



PEER-REVIEW REPORT

Name of journal: World Journal of Clinical Cases

Manuscript NO: 42995

Title: CONGENITAL ANALBUMINEMIA IN A PATIENT AFFECTED BY HYPERCHOLESTEROLEMIA

Reviewer’s code: 03531928

Reviewer’s country: China

Science editor: Fang-Fang Ji

Date sent for review: 2018-10-18

Date reviewed: 2018-10-19

Review time: 1 Day

SCIENTIFIC QUALITY	LANGUAGE QUALITY	CONCLUSION	PEER-REVIEWER STATEMENTS
<input type="checkbox"/> Grade A: Excellent	<input type="checkbox"/> Grade A: Priority publishing	<input type="checkbox"/> Accept	Peer-Review:
<input checked="" type="checkbox"/> Grade B: Very good	<input checked="" type="checkbox"/> Grade B: Minor language polishing	(High priority)	<input type="checkbox"/> Anonymous
<input type="checkbox"/> Grade C: Good		<input checked="" type="checkbox"/> Accept	<input checked="" type="checkbox"/> Onymous
<input type="checkbox"/> Grade D: Fair	<input type="checkbox"/> Grade C: A great deal of language polishing	(General priority)	Peer-reviewer’s expertise on the topic of the manuscript:
<input type="checkbox"/> Grade E: Do not publish	<input type="checkbox"/> Grade D: Rejection	<input type="checkbox"/> Minor revision	<input checked="" type="checkbox"/> Advanced
		<input type="checkbox"/> Major revision	<input type="checkbox"/> General
		<input type="checkbox"/> Rejection	<input type="checkbox"/> No expertise
			Conflicts-of-Interest:
			<input type="checkbox"/> Yes
			<input checked="" type="checkbox"/> No

SPECIFIC COMMENTS TO AUTHORS

CAA is a rare autosomal recessive disease. There are only dozens of cases in the world. This case report describes detailedly the clinical features and laboratory examinations of a patient with congenital hypoalbuminemia presenting with hypercholesterolemia. More



**Baishideng
Publishing
Group**

7901 Stoneridge Drive, Suite 501,
Pleasanton, CA 94588, USA
Telephone: +1-925-223-8242
Fax: +1-925-223-8243
E-mail: bpgoffice@wjgnet.com
https://www.wjgnet.com

importantly, ALB gene mutation fragments were detected and analyzed in a patient's family to provide a genetic basis for the diagnosis of the disease. It was also found that the disease did not require albumin infusion, and that atorvastatin was safe and effective in controlling high cholesterol in patients with CCA, helping to reduce their risk of cardiovascular disease. The report is helpful in raising awareness of the disease, providing evidence for genetic diagnosis and related treatment experience, and is recommended for publication.

INITIAL REVIEW OF THE MANUSCRIPT

Google Search:

- The same title
- Duplicate publication
- Plagiarism
- No

BPG Search:

- The same title
- Duplicate publication
- Plagiarism
- No



PEER-REVIEW REPORT

Name of journal: World Journal of Clinical Cases

Manuscript NO: 42995

Title: CONGENITAL ANALBUMINEMIA IN A PATIENT AFFECTED BY HYPERCHOLESTEROLEMIA

Reviewer's code: 02575809

Reviewer's country: Angola

Science editor: Fang-Fang Ji

Date sent for review: 2018-10-18

Date reviewed: 2018-10-20

Review time: 2 Days

SCIENTIFIC QUALITY	LANGUAGE QUALITY	CONCLUSION	PEER-REVIEWER STATEMENTS
<input type="checkbox"/> Grade A: Excellent	<input type="checkbox"/> Grade A: Priority publishing	<input type="checkbox"/> Accept	Peer-Review:
<input type="checkbox"/> Grade B: Very good	<input checked="" type="checkbox"/> Grade B: Minor language polishing	(High priority)	<input checked="" type="checkbox"/> Anonymous
<input checked="" type="checkbox"/> Grade C: Good		<input type="checkbox"/> Accept	<input type="checkbox"/> Onymous
<input type="checkbox"/> Grade D: Fair	<input type="checkbox"/> Grade C: A great deal of language polishing	(General priority)	Peer-reviewer's expertise on the topic of the manuscript:
<input type="checkbox"/> Grade E: Do not publish	<input type="checkbox"/> Grade D: Rejection	<input checked="" type="checkbox"/> Minor revision	<input type="checkbox"/> Advanced
		<input type="checkbox"/> Major revision	<input checked="" type="checkbox"/> General
		<input type="checkbox"/> Rejection	<input type="checkbox"/> No expertise
			Conflicts-of-Interest:
			<input type="checkbox"/> Yes
			<input checked="" type="checkbox"/> No

SPECIFIC COMMENTS TO AUTHORS

Comments for the authors The article reports a case with a low incidence and with few reports in the literature. There are abbreviations that should be described the first time they appear in the text. Page 4, ... homozygous for a CA deletion... On page 4



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paragraph 3 says... This observation together with the low blood pressure might explain why, despite the high lipid levels, evident clinical signs of early atherosclerosis have not been observed in the patient[3]. In my opinion there are few diagnostic evidence in this patient, or they are not described in the text to say that there are no evident clinical signs of early atherosclerosis. In Discussion he say ... In this case the diagnosis of CCA was not...Review the abbreviation. ... In conclusion, according to our experience, safe and well tolerated hypocholesterolemic treatment with atorvastatin may be administered in dislipidemic patient with CCA should be associated... Review the abbreviation. The reference number 8, missing the title of the article

INITIAL REVIEW OF THE MANUSCRIPT

Google Search:

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BPG Search:

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