

Clinical Professor of Dermatology
Icahn School of Medicine at Mount Sinai, New York, NY
Indiana University Medical Center, Indianapolis, IN
Medical Director
Physicians Skin Care, PLLC Louisville, KY
DermResearch, PLLC, Louisville, KY
Skin Sciences, PLLC, Louisville, KY

#### I have received funding either as an investigator, consultant, or a speaker from the following pharmaceutical companies:

- Abbott
- **Acambis**
- **Aclaris**
- Allergan
- Almirall
- Amgen
- Anacor
- **Anaptys**
- **Astellas**
- Asubio
- BMS
- Berlex
- Biogen-Idec
- Biolife
- Biopelle
- Boehringer-Ingleheim
- Breckenridge Pharma
- Brickell
- Cellceutix
- Celgene
- Centocor
- Cipher
- Coherus
- Colbar
- Collagenex
- Combinatrix
- Connetics
- Coria
- Dermayant
- Dermik
- Dermira
- Dow
- **Dusa Pharmaceuticals**
- Eli Lilly
- Exeltis
- EOS
- Ferndale
- Galderma
- Genentech
- **Glaxo-Smith Kline**

- HealthPoint
- Idera
- Intendis
- Innovail
- Isdin
- Johnson & Johnson
- 3M
- Leo
- L'Oreal
- MC-2
- Maruho
- Medicis
- Merc
- Merz
- NanoBio
- Novartis
- Novan
- Nucryst
- Obagi
- **Onset Therapeutics**
- OrthoNeutrogena
- PediaPharma
- Pfizer
- **PharmaDerm**
- **Promius**
- Puracap
- QLT
- Quatrix
- Quinnova
- Serono
- SkinMedica
- Stiefel
- Sun Pharma
- Taro
- TolerRx
- **Triax Pharmaceuticals**
- UCB
- Valeant
- Warner & Chilcott
- Xenoport ZAGE

Andrew F. Alexis MD, MPH
Professor of Clinical Dermatology
Weill Cornell Medical College
New York, NY

### **Disclosures**

**Grants (funds to institution):** Leo, Novartis, Almirall, Bristol-Myers-Squibb, Amgen, Menlo, Galderma, Valeant (Bausch Health), Cara, Arcutis

Advisory board/Consulting: Leo, Galderma, Pfizer, Sanofi-Regeneron, Dermavant, Beiersdorf, Valeant, L'Oreal, BMS, Bausch health, UCB, Vyne, Arcutis, Janssen, Allergan, Almirall, Abbvie, Sol-Gel, Amgen, Lilly

Speaker: Regeneron, SANOFI-Genzyme, Pfizer, Astra Zeneca

Royalties (text books/other) - Springer, Wiley-Blackwell, Wolters Kluwer Health

**Equipment** – Aerolase (loan to institution)

## HYPERPIGMENTARY CHANGES

- •TOPICAL STEROIDS
- VS
- •NOVEL NON STEROIDALS

# EARLY TREATMENT

•DOES IT PREVENT HYPERPIGMENTATION?

# Improvement in psoriasis plaques and PIH with biologic therapy





# CULTURAL PREFERENCES

- HAIR PRODUCTS
- •VEHICLES
- •SCALP TREATMENTS

# Challenges exist with managing different body areas involved in plaque psoriasis<sup>1,2</sup>

#### Difficult-to-treat areas<sup>3</sup>



### Sensitive areas<sup>1</sup>



### Intertriginous areas<sup>1</sup>

Actual clinical trial patients at baseline





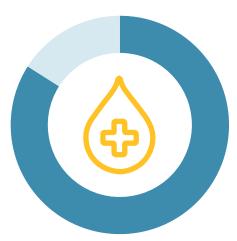
Up to 64% of patients may develop plaque psoriasis in intertriginous areas at some point throughout the course of their disease\*4

<sup>\*</sup>This survey was conducted online by The Harris Poll on behalf of Arcutis Biotherapeutics among US adults 18+ who have been diagnosed with psoriasis by a healthcare provider. The survey was conducted in 2021, among 507 plaque psoriasis patients. Figures were weighted when necessary to bring the data into line with actual proportions in the population using a multi-step weighting process.

