

SULFACETAMIDE SODIUM- sulfacetamide sodium lotion **Padagis Israel Pharmaceuticals Ltd**

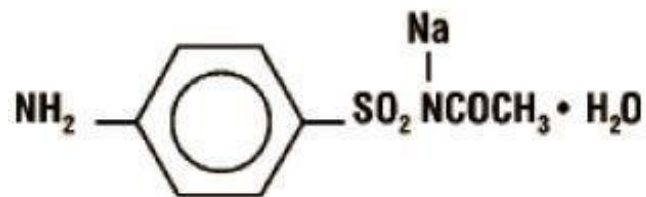
Sulfacetamide Sodium Topical Suspension USP, 10% (Lotion)

Rx Only

DESCRIPTION

Each mL of Sulfacetamide Sodium Topical Suspension USP, 10% (Lotion) contains 100 mg of sulfacetamide sodium in a vehicle consisting of diethanolamine; EDTA; hydroxyethyl cellulose; lauramide DEA; methylparaben; polyethylene glycol 400, monolaurate; propylene glycol; purified water; simethicone; sodium chloride; sodium metabisulfite and xanthan gum.

Sulfacetamide sodium is a sulfonamide with antibacterial activity. Chemically, sulfacetamide sodium is N'-[(4-aminophenyl) sulfonyl]-acetamide, monosodium salt, monohydrate. The structural formula is:



CLINICAL PHARMACOLOGY

The most widely accepted mechanism of action of sulfonamides is the Woods-Fildes theory, based on sulfonamides acting as a competitive inhibitor of paraaminobenzoic acid (PABA) utilization, an essential component for bacterial growth. While absorption through intact skin in humans has not been determined, *in vitro* studies with human cadaver skin indicated a percutaneous absorption of about 4%. Sulfacetamide sodium is readily absorbed from the gastrointestinal tract when taken orally and excreted in the urine largely unchanged. The biological half-life has been reported to be between 7 to 13 hours.

The pharmacokinetics of sulfacetamide and its major metabolite sulfanilamide in sulfacetamide sodium lotion was evaluated in adult subjects (N=14) with acne vulgaris. The subjects applied sulfacetamide sodium lotion to their face, back, chest and shoulders every 12 hours for 28 days. The percentage of the applied dose of sulfacetamide sodium lotion excreted in the urine as sulfacetamide plus sulfanilamide ranged from 0.08 to 0.33%.

INDICATIONS

Sulfacetamide Sodium Topical Suspension USP, 10% (Lotion) is indicated in the topical

treatment of *acne vulgaris*.

CONTRAINDICATIONS

Sulfacetamide Sodium Topical Suspension USP, 10% (Lotion) is contraindicated for use by patients having known hypersensitivity to sulfonamides or any other component of this preparation (see **WARNINGS** section).

WARNINGS

Fatalities have occurred, although rarely, due to severe reactions to sulfonamides including Stevens-Johnson syndrome, toxic epidermal necrolysis, fulminant hepatic necrosis, agranulocytosis, aplastic anemia, and other blood dyscrasias. Hypersensitivity reactions may occur when a sulfonamide is readministered, irrespective of the route of administration. Sensitivity reactions have been reported in individuals with no prior history of sulfonamide hypersensitivity. At the first sign of hypersensitivity, skin rash or other reactions, discontinue use of this preparation (see **ADVERSE REACTIONS** section).

Sulfacetamide Sodium Topical Suspension USP, 10% (Lotion) contains sodium metabisulfite, a sulfite that may cause allergic-type reactions including anaphylactic symptoms and life-threatening or less severe asthmatic episodes in certain susceptible people. The overall prevalence of sulfite sensitivity in the general population is unknown and probably low. Sulfite sensitivity is seen more frequently in asthmatic than in non-asthmatic people (see **CONTRAINDICATIONS** section).

PRECAUTIONS

General -

For external use only. Keep away from eyes. If irritation develops, use of the product should be discontinued and appropriate therapy instituted. Patients should be carefully observed for possible local irritation or sensitization during long-term therapy. Hypersensitivity reactions may occur when a sulfonamide is readministered irrespective of the route of administration, and cross-sensitivity between different sulfonamides may occur. Sulfacetamide sodium can cause reddening and scaling of the skin. Particular caution should be employed if areas of involved skin to be treated are denuded or abraded.

