Antiemetics

Prof. Hanan Hagar
Dr Ishfaq Bukhari
Pharmacology Department
College of Medicine

Learning objectives

- •Classify the main different classes of antiemetic drugs according to their mechanism of action.
- •Know the characteristic pharmacokinetics & dynamics of different classes of antiemetic drugs.
- •Identify the selective drugs that can be used according to the cause of vomiting.
- Learn the adjuvant antiemetics.
- •Describe the major side effects for the different classes of antiemetics.

Vomiting

- Is a complex series of integrated events culminating in the forceful expulsion of gastric contents through the mouth.
- Vomiting can be a valuable, life-saving physiological response WHY????

Consequences of vomiting

- Severe vomiting may result in :
- Dehydration
- Acid-base imbalance
- Electrolyte depletion

Causes of Vomiting

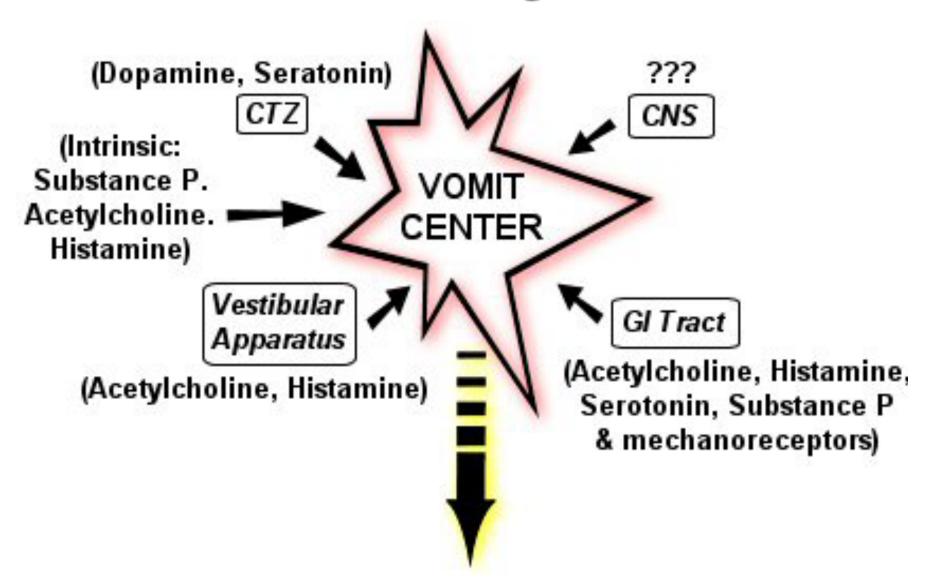
Nausea and vomiting occurs due to stimulation of vomiting center that respond to inputs from:

- Higher cortical centers stimulation (CNS)
- Chemoreceptor trigger zone (CTZ) stimulation
- Disturbance of vestibular system
- The periphery (Pharynx, GIT) via sensory nerves

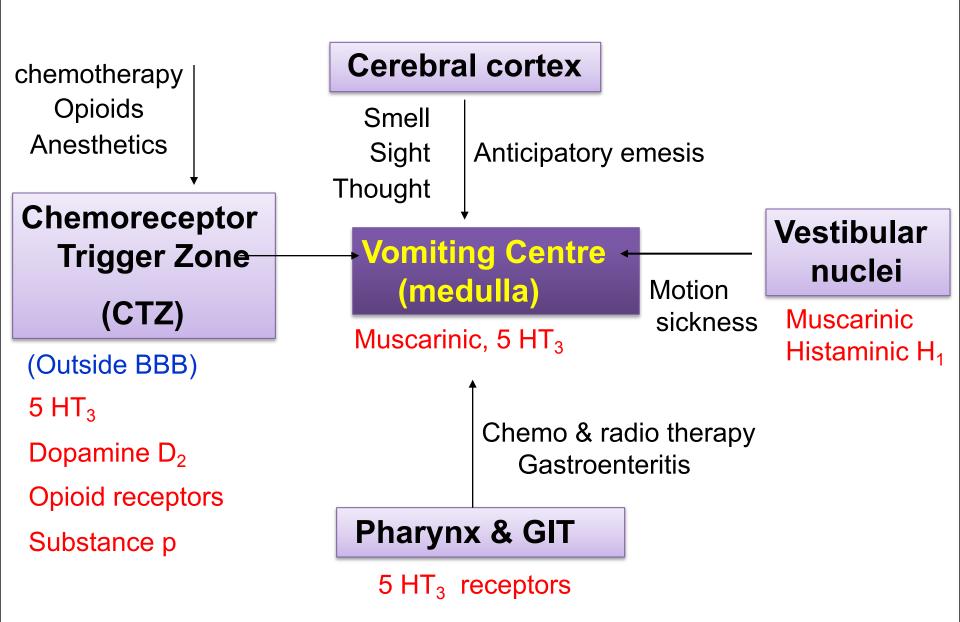
1. Stimulation of chemoreceptor trigger zone (CTZ)

