Framework for Consumer and Community Involvement (CCI) in Health Research

by CCI in Research Alliance

LAUNCH WEBINAR
17 OCTOBER 2025





Acknowledgement of Country

Health Translation Queensland acknowledges the Traditional Owners and their custodianship of the lands on which we meet.

We pay our respects to their Ancestors and their descendants, who continue cultural and spiritual connections to Country.

We recognise their valuable contributions to Australian and global society.

Recognition of lived experience

We are grateful to those who share their lived-experiences, stories and health journeys with us.

We recognise that meaningful involvement of consumers and community members adds significant value to our research in many ways, and at all stages of the research cycle.

Working together strengthens the quality of our research findings, enhances its translation into practice, and ensures our research is more impactful for the wider community



CCI in Research Alliance



Children's Health Queensland Metro North Health

Metro South Health

West Moreton Health

Gold Coast Health























Today's presenters



Dr Nancy Cloake QIMR Berghofer



Benedette Watson
Translational Research Institute



Dr Zephanie TyackQueensland University of Technology



Anja Christoffersen
Award-winning disability advocate,
social entrepreneur and consumer
co-researcher

How did we get here?

- ➤ 6 March 2023, Meeting 1, HTQ CCI in Research Alliance
 - A group of consumers, researchers and consumer engagement specialists form the Framework Project Working Group



First meeting of the HTQ CCI in Research Alliance, 6 March 2023

Framework for CCI in Health Research

- > Identified a gap in advice vs. resources:
 - Researchers should involve consumers at every stage of the research cycle
 - But resources guiding how to involve consumers meaningfully at each stage of the research cycle were missing
- > A streamlined practical tool is needed!

Co-design and Consultation

25 Aug 29 Nov **Early 2024** Late 2024 -31 Jul 2025 2025 2023 17 Oct 2025 **Early 2025** Alliance Group Alliance Framework members Draft shared Final version Workshop endorses Workshop# review first for public^ reviewing launched updated feedback draft consultation First draft version

- #Framework Project Workshop guided by an independent IAP2* facilitator
- ^145 feedback submissions received from consumers, researchers, healthcare staff, clinicians

Importance of the Framework

- Elevating consumer and community voices in research
- > A framework for all, to aid in power balance
- Providing consistency, coordination and safety – breaking down silos and setting expectations

Framework for Consumer and Community Involvement in Health Research

Why is consumer and community involvement (CCI) important?1

Effective CCI ensures that research is responsive to national consumer and community priorities. It reflects the growing expectations of the public, funding bodies and ethics committees for meaningful consumer involvement in health and medical research. Effective involvement of consumers and communities improves public trust in research and facilitates the translation of research findings into practice. The benefits of such involvement are clear. It strengthens the research process, makes it more responsive to the needs of the community, and ultimately leads to more effective and relevant health outcomes.

What is the purpose of this Framework?

To guide those conducting health research on how and when to involve consumers and community in their research based on good practice and the stages of the research cycle.

What does the Framework include?

- Principles for how to involve consumers and community in health research
- Guidance to embed meaningful involvement at each stage of the research cycle
- · Links to resources

Who should use this Framework?

Researchers

- · Clinician researchers
- Consumer researchers
- · Research institutions



¹ Taken from NHMRC and CHF Draft Statement and Consumer and Community Involvement in Health and Medical Research
Figure 1: The stages of the research cycle used for this Framework have been adapted from NHMRC's Keeping Research on Track II (2018).

We acknowledge the Traditional Owners and their custodianship of the lands on which we operate.

We pay our respects to their Ancestors and their descendants, who continue cultural and spiritual connections to Country. We recognise their valuable contributions to Australian and global society and the need to involve First Nations peoples in research to address health equity.

Who has developed the Framework?

This Framework is an initiative of the <u>Health Translation Queensland (HTQ) led Consumer and Community Involvement (CCI) in Research Alliance</u>. The group brings together consumers and CCI leads from across the HTQ partnership and beyond to deliver activities that support good practice CCI in health research in Queensland. The <u>Framework</u> and complementing <u>resources</u> are available on the HTQ website. For enquiries, please contact consumers@healthtranslationqld.org.au.

