## NR 602 FINAL EXAM QUESTIONS BANK (129 Q&A) / NR602 EXAM QUESTIONS BANK (129 Q&A):RATED A 2022 |CHAMBERLAIN



- 1. The following are risk factors for hypertension in children and teens (choose all that apply):being obese. being exposed to second-hand smoke.
- In evaluating a 9-year-old child with a healthy BMI during a well visit, a comprehensive cardiovascularevaluation should be conducted by the following methods (choose all that apply): Obtain fasting lipid profile. / Assess diet and physical activity.
- 3. At what age is it appropriate to recommend dietary changes to parents if overweight or obesity is aconcern?

12 months old

- 4. The following are risk factors for type 2 diabetes mellitus in children and teens (choose all that apply):hyperinsulinemia: abnormal weight-to-height ratio.: Native American ancestry.
- 5. Screening children with a known risk factor for type 2 diabetes mellitus is recommended at age 10 or at onset ofpuberty, and should be repeated how often?
  every year.
- 6. Prediabetes in children is defined as (choose all that apply):
  impaired fasting glucose (glucose level ≥100 mg/dl, or 6

impaired fasting glucose (glucose level ≥100 mg/dL or 6.2 mmol/L) but ≤125 mg/dL or 7 mmol/L).impaired glucose tolerance (2-hour postprandial ≥140-199 mg/dL or 7.8 mmol/L-11 mmol/L).

- 7. Risk factors for dyslipidemia in children include (choose all that apply):family history of lipid abnormalities. family history of type 2 diabetes mellitus.
- 8. Screening cholesterol levels in children with one or more risk factors begins at what age? .2 years
- 8. An acceptable level of total cholesterol (mg/dL) in children and teens is: <170 mg/dL or 9.4 mmol/L.
- low birth weight, and poor infant growth are risk factors for type 2 diabetesTrue
- Prediabetes in children is defined as impaired fasting glucose (glucose level ≥100 mg/dL or 5.6 mmol/L but

≤125 mg/dL or 7 mmol/L) or impaired glucose tolerance (2-hour postprandial ≥140-199 mg/dL or 7.8 mmol/L-11mmol/L) or an A1C of 5.7% to 6.4%.
11. Screening for type 2 diabetes begins at age _10or at onset of puberty and continues every 2 years untiladulthood; at that point, the adult guidelines should be followed.
12. The AAP screening guidelines for total cholesterol levels in children and adolescents aged 2 to 19 years old areas follows: Acceptable level is <170mg/dL (<9.4 mmol/L), borderline is 170-199 mg/dL (9.4 mmol/L-11 mmol/L), and high is >200 mg/dL (≥11.1 mmol/L)
13. Children should be screened for family history of cardiovascular disease (CVD) beginning at age _3andshould be periodically updated annually or as required by risk factors during non-urgent health visits.
14. For at-risk children, fasting lipid levels should be tested after2years of age (but no later than 10 years ofage) and should be retested in 3-5 years if the values fall within the reference range.
15 Body mass index (BMI) should be measured beginning at age 2

