

Cushing disease in children and adolescents – assessment of the clinical course, diagnostic process, and effects of the treatment – experience from a single paediatric centre

Choroba Cushinga u dzieci i młodzieży – ocena przebiegu klinicznego, procesu diagnostycznego i efektów terapeutycznych – doświadczenia pojedynczego ośrodka pediatrycznego

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Abstract

Introduction: Adrenocorticotrophic hormone-dependent Cushing syndrome, known as Cushing disease (CD), is a rare disease in paediatric patients, the signs and symptoms of which differ from those seen in adult patients.

Aim of the study: The objective of the study was to present the diagnostics and treatment results of CD in children and adolescents.

Material and methods: We included four consequent patients in the study, 7–15 years old, with CD confirmed by laboratory tests and finally by histology examinations after surgery. The data were retrospectively retrieved from the medical records of all the patients from the years 2012–2018.

Results: The median time from the onset of clinical symptoms to CD diagnosis was 1.5 years. All patients presented with rapid weight gain accompanying growth retardation, and behavioural, puberty, or menstruation disorders. The diagnostic process was carried out in accordance with the current standard. Diagnostic difficulties regarded the cyclic CD. All patients underwent transsphenoidal complete resection of the pituitary adenoma, and they all required transient glucocorticoid substitution for a period that correlated with the time from onset of symptoms to diagnosis. All subjects achieved normalisation of hormonal results and resolution of clinical symptoms, but all are still obese. Currently the patients are in remission, but there is a suspicion of the regrowth of the tumour mass in the eldest patient.

Conclusions: In each case of the coexistence of growth velocity inhibition (not necessarily short stature) with the weight gain (not necessarily obesity) and the appearance of hyperandrogenism and depressive-anxiety disorders, biochemical diagnostics of CS should be performed. Obesity could be a chronic complication of CD in childhood, even after effective neurosurgery treatment.

Key words:

Cushing disease, CD, children, adolescents, hypercortisolaemia.

Streszczenie

Wprowadzenie: Zespół Cushinga zależny od kortykotropiny (*adrenocorticotrophic hormone* – ACTH), zwany również chorobą Cushinga (*Cushing's disease* – CD), jest rzadkim schorzeniem u dzieci i młodzieży, którego objawy różnią się od obserwowanych u osób dorosłych.

Cel pracy: Ocena procesu diagnostycznego i wyników terapii CD u dzieci i młodzieży.

Materiał i metody: Do pracy włączono 4 kolejne pacjentki w wieku 7–15 lat z CD rozpoznaną na podstawie wyników badań hormonalnych i badania histopatologicznego usuniętego gruczolaka przysadki po przeprowadzeniu retrospektywnej analizy dokumentacji medycznej pacjentów leczonych w naszym ośrodku w latach 2012–2018.

Wyniki: Mediana czasu od początku objawów klinicznych do rozpoznania CD w prezentowanej grupie pacjentów wynosiła 1,5 roku. Wszystkie pacjentki miały objawy CD typowe dla wieku rozwojowego: gwałtowny przyrost masy ciała z zahamowaniem

tempa wzrastania, zaburzenia zachowania, dojrzewania bądź miesiączkowania. Diagnostyka pacjentów była przeprowadzona zgodnie z obowiązującym standardem, najwięcej trudności diagnostycznych sprawiła postać cykliczna CD rozpoznana u jednej z pacjentek. Wszystkie pacjentki przebyły przezklinową całkowitą resekcję gruczolaka przysadki. Po zabiegu wszystkie wymagały przejściowej substytucji glikokortykoidów w czasie, który wyraźnie korelował z długością czasu od wystąpienia objawów do ustalenia rozpoznania. U wszystkich uzyskano normalizację wyników hormonalnych i ustąpienie objawów klinicznych po leczeniu neurochirurgicznym, jednakże wszystkie pacjentki nadal są otyłe. W badaniach MR przysadki u 3 pacjentek nie stwierdza się odrostu masy guza, u najstarszej jest jego podejrzenie.

Wnioski: W przypadku współwystępowania zahamowania procesu wzrastania (niekoniecznie niskorosłości) z przyrostem masy ciała (niekoniecznie otyłości), cech androgenizacji i zaburzeń zachowania konieczne jest przeprowadzenie diagnostyki w kierunku CD. Otyłość może być stałym powikłaniem CD rozpoznanej w dzieciństwie mimo jej skutecznego leczenia neurochirurgicznego.

Słowa kluczowe:

choroba Cushinga, dzieci, młodzież, hiperkortyzolemia.

Introduction

After the age of five years, adrenocorticotrophic hormone (ACTH)-dependent Cushing syndrome (CS) is more common than ACTH-independent CS. In most cases ACTH is produced by the pituitary microadenoma and causes hypercortisolaemia resulting in obesity with simultaneous growth velocity inhibition, behavioural changes, and hyperandrogenism (precocious puberty, hirsutism, lack or regression of puberty due to the suppression of gonadal axis by adrenal androgens) [1, 2]. The overproduction of ACTH by the pituitary adenoma occurs in a semi-autonomous manner and retains some autoregulatory properties of the hypothalamic-pituitary-adrenal axis in the negative feedback cycle. There are no symptoms of excess aldosterone [3]. In each case of the coexistence of the above features CS should be suspected and detailed, proper biochemical diagnostics should be performed. After the diagnosis of CS, the main goal of the treatment is a rapid normalisation of serum cortisol levels to prevent the adverse effects of prolonged hypercortisolaemia. In children, this is particularly important to prevent disorders of their somatic development [4].

The aim of the study was to present the diagnostics and treatment results of CD in children and adolescents, underling the differences in the clinical outcome of CD between paediatric patients and adults.

Material and methods

Patients

Four consequent patients, 7–15 years-old, with CD confirmed by laboratory tests and finally by histology examinations of removed pituitary adenoma, treated in our department in the years 2012–2018, were included in the study. All patients underwent transsphenoidal complete resection of the pituitary adenoma, performed by the same experienced neurosurgeon. The data were retrospectively retrieved from medical records of all the patients. The parents of the patients (and the patients above 16 years old) gave their written, informed consent for the presentation of the medical data in the journal article.

Biochemical diagnostics

The diagnosis of CD was made based on standard hormonal criteria: increased urinary-free cortisol (UFC) in three 24-h urine collections, disturbances of cortisol circadian rhythm, increased or not suppressed plasma ACTH levels at 8.00 a.m. accompanying increased serum cortisol levels, and a failure to suppress serum cortisol levels to $< 1.8 \mu\text{g/dl}$ during an overnight dexamethasone suppression test (ODST; 1 mg of dexamethasone was given at 11.00 p.m.). The pituitary aetiology of CS was confirmed based on serum cortisol or UFC suppression $< 50\%$ with a high-dose dexamethasone suppression test (HDDST; 2 mg q.i.d. for 48 h) or a positive result of a corticotrophin-releasing hormone (CRH) stimulation test ($1 \mu\text{g/kg}$, maximum $100 \mu\text{g IV}$) and positive pituitary MRI.

Plasma ACTH levels were measured using radioimmunochemistry (Brahms, Germany). Serum cortisol concentrations were determined by indirect chemiluminescence using an Advia Centaur (Siemens Healthcare Diagnostics, USA). Serum levels of DHEA-S (Diasource, Belgium), PRA (DiaSorin, Italy before 2014, currently Immunotech, Czech Republic), and aldosterone (ZenTech, Belgium before 2014, currently Cisbio, France) were measured using radioimmunochemistry. Cortisol in urine was measured using isotope dilution mass spectrometry (Orion Diagnostica, Finland before 2013, in the years 2013–2017 Diasource, Belgium, currently Immunotech, Czech Republic).

Treatment and postoperative outcome

All patients underwent MRI of the pituitary (GE Signa, 1.5 Tesla) before and after IV injection of gadolinium (Gd-DTPA). A macroadenoma was defined as a tumour with at least one diameter $> 10 \text{ mm}$, and a microadenoma as a pathological mass with maximum diameter $\leq 10 \text{ mm}$.

Histopathological examinations were performed in accordance with the 2004 WHO classification. Tissue samples were routinely immune-stained for all pituitary hormones. Ki-67 (MIB1) labelling index (LI) was graded in two categories: a) < 3 and b) ≥ 3 . Corticotroph adenomas were classified on the basis of commonly accepted ultrastructural features, such as densely granulated (DG-ACTH) or sparsely granulated (SG-ACTH).

Blood samples for serum cortisol were collected from all patients at 6.00 a.m. on the first, second, and third postoperative days. Hydrocortisone replacement therapy was started after biochemical confirmation of hypocortisolaemia or when clinical manifestations of adrenal insufficiency were observed. An early, postoperative remission was recognised, when the nadir serum cortisol level taken at 6.00 a.m. on the first, second, and third postoperative day was $< 1.8 \mu\text{g/dl}$. Late biochemical remission was recognised on the basis of hormonal assessments 6 months after pituitary surgery.

Statistical analysis

Statistical analysis of the clinical data was made using Excel (Microsoft Office Professional Plus 2016, PL).

Results

Patients' clinical signs and symptoms

All patients presented with obesity and a decrease in growth velocity at various times before CD diagnosis. Moreover, significant changes of the patients' behaviour were observed, such as depression or aggressive and auto-aggressive behaviours, weakness, less tolerance of physical exercise, and ravenous hunger. Our patients presented with hyperandrogenism such as precocious or delayed puberty (an arrest of breast development, primary amenorrhoea), secondary amenorrhoea. The youngest patient had cyclic presentation of CD symptoms (Table I).

Physical examination of patients 1–3 revealed Cushingoid face (round, moon face with plethora), axial obesity, swarthy skin, red massive striae on the abdomen, hips, and limbs, and hirsutism and severe acne located on the face and chest. In patient 1 there was also bruising of the skin, thinned skin on the lower legs, widened veins, numerous red spots on the back (candidiasis), and erythematous cutaneous eruptions. Only patient 4 did not present with striae, but Cushingoid face, axial obesity, hirsutism, acanthosis nigricans on the neck, and precocious adrenarche (pubarche II stage according Tanner scale) in the absence of development of the breast glands (steatomastia) was seen. Blood pressure was elevated in two patients (130/100 mm Hg, 145/80 mm Hg), and it was within the normal range in the other two.

The results of anthropometric measurements are presented on growth and body mass charts of all patients (Figs. 1–4). Bone age was accelerated by a median of two years in patients 1–3, and in patient 4 it was slightly accelerated (by half a year).

Results of diagnostic process

Hypercortisolaemia was confirmed by a high level of UFC in three daily urine collections in all patients as well as by an abnormal daily rhythm of cortisol secretion with lack of nocturnal decline at midnight (Table II). All patients had elevated or non-suppressed ACTH level. There was no suppression of cortisol secretion in ODST in patients 1–3. In patient 2 an in-

hibition of cortisol in the test with a high dose of dexamethasone (8 mg) was observed (cortisol $1.2 \mu\text{g/dl}$). The CRH test performed in patients 2–4 showed an increase of over 35% in ACTH and in cortisol by more than 20%. Plasma renin and aldosterone activity were normal in all patients. Adrenal androgens (DHEA-S) were elevated only in patient 2 (Table II).

It is worth underling that in the youngest patient (patient 4), studies carried out at the time of her first visit in our outpatient clinic revealed the following: one-time determination of cortisol in the diurnal collection of urine, morning ACTH, cortisol and 17-OHP values were normal. MRI of the hypothalamic-pituitary area performed at the age of eight years showed no deviations. Due to the clearly cyclical course of the disease, it was decided to continue observation of the patient. After half a year, the patient was referred for re-evaluation of the hypothalamic-pituitary-adrenal axis and a repeat MRI of the hypothalamic-pituitary area because of repeated, rapid weight gain accompanied by nocturnal anxieties, exacerbation of depressive behaviour, insomnia, and excessive sweating.

Densitometry examination of the lumbar spine showed a decrease in bone mineralisation in patients 1–3 and no abnormalities in patient 4. Z-score values were as follows: -2.7 (patient 1), from -1.8 to -2.3 (patient 2), and -2.3 (patient 3).

Finally, MRI of the hypothalamic-pituitary region revealed adenoma in all patients; however, in two of them it was not visible in first MRI but was visible in the neuroimaging repeated six months after the previous examination (Table I).

Results of treatment

All patients underwent total transsphenoidal resection of the pituitary adenoma. Patient 1 and patient 4 were prepared for the neurosurgery with ketoconazole. Histopathological examination confirmed densely granulated pituitary corticotrophic adenoma with varied Ki67 expression. Patient 1 had a double pituitary adenoma composed of densely granulated corticotrophic and densely granulated somatotrophic cells, although in prior neurosurgery she had presented with no clinical features of excess growth hormone and her IGF-1 level was normal (366.5 ng/ml, normal range 214–753 ng/ml).

All patients met the criteria of early remission of CD, but they required supplementation for different times after surgery, for 0.5 – 40 months, median 4.5 months (Table I). Puberty was undergone normally, regular menstruation cycles reappeared, and normalisation of BMD was observed. In spite of the first patient, who lost the final target height due to an L2 vertebra compression fracture, the height velocity improved after surgery. However, all patients still present obesity or overweight (Figs. 1–4). Biochemical diagnostics has confirmed late remission of all patients.

Pituitary MRI examination repeated regularly after surgery has not showed signs of regrowth of the tumour mass in three patients. In the oldest one, there is a suspicion of regrowth of the tumour, although currently neither clinical nor biochemical signs have confirmed this. We plan to repeat the diagnostic process.

Table 1. Signs and symptoms before diagnosis, the treatment, and clinical outcome of Cushing disease in children and adolescents

Patient	Age at CD diagnosis [years]	Signs and symptoms	Time before diagnosis [years]	MRI of pituitary gland	Treatment	Result of histopathological examination	Steroid discontinuation after neurosurgery [months]	Observation period after neurosurgery [months]	Clinical status at present
♀ 1	13 1/12	Behavioural disorders, obesity + simultaneous growth retardation androgenisation (hirsutism, delayed puberty) typical skin lesions + massive cutaneous candidiasis Cushingoid face, hypertension, decrease of bone mineral density (BMD) Z-score -2.7	5	Macro-adenoma diameter 20 mm	Total trans-sphenoidal resection of macro-adenoma	Double pituitary adenoma – contagious type – composed of corticotroph cells (densely granulated – DG-ACTH) and somatotroph cells (densely granulated – DG-GH) Level of monoclonal antibodies against the Ki67 protein (MIB1 antibody) < 3%	40	78	Menarche 14 8/12 regular menstruation cycles, obesity, and behavioural disorders associated with the diagnosis of autism without clinical and biochemical signs and symptoms of CD relapse BMD -0.8 In MRI suspicion of tumour regrowth
♀ 2	15 7/12	Obesity + simultaneous growth retardation androgenisation (hirsutism, secondary amenorrhoea) typical skin lesions Cushingoid face, hypertension, decrease of BMD Z-score -1.8	1	Micro-adenoma	Total trans-sphenoidal resection of micro-adenoma	Atypical, DG-ACTH adenoma MIB1 > 10%	0.5	25	Recurrence of menstrual cycles 5 months after neurosurgery, currently regular menstrual cycles, obesity/overweight without clinical and biochemical signs and symptoms of CD relapse In MRI without tumour regrowth