Consumer and Community Involvement in Research Alliance members:



Children's Health Oueensland

Metro North Health

Metro South Health West Moreton Health























Principles for consumer and community involvement in health research



Authentic relationships and partnerships

- Be respectful, transparent and honest in your communication and decision-making.
- Listen and seek to understand the views of people with different values and beliefs.
- Show empathy, humility and curiosity to build trust and create an openness to new insights and ideas.
- Set clear expectations about the intent, purpose and parameters for involvement.
- Prioritise regular opportunities for open two-way communication and create feedback loops.
- Enable collaborative spaces for communication and working together.
- Prioritise power sharing and promote consumer leadership.
- Prepare and plan to enable successful partnerships that are mutually beneficial.



Inclusive of diverse perspectives

- Identify communities who are impacted by health issues for research.
- Be approachable to enable communities to self-identify (i.e. come to you with a research idea).
- · Reduce barriers to involvement:
 - Be flexible by offering more than one option for getting involved. Ask "What assistance can we provide to support your involvement?"
 - Language can empower or disempower. Using inclusive and non-judgemental language empowers.
 - When communicating, be mindful of unconscious bias and potential for stigma.
 If in doubt, ask.
 - Involve or seek guidance from formal or informal community networks.



Valuing contributions

- Value people's time. Only involve people where there is a clear prospect for them to influence the research process and outcomes.
- Recognise and acknowledge contributions by:
 - o supporting learning and growth
 - reimbursing reasonable out-ofpocket expenses
 - appropriately remunerating consumers for their contribution
 - providing a personalised letter of thanks
 - celebrating milestones together.



Trauma informed approach

- "Check in" and review involvement with consumers at regular intervals during the research journey.
- Recognise that telling or re-telling traumatic experiences can cause distress.
- Enable a supportive environment that enables physical, psychological and cultural safety.
- Offer pre-briefing, debriefing and peer support or external counselling services, such as an employee assistance program.



