CASE REPORT

Decreased Serum Adiponectin Level during Catecholamine Crisis in an Obese Patient With Pheochromocytoma

Yukiyoshi Okauchi, Chisaki Ishibashi, Kunihiko Shu, Shiro Adachi and Ikuo Mineo

Abstract:
We herein report the case of a 37-year-old man with both pheochromocytoma and visceral fat accumulation and describe the sequential changes in his adiponectin levels throughout the clinical course from catecholamine crisis until the follow-up for adrenalectomy. His adiponectin level decreased during catecholamine crisis and increased after adrenalectomy. However, his adiponectin level decreased again at two years postoperatively when his visceral fat area greatly increased. This case suggests that catecholamines and visceral fat volume may affect adiponectin metabolism in subjects with pheochromocytoma, which may precipitate cardiovascular complications in this endocrine disease.

Key words: pheochromocytoma, visceral fat area, adiponectin

Introduction
Pheochromocytoma is a catecholamine-producing tumor that is usually derived from the adrenal medulla (1). Catecholamine crisis is a life-threatening condition in this disease because of the excessive secretion of catecholamines from the tumor, which presents spontaneously or as the result of several inducing factors (2). Patients with pheochromocytoma are usually lean because of adipose tissue lipolysis by catecholamines. However, recent lifestyle changes have increased the number of obese individuals, including those with pheochromocytoma.

Adiponectin is an adipocyte-derived plasma protein with anti-diabetic, anti-inflammatory, anti-atherogenic, and anti-cardiomyopathic properties (3). Adiponectin levels are regulated by various factors and altered in different conditions. The levels have been found to be decreased in subjects with abdominal obesity, a central feature of metabolic syndrome (3). Although catecholamines are metabolic regulators of adipose tissue, their effects on adiponectin remain unclear. Furthermore, the adiponectin levels of patients with pheochromocytoma are controversial, with reports of their being increased (4), decreased (5), or unchanged (6). To date, there are no studies investigating the relationship between the adiponectin levels and visceral fat area (VFA) in pheochromocytoma patients.

We herein report a patient with both abdominal obesity and pheochromocytoma and describe the sequential changes in the adiponectin levels as well as the VFA throughout the clinical course, from catecholamine crisis to recovery.

Case Report
A 37-year-old man was diagnosed with hypertension and obesity at an annual health checkup, followed by further examinations in which a left adrenal tumor was revealed. Subsequently, he was referred to our hospital for endocrinological evaluations and was scheduled to be hospitalized. His height and body weight were 168 cm and 91 kg (body mass index [BMI]: 32.2 kg/m²), respectively. Five days later, he presented to the emergency room of our hospital with a complaint of general fatigue and left abdominal pain. He had severe hypertension with a blood pressure of 234/120 mmHg and exhibited sinus tachycardia of 146 beats per minute. He was admitted to the intensive care unit with a diagnosis of catecholamine crisis, probably due to pheochromocytoma. We started antihypertensive therapy using phenolamine, nicardipine, and doxazosin, which kept his blood pressure and heart rate under control.

1Diabetes Center, Toyonaka Municipal Hospital, Japan and 2Pathology Section, Toyonaka Municipal Hospital, Japan
Received: February 27, 2017; Accepted: September 1, 2017; Advance Publication by J-STAGE: December 27, 2017
Correspondence to Dr. Yukiyoshi Okauchi, yokauchi0713@gmail.com
metabolites in the plasma and urine (Table).

The diagnosis of pheochromocytoma was confirmed by the marked elevation of catecholamines and their metabolites in the plasma and urine (Table).

Other routine laboratory test data, including findings for the liver function, renal function, and electrolyte levels, were normal. The plasma glucose level increased over 200 mg/dL, despite a normal HbA1c value. Other routine laboratory test data, including findings for the liver function, renal function, and electrolyte levels, were normal. The diagnosis of pheochromocytoma was confirmed by the marked elevation of catecholamines and their metabolites in the plasma and urine (Table).

On day 8, his blood pressure level rose to 200 mmHg, with recurrence of left-sided abdominal pain. Computed tomography (CT) revealed a left adrenal tumor, inhomogeneous inside and surrounded by panniculitis, on the day of admission (Fig. 1-A), and necrosis in the left adrenal tumor and peri-adrenal fat stranding were also noted (Fig. 1-B). Because of the unstable blood pressure, we chose conservative therapy with α- and β-adrenoceptor blockade instead of emergency adrenalectomy. The catecholamine crisis and abdominal pain gradually resolved, and he was discharged.

