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Coordination of procedural equipment and supplies for the surgical set-up in the perioperative environment: A scoping review

Abstract

Background: Defective, incorrect or missing procedural devices from the surgical set-up contribute to delay, interruption, cancellation and patient harm in the perioperative environment.

Objective: This scoping review aims to identify evidence to guide approaches to surgical set-up used by perioperative health service personnel, organisations or teams. In addition, the review aims to describe factors that hinder or support the surgical set-up, identify gaps in the literature and determine any issues impacting the quality of available evidence.

Methods: Empirical research and grey literature were retrieved from seven electronic databases. Titles and abstracts were screened before full text screening. A mixed method appraisal tool (MMAT) and quality improvement minimum quality criteria set (QI-MQCS) were used for critical appraisal. After data extraction from included studies, key concepts were synthesised, thematically analysed and reported.

Results: Forty-nine full texts were included. Evidence generated by nurses responsible for the surgical set-up is limited. The majority of studies were quality improvement studies to reduce inefficiencies through optimisation or mathematical modelling with outcomes measured in cost and time saved. There is limited evidence exploring how optimisation or mathematical modelling impacts the work of perioperative staff.

Conclusion: Technology will continue to influence work systems and processes of the surgical set-up. Implementing surgical set-up quality indicators within policy may aid waste and cost reduction of organisations. The impact of human factors upon the surgical set-up is relatively unaddressed. Nurse-led research on the surgical set-up would be valuable as nurses are key professionals contributing to delivery of, management of and policy about surgical set up.

Introduction

The effective, safe and timely management of surgical devices is fundamental to patient outcomes. Internationally, evidence suggests problems with surgical set-up processes contribute to delay, interruption or cancellation of surgery¹⁻³. Problems include inadequate information regarding surgical supplies, waste from unused opened devices and superfluous, defective, incorrect or missing surgical equipment^{2, 4-6}.

A surgical set-up can be a dynamic, labour-intensive process fraught with complex, time sensitive challenges in a technological environment with evolving procedural techniques⁷⁻⁹. Many staff working at different times and locations contribute to surgical set-ups; these staff include technicians, medical device representatives and nurses. Confusion about equipment and procedural information has been reported with perioperative nurses being 'busy locating equipment' at the beginning of surgical lists^{4, p.3}. For example, an observational study by Rappold et al.¹⁰ in the United States of America (USA) recorded more than 4000 surgeon preference cards were unused, contributing to ineffective procurement, unused opened devices and superfluous instruments. Evidence regarding how to best approach and organise surgical set-up processes for perioperative personnel, organisations and teams would be valuable.

The aim of this review was to examine the availability of evidence to guide the surgical set-up. Primary scholarly literature was reviewed to identify and map available evidence and describe factors that hinder or support the surgical set-up. The

review also aimed to identify gaps in the literature regarding the surgical set-up and to determine issues impacting the quality of available evidence.

Methods

A scoping review guided by Joanna Briggs Institute (JBI) methodology¹¹ was conducted and is reported according to the PRISMA-ScR (preferred reporting items for systematic reviews and meta-analyses, extension for scoping reviews)¹². The JBI framework of population, concept and context (PCC)¹³ was used with key terms defined as:

- **population** – health service personnel, organisations, groups or teams responsible for the surgical set up
- **concept** – the surgical set-up which involves timely, coordinated organisation of single-use and re-usable medical devices (RMD), biomaterials and ancillary equipment. A set-up, or case assembly, is defined as assembly of physical resources needed for a procedure and may include opening and laying out surgical set-up items within the procedural room¹⁴. This includes surgical instruments, single-use isolation drapes, implants and ancillary medical equipment such as laparoscopic carbon dioxide insufflation devices¹⁵.
- **context** – the perioperative environment. The Australasian Health Facility Guidelines¹⁴ identify the perioperative environment to be an environmentally controlled area with one or more operating rooms to support patient procedural interventions under inhalation or other anaesthetic agents.

Types of evidence

Primary studies including randomised and non-randomised controlled trials, quality improvement projects and case, case-controlled, observational and cohort studies were eligible for inclusion. Literature reviews or discussion papers were excluded. Studies focused on testing safety and efficacy of surgical devices for patient outcomes, such as trials of new surgical devices were also excluded.

