

Efficacy of Adjuvant Immunotherapy After Radiofrequency Ablation for Hepatocellular Carcinoma

Stephen Andrew Johnston, MD¹; Marybeth Hughes, MD¹; Eric Feliberti, MD¹; Rachel Burke, MD¹; Ann Harper, MPH²; Winifred Lo, MD¹

1)Department of Surgery, Macon and Joan Brock Virginia Health Sciences Eastern Virginia Medical School at Old Dominion University, Norfolk, VA

2) Sentara Health Research Center, Norfolk, VA

Introduction / Methods

Hepatocellular carcinoma (HCC) is the most common primary liver cancer. While surgical resection is the mainstay of treatment, radiofrequency ablation is an appropriate alternative for tumors < 3.5 cm in size. The role for adjuvant systemic immunotherapy after ablation is not well defined.

We performed retrospective analysis on data obtained from the National Cancer Database. We examined patients diagnosed with HCC from 2020-2022 who underwent ablation, immunotherapy, or combination therapy. We excluded patients who received surgical resection, patients with distal metastasis at time of diagnosis, and patients who did not receive treatment.

Patient Characteristics

	RFA	Immunotherapy	Both	Overall
Number of patients	3920	3379	85	7384
Male	2815 (71.8%)	2685 (79.5%)	67 (78.8%)	5567 (75.4%)
Female	1105 (28.2%)	694 (20.5%)	18 (21.2%)	1817 (24.6%)
Age at diagnosis	67.0	66.7	66.4	66.9
Initial AFP	125	951	470	482
Tumor size	27.9 mm	83.0 mm	50.0 mm	51.9 mm

