Prognostic Value of the Albumin-Bilirubin Grade for the Prediction of Post-Hepatectomy Liver Failure: A Systematic Review and Meta-Analysis

Giovanni Marasco 1,2,*†, Luigina Vanessa Alemanni 1,2,†, Antonio Colecchia 3, Davide Festi 2, Franco Bazzoli 1,2, Giuseppe Mazzella 2, Marco Montagnani 1,2 and Francesco Azzaroli 1,2

1 IRCCS Azienda Ospedaliero-Universitaria di Bologna, 40138 Bologna, Italy; vanessaalemanni1@gmail.com (L.V.A.); franco.bazzoli@unibo.it (F.B.); marco.montagnani@unibo.it (M.M.); francesco.azzaroli@unibo.it (F.A.)
2 Department of Medical and Surgical Science, University of Bologna, 40126 Bologna, Italy; davide.festi@unibo.it (D.F.); giuseppe.mazzella@unibo.it (G.M.)
3 Gastroenterology Unit, University Hospital Borgo Trento, 37100 Verona, Italy; antonio.colecchia@aovr.veneto.it
* Correspondence: giovanni.marasco4@unibo.it; Tel.: +39-051-214-5265
† Equal contribution as first authorship.

Abstract: (1) Introduction: Liver resection (LR) for hepatocellular carcinoma (HCC) is often burdened by life-threatening complications, such as post-hepatectomy liver failure (PHLF). The albumin-bilirubin (ALBI) score can accurately evaluate liver function and the long-term prognosis of HCC patients, including PHLF. We aimed to evaluate the diagnostic value of the ALBI grade in predicting PHLF in HCC patients undergoing LR. (2) Methods: MEDLINE, Embase, and Scopus were searched through January 17th, 2021. Studies reporting the ALBI grade and PHLF occurrence in HCC patients undergoing LR were included. The Odds Ratio (OR) prevalence with 95% confidence intervals (CI) was pooled, and the heterogeneity was expressed as $I^2$. The quality of the studies was assessed using QUADAS-2 (Quality Assessment of Diagnostic Accuracy Studies). (3) Results: Seven studies met the inclusion criteria and were included in the analysis. A total of 5377 patients who underwent LR for HCC were considered, of whom 718 (13.4%) developed PHLF. Patients with ALBI grades 2 and 3 before LR showed increased rates of PHLF compared to ALBI grade 1 patients. The pooled OR was 2.572 (95% CI, 1.825 to 3.626, $p < 0.001$), with substantial heterogeneity between the studies ($I^2 = 69.6%$) and no publication bias (Begg’s $p = 0.764$ and Egger’s $p = 0.851$ tests). All studies were at a ‘low risk’ or ‘unclear risk’ of bias. Univariate meta-regression analysis showed that heterogeneity was not dependent on the country of study, the age and sex of the participants, the definition of PHLF used, the rate of patients in Child–Pugh class A or undergoing major hepatectomy. (4) Conclusions: In this meta-analysis of published studies, individuals with ALBI grades of 2 and 3 showed increased rates of PHLF compared to ALBI grade 1 patients.

Keywords: hepatocellular carcinoma; liver surgery; hepatectomy; liver failure; albumin-bilirubin; ALBI; PHLF

1. Introduction

Hepatocellular carcinoma (HCC) represents the second cause of cancer-related death worldwide [1]; in 90% of cases, it develops with underlying liver disease, leading to a relevant burden in morbidity and mortality in patients affected by chronic liver disease [2]. Despite several techniques for HCC management that have been developed in the last decades, liver resection (LR) still represents the main curative treatment offering the best outcome [3]. LR is often burdened by life-threatening complications, such as post-hepatectomy liver failure (PHLF) [4].
PHLF has been described with an incidence ranging from 8% to 12% [5]; this variability is related to different PHLF definitions and the severity of the underlying liver disease, the extent of surgery, and the intraoperative course [4]. The current International Study Group of Liver Surgery (ISGLS) definition for PHLF represents the current standard method for its diagnosis [5,6], defined as an acquired deterioration of the liver functions characterized by an increased International Normalized Ratio (INR) and hyperbilirubinemia after postoperative day 5 [7].

Thus, an accurate pre-operative assessment of patients undergoing LR is required to overcome the PHLF risk, through the evaluation of liver function and the assessment of portal hypertension [8–10]. Several markers have been previously proposed for this purpose, such as the Child–Pugh, liver stiffness measurement, and volumetric imaging for future liver remnant assessment, which are typically performed in the pre-operative work [9,11]. In Eastern countries a large use of the indocyanine green clearance (ICG) has been reported [12].

However, these evaluations are often expensive, time consuming, or inaccurate in providing a precise estimation of post-operative hepatic functional recovery [13]. Recently, the albumin-bilirubin (ALBI) score, a new non-invasive tool to evaluate liver function and predict survival in HCC patients [14], showed promising results in predicting the long-term prognosis of chronic liver disease [15] and HCC patients [16], including PHLF prediction. However, no definitive results are available for this latter outcome and, consequently, the ALBI grade has not been completely endorsed in the pre-operative assessment of HCC patients undergoing LR. Thus, we performed a systematic review with meta-analysis to evaluate the diagnostic value of the ALBI grade in predicting PHLF in HCC patients undergoing LR.

2. Materials and Methods

We performed a systematic review and meta-analysis following the recommendations of the Cochrane Collaboration Diagnostic Test Group [17] and according to the PRISMA (Preferred Reporting Items for Systematic Reviews and Meta-Analyses) guidelines [18].

2.1. Search Strategy and Study Selection

We searched on MEDLINE via PubMed, Ovid Embase, and Scopus, to identify relevant articles published up to 17 January 2021. The electronic search of the literature was conducted using the following keywords: ‘ALBI’ or ‘albumin bilirubin’, ‘PHLF’ or ‘post-operative liver failure’ or ‘post hepatectomy liver failure’, and ‘liver resection’ or ‘hepatectomy’ or ‘hepatic resection’.

The search was extended until 2015, when the first article published on ALBI scores was published [14]. In addition, the abstracts of the conference proceedings of Digestive Diseases Week, United European Gastroenterology Week, International Liver Congress, American Association for the Study of Liver Diseases Meeting, and Asian Pacific Association for the Study of the Liver Congress for the same period were searched electronically and by hand.

The complete search strategies are reported in Supplementary Material 2. There were no limitations on the type of study, publication date, or manuscript language. Two reviewers (LVA and GM) independently performed the initial screening and selection, based on titles and abstracts. Eligible full-text articles were separately evaluated by the two authors; in the case of discrepancies, they were resolved through discussion with a third reviewer (FA).

