

Blush Beyond The Skin: Healing Paths In Rosacea

Yerni Jyothi Kolli*, Sheeba Anitha Rani, Anusuya Devi

Medical Surgical Nursing, Driems University, Tangi, Cuttack

ABSTRACT

Rosacea is a chronic dermatological condition characterized by recurrent facial flushing, erythema, and inflammatory lesions, often influenced by genetic predisposition and environmental triggers. This report summarizes current treatment options for rosacea, drawing from published literature on its management. Patients are advised to avoid physiological and environmental stimuli that exacerbate facial redness. Topical therapies, including metronidazole, azelaic acid, and sulphur preparations, remain effective first-line options. For individuals with erythematotelangiectatic or papulopustular rosacea, oral antibiotics such as tetracycline are beneficial, while isotretinoin may be reserved for severe or refractory cases. Maintenance of remission can be achieved with continued topical therapy. Ocular rosacea requires long-term antibiotic use alongside topical metronidazole gel. Surgical interventions, particularly laser therapy, are recommended for telangiectasias and rhinophyma. Additionally, cosmetic camouflage offers psychological relief for patients distressed by visible symptoms. With the availability of diverse oral, topical, and procedural treatments, rosacea can be effectively managed, enabling patients to maintain both dermatological health and quality of life.^{1,3}

Keywords: erythematotelangiectatic, papulopustular rosacea, facial flushing, erythema.

INTRODUCTION

Rosacea is a chronic inflammatory skin disorder that primarily affects the central face, manifesting as recurrent flushing, persistent erythema, papules, pustules, and telangiectasias. It is a condition with a variable course, often triggered by environmental, physiological, and lifestyle factors. Although not life-threatening, rosacea significantly impacts patients' quality of life due to its visible nature and associated psychosocial distress. The disorder is commonly underdiagnosed and misinterpreted as acne or other dermatological conditions, leading to delayed treatment. With the availability of diverse therapeutic options, ranging from topical agents to systemic medications and procedural interventions, effective management is possible. This review aims to consolidate current evidence on treatment strategies for rosacea, highlighting their clinical utility and relevance for patient care.^{5,11}

CASE SCENARIO :

Mrs. S, a 42-year-old school teacher, presented with a five-year history of recurrent facial flushing that worsened with sun exposure and spicy food. She reported persistent redness over her cheeks and nose,

accompanied by occasional papules and pustules, along with a burning sensation after outdoor activities. Her family history was notable for similar symptoms in her mother. On examination, erythematotelangiectatic rosacea was evident with visible telangiectasias on the cheeks and papulopustular lesions scattered over the nose and chin.

BACKGROUND OF THE STUDY

Rosacea is a chronic, relapsing dermatological condition that predominantly affects the central facial region, presenting with recurrent flushing, persistent erythema, papules, pustules, and telangiectasias. Its etiology is multifactorial, involving vascular hyperreactivity, immune dysregulation, microbial factors such as *Demodex* mites, and genetic predisposition. Although rosacea is not life-threatening, its visible manifestations often result in significant psychosocial distress, reduced self-esteem, and impaired quality of life. Globally, rosacea is more frequently observed in fair-skinned populations of Northern European descent, but it can affect individuals across all ethnic groups. The condition typically appears in adulthood, with a

Relevant conflicts of interest/financial disclosures: The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

higher prevalence among women, though men are more prone to severe phymatous changes such as rhinophyma. Clinically, rosacea is categorized into four major subtypes: erythematotelangiectatic, papulopustular, phymatous, and ocular, each requiring tailored management strategies.^{5,11}

Despite the availability of multiple therapeutic options, rosacea remains underdiagnosed and often mismanaged due to its resemblance to acne and other inflammatory skin disorders. Patients frequently experience frustration with recurrent flare-ups and inadequate control. The chronicity of the disease, combined with its cosmetic implications, underscores the need for comprehensive reviews that consolidate evidence-based treatment approaches. Such reviews not only guide clinicians and nursing professionals in selecting appropriate interventions but also highlight the importance of patient education, lifestyle modification, and psychological support in holistic care.^{5,11}