Results

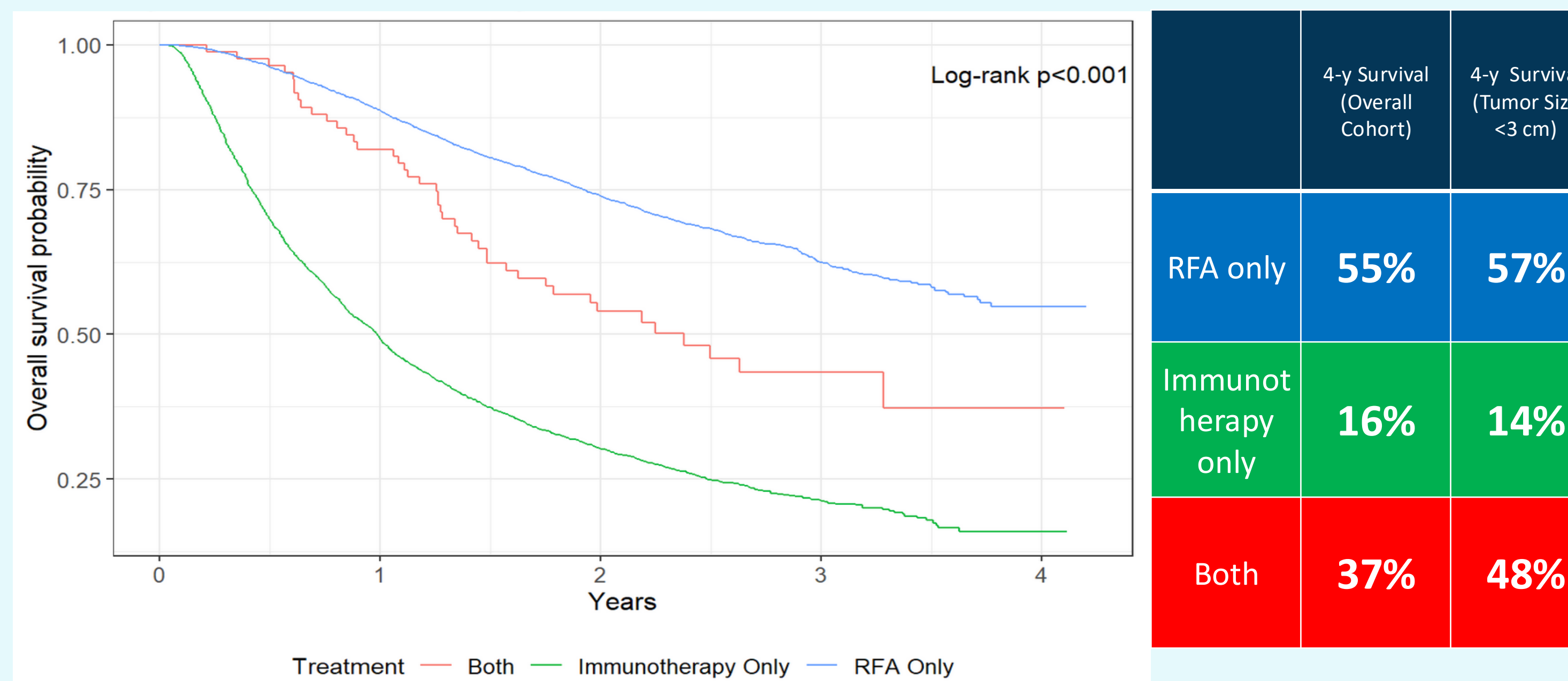


Figure 1: Kaplan-Meier Survival Curves by Treatment

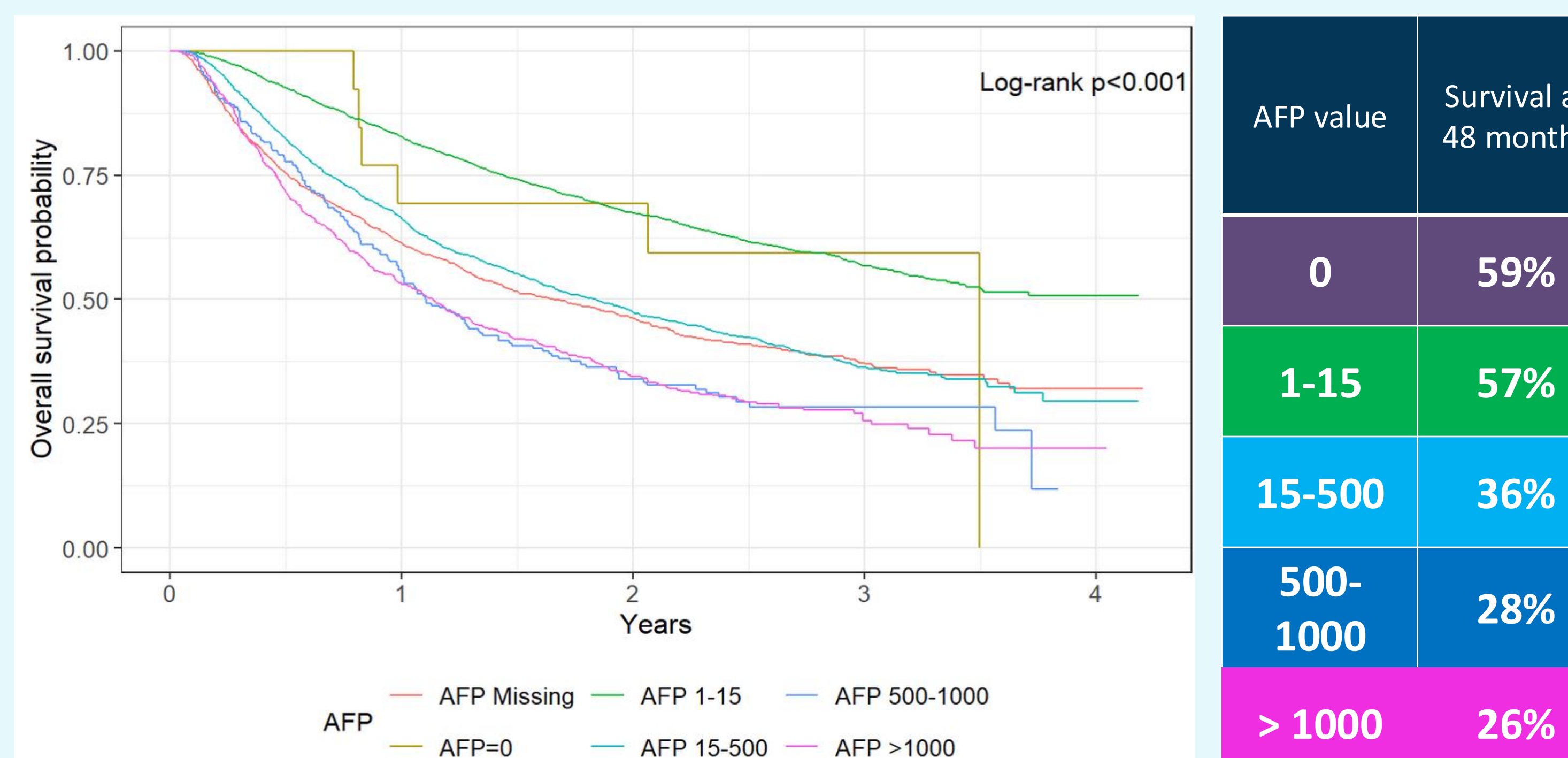


Figure 2: Kaplan-Meier Survival Curves by AFP

Discussion

RFA-only 4-year survival was 55%, whereas the lowest was in the immunotherapy only group (Figure 1). This includes all tumor sizes; subsequent subgroup analysis evaluating tumors <3 cm similarly showed that RFA-only patients experienced superior survival to those patients treated with immunotherapy.

Patients with higher AFP value were also assessed for differences in survival. AFP >500 is associated with decreased survival (Figure 2). Higher AFP value implies a larger tumor burden or higher risk-tumor, which would be associated with lower survival.

Our study does have several limitations. The National Cancer Database does not include data regarding the specific tumor biology of these patients nor degree of underlying liver dysfunction, which would contribute to survival. We also had a small sample size of patients who received combination therapy, which limits the usefulness of that comparison.

Resources

