

Common Pediatric Dermatology Diagnoses

Kimberly A. Horii MD

Associate Professor of Pediatrics

Division of Dermatology

Children's Mercy

Kansas City, Missouri



Common Pediatric Dermatology Diagnoses

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Common Pediatric Dermatology Diagnoses

- Vascular Birthmarks
 - Port wine stains
 - Infantile hemangiomas
- Dermatitis
 - Seborrheic dermatitis
 - Diaper dermatitis
 - Atopic dermatitis
- Impetigo

Common Pediatric Dermatology^{Di} Diagnoses

- Annular lesions
 - Tinea corporis
 - Granuloma annulare
- Hair loss
 - Alopecia areata
 - Tinea capitis
 - Telogen effluvium
 - Trichotillomania

Classification of Vascular Lesions

- Vascular tumors:
 - demonstrate cellular hyperplasia
 - Proliferate over time
 - Often not present at birth
 - *Infantile hemangioma*
- Vascular malformations:
 - composed of dilated/dysplastic vessels
 - Usually present at birth
 - *Capillary (port-wine stain)*
 - Lymphatic
 - Venous
 - Mixed malformations

Capillary Malformation (Port Wine Stain)

- Vascular malformation composed of dilated capillaries
- Always present at birth
- Pink to red flat patch
- Enlarges proportionately with overall growth
- Persists throughout life-may gradually darken
- May have underlying soft tissue hypertrophy
- If involves the periorbital region need to assess for glaucoma
 - Location of lesion may be important

Sturge-Weber Syndrome

- Classic triad
 - Facial capillary malformation (port-wine stain) in trigeminal nerve (V1) distribution
 - Higher risk if bilateral V1 involvement
 - Ipsilateral eye findings: glaucoma
 - Leptomeningeal and brain anomalies: leptomeningeal vascular malformation, calcifications
- May develop seizures and developmental delay
- Ophthalmologic exam and possible brain MRI

Infantile Hemangiomas

- Common vascular birthmark
- Affects approximately 4-5% of all infants
 - More common in females
 - Not usually present at birth
- Tumor composed of proliferating endothelial cells
- Rapid growth phase occurs during the first 2-6 months of life
 - Some high risk lesions may require treatment during the proliferative phase
- Gradual regression over many years

Infantile Hemangioma

- Classified as superficial, deep, or mixed
- Color can vary depending upon growth stage
 - Proliferating: bright erythema
 - Involuting: dull red, violaceous, or grey

Superficial Hemangioma



Deep Hemangioma



Mixed Hemangioma



Segmental Hemangiomas

- Newer classification system divides into localized, segmental, or indeterminate
- Segmental lesions defined as covering an “anatomic territory”
 - Can be linear/geometric
 - Usually large and plaque-like
- Lesion morphology may help predict outcome
 - Segmental lesions had poorer outcome



Localized Hemangiomas



Segmental Hemangioma



Regressing Hemangioma



Which to Worry About? Location Counts!

- Segmental
 - Increased complications & ulceration
 - Large facial: PHACE syndrome
- “Beard” distribution
 - Airway involvement
- Periocular
 - Visual axis impairment
- Multiple >5
 - Possible extracutaneous involvement
- Lumbosacral
 - Spinal dysraphism
- Perineal & Perioral
 - Risk of ulceration
 - Pain & infection
 - Feeding issues
- Disfiguring locations
 - Central face, nasal tip, perioral, & ear

PHACE Syndrome (OMIM 606519)

- Segmental facial hemangioma and ≥ 1 extracutaneous manifestation
- **P**osterior fossa brain malformations
- **H**emangioma- segmental facial or cervicofacial
- **A**rterial anomalies-cervical or cerebral arteries
- **C**ardiac defects-coarctation of aorta
- **E**ye abnormalities
- **(S)**ternal defects or supraumbilical raphe

Indications for Treatment

- Majority of hemangiomas do not require treatment
- Need for treatment depends upon
 - Functional compromise of vital structures
 - Rate of growth
 - Secondary complications
 - Ulceration
 - Location
 - Risk of disfigurement

