

SNS COLLEGE OF ALLIED HEALTH SCIENCES SNS Kalvi Nagar, Coimbatore - 35 Affiliated to Dr MGR Medical University, Chennai

DEPARTMENT OF CARDIO PULMONARY PERFUSION CARE TECHNOLOGY

COURSE NAME : Pharmacology Pathology and Clinical Microbiology II nd YEAR **TOPIC : PHARMACOLOGY OF DRUGS USED IN BROCHIAL ASTHMA AND** COPD





Pharmacology of drugs used in bronchial asthma & COPD







Bronchial Asthma

Asthma is a chronic inflammatory disorder of bronchial airways that result in airway obstruction in response to external stimuli or triggers (as pollen grains, cold air and tobacco smoke, animal fur).







Characters of airways in asthmatic patients :

- Airway hyper-reactivity: abnormal sensitivity of the airways to any external stimuli.
- Inflammation
 - ↑ edema, swelling
 - Thick mucus production.

Bronchospasm (constriction of the bronchial smooth muscles).









http://link.brightcove.com/services/player/bcpid236059233?bctid=347806802





Airway hyper-reactivity Endogenous inflammatory mediators e.g. histamine, leukotrienes







Symptoms of asthma

Asthma produces recurrent episodic attack of

- **Acute bronchoconstriction**
- **Shortness of breath**
- **Chest tightness**
- Wheezing
- **Rapid respiration**
- Cough

Symptoms can happen each time the airways are irritated by inhaled irritants or allergens.







- **Exogenous chemicals or irritants**
- **Chest infections**
- **Stress**
- **Exercise (cold air)**
- Pets
- **Seasonal changes**
- **Emotional conditions**
- Some drugs as aspirin, β-bockers







Innervation of respiratory system > Parasympathetic supply M3 receptors in smooth muscles and glands.

Bronchoconstriction

Increase mucus secretion

 \triangleright No sympathetic supply but B₂ receptors in smooth muscles and glands.

Bronchodilation

Decrease mucus secretion







Anti asthmatic drugs:

1) Quick relief medications: Bronchodilators used to relieve acute episodic attacks of asthma.

2) Control therapy (prophylactic drugs): anti-inflammatory drugs used to reduce the frequency of attacks, and nocturnal awakenings.





Anti asthmatic drugs

Bronchodilators

(Quick relief medications)

treat acute attack of asthma

- Short acting β 2-agonists
- Antimuscarinics
- **Xanthine preparations**

(**Prophylactic therapy**)

reduce the frequency of attacks

- Corticosteroids
- Mast cell stabilizers





Anti-inflammatory Agents

Leukotrienes antagonists

Anti-IgE monoclonal antibody

Long acting ß2-agonists



Bronchodilators

These drugs can produce rapid relief of bronchoconstriction.

Bronchodilators:

- **β2** adrenoreceptor agonists
- Antimuscarinics
- **Xanthine preparations**









Sympathomimetics β- adrenoceptor agonists

Classification of *β* **agonists** \triangleright Non selective β agonists: epinephrine - isoprenaline

 \triangleright Selective $\beta 2$ – agonists (Preferable). **Salbutamol (albuterol) Terbutaline Salmeterol Formeterol**





Mechanism of Action

 \succ direct β_2 stimulation \longrightarrow stimulate adenyl cyclase $\longrightarrow \uparrow cAMP \rightarrow bronchodilation.$

Increase mucus clearance by (increasing ciliary) activity).

>Stabilization of mast cell membrane.





Non selective β-agonists. Epinephrine

- **Potent bronchodilator**
- Given subcutaneously, S.C.
- rapid action (maximum effect within 15 • min).
- Has short duration of action (60-90 min) •
- **Drug of choice** for acute anaphylaxis • (hypersensitivity reactions).





Disadvantages >Not effective orally. >Hyperglycemia >Skeletal muscle tremor **CVS side effects:** tachycardia, arrhythmia, hypertension >Not suitable for asthmatic patients with hypertension or heart failure.

