<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		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.	Materials and methods/Paragraph 3	
Provide accession number in repository OR supplier name, catalog number, clone number, OR RRID	Page3/Line 89-90 and Page4/Line 91-92	
Primary cultures: Provide species, strain, sex of origin, genetic modification status.	x of Materials and methods/Paragraph 4 and paragraph 5 Page 4/Line 96-108	

Experimental animals	Yes (indicate where provided: section/paragraph)	n/a
Laboratory animals: Provide species, strain, sex, age,		n/a
genetic modification status. Provide accession		
number in repository OR supplier name, catalog		
number, clone number, OR RRID		
Animal observed in or captured from the		n/a
field: Provide species, sex and age where		
possible		
Model organisms: Provide Accession number		n/a
in repository (where relevant) OR RRID		

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)		n/a
Microbes: provide species and strain, unique accession number if available, and source		n/a

Human research participants	Yes (indicate where provided: section/paragraph)	n/a
Identify authority granting ethics approval (IRB or	Materials and methods/Paragraph 1	
equivalent committee(s), provide reference number	Page 3/Line 78-87	
for approval.		
Provide statement confirming informed consent	Materials and methods/Paragraph 1	
obtained from study participants.	Page 3/Line 78-87	
Report on age and sex for all study participants.	Results/Paragraph 1	
	Page 7/Line 188-193	

Design

Study protocol	Yes (indicate where provided: section/paragraph)	n/a
For clinical trials, provide the trial registration		n/a
number OR cite DOI in manuscript.		
Laboratory protocol	Yes (indicate where provided: section/paragraph)	n/a
Provide DOI or other citation details if detailed step-	Materials and methods/Paragraph 3-10	,
by-step protocols are available.	Page 3-6/Line 89-175	
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	Materials and methods/Paragraph 1	
Randomisation	Materials and methods/Paragraph 1,2	
Blinding	Materials and methods/Paragraph 1,2	
Inclusion/exclusion criteria	Materials and methods/Paragraph 1,2	
Sample definition and in-laboratory replication	Yes (indicate where provided: section/paragraph)	n/a
State number of times the experiment was	Materials and methods/Paragraph 6-9	
replicated in laboratory		
Define whether data describe technical or biological	Materials and methods/Paragraph 11-13	
replicates		
Ethics	Yes (indicate where provided: section/paragraph)	n/a
Studies involving human participants: State details of	Materials and methods/Paragraph 1,2	
authority granting ethics approval (IRB or equivalent		
committee(s), provide reference number for		
approval.		
Studies involving experimental animals: State details		n/a
of authority granting ethics approval (IRB or		
equivalent committee(s), provide reference number		
for approval.		
Studies involving specimen and field samples: State if	Materials and methods/Paragraph 1,2	
relevant permits obtained, provide details of	, , , , ,	
authority approving study; if none were required,		
explain why.		
Dual Use Research of Concern (DURC)	Voc (indicate where provided costion/paragraph)	n/-
If study is subject to dual use research of concern,	Yes (indicate where provided: section/paragraph)	n/a
state the authority granting approval and reference		n/a
number for the regulatory approval		
number for the regulatory approval		1

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	Results/Paragraph 1-3	
determined and specified in advance.		

Statistics	Yes (indicate where provided: section/paragraph)	n/a	
Describe statistical tests used and justify choice of	Materials and methods/ Paragraph 13		
tests.			

Data Availability	Yes (indicate where provided: section/paragraph)	n/a
State whether newly created datasets are available,	Materials and methods/ Paragraph 13	
including protocols for access or restriction on		
access.		
If data are publicly available, provide accession	Materials and methods/ Paragraph 13	
number in repository or DOI or URL.		
If publicly available data are reused, provide		n/a
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		
for replicating the main findings of the study:		
State whether the code or software is available.	Materials and methods/ Paragraph 13	
	Results/Paragraph 4-5	
If code is publicly available, provide accession		n/a
number in repository, or DOI or URL.		

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 guidelines for publication.	
(eg., CONSORT, PRISMA, ARRIVE) is provided with		
the manuscript.		

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