

## Sheehan's Syndrome

### Common Symptoms



fatigue



inability to lactate



low blood pressure



irregular or absent menstruation



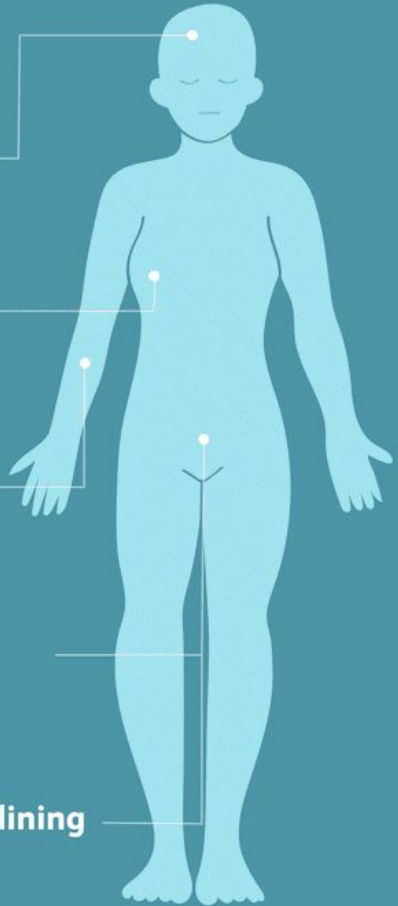
thinning of vaginal lining



loss of pubic hair



weight gain



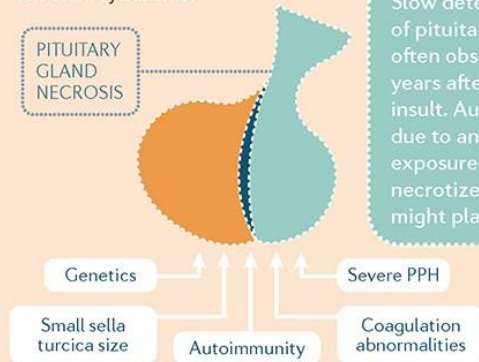
# СИНДРОМ ШИХАНА

Катамадзе Натия  
Зурабиевна

➔ Sheehan syndrome is a condition characterized by hypopituitarism due to necrosis of the pituitary gland, which is secondary to massive postpartum haemorrhage (PPH).

### EPIDEMIOLOGY

In the developed world, Sheehan syndrome is rare. The prevalence of hypopituitarism is ~40 per 1,000,000 individuals. The most common causes (~60%) of hypopituitarism are tumours (mainly in the pituitary gland) and their treatment; 6% of all hypopituitarism cases are caused by Sheehan syndrome. Prevalence of the condition is much higher in the developing world, owing to less-advanced obstetrical care. In India, ~3% of the women ≥20 years of age who have given birth are estimated to have Sheehan syndrome.

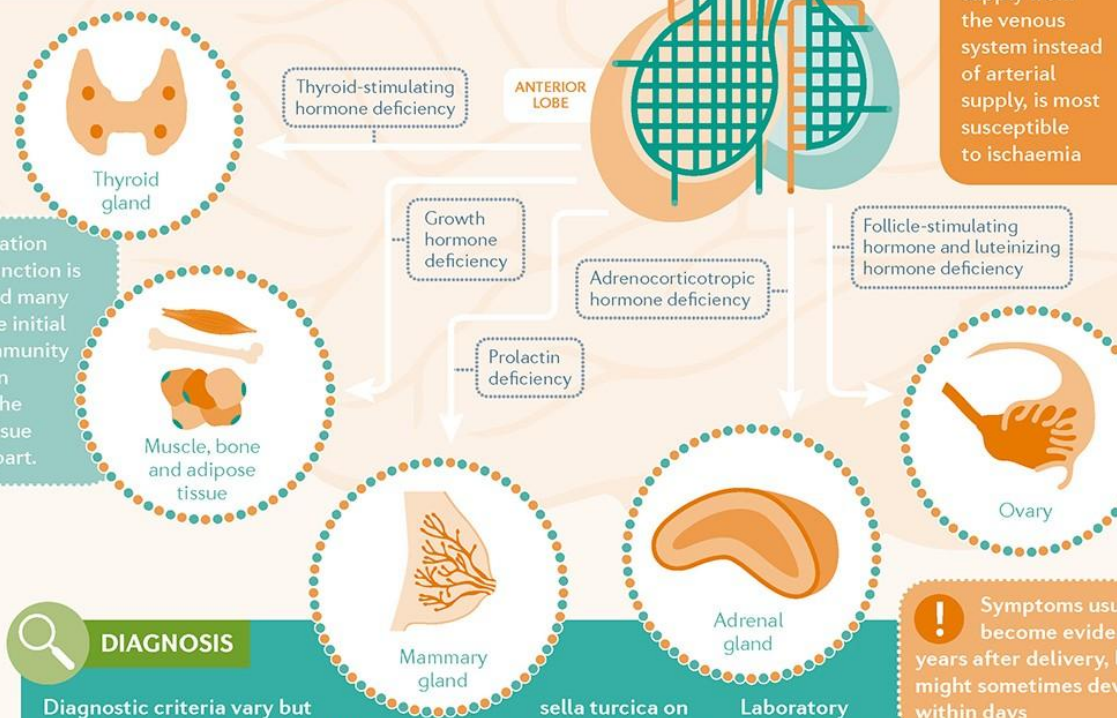


! PPH is the most important risk factor and is traditionally defined as blood loss of >500 ml after vaginal delivery or >750 ml after caesarian section. Small sella turcica (the bone enclosing the pituitary gland) size, thrombosis (associated with pregnancy or coagulation disorders), autoimmunity and/or genetics are predisposing factors.

