Exclusion of the Locus for Autosomal Recessive Pseudohypoaldosteronism Type 1 from the Mineralocorticoid Receptor Gene Region on Human Chromosome 4q by Linkage Analysis*


Department of Pediatrics, University College London Medical School, Rayne Institute (E.C., M.R., R.T., R.M.G.), London, United Kingdom WC1E 6J; the Department of Pediatrics, E. Wolfson Hospital, and Tel Aviv University, Sackler School of Medicine (A.H.), Tel-Aviv, Israel; the Medical Unit, Division of Medicine, Institute of Child Health and Renal Unit, Great Ormond Street Hospital for Children National Health Service Trust (M.D.), London, United Kingdom WC1N 1EH; the Research Institute, College of Judea and Samaria (I.H.), Ariel 44820, and Institute of Endocrinology, Sorasky Medical Center (I.H.), Tel-Aviv, Israel; the Department of Pediatrics, Assaf Harofeh Medical Center (T.B.), Zerifin, and Tel-Aviv University, Sackler School of Medicine (T.B.), Tel-Aviv, Israel; University of Munich, Children’s Hospital (U.K.), D-80337 Munich, Germany; and the Department of Medicine, University of Edinburgh, Western General Hospital (J.S.), Edinburgh, Scotland EH4 2XU

ABSTRACT

Pseudohypoaldosteronism type 1 (PHA1) is an uncommon inherited disorder characterized by salt-wasting in infancy arising from target organ unresponsiveness to mineralocorticoids. Clinical expression of the disease varies from severely affected infants who may die to apparently asymptomatic individuals. Inheritance is Mendelian and may be either autosomal dominant or autosomal recessive. A defect in the mineralocorticoid receptor has been implicated as a likely cause of PHA1. The gene for human mineralocorticoid receptor (MLR) has been cloned and physically mapped to human chromosome 4q31.1–31.2. The etiological role of MLR in autosomal recessive PHA1 was investigated by performing linkage analysis between PHA1 and three simple sequence length polymorphisms (D4S192, D4S1548, and D4S413) on chromosome 4q in 10 consanguineous families. Linkage analysis was carried out assuming autosomal recessive inheritance with full penetrance and zero phenocopy rate using the MLINK program for two-point analysis and the HOMOZ program for multipoint analysis. Lod scores of less than −2 were obtained over the whole region from D4S192 to D4S413 encompassing MLR. This provides evidence against MLR as the site of mutations causing PHA1 in the majority of autosomal recessive families. (J Clin Endocrinol Metab 80: 3341–3345, 1995)

PSEUOHYPOALDOSTERONISM type 1 (PHA1) is an uncommon inherited disorder characterized by salt-wasting in infancy and associated with failure to thrive, hyponatremia, and hyperkalemia (1, 2). Marked elevation of serum aldosterone levels is present in all cases, and PRA is increased in most (3). Patients are unresponsive to mineralocorticoids, but respond to salt supplementation which can often be discontinued after infancy. Since the first report by Cheek and Perry in 1958 (4), over 100 cases have been reported.

Clinical expression of the disease varies from severely affected infants who may die to apparently asymptomatic individuals (5). This depends in part on whether mineralocorticoid resistance is confined to renal tubules (renal form) or involves multiple target organs, including sweat and salivary glands and colon (generalized form). Inheritance is Mendelian and may be either autosomal dominant or recessive (6, 7). It has been suggested that the renal form is usually milder and is inherited as an autosomal dominant (AD) trait. Families with the generalized form are usually more severely affected and display an autosomal recessive (AR) inheritance with an increased incidence of consanguinity.

The molecular basis of PHA1 is unknown. Cheek and Perry suggested that “the disorder may be due to a refractory state on the part of the renal tubules to endogenous mineralocorticoids,” an end-organ hormone insensitivity syndrome. A primary defect in the human mineralocorticoid receptor (hMR) has remained the most plausible hypothesis for the molecular basis of PHA1. Since Albright’s description of pseudohypoparathyroidism in 1942 (8), many hormone resistance syndromes have been described. These include resistance to insulin, androgens, vitamin D, cortisol, T₄, and vasopressin. Causative mutations have been described in several members of the steroid receptor superfamily of genes, including those for androgen, calcitriol, and glucocorticoids.

