Protein losing enteropathy-hepatic fibrosis syndrome in Saguenay-Lac St-Jean, Quebec is a congenital disorder of glycosylation type Ib


Congenital disorders of glycosylation (CDG) are newly described inborn errors of metabolism involving glycan moiety synthesis. CDG Ib is caused by a deficiency of cellular phosphomannose isomerase (EC 5.4.2.28) resulting from the presence of mutations in the corresponding gene, MPI, on chromosome 15 (mutation database: http://www.med.kuleuven.ac.be/cdg/). CDG Ib differs clinically from the other CDGs. It has no neurological symptoms, presenting mainly with hepatic and gastrointestinal symptoms and is thought to be underdiagnosed. Fewer than 20 CDG Ib patients have been reported. Importantly, CDG Ib is treatable by oral mannose supplementation, in contrast with other CDGs.

The syndrome of intractable diarrhoea of infancy was first defined in 1968 by Avery et al. In 1986, a syndrome of intractable diarrhoea associated with severe protein losing enteropathy, hypoglycaemia, and congenital hepatic fibrosis, was described in four infants from the Saguenay-Lac St-Jean (SLSJ) region of north eastern Quebec, Canada. All died before 21 months of age.

In retrospect, their symptoms resembled those of CDG Ib. In this article we describe the testing of parents of these children and report on a new French family with CDG Ib. The French-Canadian parents were all heterozygous for the same deleterious MPI mutation as found in the new French patient, proving that the SLSJ patients in fact suffered from CDG Ib.

PATIENTS AND METHODS

Patients

The clinical presentation of the SLSJ infants was previously described by Pelletier et al.

The new French patient with CDG Ib, whose family originates from Nantes (Brittany, France) and whose parents have no known consanguinity, presented with diarrhoea related to protein losing enteropathy with oedema, hypoalbuminaemia, hypoglycaemia, and recurrent hypoglycaemia. No liver cytolysis was noted. Hypocholesterolaemia was found and clotting tests showed thrombocytopenia. Coagulation factor XI was decreased, resulting in an increased prothrombin time. The patient had a typical CDG I profile on western blotting of serum glycoproteins. Blood leucocyte MPI activity was deficient (0.5 U/g total protein).

Methods

DNA samples were not available from the affected SLSJ children, of whom all known cases have died; no new cases have been identified since the initial publication. EDTA blood samples were obtained from three parents of SLSJ patients, both parents of one affected child, and the mother of a second patient; the father of the second patient, who was not sampled, was distinctly related to the mother (consanguinity 1/64). All originated from the SLSJ region. Control blood samples were simultaneously obtained from two unrelated healthy adults, to control for possible loss of activity during transport. Samples were also obtained from the French CDG Ib patient and his parents.

Cytosolic MPI activity was assayed in blood mononuclear cells, according to van Schaftingen and Jaeken.

DNA was isolated from fresh blood. The eight MPI exons were sequenced from genomic DNA after PCR with intronic primers designed from published DNA sequences (GenBank AF227216, AF227217, AF227218). Primer sequences are available on request. Sequencing was performed on an ABI PRISM (Perkin Elmer). The R295H (G884A) mutation creates a Ncol restriction site, which conveniently allowed confirmation of R295H by restriction digestion of the exon 7 amplicon and the French CDG Ib family. All were typed for the extragenic polymorphic markers D15S984, D15S1026, D15S114, and D15S818. Primer sequences were obtained from the Genome Database. The PCR products were analysed on an ABI PRISM sequencer (Perkin Elmer). Three intragenic single nucleotide polymorphisms, IVS3 +15A/G, IVS5 +9G/A, and A1131G (V377V), were screened by sequencing of exons 3, 5, and 8, respectively.

