Case Report

Radiotherapy in Kimura’s Disease: A Report of Eight Cases


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Kimura’s disease is a rare condition of chronic inflammatory disorder affecting the skin and subcutaneous tissue. It is predominantly in the head and neck region. The lesion is benign but may be persistent/recurrent and difficult to eradicate. Several forms of treatment have been used, including surgical excision, intralesional and oral corticosteroid, cryotherapy and radiotherapy. The authors report eight cases with histopathology consistent with Kimura’s disease who received radiation therapy as a primary treatment or secondary treatment for recurrence after surgical excision in the Division of Therapeutic Radiology and Oncology, Chiang Mai University. The prescribed radiation doses varied from 30-40 Gy. With the mean follow-up time of 21 months, all eight patients were still free from disease at the time of analysis.

Keywords: Kimura’s disease, Head and neck, Radiotherapy

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Kimura’s disease or eosinophilic lymphfolliculoid granuloma was first described by Kimura et al in 1948(1). The most common clinical manifestation of this disease is a unilateral soft tissue mass in the head and neck area, particularly around the ear. It is also associated with regional lymphadenopathy and subcutaneous mass(2). There are some reports in other sites including oral mucosa, nasal cavity, bone, muscle, salivary glands, orbit, lacrimal gland, and tympanic membrane(4). Kimura’s disease affects young men. The incidence is high in the 2nd and 3rd decades(1,6). It appears more common in Oriental men than Caucasians(8). Kimura’s disease shares some histopathological features with angiolymphoid hyperplasia with eosinophilia (ALHE). These two lesions have repeatedly been confused and designations used interchangeably(12). The plump endothelial cells (cuboidal to dome-shaped) characteristic of ALHE remains the keys to separation of these entities(10,14). However, many clinical and histopathological features remain distinctly different(6).

Material and Method

From January 2002 to December 2004, eight patients with Kimura’s disease were treated in the Division of Therapeutic Radiology and Oncology, Chiang Mai University, Thailand. Histological examination was performed and reviewed by two pathologists. All showed the same histopathology of Kimura’s disease. The authors collected all data from the medical records including; age, sex, site of disease, primary treatment, time to recurrence (in case of recurrence disease), radiotherapy technique, total dose of radiotherapy, and the result of treatment.

Results

Patient and tumor characteristics

There were six males and two females. The
mean age was 39 years, range from 24 to 54 years. Five patients had local recurrence after surgical excision. All of them recurred at the same site with time of recurrence from 1 to 15 years. The other three patients had the first time diagnosed. Incisional biopsies were performed in all eight patients. The histopathology of all specimens revealed well circumscribed lesions with separated lymphoid follicles and prominent germinal centers. Proliferation of small vessels with flat to low cuboidal endothelial cells and tissue eosinophilia, some with eosinophilic micro-abscesses, were noted in interfollicular area. Prominent fibrosis around small vessels and lymphoid follicles was observed. The histopathology of the presented patients is shown in Fig. 1. The presenting symptoms in eight patients were painless mass in head and neck area. The most common site of involvement were parotid region in six patients, followed by post auricular region in one patient and submandibular region in the remaining patient.

All the baseline characteristics are shown in Table 1.

**Radiation therapy**

Radiotherapy was given to all eight patients, nine lesions. Seven patients were treated with appositional fields; mixed beam of electron and photon, one patient who had submandibular lesion was treated with parallel opposing two lateral techniques. The prescribed radiation doses varied from 3000 to 4000 cGy, with 200 cGy per fraction, five fractions per week.

**Outcome of treatment**

With the mean follow-up time of 21.75 months (range from 21 to 43 months), local control of the lesion was obtained in all eight patients and nine lesions. At the end of radiotherapy, the authors observed a marked reduction in tumor size in all lesions and achieved excellent local control at the time of analysis. Only one patient experienced a new lesion in the contralateral parotid region after 18 months of radiotherapy (patient No. 5). She also received radiotherapy to the left parotid area 3000 cGy, the mass disappeared and is still in remission.

