A meta-ethnography of patients' experience of chronic non-malignant musculoskeletal pain


Record Status
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Citation

Authors' objectives
The aim of this study was to increase our understanding of patients' experience of chronic non-malignant MSK pain; utilise existing research knowledge to improve understanding and, thus, best practice in patient care; and contribute to the development of methods for qualitative research synthesis.

Authors' conclusions
Our model helps us to understand the experience of people with chronic MSK pain as a constant adversarial struggle. This may distinguish it from other types of pain. Our findings call on us to challenge some of the cultural notions about illness, in particular the expectation of achieving a diagnosis and cure. Cultural expectations are deep-rooted and can deeply affect the experience of pain. We therefore should incorporate cultural categories into our understanding of pain. Not feeling believed can have an impact on a person's participation in everyday life. The qualitative studies in this meta-ethnography revealed that people with chronic MSK pain still do not feel believed. This has clear implications for clinical practice. Our model suggests that central to the relationship between patient and practitioner is the recognition of the patient as a person whose life has been deeply changed by pain. Listening to a person's narratives can help us to understand the impact of pain. Our model suggests that feeling valued is not simply an adjunct to the therapy, but central to it. Further conceptual syntheses would help us make qualitative research accessible to a wider relevant audience. Further primary qualitative research focusing on reconciling acceptance with moving forward with pain might help us to further understand the experience of pain. Our study highlights the need for research to explore educational strategies aimed at improving patients' and clinicians' experience of care.

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