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 RRID, if available.		**n/a**

Cell materials	Yes (indicate where provided: section/paragraph)	**n/a** (No Cell experiments were not involved in this study**)**
Cell lines: Provide species information, strain. Provide accession number in repository **OR** supplier name, catalog number, clone number, **OR** RRID		**n/a**
Primary cultures: Provide species, strain, sex of origin, genetic modification status.		**n/a**

Experimental animals	Yes (indicate where provided: section/paragraph)	n/a (No animal experiments were involved in this study)
Laboratory animals: Provide species, strain, sex, age, genetic modification status. Provide accession number in repository **OR** supplier name, catalog number, clone number, **OR** RRID		n/a
Animal observed in or captured from the field: Provide species, sex and age where possible		**n/a**
Model organisms: Provide Accession number in repository (where relevant) **OR** RRID		**n/a**

Plants and microbes	Yes (indicate where provided: section/paragraph)	**n/a** (No plants or microbes were not involved in this study**)**
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)	
Identify authority granting ethics approval (IRB or equivalent committee(s), provide reference number for approval.	Methods/paragraph2;page5/paragraph24,25	
Provide statement confirming informed consent obtained from study participants.	Footnote/paragraph3;page10/paragraph11-16	
Report on age and sex for all study participants.	Methods/paragraph1;page5/paragraph10-20	

Design

Study protocol	**Yes (indicate where provided: section/paragraph)**	**n/a((**No clinical trials were not involved in this retrospective study**)**
For clinical trials, provide the trial registration number **OR** cite DOI in manuscript.		**n/a**

Laboratory protocol	**Yes (indicate where provided: section/paragraph)**	**n/a((**No laboratory protocol were not involved in this retrospective study**)**
Provide DOI or other citation details if detailed step-by-step protocols are available.		**n/a**

Experimental study design (statistics details)	**Yes (indicate where provided: section/paragraph)**	
State whether and how the following have been done**, or** if they were not carried out.		
Sample size determination		n/a (This is a retrospective cohort study)
Randomisation		n/a (This is a retrospective cohort study)
Blinding		n/a (This is a retrospective cohort study)
Inclusion/exclusion criteria	Methods/paragraph1;page4/paragraph13-18	

Sample definition and in-laboratory replication	**Yes (indicate where provided: section/paragraph)**	n/a(Sample definition or in-laboratory replication were not involved in this)retrospective study
State number of times the experiment was replicated in laboratory		**n/a**
Define whether data describe technical or biological replicates		**n/a**

Ethics	**Yes (indicate where provided:**	
Studies involving human participants: State details of authority granting ethics approval (IRB or equivalent committee(s), provide reference number for approval.	Footnote/paragraph3;page10/paragraph11-16	
Studies involving experimental animals: State details of authority granting ethics approval (IRB or equivalent committee(s), provide reference number for approval.		**n/a(**No experimental animals were not involved in this retrospective study**)**
Studies involving specimen and field samples: State if relevant permits obtained, provide details of authority approving study; if none were required, explain why.		**n/a(**No specimen or field samples were not involved in this retrospective study**)**

Dual Use Research of Concern (DURC)	**Yes (indicate where provided: section/paragraph)**	**n/a(This study did not involve a dual use research)**
If study is subject to dual use research of concern, state the authority granting approval and reference number for the regulatory approval		

Analysis

Attrition	**Yes (indicate where provided: section/paragraph)**	
State if sample or data point from the analysis is excluded, and whether the criteria for exclusion were determined and specified in advance.	Methods/paragraph1;page4/paragraph13-18	

Statistics	**Yes (indicate where provided: section/paragraph)**	
Describe statistical tests used and justify choice of tests.	Methods/paragraph1;page5/paragraph9-20	

Data Availability	**Yes (indicate where provided: section/paragraph)**	
State whether newly created datasets are available, including protocols for access or restriction on access.	Data Sharing Statement	
If data are publicly available, provide accession number in repository or DOI or URL.	Data Sharing Statement	
If publicly available data are reused, provide accession number in repository or DOI or URL, where possible.		**n/a** **(** publicly available data were not used in this study **)**

Code Availability	**Yes (indicate where provided: section/paragraph)**	**n/a** **(**No c**odes** were not involved in this study**)**
For all newly generated code and software essential for replicating the main findings of the study:		
State whether the code or software is available.		**n/a**
If code is publicly available, provide accession number in repository, or DOI or URL.		**n/a**

Reporting

Adherence to community standards	**Yes (indicate where provided: section/paragraph)**	
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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