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**Table 1. Baseline characteristics**

<table>
<thead>
<tr>
<th>No.</th>
<th>Sex</th>
<th>Age</th>
<th>Site</th>
<th>Primary Rx</th>
<th>Time to recurrence (year)</th>
<th>Site of recurrence</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>M</td>
<td>50</td>
<td>Parotid Rt.</td>
<td>Superficial parotidectomy</td>
<td>2</td>
<td>Same</td>
</tr>
<tr>
<td>2</td>
<td>M</td>
<td>24</td>
<td>Postauricular Rt, Lt</td>
<td>-</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>3</td>
<td>M</td>
<td>33</td>
<td>Parotid Lt.</td>
<td>-</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>4</td>
<td>M</td>
<td>30</td>
<td>Parotid Lt.</td>
<td>-</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>5</td>
<td>F</td>
<td>40</td>
<td>Parotid Rt.</td>
<td>Surgical excision</td>
<td>2</td>
<td>Same</td>
</tr>
<tr>
<td>6</td>
<td>M</td>
<td>45</td>
<td>Parotid Rt.</td>
<td>Superficial parotidectomy</td>
<td>1</td>
<td>Same</td>
</tr>
<tr>
<td>7</td>
<td>F</td>
<td>54</td>
<td>Submandibular Rt.</td>
<td>Surgical excision</td>
<td>15</td>
<td>Same</td>
</tr>
<tr>
<td>8</td>
<td>M</td>
<td>36</td>
<td>Parotid Rt.</td>
<td>Superficial parotidectomy</td>
<td>4</td>
<td>Same</td>
</tr>
</tbody>
</table>

M = Male, F = Female, Rt. = Right, Lt. = Left
Table 2. Radiotherapy and outcome

<table>
<thead>
<tr>
<th>No.</th>
<th>RT technique</th>
<th>RT dose</th>
<th>Result</th>
<th>Follow up time (months)</th>
<th>New lesion</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Mixed beam</td>
<td>3000</td>
<td>Control</td>
<td>22</td>
<td>-</td>
</tr>
<tr>
<td>2</td>
<td>Mixed beam</td>
<td>3600</td>
<td>Control</td>
<td>22</td>
<td>-</td>
</tr>
<tr>
<td>3</td>
<td>Mixed beam</td>
<td>3000</td>
<td>Control</td>
<td>25</td>
<td>-</td>
</tr>
<tr>
<td>4</td>
<td>Mixed beam</td>
<td>4000</td>
<td>Control</td>
<td>38</td>
<td>-</td>
</tr>
<tr>
<td>5</td>
<td>Mixed beam</td>
<td>3000</td>
<td>Control</td>
<td>36</td>
<td>Contralateral parotid mass</td>
</tr>
<tr>
<td>6</td>
<td>Mixed beam</td>
<td>4000</td>
<td>Control</td>
<td>31</td>
<td>-</td>
</tr>
<tr>
<td>7</td>
<td>Photon: two lateral</td>
<td>4000</td>
<td>Control</td>
<td>43</td>
<td>-</td>
</tr>
<tr>
<td>8</td>
<td>Mixed beam</td>
<td>3000</td>
<td>Control</td>
<td>21</td>
<td>-</td>
</tr>
</tbody>
</table>

AP-PA = anteroposterior-posteroanterior

free from recurrence. Table 2 shows the result of treatment. Fig. 2 and 3 show the pictures of patients No. 5 and No. 3 before and after treatment.

Discussion

Kimura’s disease is a chronic inflammatory soft tissue disorder of unknown etiology. This condition is benign but it has been misdiagnosed as a malignant lesion or serious disease, leading to unnecessary radical surgery(10). The most common presentation is a unilateral soft tissue mass in the head and neck area. The authors' result also found the same natural history with only one patient who had bilateral lesions.

Kimura’s disease occurs predominantly in young adult males. Of the eight patients in the present study, only two patients were females, and six patients (75%) were males. The one thing that the authors found differently from the literatures was the age of the patient. The mean age of the presented patients was 39 years, and two male patients were 45 and 50 years old respectively.