Studies were selected and included in final analysis when they met the following criteria: studies conducted on patients affected by chronic liver diseases undergoing LR for HCC, reporting data on the ALBI grade in patients developing PHLF or not. All etiologies for liver disease were included. In the presence of studies reporting cohorts who underwent LR for HCC and other malignancies, further data on only HCC patients were requested.
from the authors; in the case of no response, we established a minimum of 90% of HCC within a study population to include the paper in our analysis.

Only studies reporting PHLF diagnosed according to ISGLS criteria [7] were included, as this classification has been widely endorsed [5]. As recommended by ISGLS [7], PHLF had to be diagnosed in the case of increased serum International Normalized Ratio (INR) and concomitant hyperbilirubinemia, after 5 postoperative days. The severity of PHLF was therefore graded as: grade A PHLF, requiring no specific treatment; grade B PHLF requiring essential non-invasive treatment (transfusion support, albumin supplementation, and diuretic therapy); grade C PHLF requiring invasive procedures, including mechanical ventilation, hemodialysis, or extracorporeal liver support [7].

PHLF grades B and C are, thus, considered clinically significant [7]. The ALBI score is based on the serum albumin and total bilirubin levels, calculated with the formula: (log10 bilirubin [µmol/L] \times 0.66) + (albumin [g/L] \times −0.0852). This score is further categorized into three different grades for rapid clinical use: ALBI 1 (≤ −2.60), ALBI 2 (−2.60 to ≤ −1.39), and ALBI 3 (> −1.39) [14].

We included only studies reporting the number of PHLF cases for each ALBI grade. For studies reporting the ALBI score instead of the grade, we contacted the authors in order to collect the missing data. Studies were excluded if they did not meet the inclusion criteria or when essential information was missing in the available manuscript or could not be obtained from the authors.

2.2. Data Extraction and Quality Assessment

Two authors (LVA and GM) independently extracted relevant data on the publication, study methods, and results using a standardized data extraction form. The following items were extracted from each study: type of study, year of publication, country, study design, total number of patients enrolled, age and sex of the participants, Child–Pugh classification, main etiology of liver disease, the extent of LR, the number of PHLF cases classified by the severity degrees (PHLF A, B, and C), and ALBI grade groups.

If multiple publications on a same cohort were found, the latest and most complete publication was considered. Subsequently, the methodological quality of the included studies was separately assessed by two reviewers (LVA and GM), according to the Quality Assessment of Diagnostic Accuracy Studies (QUADAS-2) tool [19] (Supplementary Material 3). QUADAS-2 is an evidence-based tool consisting of 14 items phrased as questions, each of which are scored a “yes”, “no”, or “unclear”, examining the presence of bias in the study. Disagreements were resolved through discussion or arbitration by a third reviewer (FA), when necessary.

2.3. Statistical Analysis

The rates of PHLF in patients with ALBI grade 1 and ALBI grades 2 and 3 were extracted from all studies. As we expected that only a small amount or no patients undergoing LR would have severely impaired liver function according to ALBI grade 3, we decided to consider for statistical analyses ALBI grades 2 and 3 compared to ALBI grade 1. The pooled Odds Ratios (ORs) with corresponding 95% Confidence Intervals (CI) and \( p \) were calculated to assess the association between the ALBI grade and PHLF occurrence in patients with HCC.

Heterogeneity across the studies was assessed using the Higgins I\(^2\) statistics. The value of I\(^2\) describes the percentage of variability in point estimates due to heterogeneity rather than to sampling error: low-moderate for I\(^2\) < 50% and high for I\(^2\) ≥ 50% [20]. If there was no heterogeneity (<50%, \( p > 0.1 \)), the fixed-effect model was used; otherwise, the random-effect model was applied. The ALBI grades 2 and 3 were closely associated with PHLF when the OR > 1. The publication bias was measured by Begg’s test and Egger’s test with a graph; a \( p \) value < 0.05 indicated a significant small size study effect.

Briefly, Begg’s test and Egger’s test are based on the statistical evaluation of a funnel plot, which shows the effect sizes plotted against their standard errors, instead of the
visual evaluation of asymmetry. While Begg’s test examines the correlation between the effect sizes and their variances, Egger’s test regresses the standardized effect sizes on their precisions. The Duval and Tweedie [21] non-parametric ‘trim and fill’ method was also used, accounting for publication bias in the meta-analysis [22]. Pooled ORs following adjustment for publication bias using the ‘trim and fill’ method are reported. Subgroup analyses were conducted after excluding studies with possible sources of heterogeneity (studies including subgroups of patients other than HCC or including only PHLF B and C as the case groups).

As part of the sensitivity analysis, the impact of confounding covariates (country, age of participants, sex, rate of patients in Child–Pugh class A or undergoing major hepatectomy, definition of PHLF (PHLF vs. clinically significant PHLF)) on the meta-analytic results was evaluated using meta-regression analysis [23], reporting β coefficient ± standard error (SE). Since a low number of studies was found, the p values were also re-calculated using Monte Carlo permutation [24] with 5000 permutations to obtain sufficient precision [25]. All analyses were carried out using STATA statistical software (Stata Corp., College Station, TX, USA).

3. Results
3.1. Study Selection

The electronic and manual searches provided 215 records; after duplicate elimination, 180 studies went through screening based on the title and abstracts. After the first screening, 25 records (19 [26–44] full-text papers and six [45–50] abstracts) remained for fully eligibility evaluation, of which one [29] was found during the manual search.

Among them, six [26,27,30,34,47,48] were excluded from the meta-analysis due to insufficient data or no response by the study authors; five [28,35,46,49,50] studies were excluded since the reporting cohorts were already included or best characterized in more recent studies; one study [42] was not pertinent since it was performed on patients undergoing extrahepatic surgery; a further six studies [31–33,37,39,41] included large groups of patients undergoing LR for malignancies other than HCC and/or did not use the ISGLS criteria [7] for PHLF diagnosis. Finally, a total of seven studies [29,36,38,40,43–45], all full text except one [45], met the eligibility criteria and were included in the meta-analysis, as shown in Figure 1.
3.2. Study Characteristics

The seven studies included [29,36,38,40,43–45] reported a total number of 5377 patients undergoing LR for HCC, of whom 718 (13.4%) developed PHLF, among which 502 (69.9%) were clinically significant. Only one study [44] included a negligible sub-group (9.2%) of patients who underwent LR for reasons other than HCC. Notably, two studies [38,44] reported only on PHLF grades B and C, while others [29,36,40,43,45] considered all grades of severity. The characteristics of the studies included are shown in Table 1.
### Table 1. The characteristics of the included studies in the systematic review and meta-analysis.

| Author, Year | Country | Design of the Study | Total Pts | Age | Sex (Male), n. (%) | Etiology, % | Child Pugh, n. (%) | Extent of Hepatectomy n. (%) | Outcome | ALBI Grade n., (%) | Total PHLF |
|--------------|---------|---------------------|-----------|-----|-------------------|-------------|-------------------|-------------------------------|---------|-------------------|------------|
| Wang, 2016 [43] | China | Retrospective | 1242 | >60: 223 (18%) | 1072 (86.3) | HBV 85.3% | A 1189 (95.7), B 53 (4.3), C 0 (0) | Minor 975 (78.5), Major 267 (21.5) | PHLF A + B + C | ALBI 1, 850 (68.4), ALBI 2, 390 (31.4), ALBI 3, 20 (2.2) | 166, Grade A 58, Grade B 91, Grade C 17 |
| Chong, 2018 [45] | China | Retrospective | 396 | 59.7 c | 334 (84.3) | HBV 80.3% | A 397 (97.7), B 9 (2.3), C 0 (0) | Minor 243 (61.4), Major 153 (38.6) | PHLF A + B + C | ALBI 1, 302 (76.25), ALBI 2, 93 (23.5), ALBI 3, 1 (0.25) | 109, Grade A 52, Grade B/C 57 |
| Zhang, 2018 [40] | China | Retrospective | 338 | 52 [44–66] b | 299 (88.5) | HBV 82.2% | A 308 (91.1), B 30 (8.9), C 0 (0) | ALBI 1, 302 (76.25), ALBI 2, 93 (23.5), ALBI 3, 1 (0.25) | PHLF A + B + C | ALBI 1, 134 (39.6), ALBI 2, 198 (58.6), ALBI 3, 6 (1.8) | 26, Grade A 8, Grade B/C 57 |
| Zou, 2018 [36] | China | Retrospective | 473 | 52 (18–77) a | 411 (86.9) | HBV 85.4% | A 427 (90.3), B 46 (9.7), C 0 (0) | Minor 356 (75.3), Major 117 (24.7) | PHLF A + B + C | ALBI 1, 189 (40), ALBI 2, 282 (59.6), ALBI 3, 2 (0.4) | 50 |
| Lu, 2019 [38] | China | Retrospective | 2038 | >50: 948 (46.5%) | 1810 (88.8) | HBV 88.9% | A 2038 (100), B 0 (0), C 0 (0) | Minor 1501 (73.7), Major 537 (26.3) | PHLF B + C | ALBI 1, 1570 (77), ALBI 2, 468 (23), ALBI 3, 0 (0) | 196 |
| Russolillo, 2019 [29] | Italy | Retrospective | 400 | 70 (24–86) a | 339 (84.8) | Mixed (HCV 40%) | A 385 (96.25), B 15 (3.75), C 0 (0) | Minor 299 (74.7), Major 101 (25.3) | PHLF A + B + C | ALBI 1, 208 (52), ALBI 2, 188 (47), ALBI 3, 4 (1) | 82, Grade A 48, Grade B/C 34 |
| Sposito, 2020 [44] | Italy | Prospective | 490 | 68.6 [61.4–74.7] b | 360 (73.5) | Mixed (HCV 59%) | A 463 (94.5), B 27 (5.5), C 0 (0) | Minor 457 (93.3), Major 33 (6.7) | PHLF B + C | ALBI 1, 217 (44.3), ALBI 2, 268 (54.7), ALBI 3, 5 (1.0) | 89 |

**Abbreviations:** pts: patients; n.: number; ALBI: Albumin-Bilirubin score; HBV: Hepatitis B Virus; HCV: Hepatitis C Virus; N/A: Not Available; and PHLF: Post-Hepatectomy Liver Failure. a = median, (range); b = median, [interquartile range]; and c = median.
In particular, five studies [36,38,40,43,45] were conducted in China, whereas two [29,44] were conducted in Italy. Five studies [29,36,40,44,45] reported the median age of the participants, which ranged from 52 [36,40] to 70 [29] years. In all studies, a greater rate of male patients were included, with the proportion ranging from 73.5% [44] to 88.8% [38]. All the studies had a retrospective design, with the exception of one prospective study [44]. Six studies [29,36,38,43–45] reported the extent of the hepatectomy. Notably, the rate of major hepatectomy ranged from 6.7% [44] to 38.6% [45].

Regarding liver function, the Child–Pugh A was reported in variables rates, ranging from 90.3% [36] to 100% [38], whereas the Child B was reported in rates ranging from 0% [38] to 9.7% [36]. Asian studies [36,38,40,43,45] presented mainly HBV-related liver disease (from 80.3% [45] to 88.9% [38]), while, in European studies, different etiologies were reported (from 40% [29] to 59% [44] HCV-related). Within the included studies, patients were categorized according to an ALBI grade as follows: grade 1 (3470 pts, 64.53%), grade 2 (1887 pts, 35.1%), and grade 3 (20 pts, 0.37%). The PHLF rates for each ALBI category were as follows: ALBI 1 (330 pts, 9.5%), ALBI 2 (376 pts, 19.9%), and ALBI 3 (12 pts, 60%).

### 3.3. Quality Assessment

The evaluation of the methodological quality of the included studies is reported in Supplementary Material 4 and in Table 2. The studies considered in the meta-analysis had an overall low risk of bias according to QUADAS-2. However, all studies presented ‘unclear risk’ concerning the risk of bias regarding the ‘reference test’ and the ‘flow and timing’. Indeed, in all studies, it was not reported whether the diagnosis of PHLF was blinded to the ALBI grade. The exact timing of the pre-operative functional tests was also not specified in the studies included. An ‘unclear risk’ of bias in the ‘patient selection’ was present in three studies [36,38,40]. Concerns regarding the applicability of the reference standard were raised for two studies [38,44], as they reported only clinically significant PHLF.

**Table 2.** The risk of bias and applicability concerns of the included studies.

| Study                  | Patient Selection | Index Test | Reference Standard | Flow and Timing |
|------------------------|-------------------|------------|-------------------|-----------------|
|                        | Risk of Bias      | Concerns about Applicability | Risk of Bias | Concerns about Applicability | Risk of Bias | Concerns about Applicability | Risk of Bias |
| Wang, 2016 [43]        | L                 | L          | L                 | U               | L               | L                            | U              |
| Chong, 2018 [45]       | L                 | L          | L                 | U               | L               | L                            | U              |
| Zhang, 2018 [40]       | U                 | L          | L                 | U               | L               | H                            | U              |
| Zou, 2018 [36]         | U                 | L          | L                 | U               | L               | L                            | U              |
| Lu, 2019 [38]          | U                 | L          | L                 | U               | L               | H                            | U              |
| Russolillo, 2019 [29]  | L                 | L          | L                 | U               | L               | L                            | U              |
| Sposito, 2020 [44]     | L                 | L          | L                 | U               | H               | L                            | U              |

L, low; H, high; and U, unclear.

