



Adrenaline (Epinephrine) Device Prescription

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The aim of these guidelines is to outline the appropriate prescription and dose of adrenaline (epinephrine) devices for use in non-medical settings, for the emergency first aid treatment of potentially life-threatening severe allergic reactions (anaphylaxis). These guidelines should be used in conjunction with clinical judgement, with consideration of issues that include training on a new device and preference of device from the patient or their carer.

These guidelines include adrenaline devices that are currently available in Australia and New Zealand, and are based on published evidence regarding:

- Recommended doses of adrenaline devices to treat severe symptoms, reduce the risk of serious morbidity (e.g. cerebrovascular damage) and prevent fatalities due to anaphylaxis.
- Epidemiology of allergic reactions triggers that may be difficult to avoid.
- Risk factors that have been associated with fatal anaphylaxis.

ASCIA Adrenaline Device Recommendations

Adrenaline rapidly reverses the effects of anaphylaxis and adrenaline devices are the first line emergency first aid treatment for anaphylaxis. Each adrenaline device contains a single fixed dose of adrenaline that is administered intramuscularly into the outer mid-thigh, for safe and rapid absorption of adrenaline. Adrenaline ampoules and syringes are not considered by ASCIA to be suitable for non-medical settings such as schools, children's education/care centres and workplaces.

ASCIA recommends the use of adrenaline as the first line treatment for anaphylaxis using either of the following two brands of adrenaline devices:

- **EpiPen® Jr (150 microgram) and EpiPen® (300 microgram)** - TGA approved, available in Australia on the PBS (up to two devices per prescription), and New Zealand (Pharmac listed 2023).
- **Anapen® 500 (500 microgram)** - TGA approved and available in Australia on the PBS (up to two devices per prescription).

It is important to specify brand and tick box on PBS prescription to ensure that brand is not substituted. EpiPen® and Anapen® are also available without prescription. Both EpiPen® and Anapen® devices are widely used in other countries. Most countries have multiple brands of adrenaline devices available, and this is important for the following reasons:

- To ensure continued supply of life saving adrenaline, particularly if one brand has stock shortages.
- To provide doctors with a choice of dose, who may prefer to prescribe a higher dose (500 microgram device) for people over 50kg.
- A 500 microgram dose can potentially prevent the need for further doses of adrenaline (which is important due to increasing ambulance delays and many people only carrying one device).
- To encourage suppliers to provide devices with longer shelf life.
- To provide choice for consumers to access devices with points of difference to best suit their needs.

It is essential that patients and carers are trained to use the prescribed device (EpiPen® or Anapen®) and provided with an ASCIA Action Plan for Anaphylaxis, including the prescribed device instructions.

ASCIA Adrenaline Device Dose Recommendations

ASCIA recommends the adrenaline device doses listed below, based on expert consensus and standard practice by ASCIA members, which vary to the product information.

These doses are consistent with the Acute Anaphylaxis Clinical Care Standard for Australia, Australian Prescriber Anaphylaxis wallchart, Australian Immunisation Handbook and international recommendations from the World Allergy Organisation (WAO), Canada and the UK (refer to weblinks below).

Children 7.5-20kg (aged around one to five years): EpiPen® Jr (150 microgram)

Children over 20kg and adults: EpiPen® (300 microgram)

Children and adults over 50kg: Anapen® 500 (500 microgram)*

*The dose of adrenaline in Anapen 500 is consistent with the intramuscular injection (IMI) dose recommendations for people who weigh more than 50kg in the publications listed below:

<https://www.safetyandquality.gov.au/standards/clinical-care-standards/acute-anaphylaxis-clinical-care-standard>

<https://www.nps.org.au/australian-prescriber/articles/anaphylaxis-emergency-management-for-health-professionals>

<https://immunisationhandbook.health.gov.au/resources/handbook-tables/doses-of-intramuscular-11000-adrenaline-for-anaphylaxis>

[https://www.worldallergyorganizationjournal.org/article/S1939-4551\(20\)30375-6/fulltext](https://www.worldallergyorganizationjournal.org/article/S1939-4551(20)30375-6/fulltext)

[https://www.resuscitationjournal.com/article/S0300-9572\(21\)00150-7/fulltext](https://www.resuscitationjournal.com/article/S0300-9572(21)00150-7/fulltext)

<https://www.canada.ca/en/public-health/services/publications/healthy-living/canadian-immunization-guide-part-2-vaccine-safety/page-4-early-vaccine-reactions-including-anaphylaxis.html#shr-pg0>

<https://onlinelibrary.wiley.com/doi/full/10.1111/cea.14055>

Children less than 7.5kg

Adrenaline devices are not usually recommended for children less than 7.5kg as the risk of fatal anaphylaxis in children this age is very low. While the 'optimal' dose of adrenaline is unknown there is a risk that the lower dose adrenaline device could provide a significant overdose.

