

Peer-to-Peer Educational Toolkit

A compilation of key content from select presentations at the LiVDerm Deep Dive: Racial Disparities in Dermatology program.

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Structural Racism in Dermatology Workforce What is the Magnitude of the Problem?

Black dermatologists

3% of dermatologists but

12.8% of US population

4.2% of dermatologists but16.3% of US population

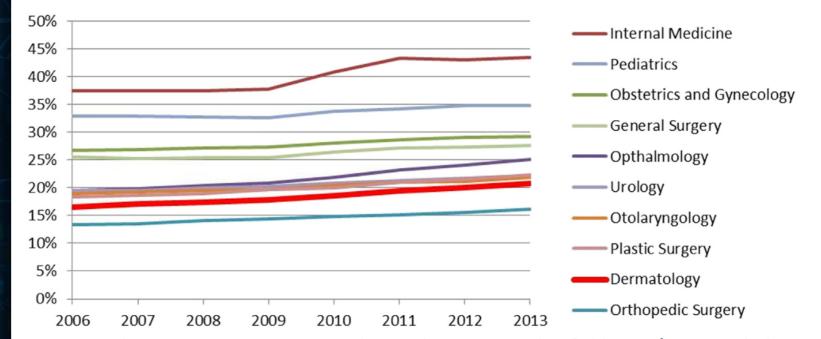


Fig 1. Total minority representation in dermatology versus other fields, 2006-2013, including Hispanics, African Americans, Asians, American Indians/Alaskan Natives, and Native Hawaiians/Pacific Islanders.

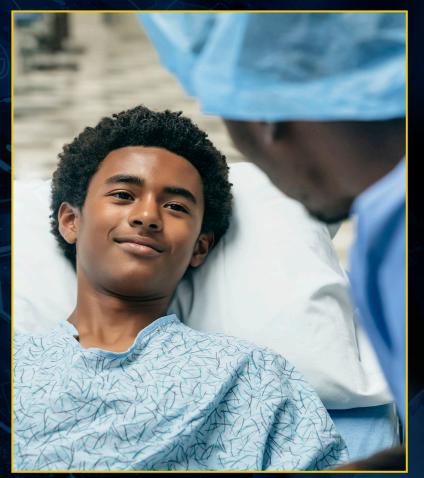
Dr. Susan Taylor



Pandya A, et al. Increasing racial and ethnic diversity in dermatology: A call to action. J Am Acad Dermatol March 2016;74(3):584-587.

Structural Racism in Dermatology Workforce Why Does Workforce Diversity Matter?

- Diversity in the medical workforce improves patient care and raceconcordant visits are longer and have higher ratings
- Minority physicians are more likely to
 - Care for patients of their own race or ethnicity
 - Practice in areas that are underserved
 - Care for patients who report poor health status
 - Have patients with Medicaid insurance or none







L.A. Cooper, D.L. Roter, R.L. Johnson, D.E. Ford, D.M. Steinwachs, N.R. Powe Patient-centered communication, ratings of care, and concordance of patient and physician race Ann Int Med, 139 (2003), pp. 907-915

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2 Letter

J AM ACAD DERMATOL 2020

Table II. Representation of dermatologic diseases in dark skin*

Diagnosis	Sauer's	Rook	Bolognia		Fitzpatrick 8th edition	Habif
				Kang		
Common dermatologic diseases						
Acne vulgaris	1/6	1/4	1/17	0/5	0/3	0/34
Alopecia	10/22	1/12	11/29	3/11	2/12	0/10
Atopic dermatitis	0/13	6/24	8/43	2/4	2/7	2/29
Pityriasis rosea	2/9	0/2	2/7	0/9	0/2	0/8
Psoriasis	2/11	2/28	3/41	3/18	1/14	0/16
Tinea infection	3/14	3/8	9/43	4/14	5/25	0/28
Syphilis, secondary	7/12	2/4	3/16	10/13	2/6	2/3
General medical diseases with racial or ethnic predisposition						
Sarcoidosis	1/3	1/13	9/13	0/2	1/5	0/0
Erythema dyschromicum perstans	0/0	0/0	0/2	0/0	0/0	0/0
Cutaneous amyloid	0/0	1/8	1/6	0/0	0/6	0/0

^{*}Values are reported as dark skin pictures/total number of pictures for each disease. See Table I for full citation details of the analyzed textbooks. Diseases with no dark skin representation are marked in bold.

Ademide Adelekun et al. J Am Acad Dermatol 2020 Apr 23;S0190

Dr. Andrew Alexis



LivDerm



Acne

Dr. Susan Taylor



Acne Vulgaris Treatment Considerations

- Early and effective treatment is key to reducing longterm sequelae.
- Waiting to get acne under control first before chasing down PIH is a disservice to patients.
- Acne can be multifactorial, consider therapies that effectively manage underlying diseases.
- Patients may be prone to PIH from irritation of topical medications.





Post-inflammatory Hyperpigmentation

- The most important and distinguishing sequela of acne in skin of color is post-inflammatory hyperpigmentation (PIH)
 - Results from an abnormal release or overproduction of melanin after cutaneous inflammation.
- Studies demonstrate PIH occurs more frequently and appears more noticeably in patients with darker skin hue (Fitzpatrick skin types III–VI).
- A psychological factor is persuading the patient that treatment is needed for acne as well as PIH.

