

## PEER-REVIEW REPORT

**Name of journal:** *World Journal of Gastroenterology*

**Manuscript NO:** 89452

**Title:** Hepatic arterial infusion chemotherapy with/without anti-angiogenesis agents and immune checkpoint inhibitions for unresectable hepatocellular carcinoma and meta-analysis

**Provenance and peer review:** Unsolicited Manuscript; Externally peer reviewed

**Peer-review model:** Single blind

**Reviewer's code:** 05747976

**Position:** Peer Reviewer

**Academic degree:** MD

**Professional title:** Assistant Professor, Doctor, Postdoc, Postdoctoral Fellow, Researcher, Senior Researcher

**Reviewer's Country/Territory:** United States

**Author's Country/Territory:** China

**Manuscript submission date:** 2023-11-01

**Reviewer chosen by:** AI Technique

**Reviewer accepted review:** 2023-11-20 23:25

**Reviewer performed review:** 2023-11-30 03:28

**Review time:** 9 Days and 4 Hours

Scientific quality	<input checked="" type="radio"/> Grade A: Excellent <input type="radio"/> Grade B: Very good <input type="radio"/> Grade C: Good <input type="radio"/> Grade D: Fair <input type="radio"/> Grade E: Do not publish
Novelty of this manuscript	<input type="radio"/> Grade A: Excellent <input checked="" type="radio"/> Grade B: Good <input type="radio"/> Grade C: Fair <input type="radio"/> Grade D: No novelty

<b>Creativity or innovation of this manuscript</b>	<input type="checkbox"/> Grade A: Excellent <input checked="" type="checkbox"/> Grade B: Good <input type="checkbox"/> Grade C: Fair <input type="checkbox"/> Grade D: No creativity or innovation
<b>Scientific significance of the conclusion in this manuscript</b>	<input checked="" type="checkbox"/> Grade A: Excellent <input type="checkbox"/> Grade B: Good <input type="checkbox"/> Grade C: Fair <input type="checkbox"/> Grade D: No scientific significance
<b>Language quality</b>	<input checked="" type="checkbox"/> Grade A: Priority publishing <input type="checkbox"/> Grade B: Minor language polishing <input type="checkbox"/> Grade C: A great deal of language polishing <input type="checkbox"/> Grade D: Rejection
<b>Conclusion</b>	<input type="checkbox"/> Accept (High priority) <input checked="" type="checkbox"/> Accept (General priority) <input type="checkbox"/> Minor revision <input type="checkbox"/> Major revision <input type="checkbox"/> Rejection
<b>Re-review</b>	<input checked="" type="checkbox"/> Yes <input type="checkbox"/> No
<b>Peer-reviewer statements</b>	Peer-Review: <input checked="" type="checkbox"/> Anonymous <input type="checkbox"/> Onymous
	Conflicts-of-Interest: <input type="checkbox"/> Yes <input checked="" type="checkbox"/> No

## SPECIFIC COMMENTS TO AUTHORS

This study had two major original components: A network meta-analysis suggesting HAIC-based regimens as first-line treatment for advanced HCC A real-world study finding significantly improved efficacy (PFS, OS) with acceptable safety for triple therapy (HAIC + angiogenesis inhibitors + PD(L)1 blockades) compared to AIPB The key hypothesis that was confirmed was that triple therapy would outperform AIPB for advanced HCC in a real-world Chinese population even though AIPB is guideline-recommended. This is a high quality study using appropriate methods and rigorous analyses. The findings are important given the lack of real-world evidence for triple therapy and help provide direction for first-line treatment selection. Unique insights: HAIC-based regimens, not just AIPB, should be considered for advanced HCC Efficacy benchmarks for triple therapy in real-world practice Factors like Child-Pugh score and number of HAIC cycles impact outcomes The conclusions accurately reflect

the data presented. This moves the field forward by validating an emerging combination therapy. Limitations are a single-center retrospective study with inherent biases. Additional prospective trials should evaluate optimal regimens, biomarkers, and safety. Future research can assess triple therapy in populations mirroring real-world demographics and disease characteristics. This could alter clinical practice by providing an evidence-based alternative to AIPB with improved outcomes.