

## PEER-REVIEW REPORT

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

**Manuscript NO:** 82025

**Title:** Clinical features, Diagnosis and Treatment Experience of Chinese 566 Patients with Peutz-Jeghers Syndrome

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

**Peer-review model:** Single blind

**Reviewer's code:** 06135158

**Position:** Peer Reviewer

**Academic degree:** MD, PhD

**Professional title:** Associate Professor

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

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

**Manuscript submission date:** 2022-12-12

**Reviewer chosen by:** AI Technique

**Reviewer accepted review:** 2022-12-16 10:04

**Reviewer performed review:** 2023-01-07 13:59

**Review time:** 22 Days and 3 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 Grade E: Do not publish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 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
Re-review	<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**

Mucocutaneous pigmentation generally does not require specific treatment, but PJS polyposis is clinically serious. Gastrointestinal polyps cause secondary rupture, bleeding, intussusception, intestinal obstruction, abdominal pain, abdominal distension, hematochezia and other symptoms, and their further development causes enteric necrosis, intestinal perforation and even cancer. In this study, the authors summarize and analyzes such experience in order to explore the clinicopathological features, diagnosis and treatment experience in PJS from a Chinese single-center. The study is well designed and the results are very attractive. The sample size is very large, and the data in the results are well discussed. Minor comments: 1. There are some minor language polishing which should be corrected. 2. The data in the results, include the general data of the patients, the first treatment age, etc., should be included in tables. 3. The references should be edited and updated. 4. The figures should be replaced with high resolution ratio images.

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**Reviewer's code:** 06110579

**Position:** Peer Reviewer

**Academic degree:** MD

**Professional title:** Doctor

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

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

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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
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<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 is a very interesting study of the clinical features, diagnosis and treatment experience of Peutz-Jeghers Syndrome. The sample is very big, and the experiences the authors shared are very important to the clinicians. The reviewer recommends to accept and publish this manuscript after a minor editing.