Learn more about the once-daily oral treatment option LUMAKRAS®[1]



What is the recommended dose?

- **LUMAKRAS® recommended once-daily, starting dosage: 960 mg orally[1]**
- **LUMAKRAS® 240 mg tablets have a yellow color. They come in one bottle containing 120 tablets.[1]**
- **LUMAKRAS® tablets are comparable in size to a dime.[1,2]** Speak with your doctor if you have any questions or if you experience a side effect.



Tablet as seen may not reflect actual size.



How to take LUMAKRAS®

Take your prescribed dose of LUMAKRAS®:



Once daily by mouth[3]



At the same time every day[3]



With or without food[3]

When taking your daily dose of LUMAKRAS®

- ○ Swallow LUMAKRAS® tablets whole. **Do not chew, crush, or split tablets[3]**
- ○ Take LUMAKRAS® exactly as your doctor tells you to take it. Do not change your dose or stop taking LUMAKRAS® **unless your doctor tells you to[3]**
- ○ Take the number of tablets (pills) your doctor prescribes. **If you have a side effect, talk to your doctor[3]**

What is LUMAKRAS®?

LUMAKRAS® is a prescription medicine used to treat adults with non-small cell lung cancer (NSCLC):

- that has spread to other parts of the body or cannot be removed by surgery, and
- whose tumor has an abnormal KRAS G12C gene, and
- who have received at least one prior treatment for their cancer.

Your healthcare provider will perform a test to make sure that LUMAKRAS® is right for you. It is not known if LUMAKRAS® is safe and effective in children.

Important Safety Information

What should I tell my healthcare provider before taking LUMAKRAS®?

- Before taking LUMAKRAS®, tell your healthcare provider about all your medical conditions, including if you:
  - have liver problems
  - have lung or breathing problems other than lung cancer
  - are pregnant or plan to become pregnant. It is not known if LUMAKRAS® will harm your unborn baby.
  - are breastfeeding or plan to breastfeed. It is not known if LUMAKRAS® passes into your breast milk. Do not breastfeed during treatment with LUMAKRAS® and for 1 week after the last dose.
- Tell your healthcare provider about all the medicines you take, including prescription and over-the-counter medicines, vitamins, dietary and herbal supplements. LUMAKRAS® can affect the way some other medicines work, and some other medicines can affect the way LUMAKRAS® works.
- Especially tell your healthcare provider if you take antacid medicines, including Proton Pump Inhibitor (PPI) medicines or H_2 blockers during treatment with LUMAKRAS®. Ask your healthcare provider if you are not sure.

Please see additional Important Safety Information for LUMAKRAS® on the next page.

ONCE-DAILY ORAL
LUMAKRAS®
(sotorasib) 240 mg tablets



If you cannot swallow LUMAKRAS® tablets whole:[3]

- Place your prescribed dose of LUMAKRAS® in a glass of 4 ounces (120 mL) of non-carbonated, room temperature water without crushing the tablets. Do not use any other liquids.
- Stir or swirl the cup for about **3 minutes** until the tablets are in small pieces (**the tablets will not completely dissolve**). The color of the mixture may be pale yellow to bright yellow.
- Drink the LUMAKRAS® and water mixture right away or within 2 hours of preparing. Do not chew pieces of the tablet.
- Rinse the glass with an additional 4 ounces (120 mL) of water and drink to make sure that you have taken the full dose of LUMAKRAS®.
- If you do not drink the mixture right away, stir or swirl the mixture again before drinking.

Important things to remember when taking LUMAKRAS®.[3]

- If you take an **antacid medicine** such as Tums®, take LUMAKRAS® either 4 hours before or 10 hours after the antacid. Speak to your doctor if you are on PPIs or H_2 blockers.
- If you **miss a dose** of LUMAKRAS®, take the dose as soon as you remember. If it has been more than 6 hours, do not take the dose. Take your next dose at your regularly scheduled time the next day. Do not take 2 doses at the same time to make up for a missed dose.
- If you **vomit after taking a dose** of LUMAKRAS®, do not take an extra dose. Take your next dose at your regularly scheduled time the next day.

Important Safety Information (cont'd)

LUMAKRAS® may cause serious side effects, including:

- Liver problems: LUMAKRAS® may cause abnormal liver blood test results. Your healthcare provider should do blood tests before starting and during treatment with LUMAKRAS® to check your liver function. Tell your healthcare provider right away if you get any signs or symptoms of liver problems, including: your skin or the white part of your eyes turns yellow (jaundice), dark or "tea-colored" urine, light-colored stools (bowel movements), tiredness or weakness, nausea or vomiting, bleeding or bruising, loss of appetite, and pain, aching, or tenderness on the right side of your stomach-area (abdomen).
- Lung or breathing problems: LUMAKRAS® may cause inflammation of the lungs that can lead to death. Tell your healthcare provider or get emergency medical help right away if you have new or worsening shortness of breath, cough or fever.
- Your healthcare provider may change your dose, temporarily stop, or permanently stop treatment with LUMAKRAS® if you develop side effects.

The most common side effects

- The most common side effects of LUMAKRAS® include diarrhea, muscle or bone pain, nausea, tiredness, liver problems, cough, changes in liver function tests, and changes in certain blood tests.
- These are not all the possible side effects of LUMAKRAS®. Call your doctor for medical advice about side effects.

Please see LUMAKRAS® Patient Information.

References: 1. LUMAKRAS® (sotorasib) Prescribing Information, Amgen. **2.** Data on file, Amgen; [Sotorasib Tablet Size]. **3.** LUMAKRAS® (sotorasib) Patient Information, Amgen.

For additional information visit LUMAKRAS.com or use the QR code

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ONCE-DAILY ORAL LUMAKRAS® (sotorasib) 240 mg tablets