



#### SUGGESTED FORMULA

Chloramphenicol 1%/ Betamethasone Dipropionate 0.05%/ Ketoconazole 1% Ointment

Version number: 1.0

Volume or quantity: 60 gm

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Chloramphenicol, USP	0.6gm
Betamethasone Dipropionate, Micronized, USP (BE181)	0.03gm
Ketoconazole, USP (K1149)	0.6gm
Mineral Oil, Heavy, Oil, USP (MI500)	XmL
Petrolatum, White, USP (PE125)	Q.S. 60gm

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#### SUGGESTED COMPOUNDING PROCEDURES

1. Calculate the required quantity of each ingredient for the amount to be prepared
2. Accurately weigh and/or measure each ingredient
3. Triturate all the dry ingredients together in mortar and pestle
4. Use enough Mineral oil to form a paste and continue to triturate
5. Geometrically mix in Petrolatum until homogenous
6. Use of an Ointment mill maybe necessary to reduce particle size
7. Package and label; refrigerate after compounding
  - a. color
  - b. texture
  - c. container
  - d. Label - auxiliary labels, storage, BUD, compounded medication

Store in air-tight container, 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.

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