

ZINCUM METALLICUM 12X- zincum metallicum liquid
Natural Creations, 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.

Zincum Metallicum 12X

ACTIVE INGREDIENTS (*HPUS) - Zincum Metallicum 12X

USES: Temporarily relieves symptoms of itchy skin.**

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DIRECTIONS: Adults & children above 12 years: 10 drops orally 3 times daily, or as directed by a health care professional.

KEEP OUT OF THE REACH OF CHILDREN. In case of overdose (or accidental ingestion) get medical help or contact a Poison Control Center right away.

WARNINGS:

- Consult a physician for use in children under 12 years of age.
- **IF PREGNANT OR BREAST-FEEDING**, ask a health care professional before use.
- **KEEP OUT OF THE REACH OF CHILDREN.** In case of overdose (or accidental ingestion) get medical help or contact a Poison Control Center right away.
- Do not use if **TAMPER EVIDENT** seal is broken or missing.

INACTIVE INGREDIENTS: Ethyl Alcohol USP, Purified Water.

QUESTIONS & COMMENTS?

Natural Creations, Inc. / Woodbine, IA 51579 / 712-647-1600

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

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

NDC:43406-0713-1

Zincum Metallicum 12X

HOMEOPATHIC

1 fl oz (30mL) / 20% Alcohol





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**M
CUM**



THIC
10% Alcohol

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HOMEOPA

1 fl oz (30 mL) / 20

ZINCUM METALLICUM 12X

zincum metallicum liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:43406-0713
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ZINC (UNII: J41CSQ7QDS) (ZINC - UNII:J41CSQ7QDS)	ZINC	12 [hp_X] in 1 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:43406-0713-1	30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	08/13/2020	

Marketing Information

Marketing	Application Number or Monograph	Marketing Start	Marketing End
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Category	Citation	Date	Date
unapproved homeopathic		08/13/2020	

Labeler - Natural Creations, Inc (018022074)

Establishment

Name	Address	ID/FEI	Business Operations
OHM Pharma, Inc		030572478	manufacture(43406-0713)

Revised: 12/2025

Natural Creations, Inc