Kurdistan Regional Government
Ministry Higher Education & Scientific Research
Salahaddin University-Erbil
College of Education
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## **Psoriasis Disease**

A Project Submitted to The Council of The College of Education at Salahaddin University-Erbil in Partial Fulfillment of the Requirements for the Degree of B.Sc. at Biology Department.

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April — 2023

## **ACKNOWLEDGAMENTS**

First of all, I wish to express my thanks to the most gracious "ALLAH", the facilitator in every step of my life and work.

I would like thanks to my supervisor Dr. Sarhang for suggesting this topic and giving me useful instruction throughout studying period.

I express my deepest apprection for this academic staff of biology department, collage of Education Salahaddin University-Erbil

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#### **Abstract**

Psoriasis is a skin condition that causes flaky patches of skin which form scales. On brown, black and white skin the patches can look pink or red, and the scales white or silvery. On brown and black skin the patches can also look purple or dark brown, and the scales may look grey, Numerous studies have reported the coexistence of psoriasis and other serious systemic diseases, most often mentioned are cardiovascular diseases, metabolic syndrome, including hypertension, dyslipidemia and diabetes mellitus, and Crohn's disease, psoriasis can be associated with an inflammatory arthritis known as psoriatic arthritis, which involves the joints of the spine and other joints. This occurs without presence of specific antibodies in the blood, Based on the type of skin lesions, location, the age of onset and course of disease, several clinical classifications of psoriasis,

Key word: Psoriasis Disease, Incidence, Management

#### Introduction

Psoriasis is a chronic, noncommunicable, painful, disfiguring and disabling disease for which there is no cure and with great negative impact on patients' quality of life (QoL). It can occur at any age, and is most common in the age group 50-69 which there is no cure and with great negative impact on patients' quality of life (QoL). It can occur at any age, and is most common in the age group 50–69 (1). The reported prevalence Psoriasis is a chronic, noncommunicable, painful, disfiguring and disabling disease for of psoriasis in countries ranges between 0.09% (2) and 11.4% (3), making psoriasis a serious global problem. The etiology of psoriasis remains unclear, although there is evidence for genetic predisposition (4). The role of the immune system in psoriasis causation is also a major topic of research. Although there is a suggestion that psoriasis could be an autoimmune disease, no autoantigen that could be responsible has been defined yet. Psoriasis can also be provoked by external and internal triggers, including mild trauma, sunburn, infections, systemic drugs and stress (5). Psoriasis involves the skin and nails, and is associated with a number of comorbidities. Skin lesions are localized or generalized, mostly symmetrical, sharply demarcated, red papules and plaques, and usually covered with white or silver scales. Lesions cause itching, stinging and pain. Between 1.3% (6) and 34.7% (7) of individuals with psoriasis develop chronic, inflammatory arthritis (psoriatic arthritis) that leads to joint deformations and disability. Between 4.2% and 69% of all patients suffering from psoriasis develop nail changes (8-10). Individuals with psoriasis are reported to be at increased risk of developing other serious clinical conditions such as cardiovascular and other noncommunicable diseases (NCDs) (5,11,12). Psoriasis causes great physical, emotional and social burden (13-15). QoL, in general, is often significantly impaired (16–23). Disfiguration, disability and marked loss of

productivity are common challenges for people with psoriasis. There is also a significant cost to mental well-being, such as higher rates of depression, leading to negative impact for individuals and society (24,25). Social exclusion, discrimination and stigma are psychologically devastating for individuals suffering from psoriasis and their families. It is not psoriasis causing the exclusion – it is largely society's reaction to it and this can change. Treatment of psoriasis is still based on controlling the symptoms. Topical and systemic therapies as well as phototherapy are available. In practice, a combination of these methods is often used. The need for treatment is usually lifelong and is aimed at remission. So far, there is no therapy that would give hope for a complete cure of psoriasis. Additionally, care for patients with psoriasis requires not only treating skin lesions and joint involvement, but it is also very important to identify and manage common comorbidity that already exists or may develop, including cardiovascular and metabolic diseases as well as psychological conditions.

In 2015, the notion is that psoriasis is a complex disease leading to numerous consequences for patients' lives (Figure 1).

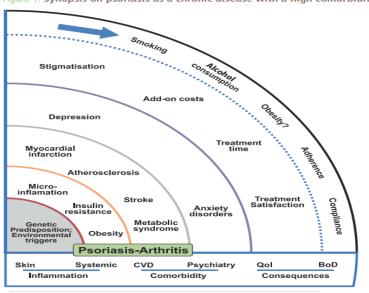


Figure 1. Synopsis on psoriasis as a chronic disease with a high comorbidity

BoD, burden of disease; CVD, cardiovascular disease; QoL, quality of life

Source: Mrowietz et al. 2014 (26).

