# **PSORIASIS**

## **Background**

- 1. Definition: Chronic relapsing dermatitis of unknown etiology typically characterized by erythematous, sharply demarcated papules and plaques with silvery white scales (plaque psoriasis) although other forms exist (see below).
- 2. General Information: 1,2
  - o Common form: Plaque Psoriasis
    - 80-90% of Psoriasis cases. Fig. 1
    - Occurs at any age. Most common between ages 20-30 and 50-60
    - Classic erythematous oval plaques with silvery scale on extensor surfaces
    - Chronic once develops; follows waxing and waning course
  - o Less common forms:
    - Eruptive (aka Guttate Psoriasis)
      - ~ 10% of psoriasis cases. Fig. 2
      - Occurs alone or in combination with Plaque Psoriasis
      - Acute onset, association with stress/trauma/infection (Streptococcus infection most common)
      - 100s of plaques all over body with torso most common.
    - Pustular Psoriasis
      - Rare
      - Localized (Palmoplantar & Acropustulosis)
      - Generalized (Von Zumbusch)
    - Inverse Psoriasis
      - Rare; usually in obese patients with other psoriasis types
      - Shiny erythematous well defined plaque typically with macerated surface
      - Intertriginous areas (armpits, groin, under breasts)
    - Erythrodermic Psoriasis
      - Least common, 1-2% of psoriasis patients. Fig. 3
      - Seen in poorly controlled plaque psoriasis patients
      - Medical Emergency, treat like a burn
      - Severely painful and pruritic

### Pathophysiology & Epidemiology

- 1. Pathology of Disease: 1,3
  - o Not completely known, multifactorial
  - Abnormal T Lymphocyte function
  - Accelerated epidermopoiesis
- 2. Incidence, Prevalence:<sup>4</sup>
  - o Incidence = 200,000 (+/- 50,000) new cases in US yearly
  - United States Prevalence = Adults 2.2% (4.5million)
- 3. Risk Factors
  - No gender preference
  - o Family History. 1/3 patients have affected family members
  - o Caucasians 2x risk than African Americans

- 4. Morbidity / Mortality:
  - o Rare mortality w/ Plaque Psoriasis; typically from systemic therapy effects<sup>5</sup>
    - Psoriasis independent cardiovascular risk factor<sup>6,7</sup>
    - With exception of severe psoriasis and psoriatic arthritis, no affect on life expectancy or general health.<sup>8</sup>
  - o Varying morbidity:
    - Pruritis, psoriatic flares vary per patient
    - Psychosocial impact can be severe
      - Increased depression & suicidal ideation 9,10
    - Psoriatic arthritis

# Diagnostics<sup>1,2</sup>

- 1. History:
  - Typically, young adult with symmetric, asymptomatic (pruritis may be present) erythematous oval plaques (1-10cm) on scalp, bilateral extensor surfaces and back.
- 2. Physical Examination:
  - Skin Findings
    - Erythematous, well defined papules that group to form stable plaques. Fig. 1
    - Extensor (elbows, knees) and back most common. Also check intertriginous areas and external ear canals. Palms/Soles/Face typically spared.
    - Silvery white scale yields bleeding spots when removed (Auspitz sign)
  - Nail Findings
    - Diagnostic support of psoriasis if nail findings also noted
    - Pitting nail plates most common. Fig. 4
    - Also see Onycholysis, Beau lines, leukonychia, splinter hemorrhages
    - Oil drop sign specific for Psoriasis<sup>11</sup>
  - Joint Disease
    - Sausage Finger and nail findings. Fig. 4
    - Varying prevalence; ranges from 5-30% of psoriasis patients 1,12,13
    - Rheumatoid factor negative
    - Typically skin involvement precedes joint involvement.
    - Several types:
      - Asymmetric oligoarticular. Most common at 70%
      - Spinal Type. Debilitating. Approximately 20%
      - Others: Mutilating type, Distal Interphalangeal, Symmetric Polyarthritis
- 3. Diagnostic Testing
  - o Typically clinical diagnosis
  - o Rarely skin biopsy needed
    - Uncertain clinical diagnosis
    - Small punch biopsy adequate
- 4. Laboratory evaluation:
  - Rarely Indicated
  - o Severe cases, check uric acid (elevated) and folate (decreased)

