

A nationwide assessment of hepatocellular adenoma resection: indications and pathological discordance

Haring, M.P.D.; Elfrink, A.K.E.; Oudmaijer, C.A.J.; Andel, P.C.M.; Furumaya, A.; Jong, N. de; ...; Dutch Benign Liver Tumor Group

Citation

Haring, M. P. D., Elfrink, A. K. E., Oudmaijer, C. A. J., Andel, P. C. M., Furumaya, A., Jong, N. de, ... Meijer, V. E. de. (2022). A nationwide assessment of hepatocellular adenoma resection: indications and pathological discordance. *Hepatology Communications*, 7(1). doi:10.1002/hep4.2110

Version: Publisher's Version

License: Creative Commons CC BY-NC-ND 4.0 license

Downloaded from: https://hdl.handle.net/1887/3567038

Note: To cite this publication please use the final published version (if applicable).

DOI: 10.1002/hep4.2110

ORIGINAL ARTICLE





A nationwide assessment of hepatocellular adenoma resection: Indications and pathological discordance

```
Martijn P. D. Haring<sup>1</sup> | Arthur K. E. Elfrink<sup>2,3</sup> | Christiaan A. J. Oudmaijer<sup>4</sup> | |
Paul C. M. Andel<sup>5</sup> | Alicia Furumaya<sup>6</sup> | Nenke de Jong<sup>7</sup> | Colin J. J. M. Willems<sup>8</sup> |
Thiis Huits<sup>9</sup> | Julie M. L. Siimons<sup>10</sup> | Eric J. T. Belt<sup>11</sup> | Koop Bosscha<sup>12</sup> | I
Esther C. J. Consten<sup>1,13</sup>  | Mariëlle M. E. Coolsen<sup>8</sup>  | Peter van Duijvendijk<sup>14,15</sup>  |
Joris I. Erdmann<sup>6</sup>  Paul Gobardhan<sup>16</sup>  Robbert J. de Haas<sup>17</sup>  |
Tjarda van Heek<sup>18</sup> | Hwai-Ding Lam<sup>19</sup> | Wouter K. G. Leclercg<sup>10</sup> |
Wouter W. te Riele<sup>5</sup> | Jeroen Hagendoorn<sup>5</sup> | Alexander F. M. Schaapherder<sup>19</sup> | |
Koert P. de Jong<sup>1</sup> | Vincent E. de Meijer<sup>1</sup> | on behalf of Dutch Benign Liver Tumor
Group
```

Supplemental Digital Content is available for this article. Direct URL citations appear in the printed text and are provided in the HTML and PDF versions of this article on the journal's website, www.hepcommjournal.com.

This is an open access article distributed under the terms of the Creative Commons Attribution-Non Commercial-No Derivatives License 4.0 (CCBY-NC-ND), where it is permissible to download and share the work provided it is properly cited. The work cannot be changed in any way or used commercially without permission from the

Copyright © 2023 The Author(s). Published by Wolters Kluwer Health, Inc. on behalf of the American Association for the Study of Liver Diseases.

¹Department of Surgery, University Medical Center Groningen, University of Groningen, Groningen, the Netherlands

²Scientific Bureau, Dutch Institute for Clinical Auditing, Leiden, the Netherlands

³Department of Surgery, Spaarne Gasthuis, Haarlem, the Netherlands

⁴Division of Hepatobiliary and Transplantation Surgery, Department of Surgery, Erasmus Medical Center Transplant Institute, Erasmus University Medical Center, Rotterdam, the Netherlands

⁵Department of Surgery, Regional Academic Cancer Center Utrecht, Utrecht Medical Center Utrecht and St. Antonius Hospital Nieuwegein, University of Utrecht, Utrecht, the Netherlands

⁶Department of Surgery, Amsterdam Gastroenterology Endocrinology Metabolism, Amsterdam University Medical Center, University of Amsterdam, Amsterdam, the Netherlands

⁷Department of Surgery, Medisch Spectrum Twente, Enschede, the Netherlands

⁸Department of Surgery, Maastricht University Medical Center+, University of Maastricht, Maastricht, the Netherlands

⁹Department of Surgery, Onze Lieve Vrouwe Gasthuis, Amsterdam, the Netherlands

¹⁰Department of Surgery, Máxima Medisch Centrum, Veldhoven, the Netherlands

¹¹Department of Surgery, Albert Sweitzer Ziekenhuis, Dordrecht, the Netherlands

¹²Department of Surgery, Jeroen Bosch Ziekenhuis, 's-Hertogenbosch, the Netherlands

¹³Department of Surgery, Meander Medisch Centrum, Amersfoort, the Netherlands

¹⁴Department of Surgery, Gelre Ziekenhuis, Apeldoorn, the Netherlands

¹⁵Department of Surgery, Isala Klinieken, Zwolle, the Netherlands

¹⁶Department of Surgery, Amphia Ziekenhuis, Breda, the Netherlands

Correspondence

Vincent E. de Meijer, Department of Surgery, University Medical Center Groningen, University of Groningen, Hanzeplein 1, 9700 RB, Groningen, the Netherlands.

Email: v.e.de.meijer@umcg.nl

Abstract

Hepatocellular adenomas (HCAs) are benign liver tumors associated with bleeding or malignant transformation. Data on the indication for surgery are scarce. We analyzed indications and outcome of patients operated for HCAs < 50 mm compared to HCAs > 50 mm. Changes in final postoperative diagnosis were assessed. We performed a retrospective study that included patients who underwent resection for (suspected) HCAs in the Netherlands from 2014 to 2019. Indication for resection was analyzed and stratified for small (<50 mm) and large (≥50 mm) tumors. Logistic regression analysis was performed on factors influencing change in tumor diagnosis. Out of 222 patients who underwent surgery, 44 (20%) patients had a tumor <50 mm. Median age was 46 (interquartile range [IQR], 33-56) years in patients with small tumors and 37 (IQR, 31–46) years in patients with large tumors (p = 0.016). Patients with small tumors were more frequently men (21% vs. 5%, p = 0.002). Main indications for resection in patients with small tumors were suspicion of (pre) malignancy (55%), (previous) bleeding (14%), and male sex (11%). Patients with large tumors received operations because of tumor size >50 mm (52%), suspicion of (pre)malignancy (28%), and (previous) bleeding (5.1%). No difference was observed in HCA-subtype distribution between small and large tumors. Ninety-six (43%) patients had a postoperative change in diagnosis. Independent risk factors for change in diagnosis were tumor size <50 mm (adjusted odds ratio [aOR], 3.4; p < 0.01), male sex (aOR, 3.7; p = 0.03), and lack of hepatobiliary contrast-enhanced magnetic resonance imaging (CE-MRI) (aOR, 1.8; p = 0.04). Resection for small (suspected) HCAs was mainly indicated by suspicion of (pre)malignancy, whereas for large (suspected) HCAs, tumor size was the most prevalent indication. Male sex, tumor size < 50 mm, and lack of hepatobiliary CE-MRI were independent risk factors for postoperative change in tumor diagnosis.

