

HIGHLIGHTS OF PRESCRIBING INFORMATION

These highlights do not include all the information needed to use ZORYVE cream safely and effectively. See full prescribing information for ZORYVE cream.

ZORYVE® (roflumilast) cream 0.3% and 0.15%, for topical use
Initial U.S. Approval: 2011

RECENT MAJOR CHANGES

Indications and Usage, Plaque Psoriasis (1.1) 10/2023
Indications and Usage, Atopic Dermatitis (1.2) 7/2024

INDICATIONS AND USAGE

ZORYVE cream is a phosphodiesterase 4 inhibitor:

- ZORYVE cream, 0.3%, is indicated for the topical treatment of plaque psoriasis, including intertriginous areas, in adult and pediatric patients 6 years of age and older. (1.1)
- ZORYVE cream, 0.15%, is indicated for the topical treatment of mild to moderate atopic dermatitis in adult and pediatric patients 6 years of age and older. (1.2)

DOSAGE AND ADMINISTRATION

- Apply ZORYVE cream: (2)
 - 0.3%, once daily to affected areas of plaque psoriasis.
 - 0.15%, once daily to affected areas of mild to moderate atopic dermatitis.
- For topical use only. (2)
- Not for ophthalmic, oral, or intravaginal use. (2)

DOSAGE FORMS AND STRENGTHS

- Cream, 0.3%: 3 mg of roflumilast per gram in 60-gram tubes. (3)
- Cream, 0.15%: 1.5 mg of roflumilast per gram in 60-gram tubes. (3)

CONTRAINDICATIONS

Moderate to severe liver impairment (Child-Pugh B or C). (4)

ADVERSE REACTIONS

The most common adverse reactions (reported in $\geq 1\%$ of subjects) are:

- **Plaque Psoriasis:** diarrhea, headache, insomnia, nausea, application site pain, upper respiratory tract infections, and urinary tract infections. (6.1)
- **Atopic Dermatitis:** headache, nausea, application site pain, diarrhea, and vomiting. (6.1)

To report SUSPECTED ADVERSE REACTIONS, contact Arcutis Biotherapeutics, Inc. at 1-844-692-6729 or FDA at 1-800-FDA-1088 or www.fda.gov/medwatch.

DRUG INTERACTIONS

- Co-administration of roflumilast with systemic CYP3A4 inhibitors or dual inhibitors that inhibit both CYP3A4 and CYP1A2 simultaneously may increase roflumilast systemic exposure and may result in increased adverse reactions. If these products are co-administered with ZORYVE cream, weigh the potential for increased adverse reactions against benefit. (7.1)
- Co-administration of roflumilast with oral contraceptives containing gestodene and ethinyl estradiol may increase roflumilast systemic exposure and may result in increased adverse reactions. If these products are co-administered with ZORYVE cream, weigh the potential for increased adverse reactions against benefit. (7.2)

See 17 for PATIENT COUNSELING INFORMATION and FDA-approved patient labeling.

Revised: 7/2024

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FULL PRESCRIBING INFORMATION

1 INDICATIONS AND USAGE

1.1 Plaque Psoriasis

ZORYVE[®] cream, 0.3%, is indicated for topical treatment of plaque psoriasis, including intertriginous areas, in adult and pediatric patients 6 years of age and older.

1.2 Atopic Dermatitis

ZORYVE cream, 0.15%, is indicated for topical treatment of mild to moderate atopic dermatitis in adult and pediatric patients 6 years of age and older.

2 DOSAGE AND ADMINISTRATION

Use ZORYVE cream, 0.3%, for the treatment of plaque psoriasis.

Use ZORYVE cream, 0.15%, for the treatment of mild to moderate atopic dermatitis.

Apply ZORYVE cream to affected areas once daily and rub in completely. Wash hands after application, unless ZORYVE cream is for treatment of the hands.

ZORYVE cream is for topical use only and not for ophthalmic, oral, or intravaginal use.

3 DOSAGE FORMS AND STRENGTHS

- Cream, 0.3%: 3 mg of roflumilast per gram of white to off-white cream in 60-gram tubes.
- Cream, 0.15%: 1.5 mg of roflumilast per gram of white to off-white cream in 60-gram tubes.

4 CONTRAINDICATIONS

ZORYVE cream is contraindicated in patients with moderate to severe liver impairment (Child-Pugh B or C) [see *Use in Specific Populations (8.6) and Clinical Pharmacology (12.3)*].

6 ADVERSE REACTIONS

6.1 Clinical Trials Experience

Because clinical trials are conducted under widely varying conditions, adverse reaction rates observed in the clinical trials of a drug cannot be directly compared to rates in the clinical trials of another drug and may not reflect the rates observed in practice.

Plaque Psoriasis

In two multicenter, randomized, double-blind, vehicle-controlled trials (DERMIS-1 and DERMIS-2), 881 adult and pediatric subjects 6 years of age or older with plaque psoriasis were treated with ZORYVE cream, 0.3%, or vehicle cream once daily for 8 weeks.

The median age was 47 years (range 6 to 88). The majority of the subjects were male (64%) and White (82%). The median body surface area (BSA) affected was 5.5% (range 2% to 20%). The proportion of subjects who discontinued treatment due to an adverse reaction was 1.0% for subjects treated with ZORYVE cream, 0.3%, and 1.3% for subjects treated with vehicle cream. The most common adverse reaction that led to discontinuation of ZORYVE cream, 0.3%, was application site urticaria (0.3%).

Table 1 presents adverse reactions that occurred in at least 1% of subjects treated with ZORYVE cream, 0.3%, and for which the rate exceeded the rate for vehicle cream.

Table 1: Adverse Reactions Reported in ≥1% of Subjects with Plaque Psoriasis Treated with ZORYVE Cream, 0.3%, (and More Frequently than Vehicle Cream) for 8 Weeks in Trials DERMIS-1 and DERMIS-2

Adverse Reaction	ZORYVE Cream, 0.3% (N=576) n (%)	Vehicle Cream (N=305) n (%)
Diarrhea	18 (3.1)	0 (0.0)
Headache	14 (2.4)	3 (1.0)
Insomnia	8 (1.4)	2 (0.7)
Nausea	7 (1.2)	1 (0.3)
Application site pain	6 (1.0)	1 (0.3)
Upper respiratory tract infection	6 (1.0)	1 (0.3)
Urinary tract infection	6 (1.0)	2 (0.7)

In 594 subjects with plaque psoriasis who continued treatment with ZORYVE cream, 0.3%, for up to 64 weeks in open-label extension trials, the adverse reaction profile was consistent with that observed in vehicle-controlled trials.

