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Mapping the Gene Causing X-linked Recessive Idiopathic Hypoparathyroidism to Xq26–Xq27 by Linkage Studies

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Abstract

Idiopathic hypoparathyroidism has been reported to occur as an X-linked recessive disorder in two multigeneration kindreds. Affected individuals, who are males, suffer from infantile onset of epilepsy and hypocalcemia, which appears to be due to an isolated congenital defect of parathyroid gland development; females are not affected and are normocalcemic. We have performed linkage studies in these two kindreds (5 affected males, 11 obligate carrier females, and 44 unaffected members) and have used cloned human X chromosome sequences identifying restriction fragment length polymorphisms to localize the mutant gene causing this disorder. Our studies established linkage between the X-linked recessive idiopathic hypoparathyroidism gene (HPT) and the DXS98 (4D.8) locus, peak LOD score = 3.82 (θ = 0.05), thereby mapping HPT to the distal long arm of the X chromosome (Xq26–Xq27). Multilocus analysis indicated that HPT is proximal to the DXS98 (4D.8) locus but distal to the F9 (Factor IX) locus, thereby revealing bridging markers for the disease. The results of this study will improve genetic counseling of affected families, and further characterization of this gene locus will open the way for elucidating the factors controlling the development and activity of the parathyroid glands. (J. Clin. Invest. 1990. 86:40–45.)

Key words: DNA linkage studies • X-linked hypoparathyroidism

Introduction

Hypoparathyroidism is an endocrine disorder in which hypocalcemia and hyperphosphatemia are the results of a deficiency in parathyroid hormone (PTH) secretion. There are a variety of causes of hypoparathyroidism and the disorder may occur after trauma to the parathyroids during neck surgery, as part of a pluriglandular autoimmune disorder, or as a congenital defect, for example in the DiGeorge syndrome. In addition, hypoparathyroidism may develop as a solitary endocrinopathy, and this form has been called isolated or idiopathic hypoparathyroidism. Familial occurrences of idiopathic hypoparathyroidism have been reported and autosomal dominant (1), autosomal recessive (2), and X-linked recessive (3, 4) inheritances have been established (5). The investigation of the genetic defects in these hereditary disorders of isolated PTH deficiency has been facilitated by the recent advances in molecular biology. Human PTH, which is an 84 amino acid polypeptide (6), is encoded by a single gene located on the short arm of chromosome 11 (7). It has been postulated that mutations either within or near the PTH gene, or at other loci, which affect the embryological development, cell structure, or regulation of the parathyroids, may cause this reduction in parathyroid gland activity (8). An abnormality within the PTH gene itself has been identified in a patient with autosomal dominant idiopathic hypoparathyroidism (9). However, in other patients with inherited forms of idiopathic hypoparathyroidism, genetic abnormalities within the PTH gene have not been detected (8, 10) and mutations at other loci need to be sought. We have undertaken family linkage studies using cloned human X chromosome sequences identifying restriction fragment length polymorphisms (RFLPs), and have mapped the gene causing X-linked recessive idiopathic hypoparathyroidism (HPT). This localization of the mutation represents a major step toward defining the primary defect causing this X-linked recessive disorder of calcium homeostasis and elucidating the genetic factors involved in the embryological development of the parathyroid glands.

