Palliative Care in Surgery: Defining the Research Priorities

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Objective: To describe the existing science of palliative care in surgery within three priority areas and expose specific gaps within the field.

Background: Given the acute and often life-limiting nature of surgical illness, as well as the potential for treatment to induce further suffering, surgical patients have considerable palliative care needs. Yet these patients are less likely to receive palliative care than their medical counterparts and palliative care consultations often occur when death is imminent, reflecting poor quality end-of-life care.

Methods: The National Institutes of Health and the National Palliative Care Research Center convened researchers from several medical subspecialties to develop a national agenda for palliative care research. The surgeon work group reviewed the existing surgical literature to identify critical knowledge gaps.

Results: To date, evidence to support the role of palliative care in surgical practice is sparse and palliative care research in surgery is encumbered by methodological challenges and entrenched cultural norms that impede appropriate provision of palliative care. Priorities for future research on palliative care in surgery include: 1) measuring outcomes that matter to patients, 2) communication and decision making, and 3) delivery of palliative care to surgical patients.

Conclusions: Surgical patients would likely benefit from early palliative care delivered alongside surgical treatment to promote goal-concordant decision making and to improve patients' physical, emotional, social and spiritual well-being and quality of life. We propose a research agenda to address major gaps in the literature and provide a road map for future investigation.

Keywords: palliative care, palliative surgery, surgical research

(Ann Surg 2017;xx:xxx–xxx)

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Disclosure: Dr Schwarze is supported by the Grants for Early Medical/Surgical Specialists’ Transition to Aging Research Award (GEMSTAR R03AG047920) and the American Geriatrics Society/Society of Vascular Surgery, Jahngen Career Development Award. Dr Schwarze also received funding during this time but not for support of this project from PCORI(1502-27462), the Greenwall Foundation, and the National Palliative Care Research Center. Dr Cooper is supported by the Paul B. Beeson Emerging Leaders Career Development Award in Aging (1K76AG054859-01) and the Cambia Foundation. Dr Cooper also received funding during this time, but not for support of this project, from PCORI (1502-27462), the American Geriatrics Society Geriatrics for Specialists Initiative, National Cancer Institute (1R35CA197730-01), and the National Institute on Aging (95R01AG04518-02). Dr Mosenthal received funding during this time, but not for support of this project, from PCORI (1502-27462). Dr Lilley reports no disclosures.

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ISSN: 0003-4932/16/XXXX-0001
DOI: 10.1097/SLA.0000000000002253

Palliative care is a multidisciplinary specialty which aims to relieve suffering and support quality of life for seriously ill patients and their families. In a statement on the principles of palliative care in surgery, the American College of Surgeons recognized the life-affirming role of palliative care in the management of surgical patients with serious illness, and emphasized the need to provide palliative care alongside life-prolonging and curative surgical treatments. Despite the burdens of surgical treatment and frequently life-limiting nature of surgical illness, palliative care delivery remains insufficient for surgical patients.

In 2003, the American College of Surgeon Palliative Care Workgroup identified 7 priority areas to build the science around palliative care in surgery, including surgical decision making, patient-oriented decision making, end-of-life decision making, symptom management, communication, processes of care, and surgical education about palliative care. A systematic review of the literature from 1994 to 2014, however, reveals only 25 studies focused on palliative care interventions for surgical patients. Like other assessments of palliative care interventions for nonsurgical patients, these studies suggest that palliative care interventions for surgical patients may reduce healthcare utilization and improve advance care planning without increasing mortality. Nonetheless, interventions to promote the alignment of surgical treatment decisions with patients’ goals of care and research on how to integrate palliative care principles into rescue-oriented surgical culture are notably absent. Furthermore, existing research is difficult to interpret due to the array of heterogeneous outcomes targeted. Additional methodological weaknesses include small sample size, single-center studies, and inadequate follow-up.

