

**ESPS Peer-review Report**
**Name of Journal:** World Journal of Gastroenterology

**ESPS Manuscript NO:** 7393

**Title:** Pathology of Pancreatic Ductal Adenocarcinoma: Facts, Challenges and Future Developments

**Reviewer code:** 01557573

**Science editor:** Gou, Su-Xin

**Date sent for review:** 2013-11-16 16:32

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CLASSIFICATION	LANGUAGE EVALUATION	RECOMMENDATION	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"/> Existed	<input type="checkbox"/> High priority for publication
<input checked="" type="checkbox"/> Grade C (Good)	<input type="checkbox"/> Grade C: a great deal of language polishing	<input type="checkbox"/> No records	<input type="checkbox"/> Rejection
<input type="checkbox"/> Grade D (Fair)	<input type="checkbox"/> Grade D: rejected	<input type="checkbox"/> Existed	<input checked="" type="checkbox"/> Minor revision
<input type="checkbox"/> Grade E (Poor)		<input type="checkbox"/> No records	<input type="checkbox"/> Major revision

**COMMENTS TO AUTHORS**

Pancreatic carcinoma remains one of the most lethal cancer type, its prognosis is so poor that the mortality is almost as same as the incidence. Currently, the most practical way to achieve a better prognosis is early diagnosis and the following curative resection. This manuscript reviews some very important issues that relevant to the points, including the pre-cancerous lesions, standardized pathological reporting regime and subtypes based on the genetic profiles. The language is good except sporadic mistakes. Clinicians, esp. pancreatic surgeons can obtain useful information from this manuscript.

# ESPS Peer-review Report

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

**ESPS Manuscript NO:** 7393

**Title:** Pathology of Pancreatic Ductal Adenocarcinoma: Facts, Challenges and Future Developments

**Reviewer code:** 01734797

**Science editor:** Gou, Su-Xin

**Date sent for review:** 2013-11-16 16:32

**Date reviewed:** 2013-11-26 17:52

CLASSIFICATION	LANGUAGE EVALUATION	RECOMMENDATION	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"/> Existed	<input checked="" 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"/> No records	<input type="checkbox"/> Rejection
<input type="checkbox"/> Grade D (Fair)	<input type="checkbox"/> Grade D: rejected	<input type="checkbox"/> Existed	<input type="checkbox"/> Minor revision
<input type="checkbox"/> Grade E (Poor)		<input type="checkbox"/> No records	<input type="checkbox"/> Major revision

## COMMENTS TO AUTHORS

The authors described an importance of pathological point of view at the interface between basic research and clinical management of pancreatic cancer. They raised three points: understanding precursor lesions, atypical flat lesions, evaluating whether R0 or R1 resection by axial slicing technique, and understanding three molecular subtypes of pancreatic ductal adenocarcinoma. This might be a useful review for both, basic researchers and clinical physicians and surgeons, to tackle the difficult cancer from a bit new, different perspectives. In the discussion of precursor lesions, the authors mentioned about IPMN and its genetic alteration, "the presence of KRAS mutations as common characteristics of PDAC precursors." However, it is known that gastric subtype IPMN, which is mostly low-grade, has more frequent KRAS mutation than intestinal subtype IPMN, which is frequently high-grade or sometimes IPMC. Moreover, branch duct IPMN patients (most of them are low-grade gastric subtype) frequently have PDAC at the distinct area in the same pancreas. Therefore, the relationship of IPMN and PDAC is not so simple. The authors should have comments on this. The authors mentioned about three molecular subtypes of PDAC, classical, quasi-mesenchymal, and exocrine-like subtypes. They discussed differences between the classical and quasi-mesenchymal subtypes, but not exocrine-like subtype at all. The authors should explain a little more about the exocrine-like subtype for readers to understand.