A point mutation in the human serum albumin gene results in familial dysalbuminaemic hyperthyroxinaemia

Charles E Petersen, Alfred G Scottolini, Linda R Cody, Morton Mandel, Neil Reimer, Nadhipuram V Bhagavan

Abstract
Using DNA samples obtained from two unrelated patients, diagnosed as having familial dysalbuminaemic hyperthyroxinaemia (FDH), exons 1–14 which span the entire coding region of the human serum albumin (HSA) gene were amplified by the polymerase chain reaction. The sequence of each of the 14 DNA fragments was then determined. In each case a point mutation was identified at nucleotide 653 which causes an Arg to His substitution at amino acid position 218. The substitution was confirmed by amino acid sequencing of a mutant peptide resulting from tryptic digestion of the protein. Abnormal affinity of FDH HSA for a thyroxine (T4) analogue was verified by an adaptation of the procedure used in routine free T4 measurement. The location of the mutation is discussed in relation to other studies on the binding properties of HSA.

(F J Med Genet 1994;31:355–359)

Familial dysalbuminaemic hyperthyroxinaemia (FDH) is an autosomal dominant syndrome in which, although patients have a raised total serum thyroxine concentration (TT4), they are clinically euthyroid. The euthyroid state of FDH patients is confirmed by normal serum levels of free thyroxine (FT4), thyroid stimulating hormone, and triiodothyronine. The prevalence of FDH has been estimated in our earlier study to be about 4% of hyperthyroxinaemic patients. The two FDH patients discussed in this paper are new case findings, unrelated to our previous study. The major hazard for FDH patients is that they can be misdiagnosed as hyperthyroid and subjected to unwarranted treatment. The raised serum TT4 in FDH patients results from the presence of an abnormal human serum albumin (HSA) which exhibits enhanced binding to thyroxine (T4). Our study on intrinsic fluorescence properties of purified normal HSA and HSA obtained from an FDH patient, hereafter referred to as patient 1, using both steady state and time resolved methodologies, showed that the patient had two HSA species with different affinities for T4. Until now the specific molecular defect of FDH HSA was unknown. This study describes a specific amino acid substitution resulting from the same nucleotide change in the HSA gene in two unrelated FDH patients.

We have developed a rapid method for identifying FDH patients with the mutation we describe by using polymerase chain reaction (PCR) amplification of exon 7 of the HSA gene followed by restriction endonuclease digestion and gel electrophoresis.

Materials and methods
SOURCE OF WHOLE BLOOD
Whole blood drawn up in EDTA was obtained from patient 1 and patient 2, diagnosed as FDH by methods described previously. Whole blood was also obtained from a normal volunteer.

PURIFICATION OF NORMAL AND FDH HSA
HSA was purified from the plasma of patient 1 and the normal volunteer using identical methods. Ammonium sulphate was added to the plasma to 50% saturation. HSA was precipitated from the supernatant solution by lowering the pH to 4.4 at 2°C. The precipitate was dissolved in water and the lipid removed by reaction with Norit A activated charcoal at pH 3-0 followed by dialysis in water. The HSA was further purified by affinity chromatography using cibacron blue linked to sepharose CL-6B (Sigma). The matrix was equilibrated in 0-04 mol/l sodium phosphate, pH 7.5, 0-15 mol/l NaCl (PBS). After washing the column with PBS, the bound HSA was eluted with 3 mol/l NaCl. The eluent was dialysed in water. The purity of the HSA was determined by SDS-PAGE.

TRYPTIC DIGESTION OF HSA
Purified HSA from patient 1 and purified commercial HSA (Sigma, fatty acid free, globulin free) dissolved in water were lyophylised and treated identically.

The purified HSA was reduced and alkylated to pyridyl-ethyl cystinyl HSA (PEC-HSA) and desalted by preparative C-8 reverse phase high pressure liquid chromatography (HPLC) (0-1% trifluoroacetic acid to acetonitrile). The desalted PEC-HSA was digested with trypsin (2%) in 100 mmol/l sodium bicarbonate, 2 mol/l urea, 1 mmol/l calcium chloride, pH 8-0 for 24 hours. The resultant peptide mixture was subjected to C-18 reverse phase HPLC using a linear gradient (0-1% trifluoroacetic acid, 9% acetonitrile to
DIGESTION OF EXON 7 CONTAINING FRAGMENT

DNA fragments containing exon 7 were generated by the same procedure used for sequencing. The fragment containing exon 7 from a normal HSA gene does not contain an HphI site whereas a mutation identified in both FDH patients creates an HphI site. The amplification product was precipitated in 75% absolute ethanol and resuspended in 20 μl of HphI digestion buffer. Ten units of HphI (New England Biolabs) was added and the reaction was incubated for six hours at 37°C. For each sample a control was incubated without HphI. Samples were electrophoresed in a 3% agarose (FMC BioProducts, Metaphor Agarose) gel in 1 x TAE (0.04 mol/l Tris-acetate, 0.001 mol/l EDTA, pH 8.0) and bands were visualised by staining with ethidium bromide.

