DO NOT OPEN THIS TEST BOOKLET UNTIL YOU ARE ASKED TO DO SO

TEST BOOKLET AP (AS&H) CHEMISTRY—2016

Time	Allowed : 2 Hours	[Maximum Ma	arks : 100
	All questions carry equal marks.		•
	INSTRUCTIONS		
1.	Immediately after the commencement of the examination, you sho does not have any unprinted or torn or missing pages or items		
	by a complete test booklet.		
2.	Write your Roll Number only in the box provided alongside.		
	Do not write anything else on the Test Booklet.		

- This Test Booklet contains 100 items (questions). Each item comprises four responses (answers). Choose only one response for each item which you consider the best.
- 4. After the candidate has read each item in the Test Booklet and decided which of the given responses is correct or the best, he has to mark the circle containing the letter of the selected response by blackening it completely with Black or Blue ball pen. In the following example, response "C" is so marked:
 - (A) (B) (D)
- 5. Do the encoding carefully as given in the illustrations. While encoding your particulars or marking the answers on answer sheet, you should blacken the circle corresponding to the choice in full and no part of the circle should be left unfilled. After the response has been marked in the ANSWER SHEET, no erasing/fluid is allowed.
- You have to mark all your responses ONLY on the ANSWER SHEET separately given according to 'INSTRUCTIONS FOR CANDIDATES' already supplied to you. Responses marked on the Test Booklet or in any paper other than the answer sheet shall not be examined.
- All items carry equal marks. Attempt all items. Your total marks will depend only on the number of correct responses marked by you in the Answer Sheet. There will be no negative marking.
- Before you proceed to mark responses in the Answer Sheet fill in the particulars in the front portion of the Answer Sheet as per the instructions sent to you.
- If a candidate give more than one answer, it will be treated as a wrong answer even if one of the given answers happens to be correct.
- 10. After you have completed the test, hand over the Answer Sheet only, to the Invigilator.

DO NOT OPEN THIS TEST BOOKLET UNTIL YOU ARE ASKED TO DO SO

A.P (AS&H) CHEMISTRY 2016

Time Allowed: 2 Hours]

[Maximum Marks: 100

1. The effect of ionic strengths on reaction velocity of ionic reactions has been studied and found that a plot of $\log \frac{k}{k_0}$ $v/s \sqrt{\mu}$ gives a straight line with a slope equals to Z_a . Z_b .

Which of the following reactions has reaction velocity independent of ionic concentrations (The value of $Z_a.Z_b$ of each reaction is given in parenthesis)

(A)
$$2[Co(NH_3)_5 Br]^{++} + Hg^{+2} + 2H_2O \rightarrow 2[Co(NH_3)_5 (H_2O)]^{+3} + HgBr_2$$

 $(Z_a \cdot Z_b = 4)$

- (B) $NH_4^+ + CN\overline{O} \rightarrow NH_2CONH_2 (Z_a \cdot Z_b) = -1$
- (C) $[Cr(NH_2CONH_2)_6]^{+3} + 6H_2O \rightarrow [Cr(H_2O)_6]^{+3} + 6NH_2CoNH_2 (Z_a \cdot Z_b = 0)$
- (D) $[Co(NH_3)_5 Br]^{+2} + O\overline{H} \rightarrow [Co(NH_3)_5(OH)]^{+2} + B_r^- (Z_a \cdot Z_b = -2)$
- 2. The rate equation in terms of product concentration for zero order reaction is:

(A)
$$\frac{1}{[A_0]-[x]}-\frac{1}{[A_0]}=kt$$

(B) [x] = kt

(C) 2.303 log
$$\frac{[A_0] - [x]}{[A_0]} = -kt$$

(D)
$$\frac{1}{\{[A_0] - [x]\}^2} - \frac{1}{[A_0]^2} = 2kt$$

[Rate = $k[A]^n$; [A] = concentration of reactant at a given time t(s).

[A0] = initial concentration of the reactant

[x] = concentration of the product at a given time t(s)

 For the reaction A → C + D, the initial concentration of A is 0.010 M. After 100 secs the concentration of A is 0.0010 M. The rate constant has a numerical value 9.0.

The order of the reaction is:

(A) first order

(B) second order

(C) third order

- (D) zero order
- 4. Which of the following observations can be explained without the knowledge of hydrogen voltage ?
 - (A) The reversible electrode potential (E°) of H₂ and Zn are ± 0.000 V and -0.76 V respectively. Hydrogen alone should be liberated at cathode but Zn starts depositing along with liberation of H₂ at cathode inspite of their large difference in potential.
 - (B) The reversible electrode potential of H₂ at pH 7 (neutral solution) is - 0.41 V. This value is very close to that of Cd 0.48 V. Hence Cd and H₂ are expected to evolve simultaneously but only Cd is deposited at cathode.
 - (C) The standard electrode potential of Pb is -0.13 V and that of Zn as -0.76 V. Both should dissolve in dilute HCl evolving H₂. In actual practice neither Pb nor pure Zn dissolves in HCl and no evolution of H₂ occurs.
 - (D) The standard electrode potential of Ni and Cd are -0.24 and 0.76 V respectively. Ni⁺² are discharged first and metallic Ni will be deposited at cathode till all Ni⁺² present in solution is deposited. If cathode potential is made more negative and brought to the level of -0.76 V deposition of Zn takes place.

