

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

Item	Item: Optimized menu	Where lo	ocated **
number		Primary paper	Other † (details)
		(page or appendix	
		number)	
	BRIEF NAME		
1.	Provide the name or a phrase that describes the intervention.	pp.108-109	
	WHY		
2.	Describe any rationale, theory, or goal of the elements essential to the intervention.	pp.118-122	
	WHAT		
3.	Materials: Describe any physical or informational materials used in the intervention, including those		
	provided to participants or used in intervention delivery or in training of intervention providers.	N/A	
	Provide information on where the materials can be accessed (e.g. online appendix, URL).		
4.	Procedures: Describe each of the procedures, activities, and/or processes used in the intervention,	pp.149-214	
	including any enabling or support activities.		
	WHO PROVIDED		
5.	For each category of intervention provider (e.g. psychologist, nursing assistant), describe their	pp. 219-220	
	expertise, background and any specific training given.		
	HOW		
6.	Describe the modes of delivery (e.g. face-to-face or by some other mechanism, such as internet or	pp. 218-219	
	telephone) of the intervention and whether it was provided individually or in a group.		
	WHERE		

7.	Describe the type(s) of location(s) where the intervention occurred, including any necessary	pp. 218-219	
	infrastructure or relevant features.		
	WHEN and HOW MUCH		
8.	Describe the number of times the intervention was delivered and over what period of time including	pp. 113-116	
	the number of sessions, their schedule, and their duration, intensity or dose.		
	TAILORING		
9.	If the intervention was planned to be personalised, titrated or adapted, then describe what, why,	pp. 177-179	
	when, and how.	pp. 202-205	
	MODIFICATIONS		
10.‡	If the intervention was modified during the course of the study, describe the changes (what, why,	pp. 215-218	
	when, and how).		
	HOW WELL		
11.	Planned: If intervention adherence or fidelity was assessed, describe how and by whom, and if any	pp. 244-253	
	strategies were used to maintain or improve fidelity, describe them.		
12. [‡]	Actual: If intervention adherence or fidelity was assessed, describe the extent to which the	pp. 322-327	
	intervention was delivered as planned.		

^{**} **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 **Item 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 **Item 11 of the SPIRIT 2013**TIDieR checklist