Atopic Dermatitis

In two multicenter, randomized, double-blind, vehicle-controlled trials (INTEGUMENT-1 and INTEGUMENT-2), 1336 adult and pediatric subjects 6 years of age or older with mild to moderate atopic dermatitis were treated with ZORYVE cream, 0.15%, or vehicle cream once daily for 4 weeks.

The median age was 20 years (range 6 to 91). The majority of the subjects were female (57%) and White (60%). The median body surface area (BSA) affected was 10% (range 3% to 88%).

The proportion of subjects who discontinued treatment due to an adverse reaction was 1.6% for subjects treated with ZORYVE cream, 0.15%, and 1.1% for subjects treated with vehicle cream.

Table 2 presents adverse reactions that occurred in at least 1% of subjects treated with ZORYVE cream, 0.15%, and for which the rate exceeded the rate for vehicle cream.

Table 2: Adverse Reactions Reported in ≥1% of Subjects with Atopic Dermatitis Treated with ZORYVE Cream, 0.15%, (and More Frequently than Vehicle Cream) for 4 Weeks in Trials INTEGUMENT-1 and INTEGUMENT-2

Adverse Reaction	ZORYVE Cream, 0.15% (N=885) n (%)	Vehicle Cream (N=451) n (%)
Headache	26 (2.9)	4 (0.9)
Nausea	17 (1.9)	2 (0.4)
Application site pain	13 (1.5)	3 (0.7)
Diarrhea	13 (1.5)	2 (0.4)
Vomiting	13 (1.5)	2 (0.4)

The adverse reaction of insomnia was reported in fewer than 1% of subjects treated with ZORYVE cream, 0.15%.

7 DRUG INTERACTIONS

7.1 Drugs that Inhibit Cytochrome P450 (CYP) Enzymes

The co-administration of roflumilast with systemic CYP3A4 inhibitors or dual inhibitors that inhibit both CYP3A4 and CYP1A2 simultaneously (e.g., erythromycin, ketoconazole, fluvoxamine, enoxacin, cimetidine) may increase roflumilast systemic exposure and may result in increased adverse reactions. If these products are co-administered with ZORYVE cream, weigh the potential for increased adverse reactions against benefit [see *Clinical Pharmacology* (12.3)].

7.2 Oral Contraceptives Containing Gestodene and Ethinyl Estradiol

The co-administration of roflumilast with oral contraceptives containing gestodene and ethinyl estradiol may increase roflumilast systemic exposure and may result in increased adverse reactions. If these products are co-administered with ZORYVE cream, weigh the potential for increased adverse reactions against benefit [see *Clinical Pharmacology* (12.3)].

8 USE IN SPECIFIC POPULATIONS

8.1 Pregnancy

Risk Summary

There are insufficient data available on the use of ZORYVE cream in pregnant women to inform a drug-associated risk of major birth defects, miscarriage, or other adverse maternal or fetal outcomes. In animal reproduction studies, roflumilast administered orally to pregnant rats and rabbits during the period of organogenesis produced no fetal structural abnormalities at doses up to 36 and 31 times the maximum recommended human dose (MRHD), respectively. Roflumilast induced post-implantation loss in rats at oral doses greater than or equal to 12 times the MRHD. Roflumilast induced stillbirth and decreased pup viability in mice at oral doses 19 and 59 times the MRHD, respectively. Roflumilast has been shown to adversely affect pup post-natal development when dams were treated with an oral dose 59 times the MRHD during pregnancy and lactation periods in mice (see *Data*).

The background risk of major birth defects and miscarriage for the indicated population is unknown. In the U.S. general population, the estimated background risk of major birth defects and miscarriage in clinically recognized pregnancies is 2% to 4% and 15% to 20%, respectively.

Clinical Considerations

Labor and delivery

Avoid using ZORYVE cream during labor and delivery. There are no human studies that have investigated effects of ZORYVE cream on preterm labor or labor at term; however, animal studies showed that oral roflumilast disrupted the labor and delivery process in mice.

Data

Animal data

In an embryo-fetal development study, pregnant rats were dosed orally during the period of organogenesis with up to 1.8 mg/kg/day roflumilast (36 times the MRHD on a mg/m² basis). No evidence of structural abnormalities or effects on survival rates were observed. Roflumilast did not affect embryo-fetal development at a maternal oral dose of 0.2 mg/kg/day (4 times the MRHD on a mg/m² basis).

In a fertility and embryo-fetal development study, male rats were dosed orally with up to 1.8 mg/kg/day roflumilast for 10 weeks and females for 2 weeks prior to pairing and throughout the organogenesis period. Roflumilast induced pre- and post-implantation loss at maternal oral

doses greater than or equal to 0.6 mg/kg/day (12 times the MRHD on a mg/m² basis). Roflumilast did not cause fetal structural abnormalities at maternal oral doses up to 1.8 mg/kg/day (35 times the MRHD on a mg/m² basis).

In an embryo-fetal development study in rabbits, pregnant does were dosed orally with 0.8 mg/kg/day roflumilast during the period of organogenesis. Roflumilast did not cause fetal structural abnormalities at the maternal oral doses of 0.8 mg/kg/day (31 times the MRHD on a mg/m² basis).

In pre- and post-natal developmental studies in mice, dams were dosed orally with up to 12 mg/kg/day roflumilast during the period of organogenesis and lactation. Roflumilast induced stillbirth and decreased pup viability at maternal oral doses greater than 2 mg/kg/day and 6 mg/kg/day, respectively (19 and 59 times the MRHD on a mg/m² basis, respectively). Roflumilast induced delivery retardation in pregnant mice at maternal oral doses greater than 2 mg/kg/day (19 times the MRHD on a mg/m² basis). Roflumilast decreased pup rearing frequencies at a maternal oral dose of 6 mg/kg/day during pregnancy and lactation (59 times the MRHD on a mg/m² basis). Roflumilast also decreased survival and forelimb grip reflex and delayed pinna detachment in mouse pups at a maternal oral dose of 12 mg/kg/day (116 times the MRHD on a mg/m² basis).