♀ 3	12 4/12	Obesity + simultaneous growth retardation, behavioural disorders, striae only on breasts, Cushingoid face, decrease of BMD Z-score -2.3	1.5	diameter 5 mm	Total trans-sphenoidal resection of micro-adenoma after bilateral inferior petrosal sinus sampling	DG-ACTH adenoma MIB1 < 1%	5	22	Menarche at the age of 13 2/12 years, regular menstrual cycles Overweight/obesity Without clinical and biochemical hypercortisolaemia In MRI without recurrence
♀ 4	7 8/12	Cyclic episodes of obesity + simultaneous growth retardation + behavioural disorders + androgenisation + acanthosis nigricans + Cushingoid face Z-score -1.3	1.5	Micro-adenoma diameter 4 mm	Total trans-sphenoidal resection of micro-adenoma	Atypical, DG-ACTH adenoma MIB1 > 3%	4	12	Improved growth rate (10 th percentile) overweight/obesity, without clinical and biochemical hypercortisolaemia In MRI without recurrence
Median	12 8/12	Obesity + simultaneous growth retardation behavioural disorders, disturbances of menstruations hypertension Cushingoid face	1.5	Micro-adenoma diameter 5 mm	Total trans-sphenoidal resection of micro-adenoma	Pituitary corticotroph adenoma in two cases (50%) MIB1 > 3%	4.5	16.5	Obesity without clinical and biochemical signs and symptoms of CD relapse in MRI without tumour regrowth

A

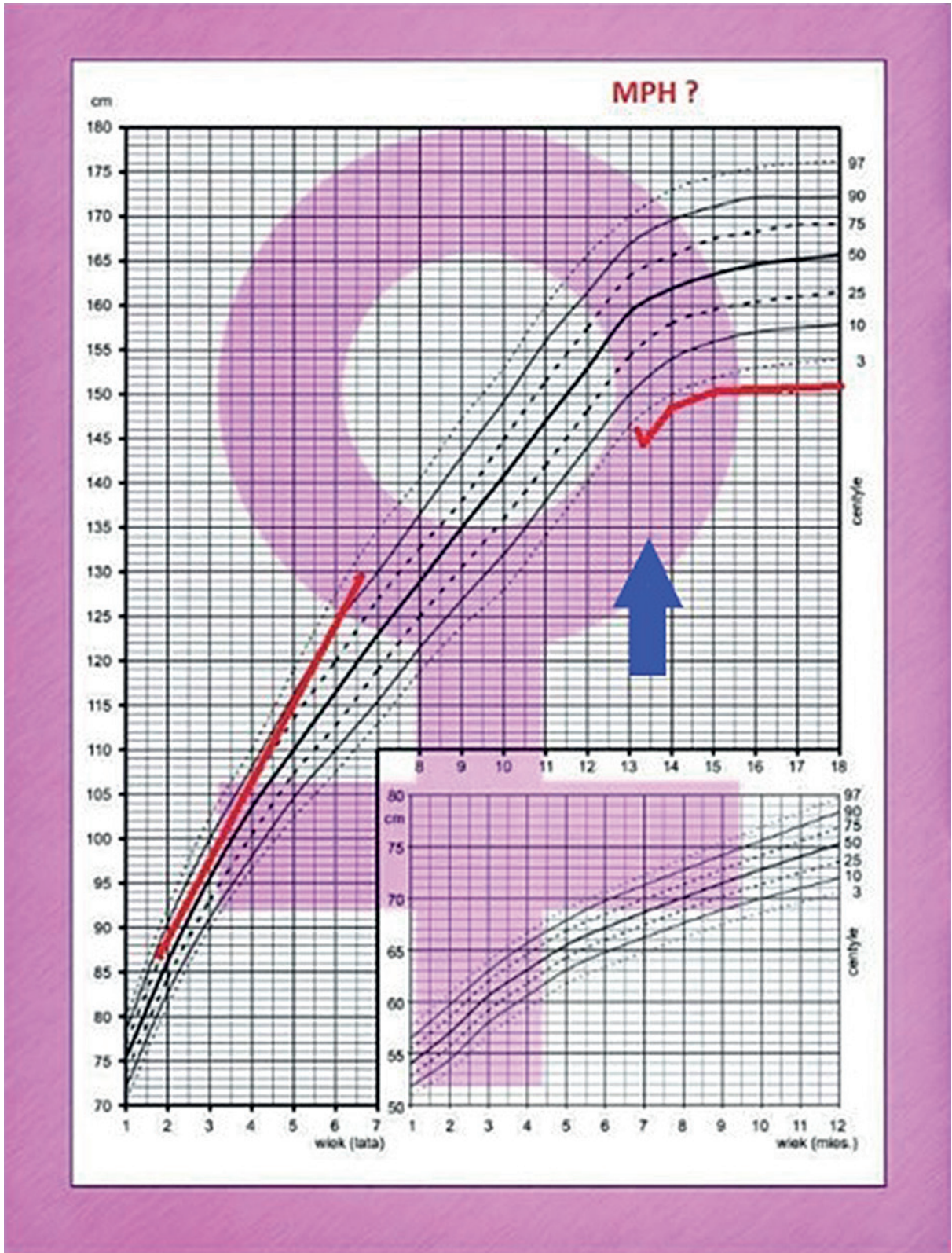


Figure 1. Growth chart (A) and body mass chart (B) of patient 1. Blue arrow indicates the time of neurosurgery

B

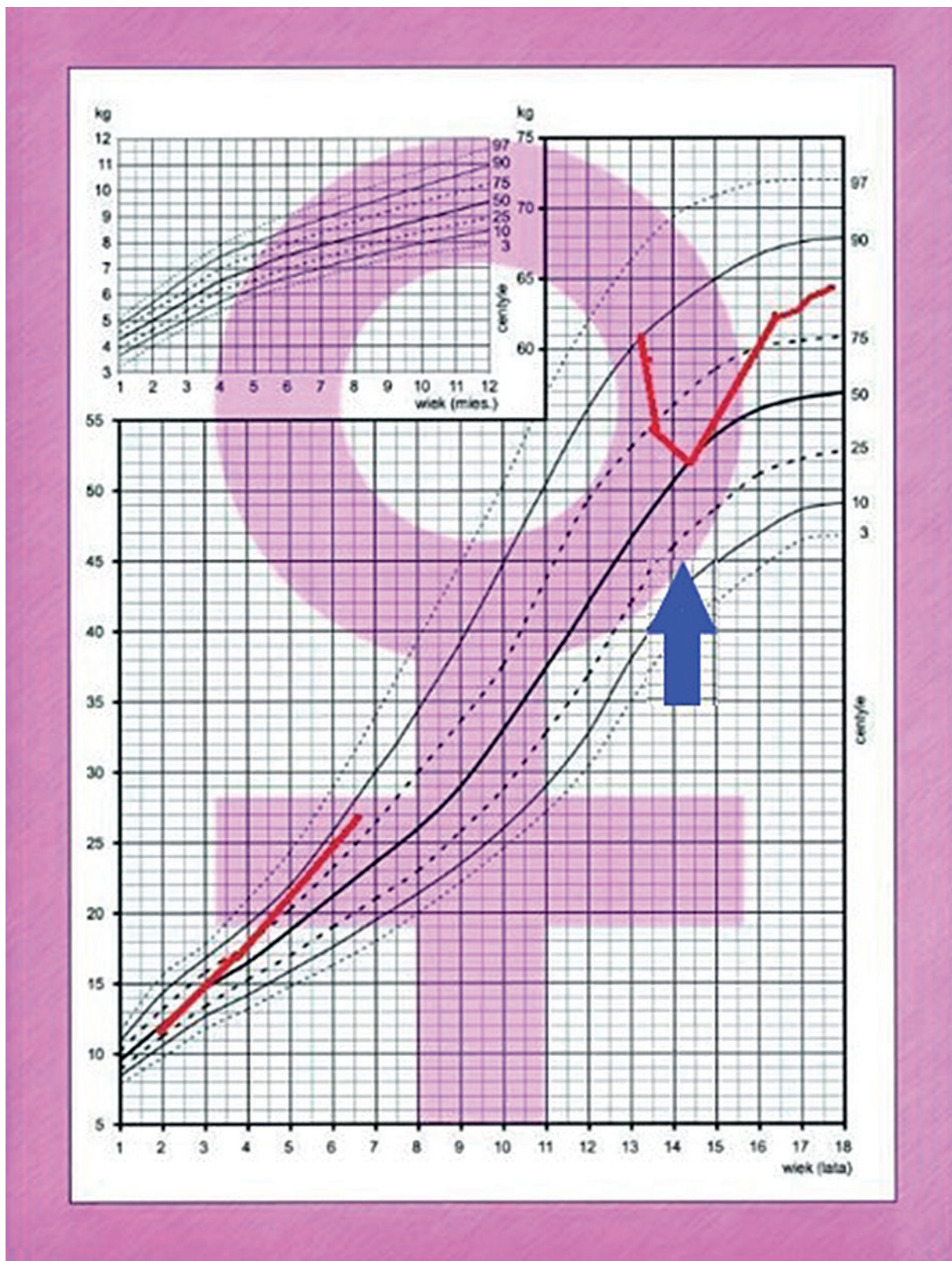


Figure 1. Growth chart (A) and body mass chart (B) of patient 1. Blue arrow indicates the time of neurosurgery

