A Review of Scientific Topics and Literature in Abdominal Radiology in Germany – Part 2: Abdominal Parenchymal Organs

Zusammenfassung


Key Points:
▶ Different subtypes of liver adenomas with different therapeutic consequences can be differentiated by MRI
▶ Most recently published studies focus on liver imaging with extracellular liver specific contrast media as well as diffusion weighted imaging. They consider this new method having a high diagnostic potential.
▶ For pancreatic neoplasm diagnosis diffusion – as well as perfusion – imaging is considered as a highly promising method.

Abstract

The working group for abdominal and gastrointestinal diagnosis is a group of the German Radiological Society (DRG) focusing clinically and scientifically on the diagnosis and treatment of the gastrointestinal tract as well as the parenchymal abdominal organs. In this article we give an up-to-date literature review of scientific radiological topics especially covered by German radiologists. The working group experts cover the most recent relevant studies concerning liver-specific contrast media with emphasis on a new classification system for liver adenomas. Additionally studies regarding selective internal radiotherapy (SIRT) are reviewed. For the pancreas the most important tumors are described followed by an introduction to the most recently introduced functional imaging techniques. The manuscript concludes with some remarks on recent studies and concerning chronic pancreatitis as well as autoimmune pancreatitis.

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▶ Most recently published studies focus on liver imaging with extracellular liver specific contrast media as well as diffusion weighted imaging. They consider this new method having a high diagnostic potential.
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Citation Format:
Introduction

In part one of our literature review regarding the gastrointestinal tract we already made the expertise of the members of the working group for abdominal and gastrointestinal diagnosis of the German Radiological Society available to all members of the DRG. The working group is a representative of German abdominal radiology and a comprehensive platform for interested researchers and clinicians with a focus on abdominal diagnosis and therapy. This second part includes an overview of the current literature with a focus on the developments and trends related to the abdominal parenchymal organs such as the liver and the pancreas.

The use of liver-specific contrast agents with uptake in the hepatocytes via an organic anion-transporting polypeptide (OATP) has resulted in a significant improvement in differential diagnosis. After the original euphoria regarding the exact differentiation of the hepatocarcinogenesis, a certain amount of sobering has occurred primarily as a result of several Japanese studies. A liver-specific contrast agent seems to be helpful for the subtyping of liver cell adenomas and their differentiation from FNH (focal nodular hyperplasia). The current status for the subtyping of liver cell adenomas is therefore briefly discussed in this overview. In addition, the value of SIRT (selective internal radiotherapy) is presented on the basis of the current literature.

The second part of this overview addresses the pancreas. In addition to the high mortality rate compared to other tumor entities, the pancreas has increasingly become a topic of interest in recent years in the literature in Germany are largely from members of the working group for abdominal and gastrointestinal diagnosis or were conducted with their active participation.

The value of the liver-specific contrast agent Gd-EOB-DTPA (Primovist) regarding the diagnosis of focal liver lesions was further examined [1–4]. A large study proved, that the uptake of Gd-EOB-DTPA in the hepatobiliary contrast phase is the best criterion in the differentiation between FNH and hepatocellular adenoma and this criterion is superior to all morphological and dynamic criteria in the differentiation of the two lesions [1]. The nomenclature of adenomatous lesions was constantly changing in the past. In 2010a new standard was presented by a French working group in a newly published classification system [5]. There are currently 3 adenoma subtypes with varying frequencies (see parentheses) and an unclassified subgroup to which various tendencies for malignant change and thus divergent treatment approaches can be assigned:

- HCA with HNF1α mutation: No tendency for malignant change (35 %)
- HCA with β-catenin mutation: High tendency for malignant change (10 %)
- Inflammatory HCA (IHCA) (45 %)
- Unclassified HCA (10 %)

In a recent study presented at ESGAR 2014 including a total of 39 systematically analyzed and histologically verified liver cell adenomas, characteristics for the subtyping of liver cell adenomas using Gd-EOB-DTPA were developed [6]; All adenomas showed a hypointense signal behavior in the hepatobiliary phase after 10 min p. i. in contrast to the regular hyperintensity of FNH in this phase. Under consideration of the qualitative and quantitative results of this study, the unenhanced T1 sequence and the venous contrast phase seem to be diagnostically valid for the differentiation between inflammatory HCA and HNF1α adenoma.
Intra-arterial treatment of liver tumors

