CORTICOSTEROIDS

DR SHAMS SULEMAN

LEARNING OBJECTIVES

- Describe the role of corticosteroids in General
- Describe the role of corticosteroids in particular as an immunosuppressant
- Describe clinical uses and adverse effects of corticosteroids

ADRENOCORTICAL HORMONES

Basal Secretion

Group	Hormone	Daily
Glucocorticoids	Cortisol Corticosterone	5 – 30 mg 2 – 5 mg
Mineralocorticoids	Aldosterone 11- deoxycorticosterone	5 – 150 mcg Trace
Sex Hormones		
Androgen	DHEA	15 – 30 mg
Progestogen	Progesterone	0.4 – 0.8 mg
Oestrogen	Oestradiol	Trace

ADRENAL HORMONES: SOURCES

Natural

- Glucocorticoids
 - Cortisol (Hydrocortisone)
 - Corticosterone
- Mineralocorticoids
 - v11-Desoxycorticosterone Acetate (Doca)
 - Aldosterone

Synthetic

GLUCOCORTICOIDS

Signs and Symptoms of Adrenal Crisis

- Headache / dizziness
- Low back pain
- Stomach / leg pain
- Pale skin/shivering
- Severe vomiting / diarrhea
- Lethargy / listlessness
- Loss of appetite
- Neurological deficits
- Confusion
- Low blood sugar
- Low blood pressure
- Seizures
- Cardiovascular collapse
- May present with shock-like symptoms

- 1. Short acting
- 2. Intermediate acting
- 3. Long acting
- 4. Inhalational

Short acting (8-12 hours)

> Hydrocortisone (cortisol)> Cortisone

Intermediate acting (12-36 hours)

- > Prednisone
- > Prednisolone
- Methylprednisolone
- Fluprednisolone
- Paramethasone
- Triamcinolone

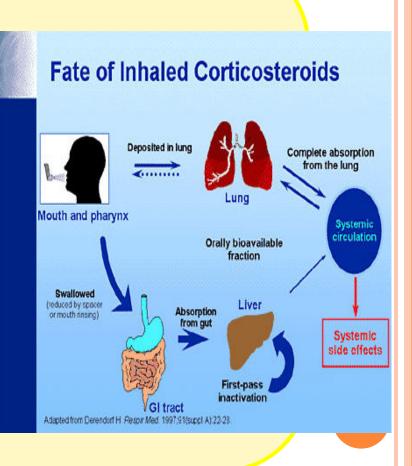
Long acting (36-72 hours)

- > Betamethasone
- > Dexamethasone
- Beclomethasone
- > Budesonide
- Fluticasone



Inhalational

- Triamcinolone
 Declamathered
- Beclomethasone
- > Budesonide
- Fluticasone
- Ciclesonide
- Flunisolide
- Mometasone



PHARMACOLOGICAL EFFECTS

- Corticosteroids are a class of steroid hormones that are produced in the adrenal cortex.
- Corticosteroids are involved in a wide range of physiologic systems such as
- stress response,
- immune response and regulation of inflammation,
- carbohydrate metabolism,
- protein catabolism,
- blood electrolyte levels, and
- behavior.

PHARMACOLOGICAL EFFECTS

(Extensions of physiological effects)

1. PHYSIOLOGIC EFFECTS

- > Direct effects
- > Permissive effects
 - Response of vascular & bronchial smooth muscle to catecholamines
 - Lipolytic response of fat cells to ACTH, catecholamines, growth hormones

PHARMACOLOGIC EFFECTS

2. METABOLIC EFFECTS

- Carbohydrate gluconeogenesis, glycogen, decrease glucose utilization, increase blood glucose levels
- Protein (catabolic & anti-anabolic effects) – in lymphoid tissue, muscle, fat, skin
- Lipid lipolysis, redistribution of body fat (different tissue sensitivity

PHARMACOLOGIC EFFECTS

- **3. ANTI-INFLAMMATORY EFFECTS4. IMMUNOSUPPRESSIVE EFFECTS**
- 5. CNS sense of well being, insomnia, restlessness, depression, psychosis
- 6. Skeletal muscle: required for normal function
- 7. CVS: mineralcorticosteroid –induced changes, enhance vascular reactivity to other vasoactive substances

ANTI-INFLAMMATORY & IMMUNOSUPPRESIVE ACTION

Reduced manifestations of inflammation

- Decreased release of vasoactive & chemo attractive factors
- Decreased secretion of lipolytic & proteolytic enzymes
- Decreased extravasation of leukocytes (increased neutrophils in blood, decreased lymphocytes)
- Decreased fibrosis
- Decreased expression of pro-inflammatory cytokines: COX2

Immunosuppressive & anti-allergic actions

- Suppresses all types of hypersensitivity & allergic phenomenon
- At High dose: Interfere with all steps of immunological response
- Causes greater suppression of CMI (graft rejection & delayed hypersensitivity)
- Transplant rejection: ↓ antigen expression from grafted tissues, delay revascularization, ↓ sensitisation of T lymphocytes etc.

