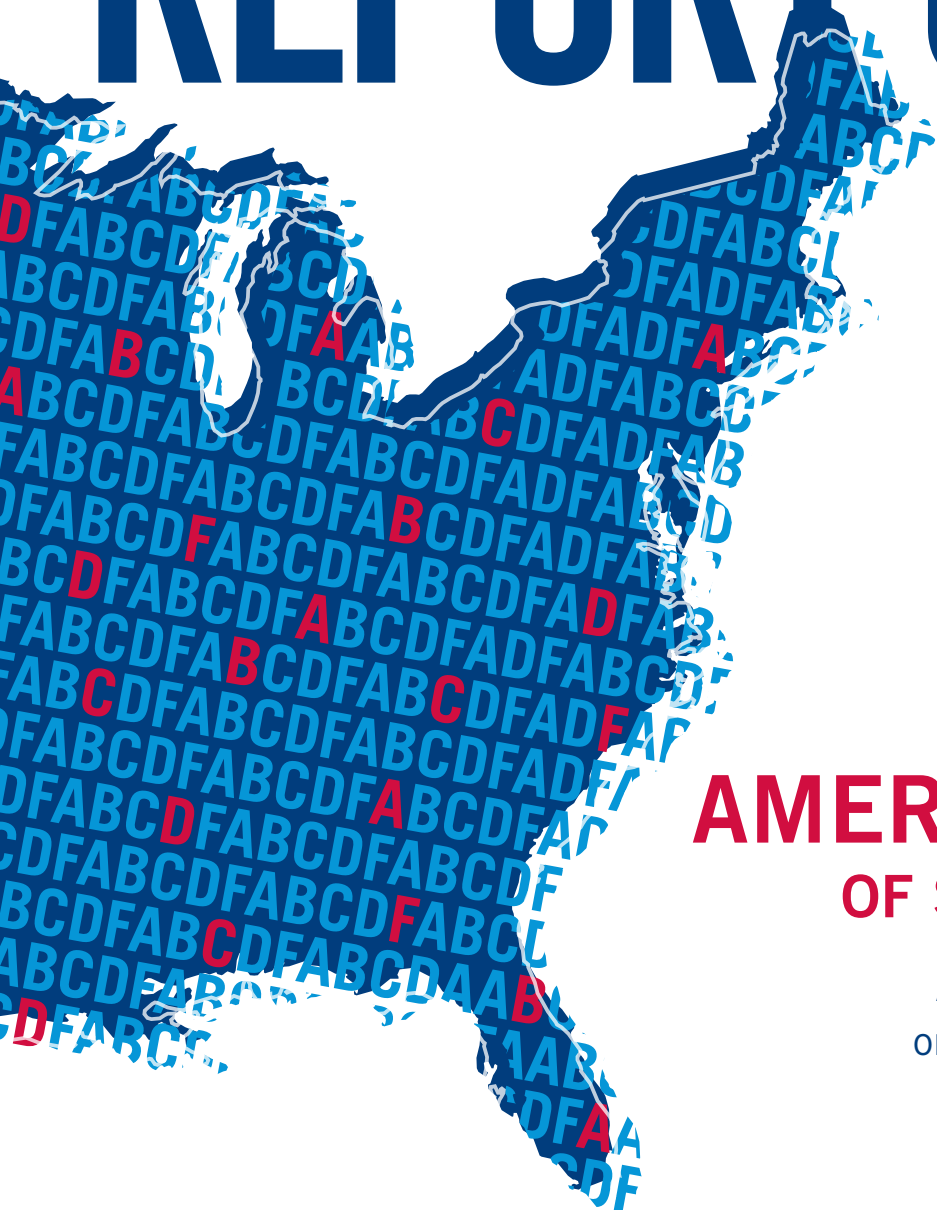




# REPORT CARD



## AMERICA'S CARE OF SERIOUS ILLNESS

A State-by-State Report Card  
on Access to Palliative Care in  
Our Nation's Hospitals

# REPORT CARD

## Preface

This report, an update of the 2008 publication *America's Care of Serious Illness: A State-by-State Report Card on Access to Palliative Care in Our Nation's Hospitals*, is the result of collaboration between the Center to Advance Palliative Care (CAPC) and the National Palliative Care Research Center (NPCRC). Highlighting the current prevalence of hospital palliative care programs, the report was undertaken in an effort to understand changes in access to such programs in the United States.

Drawing on the clinical and operational expertise of leaders in the field of palliative care, the research team was led by R. Sean Morrison, MD, Director of the National Palliative Care Research Center, and Diane E. Meier, MD, Director of the Center to Advance Palliative Care. Rachel Augustin, MPH, research officer for the Center to Advance Palliative Care, also contributed to the analyses contained in this report.

We would also like to acknowledge our funders, whose support enabled this study: Archstone Foundation, Brookdale Foundation, Donaghue Foundation, the Emily Davie and Joseph S. Kornfeld Foundation, the John A. Hartford Foundation, Mill Park Foundation, the Olive Branch Foundation, Partnership for Palliative Care, the Robert Wood Johnson Foundation, the United Hospital Fund and the Y. C. Ho/Helen and Michael Chiang Foundation.

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### **Diane E. Meier, MD**

*Director, Center to Advance Palliative Care*

May 2011

### **R. Sean Morrison, MD**

*Director, National Palliative Care Research Center*

May 2011

**Lead Author:** R. Sean Morrison, MD

**Coauthor:** Diane E. Meier, MD

**Lead Analysts:** Rachel Augustin, MPH, and Phomdaen Souvanna, MPP

**Additional Contributions:** Marian Appellof, Lisa Morgan, Carolyn Tieger

The major findings contained in this report build on research first described in “Variability in access to hospital palliative care in the United States” (*Journal of Palliative Medicine*, October 2008). The data were obtained primarily from the American Hospital Association (AHA) Annual Survey Database™ for fiscal year 2009. Additional data for the 2011 report were obtained from the American Board of Medical Specialties (ABMS), American Academy of Hospice and Palliative Medicine (AAHPM), National Board for Certification of Hospice and Palliative Nurses and the Dartmouth Atlas.

**Center to Advance Palliative Care**

1255 Fifth Avenue, Suite C-2

New York, New York 10029

212.201.2670

[www.capc.org](http://www.capc.org)

**National Palliative Care Research Center**

Box 1070

Mount Sinai School of Medicine

New York, New York 10029

212.241.7447

[www.npcrc.org](http://www.npcrc.org)

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# SECTION:

# 1

## INTRODUCTION

### *Palliative Care: The New Paradigm for Managing Serious Illness*

Today, approximately ninety million Americans are living with serious illness, and this number is expected to more than double over the next twenty-five years.<sup>1</sup>

We are also an aging nation. By the year 2030, according to the Administration on Aging (AoA), there will be more than 72.1 million Americans over age sixty-five in the U.S. (20 percent of the total U.S. population), and more than twice the number in 2000.<sup>2</sup>

While baby boomers will experience unprecedented longevity (those who reach sixty-five can expect to live well into their mid-eighties), these additional years carry a significant burden of chronic illness. Most will live with multiple, long-term medical conditions and need expert, compassionate and coordinated delivery of health care.

Yet despite the highest per-capita spending on health care in the world, more than 50 percent of caregivers of Americans hospitalized with a serious illness report less than optimal care,<sup>3</sup> and more than 30 percent of families lose most or all of their savings while caring for a loved one with a serious illness.<sup>4</sup>

Of the 1.5 million Americans who die of a chronic illness each year, more than 70 percent are admitted to a hospital during the last six months of life.<sup>5</sup> Recent studies also show that most people living with a serious illness experience inadequately treated symptoms, fragmented care, poor communication with

- 1 The Dartmouth Institute for Health Policy and Clinical Practice. The Dartmouth Atlas of Health Care website, [www.dartmouthatlas.org](http://www.dartmouthatlas.org). Accessed July 14, 2008.
- 2 Aging Statistics. The Administration on Aging (AoA) website, [www.aoa.gov](http://www.aoa.gov). Accessed June 9, 2011.
- 3 Teno JM, Clarridge BR, Casey V, et al. Family perspectives on end-of-life care at the last place of care. *JAMA*. 2004 Jan 7;291(1):88–93.
- 4 Covinsky KE, Goldman L, Cook E, et al. The impact of serious illness on patients' families. *JAMA*. 1994 Dec 21;272(23):1839–44.
- 5 Dartmouth Atlas of Health Care.

their doctors and enormous strains on their family caregivers. Consider the following:

- > *1 in 4 patients report inadequate treatment of pain and shortness of breath.<sup>6</sup>*
- > *1 in 3 families report inadequate emotional support.<sup>7</sup>*
- > *1 in 3 patients report that they receive no education on how to treat their pain and other symptoms following a hospital stay.<sup>8</sup>*
- > *1 in 3 patients are not provided with arrangements for follow-up care after hospital discharge.<sup>9</sup>*

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Palliative care is specialized medical care focused on providing relief from the symptoms, pain and stress of a serious illness—whatever the diagnosis. It is appropriate at any age and at any stage of a serious illness, and can be provided together with curative treatment. The goal is to improve quality of life for both the patient and the family.