Primary and secondary liver tumors are often part of diagnostic as well as therapeutic radiology in the clinical routine. Depending on the extent of the tumor invasion, different treatment concepts can be used. In addition to surgical and medication-based treatments, minimally invasive local or locoregional methods, such as radiofrequency ablation (RFA) or transarterial chemoembolization (TACE), are often used. A newer form of endovascular treatment is selective internal radiotherapy (SIRT). This is a transarterial, catheter-based method in which the beta-emitter yttrium-90 is introduced into the liver. The radionuclide is bound to glass or resin microspheres. As a result of increased accumulation in particular in hypervascularized liver tumors, a focused radiotherapeutic effect is achieved.

Although SIRT is similar to TACE in many respects, precautionary safety measures must be taken to avoid a transfer of yttrium to extrhepatic tissue. In recent years the handling of “critical” vessels has been a topic of research. Such critical vessels include for example the gastroduodenal artery, the cystic artery, and the falciform artery, which usually arise from the vascular bed of the hepatic artery and provide the arterial supply to the GI tract, the gallbladder and the abdominal wall, respectively. For example, the gastroduodenal artery is routinely closed prior to SIRT in many centers. In a study by Schelhorn et al. [13], 86 patients in whom coil embolization of the gastroduodenal artery was performed were followed up. It was able to be shown that new collaterals into the flow region of the gastroduodenal artery that had to be closed prior to yttrium administration formed in 28 of 86 patients after vascular occlusion. However, these collaterals could not be probed and occluded in 3 patients so that these patients could not undergo SIRT. The authors concluded that the gastroduodenal artery should not be occluded and instead segmental or lobar SIRT should be performed with the microspheres being applied distal to the origin of the gastroduodenal artery.

Another vessel that is a focus of SIRT is the cystic artery. If yttrium particles spread to the gallbladder, radiation-induced cholecystitis can occur. Theysohn et al. [14] reported on a series of 20 patients in whom special measures were necessary prior to yttrium therapy in order to prevent or reduce the spread of yttrium to the gallbladder wall. The cystic artery was occluded temporarily with gel particles or permanently with coils. Ischemia of the gallbladder wall did not occur in the patients. However, it was able to be ensured by these measures that no microspheres entered the gallbladder bed.

The falciform artery is categorized as problematic less frequently. Since the falciform artery often originates far in the periphery of the hepatic artery, selective occlusion is often not possible. Schelhorn et al. examined 19 SIRT patients with an open falciform artery [15]. It was able to be shown that a large-caliber, open falciform artery is not a contraindication for SIRT. The authors recommended placing cold packs on the abdominal wall during yttrium administration in cases in which the falciform artery cannot be occluded. This reduces the arterial blood flow into this area and the perfusion with yttrium is minimized.

In addition to the rather technical studies mentioned above, a further focus of scientific interest is the question as to whether it is useful to combine SIRT with other therapeutic methods. A multicenter study (SIRFLOX study) examines patients with liver metastases in colorectal cancer [16]. The extent to which SIRT in addition to chemotherapy with FOLFIRI extends the progression-free survival of patients is examined. The SORAMIC study has a
In the case of early diagnosis, tumor resection seems possible. However, approximately 70% of patients have an inoperable tumor at the time of diagnosis. Under clinical conditions, a positive effect of sorafenib prior to SIRT could be described [18]. Thaysen et al. examined patients in whom SIRT was initially not possible due to a high hepatopulmonary shunt fraction. Sorafenib therapy resulted in a significant reduction of the pulmonary shunt so that SIRT could consequently be performed in select patients.

**Pancreas**

**Solid pancreatic tumors – ductal adenocarcinoma**

The most common solid tumor of the pancreas is the ductal adenocarcinoma. Approximately 15,000 patients are diagnosed each year in Germany with this tumor and the annual mortality rate for this entity is similarly high. Despite the advances in diagnosis and treatment, the prognosis remains unfavorable. 80–90% of patients have an inoperable tumor at the time of diagnosis. The recommendation of the S3 guidelines updated in 2013 regarding imaging methods for primary diagnosis is: “The first-choice diagnostic methods for detecting pancreatic cancer are upper abdominal ultrasound, endoscopic ultrasound, MD-CT examination, and MRI in combination with MRCP” [19]. A definitive recommendation for or against one of the methods is not made. The method with which the examiner has the greatest experience should be used. Imaging in adenocarcinoma of the pancreas has 3 main goals:

- Detection or exclusion of a tumor
- Evaluation of the resectability of a carcinoma
- Differentiation of ductal adenocarcinoma from other tumor types or from chronic pancreatitis

In the case of early diagnosis, tumor resection seems possible. This means a possible improvement of the survival rate for a select group of patients [20, 21]. Since these pancreatic tumors have a ductal origin, an obstruction and duct dilatation are the earliest signs of a tumor. Tumors with a size of less than 1 cm can be detected solely on the basis of duct dilatation in approximately 60% of cases [22]. Approximately 70% of ductal adenocarcinomas are located in the head of the pancreas which is why obstruction of the distal bile duct (double duct sign) occurs relatively frequently and early [23]. After contrast administration, pancreatic cancer is typically less perfused than the surrounding tissue on CT and MRI. Around the tumor the pancreas loses the typical lobulated contour. In addition, tumors show direct infiltration into the surrounding retroperitoneal fatty tissue very early. Pancreatic cancer spreads quickly lymphatically and perineurally into the retroperitoneal fatty tissue and grows around the mesenteric vessels and the celiac trunk. After the diagnosis of pancreatic cancer, the assessment of resectability or irresectability is the greatest challenge for imaging. Irresectability can be determined more reliably. The S3 guidelines specify the following recommendation regarding preoperative diagnosis: “MD-CT and endoscopic ultrasound are recommended for the preoperative evaluation of local tumor extension and for evaluating resectability. Abdominal ultrasound is mandatory for evaluating systemic tumor spread. Abdominal MD-CT is mandatory when systemic formation of metastases cannot be detected on abdominal ultrasound or when imaging according to RECIST criteria (“response evaluation criteria in solid tumors”) is desired in study situations [19].

The following irresectability criteria are defined:
- Extensive infiltration of the celiac trunk and/or of the superior mesenteric artery
- Distant metastases

While every vascular contact of the tumor was previously evaluated as a sign of infiltration, surrounding of the superior mesenteric artery or the celiac trunk of more than 180° is now considered a reliable sign of irresectability (Fig. 2) [24].

**Endocrine pancreatic tumors**

Rare endocrine pancreatic tumors include a heterogeneous group of diseases with different clinical pictures, different morphologies, and varying degrees of malignancy. The diagnosis of these tumors is typically based on the combination of characteristic clinical symptoms and lab findings, particularly in the case of hormonally active tumors. The goal of imaging is to locate tumors and detect metastases. The two most common tumors, insulinoma and gastrinoma, have an annual incidence of 0.3 – 3 per million people while the remaining tumor types occur much less frequently.

**Cystic pancreatic tumors**

Cystic pancreatic tumors are discovered using high-resolution imaging methods with increasing frequency and are responsible for the increase in surgical interventions involving the pancreas. The following tumors represent approximately 90% of pancreatic lesions with a cystic appearance [25, 26]:

- Intraductal papillary mucinous neoplasms (IPMN)
- Mucinous cystic neoplasms (previously referred to as mucinous cystadenomas)
- Serous cystic neoplasms (previously referred to as serous cystadenomas)
- Solid pseudopapillary tumors (Frantz’s tumors)
- Neuroendocrine cystic tumors
Cystic pancreatic tumors typically have a series of clinical and morphological features that allow characterization in most cases.

**Intraductal papillary mucinous neoplasms (IPMN)**
Histologically speaking, an IPMN is based on the intraductal epithelial proliferation of cells that are polypl-like or flat and are usually associated with an excessive formation of mucus. The excessive viscous mucus displaces the pancreatic ducts and causes them to dilate thus giving these tumors their cystic appearance. Strictly speaking, this is not a “cystic” tumor of the pancreas but only cystic dilatation of the pancreatic duct system. A communicating system of cystically dilated ducts that look like cysts on cross-sectional images but differ clearly from the grape or honeycomb-like appearance of serous cystic neoplasms typically form. The papillary tumors can seem like small polyps and can be identified starting at a size of approximately 3 mm, particularly after contrast administration.

