# Physostigmine



Short acting IV acetylcholinesterase inhibitor useful in Rx of anticholinergic delirium. Avoid in patients with seizures or evidence of cardiac toxicity.

**Physostigmine** is the only IV acetylcholinesterase inhibitor that crosses the blood-brain barrier

#### **Indications**

- Patients with anticholinergic delirium, where urinary retention has been excluded, and behavioural disturbance is not controlled with benzodiazepines
- In cases where anticholinergic poisoning is suspected, administration of physostigmine with resolution of delirium may negate the need for further investigations (e.g., including CT brain / lumbar puncture)

#### **Contraindications**

- Seizures occurring as part of presentation
- Heart rate < 60 bpm
- QRS duration > 120 milliseconds
- Any evidence of AV block on ECG
- Bronchospasm / history of brittle asthma

#### Presentation

-1 vial of 5 mL = 2mg

## Dose and Administration (discuss use with a clinical toxicologist)

- Administration requires cardiac monitoring and access to full resuscitative care
- dilute 2 mg ampoule in 100 mL of normal saline
- infuse at 100 mL / hour
- cease infusion once therapeutic endpoint (resolution of delirium and / or control of behavioural disturbance) is achieved to avoid cholinergic toxicity
- Maximum dose over an hour = 2 mg (child: 0.02 mg/kg up to 0.5 mg)
- Further dosing may be required after 2 hours as physostigmine is short-acting
- Consider alternative diagnosis if no clinical effect is observed after 2 mg has been administered

## Therapeutic Endpoint:

- desired clinical effect is achieved i.e., resolution of delirium and / or control of behavioural disturbance

### Adverse effects: (usually occur if doses are given too frequently or rapidly)

- Cholinergic toxicity: peripheral muscarinic effects (hypersecretion, **bronchospasm, bradycardia**, nausea, vomiting), peripheral nicotinic effects (neuromuscular weakness) and CNS effects (**seizures**)