Can Multi-Detector Computed Tomography (MDCT) Help in Differentiation of Neoplastic Parotid Lesions?

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Abstract

Background: Parotid lesions are uncommon issue in clinical practice with wide differential diagnosis based upon multiple pathological varieties and limited clinical values. Computed tomography (CT) try to solve this issue as possible.

Objectives: To evaluate the diagnostic reliability of multi-detector computed tomography (MDCT) in characterization of benign and malignant parotid lesions using a systematic approach in comparison to final pathological data.

Patients & Methods: A retrospective study includes 136 patients diagnosed pathologically as parotid tumors underwent multi-detector computerized tomography (MDCT) examination using 64-row multi-detector computerized tomography (CT) scanner. All the images were evaluated for certain multi-detector computerized tomography (MDCT) criteria in a systematic manner.

Results: Our study revealed primary malignant tumors in 88 patients, secondary metastatic lesions in 3 patients and benign tumors in 45 patients. Five multi-detector computerized tomography (MDCT) criteria were statistically significant in differentiating between benign and malignant parotid lesions, they were; lesion margin (p=0.001), location (p=0.05), enhancement pattern (p=0.001), central necrosis (p=0.001) and para-pharyngeal space extension (p=0.001).

Conclusion: Multi-detector computerized tomography (MDCT) can be valuable in differentiating between benign and malignant parotid lesions and leading the management strategy aiming for best surgical outcome.

Keywords: Parotid; MDCT; Benign; Malignant; Tumors

Key Points

a) Parotid lesions show unsatisfactory discriminating imaging criteria to guide the surgical approach.

b) Rapid advances of MDCT techniques allow optimum parotid lesions characterization.

c) Imaging is forming a crucial triad with clinical data and surgical approach.

Introduction

Parotid lesions are uncommon in clinical practice with unsatisfactory discriminating imaging criteria to guide the surgical approach. They account for 1% of head and neck tumors and about 80-85% of parotid neoplasm are benign, 15-20% of parotid lesions are malignant [1,2]. The clinical challenge of parotid lesions is that, they grow slowly over years before the patients seek medical consultation, only few cases may have represented by facial nerve palsy. Also the fine needle aspiration results may be inaccurate and non-conclusive in many cases [3]. The availability and low cost of computed tomography (CT) in conjunction with rapid advance of CT technology including dual source, prospective triggering and high pitch helical acquisition should be applied for better characterization of various parotid lesions. All benign and malignant parotid lesion are indicated for surgery and our rule is to get well preoperative differentiation between benign and malignant lesions and leading the surgical approach to be as conservative as possible especially for facial nerve and lymph node dissection as well as improve the post-operative management strategy [4-6].

Patients and Methods

Patients

This retrospective study was approved by the ethics committee of our institution during the period between May 2014 and February 2016. It included 136 patients who were pathologically diagnosed by biopsy or fine needle aspiration biopsy (FNAB) as parotid neoplasm.
MDCT technique

Multi-detector CT examination of the parotid region was performed for all the 136 patients included in this study using 64-row multi-detector CT scanner (Aquilion 64; Toshiba Medical Systems Corporation, Otawara, Japan), which is a 64x0.5mm collimation scanner with a gantry rotation speed of 400ms/rotation. The examination was performed with the patient supine in quite respiration. A pad is placed beneath the patient scapula to produce mild hyperextension of the neck and provide consistent images perpendicular to long axis of the neck, that minimizing dental artifacts. The scanning range will be individually adapted and included from the external auditory canal to the level of the hyoid bone. Contrast material is injected through an 18-20 gauge catheter in to the ante-cubital vein with flow rate 4ml/sec. The contrast used according to body weight 1.5ml/kg. A non ionic contrast material (Ultravist 370) was used. A timing bolus tracking technique was employed. The acquisition parameters were 120kvp, 350mAs, a helical pitch of 0.983: 1, 7.5 second scan time, 7.5 second total exposure time, 1.25mm helical slice thickness, and 1.25mm reconstruction interval with a large FOV.

