WORLD JOURNAL OF PHARMACEUTICAL AND MEDICAL RESEARCH

www.wjpmr.com

SJIF Impact Factor: 5.922

Review Article
ISSN 2455-3301
WJPMR

A SYSTEMATIC REVIEW OF PATHOPHYSIOLOGY, CLINICAL PRESENTATION, TYPES AND TREATMENT OF PSORIASIS

Neha Kushwaha, Monika Rai, Basant Khare*, Prateek Kumar Jain and Anushree Jain

Adina College of Pharmacy, ADINA Campus Rd, Lahdara, Sagar, MP, 470001.

*Corresponding Author: Basant Khare

Adina College of Pharmacy, ADINA Campus Rd, Lahdara, Sagar, MP, 470001.

Article Received on 31/09/2022

Article Revised on 21/10/2022

Article Accepted on 11/11/2022

ABSTRACT

Psoriasis is a chronic, multisystem inflammatory disease with predominantly skin and joint involvement. It has a bimodal age of onset (16 to 22 and 57 to 60 years)2 and affects both sexes equally. Beyond the physical dimensions of disease, psoriasis has an extensive emotional and psychosocial effect on patients, affecting social functioning and interpersonal relationships. As a disease of systemic inflammation, psoriasis is associated with multiple comorbidities, including cardiovascular disease and malignancy. The diagnosis is primarily clinical and a skin biopsy is seldom required. Depending on the severity of disease, appropriate treatment can be initiated. For mild to moderate disease, first-line treatment involves topical therapies including corticosteroids, vitamin D3 analogues, and combination products. In this review we have discussed about, etiology, pathophysiology, types, diagnostic methods and available treatment strategies for Psoriasis.

KEYWORDS: Psoriasis, Skin disorders, Chronic skin disease.

INTRODUCTION

Psoriasis chronic proliferative inflammatoryconditionofthe skin. Inflammation is a part of the body's immune response and is the end result of oxidative stress in any bodypart. Among the various inflammatory diseases psoriasis is found to be more severe in form, thoughit isnot infectious. The mostlyaffected partsinpsoriasisare the skin, nailsand joints. It comes under papulo-squamous disorders. Here, the outer layer of skin i.e. the epidermis moves towardsthe surface and thencontinuallyshed fromskin. The skin formationtouches a dramatically higher turnover rate. The name psoriasis is from the Greek language, meaning "roughly itching condition" (psora: "itch", sis: "action"). Psoriasis is an immune mediated disorder, where a normal skin cell mistakes for a pathogen, and sends a faulty signal that causes over production of new skin cell. It is also a hereditary condition but the way itinherits is still not predictable. It is a typically lifelong condition, which is not having a permanent cure, but various treatments can be implemented for controlling the severity of symptoms produced by it.^[1] The eye is involved in about 10% of patients, mostly women. In general, the eye is rarely involvedalone;

it is almost always associated with skin features.

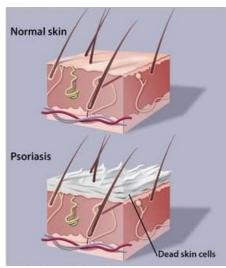


Figure 01: Psoriasis Skin.

Etiology of Psoriasis

Psoriasis has a prevalence ranging from 0.2% to 4.8%. The exact etiology is unknown, but it is considered to be an autoimmune disease mediated by T lymphocytes. There is an associationofHLA antigensseen in manypsoriatic patients, particularly in various racial and ethnic groups. Familial occurrence suggests its genetic predisposition. Injury in the form of mechanical, chemical, and radiational trauma induces lesions of psoriasis. Certain drugs like chloroquine, lithium, betablockers, steroids, and NSAIDs can worsen psoriasis. Generally, summer improves psoriasis while winter

www.wjpmr.com | Vol 8, Issue 12, 2022. | ISO 9001:2015 Certified Journal | 124

aggravates it. Apart from the above factors infections, psychological stress, alcohol, smoking, obesity, and hypocalcemia are other triggering factors for psoriasis. Psoriasis occurs worldwide, and its prevalence varies. In the United States, about 2% of the population is affected. High rates of psoriasis have been reported in the Faroe Islands. The prevalence of psoriasis is low in Japan and may be absent in Aboriginal Australians and Indians from South America. [2]

Disease initiation of complex diseases, such as psoriasis, takes place in genetically predisposed individuals in which advsregulated immune response occurs following exposure to certain environmental triggers. Although mechanistic associations linking distinct environmental factors with specific genetic determinants and dysregulated have been identified (Fig. 2).^[3]

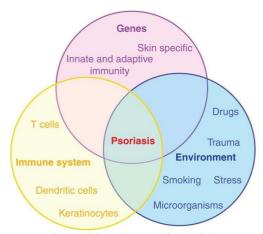


Figure 02: Etiology of Psoriasis.

Psoriasis can present at anyage. A bimodal age of onset has been recognized. The mean age of onset for the first presentation of psoriasis can range from 15 to 20 years of age, with a second peak occurring at 55 to 60 years.

Pathophysiology of Psoriasis

Thepathophysiologyofpsoriasis involves infiltrationoftheskinbyactivated Tcellswhich stimulate the proliferation of keratinocytes. This dysregulation in keratinocyte turnover results in the formation of thick plaques. Other associated features include epidermal hyperplasia and parakeratosis. In addition, the epidermal cells fail to secrete lipids which results in flaky and scaly skin, which is typical of psoriasis.^[4]

The pathophysiology of psoriasis is multi factoria land involves epidermal hyperproliferation, abnormal differentiation of epidermal keratinocytes, inflammation with immunologic alterations in the skin. The hyperproliferation is characterized by increased DNA synthesis and a markedly decreased turnover rate for the epidermis. Abnormal keratinocyte differentiation involves increased expression of certain keratins (6 and 16) and a delay in expression of other keratins (1 and 10) that are expressed in normally differentiating skin. [5]

Inflammation results from an infiltrate of neutrophils in the epidermis and super ficial dermis and an infiltrate of T lymphocytes in the dermis with a predominance of CD8+ cells.

