



UNIVERSITY OF LEEDS

This is a repository copy of *Patient and surgeon perspectives of a large-scale system for automated, real-time monitoring and feedback of shared decision-making integrated into surgical practice: a qualitative study.*

White Rose Research Online URL for this paper:

<https://eprints.whiterose.ac.uk/id/eprint/227561/>

Version: Accepted Version

Article:

Hoffman, C., Avery, K., Macefield, R. et al. (11 more authors) (Accepted: 2025) Patient and surgeon perspectives of a large-scale system for automated, real-time monitoring and feedback of shared decision-making integrated into surgical practice: a qualitative study. BMJ Open. ISSN 2044-6055 (In Press)

This is an author produced version of an article accepted for publication in BMJ Open, made available under the terms of the Creative Commons Attribution License (CC-BY), which permits unrestricted use, distribution and reproduction in any medium, provided the original work is properly cited.

Reuse

This article is distributed under the terms of the Creative Commons Attribution (CC BY) licence. This licence allows you to distribute, remix, tweak, and build upon the work, even commercially, as long as you credit the authors for the original work. More information and the full terms of the licence here: <https://creativecommons.org/licenses/>

Takedown

If you consider content in White Rose Research Online to be in breach of UK law, please notify us by emailing eprints@whiterose.ac.uk including the URL of the record and the reason for the withdrawal request.



eprints@whiterose.ac.uk
<https://eprints.whiterose.ac.uk/>

Hoffman C, Avery K, Macefield R, Snelgrove V, Rooshenas L, Bekker HL, Hopkins D, Cabral C, Blazeby J, Gibbison B, Hickey S, Williams A, Aning J, McNair AGK. *Patient and surgeon perspectives of a large-scale system for automated, real-time monitoring and feedback of shared decision-making integrated into surgical practice: a qualitative study*. (Submitted 14th Jan 2025 BMJ Open; accepted 4/6/2025)

Title:

Patient and surgeon perspectives of a large-scale system for automated, real-time monitoring and feedback of shared decision-making integrated into surgical practice: a qualitative study

Authors:

Christin Hoffmann¹, PhD, 0000-0002-6293-3813
Kerry Avery¹, PhD, 0000-0001-5477-2418
Rhiannon Macefield¹, PhD, 0000-0002-6606-5427
Val Snelgrove², HNC
Leila Rooshenas¹, PhD
Hilary L Bekker^{9, 10}, PhD, 0000-0003-1978-5795
Della Hopkins³, PhD, 0009-0000-6390-4742
Christie Cabral⁵, PhD, 0000-0002-9884-0555
Jane Blazeby¹, MD, 0000-0002-3354-3330
Ben Gibbison^{1,6}, MD, 0000-0003-3635-6212
Shireen Hickey⁴, MBChB
Adam Williams³, MBChB
Jonathan Aning^{7, 8}, BMedSci, DM, 0000-0001-6189-4720
Angus GK McNair^{1,3}, PhD, 0000-0002-2601-9258

Collaborators:

Andy Judge¹;
Andrew Smith³;
Archana Lingampalli⁶;
Barnaby Reeves¹¹;
Jessica Preshaw³;
Michael R Whitehouse^{1,3};
Paul Cresswell³;
Philip Braude³;
Shelley Potter³;
Timothy Beckitt³;
Timothy Whittlestone³

Author affiliations:

¹ NIHR Bristol Biomedical Research Centre, University Hospitals Bristol and Weston NHS Foundation Trust and University of Bristol, Bristol Medical School: Population Health Sciences, University of Bristol, Bristol, BS8 2PS, UK

² Patient Representative

³ North Bristol NHS Trust, Bristol, UK

⁴ Bradford Institute for Health Research, Bradford Teaching Hospitals NHS Foundation Trust, Bradford, UK

⁵ Centre for Academic Primary Care, Bristol Medical School, University of Bristol, Bristol, UK

⁶ University Hospitals Bristol and Weston NHS Foundation Trust, Bristol, UK

⁷ Bristol Urological Institute, Southmead Hospital, North Bristol Trust, Bristol, UK

⁸ Population Health Sciences, Bristol Medical School, University of Bristol, Bristol, UK

⁹ Leeds Unit of Complex Intervention Development (LUCID), Leeds Institute of Health Sciences, School of Medicine, University of Leeds, Leeds, UK

¹⁰ The Research Centre for Patient Involvement (ResCenPI), Department of Public Health, Aarhus University, Central Denmark Region, DK

¹¹ Bristol Trials Centre, Population Health Sciences, Bristol Medical School, University of Bristol, Bristol, UK

Corresponding author: Christin Hoffmann (c.hoffmann@bristol.ac.uk), National Institute for Health and Care Research Bristol Biomedical Research Centre, Bristol Centre for Surgical Research, Bristol Medical School: Population Health Sciences, University of Bristol, Bristol, BS8 2PS, UK

Data availability statement: Summaries and examples of anonymised data have been included as supplementary material in this manuscript. Original pseudo-anonymised datasets are stored in a non-publicly available repository. Pseudo-anonymised datasets can be made available via the University of Bristol's Research Data Repository, data.bris to bona fide researchers, subject to a legally binding data access agreement. Any applications to access data will involve a case-by-case review by the University of Bristol Data Access Committee. Qualifying researchers will be required to sign a data access agreement and closely liaise with study team members to ensure that the data they plan to make public are sufficiently anonymised. Generally, data will be made available for non-commercial use, only for the purpose of health and care research and with appropriate approvals in place.

Abstract

Objective: To explore patient and healthcare professional perceptions about the acceptability and impact of a large-scale system for automated, real-time monitoring and feedback of shared decision-making (SDM) that has been integrated into surgical care pathways.

Design: Qualitative, semi-structured interviews were conducted with patients and healthcare professionals between June and November 2021. Data was analysed using deductive and inductive approaches.

Setting: Large-scale measurement of SDM has been integrated in NHS surgical care across two large United Kingdom National Health Service Trusts.

Participants: Adult surgical patients (N=18, 56% female), following use of an SDM real-time monitoring and feedback system, and healthcare professionals (N=14, 36% female) involved in their surgical care. Patient recruitment was conducted through hospital research nurses and professionals by direct approach from the study team to sample individuals purposively from seven surgical specialties (general, vascular, urology, orthopaedics, breast, gynaecology, urgent cardiac).

Results: Ten themes were identified within three areas of exploration that described factors underpinning: (1) the acceptability of large-scale automated, real-time monitoring of SDM experiences, (2) the acceptability of real-time feedback and addressing SDM deficiencies, (3) the impact of real-time monitoring and feedback. There was general support for real-time monitoring and feedback because of its perceived ability to efficiently address deficiencies in surgical patients' SDM experience at scale, and its perceived benefits to patients, surgeons and the wider organisation. Factors potentially influencing acceptability of large-scale automated, real-time monitoring and feedback were identified for both stakeholder groups, e.g. influence of survey timing on patient-reported SDM scores, disease-specific risks, patients' dissatisfaction with hospital processes. Factors particularly important for patients included concerns over digital exclusion exacerbated by electronic real-time monitoring. Factors unique to professionals included the need for detailed, qualitative feedback of SDM to contextualise patient-reported SDM scores.

Conclusion: This study explored factors influencing the acceptability of automated, real-time monitoring and feedback of patients' experiences of SDM integrated into surgical practice, at scale amongst key stakeholders. Findings will be used to guide refinement and implementation of SDM monitoring and feedback prior to formal development, evaluation, and implementation of an SDM intervention in the NHS.

Trial registration: ISRCTN17951423

Strengths and limitations of this study

- A strength of this study is its methodologically robust approach to explore the perspectives of both patients and healthcare professionals
- Factors underpinning acceptability and impact of a system for automated, real-time monitoring and feedback of shared decision making (SDM) were identified to inform its large-scale implementation and intervention development to improve SDM at scale
- Limitations with regards to diversity and inclusivity have been identified that are being addressed in ongoing work, which examines the views of under-served groups on real-time SDM monitoring and feedback

The original protocol: Supplementary file 2.