### 3.4. ALBI and PHLF Occurrence

Patients with ALBI grades 2 and 3 before LR showed increased rates of PHLF compared to ALBI grade 1 patients. The pooled OR using a was 2.572 (95% CI, 1.825 to 3.626) (Figure 2). This difference was statistically significant ($p < 0.001$). There was substantial heterogeneity between the studies ($I^2 = 69.6$%). No publication bias was found using Begg’s ($p = 0.764$) and Egger’s ($p = 0.851$) tests (Figure 3). Due to an asymmetrical appearance of the funnel plot, the ‘trim and fill’ method was applied, indicating no missing studies.

The re-estimated OR slightly increased but remained significantly different among the two groups (OR 2.997, 95% CI 2.193 to 3.801, $p < 0.001$). The subgroup analyses after removing one study [44], including a small subgroup of patients other than HCC (<10%) (OR 2.564, 95% CI 1.708 to 3.849, $I^2 = 74.6$, $p < 0.001$) (Figure 4), and after removing two
studies [38,44], including only PHLF B and C as the case groups (OR 2.543, 95% CI 1.446 to 4.471, $I^2$ 79.3%, $p < 0.001$) (Figure 5), showed a slight variation in the ORs.

| Study      | Events, ALBI grade 2-3 | OR (95% CI) | Events, ALBI grade 1 |
|------------|------------------------|-------------|----------------------|
| Wang (2018)| 97/392                 | 3.72 (2.66, 5.21) | 69/850               |
| Chong (2018)| 28/94                  | 1.16 (0.70, 1.93) | 81/302               |
| Zou (2018) | 42/284                 | 3.93 (1.80, 8.57) | 8/189                |
| Zhang (2018)| 23/204                 | 5.55 (1.63, 18.87) | 3/124                |
| Lu (2019)  | 84/468                 | 2.85 (2.10, 3.86) | 112/1570             |
| Russelillo (2015) | 48/192       | 1.71 (1.04, 2.79) | 34/298               |
| Spisolo (2020) | 60/273          | 2.69 (1.61, 4.49) | 23/217               |
| Overall    | 388/1907              | 2.57 (1.82, 3.63) | 330/3470             |

**Figure 2.** Forest plot of the pooled Odds Ratio (OR) for post-hepatectomy liver failure (PHLF) in ALBI grades 2 and 3 compared to ALBI grade 1. Events: PHLF/No PHLF; and CI: Confidence Interval.

**Figure 3.** Funnel plot visual to asymmetry. Legend: OR: Odd Ratio. se(logOR): Standard Error of log OR. Dotted black line: the line of pseudo 95% confidence limits. Solid black line: the line of overall effect. Blue point: each study included. Orange line: Egger’s test.
Univariate meta-regression analysis was used to explore and explain potential sources of heterogeneity among the studies. None of the variables assessed was able to explain the high heterogeneity found (Table 3) even after 5000 permutations.

Table 3. Results of the univariable meta-regression analysis.

| Covariates           | Number of Studies | Beta Coefficient ± SE | Adjusted R² (%) | p Value | p Value ± SE after Montecarlo Permutation |
|----------------------|-------------------|-----------------------|-----------------|---------|-----------------------------------------|
| Country              | 7                 | 0.308 ± 0.369         | -2.03           | 0.371   | 0.432 ± 0.007                           |
| Age                  | 5                 | 0.871 ± 0.074         | 29.67           | 0.202   | 0.283 ± 0.006                           |
| Sex (Male)           | 7                 | 1.097 ± 0.126         | -9.15           | 0.460   | 0.485 ± 0.007                           |
| Child–Pugh A         | 7                 | 1.014 ± 0.017         | -6.57           | 0.442   | 0.437 ± 0.007                           |
| Major hepatectomy    | 5                 | 0.956 ± 0.046         | -3.46           | 0.424   | 0.316 ± 0.007                           |
| PHLF definition      | 7                 | 1.462 ± 1.900         | -21.97          | 0.782   | 0.852 ± 0.005                           |

SE = Standard Error; R² = Relative reduction in between-study variance: the value indicates the proportion of between-study variance explained by covariate; and PHLF: Post-Hepatectomy Liver Failure.

4. Discussion

PHLF represents a major event in patients undergoing LR and mostly affects patients with chronic liver disease complicated by HCC development [4]. To date, the selection of patients undergoing LR according to the risk of post-operative complications, such as PHLF, is unsatisfactory [13]. This systematic review and meta-analysis included six studies reporting data on the ALBI grade in patients developing PHLF. The pooled data available from these studies showed that patients with ALBI grades of 2 and 3 had an increased rate of developing PHLF compared to ALBI grade 1 patients (OR of 2.572). To our knowledge, this is the first meta-analysis aiming to assessing the association between the ALBI score and PHLF occurrence using the QUADAS-2 tool for a correct evaluation of the methodological quality of the studies included. The association between

**Figure 4.** Forest plot of the pooled Odds Ratio (OR) for post-hepatectomy liver failure (PHLF) in ALBI grades 2 and 3 compared to ALBI grade 1 after removing one study that included a small subgroup of patients other than HCC. Events: PHLF/No PHLF; and CI: Confidence Interval.

**Figure 5.** Forest plot of the pooled Odds Ratio (OR) for post-hepatectomy liver failure (PHLF) in ALBI grades 2 and 3 compared to ALBI grade 1 after removing two studies that included only PHLF B and C as case groups. Events: PHLF/No PHLF; and CI: Confidence Interval.
Univariate meta-regression analysis was used to explore and explain potential sources of heterogeneity among the studies. None of the variables assessed was able to explain the high heterogeneity found (Table 3) even after 5000 permutations.

Table 3. Results of the univariable meta-regression analysis.

| Covariates         | Number of Studies | Beta Coefficient ± SE | Adjusted R² (%) | p Value | p Value ± SE after Montecarlo Permutation |
|--------------------|-------------------|-----------------------|-----------------|---------|-------------------------------------------|
| Country            | 7                 | 0.308 ± 0.369         | −2.03           | 0.371   | 0.432 ± 0.007                             |
| Age                | 5                 | 0.871 ± 0.074         | 29.67           | 0.202   | 0.283 ± 0.006                             |
| Sex (Male)         | 7                 | 1.097 ± 0.126         | −9.15           | 0.460   | 0.485 ± 0.007                             |
| Child–Pugh A       | 7                 | 1.014 ± 0.017         | −6.57           | 0.442   | 0.437 ± 0.007                             |
| Major hepatectomy  | 5                 | 0.956 ± 0.046         | −3.46           | 0.424   | 0.316 ± 0.007                             |
| PHLF definition    | 7                 | 1.462 ± 1.900         | −21.97          | 0.782   | 0.852 ± 0.005                             |

SE = Standard Error; R² = Relative reduction in between-study variance: the value indicates the proportion of between study variance explained by covariate; and PHLF: Post-Hepatectomy Liver Failure.

4. Discussion

PHLF represents a major event in patients undergoing LR and mostly affects patients with chronic liver disease complicated by HCC development [4]. To date, the selection of patients undergoing LR according to the risk of post-operative complications, such as PHLF, is unsatisfactory [13]. This systematic review and meta-analysis included six studies reporting data on the ALBI grade in patients developing PHLF. The pooled data available from these studies showed that patients with ALBI grades of 2 and 3 had an increased rate of developing PHLF compared to ALBI grade 1 patients (OR of 2.572).

To our knowledge, this is the first meta-analysis aiming to assessing the association between the ALBI score and PHLF occurrence using the QUADAS-2 tool for a correct evaluation of the methodological quality of the studies included. The association between PHLF and ALBI is undoubtedly explained by the accuracy of the ALBI grade in non-invasively mirroring the liver function [14] even in patients with mild or early stage liver disease. The prevention of PHLF is achievable mostly by a careful liver function assessment in preoperative examinations [3].

The Child–Pugh classification remains the most applied method for the evaluation of the liver reserve in the preliminary evaluations for LR [3]. However, in recent years, concerns regarding the adequacy of the Child–Pugh classification have emerged due to the subjectivity and insufficient ability in stratifying the individual risks of patients with mild severity liver diseases [14,51,52]. Instead, the ALBI grade showed a greater accuracy in further stratifying the prognosis of HCC patients belonging to Child–Pugh class A [14,52,53]. Two recent meta-analyses including HCC patients [16,51] reported a higher predictive value of the ALBI grade compared to the Child–Pugh class for stratifying patient survival.

Indeed, higher ALBI grades were associated with poor overall survival (OS) (HR = 2.060, 95% CI: 1.909–2.211, p = 0.000) [16] even in HCC patients undergoing LR [51]. Another recent meta-analysis [54] confirmed that the ALBI grade was able to better stratify the prognosis of HCC patients undergoing treatments. Specifically, among Child–Pugh class A patients, those with ALBI grade 1 showed a higher OS rate compared to ALBI grade 2 [54], even after surgical resection. However, none of these previous pooled data analyses was focused on PHLF as the main outcome.

Since liver function impairment is the main determinant of PHLF development and the vast majority of candidates to LR belonged to Child–Pugh class A [54,55], we expected that the ALBI grade could be a valuable tool for PHLF risk stratification. Our meta-analysis, evaluating a population almost entirely stratified as Child–Pugh class A (96.5%), confirmed its good performance in this setting, suggesting that a further stratification, over the Child–Pugh classification, could be safely and non-invasively applied in clinical practice without other time-consuming examinations.
Further supporting the ALBI superiority in evaluating liver function and patient prognoses, one [36] of the studies included in the present meta-analysis showed that the ALBI score (AUC 0.745) was more accurate than the Child–Pugh classification (AUC 0.665), ICG R15 (AUC 0.668), and MELD score (AUC 0.649) in predicting PHLF. However, the MELD score was specifically designed for end-stage cirrhotic patients [56], and thus a low accuracy in predicting PHLF in compensated Child–Pugh A patients undergoing LR was expected.

This meta-analysis has some weaknesses. The small number of studies included could have led to an underestimation of the association between ALBI and PHLF; however, we showed no publication bias and, using the ‘trim and fill’ methods to further strengthen our results, we found that no hypothetical studies were missing in our analysis. Concerning the reference standard, two studies [38,44] considered as case groups only PHLF grades B and C, thus, introducing a misclassification bias, in particular for the definition of patients with PHLF, which could have been underestimated.

However, we carried out a sensitivity analysis after excluding these two studies [38,44], which showed no significant differences with our initial results. Most of the studies included considered the overall rate of PHLF, without distinguishing between PHLF grades; therefore, it was not possible to further stratify according to the presence of clinically significant PHLF, which could be more relevant in clinical practice [7]. At the same time, there were insufficient data to perform a subgroup analysis according to the extent of LR, which represents one of the other most relevant risk factors for PHLF.

Another weakness of our meta-analysis was the substantial heterogeneity between the studies included. Among the differences found within the included studies, one study [44] included a small subgroup of patients undergoing LR for reasons other than HCC. We performed a sensitivity analysis excluding this latter study [44], showing no significant differences in the estimated OR. In addition, we found variability in the extension of hepatectomy, which, as mentioned above, could have also influenced the occurrence rate of PHLF.

However, we further addressed the heterogeneity by performing a univariate meta-regression analysis that showed that none of the variables tested, including an extension of the hepatectomy, was able to explain the heterogeneity found. Last, most studies of the included in the present meta-analysis were carried out in China, and thus reported on HBV patients. The race and the etiology of the underlying liver disease may influence the tumor biology, thus, adding a further bias to the surgical outcomes.

Our meta-analysis has several strengths supporting its value, as it provided for first-time pooled estimates of studies assessing the association between the ALBI grade and the occurrence of PHLF. Among the strengths of this meta-analysis, we performed a comprehensive literature search that minimized the risk of missing studies and, in the case of missing data, we contacted the authors to improve the data extraction. Another strength of our meta-analysis was the good methodological quality of the studies included. Despite the inclusion of only seven studies, we were able to include a large number of patients (5377) who underwent LR, with a reported PHLF rate of 13.4%. Of these, 69.9% were clinically significant, which is, thus, in line with other studies reporting the occurrence of PHLF [5,57].

In conclusion, our results provide additional evidence that the pre-operative ALBI grade is associated with the occurrence of PHLF. This has prognostic value for predicting this severe complication. The ALBI grade is a non-invasive, blood-test-based simple score that is able to further stratify the individual prognosis of chronic liver disease patients undergoing LR and reduce post-operative complications, such as PHLF. Further well-designed high-quality studies for evaluating the accuracy of the ALBI grade in the prediction of PHLF are needed.

Supplementary Materials: The following are available online at https://www.mdpi.com/article/10.3390/jcm10092011/s1, Supplementary material 1: PRISMA Checklist; Supplementary material 2: Electronic search strategy of the literature, Supplementary material 3: Criteria for rating the method-
ological quality of the included studies (QUADAS-2), Supplementary material 4: Methodological quality assessment of the included studies.

