



Detection of new potentially pathogenic mutations in two patients with primary pigmented nodular adrenocortical disease (PPNAD) — case reports with literature review

Wykrycie nowych, potencjalnie patogennych mutacji u dwóch pacjentów z pierwotną pigmentową chorobą guzkową nadnerczy (PPNAD) — opis przypadków wraz z przeglądem piśmiennictwa

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Abstract

Introduction: Primary pigmented nodular adrenocortical disease (PPNAD) is a rare form of ACTH-independent Cushing's syndrome (CS). Half of patients with PPNAD are sporadic cases and the other half familial.

Material and methods: We present two patients with PPNAD confirmed by genetic analysis.

Results: In both patients there were no abnormal findings on diagnostic imaging of both adrenals and heart. Patients underwent bilateral two-stage adrenalectomy. Histopathological examination confirmed PPNAD. Genetic testing showed the following mutations in the *PRKARIA* gene coding for the regulatory subunit type 1A of the protein kinase A enzyme: c.125dupG (patient 1) and c.15dupT (patient 2). Both these defects lead to inactivation of the *PRKARIA* protein and are consequently causative of PPNAD in these patients.

Conclusions: The novel mutations presented in this article are considered to be pathogenic for PPNAD. (*Endokrynol Pol* 2018; 69 (6): 675–681)

Key words: Cushing's syndrome; nodular adrenocortical disease; Carney complex; PPNAD

Streszczenie

Wstęp: Pierwotna pigmentowa choroba guzkowa nadnerczy (PPNAD) jest rzadką formą ACTH-niezależnego zespołu Cushinga (CS). W połowie przypadków PPNAD występuje w postaci sporadycznej, w pozostałej połowie rodzinie.

Materiały i metody: Przedstawiono opis przypadków 2 pacjentów z PPNAD potwierdzonym badaniem genetycznym.

Wyniki: U obojga pacjentów nie znaleziono nieprawidłowości w badaniach obrazowych nadnerczy i serca. Chorzy przeżyli obustronną etapową adrenalectomię. W badaniu histopatologicznym potwierdzono PPNAD. W badaniu genetycznym wykazano następujące mutacje w genie *PRKARIA* kodującym jednostkę regulatorową typu 1A kinazy proteinowej A: c.125dupG (pacjent 1) i c.15dupT (pacjent 2). Oba te defekty prowadzą do inaktywacji białka *PRKARIA* i w konsekwencji są przyczyną PPNAD u opisywanych pacjentów.

Wnioski: Nowe mutacje przedstawione w niniejszym artykule są rozważane jako patogene w PPNAD. (*Endokrynol Pol* 2018; 69 (6): 675–681)

Słowa kluczowe: zespół Cushinga; guzkowa choroba nadnerczy; zespół Carneya; PPNAD

Introduction

Primary pigmented nodular adrenocortical disease (PPNAD) is a rare form of ACTH-independent Cushing's syndrome (CS). Primary adrenal hypercortisolism is responsible for 10–20% of all cases of CS caused by adrenal adenomas (unilateral) and less often adrenal carcinomas (usually unilateral) and bilateral adrenal hyperplasia (BAHs) [1]. The latter is classified by the size of the nodules on high-resolution computed tomography (CT) to:

smaller than 1 cm (micronodular hyperplasia) or larger than 1 cm (macronodular hyperplasia) and the presence or absence of pigmentation on pathological examination [2]. The most common forms of adrenal hyperplasia are: primary bilateral macronodular adrenal hyperplasia — PBMAH (also known as massive macronodular adrenal hyperplasia, MMAD) with nodules usually > 1 cm, PPNAD, and isolated micronodular adrenocortical disease (iMAD) with nodules < 1 cm. PPNAD may be isolated or may occur as part of Carney complex (CNC)



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with co-existent heart and skin myxomas, lentiginos of skin and mucosa, breast fibroadenomas, and other endocrine gland tumours [3]. Mutations of the *PRKARIA* gene coding for the regulatory subunit type 1A (R1A) of the protein kinase A (PKA) enzyme are responsible for the disease in most affected patients [4, 5].

