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	Not applicable.	NA

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	Not applicable.	NA
Primary cultures: Provide species, strain, sex of origin, genetic	Not applicable.	NA

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	Not applicable.	NA
Animal observed in or captured from the field: Provide species, sex and age where possible	Not applicable.	NA
Model organisms: Provide Accession number in repository (where	Not applicable.	NA

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	Not applicable.	NA
Microbes: provide species and strain, unique accession number if available,	Not applicable.	NA

Human research participants	Yes (indicate where provided:	n/a
Identify authority granting ethics approval (IRB or equivalent committee(s), provide reference number for approval.	Materials and methods/paragraph2	
Provide statement confirming informed consent obtained from study participants.	Materials and methods/paragraph2	
Report on age and sex for all study	Materials and methods/paragraph1	

Design

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

Laboratory protocol	Yes (indicate where provided:	n/a
Provide DOI or other citation details if detailed step-by-step protocols are	Not applicable.	NA

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

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

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.	Materials and methods/paragraph2	
Studies involving experimental animals: State details of authority granting ethics approval (IRB or equivalent committee(s), provide reference number for approval.	Not applicable.	NA
Studies involving specimen and field samples: State if relevant permits obtained, provide details of authority approving study; if none were required,	Not applicable.	NA

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	Not applicable.	NA

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	Not applicable.	NA

Statistics	Yes (indicate where provided:	n/a
Describe statistical tests used and justify choice of tests.	Materials and methods/paragraph 6-8	

Data Availability	Yes (indicate where provided:	n/a
State whether newly created datasets are available, including protocols for access or restriction on access.	Not applicable.	NA
If data are publicly available, provide accession number in repository or DOI or	Not applicable.	NA
If publicly available data are reused, provide accession number in repository or DOI or URL, where possible.	Not applicable.	NA

Code Availability	Yes (indicate where provided:	n/a
For all newly generated code and software essential for replicating the main findings		
State whether the code or software is	Not applicable.	na
If code is publicly available, provide accession number in repository, or DOI or	Not applicable.	na

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.	ICMJE guidelines were followed, as the journal follows ICMJE recommendations for publication.	

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