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.	They were state in the "Methods" when the reagents was mentioned.	

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	The cells in this study were purchased from Fuheng biology, China. Catalog number:H1975: FH0086; HCC827: FH0048; BEAS-2B:FH0319; A549:FH0045; H 1299:FH0908; H520: FH0081; THP -1; FH0112; PC9: FH0083.	
Primary cultures: Provide species, strain, sex of origin, genetic modification status.	There is no primary culture cell used in this study.	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	There is no animal experiment in this study.	n/a
Animal observed in or captured from the field: Provide species, sex and age where possible	There is no animal experiment in this study.	n/a
Model organisms: Provide Accession number in repository (where relevant) **OR** RRID	There is no animal experiment 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)	No plants and microbes were used in this study.	n/a
Microbes: provide species and strain, unique accession number if available, and source	No plants and microbes were used 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.	This is not a human research.	n/a
Provide statement confirming informed consent obtained from study participants.	This is not a human research.	n/a
Report on age and sex for all study participants.	This is not a human research.	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 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 more detailed 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	**It was stated in the 7th section of "Methods".**	
Randomisation	**It was stated in the 7th section of "Methods".**	
Blinding	This is not a clinical trial.	n/a
Inclusion/exclusion criteria	**It was stated in the 7th section of "Methods".**	

Sample definition and in-laboratory replication	Yes (indicate where provided: section/paragraph)	n/a
State number of times the experiment was replicated in laboratory	**It was stated in the 4th, 5th, 6th, 8th, 9th, 10th section of "Methods".**	
Define whether data describe technical or biological replicates	**It was stated in the 4th, 5th, 6th, 8th, 9th, 10th section of "Methods".**	

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.	This is not a human research.	n/a
Studies involving experimental animals: State details of authority granting ethics approval (IRB or equivalent committee(s), provide reference number for approval.	There is no animal experiment 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.	This study was reviewed and approved by the medical ethics committee of Affiliated Cancer Hospital of Guangzhou Medical University, China.	

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 is not a dual use research.	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.	**It was provided in the 7th section of "Methods".**	

Statistics	Yes (indicate where provided: section/paragraph)	n/a
Describe statistical tests used and justify choice of tests.	**It was provided in the 11th section of "Methods".**	

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 dataset was created.	n/a
If data are publicly available, provide accession number in repository or DOI or URL.	There is no data publicly available.	n/a
If publicly available data are reused, provide accession number in repository or DOI or URL, where possible.	There is no data publicly available.	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:	There is no newly generated code or software.	n/a
State whether the code or software is available.	There is no newly generated code or software.	n/a
If code is publicly available, provide accession number in repository, or DOI or URL.	There is no newly generated code or software.	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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