Materials Design Analysis Reporting (MDAR) Checklist for Authors

The MDAR framework establishes a minimum set of requirements in transparent reporting applicable to studies in the life sciences (see Statement of Task: doi:10.31222/osf.io/9sm4x.). The MDAR checklist is a tool for authors, editors and others seeking to adopt the MDAR framework for transparent reporting in manuscripts and other outputs. Please refer to the MDAR Elaboration Document for additional context for the MDAR framework.

Materials

Antibodies	Yes (indicate where provided:	n/a
For commercial reagents, provide supplier name, catalogue number and RRID, if available.	Page 5, Line 6 Page 5, Line 10	

Cell materials	Yes (indicate where provided:	n/a
Cell lines: Provide species information, strain. Provide accession number in repository **OR** supplier name, catalog number, clone number, **OR** RRID		Cell lines were not used in this study
Primary cultures: Provide species, strain, sex of origin, genetic modification status.		Primary cultures were not used in this study

Experimental animals	Yes (indicate where provided:	n/a
Laboratory animals: Provide species, strain, sex, age, genetic modification status. Provide accession number in repository **OR** supplier name, catalog number, clone number, **OR** RRID		Laboratory animals were not used in this study
Animal observed in or captured from the field: Provide species, sex and age where possible		Animal observed in or captured from the field were not used in this study
Model organisms: Provide Accession number in repository (where relevant) **OR** RRID		Model organisms were not used in this study

Plants and microbes	Yes (indicate where provided:	n/a
Plants: provide species and strain, unique accession number if available, and source (including location for collected wild specimens)		Plants were not used in this study
Microbes: provide species and strain, unique accession number if available, and source		Microbes were not used in this study

Human research participants	Yes (indicate where provided:	n/a
Identify authority granting ethics approval (IRB or equivalent committee(s), provide reference number for approval.	Page 1, Line 40-44	
Provide statement confirming informed consent obtained from study participants.	Page 1, Line 40-44	
Report on age and sex for all study participants.	Page 7, Line 8	

Design

Study protocol	Yes (indicate where provided:	n/a
For clinical trials, provide the trial registration number **OR** cite DOI in manuscript.		Not a clinical trial

Laboratory protocol	Yes (indicate where provided:	n/a
Provide DOI or other citation details if detailed step-by-step protocols are available.		No detailed step-by-step protocols

Experimental study design (statistics details)	Yes (indicate where provided:	n/a
State whether and how the following have been done**, or** if they were not carried out.	Page 4, Line 21-22 Page 5, Line 1-2 Page 6, Line 3 Page 4, Line 22 Page 5, Line 1	
Sample size determination	Page 4, Line 21-22 Page 5, Line 1-2	
Randomisation	Page 6, Line 3	
Blinding	No blinding studies	No blinding studies
Inclusion/exclusion criteria	Page 4, Line 22 Page 5, Line 1	

Sample definition and in-laboratory replication	Yes (indicate where provided:	n/a
State number of times the experiment was replicated in laboratory	Page 5, Line 13	
Define whether data describe technical or biological replicates	Page 5, Line 13	

Ethics	Yes (indicate where provided:	n/a
Studies involving human participants: State details of authority granting ethics approval (IRB or equivalent committee(s), provide reference number for approval.	Page 1, Line 40-44	E
Studies involving experimental animals: State details of authority granting ethics approval (IRB or equivalent committee(s), provide reference number for approval.		No study involving experimental animals
Studies involving specimen and field samples: State if relevant permits obtained, provide details of authority approving study; if none were required, explain why.	Page 5, Line 2	

Dual Use Research of Concern (DURC)	Yes (indicate where provided:	n/a
If study is subject to dual use research of concern, state the authority granting approval and reference number for the regulatory approval		Not a dual use research

Analysis

Attrition	Yes (indicate where provided:	n/a
State if sample or data point from the analysis is excluded, and whether the criteria for exclusion were determined and specified in advance.		No sample or data point from the analysis is excluded

Statistics	Yes (indicate where provided:	n/a
Describe statistical tests used and justify choice of tests.	Page 6, Line 16-19	Materials and Methods, Paragraph 4

Data Availability	Yes (indicate where provided:	n/a
State whether newly created datasets are available, including protocols for access or restriction on access.		No newly created datasets
If data are publicly available, provide accession number in repository or DOI or URL.		No public data
If publicly available data are reused, provide accession number in repository or DOI or URL, where possible.		No public data

Code Availability	Yes (indicate where provided:	n/a
For all newly generated code and software essential for replicating the main findings of the study:		No newly generated code and software
State whether the code or software is available.		No newly generated code and software
If code is publicly available, provide accession number in repository, or DOI or URL.		No newly generated code and software

Reporting

Adherence to community standards	Yes (indicate where provided: section/paragraph)	n/a
MDAR framework recommends adoption of discipline-specific guidelines, established and endorsed through community initiatives. Journals have their own policy about requiring specific guidelines and recommendations to complement MDAR.		n/a
State if relevant guidelines (eg., ICMJE, MIBBI, ARRIVE) have been followed, and whether a checklist (eg., CONSORT, PRISMA, ARRIVE) is provided with the manuscript.	ICMJE guidelines were followed, as the journal follows ICMJE recommendations for publication.	

Article Information: http://dx.doi.org/10.21037/jgo-20-344