# **Incidence and prevalence**

It affects men and women of all ages, regardless of ethnic origin, in all countries (1). Published data on the prevalence of psoriasis in countries vary between 0.09% (2) and 11.4% (3). In most developed countries, prevalence is between 1.5 and 5% (27). There is also evidence to suggest that the prevalence of psoriasis may be increasing (3). Many studies have demonstrated that psoriasis can impact substantially on QoL, even when a relatively limited body surface area (BSA) is affected (16,17,22,28–32). There are very few studies on the incidence of psoriasis. Registration of psoriasis cases is not compulsory, meaning reliable data are difficult to find. A review of published literature revealed only a handful of credible studies on the incidence of psoriasis. One study showed that the overall sex- and age-adjusted incidence rate of psoriasis in Minnesota in the United States, between 1980 and 1983, was estimated at 0.60 per 1000 person-years (33). A study of 511 532 individuals in Italy between 2001 and 2005 reported an incidence of psoriasis (adults receiving a first-ever diagnosis of psoriasis) of 2.30-3.21 cases per 1000 person-years (12). In 2012, a 2-week psoriasis screening study via medical consultation was performed in three countries simultaneously - Algeria, Tunisia and Morocco, where incidence of psoriasis was estimated at 10.36, 13.26 and 15.04 per 1000, respectively (34). Relatively more studies have focused on the prevalence of psoriasis. A review of published literature found 68 full articles and reports estimating prevalence rates for 20 countries (Table 1). It should be noted, however, that the data they contain are extremely difficult to compare with each other, due to the different methodologies of the studies and their limitations. The main problems are differences in the definition of prevalence (point prevalence, cumulative prevalence, period prevalence), case definition of psoriasis (selfreported, physician diagnosed), the population ages studied (children only,

adults only, any age group) and the sampling techniques (questionnaires, clinical examination, combination of clinical examination and questionnaire, registry data). This difference in methodology clearly impacted on the prevalence rates. Depending on the region, the prevalence studies varied from 0.09% in the United Republic of Tanzania (2) to 011.4% in Norway (3). A very weak correlation between geographic latitude and psoriasis prevalence was found (35). Psoriasis appears to occur most commonly in populations of northern Europe (3,36) and least in populations of eastern Asia (37–45) Some studies investigated the ethnic differences in the prevalence of psoriasis. According to a 2001 study in the United States, people with Caucasian or Black ancestry and others had a prevalence of 2.5%, 1.3% and 1.0%, respectively (14). In another United States study from 2009–2010, these differences were higher, with the prevalence for Caucasians, Blacks, Hispanics and others at 3.6%, 1.9%, 1.6% and 1.4%, respectively (46). Psoriasis can occur at any age. While some studies indicated the average age of onset for psoriasis was 33 years of age, and 75% of cases occurred before 46 years of age (50), others suggested that the onset of psoriasis was bimodal with two peaks of the disease – the first between 16 and 22 and the second between 57 and 60 years of age (51). Psoriasis also occurs in children. However, there are few studies on the incidence or prevalence of psoriasis in children, and those that do exist reveal variations between almost absence of juvenile psoriasis in Taiwan, China (38,39) and 1.37% lifetime prevalence in 0-17-year-old children in Germany (49). The largest study on prevalence among children was carried out in Germany in 2007 (48). Data collected from a health insurance company database of about 1.3 million individuals showed the prevalence of psoriasis in children younger than 18 years of age was 0.40%, and increased roughly linearly over the life course. In 2008–2009, a study of 2194 children in Egypt (47) found that the prevalence of psoriasis among people 18 years of age and younger was 0.05%.

It can manifest in many different forms. In addition to the involvement of skin and nails, inflammatory arthritis (psoriatic arthritis) may develop. Patients suffering from psoriasis are at higher risk of developing cardiovascular and other NCDs (52). Moreover, psoriasis affects mental health and people suffering from the disease experience significant social stigma. In assessing the severity of psoriasis, more than 40 different tools are being used (53). Commonly used measures for scoring the severity of psoriasis include the Psoriasis Area and Severity Index (PASI), and the Physician Global Assessment. Clinicians assess the severity of the disease, taking into account the degree of scaling, redness, thickness of the skin lesions or the size of the BSA occupied by psoriasis. QoL measures are also important.

# Risk factor of psoriasis

- **Tobacco**. The risk of onset of psoriasis in smokers is twice that of non-smokers. In addition, it seems that the more quantity and the longer you smoke, the higher the risk.(54).
- **Alcohol.** As with smoking, alcohol use also seems to play a role in the onset of psoriasis. Furthermore, alcohol increases severity and may decrease treatment efficacy.(55).
- **Obesity.** Or weight gain has been shown to be an independent risk factor for psoriasis. There are two suspected reasons for this. Firstly, obesity promotes inflammation and, therefore, the onset of psoriasis. Secondly, psoriasis affects mood and can hinder physical activity, all of which favors the development of obesity.(56)

- **Infections**. Infections can trigger and exacerbate psoriasis. Pharyngitis caused by a bacterium called streptococcus is a typical trigger of guttate psoriasis. Also, the human immunodeficiency virus (HIV) can trigger psoriasis.(57).
- **Drugs**. Some drugs can trigger psoriasis, for instance,  $\beta$  blockers (used to treat heart disease) and lithium (a drug for bipolar disorder).(58).
- **Stress.** Stress is associated with both the onset of psoriasis and flare-ups in people previously diagnosed with psoriasis.(59).
- Weather. Skin psoriasis and psoriatic arthritis usually worsen in the winter and improve throughout summer. The National Psoriasis Foundation explains that this is due to air dryness, poor sunlight and cold weather. (60)
- Other skin lesions. Any alteration of the skin—such as scratches, piercings or sunburns—can trigger psoriasis lesions to appear. This is called Koebner's phenomenon.(60-61).

