# **PSORIASIS UNCOVERED**

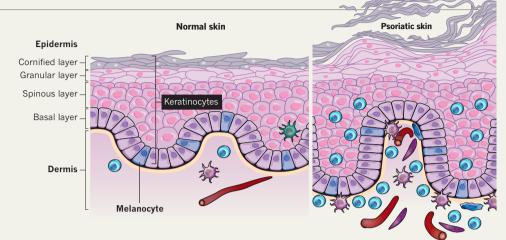
Science is finally getting to grips with this enigmatic autoimmune disease. By James Mitchell Crow.

#### SKIN DEEP

Psoriasis is a non-contagious chronic skin disease affecting the keratinocytes, the cells that predominantly form the epidermis. During outbreaks of the disease, the keratinocytes proliferate at ten times the rate of non-diseased skin cells and fail to mature properly, resulting in raised, inflamed, scaly red skin lesions known as plaques, which can be itchy and painful.

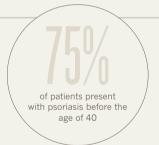


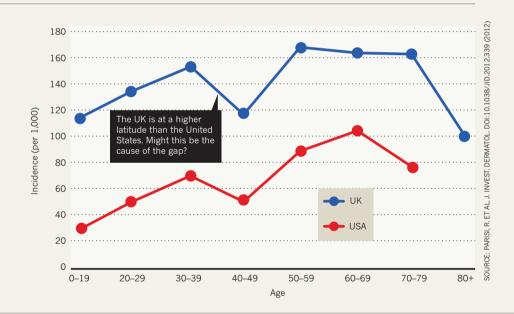




#### **AGE OF ONSET**

Although plaque psoriasis is considered a single disease, its severity, progression and response to treatment can vary markedly between patients - suggesting the existence of multiple underlying subtypes. For example, incidence data by age show two spikes, suggesting early- and late-onset forms of the disease.





# **TREATMENT**

A broad spectrum of therapies are available to treat psoriasis, depending on the severity of the disease. Newer biologic drugs are assessed by a 'PASI 75' score, representing the percentage of patients achieving at least a 75% reduction in their Psoriasis Area and Severity Index (PASI). Despite considerable progress (see 'Silencing psoriasis', page S58), a cure remains elusive.





Systemic small molecule



## >100 YEARS AGO **COAL TAR**

Tars have been used for nearly 2,000 years to treat skin diseases. They are often messy and smelly, but effective.

# GOECKERMAN THERAPY

A combination of coal tar and ultraviolet (UV) irradiation. A course of treatment takes several weeks, and is now less commonly used.

# CORTICOSTEROIDS

Very effective as a short-term treatment. Unsuitable for long-term use because of the growing risk of

1950s **METHOTREXATE** Safe and highly

effective for chronic plaque psoriasis. In common use for many years before official FDA approval in 1972

>100 YEARS AGO

1920

1930

1940

1950

1960

# **PSORIASIS IS COMMON**

Global epidemiological data are sparse. There is a lack of uniformity of data collection, so worldwide incidence and prevalence are poorly documented.

Some trends do emerge — for example, prevalence is higher closer to the poles. A combination of genetic and environmental factors are thought to be behind this pattern.

Temperate (latitude above 40°) Subtropical (23-40°)

Tropical (<23°)

#### Study type

1038/JID.2012.339

DOI:10.1

INVEST

PARISI.

SORIATIC ARTHRITIS POCKET GUIDE, NATIONAL PSORIASIS

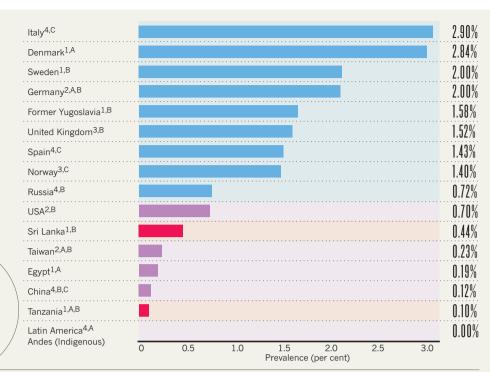
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SOURCE

- 1 Point prevalence
- 2 Period prevalence
- 3 Lifetime prevalence 4 Not specified

### Diagnostic method

- A Dermatologist
- **B** Physician
- C Self-reported



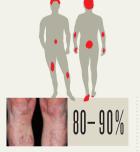
# **PSORIASIS IS HIGHLY HETEROGENEOUS**

average global

prevalence of

psoriasis

Psoriasis comes in several distinct forms. Patients usually display a single form at any one time, although forms can coexist, and one form can be followed by another. Around 80% of psoriasis cases are classified as mild.



#### Chronic plaque psoriasis

Red, scaly plaques in discrete patches. The extent of body surface area covered varies widely from patient to patient.



#### **Guttate psoriasis**

Multiple small, red spots, usually on the trunk and limbs.



#### Inverse/flexural psoriasis

Very red scale-free lesions that form in skin folds.



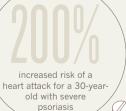
#### Pustular psoriasis

White blisters surrounded by red skin, which may be localized to a particular area, but which can also cover the whole body.



### **Erythrodermic psoriasis**

Severe red inflammation and skin shedding covering most of the body. This is a rare but dangerous form of the disease that can cause patients to lose excessive amounts of heat through the skin.



# 1970s 🚕 UVB LIGHT

Can clear mild cases of the disease.

#### **PSORALEN AND UVA** LIGHT (PUVA)

Possibly slightly more effective, although less convenient, than UVB therapy. A second-line treatment.



Effective for generalized pustular and erythrodermic psoriasis, particularly in combination with UVB or PUVA therapy.

# 2004 **ENBREL (ETANERCEPT)** PASI 75 at 24 weeks: 59%

REMICADE (INFLIXIMAB)

PASI 75 at 26

weeks: 50%

2003 **AMEVIVE** (ALEFACEPT) PASI 75 at 14 weeks: 21%

2008 HUMIRA (ADALIMUMAB)

PASI 75 at 16 weeks: 71%

> STELARA (USTEKINUMAB) PASI 75 at 28 weeks: 71-78%

VITAMIN D3 ANALOGUES Well tolerated and effective

for long-term use with minimal side effects.

1970

1980

1990

2000

2010