

Physician Engagement in Palliative Care

Jim McGregor, MD, ACQHPM Catherine Russell-McGregor, RN, FNS,C



Disclosures

The following presenters have no relevant financial relationships with commercial interests to disclose:

- Jim McGregor, MD, ACQHPM
- Catherine Russell-McGregor, RN, FNS,C



Objectives:

Participants will be better able to:

- Differentiate between Palliative Care and Hospice.
- Integrate Advance Care Planning conversations in daily practice
- Demonstrate a strategies to encourage conversations around Advance Care Planning and goals of care.
- Describe a process in which to engage physicians and allied health professionals from different settings in transitions of care for palliative patients.



Palliative Care

- Medical care provided by an interdisciplinary team of health care professionals focusing on relief of suffering and support for the best possible QoL for patients facing serious lifethreatening illness and for their families
- Team members include physicians, nursing, social work, chaplaincy and other allied health professionals
- Can be in concert with disease modifying therapy



Benefits of Palliative Care

- Extends principles of hospice care to a broader population
- Earlier in disease course than hospice
- Comprehensive and specialized
- Pain and symptom management, advance care planning, psychosocial and spiritual support, coordination of care
- Definition may be able to be expanded to all aspects of medical care



Hospice

- Comprehensive and coordinated care for patients with limited life expectancy of 6 months
- Provided both at home and in institutions
- Bio-psycho-social-spiritual model
- Focused on comfort, dignity and healing at the end-of-life
- Medicare benefit

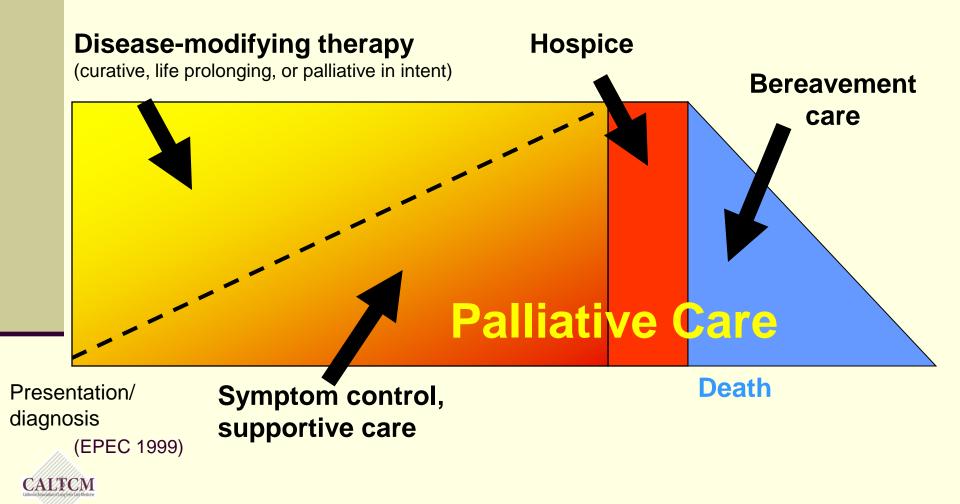


How Hospice Can Help?

- Nursing home residents who had a "hospice informational visit" had fewer acute care admissions (mean 0.28 vs. 0.49; p = .03) and fewer acute care days (mean 1.2 vs. 3.0; p = .03) than those who did not. Casarett et al. JAMA 294:211-217, 2005
- Patients with advanced lung cancer lived up to 3 months longer on hospice than those receiving "standard care". NEJM 2011
- Patients receiving hospice live longer. NHPCO



Advance Care Planning in the Continuum of Palliative Care



What % of patients if seriously ill want to talk about end-of-life care?

■ A:93%

■ B:12%

C:80%

■ D:54%

■ E:42%



The Conversation:

80% of people say that if seriously ill, they would want to talk to their doctor about end-of-life care

7% report having had an end-of-life conversation with their doctor





Requires physician engagement to be successful...