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To address these gaps in care, hospitals have been quietly establishing, over the last decade, what has become a growing trend—palliative care programs. Palliative care is specialized medical care focused on providing relief from the symptoms, pain and stresses of a serious illness—whatever the diagnosis. It is appropriate at any age and at any stage of a serious illness, and can be provided together with curative treatment. The goal is to improve quality of life for both the patient and the family.

Palliative care is team-based care that includes specialized physicians, nurses, social workers and others (such as chaplains) who work with a patient's other doctors to provide an extra layer of support. These specialists treat pain and other symptoms; provide time-intensive communication; support complex medical decision making; ensure practical, spiritual and psychological support; and coordinate care across all the settings where patients receive medical care. An important focus is the plan of care, which is developed after in-depth consultation to ensure that medical treatments align with the needs, goals and priorities of the patient and the family.

Palliative care is not defined by a person's life expectancy. Indeed, patients receiving palliative care from the point of diagnosis, or may live for years with one or more chronic diseases.



The good news is that in the last ten years the number of palliative care teams in hospitals has more than doubled in response to the increasing numbers and needs of Americans living with serious and chronic illnesses, and the caregiving realities faced by their families.

The bad news is that despite the myriad benefits of palliative care, as well as its recent growth, more progress is needed. Millions of Americans with serious illness do not yet have access to palliative care from the point of diagnosis throughout the course of illness. Availability varies considerably by region and by state.

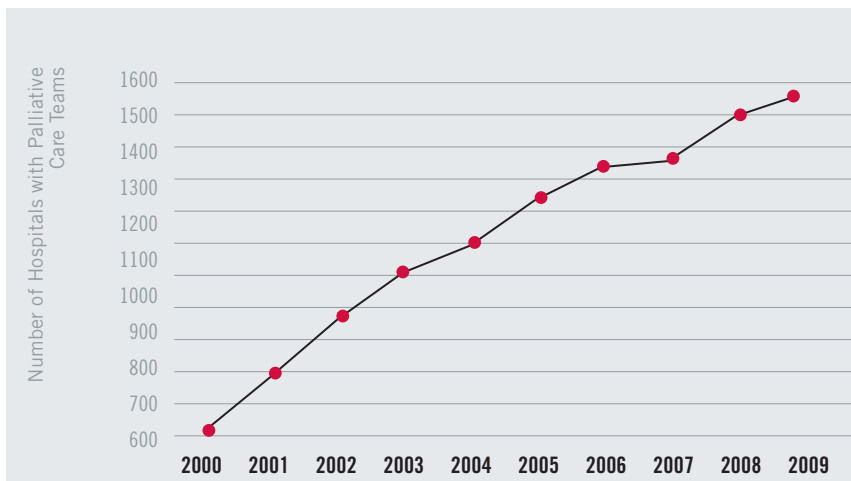
This Report Card examines variation in access to palliative care at the state level in order to help both the public and policymakers increase the availability of these services for all Americans in need.

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The 2011 Report Card website can be found at [www.capc.org/reportcard](http://www.capc.org/reportcard).

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### Prevalence of U.S. Hospital Palliative Care Teams 2000–2009



Source: Center to Advance Palliative Care, March 2011

- 6 Teno et al. Family perspectives.
- 7 Ibid.
- 8 The Commonwealth Fund. Care coordination. *Quality Matters*. 2007 May/June;24.
- 9 Ibid.



# SECTION:

# 2

## POLICY IMPERATIVES

### *Initiatives to Support Palliative Care*

Over the last ten years palliative care has been one of the fastest-growing trends in health care. In fact, the number of palliative care programs within hospital settings has increased by 138 percent since 2000. Surprisingly, this development has occurred with little notice and within the current health care system. The reason for this quiet but steady growth is that palliative care has been embraced by patients and their families for the simple reason that it gives them control and choice over their own care and responds to their needs as they themselves have defined them. The strong partnership of patient, family and the palliative care team ensures that treatment goals are aligned and coordinated, and that open and full communication is maintained throughout the long, complex course of a serious illness.

Recent public opinion research by the national polling firm Public Opinion Strategies has shown that once informed about palliative care, 92 percent of the American public are highly likely to consider palliative care for themselves or their families if they have a serious illness. Ninety-two percent also said it is important that palliative care services be made available at all hospitals for patients with serious illness and their families throughout the United States.<sup>10</sup> The data also show a positive response to palliative care regardless of party affiliation.

To meet the growing demand for and access to palliative care, specific changes in our health care system are needed now.

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Recent public opinion research shows that 92 percent of people said it is important that palliative care services be made available at all hospitals for patients with serious illness and their families.

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<sup>10</sup> Public Opinion Strategies Poll, April 2011.

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## Policy initiatives to support palliative care:

- > Workforce
  - > Research
  - > Access and Quality
- 

While this Report Card demonstrates considerable growth in the number of palliative care programs across the United States, barriers in three key areas—workforce, research and access—currently prevent full availability of palliative care for all patients and families facing serious or life-threatening illness. Specifically, three key policy initiatives are needed:

- > *Investment in a trained workforce to ensure sufficient numbers of specialists both to teach all clinicians the fundamentals and to directly provide high-quality palliative care for the highest-risk and most complex patients;*
- > *Investment in the research necessary to establish a strong science base for palliative care and to expand palliative care's ability to improve both quality and length of life;*
- > *Investment in health care system capacity by requiring delivery of high-quality palliative care in hospitals, nursing homes and community settings through changes in measurement, payment and accreditation standards.*

## Workforce

A major barrier facing the expansion of palliative care services is the lack of palliative medicine physicians. Where there is approximately one cardiologist for every 71 persons experiencing a heart attack and one oncologist for every 141 newly diagnosed cancer patients, there is only one palliative medicine physician for every 1,200 persons living with a serious or life-threatening illness.

There are two principal reasons for the shortage of palliative medicine providers. First, palliative medicine is a new specialty, formally recognized in 2007. Thus, although the number of training programs is increasing, as of 2010 there were only 73 accredited allopathic subspecialty training fellowship programs in the United States, collectively producing approximately 86 new palliative medicine physicians per year.<sup>11</sup> Second, Medicare funding, which currently supports the majority of post-medical school residency training for physician specialties in the United States, does not support palliative medicine specialty training. This is a consequence of the 1997 Balanced Budget Act's cap on Medicare-funded graduate medical education (GME) slots. The majority of academic medical centers—the site of training for medical students, residents and fellows—currently are at or exceed their GME residency cap. As a result, Medicare dollars are unavailable for training

<sup>11</sup> Accreditation Council for Graduate Medical Education website, [www.acgme.org](http://www.acgme.org). Accessed June 16, 2011.

palliative medicine physicians, and current palliative medicine fellowship training programs are inadequately supported by scarce philanthropic dollars.

Three initiatives are necessary to ensure that Americans with serious illness and their families have access to quality palliative care. First, future generations of palliative medicine specialists require support and expansion of palliative medicine fellowship training programs. Demand for this training is high, as palliative medicine is the fastest-growing medical subspecialty in the United States and expansion and support of fellowship programs is needed. Second, a substantial number of midcareer physicians are seeking training in order to transition from their current field of practice into palliative medicine. Third, all physicians who care for patients with serious illness should be trained in the core principles of palliative medicine to ensure that they know how to communicate with seriously ill patients and treat pain and other distressing symptoms.

Several policy initiatives are likely to have a major impact on increasing the current number of palliative care clinicians, including proposals to:

- > *Lift the GME cap on Medicare-funded residency positions and redistribute unused GME slots to support Accreditation Council for Graduate Medical Education (ACGME)-approved palliative medicine fellowship training.*
- > *Establish loan-forgiveness programs for palliative care physicians and nurses through the Health Resources and Services Administration (HRSA) to promote palliative care as a viable career path for young health care professionals.*
- > *Create HRSA Title VII-supported career development awards (similar to Geriatric Health Professions Training Programs) to support clinician-educators who can integrate palliative care into medical, nursing school and postgraduate training curricula.*
- > *Establish HRSA awards to support retraining of midcareer current medical and nursing workforce for this new specialty.*
- > *Require CME training for referring physicians as a condition of licensure at the state level—similar to California's provision for pain management training—that would ensure physician competency in the core principles and practice of palliative medicine.*

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There is only one palliative medicine physician for every 1,200 persons living with a serious or life-threatening illness.

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The National Institutes of Health (NIH) and the Institute of Medicine have repeatedly called for substantial investments in palliative care research.

