



By the end of this lecture you will be able to:

- Perceive the differences between anaphylactic shock and other types of shock
- Recognize its nature, causes & characteristics.
- Specify its diagnostic features
- Identify its standard emergency management protocol
- Justify the mechanism of action and method of administration of each of the different used drugs to limit its morbid outcomes

ANAPHYLAXSIS

Is a sudden, severe allergic reaction affecting the whole body symptoms including:

- > Rash
- Mucosal swelling
- Difficulty breathing
- Reduced blood pressure



ANAPHYLACTIC SHOCK

A life-threatening allergic reaction that causes shock (hypoperfusion) and airway swelling

What TYPE of shock is it ???



Generalized circulatory derangement causing multiple organ HYPOPERFUSION [Inadequate oxygen delivery to meet metabolic demands] & strong sympathetic activation

⇒ when intense or sustained enough, irreversible derangements sets ⇒ permanent functional deficit or death

- Hypovolemic Haemorrhage / fluid loss (plasma, ECF)
- **Cardiogenic**Inability to contract & pump→ myocardial infarction
- **Ubstructive**Extracardiac obstruction → Pul. embolism, cardiac tamponade
- Distributive
 → PR → septic shock, neurogenic, anaphylactic shock

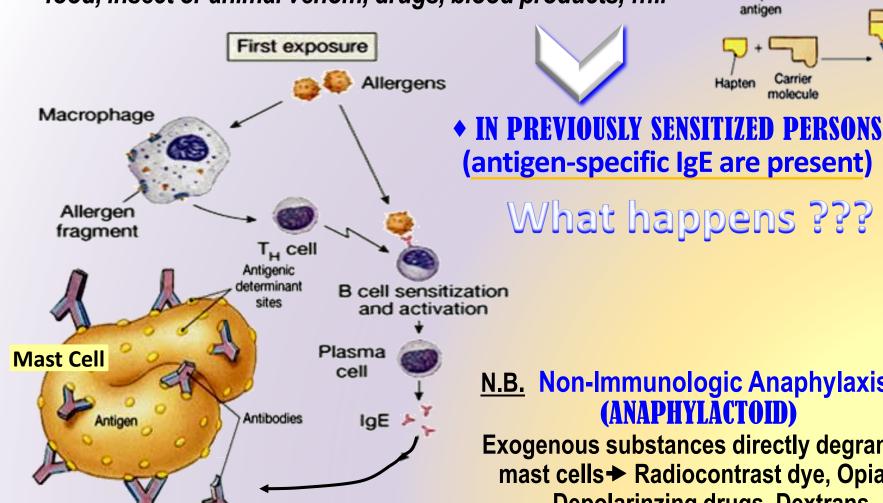
Severe, life-threatening, generalized or systemic hypersensitivity reaction in response to allergen

ANAPHYLACTIC SHOCK

Nature

Belong to TYPE I HYPERSENSITIVITY REACTION

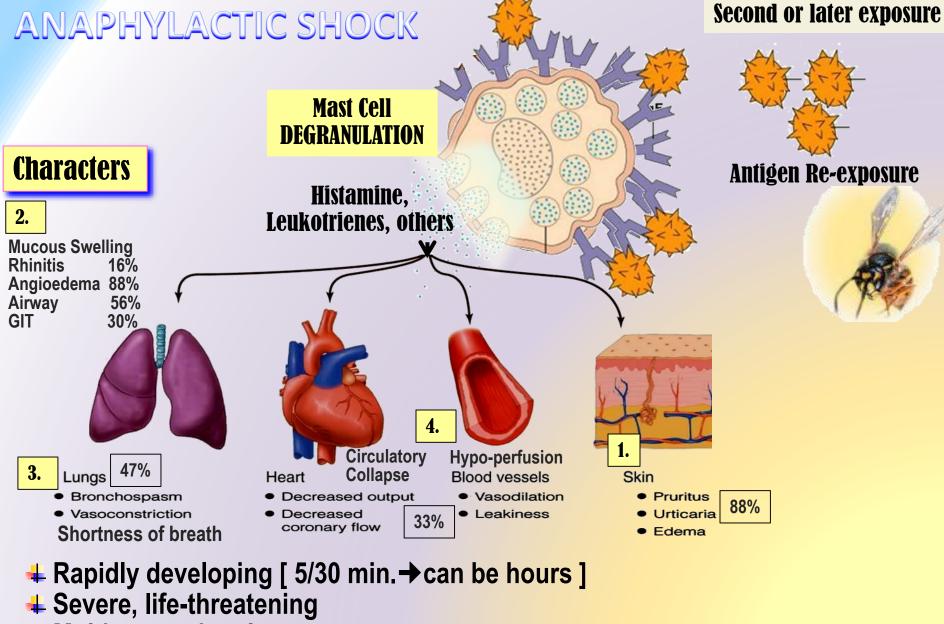
Occurs after exposure to foreign substances [antigen]; food, insect or animal venom, drugs, blood products,



