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Liver resection versus radiofrequency ablation for solitary

small hepatocellular carcinoma measuring≤3cm: a

systematic review and meta-analysis

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Summary Statement

Curative intent surgical resection is the best option for achieving long-term survival outcomes in patients with solitary HCC \leq 3cm.

Author's contribution

Ming, Gangjun, Ting and Wentao designed the study and performed the literature search, study selection, data extraction and statistical analyses. Ming, Guangjun and Youwei performed the statistical analyses. Gangjun, Kunlin and Ming wrote the first draft of the manuscript. All authors contributed to the interpretation of the data and critically reviewed the manuscript.

Conflicts of interest

The authors disclose no potential conflicts of interest.

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Abstract:

Background:

Controversy remains regarding liver resection (LR) and radiofrequency ablation (RFA) for patients with single hepatocellular carcinomas (HCCs) measuring 3 cm or less. The purpose of our study was to compare the prognosis between LR and RFA in patients with solitary HCCs \leq 3 cm.

Methods:

The meta-analysis followed the PRISMA guidelines and the Cochrane Handbook. All RCTs and cohort studies that compared LR versus RFA in patients with solitary HCCs≤3 cm were comprehensively searched in the PubMed, Cochrane Library, Embase, and Web of Science databases up to January 30, 2024. The primary endpoints were overall survival (OS), recurrence-free survival (RFS), and disease-free survival (DFS).

Results:

A total of 6356 patients with solitary HCCs \leq 3 cm and 5829 patients with solitary HCCs \leq 2 cm from 39 included studies were analyzed (LR = 5759, RFA = 6426). The present meta-analysis of two RCTs showed no statistically significant difference in OS between LR and RFA. However, the meta-analysis of cohort studies revealed that, compared with RFA, LR conferred a superior OS advantage (HR = 0.80, 95% CI: 0.68–0.93, P = 0.005). There was a significant improvement in the DFS rate with LR over RFA (HR=0.63, 95% CI: 0.49–0.81) and in the RFS rate (HR=0.65, 95% CI: 0.55–0.76). Compared with RFA, LR resulted in better OS (HR=0.73, 95% CI: 0.54–0.97), DFS (HR=0.74, 95% CI: 0.67–0.82) and RFS (HR=0.71, 95% CI: 0.57–0.90) in patients with a solitary HCC lesion \leq 2 cm.

Conclusions:

Evidence from cohort studies suggested that in patients with a solitary HCC lesion ≤ 3 cm, LR is preferable to RFA. Additional RCTs are needed to confirm the validity of

this evidence.

Keywords:

Hepatocellular carcinoma; Radiofrequency ablation; Liver resection; Survival

Introduction

There were close to 800,000 deaths from liver cancer in 2022^{1} . Hepatocellular carcinoma (HCC) is the dominant histological subtype of liver malignancy and accounts for approximately 75% of the total liver cancer burden worldwide². The management of HCC involves a multidisciplinary approach that can include surgery, chemotherapy/targeted therapy, and radiation therapies. Surgical intervention continues to be the foremost therapeutic approach for these patients, encompassing a range of options such as liver resection (LR), radiofrequency ablation (RFA), or liver transplantation (LT). The Barcelona Clinic Liver Cancer (BCLC) system was used for tumor staging and to guide treatment strategies³. Patients with solitary HCCs ≤ 2 and 3 cm in size were defined as BCLC 0 and A, respectively. According to the 2022 BCLC guidelines described here, for patients with a solitary HCC <2 cm, LT is recommended as the primary therapy, with RFA recommended as a secondary treatment option; for patients with a solitary HCC ≤3 cm, LR is recommended. LT is an effective treatment for cirrhosis and early-stage tumors because it eliminates the tumor and the cirrhotic tissue simultaneously. However, there is an organ shortage and a large number of ineligible patients.

Both LR and RFA are recommended as primary treatment strategies in several international guidelines for solitary small HCC, including the 5th JSH-HCC Guidelines ⁴, AASLD Practice Guidance⁵, and 2022 Chinese clinical guidelines⁶. Analyses were performed for the subgroup of patients with solitary small HCC lesions in three randomized controlled trials (RCTs)⁷⁻⁹, and the findings did not confirm that LR or RFA was associated with better outcomes. Furthermore, the results of direct comparisons in terms of survival between LR and RFA in patients with single small HCC lesions remain unclear. Wu et al. reported that, compared with RFA, LR offers the best chance for better recurrence-free survival (RFS)¹⁰. Li et al. reported no statistically significant differences in disease-free survival (DFS) between patients who underwent LR and those who underwent RFA¹¹.

Previous meta-analyses compared the outcomes between LR and RFA in treating small HCCs, but these studies included a heterogeneous population, ranging from solitary nodules <5 cm to up to 3 nodules with a maximum diameter of <3 cm for each nodule^{12,13}. The purpose of this study was to retrieve the available evidence and perform a meta-analysis comparing the prognoses of LR and RFA for solitary small HCCs≤3 cm to improve the quality of evidence.

Materials and methods

This study was conducted in strict adherence to the Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA)¹⁴, and Assessing the

Methodological Quality of Systematic Reviews (AMSTAR) guidelines¹⁵. The protocol of this meta-analysis was registered on PROSPERO.

Data sources and search strategy

Two reviewers independently conducted a comprehensive search of major databases, including PubMed, Embase, Web of Science, and the Cochrane Library, from database inception to March 31, 2024. We used a combination of the terms "single hepatocellular carcinoma", "early-stage liver cancer", and "early-stage liver cancer" or "radiofrequency ablation". A limit was set on cohort studies that were designed to compare outcomes between RFA and surgical resection for solitary HCCs≤3 cm, and only English-language articles were included. No restrictions were placed on publication dates or regions during the search.

During the initial screening phase, researchers excluded articles that were obviously not relevant to the research topic by reading their titles and abstracts. The full texts of the articles were screened for eligibility. Any disagreements were resolved through consultation and discussion among all the authors.

Inclusion criteria

Eligible studies were screened on the basis of the following inclusion criteria: (1) study design: the trials compared liver resection with radiofrequency ablation for the treatment of HCC; (2) the diagnosis of HCC was established on the basis of pathological or clinical findings; (3) the studies included in this meta-analysis focused on solitary HCC, with each lesion having a maximum diameter of 3 cm or less; (4) there was no prior treatment history for HCC; (5) there was no evidence of extrahepatic tumor metastasis or macrovascular invasion; (6) there was no concomitant malignant disease other than HCC; (7) the patients were all aged 18 years or older; and (8) the evaluation of treatment outcome endpoints was conducted via the following standardized definitions: overall survival (OS), disease-free survival (DFS), and recurrence-free survival (RFS).

Exclusion criteria

The exclusion criteria were as follows: (1) studies included patients with recurrent HCC; (2) medical records were not available for analysis; and (3) the study types included comments, case reports, reviews, conference abstracts and letters.

Data extraction

Two authors independently extracted relevant study characteristics and data via a data extraction form. The following data were recorded: baseline information (author, year, location, study period, study design), participant characteristics (number of patients, age, sex, etiology of liver disease, Child–Pugh score), tumor size, and tumor location. We subsequently extracted hazard ratios (HRs) with 95% confidence intervals (CIs) associated with the primary outcomes (OS, DFS and RFS). We resolved any disagreements in the opinion of the data extracted through discussion. If the HR and 95% CI could not be directly obtained, they were estimated from Kaplan–Meier

curves¹⁶.

Quality assessment

The risk of bias of the included trials was assessed in accordance with guidelines from the Cochrane Handbook for Systematic Reviews of Interventions¹⁷. We used the Newcastle–Ottawa scale (NOS)¹⁸, which is a nine-point scale comprising three items, patient selection, study group comparability, and outcome, for retrospective cohort studies to assess study quality. Studies scoring 7–9 points were considered to have a low risk of bias.

