Acquired Bartter-like Syndrome Presenting with Polyuria and Reversible Hypokalemia Associated with Colistin Use in a Critically Ill Pediatric Patient

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ABSTRACT

We report a case of an acquired Bartter-like syndrome (BLS) after 3 days of treatment initiation and improved after discontinuation of colistin therapy in pediatric intensive care unit. A 2-month-old girl with spinal muscular atrophy type 1 who had respiratory distress received colistin therapy with a dose of 5 mg/kg/day for Acinetobacter baumannii complex isolation from endotracheal aspirate on the 12th day follow-up. Polyuria (6 mL/kg/hour) in the presence of normal blood pressure and hypokalemic metabolic alkalosis were developed on the 3rd day of colistin treatment. Colistin was stopped on the 4th day, and 2 days after discontinuation of colistin, polyuria improved dramatically. Her metabolic alkalosis and hypokalemia discontinued after 2 and 4 days, respectively.

There are very few reports about colistin-induced BLS. The onset of polyuria, hypokalemia, and metabolic alkalosis during treatment with colistin and resolution after interruption suggest a causative relationship.

Keywords: Bartter-like syndrome, Colistin, Hypokalemia, Pediatric, Polyuria.

Indian Journal of Critical Care Medicine (2021): 10.5005/jp-journals-10071-23898

INTRODUCTION

Colistin is a polypeptide antibiotic that belongs to the polymyxin class of cationic polypeptide antibiotics.1 The emergence of multidrug-resistant, gram-negative bacterial infections has resulted in a significant increase in the use of intravenous colistimethate sodium.2 Colistin has dose-dependent bactericidal effect and also postantibiotic effect against Acinetobacter baumannii, Pseudomonas aeruginosa, and Klebsiella pneumonia.3,4 Although colistin-associated nephrotoxicity has been previously reported on a wide range in the literature, there are very few reports about colistin-induced Bartter-like syndrome (BLS). BLS develops during treatment with different classes of drugs and is presented with hypokalemic metabolic alkalosis, hypomagnesemia, hypocalcemia, and normal serum creatinine levels.5 Herein we report a 2-month-old infant of an acquired BLS after 3 days of treatment initiation and improved after discontinuation of colistin therapy in pediatric intensive care unit (ICU).

CASE DESCRIPTION

A 2-month-old girl with spinal muscular atrophy type 1, receiving high-flow nasal cannula (HFNC) due to respiratory distress, was transferred from another hospital to our pediatric ICU. On admission, she was slightly tachycardic with a heart rate of 136/minute, blood pressure was 111–63 mm Hg, SpO2 was 100%, and capillary refilling was 2 seconds. On physical examination, she had tachypnea and her intercostal retractions and auscultation of the lungs revealed bilateral secretory rales. Preliminary blood gas and complete blood count were as follows: pH: 7.38, pCO2: 55 mm Hg, pO2: 83 mm Hg, HCO3: 26, lactate: 1.3 mmol/L, Hb: 9.4 g/dL, WBC: 8.9 × 103 μL, platelets: 403 × 103 μL, BUN: 1.2 mg/dL, creatinine: 0.06 mg/dL, albumin: 34 g/L, sodium: 136 mEq/L, potassium: 4 mg/dL, phosphate: 4.2 mg/dL, calcium: 9.8 mg/dL, magnesium: 2 mg/dL, and procalcitonin: 0.2 ng/mL (0–0.5 normal range).

On the 4th day, we noted decreased respiratory distress and a complete resolution of secretory rales. Chest X-ray showed new-onset infiltrates in the lower lobe of the left lung. Lung ultrasound showed increased localized B-lines in the zone 1 of the right lung. Meropenem and amikacin were initiated to extend the antimicrobial spectrum, and cefotaxime was discontinued.

On the 8th day, polyuria (6 mL/kg/hour) in the presence of normal blood pressure and hypokalemia was observed. Colistin therapy was stopped on the 10th day follow-up, due to the persistence of recurrent apnea and worsening of respiratory failure while in noninvasive ventilation, patient underwent tracheostomy. Two days after tracheostomy procedure, patient’s ventilatory pressures needed to be increased and she developed worsening respiratory distress.

We continued HFNC oxygen therapy and cefotaxime treatment for her pneumonia that she was already receiving. On the 2nd day of intensive care hospitalization, she required noninvasive ventilation through bilevel positive airway pressure due to the recurrent apneic episodes. The patient developed progressively increasing respiratory distress, and her chest X-ray showed new-onset infiltrates in the lower lobe of the left lung. Lung ultrasound showed increased localized B-lines in the zone 1 of the right lung. Meropenem and amikacin were initiated to extend the antimicrobial spectrum, and cefotaxime was discontinued.

On the 10th day follow-up, due to the persistence of recurrent apnea and worsening of respiratory failure while in noninvasive ventilation, patient underwent tracheostomy. Two days after tracheostomy procedure, patient’s ventilatory pressures needed to be increased and she developed worsening respiratory distress.
Acquired Bartter-like Syndrome associated with Colistin Use

Despite invasive mechanical ventilation, A. baumannii complex (10^5 colony) was isolated from endotracheal aspirate that is sensitive only to colistin. Intravenous colistin was initiated with a dose of 5 mg/kg/day and divided into two equal doses. The patient received amikacin with colistin, and meropenem therapy was discontinued.

