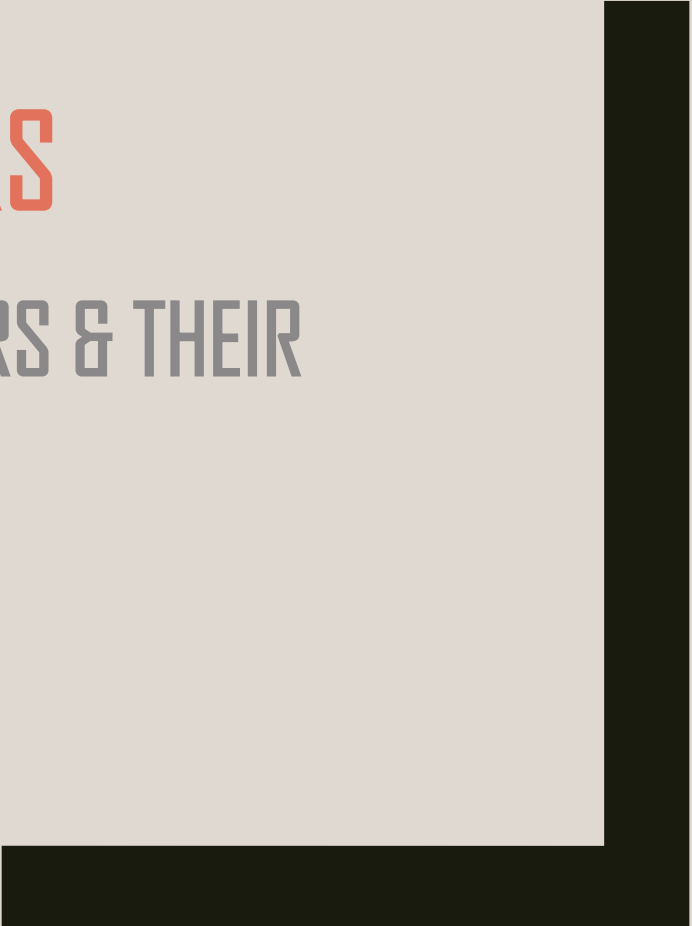




VITILIGO AND MIMICKERS

EXPLORING DEPIGMENTATION DISORDERS & THEIR DIAGNOSIS

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Distinguishing Vitiligo from Its Mimickers

Several skin conditions can **mimic vitiligo**, leading to potential **misdiagnosis**. Accurate differentiation is crucial for appropriate treatment.

Common Mimickers:

- **Tinea Versicolor:** Fungal infection with scaly patches
- **Post-Inflammatory Hypopigmentation:** Follows skin trauma or inflammation
- **Pityriasis Alba:** Well-defined hypopigmented patches, common in children
- **Idiopathic Guttate Hypomelanosis:** Small, white macules, typically in older adults
- **Progressive Macular Hypomelanosis:** Ill-defined hypopigmented macules, often in young adults
- **Hypopigmented Mycosis Fungoides:** Hypopigmented patches or plaques, more common in younger individuals with darker skin tones



Understanding Vitiligo & Its Clinical Relevance

What is Vitiligo?

A chronic skin condition characterized by the loss of melanocytes, leading to patchy depigmentation.

Relevance in Clinical Dermatology:

- Impacts 0.5–2% of the global population.
- Associated with psychological and social challenges for patients.
- Often confused with mimickers like Tinea Versicolor or Post-inflammatory Hypopigmentation.

Importance of Diagnosis and Management:

- Early recognition can prevent misdiagnosis.
- Tailored therapies improve quality of life.

Poll: How often do you encounter or diagnose vitiligo in your clinic?

Options:

1. Frequently (at least once a month)
2. Occasionally (a few cases a year)
3. Rarely (once every few years)
4. Never



Key Factors In the Development of Vitiligo

Autoimmune Factors: The body's immune system mistakenly attacks melanocytes, leading to depigmentation. This autoimmune process is often associated with other autoimmune conditions such as thyroid disease (Hashimoto's thyroiditis) and alopecia areata.

Genetic Predisposition: Family history plays a significant role, with genetic studies identifying specific polymorphisms in genes regulating the immune response, such as NLRP1 and PTPN22.

