Value of SPET/CT Image Fusion in the Assessment of Neuroendocrine Tumours with $^{111}$In-Pentetreotide Scintigraphy

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Summary.—Objective: The purpose of this study was to evaluate the impact and clinical value of anatomical-functional image fusion in the study interpretation and clinical management of patients with neuroendocrine tumours (NET) using somatostatin receptor scintigraphy (SRS) and combined transmission and emission tomography – single-photon emission tomography/CT (SPET/CT).

Material and methods: Twelve patients (8 female and 4 male; age range 32-74 years, mean 66 years) with proven or clinically suspect NET were studied with routine planar SRS and SPET/CT at 2 and 24 hours after injection of $^{111}$In-Pentetreotide. Seven patients came for staging/follow-up and 5 patients for primary tumour localization with staging. Analysis of fused images (SPET/CT) was done on a patient basis, with separated evaluation of SPET, low-dose CT images and fusion images. The gold standard for presence or absence of malignancy was pathology or clinical and imaging follow-up data.

Results: SRS was negative in 6 patients and positive in 6. SPET/CT provided no additional information in 6 patients, including all 6 negative studies. SPET/CT improved localization of SPET detected lesions in 6 positive studies. It defined the extent of the disease and showed bone involvement in 3 of the 6 positive studies. SPET/CT affected the diagnostic interpretation in 6 patients (50%) and induced changes of management in 3 (25%).

Conclusion: The results of this study indicate that combined anatomical-functional imaging with SPET/CT significantly improves tumour localization and characterization, contributing to a better therapeutic management of patients with NET.


INTRODUCTION

Neuroendocrine tumours (NET) are rare neoplasms corresponding to a very heterogeneous group, which differ in histology, hormone production and clinical behaviour. Nuclear Medicine plays an important role in the management of patients with NET. The outcome of patients with NET is significantly affected by accurate tumour localization and precise determination of the extent of disease. Management of these patients is primarily based on surgical removal of the primary lesion and single...
metastases, as these slow growing tumours are only minimally responsive to systemic chemotherapy. In cases of evidence of multifocal metastatic spread, the differentiated tumour may benefit from treatment with therapeutic doses of cold or radio-labelled ligand. Recent studies indicate that somatostatin receptor scintigraphy (SRS) is more sensitive than sonography, CT, MRI and angiography, for identifying these tumours and their metastases. However, the shortcomings of SRS in providing precise anatomical localisation of lesions often necessitate correlation with high-resolution anatomical imaging modalities.

SRS is a functional imaging modality based on preferential binding of 111In-pentetreotide to somatostatin receptors subtypes II and V. It has the potential to define the extent of metastatic disease, to localize the primary occult tumour site, and to identify the receptor status of metastases for potential octreotide treatment. However, the detection of NET remains challenging, due to their usually reduced dimensions and poorly defined anatomical localization. Its specificity may also be affected by tracer uptake at physiological sites or benign conditions. Because of these limitations, SRS may benefit from co-registration with CT, with acquisition of nuclear medicine and CT data during a single imaging session. The implementation of these SPET/CT imaging systems has the advantages of minimizing the errors described with co-registration of independently performed studies and optimizing anatomic localization of foci of increased single photon emitting tracer uptake, in addition to a significant improvement in attenuation correction.

The purpose of this study was to evaluate the contribution of hybrid SPET/CT imaging to scintigraphic evaluation, assessing as well fusion related changes and the clinical impact in patients with NET.

**MATERIAL AND METHODS**

Twelve patients with proven or clinical suspicion of NET were referred for SRS between February 2003 and January 2004. The patient population included 8 females and 4 males, aged from 32 to 74 years (mean 57 ± 13 years).

