



# Scirrhou Hepatocellular Carcinoma: Systematic Review and Pooled Data Analysis of Clinical, Radiological, and Histopathological Features


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**Background:** Aberrant subtypes of hepatocellular carcinoma (HCC) account for 20–30% of all HCCs and habitually present a challenge in diagnosis and treatment. Scirrhou hepatocellular carcinoma (s-HCC) is often misdiagnosed as cholangiocarcinoma, fibrolamellar hepatocellular carcinoma, or metastasis.

**Methods:** Electronic databases (PubMed, Web of Knowledge, Google Scholar, Cochrane Library, and WHO International Clinical Trials Registry Platform) were searched for publications on scirrhou hepatocellular carcinoma without date or language restrictions. Quality assessment was performed using a tool proposed by Murad et al for case reports and series. For observational studies, MINORS quality assessment tool was used. This study was registered at PROSPERO (CRD42020212323).

**Results:** S-HCC arises in patients with chronic hepatitis (hepatitis B in 60% and hepatitis C in 21%). S-HCC primarily affects men with a mean age of 55.8 years. Serum AFP is elevated above 20IU/mL in 66.7% of the patients. On ultrasound, s-HCC presents as hypoechoic or mosaic pattern lesions (47.6% each) and causes a retraction of the liver surface (70%) when near the capsule. Delayed enhancement of the tumor is evident in 87.0%. On MRI, 65.0% of s-HCCs show a target appearance. Histopathologic pattern is mostly irregular (97.6%). Lesions show a bulging appearance (100%), septae (85.6%) and a central scar (63.5%), and usually lack central necrosis (75%). Immunohistochemistry shows HepPar 1 positivity in 64.6%, CK7 in 40.7%, and EMA in 41.9%. The 5-year overall survival rate estimates 45.2% and 45.5% of the patients experience a recurrence after hepatectomy.

**Conclusion:** S-HCC is a rare subtype of HCC primarily arising in hepatitis- or cirrhosis-afflicted livers and incorporates atypical radiological and histopathological HCC features. Despite lower recurrence rates, overall survival of patients with s-HCC is poorer than generally for HCC, underlining the need for individualized treatment. Patients with atypical lesions of the liver should be referred to tertiary hospitals for interdisciplinary assessment and treatment.

**Keywords:** scirrhou HCC, s-HCC, rare liver tumors, RELIVE initiative

## Introduction

Scirrhou hepatocellular carcinoma (s-HCC) is a rare morphologic subtype of hepatocellular carcinoma accounting for an estimated 4% of all HCCs. Only recently has it been recognized as a subtype of HCC in the WHO classification.<sup>1</sup> HCC poses high morbidity and mortality risks, becoming the fourth most common cause of cancer-related mortality.<sup>2</sup> Atypical HCCs comprise about 20–30% of all HCC cases; however, this estimation is based on limited data.<sup>3</sup> Further analysis in order to estimate the

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impact of morphological subtypes is needed to steer HCC treatment forward, towards individual cancer therapy.

S-HCC is often misdiagnosed radiologically as cholangiocarcinoma, fibrolamellar hepatocellular carcinoma (FL-HCC), or even metastasis.<sup>4–7</sup> Yet, for adequate treatment, correct diagnosis is indispensable. Although morphologically there are some similarities between scirrhous and fibrolamellar HCCs, only an adequate histological work-up can lead to the correct diagnosis. S-HCC shows abundant fibrous stroma, which accounts for 30–50% of the tumor mass, concentrated at the center, with nests of densely associated tumor cells.<sup>3</sup> While FL-HCC occurs primarily in young patients without liver disease,<sup>8</sup> s-HCC tends to affect patients with chronic hepatitis or cirrhosis. A crucial aspect of diagnosis is molecular testing, as DNAJB1-PRKACA oncogenic fusion is a sensitive marker for FL-HCC, albeit potentially non-specific.<sup>9</sup> Detailed future clinical characterization of patients may be revealing for the differentiation between s-HCC and other lesions.