Statistical analysis

The primary endpoints of the meta-analysis were OS, RFS, and DFS. RevMan software (version 5.3, Cochrane Collaboration) was used to calculate the hazard ratio (HR) and 95% confidence interval (CI) in this meta-analysis. Heterogeneity was evaluated by the Cochran Q statistic and I2 test. If the P value of Cochran's Q statistic was less than or equal to 0.10, heterogeneity was present. A low inconsistency was indicated by an I² statistic of less than 25%. A value of 25% to 75% indicated moderate inconsistency, and a value greater than 75% indicated high inconsistency among studies. When statistical heterogeneity was evident, a random effects model (Cochrane Handbook) was adopted.

Publication bias

Publication bias was assessed for the primary endpoints via funnel plots. Funnel plot asymmetry was estimated by Egger regression, with a p value<0.05 indicating asymmetry. Funnel plots and Egger regression were generated with RevMan 5.3 and Stata 15.1 software, respectively.

Results

Included studies and study characteristics

The PRISMA flow chart describing the inclusion process is illustrated in Figure 1. We initially retrieved a total of 10225 records from the abovementioned databases: PubMed (4851 records), Embase (213 records), Web of Science (4991 records), and the Cochrane Library (170 records). After duplicate removal, 5013 records were further reviewed. Altogether, 4140 records were excluded at the title/abstract level, and 748 were summarized literature (e.g., review, report and commentaries) and therefore excluded.

The full-text articles of the remaining records were retrieved and assessed for inclusion by all reviewers. During further screening. 64 studies were excluded because they reported data on either single HCCs \leq 5 cm in diameter or multiple HCCs \leq 5 cm in diameter, with the inability to analyze data specifically for single HCCs < 3 cm on the basis of the original text (60 retrospective studies; 4 randomized controlled trials).

A total of 6356 patients with solitary HCCs≤3 cm and 5829 patients with solitary

HCCs≤2 cm from the 39 included studies were analyzed (LR = 5759, RFA = 6426)^{7-11,19-52}. No study included patients with vascular invasion or distant metastases. Eight studies^{29,34,35,37,40,45,47,52} used laparoscopic or robotic liver resection in the LR group, and two^{37,47} used laparoscopic RFA. Three studies were randomized controlled trials (RCTs)⁷⁻⁹; the remaining 36 were retrospective studies (RSs)^{10,11,19-52}. The studies included in the meta-analysis were from China^{7,8,10,11,24,32,39,42-46,50,52}, Korea^{21,25-28,30,31,34,37,38,40,49,51}, Japan^{9,19,20,22,41,48}, Italy^{23,33,35,47}, France²⁹ and Canada³⁶. The Cochrane risk-of-bias tool revealed no high bias in 3 RCTs⁷⁻⁹. The overall quality of the 36 RSs^{10,11,19-52} was moderate; was moderate; the Newcastle–Ottawa Scale scores ranged from 7 to 9. A summary of the baseline information of patients in the included studies is presented in Table 1.

A total of 17 studies^{11,28,32-35,37-39,42,43,46-51} employed propensity score matching analysis to balance treatment-related characteristics, including patient age, sex, body mass index, race, Child–Pugh class, MELD score, liver function indicators (serum albumin, total bilirubin levels, aspartate aminotransferase, alanine aminotransferase), platelet count and tumor markers (serum alpha-fetoprotein and DCP level), and tumor size.

Survival outcomes of patients with solitary HCCs≤3 cm based on RCT data.

Few RCTs have compared survival between LR and RFA in patients with solitary HCC. The study population consisted of patients with HCC diameters ≤ 3 cm and ≤ 3 HCC nodules in the RCT of Takayama et al.⁹. In the study of Jiwei et al., the study population consisted of patients with a single HCC ≤ 5 cm or up to 3 nodules, each <3 cm⁸. The participants in the RCT of Minshan et al. consisted of patients with a solitary HCC smaller than 5 cm in diameter.

First, we reported the pooled OS from two RCTs^{7,8}. There was no significant difference in OS between the LR group and the RFA group (HR = 0.73, 95% CI: 0.26–2.05, P = 0.55), with no heterogeneity (I² = 56%, P = 0.90) (Supplementary Figure 1). A meta-analysis of RFS was not possible because only one RCT reported RFS⁹. In this RCT⁹, the median (range) post-enrollment follow-up duration was 5.04 (0.36–9.49) years for the surgery group and 4.99 (0.00–8.70) years for the RFA group. There were 135 patients with solitary HCC lesions \leq 3 cm who underwent surgical treatment, among whom 72 had postoperative recurrence; and there were 136 patients with solitary HCC lesions \leq 3 cm who received RFA treatment, among whom 74 had postoperative recurrence. Prognostic analysis demonstrated no significant difference in RFS between the two groups of patients.

Survival outcomes of patients with solitary HCCs≤3 cm based on RSs data. Overall survival

Propensity score matching was employed in 17 included RSs^{11,28,32-35,37-39,42,43,46-51} to reduce confounders of treatment allocation, and the remaining 19 included RSs^{10,19-27,29-31,36,40,41,44,45,52} that did not. Thus, the pooled data from the entire cohort of all the RSs^{10,11,19-52} indicated that the patients in the LR group had better OS than those in the RFA group (HR = 0.80, 95% CI: 0.68–0.93, P = 0.005), with high heterogeneity (I² =

58%, P<0.0001) (Figure 2).

Next, we assessed the OS of patients in all propensity score-matched cohorts. The pooled data from 14 RSs^{11,23,30,32,33,35,38,39,42,46,48-51} revealed a longer OS in the LR group (HR = 0.82, 95% CI: 0.72–0.95, P=0.006), with low heterogeneity (I2 = 23%, P=0.21) (Supplementary Figure 2).

Disease-Free Survival

Twelve $RSs^{11,20,21,28,31,34,37,39,41,43,50,52}$ reported data for the meta-analysis of DFS. A random-effects model was used to pool those RSs and demonstrated that LR was associated with better DFS (HR=0.63, 95% CI: 0.49–0.81, P = 0.0003), with high heterogeneity (I²=81%, P<0.00001) (Figure 3).

A meta-analysis encompassing six propensity score-matched cohorts^{11,27,33,35,43,50} revealed that SR significantly outperforms RFA in terms of RFS (HR = 0.64, 95% CI: 0.48–0.85, P = 0.002), with low heterogeneity ($I^2 = 0\%$, P=0.86) (Supplementary Figure 3).

Recurrence-free survival

Thirteen RSs^{10,20,21,28,31,34,37,39,44-46,48,51} reported data for the meta-analysis on RFS. A random-effects model was used to pool those data and demonstrated that LR was associated with better DFS (HR=0.65, 95% CI: 0.55–0.76, P<0.00001), with high heterogeneity ($I^2 = 81\%$, P=0.02) (Figure 4).

The RFS was also evaluated in eight propensity score-matched cohorts^{28,34,37-39,46,48,51}. RFS was reported to be longer in the LR group than in the RFA group (HR=0.64, 95% CI: 0.57–0.71, P<0.00001), with low heterogeneity ($I^2 = 0\%$, P=0.52) (Supplementary Figure 4).

Survival outcomes of patients with solitary HCCs≤2 cm

Overall survival

Fourteen RSs^{10,11,22,23,32,42,44,46-52} reported the prognostic data on patients with solitary HCCs \leq 2cm. On the basis of the results of the meta-analysis, patients with solitary HCCs \leq 2 cm in size who received LR vs. RFA had better OS (HR=0.73, 95% CI: 0.54–0.97, P = 0.03), with high heterogeneity (I² =69%, P<0.0001) (Supplementary Figure 5).

Ten propensity score-matched cohorts^{11,23,32,38,42,46,48-51} provided information on OS. Compared with the RFA group, the LR group had a longer OS (HR=0.78, 95% CI: 0.67–0.91, P=0.002), with low heterogeneity (I^2 =6%, P=0.38) (Supplementary Figure 6).