INTRODUCTION

Hepatocellular adenomas (HCAs) are benign liver tumors that are frequently associated with chronic oral contraceptive pill use and obesity. [1,2] Complications are associated with tumor size \geq 50 mm. Large HCAs (\geq 50 mm) are associated with hemorrhage (15%–20%) and a small chance of malignant transformation to hepatocellular carcinoma (HCC; 1.6%), whereas both of these

complications are rare in HCAs $< 50\,\mathrm{mm.}^{[3-6]}$ According to current international guidelines, size $> 50\,\mathrm{mm}$ is an indication for resection. $^{[7,8]}$ The role of liver resection for the treatment of smaller HCAs, however, remains unclear. $^{[7]}$

The key to noninvasive HCA management is the HCA's ability to stabilize or regress in size after estrogen lowering and lifestyle advice (oral contraceptive pill cessation and weight loss).^[2,9] Since 2016, European guidelines recommend lifestyle changes for 6 months

¹⁷Department of Radiology, University Medical Center Groningen, University of Groningen, Groningen, the Netherlands

¹⁸Department of Surgery, Ziekenhuis Gelderse Vallei, Ede, the Netherlands

¹⁹Department of Surgery, Leiden University Medical Center, University of Leiden, Leiden, the Netherlands

²⁰Department of Surgery, Amsterdam Gastroenterology Endocrinology Metabolism, Amsterdam University Medical Center, Vrije Universiteit, Amsterdam, the Netherlands

before surgery in HCAs \geq 50 mm in women, after which HCA response is evaluated. Invasive treatment is recommended if HCA size remains \geq 50 mm, whereas a noninvasive approach is advocated in HCAs < 50 mm. The 50-mm diameter cutoff for invasive treatment is regardless of any significant response in terms of regression in size. International guidelines unequivocally recommend intervention for HCAs in all male patients because of the high rate of malignant transformation, regardless of any co-occurring metabolic disease. In HCA-related symptoms include nausea, fatigue, bloating, and pain. These symptoms have been related to significant quality of life (QoL) impairment, and surgical resection may be an effective treatment relief.

Indications for resection of HCAs are clear and concise in current European guidelines, which discourage resection of HCAs $<50\,\mathrm{mm}$ in women. However, up to one third of all resected benign liver tumors, including HCAs, are $<50\,\mathrm{mm}$. Data on indications for resection in this specific group remain scarce. [13–15] Evaluation of indications for resection of HCAs, and especially HCAs $<50\,\mathrm{mm}$, could assist clinicians and patients in future treatment decisions. In the current study, we aimed to provide an evaluation of resection indications for small (suspected) HCAs $<50\,\mathrm{mm}$ in comparison to larger HCAs ($\geq50\,\mathrm{mm}$) in a nationwide cohort. We also analyzed changes in final postoperative diagnosis.

MATERIALS AND METHODS

A nationwide observational cohort study was performed in the Netherlands. Data were retrieved from the Dutch Hepato Biliary Audit, which is a mandatory nationwide registry in which all Dutch liver surgery centers record all liver resections performed. Data verification was performed by a trusted third party to provide insight into data completeness and quality.[16] Additional data, including indications for resection, were collected from local electronic patient files. The Strengthening the Reporting of Observational Studies in Epidemiology (STROBE) guidelines were adhered to in study design and manuscript preparation.[17] The Medical Ethical Committee of the University Medical Center Groningen (UMCG) confirmed that the Law on Medical Scientific Research involving human subjects did not apply (MEC 2020-004). All local ethical and scientific committees were consulted for study approval. The study was registered before initiation in the UMCG research registry (UMCG RR#201900849) and in all local research registries when obliged.

Patient selection

Included were patients who underwent liver resection in the Netherlands for (presumed) HCAs (i.e., patients with suspected HCA but later with a proven alternate diagnosis, for example, focal nodular hyperplasia [FNH] or malignancy, were also included). Inclusion period was defined as patients having an operation between January 1, 2014, and December 31, 2019, and registered in the mandatory audit before April 1, 2020. Patients were excluded when surgery was indicated by (suspicion of) hepatic malignancy, but definitive post-operative pathological tumor diagnosis was HCA. Patients were also excluded if information regarding date of birth, date of surgery, or type of intervention was missing.

Outcomes

Primary outcome was the main indication for resection for presumed HCAs as determined by the local (multidisciplinary tumor board) practitioner(s), stratified for small HCAs (largest tumor diameter $<50\,\text{mm}$) and large HCAs (largest tumor diameter $\geq50\,\text{mm}$). Indications were compared between regional hospital versus tertiary referral hospitals. When multiple HCAs were resected, the largest tumor diameter was registered.

Secondary outcomes included 30-day major morbidity, defined as a Clavien-Dindo grade Illa or greater complication (i.e., requiring re-intervention, medium care or intensive care management, organ failure, or death) within 30 days of surgery, and 30-day mortality defined as death within 30 days of surgery. Changes in preoperative diagnosis and diagnosis after final postoperative histopathological analysis were scored. Patients for whom there was doubt on preoperative diagnosis were also scored as such (e.g., preoperative doubt on HCA/FNH). In this analysis, all (suspected) primary and secondary malignancies were characterized as "malignancy."

Variables

Patient characteristics included age, sex, American Society of Anesthesiologists classification, comorbidity score according the Charlson Comorbidity Index, previous medical history of liver disease, and a history of previous liver resection. Tumor characteristics included number of HCAs and diameter of largest HCA before treatment as well as subtype of HCA. Treatment characteristics included surgical approach (i.e., open or minimally invasive approach), major (three or more adjacent Couinaud hepatic segments) or minor resection, and type of hospital (i.e., tertiary referral hospital or regional hospital) where treatment was performed. The conclusion on any (pre)malignant tumor was derived from the original radiology and histopathology reports. The indication "suspicion of (pre)malignancy" was only scored as such if the radiology and/or histopathology reports regarded the tumors as such. Imaging and/or pathology were not centrally re-reviewed.

