

Mortality Rates Do Not Differ Among Patients on Peritoneal Dialysis Prescribed Different Active Vitamin D Agents

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INTRODUCTION

There is little well-controlled research examining the impact of the different formulations of oral vitamin D on clinical outcomes in dialysis patients. There is clear evidence that some providers have strong preference and nearly exclusive use of one oral vitamin D over another. For this retrospective analysis, we used physician preference as an instrumental variable to reduce or eliminate confounding.

METHODOLOGY

- We examined two years (7/1/2008 to 6/30/2010) of oral medication records of peritoneal dialysis (PD) patients from a large US dialysis organization.
- Patients were identified whose physicians prescribed only one form of vitamin D (calcitriol, paricalcitol, or doxercalciferol) to ≥90% of their PD patients.
- We excluded incident patients (treated <90 days) and patients whose physicians treated <5 PD patients.

RESULTS

Table 1. Vitamin D Usage: All PD Patients Across the LDO

Vitamin D usage	Patient- months	All months with use of any vitamin D (%)	All months with use of a single vitamin D (%)
Calcitriol only	44945	24.73%	30.35%
Doxercalciferol only	54219	29.83%	36.61%
Paricalcitol only	48949	26.93%	33.05%
Multiple formulations in month	4386	2.41%	
No oral vitamin D	29257	16.10%	
All patient-months	181756	100.00%	

Table 2. Vitamin D Usage: PD Patients of Physicians with 1 Formulation and Mortality Risk*

Physician formulation	Physicians	Patients	Use (%)	Deaths	Overall mortality	Deaths per 100 patient-years	95% CI
Calcitriol	74	643	39.5%	65	10.1%	9.33	7.06, 11.60
Doxercalciferol	53	516	31.4%	70	13.6%	12.20	9.34, 15.06
Paricalcitol	61	548	29.1%	64	11.7%	12.27	9.27, 15.28
Total	188	1707		199	11.7%	11.11	

^{*} limited to physicians prescribing only 1 form of vitamin D to ≥90% of their PD patients



	Formulation			
Patient Characteristics	Calcitriol (n=643)	Doxercalciferol (n=516)	Paricalcitol (n=548)	AII (N=1707)
Gender				
Male	354 (55.1%)	294 (57.0%)	270 (49.3%)	918 (53.8%)
Female	289 (44.9%)	222 (43.0%)	278 (50.7%)	789 (46.2%)
Race/Ethnicity				
African-American	215 (33.4%)	180 (34.9%)	150 (27.4%)	545 (31.9%)
Caucasian	240 (37.3%)	207 (40.1%)	297 (54.2%)	744 (43.6%)
Hispanic	82 (12.8%)	82 (15.9%)	68 (12.4%)	232 (13.6%)
Asian/Pacific Islander	74 (11.5%)	33 (6.4%)	17 (3.1%)	124 (7.3%)
Native American	2 (0.3%)	1 (0.2%)	1 (0.2%)	4 (0.2%)
Other/Unknown	30 (4.7%)	13 (2.5%)	15 (2.7%)	58 (3.4%)
Mean age (yr)	53.03	54.96	55.14	54.30
Mean vintage (yr)	2.99	2.90	2.49	2.80
Mean BMI	27.14	27.87	28.72	27.88
Cause of ESRD				
Diabetes	200 (31.1%)	176 (34.1%)	198 (36.1%)	574 (33.6%)
Hypertensive KD	148 (23.0%)	182 (35.3%)	137 (25.0%)	467 (27.4%)
Polycystic KD	21 (3.3%)	25 (4.8%)	38 (6.9%)	84 (4.9%)
Other	274 (42.6%)	133 (25.8%)	175 (31.9%)	582 (34.1%)
Mean Charlson Index	4.15	4.29	4.32	4.25

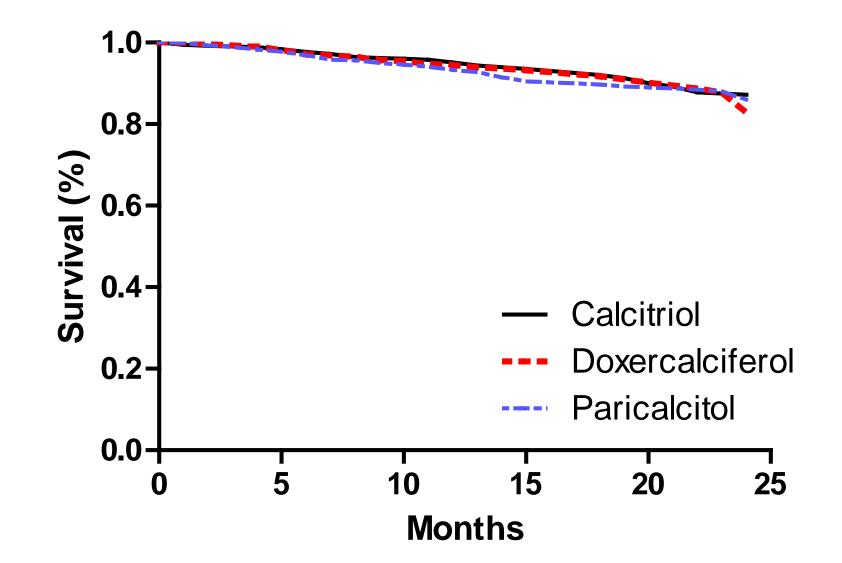


Figure 1. Survival Curve among the 3 Vitamin D Formulations. The survival curve has been adjusted for demographic factors, ERSD cause and comorbid conditions.

Table 4. Adjusted Hazards Ratios and 95% Risk Limit Intervals between Active Vitamin D Formulations

Comparison	Adjusted HR	95% Risk Limit Interval
Doxercalciferol vs. calcitriol	1.186	(0.795, 1.768)
Paricalcitol vs. calcitriol	1.213	(0.818, 1.797)
Paricalcitol vs. doxercalciferol	1.023	(0.696, 1.502)

SUMMARY of RESULTS

- Calcitriol is used in 30.35% of all patient-months with 1 vitamin D formulation (Table 1) but the patient-time observed among physicians identified to prescribe calcitriol is 39.5% (Table 2).
- Patients with physicians in the calcitriol group had a lower mortality rate (9.33 deaths per 100 patient-years) than patients in the other 2 groups (Table 2), however, results did not reach statistical significance.
- Patients in the calcitriol group had a mean age approximately 2 years lower than the other vitamin D groups (Table 3) and a lower prevalence of any cardiac or vascular disease: 9.3% calcitriol versus 12.6% and 15.0% for doxercalciferol and paricalcitol respectively.
- A Cox proportional hazards model, adjusting for differences in age, vintage, gender, race, BMI, and comorbidities showed no significant differences among the formulations (Figure 1 and Table 4).

KEY LEARNINGS

Among PD patients, instrumental variables analyses showed no differences in mortality among patients receiving any of the 3 currently available oral vitamin D formulations.

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