Active Non-Intervention

- A technique for small lesions with good prognosis for spontaneous resolution
- Actively discuss expectations with the parents
- Close observation
- Discuss possible complications
 - Ulceration
- Patients with “high risk” hemangiomas should be followed closely during the early rapid growth period

Treatment of Infantile Hemangiomas

- Treatment may be required for life or function threatening complications or risk of disfigurement
- Treatment should ideally be instituted during the early proliferative stage
- No FDA approved treatment for hemangiomas
 - Oral propranolol
 - Topical timolol
 - Oral steroids
 - Intralesional steroids
 - Surgery

Seborrheic Dermatitis

- Commonly occurs in infants within the first three months of life
- Also known as “cradle cap”
- May be associated with proliferation of *Malassezia* species (yeast)
- Thick greasy scale on the scalp
- Greasy erythema and mild scaling behind the ears, on the central face, flexural folds, & diaper area
- Usually non-pruritic & self limited





Treatment

- Mineral oil for scalp
- Antifungal shampoos
 - Ketoconazole shampoo
 - Selenium sulfide shampoo
- Antifungal creams (effective against yeast)
 - Ketoconazole cream
- Low potency topical steroids
 - 1-2.5% Hydrocortisone ointment

Diaper Dermatitis

- Common dermatologic problem in infancy
- Variants of diaper dermatitis
 - Common
 - *Irritant contact diaper dermatitis*
 - *Infectious-candida, staph, strep*
 - Seborrheic dermatitis
 - Uncommon
 - Psoriasis
 - Zinc deficiency
 - Histiocytosis

Irritant Contact Dermatitis

- Affects up to 25% of infants wearing diapers
- Due to increased skin hydration, exposure to chemical irritants, & friction beneath the diaper
 - Chronic stooling can exacerbate
- Erythema involving the convex surfaces of the buttocks, perineum, lower abdomen, & thighs
 - Commonly spares the skin folds
- Severe cases may have superficial erosions

Treatment of Irritant Contact Dermatitis

- Frequent diaper changes
- Gentle cleansing
- Topical barriers ointments/cream applied thickly
 - Zinc oxide
 - White petrolatum

Treatment of Infectious Diaper Dermatitis

- Candidal diaper dermatitis
 - Antifungal cream (effective for yeast) or ointment
 - Azoles (clotrimazole, miconazole, ketoconazole)
- Impetigo
 - Topical or oral antibiotics
- Perianal Strep
 - Oral antibiotic
- All types benefit from topical barriers

Atopic Dermatitis

- Also known as eczema or “the itch that rashes”
- Affects at least 10-15% of children & adolescents
- Chronic inflammatory skin disease
- Characterized by
 - Xerosis
 - Pruritus
 - Recurrent skin lesions in a specific distribution
- Usually strong family history of “atopy”

Atopic Dermatitis

- Atopic dermatitis is often the first manifestation of atopy
 - Approximately 80% of children with atopic dermatitis will develop asthma or allergic rhinitis
- Onset usually within the first 5 years of life
 - 60% of cases begin by 1 year of age
 - 90% of cases begin by 5 years of age
- Prevalence decreases with age, though persistent or recurrent disease is common

Phases of Atopic Dermatitis

- Infantile phase (1 month-18 months)
- Childhood phase (18 months-puberty)
- Adolescent and adult phase

Infantile Phase Features

- Xerosis and evidence of pruritus
- Dermatitis begins on the cheeks or scalp
- Eventually involves the trunk and extensor extremities
- Diaper area usually spared
- “Rubbing” of face











Childhood/Adolescent Phase Features

- Chronic dermatitis
- Lichenification and excoriation
- Flexural surfaces of neck, arms, wrists, ankles, and legs
- Complaints of pruritus









Atopic Dermatitis Treatment

- Appropriate skin care regimen
- Eliminate or avoid triggering agents
- Treat
 - Active inflammation (red, rough areas)
 - Secondary infection
 - Pruritus
- Extensive patient & family education