### MECHANISMS

The enlargement of the pituitary gland during pregnancy and the early postpartum period increases its vulnerability to changes in blood flow

Postpartum necrosis of the pituitary gland



Slow deterioration of pituitary function is often observed many years after the initial insult. Autoimmunity due to antigen exposure by the necrotized tissue might play a part.

### DIAGNOSIS

Diagnostic criteria vary but usually involve a history of PPH, absence of postpartum lactation, failure to resume regular menstruation, varying degrees of anterior pituitary hormone insufficiencies and an empty sella turcica on imaging. Symptoms are caused by a decrease or absence of at least one pituitary hormone, and vary from nonspecific symptoms (such as fatigue) to severe adrenal crisis.

Laboratory tests (hormone and electrolyte levels and blood cell counts), hormone stimulation tests and imaging of the pituitary gland are used to confirm diagnosis.

! Symptoms usually become evident years after delivery, but might sometimes develop within days

### MANAGEMENT

Hormone replacement therapy is the only available management option, although it does not improve pituitary function or halt the progression of pituitary gland necrosis. In accordance with the location of hormone-secreting cells relative to the vasculature, the secretion of growth hormone and prolactin is most commonly affected, followed by follicle-stimulating hormone and luteinizing hormone. Severe pituitary gland necrosis also affects the secretion of thyroid-stimulating hormone and adrenocorticotrophic hormone. Although rare, involvement of the posterior lobe of the pituitary gland can result in arginine vasopressin deficiency, with diabetes insipidus as a consequence.

The anterior lobe, which mainly receives blood supply from the venous system instead of arterial supply, is most susceptible to ischaemia

### PREVENTION

Prevention of Sheehan syndrome requires aggressive management (for example, by giving intravenous fluids) and minimizing the risk of PPH. The WHO recommends a combination of interventions such as umbilical cord clamping and cutting, controlled cord traction (to minimize retention of the placenta) and the use of a uterotonic agent to stimulate uterine contraction. Other risk factors for haemorrhage, including anaemia, should be prevented, for example, via iron supplementation.