Funding statement: The ALPACA Study is funded by the National Institute for Health and Care Research (NIHR) Programme Development Grant (NIHR205174). This study was also supported by an NIHR Clinician Scientist award to AM (NIHR CS-2017-17-010) and delivered through the NIHR Biomedical Research Centre (BRC) at the University Hospitals Bristol and Weston NHS Foundation Trust and the University of Bristol (BRC-1215-20011, NIHR203315). The views expressed in this publication are those of the authors and not necessarily reflect those of the NIHR, the Department of Health and Social Care or the UK National Health Service.

Competing interest statement: All authors declare no conflict of interests.

Introduction

Shared decision-making (SDM) is a key pillar of patient-centred care [1,2] and can impact health outcomes [3]. High-quality SDM discussions for surgical treatment decisions is particularly important because consequences are usually irreversible. About one-third of surgical patients, however, report deficiencies with SDM experiences [4]. Improving surgical care pathways to enable consistent SDM practices SDM remains a key patient and policy priority in many modern healthcare systems [1,5,6].

Health policy recommends a comprehensive approach to improving SDM in practice. It suggests a combination of interventions to support patients, professionals and organisations, e.g. use of patient decision aids, provision of healthcare professional training and supportive leadership [7,8]. Routine measurement and evaluation of patient-reported SDM within healthcare organisation systems has been identified as essential for driving these improvements, with financial incentives promoting its uptake [9,10]. Although previous United Kingdom (UK) service improvement programmes incorporated SDM measurement in specific healthcare contexts [11–13], limited evidence exists for how to effectively integrate, monitor and sustain SDM measurement at scale [3,8]. Feedback has shown to positively impact clinical practice [14], and that learning from patient-reported experiences leads to improvements in care [15,16]. For example, integrated, real-time symptom monitoring has shown to improve overall two-year survival in patients with advanced cancer compared to usual care. There is uncertainty about how best to embed feedback of patient-reported SDM scores to drive meaningful and sustained behaviour change within clinical teams and organisations along the care pathway to affect national practice improvements [17–19].

We integrated within surgical care pathways of two NHS trusts, a novel electronic system to monitor and feedback surgical patients' experience of SDM automatically and in real-time [20,21]. The system has the potential to support large-scale interventions to enhance patients' experiences of SDM before surgery and to ultimately improve patient and health service outcomes [22,23]. In-depth exploration, using qualitative methods, of stakeholders' views towards the system is important to inform organisation-wide implementation of surgical patient-reported measurement systems [24–27]. Surgical patients and surgeons are key stakeholders and primary intended users of real-time SDM monitoring and feedback. Improved understanding of their perceptions of the system and factors influencing its acceptability, is necessary to facilitate its effective implementation and intervention development to improve SDM.

Aim

To explore patient and healthcare professional perceptions about the acceptability and impact of a system for automated, real-time monitoring and feedback of SDM integrated into elective surgical care pathways in the NHS.

Methods

This qualitative study was guided by the interpretivist paradigm which epistemologically and ontologically acknowledges multiple realities [28]. It employed semi-structured interviews to elicit views and experiences of patient and healthcare professional participants and applied codebook thematic analysis to generate themes [29]. Conduct and reporting of this study followed the consolidated criteria for reporting qualitative research (COREQ) checklist [30].

Study context

This study was part of a larger programme of work (the ALPACA Study) that seeks to co-develop a decision support intervention to improve SDM practices within surgical pathways[20]. The intervention is expected to use a large-scale system for automated and real-time monitoring and feedback of patients' experiences of SDM before surgery. Details about the system and processes for its integration into surgical practice are available elsewhere [20,21] and briefly described below.

Integrating real-time SDM measurement in surgical care pathways was achieved through procuring and customising an off-the-shelf electronic patient-reported outcome software (Cemplicity Ltd., New Zealand) in two NHS Trusts in England (North Bristol Trust/NBT, University Hospitals Bristol and Weston Foundation Trust/UHBW) from April 2021. In collaboration with the software provider, Structured Language Queries were developed to enable secure, automated, daily data transfers between the system and hospital data warehouse to monitor and feedback surgical patients' experiences. Monitoring involved the system automatically administering validated, electronic patient-reported SDM measurement instruments via text message or e-mail to patients within 24 hours of elective surgery booking, following SDM discussions. Patients' responses were received in real-time and securely returned to NHS Trust Electronic Patient Records. Feedback involved the system processing and displaying patients' responses in an electronic dashboard accessible to the clinical teams.

The participating hospitals are large tertiary care centres in the South West of England, UK. NBT is among the largest acute NHS trusts in the UK, offering a comprehensive range of acute clinical care

to both local and regional clinical commissioning groups in South West of England. Surgical departments where the system for real-time SDM monitoring and feedback has been integrated included services in general surgery, urology, gynaecology, orthopaedic, breast and vascular surgery. One of UHBW's departments was included as the regional cardiac surgical centre for South West of England.

Data were collected for NHS quality improvement purposes with approval from Trust clinical governance committees. Ethical approval to conduct interviews was granted by the NHS HRA North West - Liverpool Central Research Ethics Committee.

Participants, sampling and recruitment

Patient participants

Patient participants were recruited from the two NHS Trusts (NBT, UHBW) where the system for real-time monitoring and feedback of SDM had already been integrated into surgical practice.

Patients were eligible if they used the electronic platform to complete an SDM measure as part of the wider study. The electronic platform invited all adult patients booked for elective surgery across general, gynaecological, breast, urological, orthopaedic or vascular surgical departments (NBT) and urgent cardiac surgery (UHBW). Patients were excluded if they lacked decisional capacity for medical treatments, had undergone unplanned (emergency) surgery or were booked for endoscopic procedures these include large volumes of diagnostic procedures. A sub-set of those patients were sampled for the current study.

A purposive sampling strategy was adopted to ensure that insights are drawn from a range of perspectives. Sampling characteristics included variation by i) experiences of different types of surgery (e.g. general, vascular, urology, orthopaedics, breast, gynaecology, urgent cardiac), ii) experience of SDM process (i.e. good/poor experience as determined by SDM scores), and iii) socio-demographic characteristics (i.e. sex, age). Participant characteristics were assessed as the study progressed and recruitment efforts targeted under-represented patients. No further socio-demographic details were collected to retain anonymity of participants. Data on the number of participants declining to take part in interviews were not collected.

Potential patient participants were identified and contacted by a member of hospital staff (e.g. research nurse) with authorised access to patient data and the system for real-time SDM monitoring

and feedback. Contact was initiated via email or telephone to explain the study, send the patient information sheet and consent form, and to ask for permission to pass on their contact details to a researcher. If patients agreed and expressed interest in participating, a member of the study team (CH) followed up with a phone call to answer any queries and arrange a suitable interview date.

Recruitment stopped when no new codes or meaning were identified through analysis of additional transcripts. This involved multi-disciplinary team discussions reviewing identified codes/themes provided sufficient conceptual depth to answer the research question (see analysis section for more details) [31–33].

Professional participants

Healthcare professionals were consultant surgeons working in the participating surgical departments. Included were those that: 1) booked eligible patients for surgery, 2) were involved in SDM discussions with eligible patient participants, or 3) had overall responsibility for eligible patients' care.

Eligible healthcare professionals were identified through their involvement in surgical teams within the two NHS Trusts and approached by the principal investigator (AM). Contact was initiated through face-to-face conversations in the hospital or clinical departmental meetings where the project was presented, inviting healthcare professionals to take part in interviews. Follow-up emails including a participant information sheet and consent form were sent to healthcare professionals and teams expressing interest to arrange a suitable time and date for an interview.

A purposive sampling approach sought variation by surgical specialty and sex. As above, no information about other personal details (e.g. ethnicity, age) or those declining the interview were recorded. Recruitment continued until sufficient conceptual depth was achieved as outlined above.

Data collection

Semi-structured, one-to-one interviews with participants were undertaken remotely using video conferencing software (Zoom, MS Teams) or telephone. All interviews were audio-recorded and conducted in English by one of two authors (AM, a methodologist and academic surgeon not involved in the care of participants; CH a social scientist with extensive experience of health services

research). Both interviewers are trained in qualitative methods and have extensive experience conducting qualitative research in secondary care settings and for health services research.

Patients received a participant information sheet and completed an electronic consent form prior to the interview. Participants were made aware of the study purpose, aims and why they were invited to participate. Interviewers introduced themselves to explain their occupation and role in the study. It was emphasised that in case the patient participant raises any issues about their clinical care, the interviewees were unable to address these, but could direct them back to the clinical team.

