BY PHARMICELL LAB MOISTURIZING FACE TONING MIST- allantoin spray Pharmicell 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 : ALLANTOIN 0.5%

INACTIVE INGREDIENT

Inactive ingredient : WATER(GLACIER WATER), WATER, BUTYLENE GLYCOL, GLYCERIN, HUMAN BONE MARROW STEM CELL CONDITIONED MEDIA, PPG-26-BUTETH-26, PEG-40 HYDROGENATED CASTOR OIL, 1,2-HEXANEDIOL, PHENOXYETHANOL, CITRUS PARADISI (GRAPEFRUIT) FRUIT EXTRACT, GANODERMA LUCIDUM (MUSHROOM) EXTRACT, GRIFOLA FRONDOSA EXTRACT, INONOTUS OBLIQUUS (MUSHROOM) EXTRACT, SPARASSIS CRISPA EXTRACT, PHELLINUS LINTEUS EXTRACT, POLYGONUM MULTIFLORUM ROOT EXTRACT, ACANTHOPANAX SENTICOSUS (ELEUTHERO) ROOT EXTRACT, PORTULACA OLERACEA EXTRACT, PANAX GINSENG ROOT EXTRACT, ASPARAGUS COCHINCHINENSIS ROOT EXTRACT, LAVANDULA ANGUSTIFOLIA (LAVENDER) OIL, MELALEUCA ALTERNIFOLIA (TEA TREE) LEAF OIL, EUCALYPTUS GLOBULUS LEAF OIL, GERANIUM MACULATUM OIL, ROSMARINUS OFFICINALIS (ROSEMARY) LEAF OIL, MENTHA PIPERITA (PEPPERMINT) OIL, TRISODIUM EDTA

PURPOSE

Purpose : Skin Protectant

WARNINGS

Warning : Keep out of reach of children and babies. For external use only. Avoid contact with eyes. Discontinue use if signs of irritation and/or rash appear.

KEEP OUT OF REACH OF CHILDREN

Keep out of reach of children and babies.

INDICATIONS AND USAGE

Indications and usage : Before and after makeup, spray the mist frequently $10\sim20$ cm away from the face in a dry indoor condition. This product can be used as an alternative to the toner, and shaking the product before use is not necessary as it has a compressed injection form.

DOSAGE AND ADMINISTRATION

Dosage and administration : Before and after makeup, spray the mist frequently 10~20 cm away from the face in a dry indoor condition.

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL



BY PHARMICELL LAB MOISTURIZING FACE TONING MIST

allantoin spray

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Product Information									
Product Type		HUMAN OTC DRUG	Item Co	ode (Source)	Ν	NDC:60949-100			
Route of Administrati	on	TOPICAL							
Active Ingredient/	Active Moi	etv							
Ingredient Name Basis of Streng						Strength			
ALLANTOIN (UNII: 344S277G0Z) (ALLANTOIN - UNII:344S277G0Z)			Z)	ALLANTOIN					
Inactive Ingredients									
		Strength							
WATER (UNII: 059QF0K									
BUTYLENE GLYCOL (
Packaging									
# Item Code	Pac	kage Description	Marketing Start Date		Ma	Marketing End Date			
1 NDC:60949-100-01	120 mL ii	n 1 CARTON							
Marketing Information									
Marketing Category	Application Number or Monograph Citation			Marketing Star	t Date	Marketing End Date			
OTC monograph final	part347	347 1		10/01/2013					

Labeler - Pharmicell Co., Ltd. (687744110)

Registrant - Pharmicell Co., Ltd. (687744110)

Establishment

Name	Address	ID/FEI	Business Operations
Pharmicell Co., Ltd.		687744110	manufacture(60949-100)

Revised: 7/2013

Pharmicell Co., Ltd.