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: section/paragraph) | n/a |
|------------|-------------------------------------------------|-----|
| For commercial reagents, provide supplier name, catalogue number and RRID, if available. | Methods, Western blotting, Immunofluorescence | |

| 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 | Methods, Cell culture and treatment | |

| Primary cultures: Provide species, strain, sex of origin, genetic modification status. | No primary cultures | n/a |

| 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 experimental animals | n/a |

| Animal observed in or captured from the field: Provide species, sex and age where possible | No experimental animals | n/a |

| Model organisms: Provide Accession number in repository (where relevant) OR RRID | No experimental animals | 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) | No plants | n/a |

| Microbes: provide species and strain, unique accession number if available, and source | No microbes | 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. | Methods, Patients and sampling | |

| Provide statement confirming informed consent obtained from study participants. | Methods, Patients and sampling | |

| Report on age and sex for all study participants. | Relevant information had been published before | 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. | Not clinical trials | 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. | No step-by-step protocols was cited | 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 | Methods, Patients and sampling | |
| Randomisation | Methods, Patients and sampling | |
| Blinding | Methods, Patients and sampling | |
| Inclusion/exclusion criteria | Methods, Patients and sampling | |

| Sample definition and in-laboratory replication | Yes (indicate where provided: section/paragraph) | n/a |
|-------------------------------------------------|-------------------------------------------------|-----|
| State number of times the experiment was replicated in laboratory | Methods, Statistical analysis | |
| Define whether data describe technical or biological replicates | Methods, 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. | Methods, Patients and sampling | |
| Studies involving experimental animals: State details of authority granting ethics approval (IRB or equivalent committee(s), provide reference number for approval. | No experimental animals | 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. | No specimen and field samples | 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 | Not dual use research of concern | n/a |
### Analysis

| Category               | Yes (indicate where provided: section/paragraph) | n/a |
|------------------------|--------------------------------------------------|-----|
| 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. | No sample or data was excluded | 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. | No sample or data was excluded | n/a |
| Statistics             | Yes (indicate where provided: section/paragraph) | n/a |
| Describe statistical tests used and justify choice of tests. | Methods, Statistical analysis | |
| 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. | No newly created datasets | n/a |
| If data are publicly available, provide accession number in repository or DOI or URL. | No accession number | n/a |
| If publicly available data are reused, provide accession number in repository or DOI or URL, where possible. | Results, CLPP is lowly expressed in EOC | |
| 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: | No code or software | n/a |
| State whether the code or software is available. | No code or software | n/a |
| If code is publicly available, provide accession number in repository, or DOI or URL. | No code | 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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