- CTZ is an area of medulla that communicate with vomiting center to initiate vomiting.
- CTZ is physiologically outside BBB.
- CTZ contains D₂ receptors, 5 HT₃ receptors & opioid receptors.
- stimulated by:
- Emetogenic drugs (opioids, general anesthetics, digitalis, L-dopa).
- chemicals and toxins (blood, CSF).
- Radiation.
- Uremia, estrogent (vomiting of pregnancy)

Receptors Associated with Nausea and Vomiting



Pathophysiology of Emesis



Chemical transmitters & receptors involved in vomiting and drug tragets:

- Ach (Muscarinic receptors)
- Dopamine (D2)
- Histamine (Histaminergic receptors H1)
- Serotonin (5-HT3)
- Substance P (Neurokinin receptors, NK1)
- Opioid (Opioid receptors)

Classification of Antiemetic Drugs

- 1. 5-HT3 antagonists
- 2. D₂ receptor <u>antagonists</u>
- 3. NK₁ antagonists
- 4. H₁-receptor <u>antagonists</u>
- 5. Muscarinic receptor antagonists
- 6. Glucocorticoids

Serotonin (5-HT3) antagonists

- Drugs as
 - Ondansetron
 - Granisetron
- Orally or parenterally,
- have long duration of action, first pass effect
- The most potent antiemetic drugs
- Act by blocking 5-HT3 receptor centrally (in vomiting center, CTZ) and peripherally (5HT3 receptors on GI vagal afferents).

Uses of 5-HT3 antagonists

- First choice for prevention of moderate to severe emesis:
 - -Chemotherapy-induced nausea and vomiting (CINV) especially cisplatin
 - -Post-radiation NV& Post-operative NV
 - Their effects is augmented by combination with corticosteroids and NK₁ antagonists.

Side effects

- Well tolerated
- Headache, dizziness and constipation
- o minor ECG abnormalities (QT prolongation)

D₂ receptor antagonists

- block D₂ dopamine receptors in the CTZ
- Two types exist:
 - Prokinetics drugs
 - Neuroleptics (antipsychotics)

D2 receptor antagonists

Prokinetics drugs

- Domperidone: oral
- Metoclopramide: oral, i.v
- Are prokinetic agents (increased GI motility & gastric emptying).

Uses

- Antiemetics (blocking D2 receptors in CTZ)
 - Effective against vomiting due to cytotoxic drugs, gastroenteritis, surgery, toxins, uremia, radiation
- Prokinetic (5 HT4 agonist activity)
 - Gastroesophageal reflux disease (GERD)
 - Gastroparesis (impaired gastric emptying after surgery).

Metoclopramide crosses BBB but domperidone cannot (both have antiemetic effects as CTZ is outside BBB).

Side effects (only for metoclopramide):

- Dyskinesia (extra-pyramidal side effects),
- Galactorrhea, menstrual disorders, impotence
- \checkmark Postural hypotension (α -blocking action).
- Sedation, drowsiness

Other D2 receptor antagonists

Neuroleptics (Antipsychotics)

- Chlorpromazine (CPZ), droperidol
- used for postoperative vomiting and chemotherapy-induced emesis.