NK cells and NKT cells in cutaneousdis orders:-

NKTcells

There is considerable interest in NKT cells and their role inthepathophysiologyofpsoriasis. Increased NKT cells were consistently observed in psoriasis lesions by different groups, although the exact role played by these cells is yet to be defined precisely. [7-10] Nickoloffandcoworkers demonstrated that psoriatic keratinocytes (KCs) overexpressed CD1d and NKT cells can be activated to elaborate IFNy when cultured with CD1d overexpressing KCs. These results provided a pathogenetic link between psoriatic KCs, which over expressCD1d and NKTcell sinfiltrating the lesions. Experiments in severe combined immunodeficient mice have also shown that injection of human cells of NKT characteristics into transplanted psoriatic skin could drive lesion development. Zhao et al. showed increased densities of NKT cells using a set ofprecise markers for classicalNKT cells, anti-Vα24 and anti-Vβ11 mAbs, inpsoriatic lesions, especially in thee pidermis, compared with healthy adult skin. Thesedata wereconfirmed byreal-timepolymerasechainreaction (PCR)too. Also, CD1dexpressionwas more extensive in psoriasis than in normal skin.

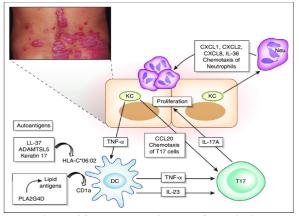


Figure 03: Pathophysiology of Psoriasis.

Symptoms and triggers of Psoriasis^[11-12]

Common signs and symptoms of psoriasis include:-

- A patchy rash that varies widely in how it looks fromperson to person, ranging from spots of dandruff-like scaling to major eruptions over much of the body.
- 2. Rashes that vary in color, tending to be shades of purple with gray scale on brown or Black skin and pink or red with silver scale on white skin.
- 3. Smalls calings pots(commonly seen in children).
- 4. Dry, cracked skin that may bleed.
- 5. Itching, burningor soreness.

Psoriasis triggers Many people who are predisposed to

www.wjpmr.com | Vol 8, Issue 12, 2022. | ISO 9001:2015 Certified Journal | 125

psoriasis may be free of symptoms for years until the disease is triggered by some environmental factor. Common psoriasis triggers include:

- Infections, suchasstrepthroat or skin in fections
- Weather, especially cold, dry conditions
- Injurytotheskin, suchasacut orscrape, abugbite, oraseveresunburn
- Smokingandexposuretosecondhandsmoke
- Heavyalcoholconsumption
- Certainmedications—includinglithium, highbloodpressuredrugsandantimalarialdrugs
- Rapidwithdrawaloforalorinjectedcorticosteroids

Complications^[13-15]

- Secondaryinfections
- Poorcosmesis
- Psoriaticarthritis
- Risk of lymphoma
- Increasedriskofadversecardiac events.
- Psoriaticarthritis, which causespain, stiffness, and swelling in and around the joints.
- Temporary skin color changes (postinflammatoryhypopig mentation or hyperpigmentation) where plaques have healed.
- Eyeconditions, suchasconjunctivitis, blepharitis and uveitis.
- Obesity.
- Type2 diabetes.
- Cardiovasculardisease.
- Other auto immunediseases, such as celiac disease, sclerosis and the inflammatory bowel disease called Crohn's disease.
- Mentalhealthconditions, such as lowself-esteemand depression.

Typesof Psoriasis

A) Plaque psoriasis: -The most common type of psoriasis, plaque psoriasis causes dry, itchy, raised skin patches (plaques) covered with scales. There may be few or many. They usually appear on the elbows, knees, lower back and scalp. The patches vary in color, depending on skin color. The affected skin might heal with temporary changes in color (post inflammatory hyperpigmentation), particularly on brown or Black skin. [16]



Figure 04: Plaque Psoriasis.

Symptoms

Plaquepsoriasis causesraised, inflamed, red skin coveredwith silvery, whitescales. These patches mayitchandburn. Itcanappearanywhereonyourbody, butitoftenpopsupintheseareas:

- Elbows
- Knees
- Scalp
- Lowerback.

Treatments

- Topicaltreatments: Thesego onyourskinandare usuallythe first thingdoctorstry. Some have steroids; others don't. Prescription products slow skin cell grow thand ease inflammation.
- **Phototherapy:** This treatment usesultravioletlight. You'llget it at your doctor's office or at home witha phototherapyunit.
- Systemic medications: These prescription drugs work throughout your body. You'llget them if you have moderate to severe psorias is that doesn't respond to other treatments. You could take the mbymouthorgetthemasashotor IV. Thiscategoryincludesdrugs called biologics, whichtarget specificpartsofyour immunesystemthat playarole inthe inflammatoryprocess. Learnmore about systemictreatments for psoriasis.
- B) Nail psoriasis: -Psoriasis can affect fingernails and toenails, causing pitting, abnormal nail growth and discoloration. Psoriatic nails might loosen andseparate from the nail bed (onycholysis). Severe disease may cause the nail to crumble When psoriasis affects the nails, you may notice:^[17]
- Tinydentsinyournails(called"nailpits")
- White, yellow, or brownd is coloration under oneormorenails
- Crumbling, roughnails
- A nail lifting up so thatit's no longerattached
- Buildup of skin cells beneath one or morenails, which lifts up the nail Treatment and proper nail care can help you control nail psoriasis.



Figure 05: Nailpsoriasis.

C) Guttate psoriasis:-Guttate psoriasis primarily affects young adults and children. It's usually triggered by a bacterial infection such as strep

throat. It's marked bysmall, drop-shaped, scalings pots on the trunk, armsorlegs. Classically, guttate psoriasis occurs shortly after an acute group B haemolytic streptococcal infection of thepharynx or tonsils and can be the presenting episode of psoriasis in children or, occasionally, adults. Thenumber of lesions mayrangefromfiveor 10toover Guttatepsoriasisaccounts for 2% ofthetotalcases ofpsoriasis. Inchildren, anacute episode of guttate psoriasis is usually self limiting; in adults, guttate flares may complicate chronic plaque disease. Although few studies haveassessedthelongtermprognosis ofchildren withacute guttatepsoriasis, onesmall study revealed that 33% of patients with acute guttate psoriasis eventually developed chronic plaque disease. [18]



Figure: 06 Guttate Psoriasis.

Guttatepsoriasishappensaftercertaintriggers. These triggers include:

- Strepthroat
- Stress
- Skininjury
- Infection
- Medication

Treatments Totreat guttate psoriasis, a doctor may prescribe steroid creams, light therapy, andoral medications. Determining the under lying cause of the infection can also help clear guttate psoriasis. If a bacterial infection caused the condition, antibiotics may help.

- D) Inversepsoriasis:-Inverse psoriasis mainlyaffects the skin folds of the groin, buttocks and breasts. It causes smooth patches ofinflamed skin that worsen with friction and sweating. Fungal infections may trigger this type of psoriasis. Where the inversepsoriasis appears, you're likely to notice: [19]
- Smooth, red patches of skin that look raw
- Little, ifany, silvery-whitecoating

Symptoms: Patches of skin that are bright red, smooth, and shiny, but don't have scales

Getting worse with sweating and rubbing Common triggers

are:

- Friction
- Sweating
- Fungal infections
- Soreorpainfulskin.



Figure 07: Inverse Psoriasis.

- E) Pustular psoriasis. Pustular psoriasis, a rare type, causes clearly defined pus-filled blisters. It can occur in widespread patches or on small areas of the palms or soles. Where pustular psoriasis appears, you tend to notice: [20]
- Red, swollen skin that is dotted with pus-filled bumps
- Extremelysoreor painfulskin
- Browndots(andsometimesscale)appear asthepusfilledbumpsdry

Pustularpsorias is can make just about any activity that requires your hands or feet, such as typing or walking, unbearably painful.

Symptoms include

- Fever
- Chills
- Nausea
- Fastheartrate
- Muscleweakness

Triggers include

- Topicalmedicine(ointments you put on your skin) or systemic medicine(drugs that treat your whole body), especially steroids
- Suddenlystoppingsystemic drugs or strong topical steroids that you used over a largearea of your body
- Getting too much ultraviolet(UV)light with out using sunscreen
- Pregnancy
- Infection
- Stress
- Exposuretocertain chemicals

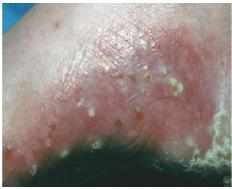


Figure: 08 Pustular psoriasis.

F) Erythrodermic psoriasis. The least common type of psoriasis, erythrodermic psoriasis can cover the entire **body** with a peeling rash that can itch or burn intensely. It can be short-lived (acute) or long-term (chronic). [21]

Symptomsinclude

- Severeitching, burning, orpeeling
- Afaster heartrate
- Changes inbodytemperature

If you have the sesymptoms, seeyour doctor rightaway. You may need to get treatedina hospital. This type of psoriasis can cause severe illness from protein and fluidloss. You may also get an infection, pneumonia, or congestive heart failure.

Triggers include

- Suddenly stopping your systemic psoriasis treatment
- Anallergicdrugreaction
- Severesunburn
- Infection
- Medications such as lithium, antimalarial drugs, cortisone, or strong coaltar products



Figure 09: Erythrodermic psoriasis.

G) Scalp Psoriasis

Scalppsoriasis is a common skindis order that makesraised, reddish, often scalypatches. It can popupasasing lepatch or several, and can even affect your entire scalp. It can also spread to your fore head,

thebackofyourneck, orbe hind and inside you rears. For some people, it may cause severe dandruff. For others, it can be painful, itchy, and very noticeable at the hairline. Scalp psoriasis can extend to your neck, face, and ears in one large patch or many smaller patches. In some cases, scalp psoriasis can complicate regular hair hygiene. Excessive scratching can cause hair loss and scalp infections. The condition may also cause feelings of social stress.^[22]

Symptoms

- Scaly, red, bumpypatches
- Silvery-whitescales
- Dandruff-likeflaking
- Dryscalp
- Itching
- Burningorsoreness



Figure: 10 Scalp Psoriasis.

Treatments Topical treatments are most commonly used for scalp psoriasis. They may requireaninitial 2months of intensive applications, plus permanent, regular maintenance. Treatment options include:

- Medicatedshampoos
- Tarpreparations
- Topical application of vitamin D, know nas calcipotriene (Dovonex)

Light therapy, oral medications, and biologics also may be recommended depending on the responsiveness to treatment.

H) Psoriatic arthritis Psoriatic arthritis (PsA) is a painful and physically limiting conditionthat affectsbetween30% and 33% of people with psoriasis. There are fivetypes of PsA with varying symptoms. There's also no cure for this condition.

