



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

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

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# ANAPHYLAXSIS



Is a sudden, severe allergic reaction affecting the whole body

### symptoms including:

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



What TYPE of shock is it ???

### ANAPHYLACTIC SHOCK

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



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)
- **4 Cardiogenic** Inability to contract & pump→ myocardial infarction
- **♣ Obstructive**Extracardiac obstruction **→** Pul. embolism, cardiac tamponade
- **Distributive**Septic shock, neurogenic, anaphylactic shock.

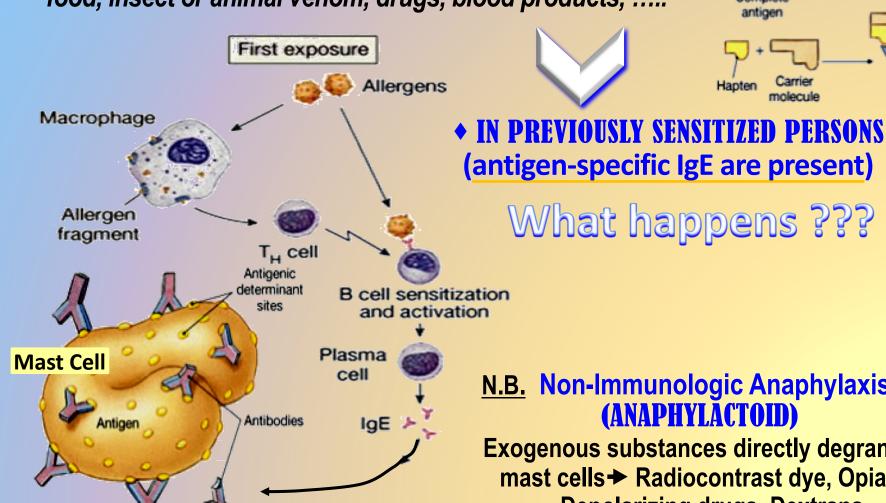
Severe, life-threatening, generalized or systemic hypersensitivity reaction in response to allergen.

### ANAPHYLACTIC SHOCK

**Nature** 

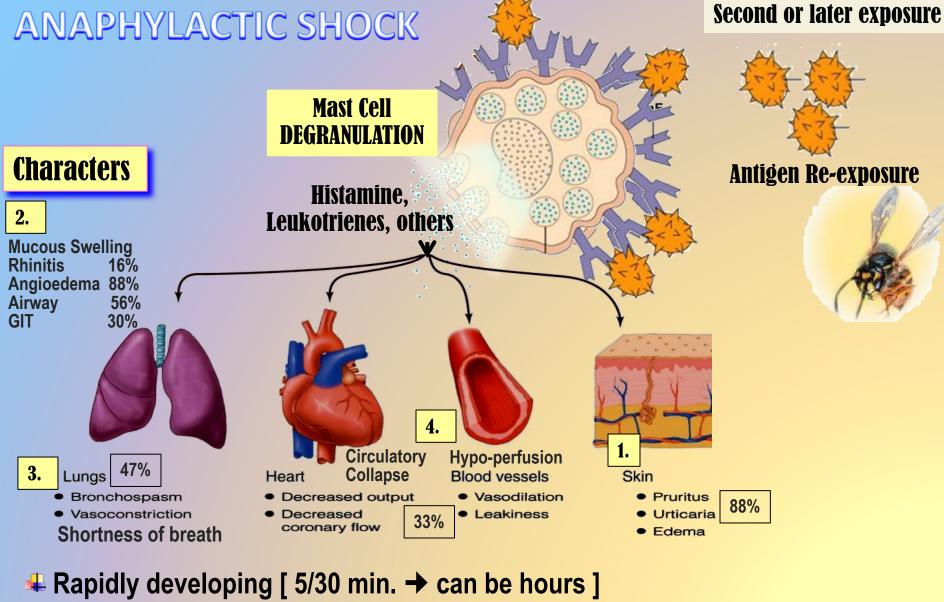
Belong to TYPE I HYPERSENSITIVITY REACTION