Search strategy

A three-step search strategy included an initial search of Cumulative Index for Allied Health Literature (CINAHL) and Scopus identifying medical subject headings (MeSh) for key terms within titles and abstracts^{16,17}. Seven electronic databases were subsequently searched using MeSh terms: CINAHL, Joanna Briggs Institute EPD (via OPD), Scopus, PubMed, Cochrane Central Register of Controlled Trials (CENTRAL). Grey literature was sought via Overton and ProQuest Dissertations and Theses Global (PQDT)TM. The search strategy used for Joanna Briggs Institute EPD database is presented as Supplement 1. With a lack of access to translators, only papers in English were included. The publication timeframe was from database inception to 25 March 2023 to permit capture of trends over time. Reference lists from included sources were examined for additional relevant literature. A PRISMA-ScR flowchart is presented in Figure 1.

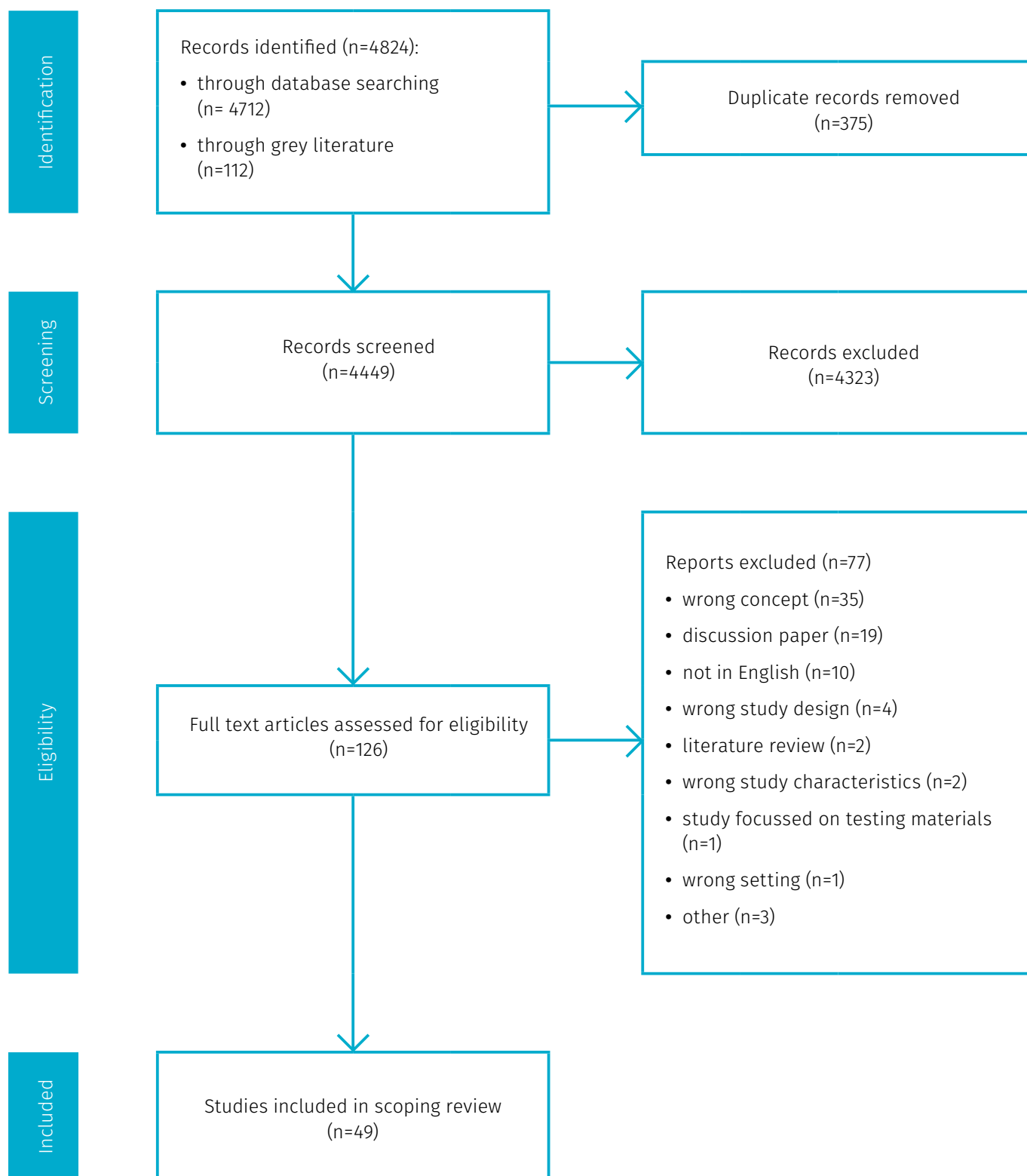


Figure 1: PRISMA flow diagram of paper selection process

Selection of evidence

Piloting of the eligibility criteria was undertaken by three reviewers (ML, JD, JM) screening three full texts followed by discussion (see Supplement 2). The eligibility criteria were rephrased for clarity prior to screening. Search results were imported to EndnoteTM and duplicates removed, then into CovidenceTM for review. Titles and abstracts were screened against the eligibility criteria by three reviewers (ML, JD, JM), then full texts were screened for eligibility by two independent reviewers (ML, JM). Disagreements were resolved through consensus. Reasons for exclusion are summarised in Figure 1.

Data charting process

An adapted JBI data extraction instrument (Supplement 3) was developed and pilot tested. Data was extracted independently from the aims of each study, and included the population, concept, context, type of evidence, citation, participants, country of origin and approaches used for the surgical set up. Factors that hinder or support a surgical set-up were also extracted from the results of each paper.