Barriers to Advance Care Planning Discussions

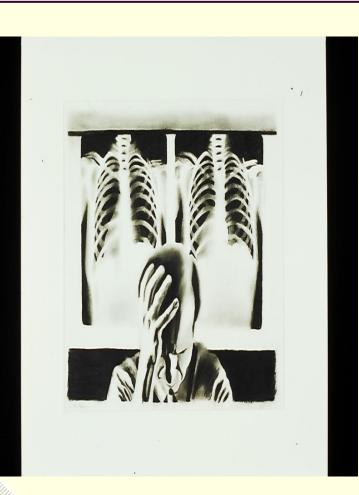
What do you think some barriers are?







Barriers to Communicating with Patients and Their Families



- Lack formal training
- Perceived lack of time
- Lack of reimbursement
- Fear of showing emotions
- Difficulties dealing with family responses in bearing "bad news"
- Provider's personal fear of end-oflife
- Healthcare professionals believe the patient should initiate the conversation
- Patient expect the physician to initiate



Communication Challenges...

- Culture
- Age
- Gender
- Personal & family experiences





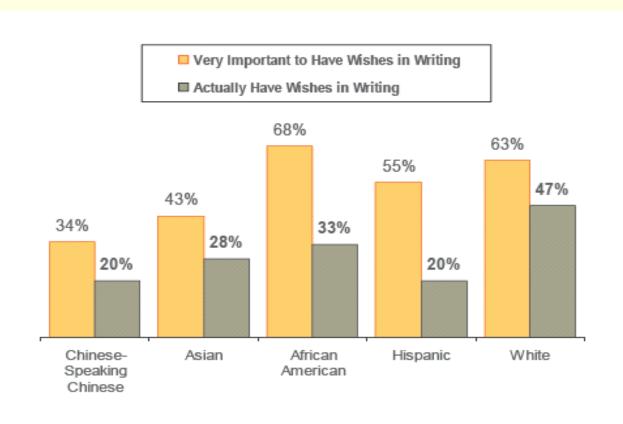








Variance in putting end-of-life wishes in writing





Advance Care Planning in SNF

- Lack of centralized document location
- Inconsistent documentation
- Habits
- Dispersion of responsibility
- Lack of time
- Under-recognition of team members efforts



Advance Care Planning in SNF

Patient with cognitive impairment

- Lack of family involvement
- Asymmetry of knowledge between professional and patient/family regarding the indications and effectiveness of treatments



Results of Discussions

What do you think the results of these discussions might be?



Benefits of ACP Discussions: The Patient's Perspective

Facing and preparing for end-of-life

- Achieving a sense of control
- Strengthening relationships
- Relieving burdens on loved ones
- Provides opportunities to work on life closure



Results of Discussions

- Leads to less depression, pain and anxiety
- Empowerment
- Autonomy
- Right care in the right setting for the patient
- Decreased rates of readmissions
- Access to outpatient resources
- Increased completion of AD or POLST
- Patient/family satisfaction



What Americans Want...

- Freedom from pain
- To die at home
- To be in the presence of their loved ones
- Not to be a burden to their family
- Spiritual well-being
- Honest dialogue from healthcare providers



Current ACP Practice: Causing Harm?

- Patients and families betrayed, misled?
- Health professionals pressured to complete forms, ask for decisions
- Patient and families stigmatized if they refuse to participate/make "wrong" choices
- Health professionals disengage due to experience of AD's not being helpful

J Pall Med Vol 5, 2002



% of Americans who would choose life prolongation if unlikely to regain consciousness?

■ A: 12%

■ B: 96%

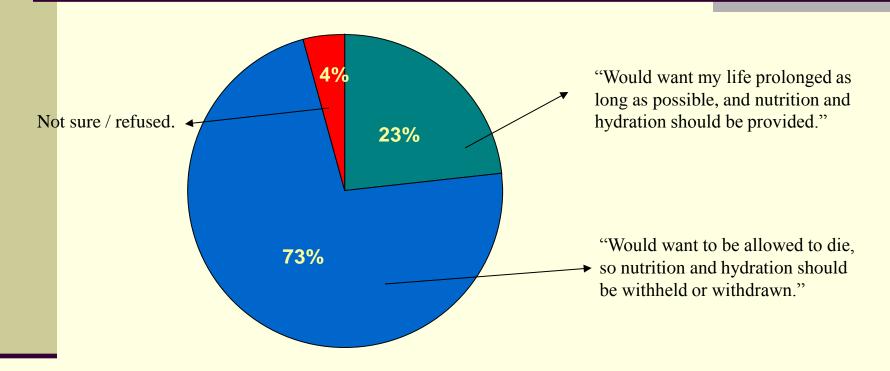
■ C: 50%

■ D: 23%

■ E: 32%



Preferences of Americans for receiving life-sustaining treatment if unlikely to regain consciousness



