NATIVEREMEDIES BACKACHE EASE- arnica, bryonia, cimicifuga, hypericum, ignatia, kali carb, nux vom, rhus tox, ruta tablet Silver Star Brands

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.

NativeRemedies® Backache Ease Tablets

HPUS Active Ingredients

HPUS Active Ingredients: Equal volumes of each ingredient in a 12C potency: Arnica, Bryonia, Cimicifuga, Hypericum, Ignatia, Kali carb, nux vom, rhus tox, ruta.

Uses*:

Uses*: Homeopathic relief of symptoms of backache pain.

*Claims based on traditional homeopathic practice, not accepted medical evidence. Not FDA evaluated.

Warnings

Warnings: If symptoms persist or worsen, consult a healthcare professional. If pregnant or breastfeeding, ask a healthcare professional before use. keep this and all medication out of reach of children. In case of overdose, get medical help or contacta Poison Control Center right away.

Directions:

Directions: chew or dissolve tablets in the mouth. For Adults and Children 12+: Take 2 tablets every 15-20 minutes up to 4 doses, then every 4 hours until symptoms subside.

Contains no artificial flavors or colorants. No gluten added. Lactose Free.

Tamper resistsant for your protection. use only if saftey seal is intact.

Inactive Ingredients:

Inactive Ingredients: Magnesium Stearate, Maltodextrin, Sucrose.

Keep this and all medication out of reach of children.

If pregnant or breastfeeding, ask a health professional before use.

Temporarily relieves symptoms of backache pain.*

*claims based on traditional homeopathic practice, not accepted medical evidence. Not FDA evaluated.

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

NDC# 68703-373-80

All Native Remedies health products are especially formulated by experts in the field of natural health and are manufactured according to the highest pharmaceutical standards for maximum safety and effectiveness. For more information, visit us at www.nativeremedies.com.

Contains no artificial flavors or colorants. No gluten added. Lactose-free.

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SSB 01 r06-21 8 18837

NativeRemedies®



Backache Ease™

Temporarily relieves symptoms of backache pain*

180 TABLETS

Drug Facts

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Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:68703-373

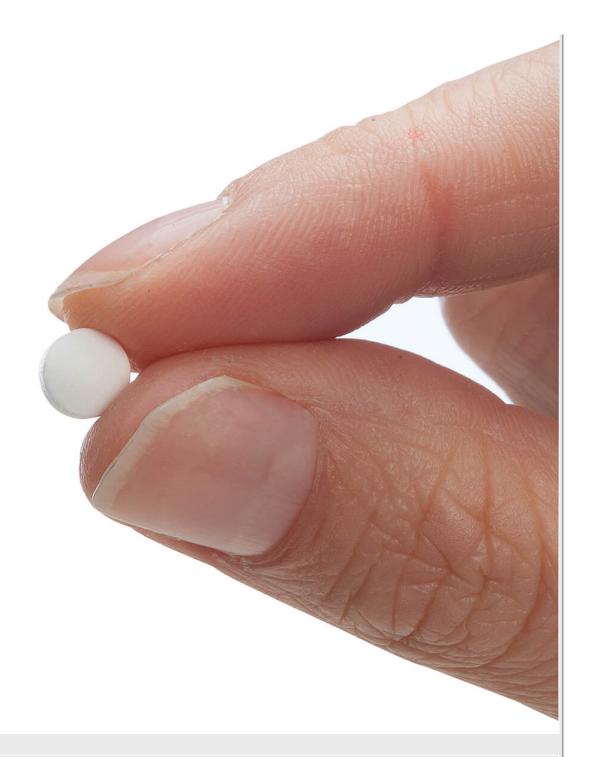
Route of Administration ORAL

Active Ingredient/Active Moiety Ingredient Name Basis of Strength Strength RUTA GRAVEOLENS FLOWERING TOP (UNII: N94C2U587S) (RUTA **RUTA GRAVEOLENS** 12 [hp C] GRAVEOLENS FLOWERING TOP - UNII:N94C2U587S) FLOWERING TOP in 180 HYPERICUM PERFORATUM (UNII: XK4IUX8MNB) (HYPERICUM PERFORATUM -**HYPFRICUM** 12 [hp_C] UNII:XK4IUX8MNB) **PERFORATUM** in 180 STRYCHNOS NUX-VOMICA SEED (UNII: 269XH13919) (STRYCHNOS NUX-STRYCHNOS NUX-12 [hp C] in 180 VOMICA SEED - UNII:269XH13919) VOMICA SEED ARNICA ANGUSTIFOLIA FLOWER (UNII: NG0U89QRD8) (ARNICA ANGUSTIFOLIA ARNICA ANGUSTIFOLIA 12 [hp_C] FLOWER - UNII:NG0U89QRD8) **FLOWER** in 180 BRYONIA ALBA ROOT (UNII: T7J046YI2B) (BRYONIA ALBA ROOT -12 [hp_C] **BRYONIA ALBA ROOT** UNII:T7J046YI2B) in 180 POTASSIUM CARBONATE (UNII: BQN1B9B9HA) (CARBONATE ION -**POTASSIUM** 12 [hp_C] UNII:7UJQ5OPE7D) **CARBONATE** in 180 STRYCHNOS IGNATII SEED (UNII: 1NM3M2487K) (STRYCHNOS IGNATII SEED -STRYCHNOS IGNATII 12 [hp C] UNII:1NM3M2487K) SEED in 180 ACTAEA CIMICIFUGA ROOT (UNII: 65/267BAZT) (ACTAEA CIMICIFUGA ROOT -ACTAEA CIMICIFUGA 12 [hp_C] UNII:65J267BAZT) ROOT in 180 **TOXICODENDRON** TOXICODENDRON RADICANS LEAF (UNII: CDH3461U7L) (TOXICODENDRON 12 [hp C] RADICANS LEAF - UNII:CDH3461U7L) RADICANS LEAF in 180

Inactive Ingredients			
Ingredient Name	Strength		
SUCROSE (UNII: C151H8M554)			
MAGNESIUM STEARATE (UNII: 70097M6I30)			
MALTODEXTRIN (UNII: 7CVR7L4A2D)			

Product Characteristics			
Color	white	Score	score with uneven pieces
Shape	ROUND	Size	3mm
Flavor		Imprint Code	none
Contains			

P	ackaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68703- 373-80	180 in 1 BOTTLE, GLASS; Type 0: Not a Combination Product	09/07/2021	



Category Citation Date Date unapproved	Marketing Information			
unapproved 09/07/2021		• • • • • • • • • • • • • • • • • • • •	_	Marketing End Date
nomeopatric	unapproved homeopathic		09/07/2021	

Labeler - Silver Star Brands (006070379)

Registrant - Silver Star Brands (006070379)

Establishment

Name	Address	ID/FEI	Business Operations
OHM Pharma, Inc.		030572478	manufacture(68703-373)

Revised: 1/2025 Silver Star Brands