Seven patients came for staging/follow-up and five came for primary tumour localization with staging (table I). Scintigraphy was performed after i.v. administration of 111-222 MBq of 111In-Pentetreotide (Octreoscan, Mallinckrodt Medical). Planar views and SPET/CT were obtained using a dual head, variable angle gamma camera equipped with a medium energy collimator and a low-dose X-ray CT transmission system, mounted on the same slip-ring gantry (Discovery VH, with CoDe VC & Hawkeye options, GE Medical Systems). The combined system that provides hybrid functional (SRS) and anatomic (low-dose CT) images allows for acquisition of both procedures in the same setting, without changing the patient’s positioning. Planar views were recorded for 15 minutes or up to 500 Kcts from skull to mid-thigh at 2 h and 24 h after injection. SPET/CT of abdomen and/or chest and/or pelvis was performed at 24 h after injection.

For the CT component of the study, a cross-sectional anatomic transmission map was acquired as the system rotated around the patient, with a rotation speed of 2.6 r.p.m. Each slice acquisition, termed “half-scan” was performed over 240°, taking 16 sec. acquisition time for each transaxial slice. The x-ray tube is operated at 140 KVp, 2.5 mA. Multiple slices were obtained by moving the table by a slice step (10 mm) before acquiring the next slice. The full field-of-view, consisting of 40 slices, was completed in 10 minutes. The transmission data were reconstructed using filtered back-projection to produce cross-sectional attenuation images in which each pixel represents the estimated attenuation of the imaged tissue. Radiation exposure of the patient following a transmission scan, calculated on the basis of phantom studies, ranged from 1,3-3,5 mGy at the centre to 5 mGy at the surface (CTDIw = 3.54 mGy).

Following transmission, the emission part of the study was acquired. Acquisition parameters included 15 minutes or up to 500 Kcts from skull to mid-thigh at 2 h and 24 h after injection.

<table>
<thead>
<tr>
<th>Tumour type</th>
<th>Indication for SRS</th>
</tr>
</thead>
<tbody>
<tr>
<td>Primary tumour staging</td>
<td>Staging follow-up</td>
</tr>
<tr>
<td>Total</td>
<td></td>
</tr>
<tr>
<td>Pancreatic</td>
<td>2</td>
</tr>
<tr>
<td>Gastrointestinal carcinoid</td>
<td>1</td>
</tr>
<tr>
<td>Bronchopulmonary carcinoid</td>
<td>1</td>
</tr>
<tr>
<td>Metastatic NET (suspected)</td>
<td>1</td>
</tr>
<tr>
<td>Total</td>
<td>5</td>
</tr>
</tbody>
</table>

SRS: somatostatin receptor scintigraphy; NET: neuroendocrine tumour.
a 360° rotation, 128 × 128 matrix, 6° angle step and acquisition time of 45 sec per frame for a total 33 min and 45 sec per study. Reconstruction was performed using the ordered subsets expectation maximization (OSEM) technique (2 iterations), to avoid the artefacts seen with filtered back projection in areas adjacent to intense tracer activity. The resultant emission images were inherently registered to the anatomical maps. These matching pairs of X-ray and nuclear medicine images were fused using the eNTEGRA workstation software (General Electric Medical Systems) and images of overlying transmission (CT) and emission (SPET) data were generated. Data acquired during SPET/CT were used for anatomic localization of the scintigraphic findings. 

SRS, including planar and SPET study were interpreted independently of SPET/CT images by a team of nuclear medicine physicians. Subsequently, data obtained from SPET/CT images analysis were compared to the reports of SRS from the independent analyses. SPET/CT data were confirmed with pathological findings, when available, or with clinical and radiological follow-up, during a 4 to 12 month period. SPET/CT was considered to improve image interpretation once it provided a better localization of sites of increased tracer uptake, or when it excluded disease in sites of physiological or benign tracer uptake. Any change or adjustment in the patient management, whether diagnostic or therapeutic, due to the additional information provided by SPET/CT was recorded.

RESULTS

Six patients had a negative SRS study, and six had receptor-positive findings.

Among the 6 patients with negative SRS, no further evidence of disease was documented. They included 1 patient with bronchopulmonary NET, 1 patient with rectal NET (excised) and 2 patients with pancreatic NET, who came for staging/follow-up. Two patients with suspected carcinoid came for primary tumour localization and staging. SPET/CT provided no added value to image interpretation and to clinical decision making in all 6 patients with negative SRS.

