2013年02月22日 中華民國放射線醫學會 住院醫師閱片測驗-答案

出題醫院 台中榮民總醫院放射線部 Q01-1 性別: Male 年齡: 62 year-old 主訴或重要病史或物理檢查: Sudden right side limbs clumsine





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# ANS: Left type type IIA +IIB dural AV fistula, s/p ONYX embolization with successful obliteration of the fistula.

### Q02-1 性別:female 年齡:26 主訴或重要病史或物理檢查: Frequent seizure attack







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# ANS: Heterotopic gray matter and callosal dysgenesis



性別:Female

年齡:5 day-old, Gestation period 40+4 weeks

主訴或重要病史或物理檢查: Much oral meconium, general cyanosis and apnea after birth





# ANS: Hypoxic ischemic encephalopathy, full-term

## Hypoxic ischemic encephalopathy

- Brain maturation
- Hypoperfusion: severity, duration
- Mild to moderate hypoperfusion: blood redistribution, damage of watershed regions
- Severe hypoperfusion: high metabolic regions injury.
  - Preterm: thalamus, brainstem
  - Fullterm: lateral thalamus, globus pallidus, posterior putamina, hippocampi, brainstem, sensorimotor cortex (perirolandic cortex)

### Q04

性別: Male 年齡: 10 y/o 主訴或重要病史或物理檢查: General lethargy for 16 days, frequent vomiting, headache, fever up to 38C According to mother's statement, patient was used to kissed 蛞 蝓





### ANS: Eosinophic meningoencephalitis Caused by Angiostrongylus cantonensis

- Angiostrongylus cantonensis, also known as the rat lungworm, is prevalent in the Pacific Islands and southeast Asia and is the most common cause of eosinophilic meningitis in humans.
- Rats serve as the definitive host of the nematode.
- Humans are infected by ingestion of freshwater and terrestrial snails and slugs, or transport hosts, such as freshwater prawns, frogs, fish.
- Taiwan has recorded hundreds of cases of eosinophilic meningitis. Most of the cases occurred among children.
- There was usually a history of eating or playing with snails or slugs.

# MRI

- Multiple enhancing nodules in the brain and linear enhancement in the leptomeninges, accompanied by stick-shaped enhancement.
- Leptomeningeal enhancement, ventriculomegaly and abnormal enhancement within the cerebral and cerebellar hemisphere.
- Significant correlation between the severity of headache, cerebrospinal fluid (CSF) pleocytosis, and CSF and blood eosinophilia with MRI signal intensity in T1-weighted imaging.

#### Q05 性別:male 年齡:22 主訴或重要病史或物理檢查:<u>Hearing deteriorated recently</u>





# ANS: Mondini syndrome

Q06-1 性別:Female 年齡:35 y/o 主訴或重要病史或物理檢查:Severe headache for 2 weeks

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FLAIR

T1/C







### Q06-2

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# ANS: Gliomatosis cerebri

#### Q07 性別:Male 年齡:80 y/o 主訴或重要病史或物理檢查: Involuntary movement of left side limbs for 2 weeks





# ANS: hyperglycemic hemichorea

# hemichorea-hemiballism

- Since hemiballism often evolves into hemichorea: hemichorea-hemiballism (HCHB)
- The involuntary movements of chorea consist of random and fast jerking motions in distal parts of the limbs, whereas those of ballism consist of larger-amplitude, random, and violent flinging or kicking, mainly in the proximal joints.
- The most common cause : vascular lesion, hyperglycemia
- HCHB that accompanies hyperglycemia may exhibit a hyperintense putamen on T1WI and transient presence of high density in the corresponding regions on CT scans.



A, hyperintensities in the right caudate nucleus, the putamen, globus pallidus (arrow).B, Sagittal section shows hyperintensities extending from the anteroventral part of the right lentiform nucleus to the anterior part of the midbrain (arrow).C, Axial section at the level of the midbrain shows hyperintensity in medial part of the

C, Axial section at the level of the midbrain shows hyperintensity in medial part of the right cerebral peduncle (arrow).