ASCIA does not recommend the use of adrenaline ampoules and syringes for children less than 7.5kg in non-medical settings, such as childcare centres. Even if ampoules are injected by 'trained' non-medical personnel such as parents, there may be a risk of a serious dosing error.

If there is a concern regarding an infant under 7.5kg requiring adrenaline, they should be referred to a clinical immunology/allergy specialist for assessment. The issues should be discussed with the parents based on a risk assessment.

Where it is felt that it is essential for emergency adrenaline to be prescribed for a child less than 7.5kg, the risk of administering a 'fixed' overdose via a 150 microgram device is considered to be lower than the risk of a dose error with an adrenaline ampoule and syringe.

ASCIA Adrenaline Device Prescription Guide

Adrenaline device prescription is recommended for patients with:

- **History of anaphylaxis** - If the patient is considered to be at continuing risk from allergic reactions to identified triggers (confirmed allergen/s) or unidentified triggers (idiopathic anaphylaxis).
- **Food allergy (excluding oral allergy syndrome) and co-existing unstable or moderate to severe, persistent asthma.** Rationale: Most food allergy related fatalities occur in those with unstable asthma.
- **Underlying mast cell disorders** (e.g. systemic mastocytosis or elevated baseline serum tryptase concentrations) together with any previous systemic allergic reactions to insect stings, including patients undergoing venom immunotherapy.

Adrenaline device prescription is sometimes recommended for patients with a history of a generalised allergic reaction, with one or more of the additional risk factors listed below.

Age:

- **Teenagers and young adults with food allergy.** While food allergy is most common in young children aged five years or less, most recorded fatal reactions to foods occur in teenagers and young adults. This may in part relate to greater risk taking behaviour in this age group, but may also reflect greater likelihood of accidental exposure to food allergens when eating away from home, or while not under parental supervision.

Specific allergic triggers:

- **Peanut, tree nuts and seafood.** Fatal anaphylaxis may arise from any food, but most fatalities arise from food allergy that persists into adolescence and adult life (e.g. peanut, tree nut, sesame seed and seafood allergies). Allergic reactions to these foods may occur following ingestion of relatively small amounts, and the risk of reaction is unlikely to be reduced by cooking or food processing.
- **Generalised urticaria alone without anaphylaxis following insect stings** (e.g. bee, wasp or Jack Jumper ant stings) or following tick bites is not a routine indication for adrenaline device prescription but may be considered (in conjunction with allergen specific immunotherapy if available) in selected cases. Decisions regarding immunotherapy will take into consideration factors such as the risk of progression to anaphylaxis (based on follow-up studies), patient age (more likely in adults than children), co-morbidity (significant cardiorespiratory disease) or living or working in remote areas (where access to emergency medical care may be more problematic) or occupational (e.g. bee keeping) or even recreational exposure to stinging insects (e.g. hiking in areas where Jack Jumper ants are endemic).

Co-morbid conditions:

- **Asthma.** Unstable or moderate to severe, persistent asthma increases the risk of respiratory compromise in those allergic to food. Treatment to control asthma symptoms is important in this group (e.g. medication, allergen immunotherapy).
- **Cardiovascular disease** (hypertension, ischaemic heart disease or arrhythmia) is associated with a relatively greater risk of fatal anaphylaxis from insect stings.
- **Systemic Mastocytosis**

This list is not comprehensive and if there is a concern, patients should be referred to a clinical immunology/allergy specialist for assessment.

Limited access to emergency medical care:

- **Remote residential locations.** In some remote residential locations (e.g. remote rural areas), access to medical care and early administration of adrenaline may not be possible unless an adrenaline device is provided to the patient or their carers for administration. It is important to distinguish this situation of permanent risk from those at short-lived risk (e.g. bushwalking, school camps).
- **Prolonged travel abroad.** Consideration of temporary availability to patients considered at lower risk, who are travelling abroad may also be considered, where language barriers and lesser control over food preparation may increase the risk of accidental exposure and access to medical care may also be limited.

