ALOE- aloe ferox leaf 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.

Aloe 30C

Aloe 30C

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

Less than 10⁻¹² mg anthraquinone derivatives per pellet

Urgent diarrhea or hemorrhoids*

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



HUMEUPATHIC MEDICINE Made in France

HPUS NDC 0220-0187-41

Less than 10 ⁻¹² mg anthraquinone derivatives per pellet Do not use if pellet dispenser seal is broken



LOT: EXP:

3 06960 03713 2

Contains approx. 80 pellets. US Peel for Drugs Facts and instructions for use.



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)

Questions or comments?

BoironUSA.com Info@Boiron.com
1-800-BOIRON-1 (1-800-264-7661)

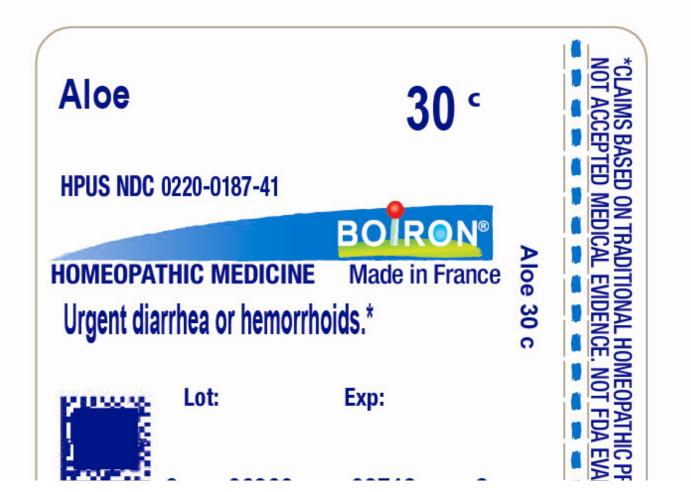
**C, K, CK, and X are homeopathic dilutions: see
BoironUSA.com/info for details.

Distributed by Boiron Inc.

Newtown Square, PA 19073

Turn tube upside down.

Turn tube upside are dispensed and use it to pour pellets under the tonque.





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.

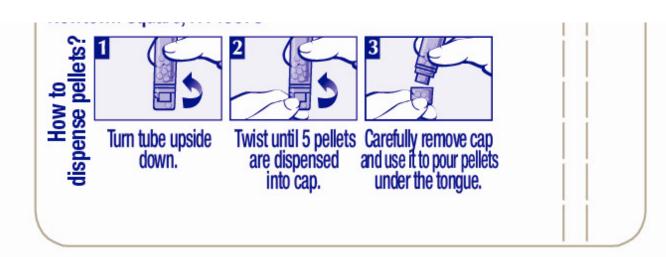
Other information: ■ Do not use if pellet dispenser seal is broken.

Drug Facts (continued) **Inactive ingredients:** lactose, sucrose

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



ALOE

aloe ferox leaf pellet

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:0220-0187

Route of Administration ORAL

Active Ingredient/Active Moiety Ingredient Name Basis of Strength ALOE FEROX LEAF (UNII: 0D145J8EME) (ALOE FEROX LEAF - UNII: 0D145J8EME) ALOE FEROX LEAF (UNII: 0D145J8EME) 30 [hp_C] in 30 [hp_C]

Inactive Ingredients		
Ingredient Name	Strength	
LACTOSE, UNSPECIFIED FORM (UNII: J2B2A4N98G)		
SUCROSE (UNII: C151H8M554)		

Product Characteristics			
Color	white	Score	
Shape	ROUND	Size	4mm
Flavor		Imprint Code	
Contains			

F	Packaging			
#	tem Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0220-0187- 41	30 [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(0220-0187)

Revised: 3/2023 Boiron