CELL TYPE	FACTOR	COMMENTS
Macrophages and monocytes	Arachidonic acid and its metabolites (prostaglandins and leukotrienes)	Mediated by glucocorticoid inhibition of COX-2 and PLA ₂ .
	Cytokines, including: interleukin (IL)-1, IL-6, and tumor necrosis factor-α (TNF-α)	Production and release are blocked. The cytokines exert multiple effects on inflammation (e.g., activation of T cells, stimulation of fibroblast proliferation).
	Acute phase reactants	These include the third component of complement.
Endothelial cells	ELAM-1 and ICAM-1	ELAM-1 and ICAM-1: critical for leukocyte localization.
	Acute phase reactants	Same as above, for macrophages and monocytes.
	Cytokines (e.g., IL-1)	
	Arachidonic acid derivatives	
Basophils	Histamine, LTC ₄	IgE-dependent release inhibited by glucocorticoids.
Fibroblasts	Arachidonic acid metabolites	Same as above for macrophages and monocytes. Glucocorticoids also suppress growth factor– induced DNA synthesis and fibroblast proliferation.
Lymphocytes	Cytokines (IL-1, IL-2, IL-3, IL-6, TNF-α, GM-CSF, interferon-1)	Same as above for macrophages and monocytes.

Glucocorticoids - Pharmacokinetics

- Therapeutically given by various routes orally, IM, IV, topically
- Hydrocortisone undergoes high first pass metabolism
- Oral bioavailability of synthetic corticoids is high
- Both, endogenous and therapeutically administered GC are bound to Corticosteroid Binding Globulin (CBG)
- Synthetic steroids have to undergo reduction in liver to active compounds
- Metabolized in liver and excreted in urine
- Exogenously administered hydrocortisone has t1/2 of 1.5 Hrs

PREPARATIONS

Oral: Tablets
Parenteral: Dexame Hudroo

Dexamethasone Hydrocortisone Pulse therapy = Methyl prednisolone 1 gm IV OD * 3 days Depot = Triamcinolone Acetonate

Respiratory:

Nebulizers Revolizers Inhalers

Topical:

Creams/Lotions/Solutions

🗅 Eye, ear & nasal

Creams/Lotions/Solutions

POSOLOGY & BIO EQUIVALENCE

- Usual dosage ==== 1 mg/kg/day
- Preferably in a single morning dose
 - Impractical !!!!!!!!!
- Interconversions
- 1 mg Dexamethasone = 10 mg Prednisolone
- 1 mg Prednisolone = 25 mg Hydrocortisone
- Injection Hydrocortisone == 100mg/250mg/500mg
- Tab. Prednisolone == 5 mg
- Tab. Dexamethasone == 0.5 mg
- Injection 1ml Dexamethasone == 4 mg
 - = 8 Tablets each of Prednisolone & Dexamethasone

Glucocorticoids - MOA

Not stored:

rate of synthesis = rate of release

- Synthesize rhythmically and controlled by irregular pulses of ACTH, influenced by light and major pulses occur early in the morning and after meals
- Glucocorticoids act via their receptors located in nucleus (GR)
- GRs are widely distributed and located almost in all cells of the body
- They are made up of almost 800 amino acids

