Knowledge, Attitudes, and Practice Behaviors of Oncology Advanced Practice Nurses Regarding Advanced Care Planning for Patients With Cancer

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Cancer is the second-leading cause of death in the United States (American Cancer Society [ACS], 2009). Unfortunately, many Americans with advanced cancers are dying with unmet needs, and they spend their last days of life in physical, psychological, emotional, social, and spiritual pain (Robert Wood Johnson Foundation, 2002). Nationally, an increasing trend has been seen in hospitalizations and intensive care unit admissions toward the end of life in patients with cancer (Earle et al., 2004, 2008; Sharma, Freeman, Zhang, & Goodwin, 2008). Eighteen to 20% of patients with cancer are given toxic chemotherapy during their last two weeks of life (Earle et al., 2004, 2008), and 14% of patients had new chemotherapy initiated during the last month of their lives (Braga et al., 2005). On the other hand, only 20%–40% of dying patients received hospice services (Earle et al., 2008; McCarthy, Burns, Ngo-Metzger, Davis, & Phillips, 2003). For those receiving hospice care, the average hospice stay was rather short (around 16 days) (Darwish-Yassine, Towns, & Finn, 2008; Temel et al., 2008), and many hospice referrals were made within three days of death (Earle et al., 2008).

Advanced care planning is a patient-centered communication process during which clinicians gently explore patients’ values, goals, and preferences regarding their future care needs, particularly concerning end-of-life care needs, to ensure their wishes are met should they lose capacity to make decisions (Kass-Bartelmes, Hughes, & Rutherford, 2003; Teno, 2000). For definitions of terms, see Figure 1. A study by Zhang et al. (2009) indicated that only 31% of patients with advanced cancer reported having baseline discussions about end-of-life care issues with oncology clinicians. However, 60%–70% of Americans are willing to discuss advanced care planning if asked by clinicians (Eidsness, Schelling, Young, ...