8.2 Lactation

Risk Summary

There are no data on the presence of roflumilast or its metabolite in human milk, the effects on the breastfed infant, or the effects on milk production.

Roflumilast and/or its metabolites are excreted into the milk of lactating rats (*see Data*). When a drug is present in animal milk, it is likely that the drug will be present in human milk. The developmental and health benefits of breastfeeding should be considered along with the mother's clinical need for ZORYVE cream and any potential adverse effects on the breastfed infant from ZORYVE cream or from the underlying maternal condition.

Clinical Considerations

To minimize potential exposure to the breastfed infant via breast milk, use ZORYVE cream on the smallest area of skin and for the shortest duration possible while breastfeeding. To avoid direct infant exposure, advise breastfeeding women not to apply ZORYVE cream directly to the nipple or areola. If applied to the patient's chest, avoid exposure via direct contact with the infant's skin.

Data

Animal data

Roflumilast and/or its metabolite concentrations measured 8 hours after an oral dose of 1 mg/kg given to lactating rats were 0.32 and 0.02 mcg/g in the milk and pup liver, respectively.

8.4 Pediatric Use

Plaque Psoriasis

The safety and effectiveness of ZORYVE cream, 0.3%, for the topical treatment of plaque psoriasis, including intertriginous areas, have been established in pediatric patients 6 years of age and older. Use of ZORYVE cream, 0.3%, in pediatric patients 6 years of age and older is supported by data from two 8-week, vehicle-controlled, safety and efficacy trials which included 18 subjects 6 to 17 years of age, of whom 11 received ZORYVE cream, 0.3%. Use of ZORYVE cream, 0.3%, in pediatric patients 12 years of age and older is also supported by data from

open-label trials of 2 and 24 weeks duration which included 18 subjects 12 to 17 years of age treated with ZORYVE cream, 0.3%. Use of ZORYVE cream, 0.3%, in pediatric patients 6 to less than 12 years of age is also supported by data from one 4-week, open-label, safety and pharmacokinetic (PK) study which included 20 pediatric subjects 6 to less than 12 years of age [see *Adverse Reactions (6.1)*, *Clinical Pharmacology (12.3)*, and *Clinical Studies (14.1)*].

The safety and effectiveness of ZORYVE cream, 0.3%, have not been established in pediatric patients younger than 6 years of age.

Atopic Dermatitis

The safety and effectiveness of ZORYVE cream, 0.15%, for the topical treatment of mild to moderate atopic dermatitis have been established in pediatric patients 6 years of age and older. Use of ZORYVE cream, 0.15%, in this age group is supported by data from two 4-week, vehicle-controlled, safety and efficacy trials which included 615 subjects 6 to 17 years of age, of whom 406 received ZORYVE cream, 0.15%. Use of ZORYVE cream, 0.15%, in pediatric patients 6 years of age and older is also supported by data from 481 pediatric subjects treated with ZORYVE cream, 0.15%, in open-label trials, of which 104 were treated for 52 weeks [see *Adverse Reactions (6.1)*, *Clinical Pharmacology (12.3)*, and *Clinical Studies (14.2)*].

The safety and effectiveness of ZORYVE cream, 0.15%, have not been established in pediatric patients younger than 6 years of age.

8.5 Geriatric Use

There were 184 patients 65 years of age and older in clinical studies for plaque psoriasis and atopic dermatitis [see *Clinical Studies (14.1, 14.2)*].

Plaque Psoriasis

Of the 576 patients treated with ZORYVE cream, 0.3%, in the 2 controlled clinical studies for plaque psoriasis, 56 (9.7%) were 65 to 74 years of age and 21 (3.7%) were 75 years of age and older. No overall differences in safety or effectiveness were observed between these subjects and younger subjects. Other reported clinical experience has not identified differences in responses between the geriatric and younger patients, but greater sensitivity of some older individuals cannot be ruled out.

Atopic Dermatitis

Of the 885 patients treated with ZORYVE cream, 0.15%, in the 2 controlled clinical studies for atopic dermatitis, 36 (4.1%) were 65 to 74 years of age and 8 (0.9%) were 75 years of age and older. No overall differences in safety or effectiveness were observed between these subjects and younger subjects. Other reported clinical experience has not identified differences in responses between the geriatric and younger patients, but greater sensitivity of some older individuals cannot be ruled out.

8.6 Hepatic Impairment

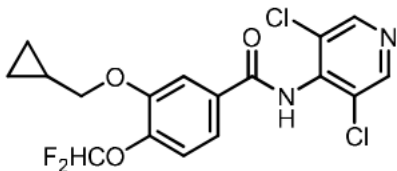
Oral roflumilast 250 mcg once daily for 14 days was studied in subjects with hepatic impairment. The systemic exposure of roflumilast and roflumilast N-oxide were increased in subjects with moderate (Child-Pugh B) hepatic impairment. ZORYVE cream is contraindicated in patients with moderate to severe liver impairment (Child-Pugh B or C). No dosage adjustment is needed in patients with mild (Child-Pugh A) hepatic impairment [see *Contraindications (4)* and *Clinical Pharmacology (12.3)*].

11 DESCRIPTION

ZORYVE (roflumilast) cream is a white to off-white cream for topical use. The active ingredient, roflumilast, is a phosphodiesterase 4 (PDE4) inhibitor.

Roflumilast is described chemically as 3-cyclopropylmethoxy-*N*-(3,5-dichloropyridin-4-yl)-4-(difluoromethoxy)benzamide. The empirical formula is C₁₇H₁₄Cl₂F₂N₂O₃, and the molecular weight is 403.21.

The structural formula is represented below:



Roflumilast is practically insoluble in water and hexane, sparingly soluble in ethanol, and freely soluble in acetone.

