Glucokinase activating mutation causing hypoglycaemia diagnosed late in adult who fasts for Ramadhan

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Summary

Activating mutation of glucokinase gene (GCK) causes resetting of insulin inhibition at a lower glucose threshold causing hyperinsulinaemic hypoglycaemia (GCK-HH). This is the first reported case who tolerated years of regular fasting during Ramadhan, presenting only with seizure and syncope now. We describe a case with GCK gene variant p.T65I diagnosed in a 51-year-old woman with hypoglycaemia unawareness even at glucose level of 1.6 mmol/L. Insulin and C-peptide levels during hypoglycaemia were suggestive of hyperinsulinism, but at a day after intravenous glucagon, hypoglycaemia occurred with low insulin and C-peptide levels, pointing against insulinoma as the underlying aetiology. Imaging studies of the pancreas and calcium arterial stimulation venous sampling were unremarkable. A review of old medical records revealed asymptomatic hypoglycaemia years ago. Genetic testing confirmed activating mutation of GCK. Hypoglycaemia was successfully controlled with a somatostatin analogue. This case highlights the importance of consideration of genetic causes of hypoglycaemia in adulthood, especially when imaging is uninformative.

Learning points:

- Consider genetic causes of endogenous hyperinsulinism hypoglycaemia in adulthood, especially when imaging is uninformative.
- Late presentation of activating mutation of GCK can occur because of hypoglycaemia unawareness.
- Long-acting somatostatin analogue may be useful for the treatment of activating mutation of GCK causing hypoglycaemia.
- Depending on the glucose level when the blood was taken, and the threshold of glucose-stimulated insulin release (GSIR), the serum insulin and C-peptide levels may be raised (hyperinsulinaemic) or low (hypoinsulinaemic) in patients with activating mutation of GCK.
- Glucagon may be useful to hasten the process of unmasking the low insulin level during hypoglycaemia below the GSIR level of which insulin released is suppressed.

Background

We present here a unique case of activating mutation of glucokinase gene (GCK) causing hyperinsulinism hypoglycaemia (GCK-HH). This is the first reported case who tolerated decades of regular fasting during Ramadhan, presenting only with seizure and syncope now. Her hypoglycaemia unawareness is severe as she...
remains asymptomatic even with glucose of 1.6 mmol/L, which can partially explain her late diagnosis of GCK-HH. Hypoglycaemia unawareness due to downregulation of counter-regulatory hormone response contributed to her delayed presentation and absence of Whipple’s triad. She had left-sided campodactyly with right hemisphere atrophy, possibly secondary to hypoglycaemia during brain development. The use of intravenous glucagon during the 72 h fast appeared to hasten the process of unmasking the low insulin level during hypoglycaemia below the glucose-stimulated insulin release (GSIR) level of which insulin released is suppressed.

**Case presentation**

A 51-year-old female of Muslim religion was admitted to our hospital after she sustained a right femur fracture following a syncope. On the second day of admission, she had an episode of generalised tonic-clonic seizure. She was found to have multiple asymptomatic hypoglycaemia readings 2.3–3 mmol/L occurring typically in the morning and postprandially (3–3.5 mmol/L). Urine toxicology screens for sulphonylurea and meglitinides were negative.

She had abnormal developmental history. She had campodactyly of her left hand since she was an infant. Although she had below average IQ, she was community-ambulant, had a friendly personality and independent in her daily activities. She was born at full term in a village along with her family, religiously fasted for 14 h during Ramadhan month every year since she was a teenager. She weighed 64 kg, her height was 1.53 m and her BMI was 27 kg/m².

Her blood tests were normal for full blood count, renal panel, liver panel, thyroid function, lipid panel and serum ammonia. A 250 µg synacthen test was normal with a peak cortisol level of 577 mmol/L. She did not have growth hormone deficiency as evident by her normal glucagon stimulation test, achieving a peak of 59.2 MU/L.