Post-processing and image reconstructions

The axial source images with a 1.25mm slice were transferred to an external work station (Vitrea) to perform multi-planar reformatted images (MPR) images in coronal and sagittal planes with a section thickness of 3-5mm.

Image analysis

The MDCT images were interpreted by two expert radiologists in head and neck imaging 5 and 10 years of experience who were blinded to results of each other. All the MDCT images were interpreted for the MDCT criteria of the parotid lesions that included the following items; size, margin, location, enhancement pattern, multiplicity, central necrosis, parapharyngeal fat extension and calcifications as well as associated lymph node enlargement.

Statistical analysis

Data entry was done by SPSS version 11. Frequency distribution, descriptive statistics and correlation analysis were done using Chi2 and Fisher exact tests for qualitative data. The probability (p value) of less than 0.05 is used as a cut off point for all significant tests.

Pathological examination

Biopsy was performed for 79 lesions, fine needle aspiration biopsy (FNAB) for 57 lesions and 96 patients had surgical excision. Histo-pathologic examination was performed for the entire biopsied specimen.

Results

The patient sages of the studied group ranged from 5 to 77 years old with a mean age of 44.4± 22.2. They were 46 males (57.5%) and 34 females (42.5%). Our biased study shows higher rate of malignant parotid lesions with lower incidence of benign tumors. Table 1 listed the pathologic types of the parotid lesions according to final pathological examination. The benign lesions were detected in 45 (33.1%) out of 136 patients. The most common benign lesion was pleomorphic adenoma which represents 55.6% of all benign lesions. The malignant lesions were further classified into primary and metastatic lesions. The primary lesions were detected in 88(64.7%) out of 136 patients, the most common primary malignant lesion was mucoepidermoid carcinoma which represents 28.4%. The metastatic lesions were detected in 3(2.2%) out of 136 patients, 2 of them were metastasis from malignant scalp lesion and 1 of them was metastasis from nasopharyngeal squamous cell carcinoma.

### Table 1: Pathological causes of enlarged parotid gland (n=136).

<table>
<thead>
<tr>
<th>Pathological cause</th>
<th>Number</th>
<th>Percent</th>
</tr>
</thead>
<tbody>
<tr>
<td>Benign neoplastic causes</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Pleomorphic adenoma</td>
<td>25</td>
<td>55.50%</td>
</tr>
<tr>
<td>Warthin’s tumor</td>
<td>10</td>
<td>22.20%</td>
</tr>
<tr>
<td>Lymphoepithelial lesion</td>
<td>8</td>
<td>17.70%</td>
</tr>
<tr>
<td>Myoepithelioma</td>
<td>2</td>
<td>4.40%</td>
</tr>
<tr>
<td>Total</td>
<td>45</td>
<td>100%</td>
</tr>
<tr>
<td>Primary malignant causes</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Mucoepidermoid carcinoma</td>
<td>25</td>
<td>28.40%</td>
</tr>
<tr>
<td>Lymphoma</td>
<td>20</td>
<td>22.70%</td>
</tr>
<tr>
<td>Acinic cell tumor</td>
<td>18</td>
<td>20.40%</td>
</tr>
<tr>
<td>Carcinoma</td>
<td>11</td>
<td>12.50%</td>
</tr>
<tr>
<td>Adenoid carcinoma</td>
<td>6</td>
<td>6.80%</td>
</tr>
<tr>
<td>Mixed salivary gland tumor</td>
<td>5</td>
<td>5.60%</td>
</tr>
<tr>
<td>Total</td>
<td>88</td>
<td>100%</td>
</tr>
<tr>
<td>Secondary malignant causes (metastatic)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>-From malignant scalp lesion</td>
<td>2</td>
<td>66.60%</td>
</tr>
<tr>
<td>Squamous cell carcinoma</td>
<td></td>
<td></td>
</tr>
<tr>
<td>-From nasopharyngeal</td>
<td>1</td>
<td>33.30%</td>
</tr>
<tr>
<td>Squamous cell carcinoma</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Total</td>
<td>3</td>
<td>100%</td>
</tr>
</tbody>
</table>