- 5. Diagnostic imaging
  - Plain films only if joint involvement
- 6. Diagnostic Criteria:
  - o No formal diagnostic criteria
  - o Clinical diagnosis based on exam or, if uncertainty exists, skin biopsy
  - o Consider Dermatology consult if diagnosis in doubt
  - Psoriasis Area and Severity Index (PASI)
    - Quantifies subjective severity typically for research purposes
    - Clinical utility limited

## **Differential Diagnosis**

- 1. Key Differential Diagnoses: 1,2
  - o Seborrheic dermatitis (typically more face involvement)
  - o Eczema
  - o Tinea infection (KOH to rule out)
  - o Pityriasis rosea (Herald patch / Christmas tree pattern)
  - Candidiasis (KOH to rule out)
  - Lichen planus
  - o Drug eruption
- 2. Extensive Differential Diagnoses
  - o Paget's disease
  - o Cutaneous lupus erythematosus
  - o Bowen disease (Squamous Cell Carcinoma in situ)

# Therapeutics 14,15,16

- 1. Three categories (Topical, Phototherapy, Systemic) typically tried in succession but can be combined
  - Topical Therapy
    - Emollients
      - Critical therapy for skin hydration and protection
      - Normalizes hyperproliferation and apoptosis
      - Used for mild to moderate plaque psoriasis
      - BID dosing, best after bathing. Allow 10-15 minutes to penetrate prior to other topical applications.
      - Excellent safety profile
      - Examples include lotions, creams, ointments (each with increasing lipid:H<sub>2</sub>O ratio and thus viscosity). Compliance is key; therefore, tailor to patient preference.
    - Keratinolytic Agents
      - Adjunctive therapy; limited data as monotherapy
      - Softens / removes scales
      - Increases absorption of other topical agents
      - Especially useful on thick scalp lesions
      - Representative example: Salicylic Acid

- Cautions: Do not use with other salicylate drugs or prior to UVB phototherapy.
  - Potential for systemic absorption and side effects if
    >20% body surface application &/or abnormal hepatic or renal function.
  - Not for children.

#### Steroids

- Mainstay of therapy. Monotherapy or combined with other topical, UV light or systemic agents
- Class II-VII steroids typically BID, duration of therapy unknown
- Class I (super potent) steroids 2-4 weeks of treatment (50g or less per week for Class I). (SOR:A)<sup>16</sup>
- Gradually reduce use based on clinical response
- Clinical trials typically of short duration; therefore, longterm efficacy and risks largely unknown.
- Tachyphylaxis to topical steroids well documented, so pulse steroids often utilized<sup>17</sup>
- Side effects:
  - o Local: Skin atrophy, telangiectasias, striae, rosacea
  - Systemic: Hypothalamic-Pituitary Axis (HPA) suppression (medium to high potency topical steroids).

## Vitamin D Analogues

- Examples: calcipotriene (Calcitrene®, Dovonex®), calcitriol (Vectical®)
- Monotherapy more effective than placebo (SOR:A)
- Combine with topical steroids for added benefit (SOR:A)
- No corticosteroid side effects.
- Transient skin irritation can occur.
- Rarely, elevated serum calcium at higher dosing.
- BID dosing to affected areas. Custom combinations with steroids and once daily also used.
- Calcitriol less irritating to skin than calcipotriene, especially intertriginous areas. Safe to use on face and genitals.
- Inactivated by acids; avoid use with salicylic acid and some steroid preparations with acids
- Use after phototherapy.
  - o Ultraviolet (UV) A inactivates calcipotriene,
  - UVA and UVB inactivates calcitriol<sup>18</sup>

#### Retinoids

- Adjunct to topical steroids or phototherapy for stable plaque psoriasis up to 20% body surface area (BSA)
- Apply once daily, typically evenings
- Causes photosensitivity and skin irritation
- If combining with phototherapy, reduce UV dose by >1/3
- Pregnancy category X

