The "How Tos":

Incorporating Patient Partners in the Dissemination/Implementation of Findings from Clinical Pain Research

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INITIATIVE ON METHODS, MEASUREMENT, AND PAIN ASSESSMENT IN CLINICAL TRIALS (IMMPACT) – XXV

Patient Engagement in Planning, Conduct & Implementation/Dissemination of Clinical Pain Research





















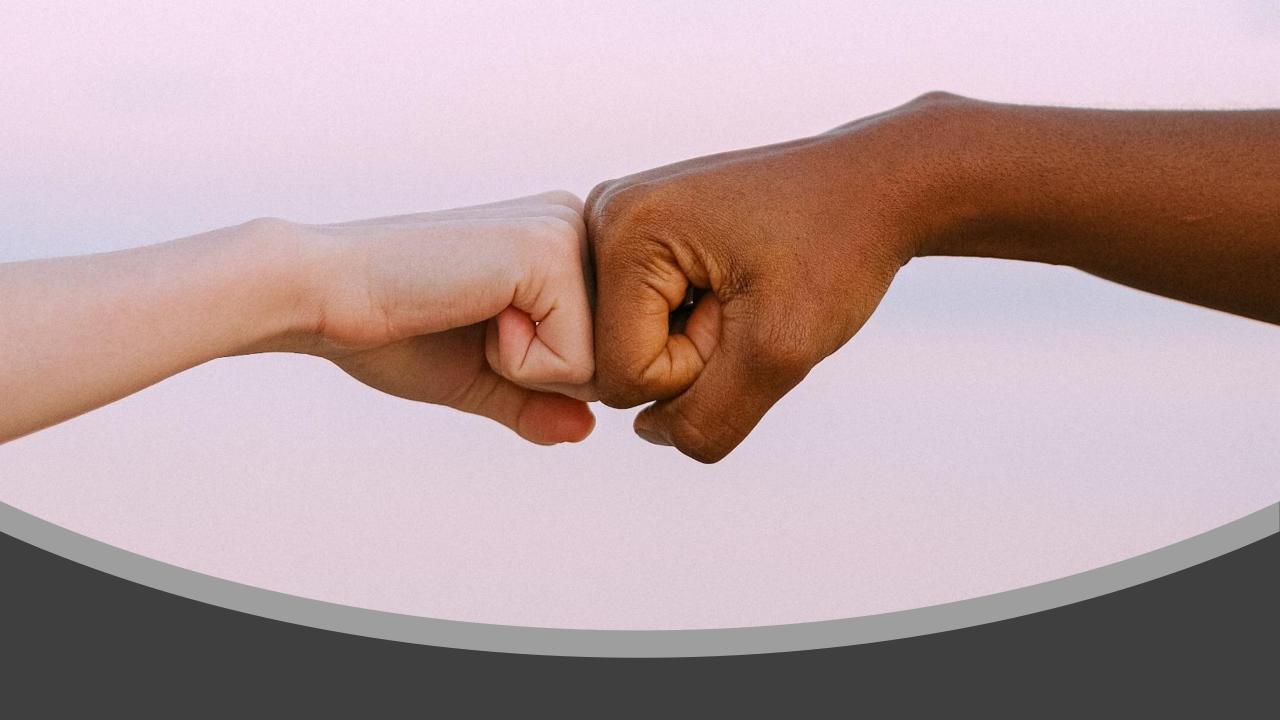


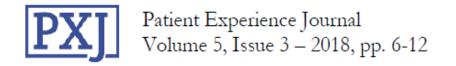
What is your why?





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Commentary

Patient partner compensation in research and health care: the patient perspective on why and how

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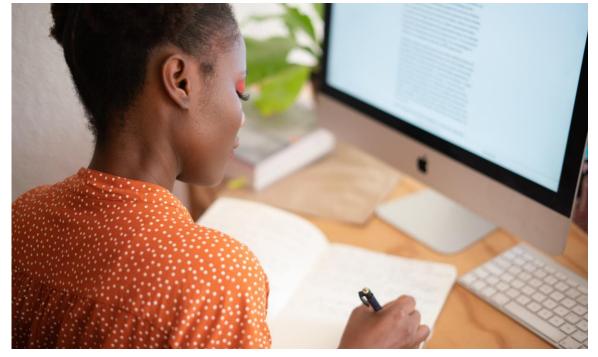
Guidance on authorship with and acknowledgement of patient partners in patient-oriented research



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Abstract

The Strategy for Patient-Oriented Research Chronic Pain Network was founded in 2016 and is a patient-oriented research network funded by the Canadian Institutes of Health Research. The Network incorporates patient partners throughout its governance and operations meaning that patient partners may contribute to research projects in ways that warrant scientific authorship as defined by the International Committee of Medical Journal Editors. The Network did a brief informal review of guidance on patient authorship in 2019, but could not find any practical



It's okay to start small...

Just start





Patient engagement takes a village...



While the path to good engagement isn't always clear, the journey can yield incredible results

