while 30.5% and 28.8% belonged to average and high grades respectively. However, Tajudeen et al.<sup>(9)</sup> reported 35% of their cases as high-grade sarcomas.

## Conclusion

The head and neck sarcomas are rare tumors that demonstrate a high variability in histology. In the current study, soft tissue sarcomas were generally much more prevalent than hard tissue sarcomas, among which rhabdomyosarcoma was the most common soft tissue sarcoma and osteosarcoma was the most common hard tissue sarcoma. Moreover, the age and gender prevalence, as well as the involved anatomic location in the studied population was similar to those of most other studies. However, our results, for the first time, provide an insight into the prevalence of head and neck sarcomas in Yazd, Iran.

The weak points of this study were the incomplete medical and pathological evidence of the patient and the patients were referred to other treatment centers and IHC (ImmunoHistoChemistry) results were unavailable in some cases.

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10. Levi F, Randimbison L, Maspoli-Conconi M, et al. Incidence of second sarcomas: a cancer registry-based study. *Cancer Causes Control* 2014; 25(4):473-7. doi: 10.1007/s10552-014-0349-7. Epub 2014.
11. Salari AA, Binesh F, Valizadeh S. Epidemiology of Soft Tissue Sarcomas in Shahid Sadoughi University of Medical Sciences of Yazd during 1994-2005. *Iranian J Surge* 2010; 18(4):39-44. Persian.
12. Neville BW, Damm DD, Allen CM, Bouquot JE. *Oraland Maxillofacial Pathology*. St Louis: Sanders; 2009.P.771-774.
13. Seddighi S, Rafat J. 1470 cases of sarcoma referring to Imam Khomeini hospital during an 11-year period. *Med Sci J* 2005; 15 (3):131-136 . Persian.
14. Huber GF, Matthews TW, Dort JC. Radiation-induced soft tissue sarcomas of the head and neck. *J Otolaryngol* 2007; 36(2):93-7.
15. Lahat G, Lazar A, Lev D. Sarcoma epidemiology and etiology: potential environmental and genetic factors. *Surg Clin North Am*. 2008 ;;88(3):451-81,
16. Pacheco IA, Alves AP, Mota MR, et al. Clinicopathological study of patients with head and neck sarcomas. *Braz J Otorhinolaryngol* 2011; 77(3):385-90.
17. Rapidis AD. Sarcomas of the head and neck in adult patients: current concepts and future perspectives. *Expert Rev Anticancer Ther* 2008; 8(8):1271-97.
18. Aljabab AS, Nason RW, Kazi R, Pathak KA. Head and neck soft tissue sarcoma. *Indian J Surg Oncol* 2011 2(4):286-90.
19. Hagiwara AI, Inoue Y, Nakayama T, et al. The 'botryoid sign': a characteristic feature of rhabdomyosarcomas in the head and neck. *Neuroradiology* 2001; 43(4):331-5.
20. Lee JH, Lee MS, Lee BH, et al. Rhabdomyosarcoma of the head and neck in adults: MR and CT findings. *AJNR Am J Neuroradiol* 1996; 17(10):1923-8.
21. Lioyd C, McHugh K. The role of radiology in head and neck tumours in children. *Cancer Imaging* 2010 Mar 3;10:49-61. doi: 10.1102/1470-7330.2010.0003.
22. Franco T, La Boria A, Domanico R, et al. Rare adult masseteric rhabdomyosarcoma and a review of the literature. *Case Rep Oncol* 2013; 6(3):472-9.
23. Penel N, Van Haverbeke C, Lartigau E, et al. Head and neck soft tissue sarcomas of adult: prognostic value of surgery in multimodal therapeutic approach. *Oral Oncol* 2004; 40(9):890-7.
24. Bentz BG, Singh B, Woodruff J, et al. Head and neck soft tissue sarcoma: a multivariate analysis of outcome. *Ann Surg Oncol* 2005; 11(6):619-28.
25. Chen AS, Morris CG, Andur RJ, et al. Adult head and neck soft tissue sarcomas. *Am J Clin Oncol* 2005; 28(3):259-63.
26. Yamaguchi S, Nagasawa H, Suzuki T, et al. Sarcomas of the oral and maxillofacial region: a review of 32 cases in 25 years. *Clin Oral Invest* 2004; 8(2):52-5.
27. Bree R, Valk P, Kuik DJ, et al. Prognosis factors in adult soft tissue sarcomas of the head and neck: a single centre experience. *Oral Oncol* 2006; 42(7):703-9.
28. Penel N, Mallet Y, Robin YM, et al. Prognostic factors for adult sarcomas of head and neck. *Int J Oral Maxillofac Surg* 2008; 37(5):428-32.
29. Le QT, Fu KK, Kröll S, et al. Prognostic factors in adult soft tissue sarcoma of the head and neck. *Int J Radiat Oncol Biol Phys* 1997; 37:975-84.
30. Dudhat SB, Mistry RC, Varughese T, et al. Prognostic factors in head and neck soft tissue sarcomas. *Cancer* 2000; 89(4):868-72.
31. Singh RP, Grimer RJ, Bhujel N, et al. Adult head and neck soft tissue sarcomas: treatment and outcome. *Sarcoma* 2008;2008:654987. doi: 10.1155/2008/654987.
32. Kraus DH, Dubner S, Harrison LB, et al. Prognostic factors for recurrence and survival in head and neck soft tissue sarcomas. *Cancer* 1994; 74(2):697-702.
33. Gorsky M, Epstein JB. Craniofacial osseous and chondromatous sarcomas in British Columbia - a review of 34 cases. *Oral Oncol* 2000; 36(1):27-31.
34. Mendenhall WM, Mendenhall CM, Werning JW, et al. Adult head and neck soft tissue sarcomas. *Head Neck* 2005; 27(10):916-22.
35. Patel SG, Shaha AR, Shah JP. Soft tissue sarcomas of the head and neck: an update. *Am J Otolaryngol* 2001;22(1):2-18.
36. Vincent T. Devita Jr. Samuel Hellman, et al, *Sarcoma of the Soft tissue and bone, cancer principles and practice of oncology. USA: Lipincott Williams & Wilkins; 2001.p. 1841-44.*
37. Weiss SW, Goldblum JR. *Enzinger and Weiss's Soft Tissue Tumor*. 4th ed. St. Louis: Mosby; 2001.p. 1-17.

