

The University of Chicago Genetic Services Laboratories



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KCNJ11 analysis for permanent neonatal diabetes and hyperinsulinemia of infancy

Clinical Features:

- **Permanent neonatal diabetes:** Neonatal diabetes is hyperglycemia that presents at 0-6 months of age (rarely later) and requires insulin. Approximately 50-60% of cases are considered transient neonatal diabetes [OMIM #601410] and resolve within 18 months. The remaining cases are considered permanent neonatal diabetes (PNDM) [OMIM #606176]. These patients have low birth weight and require insulin treatment throughout life. PNDM does not include autoimmune diabetes that almost never occurs before 6 months of age. Most patients have isolated diabetes, but approximately 20% will also have neurological findings. Patients have been described with developmental delay, epilepsy, and neonatal diabetes (DEND syndrome), relapsing and remitting diabetes, or intermediate phenotypes [1].
- **Hyperinsulinemia of infancy (HI):** HI, also known as familial hyperinsulinism and persistent hyperinsulinemic hypoglycemia of infancy, is inappropriate oversecretion of insulin despite hypoglycemia. It usually presents with seizures, hypotonia, poor feeding, apnea, and coma in the neonatal period or infancy, along with high birth weight. In the absence of treatment, HI can result in permanent brain damage [1,2].

Molecular and Biochemical Genetics:

- **Permanent neonatal diabetes:** Mutations of the *KCNJ11* [OMIM #600937] gene have been identified in patients with PNDM [3]. Studies have shown that approximately one-third to one-half of all cases of PNDM are due to activating mutations in *KCNJ11*. Genotype-phenotype correlations have been reported [1].
- **Hyperinsulinemia of infancy (HI):** Mutations of the *KCNJ11* gene have also been identified in patients with HI [4]. Approximately 5% of individuals with HI have inactivating mutations in *KCNJ11* [1].

KCNJ11 encodes Kir6.2, an essential subunit of the beta-cell K_{ATP} channel. It has one coding exon, and more than 32 mutations have been identified in patients with PNDM, HI, DEND and some instances of transient neonatal diabetes. A majority of mutations are missense mutations [1].

Management:

Patients with diabetes caused by mutations in the *KCNJ11* gene have K_{ATP} channels with decreased sensitivity to ATP. Thus, they remain open in the presence of glucose, decreasing the secretion of insulin. The most common mutation (R201H) associated with PNDM leads to a 40-fold reduction in sensitivity to ATP and failure of the channel to close. These patients have low levels of circulating insulin, and are therefore treated with insulin. Oral sulfonylureas are a class of drugs that close K_{ATP} channels independent of ATP, inducing secretion of insulin. Most patients with *KCNJ11* mutations can get better glycemic control without increasing hypoglycemic events by switching from insulin treatment to oral sulfonylureas [5,6].

Inheritance:

- **Permanent neonatal diabetes:** *KCNJ11* mutations that cause PNDM are gain-of-function mutations inherited in an autosomal dominant pattern. Most cases are sporadic and have no family history of this condition. Approximately 90% of autosomal dominant cases are spontaneous mutations [6]. Paternal germline mosaicism for the R201C mutations has been reported in a family with PNDM [1].
- **Hyperinsulinemia of infancy (HI):** *KCNJ11* mutations that cause HI are loss-of-function mutations inherited in an autosomal recessive pattern.

Additional Resources:

Dr. Louis Philipson is available for consultation prior to genetic testing. He is Director of the Comprehensive Diabetes Center at The University of Chicago. Patient's that do not have an identified *KCNJ11* mutation may be eligible for enrollment in Dr. Philipson's research project. Please contact him at lphilip@bsd.uchicago.edu.

Test methods:

We offer full sequencing of the coding exon and intron/exon boundaries.

Mutation analysis (sequencing)

Sample specifications:	3 to10 cc of blood in a purple top (EDTA) tube
Cost:	\$400
CPT codes:	83891, 83898, 83904, 83912
Turn-around time:	4 weeks

Testing for a known mutation in additional family members

Sample specifications:	3 to10 cc of blood in a purple top (EDTA) tube
Cost:	\$390
CPT codes:	83891, 83898 x 2, 83894, 83912
Turn-around time:	3-4 weeks

Prenatal testing for a known mutation

Sample specifications:	2 T25 flasks of cultured cells from amnio or CVS or 10ml of amniotic fluid
Cost:	\$590
CPT codes:	83891, 83898 x 2, 83894, 83912, 99051
Turn-around time:	1-2 weeks

Results

You will be informed of the results of your case as soon as it has been completed. Results, along with an interpretive report, will be faxed and mailed to the referring physician. Additional reports will be provided as requested. All abnormal results will be reported by telephone.

Laboratory Faculty and Staff:

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References:

1. Gloyn AL, Siddiqui J, Ellard S. Mutations in the genes encoding the pancreatic beta-cell K_{ATP} channel subunits Kir6.2 (*KCNJ11*) and SUR1 (*ABCC8*) in diabetes mellitus and hyperinsulinism (2006) *Human Mutation* 27(3): 220-231.
2. Glaser B. Familial hyperinsulinism (2005) *GeneReviews*. www.genetests.com
3. Gloyn AL, et al. Activating mutations in the gene encoding the ATP-sensitive potassium-channel subunit Kir6.2 and permanent neonatal diabetes (2004) *N Engl J Med* 350: 1838-49.
4. Thomas P, Ye Y, Lightner E. Mutation of the pancreatic islet inward rectifier Kir6.2 also leads to familial persistent hyperinsulinemic hypoglycemia of infancy (1996) *Human Molec Genet* 5(11): 1809-12.
5. Pearson ER, et al. Switching from insulin to oral sulfonylureas in patients with diabetes due to Kir6.2 mutations (2006) *N Engl J Med* 355: 467-77.
6. Hattersley A, et al. ISPAD clinical practice consensus guidelines 2006-2007: The diagnosis and management of monogenic diabetes in children (2006) *Pediatric Diabetes* 7:352-360.

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