Contraindications: CVS patients, diabetic patients







Selective β2 –agonists

- Are mainly given by inhalation by (metered dose inhaler or nebulizer).
- Can be given orally, parenterally.
- Short acting B2 agonists
 - e.g. salbutamol, terbutaline
- Long acting B2 agonists
 - e.g. salmeterol, formoterol







Nebulizer







Inhaler









Short acting B₂ **agonists Salbutamol**, inhalation, orally, i.v. **Terbutaline**, inhalation, orally, s.c. > Have rapid onset of action (15-30 min). > short duration of action (4-6 hr) > used for acute attack of asthma (drugs of choice).





Long acting selective B₂ agonists **Salmeterol & formoterol** ➤ are given by inhalation Long acting bronchodilators (12 hours) due to high lipid solubility (creates depot effect). *▶*<u>are not used to relieve</u> acute episodes of asthma ≻used for nocturnal asthma. >combined with inhaled corticosteroids to control asthma (decreases the number and severity of asthma attacks).







Advantages of B₂ agonists > Minimal CVS side effects > suitable for asthmatic patients with CV disorders as hypertension or heart failure. **Disadvantages of B**₂ **agonists** Skeletal muscle tremors. > Nervousness

- > Tolerance (β -receptors down regulation).
- > Overdose may produce tachycardia due to β_1 stimulation.





Muscarinic antagonists Ipratropium – Tiotropium >Act by blocking muscarinic receptors . >given by aerosol inhalation >Have delayed onset of action. >Quaternary derivatives of atropine (polar). >Does not diffuse into the blood >Does not enter CNS. >Have minimal systemic side effects >Ipratropium has short duration of action 3-5 hr **Tiotropium** has longer duration of action (24) h).





Pharmacodynamics

- Inhibit bronchoconstriction and mucus secretion
- > Less effective than β_2 -agonists.
- > No anti-inflammatory action only bronchodilator

Uses

- > Main choice in chronic obstructive pulmonary diseases (COPD).
- > In acute severe asthma combined with β_2 agonists & corticosteroids.
- > Never use as a rescue medication.





Methylxanthines

Theophylline - aminophylline

Mechanism of Action > are phosphodiestrase inhibitors \succ \uparrow cAMP \rightarrow bronchodilation > Adenosine receptors antagonists (A1) > Increase diaphragmatic contraction > Stabilization of mast cell membrane









Adenyl cyclase

Phosphodiesterase

Theophylline



Pharmacological effects : ➢Bronchial muscle relaxation \rightarrow f contraction of diaphragm \rightarrow improve ventilation **CVS:** ↑ heart rate, ↑ force of contraction **GIT:** ↑ gastric acid secretions Kidney: *renal blood flow, weak divertic action* **CNS** stimulation

- * stimulant effect on respiratory center.
- * decrease fatigue & elevate mood.
- * overdose (tremors, nervousness, insomnia, convulsion)





Pharmacokinetics Theophylline is given orally >Aminophylline, is given as slow infusion > metabolized by Cyt P450 enzymes in liver $T^{1/2} = 8$ hours >has many drug interactions > Enzyme inducers: ➤ as phenobarbitone & rifampicin > \uparrow metabolism of the ophylline $\rightarrow \downarrow T \frac{1}{2}$. > Enzyme inhibitors: > as erythromycin \downarrow metabolism of the ophylline $\rightarrow \uparrow T \frac{1}{2}$.





Uses

Second line drug in asthma (theophylline). For status asthmatics (aminophylline, is given as slow infusion).