The examined MDCT criteria for benign and malignant parotid lesions were shown in Table 2. The size criteria were determined by measuring the longest dimension of the lesion, consider, the lesions were classified accordingly to small lesions.
if their size is less than 2cm, intermediate size if they were from 2-4cm and large lesions if their sizes were more than 4cm. This classification was based on National Comprehensive Cancer Network [7]. The margins of enlarged parotid gland in this study were either sharp well-defined that can be delineated along its all margins or ill-defined which means ill definition of one point at least. Thirty-nine out of 45 benign lesions shows well-defined margins, while 54 out of 91 of malignant lesions show ill-defined margins. The parotid lesions were located either in the superficial lobe, deep lobe or both lobes. Most parotid lesions that involved both lobes were malignant which found in 44(50%) and most of the parotid lesions that involve the superficial lobe were benign which found in 27 patients (60%).

Most of benign lesions were single and detected in 30 patients and most of malignant lesions were multiple and detected in 59 patients. The enhancement pattern of the lesions was varying from no enhancement, homogenous enhancement, and heterogeneous enhancement to peripheral or marginal enhancement. It was found that the most common enhancement pattern in the benign lesions were homogenous pattern, as it was determined in 29 out of 45 patients. Most of the malignant parotid lesions exhibited heterogeneous pattern, as it was determined in 57 out of 91 patients [8-10].

Central necrosis was present in 8 benign lesions and in 33 malignant lesions and was highly significant. Another significant MDCT criterion was the extension of the parotid lesions to the para- pharyngeal fat space which was determined in 48 out of 91 malignant lesions and in 20 out of 45 benign lesions. Calcifications were detected in 4 benign lesions and in 12 malignant lesions. The lastly evaluated MDCT criterion was the associated lymph node enlargements which were detected in association with 52malignant lesions and 9 benign lesions.

Each one of the MDCT criteria was statistically tested for its reliability for diagnosis and for differentiating between benign and malignant causes using the pathological diagnosis as a gold standard. In this study, the most reliable MDCT criteria that show statistical significance for diagnosis of parotid gland lesions (significant p values) were the margin (0.001), central necrosis (0.001), enhancement pattern (0.001) and extra-parotid extension (0.001) as well as lesion location (0.05) (Table 2).

Table 2: MDCT criteria of parotid gland lesions with pathological verification (n=136).

<table>
<thead>
<tr>
<th>MDCT Criteria</th>
<th>Pathological Verification</th>
</tr>
</thead>
<tbody>
<tr>
<td>1- Size:</td>
<td></td>
</tr>
<tr>
<td>- Small &lt;3cm</td>
<td>Benign: 9(20%) Primary Malignant 32(36.3%) Secondary Malignant 2(66.6%) P Value 0.5</td>
</tr>
<tr>
<td>- Intermediate 3-5cm</td>
<td>24(53.3%) 36(40.9%) 1(33.3%)</td>
</tr>
<tr>
<td>- Large &gt;5cm</td>
<td>12(26.6%) 20(22.7%) 0</td>
</tr>
<tr>
<td>2-Margin:</td>
<td></td>
</tr>
<tr>
<td>- Well defined</td>
<td>39(86.6%) 36(40.9%) 1(33.3%) P Value 0.001*</td>
</tr>
<tr>
<td>- Ill defined</td>
<td>6(13.3%) 52(59%) 2(66.6%)</td>
</tr>
<tr>
<td>3-Lesion location:</td>
<td></td>
</tr>
<tr>
<td>- Superficial lobe</td>
<td>27(60%) 32(36.3%) 2(66.6%) P Value 0.05*</td>
</tr>
<tr>
<td>- Deep lobe</td>
<td>8(17.7%) 12(13.6%) 1(33.3%)</td>
</tr>
<tr>
<td>- Both lobes</td>
<td>10(22.2%) 44(50%) 1(33.3%)</td>
</tr>
<tr>
<td>4-Multiplicity:</td>
<td></td>
</tr>
<tr>
<td>- Single</td>
<td>30(66.6%) 31(35.2%) 1(33.3%) P Value 0.5</td>
</tr>
<tr>
<td>- Multiple</td>
<td>15(33.3%) 57(64.7%) 2(66.6%)</td>
</tr>
<tr>
<td>5-Enhancement:</td>
<td></td>
</tr>
<tr>
<td>- No enhancement</td>
<td>4(8.8%) 4(4.5%) 0</td>
</tr>
<tr>
<td>- Homogenous</td>
<td>29(64.4%) 28(31.8%) 1(33.3%)</td>
</tr>
<tr>
<td>- Heterogeneous</td>
<td>8(17.7%) 56(63.3%) 1(33.3%)</td>
</tr>
<tr>
<td>- Peripheral</td>
<td>4(8.8%) 0 1(33.3%)</td>
</tr>
<tr>
<td>6-Central necrosis</td>
<td>8(17.7%) 32(36.3%) 1(33.3%) P Value 0.001*</td>
</tr>
<tr>
<td>7-Para pharyngeal fat extension</td>
<td>20(44.4%) 48(54.5%) 0 P Value 0.001*</td>
</tr>
<tr>
<td>8-Calcification</td>
<td>4(8.8%) 12(13.6%) 0 P Value 0.4</td>
</tr>
<tr>
<td>9-Associated LN enlargement</td>
<td>9(20%) 49(55.7%) 3(100%) P Value 0.2</td>
</tr>
</tbody>
</table>