Critical appraisal

Critical appraisal was undertaken using the mixed method appraisal tool (MMAT)¹⁸ and quality improvement minimum quality criteria set (QI-MQCS)¹⁹ relevant to the study design. Studies were evaluated by methodology to identify trends and strengths or weaknesses.

Synthesis of results

Extracted data was synthesised into narrative and tabulated results addressing the population, concept and context outlined above. Approaches to the surgical set-up were mapped with key themes identified and narratively summarised. Factors that hinder or support the surgical set-up were thematically analysed and classified.

Results

Forty-nine papers are included in this scoping review²⁰⁻⁶⁸. Most studies were conducted in the United States of America (USA)^{20-22,25-27,30-32,34-44,47,48,50,56,57,60-62,64,65,67} (n = 31). The remainder were conducted in Europe^{24,45,46,51-54,58,66} (n = 9), Singapore^{29,33,63} (n = 3), Brazil^{49,68}

(n = 2), Canada^{23,28} (n = 2), Australia⁵⁵ (n = 1) and Australia and Brazil binationally⁵⁹ (n = 1). Included studies were published over 35 years from 1986 to 2023. From 2005 the number of publications increased, with a sharp rise from 2015 (see Figure 2).

Characteristics of included studies

Supplement 4 summarises the characteristics of the included studies. Over half of included papers were quality improvement projects focused on waste minimisation²⁰⁻⁴⁵ (54%, n = 26). Of these, more than three quarters aimed to eliminate inefficiencies, reduce costs and comply with the Patient Protection and Affordable Care Act⁶⁹ in the USA^{20-22,25-27,30-32,34-44} (77%, 20/26).

Four mixed methods studies explored hazards or work systems responsible for re-usable medical devices, often within a human factors or failure effects model⁴⁶⁻⁴⁹. One mixed method study examined how physician preference card planning and communication influenced unplanned costs⁵⁰. Nine observational studies sought to evaluate resource inefficiencies⁵¹⁻⁵⁹. Four observational studies modelled the optimal number of resources needed

