

collection of primary data (through methods such as surveys and interviews) and secondary data (including historical and current census data). Primary data analysis for primary will use conventional quantitative and qualitative methods. Finally, researchers will use the analyzed data to develop a context-appropriate initial model of care for the ERC health clinic and a subsequent evaluation plan to assess the model's effectiveness and sustainability. RESULTS/ANTICIPATED RESULTS: All primary and secondary data will be synthesized into a report that will inform the development and implementation of two main deliverables. The first deliverable will be a framework for the clinic's initial model of care that is context-appropriate to the current needs of the Keith Neighbourhood. The model of care will be culturally sensitive and trauma-informed. The second deliverable will be an evaluation plan for the clinic that can be used to continuously iterate on the initial model, ensuring its sustainability. Furthermore, the project's process may be extrapolated into a framework that could be used to establish primary care clinics within other priority communities. DISCUSSION/SIGNIFICANCE: The proposed model of care will enable practitioners to deliver relevant and timely health services while being adaptable to the community's evolving needs. It will help improve the Keith Neighbourhood residents' long-term health and social outcomes. This project will contribute to and inform the development of the field of translational science.

268

The role of family and filial piety in serious illness and end-of-life decision making in the Chinese diaspora: an exploratory qualitative study

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OBJECTIVES/GOALS: The process of decision making in serious illness and end-of-life care needs to be culturally appropriate, relevant, and equitable. For Chinese Americans, family and filial piety have been shown to impact this process, yet it remains poorly understood how. The purpose of our study is to explore this decision-making process qualitatively. METHODS/STUDY POPULATION: We created semi-structure interview guides, based on a literature review of values salient to serious illness and end of life care for Chinese Americans. The guides will be used to conduct three focus groups, each with ten participants who identify as Chinese American stratified by highest attained familial role (grandparents, parents, and adult children). This will be done in partnership with Greater Boston Chinese Golden Age Center, a local community organization, in English, Cantonese, and Mandarin using our interview guides with live interpretation. Focus groups will be audiotaped, transcribed, and analyzed using thematic content analysis and modified grounded theory. RESULTS/ANTICIPATED RESULTS: We anticipate that we will identify how, by whom, and why Chinese Americans in Boston's Chinatown make decisions during serious illness and end-of-life. We anticipate that we will clarify cultural values, the balance between individual and collective values, intergenerational processes, individual and family suffering, and themes salient to this process. We expect to elucidate the role of family and filial piety during serious illness and end-of-life decision making and compare how these processes differ by generation and familial role for this

population. DISCUSSION/SIGNIFICANCE: Using the family as a unit of investigation and focusing on intergenerational processes represent a novel approach to understanding decision making during serious illness and end of life care for Chinese Americans.

269

Benefits of leveraging community-academic partnerships to plan and implement the Great Plains IDeA CTR Annual Community-Engaged Research Institute

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OBJECTIVES/GOALS: Building community-engaged research capacity is imperative to improve translation, but not everyone exhibits capacity to conduct research, especially community. We modified the research institute planning and implementation process between 2022 and 2023 to increase community appeal and engagement. METHODS/STUDY POPULATION: The 2022 and 2023 Community-Engaged Research Institutes (CERI) varied in their formats. For 2022, we hosted a one-hour planning session with community-academic dyads from our Community Advisory Board facilitated by the CEO director and co-director. The 2022 CERI consisted of five sessions totaling 12 hours. The first session was hybrid, followed by four virtual sessions. For 2023, we hosted two, facilitated one-hour planning sessions with community partners who attended the 2022 CERI, based on feedback and attendance from 2022. The 2023 CERI was consolidated to a seven-hour, one-day hybrid session. RESULTS/ANTICIPATED RESULTS: In 2022, two community partners participated as CERI panelists. There were 95 unique attendees spanning five days. Of the 95 attendees, 67% (n=64) were researchers and 33% (n=31) were community members. In 2023, eight community partners participated as CERI panelists and presenters. There were 57 unique attendees, of which 61% (n=34) were researchers and 39% (n=23) were community members. When comparing attendance for 2022 (29%, n=28) and 2023 (86%, n=49), in-person attendance was increased by 57%. DISCUSSION/SIGNIFICANCE: In 2023, we focused on research capacity building for community partners to align with community engagement principles and increasing research impact across the translational spectrum. Partnering with community elevated appeal for community participation and an increase in-person attendance.

270

Four Pillars of Community Health and Research Engagement: The UT Southwestern CTSA Program's Innovative Approach to Putting the Community Health Science Model into Practice

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OBJECTIVES/GOALS: Goals: Engage in collaborative approaches that target health disparities, especially among communities experiencing poverty or ethnic/racial minorities. Develop and sustain effective community-academic partnerships that foster trust through