

# Inpatient Resource Use and Cost of Hospitalization for KTP Surgery Among Recipients with vs. without Autosomal Dominant Polycystic Kidney Disease

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## Background

- Autosomal dominant polycystic kidney disease (ADPKD) is a genetic condition accounting for 5-10% of patients diagnosed with end-stage renal disease (ESRD) in the US and Europe.<sup>1-3</sup>
- While dialysis prolongs life, kidney transplant (KTP) is the first-line treatment for patients with ESRD due to ADPKD.<sup>4</sup>
- ADPKD patients have a higher rate of kidney transplantation (KTP) within the first year of initiating dialysis compared to the total ESRD population in the US.<sup>1</sup>
- Inpatient resource use and cost outcomes of patients with ADPKD and receiving KTP are limited.

## Objectives

- To assess differences in patient demographics, comorbidities, inpatient resource use, and cost outcomes among KTP recipients with vs. without ADPKD at date of KTP.

## Methods

### Study Design & Data Source

- A case-cohort analysis of patients with hospitalization for KTP surgery between 01Jan2018-31Dec2018 in the Premier Healthcare Database (PHD). (Figure 1)

### Sample Population

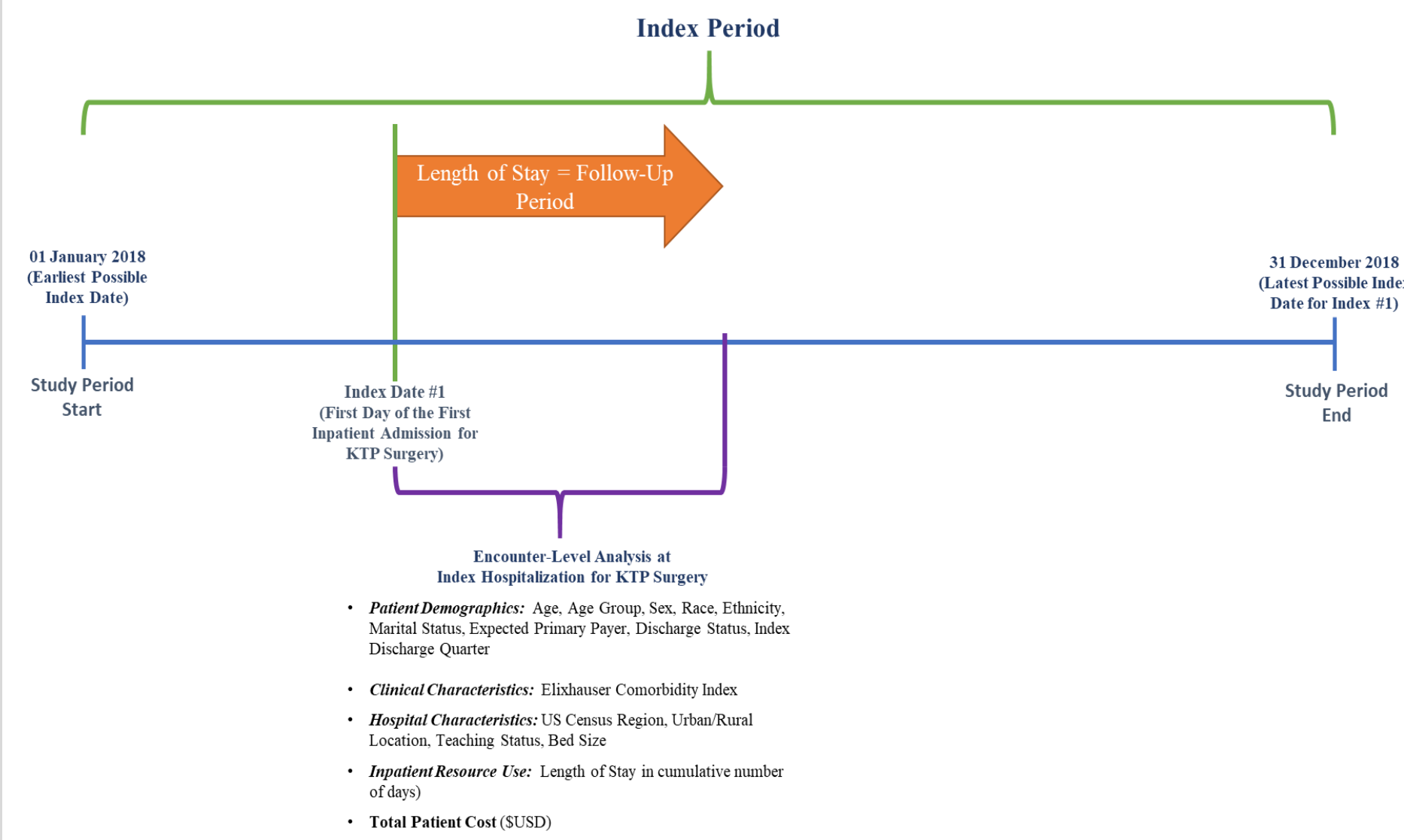
- Inclusion Criteria:** Inpatients  $\geq 18$  years old at KTP and distinguished as cases if presence of ADPKD and/or PKD-Unspecified was observed.
- Exclusion Criteria:** Inpatients were excluded if autosomal recessive polycystic kidney disease (ARPKD) diagnosis was observed.

