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Clinical and molecular characterisation of hyperinsulinaemic hypoglycaemia in infants born small-for-gestational age

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ABSTRACT

Objective To characterise the phenotype and genotype of neonates born small-for-gestational age (SGA; birth weight <10th centile) who developed hyperinsulinaemic hypoglycaemia (HH).

Methods Clinical information was prospectively collected on 27 SGA neonates with HH, followed by sequencing of KCNJ11 and ABCC8.

Results There was no correlation between the maximum glucose requirement and serum insulin levels. Serum insulin level was undetectable in five infants (19%) during hypoglycaemia. Six infants (22%) required diazoxide treatment >6 months. Normoglycaemia on diazoxide <5 mg/kg/day was a safe predictor of resolved HH. Sequencing of KCNJ11/ABCC8 did not identify any mutations.

Conclusions Serum insulin levels during hypoglycaemia taken in isolation can miss the diagnosis of HH. SGA infants may continue to have hypofattyacidaemic hypoketotic HH beyond the first few weeks of life. Recognition and treatment of this group of patients are important and may have important implications for neurodevelopmental outcome of these patients.

INTRODUCTION

Hyperinsulinaemic hypoglycaemia (HH) is characterised by the unregulated secretion of insulin from pancreatic β-cells in relation to the blood glucose concentration. It is the commonest cause of severe and persistent hypoglycaemia in the neonatal and infancy periods.1 The genetic basis of congenital forms of HH involves defects in key genes (com- monen—ABCC8 and KCNJ11 encoding the two subunits SUR1 and Kir6.2, respectively), which regulate insulin secretion from pancreatic β-cells.1 HH may be secondary to certain risk factors (such as maternal diabetes mellitus, perinatal asphyxia and small-for-gestational age (SGA)). In 1984, Collins and Leonard first described transient HH in SGA infants.2 It has not previously been documented whether the HH observed in infants born SGA is due to defects in the ABCC8/KCNJ11 genes. The phenotype of a large cohort of SGA neonates who developed HH has not been studied previously. The purpose of the present study was to report on the phenotype, clinical course and results of the sequencing of ABCC8/KCNJ11 in a cohort of infants born SGA who developed HH.

PATIENTS AND METHODS

We prospectively studied the clinical and biochemical features of 27 SGA (defined as birth weight <10th centile) neonates with HH who were admitted to two tertiary paediatric referral centres (Great Ormond Street Hospital and Bristol Royal Hospital for Children). Neonates with history of perinatal asphyxia, Rhesus isoimmunisation and syndromic forms such as Beckwith-Wiedemann syndrome were excluded from the study. All were investigated at the time of clinical and biochemical hypoglycaemia to establish the diagnosis of HH and were commenced on diazoxide (>5–20 mg/kg/day). Diazoxide responsiveness was defined as ability to come off intravenous glucose and maintain normoglycaemia during a physiological fast. Following discharge, they were followed up at 3-monthly intervals in outpatient clinic, where glycaemic control and diazoxide dose were reviewed. These infants were readmitted for a 24-h glucose profile and a controlled fast after stopping diazoxide for 3 days, if home glucose monitoring indicated no recurrence of hypoglycaemia on <5 mg/kg/day of diazoxide.1 If home glucose monitoring revealed hypoglycaemic episodes, the dose of diazoxide was adjusted and the trial off diazoxide was postponed until the dose was <5 mg/kg/day.
Glucose profile and fasting responses at resolution of hyperinsulinism

In all, 17 infants (63%) underwent a 24 h glucose profile and controlled fast after stopping diazoxide to demonstrate resolution of hyperinsulinism. None of these infants recorded hypoglycaemia during a 24-h glucose profile. All infants maintained blood glucose concentrations above 3.0 mmol/l at the end of fast appropriate for their age, with 13 infants maintaining blood glucose concentrations >3.5 mmol/l. All infants showed appropriate plasma insulin suppression and elevation of plasma β-hydroxybutyrate/non-esterified fatty acids as evidence of resolution of HH.

Genetic results

Sequencing did not reveal mutations in ABCC8/KCNJ11 genes in 25/27 DNA samples analysed.

DISCUSSION

This report is the first case series of SGA neonates with HH that reports the clinical characteristics and ABCC8/KCNJ11 sequencing results in a cohort of 27 subjects. Collins et al first described transient HH in three SGA infants and subsequently reported HH in five of 27 SGA neonates studied, suggesting it is common in SGA neonates. It is important to recognise this as neonates with HH are especially prone to the complications of hypoglycaemia because of their inability to generate alternative fuels, such as ketone bodies.

In our study, despite other evidence of hyperinsulinism, plasma insulin levels were undetectable during hypoglycaemia in 5/27 SGA neonates. We did not find any correlation between the maximum glucose infusion rate required to maintain normoglycaemia and plasma insulin levels. This is explained by the short half-life of insulin of 4–5 min and rapid clearance by the liver before it reaches peripheral circulation. In a study by Hoe et al, plasma insulin was inappropriately elevated in only 11 of 24 neonates who had transient prolonged neonatal hyperinsulinism. Our results corroborate these findings and highlight the value of other parameters (apart from plasma insulin) in the diagnosis of HH.

In this case study of SGA infants referred to two tertiary hospitals with HH, all infants responded to diazoxide, though the dose was variable ranging from 5 to 20 mg/kg/day. Diazoxide was successfully stopped in 21 (78%) patients by 6 months of age. In the remaining six infants, five needed diazoxide until 10 months of age and one infant needed prolonged treatment until 22 months. In a study of 26 neonates with hyperinsulinism which included 7 SGA neonates, HH resolved by a median age of 181 days (range, 18–403 days). Fafoula et al also reported prolonged transient HH up to 9 months in SGA infants.

In our cohort, no recurrence of HH was noticed after stopping diazoxide. This highlights that with the management protocol whereby diazoxide was stopped when good glycaemic control was maintained with diazoxide dose <5 mg/kg/day, the risk of recurrence of hypoketotic hypoglycaemia brain injury to these vulnerable infants is avoided.

To our knowledge, no previous study has reported on the genetic results of SGA neonates who developed HH. In our study, no mutations in ABCC8/KCNJ11 were identified in these SGA infants. However, rarer genetic causes of HH due to mutations in GLUD1, HADH, GCK and HNF4A were not excluded.

CONCLUSIONS

SGA infants may continue to have hypofattyacidemic hypoketotic HH beyond the first few weeks of life. Recognition and
treatment of this group of patients are important and may have important implications for neurodevelopmental outcome of these patients. Plasma insulin levels during hypoglycaemia taken in isolation can miss the diagnosis of HH and hence other parameters (such as glucose infusion rates and production of free fatty acids/ketone bodies during hypoglycaemia) must be considered in the diagnosis. Withdrawal of diazoxide treatment when glycaemic control is maintained at a dose of <5 mg/kg/day can be safely recommended.

Finally, the genetic aetiology of HH in SGA infants is not understood and, in this study, mutations in the common genes implicated in the aetiology of congenital hyperinsulinism were not identified. Further studies are required to understand the underlying mechanism of HH in these infants.

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Contributors VBA collected and analysed the data and wrote the manuscript. SEF performed the genetic studies and reviewed the manuscript. AK and JPS collected the data and critically reviewed the manuscript. SE and KH conceptualised the study and critically reviewed the manuscript. All authors contributed to the approval of the final version of the manuscript. RRK is the guarantor of this work.

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Competing interests None.

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