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Benign Hepatic Neoplasms

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1. Introduction

Historically benign liver tumors were encountered incidentally during laparotomy or more recently during laparoscopy at which time definitive histological diagnosis can be established. However, with the utilization of advanced imaging modalities hepatic neoplasms have been increasingly identified, with a prevalence rate of up to 50% reported among the general population [1]. Among these incidental lesions, 83% were characterized as benign neoplasms, as outlined in Table 1 [1-3]. Benign hepatic neoplasms represent a diverse group of tumors that develop from either epithelial or mesenchymal cell lines (Table 2), and while the frequency of such lesions is not well documented, more than 50% are classified as hemangiomas [1]. Focal nodular hyperplasia (FNH) and hepatic adenomas represent the next most frequently diagnosed benign tumors. A variety of additional exceedingly rare benign lesions have also been described most of which are sufficiently infrequent enough to be classified as “fascinomas” [1].

Neoplasm	Relative frequency
Hemangioma	52%
Focal nodular hyperplasia	11%
Metastatic tumor (T _x N _x M1)	11%
Hepatocellular adenoma	8%
Focal fatty infiltration	8%
Hepatocellular carcinoma	6%
Extrahepatic process (eg., abscess, adrenal tumor)	3%
Other benign hepatic process	1%

Table 1. Diagnostic frequency of incidentally identified solid liver neoplasms^{1,2,9}

Cell of origin	Tumors
Epithelial	
Hepatocellular	Focal nodular hyperplasia (FNH)
	Hepatocellular adenoma (HA)
	Regenerative nodule
Cholangiocellular	Biliary adenoma
	Biliary cystadenoma
Other	Epithelioid leiomyoma
Mesenchymal	
Endothelial	Hemangioma
	Cavernous
	Capillary
	Hemangioendothelioma
	Adult
	Infantile
Mesothelial	Solitary fibrous tumor
	Benign mesothelioma
	Fibroma
Adipocyte	Lipoma
	Myelolipoma
	Angiomyelipoma
Miscellaneous	
Tumors	Biliary hamartoma

Table 2. Benign solid liver neoplasms^{1,9}

Most benign tumors are asymptomatic which makes standardizing the work-up difficult. The evaluation of incidental solid hepatic tumors should be individualized based upon the patient’s age, sex, past medical history, medications, and associated clinical signs. Although physical examination of the abdomen is typically unremarkable it may rarely reveal localized tenderness and/or a palpable mass. Liver function tests are indicated though are seldom abnormal in asymptomatic patients. Additional laboratory testing such as alpha-fetoprotein (AFP), carcinoembryonic antigen (CEA), carbohydrate antigen (CA) 19-9 and, lactate dehydrogenases may also be ordered depending on the clinical scenario.

Substantial advancements and the widespread availability and use of modern imaging modalities to diagnose and treat abdominal pain, has led to a marked increase in the identification of benign liver tumors. A full discussion of the advantages and disadvantages of

individual imaging techniques is beyond the scope of this chapter but is outlined in Table 3. Briefly, B-mode ultrasonography (US) can effectively differentiate cystic and solid neoplasms and is usually the initial study of choice [4,5]. Contrast-enhanced computed tomography (CT) provides greater sensitivity than US for determination of lesion number, size, and location [5, 6]. Magnetic resonance imaging (MRI) represents the most sensitive and specific study to discriminate between various benign liver lesions, particularly when contrast agents are used [5-7]. Finally, fluorodeoxyglucose positron emission tomography (^{18}F FDG-PET) can aid in the differentiation of benign versus malignant tumors based on the metabolic activity of the lesion [8]. Although modern imaging techniques can precisely diagnose the vast majority of incidental benign tumors, laparoscopic or open biopsy is necessary to exclude malignancy when precise diagnosis remains elusive.

