Advance Care Planning in a Primary Care Setting

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Abstract

Purpose: Advance care planning (ACP) allows an individual to discuss and document their personal preferences at end-of-life. ACP has been shown to improve communication and reduce discomfort for patients and their families. The literature supports utilizing formalized, multimodality training programs for healthcare providers in order to increase their confidence in initiating ACP discussions. These findings led to the initiation of an evidence-based practice project in a primary care setting with the purpose of increasing advance care planning discussions between providers and patients with the use of a standardized education tool.

Background and Significance: National regulations mandate that patients are provided information about advance directives in the healthcare setting, but completion rates are not monitored and continue to be low. ACP is now a billable service for healthcare providers, but it has not provided enough incentive to increase completion rates. Barriers for healthcare providers in the outpatient setting include lack of time, protocols, and lack of education on how to initiate and foster advance care planning discussions.

Methods: Healthcare providers in a primary care office attended a 15-minute structured educational session with and a toolkit was provided on the importance of ACP, how to initiate conversations with patients, and bill for the service. Participants completed a portion of the *Knowledge, Attitudinal, and Experiential Survey on Advance Directives* (KAESAD) survey assessing their confidence in ACP before and three months post intervention. Participant confidence (N = 6) in ACP was analyzed using the Wilcoxin test and descriptive statistics. The number of billed ACP services for the office was collected for four months post intervention and compared to the previous four months.

Outcomes: A significant increase in provider confidence after participating in a multimodality

education program was found in the results (Z = -2.21, p = .03). There was a 42.1% increase in the number of billed ACP discussions for the office in the four months post intervention.

Conclusion: The future desired state is that ACP discussions become standard practice in primary care leading to the completion of advance directives. This can be accomplished through formalized education sessions and resources for providers in order to increase their confidence in initiating ACP discussions with patients. The ultimate goal is to decrease unnecessary spending at end-of-life while improving patient and family satisfaction with the quality of care received at end-of-life.

Keywords: Advance directive, advance directives, advance care planning, primary care, intervention, education, communication

Advance Care Planning in a Primary Care Setting

Death is inevitable for all, but the discussions surrounding this topic continue to be difficult and are often postponed until it is too late. An advance directive is a written statement by a competent person that indicates their wishes regarding medical decisions at end-of-life if they are unable to communicate due to mental infirmity or unconsciousness (Congressional Research Services (CRS) & Library of Congress (LC), 2016). The two parts of an advance directive are the living will and the health care power of attorney. A living will is utilized when an individual is unable to communicate their wishes and inform health care provider about the type of medical care they wish to receive at the end-of-life (CRS & LC, 2016). The health care power of attorney identifies a health care proxy or decision maker in the event that the patient is unable to make decisions for themselves (CRS & LC, 2016). Advance directives promote patient autonomy, dignity, reassurance, and empowerment (Wissow et al., 2004).

Problem Statement

In the United States (U.S.), advance care planning rates are less than 15%, and healthcare providers have not made progress in improving these rates (Ramsaroop, Reid, & Adelman, 2007; Tung et al., 2014). It is estimated that 40% to 70% of medical inpatients are not capable of making medical decisions at end-of-life (Sullivan & Dickerson, 2016). If a patient has not previously documented their end of life wishes and is unable to do so, it is left up to the patient's family to make those difficult decisions. This can cause stress and uncertainty for those involved. Advance care planning (ACP) has been shown to improve communication, increase knowledge and respect for a patient's wishes at end-of-life, and improve patient and family satisfaction with care (Detering et al., 2014; Sullivan & Dickerson, 2016).

Laws regarding advance directives are individualized by state, but there have been a few

regulations passed by Congress regarding advance care planning (CRS & LC, 2016). In 1991, Congress approved the Patient Self-Determination Act (PSDA) which requires that all Medicare-participating healthcare facilities provide information to patients on advance directives to give patients autonomy over their care (Ramsaroop, Reid, & Adelman, 2007; Tung et al., 2014). These facilities are also required to provide community education on advance directives, provide information about health care decision making rights, ask all patients if they have an advance directive, educate staff and community on advance directives, and not discriminate against patients based on their advance directive state (Ramsaroop, Reid, & Adelman, 2007). The PSDA mandates that providers distribute advance directives at time of admission to the hospital and on initial receipt of care or plan of enrollment (CRS & LC, 2016). If advance directives are not distributed and documented per the PSDA, the Secretary of Health and Human Services can withhold payment of services to Medicare-certified providers (CRS & LC, 2016).

Additional laws include The National Defense Authorization Act which established a federal advance directive for military personnel and the Medicare Improvements for Patients and Providers Act (MIPPA) (CRS & LC, 2016). The MIPPA added end-of-life planning to the initial preventive physical exam for new Medicare patients (CRS & LC, 2016). The Centers for Medicare and Medicaid Services (CMS) rely on state agencies and internal contract review to monitor advance directive requirements (CRS & LC, 2016). The Institute of Medicine prioritized advance care planning in 2014. Healthcare providers can now bill Medicare for ACP discussions, but this incentive has not provided enough motivation to overcome the barriers (Sullivan & Dickerson, 2016). How advance directive information is addressed can vary, but there are common challenges such as the providers discomfort in talking about end-of-life issues as well

as the lack of time for discussion (Detering et al., 2014; Lund, Richardson, & May, 2015; Spoelhof & Elliott, 2012).

Purpose and Rationale

Advance care planning is an ongoing process that requires multiple conversations with family members and healthcare professionals (Lum et al., 2016). Primary care clinics have the ability to implement an intervention to engage patients in advance care planning (Lum et al., 2016). However, many providers find advance care planning conversations difficult and feel that they have inadequate training to assist in the process of end-of-life care communication (Clayton et al., 2012; Detering et al., 2014; Tung et al., 2014).

Advance directives are associated with lower levels of Medicare spending, decreased likelihood of in-hospital deaths, and increased hospice use in areas that have high levels of end-of-life spending (Rao, Anderson, Lin, & Laux, 2014). An advance directive allows a patient to have high quality care at end-of-life without unwanted or invasive treatments (Oczkowski, Chung, Hanvey, Mbuagbaw, & You, 2016). The purpose of this evidence-based practice project was to utilize a multimodality educational intervention for healthcare providers in a primary care practice to improve their confidence in their ability to have advance care planning discussions with patients and bill for the service.

Background and Significance

Many barriers that have been identified that may account for the low completion rates of advance directives. Prior to the 17th century, death was accepted as an inevitable part of life and people were prepared, however, life-prolonging technology in the 19th and 20th centuries improved the lives of humankind and attitudes changed (Sullivan & Dickerson, 2016). Death and dying became something to be ashamed of and was hidden to imply the possibility of

immortality (Sullivan & Dickerson, 2016). Federal funding for research is often focused on finding a cure instead of determining ways in which to manage chronic conditions (Sullivan & Dickerson, 2016). Living wills were initially developed by advocates of euthanasia, and advance care planning may be looked at by some as murder (Sullivan & Dickerson, 2016). Religious or spiritual beliefs may increase the rate of life-prolonging care along with the fee-for-service medical paradigm in the United States (Sullivan & Dickerson, 2016).

Effective communication is a major barrier to end-of-life discussion and can be attributed to a healthcare provider's lack of time and communication skills, lack of education on how to initiate discussions, lack of protocols, and inadequate mutual understanding regarding values and goals (Houben, Spruit, Groenen, Wouters, & Janssen, 2014; Spoelhof & Elliot, 2012). Historical, political, economic, and social beliefs all contribute to the low rates of advance care planning in the United States, and it is important to train healthcare providers on the ways in which to initiate these discussions in the primary care setting (Sullivan & Dickerson, 2016). Factors that may increase the likelihood of having an advance directive include presence of a chronic illness or condition, being 65 years or older, being Caucasian, being female, and having a higher income and education level (Rao et al., 2014). In addition to demographic barriers to completion of ACP, there are legal barriers to consider. Barriers include poor health literacy, health care agent restrictions, execution requirements, inadequate reciprocity, and religious, cultural, and social inadequacies (Hinders, 2012).

A systematic review and meta-analysis compared communication tools such as decision aids, structured meeting plans, and educational interventions to determine the impact on end-of-life decisions of adult patients in the ambulatory care setting (Oczkowski et al., 2016). Low quality evidence was found that supported structured communication tools to assist with end-of-

life discussions in the outpatient setting (Oczkowski et al., 2016). The lack of supporting evidence was thought to be related to the need for a large sample size over a prolonged period of time with follow-up (Oczkowski et al., 2016). A standardized approach to measuring outcomes and knowledge of advance care planning after implementing an intervention was found to be lacking and made it difficult to objectively compare the interventions (Oczkowski et al., 2016). In another systematic review, it was determined that a combined written, verbal, and video education was significantly more effective than a single written intervention (Durbin, Fish, Bachman, & Smith, 2010).