Topic guides, developed separately for patient and professional stakeholders, were piloted and refined before interviews commenced (Supplemental File 1). Topic guides focused on two areas of exploration aligned with the study aim: acceptability of real-time monitoring and feedback and impact of this on clinical care. These were defined *a priori* as topics of interest to support the wider ALPACA study [20].

Analysis

Anonymised interview transcripts were uploaded to a qualitative data management software (Nvivo, Version 14). The analytical process followed principles of deductive and inductive coding [34] and template analysis [35] and was performed in parallel to data collection to prospectively inform ongoing participant sampling. Data analyses were performed separately and sequentially for each stakeholder group (healthcare professionals, patients) to allow exploration of any potential differences within their experiences and context. Analyses were performed by two trained and experienced qualitative researchers (AM, CH).

The analytical process involved the two researchers separately (i) reading the transcripts several times; (ii) deductively coding participants' responses according to the two broad areas of exploration (i.e. the acceptability of real-time SDM monitoring and feedback and impact on clinical practice); (iii) generating initial codes within the two broad areas of exploration, by assigning latent and semantic labels to relevant excerpts and drawing on constant comparison [36]; (iv) generating early themes by collating similar codes and defining an initial coding template [35]; and (v) iteratively refining the coding template and early themes through interpretation of the data and by triangulating views through multi-disciplinary study team discussions. Themes were further developed through analysis of more sets of transcripts, applying and further modifying the coding template as necessary.

Summaries (descriptive reports) of findings were written for each stakeholder group following rounds of transcript analyses and discussed within the multi-disciplinary study team and patient

advisory group. Discussions were used to enrich interpretations of the data by exploring new lines of enquiry considered important to addressing the study objectives. The summaries were iteratively developed as analyses proceeded. The separate reports were then synthesised to form the basis for research outputs by comparing and contrasting participants' accounts within and between stakeholder groups. Thereby, themes with similar conceptual meaning were combined and theme labels harmonised; themes that were identified as conceptually distinct and unique to either patient or professional participants were kept separate.

Patient and Public Involvement statement

A dedicated patient and public advisory group was established for the wider ALPACA study [20]. Responsibilities included oversight of study and PPIE activities to ensure these are aligned with patient priorities. The group was involved in interpreting the results following initial analyses. The patient co-author (VS) co-developed topic guides and recruitment materials for this study, and was involved in writing this manuscript.

Results

Participant characteristics

A total of 18 patient and 14 professional participants were interviewed. Interviews were conducted between June and November 2021 and lasted between 25 and 57 minutes. One patient participant consented, but did not attend the interview. All professional participants who consented to take part proceeded with the interview.

Participant characteristics are summarised in Table 1. Patient participants were 56% female, and all from a white British background. They were on average 52 years old and their surgical experiences spanned a total of seven surgical specialties. Healthcare professionals were 64% male and represented consultant surgeons from three surgical specialties.

Table 1. Participant characteristics

Characteristic	Patient participants (N=18)	Professional participants (N=14)
Age (years), Mean (\pmSD)	52 (\pm 14)	Not reported
Sex, n		
Female	10	5
Male	8	9
Ethnicity, n		
White British	18	Not reported
Specialty, n		
Urology	5	7
General	4	3
Breast	3	-
Orthopaedics	2	-
Urgent cardiac	2	-
Vascular	1	4
Gynaecology	1	-

Themes identified from interviews

A total of 10 themes relating to participants' perceptions were identified across three areas of exploration (acceptability of real-time monitoring, acceptability of real-time feedback and addressing SDM deficiencies, impact on clinical practice). An overview of themes and summaries of findings per theme are displayed in Table 2. Detailed descriptions of each theme, including illustrative quotes are provided below. Anonymised quotes were labelled with a unique participant ID, using acronyms 'PAT' (patient participant) and 'HCP' (healthcare professional) to distinguish their participant groups.

Table 2. Overview of areas of exploration, identified themes and summary of findings

Area of exploration		Themes	Summary of findings
Acceptability	Real-time monitoring of SDM experiences	Benefits and challenges in identifying SDM deficiencies	All participants expressed support for a real-time monitoring system to assess SDM experiences and its ability to detect deficiencies in SDM experiences. Professionals discussed engagement-related challenges.
		Complexities of SDM measurement	Both participant groups discussed the complexities of real-time SDM monitoring and pointed out common sources of variability of their SDM experience that might influence patient-reported SDM scores.
		Challenges of distributed decision making	Some participants in both participant groups noted that a poor SDM experience can depend on patients' holistic care experience and multi-professional care influencing SDM measurement.
		Limited inclusivity of real-time monitoring	Both participant groups expressed concerns regarding digital and language inclusivity, with patients frequently highlighting limitations for elderly, ethnic minority and disabled patients.
	Real-time feedback and addressing SDM deficiencies	Balanced and informative feedback	Both participant groups highlighted the importance of sharing positive and negative feedback, with professionals emphasising the need for details about reasons for poor SDM experiences.
		Approaches to providing feedback to clinical teams	Professionals discussed a desire for 'triage process' where feedback is initially handled by intermediaries before involving surgeons.
		Approaches to responding to feedback	Participants favoured follow-up encounters to address deficiencies in patient-reported SDM experiences, emphasising that tailored discussions need to address individual concerns.
	Impact on clinical practice	Perceived impact on patients	All participants stressed the important role of the system in promoting patient-centred care, with patients discussing specific benefits of decision satisfaction and reassurance.
		Perceived impact on surgeons	Participants perceived the system as beneficial for identifying practice improvements and training needs, with professionals expressing concern of workload and emotional burden.
		Perceived impact on department or organisation	Professionals discussed the potential of the system for identifying and addressing systemic performance issues, highlighting benefits such as trend analysis and organisational reputation.

1. Acceptability of real-time monitoring of SDM experiences

1.1 Benefits and challenges in identifying SDM deficiencies

Participants expressed overall support for electronic, automated real-time measurement of patients' SDM experiences. Participant highlighted benefits of the electronic survey, such as *"it was quite quick"* (PAT004), *"very easy to complete"* (PAT015) and *"seems efficient"* (HCP004), and *"straightforward and simple to use"* (HCP013), which support acceptability of SDM monitoring.

It was widely recognised that monitoring (i.e., the survey) has the potential to detect SDM deficiencies and provides opportunities to improve SDM experiences.

"I want that opportunity to be able to discuss exactly what's involved and you know if that can happen as a result of this survey then that's great." (PAT003)

"I received the survey without knowing they had made a decision to have an operation. [...]" (PAT018)

"It seemed one of those projects that you think how do we not know that? How have we not asked this before? Why are we just steaming ahead with information leaflets and bespoke consent forms with all the risks written out and yet we didn't ask them whether it was what they wanted rather than what a medico legal team thought we should do. So it's a critical project." (HCP006)

Patient participants often cited an interest in giving feedback and thereby potentially improving care for future patients as important drivers for completing SDM monitoring. Interviewees often noted that reporting SDM experience can lead to feeling valued and listened to.

"I think most people would fill out a survey like this, especially for, if it's on their own health." (PAT004)

"Unless you were part of doing a survey on something like this, you never really get to tell people your experience, whether it be a good one, whether it be a bad one, you might get somebody who has had the most horrendous time" (PAT007)

"Again, being taken seriously, so if there is a real, serious problem, sometimes it's not heard, it's not listened to and I feel [...] that it's going to help in the future" (PAT002)

Some professional participants also commented on the wide applicability real-time SDM monitoring beyond surgery, demonstrating the perceived potential to enhance SDM in various clinical settings.

"It could be extended to the non-operative cases" (HCP008).

"So what's good about it is it's rapid, the questionnaire is short, [...] it's not complex and the questions are quite well designed. If you look at them you think you know what, actually they have wide applicability." (HCP001)

Professional participants emphasised the importance of healthcare professional buy-in for the successful uptake and acceptance of a system for real-time SDM monitoring. They highlighted the

need for engagement strategies that address concerns such as time pressure and fear of SDM monitoring being perceived as tool for assigning blame for deficiencies in decision-making.