REVIEW OF LITERATURE

Literature Review related to Procedural and Supportive Interventions

The research study reviewed procedural options such as laser therapy and intense pulsed light (IPL) for the management of telangiectasias and rhinophyma. Their analysis demonstrated that these interventions are particularly effective in reducing vascular lesions and improving cosmetic appearance. Furthermore, the study highlighted the importance of psychological support and cosmetic camouflage in patients experiencing distress due to visible symptoms. The authors stressed that combining medical treatment with supportive measures enhances both dermatological control and quality of life.^{2,9}

Literature Review related to Topical and Systemic Therapies

A study emphasized the effectiveness of topical agents such as metronidazole and azelaic acid in reducing erythema and inflammatory lesions in rosacea. Their findings highlighted that topical therapy remains the cornerstone of initial management, particularly in mild to moderate cases. In addition, systemic antibiotics like tetracyclines were shown to provide significant improvement in

papulopustular rosacea, with isotretinoin reserved for severe or refractory cases. The authors concluded that a stepwise approach, beginning with topical agents and progressing to systemic therapy when necessary, ensures optimal patient outcomes.^{2,9}

A systematic review analyzed 43 clinical trials involving 18,347 patients with rosacea. The study found that oral isotretinoin was the most effective therapy for reducing inflammatory lesions, particularly in severe and recalcitrant cases. Topical agents such as ivermectin, metronidazole, and azelaic acid demonstrated consistent efficacy in mild to moderate disease, while oral antibiotics like minocycline and doxycycline were effective for papulopustular rosacea. The review emphasized the role of *Demodex folliculorum* mites in pathogenesis and highlighted the need for combination therapies to improve long-term outcomes.^{2,9}

Literature Review related to Advancement in Targeted and Emerging Therapies

A systematic review evaluated 15 studies with 537 rosacea patients and 77 controls. The review highlighted vascular-targeted therapies (oxymetazoline, pulsed-dye laser, platelet-rich plasma) and microbiome-oriented approaches as promising new strategies. Established agents like ivermectin, azelaic acid, dapsone, sulphur preparations, and metronidazole continued to show benefit, while tofacitinib (a JAK inhibitor) and oral ivermectin emerged as potential options for refractory cases. The authors concluded that a multimodal, mechanism-based approach—addressing vascular, inflammatory, immunological, and microbial pathways—offers the best outcomes. However, they stressed the need for large randomized controlled trials to validate long-term efficacy and safety. A systematic review analyzed 43 clinical trials involving 18,347 patients with rosacea. The study found that oral isotretinoin was the most effective therapy for reducing inflammatory lesions, particularly in severe and recalcitrant cases. Topical agents such as ivermectin, metronidazole, and azelaic acid demonstrated consistent efficacy in mild to moderate disease, while oral antibiotics like minocycline and doxycycline were effective for papulopustular rosacea. The review emphasized the role of *Demodex folliculorum* mites in pathogenesis

and highlighted the need for combination therapies to improve long-term outcomes.^{2,9}

A systematic review evaluated 15 studies with 537 rosacea patients and 77 controls. The review highlighted vascular-targeted therapies (oxymetazoline, pulsed-dye laser, platelet-rich plasma) and microbiome-oriented approaches as promising new strategies. Established agents like ivermectin, azelaic acid, dapsone, sulfur preparations,

and metronidazole continued to show benefit, while tofacitinib (a JAK inhibitor) and oral ivermectin emerged as potential options for refractory cases. The authors concluded that a multimodal, mechanism-based approach addressing vascular, inflammatory, immunological, and microbial pathways offers the best outcomes. However, they stressed the need for large randomized controlled trials to validate long-term efficacy and safety.^{2,9}

