PLANNING NEEDED FOR DEMENTIA PATIENTS

A study into hospital discharge planning for patients with dementia has uncovered a number of cases in which the system failed to take into account the needs of the family.

A study funded by Alzheimer’s Australia and undertaken by researchers from La Trobe University looked at the experiences of 25 carers of elderly family members who have the degenerative brain disease.

‘What usually happens with dementia – whether someone is living alone, with someone else or in a nursing home – is that a son, daughter or other family member becomes the primary carer,’ said Dr Les Fitzgerald, lead author of the study and acting Head of Nursing & Midwifery at La Trobe Rural Health School. ‘They are the ones who call around regularly and do the bulk of the caring.

‘This also means that they become heavily involved in care when their family member is hospitalised with, for instance, a broken hip.’

Dr Fitzgerald and his team wanted to know how well carers thought hospitals professionals prepared them for the ongoing care of their family member after discharge.

‘The critical issue that many of the carers we interviewed had was that they were not necessarily involved in planning for post-hospital care. They were often unaware of the existence of a discharge plan.’

Dr Fitzgerald said this complicated care once a patient left hospital.

‘For instance there were several examples of hospital staff not following routines that carers had set up around managing incontinence. Family carers often had spent a lot of time trying to establish routines around going to the bathroom at specific times. In hospital those routines were often not being followed and as a result incontinence became a problem for the family carer to manage after the person with dementia was discharged.

Another problem was that carers found it difficult to keep track of recommendations from different specialists who might see a patient.

‘Carers are extremely committed people who want to know what they can and need do to help their family member have a better quality of life but to do this they need to be more involved in the planning of in hospital care as well as better informed about post hospital care.’

Overall, Dr Fitzgerald and his team are recommending that discharge planning for family carers of people with dementia can still be improved. The major recommendation is to involve the family carer more in the plan of care.

‘When it comes to treatment and discharge plans, hospitals are good at treating the primary diagnosis, such as the fractured hip. What our study has identified is that managing the complicating factor of dementia on top of the primary admission diagnosis is difficult and family carers can assist. We are asking hospitals to consider that a key element in creating discharge plans for these patients is involvement of the family carer.’

Further information can be found at the La Trobe Rural School of Health: latrobe.edu.au/health/about/schools/la-trobe-rural-health-school