Imaging of Adrenal Incidentalomas with PET Using $^{11}$C-Metomidate and $^{18}$F-FDG

Heikki Minn, MD1,2; Anna Salonen, BSc1; Johan Friberg, MD1; Anne Roivainen, PhD1; Tapio Viljanen, MS1; Jaakko Långsjö, MD1; Jorma Salmi, MD1; Matti Välimäki, MD4; Kjell Någren, PhD1; and Pirjo Nuutila, MD1,5

1Turku PET Centre, University of Turku, Turku, Finland; 2Department of Oncology and Radiotherapy, Turku University Central Hospital, Turku, Finland; 3Department of Internal Medicine, Tampere University Hospital, Tampere, Finland; 4Department of Internal Medicine, Helsinki University Hospital, Helsinki, Finland; and 5Department of Internal Medicine, Turku University Central Hospital, Turku, Finland

Our aim was to evaluate the use of PET with $^{11}$C-metomidate and $^{18}$F-FDG for the diagnosis of adrenal incidentalomas. **Methods:** Twenty-one patients underwent hormonal screening before dynamic imaging of the upper abdomen with $^{11}$C-metomidate, and for 19 of these 21 patients, static $^{18}$F-FDG imaging followed. Uptake of $^{11}$C-metomidate and $^{18}$F-FDG in incidentalomas was quantified and correlated with the hormonal work-up and the mass size on CT (median, 2.5 cm; range, 2–10 cm).

**Results:** The final diagnoses were hormonally active adenoma ($n = 7$), nonsecretory adenoma ($n = 5$), adrenocortical carcinoma ($n = 2$), pheochromocytoma ($n = 2$), benign noncortical tumor ($n = 3$), normal adrenal ($n = 1$), and malignant noncortical tumor ($n = 3$). Diagnosis was established at surgery ($n = 9$), percutaneous biopsy ($n = 4$), or follow-up ($n = 8$). The highest uptake of $^{11}$C-metomidate, expressed as standardized uptake value (SUV), was found in adrenocortical carcinoma (SUV = 28.0), followed by active adenomas (median SUV = 12.7), nonsecretory adenomas (median SUV = 12.2), and noncortical tumors (median SUV = 5.7). Patients with adenomas had significantly higher tumor–to–normal-adrenal $^{11}$C-metomidate SUV ratios than did patients with noncortical tumors. $^{18}$F-FDG detected 2 of 3 noncortical malignancies but failed to detect adrenal metastases from renal cell carcinoma. At all inactive and most active adenomas were difficult to detect with $^{18}$F-FDG, and the mass size on CT (median, 2.5 cm; range, 2–10 cm).

**Conclusion:** $^{11}$C-Metomidate is a promising PET tracer to identify incidentalomas of adrenocortical origin. $^{18}$F-FDG should be reserved for patients with a moderate to high likelihood of neoplastic disease.

**Key Words:** adrenal gland; PET; radionuclide imaging; metomidate; $^{18}$F-FDG

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The rate of finding unexpected adrenal masses, so-called incidentalomas, has been increasing because of increased use of CT, MRI, and ultrasound to study symptoms potentially originating from the abdomen (1). Patients with incidentalomas are subjected to hormonal screening and, sometimes, an extensive diagnostic work-up, especially if malignancy is suspected (2). Patients with a history of oncologic disease clearly have a high risk (32%–75%) for adrenal metastasis (2–4), and adrenocortical adenoma accounts for 36%–94% of incidentalomas diagnosed in patients without a history of cancer (1,2,5). Other final diagnoses for incidentalomas include pheochromocytomas, lipomas, cysts, or false adrenal masses actually arising from a nearby organ such as the liver, kidney, or stomach (1,2).

Adrenal tumors are found incidentally in up to 5% of patients, illustrating the need for an effective strategy to determine whether a patient should be treated surgically, pharmacologically, or not at all (1,2). Besides clinical history and hormonal profile, mass size and imaging characteristics play an important role in the diagnostic algorithm, which by no means has matured to a universally approved form (6). In some patients, the size, morphologic appearance, attenuation on CT, and growth pattern of the adrenal mass do not disclose its nature. A positive intracellular lipid signal on MRI suggests a benign adenoma, but in the absence of characteristic features on MRI or CT, functional imaging with radionuclides should be considered for differential diagnosis (7,8).

Recently, PET using $^{11}$C-labeled metomidate was introduced for the identification of indeterminate adrenal masses (9,10). Metomidate is an inhibitor of 11β-hydroxylase, a key enzyme in the biosynthesis of cortisol and aldosterone by the adrenal cortex. Our goal was to evaluate $^{11}$C-metomidate PET in the diagnosis of incidentalomas and to study whether uptake of tracer is associated with adrenal cortex function.

**MATERIALS AND METHODS**

**Patients**

Twenty-one patients (14 female, 7 male) aged 21–79 y participated in the study. Written informed consent to undergo PET...
imaging was obtained from all patients, and the consent form and study protocol were approved by the Ethical Committee of the Local Hospital District. Pertinent clinical characteristics of the patients and their final diagnoses after PET are given in Table 1. The patients were referred by a certified specialist in either clinical endocrinology or oncology because of unilateral (n = 19) or bilateral (n = 2, patients 19 and 21) adrenal masses incidentally discovered during an abdominal imaging procedure (ultrasound or CT). The axial diameter of the mass was required to be at least 2 cm in one direction, and all patients with clear evidence of hormonally active tumor were excluded. Patient 7 had previously undergone a left adrenalectomy because of hormonally silent adenoma, and 6 patients had a history of malignancy without evidence of recurrence or metastatic disease.

All patients whose incidentaloma was found at ultrasound also underwent abdominal CT, which served as a morphologic gold standard for measurement of lesions. The median diameter at CT was 2.5 cm (range, 2–10 cm). Before imaging, the patients underwent hormonal evaluation including serum cortisol, plasma adrenocorticotropic hormone, plasma renin, serum aldosterone, serum sodium and potassium, serum dehydroepiandrosterone sulfate, nocorticotropic hormone, plasma renin, serum aldosterone, serum sodium and potassium, serum dehydroepiandrosterone sulfate, normetanephrine, and 24-h urine metanephrine and normetanephrine. An overnight oral 1-mg dexamethasone suppression test was performed to assess adrenocortical hormonal activity in all but 3 patients (patients 8, 14, and 19). This study was part of a larger multicenter trial performed under European Cooperation in the Field of Scientific and Technical Research (COST) action B12.

### Synthesis of Radiopharmaceuticals


11C-Metomidate was prepared by a modification of the published procedure (9). 11C-Methyl triflate, prepared by a standard procedure (11), was trapped at 0°C in 100 mL of acetone containing 0.5–1.0 mg of O-desmethyl precursor (R028141; Janssen Research Foundation) and 2.0–4.0 mL of freshly prepared tetrabutylammonium hydroxide (1 mol/L, aqueous). When trapping was completed, the reaction mixture was purified on a μBondapak C-18 column (125Å, 10 μm, 7.8 × 300 mm in internal diameter; Waters) using an eluent of methanol:water, 60:40, and a flow of 4 mL/min. The purified product was collected in a vessel containing 0.8 mL of sterile propylene glycol:ethanol, 7:3, evaporated, redissolved in sterile phosphate buffer (0.1 mol/L, pH 7.4), and filtered through a Gelman Acrodisc sterile filter (PN 4192, 0.2 mm; Pall Corp.). The final radiochemical yield of 11C-metomidate from 11C-methyl triflate was 40%–80%, with a specific radioactivity of 60 ± 20 Bq/μmol. The radiochemical purity was higher than 98% and stable for at least 2 h. Omission of propylene glycol and ethanol during evaporation decreased radiochemical purity (80%–90%), showing that the radiochemical impurities were formed by radiolysis.

