

BY PHARMICELL LAB THE PRESTIGE ESSENCE- allantoin cream
Pharmacell 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, HUMAN BONE MARROW STEM CELL CONDITIONED MEDIA, GLYCERIN, DIPROPYLENE GLYCOL, HYDROGENATED POLYDECENE, NEOPENTYL GLYCOL DICAPRATE, CYCLOPENTASILOXANE, CYCLOHEXASILOXANE, CAPRYLIC/CAPRIC TRIGLYCERIDE, CETEARYL ALCOHOL, GLYCERYL STEARATE SE, HYDROGENATED LECITHIN, CITRUS PARADISI (GRAPEFRUIT) FRUIT EXTRACT, BEESWAX, POLYGONUM MULTIFLORUM ROOT EXTRACT, ACANTHOPANAX SENTICOSUS (ELEUTHERO) ROOT EXTRACT, PORTULACA OLERACEA EXTRACT, PANAX GINSENG ROOT EXTRACT, ASPARAGUS COCHINCHINENSIS ROOT EXTRACT, HYDROGENATED LECITHIN, TOCOPHERYL ACETATE, CARBOMER, PHENOXYETHANOL, 1,2-HEXANEDIOL, XANTHAN GUM, POTASSIUM HYDROXIDE, LITHOSPERMUM ERYTHRORHIZON ROOT OIL, DIPOTASSIUM GLYCYRRHIZATE, ALTHAEA ROSEA ROOT EXTRACT, ALOE BARBADENSIS LEAF EXTRACT, CLADOSIPHON OKAMURANUS EXTRACT, CERAMIDE 3, NEOFINETIA FALCATA CALLUS CULTURE EXTRACT, ZANTHOXYLUM PIPERITUM FRUIT EXTRACT, PULSATILLA KOREANA EXTRACT, USNEA BARBATA (LICHEN) EXTRACT, LECITHIN, ADENOSINE, SODIUM HYALURONATE, DEXTRIN, 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

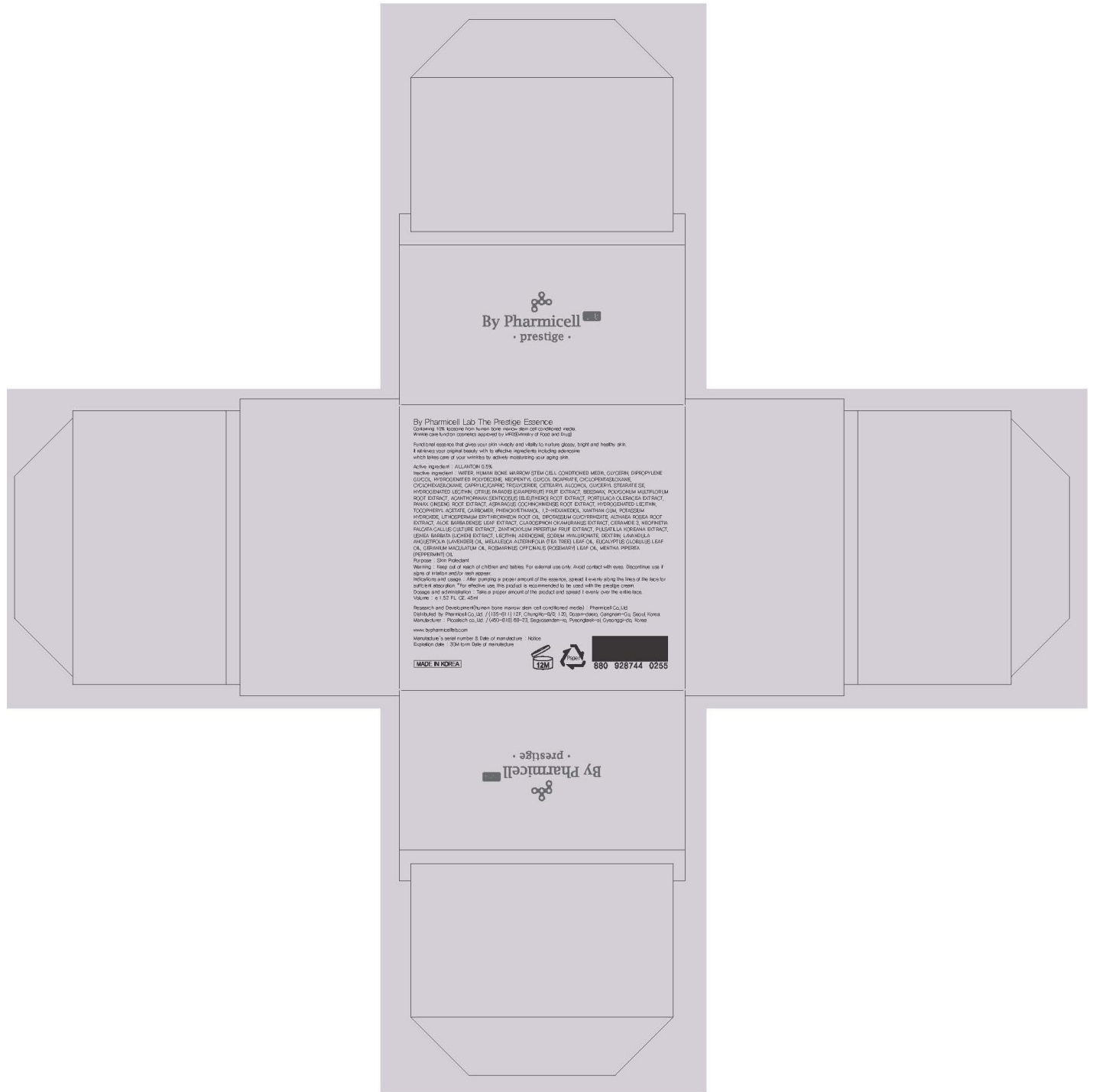
INDICATIONS AND USAGE

Indications and usage : After pumping a proper amount of the essence, spread it evenly along the lines of the face for sufficient 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			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:60949-020
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALLANTOIN (UNII: 344S277G0Z) (ALLANTOIN - UNII:344S277G0Z)	ALLANTOIN	0.22 mg in 45 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:60949-020-01	45 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(60949-020)

Revised: 11/2013

Pharmicell Co., Ltd.