“[D]o it in a way that doesn’t make time pressured, angry surgeons more angry” (HCP003)

“What we’ve got to do is engage the clinicians in the fact that it’s not a blame culture” (HCP001)

The importance of engaging all clinical team members during the implementation process was also apparent in one patient’s experience who reported: *“The ward doctor was surprised I got the survey because he didn’t know you did surveys” (HCP018).*

1.2 Complexities of SDM measurement

Both participant groups discussed complexities of real-time SDM monitoring. Participants shared their concern about SDM being influenced by the context in which care is provided and varying preferences for decision styles. Most commonly, procedure/disease and specialty-dependency were highlighted. In particular, participants stressed the importance of SDM for elective surgery and operations that carry the greatest risks and impacts on patients’ quality of life.

“All of us [patients] don’t necessarily want to question the decisions of those who have positions which are not equal to our own do we. [...] I think it will depend on how patients feel about their consultations and where in the cycle of treatment they get it because there’s no doubt at all if you’re looking at cancer it’s very emotive. Surgery often takes place very quickly [...] so it’s how you catch people and that will be quite individual in some ways.” (PAT001)

“[...]bigger operations have higher risk, it’s more important that those patients are engaged in the process, particularly for what we do. Carotid endarterectomy [surgery to unblock a main blood vessel], an operation that prevents something that might or might not happen, shared decision-making is crucial.” (HCP001)

About half of patient participants also mentioned other sources of potential variability in patient-reported SDM experiences which may influence responses to real-time data capture. Participants discussed the importance of timing of the SDM survey, suggesting that patients might feel too unwell or distressed at various points in time before surgery.

“I think probably if its straight after the consultation or before the operation, I mean a lot of these people are... well I know I was, I was really ill and probably giving your time to do that is probably not a high priority.” (PAT005)

Patients not wanting to criticise their care team or potentially affecting care (see also theme 2.1) was mentioned as another factor that may influence responses.

“My mum is a very quiet person and she wouldn’t push forward for anything, you know, she wouldn’t go, well actually I’m not happy about that, and there are lots of people like that and they may not want it to rock the boat or cause problems when they have surgery.” (PAT002)

All professional participants expressed concerns about whether patient-reported poor experiences always reflected SDM deficiencies, potentially affecting surgeons' acceptability of the measurement system. Interviewees discussed several factors they believed were unrelated to patients' SDM experience but could influence responses to SDM monitoring. These included the possibility that patients may have misinterpreted the survey questions or conflated other experiences (e.g., dissatisfaction with hospital processes), which could lead patients to report poorer experiences through the survey.

"I think patients can just lump everything [...] into one big umbrella and [...] saying that they're not happy about shared decision-making, actually if you unpick it, it might be more about the practicalities of the booking process or the kind of logistics" (HCP010)

"It's not always that it is a poor experience as such, it's just that you might have worked on a slightly different agenda to them or you may not have said what they wanted you to say." (HCP003)

"[I]f a patient's told that they have to have an operation, then they have to think about the consequences [but] they might not instantly think about all that and have all their questions ready and then they might become unhappy that they've forgot to or it was sprung on them." (HCP008)

Professional participants also highlighted broader challenges in capturing patients' SDM experiences by acknowledging *"It can be a difficult thing to measure"* (HCP007). Interviewees drew comparisons to other patient-reported outcome measures and associated potential difficulties with interpreting scores.

"I still think it is difficult because if you give three patients a pain score who have the same pain, they will score differently on the pain scales because some of them will be more bothered by it and others will be less bothered by it." (HCP012)

1.3 Challenges of distributed decision making

Though discussed by fewer participants, both participant groups talked about challenges in SDM monitoring in distributed decision-making contexts that could affect stakeholder acceptability of SDM monitoring. This means care pathways where patients are under the care of multiple healthcare professionals (e.g., general practitioner, oncologist, nurse, surgeon) and where the SDM experience cannot be isolated to a single event. For example, patients may be referred from other hospitals or are on pooled operating lists where multiple healthcare encounters influence the decision-making experience.

"The whole feeling with this whole gynae thing, is that it doesn't feel very centralised, you're dealing with the physio, you go and see a nurse for the pessary, you just don't feel like it's very joined together." (PAT017)

"My worry about urology for example is about our pooled list system [...] I don't really want to be getting alerts when one of my other colleagues has not had a good interaction with a patient in the one stop clinic." (HCP010)

A perceived lack of communication between institutions and teams ultimately led to a feeling of uncertainty about whether their health problems were understood and doubt whether surgery was really the best option. A small number of patients explained how discrepancies or delays in information provision caused by different teams or institutions may impact their SDM experience which may affect patient-reported experiences.

“To be honest, up until the phone call - up until six weeks before I was due for my op, I've had a little contact with the doctor, I've had very little information. I've been given probably three different diagnoses of what was wrong with my knee.” (PAT015)

1.4 Limited inclusivity of real-time monitoring

Both participant groups commonly discussed barriers to digital and language inclusivity common to accessing electronic, real-time monitoring systems in a healthcare setting.

Patient participants frequently highlighted limitations in relation to patients' age, preferred language and disability status which can increase difficulty engaging in real-time SDM monitoring without the synchronous provision of additional support.

“I think the challenges would be people's ability to use technology. Moving forward there are people of any age group who still can't, especially if you have got people with special needs. [...] There are obviously sectors in society where they need more care in a situation like that, because it is very frightening, so filling out questionnaires beforehand, they are always gonna need help. [...] when you have got elderly or people with a disability or a learning disability, their interpretation of things can cause a stressful situation” (PAT007)

Professional participants also raised concerns about potentially encouraging health-related inequalities for a range of under-served groups.

“People with learning disabilities, people with mental health issues, people whose literacy levels are not high might struggle with this.” (HCP004)

“One of the concerns that I have with any form of patient reported measures [...] they lead to an increase in health inequality and I think there is a little bit of evidence for that as well is that those who shout loudest, tend to get the most input and those who are quietest who are often those that need the most input” (HCP011)

Some suggested the lack of inclusivity needs addressing in later stages of implementing real-time monitoring of SDM to avoid exacerbating inequalities in future healthcare provision.

“I think the only downside for me is the health and inequalities possibly, that you're not reaching the hard to reach groups, so that's the only downside but it's still better than doing nothing which is what we do now so, even if you don't reach everyone, it's still better than nothing.” (HCP004)

“The problem of course is that we're using an electronic format and we are therefore serving ourselves up to patients who've got Smartphones, who are electronically engaged, who speak good English and therefore largely the middle-classes. [...] So we by definition have a huge sampling error in this and [...] it's going to have to be a project priority at some point to identify those patients and think about another intervention” (HCP001)

2. Acceptability of real-time feedback and addressing SDM deficiencies

2.1 Balanced and informative feedback

Both participant groups considered it appropriate to provide feedback on patients' personal SDM experience and stressed the importance that SDM deficiencies should be known.

"Yes, I would like them to know how I feel about it and my concern." (PAT008)

"I'm terribly concerned about the individual. So, if a person hadn't had a good experience, I would want to put it right for that individual." (HCP009)

"I think from a personal perspective, if you'd seen that patient, you'd want to know which areas they felt the decision making was deficient in." (HCP002)

The need to share positive alongside negative feedback was highlighted by stakeholders in different ways. Patient participants raised concerns that providing negative feedback may cause distress and additional workload for healthcare professionals, or could negatively affect their own care.

"I don't want to put pressure on anybody that's under pressure as it is. [...] It's a worry because you do feel, so if I say this, will they not do my surgery? Will they take it to heart and you know, or will I cause stress and anguish with the surgical team?" (PAT002)

Professional participants also suggested that balancing positive and negative messages can impact acceptability in real-time monitoring and feedback amongst surgeons who demonstrate lower engagement.

"I think the other thing is not to forget positive feedback can be as important as negative. So actually, as a clinician, occasionally getting a positive feedback is also very nice. I think, particularly as surgeons, we are very good at picking up when someone else has done something wrong."" (HCP005)

"The problem is that you get a lot of egotistical characters, and I think that a lot of times the base response will be 'I didn't do anything wrong.' So, you've got to make sure that information is coming across in a way that you're not reinforcing that that initial reaction." (HCP007)

Professional participants unanimously agreed that receiving patient-reported scores from real-time SDM monitoring is helpful to contextualise responses, but only receiving numerical data is insufficient to address any deficiencies in patients' SDM experience. Participants suggested that additional, qualitative detail about the reasons for patients' low scores is needed to be able to contextualise patients' responses. Some interviewees also preferred if feedback also contains a call to action to facilitate further action.