In a study of community-dwelling adults, lack of awareness was the most frequently reported reason for not having an advance directive but 67.8% reported having concerns about end-of-life care (Rao et al., 2014). The data suggested that most people prefer receiving this information from healthcare providers but their attitude and comfort levels could be a barrier (Rao et al., 2014). Patients have indicated that they would prefer to discuss this topic with a healthcare provider who has known them for some time and for the provider to initiate the advance care planning when they are in good health (Ramsaroop, Reid, & Adelman, 2007; Wissow et al., 2004). In an evaluation of a multimodality education program for healthcare providers it was determined that providers were significantly more confident in their abilities to discuss advance care planning after reading a training manual, viewing a DVD, and participating in an interactive e-simulation (Detering et al., 2014). In a study of internal medicine resident physicians, 70% of the residents reported a lack of confidence in discussing advance directives and avoided the conversation (Tung et al., 2014). Enhanced training for providers, workshops for all members of the health care team, non-physician ACP champions, and standardized documentation were proposed interventions (Tung et al., 2014).

Interactive interventions in which patients can ask questions have also been shown to be more effective than didactic interventions (Spoelhof & Elliott, 2012). Group visits to discuss advance care planning are an alternative option which allow patients to interact with one another and discuss end-of-life decisions in a safe and supportive environment (Lum et al., 2016). An intervention to improve advance directive completion rates in a primary care setting will need to first focus on provider education and comfort levels with the material. Overcoming barriers to starting these difficult conversations can be done through interactive education and standardized workflows. An example of a comprehensive resource is The Conversation Project (Bisognano & Goodman, 2013). When implemented within several health systems in La Crosse, Wisconsin, patients and families were routinely engaged in conversations regarding their wishes (Bisognano & Goodman, 2013). In a study of 540 adults who died from April 1995 to March 1996 in La Crosse, 85% of them had written advance directives (Bisognano & Goodman, 2013). Impacting advance directive rates will require increased education for providers to initiate the conversation and patient-friendly materials to further the discussion.

Internal Evidence

In a primary care setting within an Accountable Care Organization in Phoenix, Arizona, there were new efforts to increase advance care planning discussions during annual physical exams. The plan was to empower providers to not only initiate these conversations, but also properly document and bill for the service. The goal was that these conversations would meet quality measures, increase revenue, decrease end-of-life care costs, and improve quality of care for their patients at end-of-life. Within the Accountable Care Organization, there were 16,415 patients on Medicare, but only 1,590 advance care planning discussions were billed in 2016 (L. Donkerbrook, personal communication, April, 18, 2017). Only 9.68% of eligible patients

participated in advance care planning discussions, and there is a need for further evaluation and change in current practice (L. Donkerbrook, personal communication, April, 18, 2017). After review of the background and clinical significance of advance care planning along with effective interventions, the clinically relevant PICOT question was developed, "In patients in a primary care setting, does a multimodality training program for providers in comparison to current practice impact the completion rates of advanced directives after the completion of the program?".

Search Strategy

Electronic database searches were completed to obtain relevant studies to answer the clinical PICOT question. The databases searched for this literature review included CINAHL (Appendix A), PubMed (Appendix B), Web of Science (Appendix C) and Cochrane Library (Appendix D). Keywords included; advance directive, advance directives, advance care planning, primary care, evidence-based interventions, intervention, education, communication, conversation project, respecting choices, next steps, and prepared. The initial search of the keywords ('advance directive' or 'advance care planning') yielded 6,463 results in CINAHL (Appendix A), 953 in PubMed (Appendix B), 9,204 in Web of Science (Appendix C) and 752 in Cochrane Library (Appendix D). Keywords were searched independently and then combined using "AND" with additional keywords to obtain a manageable yield in each database.

The initial yield in CINAHL was narrowed by searching the original terms with AND 'education' AND 'primary care'. Limitations included published dates before 2012 and studies outside of the U.S. The final yield was 25 articles that were then reviewed for inclusion in the literature review (Appendix A).

The initial PubMed search was limited by searching the original terms with AND 'primary care' AND ('provider education' OR 'evidence-based interventions') with a yield of 17 articles (Appendix B). An additional search was conducted within PubMed using the original key terms AND ('conversation project' OR 'respecting choices' OR 'next steps' OR 'prepared') with the limitation of being published in the last 5 years. This resulted in a yield of 85 articles.

The Web of Science database search was limited by using the original key terms AND ('intervention' AND 'communication') and a timespan from 2012 to 2017 yielding 182 results (Appendix C). Article titles and abstracts were reviewed for inclusion in literature review. A search of the Cochrane Library initially yielded 25 results (Appendix D). Exclusions included published dates prior to 2012 and study protocols with a yield of 18. The only article that was relevant to the PICOT was excluded because it focused solely on hemodialysis patients.

Exclusion criteria in each database included published dates before 2012 apart from one landmark systematic review. Additional exclusion criteria were studies written in a language other than English, study protocols, doctoral dissertations, studies focused on patients under 18 years of age, or studies focused on specific patient populations (i.e. congestive heart failure, dialysis). A hand ancestry search was also completed and yielded one study. The majority were found to be irrelevant or outside of the published date guidelines. The ten studies chosen for inclusion were relevant to the PICOT question. They were independently reviewed for validity, reliability, and applicability (Appendix E).

Critical Appraisal and Synthesis

Ten studies were selected in this literature review. Each study was evaluated using Melnyk and Fineout-Overholt's (2015) rapid critical appraisal tool and was presented in evidence tables for analysis of data (Appendix E). The strength of the studies varied from level

one through level six evidence. There were three systematic reviews (SR) with two of them also being a meta-analysis (MA), three quasi-experimental studies (QE), one retrospective chart review (RCR), one qualitative systematic review, and two mixed-method research studies (MMR). There was minimal bias reported amongst the studies with only one study reporting that an author received royalties from the educational tool (Appendix E). Additional biases to consider were that several of the study's participants were volunteers and may have already had positive beliefs about advance care planning or wanted to provide socially desirable answers on surveys. There was moderate heterogeneity in the measurable outcomes for each study but homogeneity in the fact that they each assessed the effect of a structured tool to address advance care planning (Appendix F).

The sample size for each study was adequate and patient demographics were consistent in focusing on patients over 21 years of age and excluding studies or interventions that were disease-specific or involved psychiatric care planning (Appendix E). Interventions across all studies had at least two components, but there was a wide variety of measurement tools utilized. Eight of the studies did not report the specificities or sensitivities of their measuring tools (Appendix E). Seven studies assessed patient or provider satisfaction and confidence through pre and post questionnaires or phone interviews. Two studies focused on advance directive completion rates, and two studies measured both outcomes (Appendix F). Transferability may be impacted by the fact that only four of the studies were completed in the United States. Other countries may have different views on end of life, but the patient population and clinical settings are similar. There were significant results or positive outcomes in each study to support the use of multimodality education tools for advance care planning.

There is currently not a standardized approach for discussing or completing advance

directives. Healthcare providers are encouraged to initiate advance care planning conversations with their patients, but studies have indicated that lack of provider comfort and time are barriers. Healthcare providers currently do not receive additional training or tools to help them facilitate these important discussions. The evidence indicates that standardized, multimodality education tools can improve provider confidence in having advance care planning discussions. Additionally, standardized education tools for patients are shown to improve patient satisfaction and can positively impact the number of advance directives completed.

Conceptual Framework and Evidence-Based Practice Model

The self-efficacy theory was the conceptual framework that guided this evidence-based project (Appendix G). Self-efficacy is an individual's beliefs in their ability to complete an action (Resnick, 2014). Behavior is believed to be influenced by environment, person, and informational sources as well as self-efficacy and outcome expectations (Resnick, 2014). This theory was appropriate for this project design because the project was focused on increasing healthcare provider's confidence in having advance care planning discussions with patients. The project was designed to promote self-efficacy for healthcare providers while addressing current behavior and potential barriers.

The Model for Evidence-Based Practice Change facilitated this proposed practice change (Appendix H). The model guides healthcare professionals through a systematic process for change and supports a combination of quantitative and qualitative data, clinical expertise, and contextual evidence (Rosswurm & Larrabee, 1999). This model provides a six-step approach for implementing an evidence-based practice (EBP) that starts with identifying a need for change and ends with implementing and evaluating that change (Melnyk & Fineout-Overholt, 2015). The model provided a solid framework to guide this project through the steps of evaluating,

analyzing, and synthesizing the evidence for the purpose of designing a practice change (Melnyk & Fineout-Overholt, 2015). A need for increasing advance care planning discussions was identified within the Accountable Care Organization and the goal was to implement an intervention that would assist providers in having advance care planning discussions with patients and evaluate the outcomes.