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## Research

Increased funding for palliative care research is needed to strengthen clinical practice and improve health care delivery that will maximize quality of life for patients and families facing serious and chronic illnesses.

The National Institutes of Health (NIH) and the Institute of Medicine have repeatedly called for substantial investments in palliative care research. Yet, as of 2009, there were only 114 active awarded NIH grants supporting palliative care research. While the growth of the palliative care field has been remarkable, there is a need to strengthen the knowledge base that supports basic elements of clinical practice in pain and symptom management, communication skills and care coordination. To that end, specific focus areas and funding mechanisms for palliative care research should be developed in all NIH institutes, the Agency for Healthcare Research and Quality (AHRQ) as well as the Veterans Administration.

Policy initiatives to address this knowledge gap are straightforward and easily integrated within current biomedical research funding structures:

- > *NIH/AHRQ should allocate 2 percent (increase from 0.01 percent between 2003–2005) of their current budgets to focus on symptom relief, communication in the setting of serious illness and research focused on the impact of concurrent palliative care along with disease-directed therapies for patients with serious illness.*
- > *NIH should establish an Office of Palliative Care Research modeled after the Office of AIDS Research to oversee and ensure appropriate distribution of research funding.*
- > *Existing NIH career development award mechanisms should be utilized to support junior investigators and midcareer palliative care investigators in order to build a critical mass of established palliative care researchers.*

## Access and Quality

While physician and (in some states) advanced practice nursing services provided by a palliative care team are covered by most private insurance, as well as Medicare and Medicaid, palliative care is not readily accessible to all Americans. It is offered

primarily in hospital settings, with only a few major medical centers providing outpatient or home-delivered palliative care programs. As of 2009, 63 percent of U.S. hospitals with at least fifty beds and 85 percent of hospitals with more than 300 beds reported the presence of a palliative care team. While this represents an increase of 138 percent from 2000, it is not good enough. Currently, there are only about 1,568 hospitals out of a total of 2,489 (American Hospital Association estimate for 2010) nationwide that offer some type of palliative care program.

Part of the problem is that the current business model for palliative care is based on cost avoidance (reducing unwanted and unnecessary utilization) rather than on revenue generation. This model is unusual in health care, requires sophisticated analytic methods to employ successfully and is thus difficult to integrate into hospitals' current operating metrics. Additionally, accreditation standards for hospitals and nursing homes do not currently require the presence of a quality palliative care program despite publication of consensus standards by the National Quality Forum (NQF). Near-term policy solutions that could increase access to quality palliative care are as follows:

- > *The reimbursement structure for palliative care should enable hospitals and nursing homes to support provision of palliative care services. Palliative care specialists, like critical care physicians, should be compensated at a level commensurate with the complexity and seriousness of illness in the patients they are serving. That includes the work of the entire interdisciplinary team (doctor, nurse, social worker and others). Advanced practice nurses delivering palliative care are allowed to bill fee for service in some states, but not all. To improve access to palliative care services throughout the country, advanced practice nurse compensation should be standardized.*
- > *Given the high costs and poor quality of care received by patients with serious illnesses, new delivery and payment models rewarding whole-person care instead of fragmented fee-for-service care should ensure access to palliative care as a core component of available services.*
- > *Palliative care teams meeting quality standards should be a condition of accreditation for all U.S. hospitals and nursing homes.*

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Approximately 63 percent of hospitals report a palliative care program.

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# SECTION:

# 3

## THE CASE FOR PALLIATIVE CARE

### *Providing Care That People Want*

Palliative care teams are tackling a perfect storm. Unprecedented growth of an aging population; the failure to recognize and treat pain and other distressing symptoms experienced by seriously ill people; the widespread availability and use of costly medical technologies that fail to restore health and functional independence, or a quality of life acceptable to the patient; exponential cost increases; and dissatisfaction with and confusion about the medical care system for those with serious illness—all form the context for the rapid growth of the field of palliative care.

Expert attention to physical and psychological symptoms and improved communication and coordination—the domains of palliative care—form the foundation for effective health care. Yet, this is not what seriously ill patients usually get.

Typically, doctors generate a list of medical problems—based upon discrete organ systems and diseases—and approach the “problem list” as a series of items on a checklist. Although effective for single disease states, this approach falls apart in the setting of multiple diseases and serious conditions interacting and influencing one another. Emotional and spiritual distress and enormous caregiver needs make the situation even more complex. In New York City, for example, the average person with a serious illness is receiving care from twelve different specialist physicians, with no single individual coordinating care.<sup>12</sup>

<sup>12</sup> Dartmouth Atlas of Health Care.

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Palliative care ensures that the person is viewed in his or her entirety, not as a collection of organs and medical problems.

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In contrast, palliative care teams begin by interviewing patients about their goals for care, the symptoms and other issues that bother them and the needs of their families. By starting with the patient's and family's priorities and goals, palliative care teams ensure that the care received is aligned with patient and family needs and is driven by their agenda, not that of the health care system or the physician. Furthermore, and perhaps most importantly, palliative care ensures that the person is viewed in his or her entirety, not as a collection of organs and medical problems.

By following this approach:

***Palliative care teams reduce high levels of suffering and distress among patients with serious illness, at any age and at any stage of disease.*** People with serious and life-threatening illnesses live not only with multiple diseases (e.g., advanced heart and lung disease, diabetes, arthritis, cancer), but also with multiple symptoms (e.g., pain, anxiety, shortness of breath, fatigue, depression, sleep disorders). Palliative care reduces pain and other debilitating symptoms, and it can be provided at any age and any stage in the course of an illness, along with curative treatment.

***Palliative care teams improve communication among patients, families and health care providers.*** Palliative care results in the identification of achievable care goals, better patient and family understanding of disease and treatment choices, and improved communication among patients, families and all health care providers.

***Palliative care teams address the needs of family caregivers as they cope with and care for a loved one with serious illness.*** By identifying family needs, fears and knowledge gaps, palliative care programs have been shown to improve a family's feeling of confidence and security in the setting of a loved one's serious illness. People with serious illnesses live with enormous caregiving requirements (e.g., assistance with feeding, bathing, toileting, walking); complicated transitions (e.g., rehabilitation, visiting-nurse services, nursing homes, care coordination) and many other complex needs. Caregivers need practical and emotional support, and palliative care ensures both.

***Palliative care teams reduce unwanted, unnecessary and painful interventions.*** By establishing patients' goals and focusing on what are important and achievable outcomes for medical treatment, palliative care teams ensure that patients receive the most effective and timely treatments and that their valuable time and energy is protected from burdensome tests and procedures that will not lead to improvements or changes in care.

***Palliative care can improve survival.*** A recent study published in the *New England Journal of Medicine* showed that receiving palliative care early in the treatment of advanced lung cancer led to significant improvements in both quality of life and mood. As compared with patients receiving only disease-directed care, patients receiving early palliative care required less emergency hospital and ICU care, and survived approximately 2.7 months longer.<sup>13</sup>

***Palliative care teams improve patient and family satisfaction.*** Surveys of patients cared for by palliative care teams routinely reveal high patient and family satisfaction—among the highest rates observed in any study of hospitalized and seriously ill patients. A recent poll conducted by Public Opinion Strategies also shows that palliative care provides what people want: quality of life; relief from suffering; support for family caregivers; control and choice.<sup>14</sup>

***Palliative care improves quality of care while reducing costs.*** About 10 percent of all Medicare beneficiaries have five or more chronic conditions and two-thirds of Medicare spending goes to cover their care. This patient population is also the most likely to benefit from palliative care. Recent studies indicate, in fact, that by focusing on the needs of our most complex patients by matching treatments with their goals, improving their quality of life and providing support for care at home—thus helping to avoid preventable emergency hospitalizations—palliative care teams can contribute to substantial cost savings.<sup>15</sup> Savings are the result of patients' decisions to shift their care out of the hospital and into the community, usually into their own home, ultimately reducing costs.

13 Temel JS, Greer JA, Muzikansky A, et al. Early palliative care for patients with metastatic non-small-cell lung cancer. *N Engl J Med*. 2010 Aug 19;363 (8):733–42.

14 Public Opinion Strategies Poll, April 2011.

15 Morrison RS, Penrod JD, Cassel JB, et al. Cost savings associated with US hospital palliative care consultation programs. *Arch Intern Med*. 2008 Sep 8;168(16):1783–90.

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If palliative care teams were fully integrated into the nation's hospitals, total savings could exceed \$6 billion per year.

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As mentioned earlier in this report, once informed about palliative care, 92 percent of the American public wants access to palliative care programs in hospitals nationwide.<sup>16</sup> If this demand is met, quality of care for those patients with serious and chronic illnesses will go up but costs of care will go down. In fact, according to a recent report published by the Institute of Medicine, if palliative care teams were fully integrated into the nation's hospitals, total savings could exceed \$6 billion per year.<sup>17</sup>

In summary, coordinated care by an interdisciplinary team is at the heart of palliative care and represents the best response to the long-term nature of chronic, multifaceted illnesses. Studies demonstrate that when patients have control of their care by working with palliative care teams to set treatment goals, quality of care is improved as well as patient and family satisfaction. Such an approach reduces the need for hospitalization and duplicative or unnecessary testing and medication and ultimately reduces costs for patients, families and health care providers alike.

