A Point Mutation in the Bulge of the Iron-Responsive Element of the L Ferritin Gene in Two Families With the Hereditary Hyperferritinemia-Cataract Syndrome

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The molecular basis for the recently described hereditary hyperferritinemia-cataract syndrome is the presence of a mutation in the iron-responsive element (IRE) of the L ferritin gene, located on chromosome 19q13.3-13.4. Two mutations have been reported so far, altering adjacent nucleotides in the IRE loop, in a region that has been extensively studied in vitro and shown to mediate high affinity interaction with the iron-responsive protein. In this report, we describe two families with a new mutation in the bulge of the IRE stem, and we show that this mutation alters the protein-binding affinity of the IRE in vitro to the same extent as the loop mutation. In addition, we present evidence that some variability in the age of onset of cataract can be associated with this genetic syndrome, probably because of additional genetic or environmental factors that modulate the penetrance of the L ferritin defect in the lens. We confirm that the patients do not have increased iron stores despite the persistence of elevated serum ferritin levels and that, accordingly, they do not tolerate well venesection therapy. Further studies will be necessary to elucidate the mechanism responsible for the onset of cataract.

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MATERIALS AND METHODS

Case Report

Family 1. The proband (I-1) (Fig 1), a 77-year-old woman, was hospitalized for a sudden episode of generalized edema accompanied by reduced serum albumin levels. In her past clinical history a bilateral cataract was reported. Hemoglobin, leukocyte, and platelet counts were in the normal range, serum iron was 12.8 µmol/L, transferrin 3.14 g/L, and ferritin 1,600 µg/L. Sedimentation rate was elevated, whereas the other acute phase reactants were within normal range. Because a kidney and hepatic origin of the edema was ruled out, as well as malnutrition and gastrointestinal protein loss, the patient was subjected to a liver biopsy, which showed a normal liver structure and a scant amount of iron. A monoclonal IgG gammapathy was present, but the bone marrow biopsy was normal with no evidence of iron overload. After health recovery, the patient was discharged from the hospital with normal blood tests, with the exception of serum ferritin, which was persistently elevated. In an attempt to reduce the hyperferritinemia, the patient was started on a program of phlebotomies, but after only three phlebotomies of 350 mL each, she developed anemia (hemoglobin [Hb] decreased from 11.7 to 8.4), transferrin saturation decreased from 20%
...to 5%, but serum ferritin did not change. A family study was performed, and two sons and a grandchild were tested.

The elder son (II1), 52 years old, did not have any pathological report but did have a bilateral cataract that had been operated on 10 years before. All his tests were normal (serum iron 21.4 µmol/L and transferrin 2.1 g/L), but his ferritin was 1,200 µg/L. He was phlebotomized, and after six phlebotomies Hb turned from 15.6 to 13, transferrin saturation decreased, and serum ferritin did not vary. Later, the patient moved to another city and he was again subjected to several phlebotomies for the high values of ferritin and consequently developed a severe anemia without any modification of ferritin values. He was also subjected to liver biopsy, which showed a completely normal liver without evidence of iron or inflammatory cells. However, some granular, yellowish pigment of unknown composition was also observed. His 16-year-old daughter was examined; blood cell counts and iron indices were all in the normal range. The younger son (II2), 45 years old, had increased serum ferritin (2,264 µg/L) and a normal transferrin saturation. He was subjected to 14 phlebotomies, once a week, with a decrease of Hb from 15 to 12.5 g/dL, a decrease of transferrin saturation, but no modification of ferritin levels. He had no history of cataract, and a recent control ruled out the presence of cataract.

Family 2. The proband (III1) (Fig 1), a 24-year-old woman whose sole previous medical problem was a bilateral cataract, presented in 1995 with constant hyperferritinemia. This biochemical abnormality was detected in the frame of a family study initiated from her sister who had been suspected from having genetic hemochromatosis. She was otherwise clinically asymptomatic. Serum ferritin levels ranged between 1,250 and 1,400 µg/L (upper normal limit 250). Serum iron was normal (16 to 19 µmol/L), and transferrin saturation (19% to 28%) slightly decreased. Blood cell counts, sedimentation rate, serum hepatic enzyme activities (aspartate transaminase [AST]; alanine transaminase [ALT]; γ-glutamyl transferase [GGT]) were normal. Liver biopsy showed no abnormalities, and particularly Perl’s staining was negative.