IPMNs are categorized according to their location and type of growth as main-duct types, side-branch types, and mixed types. The mixed types seem to be neoplasms that arise from the main duct and spread secondarily into the side branches [27]. The side-branch type is usually located in the head of the pancreas and preferably in the uncinate process. A side-branch IPMN histologically corresponds to a gastric type while the main-duct IPMN is differentiated as a pancreato-biliary or intestinal type. The main-duct type has a significantly higher tendency for malignant change than the side-branch type. A differentiation is made between benign tumors, borderline types, and malignant tumors. A cancer in situ is found in up to 27% of cases and an invasive cancer in up to 40% of cases. Histologically speaking, invasive types are either mucinous or ductal carcinomas. The 5-year survival rate is 98–100% for noninvasive tumor types, 89% for minimally invasive cancers, and 58% for invasive cancers [28].

Depending on the type of spread, the following changes are characteristic on CT and MRI:

- **Cystic dilatation of the side branches**
- **Circumscribed or diffuse dilatation of the main duct**
- **Small intraductal papillary tumors**

In the case of the side-branch type, the detection of communication with the pancreatic duct system is essential for the delimitation from other tumors (Fig. 3). The introduction of thin-slice CT made it possible to show this connection in over 80% of cases. However, reconstruction methods must be used. This detection is possible in approximately 90% of cases with MRCP.

In the case of the main-duct type, segmental or diffuse dilatation of the pancreatic duct is pathogenic and is frequently associated with polyposy intraductal changes (Fig. 4).

In principle, it is possible to differentiate between benign and malignant types of IPMN [29]. Main-duct IPMNs must be fundamentally classified as potentially malignant. Patients with dilatation of the main duct and a corresponding physical condition should undergo surgery. In contrast, patients with side-branch IPMN without high-risk stigmata (obstructive jaundice in patients with a cystic lesion in the head of the pancreas, solid portion and/or diameter > 10 mm) can be monitored [30].

**Mucinous cystic neoplasm**
According to the new WHO classification, mucinous cystic neoplasms are characterized as follows: "Tumors without a connection to the pancreatic duct system that have a multicellular epithelium and often have an ovarian-like stroma" [31]. The histological detection of an ovarian-like stroma is because this tumor type is found almost exclusively in women. The age of onset is usually between the 40th and 50th year of life. Mucinous cystic tumors are typically solitary and are comprised of individual or a few large, thick-walled cysts that have a chambered appearance in some cases. Almost all tumors (>95%) are in the corpus and cauda of the pancreas. The cysts contain mucinous, hemorrhagic or necrotic material. Nodular changes in the wall and papillary projections can be indications of a malignant change [32]. Imaging is dependent on the tumor architecture and cyst contents. In particular, the detection of a hemorrhage is a strong indicator of this type of tumor. High signal intensity caused by bleeding and protein-rich mucus is seen on T1-weighted MRI sequences. The cyst contents are signal-intense on T2-weighted images and even thin cyst walls are sharply defined in the fluid. Cyst walls and septae visibly absorb contrast agent. Although calcifications can be better detected with CT, septations usually cannot be delimited as well as with MRI.

**Serous cystic neoplasm**
Serous cystic neoplasms include the following tumors:

- **Serous microcystic adenoma**
- **Serous oligocystic adenoma**
- **Cystic neoplasms in Von Hippel-Lindau syndrome**
- **Rare serous cystadenocarcinoma**

The most common subtype is microcystic adenoma comprised of numerous small cysts with a diameter of 1–5 mm. A central scarred connective tissue structure from which radial septae arise and that can contain calcifications is common. Serous oligocystic adenomas are slightly rarer and comprise approximately 7–10% of all serous cystic tumors. They are comprised of individual or a few larger cysts with a size of 2–25 mm. They have no central scar and no calcifications but have the same histological signs as serous microcystic adenomas. Cystic neoplasms in Von Hippel-Lindau syndrome are usually numerous and are distributed in a diffuse or multifocal manner over the entire pancreas [33].

The morphological visualization of these tumors depends on their architecture, e.g. the size of individual cysts, and the spatial resolution of the technique being used. If the cysts are very small and the septae are delicate, they can only be sufficiently visualized with very high resolution. In the case of insufficient resolution, these tumors can give the appearance of a solid aspect.