Side effects:

- Extra pyramidal symptoms
- Sedation
- Postural hypotension

Neurokinin1 (NK1) receptor antagonists Aprepitant

- Acts centrally as <u>substance P antagonist</u> by blocking neurokinin 1 receptors in vagal afferent fibers.
- Orally
- Usually combined with 5-HT₃ antagonists and corticosteroids in prevention of chemotherapy-induced nausea and vomiting and post- operative NV.

H₁-receptor antagonists

Include drugs as

- diphenhydramine, promethazine
- meclizine, cyclizine

Used for

- Motion sickness
- Morning sickness in pregnancy
- Promethazine: severe morning sickness of pregnancy (if only essential).

Side effects:

- -Prominent sedation
- -Hypotension
- -Anticholinergic effects or atropine like actions (dry mouth, dilated pupils, urinary retention, constipation).

Muscarinic receptor antagonists

- Hyoscine (scopolamine)
- Orally, injection, patches
- Used as transdermal patches in motion sickness (applied behind the external ear).
- Reduce impulses from vestibular apparatus
- Not in chemotherapy-induced vomiting

Side effects:

- Sedation
- Tachycardia, blurred vision, dry mouth, constipation, urinary retention (atropine-like actions).

Glucocorticoids

- Dexamethasone methylprednisolone
- Used in chemotherapy-induced vomiting
- combined with 5-HT₃ antagonists or NK1 receptor antagonists.

Glucocorticoids

Side effects if used for long term:

- Hyperglycemia
- Hypertension
- Cataract
- Osteoporosis
- Increased intraocular pressure
- Increased susceptibility to infection
- Increased appetite & obesity

Summary

The choice of antiemetic depends on the etiology

Motion sickness

Muscarinic antagonists

Antihistaminics

Vomiting with pregnancy (morning sickness)

avoid all drugs in the first trimester

Pyridoxine (B6)

Promethazine (late pregnancy).

Drug- induced vomiting (CTZ), uremia, gastritis Dopamine antagonists

Post operative nausea & vomiting Dopamine antagonists

Vomiting due to cytotoxic drugs.

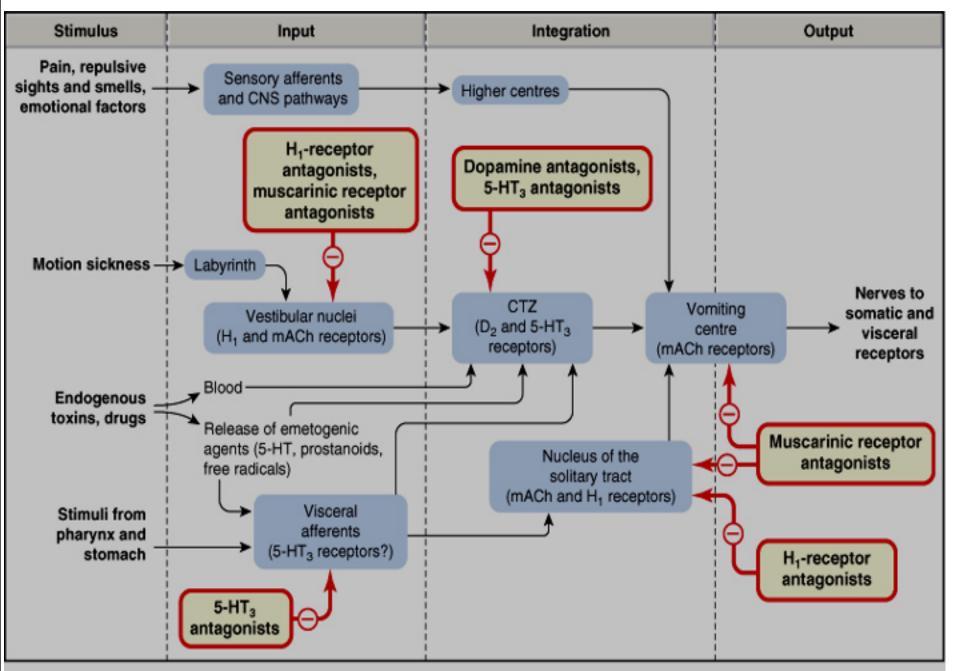
5-HT₃ antagonists

NK₁ antagonists

D₂- antagonists

Glucocorticoids

Cannabinoids



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