

INI CET Paediatrics

Sample Paper – 1

Duration: 14 Minutes

Maximum Marks: 15

Instructions

- This paper contains **15** Multiple Choice Questions (Single Best Response), modelled on the Paediatrics content of **INI CET** (Institutes of National Importance Combined Entrance Test) conducted by AI-IMS New Delhi.
- The **15-question** length matches the number of Paediatrics items you actually meet in the real 200-question paper.
- Each correct answer carries **+1 mark**. There is **negative marking of one-third (0.33) mark** for every incorrect answer; unattempted questions carry no penalty.
- Every question has **exactly one best answer**. Choose the single most appropriate option.
- Attempt this practice paper in one timed sitting of about **14 minutes**, the pace the real computer-based test demands.

Neonatology

- Q1.** The APGAR score used to assess a newborn at 1 and 5 minutes is built from five clinical signs. Which of the following is **NOT** one of its five components?
- (A) Heart rate
(B) Muscle tone
(C) Birth weight
(D) Reflex irritability (response to stimulation)
- Q2.** In a healthy term newborn, physiological jaundice characteristically:
- (A) Appears after 24 hours of life and peaks around day 3 to 5



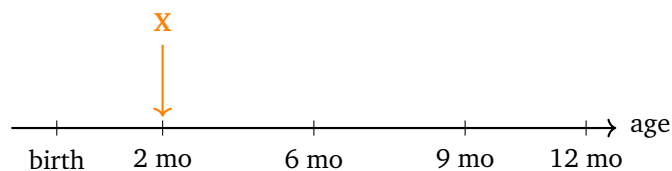
- (B) Appears within the first 24 hours of life
- (C) Appears only after the first week of life
- (D) Never occurs in a term infant

Q3. Respiratory distress syndrome (hyaline membrane disease) in a preterm newborn is caused by deficiency of:

- (A) Alpha-1 antitrypsin
- (B) Fetal haemoglobin
- (C) Pulmonary surfactant
- (D) Vitamin K

Growth, Development and Nutrition

Q4. On the developmental timeline below, the arrow **X** marks the age at which a normal infant first shows a *social smile* in response to a caregiver. That age is:



- (A) About 6 to 8 weeks (around 2 months)
- (B) About 6 months
- (C) About 9 months
- (D) About 12 months

Q5. A 2-year-old presents with bilateral pitting pedal oedema, a distended abdomen, sparse depigmented hair and a low serum albumin. This picture is the hallmark of:

- (A) Marasmus
- (B) Kwashiorkor
- (C) Nutritional rickets



(D) Infantile scurvy

Q6. Bitot spots on the conjunctiva together with night blindness in a mal-nourished child indicate deficiency of:

(A) Vitamin C

(B) Vitamin D

(C) Vitamin B12

(D) Vitamin A

Immunisation and Infectious Disease

Q7. Under India's Universal Immunisation Programme (UIP), the BCG vaccine is scheduled to be given:

(A) At 6 weeks of age

(B) At birth, or as early as possible up to one year of age

(C) At 9 months of age

(D) At 5 years of age

Q8. Koplik spots, small bluish-white lesions on the buccal mucosa appearing before the rash, are pathognomonic of:

(A) Chickenpox

(B) Rubella

(C) Mumps

(D) Measles

Respiratory and Cardiovascular Paediatrics

Q9. The single most common causative organism of acute bronchiolitis in infants under 2 years is:

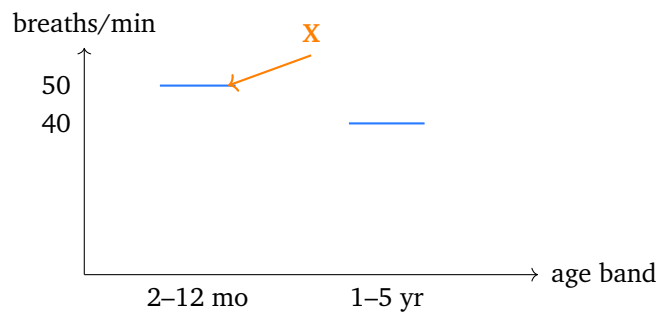
(A) Streptococcus pneumoniae

(B) Influenza virus

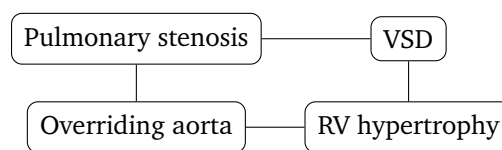


- (C) Respiratory syncytial virus (RSV)
- (D) Staphylococcus aureus

Q10. The chart below plots the WHO IMNCI cut-off for *fast breathing*. For an infant aged 2 to 12 months, the arrow **X** marks the threshold respiratory rate that counts as fast breathing, namely:



- (A) 50 breaths per minute or more
 - (B) 30 breaths per minute or more
 - (C) 20 breaths per minute or more
 - (D) 70 breaths per minute or more
- Q11.** The schematic shows the four features that together define one congenital lesion: pulmonary stenosis, a ventricular septal defect, an overriding aorta and right ventricular hypertrophy. This most common cyanotic congenital heart disease of childhood is:



- (A) Ventricular septal defect alone
- (B) Patent ductus arteriosus
- (C) Tetralogy of Fallot
- (D) Atrial septal defect

Haematology, Genetics and Systemic Disease

- Q12.** A 2-year-old has severe microcytic hypochromic anaemia needing repeated transfusions, hepatosplenomegaly and a 'hair-on-end' appearance of the skull on X-ray. The most likely diagnosis is:
- (A) Iron deficiency anaemia
 - (B) Sickle cell trait
 - (C) G6PD deficiency
 - (D) Beta-thalassaemia major
- Q13.** Trisomy 21 (Down syndrome) is most commonly associated with which congenital heart defect?
- (A) Atrioventricular septal defect (endocardial cushion defect)
 - (B) Coarctation of the aorta
 - (C) Bicuspid aortic valve
 - (D) Ebstein anomaly

Emergency, Fluids and Electrolytes

- Q14.** The current WHO low-osmolality oral rehydration solution (ORS) used to treat childhood dehydration has a total osmolality of approximately:
- (A) 311 mOsm/L
 - (B) 245 mOsm/L
 - (C) 90 mOsm/L
 - (D) 400 mOsm/L
- Q15.** During single-rescuer cardiopulmonary resuscitation of a child, the recommended chest compression to ventilation ratio is:
- (A) 15 compressions to 2 breaths
 - (B) 30 compressions to 2 breaths
 - (C) 5 compressions to 1 breath
 - (D) 3 compressions to 1 breath



Detailed Solutions**Q1.****Solution**

Concept – the five components of the APGAR score: The APGAR score is a rapid bedside assessment of a newborn's condition at 1 and 5 minutes.

Step 1 – list the five signs: The mnemonic APGAR stands for Appearance (colour), Pulse (heart rate), Grimace (reflex irritability), Activity (muscle tone) and Respiration.

Step 2 – spot the odd one out: Birth weight is recorded separately at delivery but is *not* scored in APGAR.

Why the other options are wrong:

- A, heart rate: this is the 'Pulse' component (0, less than 100, or 100 or more).
- B, muscle tone: this is the 'Activity' component.
- D, reflex irritability: this is the 'Grimace' component.

Final Answer: Birth weight is not part of APGAR ⇒ C

Answer: (C) [Go Back to Q1](#)

Q2.**Solution**

Concept – physiological versus pathological neonatal jaundice: Timing of onset is the key to separating benign physiological jaundice from a dangerous pathological cause.

Step 1 – recall the rule: Physiological jaundice appears **after 24 hours** of life, peaks on day 3 to 5 in a term baby, and then fades.