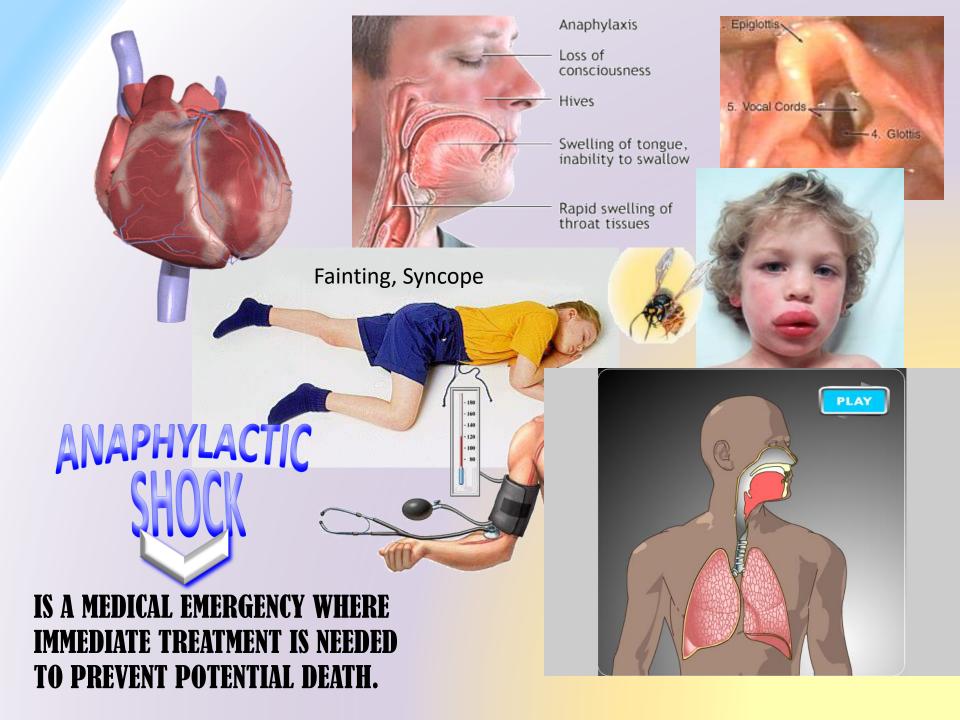
N.B. Non-Immunologic Anaphylaxis (ANAPHYLACTOID)

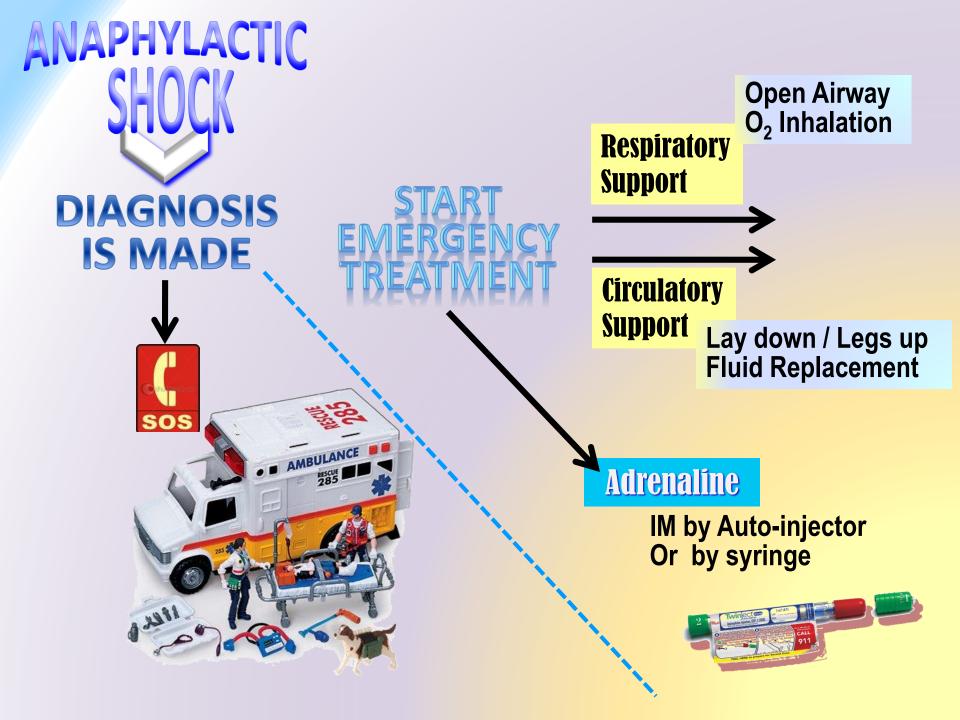
Exogenous substances directly degranulate mast cells→ Radiocontrast dye, Opiates, **Depolarinzing drugs, Dextrans**