- Tar
  - Confine tar to plaques only
  - Tar irritates locally, stains clothes, malodorous
  - Falling out of favor due to compliance issues
- Phototherapy
  - Dermatologist guided therapy
  - Typically used for extensive psoriasis
  - Broad-band UVB (bbUVB) (SOR:C)
    - 290-320 nm. Causes skin burning
    - Typically combined with topical or systemic agents (SOR:B)
    - Frequency 3-5 times per week typically for 20-25 treatments
    - Improvement seen within one month; maintenance therapy needed by some
  - Narrow-band UVB (nbUVB) (SOR:B
    - 311 nm. Burns greater than bbUVB
    - Superior to broad-band UVB, safer than PUVA
    - Expense limits use
  - Photochemotherapy (PUVA)
    - Methoxsalen or psoralen (either orally or via bath) + UVA = PUVA
    - 2-3 times per week for 20-30 treatments
    - One of the most effective treatments for plaque psoriasis
    - Reports of cutaneous malignancy in Caucasians has limited use
- Systemic Therapy
  - Best managed by Dermatologist
  - Consider for >20% body surface involvement &/or pt very uncomfortable
  - Useful guide on systemic therapy including biologics, their use, monitoring and side effects: <u>Psoriasis.org Pocket Guide</u>
  - Biologics:
    - TNF alpha inhibitors (Adalimumab, Etanercept, Golimumab, Infliximab)
    - T Cell modulators (Alefacept)
    - Cytokine modulators (Ustekinumab)
  - Systemics:
    - Synthetic retinoids Acitretin (*Soriatane*®)
    - Immunosuppresives Cyclosporine, Methotrexate
- 2. Further Management (within 24 hrs)
  - o Rarely indicated
  - Most treatments are chronic in nature
  - Initial high dose therapies for psoriasis flare warrant office follow up to monitor response and adjust dosing.
- 3. Long-Term Care
  - Dependent on therapy 14,16
  - o Remission is goal, adjust therapy accordingly

## Follow-Up

- 1. Return to Office
  - o Interval of "regular" or "periodic" in-office follow-up poorly defined in guidelines.
  - o Therapy dictates follow-up in office
  - o Therapy side effects warrant sooner follow-up
- 2. Refer to Specialist
  - Rheumatologist if Psoriatic arthritis
  - o Consider cardiology consult if cardiovascular co-morbidities
  - Dermatology once phototherapy, biologics or systemic therapy considered, or if patient not responding to therapies.
- 3. Admit to Hospital
  - o Rarely indicated
  - Generalized pustular psoriasis & Erythrodermic Psoriasis immediate medical attention

# **Prognosis**

- 1. No cure available
- 2. Relapse, response to therapy, severity individualized.
- 3. No consensus prognostic tools.
- 4. Early onset and family history are poorer prognostic factors.
- 5. Treatment aimed at reducing or eliminating symptoms for as long as possible<sup>2</sup>

#### **Prevention/Exacerbation**

- 1. No known primary prevention
- 2. Exacerbating factors<sup>1,2,3</sup>
  - o Human Immunodeficiency Virus (HIV)
  - Physical Trauma
    - Koebner phenomenon: Psoriasis plaques forming at site of physical trauma.
  - Infection (Streptococcus and Candida)
  - Drugs (Lithium, beta blockers, chloroquine, ACE inhibitors, Corticosteroid withdrawal, terbinafine)
  - Winter season (cold/dry environment)
  - o Sunburn
  - Alcohol consumption
  - o Emotional Stress

### **Patient Education**

- 1. Psoriasis Patient handout from familydoctor.org:
  - o <u>http://familydoctor.org/familydoctor/en/diseases-conditions/psoriasis.printerview.all.html</u>
- 2. National Psoriasis Foundation:
  - o http://www.psoriasis.org

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Fig. 1: Plaque Psoriasis. Used with permission. LTC Kimberly Wenner MD, Madigan Army Medical Center Dermatology. 2012.



Fig. 2: Guttate Psoriasis. Used with permission. LTC Kimberly Wenner MD, Madigan Army Medical Center Dermatology. 2012.



Fig. 3: Erythrodermic Psoriasis. Used with permission. MAJ Drew Reese MD, Madigan Army Medical Center Dermatology. 2012.



Fig. 4: Psoriatic Arthritis with Sausage Finger and Pitting Nail of 3<sup>rd</sup> digit. Used with permission. LTC Kimberly Wenner MD, Madigan Army Medical Center Dermatology. 2012.

Psoriasis.org Pocket Guide <a href="http://www.psoriasis.org/document.doc?id=354">http://www.psoriasis.org/document.doc?id=354</a>

Authors: Matthew T. Porter, MD, & Thomas C. Michels, MD, Madigan Army Medical Center, Tacoma, WA

Editor: Robert Marshall, MD, MPH, MISM, CMIO, Madigan Army Medical Center, Tacoma, WA