

Hypoglycaemia (PIP)

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The following guidance is taken from the Partners In Paediatrics (PIP)

Date	Key Amendment	Approved by:
9 th March 2024	Document reviewed and amended:	Paediatric Governance
	 Flow chart (Differential diagnosis SI not detected) Glucose infusion rate calculator (link) 	Meeting



Hypoglycaemia 2018-20

HYPOGLYCAEMIA

Management of unexplained and prolonged hypoglycaemia

RECOGNITION AND ASSESSMENT

Definition

Blood glucose <2.6 mmol/L in child aged >1 month

Symptoms and signs

- Lethargy
- Tremulousness
- Loss of consciousness
- Seizure
- Autonomic effects
- sweating
- shaking
- tachycardia
- anxiety
- hunger

Previous history

- Ask about:
- antenatal history e.g. small-for-dates, gestational diabetes
- prematurity
- history of neonatal hypoglycaemia
- early or prolonged jaundice
- family history of sudden infant death
- development, especially developmental regression
- medication (steroids)
- access to glycopaenic agents (e.g. metformin, insulin)
- onset and frequency of hypoglycaemia
- history of infection/food intake

Investigations

Certain pointers to cause of unexplained hypoglycaemia are detectable only during episode. Take blood samples BEFORE correcting blood glucose

Immediate samples

- Before treating, take blood samples (Table 1)
- Bloods must arrive in laboratory within 30 min
- Include clear clinical details on request form
- If sample volumes limited prioritise glucose, insulin and C-peptide
- Request urgent analysis of insulin and C-peptide (discuss with duty biochemist)
- routine analysis for obese child with insulin resistance
- Blood ketones for ketone bodies on ward (glucometer)
- Once samples obtained, correct hypoglycaemia. See Immediate treatment
- Collect first urine voided after correction. Check for ketones using urine dipstick, send remaining urine for organic/amino acid metabolites and reducing substances

Table 1: Total blood requirement (5 mL minimum)

Fluoride (grey top)	1.3 mL (1 bottle)	Glucose, lactate, beta-
		hydroxybutyrate, free fatty acids
Lithium heparin (green	2.6 mL (2 bottles –	U&Es, LFTs, blood amino acids,
top)	1 bottle on ice)	Acylcarnitines, ammonia



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Clotted (red top)	2.6 mL (2 bottles)	Insulin, C-peptide, growth hormone,	
		cortisol	ĺ

Investigations

- In all prolonged unexplained hypoglycaemia:
- glucose point of care
- ketones. Urine dipstick or blood ketones (glucometer)
- capillary blood gas
- laboratory glucose to confirm hypoglycaemia
- insulin
- C-peptide
- U&E
- growth hormone
- cortisol
- 17-hydroxyprogesterone in infant if hyponatraemia present

- if urgent analysis required contact duty biochemist

- Further investigations may be required, depending on results from above:
- IGF-1
- beta-hydroxybutyrate
- free fatty acids
- carnitines
- urinary reducing substances
- urine organic acids
- urine and plasma amino acids

Physical examination

- Height and weight
- Midline defects, micropenis, optic nerve hypoplasia (pituitary disorder)
- Dysmorphic features: macroglossia, macrosomia, ear lobe crease (Beckwith-Wiedemann)
- Skin hyperpigmentation (adrenal insufficiency)
- Hepatomegaly (glycogen storage disorder)

Differential diagnosis





Algorithm: Ketones present (blood/urine)



IMMEDIATE TREATMENT





SUBSEQUENT MANAGEMENT



To calculate the amount of mg glucose/kg/min:

% glucose x 10 x mL/hr 60 x wt (kg)

Glucose infusion rate calculator: <u>https://www.pediatriconcall.com/calculators/glucose-infusion-rate-gir-</u> calculator