Malignant minor salivary gland tumors: A retrospective study of 28 cases: Clinical experience of a single institution

Abeer Hussien Anter, Rash Hamdy Hamed

Department of Clinical Oncology and Nuclear Medicine, Faculty of Medicine, Mansoura University, Egypt

Received October 29, 2014; Revised December 21, 2014; Accepted December 29, 2014; Published Online January 01, 2015

Original Article

Abstract

Purpose: This retrospective study was done to report our experience with the management of patients with primary malignant tumors of minor salivary glands that were treated with various combinations of surgery, radiation, and chemotherapy and to review treatment outcome. Methods: The records of all patients with malignant minor salivary gland tumors presenting for treatment at our department between 2000 and 2010 were retrospectively reviewed. Variables were collected and outcome measures were defined in terms of overall survival, disease-free survival. Descriptive statistics were compiled and statistically evaluated. Survival was described using the Kaplan-Meier method. Results: Twenty-eight patients (10 males and 18 females) ranging in age from 18 to 80 years (median, 64 years) met the criteria for inclusion in the study. The majority of tumors were located in the oral cavity (n = 20), followed by nasal cavity and paranasal sinuses (n = 8). Adenoid cystic carcinoma was the most common neoplasm (n = 18). All patients were treated with surgery as the primary modality. Neck dissection was performed in 18% of patients, and all patients (n = 28) were treated with adjuvant external beam radiation therapy to a dose of 50 to 60 Gy. The disease-free survival rate and overall survival at 5 years were 80% and 85%, respectively. Conclusion: Postoperative radiation therapy is effective in preventing local recurrence in most patients with minor salivary gland tumors after gross total excision.

Keywords: Minor Salivary Gland tumours; Postoperative Radiation; Adenoid Cystic Carcinomas

Introduction

Minor salivary glands (MSG) are numerous (700 to 900) exocrine glands, which are distributed in the submucosa of the oral cavity, nasopharynx, tonsils, hypopharynx, and larynx. Heterotopic sites include lymph nodes, the thyroid gland, facial bones, and the hypophysis.1-3 Tumors of the intra-oral minor salivary glands constitute 2-3% of all malignant neoplasms of the upper aerodigestive tract and less than 25% of all salivary gland tumors.4 Tumors arising at major salivary glands due to their well-defined anatomic borders and their higher incidence cannot be directly compared to minor salivary gland tumors. Most tumors arising at minor salivary glands are malignant.4,5

Surgery is the accepted primary modality of treatment for most of these tumors.6 The role of radiation in the post-surgical setting is less clear because there are few large series with long follow-up. Thus, the treatment approach to minor salivary gland tumors is based on principles of treatment of major salivary gland tumors, where a larger experience with postoperative radiation has established its value in patients with locally advanced disease; recurrent disease in an unirradiated site; high grade histological type; perineural spread; close, uncertain, or positive surgical margins; or neck nodal disease.7

These features are relatively more common in minor salivary gland tumors because they have a higher incidence of malignancy than do parotid tumors, adenoid cystic carcinomas are more common, and it is frequently harder to get clear margins, particularly in tumors arising from the paranasal sinuses. Thus, it is reasonable to expect that postoperative radiation would have a correspondingly greater role in their management.8 Malignant tumors of minor salivary glands are very rare. Therefore, design of management protocols is challenging. So our study that reports our experience in management of malignant MSG tumors beside other retrospective studies may help to design a protocol of management for those patients.

This study aims to retrospectively review our experience with the management of patients with primary malignant tumors of minor salivary glands that were treated with various combinations of surgery, radiation, and chemotherapy and to review treatment outcome, over ten years within a
single radiotherapy (RT) unit in a large tertiary referral centre in our locality.

Methods and Materials

This study is a retrospective study of patients with primary malignant tumors of MSG that has been treated by post-operative radiotherapy at Clinical Oncology & Nuclear Medicine Department, Mansoura University Hospital during the 10-year period (2000 to 2010).

A total 38 patients with primary malignant tumors of minor salivary gland were identified. The medical records of all patients with primary malignant tumors of MSG, were retrieved and reviewed. Demographic data, clinical characteristics, diagnostic investigations, histological typing, treatment modalities including type of surgery, radiotherapy and chemotherapy, follow-up, recurrence, and survival were noted.

