

SUMMARY OF PRODUCT CHARACTERISTICS

1 NAME OF THE MEDICINAL PRODUCT

Magnesium Sulphate Injection 50%w/v

2 QUALITATIVE AND QUANTITATIVE COMPOSITION

Magnesium Sulfate	5g/	2.5g/	1g/
Heptahydrate	10ml	5ml	2m
			1

For full list of excipients, see section 6.1

3 PHARMACEUTICAL FORM

Sterile Solution for Injection

A clear, colourless solution practically free from visible particles

4 CLINICAL PARTICULARS

4.1 Therapeutic indications

- Treatment of Magnesium deficiency in hypomagnesaemia where the oral route of administration may be inappropriate.
- To prevent further seizures associated with eclampsia.

4.2 Posology and method of administration

Dosage should be individualised according to patient's needs and responses. Plasma levels should also be monitored throughout therapy.

a) Treatment of magnesium deficiency in hypomagnesaemia:

For intravenous administration, a concentration of 20% or less should be used; the rate of injection not exceeding 1.5ml/minute of a 10% solution or its equivalent.

Up to 40g MgSO₄ (equivalent to 160mmol Mg²⁺) by slow intravenous infusion (in glucose 5%) over up to 5 days, may be required to replace the deficit (allowing for urinary losses).

Mild magnesium deficiency

1g intramuscularly every 6 hours for 4 doses.

Severe magnesium deficiency

Up to 250mg/kg intramuscularly given within a period of 4 hours or 5g/litre of infusion solution intravenously over 3 hours

Paediatric population

It is recommended that the solution be diluted to 20% w/v prior to intramuscular injection

Elderly

No special recommendation except in renal impairment, see below

Renal impairment:

Dosage should be reduced in renal impairment. Plasma magnesium concentrations should be monitored throughout therapy

b) To prevent further seizures associated with eclampsia:

An initial intravenous (IV) loading dose is followed for 24h by either an IV infusion, or regular intramuscular (IM) injections.

Intramuscular Maintenance Regimen

A loading dose of 4g MgSO_4 (approx. 16mmol Mg^{2+}) IV (usually in 20% solution) over 5min (minimum, preferably 10-15 min) is followed immediately by 5g MgSO_4 (approx. 20mmol Mg^{2+}) (usually in 50% solution) as a deep IM injection into the upper outer quadrant of each buttock.

Maintenance therapy is a further 5g MgSO_4 (approx. 20mmol Mg^{2+}) IM every 4h, continued for 24h after the last fit (provided the respiratory rate is $\geq 16/\text{min}$, urine output $\geq 25\text{ml}/\text{h}$, and knee jerks are present).

Intravenous Maintenance Regimen

A loading dose of 4g MgSO_4 (approx. 16mmol Mg^{2+}) IV (or in some cases 5g MgSO_4 (approx. 20mmol Mg^{2+}) IV), as described above, is followed by an infusion of 1g/h continued for 24h after the last fit.

Recurrent Convulsions: In both the IM and IV regimens, if convulsions recur, a further 2-4g MgSO_4 (approx. 8 - 16mmol Mg^{2+}) (depending on the woman's weight, 2g MgSO_4 (approx. 8mmol Mg^{2+}) if less than 70Kg) is given IV over 5 min.

**** The Eclampsia Trial Collaborative Group (Duley L et al) (1995) Which anticonvulsant for women with eclampsia? Evidence from the Collaborative Eclampsia Trial., The Lancet, Vol. 345, pp. 1455-1463.***

Appropriate reductions in dosage should be made for patients

with renal impairment; a suggested dose reduction in severe renal impairment is a maximum of 20g MgSO₄ (approx. 80mmol Mg²⁺) over 48 hours.

Method of administration

Magnesium sulfate injection may be administered by intramuscular or intravenous routes.

Intramuscular therapy should be used only when peripheral venous access is impossible.

4.3 **Contraindications**

Hypersensitivity to magnesium and its salts or to any of the excipients listed in section 6.1.

Magnesium sulfate is contraindicated in patients with severely impaired renal function.

4.4 **Special warnings and precautions for use**

Magnesium sulfate must be used with caution in patients suspected of or known to have renal impairment.

Magnesium sulfate should not be used in hepatic coma if there is a risk of renal failure.

Parenteral magnesium salts should be used with caution in patients with myasthenia gravis.

Serum calcium levels should be routinely monitored in patients receiving magnesium sulfate.

4.5 **Interaction with other medicinal products and other forms of interaction**

Administer with caution to patients receiving digitalis glycosides. Magnesium sulfate should not be administered concomitantly with high doses of barbiturates, opioids or hypnotics due to the risk of respiratory depression

The action of non-depolarising muscle relaxants such as tubocurarine is potentiated and prolonged by parenteral magnesium salts.

Concomitant use of calcium channel blockers such as nifedipine or nimodipine may rarely lead to a calcium ion imbalance and could result in abnormal muscle function.

The neuromuscular blocking effects of parenteral magnesium and aminoglycoside antibacterials may be additive.

4.6 **Fertility, pregnancy and lactation**

Pregnancy

Safety in human pregnancy has not been established, however, in the medical emergency of a patient having Eclampsia, Magnesium Sulfate can be administered to relieve this condition, which may be life threatening to both mother and baby.

Magnesium crosses the placenta. When used in pregnant women, foetal heart rate should be monitored and use within 2 hours of delivery should be avoided.

Magnesium sulfate can cause skeletal adverse effects when administered continuously for more than 5 to 7 days to pregnant women. There are retrospective epidemiological studies and case reports documenting fetal adverse effects including hypocalcaemia, skeletal demineralization, osteopenia and other skeletal adverse effects with maternal administration of magnesium sulfate for more than 5 to 7 days. The clinical significance of the observed effects is unknown.

If prolonged or repeated exposure to magnesium sulfate occurs during pregnancy monitoring of neonates for abnormal calcium or magnesium levels and skeletal adverse effects should be considered.

Breast-feeding

As with all drugs it is not advisable to administer magnesium sulfate during pregnancy or breastfeeding unless considered essential, and it must be administered under medical supervision.

Fertility

No studies and/or data are available on the effects on fertility.

4.7 Effects on ability to drive and use machines

No studies on the effects on the ability to drive and use machines have been performed.

4.8 Undesirable effects

Metabolism and nutrition disorders

Electrolyte/fluid abnormalities (hypophosphataemia, hypertonic dehydration) Hypersensitivity reactions.

Hypocalcaemia.

Pain or burning at the injection site following IV/IM administration.

Hypermagnesaemia characterised by flushing, sweating, thirst, hypotension, drowsiness, dizziness, headache, risk of itching and tingling, nausea, vomiting, confusion, slurred speech, double vision, loss of tendon reflexes due to neuromuscular blockade, muscle weakness, respiratory depression, electrolyte/fluid abnormalities (hypophosphataemia, hyperosmolar dehydration), ECG changes (prolonged PR, QRS and QT intervals), bradycardia, tachycardia cardiac arrhythmias, coma and cardiac arrest.

There have been isolated reports of maternal and fetal hypocalcaemia with high doses of magnesium sulfate (see section 4.6).

Reporting of suspected adverse reactions

Reporting suspected adverse reactions after Authorisation of the medicinal product is important. It allows continued monitoring of the benefit/risk balance of the medicinal product. Healthcare professionals are asked to report any suspected adverse reactions via Pharmacovigilance Electronic Reporting System (PvERS) <https://pv.pharmacyboardkenya.org/>

4.9 Overdose

Appropriate action should be taken to reduce the blood level of magnesium to avoid hypermagnesaemia. Neuromuscular blockade associated with hypermagnesaemia may be reversed with calcium salts, such as Calcium Gluconate, which should be administered intravenously in a dose equivalent to 2.5 to 5mmol of calcium.

5 PHARMACOLOGICAL PROPERTIES

5.1 Pharmacodynamic properties

Pharmacotherapeutic group: Mineral Supplements, ATC code: A12CC 02.

Magnesium is the second most abundant cation in intracellular fluid and is an essential body electrolyte. Magnesium is a factor in a number of enzyme systems and is involved in neurochemical transmission and muscular excitability. Parenterally administered magnesium sulfate exerts a depressant effect on the central nervous system and acts peripherally to produce vasodilation.

5.2 Pharmacokinetic properties

Absorption

When magnesium is administered intravenously, the onset of action is immediate and last for approximately 30 minutes. Following intramuscular administration, the onset of action occurs within 1 hour and the duration of action is 3 to 4 hours. Upon intramuscular administration, the plasma concentration of magnesium sulfate shows a slow increase that reaches a plateau within 1 to 2 hours and is followed by a slow decline back to baseline within the next 6 to 8 hours. At the end of 4 hours, after another intramuscular injection of magnesium sulfate, the plasma concentration remains constant.

Distribution

Plasma protein binding of injected magnesium is comparable to

endogenous magnesium. About 40% to 50% of plasma magnesium is protein bound and not ultrafiltrable.

