LOKELMA is a potassium binder indicated for the treatment of hyperkalemia in adults. (1)

Limitation of Use
LOKELMA should not be used as an emergency treatment for life-threatening hyperkalemia because of its delayed onset of action. (1)

Dosage and Administration

For initial treatment of hyperkalemia, the recommended dose of LOKELMA is 10 g administered three times a day for up to 48 hours. (2.1)

For maintenance treatment, recommended dose is 10 g once daily. (2.1)

Adjust dose at one-week intervals as needed (by 5 g daily) to obtain desired serum potassium target range. (2.1)

For oral suspension: 5 g per packet (3)
For oral suspension: 10 g per packet (3)

Adverse Reactions

Most common adverse reactions with LOKELMA: mild to moderate edema. (6.1)

In general, other oral medications should be administered at least 2 hours before or 2 hours after LOKELMA. (2.2, 7, 12.3)

Adverse Reactions

• Gastrointestinal Adverse Events in Patients with Motility Disorders. (5.1)
• Edema. (5.2)

Drug Interactions

None. (4)

Warnings and Precautions

• Gastrointestinal Adverse Events in Patients with Motility Disorders. (5.1)

To report SUSPECTED ADVERSE REACTIONS, contact AstraZeneca at 1-800-236-9933 or FDA at 1-800-FDA-1088 or www.fda.gov/medwatch.

Dosing Forms and Strengths

• For oral suspension: 5 g per packet (3)
• For oral suspension: 10 g per packet (3)
5.2 Edema
Each 5 g dose of LOKELMA contains approximately 400 mg of sodium. In clinical trials of LOKELMA, edema was generally mild to moderate in severity and was more commonly seen in patients treated with 15 g once daily. Monitor for signs of edema, particularly in patients who should restrict their sodium intake or are prone to fluid overload (e.g., heart failure or renal disease). Advise patients to adjust dietary sodium, if appropriate. Increase the dose of diuretics as needed [see Adverse Reactions (6)].

6 ADVERSE REACTIONS
The following adverse reactions are discussed in greater detail elsewhere in the label: 
- Edema [see Warnings and Precautions (5.2)].

6.1 Clinical Studies Experience
Because clinical studies are conducted under widely varying conditions, adverse reaction rates observed in the clinical studies of a drug cannot be directly compared to rates in the clinical studies of another drug and may not reflect the rates observed in practice.

The total exposure to LOKELMA in the safety and efficacy clinical trials of patients with hyperkalemia was 1,760 patients with 652 patients exposed to LOKELMA for at least 6 months and 507 patients exposed for at least one year.

The population (n=1,009) in the placebo-controlled trials included patients aged 22 to 96 years, females (n=454), Caucasians (n=859) and Blacks (n=130). Patients had hyperkalemia in association with comorbid diseases such as chronic kidney disease, heart failure and diabetes mellitus.

In placebo-controlled trials in which patients were treated with once daily doses of LOKELMA for up to 28 days, edema was reported in 4.4% of patients receiving a dose of 5 g and 5.9% of patients receiving 10 g of LOKELMA compared to 2.4% of patients receiving placebo. In longer-term uncontrolled trials in which most patients were maintained on doses <15 g once daily, adverse reactions of edema (generalized edema and peripheral edema) were reported in 8% to 11% of patients.

Laboratory Abnormalities
In clinical trials, 4.1% of LOKELMA-treated patients developed hypokalemia with a serum potassium value less than 3.5 mEq/L, which resolved with dosage reduction or discontinuation of LOKELMA.

7 DRUG INTERACTIONS
LOKELMA can transiently increase gastric pH. As a result, LOKELMA can change the absorption of co-administered drugs that exhibit pH-dependent solubility, potentially leading to altered efficacy or safety of these drugs when taken close to the time LOKELMA is administered. In general, other oral medications should be administered at least 2 hours before or 2 hours after LOKELMA [see Dosage and Administration (2.2) and Clinical Pharmacology (12.3)]. LOKELMA is not expected to impact systemic exposure of drugs that do not exhibit pH-dependent solubility and so spacing is not needed if it has been determined that the concomitant medication does not exhibit pH-dependent solubility.

8 USE IN SPECIFIC POPULATIONS

8.1 Pregnancy
Risk Summary
LOKELMA is not absorbed systemically following oral administration and maternal use is not expected to result in fetal exposure to the drug.

8.2 Lactation
Risk Summary
LOKELMA is not absorbed systemically following oral administration, and breastfeeding is not expected to result in exposure of the child to LOKELMA.