Side Effects

- > Low therapeutic index (narrow safety margin) monitoring of theophylline blood level is necessary.
- ➢ GIT effects: nausea & vomiting
- **CVS effects:** hypotension, arrhythmia.
- **CNS** side effects: tremors, nervousness, insomnia, convulsion





Prophylactic therapy

Anti - inflammatory drugs include: Glucocorticoids >Leukotrienes antagonists >Mast cell stabilizers >Anti-IgE monoclonal antibody e.g. omalizumab

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Anti - inflammatory drugs: (control medications / prophylactic therapy)

- ↓ bronchial hyper-reactivity.
- I reduce inflammation of airways
- \downarrow reduce the spasm of airways





Glucocorticoids **Mechanism of action**

Anti-inflammatory action due to: Inhibition of phospholipase A2 ↓ prostaglandin and leukotrienes ↓ Number of inflammatory cells in airways. Mast cell stabilization $\rightarrow \downarrow$ histamine release. ↓ capillary permeability and mucosal edema. Inhibition of antigen-antibody reaction. Upregulate β_2 receptors (have additive effect to B_2) agonists).











Pharmacological actions of glucocorticoids

- > Anti-inflammatory actions
- Immunosuppressant effects
- > Metabolic effects
 - Hyperglycemia
 - protein catabolism,
 protein anabolism
 - Stimulation of lipolysis fat redistribution
- > Mineralocorticoid effects:
 - sodium/fluid retention
 - Increase potassium excretion (hypokalemia).
 - Increase blood volume (hypertension).











Routes of administration Inhalation:

- e.g. Budesonide & Fluticasone, beclometasone
 - Given by inhalation (metered-dose inhaler).
 - Have first pass metabolism
 - Best choice in asthma, less side effects
- Orally: Prednisone, methyl prednisolone
- Injection: Hydrocortisone, dexamethasone



eclometasone ose inhaler).

effects ednisolone xamethasone



Glucocorticoids in asthma Are not bronchodilators Reduce bronchial inflammation Reduce bronchial hyper-reactivity to stimuli •Have delayed onset of action (effect usually attained after 2-4 weeks). Maximum action at 9-12 months. •Given as prophylactic medications, used alone or combined with β_2 agonists. Effective in allergic, exercise, antigen and irritant-induced asthma,





Systemic corticosteroids are reserved for: – Status asthmaticus (i.v.).

Inhaled steroids should be considered for adults, children with any of the following features using inhaled β_2 agonists three times/week symptomatic **three times**/ week or more; or waking **one night**/week.







Clinical Uses of glucocorticoids

- Treatment of inflammatory disorders (asthma, 1. rheumatoid arthritis).
- Treatment of autoimmune disorders (ulcerative 2. colitis, psoriasis) and after organ or bone marrow transplantation as immunosuppressants.
- **3.** Antiemetics in cancer chemotherapy.





Side effects due to systemic corticosteroids

- Adrenal suppression —
- Growth retardation in children
- Susceptibility to infections —
- Osteoporosis
- Fluid retention, weight gain, hypertension
- Hyperglycemia
- Fat distribution
- Cataract
- Psychosis







Inhalation has very less side effects: – Oropharyngeal candidiasis (thrush). – Dysphonia (voice hoarseness).

Withdrawal of systemic corticosteroids Abrupt stop of corticosteroids should be avoided and dose should be tapered (*adrenal*) insufficiency syndrome).





Mast cell stabilizers

e.g. Cromoglycate - Nedocromil > act by stabilization of mast cell membrane. > given by inhalation (aerosol, nebulizer). ≻Have poor oral absorption (10%)







Pharmacodynamics

- are Not bronchodilators
- Not effective in acute attack of asthma.
- Prophylactic anti-inflammatory drug
- Reduce bronchial hyper-reactivity.
- Effective in exercise, antigen and irritantinduced
 - asthma.
- Children respond better than adults





Uses

- Prophylactic therapy in asthma especially in children.
- Allergic rhinitis.
- Conjunctivitis.