#### Skin and nails

Based on the type of skin lesions, location, the age of onset and course of disease, several clinical classifications of psoriasis are used (Table 2). The most frequently reported symptoms connected to psoriasis are (62): n scaling of the skin in 92% n itching in 72% n erythema in 69% n fatigue in 27% n swelling in 23% n burning in 20% n bleeding in 20% of individuals. According to another study, flaking or scaling in the non-scalp area occurred in 89% and flaking or scaling of scalp areas in 62% of patients. Itching or scratching were observed in 87%, rash in 74%, skin pain in 62%, bleeding in 58%, redness in 57%, flare-ups in 49%, joint pain in 42%, skin cracking in 39%, dry skin in 34%, physical discomfort in 32%, burning in 28% and nail problems in 22% (7).

Table 2. Common types of psoriasis and their manifestations

Ps orias is vulgaris (plaque psoriass)	The most common type of gsoria is, a ffects between 58% (39) and 97% (45) of all patients. Inflammatory red, sharply demacated, raised, dry, differently sized plaques, usually covered by silvery or white scales. Involves the scale and the are abehind the earth side of the extensors unfaces of the fore arms and shins (expecially elbows and knees), trunk, face, pains, soles and nails.	Pronasis wigaris in adult patients. Clutish Mrowietz, Psonasis Center Ref.
Intert if gin ous psoriasis (psoriasis in folds and genital areas)	Affects between 12% (105) and 26% (21) of all cases of products.  Deep-red or white, flat, sharply demancated, wet part dues or plaques, scales are usually absent.  Affects almost exclusively fleoural body sites—ad fae, antecubitations, informammary or eases, umbilizing, sprins, genitari area, glubari cleft, popiliteal forsase and other body folds.	Inter triginous psoria is in adult partients. ©Utrich Mrowletz, Psoriasis Center Kiel.
<b>Gutta te p sori asis</b> (drop let psoriasis)	Affects between 0.6% (45) and 20% (8) of individuals diagnosed with provisits and usually occurs in childhood and adolescence.  Reddish, drop-like pagules and plaques, mainly involving the trunk, arms and legs.  Onset is associated with streptococcol infection of the upper respirator yit act and prior skin symptoms (106).	Guttate psoria sis in an 8-year-old girl. GWidhael 8 Sidrin, Universit absmedizin Göttlingen.
Pustular ps orias is	Affects between 1.1% (45) and 1.2% (105) of all cases of psorfiasis. Coale string pussibles, filled with non-infectious pussion involves eithers mail areas such as pairns of the hands, fingertips, nails and soles of the feet, or the entire body surface can occur as a single episode after a trigger.	Generalized pustular psoria sis in an adult female patient. CMichael P. Schön, Universität smedizin Göttingen.
Erythrode mi c psori asis	Affects between 0.4% (45) and 7% (105) of all cases of psoriasis. Fery redness and oxfoliation of most of the body surface. The most serious type of psoriasis, potentially lifethreatening, because it can lead to hypothermia, hypoalbuminemia and high output cardiac failure.	Eysthrodermic psoriasis in adult male patient: © Matthias Augustin, University Medical Center Hamburg.

Based on a review of the literature, the prevalence of nail psoriasis ranges between 4.2% (8) and 69% (9) of all patients suffering from psoriasis. Nail psoriasis may occur with the involvement of the skin or it may occur alone, being the only symptom of psoriasis. Nail psoriasis is not only a problem of an aesthetic nature, but can also restrict manual dexterity. The nail disease may be acute or chronic, with varied severity. There may be involvement of only a single nail or of all nails associated with severe nail destruction or loss (Figure 5).

Figure 5. Nail psoriasis





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Nail psoriasis may be a predisposing factor for fungal or bacterial infections, which occur in 4.6% to 30% of cases of nail psoriasis (63). Patients suffering from nail psoriasis have significantly worse psoriasis severity scores, compared to patients without nail involvement (64). In addition, they report poorer QoL and a greater number of days unfit for work (64). These patients were also more likely to be admitted to the hospital and more often suffered from psoriatic arthritis.

#### **Psoriatic arthritis**

In addition to the skin, psoriasis can be associated with an inflammatory arthritis known as psoriatic arthritis, which involves the joints of the spine and other joints. This occurs without presence of specific antibodies in the blood (seronegative spondyloarthropathy). The rheumatoid factor (antibody occurring in rheumatoid arthritis) is also negative. A review of the literature showed that psoriatic arthritis affects between 1.3% (6) and 34.7% (7) of patients diagnosed with psoriasis. There are no data on sex predilection. In a United States population, it was observed that psoriatic arthritis occurred more frequently in Caucasian patients than in other ethnic groups (65). Two large consecutive German studies from dermatological practices that assessed the prevalence of arthritis from examinations by rheumatologists in 2005 and 2007 was 20.6% (10) and 19.6% (66), respectively.