Recent reports from the Institute of Medicine and the National Institutes of Health (NIH) identify key contributions of palliative care for the management of seriously ill patients, including improved symptom management, better quality of life, reduced healthcare costs, and higher quality physician-patient communication. In light of these benefits, the Institute of Medicine and NIH reports called for increased research and support for the unmet needs of patients and families. The NIH and the National Palliative Care Research Center convened subspecialty work groups to develop a national agenda for palliative care research in several medical disciplines. Herein, we provide an overview of major gaps in the current evidence (Table 1) and identify 3 priority areas for research on palliative care in surgery targeted to fill these gaps (Table 2).

PRIORITY AREA 1: MEASURING OUTCOMES THAT MATTER TO PATIENTS

Defining Outcomes That Patients Value

A major pitfall in measurement to improve quality is that measured and reported outcomes, such as 30-day mortality, fall short of measuring outcomes most meaningful to patients and can impede patient access to palliative and end-of-life care. Survival is frequently measured in surgical research, but reporting the quantity
of days without concurrent reporting of quality of life does not attest to the patient experience. Furthermore, defining surgical quality and value based solely on survival duration incentivizes surgeons to prolong life, not improve it, and can impede integration of palliative care. Alternatively, measures of functional independence, disability-free survival, days spent at home, or freedom from pain after surgery provide information on outcomes that are both clinically meaningful and important to patients. Likewise, measures of health-care utilization required to achieve specific outcomes [i.e., surgical intensive care unit (SICU) admission, days on a ventilator, discharge to skilled nursing facilities, or long-term acute care hospitals] provide information regarding the burdens of treatment. In addition, survivors of postoperative complications, major trauma, and critical surgical illness are likely to have postacute palliative care needs; yet their long-term symptom burden, impairment, social concerns (i.e., need for assistance at home), and overall well-being are not well described in the current literature. Future studies need to characterize patients’ perspectives on the benefits, burdens, and tradeoffs associated with surgery and how best to measure the outcomes that are most meaningful to them.

Existing patient-reported outcomes measures used in palliative care were designed for patients with chronic, progressive illnesses, such as cancer, and are not readily translated to surgical patients because they do not account for the expected pain and disability that frequently accompany surgical recovery, nor do they distinguish acute postoperative symptoms from those which are chronic or refractory. Furthermore, some instruments, such as the Edmonton Symptom Assessment Scale, have only been validated in cancer patients receiving palliative care who forego disease-directed treatment. There are a few excellent examples of patient-reported outcomes measures developed for surgical problems in the literature; 1 study used qualitative data from adult trauma survivors and their caregivers to develop a questionnaire assessing aspects of quality of life that were specifically related to the trauma experience. Additional examples of surgery-specific instruments have been developed to measure patient-reported outcomes after breast surgery, bariatric surgery, colorectal surgery, and cosmetic surgery. Recent studies have also used measures from the NIH Patient-Reported Outcomes Measurement System to assess physical, mental, and social health in surgical patients. Although there are several instruments, which are appropriate for measuring patient-reported outcomes after surgery, the evidence remains thin due to their underuse in research. Observational studies measuring patient-reported outcomes are needed for a broad range of surgical subspecialties, including surgical oncology, neurosurgery, vascular surgery, and trauma.

In addition to patient-reported outcomes measures, there are other clinically relevant measures that are more aligned with outcomes that patient value than 30-day morbidity and mortality, such as longer-term survival, SICU days, and postacute care needs.
TABLE 2. Research Priorities for Palliative Care in Surgery