Callender VD, Alexis AF, Daniels SR, et al. Racial differences in clinical characteristics, perceptions and behaviors, and psychosocial impact of adult female acne. J Clin Aesthet Dermatol. 2014;7:19-31.





SPF for Post-inflammatory Hyperpigmentation

- Mineral sunscreen ingredients form a physical barrier on the skin to help reflect UV rays
- Chemical sunscreen ingredients penetrate the top layers of the skin to absorb UV rays before they can damage the skin
- Non-comedogenic formulations can help PIH and reduce acne exacerbation

FDA Approved Ingredients

Mineral (physical) sunscreen:

- Zinc oxide
- Titanium oxide

Chemical sunscreen:

- Oxybenzone
- Avobenzone
- Octisalate
- Octocrylene
- Homosalate
- Octinoxate





Tolerability of Topical Agents

- It is important to consider the potential risk of irritant contact dermatitis, which may also result in PIH
- To address this concern in patients of color, it is advisable to start with low concentrations and more tolerable formulations
- Typically, the use of a cream vehicle is more tolerable than an alcohol-based gel, especially in dry/sensitive skin types
- Another approach is alternate-day dosing, usually at night which can then be titrated up as tolerated





Topical Retinoids

- Treat acne and PIH
- Approved retinoids: tretinoin, adapalene, tazarotene, trifarotene
 - Adapalene gel 0.1% is OTC for ~ \$12-15
- Modifies abnormal follicular keratinization, may be anti-inflammatory
- Dose and formulation dependent adverse effects
 - Irritation may occur initially
 - Erythema, dryness, peeling, burning, and cutaneous edema
- Slowly titrate upward beginning with application only on M-Th nights for 4 weeks, then increase to M-W-F nights for 4 weeks, then to M through F
- Apply a pea-size dot to the entire face (1 pea, not 4)





Antibiotics for Acne

- Oral antibiotics
 - Minocycline and doxycycline are the most prescribed antibiotics for acne
 - Not FDA approved for acne
 - Inhibit *P. acnes*
 - Anti-inflammatory: inhibits neutrophil chemotaxis, cytokines, and (MMP)-9
 - Dosage range of 100-200 mg/day
- Adverse effects:
 - GI intolerance
 - Drug eruption
 - Tetracyclines may have photosensitivity
 - Minocycline rare autoimmune disorders (lupus- like changes, hepatitis, arthritis, thyroiditis, or polyarteritis nodosa)

- Topical antibiotics
 - Erythromycin (solution, gel, pads)
 - Clindamycin (solution, gel, pledgets)
 - Slow to act, best in combination with BPO or retinoids
 - Resistance more likely with single agent



Salicylic Acid Peels may improve Acne and PIH

- Asian patients with mild-moderate acne, N=45
- Three study groups treated biweekly for 6 sessions:
 - 35% glycolic acid
 - 20% salicylic-10% mandelic acid
 - phytic acid
- Reduction in acne score at the end of 12 weeks in the three study groups was 70.55%, 74.14%, and 69.7%, respectively.
- A significant decline was observed in the postacne hyperpigmentation index in all the three study groups at the end of 12 weeks

Sarkar R, Ghunawat S, Garg VK. Comparative Study of 35% Glycolic Acid, 20% Salicylic-10% Mandelic Acid, and Phytic Acid Combination Peels in the Treatment of Active Acne and Postacne Pigmentation. J Cutan Aesthet Surg. 2019 Jul-Sep;12(3):158-163. doi: 10.4103/JCAS.JCAS_135_18. PMID: 31619887; PMCID: PMC6785964.







Psoriasis

Dr. Andrew Alexis



Clinical Differences

- Survey to 29 dermatologists opinion leaders in psoriasis – re: clinical characteristics of psoriasis in African Americans
- 66% (19/29) of respondents reported clinical differences in African Americans (including dyspigmentation, thicker plaques, less erythema)
- Clinical features: red-brown color





- NPF Survey 4,725 respondents (from 2004-2009)
- Greater psychosocial impact of psoriasis in African-Americans than in Caucasians
 - 72% said that psoriasis interfered with their capacity to enjoy life (vs. 54% of Caucasians)
- More severe disease in African-Americans
 - 23% of African American respondents had very severe psoriasis (vs. 8% of Caucasians)





Scalp Psoriasis

- Select treatment regimen that is compatible with patient's hair care practices
- Less frequent hair washing frequency in women of African descent (typically once per week to once every other week)
- Daily hair washing, especially with most prescription shampoos, is often associated with increased hair dryness and breakage; it is also very time consuming for most women of African descent due to common styling practices

- Non-topical therapies for severe cases:
 - Etanercept
 - Adalimumab
 - Secukinumab
 - Ixekizumab
 - Apremilast
 - Methotrexate
 - Cyclosporine
 - Acitretin*avoid in women of childbearing potential
 - Excimer laser