Methods

Ethical Considerations

An approval letter from the Vice President of Case Management of the Accountable Care Organization was obtained (Appendix G). Arizona State University Institutional Review Board approved the project as an Initial Study (Appendix H). The practice was selected by the Case Management team at the Accountable Care Organization and the practice verbally consented to host the project. Informed consent was obtained from participants prior to their engagement in project (Appendix I).

Practice Setting and Participants

The project site was a primary care clinic in a metropolitan area in the southwestern United States. The office had two physicians, one nurse practitioner, two physician assistants, one registered nurse, one office manager, and several front and back office staff members. The educational session time was planned with the office manager and all members of the staff were invited to participate in the educational session over a lunch break.

The participants donated their time and did not receive any direct compensation for participating in this educational program. Inclusion criteria included providers and clinical staff who were English speaking and at least 18 years of age. Exclusion criteria were minors, adults unable to consent, and were non-English speaking. There were six participants.

Intervention

This project consisted of a pre-post design with an educational component and follow up data collection on billable advance care planning rates in an accountable care organization. Staff in an internal medicine office voluntarily attended a 15-minute structured educational session about Advance Care Planning. that included verbal and written material. The written materials included the *Five Wishes Physician Toolkit* and an additional information about the laws and regulations surrounding ACP, barriers, and a list of additional resources. Participants completed a Level of Confidence subscale of the *Knowledge*, *Attitudinal*, *and Experiential Survey on Advance Directives* (KAESAD) survey to assess their confidence in Advance Care Planning before the intervention and three months post intervention (Jezewski et al., 2005). The total time required to complete the survey before and after the completion of the project was approximately five minutes each.

The educational session provided participants with information about the background and importance of ACP, how to initiate conversations with patients, implement into practice, and bill for the service. The proposed program educated participants through verbal instruction and written materials. There was additional time allowed to answer questions about the education material and surveys.

Outcome Measures

The primary outcome of this project was to assess the change in participant confidence in advance care planning after the educational session. The secondary outcome was tracking the number of billed Advance Care Planning services for each provider. This data was collected from quality data reported to the organization from Medicare for four months post intervention and compared to the data four months before the intervention.

Instruments

A demographic questionnaire was developed to collect information on participant's demographics, role, and years of experience (Appendix J).

Staff completed a Level of Confidence subscale of the *Knowledge, Attitudinal, and Experiential Survey on Advance Directives* (KAESAD) originally developed by Mary Ann Jezewski, RN, PhD, FAAN to assess their confidence in Advance Care Planning (Jezewski et al., 2005). Permission was given by Dr. Jezewski to reproduce the instrument for the project with the understanding that it would not be published or attached to this final report. The portion of the survey utilized consisted of 10 questions on a five-point Likert scale designed to measure confidence in Advance Care Planning. The reliability and validity of the entire survey was established for use by registered nurses and not the general public.

Data Collection and Analysis

Post surveys were administered, and data collected from subjects before the intervention and three months post intervention. The rate of advance care planning codes billed by the providers was tracked through quality data reported to the organization's chief financial officer each month. All data analysis was performed using IBM SPSS® Statistics Version 24. Data was analyzed using the Wilcoxin test with intent to treat. Descriptive statistics and cross-tabulation were used to describe the sample and outcome variables. Non-parametric statistics were used to analyze the data using a two-tailed test and the critical value was set at p > .05.

Budget

Total costs for the project were estimated to be \$1365.00 USD (Appendix K). In-kind support in the amount of \$1215.00 USD was provided by the project site and Accountable Care

Organization. Education materials were \$5.00 per person. Lunch was provided by the DNP student during the education session and was \$10.00 per person.

Results

Demographic data

The respondents were clinic personnel at a primary care practice in Arizona (N = 6). The respondents were 2 (33.3%) males, 4 (66.7%) females and all Caucasian (100%). The average years of age of the respondents was 42.5 (SD = 19.25) and ranged from 25 to 68 years. The respondents were 1 (16.7%) Nurse Practitioner (NP), 2 (33.3%) Physician Assistants (PA), 1 (16.7%) Medical Doctor/Doctor of Osteopathic Medicine (MD/DO), 1 (16.7%) Office Manager, and 1 (16.7%) Medical Student. The level of education of the respondents were 2 (33.3%) Bachelor's degrees, 3 (50%) Master's degrees and 1 (16.7%) Doctorate degree. The average years of experience and the years in current role were both 13 (SD = 17.05) years with the same range of 1 to 35 years.

Outcomes

Confidence. A Wilcoxon test examined the results of the pre and post survey conducted to measure the provider's level of confidence related to advance care planning (Appendix L). A significant increase in provider confidence was found in the results (Z = -2.21, p = .03). Provider confidence in advance care planning improved after participating in a multimodality education program. The NP had a high initial score of 88% and the same score post intervention. The PAs and MD/DO all had improved scores post intervention. The PAs had initial scores of 62% and 86% which increased to 72% and 100% confidence respectively. The MD/DO had an initial score of 80% that increased to 84%. Years of experience or time in current role did not have any clinical significance.

Billed ACP codes. The billed ACP codes were measured from quality data reported to the organization from Medicare for four months post-intervention and compared to the data four months pre-intervention. The practice had billed for 19 ACP codes between June 2017 to September 2017 and 27 ACP codes between October 2017 to January 2018. Providers within the Accountable Care Organization (ACO) averaged 4.48 billed ACP codes during the four months pre-intervention. The selected practice had billed for 19 ACP codes accounting for 4.49% of the total codes billed within the organization. During the four months post intervention, providers within the ACO averaged 4.22 billed codes and the selected practice billed for 27 accounting for 6.63% of the total billed codes. There was a 42.1% increase in the number of billed ACP discussions for the selected practice in the four months post intervention.

Discussion

The evidence indicated that a formalized, multimodality training program would increase provider confidence in advance care planning. This evidence-based practice project was successful in significantly increasing provider confidence regarding advance care planning after a formalized education session. The rate of billed advance care planning rates within the practice increased post intervention. ACP is a billable service but was underutilized within the entire organization and can be attributed to lack of time, provider confidence, and patient resistance. This project was successful in addressing provider confidence surrounding advance care planning. The educational session provided them with the resources to help them quickly discuss the topic and ways in which to address a patient's resistance or concerns.

The goal of the organization was to meet quality measures, increase revenue, and decrease end-of-life care costs while improving quality of care at end-of-life. This project suggests that if providers receive education on advance care planning their confidence in

initiating conversations with patients improves and more patients will receive this important information. The developed education session is sustainable and could be easily implemented at other practices within the organization as the materials are both cost effective and time efficient. This project directly impacted the organization and providers and indirectly impacted patients and their families.

Limitations and Implications

One of the limitations of this project was a small sample size. The practice that was selected by the ACO to participate in the intervention was a small practice and each of the providers on site the day of the presentation participated. The initial aim was that a well performing practice would be selected and this project would be conducted as a pilot project for a larger roll-out within the organization. The educational session was short but the time with the participants was rushed because coordinating schedules over a lunch hour on a Friday afternoon was difficult. However, this time was selected by the practice as they felt it would be the best time.

The ACP billed rates were also only tracked by the organization for Medicare patients. ACP discussions may have occurred with patients who had a different insurance and there was not a way to track this information at this time. A limitation of the project was that there was a significant delay in how the data is reported back to the organization from Medicare so that only four months post data could be collected with the timeline of the project. A better measure of the impact of the project would have been to examine at data six months pre and six months post intervention. Also, implementing the project at the beginning of a calendar year and monitoring for a longer period of time would have allowed the data to be collected for a full year of annual Medicare physicals and compared to the previous year.

There is also the possibility that ACP discussions were initiated with patients but that the providers did not remember to bill for the service. While provider confidence increased in ACP it is difficult to assess if this directly impacted their practice. Future practice sites may benefit from weekly check-ins with providers and staff to answer questions, audit charts, and assess facilitators or barriers to ACP conversations with patients.

Material cost were minimal but may be a limiting factor for the organization or future practices. The educational packets were \$5 each and the organization would have to pay staff to participate in the educational session and block appointment times. The materials could be redesigned and reproduced at a lower cost but there would be initial costs associated with developing the material. It is important for the organization and providers to recognize billing for ACP has the potential to both increase revenue for the practice as well as improve quality of care for their patients at end-of-life.