Glucocorticoids - MOA

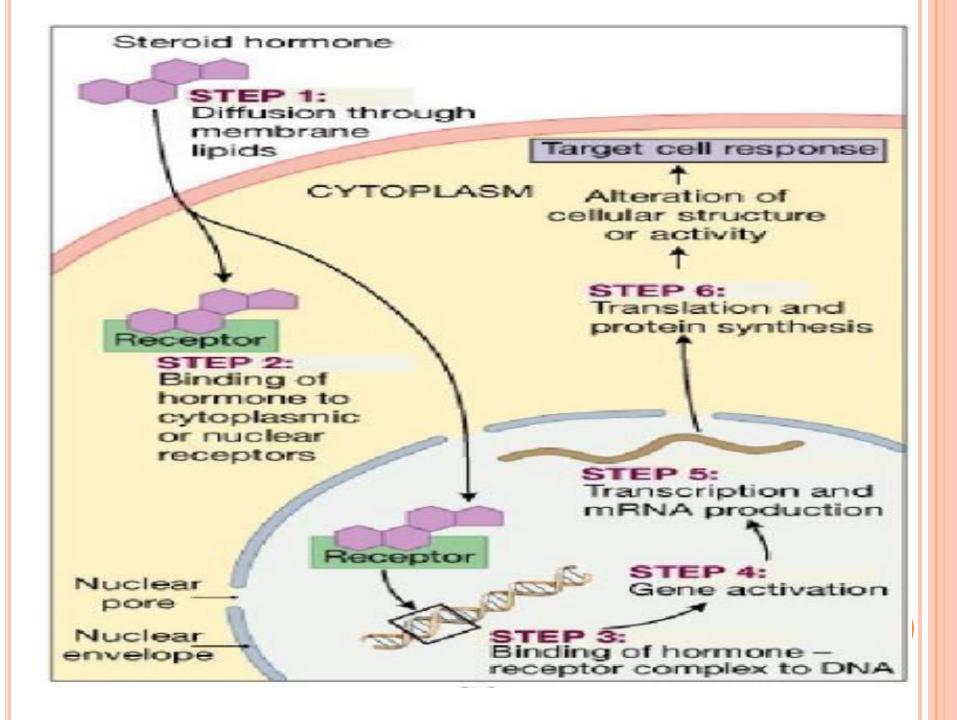
- GR receptors are located in the cytoplasm
- One GR receptor has a DNA binding domain and a ligand binding domain along with stabilizing proteins (HSP 90 and HSP 70)
- This receptor is incapable of activating transcription
- Binding of free steroid molecule to GR forms an unstable compound
- Therefore HSP and other proteins get dissociated
- The S+GR complex enters the nucleus and binds to Glucocorticoids response element (GRE) on gene and regulate transcription by RNA polymerase II and others
- The resulting mRNA is transported to cytoplasm for production of protein and bring about final response

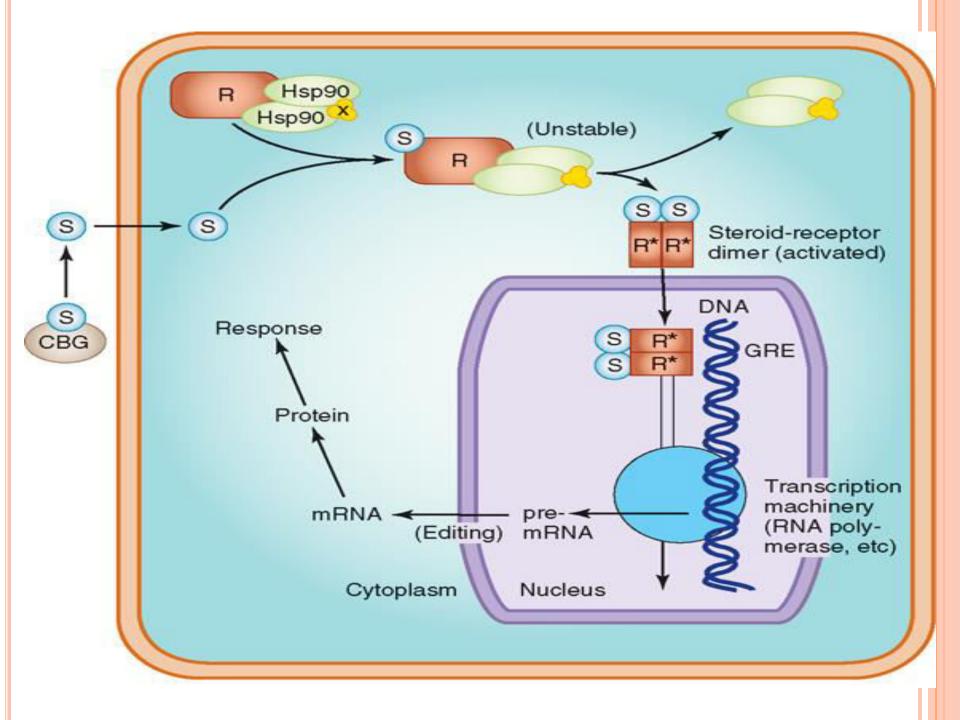
Hormone enters cell & binds to glucocorticoid receptor

