ABROPERNOL - artemisia abrotanum flowering top and pulsatilla vulgaris and calcium fluoride and hamamelis virginiana root bark/stem bark and amanita muscaria var. muscaria fruiting body and nitric acid and kerosene tablet
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.

Abropernol Tablet

KEEP OUT OF REACH OF CHILDREN
Keep this and all medications out of the reach of children.

INDICATIONS AND USAGE
For the temporary relief of:
- Dermatitus
- Minor skin irritations
- Eczema, hyperhidrosis
- Hyperkeratosis

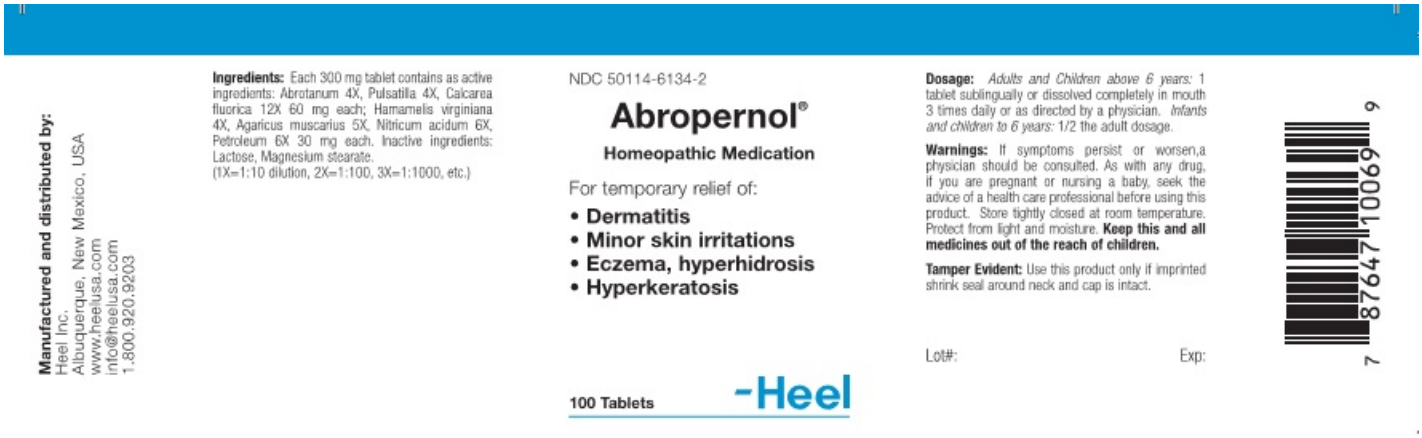
WARNINGS
If symptoms persist or worsen, a physician should be consulted. As with any drug, if you are pregnant or nursing a baby, seek the advise of a health care professional before using this product.

DOSAGE AND ADMINISTRATION
Adults and children above 6 years: 1 tablet sublingually or dissolved completely in mouth 3 times daily or as directed by a physician.
Infants and children to 6 years: 1/2 the adult dosage.

ACTIVE INGREDIENT
Each 300mg tablet contains as active ingredients: Abrotanum 4X, Pulsatilla 4X, Calcarea fluorica 12X 60 mg each; Hamamelis virginiana 4X, Agaricus muscarius 5X, Nitricum acidum 6X, Petroleum 6X 30 mg each.

INACTIVE INGREDIENT
Inactive ingredients: Lactose, Magnesium Stearate

PURPOSE
Dermatitus, Minor skin irritations, Eczema, hyperhidrosis, Hyperkeratosis



ABROPERNOL

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Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50114-6134
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ARTEMISIA ABROTANUM FLOWERING TOP (UNII: QG07G580U0) (ARTEMISIA ABROTANUM FLOWERING TOP - UNII:QG07G580U0)	ARTEMISIA ABROTANUM FLOWERING TOP	4 [hp_X] in 300 mg
PULSATILLA VULGARIS (UNII: I76KB35JEV) (PULSATILLA VULGARIS - UNII:I76KB35JEV)	PULSATILLA VULGARIS	4 [hp_X] in 300 mg
CALCIUM FLUORIDE (UNII: O3B55K4YKI) (CALCIUM FLUORIDE - UNII:O3B55K4YKI)	CALCIUM FLUORIDE	12 [hp_X] in 300 mg
HAMAMELIS VIRGINIANA ROOT BARK/STEM BARK (UNII: T7S323PKJS) (HAMAMELIS VIRGINIANA ROOT BARK/STEM BARK - UNII:T7S323PKJS)	HAMAMELIS VIRGINIANA ROOT BARK/STEM BARK	4 [hp_X] in 300 mg
AMANITA MUSCARIA VAR. MUSCARIA FRUITING BODY (UNII: DIF093I037) (AMANITA MUSCARIA VAR. MUSCARIA FRUITING BODY - UNII:DIF093I037)	AMANITA MUSCARIA VAR. MUSCARIA FRUITING BODY	5 [hp_X] in 300 mg
NITRIC ACID (UNII: 411VRN1TV4) (NITRIC ACID - UNII:411VRN1TV4)	NITRIC ACID	6 [hp_X] in 300 mg
KEROSENE (UNII: 1C89KKC04E) (KEROSENE - UNII:1C89KKC04E)	KEROSENE	6 [hp_X] in 300 mg

Inactive Ingredients

Ingredient Name	Strength
LACTOSE (UNII: J2B2A4N98G)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	

Product Characteristics

Color	white	**Score**	no score
Shape	ROUND (Tablet)	**Size**	9mm
Flavor		**Imprint Code**	Heel

Contains	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50114-6134-2	30000 mg in 1 BOTTLE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		01/31/1984	

Labeler - Heel Inc (102783016)

Establishment

Name	Address	ID/FEI	Business Operations
Heel Inc		102783016	manufacture

Revised: 12/2011 Heel Inc