



For Adults With Moderate to Severe Plaque Psoriasis

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THE OPPORTUNITY FOR NOTHING ON MY SKIN MEANS EVERYTHING TO ME

NOTHING IS EVERYTHING

IN CLINICAL TRIALS AT 1 YEAR, NEARLY
6 OUT OF 10 PEOPLE ACHIEVED 100% CLEAR SKIN

Also, 3 out of 4 people achieved 90% clearer skin at 4 months.

See what SKYRIZI can do

- RESULTS WITH SKYRIZI
- BEFORE & AFTER PHOTOS
- DOSING FOR SKYRIZI
- SAVE ON SKYRIZI

Coronavirus Update

USE

SKYRIZI is a prescription medicine used to treat adults with moderate to severe plaque psoriasis who may benefit from taking injections or pills (systemic therapy) or treatment using ultraviolet or UV light (phototherapy).

IMPORTANT SAFETY INFORMATION

SKYRIZI may cause serious side effects, including infections. SKYRIZI is a prescription medicine that may lower the ability of your immune system to fight infections and may increase your risk of infections. Your healthcare provider should check you for infections and tuberculosis (TB) before starting treatment with SKYRIZI and may treat you for TB before you begin treatment with SKYRIZI if you have a history of TB or have active TB. Your healthcare provider should watch you closely for signs and symptoms of TB during

See the data

View real results

Get dosing details

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Share your SKYRIZI story

Share your experience with SKYRIZI to help inform and inspire others. To participate, please email us at info@SPEAKnetwork.net or call 877-861-6180.



Understanding plaque psoriasis

What it is. What it isn't. What to know about it, from diagnosis to treatment.

[Find out more](#)



Your dedicated Nurse Ambassador*

Your Skyrizi Complete Nurse Ambassador is committed to getting to know you and your individual SKYRIZI needs.

[Get to know the Nurse Ambassador](#)

*Nurse Ambassadors do not give medical advice and will direct you to your healthcare professional for any treatment-related questions, including further referrals.



Coronavirus Update

USE

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USE

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IMPORTANT SAFETY INFORMATION

What is the most important information I should know about SKYRIZI™ (risankizumab-rzaa)?

SKYRIZI may cause serious side effects, including infections. SKYRIZI is a prescription medicine that may lower the ability of your immune system to fight infections and may increase your risk of infections. Your healthcare provider should check you for infections and tuberculosis (TB) before starting treatment with SKYRIZI and may treat you for TB before you begin treatment with SKYRIZI if you have a history of TB or have active TB. Your healthcare provider should watch you closely for signs and symptoms of TB during and after treatment with SKYRIZI.

- Tell your healthcare provider right away if you have an infection or have symptoms of an infection, including:
 - fever, sweats, or chills
 - muscle aches
 - weight loss
 - cough
 - warm, red, or painful skin or sores on your body different from your psoriasis
 - diarrhea or stomach pain
 - shortness of breath
 - blood in your mucus (phlegm)
 - burning when you urinate or urinating more often than normal

Before using SKYRIZI, tell your healthcare provider about all of your medical conditions, including if you:

- have any of the conditions or symptoms listed in the section "What is the most important information I should know about SKYRIZI?"
- have an infection that does not go away or that keeps coming back.
- have TB or have been in close contact with someone with TB.
- have recently received or are scheduled to receive an immunization (vaccine). You should avoid receiving live vaccines during treatment with SKYRIZI.
- are pregnant or plan to become pregnant. It is not known if SKYRIZI can harm your unborn baby.
- are breastfeeding or plan to breastfeed. It is not known if SKYRIZI passes into your breast milk.

Tell your healthcare provider about all the medicines you take, including prescription and over-the-counter medicines, vitamins, and herbal supplements.

What are the possible side effects of SKYRIZI?



What are the possible side effects of SKYRIZI?

SKYRIZI may cause serious side effects. See "What is the most important information I should know about SKYRIZI?"

The most common side effects of SKYRIZI include upper respiratory infections, fungal skin infections, headache, feeling tired, and injection site reactions.

These are not all the possible side effects of SKYRIZI. Call your doctor for medical advice about side effects.

Use SKYRIZI exactly as your healthcare provider tells you to use it.

US-RISN-180038

Please see the **Full Prescribing Information**, including the **Medication Guide**, for SKYRIZI.