Tumor	US	CT	MRI	Tc 99 RBC scan	Tc 99 SC scan
<i>Hemangioma</i>	Hyperechoic Well-demarcated Increased vascular flow Central venous pooling	Highly sensitive Non-contrast: Isodense Contrast: Hypoense Irregular peripheral enhancement with delayed central filling	Highly sensitive Isodense on T1 Hyperdense on T2 Gadolinium enhanced scan shows similar findings to contrast CT	Blood pooling of radionuclide	Not indicated
<i>Focal Nodular Hyperplasia (FNH)</i>	Non-specific Hyperechoic	Highly specific Non-contrast: isodense Contrast: isodense and well-demarcated with central scar	Highly specific Isodense of T1 & T2 Early hyperdense after gadolinium	Not indicated	Takes up Tc99 Tc99 SC contains bile ducts and Kupffer cells
<i>Hepatic Adenoma</i>	Non-specific Hyperechoic Increased blood flow on duplex scanning	Non-specific Non-contrast: hypo to isodense Contrast: isodense with peripheral enhancement with subsequent centripetal flow	Non-specific Iso- hypointense T1 & T2 Uniform enhancement after gadolinium	Not indicated	Generally does not take Tc99 SC because of the lack of bile ducts and Kupffer cells

US = ultrasonography; CT = computed tomography; MRI = magnetic resonance imaging; T1 = T1-weighted MRI; T2 = T2-weighted MRI; Tc-99m RBC = technetium-99m-labeled red blood cell; Tc-99m SC = technetium-99m sulfur colloid.

Table 3. Radiographic appearance of benign liver neoplasms^{1,9}

Accurate diagnosis is essential to the appropriate management of hepatic neoplasms. Although patients may require surgical intervention for diagnostic purposes, few benign tumors require surgical management for symptomatic relief. As such, surgical intervention for benign tumors is primarily indicated (1) for definitive diagnosis when imaging is inconclusive, (2) to prevent malignant transformation, such as in the case of hepatic adenoma, (3) to reduce the risk of rupture and, (4) for the treatment of rare life-threatening complications as a result of rupture or haemorrhage [9].

2. Hemangioma

Hemangioma is the most common benign mesenchymal neoplasm of the liver and occurs in two variants, capillary and cavernous. Hepatic hemangiomas are identified in 0.4% to 20% of all imaging studies performed [10-14]. Hemangiomas are frequently discovered incidentally

on autopsy studies with 60%-80% identified in individuals in their 4th- 6th decade of life [12-16]. The precise etiology of hemangiomas is poorly understood but they are generally considered to be benign congenital hamartomas composed of disorganized venous vasculature separated by intervening fibrous tissue [17]. Hemangiomas vary greatly in size from a few millimeters to over 50 cm, with the majority (up to 80%) less than 4 cm [1,12,18]. Although most commonly solitary, up to 40% of patients with hemangiomas have multiple tumors [19].

Capillary hemangiomas are more prevalent than are cavernous hemangiomas [1,20]. However, these hypervascular lesions are typically small (2 cm) and are rarely clinically significant [1]. As such, the management of capillary hemangiomas requires the exclusion of malignancy and patient reassurance that routine surveillance is not necessary in the absence of symptoms [9].

Cavernous hemangiomas are far more often clinically relevant than capillary hemangiomas. The incidence of cavernous hemangiomas is 3 times greater among women than men, with a mean age of 45 years [12,16]. Whether this reflects a true increase in incidence or a result of more frequent imaging amongst females remains unclear as evident by one autopsy series in which there was a nearly equal sex incidence [1,21]. Although no link between oral contraceptive pill (OCP) use and hemangioma incidence has been established, early studies suggest a link between OCP use and increased hemangioma size at initial presentation [18].

3. Clinical presentation

The most frequently reported symptoms of liver hemangiomas include abdominal pain, nausea, vomiting, early satiety, and prolonged fever [1,22]. Most symptoms of hepatic hemangioma are attributable to rapid expansion, thrombosis, or infarction, resulting in inflammation or stretching of Glisson's capsule [1]. Large hemangiomas (> 10 cm) may occasionally present as a non-tender palpable mass in the right upper quadrant, however physical exam more often reveals only vague abdominal tenderness without a mass [1,23]. Occasionally, a bruit may be detected over the liver. Evidence of intratumoral or intraperitoneal rupture may be reflected by hemoperitoneum and subsequent shock, which requires emergent surgical intervention. Rarely biliary colic, obstructive jaundice, gastric obstruction, torsion of a pedunculated lesion, pulmonary embolism, spontaneous intraperitoneal hemorrhage, and consumptive coagulopathy have been reported [22,24,25]. Kasabach-Merritt syndrome, which was originally used to describe thrombocytopenia and afibrinogenemia associated with hemangiomas on the skin and spleen of infants, is frequently used to define hepatic hemangioma patients with severe thrombocytopenia and concomitant consumptive coagulopathy [26].

