Alopecia

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| Non-Cicatricial | Alopecia | | | | |
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| Disease (Synonyms) | Pathogenesis | Clinical features | Pathology | Treatment | Notes and Associations |
| Androgenic Alopecia (pattern baldness) | Androgen-dependent hereditary disorder; increased dihydrotestosterone with elevated 5-alpha reductase activity | Male: Frontotemporal recession and vertex balding; Female: Diffuse thinning over the vertex and parietal scalp; "christmas-tree" | Normal total number of hair follicles; increased number of vellus hairs | Finasteride, minoxidil, hair transplantation, spironolactone | Male: Hamilton-Norwood classification Female: Ludwig classification |
| Alopecia Areata | T-lymphocyte interaction with follicular antigens; HLA-A, B, C, and DR become expressed by follicle allowing T-cell cytotoxicity | Round, oval patches of hairloss; short- exclamation point hairs Variant: hair turning white (pigmented hairs lost) | Peribulbar lymphohistiocytic infiltrate; "swarm of bees" | Intralesional or topical corticosteroids, prednisone, topical immunotherapy | Totalis: entire scalp Universalis: whole body Ophiasis: band-like hair loss along periphery of scalp Associations: thyroid disease, vitiligo, type 1 diabetes mellitus, pernicious anemia, systemic lupus erythematous |
| Temporal Triangular Alopecia | Congenital disorder | Present at birth or acquired during first decade; triangular patch of hairloss or vellus hairs in temporal region | Normal number of follicles; almost all vellus hairs | None, usually persistent | Frequently bilateral |
| Telogen Effluvium | Large number of hairs enter telogen phase simultaneously; reaction to physical or mental stressors | Women age 30 to 60y; thinning of the entire scalp; may also affect axillary and pubic hairs | Increased number of telogen hairs (>20%); normal total number of hairs | Spontaneous recovery occurs within 6 months | Associated causes: Post-partum, severe infection, chronic illness, hypothyroidism, post-febrile |
| Anagen Effluvium | Secondary to insult to hair follicle, impairing mitotic activity | Hair follicles broken off diffusely due to tapering of hair shafts | Normal hair follicles | No specific treatment | Associated with radiation therapy and chemotherapy agents; Other causes: mercury, boric acid, thallium, colchicine, severe protein deficiency |
| Trichotillo- mania | Compulsive hair pulling; self-induced | Children age 5 to 12y; patchy hair loss of scalp, eyebrows, and/ or pubic hairs; varied lengths of regrowth; irregular borders | Incomplete disrupted follicular anatomy; trichomalacia and pigment casts; increased number of terminal catagen and/or telogen hairs | No specific treatment; hypnosis, behavioral modification, clomipramine, SSRIs. | |
| Pressure- Induced Alopecia | Pressure-necrosis of hair follicles with synchronized conversion of terminal hair follicles to catagen/ telogen phase | Solitary, rough, oval patch at site of pressure, usually upper occiput | Early: Vascular thrombosis, dermal inflammation Later: Increased catagen and/or telogen hairs, pigment casts, trichomalacia | Usually complete hair regrowth occurs | Most commonly in women after lengthy surgery |
| Lipedematous Alopecia | Not clear | Women with darkly pigmented skin; thick boggy scalp and hair loss | Approximate doubling of scalp thickness with increased edema and expansion of subcutaneous fat layer | No specific treatment | Follicular structures appear normal +/- ectatic lymphatic vessels seen |
| Cicatricial Alon | ecia | | | | |
| Disease (Synonyms) | Pathogenesis | Clinical Features | Pathology | Treatment | Notes and Associations |
| Lichen Planopilaris | Unknown, related to lichen planus | Most common in Caucasian women; scattered foci of partial hair loss associated with perifollicular erythema, follicular spines and scarring | Lichenoid interface dermatitis of the superficial follicular epithelium; inflammation affects upper portion of hair follicle | Antimalarial drugs and corticosteroids; difficult to treat | Greater than 50% associated with cutaneous or oral lichen planus; Frontal fibrosing alopecia: Frontotemporal hairline recession and eyebrow loss in postmenopausal women associated with perifollicular erythema, especially along the hairline |
| Discoid Lupus Erythematous | Chronic cutaneous lupus erythematous | Most common in adult women; scalp lesions have alopecia with erythema, epidermal atcopby, dilated and | Vacuolar interface alteration of epidermis and follicular epithelium; chronic inflammation with dermal mucin | Topical, intralesional or oral steroids; systemic retinoids; antimalarial medication | 50% of patients with discoid lupus have scalp involvement |

