# BY PHARMICELL LAB LUXURY CELL PERFORMANCE TONER- allantoin cream 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.

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#### ACTIVE INGREDIENT

Active ingredient: ALLANTOIN 0.5%

## INACTIVE INGREDIENT

Inactive ingredient: WATER, GLYCERIN, ALCOHOL, HUMAN BONE MARROW STEM CELL CONDITIONED MEDIA, DIPROPYLENE GLYCOL, CITRUS PARADISI (GRAPEFRUIT) FRUIT EXTRACT, BETAINE, TREHALOSE, METHYL GLUCETH-20, POLYGONUM MULTIFLORUM ROOT EXTRACT, ACANTHOPANAX SENTICOSUS (ELEUTHERO) ROOT EXTRACT, PORTULACA OLERACEA EXTRACT, PANAX GINSENG ROOT EXTRACT, ASPARAGUS COCHINCHINENSIS ROOT EXTRACT, 1,2-HEXANEDIOL, ACRYLATES/C10-30 ALKYL ACRYLATE CROSSPOLYMER, PHENOXYETHANOL, XANTHAN GUM, ALTHAEA ROSEA ROOT EXTRACT, ALOE BARBADENSIS LEAF EXTRACT, GANODERMA LUCIDUM (MUSHROOM) EXTRACT, GRIFOLA FRONDOSA EXTRACT, INONOTUS OBLIQUUS (MUSHROOM) EXTRACT, SPARASSIS CRISPA EXTRACT, PHELLINUS LINTEUS EXTRACT, PEG-40 HYDROGENATED CASTOR OIL, POTASSIUM HYDROXIDE, ADENOSINE, SODIUM HYALURONATE, 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

#### **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: After wash up, take a proper amount of toner, and spread it evenly over the entire face using a cotton pad for absorption.

#### DOSAGE AND ADMINISTRATION

Dosage and administration: Take a proper amount of the product and spread it evenly over the entire face.

## PACKAGE LABEL.PRINCIPAL DISPLAY PANEL





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allantoin cream

# **Product Information**

HUMAN OTC DRUG NDC:60949-090 Product Type Item Code (Source)

**Route of Administration** TOPICAL

# **Active Ingredient/Active Moiety**

**Ingredient Name Basis of Strength** Strength

ALLANTO IN (UNII: 344S277G0Z) (ALLANTO IN - UNII: 344S277G0Z) ALLANTOIN 0.65 mg in 130 mL

# **Inactive Ingredients**

Ingredient Name	Strength	
WATER (UNIII) 0500 F0 KOOD)		

WATER (UNII: 059QF0KO0R) GLYCERIN (UNII: PDC6A3C0OX)

# **Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:60949-090-01	130 mL in 1 CARTON		

# **Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part347	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(60949-090)			

Revised: 7/2013 Pharmicell Co., Ltd.