"[These] scores mean nothing to the clinician." (HCP001)

"I think it needs to be more descriptive. Yes, you can give the overall number as well but I still think you need more description one way or another." (HCP012)

"I suppose some detail as to what made that an unsatisfactory experience for the patient and also what to do next." (HCP008)

2.2 Approaches to providing feedback to clinical teams

Professional participants discussed in detail how to provide feedback of patient-reported SDM scores to clinical teams, evaluating the acceptability of different approaches to using feedback. Interviewees initially discussed surgeons receiving feedback as personal notification (email or text message) about poor patient-reported SDM experiences to prompt further intervention. This was found acceptable in case of low number of patients requiring attention and sometimes preferred by some participants.

"[T]he actual practicalities of how I want to be fed back, I wouldn't be averse to having an email alert to say that somebody that's under my care is worried about their surgery. [...] I know we have a lot of emails, but if we had... I think because the numbers would be relatively low and we want to obviously nip those sort of issues in the bud, I think it's a reasonable" (HCP010)

"I wouldn't want it [feedback] to go anywhere else first. Especially if I've listed the patient and I've seen the patient, then it's feedback to me really and I need to explore that further." (HCP013)

There was overall strongly support for a process where feedback is initially provided to someone other than the surgeon. Professional participants frequently discussed the need for care pathways changes whereby introducing a "filter" (HCP011), a "triage kind of mechanism" (HCP010) or "bringing someone in the middle" (HCP003).

"I think having a process delivered by someone who's not the consultant as a first step would be more efficient and reliable." (HCP004)

Several reasons were mentioned why this solution was deemed acceptable. For example, interviewees noted different communication styles or failed doctor-patient relationship building may mean patients want to speak to another staff member. Further, absence or competing priorities may lead to surgeons missing feedback provided to them directly. Participants also felt that other professional roles could often better address issues not requiring clinical input. There was no consensus on the which professional role would be most acceptable to surgeons to take on the responsibility of receiving feedback. Professional participants typically felt that specialist nurses were most appropriate, but also suggested administrative professions (e.g. waiting list coordinators, secretaries) for surgical departments without specialist nurse roles.

"Another one that they might like to talk to a different doctor or a specialist nurse or some other member of staff because they didn't get along with that particular doctor." (HCP008)

"[M]y ideal would be one of your team, or one of my team has spoken to the patient and explored it, and said actually, these were the concerns, because then I can address them." (HCP005)

"[Y]ou could weed out people that have just either made a mistake with the survey and got their scale the wrong way round." (HCP004)

"I wouldn't be averse to having an email alert to say that somebody that's under my care is worried about their surgery [...] because obviously I'd want to act on it. [...] I suppose we need to think about what happens if the person receiving that email isn't around and [...] what the sort of process is to have a backup plan in terms of our maybe admin team. [...] I suppose sort of looking at it in a sort of similar way to how we manage our kind of complaints, is that essentially the default is that they're all for our admin managerial team to look at in the first instance and only come to us with the clinical stuff that can't be dealt with by them." (HCP010)

2.3 Approaches to responding to feedback

Both stakeholder groups frequently talked about the need to refine processes for responding to feedback, exploring various approaches in terms of their acceptability and effectiveness. All participants indicated that a process which provided a follow-up encounter (e.g., a conversation or consultation encounter) would be acceptable to them to address deficiencies in patient-reported SDM experiences. There was emphasis from both participant groups that follow-up encounters should be tailored to the individual and address the specific issue underlying the poor SDM experience.

"I want that opportunity to be able to discuss exactly what's involved and you know if that can happen as a result of this survey then that's great." (PAT003)

"I would assume that the next step would be a follow up consultation to go through and make sure that the things that mattered most were discussed if they didn't feel they had been already." (HCP013)

Most frequently, participants in both stakeholder groups thought that telephone conversations would be an appropriate and *"more convenient"* (HCP004) method for a follow-up encounter. Some interviewees expressed a preference for face-to-face or virtual conversations because *"you can get a lot more information across"* (PAT006), but acknowledged the associated resource implications.

Whilst some patient participants preferred to speak to a surgeon, most felt that additional communication can be with *"anyone from that department really that has the authority to be able to answer those questions"* (PAT003). Most often, patients mentioned nurses or specialist nurses as appropriate roles to address SDM deficiencies.

"I think it's a bit much to expect to speak to the surgeon, I know that's what they're there for, but you don't want to take them away from what they're meant to be doing. But it would have been nice to just have said, 'If you want talk about your op...' even if it had been 10 minutes, just to explain to me about exactly what I'm having done." (PAT017)

"I feel that if I had put that survey score back in, but I still didn't feel that I had the opportunity to discuss why and if somebody had have contacted me immediately and then said look is there anything we can do, it could have even been with the Breast Care Nurse, a meeting to discuss the specific concerns because I think that would have helped me, just even her acting as a conduit between the Consultant who didn't have much time, but it would have only been a short discussion really." (PAT010)

Improving SDM experiences during follow-up encounters was widely viewed as essential to ensure real-time monitoring and feedback is generally perceived as credible and effective. Both participant groups thought that the responsibility for this improvement predominantly lies with healthcare professionals, who should adapt their communication styles. Some patient participants also described the importance of being better prepared to discuss their treatment.

"I think maybe there's somebody available, where you can just say, 'Can I have a quick chat and ask more questions?' because you go to your appointment, you're not really prepared, and maybe I should have had questions, but I didn't at the time" (PAT017)

"I'm trying to understand what parts of the conversation are difficult to understand. Which parts are relatively straightforward to understand, and indeed which parts may need to be presented in a different way?" (HCP009)

Successful engagement of staff in responding to feedback was a key concern mentioned by professional participants. Sharing data that reflects current practices, for example, was identified as important for encouraging their involvement in follow-up encounters.

"I mean it's an extra step isn't it? So I think if you've not engaged everybody into the process then getting people to act on the results might initially be difficult but if there's a consistent trend, like I said, the reality is the data speaks for itself." (HCP002)

3. Impact of real-time monitoring and feedback

3.1 Perceived impact on patients

Positive patient-level impacts of real-time SDM monitoring and feedback were commonly discussed by patient participants and by about half of professional participants. Both participant groups emphasised its value in realising patient-centred care.

"what's important for one person is really irrelevant to somebody else and unless you ask them what's important for them, you don't know" (PAT010)

"I think it's incredibly important, the project [real-time monitoring and feedback of SDM]. I admire it because it is patient focused. Often, we tend to be process focused rather than patient focused, and this will just provide that additional insight into the patients experience that actually can modify things that sometimes we tend to ignore" (HCP009)

Patient participants often felt that monitoring and feedback of SDM would positively impact their general satisfaction and happiness. Some interviewees emphasised how improved SDM may provide reassurance about their treatment decision and thereby decrease their worry about having surgery.

"Just to get some clearer explanation as to you know, what exactly the surgery entails, somebody to listen to my concerns just so that I feel happier overall. [...] I think the impact is you know, reassurance and peace of mind. I would just feel a lot more comfortable knowing what is going to happen to me than going in blindly putting literally my life in the hands of another person." (PAT003).

Similarly, professional participants mentioned increased satisfaction and improved treatment outcomes as potentially positive impacts.

"[I]f the patient feels they've made an active decision with you, they will be much happier with the treatment outcome, whatever that is." (HCP007)

3.2 Perceived impact on surgeons

Potential impacts on surgeons were mentioned by both participants groups.

A few patient participants stated that real-time monitoring and feedback of SDM could help surgeons identify training needs or areas where they could provide better support.

"It will help the surgeons then, they'll think, 'Oh, hang on I've got to help out a bit more,'" (PAT016)

"I think they should learn from it and teach the solution, do you know what I mean? Not my individual solution but I'm sure [there are] courses and procedures where you have to go by what's been said and how to deal with it." (PAT002)

Professionals often discussed positive as well as negative impacts of real-time SDM monitoring and feedback on their own practice. Interviewees also perceived real-time monitoring and feedback of SDM as a crucial part of their professional development and highlighted they would *"be open to learning from that sort of data"* (HCP010). Specifically, the majority felt that receiving information about patients' experience with SDM were opportunities for reflection and improvement in communication skills. Interviewees were interested in personal performance trends and felt there was value in receiving analytical feedback from aggregate scores over time.

