# TUBERCULINUM RESIDUUM- tuberculinum residuum 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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#### TUBERCULINUM RESIDUUM 30C MD

**TUBERCULINUM RESIDUUM 30C** 

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

Relieves stiffness and pain associated with minor arthritis\*

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.

Do not use if pellet dispenser seal is broken.

Contains approx 80 quick dissolving pellets per tube.

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.



### **TUBERCULINUM RESIDUUM**

tuberculinum residuum pellet

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0220-5066
Route of Administration	ORAL		

Active Ingredient/Active Moiety			
Ingredient Name	_	asis of Streng	th
TUBERCULINUM RESIDUUM (UNII: X4R8E2P3V3) (TUBER - UNII:X4R8E2P3V3)	CULINUM RESIDUUM TUBER RESIDU		[]

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

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

l	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:0220-5066- 41	30 [hp_C] in 1 TUBE; Type 0: Not a Combination Product	10/14/2022	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		10/14/2022	

### **Labeler -** Boiron (282560473)

## Registrant - Boiron, Inc. (014892269)

Establishment				
Name	Address	ID/FEI	Business Operations	
Boiron		282560473	manufacture(0220-5066)	

Revised: 1/2023 Boiron