38. Hirshberg A, Buchner A. Metastatic tumours to the oral region. An overview. *Eur J Cancer B Oral Oncol* 1995;31B(6):355-60.
39. Srivastava A, Ghosh A, Saha S, et al. Sarcomas of head and neck - A 10 yrs experience. *Indian J Otolaryngol Head Neck Surg* 2007;59(4):322-6.
40. Gullane P, Kraus D, Weber R. Soft tissue sarcoma. *Head Neck* 2002; 24(3):296-300.
41. Borden EC, Baker LH, Bell RS, et al. Soft tissue sarcomas of adults: state of the translation science. *Clin Cancer Res* 2003; 9(6):1941-56
42. Wunder JS, O Nielsen T, Maki RG, et al. Opportunities for improving the therapeutic ratio for patients with sarcoma. *Lancet Oncol* 2007;8(6):513-24.
43. Shellenberger TD, Sturgis EM. Sarcomas of the head and neck region. *Curr Oncol Rep* 2009; 11(2):135-42.
44. Piñeiro Aguin Z, León Vintó X, García Lorenzo J, et al. [Head and neck sarcomas. Our experience]. *Acta Otorinolaringol Esp* 2011 Nov-Dec;62(6):436-42. doi: 10.1016/j.otorri.2011.05.005. Epub 2011 Aug 5. Spanish.
45. Tran LM, Mark R, Meier R, et al. Sarcomas of the head and neck. Prognostic factors and treatment strategies. *Cancer* 1992;70(1):169-77.
46. Stojadinovic A, Leung DH, Hoos A, et al. Analysis of the prognostic significance of microscopic margins in 2,084 localized primary adult soft tissue sarcomas. *Ann Surg* 2002; 235(3):424-34.