

PETECHIAE AND PURPURA

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Objectives

- Identification of petechiae and purpura
- Think through broad differential
- Inpatient vs Outpatient management
- History and exam to narrow differential and workup

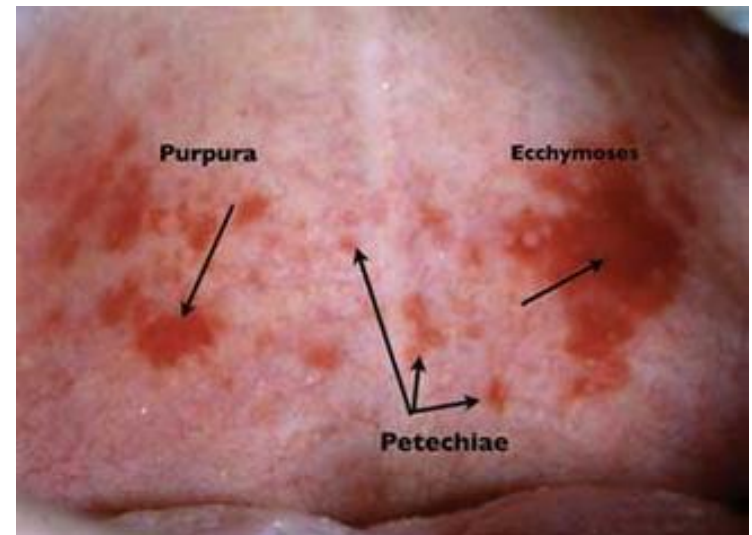
Petechiae and Purpura

- **Petechiae**

- Red/purple dots that represent bleeding from leaking capillaries
- Ecchymosis occur deeper in the dermal layers

- **Purpura**

- Petechiae that have coalesced and become bigger



Differential is BROAD

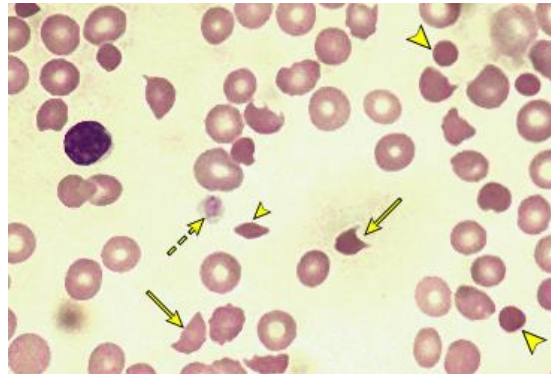
- Vasculitis
- Hemolytic-Uremic syndrome
- Immune thrombocytopenic purpura (ITP)
- Aplastic anemia
 - All cell lines will be down (infectious, medication, or leukemia)
- Trauma
- Hematologic
 - von Willebrand disease or hemophilia

Work-Up

- Labs to consider:
 - CBC - Low platelets? All cell lines down?
 - Coags/fibrinogen/bleeding time - DIC?
 - CRP/ESR - if CRP normal, sepsis is less likely
 - Blood cultures - in ANY febrile patient with petechiae
 - UA - if suspect HSP

Life threatening causes of petechiae

- Meningococcal septicemia
- Pneumococcal septicemia
- Disseminated intravascular coagulation (DIC)
- Rocky Mountain Spotted Fever



Non-life threatening causes of petechiae

- EBV, Adenovirus, other viruses
- Pertussis
 - Disruption of capillaries due to inc intravascular pressure from coughing or vomiting (above the nipple line)
- Strep pharyngitis
 - Sore throat without cough and petechiae on the soft palate



Neisseria meningitidis

- A leading source of community-acquired sepsis and meningitis
 - Serogroup B in < 5 years
 - Serogroups C, Y, and W135 in adolescents and adults
 - US has historic low since quadrivalent conjugate vaccine
- Transmission through respiratory droplets or secretions
- Risk factors
 - Age (younger than 1 year or between 15 and 24 years)
 - Crowded living conditions (military barracks, dormitories)
 - Cigarette smoking (active or passive)
 - Prior viral respiratory infection (especially influenza)
 - Family/household contact with meningococcal disease
 - Immunodeficiency

Neisseria meningitidis

- Fever followed by **petechial rash** (macules/papules → **petechiae**) and rapid deterioration
- Pallor, mottling, leg pain, or cold extremities are early sensitive signs
- **DIC**: Increasing petechiae, ecchymosis, or bleeding
- **Petechial rash can progress to Purpura fulminans**
 - Can lead to limb ischemia
- Adrenal insufficiency
- Shock



Neisseria meningitidis

- Dx:
 - History and physical exam
 - Blood and CSF culture are gold standard
- Treatment:
 - Favorable with early antibiotics (CTX) and correction of shock
 - Chemoprophylaxis for close contacts (>8hr and < 3ft), exposed to oral secretions (kissing, sharing drinks)
 - Rifampin, CTX, or Ciprofloxacin

Henoch-Schonlein Purpura (HSP)

- Most common vasculitis of childhood, < 10yo (peak 4-6 yo)
- Often follows URI, greatest during fall/winter
- IgA deposition in glomerulus, skin, and GI tract blood vessels
- GI
 - Colicky pain
 - Upper/lower GI tract bleeding —> +heme stools
 - Intestinal edema —> intussusception (throughout, not just ileocecal)
- Kidney: greatest risk for **morbidity**
 - Hematuria, proteinuria, azotemia or hypertension
 - Renal bx with suspicion of nephritic or nephrotic syndrome
 - IgA immune complex deposition in renal mesangium
- Arthritis/Arthralgia