Complete



- Multisystem involvement
- ♣ Mortality: due to respiratory (70%) or cardiovascular (25%)





ANAPHYLACTIC SHOCK THERAPY PROTOCOL

1 Life-threatening problems:

Airway: swelling, hoarseness, stridor

Breathing: rapid breathing, wheeze, fatigue, cyanosis, SpO, < 92%, confusion

Circulation: pale, clammy, low blood pressure, faintness, drowsy/coma

2 Adrenaline (give IM unless experienced with IV adrenaline) IM doses of 1:1000 adrenaline (repeat after 5 min if no better)

 Adult. 500 micrograms IM (0.5 mL)

Child more than 12 years: 500 micrograms IM (0.5 mL)

 Child 6 -12 years: 300 micrograms IM (0.3 mL)

Child less than 6 years: 150 micrograms IM (0.15 mL)

Adrenaline IV to be given only by experienced specialists

Titrate: Adults 50 micrograms; Children 1 microgram/kg

3 IV fluid challenge:

Adult - 500 - 1000 mL

Child - crystalloid 20 mL/kg

Stop IV colloid if this might be the cause of anaphylaxis

Adult or child more than 12 years

Child 6 - 12 years

Child 6 months to 6 years

Child less than 6 months

4 Chlorphenamine (IM or slow IV) 10 mg 5 mg 2.5 ma

250 micrograms/kg

5 Hydrocortisone (IM or slow IV) 200 mg 100 ma 50 mg 25 mg

6. Bronchodilators 7. Glucagon 8. H₂ Blockers

ANAPHYLACTIC SHOCK THERAPY PROTOCOL

ADJUVANT TO 2ND LINE

Bronchodilators

Salbutamol nebulizer / Ipratropium nebulizer / Aminophylline IV

Glucagon

For patients taking β-blockers & with refractory hypotension → 1 mg IV q 5 minutes until hypotension resolves

H₂ blocker

Ranitidine 50 mg IV / No cimetidine in elderly, renal/hepatic failure, or if on β -blockers

ANAPHYLACTIC SHOCK THERAPY PROTOCOL

ADJUVANT TO 2ND LINE

- **+To support the respiratory & circulatory deficits**
- To halt the existing hyper-reaction

Objective of Therapy

♣To prevent further hyper-reaction of immune system

Biphasic phenomenon

2nd release of mediators without re-exposure to antigen (in up to 20%) Clinically evident 3-4h after the initial manifestations clear

A Sympathomimetic.



Mechanism

A nonselective AD agonist $[\alpha_1, \alpha_2, \beta_1, \beta_2]$

Actions

As an α -AD agonist \rightarrow

- ♣Reverses peripheral vasodilation → maintains BP & directs blood flow to major organs
- **♣** dema **→** reverse hives, swelling around face & lips & angioedema in nasopharynex & larynx

As a β -AD agonist \rightarrow

- ♣ Dilates bronchial airways + ♣ histamine & leukotriene release from mast cells $\rightarrow \beta_2$ effect
- $+ \uparrow$ force of myocardial contraction $\rightarrow \beta_1$ effect

Contraindications

Rare in a setting of anaphylaxsis Not given > 40 y cardiac patient

ADRS

Dysrrhythmias

PHYSIOLOGICAL ANTAGONIST

Attenuates the severity of IgEmediated allergic reactions.

