Neonatal hyperbilirrubinemia and prematurity as cause of green deciduous teeth.

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Abstract

The green pigmentation of the teeth is a rare consequence of neonatal hyperbilirrubinemia. About 50 cases of green dental discolouration have been registered in literature, most of which affected children with hyperbilirrubinemia caused by biliary atresia. Bilirubin can be deposited in multiple tissues. The pigmentation of skin and mucous membranes is caused by the high cell turnover, contrary to mineralized tissues as dentin and enamel where bilirubin is permanently trapped.

The severity and duration of jaundice do not directly correlate with dental changes intensity and tooth discolouration. There are additional predisposing risk factors to the eruption of green teeth in later infancy.

This article presents a case involving a 3 years old child with green primary teeth who had a history of neonatal cholestasis and a prolonged conjugated hyperbilirrubinemia associated to Alagille syndrome.

Keywords: Green teeth, Hyperbilirrubinemia, Neonatal cholestasis.

Accepted April 12, 2017
During the first hours of life in the patient, generalized hypotonia was observed along with hypotension, requiring dopamine as ionotropic support. Facial anomalies such as epicanthus, hypertelorism and broad front were observed. During her admission, it presented a sustained low level of serum bicarbonate (minimum 11.5 mmol/L) and metabolic acidosis and the following analytical results were obtained: urea 22.3 mmol/L, creatinine 274 mcmol/L, potassium 6 mEq/L, 124 mEq/L sodium, 1.85 mmol/L calcium, 91 mEq/L chlorides, total protein 33 g/L. As a result, peritoneal dialysis was needed from the third day of life, maintaining it up to 21 days. The renal Doppler ultrasound showed peripheral and bilateral arterial stenosis. The isotopic renogram indicated reduced cortical uptake. Throughout her evolution she presented a Disseminated Intravascular Coagulation State (DIC) requiring several transfusions of packed red blood cells, fresh plasma and platelets. From the respiratory point of view she needed mechanical ventilation for 28 days. The cardiologic examination revealed the presence of pulmonary artery stenosis along with foramen ovale with left to right shunt. High levels of direct bilirubin (427.5 mcmol/L, N: 0-3.4 mcmol/L) and of ferritin (6,600 mcg/L, N: 25-200 mcg/L) were observed before the peritoneal dialysis was removed, maintaining a cholestatic syndrome for several months, which needed treatment with chelating agents and vitamins. The patient suffered Enterobacter cloacae sepsis at 24 days of life, which increased levels of GGT, bilirubin and factor V.

With the available data up to that time, it was thought that there was neonatal haemochromatosis which was treated with a total blood transfusion and intravenous immunoglobulin (1 g/kg). The hepatic Doppler ultrasound showed mild intrahepatic biliary dilatation. Liver scanning showed delayed biliary removal and liver biopsy presented a shortage of bile duct, without evidence of hemosiderin deposits. Regarding hydroelectrolytes, potassium, sodium and chloride levels were normal, but she needed oral administration of 1.25 hydroxyalkyl-calciferol due to hypocalcemia (1.25 mmol/L) and hyperphosphatemia (3.68 mmol/L). High blood pressure (154/86 mmHg) required administration of losartan (0.3 mg/kg/day) and captopril (0.5 mg/kg/8 h).

At discharge, the patient was 3 months old; she presented icteric colouration and 3 cm hepatomegaly without splenomegaly. She weighed 3,200 g. Treatment with ursodeoxycholic acid and phenobarbital is given during several weeks after discharge. Liver enzyme values were tested at the periodic follow-ups and GGT, ALT and AST were all normal. In the genetic study of our patient, a heterozygous duplication was confirmed in the exon 6 area of the JAG1 gene located at 20p12.1-p11.23 for Alagille syndrome with an autosomal dominant mode of inheritance.

At 3 years of age, the patient presented profound hearing loss in addition to chronic renal failure at the predialysis stage, pending a kidney transplant. The intraoral exam showed a dark green pigmentation of her fully erupted primary teeth, although the jaundiced skin had disappeared (Figure 1).
Discussion

The exact prevalence of the association of hyperbilirubinemia and green spotted teeth is unknown. In the literature we can find about 50 cases of green dental discoloration, most of which linked with biliary atresia. The fact that conjugated bilirubin is more soluble in water and is capable of being incorporated during the formation of the developing tooth matrix in dentin and enamel could be a reason for this.

Our patient had a medical history of prolonged conjugated hyperbilirubinemia caused by anatomic alterations in the biliary tract. Infections or metabolic, toxic, genetic and anatomic abnormalities can be the causes of neonatal through mechanical obstruction of bile flow or functional impairment of any processes involved in hepatic excretory function and bile secretion. Few cases of green teeth have been found linked with non-conjugated hyperbilirubinemia that can be a result of Rh incompatibility, ABO incompatibility, hemolytic disease, and sepsis. In a review of 48 children with green stained teeth secondary to hyperbilirubinemia the total serum bilirubin level was 20 to 90 mg/dL and the average duration of jaundice was 24.6 weeks [5]. However, green staining can not directly correlate with the duration and intensity of jaundice; it seems to be other factors that determine the appearance of this staining, such as the density of the matrix or alterations in the absorption of phosphate, calcium and fat-soluble vitamins. Low birth weight or extreme preterm birth may be additional risk factors [6]. Primary teeth are the most affected, although permanent dentition staining has been observed related to enamel hypoplasia as a consequence of biliary atresia [7].

Differential diagnosis of teeth discolouration includes extrinsic causes such as chromogenic bacteria or medicaments and intrinsic causes such as dentinogenesis imperfecta, amelogenesis imperfecta, tetracycline, and congenital erythropoietic porphyria.

Parents can be unsettlingly surprised by the appearance of a discolored tooth. Therefore, it is important for families with children with severe hyperbilirubinemia to have access to anticipatory guidance. Treatment of pigmented teeth is mainly cosmetic and its importance increases as the child grows older. In spite of this, since primary stained dentition is most often replaced with permanent teeth, primary teeth treatment is often not advisable.

References


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