### TABLE 1

Patient Characteristics, Final Diagnosis, and PET Findings Expressed as SUV of the Adrenal Mass

<table>
<thead>
<tr>
<th>Patient no.</th>
<th>Sex (y)</th>
<th>Age</th>
<th>Previous malignancy</th>
<th>Oral dexamethasone test</th>
<th>Mass diameter (cm) on CT</th>
<th>Final diagnosis</th>
<th>Operation</th>
<th>11C-MTO</th>
<th>18F-FDG</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>F</td>
<td>57</td>
<td>Uterine cancer</td>
<td>Positive</td>
<td>2.5</td>
<td>Adenoma, active</td>
<td>Yes</td>
<td>20.9</td>
<td>2.2</td>
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<tr>
<td>2</td>
<td>F</td>
<td>59</td>
<td>None</td>
<td>Positive</td>
<td>3</td>
<td>Adenoma, active</td>
<td>Yes</td>
<td>11.9</td>
<td>3.9</td>
</tr>
<tr>
<td>3</td>
<td>F</td>
<td>74</td>
<td>Laryngeal cancer</td>
<td>Positive</td>
<td>2.5</td>
<td>Adenoma, active</td>
<td>No*</td>
<td>26.1</td>
<td>2.5</td>
</tr>
<tr>
<td>4</td>
<td>M</td>
<td>67</td>
<td>None</td>
<td>Positive</td>
<td>2.6</td>
<td>Adenoma, active</td>
<td>No*</td>
<td>12.7</td>
<td>1.8</td>
</tr>
<tr>
<td>5</td>
<td>F</td>
<td>68</td>
<td>None</td>
<td>Positive</td>
<td>4</td>
<td>Adenoma, active</td>
<td>No*</td>
<td>9.1</td>
<td>1.8</td>
</tr>
<tr>
<td>6</td>
<td>F</td>
<td>48</td>
<td>None</td>
<td>Positive</td>
<td>2</td>
<td>Adenoma, active</td>
<td>Yes</td>
<td>16.0</td>
<td>3.4</td>
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<tr>
<td>7</td>
<td>F</td>
<td>74</td>
<td>Colorectal cancer</td>
<td>Positive</td>
<td>2.5</td>
<td>Adenoma, active</td>
<td>No*</td>
<td>10.2</td>
<td>2.3</td>
</tr>
<tr>
<td>8</td>
<td>F</td>
<td>59</td>
<td>None</td>
<td>Not done</td>
<td>2</td>
<td>Adenoma, inactive</td>
<td>No</td>
<td>12.2</td>
<td>1.6</td>
</tr>
<tr>
<td>9</td>
<td>M</td>
<td>66</td>
<td>Colorectal cancer</td>
<td>Negative</td>
<td>2</td>
<td>Adenoma, inactive</td>
<td>No*</td>
<td>10.0</td>
<td>1.7</td>
</tr>
<tr>
<td>10</td>
<td>M</td>
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<td>3</td>
<td>Adenoma, inactive</td>
<td>Yes</td>
<td>25.4</td>
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</tr>
<tr>
<td>11</td>
<td>M</td>
<td>54</td>
<td>None</td>
<td>Negative</td>
<td>2.5</td>
<td>Adenoma, inactive</td>
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<td>16.2</td>
<td>1.7</td>
</tr>
<tr>
<td>12</td>
<td>M</td>
<td>63</td>
<td>None</td>
<td>Negative</td>
<td>2</td>
<td>Adenoma, inactive</td>
<td>No</td>
<td>7.8</td>
<td>Not done</td>
</tr>
<tr>
<td>13</td>
<td>F</td>
<td>34</td>
<td>None</td>
<td>Negative</td>
<td>4.8</td>
<td>Adrenocortical cancer</td>
<td>Yes</td>
<td>28.0</td>
<td>2.9</td>
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<td>14</td>
<td>F</td>
<td>43</td>
<td>None</td>
<td>Not done</td>
<td>2</td>
<td>Normal adrenal</td>
<td>No</td>
<td>11.3</td>
<td>1.7</td>
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<tr>
<td>15</td>
<td>F</td>
<td>21</td>
<td>None</td>
<td>Negative</td>
<td>7</td>
<td>Cyst</td>
<td>No*</td>
<td>1.2</td>
<td>0.5</td>
</tr>
<tr>
<td>16</td>
<td>F</td>
<td>55</td>
<td>None</td>
<td>Negative</td>
<td>2</td>
<td>Lipoma</td>
<td>No§</td>
<td>13.3</td>
<td>2.2</td>
</tr>
<tr>
<td>17</td>
<td>F</td>
<td>41</td>
<td>None</td>
<td>Negative</td>
<td>2.5</td>
<td>Pheochromocytoma</td>
<td>Yes</td>
<td>10.5</td>
<td>3.0</td>
</tr>
<tr>
<td>18</td>
<td>F</td>
<td>67</td>
<td>None</td>
<td>Negative</td>
<td>4</td>
<td>Pheochromocytoma</td>
<td>Yes</td>
<td>7.8</td>
<td>2.9</td>
</tr>
<tr>
<td>19</td>
<td>M</td>
<td>69</td>
<td>Renal cell cancer</td>
<td>Not done</td>
<td>3</td>
<td>Metastasis</td>
<td>No*</td>
<td>2.0</td>
<td>1.7</td>
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<tr>
<td>20</td>
<td>M</td>
<td>71</td>
<td>Prostate cancer</td>
<td>Negative</td>
<td>6</td>
<td>Hepatocellular cancer</td>
<td>Yes</td>
<td>6.2</td>
<td>8.3</td>
</tr>
<tr>
<td>21</td>
<td>F</td>
<td>79</td>
<td>None</td>
<td>Positive</td>
<td>10</td>
<td>Lymphoma</td>
<td>No*</td>
<td>2.3</td>
<td>16.4</td>
</tr>
</tbody>
</table>

