Conclusions False positive diagnosis of advanced fibrosis in NAFLD patients can be reduced and unnecessary liver biopsy can potentially be avoided by repeat LSM.

**HEPATIC RESECTION VERSUS TRANSCATHETER ARTERIAL CHEMOEMBOLIZATION IN RESECTABLE INFILTRATIVE HEPATOCELLULAR CARCINOMA: A PROPENSITY SCORE WEIGHTED LANDMARK STUDY**

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Background Patients with infiltrative hepatocellular carcinoma (iHCC) have a worse prognosis than those with other types of HCC and limited choices of treatments. The efficacy of hepatic resection in iHCC patients was unclear, especially comparing with other treatments. We aim to compare the efficacy of hepatic resection (HR) and transcatheter arterial chemembolization (TACE) for patients with resectable iHCC.

Methods We retrospectively enrolled patients with resectable iHCC who were treated by HR or TACE from four clinical centers. Their overall survival (OS) time was calculated and compared by Log-Rank test. A propensity score-matched (PSM) analysis was performed to reduce selection bias.

Results From January 2010 to December 2017, 178 patients with resectable iHCC were collected (124 patients received HR and 54 patients received TACE) and entered into 6, 9, 12-month landmark analysis. The median overall survival (OS) time was significantly longer in patients treated by HR than TACE (19 vs 11 months, p=0.0041). Landmark analysis limiting survivors after 6, 9, 12 months also showed the benefit of HR over TACE in multi-variables COX regression (all p<0.01). Patients with tumors located in both liver lobes or portal hypertension tended to receive HR rather than TACE. After propensity score matched, 46 pairs were compared and HR obtained better overall survival than TACE (median OS time: 29 vs 11 months, p=0.001). The multi-variables analysis indicated vascular invasion as an independent predictor for worse OS in 6-month landmark subgroup (HR: 2.231, 95%CI: 1.364-3.649, p=0.001), especially for patients with the major trunk of portal vein involved.

Conclusions HR might be an optimal choice for patients with resectable iHCC due to its better survival benefit than TACE. Patients with trombosis in the major trunk of portal vein need multiple department team discussion to decide which therapy to be performed.

**PREDICTORS OF SURVIVAL FOR HEPATOCELLULAR CARCINOMA TREATED WITH ULTRASOUND-GUIDED PERCUTANEOUS RADIOFREQUENCY ABLATION**

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Background Radiofrequency ablation (RFA) is a widely used technique for treating hepatocellular carcinoma (HCC). Based on current data, tumor size and liver function are predictors of survival for HCC. Predictors of Survival in the Philippines may be different from other countries. We aimed to determine the independent predictors of survival for HCC patients undergoing RFA.

Methods RFA was performed on 181 consecutive HCC patients from June 2007 to November 2018. Survival was estimated using Kaplan-Meier and multivariate analysis of survival predictors were analyzed using Cox regression.

Results After a median follow-up of 19.9 months, the 1-, 3- and 5-year overall survival rates were 81.4%, 53.7% and 30.4%, respectively. Local tumor progression and intrahepatic distant recurrence were observed in 42.5% (77/181) and 30.9% (56/181), respectively. Patients with serum albumin ≥3.5 g/L had better survival compared to serum albumin<3.5 g/L (1,3&5 years: =69.5%,47.2%,38.9% vs 39.1%,0%,0%; p<0.001 – figure 1). Other factors associated with survival on univariate analysis included: Cirrhosis(p=0.0001), Child-pugh class(p=0.026), RFA time (p=0.011), Platelet count<150×10^9/L (p=0.002), INR (p<0.001) and total bilirubin(p=0.026). The sole independent predictor of survival on multivariate analysis was serum albumin(OR=3.40;95%CI= 0.196-0.442;p<0.001).

Conclusions In patients with HCC treated with RFA, serum albumin ≥3.5 g/L appear to result in better survival.

**PREDICTORS OF TECHNIQUE EFFECTIVENESS FOR ULTRASOUND-GUIDED PERCUTANEOUS RADIOFREQUENCY ABLATION FOR HEPATOCELLULAR CARCINOMA**

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Background Radiofrequency ablation (RFA) is a widely used technique for treating hepatocellular carcinoma (HCC),