<sup>1.</sup> Dopytalska K, et al. Reumatologia. 2018;56(6):392-398. 2. Merola JF, et al. Dermatol Ther. 2018;31(3):e12589. 3. Aldredge LM, et al. J Dermatol Nurses Assoc. 2018;10(4):189-197. 4. Data on File. Arcutis Biotherapeutics, Inc.

### Recent patient survey highlights gaps in topical treatment options

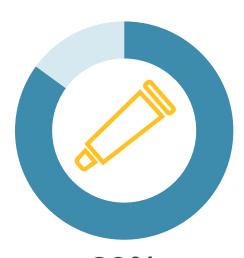
A 2021 survey of plaque psoriasis patients in the US:



**81%**Want more topical alternatives to steroids (n=507)



43%
Use treatments on body
areas not discussed
with their HCP
(n=168)\*



89%
Are interested in trying a new topical treatment for their plaque psoriasis (n=507)

This survey was conducted online by The Harris Poll on behalf of Arcutis Biothetapeutics among US adults 10: who have been diagnosed with psoriasis by a healthcare provider. The survey was conducted in 2021, among 507 plaque psoriasis patients. Figures were weighted when necessary to bring the data into line with actual proportions in the population using a multi-step weighting process.

\*For subset of patients who use different treatments for different parts of their body. HCP. healthcare provider.

HCP, healthcare provider.

c/o Andrew Alexis MD, MPH - Not to be used without written permission



c/o Andrew Alexis MD, MPH - Not to be used without written permission

# Scalp Psoriasis Nuances to Treatment

- Select treatment regimen that is compatible with patient's hair care practices
  - Less frequent hair washing frequency in women of African descent (typically once per week to once every other week)
  - Daily hair washing, especially with most prescription shampoos, is often associated with increased hair dryness and breakage; it is also very time consuming for most women of African descent due to common styling practices

c/o Andrew Alexis MD, MPH - Not to be used without written permission

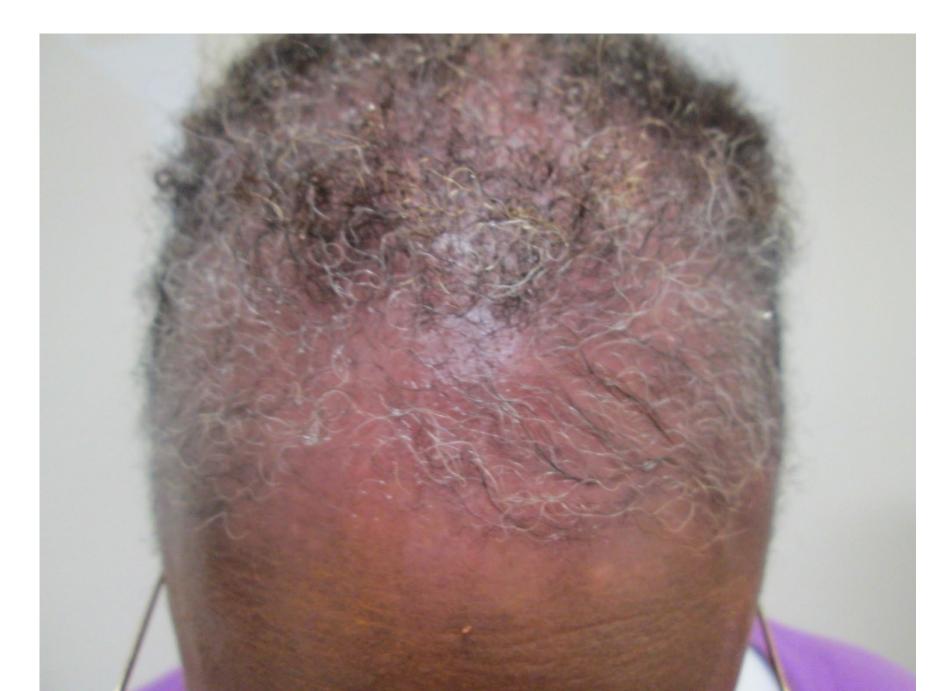
#### Clinical trial

A calcipotriene/betamethasone dipropionate two-compound scalp formulation in the treatment of scalp psoriasis in Hispanic/Latino and Black/African American patients: results of the randomized, 8-week, double-blind phase of a clinical trial

