

ZIRCONIUM CARBONICUM- zirconium carbonicum pellet
Hahnemann Laboratories, 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.

Single label image contains content for sections 1-17.

**Zirconium Carbonicum
30C**

EST 63545
Contents: 100g

HOMEOPATHIC MEDICINE.

Dose: Adult and child take 2 - 3 pellets once or as directed by your prescriber. **Warning: If pregnant, nursing, or if symptoms persist consult your practitioner.**

USE according to standard Homeopathic indications for self-limiting conditions, such as listed above.
DIRECTIONS: Dissolve one dose under tongue. Repeat as needed or as directed. Stop taking if symptoms worsen. **Keep out of reach of children.**
ACTIVE INGREDIENT listed above.
INACTIVE INGREDIENTS: sucrose.
TAMPER EVIDENCE: Use only if paper seal over cap is intact.
HAIHNIENJANNI LABS, INC., S.Rafael, CA 94901
QUESTIONS: Toll free 888-427 6422



Hahnemann Laboratories Inc. 1940 Fourth St. San Rafael, CA 94901
TRUST TO OPEN/USE ONLY IF SEALED

zirconium carbonicum pellet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63545-692
Route of Administration	BUCCAL, ORAL, SUBLINGUAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ZIRCONIUM CARBONATE HYDROXIDE OXIDE (UNII: JNE0 OSD75P) (ZIRCONIUM - UNII: C6 V6 S9 2N3C)	ZIRCONIUM CARBONATE HYDROXIDE OXIDE	30 [hp_C] in 1 [hp_C]

Inactive Ingredients

Ingredient Name	Strength
SUCROSE (UNII: C151H8M554)	

Product Characteristics

Color	white	Score	no score
Shape	ROUND	Size	3mm
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63545-692-01	240 [hp_C] in 1 VIAL, GLASS; Type 0: Not a Combination Product	05/08/2017	

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necessary to test for other microorganisms depending on the nature of the materials and the manufacturing process. If it has been shown that none of the prescribed tests will allow for detection of microorganisms at the level prescribed, a validated method will allow for detection as close as possible to the indicated acceptance criteria. In addition to the microorganisms listed in Table 1, the significant microorganisms recovered should be evaluated in terms of the fit

- The use of the product: hazard varies according to the route of exposure (eye, nose, respiratory tract).
- The nature of the product: does the product support growth? does it have adequate antimicrobial preservatives?
- The method of application.
- The intended recipient: risk may differ for neonates, infants, the debilitated.
- Use of immunosuppressive agents, corticosteroids.
- The presence of disease, wounds, organ damage.

Where warranted, a risk-based assessment of the relevant factors is conducted by personnel with specialized training in microbiology and in the interpretation of microbiological data. For raw materials, the assessment takes account of the processing to which the product is subjected, the current technology of testing, and the availability of materials of the desired quality.

Topic/Question	Contact	Expert Committee
General Chapter	Lisison Principal Scientist (301) 816-8339	Chapters-Microbiology (CCM2015) General 2015

USP92-WF34 Page 1321
Pharmacopoeial Forum: Volume No. 29(2) Page 1733

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
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unapproved homeopathic

05/08/2017

Labeler - Hahnemann Laboratories, Inc. (147098081)

Establishment

Name	Address	ID/FEI	Business Operations
Hahnemann Laboratories, Inc.		147098081	manufacture(63545-692)

Revised: 5/2017

Hahnemann Laboratories, Inc.