4. Pathology

Hemangiomas are typically well demarcated from surrounding hepatic tissue, which often permits surgical enucleation [27]. In tumors not well demarcated, the tumor-parenchymal interface defines the ease with which enucleation versus formal resection is required. Four

interface variants between the hemangioma and hepatic parenchyma have been described. The “fibrolamellar” interface is characterized by a capsule-like fibrous ring of various thickness and is the most common [9]. The involved veins parallel the periphery of the hemangioma or traverse the fibrous lamella. The healthy hepatic parenchyma is often atrophic and a plane between the hemangioma and uninvolved liver tissue is well defined. A second variant, the “compression” interface consists of a hemangioma in which the periphery of the neoplasm is well demarcated despite the absence of a fibrous lamella [1]. An “interdigiting” pattern lacks a fibrous lamella and instead is replaced by an ill-defined plane between the vascular channels of the hemangioma and uninvolved hepatic parenchyma [1]. Finally, an “irregular” or “spongy” interface occur when the hemangioma appears to intercalate into the surrounding hepatic parenchyma [1]. Despite the invasive appearance of this variant, hemangiomas do not possess any malignant potential.

The diagnosis of cavernous hemangioma is generally easy to establish with modern imaging techniques. However, in some instances atypical hemangiomas may be confused for other pathology, including but not limited to, hemorrhagic telangiectasia (Osler-Rendu-Weber), hemangioendothelioma, and peliosis hepatis [9]. When diagnosis remains unclear, indeterminate lesions should be managed surgically as percutaneous biopsy may result in uncontrollable hemorrhage [1].

5. Radiographic evaluation

Accurate radiographic diagnosis of hepatic hemangioma is essential since once definitive diagnosis is established no additional intervention is typically required [9]. Radiographic evaluation is largely dictated by clinical presentation as most hemangiomas are discovered incidentally on imaging studies completed for unrelated symptomology and/or pathology. Depending on the initial degree of diagnostic certainty additional imaging maybe superfluous.

B-mode ultrasonography is typically the initial imaging study performed [1]. On US hemangiomas appear as a homogenous hyperechoic mass that is well demarcated from surrounding liver parenchyma [1,28,29]. The addition of duplex US provides additional information regarding peripheral blood flow and central pooling of venous blood [1,28]. As malignant lesions may demonstrate similar acoustic patterns, additional imaging modalities are often required for definitive confirmation. On contrast enhanced compute tomography (CE-CT) hemangiomas initially appear as hypodense masses with a pattern of irregular peripheral nodular enhancement following initial injection of contrast [30,31]. Delayed venous images subsequently demonstrate characteristic central venous filling of the hypodense mass [30,31]. Magnetic resonance imaging (MRI), though rarely needed for diagnosis of most hemangiomas, is the most sensitive and specific modality for the detection and diagnosis of hemangioma [6,32]. T-1 weighted images reveal a smooth well-demarcated homogenous isodense mass, whereas T-2 weighted studies demonstrate a hyperdense pattern [33,34]. The administration of intravenous gadolinium diethylenetriaminepentaacetic acid (Gd-DTPA) contrast results in the pathognomonic pattern of peripheral nodular enhancement with central filling on delayed

images [1,35,36]. This enhancement pattern is typical of most hemangiomas > 2 cm [37]. Hemangiomas < 2 cm may demonstrate rapid uniform enhancement which is indistinguishable from hypervascular hepatocellular carcinoma (HCC) [37]. ^{18}F -FDG PET scan may be useful for differentiation between benign and malignant hepatic tumors [38]. Studies have shown that the activity of both glucose-6-phosphatase and glucose transporters are increased in HCC resulting in decreased uptake of ^{18}F -FDG in hemangiomas as compared to HCC [8]. Historically, technetium-99 labeled red blood cells scintigram (Tc-99 RBC scan) was the gold standard for the diagnostic evaluation of hemangiomas, but technological advancements in axial imaging has led to a decline in the reliance on RBC scintigraphy [31,39]. Finally, selective hepatic angiography typically yields a characteristic neovascular “corkscrewing” appearance with rapid central filling from the neovascular periphery described as “cottonwool” [1]. Despite these characteristic findings, the high diagnostic yield of less invasive modalities makes arteriography rarely necessary.