Conclusion

Advance care planning has been shown to decrease unnecessary spending while improving patient and family satisfaction with the quality of care received at end-of-life. The future desired state is that advance care planning will become a standard practice in primary care. Death is both inevitable and unpredictable. Advance care planning discussions need to be initiated in the primary care setting long before a patient's health declines or they have an unexpected event. American society is focused on planning for the future but fails to plan for the end. Healthcare providers have a unique opportunity to encourage their patients to make decisions surrounding end-of-life. This will ensure that the care that each patient receives is within their wishes and can decrease the amount of stress and burden on friends and family. The literature and this project indicate that all healthcare providers can benefit from additional

education and training about advance care planning on how to initiate and have these critical conversations with patients.

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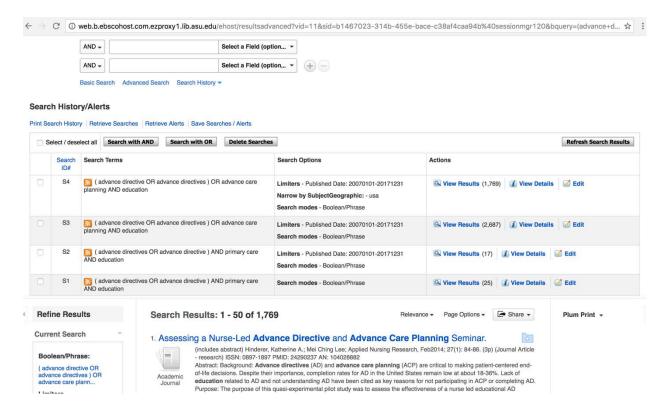
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Appendix A

Search Strategy 1

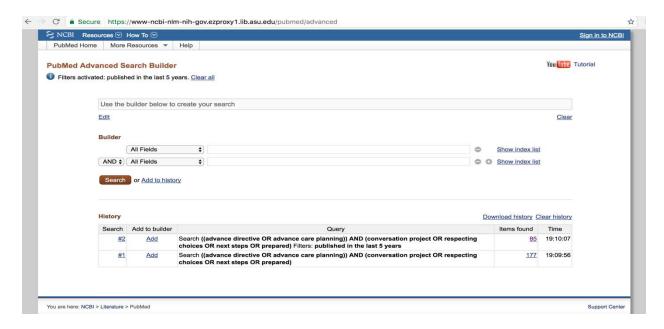
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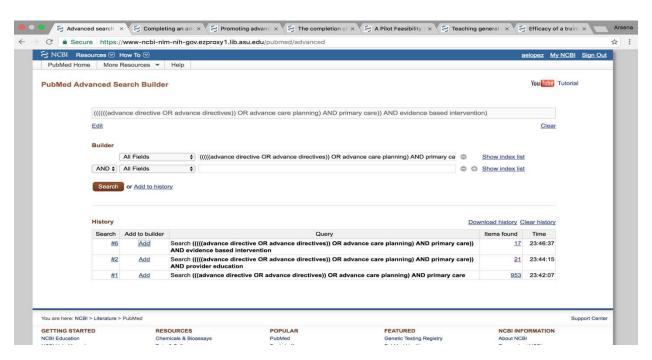
Appendix B

Search Strategy 2

PubMed 1



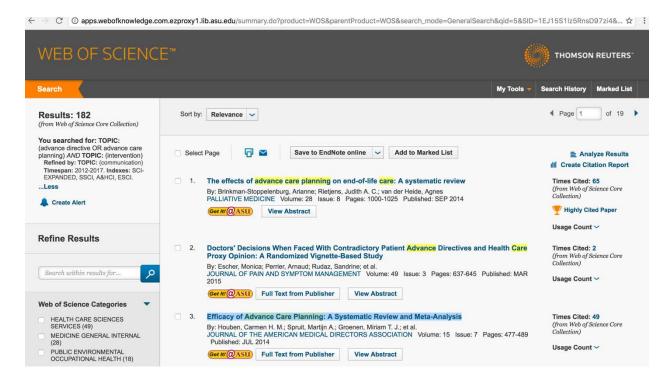
PubMed 2



Appendix C

Search Strategy 3

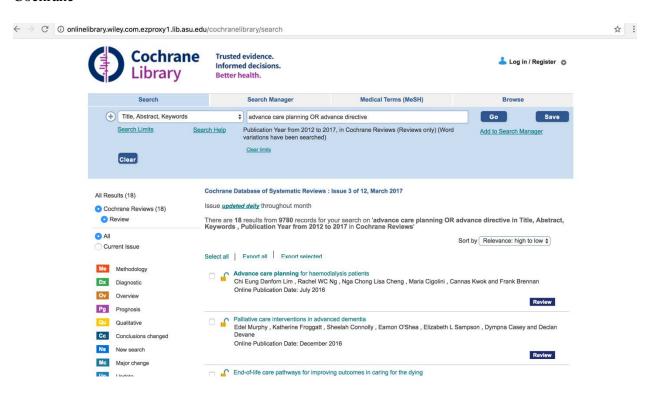
Web of Science



Appendix D

Search Strategy 4

Cochrane



Appendix E

Evaluation Table

Citation	Conceptual Framework	Design/ Method	Sample/ Setting	Major Variables & Definitions	Measurement	Analysis	Finding	Decision for Use
Butler et al.,	CLT	Design:	N= 16 studies	IV:	Semi-	NA	DV1: PF to	LoE: VI
(2014).		MMR	n=9 RCT	MyDirectives	structured		support	
			n=7 case	IV1: Five	telephone		decisions aids	Strength:
Decision Aids for		Method:	series	Wishes	interviews		(12)	Evaluated
advance care		Review of		IV2:				multiple ACP
planning: An		both gray and	Demographi	Consumer's	Standardized		DV2: PF to	tools
overview of the		published	c: Adult PTs	Toolkit for	data extraction		support clarity	
state of science		literature	with and	ACP	form by		(5- all video	Weakness:
		along with	without	IV3: EOL	IPDAS to		interventions)	Technical brief
Funded by MN		semi-	serious or	Decisions	collect			methods used
EPC under		structured	advance	IV4: Caring	information		DV3:	and outcomes
contract HHSA to		interviews	illness.	Conversations	about		Knowledge of	not
AHRQ		for technical		IV5: ACP	population,		AD or disease	synthesized,
		brief	Setting: OP	Conversation	decision aids,		process (9)	risk of bias not
Bias: Author			(15), IP(1)	Guide	and outcomes			assessed or
affiliations at		Purpose:		IV6: The			DV4: Medium	strength of
acponline.org/auth		Overview of	Exclusion:	Conversation	Technical		effect	evidence.
ors but unable to		advance care	Studies with	Project	brief methods		PREPARE and	
be viewed		planning	children,	IV7: The One	used		Making Your	Individual
		decision aids	psychiatric	Slide Project			Wishes Known	studies
U.S.		for adults	care	IV8:				measured
			planning, or	CRITICAL				satisfaction
			current EOL	IV9:				rather than
			decisions,	Preferred				IPDAS
			interventions	Priorities of				effectiveness
			without	Care				measures.

Key: ASA - agreed or strongly agreed; ACP - advance care planning; AD - advance directive; AHRQ - Agency for Healthcare Research and Quality; AR - attrition rate; AU - Australia; CLT - cognitive learning theory; CI - confidence interval; CST - Conversation Starter Tool; DN - authors declared none; EB - evidence based; ED - emergency department; EOL - end of life; EMR - electronic medical record; EPC - evidence-based practice center; ERA - Elder Risk Assessment; ES - effect size; FRNR - funders report no role in study design, data collection or analysis; GHS - Gundersen Health System; GRADE - grading recommendations, assessment, development, and evaluation; HC - healthcare; HCP - health care provider; HCP - healthcare provider; IM - internal medicine; IP - inpatient; IPDAS - international patient decision aids standards; LQE - low quality evidence; LoE - level of evidence; LS - Likert Scale; MA - meta-analysis; MBI - Maslach Burnout Scale; MMR - mixed methods research; MN - Minnesota; MNT - McNemar Test; N/A - not applicable; ND - not disclosed; NIHR CLAHRC - National Institute for Health Research Collaboration for Leadership in Applied Health Research and Care; NP - nurse practitioner; NPT - Normalization process theory; NR - narrative review; NRT - nornandomized trials; OP - outpatient; OR - odds ratio; PASW - Predictive Analytics Software; PC - palliative care; PEDRo - Physiotherapy evidence based database; PGY - post-graduate year; PF - positive findings; PT - patient; QE - quasi-experimental; QOE - quality of evidence; RC - Respecting Choices; RCR - retrospective chart review; RCT - randomized controlled trials; RN - nurse; SE - standard error; SR - systematic review; U.S. - United States; UTD - unable to determine; WSRT - Wilcoxon Signed Ranks Test

			educational component Inclusion: English studies that assessed effect of decision aid from 1990-2014 AR: n/a	IV10: PREPARE DV1: Satisfaction with decision aid DV2: Clarity of patient preferences DV3: Knowledge of AD and				Feasibility: Tools assessed were publically available and appropriate in ambulatory setting
Citation	Conceptual Framework	Design/ Method	Sample/ Setting	disease process DV4: Effectiveness of decision aids. Major Variables &	Measurement	Analysis	Finding	Decision for Use
Blackford et al., (2013). Facilitating advance care planning in community palliative care:	PC practice framework	Design: MMR Method: Multi-site action research with observation,	N=7 service locations n= 390 medical record audits n=27 semi-	Definitions IV:CST-ACP DV: PC team initiating ACP discussions in practice	60 hours of participant observations Medical record audits to identify potential cues	Directed content analysis NVivo Ver 8 (QSR International)	The tool was helpful reminder for PC team to initiate ACP. Experienced staff relied more on	LoE: VI Strength: Useful reminder and educational tool for HCP to initiate ACP
Conversation starters across the client journey		semi- structured interviews, medical record audits,	structured interviews n=35 focus group		for ACP Semi- structured interview with		clinical judgment and less- experienced PC staff	Weakness: May not be applicable to other