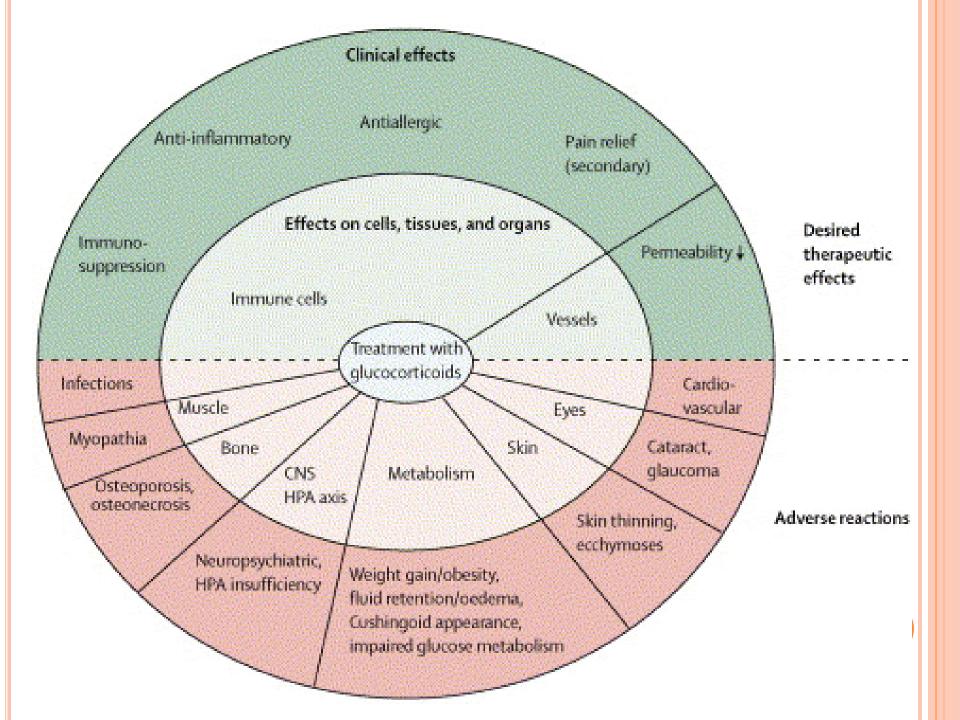
Conformational changes that allow it to dissociate from the Heat shock protein(HSP)

Dimerization - dimeric ligand-bound receptor complex is transported into the nucleus

Interacts with DNA & nuclear proteins through GREs in the promoter of responsive genes







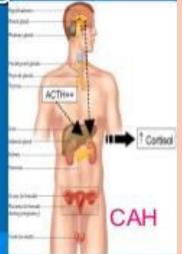
THERAPEUTIC USES

- 1. Diagnosis & treatment of disturbed adrenal function
- 2. Prevention of infant respiratory distress syndrome
- 3. Non-adrenal disorders

Replacement Therapy

Adrenal insufficiency – acute/chronic

- Abrupt withdrawal of steroid therapy
- Chronic infections Tuberculosis
- Autoimmune adrenal disease
- Surgery, Hemorrhage and AIDS
- Congenital adrenal hyperplasia
 - Congenital disorder due to deficiency of 21hydroxylse enzyme – no cortisol but ACTH – increased androgen production



Addisonian Crisis

Features:

- Severe shock hypotension, tachycardia
- Fever, abdominal pain, nausea & vomiting
- Hyponatraemia/hyperkalaemia ±hypercalcaemia, hypoglycaemia

Management:

ABCDE assessment

- Correct volume depletion
- Replace glucocorticoids
- Correct metabolic abnormalities
- Treat underlying cause

100mg hydrocortisone I/V every 8hrs

Replacement Therapy

Acute adrenal insufficiency

- IV replacement of sodium chloride and fluid
- IV hydrocortisone 100 mg stat followed by100 mg every 8 Hrs – maximal daily rate of secretion

(alternatively, dexamethasone can be used)

- Chronic adrenal insufficiency
 - Hydrocortisone
 - Prednisolone or dexamethasone long acting
 - Fludrocortisone for mineralocorticoid effects
- Congenital adrenal hyperplasia
 - Hydrocortisone 0.6 mg/kg in divided doses to maintain feedback suppression

EMERGENCY USES

- ✤ Acute severe asthma
- Raised intracranial pressure
- Septic shock
- ✤ Acute adrenal insufficiency
- Anaphylactic shock -2^{nd} choice

NON-ADRENAL DISORDERS

1. Allergic reactions: □ Urticaria □ Angio -neurotic edema **Contact** dermatitis □ Bee stings □ Serum sickness, Drug reactions