## MECHANISMS

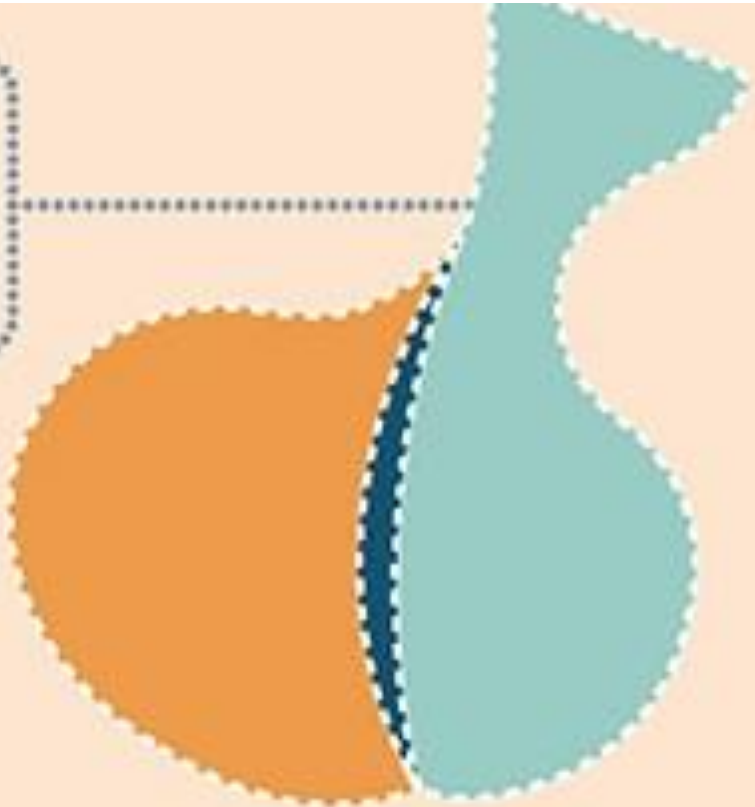


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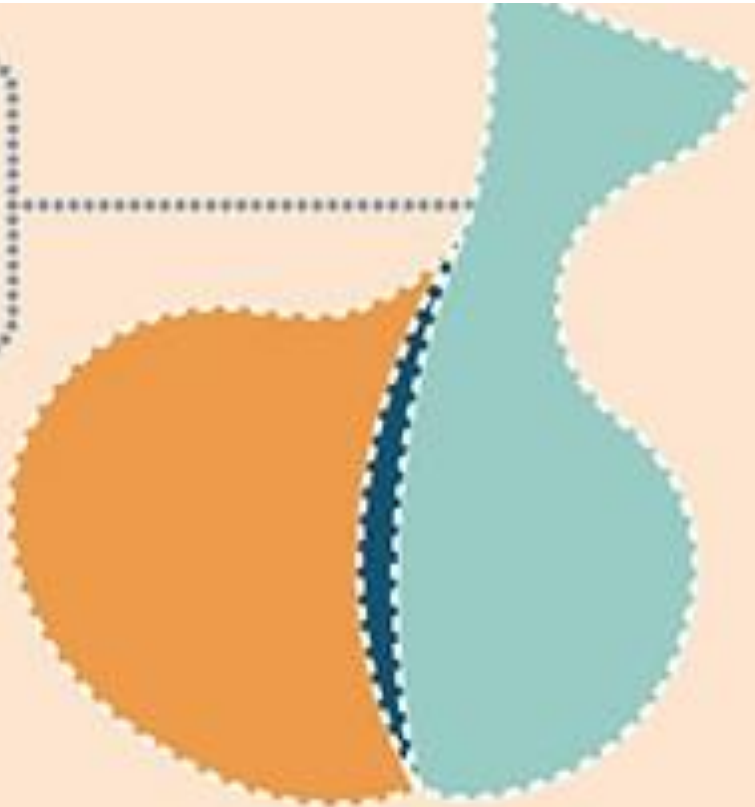
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sella turcica on imaging. Symptoms are caused by a decrease or absence of at least one pituitary hormone, and vary from nonspecific symptoms (such as fatigue) to severe adrenal crisis.



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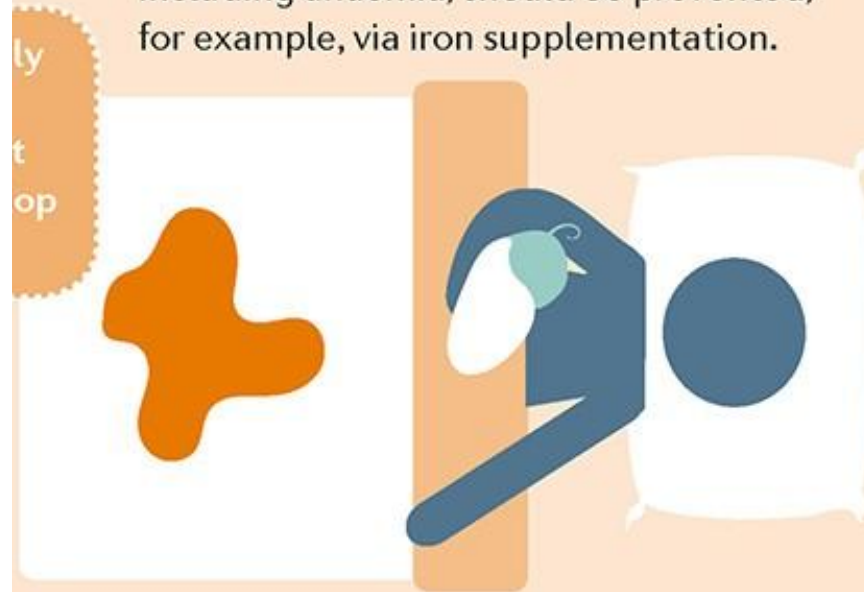


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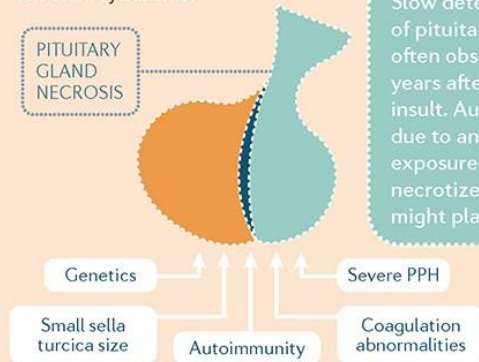
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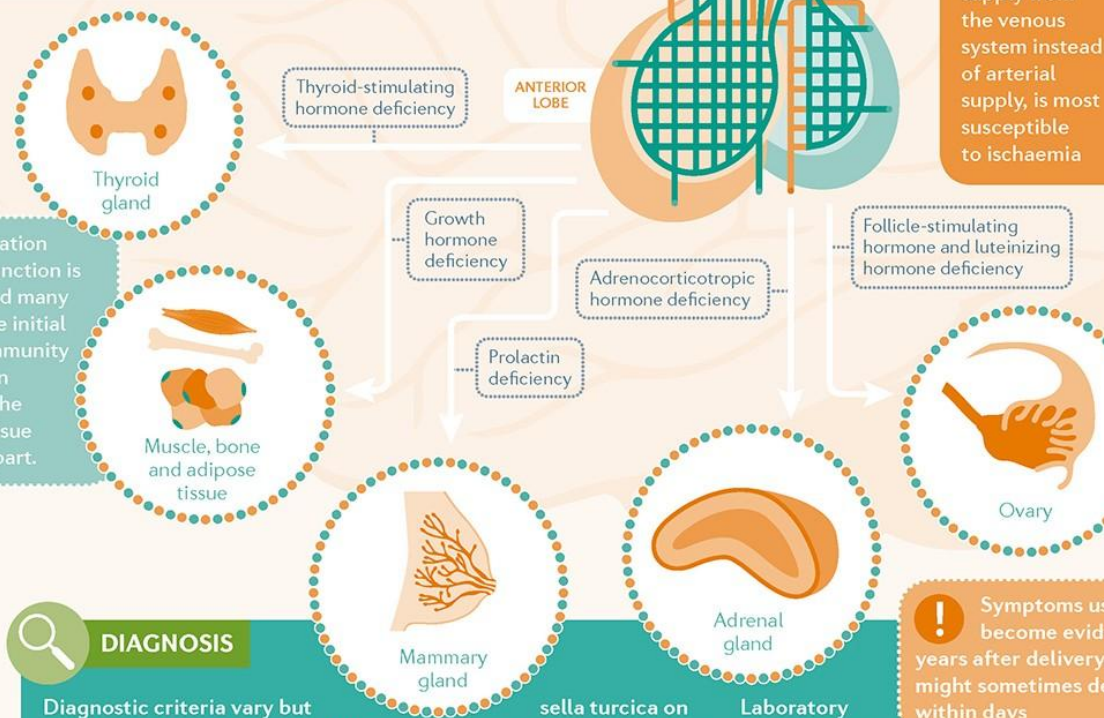
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# Extensive investigation of 114 patients with Sheehan's syndrome: a continuing disorder

