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PEER-REVIEW REPORT

Name of journal: World Journal of Gastroenterology

Manuscript NO: 88517

Title: Transcatheter arterial chemoembolization combined with PD-1 inhibitors and

Lenvatinib for hepatocellular carcinoma with portal vein tumor thrombus

Provenance and peer review: Unsolicited Manuscript; Externally peer reviewed

Peer-review model: Single blind

Reviewer's code: 06195974 Position: Editorial Board Academic degree: MD, PhD

Professional title: Assistant Professor

Reviewer's Country/Territory: United States

Author's Country/Territory: China

Manuscript submission date: 2023-09-27

Reviewer chosen by: AI Technique

Reviewer accepted review: 2023-10-13 18:38

Reviewer performed review: 2023-10-13 19:12

Review time: 1 Hour

	[] Grade A: Excellent [Y] Grade B: Very good [] Grade C:
Scientific quality	Good
	[] Grade D: Fair [] Grade E: Do not publish
Novelty of this manuscript	[Y] Grade A: Excellent [] Grade B: Good [] Grade C: Fair [] Grade D: No novelty
Creativity or innovation of	[Y] Grade A: Excellent [] Grade B: Good [] Grade C: Fair
this manuscript	[] Grade D: No creativity or innovation



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Scientific significance of the conclusion in this manuscript	[Y] Grade A: Excellent [] Grade B: Good [] Grade C: Fair [] Grade D: No scientific significance
Language quality	[Y] Grade A: Priority publishing [] Grade B: Minor language polishing [] Grade C: A great deal of language polishing [] Grade D: Rejection
Conclusion	[] Accept (High priority) [] Accept (General priority) [Y] Minor revision [] Major revision [] Rejection
Re-review	[Y] Yes [] No
Peer-reviewer statements	Peer-Review: [Y] Anonymous [] Onymous Conflicts-of-Interest: [] Yes [Y] No

SPECIFIC COMMENTS TO AUTHORS

The authors assessed efficacy and safety of the combined treatment based on programmed cell death-1 (PD-1) inhibitor, transcatheter arterial chemoembolization (TACE), and Lenvatinib (PTL regimen) in a cohort of Hepatocellular carcinoma (HCC) subjects with portal vein tumor thrombus (PVTT). Overall, 41 eligible HCC patients with PVTT types I-IV were retrospectively enrolled. They were distributed to either the PTL (n=18) or TACE/Lenvatinib (TL) (n=23) regimen. Primary end-point was the median progression-free survival (mPFS), while median overall survival (mOS), objective response rate (ORR), disease control rate (DCR), and toxicity level served as secondary endpoints. After a median follow-up of 21.8 months, the DCRs were 88.9% and 60.9% in the PTL and TL groups (P=0.046), respectively. mPFS indicated significant improvement (HR=0.25; P