Step 2 – the red flag: Any jaundice appearing *within* the first 24 hours is pathological until proven otherwise (for example haemolysis from Rh or ABO incompatibility).

Why the other options are wrong:

- B, within 24 hours: this defines pathological, not physiological, jaundice.
- C, only after the first week: this suggests prolonged or pathological jaundice.
- D, never in a term infant: physiological jaundice is in fact very common in term babies.



Final Answer: Appears after 24 hours, peaks day 3 to 5 $\Rightarrow$ A

Answer: (A) [Go Back to Q2](#)

Q3.

Solution

Concept – pathogenesis of respiratory distress syndrome (RDS): RDS, or hyaline membrane disease, is the classic lung disease of prematurity.

Step 1 – identify the deficient substance: It is caused by deficiency of **pulmonary surfactant**, produced by type II pneumocytes, which normally lowers alveolar surface tension.

Step 2 – the consequence: Without surfactant the alveoli collapse at end-expiration, causing widespread atelectasis, hypoxia and grunting respiration.

Why the other options are wrong:

- A, alpha-1 antitrypsin: its deficiency causes emphysema and liver disease, not neonatal RDS.
- B, fetal haemoglobin: this is normally high in newborns and is unrelated to RDS.
- D, vitamin K: its deficiency causes haemorrhagic disease of the newborn, not RDS.

Final Answer: Pulmonary surfactant deficiency $\Rightarrow$ C

Answer: (C) [Go Back to Q3](#)

Q4.

Solution

Concept – the earliest social milestone: Developmental milestones follow a predictable age sequence, and the social smile is one of the first.

Step 1 – read the timeline: The arrow X points to the 2-month mark on the age axis.

Step 2 – match the milestone: A responsive social smile normally appears by **6 to 8 weeks** (about 2 months) of age.

Why the other options are wrong:

- B, 6 months: by now the baby sits with support and transfers objects, well



past the social smile.

- C, 9 months: the age of crawling and stranger anxiety.
- D, 12 months: the age of first words and standing, far later than the social smile.

Final Answer: About 6 to 8 weeks ⇒ A

Answer: (A) [Go Back to Q4](#)

Q5.

Solution

Concept – classifying protein-energy malnutrition: Severe malnutrition splits into two classic patterns, and oedema is the discriminator.

Step 1 – read the picture: Pitting oedema with a distended abdomen, sparse depigmented hair and hypoalbuminaemia is the hallmark of **kwashiorkor** (severe protein deficiency with relatively preserved calories).

Step 2 – contrast with marasmus: Marasmus is severe wasting from a total calorie deficit and, crucially, has *no* oedema.

Why the other options are wrong:

- A, marasmus: presents as gross wasting with an old-man face and no oedema.
- C, nutritional rickets: causes bony deformities from vitamin D deficiency, not oedema.
- D, infantile scurvy: causes bleeding gums and painful subperiosteal haemorrhages from vitamin C deficiency.

Final Answer: Kwashiorkor ⇒ B

Answer: (B) [Go Back to Q5](#)

Q6.

Solution

Concept – ocular signs of vitamin deficiency: Certain eye findings point to a single specific micronutrient lack.

Step 1 – match the signs: Bitot spots (foamy keratinised patches on the conjunctiva) plus night blindness (nyctalopia) are classic features of **vitamin A** deficiency.

Step 2 – why: Vitamin A (retinal) is essential for rhodopsin in the rods, so its



lack impairs dim-light vision first, followed by conjunctival and corneal changes (xerophthalmia).

Why the other options are wrong:

- A, vitamin C: deficiency causes scurvy (bleeding gums), not Bitot spots.
- B, vitamin D: deficiency causes rickets, not night blindness.
- C, vitamin B12: deficiency causes megaloblastic anaemia and neuropathy.

Final Answer: Vitamin A deficiency $\Rightarrow$ **D**

Answer: (D) [Go Back to Q6](#)

Q7.

