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Additional Acknowledgements

Supplemental Table 1. Summary of screening categorization for participants ineligible to

participate in the trial.

Consort Diagram Category	Reason	Number
1945 did not meet	• Patients without CKD stages 3b and 4	
eligibility criteria	(eGFR between 14 and	
	45ml/min/1.73m2)	527
	• Urinary albumin/creatinine ratio not	
	>10mg/mmol	97
	• Serum phosphate not >1.00mmol/L	704
	• Age less than 18 years	6
	• Unable to give informed consent	27
	• Patients with a history of psychological	
	illness or condition which interferes	
	with their ability to understand or	
	comply with the requirements of the	
	study	90
	Kidney transplantation	39
	• Recent (within 1 month) hospitalisation	
	or cardiovascular event	18
	 Pregnancy or breast feeding 	1
	 Medical conditions that impact on 	
	phosphate metabolism (apart from	
	CKD), eg. primary hyperparathyroidism	
	or hypoparathyroidism; previous	
	subtotal parathyroidectomy;	
	gastrointestinal malabsorption disorders	
	such as Crohn's disease, ulcerative	
	colitis, coeliac disease or severe liver	
	dysfunction	53
	• Malnutrition, defined as serum albumin	10
	<30g/L	10
	• Atrial fibrillation as documented on	4.5
	ECG performed at screening	45
	• Unable to obtain a pulse wave velocity	7
	• Unknown	321
814 other reasons	• Eligible but not approached for consent	426
	(various reasons)	
	• Insufficient information	388

Abbreviations: CKD, chronic kidney disease; ECG, electrocardiogram; eGFR, estimated glomerular

filtration rate

N=138) (32.6%) (44.2%)
(44.2%)
6 (33.3%)
(20.3%)
(55.8%)
1 (8.0%)
1 (73.2%)
2 (8.7%)
(19.6%)
(6.5%)

Supplemental Table 2. Medications at baseline for study participants, by treatment group.

Abbreviations: ACE, angiotensin-converting enzyme; ARB, angiotensin II receptor blocker

Results are presented as number (percentage)

Supplemental Table 3. Summary of missing data on primary and main secondary outcome variables

Outcome variable	No data	Baseline but no follow-up measures	No baseline but at least one follow-up measure§	Total participants with missing data
PWV (rated)	6	24	14	44
Agatston score	25	50	14	89
Serum phosphate	4	12	1	17
Serum calcium	4	13	2	19
Serum calcium- phosphate product	4	13	2	19
РТН	33	19	9	61
cFGF-23	27	27	19	73
iFGF-23	24	17	29	70
cFGF-23/ iFGF-23 ratio	29	28	29	86
Urinary phosphate excretion	15	27	20	62
eGFR	4	12	0	16
Creatinine clearance	14	31	26	71
Urine phosphate/creatinine ratio	21	26	52	99

§ As specified in the statistical analysis plan, mean imputation of missing baseline data permitted inclusion of these participants in the analysis.

<u>Supplemental Table 4.</u> On-treatment and compliance-adjusted comparisons of lanthanum and placebo groups at week 96 for intermediate cardiovascular parameters and biochemical parameters.

	On-treatment		Compliance-adjusted	
Outcome variable	Difference [95% CI] [#]	Р	Difference [95% CI] [#]	Р
		value		value
PWV (m/s)	+0.63 [-0.33, 1.59]	0.20	+0.59 [-0.32, 1.51]	0.20
Agatston score	+203 [-207, 612]	0.33	+176 [-195, 547]	0.35
Serum phosphate	-0.09[-0.13, -0.01]	0.03	-0.03 [-0.11, -0.05]	0.41
(mmol/L)				
cFGF-23 (RU/mL)*	-0.09 [-0.31, 0.12]	0.39	-0.03 [-0.24, 0.17]	0.74
iFGF-23 (pg/mL)*	-0.20 [-0.46, 0.06]	0.13	-0.08 [-0.33, 0.16]	0.51
PTH (pmol/L)	-0.10 [-0.29, 0.08]	0.27	0.01 (-0.16, 0.18]	0.91
Urinary phosphate	1.62 [-4.88, 1.63]	0.33	-1.17 [-4.21, 1.87]	0.45
excretion				
(mmol/24hrs)				

*Log transformed

[#]Lanthanum relative to placebo

Variables in the models include treatment group, visit, treatment group x visit interaction, baseline variable

Abbreviations: cFGF-23, c-terminal fibroblast growth factor 23; iFGF-23, intact fibroblast growth factor 23; PWV, pulse wave velocity

Additional Acknowledgements

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