Abstract

Background. Despite recognized risks associated with hyperphosphataemia in patients with chronic kidney disease (CKD) Stage 5 on dialysis, the achievement of target levels of serum phosphate is poor. It is likely that this is partly due to poor adherence by patients to their phosphate-binder treatment regimens, which often comprise large daily tablet burdens.

Methods. In this multicentre, open-label trial, patients on a stable dialysis regimen were screened while receiving phosphate-binder therapy, then entered into a washout phase. Patients with serum phosphate > 1.78 mmol/L after washout entered into the main 12-week treatment phase (N = 367), during which they were treated to target [Kidney Disease Outcomes Quality Initiative (K/DOQI): 1.13–1.78 mmol/L; 3.5–5.5 mg/dL] with lanthanum carbonate monotherapy. Efficacy variables included serum phosphate levels and the percentage of patients with serum phosphate control. Safety and tolerability assessments were also conducted.

Results. Mean serum phosphate levels were significantly reduced following 12 weeks of lanthanum carbonate monotherapy versus previous phosphate-binder therapy. The mean number of phosphate-binder tablets being taken per day at screening was 7.6, but during treatment with lanthanum carbonate, most patients were taking doses of up to 3000 mg/day, achievable with 3 × 1000 mg tablets per day (maximum of 6).

Conclusion. These findings suggest that lanthanum carbonate monotherapy offers effective control of serum phosphate and, due to a low tablet burden, may help to simplify the management of hyperphosphataemia in patients with CKD Stage 5.

Keywords: dialysis; hyperphosphataemia; lanthanum carbonate; phosphate binder; tablet burden

Introduction

Hyperphosphataemia is an established risk factor for cardiovascular mortality, renal osteodystrophy and secondary hyperparathyroidism in patients with chronic kidney disease (CKD) Stage 5 on dialysis, and the need for adequate control of serum phosphate is well recognized [1–3]. Despite this, the evidence suggests that target levels of serum phosphate are not consistently being met. Indeed, the Dialysis Outcomes and Practice Patterns Study (DOPPS) revealed that less than half of the patients (40.8%) achieved the Kidney Disease Outcomes Quality Initiative (K/DOQI) target range for serum phosphate (1.13–1.78 mmol/L; 3.5–5.5 mg/dL) [4]. Given the demonstrated efficacy of phosphate binders in clinical trials [5–9], it is unlikely that the failure to achieve targets is entirely due to a lack of availability of effective treatments. A number of other factors may contribute to the poor achievement of serum phosphate targets, including lack of patient understanding of the consequences of hyperphosphataemia, variable dietary management and poor adherence to phosphate-binder treatment regimens.

Poor adherence to medication is an increasingly recognized problem among patients with CKD Stage 5 [10]. These patients are subject to large daily tablet burdens and may be prescribed > 10 different medications [11]. Typically, the tablet burden can be expected to be two- to three-fold higher than the total number of prescribed medications per day [12]. Without doubt, the cumulative effect of multiple dosing regimens can impose a confusing and possibly overwhelming burden on a patient. In addition, patients with CKD Stage 5 are often elderly and the condition has been associated with an increased rate of cognitive decline compared with individuals with preserved renal function [13].

Phosphate-binder therapy contributes a considerable proportion of the daily tablet burden for patients with CKD Stage 5, and for some binders, nine or more tablets are required per day to treat elevated serum phosphate levels in certain patients [14,15]. Furthermore, combination phosphate-binder therapy is commonly adopted in an attempt to minimize costs, avoid excessive elevation of serum phosphate and, due to a low tablet burden, may help to simplify the consequences of hyperphosphataemia, variable dietary management and poor adherence to phosphate-binder treatment regimens.

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Switching to lanthanum carbonate monotherapy provides effective phosphate control with a low tablet burden

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calcium levels and achieve control of serum phosphate to target levels [1]. However, this practice may be counterproductive if it reduces the adherence of patients to their phosphate-binder regimens as a result of increased tablet burden or complexity of the treatment regimen. Lanthanum carbonate (FOSRENOL®, Shire Pharmaceuticals, Basingstoke, UK) is an effective non-calcium-based phosphate binder [16,17], for which efficacy, safety and tolerability profiles have been demonstrated in both short- and long-term clinical studies [18–22]. Importantly, it has proven to be effective with a relatively low daily tablet burden, with the majority of patients requiring a single tablet taken during each meal [20].