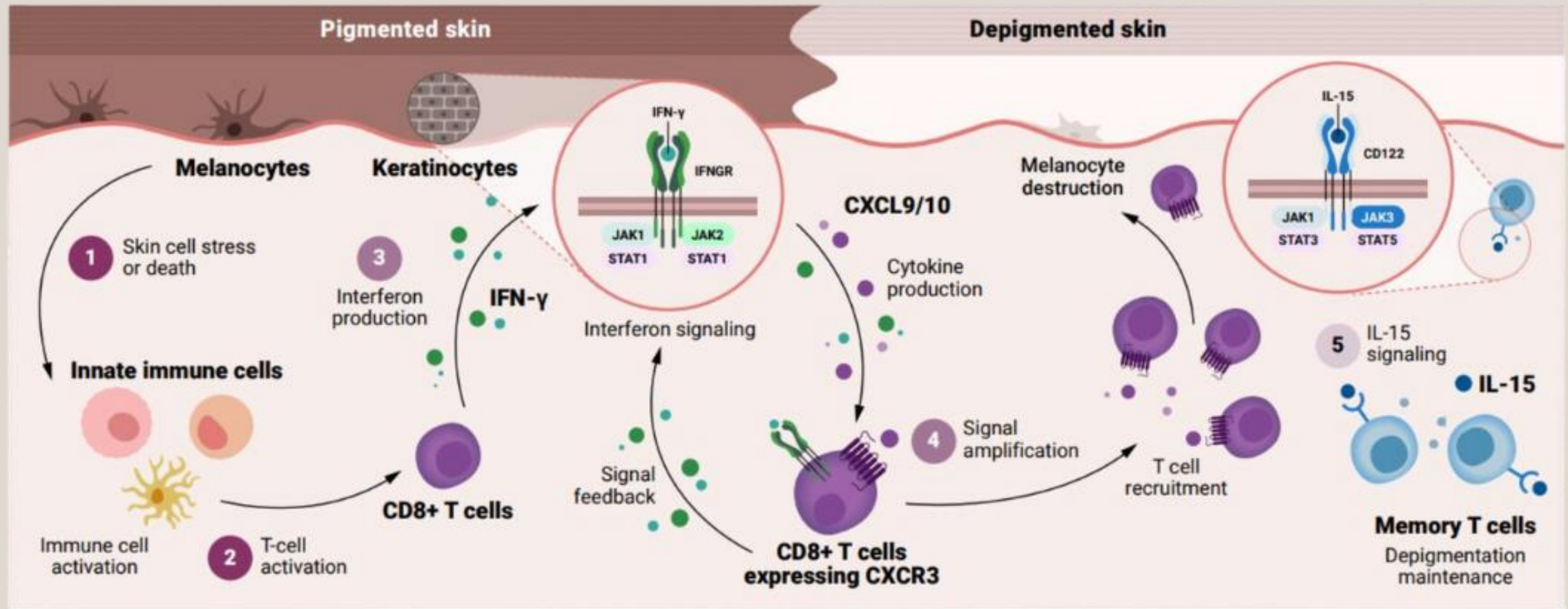
Environmental Triggers: Exposure to certain triggers, including stress, trauma, sunburn, and chemical irritants, can precipitate or exacerbate the onset of vitiligo.



INITIATION

PROGRESSION

PERSISTENCE



Mechanisms Behind Vitiligo

Melanocyte Destruction and Immune Dysregulation

Melanocyte Destruction:

- The progressive loss of **melanocytes** in affected areas leads to depigmentation.
- Linked to **oxidative stress** and intrinsic cellular **vulnerabilities** in melanocytes.

Immune Response:

- **Cytotoxic T-cells** target and destroy melanocytes.
- Autoimmune pathways involving interferon-gamma and CXCL10 contribute to disease progression.
- Persistent **inflammation** hinders repigmentation efforts.

Tools and Techniques for Diagnosing Vitiligo

Wood's Lamp Examination:

- Highlights depigmented areas by causing a bright white fluorescence under UV light.
- Useful for distinguishing vitiligo from mimickers like Tinea Versicolor or post-inflammatory hypopigmentation.