Considerations for African American Patients

- Less conspicuous erythema—may appear violaceous, gray, or hyperpigmented
- Postinflammatory hypo- or hyperpigmentation
- Consider potential clinical mimickers of psoriasis: lichen planus, cutaneous lupus erythematosus, sarcoidosis
- Scalp psoriasis in African Americans: Impact of hair texture, styling practices, and washing frequency on selection of topical therapy and severity
- Potential traditional/cultural therapies used before seeking dermatological consultation







Hidradenitis Suppurativa

Dr. Ginette Okoye



Hidradenitis Suppurativa Pathophysiology

- Chronic, debilitating inflammatory disease of the hair follicle
 - Apocrine sweat gland involved secondarily
- Post-pubertal onset of painful, recurrent abscesses/boils in the axillae, groin, and anogenital regions
- Tender subcutaneous nodules
 - can persist for days to weeks until they rupture, draining purulent malodorous material
 - after rupture, can remain open and draining for weeks to months
 - frequently reappear in the same location
- Recurrent flares can lead to sinus tract formation, hypertrophic or keloidal scarring and dermal contractures
- Chronic inflammation of HS can lead to lymphatic obstruction and squamous cell carcinoma







Hidradenitis Suppurativa Pathophysiology

Pathogenesis of HS is likely multifactorial

- Hyperkeratinization of the follicular infundibulum leading to follicular occlusion
 - The dilated follicle ruptures, spilling keratin and bacteria into the dermis

 inflammation
- Bacterial colonization/superinfection
- Genetic predisposition
- Obesity
- Smoking
- Hormonal influences: androgens
- Defects or alterations in host immunity

- Comorbidities:
 - Pyoderma gangrenosum
 - Inflammatory bowel disease
 - Cardiovascular disease
 - Musculoskeletal symptoms
 - Metabolic syndrome
 - Depression, Anxiety, Suicidality
 - Squamous cell carcinoma





Hurley Stage I Acute nodule / "abscess"



Hurley Stage II
Sinus tracts



Hurley Stage III

Sinus tracts and nodules involving an entire anatomic area; Hypertrophic scars



https://www.clinicaladvisor.com/hidradenitis-suppurativa/slideshow/277/



Disparities in HS

- Higher prevalence in African-Americans
 - Prevalence is 2-3 fold higher than among white patients
 - Black patients are more likely to have severe disease, and are more likely to be hospitalized for HS
- HS associated with low socioeconomic status²
 - Black patients with HS are more likely to have lower SES than other groups³
 - Patents with more severe disease are more likely to have a lower SES than patients with mild disease





Race, Socioeconomic Status, and HS

 Do patients with low SES have HS due to factors related to SES, e.g. access to care, differences in diet / nutrition > obesity, microbiome differences, etc.?

AND / OR...

• Do patients with HS have low SES due to the loss of educational opportunities and employment secondary to pain, drainage, and the medical & psychiatric comorbidities associated with HS?





Management of HS



Microbiome support?

Laser hair removal

HS Treatment Pyramid

Weight reduction: nutrition consult /
bariatric surgery
Smoking cessation

Prednisone

Surgery / Deroofing

Oral retinoids: isotretinoin, acitretin

Biologics +/- MTX

Hormonal therapy: Spironolactone, Finasteride / Metformin

Other antibiotics: metronidazole, cipro

Clindamycin 300 TID + Rifampin 300 TID

Doxycycline / Minocycline

Clindamycin 1% lotion/gel/solution Bleach baths

Chlorhexidine 4% wash Benzoyl peroxide 10% wash





Atopic Dermatitis

Dr. Andrew Alexis



Eczema Prevalence in the United States: Data from the 2003 National Survey of Children's Health

Tatyana E. Shaw¹, Gabriel P. Currie¹, Caroline W. Koudelka² and Eric L. Simpson¹

- National Survey of Children's Health (NSCH), a large population-based survey of >100,000 families representing all 50 states
- African American children are 1.7 times more likely to have AD than their white counterparts, even when adjusting for household income, parental education level, metropolitan versus rural environment, and health insurance coverage status

Shaw TE et al. J Invest Dermatol. 2011 Jan;131(1):67





Genetic Differences

- FLG mutations have a significant association with atopic dermatitis in European populations, however populations differences in prevalence have been observed.
- FLG mutations are found in up to:
 - 50% of all European patients with AD
 - 27% of Asian patients with AD

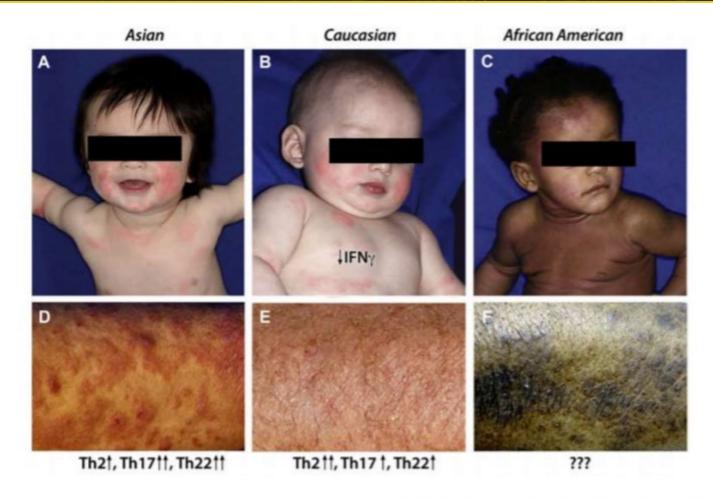
Korean, Japanese, Chinese, Singaporean, and Taiwanese populations all have specific FLG null mutations unique to their ethnic group, and they rarely exhibit the mutations commonly observed in Caucasians

