



ORIGINAL ARTICLE

Nebulized salbutamol for asthma: Effects on serum potassium and phosphate levels at the 60 min

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KEYWORDS

Salbutamol;
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Abstract

Objective: We conducted this prospective study to expand available information in relation to serum phosphate levels in treatment of acute asthma. A β -adrenergic agonist, salbutamol, was used for this purpose.

Material and methods: Twenty-six patients who met the inclusion criteria as; age over 16 years, asthma history, and an acute exacerbation were included. Serum blood urea nitrogen, creatinine, glucose were within normal limits in all the patients. None of the patients were on chronic theophylline therapy. Baseline serum phosphate and potassium levels were measured. Nebulized salbutamol (2.5 mg) was used three times at every hour. After 60 min, serum phosphate and potassium levels were measured.

Results: Serum phosphate levels decreased from 3.7 ± 0.9 mg/dL (baseline) to 3.6 ± 0.9 mg/dL at 60 min. This decrease was not statistically significant ($p = 0.373$). Serum potassium levels decreased significantly ($p < 0.001$) from 4.6 ± 0.7 mmol/L (baseline) to 4.3 ± 0.7 mmol/L (60 min).

Conclusion: Administration of nebulized salbutamol during the emergency treatment of acute exacerbation of asthma is not associated with a statistical decrease in serum phosphate. There was significant hypokalemia. This study indicates that a further study is needed to elucidate the clinical significance of nebulized salbutamol on serum phosphate.

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PALAVRAS-CHAVE

Salbutamol;
Fosfato;
Asma

Salbutamol Nebulizado na Asma: Efeitos nos níveis de Potássio e Fosfato Sérico aos 60 minutos**Resumo**

Objetivo: Levámos a cabo este estudo prospetivo para ampliar a informação disponível relativamente aos níveis de fosfato sérico no tratamento de asma aguda. Foi utilizado um agonista β -adrenérgico, salbutamol, para este efeito.

Materiais e métodos: Foram incluídos 26 doentes que cumpriam os critérios de inclusão de: idade superior a 16 anos, história de asma, incluindo uma exacerbação aguda. A ureia do soro sanguíneo, creatinina e glucose estavam nos limites normais em todos os doentes. Nenhum dos doentes era submetido a terapia crónica de teofilina. Os níveis da base de referência de fosfato e potássio sérico foram medidos. O nebulizador de salbutamol (2,5 mg) era usado 3 vezes por hora. Após 60 min, os níveis da base de referência de fosfato e potássio sérico eram medidos.

Resultados: Os níveis de fosfato sérico diminuíram de $3,7 \pm 0,9$ mg/dL (base de referência) para $3,6 \pm 0,9$ mg/dL aos 60 min. Esta redução não era estatisticamente significativa ($p = 0,373$). Os níveis de potássio sérico diminuíram significativamente ($p < 0,001$) de $4,6 \pm 0,7$ mmol/L (base de referência) para $4,3 \pm 0,7$ mmol/L aos 60 min.

Conclusão: A administração de salbutamol nebulizado durante o tratamento de urgência de uma exacerbação aguda de asma não está associada a uma diminuição estatística do fosfato sérico. Ocorreu uma hipocalcemia significativa. Este estudo indica que é necessário um estudo adicional para esclarecer o significado clínico do salbutamol nebulizado no fosfato sérico.

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Introduction

Asphyxia is responsible for the majority of deaths in patients with acute asthmatic crisis, and the adverse effects of medical treatments are also important in terms of the deaths.¹ Nebulized β -adrenergic agonists are the mainstay of treatment for acute exacerbation of asthma in the emergency setting.² β -Adrenergic agonists (BAA) have side effects such as tremor, palpitations, tachycardia, and anxiety.³ There are several studies that demonstrate a statistically significant decrease in serum potassium and magnesium after BAA administration.⁴⁻⁷ In addition, there are also some studies regarding the BAA induced hypophosphatemia.⁸⁻¹¹ In the current research, our aim is to investigate Serum Potassium and Phosphate Levels at the 60 min after administration of Nebulized Salbutamol for Asthma.

Materials and methods

This prospective study is conducted in patients with asthmatic attack in the emergency department. Patients were treated with a BAA-salbutamol – according to a study protocol that was approved by the local ethical committee. Informed consent was obtained from all the patients. Baseline data including age, sex, clinical history, and the medications used were recorded. All patients included in the study met the criteria as: age >16 years, a history of acute exacerbation of asthma (dyspnea, cough and, on examination, wheezing). Patients with respiratory failure, altered mental status, renal failure, diabetes mellitus, and cirrhosis were excluded. Patients using alcohol, diuretics, theophylline, aminophylline, antidiabetic, and antihypertensive drugs were also excluded. The samples had BAA at

least four hours before the admission. The severity of the exacerbation determines the treatment modality. Indices of severity including particularly PEF, pulse rate, respiratory rate, and pulse oximetry, had to be monitored during treatment. The exacerbation severity was defined according to the Global Strategy for Asthma Management and Prevention 2011.¹²

All the patients were monitored by continuous electrocardiography, and arterial oxygen saturation by pulse oximetry with a finger oximeter. The baseline peak expiratory flow rate was measured. An intravenous (i.v.) line for repeated blood sampling was established and baseline laboratory assay which included serum phosphate, creatinine, blood urea nitrogen, glucose, potassium levels was performed. Patients were given initial nebulized treatment of salbutamol (2.5 mg).

Nebulized salbutamol (2.5 mg) was used three times during the first hour. After the third administration of salbutamol, serum phosphate and potassium levels were assessed.

None of the patient received magnesium, potassium, phosphate, corticosteroids, theophylline, and i.v. dextrose or saline during the study period.

The mean and standard deviation were calculated for baseline and subsequent measure of phosphate and potassium. Paired-*t* test was used for analyzing the changes in case of phosphate and potassium levels. *p*-Value <0.05 was considered statistically significant.

Results

Twenty-six patients were enrolled during the study period. The mean patient age was 58.8 ± 10.3 years (range 35–74 years), with 11 female and 15 male asthmatic patients.

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