It is unclear, how the outcomes of s-HCC compare to the conventional subtype and therefore patients receive the same recommendations despite striking differences in the pathology of the lesions and associated comorbidities. Comparative studies on chemotherapy response in different subtypes of HCCs are also lacking. Comprehensive molecular analysis of each subtype of HCC is necessary to achieve a satisfactory level of expertise and with it structure the targeted approach for adequate patient-oriented treatment. International clinical registries for atypical HCCs could close the informational gap and generate data for an evidence-based approach. The aim of this analysis is to examine available data on clinical, radiological, and histopathological features of s-HCC to summarize the currently available knowledge and guide further research into HCC subtypes.

## Methods

The systematic review is reported in accordance with the current PRISMA guidelines.<sup>10</sup> The review methodology was established before commencement. The protocol of the study was registered prior to data extraction on PROSPERO platform (CRD42020212323).<sup>11</sup>

## Literature Search

A literature search was systematically conducted according to the recent recommendations of the Cochrane Collaboration.<sup>12</sup> The aim of the search was to identify all reports on scirrhous hepatocellular carcinoma. The searches

were performed using databases MEDLINE via PubMed, Web of Knowledge, Google Scholar, Cochrane Library, and WHO International Clinical Trials Registry Platform. An additional hand search was completed using the references of included studies. The last search was performed on February 12, 2021. No language or data restriction was implemented. The details of the search strategy for MEDLINE are provided in the [Supplementary Materials](#).

## Study Selection

No restriction was applied to the study design of the publications as a scarcity of studies on s-HCC was anticipated. All studies describing histological, radiological, or clinical features of s-HCCs were included. Publications such as comments, editorials, meeting abstracts, correspondence, and reviews were excluded. The screening of titles, abstracts, as well as of full texts was carried out by two independent reviewers (AML and JF). All disagreements were resolved by consensus.

## Data Extraction

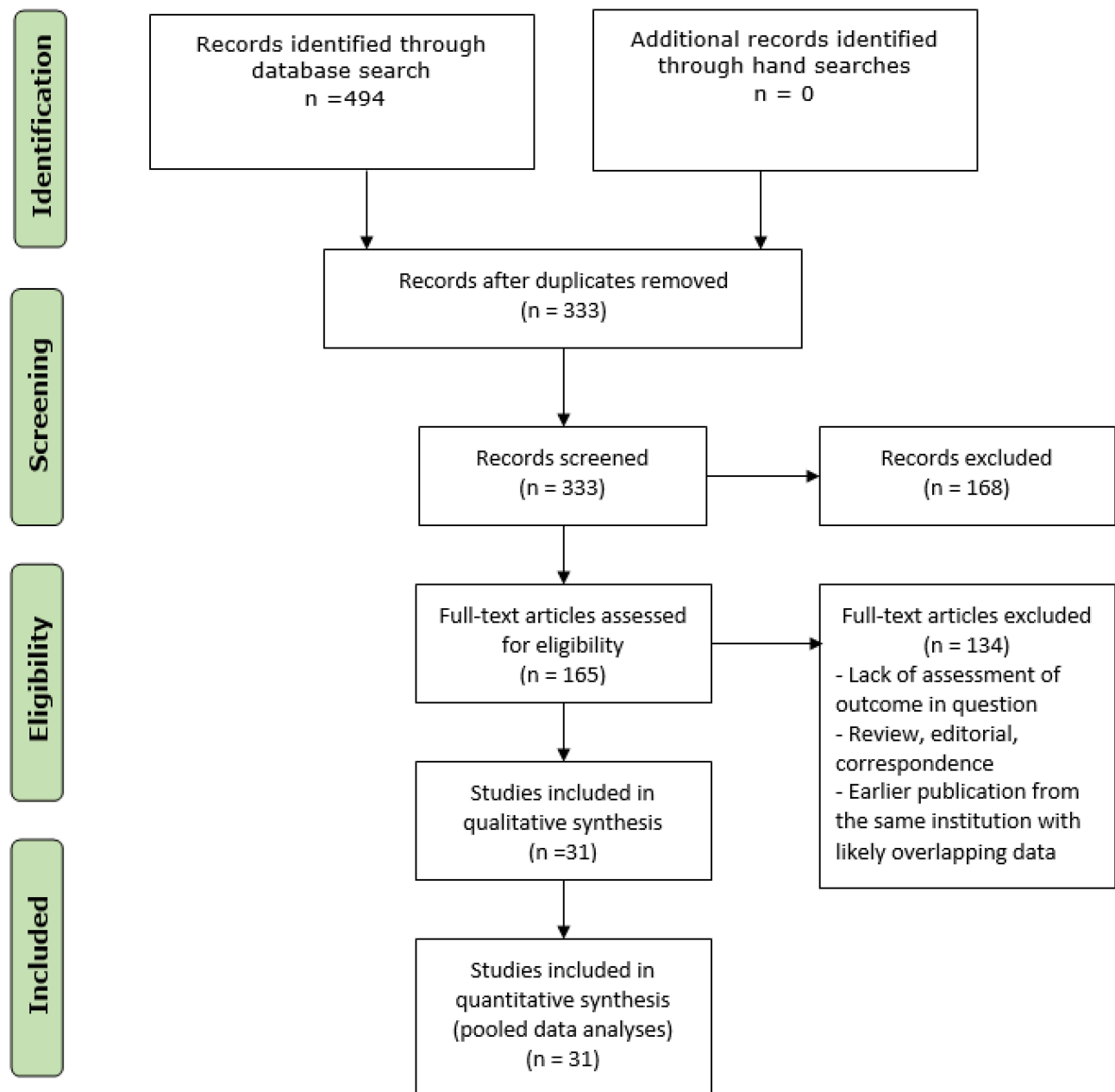
Data extraction of included studies was performed independently by two reviewers (AML and JF) using a standardized form prepared beforehand. Following data were extracted for each study: title of the publication, year, authors, country, journal, source of funding, study design, number in the cohort, patient characteristics, intervention, radiological description of the tumors, histological classifications and properties of the tumors, and clinical outcomes, including follow-up and recurrence rates.