Material and methods

In the article we characterize two patients with PPNAD, including the genetic testing.

Results

Case report 1

A 15-year-old Caucasian female patient was referred to the Endocrinology Department CMHI due to slow growth rate and excessive weight gain for about three years. She was the third child of healthy parents and was born full-term after an uneventful pregnancy. During early childhood there were no health problems, growth and development proceeded normally. At the age of eight years she had excision of a myxoma of the external auditory canal and several years later excision of a perineal myxoma.

On admission, typical features of hypercortisolism were found (central obesity, hirsutism, bruises, and petechiae), as well as scars after surgeries and melanosis of the anterior segment of the eye. Pubertal development was at Tanner stages: pubic hair 4, breast 3, no menarche [6].

Laboratory tests showed (Table I and II): normal morning cortisol level, loss of a circadian cortisol rhythm, slight increase of plasma cortisol after low-dose (0.5 mg every six hours for two days) and high-dose (2 mg every six hours for two days) dexamethasone suppression test (LDDST, HDDST-Liddle's test), and low plasma ACTH level (4.73 pg/ml; N: 10–60). The urinary steroid profile showed increased excretion of free cortisol and cortisol metabolites and no inhibition after both LDDST and HDDST, there was a slight increase of cortisol metabolites after HDDST. In the abdominal USG tumour of the right adrenal gland was suspected, and the left adrenal gland was enlarged. Repeated ultrasound examination and CT showed no abnormalities of adrenals. The echocardiogram was normal. Densitometry showed low bone mineral density (BMD) (total Z-score: -2,3 SD; L2-L4: -3,4 SD) [7, 8]. ACTH-independent CS was diagnosed with suspicion of CNC. The patient underwent right-sided laparoscopic adrenalectomy. Histopathological examination revealed typical PPNAD features (Fig. 1). Perioperative hydrocortisone (HC) therapy was performed, and HC was discontinued. Despite normal cortisol level, symptoms of adrenal

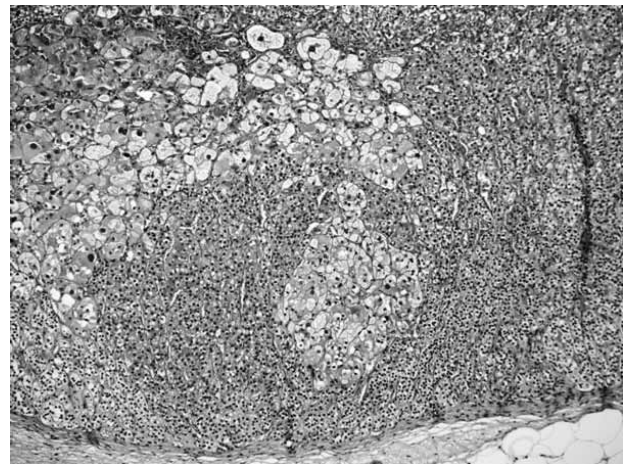


Figure 1. PPNAD, adrenal gland (patient 1). Original magnification 200×, haematoxylin and eosin staining

Rycina 1. PPNAD, nadnercze (pacjent 1). Oryginalne powiększenie 200×, barwienie hematoksyliną i eozyną

insufficiency were observed (vomiting, hypotension) subsequently. The patient remained on substitutional HC therapy. Clinical improvement and gradual withdrawal of hypercortisolism symptoms were observed for six months, then she again developed symptoms of CS: weight gain (4 kg) and hirsutism. Laboratory tests revealed loss of a circadian cortisol rhythm (Table I). Left-sided adrenalectomy was performed 10 months after right-sided surgery, and histopathological examination revealed also PPNAD features. The patient has been on HC and fludrocortisone replacement therapy since then. When she was 23 years old, genetic testing was performed, and a mutation (c.125dupG) in the *PRKARIA* gene was found.

CNC was diagnosed on the basis of three main criteria (PPNAD, myxomas, melanosis of the anterior segment of the eye).

