Long-term response to sulfonylurea in a patient with diabetes due to mutation in the \textit{KCNJ11} gene

Marcio F. Vendramini, Lucimary C. Gurgel, Regina S. Moisés

ABSTRACT

Objective: To report the long-term (30-month) effect of the switch from insulin to sulfonylurea in a patient carrying the p.G53D (c.158G>A) mutation in \textit{KCNJ11} gene. Subject and method: A 29-year-old male patient was diagnosed with diabetes in the third month of life and after identification of a heterozygous p.G53D mutation in the \textit{KCNJ11} gene, the therapy was switched from insulin to sulfonylurea. Results: Long-term follow-up (30 months) showed that good metabolic control was maintained (HbA1c: 6.6%) and the glibenclamide dose could be reduced. Conclusion: Long-term therapy with sulfonylureas in patients with neonatal diabetes due to mutation in the \textit{KCNJ11} gene is safe and promotes sustained improvement of glycemic control.

Keywords

Neonatal diabetes; \textit{KCNJ11} mutation; sulfonylurea therapy

INTRODUCTION

Neonatal diabetes mellitus is a rare condition characterized by hyperglycemia within the first months of life, which may either be permanent or transient. Activating mutations in the genes \textit{KCNJ11} and \textit{ABCC8}, which encode the two subunits Kir6.2 and SUR1 of the pancreatic ATP-sensitive potassium channel, are the most frequent cause of permanent neonatal diabetes (1-3). Approximately 30% of individuals with activating mutations in \textit{KCNJ11} have neurologic features in addition to neonatal diabetes (4). The main neurologic findings are developmental delay, muscle weakness and epilepsy and this has been proposed to be part of a syndrome referred as DEND (developmental delay, epilepsy and neonatal diabetes) syndrome (5). A less severe form without epilepsy is called intermediate DEND syndrome.

Successful switch from insulin to sulfonylurea (SU) therapy has been reported in the majority of the patients with \textit{KCNJ11} mutations (6-8); however, there are few reports on the long-term efficacy and safety of this change (9-12).

Here we report the long-term (30-month) effect of the switch from insulin to sulfonylurea in a patient carrying the p.G53D mutation in the \textit{KCNJ11} gene.
**PATIENT PRESENTATION**

The patient, who is currently 29 years of age, was diagnosed with diabetes in the third month of life during a ketoacidosis episode. Insulin treatment was initiated at that time and even on intensive insulin therapy, his glucose control was erratic with marked hyperglycemia. His growth curve was consistently along the 10th percentile for height. During follow-up a delay in motor milestones was observed: the proband started walking autonomously only at 28 months. No muscle weakness or lower limb hypotonia was present. Crisis of generalized seizures started at age 5 during episodes of hypoglycemia, mainly in nocturnal period. His EEG was normal. He also showed severe learning difficulties and very poor attention and despite receiving special education little improvement was achieved.