Statistical analyses

Dichotomous data were presented as proportions. Continuous variables were reported as median with interguartile range (IQR). Variable distribution was assessed by plotting histograms. Categorical variables were expressed as number and percentage. Variables were analyzed using appropriate statistical tests for variable type and distribution. Multivariate analysis using logistic regression was performed, with reporting of odds ratio (OR), adjusted OR (aOR), and 95% confidence interval (95% CI). Covariates were included if $\rho < 0.10$ after univariate analysis and were corrected for interaction when necessary. Parameters with twotailed p < 0.05 were considered statistically significant. All analyses were performed in R version 4.1.0. (R Core Team, 2021; R Foundation for Statistical Computing, Vienna, Austria). Study data were collected and managed using REDCap electronic data capture tools hosted at the UMCG.[19,20]

RESULTS

Baseline characteristics

A total of 222 patients who underwent surgery for (suspected) HCAs were included, of whom 44 (20%) patients had small tumors (< 50 mm) and 178 (80%) patients had large tumors (\geq 50 mm) (Figure S1). Patients with small tumors were older (46 years vs. 38 years, p = 0.016) and more frequently men (20%) vs. 5.1%, p = 0.002) than patients with large tumors. In both groups, 33% of patients had hepatic steatosis. Median tumor diameter was 30 (21-40) mm in patients with small tumors compared to 83 (64–110) mm in patients with large tumors (Table 1; p < 0.001). Bilobar presence of tumors was approximately 35%-40% in both groups. The number of tumors was comparable in both groups, with 60% of patients diagnosed with one tumor and 20% with two tumors (Table 1).

Overall use of preoperative magnetic resonance imaging (MRI) was similar between the two groups (86% vs. 92%, p=0.37), although hepatobiliary contrast enhanced MRI (CE-MRI) was less often used in patients with small tumors (58% vs. 71%, p=0.008). Preoperative histopathology was obtained in five (11%) patients with small tumors compared to 42 (24%) patients with large tumors (p=0.12; Table 1).

Discussion of the indication for surgery in a multidisciplinary team meeting occurred in 91% of patients with small tumors and in 89% of patients with large tumors (p = 0.97). A multidisciplinary team meeting, however, was more often consulted in tertiary referral centers (94%) than in regional hospitals (75%; p < 0.001) before surgery.

Indications for surgery

Indications for resection differed between patients with small and large tumors (p < 0.001). In patients with small tumors <50 mm, the most common indication for resection (55%) was suspicion of (pre)malignancy (either on imaging or on histopathological analyses) (Figure 1A). Other indications for resection of small tumors were (previous) tumor hemorrhage (14%), male sex (11%), pregnancy wish (4.5%), tumor growth (4.5%), and patient uncertainty (4.5%) (Table S1). The main indication for resection of large tumors \geq 50 mm was tumor size (52%), followed by suspicion of (pre)malignancy (27%), histopathological features of beta-catenin-mutated HCAs (b-HCAs; 5.6%), and (previous) hemorrhage (5.1%) (Figure 1B). Other reasons were abdominal complaints (i.e., pain, bloating, or tiredness), exophytic tumor growth, HCA-induced amyloidosis, and HCA-induced anemia. In regional hospitals, more HCAs were resected because of previous hemorrhage (12% vs. 5.3%) or male sex (12% vs. 1.2%) when compared to tertiary referral centers (p = 0.004) (Table 2). All patients receiving an operation due to male sex received an MRI, and no difference in MR contrast was observed (p = 0.11).

Preoperative histopathological and imaging characteristics

A total of 47 patients (21%) had undergone preoperative biopsy, more often in large tumors (42/178) than in small tumors (5/44), albeit not significantly (p=0.12; Table 1). Eventually, 11 patients (23%) with a preoperative biopsy underwent resection because of cellular atypia (Table S1). At final pathology, 10 of those were diagnosed as HCAs and one as FNH. Sixty-one out of 175 (35%) patients without preoperative biopsy were operated on because of suspected (pre)malignancy on MRI. Of those, 34 patients (56%) had undergone a preoperative CE-MRI with liver-specific contrast agent (7/20 [44%] patients with tumors <50 mm and 27/41 [66%] patients with tumors >50 mm [p=0.22]).

Final histopathological outcomes and risk factors for change in diagnosis

At final pathology, no differences were observed for HCA subtypes between tumor size groups (Table 3). However, FNH was diagnosed in 24 patients, comprising 21% of the smaller tumors vs. 8.5% of the resected larger tumors (p = 0.11). Of all 24 patients with FNH at final pathology, 22 patients (92%) had undergone MRI in the preoperative workup, and in 14 patients (64%), a liver-specific contrast agent was administered. In the total cohort, use of hepatobiliary CE-MRI was similar for male and female patients (p = 0.10). In patients with tumors <50 mm,

TABLE 1 Baseline characteristics of patients with (suspected) HCA, stratified by tumor diameter

Characteristic	Tumor < 50 mm	n (n = 44)	Tumor ≥ 50 mm	(n = 178)	p value ^a
Female sex	35	(80)	169	(95)	0.002
Age at surgery (years)	46	(33–56)	38	(31–46)	0.016
Body mass index (kg/m2)	27	(22–32)	28	(24–32)	0.68
Charlson Comorbidity Index					0.70
CCI 0/1	43	(98)	169	(95)	
CCI ≥ 2	1	(2)	9	(5)	
American Society of Anesthesiology					0.013
ASA I/II	33	(75)	162	(91)	
$ASA \ge III$	10	(23)	15	(8.4)	
Missing	1	(2.3)	1	(7.9)	
Preoperative MRI	38	(86)	164	(92)	0.37
MRI contrast agent					0.008
Liver-specific contrast agent	7	(16)	36	(20)	
Extracellular contrast agent	22	(50)	115	(65)	
No contrast administered	9	(21)	11	(6.2)	
Missing	6	(14)	16	(9.0)	
Number of tumors					0.52
1 tumor	26	(59)	107	(60)	
2–5 tumors	9	(21)	36	(20)	
6–9 tumors	2	(4.5)	9	(5.1)	
≥ 10 tumors	2	(4.5)	2	(1.1)	
Missing	5	(11)	24	(14)	
Diameter of largest tumor (mm)	30	(21–40)	83	(64–110)	< 0.001
Bilobar tumor occurrence	15	(34)	74	(42)	0.43

Note: Data show number (%). Continuous values are provided as median and interquartile range.

