CLORRELIEF- chlorpheniramine maleate aerosol, spray
FERRER MEDICAL INNOVATIONS LLC

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

ClorRelief (Chlorpheniramine Maleate 0.5%) Antihistamine Oral Spray

Active ingredient

Chlorpheniramine Maleate 0.5%

Purpose

Antihistamine

Uses

Temporarily relieves symptoms of hay fever or other upper respiratory allergies:

- runny nose
- itchy nose or throat
- sneezing itchy, watery eyes

Warnings

Ask a doctor before use if you have

glaucoma a breathing problem such as emphysema or chronic bronchitis trouble urinating due to an enlarged prostate gland

Ask a doctor or pharmacist before use if

you are taking sedatives or tranquilizers

When using this product

- you may get drowsy
- avoid alcoholic drinks, sedatives & tranquilizers, may increase drowsiness
- be careful when driving a motor vehicle or operating machinery
- excitability may occur, especially in children

If pregnant or breastfeeding,

ask your doctor before use.

Keep out of reach of children

Do not use to make a child sleepy. In case of overdose, getmedical help or contact a Poison Control Center right away.

Inactive ingredients

Aloe Vera, glycerin, grapefruit seed extract, hyaluronic acid, peppermint flavor, purified water, stevia, xylitol.

Directions

Adults and children 12 years and over:

For each application, spray 4 times every 4 to 6 hours,

do not exceed 6 applications in 24 hours.

Children 6 to under 12 years of age:
Should be supervised in the use of this product. For each application, spray 2 times every 4 to 6 hours. Do not exceed 3 applications in 24 hours.

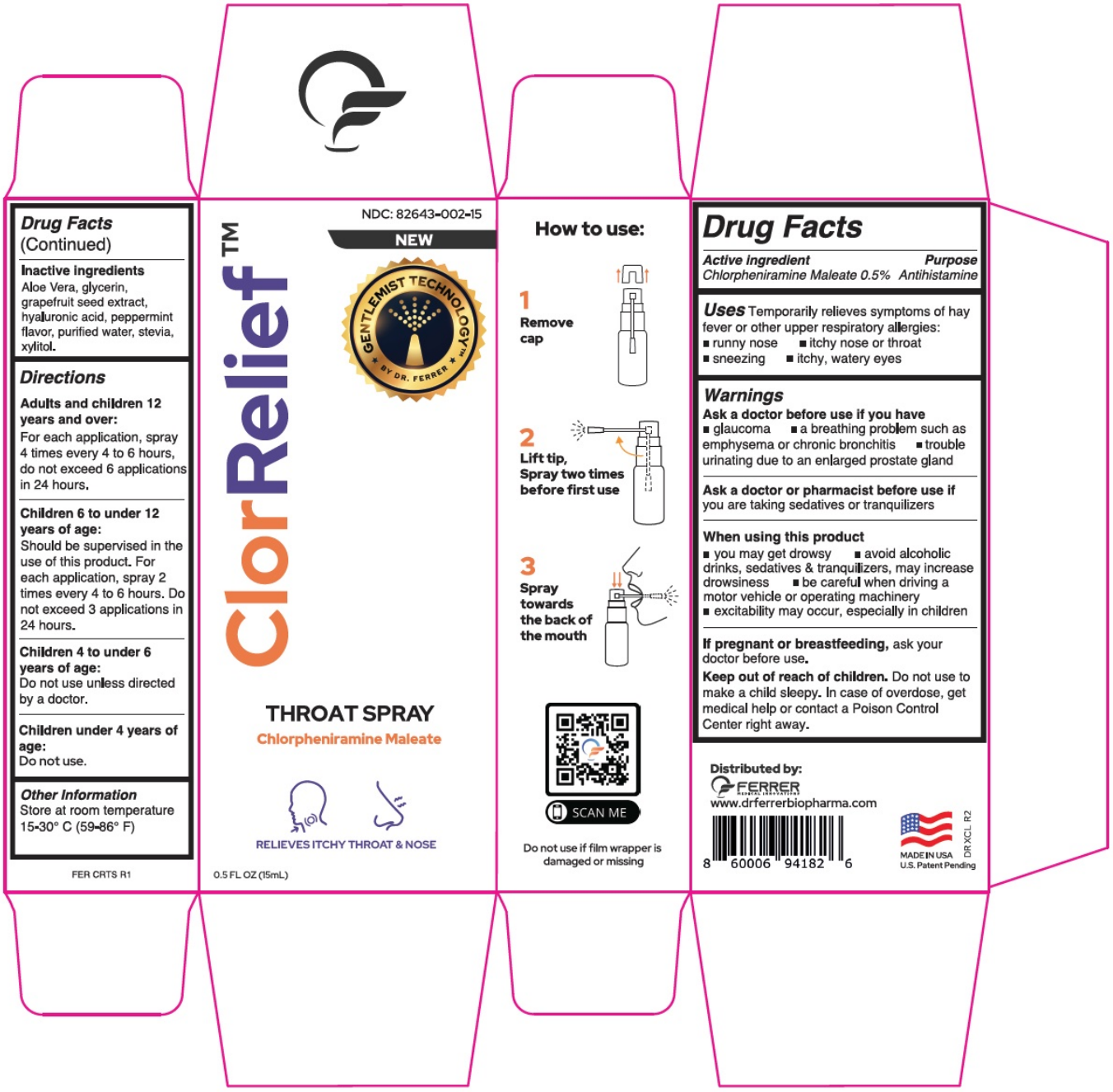
Children 4 to under 6 years of age:
Do not use unless directed by a doctor.

Children under 4 years of age:
Do not use.

Other Information

Store at room temperature 15-30° C (59-86° F)

ClorRelief Box



CLORRELIEF

chlorpheniramine maleate aerosol, spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:82643-002
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**

CHLORPHENIRAMINE MALEATE (UNII: V1Q0O9OJ9Z) (CHLORPHENIRAMINE - UNII:3U6IO1965U)	CHLORPHENIRAMINE MALEATE	5 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
XYLITOL (UNII: VCQ006KQ1E)	
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
ALOE VERA WHOLE (UNII: KIZ4X2EHYX)	
PEPPERMINT (UNII: V95R5KMY2B)	
GRAPEFRUIT SEED OIL (UNII: 598D944HOL)	
HYALURONIC ACID (UNII: S270N0TRQY)	
STEVIA LEAF (UNII: 6TC6NN0876)	

Product Characteristics

Color		**Score**	
Shape		**Size**	
Flavor	SPEARMINT, PEPPERMINT	**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82643-002-15	1 in 1 CARTON	04/24/2023	
1		15 mL in 1 BOTTLE, WITH APPLICATOR; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part341	04/21/2023	

Labeler - FERRER MEDICAL INNOVATIONS LLC (041608434)

Registrant - FERRER MEDICAL INNOVATIONS LLC (041608434)

Establishment

Name	Address	ID/FEI	Business Operations
Xlear Inc.		839884058	manufacture(82643-002)

Revised: 4/2023 FERRER MEDICAL INNOVATIONS LLC