The clinical symptoms are variable, however, peripheral arthritis, spondylitis, enthesitis (inflammation of the sites where tendons insert into the bone), arthritis in the fingers and dactylitis (profuse swelling of the fingers or toes) are considered to be most common (Figure 6). Psoriatic arthritis can lead to chronic pain and change in physical appearance. Patients suffering from psoriatic arthritis have reduced physical fitness, compared to psoriasis patients without it (22). Typically, psoriatic arthritis occurs in conjunction with longstanding skin lesions, although rarely it occurs alone, in the absence of psoriasis.

Figure 6. Psoriasis arthritis





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#### **Associated diseases**

Numerous studies have reported the coexistence of psoriasis and other serious systemic diseases, most often mentioned are cardiovascular diseases, metabolic syndrome, including hypertension, dyslipidemia and diabetes mellitus, and Crohn's disease. Even children show increased rates of comorbidity compared to unaffected infants, or those with atopic eczema (11,49). Most publications discuss the association between cardiovascular disease and psoriasis. Patients diagnosed with psoriasis have an increased burden of subclinical atherosclerosis and vascular inflammation (67). They also have significantly higher levels of serum lipids, including triglycerides and total cholesterol, compared to healthy individuals (68).

Moreover, psoriasis is associated with atrial fibrillation and stroke, which may be aggravated in young patients (69). However, it should be noted that at present it is not known whether psoriasis is an independent risk factor for the development of cardiovascular disease. Obesity or weight gain has been shown to be an independent risk factor for psoriasis. Tobacco smoking is another risk factor (70). Frequency of metabolic syndrome, depression and erectile dysfunction has also been found to be significantly higher in patients diagnosed with psoriasis (71). In some diseases and subgroups of patients, psoriasis has been shown to be an independent risk factor for non-alcoholic fatty liver disease (72). In spite of a high number of studies on the association of psoriasis with comorbidity, the causality and independence on some associated diseases remain unclear and need further research (73,74).

# Psychological and mental health

Psoriasis is not only a disease that causes painful, debilitating, highly visible physical symptoms. It is also associated with a multitude of psychological impairments. For many reasons, psoriasis can be psychologically devastating. Patients' lives become especially difficult when psoriasis is present in highly visible areas of the skin such as the face and hands. Related psychological problems can affect every day social activities and work. It causes embarrassment, lack of self-esteem, anxiety and increased prevalence of depression (24,25). Patients with psoriasis report experiencing anger or helplessness, and they disclose a higher rate of suicidal ideations than other patients. In a study of 127 patients with psoriasis, 9.7% reported a wish to be dead and 5.5% reported active suicidal ideation at the time of study (75).

# Influences from the workplace

In the workplace, psoriasis may be triggered or aggravated by mechanical or other physical impact on unprotected body locations (76). In patients who claimed occupational hand dermatitis, psoriasis was found to be the cause of the disease in 3.8% (77)–6.5% (78) of cases. Special gloves and other personal protective equipment may reduce lesions and enable the person to continue working, which otherwise may be jeopardized (77,79).

# **Social participation**

Psoriasis can affect relationships at home, school or work as well as sexual relationships and thus reduce QoL and cause psychological strain (15,17–19,32,80–84). Patients are frequently stigmatized and excluded from normal social environments, including schools, workplaces and swimming pools. As a result, they often avoid social activities and commonly report experiencing loneliness, isolation, feelings of being unattractive and frustration. A study conducted in the United States evaluated which spheres of patients' lives suffered most (7). Seven impact areas were considered: emotional (mood, feelings); social (friends, activities); the family (activities, responsibilities); professional (work, career); physical functioning; sexual intimacy; and educational life. The results showed that 98% of patients reported that psoriasis impacted on their emotional life, 94% on their social life, 70% on family life, 68% on their professional career, 38% on physical functioning, 17% on sexual intimacy, and 21% on their educational life. These values were all higher for patients with psoriatic arthritis.

# Principles of psoriasis management

Psoriasis is by nature a chronic, incurable disease with an unpredictable course of symptoms and triggers. The consequence is often life-long treatment, therefore, all treatments must meet high quality criteria that are not only efficacious, but also safe over long periods. As the cause of psoriasis is still unknown, treatment is only available to control symptoms. Treatments include a range of topical and systemic therapies as well as phototherapy. It also involves treatment for reducing pain and disability from arthritis and other manifestations. Care for patients with psoriasis requires more than management of the skin lesions and joint involvement. The complexity of psoriasis means that prescribing drugs in isolation is insufficient to control the disease and a holistic, whole person approach to care is needed. Management of psoriasis also includes screening for associated diseases such as hypertension, dyslipidemia, diabetes mellitus and cardiovascular disorders as well as their complications such as myocardial infarction and stroke (Figure 7). Psoriasis patients are more likely to suffer from depression and anxiety disorders and have an increased rate of suicidal ideation. Screening at regular intervals for these associated diseas[es and for co-medication to prevent drug-drug interactions or drug-triggered psoriasis as well as recognition of trigger factors and their treatment are an essential part of psoriasis management.