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Funding:	and focus	interviews	key	required more	cultures/setting
Department of	group	with PC staff	informants	ACP education	S
Health and	interviews		comprising	to develop	
Ageing,		Demographi	medical staff,	skills and	Practice
Commonwealth of	Purpose:	c:	RN managers,	confidence.	relevant tool
AU	Development	Majority of	NPs, and		but not
	and	care was	educators	Tool is a	validated
Bias: DN	implementati	provided by		starting point	research tool
	on of a tool	PC RNs. RN	Focus group	for staff.	
AU	to facilitate	managers (6),	interviews		Feasibility:
	ACP in	NP (6), PC	with PC staff	Participative	CST-ACP tool
	community	RN		natures of	can assist HCP
	palliative	specialists		action research	in developing
	care practices	(45), RN		the CST was a	relationship
		(33), and		practice-	with PTs to
		enrolled RN		relevant tool	facilitate ACP
		(14)		rather than a	discussions
				validated	
		Setting:		research tool	
		Community		and would	
		PC locations:		require more	
		eastern states		formal testing.	
		(5), urban (3),		_	
		regional sites			
		(2); northern			
		AU (2) with			
		service to			
		regional and			
		remote areas			
		Exclusion:			
		ND			
		Inclusion:			
		ND			

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			AR: ND					
Citation	Conceptual Framework	Design/ Method	Sample/ Setting	Major Variables & Definitions	Measurement	Analysis	Finding	Decision for Use
Clayton et al., (2012).	CLT	Design: QE	N=21	IV: Brief training	Pre and post video	PASW	DV1: Communicatio	LoE:III
Evaluation of a novel individualized communication-skills training intervention to improve doctors' confidence and skills in end-of-life communication Funding: Cancer Institute NSW Clinical Research Fellowship for author. Bias: ND AU		Purpose: Determine if an individualize d training program to discuss EOL issues will improve MDs confidence, communicati on skills, and attitudes	Demographi c: Junior- doctors Setting: Teaching hospital in AU Exclusion: MDs with competing commitments Inclusion: MDs who volunteered to internal advertisement s about study AR: 9%	program (3 teaching sessions, 2 individual sessions with expert facilitator and simulation) DV1: MD communicatio n skills DV2: MD confidence in EOL discussion DV3: Attitude towards psychosocial care DV4: MD sense of accomplishm ent	recorded consultation and deidentified questionnaires pre/post training measuring self-assessed confidence, attitudes to psychosocial care, and MBI Communication-21 specific skills rates on four-point scale and recoded using Cohen's Kappa with Landis and Koch's	DS to summarize demographics NPT to analyze pre and post self-reports (WSRT) and behavior/skills WSRT –ordinal items MNT-dichotomous items	communication is significantly improved Clarity of information (p<0.001), pacing of information (p=0.002), empathy and support (p=0.022) DV2: Confidence significantly increased (Z=-3.923, p<0.001) DV3: Attitude significantly improved (Z=-2.155,	Strength: Short total teaching time in comparison to other workshops with one-on- one and face- to-face interaction Weakness: Small, non- randomized sample, involvement of only one institution and lack of longer follow-up, participants were volunteers
				Cit	standard for interpretation		p=0.031) DV4: Personal accomplishme	Feasibility: One-to-one training intervention (1

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					Satisfaction: Four-point LS Self-assessed confidence: 15-item, five-point LS (Cronbach's alpha 0.934) Attitude: 20-item questionnaire using five-point LS Stress and burnout: 22-item MBI		nt significantly improved (p=0.043); no significant differenced in emotional exhaustion (p=0.115) or depersonalizati on (p=0.48)	hr) total teaching (3 hrs face-to-face and 2 hrs reviewing material); difficult to schedule times
Citation	Conceptual Framework	Design/ Method	Sample/ Setting	Major Variables & Definitions	Measurement	Analysis	Finding	Decision for Use
Detering, et al., (2014). Teaching general practitioners and	CLT	Design: QE Sampling: Convenience	N=148 n=69 HCP who participated	IV1: Completion of the multimodality education	Pre-education and post- education questionnaires	X ² tests- demographics Paired sample t test- comparing	DV1: Improved confidence in six of eight area surveyed	LoE: III Strength: Significant results with
doctors-in-training to discuss advance		Purpose: Develop and	in training and	ACP program (DVD,	E-simulation scores	knowledge	(p<0.001)	strong design
care planning: Evaluation of a brief multimodality education programme		evaluate an interactive ACP educational program for HCP	completed pre and post workshop evaluations	interactive e- simulation, 2 hr workshop and training manual)		MNT- general knowledge and attitudes/confide nce	DV2: Significant improvement of knowledge of ACP (p=0.001)	Weakness: Amount of missing data from surveys, participants volunteered,

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	Demographi	DV1: HCP	Wilcoxon-		unsure if less
	cs: Majority	confidence in	assessed e-	DV3: No	experienced
Funding:	40 years in	having ACP	simulation scores	significant	facilitators
Development and	age with	conversations		changes in	would produce
evaluation of Next	more than 10		Bonferroni	attitude	similar results
Step programme	years of	DV2: Change	correction to	(p>0.05)	
by Victorian	experience.	in ACP	adjust for		Feasibility:
Quality Council,	63% not	knowledge	multiple	DV4: 90% of	Practical,
Department of	trained in		statistical	participants	relatively
Health Victoria	AU.	DV3: Attitude	comparisons	ASA the	short, need
				workshop was	facilitators
Bias: ND	Setting: 8	DV4:	$\alpha \leq 0.05$	well presented,	
	workshops:	Satisfaction		88% pace was	
AU	Metropolian	with program		appropriate,	
	(4)			85% group	
	Rural (4)	DV5: E-		discussion	
		simulation		helpful	
	Inclusions:	performance			
	HCP and			65% ASA that	
	doctors-in-			reading	
	training			material was	
				useful, 85%	
	Exclusion:			DVD was	
	Not			valuable, 69%	
	completing			that e-	
	both pre and			simulation was	
	post survey			helpful	
	AR : n/a			DV5: (n=64)	
				completed e-	
				simulation at	
				least once,	
				(n=46)	
				completed e-	
				simulation	

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more than once

							more than once Score pre- education (7) and post- education (60) (max score=85) (p<0.001)	
Citation	Conceptual Framework	Design/ Method	Sample/ Setting	Major Variables & Definitions	Measurement	Analysis	Finding	Decision for Use
Durbin et al., (2010). Systematic review of educational interventions for improving advance directive completion Funding: ND Bias: ND U.S.	Clinical decision-making theory	Design: Systematic review Purpose: Analyze effectiveness of educational interventions for improving advance directive completion	N=12 studies n= 8 RCT n= 4 NRT Demographi c: Ages 26 to 93. Eight of 12 samples predominantl y women. Three studies Caucasian only, 2 majority African American, and 4 included both races. Setting: Two studies IP, 9	IV: Educational interventions IV1: written IV2: verbal discussions IV3: videos DV: AD completion	Systematic review based on Cochrane review and grading criteria Effectiveness of types of educational interventions	ND for individual studies $\alpha < 0.05$	In 3 RCTs combined written and verbal interventions were more effective than single written interventions in completion of AD (p<0.05)	LoE: I Strength: High quality evidence Weakness: Low number of studies for types of educational interventions Feasibility: Findings can be generalized to adult clinic outpatients and hospitalized elderly

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Citation	Conceptual Framework	Design/ Method	AD for general population AR: n/a Sample/ Setting	Major Variables &	Measurement	Analysis	Finding	Decision for Use
			nursing, medicine, and social work written in English, educational intervention, AD completion as outcome, and					
			(i.e. HIV). RCTs that lacked sufficient data. Inclusion: Studies in					
			Exclusion: Only dates 1991 -2009. Specific PT populations					
			OP, 1 community based.					