A review of her old medical records revealed that she was seen in outpatient clinic 11 years ago for an incidental finding of asymptomatic low venous glucose of 1.9 mmol/L during a general health check. An inpatient fast revealed a nadir glucose of 2.4 mmol/L at 44 h after initiation of fast, with hyperinsulinism hypoglycaemia (venous glucose 2.4 mmol/L, insulin 2.8 mIU/L, C-peptide 430 pmol/L) albeit a borderline insulin level. A CT scan of her abdomen was normal. She was lost to follow-up as she was asymptomatic. In view of the significant time-lapse from her 1st asymptomatic presentation years ago and the absence of Whipple’s triad, an inpatient 72 h fast was repeated (Fig. 1).

At 16 h of the fast, venous glucose of 2.5 mmol/L was reached with C-peptide 348.5 pmol/L and insulin 2.7 mIU/L, suggestive of hyperinsulinism hypoglycaemia, albeit another borderline insulin level. Her venous glucose increased by >1.4 mmol/L from 2.5 to 5.5 mmol/L, indicating she had hepatic glycogen stores from insulin excess at this point (1). In view of the borderline fasting insulin level, fasting was continued with hourly glucose monitoring with the aim to reach venous glucose <2.5 mmol/L. Capillary glucose remained at 2.6–3.5 mmol/L until at 29 h later after the glucagon injection, her venous glucose dropped to 1.6 mmol/L, with undetectable serum insulin levels and low C-peptide levels (insulin <1 mIU, C-peptide 154 pmol/L), Fig. 1. At this point, cortisol and growth hormones were taken which showed suboptimal responses, suggesting downregulation of counter-regulatory hormones in hypoglycaemia unawareness (2).

At 0, 30, and 60 min after the onset of hypoglycaemia 1.6 mmol/L was detected, cortisol levels were 483, 416, 389 mmol/L, GH levels were 0.2, 0.5, and 3.5 MU/L, and IGF-1 levels were 34.4, 32, and 30.9 µg/L (NR 100–314 µg/L), respectively. The suppressed serum ketones during the fast and only slightly increased to 1.1 mmol/L but still much lower than 2.7 mmol/L (1) reflect that ketoacid formation

**Investigation**

- **Insulin level of 0 signifies undetectable insulin level. The lowest detection limit <1 mIU/L**

*Fig. 1 Venous glucose, serum insulin and C-peptide levels at 16th h of 72 h fast (A) and at 29 h after IV glucagon was given illustrated in (B).*

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has initiated but not yet at starvation ketosis state. She remained asymptomatic throughout the whole fast. In view of her borderline insulin levels during hypoglycaemia despite raised C-peptide values, proinsulin level was measured and was raised at 48 pmol when glucose was 2.6 mmol/L (proinsulin NR < 5 pmol/L (1)).

The pancreatic imaging scan using CT and MRI were negative for insulinoma. Calcium arterial stimulation venous sampling of the pancreas did not localise hypersecretion of insulin to any pancreatic region (Fig. 2). Insulin antibodies were negative and IGF-2: IGF-1 ratio was 4.8 (NR < 10). An activating GCK missense variant (NM_000162.5 (GCK): c.194C>T p.[Thr65Ile];[=]) known to be associated with GCK-HH (3) was identified by targeted next-generation sequencing using the Ion Torrent PGM™ (Life Technologies). All protein-coding regions and their flanking splice sites, the 5’UTR and 3’UTR regions, were sequenced for 16 maturity-onset diabetes of the young (MODY)-associated genes, including GCK. Variants identified were annotated employing Alamut® software, classified by American College of Clinical Genetics and Genomics (ACMG) guidelines (4), and those classified as pathogenic/likely pathogenic were verified by bi-directional Sanger sequencing. The rare variant p.Thr65Ile is found in exon 2 of the GCK gene (RefSeqGene NG_008847.2) and resides in the allosteric activator site, which is involved in transforming the enzyme from the active to non-active state. Her mother and brother were tested negative for the genetic mutation. We could not confirm if this was a de novo variant as her father could not be tested. He died from ruptured appendicitis in his 60s, and he was not known to have hypoglycaemia before.