# Treating the skin manifestations

There are three major forms of therapy – topical therapy; phototherapy; and systemic therapy (Table 3). Treatment is based on psoriasis severity at the time of presentation. Mild psoriasis usually is treated with topical therapy, progressing to phototherapy in the case of insufficient response. Moderate to severe psoriasis requires systemic therapy. Commonly used first-line drugs include methotrexate, ciclosporin, acitretin and etretinate. In some countries, other systemic therapies such as biologic agents and fumaric acid esters are available (8). All treatments for psoriasis, apart from retinoids, are primarily anti-inflammatory, and subsequently lead to slowed epidermal keratinocyte turnover and a flattening of plaques. In many countries, other treatments may play a prominent role, including traditional Chinese medicine, self-treatment with over-the-counter products (non-prescription drugs) and climatotherapy.

Table 3. Treatment options for psoriasis

Topical therapies (ointments, creams,	Vitamin D, analogues
lotions, gels, or foams applied to the skin)	Corticosteroids
	e.g. betamethasone and hydrocortisone
	Anthralin/dithranol
	Topical retinoids
Phototherapy (UV-light therapy)	
Systemic therapies (tablets or	Methotrexate
injections/infusion)	Ciclosporin
	Acitretin
	Biologic agents
	Oral small molecules

#### **Reference:**

- 1. Institute for Health Metrics and Evaluation (IHME). Global Burden of Disease Study 2010: Results by Cause 1990–2010. Seattle: IHME; 2012.
- 2. Gibbs S. Skin disease and socioeconomic conditions in rural Africa: Tanzania. Int J Dermatol. 1996;35(9):633–9.
- 3. Danielsen K, Olsen AO, Wilsgaard T, Furberg AS. Is the prevalence of psoriasis increasing? A 30–year follow-up of a population–based cohort. Br J Dermatol. 2013;168:1303–10.
- 4. Harden JL, Krueger JG, Bowcock AM. The immunogenetics of psoriasis: a comprehensive review. J Autoimmun. 2015;64:66–73.
- 5. Boehncke W-H, Schön MP. Psoriasis. Lancet. 2015;386(9997):983–94.
- 6. Bedi TR. Clinical profile of psoriasis in North India. Indian J Dermatol Venereol Leprol. 1995;61(4):202–5.
- 7. Pariser D, Schenkel B, Carter C, Farahi K, Brown TM, Ellis CN, and Psoriasis Patient Interview Study Group. A multicenter, non-interventional study toevaluate patient-reported experiences of living with psoriasis. J Dermatol Treat. 2015;1–8.
- 8. Alshami MA. Clinical profile of psoriasis in Yemen, a 4-year retrospective study of 241 patients. J Eur Acad Dermatol Venereol. 2010;24(Suppl. 4):14.
- 9. Falodun OA. Characteristics of patients with psoriasis seen at the dermatology clinic of a tertiary hospital in Nigeria: a 4-year review 2008–2012. J Eur Acad Dermatol Venereol. 2013;27(Suppl. 4)
- 10.Reich K, Krüger K, Mössner R, Augustin M. Epidemiology and clinical pattern of psoriatic arthritis in Germany: a prospective interdisciplinary epidemiological study of 1511 patients with plaque-type psoriasis. Br J Dermatol. 2009;160(5):1040–7.

- 11. Augustin M, Radtke MA, Glaeske G, Reich K, Christophers E, Schaefer I et al. Epidemiology and Comorbidity in Children with Psoriasis and Atopic Eczema. Dermatology. 2015;231(1):35–40.
- 12. Vena GA, Altomare G, Ayala F, Berardesca E, Calzavara-Pinton P, Chimenti S et al. Incidence of psoriasis and association with comorbidities in Italy: a 5-year observational study from a national primary care database. Eur J Dermatol. 2010;20(5):593–8.
- 13. Fuji R, Mould JF J, Tang B, Brandt H, Pomerantz D, Chapnick J et al. Burden of disease in patients with diagnosed psoriasis in Brazil: results from 2011 national health and wellness survey (NHWS). Value Health. 2012;15(4):A107.
- 14.Stern RS, Nijsten T, Feldman SR, Margolis DJ, Rolstad T. Psoriasis is common, carries a substantial burden even when not extensive, and is associated with widespread treatment dissatisfaction. J Investig Dermatol Symp Proc. 2004;9(2):136–9.
- 15.Kimball AB, Jacobson C, Weiss S, Vreeland MG, Wu Y. The psychosocial burden of psoriasis. Am J Clin Dermatol. 2005;6(6):383–92.
- 16.De Korte J, Sprangers MA, Mombers FM, Bos JD. Quality of life in patients with psoriasis: a systematic literature review. J Investig Dermatol Symp Proc. 2004;9(2):140–7.
- 17. Zachariae H, Zachariae R, Blomqvist K, Davidsson S, Molin L, Mørk C et al. Quality of life and prevalence of arthritis reported by 5,795 members of the Nordic Psoriasis Associations. Data from the Nordic Quality of Life Study. Acta Derm Venereol. 2002;82(2):108–13.
- 18.Krueger G, Koo J, Lebwohl M, Menter A, Stern RS, Rolstad T. The impact of psoriasis on quality of life: results of a 1998 National Psoriasis Foundation patient-membership survey. Arch Dermatol. 2001;137(3):280–4.

