Impact of marital status on overall survival in patients with early-stage hepatocellular carcinoma

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The purpose of the present research was to assess the prognostic impact of marital status in hepatocellular carcinoma (HCC) patients with tumors ≤ 2 cm (stage Ia) based on the data from the Surveillance, Epidemiology, and End Results (SEER) database. Patients who received a histopathologic HCC diagnosis between 2004 and 2016 were recruited. Overall survival (OS) was the major outcome measure. The Cox regression model and the Fine-Gray regression model were used for the purpose of comparing and examining the prognostic value of marital status for OS. The data for a total of 2446 stage Ia HCC patients were extracted from the database. The median overall survival time was 96.0 months, with 5-year and 10-year overall survival rates of 58.2% and 45.8%, respectively. In both the Fine-Gray regression model and Cox regression model, marital status [married vs. unmarried and others, both \( P < 0.001 \), hazard ratio (HR) = 1.389 for Cox and HR = 1.378 for Fine-Gray], age at diagnosis, tumor grade, and surgery at the primary site independently served as prognostic indicators associated with OS. In conclusion, positive marital status was independently associated with better OS for stage Ia HCC patients, and its prognostic influence should be validated in the near future.

Primary liver cancer has been ranked as the seventh most prevalent malignant neoplasm and the second major contributor to cancer-related deaths on a global scale1–3. Among all types of liver cancers, hepatocellular carcinoma (HCC) is the dominant type and accounts for more than 75% of primary liver cancers4,5. Benefiting from early detection and timely treatment of some major risk factors, including alcoholic and/or nonalcoholic fatty liver disease and chronic HCV and/or HBV infection, the incidence rate of HCC has recently slowed in some areas6–8. Additionally, encouraging clinical results over the past three decades indicate that the overall survival (OS) rate of HCC has increased slightly9,10. Further, curative surgical resection, including liver transplantation or local ablation, for treating early-stage HCC has made great contributions11.

In addition, amid all of the indicators to defeat this disease, marital status has long been explored as a significant prognostic variable in a wide range of tumors. In 2013, Aizer et al. examined the influences of marital status on clinical outcomes among 10 major contributors to cancer-associated fatality in the United States using the Surveillance, Epidemiology and End Results (SEER) program12. For liver or intrahepatic bile duct cancers specifically, their results demonstrated that married patients exhibited a greater possibility of receiving definitive therapy \( (P < 0.01) \) and have significant survival benefits \( (P < 0.01) \) than unmarried persons (including divorced/separated, widowed, and single). However, in another large sample retrospective analysis conducted in Italy, a decreased risk of liver cancer was observed in unmarried patients13. Similarly, for HCC patients with poor or anaplastic differentiation who underwent surgical resection, a 2019 SEER report suggested that marital status had a non-significant benefit on survival outcomes14.

Considering that HCC patients with tumors ≤ 2 cm (stage Ia) could enjoy long-term survival and the unknown correlation between stage Ia HCC and marital status, we extracted and analyzed data from the SEER database to further evaluate the influences of marital status on survival status in this setting.

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Methods

Patient selection. This retrospective study obtained data from the SEER 18 population-based registries (SEER® Stat 8.3.6), which includes approximately 30% of the United States population. Data of 88,559 patients with confirmed liver cancer diagnoses from 2004 to 2016 was obtained from the database (ICD-O-3 Histology recodes 8140-8389). The following were the inclusion criteria used in recruiting participants for the present study: (1) patients who have a histopathological diagnosis of HCC; (2) tumors less than 2 cm without lymph node metastasis; and (3) upon HCC diagnosis, the patient must be no younger than 18 years of age. Below were some of the major exclusion criteria: (1) patients’ records of other cancers or with metastatic diseases and for whom HCC was not the first diagnosed cancer, and (2) HCC patients whose survival time was less than 1 month or who had untraced data (Fig. 1).

