



REGIONAL PEDIATRIC PARENTERAL DRUG MONOGRAPH

GENERIC NAME

glucagon

Effective Date: Dec 2011 CLASSIFICATION OTHER NAMES PAGE

Revised Date: Dec 2022 Hyperglycemic Agent

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ADMINISTRATION POLICY:

IV infusion — Administration restricted to nurses experienced in ED/CARDIAC ROOM/ICU/PACU

IV bolus - May be administered by a nurse
 IM injection - May be administered by a nurse
 Subcutaneous - May be administered by a nurse

RECONSTITUTION/DILUTION/ADMINISTRATION:

Available as: 1 mg glucagon powder with 1 mL diluent to prepare a 1 mg/mL solution

Dilute with entire contents of prefilled syringe of diluent or 1 mL sterile water or D5W. Swirl gently until dissolved. Solution should be clear with a water-like consistency.

IV Bolus: Administer undiluted over 1-5 minutes. Flush tubing with D5W before and after glucagon.

IM/ Subcut: Administer undiluted

IV Infusion: Pump Library:

NOTE: If 100 mL D5W unavailable:

Use 250 mL D5W IV bag, Remove 150 mL from bag and discard; then add 10 mg (10 mL of 1 mg/mL glucagon)

Drug Library	Dose Rate	Short Name	Care Unit
Yes	mg/h	gluca10p	Pediatric
Drug	Diluent	Final Volume (VTBI)	Final Concentration
Drug 10 mg (10 mL of 1 mg/mL)	Diluent 100 mL D5W	Final Volume (VTBI) 110 mL	Final Concentration 0.091 mg/mL

Clinical Advisory:

Soft Low Dose Limit: 0.3 mg/h

Soft High Dose Limit: 5 mg/h

Maximum rate: 5 mg/hour **Maximum concentration:** 1 mg/mL

DOSAGE:

Hypoglycemic episode

IV/IM/Subcutaneous: 0.02 – 0.03 mg/kg per dose (maximum 1 mg per dose) given once

and followed by administration of Dextrose IV or PO to prevent secondary hypoglycemia

If no response in 15 to 20 min, repeat dose

Usual maximum: 3 doses

Beta Blocker or Calcium Blocker Overdose:

IV Bolus: 0.05 mg/kg x 1. If no response, may repeat dose

IV infusion: 1-5 mg/hour; titrate every 30 minutes to desired heart rate and blood pressure

Anaphylaxis: For patients on beta blockers not responding to epinephrine:

IV Bolus: 20 - 30 mcg/kg (0.02-0.03mg/kg) maximum 1 mg IV over 5 minutes, may repeat dose x1

IV infusion: 0.3 - 0.9 mg/hour





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STABILITY/COMPATIBILITY:

Stability of reconstituted vial: Use immediately after reconstitution; discard any unused portion.

Stability of Final Admixture: 24 hours at room temperature

Compatibility: Compatible with D5W

Incompatible with normal saline

PRECAUTIONS, POTENTIAL ADVERSE REACTIONS:

- Nausea, vomiting (may be prevented by pre-treating with prochlorperazine)
- Hypokalemia, hyperglycemia and hypersensitivity reactions
- Transient hypertension and heart rate elevation, hypotension and with rapid administration of large doses
- Hypersensitivity to urticaria, respiratory distress, hypotension
- CV: Transient hypertension and tachycardia. Hypotension with rapid administration of large doses (greater than 1 mg)
- GI: Nausea and vomiting (action: position patient in lateral position prior to dose; consider pre-medication with antiemetic)
- Metab: Hypokalemia, hyperglycemia. Paradoxical hypoglycemia after abrupt discontinuation of IV infusion (action: continue to infuse dextrose)
- Caution in patients with insulinoma (glucagon increases insulin release) or pheochromocytoma (glucagon increases catecholamine release from tumor)
- Hypoglycemia secondary to starvation, adrenal insufficiency or chronic hypoglycemia (insufficient glycogen stores)
- Acute or chronic alcohol ingestion (decreased response due to glycogen depletion or decreased gluconeogenesis)
- Contraindicated in hypersensitivity to glucagon or any component

ADDITIONAL NOTES AND NURSING CONSIDERATIONS:

- In hypoglycemic events, if a patient fails to respond to glucagons, give IV dextrose
- Administer carbohydrates and protein containing foods as soon as possible to prevent recurrence
- Use caution in patients with a history of pheochromocytoma and insulinoma
- Canadian product contains no phenol
- Required monitoring

Blood pressure, HR: Baseline and every 15 minutes x 2, then as required for indication Blood glucose: Baseline and every 30 minutes x 2, then as required for dose and indication

Serum potassium: Baseline and repeat 30 minutes after dose, then as required for dose and indication