<sup>16</sup> Public Opinion Strategies Poll, April 2011.

<sup>17</sup> Morrison RS, Meier D, Carlson M. Palliative care, access, quality and costs, in chapter 16, Medically complex patients. In: Yong PL, Saunders RS, Olsen LA, eds. *The Healthcare Imperative: Lowering Costs and Improving Outcomes*. Washington, DC: National Academies Press; 2010: 498–504.

# SECTION:

# 4

## HOW DOES OUR NATION FARE?

### *A National and State-by-State Review and Report Card*

#### **The 2011 State-by-State Report Card Shows Improvement**

The 2011 State-by-State Report Card updates our 2008 Report Card.<sup>18</sup> Reflecting data from the American Hospital Association Annual Survey Database™ for fiscal year 2009,<sup>19</sup> we again examine prevalence and geographic variations in access to palliative care in U.S. hospitals. Specifically, in 2011 we examine:

1. *Patient access to palliative care services in hospitals; and*
2. *Patient access to board-certified palliative care professionals (e.g., physicians and nurses).*

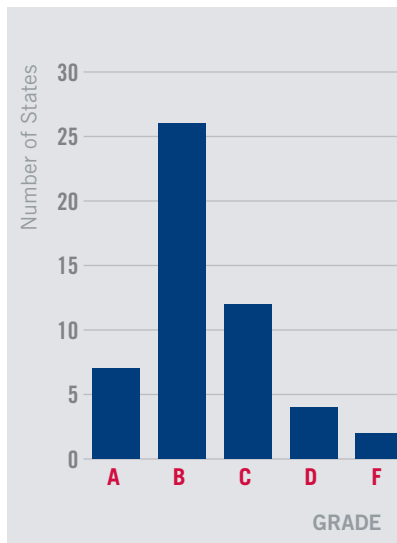
Hospitals containing fifty or more beds are the primary focus of the 2011 Report Card. Hospitals with fewer than fifty beds see only a small number of patients with serious or life-threatening illnesses, and are therefore unlikely to be able to support the model of an interdisciplinary palliative care consultation team that includes, at minimum, a specialty-level palliative care physician, nurse and social worker. Also excluded from the study are rehabilitation hospitals; psychiatric hospitals; subacute and chronic-care facilities; eye, ear, nose and throat hospitals; pediatric hospitals; hospitals under federal control (e.g., V.A. and military hospitals); hospitals located outside the fifty states and the District of Columbia; and hospitals that did not respond to the AHA Annual Survey Database™.

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<sup>18</sup> Goldsmith BA, Dietrich J, Du Q, Morrison RS. Variability in access to hospital palliative care in the United States. *J Palliat Med.* 2008 Oct;11(8):1094–1102.

<sup>19</sup> American Hospital Association Annual Survey™ for fiscal year 2009.

### Distribution of State Grades



Findings in 2011 demonstrate considerable improvement since the 2008 Report Card. However, significant variation still persists from state to state. The 2011 analysis also demonstrates fluctuations in the number of hospital palliative care teams within a given state from year to year.

Table 1 of the Appendix presents the state-by-state findings of hospitals evaluated in this study—hospitals with fifty or more beds. Table 1 also shows the prevalence of palliative care teams among sole community provider hospitals, larger hospitals with 300 or more beds and small hospitals with fewer than fifty beds.

#### ***What Are the Key Predictors of Access to Palliative Care?***

As we have seen in other areas of our health care system, we found wide geographic variation in access to palliative care services. Factors predicting the presence of a hospital palliative care team have not changed markedly since the 2008 Report Card.

- > *Large hospitals with 300 or more beds are more likely to report a palliative care team (85 percent).*
- > *Public hospitals (54 percent), for-profit hospitals (26 percent), and sole community provider hospitals (37 percent) are less likely to report a palliative care team.*
- > *On average, midsize (50–300 beds) and large hospitals (over 300 beds) are more likely to be not-for-profit.*

#### **Improvements Since the 2008 Report Card**

Since our last report, the overall prevalence of hospital palliative care teams among hospitals with fifty or more beds increased 13.3 percent in the Midwest, 21.7 percent in the Northeast, 23.7 percent in the South and 29.3 percent in the West. The cumulative national average is 63 percent (1,568 out of 2,489 study hospitals).

There have also been improvements at the state level. In 2008, the eleven states with the lowest prevalence rates were: Mississippi (10 percent), Alabama (16 percent), and Oklahoma (19 percent), Nevada (23 percent), Wyoming (25 percent), Louisiana (27 percent), South Carolina (30 percent), Texas (33 percent), New Mexico (33 percent), Kentucky (37 percent) and Georgia (38 percent).

Today, only Delaware and Mississippi get an F (20 percent). However, Mississippi demonstrated substantial improvement since 2008, doubling its prevalence from 10 to 20 percent. Alabama increased from 16 to 28 percent, and Oklahoma increased from 19 to 30 percent. Seven states improved their grades from a D to a C: Georgia, Kentucky, New Mexico, Texas, South Carolina, Louisiana and Wyoming. Nevada saw dramatic gains, rising from a D to a B grade.

States getting an A nearly tripled from three in the 2008 Report Card to seven plus the District of Columbia in the 2011 Report Card. Along with the District of Columbia (100 percent), states receiving an A grade now include: Vermont (100 percent), Nebraska (93 percent), Maryland (90 percent), Minnesota (89 percent), Oregon (88 percent), Rhode Island (88 percent) and Washington (83 percent).

Fifty percent of states received a B grade. The top nine states with B grades, ranging from 75 to 80 percent, include: New Jersey (80 percent), Ohio (80 percent), Virginia (78 percent), South Dakota (78 percent), New Hampshire (77 percent), Michigan (76 percent), and Missouri, New York and North Carolina (each with 75 percent).

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Our nation receives an overall grade of B.

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## Does Your State Make the Grade?

Despite the rapid growth of palliative care teams in our nation's hospitals and improvement since the last Report Card, access to palliative care services in the United States must improve if we are to adequately care for our sickest patients. In 2008, our nation received an overall grade of C. In 2011, the country receives an overall grade of B. Seven states plus the District of Columbia now receive a grade of A, with more than 80 percent of hospitals reporting palliative care services. More than half of the fifty states receive a grade of B. Fewer than 25 percent of states now need significant improvement (C). Approximately 12 percent receive nonpassing grades of D or F.

### Top performers (programs in 83% to 100% of hospitals): States receiving an A grade

District of Columbia	100%
Maryland	90%
Minnesota	89%
Nebraska	93%
Oregon	88%
Rhode Island	88%
Vermont	100%
Washington	83%

### On their way (programs in 61% to 80% of hospitals): States receiving a B grade

Arizona	69%
California	67%
Colorado	73%
Connecticut	72%
Florida	62%
Idaho	63%
Illinois	67%
Indiana	63%
Iowa	61%
Maine	71%
Massachusetts	67%
Michigan	76%
Missouri	75%
Montana	67%
Nevada	69%
New Hampshire	77%
New Jersey	80%
New York	75%
North Carolina	75%
North Dakota	67%

