Pancuronium
Newborn use only

2021

Alert	High-risk medicine: High risk of causing significant patient harm when used in error. This drug should be administered in the presence of personnel trained in advanced airway management. Suggest regular cessation of infusion for a few to several hours, possibly every 24 hours (commonly referred to as 'drug holiday'[7]) to assess the need for continued paralysis and adequacy of sedation or analgesia. Line should be adequately flushed to avoid unintended paralysis during later use of the line.
Indication	1. Skeletal muscle relaxation or paralysis in mechanically ventilated infants 2. For elective endotracheal intubation.
Action	Long acting non-depolarising muscle relaxant that competitively antagonises acetylcholine antagonist at nicotinic acetylcholine receptors at neuromuscular junctions. Also has autonomic anticholinergic effect resulting in increase in heart rate. Onset of action: 1–2 minutes. Duration of action: 45–60 minutes.
Drug type	Long acting non-depolarising neuromuscular blocking agent.
Trade name	Pancuronium Bromide Injection BP – Astra Zeneca Unregistered SAS products are available
Presentation	4 mg/2 mL ampoule.
Dose	**Muscle relaxation** IV bolus: 100 microgram/kg (50-100 microgram/kg) followed by intermittent IV boluses 50 microgram/kg (50-100 microgram/kg) every 1–2 hours as needed. **Intubation** IV bolus: 100 microgram/kg.
Dose adjustment	Therapeutic hypothermia (TH) –Definite dose adjustment is not yet clear. Dose is to be adjusted to the effect. ECMO –Definite dose adjustment is not yet clear. Dose is to be adjusted to the effect. Renal impairment- Prolonged duration of blocking effect.(MIMS) Hepatic impairment – Effect variable. Adjust the dose to the effect.(MIMS)
Maximum dose	IV bolus: 100 microgram/kg/dose.
Total cumulative dose	
Route	IV
Preparation	Draw up 2 mL (4000 microgram pancuronium) and add 6 mL water for injection to make a final volume of 8 mL with a final concentration of 500 microgram/mL
Administration	IV bolus: Rapid injection over several seconds. Line should be adequately flushed upon cessation of treatment to avoid unintended paralysis during later use of the same line.
Monitoring	Continuous cardio-respiratory and pulse oximetry monitoring. Close monitoring of neuromuscular function, sedation and blood pressure (invasive or non-invasive) is essential. Fluid balance is essential due to of risk of fluid retention. Hepatic and renal function with prolonged use.
Contraindications	Known hypersensitivity to pancuronium bromide or to the bromide ion.
Precautions	Avoid prolonged usage. Suggest regular cessation of infusion, possibly every 24 hours (commonly referred to as 'drug holiday') to assess the need for continued paralysis and adequacy of sedation or analgesia. Pre-existing tachycardia, hypertension (including that associated with renal failure or phaeochromocytoma)—consider an alternative agent. **Renal:** Prolonged neuromuscular blockade may occur in renal impairment; reduction in maintenance dose may be necessary. **Hepatic:** Increased onset time and prolonged duration of action may occur in impairment; consider using alternative agent. Myasthenia gravis—prolongs paralysis; avoid neuromuscular blocking agents if possible. Neuromuscular diseases (e.g. dystrophia myotonica, history of polio), severe obesity—unpredictable effect; use cautiously and monitor neuromuscular function closely.

