

تشكر لجنة الطب والجراحة في جامعة مؤتة الزملاء : نور ظاهر الحجاج رعد بسام بني عامر & طارق نظمي أبولبدة لجهدهم الكبير في إعداد هذا الملف ، والذي يحتوي على أهم النقاط التي تأتي في امتحان الـ Mini – OSCE لمادة الأطفال .. وسيتم التحديث عليه بشكل مستمر ان شاء الله ..

Content

Α.	Developmental assessment (4)		
В.	Hematology (43)	Q	. Past Years Questions (333)
С.	Respiratory (65)		
D.	Endocrinology (102)		
Ε.	Nephrology (112)		
F.	Birth trauma <mark>(128)</mark>	-	5 th year – 14/8/2018 (334)
G.	Dermatology (133)	-	5 th year – 16/10/2018 (355)
н.	Chromosomal and congenital anomalies (174)	-	5 th year - 11/12/2018 (360)
Ι.	GI (208)	-	5 th year – 19/2/2019 (374)
J.	Growth chart (218)	-	5 th year – 14/4/2019 (392)
к.	CNS (228)	-	6 th year – 22/4/2019 (411)
L.	Skeletal (255)	-	5 th year – 6/8/2019 (421)
м.	Vaccines (280)	-	5 th year – 15/9/2019 (444)
N.	Jaundice (291)	-	5 th year – 12/11/2019 (475)
Ο.	Others (299)	-	5 th year - 19/2/2020 (491)
Ρ.	Important photos (321)		
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Developmental assessment



1- what is that skill ? Reaching out object

2- What is the age ?4 months

3- what is the skill he has on prone position ? Raises half of his chest



1- What is that skill ?Scribbling

2- what is the age ?14 months



1- what part of development is assessed in this image ? gross motor

2- what is the developmental age ?3 years



1- What is the age ? 18 months

2- No. of cubes can build ?3 cubes

3- How he tell his mother thing thathe wants ?Pointing to it.



Holds a spoon and gets food safely to mouth

1-What's the name of this skill? Reach out for toys

2-The developmental age is ? at least 4 months



What are the ages of these children?





3 Years

10 Months

1-What's the name of this skill? Play in role ...take turn

2-The developmental age is ? At least 3 years



1-The age of this baby not more than ?6 Months

2-why? Because he sits with rounded back



What is the age of this baby ? 8 months (mouthing)



what's the approximate age for these children ?





10 months

5 years

1-What's the name of this skill ? Pencil scribbling

2-What's his estimated developmental age?At least 14 months



1-What is his developmental age?18 months

2-How many words can he speak? 10 words



1-What can you see in the photo? Phobia to a stranger

2-What is the approximate age of the child?

9 months

تخيل الصورة التالية:

الدكتور عمر حامل ولد بعد ما أخذه من أمه اللي قاعده جنبهم. الولد بده يموت من الصياح

1-What is this milestone? Mature pincer grasp

2-At which age its expected to be found?10 months



1-What's the developmental milestone in each pic.? A-Walks around furniture. B-Mature pincer grasp.

2-Estimate the age of this child. (No range, Same child) 10 months.





The age of this child is at least?

A) 3 years.B) 3 months (raises half of the chest).C) 10 months (say bye bye).



What's the age of this baby? 8 months (mouthing).



1-What's the name of this test? Distraction test.

2-What does it test for?Auditory function, hearing, cochlearbranch of vestibulo-cochlear nerve.



What's the age of this baby? 9 months (stranger Anxiety).



What's the age of this baby? Around (10) months.





Age: 18 months





Age: 10 months

مش موجود ضمن مادة الدكتور عمر





Reflex: ATNR, disappears at 3-4 months

Age: 6 weeks

*ATNR: Asymmetrical Tonic Neck Reflex.

مش موجود ضمن مادة الدكتور عمر







Age: 10 months "Mature Pincer Grasp"

Age: 9 months "Prefers mother" (Not sure!!) Age: 8-9 months "crowling" 1-What is the Name of this reflex ? Moro reflex

2-When does it disappear ? 4-6 months

3-What causes it to be absent unilateral ? Brachial plexus injury





مش موجود ضمن مادة الدكتور عمر



Parachute Reflex



Symmetric Tonic Neck Reflex Appear at 6 – 7 months







Palmar grasp Reflex Disappears at 5-6 months.

Landau reflex. Develops at 3-10 months & lost at 36 months.

What is the age of this child ? 8 months (sit with straight back)

What is the age of this child ? 10 months (say bye-bye)





This child is drawing a circle, what's the age ? 3 years



What is the developmental age of this child? 4 years





What is the child doing in both pictures and what is the developmental age?





Responsive smile 6 weeks Spontaneous smile newborn

1-Masa can draw a triangle ,how old is she?

5years



Name of each skill ?

3 years (interactive play)



mature pincer grasp (10 months)


what is the developmental age for each one?

5 years





3 years



What is the age ?



(10 months)

1-Omar rides a tricycle what is his age?3 years

2-Masa lefts her head, says goo, ahhwhat is her age?3 months "vocalization"





A boy who says few words other than (mama, dada, baba), has just started to walk and has mature pincer grasp .What is the age of this child? One year old

What is the age for each child?





8 months

(many thought that she sits supported to chair but the doctor said she is unsupported, so if not clear ask about it.)

3 years old

1-At what age would you expect a child to draw a triangle?

5 years

2-At what age you expect a child to walk independently (Give a range)?12-18 months





Hematology

رمز النجمة يعني : موجود ضمن مرادة هذه السنة (مهم)

1-this blood film shows ? sickle cell RBCs (SCA)

2-type of inheritance ? AR

3-give one complication ? Vaso-occlussive crises ,, hemolytic crises



1-What's the abnormality in this blood film? Fragmented RBCs

2-Write 2 deferential diagnosis make this ?

A) G6BD

B) HUS

C) DIC

(u can write any hemolytic disease)





Patient diagnosed with anemia and treated with iron supplements for 3 months then blood film performed.

1- What is your diagnosis ?Thalassemia

2- how to confirm the diagnosis ?Hb electrophoresis





1-Mention two needed investigation To diagnose: Urine analysis & Renal biopsy ?!

2- DDx: HSP ,ITP







1-Mention the characteristic findings on this blood film.

Helmet cell, fragmented cells.

2-Mention 2 causes. HUS, DIC, G6PD



A blood film for a exclusively breast fed baby (Left for the patient and Right is normal)





1-What is the type of this anemia? Iron Deficiency Anemia

2-What other nutritional deficiency cause this pic? Vitamin D

السؤال غلط المفروض يسأل كالآتي

What other nutritional deficiencies might be seen in this patient?

1-What is the name of the cell on the biopsy? Reed sternberg cell

2-What is your diagnosis? Hodgkin's lymphoma





1-Describe what you see in the picture. Hyper-segmented Neutrophil.

2-What's this condition?

Macrocytic anemia due to Vit.B12 deficiency.





What's the type of this anemia? Sickle cell anemia.



Low mcv, low mch, low retics



1-What's the type of this anemia? Microcytic Hypochromic.

2-What 2 investigations you want to order? Hemoglobin Electrophoresis, Ferritin Level.

What's the next step in investigation? Hb Electrophoresis.



1-Findings in blood film: Microcytic & Hypochromic RBCs.

2-How do differentiate between IDA & Minor thalasemia: RDW index & Mentzer's index



This is a peripheral blood smear, What is the deficient nutrient? Vitamin B12



1-What is your diagnosis ? Thalassemia

2-Write two methods used in

treatment?

Blood Transfusion, Bone marrow transplant





A 6 month old child, his mother brought him when he had swollen fingers.

1-What is the sign ? Dactylitis

2-What's the diagnosis? Sickle cell anemia



1-What is your diagnosis ? Fanconi anemia



2-What other abnormalities you may find in this patient ?

Short stature

Abnormal thumb

Renal, cardiac, anomalies



1-What is the name of this Abnormality ? Fanconi Anemia

2-What's the definitive diagnostic test ?? White cell chromosomal fragility , high HbF





1-Describe the abnormality in the picture. Absent thumb.

2-Name the condition associated with this abnormality.Fanconi's anemia.





Anemic baby with splenomegaly .

1-What is the type of his anemia? Thalassemia Anemia.

2-What is the diagnostic test in this case? Hb-electrophoresis.



1-What's this x-ray sign? Sun-ray or Hair-on-end or crew-cut appearance.

2-What does it indicate? Thalassemia major

3-What other 2 findings in the face you look for? Frontal bossing Protruded maxilla.



A 5 month old boy is seen because of failure to thrive. As part of the investigation the following blood tests are done:

```
Hb 4.2 g/dl
WBC 12.3 x 10^9/l
Plt 211 x 10^9/l
Haemaglobin electrophoresis:
HbA - 0%
HbA<sub>2</sub> - 9%
HbF - 91%
```

1-What is the diagnosis? Beta Thalassaemia Major

2-What are the main treatment options?

1. Regular blood transfusions with iron chelation therapy; or

2. Bone marrow transplantation.

Respiratory system

1-What are the abnormalities in this X-ray film ?

Area of consolidation in the right middle lope with massive pleural effusion

2-Write 2 complications ?1-sepsis 2-meningitis



1- Give 1abnormality in this X_ray? Right tension pneumothorax.

2- What is the treatment?Chest tube or thoracostomy.



Newborn presented with respiratory distress & scaphoid abdomen. X-ray of patient is shown.

- 1- What is the diagnosis ? Diaphragmatic hernia
- 2- What is your next step for management ? Intubation then o2



6-month old baby presented with O2.sat 60%

1)Give one finding in the X-Ray? Boot shape, hyperlucent lung

2)Your Dx.? TOF





7 year-old male complained of SOB and cough , the vitals : fever, hypotension , the patient look sick, CXR is shown.



1- What are CXR findings ? Right middle lobe pneumonia with pleural effusion .

2- what is the treatment ?

Ceftriaxone with vancomycin



1-What is your diagnosis ? Tension pneumothorax

2-What is your immediate emergent management ?Needle decompression / chest tube



A 4 year old Child came with drooling and dysphagia.

1)What is your Dx? Foreign body ingestion "coin"

2)What is the management? Endoscopy


what is the name for this chest deformity ? pectus excavatum



1-Mention 2 clinical findings in this pt. Hand Clubbing, Cyanosis.

2-Mention 2 diseases cause it. Cystic Fibrosis, Infective endocarditis.

3-What system do you want to examine for this pt? CVS or RS.



name 3 causes for the sign seen here ? cystic fibrosis , infective endocarditis , cyanotic heart diseases



1-What's the name of this test ? Tuberculin skin test

2-What's your diagnosis?

Tb

3-When to interpret the result of this test

After 48-72 hours





This child did not take any vaccine till this age, Now he is presented with paroxysmal cough and coryza

1-What is your spot Dx? Pertussis , Whooping cough.

2-What is the cause of this condition? Bordetella pertussis.