The associated lymph node (LN) enlargements were interpreted for their number and bi-laterality. Most of the enlarged LN (9) that associated benign parotid lesions were unilateral and multiple, and most of the enlarged LN (52) that associated malignant parotid lesions were bilateral and multiple (Table 3). Illustrative cases Figure 1-5.

Table 3: MDCT criteria of associated lymph node enlargement regarding to number and bilaterality (n=136).

<table>
<thead>
<tr>
<th>Causes</th>
<th>Unilateral-Single</th>
<th>Unilateral-Multiple</th>
<th>Bilateral-Single</th>
<th>Bilateral-Multiple</th>
<th>No Involvement</th>
</tr>
</thead>
<tbody>
<tr>
<td>Benign</td>
<td></td>
<td>7 (15.5%)</td>
<td>-</td>
<td>2 (4.5%)</td>
<td>36 (80%)</td>
</tr>
<tr>
<td>Primary malignant</td>
<td>4 (4.5%)</td>
<td>10 (11.4%)</td>
<td>-</td>
<td>35 (39.8%)</td>
<td>39 (44.3%)</td>
</tr>
<tr>
<td>Secondary malignant</td>
<td></td>
<td>1 (33.3%)</td>
<td>-</td>
<td>2 (66.6%)</td>
<td>-</td>
</tr>
</tbody>
</table>

p value is considered significant if less than 0.05.

Figure 1 (a, b & c): 24 years old female patient presented by unilateral parotid mass diagnosed pleomorphic adenoma. (a & b) axial and (c) coronal contrast enhanced MDCT images show well defined hypodense homogenous parotid lesion at the right side involving superficial lobe mainly. The condition is associated with bilateral lymph node enlargement (black arrow in a) and intra parotid lymph node enlargement (black arrow in b). Intact Para pharyngeal fat space (white arrow in b). No calcifications.

Figure 2 (a, b & c): 60 years old man presented by unilateral mass in the parotid region diagnosed Warthin’s tumor. (a & b) Axial & (c) coronal contrast enhanced MDCT images at the level of the parotid gland. They show a well-defined parotid lesion involving both superficial and deep lobes (white arrow in a) with slight encroachment upon the related Para-pharyngeal fat space. It shows a heterogeneous enhancement with areas of central necrosis (white arrow in b). No loco-regional lymphadenopathy. These imaging criteria are characteristic to Warthin’s tumor.