Ohio	80%
Pennsylvania	67%
South Dakota	78%
Virginia	78%
Wisconsin	74%

### States in the middle (programs in 42% to 60% of hospitals): States receiving a C grade

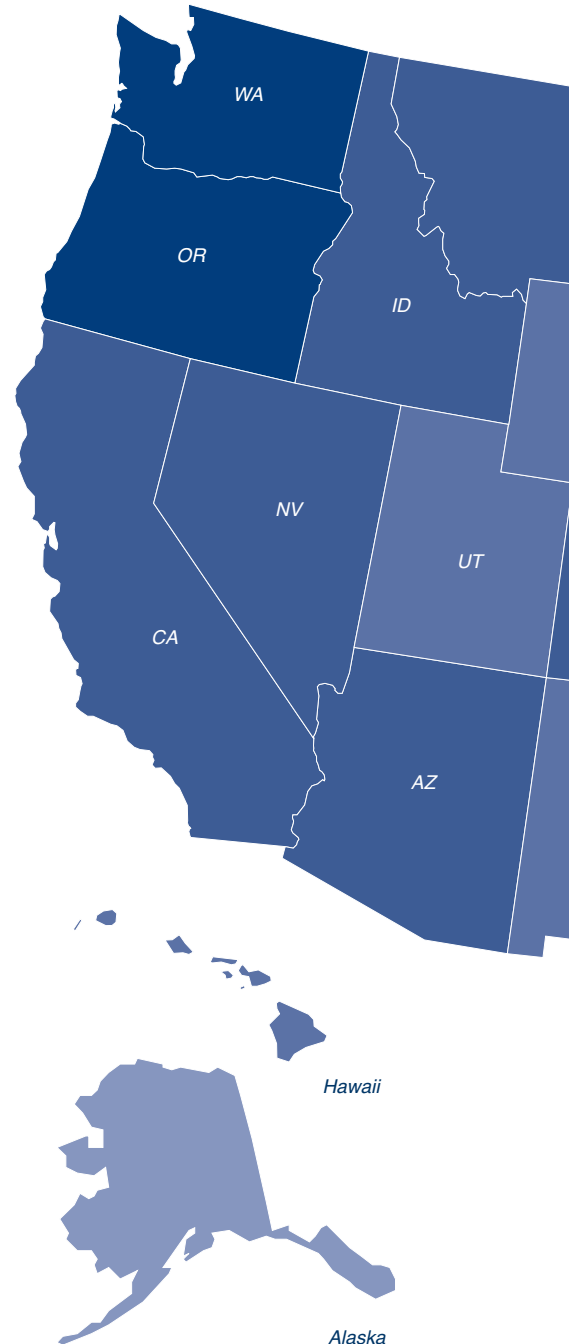
Georgia	43%
Hawaii	58%
Kansas	47%
Kentucky	55%
Louisiana	43%
New Mexico	44%
South Carolina	51%
Tennessee	52%
Texas	42%
Utah	60%
West Virginia	55%
Wyoming	50%

### States that need significant improvement (programs in 28% to 38% of hospitals): States receiving a D grade

Alabama	28%
Alaska	29%
Arkansas	38%
Oklahoma	30%

### States with little or no access (programs in 0% to 20% of hospitals): States receiving an F grade

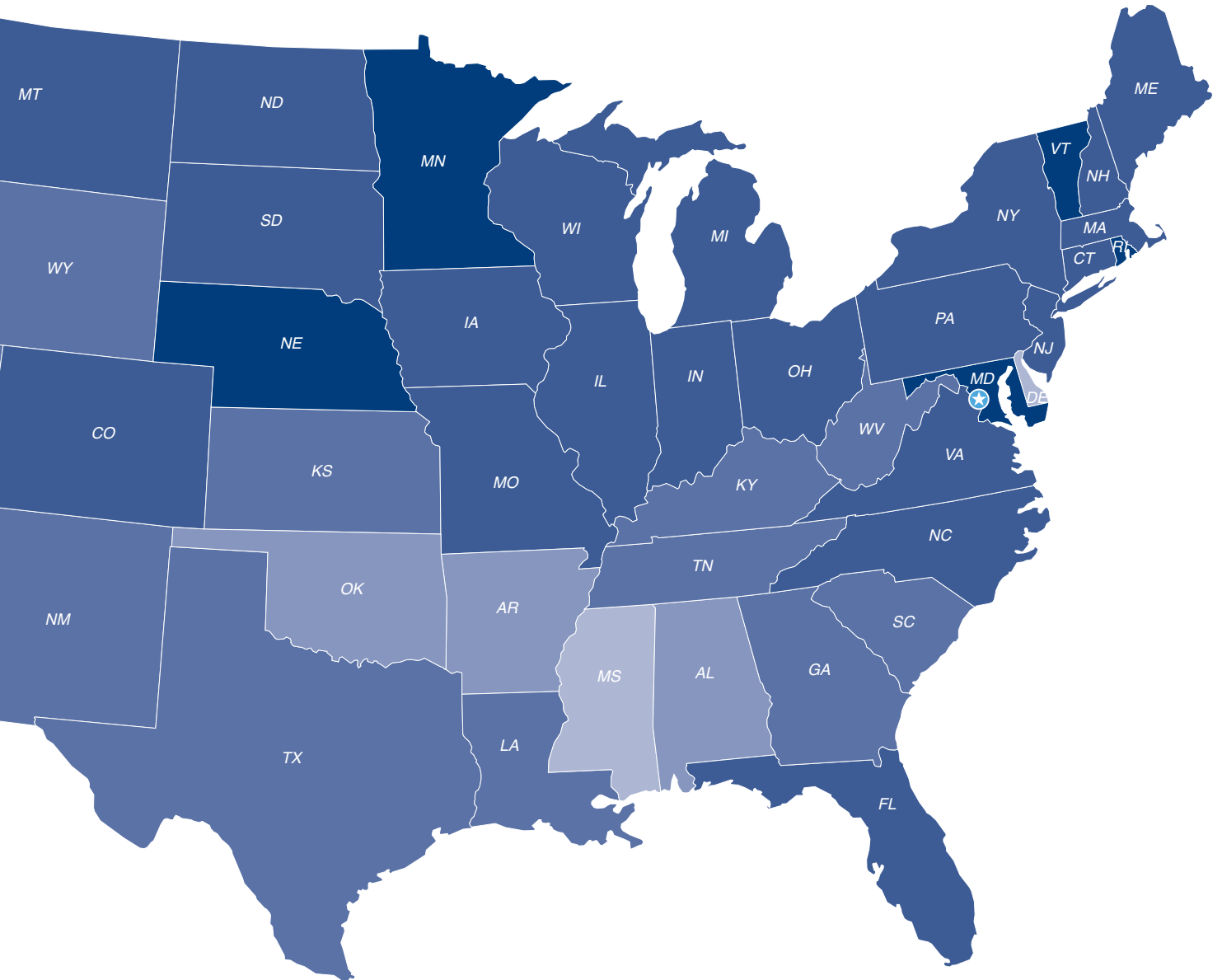
Delaware	20%
Mississippi	20%





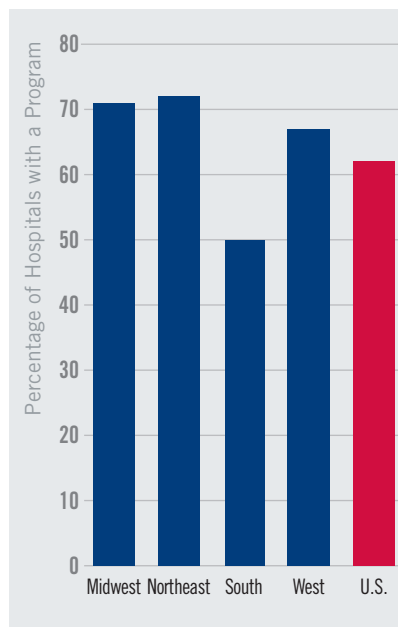
AMERICA'S CARE OF SERIOUS ILLNESS:

A State-by-State Report Card on Access to Palliative Care in Our Nation's Hospitals



- A** Top Performers
- B** On Their Way
- C** In the Middle
- D** Need Significant Improvement
- F** Little or No Access

Distribution of Palliative Care Programs by Region



### Where You Live Matters

“Geography is destiny,” as the Dartmouth Atlas researchers have often pointed out,<sup>20</sup> appears to hold true when speaking of access to hospitals offering palliative care. In the Northeast, 73 percent of hospitals with fifty or more beds report a palliative care team, compared to only 51 percent in the South. Among hospitals with fewer than fifty beds, your chances of having access to a palliative care team are extremely limited, particularly in the South, where only 14 percent of small hospitals (fewer than fifty beds) report having palliative care services.

Nationally, the prevalence of palliative care in large hospitals with 300 or more beds is 85 percent, ranging from 50 percent (Alabama and Delaware) to 100 percent in nineteen states. The lowest prevalence rates of palliative care in large hospitals were found in Alabama (50 percent) and Delaware (50 percent). The highest rates in large hospitals were found in nineteen states with 100 percent prevalence (see Appendix Table 1).

The number of small hospitals (under fifty beds) varies widely by state. For example, four states (Connecticut, Delaware, New Jersey and Rhode Island) and the District of Columbia do not report any hospitals of this size. The 2011 Report Card shows that the national prevalence rate for palliative care services in small hospitals is 22 percent. The lowest prevalence rates in small hospitals are found in Louisiana (0 percent), Mississippi (0 percent) and Alabama (4 percent). The highest prevalence rates for small hospitals are found in New Hampshire (62 percent) and Maine (58 percent).

### Underserved Populations Have Less Access to Palliative Care

In addition to marked disparities in geographic availability, we observed low rates of access to palliative care in public and sole community provider hospitals. Public and sole community provider hospitals often serve as the only option for medical care for the forty-seven million Americans lacking health care coverage or living in geographically isolated communities. Therefore, our finding that the majority of these institutions continue to lack palliative care services speaks to a disparity in access to comprehensive care for America’s most vulnerable patient populations. Notably:

<sup>20</sup> The Dartmouth Institute for Health Policy and Clinical Practice. The Dartmouth Atlas of Health Care website, [www.dartmouthatlas.org](http://www.dartmouthatlas.org). Accessed January 13, 2011.