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Departments of <sup>1</sup>Endocrinology and <sup>2</sup>Radiology, Erciyes University Medical School, 38039 Kayseri, Turkey and

<sup>3</sup>Department of Endocrinology, Ondokuz Mayıs University Medical School, Samsun, Turkey

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## **Abstract**

*Objective:* Sheehan's syndrome (SS) is a well-known cause of hypopituitarism resulting from *postpartum* pituitary necrosis. Because of its rarity in Western society, its diagnosis is often overlooked. We aimed to investigate the clinical, laboratory, and radiological aspects of SS in a large number of patients.

*Study design:* A retrospective assessment of the medical records of 114 patients with SS was conducted. In addition, sella turcica volumes of 29 healthy women were compared with those of patients by magnetic resonance imaging examinations.

*Results:* The mean period of diagnostic delay was 19.7 years in patients with SS. It was found that 52.6% of patients had nonspecific complaints, 30.7% had complaints related to adrenal insufficiency, and 9.6% had complaints related to hypogonadism when diagnosed. At the time of diagnosis, 55.3% of the patients had panhypopituitarism, while 44.7% had partial hypopituitarism. The number of deficient hormones was found to be increased over the years. None of the patients whose basal prolactin was below 4.0 ng/ml had adequate prolactin responses to TRH test, while all patients whose basal prolactin was above 7.8 ng/ml had adequate responses. Mean sella volume was found to be significantly lower in the SS group ( $340.5 \pm 214 \text{ mm}^3$ ) than that in the healthy group ( $602.5 \pm 192 \text{ mm}^3$ ).

*Conclusions:* SS is a common cause of hypopituitarism in underdeveloped and developing countries. The main reasons for diagnostic delay seem to be the high frequency of patients with nonspecific complaints and neglect of SS. In addition, the TRH stimulation test was found to have a high sensitivity and specificity to recognize PRL deficiency. Furthermore, small sella size may have an important contributing role in the etiopathogenesis of SS.



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## Results

The mean age of 114 patients previously diagnosed with SS between 1985 and 2013 was  $63.2 \pm 12.5$  years, and age at diagnosis was  $52.1 \pm 12.7$  years. When the past histories of patients with SS were analyzed, mean age at the last delivery was  $32.4 \pm 6.5$  years and thereby the period of diagnostic delay was  $19.7 \pm 10.2$  years. After diagnosis, the follow-up time was  $7.4 \pm 6.9$  years. Nine (7.9%) cases had died and 55 (48.2%) were alive. The other 50 (43.9) patients did not attend follow-up visits in the last 3 years, and therefore it was not known whether they were still alive.

**Table 3** Deficient hormones of anterior pituitary gland in patients with SS.

<b>Deficient hormones</b>	<b>Number (%) of patients</b>
All hormones	63 (55.3)
FSH-LH+GH+ACTH+TSH	17 (14.9)
FSH-LH+GH+TSH+PRL	12 (10.5)
FSH-LH+GH+TSH	11 (9.6)
FSH-LH+GH+PRL	5 (4.4)
FSH-LH+GH	4 (3.5)
FSH-LH+GH+ACTH+PRL	1 (0.9)
FSH-LH+GH+ACTH	1 (0.9)
Total	114 (100)



**Table 3** Deficient hormones of anterior pituitary gland in patients with SS.

**Deficient hormones**

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FSH-LH+GH+ACTH+PR
FSH-LH+GH+ACTH
Total

**Table 4** Progression of hormonal deficiency in ten patients with SS.

<b>Patient nos</b>	<b>Intact hormones at date of diagnosis</b>	<b>After a period of (years)</b>	<b>Intact hormones in last clinical visit</b>
21	ACTH	18	None
31	ACTH+TSH+PRL	15	ACTH+PRL
35	ACTH	22	None
60	ACTH	6	None
64	ACTH+PRL	9	None
66	PRL+TSH	12	None
81	PRL	22	None
90	ACTH	11	None
95	ACTH	10	None
109	ACTH+TSH	8	TSH

