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**Acute pancreatitis with hypercalcemia caused by primary hyperparathyroidism associated with paraneoplastic syndrome: A case report and review of literature**

Yang L *et al*. Pancreatitis with hyperparathyroidism and paraneoplastic syndrome

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**Abstract**

BACKGROUND

Although acute pancreatitis associated with hyperparathyroidism has occasionally been reported, acute pancreatitis with metabolic encephalopathy caused by hyperparathyroidism combined with paraneoplastic syndrome is an extremely rare entity and poorly described in the literature.

CASE SUMMARY

We present a case of a 56-year-old female with upper abdominal discomfort and intermittent nausea and vomiting for 1 wk, without apparent abdominal pain or bloating, no jaundice and decreased blood pressure at the outset. The patient was ultimately diagnosed with moderately severe acute pancreatitis (according to the revised Atlanta classification of acute pancreatitis) combined with metabolic encephalopathy secondary to hypercalcemia caused by primary hyperparathyroidism associated with paraneoplastic syndrome. After active treatment of acute pancreatitis, massive fluid resuscitation, resection of parathyroid and uterine malignant tumors, neoadjuvant chemotherapy and other treatments, her serum calcium eventually returned to the normal level. The patient was successfully discharged from hospital.

CONCLUSION

This is the first case of acute pancreatitis caused by primary hyperparathyroidism associated with paraneoplastic syndrome.

**Key Words:** Acute pancreatitis; Humoral hypercalcemia; Primary hyperparathyroidism; Paraneoplastic syndrome; Case report

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**Core Tip:** This is the first case of acute pancreatitis caused by primary hyperparathyroidism associated with paraneoplastic syndrome successfully treated with timely surgery and chemotherapy. This further raises concern for women with refractory hypercalcemia and abnormal vaginal bleeding or abdominal symptoms.

**INTRODUCTION**

Acute pancreatitis is an inflammatory disease of the pancreas[1,2]. Although several causes of acute pancreatitis have been described, including toxins or drugs, neoplastic obstruction of the biliary tract, hyperparathyroidism, hypercalcemia, and trauma[3], primary hyperparathyroidism with paraneoplastic syndrome is rare and infrequently reported in the literature. Here we report a patient with acute pancreatitis associated with refractory hypercalcemia, hyperparathyroidism, and paraneoplastic syndrome. Due to refractory hypercalcemia, the diagnosis is difficult. Only by fully understanding the characteristics of such diseases, making an accurate diagnosis, and providing timely treatment can we avoid fatal consequences.

**CASE PRESENTATION**

***Chief complaints***

A 56-year-old woman visited the emergency department of an affiliated hospital with progressive upper abdominal discomfort, intermittent nausea, and abnormal vaginal bleeding, for 1 wk.

***History of present illness***

Initially, there was no apparent jaundice or changes in blood pressure. The patient denied any fever, dizziness, headache, or disturbance of consciousness at the beginning of the illness.

***History of past illness***

The patient had a history of *Helicobacter pylori* infection for more than 10 years. She had no relevant past interventions.

***Personal and family history***

She denied family or psycho-social history.

***Physical examination***

The patient had stable vital signs: body temperature was 37.0 °C, blood pressure was 168/87 mmHg, heart rate was 74 bpm, and respiratory rate was 25 breaths/min, with obvious upper abdominal tenderness, slight rebound pain, and muscle tension.

***Laboratory examinations***

Laboratory examination results were as follows: white blood cell count 26.5 × 109/L, neutrophil percentage 95%, C-reactive protein 109 mg/L, procalcitonin 4.35 ng/mL, serum amylase 1222.5 U/L, creatinine 172.3 µmol/L, urine amylase 1433.0 U/L, platelet 214 × 109/L, hemoglobin 156 g/L, albumin 26.8 g/L, alanine aminotransferase 19.5 U/L, aspartate aminotransferase 26.6 U/L, lactate dehydrogenase 457.3 U/L, ferritin 747 ng/mL, and 25-hydroxyvitamin D < 3 ng/mL. Serum calcium increased significantly to 4.72 mmol/L, with parathyroid hormone (PTH) rising to 366.90 pg/mL and carbohydrate antigen 125 to 142.5 U/mL. An overview of selected initial laboratory values is shown in Table 1.