A

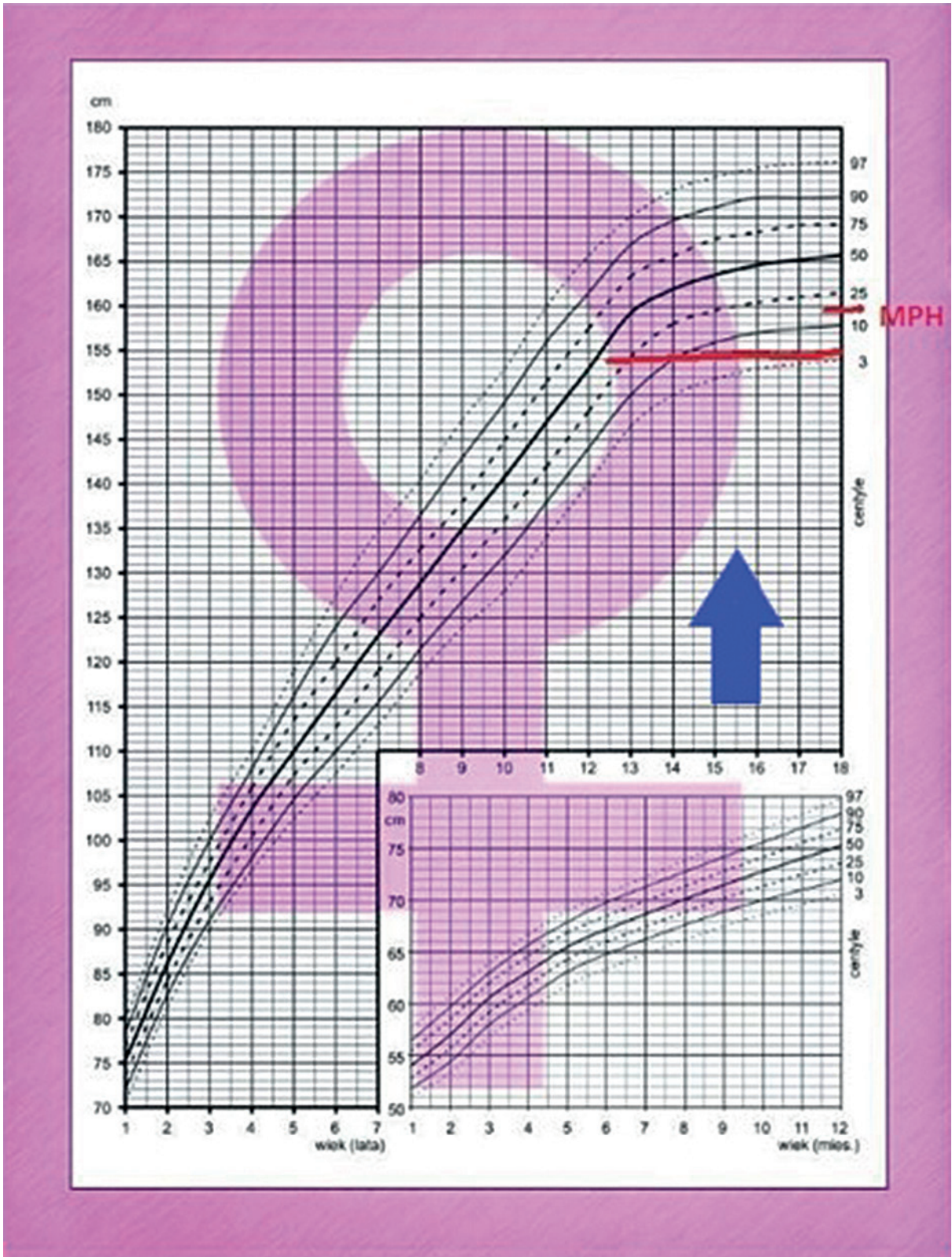


Figure 2. Growth chart (A) and body mass chart (B) of patient 2. Blue arrow indicates the time of neurosurgery

B

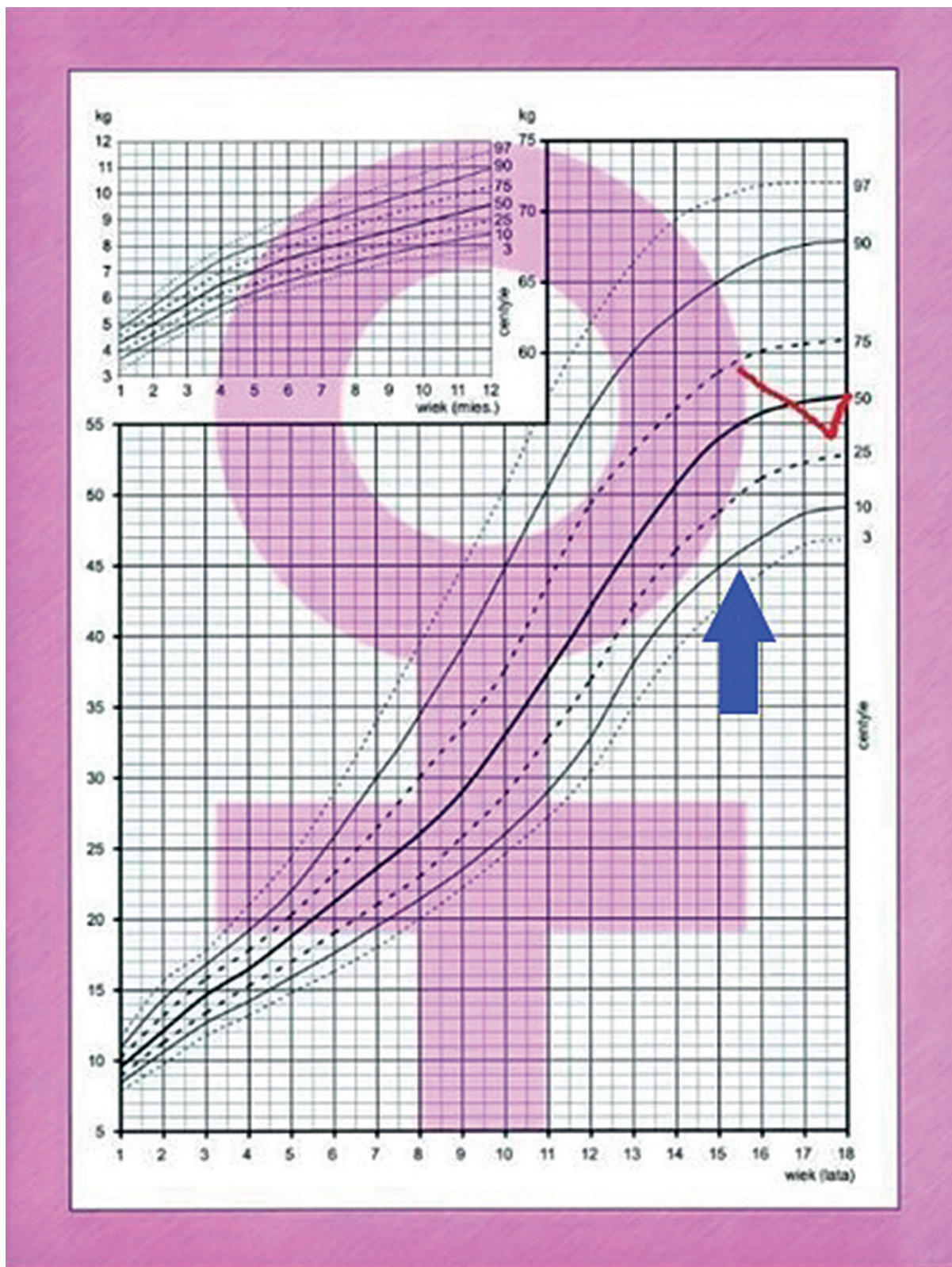


Figure 2. Growth chart (A) and body mass chart (B) of patient 2. Blue arrow indicates the time of neurosurgery

A

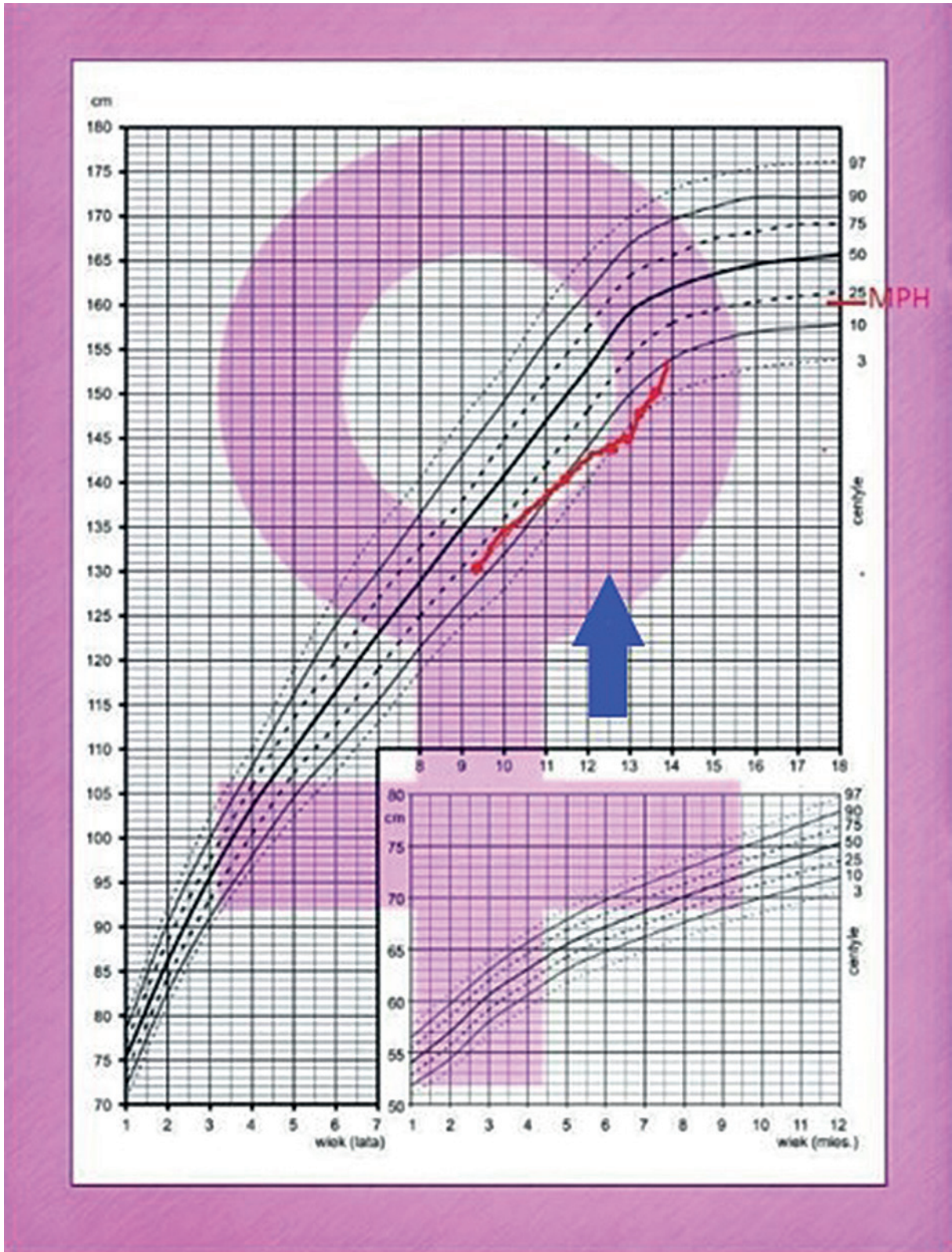


Figure 3. Growth chart (A) and body mass chart (B) of patient 3. Blue arrow indicates the time of neurosurgery

B

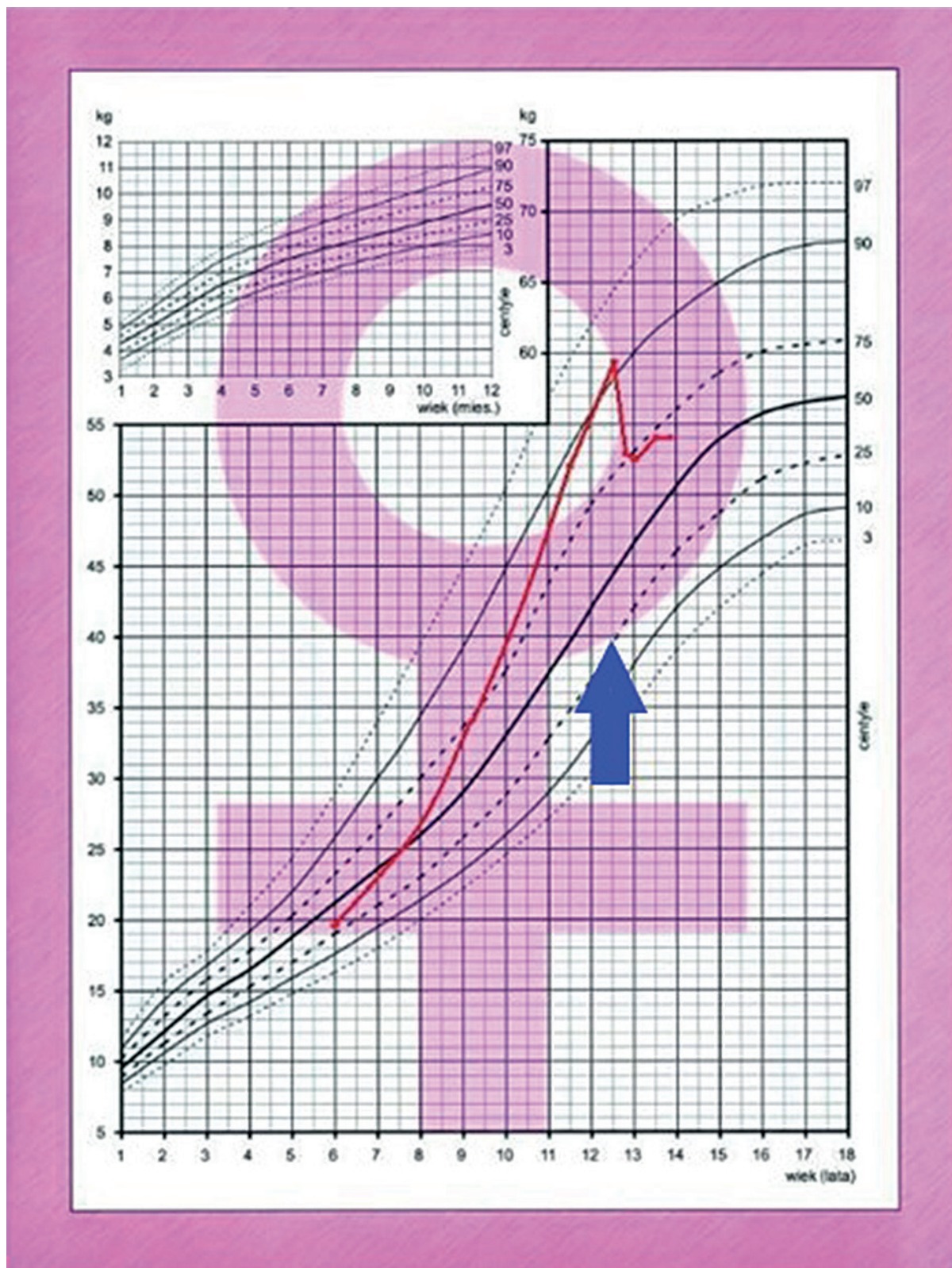


Figure 3. Growth chart (A) and body mass chart (B) of patient 3. Blue arrow indicates the time of neurosurgery