Q08-1 性別:Male 年齡:13 y/o 主訴或重要病史或物理檢查:Acute left eye visual decline. He had another attack five months ago









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# ANS: Rosai-Dorfman disease

# Rosai-Dorfman disease

- Sinus histiocytosis with massive lymphadenopathy (SHML), is a rare histiocytosis of unknown cause.
- Epstein-Barr virus and human herpes virus 6 have been isolated in a few patients, but no clear association has been identified.
- Extranodal involvement occurs in 43% cases, with 75% occurring in the head and neck, including upper respiratory tract (73%), Orbit (50%), and salivary glands (25%).
- The symptoms include nasal obstruction, rhinitis, and epistaxis, and clinical signs are exophthalmos and nasal polyps according to the location of the lesion

- The diagnosis is base on the pathologic findings of emperiopolesis and S-100 protein positivity.
- There is no ideal treatment, and the disease is often self-limited.
- Steroid, chemotherapy, radiotherapy, and surgery have been reported for the management of Rosai-Dorfman disease.



#### 性別:Male 年齡:16 y/o 主訴或重要病史或物理檢查:mild dizziness









### ANS: Wilson disease

# Lab

- Ceruloplasmin:0.097 (0.21-0.42)
- Urine cu: 134 (15-50)
- Blood cu: 47 (70-150)

# Wilson disease

- Wilson's disease is an autosomal-recessive disorder caused by mutation in the ATP7B gene, with resultant impairment of biliary excretion of copper
- Diagnosis:
- -- Symptomatic pt: presence of Kayser-Fleischer rings and ceruloplasmin levels of less than 20 mg/dL
- -- Asymptomatic: isolated liver disease, and lacks corneal rings, the coexistence of a hepatic copper concentration of more than 250 mg/g of dry weight and a low serum ceruloplasmin level





# Wilson disease

- Bilateral symmetric involvement of the putamen, caudate nuclei, thalamus, globus pallidus, dentate nucleus, pons, substantia nigra, periaqueductal gray matter, tectum, and red nucleus can be seen.
- Subcortical and centrum semiovale white matter involvement may also be seen.
- Hypointensity on T2-weighted images may be seen sometimes, secondary to copper deposition or iron deposition.
- Restricted diffusion : early in the disease process due to cytotoxic edem, or inflammation due to excessive copper deposition. However, this restricted diffusion is not seen in chronic cases. which are characterized by necrosis, spongiform degeneration, and demyelination.
## Wilson disease

- The lentiform nuclei are involved most often, followed by the thalami, pons, midbrain, superior and middle cerebellar peduncles, and cerebellar nuclei.
- Cause: edema, gliosis, demyelinization, neuronal necrosis, or cystic degeneration
- Some of the lesions showed reversible changes following copper chelating Tx



T2-weighted axial MR image shows bilateral symmetric high signal intensity in the putamen (*black arrows*) and caudate nuclei (*white arrows*). Slightly increased signal intensity in both thalami (*black arrowheads*) and globus pallidi (*white arrowheads*) is also noted.

## Wilson disease

- Dark SI in the globus pallidus, putamen, or caudate nucleus, surrounded by peripheral high signal intensity on T2WI
- Cause: increased iron deposition in the areas with accumulation of copper.
- The reversible process of copper metabolism can be a possible explanation for the reversibility of hypointensity after copper chelating therapy,



T2-weighted axial MR image shows bilateral high signal intensity with central dark signal intensity in the putamen (*arrows*) and caudate nuclei (*arrowheads*).

AJNR Am J Neuroradiol 27: 1373-1378

## Wilson disease

- bilateral T1 hyperintensity in the GP, putamen, midbrain, and pons, without signal-intensity abnormality on T2WI.
- result from the toxic effect of copper overload in the brain and indirectly from liver failure as a result of copper intoxication.



T1-weighted axial MR images show bilateral increased signal intensity in the globus pallidus (*arrows*) and midbrain (*arrowhead*).

AJNR Am J Neuroradiol 27: 1373-1378

Q10 性別: Male 年齡: 4 m/o 主訴或重要病史或物理檢查: Poor head control was noted. Developmental delay with obvious head lag





# ANS: Leigh syndrome

- Bilateral symmetric hyperintense on T2WI over bilateral dorsal putamina and periaqueductal gray matter.
- Thalami, dentate nuclei.
- Progressive neurodegeneration leading to respiratory failure and death in childhood



Symmetrical involvement of head of caudate nuclei, putamina and dentate nuclei.