- 19. Weiss SC, Kimball AB, Liewehr DJ, Blauvelt A, Turner ML, Emanuel EJ. Quantifying the harmful effect of psoriasis on health-related quality of life. J Am Acad Dermatol. 2002;47(4):512–8.
- 20. Kimball AB, Gieler U, Linder D, Sampogna F, Warren RB, Augustin M. Psoriasis: Is the impairment to a patient's life cumulative? J Eur Acad Dermatol Venereol. 2010;24(9):989–1004.
- 21. Moradi M, Rencz F, Brodszky V, Moradi A, Balogh O, Gulácsi L. Health status and quality of life in patients with psoriasis: an Iranian cross-sectional survey. Arch Iran Med. 2015;18(3):153–9.
- 22. Tang MM, Chang CC, Chan LC, Heng A. Quality of life and cost of illness in patients with psoriasis in Malaysia: a multicenter study. Int J Dermatol. 2013;52(3):314–22.
- 23. Augustin M, Krüger K, Radtke MA, Schwippl I, Reich K. Disease severity, quality of life and health care in plaquetype psoriasis: a multicenter cross-sectional study in Germ
- 24.Russo PAJ, Ilchef R, Cooper AJ. Psychiatric morbidity in psoriasis: a review. Australas J Dermatol. 2004;45(3):155–9; quiz;160–1.
- 25. Sampogna F, Tabolli S, Abeni D, IDI Multipurpose Psoriasis Research on Vital Experiences (IMPROVE) investigators. Living with psoriasis: prevalence of shame, anger, worry, and problems in daily activities and social life. Acta Derm Venereol. 2012;92(3):299–303.
- 26.Mrowietz U, Steinz K, Gerdes S. Psoriasis: To treat or to manage? Exp Dermatol. 2014;23(10):705–9.
- 27. Parisi R, Symmons DPM, Griffiths CEM, Ashcroft DM, and the Identification and Management of Psoriasis and Associated ComorbidiTy (IMPACT) project team. Global epidemiology of psoriasis: a systematic review of incidence and prevalence. J Invest Dermatol. 2013;133(2):377–85.

- 28.Gelfand JM, Feldman SR, Stern RS, Thomas J, Rolstad T, Margolis DJ. Determinants of quality of life in patients with psoriasis: a study from the US population. J Am Acad Dermatol. 2004;51(5):704–8.
- 29. Nijsten T, Meads DM, de Korte J, Sampogna F, Gelfand JM, Ongenae K et al. Cross-cultural inequivalence of dermatology-specific health-related quality of life instruments in psoriasis patients. J Invest Dermatol. 2007;127(10):2315–22.
- 30. Augustin M, Radtke MA. Quality of life in psoriasis patients. Expert Rev Pharmacoecon Outcomes Res. 2014;14(4):559–68.
- 31.Prins M, Krabbe PFM, Swinkels QOJ, de Boo T, van de Kerkhof PCM, van der Valk PGM. The effect of treatment on quality of life in psoriasis patients. Acta Derm Venereol. 2005;85(4):304–10.
- 32. Vardy D, Besser A, Amir M, Gesthalter B, Biton A, Buskila D. Experiences of stigmatization play a role in mediating the impact of disease severity on quality of life in psoriasis patients. Br J Dermatol. 2002;147(4):736–42.
- 33.Bell LM, Sedlack R, Beard CM, Perry HO, Michet CJ, Kurland LT. Incidence of psoriasis in Rochester, Minn, 1980–1983. Arch Dermatol. 1991;127(8):1184–7.
- 34. Ammar-Khodja A, Benkaidali I, Bouadjar B, Serradj A, Titi A, Benchikhi H et al. EPIMAG: International CrossSectional Epidemiological Psoriasis Study in the Maghreb. Dermatology. 2015;231(2):134–44
- 35.Jacobson CC, Kumar S, Kimball AB. Latitude and psoriasis prevalence. J Am Acad Dermatol. 2011;65(4):870–3.
- 36.Bø K, Thoresen M, Dalgard F. Smokers report more psoriasis, but not atopic dermatitis or hand eczema: results from a Norwegian population survey among adults. Dermatol Basel Switz. 2008;216(1):40–5.

- 37.Shao CG, Zhang GW, Wang GC. Distribution of psoriasis in China: a nationwide screening. Proc Chin Acad Med Sci Peking Union Med Coll. 1987;2(2):59–65.
- 38.Chen G-Y, Cheng Y-W, Wang C-Y, Hsu T-J, Hsu MM-L, Yang P-T et al. Prevalence of skin diseases among schoolchildren in Magong, Penghu, Taiwan: a community-based clinical survey. J Formos Med Assoc. 2008;107(1):21–9.
- 39. Yang Y-C, Cheng Y-W, Lai C-S, Chen W. Prevalence of childhood acne, ephelides, warts, atopic dermatitis, psoriasis, alopecia areata and keloid in Kaohsiung County, Taiwan: a community-based clinical survey. J Eur Acad Dermatol Venereol. 2007;21(5):643–9.
- 40.Li M-J, Wang P, Wu W-W, Fu L, Cai M, Chen M-X et al. An epidemiological survey of psoriasis in 18 cities in Hainan province of China. J Dermatol. 2012;39(Suppl. 1):243–4.
- 41. Chang Y-T, Chen T-J, Liu P-C, Chen Y-C, Chen Y-J, Huang Y-L et al. Epidemiological study of psoriasis in the national health insurance database in Taiwan. Acta Derm Venereol. 2009;89(3):262–6.
- 42. Yip SY. The prevalence of psoriasis in the Mongoloid race. J Am Acad Dermatol. 1984;10(6):965–8.
- 43.Ding X, Wang T, Shen Y, Wang X, Zhou C, Tian S et al. Prevalence of psoriasis in China: a population-based study in six cities. Eur J Dermatol. 2012;22(5):663–7.
- 44. Wang R, Cao L, Zhou C, Zhang J. Prevalence of 15 skin diseases in adolescents from Liangshan prefecture in Sichuan Province. Chin J Dermatol. 2012;45(4):270–2.