Case report 2

A 15-year-old male patient has been observed in the Endocrinology Outpatient Department since the age of 13 years due to slow growth rate and excessive weight gain. He was born preterm (36 week of pregnancy) as the fourth child of healthy parents. His psychomotor development was delayed (sitting — nine months, walking — two years). Since his second year of life deteriorating visual acuity has been observed. When he was five years old, atrophy of both optic nerves of unknown aetiology was diagnosed. On admission to the Endocrinology Department CMHI, red striae on his arms and lower abdomen and acne on his face were found. Pubertal development was at the following Tanner stages: genitals 3 and pubic hair 3 [6]. The testicular volume was 6 ml/8 ml. Laboratory tests showed (Table I and II): loss

Table I. Cortisol circadian rhythm at the diagnosis and during re-manifestation of the disease after one-side adrenalectomy [$\mu\text{g/dl}$]**Tabela I. Dobowy rytm kortyzolu w momencie diagnozy i podczas nawrotu choroby po jednostronnej adrenalectomii [$\mu\text{g/dl}$]**

	8.00 a.m.	08.30 a.m.	00.00 a.m.	00:30 a.m.
At the diagnosis				
Patient 1	15.2	16.2	13.7	12.6
Patient 2	23.7	22.6	27.3	22.8
During re-manifestation				
Patient 1	8.6	7.3	7.3	8.0
Patient 2	12.5	13.8	9.0	14.5

Table II. Basic hormonal investigations at the diagnosis and hormonal values after low- and high-dose dexamethasone (Dx) test (Liddle's test)**Tabela II. Podstawowe badania hormonalne w momencie diagnozy i wartości hormonów po teście z małą i dużą dawką deksametazonu (Dx) (test Liddle'a)**

	8.00 a.m.	Reference	After 0.5 mg of Dx every 6 hours for 2 days	After 2 mg of Dx every 6 hours for 2 days
Patient 1				
Cortisol [$\mu\text{g/dl}$]	15.2	5–20	15.4	15.9
ACTH [pg/ml]	4.73	10–60	2.0	< 2.0
Androstenedione [ng/dl]	409	50–370	409	512
DHEAS [ng/ml]	716	350–4300	725	917
Testosterone [pg/ml]	869	88.4–964	869	618
Patient 2				
Cortisol [$\mu\text{g/dl}$]	20.7	5–25	28.2	26.2
ACTH [pg/ml]	< 10	10–50	< 10.0	< 10.0
Androstenedione [ng/dl]	338	60–380	362	452
DHEAS [ng/ml]	892	800–5600	970	—
Testosterone [pg/ml]	443	88.4–964	—	—

ACTH — adrenocorticotrophic hormone; DHEA — dehydroepiandrosterone

of a circadian cortisol rhythm, low plasma ACTH level (< 10 pg/ml), increase of plasma cortisol after LDDST and HDDST, increase of 17-hydroxycorticosteroids in urine after LDDST (8%) and HDDST (36%). CNC was suspected on the basis of clinical picture and suspicion of heart myxoma in the echocardiogram. CRH stimulation test showed no increase of ACTH and cortisol after stimulation. Echocardiography showed no abnormalities, MRI showed adrenal glands with uneven margins, without thickening. On the basis of densitometry low BMD was diagnosed (total Z-score: -2.7 SD; L2-L4: -4.3 SD). Right-sided laparoscopic adrenalectomy has been performed (with no need of subsequent HC substitutional therapy). After five months acceleration of growth rate (6 cm/year), weight loss (6 kg), and withdrawal of stretch marks and acne were observed. During the following two months weight gain (4.6 kg) and pale-pink stretch marks appeared again. Laboratory: loss of a circadian cortisol rhythm (Table I). Steroid profile in

urine revealed slightly increased cortisol metabolites with marked deficiency of 5α -reductase and increased ratio of free cortisol to cortisone. Androgens plasma levels were normal, ACTH plasma level was below the normal range. The patient underwent a left-sided adrenalectomy eight months after removal of the right adrenal gland. The diagnosis of PPNAD was confirmed by histopathological examination of both adrenals (Fig. 2). The patient is on HC and fludrocortisone replacement therapy. Genetic testing revealed a mutation (c.15dupT) in the *PRKARIA* gene.