Each gram of cream, 0.15% or 0.3%, contains 1.5 mg or 3 mg of roflumilast, respectively, in a cream base containing cetareth-10 phosphate, cetaryl phosphate, cetostearyl alcohol, diethylene glycol monoethyl ether, hexylene glycol, isopropyl palmitate, methylparaben, propylparaben, purified water, sodium hydroxide, and white petrolatum. Hydrochloric acid may have been added to adjust pH.

12 CLINICAL PHARMACOLOGY

12.1 Mechanism of Action

Roflumilast and its active metabolite (roflumilast N-oxide) are inhibitors of PDE4. Roflumilast and roflumilast N-oxide inhibition of PDE4 (a major cyclic 3',5'-adenosine monophosphate [cyclic AMP] metabolizing enzyme) activity leads to accumulation of intracellular cyclic AMP. The specific mechanism(s) by which roflumilast exerts its therapeutic action is not well defined.

12.2 Pharmacodynamics

Pharmacodynamics of ZORYVE cream in the treatment of plaque psoriasis and atopic dermatitis are unknown.

12.3 Pharmacokinetics

Absorption

Following application of ZORYVE cream, the plasma concentration versus time profile was relatively flat, generally with a peak-to-trough ratio less than 2.

Plaque Psoriasis

The PK of ZORYVE cream, 0.3%, was investigated in 18 adult and 6 pediatric subjects 13 to 16 years of age with plaque psoriasis and a mean \pm SD body surface area (BSA) involvement of $26.8 \pm 6.80\%$ and $13.0 \pm 3.58\%$ in adult and pediatric subjects, respectively. In this study, on average, subjects applied 3 to 6.5 g of ZORYVE cream, 0.3%, once daily for 15 days. Plasma concentrations of roflumilast and roflumilast N-oxide (*see Metabolism*) were quantifiable in all but two subjects at Day 15.

In adults, the mean \pm SD systemic exposure (AUC_{0-24}) was 72.7 ± 53.1 and 628 ± 648 h·ng/mL for roflumilast and the N-oxide metabolite, respectively. In pediatric subjects, the mean \pm SD AUC_{0-24} was 25.1 ± 24.0 and 140 ± 179 h·ng/mL for roflumilast and the N-oxide metabolite, respectively.

The PK of ZORYVE cream, 0.3%, was investigated in 10 pediatric subjects (6 to less than 12 years of age) with at least mild plaque psoriasis and a mean \pm SD BSA involvement of $10.9 \pm 6.56\%$. The 14-day mean \pm SD extrapolated AUC_{0-24} was 75.6 ± 87.3 and 693 ± 986 h·ng/mL for roflumilast and the N-oxide metabolite, respectively. In another study, the PK of ZORYVE cream, 0.3%, was investigated in 9 pediatric subjects (2 to less than 6 years of age) with at least mild plaque psoriasis and a mean \pm SD BSA involvement of $9.44 \pm 5.57\%$. The 14-day mean \pm SD extrapolated AUC_{0-24} was 51.6 ± 29.9 and 539 ± 372 h·ng/mL for roflumilast and the N-oxide metabolite, respectively.

Atopic Dermatitis

The PK of ZORYVE cream, 0.15%, was investigated in 12 pediatric subjects 12 to 16 years of age and 13 pediatric subjects 6 to 11 years of age with atopic dermatitis and a mean \pm SD BSA involvement of $33.7 \pm 14.8\%$ and $43.5 \pm 10.5\%$, respectively. On average, subjects applied 8.2 to 10.5 g of ZORYVE cream, 0.15%, once daily.

In pediatric subjects 12 to 16 years of age, the Day 14 mean \pm SD systemic exposure (AUC_{0-24}) was 62.9 ± 53.0 and 336 ± 310 h·ng/mL for roflumilast and the N-oxide metabolite, respectively. In pediatric subjects 6 to 11 years of age, the Day 14 mean \pm SD AUC_{0-24} was 102 ± 96.5 and 743 ± 710 h·ng/mL for roflumilast and the N-oxide metabolite, respectively.

Distribution

Plasma protein binding of roflumilast and its N-oxide metabolite is approximately 99% and 97%, respectively.

Elimination

The plasma clearance after short-term intravenous infusion of roflumilast is on average about 9.6 L/h. Following topical administration, the half-lives of roflumilast and the N-oxide metabolite were 4.0 and 4.6 days, respectively.

Metabolism

Roflumilast is extensively metabolized via Phase I (cytochrome P450) and Phase II (conjugation) reactions. The N-oxide metabolite is the only major metabolite observed in the plasma of humans. Following oral administration, roflumilast and roflumilast N-oxide account for the majority (87.5%) of total dose administered in plasma. Roflumilast was not detectable in urine, while roflumilast N-oxide was only a trace metabolite (less than 1%). Other conjugated metabolites such as roflumilast N-oxide glucuronide and 4-amino-3,5-dichloropyridine N-oxide were detected in urine.

While roflumilast is 3 times more potent than roflumilast N-oxide at inhibition of the PDE4 enzyme *in vitro*, the plasma AUC of roflumilast N-oxide on average is approximately 8-fold greater than the plasma AUC of roflumilast following topical administration. A similar ratio was observed following intravenous administration, whereas following oral administration the N-oxide metabolite circulated on average about 10-fold higher than the parent drug.

Specific Populations

Following topical administration, no clinically significant differences in the pharmacokinetics of roflumilast and roflumilast N-oxide were observed based on age, sex, race, or ethnicity.

Patients with Hepatic Impairment

No studies were conducted with topical roflumilast in subjects with hepatic impairment; however, oral roflumilast 250 mcg once daily for 14 days was studied in subjects with mild to moderate hepatic impairment classified as Child-Pugh A and B (8 subjects in each group). The AUC of roflumilast and roflumilast N-oxide were increased by 51% and 24%, respectively, in Child-Pugh A subjects and by 92% and 41%, respectively, in Child-Pugh B subjects, as compared to age-, weight-, and gender-matched healthy subjects. The C_{max} of roflumilast and roflumilast N-oxide were increased by 3% and 26%, respectively, in Child-Pugh A subjects and by 26% and 40%, respectively, in Child-Pugh B subjects, as compared to healthy subjects [see *Contraindications (4)*].

