

Magnesium supplementation to prevent recurrence of renal stones

Ivo Laranjinha, Patricia Matias, Jorge Dickson

Nephrology Department, Hospital de Santa Cruz/Centro Hospitalar de Lisboa Ocidental

ABSTRACT

Stone formers have lower urinary magnesium than healthy people. Higher urinary magnesium levels are associated with lower incidence of kidney stones, and hypomagnesuria has been described as a lithogenic risk factor. Magnesium can have direct and indirect inhibitory effects on lithogenesis: decreasing the absorption of oxalates in the intestine; forming magnesium oxalate complexes which reduces the saturation of calcium oxalate; increasing the urinary citrate and inhibiting the conversion of the calcium oxalate in its monohydrated form. Oral supplementation with magnesium is an effective way to correct hypomagnesuria. However, oral magnesium supplementation in recurrent stone formers with hypomagnesuria is still a subject of ongoing debate, and physicians treating these patients underestimate its potential preventive effects. Oral magnesium supplementation can be used as an adjuvant therapy to the standard prophylactic therapy, mainly in association with an alkali salt. It is well tolerated and has few adverse effects.

Keywords: Nephrolithiasis; Recurrent Stone Formers; Hypomagnesuria; Magnesium supplementation.

INTRODUCTION

Patients with recurrent stone disease should perform a complete metabolic evaluation. The modifiable lithogenic factors identified in the 24-hour urine analysis will guide therapeutic recommendations to prevent future stone episodes^{1,2}. Apart from the efforts to lower calcium (Ca) and oxalate and to increase citrate in the urine, we should look beyond these widely known lithogenic factors.

Low urinary magnesium (UMg) is a quite common finding among recurrent stone formers³. In the literature, the prevalence of hypomagnesuria among stone formers varies between 4.3 and 11%⁴⁻⁸, but in some studies it can reach one third⁹. This wide range of hypomagnesuria rate found in the literature is related to the different definitions of hypomagnesuria used.

When compared with healthy controls, stone formers have lower UMg excretion^{4,10,11}. Epidemiologic studies have found an inverse association between an increased magnesium (Mg) intake and the risk of symptomatic kidney stones¹².

Healthy children have higher UMg concentrations than healthy adults – mean UMg in adults is 90-104 mg/day and in children is between 120 and 150 mg/day¹³⁻¹⁵. The higher UMg levels in children's urine has been presented as a justification for the lower prevalence of nephrolithiasis in childhood¹⁶.

Since UMg is a reasonable surrogate for dietary Mg intake¹⁷, many physicians have hypothesized that increasing Mg intake could increase UMg and consequently reduce the risk of stone recurrence.

Several studies show that Mg ingestion in the diet is below the recommended levels – it is estimated that 50-60% of adults do not

reach the average dietary intake^{18,19}. A growing body of evidence from epidemiologic and clinical studies has confirmed the association of magnesium deficiency with several diseases such as electrolyte, neurologic, musculoskeletal and inflammatory disorders; cardiovascular diseases and diabetes^{15,20,21}.

Increasing dietary Mg intake can be an effective way to correct a degree of Mg deficiency; however most of the dietary sources of Mg also contain important amounts of oxalate, precluding the recommendation of a diet rich in Mg to all stone formers²².

However, oral Mg supplementation to prevent renal stone recurrence is still a subject of ongoing debate, and its potential preventive effects are not proven.

This review summarizes our current understanding of Mg physiology, the risks and benefits of supplementing stone formers with Mg and our suggestions for correcting hypomagnesuria in recurrent stone formers.

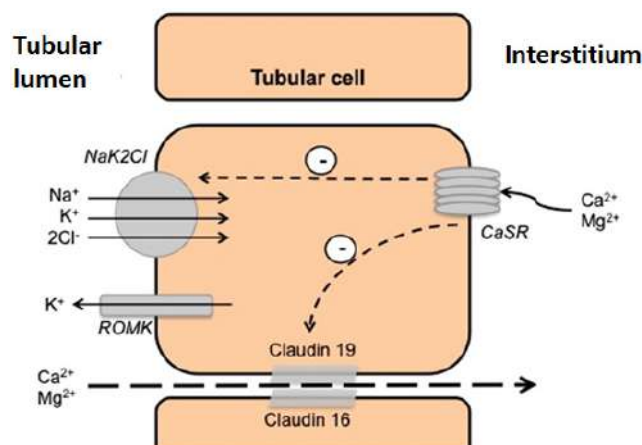
PHYSIOLOGY OF MAGNESIUM

Magnesium is mainly eliminated by the kidneys. Although 80% of the serum Mg is filtered at the glomerulus, less than 5% of it is excreted in the urine^{23,24}. The renal excretion of Mg is determined by the rate of filtration and its tubular reabsorption.