ASSESSMENT OF HSA BINDING IN THE ABSENCE OF OTHER SERUM COMPONENTS

The binding properties of HSA from the normal volunteer and patient 1 were compared by using a radioimmunoassay kit (Clinical Assays, Stillwater, MN) designed to measure the free T4 level in serum samples. Commercial HSA was also assayed as a control. The assay is designed to measure free T4 by competitive binding. T4 antibody is immobilised on the lower inner surface of a tube. A serum sample is added along with a fixed amount of an [125I]T4 tracer. The tracer and the free T4 in the serum sample compete for binding sites on the antibody giving an inverse correlation between free T4 concentration and radioactivity associated with antibody.

The assay was adapted to measure HSA affinity for the T4 tracer as follows. To each assay tube 1 ml of tracer buffer and the T4 tracer were added followed by 250 μg of HSA in 50 μl H2O. The tubes were incubated for 90 minutes at 37°C and the fluid was aspirated. The amount of radioactivity remaining bound to the assay tubes was measured in an Iso-Data gamma counter.

Results

AMPLIFICATION AND SEQUENCING OF PCR FRAGMENTS

DNA fragments containing exons 1–14, which span the entire coding region of the HSA gene, were successfully amplified and sequenced using genomic DNA obtained from the normal volunteer and patient 1. The sequence obtained for the normal volunteer matched a previously published HSA sequence at all coding nucleotides. Sequence obtained from patient 1 was identical to the sequence at all positions except for a G to A transition at nucleic acid position 653 (fig 1) causing an Arg/GGC to His/CAC substitution at amino acid position 218 in mature HSA.

Although an adenosine appears at position 653 in the FDH sequence the guanosine is still present but at half the intensity of the corresponding guanosine in the normal sample. This indicates that the condition is heterozygous as the PCR fragment represents an amplification of both alleles. The mutation was verified by observing a C to T transition on the opposite strand (fig 1). Amplification and sequencing of exon 7 using genomic DNA obtained from patient 2 gave a result for exon 7 identical to that of patient 1.

HphI DIGESTION OF EXON 7 CONTAINING FRAGMENT

The exon 7 containing DNA fragments from both FDH patients were subjected to HphI digestion. Only partial digestion was observed for both heterozygotes, as expected (fig 2). A
A point mutation in the human serum albumin gene results in familial dysalbuminaemic hyperthyroxinaemia

Figure 1 Alignment of normal and (FDH) sequencing termination reactions for exon 7 showing location of mutation. G, C, T, and A refer to the termination reactions containing ddGTP, ddCTP, ddTTP, and ddATP respectively. Lanes labelled (n) and (v) indicate sequencing termination reactions for normal and variant (FDH) samples respectively. The position of the mutation is indicated by an arrow.

Figure 2 HphI digestion of exon 7 containing DNA fragments from the normal volunteer and FDH patients 1 and 2. Fragments from the normal person and FDH patients 1 and 2 are labelled as normal, variant 1, and variant 2, respectively. The addition of 10 U HphI is indicated as +, – indicates that no HphI was added.

Figure 3 Comparison of the chromatograms derived from a tryptic digest of commercial HSA and FDH HSA clearly showed an additional peak in the FDH chromatogram (fig 3, panels A and B). Sequencing of the material from the extra peak gave a homogeneous peptide with a sequence of AWAVAHLSQR, indicating an Arg substitution at position 218 (fig 4). The amino acid percent composition of the peptide confirmed the above sequence.

Figure 4 HphI digestion of exon 7 containing DNA fragments from the normal volunteer and FDH patients 1 and 2. Fragments from the normal person and FDH patients 1 and 2 are labelled as normal, variant 1, and variant 2, respectively. The addition of 10 U HphI is indicated as +, – indicates that no HphI was added.

single thick band is seen at a position corresponding to a 174 base pair marker. The size of the two fragments expected after digestion are 173 and 182 base pairs which migrate in the agarose gel as an unresolved doublet. The fragment derived from the normal volunteer is not digested by HphI.