Psoriasis is anautoimmune disease, so it cantrigger your body to attack your joints and skin. It can affect many joints and often becomes quite severe in the hands and affect the nails.

Skin symptoms usually appear before joint symptoms. [23]



Figure 11: Psoriatic arthritis.

Treatments

Treatments for psoriatic arthritis can include non steroidal anti-inflammatory drugs(NSAIDs), such as ibuprofen(Advil) and naproxen sodium(Aleve). NSAID scan help reduce the incidences of swelling and pain associated with psoriatic arthritis.

Prescription medications, such as an oral corticosteroid, may also help reduce inflammation that leads to psoriatic arthritis. Prescription topical medications used to treat psoriatic arthritis include salicylic acid, calciopotriene, and tazarotene.

A unique category of medications known as disease-modifying antirheumatic drugs (DMARDs) can help reduce inflammation and joint damage. Biologics, which are a sub category of DMARDs, maybe prescribed to reduce inflammation at a cellular level.

Causes:-Psoriasis is thought to be an immune system problem that causes skin cells to growfaster thanusual. In the most commontype of psoriasis, knownasplaque psoriasis, this rapid turnover of cells results in dry, scaly patches.

The cause of psoriasis isn't fully understood. It's thought to be an immune system problem where infection-fighting cells attack healthy skin cells by mistake.

Diagnosis

Your health care provider will ask questions about your health and examine your skin, scalp and nails. Your health care provider then might take a small sample of skin (biopsy) for examination under a microscope. This helps determine the type of psoriasis and rule out other disorders. The diagnosis ofpsoriasis is primarilyclinical. There are different clinicaltypes of psoriasis, the most commonofwhich is chronic plaque psoriasis, affecting 80% to 90% of patients with psoriasis. The hallmark of classic plaque psoriasis is well-demarcated, symmetric, and erythematous plaques with overlying silvery scale. Plaques are typicallylocatedonthe scalp, trunk, buttocks, and extremities but canoccuranywhere onthe body.

Patients might demonstrate nail involvement, which can present without concomitant plaques. Active lesions might be itchy or painful. Psoriasis can also present as an isomorphic response, where new lesions develop on previously normal skin that has sustained trauma or injury. The severityof disease can be helpful in guiding management and is classified as mild, moderate, and severe.

Treatment of Psoriasis

Psoriasis treatments aim to stop skin cells from growing so quickly and to remove scales. Options include creams and ointments (topical therapy), light therapy (phototherapy), andoral or injected medications. Which treatments you use depends on how severe the psoriasisis and how responsive it has been to previous treatment and self-care measures. You might need to try different drugs or a combination of treatments before you find an approach that works. Even with successful treatment, usually the disease returns.Researchers believe that both genetics and environmental factors play a role. [24-27]

Topicaltherapy

- Corticosteroids. These drugs are the most frequently prescribed medications for treating mild to moderate psoriasis. They are available as oils, ointments, creams, lotions, gels, foams, sprays and shampoos. Mild corticosteroid ointments (hydrocortisone) are usually recommended for sensitive areas, such as the face or skin folds, and for treating widespread patches. Topical corticosteroids might be applied once a day during flares, and on alternate days or weekends during remission.
- Vitamin D analogues. Synthetic forms of vitamin D- such as calcipotriene (Dovonex, Sorilux) and calcitriol(Vectical) slow skincellgrowth. This type ofdrug may be used alone or withtopicalcorticosteroids. Calcitriolmay cause less irritation insensitive areas. Calcipotriene and calcitriol are usually more expensive than topical corticosteroids.
- Retinoids. Tazarotene (Tazorac, Avage, others) is available as a gel or cream. It's applied once or twice daily. The most common side effects are skin irritation and increased sensitivity to light. Tazarotene isn't recommended when you're pregnant or breastfeeding or if you intendto become pregnant.
- Calcineurin inhibitors. Calcineurin inhibitors —
 such as tacrolimus (Protopic) and pimecrolimus
 (Elidel) calm the rash and reduce scaly buildup.
 They can be especiallyhelpful inareasofthinskin,
 suchasaround theeyes, wheresteroid creamsor
 retinoids are irritating or harmful.
- Salicylic acid. Salicylic acid shampoos and scalp solutions reduce the scaling of scalp psoriasis. They are available in nonprescription or prescription strengths. This type of product may be used alone or with other topical therapy, as it prepares the scalp to absorb the medication more easily.

- Coal tar. Coal tar reduces scaling, itching and inflammation. It's available in nonprescriptionand prescriptionstrengths. It comes invarious forms, suchasshampoo, cream and oil. These products can irritate the skin. They're also messy, stain clothing and bedding, and can have a strong odor.
 - Coaltartreatmentisn'trecommended when you'repregnant or breast feeding.
- Anthralin. Anthralin is a tar cream that slows skin cell growth. It can also remove scales and make skin smoother. It's not intended for use on the face or genitals. Anthralin can irritate skin, and it stains almost anything it touches. It's usually applied for a short time and then washed off.
- **Light therapy** -Light therapy is a first line treatment for moderate to severe psoriasis, either alone or in combination with medications. It involves exposing the skin to controlled amounts of natural or artificial light. Repeated treatments are necessary.
- **Sunlight-**Brief, daily exposures to sunlight (heliotherapy) might improve psoriasis. Before beginning a sunlight regimen, ask your health care provider about the safest way to use natural light for psoriasis treatment.
- Goeckerman therapy. Anapproachthat combinescoaltartreatment with light therapy is called the Goeckerman therapy. This can be more effective because coal tar makes skin more responsive to ultraviolet B (UVB) light.
- UVB broadband. Controlled doses of UVB broadband light from an artificial light source cantreat single psoriasis patches, widespread psoriasis and psoriasis that doesn't improve with topical treatments. Short-term side effects might include inflamed, itchy, dry skin.
- UVB narrowband.