1-Diagnosis : Meconium aspiration

2-Initial management:
Intubation & Oxygen
يوجد تفاصيل اكثر في المنجمنت موجود في)
المحاضرة ، يفضل الاجابة من المحاضرة)

3-Mention two complication: ARDS & Pulmonary HTN



1-your findings in this xray? Middle lobe pneumonia (lobar pneumonia)

2-what Is your choice of treatment? 3rd gen. cephalosporin + vancomycin





6 year old boy came to your clinic complaining of cough and fever ...

- 1- mention the prominent finding ? Heterogenous opacification on the right lung field
- 2-What is the most likely Diagnosis? Right interstitial pneumonia (atypical pneumonia)
- 3- name the most common microorganism .. ? M.pneumonia , C.pneumonia



1-What's your Dx? Tension Pneumothorax.

2-Give 2 signs on CXR.Hyper-lucent Rt. lung field.Shifted mediastinum (Tracheal deviation to Lt).Heart shadow shifted to Lt.

3-What's the treatment? Needle Thoracostomy & Chest tube.



Fever and cough RR =33



1-What are 2 physical finding on chest exam without using stethoscope?
 Dullness in percussion in Rt. Side.
 Asymmetrical chest expansion.

2-What are 2 findings in CXR?Hyper-dense or consolidation on Rt. side.Costo-phrenic angle obliterated or absent.

3-What's the most likely Dx.?

Rt. Lower and middle lobe Pneumonia.

4-What's the most common microorganism? Strep. Pneumonia.



Newborn (1st hour of life) with respiratory distress and cyanosis

1-Mention 2 prominent findings. Gas at Lt. chest side, Tracheal deviation to Rt.

2-What's your Dx? Congenital Diaphragmatic hernia.

3-Mention 2 medical lines of management.
 Intubation,
 NG tube (decompression).







Premature 34 weeks

1-What's the name of this sign? Ground glass appearance (reticulogranular pattern).

2-What's the most likely Dx.? Respiratory Distress Syndrome (RDS).



1-What's the name of this sign? Sail Sign of thymus.

2-What's the most likely Dx.? Normal X-ray.



A 6 YO child , previously healthy , started to complain from fever , SOB , shoulder pain 6 days ago

1-What's the Dx? Pericardial effusion.

2-Mention 2 things you'll hear by auscultation.Friction rub, Muffled heart sounds.Others: distended neck veins, hypotension.



Dx.: Cystic Adenomatoid Malformation!



<u>Dx.: Rt. Middle lobe pneumonia</u>



Dx.: Rt. Upper lobe pneumonia



<u>Dx.: Rt. Lower lobe pneumonia</u>



Q.32: Child with acute stridor.

1-What's the abnormality? Increased or swelling of the retro-pharyngeal space.

2-What's your Dx.? Retropharyngeal Abscess.

3-What's the tt? IV antibiotics & drainage.

4-What's the most serious complication for this condition?Mediastinitis.





Premature newborn with respiratory symptoms The 2nd pic after treatment

1-What's the management he received? Surfactant.

2-Mention 2 radiological signs.Ground glass appearance.Air bronchogram.





1-What's your Dx? Rt. sided pleural effusion.

2-Give 2 findings in the chest exam (not by auscultation).Stony dull percussion.Decreased chest expansion on Rt. Side (on palpation).

3-Give auscultation findings. Diminished vesicular breathing on Rt. Side.

4-What's the tt? Chest tube, Antibiotic.



Hint : premature 34 weeks....

1- mention the name of this sign?

Ground glass appearance and air bronchogram

2-What is the most likely Diagnosis? RDS





A 3 month old boy presented to the ER with wheezing and cough and the following x-ray.

What do you see in this x-ray and what is your diagnosis?

Hyperinflated chest Bronhiolitis





A full term newborn baby born by cesarean section with this xray .

1-What is your diagnosis?
(notice arrow pointing to fluid in the fissure)
Transient tachypnea of the newborn





A patient presents with barking cough and fever he was non toxic .

1-What is the diagnosis? Croup

2-What is the most common causative organism?

Parainfluenza virus



1-What are 3 findings during ear exam of this pt?

- 1. Erythematous
- 2. Bulging with fluid
- 3. Landmarks obscured
- 4. Likely NOT mobile on pneumatic otoscopy

2-What are the complication of AOM?

- 1. TM perforation
- 2. Hearing loss
- 3. Mastoiditis
- 4. Meningitis
- 5. Epidural abscess





Hx of boy who had URTI ,then developed bilateral nodular lesion in his legs

1-What's your Dx.? Erythema Nodosum.

Give a non-infectious cause. Sarcoidosis.

2-Give 2 microorganism causes this condition.
Group A strep.
Mycoplasma tuberculosis.
Chlamydia.



question about a patient(10 years old) who had recurrent chest infections , and FTT

1-what confirms your diagnosis ? chloride sweat test >60 on 2 separated days or genetic studies(more than 2 mutations) or abnormal nasal potential discharge

2-mention two microorganisms could cause this picture at his age ? strep.pneumonia,mycoplasma are common In his age,butwe don't know if the question wants more specific answer about m.cmicroorganisms regarding his condition ,too . Pseudomonads for example.



1-What is your immediate management for this child ? emergent upper endoscopy







Endocrinology

1-Mention 2 signs rather than signs of dehydration? clitoromegaly, ambiguous genitalia.

2-Mention one diagnostic test.Definitive test: measure 17- OH progesterone.

* Note: it's a case of congenital adrenal hyperplasia.





1-What is the diagnosis ?
21-beta hydroxylase deficiency
2-Investigation to confirm ur diagnosis ?
measure 17- OH progesterone





Figure 576-2 Three virilized females with untreated congenital adrenal hyperplasia. All were erroneously assigned male sex at birth, and each had a normal female sex-chromosome complement. Infants A and B had the salt-wasting form and received the diagnosis early in infancy. Infant C was referred at 1 yr of age because of bilateral cryptorchidism. Notice the completely penile urethra; such complete masculinization in females with adrenal hyperplasia is rare; most of these infants have the salt-wasting form.

-



1-Diagnosis: Congenital hypothyroidism

2-investigations : Serum TSH & T4



1-Mention 2 signs. Macroglossia, Umbilical hernia.

2-What's your Dx.? Congenital Hypothyroidism.

3-What lab test you want to do? TFT (serum T 3,T4, TSH).





4-What's the expected result? High TSH & low T3 & T4.





1-Mention 2 abnormalities:Umbilical hernia, protruded tongue

2-What's your diagnosis: Congeintal hypothyroidism





1 month old infant presented to you with a history of week cry and hypoactivity since birth

1- name 3 signs ?
Depressed nasal bridge
Plus the previous 2 findings
2- What is the most likely Diagnosis?
Congenital hypothyroidism
Hint : TSH ,↓T4






Mention 2 causes for this condition ? congenital rubella Galactosemia (this is cataract)



1-What is the finding ? Leukocoria / absent red reflex / white reflex

2- mention two causes of such condition ?
A- Catarcts , conginetal rubella ,
galactosemia (all are considered to be
one answer which is cataract),
B- Retinoblastoma



1-What is the name of this sign ? Leukocoria

2-Give 2 causes. Retinopathy of prematurity , retinoblastoma



Nephrology



Mention two findings in dipstick are useful to diagnose UTI:

Leukocyte esterase & Nitites





1-Findings is the photo ?
Dilation in ureters , dilation of renal pelvis and calyces , mild tortuosity on the left
2-Write down 2 complications ?
Recurrent UTI reflux nephropathy



patient with chronic renal failure presented with this ECG

1-What immediate test you should do for this patient? Serum potassium level

2-What immediate drug you should give? Calcium gluconate



1-Name the finding on ECG ? Peaked T wave

2-Give 2 modalities for treatment. Insulin, Calcium gluconate, B-agonist



1-What is this technique called? MCUG or VCUG (Voiding cystourethrogram)



2-What is your diagnosis? What is the usual presentation of this disease? Vesicoureteral reflux, UTI





1-What's the name of this test? VUCG or MCUG.

2-What's your next test? Late DMSA scan.



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Hx of URTI 10 days ago , BP was normal

1-Describe what you see. Bilateral peri-orbital edema.

2-What 1st & rapid test you want to do? Urine dipstick for proteinuria.

3-Mention 2 other things you want to examine.
Abdomen "ascites".
Scrotal swelling.
Lower limb edema.



4- 2weeks ago, this boy started to develop this clinical picture.Mention 2 tests you will order to support your diagnosis.urinalysis

serum albumin level

2 month female infant presented to you with non pitting edema

1- name this sign ?
Lymphedema
2- What is the syndrome that causes this feature?
Turner





This is a CT scan for a 3 year old male boy presented with unilateral kidney mass?

Describe the mass and what is the most likely diagnosis ? Left abdominal mass , heterogeneous , crossing midline , irregular shape

Neuroblastoma



After doing hearing tests, what's the next step you must do? Renal US, KFT.



1- What is the name of the structure labeled as A? Glomerulus

2- What's the site of action of Furosemide? D (Loop of henle)

3- What diuretic acts on part E? Thiazides



1- What is the electrolyte abnormality in this ECG? Hyperkalemia

2- What is the first line management? Ca gluconate



Hematuria & Flank pain

1-Mention two DDx: Neuroblastoma & Wilm's

2-Mention associated syndrome: Blackwith-Wiedmann syndrome & WAGR syndrome



abdominal CT scan





A 12 week old boy is admitted with a fever and being generally unwell. He is found to have a urinary tract infection. He is treated with antibiotics and several weeks after the infection resolves he has a number of investigations performed:

1-What kind of investigation is radiological investigation 1 and what does is show?

Micturating cystourethrogram (MCUG). This shows vesicoureteric reflux on the right side with right sided hydoureter and hydronephrosis

2-What kind of investigation is radiological investigation 2 and what does is show?
A nuclear medicine investigation called a DMSA (dimercaptosuccinic acid) renal scan. It shows that the right kidney is small and scarred with lesions at the upper pole and low/mid lateral territory



1-What's your diagnosis ? nephrotic –nephritic presentation "hematuria"

2-mention two lab tests to support your diagnosis ? Urineanalysis, Serum albumin level, 24-hour urine collection







Birth trauma

1-What's the Dx.? Capput saccidanum.

2-Mention one complication that may occur in that pt. Jaundice

3-Mention another DDx for this case. Cephalhematoma.







Name the finding you see in this newly born infant? Pitting swelling





1-What's the injured nerve and what's the name of this deformity? Brachial plexus...klumpky palsy

2-What are the root that are affected ? C7,C8, T1



1-Diagnosis: Left LMN lesion of facial nerve

2-Findings:

1) asymmetrical facies with crying.

2) The mouth is drawn towards the normal side,

3) wrinkles are deeper on the normal side,





Dermatology (Rash)

1- What is the most common organism ? Group A streptococci

2- Toxin mediated disease of that organism ?Scarlet fever

3- immunomediated disease of that organism ?Rheumatic feveror Post-streptococcal glomerulonepthritis



1- mention 2 DDx ? Scarlet fever and Kawasaki disease

2- the organ which affected by two DDx ?Heart







He has fever since 1 week , and conjunctivitis.