***Imaging examinations***

Non-contrast enhanced computed tomography (CT) of the abdomen showed exudation around the pancreas (Figure 1A and B). Ultrasound examination of the neck revealed an inferior thyroid nodule and mild hyperplasia of the parathyroid gland. Head magnetic resonance imaging (MRI) was performed, and abnormal signals were found in the bilateral fronto-parietal-temporal-occipital cortex-medullary junction area and bilateral paraventricular area, which were likely due to metabolic encephalopathy related to pancreatitis (Figure 1C and D). A repeat non-contrast enhanced CT scan of the abdomen (Figure 1E and F) and head MRI (Figure 1G and H) showed a reduction in pancreatic exudation and abnormal head signals after effective treatment. Contrast enhanced CT of the neck (Figure 1I and J) showed nodules located at the junction of the left lobe of the thyroid and parathyroid gland. MRI of the pelvis (Figure 1K and L) suggested malignant lesions of the uterus and multiple uterine fibroids.

**MULTIDISCIPLINARY EXPERT CONSULTATION**

Due to the unknown nature of the disease and refractory hypercalcemia, a multidisciplinary expert consultation was conducted, and further workup was proposed. General practitioners believed that refractory hypercalcemia might be caused by parathyroid disease, and exploratory resection of the parathyroid gland was necessary.

**FINAL DIAGNOSIS**

Moderately severe acute pancreatitis with refractory humoral hypercalcemia caused by primary hyperparathyroidism associated with paraneoplastic syndrome.

**TREATMENT**

Based on the above laboratory and imaging examinations (initially we did not consider gynecological disease and did not perform pelvic MRI), the preliminary diagnosis was acute pancreatitis secondary to hypercalcemia caused by primary hyperparathyroidism. She was transferred to the emergency intensive care unit and treated with salmon calcitonin, zoledronic sodium, and anti-infection and intravenous fluids. During the hospitalization period, the patient became more lethargic with dry mucous membranes and mild coma. Early use of imipenem and extensive resuscitation effectively controlled her inflammation. As serum calcium treatment was not satisfactory, hemofiltration was then performed twice. Although acute pancreatitis and consciousness gradually improved, serum calcium did not return to normal level (Figure 2).

To determine the cause of the elevated calcium, a multidisciplinary expert was consulted, and further workup was proposed. Finally, although the nature of the parathyroid lesions was not yet clear, to reduce serum calcium, exploratory parathyroidectomy was performed on April 14, 2020 (Figure 2). The pathological result of the removed tissue suggested nodular goiter.

Unexpectedly, removal of the parathyroid gland did not decrease serum calcium; therefore, we speculated that high calcium might not be caused by hyperparathyroidism alone. To verify this hypothesis, an MRI of the pelvis was performed, and the results showed ambiguous, irregular masses and multiple uterine fibroids in the uterine/cervix region. There were multiple peritoneal and omental masses, and the pelvic and para-aortic lymph nodes were enlarged. The pathological results of the vaginal curettage suggested poorly differentiated uterine cancer (Figure 3). She was immediately transferred to the gynecological ward. She was treated with cytoreductive surgery and neoadjuvant chemotherapy (Figure 2). After surgery, supportive care was provided in the intensive care unit (ICU) and she received 3 cycles of carboplatin and paclitaxel.

**OUTCOME AND FOLLOW-UP**

She did not experience any serious complications or neurologic deficits in the ICU. The patient’s serum calcium eventually decreased to normal levels, and she was successfully discharged from hospital 2 mo later.