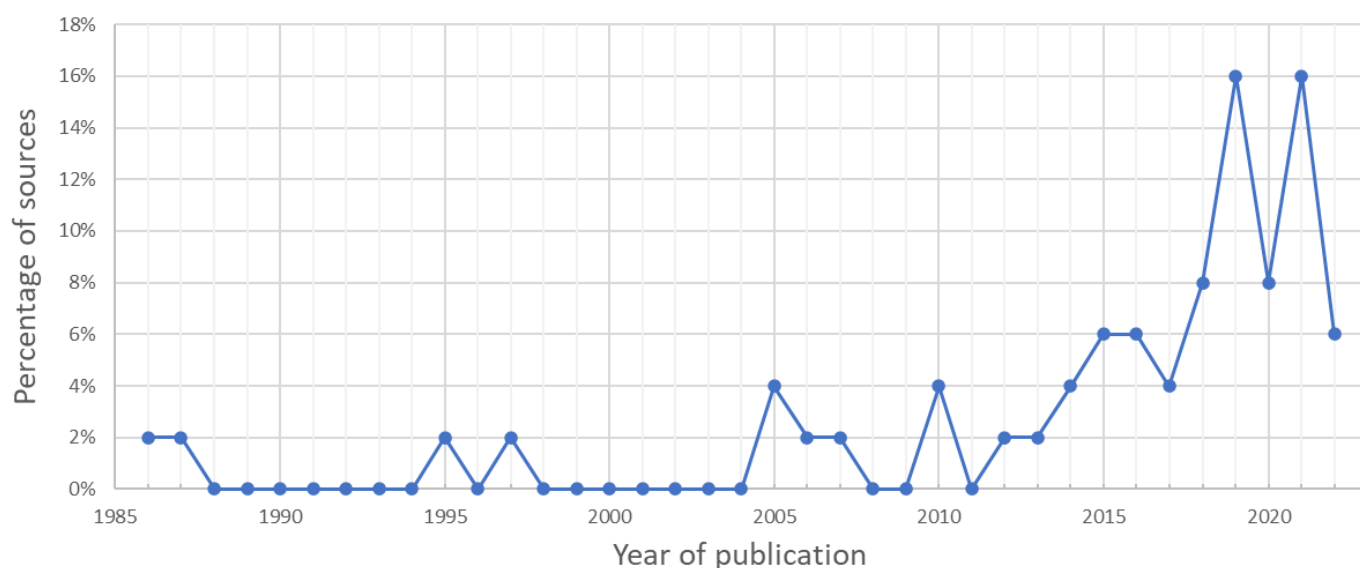


Figure 2: Distribution of published sources 1986 to 2022

to deliver surgical services⁶⁰⁻⁶³. Of three experimental studies, one compared costs between streamlined procedural and standard operating room packs⁶⁴, one compared sterility for procedural packs transported between hospital sites⁶⁵ and one analysed instrument descriptions used by nurses⁶⁶. One qualitative study explained organisational strategies for influencing stakeholders involved in medical device procurement⁶⁷. One case study mapped perioperative flow of instruments⁶⁸.

Factors that hinder the surgical set-up

Factors that hinder the surgical set-up are multidimensional, occurring at different times and locations throughout procedural departments. Three themes identified were waste, lack of governance and human factors.

Waste

The included studies focused on three sources of waste: medical device defects, unused opened medical devices and inefficient use of time.

Medical device defects

Eighteen types of defects were identified across seven studies^{34,37,40,47,48,55,57}. The defects were classified as either sterile or non-sterile (see Table 2), according to the classification used by Palo et al.³⁷ where sterile defects are any problem compromising the sterile integrity of an RMD, and non-sterile defects are any problem influencing the accuracy, functionality or availability of an RMD.

Sterile defects were less frequent than non-sterile defects; the incidence of sterile defects ranged from two per cent²¹ to six per cent³⁷ while the incidence of non-sterile defects ranged from 10.9 per cent⁴⁸ to 52.0 per cent³⁷.

Missing instruments was the most problematic non-sterile defect, with incidence ranging from 17.6 per cent⁴⁸ to 77 per cent³⁷. One observational study⁵⁷ reported that the incidence of missing, broken or unplanned instruments or tray errors was higher (49%) when trays had over 40 instruments compared to when trays had less than 40 instruments (13%).

Unused opened medical devices

In a study of 23 commonly used orthopaedic instrument trays, Cichos et al.²² reported low instrument utilisation resulting in waste – 23 per cent (n = 182/792) of all opened RMD were used. Across the studies, the incidence of unused opened RMDs for total knee arthroplasties varied from 13.0 per cent⁵⁷ to 54.5 per cent³² (n = 47/87). Harris⁶² reported that 70748 instruments were opened and not used annually in a level three trauma centre with eight procedural rooms servicing 6000 procedures. A quality improvement project by Levine³¹ found unused opened medical devices also included prosthetics, with 400 unused opened orthopaedic implants resulting in

\$425 000 lost over three years. An observational study by Chasseigne et al.⁵⁴ identified nurses’ perceptions about why medical devices in the operating room were opened and unused; reasons included anticipation of surgeon needs (33%, 52/152), wrong choice or unsuitable supplies (20%, n=30/152) and aseptic mistakes (18%, n=27/152).