6. Diagnosis & treatment

The majority of hemangiomas are asymptomatic, particularly those lesions < 1.5 cm in size [1]. Although hemangiomas can grow to great sizes, they generally do not compromise liver function and as such liver function tests are often normal. In rare instances thrombosis or intraparenchymal hemorrhage may occur acutely affecting liver function tests. Spontaneous rupture of hepatic hemangiomas is an exceptionally rare event with a review of the literature revealing less than 30 cases of spontaneous rupture since 1898. Given the low yet significant risk of bleeding, fine needle aspiration (FNA) should be avoided [1]. As a rule, biopsy is only indicated if a histologic diagnosis is unclear or will alter planned treatment, thus in the absence of clinical symptoms the most appropriate treatment strategy is careful observation [1].

Surgical resection should be considered in patients with disabling pressure or pain suggestive of extrinsic compression of adjacent structures, in those experiencing acute symptoms related to rupture, or when malignancy cannot be ruled out [22,40]. In general clinical symptoms increase concurrently with tumor size, with most symptomatic tumors having a mean size of 10 ± 8 cm as compared with 6.8 ± 5.8 cm for asymptomatic lesions [41].

Surgical intervention should be approached no differently than for treatment of other hepatic tumors. It is essential that surgeons possess an extensive knowledge of the anatomy and vascular supply of the liver. The extent of hepatic resection required is directly related to the anatomic location of the lesion and its proximity to surrounding vasculature. Thus, the location of the lesion will largely dictate the operative approach hence a full evaluation of the tumor's extent is critical. Large central lesions which border the inferior vena cava, hepatic outflow tract, or the portal vein, may pose an exorbitant surgical risk and as such may not allow for resection [1].

While enucleation is often indicated, formal resection is required in certain instances. Recall it is the histological features of the tumor-parenchymal interface which defines how easily a parenchymal-sparing technique may be utilized. Unlike malignant lesions, resection of

hemangiomas does not necessitate removal of a margin of normal tissue with the tumor. Enucleation is carried out by careful dissection within the proper plane between the hepatic parenchyma and tumor. Division and ligation of the principal hepatic artery should be completed early in the operation as this often results in significant tumor decompression thereby facilitating resection [1]. The majority of hemangiomas are contained within a tough fibrous capsule which can be clamped and used for retraction purposes [1]. As hepatic venous branches are encountered extending from the lesion they should be controlled with clips or ties [1]. Presently, mortality outcomes for resection and enucleation are comparable [42].

Hepatic artery ligation for treatment of hemangioma has also been described anecdotally [9]. Although its benefits are likely transient, hepatic artery embolization and/or ligation play a pivotal role only in the temporary management of uncontrolled hemorrhage from rupture [43,44]. Finally, radiation therapy for symptomatic hemangiomas has also been reported. Though data validating the use of radiotherapy is limited, it seems a reasonable approach for symptomatic hemangioma where surgical intervention is clearly contraindicated.

7. Special issue: Hemangioma in children

Hepatic hemangiomas of infancy and childhood differ substantially in their appearance, presentation, and progression than those in adults [1]. These lesions are frequently large and symptomatic. In contrast to adult hemangiomas, the risk of spontaneous rupture in infancy is greater [1]. Similarly, Kasabach-Merritt syndrome occurs more frequently and results more often in death among affected infants. As a result of the numerous venous lakes within these lesions, which serve as siphons for a large proportion of the total cardiac output, severe congestive heart failure and death may result. Initial treatment of high output cardiac failure in children includes oxygen, diuretics, digitalis, corticosteroids, hepatic artery ligation, and radiation therapy [2, 45-48]. Contrary to the conservative management of adult hemangiomas, hemangiomas of infancy and childhood more frequently require life-saving surgical intervention.

8. Focal nodular hyperplasia

Focal nodular hyperplasia (FNH) is the second most common benign hepatic lesion [20]. FNH is found predominately in women (in a ratio of 8-9:1) between the ages of 20-50 years, and has a prevalence of 4 - 8% in the general population [49,50]. Similar to hemangiomas, the prevalence of FNH has markedly increased over the past several decades, which likely reflects the proficiency and widespread use of advanced imaging modalities [1].