- 45. Kubota K, Kamijima Y, Sato T, Ooba N, Koide D, Iizuka H et al. Epidemiology of psoriasis and palmoplantar pustulosis: a nationwide study using the Japanese national claims database. BMJ Open. 2015;5(1):e006450.
- 46.Rachakonda TD, Schupp CW, Armstrong AW. Psoriasis prevalence among adults in the United States. J Am Acad Dermatol. 2014;70(3):512–6.
- 47. Yamamah GA, Emam HM, Abdelhamid MF, Elsaie ML, Shehata H, Farid T et al. Epidemiologic study of dermatologic disorders among children in South Sinai, Egypt. Int J Dermatol. 2012;51(10):1180–5.
- 48. Augustin M, Glaeske G, Radtke M, Christophers E, Reich K, Schaefer I. Epidemiology and comorbidity of psoriasis in children. Br J Dermatol. 2010;162:633–6.
- 49. Schmitt J, Apfelbacher C. Epidemiology of pediatric psoriasis: a representative German cross-sectional study. Exp Dermatol. 2010;19(2):219.
- 50.Nevitt GJ, Hutchinson PE. Psoriasis in the community: prevalence, severity and patients' beliefs and attitudes towards the disease. Br J Dermatol. 1996;135(4):533–7.
- 51.Henseler T, Christophers E. Psoriasis of early and late onset: characterization of two types of psoriasis vulgaris. J Am Acad Dermatol. 1985;13(3):450–6.
- 52. Abuabara K, Azfar RS, Shin DB, Neimann AL, Troxel AB, Gelfand JM. Cause-specific mortality in patients with severe psoriasis: a population-based cohort study in the U.K. Br J Dermatol. 2010;163(3):586–92.
- 53.Naldi L, Svensson A, Diepgen T, Elsner P, Grob J-J, Coenraads P-J et al., and the European DermatoEpidemiology Network. Randomized clinical trials for psoriasis 1977–2000: the EDEN survey. J Invest Dermatol. 2003;120(5):738–41.

- 54.Kamiya, K., Kishimoto, M., Sugai, J., Komine, M., Ohtsuki, M. Risk Factors for the Development of Psoriasis. 2019 Sep 5;20(18)
- 55.Naldi, L. Psoriasis and smoking: links and risks. Dovepress [Internet]. 2016 May 27;6:6571.
- 56. National Psoriasis Foundation (NPF). Frequently Asked Questions: Psoriasis in spring, summer, fall and winter [Internet]. 2019 [Cited 2020 Feb 22]
- 57. Ford, A.R., Siegel, M., Bagel, J., Cordoro, K.M., Garg, A., Gottlieb, A., et al. Dietary Recommendations for Adults With Psoriasis or Psoriatic Arthritis From the Medical Board of the National Psoriasis Foundation: A Systematic Review. JAMA Dermatol [Internet]. 2018;154(8):934-950.
- 58. World Health Organization (WHO). Global report on psoriasis [Internet]. 2016.
- 59. Gibson, L.E. Psoriasis flare-ups: How can I recognize my unique triggers? Mayo Clinic [Internet]. 2019 Jan 03.
- 60. American Academy of Dermatology (AAD). Psoriasis triggers: How to find and manage yours (Internet).
- 61. European Academy of Dermatology and Venereology (EADV). Psoriasis can be hereditary [Internet].
- 62. Dubertret L, Mrowietz U, Ranki A, van de Kerkhof PC, Chimenti S Lotti T et al. European patient perspectives on the impact of psoriasis: the EUROPSO patient membership survey. Br J Dermatol. 155(4):729–36.
- 63.Owen CM, Chalmers RJ, O'Sullivan T, Griffiths CE. Antistreptococcal interventions for guttate and chronic plaque psoriasis. Cochrane Database Syst Rev. 2000;(2):CD001976.
- 64. Natarajan V, Nath AK, Thappa DM, Singh R, Verma SK. Coexistence of onychomycosis in psoriatic nails: a descriptive study. Indian J Dermatol Venereol Leprol. 2010;76(6):723.
- 65. Augustin M, Reich K, Blome C, Schäfer I, Laass A, Radtke MA. Nail psoriasis in Germany: epidemiology and burden of disease. Br J Dermatol. 2010;163(3):580–5.
- 66.Kerr GS, Qaiyumi S, Richards J, Vahabzadeh-Monshie H, Kindred C, Whelton S et al. Psoriasis and psoriatic arthritis in African-American patients: the need to measure disease burden. Clin Rheumatol. 2014.

