

Sodium Sulfacetamide-Sulfur Therapeutic Cheat Sheet

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TRADE NAME

- Plexion, Clenia, Rosanil, Rosac, Sulfacleanse

MECHANISM OF ACTION^{1,2}

- Sodium sulfacetamide is a sulfonamide antimicrobial agent that competitively inhibits para-aminobenzoic acid utilization, impairing bacterial folate synthesis and thereby limiting proliferation of susceptible cutaneous organisms, including *Cutibacterium acnes*. In addition to its antibacterial activity, sulfacetamide exhibits anti-inflammatory effects that contribute to clinical improvement in inflammatory acne and papulopustular rosacea. Sulfur provides complementary keratolytic and desquamative activity through interactions with keratin and sulfur-containing amino acids, facilitating stratum corneum shedding and reduction of follicular obstruction. The combination of sulfacetamide and sulfur has demonstrated efficacy in reducing inflammatory lesion counts and background erythema in rosacea, with utility in patients with overlapping seborrheic dermatitis phenotypes.

FDA-APPROVED INDICATIONS

- Topical sulfur–sulfacetamide formulations are FDA approved for the treatment of acne vulgaris, rosacea (acne rosacea), and seborrheic dermatitis.

OFF-LABEL DERMATOLOGIC USES^{1,2,3,4}

- Perioral (periorificial) dermatitis, particularly in mild or chronic cases where topical antibiotics or steroids are being avoided
- Pityriasis (tinea) versicolor, as a nonstandard option in mild cases when conventional antifungals are not tolerated or are ineffective
- Other inflammatory dermatoses with scale, where sulfur’s keratolytic properties are leveraged based on clinical experience rather than controlled trials.

DOSING^{1,3}

- Leave-on formulations containing sulfacetamide 10% and sulfur 5% are typically applied once to three times daily, with many clinicians initiating once-daily use and titrating upward based on tolerability.
- Cleanser formulations are generally used once or twice daily, with avoidance of contact with the eyes, lips, and mucous membranes.

WARNINGS AND PRECAUTIONS^{1,5}

- Sodium sulfacetamide carries class sulfonamide warnings, including rare but potentially serious hypersensitivity reactions such as agranulocytosis, hemolytic anemia, drug fever, and jaundice.
- Although systemic absorption is minimal with topical use, caution is advised in patients with known sulfonamide hypersensitivity, particularly when applied to inflamed or compromised skin.
- Local irritation, dryness, and erythema are common early in therapy and may necessitate dose reduction or discontinuation.

SIDE EFFECTS^{1,2}

- Local cutaneous reactions including burning, stinging, erythema, scaling, and peeling are most common.
- Allergic contact dermatitis and hypersensitivity reactions are uncommon but reported.
- Severe systemic reactions are exceedingly rare but included in labeling due to sulfonamide class effects.

DRUG INTERACTIONS^{3,10}

- Clinically significant systemic drug interactions are unlikely with topical therapy.
- Additive irritation may occur with concurrent use of retinoids, benzoyl peroxide, or alcohol-based topical agents.

CONTRAINDICATIONS^{1,2}

- Known hypersensitivity to sulfonamides, sulfur, or any formulation component.
- Some product labels advise avoidance in patients with underlying renal disease.

PREGNANCY AND BREASTFEEDING

- Pregnancy category C under legacy FDA labeling; use only if potential benefit justifies potential risk.
- Data on topical use during lactation are limited; caution is advised due to known oral sulfonamide excretion in breast milk.

MONITORING

- Routine laboratory monitoring is not required.
- Clinical monitoring should focus on cutaneous tolerability and signs of hypersensitivity.
- Treatment response and adverse effects are typically reassessed after 6–12 weeks, consistent with clinical rosacea studies.

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