The amount of ionised magnesium (the active form of magnesium according to both in vivo and in vitro studies) rose proportionately to the total serum magnesium concentration. In a pharmacokinetic study in women with preeclampsia, demonstrated lack of correlation between total magnesium and ionised magnesium levels in both physiologic and hypermagnesmic (pharmacologic) ranges.

Several attempts have been made to estimate the apparent volume of distribution (Vd) of magnesium. Unlike non-pregnant healthy volunteers, where Vd reaches constant values within 2 hours after administration, the apparent Vd in pregnant women

becomes constant within 3 to 4 hours and ranges from 0.250 to 0.442 L/kg. Another study estimated the Vd in the range of 0.370 to 0.430 L/kg 24 hours after administration. The pharmacokinetic profile of magnesium sulfate after intravenous administration can be described by a 2-compartment model with a rapid distribution

(a) phase, followed by a relative slow beta phase of elimination, where central compartment (Vc) and terminal phase (V β) volumes were 0.250 ± 0.010 L/kg and 0.570 ± 0.022 L/kg, respectively.

The unbound magnesium ions diffuse into the extravascular-extracellular space, bones, cross the placenta and are rapidly taken up by foetal tissues and thus magnesium in amniotic fluid, the foetus and in neonates of mothers treated with magnesium sulfate show increased concentrations as compared to untreated mothers.

When the mother receives 1 to 2 g/h magnesium infusions, foetal serum magnesium concentrations are twice as high as control concentrations, the highest concentrations observed in umbilical and venous and arterial blood. Foetal magnesium plasma concentrations equalise with the maternal within 2 hours, while the increase in amniotic fluid occurs more slowly. The mean baseline cerebrospinal fluid (CSF) magnesium level in preeclamptic women was 1.1 ± 0.05 mmol/L.

Intrapartum magnesium sulfate treatment increased breast milk/colostrum magnesium levels significantly only for 24 hours. After 24 hours magnesium levels in breast milk comes back to normal. Breast milk/colostrum calcium levels were not affected by magnesium sulfate therapy.

Biotransformation and Elimination

Magnesium sulfate is not metabolised and is excreted solely by the kidney.

Urinary excretion is very rapid with a 20-fold increase during magnesium sulfate infusion. About 38 to 53% of the total injected magnesium is excreted 4 hours after the end of the infusion and >90% are eliminated within 24 hours after the

infusion.

In patients with normal renal function, the increase in magnesium clearance is linear to the serum magnesium concentrations and the half-life of magnesium is 4 hours. Half-life increases with decrease in glomerular filtration rate. Urinary calcium concentration increases 4.5-fold during infusion of magnesium sulfate and there is a 3-fold increase in urinary calcium excretion rate which is likely due to competition for common reabsorption sites.

5.3 Preclinical safety data

This product has been available for many years, and its side effects and clinical profile are well-understood, therefore no further data is provided.

6 PHARMACEUTICAL PARTICULARS

6.1 List of excipients

Water for
Injections
Hydrochloric
Acid

Sodium Hydroxide

6.2 **Incompatibilities**

Streptomycin sulfate and tetramycin sulfate activity is inhibited by magnesium ions

6.3 **Shelf life**

3 years

Discard any unused solution immediately after first use.

From a microbiological point of view, the product should be used immediately. If not used immediately, in-use storage time and conditions prior to use are the responsibility of the user and would normally not be longer than 24 hours at 2 to 8°C, unless dilution has taken place in controlled and validated aseptic conditions.

6.4 **Special precautions for storage**

Do not store above 25°C.

6.5 **Nature and contents of container**

Neutral Type 1 glass ampoules 2, 5 and 10ml, containing a 50% w/v sterile solution for injection of Magnesium Sulfate.

Not all pack sizes may be marketed.

6.6 **Special precautions for disposal**

For intramuscular use a 25% or 50% solution is used. For intravenous use this solution must be diluted before use. Concentrations of up to 20% are usually employed.

Discard any unused solution at the end of the session in the appropriate manner.

7 **MARKETING AUTHORISATION HOLDER**

Aurum Pharmaceuticals Ltd
Bampton Road
Harold Hill

Romford
Essex
RM3
8UG
United Kingdom

8 MARKETING AUTHORISATION NUMBER(S)

H2006/17754/R1

**9 DATE OF FIRST AUTHORISATION/RENEWAL OF THE
AUTHORISATION**

26/02/2026

10 DATE OF REVISION OF THE TEXT

24/08/2026