An unusual presentation of pancreatic neuroendocrine tumour (PNET)

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Insulinoma is a rare pancreatic endocrine tumour derived from β cells that secrete insulin, which results in recurrent hypoglycaemia. Most are benign and solitary. The age range for peak incidence is between 30 and 60 years (median 47 years) and it is more prevalent in women. Diagnosis relies on clinical features along with laboratory tests and imaging. However, non-specific symptoms and the small size of the tumour can lead to misdiagnosis and difficult localisation. Clinical presentation is due to hypoglycaemia secondary to excessive and uncontrolled secretion of insulin and involves neuroglycopaenic (neuronal glucose deprivation, can cause death) and neurogenic (autonomic nervous system discharge) symptoms.

Case report
A 64-year-old woman was admitted to the emergency department (ED) with history of abnormal behaviour (facial grimacing, bouts of crying/shouting, bizarre uncoordinated limb movements) over the past 6 years. She was previously misdiagnosed as dissociative disorder and treated with antipsychotics and electroconvulsive therapy (ECT).

During initial evaluation at ED, random blood glucose was documented to be 27 mg/dL which reversed with intravenous (IV) 25% dextrose bolus. Screening for sulfonylureas was negative. Further evaluation revealed increased levels of fasting serum insulin 28.60 μU/mL (normal, 2.0–25.0) and serum C-peptide 6.08 ng/mL (normal, 0.81–3.85). Triple-phase computed tomography (CT) showed a 15 mm discrete lesion projecting superiorly from the proximal body of the pancreas (Fig 1). However, a 68 Ga DOTA-TATE scan revealed no focal lesion with octreotide receptor expression within the pancreas. She was managed with IV 10% dextrose infusion and taken up for surgical excision of the exophytic pancreatic mass. Following surgery, the patient has been euglycaemic and is off dextrose infusion presently, with no neuroglycopaenic or autonomic symptoms.

Conclusion
Insulinoma remains a diagnostic challenge since symptoms are non-specific and may lead to incorrect diagnosis. As in this case, a psychiatric illness might be wrongly considered as the culprit. Neuropsychiatric symptoms are a common clinical presentation of an insulinoma. A high level of clinical expertise is crucial to avoid misdiagnoses with psychiatric illnesses before insulinoma is recognised. It can be easily confirmed if it fulfils Whipple’s triad, standard endocrine tests and is curable by surgery. On the other hand, severe and sustained hypoglycaemia due to misdiagnosis can lead to disability/death.

Conflict of interest statement
None declared.
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References