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PEER-REVIEW REPORT

Name of journal: World Journal of Clinical Cases

Manuscript NO: 77809

Title: Familial mitochondrial encephalomyopathy, lactic acidosis, and stroke-like

episode syndrome: Three case reports

Provenance and peer review: Unsolicited Manuscript; Externally peer reviewed

Peer-review model: Single blind

Reviewer's code: 04543416 Position: Peer Reviewer Academic degree: MD

Professional title: Chief Doctor, Professor

Reviewer's Country/Territory: Turkey

Author's Country/Territory: China

Manuscript submission date: 2022-05-21

Reviewer chosen by: AI Technique

Reviewer accepted review: 2022-05-22 20:13

Reviewer performed review: 2022-05-28 19:12

Review time: 5 Days and 22 Hours

Scientific quality	[] Grade A: Excellent [] Grade B: Very good [Y] Grade C: Good [] Grade D: Fair [] Grade E: Do not publish
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



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Peer-reviewer

Peer-Review: [Y] Anonymous [] Onymous

statements Conflicts-of-Interest: [] Yes [Y] No

SPECIFIC COMMENTS TO AUTHORS

On MRI figure legends, axial plane MR should be used instead of "Cross-sectional" expression. In the first sentences describing the images of the cases in the imaging examinations section, the expressions "hypointense T1 or hyperintense T2" are preferred instead of "long T1 and long T2 signals". Please make the necessary changes.



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Reviewer's code: 05477765 Position: Peer Reviewer Academic degree: PhD

Professional title: Assistant Professor, Senior Researcher

Reviewer's Country/Territory: Russia

Author's Country/Territory: China

Manuscript submission date: 2022-05-21

Reviewer chosen by: Dong-Mei Wang

Reviewer accepted review: 2022-07-19 06:10

Reviewer performed review: 2022-07-19 18:09

Review time: 11 Hours

Scientific quality	[] Grade A: Excellent [Y] Grade B: Very good [] Grade C: Good [] Grade D: Fair [] Grade E: Do not publish
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	[Y]Yes []No



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SPECIFIC COMMENTS TO AUTHORS

The manuscript of "Familial mitochondrial encephalomyopathy with lactic acidosis and stroke-like episode syndrome : a case report" by Xiao Yang and Le-Jun Fu aims to review clinical manifestations, muscle pathological examination, and genetic testing in three clinical cases of familial MELAS syndrome. The manuscript is well written, and it describes some recommendations for the management of patients with MELAS syndrome and their families. The manuscript may be accepted for publication in its Comment: In the Discussion section, the role of mtDNA A3243G, current form. A8344G and T8933C (G) gene mutations found in the patients can be discussed in more detail.