

Original Article

Exploring Perception of Terminally Ill Cancer Patients about the Quality of Life in Hospice based and Home based Palliative Care: A Mixed Method Study

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ABSTRACT

Objectives: The objectives of the study were to evaluate the perceptions and performance of terminally ill cancer patients regarding the quality of palliative care at different settings and to measure their quality of life (QOL) at the end of life.

Material and Methods: This comparative, parallel and mixed method study was conducted at the Community Oncology Centre, Ahmedabad, among 68 terminally ill cancer patients as per inclusion criteria who were receiving hospice (HS)-based and home (HO)-based palliative care under 2 months permitted by the Indian Council of Medical Research. In this parallel and mixed method study, qualitative findings were supplemented by quantitative data with both components executed simultaneously. Interview data were recorded by taking extensive notes during interviews along with an audio recording. Interviews were transcribed verbatim and a thematic approach was adopted. QoL questionnaire (FACT-G System) was used for the assessment of QOL in terms of four dimensions. Data were analysed using the appropriate statistical test on Microsoft Excel.

Results: The qualitative data (primary component) analysed under five themes – staff behaviour, comfort and peace, enough and consistent care, nutrition and moral support, in the present study favours a HS-based setting more than a HO-based setting. Among all four subscale scores, physical well-being and emotional well-being subscale scores were statistically significantly associated with the place of palliative care. Functional Assessment of Cancer Therapy-General (FACT-G) total score was high among patients getting HO-based palliative (mean = 67.64) care than HS-based palliative care (mean = 56.56) and the difference between total FACT-G scores was statistically significant (unpaired t-test = 2.20, P = 0.03).

Conclusions: Overall, with the primary component favouring HS care and higher scores obtained in HO-based patients, the present study advocates the necessity for palliative services to expand their coverage regardless of whether they are provided at HS or HO, as it has improved the QOL of cancer patients significantly.

Keywords: Cancer, Home-based care, Hospice, Palliative care, Quality of life

INTRODUCTION

Cancer is the second leading cause of death globally and is responsible for an estimated 10 million deaths in 2020.^[1] Globally, about one in six deaths is due to cancer. Approximately 70% of deaths from cancer occur in low- and middle-income countries.^[2] There are estimated 2.5 million people living with cancer in India, with every year about 700,000 new cancer cases registered.^[3] The International Agency for Research on Cancer GLOBOCAN project has predicted that India's cancer burden will nearly double in the next 20 years, from slightly over a million new cases in 2012 to more than 1.7 million by 2035.^[4]

It is estimated that over 40 million people require palliative care each year out of which over 20 million require palliative care near the end of life (EOL).^[5] It is estimated that in India, the total number of people who need palliative care is likely to be 5.4 million people a year.^[6] Although palliative care services have been in existence for many years, India ranks at the bottom of the Quality of Death Index in overall score and lacks the awareness and existence of hospice (HS) care. According to the Worldwide Palliative Care Alliance, although >100 million people across the world would benefit from HS and palliative care annually.^[7]

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Quality Of Care For The Terminally Ill

CO Houle



Quality Of Care For The Terminally Ill:

Unveiling the Energy of Verbal Artistry: An Emotional Sojourn through **Quality Of Care For The Terminally Ill**

In a world inundated with screens and the cacophony of quick connection, the profound power and mental resonance of verbal art frequently diminish in to obscurity, eclipsed by the continuous barrage of noise and distractions. However, nestled within the musical pages of **Quality Of Care For The Terminally Ill**, a interesting function of literary elegance that impulses with raw thoughts, lies an remarkable trip waiting to be embarked upon. Written by a virtuoso wordsmith, this magical opus manuals readers on a mental odyssey, lightly revealing the latent possible and profound impact embedded within the complex internet of language. Within the heart-wrenching expanse of this evocative examination, we can embark upon an introspective exploration of the book is main themes, dissect their captivating writing model, and immerse ourselves in the indelible impression it leaves upon the depths of readers souls.

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