The filtered Mg is reabsorbed by the renal tubules: 10-25% in the proximal tubule, 60-70% in the thick ascending limb (TAL) and 5% in the distal tubule. Transport of Mg in TAL is primarily passive (moving from the lumen to the interstitium) via the paracellular channels, due to a transepithelial gradient generated by the apical NaK2Cl cotransporter (Figure 1).

Figure 1

Physiology of magnesium in the kidney.



Claudins are transmembrane proteins, an important component of the tight junctions, that regulate paracellular permeability to Mg and Ca. Claudins variants may change tight junctions' permeability and paracellular reabsorption of Mg and Ca driven by the electric gradient generated by potassium-channel (ROMK) and NaK2Cl co-transporter. The calcium-sensing receptor (CaSR) inhibits the expression of claudins in tight junctions and the activity of NaK2Cl, thus decreasing paracellular reabsorption of Ca and Mg.

The selectivity of the paracellular pathway is determined by claudins which form a cation-selective tight junction through which the Mg paracellular transport is done^{23,24}. Three claudin proteins, claudin-14, claudin-16 and claudin-19, make the cation-selective paracellular pathway for Ca and Mg. Claudin-16 and claudin-19 form the pores and claudin-14 inhibits the cation selectivity of that pore^{25,26}.

Tubular Mg transport is modulated by the extracellular fluid volume and by calcium receptors (CaSR) located in the basal pole of the tubular cells of the TAL, which are sensitive to serum Ca and Mg²⁷. The activation of these receptors inhibits the NaK2Cl cotransporter, dissipating the positive transepithelial gradient and decreasing the passive Ca and Mg reabsorption, which leads to the increase of Ca and Mg urinary losses^{23,24}. CaSR also inhibits the phosphorylation of claudins (and unphosphorylated claudins are not expressed in tight junctions) reducing the tight junction permeability to Ca and Mg²⁷.

■ GENETICS OF MAGNESIUM IN STONE FORMERS

Renal stones tend to cluster in families. Studies in twins and families have confirmed the heritability of the nephrolithiasis risk. Genome-wide association studies have identified several candidate genes implicated in renal tubular handling of lithogenic elements, such as calcium, phosphate and oxalate, and inhibitors of lithogenesis, such as Mg and citrate²⁸.

Growing evidence has shown the relevance of the tight junctions (made of claudin paracellular channels) in the regulation of Ca and Mg paracellular uptake. Lieske *et al.* found for the first time a heritable component of the Mg excretion²⁹.

A genome-wide association study suggested that variants in the claudin-14 gene are associated with the ratio of urinary excretion of Mg and to the urinary stone risk^{30,26}.

Claudin-14 knockout mice presented significantly higher serum Mg and lower fractional excretion rate for Mg and Ca³¹. These experimental findings corroborate the genetic data that points to claudin-14 as being a potential culprit of hypomagnesuria in stone formers.

These observations support the fact that genetic defects in claudins expressed in the TAL are part of a spectrum of tubular dysfunction found in stone formers.

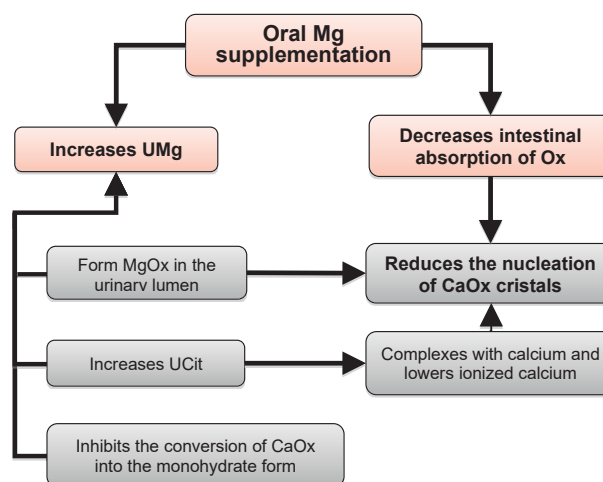
■ RATIONALE FOR MAGNESIUM SUPPLEMENTATION

Magnesium can have direct and indirect inhibitory effects on lithogenesis (Figure 2). The main mechanisms through which magnesium can inhibit the stone growth are:

- Oral Mg decreases intestinal absorption of oxalate and consequently urinary excretion of oxalate. Oral administration of Mg binds oxalate in the intestinal lumen, working as an effective oxalate-chelating agent³²⁻³⁴.
- Magnesium can also bind oxalate in the urinary lumen and form magnesium oxalate complexes, which are 100 times more soluble than calcium oxalate (0.07g/100ml versus 0.0007g/100 ml, respectively)³⁵, lowering the urinary saturation of calcium oxalate (CaOx)^{36,33,17,37,38}. It was demonstrated *in vitro* that Mg reduces the rate of growth and nucleation of CaOx crystals^{3,37}.
- Magnesium supplementation can increase urinary citrate excretion^{38,33,39,40,34}. It has been assumed that magnesium

Figure 2

The potential mechanisms through which oral magnesium supplementations inhibit calcium stones lithogenesis.



CaOx, calcium oxalate; Mg, magnesium; MgOx, magnesium oxalate; Ox, oxalate and UMg, urinary magnesium.

supplementation corrects the intracellular acidosis secondary to relative Mg deficiency leading to the reduction of the proximal tubular re-absorption of citrate^{41,34}. Citrate, a well-recognized lithogenic inhibitor, complexes with Ca and lowers ionized Ca, inhibiting Ca salts precipitation³⁷.

- Mg can directly inhibit crystal conversion of CaOx from the CaOx dehydrate form into the monohydrate form⁴²⁻⁴⁴. CaOx monohydrate has a lower solubility, is more stable and more resistant to fragmentation by lithotripsy⁴⁵⁻⁴⁸.

■ IMPACT OF MAGNESIUM SUPPLEMENTATION IN LITHOGENIC FACTOR AND STONE FORMATION

■ Urinary magnesium excretion

Oral supplementation with Mg salts is an effective way to increase UMg (Table 1) – confirmed in several studies performed in animal models, healthy humans and in stone formers¹⁷. All the studies in stone formers found a significant increase in the UMg excretion after each Mg supplementation (Table 1).

In animal models, changes in Mg amount in diet (with stable Ca content) significantly modified the Mg urinary excretion. Claudin-14 expression is regulated by Mg intake. *Corre et al.* found that a dietary Mg loading leads to an increase of claudin-14 expression, which blocks the paracellular cation channels made by claudin-16 and claudin-19²⁶ – the mechanism through which Mg supplementation increases the excretion of Mg in the urine.

The rise in UMg after supplementation can be amplified if the Mg salt is given with meals instead of taking it on an empty stomach³⁴. Also, the combination of Mg and citrate salts increases the urinary excretion of Mg more than either one supplement alone^{33,49}.

The water solubility of an Mg salt is important for its bioavailability. Organic Mg salts (for instance, citrate, gluconate, lactate and aspartate) have a higher water solubility than inorganic salts (for instance oxide and chloride), resulting in a greater intestinal absorption^{22,50,51}. In spite of the lack of studies in stone formers, studies in healthy volunteers found that the urinary excretion of Mg increases more significantly after administration of one of the organic Mg salts (citrate, lactate and aspartate, were tested) than with Mg oxide^{22,51}. However, even a relatively insoluble Mg salt can reduce oxalate absorption.

■ Urinary citrate and pH

Epidemiologic studies found that patients with lower UMg excretion had significantly lower urinary citrate excretion⁷.

Three of the four studies that evaluate the effect of Mg supplementation on urinary citrate found that apart from an increase in UMg, there was a concomitant increase in urinary citrate and in urinary pH (Table 1).

Unsurprisingly, Lindberg *et al.*³⁴ and Pak *et al.*³⁹ found that Mg citrate causes a higher increase in the urinary citrate than Mg oxide.

Table 1

Studies on magnesium supplementation in stone formers.