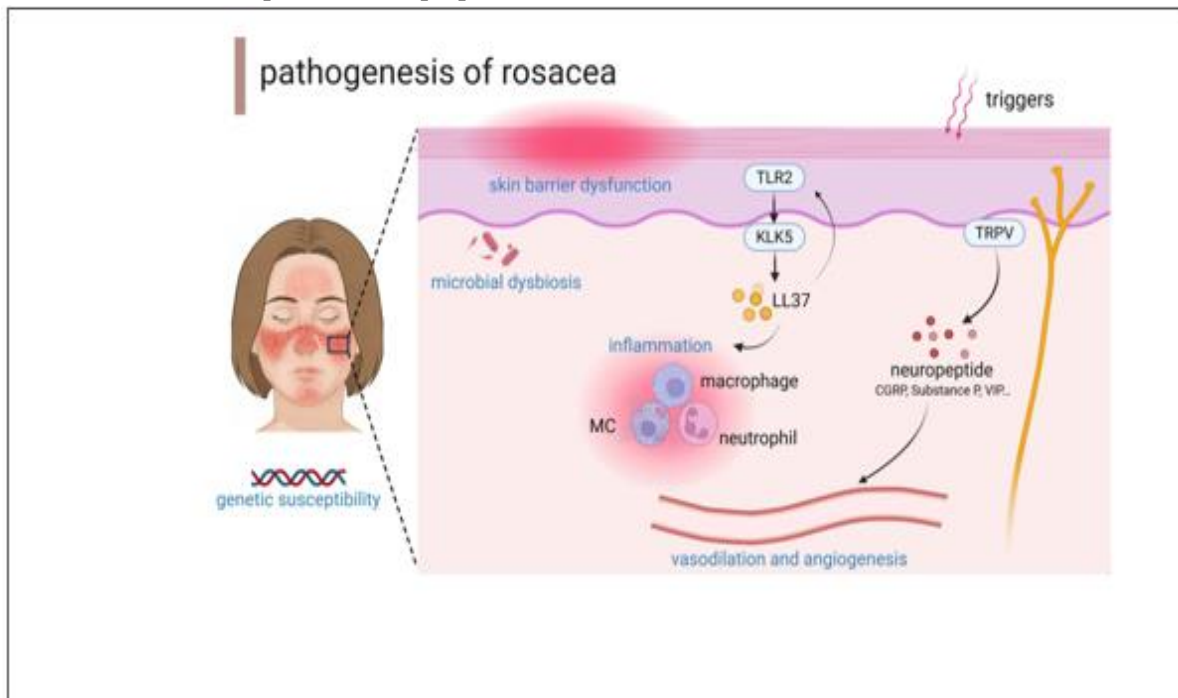


Figure 1: Pathogenesis of Rosacea

Rosacea is a common, chronic inflammatory skin condition that primarily affects the central face, causing persistent redness, visible blood vessels, and acne-like breakouts. It most frequently occurs in adults over 30, particularly those with fair skin and with global prevalence of around 5.1%. Causes psychosocial distress and reduced quality of life.

Genetic Factors: Twin studies show ~46% heritability. GWAS identified HLA variants and STAT1 mutations. Other implicated genes include LRRC4, SH3PXD2A, and SLC26A8. Vitamin D receptor polymorphisms also influence susceptibility.

Neurovascular Dysregulation: TRP channels (TRPV1, TRPV4) mediate flushing and burning. Neuropeptides (PACAP, CGRP, VIP) drive vasodilation and inflammation. Dysregulated amino acid metabolism (glutamate, aspartate) enhances neurovascular reactivity.

Immune Dysregulation:

- **Innate:** TLR2/LL-37/mTORC1 axis central to inflammation. Mast cells, macrophages, and neutrophils are overactive.
- **Adaptive:** Increased Th1/Th17 infiltration, elevated cytokines (IL-17A, IL-22, IFN- γ), angiogenesis, and impaired Treg function.