In this prospective, multicentre, open-label study, we switched patients from their previous phosphate-binder therapy (which frequently consisted of more than one agent) to lanthanum carbonate monotherapy, in order to investigate the effect on serum phosphate control and dose requirements. Although this study was designed to assess phosphate control using higher doses of lanthanum carbonate in a clinical practice setting, other data were captured, including the tablet burden attributable to phosphate binders.

**Subjects and methods**

*Inclusion/exclusion criteria*

The study included male and female CKD Stage 5 patients aged at least 18 years, who had received dialysis for two consecutive months prior to the study and had a current requirement for treatment of hyperphosphataemia. Females of childbearing potential were required to have a negative serum pregnancy test prior to entering the study. Patients were excluded from the study if they required continued treatment with aluminium-, calcium- or magnesium-containing compounds into the study treatment period. Exclusion criteria also included: significantly abnormal laboratory values, serum transaminases [alanine aminotransferase (ALT) and aspartate aminotransferase (AST)] elevated to more than three times the upper limit of normal, intact parathyroid hormone (PTH) >85 pmol/L (800 pg/mL) or clinically significant, uncontrolled concurrent illness.

*Study design*

This was a multicentre, open-label trial carried out in 49 centres across Europe, Israel and Canada, designed to assess whether daily doses of lanthanum carbonate > 3000 mg offer improved rates of serum phosphate control. Patient eligibility was assessed during a screening period of up to 1 week. A complete medical history was taken and demographic information recorded at this point, along with vital signs, pre-dialysis blood profile and concomitant medications.

The study design is summarized in Figure 1. Patients receiving prior phosphate binder(s) discontinued their treatment and entered a 1- to 2-week washout period. Serum phosphate levels were reviewed after each week. Patients whose serum phosphate levels rose above 1.78 mmol/L were eligible to begin lanthanum carbonate treatment and considered to be at baseline at this point. Those who did not demonstrate this level of serum phosphate after 2 weeks were excluded from the study. Patients who were not receiving any phosphate-binder treatment and had serum levels above 1.78 mmol/L entered the treatment period immediately following the screening period.

Patients began lanthanum carbonate treatment at a total daily recommended dose of 1500 mg/day (tablets containing 250, 500, 750 and 1000 mg elemental lanthanum were available). The daily dose was divided equally between meals, with tablets to be chewed and taken immediately after food, and could be increased by 750 mg/day each week to achieve optimal control of serum phosphate levels, to a maximum of 3000 mg/day by week 3 and 4500 mg/day by week 5 (daily dose could also be decreased). The target serum phosphate range was 1.13–1.78 mmol/L (3.5–5.5 mg/dL), with investigators requested to aim for the middle of this range. Once target levels were reached, patients received stable dosage for the remainder of the treatment period.

The primary efficacy variable was the percentage of patients with controlled serum phosphate. The primary comparison was the control rate at week 3, when patients could have been titrated up to 3000 mg/day, compared with the rate at week 5, when patients could have been titrated up to 4500 mg/day.

Patients who completed the main 12-week study were eligible to continue into an optional 24-week extension phase. Patients entering the extension phase maintained their dose at week 12 levels but dosage could be adjusted at any time to achieve optimal control of serum phosphate levels.

Pre-dialysis vital signs, post-dialysis weight, concomitant medications and adverse events (AEs) were assessed at baseline, weekly for treatment weeks 1–6 and at weeks 9, 12 (end of main study), 24 and 36 (end of extension phase). For haemodialysis patients, blood samples were collected pre-dialysis at the first dialysis session of the week. For peritoneal dialysis patients, study visits were scheduled for a Monday or Tuesday. Full blood profiles were conducted and pre-dialysis plasma lanthanum levels assessed at baseline and weeks 3, 5, 12 and 36 (partial blood profiles were conducted at all other visits).

