Neonatal Hypoglycemia, Early-Onset Diabetes and Hypopituitarism Due to the Mutation in *EIF2S3* Gene Causing MEHMO Syndrome

J. STANIK^{1,2,3}, M. SKOPKOVA², D. STANIKOVA^{1,2,4}, K. BRENNEROVA¹, L. BARAK¹, L. TICHA¹, J. HORNOVA¹, I. KLIMES², D. GASPERIKOVA²

¹Department of Pediatrics, Medical Faculty of Comenius University and Children Faculty Hospital, Bratislava, Slovakia, ²DIABGENE and Laboratory of Diabetes and Metabolic Disorders, Institute of Experimental Endocrinology, Biomedical Research Center, Slovak Academy of Sciences, Bratislava, Slovakia, ³Center for Pediatric Research Leipzig, University of Leipzig, Germany, ⁴Institute of Social Medicine, Occupational Health and Public Health, University of Leipzig, Germany

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Summary

Recently, the genetic cause of several syndromic forms of glycemia dysregulation has been described. One of them, MEHMO syndrome, is a rare X-linked syndrome recently linked to the *EIF2S3* gene mutations. MEHMO is characterized by Mental retardation, Epilepsy, Hypogonadism/hypogonitalism, Microcephaly, and Obesity. Moreover, patients with MEHMO had also diabetes and endocrine phenotype, but detailed information is missing. We aimed to provide more details on the endocrine phenotype in two previously reported male probands with MEHMO carrying a frame-shift mutation (I465fs) in the *EIF2S3* gene. Both probands had a neonatal hypoglycemia, early onset insulin-dependent diabetes, and hypopituitarism due to dysregulation and gradual decline of peptide hormone secretion. Based on the clinical course in our two probands and also in previously published patients, neonatal hypoglycemia followed by early-onset diabetes and hypopituitarism may be a consistent part of the MEHMO phenotype.

Key words

MEHMO syndrome • *EIF2S3* • Diabetes • Hypopituitarism

Corresponding authors

D. Gasperikova, DIABGENE and Laboratory of Diabetes and Metabolic Disorders, Institute of Experimental Endocrinology, Biomedical Research Center, Slovak Academy of Sciences, Dubravska cesta 9, 84505, Bratislava, Slovakia. E-mail:

daniela.gasperikova@savba.sk and J. Stanik, Department of Pediatrics, Medical Faculty of Comenius University, Limbova 1, 833 40 Bratislava, Slovakia. E-mail: juraj.stanik@savba.sk

Introduction

Monogenic diabetes is a heterogeneous group of disorders caused by a mutation of a single gene involved to the insulin secretion or action (Rubio-Cabezas *et al.* 2014). The highest prevalence of monogenic diabetes is in patients with neonatal (<6 months) and infancy onset (<1 year) diabetes mellitus (Rubio-Cabezas *et al.* 2014, Stanik *et al.* 2007). There are several subtypes of monogenic neonatal and infancy onset diabetes, i.e. 1) disorders with abnormal pancreatic development including pancreatic agenesis and imprinting abnormalities, 2) disorders with abnormal β -cell function, particularly including *INS*, *KCNJ11*, and *ABCC8* mutations, and 3) disorders causing destruction of β -cells by dysfunction in the unfolded protein response pathway (Walter and Ron 2011, Wang and Kaufman 2012) including recently published Ile465Serfs mutation in the *EIF2S3* gene causing the MEHMO syndrome (Skopkova *et al.* 2017). The *EIF2S3* gene encodes the γ subunit of eukaryotic translation initiation factor 2 (eIF2), crucial for initiation of protein synthesis and regulation of the integrated stress response. MEHMO syndrome

(OMIM# 300148) is characterized by X-linked intellectual disability, epileptic seizures, hypogonadism, hypogenitalism, microcephaly, and obesity (Leshinsky-Silver *et al.* 2002). Diabetes or endocrine phenotypes have not been included to the main signs of the MEHMO syndrome. Nevertheless, majority of the previously published MEHMO patients had hypoglycemia or diabetes and/or endocrine phenotype, but detailed information is missing. Therefore, we aimed to provide more details on the endocrine phenotype in two male probands with MEHMO carrying a frame-shift mutation (I465fs) in the *EIF2S3* gene (Skopkova *et al.* 2017).

Patients and Methods

We refer on two probands with MEHMO syndrome of Slovak origin. Glucose, serum C-peptide, and hormone levels were measured in local labs by standardized methods (Stanikova *et al.* 2015). HbA1c was evaluated from whole blood by LPLC DiaSTAT analyzer (Bio-Rad, Hercules, USA). All steps of this study were approved by the Faculty Hospital Ethics Committees in Bratislava. Parents of both patients had signed the informed consent.

Results

Diabetes and endocrine phenotypes of the affected probands with MEHMO syndrome are summarized in Table 1.

Patient 1 (MEHMO syndrome with diabetes and panhypopituitarism)

The proband is a second child of non-consanguineous parents of Slovak origin and the only male offspring in the mother's family. His mother who carried the same *EIF2S3* gene mutation as her son had sideroblastic anemia; father and older sister are healthy. The proband had prenatally diagnosed microcephaly at 28th week of gestation. He was delivered in 39+6 gestational week with birth length of 47 cm (-2.0 SDS) and weight 2,920 g (-1.4 SDS) and head circumference 32 cm (-2.6 SDS). Diagnosis of MEHMO was made at the age of 10 months based on the microcephaly, unstable partial complex epileptic seizures resistant to anticonvulsives, hypogenitalism, severe mental and motor delay, central obesity, and typical dysmorphic features (for photo see Skopkova *et al.* 2017). Currently he is 6 years old.

