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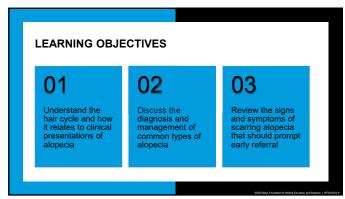
DISCLOSURE OF RELEVANT FINANCIAL RELATIONSHIP(S) WITH INELIGIBLE COMPANIES

Nothing to disclose

REFERENCES TO OFF-LABEL USAGE(S) OF PHARMACEUTICALS OR INSTRUMENTS

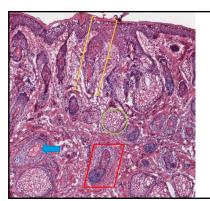
Discussing off label treatments for both scaring and non-scarring alopecia—all generic medications.

All relevant financial relationships have been mitigated.



BACKGROUND BASICS

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ALOPECIA

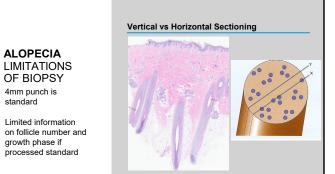
DISORDER OF HAIR FOLLICLES

A hair follicle is a tube-like structure that is continuous with the epidermis and associated with sebaceous glands

Hair is produced through successive keratinization in hair bulb Average growth is 2-6 years 100 hairs shed normally Metabolically inert

5 million hair follicles
Density varies by anatomic site
100,000 on the scalp
30,000 on the face
Remainder on the body

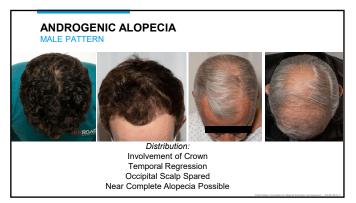




ALOPECIA LIMITATIONS OF BIOPSY Horizontal sectioning requires special processing protocol Necessary in many cases for proper diagnosis

NON-SCARRING ALOPECIA

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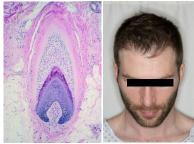


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ANDROGENIC ALOPECIA FEMALE PATTERN Distribution: Initially Hair Part Widening Then Progressive Involvement of Crown Frontal Hair Line Preserved Absence of Complete Alopecia

ANDROGENIC ALOPECIA

- · Hair diameter influenced by
- Anatomical location
 - Mesenchyme imprinting is the basis of hair transplantation
- Androgens
 - In men on the face, promotes vellus to terminal hairs ("virilization") yielding a beard
 - On scalp with permissive genetics, promotes terminal to vellus ("miniaturization") hairs producing alopecia



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ANDROGENIC ALOPECIA

PROLONG HAIR GROWTH

Minoxidil - mechanism unknown Topical

- 5% foam daily OTC
- · Irritant and contact dermatitis not uncommon

Oral

- Females: 1.25 mg to 2.5 daily
 Males: 2.5 mg to 5 mg daily
 Titrate higher after 12 wks based on response
- · LE edema possible in some

MINIMIZE ANDROGEN SENSITVITY

Finasteride – 5-alpha reductase inhibitor Oral

- 1 mg daily
- Sexual dysfunction rare
- Implications on PSA screening
- · Differential impact/import
 - Males : "Freeze effect"; more important
 - Females : less benefit given lower baseline androgen levels; postmenopausal severe only. Consider spironolactone 100 mg daily as alternative.

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A 63-year-old African American female with progressive scalp alopecia for over a decade. No improvement with Nizoral shampoo. Denies erythema or scaling.

Which diagnostic study would you obtain?

- A) Serum testosterone & DHEA-S
- B) VDRL and RPR
- C) Hair Trichogram
- D) Serum TIBC and Ferritin
- E) Scalp Biopsy



A) Serum testosterone and DHEA-S → Androgenic alopecia reflects ↑ sensitivity not ↑ levels; helpful if other signs of virilization.

B) VDRL and RPR → Syphilic alopecia is patchy & focal ("moth-eaten")

C) Hair Trichogram → Hair shaft disorders or to exclude Tinea Capitis with KOH prep

D) Serum TIBC and ferritin → history, extent and pattern not consistent with TE

E) Scalp biopsy → rule out scarring alopecia which will change treatment; suspect in any female patient of color w/ "extreme" or early onset male distribution androgenic pattern.

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CENTRAL CENTRIGUICAL CICATRIAL ALOPECIA

African Americans Females
Alopecia Begins at Vertex and Expands
Scarring but usually does not appear inflammatory



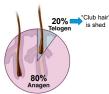


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TELOGEN EFFLUVIUM



Diagnosis is mainly historical "More hair on the pillow, with combing or with washing" Extensive or discrete alopecic patches are rare No involvement outside the scalp



Alternations to the normal ratio of anagen to telogen hairs produces alopecia

TELOGEN EFFLUVIUM

TRIGGERS

Febrile illness Chronic systemic illness Medication changes involving hormones Extreme dieting Pregnancy

WORKUP

Biopsy frequently not diagnostic Ferritin, Vitamin D, Zinc, TSH – less may be more

OUTCOMES

Self limited in three to six months

- Rare chronic variant in elderly

TREATMENT

Reassure

– Will not go bald

Minoxidil - short term

- 5% foam x 12 weeks 1.25 mg PO x 12 weeks

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43-year-old female presents with visible areas of hair loss first noticed 3 months ago. She also reports increased hair shedding, but her scalp is otherwise asymptomatic.

What is the most likely diagnosis?