Inefficient use of time

A work sampling study by Ikuma⁵⁶ reported that, for 12 knee arthroplasties observed, 68 per cent (124/182 minutes) of surgical time was dedicated to preparing instruments, preparing the operating room and clean-up, compared to 54 per cent (100/182 minutes) dedicated to performing the procedure. However, authors noted the researcher was not always present when instrument preparation commenced, so instrument preparation time may be longer than reported⁵⁶. An observational study by Chasseigne et al.⁵⁴ identified unintentional absence of the circulating nurse for up to one quarter of procedural time. Reasons for absences included

Table 2: Sterile and non-sterile re-usable medical device defects^{34,37,40,47,48,55,57}

Sterile defects	Non-sterile defects
bioburden (microscopic or foreign body)	broken
contamination	damaged
instrument not disassembled	expired
missing chemical indicator	incorrect
non-bioburden debris (e.g. pen)	incorrect device pulled for set-up
	malfunctioning
	mislabelled
	mismatched instrument/set
	misplaced
	missing
	paperwork/turnover issue
	wrong storage location

additional surgeon demands (30%, n = 16/53), surgical set-up incompleteness (25%, n = 13/53), new supplies required (23%, n = 12/53), defects (19%, n = 10/53) and implant size error (4%, n = 2/53). Of 49 procedures observed by Stockert and Langerman⁵⁷ the surgeon was idle during non-operative time for 29 per cent of procedures (n = 14) due to instrument errors, with each interruption lasting eight minutes on average.

Lack of governance

The included studies highlighted a lack of governance for the surgical set-up. A health care failure model and effects analysis at two hospital sites by Guédon et al.⁴⁶ reported up to 172 hazards in the delivery of loaned orthopaedic instruments. One quarter of hazards (26%, n = 41/158) were not managed; rather, organisations reportedly accepted that adverse events may occur, with up to 31 per cent (n = 49/158) deemed high risk.⁴⁶ High risk hazards included incomplete pre-operative information in digital planning systems.⁴⁶ Only one per cent (n = 1/172) to five per cent (n = 8/158) of hazards were controlled in the delivery of loaned orthopaedic instruments across both hospitals.⁴⁶

A cross-sectional study undertaken in Australia and Brazil by Tripple et al.⁵⁹ identified loaned devices did not conform to a recommended arrival time of 48 hours prior to surgery due to high loan turnover among health services, with approximately 63 per cent (n = 141/221) of loan devices arriving less than 24 hours prior to surgery. Alfred et al.⁴⁸ identified that the absence of instrument descriptions and photographs during sterile reprocessing resulted in incorrect or omitted instruments from trays. A quality improvement project by Prephan⁴⁰ identified instrument

availability was reduced in the absence of repair and maintenance schedules.

Four studies reported routine purchasing, with no systematic data analysis to inform decision-making, encouraged excess quantities and wastage from expiration or obsolescence^{22–24, 42}. Similarly, Levine et al.³¹ found no records of inventory for orthopaedic implants, with unused opened implants costing \$25 000 a month. A quality improvement project to standardise surgeon pick lists by Simon et al.⁴¹ found duplicated products: five comparable laparoscopic clip appliers were stocked from three manufacturers, despite no clear clinical benefit of similar products. Del Carmen et al.²⁴ identified the need to address items being out of stock, stock mismatch and urgent restocking using technological inventory systems. A study modelling surgical instrument distribution for ad hoc orders⁶³ found that even when inventory systems were available, pre-procedural time constraints inhibited the documentation of last-minute device changes.

Human factors

Various human factors were observed to influence the surgical set-up, with themes of unaddressed communication issues and ineffective collaboration. A quality improvement project to improve instrument availability⁴⁰ identified skilled labour shortages coupled with inadequate orientation led to performance deficits for sterilisation technicians. A hazard analysis for delivery of orthopaedic loaned devices by Guédon et al.⁴⁶ found instruments were occasionally double booked suggesting a lack of multidisciplinary communication.