-write down 2 modalities of treatment of this patients? IVIG , aspirine



(Dx : Kawasaki)



1-diagnosis ? Hand-foot mouth disease

2-most causative organism? Coxsackie A





1-What is the diagnosis ? Hand-foot mouth disease

2- what is the organism ? Coxsackie A



1-What's the Dx.? Impitigo

2-what's the treatment? Antibiotic " penicillin, cefotaxim,....."



1-What's the Dx.? Mangolian spots.



2-What you want to do for this pt, & why? Nothing, because it normally disappears 3–5 years after birth.



1-What's the Dx? Hand-Foot-Mouth disease.

2-What's the causative micro-organism? Coxsackie virus A & enterovirus.









1-What's the name of this disease? Chickenbox.

2-What's the virus? Varicella zoster virus.



1-Name the disease. Shingles.



2-Mention 2 specific features for this disease.

Dermatomal distribution, cluster of vesicles on an erythematous base.





three days duration fever treated with amoxicillin and after that this rash appeared

What's your Dx.? Roseola Infantum.




What's the Most likely nutrient to be deficient? (Acrodermatitis enteropathica) give supplementation of Zinc.



Dx.= Measles (Koplik's Spots)



<u> Dx.: Orbital Cellulitis</u>





This boy presented with Cervical lymphadenopathy + splenomegaly

1-What investigation would you like to ask for?Infectious mononucleosis.Diagnosed by +ve Paul-Bunnell heterophile antibody test, Mono test, PCR.





1- whats the name of this lesion? Napkin rash

2-what's the treatment? Some thing related to zink

3-How to prevent? Keep dry , frequent changing of diaper , using cream











1-What is the spot diagnosis? Chicken pox

2-Mention one complicationBacterial infection (cellulitis)Pneumoniaencephalitis



1-What's the name of rash? Purpuric rash

2-Give two other symptom? Abdominal pain, arthralgia





Name two differential diagnosis for the child's eye condition? Kawasaki disease Measles



Name four possible complications of this condition:

- 1-GI: Intussusception + Hepatosplenomegaly + bowel perforation
- 2-GU: renal involvement with development of the nephrotic syndrome. + An infrequent complication of scrotal edema is testicular torsion, which may be suggested by pain and must be treated promptly
- **3-HLS: Lymphoadenopathy**
- 4-CNS: "A rare but potentially serious outcome of central nervous system (CNS) involvement is the development of seizures, paresis, or coma."





1-What's the Dx? Scarlet fever.



2-What's the tt? Penicillin (Note; the scarlet fever is caused by streptococcus pyogenes, GABHS).





picture of patient with purpuric rash in the lower limbs?

1-what is the diagnosis? HSP

1-mention 2 other clinical findings in this patient?ArthralgiaAbdominal pain







1-What is your diagnosis ? Erythema infectiosum

2-What is the causative organism ? Parvovirus B19

Erythema infectiosum



"Slap cheek" rash on the face, lacy rash on the extremities.

This patient presented with history of sore throat and fever 3 weeks ago and rash

1-What is the causative microorganism? Group A strep

2-What is the most serious complication? Either glomerulonephritis or rheumatic fever



1-Diagnosis?

Scarlet fever.



2-Name two complications?

peritonsillar abscess, sinusitis, bronchopneumonia and meningitis, or problems associated with immune system as rheumatic fever or glomerulonephritis.







This young boy presented with acute abdominal pain , hematuria and joint swelling , what is your diagnosis ? HSP



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1-What's the Dx.? Neonatal acne.

2-Why does it occur? Response to maternal androgens, (occurs in about 20% of normal newborns).



1-Describe what you see. Slapped cheeks appearance.

2-What's you Dx.? Erythema infectiosum.

3-What's the causative agent? Parvovirus B-19.





6 days fever

1-What's Your Dx.? Kawazaki Disease.

2-Mention 2 other organs you want to examine. Hands & feet (peeling), Trunk (rash), Tongue (strawberry).

3-Name the sign observed at the eyes. Non suppurative conjunctivitis.

4-What is the most serious complication? Inflammation of coronary arteries.





1-What is the diagnosis, Scarlet fever

2-what is the treatment ?? Penicillin

Note; the scarlet fever is caused by streptococcus pyogenes, GABHS



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1-What is the diagnosis? Hand-Foot-Mouth disease

2-what is the causative micro-organism ? Coxsackie virus A









Name the characteristic rash you see and mention the causan organism? Slapped cheeks rash Erythema infectious Parvovirus B-19





This child presented with a history of 3 days fever and upper respiratory symptoms the fever was documented as 40 degrees followed by this rash

1-What is your diagnosis? Roseola infantum (6th disease



1- Describe the skin findings.Café au lait spots

2- What's the mode of inheritance? AR





1- What is the most likely Diagnosis?
Hand – foot – mouth Disease
2- name the causing microorganism .. ?
Coxackie A (and EV 71)









1-What is the most likely Diagnosis?Mangolion spot2-What is the treatment ?None





1-What's your Dx.? Henoch-Schonlein purpura (HSP).

2-Mention other 3 symptoms.Abdominal pain.Arthralgia.Renal manifestation (hematuria).

3-If the pt has a toxic appearance with a rapidly evolving rash. What will be your most probable Dx.?Acute Meningococcemia.

4-What other body organs you will examine? Joints, abdomen, eyes.

5-Give 2 important lab tests you will order. KFT, Skin biopsy from the lesion, Urine analysis.



1-What is the criteria for the diagnosis of this disease?
Fever > 5 days & 4 out of 5: 1.Polymorphous rash, 2. Cervical lymphadenitis, 3. Changes in the lips and mucus membranes, 4. Extremity skin changes (redness, swelling, peeling of the skin), 5. Non-purulent bulbar conjunctivitis

2-What are the 2 most important drugs for the treatment of this patient? Aspirin & IVIG







4 year patient presented with rash, abdominal pain and ankle pain

What is the most likely Diagnosis? HSP





Chromosomal and congenital anomalies

1-What is the hematological disorder in this patient ?ALL , AML2-name the heart defect u see ?

Endocardial cushion defect







1-What is your interpretation for this x-ray ? Cardiomegaly

2-What is the most common cause for it in this child ?

Endocardial cushion defect







1-Mention 2 dysmorphic features ?1-webbed neck2- short stature Etc

2- What is the abnormality in the kidney ?Horse shoe kidney



1- What is the most specific cardiac anomaly? AV canal (endocardial cushion)

2- what is the lab test you want to check in 2nd trimester ? MS-AFP , (dec) Estriol , (dec) HCG , (inc) inhibin A (inc)



This image represent a case of untreated VSD. Now, she has developed this complication.

1-This complication is:

"Central cyanosis due to Eisenmenger syndrome"

N.B. Answering: "Central cyanosis" alone is not enough.

2-Mention two signs that you will see in the hands of this child? Peripheral cyanosis

Finger clubbing

Imagine:

A close view of the lips and mouth of a girl aged approximately 3 years. She has bluish discoloration of her lips. 1-Mention two cardiac abnormalities that cause early cyanosis:TOF, Tricuspid atresia

2-How do differentiate between respiratory Andy cardiac hypoxia: Hyperoxic test




Name the congenital anomaly you see in this infant? Low set and malformed ears



1-What is the cardiac anomaly? PDA

2-give one therapeutic intervention ? indomethacin Surgical ligation of PDA



1-What's your Dx.? Transposition of great vessels.

2-What's the most common presentation in neonates? Cyanosis.

3-What do you give immediately after birth? Prostaglandin (PG E1).





1 day old neonate he was cyanosed with o2 sat of 75% and pao2 = 85 mmhg

1-What's the CXR finding? Egg-on-string.

2-What's the cause of his cyanosis? TGA (Two parallel circuits)





1-What's your Dx? PDA.

- 2-Mention 2 signs on physical exam.
 - Hypertension.
 - Radio-femoral delay.
 - Machinery murmur at infra-clavicular area. Bounding pulse.



4 years healthy boy

1-What's this? ASD.

2-Give 2 findings upon examination? Ejection Systolic murmur. Fixed splitted S2.



1-What is the disease? TOF.

2-Give 3 findings of CVS physical exam. Ejection systolic murmur. Thrills. Single S2.

Cyanosis.

3-Give 2 complications. Clubbing, FTT.



1-A child with down syndrome has this CXR , on examination there a systolic murmur with no S3. What is the radiological diagnosis? Cardiomegaly

2-What is the most likely cause? Endocardial cushion defect (AV canal, VSD, ASD)







A 3 months old baby presented with tachypnea , and failure to gain weight , a continuous murmur is heard , what is the most likely diagnosis ?

PDA



A 16 year old girl came to you r clinic with primary amenorrhea & delayed puberty

1-What is this syndrome, what is the chromosomal pattern? Turner Syndrome, 45 XO

2-Which cardiac lesion do you want to rule out? Coarctation of the Aorta



In the following picture

1-what is the name of this syndrome? Turner syndrome.

2-what is the main cardiovascular abnormality in this patient? coarctation of the aorta



Source: Nat Cin Pract Endocrinol Metable 2008 Nature Publishing Group

A junior doctor examines the first child of a 28 year old woman as part of a routine "babycheck" prior to discharge from hospital. The baby is 20 hours old. The doctor notices that the baby is hypotonic and also finds a systolic murmur on auscultation of the heart. After further examination by a senior paediatrician the baby's chromosomes are analysed

1- What is the diagnosis? Trisomy 21 (Down syndrome)

2-What is the most likely cardiac defect? Atrioventricular septal defect (AVSD

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A question about downsyndrome pt , what is the chromosomal abnormality during meiosis ?

non disjunction of chromosome 21

1-What's your Dx.? Unilateral complete cleft lip.



2-What are the risk factors to have this condition? Genetic (1st degree relatives, monozygotics "60% concordance"). Environmental (drugs "phenytoin, valproic acid, thalidomide", maternal alcohol & tobacco use, dioxins & other herbicides, & possibly high altitude).

3-What's the tt? & when to be performed? Surgical closure; is usually done by 3 months of age.



Turner syndrome.



1-Mention 3 signs that you can see. Webbed neck, Wide-spaced nipples, Lymphedema of the limbs, Low hair lime

2-Mention Single best test to diagnose. Karyotyping.









1-What's the name of this syndrome? Turner syndrome.

2-What's the main CVS abnormality in this pt? Coarctation of the aorta (most common).

3-Name other associated congenital defects. Bicuspid aortic valve; later in life, post-stenotic aortic dilation with aneurysm may develop. Also renal anomalies e.g. horseshoe kidney.





1-What's the most common cardiac anomaly associated with this problem? AV canal.

2-Mention 3 signs you can find at the hand. Clindactaly, Cemian creases, Short fingers.

3-What syndrome is this? Down's Syndrome.

4-What is the chromosomal defect here?

Presence of an extra copy of genetic material on the 21st chromosome, either in whole (trisomy 21) or part (such as due to translocations).

5-Name 2 congenital defects that are associated with this condition. AV canal (aka endocardial cushion defect), VSD, ASD, valvular disease, duodenal atresia, annular pancreas, & imperforate anus.



1-What is the abnormality in the picture? Brush-field spots.