# Clinical Report of 28 Patients with Sheehan's Syndrome

MURAT SERT, TAMER TETIKER, SINAN KIRIM AND MUSTAFA KOCAK

*Cukurova University, Medical Faculty, Department of Internal Medicine, Endocrinology Division, 01330 Adana, Turkey*

**Abstract.** The aim of the present study was to determine the clinical and hormonal characteristics with Sheehan's syndrome in 28 cases that we had diagnosed and followed in the last 20 years. Twenty-eight patients with Sheehan's syndrome, diagnosed and followed at our University Endocrinology Clinic in the last 20 years were reported in the study. Medical history, physical examination, routine laboratory examinations, pituitary hormone analysis, CT and/or MRI scan of the sella of the patients were reviewed. All patients had a history of massive hemorrhage at delivery and physical signs of Sheehan's syndrome. Twenty-six of them lacked postpartum milk production, followed by failure of resumption of menses. There were 9 subjects with disturbances in consciousness associated with hyponatremia on admittance. All 28 patients had secondary hypothyroidism, adrenal cortex failure, hypogonadotropic hypogonadism and growth hormone deficiency. Diabetes insipidus has not been found in any patient. Empty sellae were revealed in 8 patients by CT and/or MRI scan. Sheehan's syndrome is still encountered in clinical practice occasionally. If not diagnosed early, it could cause increased morbidity and mortality. The most important clues for diagnosis of Sheehan's syndrome are lack of lactation and failure of menstrual resumption after a delivery complicated with severe hemorrhage.

*Key words:* Sheehan's syndrome, Pituitary failure, Hyponatremia, Empty sellae

*(Endocrine Journal 50: 297-301, 2003)*

**Table 1.** The characteristics of the patients with Sheehan's syndrome.

Patients	Age at diagnosis (year)	Duration of illness (year)	Total number of pregnancy	Lactation	Serum sodium (mEq/L)	Consciousness on admittance	Drugs of substitution	CT and/or MRI scan of sellae
1	45	8	3	-	N	N	L, P, HR	N
2	60	17	8	-	110	Coma	L, P	ES
3	70	20	6	-	N	N	L, P	N
4	60	22	3	-	132	Confusion	L, P	N
5	30	7	1	-	128	N	L, P, HR	N
6	34	6	2	-	N	N	L, P, HR	N
7	64	9	2	-	N	N	L, P	N
8	35	8	1	-	N	N	L, P, HR	ES
9	51	19	1	-	110	Stupor	L, P, HR	ES
10	48	10	2	-	N	N	L, P, HR	N
11	38	9	5	-	N	N	L, P, HR	N
12	50	15	4	-	N	N	L, P, HR	N
13	35	6	4	-	N	N	L, P, HR	N
14	60	18	9	-	N	N	L, P, HR	N
15	35	6	1	-	N	N	L, P, HR	N
16	39	8	1	+	N	N	L, P, HR	N
17	45	12	1	-	N	N	L, P, HR	N
18	50	17	3	-	N	N	L, P, HR	N
19	46	20	1	-	N	N	L, P, HR	N
20	45	14	1	-	N	N	L, P, HR	N
21	43	15	1	+	N	N	L, P, HR	N
22	58	10	4	-	126	Coma	L, P, HR	ES
23	67	30	7	-	119	Coma	L, P	ES
24	47	20	5	-	101	Confusion	L, P, HR	ES
25	57	20	3	-	N	Confusion	L, P	ES
26	49	19	5	-	N	N	L, P, HR	ES
27	44	12	6	-	110	Coma	L, P, HR	N
28	45	13	4	-	114	Coma	L, P, HR	N

+ Present, - Absent, N: Normal, ES: Empty Sellae, L: Levothyroxine (100-150 ug/day), P: Prednisolon (7.5-10 mg/day), HR: Hormone replacement (estrogen plus progesterone)



Secondary hypothyroidism, secondary adrenal failure, hypogonadotropic hypogonadism and growth hormone deficiency were present in all patients. Serum hormone results (means  $\pm$  SD) were found as follows: TSH:  $1.32 \pm 0.85$  mIU/ml (R: 0.37 to 4.2, normal range [N]: 0.47–5.01), FT<sub>3</sub>:  $2.09 \pm 1.7$  pg/dl (R: 0.17–2.70, N: 2.3–4.2), FT<sub>4</sub>:  $0.43 \pm 0.28$  ng/dl (R: 0.11–0.80, N: 0.89–1.80), FSH:  $5.01 \pm 3.21$  mIU/ml (R: 0.4–11, normal range for menopause 40–200), LH:  $2.07 \pm 1.95$  mIU/ml (R: 0.08–7.3, N for menopause 40–200), GH:  $0.23 \pm 0.25$  ng/ml (R: 0–0.9, N: >7 after insulin induced hypoglycemia), prolactin:  $2.38 \pm 1.53$  ng/ml (R: 0.1–4.5, N: 2.7–26), ACTH:  $17.15 \pm 11.03$  pg/ml (R: 0.1–43.5, N: 10–60), cortisol:  $2.96 \pm 3.37$   $\mu$ g/dl (R: 0.1–11.4, N: >20 after insulin induced hypoglycemia). Individual value of each hormone is

presented (see Fig. 1). Diabetes insipidus was not observed in any patient.

