PRUNUS SPINOSA E SUMM. 2X- prunus spinosa e summ. 2x liquid Uriel Pharmacy, Inc

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.

Prunus spinosa e summ. 2X

Directions: FOR ORAL USE ONLY.

Take 3-4 times daily. Ages 12 and older: 10 drops. Ages 2-11: 5 drops. Under 2: Consult a doctor.

Active Ingredient: Prunus spinosa e summ. (Blackthorn) 2X

Inactive Ingredients: Distilled water, 30% Organic cane alcohol

Prepared using rhythmical processes.

Use: Temporary relief of fatigue.

KEEP OUT OF REACH OF CHILDREN.

Warnings: Claims based on traditional homeopathic practice, not accepted medical evidence. Not FDA evaluated. Do not use if allergic to any ingredient. Consult a doctor before use for serious conditions or if conditions worsen or persist. If pregnant or nursing, consult a doctor before use. Do not use if safety seal is broken or missing.

Questions? Call 866.642.2858 Made with care by Uriel, East Troy, WI 53120 shopuriel.com Lot:





Homeopathic Liquid net vol. 2 fl. oz (60ml) KEEP OUT OF REACH OF CHILDREN.

Warrings: Claims based on traditional homespodisic prodice, not accepted medical evidence. Not FDA evaluated. Do not use if allergic to any ingredient. Com a doctor before use for selious conditions or if conditions unemen or persist. If pregnant or musing, consult a doctor before use. Do not use if safety seal is broken or missing. Question? Call 866,6422858

Made with care by Utiel, East Troy, WI 59120 shopuriel.com. Lot:

PRUNUS SPINOSA E SUMM. 2X

prunus spinosa e summ. 2x liquid

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:48951-8422	
Route of Administration	ORAL			

Active Ingredient/Active Moiety

Ingredient Name Basis of Strength Strength 2 [hp_X] in 1 mL

SLOE (UNII: 3MLB4858X7) (SLOE - UNII:3MLB4858X7) SLOE

Inactive Ingredients

Ingredient Name	Strength

WATER (UNII: 059QF0KO0R) ALCOHOL (UNII: 3K9958V90M)

Packaging

# Item Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC:48951- 8422-3	60 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	09/01/2009	

Marketing Information

Marketing	Application Number or Monograph	Marketing Start	Marketing End
Category	Citation	Date	Date
unapproved homeopathic		09/01/2009	

Labeler - Uriel Pharmacy, Inc (043471163)

Establishment				
Name	Address	ID/FEI	Business Operations	
Uriel Pharmacy, Inc		043471163	manufacture(48951-8422)	

Revised: 11/2024 Uriel Pharmacy, Inc