### Outcomes

- Patient Demographics:** Age, Age Group, Sex, Race, Ethnicity, Marital Status, Expected Primary Payer, Discharge Status, Index Discharge Quarter
- Clinical Characteristics:** Elixhauser Comorbidity Index
- Inpatient Resource Use:** Length of Stay in cumulative number of days
- Total Patient Cost** (\$USD)

## Methods (Continued)

Figure 1. Study Schematic



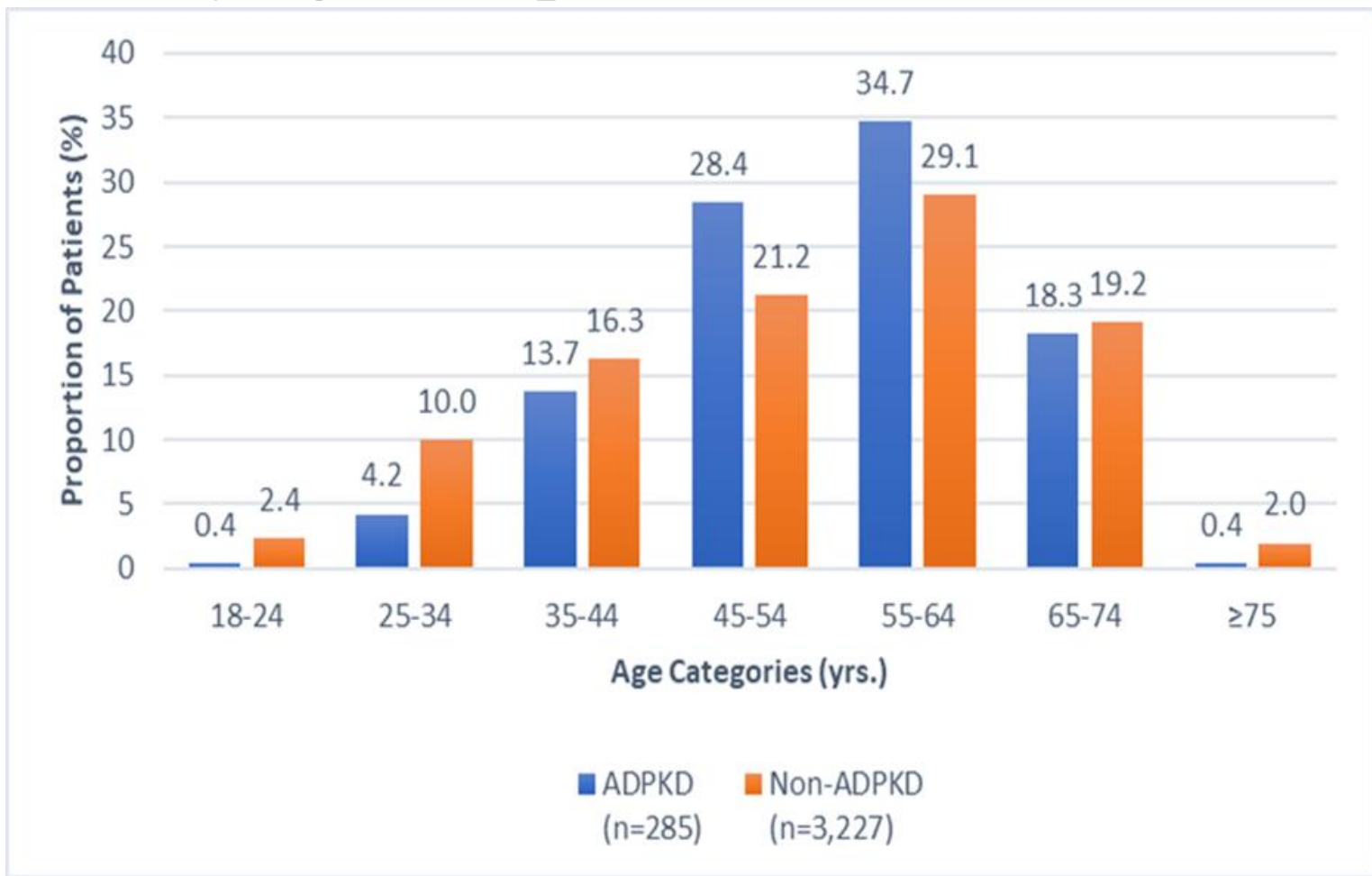
### Descriptive Analyses

- Demographics, comorbidities, length of stay [LOS], and total patient cost at hospitalization for KTP surgery were compared for those with vs. without ADPKD using the chi-square to test proportional differences and the Wilcoxon Signed-Rank Sum test to test median differences with alpha level set at  $\leq 0.05$ .

## Results

- Among 3,512 KTP recipients (ADPKD=285 vs. non-ADPKD=3,227), there was no difference in median (IQR) age (56 [47-62] vs. 55 [43-63] years old;  $p = 0.1658$ ).
- A higher proportion of KTP recipients with ADPKD were aged 55-64 (35% vs. 29%) and 45-54 (28% vs. 21%) years old ( $p < 0.0001$ ). (Figure 2)