## Statistical Analysis

Pooled data analysis of characteristics involved descriptive statistics and most results were presented as percentages. Continuous variables were presented as means with standard deviation. Statistical analysis was performed using R version 4.0.3.<sup>13</sup>

## Results

A total of 494 records were identified through databases and hand-searches. After exclusion of duplicated, 333 records were reviewed based on title and abstract by two independent reviewers. A total of 165 articles were further assessed by full-text review by two independent reviewers. Thirty-one articles were included in the qualitative synthesis and a total of 31 articles were included in various aspects of quantitative analyses. Figure 1 illustrates the study selection process.



**Diagram 1** Flow chart of the study selection process.

All included publications were of retrospective design with the majority being case reports and reports of series.<sup>4,5,14–41</sup> Due to scarcity of publications from the US and Taiwan, National Registry analyses of 161 and 30 patients, respectively, were included in the analysis.<sup>39,42</sup>

### Critical Appraisal of Included Studies

The methodological quality assessment of all included studies was performed using a tool proposed by Murad et al<sup>43</sup> due to the nature of the review, predominantly case reports and series, lack of a comparison group, and for

comparability of studies. Detailed information about assessed qualities is provided as supplementary material ([Supplementary Table 1](#)). For observational studies, methodological index for non-randomized studies (MINORS)<sup>44</sup> was additionally used to assess risk of bias ([Supplementary Table 2](#)).

### Demographics and Clinical Characteristics

The distribution of available literature on s-HCC shows a certain predominance. Countries largely contributing

articles on this subtype of HCC are Japan and South Korea (Figure 1).

The proportions are somewhat different in population sizes described (Figure 2). Although over 45% of all articles on s-HCC are published from Japanese hospitals, 25% of all patients identified in this review were treated in Japan. Only three publications originated from the US, the largest one, deriving its information from the National Registry, involved 161 patients, making it the second-largest contributor in terms of population.<sup>39</sup>

Cumulatively, South Korea describes the largest population in their publications, with Japan being the third largest contributor.

