

Palliative Care: A focus on identification – Workshop 4 SPICT4-All & Carers Identification

This workshop offered delegates information on identifying those who need more help to care for a family member or friend living with a terminal illness.

For more information on SPICT4-All and Carers Identification please contact:

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A key aspect of this workshop was to group delegates and ask them to work together to discuss a number of questions.

How will you use the tool?

- Identify expectations and plan forward
- To provide a baseline for continued clear assessment & reassessment
- Language used in the tool will help carers communicate with health professionals

What do you plan to do next after this workshop?

- Promote access to support staff when carers need a question answered
- CHAT structure for carers to use when phoning for help
- Anxiety is high – how can we reduce?
- SPICT 4All can help carers feel more in control
- Useful tool for nursing/care homes to use when new members of staff to consider patient’s needs
- Energise the team and move forward

What are the indicators?

Key messages and points identified

<ul style="list-style-type: none"> • 4 admissions to hospital in 12 months • Lack of continuity - different hospitals • “Carer” always with patient, not identified as such • Wheelchair for 4 months • Dichotomy – diagnosis increase in weight gain, decrease in muscle • Wife – despite ongoing support never identified as “carer” • Never entered into palliative care discussion despite deteriorating health • Description of state of health, variable within hours • Discussion with “carer” behind closed doors • Medical professionals did not join up dots (did not need tools) • Stats available, relate to over 65yrs • Diagnosis (late in day) – superb support thereafter • Lack of support with patient and carer - lead to potential crisis (financial, social, marriage) • Life on hold • “normalisation” should be everyday practice avoids crisis, use familiar language in conversation • Carers often ignore own health issues, following transition they are no longer “carer” • Carers assessment should be formalised, employability issues, peer support 	<ul style="list-style-type: none"> • Knowing what to do next after you have the information • Conversation around diagnosis should be normalised despite tools in place, symptoms not identified • Peer support is imperative • Education • Communication – needs to be early and meaningful • Language
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Challenges barriers identifying carers

- They don’t see themselves as carers – wording has been changed to “person you are looking after”.
- There is a drive to encourage patients to attend appointments with a supporter and acknowledging the potential of sharing the burden of information.
- The caring role is very gradual process.

For more information on identification tools please follow the link to view the Identification Tools comparator:

