Psoriatic Arthritis: Disease State



Module 3 Diagnosis, Classification, and Risk Factors of PsA

How Is PsA Diagnosed?

Diagnosis of PsA

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There are no diagnostic tests; most often, it is a clinical diagnosis¹



Joints are assessed in a manner similar to that in rheumatoid arthritis²



Most patients do not demonstrate the presence of serological markers (ie, are seronegative)¹

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In clinical trials, the CASPAR criteria may be used to classify PsA¹



Detailed patient history, physical examination, laboratory findings, and imaging results are highly important¹



Imaging studies (ultrasonography and MRI) allow more precise visualization of joint damage and tissue inflammation¹

CASPAR=Classification Criteria for Psoriatic Arthritis; MRI=Magnetic Resonance Imaging; PsA=Psoriatic Arthritis. 1. FitzGerald O, et al. *Nat Rev Dis Primers*. 2021;7(1):59. 2. Rida MA, Chandran V. *Clin Immunol*. 2020;214:108390.

Differential Diagnosis of PsA

Clinical, Laboratory, and Radiographic Features That Distinguish PsA From RA and OA^{1,2}

| Feature | PsA | RA | OA |
|---|-------------|-------------|-------------------------|
| Inflammatory arthritis | Very common | Very common | Occasional |
| Peripheral involvement | Very common | Very common | Very common |
| Axial involvement (sacroiliac joint ^a , spine) | Common | No | No |
| Symmetrical involvement | Occasional | Common | Common |
| DIP joint involvement | Common | Rare | Common |
| Enthesitis | Common | Rare | No |
| Dactylitis | Common | No | No |
| Bone erosion ^a | Common | Very common | Occasional ^b |
| New bone formation ^a | Common | Rare | Common |
| Skin involvement | Very common | Rare | Rare |
| RF positive | Occasional | Very common | Rare |
| aCCP positive | Rare | Very common | Rare |
| Nail dystrophy | Very common | No | No |

^aIn disease of >2 years duration; ^bInflammatory OA may manifest as focal erosion and new bone formation. Very common=60–90%; Common=30-60%; Occasional=10–30%; Rare <10%; No=Not found. aCCP=Anti-cyclic Citrullinated Peptide; DIP=Distal Interphalangeal; OA=Osteoarthritis; PsA=Psoriatic Arthritis; RA=Rheumatoid Arthritis; RF=Rheumatoid Factor. 1. FitzGerald O, et al. *Nat Rev Dis Primers*. 2021;7(1):59. 2. Rida MA, Chandran V. *Clin Immunol*. 2020;214:108390

CASPAR: Classification Criteria for Psoriatic Arthritis

 To be classified as having PsA, a patient should have inflammatory articular disease (joint, spine, or entheseal) with a score of ≥3 points from the 5 categories listed in the following table^{1,2}

| Criterion ^{1,2} | Description ¹ | | | |
|--|---|--|--|--|
| 1. Evidence of psoriasis (one of a, b, or c; a maximum of 2 points for this criterion) | | | | |
| a) Current psoriasis ^a (2 points) | Psoriatic skin or scalp disease present currently, as is judged by a rheumatologist or dermatologist | | | |
| b) Personal history of psoriasis (1 point) | A history of psoriasis obtained from the patient, family physician, dermatologist, rheumatologist, or other qualified healthcare professional | | | |
| c) Family history of psoriasis (1 point) | A history of psoriasis in a first- or second-degree relative per the patient | | | |
| 2. Psoriatic nail dystrophy (1 point) | Typical psoriatic nail dystrophy, including onycholysis, pitting, and hyperkeratosis, as observed during current physical examination | | | |
| 3. Negative test result for RF (1 point) | By any method except latex, but preferably by ELISA or nephelometry, according to the local laboratory reference range | | | |
| 4. Dactylitis (one of a, b; 1 point) | | | | |
| a) Current | Swelling of an entire digit | | | |
| b) History | A history of dactylitis recorded by a rheumatologist | | | |
| 5. Radiologic evidence of juxta-articular new bone formation (1 point) | Ill-defined ossifications near joint margins (excluding osteophyte formation) on plain X-ray films of hand or foot | | | |

^aCurrent psoriasis is assigned a score of 2, and all other items a score of 1.