Disease-Free Survival

DFS in patients with a solitary HCC lesion ≤ 2 cm was reported in 6 RSs^{11,22,41,43,50,52}. The pooled data indicated that patients in the LR group had a better DFS (HR=0.74, 95% CI: 0.67–0.82, P<0.00001), with low heterogeneity (I² = 0%, P=0.47) (Supplementary Figure 7).

Three RSs provided data after propensity score matching, and the pooled result

revealed that LR was associated with increased RFS (HR=0.61, 95% CI: 0.46–0.80, P=0.0005), with high heterogeneity (I^2 =56%, P=0.10) (Supplementary Figure 8).

Recurrence-free survival

Six RSs⁵¹ provided information on RFS. Patients with solitary HCCs ≤ 2 cm in size in the LR group had longer RFS than those in the RFA group (HR=0.71, 95% CI: 0.57–0.90; P=0.004), with high heterogeneity (I² =50%, P=0.07) (Supplementary Figure 9).

A meta-analysis encompassing three propensity score-matched cohorts^{46,48,51} revealed that LR significantly outperforms RFA in terms of RFS (HR=0.61, 95% CI: 0.46–0.80, P=0.0005), with low heterogeneity ($I^2 = 56\%$, P=0.10) (Supplementary Figure 10).

Sensitivity analysis and publication bias

In the present meta-analysis, the funnel plots for survival outcomes of patients with solitary $HCCs \le 3$ cm from all propensity score-matched cohorts showed basic symmetry (Figure 5). No significant publication bias was further determined by Eegg's test, and all P values were greater than 0.05.

Discussion

To our knowledge, this meta-analysis included the largest number of studies comparing the prognosis between patients with solitary HCCs \leq 3 cm who underwent LR and those with RFA. Using the results of 36 RSs, we found that, compared with RFA, LR offers a better prognosis for patients with a solitary HCC lesion \leq 3 cm, including longer OS, DFS and RFS. A total of 7 RCTs^{7-9,53-56} on small HCC lesions were identified through literature screening, among which 3 provided data on single tumors \leq 3 cm. Therefore, only the data from these three studies were analyzed in this research. In addition, the results of the subgroup analysis (solitary HCC \leq 2 cm) suggested that LR at a very early stage (according to the BCLC system) provides a better prognosis, including longer OS, DFS and RFS.

Propensity score matching is a method that can reduce bias and help ensure the homogeneity of baseline data between groups⁵⁷. Propensity score matching can achieve reasonable matching between two groups of patients, yielding results similar to those of randomized controlled trials. Consequently, we concurrently analyzed the outcomes after propensity score matching to minimize the intergroup differences in variables. In these studies, propensity score matching was performed on the basis of demographic characteristics (such as age, sex, and body mass index) and preoperative features (such as liver function indices and tumor location). Despite the varying number of variables matched across these studies, the meta-analysis results indicate that propensity score matching has reduced the heterogeneity of the combined outcomes and that the trends observed are consistent with those without matching. As propensity score matching reduces the heterogeneity among the included studies, we believe that our research findings are more convincing than those of previous studies.

HCC is the main subtype of primary liver cancer and has a poor prognosis. Surgical resection remains the best opportunity for a cure, and only relatively few patients are

suitable for surgical treatment⁵⁸. According to the BCLC staging system and treatment guidelines, transplantation is the recommended treatment of choice for patients with BCLC early-stage disease³. When these patients are not given priority for liver transplantation, LR or RFA is often further performed. The long-term DFS and OS of patients with a solitary HCC lesion <3 cm are important when optimal treatment strategies are selected. The previous Cochrane meta-analysis by Majumdar et al. included four studies and revealed no benefit of LR or RFA on the survival of patients with early-stage HCC⁵⁹. One meta-analysis revealed that LR and RFA had similar effects on long-term survival in patients with very early-stage HCC⁶⁰. One retrospective cohort study reported that long-term OS was not significantly different between LR and RFA in patients with solitary HCCs≤3 cm³⁹. However, our metaanalysis including two RCTs revealed that the LR improved the OS of patients with solitary HCCs≤3 cm and solitary HCCs≤2 cm. This seemingly conflicts with conclusions drawn by previous studies. Previous studies have reported that the preoperative neutrophil-lymphocyte ratio (NLR) and the presence of microvascular invasion (MVI) are associated with long-term prognosis after surgical resection in patients with single and <5 cm HCC lesions^{61,62}. Therefore, the impact of pretreatment liver function and tumor marker levels on the prognosis of patients with solitary HCCs \leq 3 cm receiving different treatment regimens requires further analysis.

RFA is less invasive than LR and has great potential for local tumor control with minimal damage to healthy parenchyma⁶³. As a result, hospital length of stay and complication rates are also significantly lower than those of surgical resection⁶⁴. RFA is limited by an increased rate of local recurrence because of incomplete ablation⁵⁵. The efficacy of RFA can be affected by a variety of factors. A correlation between ablation margins and local tumor progression was noted for the first time by Kei et al. in 2008⁶⁵. The geometry of the local ablation zone and tumor location, such as the peritumoral ablative margin, adjacent blood vessel, and subcapsular location, are closely related to local tumor progression⁶⁶. Tissue perfusion and vascular-mediated cooling can lead to a reduced thermal effect of local ablation⁶⁶.

For patients with HCC who fulfill the Milan criteria and exhibit no vascular invasion, transplantation or LR are deemed the optimal therapeutic modalities⁶⁷. In a prospective randomized 168-patient trial, curative resection of small HCC lesions achieved a 3-year survival rate of 74.8%⁵⁵. The advantage of LR over RFA is that LR can completely remove the lesions where peritumoral micrometastases and microscopic vascular thromboses are detected⁶⁸. Our analysis of LR and RFA treatments for solitary small HCC lesions ($\leq 3 \text{ cm}/2 \text{ cm}$) indicated that LR was associated with significantly longer DFS and RFS. The low relapse rate of HCC may be attributed to total excision of the tumor. In addition, as the surgical technique has improved over time, the incidence of surgical complications has decreased. Several procedures have been proposed to improve the accuracy and safety of LR, such as real-time visualization by indocyanine green, computer-assisted surgical navigation, and robot-assisted surgery^{69,70}. Additionally, postoperative pathological analysis provides the necessary information to direct adjuvant therapy patients with a high risk of recurrence⁷¹. However, it is not easy to obtain complete pathological specimens

after RFA.

The findings of this meta-analysis should be interpreted with due caution, considering multiple factors that necessitate prudence. First, the number of RCTs included in this study was significantly lower than that of retrospective studies, and the results from the RCT subgroup analysis still require further evaluation through more well-defined and prospective clinical trials. Second, most of the data included in the present study were extracted from RSs. Despite the use of the propensity score matching method, deviations could not be eliminated as they can be with RCTs. Due to the absence of data regarding patients' liver function, oncological markers, and other pertinent parameters, a further subgroup analysis was not feasible. Third, the types of surgical approaches of the included studies included open, laparoscopic, or robotic surgery. Furthermore, RFA was performed via percutaneous, laparoscopic approaches. It has not been determined how heterogeneity influences the results. We expect that more RCTs will be designed to guide clinical treatment.

Conclusion

In conclusion, this meta-analysis revealed that the LR provided better OS, DFS, and RFS for patients with solitary small HCCs measuring ≤ 3 cm. However, these findings should be interpreted with caution because of the limited number of eligible RCTs included in this meta-analysis. Thus, large-scale, multicenter studies and well-designed RCTs on this topic are needed for further evaluation.

Availability of data and materials

The data sets used during the current study are available from the corresponding author on a reasonable request.

Provenance and peer review

Not commissioned, externally peer-reviewed.

Reference

- 1. Bray F, Laversanne M, Sung H, et al. Global cancer statistics 2022: GLOBOCAN estimates of incidence and mortality worldwide for 36 cancers in 185 countries. *CA: a cancer journal for clinicians*. 2024;74(3):229-263.
- 2. Petrick JL, Florio AA, Znaor A, et al. International trends in hepatocellular carcinoma incidence, 1978-2012. *Int J Cancer*. 2020;147(2):317-330.
- 3. Reig M, Forner A, Rimola J, et al. BCLC strategy for prognosis prediction and treatment recommendation: The 2022 update. *Journal of hepatology*. 2022;76(3):681-693.
- Hasegawa K, Takemura N, Yamashita T, et al. Clinical Practice Guidelines for Hepatocellular Carcinoma: The Japan Society of Hepatology 2021 version (5th JSH-HCC Guidelines). *Hepatol Res.* 2023;53(5):383-390.
- Singal AG, Llovet JM, Yarchoan M, et al. AASLD Practice Guidance on prevention, diagnosis, and treatment of hepatocellular carcinoma. *Hepatology*. 2023;78(6):1922-1965.

- 6. Xie DY, Zhu K, Ren ZG, Zhou J, Fan J, Gao Q. A review of 2022 Chinese clinical guidelines on the management of hepatocellular carcinoma: updates and insights. *Hepatobiliary Surg Nutr.* 2023;12(2):216-228.
- 7. Chen MS, Li JQ, Zheng Y, et al. A prospective randomized trial comparing percutaneous local ablative therapy and partial hepatectomy for small hepatocellular carcinoma. *Ann Surg.* 2006;243(3):321-328.
- 8. Huang J, Yan L, Cheng Z, et al. A randomized trial comparing radiofrequency ablation and surgical resection for HCC conforming to the Milan criteria. *Ann Surg.* 2010;252(6):903-912.
- 9. Takayama T, Hasegawa K, Izumi N, et al. Surgery versus Radiofrequency Ablation for Small Hepatocellular Carcinoma: A Randomized Controlled Trial (SURF Trial). *Liver Cancer*. 2022;11(3):209-218.
- Wu CC, Tseng CW, Tseng KC, et al. Radiofrequency ablation versus surgical resection for the treatment of solitary hepatocellular carcinoma 2 cm or smaller: A cohort study in Taiwan. J Formos Med Assoc. 2021;120(5):1249-1258.
- 11. Li YC, Chen PH, Yeh JH, et al. Clinical outcomes of surgical resection versus radiofrequency ablation in very-early-stage hepatocellular carcinoma: a propensity score matching analysis. *BMC Gastroenterol*. 2021;21(1):418.
- 12. Wang K, Wang R, Liu S, Peng G, Yu H, Wang X. Comparison of the safety and efficacy of hepatic resection and radiofrequency ablation in the treatment of single small hepatocellular carcinoma: systematic review and meta-analysis. *Transl Cancer Res.* 2022;11(3):580-590.
- Xuan D, Wen W, Xu D, Jin T. Survival comparison between radiofrequency ablation and surgical resection for patients with small hepatocellular carcinoma: A systematic review and meta-analysis. *Medicine (Baltimore)*. 2021;100(7):e24585.
- Moher D, Liberati A, Tetzlaff J, Altman DG. Preferred reporting items for systematic reviews and meta-analyses: the PRISMA statement. *PLoS Med.* 2009;6(7):e1000097.
- 15. Shea BJ, Grimshaw JM, Wells GA, et al. Development of AMSTAR: a measurement tool to assess the methodological quality of systematic reviews. *BMC medical research methodology*. 2007;7:10.
- 16. Tierney JF, Stewart LA, Ghersi D, Burdett S, Sydes MR. Practical methods for incorporating summary time-to-event data into meta-analysis. *Trials*. 2007;8:16.
- 17. Higgins JP, Altman Dg Fau Gøtzsche PC, Gøtzsche Pc Fau Jüni P, et al. The Cochrane Collaboration's tool for assessing risk of bias in randomised trials.343:5928.
- Stang A. Critical evaluation of the Newcastle-Ottawa scale for the assessment of the quality of nonrandomized studies in meta-analyses. *Eur J Epidemiol.* 2010;25(9):603-605.
- 19. Hiraoka A, Horiike N, Yamashita Y, et al. Efficacy of radiofrequency ablation therapy compared to surgical resection in 164 patients in Japan with single

hepatocellular carcinoma smaller than 3 cm, along with report of complications. *Hepatogastroenterology*. 2008;55(88):2171-2174.

- 20. Nishikawa H, Inuzuka T, Takeda H, et al. Comparison of percutaneous radiofrequency thermal ablation and surgical resection for small hepatocellular carcinoma. *BMC Gastroenterol.* 2011;11:143.
- 21. Yun WK, Choi MS, Choi D, et al. Superior long-term outcomes after surgery in child-pugh class a patients with single small hepatocellular carcinoma compared to radiofrequency ablation. *Hepatol Int.* 2011;5(2):722-729.
- 22. Imai K, Beppu T, Chikamoto A, et al. Comparison between hepatic resection and radiofrequency ablation as first-line treatment for solitary small-sized hepatocellular carcinoma of 3 cm or less. *Hepatol Res.* 2013;43(8):853-864.
- Pompili M, Saviano A, de Matthaeis N, et al. Long-term effectiveness of resection and radiofrequency ablation for single hepatocellular carcinoma ≤3 cm. Results of a multicenter Italian survey. *Journal of hepatology*. 2013;59(1):89-97.
- 24. Wong KM, Yeh ML, Chuang SC, et al. Survival comparison between surgical resection and percutaneous radiofrequency ablation for patients in Barcelona Clinic Liver Cancer early stage hepatocellular carcinoma. *Indian J Gastroenterol.* 2013;32(4):253-257.
- 25. Kim JM, Kang TW, Kwon CH, et al. Single hepatocellular carcinoma \leq 3 cm in left lateral segment: liver resection or radiofrequency ablation? *World journal of gastroenterology*. 2014;20(14):4059-4065.
- 26. Yang HJ, Lee JH, Lee DH, et al. Small single-nodule hepatocellular carcinoma: comparison of transarterial chemoembolization, radiofrequency ablation, and hepatic resection by using inverse probability weighting. *Radiology*. 2014;271(3):909-918.
- Kang TW, Kim JM, Rhim H, et al. Small Hepatocellular Carcinoma: Radiofrequency Ablation versus Nonanatomic Resection--Propensity Score Analyses of Long-term Outcomes. *Radiology*. 2015;275(3):908-919.
- 28. Kim GA, Shim JH, Kim MJ, et al. Radiofrequency ablation as an alternative to hepatic resection for single small hepatocellular carcinomas. *The British journal of surgery*. 2016;103(1):126-135.
- 29. Vitali GC, Laurent A, Terraz S, et al. Minimally invasive surgery versus percutaneous radio frequency ablation for the treatment of single small (≤3 cm) hepatocellular carcinoma: a case-control study. Surg Endosc. 2016;30(6):2301-2307.
- 30. Lee S, Kang TW, Cha DI, et al. Radiofrequency ablation vs. surgery for perivascular hepatocellular carcinoma: Propensity score analyses of long-term outcomes. *Journal of hepatology*. 2018;69(1):70-78.
- 31. Cha DI, Song KD, Kang TW, Lee MW, Rhim H. Small masses (≤3 cm) diagnosed as hepatocellular carcinoma on pre-treatment imaging: comparison of therapeutic outcomes between hepatic resection and radiofrequency ablation. *Br J Radiol.* 2020;93(1105):20190719.