*Operation withheld because of intercurrent morbidity.

†Core biopsy without adrenalectomy.

‡Fine-needle biopsy.

§CT findings consistent with lipoma.

MTO = metomidate.
The synthesis of $^{18}$F-FDG followed a previously described nucleophilic substitution procedure (12). The radiochemical purity of $^{18}$F-FDG was at least 95%, and the specific radioactivity at the end of synthesis was more than 75 GBq/μmol.

**Patient Preparation**

$^{11}$C-Metomidate and $^{18}$F-FDG PET studies were performed either on the same day or within 1 wk of each other. For both PET studies, the subjects fasted at least 6 h or overnight. Drinking of water was allowed ad libitum. Before $^{11}$C-metomidate PET a 2-wk break was required from drugs that could affect uptake of tracer (ketocamazole, metyrapone). For the $^{11}$C-metomidate study, 2 catheters were inserted into contralateral forearm veins to inject tracer and to draw blood samples during imaging. The blood-sampling arm was wrapped in an electric blanket to heat the arm for arterIALIZATION of blood samples. The subject was placed supine in the PET scanner with the arms held upright, and correct positioning to image the adrenals was secured with anatomic landmarks and CT images. For repositioning during the second PET study, laser lines were marked on the skin with a felt-tip pen. At least 3 h separated injection of the 2 radiopharmaceuticals, with $^{11}$C-metomidate always preceding $^{18}$F-FDG. Two patients (patients 10 and 12) did not undergo $^{18}$F-FDG imaging for logistic reasons.

**Image Acquisition and Processing**

An 18-ring 2-dimensional whole-body PET scanner (Advance; General Electric Medical Systems) operated in 2-dimensional mode was used. The camera has bifhusth germate detectors, which image 35 planes at 4.25-mm intervals in a single session. The diameter of the field of view is 55 cm, and the axial length is 15.2 cm. Both PET studies were corrected for photon attenuation with 10- to 12-min preinjection transmission scans using robotically operated rods containing $^{68}$Ge/$^{68}$Ga. For cases in which $^{18}$F-FDG imaging was performed in whole-body mode, 2-min postinjection transmission scans were obtained.