**DISCUSSION**

Acute pancreatitis is a common disease associated with significant morbidity and mortality. Alcohol and biliary disease are the causes of almost all such cases, with an incidence of approximately 80%-90%[4]. Additionally, uncommon causes include toxic substances, trauma, infection, autoimmune diseases, or metabolic disorders secondary to hypercalcemia, such as primary hyperparathyroidism or malignant tumors[3]. Generally, acute pancreatitis is associated with a decrease in serum calcium, but pancreatitis caused by primary hyperparathyroidism or malignancies is usually associated with hypercalcemia. These two causes are rare in hypercalcemia. Although the relationship and the pathophysiology are unclear, it is possible that the connection between them is not accidental. Inappropriate activation of digestive enzymes in the pancreas, especially trypsinogen in the acetabulum may play an important role in the development of acute pancreatitis[5,6]. An excessive increase in intracellular calcium concentration can lead to over-activation of digestive enzymes and block the pancreatic ducts, causing inflammatory exudation of the pancreas[7].

The above two rare, but well-known causes of hypercalcemia, are discussed in the literature. Many cases of acute pancreatitis caused by hyperparathyroidism with parathyroid adenoma or adenocarcinoma have been described[8-11]. With regard to paraneoplastic syndrome, several cases have shown that pancreatitis was related to Zollinger-Ellison syndrome[12-14]. Four cases diagnosed as pancreatitis were associated with lung cancer[15-18]. One case had pancreatic adenocarcinoma[19]. One case with myelodysplastic syndromes finally led to pancreatitis[20]. One case had breast cancer[21]. One case was associated with an intraductal papillary neoplasm of the bile duct[22]. Another case of Hodgkin lymphoma with paraneoplastic hypercalcemic pancreatitis was reported[23]. Two similar cases of autoimmune-like pancreatitis with a thymoma and myasthenia gravis were described[24,25]. In addition, two rare cases of pancreatitis and ovary carcinoma were reported, where pancreatitis was caused by the gynecological malignancies[26,27]. An overview of previously described cases of pancreatitis associated with malignant paraneoplastic syndromes is shown in Table 2.

On the one hand, primary hyperparathyroidism is rare, with a documented incidence of 1.5%-8%[8]. Compared with healthy individuals, patients with hyperparathyroidism have an increased risk of acute pancreatitis. The prevalence of pancreatitis among patients with primary hyperparathyroidism is between 1.5% and 13%[28]. Acute pancreatitis caused by hyperparathyroidism can now be diagnosed by the continuous increase in serum calcium and elevated PTH levels[10]. Primary hyperparathyroidism is mainly associated with a solitary parathyroid adenoma (85%-90%), but it is also associated with parathyroid carcinoma (< 1%)[9]. Malignant tumors of the parathyroid gland are usually rare, with an incidence of less than 0.5%[29]. The clinical manifestations are traditionally secondary to hypercalcemia, including non-specific gastrointestinal symptoms and cardiovascular or neuromuscular dysfunction[30,31]. Imaging studies including ultrasound imaging, radionuclide scanning, CT, and MRI have advantages in detecting ectopic parathyroid lesions and are helpful in diagnosis[32]. Surgery is the most effective and only way to treat parathyroid carcinoma, which provides the best chance for cure and long-term survival[33,34].

On the other hand, paraneoplastic hypercalcemia occurs in patients with malignant tumors, with an incidence of about 0.3%-4.0%, which means that the immune system’s abnormal response to normal tissues is either through the production of autoantibodies or T cell attack. Dysfunction of immune regulation and cross-reactivity of the immune system to tumor and normal tissues are involved in the pathogenesis of such reactions[21]. It can occur in different types of malignant tumors, and the most common types are lung cancer, head and neck cancer, renal cancer, or multiple myeloma, but it rarely occurs in gynecological malignancies[35,36]. There are also reports in ovarian cancer, sarcoidosis, and multiple endocrine neoplasia syndrome[37]. Timely surgery and appropriate chemotherapy are necessary[36].

In our patient, examinations initially suggested that moderately severe acute pancreatitis was caused by primary hyperparathyroidism. Firstly, the major etiologies of acute pancreatitis include alcohol consumption and biliary stones, which were not present in this patient. Secondly, she did not have a family history of pancreatitis. In addition, PTH and serum calcium were significantly increased, and treatment did not decrease refractory serum calcium. In addition, ultrasound and contrast enhanced CT of the neck revealed nodules located at the junction of the left lobe of the thyroid and parathyroid gland. Therefore, these findings suggested that the pathogenesis of acute pancreatitis was the result of hypercalcemia caused by hyperparathyroidism. Although no adenoma or adenocarcinoma was found, an exploratory resection was performed to decrease refractory serum calcium. Pathological findings of the removed tissue suggested nodular goiter.