Table. Laboratory Tests on Admission.

| Blood Chemistry | Reference ranges | Hormone | Reference ranges |
|-----------------|-----------------|---------|-----------------|
| Complete blood count |                   | ACTH (pg/mL) | 17.4 | 7.2-63.3 |
|                  |                  | Cortisol (μg/dL) | 20.8 | 4.0-19.3 |
|                  |                  | TSH (μIU/mL) | 0.49 | 0.35-4.94 |
|                  |                  | FT4 (ng/mL) | 0.97 | 0.70-1.48 |
|                  |                  | Intact PTH (pg/mL) | 107 | 10-65 |
|                  |                  | Calcitonin (pg/mL) | 17.0 | 15.0-86.0 |
|                  |                  | renin activity (ng/mL/hr) | 12.3 | 0.2-2.3 |
|                  |                  | Aldosterone (pg/mL) | 314 | 30-159 |
|                  |                  | DHEA-S (μg/dL) | 64 | 98-516 |
|                  |                  | Adrenaline (ng/mL) | 3.8 | 0.0-0.17 |
|                  |                  | Noradrenaline (ng/mL) | 6.7 | 0.15-0.57 |
|                  |                  | Dopamine (ng/mL) | 0.03 | <0.03 |
|                  |                  | urinary metanephrine (mg/day) | 11 | 0.05-0.23 |
|                  |                  | urinary normetanephrine (mg/day) | 5.6 | 0.07-0.26 |
|                  |                  | urinary VMA (mg/day) | 49.4 | 1.5-4.3 |

Urinalysis

|                        | Reference ranges |
|------------------------|-----------------|
| Protein                | (1+)            |
| Glucose                | (1+)            |
| Ketone                 | (1+)            |
| Occult Blood           | (2+)            |

His white blood cell count on admission increased with a slight rise in C-reactive protein (Table). The plasma glucose level increased over 200 mg/dL, despite a normal HbA1c value. Other routine laboratory test data, including findings for the liver function, renal function, and electrolyte levels, were normal. The diagnosis of pheochromocytoma was confirmed by the marked elevation of catecholamines and their metabolites in the plasma and urine (Table).

On day 8, his blood pressure level rose to 200 mmHg, with recurrence of left-sided abdominal pain. Computed tomography (CT) revealed a left adrenal tumor, inhomogeneous inside and surrounded by panniculitis, on the day of admission (Fig. 1-A), and necrosis in the left adrenal tumor and peri-adrenal fat stranding were also noted (Fig. 1-B). Because of the unstable blood pressure, we chose conservative therapy with α- and β-adrenoceptor blockade instead of emergency adrenalectomy. The catecholamine crisis and abdominal pain gradually resolved, and he was discharged.

Figure 1. At the time of admission, abdominal CT imaging showed a left adrenal tumor, inhomogeneous inside and surrounded by panniculitis [diamond 65 mm (arrow)] and visceral fat area of 153 cm² (A). On day 8, CT showed necrosis in the left adrenal tumor (arrow) and periadrenal fat stranding (B).
from the hospital on day 23. On day 37, he underwent successful left adrenalectomy at the Department of Urology in our hospital. Macroscopic (Fig. 2-A) and microscopic (Fig. 2-B) examinations of the resected tumor revealed massive necrosis, probably due to ischemia rather than tumor necrosis. The limited viable tumor tissue revealed no malignant cells. Browning of adipocytes in the adipose tissue surrounding the tumor was not observed.

The VFA was measured by CT at the umbilical level using the FatScan software program (N2system, Osaka, Japan) (7). The serum adiponectin levels were measured using an enzyme-linked immunosorbent assay kit (Otsuka Pharmaceutical Co., Ltd., Tokyo, Japan) (8). On the day of admission at the time of the catecholamine crisis, his VFA was 153 cm², and his serum adiponectin level was 6.9 μg/mL (Fig. 3). After adrenalectomy, his VFA decreased slightly, and his adiponectin level increased to 7.8 μg/mL. The VFA at 6 months postoperatively remained unchanged at 148 cm², and the adiponectin level at 12 months postoperatively also did not increase (8.0 μg/mL). At 2 years postoperatively, the VFA increased to 224 cm², and the serum adiponectin level decreased to 7.4 μg/mL (Fig. 3).

Homeostasis model assessment of insulin resistance (HOMA-IR), a marker of insulin resistance, decreased after adrenalectomy and increased at two years postoperatively (Fig. 3). Homeostasis model assessment of β-cell function (HOMA-β), a marker of insulin secretion, increased with the increase in HOMA-IR.

**Discussion**

In this case report, we describe changes in the serum adiponectin levels and VFA throughout the clinical course from catecholamine crisis to the postoperative follow-up period. To our knowledge, there have been no such reports in the literature. Furthermore, our patient was obese with a marked accumulation of visceral fat. Most patients with pheochromocytoma are lean and rarely develop visceral fat obesity (9), as catecholamines induce lipolysis. The effects of catecholamines are greater in visceral fat than in subcutaneous fat, possibly due to different quantities of β-adrenergic receptors expressed in the adipose tissue (10). However, because of recent lifestyle changes, such as the adoption of fat-rich, high-energy diets and physical inactivity, the number of obese subjects—including those with pheochromocytoma—has increased. Because both catecholamine and visceral fat volume (3) can be regulators of adipokine metabolism, we were very interested in the serum adiponectin levels in our obese patient with pheochromocytoma.

