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OHRP meets ToS: Cloud-based technologies in human subject research

Assya Pascalev

Georgetown - Howard Universities

OBJECTIVES/SPECIFIC AIMS: To identify new ethical challenges in human subject research related to the use of cloud-based platforms for data collection. **METHODS/STUDY POPULATION:** Ethical analysis. **RESULTS/ANTICIPATED RESULTS:** The OHRP regulations protecting the data, privacy and confidentiality of human subjects and the Terms of Service regulations governing data use by cloud-based platforms are vastly different. The gap between these 2 sets of laws and regulations leaves human subjects vulnerable to harm during the data collection process via cloud-based tools. **DISCUSSION/SIGNIFICANCE OF IMPACT:** Recognizing the risks related to data gathering via cloud-based platforms, and educating researchers and research subjects about these risks and how to minimize them will strengthen the protections of participants and will enhance the informed consent process resulting in increased trust and greater willingness to participate in human subject research.

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Patient navigation training: Community-engaged workforce development

Nirmal Ahuja, Joanne Sullivan and Eugene Lengerich

Clinical and Translational Science Institute, Penn State University

OBJECTIVES/SPECIFIC AIMS: The goal of this initiative was to address this cancer health disparity in the Appalachian counties and help participants develop, implement and evaluate evidence-based "PN" that effectively and positively impacts patient and outcomes of the HealthyWomen Program. Following were the objectives of this training program: (1) To understand the broad range of roles and responsibilities associated with "PN", including care coordination and case management, in the Pennsylvania HealthyWoman Program in Pennsylvania and the National Breast and Cervical Cancer Early Detection Program. (2) To identify and assess local resources and expertise for evidence-based "PN" in the HealthyWoman Program. (3) To utilize "PN" in association with public education and targeted outreach initiatives in the HealthyWoman Program. (4) To implement strategies to manage and evaluate "PN" for the HealthyWoman Program. **METHODS/STUDY POPULATION:** The series of PN training was held at Pittsburgh, Camp-Hill, Wilkes-Barre and Philadelphia during June 2017. In total, 86 participants attended the training program at one of these 4 locations. Attendees represented organizations that provided breast, cervical and colorectal cancer outreach, screening and treatment. The participants of the training were solicited by regional program managers of the HealthyWoman Program of the PA Department of Health. The Harold Freeman model for patient navigation model was used to train the participants on the concepts of patient navigation. The training was built upon the Health Belief Model and Chronic Care Model, which defined the specific program constructs. The curriculum covered 2 important aspects, that is, clinical knowledge related to breast and cervical cancer along with aspects of patient navigation. Participants represented small, and large academic institutions/health care systems, cancer centers, federally qualified health centers, health departments as well as community-based screening programs and organizations. **RESULTS/ANTICIPATED RESULTS:** A total of 86 participants were trained; 78% had formal education and training in health-related field. In total, 62% of the participants had previous experience of patient navigation; 42% had training in social service related field and 50% had prior experience as community health worker. The demographic details reflected that majority of the participants (94%) were female. Most of the participants (30%) belonged to 50–64 years of age group followed by 30–39 years (23%) and 40–49 years (22%) of age group, respectively. As part of ethnic distribution, 70% of the participants were White Americans followed by Black/African Americans (17%). Furthermore, association of previous training in health and social service field with and without experience as a community health worker (n=84) and Navigator (n=86) was also analyzed. Among the participants, 44% had both community health worker experiences along with a prior training in social service related field whereas 42% of the respondents only had prior social service related training. This association of previous training in social service related field and prior community health worker experience was statistically significant with a p value of <0.05. Additionally, 81% of the participants who had previous experience in health-related training also possessed the prior experience as community health worker. Also, 81% of the participants who had previous training in health-related field also had a prior experience of patient navigation. In all, 38% of the participants who had a previous experience in social service field also had a prior experience of patient navigation. **DISCUSSION/SIGNIFICANCE OF IMPACT:** The training program established a pool of patient navigators which will contribute towards reducing the cancer health

