

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](https://doi.org/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.

For all that apply, please note where in the manuscript the required information is provided.

Materials:

Newly created materials	indicate where provided: page no/section/legend)	
The manuscript includes a dedicated "materials availability statement" providing transparent disclosure about availability of newly created materials including details on how materials can be accessed and describing any restrictions on access.	Page 17/Data and Materials availability section	
Antibodies	indicate where provided: page no/section/legend)	n/a
For commercial reagents, provide supplier name, catalogue number and RRID , if available.		n/a
DNA and RNA sequences	indicate where provided: page no/section/legend)	
Short novel DNA or RNA including primers, probes: Sequences should be included or deposited in a public repository.	Page 32/Cloning section	
Cell materials	indicate where provided: page no/section/legend)	n/a
Cell lines: Provide species information, strain. Provide accession number in repository OR supplier name, catalog number, clone number, OR RRID.		n/a
Primary cultures: Provide species, strain, sex of origin, genetic modification status.		n/a
Experimental animals	indicate where provided: page no/section/legend)	n/a
Laboratory animals or Model organisms: Provide species, strain, sex, age, genetic modification status. Provide accession number in repository OR supplier name, catalog number, clone number, OR RRID.		n/a
Animal observed in or captured from the field: Provide species, sex, and age where possible.		n/a
Plants and microbes	indicate where provided: page no/section/legend)	
Plants: provide species and strain, ecotype and cultivar where relevant, unique accession number if available, and source (including location for collected wild specimens).	Pages 35-36/Production and purification of v-cADPR and v2-cADPR section Page 39-40/Phytobacterial challenges, ROS assay, In vivo chemiluminescence imaging, Chlorophyll fluorescence, Production of 2',3'-cAMP/cGMP by HopAM1 sections	
Microbes: provide species and strain, unique accession number if available, and source.	Page 32-33/Cloning, Site directed mutagenesis and Protein expression sections	
Human research participants	indicate where provided: page no/section/legend) or state if these demographics were not collected	n/a
If collected and within the bounds of privacy constraints report on age, sex and gender or ethnicity for all study participants.		n/a

Design:

Study protocol	indicate where provided: page no/section/legend)	n/a
If study protocol has been pre-registered, provide DOI. For clinical trials, provide the trial registration number OR cite DOI.		n/a

Laboratory protocol	indicate where provided: page no/section/legend)	n/a
Provide DOI OR other citation details if detailed step-by-step protocols are available.		n/a

Experimental study design (statistics details)		
For in vivo studies: State whether and how the following have been done	indicate where provided: page no/section/legend. If it could have been done, but was not, write not done	
Sample size determination		n/a
Randomisation	X-ray crystal structures were refined with a randomly selected R-free reflection set based on automatic selection in Phenix 1.19. See pages 37-38	
Blinding		n/a
Inclusion/exclusion criteria		n/a

Sample definition and in-laboratory replication	indicate where provided: page no/section/legend	
State number of times the experiment was replicated in laboratory.	Page 24/ Fig. 3 legend Page 30/ Fig. 6 legend Page 42/ Fig. S1 legend Page 46/ Fig. S3 legend Page 57/ Fig. S8 legend. Page 60/ Fig. S9 legend	
Define whether data describe technical or biological replicates.	Fig. 3, 6, S8, S9: biological Fig. S1, S3: technical	

Ethics	indicate where provided: page no/section/legend	n/a
Studies involving human participants: State details of authority granting ethics approval (IRB or equivalent committee(s), provide reference number for approval.		n/a
Studies involving experimental animals: State details of authority granting ethics approval (IRB or equivalent committee(s), provide reference number for approval.		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.		n/a

Dual Use Research of Concern (DURC)	indicate where provided: page no/section/legend	n/a
If study is subject to dual use research of concern regulations, state the authority granting approval and reference number for the regulatory approval.		n/a

Analysis:

Attrition	indicate where provided: page no/section/legend	n/a
Describe whether exclusion criteria were preestablished. Report if sample or data points were omitted from analysis. If yes report if this was due to attrition or intentional exclusion and provide justification.		n/a

Statistics	indicate where provided: page no/section/legend	n/a
Describe statistical tests used and justify choice of tests.		n/a

Data availability	indicate where provided: page no/section/legend	n/a
For newly created and reused datasets, the manuscript includes a data availability statement that provides details for access or notes restrictions on access.	Page 17/Data and Materials Availability section	
If newly created datasets are publicly available, provide accession number in repository OR DOI OR URL and licensing details where available.	Protein Data Bank, Electron Microscopy Data Bank Page 74-75/ Table S4	
If reused data is publicly available provide accession number in repository OR DOI OR URL, OR citation.	PDB coordinates: Page 4/ Self-association and ligand-free structures of bacterial TIR domains Page 7/ 3'cADPR (v2-cADPR) binds to the ThsA SLOG domain, 3'cADPR (v2-cADPR) changes ThsA tetramer organization Page 22/ Fig.2 legend. Page 38/ Crystallographic data collection. Page 48/ Fig. S4 legend. Page 60/ Fig. S9 legend. Page 66/ Fig. S12 legend.	

Code availability	indicate where provided: page no/section/legend	n/a
For all newly generated custom computer code/software/mathematical algorithm or re-used code essential for replicating the main findings of the study, the manuscript includes a data availability statement that provides details for access or notes restrictions.		n/a
If newly generated code is publicly available, provide accession number in repository, OR DOI OR URL and licensing details where available. State any restrictions on code availability or accessibility.		n/a
If reused code is publicly available provide accession number in repository OR DOI OR URL, OR citation.		n/a

Reporting

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.

Adherence to community standards	indicate where provided: page no/section/legend	n/a
State if relevant guidelines (e.g., ICMJE, MIBBI, ARRIVE) have been followed, and whether a checklist (e.g., CONSORT, PRISMA, ARRIVE) is provided with the manuscript.		n/a