Reintroduction of Diazoxide after Diagnosis of Pulmonary Hypertension in a Patient with Transient Hyperinsulinism

Bahareh Schweiger1 Pedro A. Sanchez-Lara1 Dor Markush2 Pooja Nawathe1

1Pediatrics Department, Cedars-Sinai Hospital, Los Angeles, California, United States
2Guerin Congenital Heart Program, Smidt Heart Institute, Cedars-Sinai Medical Center, Los Angeles, California, United States

Abstract

Our case describes the reintroduction of diazoxide despite life-threatening pulmonary hypertension in our infant due to lack of therapeutic options for congenital hyperinsulinism.

Keywords
- congenital hyperinsulinism
- diazoxide
- pulmonary hypertension
- chlorothiazide
- side effects

Introduction

A 37-day-old 35-week premature infant with normal APGARs at birth presented to the endocrinology clinic for follow-up after being on diazoxide 13 mg/kg/day divided every 8 hours for hyperinsulinism (HI). On presentation to the clinic, a rapid response code was called as the patient appeared cyanotic and had abdominal distension and suspected necrotizing enterocolitis. He was urgently intubated for impending respiratory failure. Chest radiograph showed cardiomegaly and mild pulmonary edema. The initial echocardiogram showed a small ductus arteriosus with exclusive right-to-left shunting consistent with severely elevated (suprasystemic) pulmonary artery pressures, in addition to moderate right heart enlargement. This was in contrast to the neonatal intensive care unit (NICU) discharge echocardiogram from a month prior, where no ductus arteriosus was visualized and pulmonary pressures were estimated as only mildly elevated.

After admission, the patient remained on high ventilator settings requiring the addition of nitric oxide and had recurrent right upper lobe collapse likely from right atrial enlargement needing therapeutic bronchoscopy twice to maintain appropriate gas exchange. The patient continued to be sedated, paralyzed on mechanical ventilation, inhaled nitric oxide, diuretics, antibiotics, with multiple pulmonary hypertensive crises. The B type natriuretic peptide on admission was elevated at 2,252 picogram/milliliter (normal is <100 pg/mL) and improved to 242 pg/mL by day 5, predicted time to diazoxide elimination. He was transitioned from nitric oxide to oral sildenafil (for ongoing pulmonary vaso-dilation) on day 8 and extubated on pediatric intensive care unit day 18. Diazoxide was discontinued since it was thought to be the inciting agent for the development of significant pulmonary hypertension (PH), but the patient continued to need high glucose infusion rates through total parenteral nutrition to maintain normoglycemia. Since the patient was critically ill and requiring a high glucose infusion rate, and...
surgery was entertained as a possible option, a decision was made to obtain imaging with a Dopa positron emission tomography scan, which showed findings compatible with diffuse islet cell hyperplasia. A karyotype and chromosome microarray were normal ([arr(1–22)x2, (XY)x1]). Methylation testing for Beckwith-Wiedemann syndrome [IC1 (H19) IC2 (LIT1)] was also normal. Rapid exome trio sequencing was completed and was non-diagnostic. Targeted ABCB8 sequencing with deletion/duplication testing identified only a single heterozygous missense variant of uncertain significance in the ABCB8 gene (c.1252T > C (p.Cys418Arg)). This variant has been reported in both adult-onset diabetes and HI, yet pathogenicity has not been fully established.

Due to the patient’s abdominal distention and suspected necrotizing enterocolitis, treatment with octreotide was not thought to be a safe alternative because of potential for serious gastrointestinal side effects and also its risk for PH. After a multidisciplinary discussion with HI experts due to the options of glucagon infusion gastrostomy tube with serious gastrointestinal side effects and also its risk for PH. Diazoxide was successfully discontinued at that time. Blood glucose remained greater than 70 mg/dL on diazoxide dose of 11.38 mg/kg/day and his diazoxide was weaned 5 days after initiation of diazoxide therapy, and continued thereafter as routine surveillance during the treatment. It has been suggested that all infants being considered for treatment with diazoxide receive an echocardiogram prior to and 5 days after initiation of diazoxide therapy.

