10th National Patient Reported Outcome Measures (PROMs) Research Conference

Programme

University of Sheffield
The Edge, Endcliffe Village, The University of Sheffield,
34 Endcliffe Crescent, Sheffield S10 3ED

Tuesday 23rd June 2026

Hosted by the Sheffield Centre for Health and Related Research

PROGRAMME OVERVIEW

Tuesday 23rd June 2026

Programme summary		
09:00 - 09:45	**Registration/refreshments**	LOCATION
10:00 – 10:15	Opening remarks: Jill Carlton, University of Sheffield.	High Tor 2
Plenary session 1		
10:15 – 10:45	**Outcome Measurement for Decision Making: Can 'Quality of Life' Be Standardised and Still Meaningful for Patients?** Chaired by Jill Carlton, University of Sheffield. **Andrew Lloyd**, Acaster Lloyd Consulting; **Alix Smith**, Clinical Outcomes Lead, Sheffield Health Partnership University NHS Foundation Trust, and Practice Facilitator, Rotherham, Doncaster & South Humber NHS Foundation Trust	High Tor 2
10:45 – 11:15	Poster viewing and refreshments	Dining room, Ground floor
Parallel session 1		
11:15 – 12:15		
1a Room: High Tor 3	1b Room: High Tor 4	1bc Room: High Tor 2
12:15 – 13:15	Lunch, poster viewing and networking	Dining room, Ground floor
Plenary session 2		
13:15 – 14:15	**Celebrating 10 Years of the PROMs Conference: Where We've Come From and Where We're Headed.** Chaired by Tessa Peasgood, Senior Research Fellow, University of Sheffield. **Mike Horton**, Senior Research Fellow, and Director of Leeds Psychometric Laboratory for Health Sciences, University of Leeds; **Kate Absolom**, Associate Professor in Applied Health Research, University of Leeds; **Sam Llewellyn**, Director of Patient-Centred Research, Vitaccess	High Tor 2
Parallel session 2		
14:20 – 15:20		
2a Room: High Tor 3	2b Room: High Tor 4	2c Room: High Tor 2
15:20 – 15:45	Poster viewing and refreshments	Dining room, Ground floor
Parallel session 3		
15:45 – 16:45		
3a Room: High Tor 3	3b Room: High Tor 4	3c Room: High Tor 2
16:50 – 17:00	Closing remarks and prize presentation: Jill Carlton, Professor of Health Outcomes Research, University of Sheffield. **Dr. David Churchman**, InSpired Health Outcomes Limited	High Tor 2
17:00	Networking	

PARALLEL SESSION 1 DETAILS

11:15-12:15

Session 1a: High Tor 3 Chair: Victoria Gale (University of Sheffield)		
11:15	**Joana Monteiro** Abstract No. 1	Lived experiences of tooth hypersensitivity in children with molar incisor hypomineralisation
11:35	**Hannah Scott** Abstract No. 2	Enabling children often excluded from research to participate in qualitative research to inform outcome measure development
11:55	**Tim Pickles** Abstract No. 3	The Children's Attitude to Grief (CAG) scale: interim results from Rasch measurement theory and confirmatory factor analyses
Session 1b: High Tor 4 Chair: Tara Wickramasekera (University of Sheffield)		
11:15	**Steve Wilkinson** Abstract No. 4	openOutcomes - a PROMs platform for the real world
11:35	**Nirthika Balakumar** Abstract No. 5	Integrating Patient-Reported Outcomes and Experience Within a Primary Care Oral Surgery Service
11:55	**Rebecca Capel** Abstract No. 6	Self-Reported Outcomes in Heart Failure Patients: Real World Evidence from Heart Failure Pathways in Wales
Session 1c: High Tor 2 Chair: Donna Rowen (University of Sheffield)		
11:15	**David Churchman** Abstract No. 7	Responsiveness and Minimal Clinically Important Difference of the Recurrent Urinary Tract Infection Symptom Scale (RUTISS)
11:35	**Ling Jie Cheng** Abstract No. 8	Measurement properties of the experimental EQ-TIPS (EV3.0) in Very Young Children with Acute Infections in Singapore
11:55	**Rachael Pattinson** Abstract No. 9	New developments in the psychometric testing of the Patient-Reported Impact of Dermatological Diseases (PRIDD) questionnaire: responsiveness and minimally important change