 UVBnarrowbandlighttherapymightbemoreeffective thanUVBbroadband treatment. In many places it has replaced broadband therapy. It's usually administered two or three times a week until the skin improves and then less frequently for maintenance therapy
- PsoralenplusultravioletA(PUVA). Thistreatment involvestakinga light-sensitizing medication (psoralen)before exposing the affected skin to UVAlight.UVAlight penetrates deeper into the skin than does UVBlight, and psoralen makes the skin more responsive to UVA exposure.
- Excimer laser. With this form of light therapy, a strong UVB light targets only the affected skin.
 Excimer laser therapy requires fewer sessions than does traditional phototherapy because morepowerfulUVB light is used. Side effects might include inflammation and blistering.

Drugs for treatment of psoriasis

During 18th & 19th centuries some flower solutions containing poisonous and carcinogenic arsenic compound was used by dermatologist as a treatment for psoriasis. Grenz ray (ultra soft x-ray) was popular

- treatment for psoriasis during middle age of 20th century. Undencylenic acid was investigated and used for psoriasis some 40 year ago. [28]
- Antibiotics:-Tetracycline and Penicillin Use of systemic antibiotics and induction of gh reduction of intracellular cAMP and bythe interaction with arachidonic acid and its metabolites.
- β-blockers:-It is a very popular class of drug to treat cardiovascular diseases like Arrhythmias, hypertension, ischemic heart disease, heart failure, hyperthyroidism, glaucoma and anxiety. Their action is exerted by blocking the beta receptor or non selective β2 receptor, mainly found in the keratinocyte and on the surface of the macrophages. een theorized that tetracyclines accumulate in higher concentrations in psoriatic lesions compared to uninvolved skin.
- **Lithium:**-Lithium is used for the treatment of manic depressive disorder. In 1972, the first lithium induced psoriasis was reported. There are several therapies purported to explain the pathogenesis of lithium provoked psoriasis.
- Antimalarial Drug:-The most commonly used antimalarial drugs are chloroquine and hydroxychloroqine. The mechanism of causing psoriasis is through inhibition of transglutaminase enzyme in skin.

Oral or injected medications

Ifyou have moderateto severepsoriasis, orifothertreatmentshaven't worked, your health care provider may prescribe oral or injected (systemic) drugs. Some of these drug sare used for only brief periods and might be alternated with other treatments because they have potential for severe side effects. [29-30]

- **Steroids.** If you have a few small, persistent psoriasis patches, your health care provider might suggest an injection of triamcinolone right into them.
- Retinoids. Acitretin an do ther retinoids are pills usedt or educe the production of skin cells. Side effects might include dryskin and muscle soreness. These drugs are not recommended when you're pregnant or breastfeeding or if you intend to become pregnant.
 - Biologics. These drugs, usually administered by injection, alter the immune systemina way that disrupts the disease cycle and improves symptoms and signs of disease within weeks. Several of these drugs are approved for the treatment of moderate to severe psoriasis in people who haven't responded to therapies. first line Options include apremilast(Otezla), etanercept(Enbrel), infliximab(Remicade), adalimumab(Humira), ustekinumab (Stelara), secukinumab (Cosentyx), ixekizumab (Taltz), guselkumab (Tremfya), tildrakizumab (Ilumya) and certolizumab (Cimzia). Three of them etanercept, ixekizumabandustekinumab—are approved children. These types of drugs are expensive and

may or may not be covered by health insurance plans.

Biologics must be used with caution because they carrythe risk of suppressing the immune system in ways that increase the risk of serious in fections. People taking these treatments must be screened for tuberculosis.

- Methotrexate. Usually administered weekly as a single oral dose, methotrexate (Trexall)decreases the of skin cells and inflammation. It'sless effective than adalimumab and infliximab. It might cause upset stomach, loss of appetite and fatigue. People taking methotrexate long-term need ongoing testing to monitor their blood counts and liver function.
 - People need to stop taking methotrexate at least three months before attempting to conceive. This drug is not recommended for those who are breastfeeding.
- Cyclosporine. Taken orally for severe psoriasis, cyclosporine (Gengraf, Neoral, Sandimmune) suppresses the immune system. It's similar to methotrexate in effectiveness but cannot be used continuously for more than a year. Like other immune suppressant drugs, cyclosporine increases the risk of infection and other health problems, including cancer. People taking cyclosporine longterm need on going testing to monitor their blood pressure and kidney function.
 - These drugs aren't recommended when you'r epregnant or breast feeding or if you intend to become pregnant.
- Other medications. Thioguanine (Tabloid) and hydroxyurea (Droxia, Hydrea) are medications that can beused when you can't take other drugs. Talk with your health care provider about possible side effects of these drugs.

Treatment considerations

You and your health care provider will choose a treatment approach based on your needs and the type and severity of your psoriasis. You'll likely start with the mildest treatments - topical creams and ultraviolet light therapy (phototherapy). Then, if your condition doesn't improve, you might move on to stronger treatments.

People with pustular or erythrodermic psoriasis usually need to start with stronger(systemic) medications.

In any situation, the goal is to find the most effective way to slow cell turn over with the fewest possible side effects.