In September 2015, the FDA released a statement warning that PH had been reported in 11 cases of newborns and infant treated with diazoxide over many years. In retrospective studies of HI at congenital HI centers treated with diazoxide, development of PH with its use is a known but rare adverse event and after its initiation was observed in 2 to 4.8% of patients. In addition to risk factors including prematurity, respiratory failure, and congenital heart disease, delayed chlorothiazide initiation was a risk factor for PH. Most diazoxide-induced PH are thought to be reversible after discontinuation of diazoxide unless there is pre-existing PH. Cases of patent ductus arteriosus reopening have been reported, and sodium and water retention with an increase in pulmonary volume and pressures have been postulated to be related to reopening of the ductus arteriosus and worsening PH.

In infants, the incidence of PH may be higher as prevalence of diazoxide exposure in the NICU has increased along with the frequency of hypoglycemia as a diagnosis for infants in the NICU, doubling in the last 6 years being attributed to potentially the American Academy of Pediatrics in 2011 on criteria and treatment of hypoglycemia in newborns. Furthermore, an echocardiogram is not currently routinely ordered after the initiation of diazoxide. More recently it has been suggested that all infants being considered for diazoxide therapy receive an echocardiogram prior to and 5 days after initiation of diazoxide therapy, and continued thereafter as routine surveillance during the treatment. It has been also recommended that concurrent treatment with a thiazide diuretic be implemented. Also, parents should be informed about this possible side effect and to report any symptoms that might raise suspicion for PH, such as respiratory distress, poor feeding, pallor, or cyanosis. It has been more recently found that serious adverse events including PH are significantly higher in newborns with perinatal stress HI than that of otherwise healthy babies with genetic forms of HI. This suggests that more caution should be used when prescribing diazoxide in this population as well as other drugs that have been associated with increased risk of PH in the newborn period.

The mechanism of diazoxide-induced PH is unclear but direct toxic vascular reaction and KATP channel agonism have been postulated and it has been reportedly dose related.

Discussion

Diazoxide is benzothiadiazide derivative previously studied as an antihypertensive medication and approved by the U.S. Food and Drug Administration (FDA) for use in children and infants for a specific subset of conditions, including symptomatic hyperinsulinemic hypoglycemia and the only therapy approved by the FDA for congenital HI. Diazoxide works by activation of the ATP-sensitive potassium channels. It suppresses insulin release from the pancreatic β-cell by maintaining a hyperpolarized plasma membrane hyperpolarized.
Conclusion

Our case describes the successful reintroduction of diazoxide in an infant despite life-threatening PH, undertaken due to lack of therapeutic options for congenital HI for this patient. This was achieved using a multidisciplinary approach involving cardiology, endocrinology, and the family, and undertaken using serial echocardiographic follow-up and close clinical surveillance.

Authors’ Contributions
P.N. has made substantial contributions to the conception and design of the work; and helped in drafting the work and substantively revised it and has approved the submitted version and has agreed both to be personally accountable for the author’s own contributions and to ensure that questions related to the accuracy or integrity of any part of the work, even ones in which the author was not personally involved, are appropriately investigated, resolved, and the resolution documented in the literature.

B.S. made substantial contributions to the conception and design of the work; and helped in drafting the work and substantively revised it and has approved the submitted version and has agreed both to be personally accountable for the author’s own contributions and to ensure that questions related to the accuracy or integrity of any part of the work, even ones in which the author was not personally involved, are appropriately investigated, resolved, and the resolution documented in the literature.

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Conflict of Interest
None declared.

References

7 FDA Drug Safety Communication FDA Warns about a Serious Lung Condition in Infants and Newborns Treated with Proglycem (Diazoxide) [press release]. Food and Drug Administration. Maryland, USA: 2015