Due to this surprising finding, we conducted a further diagnostic workup and finally found gynecological malignancy. A literature review indicated that some malignant tumors might secrete parathyroid hormone-related protein (PTHrP), instead of PTH, causing pancreatitis. The amino terminus of PTHrP has a similar structure to that of PTH. Both activate PTH/PTHrP receptor 1 and decrease the renal clearance of calcium[36].

According to the results of the above analysis, the final diagnosis was moderately severe acute pancreatitis caused by primary hyperparathyroidism associated with paraneoplastic syndrome. Timely surgery and neoadjuvant chemotherapy were performed which extended the survival time of this patient. She was satisfied with the treatment outcome and was successfully discharged from hospital.

**CONCLUSION**

To our knowledge, this is the first case of moderately severe acute pancreatitis caused by primary hyperparathyroidism associated with paraneoplastic syndrome. Despite its rare occurrence, hypercalcemia secondary to primary hyperparathyroidism or malignancies usually manifests in the advanced stage of malignancy and has a poor prognosis. Early accurate management can avoid fatal consequences and extend survival time. Therefore, more attention should be paid to the differential diagnosis in women with hypercalcemia and abnormal vaginal bleeding or abdominal symptoms.

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**Footnotes**

**Informed consent statement:** Informed consent was obtained from the patient.

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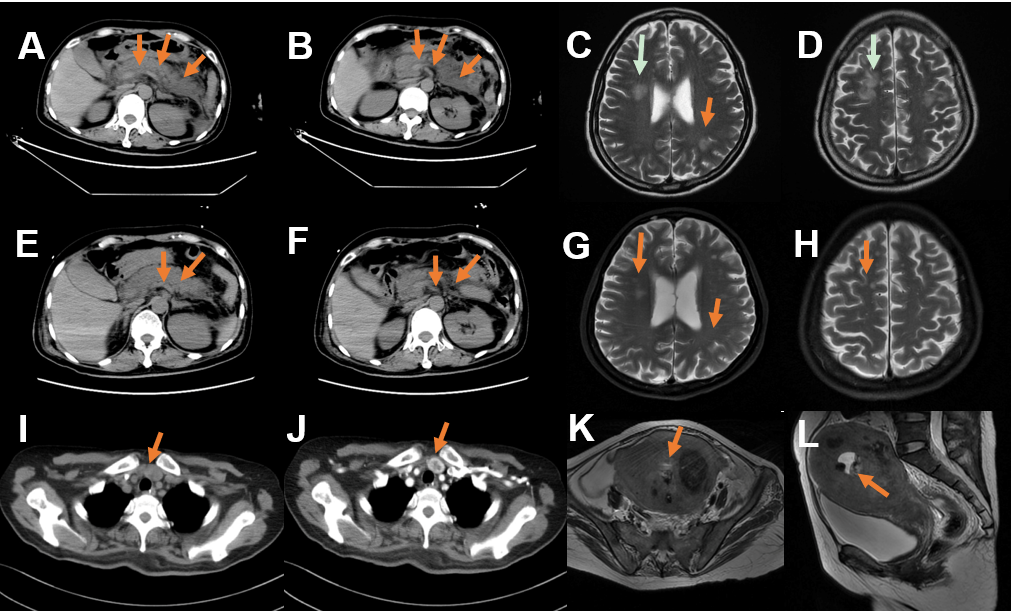
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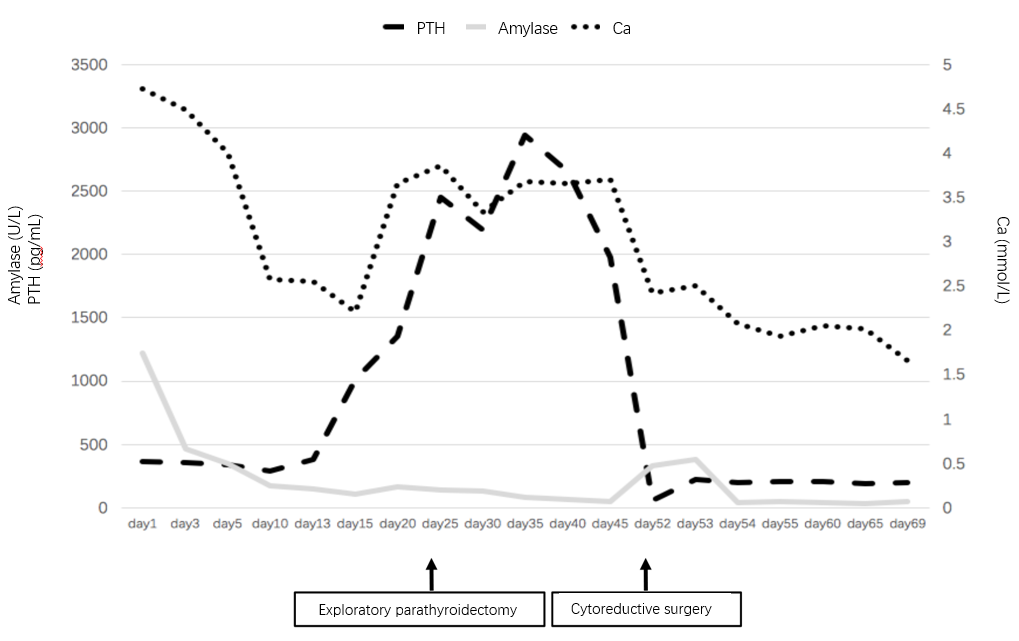
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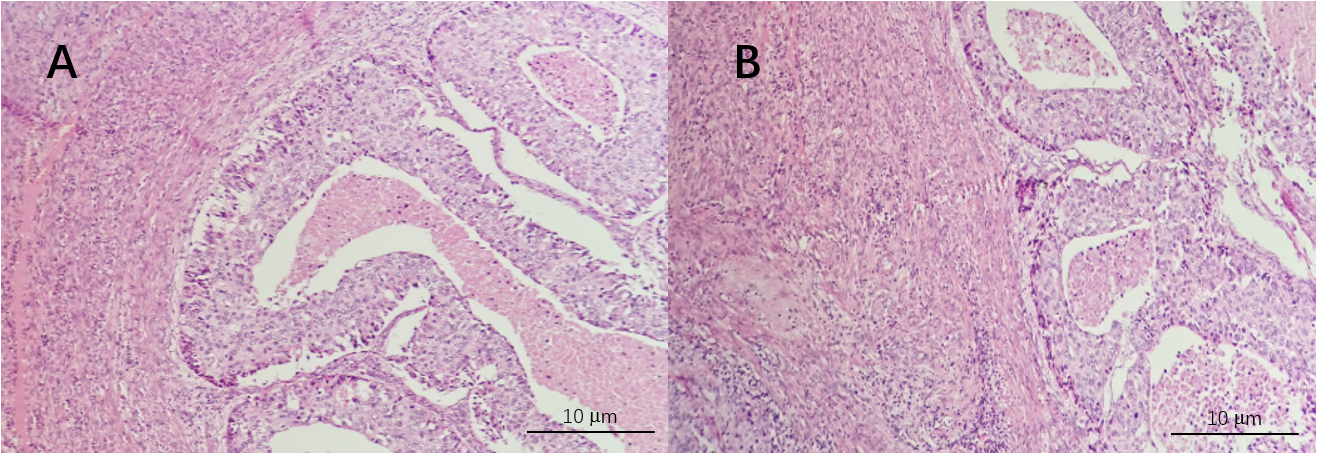
**Figure Legends**



**Figure 1 Computed tomography.** A and B: A non-contrast enhanced computed tomography (CT) scan of the abdomen revealed exudation around the pancreas (arrow); C and D: Head magnetic resonance imaging (MRI) revealed abnormal signal in bilateral fronto-parietal-temporal-occipital cortex-medullary junction area and bilateral paraventricular; E-H: A repeated non-contrast enhanced CT scan of the abdomen and head MRI showed reduction of pancreatic exudation and abnormal head signals after effective treatment; I and J: A contrast enhanced CT of the neck showed nodules located on the junction of left lobe of the thyroid and parathyroid gland; K and L: MRI of the pelvis suggested malignant lesions of the uterus and multiple uterine fibroids.