At the age of 26 years, informed consent for genetic analysis was obtained from the proband’s parents. A blood sample was obtained and genomic DNA was extracted from peripheral blood leukocytes using a commercial kit (Puregene DNA Isolation Kit, Gentra System, Minneapolis, MN, USA). The single exon of KCNJ11 was amplified by PCR in three overlapping fragments. The PCR products were directly sequenced with the use of Big Dye Terminator Cycle Sequencing Reaction Kit version 3.1 and analyzed on an ABI Prism 3100 Genetic Analyzer (Applied Biosystems, CA, USA). The proband was found to have a heterozygous change of guanine to adenine on nucleotide 158, which resulted in glycine to aspartic acid substitution at codon 53 (p.G53D, c.158G>A mutation). Both parents were negative for this mutation. On the basis of the genotype and clinical features consisting of neonatal diabetes with developmental delay but not epilepsy, the patient was given a diagnosis of intermediate DEND syndrome. Following the identification of the mutation the patient was transferred from insulin to glibenclamide. The initial response to SU therapy was previously reported (13). After 4 weeks the patient was independent of insulin and using 0.8 mg/kg/day of glibenclamide divided into three doses. During the following 3 months, as the patient experienced mild hypoglycemic episodes, the glibenclamide dose was reduced to 0.68 mg/kg/day. A 72-hour continuous glucose monitoring system (CGMS, Medtronic Minimed, Sylmar CA, USA) showed mean glucose of 108 mg/dL with 71% of glycemic values between 80 and 200 mg/dL and 1% values above 200 mg/dL (Figure 1A). HbA1c levels reduced from 8.2% to 7.6% 3 months after the transfer. A weight loss of 3.8 kg was observed in the first 8 weeks of SU therapy with subsequent weight stabilization. Currently, 30 months after the introduction of SU, good metabolic control is maintained (HbA1c: 6.6%) and the glibenclamide dose was reduced to 0.55 mg/kg/day. During the course of SU therapy no severe hypoglycemic episodes were reported. Another 72-hour continuous glucose monitoring was performed and showed mean glucose of 107 mg/dL with 80% of glycemic values between 80 and 200 mg/dL and none above 200 mg/dL (Figure 1B). Incipient diabetic nephropathy was diagnosed 9 years ago and therapy with ACE-inhibitors was initiated at that time. Fourteen months after the introducing SU, microalbuminuria in nocturnal collection of urine decreased from 77 ug/min to 32 ug/min and remained stable through the follow-up period.

The investigation was approved by the Ethics Committee of Escola Paulista de Medicina, Universidade Federal de São Paulo.

**Figure 1.** Glucose levels captured by the 72-h continuous glucose monitoring system 3 months (A) and 30 months (B) after switching from insulin to glibenclamide therapy. Each line represents a 24-h period.
DISCUSSION

Activating mutations in the genes encoding the ATP-sensitive potassium channel causes failure of this channel to close in response to increased intracellular ATP, thus inhibiting insulin secretion. However, sulfonylureas bind to the SUR1 subunit and close the ATP-sensitive channel by an ATP-independent mechanism resulting in insulin release even in cells with activated mutations in KCNJ11 (14).

We describe herein a 30-month experience of sulfonylurea treatment in an adult patient with intermediate DEND syndrome due to p.G53D mutation in KCNJ11. Following the transfer from insulin to SU we observed marked improvement in glycemic control that was maintained over the follow-up period, despite decreasing SU doses. No side effects were reported. The successful transfer to SU is consistent with the previous report of a patient with permanent neonatal diabetes carrying the same mutation (15). To examine the consequences of this mutation, Koster and cols. (15), coexpressed a mutant Kir6.2 subunit containing the p.G53D mutation with the wild-type SUR1 subunit. Channel activity in intact cells was screened by ⁸⁶Rb⁺ efflux and ATP sensitivity was measured directly in excised membrane patches (15). The authors have demonstrated an increased channel activity and reduced ATP sensitivity by approximately 20 fold in homomeric channels.

There are few reports of chronic diabetes complications in patients with neonatal diabetes and the role of SU treatment in their evolution. Klupa and cols. (10) reported progression of a pre-existing diabetic retinopathy in a patient whose treatment was transferred from insulin to SU (10). Despite no causal relationship between the retinal changes and SU treatment could be proven, the authors recommend special ophthalmological attention in subjects with pre-existing advanced diabetic retinopathy. Our patient had no diabetic retinopathy when sulfonylurea was initiated and no evidence of this complication was found during the follow-up period. Reduction in microalbuminuria was observed during SU therapy possibly related to the improvement of metabolic control. Klupa and cols. (12) also observed some regression in the albumin/creatinine ratio in one patient.

In summary, this case illustrates that switching from insulin to sulfonylurea is safe and promotes sustained improvement of glycemic control on long-term follow-up.

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REFERENCES