Biopsy:

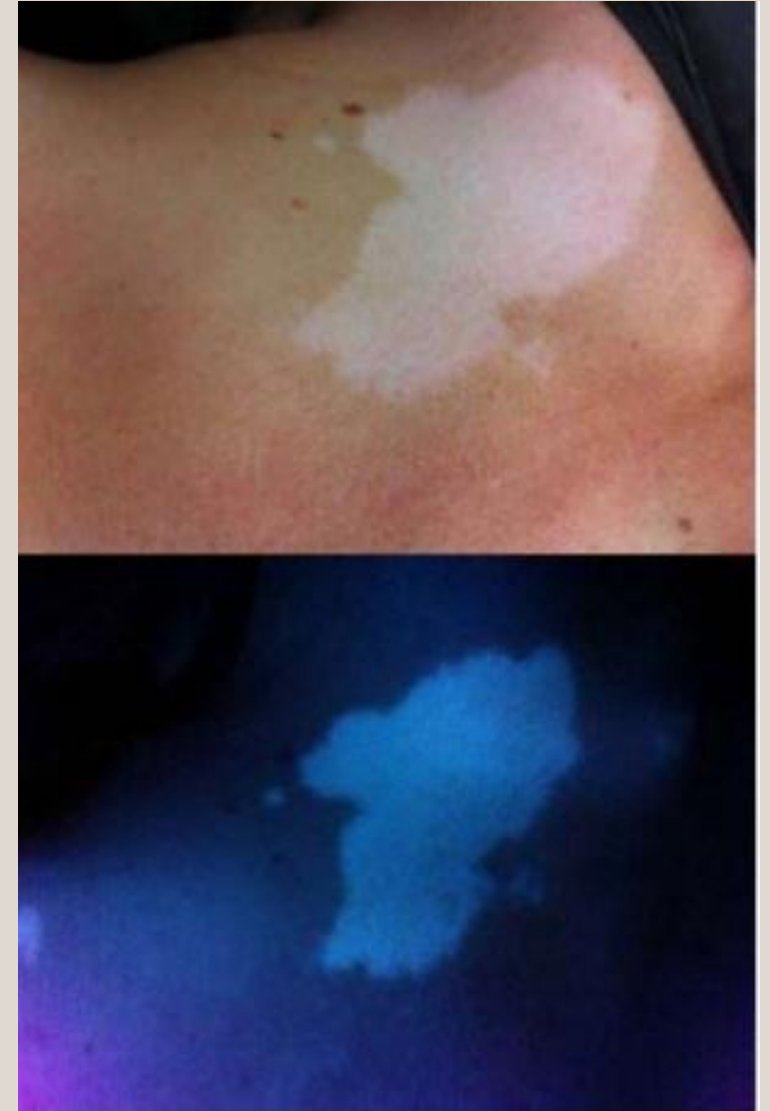
- Typically, unnecessary but may confirm the diagnosis in uncertain cases.
- Reveals the absence of melanocytes in affected skin and inflammatory markers.

Clinical Signs:

- Sharply demarcated white patches, often symmetrical.
- Frequently observed on the face, hands, and areas prone to friction or trauma.
- Look for associated leukotrichia (white hair in affected areas).

Wood's Lamp Examination

- ✓ Highlights depigmented patches, especially in lighter skin tones, often missed under normal light.
- ✓ UV light reveals bright blue-white fluorescence, contrasting with healthy skin.
- ✓ Fluorescence occurs due to the absence of melanin, exposing dermal collagen.
- ✓ Enhances lesion visibility, aiding early detection and accurate assessment.



Effective Strategies for Managing Vitiligo

Topical Steroids:

- First-line treatment for localized vitiligo.
- Reduces inflammation and promotes repigmentation.
- Common agents: Clobetasol, Betamethasone.
- **Caution:** Long-term use may lead to skin thinning or striae.

Calcineurin Inhibitors:

- Ideal for delicate areas like the face, neck, and intertriginous zones.
- Reduce local immune activity without causing skin atrophy
- Common agents: Tacrolimus (Protopic) and Pimecrolimus (Elidel).

Light Therapy:

- Narrowband UVB (NB-UVB) is the gold standard for widespread vitiligo.
- Helps stimulate melanocyte activity and suppress immune-mediated destruction.
- Typically requires 2–3 sessions per week for several months.