- A)Telogen Effluvium
 B) Alopecia Areata
 C) Female Pattern Alopecia
 D) Mixed Non-Scarring Alopecia
- E) Scarring Alopecia



Mixed Non-Scarring Alopecia Subclinical/mild female pattern alopecia "unmasked" by acute (chronic) telogen effluvium

- A)Telogen Effluvium → inconsistent with patterned loss; alopecia extent
- B) Alopecia Areata → female pattern distribution excludes
- C) Accelerated Onset Female Pattern Alopecia → follicular minimization is gradual and slow (years)
- E) Scarring Alopecia → unlikely given an acute to subacute hair loss hx with no scale

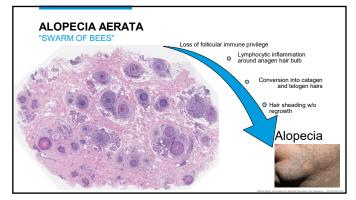
ALOPECIA AREATA

- The most prevalent autoimmune disorder
 - 2% of the global population Increasing incidence
- · Adults > Children
- Females > Males
- Asian > Black > Hispanic > White
- Darker hair types, sedating anesthesia
- Other associated conditions:
 Vitiligo
 Thyroid disease
 Anxiety/Depression



Ovoid smooth hairless patches on scalp

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ALOPECIA AREATA

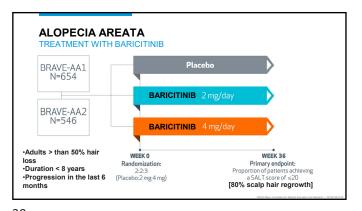
LOCALIZED, LIMITED - treatment responsive, remission common in 1 year

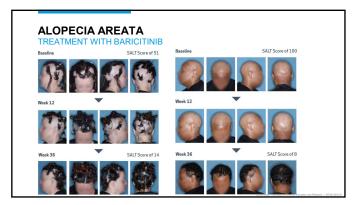
- IL Kenalog (5-10 mg/cc) q 4-6 weeks for up to 6 months
 Topical high potency corticosteroids daily
- Immunotherapy

EXTENSIVE, PROGRESSIVE - treatment resistant, full recovery rare

- Prednisone high response rate; rapid relapse with discontinuation
 Pulsed weekly prednisone + MTX, mycophenolate mofetil, azathioprine low efficacy (33% partial response); poor steroid sparing effect
- · JAK Kinase inhibitors new therapeutic class
 - · Remission of alopecia totalis in patient with rheumatoid arthritis treated with tofacitinib
 - Baricitinib selective JAK1 and JAK2 inhibitor; FDA approved for AA

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ALOPECIA AREATA

TREATMENT WITH BARICITINIB

> Br J Dermatol. 2023 Feb 10;188(2):218-227. doi: 10.1093/bjd/ljac059.

Integrated safety analysis of baricitinib in adults with severe alopecia areata from two randomized clinical trials



- N=1303; median 532 days of exposure
 - No deaths
- Drug related treated events vs placebo
 - Acne
 - Elevated CPK
- Immune related events vs placebo
 - Herpes Zoster (N=34)
 - MI (N=1), PE (N=1)

SCARRING ALOPECIA

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SCARRING ALOPECIA

- Inflammatory process destroys the hair follicle producing a scar

 Hair loss is permanent and irreversible

 Chronic slow progressive time course with spontaneous "remission"
- Difficult to treatment
- Early intervention has large impact on outcome

LYMPHOCYTIC GROUP

Discoid Lupus

Lichen Planopilaris

Frontal Fibrosing Alopecia

Central Centrifugal Cicatricle Alopecia

NEUTROPHILIC GROUP

Folliculitis Decalvans

Dissecting Cellulitis of the Scalp

Acne Keloidalis Nuchae

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DISCOID LUPUS

MOST COMMON SCARRING ALOPECIA

Skin limited (ANA-) and in association with SLE (ANA+)
Produces skin scarring and permanent alopecia in hair bearing areas





ACTIVE DLE Scaling red partially alopecic plaque

INACTIVE DLE Smooth scared nearly completely alopecic patch



- Vertex involvement usually
 Multi-focal with variable confluence
 Erythema, perifollicular scaling
 Pruritus, pain



FRONTAL FIBROSING ALOPECIA A CLINICAL VARIANT OF LICHEN PLANOPILARIS Post-menopausal females almost exclusively LLP changes limited to frontal hairline



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FOLLICULITIS DECALVANS

- White males
- · Hypersensitivity to Staph Aureus antigens
- Pustules with crusted and tufted folliculitis admixed into patches of scarring alopecia



OTHER NEUTROPHILIC SCARRING ALOPECIAS

Predominate in patients of color
Postulated follicular keratinization defect leading to occlusion and rupture





Dissecting Cellulitis of the Scalp - boggy nodules with alopecia and draining sinuses

Acne Keloidalis Nuchae – pustules with firm papules and plaques on occipital scalp

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CLUES TO EARLY SCARRING ALOPECIA

Erythema, scale >> overt alopecia in early disease



"Seborrheic dermatitis"-appearing process that fails to respond to topical steroids or OTC anti-dandruff shampoos

- 1/3 early scarring alopecia
- 2/3 scalp psoriasis

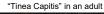
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CLUES TO EARLY SCARRING ALOPECIA

WHEN TO REFER

Diminished lateral eyebrows









Seen with many lymphocytic scarring alopecias

"Kerion"-like plaques common in neutrophilic scarring alopecias

THANK YOU SLUZEVICH.JASON@MAYO.EDU	
SSS two freedom to Mark South and Server I WESS OF	