Methods

Families. Two kindreds from the state of Missouri, in whom idiopathic hypoparathyroidism had been inherited in an X-linked recessive manner in five or more generations (3, 4), were ascertained for linkage studies. These two kindreds appeared to be unrelated, as a common ancestor was not identified after extensive genealogy dating to the mid-1800s. The family reported by Peden in 1960 (3) was designated family P/60, and that reported by Whyte and Weldon in 1981 (4) was designated family W/81. The disorder occurred in males only, who suffered in the neonatal or early infantile periods from hypocalcemic seizures that were invariably fatal if the hypocalcemia remained uncorrected. Further investigations of these male patients revealed undetectable circulating immunoreactive PTH concentrations and a normal renal response to bovine PTH extract. In addition, autopsy of an affected teenage boy from family P/60 indicated that the deficiency in PTH was the result of parathyroid gland agenesis (1). Clinical hypocalcemia was not observed in any of the affected individuals. Unaffected relatives had no history of epilepsy and were normocalcemic. Females were classified as carriers of X-linked recessive idiopathic hypoparathyroidism only if males in subsequent generations manifested the disease. Venous blood was obtained, after informed consent, from 60 family members; 5 were affected males, 11 were carrier females, and 44 were unaffected (26 males, 18 females). Of the 60 members, 50 (4 affected, 7 carriers, and 39 unaffected) were from four generations of kindred W/81 and 10 (1 affected, 4 carriers, and 5 unaffected) were from three generations of kindred P/60.

Methods. Venous blood samples were collected in tubes containing EDTA and kept frozen at −70°C. Leukocyte DNA was prepared by

1. Abbreviations used in this paper: HPT, X-linked recessive idiopathic hypoparathyroidism; LOD score, log10 odds ratio favoring linkage; RFLP, restriction fragment length polymorphism.
Figure 1. Map of some clinically useful DNA probes on the human X chromosome, which is schematically represented with Giemsa bands. The short arm is designated 'p' and the long arm is designated 'q'. The DNA probes are cloned human X chromosome sequences and are shown juxtaposed to their region of origin. The region from which each DNA probe is derived is ascertained by situ hybridization, by the use of somatic cell hybrids, or by linkage studies. Each DNA probe of unknown function is assigned a locus number; for example, the DNA probe 4D.8 is assigned DXS98. This indicates that DXS98 is a DNA sequence (D) from the X chromosome (X) detecting a single (S) DNA segment in the haploid genome, and is designated the number 98 by the Human Gene Mapping Committee (16). DNA probes of known function are allocated gene symbols; for example, the genes encoding ornithine carbamoyltransferase and coagulation Factor IX are assigned OTC and F9, respectively. These DNA probes, which reveal RFLPs, were used as genetic markers in linkage studies of families affected with X-linked recessive idiopathic hypoparathyroidism. Linkage between HPT and the DXS98 (4D.8) locus was established, thereby mapping HPT to the distal region of the long arm of the X chromosome (Xq26–Xq27).

Table I. LOD Scores for Linkage of X-linked Markers and HPT

<table>
<thead>
<tr>
<th>Locus</th>
<th>Probe</th>
<th>Peak</th>
<th>LOD scores Z (θ)</th>
</tr>
</thead>
<tbody>
<tr>
<td>DXS85</td>
<td>782</td>
<td>0.500 0.00</td>
<td>-4.99 -1.60 -1.01 -0.68 -0.46 -0.19 -0.05</td>
</tr>
<tr>
<td>DXS41</td>
<td>99.6</td>
<td>0.154 1.46</td>
<td>-2.13 1.02 1.38 1.46 1.41 1.13 0.66</td>
</tr>
<tr>
<td>DXS84</td>
<td>754</td>
<td>0.280 0.31</td>
<td>-8.08 -1.55 -0.49 -0.03 0.20 0.30 0.16</td>
</tr>
<tr>
<td>DXS7</td>
<td>L1.28</td>
<td>0.091 1.82</td>
<td>0.26 1.75 1.82 1.74 1.61 1.21 0.67</td>
</tr>
<tr>
<td>DXYS1</td>
<td>pDp34</td>
<td>0.406 0.03</td>
<td>-4.39 -1.09 -0.58 -0.33 -0.17 -0.02 0.03</td>
</tr>
<tr>
<td>DXS37</td>
<td>30Rb</td>
<td>0.226 0.88</td>
<td>-7.04 -0.53 0.36 0.73 0.86 0.79 0.49</td>
</tr>
<tr>
<td>DXS51</td>
<td>52A</td>
<td>0.000 0.73</td>
<td>0.73 0.68 0.62 0.56 0.50 0.35 0.19</td>
</tr>
<tr>
<td>F9</td>
<td>Factor IX</td>
<td>0.096 1.62</td>
<td>0.05 1.54 1.62 1.56 1.44 1.08 0.60</td>
</tr>
<tr>
<td>DXS98</td>
<td>4D.8</td>
<td>0.050 3.82</td>
<td>2.52 3.82 3.68 3.38 3.00 2.05 0.92</td>
</tr>
<tr>
<td>DXS52</td>
<td>St14</td>
<td>0.156 2.33</td>
<td>-4.81 1.49 2.17 2.33 2.26 1.76 0.95</td>
</tr>
<tr>
<td>DXS15</td>
<td>DX13</td>
<td>0.128 1.81</td>
<td>-1.63 1.49 1.78 1.80 1.69 1.26 0.63</td>
</tr>
</tbody>
</table>
markers was expressed as a location score, which was twice the natural logarithm of the odds ratio. The probability of the most likely gene order and the relative likelihoods of other gene orders were then ascertained from this location score curve. If, for a location score curve with marker framework A-B-C, the peak location scores x and y are obtained for the intervals A-B and B-C, respectively, then the relative likelihood that the disease maps in the interval A-B as opposed to B-C is given by \( \frac{e^y}{e^x} \).