Mean pooled population age was 55.8 years ( $\pm 9.2$  years), ranging from 24 to 78 years. The pooled cohort consisted of 480 males and 154 females, equivalent to a male:female ratio of 3:1. Only five case reports<sup>14,26,32,38,40</sup> described initial clinical presentations: two patients had no symptoms and presented for hepatitis C follow-up, two patients experienced upper abdominal pain, and one presented with back pain due to metastatic disease. A total of 474 patients had preoperative information on the presence of an underlying liver disease: 59.7% (283 of 474 patients) had hepatitis B and 20.8% (99 of 474 patients) had a history of hepatitis C. Information on NASH was provided in two publications – in a cumulative cohort of 61 patients, NASH was present in 10 cases.<sup>31,37</sup> Alcohol overconsumption was described for

10 patients in 7 publications reporting on a total of 139 patients, amounting to 7.2%. AFP was provided in 15 publications. Serum AFP was increased above 20 IU/mL in 70.3% (26 patients) with maximum reaching 2372 IU/mL in a pooled cohort of 37 patients (Table 1).

## Radiological Features

In total, 14 articles reported radiological features of the analyzed s-HCCs.<sup>5,14–18,21,23,25,27,33,37,38,41</sup> Most frequently reported prominent radiological features on ultrasound, CT, and MRI were selected and quantified (Table 2). On ultrasound, most s-HCC lesions presented as lesions with hypochoic or mosaic patterns (each 47.6%), while an isochoic presentation was uncommon (4.8%). For lesions described as located near the capsule of the liver, retraction of the liver surface was present in 70% (14 of pooled 20 lesions). On CT scans, most lesions were well demarcated (66 of 122 cases [54.1%]). Sixty-seven percent presented with lobulated contours (41 of 61 cases). Persistent hyperdensity of the tumor in the delayed phase was reported in 87.0% of the lesions (80 of 92 cases). On MRI, target appearance was described in 65.0% of the patients (97 of 149 cases) in the hepatobiliary phase.

## Histopathological Features

Twenty-four articles<sup>4,5,14–16,18–27,29–33,36,38,41,42</sup> provided details on histopathological features of the tumors, summarized in Table 3.

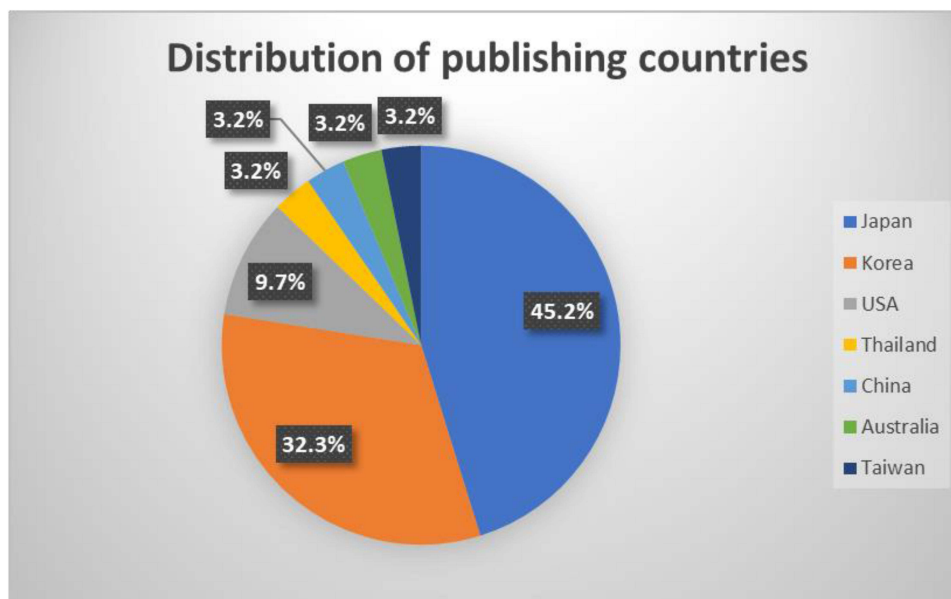
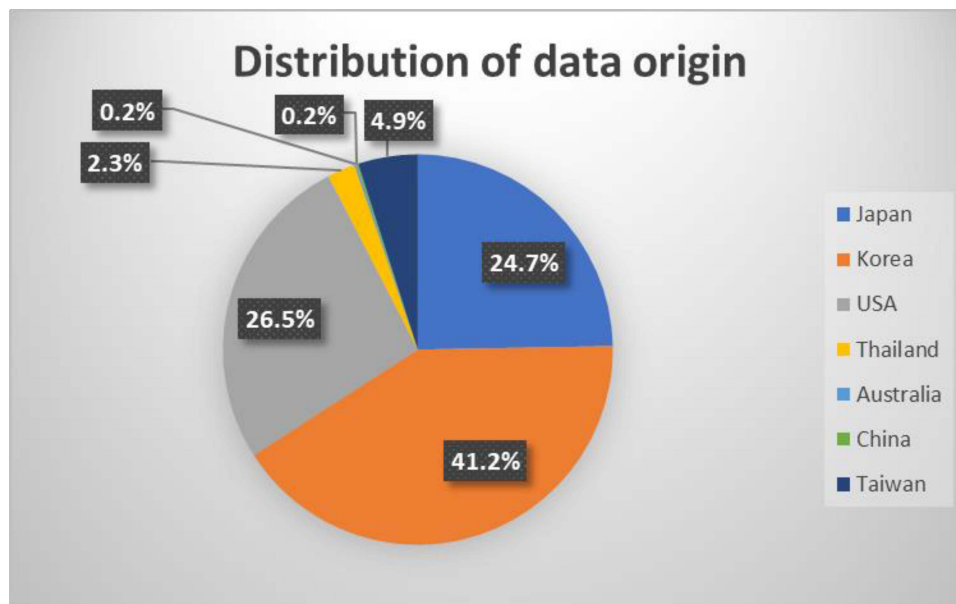