Skin Care

- Patients with atopic dermatitis have abnormal skin barrier function
- Ointments or creams emolliate better than lotions
- Moisturizers may help repair the skin barrier
 - Promote skin hydration
 - Decrease pruritus
 - Baths may help hydrate the skin if used in conjunction with moisturizers

Inflammation



- When the skin is actively inflamed, anti-inflammatory therapy is necessary
- Topical steroids are still considered first line anti-inflammatory therapy for the treatment of atopic dermatitis in children
- Choose the lowest potency/strength topical steroid which will be effective

Topical Steroids

- Topical steroids should be used no more than twice daily
- Applied in combination with an emollient
- As inflammation subsides, attempt to decrease the strength/potency of topical steroid and/or frequency of use
- When inflammation recurs, restart topical steroid

Associated Co-morbidity of Atopic Dermatitis

- Children with atopic dermatitis have notable differences in sleep
 - Greater difficulty falling asleep
 - Frequent night awakening
 - Daytime sleepiness, behavior problems
- Psychosocial effects
 - Quality of life

Infections and Atopic Dermatitis

- Patients with atopic dermatitis are more susceptible to cutaneous viral & bacterial infections
 - *Herpes simplex* (eczema herpeticum)
 - Molluscum contagiosum
 - Human papilloma virus (warts)
 - *Staph aureus* (impetiginization)

Eczema Herpeticum

- Usually due to generalized type 1 Herpes simplex infection in patients with underlying atopic dermatitis
- Can masquerade as a severe sudden flare of atopic dermatitis or secondary bacterial infection
- Multiple superficial vesicles that evolve to form punched out erosions



Treatment of Eczema Herpeticum

- Recommend obtaining viral culture, DFA, HSV PCR
- Stop topical steroids or topical calcineurin inhibitors
- Systemic acyclovir at high doses
- May require IV therapy and hospitalization
- Often requires systemic antibiotics due to secondary impetiginization
- Bland emollients
- Contact isolation.

Bacterial Infections

- 90% of patients with atopic dermatitis are colonized with *Staph aureus*
- *Staph aureus* has increased adherence to the skin of atopics
- Presence of *Staph aureus* can be associated with an exacerbation of atopic dermatitis
 - Oral antibiotics are often necessary to treat secondary infection



Impetigo

- Most common bacterial skin infection in children
- Predominant organisms: *Staph aureus* (most common) & *Strep pyogenes* (GAS)
- More common in hot humid climates
- May occur at sites of trauma
- Can spread by direct skin contact

Non-bullous Impetigo

- Non-bullous impetigo-70% of cases
- Begins as erythematous macules and papules which develop into pustules
- Eventually pustules rupture leaving erosions covered by honey-colored crust
- Associated pruritus or pain
- Common sites: perinasal, perioral, & extremities



Bullous Impetigo

- Bullous impetigo-30% of cases
- Flaccid blisters with cloudy fluid
 - Rupture easily leaving shallow erosions and well demarcated collarettes of scale
- *Staph aureus* present in blister fluid
 - Releases exfoliative toxin that leads to blister formation
- Obtain culture from blister fluid





Work-up & Treatment of Impetigo

- For small isolated areas-topical antibiotic
- Systemic antibiotics are often necessary for larger areas of involvement or bullous impetigo
 - Cephalexin
 - Dicloxacillin
- Concern of resistant organisms (MRSA)
 - Bacterial wound cultures identify antibiotic susceptibilities
 - Clindamycin or Trimethoprim sulfamethoxazole may be options
- Local skin care

Tinea Corporis

- Well demarcated, annular, erythematous scaly plaques
- May have central clearing
- May have inflammatory papules or pustules in the advancing edge
- Usually pruritic





Tinea Corporis

- Topical treatment used twice daily to lesions and surrounding 1 cm area for 2-4 weeks
- Allylamines
 - Terbinafine 1% cream
- Imidazoles
 - Econazole 1% cream
 - Ketoconazole 2% cream
 - Clotrimazole cream
- Hydroxypyridone
 - Ciclopirox cream



