A Case of Autoimmune Hypoglycemia in Switzerland

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Abstract

In this report, we describe a white man with symptomatic hypoglycemia whose medical work-up revealed an excessively elevated serum insulin level (867 mIU/l). The need to measure IAA in patients with extraordinarily high measured levels of total insulin is essential because of the possible artifactual elevation (in a conventional insulin immunoassay) caused by IAA. In the present case, the elevated IAA level (>50 kIU/l) confirmed the diagnosis of autoimmune hypoglycemia and allowed to avoid unnecessary surgical intervention.

Keywords: Insulin autoimmune syndrome; Hirata disease; Insulin autoimmune antibody

Introduction

The insulin autoimmune syndrome (IAS), also known as Hirata disease, is a rare cause of hypoglycemia, particularly in non-Asian populations, characterized by spontaneous hypoglycemia, extremely high insulin levels and the presence of circulating native insulin autoimmune antibody (IAA) in patients who have never been exposed to exogenous insulin [1]. These antibodies first bind insulin, forming a complex, and subsequently, regardless of the glucose levels, release it causing hypoglycemia. IAS is considered to be the third greatest cause of hypoglycemia in Japan following insulinoma and extrapancreatic neoplasms and is increasingly being recognized worldwide in non-Asian populations [2]. The cases reported in Japan have shown strong association between IAS and HLA-DR4 status, haplotype 20 times more common in Asians compared with non-Asian subjects, or the use of medications containing sulfhydryl (thiol) group. Cases reported from other parts of the world are more frequently associated with autoimmune disorders and plasma cell dyscrasias [3,4]. The majority of the cases reported outside Asia come from Europe (50%) and the United States (41%). Overall, IAS affects men and women equally and is seen more frequently in patients older than 40 years of age [5].

Case Report

An 88-year-old retired Caucasian man with Hispanic origins, presented an episode of symptomatic hypoglycemia at home. He was found by a family member outside his bed, on the floor, in a confused and agitated state. When the emergency medical service team arrived, they detected a severe hypoglycemia at 1.7 mmol/l (0.31 g/l). After intravenous administration of 20 g of glucose (40 ml of Glucose 50%) the patient normalized his capillary blood glucose value (5.2 mmol/l or 94 mg/dl) and became less agitated. He fully regained consciousness after a couple of minutes but he was experiencing a complete circumstantial amnesia. The patient was taken to Lausanne University Hospital for surveillance and a complete blood test. He was not known as diabetic and was not taking any drug known to cause hypoglycemia. He and his family confirmed that he had not been using exogenous insulin or other antidiabetic drugs. His medical history was significant for autoimmune disorders and plasma cell dyscrasias [3,4]. The majority of the cases reported outside Asia come from Europe (50%) and the United States (41%). Overall, IAS affects men and women equally and is seen more frequently in patients older than 40 years of age [5].

Table 1: Laboratory test results with the most relevant results of the 72 hours fast test

| Laboratory data                  | Patient’s value | Reference range   | 72 hours fast test |
|----------------------------------|-----------------|-------------------|--------------------|
|                                  |                 | At 2 hours        | At 16 hours        | At 70 hours        |
| Glucose (mmol/l)                 | 5               | 3.7-5.6           | 3.3*               | 2.4*               | 2.4*               |
| Insulin level (mIU/l)            | 269             | 3-13              | 867                | >1000              | 520                |
| C-peptide µg/l                  | 10.5            | 1-3               | 4.9                | 6.8                | 3.9                |
| Insulin auto-antibodies (kIU/l)  | >50             | <0.4              |                   |                    |                    |
| IGF II (ng/ml)                   | 561             | 373-1000          |                   |                    |                    |
| Beta-OH-butyl rate µg/l          | 1468            | 58-170            |                   |                    |                    |
| Hba1c (%)                       | 5.5             | 4.9-6.5           |                   |                    |                    |
| TSH (mIU/l)                      | 1.59            | 0.27-4.2          |                   |                    |                    |
| ACTH (ng/l)                      | 9               | 4-30              |                   |                    |                    |
| Cortisol (nmol/l)                | 357             | 210-560           |                   |                    |                    |
| Hemoglobin (g/l)                 | 98              | 133-177           |                   |                    |                    |
| AST (IU/l)                       | 67              | 14-50             |                   |                    |                    |
| ALT (IU/l)                       | 80              | 11-60             |                   |                    |                    |
| Creatinine (µmol/l)              | 80              | 62-106            |                   |                    |                    |

* The patient had no neuroglycopenic symptoms

Abbreviations: ACTH: Adrenocorticotropic Hormone; ALT: Alanine Aminotransferase; AST: Aspartate Aminotransferase; Beta-OH-Butylate: Beta-Hydroxy-Butyrate; C-Peptide: Connecting Peptide; Hba1c: Hemoglobin A1c; IGF II: Insulin-Like Growth Factor II; TSH: Thyroid Stimulating Hormone.

Table 1: Laboratory test results with the most relevant results of the 72 hours fast test.