Abbreviations: ASA, American Society of Anesthesiology; CCI, Charlson Comorbidity Index; FNH, focal nodular hyperplasia; HCA, hepatocellular adenoma; HCC, hepatocellular carcinoma; MRI, magnetic resonance imaging.

*p < 0.05 is significant.

hepatobiliary CE-MRI was used in the diagnostic workup of three men (33%) and in 19 women (54%) (p=0.46). At final pathology, 6/9 male patients were diagnosed with FNH compared to 3/35 female patients; p<0.001. From the six male patients with resected FNH <50 mm, none had preoperative histopathology analyzed, and indications for resection were suspicion of (pre)malignancy on MRI (n=3) and because of male sex with HCA suspicion (n=3). Three resected tumors turned out to be HCCs, all in female patients with tumors ≥ 50 mm (Figure 1B). Indication for resection of two HCCs was because of tumor size ≥ 50 mm (with co-occurring tumor growth), and one HCC was resected because of suspicion of (pre)malignancy.

Analysis of significant changes in preoperative and postoperative diagnosis revealed 96 (43%) changed diagnoses (Figure 2). A change in diagnosis was observed in 14 (78%) male patients compared to 82 (40%) female patients (p < 0.01), in 31 (70%) small tumors compared to 65 (37%) large tumors (p < 0.001), and in 46 (54%) patients without preoperative hepatobiliary CE-MRI compared to patients with hepatobiliary

CE-MRI available (p < 0.05). No differences were seen between patients with or without MRI or with or without percutaneous biopsy. These observations were similar in univariate logistic regression, which demonstrated an increased risk of diagnostic change in male patients (OR, 5.2; 95% CI, 1.8–18.9; p < 0.01), in tumors <50 mm (OR, 4.1; 95% CI, 2.1–8.7; p < 0.001), and in patients without hepatobiliary CE-MRI (OR, 2.1; 95% CI, 1.2–3.6; p < 0.05). Use of MRI regardless of use of contrast (type) did not influence diagnostic change (p = 0.44) and neither did use of percutaneous biopsy (p = 0.27). A model was constructed including sex, tumor size category, and use of hepatobiliary CE-MRI (Figure 3). All included variables proved independent risk factors for change in diagnosis: tumors <50 mm (aOR, 3.4; 95% CI, 1.7–7.4; p < 0.01), male sex (aOR, 3.7; 95% CI, 1.2–13.8; p = 0.03), and lack of hepatobiliary CE-MRI (aOR, 1.8; 95% CI, 1.0-3.3; p = 0.04). Influence of a sex-tumor size category interaction was explored, including the three aforementioned variables, but did not improve the model (aOR, 2.6; 95% CI, 0.2–66.3; p = 0.48).

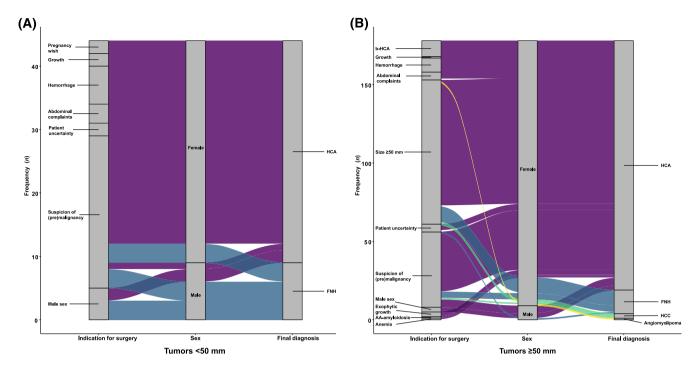


FIGURE 1 Alluvial plots depicting the individual distribution of indication for resection in patients with (A) HCAs < 50 mm or (B) HCAs ≥ 50 mm, grouped by sex and final diagnosis. AA-amyloidosis, amyloid A amyloidosis; b-HCA, beta-catenin mutated hepatocellular adenoma; FNH, focal nodular hyperplasia; HCA, hepatocellular adenoma; HCC, hepatocellular carcinoma.

Surgical outcomes

No difference in frequency of surgery was observed throughout the years. During the inclusion period, there was a trend toward more frequent laparoscopic resections in patients with smaller tumors (67% vs. 53%; p=0.13). Major resections (n=56) were more often performed in patients with large tumors. Sixteen (29%) major resections were performed through laparoscopy. Postoperative outcomes were similar for patients who underwent surgery for small or large tumors, with 30-day major morbidity <3% and 30-day mortality <1% (Table 3).

DISCUSSION

In this study, indications for resection of small (suspected) HCAs compared to large HCAs were investigated in a nationwide cohort. The study included 222 patients, of

whom 44 patients (20%) underwent surgery for small tumors (<50 mm). Half of patients with small tumors were operated on because of suspicion of (pre)malignancy, and the remaining patients mainly underwent surgery because of (previous) hemorrhage or male sex; for patients with large tumors, the most prevalent indication was tumor size itself. A logistic regression model showed that male sex (aOR, 3.7), small tumor size (aOR, 3.4), and lack of hepatobiliary CE-MRI (aOR, 1.8) were independent risk factors for a postoperative change in diagnostic.

The diagnostic process for benign liver tumors is complex because of the distinct clinical and risk profiles between and within benign liver tumors in often relatively young and healthy patients. When it comes to decision making to proceed to surgery, some indications are stronger because of a clear tradeoff between the benefit and (potential) harm of surgery. Risk of malignant transformation is such an indication, which is reflected in the observed indications in our

TABLE 2 Grouped indications for surgery and postoperative diagnosis of patients with hepatocellular adenoma, stratified for hospital type

Characteristic	Regional he	Regional hospital $(n = 52)$		Tertiary referral hospital $(n = 170)$	
Indication of surgery, n (%)					0.004**
Hemorrhage (old or new)	6	(12)	9	(5.3)	
Abdominal complaints	1	(1.9)	7	(4.1)	
Size ≥ 50 mm	22	(42)	70	(41)	
Atypia tumor	15	(29)	67	(39)	
Male sex	6	(12)	2	(1.2)	
Other	2	(3.8)	15	(8.8)	

^{**}p < 0.01.