Tinea Corporis

- Systemic therapy
 - Indicated for diffuse infection, not responsive to topical therapy
 - Immunocompromised patients
 - Always needed to treat tinea capitis
- Griseofulvin
- Itraconazole
- Terbinafine
- Fluconazole



Granuloma Annulare

- Skin colored subcutaneous papules or nodules often grouped in a ring configuration
 - No scale! (often misdiagnosed as ring worm)
 - Borders may be elevated
- Most commonly located on the feet, ankles, shins, and dorsal hands
- Usually asymptomatic
- Enlarge over time with central clearing
 - Size range from 0.5 cm to 3-5 cm



Granuloma Annulare

- Unknown etiology
- Histology is diagnostic
- Usually resolve spontaneously over many months to years
- Limited therapeutic options

Alopecia Areata

- Acquired non-scarring alopecia (bald spots)
- Cause is unknown, but autoimmune basis is hypothesized
- Males=females
- 20% of all cases occur in children
- Family history of alopecia areata is common
- Commonly seen in families with autoimmune diseases
 - Vitiligo, thyroid disease, rheumatoid arthritis, diabetes

Alopecia Areata

- Hair loss in circumscribed areas
 - May have several patchy oval or round areas
- Frontal, parietal areas commonly affected
- No underlying skin changes (no scale, erythema, or pustules)
- Usually asymptomatic





Alopecia Areata

- Prognosis:
 - Spontaneous remission is common with limited patchy hair loss if <1 year duration
 - 1/3 will have future episodes
 - ~10% will have chronic course
 - Worse prognosis if more diffuse involvement upon initial presentation
- Support Group and Information
 - National Alopecia Areata Foundation
 - www.naaf.org

Alopecia Areata Treatment

- Treatment options: (not FDA approved)
 - Active nonintervention
 - Supportive psychotherapy
 - Wigs/hair bands
 - Locks of Love www.locksoflove.org
 - Topical steroids
 - Intralesional steroids
 - Contact sensitization
 - Squaric acid, Anthralin

Alopecia Areata

- Differential diagnosis includes
 - *Tinea capitis*
 - *Telogen effluvium*
 - *Trichotillomania*
 - Traction alopecia

Tinea Capitis

- *Trichophyton tonsurans* is the most common dermatophyte to cause tinea capitis in the United States
- Humans are the main reservoir
 - More common in African Americans
 - Most common in 3-7 year olds
- “Classic clinical triad”
 - Scalp scaling, alopecia, & cervical adenopathy

Clinical Features

- Seborrheic type:
 - Diffuse scaling/dandruff, may have subtle hair loss
- “Black dot” type:
 - Patches of hair loss with broken hairs at follicular orifice
- Inflammatory type:
 - Pustules, abscesses, or kerions
 - Higher risk of scarring





Tinea Capitis Treatment

- Requires systemic treatment
- Griseofulvin
 - Gold standard
 - Good safety profile
 - Due to resistance, dosing may need to be higher than recommended on the package insert for 6-8 weeks
 - Absorption dependent on dietary fat intake
- Terbinafine
 - Another possible option with shorter treatment duration

Telogen Effluvium

- Acquired hair thinning (can be diffuse)
- Rapid conversion of scalp hairs
 - Growing phase → Resting phase (>25%)
- Normally: 85-90% is growing (anagen)
 - 10-15% is resting (telogen)
- Acute stressful events act as trigger
- No areas of focal alopecia, scale, or erythema
- May develop several months after a high fever, illness, surgery, traumatic or stressful event

Telogen Effluvium

- Diagnosis:
 - History of preceding event
 - Clinical exam
 - Consider obtaining CBC, iron studies, thyroid studies
- Treatment:
 - Reassurance & time

Trichotillomania

- Self induced hair loss resulting from pulling, rubbing, or twisting
- Individual often denies pulling hair
- Preadolescence is most common age of onset
- Hairs of varying lengths often in an unusual pattern
- Scalp>eyelash>eyebrow

Trichotillomania

- Treatment
 - Psychiatric referral
 - Cognitive behavioral therapy by an experienced therapist
 - Medications
 - Antidepressants

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