

# PEER-REVIEW REPORT

Name of journal: World Journal of Clinical Cases

Manuscript NO: 86172

Title: Multi-organ hereditary hemorrhagic telangiectasia: A case report

Provenance and peer review: Unsolicited Manuscript; Externally peer reviewed

**Peer-review model:** Single blind

Reviewer's code: 03853274

**Position:** Peer Reviewer

Academic degree: MD

Professional title: Academic Fellow

Reviewer's Country/Territory: Turkey

Author's Country/Territory: China

Manuscript submission date: 2023-07-11

Reviewer chosen by: AI Technique

Reviewer accepted review: 2023-07-18 04:47

Reviewer performed review: 2023-07-20 11:16

**Review time:** 2 Days and 6 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	[ ] Grade D: Fair [ ] Grade E: Do not publish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 this manuscript	[Y] Grade A: Excellent [] Grade B: Good [] Grade C: Fair [] Grade D: No creativity or innovation



# Baishideng

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

I have carefully read the case report. It is about a rare disease with q novel mutation. I have really enjoyed reading the case report however there are some issues to be touched. The case report definitely has novelty and it is appropriately summarized. I will make several article suggestions to enrich the Discussion section. 1. Olivieri C, Pagella F, Semino L, Lanzarini L, Valacca C, Pilotto A, et al. Analysis of ENG and ACVRL1 genes in 137 HHT Italian families identifies 76 different mutations (24 novel). Comparison with other European studies. J Hum Genet. 2007;52:820-9. 2. Karlsson T, Cherif H. Mutations in the ENG, ACVRL1, and SMAD4 genes and clinical manifestations of hereditary haemorrhagic telangiectasia: experience from the Center for Osler's Disease, Uppsala University Hospital. Ups J Med Sci. 2018;123:153–7. 3.Bossler AD, Richards J, George C, Godmilow L, Ganguly A. Novel mutations in ENG and ACVRL1 identified in a series of 200 individuals undergoing clinical genetic testing for hereditary hemorrhagic telangiectasia (HHT): correlation of genotype with phenotype. Hum Mutat. 2006;27:667-75 4.. Letteboer TG, Zewald RA, Kamping EJ, de Haas G, Mager JJ, Snijder RJ, et al. Hereditary hemorrhagic telangiectasia: ENG and ALK-1 mutations in Dutch patients.



Hum Genet. 2005;116:8–16. 5. Lesca G, Burnichon N, Raux G, Tosi M, Pinson S, Marion MJ, et al. Distribution of ENG and ACVRL1 (ALK1) mutations in French HHT patients. Hum Mutat. 2006;27:598. 6.Genetic Diagnosis of Hereditary Hemorrhagic Telangiectasia: Four Novel Pathogenic Variations in Turkish Patients Mehmet Baysal 7.A Novel Variation in the ACVRL1 Gene in a Patient with Cirrhosis and Hereditary Hemorrhagic Telangiectasia Mehmet Baysal These are important and pioneer works and ion my opinion should be cited in the case report.



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Provenance and peer review: Unsolicited Manuscript; Externally peer reviewed

**Peer-review model:** Single blind

Reviewer's code: 01221789

**Position:** Editorial Board

Academic degree: MBBS, MD

Professional title: Associate Professor

Reviewer's Country/Territory: India

Author's Country/Territory: China

Manuscript submission date: 2023-07-11

Reviewer chosen by: Geng-Long Liu

Reviewer accepted review: 2023-07-22 04:30

Reviewer performed review: 2023-07-30 11:41

**Review time:** 8 Days and 7 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 [ ] Grade B: Good [Y] Grade C: Fair [ ] Grade D: No novelty
Creativity or innovation of this manuscript	<ul> <li>[ ] Grade A: Excellent [ ] Grade B: Good [ Y] Grade C: Fair</li> <li>[ ] Grade D: No creativity or innovation</li> </ul>



Scientific significance of the conclusion in this manuscript	<ul> <li>[ ] Grade A: Excellent [Y] Grade B: Good [ ] Grade C: Fair</li> <li>[ ] Grade D: No scientific significance</li> </ul>
Language quality	[ ] Grade A: Priority publishing [Y] Grade B: Minor language polishing [ ] Grade C: A great deal of language polishing [ ] Grade D: Rejection
Conclusion	<ul> <li>[ ] Accept (High priority) [Y] Accept (General priority)</li> <li>[ ] Minor revision [ ] Major revision [ ] Rejection</li> </ul>
Re-review	[Y]Yes []No
Peer-reviewer statements	Peer-Review: [Y] Anonymous [] Onymous Conflicts-of-Interest: [] Yes [Y] No

### SPECIFIC COMMENTS TO AUTHORS

Please edit some of the spelling and grammatical errors.