

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](https://doi.org/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 name, catalogue number and RRID, if available.	Section of Cell culture, Cell transfection, Dual-luciferase gene system, Transwell Assay, Wound healing analysis, Real-time PCR, Western blot, RIP	
Cell materials	Yes (indicate where provided: section/paragraph)	n/a
Cell lines: Provide species information, strain. Provide accession number in repository OR supplier name, catalogue number, clone number, OR RRID	Section of Cell culture/paragraph 1.	
Primary cultures: Provide species, strain, sex of origin, genetic modification status.	Section of Cell culture/paragraph 1.	
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, catalogue number, clone number, OR RRID	There is no Experimental animal in this study.	n/a
Animal observed in or captured from the field: Provide species, sex and age where possible	There is no Experimental animal in this study.	n/a
Model organisms: Provide Accession number in repository (where relevant) OR RRID	There is no Experimental animal in this 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 are no Plants and microbes in this study.	n/a
Microbes: provide species and strain, unique accession number if available, and source	There are no Plants and microbes in this study.	n/a
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.	There are no Human research participants in this study.	n/a
Provide statement confirming informed consent obtained from study participants.	There are no Human research participants in this study.	n/a
Report on age and sex for all study participants.	There are no Human research participants in this study.	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.	There is no step-by-step protocol.	n/a
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	Not applicable.	n/a
Randomisation	Not applicable.	n/a
Blinding	Not applicable.	n/a
Inclusion/exclusion criteria	Not applicable.	n/a
Sample definition and in-laboratory replication	Yes (indicate where provided: section/paragraph)	n/a
State number of times the experiment was replicated in laboratory	Section of Transwell Assay/paragraph 1, Wound healing analysis/paragraph 1.	
Define whether data describe technical or biological replicates	Section of Transwell Assay/paragraph 1, Wound healing analysis/paragraph 1.	
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 are no Human research participants in this 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 are no experimental animals in this 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 are no specimen and field samples in this 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	This 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 excluded, and whether the criteria for exclusion were determined and specified in advance.	There are no Human research participants or experimental animals in this study.	n/a
Statistics	Yes (indicate where provided: section/paragraph)	n/a
Describe statistical tests used and justify choice of tests.	Section of Statistical analysis/paragraph 1.	
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 are no newly created datasets.	n/a
If data are publicly available, provide accession number in repository or DOI or URL.	There are no newly created datasets.	n/a
If publicly available data are reused, provide accession number in repository or DOI or URL, where possible.	The publicly available data are not reused in this study.	n/a
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.	The software using for statistical analysis is shown in Section Statistical analysis/ paragraph 1.	
If code is publicly available, provide accession number in repository, or DOI or URL.	There is no publicly available code in this 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, 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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