**Author Contributions:** Conceptualization, G.M. (Giovanni Marasco); methodology, G.M. (Giovanni Marasco); software, G.M. (Giovanni Marasco); validation, G.M. (Giovanni Marasco) and L.V.A.; formal analysis, G.M. (Giovanni Marasco) and L.V.A.; investigation, G.M. (Giovanni Marasco), L.V.A., A.C., D.F., F.B., G.M. (Giuseppe Mazzella), M.M., F.A.; resources, F.A.; data curation, G.M. (Giovanni Marasco), L.V.A., A.C., D.F., F.B., G.M. (Giuseppe Mazzella), M.M., F.A.; writing—original draft preparation, G.M. (Giovanni Marasco) and L.V.A.; writing—review and editing, G.M. (Giovanni Marasco), L.V.A., A.C., D.F., F.B., G.M. (Giuseppe Mazzella), M.M., F.A.; visualization, G.M. (Giovanni Marasco) and F.A.; supervision, A.C., D.F., F.B., G.M. (Giuseppe Mazzella), M.M., F.A.; project administration, G.M. (Giovanni Marasco); funding acquisition, F.A. All authors have read and agreed to the published version of the manuscript.

**Funding:** This research received no external funding.

**Institutional Review Board Statement:** Ethical review and approval were waived for this study, due to the use of already available published data.

**Informed Consent Statement:** Informed consent was obtained from all subjects involved in the study by the investigator of each published study included in the present systematic review and meta-analysis.

**Data Availability Statement:** The data presented in this study are openly available in Medline and Embase.

**Conflicts of Interest:** The authors declare no conflict of interest.

**References**

1. McGlynn, K.A.; Petrick, J.L.; London, W.T. Global Epidemiology of Hepatocellular Carcinoma: An Emphasis on Demographic and Regional Variability. *Clin. Liver Dis.* 2015, 19, 223–238. [CrossRef]

2. Kulik, L.; El-Serag, H.B. Epidemiology and Management of Hepatocellular Carcinoma. *Gastroenterology* 2019, 156, 477–491.e1. [CrossRef]

3. Galle, P.R.; Forner, A.; Llovet, J.M.; Mazzaferro, V.; Piscaglia, F.; Raoul, J.-L.; Schirmacher, P.; Vilgrain, V. EASL Clinical Practice Guidelines: Management of post-hepatic post-hepatic carcinoma. *J. Hepatol.* 2018. [CrossRef] [PubMed]

4. Van Den Broek, M.A.J.; Olde Damink, S.W.M.; Dejong, C.H.C.; Lang, H.; Malagó, M.; Jalan, R.; Saner, F.H. Liver failure after partial hepatic resection: Definition, pathophysiology, risk factors and treatment. *Liver Int.* 2008, 28, 767–780. [CrossRef]

5. Søreide, J.A.; Deshpande, R. Post hepatectomy liver failure (PHLF)—Recent advances in prevention and clinical management. *Eur. J. Surg. Oncol.* 2020, 47, 216–224. [CrossRef]

6. Sultana, A.; Brooke-Smith, M.; Ullah, S.; Figueras, J.; Rees, M.; Vauthey, J.N.; Conrad, C.; Hugh, T.J.; Garden, O.J.; Fan, S.T.; et al. Prospective evaluation of the International Study Group for Liver Surgery definition of post hepatectomy liver failure after liver resection: An international multicentre study. *HPB* 2018, 20, 462–469. [CrossRef]

7. Rahbari, N.N.; Garden, O.J.; Padbury, R.; Brooke-Smith, M.; Crawford, M.; Adam, R.; Koch, M.; Makuuchi, M.; Dematteo, R.P.; Christophi, C.; et al. Posthepatectomy liver failure: A definition and grading by the International Study Group of Liver Surgery (ISGLS). *Surgery 2011, 149, 713–724. [CrossRef] [PubMed]

8. Marasco, G.; Colecchia, A.; Milandri, M.; Rossini, B.; Alemanni, L.V.; Dajti, E.; Ravaioi, F.; Renzulli, M.; Golferi, R.; Festi, D. Non-invasive tests for the prediction of post-hepatectomy liver failure in the elderly. *Hepatoma Res.* 2020, 2020. [CrossRef]

9. Cucchi, A.; Cescon, M.; Golferi, R.; Piscaglia, F.; Renzulli, M.; Neri, F.; Cappelli, A.; Mazzotti, F.; Mosconi, C.; Colecchia, A.; et al. Hepatic venous pressure gradient in the preoperative assessment of patients with resectable hepatocellular carcinoma. *J. Hepatol.* 2016, 64, 79–86. [CrossRef] [PubMed]

10. Marasco, G.; Colecchia, A.; Dajti, E.; Ravaioi, F.; Cucchi, A.; Cescon, M.; Festi, D. Prediction of posthepatectomy liver failure: Role of SSM and LSPS. *J. Surg. Oncol. 2019, 119, 400–401. [CrossRef] [PubMed]

11. Ray, S.; Mehta, N.N.; Golhar, A.; Nundy, S. Post hepatectomy liver failure—A comprehensive review of current concepts and controversies. *Ann. Med. Surg. 2018, 34, 4–10. [CrossRef]

12. Wang, Y.-Y.; Zhao, X.-H.; Ma, L.; Ye, J.-Z.; Wu, F.-X.; Tang, J.; You, X.-M.; Xiang, B.-D.; Li, L.-Q. Comparison of the ability of Child-Pugh score, MELD score, and ICG-R15 to assess preoperative hepatic functional reserve in patients with hepatocellular carcinoma. *J. Surg. Oncol. 2018, 118, 440–445. [CrossRef]

13. Cescon, M.; Colecchia, A.; Cucchi, A.; Peri, E.; Montrone, L.; Ercolani, G.; Festi, D.; Pinna, A.D. Value of Transient Elastography Measured With Fibroscan in Predicting the Outcome of Hepatic Resection for Hepatocellular Carcinoma. *Ann. Surg. 2012, 256, 706–713. [CrossRef] [PubMed]
14. Johnson, P.J.; Berhane, S.; Kagayashiki, C.; Satomura, S.; Teng, M.; Reeves, H.L.; O’Beirne, J.; Fox, R.; Skowronska, A.; Palmer, D.; et al. A assessment of liver function in patients with hepatocellular carcinoma: A new evidence-based approach—The albi grade. *J. Clin. Oncol.* 2015. [CrossRef] [PubMed]

15. Wang, J.; Zhang, Z.; Yan, X.; Li, M.; Xia, J.; Liu, Y.; Chen, Y.; Jia, B.; Zhu, L.; Zhu, C.; et al. Albumin-Bilirubin (ALBI) as an accurate and simple prognostic score for chronic hepatitis B-related liver cirrhosis. *Dig. Liver Dis.* 2019, 51, 1172–1178. [CrossRef] [PubMed]

16. Xu, Y.X.; Wang, Y.B.; Tan, Y.L.; Xi, C.; Xu, X.Z. Prognostic value of pretreatment albumin to bilirubin ratio in patients with hepatocellular cancer: A meta-analysis. *Medicine* 2019, 98, e14027. [CrossRef] [PubMed]

17. Handbook for DTA Reviews | Cochrane Screening and Diagnostic Tests. Available online: http://methods.cochrane.org/sdt/handbook-dta-reviews (accessed on 29 May 2018).