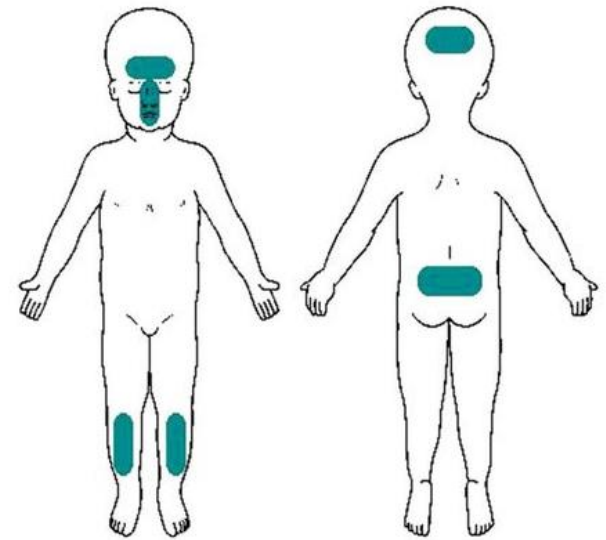
Henoch-Schonlein Purpura (HSP)

- Rash (not always first) followed by abdominal symptoms and arthralgia
- **Palpable Purpura** in pressure-dependent areas
- Dx: Clinical
 - No thrombocytopenia, normal PT/PTT
 - Skin bx: leukocytoclastic vasculitis with IgA deposition in the vessel walls
- Self-limiting illness
 - CKD and HTN observed up to 10 years

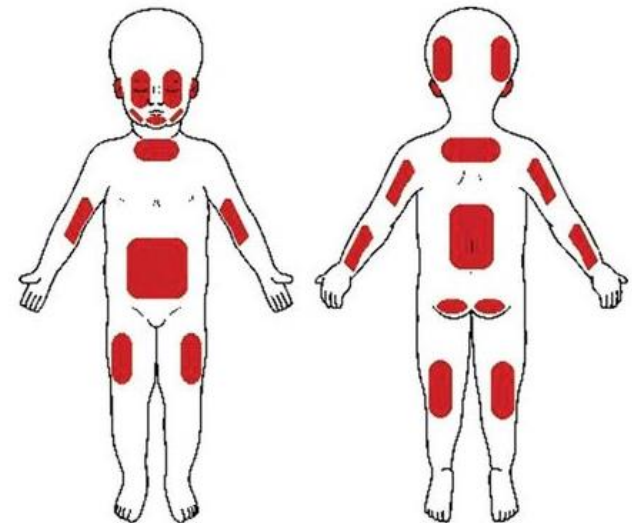


Trauma

- Bruise: bleeding into the dermis or subcutaneous tissue
- Accidental bruises over bony prominences (foreheads, knees, shins, and elbows)
- NAT
 - “Those who don’t cruise, don’t bruise” (exception: walkers)
 - Bruises over the upper arms, trunk, upper anterior legs, sides of face, ears and neck, flanks, genitalia, and buttocks
 - Shape of an instrument (belts, extension cords, human hand)
 - Petechiae in eye or mouth in suffocation (i.e. SI via hanging)



Accidental bruising pattern



Abusive bruising pattern

Trauma

- Infants and toddlers, disabilities at highest risk
- “Coining” and “cupping” (mistaken for NAT)
- Dx: whenever NAT concerned, need full workup
 - CBC (Hb, platelet count)
 - PT/PTT
 - Family history to screen for coagulation disorders
 - Others: (eg, bleeding time, coagulation factors)



Immune Thrombocytopenia (ITP)

- Most common cause of isolated thrombocytopenia
 - Generally 2-5 yo
- Autoantibody (IgG) to surface of platelets resulting in destruction in spleen and liver
- 50% following viral infection
 - Some can follow live virus vaccine, i.e. MMR
- Diagnosis of exclusion:
 - Platelet count $< 100k$ with no other cytopenias or abnormalities on peripheral blood smear
 - Absence of other clinical conditions

Immune Thrombocytopenia (ITP)

- Sudden onset widely spread petechiae/bruising over entire body, not-gravity dependent
- Mucocutaneous bleeding (petechiae, bruising, oral bleeding, epistaxis) usually $< 20k$



Immune Thrombocytopenia (ITP)

- Labs:
 - Peripheral blood smear (large and immature platelets)
 - Chem panel with LDH and Uric acid (Tumor Lysis)
 - DIC profile
 - Reticulocyte count
 - Direct Coombs (looks for associated hemolytic anemia)
 - Viral serologies: EBV, CMV, Parvovirus titers
 - Autoimmune etiologies (SLE, ALPS): ANA, dsDNA
 - Bone marrow not necessary with typical ITP features

Immune Thrombocytopenia (ITP)

- Usually short-lived (< 6 mo) without treatment
 - > 6 months is Chronic ITP, consider global immune problems
 - Severe complications (intracranial hemorrhage) in 0.1-0.5%
- Treatment (based on symptoms):
 - Observation
 - Corticosteroids (oral)
 - IVIG (fast 24-72 hours)
 - Anti-D immunoglobulin (WinRho)
 - Avoid NSAIDs
 - Platelets only if life threatening bleed