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Evidence Table 19. Managing Exacerbations: Magnesium Sulfate

Abbreviations used in table:

AE	adverse event
ARR	absolute risk reduction
ED	emergency department
FEV ₁	forced expiratory volume in 1 second
FVC	forced vital capacity
ICS	inhaled corticosteroid
ICU	intensive care unit
IVMg	intravenous magnesium sulfate
MgSO ₄	magnesium sulfate
Р	placebo
PEFR	peak expiratory flow rate
RR	relative risk
SAE	serious adverse event
SBP	systolic blood pressure
SMD	standardized mean difference
WMD	weighted mean difference

* indicates primary outcome

Evidence Table 19. Managing Exacerbations: Magnesium Sulfate

Magnesium by Nebulizer

Citation (Sponsor) Study I	Design Purpose/Objective	Study N (Number Evaluable)	Asthma Severity at Baseline (if reported)	Treatment	Lung Function	Severity/Admissions	Safety
Blitz et al. Inhaled magnesium sulfate in the treatment of acute asthma. Cochrane Database Syst Rev 2005;(2):CD003898. (Alberta Cancer Board, Canada; Canadian Institutes of Health Research, Ottawa, Canada; Department of Emergency Medicine, University of Alberta, Edmonton, Alberta, Canada)	nalysis of iized ed trials ed n 1966 03 administered in acute asthma on pulmonar functions and admissions	Six trials conducted in the United States, India, New Zealand, Turkey, and Argentina with 296 patients; three included adults, one included adults and pediatric patients, two enrolled pediatric patients. Trials were published between 1995 and 2003. Methodological quality was high: five trials scored 3 on the Jadad scale. All rated a B in concealment of allocation.	Three studies enrolled severe asthmatics (FEV ₁ or PEF <50% predicted). Five studies enrolled patients presenting to the emergency department (ED). Two studies excluded patients who had taken asthma medication within 12 hours. One excluded patients who had received corticosteroids in previous 7 days; one excluded patients who had received steroids, theophylline, or ipratropium bromide within 3 days of presenting to ED.	Four studies compared MgSO ₄ with beta ₂ -agonist to beta ₂ - agonist with placebo; two studies compared MgSO ₄ to beta ₂ -agonist.	MgSO ₄ with beta ₂ -agonist vs.beta ₂ - agonist alone Pulmonary functions were improve for MgSO ₄ with beta ₂ -agonist vs. beta ₂ - agonist alone (SMD 0.23, 95% Cl –0.03 to 0.50) with no difference between results from adults and those with children. There was a significant difference in results from severe asthma trials (SMD 0.55, 95% Cl 0.12 to 0.98). MgSO ₄ vs. beta ₂ -agonist alone There was no advantage for MgSO ₄ alone (SMD 0.17, 95% Cl –0.51 to 0.86).	MgSO ₄ with beta ₂ -agonist vs. beta ₂ -agonist alone There was no clear reduction in probability of admission for MgSO ₄ with beta ₂ -agonist (relative risk (RR) 0.69, 95%Cl 0.42 to 1.12). There was no difference between adults and children or between severe and less severe asthma. MgSO ₄ vs. beta ₂ -agonist alone There was no difference between MgSO ₄ and beta ₂ -agonist alone based on a single trial (RR 0.50, 95% Cl 0.04 to 6.12).	All studies reported no SAE in either arm.