- 32. Zheng L, Zhang CH, Lin JY, Song CL, Qi XL, Luo M. Comparative Effectiveness of Radiofrequency Ablation vs. Surgical Resection for Patients With Solitary Hepatocellular Carcinoma Smaller Than 5 cm. *Front Oncol.* 2020;10:399.
- Delvecchio A, Inchingolo R, Laforgia R, et al. Liver resection vs radiofrequency ablation in single hepatocellular carcinoma of posterosuperior segments in elderly patients. *World J Gastrointest Surg.* 2021;13(12):1696-1707.
- 34. Lee DH, Kim JW, Lee JM, et al. Laparoscopic Liver Resection versus Percutaneous Radiofrequency Ablation for Small Single Nodular Hepatocellular Carcinoma: Comparison of Treatment Outcomes. *Liver Cancer*. 2021;10(1):25-37.
- 35. Conticchio M, Delvecchio A, Ratti F, et al. Laparoscopic surgery versus radiofrequency ablation for the treatment of single hepatocellular carcinoma ≤3 cm in the elderly: a propensity score matching analysis. *HPB (Oxford)*. 2022;24(1):79-86.
- 36. Ivanics T, Rajendran L, Abreu PA, et al. Long-term outcomes of ablation, liver resection, and liver transplant as first-line treatment for solitary HCC of 3 cm or less using an intention-to-treat analysis: A retrospective cohort study. *Ann Med Surg (Lond).* 2022;77:103645.
- 37. Ko SE, Lee MW, Ahn S, et al. Laparoscopic Hepatic Resection Versus Laparoscopic Radiofrequency Ablation for Subcapsular Hepatocellular Carcinomas Smaller Than 3 cm: Analysis of Treatment Outcomes Using Propensity Score Matching. *Korean J Radiol.* 2022;23(6):615-624.
- 38. Lee J, Jin YJ, Shin SK, et al. Surgery versus radiofrequency ablation in patients with Child- Pugh class-A/single small (≤3 cm) hepatocellular carcinoma. *Clin Mol Hepatol.* 2022;28(2):207-218.
- 39. Zhang C, Gao R, Guo S, et al. Anatomic resection versus radiofrequency ablation with an ablative margin ≥ 1.0 cm for solitary small hepatocellular carcinoma measuring ≤ 3 cm: Comparison of long-term outcomes using propensity score matching analysis. *European journal of radiology*. 2022;155:110498.
- 40. Kang M, Cho JY, Han HS, et al. Comparative Study of Long-Term Outcomes of Laparoscopic Liver Resection versus Radiofrequency Ablation for Single Small Hepatocellular Carcinoma Located in Left Lateral Segments of the Liver. *Medicina (Kaunas).* 2023;59(6).
- 41. Takayama T, Makuuchi M, Hasegawa K. Single HCC smaller than 2 cm: surgery or ablation?: surgeon's perspective. *J Hepatobiliary Pancreat Sci.* 2010;17(4):422-424.
- 42. Hung HH, Chiou YY, Hsia CY, et al. Survival rates are comparable after radiofrequency ablation or surgery in patients with small hepatocellular carcinomas. *Clin Gastroenterol Hepatol.* 2011;9(1):79-86.
- 43. Wang JH, Wang CC, Hung CH, Chen CL, Lu SN. Survival comparison between surgical resection and radiofrequency ablation for patients in BCLC

very early/early stage hepatocellular carcinoma. *Journal of hepatology*. 2012;56(2):412-418.

- 44. Peng ZW, Lin XJ, Zhang YJ, et al. Radiofrequency ablation versus hepatic resection for the treatment of hepatocellular carcinomas 2 cm or smaller: a retrospective comparative study. *Radiology*. 2012;262(3):1022-1033.
- 45. Song J, Wang Y, Ma K, et al. Laparoscopic hepatectomy versus radiofrequency ablation for minimally invasive treatment of single, small hepatocellular carcinomas. *Surg Endosc.* 2016;30(10):4249-4257.
- 46. Liu PH, Hsu CY, Hsia CY, et al. Surgical Resection Versus Radiofrequency Ablation for Single Hepatocellular Carcinoma ≤ 2 cm in a Propensity Score Model. Ann Surg. 2016;263(3):538-545.
- 47. Santambrogio R, Bruno S, Kluger MD, et al. Laparoscopic ablation therapies or hepatic resection in cirrhotic patients with small hepatocellular carcinoma. *Digestive and liver disease : official journal of the Italian Society of Gastroenterology and the Italian Association for the Study of the Liver.* 2016;48(2):189-196.
- 48. Takayasu K, Arii S, Sakamoto M, et al. Impact of resection and ablation for single hypovascular hepatocellular carcinoma ≤2 cm analysed with propensity score weighting. *Liver international : official journal of the International Association for the Study of the Liver.* 2018;38(3):484-493.
- 49. Kim TH, Chang JM, Um SH, et al. Comparison of 2 curative treatment options for very early hepatocellular carcinoma: Efficacy, recurrence pattern, and retreatment. *Medicine (Baltimore)*. 2019;98(26):e16279.
- 50. Wang G, Zhang W, Tan Y, et al. The risk factors for long-term survival outcome in solitary hepatocellular carcinoma up to 2 cm: Propensity score matching analysis in a population cohort with a high rate of HBV infection. *International journal of surgery (London, England)*. 2019;72:1-6.
- 51. Chu HH, Kim JH, Kim PN, et al. Surgical resection versus radiofrequency ablation very early-stage HCC (≤2 cm Single HCC): A propensity score analysis. *Liver international : official journal of the International Association for the Study of the Liver.* 2019;39(12):2397-2407.
- 52. Lin CH, Ho CM, Wu CH, et al. Minimally invasive surgery versus radiofrequency ablation for single subcapsular hepatocellular carcinoma ≤ 2 cm with compensated liver cirrhosis. Surg Endosc. 2020;34(12):5566-5573.
- 53. Song J, Cao L, Ma K, et al. Laparoscopic liver resection versus radiofrequency ablation for small hepatocellular carcinoma: randomized clinical trial. *The British journal of surgery*. 2024;111(4).
- 54. Lee HW, Lee JM, Yoon JH, et al. A prospective randomized study comparing radiofrequency ablation and hepatic resection for hepatocellular carcinoma. *Annals of surgical treatment and research*. 2018;94(2):74-82.
- 55. Feng K, Yan J, Li X, et al. A randomized controlled trial of radiofrequency ablation and surgical resection in the treatment of small hepatocellular carcinoma. *Journal of hepatology*. 2012;57(4):794-802.

- 56. Fang Y, Chen W, Liang X, et al. Comparison of long-term effectiveness and complications of radiofrequency ablation with hepatectomy for small hepatocellular carcinoma. *Journal of gastroenterology and hepatology*. 2014;29(1):193-200.
- 57. Austin PC. Optimal caliper widths for propensity-score matching when estimating differences in means and differences in proportions in observational studies. *Pharmaceutical statistics*. 2011;10(2):150-161.
- 58. Llovet JM, Hernandez-Gea V. Hepatocellular carcinoma: reasons for phase III failure and novel perspectives on trial design. *Clinical cancer research : an official journal of the American Association for Cancer Research*. 2014;20(8):2072-2079.
- 59. Majumdar A, Roccarina D, Thorburn D, Davidson BR, Tsochatzis E, Gurusamy KS. Management of people with early- or very early-stage hepatocellular carcinoma: an attempted network meta-analysis. *Cochrane Database Syst Rev.* 2017;3(3):Cd011650.
- 60. Hu L, Lin J, Wang A, Shi X, Qiao Y. Comparison of liver resection and radiofrequency ablation in long-term survival among patients with early-stage hepatocellular carcinoma: a meta-analysis of randomized trials and high-quality propensity score-matched studies. *World journal of surgical oncology*. 2024;22(1):56.
- 61. Liao R, Tang ZW, Li DW, Luo SQ, Huang P, Du CY. Preoperative neutrophilto-lymphocyte ratio predicts recurrence of patients with single-nodule small hepatocellular carcinoma following curative resection: a retrospective report. *World journal of surgical oncology.* 2015;13:265.
- 62. Wei X, Jiang Y, Feng S, et al. Neoadjuvant intensity modulated radiotherapy for a single and small (≤5 cm) hepatitis B virus-related hepatocellular carcinoma predicted to have high risks of microvascular invasion: a randomized clinical trial. *International journal of surgery (London, England)*. 2023;109(10):3052-3060.
- 63. Lau WY, Leung TW, Yu SC, Ho SK. Percutaneous local ablative therapy for hepatocellular carcinoma: a review and look into the future. *Ann Surg.* 2003;237(2):171-179.
- 64. Lau WY, Lai EC. The current role of radiofrequency ablation in the management of hepatocellular carcinoma: a systematic review. *Ann Surg.* 2009;249(1):20-25.
- 65. Kei SK, Rhim H, Choi D, Lee WJ, Lim HK, Kim YS. Local tumor progression after radiofrequency ablation of liver tumors: analysis of morphologic pattern and site of recurrence. *AJR Am J Roentgenol.* 2008;190(6):1544-1551.
- 66. Hendriks P, Boel F, Oosterveer TT, et al. Ablation margin quantification after thermal ablation of malignant liver tumors: How to optimize the procedure? A systematic review of the available evidence. *Eur J Radiol Open*. 2023;11:100501.
- 67. EASL-EORTC clinical practice guidelines: management of hepatocellular carcinoma. *Journal of hepatology*. 2012;56(4):908-943.

