Compound heterozygosity for a whole gene deletion and p.R124C mutation in CYP21A2 causing nonclassic congenital adrenal hyperplasia

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We present a family with 2 members who received long-term steroid treatment for presumed classic congenital adrenal hyperplasia (CAH) due to 21-hydroxylase deficiency, until molecular testing revealed nonclassic CAH, not necessarily requiring treatment. A 17-year-old male presented to our clinic on glucocorticoid and mineralocorticoid treatment for classic CAH. He was diagnosed at 4 years of age based on mild-moderate elevations of 17-hydroxyprogesterone (17-OHP) and adrenocorticotropic hormone (ACTH), but without evidence of precocious adrenarche/puberty. Due to his diagnosis, his clinically asymptomatic 3-year-old sister was tested and also found to have elevated ACTH and 17-OHP levels and was started on glucocorticoids for classic CAH. Family history revealed a healthy sibling who had no biochemical evidence of CAH and consanguineous healthy parents. We questioned the diagnosis of classic CAH and performed an ACTH1-24 stimulation test, which showed a level of 17-OHP in the borderline range between classic and nonclassic CAH. Molecular testing, using sequencing and multiplex ligation-dependent probe amplification analysis of CYP21A2, revealed that both affected siblings were compound heterozygotes for a whole-gene deletion and p.R124C mutation in CYP21A2 causing nonclassic congenital adrenal hyperplasia.

We present a family with 2 members who received long-term steroid treatment for presumed classic congenital adrenal hyperplasia (CAH) due to 21-hydroxylase deficiency, until molecular testing revealed nonclassic CAH, not necessarily requiring treatment. A 17-year-old male presented to our clinic on glucocorticoid and mineralocorticoid treatment for classic CAH. He was diagnosed at 4 years of age based on mild-moderate elevations of 17-hydroxyprogesterone (17-OHP) and adrenocorticotropic hormone (ACTH), but without evidence of precocious adrenarche/puberty. Due to his diagnosis, his clinically asymptomatic 3-year-old sister was tested and also found to have elevated ACTH and 17-OHP levels and was started on glucocorticoids for classic CAH. Family history revealed a healthy sibling who had no biochemical evidence of CAH and consanguineous healthy parents. We questioned the diagnosis of classic CAH and performed an ACTH1-24 stimulation test, which showed a level of 17-OHP in the borderline range between classic and nonclassic CAH. Molecular testing, using sequencing and multiplex ligation-dependent probe amplification analysis of CYP21A2, revealed that both affected siblings were compound heterozygotes for a whole-gene deletion and a, likely pathogenic (nonclassical), sequence variant, p.R124C. The asymptomatic father had the same genotype, while the mother showed one deleted copy and 2 active copies, making her an asymptomatic carrier. Our report demonstrates the importance of molecular testing in atypical cases of CAH, as well as the importance of both sequencing and deletion analysis. The results of molecular testing should be interpreted in clinical context, and treatment should be prescribed according to guidelines when available.

Keywords: CYP21A2, Genetic testing, Nonclassic congenital adrenal hyperplasia

Introduction

Congenital adrenal hyperplasia (CAH) is a group of autosomal recessive disorders characterized by the deficiency of enzymes required for the conversion of cholesterol to glucocorticoids, mineralocorticoids, and adrenal androgens. 21-hydroxylase deficiency (21-OHD) CAH (OMIM #201910) accounts for >90% of all cases, with an incidence rate of approximately 1 in every 10,000 to 20,000 live births, for all forms of CAH. It is caused by homozygous, or compound heterozygous, mutations in CYP21A2, on chromosome 6p21.3. CAH can be broadly divided into early onset classic CAH and late onset nonclassical CAH (NC CAH).

The late-onset phenotype (NC CAH) is characterized by normal serum cortisol and aldosterone levels and absence of early childhood virilization. Enzyme activity in this form ranges from 10% to 40%. Biochemically, variable degrees of hyperandrogenemia are present;
however they most often do not cause symptoms until late childhood or adolescence, with many patients, particularly males, remaining asymptomatic throughout their lives.²³

Biochemical testing is usually the initial diagnostic step in the workup of individuals suspected of having CAH, with 17 hydroxyprogesterone levels being elevated in all forms of 21-OHD. Although the different forms of 21-OHD are distinguished on clinical grounds, molecular testing is being used more often, especially in cases picked up via newborn screening before the clinical phenotype is evident, or where biochemical testing is ambiguous.⁴ Here we present 2 individuals in a family who underwent long-term treatment for classic CAH, before revaluation with molecular testing revealed NC CAH not requiring treatment.

