CYCLOSPORINE/CHONDROITIN PF- cyclosporine/chondroitin sulfate pf emulsion
Imprimis NJOF, LLC

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.

Store at 20° to 25° C (68° to 77° F)

Sterile 5.5 ml Bottle
Cyclosporine 0.1% in Klarity®
Preservative Free
Ophthalmic Emulsion

Compounded for a licensed professional or patient use by

imprimis Rx

Imprimis NJOF
1705 Route 46 West, Suite 6B
Ledgewood, NJ 07852 (844) 446-6979

NDC 71384-514-05

Each mL contains: Cyclosporine USP 1mg
Inactive ingredients: Chondroitin Sulfate USP, Glycerin USP, Dextran 40 USP, Methocel E4M, Poloxamer 407 NF, Polysorbate 80 NF, Cremophor® EL, Ethyl Alcohol in Balanced Salt Solution and Sterile Water for injection. Sodium Hydroxide may have been used to adjust pH.

Store at controlled room temperature 20-25°C (68 -77°F). This medicine was compounded for you at the direction of your prescriber. Protect from light. Rx Only - Not for resale

Lot#
Date Compounded:
Use By:

CYCLOSPORINE/CHONDROITIN PF

cyclosporine/chondroitin sulfate pf emulsion

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:71384-514
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
CYCLOSPORINE (UNII: 83HN0GTJ6D) (CYCLOSPORINE - UNII:83HN0GTJ6D)	CYCLOSPORINE	1 mg in 1 mL

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:71384-514-05	5.5 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	07/01/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		07/01/2018	

Labeler - Imprimis NJOF, LLC (080431967)

Revised: 2/2020 Imprimis NJOF, LLC