

AnnRheumDis2005;64(SupplII):ii18–ii23.doi:10.1136/ard.2004.033217 The molecular genetic basis of psoriasis is complex with Psoriasis a common chronic, recurrent, immunemediated evidence that multiple genes are involved.

Seven major disease of the skin and joints.

It can have a significant psoriasis susceptibility loci have been reported.

Many negative impact on the physical, emotional, and, psycho- investigators have established that a major susceptibility social wellbeing of affected patients.

Psoriasis is found locus for psoriasis is at 6p21, referred to as PSORS1 and is worldwide but the prevalence varies among different ethnic overrepresented in all populations tested.10–15 As noted, an groups. It has a strong genetic component but environmental association between psoriasis and other loci has also been factors such as infections can play an important role in the

reported on chromosomes 1p(PSORS7), 141q(PSORS4), 163q

presentation of disease. There are several clinical cutaneous (PSORS5), 17 4q (PSORS3), 18

17q (PSORS2), 19 and 19p manifestations of psoriasis but most commonly the disease

(PSORS6).20 The strength of associations between such genes presents as chronic,

symmetrical, erythematous, scaling

and susceptibility to psoriasis, apart from PSORS1, is variable papules and plaques.

The epidemiology, clinical features, as replication of these findings has been incomplete.

The and impact on quality of life of psoriasis are reviewed.

difficulty of confirming psoriasis susceptibility loci may

relate, in part, to heterogeneity among different populations.

Whereas the existence of a genetic component in psoriasis is

certain, the exact location of the genes involved remain to

This paper reviews the epidemiology and clinical features be definitely determined.

of psoriasis and its impact of patients' quality of life.

CLINICAL FEATURES **EPIDEMIOLOGY** Psoriasis is a papulosquamous disease with

variable mor- Although psoriasis occurs worldwide, its prevalence varies phology, distribution, severity, and course.

Papulosquamous considerably.

In the USA, approximately 2% of the popula- diseases are characterised by scaling papules (raised lesions tion is affected. High rates of psoriasis have been reported in , 1 cm in diameter) and plaques (raised lesions .1 cm in people of the Faroe islands, where one study found 2.8% of diameter).

Other papulosquamous diseases that may be the population to be affected.1

The prevalence of psoriasis is considered in the differential diagnosis include tinea infection, certain ethnic groups such as the Japanese, and may tions, pityriasis rosea, and lichen planus.

The lesions of be absent in aboriginal Australians2 and Indians from South psoriasis are distinct from these other entities and are America.3 classically very well circumscribed, circular, red papules or Psoriasis can present at any age and has been reported at plaques with a grey or silvery-white, dry scale.

In addition, birth and in older people of advanced age.

Accurate the lesions are typically distributed symmetrically on the

determination of the age of onset of psoriasis is problematic,

scalp, elbows, knees, lumbosacral area, and in the body folds as studies which do so typically rely on a patient's recall of (fig 1).

Psoriasis may also develop at the site of trauma or the onset of lesions or determine the onset from the injury, known as Koebner's phenomenon.

If psoriasis is physician's diagnosis as recorded on the initial visit.

Data progressive or uncontrolled, it can result in a generalised

based on patient recall can be inaccurate; determining onset exfoliative erythroderma.

Nail involvement may be present, based on first visit to a physician could underestimate