NUX VOMICA- strychnos nux-vomica seed pellet Laboratoires 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.

Nux vomica 30C B2G1

Active ingredient** (in each pellet)

Nux Vomica 30C HPUS (**contains 0.44 mg of the active ingredient per pellet)

(contains less than 10⁻¹² strychnine alkaloids)

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

Relieves upset stomach, nausea, heartburn, and acid indigestion due to excessive drinking or eating, irritability

temporarily relieves upset stomach associated with nausea or fullness due to overindulgence in drink or food

temporarily relieves occasional heartburn, acid indigestion, or sour stomach

temporarily relieves irritability due to hypersensitivity to light, noise, odor or touch

Stop use and ask a doctorif hangover lasts for more than 2 days.

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

Keep out of reach of children. In case of accidental overdose, get medical help or contact a Poison Control Center right away.

Adults 21 years of age or older: 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.

Adults under 21 years of age: Not recommended.

lactose, sucrose

do not use if glued carton end flaps are open or if pellet dispenser seal is broken

contains 0.25 g of sugar per dose retain this carton for full directions Turn tube upside down. Twist until 5 pellets are dispensed. Remove cap and pour pellets under the tongue.

Approx. 80 pellets each

Total 240 pellets

3 Tubes

Hangover Relief*

On-the-Go Hangover Relief*

No Known Drug Interactions

Upset Stomach Nausea Heartburn Irritability*

*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.

Questions, Comments?

BoironUSA.com info@boiron.com

1-800-BOIRON-1 (1-800-264-7661)

Distributed by Boiron Inc. Newtown Square, PA 19073-3267



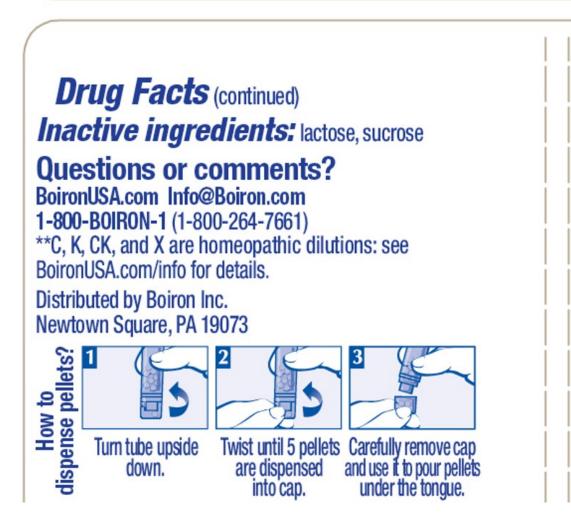
Active ingredient***: See product name on front panel (contains 0.443 mg of the active ingredient per pellet).

