Amplifying of the Patient Voice in PCOR Design and Implementation

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Amplifying of the Patient Voice in PCOR Design and Implementation

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Background:
• CRC screening rates are relatively low among Hispanics as compared to non-Hispanic whites and African Americans.
• Effective interventions are needed to raise CRC screening rates, reduce mortality, and increase survival in this growing segment of the population.
• Bidirectional engagement empowers patients to learn about research processes and inform the research question, design, intervention and dissemination.

Objective:
To engage patients in the design, development and implementation of patient-centered outcomes research (PCOR), specifically a multi-level decision support and navigation intervention aimed to increase colorectal cancer (CRC) screening in the Hispanic community.

Engagement Design:
Multi-stage Focus Groups (FGs) design:
• (4) grant development
• (2) pre-intervention
• (2) post-intervention
Continuous intervention oversight is conducted by a patient-led Patient and Stakeholder Advisory Committee (PASAC).

Setting:
Healthcare, Community and Church locations, urban community in Eastern Pa, served by large health network system.

Participants:
Autonomous patients and community members, self-reporting Hispanic, >/=50 years of age.

Data Collection/Analysis:
Audio recordings are transcribed and translated prior to coding process. Output is reviewed to identify common themes and responses. Basic descriptive statistics (frequencies) are used to identify key themes.

Ongoing Process:
While each focus group was a single meeting, the PASAC is convened 3x’s/yr for 2 hour meetings, and is sent monthly study updates via email. To date, 5 meetings have been held.

Preliminary Results:
An integral role in the research team, PASAC members report feeling that they are an extension of their community and have a duty to share their study knowledge with that community. As such, the committee provided strategies on how to disseminate findings in the larger community via various outlets. The PASAC members report feeling comfortable in the group and that their opinions are both heard and acted upon.

While members have provided feedback on study materials and protocols, most felt that their primary role was in disseminating the results to educate the Hispanic community about colorectal cancer screening. Fewer members acknowledged the role of the group in research design and implementation though several described themselves as part of the team.

Next Steps:
• The PASAC will continue to meet for the duration of the study.
• Two post-intervention focus groups will be conducted eliciting participant perceptions about intervention contacts and barriers to and facilitators of screening.

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