Houben, et al., (2014).	CLT	Design: SR and MA	N=56 RCT	IV1: Structured	Study details recorded using	X ² test- relationship	Agreement between 2	LoE: I
			Demographi	intervention	predefined	between	reviewers=0.8	Strength: SR
Efficacy of		Purpose:	c : Cancer (3),	tools focused	data	interventions	9	and MA of
advance care		Systematicall	Cardiac	on AD	abstraction		95% CI=0.86-	RCTs; Good
planning: A		y review the	diseases (4),	completion	form	IBM SPSS	0.93; p < 0.001	validity
systematic review		efficacy of	COPD (1),			statistics 21.0		
and meta-analysis		ACP	CRF (4),	IV2:	Quality-		DV1: OR 3.26;	Weakness:
		interventions	Dementia (1),	Structured	PEDro scale	Meta-analytic	95% CI 2.00-	Multiple
Funding: Grant of		in various	HIV (2),	tools to		technique used	5.32;	instruments
Lung Foundation		adult patient	Comorbiditie	initiate		random effects	p<0.00001	available to
Netherlands,		populations	s (24),	conversations		model in		assess quality
Leusden. FRNR.			nursing home	about EOL		RevMan 5.	DV2: OR 4.66;	of RCTs,
			residents (4),	planning			95% CI 1.20-	nature of
Bias: DN			elderly &			Pooled OR with	18.08; P=0.03	interventions
			healthy (8),	DV1:		95% CI		makes it
Netherlands			HCP(4), and	Completion			DV3: OR 2.82;	difficult to
			unknown (1)	of AD and		$\alpha \leq 0.05$	95% CI 2.09-	blind patients,
				EOL			3.79; p<0.0001	
			Setting: IP	discussions				Feasibility:
			(15), OP (37),					Applicable to
			both (4)	DV2:				practice
				Concordance				
				of pt				
			Inclusion:	preferences				
			Described	and delivered				
			original data,	care,				
			RCT, and	knowledge of				
			written in	ACP, EOL				
			English.	preferences,				
				quality of				
			Б.1.	communicatio				
			Exclusion:	n, satisfaction				
			Pts <21 years	with HC,				
			and studies	decisional				

Citation	Constant	Destroy	about psychiatric AD AR: n/a	conflict, use of HC services and symptoms DV3: Occurrence of EOL discussions	Management	Annalanda		Decision for
Citation	Conceptual Framework	Design/ Method	Sample/ Setting	Major Variables & Definitions	Measurement	Analysis	Finding	Use Use
Lund et al., (2015).	Normalizati on Process Theory	Design: Explanatory SR of	N=13 studies Demographi	IV: Trained staff utilizing a structured	Variable per study: individual	Data extraction developed by NPT	Data from individual articles was	LoE: V Strength: SR
Barriers to	Theory	qualitative	c: Adults in	approach to	measurement	111 1	not disclosed.	strength. Six
advance care		implementati	healthcare	ACP	tools not listed	Coding frame		Weakness:
planning at the		on studies	setting		but reported	and taxonomy	Interventions	Information is
end of life: An			without	DV: Effective	that they were	items related to	most likely to	lacking about
explanatory		Purpose:	mental health	ACP	highly	intervention	facilitate ACP	the process of
systematic review		Investigate	problems	discussions	structured	types, clinical	are those that	implementing
of implementation		barriers and	g wt HG			contexts, and HC	provide a	and delivering
studies		facilitators to implementin	Setting: HC: OP (8), IP (5)			systems	simple, structured tool	ACPs, inadequate
Funding: Author		g ACPs in	T. 1 .				for HCP to	reporting of
(Lund) is a post doctoral research		clinical	Exclusion: Interventions				take action.	intervention
fellow supported		practice	aimed at					design and methods of
by South Central			children and					evaluation
Strategic Health			those with					- , u1uu11011
Authority.			mental health					Feasibility:
(Richardson &			problems					Requires time
May) partly								and trained
supported by			Inclusion:					staff
NIHR CLAHRC			Implementati					

Decision for

ADVANCE CARE PLANNING

with a partnership on of between Wessex intervention and NHS intended to organizations and support ACP partners of University of AR: n/aSouthhampton. FRNR.

Bias: DN

United Kingdom

(6) Canada (2), AU (4) U.S. (1) Citation Design/ Major Measurement **Finding** Conceptual Sample/ Analysis Framework Method Setting Variables &

	Framework	Method	Setting	variables &				Use
				Definitions				
Oczkowski et al.,	CLT	Design: SR	N=76 studies	IV: Structured	SR using	Revman 5.3	Structured	LoE: I
(2016).		and MA	n= 46 RCTs	communicatio	GRADE with	software	communicatio	
				n tools for	GradePRO		n tools	Strength:
Communication		Purpose:	Demographi	EOL decision	software	95% CI	increased	Rigorous
tools of end-of-life		Determine	c: Patients	making and			DV1: ACP	search strategy
decision making in		effects of	age >50 with	ACP	Ottawa-	Clinical	(RR 2.31, 95%	with GRADE,
ambulatory care		structured	no specific		Newcastle	heterogeneity	CI 1.25-4.26,	wide variety of
settings: A		communicati	medical	IV1: Verbal	Scale for	reported using I ²	p=0.007, LQE)	interventions
systemic review		on tools for	conditions.	discussions	Cohort studies	calculations		
and meta-analysis		end of life	Many	IV2: Paper			Completion of	Weakness:
		decision	specifically	tools			AD (RR 1.92,	Lack of
Funding: Grant		making on	evaluated	IV3: Verbal			95% CI 1.43-	consistent
from Technology		completion	populations	discussion			2.59, p<0.001,	terminology
Evaluation in the		of ACP	with cancer,	with paper			LQE)	for
Elderly Network.			lung, heart,	tools				interventions
FRNR.			neurologic, or	IV4: Videos			DV2: AD care	which may
			renal disease.				desired and	have led to

Key: ASA - agreed or strongly agreed; ACP - advance care planning; AD - advance directive; AHRQ - Agency for Healthcare Research and Quality; AR - attrition rate; AU - Australia; CLT cognitive learning theory; CI - confidence interval; CST - Conversation Starter Tool; DN - authors declared none; EB - evidence based; ED - emergency department; EOL - end of life; EMR electronic medical record; EPC - evidence-based practice center; ERA - Elder Risk Assessment; ES - effect size; FRNR - funders report no role in study design, data collection or analysis; GHS -Gundersen Health System: GRADE - grading recommendations, assessment, development, and evaluation: HC - healthcare: HCP - health care provider: HCP - healthcare provider: HCP - health medicine; IP - inpatient; IPDAS - international patient decision aids standards; LQE - low quality evidence; LoE - level of evidence; LS - Likert Scale; MA - meta-analysis; MBI - Maslach Burnout Scale; MMR - mixed methods research; MN - Minnesota; MNT - McNemar Test; N/A - not applicable; ND - not disclosed; NIHR CLAHRC - National Institute for Health Research Collaboration for Leadership in Applied Health Research and Care; NP - nurse practitioner; NPT - Normalization process theory; NR - narrative review; NRT - nonrandomized trials; OP - outpatient; OR - odds ratio; PASW - Predictive Analytics Software; PC - palliative care; PEDRo - Physiotherapy evidence based database; PGY - post-graduate year; PF - positive findings; PT - patient; QE - quasi-experimental; QOE - quality of evidence; RC - Respecting Choices; RCR - retrospective chart review; RCT - randomized controlled trials; RN - nurse; SE - standard error; SR - systematic review; U.S. - United States; UTD - unable to determine; WSRT - Wilcoxon Signed Ranks Test