Determination of study variables and outcomes. We queried the database for patient information including demographics, treatment history and outcomes (age, sex, race, marital status, local surgery, use of radiotherapy (RT), use of chemotherapy (CT), survival data including status (alive or dead), cause of death (COD) and COD to site recode, survival time in months, and cause-specific death classification). For HCC patients registered in 2016, eligible tumors were restaged according to the definitions in the AJCC 8th edition. We defined marital status as a binary factor by dividing married (including common law) by unmarried and others, which included divorced, separated, single (never married), widowed, unmarried or domestic partners, and unknown coded in the SEER data, as has been reported in other studies. The items COD to site recode and cause-specific death classification indicated if a patient died from HCC (HCC-DSD: disease-specific death) or of causes other than HCC (HCC-NDSD: nondisease-specific death).

Data analysis. As the main endpoint in the present research, we measured OS, which was defined as the duration between the day of HCC diagnosis and the day of death or the final follow-up recorded in the database. CSS (cause-specific survival) was described as the duration between the day of initial diagnosis and the point of death attributable to HCC or the date of the final follow-up recorded in the program, whichever came first. Patients’ baseline characteristics were summarized by descriptive statistics and frequency tables. One-way analysis of variance (ANOVA) was used to compare the proportions of different groups. The non-linear correlation between age at diagnosis and all CODs was investigated by means of a restricted cubic spline (RCS). Methods for multivariate and univariate analysis by Cox regression model were described in our previous studies. In addition, considering that stage Ia HCC patients could enjoy long-term survival, we further regarded HCC-NDSD as a competing event in this cohort. In this model, the cumulative incidence function (CIF) was employed to determine the possibility of each factor in the univariate analysis and was checked with Gray’s test. According to the results of the univariate analysis, variables with a p-value < 0.05 were selected and incorporated into a multivariate competing-risks survival analysis with the aid of a proportional subdistribution hazard model, as determined by the Fine-Gray test. Hazard ratios (HRs) and their 95% confidence intervals (CIs) were also calculated in the analysis. A two-sided P value < 0.05 was established as a criterion of statically significant difference. Furthermore, to better illustrate the effects of marital status on liver cancer, the following librarians/databases were searched to trace eligible studies: Ovid MEDLINE®, PubMed, and Google Scholar updated until 31 December 2021 (present in the discussion section). The search strategy was developed and consisted of 2 main concepts: (1) liver cancer and (2) marital status. Methods for article selection were also described in our previ-
ous studies22,23 and we confirmed that the selection was performed in accordance with the PRISMA guidelines for systematic reviews.

The R software (version: 3.6.2; Institute for Statistics and Mathematics, Vienna, Austria; https://www.r-project.org) and SPSS software (version: 25.0 IBM Corporation, Armonk, NY, USA) was utilized to perform all analyses of statistical data.

Ethics statement. It was not necessary to get written informed consent for participating in the present research as the information contained in the SEER database has been de-identified and is publically available following authorization. The present research was exempted from ethical assessment by the Institutional Review Board of Zhejiang Provincial People’s Hospital. We hereby certify that the present research was conducted in conformity with the Declaration of Helsinki.

Results

Demographic and baseline characteristics. We extracted the data from 2446 eligible stage Ia HCC patients from the SEER database between 2004 and 2016. Table 1 presents the demographics and baseline features of the patients who were included in the present research. The ages of patients at diagnosis ranged between 20 and 90 years old with 59 years old as the median age. We further applied RCS with 3 knots (5th, 50th, and 95th centiles) to evaluate the association between age at diagnosis and all CODs (Fig. 2A). Based on the result, the appropriate inflection point to age at diagnosis was also 59 years old. Among all enrolled patients, 1397 (57.1%) patients were married, and 1049 (42.9%) HCC patients were classified as unmarried or other. A total of 1860 (76.0%) of the patients underwent cancer surgery. Only 64 (2.6%) HCC patients were treated with RT. The correlation coefficient of different variables is presented in Fig. S1. No factors were highly correlated with other baseline characteristics. Additionally, male patients were observed to have significant higher proportion with positive marital status than unmarried and others (45.0% vs. 29.4%, \( P < 0.001 \)), while age had no significance with marital status.

