Incidental findings in cancer patients: abdominal organs

RöKo INT 103

RöKo International Mittwoch, 24.05.2017 von 15:30 bis 16:50 Uhr im Raum: Donner

Beschreibung
in Kooperation mit ICIS

Zusatzinfo
in Kooperation mit ICIS

Vorsitz / Moderation: Diederich S / Düsseldorf, Schima W / Wien

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<th>Adrenal and retroperitoneum</th>
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<td>15:30 Uhr</td>
<td>Referent(en): Francis I</td>
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Kurzfassung: Introduction

Incidental adrenal masses are commonly seen in cross-sectional studies such as CT and MRI, which are being used with increasing frequency in daily practice in the last several years. Different studies have shown that the incidence ranges from 3-4%. The incidence rises in the older population and it has shown that in subjects with hypertension and diabetes that the incidence rises above 10%.

But in oncology patients, even though adenomas are common, metastases are also more frequent especially for example in subjects with lung cancer. Therefore, it is essential to differentiate between adenomas and metastases and other non-adenomatous masses.

Other benign masses

Other benign adrenal masses that are less commonly seen include adrenal cysts, myelolipomas and adrenal hemorrhage. These lesions at times have typical features on imaging that may permit their diagnosis. But at times for example cysts when complicated by hemorrhage or infection, may have imaging features that make it impossible to distinguish from cystic malignant tumors.

Adrenal adenomas and malignant masses

Differentiation using CT, MRI and PET

In the past decade there have been multiple studies that have shown with high accuracy that most adenomas can be distinguished from malignant lesions (mostly metastases) by density measurements (HU) on unenhanced CT and using adrenal CT washout features. Similarly using chemical shift MR imaging this differentiation between the entities has been made. These techniques have been also validated on a few studies distinguishing between adenomas and primary malignant adrenal masses i.e. adrenal cortical carcinomas and pheochromocytomas. PET imaging has also been shown to capable of differentiating benign from malignant masses with a high degree of accuracy.

Pitfalls

At times metastases from lipid-containing primary malignancies such as clear cell renal carcinoma, and well differentiated HCC, as well as primary adrenal malignancies such as ACC and pheochromocytomas can demonstrate findings on density measurements, washout and CSI-MRI that are similar to adenomas. In a small percentage of cases, adenomas can demonstrate increased tracer uptake mimicking malignant tumors.

Adrenal biopsy

The role for adrenal biopsy has diminished over the last decade being currently only rarely used to confirm diagnosis of a solitary site of metastasis and in rare instances where imaging is indeterminate.

**Lernziele:**
1. Wissen über die Prävalenz von incidentalen fokalen Leberläsionen (FLL).
2. Lernen, wie man FLL zuverlässig charakterisieren kann.
**Kurzfassung:** Pancreatic incidentalomas (PI) are discovered with increasing frequency as the use of high quality cross sectional imaging is becoming more widespread. They cover a wide spectrum of pathology from benign simple cysts through potentially malignant lesions such as intraductal papillary mucinous neoplasia, to frankly malignant adenocarcinoma. The incidence of malignancy in PI is lower than in symptomatic pancreatic lesions (31% vs 76%), however, malignant histology is found up to 30 % and pre-malignant up to 50% in PI in surgical series. A simple classification of PI subdivides lesions in SOLID LESIONS (NON ENHANCING or ISOENHANCING; HYPOENHANCING; HYPERENHANCING) and CYSTIC LESIONS (NON COMUNICATING or COMUNICATING with main pancreatic duct). Fatty infiltration, pancreatic lipoma, intrapancreatic spleen, pancreatic anomalies can give a solid appearance of the pancreas. However, all different solid tumors of the pancreas can be incidentally discovered. Among CYSTIC LESIONS of the pancreas, frequency of detection of CPL ranges from 2.4% to 19.6%. Pseudocysts are usually present in Pts with previous Hx of pancreatitic. Serous cystic neoplasm (SCN), mucinous cystic neoplasm (MCN), solid pseudopapillary tumor (SPT) can all have cystic appearance, the latter being more frequent in young patients: MCNs occur almost exclusively in the body-tail of the gland, whereas SCNs have no site predilection; cystic lesions in males and those in the head of the pancreas are unlikely to be MCN. A multicystic pattern of pancreatic lesions is more frequently observed in SCN, whereas an olio- and/or macrocystic pattern is more frequently observed in MCN. Serous oligocystic adenoma has a multicystic or lobulated contour with or without septation, whereas mucinous cystadenoma has a smooth contour with or without septation. Finally IPMN are cystic lesions communicating with main pancreatic duct, whose malignant potential is related to the involvement of the main pancreatic duct, size of the cyst and presence of solid component inside the lesions.

**Lernziele:**
- to learn about the incidence of incidental pancreatic lesions
- to illustrate the different imaging findings of pancreatic incidentalomas
- to discuss about the best management of PI

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**RöKo INT 103.4**  
**Female pelvis**

16:30 Uhr  
Referent(en): Rockall A

**Kurzfassung:** Pelvic masses are relatively common in women and the vast majority are benign in nature. These can include benign uterine leiomyoma, physiological ovarian cysts or benign ovarian lesions which may be cystic, such as cystadenoma/adenofibroma or solid, such as fibroma/fibrothecoma. When a pelvic mass is detected incidentally on imaging for cancer staging of a non-gynae cancer, comparison with any previous imaging can be very helpful in confirming whether a mass is longstanding, such as a leiomyoma. However, where there is concern for possible significant pelvic disease, such as an ovarian metastasis, it is important to undertake appropriate investigations to confirm the diagnosis. Ultrasound is the initial investigation and where this is indeterminate, MRI may be used for problem-solving.