

PATIENT ID LABEL

Procedure:

Hospital where reaction occurred:

Date of reaction: Time of induction (24 hour clock):

Time reaction first noted: Date of referral:

Type of anaesthesia: General Regional Local IV sedation

The patient was exposed to the following medications PRIOR to the reaction (indicate time of exposure):

Agent Administered	Time	Agent Administered	Time

Please tick if the patient was exposed to the agents listed below (indicate time of exposure): Time

<input type="checkbox"/> Chlorhexidine <input type="checkbox"/> Wipes <input type="checkbox"/> Skin prep <input type="checkbox"/> Other (specify):	
<input type="checkbox"/> Skin preparation Type:	
<input type="checkbox"/> Latex <input type="checkbox"/> Gloves <input type="checkbox"/> Other (specify):	
<input type="checkbox"/> Contrast agent Type:	
<input type="checkbox"/> Methylene Blue <input type="checkbox"/> Patent Blue	
<input type="checkbox"/> Colloid Type:	
<input type="checkbox"/> Blood products Type:	
<input type="checkbox"/> Antibiotics Type:	
<input type="checkbox"/> Central venous line <input type="checkbox"/> Chlorhexidine coated <input type="checkbox"/> Antibiotic coated <input type="checkbox"/> Other:	
<input type="checkbox"/> Vaginal packing Type:	
<input type="checkbox"/> Urinary catheter Type:	
<input type="checkbox"/> Lubricant Type:	
Other	

BINDING MARGIN – NO WRITING

PATIENT ID LABEL

BARCODE AREA

BINDING MARGIN – NO WRITING

Symptoms and Signs of Reaction

Tachycardia >100 bpm: Yes No
(before adrenaline administered)

Bradycardia < 60 bpm: Yes No

Arrhythmia: Yes No Type:

Cardiac arrest: Yes No

Hypotension: Yes No Time with systolic < 60 mmHg _____ mins

Cough: Yes No

Bronchospasm: Yes No
 Mild wheeze Dyspnoea reported by patient
 Moderate wheeze Difficult to ventilate
 Severe wheeze Very difficult to ventilate

Low oxygen saturations: Yes No SpO₂ 80–90 SpO₂ < 80

Flushing/erythema: Yes No Localised or Generalised

Urticaria: Yes No Localised or Generalised

Piloerection: Yes No

Angioedema: Yes No

Swelling: Yes No Site:
Duration:

Other cutaneous signs: Yes No Specify:

Gastrointestinal signs: Yes No Nausea Vomiting
 Abdominal cramps/pain
 Other

What was the first symptom you noticed?

What was the predominant symptom?

Comments:

Blank lines for writing comments.

ANESTHETIC ALLERGY TESTING REFERRAL FORM

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Details of Treatment

Airway management

Assisted/mechanical ventilation: Yes No Planned Unplanned

Endotracheal intubation: Yes No Before onset After onset

Bronchospasm treatment? Yes No

Specify agent/s used and dose:

Adrenaline given? Yes No IV IM SC ETT

Total dose administeredmcg

IV fluids given to resuscitation? Yes No

Specify type/s of fluid and total volume:

Cardiac compressions? Yes No How long was CPR performed mins

Cardioversion/defibrillation: Yes No Number of shocks

Vasopressors other than adrenaline given? Yes No

- Ephedrine Dose mg Metaraminol Dose mg
- Vasopressin Dose mg Phenylephrine Dose mg
- Noradrenaline Dose mg Methylene Blue Dose mg
- Other (specify):

Steroids given? Yes No

Specify steroid use and dose:

Antihistamines used? Yes No

Specify antihistamine used and dose:

Did you use the ANZAAG Anaphylaxis Management Resource? Yes No

Please comment on any ways in which you think the resource was helpful or could be improved:

Other treatments/comments:

BINDING MARGIN – NO WRITING

Investigations

Serum tryptase taken? Yes No

Recommended to take 10ml samples at the following times after reaction:

1-2 hours Result mcg/L 4 hours Result mcg/L

> 24 hours Result mcg/L

Where possible please attach results to this referral

Which pathology laboratory were the specimens sent to?

Is there a differential diagnosis other than anaphylaxis that you think may have caused the reaction?

Comments:

Outcome/Sequelae

Operation/procedure completed or Operation/procedure abandoned

Patient transferred to PACU/recovery? Yes No

Was patient admitted to hospital? Yes No Tick if admission unplanned

Postoperative care in ICU/HDU: Yes No

If yes: Was patient still intubated/ventilated or transfer? Yes No Duration

Was an inotrope infusion continued? Yes No Duration

How long was the patient in ICU?

Were there any further complications?

ECG changes Coagulopathy Troponin rise Pneumothorax Anxiety/PTSD

Other:

Severity of Allergic Reaction

Please specify the grade of allergic reaction from the categories below:

Grade I – Cutaneous-mucous signs: Erythema, urticaria with or without angioedema.

Grade II – Moderate multivisceral signs: Cutaneous-mucous signs +/- hypotension +/- tachycardia +/- dyspnoea +/- gastrointestinal disturbance.

Grade III – Life-threatening mono- or multivisceral signs: Cardiovascular collapse, tachycardia or bradycardia +/- cardiac dysrhythmia +/- bronchospasm +/- cutaneous-mucous signs +/- gastrointestinal disturbance.

Grade IV – Cardiac arrest.