CT and/or MRI scan of sella of 28 subjects revealed 20 normal and 8 empty sellae.

# Синдром Шихана у пациентки с патологией щитовидной железы



Волкова А.Р.,  
Остроухова Е.Н.,  
Дора С.В.,  
Власова К.А.,  
Семикова Г.В.

Федеральное государственное бюджетное образовательное учреждение высшего образования «Первый Санкт-Петербургский государственный медицинский университет им. акад. И.П. Павлова» Министерства здравоохранения Российской Федерации, 197022, г. Санкт-Петербург, Российская Федерация

*Пациентка П.*, 34 года, обратилась на прием 14.08.2019 с жалобами на выраженную слабость, пониженный аппетит, тошноту, сердцебиение, снижение массы тела на 1,5 кг за последнюю неделю. Описанные жалобы появились в конце мая 2019 г. и постепенно прогрессировали.

## Пациентка П., 34 года



ТТГ крови – **5,6**  
**мМЕ/л** (норма  
0,2–2,5),  
св.Т<sub>4</sub>–**12,1** **пмоль/л**  
(норма 12–22).  
Был поставлен  
диагноз:  
«субклинический  
гипотиреоз АИТ?». Назначены **L-**  
**тироксин** (50  
мкг/сут), **йодомарин**  
(200 мкг/сут)



## Пациентка П., 34 года

ТТГ 1,8  
мМЕ/л.

12.10.2018  
(6 недель)

22.11.2018  
(12 недель)

24.05.2019  
(39  
неделя)

С июня по  
август  
2019 г.

21.07.2019

11.08.2019

14.08.2019

ТТГ крови – 5,6  
мМЕ/л (норма  
0,2–2,5),  
св.Т<sub>4</sub>–12,1 пмоль/л  
(норма 12–22).  
Был поставлен  
диагноз:  
«субклинический  
гипотиреоз АИТ?». Назначены L-  
тироксин (50  
мкг/сут), йодомарин  
(200 мкг/сут)

## Пациентка П., 34 года

ТТГ 1,8  
мМЕ/л.



срочные  
роды

12.10.2018  
(6 недель)

22.11.2018  
(12 недель)

24.05.2019  
(39  
неделя)

С июня по  
август  
2019 г.

21.07.2019

11.08.2019

14.08.2019

ТТГ крови – 5,6  
мМЕ/л (норма  
0,2–2,5),  
св.Т<sub>4</sub>–12,1 пмоль/л  
(норма 12–22).  
Был поставлен  
диагноз:  
«субклинический  
гипотиреоз АИТ?». Назначены L-  
тироксин (50  
мкг/сут), йодомарин  
(200 мкг/сут)

## Пациентка П., 34 года

ТТГ 1,8  
мМЕ/л.



срочные  
роды

12.10.2018  
(6 неделя)

22.11.2018  
(12 недель)

24.05.2019  
(39  
неделя)

С июня по  
август  
2019 г.

21.07.2019

11.08.2019

14.08.2019

ТТГ крови – **5,6**  
**мМЕ/л** (норма  
0,2–2,5),  
св.Т<sub>4</sub>–**12,1** **пмоль/л**  
(норма 12–22).  
Был поставлен  
диагноз:  
«субклинический  
гипотиреоз АИТ?». Назначены **L-**  
**тироксин** (50  
мкг/сут), **йодомарин**  
(200 мкг/сут)

Через 40 мин после родов -  
нарушение гемодинамики с  
затруднением дыхания,  
слабостью, массивное  
кровотечение, **гемоглобина до**  
**59 г/л** (до родов гемоглобин  
составлял 116 г/л).  
Пациентка переведена в  
реанимационное отделение  
областной больницы –  
экстренное переливание крови.  
Выписана из стационара в  
удовлетворительном  
состоянии на 7-й день,  
гемоглобин составил **88 г/л.**



## Пациентка П., 34 года

ТТГ 1,8  
мМЕ/л.



срочные  
роды

12.10.2018  
(6 недель)

22.11.2018  
(12 недель)

24.05.2019  
(39  
неделя)

С июня по  
август  
2019 г.

21.07.2019

11.08.2019

14.08.2019

ТТГ крови – **5,6 мМЕ/л** (норма 0,2–2,5),  
св.Т<sub>4</sub>–**12,1 пмоль/л** (норма 12–22).  
Был поставлен  
диагноз:  
«субклинический  
гипотиреоз АИТ?». Назначены **L-тироксин** (50 мкг/сут), **йодомарин** (200 мкг/сут)

Через 40 мин после родов - нарушение гемодинамики с затруднением дыхания, слабостью, массивное кровотечение, **гемоглобина до 59 г/л** (до родов гемоглобин составлял 116 г/л). Пациентка переведена в реанимационное отделение областной больницы – экстренное переливание крови. Выписана из стационара в удовлетворительном состоянии на 7-й день, гемоглобин составил **88 г/л**.

