

# Anti-muscarinic anti-cholinergic drugs



• Natural agents:

• Atropine, Hyoscine

• Semi-synthetic

• Homatropine

• Synthetic

• Ipratropium, Pirenzepine, Propantheline

Atropine: *Atropa Belladonna*

Natural Agents Have 2 disadvantages

- Non-selective (block all Muscarinic receptors)

- Tertiary Amides (can pass through BBB)

will have central + side effects

\* if we want to make a substitute for Atropine that doesn't pass through BBB then we'll make quaternary drugs

# Pharmacodynamics

**Exocrine glands: at low doses reduced secretions**

Salivary

Bronchial

Sweat glands

} M<sub>3</sub> Receptor

Blocking of the M<sub>3</sub> causes dryness  
(using Atropine)

No sweating

No Nasal secretion

No lacrimal secretion

# CVS



Depending in the doses

- Central effect:
- Decrease heart rate
- Peripheral effect:
- Blockade of vagus nerve and increase heart rate
- ABP:
- No change

Blood vessels:  $M_3$  (Non-innervated / silent)  
so closing won't change anything

Toxic doses  $\rightarrow$  Flushing

Toxic U.D

# Respiratory system



- Bronchodilatation *Bronchi:  $M_3$*
- Reduced bronchial secretion
- Ipratropium (quaternary amine derivate of Atropine)  
inhalation:
- Useful in asthma and chronic obstructive pulmonary disease (COPD), also in patient who are unable to take adrenergic agonists.

$M_3$  in bronchi  $\rightarrow$  bronchoconstriction + increase secretion

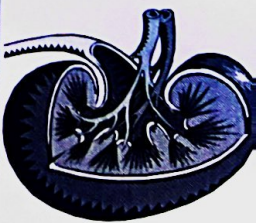
can be used in Bronchial Asthma

Atropine will inhibit the "bronchoconstriction" from happening which leaves space for ~~B2~~  $\beta_2$  to work causing bronchodilatation

\* So Atropine caused bronchodilatation but passively

genitourinary tract

# GUT M<sub>3</sub>



- Relaxation of bladder wall
- Useful in inflammatory spasm and pains of the urinary tract
- Risky in patients with **BPH** (Benign Prostatic Hypertrophy) *urine retention*  
*closed sphincters*  
*+ relaxed smooth muscles*