Magnesium by IV

Citation (Sponsor)	Study Design	Purpose/Objective	Study N (Number Evaluable)	Population Characteristics	Asthma Severity at Baseline (if reported)	Treatment	Dose	Duration of Active Treatment; Duration of Postintervention/Off-Treatment Followup	Lung Function	Vital Signs/Cardiovascular/Clinical Laboratory Values	Severity/Admissions	Safety
Boonyavorakul et al. Intravenous magnesium sulfate in acute severe asthma. Respirology 2000:5(3):221– 225.	Prospective, randomized, double- blind, placebo-controlled trial (ED in Thailand)	To determine whether intravenous MgSO₄ (IVMg) as an adjunct to a standard therapy can reduce admission rate and severity scores in patients with acute severe asthma compared with those treated by standard therapy	33 (33)	Age: ≥15 yr, mean = 39 yr Gender: 12% male, 88% female	Acute severe asthma History of intubation, 12.1% Oral steroid use, 12.1% Inhaled steroid use, 33.3% Duration of asthma attack, median 4.5 hrs Pulse, mean = 125.5 beats/min Respiration, mean = 33.1/min Fischl index, mean = 6.02	Arm 1 MgSO ₄ (n=17; 17 completers) Arm 2 Placebo (P) (n=16; 16 completers)	2 g MgSO₄ in 50 mL of 0.9% normal saline 2 mL of sterile water in 50 mL of 0.9% normal saline	All patients received 5 mg intravenous dexamethasone, 2.5 mg nebulized salbutamol at 0, 20, 40, and 60 minutes. Measurements were taken at 60, 120, 180, and 240 min.			Admission rates were 17.65% for MgSO ₄ and 25% for P (RR 0.71, 95% CI –0.19 to 2.67). Necessary to treat 14 patients with MgSO ₄ to prevent one admission. Mean severity scores of two groups were same at all time points (p=0.37).	
Rowe et al. Magnesium sulfate for treating exacerbations of acute asthma in the emergency department. Cochrane Database Syst Rev 000;(2): CD001490. (Canadian Association of Emergency Physicians: National Institute of Health; University of Alberta, Edmonton, Canada; Acute Care Research Group, Sudbury Regional Hospital, Sudbury, Ontario, Canada; and NHS Research and Development UK) NOTE: Review includes Silverman et al. study.	Meta-analysis of studies published between 1989 and 1997	To examine the effect of additional IVMg in patients with acute asthma managed in the ED	Seven randomized controlled trials with 655 subjects; five adult and two pediatric studies		Presenting to ED for treatment of acute asthma	Treatment Groups: MgSO4 Control groups: Placebo (P)			No difference in improvement in PEFR (WMD 294, 95% CI -3.4 to 62) and % predicted FEV1 (WMD 4.3, 95% CI -2.3 to 10.9).	Heart rate and respiratory rates did not change with IV MgSO ₄ . Slight but not clinically important change in systolic blood pressure (SBP).	*There was no significant difference in MgSO ₄ vs. P in hospital admission (OR 0.31, 95% Cl 0.09 to 1.02). In patients with severe asthma, admissions were lower with MgSO ₄ vs. P (OR 0.10, 95% Cl 0.04 to 0.27); no difference for those with mild to moderate asthma (OR 1.35, 95% Cl 0.72 to 2.55).	Few adverse events (AE) were reported.

Citation (Sponsor)	Study Design	Purpose/Objective	Study N (Number Evaluable)	Population Characteristics	Asthma Severity at Baseline (if reported)	Treatment	Dose	Duration of Active Treatment; Duration of Postintervention/Off-Treatment Followup	Lung Function	Severity/ Admissions
Porter et al. Intravenous magnesium is ineffective in adult asthma, a randomized trial. Eur J Emerg Med 2001;8(1):9–15.	Prospective, randomized, double-blind, placebo- controlled trial (urban ED in United States)	To test hypothesis that intravenous MgSO ₄ would improve the outcome of severe asthmatics concurrently receiving maximal conventional therapy	42 (42)	Age: ≥18 yr, mean = 35 yr Gender: 36% male, 64% female	Acute asthma exacerbation PEFR ≤100 L/min or <25% of predicted, mean = 89 L/min Heart rate, mean = 108 Respirations, mean = 31/min Mean arterial pressure = 100 mmHg Oxygen saturation, mean = 94% Borg dyspnoea scale, mean = 6.1	Arm 1: MgSO ₄ (n=18) Arm 2: Placebo (P) (n=24)	2 g in 50 mL normal saline 50 mL normal saline	Patients received 2.5 mg albuterol sulfate, 125 mg methylprednisolone, and 50 ml study solution over 20 minutes at 0, 20, 40, and 60 minutes.	*PEFR was 174 L/min in MgSO ₄ vs. 212 L/min in P (p=0.038). Controlling for age and baseline PEFR, pulse oximetry, and heart rate, PEFR at 60 minutes in MgSO ₄ averaged 75% that of P (95% CI 52% to 109%, p=0.132).	Groups did not differ in Borg dyspnoea scale score at 60 minutes in either univariate or multivariate analysis. There was no difference in hospital admission rate (28% of MgSO ₄ vs. 21% of P, p=0.72).
Cheuk et al. A meta-analysis on intravenous magnesium sulphate for treating acute asthma. Arch Dis Child 2005;90(1):74–77.	Meta-analysis of controlled clinical trials involving children below 18 years of age	To evaluate the effectiveness of IVMg in preventing hospitalization or intensive care unit (ICU) admission in children with acute asthmatic attacks, either used alone on in addition to standard therapies	Five trials involving 182 patients conducted in EDs and published between 1996 and 2000. Methodological quality was 4 or 5 on Jadad scale for all studies.	Children under 18 years of age	Inadequate response to first line treatment with three doses of beta ₂ -agonists	MgSO₄ of any therapeutic dose either alone or in additional to standard therapies.			OR of persistent PEFR <60% predicted is 0.155 (95% CI 0.057 to 0.422, p=0.00033) Difference in % improvement of PEFR at study end, 8.58 (95% CI 0.94 to 16.22, p=0.028)	IVMg was effective in avoiding hospitalization (absolute risk reduction (ARR) 0.257, 95% CI 0.124 to 0.389, p=0.0001). Number needed to treat in avoiding hospitalization is 4 (95% CI 3 to 8). Difference in clinical symptom score at study end, 1.33 (95% CI 0.31 to 2.36, p=0.011)