Figure 1 Distribution of countries to contribute publications on s-HCC.



**Figure 2** Distribution of origin of population data.

Of the patients, 82.9% had a solitary tumor (136 of 164 cases), while 17.1% presented with multiple lesions (28 of 164 cases). The tumor size ranged from 1.2 to 17.0 cm with a pooled mean and standard deviation estimated at  $4.2 \pm 2.33$  cm. Edmondson–Steiner (ES) grade was provided in 228 cases. 77.6% had ES grade I/II while 22.4% had grade III/IV. More specific ES grading was provided in 133 cases: while 30.1% had grade I, 50.4% had grade II, 18.0% had grade III and 1.5% were diagnosed with grade IV. WHO tumor grading was defined in 227 cases: 25.5% had Grade I, while 50.7% and 23.8% had Grade II and III, respectively. Due to multiple revisions of the AJCC staging system, no synthesis of TNM staging could be performed for included studies.

Background liver parenchyma was described in 19 publications. Of 314 patients, 69.0% (216 of 313 cases) had either cirrhosis or chronic hepatitis in the rest of the liver. Hepatitis was more prevalent amounting to 39.9% of all cases (117 of 293 patients), while cirrhosis was evident in 28.3% (83 of 293 cases). Normal liver parenchyma was described in 11.8% of the patients (37 of 313 cases). Steatosis was assessed by only two publications, with a total of 57 patients: 63.2% (36 of 57 cases) had steatosis on histopathology.

In 97.6% of the cases (41 of 42 patients), an irregular histopathological structure of the border was described. For cases close to the capsule, bulging appearance was described in 100% of pooled cases (51 of 51). A complete capsule was described in 26.3% of the patients (82 of 312

cases), although an incomplete capsule was present in 43 more. A septum was present in 85.6% of the patients (83 of 97 cases), while a central scar was reported for 63.5% (40 of 63 cases). At the time of diagnosis/resection, venous invasion was present in 32.6% (46 of 141 cases). Most cases (174 of 232 cases) presented without necrosis, amounting to 75%.

Only 14 studies<sup>4,5,14,15,18,20–22,24,27,30,31,35,36</sup> described immunohistochemical properties of the tumors summarized in Table 4. For targets analyzed in more than one publication, data is provided in Table 4. HepPar1 was evident in 64.6% of the patients (64 of 99 cases), while

**Table 1** Demographics of Patients

Feature	Metrics
<b>Age</b>	
Mean $\pm$ SD	55.8 $\pm$ 9.2 years
Range	24–78 years
<b>Gender</b>	
Male [n]	480
Female [n]	154
<b>Aetiology of liver disease</b>	
Hepatitis B	59.7% (283/474)
Hepatitis C	20.8% (99/474)
NASH	16.4% (10/61)
Alcohol	7.2% (10/139)
<b>AFP</b>	
>20 IU/mL	70.3% (26/37)



**Table 2** Radiological Features of s-HCCs

Imaging Modality	Criterion	Prevalence
Ultrasound	<b>Pattern:</b>	
	Hypoechoic	47.6% (10/21)
	Mosaic	47.6% (10/21)
	Isoechoic	4.8% (1/21)
CT	Liver retraction	70% (14/20)
	Well demarcated	54.1% (66/122)
	Lobulated contour	67% (41/61)
	Persistent hyperdensity in the delayed phase	87% (80/92)
MRI	Target appearance on HBP	65.0% (97/149)

CK 7 was positive in 40.7% of the patients (35 of 86 cases). CK 19 was evident in 16% (20 of 125 cases). EMA was positive in 41.9% of the patients (18 of 43 cases) (Table 4).

## Clinical Outcomes

Fifteen publications<sup>14,24,27–31,36,37,39,42</sup> provided information on follow-up. The follow-up ranged from 5 to 138 months. In 45.5% of the patients, a recurrence of s-HCC occurred during the follow-up period (70 of 154 cases) and 35.7% of the patients died of disease during the follow-up period (20 of 56 cases). A 5-year overall survival was reported between 21.1% and 73.3% with a pooled mean of 45.2%. All patients with reported outcomes have undergone hepatectomy.

## Discussion

Analysis of HCC morphological subtypes is essential to evolve treatment algorithms. With first-line treatment achieving a subpar median overall survival of merely 11–13 months<sup>45,46</sup> and recurrence rates after hepatectomy remaining high, the development of new therapies based on specific pathophysiological characteristics and a personalized approach is the only strategy to ensure no HCC patient is left behind. As numerous scores have been unable to sufficiently identify patients at risk for HCC development, recurrence, or progression,<sup>47</sup> translational research remains the only approach with potential. After identifying fundamental clinicopathological features of specific subtypes of HCC, associated genetic and molecular signal transduction elements can then be studied to improve prognostic accuracy and adjust

**Table 3** Histopathologic Features of s-HCCs

Histologic Finding	Metrics
Single tumor [%]	82.9% (136/164)
Multiple tumors [%]	17.1% (28/164)
Tumor size	
Range	1.2–17 cm
Mean ± SD	4.2 ± 2.33 cm
Edmondson grade	
I	30.1%
II	50.4%
III	18.0%
IV	1.5%
Grading	
G1	25.5% (58/227)
G2	50.7% (115/227)
G3	23.8% (54/227)
Background liver parenchyma	
Hepatitis	39.9% (117/293)
Cirrhosis	28.3% (83/293)
Normal liver parenchyma	11.8% (37/313)
Steatosis	63.2% (36/57)
Irregular histopathological structure	97.6% (41/42)
Bulging of the tumor*	100% (51/51)
Capsule	26.3% (82/312)
Septum	85.6% (83/97)
Central scar	63.5% (40/63)
Venous invasion	32.6% (46/141)
Necrosis:	
Present	25.0% (58/232)
Absent	75.0% (174/232)

**Note:** \*For tumors located near the capsule.

treatment and surveillance. The aim of this review was to summarize available knowledge on s-HCC and thus provide a solid basis for further research on s-HCCs.

The distribution of publications on s-HCC is striking. Although it is not surprising that countries with higher incidence of viral hepatitis, such as South Korea and Japan, would report on s-HCC more, the absence of publications from South America, Canada, China, Russia,

**Table 4** Immunohistochemical Features of s-HCCs

Immunohistochemical Feature	Prevalence
HepPar I positivity	64.6% (64/99)
CK7 positivity	40.7% (35/86)
CK19 positivity	16.0% (20/125)
EMA positivity	41.9% (18/43)

Africa, and Europe is surprising and may be indicative of publication bias. Additionally, s-HCC has only recently been recognized as a separate subtype of HCC, thus, some retrospective studies may perhaps still be pending.