18. Moher, D.; Liberati, A.; Tetzlaff, J.; Altman, D.G.; Altman, D.; Antes, G.; Atkins, D.; Barbour, V.; Barrowman, N.; Berlin, J.A.; et al. Preferred reporting items for systematic reviews and meta-analyses: The PRISMA statement. *Ann. Intern. Med.* 2009, 151, 264–269. [CrossRef] [PubMed]

19. Whiting, P.F.; Rutjes, A.W.S.; Westwood, M.E.; Mallett, S.; Deeks, J.J.; Reitsma, J.B.; Leeflang, M.M.G.; Sterne, J.A.C.; Bossuyt, P.M.M. Quadast-2: A revised tool for the quality assessment of diagnostic accuracy studies. *Ann. Intern. Med.* 2011, 155, 529–536. [CrossRef] [PubMed]

20. Ruzzene, A.; Del Angelis, M.; Concì, S.; Campagnaro, T.; Isà, G.; Bagante, F.; Ciangerottì, A.; Pedrazzani, C.; Capelli, P.; Iacono, C.; et al. The albumin-bilirubin score stratifies the outcomes of Child-Pugh class A patients after resection of hepatocellular carcinoma. *Surg. Oncol.* 2019, 28, 78–85. [CrossRef] [PubMed]

21. Manly, B.F.J. Randomization, Bootstrap and Monte Carlo Methods in Biology; Chapman and Hall/CRC: Boca Raton, FL, USA, 2018; Chapter 4.

22. Steichen, T. Nonparametric trim and fill analysis of publication bias in meta-analysis: Erratum. *Stata J.* 2001, 10.

23. Thompson, S.G.; Sharp, S.J. Explaining heterogeneity in meta-analysis: A comparison of methods. *Stat. Med.* 1999, 18, 2693–2708. [CrossRef] [PubMed]

24. Higgs, J.P.T.; Thompson, S.G. Controlling the risk of spurious findings from meta-regression. *Stat. Med.* 2004, 23, 1663–1682. [CrossRef] [PubMed]

25. Mai, R.Y.; Wang, Y.Y.; Bai, T.; Chen, J.; Xiang, B.D.; Wu, G.B.; Wu, F.X.; Li, L.Q.; Ye, J.Z. Combination of ALBI and APRI to predict posthepatectomy liver failure after liver resection for HBV-related HCC patients. *Cancer Manag. Res.* 2019, 11, 8799–8806. [CrossRef]

26. Cai, W.; He, B.; Hu, M.; Zhang, W.; Xiao, D.; Yu, H.; Song, Q.; Xiang, N.; Yang, J.; He, S.; et al. A radiomics-based nomogram for the preoperative prediction of posthepatectomy liver failure in patients with hepatocellular carcinoma. *World J. Surg.* 2020, 44, 4197–4206. [CrossRef] [PubMed]

27. Shi, J.Y.; Sun, L.Y.; Quan, B.; Xing, H.; Li, C.; Liang, L.; Pawlik, T.M.; Zhou, Y.H.; Wang, H.; Gu, W.M.; et al. A novel online calculator based on noninvasive variables (ALBI and APRI) for predicting post-hepatectomy liver failure in patients with hepatocellular carcinoma. *Clin. Res. Hepatol. Gastroenterol.* 2020. [CrossRef]

28. Sun, L.-Y.; Zhu, H.; Diao, Y.-K.; Xing, H.; Liang, L.; Li, J.; Zhou, Y.-H.; Gu, W.-M.; Chen, T.-H.; Zeng, Y.-Y.; et al. A novel online calculator based on albumin-bilirubin and aspartate transaminase-to-platelet ratio index for predicting postoperative morbidity following hepatectomy for hepatocellular carcinoma. *Ann. Transl. Med.* 2020, 8, 1591. [CrossRef]

29. Xu, Y.; Hu, X.; Li, J.; Dong, R.; Bai, X. An improved scoring system based on PALBI in predicting post-hepatectomy liver failure outcomes. *Dig. Dis.* 2020. [CrossRef] [PubMed]

30. Zou, H.; Wen, Y.; Yuan, K.; Miao, X.Y.; Xiong, L.; Liu, K.J. Combining albumin-bilirubin score with future liver remnant predicts post-hepatectomy liver failure in HBV-associated HCC patients. *Liver Int.* 2018, 38, 494–502. [CrossRef]

31. Zou, H.; Yang, X.; Li, Q.L.; Zhou, Q.X.; Xiong, L.; Wen, Y. A Comparative Study of Albumin-Bilirubin Score with Child-Pugh Score, Model for End-Stage Liver Disease Score and Indocyanine Green R15 in Predicting Posthepatectomy Liver Failure for Hepatocellular Carcinoma Patients. *Dig. Dis.* 2018, 36, 236–243. [CrossRef] [PubMed]
37. Wang, L.; Xie, L.; Zhang, N.; Zhu, W.; Zhou, J.; Pan, Q.; Mao, A.; Lin, Z.; Wang, L.; Zhao, Y. Predictive Value of Intraoperative Indocyanine Green Clearance Measurement on Postoperative Liver Function After Anatomic Major Liver Resection. J. Gastrointest. Surg. 2019. [CrossRef]

38. Lu, L.H.; Zhang, Y.F.; Mu-Yuan, C.; Kan, A.; Zhong, X.P.; Mei, J.; Ling, Y.H.; Li, S.H.; Shi, M.; Wei, W.; et al. Platelet-albumin-bilirubin grade: Risk stratification of liver failure, prognosis after resection for hepatocellular carcinoma. Dig. Liver Dis. 2019, 51, 1430–1437. [CrossRef]

39. Andreatos, N.; Amini, N.; Gani, F.; Margonis, G.A.; Sasaki, K.; Thompson, V.M.; Bentrem, D.J.; Hall, B.L.; Pitt, H.A.; Wilson, A.; et al. Albumin-Bilirubin Score: Predicting Short-Term Outcomes Including Bile Leak and Post-hepatectomy Liver Failure Following Hepatic Resection. J. Gastrointest. Surg. 2017, 21, 238–248. [CrossRef] [PubMed]

