

PEER-REVIEW REPORT

Name of journal: World Journal of Nephrology

Manuscript NO: 37366

Title: Is serum copeptin a modifiable biomarker in autosomal dominant polycystic kidney disease?

Reviewer's code: 02876269

Reviewer's country: Italy

Science editor: Li-Jun Cui

Date sent for review: 2017-12-18

Date reviewed: 2017-12-20

Review time: 2 Days

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 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 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 checked="" type="checkbox"/> Plagiarism	<input checked="" type="checkbox"/> Minor revision
<input type="checkbox"/> Grade E: Poor		<input 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

In the article "Is serum copeptin a modifiable biomarker in autosomal dominant polycystic kidney disease?" the Authors describe the possible role of copeptin as molecular marker for ADPKD. The topic is very interesting because little is known about the use of biomarkers for the diagnosis/prognosis of ADPKD. Unfortunately, copeptin expression is affected by different environment factors, rendering hard the use of this molecule as efficient biological marker. Copeptin, likely, could be used as a possible marker in patient selection for the treatment with Tolvaptan. There are things that can be improved. Minor Points: - In the legend of Fig. 1 the last part of the sentence is lost. "Figure adapted from...???". - Chapter "Synthesis, function and degradation of copeptin", the table cited in the sentence "As summarised in Table 2, the precursor peptide..." is wrong, therefore must be substituted with table 1. - In the

Chapter “Advantages and disadvantages of copeptin measurement in the laboratory”, Authors describe possible disadvantages for the copeptin measurement in clinical laboratories mainly in terms of costs. In my opinion, this description is incorrect because, as underline the Authors, there are insufficient clinical evidence for the use of copeptin as ADPKD marker. Therefore, the clinical use of copeptin is still too far. - Are there evidences of copeptin plasma level variations in ADPKD patients treated with Tolvaptan? The reduction or the slowed increase in copeptin plasma levels could indicate a “good” response to Tolvaptan treatment.

PEER-REVIEW REPORT

Name of journal: World Journal of Nephrology

Manuscript NO: 37366

Title: Is serum copeptin a modifiable biomarker in autosomal dominant polycystic kidney disease?

Reviewer's code: 03475636

Reviewer's country: United States

Science editor: Li-Jun Cui

Date sent for review: 2018-01-04

Date reviewed: 2018-01-07

Review time: 3 Days

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 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 checked="" type="checkbox"/> No	<input checked="" type="checkbox"/> Minor revision
<input type="checkbox"/> Grade E: Poor		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

This is well written review on serum copeptin and autosomal dominant polycystic kidney disease. However, I have a few suggestions to improve this review. 1. In introduction, please also include for information regarding use of Copeptin measurement in various other clinical indications, e.g., including the diagnosis of diabetes insipidus and the monitoring of sepsis and cardiovascular diseases. 2. Case control study by Corradi et al is an important one, I agree. The reference for citation #28 is not correct; should be Corradi et al, not Zittema et al. Corradi et al. recently demonstrated that glomerular filtration affects copeptin to a greater extent rather than its correlation to AVP [28] 3. Suggest discussing the findings by Nakajima et al. Association of arginine vasopressin surrogate marker urinary copeptin with severity of autosomal dominant polycystic kidney disease (ADPKD). PMID: 25715868



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PEER-REVIEW REPORT

Name of journal: World Journal of Nephrology

Manuscript NO: 37366

Title: Is serum copeptin a modifiable biomarker in autosomal dominant polycystic kidney disease?

Reviewer's code: 02855928

Reviewer's country: Japan

Science editor: Li-Jun Cui

Date sent for review: 2018-01-04

Date reviewed: 2018-01-11

Review time: 6 Days

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 checked="" 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 checked="" 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

Some biomarkers are crucial for the diagnosis of ADPKD and the prediction of prognosis. Expression of copeptin is 'multifactorial' (i.e., some situations have some effects on the result). This point should be mentioned.

PEER-REVIEW REPORT

Name of journal: World Journal of Nephrology

Manuscript NO: 37366

Title: Is serum copeptin a modifiable biomarker in autosomal dominant polycystic kidney disease?

Reviewer's code: 00468214

Reviewer's country: Italy

Science editor: Li-Jun Cui

Date sent for review: 2018-01-04

Date reviewed: 2018-01-15

Review time: 11 Days

CLASSIFICATION	LANGUAGE EVALUATION	SCIENTIFIC MISCONDUCT	CONCLUSION
<input type="checkbox"/> Grade A: Excellent	<input type="checkbox"/> Grade A: Priority publishing	Google 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
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		<input checked="" type="checkbox"/> No	

COMMENTS TO AUTHORS

The paper is well written and the bibliography adequate. It clearly explains uses and future developments of copeptin as possible biomarker in order to evaluate therapy response and to stratify patients with ADPKD (to early individuate those with a higher risk of ESKD). Minor language polishing required.