

Background/Aims:

- The GALAD score, which incorporates the biomarkers AFP, AFP-L3% and DCP, has excellent performance for detection of hepatocellular carcinoma (HCC).
- However, its accuracy in evaluating treatment response and predicting recurrence prediction is unknown.
- We aim to evaluate the accuracy of these biomarkers in predicting the presence of posttreatment viable tumor and HCC recurrence.

Methods:

Population: Single-institution, retrospect cohort study. Deep6 software was used to identify patients who had HCC biomarkers obtained from May 2019current.

Inclusion criteria:

- 1) History cirrhosis or chronic hepatitis B infection, HCC diagnosis, and curative surgical or locoregional treatment (Y-90, TACE, ablation).
- 2) HCC biomarkers obtained greater than 1 month post-treatment.

LAD score: Because the study cohort already has known HCC, we removed the demographic factors to calculate a score based solely on the biomarkers (the "LAD" score). The LAD score is calculated from the following formula:

$.04\times(AFP-L3) + 2.34\times\log(AFP) + 1.33\times\log(DCP)$

Evidence of viable tumor was determined by imaging, biopsy, or liver explant pathology after transplant.

Survival analysis for tumor recurrence was performed in patients with non-viable tumor on initial posttreatment imaging.

Figure 1: Patient identification and selection

Utility of LAD Score for Treatment Response Assessment and Recurrence Monitoring in Patients With Hepatocellular Carcinoma

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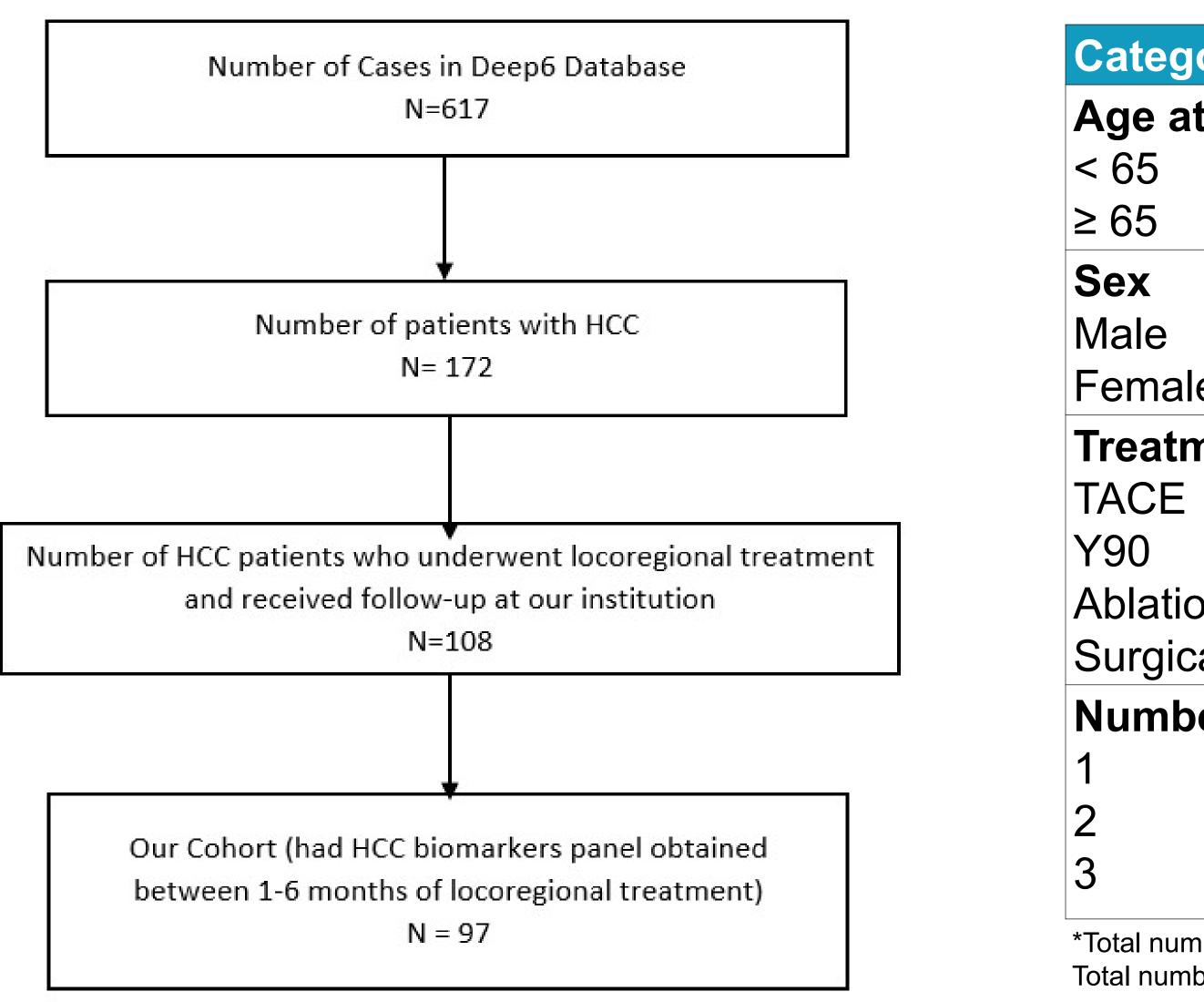


Figure 2: HCC recurrence in LAD-positive vs LAD-negative patients with non-viable tumor on initial imaging