40. Zhang, Z.Q.; Xiong, L.; Zhou, J.J.; Miao, X.Y.; Li, Q.L.; Wen, Y.; Zou, H. Ability of the ALBI grade to predict posthepatectomy liver failure and long-term survival after liver resection for different BCLC stages of HCC. Medical and Health Sciences 1112 Oncology and Carcinogenesis. World J. Surg. Oncol. 2018, 16. [CrossRef]

41. Fagenson, A.M.; Gleeson, E.M.; Pitt, H.A.; Lau, K.N. Albumin-Bilirubin Score vs Model for End-Stage Liver Disease in Predicting Post-Hepatectomy Outcomes. J. Am. Coll. Surg. 2020, 230, 637–645. [CrossRef]

42. Taylor, G.A.; Fagenson, A.M.; Kuo, L.E.; Pitt, H.A.; Lau, K.N. Predicting Outcomes of Surgery in Patients with Liver Disease: Albumin-Bilirubin Score vs Model for End-stage Liver Disease-Sodium Score. J. Am. Coll. Surg. 2020. [CrossRef]

43. Wang, Y.Y.; Zhong, J.H.; Su, Z.Y.; Huang, J.F.; Lu, S.D.; Xiang, B.D.; Ma, L.; Qi, L.N.; Ou, B.N.; Li, L.Q. Albumin-bilirubin versus Child-Pugh score as a predictor of outcome after liver resection for hepatocellular carcinoma. Br. J. Surg. 2016, 103, 723–734. [CrossRef]

44. Sposito, C.; Monteleone, M.; Aldrighetti, L.; Cillo, U.; Dalla Valle, R.; Guglielmi, A.; Ettorre, G.M.; Ferrero, A.; Di Benedetto, F.; Rossi, G.E.; et al. Preoperative predictors of liver decompensation after mini-invasive liver resection. Surg. Endosc. 2020, 35. [CrossRef]

45. Chong, C.; Wong, G.; Fung, A.; Lok, H.-T.; Fong, A.; Cheung, S.; Wong, J.; Lee, K.-F.; Lai, P. Albumin-bilirubin versus liver stiffness measurement versus child’s pugh grade as a predictor of postoperative outcome following hepatectomy for hepatocellular carcinoma. HPB 2018, 20, S316. [CrossRef]

46. Fagenson, A.M.; Gleeson, E.M.; Karhadkar, S.; Di Carlo, A.; Pitt, H.A.; Lau, K.N. Comparison of Albumin-Bilirubin and Model for End-Stage Liver Disease in Predicting Post-Hepatectomy Liver Failure. J. Am. Coll. Surg. 2019, 229, S170–S171. [CrossRef]

47. Fukutomi, S.; Nomura, Y.; Muroya, D.; Goto, Y.; Sakai, H.; Akagi, Y.; Okuda, K. The validity of albumin-bilirubin model (ALBI) in predicting surgical outcome after hepatectomy for hepatocellular carcinoma-comparison to child pugh score. J. Hepatobiliary. Pancreat. Sci. 2017, 24, A289.

48. Imai, D.; Maeda, T.; Kayashima, H. Usefulness of ALBI grade in post-hepatectomy liver failure of early stage HCC. J. Hepatobiliary. Pancreat. Sci. 2017, 24, A147.

49. Shum, J.K.; Ng, S.W.Y.; Wong, B.; Chan, W.L.; Lok, H.T.; Fung, A.K.Y.; Cheung, Y.S.; Wong, J.; Lee, K.F.; Lai, P.B.S.; et al. Higher albumin-bilirubin grade and liver stiffness measurement were associated with more morbidities and mortality after hepatectomy for hepatocellular carcinoma. Surg. Pract. 2018, 22, 20.

50. Mai, R.Y.; Zeng, J; Ye, J.Z.; Su, Q.B.; Long, Z.R.; Shi, X.M.; Huang, S.; Wu, F.X.; Li, L.Q.; Lian, F.; et al. Clinical value of preoperative aspartate aminotransferase-to-platelet ratio index in predicting liver failure after hepatectomy for primary liver cancer. Acad. J. Second Mil. Med. Univ. 2019, 40, 61–67. [CrossRef]

51. Geng, L.; Zong, R.; Shi, Y.; Xu, K. Prognostic role of preoperative albumin-bilirubin grade on patients with hepatocellular carcinoma after surgical resection: A systematic review and meta-analysis. Eur. J. Gastroenterol. Hepatol. 2020, 32, 769–778. [CrossRef]

52. Mohammadi, H.; Abuodeh, Y.; Jin, W.; Frakes, J.; Friedman, M.; Bieber, B.; Choi, J.; El-Haddad, G.; Kis, B.; Sweeney, J.; et al. Using the Albumin-Bilirubin (ALBI) grade as a prognostic marker for radioembolization of hepatocellular carcinoma. J. Gastrointest. Oncol. 2018, 9, 840–846. [CrossRef]

53. Na, S.K.; Yim, S.Y.; Suh, S.J.; Jung, Y.K.; Kim, J.H.; Seo, Y.S.; Yim, H.J.; Yeon, J.E.; Byun, K.S.; Um, S.H. ALBI versus Child-Pugh grading systems for liver function in patients with hepatocellular carcinoma. J. Surg. Oncol. 2018, 117, 912–921. [CrossRef]

54. Xu, L.; Wu, J.; Lu, W.; Yang, C.; Liu, H. Application of the Albumin-Bilirubin Grade in Predicting the Prognosis of Patients With Hepatocellular Carcinoma: A Systematic Review and Meta-Analysis. Transplant. Proc. 2019. [CrossRef] [PubMed]

55. Maluccio, M.; Covey, A. Recent progress in understanding, diagnosing, and treating hepatocellular carcinoma. CA Cancer J. Clin. 2012, 62, 394–399. [CrossRef] [PubMed]

56. Kamath, P.S.; Wiesner, R.H.; Malinchoc, M.; Kremers, W.; Therneau, T.M.; Kosberg, C.L.; D’Amico, G.; Dickson, E.R.; Kim, W.R. A model to predict survival in patients with end-stage liver disease. Hepatology 2001, 33, 464–470. [CrossRef] [PubMed]

57. Fukushima, K.; Fukushima, T.; Kuramitsu, K.; Kido, M.; Takebe, A.; Tanaka, M.; Itoh, T.; Ku, Y. Assessment of ISGLS Definition of Posthepatectomy Liver Failure and Its Effect on Outcome in Patients with Hepatocellular Carcinoma. J. Gastrointest. Surg. 2014, 18, 729–736. [CrossRef]