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

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

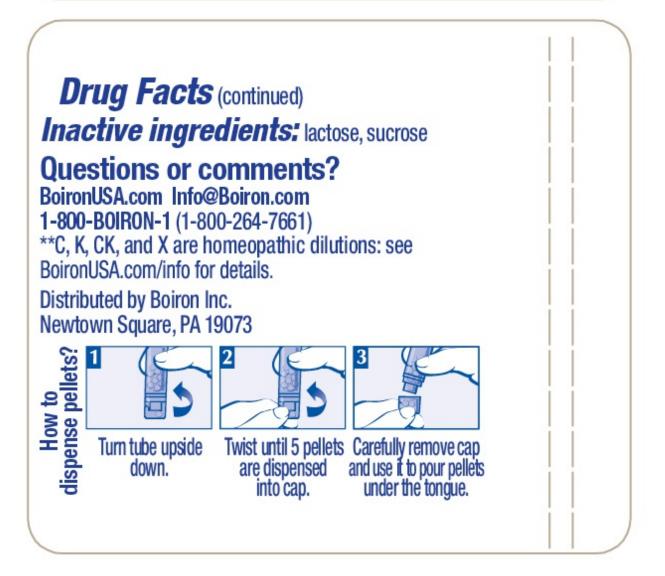
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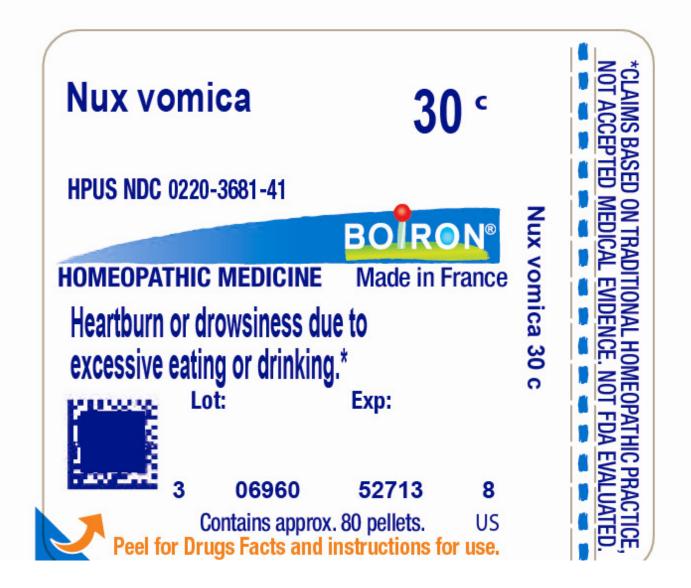
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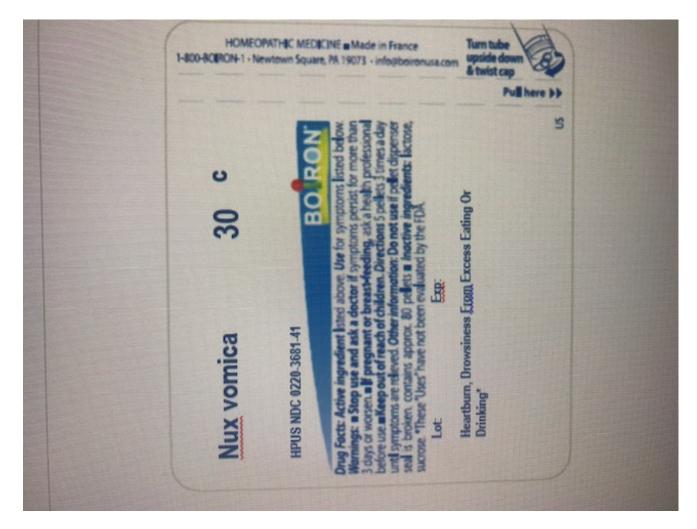
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Other information: Do not use if pellet dispenser seal is broken.

Drug Facts (continued) Inactive ingredients: lactose, sucrose Questions or comments? BoironUSA.com Info@Boiron.com 1-800-BOIRON-1 (1-800-264-7661) **C, K, CK, and X are homeopathic dilutions: see BoironUSA.com/info for details. Distributed by Boiron Inc. Newtown Square, PA 19073







NUX VOMICA trychnos nux-vomica seed p						
	allat					
	Jellet					
Product Information						
Product Type	HUMAN OTC DRUG	ltem Code (Sou	rce)	NDC:0220	-3681	
Route of Administration	ORAL					
Active Ingredient/Active Moiety						
Ingredient Name			Basis of S	trength	Strength	
STRYCHNOS NUX-VOMICA SEED (UNII: 269XH13919) (STRYCHNOS NUX- VOMICA SEED - UNII:269XH13919)			STRYCHNOS NUX- VOMICA SEED 30 [30 [hp_C]	
nostivo Ingradianta						
Inactive Ingredients				<u>.</u>	-	
Ingredient Name				Strength		
LACTOSE (UNII: J2B2A4N98G)						
SUCROSE (UNII: C151H8M554)						
Product Characteristics						

Color	white	Score	no score
Shape	ROUND (Pellet)	Size	4mm
Flavor		Imprint Code	
Contains			

Packaging

#	ltem Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0220-3681- 43	3 in 1 PACKAGE	01/01/2012	
1	NDC:0220-3681- 41	80 in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:0220-3681- 41	80 in 1 TUBE; Type 0: Not a Combination Product	01/01/2012	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		01/01/2012	

Labeler - Laboratoires Boiron (282560473)

Registrant - Boiron Inc. (014892269)

Establishment				
Name	Address	ID/FEI	Business Operations	
Boiron		282560473	manufacture(0220-3681)	

Revised: 11/2023

Laboratoires Boiron