Patients with Renal Impairment

No studies were conducted with topical roflumilast in subjects with renal impairment. Following oral administration in 12 subjects with severe renal impairment, no clinically significant differences in the pharmacokinetics of roflumilast and roflumilast N-oxide were observed.

Drug Interaction Studies

Clinical Studies

Since a major step in roflumilast metabolism is the N-oxidation of roflumilast to roflumilast N-oxide by CYP3A4 and CYP1A2, drug interaction studies were performed with oral roflumilast and systemic inhibitors of CYP3A4 and CYP1A2 [see *Drug Interactions (7.1)*].

Erythromycin: In an open-label crossover study in 16 healthy volunteers, the co-administration of CYP3A4 inhibitor erythromycin (500 mg 3 times daily for 13 days) with a single oral dose of 500 mcg roflumilast resulted in 40% and 70% increase in C_{max} and AUC for roflumilast, respectively, and a 34% decrease and a 4% increase in C_{max} and AUC for roflumilast N-oxide, respectively.

Ketoconazole: In an open-label crossover study in 16 healthy volunteers, the co-administration of a strong CYP3A4 inhibitor ketoconazole (200 mg twice daily for 13 days) with a single oral dose of 500 mcg roflumilast resulted in 23% and 99% increase in C_{max} and AUC for roflumilast, respectively, and a 38% reduction and 3% increase in C_{max} and AUC for roflumilast N-oxide, respectively.

Fluvoxamine: In an open-label crossover study in 16 healthy volunteers, the co-administration of dual CYP 3A4/1A2 inhibitor fluvoxamine (50 mg daily for 14 days) with a single oral dose of 500 mcg roflumilast showed a 12% and 156% increase in roflumilast C_{max} and AUC along with a 210% decrease and 52% increase in roflumilast N-oxide C_{max} and AUC, respectively.

Enoxacin: In an open-label crossover study in 16 healthy volunteers, the co-administration of dual CYP 3A4/1A2 inhibitor enoxacin (400 mg twice daily for 12 days) with a single oral dose of 500 mcg roflumilast resulted in an increased C_{max} and AUC of roflumilast by 20% and 56%, respectively. Roflumilast N-oxide C_{max} was decreased by 14% while roflumilast N-oxide AUC was increased by 23%.

Cimetidine: In an open-label crossover study in 16 healthy volunteers, the co-administration of a dual CYP 3A4/1A2 inhibitor cimetidine (400 mg twice daily for 7 days) with a single oral dose of 500 mcg roflumilast resulted in a 46% and 85% increase in roflumilast C_{max} and AUC; and a 4% decrease in C_{max} and 27% increase in AUC for roflumilast N-oxide, respectively.

Oral Contraceptives Containing Gestodene and Ethinyl Estradiol: In an open-label crossover study in 20 healthy adult volunteers, co-administration of a single oral dose of roflumilast with repeated doses of a fixed combination oral contraceptive containing 0.075 mg gestodene and 0.03 mg ethinyl estradiol to steady state caused a 38% increase and 12% decrease in C_{max} of

roflumilast and roflumilast N-oxide, respectively. Roflumilast and roflumilast N-oxide AUCs were increased by 51% and 14%, respectively.

In Vitro Studies

Cytochrome P450 (CYP) Enzymes: *In vitro* studies suggest that the biotransformation of roflumilast to its N-oxide metabolite is mediated by CYP1A2 and 3A4. Based on further *in vitro* results in human liver microsomes, roflumilast and roflumilast N-oxide in therapeutic plasma concentrations do not inhibit CYP1A2, 2A6, 2B6, 2C8, 2C9, 2C19, 2D6, 2E1, 3A4/5, or 4A9/11; therefore, there is a low probability of relevant interactions with substances metabolized by these P450 enzymes. In addition, *in vitro* studies demonstrated no induction of the CYP1A2, 2A6, 2C9, 2C19, or 3A4/5 and only a weak induction of CYP2B6 by roflumilast.

13 NONCLINICAL TOXICOLOGY

13.1 Carcinogenesis, Mutagenesis, Impairment of Fertility

Long-term studies were conducted in hamsters and mice with roflumilast to evaluate its carcinogenic potential. In 2-year oral gavage carcinogenicity studies, roflumilast treatment resulted in dose-related, statistically significant increases in the incidence of undifferentiated carcinomas of nasal epithelium in hamsters at doses greater than or equal to 8 mg/kg/day (12 times the MRHD on an AUC basis). The tumorigenicity of roflumilast appears to be attributed to a reactive metabolite of 4-amino-3,5-dichloropyridine N-oxide (ADCP N-oxide). No evidence of tumorigenicity was observed in mice at roflumilast oral doses up to 12 and 18 mg/kg/day in females and males, respectively (11 and 17 times the MRHD, respectively, on an AUC basis).

In a 2-year dermal mouse carcinogenicity study, no evidence of carcinogenicity was observed at topical doses of roflumilast cream up to 1% applied at 2 mL/kg/day (4 times the MRHD on an AUC basis).

Roflumilast tested positive in an *in vivo* mouse micronucleus test, but negative in the following assays: the Ames test, an *in vitro* chromosome aberration assay in human lymphocytes, an *in vitro* HPRT assay with V79 cells, an *in vitro* micronucleus test with V79 cells, a DNA adduct formation assay in rat nasal mucosa, liver, and testes, and an *in vivo* mouse bone marrow chromosome aberration assay. Roflumilast N-oxide was negative in the Ames test and an *in vitro* micronucleus test with V79 cells.

In a human spermatogenesis study, oral roflumilast 500 mcg had no effects on semen parameters or reproductive hormones during the 3-month treatment period and the following 3-month off-treatment period. In a fertility study, oral roflumilast decreased fertility rates in male rats at 1.8 mg/kg/day (35 times the MRHD on a mg/m² basis). The male rats also showed increases in the incidence of tubular atrophy, degeneration in the testis, and spermiogenic granuloma in the epididymides. No effect on rat fertility rate or male reproductive organ morphology was observed at 0.6 mg/kg/day (12 times the MRHD on a mg/m² basis). In a female fertility study, no effect on fertility was observed up to the highest roflumilast dose of 1.5 mg/kg/day in rats (29 times the MRHD on a mg/m² basis).

14 CLINICAL STUDIES

14.1 Plaque Psoriasis

Two multicenter, randomized, double-blind, vehicle-controlled trials (DERMIS-1 [NCT04211363] and DERMIS-2 [NCT04211389]) enrolled a total of 881 adult and pediatric subjects 6 years of age or older with mild to severe plaque psoriasis. Subjects were randomized 2:1 to receive ZORYVE cream, 0.3%, or vehicle cream applied topically once daily for 8 weeks. The median age of the trial population was 47 years (range 6 to 88 years of age). The trial population was 64% male and 36% female; 82.3% were White, 7.1% Asian, 3.6% Black or African American,

0.9% Native Hawaiian or Other Pacific Islander, 0.7% American Indian or Alaska Native, 0.5% more than one race, 3.3% Other, and missing in 1.6% of subjects. For ethnicity, 75.7% of subjects identified as Not Hispanic or Latino, 24.1% identified as Hispanic or Latino, and 0.2% were unknown. The median body surface area (BSA) affected of the trial population was 5.5% (range 2% to 20%). At baseline, 16% of subjects had an Investigator’s Global Assessment (IGA) score of 2 (mild), 76% had an IGA score of 3 (moderate), and 8% had an IGA score of 4 (severe). One hundred seventy-nine (20%) subjects had a baseline intertriginous IGA (I-IGA) score of 2 or higher (mild), and 678 (77%) subjects had a baseline Worst Itch-Numeric Rating Scale (WI-NRS) score of 4 or higher on a scale of 0 to 10.

The primary endpoint was the proportion of subjects who achieved IGA treatment success at Week 8 (Table 3). Success was defined as a score of “Clear” (0) or “Almost Clear” (1), plus a 2-grade improvement from baseline.

Table 3: IGA Treatment Success at Week 8 in Subjects with Mild to Severe Plaque Psoriasis in Trials DERMIS-1 and DERMIS-2

	DERMIS-1		DERMIS-2	
	ZORYVE Cream, 0.3%	Vehicle Cream	ZORYVE Cream, 0.3%	Vehicle Cream
Subjects randomized	N=286	N=153	N=290	N=152
IGA success*	41.5%	5.8%	36.7%	7.1%
Difference from vehicle (95% CI)†	39.7% (32.4%, 47.0%)		29.5% (21.5%, 37.6%)	

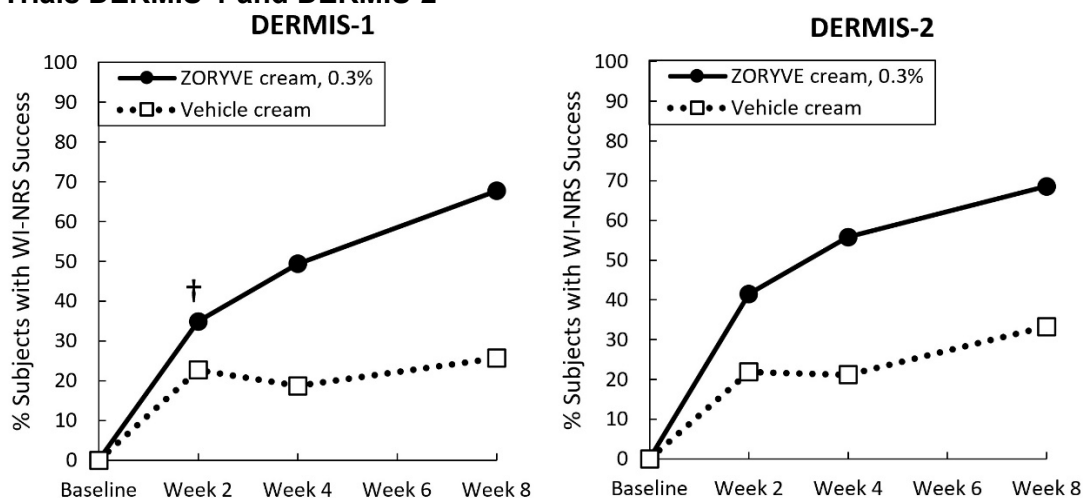
Abbreviations: CI = Confidence Interval

*IGA treatment success was defined as an IGA score of “Clear” (0) or “Almost Clear” (1), plus a 2-grade IGA score improvement from baseline at Week 8 (Multiple Imputation).

†Treatment difference and 95% CI are based on the CMH method stratified by site, baseline IGA, and baseline intertriginous involvement.

Secondary endpoints included the proportion of subjects that achieved I-IGA success at Week 8 and WI-NRS success sequentially at Weeks 8, 4, and 2 (see Figure 1). WI-NRS success was defined as a reduction of at least 4 points from baseline in subjects with a baseline WI-NRS score of at least 4.

Figure 1: WI-NRS Success Over Time in Subjects with Mild to Severe Plaque Psoriasis in Trials DERMIS-1 and DERMIS-2*



*WI-NRS success is a reduction of at least 4 points in subjects with a WI-NRS score of 4 or higher at baseline.

†The treatment difference at Week 2 in DERMIS-1 was not statistically significant.

Among subjects with an I-IGA score of at least 2 (mild) at baseline (approximately 22% of subjects in DERMIS-1 and 19% in DERMIS-2), there was a higher percentage of subjects who achieved I-IGA success at Week 8 in the group who received ZORYVE cream, 0.3%, compared to the group who received vehicle cream (DERMIS-1: 71.5% vs. 13.8%; DERMIS-2: 67.5% vs. 17.4%).

