<u>Materials Design Analysis Reporting (MDAR)</u> 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	There are no antibodies used in the study.	n/a
name, catalogue number and RRID, if available.		

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	Yes, it is provided in Materials and Methods/ Cell culture and gene knockdown/Paragraph 6	
Primary cultures: Provide species, strain, sex of origin, genetic modification status.	There is no such experiment in the study.	n/a

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	There is no animal used in the study.	n/a
Animal observed in or captured from the field: Provide species, sex and age where possible	There is no animal used in the study.	n/a
Model organisms: Provide Accession number in repository (where relevant) OR RRID	There is no model organism in the 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)	There is no plant in the study.	n/a
Microbes: provide species and strain, unique accession number if available, and source	There is no microbe used in the study.	n/a

Human research participants	Yes (indicate where provided: section/paragraph)	n/a
Identify authority granting ethics approval (IRB or	There is no human participant who participated in the	n/a
equivalent committee(s), provide reference number	study.	
for approval.		
Provide statement confirming informed consent	There is no human participant who participated in the	n/a
obtained from study participants.	study.	
Report on age and sex for all study participants.	There is no human participant who participated in the	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.	This study 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.	Yes, it is provided in Materials and Methods.	
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.	, , , , , , , , , , , , , , , , , , , ,	
Sample size determination	There is no such experiment method in the study.	n/a
Randomisation	There is no such experiment method in the study.	n/a
Blinding	There is no such experiment method in the study.	n/a
Inclusion/exclusion criteria	There is no such experiment method in the study.	n/a
Sample definition and in-laboratory replication	Yes (indicate where provided: section/paragraph)	n/a
State number of times the experiment was	Yes, it is provided in Materials and	
replicated in laboratory	Methods/Paragraph 10	
Define whether data describe technical or biological	Yes, it is provided in Materials and Methods/	
replicates	Statistical analysis/Paragraph 10	
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.	There is no human participant who participated in the study.	n/a
Studies involving experimental animals: State details of authority granting ethics approval (IRB or equivalent committee(s), provide reference number for approval.	There is no animal used in the study.	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.	There is no specimen and field samples used in the study.	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	The study is not subject to dual use research of concern.	n/a

Analysis

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

Statistics	Yes (indicate where provided: section/paragraph)	n/a
Describe statistical tests used and justify choice of	Yes, it is provided in Materials and Methods/	
tests.	Statistical analysis/Paragraph 10	

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.	There is no such experiment method in the study.	n/a
If data are publicly available, provide accession number in repository or DOI or URL.	There is no such data in the study.	n/a
If publicly available data are reused, provide accession number in repository or DOI or URL, where possible.	Yes, it is provided in Materials and Methods/ Data source collection and processing/Paragraph 1	

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:		
State whether the code or software is available.	There is no such code or software in the study.	n/a
If code is publicly available, provide accession number in repository, or DOI or URL.	There is no such code in the 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.		
State if relevant guidelines (eg., ICMJE, MIBBI,	ICMJE guidelines were followed, as the journal follows	
ARRIVE) have been followed, and whether a checklist	ICMJE recommendations for publication.	
(eg., CONSORT, PRISMA, ARRIVE) is provided with		
the manuscript.		

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