Symmetrical hyperintense lesions involved thalamic posteromedial ventral nuclei, globus pallidi and putamina.

Symmetrical involvement of reticular formation of midbrain (thin arrow in B), subthalamic nuclei (thick arrow in B), substantia nigra (thick arrow in C), dorsal midbrain (thin arrow in C) central tegmental tracts (thin arrow in D) and cerebellar nuclei region (thick arrow in D).



Q11-1

性別:male 年齡:55 主訴或重要病史或物理檢查:history of infective endocarditis











# ANS: septic emboli over bilateral lung

Q12-1 性別:female 年齡:19 主訴或重要病史或物理檢查:Asymmetric breasts



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## Q12-2

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# ANS: Poland syndrome

Q13-1 性別: Male 年齡: 68 主訴或重要病史或物理檢查: Myalgia for 40 days and fever with chillness











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# ANS: Wegener's granulomatosis

Q14-1 性別: Male 年齡: 34 主訴或重要病史或物理檢查: Hemoptysis for several months





# ANS: Pulmonary TB with Rasmussen aneurysm

Q15 性別:female 年齡:32 主訴或重要病史或物理檢查:Abdominal pain





# ANS: Neurilemmoma (Schwannoma)

Q16-1 性別:Female 年齡:64 主訴或重要病史或物理檢查: Incidentally found to have a liver tumor by sono in LMD 2 weeks ago. No abdominal discomfort

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Q16-2

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# ANS: Cholangiocarcinoma

#### Q17 性別:Male 年齡:75 主訴或重要病史或物理檢查:Abdominal fullness





# ANS: Icteric type HCC



性別:female 年齡:1 day 主訴或重要病史或物理檢查:vomiting





# ANS: Jejunal stenosis

Q19 性別:female 年齡: 56 主訴或重要病史或物理檢查:Hematuria with left flank pain





# ANS: Nutcracker syndrome, causing pelvic congestion syndrome






# ANS: Encapsulated sclerosing peritonitis

Q21 性別:Male 年齡:65 y/o 主訴或重要病史或物理檢查:Acute onset abdominal pain









## ANS: Ischemic enteritis with portal vein air

#### Q22 性別:Female 年齡:34 y/o 主訴或重要病史或物理檢查:LUQ abdominal pain off and on for 2 months





# ANS: Solid and papillary epithelial neoplasm (SPEN)

#### Q23-1 Female 14 year-old Right knee pain for 2 months

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#### Q23-2

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T2 fat-sat



T1 fat-sat Gd+





# ANS: Pigmented villonodular synovitis

#### Male 38 year-old Q24-1 Back pain after slip down



一張





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### ANS: Transvertebral fracture in AS

Q25 Male 45 year-old Left knee in recent months. Tenderness over knee joint line, Genu varus. No trauma. DM for years



### ANS: DM neuroarthropathy

#### Q26 Male 20 year-old Right ankle pain. Had gouty arthritis history since high school



#### **ANS:** Talocalcaneal coalition

#### Q27 Female 30 year-old Bilateral heel pain for 1 year



### ANS: Intra-osseous lipoma





What's your diagnosis ? 性別:Male 年齡:18





### ANS: Intra-osseous gout

#### Q29 What's your diagnosis?

- 性別:Male
- 年齡:27





### ANS: Hemophilic arthropathy, right knee

**Q30** Male 2 year-old Deformity of left lower leg for 6+ months. Unable to walk. Pain over left lower leg for weeks





### ANS: Congenital pseudoarthrosis

Q31-1性別: Male年齡: 55主訴或重要病史或物理檢查: RCC of left kidney about<br/>six years ago post left nephrectomy



## Q31-2 本題有二張投影片,第二張







## ANS: Metastatic renal cell carcinoma in the right kidney and pancreas, (Gallstone)

## Q32-1性別:Male年齡:52主訴或重要病史或物理檢查:Painless grosshematuria off and on for one month

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## ANS: Left renal UC with lymph node metastasis