"I think that the utility of this tool is in providing aggregate feedback." (HCP011)

"My view is, if you get negative feedback, that's much more important than any positive feedback because somebody's really wanted to say that, if you see what I mean. They've really felt – they've climbed and crossed a threshold in order to actually criticise somebody who's trying to help them. So, from my point of view, certainly that negative side, those themes that might come out on an analysis over a period of time were probably more helpful than the individual cases.[...] it's giving the clinician the tool to examine their general practice so that they can modify and improve matters." (HCP009)

However, challenges with regards to individual surgeon's motivation to change and improve their communication style were also acknowledged.

"I guess you've got to be wanting to improve before with this information, rather than thinking: 'Gosh, well I do a good enough job anyway and I don't care what anybody says'. But there will be people like that but there's nothing you can do about that" (HCP009)

"I'd be interested in knowing is whether they can actually change their communication style to the extent... If you feed this back to a 55 year old surgeon, can that person actually change their style in the last few years of their career, to improve their patient satisfaction? I don't know." (HCP007)

Use of SDM monitoring data for appraisal and revalidation were sometimes discussed by professionals. Participants were divided over whether the data should be used for formal performance evaluation. Some interviewees perceived it as “a much better way of doing it [revalidation]” (HCP008), whereas others were sceptical, expressing concerns about the potential misuse of SDM measurement for performance management and the overlap with existing performance reviews.

“I suppose the only thing is every time a Trust measure something there’s a risk that they’re using it for performance management [...] and is that a bad thing or is it a good thing?” (HCP004)

“I think the difficulty is whether it is going to end up being part of your revalidation I guess is something that might be considered. Yes, I think that might be one way to take it and then I guess to show reflection and if you have changed your style” (HCP012)

“We already get a lot of reviews of our performance from a cancer perspective and it [the national prostate cancer audit] already surveys the quality of information” (HCP002)

Interviewees were often concerned about possible additional burden resulting from receiving negative feedback about patients’ SDM experience. Some expressed worries about increased workload to address feedback which were often discussed in connection with existing performance pressures (e.g. reviews on cancer targets). Others viewed actions on feedback as manageable if these are offset with a reduction in complaints in the long-term.

“I know it’s only a phone call, but it is another phone call – it all adds burden, time” (HCP005)

“The tricky bit is going to be if your patient thinks that they’ve been hard done by and you’ve got to try and meet their cancer target, you’re suddenly going to go ‘how am I going to fit all that in before their operation in a week’s time?’” (HCP003)

“I would feel completely happy with that, assuming I’m not going to get 40. [...] We get feedback in lots of different ways already and again, if it can stop complaints because I think what you are trying to do is almost pre-empt that.” (HCP012)

Professional participants occasionally highlighted emotional burden as a negative consequence of receiving negative feedback.

“I pride myself on my communication and if someone said [...] that I have made no attempt to... and that is pretty awful. Of course I’d be upset about that” (HCP004)

3.3 Perceived impact on departments or the organisation

Organisation or department-level impacts of SDM monitoring and feedback were rarely mentioned by patient participants but were discussed by all professional participants. These commonly included the potential to detect and address wider systemic performance issues.

“Well, it’s making sure that any errors or problems that there might be with the system, or the procedures that are in place, any problems that they might have can be identified and eliminated”. (PAT006)

"I suspect that might be an area where we don't do so well. [...] I think if there are particular themes within our department then we could look to address that" (HCP012)

Professionals further discussed the potential for larger-scale training in connection with the importance of monitoring trends and analysing cumulative data. For around half of the interviewees, the benefit of cohort-level feedback outweighed the benefits of receiving feedback on an individual, patient-level. This extended to the desire to "see how we ranked versus other specialities, other clinicians, you know anonymised" (HCP008).

"It would be incredibly useful to have a sort of annual appraisal style report to allow you to influence practice, and I think it would be extremely helpful to flag up when there are any trends." (HCP005)

Professional participants sometimes pointed out wider impacts such as reputational benefits for the organisation resulting from potentially reduced complaints and litigation, or the value of gathering evidence to inform improvements to hospital processes (e.g., longer clinic times).

"I mean, I think from a hospital perspective and an organisation perspective, maybe the hope is that they might, you know, the great thing about this is it might avoid people complaining." (HCP002)

"Something we talk about fairly regularly is about whether we should be doing a consent clinic for the patients we're operating on in two weeks' time and there is so much pressure on clinic spaces and our time that we haven't done it, you know. I suppose we were surveyed and we were scoring low and additional phone calls weren't solving it then you can say, "Well, actually, we'll formalise this and everyone who's having an operation in the next two weeks is booked, schedule two hours for phone calls for patients next week." (HCP006)

Discussion

This qualitative study explored the acceptability and impact of real-time monitoring and feedback of patients' experiences of SDM measurement. Ten themes were identified describing the perspectives of surgical patients and healthcare professionals in relation to acceptability of real-time SDM monitoring, real-time feedback and addressing SDM deficiencies and impact on clinical practice. Findings suggest that integrating an automated system for real-time SDM measurement and feedback within surgical pathways of NHS Trusts is acceptable. It was endorsed by both groups who also felt it would be valuable to improve the process of care where deficiencies were identified. Results emphasised the complexities of SDM measurement in surgical contexts, specifically regarding the timing of the SDM monitoring, how and what feedback was implemented and actioned within the care pathways, and the differing SDM needs of individuals with various conditions. These complexities might significantly impact acceptability of real-time monitoring and feedback with some issues being of particular importance to patients (e.g., limited inclusivity of real-

time SDM monitoring) and others unique to professionals (e.g., the need for additional qualitative feedback alongside SDM scores). Insights also revealed support for real-time SDM monitoring and feedback because of the potential to improve patient-centred care and associated benefits to patients, surgeons and clinical teams, and the wider organisation. Facilitators and barriers at all levels were identified that can affect adoption of future interventions aimed at improving SDM.

Implications for research

The acceptability and impact of real-time electronic measurement and feedback has been explored in other healthcare context which examined other types of patient-reported data, such as symptoms and health-related quality of life scores [37,38], clinical outcomes [39] or safety [40]. Findings in those studies align with results reported here, demonstrating that clearly presented feedback on patient-reported outcomes and user-friendly technology can enable adoption of similar systems. To our knowledge, however, no previous research has yet investigated stakeholders' views towards integrating, at scale within a healthcare setting, the real-time monitoring and feedback of patients' experiences of SDM. This study provides an important first step to addressing the UK National Institute for Health and Care Excellence call for research into new interventions targeting inclusive, sustained, health system-level SDM improvement [10]. Results add to our understanding of how to efficiently and routinely monitor patients' SDM experiences at scale and it adds insights into factors impacting the acceptability amongst key stakeholders that are necessary to inform intervention development for organisation-wide improvements in SDM.

Insights gained from this study are crucial to inform the development, implementation and evaluation of an SDM improvement intervention in alignment with the principles of the Medical Research Council's framework for the development and evaluation of complex interventions and the Health Technology Assessment framework [27,41]. Factors were identified relevant to optimising the acceptability of a system for real-time SDM monitoring and feedback that can be used to inform the co-design of components of a future SDM intervention that meet the specific needs of key stakeholders. For example, identified themes suggest that the approach to providing feedback of SDM scores to clinical teams will require careful management along the pathway of care. Participants highlighted consequences of potential emotional burden and disengagement in cases of negative feedback. Patient and professional-facing communication may therefore consider strategies linked to positive message framing to optimise engagement with SDM monitoring and feedback.

Findings suggest that the implementation of real-time SDM monitoring and feedback is a service-wide responsibility. Careful consideration should be given to identify appropriate professional roles for receiving and responding feedback and how services should respond. Future work should further explore effective strategies for embedding SDM monitoring into care pathways. Established frameworks from behavioural and implementation sciences (e.g., Theoretical Domains Framework, Normalisation Process Theory [42,43]) could help in developing approaches systematically and transparently which incorporate stakeholder insights and map barriers and facilitators. Additionally, more work is needed to develop effective strategies for stakeholder engagement that can address identified healthcare professional concerns (e.g., worries about time pressures, performance evaluation or scrutiny) to maximise uptake of the future intervention to improve SDM.