Stephen Tyring<sup>1</sup>, MD, Natalia Mendoza<sup>1</sup>, MD, Melanie Appell<sup>2</sup>, MD, Adrian Bibby<sup>3</sup>, BSc, Richard Foster<sup>4</sup>, MSc, Tiffani Hamilton<sup>5</sup>, MD, and Mark Lee<sup>6</sup>, MD

- 177 Hispanic/Latino and Black/African American subjects
- Safe, effective, and well tolerated
- Vehicle (base of hydrogenated castor oil, PPG-11 stearyl ether, all-rac-alphatocopherol, butylhydroxytoluene, and mineral oil) compatible with Afro-textured hair and common hair styles in AA women

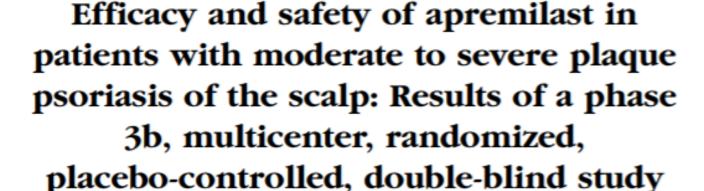
c/o Andrew Alexis MD, MPH - Not to be used without written permission



### Non-Topical Treatment Options for Scalp Psoriasis



### Non-Topical Treatment Options for Scalp Psoriasis





Abby S. Van Voorhees, MD, <sup>a</sup> Linda Stein Gold, MD, <sup>b</sup> Mark Lebwohl, MD, <sup>c</sup> Bruce Strober, MD, <sup>d,c</sup> Charles Lynde, MD, <sup>f</sup> Stephen Tyring, MD, <sup>g</sup> Ashley Cauthen, MD, <sup>h</sup> Howard Sofen, MD, <sup>i</sup> Zuoshun Zhang, PhD, <sup>j</sup> Maria Paris, MD, <sup>j</sup> and Yao Wang, MD, <sup>j</sup> Norfolk, Virginia; West Bloomfield, Missouri; New York, New York, New Haven and Cromwell, Connecticut; Markbam, Ontario, Canada; Houston, Texas; Ocala, Florida; Los Angeles, California; and Summit, New Jersey

# PSORIATIC ARTHRITIS

# SKIN MANIFESTATIONS USUALLY PRECEDE JOINT SYMPTOMS BY ABOUT 10 YEARS IN PATIENTS WITH PSORIATIC ARTHRITIS<sup>1-8</sup>

**UP TO 30%**OF PSORIASIS PATIENTS may actually have psoriatic arthritis<sup>5-8</sup>



- Skin symptoms occur first in ~70% of patients<sup>1,3,4</sup>
- Joint and skin symptoms occur together in ~15% of patients with psoriatic arthritis<sup>1,3,4</sup>
- Joint symptoms occur first in ~15% of patients with psoriatic arthritis<sup>1-4</sup>
- Diagnosis received ≥2 years from symptom onset in 29% of patients with psoriatic arthritis<sup>8</sup>

# Approximately 50% of Psoriatic Arthritis (PsA) Patients Have Joint Damage\*,1,2

Up to 30% of psoriasis patients may have PsA<sup>3-5</sup>







Note: Images are not of the same patient. The image on the right is not a real x-ray and is for illustrative purpose only.