Allergic Disorders

- Exhibit a delayed response in allergies (1-2 hrs even in IV injection)
- In anaphylaxis, angioneurotic oedema and serum sickness etc. – adrenaline is the choice
- Seasonal allergies, bee sting, drug allergies
 - Allergic reactions can be suppressed by corticosteroids as supplements
- Intranasal administration in allergic rhinitis budesonide and flunisolide

NON-ADRENAL DISORDERS

2. Anti-Inflammatory& Immunosuppression:

- Inflammatory Arthritides; RA, SLE, Gout, Reactive Arthritis
- Bronchial Asthma
- Inflammatory Bowel Disease
- Bechet's syndrome
- Nephrotic syndrome
- Organ transplant
- **G** Subacute thyroiditis
- Sarcoidosis, asthma

Anti-inflammatory Uses

For suppression of inflammatory components in

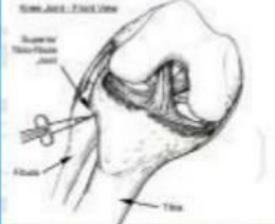
- Rheumatoid arthritis as adjuvant with NSAIDs in severe cases
- Osteoarthritis NSAIDs, intra-articular injection
- Rheumatic fever severe cases with carditis and CHF
- Gout NSAID failed cases and colchicine failed cases – intra-articular injection
- Vasculitic disorders: Polyarteritis nodosa

Intra-articular Steroids

Can be used in inflammatory Non-inflammatory diseases

- Knee joint
- Shoulder joint
- Tennis elbow
- Carpal tunnel syndrome





Bronchial Asthma

- The increased recognition of the immunological and inflammatory nature of Bronchial asthma has led to the use of corticosteroids
- In severe asthma attacks IV hydrocortisone Methylprednisolone Oral prednisolone
 - Acute attacks:

>



*Inhaled beclmethasone, budesonide, flunisolide alone or combined with beta-2 agonists/ipratropium *Oral steroids

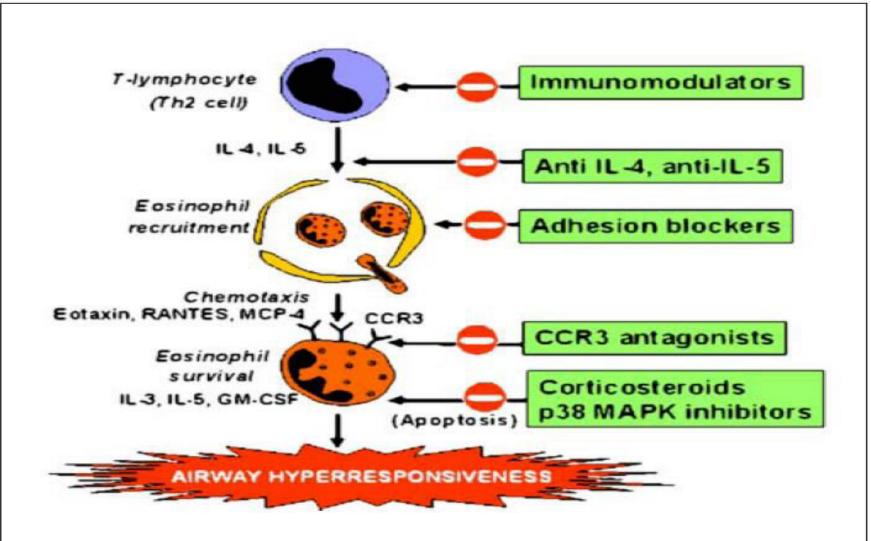


Figure 1. Inhibition of eosinophilic inflammation. Several strategies are possible to inhibit eosinophil inflammation in tissues, including immunomodulators (e.g., cyclosporin), inhibitors of proinflammatory cytokines (e.g., IL-4 and IL-5), inhibition of critical adhesion molecules (e.g., VLA4), blockade of chemokine receptors on eosinophils (e.g., CCR3) and induction of apoptosis (e.g., by corticosteroids and p38 MAP kinase inhibitors).