Strengths and limitations of this study

Strengths included use of robust qualitative methods which explored and compared perspectives from both patients and healthcare professionals. Investigating perspectives of key stakeholders who had the opportunity to engage with the electronic system obtained new insights about important issues impacting acceptance and implementation of a real-time SDM monitoring and feedback. The study was conducted in a real-world clinical setting and will likely inform wider implementation of the system in UK NHS Trusts.

Limitations include a self-selected sample that lacked diversity. Consideration of ethnicity, diversity and inclusion principles is particularly important in the design and development of digital health interventions [44] and for improving SDM [45]. Recruitment of patient participants from a more diverse of population groups was hampered because hospital electronic patient record systems did not comprehensively recording ethnicity data. Similarly, governance approvals restricted the eligibility criteria to patients who have already completed SDM monitoring. It may be that patient participants in this study were comfortable with using digital technologies and interested in enhancing patient involvement in their care. Likewise, the self-selection of professional interviewees poses a further bias. Healthcare professionals represented three of seven surgical specialties. Recruitment relied on convenience methods and depended on surgeons' availability and willingness to participate in interviews. While participation from all surgeons within departments was actively encouraged, including those who might hold opposing views, it is possible that individuals less interested in improving SDM or those constrained by heavy workloads lacked the motivation or capacity to take part. These limitations may help to explain why themes frequently reflected positive perspectives of the system and might mean additional critical viewpoints were not captured.

The use of digital systems to monitor patient outcomes in surgery has accelerated since the COVID-19 pandemic, but the need to prioritise health equity and research to improve language and digital inclusivity is widely acknowledged [46]. Our related studies have found a range of factors impacting the usability of the system for real-time SDM monitoring and the assessment of SDM in underserved population groups (ethnic minority, elderly and economically disadvantaged individuals) including barriers around accessibility and unwillingness to engage in real-time monitoring [20,21]. This study explored the views of key stakeholders and reports insights gained above and beyond other related work. However, the sample did not include other relevant professions (e.g., NHS managers, administrative staff) important to understanding the implementation of a system for real-time monitoring and feedback. Planned future intervention development work will incorporate views of a wider variety of professional stakeholders and will address relevant factors identified through formative work that may impact acceptability and usability of real-time SDM monitoring and feedback. For example, it is anticipated that non-English versions of the SDM survey will be provided and supported completion of the electronic system (e.g., on tablet computers in clinic, supported by staff/family) will be explored to meet accessibility needs and maximise engagement.

Clinical implications

International health policy emphasises the importance of high-quality SDM processes in patient-centred care [2,5,6]. Results have direct relevance for health service or quality improvement managers seeking to enhance SDM in practice. Implementing real-time monitoring and feedback of SDM addresses a key policy recommendation to track SDM at large-scale [8]. It can therefore be a critical mechanism to guiding new interventions for targeted sustained improvement activities at organisation-level.

Sustained improvements in SDM, however, require complex interventions involving multiple stakeholders at different levels of the organisation [7,11,27,47]. Although the findings identified several factors influencing the acceptability of real-time SDM monitoring and feedback within clinical practice, there are many unknowns as to what components and mechanisms are needed for all stakeholders to ensure its effective adoption and impact on service outcomes and patient benefit. Research to formally develop and evaluate a decision-support intervention using real-time SDM monitoring and feedback is needed to improve evidence base. Feedback components, for example, have been widely used to prompt enhancements professional practice, but the mechanisms through which consistent improvements can be achieved are not well understood [14,19]. Findings highlighted real-time SDM monitoring and feedback provided opportunities for healthcare

professional reflection, potentially prompting practice improvements. This can be crucial for enhancing clinical reasoning that underpins high-quality SDM [48] and may be a key mechanism through which decision support interventions help achieve higher standards of care in this field. Future intervention development work should draw on specific theories (e.g., Clinical Performance Feedback Intervention Theory [19]) to help explain the specific factors that influence feedback success to enhance effectiveness of an intervention that uses real-time SDM monitoring and feedback.

Conclusion

This study explored patient and healthcare professional views towards a novel system for automated, real-time monitoring and feedback of surgical patients' experiences of SDM integrated within elective surgical care in two English NHS Trusts. The system was acceptable to the sampled population and well-positioned to form the basis for future intervention development to improve SDM. Ongoing work addresses the need to understand views of a more diverse sample to ensure the acceptability of SDM to the general UK population. Findings will guide effective strategies for wider implementation of real-time SDM monitoring and feedback in the NHS.

Contributors AGKM developed the original idea for this study along with KNLA and JMB. AGKM, KNLA, CH, RCM, JMB, DH, SH, CC, JH, BG, LR, AW, JA and HLB contributed to the development of the research question and objectives and were involved in the design of the study. KNLA, RCM, CC, LR, JH and HLB provide methodological expertise. CH and AGKM wrote the first draft of the manuscript, and all co-authors reviewed and critically appraised the manuscript. AGKM (guarantor) has overall responsibility for the content and project with strategic oversight from JMB. VS contributed to the PPIE strategy. Collaborators part of the ALPACA Study team (AJ, AS, AL, BR, JP, MRW, PC, PB, SP, TB, TW) provide clinical liaison and subject expertise that have shaped the study design. All collaborators have critically reviewed the manuscript and approved the final version.

References

- 1 NHS England and NHS Improvement. Personalised Care Shared Decision Making Summary guide. 2019.
- 2 National Health Service. The NHS Long Term Plan. Published Online First: 2019.
- 3 Légaré F, Adekpedjou R, Stacey D, *et al.* Interventions for increasing the use of shared decision making by healthcare professionals. *Cochrane Database Syst Rev.* 2018;7:CD006732. doi: 10.1002/14651858.CD006732.PUB4
- 4 de Mik SML, Stubenrouch FE, Balm R, *et al.* Systematic review of shared decision-making in surgery. *British Journal of Surgery.* 2018;105:1721–30. doi: 10.1002/bjs.11009
- 5 Légaré F, Stacey D, Forest PG, *et al.* Milestones, barriers and beacons: Shared decision making in Canada inches ahead. *Z Evid Fortbild Qual Gesundheitswes.* 2017;123–124:23–7. doi: 10.1016/J.ZEFQ.2017.05.020
- 6 Australian Commission on Safety and Quality in Health Care. National Safety and Quality Health Service Standards (second edition). 2021.
- 7 Carmona C, Crutwell J, Burnham M, *et al.* Shared decision-making: summary of NICE guidance. *BMJ.* 2021;373. doi: 10.1136/BMJ.N1430
- 8 National Institute for Health and Care Excellence. Shared decision making [B] Evidence review for interventions to support effective shared decision making NICE guideline NG197 Evidence reviews underpinning recommendations 1.2.1 to 1.2.18 and research recommendations in the NICE guideline. 2021.
- 9 NHS England. Commissioning for Quality and Innovation (CQUIN): 2022/23. 2022.
- 10 National Institute for Health and Care Excellence. Shared decision making [NG197]. Evidence reviews 2021. NICE 2021.
- 11 Joseph-Williams N, Lloyd A, Edwards A, *et al.* Implementing shared decision making in the NHS: lessons from the MAGIC programme. *BMJ.* 2017;357:j1744. doi: 10.1136/bmj.j1744
- 12 NHS Rightcare. Your Health – Your Decision. London 2013.
- 13 NHS Rightcare. Measuring Shared Decision Making: A review of research evidence - A report for the Shared Decision Making programme in partnership with Capita Group Plc. London 2012.
- 14 Ivers N, Jamtvedt G, Flottorp S, *et al.* Audit and feedback: Effects on professional practice and healthcare outcomes. *Cochrane Database of Systematic Reviews.* 2012;2012. doi: 10.1002/14651858.CD000259.pub3
- 15 Lee R, Baeza JI, Fulop NJ. The use of patient feedback by hospital boards of directors: a qualitative study of two NHS hospitals in England. *BMJ Qual Saf.* 2018;27:103–9. doi: 10.1136/BMJQS-2016-006312
- 16 NIHR Dissemination Centre. Improving Care by Using Patient Feedback. *Health and Social Care Services Research.* Published Online First: December 2019. doi: 10.3310/THEMEDREVIEW-04237

