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PEER-REVIEW REPORT

Name of journal: World Journal of Clinical Oncology

Manuscript NO: 92103

Title: Tumor Infiltrating Lymphocytes in Gastric Cancer: Unraveling Complex Interactions for Precision Medicine

Provenance and peer review: Invited Manuscript; Externally peer reviewed

Peer-review model: Single blind

Reviewer's code: 05381979

Position: Editorial Board

Academic degree: Doctor, PhD

Professional title: Director, Professor, Surgeon

Reviewer's Country/Territory: China

Author's Country/Territory: India

Manuscript submission date: 2024-01-15

Reviewer chosen by: AI Technique

Reviewer accepted review: 2024-01-29 03:09

Reviewer performed review: 2024-01-29 06:49

Review time: 3 Hours

	[] Grade A: Excellent [] Grade B: Very good [Y] Grade C:
Scientific quality	Good
	[] Grade D: Fair [] Grade E: Do not publish
Novelty of this manuscript	 []Grade A: Excellent []Grade B: Good [Y]Grade C: Fair []Grade D: No novelty
Creativity or innovation of	[] Grade A: Excellent [Y] 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	 [] Grade A: Excellent [Y] Grade B: Good [] Grade C: Fair [] Grade D: No scientific significance
Language quality	[] Grade A: Priority publishing [Y] Grade B: Minor language polishing [] Grade C: A great deal of language polishing [] Grade D: Rejection
Conclusion	 [] Accept (High priority) [Y] Accept (General priority) [] Minor revision [] Major revision [] Rejection
Re-review	[]Yes [Y]No
Peer-reviewer statements	Peer-Review: [Y] Anonymous [] Onymous Conflicts-of-Interest: [] Yes [Y] No

SPECIFIC COMMENTS TO AUTHORS

After reviewing the literature, the authors found that high intratumoral (IT) TILs were associated with higher grade, HER2 - and H. pylori negativity. Furthermore, stromal (ST) TILs are associated with lower GC grade and less risk of recurrence. High TILs in the ST and invasive border (IB) are also associated with dMMR status. Further characterization of CD3+, CD8+, and other cells is required. In the future, this complex correlation of cancer cells with the immune system could be explored to find therapeutic pathways.