

PEER-REVIEW REPORT

Name of journal: World Journal of Gastroenterology

Manuscript NO: 41105

Title: Ductopenia and cirrhosis in a 32-year-old woman with progressive familial intrahepatic cholestasis type 3: A case report

Reviewer's code: 02942549

Reviewer's country: Greece

Science editor: Xue-Jiao Wang

Date sent for review: 2018-08-01

Date reviewed: 2018-08-03

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 checked="" type="checkbox"/> Grade B: Very good	<input checked="" type="checkbox"/> Grade B: Minor language	(High priority)	<input type="checkbox"/> Anonymous
<input type="checkbox"/> Grade C: Good	polishing	<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	(General priority)	Peer-reviewer's expertise on the
<input type="checkbox"/> Grade E: Do not	language polishing	<input type="checkbox"/> Minor revision	topic of the manuscript:
publish	<input type="checkbox"/> Grade D: Rejection	<input type="checkbox"/> Major revision	<input type="checkbox"/> Advanced
		<input type="checkbox"/> Rejection	<input checked="" type="checkbox"/> General
			<input type="checkbox"/> No expertise
			Conflicts-of-Interest:
			<input type="checkbox"/> Yes
			<input checked="" type="checkbox"/> No

SPECIFIC COMMENTS TO AUTHORS

It is an interesting case study. The authors analyzed every aspect of it by the most appropriate way.



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INITIAL REVIEW OF THE MANUSCRIPT

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- ☐ The same title
- ☐ Duplicate publication
- ☐ Plagiarism
- ☐ No

BPG Search:

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PEER-REVIEW REPORT

Name of journal: World Journal of Gastroenterology

Manuscript NO: 41105

Title: Ductopenia and cirrhosis in a 32-year-old woman with progressive familial intrahepatic cholestasis type 3: A case report

Reviewer's code: 02943023

Reviewer's country: South Korea

Science editor: Xue-Jiao Wang

Date sent for review: 2018-07-25

Date reviewed: 2018-08-12

Review time: 17 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 type="checkbox"/> Grade B: Minor language	(High priority)	<input type="checkbox"/> Anonymous
<input checked="" type="checkbox"/> Grade C: Good	polishing	<input type="checkbox"/> Accept	<input type="checkbox"/> Onymous
<input type="checkbox"/> Grade D: Fair	<input checked="" type="checkbox"/> Grade C: A great deal of	(General priority)	Peer-reviewer's expertise on the
<input type="checkbox"/> Grade E: Do not	language polishing	<input checked="" type="checkbox"/> Minor revision	topic of the manuscript:
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		<input type="checkbox"/> Rejection	<input checked="" type="checkbox"/> General
			<input type="checkbox"/> No expertise
			Conflicts-of-Interest:
			<input type="checkbox"/> Yes
			<input checked="" type="checkbox"/> No

SPECIFIC COMMENTS TO AUTHORS

In this manuscript, authors reported a case of PFIC3 in a 32-year-woman based on clinical symptoms, pathological findings, and gene mutation detection. It is interesting case suggesting important differential diagnosis in a patients with recurrent cholestasis. I

would like to indicate some minor comments. 1. Since the clinical features of PFIC3 overlap with many other forms of liver disease in childhood, definitive diagnosis may be problematic or delayed. In addition, impaired copper secretion and copper accumulation can be seen in all chronic cholestatic disorders. In this case report, liver histology showed copper-associated protein sinking in hepatocyte cytoplasm (Figure 1C). Therefore, please describe the findings for differential diagnosis with Wilson disease (WD) in this case, such as serum ceruloplasmin, serum and 24-hour urinary copper level, and liver histology etc. In addition, it would be better to comments about differential diagnosis or overlapping with WD in young patients with recurrent cholestasis and/or cirrhosis in 'Discussion'. 2. Please correct the errors in English.

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Name of journal: World Journal of Gastroenterology

Manuscript NO: 41105

Title: Ductopenia and cirrhosis in a 32-year-old woman with progressive familial intrahepatic cholestasis type 3: A case report

Reviewer's code: 00030603

Reviewer's country: Greece

Science editor: Xue-Jiao Wang

Date sent for review: 2018-08-01

Date reviewed: 2018-08-13

Review time: 12 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 checked="" type="checkbox"/> Grade B: Very good	<input checked="" type="checkbox"/> Grade B: Minor language	(High priority)	<input checked="" type="checkbox"/> Anonymous
<input type="checkbox"/> Grade C: Good	polishing	<input checked="" type="checkbox"/> Accept	<input type="checkbox"/> Onymous
<input type="checkbox"/> Grade D: Fair	<input type="checkbox"/> Grade C: A great deal of	(General priority)	Peer-reviewer's expertise on the
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		<input type="checkbox"/> Rejection	<input checked="" type="checkbox"/> General
			<input type="checkbox"/> No expertise
			Conflicts-of-Interest:
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SPECIFIC COMMENTS TO AUTHORS

interesting case reminding a rare cause of cholestasis

INITIAL REVIEW OF THE MANUSCRIPT



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Name of journal: World Journal of Gastroenterology

Manuscript NO: 41105

Title: Ductopenia and cirrhosis in a 32-year-old woman with progressive familial intrahepatic cholestasis type 3: A case report

Reviewer's code: 03024199

Reviewer's country: Japan

Science editor: Xue-Jiao Wang

Date sent for review: 2018-08-01

Date reviewed: 2018-08-19

Review time: 18 Days

SCIENTIFIC QUALITY	LANGUAGE QUALITY	CONCLUSION	PEER-REVIEWER STATEMENTS
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			Conflicts-of-Interest:
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			<input checked="" type="checkbox"/> No

SPECIFIC COMMENTS TO AUTHORS

The authors described a rare case of progressive familial intrahepatic cholestasis type 3, by confirming a mutation in the ABCB4 gene encoding multidrug resistance protein 3. Interesting case presentation addressing a rare topic. I think it should be published in the



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