

YEAST GARD ADVANCED- berberis vulgaris, borax, collinsonia canadensis, hamamelis virginiana douche

Wisconsin Pharmacal Company

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.

Active Ingredients

Borax 3X HPUS

Collinsonia Canadensis 3X HPUS

Berberis Vulgaris 6X HPUS

Hamamelis Virginiana 6X HPUS

Purpose

Temporary relief from vaginal itching, burning and irritation

Uses

Uses For the temporary relief of symptoms associated with vaginal yeast infection, including:

- itching
- burning
- irritation

Warnings

- Use only as directed
- For vaginal use only

Ask a doctor before use if you have

- Vaginal itching and discomfort for the first time
- Lower abdominal, back or shoulder pain, fever, chills, nausea, vomiting, or foul-smelling vaginal discharge. You may have a serious condition.
- Vaginal yeast infections often (such as once a month or 3 in 6 months). You could be pregnant or have a serious underlying medical cause for your symptoms, including diabetes or a weakened immune system.
- Been exposed to the human immunodeficiency virus (HIV) that causes AIDS
- You have an STD (sexually transmitted disease) or PID (Pelvic inflammatory disease)
- You have frequent and painful urination

Stop use and ask a doctor if:

- Symptoms last more than 7 days
- You get a rash or hives, abdominal pain, fever, chills, nausea, vomiting, or foul-smelling vagina.

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away

Do not use

- if you have never had a vaginal yeast infection diagnosed by a doctor

When using this product

- if you do not get complete relief ask a doctor before using another product

Directions**Directions**

Tamper Evident: If the foil seal on bottle opening is missing, torn or damaged in any way, do not use.

- Wash hands thoroughly with soap and water before and after use
- remove cap and safety seal from bottle. Carefully open the packet and pour homeopathic contents into the douche bottle. Screw cap back onto bottle and shake thoroughly to fully mix powder into solution. After fully mixed, remove cap. Remove sanitary wrap from nozzle. Screw nozzle onto douche bottle.
- Slowly insert nozzle into vagina about 3 inches. If needed, a small amount of personal lubricant can be applied to nozzle to ease insertion of nozzle into vagina. Gently squeeze bottle until solution is dispensed. Use while sitting on the toilet, standing in the shower or in the tub. After douching discard bottle and nozzle.

Other information

- a douche is not a contraceptive and should not be used to prevent pregnancy
- **do not** use if foil seal on bottle opening is missing, torn damaged in any way. use if homeopathic packet is torn, open or incompletely sealed. purchase if carton is open
- Store at room temperature
- Lot number and expiration date: see upper back panel of box
- There is no scientific evidence that this product works. The product claims are based on theories of homeopathy from the 1700's that are not accepted by most modern medical experts.

Inactive ingredients

Bacillus coagulans, citric acid, edetate disodium, maltodextrin, purified water, sodium benzoate, sodium lauryl sulfate, trisodium phosphate

Questions or comments? 1-800-635-3696



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Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68093-0668
Route of Administration	VAGINAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
HAMAMELIS VIRGINIANA ROOT BARK/STEM BARK (UNII: T7S323PKJS) (HAMAMELIS VIRGINIANA ROOT BARK/STEM BARK - UNII:T7S323PKJS)	HAMAMELIS VIRGINIANA ROOT BARK/STEM BARK	6 [hp_X] in 1 g
BERBERIS VULGARIS ROOT BARK (UNII: 1TH8Q20J0U) (BERBERIS VULGARIS ROOT BARK - UNII:1TH8Q20J0U)	BERBERIS VULGARIS ROOT BARK	6 [hp_X] in 1 g
SODIUM BORATE (UNII: 91MBZ8H3QO) (BORATE ION - UNII:44OAE30D22)	SODIUM BORATE	3 [hp_X] in 1 g
COLLINSONIA CANADENSIS ROOT (UNII: O2630F3XDR) (COLLINSONIA CANADENSIS ROOT - UNII:O2630F3XDR)	COLLINSONIA CANADENSIS ROOT	3 [hp_X] in 1 g

Inactive Ingredients

Ingredient Name	Strength
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
BACILLUS COAGULANS (UNII: ISK1LOY57E)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
MALTO DEXTRIN (UNII: 7CVR7L4A2D)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
WATER (UNII: 059QF0K00R)	
SODIUM PHOSPHATE, TRIBASIC, ANHYDROUS (UNII: SX01TZO3QZ)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68093-0668-1	2 in 1 CARTON	10/01/2007	
1		133.07 g in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		10/01/2007	

Labeler - Wisconsin Pharmacal Company (800873986)

Establishment

Name	Address	ID/FEI	Business Operations
Wisconsin Pharmacal Company		800873986	manufacture(68093-0668)

Revised: 12/2018

Wisconsin Pharmacal Company