2-Name the condition associated with this. Down Syndrome.





1-What's the name of this syndrome? William's syndrome.

2-What's the chromosomal defect here? A small deletion of chromosome 7q11.

3-Name 2 associated congenital defects. Supra-valvular aortic & pulmonic stenosis & peripheral pulmonic stenosis.





1-What's the name of this syndrome? Noonan's Syndrome.

2-What's the chromosomal defect here? Autosomal dominant (AD) congenital disorder.

3-What's the male:female ratio of incidence? 1:1... as it's AD.

* Sometimes; described as "the male version of Turner's syndrome".



1- what is the most common congenital anomaly in this patient? Endocardial Cushing defects (ASD,VSD,AV canal)

 2-Name a gastrointestinal abnormality they have.
 Duodenal atresia , annular pancreas , hirschsprung disease, imperforated anus



1-Write down 2 dysmorphic features you see? Clinodactyly, Micrognathia, Macroglossia

2-what test you do to confirm dx? Karyotyping









This Karyotype indicates which syndrome Turner syndrome



1-What's the name of this sign? Short sternum.



2-What's the most likely Dx.? Edwards syndrome (Trisomy 18).

3-Give other 2 physical findings related to this disorder? (not seen in the image) Microcephaly.

Ocular hypertelorism. Low-set, malformed ears.



1-what is the sign seen? Double bubble sign

2-Give two possible GI diagnosis ?Duodenal atresia ,Annular Pancreas



1-What's the name of this sign? Rocker bottom feet.

2-What's the most likely Dx.? Edwards syndrome (Trisomy 18).

3-Give other 2 physical findings related to this disorder.
Microcephaly.
Ocular hypertelorism.
Low-set, malformed ears.



GI

What is your Dx?





Gastroschisis.

Omphalocele

1-What's your Dx.? Omphalocele.

2-What other findings could be found in this pt?
Cardiac defects.
Beckwith-Wiedemann syndrome (somatic overgrowth, hyper-insulinemic hypoglycemia, risk for Wilm's tumor).
Intestinal complications.



1-What's your Dx.? Gastroschisis.

2-What's the importance of this condition? Association with intestinal necrosis; although it's not associated with extraintestinal anomalies, but segments of intestinal atresia are common.



1-Give 2 DDx. Gastroschisis, Omphalocele.

2-What's the most important pre-op step of management? Covering.



infant on babylac formula and has diarrhea .
1-diagnosis?
Cow-milk allergy
2-treatment ?
Amino acid based formula





4.3 kg neonate presented with hypoglycemia and failure to pass meconium , barium enema was performed.

1- What is the diagnosis ? Left small bowel syndrome

2- What is the underlying pathology ? Infant of diabetic mother



1-What's this sign? Double-bubble sign.

2-What's the most probable Dx? Duodenal Atresia.

3-What other signs do you expect on examination?Abdominal distension, Jaundice.

4-What would be the typical presentation? Bilious vomiting after the 1st feed.



A 7 year old boy presented with intermittent abdominal pain and difficulty walking due to painful knees. He had been previously well except for a mild upper respiratory tract infection approximately one week ago. On examination he seems well but a non-blanching rash was seen on the buttocks and the legs (see photo). The rest of the body was spared

1-What is the diagnosis? Henoch-Schönlein Purpura (HSP).

2-What renal complication can occur? Glomerulonephritis


A 7 years child with sever gastroentritis What's the most important sign seen in the picture?

Sunken eyes.



Growth chart

12 year old male complained of chronic diarrhea , with positive family history of DM type 1. the growth chart of patient presented.

1- What is the abnormality at growth chart? Short stature

2-what is the most sensitive test for diagnosis of that patient? Anti-tissue glutaminase and endomysial IgA abs

3- what is the treatment? Gluten free diet

2 to 20 years: Boys NAME Stature-for-age and Weight-for-age percentiles RECORD # 12 13 14 15 16 17 18 19 20 lother's Statur Father's Stature AGE (YEARS) Date BMI* 76 Weight Stature Age 72 68 To Calculate BMI: Weight (kg) ÷ Stature (cm) ÷ Stature (cm) x 10,00 -66 65 -64 60 60 62 -62 155 155 s -60т 150 150 А -58 т -56 105-230 R -54 Е 100-220 135 -52 210 130 -50 -200 125 -48 190 120 85 180 -46 115 80 -170-44-110 160 -42-150-W -40-65 140 -38 60 130 -36-55-120 -34-50-110 -32-45 100 -30 -90 -80 35 -70: 30 Е -60 -60 25 -50 -50 G -20 15 -30 10 AGE (YEARS kg Ib 13 14 15 16 17 18 19 20 Published May 30, 2000 modified 11/21/00) SOURCE: Developed by the National Center for Health Statistics in collaboration with the National Center for Chronic Disease Prevention and Health Promotion (2000) ttp://www.cdc.gov/growthcharts SAFER + HEALTHIER + PEOPL

patient who didn't receive any treatment. What is the cause behind his growth delay?

Constitutional delay growth



This chart for pt. not receiving any treatment...

What's the cause for this change in the chart? constitutional growth delay



what is your interpretation for this growth chart for healthy child ?

Constitutional delay



What is the cause of the change in the growth of this child at the age of (catch up)?

Receiving Growth hormone.



3 years old boy

1-What's the abnormality in this Growth chart? short stature

2-What's the most appropriate Dx.? Celiac disease.



a 15 year old female presented with short stature, she also has constipation, skin changes and neck swelling, what is the most likely diagnosis?

hypothyroidism



What is your diagnosis ??

Cystic fibrosis





Birth 3 6 9

9 m/o pt , HC > 97th percntile

1- Give 2 DDx?HydrocephalusMacrocephaly (Hurlar syndrome or any pathological cause)

2- Mention one therapeutic measure.VP shunt



CNS

1-what's the causative organism ? Neisseria meningitidis (meningococcemia)

2-what dose it stain ? red





Name two differential diagnosis:

Meningitis

? Sepsis? Encephalitis? Subarachnoid hemorrage?



1-Describe: Skin glass test

2-Use:
differentiate between
blanching/nonblanching purpura
Nonblanching purpura:
Meningiococcemia &
Thrombocytopenia



1-What is your diagnosis? Meningococcemia

2-What are the lines of treatment? Antibiotics, Fluids, Inotropes (Any 2 of the 3 is enough)





Gram stain for CSF for 5 y/o Pt with meningitis.

1- Identify the microorganism?Strep. Pneumonia

2- What is the treatment for it ?
vancomycin
3rd generation cephalosporin
(Both drugs should be written)





1-Describe the study of CSF: CSF stain

2-Finding: Intracellular gram – diplococci (N.Meningiditis)





1-What's the abnormality in this child ? hearing difficulties

2-What is other abnormality you think that you will find in this baby ? Cerebral palsy with speech difficulties



1- What is the aspect of developmental assessment is affected by this patient ?Hearing and speech and language

2- What is the infection at infancy cause this problem ?Meningitis



in the following image:

1-what is the name of this sign? sunset eyes (setting sun sign).

2-what is the next examination Head circumference





Scenario of child with head trauma and the following lab results Na+: 110 mEq/L ,

Urine output: 3ml/kg/hr

1-What's your diagnosis: SIADH

2-Mention 2 lines of management:Fluid restriction,Hypertonic saline

Q.80: CSF results for a 5-day old neonate who complained of vomiting.

- WBC: 155/μL
- Neutrophils: 70%
- RBC: 0/ μL
- Lymphocytes: 30%
- Serum Glucose: 5 mmol/L
- CSF glucose: 2 mmol/L
- Protein: 80 mg/dL

1-What's your interpretation? Leukocytosis, High neutrophiles, High protein & glucose concentration.

2-What is the Dx? Acute bacterial meningitis.

3-Mention the most specific tt. Ampicillin + a 3rd generation cephalosporin. Csf result for 5 days neonate : WBCs = 22, RBC= 0, proteins= 50, Sugar= 3, Blood sugar= 5.

1-What's your interpretation? Normal CSF "for neonate".

2-Give 2 other CSF tests you will order.Gram stain.CSF Culture.PCR.Latex agglutination.

* Remember .. In neonates (<7 days) WBC's: up to 30, RBC: up to 50.



Hx of abdominal pain

1-What other systems would you like to examine?

CNS, Respiratory or Gl.

2-What's the most likely Dx.? Neuroblastoma.

3-Mention 2 non- radiological investigations to do.

VMA, Bone marrow biopsy.







This procedure is used to rule out what?

Meningitis or CSF infection.



This baby present with hydroceph , which device is used to decrease intracranial pressure

1-What is this device? External Ventricular Drain (EVD).

2-Mention one indication to use it. Infected shunt or obstruction.



1-What's the name of this test? Babinski sign.

2-Mention 2 abnormalities you expect to find in the lower extremities of this child.

Clonus, Hyper-reflexia «brisk tendon reflexes».





Tearing for the Habinetici Sign.

What is your diagnosis ?

Facial Nerve palsy





1-Give 2 findings seen in the picture. Café Au Lait Spot, Neurocatunios Nodules.

2-What's your Dx.? Neurofibromatosis.





Mentally retarded his brother has same condition (autosomal dominant)

1-Mention 2 signs. Ungual fibromas, Adenoma sebaceum, Shagreen patch.

2-What's the Dx.? Tuberous sclerosis.

3-Name 2 other investigations you will order to look for more signs. Brain MRI , kidney CT.



1-What's your Dx.? Hydrocephalus.

2-Mention 2 signs.Increased head circumference.Bulging anterior fontanelle.Sun set eyes





1-What is this sign? Sunset eyes.

2-Mention 2 things you'll find in examination.Increase HC.Bulging fontanel.Dilated scalp veins

3-What's the cause?

Increased intracranial pressure "hydrocephalus".



Growth chart of head circumference to age , the HC at 11th month became above 97th centile

1-What's your comment (what's that called)? Macrocephaly.

2-What's the most likely cause of this presentation? Hydrocephalus.

3-Mention 2 signs.Sun set eyes, Papillodema.

4-Mention 2 symptoms.Vomiting, Headache.



A child presented with this Ct scan And his head circumference was at 97th percentile.

1-Mention 2 signs
(Not sure if they were asking about signs on CT or signs on physical examination)
On CT: Widening of the ventricles
(ventriculomegaly) and effacement of sulci
On Physical examination: Sunset eyes & bulging
frontanelle

2-Mention 2 symptoms Headache & Projectile vomiting



1-what's the most probable diagnosis ? peri-Ventricular calcifications

2-Describe what you see ? congenital CMV some answered Tuberous sclerosis, since it causes brain calcifications.but remember : congenital Cmv >> C- shaped"periventricular calcifications congenital toxoplasmosis >> diffuse"scattered" intracranial calcifications


1-What is the spot diagnosis? Tuberous Sclerosis

2-What is the name of the skin lesion? "Ash Leaf Spots" (hypomelanic macules)





What is your deferential diagnosis ?