From the full blood profiles conducted at screening, baseline and at weeks 3, 5, 12 and 36, primary (serum phosphate) and secondary [calcium, intact PTH and calcium × phosphate product (Ca × P)] efficacy variables were assessed. For all patients, serum-intact PTH was measured using a DPC Immulite iPTH kit. Routine physical examinations and laboratory measurements were taken. Liver function tests were carried out, measuring ALT, AST, gamma glutamyl transferase (GGT), alkaline phosphatase and bilirubin levels.

Adverse events were recorded from the time that informed consent was signed until the end of treatment exposure and for the 30 days following the last exposure to the study medication. Treatment-emergent adverse events (TEAEs) were defined as those that occurred during this period and were not present prior to the first dose of study medication, or were present prior to the first dose of study medication but the severity increased during treatment.
### Patient populations

**Safety population.** Safety population included all patients who received at least one dose of study medication. Screening and baseline characteristics were reported for this population, as well as assessments of tolerability and safety.

**Intention-to-treat (ITT) population** The primary efficacy population included all patients who received at least one dose of lanthanum carbonate and had at least one post-baseline phosphate measurement. Primary and secondary efficacy data were reported for this population.

### Statistics

It was anticipated that at week 3 of lanthanum carbonate treatment (maximum dose 3000 mg/day), ~50% of patients would be controlled, and that at higher doses (maximum dose 4500 mg, week 5), a clinically important effect would result in 55–60% of patients being controlled. It was estimated that 388 patients would be needed at week 3 and week 5 to provide 90% power to detect a minimum clinically important difference of 5% in control rates between doses, with 5% statistical significance determined using a two-sided test. Continuous variables were summarized using the number of observations, number of missing observations, mean, standard deviation (SD), 95% confidence interval (95% CI), median, minimum and maximum. The standard error (SE) was presented where appropriate. Categorical data were summarized using the number of observations, number of missing observations and percentages. Changes from screening were analysed using the one-sample t-test. McNemar’s test of paired proportions was used to determine differences in control rates between weeks 3 and 5.

### Results

#### Patient disposition

In total, 477 patients were screened, 367 of whom entered the main study. Of these, 366 received treatment, 274 completed the 12-week period, while 93 withdrew. Reasons for discontinuation in the main study are presented in Table 1.

Two hundred and eleven patients entered the 24-week extension phase. Of these, 210 received treatment, 159 completed the study, while 52 withdrew. The most common reasons for discontinuation were AEs/SAEs (17 patients), patient request (13 patients) and kidney transplant (9 patients). TEAEs resulting in withdrawal from the study were reported for 9.5% of patients during the extension phase.

#### Patient demographics and other baseline characteristics

Baseline characteristics are shown in Table 2. The majority of patients entering the main study were male, Caucasian.
Table 2. Baseline characteristics for all patients entering the main study and extension phase

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>Main study n = 366</th>
<th>Extension phase n = 211</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sex, n (%)</td>
<td>Female 115 (31.4)</td>
<td>57 (27.0)</td>
</tr>
<tr>
<td></td>
<td>Male 251 (68.6)</td>
<td>154 (73.0)</td>
</tr>
<tr>
<td>Ethnic origin, n (%)</td>
<td>Caucasian 307 (83.9)</td>
<td>183 (86.7)</td>
</tr>
<tr>
<td></td>
<td>Black 4 (1.1)</td>
<td>4 (1.9)</td>
</tr>
<tr>
<td></td>
<td>Hispanic 5 (1.4)</td>
<td>4 (1.9)</td>
</tr>
<tr>
<td></td>
<td>Asian 12 (3.3)</td>
<td>4 (1.9)</td>
</tr>
<tr>
<td></td>
<td>Other 2 (0.5)</td>
<td>1 (0.5)</td>
</tr>
<tr>
<td></td>
<td>Missing* 36 (9.8)</td>
<td>15 (7.1)</td>
</tr>
<tr>
<td>Age and height, mean (SD) [range]</td>
<td>Age (years) 59.5 (13.81) [24–85]</td>
<td>57.9 (14.53) [27–85]</td>
</tr>
<tr>
<td></td>
<td>Weight (kg) 73.1 (15.47) [33–128]</td>
<td>74.1 (11.53) [33–128]</td>
</tr>
</tbody>
</table>

*Local legislation prevented study centres in France from recording ethnic or racial information.

and had been on dialysis for 2 years or more [median (range) time since starting dialysis, 2.42 (0.2–33.3) years]. Approximately one-sixth had received renal transplants (16.1%) and the most frequent known causes underlying CKD Stage 5 were diabetes (23.2%) and glomerulonephritis (21.6%).

