

8226 Regency Drive, Pleasanton, CA 94588, USA
Telephone: +1-925-223-8242
E-mail: bpgoffice@wjgnet.com http://www.wjgnet.com

ESPS PEER-REVIEW REPORT

Name of journal: World Journal of Gastroenterology

ESPS manuscript NO: 29010

Title: Effects of asymmetrical dimethylarginine on renal arteries in portal hypertension

and cirrhosis

Reviewer's code: 00068278 Reviewer's country: Turkey

Science editor: Ze-Mao Gong

Date sent for review: 2016-07-30 19:18

Date reviewed: 2016-08-09 18:53

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

COMMENTS TO AUTHORS

Asymmetric dimethylarginine (ADMA) is a new molecule that tested its value as a marker is being tested for many diseases and situations; cardiovascular diseases, statin usage, etc. In the presented study the effects of ADMA on basal and induced releases of nitric oxide (NO) in renal arteries from portal hypertensive and cirrhotic rats were investigated and was shown that ADMA inhibited both basal and induced NO in renal arteries. This inhibition may prevent the relaxation of renal arteries and may contribute to renal impairement. The study is a well designed and conducted one. It may contribute to the pathophysiology and to the development strategies to prevent/treat of hepatorenal syndrome.



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ESPS PEER-REVIEW REPORT

Name of journal: World Journal of Gastroenterology

ESPS manuscript NO: 29010

Title: Effects of asymmetrical dimethylarginine on renal arteries in portal hypertension

and cirrhosis

Reviewer's code: 00068215 Reviewer's country: Romania Science editor: Ze-Mao Gong

Date sent for review: 2016-07-30 19:18

Date reviewed: 2016-08-16 01:45

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

COMMENTS TO AUTHORS

Please, check the english language



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ESPS PEER-REVIEW REPORT

Name of journal: World Journal of Gastroenterology

ESPS manuscript NO: 29010

Title: Effects of asymmetrical dimethylarginine on renal arteries in portal hypertension

and cirrhosis

Reviewer's code: 00182864 Reviewer's country: Turkey Science editor: Ze-Mao Gong

Date sent for review: 2016-07-30 19:18

Date reviewed: 2016-08-19 16:14

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

COMMENTS TO AUTHORS

Dear AUTHORS, This study seems to be good enough. Sincerely



8226 Regency Drive, Pleasanton, CA 94588, USA
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ESPS PEER-REVIEW REPORT

Name of journal: World Journal of Gastroenterology

ESPS manuscript NO: 29010

Title: Effects of asymmetrical dimethylarginine on renal arteries in portal hypertension

and cirrhosis

Reviewer's code: 00050424 Reviewer's country: Greece Science editor: Ze-Mao Gong

Date sent for review: 2016-07-30 19:18

Date reviewed: 2016-08-22 05:49

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

COMMENTS TO AUTHORS

It is a well written article. The study is well designed. I do not know the clinical significance of these results.



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ESPS PEER-REVIEW REPORT

Name of journal: World Journal of Gastroenterology

ESPS manuscript NO: 29010

Title: Effects of asymmetrical dimethylarginine on renal arteries in portal hypertension

and cirrhosis

Reviewer's code: 00034635 Reviewer's country: Spain

Science editor: Ze-Mao Gong

Date sent for review: 2016-07-30 19:18

Date reviewed: 2016-08-24 12:29

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

COMMENTS TO AUTHORS

In this manuscript Segarra et al studied the effect of asymmetric dimethyl-arginine (ADMA) on the renal arteries of two models of murine portal hypertension, one of the pre-hepatic and the other one biliary cirrhosis. Whilst ADMA competes with L-Arginine (the precursor of endothelial NO) and competitively inhibits NO-synthase, its role in the renal dysfunction of cirrhosis and even in human pathology remains unclear. These results show that ADMA may induce renal arterial vasoconstriction in BDL cirrhosis but not in pre-hepatic portal hypertension animals, thereby it suggests the importance of liver dysfunction in endothelial-dependent renal vasoconstriction of cirrhosis. Actually, ADMA is partly catabolized by hepatic DDAH and this enzyme activity was lower in the BDL group. Although of clinical interest, ADMA pathogenetic role in cirrhosis and HRS is still uncertain as its circulating concentration may be extremely low as to produce clinically relevant effects. Overall, its role in human pathology have yielded inconsistent results (Loscalzo et al 2004). Minor comments: 1. Bile duct ligation is the preferred term for this murine model of cirrhosis and PHT. 2. Obstructive jaundice it-self may induce renal dysfunction in humans. Conceivably,



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hyperbilirubinaemia may sensitize arterial vasculature to endogenous vasoconstrictors. Please, discuss.