Inflammatory conditions of intestine like

- Ulcerative colitis
- Crohn`s disease
- Coeliac disease

(oral therapy or retention enema with hydrocortisone)

May mask the major complications like perforation and peritonitis

Autoimmune diseases



 Autoimmune haemolytic anaemia
 Idiopathic thrombocytopenic purpura
 Active chronic hepatitis, alcoholic hepatitis
 (Prednisolone 1-2 mg/kg/day given till remission followed by gradual withdrawal or low dose maintenance)

Renal diseases



Nephrotic syndrome in children Renal disease secondary to SLE Renal sarcoidosis Glomerulonephritis – membranous type (Life saving importance - usually given in large doses followed by tapering to maintenance dose)

Ocular Diseases



Important drug therapy for suppressing inflammation in eye and preservation of sight

- Topical instillations are used for conditions of the anterior chamber – allergic conjunctivitis, iritis, iridocyclitis and keratitis etc.
- Systemic steroids for the posterior chamber
- Dexamethasone topical 0.1%
- Prednisolone oral
- Contraindicated in viral, fulminant bacterial infections, fungal infections and injuries

NON-ADRENAL USES

3. Stimulation of fetal lung maturation (RESPIRATORY DISTRESS SYNDROME)

Treatment of mother with large dose of glucocorticoids

> BETAMETHASONE I/M to mother

NON-ADRENAL USES

- 4. Infections: Gram negative septicemia
- 5. Leukemia, lymphoma
- 6. Pemphigus and other skin diseases
- 7. Use in raised intracranial pressure
- 8. Mountain sickness
- 9. Hypercalcemia
- 10. Multiple sclerosis

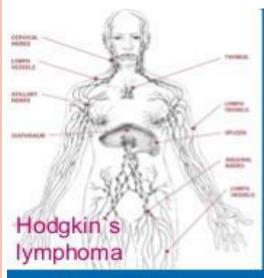
Infectious Diseases

- Indicated only in severe infective diseases to tide over crisis or prebent complications
 - AIDS and pneumocystis carinii pneumonia
 - In haemophilus influenza meningitis to reduce neurological complications
 - Tubercular meningitis
 - Lepra reaction
 - Scepticaemia

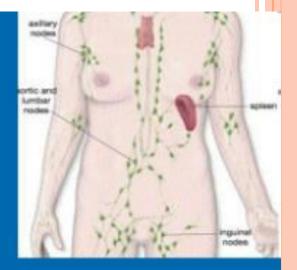


INDICATIONS OF CORTICOSTEROIDS (ALONG WITH ATT) IN T.B

• Massive T.B pleural effusion • Massive T.B pericardial effusion • T.B ascites (massive) • T.B meningitis • T.B choroiditis



Malignancy



Essential for combined chemotherapy of

- Acute lymphatic leukemia
- Hodgkin's and other lymphomas
- Hormone responsive breast carcinoma

Symptomatic relief in other advance malignancies by improving appetite and controlling secondary hypercalcaemia

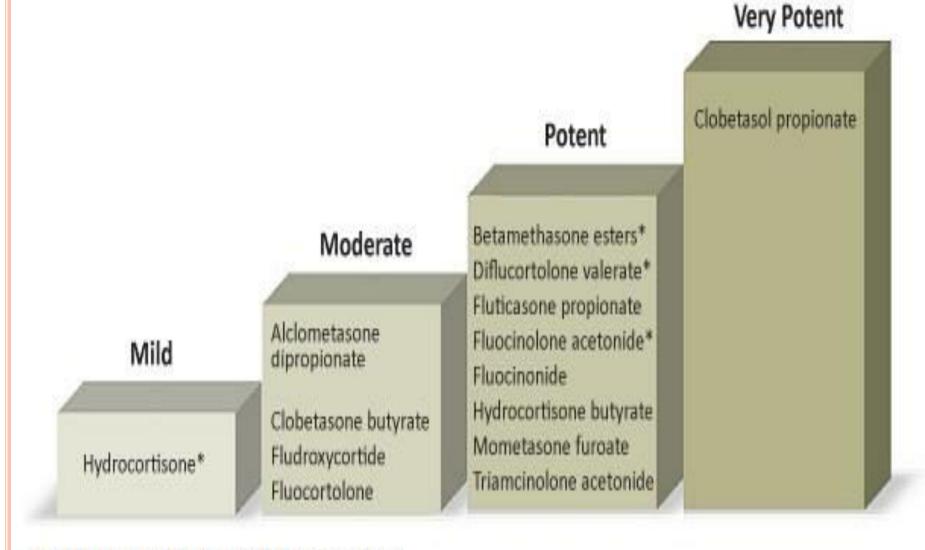
Skin Diseases

Pemphigus vulgaris

ADAM

 The largest application of steroid therapy
 Topical forms are widely used in many eczematous skin diseases
 Systemic therapy are also required and

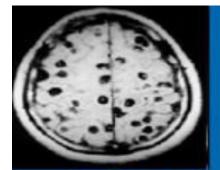
- may be life saving in
 - Pemphigus vulgaris
 - Exfoliative dermatitis
 - Stevens-Johnson syndrome



