



SUGGESTED FORMULA

Hydroquinone 8% / Kojic Acid 1% / Retinoic Acid 0.1% / Fluocinolone 0.025%/ Niacinamide 4% Topical Solution

Version number: 1.0

Volume or quantity: 100 gm

Hydroquinone, USP (HY113)	8gm
Kojic Acid	1gm
Retinoic Acid, USP (R1022)	0.1gm
Fluocinolone, USP	0.025gm
Niacinamide, USP (NI105)	4gm
Butylated Hydroxytoluene, NF (B1196)	0.4gm
Sodium Metabisulfite, NF (SO182)	0.3gm
Alcohol 95%, USP (ET108)	XmL
Polyethylene Glycol 300, NF (PO108)	Q.S. 100gm

SUGGESTED COMPOUNDING PROCEDURES

- 1. Calculate the required quantity needed for the total amount to be prepared
- 2. Weigh and/or measure all ingredients
- 3. Calibrate a beaker (with spin bar) to final volume
- 4. Add all dry chemicals to calibrated beaker
- 5. Add enough alcohol 95% to dissolve all powders
- 6. Bring to volume with Polyethylene Glycol 300 and mix until homogenous
- 7. Package and label
- 8. Suggested Quality assessments:
 - a. colorb. texturec. 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 based on the current USP <795> Standards

Precautions:

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 protective equipment. Use safety enclosures (hoods) when weighing and mixing.

Hazardous drugs should be handled according to USP <800>, NIOSH standards and pharmacy SOPs. This preparation should be compounded in a biological safety cabinet or compounding aseptic containment isolator. 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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