Effective Strategies for Managing Vitiligo

New and Emerging Treatments:

- **JAK Inhibitors:** Topical or oral treatments targeting Janus kinase pathways to reduce immune response.
- **Cellular Transplantation:** Melanocyte keratinocyte transplantation procedure (MKTP) for stable vitiligo.
- **Targeted Biologics:** IL-15 inhibitors are currently in clinical trials.



MIMICKERS OF VITILIGO

RECOGNIZING THE IMPOSTERS



Post-Inflammatory Hypopigmentation

Occurs when pigment production decreases after inflammation or injury.

Causes:

- Eczema
- Psoriasis
- Lichen Planus
- Burns (Trauma/Chemical)
- Infections (e.g., Tinea Versicolor)

Features:

- Pale patches/spots
- Gradual repigmentation

Management:

- Reassurance
- Treat inflammation
- Sun protection



Tinea Versicolor vs. Vitiligo

What is Tinea Versicolor?

- A superficial fungal infection caused by *Malassezia* species
- Presents as hypo- or hyperpigmented patches, often on the trunk and shoulders

Clinical Features:

- Patches may have slight scaling
- Common in humid or warm environments
- Can worsen with sweating

Management:

- Antifungal treatments (ketoconazole or selenium sulfide)
- Educate on recurrence prevention and differentiation from vitiligo



Features	Tinea Versicolor	Vitiligo
Pigmentation	Hypo- or hyper-	DE-pigmented
Scaling	Often present	Absent
Wood's Lamp	No change	Fluoresces
Treatment	Antifungals	Varies

Idiopathic Guttate Hypomelanosis vs. Vitiligo

What is Idiopathic Guttate Hypomelanosis (IGH)?

- Small, white, round or oval macules
- Commonly in older adults and sun-exposed areas like arms and legs

Clinical Features:

- Small, flat, depigmented spots (2–5 mm)
- Frequently occurs with age and prolonged sun exposure
- Asymptomatic, not progressive

Management:

- Cosmetic camouflage for aesthetic concerns
- Sun protection to prevent further contrast
- Educate on the benign nature of IGH



Features	Idiopathic Guttate Hypomelanosis (IGH)	Vitiligo
Size/Shape	Small, uniform macules	Larger, irregular patches
Progression	Stable	Often progresses
Cause	Aging, sun exposure	Autoimmune
Treatment	Cosmetic camouflage, sun protection	Varies

Progressive Macular Hypomelanosis vs. Vitiligo

Definition:

- Development of red-brown macules that gradually lighten to hypopigmented areas

Clinical Features:

- Seen on the back, especially along the trunk
- Lesions may merge to form larger hypopigmented patches
- Common in adolescents and young adults, particularly in tropical climates

Management:

- Topical Antibacterial Agents: Clindamycin or benzoyl peroxide
- Phototherapy: Narrow-band UVB or sunlight exposure
- Reassure patients that it's a benign and manageable condition



Features	Progressive Macular Hypomelanosis	Vitiligo
Lesion Evolution	Red-brown to hypopigmented	Direct depigmentation
Distribution	Primarily on the back	Widespread or localized
Wood's Lamp	May fluoresce orange-red	Blue-white
Treatment	Antibacterials, phototherapy	Varies

Pityriasis Alba vs. Vitiligo

What is Pityriasis Alba?

- Mild, scaly, hypopigmented patches,
- Commonly on face, neck, and upper arms
- Primarily affecting children and adolescents

Clinical Features:

- Round or oval, poorly defined patches with slight scaling
- Hypopigmented patches
- Follows mild eczema or dry skin

Management:

- Use of moisturizers to address dryness
- Mild topical steroids and sun protection
- Educate about the benign and self-limiting nature



Features	Pityriasis Alba	Vitiligo
Borders	Ill-defined	Sharply demarcated
Scaling	Mildly scaly	Absent
Progression	Resolves with time	May progress
Treatment	Moisturizers, mild topical steroids, sun protection	Varies

Hypopigmented Mycosis Fungoides

A Rare Presentation of T-Cell Lymphoma

What is Hypopigmented Mycosis Fungoides?

