<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	Methods	
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 (Page 3, Cell culture and treatment section)	
Primary cultures: Provide species, strain, sex of origin, genetic modification status.	Yes. (Page 3, Cell culture and treatment 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	Not applicable. No animals were used in the experiment.	
Animal observed in or captured from the field: Provide species, sex and age where possible	Not applicable. No animals were used in the experiment.	
Model organisms: Provide Accession number in repository (where relevant) OR RRID	Not applicable. No animals were used in the experiment.	

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)	Not applicable. No plants or microbes were used in the experiment.	
Microbes: provide species and strain, unique accession number if available, and source	Not applicable. No plants or microbes were used in the experiment.	

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.	Not applicable. This study did not involve humans.	
Provide statement confirming informed consent obtained from study participants.	Not applicable. This study did not involve humans.	
Report on age and sex for all study participants.	Not applicable. This study did not involve humans.	

<u>Design</u>

Study protocol	Yes (indicate where provided: section/paragraph)	n/a
For clinical trials, provide the trial registration number OR cite DOI in manuscript.	Not applicable.	
Laboratory protocol	Yes (indicate where provided: section/paragraph)	n/a
Provide DOI or other citation details if detailed step- by-step protocols are available.	Not applicable.	
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.	Yes (Page7, statistical analysis section)	
Sample size determination	Yes (Page7, statistical analysis section)	
Randomisation		
Blinding		
Inclusion/exclusion criteria		
Sample definition and in-laboratory replication	Yes (indicate where provided: section/paragraph)	n/a
State number of times the experiment was replicated in laboratory	3 times. (Page 6-7, statistical analysis)	
Define whether data describe technical or biological replicates	Technical and biological replicates. (Page 6-7, statistical analysis)	
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.	Not applicable. This study did not involve humans or animals.	iiy a
Studies involving experimental animals: State details of authority granting ethics approval (IRB or equivalent committee(s), provide reference number for approval.	Not applicable. This study did not involve humans or animals.	
Studies involving specimen and field samples: State if relevant permits obtained, provide details of authority approving study; if none were required, explain why.	Not applicable. This study did not involve humans or animals.	
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	Not applicable. This study did not involve humans or animals.	

<u>Analysis</u>

Attrition	Yes (indicate where provided: section/paragraph)	n/a
State if sample or data point from the analysis is	Not applicable.	
excluded, and whether the criteria for exclusion were		
determined and specified in advance.		

Statistics	Yes (indicate where provided: section/paragraph)	n/a	l
Describe statistical tests used and justify choice of	Yes (Page 6-7, statistical analysis)	1	l
tests.		1	l

Data Availability	Yes (indicate where provided: section/paragraph)	n/a
State whether newly created datasets are available,	The datasets analyzed during the current study are	
including protocols for access or restriction on	available from the corresponding author on reasonable	
access.	request.	
If data are publicly available, provide accession	Not applicable.	
number in repository or DOI or URL.		
If publicly available data are reused, provide	Not applicable.	
accession number in repository or DOI or URL, where		
possible.		

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

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.		

Article information: https://dx.doi.org/10.21037/atm-22-1894