Case Report

Missed Case of Pancreatogenic Diabetes Diagnosed Using Ultrasound

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Abstract

Nonalcoholic pancreatogenic diabetes mellitus (type 3c DM) is an often-misdiagnosed entity usually seen in young men of tropical countries. Although most of the patients present with abdominal pain and symptoms of exocrine pancreatic insufficiency, there is still a subset that does not present with these classical symptoms, which emphasizes the need for special diagnostic considerations. The significance of identifying this subset of diabetic lies not only in the change in management of the disease but also in early detection for pancreatic carcinoma that is more common among those patients. In our case, ultrasound with X-ray played a vital role in diagnosis, prompting us to consider it as an essential part of the investigation panel in all newly diagnosed nonobese diabetic individuals.

Keywords: Fibrocalculous pancreatogenic diabetes, nonalcoholic pancreatogenic diabetes mellitus, tropical pancreatic diabetes

INTRODUCTION

Diabetes mellitus (DM) is a common entity encountered by physicians from all specialties and usually classified as either type 1 or 2. Pancreatic diabetes type 3c DM is also an important subtype that is often misdiagnosed or underdiagnosed.[1] It is characterized by the damage of pancreatic exocrine function due to acute relapsing or chronic pancreatitis (CP) of any etiology that eventually leads to impairment of pancreatic endocrine function causing DM. The occurrence of pancreatic DM is more common than was generally thought. Its early identification is very important to assess the need for some special diagnostic and treatment plan.[2] We present an unusual case of pancreatic DM that was initially diagnosed and treated as type 2 DM for more than 6 years.

CASE REPORT

A 43-year-old nonalcoholic male, on treatment for type 2 diabetes for more than 6 years with oral hypoglycemic drugs, presented with worsening of glycemic control, weight loss (5 kg), and fatigability. However, due to these symptoms, he stopped all his medications about 6 months ago. The patient’s body mass index was 19.5 kg/m² with HbA1c ~ 12. Other investigations done showed that hemoglobin level, total leukocyte count, lipid profile, and renal function test were within normal limits except for mild elevation of liver function test (total bilirubin – 1.7; direct – 0.7; indirect – 1.0, and serum glutamic oxaloacetic transaminase – 54).

Since he was found to be a lean diabetic with no family history of type 2 DM, C-peptide level along with other routine investigations was done. Serum C-peptide at fasting (using chemiluminescence Immunoassay method) was 0.37 ng/ml (normal value: 0.8–3.5 ng/ml), so a possibility of autoimmune DM was thought and anti-insulin and anti-islet cell antibodies were tested which turned out to be negative. No excessive insulin resistance was noted as evidenced by HOMA-IR of 0.4. Further evaluation of pancreatic function revealed normal amylase and lipase levels. Deranged liver function prompted us to do an ultrasound abdomen to rule out the possibility of hepatic or gallbladder pathologies [Figure 1].

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Surprisingly, ultrasound showed atrophic pancreas, multiple tiny hyperechogenic foci with postacoustic shadowing noted in the head and body of the pancreas (suggestive of calcifications). The main pancreatic duct appeared dilated with an intraductal calculus seen [Figure 2]. No peripancreatic fat stranding or fluid collection was seen. Computed tomography (CT) confirmed chronic calcific pancreatitis [Figures 3 and 4].

The expensiveness of fecal elastase for evaluation of the exocrine function of the pancreas precluded the patient from doing it. However, indirect tests of pancreatic functions showed low Vitamin D level (25-hydroxycholecalciferol), i.e., 10.1 ng/ml (normal: 30–100), slightly increased prothrombin time (an indirect measure of Vitamin K) ~14.5 s (normal 10–14), and marginally lower spectrum of triglyceride to high-density lipoprotein ratio ~ 1.1, indicating fat malabsorption.