<table>
<thead>
<tr>
<th>Research Priority</th>
<th>Study Objective</th>
<th>Study Setting</th>
<th>Sample</th>
<th>Study Design</th>
</tr>
</thead>
<tbody>
<tr>
<td>Defining outcomes that matter to patients</td>
<td>Evaluate patient-reported outcome measures that reflect palliative care in surgery</td>
<td>Outpatient, inpatient, ICU, home</td>
<td>Patients who have major surgery and their caregivers</td>
<td>Qualitative and mixed method studies; psychometric research</td>
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<tr>
<td>Develop and validate instruments for patient-reported measures of palliative outcomes relevant to surgical patient populations</td>
<td>Develop measurable processes of care to deliver high quality palliative care to surgical patients</td>
<td>Outpatient, inpatient, ICU</td>
<td>Seriously ill surgical patients and their caregivers</td>
<td>Randomized controlled trials; cohort studies</td>
</tr>
<tr>
<td>Communication and decision making</td>
<td>Determine the effectiveness and comparative effectiveness of communication interventions versus usual care</td>
<td>Outpatient, inpatient, emergency room</td>
<td>Patients with serious illness and caregivers who are considering major surgery</td>
<td>Randomized controlled trials; Prospective and retrospective studies; quasi-experimental designs</td>
</tr>
<tr>
<td>Design and conduct large, multicenter trials assessing effectiveness of communication tools to disclose effectiveness in the perioperative period</td>
<td>Evaluate interventions to improve perioperative decision making and align surgical treatments with outcomes patients value</td>
<td>Inpatient, emergency room, ICU</td>
<td>Patients who experience complications and their families</td>
<td>Randomized controlled trials; prospective and retrospective studies; quasi-experimental designs</td>
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<tr>
<td>Delivery of palliative care to surgical patients</td>
<td>Increase acceptance and utilization of palliative care in surgical culture and practice</td>
<td>Academic centers and community hospitals</td>
<td>Surgical clinicians (ie, surgeons, nurses, anesthesiologists)</td>
<td>Randomized controlled trials; prospective and retrospective studies; quasi-experimental designs</td>
</tr>
<tr>
<td>Determine effectiveness and comparative effectiveness of targeted early versus late palliative care on healthcare cost, symptom management, quality of life, and caregiver burden</td>
<td>Increase timeliness of palliative care interventions for surgical patients with complex palliative care needs</td>
<td>Community, outpatient, inpatient</td>
<td>Surgical patients with poor prognosis and their caregivers</td>
<td>Randomized controlled trials; prospective and retrospective studies; quasi-experimental designs</td>
</tr>
<tr>
<td>Design and conduct large, multisite studies to compare palliative surgery versus medical management on symptom burden and quality of life.</td>
<td>Examine the effect of palliative surgical procedures on patient-reported outcomes</td>
<td>Outpatient, inpatient</td>
<td>Patients with oncologic, vascular, cardiac surgical problems, and their caregivers</td>
<td>Mixed-methods studies; randomized controlled trials; prospective and retrospective studies; quasi-experimental designs</td>
</tr>
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ACP indicates advance care planning; DNR, do-not-resuscitate; ICU, intensive care unit; LST, life-sustaining treatment.

Future work will also need to determine the feasibility and validity of incorporating these metrics into assessment and reporting of surgical quality.

**Measures to Evaluate High-quality Palliative Care in Surgery**

Measures that reflect the timely and appropriate delivery of high-quality palliative care in surgery are needed to promote accountability and identify targets for improvement. These metrics within the scope of surgical practice should address 2 separate issues:

1. Palliative care provision for seriously ill surgical patients and (2) management of patients undergoing palliative surgery.

Evaluating and improving processes of care, such as documentation of advance directives, and quality indicators for care at the end of life, such as hospice enrollment and death on life-sustaining treatments, are particularly pertinent for surgical patients at high risk of death. The National Quality Forum has endorsed 24 measures to assess the utilization and adequacy of palliative care in multiple settings. These measures were, however, designed around the needs of patients afflicted with suffering as they approach the final stages of...
illness, and may not be relevant for the management of seriously ill surgical patients, many of whom have a high symptom burden and a high risk of mortality but are not clearly expected to die in the short term.

Efforts are underway to collect national data to analyze patterns and utilization of do-not-resuscitate orders, palliative care consultation, and hospice referral in older surgical patients. These data, however, do not provide in-depth understanding of how and why these processes occur, nor whether they are beneficial from a patient’s perspective; both are necessary to inform the development of surgery-specific indicators for high-quality palliative care (ie, quality of communication, adherence to treatment preferences, and quality of death and dying). Multi-institutional prospective cohort studies are needed to define and measure palliative care process measures for surgical practice and correlate them with patients’ perceptions, experiences, and outcomes of care.