disparity in the Appalachian region of Pennsylvania. The participants reflected a wide diversity in the navigators' backgrounds and differences across programs in their choices of patient navigators. It is important to consider this diversity when designing curricula materials and the methods of delivery in a patient navigation training program. As PN training programs are developed and implemented, further data is needed to guide practitioners and administrators in their efforts to include separate curriculum and materials for experienced and lay navigators. In addition, it is also important to assess the role and involvement of patient navigators in research and clinical trials. In total, 82% of the participants when asked agreed to be contacted for participation in research studies. Specific curriculum which includes research could be designed for further development of patient navigators. PN training and implementation knowledge is critical to the development of standards and best practices in this emergent area of cancer care.

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Perceptions of "translation" and the application of research across disciplines at the University of Michigan

Misty Gravelin, Meagan Ramsey, Kanchan Lota and George Mashour
University of Michigan School of Medicine

OBJECTIVES/SPECIFIC AIMS: There is no consensus on what constitutes translational research. To effectively support translation of research into practical settings, universities must determine who is involved, in which disciplines, and what results. In addition, it is unclear whether these researchers would see "translational research" as describing their work. **METHODS/STUDY POPULATION:** A survey assessing perceptions, successes, and barriers to the application of research was distributed to faculty, fellows, and graduate students within the University of Michigan. This survey included a question on the definition of translational research. **RESULTS/ANTICIPATED RESULTS:** Investigators of every rank and school participated (n=865), and all schools reported forms of applied research. Over 70% of participants said it was important to use research results beyond academia, and those responses represented diverse successes ranging from product development to artistic endeavors. Common barriers to such as lack of time and funding were also widely experienced. The definitions of translational research were divided between strictly health-oriented or broadly focused application. However, both definitions and familiarity with the term differed by field. **DISCUSSION/SIGNIFICANCE OF IMPACT:** Translation of research is widespread throughout the university, and many would define translational research to include their research discipline. Strategic university policies could benefit society by enhancing translation and application across many disciplines.

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Polypharmacy and patterns of prescription medication use among cancer survivors

Caitlin Murphy, Hannah Fullington, Carlos Alvarez, Simon C. Lee,
Andrea Betts, David Haggstrom and Ethan Halm
University of Texas Southwestern Medical Center Dallas

OBJECTIVES/SPECIFIC AIMS: The population of cancer survivors is rapidly growing in the United States. Long term and late effects of cancer, combined with ongoing management of other chronic conditions, make cancer survivors particularly vulnerable to polypharmacy and its adverse effects. We examined patterns of prescription medication use and polypharmacy in a population-based sample of cancer survivors. **METHODS/STUDY POPULATION:** Using data from the Medical Expenditure Panel Survey (MEPS), we matched cancer survivors (n=5216) to noncancer controls (n=19,588) by age, sex, and survey year. We defined polypharmacy as using 5 or more unique medications. We also estimated proportion of respondents prescribed specific medications within therapeutic classes and total prescription expenditures. **RESULTS/ANTICIPATED RESULTS:** A higher proportion of cancer survivors were prescribed 5 or more unique medications (64.0%, 95% CI 62.3%–65.8%) compared with noncancer controls (51.5%, 95% CI 50.4%–52.6%), including drugs with abuse potential. Across all therapeutic classes, a higher proportion of newly (≤ 1 year since diagnosis) and previously (> 1 years since diagnosis) diagnosed survivors were prescribed medications compared to controls, with large differences in central nervous system agents (65.8% vs. 57.4% vs. 46.2%), psychotherapeutic agents (25.4% vs. 26.8% vs. 18.3%), and gastrointestinal agents (31.9% vs. 29.6% vs. 22.0%). Specifically, nearly 10% of cancer survivors were prescribed benzodiazepines and/or opioids compared to about 5% of controls. Survivors had more than double prescription expenditures (median \$1633 vs. \$784 among noncancer controls). Findings persisted similarly across categories of age and comorbidity. **DISCUSSION/SIGNIFICANCE OF IMPACT:** Cancer survivors were frequently prescribed