- > *Only 54 percent of public hospitals provide their patients access to palliative care.*
- > *Fewer than 40 percent of sole community provider hospitals offer their patients access to palliative care.*

### **Lack of Board-Certified Palliative Medicine Physicians**

Improving access to palliative care for America's patients and families requires a workforce highly trained in the fundamentals of palliative medicine. Today in the United States there are 2,887 physicians board-certified in palliative medicine.

Prevalence of board-certified palliative care physicians varies across states. The highest rates are to be found in Hawaii (1 per 154 Medicare deaths), New Mexico (1 per 222 Medicare deaths) and Colorado (1 per 244 Medicare deaths). The lowest rates are in Mississippi (1 per 1,698 Medicare deaths), Rhode Island (1 per 1,267 Medicare deaths), Montana (1 per 1,218 Medicare deaths), Delaware (1 per 1,060 Medicare deaths) and South Dakota (1 per 1,037 Medicare deaths).

Table 2 of the Appendix presents a state-by-state comparison of the number of board-certified physicians as well as advanced practice nurses and registered nurses.

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Hawaii, New Mexico and Colorado have the greatest number of board-certified palliative care physicians.

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# SECTION:

# 5

## CONCLUSION

### *Equal Care Requires Equal Access*

Despite rapid improvements over the last few years, people living with a serious illness still experience significant disparities in access to a hospital that provides palliative care. Some areas of the United States are better equipped than others, and some hospitals are more committed than others to providing these services.

The ultimate goal of palliative care is to improve overall quality of life and quality of care for patients experiencing serious and chronic illnesses and their families. To benefit from palliative care, however, patients and their caregivers must be able to access these services in their local hospital. Physicians in training must learn from direct experience at the bedside with high-quality hospital palliative care teams.

Focused efforts by hospital administrators, the health care community and policymakers are required to promote the development of quality palliative care programs in all hospitals, with special attention needed in small, rural, public and for-profit hospitals (see “Policy Imperatives”). This will result in a more efficient and effective use of hospital resources, and will enhance the quality of care delivered to our nation’s most seriously ill patients and their families.

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Policy initiatives that address workforce, research and patient access could rapidly bring palliative care to scale in the United States.

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Studies over the past twenty years have conclusively demonstrated that too many seriously ill Americans experience treatable pain and distress and are financially devastated because of the high costs of medical care. At the same time, rising costs threaten Medicare just as millions of boomers are turning to it for health insurance coverage. Palliative care offers a logical and patient-centered approach to improving medical care by focusing on quality of life and ensuring that care is matched to patient and family goals.

Patients who have the information they need to choose care plans that meet their personal goals mean a substantial reduction in harmful and unnecessary hospitalizations for the highest-risk, most vulnerable patient population. Policy initiatives that address workforce, research and patient access could rapidly bring palliative care to scale in the United States. The implementation of such policies would help meet the needs of our sickest children and a growing population of older Americans with long-term chronic conditions.

# APPENDIX:

# 6

## TABLES AND GLOSSARY

### *Grades, Prevalence, Distributions and Terms*

Although this report assigned grades to states, regions and the nation based on the percentage of hospitals (with fifty or more beds) that reported a palliative care team to the American Hospital Association, it may be useful in the future to look more closely at how hospitals of various sizes and types rate.

**Table 1**, “Prevalence and Distribution of Palliative Care in U.S. Hospitals,” lists percentages and raw numbers of hospitals reporting palliative care programs by hospital size (50-plus, 300-plus, and fewer than 50 beds) and type within each state and region and for the nation overall. *Note: The number of public and for-profit hospitals is a subset of 50-plus hospital beds.*

**Table 2**, “Prevalence of Palliative Care Health Care Professionals,” lists numbers of physicians,<sup>21</sup> advanced practice nurses and registered nurses who are certified in palliative care. It also shows the numbers of Medicare deaths (overall and in the hospital) per certified palliative care clinician to indicate how the distribution of qualified professionals affects quality of care for our sickest hospitalized patients. These numbers are also listed by state, region and the entire country for comparison.

Please refer to the 2011 Report Card website ([www.capc.org/reportcard](http://www.capc.org/reportcard)) for the methodology used in this report.

The Glossary contains terms used throughout this report, along with others that are common in similar publications.

<sup>21</sup> Physicians are certified in hospice and palliative medicine in each state by either the American Board of Medical Specialties (ABMS) or the American Board of Hospice and Palliative Medicine (ABHPM). The information includes results from the 2008 ABMS certification examination. As of the writing of this report, the 2010 exam results were unavailable. Statistics are derived from the *Official ABMS Directory of Board Certified Medical Specialists* provided by Elsevier, and the ABHPM directory database of board-certified diplomates provided by the American Academy of Hospice and Palliative Medicine (AAHPM).

Table 1 Prevalence and Distribution of Palliative Care in U.S. Hospitals

PERCENT OF HOSPITALS REPORTING A PALLIATIVE CARE PROGRAM							
	Grade	50+ Beds	For-Profit	Public	Sole Community Provider	300+ Beds	<50 Beds
Illinois	B	67% (76/113)	0% (0/5)	29% (2/7)	63% (5/8)	87% (26/30)	13% (6/45)
Indiana	B	63% (39/62)	45% (5/11)	63% (10/16)	33% (1/3)	71% (10/14)	24% (9/37)
Iowa	B	61% (20/33)	-(0/0)	50% (3/6)	57% (4/7)	100% (5/5)	29% (24/84)
Kansas	C	47% (16/34)	13% (1/8)	50% (3/6)	31% (4/13)	80% (4/5)	15% (15/98)
Michigan	B	76% (65/85)	50% (1/2)	33% (2/6)	56% (10/18)	97% (28/29)	44% (20/45)
Minnesota	A	89% (31/35)	-(0/0)	100% (3/3)	67% (4/6)	100% (11/11)	37% (22/60)
Missouri	B	75% (51/68)	42% (5/12)	75% (9/12)	50% (9/18)	100% (19/19)	16% (7/45)
Nebraska	A	93% (13/14)	-(0/0)	100% (1/1)	67% (2/3)	100% (2/2)	21% (8/38)
North Dakota	B	67% (4/6)	-(0/0)	-(0/0)	67% (4/6)	100% (1/1)	46% (12/26)
Ohio	B	80% (87/109)	100% (1/1)	73% (8/11)	100% (3/3)	97% (29/30)	27% (9/33)
South Dakota	B	78% (7/9)	100% (1/1)	-(0/0)	56% (5/9)	67% (2/3)	17% (7/41)
Wisconsin	B	74% (42/57)	100% (1/1)	-(0/0)	50% (3/6)	78% (7/9)	27% (17/64)
<b>MIDWEST</b>	<b>B</b>	<b>72% (451/625)</b>	<b>37% (15/41)</b>	<b>60% (41/68)</b>	<b>54% (54/100)</b>	<b>91% (144/158)</b>	<b>25% (156/616)</b>
Connecticut	B	72% (18/25)	-(0/0)	0% (0/1)	-(0/0)	100% (9/9)	-(0/0)
Maine	B	71% (10/14)	-(0/0)	-(0/0)	75% (3/4)	100% (2/2)	58% (11/19)
Massachusetts	B	67% (32/48)	67% (2/3)	50% (1/2)	100% (1/1)	93% (13/14)	43% (3/7)
New Hampshire	B	77% (10/13)	50% (1/2)	-(0/0)	-(0/0)	100% (1/1)	62% (8/13)
New Jersey	B	80% (45/56)	-(0/0)	50% (1/2)	-(0/0)	89% (24/27)	-(0/0)
New York	B	75% (103/138)	-(0/0)	89% (17/19)	53% (8/15)	89% (55/62)	33% (3/9)
Pennsylvania	B	67% (79/118)	14% (1/7)	-(0/0)	50% (5/10)	92% (33/36)	26% (5/19)
Rhode Island	A	88% (7/8)	-(0/0)	-(0/0)	-(0/0)	100% (2/2)	-(0/0)
Vermont	A	100% (4/4)	-(0/0)	-(0/0)	100% (2/2)	100% (1/1)	50% (3/6)
<b>NORTHEAST</b>	<b>B</b>	<b>73% (308/424)</b>	<b>33% (4/12)</b>	<b>79% (19/24)</b>	<b>59% (19/32)</b>	<b>91% (140/154)</b>	<b>45% (33/73)</b>
Alabama	D	28% (16/58)	6% (1/18)	38% (10/26)	11% (1/9)	50% (6/12)	4% (1/28)
Arkansas	D	38% (15/39)	14% (2/14)	67% (2/3)	19% (3/16)	71% (5/7)	13% (4/32)
Delaware	F	20% (1/5)	-(0/0)	-(0/0)	-(0/0)	50% (1/2)	-(0/0)
District of Columbia	A	100% (6/6)	100% (1/1)	-(0/0)	-(0/0)	100% (3/3)	-(0/0)
Florida	B	62% (69/112)	29% (10/34)	79% (15/19)	17% (1/6)	87% (39/45)	31% (4/13)