**Results**

Members of family W/81 proved informative for 11 X-linked genetic markers, 4 from the short arm and 7 from the long arm, and the results of two-point linkage analysis are shown in Table I. The members from family P/60 proved uninformative and are therefore not included in the results shown in Table I. Linkage between HPT and the DXS98 (4D.8) locus was established with a peak LOD score of 3.82, a recombination fraction (\( \theta \)) of 0.05, and a 95% confidence interval (0.001, 0.22), thereby localizing HPT to the distal long arm of the X chromosome (Xq26-Xq27). All the other X-linked RFLP loci gave negative or low LOD scores.

An analysis of recombination events within this distal segment of the long arm of the X chromosome helped to further localize the HPT locus. The pedigree in Fig. 2 shows 40 members (29 surviving and 11 deceased) in five generations from family W/81 with genetic marker data. The pedigree is informative for five X-linked RFLP loci, whose order in the region Xq25-Xq28 has been established as Xcen-DXS37-F9-DXS98-DXS52-DXS15-Xqter, and multipoint crosses exist. Individual IV.4 is a carrier mother who is heterozygous for F9, DXS98, DXS52, and DXS15, and the alleles which she has inherited from her mother (III.2) and father (III.1) can be ascertained by examination of her mother’s (III.2) and unaffected brother’s (IV.1) genotypes. Her affected son, V.2, shows

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**Figure 2.** Pedigree from family W/81, segregating for X-linked recessive idiopathic hypoparathyroidism and distal long arm RFLP loci, whose respective alleles are indicated in parenthesis: DXS37 (Ee), F9 (NN), DXS98 (Aa), DXS52 (2, 3, 6, 8), and DXS15 (Tt). The loci are shown in the correct order but not the correct distances apart. Individuals are represented as: unaffected male (c), affected male (e), unaffected female (o), and carrier female (o). In some females, the inheritance of paternal and maternal alleles can be ascertained, and in these the paternal X chromosome is shown on the left. Recombinants between HPT and each allele are indicated by an asterisk. Deduced genotypes are shown in brackets.