Her family members presented as follows: (1) her mother (II1) had a bilateral congenital cataract and hyperferritinemia; (2) her maternal grandfather had a bilateral cataract; (3) her brother (III2) had a bilateral cataract, no serum ferritin could be checked; and (4) her sister (III3), born in 1953, had been explored in 1993 to 1994 for chronic hyperferritinemia discovered after a malaise in 1992. Serum ferritin levels were comprised between 1,025 and 1,330 µg/L, with erythrocyte sedimentation rate, serum hepatic enzyme activities, and particularly Perl’s staining was negative. The younger son (II1) (Fig 1) had a bilateral congenital cataract and hyperferritinemia; (3) her brother (III2) had a bilateral cataract, no serum ferritin could be checked; and (4) her sister (III3), born in 1953, had been explored in 1993 to 1994 for chronic hyperferritinemia discovered after a malaise in 1992. Serum ferritin levels were comprised between 1,025 and 1,330 µg/L, with erythrocyte sedimentation rate, serum hepatic enzyme activities, and particularly Perl’s staining was negative.

Results

Genomic DNA Analysis

Genomic DNA was extracted from blood samples collected on citrate-EDTA and subjected to polymerase chain reaction (PCR) amplification. A first round of amplification was performed using Lprom and Lex2 (see below) as 5’ and 3’ primers, and 30 cycles with 95°C for 30 seconds, 88°C for 30 seconds, and 72°C for 1 minute. The resulting 610-bp fragment was then purified on agarose gel and further amplified using seminested primers (Lex1 and Lex2), mapping to exon 1 and 2. The resulting 570-bp fragment was purified and sequenced.

Primers used for PCR amplification were (see Fig 2A): Lprom: 5’ CGGCGCACCCATAAAGAAGGCC (upper primer); Lex2: 5’ GCTG-GTTTTGCA TCTTCAG (lower primer); and Lex1: 5’ AGTTCGGCG-GTCCCGCGGGTC (upper primer).

Gel Retardation Assay

Labeled RNAs were transcribed in vitro from oligonucleotide template using T7 RNA polymerase, as previously described. The oligonucleotide used to synthesize the normal L ferritin IRE probe was: 5’ GTGTTCCGTCCAAAACACTGTTGAGACGACAAGAGACATCG-GCTATAGTGATCTGATATTAGC (upper primer).

RESULTS

We performed PCR amplification of genomic DNA from affected members of both families. Two rounds of PCR with a different primer pair were necessary to avoid the amplification of the ferritin L subunit intronless pseudogenes. A first round was performed using a sense primer mapping to the proximal promoter region of the L ferritin gene and an antisense primer mapping to exon 2. The resulting 610-bp fragment was gel-purified, and a nested PCR was performed using a combination of oligonucleotides mapping to the beginning of exon 1 and to exon 2. Sequencing of the PCR fragment revealed that affected members were heterozygous for a point mutation, consisting in a G to T transition in the bulge of the IRE (Fig 2B and C).

To confirm that the mutation was responsible for a reduced binding affinity of the IRE to the IRP, we performed gel retardation assay. Normal and mutated sense RNA probes were generated by in vitro transcription in the presence of 32P-UTP.
In the presence of K562 cytoplasmic extracts, the normal probe generated a high-affinity protein RNA complex (Fig 3, lane 2). Mutated probes corresponding to the previously described loop mutation (mut1, lane 4) or to the new bulge mutation (mut2, lane 6) did not generate any specific complex. In addition, the IRE/IRP complex generated by the normal IRE probe was partially displaced by a 50-fold molar excess of the cold probe (lane 8) and almost entirely by a 100-fold excess (lane 9), whereas similar excess of the cold probe bearing the loop mutation (mut1, lanes 10 to 12) or the bulge mutation (mut2, lanes 13 to 15) did not compete for binding to the normal IRE probe.

The results confirm that the G to T mutation in the bulge of the L ferritin IRE fully impairs the protein-binding affinity of the IRE and is likely to be responsible for the increased serum ferritin levels in patients of the two families and probably for the associated cataract.

