Do we really know what our patients want and do we respect it?

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Director, Respecting Patient Choices Program
Intensive Care Specialist, Austin Hospital, Melbourne
What are we talking about?

• A 76 yo father with disseminated cancer with progressive pneumonia in ICU on a ventilator after a MET call

• A 93 yo mother with dementia, from a nursing home, in hospital for 6 weeks with # NOF

• An 84 yo mother, bedbound & unable to speak after a stroke, heading for a PEG and NH that she has told her family she doesn’t want

• A 62 yo father with advanced cancer, who has requested NFR, but is resuscitated no form is completed
What is advance care planning?

... ‘a process, whereby a patient, in consultation with health care providers, family members and important others, makes decisions about his or her future health care, should he/she become incapable of participating in medical treatment decisions’.

P Singer et al 1996

Ethical principles

- autonomy
- informed consent
- beneficence vs non-maleficence
- dignity
- prevent suffering
The classic statement was by Justice Cardozo in the USA in 1914

“Every human being of adult years and sound mind has a right to determine what shall be done with his own body; and a surgeon who performs an operation without his patient’s consent commits an assault.”
Why is advance care planning important?

Most people (~ 85%) will die after chronic illness, not a sudden event.

Nearly half of us are not in a position to make our own decisions when we are near death.

Our family have a significant chance of not knowing our views without discussion.

A doctor who is uncertain about what to do, and who has to make a decision, will often treat aggressively.

Many of us will be kept alive under circumstances that are not dignified, frequently suffering and in a way that we would not have wanted.
The aims of advance care planning

- **Initiate conversations with adults regarding views about future medical care**
- **Assist individuals with advance care planning**
- **Make sure plans are clear**
- **Ensure plans are available**
- ** Appropriately follow plans**
Key Elements of the RPC Program

• We train doctors and non-medical staff
  – on-line and 1 day face-to-face training course
  – to discuss and facilitate advance care planning

• Implement system changes
  – medical records
  – process changes

• Complement existing legislation

• Health professional education
  – GP education toolkit
  – RACGP and RACP website
Case study of Patient AG

Patient Background

81yo man lives with wife, diagnosed with lung fibrosis in July 2004, and treated with steroids and oxygen. Poor response to treatment with increasing breathlessness, and increasing oxygen requirements.

Respecting Patient Choices

RPC was introduced.

• At his initial contact in July 2004 he declined.

• In January 2005 he was approached again, and he was able to state some of his wishes, including:
  • Not for intubation
  • Not for ICU and other aggressive management
• He discussed his wishes with his GP, and these were documented on the discussion card, but he did not complete a MEPOA (was in process of doing this) or SOC.

• He subsequently became acutely breathless at home and asked his wife to ring GP, not ambulance (as he would have previously done).

• His GP attended him and provided comfort care at home and he died.
Pilot study: impact

Patients recognise their right to make informed decisions now rather than just informing future requests

• "I want no further chemotherapy now",

• "I would like to have the tracheostomy out now and to stop ventilation”

• "I do not want the tracheostomy that is booked for tomorrow”

• "I want no further transfusions"

• "I want you to make me as well as possible so that I can get home for my daughter's 21st birthday before I die"
The impact of advance care planning on end of life care in elderly patients: randomised controlled trial

Karen M Detering, respiratory physician and clinical leader,1 Andrew D Hancock, project officer,1 Michael C Reade, physician,2 William Silvester, intensive care physician and director1

ABSTRACT
Objective To investigate the impact of advance care planning on end of life care in elderly patients.
Design Prospective randomised controlled trial.
Setting Single centre study in a university hospital in Melbourne, Australia.
Participants 309 legally competent medical inpatients aged 80 or more and followed for six months or until death.
Interventions Participants were randomised to receive
Randomised controlled trial

- English speaking, competent patients aged \( \geq 80 \)YO, admitted to hospital
- Intervention – ACP using the RPC model
- Primary outcome
  - compliance with patients EOL wishes
    - Wishes known & respected
Randomised Controlled Trial (Aug 07 – Mar 08)

1044 files / patients reviewed

↓

877 individual patients

↓

309 patients - informed consent obtained (35%)
154 intervention, 155 control

Excluded patients

• **not competent**, NESB, Prior ACP,
• expected to be discharged or die ,
• no family, refused
ACP patients (n=154)

- 80% Completed ACP
  - 38% appointed SDM + wishes
  - 32% Wishes alone
  - 15% SDM alone – ½ left doctor to decide

- Family involved – Yes in 74%
- Average time taken - 64 minutes
What decisions were made?

