HAMSOA ATOBY MEDI- witch hazel lotion
HAMSOA PHARMACEUTICAL CO., LTD.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Active ingredient

Active ingredient: Witch hazel(Hamamelis Virginiana) 0.1%

Inactive ingredient

Inactive ingredients:
Water, Sorbitol, Lavandula Angustifolia (Lavender) Extract, Ocimum Basilicum (Basil) Extract, Althaea Officinalis Extract, Houttuynia Cordata Extract, Anthemis Nobilis Flower Extract, Rosmarinus Officinalis (Rosemary) Leaf Extract, Foeniculum Vulgare (Fennel) Fruit Extract, Hydrogenated Polydecene, Glycerin, Cetyl Ethylhexanoate, Pentylene Glycol, Stearic acid, Betaine, Glyceryl Stearate, PEG-100 Stearate, Sorbitan Stearate, Cetearyl Alcohol, Cetearyl Glucoside, Hydrogenated Lecithin, Butyrospermum Parkii (Shea Butter), Dimethicone, Allantoin, Carbomer, Arginine, Tremella Fuciformis Sporocarp Extract, Butylene Glycol, 1,2-Hexanediol, Caprylyl Glycol, Tropolone, Citrus Unshiu Fruit Oil, Disodium EDTA, Portulaca Oleracea Extract, Coptis Japonica Root Extract, Salvia Miltiorrhiza Root Extract, Schizandra Chinensis Fruit Extract, Rhus Semialata Gall Extract, Glycyrrhiza Glabra (Licorice) Root Extract, Vigna Radiata Seed Extract, Scutellaria Baicalensis Root Extract, Sophora Angustifolia Root Extract, Paeonia Suffruticosa Root Extract, Paeonia Albiflora Root Extract, Citrus Grandis (Grapefruit) Peel Oil, Citrus Nobilis (Mandarin Orange) Peel Oil, Mentha Viridis (Spearmint) Leaf Oil, Ceramide 3, PEG-10 Rapeseed Sterol, Glycereth-20, Dipropylene Glycol, Malt Extract Concentrated Powder, Cananga Odorata Flower Oil, Lavandula Angustifolia (Lavender) Oil, Eucalyptus Globulus Leaf Oil, Pelargonium Graveolens Flower Oil, Citrus Medica Limonum (Lemon) Peel Oil, Citrus Aurantium Dulcis (Orange) Fruit Extract, Abies Sibirica Oil

Purpose

Purpose: Skin moisturizer

Warnings

Warnings
1. Stop use immediately if the following symptoms occur, and consult a dermatologist.
A) If use of the product causes red spots, swelling, itching, or other irritation
B) If direct sunlight causes the symptoms listed above
2. Do not use on areas of injury, rash, or skin inflammation.
3. Keep out of eyes when using the products. Immediately rinse with water in case of eye contact.
4. Storage and safety precautions
A) Close lid tightly when not in use.
B) Keep out of reach of children and infants.
C)Do not store in high/low temperatures, and keep away from sunlight.
5. Try patch test on small area of skin before use or when switching to a new product.
6. Do not place product contents back into container after use to avoid deterioration of quality.
7. This is a cosmetic product not intended to be used by children.

Keep out of reach of children

Keep out of reach of children:
Keep away from children and infants.
This is a cosmetic product not intended for consumption.

DOSAGE AND ADMINISTRATION

Recommended amount per use: 1-2g

INDICATIONS AND USAGE

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Use several times a day after taking a bath or washing face.
Apply an appropriate amount and massage gently onto skin for better absorption.
Be thorough for dried skin areas such as the face, neck, inner arm, knees, and calves to offer better protection for sensitive, dried skin.
Use both Atoby Medi Lotion and Atoby Medi Cream for maximum effectiveness in areas of severe dryness and in sensitive/troubled areas.

PACKAGE LABEL. PRINCIPAL DISPLAY PANEL

HAMSOA ATOBY MEDI

witch hazel lotion

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50964-040
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
WITCH HAZEL (UNII: 101I4J0U34) (WITCH HAZEL - UNII:101I4J0U34)	WITCH HAZEL	0.15 g in 150 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
SORBITOL (UNII: 506T60A25R)	
LAVENDER OIL (UNII: ZBP1YXW0H8)	
BASIL (UNII: 2U0KZP0FDW)	
GLYCERIN (UNII: PDC6A3C0OX)	
CETYL ETHYLHEXANOATE (UNII: 134647WMX4)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50964-040-01	150 mL in 1 CARTON		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part347	03/01/2012	

Labeler - HAMSOA PHARMACEUTICAL CO., LTD. (690417753)

Registrant - HAMSOA PHARMACEUTICAL CO., LTD. (690417753)

Establishment

Name	Address	ID/FEI	Business Operations
HAMSOA PHARMACEUTICAL CO., LTD.		690417753	manufacture(50964-040)

Revised: 8/2012 HAMSOA PHARMACEUTICAL CO., LTD.