## NUX MOSCHATA- nutmeg pellet Boiron

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.

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## Nux moschata 6C

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(\*\*contains 0.443 mg of the active ingredient per pellet)

Abdominal bloating accompanied by constipation\*

Stop use and ask a doctor if symptoms persist for more than 3 days or worsen

If pregnant or breast-feeding ask a health professional before use

Keep out of reach of children

Do not use if pellet dispenser seal is broken.

Contains approx 80 pellets.

How to dispense pellets? Turn tube upside down. Twist until 5 pellets are dispensed into cap. Carefully remove the cap and use it to pour pellets under the tongue. \*CLAIMS BASED ON TRADITIONAL HOMEOPATHIC PRACTICE NOT ACCEPTED MEDICAL EVIDENCE. NOT FDA EVALUATED.

\*C,K,CK, and X are homeopathic dilutions: see BoironUSA.com/info for details.

lactose, sucrose

Adults and children: At the onset of symptoms, dissolve 5 pellets under the tongue 3 times a day until symptoms are relieved or as directed by a doctor.

1-800-BOIRON-1 (1-800-264-7661), BoironUSA.com Info@boiron.com Distributed by Boiron, Inc. Newtown Square, PA 19073





## **Drug Facts**

**Active ingredient**\*\*: See product name on front panel (contains 0.443 mg of the active ingredient per pellet).

**Uses:** See symptoms on front panel.

**Warnings:** Stop use and ask a doctor if symptoms persist for more than 3 days or worsen. If pregnant or breastfeeding, ask a health professional before use. Keep out of reach of children.

**Directions:** Adults and children: At the onset of symptoms, dissolve 5 pellets under the tongue 3 times a day until symptoms are relieved or as directed by a doctor.

**Other information:** Do not use if pellet dispenser seal is broken.

Drug Facts (continued) Inactive ingredients: lactose, sucrose



NUX MOSCHATA				
nutmeg pellet				
Product Information				
Product Type	HUMAN OTC DRUG	ltem	Code (Source)	NDC:0220-3677
Route of Administration	ORAL			
Active Ingredient/Active	e Moiety			
Ingredie	ent Name		<b>Basis of Strength</b>	Strength
NUTMEG (UNII: AEE24M3MQ9) (N	NUTMEG (UNII: AEE24M3MQ9) (NUTMEG - UNII:AEE24M3MQ9) NUTMEG			6 [hp_C] in 6 [hp_C]
Inactive Ingredients				
	Strength			
SUCROSE (UNII: C151H8M554)				
LACTOSE, UNSPECIFIED FORM (UNII: J2B2A4N98G)				
Product Characteristics				
Color	white	Score		
Shape	ROUND	Size		4mm
Flavor	Imprint Code			

Co	ontains						
Packaging							
#	ltem Code	Package Description	Marketing Start Date	Marketing End Date			
1	NDC:0220-3677- 41	6 [hp_C] in 1 TUBE; Type 0: Not a Combination Product	03/03/1983				
Marketing Information							
	Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
	approved meopathic		03/03/1983				

Labeler - Boiron (282560473)

Registrant - Boiron, Inc. (014892269)

Establishment							
Name	Address	ID/FEI	<b>Business Operations</b>				
Boiron		282560473	manufacture(0220-3677)				

Revised: 7/2023

Boiron