

Abridged Prescribing Information

Belumosudil Tablets 200mg

REZUROCK®

Composition: Each film-coated tablet contains 200 mg belumosudil (equivalent to 242.5 mg belumosudil mesylate)

Indication: Rezurock® is indicated for the treatment of patients 12 years and older with chronic graft-versus-host disease (chronic GvHD) after failure of at least two prior lines of systemic therapy.

Dosage and administration: The recommended dose of Rezurock® is 200 mg given orally once daily at approximately the same time each day with a meal. If the patient misses a dose of Rezurock®, instruct the patient not to take extra doses to make up the missed dose.

Safety related information

Special populations: The safety and effectiveness of belumosudil tablets in pediatric patients aged less than 12 years have not been established. No dose adjustment is recommended over the age of 65 years, patients with mild or moderate renal impairment and patients with mild or moderate hepatic impairment (Child-Pugh A and B).

Contraindications: Hypersensitivity to this drug or to any of the excipients.

Warnings and precautions: Rezurock® can cause fetal harm when administered to a pregnant woman based on animal findings and mechanism of action. Advise pregnant women of the potential risk to a fetus. Advise females of reproductive potential and males with female partners of reproductive potential to use effective contraception during treatment with belumosudil and for at least one week after the last dose of belumosudil

Interactions: Coadministration of Rezurock® with strong CYP3A inducers and proton pump inhibitors decreases belumosudil exposure, which may reduce the efficacy of belumosudil. For detailed information, please refer to the interaction section in the prescribing information.

Effects on ability to drive: Rezurock® has no influence on the ability to drive and use machines.

Pregnancy and Lactation: Rezurock® is not recommended during pregnancy or breast-feeding.

Undesirable effects: The most common side effects of Rezurock® include tiredness or weakness, diarrhea, infections, nausea, shortness of breath, cough, swelling, high blood pressure, headache, muscle or bone pain and stomach (abdominal) pain.

For full prescribing information please contact: Sanofi Healthcare India Pvt. Ltd., Sanofi House, CTS No. 117-B, L&T Business Park, Saki Vihar Road, Powai, Mumbai – 400072

Updated: Dec 2024

Source: CCDS v2 dated 3rd Aug 2023