

education for acute hospitals

## General strategies to facilitate hope and coping during prognostic and end-of-life discussions

Recommendation	Useful phrases (where applicable)
<ul> <li>Reassure the patient or caregiver that you (or a support system, service or team) will be there for them throughout the illness trajectory. If this is not possible, it is critical that alternative arrangements are spelt out and are reliable.</li> </ul>	"Our team [or whoever applicable] will do our best to support you throughout this illness."  "I [or our team or whoever applicable] will do whatever [I/we] can to assist you in whatever lies ahead for you." "You will not be abandoned." If the patient is no longer able to come for outpatient follow-up:  "I will be available to discuss your care with your [GP/community nurse/ palliative care team] even if you are not able to come and see me in the clinic."
Emphasise the available support, such as the palliative care team (or whatever other services are available in the local area).	"We've been talking about some treatments that are really not going to be effective now and that we don't recommend you use. But there are a lot of other things we can still do to help and support you and make sure you are as comfortable as possible." "Although this type of chemotherapy has not been useful, there are other treatments we can use to make you more comfortable." "The aim of treatment is changing more towards maximising your function and comfort." "I cannot give you any specific treatment to make this illness go away, but there is a lot we can offer to help you cope with it."
<ul> <li>Reassure the patient or caregiver that many treatments are available for controlling pain and other symptoms, where applicable.</li> <li>Some symptoms may be difficult to control, and therefore it is also important not to make unrealistic promises (such as to make the person free of pain at all times).</li> </ul>	"We have a lot of ways to relieve [pain/nausea/dyspnoea] and other symptoms."  "We will do everything we can to ensure you are as comfortable as possible." "We cannot promise that you will have no pain. However, we can control almost all of the pain almost all of the time."
Emphasise the available support, such as the palliative care team (or whatever other services are available in the local area).	

Clayton JM, Hancock KM, Butow PN, et al. Clinical practice guidelines for communicating prognosis and end-of-life issues with adults in the advanced stages of a life-limiting illness, and their caregivers. Med J Aust 2007; 286(12): S79-S108. © Copyright 2007 The Medical Journal of Australia – reproduced with permission.







## Useful phrases (where applicable) Recommendation "What are your most important [hopes/expectations] about the Where appropriate, explore and discuss realistic goals and future?" expectations: facilitate realistic goals and reframe the patient's "As you think about the future and that you may not have a very long and family's expectations. time to live, what is most important to you? Are there any aspects of your life that you want to attend to?"<sup>2</sup> "Have you any unfinished business?" "What are the things you most want to invest your time and energy in?" "What are the things you want to do in the time you have?" "Is there any particular event that you are looking forward to?" Identify areas where control can be fostered [eg, advance "What is your understanding of your health situation and what is care planning, enduring power of attorney (see also Advance likely to happen?" Care Planning recommendations), tidying up unfinished "Do you have thoughts about where things are going with your business, arranging to see people to complete conversations]. illness?" "Some of my patients tell me it helps to try to take one day at a Where appropriate, discuss ways of coping on a day-to-day time, and live for the moment as much as possible." basis (eg, taking one day at a time, using strategies that allow "You will have good days and bad days." an ongoing relationship with the family, such as writing letters "Many people find that it helps them to cope by trying to maintain or recording thoughts, focusing on important relationships). some [sense of normality/normal life] or having a routine." It is important to respect and be sensitive to the patient's "Be kind to yourself and do the fun things — it doesn't have to be ways of coping (eg, denial can be a useful coping mechanism). all about your illness." "We can prepare for the worst while hoping for the best." Recognise the spectrum of hope and that patients may simultaneously hope for cure as well as acknowledge the terminal nature of their illness. "I have no problem with you exploring alternative treatments. They Respect the patient's wishes to explore alternative or are not my area of expertise. However, if the treatments were a experimental treatments aimed at controlling the underlying magic bullet, we would be aware and we would also use them. I disease (provided adequate information has been provided to know they are usually expensive." enable the patient to make an informed decision: see "Everything has benefits and burdens. What are you hoping this 'Commencing or changing disease-specific treatments' and therapy will achieve?" 'Cessation of disease-specific treatments' regarding "You must do whatever you feel is important, but remember that treatment decisions). none of these things have been proven to work (or we would be using them) and many are very expensive."

- 1. Tulsky JA. Beyond advance directives: importance of communication skills at the end of life. JAMA 2005; 294: 359-365.
- 2. Schofield P, Carey M, Love A, et al. 'Would you like to talk about your future treatment options?' Discussing the transition from curative cancer treatment to palliative care. Palliat Med 2006; 20: 397-406.

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