Side effects

- Bitter taste
- minor upper respiratory tract irritation (burning sensation, nasal congestion)





Leukotrienes antagonists Leukotrienes

- •synthesized by inflammatory cells found in the airways (eosinophils, macrophages, mast cells).
- produced by the action of <u>5-lipoxygenase</u> on arachidonic acid.
- Leukotriene B4: chemotaxis of neutrophils ► Cysteinyl leukotrienes C4, D4 & E4: bronchoconstriction increase bronchial hyper-reactivity – ↑ mucosal edema, ↑ mucus secretion











Leukotriene receptor antagonists

- e.g. zafirlukast, montelukast, pranlukast
 - are selective, reversible antagonists of cysteinyl leukotriene receptors
 (CysLT₁receptors).
- ➤ Taken orally.
- Are bronchodilators
- Have anti-inflammatory action
- Less effective than inhaled corticosteroids
- Have glucocorticoids sparing effect (potentiate corticosteroid actions).

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ticosteroids effect ns).



Uses of leukotriene receptor antagonists

- Not effective in acute attack of asthma.
- **Prophylaxis** of mild to moderate asthma.
- Aspirin-induced asthma
- Antigen and exercise-induced asthma
- > Can be combined with glucocorticoids (additive effects, low dose of glucocorticoids can be used).

Side effects:

Elevation of liver enzymes, headache, dyspepsia





Anti-IgE monoclonal antibody e.g. Omalizumab

- •is a monoclonal antibody directed against human IgE – given by injection (s.c.) • prevents IgE binding with its receptors on mast cells & basophiles.
- release of allergic mediators.
- Expensive-not first line therapy.
- •used for treatment of moderate to severe allergic asthma which does not respond to high doses of corticosteroids.















Figure 1. Mechanisms of action of omalizumab in allergic asthma. Reprinted by permission from Macmillan Publishers Ltd: Nat Rev Immunol,14 copyright 2008. Abbreviation: Fc RI, high-affinity IgE receptor.



Drugs used in chronic obstructive pulmonary disease (COPD)

COPD is a chronic irreversible airflow

obstruction, lung damage and inflammation of the air sacs (alveoli).

Smoking is a high risk factor but air pollution and genetic factors can contribute.





Bronchioles lose their shape and become clogged with mucus

Bronchioles (tiny airways)

Left lung

Treatment:

- Inhaled bronchodilators – Inhaled glucocorticoids -Oxygen therapy -Antibiotics specifically macrolides such as azithromycin to reduce the number of exacerbations.
- -Lung transplantation

Inhaled bronchodilators in COPD >Inhaled antimuscarinics > Ipratropium & tiotropium. \triangleright are superior to $\beta 2$ agonists in COPD $>\beta_2$ agonists > these drugs can be used either alone or combined – salbutamol + ipratropium - salmeterol + Tiotropium (long acting-less dose frequency).

Summary

Exposure to antigen (dust, pollen, etc)

Antigen and IgE on mast cells

Mediators (leukotrienes, cytokines, etc)

β Agonists, theophylline, muscarinic antagonists, zafirlukast

Early response: bronchoconstriction

Late response: inflammation

Acute symptoms

Cromolyn, steroids, zileuton

Steroids, cromolyn, leukotriene antagonists

Bronchial hyperreactivity

Bronchodilators (relievers for bronchospasm)

Drugs			
B2 agonists	- Short acting	↑Adenyl	INS
Salbutamol, terbutaline	– main choice in acute attack of asthma	cyclase	
	- Inhalation		
Salmeterol, formoterol	Long acting, Prophylaxis		
	Nocturnal asthma		
Antimuscarinics	Main drugs For COPD	Blocks M	
Ipratropium (Short)	Inhalation	recepttors	
Tiotropium (long)	Inhalation		
Xanthine derivatives		Inhibits	
Theophylline	(orally)	phosphodi	
Aminophylline	(parenterally)	esterase	
		↑ cAMP	

Anti-inflammatory drugs (prophylactic)

Corticosteroids

(Inhibits phospholipase A2)

Dexamethasone, Fluticasone, budesonide

prednisolone

Hydrocortisone

Mast stabilizers

Cromoglycate (Cromolyn), Nedocromil

Cysteinyl antagonists (CyLT1 antagoist) Zafirlukast, montelukast

Omalizumab (Anti IgE antibody)

Inhalation

Orally

parenterally

Inhalation, prophylaxis in children

orally

Injection, SC

THANK YOU