• On CPR and life prolonging treatments
  – Before discussions, 30 to 45% said no
  – After discussion:
    » “yes” changed to - “yes depending on outcome”
    - “I delegate the decision”
    - less “don’t know”

• Requests “If I become very unwell”
  for palliation and no aggressive treatment
  call my family, the priest, don’t let me suffer
Deceased patients

• 56 patients (18%) died within 6 months
  • 29 intervention, 27 control patients
• Patient’s wishes known and respected
  • Intervention 86%
  • Control 30%  \( p < 0.001 \)
• No difference in mortality between groups
• Location of death
  • Acute hospital: 16 in both groups
  • ICU: 0 intervention pt, 4 control pt \( (p = 0.03) \)
Patient / family feedback

• Discharge questionnaire
  • Control patients- negative comments
    • *the doctors don’t listen*
    • *I felt ignored and in the way*
    • *They don’t want me as I am too old*
    • *They wouldn’t speak to me, and kept discussing things with my family*
Impact of death on surviving relatives

- Death of a relative can cause significant anxiety, depression and post-traumatic stress

- How do you quantify the impact?
  - IES: Impact of Event Score (to assess PTSD)
  - HADS: Hospital Anxiety & Depression Score
## Deceased patients (56 patients)

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Deceased questionnaire

**Intervention group**

- *He had a very peaceful death, just as it should have been, & I would like to thank all staff for this.*
- *Even though we already knew what he wanted it was great to be able to talk about it so openly.*

**Control group**

- *Mum didn’t want heroics. I was horrified to hear she received 45 minutes of CPR. She didn’t want it. All anyone had to do was ask.*
- *The doctors kept asking if dad should be resuscitated. I didn’t think they should keep asking, as they also told us it wouldn’t help him. It was obvious to us he was dying.*
Conclusions from randomised controlled trial

- First RCT - coordinated ACP improves EOL care
- No difference in mortality
- Success of the RPC model (5 key elements)
  - Trained non-physician facilitators
  - Patient centred discussions
  - Involvement of family
  - Correctly filed documentation
  - Systematic doctor education
Benefits of ACP

- ACP improves EOL care and patient satisfaction
- ACP assists family to:
  1. know patient wishes, be involved in ACP discussions
     - More able to make decisions
     - Less burdened
  2. Have less risk of stress, anxiety and depression
  3. Be more satisfied with quality of patient’s death
RPC in 17 Hostels & Nursing Homes

- Began February 2004, 2 year evaluation
- 1108 residents
- Median age 86 (range 31-101)
- 76% female
- 37% competent
- 38% not competent
- 25% uncertain
What was the impact of RPC Program?

• 51% of 1108 residents were introduced
  – Of 565 introduced, only 12 residents refused further discussion
• Of those introduced 52% residents and/or families completed advance care plans
  – cf 3% of those not introduced ($p < 0.0001$)

• 42% completed by resident
• 58% completed by family on behalf of non-competent resident
What was requested?

- **90%** requested to receive **no life-prolonging measures**
- **87%** requested symptom and pain management
- **34%** requested to be **cared for at the facility** at end-of-life
  - 17% requested brief admission to hospital
  - 6% requested admission to hospital for aggressive treatment

- Many other personal requests
What were the outcomes?

• 16% of residents died during the 2 year evaluation
• 58% of the deceased residents had been introduced to RPC
• Of these, 89% had advance care plans (ACP)
  – cf 42% of those not introduced to RPC ($p < 0.0001$)
• 96-100% of their wishes were respected at EOL
• 85% of those with ACPs received EOL care in their facility
  – cf 33% of those without ACPs ($p < 0.0001$)
• Likelihood of dying in hospital is much greater without RPC introduction: 46% vs 18% ($p=0.0002$)
What was the GP involvement?

- GP involvement was encouraged, not mandatory
  - signed / witnessed 49% of 296 advance care plans
  - witnessed 60 MEPOAs
  - recorded advance care planning in 90 clinical records
  - actively participated in discussions with residents, NOK

- Involvement was more organised in some RACFs
  - benefit of aged care program managers

- Varied between GPs
What was the GP feedback?

• “this program is long overdue”
• “of practical value…involves the RACF staff in discussion”
• “having clarity of a resident’s wishes made decision making with the family much easier”
• “the GP information kit is very useful”
• “avoids residents being sent to hospital for EOL care”
Respecting Patient Choices (RPC)

- An Australian advance care planning program
- Commenced in 2002
- DoHA funding
  - Multiple aged care facilities
  - Community palliative care services
  - Health services in each state and territory
- Victorian Department of Health
  - 8 Victorian health services
  - Statewide ACP policy
International Society of Advance Care Planning and End of Life Care (ACPEL)

It gives me great pleasure to welcome you to the International Society of Advance Care Planning and End of Life Care. The idea of this international society was inspired by the exciting collaboration between like-minded people that arose during the preparation for the Inaugural International Conference on Advance...
International Society of Advance Care Planning and End of Life Care Conference 2011

22 - 24 June 2011
London, UK