CORTANE-B- hydrocortisone, pramoxine hcl, chloroxylenol lotion
BLANSETT PHARMACAL CO

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

CORTANE-B LOTION

DOSAGE AND ADMINISTRATION

APPLY TO THE AFFECTED AREA 3 TO 4 TIMES DAILY. SEE INSERT FOR COMPLETE INSTRUCTIONS.

EACH 1 ML CONTAINS:

HYDROCORTISONE 10 MG

PRAMOXINE HCL 10 MG

CHLOROXYLENOL 1 MG

IS A BLAND VEHICLE WITH BENZALKONIUM CHLORIDE AS A PRESERVATIVE.

WARNING:

KEEP THIS AND ALL MEDICATION OUT OF THE REACH OF CHILDREN.

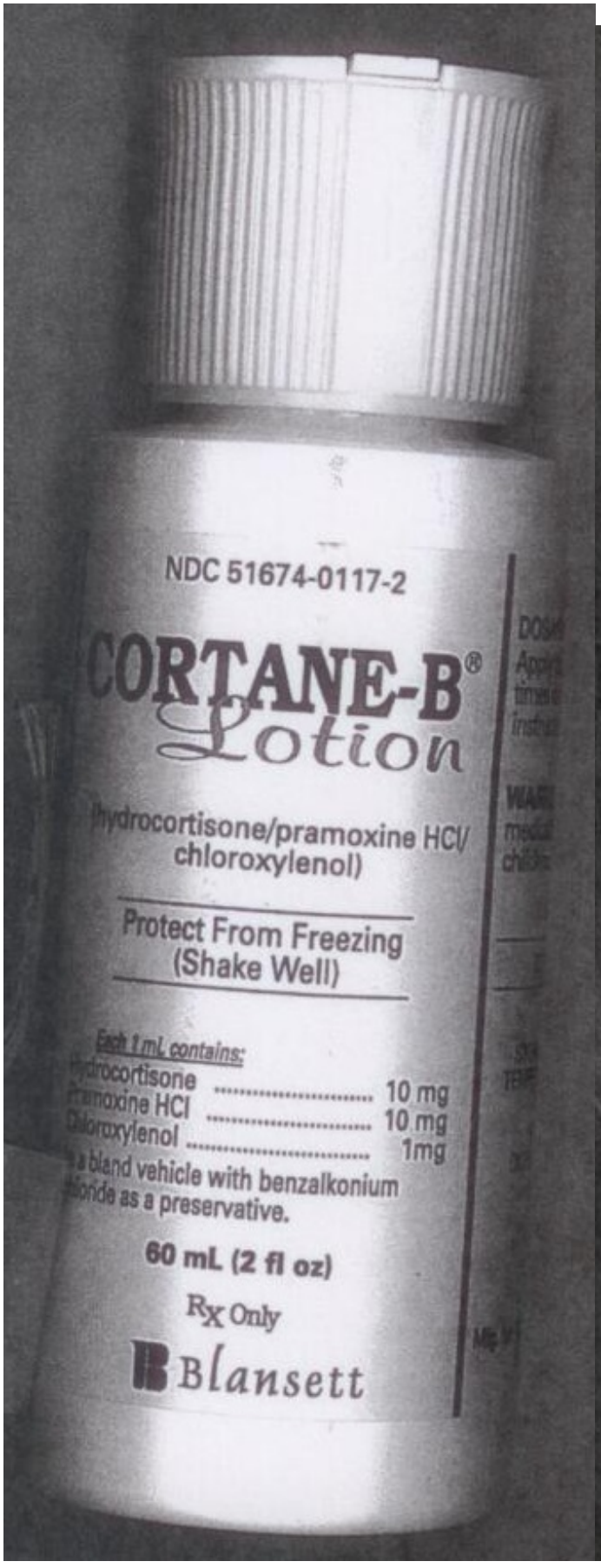
PROTECT FROM FREEZING (SHAKE WELL)