All patients provided a history and underwent physical examination. Radiologic imaging included chest x-ray, and computed tomography or magnetic resonance imaging scans. Computed tomography was preferred for cases with bone involvement, whereas MRI was reserved solely for soft tissue invasion.

Radiotherapy

All patients received their radiation therapy with or without cisplatin after a surgical procedure for removing a primary minor salivary gland cancer or after surgery for locally recurrent disease.

![FIG. 1: A 58 year-old man with T4N0M0 Adenoid cystic carcinoma of Rt maxillary sinus, for post-operative radiotherapy to tumor bed only (nodal regions not electively treated). (a) Axial. (b) Sagittal images displaying isodose distributions. Shaded blue is PTV.](image)

The interval between the date of last surgery and start of radiation therapy ranged from 2 to 8 weeks. Radiation techniques were dependent on the primary site. The techniques were grouped as follows: single appositional fields, parallel opposed fields, three field techniques, and wedged paired fields (Figure 1), with megavoltage equipment, using cobalt-60 gamma rays, 6 MeV linear accelerator or 6-20 MeV electrons. Doses to the primary sites ranged from 5000 to 6000 cGy. The median number of fractions per course of treatment was 30. Concurrent chemotherapy consisted of cisplatin, which was typically administered weekly with a dose of 40 mg/m².

Statistical analysis

The Kaplan-Meier method was used to estimate survival. Overall survival was defined as the interval between diagnosis and death or last follow-up. Disease-free survival (DFS) was defined as the interval between from the date of surgery to the earliest time of recurrence in local or distant sites, death from any cause or last follow-up.

A p-value ≤0.05 was considered significant. All analyses were conducted with SPSS version 15.0 (SPSS for Windows, Rel. 13.0 2004. Chicago: SPSS Inc.).

Results

In the 10-year period of 2000 to 2010, 38 patients with minor salivary gland malignancy were treated in our institution. Patients with unresected or gross residual disease after attempted resection (6 patients) or with distant metastatic disease at the time of their radiation (4 patients) were excluded from the study. This left 28 patients between the ages of 18 and 75 years (median, 64 years) for inclusion in this analysis. Eighteen patients were female, and 10 were male.

The primary tumor site was in the oral cavity in 20 patients and in nasal cavity or paranasal sinuses in 8 patients (Table 1). The chief complaint on presentation was a lump (60%), followed by an ulcerated lump (22%). Overall 82% of the patients presented a lump with or without ulceration of the surface.

<table>
<thead>
<tr>
<th>Site</th>
<th>No. of Patients</th>
</tr>
</thead>
<tbody>
<tr>
<td>Maxillary sinus</td>
<td>2</td>
</tr>
<tr>
<td>Nasal cavity</td>
<td>4</td>
</tr>
<tr>
<td>Ethmoid sinus</td>
<td>2</td>
</tr>
<tr>
<td>Palate</td>
<td>4</td>
</tr>
<tr>
<td>Buccal mucosa</td>
<td>4</td>
</tr>
<tr>
<td>Lip</td>
<td>2</td>
</tr>
<tr>
<td>Tongue</td>
<td>2</td>
</tr>
<tr>
<td>Floor of mouth</td>
<td>4</td>
</tr>
<tr>
<td>Gingiva/alveolar ridge</td>
<td>2</td>
</tr>
<tr>
<td>Retromolar trigone</td>
<td>2</td>
</tr>
</tbody>
</table>

Most patients (18) had a histologic diagnosis of adenoid cystic carcinoma. 6 mucoepidermoid carcinoma and 4 patients had adenocarcinoma (Table 2). Surgical procedures varied dependent on the size and site of the primary lesion. 2 patients had excisional biopsies, 14 patients had wide local
excisions, and the remaining 12 patients had more extensive resections, including 3 partial or total maxillectomies. After these procedures, 11 (39%) patients had pathologic evidence of microscopic positive margins, and an additional 6 (21%) had close (less than 5 mm) or uncertain margins. A neck dissection was performed on 5 (18%) patients, with pathologically positive lymph nodes being found in 3 patients. Thirteen patients (46.6%) had pathologic evidence of perineural invasion. Bone invasion by the primary was present in 6 (21%) patients.