Discussion

PPNAD is a histologically benign form of BAH [9], which causes CS that can be clinically evident or sub-clinical, constant, or cyclic [10]. It is the most frequent presentation of CNC in teens and young adults [11, 12], with the peak of occurrence in the third decade of life

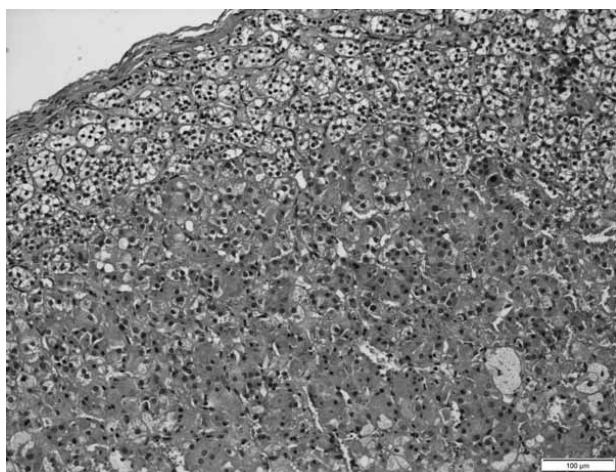


Figure 2. PPNAD, adrenal gland (patient 2). Original magnification 200×, haematoxylin and eosin staining

Rycina 2. PPNAD, nadnercze (pacjent 2). Oryginalne powiększenie 200×, barwienie hematoksyliną i eozyną

[13]. Clinical signs of CS are quite similar to those seen in patients presenting with other causes of hypercortisolism [13–15]. The most frequent symptoms of CS are: weight gain, central obesity, and slower growth rate during childhood [16]. PPNAD usually presents at a young age but not in infancy, whereas adrenal hyperplasia due to McCune-Albright syndrome usually presents in the infantile period. Other adrenocortical tumours may present before the age of five years, and they often manifest as precocious puberty [11, 16–18]. Hypogonadism is frequently observed in patients with adrenal lesions that do not secrete androgens [19].

Among 72 children with CS due to PPNAD 29% presented with short stature [13]. However, growth arrest is not a constant finding in children with PPNAD as emphasised by Groussin et al., and the classical features of CS in patients with PPNAD can be absent in childhood [14]. Güemes et al. in their analysis of 30 paediatric patients with CS showed that a decreased growth rate was observed in 36.6% of patients [10]. Growth failure was common in pituitary CD patients but not in ACTH-independent CS. The authors explain that while pure cortisol-secreting pituitary CD would inhibit growth (cortisol suppresses GH production [21], combined with a possible pituitary mass effect leading to GH deficiency), adrenal tumours causing CS may cosecrete androgens, and hence stature would be less affected.

According to the case reports from literature, the mean time between diagnosis of endogenous CS and detection of the first component of the disease is about 2–3 years (range between 0.2 to as much as 9 years) [11, 22–24]. The symptoms and signs of CS in PPNAD are insidious, the clinical manifestation may be subtle, and

most patients come to medical attention several years after their onset [25–27]. In the presented patients this time was 3 and 3.5 years. In the case of late diagnosis of PPNAD, osteoporosis is common [14]; in both analysed patients low BMD was diagnosed.

In most patients with PPNAD a paradoxical increase of cortisol level after LDDST or HDDST is observed; an increase in urinary free cortisol excretion on the second day of HDDST > 50% of the basal level usually supports the diagnosis of PPNAD [1]. It is established that dexamethasone paradoxically stimulates cortisol release through a glucocorticoid receptor-mediated effect on PKA catalytic subunits [28]. There was a 36% increase of serum cortisol after LDDST and 27% after HDDST in patient 2, and a non-significant increase, instead of a decrease, was observed in patient 1.