**Figure 2 The serum concentration of amylase, calcium, and intact parathyroid hormone during the disease course of this patient.** The serum amylase gradually recovered to normal level whereas the intact parathyroid hormone concentration decreased after exploratory parathyroidectomy, which was performed 25 d after admission. Refractory hypercalcemia ultimately decreased to normal level after cytoreductive surgery on May 11. PTH: Parathyroid hormone.



**Figure 3 Pathological diagrams of endometrium.** Hematoxylin and eosin stain, × 200 magnification. A: Poorly differentiated endometrial carcinoma, increased lymphocytes infiltrate the cervical endometrium; B: Endometrioid carcinoma with squamous differentiation is not excepted, cytoplasm with eosinophilic keratinocytes, endometrial glandular differentiation.

**Table 1 Selected initial laboratory values**

|  |  |
| --- | --- |
| WBC (4-10) (× 109/L) | 26.5 |
| Neutrophil percentage (50-70) (%) | 95 |
| PLT (100-300) (× 109/L) | 214 |
| Hb (113-151) (g/L) | 156 |
| Albumin (35-53) (g/L) | 26.8 |
| ALT (4-40) (U/L) | 19.5 |
| AST (4-40) (U/L) | 26.6 |
| CRP (0-10) (mg/L) | 109 |
| PCT (0-0.5) (ng/mL) | 4.35 |
| Serum amylase (0-200) (U/L) | 1222.5 |
| Urine amylase (0-1000) (U/L) | 1433.0 |
| Creatinine (44-97) (mol/L) | 172.3 |
| Serum calcium (2.13-2.65) (mmol/L) | 4.72 |
| PTH (15-65) (pg/mL) | 366.90 |
| CA-125 (0-35) (U/mL) | 142.5 |
| LDH (80-250) (U/L) | 457.3 |
| Ferritin (11-306) (ng/mL) | 747 |
| 25-OH-VD (20-100) (ng/mL) | < 3 |

ALT: Alanine aminotransferase; AST: Aspartate aminotransferase; CA-125: Carbohydrate antigen 125; CRP: C-reactive protein; Hb: Hemoglobin; LDH: Lactate dehydrogenase; PCT: Procalcitonin; PTH: Parathyroid hormone; PLT: Platelet; WBC: White blood cell; 25-OH-VD: 25-hydroxyvitamin D.

**Table 2** **Overview of previously described cases of paraneoplastic syndrome and pancreatitis**

|  |  |  |
| --- | --- | --- |
| **Ref.** | **Cancer** | **Along with increased PTH** |
| Danne *et al*[12], 1985 | Gastrinoma | No |
| Yamamoto *et al*[13], 2003 | Zollinger-Ellison syndrome | No |
| Baffy *et al*[14], 2000 | Zollinger-Ellison syndrome | No |
| Belhassen-García *et al*[15], 2009 | Lung cancer | No |
| Saliba *et al*[16], 2006 | Lung carcinoma | No |
| Akinosoglou *et al*[17], 2014 | Lung cancer | No |
| Casadei Gardini *et al*[18], 2016 | Lung cancer | No |
| Leone *et al*[19], 1998 | Pancreatic adenocarcinoma | No |
| Tanvetyanon *et al*[20], 2005 | Myelodysplastic syndrome | No |
| Lekakis *et al*[21], 2012 | Breast cancer | No |
| Miyazaki *et al*[22], 2019 | Intraductal papillary neoplasm of bile duct | No |
| Mittra and Davidzon[23], 2014 | Hodgkin lymphoma | No |
| Colaut *et al*[24], 2002 | Thymoma | No |
| Tomiyama *et al*[25], 2008 | Thymoma | No |
| Wynn *et al*[26], 2004 | Ovarian carcinoma | No |
| Seifert and Seemann[27], 1967 | Ovarian carcinoma | No |

PTH: Parathyroid hormone.



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