Heterogeneity in the gene locus for steroid 21-hydroxylase deficiency

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SUMMARY DNA was analysed from 33 patients with congenital adrenal hyperplasia due to steroid 21-hydroxylase deficiency. In each case Southern blots were prepared from a number of restriction enzyme digests and hybridised with probes for both the 21-hydroxylase and the adjacent fourth component of complement (C4).

Evidence for deletion of the active 21-hydroxylase gene (CYP21B) was found in 13 cases and in 10 of these the deletion included the adjacent C4B gene, leading to a hybrid CYP21A/CYP21B gene. Deletion of CYP21B alone was found in one patient, the remaining two cases appearing to have the active gene replaced by the inactive pseudogene. Duplications of the CYP21A–C4B region and deletion of the pseudogene are also described. In a further 12 cases no gross abnormality could be found.

The genes coding for the steroid 21-hydroxylase enzyme (EC 1.14.99.10, CYP21) are located on the short arm of chromosome 6, 3' to the genes coding for the fourth component of complement, C4A and C4B. Deficiency of this enzyme leads to congenital adrenal hyperplasia (CAH) and has been shown to be caused by a heterogeneous range of defects, including deletions of the active (21-hydroxylase B) gene1,2 and possible point mutations and gene conversions.3,4 This paper summarises the findings in a group of 33 patients with 21-hydroxylase deficiency from 31 families.

Patients

The majority of patients had presented in the neonatal period as salt losers (SL, n=24) or as simple virilisers (SV, n=3). Six had presented as late-onset (LO) cases at ages ranging from five to 20 years.

FIG 1 Simplified restriction enzyme map of the C4/21-hydroxylase region of chromosome 6.

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Methods

Genomic DNA was isolated from peripheral blood leucocytes by the method of Bell et al. Samples of 10 μg were digested with restriction endonucleases according to the manufacturer's instructions (Pharmacia Ltd). Digests were separated on 0.75% agarose gels, treated according to Wahl et al., and blotted onto nitrocellulose. Hybridisation conditions and probes used were as previously described. Their location in the C4/21-hydroxylase gene region is shown in fig 1. C4 allotyping was carried out after neuraminidase treatment of EDTA plasma.

Results

Interpretation of results required the use of all probes and a variety of restriction endonucleases. The fragments these produce are shown in fig 1. Most useful was the analysis of TaqI digests with both probes A and K which gave information on the C4 (A and B) and 21-hydroxylase (CYP21A and B) genes respectively, as described by Schneider et al. In addition, a double digest of DNA with the enzymes BglII and EcoRI, as previously described, gives information on both the 5' and 3' ends of each 21-hydroxylase gene.

Twelve patients (10 SL, 2 LO) had intact C4 and 21-hydroxylase genes on both chromosomes, giving a normal pattern with all enzymes used (fig 2, lane 1).

Homozygous deletions of the active CYP21B locus in this group of patients are rare (1/33) (fig 2, lane 2). This SL patient had a deletion extending from the 3' end of the 21A gene to the 3' end of the 21B gene, including the C4B locus, and has been previously described. An apparent heterozygous deletion of the same region occurred in a further nine cases (1 SV and 8 SL) (fig 2, lane 3). This was associated with an HLA-B47 haplotype in only two of these cases.

One SL patient appeared to have a deletion of the 21B gene alone. The results of a TaqI digest using probes A and K are shown in fig 2, lane 4. No 3.7 kb band is present with probe K, while probe A shows three bands of 7 kb, 6 kb, and 5.4 kb respectively, the first being of approximately twice the intensity of each of the other two. This indicates that not only is the C4B locus intact, but that both a 'short' and 'long' C4B gene are present (bands at 6 kb and 5.4 kb). The 3 kb Kpn band was also absent. The data suggest either that both chromosomes have a single deleted CYP21B or that there are two CYP21A genes present on each chromosome and no CYP21B gene, that is, gene conversion has occurred. In this particular subject, the 3' end of the CYP21B gene appears to be unchanged, as a double digest using

![Fig 2 Southern blots of TaqI digests of DNA from six patients with 21-hydroxylase deficiency.](http://jmg.bmj.com/)

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BglII/EcoRI showed a deletion of the 2.4 kb band located at the 5' end of 21B and not of the 3' 8 kb band (fig 3a). One other SL case was heterozygous for a 'deleted' 'converted' CYP21B (fig 3b).