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resonance imaging (MRI) showed no pituitary mass or other intracranial abnormality except the ancient ischemic stroke lesions. During the first 48 hours following hospitalization, the patient continued to have mild asymptomatic hypoglycemic episodes (3.5 mmol/l or 0.5-0.9 g/l). In this case we initially had a clinical suspicion of endogenous hyperinsulinism, so we performed a 72 hours fast test. In order to have a positive test, we need to objectify the Whipple's triad and an insulin/C-peptide ratio below 1. The Whipple's triad is positive if biochemical hypoglycemia is accompanied by neuroglycopenic symptoms, with disappearance of these symptoms after correction of the hypoglycemia. In order to exclude factitious hypoglycemia, the insulin/C-peptide ratio should be determined at the time point when Whipple's triad is positive; in case of insulinoma the ratio has been reported to be <1. During the 72 hours fast test, the patient developed frequent but asymptomatic hypoglycemia with plasma glucose level varying from 2.4 and 4.9 mmol/l (0.4 and 0.8 g/l). The corresponding serum insulin levels were very high, varying from 269 to 867 mIU/l (reference range of 3-13 mIU/l); the insulin connecting peptide (C-peptide) levels were also high (10.5 μg/l for a reference range of 1-3 μg/l), and plasma beta-hydroxybutyrate was high (1468 μg/l for a reference range of 58-170 μg/l). Because the patient had very high immunoreactive insulin, free insulin was assayed using polyethylene glycol (PEG) precipitation according to the method of Nakagawa et al to rule out endogenous hyperinsulinism [6]. Recovery after precipitation with PEG was <2.6%, result in favor of the presence of IAA. We measured the level of these antibodies: they were elevated >50 kIU/l for a reference range <0.4 kIU/l. These findings were highly suggestive of insulin autoimmunity syndrome (IAS) being the cause of the hypoglycemia. The patient was advised to take frequently meals in small quantities in order to maintain a more consistent blood glucose level and to perform daily monitoring of capillary blood glucose. After the hospital discharge, a glucagon injection kit and additional treatment with steroids were prescribed. Initially at 60 mg daily dose, the oral prednisolone dose was tapered to 5 mg/day after 3 months. During the follow-up, the episodes of hypoglycemia gradually disappeared, a moderate decrease in serum insulin levels was also observed but the level was still high (88 mIU/l). Unfortunately, the patient died after 4 months of follow-up due to an infectious pneumonia.

Discussion

In this report, we describe a white man with symptomatic hypoglycemia whose medical work-up revealed an excessively elevated serum insulin level (867 mIU/l). The need to measure IAA in patients with extraordinarily high measured levels of total insulin is essential because of the possible artificial elevation (in a conventional insulin immunoassay) caused by IAA. In the present case, the elevated IAA level (>50 kIU/l) confirmed the diagnosis of autoimmune hypoglycemia and allowed to avoid unnecessary surgical intervention. Since the patient’s health status could not permit more invasive procedures (e.g. endoscopic ultrasound with pancreatic biopsy, surgery), we did not formally exclude insulinoma, but the recent abdominal CT-scans and the negative 72 hours fast test were not in favor of this diagnosis. Conservative therapy, coupled with immunosuppression, permitted to stabilize patient's blood glucose level and to avoid other episodes of severe hypoglycemia. The supposed mechanism for hypoglycemia in IAS is a mismatch between blood glucose and free insulin concentration secondary to the binding and the release of secreted insulin related to autoimmune antibodies. Following a meal or oral glucose intake, high blood glucose level induces insulin secretion. Autoimmune antibodies bind to these insulin molecules, rendering them unavailable to exert their effects. This binding reduces the initial insulin response, resulting in hyperglycemia and further insulin secretion. As the glucose concentration begins to fall, insulin secretion also decreases but the antibody-bound insulin molecules spontaneously dissociate at this time, resulting in a free insulin level inappropriate for the glucose concentration, evoking hypoglycemia [6,7].

About half of IAS patients report recent exposure to medications, with over 90% of offending agents containing a thiol group such as Methimazole (anti-thyroid agent), Penicillamine, Glutathione, Hydralazine, Procainamide and Isoniazid [4,8]. The subject described in the present report was taking two drugs that contained active thiol metabolites, namely Pantoprazole and Clopidogrel. There are case reports of IAS in patients using thiol-containing proton pump inhibitors including Pantoprazole and Omeprazole [9,10]. The mechanism of development of IAS with thiol group-containing drug is not entirely clear. Different approaches for management of IAS have been described. The first line of treatment is implementing lifestyle modification as small frequent meals with low–glycemic index carbohydrates. Any potentially incriminating medication should be discontinued. Glucocorticoid oral therapy (1 mg/kg/day) can improve glycemic control and diminish the levels of IAA. Other medications such as Acarbose can decrease carbohydrate digestion and absorption. Somatostatin, diazoxide and even partial pancreaticectomy have been employed in a attempt to limit insulin secretion [5,7] None of these therapy has been universally successful. For refractory cases, plasmapheresis or Rituximab have been used with good results [11,12]. Previous studies already showed that in type 1 diabetic patients who developed insulin antibodies to exogenous insulin, Rituximab has been successfully used to suppress the production of insulin antibodies. A single course of Rituximab was enough to obtain a significant level of insulin antibodies suppression that has been noted to last, in a subgroup of patients, up to 3 years despite continuing insulin therapy [13].

Conclusion

Here we report the case of a white man with potentially life threatening hypoglycemia due to IAS. Even though primary investigation of hypoglycemia in non-diabetic patients is always focused on insulinoma, IAA measurement should be always performed because IAS is a cause of hypoglycemia that can be cured without surgery in the majority of cases.

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