Figure 2. Proportion of Recipients with vs. without ADPKD by Age Group

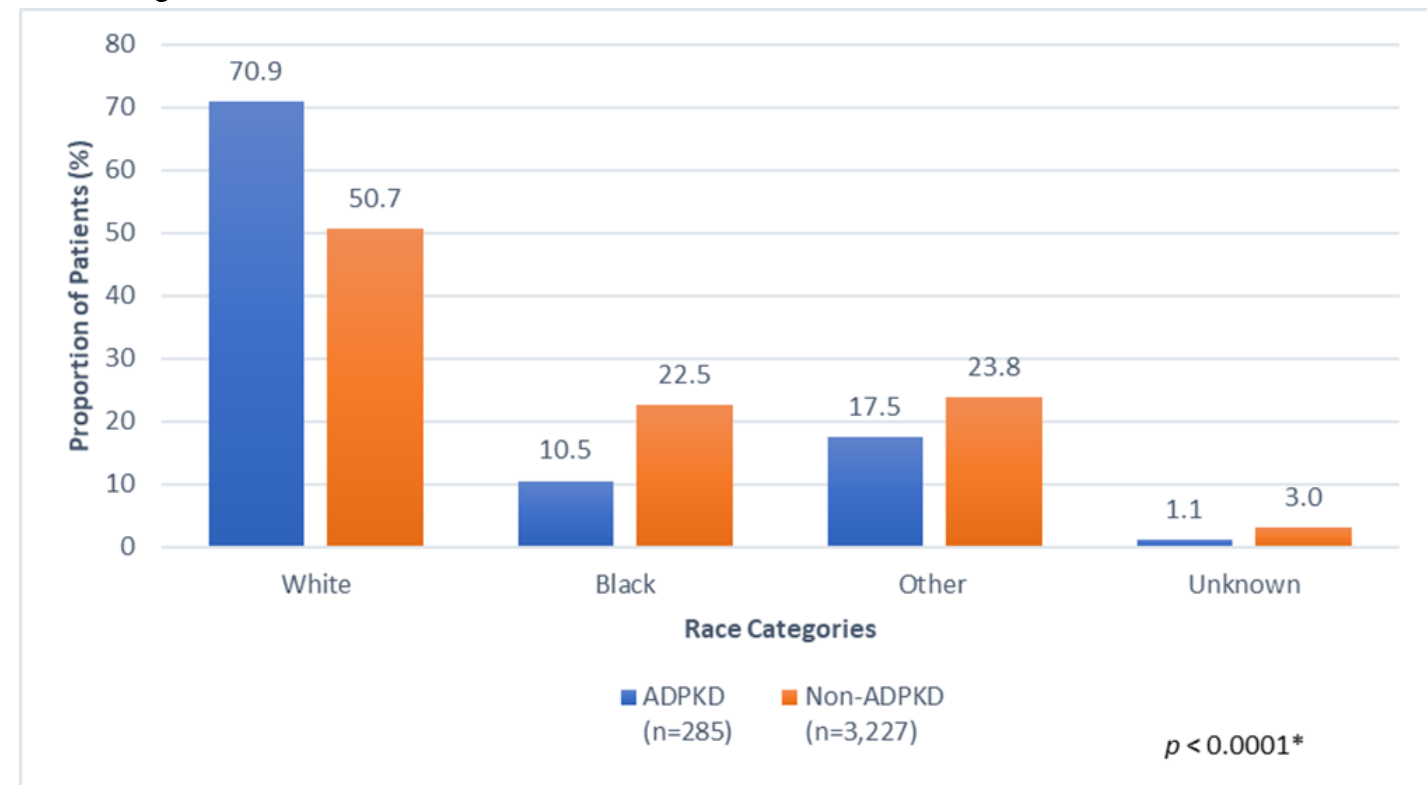


- A higher proportion of KTP recipients with ADPKD were female (46% vs. 38%;  $p = 0.0050$ ).

## Results (Continued)

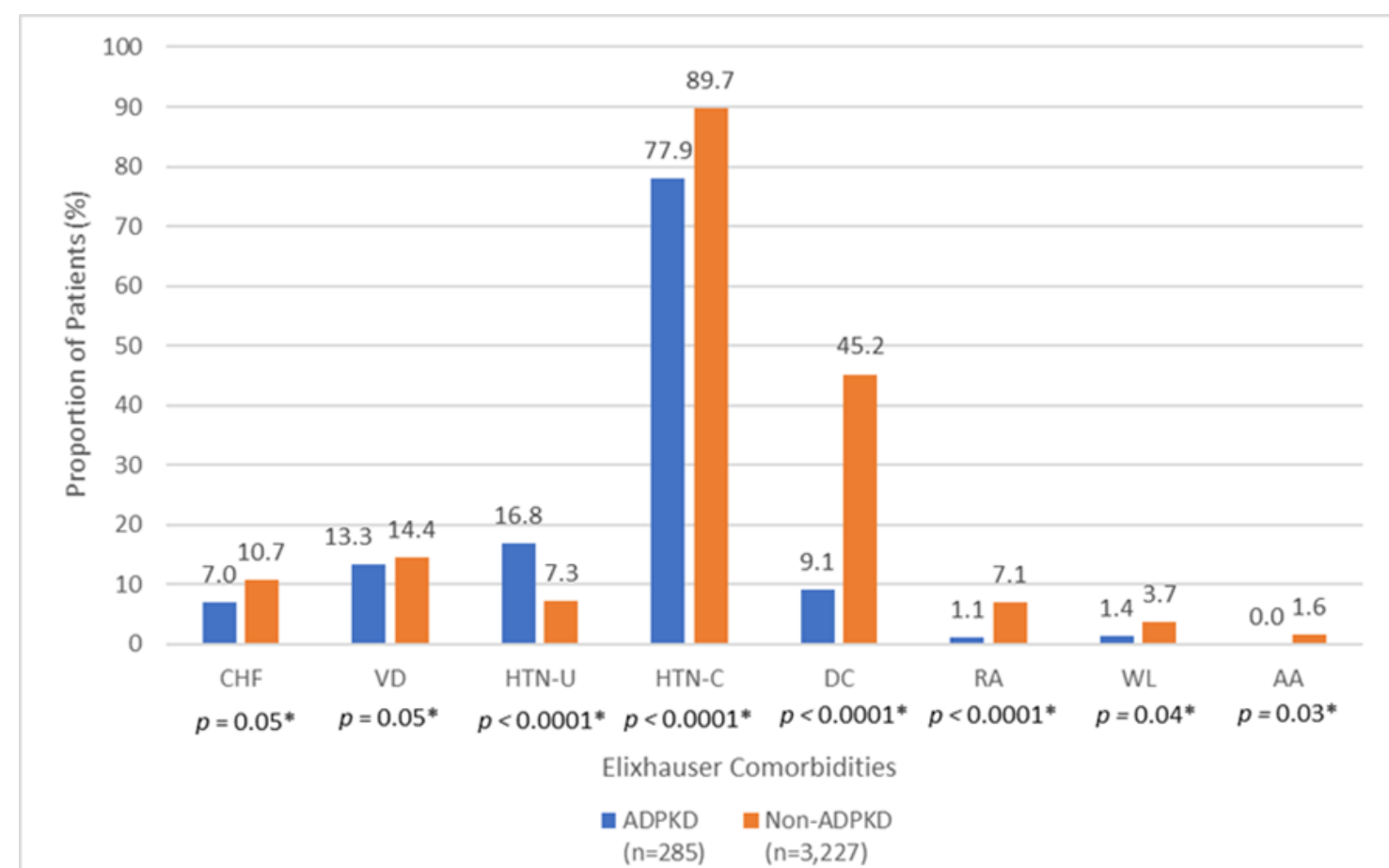
- A significantly higher proportion of recipients with ADPKD were White (71% vs. 51%). (Figure 3)
- A significantly higher proportion of recipients without ADPKD were of racial minority (Other [24 vs. 18%] and Black [23% vs. 11%] race). (Figure 3)