CT and MRI brain showed that she had right cerebral hemisphere atrophy, which was likely a developmental abnormality (Fig. 3). This could explain her left hand and left foot camptodactyly. Silver-Russell syndrome was also considered in view of the camptodactyly and reduced IQ. However, she had normal head circumference and asymmetry of limbs could not be ascertained in view of femur fracture and camptodactyly. Genetic tests for Silver-Russell syndrome were negative; methylation studies of 11p15 and chromosomal microarray analysis were normal.

**Treatment**

Management of hypoglycaemia was challenging initially. She had hypoglycaemia despite intravenous dextrose and frequent meals. Her hypoglycaemia was not controlled with frequent meals. She had severe muscle stiffness, painful joints and leg oedema after three doses of diazoxide 100 mg BD. While the oedema of her legs was tolerable, the generalised muscle stiffness and painful hand joints prohibited her movements. The adverse events resolved on stopping diazoxide. On rechallenging diazoxide 1 month

![Figure 2](http://creativecommons.org/licenses/by-nc-nd/4.0/)

**Figure 2**

Calcium arterial stimulation venous sampling showed that the rise of insulin (A) and C-peptide (B) were not localised to any particular pancreatic region, although the highest rise was in the region supplied by the splenic artery. A significant rise was taken as >two times elevation from baseline, with the symbol × denoting the number of multiplication from baseline.

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later, the same adverse events occurred after two doses, and hence diazoxide was stopped. Anti-epileptic valproate was temporarily switched to phenytoin for hyperglycaemic effect (5), but this did not help. Monthly injections of somatostatin analogue Sandostatin LAR 20 mg ameliorated hypoglycaemia <3.5 mmol/L as seen using continuous glucose monitoring. This was switched to Lanreotide LAR 120 mg for cost reasons with a good response. Her HbA1c improved from <4.3% (i.e. lower than detection limit) to 4.9% on treatment.

**Outcome and follow-up**

Four years on, her hypoglycaemia remains well controlled on somatostatin analogue, with the lowest glucose of 3.2–3.3 mmol/L occasionally.

**Discussion**

Clinically and biochemically, it can be difficult to distinguish between insulinoma and CHI but critically important to do so because the management is quite different. The biochemical criteria from 72 h fast to diagnose insulinoma are defined by glucose <3 mmol/L, insulin ≥3 mIU/L (18 pmol/L), and C-peptide >200 pmol/L (1). These criteria do not differentiate between insulinoma and GCK-HH, unless hypoglycaemia was below the glucose-stimulated insulin release (GSIR) threshold in GCK-HH. Thus, GCK-HH may have either positive or even negative asymptomatic 72 h fast (2). We found that continuing to monitor glucose and insulin levels after a glucagon injection was useful to detect hypoinsulinaemic hypoglycaemia in GCK-HH. The glucagon-induced glycogenolysis may have hasten the unmasking of her GSIR level at which endogenous hyperinsulinism is suppressed. Unlike GCK-HH, insulinoma would persistently have hyperinsulinism due to autonomous production regardless of venous glucose level or glycogen status. Glucokinase is a major controller that promotes liver glycogen synthesis (6). Perhaps, the metabolic adaptation of GCK-HH controlling glycogenesis and glycogenolysis at a lower GSIR, coupled with hypoglycaemia unawareness explains why this patient with GCK-HH could withstand fasting during Ramadan for many years, while patients with insulinoma are unable to (7). Our observation suggests there is a risk of hypoglycaemia post-glucagon, especially in those with lower glycogen reserves.

Understanding GCK-HH may provide mechanistic insight to the use of glucokinase activators (GKA) as a treatment for type 2 diabetes. Non-selective GKA studies were disappointing in terms of rapid decline of hypoglycaemia efficacy and causing hepatic steatosis (6). Hepatoselective GKA TTP399 showed promising results of glucose-lowering effect and yet without causing hepatic steatosis (8). Unlike non-selective GKA, which increased glucagon and insulin levels, the liver-selective GKA TTP399 reduced plasma glucagon without change of insulin levels (8). Since monogenic diabetes due to inactivating GCK gene mutation (MODY2) have increased glycaemic threshold for glucagon secretion, activating-mutation of GCK may have decreased glucagon counterregulatory response during hypoglycaemia (2, 9). Lower glucagon response may thus be protective yet also contributing to hypoglycaemia unawareness. Further studies are needed to understand the role of glucagon in GCK-HH and GKA.

