CONTACT DETAILS FOR **WHOLESALERS AND SPECIALTY DISTRIBUTORS**

Wholesalers and Specialty Distributors handle the distribution of specialty products to Specialty Pharmacies, health systems, and healthcare providers.

HOW SUPPLIED	NDC #
SKYRIZI 600 mg/10 mL single-dose vial	0074-5015-01

SKYRIZI® (risankizumab-rzaa) Distribution Network

All adult SKYRIZI patients with moderately to severely active Crohn's disease will receive 3 induction doses of SKYRIZI through intravenous (IV) infusion before transitioning to the SKYRIZI prefilled cartridge with On-Body Injector for maintenance dosing.

Wholesalers and Specialty Distributors authorized to supply all presentations of SKYRIZI include, but are not limited to, the following:

NAME	TELEPHONE #	WEBSITE/E-MAIL
AmerisourceBergen	1-844-222-2273	www.amerisourcebergen.com
AmerisourceBergen – ASD Healthcare	1-800-746-6273	www.asdhealthcare.com
AmerisourceBergen – Besse Medical	1-800-543-2111	www.besse.com
AmerisourceBergen – Oncology Supply	1-800-633-7555	www.oncologysupply.com
Cardinal Health Specialty Pharmaceutical Distribution	1-800-926-3161	www.cardinalhealth.com
Cardinal Health Specialty Solutions	1-855-855-0708	specialtyonline.cardinalhealth.com
Metro Medical	1-800-768-2002	www.metromedicalorder.com
McKesson U.S. Pharmaceutical	1-855-625-7385 – Independent Pharmacies 1-855-625-6285 – Retail National Account 1-855-625-4677 – Hospitals and Health System	www.mckesson.com
McKesson Medical-Surgical	1-855-571-2100	www.mms.mckesson.com
McKesson Specialty Health	1-800-482-6700	www.mckesson.com
CuraScript SD	1-877-599-7748	www.curascriptsd.com
Henry Schein, Inc.	1-800-472-4346	www.henryschein.com
Morris & Dickson Co., LLC	1-800-388-3833	www.morrisdickson.com
Smith Drug Company	1-800-572-1216	www.smithdrug.com
Value Drug Company	1-800-252-3786	www.valuedrugco.com

For a complete list of AbbVie authorized distributors, please contact Customer Service at (800) 255-5162.

SKYRIZI On-Body Injector is available via open distribution, and prescriptions may be sent to the Specialty Pharmacy of the patient's choosing.

INDICATION[1]

SKYRIZI is indicated for the treatment of moderately to severely active Crohn's disease in adults.

SAFETY CONSIDERATIONS[1]

SKYRIZI is contraindicated in patients with a history of serious hypersensitivity reaction to risankizumab-rzaa or any of its excipients. Serious hypersensitivity reactions, including anaphylaxis, have been reported with use of SKYRIZI. If a serious hypersensitivity reaction occurs, discontinue SKYRIZI and initiate appropriate therapy immediately. SKYRIZI may increase the risk of infection. Instruct patients to report signs or symptoms of clinically important infection during treatment. Should such an infection occur, discontinue SKYRIZI until infection resolves. Evaluate patients for tuberculosis infection prior to initiating treatment with SKYRIZI. Drug-induced liver injury was reported in a patient with Crohn's disease during induction dosing of SKYRIZI. For the treatment of Crohn's disease, evaluate liver enzymes and bilirubin at baseline and during induction (12 weeks). Interrupt treatment with SKYRIZI if drug-induced liver injury is suspected, until this diagnosis is excluded. Avoid use of live vaccines in SKYRIZI patients.

Please see additional Important Safety Information on the next page.

Please see accompanying full Prescribing Information.

Skyrizi®
risankizumab-rzaa

IMPORTANT SAFETY INFORMATION[1]

Hypersensitivity Reactions

SKYRIZI® (risankizumab-rzaa) is contraindicated in patients with a history of serious hypersensitivity reaction to risankizumab-rzaa or any of the excipients. Serious hypersensitivity reactions, including anaphylaxis, have been reported with the use of SKYRIZI. If a serious hypersensitivity reaction occurs, discontinue SKYRIZI and initiate appropriate therapy immediately.

Infection

SKYRIZI may increase the risk of infection. Do not initiate treatment with SKYRIZI in patients with a clinically important active infection until it resolves or is adequately treated.

In patients with a chronic infection or a history of recurrent infection, consider the risks and benefits prior to prescribing SKYRIZI. Instruct patients to seek medical advice if signs or symptoms of clinically important infection occur. If a patient develops such an infection or is not responding to standard therapy, closely monitor and discontinue SKYRIZI until the infection resolves.

Tuberculosis (TB)

Prior to initiating treatment with SKYRIZI, evaluate for TB infection and consider treatment in patients with latent or active TB for whom an adequate course of treatment cannot be confirmed. Monitor patients for signs and symptoms of active TB during and after SKYRIZI treatment. Do not administer SKYRIZI to patients with active TB.

Hepatotoxicity in Treatment of Crohn's Disease

Drug-induced liver injury was reported in a patient with Crohn's disease who was hospitalized for a rash during induction dosing of SKYRIZI. For the treatment of Crohn's disease, evaluate liver enzymes and bilirubin at baseline and during induction (12 weeks); monitor thereafter according to routine patient management. Consider an alternate treatment for patients with evidence of liver cirrhosis. Interrupt treatment if drug-induced liver injury is suspected, until this diagnosis is excluded. Instruct your patient to seek immediate medical attention if they experience symptoms suggestive of hepatic dysfunction.

Administration of Vaccines

Avoid use of live vaccines in patients treated with SKYRIZI. Medications that interact with the immune system may increase the risk of infection following administration of live vaccines. Prior to initiating SKYRIZI, complete all age appropriate vaccinations according to current immunization guidelines.

Adverse Reactions

Most common (>3%) adverse reactions associated with SKYRIZI in Crohn's disease are upper respiratory infections, headache, and arthralgia in induction and arthralgia, abdominal pain, injection site reactions, anemia, pyrexia, back pain, arthropathy, and urinary tract infection in maintenance.

Lipid Elevations: Increases from baseline and increases relative to placebo were observed at Week 4 and remained stable to Week 12 in patients treated with SKYRIZI in Crohn's disease.

Dosage Forms and Strengths: SKYRIZI is available in a 600 mg/10 mL single-dose vial for intravenous infusion and a 180 mg/1.2 mL or 360 mg/2.4 mL single-dose prefilled cartridge with on-body injector.

Please see accompanying full Prescribing Information.

Reference: 1. SKYRIZI [package insert]. North Chicago, IL: AbbVie Inc.

abbvie

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