CASPAR=Classification Criteria for Psoriatic Arthritis; ELISA=Enzyme-Linked Immunosorbent Assay; PsA=Psoriatic Arthritis; RF=Rheumatoid Factor.

^{1.} Taylor W, et al. Arthritis Rheum. 2006;54(8):2665-2673. 2. Raychaudhuri SP, et al. J Autoimmun. 2017;76:21-37.

Who Develops PsA, and What Are the Risk Factors for the Disease?

Incidence and Prevalence of PsA

- In the United States, the overall incidence of PsA in patients with newly diagnosed PsO is 2.9 (95% CI, 2.9-3.0) events per 100 PY¹
- The prevalence of PsA in patients with¹
 - Mild PsO was 2.1 (95% CI, 2.1-2.1) events per 100 PY
 - Moderate PsO was 9.9 (95% CI, 9.5-10.4) events per 100 PY
 - Severe PsO was 17.6 (95% CI, 16.9-18.3) events per 100 PY
- About 20-30% of patients with psoriasis have PsA, and the prevalence estimates range widely, from 7% to 40%¹

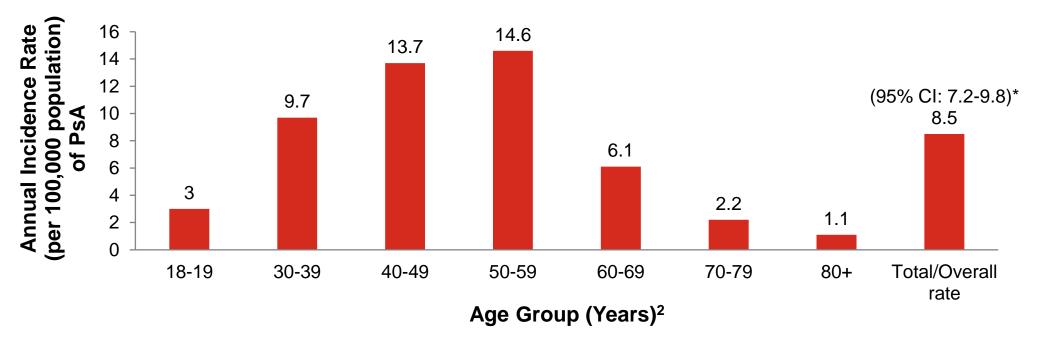


CI=Confidence Interval; PsA=Psoriatic Arthritis; PsO=Psoriasis; PY=Patient Years; US=United States.

1. Merola JF, et al. J Am Acad Dermatol. 2022;86(4):748-757. 2. Armstrong AW, et al. JAMA Dermatol. 2021;157(8):940-946. 3. Dures E, et al. Patient. 2017;10(4):455-462. 4. Inui K, et al. Modern Rheumatology. 2021;31(6):1179–1191.

Age at Onset

- The onset of PsA usually occurs in patients in their 30s and 40s (highest incidence is noted in those aged 40-59 years), and the prevalence is almost equal in men and women^{1,2}
- Median age at diagnosis is 44.8 years (interquartile range: 34.7-55.8 years)³



Annual incidence rate of PsA by age in Olmsted County, in the US state of Minnesota²