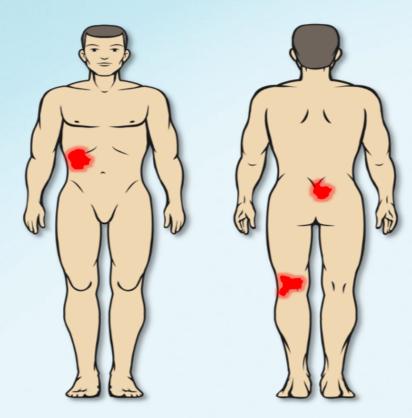
<sup>1.</sup> Kane D, et al. Rheumatology. 2003;42:1460-1468. 2. Torre Alonso JC, et al. Br J Rheumatol.1991;30:245-250. 3. NPF. About Psoriatic Arthritis. https://www.psoriasis.org/about-psoriatic-arthritis. Accessed March 25, 2016. 4. Prey S, et al. J Eur Acad Dermatol Venereol. 2010;24(Suppl 2):31-35. 5. Zachariae H, et al. Acta Derm Venereol. 2002;82:108-1136. 6. Mease P, et al. J Am Acad Dermatol. 2005;52:1-19. 7. Mease PJ. Ann Rheum Dis. 2001;60(Suppl 3):iii37-iii40. 8. Tankosic T, et al. In: Welter R, et al. Psoriasis and Associated Arthropathy. Waltham, Mass: Decision Resources, Inc. 2001:53-54. 9. Gladman DD. In: Isenberg DA, et al. Oxford Textbook of Rheumatology. 3rd ed. New York, NY: Oxford University Press. 2004:766-778.

# PSA IS CORRELATED TO LARGE BSA



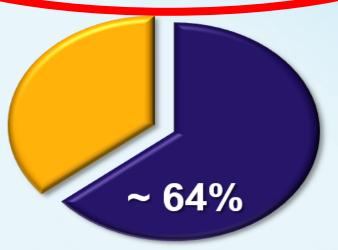
MYTH r TRUTE





1% BSA is approximately the size of the palm, including the fingers<sup>1</sup>

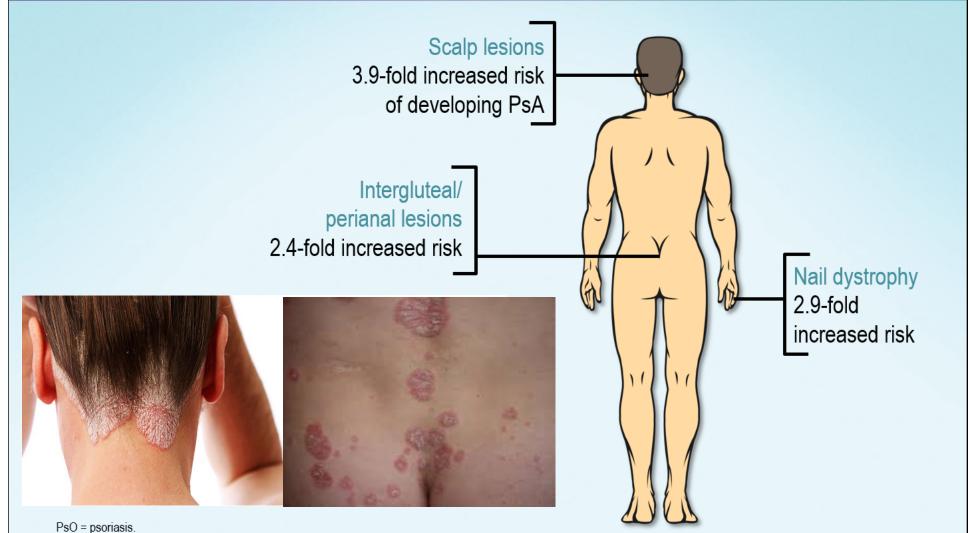
What percent BSA involvement do you typically see in patients with PsA in your office?



of patients with PsA have < 3% BSA affected\*,2

\*Data from the CORRONA registry (N = 1567).

### Clinical Characteristics Can Predict Development of PsA in **Patients With Psoriasis**



In a study of 1593 patients with PsO, the 57 who developed new-onset PsA were more likely to have ≥ 1 of the features of PsO shown than patients who did not develop PsA.

