

Evidence Series: Study

**Time on Therapy of Automated
Peritoneal Dialysis with and
without Remote Patient
Monitoring: A Cohort Study**

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BACKGROUND

- Remote patient monitoring (**RPM**) of patients undergoing automated peritoneal dialysis (**APD-RPM**) may potentially enhance time on therapy due to possible improvements in technique and patient survival

OBJECTIVES

- To evaluate the effect of **RPM** (the **Sharesource** connectivity platform) in **APD** in terms of time on PD therapy compared to **APD without RPM**

ENDPOINTS



Primary outcome was time on APD therapy

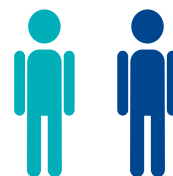


Secondary outcome was the mortality rate over a two-year period follow-up

METHODS

Retrospective, observational, multicenter, cohort study of incident patients undergoing **APD** (defined as those receiving **APD** for more than **90 days**) at the Baxter Renal Care Services network in Colombia

The patients were divided into two cohorts according to the use of RPM, which constitutes the exposure variable:



APD-RPM cohort: patients using the **Homechoice Claria** APD system device with the **Sharesource** connectivity platform (Baxter Healthcare, Deerfield, USA)



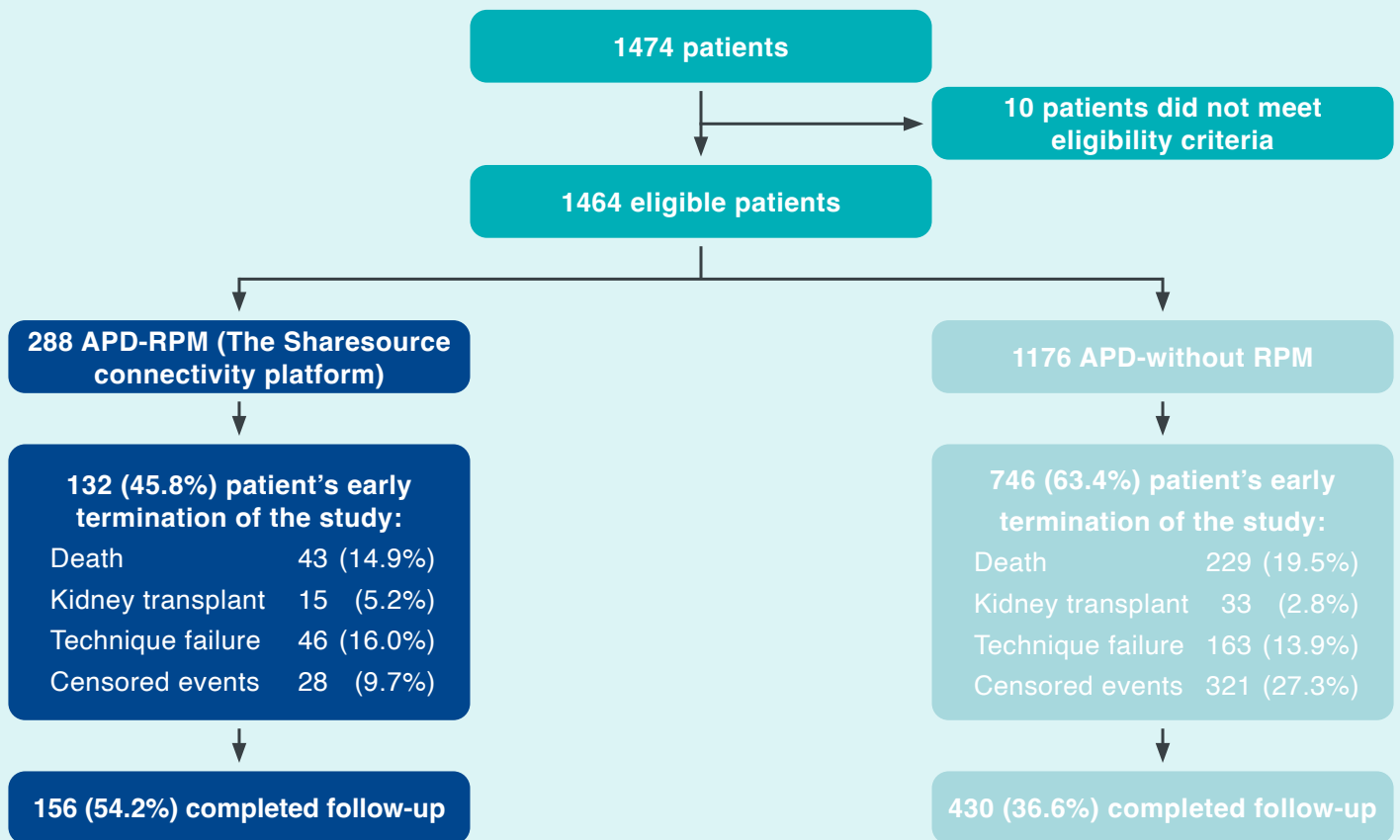
APD-without RPM cohort: patients using APD **Homechoice** without RPM

Follow-up within the study was up to two years

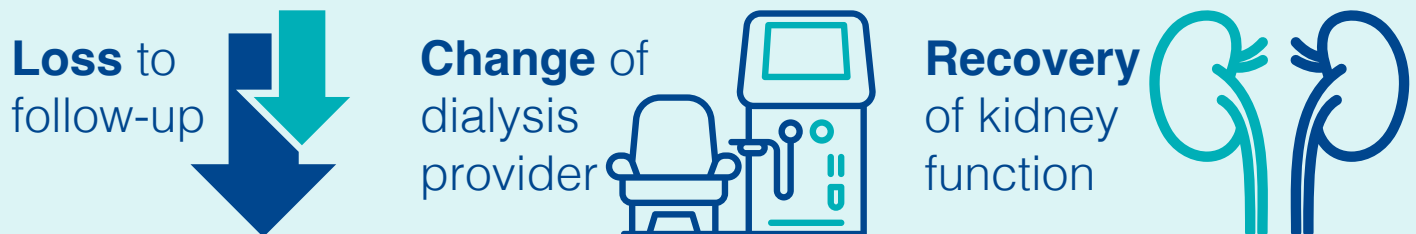


An intention-to-treat approach was used according to whether the patient used a remote monitoring program at the time of the inception of the cohort (**APD-RPM**) or not (**APD-without RPM**)

Reduction of Drop Out



Censored events were:



Change of **APD modality** (patients who started in the **APD without RPM** cohort and then switched to **APD with RPM**)

Propensity Score Matching (PSM):

- A propensity score matching (PSM) 1:1 without replacement utilising the nearest neighbor within caliper (0.035) was used and created a subpopulation in which the baseline covariates were well balanced
- The propensity score for each subject was calculated from a logistic regression model that included variables as predictors of the exposure status such as age, sex, black race, diabetes history, socioeconomic level, school level, Charlson Comorbidity Index, centre size, haemoglobin, phosphorus, potassium, albumin, urine output in mL/day, and censored events
- Fine & Gray multivariate analysis was performed to assess the effect of demographic, clinical, and laboratory variables on the risk of death, adjusting for competing risk events such as technique failure and kidney transplantation
- In addition, the cumulative incidence of deaths was estimated by adjusting for competing risk events, and Pepe and Mori's statistical test was used to compare the equality of the cumulative incidence functions (CIFs) by exposure status
- Inverse Probability of Treatment Weighting (IPTW) was approached using a propensity score to control differences between the groups and perform a sensitivity analysis for the direction of the observed effect

RESULTS

Full Sample:

- There were **272** death events with an overall rate of **0.13** events per patient-year
- The mortality incidence function adjusting for competing risks was:



APD-RPM group

7.1% for **1st year**
and **15.9%** in **2nd year**



APD-without RPM group

11% for **1st year**
and **23.6%** in **2nd year**

Matched sample:

- Statistically significant differences were observed in time on therapy when comparing APD-RPM with the APD-without RPM group. The time on therapy was greater for those with the **RPM program by 3.2 months**
- The APD-RPM group was associated with a lower mortality rate per patient-year, although this difference was not statistically significant. 0.10 events per patient-year in the APD-RPM group (95% CI: 0.07 to 0.13) vs. 0.12 in the APD-without RPM group (95% CI: 0.09 to 0.16), $p=0.325$.

Time on therapy and mortality rate in the full sample and matched sample:

	Full Sample			Matched Sample		
Outcomes	APD with RPM n=288	APD without RPM n=1176	p value	APD with RPM n=287	APD without RPM n=287	p value
Time on therapy, Months, mean (SD)	18.96 (7.32)	16.59 (8.04)	<0.001	18.95 (7.33)	15.75 (8.1)	<0.001
Difference, months (95% CI)	2.37 [1.35, 3.39]			3.2 [1.93, 4.46]		
Mortality, events/ person-year 95% CI	0.10 [0.07, 0.13]	0.14 [0.12, 0.16]	0.0013	0.10 [0.07, 0.13]	0.12 [0.09, 0.16]	0.325
IRR, 95% CI	0.67 [0.47, 0.93]			0.81 [0.54, 1.22]		

SD: Standard deviation, RPM: Remote patient monitoring, CI: Confidence interval, PS: Propensity score
IRR: Incidence rate ratio defined as APD-RPM/APD without

CONCLUSIONS

- Over two years of follow-up, APD patients supported by RPM with the **Sharesource** connectivity platform stayed 3.2 months longer on APD therapy compared to a matched group of APD patients without RPM
- This result indicates that RPM has the potential to improve the clinical effectiveness and the overall quality of APD therapy

Baxter's **Homechoice Claria** APD system is intended for automatic control of dialysis solution exchanges in the treatment of paediatric and adult renal failure patients undergoing peritoneal dialysis in the HOME HEALTHCARE ENVIRONMENT including comparable use in professional healthcare facilities.

The **Sharesource** portal is intended for use by healthcare professionals to remotely communicate new or modified treatment parameters with compatible dialysis instruments and transfer completed treatment data to a central database to aid in the review, analysis, and evaluation of patients' historical treatment results. This system is not intended to be a substitute for good clinical management practices, nor does its operation create decisions or treatment pathways.

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