Diagnostic Accuracy of 4D-CT for Parathyroid Adenomas and Hyperplasia

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BACKGROUND AND PURPOSE: 4D-CT is a novel method of multiphase CT imaging. When used to localize parathyroid adenomas and hyperplasia, this technique may allow for more robust diagnostic accuracy than traditional sonography and nuclear scintigraphy techniques. The purpose of our study is to assess the accuracy of 4D-CT for localizing pathologically proved parathyroid adenomas and hyperplasia found during surgery.

MATERIALS AND METHODS: A total of 35 pathologically proved cases of parathyroid adenoma and hyperplasia were retrospectively reviewed between January 2009 and March 2011. Inclusion criteria were availability of final surgical pathology and performance of preoperative 4D-CT. No cases were excluded. Sensitivity, specificity, and accuracy of 4D-CT were ascertained including both the side and quadrant of the pathologically proved lesion.

RESULTS: Of the 35 pathologically proved cases collected over the study period, 32 (sensitivity = 91%) patients were found positive for parathyroid disease using 4D-CT, including 3 cases of multigland disease. For lateralization of single-gland disease, 4D-CT demonstrated an accuracy of 93%. 4D-CT revealed a suboptimal 44% sensitivity, but 100% specificity, for multigland disease.

CONCLUSIONS: 4D-CT demonstrated a high diagnostic accuracy for single and multigland disease in our cohort. Importantly, 4D-CT accurately lateralized single-gland adenomas in >90% of cases, allowing the surgeon to employ a directed operative approach. 4D-CT also showed a very high specificity for the detection of multigland disease.

ABBREVIATIONS: CTDI = CT dose index; NPV = negative predictive value; PPV = positive predictive value; PTH = parathyroid hormone.
ness. Imaging extended from the carina to the bottom of the mandibu-
lar teeth in a caudal to cranial direction, in a plane parallel to the
hard palate. Each examination included a scout scanogram, noncon-
trast acquisition, and multiphase imaging following the administra-
tion of intravenous contrast.

Iohexol 300 (Omnipaque; GE Healthcare) nonionic iotaded con-
trast was administered at a dose of 2 mL per kg patient weight to
a maximum total of 120 mL. Intravenous infusion rate was 4 mL per
second via an 18-gauge catheter.

The diagnostic multiphase CT was acquired at 1.25-mm section
thickness. The helical postcontrast sequence was obtained 25 seconds
after the start of intravenous contrast administration. Following a
subsequent 30-second delay, a delayed phase sequence was acquired.
Axial, sagittal, and coronal 2.5-mm reformations were generated.

Image Analysis
Images were reviewed prospectively on a PACS workstation by a sin-
gle attending neuroradiologist with over 20 years of experience in
head and neck imaging, and a certificate of added qualification in
neuroradiology. Biochemical information was available at the time
of interpretation, including the preoperative PTH levels. Images were
reviewed and determined to be positive if a soft tissue lesion in the
expected location of parathyroid tissue (orthotopic or ectopic) dem-
onstrated early arterial enhancement and qualitatively appreciable
contrast washout on early or delayed images. The radiology re-
port provided a descriptive location for our referring surgeons, with
anatomic landmarks. Any coexistent pathology that was deemed to
potentially alter patient management was also noted.

Statistical Analysis
SAS version 9.2 (SAS Institute, Cary, North Carolina) was used for all
statistical analysis. Categoric variables are presented as frequen-
cies and percents, while continuous variables are presented as means
and standard deviations. To determine how well 4D-CT performs in
preoperative localization, we used diagnostic accuracy measures com-
paring 4D-CT to the “gold standard” of surgical pathology. Correct
localization of the lesions was ascertained by using side and quadrant
of the neck. Diagnostic accuracy measures for both side and quadrant
location were determined by using sensitivity, specificity, PPV, NPV,
and accuracy, with corresponding 95% confidence intervals calcu-
lated by using the exact binomial distribution method.