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

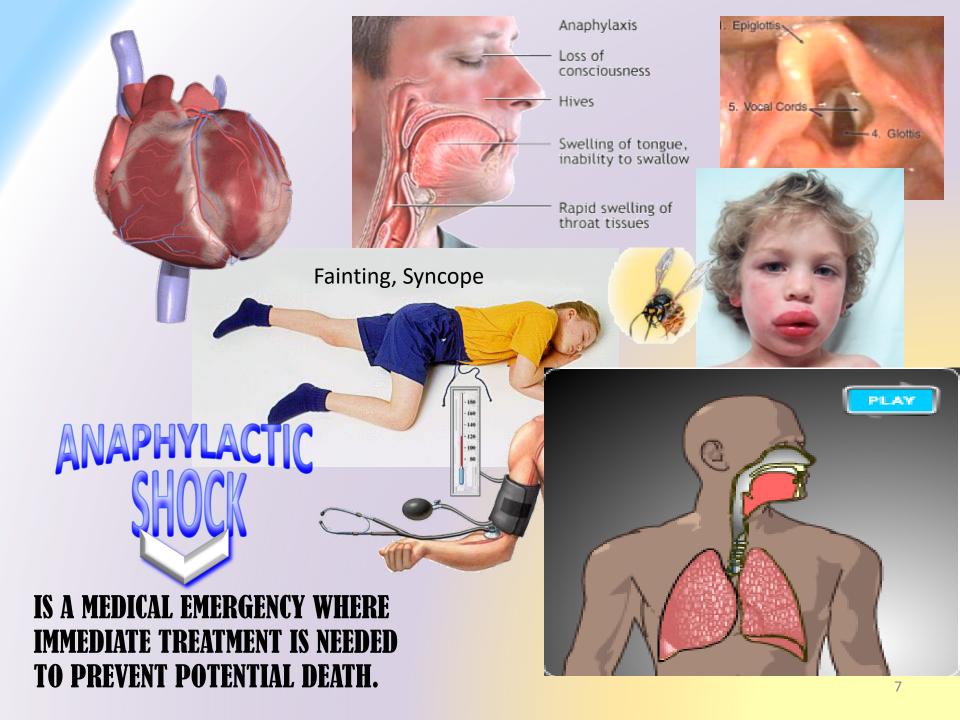


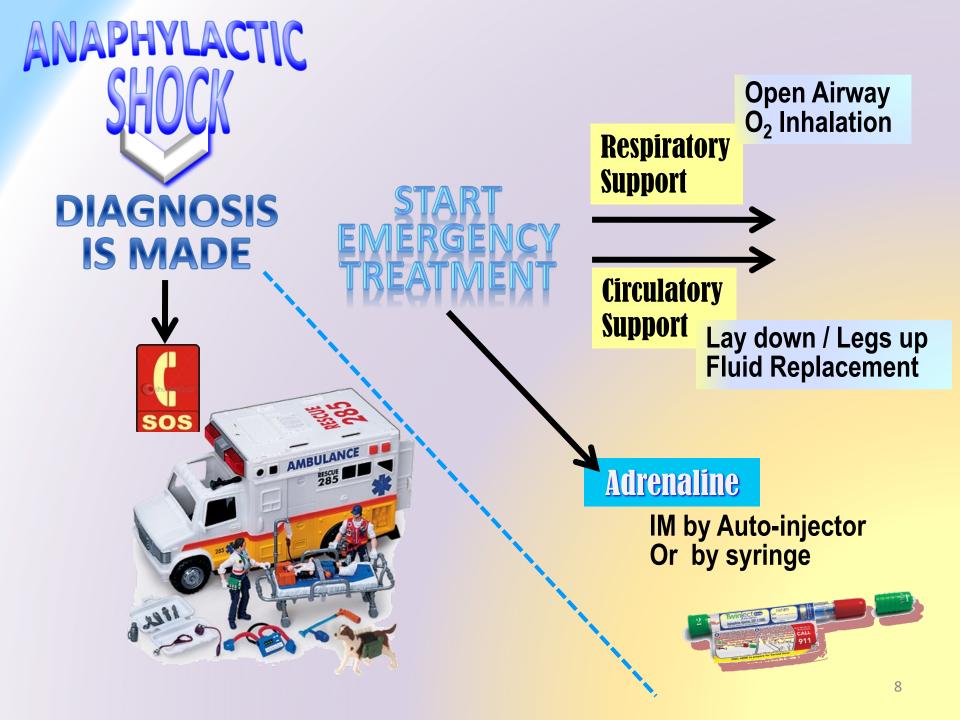
N.B. Non-Immunologic Anaphylaxis (ANAPHYLACTOID)

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



- Severe, life-threatening
- Multisystem involvement
- Mortality: due to respiratory (70%) or cardiovascular (25%).