- 67.Radtke MA, Reich K, Blome C, Rustenbach S, Augustin M. Prevalence and clinical features of psoriatic arthritis and joint complaints in 2009 patients with psoriasis: results of a German national survey. J Eur Acad Dermatol Venereol. 2009;23(6):683–91.
- 68. Shaharyar S, Warraich H, McEvoy JW, Oni E, Ali SS, Karim A et al. Subclinical cardiovascular disease in plaque psoriasis: association or causal link? Atherosclerosis. 2014;232(1):72–8.
- 69.Robati RM, Partovi-Kia M, Haghighatkhah HR, Younespour S, Abdollahimajd F. Increased serum leptin and resistin levels and increased carotid intima-media wall thickness in patients with psoriasis: is psoriasis associated with atherosclerosis? J Am Acad Dermatol. 2014;71(4):642–8.
- 70. Ahlehoff O, Gislason GH, Jørgensen CH, Lindhardsen J, Charlot M, Olesen JB et al. Psoriasis and risk of atrial fibrillation and ischaemic stroke: a Danish Nationwide Cohort Study. Eur Heart J. 2012;33(16):2054–64.
- 71. Wolk K, Mallbris L, Larsson P, Rosenblad A, Vingård E, Ståhle M. Excessive body weight and smoking associates with a high risk of onset of plaque psoriasis. Acta Derm Venereol. 2009;89(5):492–7.
- 72. Tasliyurt T, Bilir Y, Sahin S, Seckin HY, Kaya SU, Sivgin H et al. Erectile dysfunction in patients with psoriasis: potential impact of the metabolic syndrome. Eur Rev Med Pharmacol Sci. 2014;18(4):581–6.
- 73. Van der Voort EAM, Koehler EM, Dowlatshahi EA, Hofman A, Stricker BH, Janssen HLA et al. Psoriasis is independently associated with nonalcoholic fatty liver disease in patients 55 years old or older: Results from a population-based study. J Am Acad Dermatol. 2014;70(3):517–24.
- 74.Nijsten T, Wakkee M. Complexity of the association between psoriasis and comorbidities. J Invest Dermatol. 2009;129(7):1601–3.

- 75. Parisi R, Rutter MK, Lunt M, Young HS, Symmons DPM, Griffiths CEM et al. Psoriasis and the Risk of Major Cardiovascular Events: cohort study using the Clinical Practice Research Datalink. J Invest Dermatol. 2015;135(9):2189–97.
- 76.Gupta MA, Schork NJ, Gupta AK, Kirkby S, Ellis CN. Suicidal ideation in psoriasis. Int J Dermatol. 1993;32(3):188–90.
- 77. Mahler V, Diepgen T, Skudlik C, Becker D, Dickel H, Fartasch M et al., and the Work Group "Assessment of allergens in occupational disease (BK) 5101" of the Study Group Occupational and Environmental Dermatology (ABD), German Contact Dermatitis Group (DKG) of the German Dermatological Society. Psoriasis predisposition and occupational triggering factors in the appraisal of occupational medical expertises. J Dtsch Dermatol Ges. 2014;12(6):519–29.
- 78. Skudlik C, John SM. Psoriasis and work. In: Kanerva's Occupational dermatology, 2nd edition (volume 1). Heidelberg, New York, Dordrecht, London: Springer; 2012.
- 79. Agrup G. Hand Eczema and Other Hand Dermatoses in South Sweden. J Occup Environ Med. 1970;12(2):59–60.
- 80. Weisshaar E, Skudlik C, Scheidt R, Matterne U, Wulfhorst B, Schönfeld M et al., and the ROQ Study Group. Multicentre study "rehabilitation of occupational skin diseases: optimization and quality assurance of inpatient management (ROQ)"—results from 12-month follow-up. Contact Dermatitis. 2013;68(3):169–74.
- 81.Richards HL, Fortune DG, Main CJ, Griffiths CEM. Stigmatization and psoriasis. Br J Dermatol. 2003;149(1):209–11.

- 82.Picardi A, Abeni D, Renzi C, Braga M, Puddu P, Pasquini P. Increased psychiatric morbidity in female outpatients with skin lesions on visible parts of the body. Acta Derm Venereol. 2001;81(6):410–4.
- 83.Pereira MG, Brito L, Smith T. Dyadic adjustment, family coping, body image, quality of life and psychological morbidity in patients with psoriasis and their partners. Int J Behav Med. 2012;19(3):260–9.
- 84.Eghlileb AM, Davies EEG, Finlay AY. Psoriasis has a major secondary impact on the lives of family members and partners. Br J Dermatol. 2007;156(6):1245–50.
- 85. Sampogna F, Gisondi P, Tabolli S, Abeni D, and the IDI Multipurpose Psoriasis Research on Vital Experiences investigators. Impairment of sexual life in patients with psoriasis. Dermatology. 2007;214(2):144–50.
- 86.Mansouri B, Patel M, Menter A. Biological therapies for psoriasis. Expert Opin Biol Ther. 2013;13(12):1715–30.