<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 | No antibodies were used in this study. | |
| 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, | No cell materials were used in this study. | |
| OR RRID | | |
| Primary cultures: Provide species, strain, sex of | No cell materials were used in this study. | |
| origin, genetic modification status. | | |
| 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 | No animals were used in this study. | |
| Animal observed in or captured from the field: Provide species, sex and age where possible | No animals were used in this study. | |
| Model organisms: Provide Accession number in repository (where relevant) OR RRID | No animals were used in this study. | |
| 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) | No plant materials were used in this study. | |
| Microbes: provide species and strain, unique accession number if available, and source | No microbes were used in this study. | |
| Human research participants | Yes (indicate where provided: section/paragraph) | n/a |
| Identify authority granting ethics approval (IRB or | Materials and methods /paragraph 2.1(General | |
| equivalent committee(s), provide reference number for approval. | information) | |
| Provide statement confirming informed consent obtained from study participants. | Materials and methods /paragraph 2.1(General information) | |
| Report on age and sex for all study participants. | Materials and methods /paragraph 2.1(General information) | |

Design

| 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 | |
| | | 1 |
| 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. | | |
| Sample size determination | Not applicable | |
| Randomisation | No randomisation | |
| Blinding | No blinding | |
| Inclusion/exclusion criteria | No exclusion | |
| Comple definition and in laboratory realization | | |
| Sample definition and in-laboratory replication | Yes (indicate where provided: section/paragraph) | n/a |
| State number of times the experiment was replicated in laboratory | Not applicable | |
| Define whether data describe technical or biological replicates | Not applicable | |
| 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. | Materials and methods /paragraph 2.1(General information) | |
| Studies involving experimental animals: State details of authority granting ethics approval (IRB or equivalent committee(s), provide reference number for approval. | Not applicable | |
| 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 | |
| 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 | |

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. | Not applicable | |
| I | | |
| Statistics | Yes (indicate where provided: section/paragraph) | n/a |
| Describe statistical tests used and justify choice of tests. | Materials and methods /paragraph 2.2.2(Quality analysis of sequencing data). | |
| 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. | Not applicable | |
| If data are publicly available, provide accession number in repository or DOI or URL. | Not applicable | |
| If publicly available data are reused, provide accession number in repository or DOI or URL, where possible. | Not applicable | |
| 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: | | 1170 |
| State whether the code or software is available. | Materials and methods /paragraph 2.3(Software), | |
| If code is publicly available, provide accession | Not applicable | |

Reporting

number in repository, or DOI or URL.

| 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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