

PEER-REVIEW REPORT

Name of journal: *World Journal of Gastroenterology*

Manuscript NO: 80398

Title: Emerging novel targets for nonalcoholic fatty liver disease treatment: evidence from recent basic studies

Provenance and peer review: Invited Manuscript; Externally peer reviewed

Peer-review model: Single blind

Reviewer's code: 03475142

Position: Peer Reviewer

Academic degree: MD, PhD

Professional title: Assistant Professor

Reviewer's Country/Territory: Japan

Author's Country/Territory: China

Manuscript submission date: 2022-10-04

Reviewer chosen by: AI Technique

Reviewer accepted review: 2022-10-07 08:37

Reviewer performed review: 2022-10-16 15:50

Review time: 9 Days and 7 Hours

Scientific quality	<input type="checkbox"/> Grade A: Excellent <input checked="" type="checkbox"/> Grade B: Very good <input type="checkbox"/> Grade C: Good <input type="checkbox"/> Grade D: Fair <input type="checkbox"/> Grade E: Do not publish
Language quality	<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
Conclusion	<input type="checkbox"/> Accept (High priority) <input type="checkbox"/> Accept (General priority) <input checked="" type="checkbox"/> Minor revision <input type="checkbox"/> Major revision <input type="checkbox"/> Rejection
Re-review	<input checked="" type="checkbox"/> Yes <input type="checkbox"/> No



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Peer-reviewer statements	Peer-Review: [<input checked="" type="radio"/>] Anonymous [<input type="radio"/>] Onymous Conflicts-of-Interest: [<input type="radio"/>] Yes [<input checked="" type="radio"/>] No
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SPECIFIC COMMENTS TO AUTHORS

The manuscript was reviewed for publication in the journal. The manuscript was designed to review emerging novel targets for NASH treatment. It is the reviewer's opinion that the review is quite interesting and easy to follow. However, it appears that there are a couple of minor concerns in the manuscript. 1) The authors discussed the potential targets outside the liver. How about other targets such as the systemic or metabolic effects? The authors should discuss the issue. 2) The authors have discussed the cell specific effects for NASH treatment. How about the specific effects of liver immune cells such as lymphocytes, neutrophils, and other cells? The authors should discuss the issue.

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Manuscript NO: 80398

Title: Emerging novel targets for nonalcoholic fatty liver disease treatment: evidence from recent basic studies

Provenance and peer review: Invited Manuscript; Externally peer reviewed

Peer-review model: Single blind

Reviewer's code: 06360124

Position: Peer Reviewer

Academic degree: PhD

Professional title: Assistant Professor

Reviewer's Country/Territory: Iran

Author's Country/Territory: China

Manuscript submission date: 2022-10-04

Reviewer chosen by: AI Technique

Reviewer accepted review: 2022-11-01 18:36

Reviewer performed review: 2022-11-10 05:58

Review time: 8 Days and 11 Hours

Scientific quality	<input type="checkbox"/> Grade A: Excellent <input type="checkbox"/> Grade B: Very good <input checked="" type="checkbox"/> Grade C: Good <input type="checkbox"/> Grade D: Fair <input type="checkbox"/> Grade E: Do not publish
Language quality	<input type="checkbox"/> Grade A: Priority publishing <input checked="" type="checkbox"/> Grade B: Minor language polishing <input type="checkbox"/> Grade C: A great deal of language polishing <input type="checkbox"/> Grade D: Rejection
Conclusion	<input type="checkbox"/> Accept (High priority) <input type="checkbox"/> Accept (General priority) <input checked="" type="checkbox"/> Minor revision <input type="checkbox"/> Major revision <input type="checkbox"/> Rejection
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Peer-reviewer statements	Peer-Review: [<input checked="" type="radio"/>] Anonymous [<input type="radio"/>] Onymous Conflicts-of-Interest: [<input type="radio"/>] Yes [<input checked="" type="radio"/>] No
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SPECIFIC COMMENTS TO AUTHORS

This review focuses on recent researches reporting the mechanism underlying the pathogenesis of NASH. This research also tries to compile signaling pathways targeting key mechanisms that contribute to NASH development. The authors try to open new insights into explored the mechanism of NASH and its novel therapeutic targets. This study focused on basic cellular and molecular mechanisms and did not discuss subsequent pathological and also therapeutic outcomes in every research. However, it seems that approaches target signaling pathways driving disease development can use to inform new treatments for precision medicine. Therefore, it may provide a novel platform to identify and prioritize novel targets for NAFLD and NASH, creating a path for drug developers towards a first-in-class treatment.