14.2 Atopic Dermatitis

Two multicenter, randomized, double-blind, vehicle-controlled trials (INTEGUMENT-1 [NCT04773587] and INTEGUMENT-2 [NCT04773600]) enrolled a total of 1337 adult and pediatric subjects 6 years of age and older (615 subjects were 6 to 17 years of age) with mild to moderate atopic dermatitis. Subjects were randomized 2:1 to receive ZORYVE cream, 0.15%, or vehicle cream applied once daily for 4 weeks. The median age of the trial population was 20 years (range 6 to 91 years of age). The trial population was 57% female; 59.5% were White, 20.3% Black or African American, 13.2% Asian, 3.4% Other, 2.8% more than one race, 0.6% American Indian or Alaska Native, and 0.1% Native Hawaiian or Other Pacific Islander. For ethnicity, 82.8% of subjects identified as Not Hispanic or Latino, 16.6% identified as Hispanic or Latino, and 0.6% were unknown. At baseline, subjects had a mean affected BSA of 14% (range of 3% to 88%), and 42% had facial involvement. At baseline, 24% of subjects had a validated Investigator Global Assessment for Atopic Dermatitis (vIGA-AD) score of 2 (mild), and 76% had a vIGA-AD score of 3 (moderate). Eight hundred thirteen (60.8%) subjects 12 years of age and older had a baseline WI-NRS score of 4 or higher on a scale of 0 to 10.

The primary endpoint was the proportion of subjects who achieved vIGA-AD treatment success at Week 4 (Table 4). Success was defined as a score of “Clear” (0) or “Almost Clear” (1), plus a 2-grade improvement from baseline.

Secondary endpoints included the proportion of subjects that achieved vIGA-AD success at Weeks 2 and 1 (Figure 2) and WI-NRS success at Weeks 4, 2, and 1 (Figure 3). WI-NRS success was defined as a reduction of at least 4 points from baseline in subjects 12 years of age or older with a baseline WI-NRS score of at least 4.

Table 4: Efficacy Results at Week 4 in Subjects with Mild to Moderate Atopic Dermatitis in Trials INTEGUMENT-1 and INTEGUMENT-2

	INTEGUMENT-1		INTEGUMENT-2	
	ZORYVE Cream, 0.15%	Vehicle Cream	ZORYVE Cream, 0.15%	Vehicle Cream
Subjects randomized	N=433	N=221	N=451	N=232
vIGA-AD success*	32.0%	15.2%	28.9%	12.0%
Difference from vehicle (95% CI)†	17.4% (11.09%, 23.75%)		16.5% (10.61%, 22.42%)	
Subjects ≥12 years randomized with baseline WI-NRS ≥4	N=278	N=135	N=264	N=136
WI-NRS success‡	33.6%	20.7%	30.2%	12.4%
Difference from vehicle (95% CI)†	13.3% (4.01%, 22.60%)		15.0% (6.52%, 23.54%)	

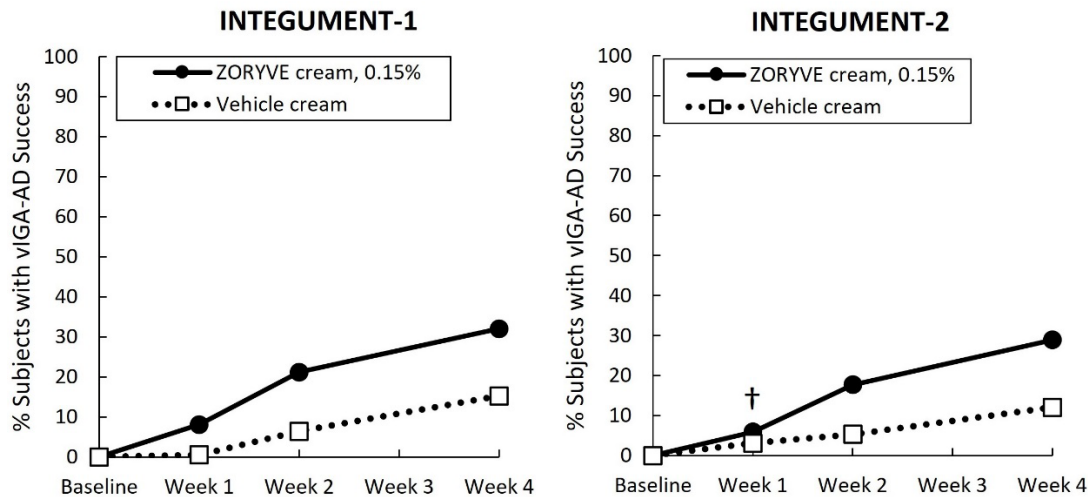
Abbreviations: CI = Confidence Interval

*vIGA-AD success was defined as a vIGA-AD score of “Clear” (0) or “Almost Clear” (1), plus a 2-grade vIGA-AD score improvement from baseline at Week 4 (Multiple Imputation).

†The difference in percent and related 95% CIs are Mantel-Haenszel estimates stratified by pooled study site and vIGA-AD randomization strata.

‡WI-NRS success was defined as a reduction of at least 4 points from baseline for subjects 12 years of age or older with a baseline score of at least 4.

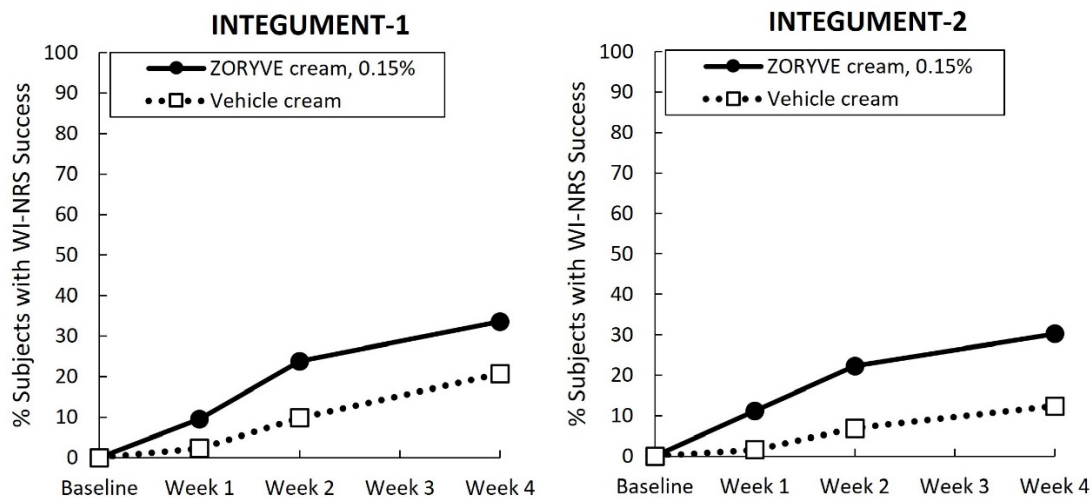
Figure 2: vIGA-AD Success Over Time in Subjects with Mild to Moderate Atopic Dermatitis in Trials INTEGUMENT-1 and INTEGUMENT-2*



*vIGA-AD success was defined as a vIGA-AD score of “Clear” (0) or “Almost Clear” (1), plus a 2-grade vIGA-AD score improvement from baseline at Week 4 (Multiple Imputation).