TABLE 3 Operative characteristics and final histopathology of patients with (suspected) HCA, stratified for tumor diameter

Characteristic	Tumors < 50 mm (n = 44)		Tumors ≥ 50 mm (n = 178)		p value ^a
Final histopathology					
Tumor diagnosis				0.11	
HCA	35	(80)	159	(89)	
FNH	9	(20)	15	(8.4)	
HCC	0	(-)	3	(1.7)	
Angiomyolipoma	0	(-)	1	(0.6)	
HCA subtype		()		0.13	
I-HCA	18	(41)	99	(56)	
H-HCA	5	(11)	14	(7.9)	
b-HCA / b-IHCA	0	(-)	11	(6.2)	
U-HCA	2	(4.5)	8	(4.5)	
No subtype analyses performed	10	(23)	27	(15)	
Alternate tumor than HCA	9	(21)	19	(11)	
Postoperative change in tumor diagnosis	31	(71)	65	(37)	< 0.001
Operative characteristics and outcomes				0.00	
Year of surgery	0	(4.0)	20	0.20	
2014	8	(18)	28	(16)	
2015	9	(21)	27	(15)	
2016	4 12	(9.1)	36	(20)	
2017		(27)	27	(15)	
2018 2019	5 6	(11)	34 26	(19)	
Type of resection	O	(14)	20	(15)	0.042
	11	(25)	29	(16)	0.042
Wedge resection	28	(25)	99	(16)	
Segment resection	0	(64)	6	(56)	
Left hemihepatectomy Right hemihepatectomy	1	(-) (2.3)	38	(3.4) (21)	
Extended left hemihepatectomy	0	(-)	1	(0.6)	
Extended right hemihepatectomy	0	(-)	1	(0.6)	
Missing	4	(-) (9.1)	4	(2.2)	
Extensiveness of resection	7	(3.1)	4	(2.2)	< 0.001
Minor resection	43	(98)	123	(69)	₹ 0.001
Major resection	1	(2)	55	(31)	
Surgical approach	•	(-)	30	0.13	
Open	14	(32)	83	(47)	
Laparoscopic	29	(66)	94	(53)	
Missing	1	(2.3)	1	(0.6)	
30-day major morbidity ^b	1	(2)	3	(1.7)	1.00
30-day mortality	0	(-)	1	(0.6)	1.00

Note: Data show number (%); continuous values are provided as median and interquartile range.

Abbreviations: b-HCA, beta-catenin mutated HCA; b-IHCA, hybrid b-HCA; FNH, focal nodular hyperplasia; HCA, hepatocellular adenoma; HCC, hepatocellular carcinoma; H-HCA, hepatocyte nuclear factor 1A inactivated HCA; I-HCA, inflammatory HCA; U-HCA, unclassified HCA.

ap < 0.05 is significant.

bDefined as Clavien-Dindo score ≥ 3a.

study cohort. In half of patients with small tumors, surgery was performed because of suspicion of (pre) malignancy on imaging or histopathology and the same holds true for almost a third of patients with large tumors. All 18 male patients in our cohort were operated

on because of their sex in combination with (suspicion of) HCA diagnosis. Male sex is an independent risk factor for malignant transformation, and the premalignant b-HCAs occur more often in men, which may justify the indication for resection.^[3,4,6] Male

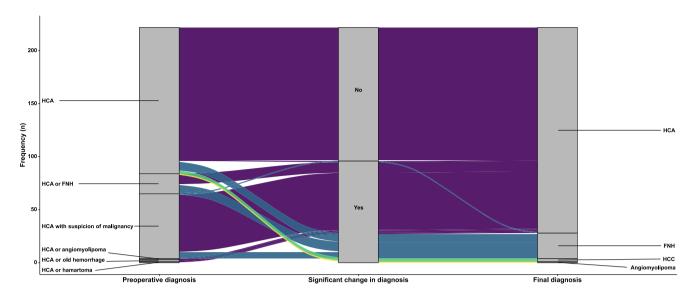


FIGURE 2 Alluvial plot depicting the change in tumor diagnosis before and after resection in the total cohort. FNH, focal nodular hyperplasia; HCA, hepatocellular adenoma; HCC, hepatocellular carcinoma.

patients with HCAs due to metabolic disease, like hepatocyte nuclear factor 1A (HNF1A) maturity-onset diabetes of the young (HNF1A-MODY) or glycogen storage disease, might be exceptions, but future research is needed for definite answers regarding oncologic safety.[21,22] HNF1A-MODY should especially be considered if multiple HNF1A-inactivated HCAs (H-HCAs), which can be preceded by diabetic symptoms, are observed in a (male) patient.[21] Because H-HCAs generally demonstrate very limited risk of bleeding or malignant transformation, a conservative approach with follow-up imaging may be warranted, also in male patients. Future research on oncologic safety is needed for definite conclusions as HNF1A mutations are rare but have been observed in 1.5% of resected HCCs, and a family with HNF1A-MODY and H-HCA-induced primary hepatic malignancies has been reported.[23,24]

Relative indications, such as abdominal complaints or patient uncertainty, were observed in only five (11%) patients with small tumors and in 10 (5.6%) patients with large tumors. It is assumed that the severity of symptoms led to the indication for resection; however, in the absence of well-developed QoL instruments for

patients with benign liver tumors, it remains difficult to assess the burden of disease.

Prevention of hepatic hemorrhage by tumor rupture, which is size dependent and is low in tumors <50 mm, is another important indication for invasive tumor treatment.[5,6] Tumor size was the deciding factor in half of patients with large tumors > 50 mm. Risk of HCA bleeding is especially increased in exophytic HCAs. [5,25] In our series, exophytic growth was an indication for only three cases, whereas previous hemorrhage was an indication in 15 cases. European guidelines do not consider previous tumor hemorrhage as an absolute indication for surgery, although in rare cases, such as bleeding in exophytic HCAs, previous hemorrhage can be a relative indication for intervention. Before deciding on HCA treatment after bleeding, HCAs should first be observed for posthemorrhagic necrosis-induced regression, which might remove the need for tumor resection.[25] Finally, current European guidelines recommend HCA resection if size remains >50 mm 6 months after lifestyle alterations (i.e., ceasing oral contraceptive pill use and weight loss). However, 6 months might be a too short interval for large HCAs

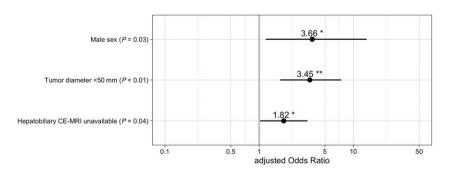


FIGURE 3 Forest plot of logistic regression analysis on risk factors for change in tumor diagnosis. CE-MRI, contrast enhanced magnetic resonance imaging. *p < 0.05, **p < 0.01.