Reference	Number of patients	Drug and Dose	Effect on urinary parameters					Effect on stone formation	
			UpH	UMg	UOx	UCa	UCit	↑ ↓	Comment
Fetner (1977) ⁵⁶	4	MgO 1680 mg/day (1000mg of Mg)	↑	↑	=	↑ (in 2 patients)	NA	=	No change in the urinary growth or formation of CaOx or CaP crystals.
Vagelli (1998) ³²	9	Mg(OH) ₂ 500mg/day	NA	↑	↓	=	NA	↓	Decreases the stone recurrence rate from 0.75 to 0.11 stones/year/patient
Johanson (1980) ⁴⁰	90	Mg(OH) ₂ 400-500mg/day	NA	↑	NA	=	NA	↓	Decreases the stone recurrence rate from 0.8 to 0.03 stones/year/patient
Johanson (1982) ⁵³	98	Mg(OH) ₂ 500mg/day	NA	↑	NA	=	↑	↓	Decreases the stone recurrence rate from 0.8 to 0.08 stones/year/patient
Ettinger (1988) ⁵⁸	82	Mg(OH) ₂ 600 or 1300mg/day	NA	↑	=	=	=	↓	73.9 and 62.3% fewer calculi formation than predicted
Lindberg (1990) ³⁴	11	MgCit 40 mEq of K/day or MgO 40 mEq of K/day	=	↑	↓	↑	↑	=	No changes in the urinary saturation of CaOx or CaP.
Reungjui (2002) ⁴¹	64	0.1 mmol/L MgSO ₄ /kg body weight	↑	↑	NA	NA	↑	NA	–
Kato (2003) ³³	14	231 mg KNa-Cit + 250 mg MgO	↑	↑	↓	=	↑	↓	Decreases the CaOx saturation (but not the CaP saturation)
Pak (1992) ³⁹	10	24.5 mEq/day of K (KMgCit)	↑	↑	=	=	↑	↓	Inhibits the crystallization of UA and CaOx in urine
Ettinger (1997) ³⁶	64	42 mEq/day of K (KMgCit)	↑	↑	↑	=	↑	↓	Reduces the 3-year recurrence rate of CaOx stone formation in 85%

UpH, urinary pH; UMg, urinary magnesium; UCit, urinary citrate; UCa, urinary calcium; UOx, urinary oxalate; MgO, magnesium oxide; Mg(OH)₂, magnesium hydroxide; MgCit, magnesium citrate; KMgCit, potassium-magnesium citrate; MgSO₄, magnesium sulphate; CaOx, calcium oxalate; CaP, calcium phosphate; “=”, non-significant change; NA, not available; K, potassium and UA, uric acid.

■ Urinary oxalate

Observational studies have found an association between higher UMg levels and lower rates of hyperoxaluria, suggesting that high Mg intake can decrease urinary oxalate levels³.

Several studies in Ca stone formers³²⁻³⁴ found a significant reduction in urinary oxalate excretion with oral Mg supplementation. In order to maximize the effect of Mg as an intestinal oxalate-chelating agent, Mg salts should be given with meals³⁴. We noted that some of the authors who did not find a significant decrease in urinary oxalate levels during Mg salts supplementation did not specify if the Mg supplements were taken with meals, which could underestimate the chelating effect of these salts.

Liebman *et al.* found that Mg salts (Mg oxide) can be as effective as Ca salts (Ca carbonate) in reducing intestinal absorption of oxalates⁵².

■ Stone formation – crystallization and stone incidence

In vitro, Mg decreases the rate of nucleation and growth of CaOx crystals^{17,38}. Table 1 shows that the great majority of the studies that evaluate the effect of Mg supplementation in stone formers found a significant inhibitory effect in urine crystallization and/or a significant drop in the stone recurrence rate.

An important fall of the stone recurrence rate was found in the studies that evaluated the stone incidence rate during Mg supplementation. They found a significant drop in the incident rate from values higher than 0.7 stones/year/patient to an incident rate lower than 0.11 stones/year/patient^{32,40,53} (Table 1).

Allie *et al.* and Rodgers *et al.* found a positive synergic effect of the combination of Mg with citrate on supersaturation of Ca phosphate (lower supersaturation in the combination vs. citrate or Mg alone)^{49,50}. The reduction of Ca phosphate saturation is favorable since it has been reported that these crystals might provide the nucleus for CaOx stone formation.

Not all intervention studies with Mg demonstrate a beneficial effect. We could find several reasons for this: small studies (all studies with less than one hundred patients); different Mg salts (with different bioavailabilities) were tested; several doses of each salt were used; studies were done on patients with different stone recurrence risk and different levels of Mg deficiency; several studies ignore the indication to take the Mg supplements with meals and others did not mention the concomitant optimization of the standard stone prophylaxis.

Regarding the effect of magnesium supplementation on calcium phosphate stones prevention, the evidence is even more scarce (or absent). The theoretical preventive effect of magnesium in CaP prophylaxis is based on *in vitro* experiments that showed a marked inhibitory effect on crystal nucleation, growth and aggregation^{54,55,53}.

The few studies that evaluated the effect on CaP formation *in vivo* (all with less than 10 patients) did not find a significant change in the

urinary saturation ratio, formation or rate of CaP crystal growth with magnesium supplementation (Table 1)^{56,34,33,39}.

■ HOW AND WHO SHOULD WE SUPPLEMENT WITH MAGNESIUM?

It is important to note that Mg supplementation cannot be recommended for all the stone formers. Most of the stone formers present UMg in the normal range, and an increase in the UMg level will present only a modest inhibitory effect⁶. As Massey⁶ explained, Mg-deficient stone formers who excrete less than 50 mg/day are more likely to benefit from Mg supplementation.