Although Klatskin (1977) and Vana (1979) each reported an association between OCP use and the development of FNH, the high frequency of FNH in the absence of OCP use suggests no causal relationship [32,51]. However, enlargement of FNH lesions has been described in the setting of pregnancy and long-term OCP use [52]. While the etiology of these lesions has not

yet been clearly delineated, it has been suggested that FNH is a hyperplastic polyclonal response of normal hepatic parenchyma to localized areas of increased arterial perfusion [53]. Expectantly, FNH has been found in association with vascular disorders and malformations including hereditary hemorrhagic telangiectasia, hemihypertrophy Klippel-Trenaunay-Weber syndrome, and congenital absence of the portal vein [49,54-57].

While typically small (< 5 cm), FNH lesions have been reported as large as 19 cm [48,50]. The majority of FNH lesions are solitary in nature (80%-95%), although up to 20% of individuals are reported to have multiple lesions [1, 48, 50]. When multifocal, FNH often occurs in conjuncture with other benign hepatic lesions including hemangiomas [58].

9. Clinical presentation

FNH is frequently asymptomatic with up to 75% of lesions discovered incidentally during radiologic workup, laparotomy, or laparoscopy for unrelated pathology [59]. Similar to hepatic hemangiomas, spontaneous rupture is extremely rare as illustrated by Chamberlain et. al (2003) management of 33 patients with FNH where no ruptures were evident [9]. Large, peripheral, pedunculated lesions may result in a palpable mass associated with abdominal pain and/or fullness, but acute symptoms associated with rupture, necrosis, or infarction are a rarity.

10. Pathology

Macroscopically FNH is a firm pale to red colored lesion with sharp margins. Lesions are typically small, pedunculated, and peripherally located. Unlike hemangiomas and hepatic adenomas, FNH lack a capsule. Histologically FNH appears as regenerative nodules making histopathological differentiation from cirrhosis difficult. Lesions contain normal hepatic elements with a haphazard arrangement of cords and sinusoids [5]. Proliferating bile ducts, fibrous septae, Kupffer cells, and sinusoids are typically present in FNH, and are characteristically absent in hepatocellular adenomas [13,50,59]. Generally FNH contain a large artery with multiple branches radiating through disorganized fibrous septa to the periphery. This radiating arterial pattern produces a spoke and wheel image on angiography and is responsible for the central scar appearance on radiographic imaging studies [60,61].

11. Radiographic imaging

Definitive diagnosis of FNH can be challenging. FNH lesions are well visualized on US but are highly variable and exhibit no distinct characteristic features. Helical CE-CT reveals a well-demarcated lesion that is often isodense [29]. However, during the portal venous phase the pathognomonic central scar may be appreciated. Distinguishing FNH on standard MRI can

prove challenging as the lesion is composed of the similar elements as the normal liver parenchyma. FNH may appear isointense with a central scar on T-1 and T-2 weighted imaging [62]. MRI with Gd-DTPA demonstrates a hyperintense lesion early, which becomes isointense with central scar enhancement on delayed imaging [63-65]. The use of reticuloendothelial agents including Ferridex, which is taken up selectively by Kupffer cells, increases the specificity of both CT and MRI imaging [1]. Technetium-99-labeled sulfur colloid scintigraphy may prove helpful in demonstrating the presence of Kupffer cells within the FNH lesion, however this finding is not specific enough for definitive diagnosis [1,66,67]. Angiography, though rarely indicated for the diagnosis of FNH, usually demonstrates a hypervascular mass with a single central artery and enlarged peripheral vessels in a “spoken wheel” appearance [66-68]. Finally, ^{18}F -FDG PET can aid in the differentiation between benign and malignant lesions, but it is neither sensitive nor specific enough for diagnosis of FNH [8,38].

12. Diagnosis & treatment

The natural course of an FNH lesion is generally indolent with minimal risk of rupture or complication. Laboratory testing generally reveals normal liver function tests and alpha-fetoprotein levels, although minor elevations in aspartate and alanine aminotransferase, alkaline phosphatase, and gamma glutamyl transpeptidase may occasionally be seen. Definitive diagnosis of FNH in an asymptomatic patient warrants conservative management and includes close observation with repeat imaging every four to six months [9]. When radiology is equivocal, most surgeons still choose close observation with follow-up studies preformed every three to four months. Biopsy is generally not indicated, as results are seldom diagnostic [69].