to regress sufficiently, and data suggest that watchful waiting can be prolonged safely.^[9,26]

The clinical decision process is dependent on the (suspected) preoperative tumor diagnosis. A postoperative change in tumor diagnosis was observed frequently (43%). Although amounting for only a small number of total cases, preoperative diagnosis was altered from HCA or (pre)malignancy to FNH in six out of the nine male patients with tumors <50 mm. A therapeutically defensive approach by resecting tumors not distinguishable between HCAs or FNH in male patients is conceivable; however, preoperative diagnostic workup should be adequate. Our findings that especially male sex, small tumors, and lack of hepatobiliary CE-MRI were independently associated with increased risk of change in diagnosis may highlight the need to improve the diagnostic process to prevent unnecessary hepatic surgery. We propose to perform a hepatobiliary CE-MRI in all patients with (suspected) HCAs, and to perform a percutaneous biopsy whenever there is doubt on tumor diagnosis after this imaging. Furthermore, all (suspected) HCAs in male patients should be confirmed through percutaneous liver biopsy. preferably through histopathological molecular analysis due to higher diagnostic sensitivity.[27] A suspicion of malignant transformation after hepatobiliary CE-MRI needs to be confirmed by liver biopsy, especially in tumors <50 mm and in male patients. In our opinion, potential complications and morbidity following unnecessary surgery outweigh the limited risk of biopsy-induced bleeding. Molecular analysis is more able to diagnose b-HCA (catenin beta 1 gene CTNNB1) and should supplement immunohistochemistry whenever a beta-catenin mutation is suspected.[27] Molecular analysis also allows for differentiation between exon 3 and exon 7/8 CTNNB1 HCA mutations, the latter having less malignant potential, although exon 7/8 mutated HCAs transforming into HCCs have been observed. [6,28]

Of note, percutaneous tumor ablation is minimally invasive, effective, and safe in treating hepatic malignancies and HCAs and could be performed in the same session directly after the histologic biopsy in tumors <50 mm.^[29,30]

A limitation of the current study is the retrospective assessment of preoperative diagnostic workup, including imaging, as the radiologic analysis often contains many nuances open to varying interpretations. In addition, the current data do not allow for analysis of indications per HCA subtype, which may have been potentially insightful and could be analyzed in future studies. Another potential limitation is the accuracy and coverage of the included registry data. Third-party data verification has deemed 97% of the data accurate, yet not all specific information concerning operative outcomes could be obtained.^[16] Because the current study reflects the historical decision-making process from 2014 to 2019 in a nationwide cohort,

substantial improvements in diagnostic workup have been made since, including identification of new HCA subtypes. For example, sonic hedgehog-mutated HCAs (sh-HCAs) have been discovered in recent years, which represent 4% of HCAs.^[6,31] Sh-HCAs are especially at increased risk of tumor bleeding.^[6] Unfortunately, immunohistochemical staining of argininosuccinate synthetase 1 or molecular characterization of inhibin beta E chain with GLI family zinc finger 1 (GLI1) was not routine practice during the study period.^[6,31,32]

Future studies on preoperative modality and final tumor diagnosis are needed to uncover potential areas of improvement of care. Although some extent of diagnostic uncertainty occurs in every modality, suboptimal use of imaging modalities or radiologic contrast agents might lead to unnecessary diagnostic ambiguity. Second, the "relative" indications for resection of (suspected) HCAs, like impact on QoL by psychological burden or abdominal complaints, should be further explored. This necessitates both development and validation of QoL tools or patient-related outcome instruments specifically for patients with benign liver tumors as well as analysis of these potential consequences in a cohort.

In conclusion, surgery for small HCAs was mainly indicated by suspicion of (pre)malignancy, whereas for large (suspected) HCAs, tumor size was the most prevalent indication. Male sex, tumor size <50 mm, and lack of hepatobiliary CE-MRI were independent risk factors for postoperative change in tumor diagnosis. Future studies should focus on evaluation of preoperative diagnostics as well as exploration of QoL-related indications, such as patient uncertainty or abdominal complaints.

AUTHOR CONTRIBUTIONS

Martijn P. D. Haring: manuscript and figure drafting, data collection, data analysis, study design. Arthur K. E. Elfrink: study design, data analysis, review of manuscript. Christiaan A. J. Oudmaijer, Paul C. M. Andel, Alicia Furumaya, Nenke de Jong, Colin J. J. M. Willems, Thijs Huits, Julie M. L. Sijmons, Joris I. Erdmann, Paul Gobardhan, Robbert J. de Haas, Tjarda van Heek, Hwai-Ding Lam, Wouter K. G. Leclercq, Mike S. L. Liem, Hendrik A. Marsman, Gijs A. Patijn, Türkan Terkivatan, Babs M. Zonderhuis, Izaak Quintus Molenaar, Wouter W. te Riele, Jeroen Hagendoorn, Alexander F. M. Schaapherder, Jan N. M. IJzermans, Carlijn I. Buis, Joost M. Klaase, Koert P. de Jong: study design, data collection, review of manuscript; Vincent E. de Meijer: study design, figure data analysis supervision, drafting, review of manuscript.

CONFLICT OF INTEREST

Nothing to report.

DATA AVAILABILITY STATEMENT

The data that support the findings of this study are available from the corresponding author on reasonable request.

ETHICAL APPROVAL

Ethical approval was obtained from the Medical Ethical Committee in the UMCG (MEC 2020-004 & UMCG Research Registry 201900849), and confimed all other partipating centers. All data was collected and processed according to relevant privacy laws.

