YEAST GARD HOMEOPATHIC GEL HOMEOPATHIC GEL TREATMENT- borax, collinsonia canadensis, berberis vulgaris, hamamelis virginiana gel 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 external vagianl discomfort including itching, burning, and irritation

Uses For temporary relief of symptoms associate with vaginal yeast infection, including: itching, burning, and irritation.

Warnings

For external vaginal use only. Use only as directed.

Keep out of reach of children. If swallowed, get medial help or contact a Poison Control Center right away. If in eyes, flush with water for 15 minutes and call a physician.

Stop use and ask doctor if

- Symptoms worsen, last more than 7 days, or clear up and occur again within a few days.
- You get a rash or hives, abdominal pain, fever, chills, nausea, vomiting, or foul-smelling vaginal discharge.

Do not use if

• you have never had a vaginal yeast infection diagnosed by a doctor.

When using this product

- Wash hands thoroughly with soap and water before and after use
- Avoid contact with eyes
- If burning sensation occurs and lasts more than a few minutes, remove gel and discontinue use
- Do not use more than one week unless directed by a doctor
- If yoyu do not get complete relief ask a doctor before using another product.

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 you 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

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

Directions

Directions

- Apply to affected area as often as needed. Because the area is irritated, you could experience a slight burning sensation. If this persists more than a few minutes, remove gel and discontinue use.
- Wash hand thoroughly with soap an dwater before and after use.
- Do not use for children under two years. Consult a doctor if condition persists.

Inactive Ingredients

Allantoin, Benzalkonium Chloride 50%, Disodium EDTA, Glucono Delta Lactone, Polysorbate 20, Propanediol, Sodium Benzoate, Sodium Hydroxide, Tea Tree Oil, Water, Xanthan Gum

Other Information

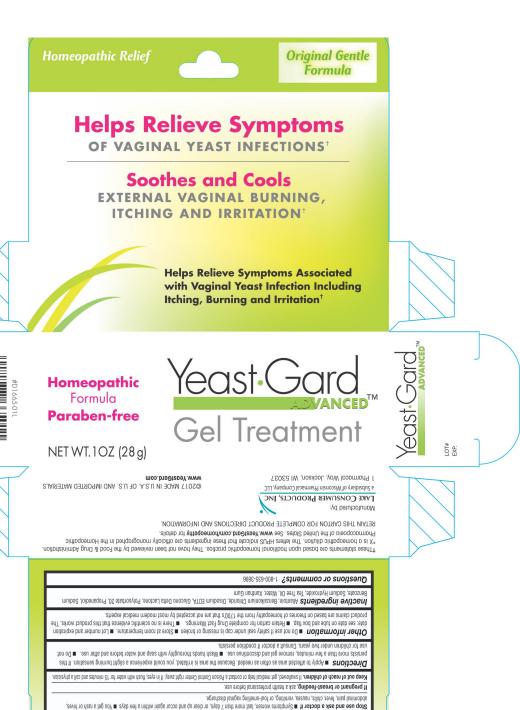
Other information

- Do not use if safety seal under cap is missing or broken
- Store at room temperature
- Lot number and expiration date: see date on tube and box flap.
- Retain carton for complete Drug Facts Warnings
- There is no scientific evidence that this product works. Ther product claims are based on theories of homeopathy from the 1700's that are not accepted by most modern medical experts.

Questions or comments? 1-800-635-3696



Active ingredient: Boxx 3X HPUS - (Sodium Borne), Colinscoin Canodensis SX HPUS - (Stone Boot), Betheris Vulgaris 6X HPUS indicates
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Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68093-7247
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		
COLLINSONIA CANADENSIS ROOT (UNII: O2630F3XDR) (COLLINSONIA CANADENSIS ROOT - UNII:O2630F3XDR)	COLLINSONIA CANADENSIS ROOT	3 [hp_X] in 1 g		
SODIUM BORATE (UNII: 91MBZ8H3QO) (BORATE ION - UNII:44OAE30D22)	SODIUM BORATE	3 [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		

Inactive Ingredients		
Ingredient Name	Strength	
PROPANEDIOL (UNII: 5965N8W85T)		
XANTHAN GUM (UNII: TTV12P4NEE)		
SO DIUM HYDRO XIDE (UNII: 55X04QC32I)		
SODIUM BENZOATE (UNII: OJ245FE5EU)		
GLUCONOLACTONE (UNII: WQ29KQ9POT)		
EDETATE DISO DIUM (UNII: 7FLD9 1C86K)		
ALLANTO IN, (+)- (UNII: XDK458 E1J9)		
BENZALKO NIUM CHLO RIDE (UNII: F5UM2KM3W7)		
WATER (UNII: 059QF0KO0R)		
POLYSORBATE 20 (UNII: 7T1F30V5YH)		
TEA TREE OIL (UNII: VIF565UC2G)		

Packaging			
# Item Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC:68093-7247-1	1 in 1 CARTON	04/01/2007	
1	28.35 g in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		04/01/2007	

Labeler - Wisconsin Pharmacal Company (800873986)

Establishment				
Name	Address	ID/FEI	Business Operations	
Wisconsin Pharmacal Company		800873986	manufacture(68093-7247)	

Revised: 12/2018 Wisconsin Pharmacal Company