Results
All studies were performed in patients with elevated PTH and
a clinical diagnosis of primary hyperparathyroidism. A total of
42 4D-CT studies were performed over the study period. Of
these, 35 patients went to surgery with final pathology avail-
able and were included in the analysis. The remaining 7 pa-
tients were lost to follow-up or refused surgery. No cases were
negative for parathyroid hyperplasia on surgical exploration.
Table 1 displays the descriptive statistics for our cohort. Of the
35 cases, 32 cases of parathyroid adenoma or multigland hy-
perplasia were detected by using 4D-CT (sensitivity = 91%, 95% CI
77%, 98%). To calculate diagnostic accuracy of
4D-CT to correctly determine side and quadrant location of
the lesions, we looked at the 32 cases that were positive by both
pathology and 4D-CT. Of these 32 cases, 5 cases were removed
because multiple glands were involved. The final single-gland
dataset contained 27 cases. Table 2 displays the distribution of
the pathology and 4D-CT findings from these 27 individuals.

<table>
<thead>
<tr>
<th>Table 1: Descriptives</th>
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</thead>
<tbody>
<tr>
<td><strong>Variable</strong></td>
</tr>
<tr>
<td>Age (yr)</td>
</tr>
<tr>
<td>Female gender (n)</td>
</tr>
<tr>
<td>BMI (kg/m²)</td>
</tr>
<tr>
<td>Weight (kg)</td>
</tr>
<tr>
<td>Lesion weight (mg)</td>
</tr>
<tr>
<td>Prior neck surgery (n)</td>
</tr>
<tr>
<td>Single-gland disease (n)</td>
</tr>
<tr>
<td>Multigland disease (n)</td>
</tr>
<tr>
<td>Peak preoperative parathyroid hormone (pg/mL)</td>
</tr>
<tr>
<td>Postoperative parathyroid hormone (pg/mL)</td>
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<table>
<thead>
<tr>
<th>Table 2: Distribution of findings</th>
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</thead>
<tbody>
<tr>
<td><strong>Lesion detection</strong></td>
</tr>
<tr>
<td>Positive</td>
</tr>
<tr>
<td>Negative</td>
</tr>
<tr>
<td><strong>Lesion location</strong></td>
</tr>
<tr>
<td>Right superior</td>
</tr>
<tr>
<td>Right inferior</td>
</tr>
<tr>
<td>Left superior</td>
</tr>
<tr>
<td>Left inferior</td>
</tr>
<tr>
<td><strong>Lesion side</strong></td>
</tr>
<tr>
<td>Right</td>
</tr>
<tr>
<td>Left</td>
</tr>
<tr>
<td><strong>Lesion z-axis</strong></td>
</tr>
<tr>
<td>Superior</td>
</tr>
<tr>
<td>Inferior</td>
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<table>
<thead>
<tr>
<th>Pathology Findings</th>
<th>4D-CT Findings</th>
</tr>
</thead>
<tbody>
<tr>
<td>n</td>
<td>27</td>
</tr>
<tr>
<td>Female gender</td>
<td>26</td>
</tr>
<tr>
<td>BMI</td>
<td>26.8</td>
</tr>
<tr>
<td>Weight</td>
<td>73.2</td>
</tr>
<tr>
<td>Lesion weight</td>
<td>566.7</td>
</tr>
<tr>
<td>Prior neck surgery</td>
<td>2</td>
</tr>
<tr>
<td>Single-gland disease</td>
<td>29</td>
</tr>
<tr>
<td>Multigland disease</td>
<td>7</td>
</tr>
<tr>
<td>Peak preoperative parathyroid hormone</td>
<td>190.3</td>
</tr>
<tr>
<td>Postoperative parathyroid hormone</td>
<td>29.5</td>
</tr>
</tbody>
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Note: Freq indicates frequency.

Fig 1 demonstrates the classic enhancement pattern seen with
a parathyroid adenoma. Figs 2 and 3 demonstrate ectopic ad-
enomas detected by using 4D-CT.