Dynamic imaging was started by bolus intravenous injection of a median dose of 432 MBq (range, 211–444 MBq) of $^{11}$C-metomidate, and a 45-min emission scan was subsequently acquired (frame rate of 5 × 60 s, 5 × 180 s, 3 × 300 s, and 1 × 600 s). Arterialized venous blood was frequently sampled for measurement of radioactivity throughout imaging and determination of metabolites of $^{11}$C-metomidate. Samples for the latter were obtained at 2, 6, 10, 20, 30, and 40 min after injection of tracer. $^{18}$F-FDG imaging (n = 11) was performed in steady state starting 45 min after intravenous injection and consisted of 2-10 min frames. Alternatively, whole-body scans (n = 8) were obtained in the craniocephal direction with 6–7 bed positions, each lasting 5 min. No blood samples were taken during $^{18}$F-FDG imaging except for plasma glucose before tracer injection. The median dose of $^{18}$F-FDG was 369 MBq (range, 251–378 MBq). All image acquisition data were corrected for dead time, decay, and photon attenuation and reconstructed with an ordered-subsets expectation maximization algorithm using 4 iterations. The final in-plane spatial resolution in reconstructed images was 5 mm, and the axial resolution was 6 mm.

**Blood Metabolite Analysis**

Venous blood samples were collected into heparinized tubes at 2, 6, 10, 20, 30, and 40 min after injection to measure the amount of unchanged $^{11}$C-metomidate and radioactive metabolites in plasma. Plasma proteins were precipitated with acetonitrile, and the supernatant obtained after centrifugation was analyzed with high-performance liquid chromatography (HPLC) using a $\mu$Bondapak C-18 reversed-phase column (125A, 10 μm, 7.8 × 300 mm in internal diameter; Waters) at a flow rate of 6.0 mL/min and a gradient of 100% acetonitrile (B) in 50 mmol of phosphoric acid per liter (A) as follows: 0 min of 75% A and 25% B, 5.5 min of 40% A and 60% B, 7.5 min of 40% A and 60% B, and 8.5–9 min of 75% A and 25% B. A LaChrom HPLC system (Hitachi/ Merck) and a Radiomatic 150TR radioactivity detector (Packard) were used.

**Measurement of $^{11}$C-Metomidate and $^{18}$F-FDG Uptake**

The adrenals were invariably seen as hot spots in the final $^{11}$C-metomidate PET frame, which was used for defining regions of interest (ROI). Trace ROI function was used to outline ROIs encompassing the whole hot spot area in normal and enlarged adrenal glands, and mean (not maximum) radioactivity in the 3 consecutive axial planes with the highest radioactivity was used for further calculations. Similarly, a standard-sized circular ROI approximately 3 cm in diameter was drawn in 3 consecutive planes in the right lobe of the liver, also an organ with high uptake of $^{11}$C-metomidate (10). In the kinetic analysis, the input function for uptake of $^{11}$C-metomidate was corrected for 2 major labeled metabolites that constituted a variable but highly significant fraction of total plasma radioactivity. The corrected plasma and tissue time–activity curves derived from ROIs as explained above were used to calculate kinetic influx constant ($K_i$) values by applying the graphical analysis first described by Patlak (13). Standardized uptake values (SUVs) were also defined from the last frame, between 35 and 45 min, correcting tissue radioactivity for patient weight and injected dose (14).

Because normal adrenals have faint uptake of $^{18}$F-FDG, definition of ROIs was much more difficult on $^{18}$F-FDG images than on $^{11}$C-metomidate PET images. Accordingly, we outlined ROIs onto $^{18}$F-FDG planes by reading $^{18}$F-FDG and $^{11}$C-metomidate axial images together and by relating findings to those on corresponding CT scans. This was facilitated by the normal $^{18}$F-FDG uptake seen in liver, spleen, kidneys, and, variably, in stomach and bowel. The second of the 2 time frames (55–65 min) was used for calculation of SUV in adrenals and liver, again using the mean value over 3 consecutive planes. For whole-body $^{18}$F-FDG studies, only one 5-min frame was available for SUV calculation.

**Statistical Analysis**

Commercially obtained software (SPSS, release 11.0.1, standard version; SPSS Inc.) for Windows (Microsoft) was used for statistical evaluation. Results are expressed mostly as median and range. Normality of quantitative data was assessed with the Kolmogorov–Smirnov test. For normally distributed data, ANOVA and the independent-samples $t$ test was used; for other data, non-parametric methods (the Kruskal–Wallis test and Mann–Whitney $U$ test) were applied. The association between $K_i$ and SUV was evaluated with the Pearson correlation coefficient. All $P$ values are 2 tailed.