Two studies^{48,66} reported that intra-operative comprehension of instruments decreased when nurses were temporarily assigned or unfamiliar with the surgery, or when one instrument had multiple names. Nonetheless, an observational study by Chasseigne et al.⁵⁴ reported nurses occasionally opened medical devices out of 'comfort' rather than patient need (12%, n=18/152). A quality improvement study by Nilsen³⁶ to determine appropriate operating theatre inventory identified that low surgical device supply generated employee stress, with staff hiding surgical cameras for fear of not having the device ready. A vicious cycle of camera unavailability persisted with impact on patients re-scheduled to an earlier start, although the exact impact was not clearly defined.

Factors that support the surgical set-up

The included studies primarily focused on optimisation – increasing procedural efficiency and reducing cost through standardisation, patient matched devices and eliminating unused medical devices^{32,41,51}. One study reported initiatives to support technicians responsible for the surgical set-up included training for bioburden inspection, testing device functionality, instrument tray completeness and sterilisation processes⁴⁸. There were no initiatives supporting professional development of perioperative nurses.

Optimisation

Twenty-one studies focused on optimisation of medical device use through eliminating unused devices, patient matched devices or standardisation, primarily in orthopaedics^{20,22,32,43,45,51,52} (n = 7), otolaryngology^{23,28,38,42,57} (n = 5) and various other specialities^{25–27, 30, 39, 41, 44, 49, 64} (n = 9). Interventions included

reducing the volume^{20–23,25,27,28,32,38,39,42–45,49,51,52} (n=17) and weight^{20–22,25,27,38} (n = 6) of devices on trays, with outcomes measured in time^{20,21,23,25,27,28,32,35,41,42,44,45,51,52} (n = 14) and costs saved^{20–22,25–28,30,32,38,39,41,42,49,51,52,64} (n = 17).

Eliminating unused devices and patient matched devices

Substantial cost savings were often achieved through eliminating unused instruments. For example, a 30 per cent reduction (n = 31616/106959) in unused opened medical devices for total knee arthroplasty saved on average USD\$191434 (\$18653–\$364216) annually³². An observational cohort study⁵¹ estimating the economic value of patient matched instrumentation saved 20 minutes per knee arthroplasty, or 7000 minutes annually, thereby increasing service capacity.

Dreyfus et al.⁵⁰ observed a curvilinear relationship between planning items needed for surgery and unplanned costs. Over two years, revisions to physician preference cards initially increased unplanned costs; however, unplanned costs dramatically fell after the sixth revision of physician preference cards⁵⁰. A \$5.83 billion waste reduction was achieved in this same study when physician preference cards were revised nine times over two years, with cost savings plateauing at 11 preference card revisions over the same timeframe⁵⁰.

Less frequently, studies assessed staff satisfaction when instruments were reduced or eliminated^{28,42,44,45,54}. Through surveys, Wannemuehler et al.⁴² identified that most scrub nurses (93.75%, n = 16) expressed satisfaction with the reduction of adenotonsillectomy instruments and, as a result, no longer needed to search through dozens of unused devices on instrument trays⁴². In their

study of optimised otolaryngology surgical trays, Fu et al.²⁸ reported that eleven (92%) participants achieved enhanced set-up efficiency without impacting education, patient safety or operating time. An optimisation pre-post satisfaction survey by Toor et al.⁴⁴ identified that the percentage of staff members who reported that 'inventory configuration is unacceptable, and I am significantly concerned that it can affect clinical operations' fell from 48 per cent (n = 29/60) before optimisation to 3.3 per cent (n = 2/60) after optimisation^{44, p.6}. Staff satisfaction surveys were conducted as part of larger studies conducted by Howard⁶⁴ and Capra et al.²⁰ but no results were reported.

Chasseigne et al.⁵⁴ found that waste prevention could be improved through effective communication between surgeons, instrument nurses and circulating nurses at the beginning of and during a procedure, followed by knowledge of surgical techniques.

Standardisation

Six studies explored medical device standardisation, with joint cost savings for hospitals and surgeons, in addition to vendor competitive bargaining^{26,29,31,37,41,67}. Montgomery and Schneller's qualitative study⁶⁷ of physician behaviour and countering suppliers' power in purchasing devices defined models of standardisation, with methods and mechanisms to achieve standardisation. A quality improvement study by Goh et al.²⁹, focussed on instrument management within the sterile stock unit, found eliminating different vendors offering the same products decreased variability and duplication, resulting in a reduction from 75 general surgery sets to 45, saving S\$64 000 per year while maintaining timely supply for surgery.

