LOKELMA™
(sodium zirconium cyclosilicate)
10 g for oral suspension

YOU ARE INVITED TO ATTEND

Rethinking the Approach to the Treatment of Hyperkalemia

October 24, 2019
06:00 PM – 08:00 PM
Central Standard Time

LOCATION
Oceans at Arthur's
16100 Chenal Parkway
Little Rock, AR 72211

PRESENTED BY
Anand Reddy, MD, MRCP
Nephrologist
Odessa Medical Center, Midland Memorial, Medical Center Hospital in Odessa, Texas

RSVP IS REQUIRED BY: 10/21/2019

To find out more information or to register, please contact
JENNIFER HANNAH
Text 501-804-1964 jennifer.hannah1@astrazeneca.com

IMPORTANT SAFETY INFORMATION

WARNINGS AND PRECAUTIONS:

- **Gastrointestinal Adverse Events in Patients with Motility Disorders:** Avoid LOKELMA in patients with severe constipation, bowel obstruction or impaction, including abnormal post-operative bowel motility disorders. LOKELMA has not been studied in patients with these conditions and it may be ineffective and may worsen gastrointestinal conditions
- **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 (eg., heart failure or renal disease). Advise patients to adjust dietary sodium, if appropriate. Increase the dose of diuretics as needed

ADVERSE REACTIONS: The most common adverse reaction with LOKELMA was mild to moderate edema. In placebo-controlled trials up to 28 days, edema was reported in 4.4%, 5.9%, 16.1% of patients treated with 5 g, 10 g and 15 g of LOKELMA once daily, respectively vs 2.4% of patients receiving placebo.

DRUG INTERACTIONS: LOKELMA can transiently increase gastric pH. In general, oral medications with pH-dependent solubility should be administered at least 2 hours before or 2 hours after LOKELMA. Spacing is not needed if it has been determined the concomitant medication does not exhibit pH-dependent solubility.

INDICATION AND LIMITATION OF USE

LOKELMA is indicated for the treatment of hyperkalemia in adults.

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

Please read accompanying full Prescribing Information.

AstraZeneca

You are encouraged to report negative side effects of AstraZeneca prescription drugs by calling 1-800-236-9933. If you prefer to report these to the FDA, either visit www.FDA.gov/medwatch or call 1-800-FDA-1088.

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