**RESULTS**

**Final Diagnosis of Incidentalomas**

The final diagnoses of 21 patients included 7 active adenomas as evidenced by a deficient dexamethasone sup-
pression test, 5 inactive adenomas, 1 adrenocortical carcinoma, 2 pheochromocytomas, 1 cyst, 1 lipoma, 1 normal adrenal falsely interpreted as tumor, and 3 noncortical malignancies (Table 1). Of the patients with active adenomas, 3 of 7 (43%) underwent adrenalectomy, whereas operation was withheld from the remaining 4 because of intercurrent disease (n = 3) or unexpected death due to coronary artery disease (n = 1). Diagnosis was confirmed histologically or cytologically for 13 patients and, for the remaining 8 patients, by characteristic radiologic appearance combined with concordant hormonal blood test findings and follow-up lasting a median of 14 mo (range, 9–21 mo). Patient 20 had well-differentiated hepatocellular carcinoma presenting as a pseudoadrenal mass (1).

Quantification of 11C-Metomidate Uptake

Identification of adrenocortical adenomas and normal adrenal glands was easy with 11C-metomidate PET because of the invariably high uptake of tracer in target organs with functional steroid hormone synthesis. In accordance with Bergström et al. (10), liver and, more variably, stomach and duodenum also showed moderate to high 11C-metomidate uptake, which did not interfere with radioactivity seen in adrenal tissues in late images. PET images of 4 patients with different types of adrenal incidentalomas are shown in Figure 1.

Two major 11C-metomidate polar metabolites with retention times of 2.3 and 3.2 min were seen in venous plasma with HPLC, and the rate of their appearance varied considerably between individual patients (Fig. 2). As a result, unchanged 11C-metomidate accounted for a median of 28% (range, 17%–40%; n = 12) of total radioactivity at 40 min from injection, and the respective figure at 20 min was 40% (range, 22%–54%; n = 12). Figure 3A shows the time course of radioactivity in uncorrected and corrected plasma samples and selected tissues in a representative patient with adrenocortical adenoma. The HPLC analysis made clear that only plasma corrected for metabolites could be used as input when applying the graphical method to evaluate the rate of 11C-metomidate uptake and metabolism in tissues. With the corrected input function, excellent fits were found for adrenal tumors and glands and for liver (Fig. 3B). A strong relationship between average SUV and graphical Kᵢ was seen for all tumor types and normal adrenal glands (r² = 0.73, P < 0.0001).

In quantitative analysis, the highest 11C-metomidate uptake among all tumors was seen in the single case of adrenocortical carcinoma, with an average SUV of 28.0 (Figs. 1E and 1F). This was followed in decreasing order of 11C-metomidate uptake, expressed as average SUV, by active adenomas (median SUV = 12.7; range, 9.1–26.1; n =

![Figure 1](image_url)

high 18F-FDG uptake of the adenoma in 1 d was misjudged as metastasis, with an average SUV of 3.9 (maximum, 6). Liver has invariably high uptake of 11C-metomidate, and tracer excreted in the urinary tract presents the highest radioactivity in 18F-FDG images.
7), inactive adenomas (median SUV = 12.2; range, 7.8–25.4; n = 5), normal adrenals (median SUV = 9.4; range, 4.6–13.8; n = 20), pheochromocytomas (SUVs = 7.8 and 10.5; n = 2), and noncortical malignancies (SUVs = 2.0, 2.3, and 6.2; n = 3). The solitary lipoma had an average SUV of 13.3, and in the cyst a very low tracer uptake was seen, with an average SUV of 1.2, the lowest value among all (Fig. 4). True uptake of 11C-metomidate into pheochromocytomas and lipoma may actually be lower, since the reported values are affected by the rim of displaced adrenal cortex surrounding the tumor (10).