 A study in 370 African Americans and 433 whites demonstrated that only 5.8% of blacks (compared to 27.5% of whites) had at least one of the four most common FLG null mutations





Immunophenotypic Differences



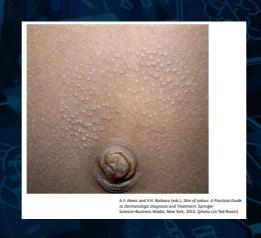
Leung DY. J Allergy Clin Immunol. 2015 Nov;136(5):1265



Clinical Differences



Erythema less visible – masked by pigment



Follicular accentuation



Lichenification



Dyspigmentation
Pruritus
— more severe?





Approach to Treatment

- Avoid under-treatment to prevent long-term sequelae (e.g. dyspigmentation)
- Aggressively target inflammation
- Effectively manage pruritus (risk of scratch induced dyspigmentation)
- Minimize duration of potent topical corticosteroids risk of hypopigmentation (among other side effects)
- Barrier protection liberal use of moisturizers and/or barrier repair "devices"







Pediatric Atopic Dermatitis

Dr. Candrice Heath



Disparities in Atopic Dermatitis

- United States:
 - 25% of children
 - Onset most commonly between 3-6 months old
 - 60% have symptoms by age 5 years
- African American children were more likely to be exposed
 - Lower household income
 - Lower parental education attainment
 - Lack of home ownership
 - Child living between to addresses
 - Exposure to smoking

- Higher prevalence in skin of color
- More treatment resistant disease in SOC
- Non-Hispanic Black children and Hispanic children had greater odds of persistent AD than non-Hispanic white children
- Atopic dermatitis prevalence and persistence were <u>highest</u> in US urban children who were **female or Black**.
- Urban children with persistent AD were more likely to have poor QOL and asthma.



Communication with Parents/Caregivers

Parents bring their frustrations with hyperpigmentation to the pediatric visit!

Verbalize:

- 1. You see the pigmentation change
- 2. Explain that controlling the AD improves PIH
- 3. Remaining pigmentation can be treated





Clinical Differences

 Papular and Follicular Eczema



 'close your eyes' technique to determine flare

Erythema may be barely / not perceptible







Pigmentary Disorders

Dr. Seemal Desai



Post-Inflammatory Hyperpigmentation:Treatment Options

- First and foremost:
 - Treat any underlying dermatoses and stress the importance of sun protection
- Topical retinoids
- Azelaic acid
- Hydroquinone
- Chemical Peels
- Cosmeceuticals
- Reassurance and Time





Melasma: Treatment Options

- Topical retinoids & combination therapy
- Azelaic acid
- Hydroquinone
- Chemical Peels
- Cosmeceuticals
- Lasers
- Dermabrasion
- Reassurance and Time



Drug Safety

Hydroquinone

- Hydroquinone remains the goldstandard, and non-HQ based therapies remain second line and/or adjunctive
- No substantial evidence to prove its carcinogenicity
- Take the time to discuss long term use with each of your patients!

Azelaic Acid

- Dicarboxylic acid that has selective cytotoxic effects
- Inhibits tyrosinase and mitochondrial respiratory enzymes
- Minimal side effects of erythema, pruritus, and mild burning



Drug Safety

Tranexamic Acid

- Contraindications:
 - Current/past DVT/PE/clotting disorder/anti-coagulant medications
 - Pregnancy/breastfeeding
 - Smoking
 - Renal/cardiac/pulmonary disease
- Dosing:
 - Most effective dosing seems to be at 250-500 mg daily
 - Topical, oral, and intradermal forms

Glutathione

- Potent anti-oxidant indirect inactivation of tyrosinase
- Assists in converting eumelanin to phaeomelanin
- Typically used oral
- Also being given IV in Asia
- Still controversial due to bioavailability, but some promising results in studies





Takeaways for Melasma Treatment

- Topical Therapy remains the first line treatment of melasma
- Hydroquinone remains the gold standard
- Second line topicals, though off-label, should be considered
- Tranexamic acid is an exciting new step in melasma treatment





Chemical Peel Cocktails

- Jessner's + TCA 20% for full face acne scars, especially when not deep boxcar or ice pick
 - Can do in skin of color
 - One layer after another
- Salicylic acid 30% and 10-20% Mandelic for Acne Vulgaris
 - Works great for acne and also rejuvenation
 - Synergy between alpha and beta hydroxy
- TCA 10% immediately after microneedling for dark circles
 - Procedure only lasts about 5 minutes
 - Depth of 0.5mm → SHALLOW





Vitiligo: Treatment Options

- Topicals including steroids, vitamin D analogues, calcineurin inhibitors
- Depigmentation
- Systemic treatment
 - JAK inhibitors (ruxolitinib 1.5% cream)
- Phototherapy
- Surgical Treatment
- Psychological therapy
- If treatments fail

 Analyze patient's desires!





