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ESPS PEER-REVIEW REPORT

Name of journal: World Journal of Gastrointestinal Oncology

ESPS manuscript NO: 19260

Title: Genomic alterations in pancreatic cancer and their relevance to therapy

Reviewer's code: 00069406

Reviewer's country: China

Science editor: Yue-Li Tian

Date sent for review: 2015-05-08 16:22

Date reviewed: 2015-05-18 11:20

CLASSIFICATION	LANGUAGE EVALUATION	SCIENTIFIC MISCONDUCT	CONCLUSION
<input type="checkbox"/> Grade A: Excellent	<input type="checkbox"/> Grade A: Priority publishing	Google Search:	<input type="checkbox"/> Accept
<input type="checkbox"/> Grade B: Very good	<input checked="" type="checkbox"/> Grade B: Minor language polishing	<input type="checkbox"/> The same title	<input type="checkbox"/> High priority for publication
<input type="checkbox"/> Grade C: Good	<input type="checkbox"/> Grade C: A great deal of language polishing	<input type="checkbox"/> Duplicate publication	<input checked="" type="checkbox"/> Rejection
<input checked="" type="checkbox"/> Grade D: Fair	<input type="checkbox"/> Grade D: Rejected	<input checked="" type="checkbox"/> Plagiarism	<input type="checkbox"/> Minor revision
<input type="checkbox"/> Grade E: Poor		[Y] No	<input type="checkbox"/> Major revision
		BPG Search:	
		<input type="checkbox"/> The same title	
		<input type="checkbox"/> Duplicate publication	
		<input type="checkbox"/> Plagiarism	
		[Y] No	

COMMENTS TO AUTHORS

The current review conclude the recent progress on genomic alterations in pancreatic cancer and their relevance to therapy. However, there was less novelty in the text to warrant published. KRAS, CDKN2A, TP53 and SMAD4 are the four genes frequently mutated in PDAC, which are already known for years. Also, there is no more progresses to be concluded on target therapy for PDAC. So the significance of review is fair. Figure 1 contains less information than which is already published in the literature.

ESPS PEER-REVIEW REPORT

Name of journal: World Journal of Gastrointestinal Oncology

ESPS manuscript NO: 19260

Title: Genomic alterations in pancreatic cancer and their relevance to therapy

Reviewer's code: 00224612

Reviewer's country: Germany

Science editor: Yue-Li Tian

Date sent for review: 2015-05-08 16:22

Date reviewed: 2015-05-08 21:12

CLASSIFICATION	LANGUAGE EVALUATION	SCIENTIFIC MISCONDUCT	CONCLUSION
<input type="checkbox"/> Grade A: Excellent	<input checked="" type="checkbox"/> Grade A: Priority publishing	Google Search:	<input type="checkbox"/> Accept
<input checked="" type="checkbox"/> Grade B: Very good	<input type="checkbox"/> Grade B: Minor language polishing	<input type="checkbox"/> The same title	<input type="checkbox"/> High priority for publication
<input type="checkbox"/> Grade C: Good		<input type="checkbox"/> Duplicate publication	
<input type="checkbox"/> Grade D: Fair	<input type="checkbox"/> Grade C: A great deal of language polishing	<input type="checkbox"/> Plagiarism	<input type="checkbox"/> Rejection
<input type="checkbox"/> Grade E: Poor	<input type="checkbox"/> Grade D: Rejected	<input checked="" type="checkbox"/> No	<input checked="" type="checkbox"/> Minor revision
		BPG Search:	<input type="checkbox"/> Major revision
		<input type="checkbox"/> The same title	
		<input type="checkbox"/> Duplicate publication	
		<input type="checkbox"/> Plagiarism	
		<input checked="" type="checkbox"/> No	

COMMENTS TO AUTHORS

The manuscript by Takai and Yachida is a well written, up-to-date and in-depth review regarding genomic alterations in pancreatic cancer and their relevance to therapy. There are some minor comments/suggestions: The authors state that "although great efforts have been made to develop small-molecular inhibitors of mutant KRAS, no clinically effective antagonist has yet been identified". In this context, the authors should cite and include: The Ras renaissance. PMID: 25877186. The authors state that "activation of the PI3K/Akt/mTOR pathway also plays an important role in maintenance of pancreatic cancer". In this context, the authors might also want to cite and include: A subset of metastatic pancreatic ductal adenocarcinomas depends quantitatively on oncogenic Kras/Mek/Erk-induced hyperactive mTOR signalling. PMID: 25601637. The authors state that "interestingly, blockade of the Hedgehog pathway has also been proposed as a means to target the tumor stroma and improve delivery of gemcitabine in vivo". The authors should also state and cite that the Saridegib trial failed. Figure 1 should be omitted since the authors do not review in depth pancreatic cancer progression models. A table depicting the core signalling pathways and current trials/targets for each pathway would be helpful.