Although it may be impossible to distinguish FNH from a well-differentiated HCC without surgical excision, FNH tumors do not undergo malignant transformation. Thus indications for surgical intervention should be limited to those situations where there is a change in the size or number of lesion(s), a change in the intensity of symptoms, or where classic imaging characteristics are absent and diagnostic dilemma remains [70]. Hence, the role of the surgeon is typically limited to patient reassurance and close observation [9].

13. Hepatic adenoma

Hepatic adenomas are identified predominately in women of reproductive age [49]. The estimated prevalence of hepatic adenomas within the general population on postmortem exams is approximately 1% [10]. Etiologically, hepatic adenomas are of epithelial origin. Unlike hepatic hemangiomas and FNH, a clear association between the use of OCPs and hepatic adenomas has been established. First described in 1973, multiple studies have documented a reciprocal relationship between OCP use and adenoma incidence based on estrogen dose and exposure time [71-75]. Approximately 90% of individuals with adenomas have previous OCP

exposure [1]. The prevalence of hepatic adenomas is estimated at 1 per 1,000,000 among women who have never used OCP as compared with 30-40 per 1,000,000 amongst long-term OCP users [72,76]. OCPs also affect the course of disease progression as lesions are generally larger, more numerous, and more likely to bleed than tumors in OCP-naïve individuals [32,75,77,78]. Adenoma regression has been observed in patients after discontinuation of OCP with recurrence ensuing during pregnancy and/or OCP re-administration [72,79,-82]. Despite these findings, the mechanism by which estrogen therapy affects the development and course of hepatic adenomas has yet to be clearly elucidated.

Hepatic adenomas are typically small (< 5 cm), soft, solitary lesions but may be multiple in up to 30% of cases [9]. Of note, hepatic adenomatosis disease, defined as the presence of >10 lesions, is a distinct disease entity from that of hepatic adenoma and as such will not be described in further detail [83]. Hepatic adenomas have been associated with type I glycogen storage disease, galactosemia, Klienfelter's syndrome, and Turner's syndrome as well as with androgen, domiphen, danazol and growth hormone use [1,84-86]. Although hepatic adenomas are benign, these lesions have been associated with spontaneous hemorrhage, rupture, and malignant transformation, making prognosis more grave than that of other benign hepatic tumors [5,87].

14. Clinical presentation

Since adenoma and FNH both present in women of reproductive age and have similar radiographic appearances they are frequently confused. Differential diagnosis is critical given that the recommended treatment of each respective lesions differs. Hepatic adenomas are most often diagnosed as a result of imaging done for unrelated pathology or following workup of a palpable abdominal mass (30% patients) [88]. Occasionally episodic pain may be evident as a result of an enlarged liver, intratumoral bleed, or tumor necrosis [9]. Up to 33% of patients with hepatic adenomas present with acute rupture and concomitant intraperitoneal bleeding [1]. The development of acute severe pain associated with hypotension reflects spontaneous rupture and carries a 20% mortality rate if not appropriately identified and treated [32,89-91].

15. Pathology

Grossly hepatic adenomas appear as smooth, soft, and pale yellow tumor on cut surface [1]. These lesions often contain prominent blood vessels that have a high potential for rupture and hemorrhage [1]. As adenomas lack a fibrous capsule intraparenchymal bleeding may occur, which frequently results in a variegated appearance.

Microscopically hepatic adenomas appear as well circumscribed lesions composed of monotonous sheets of hepatocytes laden with glycogen and lipids [5]. These lesions lack normal hepatic architecture and demonstrate thickened trabeculae interspersed with sinusoids and

prominent thin walled vessels [1,5]. Biliary ducts and portal tracts are distinctly absent from adenomas.

While the malignant potential of adenomas remains controversial, several authors have reported a low (5%) yet consistent risk of transformation [87]. Histological differentiation between well differentiated HCC and adenoma can be difficult, especially in the presence of fibrolamellar HCC which is also more common in women of reproductive age. This issue is further explained in situations in which HCC and hepatic adenoma have been found adjacent to one another [61,50,89,92,93].