Types of Vitiligo

- Active/Unstable Vitiligo
 - Depigmentation spreading more than 2% BSA in one month
- Chronic Vitiligo
 - Depigmentation present for at least 1 year with no h/o spontaneous repigmentation
- Refractory Vitiligo





Depigmentation Options

- Traditional:
 - Monobenzyl Ether of Hydroquinone (MBEH)
 - Monomethyl Ether of Hydroquinone/4-Methoxyphenol
 - 88% Phenol
 - Cryotherapy
 - Laser treatment

 Q-Switched alexandrite & Q-Switched Ruby
- Newer:
 - Imatinib
 - Imiquimod
 - Diphencyprone





Racial Disparities in Dermatology

Melanoma

Dr. Valerie Harvey



Stage-specific worsening of racial/ethnic disparity

Table II. Unadjusted hazard ratio and 95% confidence intervals for melanoma-specific survival of different racial/ethnicity groups compared with non-Hispanic whites in the Surveillance, Epidemiology, and End Results cohort and across 3 diagnostic time periods for different stages for cutaneous melanoma

	Race	<2000 HR (95% CI) for MSS	2000-2009		≥2010	
			HR (95% CI) for MSS	P for interaction	HR (95% CI) for MSS	P for interaction
Localized stage*				•		
	Hispanic	0.93 (0.79-1.09)	1.24 (1.08-1.43)	.007	1.35 (1.06-1.72)	.012
	NHB	1.62 (1.22-2.15)	2.26 (1.70-3.02)	.10	4.29 (2.84-6.47)	<.001
	NHAPI	1.18 (0.89-1.57)	1.47 (1.10-1.96)	.29	2.21 (1.42-3.43)	.020
	NHAIAN	0.82 (0.44-1.53)	1.02 (0.58-1.80)	.61	1.98 (0.94-4.15)	.076
Regional stage*						
	Hispanic	0.98 (0.81-1.18)	1.24 (1.10-1.40)	.042	1.46 (1.22-1.75)	.003
	NHB	1.72 (1.30-2.27)	1.48 (1.18-1.84)	.41	1.80 (1.28-2.52)	.84
	NHAPI	1.20 (0.90-1.59)	1.31 (1.06-1.62)	.61	1.24 (0.82-1.88)	.88
	NHAIAN	2.25 (1.24-4.07)	1.15 (0.72-1.83)	.080	2.72 (1.57-4.69)	.65
Distant stage*						
_	Hispanic	0.80 (0.64-1.00)	1.01 (0.87-1.17)	.088	1.14 (0.96-1.35)	.014
	NHB	1.18 (0.89-1.56)	0.78 (0.60-1.00)	.032	0.96 (0.70-1.32)	.34
	NHAPI	1.19 (0.87-1.63)	1.02 (0.80-1.30)	.45	1.69 (1.28-2.25)	.10
	NHAIAN	1.51 (0.72-3.16)	0.73 (0.35-1.54)	.18	0.82 (0.39-1.71)	.25
All stages*						
-	Hispanic	1.13 (1.02-1.25)	1.61 (1.49-1.73)	<.001	1.96 (1.76-2.17)	<.001
	NHB	2.32 (1.99-2.72)	2.88 (2.51-3.31)	.044	3.85 (3.16-4.68)	<.001
	NHAPI	1.77 (1.51-2.07)	2.23 (1.95-2.55)	.029	2.76 (2.27-3.36)	.001
	NHAIAN	1.30 (0.91-1.86)	1.34 (0.99-1.81)	.89	2.09 (1.45-3.01)	.067

Journal of the American Academy of Dermatology DOI: (10.1016/j.jaad.2020.08.097)





Differences in clinical presentation

Gender Majority of NHB and Hispanic patients

were female

Anatomic location of primary siteLower limb or hip most common primary

site for Hispanic, NHB, NHAPI

Histological subtype NHB have higher percentage of ALM

compared to other groups

Mucosal involvement Higher percentage of minorities with

mucosal melanoma

https://doi.org/10.1016/j.det.2019.05.009





Social determinants of melanoma disparities

- Disparities are symptoms of underlying social and economic determinants.
- Social determinants of Health (SDOH)
 - Five domains:
 - Economic stability (employment, income)
 - Education and access (higher education, literacy, language)
 - Health care access and quality (health coverage)
 - Built environment (housing, transportation, safety)
 - Social and community context (social support systems)