- 17 Sheard L, Peacock R, Marsh C, *et al.* What's the problem with patient experience feedback? A macro and micro understanding, based on findings from a three-site UK qualitative study. *Health Expectations*. 2019;22:46–53. doi: 10.1111/HEX.12829
- 18 Flott KM, Graham C, Darzi A, *et al.* Can we use patient-reported feedback to drive change? the challenges of using patient-reported feedback and how they might be addressed. *BMJ Qual Saf*. 2017;26:502–7. doi: 10.1136/BMJQS-2016-005223
- 19 Brown B, Gude WT, Blakeman T, *et al.* Clinical Performance Feedback Intervention Theory (CP-FIT): A new theory for designing, implementing, and evaluating feedback in health care based on a systematic review and meta-synthesis of qualitative research. *Implementation Science*. 2019;14:1–25. doi: 10.1186/S13012-019-0883-5/TABLES/10
- 20 Hoffmann C, Avery KNL, Macefield RC, *et al.* Real-time monitoring and feedback to improve shared decision-making for surgery (the ALPACA Study): protocol for a mixed-methods study to inform co-development of an inclusive intervention. *BMJ Open*. 2024;14. doi: 10.1136/BMJOPEN-2023-079155
- 21 Hoffmann C, Avery K, Macefield R, *et al.* Usability of an Automated System for Real-Time Monitoring of Shared Decision-Making for Surgery: Mixed Methods Evaluation. *JMIR Hum Factors*. 2024;11:e46698. doi: 10.2196/46698
- 22 Hughes TM, Merath K, Chen Q, *et al.* Association of shared decision-making on patient-reported health outcomes and healthcare utilization. *The American Journal of Surgery*. 2018;216:7–12. doi: 10.1016/J.AMJSURG.2018.01.011
- 23 Aubree Shay L, Lafata JE. Where is the evidence? a systematic review of shared decision making and patient outcomes. *Medical Decision Making*. 2015;35:114–31. doi: 10.1177/0272989X14551638
- 24 Avery KNL, Richards HS, Portal A, *et al.* Developing a real-time electronic symptom monitoring system for patients after discharge following cancer-related surgery. *BMC Cancer*. 2019;19:1–15. doi: 10.1186/S12885-019-5657-6/TABLES/6
- 25 Richards HS, Blazeby JM, Portal A, *et al.* A real-time electronic symptom monitoring system for patients after discharge following surgery: A pilot study in cancer-related surgery. *BMC Cancer*. 2020;20:1–17. doi: 10.1186/S12885-020-07027-5/FIGURES/4
- 26 Warrington L, Absolom K, Holch P, *et al.* Online tool for monitoring adverse events in patients with cancer during treatment (eRAPID): field testing in a clinical setting. *BMJ Open*. 2019;9:e025185. doi: 10.1136/BMJOPEN-2018-025185
- 27 Skivington K, Matthews L, Simpson SA, *et al.* A new framework for developing and evaluating complex interventions: update of Medical Research Council guidance. *BMJ*. 2021;374:n2061. doi: 10.1136/BMJ.N2061
- 28 Cresswell J. *Qualitative inquiry & research design: Choosing among five approaches*. 2013.
- 29 Braun V, Clarke V. Conceptual and Design Thinking for Thematic Analysis. *Qualitative Psychology*. 2021;9:3–26. doi: 10.1037/qup0000196
- 30 Tong A, Sainsbury P, Craig J. Consolidated criteria for reporting qualitative research (COREQ): A 32-item checklist for interviews and focus groups. *International Journal for Quality in Health Care*. 2007;19:349–57. doi: 10.1093/intqhc/mzm042

- 31 Hennink MM, Kaiser BN, Marconi VC. Code Saturation Versus Meaning Saturation: How Many Interviews Are Enough? *Qual Health Res.* 2017;27:591–608. doi: 10.1177/1049732316665344
- 32 Saunders B, Sim J, Kingstone T, *et al.* Saturation in qualitative research: exploring its conceptualization and operationalization. *Qual Quant.* 2018;52:1893. doi: 10.1007/S11135-017-0574-8
- 33 Nelson J. Using conceptual depth criteria: addressing the challenge of reaching saturation in qualitative research: <https://doi.org/10.1177/1468794116679873>. 2016;17:554–70. doi: 10.1177/1468794116679873
- 34 Bingham AJ. From Data Management to Actionable Findings: A Five-Phase Process of Qualitative Data Analysis. *Int J Qual Methods.* 2023;22. doi: 10.1177/16094069231183620/ASSET/IMAGES/LARGE/10.1177_16094069231183620-FIG11.JPEG
- 35 Brooks J, Mccluskey S, Turley E, *et al.* The Utility of Template Analysis in Qualitative Psychology Research. *Qual Res Psychol.* 2015;12:202–22. doi: 10.1080/14780887.2014.955224
- 36 Glaser BG, Strauss AL. *The Discovery of Grounded Theory: Strategies for Qualitative Research.* Mill Valley, CA: Sociology Press 1967.
- 37 van den Hurk CJG, Mols F, Eicher M, *et al.* A Narrative Review on the Collection and Use of Electronic Patient-Reported Outcomes in Cancer Survivorship Care with Emphasis on Symptom Monitoring. *Current Oncology 2022, Vol 29, Pages 4370-4385.* 2022;29:4370–85. doi: 10.3390/CURRONCOL29060349
- 38 Richards HS, Blazeby JM, Portal A, *et al.* A real-time electronic symptom monitoring system for patients after discharge following surgery: A pilot study in cancer-related surgery. *BMC Cancer.* 2020;20:1–17. doi: 10.1186/S12885-020-07027-5/FIGURES/4
- 39 Pickens R, Cochran A, Tezber K, *et al.* Using a Mobile Application for Real-Time Collection of Patient-Reported Outcomes in Hepatopancreatobiliary Surgery within an ERAS® Pathway. <https://doi.org/10.1177/000313481908500847>. 2019;85:909–17. doi: 10.1177/000313481908500847
- 40 Hernan AL, Kloot K, Giles SJ, *et al.* Investigating the feasibility of a patient feedback tool to improve safety in Australian primary care: a study protocol. *BMJ Open.* 2019;9:27327. doi: 10.1136/bmjopen-2018-027327
- 41 National Institute for Health and Care Research. Health technology assessment 2020. 2020.
- 42 Murray E, Treweek S, Pope C, *et al.* Normalisation process theory: A framework for developing, evaluating and implementing complex interventions. *BMC Med.* 2010;8:1–11. doi: 10.1186/1741-7015-8-63/TABLES/3
- 43 Francis JJ, O'Connor D, Curran J. Theories of behaviour change synthesised into a set of theoretical groupings: introducing a thematic series on the theoretical domains framework. *Implementation Science.* 2012;7:1–9. doi: 10.1186/1748-5908-7-35/TABLES/2
- 44 Thomson A, Gooberman-Hill R. *How can EDI be nurtured through communities and individuals?* Policy Press 2024.

- 45 Mead EL, Doorenbos AZ, Javid SH, *et al.* Shared decision-making for cancer care among racial and ethnic minorities: a systematic review. *Am J Public Health.* 2013;103. doi: 10.2105/AJPH.2013.301631
- 46 Fong Y, Melstrom L, Evans HL. Practical Guide to Remote Patient Monitoring in Surgical Patients. *JAMA Surg.* Published Online First: 8 January 2025. doi: 10.1001/JAMASURG.2024.6015
- 47 Bekker HL, Winterbottom AE, Gavaruzzi T, *et al.* Decision aids to assist patients and professionals in choosing the right treatment for kidney failure. *Clin Kidney J.* 2023;16:i20–38. doi: 10.1093/CKJ/SFAD172
- 48 Holm A, Rodkjær LØ, Bekker HL. Integrating Patient Involvement Interventions within Clinical Practice: A Mixed-Methods Study of Health Care Professional Reasoning. *MDM Policy Pract.* 2024;9. doi: 10.1177/23814683241229987/ASSET/IMAGES/LARGE/10.1177_23814683241229987-FIG4.JPEG