16. Radiological imaging

Although radiographic evaluation is important for complete workup of hepatic adenoma radiographic features are often nonspecific [94]. As such, despite the use of multiple imaging techniques, diagnosis often remains equivocal. Ultrasound exhibits a mixed echogenic pattern with an overall heterogeneous appearance [1,29]. Lesions appear hyperechoic as a result of their high lipid content with a heterogeneous pattern reflecting intratumoral hemorrhage and necrosis [95]. CE-CT imaging is frequently utilized for adenoma visualization and typically demonstrates a hypo- to isodense lesion as a result of low attenuation on non-contrast phase [1]. A variegated appearance with peripheral enhancement during the early contrast phase with subsequent centripetal flow during the venous phase may be apparent, however CT can demonstrate a spectrum of disparate findings [96]. MRI findings for hepatic adenoma are similar to those on CT. Due to the high fat and glycogen content, adenomas are usually well demarcated on MRI imaging [29]. While most adenomas appear iso- to hyperintense on both T-1 and T-2 weighted images, findings are highly variable [1,97]. The administration of contrast agents including gadolinium or gadobenate dimeglumine (Gd-BOPTA) results in early markedly uniform enhancement on arterial phase, which subsequently becomes isodense on the portal venous phase [98]. The use of ¹⁸FDG PET scan may also aid in the differentiation of benign versus malignant disease in which where adenomas demonstrate poor uptake of ¹⁸FDG as compared to HCC [8,38].

Additional imaging modalities infrequently used include technetium-99 sulfur colloid scanning. This imaging modality is particularly useful in differentiating between hepatic adenoma and FNH, as hepatic adenomas lack bile duct components and frequently appear as a “cold nodules” on imaging [99]. Occasionally however, a minority of lesions do take up the sulfur colloid, rendering them indistinguishable from FNH [99]. Although rarely utilized, angiography typically reveals hypervascular lesions with areas of hemorrhage and necrosis [1,28].

17. Diagnosis & treatment

In the absence of acute hemorrhage, serological tests rarely assist in diagnosis. Liver function tests and tumor makers including CEA, alpha-fetoprotein, and CA 19-9 are invariably normal.

Hepatic adenomas pose a greater risk for rupture (33%) and malignant transformation (5%) than do other benign hepatic lesions [9,87]. As such all patients with suspected or confirmed hepatic adenoma > 3 cm should undergo enucleation or surgical resection [1,100]. The approach to surgical excision should be as previously described. Since all adenomas are suspected to harbor malignancy an adequate margin of normal parenchyma should be taken [1]. When surgical exploration is not feasible angiographic embolization or ligation can provide temporary yet life saving relief.

As a result of the relationship between OCP and adenoma incidence, it is recommended that all individuals suspected of having an adenoma discontinue the use of OCP immediately and indefinitely [1,61]. Patients should also be advised against pregnancy until after adenoma resection, as the growth and rupture risk of hepatic adenomas is highly unpredictable during gestation [101]. Yearly follow-up with imaging is advised among all patients where a causal link between OCP use and adenoma is absent [9]. As a result of improved safety of hepatic resection and the use of minimally invasive techniques in hepatectomy it is suggested that all hepatic adenomas > 3 cm be resected [1,100]. In patients with significant contraindications to surgical intervention, OCP should be discontinued and the patient enrolled in an ongoing surveillance program [9].

18. Additional liver tumors

18.1. Epithelial tumor

Biliary hamartomas

Bile duct adenomas and hamartomas are common tumors. Bile duct adenomas appear as small, white, solitary, subcapsular masses [1]. They are defined histologically by narrow lumen bile ducts surrounded by fibrosis. Hamartomas appear as small gray-white nodules that lie just beneath the capsule of the liver [102]. Biliary hamartomas are frequently multifocal and are characterized microscopically by the presence of dilated mature bile ducts surrounded by fibrous tissue [1]. These lesions are especially important as they are frequently misinterpreted as metastatic tumor by the operating surgeon. This notion heightens the importance of confirmatory diagnosis to rule out malignancy for all hepatic lesions. Precise diagnosis is most important in situations in which the presence of a metastatic liver disease will alter the proceedings of a planned operation.