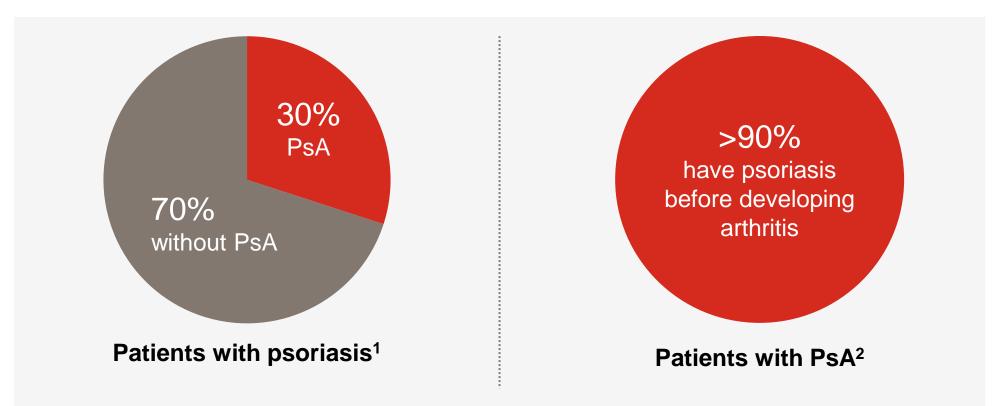
*Age- and sex-adjusted to the 2010 White population in the US.

CI=Confidence Interval; PsA=Psoriatic Arthritis.

1. Tiwari V, Brent LH. Psoriatic Arthritis. In: StatPearls [Internet]. Treasure Island, FL: StatPearls Publishing; 2022. https://www.ncbi.nlm.nih.gov/books/NBK547710/ (Accessed September 13, 2022). 2. Karmacharya P, et al. Arthritis Rheumatol. 2021;73(10):1878-1885. 3. Ogdie A, et al. Rheumatology (Oxford). 2013;52(3):568-575.

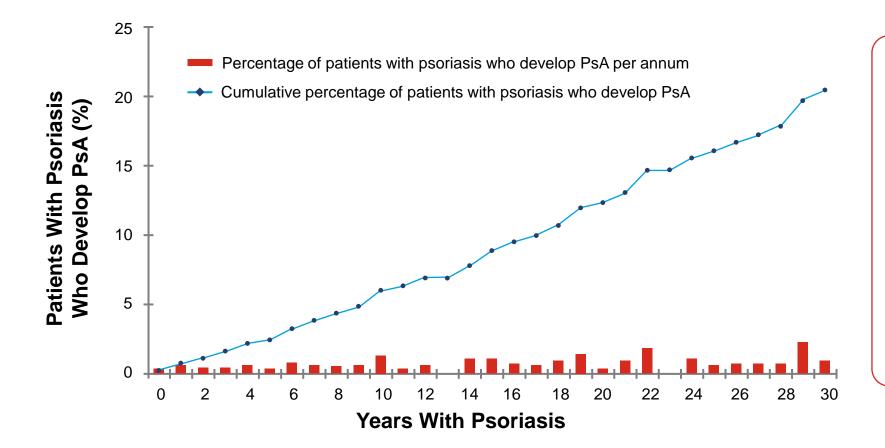
Psoriasis and PsA

Although the association between psoriasis and arthritis was first described in 1818, PsA was officially
recognized as a distinct disorder by the American Rheumatism Association only in 1964¹



Development of PsA in Relation to Psoriasis

Incidence and cumulative prevalence of PsA over time in population of patients with psoriasis

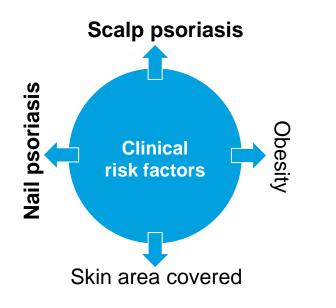


- During the 30 years of examination, incidence of PsA in the population of patients with psoriasis remained relatively constant, largely below 1% per year (74 per 1000 person-years)
- Among patients with a 30-year history of psoriasis, the prevalence of PsA increased steadily with disease duration reaching 20.5%

PsA=Psoriatic Arthritis. Christophers E, et al. *J Eur Acad Dermatol Venereol.* 2010;24(5):548-554.