A

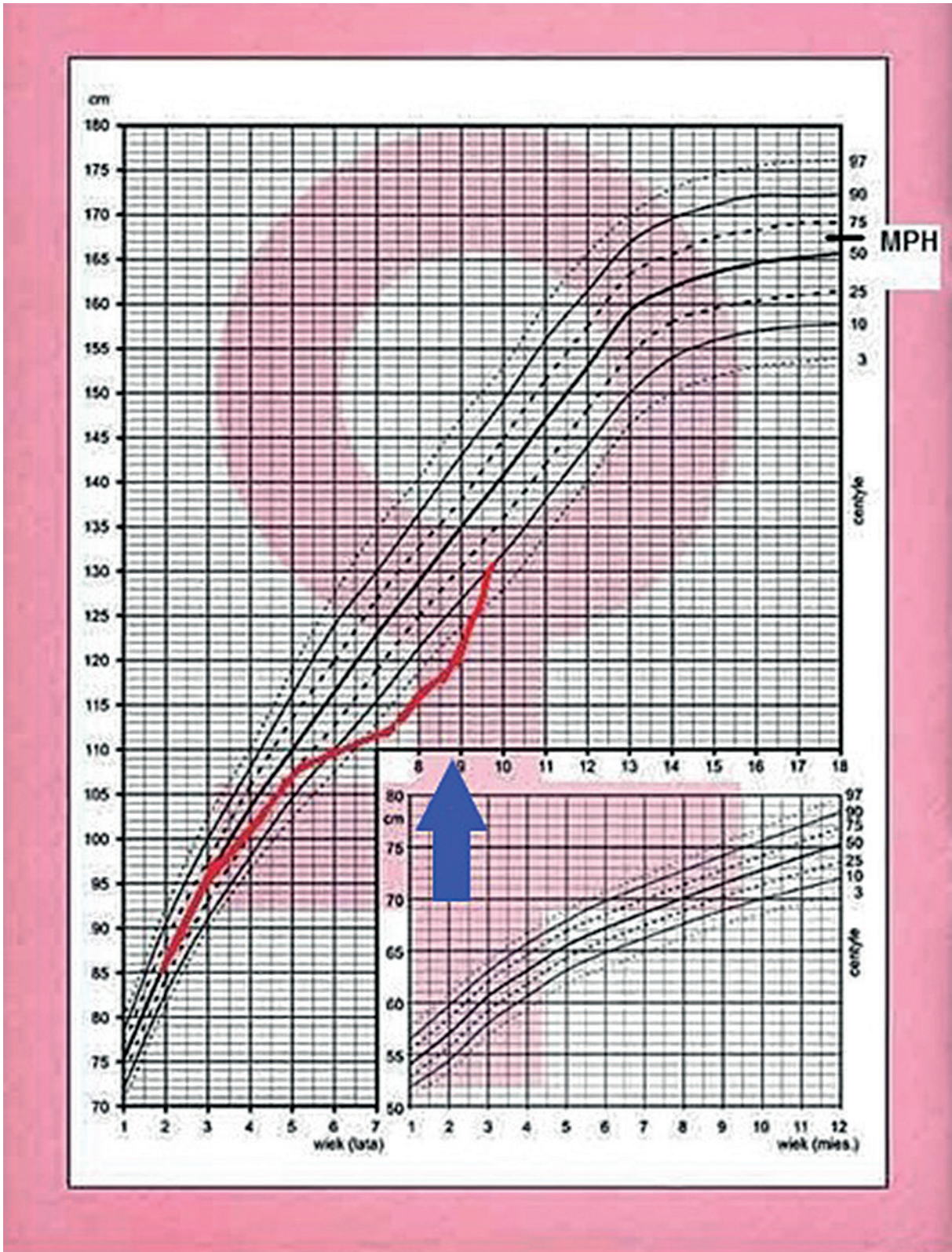


Figure 4. Growth chart (A) and body mass chart (B) of patient 4. Blue arrow indicates the time of neurosurgery

B

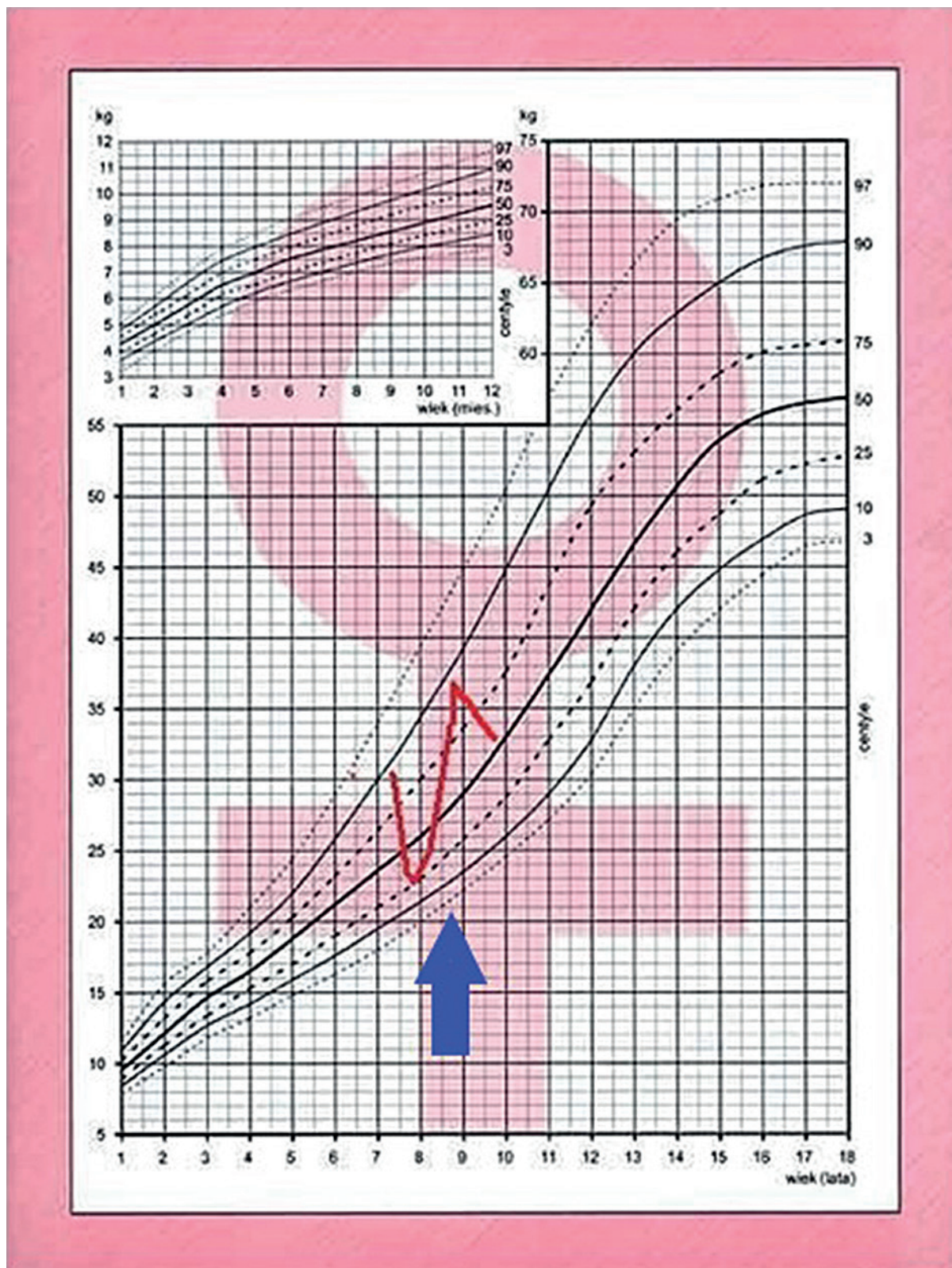


Figure 4. Growth chart (A) and body mass chart (B) of patient 4. Blue arrow indicates the time of neurosurgery