Key points

- Congenital disorder of glycosylation with hereditary phosphomannose isomerase (MPI) deficiency (CDG Ib) is a very rare disease, probably underdiagnosed. A fatal syndrome of protein losing enteropathy and congenital hepatic fibrosis described in 1986 in the Saguenay-Lac St-Jean (SLSJ) region of Quebec, Canada, clinically resembles CDG Ib, with intractable diarrhoea, hypoglycaemia, hepatomegaly, vomiting, and malnutrition.
- This prompted us to study MPI activity and the MPI encoding gene, MPI, in blood samples from three parents of two of the affected Canadian children. All the parents had partial deficiency of leucocyte MPI activity. MPI gene sequencing showed that all three were heterozygous for the same mutation in exon 7, R295H (G884A).
- Meanwhile, a newly diagnosed French CDG Ib patient was found to be homozygous for R295H and also for nearby polymorphic markers. Interestingly, an identical variant was also found for each marker on at least one chromosome of each of the SLSJ parents.
- This indicates that the SLSJ syndrome is a variant of CDG Ib. Since CDG Ib is treatable by oral mannose supplementation, awareness of this condition is imperative to allow early diagnosis and treatment.

LETTER TO JMG

In 1986, a syndrome of intractable diarrhoea of infancy was first defined in 1968 by Avery et al. In 1986, a syndrome of intractable diarrhoea associated with severe protein losing enteropathy, hypoglycaemia, and congenital hepatic fibrosis, was described in four infants from the Saguenay-Lac St-Jean (SLSJ) region of north eastern Quebec, Canada. All died before 21 months of age.

In retrospect, their symptoms resembled those of CDG Ib. In this article we describe the testing of parents of these children and report on a new French family with CDG Ib. The French-Canadian parents were all heterozygous for the same deleterious MPI mutation as found in the new French patient, proving that the SLSJ patients in fact suffered from CDG Ib.

PATIENTS AND METHODS

 Patients

The clinical presentation of the SLSJ infants was previously described by Pelletier et al.

The new French patient with CDG Ib, whose family originates from Nantes (Brittany, France) and whose parents have no known consanguinity, presented with diarrhoea related to protein losing enteropathy with oedema, hypoalbuminaemia, hypoglycaemia, and recurrent hypoglycaemia. No liver cytolysis was noted. Hypocholesterolaemia was found and clotting tests showed thrombocytopenia. Coagulation factor XI was decreased, resulting in an increased prothrombin time. The patient had a typical CDG I profile on western blotting of serum glycoproteins. Blood leucocyte MPI activity was deficient (0.5 U/g total protein).

Methods

DNA samples were not available from the affected SLSJ children, of whom all known cases have died; no new cases have been identified since the initial publication. EDTA blood samples were obtained from three parents of SLSJ patients, both parents of one affected child, and the mother of a second patient; the father of the second patient, who was not sampled, was distinctly related to the mother (consanguinity 1/64). All originated from the SLSJ region. Control blood samples were simultaneously obtained from two unrelated healthy adults, to control for possible loss of activity during transport. Samples were also obtained from the French CDG Ib patient and his parents.

Cytosolic MPI activity was assayed in blood mononuclear cells, according to van Schaftingen and Jaeken.

DNA was isolated from fresh blood. The eight MPI exons were sequenced from genomic DNA after PCR with intronic primers designed from published DNA sequences (GenBank AF227216, AF227217, AF227218). Primer sequences are available on request. Sequencing was performed on an ABI PRISM (Perkin Elmer). Three intragenic single nucleotide polymorphisms, IVS3 +15A/G, IVS5 +9G/A, and A1131G (V377V), were screened by sequencing of exons 3, 5, and 8, respectively.

Key points

- Congenital disorder of glycosylation with hereditary phosphomannose isomerase (MPI) deficiency (CDG Ib) is a very rare disease, probably underdiagnosed. A fatal syndrome of protein losing enteropathy and congenital hepatic fibrosis described in 1986 in the Saguenay-Lac St-Jean (SLSJ) region of Quebec, Canada, clinically resembles CDG Ib, with intractable diarrhoea, hypoglycaemia, hepatomegaly, vomiting, and malnutrition.
- This prompted us to study MPI activity and the MPI encoding gene, MPI, in blood samples from three parents of two of the affected Canadian children. All the parents had partial deficiency of leucocyte MPI activity. MPI gene sequencing showed that all three were heterozygous for the same mutation in exon 7, R295H (G884A).
- Meanwhile, a newly diagnosed French CDG Ib patient was found to be homozygous for R295H and also for nearby polymorphic markers. Interestingly, an identical variant was also found for each marker on at least one chromosome of each of the SLSJ parents.
- This indicates that the SLSJ syndrome is a variant of CDG Ib. Since CDG Ib is treatable by oral mannose supplementation, awareness of this condition is imperative to allow early diagnosis and treatment.
The parents share the same ancestral chromosome showing a R295H homozygote. The amplicon contains a constant (P1) are heterozygous R295H/N. The French CDG Ib patient is a R295H homozygote. The amplicon contains a constant NlaIII restriction site that serves as a control for completeness of digestion. ND=non-digested fragment.