Survival outcomes. The OS rates for 5- and 10-year were 58.2% (95% CI, 0.560–0.604) and 45.8% (95% CI, 0.431–0.485), respectively, with a median OS time of 96.0 months (95% CI, 82.9–109.1, Fig. 2B). Of the 2,446 HCC patients, 326 patients died of HCC-NDSD, such as other infectious and parasitic diseases including HIV (n = 122), other causes of death (n = 39), and diseases of the heart (n = 38), accounting for 61.0% of the total (Fig. S2). Table 1 presents the baseline features of HCC patients who died due to HCC-DSD and HCC-NDSD.

Table 1. Baseline characteristics of stage Ia HCC patients. HCC: Hepatocellular carcinoma; DSD: disease-specific death; NDSD: non disease-specific death.

| Characteristic                        | ALL (%) | HCC-DSD (%) | HCC-NDSD (%) |
|---------------------------------------|---------|-------------|--------------|
| Age (years)                           |         |             |              |
| < 59                                  | 1182 (48.3) | 271 (41.5) | 156 (47.9)   |
| ≥ 59                                  | 1264 (51.7) | 382 (58.5) | 170 (52.1)   |
| Race                                  |         |             |              |
| White                                 | 1763 (72.1) | 476 (72.9) | 240 (73.6)   |
| Non-white                             | 683 (27.9) | 177 (27.1) | 86 (26.4)    |
| Sex                                   |         |             |              |
| Female                                | 626 (25.6) | 186 (28.5) | 72 (22.1)    |
| Male                                  | 1820 (74.4) | 467 (71.5) | 254 (77.9)   |
| Marital status                        |         |             |              |
| Married                               | 1397 (57.1) | 327 (50.1) | 165 (50.6)   |
| Unmarried and others                  | 1049 (42.9) | 326 (49.9) | 161 (49.4)   |
| Differentiation                       |         |             |              |
| Well and moderately differentiated    | 1556 (63.6) | 336 (51.5) | 213 (65.3)   |
| Poorly and undifferentiated           | 164 (6.7) | 61 (9.3) | 20 (6.1)     |
| Unknown                               | 726 (29.7) | 256 (39.2) | 93 (28.6)    |
| Surgery at the primary site           |         |             |              |
| No/unknown                            | 586 (24.0) | 282 (43.2) | 84 (25.8)    |
| Cancer-directed surgery performed     | 1860 (76.0) | 371 (56.8) | 242 (74.2)   |
| Radiotherapy (RT)                     |         |             |              |
| No/unknown                            | 2382 (97.4) | 638 (97.7) | 319 (97.9)   |
| Yes                                   | 64 (2.6) | 15 (2.3) | 7 (2.1)      |
| Chemotherapy (CT)                     |         |             |              |
| No/unknown                            | 1792 (73.3) | 444 (68.0) | 259 (79.4)   |
| Yes                                   | 654 (26.7) | 209 (32.0) | 67 (20.6)    |
spondingly, with a median CSS time not achieved at the time of analysis (Fig. 2B). Among married patients, the median OS duration was 130.0 ± 7.7 months (95% CI, 114.961–145.039), and in patients classified as unmarried and other, the median OS duration was 65.0 ± 5.6 months (95% CI, 53.951–76.049).

**Fine-gray regression analysis.** A univariate analysis using Fine-Gray test suggested that age at diagnosis \( (P < 0.001) \), sex \( (P = 0.004) \), surgery at the primary site \( (P < 0.001) \), tumor differentiation \( (P < 0.001) \), and CT \( (P < 0.001) \) significantly impacted the prognosis of stage Ia HCC patients. The cumulative risk curves for marital status are shown in Fig. 2C. The CIF was found to be elevated over 36, 60, and 120 months and was elevated for advanced age, female sex, unmarried and others, poorly or undifferentiated or unknown tumor grade, no or unknown status of surgery at the primary site, and receiving CT among eight variables. The CIF values among married HCC patients were 18.8%, 25.0%, and 30.6% at 36, 60, and 120 months, respectively. The corresponding figures for patients recoded as unmarried and others were 25.4%, 33.5%, and 42.4%, correspondingly. Table 2 depicts the findings recorded from the CIF values and univariate analysis.

