ARGENTUM MURIATICUM- argentum muriaticum pellet
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.

Argentum Muriaticum

DRUG FACTS: ACTIVE INGREDIENTS: ARGENTUM MURIATICUM

CONTENT: Approx. 100 Pellets

USE: To be used according to standard homeopathic indications.

WARNINGS: Keep out of reach of children. If pregnant or breast-feeding, ask a health care professional before use. **Stop use and ask a health care professional** if symptoms persist for more than 3 days or worsen.

Keep out of reach of children.

DIRECTIONS: Dissolve 3-5 pellets under the tongue 3 times a day or as directed by a professional.

OTHER INFORMATION: Store at room temperature. **Do not use** if pellet dispenser seal is broken.

INACTIVE INGREDIENTS: Organic sucrose, lactose free.

Manufactured according to the Homeopathic Pharmacopoeia of the United States (HPUS). **PRODUCT OF USA.**

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

www.ohmpharma.com FDA Est # 3003231743

ARGENTUM MURIATICUM

The OTC Potency range from Argentum Muriaticum 6X-30X, 3C-30C, 200C, 1M.

Standard bottle sizes for dilution-form can range from 30mL to 60mL.

To be used according to standard homeopathic indications.

ARGENTUM MURIATICUM

argentum muriaticum pellet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:66096-788
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SILVER CHLORIDE (UNII: MWB0804EO7) (SILVER CATION - UNII:57N7B0K90A)	SILVER CHLORIDE	6 [hp_C] in 6 [hp_C]

Inactive Ingredients	
Ingredient Name	**Strength**
SUCROSE (UNII: C151H8M554)	

Product Characteristics			
Color	white	**Score**	
Shape	ROUND	**Size**	4mm
Flavor		**Imprint Code**	
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:66096-788-01	6 [hp_C] in 1 TUBE; Type 0: Not a Combination Product	11/01/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		11/01/2019	

Labeler - OHM PHARMA INC. (030572478)

Establishment

Name	Address	ID/FEI	Business Operations
OHM PHARMA INC.		030572478	manufacture(66096-788)

Revised: 1/2021 OHM PHARMA INC.