Palliative surgical procedures are intended to reduce suffering or support quality of life rather than prolong life or cure disease. Prior studies have described the considerable risks of postoperative complications and mortality after palliative surgery, but few have measured the impact of palliative surgery on restoration of function and quality of life, or conversely, the occurrence of adverse outcomes that further threaten quality of life, function, and ability to achieve a good death. Absence of a uniform system for designating and classifying procedures performed with palliative intent presents a barrier to studying outcomes of palliative surgery. Generation of standards for palliative surgery will permit future studies to assess the quality of palliative surgical care using criteria consistent with high-quality palliative care, rather than current metrics used in surgery, namely mortality and morbidity. Future comparative effectiveness trials are needed to compare the effectiveness of surgical procedures to non-surgical management on palliative outcomes for multiple surgical indications, including limb salvage, valve repair, and malignant obstruction.

**PRIORITY AREA 2: COMMUNICATION AND DECISION MAKING**

**Aligning Surgical Treatments With Patient-oriented Outcomes**

Surgeons and patients face high-stakes care decisions in the perioperative period, specifically whether to proceed with surgical intervention and associated, potentially burdensome, postoperative treatments. To support complex, in-the-moment decision making—often when the patient’s clinical condition is changing or has changed rapidly—patients and their family members need to clearly understand the capacity and limitations of surgical intervention and the short- and long-term effects of surgery on their functional status and quality of life. Although patients frequently pursue surgery with the intent to cure disease, the trade-offs between cure and quality of life (ie, impaired functional status and prolonged pain and suffering) are typically value sensitive. In cases in which surgery is directed toward palliation or ameliorating symptoms, evaluation of these trade-offs brings added complexity to decisions for surgery. Moreover, clinical decisions are severely hampered by the paucity of data comparing longer-term survival, quality of life, and function after operative and nonoperative management. This lack of data hinders the consideration of palliative care as an adjunct or alternative to surgery.

Although innovations to improve preoperative communication and decision making have been described, whether these strategies improve the quality of surgical decisions (ie, better patient understanding of their disease and procedure, realistic expectations of recovery, reduced decisional regret) or other patient-oriented outcomes is unknown because assessment of these interventions is fraught with multiple serious methodological challenges. Two single-institution cohort studies suggest that preoperative interventions to better clarify patients’ disease understanding and treatment preferences are associated with a decrease in surgical procedures among frail older adults and improved symptom control and reduced morbidity and mortality after palliative procedures in patients with cancer. Both studies are, however, observational and lack a control group. Future randomized clinical trials and comparative effectiveness studies are needed to test structured communication interventions for surgical decision making that emphasize quality of life, long-term survival, and quality of death and dying. Valuable palliative care outcomes should include the alignment between patients’ goals and the likely outcomes of surgery, reduction of burdensome, unwanted or nonbeneficial postoperative interventions, and improvement in physical and psychological outcomes after surgery.

**Preoperative Advance Care Planning**

Because patients who have surgery are at risk for losing decision-making capacity for prolonged periods, it is important to clarify—before surgery—the desired outcome from the patient’s perspective, treatments patients are willing to endure to achieve those outcomes, and postoperative outcomes patients find unacceptable (ie, prolonged ventilator dependence). Patients who have major surgical procedures may also have desires to limit burdensome life-supporting treatments after surgery, and those with pre-existing directives restricting specific treatments may want to suspend these restrictions during the acute, perioperative period to achieve specific goals. Despite the importance of clarifying treatment preferences before surgery, some surgeons are resistant or reluctant to pursue preoperative advance care planning and data suggest that such conversations are often incomplete, or altogether absent, which can lead to unwanted postoperative treatment and conflict between surgeons and patients. Unless preferences are clarified beforehand, surgeons and surrogates may presume that permission for surgery implies permission for all postoperative treatments necessary to avoid postoperative death.