We then employed the six variables that had statistical significance in the univariate analysis entered into the Fine-Gray model. According to the findings from the Fine-Gray regression model, age at the time of diagnosis \(< 59 \) vs. \( \geq 59 \), \( P < 0.001 \), HR = 1.419, 95% CI: 1.246–1.618), marital status (married vs. unmarried and others, \( P < 0.001 \), HR = 1.378, 95% CI: 1.212–1.568), tumor differentiation (well or moderately differentiated vs. poorly or undifferentiated, \( P = 0.007 \), HR = 1.402, 95% CI: 1.095–1.795; well or moderately differentiated vs. unknown, \( P = 0.032 \), HR = 1.169, 95% CI: 1.014–1.349), surgical resection of the primary site (no/unknown vs. yes, \( P < 0.001 \), HR = 0.340, 95% CI: 0.293–0.393) and treatment of CT (no/unknown vs. yes, \( P = 0.009 \), HR = 0.819, 95% CI: 0.706–0.952) all served as prognostic indicators that were significantly associated with OS in an independent manner (Table 3). Furthermore, multivariate analysis of NDSD also indicated that marital status (married vs. unmarried and others, \( P < 0.001 \), HR = 1.481, 95% CI: 1.308–1.680) was an independent factor associated with NDSD (Table S1).

**Cox regression analysis.** According to the results of the Cox regression analysis, four clinicopathological characteristics, namely age at the time of diagnosis, sex, tumor differentiation, marital status, and one treatment-related parameter (surgery at the primary site) were significantly associated with OS (Table S2). The findings recorded from the multivariate analysis illustrated that the significant covariates were age at the time of diagnosis \(< 59 \) vs. \( \geq 59 \), \( P < 0.001 \), HR = 1.415, 95% CI: 1.244–1.611), marital status (married vs. unmarried and others, \( P < 0.001 \), HR = 1.389, 95% CI: 1.106–1.767; well or moderately differentiated vs. poorly or undifferentiated, \( P = 0.005 \), HR = 1.398, 95% CI: 1.106–1.767; well or moderately differentiated vs. unknown, \( P = 0.055 \), HR = 1.146, 95% CI: 0.997–1.318) and surgical resection of the primary site (no/unknown vs. yes, \( P < 0.001 \), HR = 0.357, 95% CI: 0.311–0.409; Table 3). Both the Fine-Gray regression model and the Cox regression model demonstrated that marital status independently served as a prognostic indicator for OS.

**Discussion**

The objective of the present research was to evaluate the effect of marital status on OS in stage Ia HCC patients since the influence of marital status remains controversial in this setting. While using available data in the SEER database, we demonstrated that positive marital status acted as a prognostic variable in an independent manner favoring improved OS in both the Fine-Gray regression model and Cox regression model.

