Health Canada Santé Canada

Product Licence
Licence de mise en marché

Product Number/*Numéro de produit*: 80071586
Brand Name/*Marque nominative*: ZEOLITE - 4 (Tablet)
Issued to/*Émise à:*
Name of licensee/*Nom du titulaire:*

Mineral Medix Corp.
58 Albright Crescent
Richmond Hill, Ontario, L4E 4Z4
Canada

Authorized for the following/*Autorisé pour ce qui suit:*

Dosage form/*Forme posologique*: Tablet
Recommended route of administration/Voie *d'administration recommandée:*

Oral

Recommended dose/*Dose recommandée:*

Adults : Take one tablet two times each day with full glass of water.
Do not take with other medication.
Allow 1-1.5 hours before or after any medication.
Store in tightly sealed container in dry place with temperatures between -30C and +30C

Recommended duration of use/*Durée d'utilisation recommandée:*

Consult a health care practitioner for use beyond 30 days.

Recommended use or purpose/*Usage ou les fins recommandés:*

Helps to support detoxification.

Risk Information/*Renseignements sur les risques:*

Cautions and Warnings

Keep out of reach of children.
If you are pregnant or breastfeeding, consult a health care practitioner prior to use.
Individuals with kidney disease should not take this product except on the advice of a doctor.
This product is not intended for the treatment of heavy metal poisoning. If exposure to heavy metals is known or suspected, call 911 or call your local Poison Control Center.
If you are taking medication to have a medical condition, consult your health care practitioner prior to use.

Contra-Indications

Do not use if you have gastric or duodenum ulcer.

Known Adverse Reactions

Gastrointestinal discomfort has been known to occur, in which case, reduce dose.

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	.2 g	N/A	N/A	Clinoptilolite

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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: 2016-08-02	Revised/Amended/Modifié le: N/A

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