8.4 Pediatric Use
Safety and effectiveness in pediatric patients have not been established.

8.5 Geriatric Use
Of the total number of subjects in clinical studies of LOKELMA, 58% were age 65 and over, while 25% were 75 and over. No overall differences in safety or effectiveness were observed between these patients and younger patients.

11 DESCRIPTION
LOKELMA is a powder for oral suspension. The active ingredient in LOKELMA is sodium zirconium cyclosilicate, a potassium binder. Sodium zirconium cyclosilicate is a non-absorbed zirconium silicate that preferentially exchanges potassium for hydrogen and sodium. LOKELMA is a free flowing, odorless, insoluble white powder for oral suspension. It has a mean particle size of 20 µm and includes no more than 3% of particles with a diameter below 3 µm. Each 5 g of sodium zirconium cyclosilicate contains 400 mg of sodium.

12 CLINICAL PHARMACOLOGY

12.1 Mechanism of Action
LOKELMA (sodium zirconium cyclosilicate) is a non-absorbed zirconium silicate that preferentially captures potassium in exchange for hydrogen and sodium, in vitro, LOKELMA has a high affinity for potassium ions, even in the presence of other cations such as calcium and magnesium. LOKELMA increases fecal potassium excretion through binding of potassium in the lumen of the gastrointestinal tract. Binding of potassium reduces the concentration of free potassium in the gastrointestinal lumen, thereby lowering serum potassium levels.

12.2 Pharmacodynamics
In a study in healthy adult subjects, LOKELMA administered as 5 g or 10 g once daily for four days caused a dose-dependent increase in fecal potassium excretion. Corresponding dose-dependent decreases in urinary potassium excretion and serum potassium were also observed. In patients with hyperkalemia treated with LOKELMA 10 g three times a day for up to 48 hours, reductions in serum potassium were observed one hour after initiation of therapy; serum potassium concentrations continued to decline over the 48-hour treatment period [see Clinical Studies (14.2)]. In patients not continuing LOKELMA, potassium levels increased. Patients with higher starting serum potassium levels or receiving a higher dose have greater reductions in serum potassium.

LOKELMA causes a small dose-dependent increase in serum bicarbonate concentrations (1.1 mmol/L at 5 g once daily, 2.3 mmol/L at 10 g once daily and 2.6 mmol/L at 15 g once daily as compared with a mean increase of 0.6 mmol/L in patients treated with placebo). The clinical significance of this finding is unclear.

12.3 Pharmacokinetics
LOKELMA is an inorganic, insoluble compound that is not subject to enzymatic metabolism. In a clinical study in patients with hyperkalemia in which zirconium concentrations were measured in the urine and blood, zirconium concentrations were similar in treated and untreated patients (i.e., either undetectable or around the lower limit of quantification of the assay). An in vivo mass balance study in rats showed that LOKELMA was recovered in the feces with no evidence of systemic absorption.

Drug Interactions
Thirty-nine (39) drugs were tested to determine potential interactions with LOKELMA. Sixteen (16) drugs tested did not show an in vitro interaction with LOKELMA (allopurinol, apixaban, aspirin, captopril, cyclosporine, digoxin, ethinyl estradiol, lisinopril, magnesium, metformin, phenytoin, prednisone, propranolol, quinapril, spironolactone and ticagrelor).

Nine (9) of the 23 drugs that showed an in vitro interaction were subsequently tested in vivo in healthy volunteers. Losartan, glipizide and levothyroxine did not show any changes in exposure when co-administered with LOKELMA. However, there was an increase in systemic exposure to weak acids such as furosemide and atorvastatin, and a decrease in systemic exposure to weak bases such as dabigatran when co-administered with LOKELMA, as shown in Figure 2. These changes are consistent with the hypothesis that LOKELMA, by elevating gastric pH, affects the systemic exposure of co-administered drugs whose solubility is pH-dependent [see Drug Interactions (7)].
LOKELMA™ (sodium zirconium cyclosilicate) for oral suspension

3.0

had a greater response to LOKELMA. LOKELMA was effective in lowering potassium at 2.5, 5, and 10 g. In patients administered 10 g TID, the mean serum potassium from baseline, LOKELMA showed dose-dependent reductions in serum potassium < 0.001). As displayed in Table 1 for the secondary endpoint of potassium change in serum potassium levels during the initial 48 hours of study drug treatment, comparing patients receiving LOKELMA and patients receiving placebo. The study met the primary efficacy endpoint at the 5 and 10 g doses when compared with their respective placebo groups (p<0.01 and p<0.001).