Skin Barrier Dysfunction: Increased transepidermal water loss, reduced claudins, altered lipid layers, and STAT3 overexpression contribute to barrier impairment.

Microbial Dysbiosis:

- **Skin:** High density of *Demodex folliculorum*, increased *Staphylococcus aureus* and *Corynebacterium kroppenstedtii*.

- Gut: Associations with *Helicobacter pylori* and altered gut microbiota; protective roles of *Lactobacilli* and *Bifidobacteria*.

Rosacea arises from interplay of genetics, neurovascular changes, immune dysregulation, barrier dysfunction, and microbiota imbalance. Advances in genomics and immunology are paving the way for personalized therapies.

SYMPTOMS AND SUBTYPES

Rosacea typically manifests in one or a combination of four primary subtypes:

- **Erythematotelangiectatic Rosacea:** Characterized by persistent facial flushing, redness, and visible broken blood vessels (telangiectasias). Skin may feel stinging or burning.
- **Papulopustular Rosacea:** Causes red bumps (papules) and pus-filled blemishes (pustules) similar to acne, alongside facial redness and swelling.
- **Phymatous Rosacea:** Involves skin thickening and excess tissue growth, most commonly affecting the nose (rhinophyma), causing it to appear bulbous or bumpy.
- **Ocular Rosacea:** Affects the eyes, causing dryness, burning, itching, bloodshot appearance, swollen eyelids, or photosensitivity.

Common Triggers: Symptoms often flare up in response to specific environmental or lifestyle factors that increase blood flow to the skin's surface:

- Sun exposure and wind
- Hot foods and hot beverages
- Spicy foods
- Alcohol (especially red wine)
- Temperature extremes (hot baths, heavy humidity, cold weather)
- Emotional stress or anxiety
- Strenuous exercise

- Harsh skincare products (e.g., exfoliants, alcohol-based toners, menthol)

TREATMENT AND MANAGEMENT

While there is no cure for rosacea, symptoms can be managed effectively through medical treatments and lifestyle adjustments:

- **Trigger Management:** Keep a journal to identify and avoid personal flare-up triggers.
- **Sun Protection:** Apply a broad-spectrum mineral sunscreen (zinc oxide or titanium dioxide) daily with SPF 30+, as UV exposure is the most common trigger.
- **Gentle Skincare Routine:** Use mild, non-comedogenic, fragrance-free cleansers and moisturizers. Avoid scrubbing or harsh exfoliants.
- **Topical Medications:** Prescription gels or creams (e.g., azelaic acid, metronidazole, ivermectin, or oxymetazoline) reduce inflammation and facial redness.
- **Oral Medications:** Low-dose oral antibiotics (such as doxycycline) are often prescribed for inflammatory bumps and ocular symptoms.
- **Laser and Light Therapy:** Intense Pulsed Light (IPL) or vascular lasers can reduce persistent facial redness and visible blood vessels.

1. Classification & Diagnostic Paradigm

Historically categorized into four fixed subtypes (erythematotelangiectatic, papulopustular, phymatous, and ocular), modern dermatologic consensus—including the Global Rosacea Consensus (ROSCO) panel—has shifted toward a **phenotype-directed framework**. This approach emphasizes assessing individual clinical presentations (e.g., persistent centrofacial erythema, inflammatory papules/pustules, telangiectasias, or ocular features) to allow for tailored, combination therapies.

2. Etiology & Pathogenesis

Rosacea is understood as a chronic, multifactorial inflammatory disease involving immune

dysregulation, neurovascular dysfunction, and skin barrier alterations.