Polyuria (6 mL/kg/hour) in the presence of normal blood pressure and hypokalemic metabolic alkalosis developed on the 3rd day of colistin treatment. The patient did not have any diarrhea or vomiting. Serum sodium, magnesium, calcium, and serum creatinine concentrations were within the normal range. Her blood test revealed serum potassium 2.5 mg/dL, albumin 25 g/L, and serum bicarbonate 29 mEq/L that increased to 35 mEq/L on the 4th day of colistin therapy. Polyuria with urine output was around 6–8 mL/kg/hour. Urine studies showed urine spot potassium was 25 mEq/L and urine potassium creatinine ratio was 2:3 mEq/mg. Urine density was 1003. Renal ultrasonography was normal. Colistin was stopped on the 4th day; meropenem and tigecycline were initiated. Daily potassium supplementation was given. Daily intravenous fluid supplementation was increased by 30%.

Two days after discontinuation of colistin, polyuria improved dramatically. Her metabolic alkalosis and hypokalemic alkalosis discontinued after 2 and 4 days, respectively. On the 23rd day of hospitalization, her pneumonia showed regression, tracheal aspirate cultures showed any bacterial growth, and she was discharged to the general ward. The daily urine output, serum potassium, bicarbonate and pH levels were shown in Figure 1.

**Discussion**

Pharmacological studies on nephrotoxicity mechanisms of colistin have been neglected because it was off the market for a long time. Renal damage due to colistin is considered a result of its detergent activity for many years, but it induces membrane permeability, increases intracellular cations, anions, and water, and results in cell lysis, which have been proposed to be due to colistin’s nephrotoxicity mechanism. In animal studies, antioxidant therapies have shown promising results in protecting the kidney. Colistin nephrotoxicity is defined as an increase in serum creatinine to a level of >2 mg/dL, a 50% reduction in GFR, or a decline in renal function leading to a need for renal replacement therapy. It is usually reversible.

Bartter syndrome (BS) is an autosomal recessive disease characterized by hypokalemia, hypochloremia, metabolic alkalosis, and hypercalciuria. Renin and aldosterone levels are increased, but blood pressure is normal. Primary pathology in BS is in the thick ascending limb of the loop of Henle. As a result of the influence of Na/K/2Cl, apical K, and basal chloride channels in the membrane, renal wasting of Na, K, and Cl occurs. Volume depletion also develops. BLS develops during treatment with different classes of drugs, including tuberculostatics and aminoglycosides, and is characterized by hypokalemic metabolic alkalosis, hypomagnesemia, hypocalcemia, and normal serum creatinine levels. It is unclear how prolonged use of colistin can cause such tubulopathy.

In recent case reports, BLS due to colistin therapy has been reported in only one child case who is a preterm infant. Following the literature, a total of five cases who developed BLS after colistin therapy have been reported; however, only two cases reported having polyuria after a short time (3–4 days) of colistin therapy. The association of colistin with hypokalemia was reported previously. We know about aminoglycoside-induced BLS. Colistin may directly activate the calcium-sensing receptor in the thick ascending loop of Henle and the distal tubule similar to aminoglycosides and result in hypokalemia, metabolic alkalosis, hypocalcemia, hypomagnesemia, and hypercalciuria.

In our case, BLS developed on the 3rd day of colistin treatment, and she had already received amikacin for 12 days; symptoms disappeared only after colistin was discontinued. Also, amikacin treatment was continued after colistin, making it a causative factor, and not taking any diuretics suggests that colistin is the causative drug. In recent case reports, the onset of BLS after starting

![Fig. 1: Graphical representation of serum potassium, serum bicarbonate, and 24-hour urinary output trend. Day 12: Colistin started; Day 15: Colistin stopped](image)
Acquired Bartter-like Syndrome associated with Colistin Use

Acquired Bartter-like Syndrome associated with Colistin Use was observed in our patient. Colistin therapy was, respectively, 3 days, 4 days, 9 days, 2 weeks, and 4 weeks. Only hypokalemia was developed with polyuria in the patient, and other electrolytes and serum creatinine levels were normal. We did not continue colistin because of significant hypokalemia and polyuria, and the patient also showed a remarkable improvement in lung function and other markers of sepsis after changing the treatment.

CONCLUSION

Our case received colistin 5 mg/kg/day in two doses, and there is no literature about the dosage and frequency of colistin administration for tubulopathies beside serum creatinine levels. The onset of the metabolic alterations during treatment with colistin and resolution after interruption suggest a causative relationship. More pharmacological studies are needed about colistin’s nephrotoxicity and tubulopathy in critically ill children.

HIGHLIGHTS

Colistin should be considered as the causative agent for BLS. BLS due to colistin therapy has been reported in a total of five cases, and only one of them was a child, that is, a preterm infant.

Authors’ Contributions

D.P.Y., F.E., O.O.H., O.O.G., B.A., and D.Y. treated the patient and were responsible for the concept, design, and acquisition of data. D.P.Y. wrote the first draft of the manuscript; F.E., O.O.H., and D.Y. revised and edited the manuscript. O.O.G. and B.A. collected the data and were responsible for the literature search. All authors read and approved the final version of the manuscript.

Consent

Written informed consent was obtained from the patient for publication of this case report and any accompanying images.

Ethical Standard

We confirm that we have read the Journal’s position on issues involved in ethical publication and affirm that this report is consistent with those guidelines. All authors approved the final article.

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