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: section/paragraph)	n/a
For commercial reagents, provide supplier name, catalogue number and RRID, if available.	No antibodies in this study.	**n/a**

Cell materials	Yes (indicate where provided: section/paragraph)	n/a
Cell lines: Provide species information, strain. Provide accession number in repository **OR** supplier name, catalog number, clone number, **OR** RRID	No cell materials in this study.	**n/a**
Primary cultures: Provide species, strain, sex of origin, genetic modification status.	Yes, in materials and methods section.	

Experimental animals	Yes (indicate where provided: section/paragraph)	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	No laboratory animals in this study.	**n/a**
Animal observed in or captured from the field: Provide species, sex and age where possible	No laboratory animals in this study.	**n/a**
Model organisms: Provide Accession number in repository (where relevant) **OR** RRID	No laboratory animals in this study.	**n/a**

Plants and microbes	Yes (indicate where provided: section/paragraph)	n/a
Plants: provide species and strain, unique accession number if available, and source (including location for collected wild specimens)	No plants in this study.	**n/a**
Microbes: provide species and strain, unique accession number if available, and source	No plants in this study.	**n/a**

Human research participants	Yes (indicate where provided: section/paragraph)	n/a
Identify authority granting ethics approval (IRB or equivalent committee(s), provide reference number for approval.	NO, the data comes from the Internet and is anonymous.	**n/a**
Provide statement confirming informed consent obtained from study participants.	NO, the data comes from the Internet and is anonymous.	**n/a**
Report on age and sex for all study participants.	NO, data does not include age and gender.	**n/a**

Design

Study protocol	Yes (indicate where provided: section/paragraph)	n/a
For clinical trials, provide the trial registration number **OR** cite DOI in manuscript.	No, this is not a clinical trial.	**n/a**

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

Experimental study design (statistics details)	Yes (indicate where provided: section/paragraph)	n/a
State whether and how the following have been done**, or** if they were not carried out.		**n/a**
Sample size determination	Not experimental study design	**n/a**
Randomisation	Not experimental study design	**n/a**
Blinding	Not experimental study design	**n/a**
Inclusion/exclusion criteria	Not experimental study design	**n/a**

Sample definition and in-laboratory replication	Yes (indicate where provided: section/paragraph)	n/a
State number of times the experiment was replicated in laboratory	Not applicable	**n/a**
Define whether data describe technical or biological replicates	Not applicable	**n/a**

Ethics	Yes (indicate where provided: section/paragraph)	n/a
Studies involving human participants: State details of authority granting ethics approval (IRB or equivalent committee(s), provide reference number for approval.	The data comes from the Internet and is anonymous.	**n/a**
Studies involving experimental animals: State details of authority granting ethics approval (IRB or equivalent committee(s), provide reference number for approval.	The data comes from the Internet and is anonymous.	**n/a**
Studies involving specimen and field samples: State if relevant permits obtained, provide details of authority approving study; if none were required, explain why.	The data comes from the Internet and is anonymous.	**n/a**

Dual Use Research of Concern (DURC)	Yes (indicate where provided: section/paragraph)	n/a
If study is subject to dual use research of concern, state the authority granting approval and reference number for the regulatory approval	No dual-use research	**n/a**

Analysis

Attrition	Yes (indicate where provided: section/paragraph)	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.	Yes, in materials and methods section.	

Statistics	Yes (indicate where provided: section/paragraph)	n/a
Describe statistical tests used and justify choice of tests.	Statistics are generated by website calculation	

Data Availability	Yes (indicate where provided: section/paragraph)	n/a
State whether newly created datasets are available, including protocols for access or restriction on access.	The data comes from the Internet	
If data are publicly available, provide accession number in repository or DOI or URL.	The data comes from the Internet	
If publicly available data are reused, provide accession number in repository or DOI or URL, where possible.	Yes, in the footnote section we provide access to the data of this study.	

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

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/TCR-20-2309