

PROFESSIONAL INFORMATION

This unregistered medicine has not been evaluated by SAHPRA for its quality, safety or intended use.

SCHEDULING STATUS: S0

1. NAME OF THE MEDICINE: MG/K ASPARTATE

2. QUALITATIVE AND QUANTITATIVE COMPOSITION

Serving Size	2 tablets	
Servings Per Container: 60 pack size		30
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Magnesium (as magnesium aspartate)		100 mg
Potassium (as potassium aspartate)		160 mg

Other Ingredients: Microcrystalline cellulose, cellulose, stearic acid (vegetable), and croscarmellose sodium.

For full list of excipients, see section 6.1

This product is non-GMO, gluten-free, and vegetarian.

3. PHARMACEUTICAL FORM

Off-white uncoated tablet

4. CLINICAL PARTICULARS

4.1 Therapeutic indications

Mg/K Aspartate features magnesium and potassium in the form of true mineral aspartates to enhance absorption. Magnesium plays an important role in energy production, and potassium is a key electrolyte in cellular function. *

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4.2 Posology and method of administration

Take two tablets daily or as directed by your healthcare practitioner.

4.3 Contraindications

None known.

4.4 Special warnings and precautions for use

Keep out of the reach of children.

4.5 Interaction with other medicines and other forms of interaction

None known.

4.6 Fertility, pregnancy and lactation

There are no adequate data from the use of **Mg/K Aspartate** in pregnant or breast-feeding woman

4.7 Effects on ability to drive and use machines

There are no adequate data from the use of **Mg/K Aspartate** on the ability to drive and use machines.

4.8 Undesirable effects

None known.

4.9 Overdose

Treatment is symptomatic and supportive.

5. PHARMACOLOGICAL PROPERTIES

5.1 Pharmacodynamic properties

Category of medicine: D (Complementary Medicine)

Class of medicine: 34.13 Health supplements - Other

5.2 Pharmacokinetic properties

Pharmacokinetic studies have not been conducted on **Mg/K Aspartate** product

6. PHARMACEUTICAL PARTICULARS

6.1 List of excipients

Microcrystalline cellulose
Cellulose
Stearic acid (vegetable)
Croscarmellose sodium

6.2 Incompatibilities

None known.

6.3 Shelf Life

24 Months.

6.4 Special precautions for storage

Keep tightly closed in a cool, dry place
Keep tablets in original container until required for use.
KEEP OUT OF REACH OF CHILDREN.

6.5 Nature and contents of container

Packs of 60 tablets in amber glass bottles.
Pack-size: 60 tablets

6.6 Special precautions for disposal

No special requirements.

7. HOLDER OF CERTIFICATE OF REGISTRATION

Amipro Advanced Development Products (pty) Ltd
Unit 3 Eastgate Business Park, 1 South Road, Marlboro, Sandton, Johannesburg 2063
Tel: 011 802 8101

8. REGISTRATION NUMBER

Will be allocated by SAHPRA upon registration.

9. DATE OF FIRST AUTHORISATION

Will be allocated by SAHPRA upon registration.