MEDICAL SCHOOL

Patient Centered Outcomes & Uptake of Health System Portal Use in Chronic Kidney Disease

June Tome¹, Shahbaz Ahmed¹, Marcio Mourao¹, Emily Chen², Sam Harrison², Angela Fagerlin^{3,4}, Julie Wright Nunes¹

MEDICAL SCHOOL 1. Department of Internal Medicine, University of Michigan, 2. Center for Bioethics Social Sciences in Medicine, University of Michigan, 3. Department of Population Health Sciences, University of Utah, 4. VA IDEAS, Salt lake city, Utah.

Background

 Web-based resources for chronic kidney disease (CKD) care and education are increasingly emphasized.

Objectives

 Examined if patient use of a specific web-based resource (patient portal) was associated with patient CKD knowledge, perceived CKD-specific stress, and patient self-ratings of health.

Methods

- A cross sectional survey was administered to adult patients with non-dialysis CKD during nephrology follow up visits (April 2015-March 2018).
- Logistic regression was used to examine the association between patient electronic health record portal use (outcome) and demographics, eGFR, patient CKD knowledge, perceived CKD-stress, and patient self-ratings of health assessed using validated questionnaires.

Results

Characteristics of Cohort:

- 245 participants, the mean age was 60 (SD ± 17) years. Eighty percent were white, 51% were women, 94% had >8th grade formal education, and 40% had < \$50,000 income.
- Summary scores for patient self reported health status was 1.74 (between very good and good, SD 0.95), accuracy score for CKD knowledge was 0.68 (scale of 0-100%, SD 0.15), and for CKD disease specific stress was 1.1 (0 representing no concern and 3 high anxiety, SD 0.67).

Health Portal Use:

• Sixty-seven percent used the portal, with 99% using it to check lab results, 82% to send messages to providers, 62% to review or renew prescriptions, 84% to make or change clinic appointments, 81% to view their medical history, and 72% to check information including patient educational resources.

Variable	OR Unadjusted (95% CI)	OR Adjusted (95% CI)
Age	1.0 (0.98-1.01)	0.98 (0.95-1.00)
Men (compared to women)	0.99 (0.57-1.72)	1.37 (0.69-2.74)
Race (compared to white)		
African American	0.39 (0.18-0.83)	0.69 (0.28-1.73)
Other race	0.88 (0.27-3.36)	3.90 (0.69-38.65)
< HS grad (compared to ≥ HS grad)	0.06 (0.00-0.34)	0.15 (0.00-1.11)
Income (compared to > \$50,000 annual income)		
< \$25,000	0.28 (0.13-0.61)	0.32 (0.12-0.78)
\$25,000-50,000	0.25 (0.12-0.52)	0.26 (0.11-0.58)
CKD stage (compared to stage 3)		
Stages 1-2	0.74 (0.30-1.91)	0.61 (0.17-2.14)
Stage 4	0.58 (0.30-1.10)	0.65 (0.30-1.41)
Stage 5	0.50 (0.20-1.29)	0.79 (0.23-2.78)
Health status	1.33 (0.99-1.82)	1.07 (0.74-1.53)
CKD knowledge	1.02 (1.00-1.04)	1.01 (0.98-1.04)
CKD stress	0.62 (0.40-0.94)	0.54 (0.31-0.93)

Table 2: Logistic regression model adjusted for age, sex, race, education, income, CKD stage, health status, CKD knowledge, and CKD stress

Conclusion

 Patient health portals are less likely to be used by patients of lower income and those with more kidney disease specific stress.

Limitations

- Cross sectional study limiting interpretation of causality.
- Single-center study at outpatient ambulatory nephrology clinics within one academic medical center, limiting generalizability.

Future Directions

 With an increasing emphasis on the use of health portals to improve patient engagement, interventions are needed to address disparities between portal users and non-users and ensure that promotion of patient portals does not widen gaps in support for those patients who may need it most.

There are no conflict of interest disclosures for any of the investigators.