Wilson FC, et al. Arthritis Rheum. 2009;61:233-239.

### **GRAPPA RECOMENDATIONS**



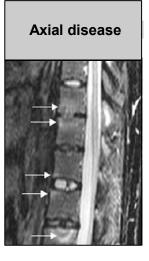
THE GROUP FOR RESEARCH AND ASSESSMENT OF PSORIASIS AND PSORIATIC ARTHRITIS

#### **Psoriatic Arthritis is a Multi-Domain Disease**

 PsA can affect one or more of the six clinical domains recognized by the Group for Research and Assessment of Psoriasis and Psoriatic Arthritis (GRAPPA)

#### **GRAPPA DOMAINS**













### Predictors of progression from psoriasis to PsA

Site of psoriasis	HR
Scalp	3.89
Nail	2.93
Intergluteal/perianal	2.35
Axilla/groin	1.40

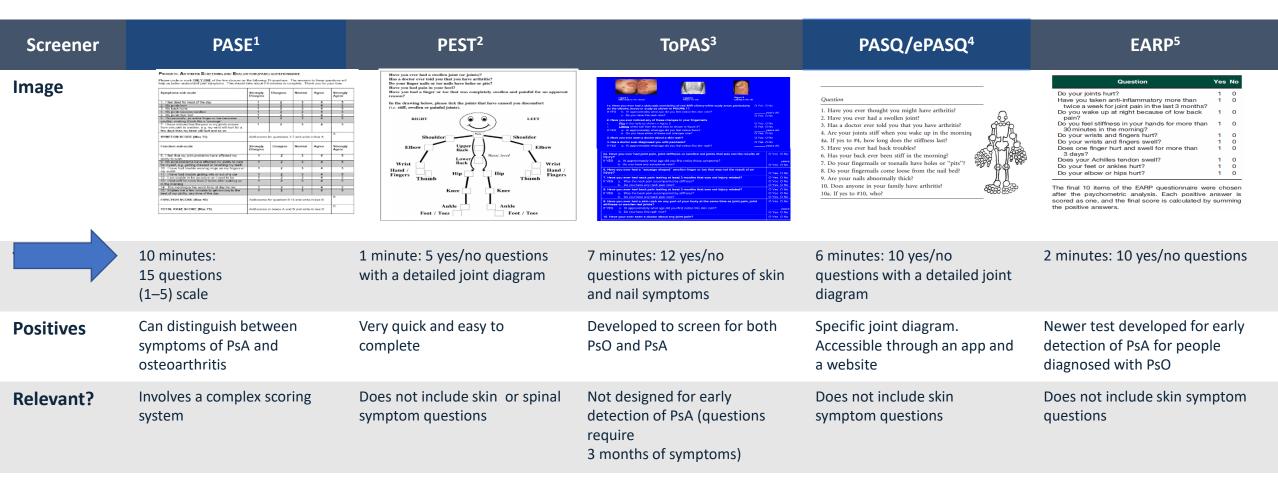
<1

1–2 3–10 >10

HR, hazard ratio; PsA, psoriatic arthritis

Psoriasis incidence cohort included 1593 subjects without PsA

### PsA screening tools



EARP, Early Arthritis Psoriatic patients; PASE, Psoriatic Arthritis Screening Evaluation; PASQ/ePASQ, electronic/Psoriatic Arthritis Screening Questionnaire; PEST, Psoriatic Epidemiological Screening Tool; PsO, psoriasis; ToPAS, Toronto Psoriatic Arthritis Screening Questionnaire

- 1. Husni E, et al. J Am Acad Dermatol 2007;57:481–487;
- 2. Ibrahim GH, et al. Clin Exp Rheumatol 2009;27:469–474;
  - 3. Gladman DD, et al. Ann Rheum Dis 2009;68:491-501;
  - 4. Khraishi M, et al. J Cutan Med Surg 2011;15:143-149;
- 5. Tinazzi I, et al. Rheumatology (Oxford) 2012;51:2058–2063