PARALLEL SESSION 2 DETAILS

14:20-15:20

Session 2a: High Tor 3 **Chair: Sam Llewelyn (Vitaccess)**		
14:20	**Ben Rimmer** Abstract No. 10	Understanding health-related quality of life after transplant to inform the development of a novel patient-reported outcome measure for solid organ transplant recipients
14:40	**Hannah Jelley** Abstract No. 11	Adaptation and Quality of Life (AdaptQoL): Development of a novel preference-based outcome measure for people using assistive technology.
15:00	**Christina Yiallouridou** Abstract No. 12	Quality of life trajectories in the year following allogeneic stem cell transplantation: a mixed-methods study
Session 2b: High Tor 4 **Chair: Mike Horton (University of Leeds)**		
14:20	**Christel McMullan** Abstract No. 13	Early insights into patients' and carers' experiences of CAR-T therapy
14:40	**Rhiannon Macefield** Abstract No. 14	Development and implementation of an electronic patient-reported outcome system for symptom management to improve post-discharge recovery after surgery for cancer: lessons learnt from a multi-centre randomised controlled trial (the ROSE study)
15:00	**Lorraine Warrington** Abstract No. 15	Mapping Psychosocial Support Pathways to Enable Effective Implementation of an ePROMs system (MyPath) in Metastatic Prostate Cancer Care
Session 2c: High Tor 2 **Chair: Matthew Franklin (University of Sheffield)**		
14:20	**Fiona Lerigo** Abstract No. 16	Using Rasch methods to develop a new measure of care worker-related quality of life in social care: the ASCK-WELL scale
14:40	**Jenna Shepherd** Abstract No. 17	The Oxford Shoulder Score is not a one-trick pony : Factor analysis of data from the National Joint Registry. Should we re-think how we use it?
15:00	**Sally Cox** Abstract No. 18	Linking Patient-Reported Outcomes With Housing Energy Efficiency Data to Inform Breathlessness Service Design in Wales

PARALLEL SESSION 3 DETAILS

15:45 – 16:45

Session 3a: High Tor 3 Chair: Tessa Peasgood (University of Sheffield)		
15:45	**Georgina Jones** Abstract No. 19	The development of a PROM to measure the impact of surgery for prolapse, incontinence, and mesh complications
16:05	**Victoria Robins** Abstract No. 20	Physical functioning after treatment for breast cancer and the impact of age
16:25	**Georgina Forshall** Abstract No. 21	A Patient-Reported Outcome Measure for Caesarean Scar Disorder (CSDi): A Qualitative Interview Study to Establish Content Validity and Inform Item Development
Session 3b: High Tor 4 Chair: Andrew Lloyd (Acaster Lloyd Consulting)		
15:45	**Linda Fenocchi** Abstract No. 22	The Recovering Quality of Life 10-item (ReQoL-10) and EQ-5D-5L instruments in people with a history of suicidal thoughts and behaviours accessing a Distress Brief Intervention (DBI)
16:05	**Emily McDool** Abstract No. 23	Understanding the relationship between the EQ-HWB-9 and age
16:25	**Matthew Franklin** Abstract No. 24	Psychometric Assessment of the EQ-5D-3L And ReQoL Measures in Secure Care Patients with an Intellectual Disability: Reliability, Agreement, Construct Validity, and Responsiveness
Session 3c: High Tor 2 Chair: Kate Absolom (University of Leeds)		
15:45	**Dimity Dutch** Abstract No. 25	The importance of Public and Patient involvement in choosing and refining appropriate tools for measuring Patient Reported Outcome Measures in early childhood obesity research
16:05	**Chryso Hadjidemetriou** Abstract No. 26	The significance of patient involvement when assessing and revising linguistic difficulties in a bronchiectasis patient-reported outcome measure