In a Spring 2005 national poll conducted shortly after the death of Terri Schiavo, nearly three fourths of those polled said that if they were unlikely to regain consciousness, they would want to be allowed to die and so nutrition and hydration should be withheld or withdrawn.



What do people say they want?

Three things that influence treatment decisionmaking:

Treatment burden

Treatment outcome

Likelihood of outcome



Patient Wishes

- Survey of Massachusetts seniors older than 50: 90% said they wanted honest answers from their doctors but fewer than 20% had discussed End-of-Life issues with their doctors (AARP 2005)
- Patient's with terminal lung cancer just slightly more than ½ had had their health care provider discuss hospice. (Archives of IM- 2009)



The approach....

Reductionist

- Switch to just comfort care
- Time to talk about pulling back
- Stop active treatment

Expansive

- Concentrate on maximizing comfort
- Do as much as possible to meet your needs
- Work as hard as possible so you can go visit your new great-grandbaby



HOPE

H: honest communication

- O: opportunities for reflection and considering options
- P: planning, illuminates one's potential

■ E: engagement, empowerment, explore



Drivers of Change

Incentive: economic and non-economic

- Actors: consumers, funders, professionals
- Mechanisms:
 regulatory,
 administrative/professional,
 market based



Changing Physician Practice

Strong research evidence

Clinical issues sharply drawn



Financial Incentives

- Pay for discussion: fear "death panels and rationing"
- Pay for performance such as decreasing length of stay, readmits, ICU days at end-oflife
- Incentivized clinical activities reached a plateau in the first year and quality of care remained higher year 2 to 4 but lower than projected



Non-economic Methods for Changing Professional Behaviors

- Minimal impact: one-off info session, intermittent performance feedback
- Bigger impact: consistent performance feedback, point of decision tools for patient and provider, training and supporting opinion lead, repeated educational intervention
- Biggest impact: strong evidence, identify and address barriers to change, multimodal approach and engagement of other health care professionals



Hospitalizations

Estimated 40% patients are sent to hospital 30 days before dying

Many in the last six months of their lives live in and out of the hospital, being treated by more than 10 physicians

30 day re-admissions cost Medicare \$12-15 billion



Transitions in Care Community Forum

Collaboration objectives...

- Hospital/SNF collaboration to establish transitions in care processes with SNF residents
- Partnership to improve staff skills in developing and implementing plans that support EOL wishes
- Spread the model to demonstrate costeffectiveness of this approach
- Establish community standard of care



Transitions in Care Community Forum

- Meetings every other month
- Open agenda
- Discuss successes and challenges
- Identify opportunities for improvement in communication
- Identify educational opportunities



Partnership for Patients

- Billion dollar quality initiative
- Support community coalitions
- Reduce 30 day readmission rate
- Significance:

cost and inefficiency of 30-day readmissions negative impact on the lives of patients with serious complex illnesses PC pt's have complex diseases



Data to support partnership...

- ■23.5% of patients admitted to SNF's were rehospitalized within 30 days
- Few nursing home residents have identified proxies for decision-making (18%)
- Many nursing home residents have a diagnosis of dementia (over 50%)
- Transfers near end of life are common & not always beneficial
- 13% of hospital deaths of people over age 65 are proceeded by a transfer from a nursing home



Significance...

- 38 % of SNF discharges are admissions to hospitals
- Up to 37% of SNF resident hospitalizations may be inappropriate/avoidable and costly
- 13% of CA hospital deaths are of residents transferred from SNFs
- SNF to hospitals transfers represent suffering and dislocation for residents
- Reducing transfers opportunity to improve quality and reduce costs



Tools...

- Advance Directives, 4/16 HCDD
- POLST document and policies
- Comfort Care guidelines
- S-B-A-R
- GO WISH
- INTERACT