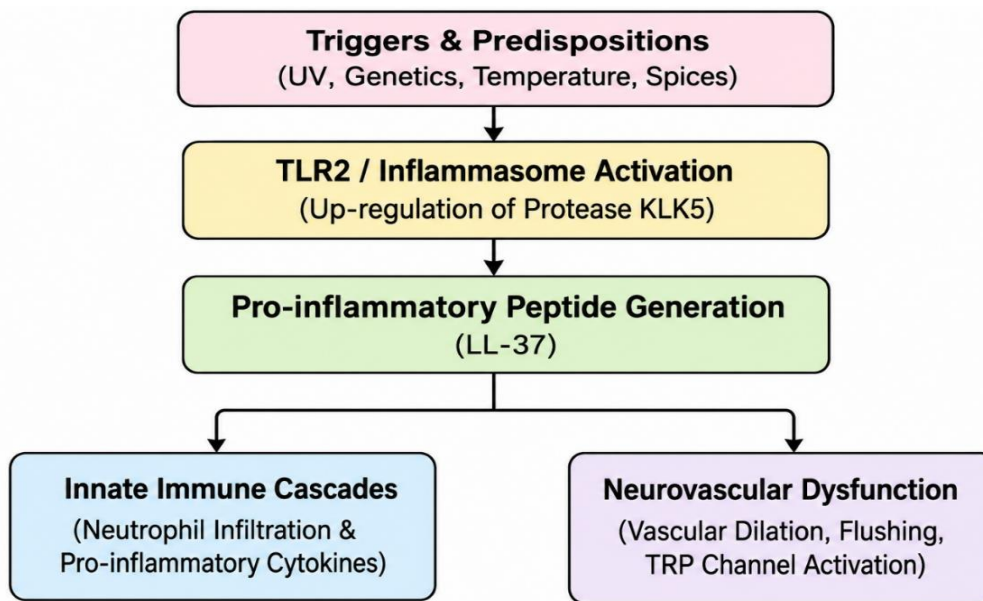


Figure 2: Pathophysiology of Rosacea

- **Innate Immunity & Cathelicidins:** A central mechanism involves activation of Toll-like receptor 2 (TLR2) and the NLRP3 inflammasome, leading to elevated serine protease (kallikrein-5, KLK5) activity. This process cleaves cathelicidin into the active pro-inflammatory peptide **LL-37**, driving vascular permeability and tissue inflammation.
- **Neurovascular Dysregulation:** Exogenous triggers (such as temperature extremes, capsaicin, and UV radiation) activate transient receptor potential (TRP) ion channels on cutaneous nerve endings, triggering the release of neuropeptides (such as CGRP and substance P) that induce flushing and erythema.
- **Barrier Dysfunction & Microbiome:** Disruption of the epidermal barrier—marked by elevated transepidermal water loss (TEWL)—interacts with microbial alterations, including *Demodex folliculorum* density and local microbiota dysbiosis, exacerbating cutaneous inflammation.

3. Therapeutic Modalities & Emerging Targets

Contemporary management combines skin barrier repair with targeted pharmacologic interventions:

- **Topical Anti-inflammatories & Vasoconstrictors:** First-line topicals include metronidazole, azelaic acid, and ivermectin for papulopustular manifestations, while topical alpha-adrenergic agonists (brimonidine, oxymetazoline) target transient or persistent erythema.
- **Systemic Interventions:** Sub-antimicrobial dose doxycycline (40 mg modified-release) provides anti-inflammatory benefit while reducing the risk of antibiotic resistance.
- **Procedural & Emerging Approaches:** Pulsed-dye laser (PDL) and Intense Pulsed Light (IPL) effectively target persistent vascular changes. Emerging research explores JAK/STAT pathway inhibitors and IL-17/TLR antagonists to address refractory barrier disruption and inflammatory cascades.