CHI, otherwise called hyperinsulinism in infancy or nesidioblastosis, is most often diagnosed at infancy because of the early symptoms of poor feeding, floppiness, and jitteriness (10). Among the 11 different genes associated with CHI, GCK-HH is less common than ABCC8 and KCNJ11 (11). About 20 GCK-HH mutations have been reported with the majority being missense mutations in 95% (11). GCK-HH presents with a wide spectrum of phenotypic features even within the same family with respect to presentation, age, severity, treatment response and hypoglycaemia awareness (2, 6, 12, 13). Although commonly presenting as neonates, late diagnosis has rarely been reported via cascade screening (6) with the oldest patient being 77 years old. Affected family members have been identified because of the autosomal dominant inheritance (2, 6). Unlike other causes of CHI, GCK-HH lowers the threshold for GSIR and usually responds poorly to the suppressive effect of insulin release by diazoxide (14). Whilst somatostatin analogue experience is limited and not effective for some patients (6), this patient presented herein had an excellent clinical response.
The absence of family history does not exclude GCK-HH because mutations could be de novo and results in a variable spectrum of symptoms (11, 13). Patients presenting with de novo mutation tend to be unresponsive to diazoxide, have macrosomia and less adult onset (11).

CHI causes neonatal-infantile hypoglycaemia leading to permanent neurodevelopmental deficits such as motor and speech deficits, epilepsy, and mental retardation in 20–50% of survivors (10, 15). Specific regions of brain injury associated with neonatal hypoglycaemia on MRI are the white matter of posterior regions, cortex, basal ganglia and pulvinar injury, depending on the age of symptomatic hypoglycaemia (16, 17, 18). Although mental retardation and speech deficits could be explained by hypoglycaemia, the asymmetrical hemisphere damage, in this case, was atypical. Generalised cerebral atrophy on MRI were found in 22.5% of patients with CHI but none with localised cerebral atrophy (19). We postulate that in this case, the neurological deficit causing hemisphere agenesis, camptodactyly, mental retardation and epilepsy were caused by hypoglycaemia during the critical brain developmental phase of neonatal-infantile period. Neonatal hypoglycaemia itself is an important cause of hypoxic-ischaemic encephalopathy (16). Whilst expectedly, the MRI changes of hypoglycaemia-induced injuries are usually symmetrical (16, 17), asymmetrical brain damages have been described in a case report (18). A 5 years old male with hypoglycaemia due to glycogen storage disease had right temporal lobe seizure causing status epilepticus and MRI brain showed atrophy of right hemisphere 1 month later (18). Asymmetrical changes of basal ganglia causing hemichorea–hemiballismus secondary to hyperglycaemia have been commonly described (20), also highlighting that biochemical glucose derangements can lead to asymmetrical brain damages.

In conclusion, GCK-HH should be considered as a differential diagnosis in an adult with hypoglycaemia, especially when imaging is uninformative. Genetic testing in such cases is invaluable as our case demonstrated that pancreatectomy could be avoided with the use of somatostatin analogue. The use of glucagon test during 72hr fast may be useful. Repeating insulin, C-peptide and proinsulin levels, if normal, are worthwhile in patients with suspected GCK-HH.

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**Patient’s perspective**

The patient and family were satisfied with the treatment and outcome because there were less need for multiple feeding throughout the day and night, less worry of hypoglycaemia during sleep and hypoglycaemia-related complications.

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20 Oh SH, Lee KY, Im JH & Lee MS. Chorea associated with non-ketotic hyperglycaemia and hyperintensity basal ganglia lesion on T1-weighted brain MRI study: a meta-analysis of 53 cases including four present cases. Journal of the Neurological Sciences 2002 200 57–62. (https://doi.org/10.1016/s0022-510x(02)00133-8)

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