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. | Materials and Methods / Western blotting | |

| Cell lines | 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 | Materials and Methods / Cell culture | |
| Primary cultures: Provide species, strain, sex of origin, genetic modification status. | NO. | The |
## Design

| Study protocol | Yes (indicate where provided: section/paragraph) | n/a |
|----------------|-------------------------------------------------|-----|
| For clinical trials, provide the trial registration number OR cite DOI in manuscript. | | N O. |

| Laboratory protocol | Yes (indicate where provided: section/paragraph) | n/a |
|---------------------|-------------------------------------------------|-----|
| Provide DOI or other citation details if detailed step-by-step protocols are available. | | |

| 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 / Human blood and colon | |
| Randomisation | N | |
| Blinding | N | |
| Inclusion/exclusion criteria | Materials and Methods / Human blood and colon | |

| Sample definition and in-laboratory replication | Yes (indicate where provided: section/paragraph) | n/a |
|-------------------------------------------------|-------------------------------------------------|-----|
| State number of times the experiment was replicated in laboratory | Figure legends/ Fig. 1, 2, 3 | |
| Define whether data describe technical or biological replicates | Figure legends / Fig. 1, 2, 3 | |

| 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. | N O. T h. |
| Studies involving experimental animals: State details of authority granting ethics approval (IRB or equivalent committee(s), provide reference number for approval. | N O. T h. |
| Studies involving specimen and field samples: State if relevant permits obtained, provide details of authority approving study; if none were required, explain why. | Materials and Methods/ Human blood and colon tissues, Footnote/ Ethical Statement |

| 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 | | |
## 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. | Materials and Methods / Human blood and colon tissues |

| Statistics | Yes (indicate where provided: section/paragraph) | n/a |
|------------|--------------------------------------------------|-----|
| Describe statistical tests used and justify choice of tests. | Materials and 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. | Footnote / Data Sharing Statement |
| If data are publicly available, provide accession number in repository or DOI or URL. | Footnote / Data Sharing Statement |
| If publicly available data are reused, provide accession number in repository or DOI or URL, where possible. | Footnote / Data Sharing Statement |

| 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. | N |
| If code is publicly available, provide accession number in repository, or DOI or URL. | N |

## 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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