**Solid pseudopapillary pancreatic tumor**
This tumor, also referred to as “Frantz’s tumor” occurs almost exclusively in young women (average age 24–39 years) and is seen in children in individual cases. However, this mass is typically classified as benign or with low malignant potential. The prog-
nosis is favorable [34]. If these tumors are seen in older patients, the probability of a malignant change increases. The most common metastatic pathway in the case of a malignant change is hematogenic spread into the liver. Tumors can be well delimited on MRI and CT and have a very varied appearance depending on their architecture. Small tumors (diameter of <3 cm) are often solid, can be well delimited and show slow contrast enhancement after intravenous contrast administration [35, 36]. Only larger tumors show the characteristic presentation with solid and cystic areas with hemorrhagic areas standing out in MRI in particular. A thick capsule that shows contrast enhancement can typically be detected. Peripheral calcifications are described in up to 30% of tumors. This tumor is often misinterpreted as a (post-) traumatic pseudocyst.

New trends in pancreas diagnosis – functional imaging
In the special research area SFB/TR125 “cognition-guided surgery”, the Heidelberg working group examined new imaging methods such as dual-energy CT perfusion (known as DECT perfusion) and diffusion MRI (DWI-MRI) regarding their value for pancreas diagnosis in two SFB projects. Using DECT perfusion on color-coded perfusion maps typical hypodense pancreatic tumors can be reliably identified and small isodense tumors can appear as focal hypoperfused areas that can be determined quantitatively as well as qualitatively [37, 38]. In addition to the two-compartment model (Patlak model) for perfusion map calculation, it is important to also use motion correction algorithms [39]. The value of DECT perfusion for the identification of pancreatic tumor relapse in early postoperative follow-up is the topic of current research. Diffusion MRI is becoming increasingly important in oncology and thus in abdominal radiology. In addition to the liver, diffusion sequences are increasingly used for pancreas diagnosis. However, the sequence technique from the liver cannot be transferred on a 1:1 basis due to the different microcirculation structures of the vessels in the pancreas. Basically, two different models are used for diffusion imaging of the pancreas: the monoexponential method and the biexponential method. The former method is faster and more widely available and typically uses two b-values for tumor detection (b=0 and 1000) and four b-values for lesion detection (0, 100, 500, 1000) [40]. The resulting ADC maps correspond to the diffusion effect in the analyzed voxel. The less known and more time-intensive biexponential method combines tissue perfusion with the diffusion component, microvascular perfusion being visualized at low b-values in particular. Low b-values of less than 100 sec/mm2 are more sensitive for perfusion and microcirculation effects. However, measurements of the up to 11 b-values needed for the acquisition in addition to the longer acquisition time are significantly more complex and also require more complex post-processing. However, a differentiation between pancreatitis, carcinoma and fibrosis is possible when there is a certain overlap of both collective [41]. Therefore, this method is currently not used in the clinical routine and is only evaluated in studies. At 1.5 T, monoexponential models as mentioned above are therefore currently recommended even if the authors recommend consideration of the biexponential method for diffusion imaging of the pancreas based on data from larger study collectives [42]. The value of DWI-MRI for the follow-up of autoimmune pancreatitis is currently being examined.

**Chronic pancreatitis**
After a long preparation period, the first German S3 guidelines regarding the diagnosis and treatment of chronic pancreatitis were developed and published in 2012 under the supervision of the German Society of Digestive and Metabolic Diseases with participation of the German Radiological Society [43]. Basically, the guideline recommends the use of ultrasound for basic diagnosis in this complex and increasingly prevalent chronic disease [44]. According to current literature, MRI in combination with MRCP is the optimal method for differentiating pancreatic cancer from focal changes in chronic pancreatitis. With respect to study design, categorization according to the modified Cambridge classification is recommended [45].

**Autoimmune pancreatitis (AIP)**
Autoimmune pancreatitis is a rare special form of idiopathic chronic pancreatitis with pain, cholestatic jaundice, and hypergammaglobulinemia of all classes primarily of IgG4, described for the first time in 1995 [46]. Since the increase in immunoglobulins results in systemic inflammatory fibrosis, the pancreas is only an effector organ. Autoimmune pancreatitis responds well to cortisone and should therefore be definitively radiologically differentiated from the most important differential diagnosis, i.e., pancreatic cancer which requires a completely different treatment [43]. In addition to histological and serum changes for reliable diagnosis, there are some typical signs that can be effectively detected with radiology [47]:

a) The “sausage sign”, a generalized sausage-shaped thickening of the entire pancreas without significant peripancreatic fluid quantities that can be diagnosed equally with CT and MRI. Differentiation from acute edematous pancreatitis is not possible here.