Clinical Risk Factors for PsA

• Risk factors for PsA development include involvement of nail and scalp psoriasis



Nail Psoriasis

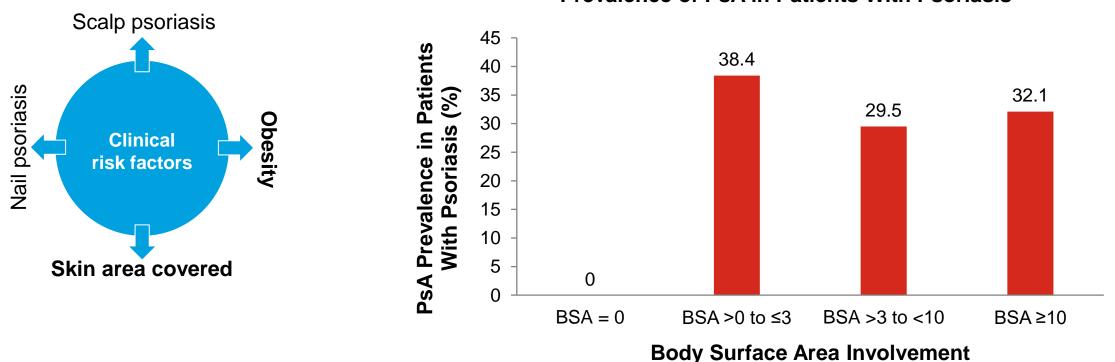


Scalp Psoriasis



Clinical Risk Factors for PsA

- Greater involvement of skin and obesity in patients with psoriasis increases the risk for PsA¹
- Obesity is more common in patients with psoriasis (36.5% vs. 22%; OR 2.1, 95% CI: 1.5-2.8, p<.01) and PsA (27.6% vs. 22%; OR 1.4, 95% CI: 1.0-1.9, p<.05) compared to non-inflammatory population²



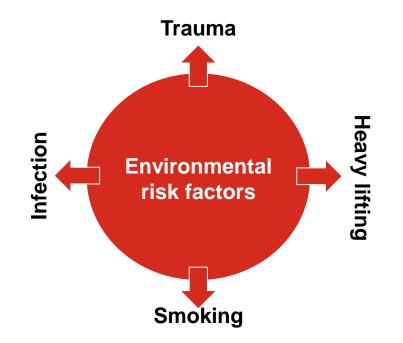
Prevalence of PsA in Patients With Psoriasis³

BSA=Body Surface Area; CI=Confidence Interval; OR=Odds Ratio; PsA=Psoriatic Arthritis.

1. Ogdie Å, Gelfand JM. Curr Rheumatol Rep. 2015;17(10):64. 2. Queiro R, et al. Medicine (Baltimore). 2019;98(28):e16400. 3. Tillett W, et al. Rheumatol Ther. 2020;7(3):617-637.

Environmental Risk Factors for PsA

- Infection, injury, smoking, and trauma may also increase the risk for development of PsA in patients with psoriasis^{1,2}
- The following factors are associated with the onset of PsA
 - Infection¹
 - Injury and heavy lifting¹
 - Smoking (increases the risk and leads to worse clinical outcomes)¹
 - Recent trauma (observed IR: 23.8 [95% CI: 22.3-25.3] per 10,000 patient-years)