	Neonates are generally more sensitive to non-depolarising neuromuscular blocking agents; duration of action may be prolonged; monitor neuromuscular function closely. Acidosis, dehydration, hypokalaemia, hypermagnesaemia, hypocalcaemia—enhances effects of neuromuscular blocking drugs; where possible correct before administration, reduce dose and monitor neuromuscular blockade. Hypothermia—decreases effect of pancuronium (unlike the rest of the neuromuscular blockers); reduce dose and monitor neuromuscular blockade. Anaphylactic reaction to neuromuscular blocking agents—allergic cross-reactivity has been reported; refer to specialist for skin testing for sensitivity to other neuromuscular blockers.
Drug interactions	<u>Drugs that POTENTIATE the effect of pancuronium:14</u> Amlodipine, Atenolol, carvedilol, diazepam, diltiazem, doxycycline, fentanyl, furosemide, gentamicin, hydrochlorothiazide, ketamine, ketoconazole, lignocaine (high dose), magnesium sulphate, metoprolol, metronidazole, miconazole, minocycline, nifedipine, nimodipine neomycin, phenytoin, piperacillin, polymixins, propranolol, protamine, suxamethonium, thiamine (high dose), thiopentone, verapamil <u>Drugs that DECREASE the effect of pancuronium</u> Adrenaline (Epinephrine), azathioprine, calcium chloride, edrophonium, hydrocortisone, neostigmine, potassium chloride, prednisone, sodium chloride, theophylline (high doses) <u>Other</u> Risk of developing arrhythmias increased when Pancuronium is used with cardiac glycosides: Digoxin
Adverse reactions	**Respiratory:** May result in prolonged apnoea or respiratory depression. **Cardiovascular:** After administration, approximately 10% of patients may exhibit mild to moderate increases in blood pressure and/or pulse rate. Dysrhythmias may occasionally occur and increased cardiac output is frequently noted. **Hypersensitivity:** Hypersensitivity reactions occur rarely (< 1%). Bradycardia, bronchospasm, hypotension and cardiovascular collapse have been reported. An occasional transient rash has been reported. Pruritus can occur, as well as rare cases of flushing, oedema and wheezing. **Skin:** A few case reports of local reactions including pain and burning at the site of injection. **Ocular:** Pancuronium decreases intraocular pressure and induces miosis. **Neuromuscular:** Prolonged paralysis, disuse atrophy and areflexia have been reported with prolonged use of pancuronium. **Other:** Hypersalivation may occur, especially if no anticholinergic premedication is given.
Compatibility	**Fluids**: Glucose 5%, glucose 5% in sodium chloride 0.9%, glucose 5% in sodium chloride 0.45%, sodium chloride 0.9%.[10] **Y-site** : Aciclovir, amikacin, aminophylline, amiodarone, amphotericin B liposome, ampicillin, atenolol, azithromycin, aztreonam, calcium chloride, calcium gluconate, cefazolin, cefepime, cefotaxime, cefoxitin, ceftazidime, ceftriaxone, cefuroxime, ciprofloxacin, chloramphenicol, clindamycin, dexamethasone, dexmedetomitidine, digoxin, diltiazem, dobutamine, dopamine, doxycycline, epinephrine, erythromycin lactobionate, fentanyl, fluconazole, fluorouracil, ganciclovir, gentamicin, glycopyrrolate, heparin, hydralazine, hydrocortisone, imipenem-cilastin, insulin, regular, ketamine, lidocaine, linezolid, lorazepam, magnesium sulfate, Meropenem, methylprednisolone sodium succinate, metronidazole, midazolam, milrinone, morphine sulfate, naloxone, nitroglycerin, nitroprusside sodium, norepinephrine, octreotide, pamidronate, pentobarbital, phenobarbital, piperacillin, piperacillin-tazobactam, potassium acetate, potassium chloride, potassium phosphates, propranolol, remifentanil, sodium acetate, sodium bicarbonate, sodium phosphates, sulfamethoxazole-trimethoprim, theophylline, ticarcillin-clavulanate, tobramycin, vancomycin, verapamil, zidovudine.[10]
Incompatibility	**Fluids** : No information **Y site** : Amphotericin B conventional colloidal, amphotericin B lipid complex, diazepam, furosemide, pantoprazole, phenytoin, thiopental.[10]
Stability	Dilutions are stable for 48 hours.[9] The stability can be extended if refrigerated. Pancuronium stored at room temperature (15–30°C) will maintain its full clinical potency for 6 months. However, if refrigerated (2–8°C), it will be stable for up to 3 years or until its expiration date, whichever comes first.
Storage	Store at 2–8°C. Do not freeze. Refrigeration is unnecessary during normal periods of use.
Excipients	Sodium chloride, sodium acetate, water for injections, acetic acid, sodium hydroxide.[20]