Subject IV.4 is a carrier mother who is heterozygous for F9, DXS98, DXS52, and DXS15. Her affected son, V.1, is recombinant for HPT and the proximal locus F9, but nonrecombinant for DXS98, DXS52, and DXS15. Subject II.4 is a deceased carrier mother whose genotype was deduced from her 11 children. Her carrier daughter, III.12, is recombinant for HPT and the distal locus DXS52 and DXS15 and for the proximal locus DXS37, but nonrecombinant for F9 and DXS98; whereas the unaffected son, III.6, is recombinant for HPT and the distal group of loci DXS98, DXS52, and DXS15, and nonrecombinant for the proximal locus DXS37. The minimum number of total recombinants in this pedigree is therefore obtained by locating HPT between F9 and DXS98.
segregation of the disease (HPT) with the alleles (N, a, 3, and t), defined respectively by the polymorphic loci F9, DXS98, DXS52, and DXS15. Her other affected son, VI.1, reveals segregation of HPT with the distal loci DXS98, DXS52, and DXS15 (alleles a, 3, and t), but demonstrates recombination between HPT and the proximal locus F9 (allele n*). This observation locates HPT distal to the F9 locus. Analysis of the children of individual II.4 further helps to localize HPT. The affected male III.15 shows that in this branch of the family HPT is segregating with the alleles (E, a, 3, and t). His carrier sister, III.12, is recombinant for the HPT and the distal loci DXS52 and DXS15 and for the proximal locus DXS37, but nonrecombinant for DXS98. This observation locates HPT proximal to DXS52 and distal to DXS37. The combined observations of multipoint crossovers from III.12 and VI.1, locate HPT distal to F9 and proximal to DXS52, i.e., in the vicinity of DXS98. Examination of the multipoint crossovers in the unaffected male, III.6, locates HPT proximal to DXS98; this individual, who has inherited the alleles (a, 3, and t) but has not inherited the disease, demonstrates recombination between HPT and the distal loci DXS98, DXS52, and DXS15, and indicates that the location of HPT is not in the chromosomal segment distal to DXS98. Thus, the combined observations from all the multipoint crosses suggest that HPT is located distal to F9 and proximal to DXS98. The likelihood of this location of HPT versus the other possible locations of HPT within the fixed order Xcen–DXS37–F9–DXS98–DXS52–DXS15–Xqter was quantitatively assessed using the LINKMAP program.

Analysis using the LINKMAP program yielded the location score curve shown in Fig. 3. There is a high peak distal to the F9 locus and proximal to the DXS98 locus, maximum location score = 25.11, 0.03 Morgans proximal to the DXS98 locus. There are five subsidiary peaks between Xcen and DXS37, between DXS37 and F9, between DXS98 and DXS52, between DXS52 and DXS15, and between DXS15 and Xqter. The location score for each peak is 6.08 at 0.20 Morgans proximal to DXS37, 19.40 at 0.05 Morgans proximal to F9, 18.14 at 0.02 Morgans distal to DXS98, –22.34 at 0.006 Morgans proximal to DXS15, and 11.05 at 0.15 Morgans distal to DXS15. The odds ratio for location of HPT in each one of these segments is shown in Table II. This reveals that the odds ratio favoring the order Xcen–DXS37–F9–HPT–DXS98–DXS52–DXS15–Xqter versus a location of HPT unlinked to this cluster of five loci is > 280,000:1. In addition, the odds ratios favoring two other locations of HPT within the framework of the five loci are also significant. The odds ratio for a location of HPT proximal to F9 and distal to DXS37 is 16,300:1, and the odds ratio for a location of HPT distal to DXS98 and proximal to DXS52 is 8,700:1. Thus, a distal as opposed to proximal location of HPT to F9 is 17 times (i.e., 283,500 ± 16,300) more likely, and a proximal as opposed to distal location of HPT to DXS98 is 32 times (i.e., 283,500 ± 8,700) more likely. These results indicate that HPT maps between F9 and DXS98. All the other odds ratios for possible locations of HPT within this framework of loci are < 1,000:1. These results of multipoint linkage analysis demonstrate that the most likely order of genetic loci is Xcen–DXS37–F9–HPT–DXS98–DXS52–DXS15–Xqter.