**Table 1** Prevalence and Distribution of Palliative Care in U.S. Hospitals *(continued)*

PERCENT OF HOSPITALS REPORTING A PALLIATIVE CARE PROGRAM							
	Grade	50+ Beds	For-Profit	Public	Sole Community Provider	300+ Beds	<50 Beds
Georgia	C	43% (29/67)	15% (2/13)	46% (6/13)	8% (1/12)	79% (15/19)	11% (4/35)
Kentucky	C	55% (29/53)	0% (0/7)	80% (4/5)	33% (2/6)	92% (12/13)	15% (4/27)
Louisiana	C	43% (22/51)	36% (4/11)	32% (8/25)	0% (0/3)	67% (8/12)	0% (0/25)
Maryland	A	90% (37/41)	100% (1/1)	-(0/0)	-(0/0)	83% (10/12)	50% (1/2)
Mississippi	F	20% (9/45)	0% (0/16)	25% (4/16)	8% (1/13)	70% (7/10)	0% (0/47)
North Carolina	B	75% (55/73)	75% (3/4)	72% (18/25)	56% (5/9)	94% (16/17)	19% (4/21)
Oklahoma	D	30% (12/40)	18% (2/11)	15% (2/13)	5% (1/21)	90% (9/10)	10% (5/51)
South Carolina	C	51% (22/43)	27% (4/15)	30% (3/10)	44% (4/9)	92% (12/13)	21% (3/14)
Tennessee	C	52% (28/54)	22% (4/18)	38% (3/8)	33% (1/3)	76% (13/17)	46% (12/26)
Texas	C	42% (90/214)	21% (22/104)	42% (11/26)	18% (9/51)	65% (36/55)	13% (23/180)
Virginia	B	78% (40/51)	22% (2/9)	100% (4/4)	56% (5/9)	93% (14/15)	45% (5/11)
West Virginia	C	55% (16/29)	20% (1/5)	50% (1/2)	44% (4/9)	100% (4/4)	23% (5/22)
<b>SOUTH</b>	<b>C</b>	<b>51% (496/981)</b>	<b>21% (59/281)</b>	<b>47% (91/195)</b>	<b>22% (38/176)</b>	<b>79% (210/266)</b>	<b>14% (75/534)</b>
Alaska	D	29% (2/7)	0% (0/2)	0% (0/1)	40% (2/5)	100% (1/1)	29% (2/7)
Arizona	B	69% (25/36)	0% (0/3)	75% (3/4)	56% (5/9)	81% (13/16)	11% (1/9)
California	B	67% (159/236)	32% (14/44)	63% (24/38)	35% (6/17)	83% (54/65)	21% (10/48)
Colorado	B	73% (24/33)	86% (6/7)	25% (1/4)	25% (3/12)	75% (6/8)	20% (6/30)
Hawaii	C	58% (7/12)	-(0/0)	0% (0/2)	25% (1/4)	100% (1/1)	20% (1/5)
Idaho	B	63% (5/8)	50% (1/2)	50% (1/2)	0% (0/1)	100% (2/2)	29% (6/21)
Montana	B	67% (6/9)	-(0/0)	-(0/0)	75% (6/8)	100% (1/1)	20% (9/46)
Nevada	B	69% (9/13)	60% (3/5)	0% (0/1)	0% (0/3)	67% (2/3)	17% (1/6)
New Mexico	C	44% (8/18)	33% (3/9)	50% (1/2)	36% (5/14)	100% (2/2)	23% (3/13)
Oregon	A	88% (23/26)	0% (0/2)	100% (2/2)	80% (4/5)	100% (7/7)	25% (8/32)
Utah	C	60% (9/15)	17% (1/6)	100% (1/1)	50% (3/6)	100% (5/5)	38% (5/13)
Washington	A	83% (33/40)	50% (2/4)	63% (5/8)	67% (2/3)	90% (9/10)	29% (8/28)
Wyoming	C	50% (3/6)	0% (0/1)	75% (3/4)	27% (3/11)	-(0/0)	11% (2/19)
<b>WEST</b>	<b>B</b>	<b>68% (313/459)</b>	<b>35% (30/85)</b>	<b>59% (41/69)</b>	<b>41% (40/98)</b>	<b>85% (103/121)</b>	<b>22% (62/277)</b>
<b>NATIONAL</b>	<b>B</b>	<b>63% (1568/2489)</b>	<b>26% (108/419)</b>	<b>54% (192/356)</b>	<b>37% (151/406)</b>	<b>85% (597/699)</b>	<b>22% (326/1500)</b>

Table 2 Prevalence of Palliative Care Health Care Professionals

	ABHPM-Certified Physicians	Medicare Deaths per Certified Physician	Medicare Hospital Deaths per Certified Physician	NBCHPN-Certified Advanced Practice Nurses	Medicare Deaths per Certified APN	Medicare Hospital Deaths per Certified APN	NBCHPN-Certified Registered Nurses	Medicare Deaths per Certified RN	Medicare Hospital Deaths per Certified RN	Medicare Deaths (2005)	Medicare Deaths Occurring Within a Hospital (2005)
<b>Illinois</b>	108	644	184	15	4,640	1,322	328	212	60	69,602	19,824
<b>Indiana</b>	39	995	266	6	6,470	1,728	167	232	62	38,819	10,366
<b>Iowa</b>	28	728	169	3	6,796	1,573	265	77	18	20,389	4,718
<b>Kansas</b>	37	472	117	8	2,181	539	121	144	36	17,448	4,313
<b>Michigan</b>	92	665	172	33	1,855	480	299	205	53	61,218	15,842
<b>Minnesota</b>	64	347	82	14	1,585	376	168	132	31	22,186	5,265
<b>Missouri</b>	54	642	189	5	6,932	2,045	208	167	49	34,662	10,225
<b>Nebraska</b>	11	987	252	—	—	—	76	143	36	10,852	2,772
<b>North Dakota</b>	8	540	124	1	4,319	990	9	480	110	4,319	990
<b>Ohio</b>	129	531	134	38	1,803	456	655	105	26	68,501	17,310
<b>South Dakota</b>	5	1,037	239	2	2,593	598	9	576	133	5,185	1,195
<b>Wisconsin</b>	70	469	109	24	1,369	318	265	124	29	32,849	7,635
<b>MIDWEST</b>	645	598	156	149	2,591	674	2,570	150	39	386,030	100,455
<b>Connecticut</b>	27	796	245	17	1,264	390	131	164	51	21,481	6,626
<b>Maine</b>	21	459	131	2	4,815	1,376	73	132	38	9,630	2,751
<b>Massachusetts</b>	84	387	115	31	1,049	311	294	111	33	32,530	9,631
<b>New Hampshire</b>	15	503	131	6	1,257	329	101	75	20	7,544	1,972
<b>New Jersey</b>	73	669	240	15	3,257	1,166	298	164	59	48,849	17,495
<b>New York</b>	268	343	128	59	1,560	581	509	181	67	92,025	34,259
<b>Pennsylvania</b>	134	564	158	32	2,364	662	505	150	42	75,640	21,175
<b>Rhode Island</b>	4	1,267	369	6	845	246	43	118	34	5,068	1,477
<b>Vermont</b>	11	351	91	3	1,286	335	45	86	22	3,859	1,006
<b>NORTHEAST</b>	637	466	151	171	1,735	1,735	1,999	148	48	296,626	96,392
<b>Alabama</b>	32	900	296	2	14,394	4,733	168	171	56	28,787	9,465
<b>Arkansas</b>	25	775	248	2	9,682	3,099	92	210	67	19,363	6,198
<b>Delaware</b>	5	1,060	291	2	2,649	727	55	96	26	5,298	1,453
<b>District of Columbia</b>	5	568	186	1	2,840	930	6	473	155	2,840	930
<b>Florida</b>	179	562	143	20	5,030	1,279	923	109	28	100,609	25,575