In 1985, Armanini and co-workers demonstrated reduced or absent binding sites for tritiated aldosterone in circulating...
mononuclear lymphocytes in three patients with PHA1 (9, 10). These observations have been confirmed in subsequent studies (11–13) and provide evidence in favor of a defect in hMR in PHA1. However, the possibility exists that they may represent a secondary effect of down-regulation due to high aldosterone levels.

Molecular cloning of a complementary DNA (cDNA) encoding hMR (14) has allowed a direct genetic approach to evaluating the hypothesis that PHA1 arises from mutations in the gene for this receptor. Analysis has been carried out by 3 independent groups on 7 patients, including the original sporadic case of Cheek and Perry. In this patient and his family, Southern blot analysis using 3 probes from the hMR cDNA revealed bands of normal size and intensity (15). In another patient and his family, hMR cDNA was amplified using 11 pairs of oligonucleotide primers, and total ribonucleic acid from peripheral blood leukocytes was amplified by reverse transcriptase-polymerase chain reaction (PCR). Sequencing of the products showed no deviation from the wild-type sequence (16). Similar results were reported in a study of 5 other patients with PHA1 (17, 18). Sequencing of the patient’s hMR cDNA identified a nonconservative homozygous base change in 4 of the 5 patients, and 1 of them had an additional conservative heterozygous base change. Both of these base changes were in the immunogenic domain and were also found to be present in a considerable number of normal individuals. It was concluded that these sequence variations did not have any pathological significance. The sequences of the first untranslated exon, 0.9 kilobase of the 5′-regulatory region, and the DNA- and ligand-binding receptor domain in these 5 patients were all normal. Full evaluation of the gene for hMR (MLR) by direct mutational analysis awaits characterization of its genomic organization. These results do not, of course, exclude MLR as the site of mutations causing PHA1, particularly in the recessive variety, as genetic ( locus or allelic) heterogeneity may exist.

Homozygosity mapping provides a powerful approach to testing the role of a candidate gene in a rare autosomal recessive disorder with a high level of parental consanguinity (19). MLR has been mapped to human chromosome 4 by somatic cell hybridization and regionally localized to 4q31.1–31.2 by in situ hybridization (20, 21). Linkage analysis was, therefore, carried out in 10 inbred families (see Fig. 1) using 3 simple sequence length polymorphisms spanning the MLR region.

Subjects and Methods

Patients and families

Ascertainment of families has been achieved through collaboration with the physicians who had reported cases of PHA1 in the literature and British pediatric endocrinologists or nephrologists who care for children with PHA1. The 10 families included in this study are shown in Fig. 1. All parents are consanguineous (all were first cousins except in 2 families), and autosomal recessive inheritance is assumed. There were 2 affected individuals in 6 of the families. The study includes a total of 39 individuals, 14 of whom were affected (5 males and 9 females).

Diagnosis was made according to standard and clinical biochemical criteria. All affected individuals had documented elevated urinary sodium in the presence of hyponatremia, hyperkalemia, increased PRA, elevated serum aldosterone level, and normal renal and adrenal func-

FIG. 1. PHA1 pedigrees used for linkage analysis. All affected individuals are offspring of marriages between first cousins, except in two families (010 and 012) as shown.