Medication use at screening

The number of medications that patients were receiving at the screening visit ranged from 2 to 27 per patient (mean, 10.2).

At screening, a large proportion of the ITT population (40.7%) were receiving more than one phosphate-binding agent. Another 57.4% were receiving monotherapy, and 1.9% were not receiving any phosphate binder (Figure 2). Calcium carbonate was the most commonly used binder (60.2%), followed by sevelamer hydrochloride (55.7%), calcium acetate (11.4%) and aluminium-based binders (10.9%). These figures demonstrate that combination phosphate-binder use was prevalent in this patient population. The mean total daily phosphate-binder tablet count for the ITT population in the main study, whether on monotherapy or combination therapy, was 7.6 tablets (range, 1–28). For patients who required one, two or three agents the mean (range) daily tablet count was 5.3 (1–15), 10.5 (2–28) and 13.1 (9–21) tablets, respectively.

Phosphate levels and control rates

At screening, 34.9% of patients met the K/DOQI serum phosphate target range, falling to 7.0% after washout. After 3 weeks of lanthanum carbonate treatment, 39.1% of patients met the K/DOQI targets, with the control rate 43.6% at week 5. The percentage of patients with controlled serum phosphate at week 5 was observed to be higher than that at week 3, but it was not statistically significant and was less than the predefined level of clinical importance. Forty-eight percent of patients were controlled to K/DOQI by week 12, and this was statistically significant (P = 0.026) when compared with screening levels. Of those patients whose serum phosphate was not controlled on previous phosphate binder(s), 26.4% achieved control on lanthanum carbonate.

At screening, when patients were still using their previous phosphate binder, the mean (95% CI) serum phosphate level was 1.99 (1.92, 2.06) mmol/L. During the 1- to 2-week washout period, serum phosphate levels increased as expected, and then decreased with lanthanum carbonate treatment. At week 12, the mean (95% CI) serum phosphate level was 1.84 (1.78, 1.90) mmol/L. The mean reduction from screening in serum phosphate levels after 12 weeks (for patients with both screening and week 12 measurements) on lanthanum carbonate treatment was significant [mean (SE) change from screening, −0.13 (0.05) mmol/L; P = 0.007 using the one-sample t-test based on the 98.1% patients receiving phosphate binders at screening]. Furthermore, mean serum phosphate levels improved with 12 weeks of lanthanum carbonate monotherapy regardless of whether patients were previously receiving monotherapy (1.83 versus 1.98 mmol/L) or combination binder therapy (1.87 versus 1.97 mmol/L) (Figure 3).
The distribution of dose levels at week 12 and percentages of patients controlled to K/DOQI targets at each dose are shown in Figure 4. Although the tablet burden was not recorded in this study, most patients (57.1%) were receiving doses that could be achievable with 3 tablets per day (≤3000 mg/day), with the remainder on doses achievable with 6 tablets per day (median dose, 3 tablets per day).

Of the 48% of patients whose serum phosphate levels fell within the target range by week 12, more than three quarters (77.1%) were taking daily lanthanum carbonate doses of ≤3000 mg.

**Other mineral metabolism parameters**

At screening, the mean (95% CI) serum total calcium level in the main study group was 2.35 (2.33, 2.37) mmol/L. Levels of calcium fell following washout to 2.27 (2.25, 2.29) mmol/L at baseline, but mean serum calcium rose back to a level similar to those observed at screening with lanthanum carbonate treatment [2.34 (2.32, 2.37) mmol/L at week 12]. Similar observations were made in patients taking part in the extension phase. Calcium levels corrected for albumin were almost identical to total calcium levels.

The mean (95% CI) level of Ca × P at screening was 4.65 (4.51, 4.80) mmol²/L². Levels increased after washout to 5.77 (5.62, 5.93) mmol²/L² at baseline, but then fell following commencement of lanthanum carbonate therapy to below those observed at screening [4.31 (4.16, 4.47) mmol²/L² at week 12]. A similar change was seen in patients taking part in the extension phase.