Staff professional development

Six studies implemented professional development opportunities for technicians responsible for the surgical set-up^{29,33,35,37,48,63}. Strategies included preceptorship, training, orientation, formal education and in-service education^{29,33,48}. Palo et al.³⁷ found technician cross-rotation, orientation and competency assessments aided reduction of non-sterile defects by 56 per cent (46.8 to 26.5 defects per 1000 cases). Staff redistribution informed by workload analysis as reported in a study by Lum et al.³³ reduced reprocessing time by five per cent (267 min/day from 89 procedures) and sterile stock room replenishment time by 29 per cent (254 minutes to 180 minutes).

Job redesign included reassignment of tasks – including delivery of instruments to operating rooms, packing, storing, decontamination and sterilisation – from nurses to technicians^{33,70}. Task reassignment was proposed to enable nurses to spend more time with patients in the operating room^{33,63}. Ngu³⁵ used weekly meetings to aid pre-operative planning for assigning preference cards, implants and medical devices to surgical cases. Goh et al.²⁹ found that supporting staff through successful implementation of instrument management systems increased workplace safety.

Critical appraisal of literature

Supplement 5 summarises critical appraisal of studies. Weaknesses apparent in observational studies^{51–57,60–62,70} (n = 13) included limited use of reporting guidelines, unclear study design and unknown risk of non-response bias (limited response or dropout rates, and reporting of reasons for non-participation). Only six of the 26

quality improvement projects were reported according to the SQUIRE guidelines^{20,23,28,37,39,71}. Patient health-related outcomes among quality improvement projects were rarely measured, despite four studies^{26,27,31,32} describing patient safety and quality as a priority.

Gaps in evidence

This scoping review identified a lack of available evidence from the perspective of perioperative nurses despite their being key professionals responsible for the surgical set-up. Studies primarily reported attempts to reduce medical device waste through optimisation or mathematical modelling to support efficiency and cost reduction^{20,25,51,60}; there was limited evaluation of impact on the perioperative environment, personnel responsible for the surgical set-up (including registered nurses) and patient outcomes. No studies examined organisational behaviours of perioperative team members responsible for the surgical set-up.

Discussion

This scoping review explored available evidence focused on the surgical set-up. Most included studies were organisational quality improvement projects, with outcomes of procedural efficiency measured by time and cost savings. Strategies to optimise procedural devices include elimination, standardisation and customised patient devices^{72–74}. Enhancing efficiency also included mathematical modelling to predict how many people or devices are needed for surgery^{60–62,75}. The review revealed the scarcity of primary research studies focusing on outcomes related to the surgical set-up, such as patient outcomes. The volume of quality improvement projects versus the lack of primary

research identifies research opportunities, particularly from the perspective of intra-operative nursing as these specialties perform key roles in surgical set up processes^{5,76}.

An increasing volume of papers from 2005 onwards focused on waste management. This may reflect the importance of surgical set-up problems or the improvement in access to data over time, with the introduction of advanced tracking and monitoring systems^{77,78}. As the complexity and diversity of procedural care evolves, solutions involving automation are increasingly common in health services. Despite numerous benefits, technology in the perioperative environment is known to negatively influence workflow⁷⁹. Impacts to workflow include additional job demands for nurses who are also expected to be abreast of technology and troubleshooting^{79–81}.

The increase in technology and specialised procedures, for example patient positioning during robotic surgery, has transformed routine nursing care into a highly technical, complex and arduous responsibility⁸¹. Mastery of surgical set-up technology is stressful and can adversely impact the health, well-being and professional efficacy of nurses^{81,82}, and this impact is worthy of consideration by management. As technology and artificial intelligence continue to evolve, exploring how technology influences the work involved in a surgical set-up will require ongoing investigation as well as policy and practice reform.

A number of studies in this review implemented professional development for sterilisation technicians about the pre- and post-procedural phase^{33,37,47,48,59}. However, there was limited focus on education for intra-operative nurses and

other perioperative professionals. Evidence-based educational approaches are crucial for patient care and safety. Intra-operative nurses learning new technologies ‘on the job’ and during real time surgery is reported to cause nurses to experience fear and anxiety about harming the patient⁸².

