

# Rapid Technical Review report

EDI0000 Job Code: Manuscript title: XXXX XXXXX Date filled: XX-XX-XXXX

> Note: Add notes in case client instructions were not followed or if you'd like to draw the

client's attention to any aspect of your review.

### **SCIENTIFIC VALIDITY**

Study Design	
1. Is the study approach consistent with research objective?	
○ Yes ● No ○ NA	
Problem identified	Action recommended
The research objective was to investigate the association between abstinence prognosis and various statuses in pathological gamblers. There are several problems with the operational definition of abstinence in this study. Abstinence was defined only with respect to the one month period of assessment, and it is known that gambling addiction is "transient and episodic"	It is recommended that standard definitions of abstinence and relapse from the literature be used and groups not be grouped together. Further attempts to evaluate all the subjects should be made. Major revision and rethinking of the research design is needed
Relevant manuscript text (optional)	Nature of revision
The present study tested the impact of Pq-LPS infection on the expression level	• Essential-Major
of key proteins supporting foam cell formation, namely CD36, LOX-1, ACAT1 and ABCG1.	<ul> <li>Optional-Major</li> <li>Optional-Minor</li> </ul>
○ Yes ● No ○ NA  Problem identified	Action recommended
It is not clear how the outcome was determined—chart review or self-report?  Was the history obtained from the medical chart obtained by someone who was blinded to the abstinence outcome?	The author should describe in detail in the manuscript Introduction why these particular methods were used and then describe in detail what each parameter stands for.
Relevant manuscript text (optional)	Nature of revision
For quantitative outcome measures using repeated measures analysis of	• Essential-Major $\circ$ Essential-Minor
variance (mixed-effects model) for processing	<ul> <li>Optional-Major</li> <li>Optional-Minor</li> </ul>
3. Is the study missing any experimental controls?  • Yes O NO NA	
Problem identified	Action recommended
Positive controls are desirable for in vivo experiments. Since the authors test an anti-inflammatory activity of CSB, a known anti-inflammatory drug (such as rolipram) could have been used.	If the authors have data on positive controls for at least some experiments they could be included.
Relevant manuscript text (optional)	Nature of revision
Reconsider interpretation of the data on changes in uric acid,	<ul> <li>Essential-Major</li> <li>Essential-Minor</li> </ul>
taking into account the differences in renal function changes seen in Groups A and B and Groups C and D.	<ul> <li>Optional-Major</li> <li>Optional-Minor</li> </ul>



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## Reporting Guidelines and Ethics

○ Yes ● No ○ NA	
Problem identified	Action recommended
Using STROBE criteria: Insufficient background information/rationale and description of setting, participants, variables, bias, statistical methods, and reasons for non-participation/handling of loss to follow-up. Questionable interpretation of results.	No mention of study protocol approval by local institution review board OR of patient consent. The authors also do not make it clear whether the study was done prospectively or
Delouant manuscript tout (autional)	Nature of revision
Relevant manuscript text (optional)  It is assumed that for the emotional process related to Aad, it is highly possible	Essential-Major
It is assumed that for the emotional process related to Aad, it is highly possible that the recipient's response in the form of the emotion generated	<ul> <li>Optional-Major</li> <li>Optional-Minor</li> </ul>
Is appropriate research ethics information (consent, ethical a	approval, patient anonymity) provided?
○ Yes ● No ○ NA	
Problem identified	Action recommended
No mention of Informed consent obtained in writing.	No clear mention of informed consent is included.
Relevant manuscript text (optional)	Nature of revision
Written informed consents from all subjects need to be included	Essential-Major     Essential-Minor
Instead of patient's family, mention that the consent was obtained from patients' parents or guardians.	Optional-Major Optional-Minor
Has the sample been adequately described (number, inclusi	on and exclusion criteria, etc.)?
Has the sample been adequately described (number, inclusi  Yes ● No ○ NA	
Has the sample been adequately described (number, inclusi Yes No NA Problem identified	Action recommended
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