**Case report**

This research related to human use has been complied with all the relevant national regulations, institutional policies and in accordance with the tenants of the Helsinki declaration, and has been approved by the author’s or equivalent committee. Informed consent was obtained from the parents of the patients. The proband was a male born to consanguineous parents (Fig. 1). He had an acute viral fever at 4 years of age, and was found to have mild hyponatremia and hyperkalemia (sodium 131 mEq/L and potassium 6.1 mEq/L, respectively). Further investigation revealed increased 17-hydroxyprogesterone (17-OHP) and adrenocorticotropic hormone (ACTH) levels (Table 1), and a suboptimal rise in cortisol levels following ACTH1-24 stimulation (13.4 μg/dL after 60 minutes). At the time of diagnosis proband was of normal growth and stature (height=107 cm, z-score=−0.4, 32nd percentile; weight=20.8 kg, z-score=+0.8, 80th percentile; body mass index [BMI]=18.2 kg/m²). Treatment was thus begun for presumed salt wasting (SW) CAH using replacement doses of hydrocortisone and fludrocortisone. He had normal male genitalia, descended testes, and no evidence of precocious adrenarche or puberty. He was seen by various providers sporadically over the years, and finally was reviewed in our pediatric endocrinology clinic at 17 years of age. He was on subphysiologic hydrocortisone replacement (7.7 mg/m²/day) as well as fludrocortisone (0.2 mg per day), without any symptoms, no history of prostration with illness, and normally-timed pubertal changes. He was obese with normal stature (height=170 cm, z-score=−0.8, 21st percentile; weight=110 kg, z-score=+2.4, 99th percentile; BMI=38.1 kg/m²), and had normal testes with Tanner stage V pubic hair.

After his diagnosis, the 3-year-old sister was screened for increased 17-OHP and ACTH levels, which were found to be mildly elevated (Table 2). At the time of her diagnosis she was also of normal growth and stature (height=94 cm, z-score=−0.05, 48th percentile; weight=14 kg, z-score=0.03, 51st percentile; BMI=15.8 kg/m²). She was also started on physiologic doses of hydrocortisone after ACTH1-24-stimulation showed suboptimal rise in cortisol levels (13.12 μg/dL after 60 minutes). She also presented to us in the Pediatric Endocrinology clinic at the age of 15 and 11 months, on subphysiologic hydrocortisone replacement (8.33 g/m²/day), with no history of virilization, precocious puberty or prostration with illness. She was obese (height=159 m, z-score=−0.7, 23rd percentile; weight=73.6 kg, z-score=1.4, 92nd percentile; BMI=29.5 kg/m²), with no hirsutism and normal female genitalia (Tanner V breasts and pubic hair).

![Fig. 1. The figure shows that the parents of the proband in a consanguineous relationship as well as his affected sister and 2 unaffected siblings. CAH, congenital adrenal hyperplasia.](image)

**Table 1. Results of biochemical testing of brother**

<table>
<thead>
<tr>
<th>Demographic</th>
<th>Analytes</th>
<th>Results (reference range)</th>
<th>Sampling times (min)</th>
</tr>
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<tbody>
<tr>
<td></td>
<td>17-OHP (ng/dL)</td>
<td>2,620.0 (&lt;100)</td>
<td>0</td>
</tr>
<tr>
<td>Baseline values (5 yr)</td>
<td>ACTH (AM; pg/mL)</td>
<td>52.9 (&lt;46)</td>
<td>-</td>
</tr>
<tr>
<td></td>
<td>Cortisol (AM; μg/dL)</td>
<td>15.1 (6–25)</td>
<td>-</td>
</tr>
<tr>
<td></td>
<td>ACTH1-24 stimulation (17 yr)</td>
<td>Cortisol (μg/dL)</td>
<td>-</td>
</tr>
<tr>
<td></td>
<td>17-OHP (ng/dL)</td>
<td>8,190.0 (10,730.0)</td>
<td>0</td>
</tr>
</tbody>
</table>

AM, morning 8:00; 17-OHP, 17 hydroxyprogesterone; ACTH, adrenocorticotropic hormone; ACTH1-24, synthetic ACTH fragment amino acids 1-24, cortrosyn; NC CAH, nonclassical congenital adrenal hyperplasia.

*Reference ranges/expected values for stimulated cortisol and 17-OHP: Cortisol: >12 μg/dL denotes a normal response; 17-OHP: <1,666 ng/dL—likely unaffected heterozygote (some overlap with NC CAH), 1000–10,000 ng/dL–NC CAH (some overlap with unaffected heterozygotes), >10,000 ng/dL—classical CAH (some minimal overlap with NC CAH).
We questioned the diagnosis of SW CAH, since they were both asymptomatic on subphysiologic hydrocortisone replacement. Work up revealed normal serum electrolytes, mildly increased ACTH and stimulated 17-OHP levels, which when combined with the lack of any symptoms, led us to suspect NC CAH (Tables 1, 2). Thus samples were taken from the family and sent for molecular testing.

Participants were tested for the presence of mutation(s) within the CYP21A2 gene and large rearrangements between CYP21A2 and the CYP21A1P pseudogene. The method used was sequencing and multiplex ligation-dependent probe amplification.

