



**PEER-REVIEW REPORT**

**Name of journal:** World Journal of Gastroenterology

**Manuscript NO:** 40896

**Title:** A challenge to overcome: Nonstructural protein 5A- 2 deletion in direct-acting antiviral-based therapy for hepatitis C virus

**Reviewer's code:** 00052928

**Reviewer's country:** Reviewer\_Country

**Science editor:** Ze-Mao Gong

**Date sent for review:** 2018-07-17

**Date reviewed:** 2018-07-18

**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 checked="" type="checkbox"/> Anonymous
<input 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 Grade E: Do not publish	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**

This editorial focuses on the problem of the recently-found super resistant-associated substitution (RAS), S5A-P32 deletion, in HCV infection. The manuscript describes in detail as to the background situation of the appearance of P32 deletion, the relationship



**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**

of P32 deletion with other RASs, and the responses to the new DAA therapies. Minor: Though the author insists that “P32 deletion and Y93 mutants and L31 mutants are mutually exclusive”, P32 deletion is often accompanied by L31F mutation actually. Therefore, excessive emphasizing of the mutual exclusiveness might cause misunderstanding.

#### **INITIAL REVIEW OF THE MANUSCRIPT**

##### ***Google Search:***

- The same title
- Duplicate publication
- Plagiarism
- No

##### ***BPG Search:***

- The same title
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- No



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**Reviewer’s code:** 02540539

**Reviewer’s country:** Reviewer\_Country

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<input type="checkbox"/> Grade C: Good	polishing	<input 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
<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 type="checkbox"/> General
			<input type="checkbox"/> No expertise
			Conflicts-of-Interest:
			<input type="checkbox"/> Yes
			<input type="checkbox"/> No

**SPECIFIC COMMENTS TO AUTHORS**

P32 deletion is the hot topic in DAA-associated mutation. Overcoming the mutation is very important. Therefore, this manuscript has high grade priority to be published. The authors well described about P32 mutation comprehensively.



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Pleasanton, CA 94588, USA  
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