Lifestyle and homeremedies

Trytheseself-care measurestobettermanageyour psoriasis:

- Takedailybaths. Washgentlyratherthanscrubbing yourskinintheshowerorbath. Use lukewarmwater and mild soaps that have added oils or fats. It might help to add bath oil, Epsom salts or oatmeal to bathwater and soak for at least 15 minutes.
- Keep your skin moist. Apply moisturizer daily. If you're moisturizing after bathing, gentlypat

dryandapplyyour preferredproductwhile your skinisstillmoist. Forvery dryskin, heavyointment-based moisturizers may be preferable — theystayon the skin longer thancreams orlotions do. Ifmoisturizing seems to improve your skin, apply the product more than once a day.Iftheair whereyouliveisverydry,

useahumidifiertoaddmoisturetotheair.

- Cover the affected areas overnight. Before going bed. apply ointment-based an moisturizertotheaffectedskinandwrapwithplasticwra p.Whenyouwake, remove the plastic and wash away
- Exposeyourskin tosmallamountsofsunlight. Ask your healthcareprovider about thebest wayto usenaturalsunlight totreat your skin. Acontrolled amount of sunlight can improve psoriasis, but too much sun can trigger or worsen outbreaks and increase the risk ofskincancer. Logyourtime inthe sun, and protect skinthat isn't affected by psoriasis with a hat, clothing or sunscreen with asunprotection factor(SPF)ofat least 30.
- Avoidscratching. Itmight helpto nonprescription anti-itch cream or ointment containing hydrocortisone or salicylicacid. If you have scalppsoriasis, try a medicated shampoo that contains coal tar. Keep your nails trimmed so that theywon't hurt your skin if you do scratch. Wear soft fabrics that don't contribute to itchiness.
- Avoidpsoriasistriggers. Notice what triggers your psoriasis, and take step stoprevent or avoid it. Infections, injuries to your skin, smoking and intense sun exposure can all worsen psoriasis.
- Stay cool. Being too hot can make your skin feel itchy. Wear light clothing if you're outside onhot days. If you have air conditioning, use it onhot days to keep cool. Keep cold packs in your freezer and apply themto itchy spots for a few minutes ofrelief. Youmight trystoring yourmoisturizing lotionintherefrigeratortoaddacoolingeffect when you apply it
- Strive to maintaina healthy Trypracticingo ther healthy-living habits to help man agepsoriasis. These include being active, eatingwell, limiting or avoiding alcohol consumption, and maintaining a healthy weight.

Alternative and natural treatments for psoriasis

Some studies claim that alter native the rapies(integrative medicine) -products and practices not part of conventional medical care or that developed outside of traditional Western practice — ease the symptoms ofpsoriasis. Examples ofalternative therapies used bypeople with psoriasis include special diets, vitamins, acupuncture and herbalproducts applied to the skin. None of these approaches is backed by strong evidence, but theyare generallysafe and might help reduce itching and scaling in people with mild to moderate psoriasis. [3]-

Aloe extract cream. Taken from the leaves of the

aloe vera plant, aloe extract cream mayreducescaling, itchingand inflammation. You might need to use the cream several times a day for a month or more to see any improvement in your skin.

- **Fish oil supplements.** Oral fish oil therapy used in combination with UVB therapy might reduce the exten to fthe rash. Applying fish oil to the affected skin and covering it with a dressing for six hours a day for four weeks might improve scaling.
- **Oregongrape.** Oregongrape—also known as barberry— is applied to the skin and may reduce the severity of psoriasis.

If you'r econsidering alternative medicine to easethesigns and symptoms of psoriasis, talk with your health care provider about the pros and cons of these approaches.^[35]

The HerbalMedicines not have more side effects as compared to synthetic drugs. The herbal medicine is easily available and easy to use in treatment. Now a day, herbal resources play a very important role in the management of the skin and inflammatory diseases. Some studies suggest that psoriasis symptoms can be relieved by change in diet and life style. Fasting food period, low energy diet and vegetarian diets have improved psoriasis symptoms. In some treatments supplemented with fish oil shows a beneficial effect dueto the presence ofomega - 3 FattyAcids and Vitamin E. Cannabis is also suggested for treating psoriasis due to Anti - inflammatory properties of its canabinoids and their regulatory effect on immune system. [36-37]

- Cayenne, its chief component being capsaicin. One hypothesis on the pathogenesis of psoriasis suggests a neurogenic inflammatory etiology mediated through substance P (SP). SP activates inflammatory cells and ultimately perpetuates vasodilatation, angiogenesis and keratinocyte hyperproliferation. In accordance, psoriatic lesions are known to be more densely innervated with higher SP content than control or uninvolved psoriatic skin.
- Aloe vera: -Aloe vera is a popular plant used in cosmetic care and first aid products in case of thermal injuries. Aloe contains anthroquinones, saponins, mucopolysaccharides salicylic acid. Syed and colleagues (1996) conducted a double - blind, placebo - controlled studyon 60 patients with psoriasis with slight to moderate plague type psoriasis and an average 8.5 year duration of their disease. Patients'self - administered topical Aloevera extract reamorve hicleplacebo three times a day without occlusion for 4 weeks to the irpsoriaticplaques. The aloegroup significantly higherrates of clearing the psoriatic plaquesinal most all patients. Anthraquin one andacemannan, the main active compoundsin Aloevera, have antibacterial activity against Staphylococcus and Strep coccus to speciesandmayprovidearationale for their

