Journal of Palliative Care & Medicine

Perspective Open Access

The Impact of Palliative Care on Managing the Emotional and Physical Challenges of Terminal Illness

Margaret Charles*

Department of Internal Medicine, Oregon Health and Science University, Portland, USA

Corresponding author: Margaret Charles, Department of Internal Medicine, Oregon Health and Science University, Portland, USA, E-mail: marchar@OHSU.edu

Received: 28-May-2024, Manuscript No. JPCM-24-145007; Editor assigned: 31-May-2024, PreQc No. JPCM-24-145007 (PQ); Reviewed: 14-Jun-2024, QC No. JPCM-24-145007; Revised: 21-Jun-2024, Manuscript No. JPCM-24-145007 (R); Published: 28-Jun-2024, DOI: 10.4172/2165-7386.1000S8010

Citation: Charles M (2024) The Impact of Palliative Care on Managing the Emotional and Physical Challenges of Terminal Illness. J Palliat Care Med 14:010.

Copyright: © 2024 Charles M. This is an open-access article distributed under the terms of the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited.

Description

A terminal illness diagnosis is a serious, life-changing situation that usually indicates that a patient has an incurable sickness or condition that will most likely cause them to pass away quickly. Neurodegenerative diseases such as Amyotrophic Lateral Sclerosis (ALS), severe respiratory illnesses, advanced-stage cancer, and latestage heart disease are common instances. Palliative care and medication are essential components of complete medical approaches due to the emotional, physical, and psychological impact that terminal illnesses impose. When a patient is diagnosed with an incurable disease, those who are closest to them will also experience significant changes. This type of information could cause a wide range of immediate reactions, from shock and disbelief to depression, anger, and fear. This traumatic experience points out the necessity of a responsive, caring support network that attends to the many needs of individuals impacted. The treatment of terminal illnesses often involves complicated symptom control. Patients may have severe pain, exhaustion, dyspnea, nausea, and other uncomfortable symptoms as their condition increases. Whatever of the diagnosis, palliative care is a specialty area of medicine that focuses on reducing the stresses, pains, and symptoms associated with a serious illness. Improving the patient's and their family's quality of life is the main objective of palliative care. Palliative care involves symptom management and assisting patients in managing their illness path with dignity and comfort, in contrast to curative treatments, which want to cure the disease. This includes psychological and emotional distress in addition to physical symptoms. For example, pain treatment varies according to each patient's needs and frequently requires a mix of prescription drugs, physical therapy, and complementary therapies like massage and acupuncture.

An individualized therapy also addresses exhaustion, respiratory problems, and nausea. Social workers, psychologists, and counselors frequently collaborate on palliative care teams, providing patients and their families' emotional support and counseling. In order to improve treatment acceptance and sustain a better quality of life, patients can benefit from early symptom management and psychosocial support. Palliative care serves as a connection between different medical professionals. Palliative care teams make sure that the patient's overall needs are effectively satisfied and that there are no gaps or overlaps in therapy by coordinating care. This includes arranging treatments, managing medications, and making sure that the patient's wishes and preferences are communicated to all healthcare providers. Advance care planning, which entails discussing and recording the patient's desires for future medical care, is made easier with the support of palliative care. This is essential to make sure that the patient's care is

in line with their values and preferences, especially if they are unable to express their choices for themselves.

The priority frequently switches from curative therapies to comfort care as an incurable disease increases. Palliative care treats physical symptoms like pain and dyspnea while also offering emotional and spiritual support to keep the patient comfortable and consoled. Hospice care, a form of palliative care intended especially for patients reaching the end of their lives, is the term used to describe this stage of treatment. When caring for patients who have reached the end of their lives, ethical dilemmas frequently come up, especially when it comes to choosing whether to continue or stop life-sustaining therapies. Palliative care teams are skilled in managing these challenging moral environments, guaranteeing that choices are made with the patient's best interests in mind and with respect for their autonomy. The perception of terminal disease and end-of-life care by patients and their families is significantly influenced by social factors. Palliative care professionals are taught to be attentive to social differences and to accept and appreciate other traditions and opinions. Cultural competency guarantees individualized care that respects the patient's values and history. Despite its advantages, palliative care has a number of drawbacks, such as inadequate patient and healthcare provider understanding and restricted access, especially in those in need or rural areas.

Furthermore, given the fact that palliative care can be helpful at any stage of a serious disease, it is frequently believed to be reserved for the last stages of life. Healthcare professionals need to receive more information and training in the concepts of palliative care in order to address these issues. Moreover, public educations initiatives can assist clarify errors and support the early incorporation of palliative care. Furthermore, increasing access requires policy changes that support the infrastructure and finance of palliative care services. Palliative care has a bright future ahead of it because of advances in science and technology that will help with better symptom management and care coordination. For example, telehealth has become an important resource for palliative care, especially for patients who live in remote locations. Patients who have reached the end of their lives may benefit from ongoing research into new medication and treatments. The treatment of terminal illness requires an extensive and sympathetic approach due to its severe difficulties. Palliative care is critical for the treatment of terminally ill patients because it prioritizes quality of life, emotional support, and symptom management. Palliative care guarantees that patients and their families have the most comfortable and dignified experience through terminal illness by collaborating with traditional medical methods and attending to their complete needs. It is our goal to improve the quality of life and support for Citation: Charles M (2023) The Impact of Palliative Care on Managing the Emotional and Physical Challenges of Terminal Illness. J Palliat Care Med. 14: 010.

Page 2 of 2

individuals dealing with the severe challenges of terminal illness as we continue to grow in our knowledge of and expertise in the provision of palliative care.