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	n/a
For commercial reagents, provide supplier name, catalogue number and RRID, if available.		No antibodies

Cell materials	Yes (indicate where	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
Primary cultures: Provide species, strain, sex of origin, genetic modification status.		No Primary cultures

Experimental animals	Yes (indicate where	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 experimental animals
Animal observed in or captured from the field: Provide species, sex and age where possible		No experimental animals
Model organisms: Provide Accession number in repository (where relevant) **OR** RRID		No experimental animals

Plants and microbes	Yes (indicate where	n/a
Plants: provide species and strain, unique accession number if available, and source (including location for collected wild specimens)		No plants and microbes
Microbes: provide species and strain, unique accession number if available, and source		No plants and microbes

Human research participants	Yes (indicate where	n/a
Identify authority granting ethics approval (IRB or equivalent committee(s), provide reference number for approval.		Have no human research participants
Provide statement confirming informed consent obtained from study participants.		Have no human research participants
Report on age and sex for all study participants.		Have no human research

Design

Study protocol	Yes (indicate where provided:	n/a
For clinical trials, provide the trial registration number **OR** cite DOI in manuscript.		No clinical trials

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

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.		No experimental study design
Sample size determination		Not carry out
Randomisation		Not carry out
Blinding		Not carry out
Inclusion/exclusion criteria		Not carry out

Sample definition and in-laboratory replication	Yes (indicate where provided:	n/a
State number of times the experiment was replicated in laboratory		No sample definition and in-laboratory
Define whether data describe technical or biological replicates		No sample definition and in-laboratory

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.	approval from the local ethics committee or written informed consent from the participants before the study was not necessary	
Studies involving experimental animals: State details of authority granting ethics approval (IRB or equivalent committee(s), provide reference number for approval.		No 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.		No specimen and field samples

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		No Dual Use Research of Concern

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 attrition

Statistics	Yes (indicate where provided:	n/a
Describe statistical tests used and justify choice of tests.	chi-square test to analyze	

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, all data has been presented in the article
If data are publicly available, provide accession number in repository or DOI or URL.		All data has been presented in the article
If publicly available data are reused, provide accession number in repository or DOI or URL, where possible.		All data has been presented in the article

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

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.		
State if relevant guidelines (eg., ICMJE, MIBBI, ARRIVE) have been followed, and whether a checklist (eg., CONSORT, PRISMA, ARRIVE) is provided with the manuscript.	SQUIRE guidelines and ICMJE guidelines were followed, as the journal follows ICMJE recommendations for publication.	

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