Neurofibromatosis 1





SKELETAL

1-What is this sign? Gower sign

2-mention 3 causes ? duchenne and becker muscular dystrophy , juvini e dermatomyositis

3-What Is the pattern of inheritance? X-linked recessive





1-What's this sign? Gower sign.

2-What's the disease that causes it? Proximal muscle wasting.



1-What's the name of this test? Scissoring

2-Name two finding in the examination? hyper reflexia clonus



Mention the name of this particular neonatal reflex? Tonic neck reflex



1-What's the sign called ? scissoring sign



2-Give one disease you can see this sign? spastic CP





1-What is the name of this sign ? Club foot

2-What is the association defect ? Meningomyelocele or spinal cord defect



<u>Dx.: Bilateral Club Feet</u> "TALIPES EQUINOVARUS"

1-What's the etiology for this condition?
Congenital (75%, usually an isolated abnormality).
Teratologic (associated with a neuromuscular disorder, such as myelomeningocele, arthrogryposis, or other syndromes).

Positional (normal foot that was held in the deformed position in utero.).

2-What are the complications of both treated or untreated pts? Untreated: severe disability. Treated: recurrence & stiffness.





1-Mention 2 clinical findings in the physical examination. Head Lag, C-shape or Drapes over in horizontal suspension.

2-What's the cause of these manifestations? Hypotonia.





1-What's the name of this sign? Frog-like Posture.

2-Mention 4 causes. Hypotonia, Congestive heart failure, Fulminant scurvey, Incorrect sleeping position.





1-What's the name of this sign?Scissoring Posture.

2-Mention 4 causes.
Spastic CP.
Cervical spondylosis with myelopathy.
Spinal cord trauma or tumors.
CVA, MS ...



What is The most Prominent feature in this child?

Muscle Wasting.



1- Mention two abnormalities in the X-ray?cupping of ulna.fraying of the metaphyseal region.

2- Your Dx? Rickets

3- Rx? Vit. D



What is the diagnosis?

Ricketts



1- What's your Dx? Rickets.

- 2-Give 2 radiological signs you can find. Cupping of distal head of ulna.
- Widening of epiphyseal plate.
- Shafts of the long bones become osteopenic & cortices become thin.
- Deformities of the shafts of the long bones are present.
- Bowing of long bones +- pathological fx.





1-What's your Dx? Rickets.

2-Give 2 abnormalities in the lower limbs.Bowing.Valgus & varus deformity.

Club foot.





1-What's your Dx.? Rickets.

2-What's the test you will ask for? Serum Ca+ & phosphate level.

3-What's the tt in this case? Vitamin D with Ca+ supplement.



A patient who is exclusively breast fed presents with the following X-ray.

1-What is the radiological diagnosis? Ricketts

2-Mention 2 abnormal labs? Low Ca+2 Low Vitamin D



In the next X-ray

1-what is the test you will ask for?

Serum Ca+ and phosphate level

2-what is the treatment in this case?

Vitamin D with Ca+ supplement



1-write one differential diagnosis for this pic ? Hemophilia

2-what u expect to see in the coagulation profile for this patient ? Prolonged PTT , normal PT , normal bleeding time



1-What's the name of this sign ? Rachitic rosary sign(*in rickets disease)



2-write 2 tests you have to do to confirm your Dx? Level V. D3 and PTH



1-Findings is this photo ?Cupping of distal head of radius and ulna.Widening of epiphyseal plate.



2-Investigations u need to do ? Vit. D , PTH



1-what is your diagnosis ? achondroplasia

2-except for sporadic cases, what is the mode of inheritance? autosomal dominant



1-What's the Dx.? Achondroplasia.

2-What's the mode of inheritance? AD.

3-Mention 3 complications.
Cervicomedullary compression.
Spinal stenosis.
Restrictive & obstructive lung disease.
Otitis media.
Tibial bowing.



1-What's this finding? Raccoon eyes.

2-What does it indicate? A closed-head injury that results in a basilar skull fracture.



Vaccines

The Jordanian National Immunization Program				
Age	Recommended Vaccines			
First Month	BCG			
61 Days	DPT-HBV-Hib	IPV		
91 Days	DPT-HBV-Hib	IPV	OPV	
121 Days	DPT-HBV-Hib	OPV		
9 Months	Measles	OPV		
18 Months	DPT	OPV	MMR	
6 Years (First Grade)	Td	OPV		
15 Years (10th Grade)	Td			

Mention 3 moderate side effects for DTP

Seizure

- Non-stop crying for 3 hours or more
- High fever

1- What vaccines you give for a 4 months old baby? (DTaP – Hib – IPV) – HBV – RVS

2- What is the route of administration ?IM

Age	1 m	61 day	91 day	121 day	10 m	12 m	18-24 m	6 Year
/accine								
BCG	G							
DTP		e	e	G			e	
Polio V.		IPU	Course and	OPU	OPU		OPU	OPU
нів		•	ø	ø				
нву		۲	e	۲				
Measles					ø			
MMR							Ð	

What are the absent vaccine by time?

1-at 91 day ... IPV and OPV 2-at 12 month... MMR 3-at 6 year... dT

Whats your Diagnosis for the following cases :

	Hbs Ag	HbsAb	HbcAb	IgM
1	Neg	Pos	Neg	Neg
2	Pos	Neg	Pos	Pos

1- immunity secondary to vaccination hepatitis B

2- Acute Hepatitis B virus infection

A Nurse is giving injection to a 2 month old baby in his Rt thigh.

What is she giving him? DTP,HIB,HBV and IPV



1-What do we call this vaccine? OPV.

2-What's the age of this child? 91 days (& more).



This baby took a vaccine . After 6 wks he developed this lesion with axillary LNs enlargement

What is your spot Dx?

Post-BCG vaccine abscess formation with regional lymphadenitis.



What's your Diagnosis for the following cases :

	Hbs Ag	HbsAb	HbcAb	IgM
1	Neg	Pos	Neg	Neg
2	Pos	Neg	Pos	Pos

- 1 immunity secondary to vaccination hepatitis B
- 2 Acute Hepatitis B virus infection
6 YO child with this scar on his abdomen

What vaccines would you like to give him? Post splenectomy pt should receive Pneumococcal vaccine or Meningococcal vaccine.



Mention 3 moderate side effect of DTP

The Jordanian National Immunization Program			
Age	Recommended Vaccines		
First Month	BCG		
61 Days	DPT-HBV-Hib	IPV	
91 Days	DPT-HBV-Hib	IPV	OPV
121 Days	DPT-HBV-Hib	OPV	
9 Months	Measles	OPV	
18 Months	DPT	OPV	MMR
6 Years (First Grade)	Td	OPV	
15 Years (10th Grade)	Td		

Seizure, High fever, Non-stop crying (Ref.: CDC.gov)

Jaundice

At second day of life of this child he presented with seizure , poor sucking , hypo tonia.

- 1- What are the tests you should perform ?
- 1- Random blood sugar
- 2- Serum bilirubin level3- CBC

2-What is the cause of seizure ? Kernicterus



Mention 2 causes for this condition seen in 2 months old baby, who has elevated indirect bilirubin levels.

hemolytic causes : (hemolytic blood diseases..G6PD,heredita ry spherocytosis, sickle cell)

conjugation disorders (crigler-najjar syndrome type 1) Hematoma



1-What's the Dx.? Neonatal jaundice.

2-When it's seen? When Bilirubin levels > 5 mg/dL.





mention two diseases requires such a treatment?



Cephalohematoma , Crigler najjar type 1 "conjugation disorder", G6pd "hemolytic disease"....



A baby on phototherapy ,mention 4 causes for jaundice.

G6PD Defeciency Hereditary Spherocytosis Hematoma

Crigglar- Najjar Syndrome.





1-What's this type of tt? Phototherapy.



2-What's the mechanism of action in the body? Transform unconjugated bilirubin to water soluble form to excrete it out of the body in Urine.



A 3 days old patient is put under phototherapy for his jaundice and otherwise he is normal.

Mention 2 causes for this condition. Any cause of indirect hyperbilirubinemia Sepsis Hemolysis Criggler Najjar Etc





Others

1-Name two signs you see in the image?

Ascites (Not abdominal distention... the later is a wrong answer)

Caput medusae

في الإمتحان... حكى الدكتور أنه ممكن نطلع 10 «ساينات» من الصورة.

2-What is a prognostic and diagnostic test that you will request?

Imagine the following picture:

An approximately 5-year-old boy with hugely distended abdomen, with dilated blood vessels, protruding umbilicus, and yellow discoloration of the eyes.

N.B. The jaundice was not visible due to the bad quality of the images by the Projector in the hall. When we complained to the doctor, he said that the yellowish discoloration was visible in his laptop and he promised to 'solve this problem' when correcting the answer sheets!!! How? God knows!

PTT

What's the estimated gestational age for both pics

Genitalia : 34 weeks Foot: 37 weeks at least (full term)





A previously well 10 year old boy presented to the emergency department following a 6 hour period of vomiting. His parents say that he has been feeling generally unwell for the last week and has lost some weight. On examination he is found to have "sighing" hyperventialtion and to have a reduced level of consciousness - only responding to a sternal pressure by moaning. Urgent blood tests show the following:

1-What is the diagnosis? Diabetic ketoacidosis

2-Why is he hyperventilating? Kussmaul respiration (secondary to acidosis.)

Blood test results			
Haematology	Biochemistry	Venous Blood Gas	
Hb 17.5 g/dl	Na 136	pH 7.06	
WBC 23.4 x 10 ⁹ /l	К 6.4	Bicarb 9.2	
Plt 429	12.5	Base Excess -14.8	
	Cr 112		
	Glucose 33.5 mmol/l		

Mention two diagnostic tests are useful to diagnose child with recurrent pneumonia and history of meconium ileus:

-Sweat chloride -Genetics



1-what is ur diagnosis ? TOF

2-write down 2 modalities of treatment ? Surgerical : VSD closure and relieve of RVOT obstruction Medical : maintain HCT 45-50% , infection , TET spells , knee chest position





1-What's the name of this machine? Incubator.

2-Mention 2 functions for it.Thermoregulation.Isolation from infections.



1-Name one diagnostic test you would perform for this patient? Sweat chloride test or Genetic testing or Nasal Potential Difference

2-What is the mode of inheritance? Autosomal recessive



A child presented with fever for 1 week

1-Give 2 lab test to help in diagnosis: CRP , CBC

2-What other sites you would like to examine: Mouth, hands and feet, heart







1-identify the MO? gram positive bacilli

2-Give one example for the MO? listeria spp. ,,, clostridium spp



1-This device is called ? self inflating bag (ambu bag)

2-used for what ? apnea ,, resuscitation ,, croup + epiglottitis



What is the name of this device ? Inhaler spacer

What drug is commonly used in it ? SABA





What are these devices





Inhaler

Spacer

This patient has been admitted to ER after a bee bite ,he was hypotensive.

1-what's your diagnosis ? Anaphylaxis

2-What's your immediate management ? Epinephrine injection



A child 11 mothns old who weighs 10 kgs presents with moderate hyponatremic dehydration signs of dehydration. His Na+ level is 125

1-Calculate the sodium maintenance Maintenance = 2-4 =mEq/kg. So approximately 30 mEq

2-Calculate the sodium deficit

Deficit = 100 mEq

Na correction = 60 mEq

Total = 100+60 = 160mEq





A patient presents with sand paper like rash & sore throat.