МРТ головного мозга и гипофиза от 21.07.2019 – размеры гипофиза в норме, имеет однородный сигнал

## Пациентка П., 34 года

ТТГ 1,8  
мМЕ/л.



срочные  
роды

12.10.2018  
(6 недель)

22.11.2018  
(12 недель)

24.05.2019  
(39  
неделя)

С июня по  
август  
2019 г.

21.07.2019

11.08.2019

14.08.2019

ТТГ крови – **5,6**  
**мМЕ/л** (норма  
0,2–2,5),  
св. Т<sub>4</sub> – **12,1** **пмоль/л**  
(норма 12–22).  
Был поставлен  
диагноз:  
«субклинический  
гипотиреоз АИТ?». Назначены **L-**  
**тироксин** (50  
мкг/сут), **йодомарин**  
(200 мкг/сут)

Через 40 мин после родов -  
нарушение гемодинамики с  
затруднением дыхания,  
слабостью, массивное  
кровотечение, **гемоглобина до**  
**59 г/л** (до родов гемоглобин  
составлял 116 г/л).  
Пациентка переведена в  
реанимационное отделение  
областной больницы –  
экстренное переливание крови.  
Выписана из стационара в  
удовлетворительном  
состоянии на 7-й день,  
гемоглобин составил **88 г/л**.

МРТ головного  
мозга и гипофиза  
от 21.07.2019 –  
размеры  
гипофиза в  
норме, имеет  
однородный  
сигнал

ТТГ крови – **0,0025**  
**мМЕ/л, св. Т<sub>4</sub> – 28,67**  
**пмоль/л** (норма  
9–19,05), св. Т<sub>3</sub> – 4,24  
**пмоль/л** (норма  
4,0–7,4 пмоль/л),  
гемоглобин – **118 г/л**.  
Обратилась к  
эндокринологу,  
поставлен диагноз  
**диффузного**  
**токсического зоба,**

## Пациентка П., 34 года

ТТГ 1,8  
мМЕ/л.



срочные  
роды

Рекомендован прием  
тирозола (по 10 мг 3 раза  
в день )  
(не принимала)

12.10.2018  
(6 недель)

22.11.2018  
(12 недель)

24.05.2019  
(39  
неделя)

С июня по  
август  
2019 г.

21.07.2019

11.08.2019

14.08.2019

ТТГ крови – 5,6  
мМЕ/л (норма  
0,2–2,5),  
св. Т<sub>4</sub>–12,1 пмоль/л  
(норма 12–22).  
Был поставлен  
диагноз:  
«субклинический  
гипотиреоз АИТ?». Назначены L-  
тироксин (50  
мкг/сут), йодомарин  
(200 мкг/сут)

Через 40 мин после родов -  
нарушение гемодинамики с  
затруднением дыхания,  
слабостью, массивное  
кровотечение, гемоглобина до  
59 г/л (до родов гемоглобин  
составлял 116 г/л).  
Пациентка переведена в  
реанимационное отделение  
областной больницы –  
экстренное переливание крови.  
Выписана из стационара в  
удовлетворительном  
состоянии на 7-й день,  
гемоглобин составил 88 г/л.

МРТ головного  
мозга и гипофиза  
от 21.07.2019 –  
размеры  
гипофиза в  
норме, имеет  
однородный  
сигнал

ТТГ крови – 0,0025  
мМЕ/л, св. Т<sub>4</sub>–28,67  
пмоль/л (норма  
9–19,05), св. Т<sub>3</sub>–4,24  
пмоль/л (норма  
4,0–7,4 пмоль/л),  
гемоглобин – 118 г/л.  
Обратилась к  
эндокринологу,  
поставлен диагноз  
диффузного  
токсического зоба,



# 14.08.2019 обратилась на прием к эндокринологу в ПСПбГМУ им. акад. И.П. Павлова.

Объективный осмотр: общее состояние удовлетворительное, кожные покровы бледные, поредение волос в паховой и подмышечной областях, рост – 161 см, масса тела – 52,8 кг, индекс массы тела – 20,4 кг/м<sup>2</sup>, пульс – 82 в минуту, удовлетворительных свойств, артериальное давление на левой руке – 90/50 мм рт.ст. Тоны сердца ясные, чистые.

Щитовидная железа не увеличена, сосудистый шум над железой не выслушивается, узловые образования пальпаторно не определяются. Глазные симптомы отрицательны.

Клинически не складывалось впечатление в пользу тиреотоксикоза.

Рекомендовано дообследование: определение в крови кортизола, адренкортиктропного гормона (АКТГ), анализ крови на антитела к тиреопероксидазе (ТПО), антитела к рецептору тиреотропного гормона (ТТГ).

**16.08.2019:** кортизол – **1,8 нмоль/л** (норма 171–536), АКТГ – 5,1 пг/мл (норма 0–46), антитела к рецептору ТТГ – 0,75 МЕ/л (норма 0–1,75), антитела к ТПО – 7,7 Ед/мл (норма 0–34).  
Поставлен диагноз: «послеродовый тиреодит, эутиреоз? Вторичная надпочечниковая недостаточность. Синдром Шихана».