- rapeuticefficacyinpsoriasis. Inaddition, salicylicacid, acomponent of Aloevera, isakeratolyticandwouldcontributetoits reported efficacy in the desquamationofpsoriaticplaques.
- Matricariarecutita: -It is commonly known as Chamomile. The chamomile flowers have a long therapeutic tradition in treating gastrointestinal ailments. The rationale for its use in psoriasis is that chamazulene, a by-product ofthe non - volatile oil extract. matricin, known haveantito inflammatoryactivityby inhibition oflipoxygenaseand asaresult, leukotriene B4 (LTB4) formation. There is evidence supporting the role of increased LTB4 formation in psoriatic plaques; therefore, inhibition results in disease improvement.
- 4) Curcumadomestica: -Turmerichasa long history of being used for infections and kidney stones. The use in psoriasis is a relatively new adjunct. The antiinflammatory components are thought to be contained in the curcuminoids and volatile oils which function through selective inhibition of phosphorylase kinase(PhK). PhKisanenzy me found intheepidermis.
- **Gaultheria procumbens:** -It is commonly known as Wintergreen. Wintergreen is a plant native to the Eastern United States and historically was used by Native Americans as an analgesic. Al though used topically for psoriasis, winter green can cause systemic effects like tinnitus, vomiting, tachypnea and acid-based is turbances. Patientsusing as pirinoraprescribed salicylic acid compound in conjunction with a salicylate herbal(forexample, winter green, aloevera, orredcl over) are more susceptible for systemict oxicities. Additionally, oil of winter green can increase prothrombin time and international normalized ratio(INR) of clotting, creating problems for patients onwarfarin. There are noinvestigation son it seffective nessin psoriasis, but have potentialanti- inflammatory effect and needs further scientific investigations for it suseinpsoriasis.

CONCLUSION

Psoriasis is a complex multifactorial disease for which various novel therapies have arisen in the past years. Many patients with psoriasis seek initial evaluation and treatment from their primary care providers. Recognition of psoriasis, as well as its associated medical and psychiatric comorbidities, would facilitate timely diagnosis and appropriate management with effective and safe topical therapies and other medical and psychological. In spite of the refinement of the targeted therapies, psoriasis remains a treatable but so far not curable disease. The targeted therapies show high clinical efficacy for the inhibition of IL-23 and IL-17. Some degree of a persistent antipsoriatic effect by these could demonstrated be discontinuation, and argue for disease modification concept. This important finding will be followed up in ongoing and future studies.

www.wjpmr.com | Vol 8, Issue 12, 2022. | ISO 9001:2015 Certified Journal | 132

REFERENCES

- 1. Elman SA, Weinblatt M, Merola JF. Targeted therapies for psoriatic arthritis: an update for the dermatologist. Semin Cutan Med Surg, Sep, 2018; 37(3): 173-181.
- 2. YiuZZ, WarrenRB. Ustekinumab for the treatment of psoriasis:anevidence update. Semin Cutan Med Surg., Sep, 2018; 37(3): 143-147.
- 3. Yang EJ, Beck KM, Sanchez IM, Koo J, Liao W. The impact of genital psoriasis on quality of life: a systematic review. Psoriasis (Auckl), 2018; 8: 41-47.
- Gamret AC, Price A, Fertig RM, Lev-Tov H, Nichols AJ. Complementary and Alternative Medicine Therapies for Psoriasis: A Systematic Review. JAMADermatol, Nov 01, 2018; 154(11): 1330-1337
- Nguyen CT, Bloch Y, Składanowska K, Savvides SN, Adamopoulos IE. Pathophysiology and inhibition of IL-23 signaling in psoriatic arthritis: A molecularinsight. Clin Immunol, Sep, 2019; 206: 15-22.
- Larsabal M, Ly S, Sbidian E, Moyal- Barracco M, Dauendorffer JN, Dupin N, Richard MA, Chosidow O, Beylot-Barry M. GENIPSO: a Frenchprospective studyassessing instantaneous prevalence, clinicalfeatures and impact onqualityoflife ofgenitalpsoriasis among patients consulting for psoriasis. Br J Dermatol, Mar, 2019; 180(3): 647-656.
- Eder L, Widdifield J, Rosen CF, Cook R, Lee KA, Alhusayen R, Paterson MJ, Cheng SY, Jabbari S, Campbell W, Bernatsky S, Gladman DD, Tu K. Trends in the Prevalenceand Incidence of Psoriasis and Psoriatic Arthritis in Ontario, Canada: A Population-Based Study. Arthritis Care Res (Hoboken), Aug, 2019; 71(8): 1084-1091.
- 8. Kahn J, Deverapalli SC, Rosmarin D. JAK-STAT signaling pathway inhibition: a role for treatment of various dermatologic diseases. Semin Cutan Med Surg, Sep, 2018; 37(3): 198-208.
- 9. Caiazzo G, Fabbrocini G, Di Caprio R, Raimondo A, Scala E, Balato N, Balato A. Psoriasis, CardiovascularEvents, and Biologics:Lights and Shadows. Front Immunol, 2018; 9: 1668.
- 10. Trayes KP, Savage K, Studdiford JS. Annular Lesions: Diagnosis and Treatment. Am Fam Physician, Sep 01, 2018; 98(5): 283-291.
- 11. Vázquez-Herrera NE, Sharma D, Aleid NM, Tosti A. Scalp Itch: A SystematicReview. Skin Appendage Disord, Aug, 2018; 4(3): 187-199.
- 12. Perez-Chada LM, Cohen JM, Gottlieb AB, Duffin KC, Garg A, Latella J, Armstrong AW, Ogdie A, Merola JF. Achieving international consensusonthe assessment ofpsoriatic arthritis in psoriasis clinical trials: an International Dermatology Outcome Measures (IDEOM) initiative. Arch Dermatol Res., Nov. 2018; 310(9): 701-710.
- 13. Schadler ED, OrtelB, MehlisSL. Biologicsforthe primarycare physician: Review and treatment of psoriasis. Dis Mon, Mar, 2019; 65(3): 51-90.
- 14. Dauden E, Blasco AJ, Bonanad C, Botella R,

