HYDRA NEB- sodium chloride liquid
Laboratoire Unither

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.

Unither -VentureSaline Solution 0.9% Sodium Chloride

Drug Facts

Active ingredient

Sodium Chloride 0.9%

Purpose

Moisturizer

Uses:

For dry nasal membranes

Warnings

NOT FOR INJECTION

Only use sealed vials do not use if vial is broken or already open

Keep out of reach of children
Do not use after expiration date

If you experience any adverse reaction discontinue use and call your physician

- **Keep out of reach of children**
- **Do not use**

after expiration date
- Discontinue if you experience any adverse reaction and call your physician

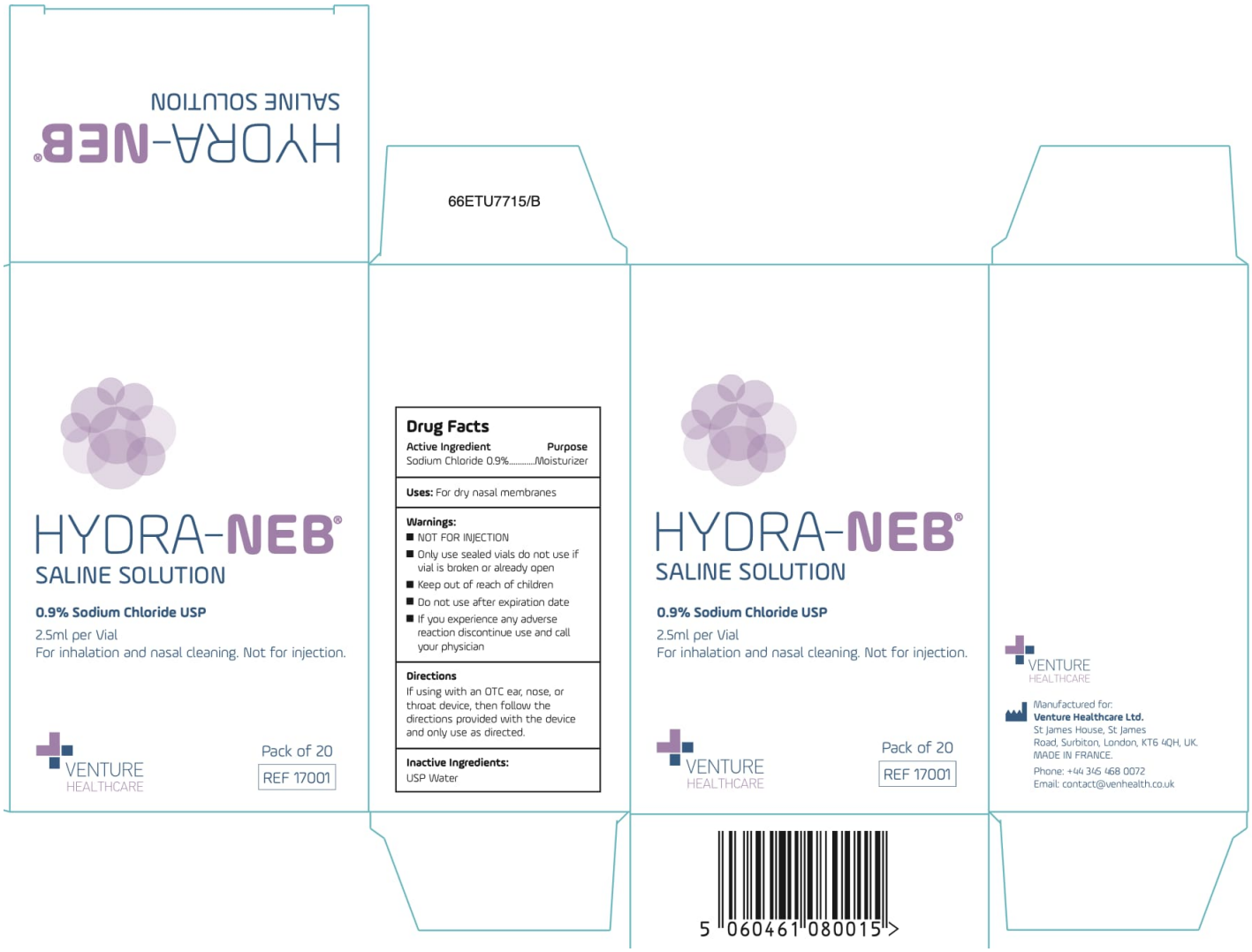
Directions

If using with an OTC ear, nose, or throat device, then follow the directions provided with the device and only use as directed.

Inactive ingredients

USP Water

Package Labelling:



2.5 mL NDC: 43014-1111-1

HYDRA NEB

sodium chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:43014-1111
Route of Administration	NASAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SODIUM CHLORIDE (UNII: 451W47IQ8X) (CHLORIDE ION - UNII:Q32ZN48698)	SODIUM CHLORIDE	9 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:43014-1111-1	20 in 1 CARTON	03/19/2025	
1		2.5 mL in 1 VIAL; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		03/19/2025	

Labeler - Laboratoire Unither (574139809)

Establishment

Name	Address	ID/FEI	Business Operations
Laboratoire Unither		574139809	manufacture(43014-1111)

Revised: 3/2025 Laboratoire Unither