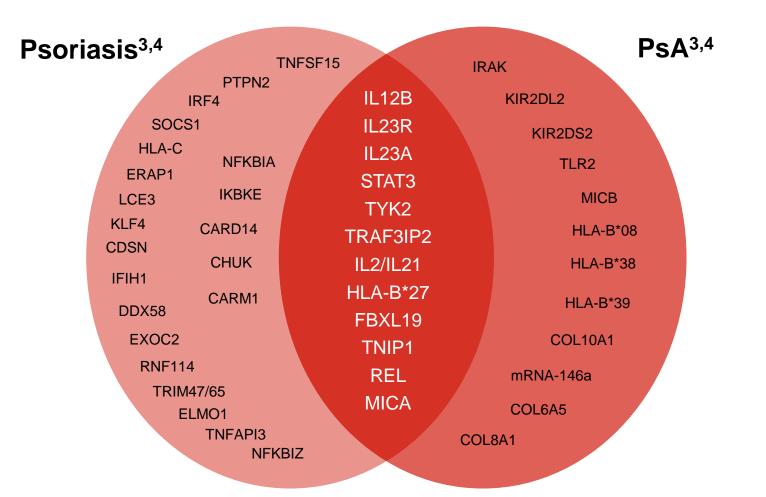


CI=Confidence Interval; IR=Incidence Rate; PsA=Psoriatic Arthritis.

1. Ogdie A, Gelfand JM. Curr Rheumatol Rep. 2015;17(10):64. 2. Thorarensen SM, et al. Ann Rheum Dis. 2017;76(3):521-525.

Genetic Risk Factors

- PsA prevalence is 49 times more likely among firstdegree relatives of patients with PsA than that in the general population¹
- Class I HLA genes are highly associated with PsA and account for ~30% of the genetic susceptibility²

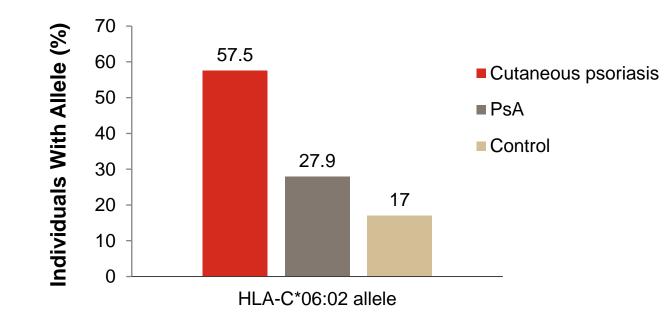


HLA=Human Leukocyte Antigen; PsA=Psoriatic Arthritis.

1. Giannelli A. Rheumatol Ther. 2019;6(1):5-21. 2. de Vlam K, et al. Acta Derm Venereol. 2014;94(6):627-634. 3. Dand N, et al. Acta Derm Venereol. 2020;100(3):adv00030. 4. Carvalho AL, Hedrich CM. Front Mol Biosci. 2021;8:662047.

Genetic Risk Factors

- PsA appears to be genetically distinct from psoriasis
- Although certain genes are shared between them, HLA-C*06:02, for example, is less common among people with PsA



Conclusions

There are no diagnostic tests for PsA, and most patients are seronegative. PsA is often a clinical diagnosis based on detailed patient history, physical examination, laboratory findings, and imaging results.¹

Clinical, laboratory, and radiographic features that distinguish PsA from RA and OA include, but are not limited to, axial involvement, dactylitis, enthesitis, and nail dystrophy.^{1,2}

Roughly 20%-30% of patients with psoriasis will have PsA,³ with the onset of PsA usually occurring in a patient's 30s and 40s.^{4,5}

Risk factors for PsA development include, but are not limited to, the presence of nail psoriasis, scalp psoriasis, PsO severity, and obesity.⁶

OA=Osteoarthritis; PsA=Psoriatic Arthritis; RA=Rheumatoid Arthritis.

1. FitzGerald O, et al. *Nat Rev Dis Primers.* 2021;7(1):59. 2. Rida MA, Chandran V. *Clin Immunol.* 2020;214:108390. 3. Merola JF, et al. *J Am Acad Dermatol.* 2022;86(4):748-757. 4. Tiwari V, Brent LH. Psoriatic Arthritis. *In: StatPearls* [Internet]. Treasure Island, FL: StatPearls Publishing; 2022. https://www.ncbi.nlm.nih.gov/books/NBK547710/ (Accessed September 13, 2022). 5. Karmacharya P, et al. *Arthritis Rheumatol.* 2021;73(10):1878-1885. 6. Ogdie A, Gelfand JM. *Curr Rheumatol Rep.* 2015;17(10):64.

US Medical Education

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