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Date	July, 2022		
Purpose	To ensure a consistent procedural approach to ondansetron administration.		
Scope	Applies to all Queensland Ambulance Service (QAS) clinical staff.		
Health care setting	Pre-hospital assessment and treatment.		
Population	Applies to all ages unless specifically mentioned.		
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# Ondansetron

July, 2022

# **Drug class**

Anti-emetic – 5-HT3 antagonist [1,2]

# **Pharmacology**

Ondansetron is a serotonin 5-HT3 receptor antagonist. It works by blocking the action of serotonin, a natural substance that may cause nausea and vomiting. [1,2]

## Metabolism

The majority of circulating ondansetron is metabolised by the liver and excreted by the kidneys.<sup>[1]</sup>

#### Indications

• Significant nausea AND/OR vomiting

## Contraindications

- **Absolute** contraindication:
  - Allergy AND/OR Adverse Drug Reaction
  - Congenital long QT syndrome
  - Current apomorphine therapy (used in severe Parkinson's disease)
  - Patients less than 2 years of age
- **Relative** contraindication:
  - First trimester pregnancy (may only be administered for extreme and uncontrolled hyperemesis)[3]

#### Precautions

- Hepatic impairment
- Elderly patients
- Intestinal obstruction
- Patients with risk factors for QT interval prolongation or cardiac arrhythmias.

#### Adverse events [1]

## Common (> 1%)

- Headache
- Constipation

#### Rare (< 0.1%)

- Hypersensitivity reactions (including anaphylaxis)
- ECG changes

## Presentation

- Ampoule, 4 mg/2 mL ondansetron
- Orally Disintegrating Tablet (ODT), 4 mg ondansetron

Onset (IV)	Duration (IV)	Half-life
5 minutes	Several hours	3-4 hours

## **Schedule**

• S4 (Restricted drugs).

#### Routes of administration

Per oral (PO)



Intramuscular injection (IM)



Intravenous injection (IV)



#### Special notes

- Ambulance officers must only administer medications for the listed indications and dosing range. Any consideration for treatment outside the listed scope of practice requires mandatory approval through the *QAS Clinical Consultation and Advice Line*.
- Ondansetron ampoules should be protected from light.
- Under no circumstances is an IV cannula to be inserted for the sole purpose of ondansetron administration. Unless contraindicated, ODT ondansetron should always be the preferred option.
- Transient adverse effects have been reported with rapid intravenous injections.
- All cannulae and IV lines must be flushed thoroughly with sodium chloride 0.9% following each medication administration.

## Adult dosages<sup>[1-5]</sup>

Significant nausea AND/OR vomiting					
ACP <sup>2</sup> CCP	PO/IM	4–8 mg Total maximum dose 8 mg. Must not be given within 8 hours of previous ondansetron administration.			
ACP2 CCP	IV	4–8 mg Slow push over 2–3 minutes. Total maximum dose 8 mg.			
		Must not be given within 8 hours of previous ondansetron administration.			



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# Paediatric dosages [1,2,4,5]

# Significant nausea AND/OR vomiting



PO

 $\geq$  5 years – 4 mg Single dose only.

2-4 years - 2 mg Single dose only.

Must not be given within 8 hours of previous ondansetron administration.



IM

≥ 2 years – 100 microg/kg (rounded up to the nearest 5 kg)

Single dose only, not to exceed 4 mg.

Must not be given within 8 hours of previous ondansetron administration.

Weight	Dose	Volume
> 15 - 20 kg	2 mg	1 mL
> 20 - 25 kg	2.5 mg	1.25 mL
> 25 – 30 kg	3 mg	1.5 mL
> 30 - 35 kg	3.5 mg	1.75 mL
> 35 kg	4 mg	2 mL



IV

≥ 2 years – 100 microg/kg

Slow push over 2-3 minutes.

Single dose only, not to exceed 4 mg.

Must not be given within 8 hours of previous ondansetron administration.



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