

EUPHRASIA- euphrasia stricta tablet
Standard Homeopathic Company

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.

EUPHRASIA 30X

Made according to the
Homeopathic Pharmacopoeia
of the United States since 1903.

Warnings

Do not use if cap band is missing or broken.

If you are pregnant or nursing, consult a licensed health care professional before using this product.

If symptoms persist for 7 days or worsen, contact a licensed practitioner.

Keep this and all medicines out of the reach of children.

To be used according to label indications and/or standard homeopathic indications.

Standard Homeopathic Company
Los Angeles, CA 90061
Questions? 800-624-9659

Directions

Adults: Dissolve 4 tablets under tongue 4 times a day.

Children: Consult a healthcare professional

In a base of Acacia Gum and Lactose N.F.

EUPHRASIA 30X

EYE IRRITATION, COLD OR HAY FEVER

Principal Display Panel - 250 Tablet Bottle Label

Standard

Homeopathic
Made in the USA Since 1903

NDC # 0360-0171-01

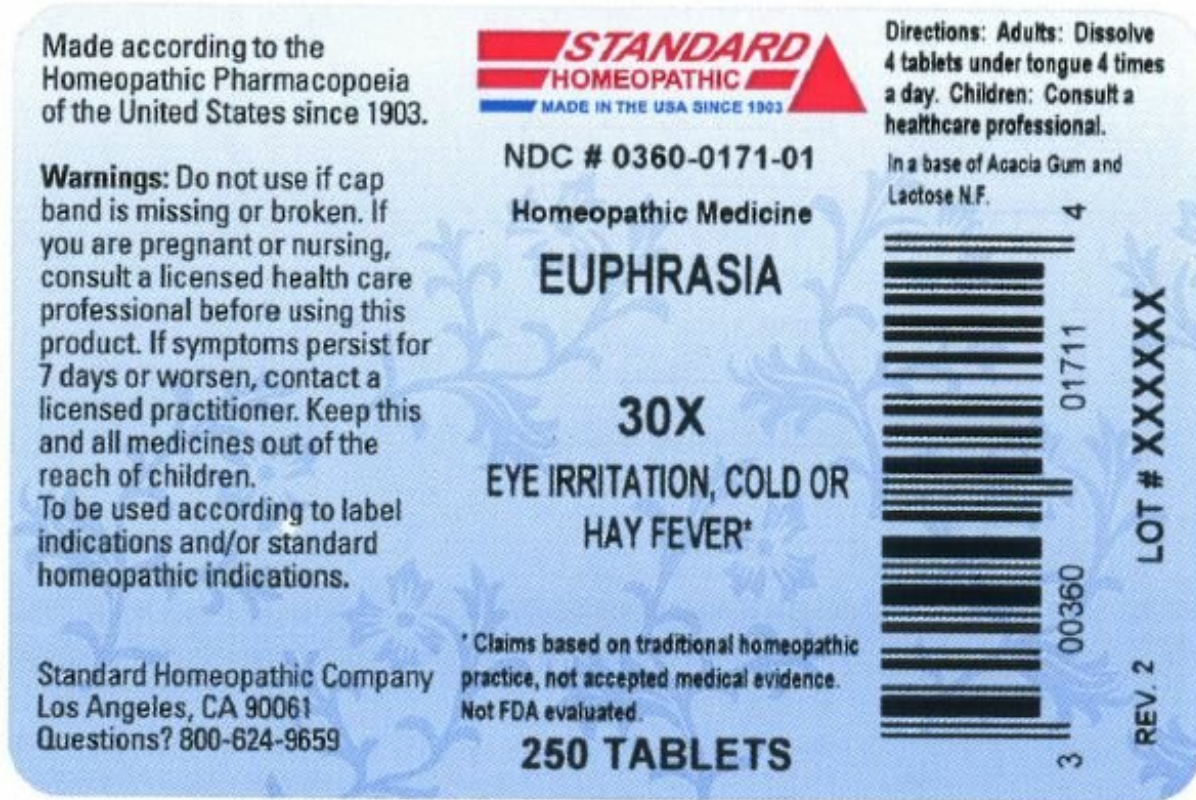
Homeopathic Medicine

EUPHRASIA
30X

EYE IRRITATION, COLD OR
HAY FEVER*

*Claims based on traditional homeopathic
practice, not accepted medical evidence.
Not FDA evaluated.

250 TABLETS



EUPHRASIA			
euphrasia stricta tablet			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0360-0171
Route of Administration	ORAL		
Active Ingredient/Active Moiety			
	Ingredient Name	Basis of Strength	Strength
	EUPHRASIA STRICTA (UNII: C9642I91WL) (EUPHRASIA STRICTA - UNII:C9642I91WL)	EUPHRASIA STRICTA	30 [hp_X]
Inactive Ingredients			
	Ingredient Name		Strength
	ACACIA (UNII: 5C5403N26O)		

LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)

Product Characteristics

Color	white	Score	no score
Shape	ROUND	Size	5mm
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0360-0171-01	250 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/17/1997	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		02/17/1997	

Labeler - Standard Homeopathic Company (008316655)

Establishment

Name	Address	ID/FEI	Business Operations
Standard Homeopathic Company		008316655	manufacture(0360-0171) , pack(0360-0171) , label(0360-0171)

Revised: 12/2018

Standard Homeopathic Company