

Contact Details for Specialty Pharmacies

SKYRIZI is broadly available via open distribution and prescriptions may be sent to the specialty pharmacy of the patient's choosing.



Specialty Pharmacy contact details

Pharmacy Name	Telephone	Rx Email	ePrescription	Fax
Absolute Pharmacy Inc.	787-892-8700	Rxsend@absolutepharmacypr.com	Absolute Pharmacy	787-496-1010
Alivia Specialty Pharmacy	787-925-1999	farmaciaespecializada@aliviahealth.com	AliviaSpecialtyPharmacy NAPB:4030057	787-925-1015 787-723-6987
Axium Healthcare Pharmacy Puerto Rico	787-780-7200 844-355-4191		Axium Healthcare de PR	800-546-2163
CVS Caremark Specialty Pharmacy	787-759-4160	N/A	CVS Specialty	787-759-4162
Farmacia San Rafael Specialty	787-724-3333	farmacia@fsanrafael.com		787-919-7058
Grafed Specialty Pharmacy	787-847-9393 939-731-7109	receta@grafet.com	Grafet Specialty Pharmacy	787-847-9292
Optima Health Pharmacy	787-883-5957 787-907-1801	farmacia@optimahealthpr.com	Optima Health Dorado	787-883-6040
Special Care Pharmacy Services	787-781-4585 1-877-899-8997	rxmail@scpspr.com		787-783-2951 1-855-230-9963
Specialty Pharmacy Services- Caguas	787-704-2025 787-704-2028	SPSpharmacy@SPScaguas.com		787-704-2027 787-743-3005
Walgreens Specialty Pharmacy	787-777-1120	rxm.15191@store.walgreens.com rxm.21385@store.walgreens.com		787-777-1124 787-758-3730
Medplus Specialty /PRRX Specialty	787-523-3888	sprx@medpluspr.com	Medplus A+	

The pharmacies listed here have not endorsed and are not affiliated with this material.

Resources, support and education are available for your patients at www.AbbvieContigo.com or by calling 1-855-266-8446



Scan this QR code or visit SkyriziHCPPR.com to learn more about SKYRIZI

Please see Indication and Important Safety Information on back page.
Please see accompanying full [Prescribing Information](#).

abbvie


Skyrizi[®]
risankizumab-rzaa

Indications and Important Safety Information for SKYRIZI

Indication¹

Plaque Psoriasis: SKYRIZI is indicated for the treatment of moderate to severe plaque psoriasis in adults who are candidates for systemic therapy or phototherapy.

Psoriatic Arthritis: SKYRIZI is indicated for the treatment of active psoriatic arthritis in adults.

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

Ulcerative Colitis: SKYRIZI is indicated for the treatment of moderately to severely active ulcerative colitis in adults.

IMPORTANT SAFETY INFORMATION¹

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 Inflammatory Bowel 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 and ulcerative colitis, 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 ($\geq 1\%$) adverse reactions associated with SKYRIZI in plaque psoriasis and psoriatic arthritis include upper respiratory infections, headache, fatigue, injection site reactions, and tinea infections.

In psoriatic arthritis phase 3 trials, the incidence of hepatic events was higher with SKYRIZI compared to placebo.

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.

Most common ($\geq 3\%$) adverse reactions associated with SKYRIZI in ulcerative colitis are arthralgia in induction, and arthralgia, pyrexia, injection site reactions, and rash 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. Lipid elevations observed in patients with ulcerative colitis were similar to those in Crohn's disease.

Dosage Forms and Strengths: SKYRIZI (risankizumab-rzaa) is available in a 150 mg/mL prefilled syringe and pen, 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.

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

Please see accompanying full **Prescribing Information** for additional information or visit https://www.rxabbvie.com/pdf/skyrizi_pi.pdf.

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Skyrizi[®]
risankizumab-rzaa