†The treatment difference at Week 1 in INTEGUMENT-2 was not statistically significant.

Figure 3: WI-NRS Success Over Time in Subjects with Mild to Moderate Atopic Dermatitis in Trials INTEGUMENT-1 and INTEGUMENT-2*



*WI-NRS success in subjects 12 years of age or older is a reduction of at least 4 points in subjects with a WI-NRS score of 4 or higher at baseline.

16 HOW SUPPLIED/STORAGE AND HANDLING

How Supplied

ZORYVE (roflumilast) cream is a white to off-white cream supplied as follows:

- Cream, 0.3%: 3 mg of roflumilast per gram supplied in 60-gram aluminum tubes (NDC 80610-130-60).
- Cream, 0.15%: 1.5 mg of roflumilast per gram supplied in 60-gram aluminum tubes (NDC 80610-115-60).

Storage and Handling

Store at 20°C to 25°C (68°F to 77°F); excursions permitted between 15°C and 30°C (59°F and 86°F). [See USP Controlled Room Temperature]

17 PATIENT COUNSELING INFORMATION

Advise the patient or caregiver to read the FDA-approved patient labeling (Patient Information).

Lactation

Advise patients to use ZORYVE cream on the smallest area of skin and for the shortest duration possible while breastfeeding. Instruct patients who are breastfeeding not to apply ZORYVE cream directly to the nipple or areola to avoid direct infant exposure. Instruct patients to avoid inadvertent contact of treated areas with infant skin [see *Use in Specific Populations (8.2)*].

Marketed by: Arcutis Biotherapeutics, Inc.

Westlake Village, CA 91361

For more information, call 1-805-418-5006 or visit <http://www.zoryve.com>.

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Patient Information
ZORYVE® (zor-EEV)
(roflumilast) cream

Important information: ZORYVE cream is for use on the skin (topical use) only. Do not use ZORYVE cream in or on your eyes, mouth, or vagina.

What is ZORYVE cream?

- ZORYVE cream 0.3% is a prescription medicine used on the skin (topical), including in areas with skin folds, to treat plaque psoriasis in people 6 years of age and older. It is not known if ZORYVE cream 0.3% is safe and effective in children with plaque psoriasis under 6 years of age.
- ZORYVE cream 0.15% is a prescription medicine used on the skin (topical) to treat mild to moderate eczema (atopic dermatitis) in people 6 years of age and older. It is not known if ZORYVE cream 0.15% is safe and effective in children with mild to moderate eczema (atopic dermatitis) under 6 years of age.

Do not use ZORYVE cream if you have certain liver problems.

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

- have liver problems.
- are pregnant or plan to become pregnant. It is not known if ZORYVE cream will harm your unborn baby.
- are breastfeeding or plan to breastfeed. It is not known if ZORYVE cream passes into your breast milk. Talk to your healthcare provider about the best way to feed your baby during treatment with ZORYVE cream. Breastfeeding women using ZORYVE cream should use it on the smallest area of the skin and for the shortest time needed. Do not apply ZORYVE cream directly to the nipple or areola to avoid contact with your baby.

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

How should I use ZORYVE cream?

- Use ZORYVE cream exactly as your healthcare provider tells you to use it.
- Apply ZORYVE cream to the affected areas 1 time a day. Rub the cream in completely until you no longer see it on your skin.
- Wash your hands after applying ZORYVE cream, unless your hands are being treated. If someone else applies ZORYVE cream for you, they should wash their hands after applying it.

What are the possible side effects of ZORYVE cream?

The most common side effects of ZORYVE cream 0.3% in people treated for plaque psoriasis include:

- diarrhea
- headache
- trouble sleeping
- nausea
- application site pain
- upper respiratory tract infections
- urinary tract infections (UTIs)

The most common side effects of ZORYVE cream 0.15% in people treated for atopic dermatitis include:

- headache
- nausea
- application site pain
- diarrhea
- vomiting

These are not all of the possible side effects of ZORYVE cream.

Call your doctor for medical advice about side effects. You may report side effects to FDA at 1-800-FDA-1088.

You may also report side effects to Arcutis Biotherapeutics, Inc. by calling 1-844-692-6729.

How should I store ZORYVE cream?

- Store ZORYVE cream at room temperature from 68°F to 77°F (20°C to 25°C).

Keep ZORYVE cream and all medicines out of the reach of children.

General information about the safe and effective use of ZORYVE cream.

Medicines are sometimes prescribed for purposes other than those listed in a Patient Information leaflet. Do not use ZORYVE cream for a condition for which it was not prescribed. Do not give ZORYVE cream to other people, even if they have the same symptoms you have. It may harm them. You can ask your healthcare provider or pharmacist for information about ZORYVE cream that is written for health professionals.

What are the ingredients in ZORYVE cream?

Active ingredient: roflumilast

Inactive ingredients: cetareth-10 phosphate, cetearyl phosphate, cetostearyl alcohol, diethylene glycol monoethyl ether, hexylene glycol, isopropyl palmitate, methylparaben, propylparaben, purified water, sodium hydroxide, and white petrolatum. Hydrochloric acid may have been added to adjust pH.

Marketed by: Arcutis Biotherapeutics, Inc.

Westlake Village, CA 91361

For more information, call 1-844-692-6729 or visit <http://www.zoryve.com>.

This Patient Information has been approved by the U.S. Food and Drug Administration.

Revised 7/2024

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