Solution

Concept – timing of BCG in the UIP schedule: The Universal Immunisation Programme fixes the age at which each vaccine is due.

Step 1 – recall BCG timing: BCG is given **at birth**, or as early as possible up to one year of age.

Step 2 – the rationale: Early BCG protects the infant against severe childhood tuberculosis, especially tuberculous meningitis and miliary TB.

Why the other options are wrong:

- A, 6 weeks: this is when the first dose of OPV, pentavalent, rotavirus and PCV are due, not BCG.
- C, 9 months: this is the timing of the first measles-rubella (MR) dose.
- D, 5 years: around this age the DPT booster is given, not BCG.

Final Answer: At birth (up to one year) $\Rightarrow$ **B**

Answer: (B) [Go Back to Q7](#)

Q8.

Solution

Concept – pathognomonic sign of measles: One mucosal sign appears before the rash and clinches the diagnosis.

Step 1 – name the disease: Koplik spots, small white lesions on an erythematous buccal mucosa, are pathognomonic of **measles** (rubeola) and appear 1 to 2 days before the skin rash.



Step 2 – the sequence: Measles then produces a maculopapular rash spreading cephalocaudally, along with the ‘3 Cs’ of cough, coryza and conjunctivitis.

Why the other options are wrong:

- A, chickenpox: gives crops of vesicles at different stages, not Koplik spots.
- B, rubella: causes a milder rash with post-auricular lymphadenopathy.
- C, mumps: causes parotid swelling, not a buccal enanthem or rash.

Final Answer: Measles ⇒

Answer: (D) [Go Back to Q8](#)

Q9.

Solution

Concept – aetiology of acute bronchiolitis: Bronchiolitis is the commonest lower respiratory infection of infancy.

Step 1 – name the organism: **Respiratory syncytial virus (RSV)** is by far the commonest cause, typically in winter epidemics in babies under 2 years.

Step 2 – the clinical picture: It causes wheeze, hyperinflation and fine crepitations from inflammation and plugging of the small airways (bronchioles).

Why the other options are wrong:

- A, *Streptococcus pneumoniae*: a leading cause of bacterial pneumonia, not bronchiolitis.
- B, influenza virus: can cause bronchiolitis occasionally but is far less common than RSV.
- D, *Staphylococcus aureus*: causes severe pneumonia with pneumatoceles, not typical bronchiolitis.

Final Answer: Respiratory syncytial virus ⇒

Answer: (C) [Go Back to Q9](#)



Q10.

Solution

Concept – WHO IMNCI cut-offs for fast breathing: The threshold respiratory rate that defines pneumonia depends on the child's age.

Step 1 – read the chart: The arrow X sits at the 2-to-12-month band, whose cut-off line is drawn at 50 breaths per minute.

Step 2 – apply the value: For an infant aged 2 to 12 months, fast breathing is **50 breaths per minute or more**.

Why the other options are wrong:

- B, 30 or more: this is closer to a normal infant rate, not the cut-off.
- C, 20 or more: this is a normal adult rate, far too low for infants.
- D, 70 or more: too high; the 40-per-minute cut-off applies to the older 1-to-5-year band, and 50 to the infant band.

Final Answer: 50 breaths per minute or more ⇒ A

Answer: (A) [Go Back to Q10](#)

Q11.

Solution

Concept – Tetralogy of Fallot (TOF): Four anatomical features together define the commonest cyanotic congenital heart disease of childhood.

Step 1 – match the four features: Pulmonary stenosis, a ventricular septal defect, an overriding aorta and right ventricular hypertrophy are the tetrad of **Tetralogy of Fallot**.

Step 2 – the clinical clue: Right-to-left shunting across the VSD causes cyanotic 'tet' spells and a boot-shaped heart on chest X-ray.

Why the other options are wrong:

- A, isolated VSD: the commonest congenital heart defect overall, but acyanotic (left-to-right shunt).
- B, patent ductus arteriosus: an acyanotic shunt with a continuous machinery murmur.
- D, atrial septal defect: an acyanotic shunt with fixed splitting of the second heart sound.