Bias: DN		IV5:	care received	relevant
	Setting:	Computer	(RR 1.17, 95%	articles being
U.S.	Ambulatory	programs	CI 1.05-1.30,	missed; review
	care	IV6:	p=0.004, LQE,	limited by
		multimodal	2 RCTs)	highly
	Inclusion:	interventions		heterogenous
	RCT or	IV7:		nature of the
	prospective	interventions		populations
	observational	directed at		and
	studies with	HCP vs PTs		interventions
	control group			
	published in	DV1:		Feasibility:
	peer-	Completion		Structured
	reviewed	of ACP		communication
	journals in			tools should be
	English. PTs	DV2: Quality		considered for
	> 18 years	of		ACP
	and evaluated	communicatio		discussions in
	a	n between PT,		practice
	communicati	family, and		
	on tool to	HCP; PT and		
	assist patients	family		
	in EOL	knowledge		
	decision	about EOL		
	making	care and		
	compared to	available		
	a control	resources; HC		
	group.	resource		
		utilization; PT		
	Exclusion:	and family		
	Duplicate	satisfaction		
	references,	with EOL		
	conference	care;		
	abstracts,	acceptability		
	study			

			protocols, outcomes of	of intervention				
			interest not	mici vention				
			specifically					
			stated,					
			pediatric					
			studies, no					
			comparison					
			arm,					
			qualitative, or					
			not based in					
			ambulatory					
			settings					
			AR: n/a					
Citation	Conceptual	Design/	Sample/	Major	Measurement	Analysis	Finding	Decision for
	Framework	Method	Setting	Variables &				Use
				Definitions				
Pecanac, K.	CLT	Design: RCR	N = 732	IV- RC in	Data collected	Stata's	DV1:	LoE: III
(2014).				community	from manual	TEFFECTS	$\beta = 0.026$	
		Purpose:	Demographi		chart review	program	SE = 0.06	Strength: RC
Respecting		Determine if	c: Decedents	DV1- AD	about		TS = 0.46	is culturally
Choices and		RC would	from 2005-	completion	prevalence	Regression	p = 0.648	sensitive and
advance directives		improve AD	2010	rates for	and utilization	analysis and	95% CI= -	respects
in a diverse		in a racially		whites	of ADs	matching to	0.09-0.14	individual and
community		and	Setting: 300-			analyze RC on		spiritual needs,
		ethnically	bed	DV2- AD	Data included	AD prevalence	DV2:	EB ACP
Funding: ND		diverse	Midwestern	completion	date of	Th	β=0.136	***
D.		community	metropolitan	rates for	admission and	Proportional	SE = 0.05	Weakness:
Bias:			hospital	racial/ethnic	discharge, age	difference tests	TS=2.53	RCR and
Author BJS is an			E1	minority	at discharge,	to compare	p = 0.011	difficult to
employee of GHS			Exclusion:	DU2. DU1	gender, length	consistency of	95% CI= 0.03-	know if
who owns the			Decedents	DV3: DV1	of hospital	wishes by	0.24	decedents were
copyright to RC.				and DV2	stay,	race/ethnicity		personally

He also a coauthor of RC and is entitled to a small royalty if a profit is made on the materials.			who died in ED Inclusion: Decedent >		diagnosis, unit where care was provided and race/ethnicity	$\alpha = 0.05$	DV3: β= 0.071 SE= 0.04 TS= 1.82 p= 0.069 95% CI= -	impacted by RC program, only included legal documents as
U.S.			18 years who died during hospitalizatio n 2005-2010 AR: n/a				9.5% CI= - 0.01-0.15	AD, data only collected in one hospital and did not include patients discharged from hospital
								Feasibility: Applicable to diverse community, cost of program that includes multiple components
Citation	Conceptual Framework	Design/ Method	Sample/ Setting	Major Variables & Definitions	Measurement	Analysis	Finding	Decision for Use
Tung et al., (2014).	CLT	Design: QE Purpose:	N= 144 Demographi	IV: Quality improvement workshop	Pre- and post intervention surveys to	Descriptive summary statistics for	Residents reported significant	LoE: III Strength:
Improved resident physician confidence with advance care planning after an		Assess internal medicine residents' ACP practices and	c: IM residents: PGY-1 (35), PGY-2 (39), PGY-3 (32).	(interactive presentation & chart audit)	assess ACP practices in clinic, barriers to effective ACP, confidence	survey results and chart audits Pearson chi- square test and 2- sample t test	improvement in confidence in finding AD in EMR (Scores 4.8 vs 6.46 on 7-point	Significant results and applicable to other settings, mandatory participation

ambulatory clinic	improve their	Avg age 28.4;	DV: IM	with ACP	Multivariate	Likert scale)	
intervention	confidence in	47% female.	resident's	tasks (8 items	logistic	p<0.001	Weakness:
	discussing		confidence	on 7-point	regression		HCP at single
Funding: Mayo	ACP	Setting:	with ACP	Likert scale)	models to	Improved	institution,
Clinic		Mayo	discussions		determine	confidence in 7	recall bias and
		Clinical		Patient's	association	of 8 ACP	respondents'
Bias: DN		ambulatory		classified	between ACP	related	tendency to
		clinic in		using ERA	completion and	practices	provide
U.S.		Rochester,		score	PT and HCP	following	socially
		MN			variables	interventions	desirable
						(p value range	answers, did
		Exclusion:			SAS version 9.3	0.028 to	no measure
		IM residents				< 0.002)	impact of
		who were			α < 0.05		intervention on
		away from					the quality of
		Mayo campus					ACP
		during					
		scheduled					Feasibility:
		sessions were					Multidimensio
		excused					nal educational
							intervention is
		Inclusion:					feasible way to
		Required for					improve HCP
		all IM					ACP
		residents					confidence.
		A.D., 00/					Generalizable
		AR: 8%					to other OP
							clinics.

Appendix F

Synthesis Table

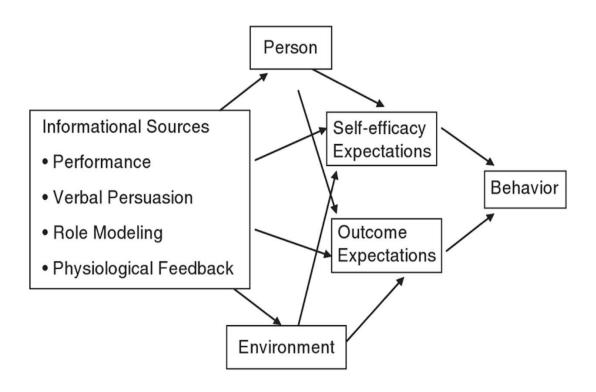
Author	Butler	Blackford	Clayton	Detering	Durbin	Houben	Lund	Oczkowski	Pecnac	Tung
Year	2014	2013	2012	2014	2010	2014	2015	2016	2014	2014
Level of Evidence	VI	VI	III	III	I	I	V	I	III	III
Design	MMR	MMR	QE	QE	SR	SR and MA	SR Qualitative	SR & MA	RCR	QE
Country	AU	AU	AU	AU	US	Netherlands	UK, CA, AU, US	US	US	US
Setting										
Inpatient	X (15)		X		X (2)	X (15)			X	
Outpatient	X (1)	X		X	X (9)	X (37)	X (8)	X		X
Community or Mixed					X (1)	X (4)	X (5)			
HCP focused education		X	X	X						X
E-Simulation			X	X						X
Lecture			X	X						X
Reading			X	X						X
Observation		X								
Individual training			X							X
Video				X						
Structured tools	X	X	X	X	X	X	X	X	X	X
Discussion	X (3)					X				
Written					X			X (9)		
Video	X (5)									
Computer Program	X (6)							X (4)		
Multiple interventions	X (2)	X	X	X	X	X	X	X (21)	X	X

Key: \mathbf{AU} – Australia; \mathbf{CA} – Canada, \mathbf{MA} - meta-analysis; \mathbf{MMR} - mixed methods research; \mathbf{RCR} - retrospective chart review; \mathbf{QE} - quasi-experimental; \mathbf{SR} - systematic review; $\mathbf{U.S.}$ - United States; \mathbf{X} - applicable to study; \uparrow - positive finding; * - significant; \neq - no significance

Outcomes Evaluated										
Patient satisfaction	^*					^*		^*		
AD Completion rates					^*	^*		^*	^*	^*
Rate of ACP Discussions		1				^*	^*			
Provider Pre/post survey			X	X						
Communication skills			^ *	^ *						
Confidence		↑	^ *	^ *						^*
Attitude			^ *	<i>≠</i>						^*

Key: \mathbf{AU} – Australia; \mathbf{CA} – Canada, \mathbf{MA} - meta-analysis; \mathbf{MMR} - mixed methods research; \mathbf{RCR} - retrospective chart review; \mathbf{QE} - quasi-experimental; \mathbf{SR} - systematic review; $\mathbf{U.S.}$ - United States; \mathbf{X} - applicable to study; \uparrow - positive finding; * - significant; \neq - no significance

Appendix G
Self-Efficacy Model



The Theory of Self-Efficacy (Resnick, 2014)

Appendix F

Model for Evidence-Based Practice Change

Model for Evidence-Based Practice Change Step 1: Assess the need for Step 2: Locate the best evidence change in practice · Identify types and sources • Include stakeholders of evidence Collect internal data about • Review research concepts • Plan the search current practice Conduct the search Compare external data with internal data Identify problem · Link problem, interventions, and outcomes Step 6: Integrate and maintain Step 3: Critically analyze the change in practice evidence · Critically appraise and weigh Communicate recommended change to stakeholders the evidence Integrate into standards of Synthesize the best evidence practice Assess feasibility, benefits, Monitor process and outcomes and risks of new practice periodically Celebrate and disseminate results of project Step 5: Implement and evaluate Step 4: Design practice change • Define proposed change change in practice Implement pilot study • Identify needed resources · Evaluate processes, outcomes, • Design the evaluation of the pilot and costs • Design the implementation plan Develop conclusions and

The Model for Evidence-Based Practice Change (Melnyk & Fineout-Overholt, 2015)

recommendations

Appendix G

Letter of Support



July 10th, 2017

To Whom It May Concern:

On behalf of Commonwealth Primary Care ACO, I am pleased to support the quality improvement project titled "Advance Care Planning in the Primary Care Setting" as proposed by Dr. Diane Nuñez, DNP, RN, ANP-BC, FNAP and Arseña Smith, BSN, RN, DNP graduate student.

