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.	ADAMTS6,Snail1, AKT,ZEB1, VIM, EGFR , TGFβ1 (Abcam, MA) methods/paragraph 11	

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	A2780 and A2780/DDP human ovarian carcinoma cell lines were purchased from the American Type Culture Collection (Shanghai, China) methods/paragraph 2	
Primary cultures: Provide species, strain, sex of origin, genetic modification status.	CD163+TAMs of ovarian cancer ascites methods/paragraph 1	

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
Animal observed in or captured from the field: Provide species, sex and age where possible		no
Model organisms: Provide Accession number in repository (where relevant) **OR** RRID		no

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
Microbes: provide species and strain, unique accession number if available, and source		no

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.	The protocol for this study was approved by the ethics committee of Tongji University. (KS1937) Footnote paragraph 3	
Provide statement confirming informed consent obtained from study participants.	Written informed consent was obtained from all participants. Footnote paragraph 3	
Report on age and sex for all study participants.	Age : 22-82 , sex :female methods/paragraph 1	

Design

Study protocol	Yes (indicate where provided: section/paragraph)	n/a
For clinical trials, provide the trial registration number **OR** cite DOI in manuscript.		no

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

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.		no
Sample size determination		n
Randomisation		n
Blinding		n
Inclusion/exclusion criteria		n

Sample definition and in-laboratory replication	Yes (indicate where provided: section/paragraph)	n/a
State number of times the experiment was replicated in laboratory	3-6	
Define whether data describe technical or biological replicates		no

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.	The protocol for this study was approved by the ethics committee of Tongji University. (KS1937) Footnote paragraph 3	
Studies involving experimental animals: State details of authority granting ethics approval (IRB or equivalent committee(s), provide reference number for approval.		No
Studies involving specimen and field samples: State if relevant permits obtained, provide details of authority approving study; if none were required, explain why.	the protocol for this study was approved by the ethics committee of Tongji University Footnote paragraph 3	

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		no

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.	The enrolled patients had not received chemotherapy, radiotherapy, surgery, immunotherapy, and other treatments, and the patients with other diseases were also excluded. methods/paragraph 1	

Statistics	Yes (indicate where provided: section/paragraph)	n/a
Describe statistical tests used and justify choice of tests.	Statistical analysis was performed using SPSS software (version 21.0; SPSS, Chicago, IL, USA). Data are presented as the mean ± standard error of the mean. The statistical significance of differences between groups was assessed by t-test or one-way analysis of variance with multiple comparisons. Statistical significance was Set at $P < 0.05$. Methods/paragraph 14	

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
If data are publicly available, provide accession number in repository or DOI or URL.		no
If publicly available data are reused, provide accession number in repository or DOI or URL, where possible.		no

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
State whether the code or software is available.		n
If code is publicly available, provide accession number in repository, or DOI or URL.		no

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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