ORCID

Martijn P. D. Haring https://orcid.org/0000-0003-4789-3910

Arthur K. E. Elfrink https://orcid.org/0000-0003-2332-2849

Christiaan A. J. Oudmaijer https://orcid.org/0000-0003-4852-6514

Alicia Furumaya https://orcid.org/0000-0001-5897-0438

Julie M. L. Sijmons https://orcid.org/0000-0002-4914-4537

Eric J. T. Belt https://orcid.org/0000-0002-1211-369X

Esther C. J. Consten https://orcid.org/0000-0002-9447-8181

Mariëlle M. E. Coolsen https://orcid.org/0000-0002-1608-6668

Peter van Duijvendijk https://orcid.org/0000-0002-0441-6180

Joris I. Erdmann https://orcid.org/0000-0001-8233-1505

Paul Gobardhan https://orcid.org/0000-0001-5888-0483

Robbert J. de Haas https://orcid.org/0000-0002-4915-6781

Hwai-Ding Lam https://orcid.org/0000-0002-4235-6916

Mike S. L. Liem https://orcid.org/0000-0001-8665-

Gijs A. Patijn https://orcid.org/0000-0001-8113-4171

Türkan Terkivatan https://orcid.org/0000-0002-2541-1530

Izaak Quintus Molenaar https://orcid.org/0000-0002-1585-7184

Wouter W. te Riele https://orcid.org/0000-0002-5256-3919

Jeroen Hagendoorn https://orcid.org/0000-0001-8737-3923

Alexander F. M. Schaapherder https://orcid.org/0000-0003-3262-9041

Jan N. M. IJzermans https://orcid.org/0000-0003-3558-1039

Carlijn I. Buis https://orcid.org/0000-0002-2379-3723

Joost M. Klaase https://orcid.org/0000-0002-4409-2280

Koert P. de Jong https://orcid.org/0000-0001-6345-653X

Vincent E. de Meijer https://orcid.org/0000-0002-7900-5917

REFERENCES

- Edmondson HA, Henderson B, Benton B. Liver-cell adenomas associated with use of oral contraceptives. N Eng J Med. 1976; 294:470–2.
- Gevers TJG, Marcel Spanier BW, Veendrick PB, Vrolijk JM. Regression of hepatocellular adenoma after bariatric surgery in severe obese patients. Liver Int. 2018;38:2134–6.
- Farges O, Ferreira N, Dokmak S, Belghiti J, Bedossa P, Paradis V. Changing trends in malignant transformation of hepatocellular adenoma. Gut. 2011;60:85–9.
- Stoot JHMB, Coelen RJS, de Jong MC, Dejong CHC. Malignant transformation of hepatocellular adenomas into hepatocellular carcinomas: a systematic review including more than 1600 adenoma cases. HPB (Oxford). 2010;12:509–22.
- Bieze M, Phoa SSKS, Verheij J, van Lienden KP, van Gulik TM. Risk factors for bleeding in hepatocellular adenoma. Br J Surg. 2014;101:847–55.
- Nault JC, Couchy G, Balabaud C, Morcrette G, Caruso S, Blanc JF, et al. Molecular classification of hepatocellular adenoma associates with risk factors, bleeding, and malignant transformation. Gastroenterology. 2017;152:880–94.e6.
- European Association for the Study of the Liver (EASL). EASL clinical practice guidelines on the management of benign liver tumours. J Hepatol. 2016;65:386–98.
- Marrero JA, Ahn J, Reddy KR, Americal College of Gastroenterology. ACG clinical guideline: the diagnosis and management of focal liver lesions. Am J Gastroenterol. 2014;109: 1328–47.
- Haring MPD, Gouw ASH, de Haas RJ, Cuperus FJC, de Jong KP, de Meijer VE. The effect of oral contraceptive pill cessation on hepatocellular adenoma diameter: a retrospective cohort study. Liver Int. 2019;39:905–13.
- Haring MPD, Cuperus FJC, Duiker EW, de Haas RJ, de Meijer VE. Scoping review of clinical practice guidelines on the management of benign liver tumours. BMJ Open Gastroenterol. 2021;8:e000592.
- Strauss E, Ferreira ASP, França AVC, Lyra AC, Barros FMR, Silva I, et al. Diagnosis and treatment of benign liver nodules: Brazilian Society of Hepatology (SBH) recommendations. Arq Gastroenterol. 2015;52(Suppl. 1):47–54.
- van Rosmalen B v., De Graeff JJ, van der Poel MJ, De Man IE, Besselink M, Abu Hilal M, et al; Dutch Benign Liver Tumour Group. Impact of open and minimally invasive resection of symptomatic solid benign liver tumours on symptoms and quality of life: a systematic review. HPB (Oxford). 2019;21:1119–30.
- Dokmak S, Paradis V, Vilgrain V, Sauvanet A, Farges O, Valla D, et al. A single-center surgical experience of 122 patients with single and multiple hepatocellular adenomas. Gastroenterology. 2009;137:1698–705.

 Hoffmann K, Unsinn M, Hinz U, Weiss KH, Waldburger N, Longerich T, et al. Outcome after a liver resection of benign lesions. HPB (Oxford). 2015;17:994–1000.

- Elfrink AKE, Haring MPD, de Meijer VE, IJzermans JNM, Swijnenburg R-J, Braat AE, et al. Surgical outcomes of laparoscopic and open resection of benign liver tumours in the Netherlands: a nationwide analysis. HPB (Oxford). 2021;23: 1230–43
- van der Werf LR, Kok NFM, Buis CI, Grünhagen DJ, Hoogwater FJH, Swijnenburg RJ, et al. Implementation and first results of a mandatory, nationwide audit on liver surgery. HPB (Oxford). 2019:21:1400–0.
- von Elm E, Altman DG, Egger M, Pocock SJ, G

