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## PEER-REVIEW REPORT

Name of journal: World Journal of Gastroenterology

Manuscript NO: 89837

**Title:** Bile acids inhibit ferroptosis sensitivity through activating farnesoid X receptor in

gastric cancer cells

Provenance and peer review: Unsolicited Manuscript; Externally peer reviewed

Peer-review model: Single blind

Reviewer's code: 07746167 Position: Peer Reviewer Academic degree: PhD

**Professional title:** Researcher

**Reviewer's Country/Territory:** Germany

Author's Country/Territory: China

Manuscript submission date: 2023-11-21

Reviewer chosen by: AI Technique

Reviewer accepted review: 2023-11-27 05:40

Reviewer performed review: 2023-12-07 01:35

**Review time:** 9 Days and 19 Hours

	[ ] 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	[ ] Grade D: Fair [ ] Grade E: Do not publish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	[ ] Grade A: Excellent [ Y] Grade B: Good [ ] 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



## **Baishideng**

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Scientific significance of the	[ ] Grade A: Excellent [ Y] Grade B: Good [ ] Grade C: Fair
conclusion in this manuscript	[ ] 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) [ ] 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

During gastric cancer development in both humans and animals, bile acids serve as signaling molecules that induce metabolic reprogramming. This confers additional cancer phenotypes, including ferroptosis sensitivity. Ferroptosis is a novel mode of cell death characterized by lipid peroxidation that contributes universally to malignant progression. However, it is not fully defined if bile acids can influence gastric cancer progression by modulating ferroptosis. In this study, the authors treated gastric cancer cells with various stimuli and evaluated the effect of bile acids on the sensitivity to ferroptosis, and aimed to reveal the mechanism of bile acids regulation in ferroptosis of gastric cancer cells. This study is well designed and performed. The results are very interesting. The reviewer recommends to accept this study after a minor revision. Comments: 1. The manuscript requires a minor editing. 2. Some Greek characters seems can't be read, please take attention about it. 3. Quality of images should be improved.