### ANAPHYLACTIC SHOCK THERAPY PROTOCOL

1 Life-threatening problems:

swelling, hoarseness, stridor Airway:

Breathing: rapid breathing, wheeze, fatigue, cyanosis, SpO<sub>2</sub> < 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

4 Chlorphenamine

(IM or slow IV)

Adult or child more than 12 years

Child 6 - 12 years

Child 6 months to 6 years

Child less than 6 months

10 mg

5 mg

2.5 mg

250 micrograms/kg

5 Hydrocortisone

(IM or slow IV)

200 mg

100 mg

50 mg

25 mg

### ANAPHYLACTIC SHOCK THERAPY PROTOCOL

### ADJUVANT TO 2ND LINE

- **4To support the respiratory & circulatory deficits**
- To halt the existing hyper-reaction

**Objective of Therapy** 

**4To prevent further hyper-reaction of immune system** 

### **Biphasic phenomenon**

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

A Sympathomimetic.



**Mechanism** 

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

### **Actions**

As an  $\alpha$ -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 $\beta$ -AD agonist $\rightarrow$

- Dilates bronchial airways + → histamine & leukotriene release from mast cells  $\rightarrow \beta$ , effect
- + 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

### **ADRENALINE**

# 1ST LINE

#### **Administration**

Best is (IM) route in anaphylaxis. 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 anaphylaxis

#### N.B. Caution

Patients taking β-blockers either are →

- 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?

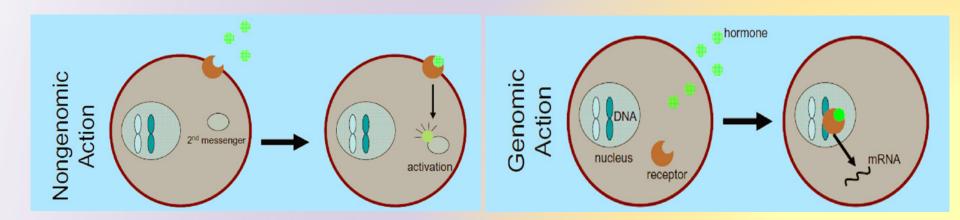
# CORTICOSTEROIDS



It can not be used alone → not life saving Given slowly IV or IM

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



May help to limit biphasic reactions → → allergic mediators.

# H<sub>1</sub> BLOCKERS



It can not be used alone → not life saving Given slowly IV or IM (e.g. pheniramine)

Though mast cells have already de-granulated, yet these drugs can still help to counteract histamine-mediated vasodilatation & bronchoconstriction

May help to limit biphasic reactions by blocking histamine receptors

# H<sub>2</sub> BLOCKERS

The significance of H2 blockers is not established, these drugs are associated with serious adverse drug interactions. Proton pump inhibitor (e.g. Pantoprazole) is safer and given once.

### 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

H<sub>2</sub> blocker: Ranitidine 150 mg IV / No cimetidine in elderly, renal/

hepatic failure, or if on β-blockers.??

## **BRONCHODILATORS**

#### **Inhalational**

- \*Salbutamol→ $\beta_2$ -AD agonist →short acting, rapid relief onset relax bronchial smooth muscle & may decrease mediators release from mast cells & 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  $\beta$ -blockers

Has both positive inotropic & chronotropic effects on heart  $\rightarrow \uparrow$  cardiac cyclic AMP  $\rightarrow$  an effect entirely independent of adrenergic receptors, that is why effective in spite of  $\beta$ -adrenergic blockade. Efficacy of acting on bronchi < heart  $\rightarrow$  no evident bronchodilation.