The results exploring the effects of marital status on liver cancer patients are summarized in Table 4 following chronological order. \(^{14,24–36}\) Fourteen studies that met the eligibility requirements were enrolled. It should be noted that marital status was not covered in the National Cancer Data Base, which is also a widely used database for the analysis of various malignancies. \(^{37}\) Most studies (13/14, 92.9%) were retrospective analyses with data extracted from the SEER database, with only one report having external validation with patients from their own cancer center. Among the 13 reports, 3 studies showed no significant association between marital status...
and survival outcomes, one study investigated the benefit of RT in HCC patients with major vascular invasion ($P = 0.834$ in univariate analysis), one study assessed the impact of RT in unresectable HCC patients ($P = 0.475$), and the other study evaluated the effects of marital status on patients developing less differentiated HCC who underwent surgical resection ($P = 0.370$). The remaining 10 studies all supported the benefit of positive marital status for liver cancer patients despite different situations of patient enrollment. A similar study was reported by Peters and colleagues$^{29}$. In this study, 13,694 HCC patients diagnosed with stage I-II disease were enrolled in the present research. Positive marital status was first demonstrated to have a significantly higher likelihood of patients receiving liver resection, and liver transplantation (both $P < 0.001$). The findings recorded from the Cox regression analysis correlated with DSS, being married was shown to have a significantly longer DSS ($P = 0.010; HR = 0.71; 95\% CI, 0.55–0.92$). In the only prospective study$^{36}$, Chiu et al. compared four models predicting quality of life

| Factors                        | Gray's test | $P$-value | Cumulative incidence function (CIF) |
|-------------------------------|-------------|-----------|------------------------------------|
|                               |             |           | 36-months | 60-months | 120-months |
| Age (years)                   | 35.156      | $< 0.001$ |           |           |            |
| $< 59$                        |             |           | 0.178     | 0.239     | 0.286      |
| $\geq 59$                      |             |           | 0.253     | 0.334     | 0.442      |
| Race                          | 0.618       | 0.432     |           |           |            |
| White                         |             |           | 0.231     | 0.290     | 0.347      |
| Non-white                     |             |           | 0.179     | 0.276     | 0.376      |
| Sex                           | 8.128       | 0.004     |           |           |            |
| Female                        |             |           | 0.252     | 0.328     | 0.412      |
| Male                          |             |           | 0.205     | 0.273     | 0.337      |
| Marital status                | 20.163      | $< 0.001$ |           |           |            |
| Married                       |             |           | 0.188     | 0.250     | 0.306      |
| Unmarried and others          |             |           | 0.254     | 0.335     | 0.424      |
| Differentiation               | 57.738      | $< 0.001$ |           |           |            |
| Well and moderately differentiated |       |           | 0.167     | 0.227     | 0.295      |
| Poorly and undifferentiated   |             |           | 0.322     | 0.395     | 0.427      |
| Unknown                       |             |           | 0.297     | 0.386     | 0.466      |
| Surgery at the primary site   | 250.974     | $< 0.001$ |           |           |            |
| No/unknown                    |             |           | 0.456     | 0.538     | 0.614      |
| Cancer-directed surgery performed |       |           | 0.142     | 0.209     | 0.278      |
| Radiotherapy (RT)             | 0.398       | 0.528     |           |           |            |
| No/unknown                    |             |           | 0.214     | 0.285     | 0.355      |
| Yes                           |             |           | 0.323     | 0.323     | 0.323      |
| Chemotherapy (CT)             | 13.425      | $< 0.001$ |           |           |            |
| No/unknown                    |             |           | 0.195     | 0.260     | 0.332      |
| Yes                           |             |           | 0.273     | 0.358     | 0.413      |

Table 2. Univariate analysis in stage Ia HCC patients by using a competing risk model.

| Factor                        | Cox regression analysis | Fine-Gray regression analysis |
|-------------------------------|-------------------------|------------------------------|
|                               | $P$ value | HR Lower | 95% CI | HR Upper | 95% CI | $P$ value | HR Lower | 95% CI | HR Upper | 95% CI |
| Age, $< 59$ versus $\geq 59$  | $< 0.001$ | 1.415    | 1.244 | 1.611 | $< 0.001$ | 1.419 | 1.246 | 1.618 |
| Race, white versus non-white  | -         | -        | -     | -     | -         | -     | -     | -     |
| Sex, female versus male       | 0.581     | 0.960    | 0.830 | 1.110 | 0.564     | 0.958 | 0.829 | 1.108 |
| Marital status, married versus unmarried and others | $< 0.001$ | 1.389    | 1.223 | 1.578 | $< 0.001$ | 1.378 | 1.212 | 1.568 |
| Grade, reference: well and moderately | -      | -        | -     | -     | -         | -     | -     | -     |
| Poorly and undifferentiated   | 0.005     | 1.398    | 1.106 | 1.767 | 0.007     | 1.402 | 1.095 | 1.795 |
| Unknown                       | 0.055     | 1.146    | 0.997 | 1.318 | 0.032     | 1.169 | 1.014 | 1.349 |
| Surgery at the prim site, no/unknown versus yes | $< 0.001$ | 0.357    | 0.311 | 0.409 | $< 0.001$ | 0.340 | 0.293 | 0.393 |
| RT, no/unknown versus yes     | -         | -        | -     | -     | -         | -     | -     | -     |
| CT, no/unknown versus yes     | -         | -        | -     | -     | -         | -     | -     | -     |