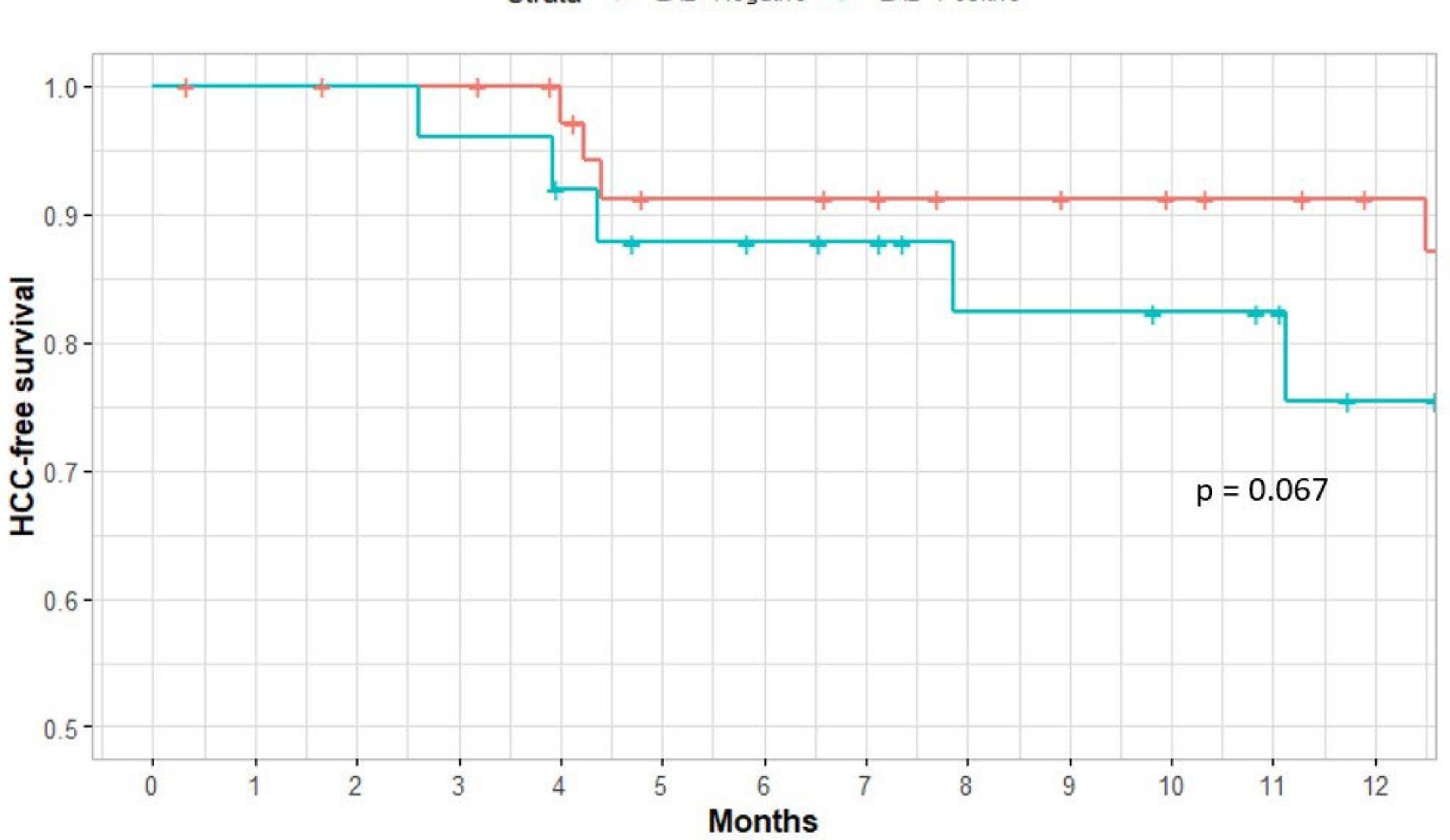


Table 1: Patient demographics

Jory	Incidence (%)
t time of treatment	
	34 (35.1%)
	63 (64.9%)
	52 (66.7%)
le	26 (33.3%)
ment modality	
	47 (48.6%)
	29 (29.9%)
on	16 (16.5%)
cal resection	7 (7.20%)
per of treatment incidents	
	63 (80.8%)
	11 (14.1%)
	4 (4.12%)

*Total number used for patient variables (sex, number of treatment incidents) is 78. Total number used for treatment variables (age at time of treatment, modality) is 97

Strata + LAD=Negative + LAD=Positive

Results:

- respectively.

Conclusions:

- viable tumors.

References:

- 9965.EPI-13-0870
- *Cancer*. 2020;2(1).

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• 78 patients (97 treatments) with surveillance imaging and biomarkers were analyzed. Mean age at time of HCC detection was 67 years, a third of patients were female, and the most common treatment modality was TACE (see **Table 1** for details).

• 34.0% of cases had a viable tumor on initial post treatment images.

• The optimal LAD cutoff was calculated at 2.23, yielding a sensitivity and specificity of 87.4% and 60.9% for detection of viable tumors,

• After median follow up of 21.3 months, 11 cases (17.2%) developed recurrent HCC (mean time to recurrence 3.5 months) among cases with post-treatment non-viable tumor.

• A total of 7 recurrence cases (64%) had a positive LAD score at negative post-treatment surveillance images. Positive LAD score had a borderline association with recurrence (HR 3.17; 95%CI [0.92, 10.92], p = 0.067 - see Fig 2).

• LAD score is increased in most HCC patients with post-treatment

It may provide benefit in risk stratification for HCC recurrence.

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3. Mendiratta-Lala M, Masch WR, Shampain K, et al. Mri assessment of hepatocellular carcinoma after localregional therapy: A comprehensive review. *Radiology: Imaging*