We then determined the diagnostic accuracy of 4D-CT to
correctly lateralize the adenoma found at surgery. For diag-
nostic accuracy, pathology was treated as the gold standard.
4D-CT test results were correlated to pathology, and each CT
result was evaluated and classified as true-positive, true-nega-
tive, false-positive, or false-negative in terms of the ability to
accurately predict laterality (right versus left). It is important
to note that false-positives and false-negatives were equally
weighted. The sensitivity of 4D-CT to correctly identify a
right-sided adenoma was 92% (61.5%, 99.8%), and the spec-
ificity of 4D-CT to correctly identify a left-sided adenoma was
93.3% (68.1%, 99.8%). The overall accuracy of 4D-CT to cor-
correctly identify adenoma side (assuming 4D-CT was positive)
was 93% (76%, 99%). Table 3 presents all diagnostic accuracy
measures for 4D-CT lateralization.

We also evaluated the ability of 4D-CT to precisely localize
an adenoma. For this analysis, we looked at the 25 patients
with single-gland disease that was correctly lateralized by 4D-
CT. Again, pathology was treated as the criterion standard,
and 4D-CT results were classified as true-positive, true-nega-
tive, false-positive, or false-negative in terms of the ability to
accurately predict location (superior versus inferior). As be-
fore, false-positives and false-negatives are equally weighted.
The sensitivity of 4D-CT to correctly determine superior ade-
oma quadrant location was 85.7% (57.2%, 98.2%), and the
specificity to correctly determine inferior adenoma quadrant

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Note: Freq indicates frequency.
location was 100% (71.5%, 100%). The overall accuracy of 4D-CT to correctly identify precise adenoma location (assuming 4D-CT was positive for adenoma and correctly identified side) was 92% (74%, 99%). Table 4 provides all diagnostic accuracy for 4D-CT quadrant localization.

Last, we looked at patients with multigland disease (including 4-gland hyperplasia and double adenomas). By pathology, 7 patients in total were determined to have multigland involvement. Of these, 2 (29%) were called negative for any parathyroid lesion on 4D-CT. We looked at diagnostic accuracy for multigland disease, both including and excluding patients called negative by 4D-CT. As before, pathology was treated as the criterion standard, with true-positives defined as those that CT correctly called lesions at multiple locations, false-positives as those that CT incorrectly called not multigland (but are truly multigland), true-negatives as those that CT correctly called not multigland, and false-negatives as those that CT incorrectly called multigland (but are truly not multigland). Looking at all 35 patients, 4D-CT has a low sensitivity of 42.9% (9.9%, 81.6%) in detecting multigland dis-
However specificity to rule out a multigland disease in this population was perfect (specificity = 100%, 95% CI, 87.7%, 100%). Looking at the 32 patients in whom both pathology and 4D-CT were positive, 4D-CT had a slightly higher sensitivity of 60% (CI, 14.7%, 94.7%) for detecting multigland disease. As before, the specificity of 4D-CT in ruling out multigland disease in this population was perfect (specificity = 100%, 95% CI, 87.2%, 100%). Table 5 displays the diagnostic accuracy of 4D-CT in detecting multigland disease, including those patients called negative by 4D-CT.

### Discussion

4D-CT demonstrated high accuracy for the determination of both lesion side and quadrant location for single-gland disease. Our series suggests that 4D-CT agrees with the pathology findings in most of the cases and is a reliable choice for parathyroid adenoma detection. Importantly, 4D-CT accurately lateralized single-gland adenomas in 93% of cases, allowing the surgeon to perform a unilateral neck dissection with associated decreased morbidity and length of hospital stay. The ability to accurately lateralize the side of disease is a commonly used end point in parathyroid imaging because it enables a directed surgical approach. By comparison, sonography has a cited sensitivity for lateralization ranging from 57% to 86%.

