

OHM MENOPAUSE- ammonium carbonicum, amyl nitrosum, cimicifuga racemosa, glonoinum, lachesis mutus, sanguinaria, canadensis. liquid
OHM PHARMA 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.

OHM Menopause

ACTIVE INGREDIENTS (HPUS*): Equal amounts of: Ammonium Carbonicum 12C, Amyl Nitrosum 6X, Cimicifuga Racemosa 3X, Glonoinum 12X, Lachesis Mutus 12X, Sanguinaria Canadensis 4X.

*The letters "HPUS" indicate that the components in this product are officially monographed in the Homeopathic Pharmacopeia of the United States.

USES: Temporarily relieves symptoms caused by common menopause, hormonal imbalances.

*** This statement has not been evaluated by the FDA. It is based on documented Homeopathic Materia Medica.*

WARNINGS: IF PREGNANT OR BREAST-FEEDING, ask a health professional before use. If symptoms worsen or persist for more than a week, discontinue use and contact a doctor

- **KEEP OUT OF REACH OF CHILDREN.**

DIRECTIONS: Shake 10 times before use. Hold close to mouth and spray 3 times directly under the tongue 2-3 times a day, or as directed by a health care professional. Not for use by children.

Do not use if cap seal is broken.

INACTIVE INGREDIENTS: Organic Alcohol 15% (Non-GMO, Kosher Certified, Corn, Gluten, Allergen Free), Purified Water, Vegetable Glycerin 15%.

Mfg. By: OHM PHARMA, INC. Mineral Wells, TX 76067

www.ohmpharma.com

HOMEOPATHIC MEDICINE

NDC:66096-156-05

MENOPAUSE

Natural - Safe - Effective

No Side Effects

2 fl oz (59mL) / 15% Alcohol

PRODUCT OF USA

DRUG FACTS

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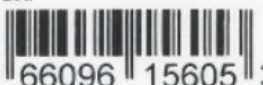

Your Partner in Homeopathy

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Lot:

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PHARMACEUTICAL GRADE PROFESSIONAL USE ONLY PRODUCT OF USA

Temporarily relieves symptoms caused by common menopause.

OHM MENOPAUSE

ammonium carbonicum, amyl nitrosum, cimicifuga racemosa, glonoinum, lachesis mutus, sanguinaria, canadensis. liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:66096-156
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
AMMONIUM CARBONATE (UNII: NJ5VT0FKLJ) (AMMONIUM CATION - UNII:54S68520I4)	AMMONIUM CARBONATE	12 [hp_C] in 59 mL
AMYL NITRITE (UNII: 22T8Z09XAK) (AMYL NITRITE - UNII:22T8Z09XAK)	AMYL NITRITE	6 [hp_X] in 59 mL
BLACK COHOSH (UNII: K73E24S6X9) (BLACK COHOSH - UNII:K73E24S6X9)	BLACK COHOSH	3 [hp_X] in 59 mL
NITROGLYCERIN (UNII: G59M7S0WS3) (NITROGLYCERIN - UNII:G59M7S0WS3)	NITROGLYCERIN	12 [hp_X] in 59 mL
LACHESIS MUTA VENOM (UNII: VSW71SS07I) (LACHESIS MUTA VENOM - UNII:VSW71SS07I)	LACHESIS MUTA VENOM	12 [hp_X] in 59 mL

SANGUINARIA CANADENSIS ROOT (UNII: N9288CD508) (SANGUINARIA
CANADENSIS ROOT - UNII:N9288CD508)

SANGUINARIA
CANADENSIS ROOT

4 [hp_X]
in 59 mL

Inactive Ingredients

Ingredient Name	Strength
ALCOHOL (UNII: 3K9958V90M)	
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:66096-156-05	59 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	08/19/2015	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		08/19/2015	

Labeler - OHM PHARMA INC. (030572478)

Revised: 12/2021

OHM PHARMA INC.