- Carrascosa JM, González-Parra E, Jodar E, Joven B, Lázaro P, Olveira A, Quintero J, Rivera R. Position statement for the management of comorbidities in psoriasis. J EurAcad Dermatol Venereol, Dec, 2018; 32(12): 2058-2073.
- 15. Luchetti MM, Benfaremo D, Campanati A, Molinelli E, Ciferri M, Cataldi S, CapeciW, Di Carlo M, Offidani AM, Salaffi F, Gabrielli A. Clinical outcomes and feasibility of the multidisciplinary management of patients with psoriatic arthritis: two-year clinical experience of a dermo-rheumatologic clinic. Clin Rheumatol, Oct, 2018; 37(10): 2741-2749.
- 16. Gelf and JM, BerlinJ, VanVoorhees A, Margolis DJ. Lymphoma rates are low but increased in patients with psoriasis: results from apopulation-basedcohort study in the United Kingdom. Arch Dermatol, 2003; 139(11): 1425–9.
- 17. Russo PA, IlchefR, CooperAJ. Psychiatric morbidityin psoriasis:areview. AustralasJ Dermatol, 2004; 45(3): 155–918.
- 18. Menter A, Korman NJ, Elmets CA, Feldman SR, Gelfand JM, Gordon KB, et al. Guidelines of care for the management of psoriasis and psoriatic arthritis. Section 3. Guidelines of care for the management and treatment of psoriasis with topical therapies. J Am Acad Dermatol, 2009; 60(4): 643–59.
- 19. MasonAR, MasonJ, CorkM, DooleyG, EdwardsG. Topical treatments for chronic plaque psoriasis. Cochrane Database Syst Rev., 2009; 2: 05-28.
- AshcroftDM, PoAL, WilliamsHC, Griffiths CE. Systematic review of comparative efficacy and tolerability of calcipotriol in treating chronic plaque psoriasis. BMJ., 2000; 320(7240): 963–7.
- 21. BakerH. The influence of chloroquine and related drugs on psorias is and kertoderm ablenorrhagium. BrJ Dermatol., 1966; 78: 161–166.
- 22. Bell AJ, Duggin G. Acute methyl salicylate toxicity complicating herbal skin treatment for psoriasis. Emergency Medicine, 2002; 14: 188-190.
- 23. Ben-AryeE, ZivM, FrenkelM, LaviI, RosenmanD. Complementary medicine and psoriasis:Linking the patient's out look with evidence-based medicine. Dermatology, 2003; 207: 302-307.
- 24. BernsteinJE, ParishLC, Rapaport M, RosenbaumMM, RoenigkHHJr. Effects of topically applied capsaicinon mode rate and severe psoriasis vulgaris. JA AcadDermatol., 1986; 15: 504-507.
- Bloomfield FJ, Young MM. Enhanced release of inflammatory mediators from lithium-stimulated neutrophils inpsoriasis. BrJDermatol., 1983; 109: 9–13.
- 26. BrownAC, Hairfield M, Richards DG, McMillinDL, MeinEA, Nelson CD. Medical nutrition therapy as a potential complementary treatment for psoriasis – Five case reports. Altern Med Rev., 2004; 9: 297-307.
- 27. Brown AC, HairfieldM, RichardsDG, McMillinDL, MeinEA, NelsonCD. Medical nutrition therapy as a potential complementary treatment for psoriasis—

- fivecasereports. AlternMedfev, 2004; 9: 297-307.
- 28. BrownDJ, DattnerAM. Phy to therapeuticapproaches to common derma to logic conditions. Archives of Dermatology., 1998; 134: 1401-1404.
- 29. CarsonC, RileyTV, CooksonBD. Efficacy and safety of tea tree oilasa topical antimicrobial agent. JHospInfect, 1998; 40: 175-178.
- 30. CarsonCF, AshtonL, DryL, SmithDW, RileyTV. Melaleuca alternifolia(teatree)oilgel(6%) for the treatment of recurrent herpes labialis. J Antimicrob Chemoth, 2001; 48: 450-451.
- 31. Carter TN. The relationship of lithium carbonate topsoriasis. Psychosomatics, 1972; 13: 325–327.
- 32. Choonhakarn C, Busaracome P, Sripanidkulchai B, Sarakarn P.A prospective, randomized clinical trial comparing topical Aloevera with 0.1% triamcinolone acetonidein mild to mode rate plaque psoriasis. JEurAcad DermatolVenereol, 2010; 24: 168-72.
- 33. CohenAD, BonnehDY, ReuveniM, VardyDA, NaqqanL, HalevyS. Drug exposure and psoriasis vulgaris: case-control and case-crossover studies. ActaDermVenereol., 2005; 85: 299-303.
- 34. CounisR, KoumanovK, RaulinJ, Antilipolyticrole of tetracycline. Inhibition of adenylatecyclaseinvitro. EurJ Biochem., 1973; 37: 244-247.
- 35. DengS, MayBH, ZhangAL, LuC, XueCCL. Topic alherbal medicine combined with pharmaco therapy for psoriasis:asystematic reviewandmetaanalysis. Arch Dermatol Res., 2013; 305: 179–89.
- 36. Dhanabal SP, Anand R, MurugananthamN, PraveenTK, Raghu PS. Screening of Wrightia tinctoria leaves for Anti-psoriatic activity. HygeiaJDMed., 2012; 4: 73-78.
- 37. DhanabalSP, Priyanka DwarampudiL, Muruganantham N, Vadivelan R. Evaluation of the anti psoriatic activity of Aloevera leaf extractusing amouse tail model of psoriasis. Phytother Res., 2012; 26: 617-19.

ISO 9001:2015 Certified Journal