Severe Hyperinsulinemic Hypoglycemia in a Neonate: Response to Sirolimus Therapy

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abstract

Hyperinsulinemic hypoglycemia (HH) is one of the most common causes of persistent hypoglycemic episodes in neonates. Current pharmacologic treatment of neonatal HH includes diazoxide and octreotide, whereas for diffuse, unresponsive cases a subtotal pancreatectomy may be the last resort, with questionable efficacy. Here we report a case of congenital diffuse neonatal HH, first suspected when severe hypoglycemia presented with extremely high serum insulin levels immediately after birth. Functional imaging and genetic tests later confirmed the diagnosis. Failure to respond to a sequence of different treatments and to avoid extensive surgery with predictable morbidity prompted us to introduce a recently suggested alternative therapy with sirolimus, a mammalian target of rapamycin inhibitor. Glucose intake could be reduced gradually while euglycemia was maintained, and we were able to achieve exclusively enteral feeding within 6 weeks. Sirolimus was found to be effective and well tolerated, with no major adverse side effects attributable to its administration.

PATIENT PRESENTATION

Our patient, a male infant, was born by normal vaginal delivery during the 37th week of gestation after an uneventful pregnancy. Birth weight was 4400 g, and 1- and 5-minute Apgar scores were 10 and 10. Both parents and the baby’s 2 older siblings had unremarkable medical histories. A right-sided clavicular fracture was noted on first examination, which did not warrant additional intervention. Two hours after delivery, tremor and irritability were observed, with severe hypoglycemia (0.5 mmol/L, 9 mg/dL). High doses of intravenous glucose (up to 20 mg/kg per minute) and occasional glucagon boluses were needed to normalize the persistently low blood glucose levels. The diagnosis of hyperinsulinemic hypoglycemia (HH) was based on the clinical picture and on the laboratory values (glucose, 0.5 mmol/L; concomitant serum insulin, 130.7 μU/mL; growth hormone, 18.3 ng/mL; thyroid-stimulating hormone, 8.2 μU/L; free thyroxine, 20.5 pmol/L; cortisol, 835 nmol/L), and samples were sent for genetic testing.

Diazoxide therapy (with hydrochlorothiazide 1 mg/kg per day) was commenced on day 4 and was gradually increased to a maximum dose of 20 mg/kg per day without any effect. Somatostatin treatment (introduced as intravenous, subsequently modified to subcutaneous) was initiated on week 2 and increased to a maximum dose (35 μg/kg per day), but only a 20% reduction in total glucose requirements was achieved. On week 4 nifedipine was added to the therapeutic regimen, but it was discontinued after a week because of lack of response (Fig 1).
A dose of 0.5 mg/m² per day, with initiated on week 8, built up from a Oaks, Surrey, United Kingdom) was gradually be discontinued, while the

pancreatectomy. A recent report\(^3\) of were organized regarding treatment discussions, involving the parents, unaffected mother. Multidisciplinary (mTOR) inhibitor in similar cases, was a mammalian target of rapamycin (mTOR) inhibitor in similar cases, was a novel indication of sirolimus, we could not attribute any significant and triglyceride and cholesterol levels mildly elevated liver function tests collected to monitor for known side effects of sirolimus, but apart from adverse events to the treatment. Regular blood (complete blood cell count, renal and liver function, lipid levels) and urine samples were collected to monitor for known side effects of sirolimus, but apart from mildly elevated liver function tests and triglyceride and cholesterol levels we could not attribute any significant adverse events to the treatment. Clinical neurologic examinations were unremarkable during the period of treatment apart from the initial tremor witnessed immediately after birth. Neurodevelopmental assessment revealed moderate axial hypotonia, which improved with physiotherapy. A brain MRI scan at 5 months was reported as normal. Weight and head circumference were both above the 98 percentile at birth and remained in this range throughout the period of observation. Although oral intake has remained the primary route for feeding, a percutaneous endoscopic gastrostomy tube was inserted at 6 months to aid supplemental feeding at regular intervals.

Follow-up assessment at the age of 1 year confirmed that the treatment had sustained results. The infant, weighing 13.5 kg, currently receives 1.5 mg sirolimus once a day, which maintains a serum sirolimus level of 9.62 ng/mL. Capillary blood sugar is checked at least ≥4 times every day, with levels ranging between 3.5 and 7.0 mmol/L, and no hypoglycemic episode (serum glucose <2.6 mmol/L) has been reported since discharge. Controlled fasting levels significantly improved from <1 hour at the time of diagnosis to >6 hours at follow-up (insulin levels were 20 μU/mL and 5.52 μU/mL after 4 and 6 hours of fasting, respectively). The liver function tests and serum lipid levels remained mildly elevated (aspartate aminotransferase, 73 U/L; alanine aminotransferase, 65 U/L; cholesterol, 5.9 mmol/L; triglyceride, 3.9 mmol/L) but showed improvement, with the reduction of sirolimus levels, and continue to be closely monitored.