Hepatitis B predominates in patients with s-HCC, afflicting approximately 60% of all reported cases, while hepatitis C causes liver disease in roughly an additional 20%. These proportions roughly mirror previously reported viral causes of all HCC subtypes,<sup>48</sup> although NAFLD and NASH have been rising as a cause of conventional HCC recently.<sup>49</sup> NAFLD was not prevalent in s-HCC, neither was alcohol overindulgence. With different causes predominating in conventional versus scirrhous HCCs, the one-for-all therapy approach appears counter-intuitive.

Individualized targeted therapy based on tumor receptor profiles has long been introduced in routine breast cancer treatment.<sup>50</sup> System-based approach and genomic profiling have identified numerous potential targets in HCC treatment, such as Wnt/ $\beta$ -catenin pathway, telomerases, and p53.<sup>51</sup> However, an individualized therapy strategy is far from implementation for HCC patients. Additionally, no profiling has thus far been made of atypical HCCs, an approach with the potential to identify targets with adequate subtype and patient specificity.

S-HCC incorporates typical and atypical radiological and histopathological HCC features. While most HCC lesions appear round and hypoechoic on ultrasound, a mosaic pattern was just as frequent in s-HCC. Commonly, this feature appears in HCC with degenerative changes.<sup>52</sup> Many imaging features of s-HCC seem to be related to its abundant desmoplastic tumor stroma. The fibrous stroma of s-HCC is presumably responsible for the frequent finding of retraction of the liver capsule, which is rare in untreated classical HCC. However, hepatic capsular retraction can occur in various other malignant and benign etiologies, including classical HCC with volume loss after locoregional therapy, other malignant liver tumors with fibrous stroma (cholangiocarcinoma, epithelioid hemangioendothelioma, FL-HCC), and confluent hepatic fibrosis, which often occurs in liver cirrhosis.<sup>53</sup> Moreover, hepatic hemangiomas are more likely to become fibrotic and decrease in size in progressively cirrhotic livers and can retract the liver capsule.<sup>54</sup> The dynamic contrast enhancement of s-HCC resembles that of cholangiocarcinomas, as both are rich in fibrous stroma.<sup>55</sup> In contrast to the fast wash-in and wash-out of untreated conventional HCC, s-HCC and

cholangiocarcinomas frequently present with a peripheral rim APHE and progressive enhancement in the venous and delayed phases, whereby the areas of delayed enhancement often closely correspond to the presence of fibrotic fibers on histopathology.<sup>56</sup> The presence of delayed enhancement in parts other than septae and the capsule in untreated HCC-lesions can be seen as indicative for a variant type of HCC, particularly s-HCC.<sup>56</sup> However, delayed enhancement is also typical for metastases of colorectal adenocarcinoma.<sup>57</sup>

Although rare, the most frequent complications of biopsy are seeding and bleeding; thus, more emphasis has been placed on radiological imaging in the diagnostic algorithm.<sup>58</sup> Yet, histopathological assessment is necessary for a definitive diagnosis. Most studies report experience with s-HCC after hepatectomy; however, since hepatitis-associated liver disease tends to precede s-HCC and many patients with HCC and liver cirrhosis can neither undergo surgery nor biopsy, a certain reporting bias must be assumed.<sup>58,59</sup>

Edmondson–Steiner classification is one of the most widely used histopathological classifications of HCC. Most s-HCC patients had grade I/II with less than 25% presenting with grade III/IV, which contrasts general reports on HCC.<sup>60,61</sup> It has been shown that ES grade may correlate with the recurrence of HCC.<sup>61</sup> With 44.9% of the patients with available data presenting with a recurrence during follow-up and generally reported lower ES grades for s-HCC, this hypothesis is, although indirectly, supported.

Specific markers for distinct lesions are crucial. Since the identification of DNAJB1-PRKACA oncogenic fusion in FL-HCC that has high sensitivity, the accuracy of diagnosis improved, and novel therapeutic targets could be proposed.<sup>62</sup> EMA is a marker of combined hepatocellular-cholangiocarcinoma<sup>63</sup> and, interestingly, was positive in nearly half of all s-HCC cases (41.9%) seeding doubt as to where s-HCC stands on the spectrum of HCC lesions. However, a sensitive marker for s-HCC is yet to be identified. Some authors suggest arginase-1 may be a more specific marker for HCC.<sup>64</sup> This may also hold true for s-HCC; Krings et al described a cohort of 20 patients with s-HCC and 11 out of 13 cases were positive for arginase, while only 5 of 19 were positive for HepPar1.<sup>4</sup> Unfortunately, this marker is widely unexplored in context of s-HCC. Additionally, polyclonal CEA<sup>65</sup> may present an interesting stain for s-HCC, but thus far unexplored. Surprisingly, most s-HCCs were

described to have a bulging appearance on histopathological examination when located near the capsule, yet, on radiological examination, surface retraction was most commonly reported – a clear contradiction, which warrants a comparative investigation.

The analyzed data showed a 5-year overall survival rate of 45.2%, which is on the lower end of survival for HCC, as typical post-resection rates range from 41% to 74%.<sup>66</sup> At the same time, the weight of the US registry significantly impacted the pooled mean, as it had the most patients and showed the lowest 5-year overall survival rates. Due to such discrepancies in reported survival rates, socioeconomic reasons and the structure of health care systems must be considered during the comparison. The recurrence rate reported was lower compared to the reported recurrence of up to 70% for all HCCs (68). With most patients presenting with hepatitis and cirrhosis and follow-up information lacking for most patients, it is vital to consider that the true recurrence and survival rates remain concealed and may lay far below currently presented.

## Certainty of Evidence

The certainty of evidence within a systematic review or a meta-analysis is contingent on the design and methods of the included studies. It can be greatly affected by each publication and its risk of bias, as well as other factors such as inconsistency, imprecision, and indirectness. Additionally, a bias considerably affecting systematic reviews of rare conditions or interventions is publication bias. Inevitably, certainty in evidence derived from case series and reports will be very low. Nevertheless, inferences from these on rare and underreported conditions can be used for decision-making and is the best evidence on such afflictions to date.

The methodological quality assessment of included studies was performed using a tool proposed by Murad et al and assessed the domains of selection, ascertainment, causality, and reporting of the included case reports and case series.<sup>43</sup> For observational studies, MINORS was additionally used as a risk of bias assessment. Overall, due to the study design and conceptual framework, the quality of evidence is very low.

## Conclusion

S-HCC is a rare subtype of HCC that primarily arises in diseased livers with hepatitis or cirrhosis but lacking

symptoms. S-HCC can often be misdiagnosed as fibrolamellar carcinoma, cholangiocarcinoma, or hepatic metastasis. Most cases of s-HCC have, perhaps, not been reported yet. Comparatively, low recurrence rate (45.5%), yet poor 5-year survival rate (45.2%) indicate that confounding factors may compromise patient life expectancy. A more comprehensive analysis of complex subtypes of HCC is possible with more reports from various parts of the world, and special emphasis must be placed on long-term outcomes after treatment, and clinical and histological features. Currently, patients with atypical lesions in the liver should be presented at tertiary referral hospitals for interdisciplinary assessment and treatment.

## Abbreviations

AFP, alpha fetoprotein; AJCC, American Joint Committee on Cancer; APHE, arterial phase hyperenhancement; CK, cytokeratin; CT, computer tomography; EMA, epithelial membrane antigen; ES, Edmondson–Steiner grading; GRADE, Grading of Recommendations, Assessment, Development and Evaluations; HBP, hepatobiliary phase; HCC, hepatocellular carcinoma; HepPar1, hepatocyte paraffin 1 antigen; IPD, independent patient data; FL-HCC, fibrolamellar hepatocellular carcinoma; s-HCC, scirrhous hepatocellular carcinoma; MRI, magnetic; NASH, non-alcoholic steatohepatitis; RELIVE, initiative for generating evidence in diagnosis and therapy of Rare LIVER disease; PRISMA, Preferred Reporting Items for Systematic Reviews and Meta-Analyses; WHO, World Health Organization.

## Data Sharing Statement

All data generated or analyzed during this study are included in this published article and its Supplementary Materials.

## Author Contributions

All authors contributed to data analysis, drafting or revising the article, gave final approval of the version to be published, agreed to the submitted journal, and agree to be accountable for all aspects of the work.

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## Disclosure

The authors declare that they have no conflicts of interest for this work.



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