There was an ambiguity in classifying him as type I or II, as his anti-insulin and anti-islet cell autoantibodies were negative and he also did not have evidence of insulin resistance. Identification of CP by ultrasound made the diagnosis of pancreatogenic DM-type 3c possible. Henceforth, the management protocol was modified with addition of insulin, withdrawing oral hypoglycemic agents. He was also started on a modified low carbohydrate diet with Vitamin D and omega-3 supplements. The patient’s glycemic control improved (HbA1c ~ 6) and weight increased by 5 kg within the next 4 months. Further follow-up after 2 months showed sustained glycemic control and weight gain.

**Discussion**

Pancreatogenic diabetes or type 3c DM refers to diabetes due to impairment in pancreatic endocrine function as a result of pancreatic exocrine damage of various causes such as acute, relapsing, or CP (of any etiology), cystic fibrosis, hemochromatosis, pancreatic cancer, as well as neonatal diabetes due to pancreatic agenesis. Of these, chronic calcific pancreatitis is the most common cause of type 3c DM which is often misdiagnosed or underdiagnosed.[1,2,4]
Patients suffering from pancreatic DM usually present with a history of pancreatitis in the past that causes abdominal pain, dyspepsia, steatorrhea, and symptoms of malabsorption. In our case report, the main reason for missed diagnosis was absence of typical clinical features of pancreatic DM such as intermittent abdominal pain and steatorrhea, but deficiency of fat-soluble vitamins and lower lipid level supported our diagnosis. A study done in Pune,[5] followed up individuals diagnosed with fibrocalculus pancreatic DM for 7 years, concluded that the two pathognomonic complaints (abdominal pain and steatorrhea) were not always present, which was the case in our patient. The biggest challenge in identifying type 3c DM is that there are no universally accepted diagnostic criteria.[6] The commonly known criteria proposed by Ewald and Bretzel[6] have disadvantage in developing countries due to the nonavailability and nonaffordability of recommended laboratory tests.

The risk factors for acquiring chronic calcific pancreatitis are alcohol, smoking, obstructive abnormality, genetic, or autoimmune. In our case, the possible cause could be a genetic or autoimmune pancreatitis. A review of literature says that 10%–30% of patients have no identifiable cause.[7] On the other side, it has been observed that 38% of patients suffering from CP develop diabetes within the next 1–2 years.[8] Hence, it becomes essential to diagnose cases of CP at the earliest to alter the natural course of the disease. However, biochemical studies are not helpful in definitive diagnosis during early stages.[9] A combination of clinical symptoms, pancreatic function tests, and radiological investigations is needed for the early diagnosis of CP in suspected patients.[10] Among the radiological investigation, endoscopic ultrasound with pancreatic function test (with intravenous synthetic secretin) is considered gold standard in the diagnosis of early exocrine insufficiency in CP. However, due to the invasive nature of the tests, CT and magnetic resonance imaging are preferred tests to detect CP at the earliest.[11] However, these tests are very expensive to be used for screening patients without any symptoms suggesting of CP. Ultrasound is a better cost-effective alternative to detect pancreatic atrophy or calcification.[12] Although it is operator dependent, standardization of technique can improve the sensitivity of the test to avoid misdiagnosis of type 3c DM.

Currently, there are no accepted guidelines regarding the treatment for type 3c DM, although this subset of people is known to require insulin. Pancreatic enzyme and fat-soluble vitamin (A, D, E, and K) supplements should be given. Occasionally, surgical treatment may be needed. Most importantly, type 3c DM is distinct from type 1 and type 2 DM, as it results from chronic inflammation of the pancreas with an increased risk of pancreatic carcinoma, thus requiring periodic evaluation of serum CA19-9 levels.[6,13]

**Conclusion**

A simple ultrasound abdomen can play a vital role in diagnosing this small subset of people with type 3c DM. We suggest the need for further research studies to assess the feasibility of using ultrasound with/without X-ray as an essential diagnostic tool for all newly diagnosed nonobese diabetic.

**Declaration of patient consent**

The authors certify that they have obtained all appropriate patient consent forms. In the form, the patient has given his consent for his images and other clinical information to be reported in the journal. The patient understands that name and initials will not be published and due efforts will be made to conceal identity, but anonymity cannot be guaranteed.

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**Conflicts of interest**

There are no conflicts of interest.

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