- 68. Zhu P, Liao W, Ding ZY, et al. Intraoperative ultrasonography of robot-assisted laparoscopic hepatectomy: initial experiences from 110 consecutive cases. *Surg Endosc.* 2018;32(10):4071-4077.
- 69. Kato Y, Sugioka A, Kojima M, et al. Minimally Invasive Anatomic Liver Resection for Hepatocellular Carcinoma Using the Extrahepatic Glissonian Approach: Surgical Techniques and Comparison of Outcomes with the Open Approach and between the Laparoscopic and Robotic Approaches. *Cancers*. 2023;15(8).
- 70. Gavriilidis P, Edwin B, Pelanis E, et al. Navigated liver surgery: State of the art and future perspectives. *Hepatobiliary Pancreat Dis Int.* 2022;21(3):226-233.
- 71. Zhou Z, Qi L, Mo Q, et al. Effect of surgical margin on postoperative prognosis in patients with solitary hepatocellular carcinoma: A propensity score matching analysis. *J Cancer*. 2021;12(15):4455-4462.

	Cou ntry	Study type	No.of Patients		Patients				Cirrhosis(Present), (Entire cohort)		HBsAg (positive)		Anti-HCV (positive)		MELD score		CP(A/B/C)		AFP(ng/ml)		N O S
Study ID			L R	RF A	LR	RFA	LR	RFA	LR	RFA	LR	RFA	LR	RFA	LR	RFA	LR	RFA			
Hiraoka A et al, 2008	Japa n	RS	59	10 5	62.4±10.6	69.4±9. 1	NA	NA	NA	NA	44(74. 6%)	80(77. 1%)	NA	NA	54/5/0	79/26/ 0	427.8±131 7.6	114.5±319 .5	7		
Nishikawa H et al, 2011	Japa n	RS	69	16 2	67.4±9.7	68.4±8. 7	50(72.4%)	124(76.5 %)	8(11.6 %)	9(5.6 %)	51(73. 9%)	135(83 .3%)	NA	NA	45/0/0	102/0/ 0	376.7±198 9.8	74.7±181. 1	7		
Yun WK et al, 2011	Kore a	RS	21 5	25 5	51.7±9.7	57.0±9. 9	139(64.7 %)	201(78.8 %)	186(86 .5%)	192(75 .3%)	11(5.1 %)	45(17. 6%)	NA	NA	215/0/ 0	255/0/ 0	476.3±120 8.1	307.4±114 8.8	7		
Imai K et al, 2013	Japa n	RS	10 1	82	63.3±9.7	67.6±8. 5	NA	NA	34(33. 7%)	8(9.9 %)	52(51. 5%)	65(79. 3%)	NA	NA	97/4/0	60/22/ 0	171.8±586 .2	51.0±106. 9	7		
Pompili M et al, 2013	Italy	RS	24 6	29 8	67(41- 83)η	69(38- 85)η	46(18.6%)	40(13.8%	24(9.8 %)	32(10. 7%)	144(58 .5%)	213(71 .5%)	8(6- 16)η	8(6- 18)η	246/0/ 0	298/0/ 0	9(1- 9000)η	29(2- 2200)η	9		
Wong KM et al, 2013	Chin a	RS	46	36	55.1±12	63.5±1 3	NA	NA	18(39 %)	13(36 %)	26(57 %)	23(64 %)	NA	NA	46/0/0	36/0/0	204.0±733	95.1±174	8		
Kim JM et al, 2014	Kore a	RS	66	67	55(27- 76)η	59(39- 85)η	NA	NA	51(78. 5%)	44(65. 7%)	4(6.2 %)	17(25. 4%)	NA	NA	NA	NA	28.5(1- 7102)η	20.0(2- 5652)η	7		
Yang HJ et al, 2014	Kore a	RS	52	79	55.7±10.6	57.2±9. 2	29(55.8%	64(81.0%	34(65. 4%)	56(70. 9%)	6(11.5 %)	18(22. 8%)	8.3± 2.2	9.4±2 .9	50/2/0	68/11/ 0	NA	NA	8		
Kang TW et al, 2015	Kore a	RS	14 2	43 8	57.8±11.7	61.6±1 3.72	91(64.1%	355(81.1 %)	120(84 .5%)	324(74 .0%)	10(7.1 %)	64(14. 6%)	NA	NA	135/7/ 0	367/7 1/0	22.2(1.0– 5517.3)η	15.4(1.0– 3772.5)η	9		
Kim GA et al, 2016	Kore a	RS*	27 3	33 1	54.4±8·5	5.73±1 0.3	163(59.7 %)	189(57.1 %)	222(81 .3%)	246(74 .3%)	18(6.6 %)	40(12. 1%)	NA	NA	273/0/ 0	331/0/ 0	NA	NA	9		
Vitali GC et al, 2016	Fran ce	RS	45	60	61.4(31- 84)η	67.3(47 -83)η	NA	NA	NA	NA	NA	NA	NA	NA	40/5/0	45/15/ 0	NA	NA	7		
Lee S et al, 2018	Kore a	RS	18 2	10 1	53.8±9.2	57.0±1 0	88(48.4%	71(70.3%	156(85 .8%)	75(74. 3%)	13(7.1 %)	15(14. 8%)	7(7– 8)η	8(8– 10)η	NA	NA	35.4(7.2– 261.5)η	15.0(5.9– 90.5)η	8		
Cha DI et al, 2020	Kore a	RS	14 5	17 8	53.3±10	56.75± 9.5	NA	NA	123(84 .8%	131(73 .6%)	12(8.3 %)	27(15. 2%)	NA	NA	131/1 4/0	156/2 2/0	376.5±119 9.0	102.2±226 .3	8		
Zheng L et al, 2020	Chin a	RS*	33 2	37 9	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	7		
Delvecchio A et al, 2021	Italy	RS*	37	40	74.98(70- 83)η	74.5(70 -87)η	37(100%)	40(100%)	10(27 %)	5(12%)	19(50 %)	21(53 %)	6(6- 16)η	8(6- 15)η	30/7/0	37/3/0	12.5(2- 3900)η	5(1- 1988)η	8		
Lee DH et al, 2021	Kore a	RS*	25 1	31 5	57.5±9.3	60.8±9. 6	NA	NA	196(78 .1%)	234(74 .3%)	21(8.4 %)	34(10. 8%)	NA	NA	251/0/ 0	315/0/ 0	218.4±904 .2	72.9±211. 6	9		
Conticchio M et al, 2022	Italy	RS*	86	98	75.7(69.5 -86.5)η	75(70- 89)η	86(100%)	98(100%)	20(23 %)	6(6%)	49(57 %)	56(57 %)	6(6– 13)η	8(6– 18)η	77/9/0	84/14/ 0	NA	NA	8		
Ivanics T et al, 2022	Cana da	RS	25	83	64(55- 67)η	60(56- 66)η	NA	NA	17(68 %)	46(55 %)	7(28%	22(27 %)	7(6- 8)η	7(6- 8)η	25/0/0	79/4/0	700(400,9 200)δ	600(400,4 200)δ	7		

