REBOOST - antimony potassium tartrate syrup Heel Inc

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.

Reboost Cough Syrup

ACTIVE INGREDIENTS

Active Ingredients : Each 5ml contains: Antimonium tartaricum 6X, Cuprum sulphuricum 6X, Drosera rotundifolia 4X, Ipecacuanha 4X, Rumex crispus 4X, Spongia tosta 8X

INACTIVE INGREDIENTS

Inactive Ingredients: Purified Water, Honey, Preserved water and Simple syrup

PURPOSE

Antimonium tartaricum 6X	Relieves chest congestion
Cuprum sulphuricum 6X	Relieves cough
Drosera rotundifolia 4X	Relieves cough
Ipecacuanha 4X	Relieves cough
Rumex crispus 4X	Relieves cough
Spongia tosta 8X	Relieves cough

DOSAGE AND ADMINISTRATION

Standard Dosage: Adults and children 12 years and older: 2 teaspoons every 4 to 6 hours.

Children 4 to 11 tears: 1 teaspoon every 3 to 4 hours.

Children 1 to 4 years, consult your healthcare provider.

Initial Dosage: Adults and children 12 years and older: 2 teaspoons every 1 to 2 hours until symptoms lessen, then continue with standard dosage. Do not exceed 36 teaspoons in 24 hours.

Children 4 to 11 tears: 1 teaspoon every 1 to 2 hours until symptoms lessen, then continue with standard dosage. Do not exceed 18 teaspoons in 24 hours.

Children 1 to 4 years, consult your healthcare provider.

WARNINGS

Do not use in infants younger than 1 year due to the presence of honey. **If pregnant or breast-feeding,** ask a healthcare provider before use. **Keep out of reach of children.** If symptoms persist or worsen, a healthcare provider should be consulted. **Do not use** if known sensitivity to Reboost or any of its ingredients exists.

KEEP OUT OF REACH OF CHILDREN

Keep out of reach of children.

INDICATIONS AND USAGE

For the temporary relief of minor:

- Chest Congestion
- Cough due to colds and flu



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REBOOST antimony potassium tartrate syrup Product Information Product Type HUMAN OTC DRUG Route of Administration ORAL

Active Ingredien	0				
	Ingredient Name			of Strength	Strength
ANTIMONY POTASS (3+) - UNII:069647RPT		FARTRATE (UNII: DL6OZ476V3) (ANTIMONY CATION ANTIMON TARTRATE TARTRATE		POTASSIUM	6 [hp_X] in 125 mL
CUPRIC SULFATE (U	RIC SULFATE (UNII: LRX7AJ16DT) (CUPRIC CATION - UNII:8CBV67279L) CUPRIC CATION			ATION	6 [hp_X] in 125 mL
DROSERA ROTUNDIFOLIA (UNII: QR44N9XPJQ) (DROSERA ROTUNDIFOLIA - UNII:QR44N9XPJQ)			ROTUNDIFOLIA	4 [hp_X] in 125 mL	
IPECAC (UNII: 62I3C8233L) (IPECAC - UNII:62I3C8233L) IPECAC				4 [hp_X] in 125 mL	
RUMEX CRISPUS ROOT (UNII: 9N1RM2S62C) (RUMEX CRISPUS ROOT - UNII:9N1RM2S62C)			ISPUS ROOT	4 [hp_X] in 125 mL	
SPONGIA OFFICINALIS SKELETON, ROASTED (UNII: 1PIP394IID) (SPONGIA SPONGIA OFFICI OFFICINALIS SKELETON, ROASTED - UNII: 1PIP394IID) SKELETON, ROASTED - UNII: 1PIP394IID)				8 [hp_X] in 125 mL	
Inactive Ingredie	nts				
Inactive Ingredients Ingredient Name					ength
WATER (UNII: 059QF0	•	L		51	cingtin
HONEY (UNII: Y9H1V5	,				
	L (UNII: 6DC9Q167V3)				
METHYLPARABEN (U					
PROPYLPARABEN (U	,				
SUCROSE (UNII: C151	H8 M554)				
Packaging					
# Item Code	Package Description	Marketing Start Date Marke		Marketing I	End Date
1 NDC:50114-8550-1	1 in 1 CARTON				
1	125 mL in 1 BOTTLE				
Marketing Inf	ormation				
Maala da a Catana	A	and Charles Ma			

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		02/13/2013	

Labeler - Heel Inc (102783016)

Establishment				
Name	Address	ID/FEI	Business Operations	
Heel Belgium		282761204	manufacture(50114-8550)	

Revised: 2/2013