Our patient had a lower serum adiponectin level during the catecholamine crisis than after adrenalectomy. Elenkova et al. similarly reported that adiponectin levels were lower in subjects with pheochromocytoma than in healthy controls, and the levels increased after adrenalectomy despite postoperative weight gain (4). Conversely, Isobe et al. reported that adiponectin levels were significantly higher in patients with pheochromocytoma who dominantly produce noradrenaline than in healthy controls (5). They speculated that noradrenaline might promote adiponectin production, thereby leading to protection against diabetes in this type of pheochromocytoma. However, our patient had a different type of pheochromocytoma that equally produced adrenaline and noradrenaline. Conversely, Bosanska et al. reported that adiponectin levels remained unchanged after adrenalectomy, as the normalization of catecholamines was canceled because of postoperative weight gain (6). Recent in vitro studies have demonstrated that the expression of adiponectin mRNAs is downregulated by β-adrenergic stimuli (11, 12) and that adiponectin release from isolated rat adipocytes was suppressed by high-dose adrenaline (13). Intratumoral hemorrhaging is a precipitating factor of catecholamine crisis (2), and the adrenaline level in our patient actually showed a 22-fold increase from the upper limit of the nor-

![Figure 2](image)

**Figure 2.** Macroscopic examinations of the resected tumor revealed massive necrosis, probably due to ischemia rather than tumor necrosis (A). Hematoxylin and Eosin staining (×100) showed necrotic tissue (N) and viable pheochromocytoma tissue (P). Limited viable tumor tissue revealed no malignant cells (B).
Physical inactivity and overeating in our patient gradually increased the visceral fat volume after adrenalectomy, and his adiponectin level decreased again two years postoperatively. A negative correlation has been shown between serum adiponectin levels and the VFA in healthy people (3, 14). Inflammatory cytokines, including TNF-α and IL-6, might be intermediary factors for this correlation; however, the details of the mechanism remain unclear. Adiponectin is an adipocyte-derived plasma protein with anti-inflammatory, anti-atherogenic, anti-diabetic, and anti-cardiomyopathic properties. Serum adiponectin levels have been found to be reduced in subjects with metabolic syndrome, which is associated with increased risks of cardiovascular disease and diabetes (3). Some reports have shown a high incidence of atherosclerotic complications in patients with pheochromocytoma (15, 16). Furthermore, various types of cardiomyopathies are known to develop frequently in pheochromocytoma (17). These cardiovascular complications in pheochromocytoma may be precipitated by not only catecholamine excess but also hypoadiponectinemia; however, further studies are required.

Plasma adiponectin levels have been reported to be 5-10 μg/mL in healthy subjects (8) and <4 μg/mL in subjects at high risk for diabetes or cardiovascular disease (18, 19). The adiponectin levels in our patient were not markedly low, despite his obesity. However, adiponectin levels vary considerably among individuals, and our patient may have had certain features protecting him from adiponectin decrease. Although the absolute values of adiponectin may not have been pathological in this particular patient, determining whether or not adiponectin levels can be changed by not only the visceral fat volume but also catecholamine status in pheochromocytoma patients would be a meaningful finding.

In this case report, we were unable to measure other adipocytokines, such as leptin and resistin. One report described a reduction in plasma leptin in a patient with pheochromocytoma (20), whereas another report suggested no changes (6). Wocial et al. reported that acute catecholamine excess affects the plasma leptin level, while chronic catecholamine excess does not (21). Further investigation into possible changes in leptin during the clinical course in patients with pheochromocytoma is required.

In conclusion, we described the sequential changes in serum adiponectin levels in a patient with both pheochromocytoma and visceral fat accumulation. The serum adiponectin level was low during catecholamine crisis and increased after adrenalectomy. However, it decreased again two years postoperatively when the visceral fat accumulation increased because of an unhealthy lifestyle. Adiponectin metabolism is
altered by catecholamines and the visceral fat volume in pheochromocytoma, which may affect the pathophysiological condition in this endocrine disease.

The authors state that they have no Conflict of Interest (COI).

Acknowledgement

We thank Dr. Tadashi Nakamura and Mr. Tohru Yoshizumi for the measurement of the visceral fat area, Dr. Norikazu Maeda for the measurement of the serum adiponectin level, and all of the medical staff of Toyonaka Municipal Hospital, particularly those in the intensive care unit, for their helpful support.

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