Racial Disparities in Dermatology

Cutaneous T-Cell Lymphoma

Dr. Ginette Okoye



Types of Cutaneous T-Cell Lymphomas

Indolent clinical behavior

- 1) Mycosis fungoides (MF)*
- 2) Mycosis fungoides variants
 - Folliculotropic MF
 - Pagetoid reticulosis
 - Granulomatous slack skin
- 3) Primary cutaneous CD30-positive lymphoproliferative disorders
 - Primary cutaneous anaplastic large cell lymphoma (C-ALCL)
 - Lymphomatoid papulosis (LyP)
- 4) Subcutaneous panniculitis-like T-cell lymphoma (SPTCL)
- 5) Primary cutaneous CD4-positive small/medium pleomorphic T-cell lymphoma

Aggressive clinical behavior

- •1) Sézary syndrome (SS)
- •2) Adult T-cell leukemia/lymphoma (ATLL)
- •3) Extranodal NK/T-cell lymphoma, nasal type
- •4) Primary cutaneous CD8-positive aggressive epidermotropic cytotoxic T-cell lymphoma
- •5) Primary cutaneous gamma/delta T-cell lymphoma (PCGD-TCL)
- •6) Primary cutaneous peripheral T-cell lymphoma (PTCL), unspecified





- Mycosis Fungoides in African Americans:
 - Higher incidence
 - Presents at a younger age
 - Diagnosed at later disease stages
 - Higher mortality rate
 - African-American women diagnosed before age 40 have a poor prognosis
- "Incidence of MF is higher among blacks (9.0 x 10⁻⁶) than among whites (6.1 x10⁻⁶) and was higher among men than among women."
- "The male-female IRR was lowest among blacks (1.5) and highest among the other racial groups (2.7)."
- Black patients with MF more likely to present with more advanced disease
 - i.e. T3 T4 (OR 1.72)
 - T3=tumors / T4=erythroderma
- Black, Asian/Pacific Islander, and Native American patients with MF more likely to present at a younger age
 - Mean age at diagnosis in white patients → 59.2 y.o.
 - A/PI \rightarrow 51.3
 - Black → 51.5
 - Native American → 53.8

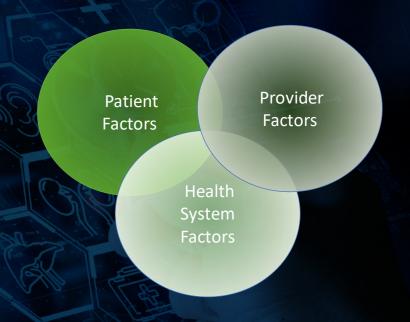
Wilson LD, Hinds GA, Yu JB. Clin Lymphoma Myeloma Leuk. 2012 Oct;12(5):291-6.





Health Disparities in CTCL

- Patient and Provider education about clinical presentation of MF in skin of color
- Access to dermatologists
 - Depends on insurance type
 - Expensive specialist co-pays
 - Too few dermatologists, varies by geography
- Cost/Availability of MF therapy
 - Expensive phototherapy co-payments
 - Access to phototherapy, extra-corporeal photopheresis (ECP), total skin electron beam radiation therapy
 - Missing work for treatment, e.g. TIW phototherapy
- Bias, attitudes and beliefs of providers





Clinical Presentation in Skin of Color

- Pigmentary changes more common/easier to appreciate than erythema
 - Hyperpigmentation, Hypopigmentation, Depigmentation, Polymorphic pigmentation, Lichenification
- Easily mistaken for other dermatoses:
 - Atopic dermatitis, psoriasis, lichen planus pigmentosus, inflammatory vitiligo, tinea versicolor
- Erythroderma more difficult to appreciate
 - May present as diffuse lichenification or diffuse skin darkening
- Facial lesions may be more common



Photo courtesy Peter Heald, MD





Treatment of Mycosis Fungoides

Management of mycosis fungoides depends on stage.

<u>Limited Stage (i.e. IA – IIA)</u>

Skin directed therapies:

- Potent topical corticosteroids (e.g. clobetasol)
- Phototherapy: Narrowband UVB, PUVA
- Topical bexarotene, topical nitrogen mustard
- Radiation therapy (*TSEB: total skin electron beam)
- <u>Advanced stage (IIB IV)</u>

Systemic agents:

- IFNa, oral bexarotene
- Extracorporeal photopheresis (*ECP)
- vorinostat, denileukin diftitox, histone deacetylase inhibitors, antifolate analogs, alemtuzumab







Racial Disparities in Dermatology

Non-Scarring Hair Disorders

Dr. Amy McMichael



Top Diagnoses in African American Patient Visits to Dermatologists

Diagnosis	ICD-9 Code	No. of Visits	% of Visits
Acne	706.1	5,720,000	22.1%
Unspec. dermatitis	692.9	3,640,000	14.0%
Seb dermatitis	690.10	1,990,000	7.7%
Atopic derm	691.8	1,590,000	6.1%
Dyschromia	709.0	1,290,000	5.0%
Psoriasis	696.1	950,000	3.6%
Alopecia	704.00	920,000	3.6%
Keloid scar	701.4	830,000	3.2%
Viral warts	078.1	780,000	3.0%
Sebaceous cyst	706.2	780,000	3.0%

Davis SA, et al. J Drugs Dermatol 2012



Damaging hair care practices

- Chemical straightening agents remove the mono-molecular layer of fatty acids covalently bound to the cuticle, including 18-methyl eicosanoic acid, which is important to prevent penetration of water into the hair shaft.
 - Most likely to cause catastrophic damage to the hair
- Hair dye less likely to lead to catastrophic hair breakage in one use but damage is more due to frequency of use leading to cumulative damage
 - Bleach > Permanent colorants > Demi-permanent > Semi-permanent > Temporary
- Thermal straightening with or without chemical process causes hair shaft weakening
 - Flat irons and curling irons can cause much more damage than blow dryers

Khalil EN (1986). Cosmet Toil 101:51-58.