b) An area that is hypointense on T1-weighted images compared to the rest of the parenchyma and is slightly hyperintense on T2-weighted images can be delimited in unenhanced MRI sequences in approximately 50% of cases of AIP. Hypointensity is seen in the arterial phase of contrast dynamics and a hyperintense signal known as “late enhancement” is seen in approx. 65% of cases in the venous phase and in the late venous phase in 74% of cases. This hyperintense signal in the venous and late venous phase indicates an AIP. Such a hyperdense behavior in the venous phase often cannot be delimited with CT. Therefore, MRI is superior to CT for the diagnosis of AIP.

Initial studies regarding the diagnosis and treatment course of AIP via DWI-MRI are the current topic of research. Cave: the two clinical types of AIP, lymphoplasmacytic sclerosing type 1 (LPSP) and idiopathic duct-centered type 2 (IDCP), cannot be radiologically differentiated from one another.

**Developments, summary and conclusion**

Concerning the current US cancer statistics, a dramatic change in the incidence and mortality in one of the fastest growing medical areas, namely oncology, is forecast by the year 2030. In addition to a doubling of the number of cases in oncology in the next 30 years, the number of cases of pancreatic cancer and hepato-biliary tumors in particular will increase significantly while the number of cases of colorectal tumors will slowly decrease and the number of cases of bronchial cancer and breast cancer will increase only slightly [48].

This has major effects on radiological diagnosis and interventional therapy, in particular in abdominal radiology. Publications regarding the use of liver-specific contrast agents for differentiating the hepatocarcinogenesis and the evaluation of the treatment success of new techniques such as SIRT are both of great interest because with an increasing number of oncological patients the number of treatments in general as well as the number of patients requiring palliative treatment will increase significantly and all involved disciplines will encounter capacity limits. Primarily systematic studies on SIRT therapy allowed optimization of the procedure. In addition to exact diagnosis determination, radiology could play an important role in an increasingly personalized medicine, namely for answering the following questions: which patient is suitable for which treatment, is this specific (and usually very expensive) treatment effective, and should the treatment be continued?

Therefore, the countless scientific activities in German literature and the continuous increase in the number of members in the working group for abdominal radiology of the German Radiological Society can be sufficiently explained.

The known paradigm changes in almost all new S3 guidelines with an increasing influence of cross-sectional imaging methods and interventional treatments are based on the good amount available data due to the high number of publications particularly in the German literature. In addition to the evaluation of whether a patient with newly diagnosed pancreatic cancer is suitable for a surgical treatment, cross-sectional imaging with the highest spatial resolution should also be used for the early diagnosis of a local relapse without distant metastases because repeat surgery yields a survival advantage of approximately 11 months [49]. The same is true for the constantly increasing incidental diagnosis of cystic pancreatic lesions, the correct diagnosis of type, the typing of a possible IPMN and the evaluation of the presence of a possible malignancy with a corresponding therapy procedure. Whether the increasing number of publications regarding functional oncological imaging will become established in the clinical routine is clarified by current publications regarding the differentiation of hypodense and isodense pancreatic tumors via DECT perfusion with color-coded perfusion maps regarding blood flow, blood volume, and permeability in the framework of German Research Foundation-supported special research areas. Whether a differentiation between pancreatitis and carcinoma via DWI-MRI is possible and what the value of diffusion MRI will be in the future are also topics of current research [32 – 35].

In summary, radiologists have become an established part of modern oncology in an interdisciplinary setting and today are much more than “image demonstrators” and will allow these regions to grow significantly according to the predicted increase in hepatobiliary tumors and pancreatic cancer with a special focus on abdominal radiologists and interventional colleagues by 2030.

Fig. 4 Malignant change of a main-duct IPMN a detection of a very large main-duct IPMN with purely cystic dilatation (arrow). b The same patient 3 years later: Clear, solid tumor portions (arrow) are signs of a malignant change.

Abb. 4 Maligene Entartung einer Hauptgang IPMN a Nachweis einer sehr großen Hauptgang IPMN mit rein zystischer Erweiterung (Pfeil). b Derselbe Patien t 3 Jahre später: Es finden sich nun deutliche, solide Tumoranteile (Pfeil) als Zeichen einer malignen Entartung.

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