Indication | DRUG OF CHOICE

1ST LINE

Administration

Best is (IM) route in anaphylaxsis. Why?

- Easily accessible
- ♣ Greater margin of safety ◆ no dysrrhythmias as with IV
- ♣ No need to wait for IV line → if present
 - **→** given by physician under monitoring

Repeat every 5-10 min as needed Patients observed for 4-6 hours. Why? Fear of biphasic anaphylaxsis

N.B. Caution

Patients taking β -blockers either are \rightarrow

- **4**Refractory; as it may antagonize β effects of adrenaline
- ♣Rebound hypertension ♣ [unopposed α effect], specially when adrenaline is repeated

Auto-injectors Kits;

Disposable, prefilled devices → automatically administer a single dose of epinephrine in emergency

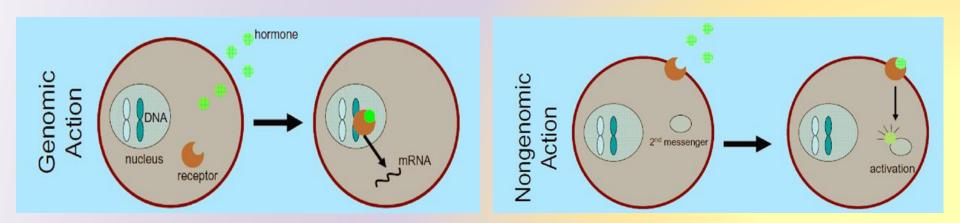
If hypotension persist → start dopamine. Why not noradrenaline?

CORTICOSTEROIDS



- It can not be used alone → not life saving Given slowly intravenously or intramuscularly.
- ■Reverse hypotension & bronchoconstriction → + release of inflammatory mediators (anti-chemotactic & mast cell stabilizing effects).
- Also decrease mucosal swelling and skin reaction.

This is through immediate GCs actions on <u>Membrane-bound receptors</u> → modulating levels of 2nd messengers → (within seconds or minutes) → Non-genomic action (genomic action is slow may take hrs to days)



May help to limit biphasic reactions → → allergic mediators

H₁ BLOCKERS



It can not be used alone → not life saving

Given slowly intravenously or intramuscularly (e.g. phenaramine).

Though mast cells have already de-granulated, yet these drugs can still help to counter act histamine-mediated vasodilation & bronchoconstriction.

H₂ BLOCKERS

The significance of H2 blockers is not established, these drugs are assocaited with serious adverse drug interactions.

BRONCHODIALATORS

Inhalational

*Salbutam0l→ β_2 -AD agonist →short acting, rapid relief onset relax bronchial smooth muscle and may decrease mediator release from mast cells and basophils.

It may also inhibit airway microvascular leakage.

***Ipratropium**→Anticholinergic →longer duration of action → ↓ secretion Less rapid in action

Parentral

Aminophylline IV → may be useful in the treatment of anaphylaxis when inhaled broncho-dilators are not effective & bronchospasm is persistent.

Given in hospital setting as levels of drug should be Therapeutically Monitored → has narrow therapeutic index



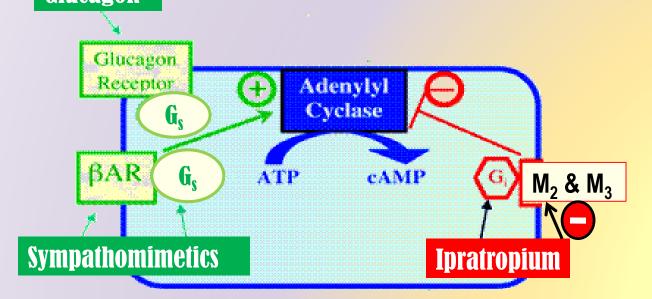


Drug of choice for severe anaphylaxis in patients taking β -blockers

Has both positive inotropic & chronotropic effects on heart → ↑ cardiac cyclic AMP → an effect entirely independent of AR

That is why effective in spite of beta-adrenergic blockade.

Efficacy of acting on bronchi < heart → no evident bronchodilation Glucagon





GOOD LUCK