REZUROCK®
(belumosudil) tablets

PRODUCT ORDERING FORM

Distributed and marketed by	Kadmon, a Sanofi Company - Phone: 800-981-2491 - Website: Sanofi.us
Product name	REZUROCK
Generic name	belumosudil
Product website	REZUROCKhcp.com
Indication[1]	REZUROCK is indicated for the treatment of adult and pediatric patients 12 years and older with chronic graft-versus-host disease (chronic GVHD) after failure of at least two prior lines of systemic therapy.
Product information[1]	**NDC:** 79802-200-30 **Description:** REZUROCK 200 mg tablets (30-count bottle)
Wholesale acquisition price	$17,874.83 per 30-count bottle
Ordering REZUROCK	**Contact any of the authorized specialty distributors below to order REZUROCK for your account.**

PHYSICIAN DISPENSING OFFICES

Cardinal Health™ Specialty Pharmaceutical Distribution
Phone: 1-877-453-3972
Fax: 1-877-274-9897
specialtyonline.cardinalhealth.com
Item number: 5699061

McKesson Specialty Health
Phone: 1-800-482-6700
Fax: 1-800-289-9285
mscs.mckesson.com
Item number: 5011227

Oncology Supply®
Phone: 1-800-633-7555
Fax: 1-800-248-8205
oncologysupply.com
Item number: 10260202

INSTITUTIONS/HOSPITALS

ASD Healthcare®
Phone: 1-800-746-6273
Fax: 1-800-547-9413
asdhealthcare.com
Item number: 10260202

Cardinal Health™ Specialty Pharmaceutical Distribution
Phone: 1-855-855-0708
Fax: 1-877-274-9897
orderexpress.cardinalhealth.com
Item number: 5699061

McKesson Plasma and Biologics
Phone: 1-877-625-2566
Fax: 1-888-752-7626
connect.mckesson.com
Item number: 2342103

Contact any of the authorized specialty pharmacies below to help get your patients started on REZUROCK.

Amber Specialty Pharmacy
Phone: 1-888-370-1724
Fax: 1-402-896-3774
amberpharmacy.com

Biologics by McKesson
Phone: 1-800-850-4306
Fax: 1-800-823-4506
biologics.mckesson.com

Onco360 Oncology Pharmacy
Phone: 1-877-662-6633
Fax: 1-877-662-6355
onco360.com

INDICATION

REZUROCK® (belumosudil) is indicated for the treatment of adult and pediatric patients 12 years and older with chronic graft-versus-host disease (chronic GVHD) after failure of at least two prior lines of systemic therapy.

IMPORTANT SAFETY INFORMATION

Warnings and Precautions

- **Embryo-Fetal Toxicity:** Based on findings in animals and its mechanism of action, REZUROCK can cause fetal harm when administered to a pregnant woman. 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 REZUROCK and for one week after the last dose

Adverse Reactions

- The most common (≥ 20%) adverse reactions, including laboratory abnormalities, were infections, asthenia, nausea, diarrhea, dyspnea, cough, edema, hemorrhage, abdominal pain, musculoskeletal pain, headache, phosphate decreased, gamma glutamyl transferase increased, lymphocytes decreased, and hypertension

Please see additional Important Safety Information on the next page. Please see full Prescribing Information.

REZUROCK®
(belumosudil) tablets

Dispensing pack dimensions	**Depth:** 1.9 inches **Height:** 3.875 inches **Width:** 1.938 inches
How supplied[1]	REZUROCK 200 mg tablets are supplied as pale yellow film-coated oblong tablets containing 200 mg of belumosudil (equivalent to 242.5 mg belumosudil mesylate). Each tablet is debossed with "KDM" on one side and "200" on the other side and is packaged as follows: 200 mg tablets in 30 count bottle; NDC 79802-200-30.
Storage and handling[1]	Store at room temperature, 20°C to 25°C (68°F to 77°F); excursions permitted from 15°C and 30°C (59°F to 86°F). Dispense to patient in original container only. Store in original container to protect from moisture. Replace cap securely each time after opening. Do not discard desiccant.
Packaging[1]	**Sales unit:** One 30-count bottle **Units per case:** 24
Product expiration	Expiration date printed on both 30-count bottle and carton.
Product returns	**Phone:** 888-379-6847 **Email:** customersupport@sanofi.com
Product information	**Phone:** 1-800-633-1610 **Website:** www.sanofimedicalinformation.com
Reimbursement and patient support	Kadmon ASSIST™ **Phone:** 1-844-KADMON1 (523-6661) **Fax:** 1-833-635-1481 **Website:** KadmonASSIST.com

IMPORTANT SAFETY INFORMATION (cont)

Adverse Reactions (cont)

- Permanent discontinuation of REZUROCK due to adverse reactions occurred in 18% of patients. The adverse reactions which resulted in permanent discontinuation of REZUROCK in > 3% of patients included nausea (4%). Adverse reactions leading to dose interruption occurred in 29% of patients. The adverse reactions leading to dose interruption in ≥ 2% were infections (11%), diarrhea (4%), and asthenia, dyspnea, hemorrhage, hypotension, liver function test abnormal, nausea, pyrexia, edema, and renal failure with (2% each)
- Monitor total bilirubin, aspartate aminotransferase (AST), and alanine aminotransferase (ALT) at least monthly

Drug Interactions

- **Strong CYP3A Inducers:** Coadministration of REZUROCK with strong CYP3A inducers decreases belumosudil exposure, which may reduce the efficacy of REZUROCK. Increase the dosage of REZUROCK to 200 mg twice daily when coadministered with strong CYP3A inducers
- **Proton Pump Inhibitors:** Coadministration of REZUROCK with proton pump inhibitors decreases belumosudil exposure, which may reduce the efficacy of REZUROCK. Increase the dosage of REZUROCK to 200 mg twice daily when coadministered with proton pump inhibitors

Use in Specific Populations

- **Pregnancy:** There are no available human data on REZUROCK use in pregnant women to evaluate for a drug-associated risk. Advise pregnant women and females of reproductive potential of the potential risk to the fetus
- **Lactation:** There are no data available on the presence of belumosudil or its metabolites in human milk or the effects on the breastfed child, or milk production. Because of the potential for serious adverse reactions from belumosudil in the breastfed child, advise lactating women not to breastfeed during treatment with REZUROCK and for one week after the last dose
- **Pediatric Use:** The safety and effectiveness of REZUROCK in pediatric patients less than 12 years old have not been established
- **Geriatric Use:** Of the 186 patients with chronic GVHD in clinical studies of REZUROCK, 26% were 65 years and older. No clinically meaningful differences in safety or effectiveness of REZUROCK were observed in comparison to younger patients
- **Renal Impairment:** Treatment with REZUROCK has not been studied in patients with pre-existing severe renal impairment. For patients with pre-existing severe renal impairment, consider the risks and potential benefits before initiating treatment with REZUROCK
- **Hepatic Impairment:** Avoid use in patients with moderate hepatic impairment (Child-Pugh B) or severe hepatic impairment (Child-Pugh C) without liver GVHD. No dose adjustment is recommended for patients with mild hepatic impairment (Child-Pugh A)

Please see additional Important Safety Information on the previous page. Please see full Prescribing Information.

Reference: 1. REZUROCK. Package insert. Kadmon Pharmaceuticals, LLC; 2023.

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