### 13.1 Carcinogenesis, Mutagenesis, Impairment of Fertility

The following tests for mutagenic potential of sodium zirconium cyclosilicate were negative: (1) the Ames (S. typhimurium and E. coli) test; (2) chromosomal aberration assay in Chinese Hamster Ovary (CHO) cells; and (3) in vivo rat micronucleus assay. Given that zirconium cyclosilicate is not genotoxic, not absorbed from the gastrointestinal tract, and did not cause local gastrointestinal alterations in a chronic toxicity study in dogs, carcinogenicity studies in animals to evaluate tumorigenic potential of sodium zirconium cyclosilicate were not deemed to be necessary.

Fertility in male and female rats has been assessed at doses up to a Human Equivalent Dose (HED) of 58 g per day (the maximum feasible dose) with no adverse effects.

### 14 CLINICAL STUDIES

#### 14.1 Study 1

The effectiveness of LOKELMA in lowering serum potassium was demonstrated in a two-part, double-blind, randomized, placebo-controlled clinical trial (NCT01737697) in patients with hyperkalemia (5 to 6.5 mEq/L, mean potassium 5.3 mEq/L), Study 1. In the first phase of the trial (the acute phase), 753 patients were randomized to receive one of four doses of LOKELMA (1.25, 2.5, 5, or 10 g of once daily LOKELMA for 12 days together with breakfast. Of the patients enrolled in the acute phase, 92% achieved a potassium level within this range and were enrolled into the second phase of the trial.

Following the acute phase of the study, there was a double-blind randomized withdrawal phase where patients who achieved potassium levels between 3.5 and 5 mEq/L were randomized to one of three doses of LOKELMA administered once-daily for 28 days, or placebo just before breakfast. Of the patients enrolled in the acute phase, 92% achieved a potassium level within this range and were enrolled into the second phase of the trial.

#### 14.2 Study 2

The efficacy of LOKELMA was also demonstrated in a two-part trial with an open-label acute phase and a month-long randomized, double-blind, placebo-controlled withdrawal phase (Study 2; NCT02088073).

In the open-label acute phase of Study 2, 258 patients with hyperkalemia (baseline mean 5.6 mEq/L, range 5.1 to 7.4 mEq/L) received 10 g of LOKELMA administered three times daily with meals for 48 hours. As shown in Figure 3, left, average serum potassium levels decreased from 5.6 to 4.5 mEq/L during treatment with LOKELMA in the acute phase.

Following the acute phase of the study, there was a double-blind randomized withdrawal phase where patients who achieved potassium levels between 3.5 and 5 mEq/L were randomized to one of three doses of LOKELMA administered once-daily for 28 days, or placebo just before breakfast. Of the patients enrolled in the acute phase, 92% achieved a potassium level within this range and were enrolled into the second phase of the trial.

Patients who achieved a potassium level between 3.5 and 5 mEq/L after receiving LOKELMA during the acute phase were re-randomized to receive once daily placebo or 1.25, 2.5, 5, or 10 g of once daily LOKELMA for 12 days together with breakfast. The primary endpoint in the maintenance phase was the difference in the exponential rate of change in serum potassium levels over the 12-day treatment interval, comparing patients receiving LOKELMA and patients receiving placebo. The study met the primary efficacy endpoint at the 5 and 10 g doses when compared with their respective placebo groups (p<0.01 and p<0.001).

### Figure 2: Effects of LOKELMA on the Pharmacokinetic Exposures of Other Orally Administered Medications

<table>
<thead>
<tr>
<th>Drug</th>
<th>AUC</th>
<th>Cmax</th>
</tr>
</thead>
<tbody>
<tr>
<td>amiodipine 5 mg</td>
<td></td>
<td></td>
</tr>
<tr>
<td>atorvastatin 10 mg</td>
<td></td>
<td></td>
</tr>
<tr>
<td>o-H atorvastatin</td>
<td></td>
<td></td>
</tr>
<tr>
<td>p-H atorvastatin</td>
<td></td>
<td></td>
</tr>
<tr>
<td>clopidogrel 75 mg</td>
<td></td>
<td></td>
</tr>
<tr>
<td>clopidogrel acid</td>
<td></td>
<td></td>
</tr>
<tr>
<td>dabigatran 75 mg</td>
<td></td>
<td></td>
</tr>
<tr>
<td>furosemide 20 mg</td>
<td></td>
<td></td>
</tr>
<tr>
<td>glipizide 5 mg</td>
<td></td>
<td></td>
</tr>
<tr>
<td>levothyroxine 50 ug</td>
<td></td>
<td></td>
</tr>
<tr>
<td>thyroxine (T4)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>triido-L-thyroxine (T3)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>losartan 25 mg</td>
<td></td>
<td></td>
</tr>
<tr>
<td>losartan acid</td>
<td></td>
<td></td>
</tr>
<tr>
<td>(R,S)-warfarin 5 mg</td>
<td></td>
<td></td>
</tr>
<tr>
<td>R-warfarin</td>
<td></td>
<td></td>
</tr>
<tr>
<td>S-warfarin</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