Small studies of preoperative advance care planning conversations with high-risk surgical patients and their surrogates suggest that interventional strategies can improve surrogate understanding of patient preferences. In these studies a total of only 28 patient-surrogate dyads, however, received an intervention, making it hard to draw definitive conclusions about intervention efficacy. Larger, hypothesis-driven studies are needed to determine the effect of preoperative advance care planning interventions to identify the patient’s preferred surrogate decision maker before surgery; elicit and document patients’ goals, expectations, and concerns about surgical treatment; and determine how much leeway patients might give to surgeons and surrogate decision makers to select the treatments needed to achieve these goals.

**Decision Making After Postoperative Complications or Critical Illness**

In the setting of postoperative complications, additional barriers to communication have been described, including surgeons’ heightened sense of duty and belief in surgical buy-in; that in accepting surgical care, patients have also agreed to any postoperative treatment deemed necessary to survive. Furthermore, in contrast to the more predictable decline in health and typical trajectory for patients with chronic, terminal illness, acutely ill surgical patients often experience minute-by-minute alterations in...
health status. When major complications occur, patients’ previously stated goals and desired surgical outcomes may become unattainable. Insofar as treatment preferences are context specific and contingent upon patients’ prognostic understanding, their priorities for treatment may change if the best possible outcome is no longer consistent with their individual values and judgments about quality of life. However, the combination of prognostic uncertainty, desire to rescue, and fear of extinguishing hope makes it difficult for surgeons and intensivists to communicate and provide accurate and precise information about outcomes for patients and their families.

Surrogate decision making in the SICU differs from most other intensive care settings in that the sudden decline to critical illness after acute surgical illness, postoperative complications, or traumatic injury is unexpected and the appointed surrogate is often poorly prepared for decision making. Given the acute shift in trajectory, this is particularly challenging for family members who often do not discuss preferences with loved ones before hospitalization, or engage in preoperative conversations between the patient and surgeon. There is little evidence about how best to support these unprepared surrogates for their decision-making role in the SICU. At the same time, few studies have examined interventions to communicate prognosis amid uncertainty and re-evaluate patients’ treatment preferences when postoperative complications or critical illness have altered the expected postoperative course. Future studies are needed to develop and refine communication tools to facilitate these conversations with patients and surrogates in the SICU and to evaluate whether rescue treatments to manage complications align with patients’ wishes in light of less desirable outcomes.

**PRIORITY AREA 3: DELIVERY OF PALLIATIVE CARE TO SURGICAL PATIENTS**

**Integrating Palliative Care Principles Into Routine Surgical Practice**

Surgical rescue culture and surgeons’ deep-seated notions about error and responsibility are frequently cited as barriers to improving palliative care for surgical patients. Interventions integrating elements of palliative care into routine surgical practice and promoting cultural changes through peer review, specifically morbidity and mortality rounds, have reported promising results. These single-institution studies, however, target high-risk patients at large academic medical centers. To establish durable improvements, dissemination and implementation studies are needed to develop scalable models of palliative care delivery and reproducible strategies for changing practice and culture. Large, multisite implementation studies of physician and systems-targeted interventions are needed to redirect treatment options so that surgery is not the default modality for patients known to have extremely poor survival due to baseline serious illness or acute surgical conditions. This requires a cultural shift promoting less aggressive treatments or comfort-directed care as reasonable adjuncts or alternatives to surgical management, rather than characterizing this high-quality care as “doing nothing.” Multidisciplinary, and multi-institutional, interventions are needed to educate surgical clinicians regarding the appropriateness of palliative care in the management of seriously ill patients. Studies will need to evaluate the adoption and maintenance of these interventions.

