

## ESPS PEER-REVIEW REPORT

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

**ESPS manuscript NO:** 14854

**Title:** Comparison of two proton pump inhibitor treatment regimens in Chinese patients diagnosed with chronic gastritis

**Reviewer's code:** 00039368

**Reviewer's country:** Estonia

**Science editor:** Ya-Juan Ma

**Date sent for review:** 2014-10-28 10:44

**Date reviewed:** 2014-12-11 18:59

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

## COMMENTS TO AUTHORS

This very well designed, performed and written multicenter, randomized, open-label study considers the comparison of symptom control between the recommended proton pump inhibitor (PPI) treatment regimens for non-erosive reflux disease (8 weeks) and chronic gastritis (2 weeks) in 305 Chinese patients who have typical reflux symptoms and negative endoscopy. This study confirms the 8-week PPI regimen provided marginally better symptom control and relief rates than the 2-week regimen, with a similar safety profile. This study is making a great contribution to randomized clinical trial studies lead to optimization of gastro-esophageal reflux disease (GERD) treatment. This is a very well written and set up clinical trial study. The authors give a well and clear overview about the study background and raised clearly the aim of the study, which is fulfilled. The authors present very detailed description of the patients included in the study, of assessments and treatments. The statistical analysis was specified and described very carefully. The material studied is large enough and allows to draw the conclusions. The two Tables and 3 figure presented in the paper are accurate, detailed and give a good overview about the studied materials and results.



## BAISHIDENG PUBLISHING GROUP INC

8226 Regency Drive, Pleasanton, CA 94588, USA

Telephone: +1-925-223-8242

Fax: +1-925-223-8243

E-mail: [bpgoffice@wjgnet.com](mailto:bpgoffice@wjgnet.com)

<http://www.wjgnet.com>

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The Results are presented clearly and have been discussed well.

## ESPS PEER-REVIEW REPORT

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

**ESPS manuscript NO:** 14854

**Title:** Comparison of two proton pump inhibitor treatment regimens in Chinese patients diagnosed with chronic gastritis

**Reviewer's code:** 00183445

**Reviewer's country:** Poland

**Science editor:** Ya-Juan Ma

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**Date reviewed:** 2014-12-16 02:05

CLASSIFICATION	LANGUAGE EVALUATION	SCIENTIFIC MISCONDUCT	CONCLUSION
<input type="checkbox"/> Grade A: Excellent	<input type="checkbox"/> Grade A: Priority publishing	PubMed Search:	<input checked="" 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		[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 aim of this study was to compare symptom control between the recommended proton pump inhibitor (PPI, omeprazole 20 mg once daily) treatment regimens: 8 weeks versus 2 weeks, in the patients who have typical reflux symptoms and a negative endoscopy. The symptom control rate showed a small but statistically significant difference in favour of the 8-week regimen. Fewer unscheduled visits and higher patients satisfaction supported the therapeutic benefits of the 8-week regimen versus the 2 week regimen. Both regimens were safe and well tolerated. This is well constructed multicenter, randomized study with high practical value. The principle of patients selection was explained in details including the broad range of exclusion. The procedures of treatment and assessment are clear. Safety assessments included monitoring of serious adverse events and discontinuation due such events of any severity. Statistical analysis is clear.