16. For children between 12 months and 2 years of age for whom overweight or obesity is a concern, the use of
REDUCEDfat milk would be appropriate.
<ul> <li>17. Beginning at age _5 if BMI is ≥ 85th percentile, intensify dietary and activity changes to the parent.</li> <li>18. Infection with Corynebacterium diphtheriae usually causes: Pseudomembranous pharyngitis</li> <li>19. The tetanus infection is caused by CLOSTRIDIUM TETANI, an anaerobic, gram-positive, spore-forming rod. This organism is found in soil and is particularly potent in manure.</li> </ul>
<ul> <li>20. Sources of lead that can contribute to plumbism include select traditional remedies such as azarcon and greta. True</li> <li>21. Patients with plumbism present with which kind of anemia? Microcytic, hypochromic</li> <li>22. Intervention for a child with a lead level of 5 to 44 mcg/dL usually includes all of the following except: Chelation therapy</li> </ul>
<ul> <li>23. Ingested lead inactivates heme synthesis by inhibiting the insertion of iron into the protoporphyrin ring. This leads to the development of what kind of anemia?         microcytic, hypochromic</li> <li>24Basophilicstippling is often noted on red blood cell morphology in lead poisoning.</li> </ul>
25. Lead is significantly toxic to the solid organs, bones, and nervous system
26. Long-term complications ofLEAD poisoning include behavior or attention problems, poor academicperformance, hearing problems, kidney damage, reduced IQ, and slowed body growth.
27. Unless deleading procedures have been performed, however, most homes built before1957 contain lead-basedpaint.
28. A diet low in calcium, iron, zinc, magnesium, and copper and high in fat, which is a typical diet for childrenliving in poverty enhances oral lead absorption
29. In older homes, the point of greatest risk is the <u>window</u> because their sills and the putty have high leadconcentration. Because toddlers (age 2 to 3) are the ideal height to reach them and are often drawn to open ones, they are at greatest risk and summer is the riskiest season.  window
30. Symptoms of elevated <u>LEAD</u> levels include abdominal pain and cramping, aggressive behavior, anemia, constipation, difficulty sleeping, headaches, irritability, loss of previous developmental skills in young children, lowappetite and energy, and reduced sensations. Very high levels can result in vomiting, staggering walk, muscle weakness, seizures, or coma.
<ul> <li>31. A measure of 5 mcg/dL is now used to identify children with elevated blood lead levels.</li> <li>32. Most children with lead levels of 5-44 mcg/dL are treated with removal from the source, improved nutrition, and IRON therapy.</li> </ul>

33. Those with lead levels of 45-50 mcg/dL are treated with a  ${\tt CHELATION}$  agent such as succimer,

inaddition to the previously listed interventions.

- 34. For children with lead levels of greater than 51 mcg/dL, hospital admission with expert evaluation is likely the most prudent course to avoid serious problems (including\_ENCEPHALOPATHY\_) associated with markedly elevated lead levels
- 35. Which of the following represents the best choice of clinical agents for a child who has had a history of penicillin allergy who requires antimicrobial therapy?

  Cefdinir

36. The clinical presentation of UTI in children can be without the classic symptoms such as frequency, dysuria, orflank pain.  True
37. In younger children, UTI often manifests as IRRITABLITY_,LETHARGEY, andFEVERwith no obvious focal infectious source.  38. Older children with UTI often present withABDOMINAL pain, unexplained fever, or both; as childrenapproach puberty, flank pain becomes more common  40. UTI should be considered in infants and young children 2 months to 2 years old with unexplained fever, particularly in boys younger than 6 months and girls younger than 2 years who have a temperature greater than or equal to 39°C (≥102.2°F).
41. AURINALYSISshould be obtained in a child with unexplained fever or symptoms that suggest aUTI; however, 20% from UTI cases return a false-negative result.
42. Any of the following findings are suggestive, although not diagnostic, of UTI: positive leukocyte esterase, positive nitrite, more than5_white blood cells (WBCs) per high-power field in spun specimen, and bacteriapresent in unspun Gramstained specimen.
43. An acceptable method because of the low rate of skin and fecal contamination is a urine specimen collection viabag or from the diaper.
False, unacceptable because of the high rate of contamination  44. a single documented UTI in a child must be taken seriously. If an infant or young child 2 months to 2 years oldwith suspected UTI is assessed as toxic, dehydrated, or unable to retain oral intake,  HOPITALIZATION  is advised.
45. Oral amoxicillin, TMP-SMX, or a second-or third-generationCEPHALOSPORINis recommended as options for initial therapy for UTI in children
46. The use of _TMP-SMXhas a small risk of treatment failure.
47. Current evidence-based practice recommendations for UTI in Children indicate a _7to14day course of antibiotics because the outcomes are superior to a 1-to 3-day course in preventing spread of infection and subsequent renal scarring.
48. Although fluoroquinolone antibiotics have not been widely used in children, ciprofloxacin is approved by the
U.S. Food and Drug Administration (FDA) for use in pediatric patients for the treatment of UTI; this use is approved starting at age1years old.
49. Urinary tract imaging should be considered for all children with UTI, particularly if this occurs before toilettraining.
true 50. The two mainstays for imaging for UTI in young children are _RENAL BLADDER ULTRACOUND(RBUS) and voiding cystourethrography (VCUG)
51. RBUS is an easily obtained, noninvasive test but can miss a small number of high-gradecases
52. The benefits ofRBUS (no radiation exposure, non-invasive, minimal discomfort for child andparents), however, outweigh the slight increase in specificity of VCUG