

The TIDieR (Template for Intervention Description and Replication) Checklist*:

Information to include when describing an intervention and the location of the information

Item	Item	Where located **	
number		Primary paper	Other † (details)
		(page or appendix	
		number)	
	BRIEF NAME		
1.	Provide the name or a phrase that describes the intervention.	Page 8	
	WHY		
2.	Describe any rationale, theory, or goal of the elements essential to the intervention.	<u>Page 11</u>	
	WHAT		
3.	Materials: Describe any physical or informational materials used in the intervention, including those	Page 11-13,	
	provided to participants or used in intervention delivery or in training of intervention providers.	<u>Supplementary</u>	
	Provide information on where the materials can be accessed (e.g. online appendix, URL).	Page 3-6	
4.	Procedures: Describe each of the procedures, activities, and/or processes used in the intervention,	Supplementary	
	including any enabling or support activities.	page 23-25	
		(Table S1)	
	WHO PROVIDED		
5.	For each category of intervention provider (e.g. psychologist, nursing assistant), describe their	<u>Supplementary</u>	
	expertise, background and any specific training given.	page 1-2, 23-25	
		(Table S1)	
	HOW		
6.	Describe the modes of delivery (e.g. face-to-face or by some other mechanism, such as internet or	<u>Supplementary</u>	
	telephone) of the intervention and whether it was provided individually or in a group.	page 3-4, 26-34	
		(Table S2)	

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WHERE		
Describe the type(s) of location(s) where the intervention occurred, including any necessary	Supplementary	
infrastructure or relevant features.	page 26-34	
	(Table S2)	
WHEN and HOW MUCH		
Describe the number of times the intervention was delivered and over what period of time including	Supplementary	
the number of sessions, their schedule, and their duration, intensity or dose.	page 23-34	
	(Table S1-S2)	
TAILORING		
If the intervention was planned to be personalised, titrated or adapted, then describe what, why,	<u>Supplementary</u>	
when, and how.	page 3-6, Figure	
	<u>S1-S5</u>	
MODIFICATIONS		
If the intervention was modified during the course of the study, describe the changes (what, why,	N/A. This will be	
when, and how).	described after	
	the study is	
	<u>completed</u>	
HOW WELL		
Planned: If intervention adherence or fidelity was assessed, describe how and by whom, and if any	Supplementary	
strategies were used to maintain or improve fidelity, describe them.	page 6, page 33	
	(Table S2)	
Actual: If intervention adherence or fidelity was assessed, describe the extent to which the	N/A. This will	
intervention was delivered as planned.	be described	
	after the study is	
	completed	
	Describe the type(s) of location(s) where the intervention occurred, including any necessary infrastructure or relevant features. WHEN and HOW MUCH Describe the number of times the intervention was delivered and over what period of time including the number of sessions, their schedule, and their duration, intensity or dose. TAILORING If the intervention was planned to be personalised, titrated or adapted, then describe what, why, when, and how. MODIFICATIONS If the intervention was modified during the course of the study, describe the changes (what, why, when, and how). HOW WELL Planned: If intervention adherence or fidelity was assessed, describe how and by whom, and if any strategies were used to maintain or improve fidelity, describe them. Actual: If intervention adherence or fidelity was assessed, describe the extent to which the	Describe the type(s) of location(s) where the intervention occurred, including any necessary infrastructure or relevant features. WHEN and HOW MUCH Describe the number of times the intervention was delivered and over what period of time including the number of sessions, their schedule, and their duration, intensity or dose. TAILORING If the intervention was planned to be personalised, titrated or adapted, then describe what, why, when, and how. Supplementary page 23-34 (Table S1-S2) MODIFICATIONS If the intervention was modified during the course of the study, describe the changes (what, why, when, and how). MIA: This will be described after the study is completed HOW WELL Planned: If intervention adherence or fidelity was assessed, describe how and by whom, and if any strategies were used to maintain or improve fidelity, describe the extent to which the intervention adherence or fidelity was assessed, describe the extent to which the intervention adherence or fidelity was assessed, describe the extent to which the intervention adherence or fidelity was assessed, describe the extent to which the intervention adherence or fidelity was assessed, describe the extent to which the intervention adherence or fidelity was assessed, describe the extent to which the intervention adherence or fidelity was assessed, describe the extent to which the intervention adherence or fidelity was assessed, describe the extent to which the intervention adherence or fidelity was assessed, describe the extent to which the intervention was delivered as planned.

TIDieR checklist

- ** **Authors** use N/A if an item is not applicable for the intervention being described. **Reviewers** use '?' if information about the element is not reported/not sufficiently reported.
- † If the information is not provided in the primary paper, give details of where this information is available. This may include locations such as a published protocol or other published papers (provide citation details) or a website (provide the URL).
- ‡ If completing the TIDieR checklist for a protocol, these items are not relevant to the protocol and cannot be described until the study is complete.
- * We strongly recommend using this checklist in conjunction with the TIDieR guide (see BMJ 2014;348:g1687) which contains an explanation and elaboration for each item.
- * The focus of TIDieR is on reporting details of the intervention elements (and where relevant, comparison elements) of a study. Other elements and methodological features of studies are covered by other reporting statements and checklists and have not been duplicated as part of the TIDieR checklist. When a randomised trial is being reported, the TIDieR checklist should be used in conjunction with the CONSORT statement (see www.consort-statement.org) as an extension of tem 5 of the CONSORT 2010 Statement. When a clinical trial protocol is being reported, the TIDieR checklist should be used in conjunction with the SPIRIT statement as an extension of tem 11 of the SPIRIT 2013
 Statement (see www.spirit-statement.org). For alternate study designs, TIDieR can be used in conjunction with the appropriate checklist for that study design (see www.spirit-statement.org). For alternate study designs, TIDieR can be used in conjunction with the appropriate checklist for that study design (see www.spirit-statement.org).

TIDieR checklist