**16.08.2019:** кортизол – **1,8 нмоль/л** (норма 171–536), АКТГ – 5,1 пг/мл (норма 0–46), антитела к рецептору ТТГ – 0,75 МЕ/л (норма 0–1,75), антитела к ТПО – 7,7 Ед/мл (норма 0–34).  
Поставлен диагноз: «послеродовой тиреодит, эутиреоз? Вторичная надпочечниковая недостаточность. Синдром Шихана».

**16.08.2019:** **кортеф** (50 мг/сут: 30 мг – утром, 20 мг – 16.00).  
Рекомендовано дообследование: ФСГ, ЛГ, эстрадиол, пролактин, соматотропный гормон (СТГ).



**16.08.2019:** кортизол – **1,8 нмоль/л** (норма 171–536), АКТГ – 5,1 пг/мл (норма 0–46), антитела к рецептору ТТГ – 0,75 МЕ/л (норма 0–1,75), антитела к ТПО – 7,7 Ед/мл (норма 0–34).  
Поставлен диагноз: «послеродовой тиреодит, эутиреоз? Вторичная надпочечниковая недостаточность. Синдром Шихана».

**16.08.2019:** **кортеф** (50 мг/сут: 30 мг – утром, 20 мг – 16.00).  
Рекомендовано дообследование: ФСГ, ЛГ, эстрадиол, пролактин, соматотропный гормон (СТГ).

На 2-й день приема глюкокортикоидов пациентка отметила улучшение самочувствия: уменьшилась слабость, появился аппетит.

**16.08.2019:** кортизол – **1,8 нмоль/л** (норма 171–536), АКТГ – 5,1 пг/мл (норма 0–46), антитела к рецептору ТТГ – 0,75 МЕ/л (норма 0–1,75), антитела к ТПО – 7,7 Ед/мл (норма 0–34).  
Поставлен диагноз: «послеродовый тиреодит, эутиреоз? Вторичная надпочечниковая недостаточность. Синдром Шихана».

**16.08.2019:** **кортеф** (50 мг/сут: 30 мг – утром, 20 мг – 16.00).  
Рекомендовано дообследование: ФСГ, ЛГ, эстрадиол, пролактин, соматотропный гормон (СТГ).

На 2-й день приема глюкокортикоидов пациентка отметила улучшение самочувствия: уменьшилась слабость, появился аппетит.

**28.08.2019:** ФСГ – **0,85 мМЕ/л** (норма 3,5–12,5), ЛГ – **0,41 мМЕ/л** (норма 1,69–15,0), эстрадиол – **12 пмоль/л** (90–860 норма), пролактин – 121 мкг/л (норма 130–540), св.Т3–3,3 пмоль/л (норма 2,6–5,7), св.Т4–8,25 пмоль/л (норма 9–19), ТТГ – **0,07 мМЕ/л** (норма 0,3–3,4).

**16.08.2019:** кортизол – **1,8 нмоль/л** (норма 171–536), АКТГ – 5,1 пг/мл (норма 0–46), антитела к рецептору ТТГ – 0,75 МЕ/л (норма 0–1,75), антитела к ТПО – 7,7 Ед/мл (норма 0–34). Поставлен диагноз: «послеродовый тиреодит, эутиреоз? Вторичная надпочечниковая недостаточность. Синдром Шихана».

**16.08.2019:** **кортеф** (50 мг/сут: 30 мг – утром, 20 мг – 16.00).  
Рекомендовано дообследование: ФСГ, ЛГ, эстрадиол, пролактин, соматотропный гормон (СТГ).

На 2-й день приема глюкокортикоидов пациентка отметила улучшение самочувствия: уменьшилась слабость, появился аппетит.

**28.08.2019:** ФСГ – **0,85 мМЕ/л** (норма 3,5–12,5), ЛГ – **0,41 мМЕ/л** (норма 1,69–15,0), эстрадиол – **12 пмоль/л** (90–860 норма), пролактин – 121 мкг/л (норма 130–540), св.Т3–3,3 пмоль/л (норма 2,6–5,7), св.Т4–8,25 пмоль/л (норма 9–19), ТТГ – **0,07 мМЕ/л** (норма 0,3–3,4).

**29.08.2019:** состояние удовлетворительное, кожные покровы обычного цвета и влажности, рост – 161 см, масса тела – 54,1 кг (**увеличение на 1,3 кг**), ИМТ – 20,9 кг/м<sup>2</sup>, артериальное давление – 110/80 мм рт.ст., пульс – 78 в минуту, ритмичный



**Диагноз:** недостаточность передней доли гипофиза. Синдром Шихана. Вторичная недостаточность коры надпочечников. Вторичный гипотиреоз. Вторичный гипогонадизм.

кортеф – 10 мг/сут (по 1 таблетке утром и по 1/2 таблетки в 16.00)

эутирокс (L-тироксин) – 25 мкг 1 раз в день, утром, натощак, за 40 мин до завтрака; фемостон 2/10.

15.09.2019: кортизол в суточной моче – 267,96 мкг/сут (норма 58–403), св.Т3–3,3 пмоль/л (норма 2,6–5,7), св.Т4–14,5 пмоль/л (норма 9–19).  
Рекомендовано дальнейшее наблюдение у эндокринолога, гинеколога.