

# POISONING

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## POISON/TOXIN:

- Substance able to produce adverse effects.
- Some of them are poisonous when taken in excessive amount, while some of them in any amount.



Poisoning:

Local

Systemic



## Severity and reverseability:

- Concentration (dose)
- Contact time
- The potency of the chemical
- Type and condition of the exposed surface
- Functional reserve of the individual/affected tissue
- Presence of secondary complications
- Coexisting illness



## Poison

- Caustic
- Neurotoxic (seizure, coma)
- Cardiotoxic
  - arrest
  - arrhythmias: bradycardia/tachycardia
- Haematotoxic
  - hemolysis
  - bone marrow depression
  - Prevents O<sub>2</sub> carrying capacity
- Nephrotoxic
  - acute tubular necrose

To evaluate the diagnosis:

- history
- physical examination
- routine/toxicology lab evaluation

## History:

- Time, route and duration of exposure
- Name/amount of each drug/chemical
- Time of onset of the symptoms
- Nature /severity of symptoms
- Post medical and psychiatric history

# Physical examination:

Vital signs

Cardiopulmonary and neurological status





# Treatment:

- Emergency
- Maintenance

# Emergency Treatment:

I) Supportive care (vital sign)

II) Prevention of further poison absorption

III) Enhancement of poison elimination

IV) Administration of specific antidotes



## I) Supportive care:

- Airway protection
- Oxygenation/ventilation
- Hemodynamic support
- Treatment of seizures
- Correction of temperature abnormalities
- Prevention of secondary complications



## **All symptomatic patients should have:**

- IV line
- O2 supplemetation
- Cardiac monitoring
- Baseline laboratory
- Continous observation

## Depression of CNS

### Stimulators

- Caffeine
- Coramine
- Na benzoat
- Cardiazol

# Depression of respiration:

- O<sub>2</sub>
- Endotracheal tube
- CNS stimulators

Seizure:

Sedatives:

- Diazepam
- Phenobarbital

# Shock:

To increase venous return

- legs are held up
- elastic band is performed to the legs
- fluid perfusion (volume expander)



TA > 85 mmHg  
urine putput > 30 ml/h

no need a dense treatment  
(fluid perfusion is enough)

TA < 85 mmHg  
urine output < 30 ml/h  
heart rate >110/min

fluid perfusion and vasopressor  
agents

(dopamin, dobutamin, adrenalin,  
noradrenalin)



## II) Prevention of further poison absorption

### **Vomiting:**

- Spontaneously
- Sirop d' ipe'ca
- Apomorphine (!!! CNS depression)
- Salt

Vomiting is **contraindicated:**

Caustic, corrosive toxins

Petroleum distillation products

Coma, seizures (Aspiration)

## II) Prevention of further poison absorption

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### **Gastric lavage:**

In Trendelenburg and left lateral decubitus position to prevent aspiration

It should be performed in first 4 hours (can be delayed to 6 hours in salicylates)

It can be performed later if the poison taken after meals



## Gastric lavage is **contraindicated**:

- corrosive poisons (acid, alkaline)
- strychnine
- petrol distillation products

\*It is too late for gastric lavage in a comatose patient; if wanted should be intubated



## II) Prevention of further poison absorption

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### Activated charcoal:

-by mouth or by a stomach tube before and after gastric lavage

-as an adsorbent for:

alcohol-atropin-morphin-opium

arsenic-barbiturate-nicotin-penicilin

salicylates



## II) Prevention of further poison absorption

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### Whole bowel irrigation:

Bowel cleansing solution (electrolytes and polyethyleneglicol).

It may be of particular benefit in patients with foreign body, drug packed and slow release medication injections

## II) Prevention of further poison absorption

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Dilution:

Ingestion of corrosive (acid-alkaline)

Water or other clean liquid



## II) Prevention of further poison absorption

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### Endoscopic or surgical removal:

- Ingestion of a potentially toxic foreign body that fails to transit the GI tractus (potentially lethal amount of a heavy metal-arsenic, iron, Hg, thallium)
- Ingestion of packets of drugs (cocaine)



## Enhancement of poison elimination

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### **A) Multiple dose activated charcoal**

A dose of 1 g/kg for every 2 to 4 hour (with sorbitol as needed to enhance GI motility)

## Enhancement of poison elimination

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### **B) Force diuresis and alteration of urinary pH**

- For the poisons that are excreted by the kidney  
(excreted by glomerular filtration and active tubular secretion)

- Renal reabsorption of poison is prevented

  - Mannitol (20%-250 ml-IV)

  - \*Contraindications: Congestive heart failure, renal failure

- Alkaline diuresis (pH>7.5):

Na HCO<sub>3</sub> / Na lactate added in fluid.

salicylates, phenobarbital, chlorpropamide

- Acide diuresis

\*\* (not used because of significant risks)

(amphetamines, cocain, quinidine)

\*Acid-base balance, Fluid and electrolyte parameters should be carefully monitored


# Enhancement of poison elimination

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## **C) Extracorporeal removal**

- Dialysis
  - Peritoneal dialysis
  - Haemodialysis
- Haemoperfusion
- Exchange transfusion

## **Dialisable molecule:**

- Low molecular weight
  - High water solubility
  - Low protein binding
  - small volume of distribution
  - prolonged elimination (long half life)
  - high dialysis clearance
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## **Dialysis is preferred:**

- in anuric cases
- The metabolites of the poison is more toxic  
(Methanol - Formic acid)
- ethanol, methanol, salicylate, lithium, heavy metals, bromide, etc.

## **Exchange transfusion:**

- Less effective, but it may be used when other procedures are not effective or are contraindicated
- removes poison affecting red blood cells  
(methemoglobinemia)

# Neutralisation

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## **- Adsorption**

Active carbon

## **- Neutralisation of the acids**

Milk of magnesia

Na HCO<sub>3</sub>

CaCO<sub>3</sub>

Ca(OH)<sub>2</sub>


## **- Neutralisation of alkaline**

Acetic acid

Lemon juice

Orange juice

\*Milk, olive oil, white of the egg and starch protect the mucosa and delay the absorption of the poison





## REFERENCES

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