Swee W, Klontz KC, Lambert LA (2000). Arch Dermatol Sep;136(9):1104-8.





Approach to Hair Breakage

- Check underlying abnormalities (Iron levels, thyroid, nutrition, etc)
- Give the hair a rest!
 - Consider stopping chemical relaxer, color, or relaxer for 6-12 months
 - Place a hair weave that is not tight and will allow hair care
 - Loose braids or wig
 - Natural hair but do not straighten with heat
- Serial trimming of hair (every 6-8 weeks)
- Use heat protectant products on the hair before styling
- Layering moisturizing regimen
 - Start with moisturizing shampoo and conditioner (should state for dry, damaged hair)
 - Next apply a leave-in conditioner with coating agents to wet hair (dimethicone-coating agents)
 - Add a leave-in conditioner (oils) to dry hair (after washing weekly and then as needed daily)
- Discuss the long wait for improvement





Olls Can Be Bad for hair and scalp

Coconut oil	 Effective hair lubricant and moisture sealant Comedogenic
Jojoba Oil	 Works well to lightly coat the hair shaft and lubricate hair Similar properties to sebum in lubricating hair shafts Can induce contact dermatitis
Argan Oil	 Can be used to protect the integrity of the hair fiber during harsh chemical processes such as hair coloring. It also gives shine and lubrication to the hair shafts. The allergenicity of argan oil is rare, but reports of it do exist.
Castor Oil	 Effective at lubricating and adding shine to hair shafts, as well as softening the hair. Known skin irritant Has been documented to cause acute hair felting in one case report
Olive Oil	 Effective at softening the hair, and providing lubrication and shine. Known skin irritant that has been .
Tea Tree Oil	 Tea tree oil is effective at soothing the scalp and improving dandruff, but given its irritating properties, it is best avoided
Shea Butter	 Good emollient Can safely be used to lock in moisture in hair shaft Should not be used for wet combing as can cause damage to the hair shaft

^{1.} Rele AS, Mohile RB. J Cosmet Sci. 2003 Mar-Apr;54(2):175-92

[.] Uwakwe L, McMichael, The Dermatologist, 2018



Seborrheic dermatitis treatment

- Discuss frequency of shampoo
- Make a contract with patient to increase washing to minimum of every 2 weeks
- Use products for "textured hair" or with "moisturizing" in the title, those developed for hair in African American patients
 - Zinc pyrithione #1
 - Selenium sulfide
 - Tar
 - Rarely use ketoconazole in African American women
- Apply medicated shampoo to the scalp only to minimize dryness of hair shafts
- Since conditioners can wash off the active ingredients in therapeutic shampoos, anti-dandruff conditioners are recommended
- Stop home use of pomades/oils to scalp





Seborrheic dermatitis treatment

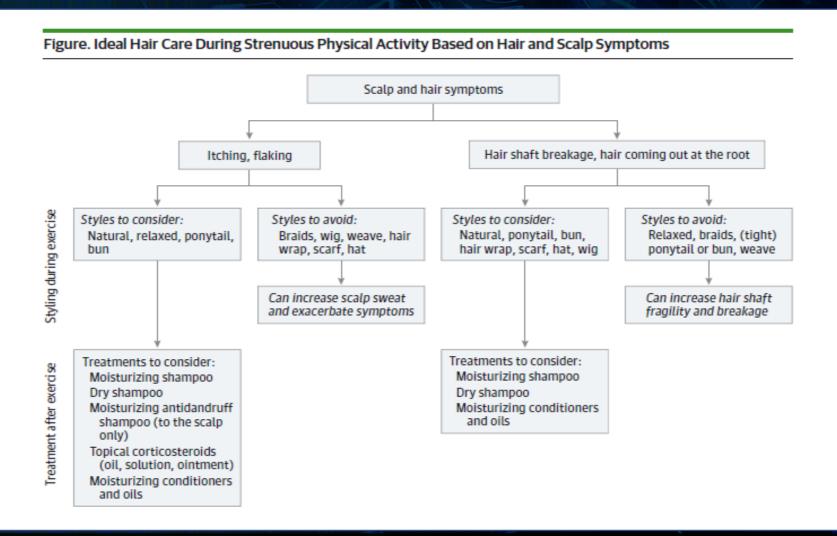
- Potent or ultrapotent topical steroids to scalp 3-4 times/week
- Mid-potency topical steroid oil weekly or more
- For face:
 - Low potency topical steroids, tapering to off
 - Topical calcineurin inhibitors
 - Anti-yeast topicals, like ketoconazole or econazole





