

Reporting Checklists for “COVID-19 Vaccine Implementation in Three Sites in Saskatchewan: Protocol for a Patient-Oriented Realist Evaluation”

1. GRIPP2-SF Reporting Checklist
2. The TIDieR (Intervention Description and Replication) Checklist

From: GRIPP2-SF Reporting Checklist: tools to improve reporting of patient and public involvement in research:

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Section and Topic	Item	Page number
1. Aims	Report the aim of PPI in the study	4
2. Methods	Provide a clear description of methods used for PPI in the study	5-12
3. Study Results	Outcomes: Report the results of PPI in the study, including both positive and negative outcomes	N/A as it is a protocol paper
4. Discussion and conclusions	Outcomes: Comment on the extent to which PPI influenced the study overall. Describe positive and negative effects	11 (As it is a protocol paper, page 11 describes patient engagement for this phase of the study)
5. Critical perspective	Comment critically on the PPI in the study, reflecting on the things that went well and those that did not, so others can learn from this experience	N/A as it is a protocol study

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Item number	Item	Where located **	
		Primary paper (page or appendix number)	Other † (details)
1.	<p>BRIEF NAME Provide the name or a phrase that describes the intervention.</p>	___ 1 ___	_____
2.	<p>WHY Describe any rationale, theory, or goal of the elements essential to the intervention.</p>	___ 4, 5 ___	_____
3.	<p>WHAT 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. Provide information on where the materials can be accessed (e.g. online appendix, URL).</p>	___ 5-12 ___	_____
4.	<p>Procedures: Describe each of the procedures, activities, and/or processes used in the intervention, including any enabling or support activities.</p>	___ 5-12 ___	_____
5.	<p>WHO PROVIDED For each category of intervention provider (e.g. psychologist, nursing assistant), describe their expertise, background and any specific training given.</p>	___ 9 ___	_____

HOW		
6.	Describe the modes of delivery (e.g. face-to-face or by some other mechanism, such as internet or telephone) of the intervention and whether it was provided individually or in a group.	___9___
WHERE		
7.	Describe the type(s) of location(s) where the intervention occurred, including any necessary infrastructure or relevant features.	___5,6___ —
WHEN and HOW MUCH		
8.	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.	___8,9, Figure 1___
TAILORING		
9.	If the intervention was planned to be personalised, titrated or adapted, then describe what, why, when, and how.	___N/A___ —
MODIFICATIONS		
10.‡	If the intervention was modified during the course of the study, describe the changes (what, why, when, and how).	___10,11___
HOW WELL		
11.	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.	___N/A___ —
12.‡	Actual: If intervention adherence or fidelity was assessed, describe the extent to which the intervention was delivered as planned.	___N/A___ —