



Product Licence Licence de mise en marché

Product Number/Numéro de produit: 80052293

Brand Name/Marque nominative: Dermakalm Scalp Psoriasis Gel

Issued to/Émise à:

Name of licensee/Nom du titulaire:

Paladin Labs Inc.
100 Boulevard Alexis Nihon, Bureau 600
Montreal, Quebec, H4M 2P2
Canada

Authorized for the following/Autorisé pour ce qui suit:

Dosage form/Forme posologique: Gel

Recommended route of administration/Voie d'administration recommandée:

Topical

Recommended dose/Dose recommandée:

Adults : Apply a thin layer of Dermakalm Scalp Psoriasis Gel onto the scalp by separating the hair. Let it dry and do not wash out. Apply 2-3 times a day; re-apply after washing your hair. Use a gentle shampoo. During the first 2-3 days of application, gently dilute a small quantity of gel with lukewarm water and massage onto the wet skin of the affected area.

Recommended duration of use/Durée d'utilisation recommandée:

For use beyond 6 weeks, consult a health care practitioner.

Recommended use or purpose/Usage ou les fins recommandés:

Helps to temporarily decrease the severity of psoriasis symptoms of the scalp.

Risk Information/Renseignements sur les risques:

Cautions and Warnings

For external use only. If symptoms persist or worsen, consult a health care practitioner. If skin irritation develops or increases, discontinue use. If irritation persists or worsens, consult a health care practitioner. If symptoms persist or worsen, consult a health care practitioner. If you are pregnant or breastfeeding, consult a health care practitioner prior to use. Avoid contact with eyes and mucous membranes; if this happens, rinse thoroughly with water.

Known Adverse Reactions

This product might produce a slight and brief burning sensation upon application to the skin lesion that may resolve with continuous usage.

Medicinal Ingredients/Ingrédients médicinaux:

Proper Name Nom propre	Common Name Nom usuel	Quantity per Dosage Unit Quantité par unité posologique	Extract Extrait	Potency Activité	Source Material Matière d'origine
Clinoptilolite	Clinoptilolite	1 % (w/w)	N/A	N/A	Clinoptilolite (Synthetic)

This licence is issued by the Minister of Health under the authority of section 7 of the Natural Health Products Regulations. Sale of the described natural health product, including any changes thereto pursuant to section 11 of the Regulations, is subject to the Food and Drugs Act and to the Natural Health Products Regulations.

Cette licence est émise par la ministre de la Santé en vertu de l'article 7 du Règlement sur les produits de santé naturels.

La vente du produit de santé naturel décrit dans la présente, y compris toute modification afférente au sens de l'article 11 du Règlement, est assujettie à la Loi sur les aliments et drogues et au Règlement sur les produits de santé naturels.

Issued/émis le: 2014-07-08

Revised/Amended/Modifié le: 2016-01-05



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***A/ Director General/ Int. Directeur général
NHPD/DPSN***