 Table 1 Patient demographics and baseline characteristics

Ko SE et al, 2022	Kore a	RS*	60	29	55.8±9	60±9.8	NA	NA	42(70. 0%)	18(62. 1%)	4(6.7 %)	2(6.9 %)	NA	NA	NA	NA	313.1±823 .2	13.9±25.0	9
Lee J et al, 2022	Kore a	RS*	23 2	15 9	>65(40%) L	>65(49 %) L	116(50.0 0%)	109(68.5 5%)	197(84 .9%)	105(66 .0%)	10(4.3 1%)	20(12, 58%)	NA	NA	232/0/ 0	159/0/ 0	129.2±357 .0	129.2±357 .0	8
Zhang C et al, 2022	Chin a	RS*	15 6	95	53.97±9.9 9	58.28± 10.04	NA	NA	140(89 .7)	6(80.0)	11(7.1 %)	13(13. 7%)	NA	NA	156/0/ 0	95/0/0	683.34±28 61.71	NA	9
Kang M et al, 2023	Kore a	RS	36	40	57.8±11.7 0	61.6±1 3.72	NA	NA	30(83. 3%)	31(77. 5%)	0(0%)	1(2.5 %)	7.36 ±1.7	8.55± 2.05	NA	NA	199.6±415 .6	29.4±68.2	7
Takayama T et al, 2010	Japa n	RS	12 35	13 15	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	1103/ 132/0	1014/ 301/0	NA	NA	7
Hung HH et al, 2011	Chin a	RS*	50	66	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	9
Wang JH et al, 2012	Chin a	RS*	52	91	NA	NA	NA	NA	34(65. 4%)	44(48. 4%)	14(26. 9%)	49(53. 8%)	NA	NA	NA	NA	NA	NA	9
Peng ZW et al, 2012	Chin a	RS	74	71	51.5±12.1	53.1±1 2.1	62(83.8%)	58(81.7%)	69(97 %)	70(95 %)	NA	NA	NA	NA	58/0/0	62/0/0	NA	NA	7
Song J et al, 2016	Chin a	RS	33	40	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	78/0/0	76/2/0	NA	NA	9
Liu PH et al, 2016	Chin a	RS*	10 9	12 8	60±13	64±12	NA	NA	65(60 %)	62(48 %)	37(34 %)	55(43 %)	7.8± 1.3	8.4±2 .5	NA	NA	145±262	92±247	9
Santambrogio R et al, 2016	Italy	RS*	76	76	66±9	68±8	76(100%)	76(100%)	NA	NA	NA	NA	8(7– 9)η	8(7– 10)η	76/0/0	76/0/0	58.9±183	101.6±498	9
Takayasu K et al, 2018	Japa n	RS*	17 6	49 1	NA	NA	NA	NA	24(13 %)	46(9%)	116(65 %)	401(89 %)	NA	NA	151/2 5/0	394/9 7/0	NA	NA	9
Kim TH et al, 2019	Kore a	RS*	52	10 2	56.4±9.1	61.6±1 0.3	36(69.2%)	92(90.2%)	38(73. 1%)	59(57. 8%)	6(11.5 %)	12(11. 8%)	7(7– 8)η	8(7– 10)η	52/0/0	102/0/ 0	19.1(4.3– 135.0)η	16.0(4.8– 77.9)η	8
Wang G et al, 2019	Chin a	RS*	19 2	81	52(42- 60)η	52(45- 62)η	NA	NA	177(92 .2%)	75(92. 6%)	NA	NA	NA	NA	176/1 6/0	62/19/ 0	11(4- 349)η	33(6- 348)η	9
Chu HH et al, 2019	Kore a	RS*	63 1	57 7	54(29- 77)η	58(29- 87)η	589(93.3 %)	528(91.5 %)	566(89 .7%)	456(79 %)	31(4.9 %)	68(11. 8%)	NA	NA	631/0/ 0	577/0/ 0	13.8(4.7- 112)η	21.2(5.1- 163)η	9
Lin CH et al, 2020	Chin a	RS	36	39	NA	NA	NA	NA	25(69. 44%)	25(64. 10%)	9(25%)	17(43. 59%)	NA	NA	36/0/0	39/0/0	1118.4±25 12.31ε	172.7±345 .81ε	8
Wu CC et al, 2021	Chin a	RS	83	73	NA	NA	40(66.7%)	47(81.0%)	26(31. 3%)	20(27. 4%)	33(39. 8%)	31(42. 5%)	NA	NA	79/4/0	66/7/0	NA	NA	8
Li YC et al, 2021	Chin a	RS*	10 3	85	57(23- 82)η	62(34- 81)η	54(52.4%)	61(71.8%)	51(49. 5%)	35(41. 2%)	49(47. 6%)	42(49. 4%)	NA	NA	100/3/ 0	79/6/0	NA	NA	9
Chen MS et al, 2006	Chin a	RCT	42	37	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	42/0/0	37/0/0	NA	NA	N A
Huang J et al, 2010	Chin a	RCT	45	57	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	N A
Takayama T et al, 2021	Japa n	RCT	13 5	13 6	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	NA	N A

Note: RS, Retrospective study; RCT, Randomized controlled trial; SR, surgical resection; RFA, radiofrequency ablation; anti-HCV, antibody to hepatitis C virus; HBsAg, hepatitis B virus surface antigen; CP, Child–Pugh classification; MELD, Model for End Stage Liver Disease; AFP, alpha-fetoprotein; NOS, Newcastle-Ottawa Scale; * Propensity score-matched; η Data were presented as median (range); ε The unit of data is u/l; δ Data were presented as median(interquartile range); \mathcal{L} The percentage of people over 65 years old; NA, Data were missing.

Figure legend

Figure 1 PRISMA Diagram showing identification of eligible studies and reasons for exclusion.

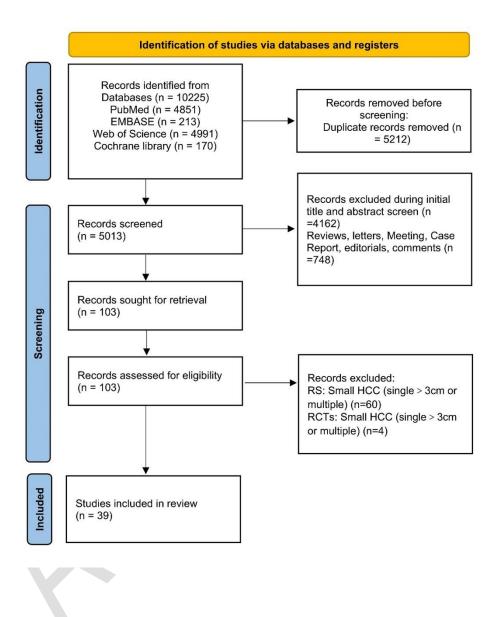


Figure2 Forest plot for hazard ratios of overall survival (OS) of patients with solitary $HCC \leq 3$ cm.