Molecular testing revealed that the proband had one deleted copy of CYP21A2, whereas the other copy harbored a likely pathogenic sequence variant (c.748C>T,p.R124C). The p.R124C variant has been reported previously in an individual with a diagnosis of CAH.3 Previous functional studies indicated that this variant has approximately 16% residual activity.3 The arginine (R) at this position is highly conserved across species and there is a large physicochemical difference between arginine (R) and cysteine (C).9 In silico analysis, based on SIFT, PolyPhen-2, MutationTaster, and Align GVGD suggests that this variant might affect protein function based on the amino acid change. SIFT categorized these changes as damaging. PolyPhen-2 scored it as probably damaging with HumDiv and HumVar scores of 1.000 and 0.997, respectively (scale: 0.000 – not damaging – to 1.000 – damaging). MutationTaster classified it as ‘disease causing’ with a score of 180 (scale: 0.00 – not disease causing – to 215 – disease causing). Finally, AlignGVGD gave it a GD score of 70.97, which puts it into the most ‘likely damaging’ prediction Class C65 (scale: Class C0 – not damaging – to C65 – most likely to be damaging).6,7 Taken together, these findings suggested that p.R124C is likely pathogenic. If p.R124C is pathogenic, the most likely phenotype due to compound heterozygosity for p.R124C and a large deletion would be NC CAH.

The proband’s sister had the same genotype. To our surprise, family studies indicated that the father had the same genotype as the 2 children, with one copy deleted and the other harboring the p.R124C mutation. The mother was heterozygous for the large deletion. Therefore, a diagnosis of NC CAH was confirmed, and consequently, we discontinued the current treatment regimen, with regular follow-up visits. Both siblings remain asymptomatic without medication, with mild, expected laboratory derangements, proband’s sister had a 17-OHP levels (9.072 ng/dL) 1 year after the cessation of medication. Follow-up testing could not be performed on the proband.

**Discussion**

Classic CAH and NC CAH differ widely in their phenotypes, and thus it is important for clinicians to be able to differentiate between the two.

The diagnosis of CAH is primarily based upon the history, physical examination and laboratory values of 17-OHP. The Endocrine Society has outlined the diagnostic criteria in its consensus statement to aid physicians in the process when evaluating patients for 21-OH CAH.2 A useful diagnostic cutoff for classic CAH is a stimulated 17-OHP level >10,000 ng/dL, 1,000–10,000 ng/dL for NC CAH and below 1,666 ng/dL for those not affected; this last category will include heterozygotes as well as some normal individuals (particularly those who are physically stressed, e.g., preterm babies). In addition, an ACTH-stimulation test and corresponding 17-OHP values are used to rule in- or rule out disease.2 However, where the above testing inconclusive, such as in this case where the 17-OHP was in the range of classic CAH in the brother despite a lack of symptoms, molecular testing can be helpful in revealing the underlying defect.3

As mentioned earlier, NC CAH is due to a mutation and/or deletion of the CYP21A2 gene that leaves the affected individual with a total enzyme activity that is less than 50% of a normal person, but larger than about 10%. More severe functional impairment is associated with classic CAH.

CYP21A2 and its associated pseudogene(s) (CYP21A1P) are located on chromosome 6 within the HLA locus, the most highly recombinant portion of the human genome. Not surprisingly, this leads to frequent genomic rearrangements between CYP21A2 and its highly homologous pseudogene, CYP21A1P. These events might create large deletions (involving parts or all of CYP21A2 and CYP21A1P, or only CYP21A2).

<table>
<thead>
<tr>
<th>Table 2. Results of biochemical testing of sister</th>
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<tbody>
<tr>
<td>Demographic</td>
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<tr>
<td>Baseline values (5 yr)</td>
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<tr>
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recombinant hybrids of gene and pseudogene, or microgene conversions of CYP21A2, where small portions of CYP21A1P get transferred into CYP21A2.\(^4\)

Guidelines set out by the Endocrine Society dictate that classic CAH be treated with mineralocorticoids and sodium chloride supplements in infancy and Hydrocortisone should be given as maintenance therapy. While, symptomatic patients with NC CAH should receive the same treatment as their CAH counterparts, asymptomatic patients should not receive any treatment. Unwarranted diagnosis and treatment not only burdens the healthcare infrastructure, but is bothersome for the patients even though they are physically healthy.

To conclude, asymptomatic patients with NC CAH do not require aggressive treatment and repetitive diagnostic testing is unhelpful.\(^8\) Genetic evaluation is important as it identifies the specific defect and is helpful when the diagnosis is uncertain. In addition, it is also vital to screen the family with genetic testing as it will assist in genetic counseling for future offspring and allow identification of asymptomatic carriers.

**Conflict of interest**

No potential conflict of interest relevant to this article was reported.

**Acknowledgments**

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**References**