Figure 3. Proportion of KTP Recipients with vs. without ADPKD by Race



- KTP recipients with ADPKD had a lower comorbidity burden and better health status even though a higher proportion had uncomplicated hypertension (17% vs. 7%;  $p < 0.0001$ ). (Figure 4)

Figure 4. Comorbidities Among KTP Recipients with vs. without ADPKD at Index Discharge for KTP Surgery



- The median (IQR) LOS (4 [4-6] vs. 5 [4-7] days;  $p = 0.0006$ ) and total patient cost (\$103,000 [\$72,000-\$128,000] vs. \$113,000 [\$75,000-\$139,000];  $p = 0.0010$ ) were significantly lower among recipients with ADPKD. (Table 1)

Table 1. Inpatient Resource Use Outcomes Among Recipients with vs. without ADPKD at Index Discharge for KTP Surgery

| Data Element  | Estimates                                 |   |   |         |
|---|---|---|---|---------|
|   | Total Sample (N=3,512)                    | ADPKD (n=285)                             | Non-ADPKD (n=3,227)                       | p value |
| LOS, cumulative number of days  |   |   |   |         |
| Median (IQR)  | 5 (4 - 7)                                 | 4 (4-6)                                   | 5 (4-7)                                   | 0.0006* |
| Total Costs, \$USD  |   |   |   |         |
| Median (IQR)  | \$112,123.83 (\$74,865.34 – \$137,314.36) | \$102,940.74 (\$72,312.10 – \$127,647.76) | \$112,940.57 (\$75,308.88 – \$138,577.07) | 0.0010* |
| Abbreviations: ADPKD-autosomal dominant polycystic kidney disease, IQR-interquartile range, KTP-kidney transplantation, LOS-length of stay, PHD-Premier Healthcare Database, USD-United States dollars. |   |   |   |         |
| Significance: $p \leq 0.05$ *, Wilcoxon Rank Sum tests were conducted to compare the medians for continuous variables (LOS and total costs).  |   |   |   |         |



## Conclusions

- A significantly higher proportion of female recipients with ADPKD at index hospitalization for KTP surgery is likely an indicator of improved access to KTP among females with ADPKD.
- A significantly larger proportion of KTP recipients without ADPKD were of a racial minority compared to those with ADPKD at index hospitalization for KTP surgery. This may be an indicator of racial disparities in access to KTP among patients with ADPKD.
- KTP recipients with ADPKD impose less inpatient resource use and cost burden on hospitals compared to those without ADPKD. This is likely due to a lower comorbidity burden among recipients with ADPKD, resulting in a shorter post-surgical observation time and utilization of inpatient resources.

## References

- Alam, A., & Perrone, R. D. (2010). Management of ESRD in Patients with Autosomal Dominant Polycystic Kidney Disease. *Advances in Chronic Kidney Disease*, 17(2), 164-172.
- Amro, O. W., & Perrone, R. D. (2015). Patients with Autosomal Dominant Polycystic Kidney Disease. *Seminars in Dialysis*, 28(5), 470-473. doi:10.1111/sdi.12397.
- K. Rangan, G., Raghubanshi, A., Chaitarvornkit, A., Chandra, A. N., Gardos, R., Munt, A., . . . Wong, A. T. Y. (2020). Current and emerging treatment options to prevent renal failure due to autosomal dominant polycystic kidney disease. *Expert Opinion on Orphan Drugs*, 8(8), 285-302. doi:10.1080/21678707.2020.1804859
- Spithoven, E. M., Kramer, A., Meijer, E., Orskov, B., Wanner, C., Abad, J. M., . . . Wgikd. (2014). Renal replacement therapy for autosomal dominant polycystic kidney disease (ADPKD) in Europe: prevalence and survival--an analysis of data from the ERA-EDTA Registry. *Nephrol Dial Transplant*, 29 Suppl 4, iv15-25. doi:10.1093/ndt/gfu017