POSTER PRESENTATIONS: Dining Room, Ground Floor

Poster No.	Presenting Author	Abstract Title
P1	Adeela Rehman	Integrating Patient-Reported Outcome Measures (PROMs) into Maternal Nutrition Research: A Reflective Analysis of Clinical Practices and Patient Perspectives in the United Kingdom
P2	Rachel Chambers	Prioritising Patient Reported Outcome Measures for use in the clinical care of young adults (aged 18-25) living with life-limiting conditions: an adapted nominal group technique
P3	Rachel Martell	Implementing PROMs at Scale: A Trust Wide Assessment of Readiness and Early Progress Through a Community of Practice and Maturity Matrix framework informed by CQC self-evaluation data
P4	Sally Sansom	Measuring outcomes of genome sequencing for rare disease diagnosis in economic evaluations: A psychometric assessment of patient- and carer-reported outcome measurement instruments
P5	Kathryn Jones	Validation of a sex-inclusive patient-reported outcome measure for evaluating lower urinary tract symptoms: The ICIQ-LUTS
P6	Tim Pickles	A systematic review of Patient Reported Outcome Measures for Lupus-specific health-related Quality-of-Life following COSMIN guidelines
P7	Megan Kirk	Quality of life measurement in people living with multiple long-term conditions: a co-produced scoping review and visual evidence map
P8	Kathleen Withers	A qualitative understanding of stakeholder preference towards questionnaires used to support person-centred care in physiotherapy and podiatry outpatients
P9	Hawys Waddington	Iaith mewn Iechyd (Language in Health): Recruiting Welsh-Speaking Volunteers for PROM Cognitive Debriefing
P10	Anna Louise Barry	Informing Patient-Reported Outcome Measurement in Cancer-Related Cachexia: Development of a Patient-Centred Conceptual Model
P11	Louis Fox	Development of an EORTC strategy to measure health-related quality of life in cancer patients receiving immune checkpoint inhibitors: Phase 1 and 2
P12	Jack Lawrence	Patient-reported impacts of atrial fibrillation: a literature-based conceptual model
P13	Nick Burke	AdopteeQOL: Co-creating a quality of life measure for young people who are adopted (adoptees)
P14	Hannah Hussain	One global item or multiple component items : Testing the Psychometric Performance of EQ-5D-5L cognition bolt-ons in the US General Population
P15	Alice Roebuck and Gwawr Evans	Understanding variation in access and response to digitally delivered patient-reported outcome measures in cancer services
P16	Dimity Dutch	How to select a Patient Reported Outcome Measure – resources and support from the COSMIN Initiative
P17	Nyantara Wickramasekera	Developing a Personalised Decision Aid: A Case Study in Ulcerative Colitis
P18	Elizabeth Hepper	A mixed methods study to co-develop a patient reported outcome measure (PROM) for adults who are nil by mouth.
P19	Anna Haire	How Patients Interpret Trade-offs: Embedding PROMs within a Discrete Choice Experiment
P20	Amy Clift	Understanding Preferences About Health States Impacted by Motor Neuron Disease (MND) using a Discrete Choice Experiment

Poster No.	Presenting Author	Abstract Title
P21	Francesca Taylor-Phillips	A qualitative interview study to identify potential barriers and facilitators to implementation of the APPRAISE PROM
P22	Christina Yiallouridou	Assessing financial toxicity among stem cell transplant and CAR T-cell therapy patients.
P23	David Mott	Spillover Effects on Informal and Family Caregivers' Utility: A Systematic Literature Review
P24	Bethany Backhouse	Patients as Data Sources, Not Decision-Makers: A Review of Patient and Public Involvement (PPI) in Women's Health Patient-Reported Outcome Measures (PROMs)
P25	Marci Kay Livingston	PROs, PROMs, and Lived Experience: Findings from a Scoping Review of Methodological Guidance for Core Outcome Set Long-List Development
P26	Ling Jie Cheng	Feasibility of Latent Scale Discrete Choice Experiments for Valuing EQ-TIPS-5L Health States: A Qualitative Study in the UK and Singapore
P27	Sophie Franckel	EHR Integrated LYMPROM Visualisations to Support Lymphoedema Care
P28	Kodchawan Doungsong	Patient-reported experience measures (PREMs) and patient-reported outcome and experience measures (POEMs) for clinical trials and health economics studies in vision disorders: a systematic review
P29	Jingni Yan	What does change mean? Meta-awareness and the interpretation of PANAS negative affect change scores
P30	Alex Blakoe	How are ePROMs used in the NHS as part of routine secondary care?
P31	Lily Nicholson	The measurement properties of menstrual blood loss measurement instruments in women with heavy menstrual bleeding: a systematic review
P32	Tessa Peasgood	Qualitative testing of potential modifications to the EQ-HWB-9 in the United Kingdom
P33	Devin Peipert	A targeted guidance and case study review of patient reported outcomes in regulatory evaluation of advanced therapies
P34	Sarah Smith	A new patient-reported experience measure (PREM) for maternity and neonatal services in England: MatNeoPREM
P35	Anastasiia Demidova	Outcome Domains and Measurement Instruments in Studies of Brain Infection Sequelae: A Scoping Review
P36	Katie Wallace	Early assessment of digital PROMs adoption in colorectal oncology
P37	Hayat Hamzeh	Mapping and content analysis of the European Organisation for Research and Treatment of Cancer (EORTC) Item Library using the World Health Organization International Classification of Functioning, Disability and Health (ICF)
P38	Georgina Jones	The development of the first adenomyosis-specific quality of life patient-reported outcome measure
P39	Alex Hind	The Paradox of Change-from-Baseline Correlation: Low Change Correlation Despite High Cross-Sectional Correlation.
P40	Rebecca Dawkins	Patient-Reported Tolerability (PRT) in Industry Sponsored Oncology Clinical Trials: Current Practices and Regulatory Implications