You are encouraged to report negative side effects of prescription drugs to the FDA. Visit www.fda.gov/medwatch or call 1-800-FDA-1088.

If you are having difficulty paying for your medicine, AbbVie may be able to help. Visit AbbVie.com/myAbbVieAssist to learn more.

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US-SKZ-200020

Coronavirus Update

SKYRIZI (risankizumab-rzaa)	Important Safety Information for Patients	Information from AbbVie
Home	Important Safety Information	Contact Us
See SKYRIZI Results	Medication Guide	Privacy Policy
What Is SKYRIZI?	Full Prescribing Information	Terms of Use
What Is Plaque Psoriasis?		Advertising Choices
Find a Dermatologist		Healthcare Professionals Site

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If you are having difficulty paying for your medicine, AbbVie may be able to help. Visit AbbVie.com/myAbbVieAssist to learn more.

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US-SKZ-200020

SKYRIZI (risankizumab-rzaa) Home See SKYRIZI Results What Is SKYRIZI? What Is Plaque Psoriasis? Find a Dermatologist Resource Center About Skyrizi Complete Sitemap	Important Safety Information for Patients Important Safety Information Medication Guide Full Prescribing Information	Information from AbbVie Contact Us Privacy Policy Terms of Use Advertising Choices Healthcare Professionals Site
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Coronavirus Update



FOR HEALTHCARE PROFESSIONALS

HOME CLINICAL EFFICACY DOSING CLINICAL SAFETY ACCESS SUPPORT & RESOURCES

The IL-23 inhibitor from AbbVie indicated for the treatment of moderate-to-severe plaque psoriasis in adults who are candidates for systemic therapy or phototherapy

Nothing less than the opportunity for durable skin clearance; for your patients, that's everything

NOTHING IS EVERYTHING



COVID-19 Update

IMPORTANT SAFETY INFORMATION AND INDICATION FOR SKYRIZI™ (risankizumab-rzaa)

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.
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.

<p>DURABLE.</p> <p>Most patients achieved PASI 90 at Week 16 and maintained it at Week 52[Ⓞ]</p> <p>VIEW TRIALS →</p>	<p>RAPID.</p> <p>Co-primary endpoints of PASI 90 and sPGA 0/1 at Week 16, including improvement 4 weeks after 1st dose[Ⓞ]</p> <p>VIEW TRIALS →</p>	<p>CLEAR.</p> <p>The majority of patients achieved PASI 100 at Week 52[Ⓞ]</p> <p>VIEW TRIALS →</p>	<p>4 DOSES PER YEAR.</p> <p>3-month dosing after 2 initiation doses at Weeks 0 and 4 (150 mg/dose)[Ⓞ]</p> <p>VIEW DOSING →</p>
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Nothing less than a commitment to working toward 100% COMMERCIAL PATIENT ACCESS

[VIEW ACCESS →](#)

PASI 90=90% improvement in Psoriasis Area and Severity Index; PASI 100=100% improvement in Psoriasis Area and Severity Index; sPGA 0/1=static Physician's Global Assessment rating of clear or almost clear.

COVID-19 Update

1 WEEK 16[Ⓞ]:
75% OF SKYRIZI PATIENTS

IMPORTANT SAFETY INFORMATION AND INDICATION FOR SKYRIZI™ (risankizumab-rzaa)[Ⓞ]
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AT WEEK 16[Ⓞ]: 75% OF SKYRIZI PATIENTS

(n=294) ACHIEVED PASI 90

in the UlimMa-2 study vs 2% placebo (n=98) patients
84% of SKYRIZI patients achieved sPGA 0/1 vs 5% placebo patients



UlimMa-1 results at Week 16
PASI 90: SKYRIZI 75% (n=304), placebo 5% (n=102)
sPGA 0/1: SKYRIZI 88% (n=304), placebo 8% (n=102)

Co-primary endpoints for UlimMa-1 and -2 were PASI 90 and sPGA 0/1 response vs placebo at Week 16.