1-What is the causative organism? Group A Strep (Streptococcus pyogenes)

2-Mention 2 complications. Glomerulonephritis

Rheumatic fever





1-What's your Dx.? Congenital Cataract.

2-Mention 2 causes. Rubella, Galactosemia.





1-What is this? Oral thrush secondary to Candida (mucosal candidiasis).

2-What's the cause? Oral Candida/ fungal infection.





Mention WHO recommendation for breastfeeding Initiation within the 1st hour of life.

- 1. Exclusive breastfeeding (that is the infant only receives breast milk without any additional food or drink, not even water).
- 2. Breastfeeding on demand (that is as often as the child wants, day & night).
- 3. No use of bottles, teats or pacifiers.

1- x ray of steeple sign(croup) :

Name the most common organism ? Parainfluenza virus

2 - pic of tuberulin test :

Name of the test? , when to read ? After 48_72 h How to measure ?Depend on the induration not erythema

- 3 pic of asymmetrical moro reflex , causes ?Fx of clavicle , humerus fx ,Erbs palsy ..
- 4 Pic of down syndrome :

Write two abnormalities ? Hypertelorism and micrognathia Karyotype ? Trisomy 21



5 - Picture of clubbing: two diffirential : cystic & celiac

6 - visicles (chicken pox) : two complicationsBacterial infection (cellulitis)encephalitis

7 - RDS x ray (air bronchogram) granular appearance : Treatment : surfactant oxygen infection control

8 - Bronchilitis x ray (not sure if it is bronchiolitis): Therapy : oxygen and fluids



10 - 6_11 months old patient with cow milk feeding history with picture of iron defiecency anemia :

Dx (IDA)

two tests to confirm : serum iron , ferritin, TIBC

(10+ 11 + 12 + 13) 4Q on developmetal assessment لازم تحفظ کل صغیرۃ و کبیر بالسلاید))



Important photos

<u>Angular Chelitis</u>

<u>Serous Otitis Media</u>
















<u>Lt. Hypoglossal Nerve Palsy</u>



<u>Lt. Vagus Nerve Paralysis</u>



Dx.= Ankyloglossia "Tongue Tied"



Hyper-telorism & Flat nasal bridge



Dx.= Acute Follicular Tonsillitis.



<u>Microcephaly</u>

<u>Dermatitis</u> <u>Herpetiformis</u>













<u>Holoprosencephaly</u>





Past Questions

Mini-OSCE Pediatrics 14/8/2018

Done by : Ammar Adaileh

Question (1)

At second day of life of this child he presented with seizure , poor sucking , hypo tonia.

1- What are the tests you should perform ?

- 1- Random blood sugar
- 2- Serum bilirubin level
- 3- CBC

2-What is the cause of seizure ?

Kernicterus



Question (2)

4.3 kg neonate presented with hypoglycemia and failure to pass meconium , barium enema was performed.

1- What is the diagnosis ? Left small bowel syndrome

2- What is the underlying pathology ? Infant of diabetic mother



Question (3)

Newborn presented with respiratory distress & scaphoid abdomen. X-ray of patient is shown.

1- What is the diagnosis ? Diaphragmatic hernia

2- What is your next step for management ? Intubation then o2



Question (4)

7 year-old male complained of SOB and cough , the vitals : fever, hypotension , the patient look sick, CXR is shown.

1- What are CXR findings ?
Right middle lobe pneumonia with pleural effusion .
2- what is the treatment ?
Ceftriaxone with vancomycin



Question (5)

Patient diagnosed with anemia and treated with iron supplements for 3 months then blood film performed.

- 1- What is your diagnosis ? Thalassemia
- 2- how to confirm the diagnosis ? Hb electrophoresis



Question (6)

1- What is that sign ? Rachitic rosary sign

2- How to confirm yourdiagnosis ?Serum vitamin D3 and serumcalcium



Question (7)

12 year old male complained of chronic diarrhea, with positive family history of DM type 1. the growth chart of patient presented.

1- What is the abnormality at growth chart? Short stature

2-what is the most sensitive test for diagnosis of that patient? Anti-tissue glutaminase and endomysial IgA abs

3- what is the treatment? Gluten free diet



Question (8)

1-Mention 2 dysmorphic features ? 1-webbed neck

2- short stature Etc

2- What is the abnormality in the kidney ? Horse shoe kidney ?!?!



Question (9)

- 1- What is the most common organism ? Group A streptococci
- 2- Toxin mediated disease of that organism ? Scarlet fever
- 3- immunomediated disease of that organism ? Rheumatic fever or Post-streptococcal glomerulonepthritis



Question (10)

1- mention 2 DDx ?
Scarlet fever and Kawasaki disease
2- the organ which affected by two
DDx ?
Heart



Question (11)

1- What is the aspect ofdevelopmental assessmentis affected by this patient ?Hearing and speech and language

2- What is the infection at infancy cause this problem?Meningitis



Question (12)

1-What is the diagnosis
?
Hand-foot mouth disease

2- what is the organism?Coxsackie A



Question (13)

1- what is that skill ?
Reaching out object
2- What is the age ?
4 months
3- what is the skill he has on
prone position ?
Raises half of his chest



Question (14)

1- What is that
skill ?
Scribbling
2- what is the
age ?
14 months



Question (15)

1- What is the age ?
18 months
2- No. of cubes can build ?
3 cubes
3- How he tell his mother thing that he wants ?
Pointing to it.



OSCE Pediatrics 15/8/2018

Done by : Ammar Adaileh

Essay station

Child comes complaining of generalized edema with +3 protein in the urine and associated with severe abdominal pain with 39 c and hypotension.

1- What is the diagnosis ? Nephrotic syndrome

2- How can we confirm dx?

- 1-24 hr urine collection
- 2- lipid profile
- 3- albumin level

3- What is the treatment?

Prednisolone for treating underlying cause , albumin and furosemide for treating acute condition.



4- If the treatment is failed , what is your next step ? Renal biopsy

5- What is the ddx of abdominal pain other than edema in acute condition of that case ? And how we treat it ?

Spontenous bacterial peritonitis and treated with ceftriaxone and vancomycin

History station

2 year-old female patient with hx of seizure came to you at ER.

Take relevant hx.

Neonatology discussion

Neonate with Irritability and poor feeding. (neonatal sepsis discussion)

Examination station

Palpation of abdomen and examine for organomegaly, and answer the qs from examiner.

Qs :

1- mention 5 causes of hepatomegaly

Infection(hepatitis)

Infectious mononucleosis....Sickle cell anemia...Thalassemia...Leukemia...Heart failure mention general signs for patient with hepatomegaly Jaundice..Palmer erythema..spider nevi..gynecomastia

Mini-OSCE Pediatrics 16/10/2018

Done by : Nebal Altarawneh & Bana Alkaraki 1 - developmental assessment

اجا عليها ٥ اسئلة واحد منهم جدول نحط فيه كم كلمه بحكي بكل عمر

- 2 Infant of diabetic mother
- 3 Hypothyroidism
- 4 Iron def anemia

5 -ECG for hyperkalemia : dagnosis and treatment الصورة نفس اللي بسلايد دسلمي بالضبط

6 -HSP

7 -Bleeding disorder

8 - Mecuniom aspiration

9 -X ray ... Pneumatocele (organism , treatment)

10 – Feeding :

11 -Side effect of phenytoin

12 -URTI وانه شو السبب tonsilكانت صورة

OSCE Pediatrics 16/10/2018

Done by : Nebal Altarawneh & Bana Alkaraki

Essay station :

للدكتورة سلمى Acute kidney injury كان فحص ونطلع الفايندنج انه هيماجوريا وبروتين يوريا والسؤال الثاني انه شو يلي بصير metabolic acidosis or alkalosis والسؤال الثالث complication

- <u>Hx</u>:rash (roseola infantum)
- <u>Examination</u>: growth chart and vaccination
- <u>Discussion</u>: neonatal sepsis

Mini-OSCE Pediatrics 11/12/2018

Done by : Hamza Wadi & Abdullah Alawneh & Tareq Abu-lebdeh
- 1-Whats the sign?
- steeple sign
- 2-Whats the diagnosis?
- (croup)
- 3-Name the most common organism ?
- Parainfluenza virus



1- Name of the test?

tuberulin test

- 2- when to read ?
- After 48_72 h
 - 3- How to measure ?
- Depend on the induration not erythema



- 1-Describe the reflux?
- asymmetrical moro reflex
- 2-causes?

Fx of clavicle , humerus fx ,Erbs palsy ..



Pic of down syndrome :

1- Write two abnormalities ?Hypertelorism and micrognathia

2- Karyotype ?

Trisomy 21



- 1-name the sign?
- Finger clubbing
- 2- two differential?
 Cystic fibrosis & celiac



- 1 visicles (chicken pox)
- 2-two complications
- Encephalitis
- Meningitis
- Reyes syndrom
- Transverse myelitis



• 1-finding?

- poor lung expansion
- air bronchogram
- reticular granular appearance.
- 2-diagnosis?
- RDS
- 3- Treatment ?
- surfactant oxygen infection control





-6_11 months old patient with cow milk feeding history with picture of iron defiecency anemia :

1-Dx ?(IDA)2-two tests to confirmserum iron , ferritin, TIBC



Bronchilitis x ray (not sure if it is bronchiolitis):
 Therapy : oxygen and fluids



Q9) 2 year old boy with jaundice (with hx and invx): Dx , specific investigation , another differential for him

(10+ 11 + 12 + 13) 4Q on developmetal assessment (لازم تحفظ کل صغیرة و کبیر بالسلاید)

OSCE Pediatrics 11/12/2019

Done by : Hamza Wadi & Abdullah Alawneh & Tareq Abu-lebdeh

Essay station :

4 years old male patient complain of diarrhea for 4 months duration his weight is 19 kg , length is 100 cm

1 - calculate BMI

19

2 - From what he suffer obisity

(because BMI percentile more than 95 percentiles

- 3 What are the percentiles of hight and weight
- 4 What is your Dx : toddlers diarrhea
- 5 How to counsil his family :

avoid fruit juices with reassurance

• <u>Station 1(Hx) :</u>

9 years old with 1 day hx of red urine (Dx PSGN)

• Station 2 (Examination) :

Examination for suspected meningitis

(Specific tests) (assesment) (most common organisms) (treatment)

• Station 3 (open discussion) : (Dx : RDS)

-27 weeks old , birth weight 1 kg , respiratory distress , central cyanosis , Granting , intercostal, supcostal, suprastarnal retraction

(What investigations to order and expected results)

(most likely dx) (Name of disease) (Acute complications) (Treatment for RDS)

Mini-OSCE Pediatrics 19/2/2019

Done by : Mamoon Saleh • What is the hematological disorder in this patient? ALL, AML

-name the heart defect u see ?
Endocardial cushion defect
-write one complication ?
endocarditis





1-write one differential diagnosis for this pic? Hemophilia

2-what u expect to see in the coagulation profile for this patient? Prolonged PTT, normal PT, normal bleeding time



He has fever since 1 week , and conjunctivitis. kawasaki write down 2 modalities of treatment of this patients? IV immunoglobuline aspirin



1-what is ur diagnosis ? TOF

2-write down 2 modalities of treatment ? Surgerical : VSD closure and relieve of RVOT obstruction Medical : maintain HCT 45-50% , infection , TET spells , knee chest position



-1-your findings in this xray? Middle lobe pneumonia (lobar pneumonia)

2-what Is your choice of treatment? 3rd gen. cephalosporin + vancomycin



1-Findings is the photo ?
Dilation in ureters ,dilation of renal pelvis and calyces ,mild tortuosity on the left
2-Write down 2 complications ?
Recurrent UTI
reflux nephropathy



What is the diagnosis ?
Scarlet fever
write Down 2 immunological complications ?
rheumatic fever
glomerulonephritis



1-Findings is this photo ?Cupping of distal head of radius and ulna.Widening of epiphyseal plate.

