

**ACONITUM NAPELLUS- aconitum napellus 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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**Aconitum napellus 9C**

Aconitum napellus 9C HPUS

Less than 10<sup>-12</sup> mg aconitine alkaloids per pellet

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

Uses: See symptoms on front panel.

Relieves high fever (up to 102f) of sudden onset with dry skin \*

**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),

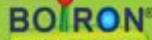
BoironUSA.com Info@boiron.com

Distributed by Boiron, Inc. Newtown Square, PA 19073

# Aconitum napellus

# 9<sup>c</sup>

Relieves high fever (up to 102F) of sudden onset with dry skin \*



Made in France HOMEOPATHIC MEDICINE

HPUS NDC 0220-0028-41

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	LOT	EXP	
	3 06960	01110	1

Contains approx. 80 pellets. US

Peel for Drugs Facts and instructions for use.

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## Drug Facts (continued)

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### Questions or comments?

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**9<sup>c</sup>**

HPUS NDC 0220-0028-41



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## ACONITUM NAPELLUS

aconitum napellus pellet

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:0220-0028
<b>Route of Administration</b>	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>ACONITUM NAPELLUS</b> (UNII: U0NQ8555JD) (ACONITUM NAPELLUS - UNII:U0NQ8555JD)	ACONITUM NAPELLUS	9 [hp_C] in 9 [hp_C]

### Inactive Ingredients

Ingredient Name	Strength
<b>LACTOSE, UNSPECIFIED FORM</b> (UNII: J2B2A4N98G)	
<b>SUCROSE</b> (UNII: C151H8M554)	

### Product Characteristics

<b>Color</b>	white	<b>Score</b>	
<b>Shape</b>	ROUND	<b>Size</b>	4mm
<b>Flavor</b>		<b>Imprint Code</b>	
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0220-0028-41	9 [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
unapproved homeopathic		03/03/1983	

**Labeler** - Boiron (282560473)

**Registrant** - Boiron, Inc. (014892269)

## Establishment

Name	Address	ID/FEI	Business Operations
Boiron		282560473	manufacture(0220-0028)

Revised: 9/2025

Boiron