Glycemia and diabetes phenotype

Hypoglycemia <1 mmol/l firstly occurred at birth and was corrected with glucose infusion. Hypoglycemia (2.6 mmol/l) reoccurred at the age of 3 months when he was admitted to the hospital with lethargy, vomiting and hypotrophy (weight of 3,830 g, -2.9 SDS). Insulin and C-peptide serum levels were in normal range (Table 1) at that time. Diabetes (2 h oral glucose tolerance test glycemia 14 mmol/l; HbA1c 7.5 %; 58 mmol/mol) was diagnosed at the age of 10 months with mild polyuria but without ketoacidosis (pH 7.37, HCO₃ 26.1 mmol/l). Type 1 diabetes specific autoantibodies were negative. Fasting C-peptide was within normal range for normoglycemic subjects at the point of diagnosis, but decreased over the time (Table 1). The proband was treated by three daily doses with both regular and NPH insulin injections since the age of 1 year. Despite intensive glycemic monitoring he had poor glycemic control with large glycemic excursions, particularly during respiratory infections. The proband was lethargic in euglycemia and needed target glycemia levels between 10-15 mmol/l to be awake.

Endocrine phenotype

Panhypopituitarism with growth hormone deficiency, central hypothyroidism, central hypocorticism, low prolactin levels, and low gonadotropin levels with micropenis and cryptorchism was diagnosed at the age of 9 months. Since then he has been treated with substitution therapy of L-thyroxin and cortisol. Most of the peptide hormone levels decreased over the time (Table 1). The brain MRI at the age of 4 months showed a normally developed pituitary gland without any pathological changes. Other features: he suffers from recurrent respiratory tract infections.

Patient 2 (MEHMO syndrome with diabetes and partial hypopituitarism)

The second proband is a first child of non-consanguineous parents of Slovak origin, and he was born in 40+0 gestational week with length of 49 cm (-0.95 SDS) and weight 2,440 g (-2.65 SDS), and head circumference 29 cm (-5.1 SDS). MEHMO features were very similar to the first proband (Skopkova *et al.* 2017) including the unstable epileptic seizures resistant to anticonvulsives. The proband inherited the *EIF2S3* gene mutation from his asymptomatic mother.

Glycemia and diabetes phenotype

The second proband had also period of neonatal hypoglycemia corrected with glucose infusion. He was diagnosed with diabetes (fasting glycemia 14.7 mmol/l; HbA1c 8.7 % (72 mmol/mol)) at the age of 10 months without ketoacidosis. Type 1 diabetes specific autoantibodies were negative. Fasting C-peptide was within the normal range for normoglycemic subjects at the point of diagnosis, but decreased over the time. The proband was treated with three daily insulin injections (<0.3 U/kg/day), but did not achieve good glycemic control (Table 1).

Endocrine phenotype

Partial panhypopituitarism with growth hormone deficiency, low prolactin levels, low gonadotropin levels with micropenis and cryptorchism was diagnosed at the age of 9 months. The decline of peptide hormone levels with aging was similar to Patient 1 (Table 1).

The second proband died at the age of 3.5 years because of infection and cardiorespiratory failure.

Discussion

We reported on detailed diabetes and endocrine phenotype in two male probands with the MEHMO syndrome. Both patients developed non-autoimmune infant-onset diabetes with decline of endogenous insulin production with age. Moreover, one of them had decline also of all hypophyseal peptide hormones; the second proband had impaired growth, thyroidal, and prolactin axis (but not ACTH production). The decline of the peptide hormone levels could be explained by the severe impairment of protein synthesis caused by the causal Ile465Serfs mutation in the *EIF2S3* gene (Skopkova *et al.* 2017). All the patients carrying this mutation had similar phenotype including growth retardation and dysregulation of insulin secretion (Skopkova *et al.* 2017, Moortgat *et al.* 2016). In both our patients, neonatal hypoglycemia preceded diabetes onset later during the life. This is similar to congenital hyperinsulinism and maturity-onset diabetes caused by dysregulation of insulin secretion due

to mutations in genes *HNF1A* and *HNF4A* (Stanescu *et al.* 2012, Rozenkova *et al.* 2015). Nevertheless, in case of MEHMO, also growth hormone deficiency could participate to the pathogenesis of hypoglycemia. Early-onset diabetes and endocrine symptomatology can also be seen in majority of other disorders caused by increased endoplasmic reticulum stress or a dysfunction in the unfolded protein response pathway, i.e. syndromic form of intellectual disability and diabetes caused by mutations in the *PPP1R15B* (Kernohan *et al.* 2015, Abdulkarim *et al.* 2015), Wolcott-Rallison syndrome caused by mutations of the *EIF2AK3* gene (Delepine *et al.* 2000), Wolfram syndrome caused by mutations in the *WFS1* gene (Fonseca *et al.* 2005), diabetes and multisystemic neurodegeneration caused by mutations of the *DNAJC3* (Synofzik *et al.* 2014), and Microcephaly, Epilepsy, and Diabetes Syndrome (MEDS) caused by mutations of the *IER3IP1* gene (Abdel-Salam *et al.* 2012).

In conclusion, we reported on two patients with MEHMO syndrome with neonatal hypoglycemia followed by early onset diabetes, and hypopituitarism presumably due to dysregulation of protein synthesis and gradual decline of peptide hormone secretion. The genetic analysis of genes involved in protein translation and its regulation needs to be considered in patients with syndromic forms of diabetes and endocrine related syndromes.

Conflict of Interest

There is no conflict of interest.

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