**Table 2** Prevalence of Palliative Care Health Care Professionals *(continued)*

	ABHPM-Certified Physicians	Medicare Deaths per Certified Physician	Medicare Hospital Deaths per Certified Physician	NBCHPN-Certified Advanced Practice Nurses	Medicare Deaths per Certified APN	Medicare Hospital Deaths per Certified APN	NBCHPN-Certified Registered Nurses	Medicare Deaths per Certified RN	Medicare Hospital Deaths per Certified RN	Medicare Deaths (2005)	Medicare Deaths Occurring Within a Hospital (2005)
<b>Georgia</b>	53	768	216	10	4,068	1,147	297	137	39	40,681	11,474
<b>Kentucky</b>	30	874	270	8	3,277	1,012	213	123	38	26,212	8,094
<b>Louisiana</b>	33	756	225	3	8,317	2,479	87	287	85	24,950	7,436
<b>Maryland</b>	51	535	164	14	1,949	597	232	118	36	27,288	8,361
<b>Mississippi</b>	11	1,698	592	—	—	—	38	491	171	18,673	6,512
<b>North Carolina</b>	89	553	172	20	2,461	766	477	103	32	49,224	15,314
<b>Oklahoma</b>	33	699	202	—	—	—	120	192	56	23,082	6,671
<b>South Carolina</b>	32	783	269	5	5,013	1,723	181	138	48	25,067	8,613
<b>Tennessee</b>	44	808	258	12	2,964	946	151	236	75	35,570	11,352
<b>Texas</b>	221	438	117	21	4,612	1,232	622	156	42	96,855	25,863
<b>Virginia</b>	76	511	162	23	1,689	536	281	138	44	38,845	12,317
<b>West Virginia</b>	15	898	290	1	13,463	4,352	110	122	40	13,463	4,352
<b>SOUTH</b>	934	618	182	144	4,006	4,006	4,053	142	42	576,807	169,980
<b>Alaska</b>	6	289	76	1	1,732	454	19	91	24	1,732	454
<b>Arizona</b>	79	270	54	9	2,369	478	405	53	11	21,321	4,300
<b>California</b>	303	336	100	41	2,482	739	722	141	42	101,756	30,308
<b>Colorado</b>	57	244	47	9	1,546	297	173	80	15	13,914	2,670
<b>Hawaii</b>	25	154	52	1	3,848	1,289	37	104	35	3,848	1,289
<b>Idaho</b>	9	758	163	3	2,274	489	56	122	26	6,823	1,467
<b>Montana</b>	5	1,218	283	3	2,030	472	36	169	39	6,089	1,417
<b>Nevada</b>	23	361	97	2	4,151	1,112	48	173	46	8,302	2,224
<b>New Mexico</b>	36	222	53	4	1,999	479	62	129	31	7,997	1,915
<b>Oregon</b>	48	298	67	10	1,428	323	218	66	15	14,284	3,225
<b>Utah</b>	25	350	60	4	2,188	378	70	125	22	8,753	1,510
<b>Washington</b>	52	530	127	15	1,837	442	274	101	24	27,559	6,626
<b>Wyoming</b>	3	920	229	—	—	—	29	95	24	2,759	686
<b>WEST</b>	671	336	87	102	2,207	2,207	2,149	105	27	225,137	58,091
<b>NATIONAL</b>	2,887	514	147	566	2,623	2,623	10,771	138	39	1,484,600	424,918

## Glossary

Here are some terms used throughout this report, along with others you are likely to encounter in similar publications.

### **Allopathic medicine**

Also known as “conventional medicine,” the system of medical practice that treats disease by using remedies that produce effects different from those produced by the disease under treatment. MDs, for example, practice allopathic medicine.

### **Hospice**

Hospice provides palliative care focused on terminally ill patients who are no longer seeking curative or life-prolonging treatments and who are expected to live for about six months or less. Hospice is provided through the Medicare hospice benefit and other payers, and can take place in the home, nursing home, hospital or at a stand-alone hospice.

### **Hospital palliative care program**

Interdisciplinary medical team focused on symptom management, intensive patient-physician-family communication, clarifying goals of treatment and coordination of care across health care settings.

### **Hospital types**

*For-profit hospitals:* Hospitals with fifty or more beds that are run by individuals, partnerships or corporations.

*Large hospitals:* Hospitals with 300 or more beds.

*Public hospitals:* Not-for-profit institutions run by a state, county, city, joint city-county, hospital district or authority, with fifty or more beds. (In this report, nongovernment, not-for-profit hospitals, such as those run by church-affiliated institutions, were excluded from this category.)

*Small hospitals:* Hospitals with fewer than fifty beds. Sole community provider (SCP) hospitals receive special designation by Medicare because they are located more than thirty-five miles from other hospitals, or they serve as the sole provider of health care services for a region due to limitations in local topography or prolonged severe weather conditions.

**Hospitalist**

A physician who is employed by the hospital and specializes in the care of hospitalized patients.

**National Consensus Project for Quality Palliative Care (NCP)**

A national consensus project that published formal definitions of palliative care in 2004 (*Clinical Practice Guidelines for Quality Palliative Care*), with a second edition released in 2009. The guidelines were intended to direct the development and structure of both new and existing palliative care teams; establish uniform definitions of the essential elements of palliative care; establish national goals for access to palliative care; promote performance measurement and quality improvement initiatives in palliative care services; and foster continuity of palliative care across settings (including home, hospital and hospice). The guidelines fall within eight defined domains of palliative care. (See [www.nationalconsensusproject.org](http://www.nationalconsensusproject.org).)

**National Quality Forum (NQF)**

A framework for preferred practices of providing quality palliative care, released in 2006. Practices are defined across the Institute of Medicine's six dimensions of quality: safe, effective, timely, efficient, equitable and patient-centered. The NQF Framework used the clinical practice guidelines of the National Consensus Project for Quality Palliative Care as a starting point for identifying expectations and best practices in palliative care. (See [www.qualityforum.org/palliative](http://www.qualityforum.org/palliative).)

**Palliative care, palliative medicine**

Palliative care (pronounced pal-lee-uh-tiv) is specialized medical care for people with serious illnesses. It focuses on providing patients with relief from the symptoms, pain and stress of a serious illness—whatever the diagnosis. The goal is to improve quality of life for both the patient and the family. Palliative care is provided by a team of doctors, nurses and other specialists who work together with a patient's other doctors to provide an extra layer of support. It is appropriate at any age

and at any stage in a serious illness and can be provided along with curative treatment.

**Palliative care team, palliative care program**

An interdisciplinary, specialized medical team that provides palliative care in the hospital. The team works together with a patient's own doctor and includes physicians, nurses and social workers. It may also include chaplains, massage therapists, rehabilitation experts, pharmacists, nutritionists and others.



### **The Center to Advance Palliative Care**

[www.capc.org](http://www.capc.org)

[www.getpalliativecare.org](http://www.getpalliativecare.org)

The Center to Advance Palliative Care (CAPC), established in 2000, is a not-for-profit national organization whose purpose is to provide health care professionals with the tools, training and technical assistance necessary to start and sustain successful palliative care programs in hospitals and other health care settings. The mission of CAPC is to increase the availability of quality palliative care services for people facing serious, complex illness.

Led by Diane E. Meier, MD, CAPC receives direction and technical assistance from Mount Sinai School of Medicine, New York City. CAPC also works in close collaboration with the National Palliative Care Research Center.

CAPC focuses primarily on the operational development of palliative care teams in hospitals, as most seriously ill patients require hospital care at one or more points during their illness. Because of the medical crises that often lead to hospitalization, hospitals are typically the places where planning for the next phase in the care continuum occurs.

CAPC is supported by funding from Archstone Foundation, Brookdale Foundation, Donaghue Foundation, the Emily Davie and Joseph S. Kornfeld Foundation, the John A. Hartford Foundation, Mill Park Foundation, the Olive Branch Foundation, Partnership for Palliative Care, the Robert Wood Johnson Foundation, the United Hospital Fund and the Y. C. Ho/Helen and Michael Chiang Foundation.



### **The National Palliative Care Research Center**

[www.npcrc.org](http://www.npcrc.org)

The mission of the National Palliative Care Research Center (NPCRC) is to improve care for patients with serious illness and address the needs of their families by promoting palliative care research. The NPCRC establishes priorities for palliative care research; develops a new generation of researchers in palliative care; and coordinates and supports studies focused on improving care for patients and families living with serious illness.

The NPCRC is led by R. Sean Morrison, MD, and is located in New York City at Mount Sinai School of Medicine. In collaboration with the Center to Advance Palliative Care, the NPCRC works to rapidly translate important research findings into clinical practice in order to improve the care received by both patients living with serious illness and their families.

The NPCRC was established in July 2005 with a grant from the Emily Davie and Joseph S. Kornfeld Foundation and has received subsequent funding from the Olive Branch Foundation, Brookdale Foundation, American Cancer Society, American Academy of Hospice and Palliative Medicine, Betty and Norman F. Levy Foundation, the Y. C. Ho/Helen and Michael Chiang Foundation, Hearst Foundations, National Institute on Aging, U.S. Cancer Pain Relief Committee, American Express Foundation, Mill Park Foundation, Hospice and Palliative Nurses Association, LIVESTRONG, the Atlantic Philanthropies and the Partnership for Palliative Care.





Center to Advance Palliative Care

1255 Fifth Avenue, Suite C-2 | New York, NY 10029 | 212.201.2670 | [www.capc.org/reportcard](http://www.capc.org/reportcard)