Palliative care education for surgeons is necessary to fully integrate palliative care into surgical care delivery; however, standardized and validated approaches for surgical palliative care education are lacking. In prior studies, surgical residents reported discomfort with conducting family meetings about end-of-life care. Others have found deficiencies in residents’ knowledge of palliative care and insufficient documentation of end-of-life care conversations. Few studies have tested the effect of palliative care education interventions on surgical patients’ outcomes. Studies are needed to delineate robust methods for teaching surgeons basic palliative care skills (ie, thoroughly exploring patients’ goals of treatment, managing acute and chronic symptoms, treating depression and anxiety), and to evaluate the effect of this training on patient outcomes. Innovative curriculum development, including simulation-based education, and competency-based assessment, is needed for surgeons in all stages of training. Incorporating core clinical and patient-reported outcomes and core processes (eg, documentation) in evaluating these interventions is essential.

**Developing Scalable Models of Primary Palliative Care Delivery for Surgical Patients**

Early integration of palliative care in the management of seriously ill nonsurgical patient populations is associated with improved quality of life, reduced healthcare costs, and longer survival. In contrast with the longitudinal management of chronic, progressive disease, surgical care typically focuses on acute conditions, with broad variation in recovery outcomes. Moreover, surgeons must quickly establish relationships with their patients and often have a finite role in their care. Thus, the approach used for integrating palliative care into other specialties does not translate into surgery specialties. There are no studies that have evaluated scalable models for delivering palliative care in the perioperative period and the optimal strategy for meeting complex palliative care needs in surgical populations. Large, multi-institutional interventional studies are needed to determine the effectiveness of palliative care interventions on patient and caregiver outcomes and healthcare cost.

**Identifying Patients Who Would Benefit From Palliative Care Specialist Consultation**

Patients with complex palliative care needs benefit from specialist consultations; however, palliative care consultations are less common among surgical patients than other patients and are often delayed until patients are within days of death. In the current treatment model, palliative care needs are typically unattended until the end of life. To attend to the palliative care needs of seriously ill surgical patients throughout all phases of care, especially as those needs gradually increase in intensity and complexity with illness progression, we need to shift our approach. Investigation is needed to identify seriously ill, but not imminently dying, surgical patients who would benefit from early palliative care intervention from surgeons with subsequent referral to palliative care specialists when their needs are beyond the scope of their primary surgical providers.

There are few studies that explore strategies for promoting timely palliative care consultation for surgical patients with unmet or complex needs. Results of 2 single-institution studies suggest that screening surgical patients for frailty and serious chronic illness may increase palliative care consultation and improved clinical outcomes. A third study, however, found no difference in the number of palliative care consultations.

Observational studies using qualitative, mixed methods and secondary dataset analyses, are needed to characterize patients with a large burden of suffering from symptoms, high postoperative morbidity and mortality, and those for whom surgery represents an inflection point in their health trajectory. Potential targets include a variety of patients with poor prognosis surgical illnesses, such as pancreatic cancer, peripheral vascular disease, and frail injured patients. Interventional studies are needed to examine whether targeted early palliative care has a similar effect on healthcare
utilization, treatment intensity, symptom management, survival, and quality of life for high-needs surgical cohorts as it does in lung cancer, advanced heart failure, and end-stage renal disease. 68,77–79

CONCLUSIONS

Seriously ill surgical patients have substantial palliative care needs that are often unrecognized and unaddressed. Although much has been accomplished since the first research agenda for palliative care in surgery was put forth in 2003, much more remains to be done. As the population ages and technical innovation advances, surgical patients will become increasingly complex as surgeons and patients navigate the blurred boundaries between technically feasible, clinically appropriate, and value-concordant care. Building the science around palliative care in surgery will require the engagement and support of stakeholders, interdisciplinary collaboration, and development of new, well-trained researchers with interest in this field. The proposed research priorities will provide evidence to support lasting improvements and establish palliative care as a core tenet of high-quality surgical care.

ACKNOWLEDGMENTS

This work was made possible by the National Institute on Aging, a division of the NIH, and the National Palliative Care Research Center, who jointly sponsored a workshop to facilitate the development of research agendas exploring the role of palliative care in medical subspecialties, including cardiology, nephrology, pulmonary/critical care, and surgery.

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