No studies have compared the different inhibitory lithogenic effect of the several oral Mg supplements available on the market. The most appropriate salts for stone formers supplementation are those that at the same time binds oxalate in the gut and increase UMg and citrate, since they are more likely to reduce the risk of stone recurrence.

Considering the higher bioavailability of organic Mg salts and the potential synergic effect of Mg and citrate, Mg citrate seems to be the most advantageous to inhibit lithogenesis³⁵.

Azarfar *et al.* found that adding Mg salts to the standard nephrolithiasis treatment (with potassium citrate) could add benefit in lowering urinary risk factors^{35,57}.

Lindberg *et al.* found that when Mg supplements were given with meals, apart from a more pronounced reduction in urinary oxalate, there was a more significant increase in UMg and citrate. This combination was found to be more effective than Mg salt alone as inhibitor of CaOx crystallization^{33,34}.

Thus, Mg supplementation should be recommended as an adjuvant therapy to the standard prophylactic therapy for those patients with high risk of stone recurrence and severe to moderate hypomagnesuria, given as an organic salt during the meal.

■ ADVERSE EFFECTS OF MAGNESIUM SUPPLEMENTATION IN STONE FORMERS

The kidney has a good capacity of Mg excretion. So hypermagnesemia generally could occur only when there is an excessive Mg intake/supplementation in combination with a moderate to severe renal insufficiency. Serum Mg is maintained within the normal range until the glomerular filtration rate falls below 20 ml/min – the remaining nephrons compensate the decline in the Mg filtered by increasing significantly its fractional excretion (reducing its tubular reabsorption)²³.

The majority of the studies in stone formers found no significant increase in serum Mg levels during long-term Mg supplementation⁴¹. However, Johansson and colleagues found in two different studies a transitory rise in serum Mg levels during the first 6-12 months of supplementation, but thereafter the mean value returned to the pre-treatment level^{40,53}. Nevertheless, no study reported any case of hypermagnesemia or any symptom that could be attributed to it^{40,53}.

The most frequently reported adverse effect of taking oral Mg supplements was minor gastrointestinal discomfort, namely diarrhea, with a reported frequency from 9 to 17%^{33,36,40}.

Apart from these common adverse effects of the Mg supplementation common to everyone, one of the main concerns regarding the use of these supplements in stone formers is the risk of hypercalciuria.

Calciuria is modulated by urinary pH and Mg. Experimental studies suggest that low urinary pH and high Mg levels increase calciuria by inhibiting TRPV5/6-mediated Ca reabsorption in the distal nephron¹⁷.

Although some human studies have found a statistically significant increase of urinary Ca excretion with Mg supplementation, this rise in the urinary Ca did not cause a significant increase in the urinary saturation or stone formation rate (Table 1)¹⁷.

Kato *et al.*³³ found, in healthy volunteers and also in stone formers, that the increase in Ca excretion did not happen if Mg supplementation was taken concomitantly with citrate. In the same direction, Bonny and colleagues¹⁷ found that although urinary Ca significantly increases after Mg salts supplementation alone, its combination with potassium-magnesium citrate did not change significantly the urinary Ca excretion.

These results suggest that Mg supplements should be provided preferentially in the form of Mg citrate or taken with concomitant urinary alkalinizing agents^{33,44}.

CONCLUSIONS

Hypomagnesuria is frequent in recurrent stone formers and higher UMG levels were associated with lower incidence of kidney stones. Oral Mg supplementation is an effective way to increase magnesuria in stone formers. Oral Mg supplements have direct and indirect lithogenic inhibitory effects in Ca stone formers, with minor adverse effects. The authors suggest that Mg supplementation, mainly in association with an alkali salt, is an underestimated weapon in the prevention of Ca stone recurrence. This combination avoids the rise in the urinary Ca and maximizes the inhibitory lithogenic effect, by a greater increase in the urinary citrate and Mg.

Larger, longer and better-designed clinical trials to evaluate the effectiveness of Mg supplementation in the prevention of stone recurrence are needed. The trial design should include patients with different UMG deficits and should require that the Mg supplement be taken with meals to maximize its benefits.

Disclosure of potential conflicts of interest: none declared

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Correspondence to:

Ivo Laranjinha, MD

Nephrology Department, Hospital de Santa Cruz/Centro Hospitalar de Lisboa Ocidental

Av. Prof. Dr. Reinaldo dos Santos 2790-134, Carnaxide, Portugal

Mobile phone: 00351 935013798;

E-mail: ivolaranjinha@gmail.com

ORCID: 0000-0001-5895-0680