Consumer and community perspectives on involvement in health research

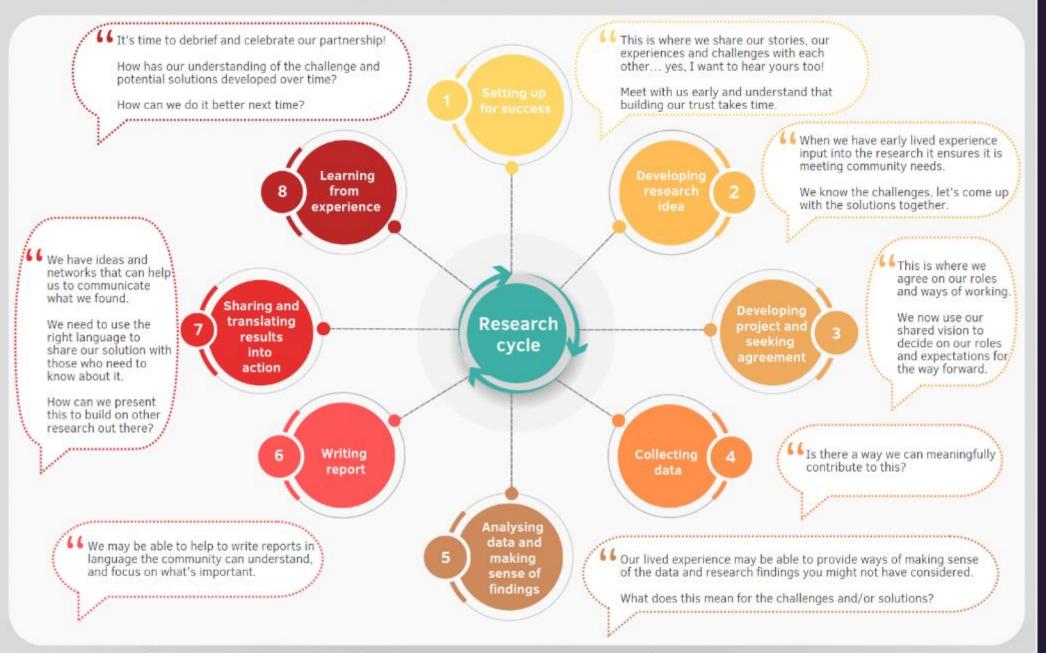


Figure 1: The stages of the research cycle used for this Framework have been adapted from NHMRC's Keeping Research on Track II (2018)

Consumer and community involvement across the research cycle

Stages of the research cycle ¹	1 Setting up for success	2 Developing research idea	3 Developing project and seeking agreement	4 Collecting data	
Consumer and community involvement (CCI) goals	 Find consumers and community representatives. Set expectations. Gather and understand consumers' and community needs 	 Establish a clear research project that will provide value to consumers and community and all other stakeholders. 	Identify how data will be collected through consumer and community involvement. Ensure the research process is accessible and inclusive.	 Gather data using appropriate methods. Have clear rules for the collection of personal data and meeting legislative requirements. Ensure a diverse sample is collected. 	
What you can ask consumers and community ²	Why do you want to be involved? How would you prefer to be involved? What supports do you need?	 What health needs are unmet and could be assisted by research? Is this the right question we need to be researching? 	Is anything missing? Are there any barriers to consumers and community being involved in this process? Are the tools we are using able to be understood?	 How do we build trust and reach the right people? How do we support consumers who are involved in data collection? 	
Examples of involvement activities ³	 Face-to-face conversations Casual group discussions Focus groups Meetings Advisory Group Steering Committee 	Citizen science Workshops Roundtables Public forums Scientific conferences Cultural/community events	 Art spaces Games/interactive play Surveys/online forms Infographics/animations Social media/media case study 	 Journal publications Conference presentations Framework analysis Debriefing sessions Reflections/reviews 	
Building relationships	 Get to know each other - "How are you?" - "What are your interests and experiences?" Share vulnerability and understanding of each other's lived experience. Establish trust with consumers and community. Work together for mutual benefit. Establish regular, two-way communication. 				

- 1 The research cycle stages used in this Framework have been taken from the NHMRC's Keeping Research on Track II (2018)
- 2 These are examples and are in no way an exhaustive list of possible options.
- 3 Consider conducting involvement methods in different attendance formats to be more inclusive for consumers and community (i.e. in-person, online or hybrid, over the phone)

Consumer and community involvement across the research cycle - continued

Stages of the research cycle ¹	5 Analysing data and making sense of findings	6 Writing report	7Sharing & translating results into action	8 Learning from experience	
Consumer and community involvement (CCI) goals	 Identify research limitations and data gaps associated with reach and access of consumers. 	 Prepare consumer-facing report that is clear and easy-to-understand outlining the key findings. Prepare a separate more detailed report for researcher stakeholders. 	Share CCI outcomes and results to integrate into existing or new research projects and/or health processes.	 Evaluate the research process and identify key learnings and improvements for CCI. Ensure consumers are acknowledged for their contributions. 	
What you can ask consumers and community ²	Is there missing community representation? Are consumer voices reflected? What does this research mean to consumers and community? Is there anything from consumers' perspectives that hasn't been considered?	 What is the best way to present the findings so that community can understand them? Have we acknowledged the contribution of consumers? 	How can we share this with community in a way that would encourage behaviour change or implementation of findings? What is the best way to get clinicians' attention?	 Do the findings identify a new research need? How could we apply the results? Can consumers and community be involved differently? 	
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Building relationships	Continue to: get to know each other and communicate regularly share vulnerability and understanding of each other's lived experience build trust and understanding work together for mutual benefit.				

GLOSSARY

Community

A group of people sharing a common interest (e.g. cultural, social, political, health, economic interests) which may or may not include a particular geographic association. Different communities are likely to have different perspectives and approaches to their involvement in research.