**DISCUSSION**

HH in neonates, formerly also described as nesidioblastosis or congenital hyperinsulinism, is characterized by unregulated and elevated insulin secretion of the pancreatic islet β cells, resulting in persistent hypoglycemia. This is a rare disease with an incidence of 1 in 50 000 live births, but significant variation has been reported.\(^4\) Early diagnosis and treatment are crucial to the appropriate management of HH. Histologic presentation may be focal or diffuse\(^5\) (and very rarely atypical\(^6\)) based on the spread of affected regions in the pancreas. Differentiation between these subgroups can be made by using \(^{18}\)F 3,4-dihydroxyphenylalanine positron emission tomography scans.\(^7\) Several congenital mutations have been described that may have a role, and in ~50% of the cases the adenosine triphosphate (ATP) sensitive K⁺ channel in pancreatic β cells seems to play a key role in the uncontrolled insulin secretion.\(^8\) 

Functional imaging with \(^{18}\)F 3,4-dihydroxyphenylalanine positron emission tomography scan performed on week 6 confirmed diffuse pancreatic enhancement. Genetic tests subsequently showed a compound heterozygote for the known mutations in the ABCC8 gene: mutation p.Y512* (c.1536C→A M/n)\(^1\) inherited from the unaffected father and IVS36-13G>A (c.4412-13G>A, p.? M/n)\(^2\) inherited from the unaffected mother. Multidisciplinary, including the parents, were organized regarding treatment options, including a subtotal pancreatectomy. A recent report\(^3\) of a novel indication of sirolimus, a mammalian target of rapamycin (mTOR) inhibitor in similar cases, was also discussed. After risks and benefits were considered carefully, a consensus decision was made to proceed with this alternative treatment option. Approval was sought from Directorate General of National Institute of Pharmacy, Hungary, and written consent from the parents was also obtained.

Sirolimus (Rapamune; Pfizer, Walton Oaks, Surrey, United Kingdom) was initiated on week 8, built up from a dose of 0.5 mg/m² per day, with gradual increments controlled by serum levels (aiming for 5–15 ng/mL) checked every 3 to 5 days (see Supplemental Information). During the next 6 weeks the patient’s glucose requirement was significantly reduced for the first time (to 8 mg/kg per minute), and somatostatin could gradually be discontinued, while the patient maintained normoglycemia. During this transitional period, an episode of sepsis with bloody stool developed at week 11, with signs of intestinal pneumatosis on the abdominal radiograph. Bell stage II A necrotizing enterocolitis (NEC) was diagnosed, enteral feed was briefly suspended, and the symptoms resolved with conservative treatment.

A brain MRI scan at 5 months was reported as normal. Weight and head circumference were both above the 98 percentile at birth and remained in this range throughout the period of observation. Although oral intake has remained the primary route for feeding, a percutaneous endoscopic gastrostomy tube was inserted at 6 months to aid supplemental feeding at regular intervals.

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KCNJ11 and ABCC8 localized on chromosome 11p15.1 are responsible
for coding the 2 subunits of these K-ATP channels.9

The primary goal of treatment is to achieve normoglycemia to prevent long-term neurologic damage.10 The treatment algorithm most widely accepted recommends the K-ATP agonist diazoxide as first-line therapy and the somatostatin analog octreotide as second-line therapy.11 The calcium channel blocker nifedipine has also been reported as an alternative in unresponsive cases.12 Focal lesions usually warrant surgical excision, whereas for therapy-resistant diffuse presentations the only treatment option available is subtotal or total pancreatectomy.13 Definitive success rates are high for the former, but about one-third of patients remain hyperinsulinemic after subtotal pancreatectomy, many develop exocrine pancreatic insufficiency, and nearly all will develop diabetes by adolescence.14

A recently published novel treatment option for HH of the newborn has been suggested including the immunosuppressant sirolimus, an mTOR inhibitor.3 No major adverse reactions were observed during the 1-year follow-up period in the 4 cases reported. This finding is reinforced by the latest case report of another infant.15 The suggested method of action of sirolimus includes the reduction of β cell proliferation, inhibition of insulin production,16 and induced peripheral insulin resistance.17

Our decision to proceed with this treatment option was prompted by the lack of response to standard therapies, with the aim to avoid the major adverse effects of subtotal pancreatectomy. The desired glycemic control was achieved within 6 weeks of initiating sirolimus treatment. A transient episode of NEC was observed after the introduction of sirolimus, which responded well to conservative treatment and subsequent termination of somatostatin therapy. Based on previously published evidence of somatostatin-related NEC,18 it was considered most likely that this adverse event was attributable to somatostatin rather than sirolimus. Although the side effects of elevated liver function, hypercholesterolemia, and hypertriglyceridemia were observed, they were mild, and improvement was seen with reduction in sirolimus dosage. Significant renal, hematologic, or pulmonary complications associated with sirolimus reported elsewhere19 were not experienced. Compliance with dietary and therapeutic recommendations took patience and full engagement with the parents.

CONCLUSIONS

We believe our case report adds to the growing evidence of published cases suggesting that sirolimus could be an effective and safe treatment option in therapy-resistant diffuse HH of the newborn.

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ABBREVIATIONS

ATP: adenosine triphosphate
HH: hyperinsulinemic hypoglycemia
mTOR: mammalian target of rapamycin
NEC: necrotizing enterocolitis

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