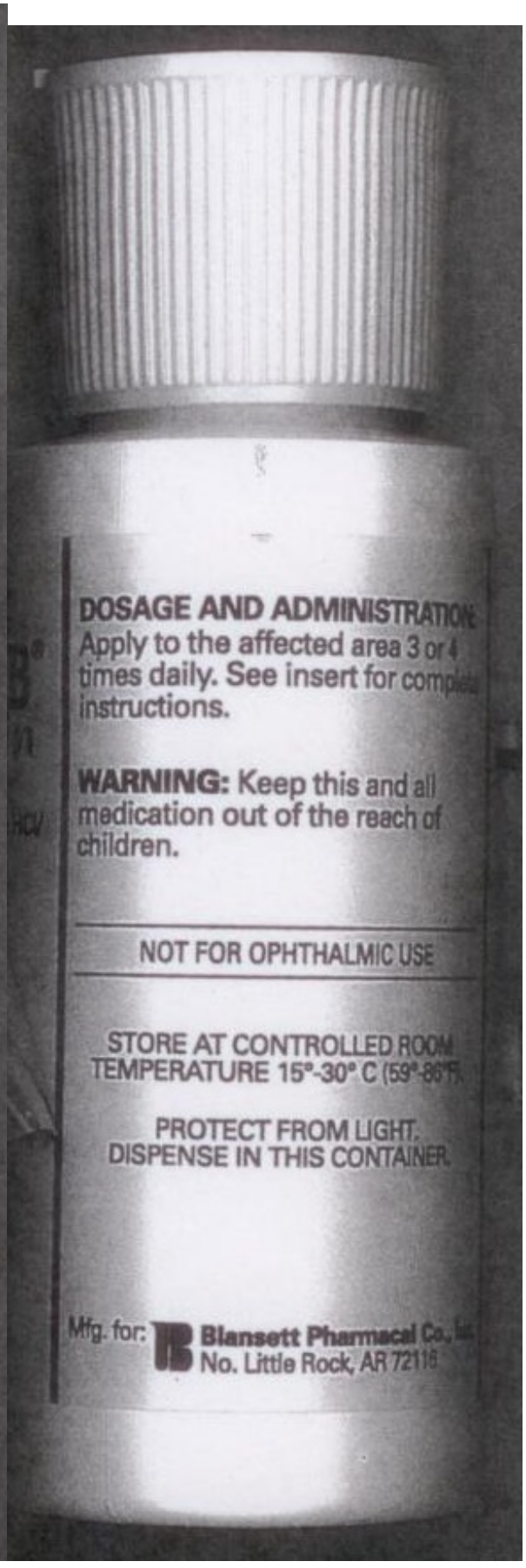
NOT FOR OPTHALMIC USE.

STORE AT CONTROLLED ROOM TEMPARATURE 15°-30°C (50°-86°F).

PROTECT FROM LIGHT.

DISPENSE IN CONTAINER.





CORTANE-B

hydrocortisone, pramoxine hcl, chloroxylenol lotion

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:51674-0117
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
HYDROCORTISONE (UNII: WI4X0X7BPJ) (HYDROCORTISONE - UNII:WI4X0X7BPJ)	HYDROCORTISONE	10 mg in 1 mL
PRAMOXINE HYDROCHLORIDE (UNII: 88AYB867L5) (PRAMOXINE - UNII:068X84E056)	PRAMOXINE HYDROCHLORIDE	10 mg in 1 mL
CHLOROXYLENOL (UNII: 0F32U78V2Q) (CHLOROXYLENOL - UNII:0F32U78V2Q)	CHLOROXYLENOL	1 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:51674-0117-2	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/20/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		03/20/2017	

Labeler - BLANSETT PHARMACAL CO (037477378)

Establishment

Name	Address	ID/FEI	Business Operations
BLANSETT PHARMACAL CO		037477378	manufacture(51674-0117)

Revised: 3/2017 BLANSETT PHARMACAL CO