Keep out of the reach of children.

Carcinogenesis, Mutagenesis, Impairment of Fertility -

Long-term studies in animals have not been performed to evaluate carcinogenic potential.

Pregnancy:

Teratogenic Effects:

Pregnancy Category C -

Animal reproduction studies have not been conducted with Sulfacetamide Sodium Topical Suspension USP, 10% (Lotion). It is also not known whether Sulfacetamide Sodium Topical Suspension USP, 10% (Lotion) can cause fetal harm when administered to a pregnant woman or can affect reproduction capacity. Sulfacetamide Sodium Topical Suspension USP, 10% (Lotion) should be given to a pregnant woman only if clearly needed.

Kernicterus may occur in the newborn as a result of treatment of a pregnant woman at term with orally administered sulfonamide. There are no adequate and well controlled studies of Sulfacetamide Sodium Topical Suspension USP, 10% (Lotion) in pregnant women, and it is not known whether topically applied sulfonamides can cause fetal harm when administered to a pregnant woman.

Nursing Mothers -

It is not known whether sulfacetamide sodium is excreted in the human milk following topical use of Sulfacetamide Sodium Topical Suspension USP, 10% (Lotion). Systemically administered sulfonamides are capable of producing kernicterus in the infants of lactating women. Small amounts of orally administered sulfonamides have been reported to be eliminated in human milk. Because many drugs are excreted in human milk, caution should be exercised in prescribing for nursing women.

Pediatric Use -

Safety and effectiveness in pediatric patients under the age of 12 have not been established.

ADVERSE REACTIONS

In controlled clinical trials for the management of acne vulgaris, the occurrence of adverse reactions associated with the use of sulfacetamide sodium lotion was infrequent and restricted to local events. The total incidence of adverse reactions reported in these studies was less than 2%. Only one of 105 patients treated with sulfacetamide sodium lotion had adverse reactions of erythema, itching and edema. It has been reported that sulfacetamide sodium may cause local irritation, stinging and burning. While the irritation may be transient, occasionally, the use of medication has to be discontinued.

DOSAGE AND ADMINISTRATION

Apply a thin film to affected areas twice daily.

HOW SUPPLIED

Sulfacetamide Sodium Topical Suspension USP, 10% (Lotion) is available as follows:
4 FL OZ (118mL) bottles (NDC 45802-**896**-26)

STORAGE

Store at 20-25°C (68-77°F) [see USP Controlled Room Temperature]. Shake well before using. Keep tightly closed.

Manufactured by Perrigo

Bronx, NY 10457

Distributed By

Perrigo[®]

Allegan, MI 49010 • www.perrigo.com

Rev. 12/12

:2C800 RC J4

PRINCIPAL DISPLAY PANEL

Rx Only

Sulfacetamide Sodium Topical Suspension USP, 10% (Lotion)

For Topical Use Only

Shake Well Before Using

4 FL OZ (118 mL)



The following image is a placeholder representing the product identifier that is either affixed or imprinted on the drug package label during the packaging operation.

S/N [insert product's serial number]
 Lot [insert product's lot number]
 Exp [insert product's expiration date]

SULFACETAMIDE SODIUM

sulfacetamide sodium lotion

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:45802-896
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SULFACETAMIDE SODIUM (UNII: 4NRT660KJQ) (SULFACETAMIDE - UNII:4965G3J0F5)	SULFACETAMIDE SODIUM	100 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
DIETHANOLAMINE (UNII: AZE05TDV2V)	
EDETIC ACID (UNII: 9G34HU7RV0)	
LAURIC DIETHANOLAMIDE (UNII: I29I2VHG38)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM METABISULFITE (UNII: 4VON5FNS3C)	
XANTHAN GUM (UNII: TTV12P4NEE)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:45802-896-26	1 in 1 CARTON	10/17/2012	
1		118 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA078649	10/17/2012	

Labeler - Padagis Israel Pharmaceuticals Ltd (600093611)

Revised: 10/2021

Padagis Israel Pharmaceuticals Ltd