# 2023–2024 Pediatrics Review Syllabus

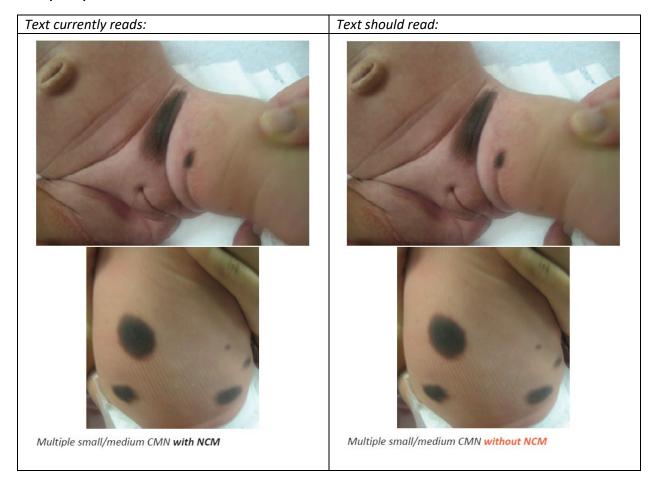
#### Adolescent Medicine & Sexual Health:

Page 24, Medical Issues in Females > Amenorrhea > Laboratory Assessment of Amenorrhea — FSH

Text currently reads:	Text should read:
<ul> <li>Laboratory Assessment of Amenorrhea — FSH</li> <li><u>High FSH</u> (secondary hypogonadism [a.k.a.</li></ul>	Laboratory Assessment of Amenorrhea — FSH <ul> <li><u>High FSH</u> (primary [hypergonadotropic]</li></ul>
hypogonadotropic hypogonadism])	hypogonadism)

#### Dermatology:

Page 154, Neonatal Dermatology > Other Birthmarks and Congenital Lesions > Congenital Melanocytic Nevi (CMN)



# MedStudy<sup>\*</sup>



# Page 157, Skin Conditions in Infants > Diaper Dermatitis

# Growth & Development:

## Page 311, Developmental Milestones > Developmental Milestones

Text currently reads:	Text should read:
<ul> <li>New checklists do not replace AAP recommendations for:         <ul> <li>Universal developmental screening with validated screening tools</li> <li>At 9-, <b>18-, and 30-month</b> well-child visits</li> <li>Autism screening</li> <li>At 18- and 24-month well visits</li> </ul> </li> </ul>	<ul> <li>New checklists do not replace AAP recommendations for:         <ul> <li>Universal developmental screening with validated screening tools</li> <li>At 9-, 18-, 24-, and 30-month well-child visits</li> <li>Autism screening</li> <li>At 18- and 24-month well visits</li> </ul> </li> </ul>

#### **Infectious Disease:**

#### Page 354, Immunodeficiency > AR 40

The test most likely to identify this patient's condition is: The t	est most likely to identify this patient's condition is:
B. Stool culture       B. St         C. Serum IgA       C. Se         D. Serum immunoglobulins to diphtheria       D. Se         E. Serum immunoglobulins to strains in the pneumococcal       E. Se	erum IgM cool culture erum IgA erum immunoglobulins to diphtheria erum immunoglobulins to strains in the pneumococcal <mark>olysaccharide</mark> vaccine

#### Metabolic Disorders:

### Page 387, Complex Molecule Defects > Lysosomal Storage Disorders (LSDs) > LSDs — Sphingolipidoses: Tay-Sachs Disease

Text currently	reads:	Text should r	ead:
Course	Clinical Signs	Course	Clinical Signs
0–6 months	Apathy, hypotonia, exaggerated startle reflex	0–6 months	Apathy, hypotonia, exaggerated startle reflex
6–12 months	Cherry-red <b>macule</b> , spasticity, rigidity, DD	6–12 months	Cherry-red macula, spasticity, rigidity, DD
12–18 months	Excessive drooling, bouts of unmotivated laughter, convulsions	12–18 months	Excessive drooling, bouts of unmotivated laughter, convulsions

#### **Ophthalmology & ENT:**

#### Page 562, Neck > Cervical Lymphadenopathy (LA) > High Yield — Cervical LA: Physical Exam

Text currently reads:	Text should read:
<ul> <li>Pharyngitis         <ul> <li>GBS</li> <li>EBV</li> <li>Adenovirus</li> <li>PFAPA</li> </ul> </li> </ul>	<ul> <li>Pharyngitis         <ul> <li>GAS</li> <li>EBV</li> <li>Adenovirus</li> <li>PFAPA</li> </ul> </li> </ul>

# **Rheumatology:**

## Page 618, Juvenile Idiopathic Arthritis (JIA)

Text currently reads:	Text should read:
<ul> <li>Definition of JIA:         <ul> <li>&lt; 16 years of age</li> <li>Persistent synovitis in 1 or more joints</li> <li>Must last for at least <u>6 weeks</u></li> <li>All other diagnoses excluded</li> </ul> </li> </ul>	<ul> <li>Definition of JIA:         <ul> <li>&lt; 18 years of age</li> <li>Persistent synovitis in 1 or more joints</li> <li>Must last for at least <u>6 weeks</u></li> <li>All other diagnoses excluded</li> </ul> </li> </ul>

# Page 618, Juvenile Idiopathic Arthritis (JIA) > PRINTO New International Classification Criteria (2019) — 5 Types

Text currently reads:	Text should read:
<ul> <li>New JIA Definition: &lt; 18 years of age, Sx &gt; 6 weeks</li> <li>1) Systemic JIA (arthritis need not be present)</li> <li>2) Early onset ANA+ JIA (two tests 3 months apart)</li> <li>3) RF+ JIA (two tests 3 months apart or CCP+)</li> <li>4) ERA/PsA JIA</li> <li>5) Other JIA (includes PsA)</li> </ul>	<ul> <li>New JIA Definition: &lt; 18 years of age, Sx &gt; 6 weeks</li> <li>1) Systemic JIA (arthritis need not be present)</li> <li>2) Early onset ANA+ JIA (two tests 3 months apart)</li> <li>3) RF+ JIA (two tests 3 months apart or CCP+)</li> <li>4) ERA/SpA JIA</li> <li>5) Other JIA (includes PsA)</li> </ul>