Schuessler et al.⁸¹ recommend universally standardised training and certification for professionals involved in robotic surgeries, rather than the duration and content of education being determined by individual hospitals resulting in education of varying quality. Evidence-based methods of teaching and learning for perioperative nurses include a range of self-directed online training, high fidelity simulation, team-focused training and practice operations involving animal cadavers^{83,84}.

As evidenced by the focus on waste found in this scoping review, governance of the surgical set-up simply cannot keep pace with technology. The variation in physician experience and skill that influences device preferences combined with unpredictability of procedures makes it difficult to create and standardise protocols¹⁰. Effective governance is also made more challenging by fiscal and time constraints¹⁰. Subsequently, perioperative departments harbour excessive, outdated and obsolete medical devices with limited systematic organisation, and this results in waste.

A lack of governance may also be influenced by perioperative efficiency measures, such as theatre utilisation representing patient intra-operative time⁸⁵. It is unclear if theatre utilisation metrics are reliable or useful to nurse managers, given that a number of quality improvement projects included in

this review attempted to reduce intra-operative waste. The findings from this review suggest that there is a lack of efficiency measures that reflect contemporary intra-operative challenges for nursing.

Incorporating quality indicators within health service policy may be the first step in aiding governance reform for the surgical set-up. By doing so, health services can effectively streamline surgical set-up processes and optimise resources to reduce waste and costs. Examples of quality indicators for the surgical set-up include the availability and usability rates of devices and equipment⁸⁶. Surgeon preference cards used to prepare surgical set-ups are often unreliable, with instruments added intra-operatively due to patient anatomy, contamination or error. Efforts to enhance the reliability of surgeon preference cards include frequent revision based on actual surgical requirements²⁵.

Without policy change, waste and inefficiencies will likely continue to impact patient outcomes such as surgical cancellations⁸⁷ and delays in emergency surgical operating lists⁸⁸. Lost time caused by medical device waste has a knock-on effect of delaying surgery for other patients; waiting for instrument availability is a logistic factor known to influence the queue of surgical cases⁸⁸. These delays reportedly lead to conflict between theatre managers and surgeons⁸⁸; however, the impact on patient outcomes is not often measured.

Studies included in this scoping review suggested that unaddressed communication failures impact surgical set-up processes^{27,50,54}. Although surgical devices are prescribed in advance, it has been argued that theatre nurses need more support and surgeons have

passive involvement in surgical set-up processes^{5,89}. Over-supply by perioperative services results in underutilisation. The volume of time and energy that perioperative nurses subsequently spend counting and managing complex medical devices is acknowledged within limited primary research^{56,90}. Procedural interruptions arising from surgical set-up problems are a distraction to the surgical team and raise concerns for patient safety^{1,91}. Apart from fixing excessive volume of surgical devices through optimisation downstream, there is limited research focused on proactively improving communication and collaboration between stakeholders to identify and effectively coordinate the surgical devices actually needed.

Chasseigne et al.⁵⁴ suggested that unused opened devices were mostly preventable through effective communication about the surgical set-up. Potential causes of perioperative communication failure include inadequate pre-operative preparation, lack of personnel and disruptive behaviours including the perception that nurses serve as 'secretaries and problem solvers for the whole team'^{92, p. e4}. Addressing communication failures and the perception of nurses as secretaries will require a comprehensive approach to improve medical, nursing and relevant stakeholder collaboration and ensure necessary procedural devices are identified and planned in advance. Collaborative approaches must consider potential variations and unforeseen circumstance to minimise errors and omissions.

Limitations

The scoping review only included studies written in English language and therefore may be limited in generalisability in countries where English is not the first language.

Conclusions

Fixing the issue of surgical set-up waste through optimisation is a short-term solution to a complex and evolving long-term problem. Most research into the surgical set-up comprises quality improvement studies, with limited primary research available. Mathematical modelling to predict the optimal number of resources to deliver a service may be helpful from a limited management perspective; however, it does not resolve unaddressed human factors, such as communication and collaboration for the surgical set up. Addressing challenges through proactive engagement could foster a culture of effective teamwork among health care providers working towards productive and efficient surgical set-up processes and ultimately improved safety and quality of procedural care.

Declaration of conflicting interests

The authors have declared no competing interests with respect to the research, authorship and publication of this article.

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