Algorithm for managing breakage and seborrheic dermatitis

ahn c, et al Jama Derm 2016





Traction alopecia

- Prevalence estimated at 1% in London population and 37% in S. African population
- Observed to be much higher prevalence in women and girls of African descent
- High risk in those using chemical relaxer and tight braiding
- More than 80% with traction alopecia have experienced pain with hair styles
- Mechanism thought to be mechanical tension on the hair shaft

Khumalo NP et al. BJD 2007 Wright DR et al. JAAD 2011 Child FJ et al BJD 1999 Khumalo NP et al Arch Derm 2006





Treatment for traction alopecia

- Decrease friction and traction behaviors to the area
- Anti-inflammatory treatments
 - Mid-potency topical steroids 3-4 times per week
 - Intralesional triamcinolone 5 mg/cc to the affected areas for 2-3 cycles
- Topical minoxidil 5% daily
- Surgical correction
- Follow improvement with photos
- Discuss the fragility of regrowth



Uwakwe L, DeSouza B, Garza-Tovar, McMichael A. JDD. 19(2):128-130







Racial Disparities in Dermatology

Scarring Hair Disorders

Dr. Andrew Alexis



Unmet Needs in Hair Disorders for SOC Patients

- Delays in diagnosis

 Long-term or permanent sequelae
- Limited coverage in educational materials

 Educational gaps
- Limited research into new therapies

 Limited treatment options





Central Centrifugal Cicatricial Alopecia

- Hot comb alopecia, Follicular degeneration syndrome
- Begins on crown
- Secondary to long-term chemical processing, thermal straightening, tension
- Primary Cicatricial Alopecia lymphocytic
- Genetic predisposition





Research Findings: CCCA and SOC

- Self report of traction associated hair styles was significantly higher in those with CCCA (ie. Grade ≥ 3 on photographic scale) vs. those without
- Mutations in PADI3 (an enzyme that post-translationally modifies other proteins important for proper hair-shaft formation) were associated with CCCA
- African-American women with CCCA had almost a five-fold increased odds of having uterine leiomyomas compared to controls

Kyei A et al. Arch Dermatol. 2011 Apr 11. Malki L et al. N Engl J Med 2019;380:833





CCCA Treatment Recommendations

- Goals of therapy
 - Stop progression of hair loss/limit further scarring
 - Promote hair growth in areas with viable follicles
 - Relief of symptoms
- Decrease potential exacerbating factors: chemical relaxers, thermal straightening/styling, traction
 - Initiate treatment regimen based on clinicopathologic correlation
 - Stop progression of hair loss/limit further scarring
 - Promote hair growth in areas with viable follicles
 - Relief of symptoms
- Therapies:
 - Anti-inflammatory agents
 - Minoxidil
 - Limit progression of hair loss/limit further scarring
 - Promote hair growth in areas with viable follicles





CCCA Treatment Recommendations

No inflammation



Limit thermal, chemical, and mechanical trauma + Topical minoxidil

Inflammation present



As above

+ corticosteroids and/or antibiotics

End-stage scarring/
 Treatment failures



Hair transplantation, Wigs, camouflage







Racial Disparities in Dermatology

Pediatric Hair and Scalp Disorders

Dr. Candrice Heath



Understanding the Common Hair Care Routine of a Child with Tightly Coiled Hair

- Remove current style (30 min 3 hrs depending on style)
- Wash
- Condition
- Detangle (15 min with large tooth comb)
- Rinse
- Leave-in-conditioner
- Detangle (1 hr)
- Style (1-3+ hrs)







Tinea Capitis

- Differential diagnosis:
 - Scalp hyperkeratosis
 - Seborrheic dermatitis
 - Atopic dermatitis
 - Psoriasis
 - Sebopsoriasis
- Treatments
 - Griseofulvin microsize (135mg/5mL)
 - 20-25mg/kg/day x 8-12 weeks
 - Terbinafine
 - 10-20 kg : 62.5 mg/day
 - 20-40 kg: 125 mg/day
 - >40 kg: 250mg/day

Gupta AK. Drummond-Main C. Meta-Analysis of Randomized, Controlled Trials Comparing Particular Doses of Griseofulvin and Terbinafine for the Treatment of Tinea Capitis. *Pediatr Dermatol* 2013;30:1-6. Chen X et al. Systemic antifungal therapy for tinea capitis in children. Cochrane Database Syst Rev. 2016





Shampoo and Tinea Capitis Treatment

- Antifungal shampoos
 - May help with household spread
 - decrease transmissible fungal spores
- Conditioners
 - May help with household spread
 - Household should use it as well
- Watch out for hair dryness → breakage
- Parent may apply antifungal shampoo directly to scalp (ex. Ketoconazole shampoo)
- → wait 5-10 minutes
- → rinse
- shampoo scalp & hair with moisturizing shampoo
- condition with moisturizing conditioner
- style the hair as desired





Traction Alopecia

- Very common, may see pustules at area of tension
- Limit hair pulling and tight styles to avoid hair loss
- When discussing hair styling practices with parents:
 - Compliment, discuss, suggest!

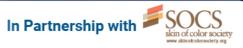












Racial Disparities in Dermatology

www.livderm.org