Figure 3 (a, b & c): 38 years old male patient with right diffuse parotid region mass on clinical examination diagnosed lymphoma. (a, b and c) multiple axial contrast enhanced MDCT images show diffuse large size right parotid lesion that seen involving both superficial and deep lobes, it has ill-defined margin, heterogeneous pattern of enhancement (white arrow in a). The lesion has and large area of central necrosis (white arrow in b) and multiple intra parotid lymph node enlargement (black arrows in c). The lesion is seen extended to right Para pharyngeal space in comparison to the left side with lost surrounding fat planes (black arrow in b). It is also extending the right infra-temporal fossa and right cheek. It is seen encasing the right Carotid sheath (black arrow in a). No calcifications.

Discussion

Parotid lesions are a major challenge for radiologists especially those of neo plastic etiologies. Parotid masses in general have relative infrequency reaching up to 1% of head and neck tumors [1] and show no definite specific clinical signs with indeterminate imaging criteria. Many modalities are used including conventional X-ray, sialography, CT and MRI to reach the clinical diagnosis, helping better surgical approach and optimum postoperative outcome.

In the past, the role of CT was limited to evaluation of salivary gland calculi and obstructive or inflammatory disease [3]. So, in our study we try to re-evaluate and update the role of multi-detector CT in diagnosing neo plastic parotid lesions by assessment of various MDCT criteria of parotid lesions that routinely included within patient's report aiming to allow helpful clues to differentiate between benign and malignant neo plastic lesions and optimize the surgical outcome.

Owing to the multiple advantages of MDCT such as the enhanced speed of scan acquisition, the high spatial resolution, the precise timing of multiphasic imaging and the multi-planar reformations (MPR), the anatomic details and localization of parotid gland and its intra and extra- glandular lesions become more sharp and accurate. It also allows good visualization of deep parotid lesions as well as reliable assessment of their size, location, margins, and areas of necrosis, hemorrhage, calcification, and extension to adjacent structures as well as determination of perineural spread especially by curved planar reconstruction images (CPR) [3].

In our study, the mean age of the patients (44.4±22.2) as well as the prevalence of the most common benign and malignant parotid tumors; pleomorphic adenoma and mucoepidermoid carcinoma respectively were in concordance with Rastogi et al. [11], who reported that in patients older than 40 years, the most causes of enlarged parotid gland are neo plastic lesions either benign or malignant. The most common benign lesions are the pleomorphic adenoma and Warthin's tumor and the most common malignant tumor are the mucoepidermoid carcinoma and adenoid cystic carcinoma.

Our patients were collected from oncology center, all studied groups proved pathologically to have parotid neo plastic lesions. The benign neoplasms were detected in 45(33%) patients, 88(64.7%) patients had primary malignant tumors and 3(2.2%) patients had secondary malignant tumors. These values are not matched with other studies [1,3,12] due to the fact that the study is biased based only upon neo plastic group of patients. Other factors may be responsible such as increased prevalence of the disease that may be related to the ethnicity; however this needs further studies which are out of the scope of our research. In this study the MDCT criteria of all parotid lesions were evaluated individually. Regarding to the size criterion, the parotid lesions were classified into small, intermediate or large sizes. However despite of these variations in tumors size, there is no statistically significant difference between the size of benign and malignant lesions. Our results are matching with Brunese et al. [13], who reported no significant differentiation between benign and malignant lesions by the lesion size.

Regarding to the margin of the parotid lesions, the margin was classified into well-defined and ill-defined on both pre- and post-contrast imaging. When the margin was blurred or ill-defined at only one point, the whole margin was deemed ill-defined. Most of benign tumors have well defined margins. The margins of malignant parotid tumors may be confusing; low grade malignancy may have well-defined margins that mimicking the benign mixed tumors where the high–grade malignancy usually...
has invasive margins with irregular outlines. Our results are
matching with Lu et al. [14], who evaluate the margin of benign
and malignant parotid lesions of 133 patients. This imaging
point is highly conclusive in our study to differentiate between
benign and malignant parotid lesions.

