

Short-Term Magnesium Oxide Monohydrate Treatment Improves Lipids and Inflammation in Healthy Subjects

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Background:

Magnesium content in food consumed in the Western world is steadily decreasing. Hypomagnesemia is associated with increased incidence of diabetes mellitus, metabolic syndrome, all-cause and coronary artery disease mortality. We investigated the impact of supplemental oral magnesium citrate versus magnesium oxide monohydrate on lipoproteins and high-sensitivity C-reactive protein (hs-CRP) in healthy subjects with no apparent heart disease.

Methods and Results:

In a randomized, prospective, double-blind, crossover study, 41 (20 women) healthy volunteers [mean age 53 ± 8 (range 31-75) years] received either magnesium oxide monohydrate tablets (520 mg/day of elemental magnesium) or magnesium citrate tablets (295.8 mg/day of elemental magnesium) for 1 month (phase 1), followed by a 4-week wash-out period, and then crossover treatment for 1 month (phase 2). Intracellular magnesium levels ($[Mg^{2+}]$), assessed from sublingual cells through x-ray dispersion (normal values 37.9 ± 4.0 mEq/L), serum lipoproteins and hs-CRP levels were assessed before and after each phase. Oral magnesium oxide monohydrate, rather than magnesium citrate, significantly increased $[Mg^{2+}]$ (34.4 ± 3 vs 36.3 ± 2 mEq/L, $p=0.001$ and 34.7 ± 2 vs 35.4 ± 2 mEq/L, $p=0.097$; respectively), reduced total cholesterol (201 ± 37 vs 186 ± 27 mg/dL, $p=0.016$ and 187 ± 28 vs 187 ± 25 mg/dL, $p=0.978$; respectively) low-density lipoprotein (LDL) cholesterol (128 ± 22 vs 120 ± 25 mg/dL, $p=0.042$ and 120 ± 23 vs 121 ± 22 mg/dL, $p=0.622$; respectively) (Figure) and hs-CRP (3.1 ± 3.0 vs 2.6 ± 2.0 mg/dL, $p=0.030$ and 3.0 ± 4.0 vs 3.5 ± 5.0 mg/dL, $p=0.438$; respectively).

Conclusions:

Short-term oral magnesium oxide monohydrate treatment significantly improved total and LDL cholesterol and hs-CRP compared with magnesium citrate in healthy subjects with no apparent heart disease.

Figure:

