

# Psoriasis and Psoriatic Arthritis Combined Clinic Electronic Medical Records Form

NAME: AGE: SEX: CHIEF COMPLAINT:	
HISTORY OF PRESENT ILLNESS: The patient presents for the initial expsoriasis for years. Areas of invo	valuation of [psoriasis / psoriatic arthritis]. The patient reports a history of lvement to date include:
	scalptrunkextremitiesnailintertriginous/inverse (body folds)genitalpalmo-plantarpustular(other):
If the patient reports "known" history of The patient reports history of psoriajoint painjoint stiffnessswelling of whole digit (dactylitis)neck painback pain Current treatment listed below. Psoriasis History Description:	of psoriatic arthritis, please complete: atic arthritis for years. The patient referes history of:
Psoriasis type:plaquepustularerythrodermainverse	

nail



### Itch severity:

How would you rate your itching over the past week? Please select a number to indicate your answer.

0:	1	2	3	4	5	6	7	8	9	10: Worst
No itch										imaginable
										itch

## Current treatment for psoriasis:

Treatment	Details
Topical(s)	
Phototherapy	
Oral DMARDs	
Biologic DMARDs	

## Past treatment for psoriasis:

Treatment	Reason for cessation: incomplete response, failed, intolerance
Topical(s)	
Phototherapy	
Oral DMARDs	
Biologic DMARDs	



# **Psoriatic Arthritis History Description:** Joints involved: \_\_\_\_ Minutes morning stiffness: \_\_\_\_ Does the stiffness improve with exercise? [YES/NO] Pain: [YES/NO] Swelling: [YES/NO] Current treatment for psoriatic arthritis: **Treatment Details** Physical therapy **NSAIDS** Oral DMARDs **Biologic DMARDs** Intra-articular Past treatment for psoriatic arthritis: Treatment Reason for cessation: incomplete response, failed, intolerance Physical therapy NSAIDS Oral DMARDs Biologic DMARDs Intra-articular



Family History of Psoriasis / Psoriatic Arthritis:
(Note: Both psoriasis and PsA show high rates of familial aggregation, suggesting that both disorders have a strong genetic basis. Of note, the prevalence of PsA among first-degree relatives was 49 times higher than the prevalence in the general population.)
HISTORY OF COMORBIDITIES / CO-PREVALENT DISORDERS:
Inflammatory bowel disease:
Uveitis: Metabolic syndrome:
Fatty liver:
Cardiovascular disease:
Depression/Anxiety:
Other:
<b>HISTORY OF RELEVANT LAB RESULTS:</b> (i.e. RF, CCP, HLA-B27, ESR, CRP; known imaging changes of inflammatory arthritis / PsA eg erosive disease on xray)
SOCIAL HISTORY:
PAST MEDICAL HISTORY:
MEDICATIONS:

### PSA SCREENING - PSORIASIS EPIDEMIOLOGY SCREENING TOOL (PEST)

If unknown history of psoriatic arthritis, administer the Psoriasis Epidemiology Screening Tool (PEST): Score 1 point for each question answered 'YES'. Note: a score of 3 or more indicates referral to rheumatology should be considered.

- 1. Have you ever had a swollen joint (or joints)? Yes/No
- 2. Has a doctor ever told you that you have arthritis? Yes/No
- 3. Do your finger nails or toenails have holes or pits? Yes/No
- 4. Have you had pain in your heel? Yes/No
- 5. Have you had a finger or toe that was completely swollen and painful for no apparent reason? Yes/No

Completed: [YES / NO]
If YES, total Score (0-5): \_\_/5



ASSESSMENT:		
General Exam:		

#### Skin Exam:

A full physical examination was performed including scalp, head, eyes, ears, nose, lips, neck, chest, axillae, abdomen, back, buttocks, bilateral upper extremities, bilateral lower extremities, hands, feet, fingers, toes, fingernails, and toenails.

Current psoriasis type: Well demarcated erythematous plaques with overlying silvery scale involving \_\_\_\_\_

Overall 5-point Physician Global Assessment:

0:Clear

1: Almost Clear

2:Mild

3: Moderate

4:Severe

The Physician's Global Assessment of Psoriasis is scored on a 5-point scale, reflecting a global consideration of the erythema, induration and scaling across all psoriatic lesions. Average erythema, induration and scaling are scored separately over the whole body according to a 5-point severity scale (0 to 4) as defined by morphologic descriptors.