Community representative

A person representing a community of interest. In a health context this can include people with shared experiences of a health condition.

Consumer

Patients and potential patients, carers and people who use health care services. Collectively, 'consumers' and 'community representatives' may be referred to as 'the public'.

Diverse/Diversity

Taking into account representation of people with different identities.

Framework

A broad system that outlines principles and structure to guide an approach.

Health and medical research

Research with a human health focus.

Involvement

When consumers and community representatives actively work with researchers and research institutions to help shape decisions about health research priorities, policy, and practice.¹

Learning and growth

Opportunities for capacity and capability building, education, training or access to conference attendance.

Lived experience

Knowledge based on someone's perspective, personal identities, and history, beyond their professional or educational experience.²

Reimbursement

The act of compensating someone for an out-of-pocket expense by giving them an amount of money equal to what was spent (Merriam-Webster Dictionary).

Remuneration

Payment provided in return for work that has been done or services that have been provided (Cambridge Dictionary).

Research

The concept of research is broad and includes the creation of new knowledge and/or the use of existing knowledge in a new and creative way so as to generate new concepts, methodologies, inventions and understandings. This could include synthesis and analysis of previous research to the extent that it is new and creative.⁴

Researcher

Person (or persons) who conducts, or assists with the conduct of, research.3

Research institution

Includes universities, independent research institutes, hospitals or any other organisation that conducts research. It may refer to one or multiple institutions.

Stages of research

The steps that a research team undertakes when conducting research.4

Trauma-informed

Understanding, recognising, and responding appropriately to signs of trauma, and seeking to prevent re-traumatisation.¹

Unconscious bias

Unconscious favouritism towards or prejudice against people of a particular race, gender or group that influences one's actions or perceptions (Oxford English Dictionary).

¹ Definition adapted from Involve Australia - Guidelines for Community Involvement in Genomic Research

² Definition from Health and Human Services - What is lived experience?

³ Definition from US Department and the Australian Code for the Responsible Conduct of Research (2018)

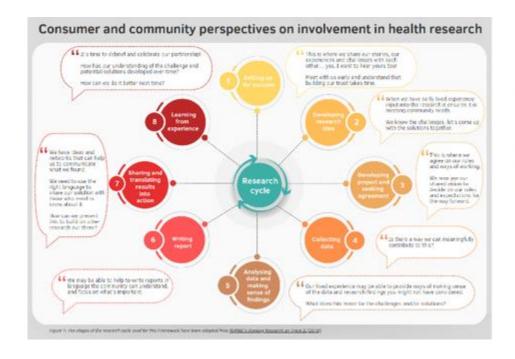
⁴ NHMRC's Keeping Research on Track II (2018)

How to use it:

To meet consumers & community where they are at









Ember to Empower is an MRFF Funded project (GNT#2040537) led by the Australian Centre for Health Services Innovation at the Queensland University of Technology, in collaboration with Monash University, Fiona Wood Foundation, Children's Health QueesnsInd Hospital and Health Service, The University of Notre Dame, K.I.D.S Foundation, and the Australia New Zealand Burns Association.

How to use it:

To think about the big picture

- Being trauma-informed!
- Prioritising open communication
- Reflecting on what worked well and what didn't



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HTQ CCI Microgrants now open!

- Established to fund CCI for design and priority setting of research projects led by a HTQ partner organisations
- ➤ Paused for evaluation in 2024, now back by popular demand bigger and bolder!
- Key improvements:
 - higher amount for each microgrant
 - regular monthly funding cycle from Feb to Nov
 - more streamlined process
- Applicants who apply before the end of 2025 and commit to using and reviewing the new **CCI Framework** can attract additional funds of \$1,000!



Thank you!

For more information:

Framework www.healthtranslationqld.org.au/cci-framework or scan the QR code

Scan here for more details

Microgrants www.healthtranslationqld.org.au/cci-microgrants



Email your questions to consumers@healthtranslationqld.org.au