 øtzsche PC, Vandenbroucke JP, et al. The Strengthening the Reporting of Observational Studies in Epidemiology (STROBE) statement: guidelines for reporting observational studies. J Clin Epidemiol. 2008:61:344–9.
- Dindo D, Demartines N, Clavien PA. Classification of surgical complications: a new proposal with evaluation in a cohort of 6336 patients and results of a survey. Ann Surg. 2004;240:205–13.
- Harris PA, Taylor R, Minor BL, Elliott V, Fernandez M, O'Neal L, et al. The REDCap consortium: building an international community of software platform partners. J Biomed Inform. 2019;95:103208.
- Harris PA, Taylor R, Thielke R, Payne J, Gonzalez N, Conde JG. Research electronic data capture (REDCap)—a metadata-driven methodology and workflow process for providing translational research informatics support. J Biomed Inform. 2009;42:377–81.
- Haring MPD, Vriesendorp TM, Klein Wassink-Ruiter JS, de Haas RJ, Gouw ASH, de Meijer VE. Diagnosis of hepatocellular adenoma in men before onset of diabetes in HNF1A-MODY: watch out for winkers. Liver Int. 2019;39:2042–5.
- Haring MPD, Peeks F, Oosterveer MH, Brouwers MCGJ, Hollak CEM, Janssen MCH, et al. High childhood serum triglyceride concentrations associate with hepatocellular adenoma development in patients with glycogen storage disease type Ia. JHEP Rep. 2022;4:100512.
- Willson JSB, Godwin TD, Wiggins GAR, Guilford PJ, McCall JL. Primary hepatocellular neoplasms in a MODY3 family with a novel HNF1A germline mutation. J Hepatol. 2013;59:904–7.
- Hechtman JF, Abou-Alfa GK, Stadler ZK, Mandelker DL, Roehrl MHA, Zehir A, et al. Somatic HNF1A mutations in the malignant transformation of hepatocellular adenomas: a retrospective analysis of data from MSK-IMPACT and TCGA. Hum Pathol. 2019;83:1–6.
- Klompenhouwer AJ, de Man RA, Thomeer MG, IJzermans JN. Management and outcome of hepatocellular adenoma with massive bleeding at presentation. World J Gastroenterol. 2017;23:4579–86.
- Klompenhouwer AJ, Bröker MEE, Thomeer MGJ, Gaspersz MP, de Man RA, IJzermans JNM. Retrospective study on timing of resection of hepatocellular adenoma. Br J Surg. 2017;104:1695–703.
- van Rosmalen BV, Furumaya A, Klompenhouwer AJ, Tushuizen ME, Braat AE, Reinten RJ, et al. Hepatocellular adenoma in men: a nationwide assessment of pathology and correlation with clinical course. Liver Int. 2021;41:2474–84.
- Klompenhouwer AJ, Thomeer MGJ, Dinjens WNM, de Man RA, IJzermans JNM, Doukas M. Phenotype or genotype: decisionmaking dilemmas in hepatocellular adenoma. Hepatology. 2019; 70:1866–8.
- Mironov O, Jaberi A, Beecroft R, Kachura JR. Retrospective single-arm cohort study of patients with hepatocellular adenomas treated with percutaneous thermal ablation. Cardiovasc Intervent Radiol. 2018;41:935–41.
- Hof J, Wertenbroek MWJLAE, Peeters PMJG, Widder J, Sieders E, de Jong KP. Outcomes after resection and/or radiofrequency ablation for recurrence after treatment of colorectal liver metastases. Br J Surg. 2016;103:1055–62.
- 31. Sala M, Gonzales D, Leste-Lasserre T, Dugot-Senant N, Paradis V, di Tommaso S, et al. ASS1 overexpression: a hallmark of

- sonic hedgehog hepatocellular adenomas; recommendations for clinical practice. Hepatol Commun. 2020;4:809–24.
- Nault JC, Couchy G, Caruso S, Meunier L, Caruana L, Letouzé E, et al. Argininosuccinate synthase 1 and periportal gene expression in sonic hedgehog hepatocellular adenomas. Hepatology. 2018;68:964–76.

How to cite this article: Haring MPD, Elfrink AKE, Oudmaijer CAJ, Andel PCM, Furumaya A, de Jong N, et al. the Dutch Benign Liver Tumor GroupA nationwide assessment of hepatocellular adenoma resection: Indications and pathological discordance. Hepatol Commun. 2023;7:e2110. 10.1002/hep4.2110

APPENDIX

Dutch Benign Liver Tumor Group: R. B. Takkenberg, Department of Gastroenterology and Hepatology, Amsterdam University Medical Center, location AMC. Amsterdam, the Netherlands: J. Verheii. Department of Medical Biology and Pathology, Amsterdam University Medical Center, location AMC, Amsterdam, the Netherlands; O. M. van Delden, Department of Radiology, Amsterdam University Medical Center, location AMC, Amsterdam, the Netherlands; J. I. Erdmann, B. V. van Rosmalen, G. Kazemier, and A. Furumaya, Department of Surgery, Amsterdam University Medical Center, location AMC, Amsterdam, the Netherlands; D. Ramsoekh, Department of Gastroenterology and Hepatology, Amsterdam University Medical Center, location VUMC, Amsterdam, the Netherlands; J. N. M. IJzermans, Department of Surgery, Erasmus Medical Center, Rotterdam, the Netherlands; R. A. de Man, Department of Gastroenterology and Hepatology, Erasmus Medical Center, Rotterdam, the Netherlands; M. D. Doukas, Department of Medical Biology and Pathology, Erasmus Medical Center, Rotterdam, the Netherlands; M. G. Thomeer, Department of Radiology, Erasmus Medical Center, Rotterdam, the Netherlands; A. J. Klompenhouwer, Department of Gastroenterology and Hepatology, Erasmus Medical Center, Rotterdam, the Netherlands; A. F. M. Schaapherder, Department of Surgery, Leiden University Medical Center, Leiden, the Netherlands; M. E. Tushuizen and M. J. Coenraad, Department of Gastroenterology and Hepatology, Leiden University Medical Center, Leiden, the Netherlands; A. S. L. P. Crobach, Department of Medical Biology and Pathology, Leiden University Medical Center, Leiden, the Netherlands; M. C. Burgmans, and S. Feshtali Department of Radiology, Leiden University Medical Center, Leiden, the Netherlands; M. M. E. Coolsen, Department of Surgery, Maastricht University Medical Center+,

Maastricht, the Netherlands; M. Kramer, Department of Gastroenterology and Hepatology, Maastricht University Medical Center+, Maastricht, the Netherlands; J. C. Beckervordersandforth and I. V. Samarska, Department of Medical Biology and Pathology, Maastricht University Medical Center+, Maastricht, the Netherlands; C. van der Leij and R. Miclea, Department of Radiology, Maastricht University Medical Center+, Maastricht, the Netherlands; J. H. W. de Wilt and P. B. van den Boezem, Department of Surgery, Radboudumc, Nijmegen, the Netherlands; E. T. T. L. Tjwa and I. Munsterman, Department of Gastroenterology and Hepatology, Radboudumc,

Nijmegen, the Netherlands; S. Vos, Department of Medical Biology and Pathology, Radboudumc, Nijmegen, the Netherlands; S. van Koeverden, Department of Radiology, Radboudumc, Nijmegen, the Netherlands; V. E. de Meijer and M. P. D. Haring, Department of Surgery, University Medical Center Groningen, Groningen, the Netherlands; F. J. C. Cuperus and R. J. de Haas, Department of Gastroenterology and Hepatology, University Medical Center Groningen, Groningen, the Netherlands; E. W. Duiker, Department of Medical Biology and Pathology, University Medical Center Groningen, Groningen, the Netherlands.