

Supraventricular tachycardia after nebulized salbutamol therapy in a neonate: Case report

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ABSTRACT

Supraventricular tachycardia (SVT) is the most common sustained arrhythmia in neonates and infants. Presentation of SVT in the neonate is usually subtle and frequently complicated by congestive heart failure. Despite the widespread use of β_2 -agonists, their safety has been questioned. Several studies have reported an increased incidence of cardiac arrhythmias in patients treated with these agents, and other studies have found increased rates of cardiovascular death associated with the use of oral and nebulized β_2 -agonists such as salbutamol, which is used to treat bronchospasm in newborns with several diseases. Herein, we report a case of SVT following administration of nebulized salbutamol in a neonate.

Keywords: tachycardia supraventricular, salbutamol, newborn.

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INTRODUCTION

Supraventricular tachycardia (SVT) includes all forms of tachycardia originated above the bifurcation of the bundle of His or have mechanisms dependent on the bundle of His. SVT is the most common sustained arrhythmia in neonates and infants. Predisposing factors such as congenital heart disease, drug administration, illness, and fever occur only in 15% of cases.¹ The presentation of SVT in the neonate is frequently subtle and it may include pallor, cyanosis, restlessness, irritability, feeding difficulty, tachypnea, diaphoresis and grunting. SVT in newborns is frequently complicated by congestive heart failure, characterized by a fixed heart rate usually greater than 230 beats per minute. Typical electrocardiogram (ECG) findings of SVT include narrow QRS complexes, regular R-R

intervals, and absent P waves or an abnormal P wave axis. Salbutamol, which has been used as bronchodilators for the prevention and treatment of chronic lung disease in preterm infants, for emergency treatment of hyperkalemia, and transient tachypnea of the newborn (TTN),²⁻⁴ is a direct-acting sympathomimetic agent that has a selective β -adrenergic effect. Because β -receptors in the heart are mainly the β -1 type, it is believed that salbutamol has minimal cardiovascular effects; however, tachycardia and cardiac arrhythmias have been reported as complications of salbutamol administration.⁵ Although well documented in adults, cardiac arrhythmias have not been widely reported in pediatric patients as a complication of salbutamol therapy.

This paper presents the case of an infant who was admitted to the neonatal intensive care unit (NICU) with a diagnosis of meconium aspiration syndrome and who developed multiple SVTs following administration of salbutamol during the recovery phase of her underlying illness.

CASE REPORT

The patient was a girl born to a 28-year-old mother via a spontaneous vaginal delivery after a term gestation; the birth weight was 3600 g. The mother was healthy and had an uneventful pregnancy. The baby was delivered through thick meconium and she was cyanotic. Her mouth was suctioned and she was intubated and suctioned for thick, particulate meconium. Her pulse fell below 100 beats per minute (bpm) and positive-pressure ventilation (PPV) was initiated. The patient's first and fifth Apgar scores were 5 and 6, respectively and she was transferred to the NICU. On admission to the NICU, the infant had a pulse of 157 (bpm), a respiration rate of 72 per minute, and a mean arterial blood pressure of 34 mmHg. She remained with 100% oxygen. A capillary blood gas test showed a pH of 7.31, a PCO_2 of 43 mmHg, and a SpO_2 of 94% at 20 minutes of life, and she was severely tachypneic. After intubation, a blood culture was taken, a sepsis evaluation was performed, and intravenous (IV) antibiotics (crystalline penicilline; 50,000

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units/kg/day in divided doses every 12 hours, and gentamicine 4 mg/kg/day every 24 hours) were begun. Chest radiography revealed dense bilateral infiltrates with prominent air bronchograms. An echocardiogram at 10 hours of life revealed septal hypertrophy with flattening of the ventricular septum, consistent with persistent pulmonary hypertension of the neonate (PPHN). Treatment was aimed at interrupting the cycle of pulmonary vasoconstriction and hypoxia. Her chest radiographs started to demonstrate a reticulogranular pattern, and there was concern that she might have an element of surfactant deficiency.

Over the next several days, the need to ventilate and oxygenate the patient gradually decreased, and on day 7 of life, she was extubated to a nasal cannula. She had coarse bilateral breath sounds with marked expiratory wheezing. The cardiovascular examination was unremarkable, the echocardiographic examination was normal, and serum potassium and the other electrolytes were in the normal ranges at this time. Inhaled salbutamol therapy was initiated at a dose of 0.15 mg/kg (0.45mg) every 4 hours. Following the inhalation of salbutamol solution (Ventolin nebuler[®] 50, Glaxo Wellcome; Greenford, UK), the infant's heart rate increased suddenly to 314 bpm. The ECG showed a narrow complex tachycardia at a heart rate of 310 bpm and retrograde conducted P waves buried in the ST segment at an RP interval (*Figure 1*). Adenosine was administered via IV rapid bolus at a dose of 100 mcg/kg which was repeated twice. However, we were not able to obtain sinus rhythm despite the raised dose for the two subsequent

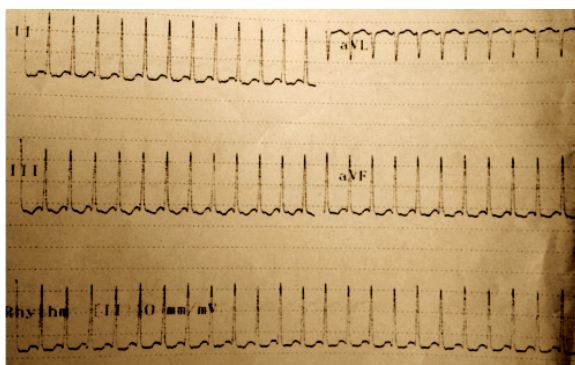
administrations of adenosine treatment and electrical cardioversion was performed, which terminated the SVT. Salbutamol treatment was ceased and SVT was never observed again. The patient was doing well and was discharged from the hospital on day 15 of life.

DISCUSSION

Despite the widespread use of β_2 -agonists, their safety has been questioned. Existing data on the effects of these agents, especially those administered in an inhaled form, on myocardial electrophysiological properties are rare.⁶ Pharmacologically, however, inhaled salbutamol can result in significant changes in cardiac electrophysiological properties.⁶ Salbutamol has been found to enhance atrioventricular nodal conduction and to decrease atrioventricular nodal, atrial, and ventricular refractoriness in addition to its positive chronotropic effects. β_2 -agonists also increase QT dispersion. All of these alterations theoretically could contribute to the generation of tachycardia and tachyarrhythmias. Several studies have reported an increased incidence of cardiac arrhythmias in patients treated with the use of oral and nebulized β_2 -agonists.⁵⁻⁷ A recent human study demonstrated that salbutamol, a selective β_2 -agonist, administered by nebulizer has significant electrophysiological effects on the atrium, nodes, and ventricle. Some studies found that salbutamol produced more evident changes on the electrophysiologic properties of the sinus node compared with the atrioventricular node.^{6,8} The cardiac effects of β_2 -agonists include tachycardia, atrial and ventricular ectopic complexes, and atrial and ventricular arrhythmias.^{6,9}

Although adverse drug reactions including tremor (90%), hypokalaemia (45.5%), and supraventricular tachycardia (21%) were reported to be common, especially after administration of continuous intravenous infusion of salbutamol,¹⁰ the estimated SVT rate in children treated with inhaled β_2 -agonists remains unknown. Tachycardia and tachyarrhythmias are very common in critically ill adult patients with various conditions.¹¹ Predisposing factors for SVT include congenital heart disease, fever, and sympathomimetics.^{8,12} Our patient had no history of congenital heart disease or fever. Although she required brief PPV in the delivery room and respiratory support and she suffered from mild PPHN, her myocardial functions and cardiac output were always within normal limits, and

FIGURE 1. Electrocardiogram indicative of supraventricular tachycardia, with no visible P wave and normal QRS morphology.



no signs were observed of myocardial ischemia, which could cause a tendency toward arrhythmia. Acute management of SVT in children involves the use of vagal maneuvers or IV adenosine. IV adenosine has been found to be safe and highly effective in the management of SVT in infants and children.⁵

Two previous patient reports have described salbutamol-induced SVT in children. In the first report, SVT in a 19-month-old baby was converted by applying facial ice, and in the second case, a 4-year-old patient recovered with IV adenosine.^{5,6} We initially administered a 100 mcg/kg dose of IV adenosine, and despite two repeated doses of 200 mcg/kg, we could not obtain sinus rhythm. Cardioversion was performed with 2j/kg followed by 4j/kg, and this treatment was successful.

Although nebulized, short-acting β_2 -agonists are effective bronchodilator drugs that are widely prescribed for the treatment of airflow obstruction and increasingly for TTN and hyperkalemia in NICUs, there is considerable concern about their potential side effects, particularly those pertaining to heart rate and arrhythmias. In addition, β_2 agonists are widely used to treat neonates with bronchopulmonary dysplasia, but there is no evidence of their efficacy.

Therefore, we suggest that all infants should be carefully monitored for tachycardia which requires urgent treatment after nebulized salbutamol in both ambulatory and inpatient care, even if there are no underlying causes. Underlying causes that might facilitate arrhythmia should be evaluated. ■

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