- Rare variant of cutaneous T-cell lymphoma (CTCL)
- Predominantly affects darker skin types
- More common in children/adolescents

Clinical Features:

- Hypopigmented, well or poorly defined
- Common on the trunk, buttocks, and extremities
- Often asymptomatic or mild itching

Diagnosis:

- **Biopsy:** Confirms the diagnosis with histopathologic examination showing atypical T-cell infiltrates
- **Immunohistochemistry:** Reveals loss of CD7 or other T-cell markers



Features	Hypopigmented Mycosis Fungoides	Vitiligo
Underlying Cause	Cutaneous T-cell lymphoma	Autoimmune
Pigmentation	Hypopigmentation	Depigmentation
Diagnosis	Biopsy essential	Often clinical
Prognosis	Variable, requires treatment	Generally benign

Vitiligo vs. Mimickers

Can you tell the difference?



Vitiligo

OR

Mimicker

Vitiligo vs. Mimickers

Can you tell the difference?



Vitiligo

OR

Mimicker

Vitiligo vs. Mimickers

Can you tell the difference?



Tinea Versicolor

Vitiligo

OR

Mimicker

Vitiligo vs. Mimickers

Can you tell the difference?



Hypopigmented
Mycosis Fungoides

Vitiligo

OR

Mimicker

Diagnostic Tips for Vitiligo vs. Mimickers

Importance of Accurate Diagnosis:

- Avoid **misdiagnosis** and **inappropriate** treatment
- Differentiate between vitiligo and conditions like post-inflammatory hypopigmentation, tinea versicolor, pityriasis alba, idiopathic guttate hypomelanosis, and hypopigmented mycosis fungoides

Clinical Tools:

- **History and Physical Exam:** Look for symmetry, progression, and other systemic findings
- **Wood's Lamp:** Critical tool for assessing depigmentation and fluorescence

Key Clinical Signs to Differentiate

Vitiligo:

- Sharply demarcated depigmented patches
- Symmetrical distribution
- Associated with autoimmune diseases.

Mimickers:

- **Post-inflammatory Hypopigmentation:** Ill-defined borders, history of inflammation or injury
- **Tinea Versicolor:** Fine scaling, hypo- or hyperpigmentation, fluoresces yellow-green under Wood's lamp
- **Pityriasis Alba:** Mild scaling, ill-defined borders, more common in children
- **Idiopathic Guttate Hypomelanosis:** Small, round, hypopigmented macules, often in sun-exposed areas
- **Hypopigmented Mycosis Fungoides:** Persistent hypopigmented plaques with biopsy showing atypical T-cells

Diagnostic Modalities: Tools and Techniques

Wood's Lamp Examination:

- **Vitiligo:** Bright blue-white fluorescence indicating depigmentation
- **Tinea Versicolor:** Yellow-green fluorescence due to fungal elements
- **Progressive Macular Hypomelanosis:** Orange-red fluorescence

Biopsy:

- **Vitiligo:** Absence of melanocytes in affected areas
- **Hypopigmented Mycosis Fungoides:** Atypical lymphocytes infiltrating the epidermis

Laboratory Workup (if needed):

- Rule out fungal infections with skin scrapings
- Autoimmune screening in suspected vitiligo cases

Primary Care in Early Diagnosis



Vitiligo is often first noticed by **patients** or their **families**.



Primary care providers play a **critical role** in identifying **early signs** (e.g., sharply demarcated white patches, often on the face, hands, or other exposed areas).



Differentiating vitiligo from mimickers like tinea versicolor, post-inflammatory hypopigmentation, and pityriasis alba prevents **misdiagnosis** and **delays** in referral.

Every Skin Tells A Story

As clinicians, it's our **duty** to uncover the right diagnosis, no matter how subtle or complex the presentation.

- ✓ **Accurate diagnosis is essential:** Impacts patient well-being, prevents mistreatment.
- ✓ **Recognize diversity:** Consider variations in skin of color.
- ✓ **Tools:** Wood's lamp, biopsies, thorough history.
- ✓ **Empower patients:** Education is key.
- ✓ **Advocate:** Stay informed, share knowledge, promote inclusion.
- ✓

Every patient deserves to feel **seen, heard, and treated**
with precision.

Let's work together to close the
diagnostic gaps
one case at a time.