Table II. Details of biochemical diagnostics of patients with Cushing disease

Patient	Results of 24-h urine collection for cortisol [$\mu\text{g}/\text{day}$]	Diurnal rhythm of cortisol [$\mu\text{g}/\text{dl}$]	Results of ODST [$\mu\text{g}/\text{dl}$]	Results of CRH test	Levels of ACTH [pg/ml]
No 1	1) 164.8 2) 560.2 3) 273.5	8 a.m. – 20.08 4 p.m. – 14.53 8 p.m. – 16.93 midnight – 22.4	10.8	Undone	156.9
No 2	1) 557.02 2) 602.82 3) 292	8 a.m. – 18.7 4 p.m. – 8.83 8 p.m. – 8.8 11 p.m. – 14.6	2.16	Increase of 62% in ACTH and 22% in cortisol	70.4
No 3	1) 119.9 2) 124.9 3) 430	8 a.m. – 15.52 8 p.m. – 13.32 11 p.m. – 16.5	12.28	Increase of 126% in ACTH and 157% in cortisol	109.7
No 4	1) 219 2) 284 3) 283	8 a.m. – 9.5 8 p.m. – 18.5 midnight – 25.4	1.02	Increase of 46% in ACTH and 45% in cortisol	37.9

Normal ranges: 24-h urine collection for cortisol: children 2–27, adolescents 5–55 $\mu\text{g}/\text{day}$; ACTH: 10–60 pg/ml ; cortisol at 8.00 a.m. – 5–23 $\mu\text{g}/\text{dl}$, midnight < 4.4 $\mu\text{g}/\text{d}$; ODST cortisol < 1.8 $\mu\text{g}/\text{dl}$; positive result of CRH test: increase of over 35% in ACTH and in cortisol of over 20%

Discussion

Cushing disease in children older than seven years accounts for about 75% of cases. The peak incidence of CD in paediatric patients is during adolescence, with a median age according to various studies ranging from 12.3 to 14.1 years [4, 5]. CD occurs very rarely and, as in adults, there is a predominance of females over males. The median age in our presented group was 12.3 years, and they were only girls. No explanation for the differences in the incidence and clinical history of CD between the sexes was found [6]. In the majority of the paediatric population with CD, germinal mutations that could be the cause of ACTH pituitary tumour production have not been identified; however, MEN1 syndrome cannot be excluded, because CD may be the first manifestation of this condition in children [7]. In addition, some patients have been identified with a somatic mutation in the USP8 deubiquitinase gene in ACTH pituitary tumour cells that may be involved in the molecular pathogenesis of the disease [8].

In children with CD, rapid weight gain accompanied by inhibition of growth velocity is an early, sensitive, and most characteristic symptom [2]. In the present work, rapid weight gain and inhibition of growth velocity in all patients were observed. This is clearly illustrated by the presentation of patient 4 with cyclic CD, in whom during hypercortisolaemic periods the growth ve-

locity inhibition was accompanied by subsequent weight gain, and during the remission periods the patient lost weight and her growth curve ran linearly parallel to normal growth curves on the growth chart (Fig. 4). Other symptoms of CD in children are (as in adults): truncal obesity, moon face with plethora, buffalo hump, hirsutism, arterial hypertension, and behavioural and mental disorders. Skin lesions are also characteristic: purple striae, acne, hirsutism, hyperpigmentation of the skin, acanthosis nigricans, fungal infections, and easy bruising. All of our patients had bruising, acne, and hirsutism, while striae, hyperpigmentation, and acne occurred in three of them. Striae, which are skin lesions most commonly associated with CS, were not visible in the youngest of the patients. In our youngest patient features of premature puberty were also found, as described in prepubertal CD patients [4, 9]. Acceleration of bone age was observed in all our patients, and it was markedly associated with the intensification of hyperandrogenaemia. Patients 1 and 2 also had hypertension. In all patients, changes in behaviour and changes in facial features were observed. According to the latest meta-analysis of symptoms in children with CD changes in facial features, which are found in all patients, followed by rapid weight gain and virilisation symptoms, hirsutism, emotional lability, fatigue, and headache are dominating. Growth inhibition was found only in 37% of cases, but in more than 95% of cases a significant difference between

SDS growth (which was below 0) and SDS BMI was observed, which was +1.5 SD [4]. The cyclical form of the disease, manifested by regression of clinical symptoms with normalisation of hormonal tests, is observed in 5–15% of patients [3, 4, 10, 11]. Due to the heterogeneity of symptoms of CD described above, sometimes many months pass from their occurrence to diagnosis. In our work the median of this time was 1.5 years. According to the literature, this time can range from three months to seven years, on average 3 years [12].

Diagnosis of CS was based on typical clinical features and a confirmation of hypercortisolaemia in biochemical tests. The first stage of biochemical diagnostics is the confirmation of hypercortisolaemia using a screening test, which is the three-fold 24-h collection of UFC. The sensitivity of a single test is 70%, and in the case of the repeated one it reaches 88%. Complementary testing may also be the determination of urinary 17-hydroxysteroides (17 OHS) in 24-h urine collections [13]. Elevated UFC exceeding the reference values for the patient's age are the confirmation of hypercortisolaemia, although false positive results may occur in cases of chronic emotional and physical stress, long-lasting, chronic obesity, depression, poorly balanced diabetes, intense physical exercise, depression, alcoholism, anorexia, malnutrition, drug withdrawal, psychogenic polydipsia, and pregnancy. These states are referred to as pseudo-Cushing [3]. Moreover, in girls during adolescence a physiological increase in the excretion of free cortisol in the urine occurs [14].

In children, a very sensitive indicator of hypercortisolaemia is the lack of typical diurnal variability of serum cortisolaemia, especially nocturnal decline, and in particular the typical suppression of cortisol secretion at midnight [3, 13], which was visible in all patients. It is considered that cortisol levels higher than 4.4 $\mu\text{g}/\text{dl}$ clearly confirm the diagnosis of CD in children. This test has 99% sensitivity and 100% specificity. This value differs between children and adults, in whom it is 7.5 $\mu\text{g}/\text{dl}$ [15]. Single ACTH and cortisol values are not diagnostically useful due to their high variability caused not only by the daily rhythm of production of these hormones, but also by the modulating effect of endo- and exogenous factors. In the paediatric population, the assessment of the cortisol concentration in the patient's saliva during the night could be useful with high sensitivity (95.2%) and specificity (100%) for hypercortisolaemia [12]. Another standard test in the diagnosis of hypercortisolaemia is an overnight dexamethasone suppression test (ODST) (1 mg, in children 15 $\mu\text{g}/\text{kg}$, maximum 1 mg) [12, 16, 17]. This test has a 15–20% risk of false positive results, especially in very obese people. In the night test with 1 mg dexamethasone, there was no inhibition of cortisol in three patients (cases 1, 2, and 3), and one patient showed inhibition of cortisol (case 4). These clinical data confirm that the suppression test results of dexamethasone in children with Cushing syndrome should be interpreted with caution.

