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.	Yes. Methods, paragraph 8/ Immunoblotting and exosome confirmation.	-

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	No cell lines were involved.	NA
Primary cultures: Provide species, strain, sex of origin, genetic modification status.	--	NA

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 lab animals were involved.	NA
Animal observed in or captured from the field: Provide species, sex and age where possible	No lab animals were involved.	NA
Model organisms: Provide Accession number in repository (where relevant) **OR** RRID	No lab animals were involved.	NA

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 involved.	NA
Microbes: provide species and strain, unique accession number if available, and source	No plants and microbes were involved.	NA

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.	Yes, Methods/the 1st paragraph.	--
Provide statement confirming informed consent obtained from study participants.	Yes, Methods/the 1st paragraph.	--
Report on age and sex for all study participants.	Yes, it was showed in Table 1 and Results/paragraph 3.	--

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 study is not a clinical trail.	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.	doi:10.1016/j.eururo.2016.08.012.	--

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.	Yes, Methods/ Statistical analysis/3rd and 4th paragraph	
Sample size determination	Methods/ 1st paragraph	--
Randomisation	Non-randomized study	--
Blinding	Non-blinding study	--
Inclusion/exclusion criteria	Yes, Methods/ 1st and 2nd paragraph	-

Sample definition and in-laboratory replication	Yes (indicate where provided: section/paragraph)	n/a
State number of times the experiment was replicated in laboratory	Yes. Methods/ Urine-derived exosomal RNA isolation and quantitative RT-PCR/1st paragraph	--
Define whether data describe technical or biological replicates	Yes. Methods/ Urine-derived exosomal RNA isolation and quantitative RT-PCR/1st paragraph	--

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.	Yes. Methods/1st paragraph	
Studies involving experimental animals: State details of authority granting ethics approval (IRB or equivalent committee(s), provide reference number for approval.	No experimental animal was involved.	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.	Yes. Methods/1st paragraph	--

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		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.	Yes. Results/2nd paragraph	--

Statistics	Yes (indicate where provided: section/paragraph)	n/a
Describe statistical tests used and justify choice of tests.	Yes. 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.	The datasets used and/or analyzed during the current study are available from the corresponding author on reasonable request.	--
If data are publicly available, provide accession number in repository or DOI or URL.	The datasets used and/or analyzed during the current study are available from the corresponding author on	--
If publicly available data are reused, provide accession number in repository or DOI or URL, where possible.	The datasets used and/or analyzed during the current study are available from the corresponding author on reasonable request.	--

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:	None	N A
State whether the code or software is available.	None	N
If code is publicly available, provide accession number in repository, or DOI or URL.	None	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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