Special comments	Dose should be individualised for each patient as there is wide variation in individual response. Inhalation agents or prior administration of suxamethonium enhance the action of pancuronium. Therapeutic: It is recommended that a peripheral nerve stimulator be used to monitor response to pancuronium to minimise the risk of overdose.
Evidence	**Efficacy** Muscle relaxation The routine use of pancuronium or any other neuromuscular blocking agent in ventilated newborn infants cannot be recommended. However, for ventilated preterm infants with evidence of asynchronous respiratory effort, neuromuscular paralysis with pancuronium seems to have a favourable effect on intraventricular haemorrhage [RR (95% CI) 0.55 (0.34, 0.89)] and possibly on pneumothorax. However, uncertainty remains regarding the long-term pulmonary and neurological effects and the safety of prolonged use of pancuronium in ventilated newborn infants.[2] (LOEI, GOR B) Intubation Thirty infants with birth weights from 580 to 3450 g (25 to 40 weeks gestation) were prospectively studied during nasotracheal intubation. The infants were randomised to receive atropine 0.01 mg/kg, atropine 10 microgram/kg plus pancuronium 100 microgram/kg or no medication (controls) prior to intubation. Pancuronium plus atropine was associated with lesser increases in intracranial pressure and with the least changes in heart rate in response to intubation.[1] (LOEII, GOR C) The dose used in RCTs for neonatal neuromuscular block in mechanically ventilated neonates is 30 microgram/kg to 100 microgram/kg.[2] There is one study reporting on use of pancuronium infusion for muscle relaxation in ventilated newborn infants with dose range 30–70 microgram/kg/hour.[8] (LOE IV GOR C) **Drug holidays** (i.e. stopping neuromuscular blocking agents until forced to restart based on the patient's condition) may decrease the incidence of post-paralytic quadriparesis.[7,18] (LOE IV GOR D) **Pharmacokinetics** Duration of action is approximately 45 to 60 minutes.[11] An average duration of action is 42 minutes following mean doses of intravenous pancuronium of 2.7 mg.[11] Following a single 50 microgram/kg intravenous pancuronium dose, the 50% recovery time was 37 minutes.[11]. Peak onset of action is at 2–3 minutes.[12] Divided doses of pancuronium may be advantageous in providing rapid, intense paralysis.[13] Pancuronium has been associated with haemodynamic effects (e.g. tachycardia, hypertension) due to blockade of cholinergic receptors outside the neuromuscular junction.[6] Recovery time after paralysis with continuous infusion is faster than that after intermittent bolus injection.[7] A prospective, open-label study conducted in 25 children receiving continuous infusions of pancuronium in ICU showed increased infusion requirement for patients requiring > 5 days treatment or for those receiving concomitant anticonvulsant therapy.[8] **Dose adjustment:** While there is evidence that hypothermia and ECMO have an impact on pharmacokinetic and pharmacodynamics properties of neuromuscular blocking agents, no definite adjusted dose regimen can be recommended and the dose should be titrated to the desired clinical effect.[19] **Safety** Prolonged administration of pancuronium during the neonatal period is associated with sensorineural hearing loss in childhood survivors of CDH.[4] Pancuronium has been associated with prolonged paralysis and muscle atrophy after 1 week when given as intermittent doses or by continuous infusion.[5] In premature infants, pancuronium has also been associated with joint contractures, specifically in the hips and knees.[6] However, this effect does not appear to persist after discontinuation of the drug and resumption of spontaneous activity.[6] Newborn infants paralysed with pancuronium, despite fluid restriction, had evidence of fluid retention and were significantly heavier that the control infants from day 3 onwards and above their birth weight by day 7. Strict attention to fluid retention is essential when newborns are treated with pancuronium.[17] (LOE III GOR C)
Practice points	
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VERSION/NUMBER	DATE
Original 1.0	10/04/2017
Current 2.0	15/07/2021
REVIEW (5 years)	15/07/2026

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