The graphical Ki was highest for the adrenocortical carcinoma (1.14), followed by active adenomas (median Ki = 0.66; range, 0.33–0.78; n = 4), normal adrenal glands (median Ki = 0.37; range, 0.15–0.58; n = 12), and inactive adenomas (Kis = 0.24 and 0.37, n = 2). The median Ki for liver was 0.33 (range, 0.12–0.55; n = 11), and the respective SUV was 6.7 (range, 2.5–12.2; n = 21). A significant association between adenoma size and uptake of 11C-metomidate could not be demonstrated with SUV or Ki as an index of tissue 11C-metomidate metabolism.

**Comparison of 11C-Metomidate and 18F-FDG**

18F-FDG was not helpful in distinguishing adrenal tumors, with the exception of 2 of the 3 noncortical malignancies, which had characteristically high glycolytic activity coupled with low uptake of 11C-metomidate (Figs. 1G and 1H). Of interest was the relatively low uptake of 18F-FDG in adrenocortical carcinoma (average SUV = 2.9 and maximum SUV = 3.8; Fig. 1F) and in bilateral adrenal metastases from renal cell cancer, which had low maximum SUVs of 2.4 and 2.3. The primary tumor in the right kidney had been previously embolized and was difficult to see in the vicinity of high 18F-FDG activity excreted in the pelvis. Although 18F-FDG was believed to generally be less informative than 11C-metomidate, active adenomas (median SUV = 2.3; range, 1.8–3.9; n = 7) and the 2 pheochromocytomas (average SUVs = 3.0 and 2.9) showed higher uptake of 18F-FDG than did the 3 inactive adenomas, with SUVs of 1.6, 1.7, and 1.7 (Figs. 1B and 1D). Again, no association between tracer uptake and mass size could be found.

To assess the relationship between adenoma metabolism and hormonal activity, ratios of tumor to normal adrenal were defined for 11C-metomidate and ratios of tumor to liver, for 18F-FDG. The 11C-metomidate ratio could distinguish both active and inactive adrenocortical adenomas from other tumors (active vs. others, P = 0.003; inactive vs. others, P = 0.019) but was not different between hormon-
ally active and inactive adenomas (Fig. 5). The $^{18}$F-FDG ratio in noncortical tumors ($n = 7$) was not different from that in any other adrenal masses, including active and inactive adenomas ($n = 12$). This lack of significance depended strongly on the low number of malignant lesions and unexpectedly low uptake of $^{18}$F-FDG in 2 of these 4 neoplasias.

**DISCUSSION**

Three major issues emerging from the discovery of adrenal incidentaloma are disclosure of malignancy, characterization of tissue type, and characterization of hormonal secretory activity. It is widely accepted that patients having abnormal hormonal screening results or adrenal masses greater than 3–4 cm in diameter should undergo surgery whereas patients with nonfunctioning tumors that show typical radiologic features suggesting benignity are candidates for follow-up and repeated imaging studies (1,2). In patients with benign normosecretory tumors, adrenalectomy may be associated with unjustified morbidity, and patients with adrenal metastasis, in turn, require systemic rather than localized treatment. Nuclear scintigraphy aids in proper management of incidentalomas by helping with discrimination between lesions derived from adrenal cortex and lesions derived from adrenal medulla (8), whereas $^{18}$F-FDG PET has an established role in the diagnostic work-up of patients likely to have metastatic disease in the adrenals and other organs (4,15). Until now, radiolabeled cholesterol derivatives have been the first choice for $\gamma$-imaging of the adrenal cortex, whereas $^{123}$I- or $^{131}$I-labeled metaiodobenzylguanidine, a norepinephrine analog, is a common tracer for depicting tumors arising from the adrenal medulla (8). Because PET is now increasingly available in the clinical setting, pheochromocytomas may be diagnosed with $^{18}$F-FDG (16) and $^{18}$F-fluorodopa (17), and inhibitors of steroid biosynthesis, such as $^{11}$C-metomidate, may become counterparts of $^{18}$F-FDG and $^{18}$F-fluorodopa for PET imaging of the adrenal cortex (9,10).