01. da a 0. karana	Is all second Daties	05	141-1-1-1	Hazard Ratio	Hazard Ratio
<u>Study or Subgroup</u> Cha DI et al, 2020	log[Hazard Ratio] -0.3496		Weight 5.0%	IV. Random. 95% C 0.70 [0.44, 1.13]	
Chu HH et al, 2020	-0.3552		5.0%	0.70 [0.44, 1.13]	-
Conticchio M et al, 2022	-0.0943		2.3%	0.91 [0.37, 2.24]	
Delvecchio A et al, 2021	-0.1278		1.5%	0.88 [0.27, 2.87]	
Hiraoka A et al, 2008		0.3353	3.5%	1.64 [0.85, 3.16]	
Hung HH et al, 2011	-0.9676		0.8%	0.38 [0.07, 2.06]	
lmai K et al, 2013	-0.3711		2.8%	0.69 [0.31, 1.54]	
lvanics T et al, 2022		0.628	1.4%	1.13 [0.33, 3.87]	
Kang M et al, 2023	-2.0326		0.9%	0.13 [0.02, 0.69]	
Kang TW et al, 2015		0.3537	3.3%	1.10 [0.55, 2.20]	
Kim JM et al, 2014		0.8876	0.8%	1.31 [0.23, 7.46]	
Kim TH et al, 2019	-0.4827		3.7%	0.62 [0.33, 1.16]	
Ko SE et al, 2022	0.802		1.4%	2.23 [0.65, 7.65]	
Lee S et al, 2018	-0.821	0.3537	3.3%	0.44 [0.22, 0.88]	
Li YC et al, 2021	-0.9465	0.256	4.7%	0.39 [0.23, 0.64]	
Lin CH et al, 2020	-2.3026	1.0818	0.5%	0.10 [0.01, 0.83]	
Liu PH et al, 2016	-0.4826	0.3272	3.6%	0.62 [0.33, 1.17]	
Nishikawa H et al, 2011	-0.0408	0.0523	8.3%	0.96 [0.87, 1.06]	1
Peng ZW et al, 2012	0.6429	0.3229	3.7%	1.90 [1.01, 3.58]	
Pompili M et al, 2013	-0.0513	0.1936	5.8%	0.95 [0.65, 1.39]	
Santambrogio R et al, 2016	-0.6505	0.2591	4.6%	0.52 [0.31, 0.87]	
Takayasu K et al, 2018	0.3646	0.166	6.4%	1.44 [1.04, 1.99]	
Vitali GC et al, 2016	-0.6733	0.6596	1.3%	0.51 [0.14, 1.86]	
Wang G et al, 2019	-0.4035	0.3098	3.9%	0.67 [0.36, 1.23]	
Wong KM et al, 2013	-0.8675	1.0859	0.5%	0.42 [0.05, 3.53]	· · · · · · · · · · · · · · · · · · ·
Wu CC et al, 2021	0.3456	0.3456	3.4%	1.41 [0.72, 2.78]	
Yang HJ et al, 2014	-0.5276	0.5781	1.6%	0.59 [0.19, 1.83]	
Yun WK et al, 2011	-1.0788	0.4905	2.1%	0.34 [0.13, 0.89]	
Zhang C et al, 2022	-0.1156	0.2102	5.5%	0.89 [0.59, 1.34]	
Zheng L et al, 2020	-0.3038	0.1976	5.7%	0.74 [0.50, 1.09]	
Total (95% CI)			100.0%	0.80 [0.68, 0.93]	•
Heterogeneity: Tau ² = 0.08; 0	Chi ² = 68.46, df = 29 (- < 0.000	01); l ² = 58	1%	
Test for overall effect: Z = 2.7					0.01 0.1 1 10 10 Favours [LR] Favours [RFA]

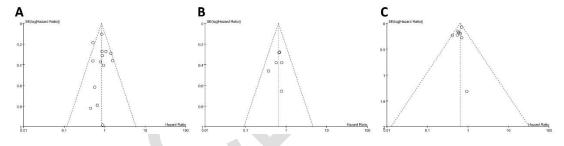
Figure3 Forest plot for hazard ratios of disease-free survival (DFS) of patients with solitary HCC≤3 cm.

				Hazard Ratio		Hazard Ratio		
Study or Subgroup	log[Hazard Ratio]	SE	Weight	IV, Random, 95% C	i	IV, Random, 95% Cl		
Cha DI et al, 2020	-0.2083	0.1788	10.9%	0.81 [0.57, 1.15]				
Kim GA et al, 2016	-0.6125	0.1157	12.4%	0.54 [0.43, 0.68]				
Ko SE et al, 2022	-0.6931	0.4426	5.3%	0.50 [0.21, 1.19]				
Lee DH et al, 2021	-0.1278	0.4023	5.9%	0.88 [0.40, 1.94]		· · · ·		
Li YC et al, 2021	-0.2992	0.2293	9.6%	0.74 [0.47, 1.16]				
Lin CH et al, 2020	-1.3093	1.1211	1.2%	0.27 [0.03, 2.43]		•		
Nishikawa H et al, 2011	-0.0121	0.0499	13.4%	0.99 [0.90, 1.09]				
Takayama T et al, 2010	-1.3093	1.1211	1.2%	0.27 [0.03, 2.43]	20 <u></u>			
Wang G et al, 2019	-0.4574	0.2303	9.6%	0.63 [0.40, 0.99]				
Wang JH et al, 2012	-0.7677	0.2898	8.1%	0.46 [0.26, 0.82]				
Yun WK et al, 2011	-0.8916	0.1594	11.4%	0.41 [0.30, 0.56]		-		
Zhang C et al, 2022	-0.512	0.1741	11.0%	0.60 [0.43, 0.84]		*		
Total (95% CI)			100.0%	0.63 [0.49, 0.81]		•		
Heterogeneity: Tau ² = 0.1	2; Chi ² = 57.26, df =	H		+	400			
Test for overall effect: Z =	3.61 (P = 0.0003)		85		0.01	0.1 1 1 Favours [LR] Favours [R	I0 FA]	100

Figure4 Forest plot for hazard ratios of recurrence-free survival (RFS) of patients with solitary HCC \leq 3 cm.

Study or Subaroup	log[Hazard Ratio]	SE	Weight	Hazard Ratio IV. Random. 95% Cl		Hazard Ratio IV. Random, 95% Cl	
Cha DI et al, 2020	-0.2083		9.8%	0.81 [0.57, 1.15]			
Chu HH et al, 2019	-0.4961	0.0776	16.0%	0.61 [0.52, 0.71]			
Kim GA et al, 2016	-0.6125	0.1157	13.6%	0.54 [0.43, 0.68]		*	
Ko SE et al, 2022	-0.6931	0.4426	2.8%	0.50 [0.21, 1.19]			
Lee DH et al, 2021	-0.1278	0.4023	3.3%	0.88 [0.40, 1.94]			
Liu PH et al, 2016	-0.7514	0.3777	3.7%	0.47 [0.22, 0.99]			
Nishikawa H et al, 2011	-0.0121	0.4023	3.3%	0.99 [0.45, 2.17]			
Peng ZW et al, 2012	-0.0101	0.2907	5.4%	0.99 [0.56, 1.75]			
Song J et al, 2016	-0.0834	0.5388	2.0%	0.92 [0.32, 2.64]			
Takayasu K et al, 2018	-0.1054	0.1282	12.7%	0.90 [0.70, 1.16]		-	
Wu CC et al, 2021	-0.5276	0.2521	6.6%	0.59 [0.36, 0.97]			
Yun WK et al, 2011	-0.8916	0.1594	10.8%	0.41 [0.30, 0.56]		.	
Zhang C et al, 2022	-0.512	0.1741	10.0%	0.60 [0.43, 0.84]		-	
Total (95% CI)			100.0%	0.65 [0.55, 0.76]		•	
Heterogeneity: Tau ² = 0.0	3; Chi² = 24.76, df = '			400			
Test for overall effect: Z =	5.38 (P < 0.00001)	20			0.01	0.1 1 10 Favours [LR] Favours [RFA]	100

Figure5 Funnel plots for comparison of all propensity score-matched cohorts at patients with solitary small HCC measuring ≤ 3 cm. (A) OS. (B) DFS. (C) RFS.



Supplementary figures 1-10 are included in the supplementary materials

SDC link: http://links.lww.com/ISJP/A6

Highlights

• In a meta-analysis of 36 observational studies, liver resection group showed a significant improvement in survival outcomes of patients with HCC measuring \leq 3 cm, compared to RFA. Results were consistent in the propensity score matched analyses.

• Merging data from 14 observational studies, liver resection exhibited a significant survival benefit on OS, DFS and RFS for patients with HCC measuring ≤ 2 cm and the results were consistent after propensity score matching.