

OXY BEE POLLEN LIQUESCENCE 3012- oxy bee pollen liqescence liquid

Professional Complementary Health Formulas

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.

T12

ACTIVE INGREDIENTS

Bee pollen 2X
Flower pollen 2X
RNA-DNA (yeast) 2X
Amygdala amara 3X
Fucus vesiculosus 3X
Laminaria thallus 3X
Medicago sativa 3X
Royal jelly 3X
Zincum metallicum 4X, 6X
B vitamins (B1, B2, B3, B5, B6, B12) 6X
Symphytum officinale 6X

QUESTIONS

Professional Formulas

PO Box 2034 Lake Oswego, OR 97035

INDICATIONS

For the temporary relief of inflammation, mild pain or stiffness in joints, muscles, or surrounding tissues, irritability, poor concentration, diminished memory, fatigue, or general weakness.*

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

WARNINGS

In case of overdose, get medical help or contact a poison control center right away.

Keep out of the reach of children.

If pregnant or breastfeeding, ask a healthcare professional before use.

DIRECTIONS

Place drops under tongue 30 minutes before/after meals. Adults and children 12 years

and over: Take one full dropper up to 2 times per day. Consult a physician for use in children under 12 years of age.

OTHER INFORMATION

Tamper resistant. If seal is broken, do not use. After opening, close container tightly and store at room temperature away from heat.

INACTIVE INGREDIENTS

20% ethanol, purified water.

LABEL

Est 1985
Professional Formulas
Complementary Health
Oxy Bee Pollen Liqueescence
Homeopathic Remedy
4 FL. OZ. (118 mL)



OXY BEE POLLEN LIQUESCENCE 3012			
oxy bee pollen liqueescence liquid			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63083-3012
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
BEE POLLEN (UNII: 3729L8MA2C) (BEE POLLEN - UNII:3729L8MA2C)	BEE POLLEN	2 [hp_X] in 118 mL
ENGLISH SOLE (UNII: 00CR8124AS) (ENGLISH SOLE - UNII:00CR8124AS)	ENGLISH SOLE	2 [hp_X] in 118 mL
DNA-DIRECTED RNA POLYMERASE SUBUNIT BETA (M. TUBERCULOSIS) (UNII: 1C627X5G1B) (DNA-DIRECTED RNA POLYMERASE SUBUNIT BETA (M. TUBERCULOSIS) - UNII:1C627X5G1B)	DNA-DIRECTED RNA POLYMERASE SUBUNIT BETA (M. TUBERCULOSIS)	2 [hp_X] in 118 mL
BITTER ALMOND (UNII: O65SFW8R9A) (BITTER ALMOND - UNII:O65SFW8R9A)	BITTER ALMOND	3 [hp_X] in 118 mL
FUCUS VESICULOSUS (UNII: 535G2ABX9M) (FUCUS VESICULOSUS - UNII:535G2ABX9M)	FUCUS VESICULOSUS	3 [hp_X] in 118 mL
LAMINARIA JAPONICA (UNII: WE98HW412B) (LAMINARIA JAPONICA - UNII:WE98HW412B)	LAMINARIA JAPONICA	3 [hp_X] in 118 mL
MEDICAGO SATIVA WHOLE (UNII: DJO934BRBD) (MEDICAGO SATIVA WHOLE - UNII:DJO934BRBD)	MEDICAGO SATIVA WHOLE	3 [hp_X] in 118 mL
ROYAL JELLY (UNII: L497I37F0C) (ROYAL JELLY - UNII:L497I37F0C)	ROYAL JELLY	3 [hp_X] in 118 mL
ZINC (UNII: J41CSQ7QDS) (ZINC - UNII:J41CSQ7QDS)	ZINC	4 [hp_X] in 118 mL
PANTOTHENIC ACID (UNII: 19F5HK2737) (PANTOTHENIC ACID - UNII:19F5HK2737)	PANTOTHENIC ACID	6 [hp_X] in 118 mL
COMFREY ROOT (UNII: M9VVZ08EKQ) (COMFREY ROOT - UNII:M9VVZ08EKQ)	COMFREY ROOT	6 [hp_X] in 118 mL

Inactive Ingredients	
Ingredient Name	Strength
ALCOHOL (UNII: 3K9958V90M)	
WATER (UNII: 059QF0KO0R)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63083-3012-4	118 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	08/15/1985	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		08/15/1984	

Labeler - Professional Complementary Health Formulas (167339027)

Registrant - Natural Pharmaceutical Manufacturing LLC (015624923)

Establishment			
Name	Address	ID/FEI	Business Operations
Natural Pharmaceutical Manufacturing LLC		015624923	manufacture(63083-3012)

Revised: 1/2026

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