Two patients (1 SL, 1 SV) in whom the CYP21B gene appeared to be intact had the C4A-CYP21A genes deleted on one chromosome. This led to a reduced intensity of the 3-2 kb TaqI, 4 kb KpnI, and 10 kb EcoRI/BglII bands with probe K, in addition to the presence of the 6-4 kb TaqI fragment detected with probe A, a marker for deletion of the C4A gene9 (fig 2, lane 5). C4 protein typing confirmed the presence of a C4A null allele.

A duplication of the 21A-C4B locus was found in two LO patients, as expected in association with HLA-B14, DR1. This gave the pattern of a heterozygous deletion of 21B, that is, reduced intensity of the 3-7 kb TaqI, 3 kb KpnI, 8 and 2-4 kb BglII/EcoRI, and 11 kb BglII fragments, as detected with probe K. With probe A, however, the bands corresponding to the 5' region of the C4B gene were of greater intensity than C4A with both the TaqI (fig 2, lane 6) and BglII digests (data not shown). C4 protein typing of one family showed the presence of three C4B genes in the affected child (C4A2,0B1,1,2), two of which had been inherited from the mother on a single chromosome. One other LO patient showed similar results with probe K, that is, reduced intensity of the 2-4 and 8 kb bands after BglII/EcoRI double digest. However, no evidence for duplication, crossover, or recombination was found either with probe A or C4 protein allotyping. One must therefore assume that this is a true deletion of the whole CYP21B gene.

The digests from the remaining four (1 LO, 1 SV, 2 SL) patients appeared to show reduced intensity of one 21-hydroxylase gene with one enzyme but not with the others. This may be the result of loss of a single restriction enzyme site which can only be confirmed by sequencing the CYP21A and CYP21B genes in these cases.

Discussion

The most important finding in this study is the range of genetic defects which can give rise to the clinical presentation of 21-hydroxylase deficiency. The interpretation of results in these patients is difficult and requires the use of more than one restriction enzyme and probes for both C4 and 21-hydroxylase.

In those cases in which a normal restriction fragment pattern is evident, one can only assume that an undetected point mutation is present in either the gene or flanking sequences. Rodrigues et al9 cloned a single intact, but presumably defective, CYP21B gene and found a total of 11 nucleotide differences, two of which were in exons leading to amino acid substitutions in the protein molecule. In the same patient, there was also some evidence for partial 'gene conversion', the defective 21B containing some changes which gave rise to the nucleotide sequence found in the CYP21A gene. Two unrelated patients in our study show evidence of possible gene conversion; in both the characteristics of the 5' end of the 21B gene are missing, the 3' end apparently remaining intact. Similar results were reported by Donohoue et al,4 Jospe et al,10 and Harada et al,11 the latter showing by partial nucleotide sequencing analysis that the 21-hydroxylase B gene has been converted to the pseudogene.

Deletion of the C4A-CYP21A locus in two of our cases may be coincidental, as subjects with homozygous deletion of CYP21A (as in HLA-B8, DR3) are negatively associated with CAH.12 An undetected fault could lie in the CYP21B genes present.

FIG 3 Southern blot of BglII/EcoRI digests of DNA from patients with 21-hydroxylase deficiency compared with controls (c).
The HLA-B14, DR1 haplotype associated with late onset CAH has been shown to have a duplication of the C4B\textsuperscript{13} and CYP21A\textsuperscript{14} loci. These findings are confirmed here. It is still not clear, however, why such a duplication should lead to defective enzyme activity in the presence of an apparently normal 21B gene.

In all patients assigned a deleted CYP21B gene in this and our previous study,\textsuperscript{1} the deletion was confirmed using several enzymes. With one exception the deletion included the C4B gene (confirmed with probe A). The gene is thus a hybrid with 5\textsuperscript{'} characteristics of the CYP21A and those of the CYP21B gene at the 3\textsuperscript{'} end, as previously described.\textsuperscript{1,15} This point was overlooked by Matteson et al\textsuperscript{16} who showed the 3-7 kb Taq fragment to be absent but after other enzyme digests failed to confirm deletion of CYP21B.

In conclusion, gene deletion, conversion, and duplication have been found in 21 of the cases of 21-hydroxylase deficiency described here, with the possibility in the remaining 12 cases of point mutations in the 21-hydroxylase gene.

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References


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