TRYPIC DIGESTION OF HSA
Comparison of the chromatograms derived from a tryptic digest of commercial HSA and FDH HSA clearly showed an additional peak in the FDH chromatogram (fig 3, panels A and B). Sequencing of the material from the extra peak gave a homogeneous peptide with a sequence of AWAVAHLSQR, indicating an Arg substitution at position 218 (fig 4). The amino acid percent composition of the peptide confirmed the above sequence.

ASSESSMENT OF HSA BINDING IN THE ABSENCE OF OTHER SERUM COMPONENTS
Adding purified HSA from the normal volunteer or commercial HSA to the T4 assay system did not significantly reduce the amount of T4 tracer associated with the T4 antibodies, relative to a standard to which no HSA was added. Adding purified HSA from patient 1 significantly reduced the amount of tracer associated with the antibodies indicating an increased affinity of the FDH HSA for the T4 tracer (table).

Discussion
The recently published high resolution x ray crystallographic structure for HSA (2.8A) gives an atomic structure divided into three homologous domains 1, 2, and 3, each divided
3B. Binding studies with some of these variants with a variety of ligands have generally shown relatively minor effects on binding affinity.23

The only Trp residue in HSA is at position 214. The position of the substitution identified is located only four amino acids downstream from Trp 214. Previous fluorescence lifetime studies based on the ability to excite Trp 214 selectively using FDH HSA obtained from patient 1 have shown enhanced quenching of tryptophan fluorescence relative to Trp 214 of normal HSA.9 This result is consistent with a substitution located relatively close to Trp 214.

Binding studies using HSA immobilised on sepharose have shown that thyroxine binding can be divided into a high and low affinity component. Only the high affinity component can be inhibited competitively with bilirubin.24 Bovine serum albumin has been shown to have a single high affinity bilirubin binding component which is retained by a proteolytic bovine serum albumin fragment (residues 186–306) which corresponds to the 2A subdomain in HSA.9 The observation that the high affinity thyroxine binding component of HSA can be competitively inhibited by bilirubin which has been localised to subdomain 2A suggests that the high affinity thyroxine binding site is located in subdomain 2A. The amino acid substitution we have described is also located in subdomain 2A.

Both FDH patients in this study are heterozygous. The plasma of such persons contains a mixture of variant and normal HSA. One of our immediate goals is to obtain accurate binding constants for the variant HSA with a number of ligands. In previous binding studies a mixture of HSA species obtained from a heterozygous FDH patient was used and although the binding data were sufficient to determine heterozygosity an accurate binding constant for thyroxine could not be determined owing to the two component nature of the system.9 Attempts to separate a mixture of HSA from a heterozygote using a variety of electrophoretic and chromatographic techniques have been unsuccessful. The observation that the mixture can not be resolved by a variety of separation methods that take advantage of differences in the shape and surface structure of proteins suggests that the variation acts locally to affect thyroxine binding rather than by transmitting a change to another location in the molecule through an effect on overall structure.

<table>
<thead>
<tr>
<th>Sample</th>
<th>CPM*</th>
</tr>
</thead>
<tbody>
<tr>
<td>No HSA</td>
<td>243551</td>
</tr>
<tr>
<td>Commercial HSA</td>
<td>24232</td>
</tr>
<tr>
<td>Normal HSA</td>
<td>26649</td>
</tr>
<tr>
<td>FDH HSA</td>
<td>12426</td>
</tr>
</tbody>
</table>

* Indicates the counts per minute remaining associated with the T4 antibodies after incubation with the HSA sample indicated.
† All assays were carried out in duplicate.
A point mutation in the human serum albumin gene results in familial dysalbuminaemic hyperthyroxinaemia

We plan to express the variant HSA in a eukaryotic expression system so that we can determine an accurate binding constant for thyroxine and other relevant ligands using a homogeneous preparation of HSA. Using site-directed mutagenesis we hope to create variants in the 2A subdomain with a range of binding properties for thyroxine, bilirubin, and warfarin. Warfarin binding has also been localised to domain 2.26

Having observed the same mutation in two unrelated FDH patients it is likely to be the most common cause of FDH. (During the treatment of hypothyroidism, a disease state in which free serum thyroxine is raised. HSA or a truncated form with increased affinity for thyroxine could be given intravenously. The thyroxine would be sequestered by the HSA, rapidly lowering the free thyroxine level. We are currently working towards the development of this clinical application.

We are grateful to Stephen Swenson for help with protein purification, Theresa Oshiro for technical assistance, and Steven Seifried and Gordon Edlin for advice and helpful suggestions.