18.2. Mesenchymal tumors

Solitary fibrous tumor (other names include benign mesothelioma or fibroma)

Solitary fibrous tumors (SFT) are rare mesenchymal tumors that are frequently mistaken for metastatic lesions as a result of their radiographic and intra-operative appearance. Grossly SFT's appear as white-to-gray lesions and can vary greatly in size ranging from 2 – 20 cm in diameter [1]. Despite their large size, most SFT's remain asymptomatic. Histologically, most have a classic short storiform pattern and display an absence of cellular atypia, mitoses,

and/or necrosis [1]. However when malignant, SFTs frequently possess a high mitotic rate and marked cellular atypia. Immunohistochemically SFTs display a strong positive staining for vimentin and CD-34 [1]. Since definitive histologic examination is required for diagnosis of either a benign or malignant SFT, surgical resection is indicated in nearly all circumstances.

Lipoma, myelolipoma, or angiomyelipoma

Similar to several other benign hepatic lesions, most benign fatty hepatic tumors are identified at the time of autopsy with only isolated reports of histological diagnosis following operative resection [13]. Multiple variants including angiolipoma, myelolipoma, and angiomyolipoma have been described [13,103]. Additionally, “pseudolipomas” have been described as lesions in which there is an extracapsular fatty tumor with involutional changes. It is probable that this lesion results when a free-floating piece of fat becomes entrapped between diaphragm and liver surface [1,10]. In most situations definitive diagnosis requires surgical resection to exclude malignancy.

Mesenchymal Hamartomas

Mesenchymal hamartomas are exceedingly rare congenital liver tumors which occur most frequently in infants under 1 year of age [9,104]. Microscopically these lesions demonstrate a myxoid background of highly cellular embryonal mesenchyme with haphazard groupings of bile ducts, cysts, and hepatic cells [105]. Generally, the cystic element is the most prominent feature resulting in a characteristic “honeycomb” appearance [106]. In contrast to biliary hamartomas, which are clinically insignificant, mesenchymal hamartomas can significantly impair hepatic function as a result of their large size [106]. Although benign, these lesions can result in death due to mass effect and/or hepatic insufficiency [1]. Thus, all suspected mesenchymal hamartomas should be completely excised when possible. If complete surgical excision cannot be achieved surgical debulking may be sufficient as there have been no reports of recurrence after an incomplete surgical resection to date [107].

Myxoma

Myxomas are exceptionally uncommon benign lesions of the liver. To date fewer than five cases have been reported [9,58,108]. These lesions arise from primitive connective tissue. Histologically myxomas demonstrate a myxoid matrix with scattered proliferation of connective tissue cells [108]. Similar to other types of hepatic tumors described above, surgical resection is generally indicated to exclude malignancy.

Teratoma

Primary teratomas are remarkably rare benign hepatic lesions. A review of the literature revealed only 7 reports to date, with the majority of lesions occurring in children [109]. Secondary hepatic teratomas have been observed following systemic chemotherapy administration for treatment of testicular cancer [1]. Teratomas arise from pluripotent cells and frequently contain components from all three germ layers. Teratomas are typically encapsulated cystic lesions that are easily resectable [1,110]. Imaging characteristics reflect tissue heterogeneity and are often non-specific [110]. Surgical resection of hepatic teratomas is indicated to exclude malignancy.

19. Conclusion

A thorough understanding of the natural history and accurate histologic diagnosis are fundamental to appropriate management of patients with benign liver tumors. Although advancements in imaging have drastically improved the detection and characterization of both benign and malignant liver neoplasms, the ultimate burden of responsibility for diagnosis and treatment remains that of the surgeon. Ongoing improvements in perioperative care and surgical techniques, coupled with increased surgical experience presently permit hepatic resection to be performed with a high level of safety. Despite these developments, a conservative approach including close observation with serial examination and imaging seems most appropriate for asymptomatic patients in which malignancy is not suspected.

Symptomatic patients without medical or anatomic contraindication to a major hepatic resection, as well as patients in whom a malignancy cannot be excluded (including individuals with adenomas > 3 cm), should be considered for surgical intervention. Preoperative needle biopsy is frequently contraindicated due to a high risk of rupture and hemorrhage, and therefore should only be considered after exclusion of hemangioma. Additionally, it is important to note that distinguishing particular lesions (especially adenoma and FNH) on needle biopsy is exceedingly difficult. As such caution should be exercised when using this information to make clinical evaluations. Excisional biopsy of small and peripheral lesions and adequate wedge incision biopsy of large lesions should permit the pathologist to make an accurate histologic diagnosis and exclude a malignancy. If doubt remains, formal hepatic resection is indicated.

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