- Alawiyia B, Constantinou C. Hepatocellular Carcinoma: a Narrative Review on Current Knowledge and Future Prospects. *Curr Treat Options Oncol.* 2023;24(7):711-724. doi:10.1007/s11864-023-01098-9
- Abdelhamed W, El-Kassas M. Hepatocellular carcinoma recurrence: Predictors and management. *Liver Res.* 2023;7(4):321-332. Published 2023 Nov 19. doi:10.1016/j.livres.2023.11.004
- Singal AG, Llovet JM, Yarrow M, et al. AASLD Practice Guidance on prevention, diagnosis, and treatment of hepatocellular carcinoma. *Hepatology.* 2023;78(6):1922-1965. doi:10.1097/HEP.000000000000466
- Qin S, Chen M, Cheng AL, et al. Atezolizumab plus bevacizumab versus active surveillance in patients with resected or ablated high-risk hepatocellular carcinoma (IMbrave050): a randomised, open-label, multicentre, phase 3 trial. *Lancet.* 2023;402(10415):1835-1847. doi:10.1016/S0140-6736(23)01796-8
- Hoepfner J, Brunner T, Schmoor C, et al. Perioperative Chemotherapy or Preoperative Chemoradiotherapy in Esophageal Cancer. *N Engl J Med.* 2025;392(4):323-335. doi:10.1056/NEJMoa2409408
- Mieog JS, van der Hage JA, van de Velde CJ. Preoperative chemotherapy for women with operable breast cancer. *Cochrane Database Syst Rev.* 2007 Apr 18;2007(2):CD005002. doi: 10.1002/14651858.CD005002.pub2. PMID: 17443564; PMCID: PMC7388837.