Among the 6 patients with positive SRS, disease was confirmed pathologically in 2 patients (1 by surgery and 1 by bronchoalveolar lavage analysis), radiologically in 5 patients (CT or MRI) and by clinical follow-up in one patient (suspected for insulinoma). SPET/CT improved the image interpretation in all 6 SRS positive patients (100 % of patients with abnormal SRS, and 50 % of the whole group). Of these, SPET/CT contributed to a better anatomic localization in 3 patients, referred for detection of primary tumour and 3 patients who came for staging/follow-up. SPET/CT detected bronchopulmonary tracer uptake (1 patient) and precisely confined tracer uptake in pancreas (2 patients) of 3 patients referred for primary tumour detection. SPET/CT also identified unsuspected bone extension in 2 patients, during follow-up.

SPET/CT results affected patient management in 3 patients, 25 % of the total population and 50 % of the cases where SPET/CT improved image interpretation. It led to a surgical therapeutic approach with chemotherapy in one patient with hepatic lesions in whom was detected a primitive pancreatic NET (fig. 1). In one patient, referred for follow-up after surgery and chemotherapy, SRS-SPET/CT detected bone metastases, which were no responsive to chemotherapy, and was submitted to cold octreotide first in mensal administrations, then weekly and fi-
nally in continued infusion due to worsening of the general condition (fig. 2). This patient died at 4 months follow-up. In another patient with hepatic lesions referred for primary tumour localization, SPECT/CT detected a bronchopulmonary carcinoid (confirmed by bronchoalveolar lavage analysis). This patient was considered not suitable for surgery and was submitted to chemotherapy.

DISCUSSION

Somatostatin receptors are present on almost all neuroendocrine cells. As a consequence, \(^{111}\)In-Pentetreotide scintigraphy has shown a high sensitivity for the diagnosis of both primary and metastatic tumour lesions\(^6,7\). Although peptide receptor scintigraphy may be useful as a tumour localizing technique in a variety of diseases where it may detect tumour localizations prior to or in addition to conventional imaging, it will never be competitive with, for instance, CT or MRI in anatomic delineation of these disease sites\(^8\). The incidence of multiple organ involvement in patients with NET is high. The knowledge of the precise extent of disease plays a major role in determining the appropriate treatment option and choice of treatment procedures such as chemotherapy, cytoreductive surgery, embolization, or liver transplantation\(^3\). Diagnosis of bone metastasis is of prognostic significance, indicating short survival and patients may also be spared unnecessary surgery.

In the present study, fusion of functional and anatomic images obtained sequentially on the same imaging device showed an additional value in the diagnostic capacities of SRS-SPECT and CT. SPECT/CT improved SRS image interpretation in 6 of the 12 study group patients (50 %), not only due to a more accurate localization of tumour sites but also due to the x-ray based attenuation correction, which results in improved image quality for SPECT images. Among these 12 patients, 6 patients were referred for detection of primary tumour and staging and 6 patients were evaluated for extent of disease and for follow-up. The primary tumour was detected in 3 patients (2 pancreatic NET and 1 bronchopulmonary carcinoid) and bone involvement in 3. SRS-SPECT/CT guided the therapeutic approach in 3 cases (25 %).

The results of our study, evaluating combined SPECT/CT in patients with NET are in agreement with the published data.

CONCLUSION

The results of this study indicate that the introduction of dual-head gamma camera with an integrated x-ray transmission system (low dose CT) for attenuation correction, anatomical mapping and image fusion, improves image interpretation and provides correct anatomic localization of SRS findings in patients with NET. Hybrid imaging is an accurate, simply achieved method of registration of functional and anatomic data which enhances the already unique role of SRS in the assessment of these tumours. It improves guidance of invasive diagnostic and therapeutic procedures and may help in monitoring therapeutic outcomes.

REFERENCES