Final Answer: Tetralogy of Fallot ⇒ C



Answer: (C) [Go Back to Q11](#)

Q12.

Solution

Concept – beta-thalassaemia major: A transfusion-dependent microcytic anaemia from infancy points to a haemoglobinopathy.

Step 1 – read the clues: Severe microcytic hypochromic anaemia needing repeated transfusions, hepatosplenomegaly and a ‘hair-on-end’ skull X-ray are classic for **beta-thalassaemia major**.

Step 2 – the mechanism: Absent beta-globin chains cause ineffective erythropoiesis; marrow expansion produces the hair-on-end skull and frontal bossing, and haemolysis causes the organomegaly.

Why the other options are wrong:

- A, iron deficiency: microcytic but responds to iron and does not need transfusions or cause a hair-on-end skull.
- B, sickle cell trait: a benign carrier state that is usually asymptomatic.
- C, G6PD deficiency: causes episodic haemolysis after oxidant stress, not chronic transfusion dependence.

Final Answer: Beta-thalassaemia major ⇒ D

Answer: (D) [Go Back to Q12](#)

Q13.

Solution

Concept – cardiac defects in Down syndrome: Trisomy 21 has a strong link to one particular congenital heart lesion.

Step 1 – name the defect: The commonest heart defect in Down syndrome is the **atrioventricular septal defect** (endocardial cushion defect / AV canal).

Step 2 – why it matters: Because of this strong association, every newborn with Down syndrome needs an echocardiogram to screen for it.

Why the other options are wrong:

- B, coarctation of the aorta: more typical of Turner syndrome.
- C, bicuspid aortic valve: also associated with Turner syndrome, not Down.
- D, Ebstein anomaly: linked to maternal lithium use, not trisomy 21.



Final Answer: Atrioventricular septal defect $\Rightarrow$ A

Answer: (A) [Go Back to Q13](#)

Q14.

Solution

Concept – WHO low-osmolarity ORS: The reduced-osmolarity formula replaced the older higher-osmolarity ORS.

Step 1 – recall the value: The current WHO low-osmolarity ORS has a total osmolarity of about **245 mOsm/L** (sodium 75, glucose 75 mmol/L).

Step 2 – the advantage: The lower osmolarity reduces stool output, vomiting and the need for unscheduled intravenous fluids compared with the old 311 mOsm/L solution.

Why the other options are wrong:

- A, 311 mOsm/L: this was the older, higher-osmolarity ORS, now replaced.
- C, 90 mOsm/L: this is the sodium-plus-glucose figure region, not the total osmolarity.
- D, 400 mOsm/L: hyperosmolar and never recommended, as it would worsen diarrhoea.

Final Answer: About 245 mOsm/L $\Rightarrow$ B

Answer: (B) [Go Back to Q14](#)

Q15.

Solution

Concept – paediatric CPR compression to ventilation ratio: The ratio depends on how many rescuers are present.

Step 1 – recall the single-rescuer ratio: For a lone rescuer performing CPR on a child, the ratio is **30 compressions to 2 breaths**, the same as for adults.

Step 2 – contrast: With two rescuers the paediatric ratio changes to 15:2, and 3:1 is used only in neonatal resuscitation.

Why the other options are wrong:

- A, 15:2: this is the *two-rescuer* paediatric ratio.
- C, 5:1: an outdated ratio no longer recommended.



- D, 3:1: this is the neonatal resuscitation ratio, not for a child.

Final Answer: 30 compressions to 2 breaths $\Rightarrow$

Answer: (B) [Go Back to Q15](#)



Answer Key

Q	Ans	Q	Ans	Q	Ans	Q	Ans	Q	Ans
1	C	2	A	3	C	4	A	5	B
6	D	7	B	8	D	9	C	10	A
11	C	12	D	13	A	14	B	15	B

