

Ethical Issues in Bone Marrow Transplantation: A Nursing Perspective

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Clinical research has paved the road for many advances in technology for the treatment of cancer. These technologies have given rise to questions about the value of treatments that simply extend life, often at considerable cost to the individual's quality of life. One such technology is bone marrow transplantation (BMT). Since the first reported successful cases in 1968 and 1969 (1), this treatment modality is now being offered to thousands of patients with advanced malignant disease. This article explores the ethical issues that are emerging with the use of autologous bone marrow transplantation (ABMT) — donation of one's own marrow — in the treatment of advanced cancers of the breast and ovary. ABMT has been considered for these types of cancers because of the poor long-term survival associated with standard therapy (2). Recent advances in the availability of colony stimulating factors to assist with marrow recovery post-chemotherapy has resulted in lower treatment-related mortality rates for this patient population (3).

A bone marrow transplant (BMT) program has been in existence at the Royal Victoria Hospital, a 600-bed McGill University teaching hospital, since 1982. In an effort to identify key ethical issues facing nurses caring for this patient population, focus group interviews were conducted with nurses on our 20-bed inpatient hematology/oncology unit and our ambulatory care setting known as the oncology day care (ODC). Approximately 20 patients are transplanted each year. The nursing care involved in the actual process of transplantation requires a three to four week hospitalization as well as numerous outpatient visits to ODC for blood count and symptom control monitoring. Patients entering the ABMT program have been previously hospitalized frequently for conventional chemotherapy followed in many instances by periods of febrile neutropenia. The demands placed on the patients and their families are high prior, during, and following the transplant.

It was within this context that focus group discussions of ethical concerns from the nursing staff occurred. The issues of informed consent, quality of life, allocation of resources, and concept of hope were highlighted and are explored in more detail as follows.

INFORMED CONSENT

Aggressive treatment protocols which include ABMT often involve serious potential complications such as infection, hemorrhage, pain, and peripheral neuropathies. Consequently, the patient and family need to have accurate information about these potentially life-threatening side effects (4). Simply, informed consent implies that the patient's permission to proceed with treatment is based on a full explanation of the available options and the risks and benefits of those options by the physician. This process should involve shared decision making between the patient and his/her physician, with the physician as information provider and the patient as decision maker (5). Carney (6) explored the process of informed consent from the perspective of both physicians and nurses caring for BMT patients. Results revealed that nurses were more concerned that patients know about the secondary effects and complications arising from transplantation, while physicians emphasized the options of treatment and possible outcomes with or without the transplant. This study also concluded that the physicians' perception of informed consent most closely matches the patients in that most patients chose the option of transplantation based on outcome, namely life or death. Downs (4) proposed that the differences in nurses' and physicians' perceptions of informed consent are in fact based on their different roles. Nurses are constantly preoccupied with management of complications during transplant while physicians are always involved with prescribing, monitoring, and evaluating the outcome of the treatment. She further states that

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