Erytehma	Induration	Scaling
0 = No evidence of erythema (post-in- flammatory hyperpigmentation and/or hypopigmentation may be present) 1 = Light pink 2 = Light red 3 = Red 4 = Dark, deep red	<ul> <li>0 = No evidence of plaque elevation</li> <li>1 = Barely palpable</li> <li>2 = Slight, but definite elevation, indistinct edges</li> <li>3 = Elevated with distinct edges</li> <li>4 = Marked plaque elevation, hard/sharp borders</li> </ul>	<ul> <li>0 = No evidence of scaling</li> <li>1 = Occasional fine scale</li> <li>2 = Fine scale predominates</li> <li>3 = Coarse scale predominates</li> <li>4 = Thick, coarse scale predominates</li> </ul>

BSA (1 patient palm = 1% BSA) = \_\_\_\_ % BSA x PGA = \_\_\_



#### PLAN:

### **Psoriasis Management:**

(Note: consider the following treat-to-target guidelines(T2T) Guidelines per the National Psoriasis Foundation:

BSA 0-3% mild, 3-10% moderate, >10% severe Target response BSA ≤1% at 3 months and 6 months Acceptable response of  $\leq 3\%$  at 3 months or 75% improvement from baseline) **Psoriatic Arthritis Management: Comorbidities Management:** Do you have a primary care physician (PCP)? [YES / NO] If YES, how frequently do you see your PCP? \_\_\_\_\_ **Major Comorbidities:** Cardiovascular Disease (CVD): (Note: Psoriatic disease is associated with an increased risk of myocardial infarction and stroke. Screening for CVD is recommended yearly by measuring blood pressure and serum lipids by PCP) Hypertension: (Note: hypertension begins at 130/80. American College of Cardiology Guidelines: http://www.acc.org/ latest-in-cardiology/ten-points-to-remember/2017/11/09/11/41/2017-guideline-for-high-blood-pressure-in-adults) Diabetes: Obesity: Depression and Anxiety:



Minor Comorbidities (i.e. osteoporosis, obesity, inflammatory bowel disease, ophtalmic disease, malignancy)

Immunization / High Risk Medication Screening  Vaccinations: (Note on vaccinations: Adminster yearly influenza vaccine to all patients, particularly those receiving immune suppression. Avoid live and live attenuated vaccines in patients receiving immunosuppression. Hepatitis B vaccinations is safe to administer in immunosuppressed patients.)
Pneumococcal Vaccines:  • Prevnar (13-valeant pneumococcal conjugate vaccine)
Pneumovax (13-valeant pneumococcal polysaccharide vaccine)
Administration recommendations: (1) A patient who has never been vaccinated for pneumococcal pneumonia should be given Prevnar, followed by Pneumovax 8–12 weeks later. The Pneumovax vaccine should be repeated 5 years later. (2) A patient who has already received Pneumovax should be given Prevnar 1 year later. (3) Patients previously vaccinated with Prevnar should receive Pneumovax 1 year later.
Hepatitis B and TB Screening:  Hepatitis B, C status:  HBV surface antigen (HBsAg)  HBV surface antibody (HBsAb)  HBV core antibody (HbcAb).
TB Screening: (Note on TB screening: screening tests include tuberculin skin test (TST); commonly referred to as a 'PPD') and the interferon gamma release assay (IGRA); commonly referred to as a T-spot or Quantiferon-Gold. If screening for TB reveals latent M. tuberculosis infection, the TB must be treated prior to initiation of immunomodulatory therapy.)
<ul><li>Date of last PPD:</li><li>Date of last T-spot:</li></ul>
Laboratory Monitoring for disease activity and Medication Toxicity Labs: BUN, Cre, LFTs, CBC with diff. Last checked:// Consider these 3 as relevant: CRP; ESR; anti-CCP, RF
Imaging Rx:
Follow up in

7

Cc

Primary care Rheumatologist