Table 3. Multivariate analysis of OS in stage Ia HCC patients with the Cox regression model and the Fine-Gray regression model. HR, Hazard ratio; CI, confidence interval.
Unfortunately, all of these enrolled studies concluded that marital status on survival outcomes was based on the findings revealed by the Cox proportional hazard models. In fact, for HCC patients diagnosed with stage Ia, 33.3% (326/979) of patients died due to various reasons other than HCC, as revealed in the current study (Fig. S2). This nonhomogeneity could certainly cause bias in calculating the effects of marital status on OS. Starting with this consideration and the knowledge that stage Ia HCC patients could enjoy prolonged survival duration, with a 10-year overall survival rate of 45.8%, we then employed the Fine-Gray regression model according to the findings recorded from the CIF analysis. Marital status still acted as a prognostic parameter affecting OS in an independent manner ($P < 0.001$, HR = 1.378). Previously, Yang et al. compared the Fine-Gray regression model with the Cox regression model in penile cancers38. Survival analysis indicated that the findings derived with the aid of the Cox regression model were different from those obtained by the Fine-Gray regression model, while the Kaplan–Meier curve analysis led to an overestimation compared to the CIF analysis of penile cancer patients. Similar findings were also observed in esophageal cancer39 and cecum cancer40 in the literature. Therefore, when comparing survival outcomes among different variables, especially for competing events in survival outcomes, it is worth considering the possibility of CIF analysis to avoid overestimation results41.

This study is first limited by its inherent retrospective nature, with some heterogeneity in the analysis. Some important information, such as patients’ baseline characteristics, including lesbian, gay, bisexual, transgender, and queer status, liver function, alcoholic and/or nonalcoholic fatty liver disease, HBV ± HCV infection, cancer location, Child–Pugh score, and treatment complications, was not available in the SEER database. Secondly, the