These factors should be considered when deciding whether an adrenaline device is prescribed, as they are known risk factors for more severe or fatal allergic reactions.

Adrenaline device prescription is not normally recommended for patients with the following:

- **Asthma without a history of anaphylaxis or generalised allergic reactions.**
- **If known allergen can be successfully avoided**, such as drug (medication) allergy. Under these circumstances the wearing of medical identification jewellery is strongly recommended.
- **Oral allergy syndrome** - The likelihood of progression to anaphylaxis is very low.
- **Elevated specific IgE only (positive blood or skin allergy test) without a history of clinical reactivity** - A positive allergy test without a history strongly suggestive of allergy is an indication for specialist allergy assessment, which will include assessment of the risk of allergy and anaphylaxis, and sometimes include challenge testing.
- **Family (rather than personal) history of anaphylaxis or allergy** - Whilst the risk for allergic disease such as asthma, allergic rhinitis and atopic eczema is in part inherited, there is not a substantial genetic contribution to food, insect sting or drug (medication) allergy risk, and the risk of anaphylaxis is not inherited.
- **Local reactions to insect stings in adults and children** - Follow up studies demonstrate that these rarely progress to anaphylaxis.
- **Generalised skin rash (only) to bee or wasp stings in children** - Follow up studies of subsequent bee stings in children presenting with local reactions or generalised skin rash (only) show that these children are at a very low risk of experiencing anaphylaxis with subsequent stings.
- **Resolved food allergy** - This should be established by a clinical immunology/allergy specialist.
- **Isolated angioedema** - The risk of fatal angioedema (unrelated to food or insect sting allergy, hereditary angioedema or use of ACE inhibitor medication) is very low. If there is a concern, patients should be referred to a clinical immunology/allergy specialist for assessment.

To access clinical definitions, refer to the ASCIA Guidelines for Acute Management of Anaphylaxis on the ASCIA website. www.allergy.org.au/hp/papers/acute-management-of-anaphylaxis-guidelines

Updated ASCIA Action Plans for Anaphylaxis, e-training courses and other resources which include EpiPen® and Anapen® instructions are available on the ASCIA website.

www.allergy.org.au/hp/anaphylaxis

Anaphylaxis Management Plan

An adrenaline device should only be prescribed within the context of a comprehensive Anaphylaxis Management Plan that includes the following:

1. Referral to a clinical immunology/allergy specialist

Review by a clinical immunology/allergy specialist should occur to:

- Ascertain if the correct trigger(s) have been identified.
- Determine whether the allergy persists.
- Advise on specific management, including suitability for allergen immunotherapy (if available).
- Advise on co-factors that may increase the risk of more serious reactions (e.g. use of beta blockers, NSAID, exercise, asthma control).

2. Identification of anaphylaxis trigger(s)

This should include a comprehensive history, clinical examination, appropriate use and interpretation of allergy testing and under some circumstances, deliberate challenge to prove or disprove allergy.

3. Education on the avoidance of trigger(s)

This is particularly important with food allergy induced anaphylaxis.

4. Provision of an ASCIA Action Plan for Anaphylaxis (emergency response plan)

This plan includes:

- Personal details - patient's photo, name, confirmed allergens and family/emergency contact.
- Symptoms and signs indicating when to use the adrenaline device.
- Instructions on how to use the adrenaline device.

5. Appropriate follow-up

Review by a patient's medical practitioner (normally their GP) approximately 12-18 monthly should occur to:

- Review any allergic reactions that have occurred since their last review.
- Examine co-factors (such as poorly controlled or persistent asthma) that may increase the risk of more serious reactions.
- Prescribe adrenaline devices. If you do not want brand substitution, it is important to specify brand and tick box on PBS prescription.
- Provide re-education on adrenaline device use (using a trainer device of the prescribed brand) and positioning (e.g. not standing or walking when experiencing anaphylaxis).
- Renew ASCIA Action Plan for Anaphylaxis - the general version includes instructions for EpiPen® and Anapen®.
- Provide ASCIA Travel Plan if required.
- Determine if specialist review is required to ascertain if the allergy persists, new allergies have developed or if more detailed review is required.

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For more information go to www.allergy.org.au/hp/anaphylaxis

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