At screening, the mean (95% CI) intact PTH level was 27.76 (25.29, 30.23) pmol/L but increased following the washout period to 38.63 (35.67, 41.59) pmol/L at baseline. Once lanthanum carbonate treatment began, mean PTH levels decreased [33.92 (31.14, 36.70) pmol/L at week 3] and remained relatively stable throughout treatment.

**Safety and tolerability**

Lanthanum carbonate treatment was generally well tolerated and caused no unexpected safety concerns. The TEAE profile was generally typical of a population with CKD Stage 5 receiving haemodialysis (Table 3). Although increasing the dose from 3000 to 4500 mg/day did not result in a statistically significant increase in the percentage of patients controlled, it had no effect on the number of TEAEs that occurred. Treatment emergent adverse events were reported for 145 (72.5%) and 114 (68.7%) patients receiving doses up to 3000 mg/day and >3000 mg/day, respectively.

In the main study and extension phase, 14.2% and 7.6% of patients, respectively, reported SAEs. These were primarily infections, cardiac disorders and gastrointestinal disorders. Eight deaths occurred during the study, seven of which were considered to be unrelated to treatment. One was documented as possibly related to treatment, the most likely cause being reported as gastrointestinal bleeding caused by non-infectious enteritis, complicated by excessive anticoagulation and possible myocardial infarction.

**Table 3. Treatment-emergent adverse events (TEAEs) with an incidence of ≥5% in the main study (0–12 weeks) and extension phase (12–36 weeks), according to MedDRA preferred term**

<table>
<thead>
<tr>
<th>MedDRA preferred term</th>
<th>Main study, N = 366</th>
<th>Extension phase, N = 211</th>
</tr>
</thead>
<tbody>
<tr>
<td>Vomiting</td>
<td>48 (13.1)</td>
<td>15 (7.1)</td>
</tr>
<tr>
<td>Nausea</td>
<td>45 (12.3)</td>
<td>9 (4.3)*</td>
</tr>
<tr>
<td>Diarrhoea</td>
<td>35 (9.6)</td>
<td>17 (8.1)</td>
</tr>
<tr>
<td>Abdominal pain upper</td>
<td>19 (5.2)</td>
<td>2 (0.9)*</td>
</tr>
<tr>
<td>Nasopharyngitis</td>
<td>24 (6.6)</td>
<td>23 (10.9)</td>
</tr>
<tr>
<td>Total patients with ≥1 TEAE</td>
<td>259 (70.8)</td>
<td>152 (72.0)</td>
</tr>
</tbody>
</table>

*Value included for completeness due to ≥5% incidence in other section of the study.
Treatment with lanthanum carbonate caused no clinically noteworthy important changes (clinical importance was determined by site investigators) in vital signs, haematological parameters or markers of liver function during the main study. Background plasma lanthanum levels were detectable at baseline and increased slightly following lanthanum carbonate treatment to 0.3–0.7 ng/mL during the main study and to 0.5–0.8 ng/mL during the extension phase. There was no evidence of dose dependence in these increases, and the levels are typical of those observed in other studies of lanthanum carbonate treatment [20,22].

**Discussion**

The data presented here demonstrate that patients can be treated with lanthanum carbonate monotherapy following either combination or monotherapy with other phosphate binders, and levels of serum phosphate are generally maintained or improved. There was no significant difference between the percentage of patients who achieved serum phosphate control at week 3 compared with week 5, when the maximum possible doses of lanthanum carbonate were 3000 mg/day and 4500 mg/day, respectively. The additional percentage of patients controlled on doses of 3750 and 4500 mg compared with 3000 mg was smaller than the pre-determined threshold for clinical importance. Hence, only a small proportion of patients benefited from increasing lanthanum carbonate doses above 3000 mg, supporting the current clinical practice of mainly utilizing doses in the region of 2250–3000 mg/day.