DNA analysis/marker typing

DNA was extracted from white cells by standard protocols (24). The primer sequences and conditions for the three markers (D4S192, D4S548, and D4S413) have been described previously (25, 26). DNA was amplified by PCR and labeled by incorporating [α-32P]deoxynucleotide. PCR was performed in 36-well microtiter plates (Hybaid). Each well contained 50–100 ng genomic DNA; 1.5 mmol/L MgCl2; 1 × reaction buffer [16 mM (NH4)2SO4, 67 mM Tris-HCl (pH 8.8 at 25°C), 0.01% Tween-20; Bioline, UK]; 200 pmol/L each of deoxy (d)-GTP, dATP, and dTTP; 20 μmol/L dCTP; 0.05 μL [α-32P]dCTP (3000 Ci/mmol); 50 ng of each primer; and 0.2 μl BIOTAG polymerase (Bioline) in a total volume of 20 μL. Thirty cycles were performed in a thermocycler (Hybaid Omnigene, Hybaid Ltd, UK). Alleles were separated by 6% polyacrylamide gel electrophoresis and detected by autoradiography.
Linkage analysis

The maps used in the chromosome 4q analysis were from the reports of NIH/CEPH Collaborative Mapping Group (25), Gyapay et al. (26), and Mills et al. (27). The relative positions of MLR and the three simple sequence length polymorphisms used (D4S192, D4S1548, and D4S413) are shown in Fig. 2. Linkage analysis was carried out assuming a fully penetrant autosomal recessive inheritance with a disease allele frequency of 0.0002 and a zero phenocopy rate. Lod scores (logarithms to the base 10 of odds that two loci are linked) were calculated using the MLINK (~5.1) program (28) for two-point analysis and the HOMOZ program (29) for multipoint analysis. The markers were assumed to have five alleles of equal frequencies. Apparently normal siblings were not included in the analysis.

Results

The alleles of the affected individuals at the three loci studied are shown in Table 1. Loci were homozygous on only 10 of 41 occasions among the 14 affected individuals typed at the 3 markers (2 for D4S192, 4 for D4S1548, and 4 for D4S413 were homozygous). The 2-point lod scores are shown in Table 2. The lod score was less than -2 for at least 10 centimorgans (CM) around D4S192 and D4S413 and for 5 CM around D4S1548. The results of the multipoint analysis are shown in Fig. 3. A lod score of less than -2 was obtained over the whole region from D4S192 to D4S413.

Discussion

These results provide strong genetic evidence against MLR as the site of mutations causing pseudohypoaldosteronism in these families. Linkage analysis in inbred families provides a powerful strategy for examining the role of a candidate gene and for identifying the true location of the gene(s) causing this disease. Aspects of the interpretation of the linkage data and consideration of the possible nonreceptor molecular defects underlying AR PHAl and strategies for future investigations are discussed below.