The current study was performed under European COST action B12, aiming to develop a noninvasive test for evaluation of adrenal incidentalomas with PET. We looked specifically at the uptake kinetics of $^{11}$C-metomidate in tumors and normal adrenal glands and in liver and venous
plasma and compared the findings with those on 18F-FDG images obtained in steady state. In line with the preclinical validation study (9) and the first clinical study (10), we could demonstrate high uptake of 11C-metomidate in adrenocortical tumors and normal adrenal glands as a sign of sustained 11β-hydroxylase activity. Decreased tracer uptake, in comparison with physiologic uptake, was seen in pheochromocytomas, cysts, and noncortical malignancies such as lymphoma and metastatic carcinomas, whereas adrenocortical carcinoma had the highest uptake of 11C-metomidate among all lesions. 18F-FDG was clearly inferior to 11C-metomidate for depicting adrenal lesions if they were adenomas and, somewhat surprisingly, also in the solitary case of adrenocortical carcinoma. The low uptake of 18F-FDG in adrenal metastases from renal cell cancer was less surprising because 18F-FDG is known to have only a limited role in the diagnosis of renal cell carcinoma (18).

The uptake kinetics of 11C-metomidate in adrenal tissue are favorable for PET imaging because irreversible binding of the tracer occurs over the first 45 min (Fig. 3). Because 2 large fractions of 11C-metomidate metabolites occur in plasma, it is necessary to correct the plasma input function for these yet unidentified radiochemical species before applying graphical analysis to assess binding of 11C-metomidate. This correction was successfully performed for 12 patients to calculate graphical influx constant K(i, Fig. 3), and in further analysis we showed that the obtained Ki's could be replaced by the robust SUV approach commonly used in clinical oncology (4,14). K(i, and SUV were not associated with mass size, and pheochromocytomas were seen to have the highest uptake in the outer rather than the inner zone of tumor—a finding compatible with the steroid-synthesizing property's being the major determinant of uptake of 11C-metomidate in adrenal tissue (9). This does not translate, however, to a direct measure of rate of adrenocortical hormone synthesis, since the SUV's in active and inactive adenomas overlapped widely, in keeping with the findings of Bergströ¨m et al. (10). Furthermore, uptake ratios for adenoma to contralateral adrenal gland were not significantly different between active and nonsecretory adenomas (Fig. 5).

Although 11C-metomidate PET does not immediately replace 131I-labeled cholesterol derivatives in the functional evaluation of adrenal cortex, a search for new tracers applicable to positron imaging is in order with the widespread expansion of dedicated PET scanners. The major advantages of PET are rapid completion of the imaging, within 1 h, and a resolution and sensitivity better than those of adrenal scintigraphy. Clearly, the value of quantitative 11C-metomidate PET will remain obscure until the results from the larger European multicenter study are at hand. In our trial, most patients were seen by a clinical endocrinologist; we hope that on that basis the multicenter trial included more patients with metastatic tumors and thus avoided the slight referral bias of the current study. Unfortunately, it seems difficult to distinguish adrenocortical adenoma from carcinoma with the current technique, but for a patient with known carcinoma, 11C-metomidate should be a specific tracer for metastatic disease (10). For patients presenting with adrenal masses and a history or strong suggestion of neoplastic disease, 18F-FDG PET may still be the study of choice because whole-body imaging may conceal other deposits of cancer in addition to adrenal metastasis (15).

CONCLUSION

We conclude that the pattern of 11C-metomidate uptake facilitates discrimination between various types of adrenal masses, suggesting that 11C-metomidate PET may be useful for functional evaluation of incidentalomas. The radiologic features of the adrenal mass, a clinical history addressing the risk for malignant disease, and the hormonal profile of the patient should be considered when assessing the advantages of 11C-metomidate relative to those of other positron-emitting tracers such as 18F-FDG and scintigraphic techniques using cholesterol derivatives or 123I-metiodobenzylguanidine. Further to be emphasized is that 11C-metomidate PET, rather than representing a new modality to replace the established methods, remains complementary to CT and MRI in the diagnosis and management of incidentalomas.

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