Potency of steroid creams

Potency	Generic name	Delivery vehicle and trade name
Class 1: Weak	Hydrocortisone acetate 1%	cream (Biocort® ^f ; Stopitch® ^g)
	Hydrocortisone acetate 0,5%	cream, ointment (Dilucort®h)
	Hydrocortisone 1%	cream (Procutan®°;Vari-Hydrocortisone®)
	Hydrocortisone 0.5%	cream (Skincalm® ^h)
Class 2: Moderately Potent	Betamethasone valerate 0.05%	cream (Betnovate Half Strength®)
Class 3: Potent	Fluticasone propionate 0.05%	Cream (Cutivate® ^k)
	Fluticasone propionate 0.005%	Ointment (Cutivate®*)
	Hydrocortisone butyrate 0.1%	cream, ointment, lotion (Locoid®) topical (Locoid Crelo®)
	Betametasone valerate 0.1%	cream (Betnovate® ⁱ ; Repivate®e; Adco- Betamethasone® ^e ; Vari-Betamethasone® ^b) ointment, lotion (Betnovate® ⁱ ; Lenovate® ^h ; Persivate® ^h)
	Fluocinolone acetonide 0,025%	ointment, gel, cream (Synalar®) cream, ointment (Cortoderm® ^h)
	Beclomethasone dipropionate 0.025%	cream (Beclate®®)
	Diflucortolone valerate 0.1%	ointment, cream (Nerisone®d)
	Methylprednisolone aceponate 0.1%	cream, ointment, milk (Advantan®d)
	Betamethasone dipropionate 0.05%	cream, ointment (Diprosone®°)
	Mometasone furoate 0.1%	cream, ointment, lotion (Elocon®°) cream (Aspen Mometasone® ^h ; Mometagen® ^p)
Class 4: Very Potent	Clobetasol propionate 0.05%	cream, ointment (Dermovate®; Dovate®h; Xenovate®h)

'Akacia HealthCare, Isando, South Africa; [®]Adcock Ingram, Bryanston, South Africa; [®]Aspen Pharmacare, Woodmead, South Africa; [®]Specpharm Holdings, Halfway House, South Africa; [®]MSD/Schering-Plough South Africa, Midrand, South Africa; [®]Sekpharma, Sandton, South Africa; [®]GlaxoSmithKline South Africa, Bryanston, South Africa; ^IAstellas Pharma, Bedfordview South Africa; [®]Cipla SA, Bellville, South Africa; ^IGlenmark Pharmaceuticals South Africa, Midrand, South Africa; [®]Bayer, Isando, South Africa; [®]Mylan South Africa, Modderfontein, South Africa



Cerebral Oedema



Cerebral oedema due to tumors (neoplasms) Traumatic and poststroke oedema (?) (Dexamethasone or betamethasone is preferred because no Na+ retaining activity) Other CNS conditions - spinal chord injury, Bell's palsy and neurocysticercosis Oral Prednisolone is the preferred drug)



Other Uses



HACE on MRI

Antiemetic – with ondansetron
 Acute mountain sickness
 Aspiration pneumonia, pulmonary oedema from drowning
 Hyperthyroidism – thyroid storm

Adverse Effects

Two types:

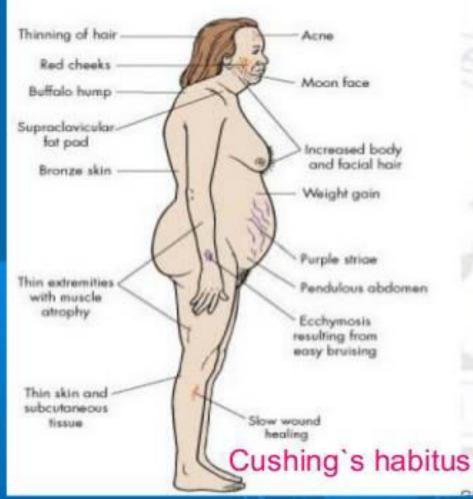
- From abrupt withdrawal
- Chronic therapeutic use of high dose
- Withdrawal
 - Flare up of underlying disease
 - Suppression of HPA axis and acute adrenal insufficiency
 - Increased ICT and papilloedema

WITHDRAWAL

- ✓ Not to be stopped abruptly if patient is on hydrocortisone [>25 mg/day] or for > 2-3 weeks
- Severe adrenal crisis & death of patient due to suppression of HPA axis
- ✓ Use short acting steroids with lowest possible doses form
- Prescribe the whole daily dose of the drugs at once in the morning
- Switch to alternate day therapy if possible
 After the long term use intermediate acting steroids allow for more flexible withdrawal