#### Table 1: Study 1 - Potassium Change from Baseline to 48 hours

<table>
<thead>
<tr>
<th>Mean Serum Potassium</th>
<th>Placebo</th>
<th>1.25 g TID</th>
<th>2.5 g TID</th>
<th>5 g TID</th>
<th>10 g TID</th>
</tr>
</thead>
<tbody>
<tr>
<td>% change mEq/L</td>
<td></td>
<td>(95% CI)</td>
<td>(95% CI)</td>
<td>(95% CI)</td>
<td>(95% CI)</td>
</tr>
<tr>
<td>All Patients</td>
<td></td>
<td>(-0.2, -0.1)</td>
<td>(-0.3, -0.2)</td>
<td>(-0.4, -0.2)</td>
<td>(-0.5, -0.4)</td>
</tr>
<tr>
<td>(n=192)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Baseline Serum Potassium &gt;5.5 mEq/L</td>
<td></td>
<td>(-0.4, -0.3)</td>
<td>(-0.5, -0.2)</td>
<td>(-0.6, -0.3)</td>
<td>(-0.6, -0.5)</td>
</tr>
<tr>
<td>(n=140)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

### Figure 3: Study 2 - Mean Serum Potassium Levels in the Acute and Randomized Withdrawal Phases

Intent-to-Treatment population includes subjects with at least one valid serum potassium measurement on or after Day 8.
14.3 Eleven-Month Extension Study
Patients who completed the 28-day randomized withdrawal phase had the option to continue treatment with LOKELMA, taken just before breakfast, in an open-label extension phase for up to 11 months (n=123; NCT02107092). Figure 4 shows that the treatment effect on serum potassium was maintained during continued therapy.

Figure 4: 11-Month Open-Label Extension Phase of Study 2 – Mean Serum Potassium (mEq/L)

14.4 Study 3
LOKELMA was evaluated in an open-label 12-month study in 751 hyperkalemic patients (NCT02163499). The mean baseline potassium level in this study was 5.6 mEq/L. Following the acute phase treatment of LOKELMA 10 g three times a day, patients who achieved normokalemia (3.5-5.0 mEq/L) within 72 hours (n=746; 99%) entered the maintenance phase. For maintenance treatment, the initial dosage of LOKELMA was 5 g once daily and was adjusted to a minimum of 5 g every other day up to maximum of 15 g once daily, based on serum potassium level. The treatment effect on serum potassium was maintained during continued therapy.

16 HOW SUPPLIED/STORAGE AND HANDLING
LOKELMA (sodium zirconium cyclosilicate) for oral suspension is supplied as a white powder in foil-lined packets as follows:

<table>
<thead>
<tr>
<th>LOKELMA (grams)</th>
<th>Single Packet</th>
<th>Box of 11 Packets</th>
<th>Box of 30 Packets</th>
</tr>
</thead>
<tbody>
<tr>
<td>5</td>
<td>NDC 0310-1105-01</td>
<td>NDC 0310-1105-39</td>
<td>NDC 0310-1105-30</td>
</tr>
<tr>
<td>10</td>
<td>NDC 0310-1110-01</td>
<td>NDC 0310-1110-39</td>
<td>NDC 0310-1110-30</td>
</tr>
</tbody>
</table>

Storage and Handling
Store LOKELMA at 15°C-30°C (59°F-86°F).

17 PATIENT COUNSELING INFORMATION

Dosing
Instruct the patient how to reconstitute LOKELMA for administration. Inform the patient that it is necessary to drink the full dose [see Dosage and Administration (2.2)].

Drug Interactions
Advise patients who are taking other oral medications to separate dosing of LOKELMA by at least 2 hours (before or after) [see Drug Interactions (7)].

Diet
Advise patients to adjust dietary sodium, if appropriate [see Warnings and Precautions (5.2)].

U.S. Patent No: 6332985, 8808750, 8877255, 8802152, 9592253
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Manufactured by: AstraZeneca Pharmaceuticals LP, Wilmington, DE 19850
07/18 US-22303 8/18