Our organization agrees to serve as the quality improvement project site for data collection, education sessions, and data analysis.

Thank you for allowing the primary care practice(s) identified by the ACO to be a part of this evidence-based project.

Sincerely,

Susie Rodriguez RN, BSN, MHA Vice President of Care Management Commonwealth Primary Care ACO

4801 S Lakeshore Dr #106 Tempe, AZ 85282

Phone: 480-749-5000

Email: srodriguez@commonwealthaco.com

Appendix H

Institutional Review Board Approval Letter



EXEMPTION GRANTED

Diane Nunez CONHI - DNP 602/496-0751 DIANE.NUNEZ@asu.edu

Dear Diane Nunez:

On 7/27/2017 the ASU IRB reviewed the following protocol:

Type of Review:	Initial Study
Title:	Advance Care Planning in Primary Care
Investigator:	Diane Nunez
IRB ID:	STUDY00006510
Funding:	None
Grant Title:	None
Grant ID:	None
Documents Reviewed:	Smith_ACP_IRBApplication.docx, Category: IRB
	Protocol;
	Smith_ACP_CoverLetter.pdf, Category:
	Recruitment Materials;
	 Smith_ACP_RecruitmentFlyer.pdf, Category:
	Recruitment Materials;
	• Smith_ACP_EducationSessionOutline.pdf,
	Category: Technical materials/diagrams;
	• Smith_ACP_LetterOfSupport.pdf, Category: Off-site
	authorizations (school permission, other IRB
	approvals, Tribal permission etc);
	• Smith_ACP_ParticipantQuestionnaire.pdf, Category:
	Measures (Survey questions/Interview questions
	/interview guides/focus group questions);
	• Smith_ACP_DemographicQuestionnaire.pdf,
	Category: Measures (Survey questions/Interview
	questions /interview guides/focus group questions);
	• Smith_ACP_InstrumentPermission.pdf, Category:
	Off-site authorizations (school permission, other IRB
	approvals, Tribal permission etc);

The IRB determined that the protocol is considered exempt pursuant to Federal Regulations 45CFR46 (1) Educational settings, (2) Tests, surveys, interviews, or observation on 7/27/2017.

In conducting this protocol you are required to follow the requirements listed in the INVESTIGATOR MANUAL (HRP-103).

Sincerely

IRB Administrator

cc: Arsena Smith Arsena Smith

Appendix I

Participant Informed Consent

Advance Care Planning in Primary Care

June 18th, 2017

Dear Participant,

I am a family nurse practitioner graduate student under the direction of Professor Dr. Diane Nuñez, DNP, RN, ANP-BC, FNAP in the College of Nursing and Health Innovation at Arizona State University. I am inviting you to participate in an educational project regarding advance care planning in primary care.

I will be providing an educational session that will include a brief review of advance care planning and tools to assist primary care providers and staff members in having advance care planning discussions with patients during scheduled office visits.

The educational session and surveys will be delivered during a scheduled meeting for all participating staff members within the practice. The total time required to complete the survey before the presentation and at the completion of the project will be approximately 5 minutes each and the presentation will be 15 minutes. There will be additional time allowed to answer questions about the education material and surveys.

Your participation in the survey and education session is voluntary. You can skip any question on the survey and if you choose to not participate or withdraw from the program at any time, there will be no penalty. It will not affect your work or team role in the agency prior to, during, or after your participation in the program. You must by 18 years of age or older to participate. Responses to the questionnaires will be to gather demographic information, evaluate knowledge, attitude, and behaviors regarding advance care planning. There are no known risks associated with participation in this education program.

Your responses to the questionnaires are anonymous. The pre and post-survey responses will be linked using an identified code of the first four numbers of your mailing address but backwards. If less than four digits, remaining numerals will be replaced with zero(s). We will not collect your name or other personal identifying information. The results of this study will be used in reports, presentations, or publications as aggregate data only.

Attending the education session and completing the pre-education and post-education surveys will be considered your consent to participate. If you have any other questions concerning this program, please contact the following team members:

Dr. Diane Nuñez, DNP, RN, ANP-BC, FNAP at (602) 496-0751 Arseña Smith, BSN, RN, DNP Student at (928) 533-5444

If you have any questions about your rights as a subject or participant in this research, or if you feel you have been placed at risk, you can contact the Chair of the Human Subjects Institutional Review Board through the ASU Office of Research Integrity and Assurance, at (480) 965-6788.

Sincerely,

Arseña Smith, BSN, RN, DNP Student Dr. Diane Nuñez DNP, RN, ANP-BC, FNAP

Data Entry _____

Appendix J

Demographic Questionnaire



ID #_ 4-digit identified code

Data Analysis _____

Advance Care Planning: Demographics and Pre-Survey

Directions: Please answer the following questions. In the top right corner, please select and fill in an identified code that you will place on all surveys and questionnaires related to this project. The 4-digit code is the first four numbers of your mailing address but backwards, if less than 4-digits, remaining numerals will be replaced with zero(s).

with zero(s).
Please check the box to the $\underline{\text{left}}$ of your response or $\underline{\text{write in number}}$.
1. Age
2. Gender Male Female
3. Race/Ethnicity
□ Caucasian □ Hispanic/Latino □ Black/African American
□ Asian □ Native American □ Other
4. Highest Educational Level
☐ High School Degree ☐ Associate's Degree ☐ Bachelor's Degree ☐ Doctorate ☐ Other
5. Role
☐ Medical Assistant ☐ Registered Nurse ☐ Nurse Practitioner ☐ Physician Assistant
□ MD/DO □ Other
6. Years of Experience
7. Years in Current Role
For Official Use Only:

Data Validation _____

Appendix K

Budget

Table 1

Projected cost

Expense Items	Expenses	In-Kind Support
Personnel		
Project director (DNP Student)		
20 hrs @ \$40/hr		\$800.00
ACO Chief Operating Officer		
1 hrs/month x 2 months @		¢70.00
\$35/hr		\$70.00
Primary Care Providers (1) Initial training 1 hr @ \$75/hr		\$75.00
Advance practice providers (3)		\$73.00
Initial training 1 hr @ \$45/hr		\$135.00
Office Manager		4100.00
Initial training and organization		
of event 3 hr @ \$20/hr		\$60.00
Equipment/Materials		
Provider resource materials		
\$5/each x 10	\$50.00	
Office/Operations		
Utilizing physician's office for		
implementation of project		\$75.00
Lunch at initial meeting &		
training 10 people at \$10 pp	\$100.00	41417.00
Total Expenses	\$150.00	\$1215.00

Appendix L

Statistical Analysis

Table 2

Demographics

	Percent (%)	Frequencies	Mean (SD)
Age			42.5 (19.25)
Gender:			
Female	66.7	4	
Male	33.3	2	
Race/Ethnicity			
Caucasian	100	6	
Hispanic/Latino	0	0	
Black/African	0	0	
American			
Asian	0	0	
Native American	0	0	
Other	0	0	
Highest Level of			
Education			
High School degree	0	0	
Associate's degree	0	0	
Bachelor's degree	33.3	2	
Master's degree	50	3	
Doctorate degree	16.7	1	
Other			
Role			
Office Manager	1	16.7	
Registered Nurse	0	0	
Nurse Practitioner	1	16.7	
Physician Assistant	2	33.3	
MD/DO	1	16.7	
Medical Student	1	16.7	
Other	0	0	
Years of Experience			13 (17.05)
Years in Current Role			13 (17.05)

Table 3

<u>Pre and Post Intervention Confidence Score Res</u>ults^a <u>POSTTS-PRETS</u>

Z -2.207^a Asymp. Sig (2-tailed) .027

a. Wilcoxin Signed Ranks Test

b. Based on negative ranks