Discussion

Our linkage study of X-linked recessive idiopathic hypoparathyroidism (HPT) using 17 polymorphic markers to explore X chromosome recombination has established linkage between HPT and the DXS98 locus defined by the polymorphic marker 4D.8, with the probability in favor of linkage > 6,500:1. The genetic marker 4D.8 has been previously localized by somatic cell hybrid and linkage studies to Xq26–Xq27 (17, 28), and a further analysis with the five genetic markers from this distal segment of Xq established that the probability favoring linkage between HPT and this group of loci was 280,000:1. Thus, the results of our study demonstrating linkage between the disease gene and these genetic markers map HPT to the distal region of the long arm of the X chromosome.

Analysis of the multipoint crosses observed within this distal region of the long arm of the X chromosome, in the pedigree in Fig. 2, suggests that HPT maps between the DXS98 and F9, which encodes coagulation Factor IX. The LINKMAP program is able to use information from a number of multipoint crosses to calculate the most likely location of one

Table II. Order of Genetic Loci and Their Respective Odds Ratios as Calculated from the Location Score Curve in Fig. 3

<table>
<thead>
<tr>
<th>Locus order</th>
<th>Peak location score</th>
<th>Odds ratio</th>
</tr>
</thead>
</table>
unmapped gene in a framework of well-mapped markers (27). Within the order Xcen–DXS37–F9–DXS98–DXS52–DXS15–Xqter, a location of HPT between DXS98 and F9 was favored above all other locations (Table II). The odds favoring the location of HPT proximal to DXS98 are 32:1 and those favoring a location distal to F9 are 17:1. Thus, the locus order is indicated and reveals two bridging markers for X-linked recessive idiopathic hypoparathyroidism, with the genetic distances between HPT and its nearest markers, DXS98 and F9, being 3 and 4 centi-Morgans (cM), respectively. These genetic distances usually correspond to physical distances of three and four million nucleotide basepairs, but HPT, DXS98, and the F9 loci are likely to be physically much nearer, as the higher frequency of genetic recombination in this region has previously led to an overestimation of physical distances (17, 26). However, the mutation causing HPT does not involve the DXS98 and F9 loci as recombinants have been observed. In addition, plasma Factor IX concentrations, which were assessed in two affected males from family W/81 subsequent to our localization of HPT, were within the normal range, further indicating that the mutation does not involve the F9 locus. The results of our study, which reveals the bridging markers DXS98 and F9, will be useful in the genetic counseling of some affected families.

The mapping of the HPT gene to Xq26–Xq27 demonstrates that a mutation at a locus distant from the PTH gene, whose location (7) is on the short arm of chromosome 11, is involved in altering parathyroid gland function. A possible role for this X-linked gene in parathyroid gland development is suggested by the neonatal or early infantile onset of hypocalcemic seizures in the two families. This suggests that the disorder may be due to parathyroid agenesis or hypoplasia, and a careful autopsy of a patient from one family has supported this (11). Thus, the HPT gene would appear to be important for the embryological development of the parathyroids and the situation may be analogous to that occurring in the DiGeorge syndrome. In this syndrome there is a failure of development of the derivatives of the third and fourth pharyngeal pouches with resulting absence or hypoplasia of the parathyroids and thymus. The syndrome may be inherited as an autosomal dominant disorder (29), and an association between the syndrome and a deletion of the proximal part of the long arm of chromosome 22 has been reported (30, 31). The deletion breakpoint for the DiGeorge syndrome has been further characterized by molecular genetic studies using in situ hybridization (32) and was found to be proximal to the locus for the immunoglobulin lambda polyepptide constant region but distal to the locus for the DNA probe D2259. The precise mapping of the HPT gene to Xq26–Xq27 by our linkage study represents an important step toward understanding this genetic component of parathyroid gland formation. Our localization of this mutant gene identifies the chromosomal segment in which a concentrated search for deletions and closer genetic markers is required to further elucidate the factors controlling parathyroid development and calcium homeostasis.

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References


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