ALEROFF- histamine liquid OKSALUD COLOMBIA S A S

Disclaimer: This homeopathic product has not been evaluated by the Food and Drug Administration for safety or efficacy. FDA is not aware of scientific evidence to support homeopathy as effective.

Active Ingredients

Histamine

Uses*

Adjuvant in the prevention and treatment of respiratory allergies. Transient allergic rhinitis. Profuse tearing. Profuse runny nose.

Warnings

Stop use and ask a doctor if symptoms persist for more than 3 days or worsen.

if pregnant or breastfeeding, ask a health proffesional before use.

Keep out of reach of children.

Directions

Adults: 15 Drops 3 times a day every 30 minutes beefore meals.

Childrens: 5 Drops 3 times a day every 30

minutes before meals.

Before use, shake the container by tapping with the palm of the hand.

Other Information

store between 20-25°C (68-77°F) in a dry place out of direct sunlight

Inactive Ingredients

Caladium (Floral essence)
Lycopersicum (Floral essence
Ranunculus bulbosus (Floral essence)
Tussilago (Floral essence)
Urtica dioica (Floral essence)

Questions?

(786) 453- 1395 Mon - Fri. 9am - 5pm EST

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KEEP OUT OF REACH OF CHILDRE IN A COOL PLACE AT A **TEMPERATURE BELOW 86°F** **ORAL DROPS 30 ML**

ALEROFF

histamine liquid

Product Information

Product Type HUMAN OTC DRUG **Item Code (Source)** NDC:81701-111

Route of Administration ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
HISTAMINE (UNII: 820484N8I3) (HISTAMINE - UNII:820484N8I3)	HISTAMINE	7 g in 7 mL

Inactive Ingredients

Ingredient Name	Strength
TUSSILAGO FARFARA FLOWER (UNII: UEE27X2Q2M)	7 mL in 7 mL
ALCOHOL (UNII: 3K9958V90M)	7 mL in 7 mL
CALADIUM BICOLOR WHOLE (UNII: 66BMR064Z9)	7 mL in 7 mL
RANUNCULUS BULBOSUS ROOT (UNII: XV11P9Y1A1)	7 mL in 7 mL
URTICA DIOICA (UNII: 710FLW4U46)	7 mL in 7 mL
SOLANUM LYCOPERSICUM WHOLE (UNII: 0243Q4990L)	7 mL in 7 mL

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	NDC:81701- 111-01	30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	03/01/2021	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		03/01/2021	

Labeler - OKSALUD COLOMBIA S A S (885622207)

Registrant - OKSALUD COLOMBIA S A S (885622207)

Establishment				
Name	Address	ID/FEI	Business Operations	
OKSALUD COLOMBIA S A S		885622207	manufacture(81701-111) , label(81701-111)	

Revised: 3/2021 OKSALUD COLOMBIA S A S