Therefore, a recognised standard in paediatrics is to perform a low-dose dexamethasone test administered for 48 hours at a dose of 30 $\mu\text{g}/\text{kg}/\text{d}$, maximum 0.5 mg per dose, for the determination of cortisolaemia in fraction 0, after 24

and after 48 hours. This test has a sensitivity of more than 90% in the diagnosis of CD [5]. This test can be replaced by a suppression test with a high dose of dexamethasone (in paediatric practice, 120 $\mu\text{g}/\text{kg}$ b.w. per dose, maximum 8 mg), which is given at 11.00 p.m. with the measurement of cortisol levels at 8 a.m. the same and the next day. Suppression of its concentration above the cortisol value from the previous day (above 50%) is confirmation of CD. This test has a high sensitivity of 88% but rather a low specificity of 57%, associated with false positive test results in obese people [18].

Diagnostic difficulties in the diagnosis of hypercortisolaemia appear in the case of the cyclic form of CD. If this form of CD is suspected, long-term observation of the patient is recommended, with particular attention paid to the course of the growth process. This CD form must also be differentiated from PPNAD, which runs cyclically, but is ACTH-independent CS with a marked inhibition of ACTH secretion with the paradoxical excessive cortisol response of the dexamethasone test [19]. Another problem in the diagnosis of CD is its differentiation with pseudo-Cushing syndrome. To distinguish these two forms, it is recommended that a combined test be performed, i.e. a dexamethasone suppression test with a CRH supply. It consists in administering eight doses of dexamethasone (30 $\mu\text{g}/\text{kg}/\text{d}$ maximum 0.5 mg per dose) every 6 hours (the test lasts 48 hours) and then performing the test with CRH (1 $\mu\text{g}/\text{kg}$, maximum 100 μg IV) with assessment of ACTH and cortisol at the following time points: –15 min, –5 min, 0 min, and +15 min. Patients with pseudo-Cushing syndrome will have uncertain or very low values of both parameters and no response to CRH stimulation [3]. In patients with CD higher initial values of cortisol and ACTH are observed in the CRH test, as well as a noticeable response of cortisol to CRH. A cortisol concentration higher than 1.4 $\mu\text{g}/\text{dl}$ 15 min after administration of CRH speaks for the diagnosis of CD. Another test that differentiates these two disease entities seems to be the measurement of cortisol at midnight. Patients with pseudo-Cushing syndrome have a cortisol level at midnight that is lower than 7.5 $\mu\text{g}/\text{dl}$, which differentiates them from CD patients [20]. These data concern adults. Therefore, in children, it seems that, similarly to the diagnosis of hypercortisolaemia, the value of 4.4 $\mu\text{g}/\text{dl}$ should be considered as differentiating the case of CD from pseudo-Cushing syndrome. However, this is a hypothesis of the authors of this work and is not confirmed by the results of a clinical trial. The occurrence of pseudo-Cushing syndrome is extremely rare in children, so it is difficult to have a sufficiently large cohort for research.

After documenting hypercortisolaemia, localisation tests should be performed to confirm CD, i.e. ACTH-dependent CS, which include ACTH concentration measurement, CRH test, and (in patients with suspected CD) imaging of the pituitary gland. If there is an absence of the visualisation of a focal lesion in the pituitary, bilateral inferior petrosal sinus catheterisation with ACTH evaluation before and after CRH stimulation should be performed [21].

Adrenocorticotrophic hormone levels above 29 ng/l, which were present in all patients, indicate ACTH-dependent CS

with high sensitivity (70%) and specificity (100%) according to the data from the literature [13]. The next stage of diagnosis is the CRH test at a dose of 1 µg/kg, maximum 100 µg IV, characterised by 91% sensitivity and 100% specificity. An increase in ACTH levels of over 35% and cortisol of more than 20% after administration of CRH confirms CD.

The presence of a microadenoma in the pituitary MRI examination, usually smaller than 3–4 mm, is found in about half of the cases. In two large studies including paediatric patients, pituitary adenomas were found in 63% and 55% [22]. However, in a multicentre analysis of 1284 paediatric patients who underwent neurosurgical treatment due to the pituitary adenoma, patients with CD (i.e. the presence of confirmed ACTH-secreting pituitary adenoma) are the largest group, constituting 43% of the whole [23]. In all patients, MRI examination showed a pituitary adenoma responsible for the CD; however, in two of them (50%) the microadenoma was revealed in MRI re-examination. In the standard procedure, in the absence of a pituitary lesion in the MRI examination, it is recommended that a catheterisation of the petrosal sinuses be performed. This procedure assesses ACTH levels from blood samples collected from the sinuses on both sides before and after CRH stimulation, but also ACTH from peripheral blood, which may help in the diagnosis of ACTH-dependent CS due to the excess of ectopic ACTH, which is very rarely diagnosed in children.

After confirming CD, the patient is qualified for surgery. The treatment of choice is transsphenoidal tumour resection. The effectiveness of the procedure depends primarily on the neurosurgeon's experience. In paediatric centres, the effectiveness of this treatment reached from 62% to 98% [24]. Remission of the CD in 200 paediatric patients was achieved in 98%, but only 3% of them had normal levels of cortisol, and the remaining 97% presented hypocortisolaemia, which is a typical response to the rapid elimination of excess ACTH, as well as earlier adrenal axis dysregulation. All our patients required transient hydrocortisone substitution, and the time of its supply clearly depended on the duration of the period from the first symptoms of the CD to its diagnosis. The symptoms of hypercortisolaemia usually disappear gradually over several months after surgery. Skin changes present at the time of diagnosis disappear gradually. According to observations of paediatric centres, acne, acanthosis nigricans, hyperpigmentation of the skin, and ease of bruising disappear up to three months after the surgery and hirsutism up to nine months after the surgery. Striae remain, which change their severity and colour from purple to pale-silver [25]. In 30% of patients successfully cured neurosurgically, disease relapse is observed.

The result of the histopathological examination supplemented with the immunohistochemical method of Ki-67 antigen labelling provides information on the degree of tumour proliferation and may have a significant clinical diagnostic significance for the course of the disease [26–28]. Tumours with invasive growth, elevated MIB-1 proliferation index, and

nuclear response to P53 are referred to as “atypical” pituitary adenomas [26, 29]. Such a diagnosis was made in two of the described patients, who, in the 1.5- and 0.5-year follow-up, have no signs of tumour recurrence. Some reports indicate that adenomas with MIB-1 proliferation index above 3% are characterised by unfavourable clinical course [27].

In patients in whom the neurosurgical treatment did not give the expected results, radiotherapy is used as the second line of therapy [30].

Despite the successful treatment of CD, in children and adolescents remaining in persistent remission of CD there are body composition disorders with visible reduction of bone mass and abdominal obesity [31]. Disorders of bone mineralisation were found in all presented patients. In the long-term observation, gradual normalisation of bone mass is observed, which is best illustrated by the densitometry results of the first of the described patients remaining in the longest observation. In the long-term observation of paediatric patients after CD treatment, an excess of total and visceral adipose tissue is observed [13, 31], which may be a risk of development of metabolic syndrome and cardiovascular disorders in the future. Obesity is still observed in all presented patients.

Due to the rare incidence of CD in children, the limitation of our study was the small number of analysed patients, which limits the strength of the conclusion.

Conclusions

Typical symptoms of CD in children, which are the inhibition of growth velocity with increasing obesity and behavioural disorders, are sometimes not noticed for a long time by their parents/careers and general practitioners. In each case of the coexistence of growth velocity inhibition (not necessarily short stature) with weight gain (not necessarily obesity), the appearance of hyperandrogenism, and depressive-anxiety disorders, the biochemical diagnostics of CS should be performed in accordance with the applicable diagnostic standard for the type of tests, their order, and interpretation of results. To diagnose CD, it is not necessary to visualise the microadenoma in MRI.

The diagnostic difficulty may be caused by the cyclical form of Cushing's disease, in which remission periods are observed with weight loss, normalisation of growth velocity, and normalisation of hormonal test results.

The treatment of choice is transsphenoidal resection of the ACTH-secreting micro/macroadenoma, which, if complete, causes a reduction of the clinical symptoms of the disease.

In most patients after neurosurgery, hydrocortisone substitution is necessary in the period depending on the duration of symptoms before the diagnosis of the disease. Obesity can be a chronic complication of CD, even after effective treatment. All patients should be carefully monitored by an experienced endocrinologist.

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