According to the protocol, titration of the dose of lanthanum carbonate up to 4500 mg/day was permitted in order to achieve control of serum phosphate to K/DOQI targets. Surprisingly, some patients did not achieve the K/DOQI target range for serum phosphate and yet were not titrated up to the maximum dose of 4500 mg/day, as permitted by and expected according to the protocol. With regard to dose titration, we consider that the behaviour of the investigators is likely to be related to their expectations for each particular patient. If one patient has achieved serum phosphate levels that the investigator considers to be a significant improvement, despite not achieving the specified target range, they may elect not to titrate further. It should also be considered that binding dietary phosphate alone cannot eradicate hyperphosphataemia. Phosphate may be released from the bone under the influence of PTH [24], and the patient must adhere to their dietary restrictions and dialysis regimen if phosphate control is to be optimal.

The medical history and disease characteristics of the patients enrolled in this study were typical of a dialysis population, but the data relating to concomitant medications highlight the considerable tablet burden that this disease generates. Patients were taking up to 27 different medications on entry to the study, many of which require administration more than once a day. Indeed, the tablet burden associated with phosphate-binder therapy alone was shown to be extensive, with between 1 and 28 tablets per day recorded at screening in this study.

Despite demonstrated use of phosphate binders in this population at screening, with a large proportion of patients receiving combination therapy, approximately two-thirds of patients (65.1%) had serum phosphate levels that were not controlled to K/DOQI targets. Given the efficacy that has been demonstrated in clinical trials [5,9], it can be speculated that this poor level of target achievement in clinical practice may be at least partly due to suboptimal levels of patient adherence to their phosphate-binder therapy. Failure of patients to adhere to their prescribed treatment regimens is believed to be related to both perceptual and practical barriers [23]. A poor understanding of what hyperphosphataemia is, and of the potential dangers of failing to control it adequately, coupled with a high daily tablet burden, would be expected to impact on the extent to which patients take their phosphate binders as prescribed. Minimizing the tablet burden associated with phosphate-binder therapy may help to improve the level of patient adherence to that treatment, as well as perhaps to other medications, by reducing the overall complexity of the daily dosing regimen.

The majority (77.1%) of patients who achieved serum phosphate control on lanthanum carbonate treatment did so on a dose of 3000 mg/day or less, which is equivalent to one tablet taken during each meal (3 per day). Given that the mean daily tablet burden attributable to phosphate-binder therapy prior to switching to lanthanum carbonate was 7.6 tablets, switching to lanthanum carbonate monotherapy clearly offered a reduction in average daily tablet burden in this study.

There is a tendency in practice to combine phosphate binders in an attempt to improve efficacy or minimize side effects, but this may be counterproductive, as combination phosphate-binder use can add to treatment complexity, which may in turn make it more difficult for patients to adhere. The reduction in tablet burden that lanthanum carbonate offers may therefore help to improve patient adherence, although a study of this design does not permit us to investigate this. Nevertheless, the data from this study support the practice of stopping other phosphate-binder therapy altogether and switching to lanthanum carbonate monotherapy, as tablet burden is reduced with no loss of efficacy or undesirable changes in serum calcium levels. Mean intact PTH levels were generally slightly above the upper limit of the K/DOQI-recommended range (16.5–33 pmol/L; 150–300 pg/mL) during treatment with lanthanum carbonate.

Treatment with lanthanum carbonate was generally well tolerated and did not generate any unexpected safety concerns compared with the screening population. The high affinity of lanthanum for phosphate ions and the low level of absorption from the gastrointestinal tract contribute to a minimal drug–drug interaction profile [17,25,26].

A limitation of this study was the lack of control group. However, a previous 6-month, randomized trial, comparing the efficacy of lanthanum carbonate with calcium carbonate, has demonstrated that lanthanum carbonate is an effective phosphate binder compared with standard therapy [19].

In conclusion, lanthanum carbonate as a monotherapy generated a statistically significant reduction in serum phosphate levels compared with previous phosphate-binder mono- or combination therapy in this study. In more than three quarters of patients who achieved serum phosphate...
control to K/DOQI targets during the main study, the dose at week 12 was not >5000 mg/day, which equates to just one tablet taken during each meal. These data demonstrate that switching to lanthanum carbonate monotherapy offers maintained or improved control of serum phosphorus with a reduced tablet burden. This may have positive implications for the adherence of patients with CKD Stage 5 to their phosphate-binder therapy.

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References


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