



Policy code	DTP_LOR_0722	
Date	July, 2022	
Purpose	To ensure a consistent procedural approach to loratadine 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.	
Source of funding	Internal – 100%	
Author	Clinical Quality & Patient Safety Unit, QAS	
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Loratadine

July, 2022

Drug class

Antihistamine (less sedating) [1]

Pharmacology

Loratadine is a long-acting, second generation peripheral histamine H1-receptor antagonistic used to treat allergies.^[1]

Metabolism

Absorbed in the gastrointestinal tract with a rapid first-pass hepatic metabolism.^[1]

Indications

Symptomatic urticaria
 (without evidence of anaphylaxis)

Contraindications

- Allergy AND/OR Adverse Drug Reaction
- Anaphylaxis
- Patients less than 8 years of age

Precautions

- Severe hepatic impairment
- Increased risk of sedation and anticholinergic effects in older people

Side effects

- Drowsiness
- Fatigue
- Headache
- Nausea
- Dry mouth

Presentation

• Tablet, 10 mg loratadine

Onset (PO)	Duration (PO)	Half-life
1–2 hours	≈ 24 hours	≈ 8 hours

Schedule

• S2 (Therapeutic poisons).

Routes of administration

Per oral (PO)



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
 via the QAS Clinical Consultation and Advice Line.
- May be administered with or without food.
- Antihistamines have no role in the treatment or prevention of respiratory or cardiovascular symptoms in anaphylaxis.

Adult dosages [1]

Symptomatic urticaria (without evidence of anaphylaxis)



PO

10 mg
Single dose only.

Not to be administered within 24 hours of previous antihistamine administration unless approval gained from the QAS Clinical Consultation and Advice Line.

Paediatric dosages [1]

Symptomatic urticaria (without evidence of anaphylaxis)



PO

 \geq 8 years – 10 mg Single dose only.

Not to be administered within 24 hours of previous antihistamine administration unless approval gained from the QAS Clinical Consultation and Advice Line.

Moter QAS officers are **NOT** authorised to administer loratadine to paediatric patients presenting with symptomatic urticarial under the age of 8.