



SUGGESTED FORMULA

Hydroquinone 8% / Retinoic Acid 0.1% Topical Cream

Version number: 1.0

Volume or quantity: 100 gm

Hydroquinone, USP (HY113)	8gm
Retinoic Acid (Tretinoin, Powder, USP)* (R1022)	0.1 gm
Butylated Hydroxytoluene, NF (B1196)	0.4gm
Polysorbate 80, NF (PO138)	3mL
Propylene Glycol, USP (PR130)	2.5mL
Sodium Metabisulfite, NF (SO182)	0.3gm
Alcohol 190Proof, USP (ET108)	XmL
Lipocream (B4520)	QS 100gm

SUGGESTED COMPOUNDING PROCEDURES

1. Calculate the required quantity for each ingredient
2. Weigh and/or measure all ingredients
3. Transfer Butylated Hydroxytoluene to mortar and triturate crystals to fine powder
4. Add Hydroquinone to mortar and triturate to fine powder
5. Add enough Propylene Glycol to form paste, then add remaining Propylene Glycol
6. Add Polysorbate 80 and mix well
7. Using geometric dilution add Lipocream and mix well
8. Process step 6 through ointment mill and/or electronic mortar and pestle to reduce grit
9. Dissolve Retinoic Acid in a small amount of Ethyl Alcohol and mix into cream by geometric dilution until homogenous
10. Package and label
11. Suggested Quality assessments:
 - a. color
 - b. texture
 - c. container
 - d. Label - auxiliary labels, storage, BUD, compounded medication

Store in light resistant air-tight container, at Controlled room temperature

No claims are made as to the safety or efficacy of this preparation. This formulation is provided solely at the unsolicited request of the pharmacist.

Beyond-Use Dates of preparations are conservative estimates from reference books, peer-reviewed literature, and intended duration of therapy, formulation from commercially available products, organoleptic observations and current USP guidelines. Compounders may have stability studies performed by a reputable laboratory if they wish to extend the Beyond-Use Date. It is recommended that you follow USP <795> recommendations for potency testing.

Beyond-Use Date should be assigned as directed by the current guidelines of USP Chapter <795>

***Precautions:**

Hazardous drugs should be handled according to USP and NIOSH standards and pharmacy SOPs. This preparation should be compounded in a vertical airflow hood, in a biological safety cabinet or barrier isolation technology. Precautions should be taken to prevent cross-contamination and exposure of ingredients to the compounder and contamination of the preparation by the compounder. Wear appropriate personal protective equipment.

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