RESULTS AND DISCUSSION

We report on a new French patient with CDG Ib and show that the SLSJ patients suffered from CDG Ib. To date, all reported CDG Ib patients have had similar clinical presentations with protein losing enteropathy, recurrent hypoglycaemia, and liver disease. CDG Ib patients have had similar clinical presentations with the SLSJ patients suffered from CDG Ib. To date, all reported CDG Ib patients have had similar clinical presentations with protein losing enteropathy, recurrent hypoglycaemia, and liver disease. CDG Ib patients have had similar clinical presentations with the SLSJ patients suffered from CDG Ib.

Blood leucocyte PMI activity of the three Canadian parents was reduced (2.7, 2.0, and 1.6 U/g total protein), compared to the Canadian (5.5 and 6.0 U/g total protein) and local (8.2 ± 3.3 U/g total protein) adult controls. Sequencing of the PMI encoding MPI cDNA of the Canadian parents showed that all were heterozygous for a single mutation in exon 7, R295H (G884A). This mutation had not been previously reported. During this period, we found that the French patient was homozygous for R295H and that his parents were heterozygous. No other sequence changes were detected in the coding regions of the samples tested, aside from a previously reported synonymous polymorphism at residue 377 (see below). The R295H mutation screened by restriction analysis was not found in 50 French controls and adult controls.

Sequencing of the PMI encoding MPI cDNA of the Canadian parents showed that all were heterozygous for a single mutation in exon 7, R295H (G884A). This mutation had not been previously reported. During this period, we found that the French patient was homozygous for R295H and that his parents were heterozygous. No other sequence changes were detected in the coding regions of the samples tested, aside from a previously reported synonymous polymorphism at residue 377 (see below). The R295H mutation screened by restriction analysis was not found in 50 French controls and adult controls. The arginine residue at position 295 is conserved in all species studied to date (S typhimurium, E coli, Candida albicans, Saccharomyces cerevisiae, A nidulans, and human MPI) consistent with the notion of a functional effect of the mutation.

Together, these data indicate that at least two of the SLSJ patients reported by Pelletier et al. suffered from CDG Ib. Given the clinical similarity of the patients, and their common geographical origin from the SLSJ region that is known to have a strong founder effect, shown, for instance, for hepatorenal tyrosinaemia and cystic fibrosis, it is likely that the great majority of CDG Ib patients from the SLSJ will be R295H homozygotes. Identification of the R295H mutation as the cause of the enteropathy-hepatic fibrosis syndrome in SLSJ children will allow for other studies to assess the frequency of this mutation in the SLSJ and surrounding regions. The lack of new cases after the report of the first cluster over 15 years ago raises questions about the frequency and distribution of the mutant allele and about disease expressivity. These questions can now be explored directly. They are particularly important because CDG Ib is treatable but can be fatal if untreated, and awareness of this disease is imperative to allow early diagnosis and therapy. CDG Ib should be considered in the differential diagnosis of patients with unexplained hypoglycaemia, chronic diarrhoea, liver disease, or coagulopathy, particularly in patients of SLSJ origin.

ACKNOWLEDGEMENTS

This work was supported by the Réseau de recherche sur les CDG (INSERM/AFM) 4MR29F.

---

**Table 1** Haplotypes of SLSJ parents (P1, P2, and P3) and the French CDG Ib parents (P4 and P5) and patient. Alleles in bold represent the alleles consistent with linkage disequilibrium with R295H. The five parents share at least one polymorphic variant at each tested site, the same as that found in the homozygous state in the French CDG Ib patient.

<table>
<thead>
<tr>
<th>Mutation</th>
<th>D155S18</th>
<th>D155S1026</th>
<th>IVS3 +15 A→G</th>
<th>IVS5 +9 G→A</th>
<th>A1131G (V377)</th>
<th>D155S114</th>
<th>D155S818</th>
</tr>
</thead>
</table>

---
REFERENCES