plugged follicular ostia present



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| Cicatricial Alop | cicatricial Alopecia (cont.) | | | | | |
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| Central Centrifugal Cicatricial Alopecia (hot-comb alopecia) | Follicular degeneration from premature desquamation of inner root sheath | Usually in African- American women; follicular loss mainly on the crown of the scalp | Premature desquamation of the inner root sheath; mononuclear infiltrate at the isthmus; loss of follicular epithelium with fibrosis | Doxycylcine or minocycline with potent topical corticosteroid; if highly inflammatory, rifampin or clindamycin | Previously thought to be due to chemical hair relaxers | |
| Aplasia Cutis Congenita | Congenital absence of skin and subcutaneous tissue; disruption of intrauterine skin development | Coin-sized defect or larger of alopecia, scalp >85% lesions | Atrophic, flattened epidermis; replacement of dermis with loose connective tissue; absence of adnexal structures | Cleansing and topical antibiotic ointment until healed; may need surgical repair | Adams Oliver syndrome: severe aplasia cutis congenita, cutis marmorata telangiectatica congenita, limb defects, and atrial septal defect Bart syndrome: aplasia cutis + epidermolysis bullosa | |
| Follicular Mucinosis | Dermal-type mucin deposited in epithelial structures; source of mucin - follicular keratinocytes or cell- mediated immune response | Children and adults age 30-40y; Primary - pink plaques composed of grouped follicular papules on face and scalp; Secondary - more widespread, chronic, older age group | Mucin in the outer root sheath and sebaceous glands; disconnected keratinocytes; perifollicular lymphohistiocytic infiltrate | Primary- many resolve spontaneously; consider topical, intralesional, systemic steroids, PUVA, dapsone, antimalarials; Secondary - treat underlying disease | Secondary type associated with cutaneous T-cell lymphoma and atopic dermatitis | |
| Acne Keloidalis | Unclear; possible foreign- body reaction to trapped hair shaft fragments | Most common in black men; follicular pustules and papules that may progress to firm, keloidal papules on occiput | Perifollicular, chronic inflammation at isthmus and lower infundibulum with lamellar fibroplasia | Chronic use of topical steroid or oral antibiotic; surgical excision | | |
| Dissecting Cellulitis of the Scalp | Follicular hyperkeratosis, often with bacterial superinfection | Most common in black men; firm scalp nodules on crown and upper occiput; develop into fluctuant ridges with purulent discharge | Perifollicular mixed inflammatory infiltrate; chronic abscess with sinus tracts | Isotretinoin, intralesional corticosteroids, oral antibiotics, and surgical approaches | Follicular occlusion tetrad: acne conglobata, hidradenitis suppurativa, pilonidal cyst | |
| Keratosis Follicularis Spinulosa Decalvans (Keratosis Pilaris Atrophicans) | X-linked recessive disorder of childhood; abnormality of follicular keratinization | Seen in childhood, often remits in puberty; alopecia involves the scalp, eyebrows, and eyelashes with perifollicular erythema and follicular hyperkeratosis | Follicular hyperkeratosis, atrophy of the follicle, and perifollicular fibrosis | Treatment with keratolytics, topical retinoids, topical and intralesional corticosteroids, oral retinoids - with limited success | Associated with eye abnormalities (corneal dystrophy); Chromosome Xp22.2-22.13 | |
| Traction Alopecia (End-Stage) | Prolonged traction by physical pressure | Most common in African-American women at 30-40y; several year history of bitemporal or frontal hair loss | Total number of terminal hairs markedly decreased; columns of connective tissue replace follicles | Permanent when traction is unrelenting (end-stage) | Associated with tight braids Biphasic form of hair loss - initially hair regrowth can occur | |
| Pseudopelade of Brocq | End stage alopecia or clinical variant of various forms of cicatricial alopecia | Affects Caucasian adults; oval or irregularly shaped atrophic patches; "footprints in the snow" | Atrophy; perifollicular inflammation at level of infundibulum, loss of sebaceous epithelium, fibrosis with absent hair follicles. | Resistant to therapy | Not a distinct disease but a clinical pattern of end-stage scarring alopecias | |

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| Syphilis | Treponema pallidum | Patchy, non-scarring areas of hair loss | Follicular plugging; a sparse, perivascular and perifollicular lymphocytic infiltrate with plasma cells | Benzathine penicillin G, intramuscular injections | "Moth-eaten" alopecia; 7% secondary syphilis associated with alopecia |
| Tinea Capitis | Dermatophyte infection: Trichophyton (endothrix) and Microsporum (ectothrix) | "Black dot" -alopecia with pinpoint black dots; kerion - boggy lesions with crust, severe inflammatory reaction | Hyphae or arthroconidia within/around hair shafts; can also be diagnosed by KOH prep | Griseofulvin, terbinafine, itraconazole | T. tonsurans most common in US Kerion - T. mentagrophytes, T. verrucosum Favus – T. schoenleinii |

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