

## ESPS PEER-REVIEW REPORT

**Name of journal:** World Journal of Gastrointestinal Surgery

**ESPS manuscript NO:** 12625

**Title:** Glucagon receptor gene mutations with hyperglucagonemia but without the glucagonoma syndrome

**Reviewer's code:** 00068250

**Reviewer's country:** China

**Science editor:** Yue-Li Tian

**Date sent for review:** 2014-07-18 10:57

**Date reviewed:** 2014-11-15 09:40

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

## COMMENTS TO AUTHORS

1 This is an interesting case of an entity not described before. The following points are provided for its improvement. 2 Serum glucagon level should be tested repeatedly and dynamically especially after the biopsy because it was not always 66 pmol/L. 3 The clinical manifestation of the patient could be described in more detail. The only symptom of abdominal pain might be compared in features with other cases and stated about the management in the last decade. 4 Had the gall stone ever be detected in the past? 5 Are there other mutations in the GCGR gene related to this condition? Future study on this entity of disease can be suggested.