✤ Long term therapy – immunosuppression HPA axis suppression Addison-like symptoms > Tapering the dose > Physiological daily replacement (5mg prednisone) until adrenal function is restored

Adverse Effects



- High blood sugar
- High blood pressure
- Vertigo
- Blurred vision
- Female balding
- •Menstrual irregularities
- Hirsutism
- Severe depression
- Cognitive impairment
- Emotional instability

Easy fatiguability







Major Side Effects Associated with Corticosteroid Therapy

Dermatologic and soft tissue

Skin thinning and purpura Cushingoid appearance Alopecia Acne Hirsutism Striae Hypertrichosis

Eye

Posterior subcapsular cataract Elevated intraocular pressure/glaucoma Exophthalmos

Cardiovascular

Hypertension Perturbations of serum lipoproteins Premature atherosclerotic disease Arrhythmias with pulse infusions

Gastrointestinal

Gastritis Peptic ulcer disease Pancreatitis Steatohepatitis Visceral perforation

Renal

Hypokalemia Fluid volume shifts Genitourinary and reproductive Amenorrhea/Infertility Intrauterine growth retardation

Intrauter in Rone

Osteoporosis Avascular necrosis

Muscle

Myopathy

Neuropsychiatric

Euphoria Dysphoria/depression Insomnia/akathisia Psychosis Pseudo tumor cerebri

Endocr ine

Diabetes mellitus Hypothalamic-pituitary-adrenal insufficiency Infectious disease Heightened risk of typical infections

Opportunistic infections Herpes zoster

- ♦ Supraphysiological doses for > 2-3 wks
 ✓ Extensions of pharmacological effects
 - Metabolic
 - Cushing's syndrome
 - > GIT
 - Muscle
 - > Bone
- ♦ Withdrawal of therapy▶ HPA axis suppression

- ≻ Eye
- Electrolytes
- Water retention
- > CNS
- > Growth

Metabolic

- > Hyperglycemia, glycosuria
- Moon facies, obesity, hyperlipidemia
- Increased appetite
- Skin Increases growth of fine hair, thinning of skin, with striae & bruising, punctate acne
- GIT peptic ulcer, pancreatitis, NV
 Muscle muscle wasting, myopathy, growth retardation in children

- CNS insomnia, depression, acute psychosis
- ✤ Eye cataract, raised IOP
- Bone Osteoporosis, aseptic necrosis of hip
- Fluid & electrolytes Water retention, hypokalemia, alkalosis, hypertension
 BLOOD – Lymphopenia, leukocytosis

CONTRAINDICATIONS

> Diabetes mellitus > Peptic ulcer > Hypertension > Pregnancy > Osteoporosis > Tuberculosis & other infection > Epilepsy Renal failure > CCF

> Glaucoma
> Cushing's syndrome
> Herpes simplex keratitis
> Psychosis

MISUSE IN SPORTS

Normal dose _____ relaxation effects on the respiratory tract
 In larger dose _____ analgesic effects.

Enables athletes for better training and sporting performance.

Glucocorticoids are catabolic steroids.

WS _____ anabolic steroids = increase muscle mass and strength.

CLINICAL PHARMACOLOGY

Adrenocorticosteroid Inhibitors

- Metyrapone: 11 beta-hydroxylase enzyme inhibitor used in Cushing's syndrome and test of pituitary efficiency
- Aminoglutethemide: Stops conversion of cholesterol to pregnelone (Medical adrenalectomy) – Breast cancers
- Mifepristone: Progesterone antagonist
- Spironolactone: Aldosterone antagonist
- Ketoconazole: Inhibits synthesis of all hormones in testes and adrenal cortex – used in Cushing's syndrome and also in hirsutism in female

Prednisolone 5mg is equivalent to Hydrocortisone 20mg Methylprednisolone 4mg Betamethasone 750µg Dexamethasone 750µg 0 Cortisone acetate 25mg Deflazacort 6mg Triamcinolone 4mg

REFERENCES

- Basic and Clinical Pharmacology: Katzung BG, Masters SB, Trevor AJ. 14th Edition.
- Katzung & Trevor's Pharmacology: Examination & Board Review. 12th Edition
- Lippincott's Illustrated Reviews: Pharmacology, Clark MA, Finkel R, Rey JA, Whalen K. 7th Edition
- Goodman & Gilman's The Pharmacological Basis of Therapeutics: Brunton LL. 12th Edition

Email address for queries on the topic

drshams11@hotmail.com