Australian Government

Department of Health

Therapeutic Goods Administration

Public Summary

Summary for ARTG Entry: 276085 Methyl BioActive

ARTG entry for Medicine Listed

Sponsor FIT-BioCeuticals Limited

Postal Address Care of Blackmores Ltd PO Box 1725, Warriewood, NSW, 2102

Australia

ARTG Start Date 1/06/2016

Product Category Medicine
Status Active

Approval Area Listed Medicines

Conditions

The sponsor shall keep records relating to this listed medicine as are necessary to: (a) Expedite recall if necessary of any batch of the listed medicine, (b) Identify the manufacturer(s) of each batch of the listed medicine. Where any part of or step in manufacture in Australia of the listed medicine is sub-contracted to a third party who is not the sponsor, copies of relevant Good Manufacturing Practice agreements relation to such manufacture shall be kept.

The sponsor shall retain records of the distribution of the listed medicine for a period of five years and shall provide the records or copies of the records to the Complementary Medicines Branch, Therapeutic Goods Administration, upon request.

Where a listed medicine is distributed overseas as well as in Australia, product recall or any other regulatory action taken in relation to the medicine outside Australia which has or may have relevance to the quality, safety or efficacy of the goods distributed in Australia, must be notified to the National Manager Therapeutic Goods Administration, immediately the action or information is known to the sponsor.

Products

1. Methyl BioActive

Product Type Single Medicine Product Effective Date 14/04/2020

Permitted Indications

Maintain/support energy levels

Maintain/support energy production

Aid/assist healthy red blood cell production

Helps maintain/support haemoglobin formation/synthesis

Helps decrease/reduce homocysteine levels

Aid/assist/helps synthesis of neurotransmitters

Maintain/support nervous system function

Maintains/support healthy foetal development

Indication Requirements

If product is indicated for weight loss, label statement: When used in conjunction with a program of reduced intake of dietary calories and increased physical activity.

Product presentation must not imply or refer to serious cardiovascular conditions.

Product presentation must not imply or refer to mental illnesses, disorders or conditions.

Label statement: Advise your doctor of any medicine you take during pregnancy, particularly in your first trimester.

Product presentation must not imply or refer to chronic fatigue syndrome.

Standard Indications

No Standard Indications included on Record

Specific Indications

No Specific Indications included on Record

Warnings

No Warnings included on Record

Additional Product information

Pack Size/Poison information

Pack Size Poison Schedule

Components

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1 . Formulation 1

Dosage Form Tablet, film coated

Route of Administration Oral

Visual Identification

Active Ingredients

 choline bitartrate
 300 mg

 levomefolate calcium
 542 microgram

 Equivalent: levomefolic acid
 500 microgram

mecobalamin (co-methylcobalamin)

pyridoxal 5-phosphate monohydrate

Equivalent: pyridoxine

riboflavin sodium phosphate

Equivalent: riboflavin

25 mg

Other Ingredients (Excipients)

calcium hydrogen phosphate dihydrate

Carnauba Wax

silicon dioxide

serine

colloidal anhydrous silica

crospovidone
hypromellose
macrogol 400
magnesium stearate
microcrystalline cellulose
potable water
povidone
purified water

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50 mg