Adrenal glands are most commonly normal or even small in size in adrenal imaging [29, 30]. Indeed, in our patients, the adrenal glands were not enlarged in CT/MR imaging.

Treatment recommendations for PPNAD vary. Many authors suggest bilateral adrenalectomy [3, 12, 31, 32]; others think that in some cases clinical and biochemical improvement can be achieved by unilateral adrenalectomy, and excision of the second adrenal gland should be considered when hypercortisolism returns [3, 26, 33]. However, the Endocrine Society clinical practice guidelines from 2015 recommend one-stage surgical resection of bilateral adrenal disorders [34].

Some rare cases have been treated with mitotane or ketoconazole. Campo et al. [35] presented 10-year observation of a 27-year-old female who did not agree to the surgical method — she was treated with mitotane with sustained regression of CS features without inducing adrenal insufficiency.

In rare patients, in whom clinical CS did not recur after unilateral adrenalectomy, loss of circadian cortisol rhythm can be observed in long-term follow-up, showing that despite apparent clinical cure the disease is indeed bilateral [26].

The reason for bilateral adrenalectomy is also the fact that in the case of unilateral surgery preserved adrenal tissue acts autonomously, which increases the risk of adrenal failure [36–39]. In our patients the need for contralateral adrenalectomy occurred in 6–7 months of follow-up, which supports the opinion of the need of bilateral adrenalectomy.

Carney complex

CNC was first reported by JA Carney in 1985 [40]. CNC is a multiple neoplasia syndrome featuring endocrine, cardiac, cutaneous, and neural tumours [41]. In patients with CNC tumours of two or more endocrine glands are reported, including PPNAD, somatotropinoma and

Table III. Diagnostic criteria for Carney complex [41]

Tabela III. Kryteria diagnostyczne dla zespołu Carneya [41]

Clinical criteria	
1. Spotty skin pigmentation with a typical distribution (lips, conjunctiva and inner or outer canthi, vaginal and penile mucosa)	
2. Myxoma (cutaneous and mucosal) ^a	
3. Cardiac myxoma ^a	
4. Breast myxomatosis ^a or fat-suppressed MRI findings suggestive of this diagnosis ^b	
5. PPNAD ^a or paradoxical positive response of urinary glucocorticosteroids to dexamethasone administration during Liddle's test ^c	
6. Acromegaly due to GH-producing adenoma ^a	
7. LCCSCT ^a or characteristic calcification on testicular ultrasonography	
8. Thyroid carcinoma ^a or multiple, hypoechoic nodules on thyroid ultrasonography, in a young patient	
9. Psammomatous melanotic schwannoma ^a	
10. Blue nevus, epithelioid blue nevus (multiple) ^a	
11. Breast ductal adenoma (multiple) ^a	
12. Osteochondromyxoma ^a	
Additional criteria	
1. Affected first-degree relative	
2. Inactivating mutation of the <i>PRKAR1A</i> gene	

^aWith histological confirmation; ^bBased on Courcoutsakis NA et al. 1997 [52]; ^cBased on Stratakis CA et al. 1999 [42]