DISCUSSION

In this report, we describe a new mutation in the IRE of the L ferritin mRNA in two families with a hereditary hyperferritinemia-cataract syndrome, one from Italy and one from France. The first two mutations previously reported affected adjacent nucleotides from the IRE loop, whereas the two families
described here have the same G to T change in the three unpaired nucleotide bulge of the IRE. This finding suggests that this syndrome is likely to be more frequent than was expected at first and reinforces the idea that the presence of a mutated IRE in the L ferritin gene, located on chromosome 19q13.3-13.4, is responsible for the association of cataract and hyperferritinemia. In addition, the position of the mutation is the first in vivo evidence for the key role of the bulge in the binding affinity of the IRE, although it has been shown that the tertiary structure of the stem-bulge region of the IRE is a critical determinant of translational regulation by iron.\(^\text{17}\) The structure of the stem varies between the different IREs\(^\text{6}\) with a three-base UGC bulge in both H and L ferritin IRE, whereas the stem of the five IREs in the transferrin receptor mRNA and of the single IRE in the erythroid ALA-S mRNA have a single cytosine bulge. An extensive in vitro study performed by Henderson et al has shown the importance for the IRE function of this highly conserved cytosine present in the bulge of all known IREs.\(^\text{18}\)

Our results present evidence that the G from the UGC bulge is important for the IRE function in vivo and for the high-affinity RNA protein interaction in vitro. This mutation abrogates the base pairing between G32 and C50 (Fig 2C), which might be necessary for the proper conformation of the IRE, as shown by studies based on nuclear magnetic resonance (NMR) spectroscopy\(^\text{19}\) or on identification of ligands with the highest affinity for the IRP.\(^\text{20}\) Alternately, the mutation might impair a specific point of protein-RNA interaction, because studies using UV radiation-induced cross-links have shown that nucleotides 31 to 43, including the UGC bulge and most of the loop, interact with the IRP and are likely to be buried within the active site of the native, iron-free protein.\(^\text{21}\)

In the two families reported here, seven individuals had a known history of cataract, and whenever serum ferritin levels were assayed, they were found to be elevated (six patients), ranging from 1,200 to 2,200 µg/L. These values are very similar to those reported in the family with an A to G mutation in the loop,\(^\text{3}\) suggesting that mutations in both the loop and the bulge affect the in vivo function of the IRE to the same extent. This is in agreement with our in vitro studies, which show that both mutated IREs have lost IRP-binding activity and have no ability to compete for IRP binding to the wild-type sequence. However, we observed a marked phenotypic variability in the age of onset of cataract between the two families reported here, despite the presence of the same mutation. In fact, whereas in Family 2 affected members developed cataract during the late childhood, the patients from Family 1 showed signs of cataract around the age of 40; surprisingly, patient H, age 45 years, still does not show any evidence of lens opacification, although he carries the mutation and his level of serum ferritin is even higher than the other members of the family (2,260 to 1,200 to 1,600). The mechanism leading to the onset of cataract is still unknown, but it is possible to speculate that abnormal deposits of ferritin molecules in the successive layers of lens epithelial cells lead progressively to lens opacification. In the case of patient H, it is possible that additional factors, either genetic or environmental, modulate the penetrance of the L ferritin defect in the lens. Further follow-up will show whether he develops cataract or not.

Altogether, five families with the hyperferritinemia-cataract syndrome described here have now been described, and it becomes clear that the elevated serum ferritin levels do not reflect the presence of increased iron stores. Whenever a liver biopsy or a bone marrow biopsy (this report, Family 1, patient I\(_1\)) was performed, there was no increased staining for iron. Accordingly, these patients did not tolerate well venesection therapy. It is also important to emphasize that recently a new syndrome has been described where patients who most often present metabolic disorders, such as increased body mass index, hyperlipidemia, glucose intolerance, and hypertension, have hyperferritinemia and normal transferrin saturation.\(^\text{22}\) However, contrary to the hereditary hyperferritinemia-cataract syndrome, these patients have iron overload and need phlebotomies. It becomes mandatory to be aware of the existence of these two syndromes because it appears from the few cases described so far that patients with the hyperferritinemia-cataract syndrome may rapidly develop severe anemia when they are phlebotomized, even if serum ferritin does not decrease. On the contrary, the phlebotomies will lead to a progressive reduction in the serum ferritin level in patients with iron overload. Therefore, a strict follow-up of these patients should be performed, including regular hemoglobin determination together with serum ferritin assay.

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REFERENCES

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