| Reference (First author) | No. of patients and study design | Study period (year) | Cancer type (HCC/ICC/Both) | Source of data (IE/Database) | Main study population and age cutoff value (CV) | Main conclusion on marital status |
|--------------------------|----------------------------------|---------------------|---------------------------|-----------------------------|-----------------------------------------------|---------------------------------|
| Chen et al.24            | 1,352, retrospective             | 1973–2013           | ICC                       | SEER                         | Patients stratified by localized/Regional/Distant stage, CV = 70 | Positive for CSS*               |
| Zhang et al.25           | 8,621, retrospective             | 1988–2007           | HCC                       | SEER                         | Patients with all stages, receiving hepatic resection, CV = 45, 60, 75 | Positive for CSS               |
| Wu et al.26              | 13,408, retrospective            | 1998–2013           | HCC                       | SEER                         | Patients with all stages, receiving hepatic resection, CV = 60 | Positive for CSS               |
| He et al.27              | 40,809, retrospective            | 2004–2012           | Both                      | SEER                         | Patients stratified by localized/Regional/Distant stage, CV = 60 | Positive for OS and CSS         |
| Wu et al.28              | 12,168, retrospective            | 2004–2012           | HCC                       | SEER                         | Patients with all stages stratified by race and gender, CV = 60, 80 | Positive for OS and CSS         |
| Peters et al.29          | 13,694, retrospective            | 2004–2012           | HCC                       | SEER                         | Stage I-II HCC, CV = 60 | Positive for DSS               |
| Lin et al.30             | 3,181, retrospective             | 2004–2013           | HCC                       | SEER                         | Patients with major vascular invasion receiving radiotherapy, CV = 65 | Negative for OS               |
| Zhang et al.31           | 1,305, retrospective             | 2004–2014           | HCC                       | SEER                         | Unresected HCC patients with all stages receiving radiotherapy, CV = 50,60,70,80 | Negative for OS               |
| Xiao et al.32            | 15,638, retrospective            | 2004–2014           | HCC                       | SEER and IE                  | Patients with all stages excluded unknown baseline characteristics, CV = 60, 70, 80 | Positive for OS and CSS        |
| Yan et al.34             | 1,581, retrospective             | 2004–2015           | HCC                       | SEER                         | All stages with less differentiated HCC receiving hepatic resection, CV = 60 | Negative for CSS               |
| Wu et al.35              | 8,677, retrospective             | 2010–2012           | Both                      | SEER                         | Compared according to metastatic status, CV = 60, 75 | Positive for OS and CSS         |
| Guo et al.36             | 1,567, retrospective             | 2010–2014           | HCC                       | SEER                         | Patients with bone metastasis, CV, not available | Positive for OS               |
| Liang et al.37           | 4,933, retrospective             | 2010–2015           | Both                      | SEER                         | Patients with all stages without hepatic resection, CV = 60, 75 | Positive for CSS               |
| Chiu et al.38            | 332, prospective                | 2012–2015           | HCC                       | IE (three centers)           | Stage I-III HCC receiving hepatic resection, median age = 60 | Positive for QoL               |

Table 4. Summary of published studies on the impact of marital status on liver cancer. HCC: Hepatocellular carcinoma; ICC: intrahepatic cholangiocarcinoma; IE: institutional experience; *: statistically significant for regional disease; QoL: quality of life.
SEER database only recorded marital status at patients’ initial diagnosis, and changes in marital status might occur during the long-term follow-up periods, which could alter the influence of marital status. Third, a high incidence of receiving CT was observed within the cohort, which was mainly explained by these patients were re-staged based on the newly AJCC 8th staging manual. Previous guidelines for treating early stage HCC with/without microvascular invasion, adjuvant CT followed by surgical resection was recommended. Finally, potential interactions among sex, generation and socioeconomic status might influence marital status, and whether the findings generated in the current study apply to other populations or demographics around the world, need to be confirmed by large, well-designed, prospective studies in the future.

In conclusion, we evaluated the impact of marital status on the OS outcomes of HCC patients with tumors ≤ 2 cm registered in the SEER database between 2004 and 2016. Through comparison between two different regression models, positive marital status can be used to act as a prognostic indicator for better OS outcomes in an independent manner. Our results are consistent with earlier major studies supporting the benefit of being married for stage Ia HCC patients. When treating localized diseases with potential medical cures, additional interventions, including but not limited to family and social support, should be given to subpopulations with negative marital status.

Data availability
The datasets produced for this work (SEER database) are accessible via the following link: https://seer.cancer.gov/data/access.html. Further inquiries can be directed to the corresponding authors.

Received: 25 March 2022; Accepted: 1 June 2022
Published online: 19 November 2022

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Acknowledgements
We received no financial support for the present research. We thank the National Cancer Institute (NCI) for providing the SEER data after permission.

Author contributions
S.Y. and F.C. conceived the study. Y.W. and H.X. searched the database and literature. F.C., T.S. and S.Y. discussed and analyzed the data. F.C. and T.S. wrote the manuscript. Y.W., H.X. and S.Y. revised the manuscript. All authors approved the final version.

Competing interests
The authors declare no competing interests.

Additional information
Supplementary Information The online version contains supplementary material available at https://doi.org/10.1038/s41598-022-14120-1.

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