Surgery, radiotherapy, and steroid therapy have been tried, but none has proved to be the optimum modality. Steroid therapy has produced marked response in reduction of tumor size, but after weaning, the tumors often increase in size(11). Surgery with complete excision is very difficult because the infiltrative nature of the tumor and swelling of regional lymph nodes(11). Five patients (62%) in the present study were treated by local excision/radical surgery before. All of them eventually had local recurrence after surgical excision from 1-15 years (mean 4.8 years). However, the authors could control this recurrent disease with radiation therapy.

Fig. 2 Photographs of case No. 5
(a) before radiotherapy
(b) after radiotherapy
Hareyama et al recommended the limited field of radiotherapy (cover only the lesion and adjacent lymph node) with a dosage of 26-30 Gy\(^{11}\). The authors gave 30-40 Gy to the lesions and swelling lymph node and had a satisfactory result in every lesion.

The present results suggest that radiotherapy is an effective alternative approach for the primary treatment of Kimura’s disease and should be applied for the recurrent disease after surgical excision. The role of adjuvant radiotherapy after surgery is still controversial and needs to be studied.

Fig. 3 (a) CT scan of case No. 5: right parotid, before radiotherapy (b) CT scan of case No. 5: right parotid, after radiotherapy; left parotid, recurrence (c) CT scan of case No. 3: left parotid before radiotherapy (d) CT scan of case No. 3: left parotid after radiotherapy

References


รังสีรักษาโรคคิมูระ: รายงานผู้ป่วย 8 ราย

อิ่มใจ ชิตาพานรักษ์, ชรินทร์ ยาอินทร์, รุ่งอรุณ กิตติเชษฐ์, พิมพ์ขวัญ กำเนิดศุภผล, วิชญ์ หล่อวิทยา, วิลชัย สุขถมยา, พิชิต สิทธิไตรย์, เอียร์ชัย ภัทรภูมิ

ภูมิหลัง: โรคคิมูระเป็นการติ่งสัณฐานของผิวหนังและเนื้อเยื่อใต้ผิวหนัง พบบ่อยที่บริเวณศีรษะและลำคอ ถือเป็นโรคก้อนทูมที่ไม่ใช่มะเร็งร้าย แต่มักกลับเป็นซ้ำ และยากในการรักษาให้หายได้ดี การรักษาหลายวิธี ที่นิยมใช้ได้แก่การผ่าตัด, การฉีดคอร์ติโคสเตียรอยด์ เข้าไปในก้อน, การกินคอร์ติโคステียรอยด์, การใช้เครื่องเย็นและการฉายรังสี

วัตถุประสงค์: เพื่อนำเสนอผลการใช้รังสีรักษาในผู้ป่วยโรคคิมูระ

วัสดุและวิธีการ: ทำการศึกษาในผู้ป่วยโรคคิมูระ ในหน่วยรังสีรักษาและมะเร็งวิทยา ภาควิชารังสีวิทยา คณะแพทยศาสตร์มหาวิทยาลัยเชียงใหม่ ที่รับการรักษาตั้งแต่เดือนมกราคม พ.ศ. 2545 ถึงเดือน ธันวาคม พ.ศ. 2547 ทั้งการรักษาครั้งแรกและเป็นการรักษาหลังจากเกิดการกลับเป็นซ้ำ จำนวน 8 ราย

ผลการศึกษา: ผู้ป่วยที่ได้รับการรักษา 8 ราย มีจำนวนร้อยละ 9 แห่ง ได้รับการรักษาด้วยการฉายรังสี 30-40 Gy หลังจากอาการดีดวยหลังการรักษาเกิดการกลับเป็นซ้ำ 21 เดือน พบว่าผู้ป่วยทั้ง 8 ราย ไม่พบมีโรคเกิดขึ้น และไม่พบภาวะแทรกซ้อนหลังการรักษาอย่างรุนแรง

สรุป: การใช้รังสีรักษาไม่ว่าจะเป็นการรักษาครั้งแรก หรือ ใช้เป็นการรักษาหลังจากที่ผู้ป่วยกลับเป็นซ้ำ ในการรักษาโรคคิมูระมีประสิทธิภาพที่ดีในการควบคุมโรคเฉพาะที่