<table>
<thead>
<tr>
<th>Type of tumour</th>
<th>Number</th>
<th>%</th>
</tr>
</thead>
<tbody>
<tr>
<td>Adenoid cystic carcinoma</td>
<td>18</td>
<td>64.3</td>
</tr>
<tr>
<td>Mucoepidermoid carcinoma</td>
<td>6</td>
<td>64.3</td>
</tr>
<tr>
<td>Adenocarcinoma</td>
<td>4</td>
<td>14.3</td>
</tr>
</tbody>
</table>

All patients received their radiation therapy with or without cisplatin after a surgical procedure for removing a primary minor salivary gland cancer. In 25 patients, radiation therapy followed the initial surgery; the remaining 3 (10.7%) were irradiated after surgery for locally recurrent disease.

Local recurrence was recorded in two patients; one after 24 months, other after 5 years. Lung metastasis was only observed in three cases; one of them developed late lung metastasis after 5 years, as confirmed by biopsy from lung lesion. Also, three patients developed wide spread metastasis. Overall, 20 patients were free of local or disseminated disease at the last follow-up. Seven patients died during the course of our study. Two patients died due to unrelated causes (pneumonia, pulmonary embolism), three of them due to disseminated metastasis and the last two patients due to respiratory failure. The 5, 10 years overall survival rates for the whole series were 85%, 66% (Figure 2). On other hand, the 5, 10 year disease-free survival was 80%, 64% respectively (Figure 3).

![FIG. 2: Survival curve for 28 patients with malignant minor salivary gland tumors treated with postoperative radiation therapy.](image)

**Discussion**

This study is a retrospective study of patients with primary malignant tumors of MSG that has been treated at Clinical Oncology & Nuclear Medicine Department, Mansoura University Hospital. The present study revealed that primary malignant tumors of MSG occurred in the old population (median, 64 years). This was also reported by others.5, 9, 10

Patients were more often female than male in our data, with a ratio of 1.8:1. Review of the literature shows a female predilection among the patients with primary malignant tumors of MSG.5, 9, 11

According to other reports,5, 13, 14, 15, the palate was the most commonly affected site, and this was also seen in our study. Moore et al.16 suggests that primary malignant tumors of MSG usually occurred at the junction of the hard and soft palates due to concentration of salivary glands in this region.

In this study, adenoid cystic carcinoma was the most common pathological type of primary malignant tumors of MSG (64.3%) followed by mucoepidermoid carcinoma (21.4%). Studies concerning west European population17–19 reported adenoid cystic carcinoma as the most common malignancy but according to other reports,11, 20, 21; mucoepidermoid carcinoma was the most common malignant MSGTs.

In the present series, a neck dissection was performed on 5 (18%) patients, with pathologically positive lymph nodes being found in 3 (10.7%) patients which abides to the general proportion of 4 to 10% recorded in other studies.10, 22, 23 Many factors usually dictate selective neck dissection, such as the site and size (T) of the tumor, the histological grade, and the age of the patient. Parsons et al. have classified MSG...
tumor sites according to the incidence of node positivity, according to this classification; low incidence (less than 10%) is observed in MSG tumors of the hard palate, nasal cavity, paranasal sinuses, lip, and buccal mucosa. The intermediate risk of positive neck at presentation should be expected in MSG tumors of the floor of the mouth and tongue. The higher risk for node metastases lies with MSG tumors of the pharynx and larynx. However, other authors advocate significantly lower lymph node infiltration than hematogenous metastases for MSG tumors.

All patients received their radiation therapy after a surgical procedure for removing a primary minor salivary gland cancer. In 25 patients, radiation therapy followed the initial surgery; the remaining 3 (10.7%) were irradiated after surgery for locally recurrent disease.

Adjuvant radiotherapy was offered to patients who presented with advanced stage, histopathologically proven positive neck metastases, close or positive margins and perineural invasion. Combination of radiotherapy and cisplatin was offered to patients with high grade histological subtype and advanced stage of the disease at presentation. Other authors reserve such a therapy for patients with distant metastases or as palliative treatment.

In this study, The 5 year survival rates for the whole series were 89% and this was also seen in another study. This trial has some limitations. One is the relatively small number of patients another factor is that it is a single institution study. Multicenter randomized Trials are necessary to clarify the role of radiotherapy in patients with MSG tumors.

Conclusion

Patients with minor salivary gland malignancies considered at high risk of local failure benefit from postoperative radiation. Local control rates are comparable to those reported for major salivary gland tumors, also late recurrences and metastases that were also noted in our study confirm the necessity of long-term follow-up of these patients, more than 5 years.

Conflict of interest

The authors declare that they have no conflicts of interest. The authors alone are responsible for the content and writing of the paper.

References