4D-CT appears to hold potential for detection of multi-
gland disease. Although this technique demonstrated suboptimal sensitivity of 44% in patients with multigland disease, it was 100% specific for ruling out multigland disease, as no cases that were thought to have multigland disease by 4D-CT had single-gland adenomas at pathology. These results compare favorably to those reported for sonography and sestamibi techniques. A review by Sugg et al., including 23 patients with multigland disease, found that the combination of sonography and sestamibi imaging correctly predicted multigland disease in 30% of patients, incorrectly called single-gland disease in 30%, did not identify abnormal parathyroid glands in 30%, and yielded discordant results in 10%. In our cohort, 2 of the 3 false-negatives for parathyroid lesion detection were patients with multigland disease. Multigland lesions are particularly challenging to detect prospectively, likely related to the small lesion size. We would further posit, as other authors have suggested, that in the case of a negative 4D-CT examination, there is presumed multigland disease. Importantly, some authors suggest a nonselective operative approach in cases of negative or discordant imaging studies due to the increased likelihood of multigland pathology.2,21,13

Unique angiographic and perfusion characteristics of parathyroid adenomas were described by Doppman and others in the 1970s, and used to identify abnormal hypervascular glands with preoperative digital subtraction angiography.9,14 A characteristic “blush” has been described following the injection of intra-arterial contrast, indicating the presence of an abnormal gland that is hypervascular compared with normal thyroid and parathyroid tissue. Similar principles are applied to contrast-enhanced CT, where hyperenhancement, location, and size are important discriminators in detecting parathyroid adenomas.4 4D-CT appears to further enhance detection, with the advantages of noncontrast imaging to distinguish the gland from surrounding thyroid tissue, early postcontrast imaging to assess for a hypervascular gland, and delayed postcontrast imaging to detect regions of altered enhancement and abnormal contrast retention.

It is important to be cognizant of the radiation dose delivered with CT, particularly multiphase CT. We use dose reduction techniques with 4D-CT, as described by Welling et al., including automatic tube current modulation and limiting the superior field of view to the mandibular teeth. Despite these techniques, 4D-CT delivers a significant radiation dose. Calculated CTDIvol measurements range from 19.8 mGy (at 150 mA) to 39.7 mGy (at 300 mA) per phase of imaging. These parameters are comparable with recently reported CTDIvol values for neck CT ranging from 23.0 to 34.7 mGy.13 We have chosen not to exclude the noncontrast examination, as we think it provides important diagnostic information, particularly in its ability to show attenuation differences in lesions contiguous with the thyroid gland. The risk-benefit discussion must weigh the cost, morbidity, and risk of formal neck exploration with the exposure from 4D-CT. The examination will likely only require performance on a single occasion.

There are several limitations to our study. First, this study is a retrospective review and was not designed to make a direct prospective comparison of the accuracy of sonography, sestamibi, and 4D-CT imaging for the detection of parathyroid pathology. Second, there is a selection bias for cases of ectopic and multigland disease because these are more difficult to diagnose on traditional imaging modalities and thus more likely to progress to 4D-CT imaging. Our patient population may be skewed by this factor, with a relatively higher proportion of these subtypes of adenomas. Third, although the technique of 4D-CT at our institution has been performed with high accuracy, variability in diagnostic accuracy may occur across institutions, secondary to interpreter experience and study technique.

Nonetheless, our series demonstrates that 4D-CT is a valuable tool for appropriate operative planning. 4D-CT also shows a higher sensitivity and accuracy than commonly cited for sonography or sestamibi imaging. We believe the accuracy of this technique is particularly impressive when considering the patients referred for 4D-CT frequently had equivocal or indeterminate findings on prior sonography and sestamibi SPECT studies. The multiplanar reconstruction capabilities provide beneficial spatial information for optimal operative planning and intervention.

Conclusions

4D-CT demonstrated a high diagnostic accuracy for single-gland disease in our cohort and high specificity in multigland disease. Importantly, 4D-CT predicted the correct side of single-gland adenomas in 93% of cases, allowing the surgeon to utilize a directed operative approach. Potential difficulties with the technique include radiation exposure and necessity of iodinated contrast media.


References