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PEER-REVIEW REPORT

Name of journal: World Journal of Clinical Cases

Manuscript NO: 73201

Title: Growth hormone ameliorates hepatopulmonary syndrome and nonalcoholic

steatohepatitis secondary to hypopituitarism in children

Provenance and peer review: Unsolicited Manuscript; Externally peer reviewed

Peer-review model: Single blind

Reviewer's code: 02959015 Position: Editorial Board Academic degree: MD

Professional title: Associate Professor

Reviewer's Country/Territory: Italy

Author's Country/Territory: China

Manuscript submission date: 2021-11-13

Reviewer chosen by: AI Technique

Reviewer accepted review: 2021-11-25 08:03

Reviewer performed review: 2021-11-26 11:54

Review time: 1 Day and 3 Hours

Scientific quality	[] Grade A: Excellent [] Grade B: Very good [Y] Grade C: Good [] Grade D: Fair [] Grade E: Do not publish
Language quality	[Y] Grade A: Priority publishing [] 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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Peer-reviewer

Peer-Review: [Y] Anonymous [] Onymous

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

SPECIFIC COMMENTS TO AUTHORS

The authors describe a case of a 13-year-old female subject who, six years from surgery for craniopharyngioma (CP) not followed by hormone replacement therapy, developed a post-NASH liver cirrhosis. 1) The authors should better highlight that NAFLD can arise in craniopharyngioma (CP) even before surgery, and it is likely that in the case described it was present at the time of surgery and then it worsened after resection of the CP in the absence of replacement therapy 2) It is necessary that they further underline how the literature shows that hormone replacement therapy is necessary even after liver Results: Why HCV has not been tested? 4) transplantation. 3) Table 1: please report normal values 5) In Figure 2 there are no dilated biliary tracts but the left branch of the portal vein so please remove or replace 6) It would be appropriate for the authors to report the MELD value and in which Child Pugh Class was the patient 7)

Discussion: In the discussion the authors state that in the past it was believed that NFLD is associated with insulin resistance then in the next paragraph they write "However, in recent years, it has been found that more and more endocrine diseases, such as polycystic ovary syndrome, hypothalamic hypopituitarism, hypothyroidism, and acanthosis nigricans, can also lead to NAFLD". The two concepts give the impression of being in contradiction, the authors should better define this concept since most of the diseases they cite (polycystic ovary syndrome, acanthosis nigricans) are related to NAFLD precisely because of insulin resistance.



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Reviewer's code: 05226306 **Position:** Editorial Board

Academic degree: FACS, MBBS, MCh, MD, MNAMS

Professional title: Additional Professor

Reviewer's Country/Territory: India

Author's Country/Territory: China

Manuscript submission date: 2021-11-13

Reviewer chosen by: Xin Liu (Online Science Editor)

Reviewer accepted review: 2022-01-20 10:21

Reviewer performed review: 2022-01-20 12:33

Review time: 2 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 [] Grade B: Minor language polishing [Y] Grade C: A great deal of language polishing [] Grade D: Rejection
Conclusion	[] Accept (High priority) [] Accept (General priority) [] Minor revision [Y] Major revision [] Rejection
Re-review	[]Yes [Y]No



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Peer-reviewer

Peer-Review: [Y] Anonymous [] Onymous

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

SPECIFIC COMMENTS TO AUTHORS

1. Introduction to be elaborate detailing the background and also include the relevant reference citations. 2. What were the anthropometrics (height, weight) with respect to age standards? 3. Is the statement valid -"The patient had no history of high blood tumor". pressure, diabetes, heart disease, The child had undergone or Craniopharyngioma surgery before. 4. The lab results can be better presented in a table format 5. Relevant details of hormones given to be included 6. Is the pt O2 dependent? ".....currently fluctuates around 92% for low flow nasal catheter oxygen." 7. What was the protocol for Growth hormone administration (dose, frequency)? 8. What are the available options for NAFLD? Are options like Life style modification / early screening to identify transaminase > 40 valid in post craniopharyngioma surgery? 9. The point of early identification / intervention is crucial and can be highlighted 10. What are the controversies in Growth Hormone therapy?



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Reviewer's code: 03262105 Position: Peer Reviewer

Academic degree: FRACP, MD

Professional title: Professor

Reviewer's Country/Territory: Turkey

Author's Country/Territory: China

Manuscript submission date: 2021-11-13

Reviewer chosen by: Xin Liu (Online Science Editor)

Reviewer accepted review: 2022-01-14 06:26

Reviewer performed review: 2022-01-27 05:59

Review time: 12 Days and 23 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 [] 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
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Peer-reviewer

Peer-Review: [Y] Anonymous [] Onymous

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

SPECIFIC COMMENTS TO AUTHORS

The name of the manuscript shall include knowledge about craniopharingioma diagnosis and surgery. English mistakes shall be corrected. Such as A 13 years and six months girl went craniopharyngioma surgery 6 years, 1 year later, she exhibited abdominal distension... A 13 years and six months girl underwent craniopharyngioma surgery 6 years ago. One year later, she exhibited abdominal distension Craniopharyngioma surgery is easy to cause hypopituitarism Craniopharyngioma frequently causes hypopituitarism.