2-Investigations u need to do? Vit. D , PTH



infant on babylac formula and has diarrhea .
1-diagnosis?
Cow-milk allergy
2-treatment ?
Amino acid based formula



1-What is the diagnosis ?21-beta hydroxylase deficiency2-Investigation to confirm ur diagnosis ?measure 17- OH progesterone



Figure 576-2 Three virilized females with untreated congenital adrenal hyperplasia. All were erroneously assigned male sex at birth, and each had a normal female sex-chromosome complement. Infants A and B had the salt-wasting form and received the diagnosis early in infancy. Infant C was referred at 1 yr of age because of bilateral cryptorchidism. Notice the completely penile urethra; such complete masculinization in females with adrenal hyperplasia is rare; most of these infants have the salt-wasting form.

-diagnosis ? Hand foot mouth disease

- most causative organism coxsackievirus



Name of each skill3 years (interactive play)

mature pincer grasp (10 month)





-some pictures of children and each one of them can say number of words , and ask to write the developmental age of each one ;

- -10 single words ? -
- 18 month -
- -50 words ?
- 24 month
- -200 words ?
- 3 years

what is the developmental age for each one ?5 years4 years3 years







OSCE Pediatrics 19/2/2019

Done by : Mamoon Saleh

Essay station :

- Didn't remember it clearly but it was about glomerulonephritis lecture .
- The diagnosis was post streptoccocal GN
- They gave a KFT labs and ask to write 6 abnormalities ?
- Ask to give other investigations ?
- Gave an another more labs , that was a typical readings for the disease , and ask what to do next ?
- Finally ask to write 3 complications ?

• History :

1.5 yr old child come to the ER with abnormal movement , take a proper history , and give the definitive diagnosis .

• **Examination :** do a full respiratory exam .

• **Discussion :** cystic fibrosis open discussion .

Mini-OSCE Pediatrics 14/4/2019

Done by :

Yazan Al-amro & Kawthar Almomani



Mention two cardiac abnormalities that cause early cyanosis:

TOF, Tricuspid atresia

How do differentiate between respiratory Andy cardiac hypoxia:

Hyperoxic test



Findings in blood film:

Microcytic & Hypochromic RBCs.

How do differentiate between IDA & Minor thalasemia:

RDW index & Mentzer's index



Diagnosis: Left LMN lesion of facial nerve Findings: Incomplete left eye closure & Mouth angle deviation



Mention two needed investigation

1-To diagnose:

Urine analysis & Renal biopsy

2-DDx:

CBC (low platelets/ITP) & PT/PTT (Protein S or C deficiency)
abdominal X-ray



abdominal CT scan



Hematuria & Flank pain Mention two DDx:

Neuroblastoma & Wilm's

Mention associated syndrome:

lackwith-Wiedmann syndrome & WAGR syndrome



Mention two findings in lipstick are useful to diagnose UTI:

Leukocyte esterase & Nitites

Mention two diagnostic tests are useful to diagnose child with recurrent pneumonia and history of meconium ileus:

> -Sweat chloride -Genetics



Diagnosis :

Meconium aspiration

Initial management:

Intubation & Oxygen

Mention two complication:

ARDS & Pulmonary HTN

Growth chart shows low stature only (Normal wight and head circumference):

> -Familial -Turner



Diagnosis: -Myelomeningiocele What expect about head circumference:

-Macrocephaly



Diagnosis:

Congenital hypothyroidism

Diagnosis:

Serum TSH & T4



Describe the study of CSF:

CSF stain

Finding:

Intracellular gram – diplococci (N.Meningiditis)

Developmental assesment

- Fork use: 18 months age
 - Hearing distraction test : 6 18 months
 - Head lag: younger than 3 months
 - Scribbling: 14 months age



Describe:

Skin glass test

Use:

differentiate between blanching/nonblanching purpura

Nonblanching purpura:

Meningiococcemia & Thrombocytopenia

OSCE Pediatrics 14/4/2019

Done by :

Yazan Al-amro & Kawthar Almomani

Essay station :

Patient with gonococcal meningitis presented by intractable seizure and not respond to antiepileptic drug

-Cause of seizure :Hyponatremia (Electrolytes showing this)-Diagnosis:Waterhouse–Friderichsen syndrome-Cause:Meningiococcemia-Initial management:Antibiotics/3rd G cephalosporin, CS, FFP-Treatment prophylaxis of contact of age less than 18 year:Rifampin-Types of vaccine for organism:Polysaccharide based and Protein based-Complication:Seizure, Hearing loss, Mental retardation

• <u>History</u>: Jaundice of 1 day age newborn

• **Examination :** CVS exam

Discussion :

-6 years old child with one week fever and limbing:

*JIA (Uveitis in oligoarthritis type and diagnosed by slit lamp exam, Hepatospleenomegaly & generlised LAD in systemic type)
*HSP (Recent URTI or vaccine or drug & Hematuria)
*Acute rheumatic fever (Migratory , Murmur, Rash)
*Leukemia (Weight loss, Anorexia, Radiation)
*Septic arthritis
*Perth's disease (Recurrent painless limbing)
*Brucellosis (History of dairy product consumption, Family member)
*Hemophilia (Family history, Bleeding)
*Septicemia (Untreated infection)
Also analyze the CC, ask about trauma, contact, drug history

CBC, ESR, CRP, PT/PTT, Urine analysis, Blood culture, X-ray

Pediatrics mini-OSCE sixth year 2019

Done By :

Noor Daher Alhijjaj

Q1 (long case)

•What do you see in this x-ray?

Double bubble sign

•What is the diagnosis ?

Deudenal atresia

•What other GI pathologies May be seen in this patient?

Annular pancreas, imperforate anus

•How to confirm dx ?

Karyotyping

•If this patient presented 3 weeks later with cyanosis and shortness of breath , what is the cause ?

AV canal "endocardiac cushion defect"

•After years this patient came with thrombocytopenia and limb pain , what is the diagnosis ?

Leukemia



Q 2 long case

Typical senario of CKD Patient came with high creatinine and high phosphate and other lab findings, What is the diagnosis ? Chronic kidney disease Mention 2 complications ? Anemia, Renal Bone Mineral Metabolism Mention importanat investigations that must be done in this patient ? Blood pressure measurement, echocardiography •How to treat the bone problem ? phosphate binders **Restriction of phosphate intake** Give active Vit D

•What is the abnormality in this chart ? Short stature

•If this girls mother had her first period at 14 years , what is the cause of her problem ? Constitutional delay

•How to confirm diagnosis ?

By wrist x-ray

•What is the treatment ?

No treatment needed , just wait



Table with blood pressure readings for child , and the growth chart , the question was to say what grade of HTN this patient have

•What is the diagnosis ? Pneumatocele •What is the cause ? Staph aureus •What is the best treatment ? Vancomycin





A pregnant complained of polyhydramnios and had this baby , what is the diagnosis ? Potter sequence (it was not mentioned in ped lectures , but they answered it from Obstetric lec)



Mention 3 abnormalities in this pic ?
Wepped neck , wide spaced nipples , wide carrying angle
What future problems will this female have ?
Short stature , infertility
What is the karyotyping
45 XO



```
•Pic of DMSA scan.
•What do you see ?
DMSA scan of one kidney
•Describe findings in this scan?
???
•What is the cause ?
Reflux nephropathy
•Other tests to do?
Biopsy, KFT
•If patient presented with hyperkalemia , how to manage ?
```

Pediatrics mini-OSCE 5th year 6/8/2019

- Whats your Dx ?
- Rosary beads sign (rickets)



Neonate with these findings and non-vigorous , what is the first step in management? الاجابة من السلايد

If the baby is not vigorous

•(defined as depressed respiratory effort, poor muscle tone, and/or heart rate < 100 bpm) \rightarrow

Use direct laryngoscopy, intubate, and suction the trachea immediately after delivery. Suction for no longer than 5 seconds. Suction befor his first breath

- → If no meconium is retrieved, do not repeat intubation and suction.
- \rightarrow If meconium is retrieved and no **bradycardia** hr <100 is present, > reintubate and suction.
- → If the heart rate is low, administer positive pressure ventilation and consider suctioning again later.





Give two DDx :

Kawasaki, scarlett fever



- 3 pictures for developmental assessment
 - All the cest on Palm .. 6 months



• 4 months.. (reach out)



• 18 months.. (spone and symbolic play)



Number of words	age
10 words	??
50 words	??

Fill boxes with "??"

10 words \rightarrow 18 months 50 words \rightarrow 2 years

1- What are the findings in this DMSA scan image?

Severe left kidney scarring which is progressive over the periode between 17 months and 9 years

2- mention 2 possible complications?

ESRD, acidosis , volume overload, elec. Disturbance ..etc (not sure)



Image taken from behind So left is left and right is right • 1- What is the deficient enzyme in this male patient?

17 hydroxylase or 3-beta-hydroxysteroid dehydrogenase deficiency

2- What is the classical presentation for congenital adrenal hyperplasia?

Salt wasting and ambaguas gentalia and hypoglycemia





1- Name 2 facial characters?

Hypertelorism and epicanthic fold

2- What is the most specific cardiac abnormality?

AV canal (endocardial cushion)







Mention 2 Acute complications of this disease?

Bleeding Acute tumor lysis syndrome Thrombosis Serious infection


What is the management in this case?

Diaphragmatic hernia Intubation + O2



1- Diagnosis? Cows milk allergy

2- Treatment ? Amino acid based formula





Patient with HX of abdominal pain and vomiting with highly elevated AST and ALT

1- what is the best investigation you would order? Anti-HAV IgM Antibodies

2- what you would like to do for his 6 years brother vaccination

3- what you would like to do for his 8 months brother IVIG and vaccination



1- wheezy chest low grade fever and x ray mention 2 indications for admission?

hypoxia, inability to take oral feedings, apnea, extreme tachypnea



3 weeks old neonate patient presented with cyanosis

What is the finding in this CXR? Cardiomegally

mention 2 heart abnormalities ? TA , TAPVR



6 years old patient presented with this CXR

1- Give the most common organism causing this finding?