Most of the benign parotid tumors were located within
the superficial lobe while most of the malignant lesions were
located within both parotid lobes. Our results were matching
with to Lu YC et al (14), who suggest that, a parotid tumor
located in the superficial lobe, with a round or oval contour and
sharp margin, is more likely to be a benign tumor; otherwise,
it might be a malignant tumor. Our study is also matching with
A. Christe et al. [15], who postulated that, the predilection of
depth lobe involvement was only indicative but not significant
for malignancy, because inflammatory disease and some benign
tumors, like schwannoma and Warthin tumors, also affected the
deep lobe. However, involvement of both lobes was significant
for malignant lesions, like the superficial lobe affinity seen with
benign tumors.

The extra-glandular extension of parotid lesions is a grave
prognostic finding, in this study it was considered when the
parotid lesions extend to the prestyloid parapharyngeal space.
Accordingly, parapharyngeal fat extension was detected in 48
malignant lesions and 20 benign lesions. This MDCT criterion
shows highly significant diagnostic reliability in our study. The
number of the parotid lesions was determined in each patient
and we found that most of benign tumors were single while most
of malignant tumors were multiple. However, this MDCT finding
shows no statistical significant difference between the benign
and malignant lesions. This sign is matched with Thoeny et al.[3],
who postulated that multiple parotid lesions may be benign or
malignant, and the most common multiple parotid gland tumors
are the Warthin’s tumor followed by acinic cell carcinoma.

The enhancement pattern of the lesion is another important
MDCT criterion for assessment of the parotid lesions. The
enhancement of the lesion seems to imply increased its vascularity
and suggests a wide differential diagnosis. In this study, most
of the malignant parotid lesions exhibited heterogeneous
enhancement pattern, as it was determined in 57 out of 91
patients, while most of the benign lesions showed homogenous
enhancement pattern, as it was determined in 29 out of 45
patients. The heterogeneous enhancement can be explained by
central necrosis and vascularity. This study is nearly symmetrical
to that of DC Howlett et al. [16]. Enhancement pattern is a very
helpful imaging criterion in our study to differentiate between
benign and malignant parotid lesions.

One of the important MDCT criterion that help in differentiating
benign from malignant parotid lesions in this study was the
central necrosis, it was found in 33 out of 91 patients with
malignant lesions and in 8 out of 45 patients with benign
lesions. Allen et al. [17], discussed five cases of pleomorphic
adenoma and conclude that central necrosis is not necessary
specified to malignant lesions especially in cases of pleomorphic
adenoma and Warthin’s tumors which have high prevalence of
cystic/necrotic regions in benign lesions. This study did not
agree with the results of our study which may be explained
by recent advances in MDCT techniques including high spatial
resolution with thin collimation that leading to better tissue
characterization and picking central necrosis more accurately.

Other MDCT criteria was the calcification of the parotid
lesions which was only demonstrated in 16 (11.7%) out of 136
patients. Izum et al. [18], reported that calcifications of the neck
are uncommon and mainly seen in cases of inflammatory lesions
and Sjögren’s syndrome.

The lastly evaluated MDCT criterion was the associated lymph
node enlargement that was found in 61 out of 136 cases. The
majority of patients (49 patients) with lymph node enlargement
had primary malignant lesions and all the 3 patients with
secondary metastases also had lymph node involvement. These
results were in agreement with Stennert et al. [19], who reported
high incidence of lymph node metastasis in major salivary gland
cancer with importance of neck dissection in surgical approach.

Limitations

The major limitation of this study is the biased nature of the
study because we were concerning about imaging of neo plastic
parotid group of patients who were directed to tertiary oncology
center for treatment. It shows different results than literature
regard the prevalence of benign and malignant parotid lesions.
Another limitation of this study is that the MDCT criteria of all
parotid lesions were evaluated individually, combined MDCT
criteria assumed to have a powerful diagnostic reliability in
distinguishing benign from malignant parotid lesions, further
study is recommended.

Conclusion

Rapid advances of MDCT techniques giving rise to get an
optimum parotid lesion characterization in a simple rapid and
low cost maneuver. Multiple promising MDCT imaging criteria
can be valuable in differentiation between benign and malignant
parotid tumors. Imaging is forming a crucial triad with clinical
data and surgical approach to get the satisfactory operative
outcome.

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