The increased incidence of rare recessive diseases in the offspring of consanguineous matings arises in general from affected individual inheriting disease alleles at a particular locus from a common ancestor. The affected individual is, therefore, said to be homozygous by descent (rather than by state) for the disease allele. This increases the power of linkage data from consanguineous families, a factor exploited in the strategy of homozygosity mapping of disease loci in rare recessive disorders (see below) (19). However, the possibility of a disease chromosome being introduced into a consanguineous family by a married-in unrelated member cannot be totally excluded. This could produce an affected individual who may be heterozygous at the disease locus. This is extremely unlikely, especially if the disease is rare, as is PHAl. Exclusion of a candidate gene is definitively achieved by following the segregation of an intragenic polymorphism, which allows identification of obligate recombinants within families with at least two affected children. As an informative
eralocorticoids allow for dysfunction in pathways that do not
response element of the nuclear DNA.
sterone-induced proteins) by binding to a number of hormone
active complex. This complex initiates protein synthesis (aldo-
corticoid deficiency (31), and pseudohypoparathyroidism
proportion of cases in several hormone insensitivity syn-
ations causing PHAl in at least some patients. This is consistent
with the observation that nonreceptor defects account for a
majority of these families or that a set of linked families
might be identified if a much larger resource of AR PHAI
families was studied. Moreover, they do not exclude MLR as
the locus accounting for PHAI in those families that appear
to display AD inheritance, although it is well recognized that
allelic heterogeneity can account for variations in the mode
of inheritance of diseases arising from a single locus. The
evidence arising from cDNA analyses in sporadic patients
does not, however, support MLR as the site of mutation
in patients who may fall into the AD category, although the
possibility that mutations exist in noncoding regulatory
regions has not been excluded.
These observations indicate that MLR is not the site of muta-
tions causing PHAI in at least some patients. This is consistent
with the observation that nonreceptor defects account for a
proportion of cases in several hormone insensitivity syn-
dromes, including androgen insensitivity (30), familial glu-
cocorticoid deficiency (31), and pseudohypoparathyroidism
(32). The likely molecular complexity underlying aldosterone
function does, of course, provide numerous possible sites at
which hormone insensitivity might arise. There is now good
evidence that aldosterone exerts an immediate effect on its
target cells via a membrane-bound receptor (now known as
nongenomic action) (33). This is followed by the classical
genomic action of aldosterone, which is thought to act by bind-
ing to the cytoplasmic mineralocorticoid receptor, forming an
active complex. This complex initiates protein synthesis (aldo-
sterone-induced proteins) by binding to a number of hormone
response element of the nuclear DNA.
Many different hypotheses for nonreceptor defects in
PHAI have been proposed. The nongenomic actions of min-
eralocorticoids allow for dysfunction in pathways that do not
include the cytoplasmic receptor. At the prereceptor level,
there could be interference with aldosterone binding, by
competition for the binding site or by cleavage of the recep-
tor. At the postreceptor level, the most obvious candidates
include the aldosterone-induced proteins, which include subunits of Na/K-adenosine triphosphatase and the amilo-
ride-sensitive epithelial Na channels. Liddle’s syndrome, or
pseudoadosteronism, has recently been shown to arise from
mutations in the β-subunit of the epithelial Na channel gene
causing excessive sodium reabsorption and hypertension at
normal or low levels of circulating aldosterone (34). In so far
as these features represent a mirror image of PHAI, it is
conceivable that mutations involving these sodium channels
may render them nonfunctional or unresponsive to aldoste-
rone.
Homzygosity mapping represents a powerful strategy
for examining the role of additional candidate genes in
PHAI. This is being pursued using the present family re-
source in parallel with a genome search using anonymous
microsatellite loci in case the causative locus is an as yet
unidentified gene. Identification of the gene for PHAI will
allow improved diagnosis and treatment in affected indi-
viduals. It should also provide new insights into the biology
of mineralocorticoid function at a molecular level, informa-
tion of potential value in understanding more common dis-
ase states such as hypertension.

Acknowledgments
This study would have been impossible without the cooperation of
the patients and their families and the collaborating doctors, including
Dr. J. W. T. Bevson (Blackburn, UK), Dr. D. M. Johnston (Petersborough,
UK), and Dr. P. G. F. Swift (Leicester, UK).

References
1. Komesaroff PA, Funder JW, Fuller PJ. 1994 Mineralocorticoid re-
2. Dillon MJ, Leonard JV, Buckler JM, et al. 1980 Pseudohypoal-
3. Pasiol-Vinay MC, Alhouri CM, Ricour C, Limal JM, Rappaport R,
Royer P. 1974 Pseudohypoaldosteronism: persistence of hyperal-
dosteronism and evidence for renal tubular and intestinal responsiv-
ness to endogenous aldosterone. J Clin Endocrinol Metab. 39:1038–1044.
5. Chitayat D, Spiter Z, Ayalon D, Golander A. 1985 Pseudohypo-
aldosteronism in a female infant and her family: diversity of
74:619–622.
dosteronism in eight families: different forms of inheritance are evident
7. Hanakoglu A. 1991 Type I pseudohypoaldosteronism includes two
clinically and genetically distinct entities with either renal or multi-
8. Albritton J, Burnett CH, Smith PH, Parson W. 1942 Pseudohypo-
parathyroidism-an example of 'Seabright-Bantam syndrome'. J Clin
Endocrinol Metab. 30:922–932.
9. Armanini D, Strasser T, Weber PC. 1985 Characterisation of aldo-
sterone binding sites in circulating human mononuclear leukocytes.
10. Armanini D, Kuhnle U, Strasser T, et al. 1985 Aldosterone-receptor