P < 0.001 for comparisons of SKYRIZI vs placebo

STUDY DESIGN:

UlimMa-1 (N=506) and UlimMa-2 (N=491) were replicate phase 3, randomized, double-blind, placebo- and active-controlled studies to evaluate the efficacy and safety of SKYRIZI (150 mg) vs placebo over 16 weeks and biologic active control over 52 weeks in adult patients with moderate to severe plaque psoriasis. [Ⓞ]

VIEW STUDY DESIGN AND BASELINE CHARACTERISTICS >

SEE CLINICAL TRIAL RESULTS INCLUDING PASI 100 DATA →

IMPORTANT SAFETY INFORMATION AND INDICATION FOR SKYRIZI™ (risankizumab-rzaa)[Ⓞ]

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COVID-19 Update

SEE CLINICAL TRIAL RESULTS INCLUDING PASI 100 DATA →



AT WEEK 52^(*):
81% OF SKYRIZI PATIENTS

(n=294)
ACHIEVED PASI 90

in the UtiIMMa-2 study; placebo did not continue beyond Week 16

UtiIMMa-1 results at Week 52
 PASI 90: SKYRIZI 82% (n=304), placebo n/a

Co-primary endpoints for UtiIMMa-1 and -2 were PASI 90 and sPGA 0/1 response vs placebo at Week 16.

*p<0.001 for comparisons of SKYRIZI vs placebo

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[VIEW STUDY DESIGN AND BASELINE CHARACTERISTICS >](#)

SEE CLINICAL TRIAL RESULTS INCLUDING PASI 100 DATA →

COVID-19 Update

IMPORTANT SAFETY INFORMATION AND INDICATION FOR SKYRIZI™ (risankizumab-rzaa)^(*)

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SEE CLINICAL TRIAL RESULTS INCLUDING PASI 100 DATA →

4 DOSES PER YEAR[®]


3-month dosing after 2 initiation doses at Weeks 0 and 4 (150 mg/dose)[®]

VIEW DOSING SCHEDULE →

COVID-19 Update

IMPORTANT SAFETY INFORMATION AND INDICATION FOR SKYRIZI™ (risankizumab-rzaa)[®]
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SKYRIZI™ (risankizumab-rzaa): ... SKYRIZI (risankizumab-rzaa) ...



NOTHING LESS THAN A COMMITMENT TO WORKING TOWARD 100% COMMERCIAL PATIENT ACCESS

Skyrizi™ COMPLETE

Eligible patients may be able to access their SKYRIZI at no charge until their insurance plan covers SKYRIZI

[VIEW SKYRIZI COMPLETE →](#)

COVID-19 Update

IMPORTANT SAFETY INFORMATION AND INDICATION FOR SKYRIZI™ (risankizumab-rzaa)®

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DURABLE.

Most patients achieved PASI 90 at Week 16 and maintained it at Week 52 ⁽²⁾

[VIEW TRIALS →](#)

RAPID.

Co-primary endpoints of PASI 90 and sPGA 0/1 at Week 16, including improvement 4 weeks after 1st dose ⁽³⁾

[VIEW TRIALS →](#)

CLEAR.

The majority of patients achieved PASI 100 at Week 52 ⁽²⁾

[VIEW TRIALS →](#)

4 DOSES PER YEAR.

3 month dosing after 2 initiation doses at Weeks 0 and 4 ⁽²⁾

[VIEW DOSING →](#)

Nothing less than a commitment to working toward

100% COMMERCIAL PATIENT ACCESS

[VIEW ACCESS →](#)

COVID-19 Update

IMPORTANT SAFETY INFORMATION AND INDICATION FOR SKYRIZI™ (risankizumab-rzaa) ⁽⁴⁾

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

Important Safety Information

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.

site reactions, and tinea infections.

Please see [Full Prescribing Information](#).

US-SKZD-190350

REFERENCES

1. SKYRIZI [package insert]. North Chicago, IL: AbbVie Inc.
2. Gordon KB, Strober B, Lebwohl M, et al. Efficacy and safety of risankizumab in moderate-to-severe plaque psoriasis (UltIMMa-1 and UltIMMa-2): results from two double-blind, randomised, placebo-controlled and ustekinumab-controlled phase 3 trials. *Lancet*. 2018;392(10148):650-661.
3. Lebwohl M, Bachelez H, Valdecantos WC, Wu T, Gordon K. Efficacy and safety of risankizumab in moderate-to-severe plaque psoriasis: an integrated analysis of UltIMMa-1 and UltIMMa-2. Poster presented at: American Academy of Dermatology Annual Meeting; March 1-5, 2019; Washington, DC.