#### Q33 性別:Female 年齡:30 主訴或重要病史或物理檢查: Supra- renal lesion was noted at LMD





## ANS: Retroperitoneal mature teratoma

#### Q34 性別:Male 年齡:45 主訴或重要病史或物理檢查: Incidental found left adrenal tumor on CT scan





T2 with fat saturation







## ANS: Cortical adenoma of left adrenal gland

Q35-1 性別:male 年齡:23 y/o 主訴或重要病史或物理檢查:chest tightness and dyspnea on exertion for months




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Ans: Kawasaki disease with aneurysm formation of left main coronary artery Q36-1 性別:male 年齡:1 y/o 主訴或重要病史或物理檢查:mild tachypnea since birth



Q36-2

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### Q36-3

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## Ans: Hypoplastic Left Heart Syndrome (HLHS) with ASD



#### Q37-2





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#### View form back of heart

# Ans: PAPVR (partial anomalous pulmonary venous return)



性別:female 年齡:1m/o 主訴或重要病史或物理檢查:tachypnea,mild suprasternal retraction





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# ANS: TAPVR (Total anomalous pulmonary venous return)

Q39-1 · 52 years old female 超音波所見病灶為何?

#### • Ca of right breast post OP



TIS<0.4 MI=0.8 A0=100%

TIS=2.2 MI=1.1 A0=100%





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# ANS: Rib metastasis, left first (from breast Ca)



#### 55 years old male Routine physical examination







## ANS: Horseshoe kidney

性別:Female 年龄:57

主訴或重要病史或物理檢查: RLQ pain, off and on for more than 5 days



CN0 6cm DR54 G 50

CN0 6cm DR54 G 50

Q41









# ANS: Acute appendicitis with rupture

#### Q42-1 性別:Female 年齡:87 主訴或重要病史或物理檢查:Hematuria for several days



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## ANS: Synchronous urothelial cell carcinoma in the right renal pelvis and urinary bladder





- 1 d/o male infant, birth hx: GA 40+2wk, BBW 2506gm
- Prenatal exams showed pleural effusion since GA 21wk
- Respiratory distress with cyanosis noted 3 hours after birth

#### Q43-2

#### 本題有二張投影片,第二張









3:39:01 pm #43 8L5 7.0MHz 8L5 CHEST 55mm **NTHI General** 

78dB S1/+1/3/4 Gain= 0dB ▲=2



Dist	=	-1 2.61cm	
		-2 2.99cm	

	3:40:09 pm
8L5	#76
7.0MHz	55mm
8L5 CH	EST
Genera	l









## ANS:

• OP finding: congenital lung cyst, LLL

 Pathology: Congenital cystic adenomatoid malformation, type 1



1 d/o female newborn, presented with bradycardia and cyanosis after birth



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 After ambu-bagging and ventilation for 1 hour





## ANS: Congenital diaphragmatic hernia, right side





- A one day old baby boy
- GA 39Wk BW
  3286g
- NSD with Apgar score 3'->7'.



## ANS: Gastroschisis

Q46-1 性別:male 年齡:3 主訴或重要病史或物理檢查:Palpable abdominal mass










## ANS: Hepatoblastoma



- Q1: BI-RADS ?
- Q2: Diagnosis ?



Rt Breast L 3 /SAI



T



- Ans (1): BIRADS category 5 or 4c
- Ans (2): Breast cancer

Q48 性別:Female 年齡:59 year-old 主訴或重要病史或物理檢查:palpable right breast nodule for 1 year

## Q1: BI-RADS ? Q2: Diagnosis ?

Pu

Т





- Ans (1): BIRADS category 5 or 4c
- Ans (2): Breast cancer

Q49-1 性別:Female 年齡:42 year-old 主訴或重要病史或物理檢查:Breast mass noted for 1 week



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L-MLO



Q2: What is the diagnosis?



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- Ans (1): BIRADS category 0, 5 or 4C
- Ans (2): Breast cancer with axillary metastatic adenopathy

## Q50-1 Q1: On which side is/are salient abnormality located? Q2: BI-RADS ?

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Rt breast Lt breast 59 Y/O female. No previous history of operation of bil. Breasts





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- Ans (1): Bilateral
- Ans (2): **BI-RADS 5**