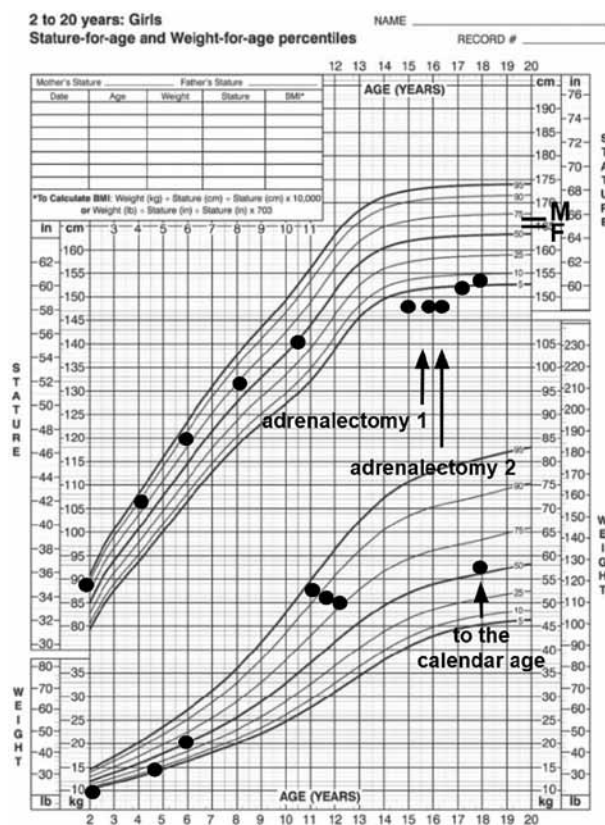


Figure 3. Growth and weight chart of patient 1
Rycina 3. Siatka centylowa wzrostu i masy ciała pacjenta 1

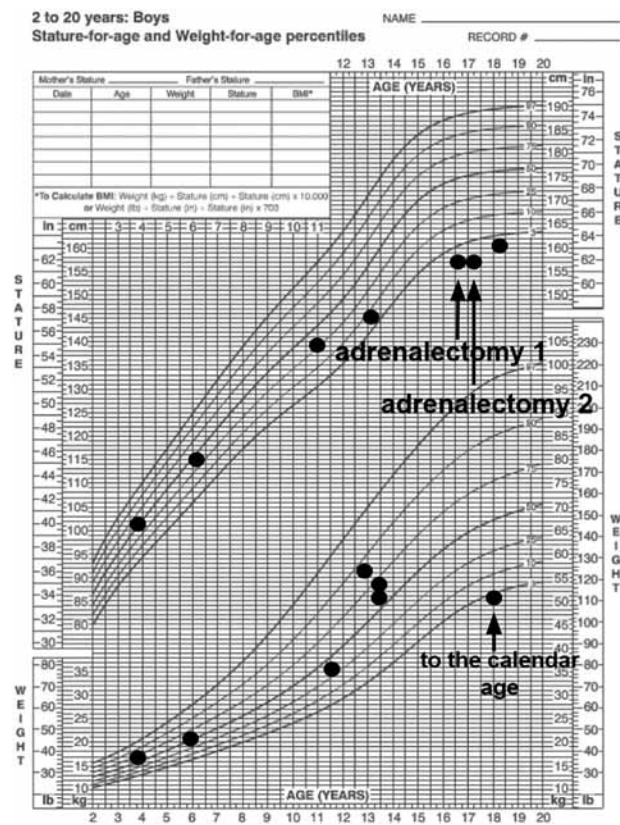


Figure 4. Growth and weight chart of patient 2
Rycina 4. Siatka centylowa wzrostu i masy ciała pacjenta 2

prolactinoma, testicular neoplasms, thyroid adenoma or carcinoma, and ovarian cysts [12, 42–48]. The diagnosis of CNC is based on clinical criteria (Table III) — it requires two main criteria or one main and one additional criterion.

Genetics of Carney complex

Two distinct loci for CNC are described: one on chromosome 17q22–24 (CNC1) and the other on chromosome 2p16 (CNC2) [4, 49]. Inactivating mutations of the *PRKAR1A* gene responsible for CNC1 are more fre-

quent [4]. Loss of R1 α function leads to increased cAMP stimulated total kinase activity [5]. The gene responsible for CNC2 is unknown [15]. At this point, there are no clear phenotypic differences between families mapping to one or the other locus.

To our knowledge, it is the first report of such types (c.125dupG and c.15dupT) of detected variants in the *PRKAR1A* gene. Both c.125dupG and c.15dupT variants are predicted to cause loss of normal protein function either through protein truncation or nonsense mediated mRNA decay. Based on currently available evidence, c.125dupG and c.15dupT are considered to be candidates for pathogenic *PRKAR1A* gene variants [4, 50, 51]. However, the possibility they are rare benign variants cannot be excluded.

Conclusions

Early diagnosis of endogenous hypercortisolism still poses a diagnostic challenge. Novel mutations presented in this article are considered to be pathogenic for PPNAD.

Acknowledgements

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