AMMONIUM BENZOICUM- ammonium benzoicum pellet
Rxhomeo Private Limited d.b.a. Rxhomeo, 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.

ACTIVE INGREDIENT

AMMONIUM BENZOICUM HPUS 2X and higher

USES

Gout

INDICATIONS

Condition listed above or as directed by the physician

DOSAGE

Adults- Take 4 or 6 Pellets by mouth, three times daily or as suggested by physician. Children 2 years and older- take 1/2 the adult dose.

WARNINGS

This product is to be used for self-limiting conditions

If symptoms do not improve in 4 days, or worsen, discontinue use and seek assistance of health professional

As with any drug, if you are preganant, or nursing a baby, seek professional advice before taking this product

Keep this and all medication out of reach of children

Do not use if capseal is broken or missing.

Close the cap tightly after use.

INACTIVE INGREDIENTS

Sucrose

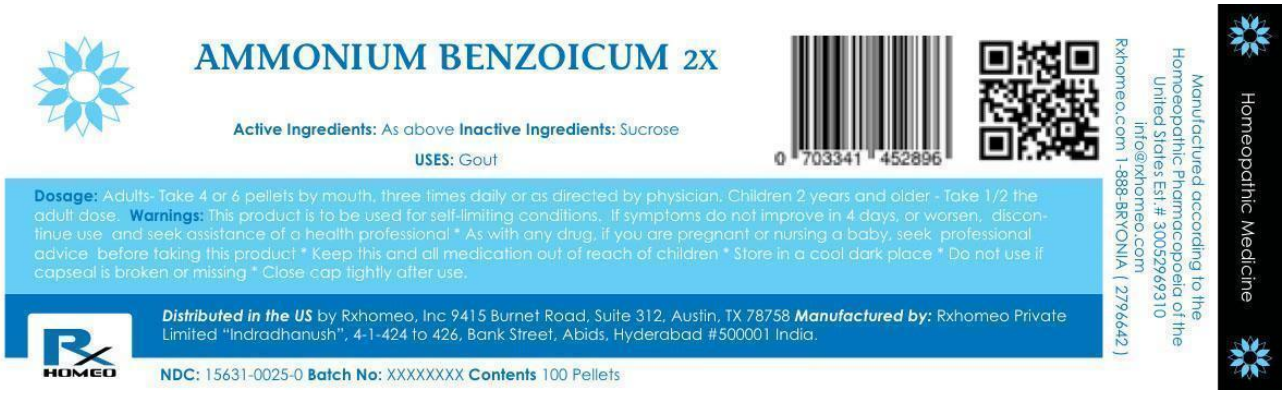
STORAGE

Store in a cool dark place

QUESTIONS OR COMMENTS

www.Rxhomeo.com | 1.888.2796642 | info@rxhomeo.com

Rxhomeo, Inc 9415 Burnet Road, Suite 312, Austin, TX 78758



AMMONIUM BENZOICUM

ammonium benzoicum pellet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:15631-0025
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
AMMONIUM BENZOATE (UNII: AC80WD7GPF) (AMMONIUM BENZOATE - UNII:AC80WD7GPF)	AMMONIUM BENZOATE	2 [hp_X]

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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:15631-0025-0	100 in 1 PACKAGE; Type 0: Not a Combination Product	01/01/2018	
2	NDC:15631-0025-1	200 in 1 PACKAGE; Type 0: Not a Combination Product	01/01/2018	
3	NDC:15631-0025-2	400 in 1 PACKAGE; Type 0: Not a Combination Product	01/01/2018	
4	NDC:15631-0025-3	750 in 1 PACKAGE; Type 0: Not a Combination Product	01/01/2018	
5	NDC:15631-0025-4	2500 in 1 PACKAGE; Type 0: Not a Combination Product	01/01/2018	
6	NDC:15631-0025-5	12500 in 1 PACKAGE; Type 0: Not a Combination Product	01/01/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		09/06/2015	

Labeler - Rxhomeo Private Limited d.b.a. Rxhomeo, Inc (650833994)

Establishment

Name	Address	ID/FEI	Business Operations
Rxhomeo Private Limited d.b.a. Rxhomeo, Inc		650833994	manufacture(15631-0025) , label(15631-0025)