S.Aureus S.pneumonia

2- whats your management?
3rd generation cephalosporin (Ceftriaxone) and vancomycin



7 years old patient, previously free , 2 days ago he started to complain of facial bluffness and hematuria

Investigation :

Many investigation were provided but the most important are: Elevated Cr and Urea RBC cast ,

1- what is the most likely Dx ?

GN

- (بأي شكل اخذ علامة السؤال GN الي كتب)
- 2- if there was a rash (as in image) whats the Dx ? HSP
- 3- what do you expect the level of complement in this pt.? Normal c3 and c4
- 4- next best investigation if protein is +2 in dipstick?
- 24-hour urine collection OR renal biopsy
- لكن الي جاوب اي وحدة من Urine collection is the best answer الاجابتين اخذ العلامة
- 5- mention 3 acute complications?

Fluid overload, uremia, acidosis, electrolyte disturbance ... etc

- 6- mention 2 management in hyperkalemia?
- Glucose + insulin , Beta agonist, calcium gluconate, bicarbonate
- 7- give 2 prognostic factors
- Proteinurea, HTN, crescents on biopsy

#Long case



Pediatrics OSCE 5th year 7/8/2019

History:

- 6 years old patient presented with fever and skin rash , take a relevant Hx
- (((التشخيص كان))) : Kawasaki Investigation ? : echocardiography to rule out coronary aneurysm)))

Examination :

Asses dehydration for a child with vomiting and diarrhea from 2 days?
 Sign, symptoms, vitals and degree of dehydration
 urine output وال mental status(place person time)

Disscusion:

• Neonate presented with seizure and cyanosis his blood was sugar 25 NICU...

1- what is your 1st step in management? ABC

- A: Airway \rightarrow oral airway {mouth piece}
- B: Breathing \rightarrow oxygenation
- C: Cannulation \rightarrow two large bore IV cannulas

2- how to correct his blood sugar?!

الاجابة حرفية وبأدق التفاصيل من سلايد infant of diabetic mother جميع الخطوات بالتفصيل من البداية حتى subtotal pancreatectomy

MINI-OSCE Of Pediatric 15-16/9/2019

Done By : Amr Mohammed Al-Khattab

Dr. Amjad Altarawneh



1- Your DDx

Left clavicle fracture

- 2- Mention 2 complications
- Erbs palsy (C5,C6) , klumpke palsy (C7,C8,T1)



1- Your DDx

Intraventricular hemorrhage

2- Mention 2 complications double diplagia spastic CP , seizure

Dr. Salma Ajarmeh



1- Mention 2 causes

Nerogenic bladder, Posterior urethral valve

2- what is the next investigation

VCUG



Hematuria and Hx of URTI 2 weeks ago

- 1- Mention 2 investigations to confirm Yr Dx
- C3 complement level , anti-DNAase
- 2- Mention 2 acute complications

HTN, fluid overload, hyperkalemia, hyponatremia, acidosis

Dr. Omar Nafi



1-What is the name of this test?

Object Permanence

2- what is the estimated age ?

9 months



If this baby can stoop down to pick up an object what is the estimated age ? 18 months تخيل صورة رسمة لطفل راسم فيها واحد وسال:

1- Estimated age according to paint

2- The formula that you estimate the age by it
3+n/4
n= عدد الاجزاء اللي رسمها الطفل



If the baby take this vaccine , what is the estimated age ? 9 months

Dr. Haitham



1- what is the abnormality ?

Short stature

2- Mention 2 DDx

Celiac disease , Hypothyroidism





What is the type and indication for each formula?

- A : Anti-regurgitation (starch based), GERD
- B: Amino acid based , cows milk protein allergy

Dr. Tariq



1- What is the type of insulin?

Long acting

2- what is the duration of action ?

24 hours

Dr. Randa



1- mention 2 differential DDx
Neuroblastoma , wilims tumor
2- one investigation to confirm the DDx
VMA



1- Mention 2 complications

Pneumonia, Encephalitis

2- At which age should receive the vaccine

9 months

Dr. Abed Alrahman



1- what is yr DDx Croup

2- Mention 2 line of management

Nebulized racemic epinephrine ,oral corticosteroids , helium-oxygen mixture



1- Your DDx

TOF

2- Mention 2 line of management

Knee-chest position (squatting), morphine, phenylephrine, Oxygen

Dr. Lina



1- What is the X-ray finding مش نفس الصورة بس انه كان فيها

Cavitation + widening of mediastinum

و الدكتورة بدها TB

2- Mention 2 DDx

TB, pneumatocele


15 months old patient with history of vomiting and fever WBC = 22 , Platelets = 298 , RBC = 4.5 , Na = 136 , K = 3.6 , glucose = 95 urine :

```
RBC = 2-4, protein = +1, PH = 5.5
```

CSF :

```
protein = 110 , glucose = 25 , WBC = 1000 ( 90 % neutrophile )
```

1- What is your DDx

Bacterial meningitis

- 2- The most common 3 organism
- Strep. pneumonia
- N. meningitidis
- H . Influenza
- 3- if the culture show gram + diplococci , what is the organism ?
- Strep. pneumonia

4- whats your treatment

3ed generation cephalosporine + vancomycin

Corticosteroid (dexamethasone)

5- tow organisms prevented by vaccination

H.Influenza vaccine , pneumococcal vaccine

6- If you repeat KFT and the Na become 127 , what is your DDx ? SIADH

7- Mention 3 long term complication deafness , other cranial nerve deficits , cerebral infarction ,

recurrent seizures or mental retardation

OSCE

Station 1 History

HX for baby 3 hour after delivery develop poor feeding and irritability and lethargy

(neonatal sepsis)

Station 2 Examination Ex the patient for respiratory system Station 3 Discussion Fever , skin rash , arthritis (Rheumatic fever)

Pediatric osce / mini-osce group 2

Done by: Mahmoud M. younis

History station

• One year male patient suffer from wet cough and fever for two days (Dx : CF)

Discussion station

• Recurrent vomiting in neonate (GERD)

1-risk factor >>neurological imairment....Eosophafeal atrasea...Asthma ...Prematurity obesity

2-complication

Recurrent pneumonia..asthma....

3-differnce between it and benign condition

- Irritability
- Not gaining weight (FTT)
- Associated complication
- Vomitus may be bloody
- Family history
- Predisposing factor (Secondary)
- Not improving with time

Examination station

• Anemia (Hb = 6)

G6PDالحالة كانت



- RBC cast/protein +2/creatinine 1.7
- 1-if patient come with this scenario after 2W of URTI, your Dx :
- Post streptococcal GN
- 2-life threatening complication in case
- Hypertision encaphalopathy
- 3-two modalities for treatment of hyperkalemia
- Ca gluconate
- Insuline and glucose
- 4-two modalities for treatment of HTN in this case

CCB, duiretics

5-if this patient come after 4 W with protein +4 /edema , your next step

24h urine collection if nephrotic proteinuria do kidney biopsy



- 1- finding : cavity (pneumatocele)
- 2-if pateint come with night sweat and weight loss for one month .your Dx : Tb
- 3-gold standard Dx : culture



- **mention two complication of this condition (meconium aspiration)
- ARDS & Pulmonary HTN



- 1-What's the CXR finding?
- Egg-on-string.
- 2-What's the cause of his cyanosis?
- TGA (Two parallel circuits)



- **mention tow Differential diagnosis of this condition
- 1. Scarlet fever
- 2. kawasaki



- 177 Erb's palsy.
- 1-name of injury
- Brachial plexus injury...klumpky palsy
- 2-two risk factors :
- Shoulder dystocia
- Prolonged labor
- macrosomia

Growth chart

- Your assessment :
 - short stature
- 2- give two causes of this case
- Celiac disease.
- familial (genetic)



- Two complication of this case
- Recurrent UTI reflux nephropathy

- History of painless abdomen mass and hematuria
- Your dignosis

((wilm tumor))

• What you see in his eyes aniridia

Developmet assesment

- 1-raise chest on palms...6 months
- 2-reah out toys..4 months
- 2-responsive smile ...6 weeks
- 4-identify body parts..18months



• 1-your Dx :

turner syndrome

• 2-mention two complication

Coarctation of the aorta

horseshoe kidney.



- Mention 3 findings in this test suggest for UTI
- Leukocyte esterase & Nitites

Pediatric mini-osce and osce 18-19/2/2020

Done by: Rayan Nihad



What's the age of each skill? Both of them at 3 years







How many words can he say

- 10 words



Q3 What age? 10 months





What age ? 4 years

What's his pencil skill ? Draws a cross



Q5

What's the vaccines taken at this developmental age?

MMR2

vit A 200 000 U

Booster PaTd

Booster OPV



Holds a spoon and gets food safely to mouth



2 months baby with head circumference 42 (growth chart)

- 1- blot HC on growth chart? Above 95 percentile
- 2-name of the sign? Macrocephaly

3- causes?

Familial , TORCH INFECTION , Hydrocephalus (increase ICP)

Q7

When the vaccine is given for this rash? 10 -12 -18 months(measles)



Q8

- 5 days old baby with late passage of meconium and no gas in the rectum
- 1- what caused these signs
- Hirschsbrung disease
- 2- other 2 complications with this syndrome ?
- Duodenal atresia, VSD



Q9 1-What is the diagnosis ? 21-beta hydroxylase deficiency 2-Investigation to confirm your diagnosis ? measure 17- OH progesterone



Figure 576-2 Three virilized females with untreated congenital adrenal hyperplasia. All were erroneously assigned male sex at birth, and each had a normal female sex-chromosome complement. Infants A and B had the salt-wasting form and received the diagnosis early in infancy. Infant C was referred at 1 yr of age because of bilateral cryptorchidism. Notice the completely penile urethra; such complete masculinization in females with adrenal hyperplasia is rare; most of these infants have the salt-wasting form.

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Q10 Describe the finding?

Cupping in the distal ends for radius and ulna

2- lab test?

Serum ca, vit D, PTH



Q11

year male acute illness sever sepsis

Shistocytes pic

Lab findings suggest the picture

Prolonged pt ptt INR

increased D dimer

Thrombocytopenia





Vomiting diarrhea seizure with stool test

Brown, blood, mucus

No cyst or trophozoites

1-What is the Organism: shigella •

2-Tx of choice: ceftriaxone •



A pic of left lower lobe pneumonia with increased vascular markings Finds on CXR (written above) Most common bacteria for this case ? Staph aureus (not sure)

Q14 questions in the next slide



Q14 CONT

Protein 90 WBC 5000 neutrophil 90% glucose 40

- Diagnosis : bacterial meningitis
- Name of test: gram stain
- Organism . N. Meningitides
- Hypotension we give normal saline not respond why?
- Waterhouse fridrechson syndrome---> adrenal hemorrhage >> no cortisol
- Management: iv fluid cortisol dopamine
- Antibiotics of choice
- 3rd generation cephalosporin and vancomycin
- Seizure not responding to antiepileptic:

SIADH as a complication

OSCE

Station 1 History (pyelonephritis) Station 2 Examination (Dehydration / gastroenteritis) Station 3 Discussion (celiac disease)