10,13

Nursing Process for Rosacea ^{8,13}

Assessment	Nursing Diagnosis	Goal/Expected Outcome	Nursing Interventions	Evaluation
Facial redness, flushing, visible small blood vessels, papules/pustules; patient reports burning or stinging	Impaired skin integrity related to inflammatory skin changes	Skin remains intact with reduced redness and inflammation	➤ Assess skin condition regularly ➤ Maintain gentle skin hygiene ➤ Advise lukewarm water and mild cleanser ➤ Avoid rubbing/scrubbing ➤ Administer prescribed topical medications	Skin integrity maintained; redness and lesions reduced
Patient reports burning, stinging, itching or discomfort of facial skin	Acute pain/discomfort related to skin inflammation	Patient reports reduced discomfort	➤ Assess intensity and triggers ➤ Apply cool compresses as appropriate ➤ Avoid hot water and irritating products ➤ Administer prescribed medication	Patient reports decreased burning/stinging
Patient expresses embarrassment or concern about facial appearance	Disturbed body image related to visible facial changes	Patient demonstrates improved acceptance of appearance	➤ Encourage expression of feelings ➤ Provide emotional support ➤ Explain that rosacea can be controlled with treatment ➤ Encourage appropriate cosmetic camouflage if desired	Patient verbalizes improved self-confidence
Patient may be unaware of triggers such as sunlight, heat, spicy food, alcohol, stress and hot beverages	Deficient knowledge regarding disease management	Patient demonstrates adequate knowledge of rosacea care	➤ Explain disease and treatment regimen ➤ Teach avoidance of individual triggers ➤ Advise use of broad-spectrum sunscreen ➤ Explain correct use of prescribed medications	Patient correctly explains self-care measures and triggers

Assessment	Nursing Diagnosis	Goal/Expected Outcome	Nursing Interventions	Evaluation
Recurrent flushing and inflammatory lesions; risk of worsening symptoms	Risk for exacerbation of symptoms related to exposure to known triggers	Patient remains free from frequent exacerbations	➤ Help identify personal triggers ➤ Maintain a symptom/trigger diary encourage sun protection ➤ Advise avoidance of extreme temperatures and irritating skincare products ➤ Reinforce follow-up	Frequency and severity of flare-ups are reduced

Important Health Education

- Use gentle, fragrance-free skincare products.
- Avoid scrubbing, exfoliation and harsh cosmetics.
- Use sun protection (SPF 30 or higher) regularly.
- Identify and avoid individual triggers such as sunlight, heat, spicy foods, hot drinks, alcohol and stress.
- Take prescribed topical/oral medicines regularly.
- Avoid applying steroid creams to the face unless specifically prescribed by the healthcare provider.
- Seek medical advice if there is eye redness, pain, blurred vision or persistent eye irritation, as rosacea can sometimes involve the eyes.^{8,13}

CONCLUSION

Nursing management is a multidimensional concept that encompasses both direct patient care and leadership responsibilities within healthcare organizations. Clinically, the nursing process (ADPIE) provides a systematic framework to assess, diagnose, plan, implement, and evaluate patient care, ensuring safe and effective outcomes. This structured approach empowers nurses to deliver evidence-based

interventions, monitor progress, and adapt care plans to meet evolving patient needs. On the other hand, administrative nursing management focuses on planning, organizing, staffing, directing, and controlling healthcare services. These functions are essential for maintaining quality standards, optimizing resource utilization, and fostering a supportive work environment. Together, clinical and administrative management ensure holistic patient care while strengthening nursing leadership. Effective nursing management not only improves patient outcomes but also enhances staff satisfaction, teamwork, and organizational efficiency. By integrating clinical expertise with managerial skills, nurses contribute to both bedside care and broader healthcare delivery. Ultimately, nursing management is the cornerstone of professional practice, bridging patient care and healthcare leadership to achieve excellence in service.¹⁶

REFERENCES

1. Fisher GW, Travers JB, Rohan CA. Rosacea pathogenesis and therapeutics: current treatments and a look at future targets. *Front Med.* 2023;10:1292722. doi:10.3389/fmed.2023.1292722
2. Geng RSQ, Bourkas AN, Mufti A, Sibbald RG. Rosacea: Pathogenesis and therapeutic correlates. *J Cutan Med Surg.* 2024;28(2):178–89. doi:10.1177/12034754241229365
3. Van Zuuren EJ. Rosacea. *N Engl J Med.* 2017;377:1754–64. doi:10.1056/NEJMcp1506630

4. Khalil S, Kurban M, Abbas O. Red scrotum syndrome: An update on clinicopathologic features, pathogenesis, diagnosis, and management. *J Am Acad Dermatol.* 2022;87:614–22. doi:10.1016/j.jaad.2020.05.113
5. Gether L, Overgaard LK, Egeberg A, Thyssen JP. Incidence and prevalence of rosacea: a systematic review and meta-analysis. *Br J Dermatol.* 2018;179:282–9. doi:10.1111/bjd.16481
6. Saurat JH, Halioua B, Baissac C, Cullell NP, Ben Hayoun Y, Aroman MS, et al. Epidemiology of acne and rosacea: A worldwide global study. *J Am Acad Dermatol.* 2024;90:1016–8. doi:10.1016/j.jaad.2023.12.038
7. Yang F, Zhang Q, Song D, Liu X, Wang L, Jiang X. A cross-sectional study on the relationship between rosacea severity and quality of life or psychological state. *Clin Cosmet Investig Dermatol.* 2022;15:2807–16. doi:10.2147/CCID.S390921
8. Halioua B, Cribier B, Frey M, Tan J. Feelings of stigmatization in patients with rosacea. *J Eur Acad Dermatol Venereol.* 2017;31:163–8. doi:10.1111/jdv.13748
9. Schaller M, Almeida LMC, Bewley A, Cribier B, Del Rosso J, Dlova NC, et al. Recommendations for rosacea diagnosis, classification and management: update from the global ROSacea COnsensus 2019 panel. *Br J Dermatol.* 2020;182:1269–76. doi:10.1111/bjd.18420
10. Clanner-Engelshofen BM, Bernhard D, Dargatz S, Flaig MJ, Gieler U, Kinberger M, et al. S2k guideline: rosacea. *J Dtsch Dermatol Ges.* 2022;20:1147–65. doi:10.1111/ddg.14849
11. Aldrich N, Gerstenblith M, Fu P, Tuttle MS, Varma P, Gotow E, et al. Genetic vs environmental factors that correlate with rosacea: A cohort-based survey of twins. *JAMA Dermatol.* 2015;151:1213–9. doi:10.1001/jamadermatol.2015.2230
12. Xiao W, Li J, Huang X, Zhu Q, Liu T, Xie H, et al. Mediation roles of neutrophils and high-density lipoprotein (HDL) on the relationship between HLA-DQB1 and rosacea. *Ann Med.* 2022;54:1530–7. doi:10.1080/07853890.2022.2077427
13. Chang ALS, Raber I, Xu J, Li R, Spitale R, Chen J, et al. Assessment of the genetic basis of rosacea by genome-wide association study. *J Invest Dermatol.* 2015;135:1548–55. doi:10.1038/jid.2015.53
14. Deng Z, Chen M, Zhao Z, Xiao W, Liu T, Peng Q, et al. Whole genome sequencing identifies genetic variants associated with neurogenic inflammation in rosacea. *Nat Commun.* 2023;14:3958. doi:10.1038/s41467-023-39761-2
15. Yamasaki K, Kanada K, Macleod DT, Borkowski AW, Morizane S, Nakatsuji T, et al. TLR2 expression is increased in rosacea and stimulates enhanced serine protease production by keratinocytes. *J Invest Dermatol.* 2011;131:688–97. doi:10.1038/jid.2010.351
16. Holmes AD, Steinhoff M. Integrative concepts of rosacea pathophysiology, clinical presentation and new therapeutics. *Exp Dermatol.* 2017;26:659–67. doi:10.1111/exd.13143

HOW TO CITE: Yerni Jyothi Kolli*, Sheeba Anitha Rani, Anusuya Devi, Blush Beyond The Skin: Healing Paths In Rosacea, *Int. J. Sci. R. Tech.*, 2026, 3 (9), 103-110. <https://doi.org/10.5281/zenodo.22303407>