5. The specific conductivity (k) of a decinormal NaCl is 0.00918 mhos cm⁻¹. The, ionic conductance of Na and Cl ions at the same temperature are 43.2 and 65.3 respectively at infinite dilution. The degree of ionisation of NaCl at this temperature:

(A) 0.846

(B) 0.096

(C) 0.0109

(D) 1.009

6. Which of the following statements is not correct ?

- (A) Corrosion is an electrochemical process involving anode where the electron leaves the metal and cathode where reduction occurs, an electrolyte to conduct the ionic current between anode and cathode and a metallic path to conduct the electrons outside the electrolyte.
- (B) The reaction is thermodynamically feasible because of higher the enthalpy of certain compounds as compared to the free metal.
- (C) Both O₂ and H₂O must be present. The rust has the formulae Fe₂O₃.xH₂O.
- (D) The acid generated speeds the process of rusting. The process is autocatalytic:
 - (i) Fe \rightarrow Fe⁺² + 2e
 - (ii) $H^+ + e \rightarrow H$; $4H + O_2 \rightarrow 2H_2O$
 - (iii) $4Fe^{+2} + O_2 + (4 + 2x)H_2O \rightarrow 2(Fe_2O_3 \cdot xH_2O) + 8H^+$

- 7. Which of the following statements is not correct with regard to nmr spectral studies?
 - (A) All protons in tetramethyl silane [TMS, (CH₃)₄Si] are chemically equivalent but magnetically non-equivalent .
 - (B) The number of absorption bands gives the number of groups of nuclei in different molecular environments.
 - (C) The relative areas under the absorption peak for the various bands give the relative number of nuclei in each different environment.
 - (D) The finer details in the absorption band give relationship of one group of nuclei to other nearby groups of nuclei.
- 8. Which of the following statements is not correct?
 - (A) The infrared absorption bands are slightly broader since vibrational transitions are also accompanied by rotational transitions
 - (B) The UV absorption bands are broad since the electronic transitions are accompanied by vibrational as well as rotational transitions.
 - (C) Glass and quartz cells are used in infrared absorption spectral studies as these are transparent to infrared radiations.
 - (D) H₂O is a non-linear molecule and shows three absorption bands at about 3756, 3652 and 1545 cm⁻¹ due to asymmetric, symmetric and bending vibrations respectively.

- 9. Which of the following statements is not correct ?
 - (A) UV absorption spectral studies exhibit vibrational structure with well defined maxima.
 - (B) The wavelength (λ_{max}) of the absorption maxima increases regularly through the series and vary from UV region for C₆H₆,C₁₀H₈, C₁₄H₁₀ to the visible region yellow, blue and green for higher members naphthacene, pentacene and hexacene
 - (C) A ring in hexacene molecule (linearly conjugated aromatic hydrocarbon molecule, green in colour, 580 mμ) is insulated by introducing two -CH₂ groups as :

i.e. the linear conjugation is broken and absorption maxima shifts to a shorter wavelength in UV region.

(D) The absorption spectra of angularly annulated condensed ring hydrocarbons are at longer wavelength than those of linear compounds with the same number of rings for eg. phenanthrene though isomeric with anthracene absorbs in visible region.

- 10. Which of the following statements is not correct?
 - (A) In an esr spectrum the total number of signals expected for an electron delocalised over n equivalent protons is given as (2n I + 1) lines. Thus,

a frequency 9500 MHz and a magnetic field at about 0.34T gives five line spectrum with intensity 1:2:3:2:1.

- (B) The esr spectrum for the unpaired electron of the radical ion (SO₃)₂NO⁻ shows a three line spectrum of equal intensity.
- (C) An electron is delocalised over sets of two non-equivalent nuclei with I_i and I_j , the number of lines expected in the spectrum $(2nI_i+1)$ $(2mI_j+1)$ i.e. $(2\times 1\times 1+1)(2\times 1\times 1+1)=9$ lines for two non-equivalent nuclei.
- (D) Naphthalene contains two different sets of four equivalent protons.
 In naphthalene negative ion 4 × 4 i.e sixteen lines spectrum is expected.

11. Borazene (B₃N₃H₆) is isolectronic and isostructural with C₆H₆. There is a delocalised π-bonding just as there is in benzene. The number of electrons is the same as in benzene. Each nitrogen has one more electron than carbon but each boron has one fewer. The disubstituted borazene molecule B₃N₃H₄X₂ can have the following isomers:

Which of the following structures are not the same despite the X atom being one atom apart on each ring?

(A) II and IV

(B) I and II

(C) III and IV

(D) I and III

- 12. Which of the following statements is not correct?
 - (A) When steam is passed over red hot iron, the reaction never goes to completion $[4H_2O(g) + 3Fe(s) \rightleftharpoons Fe_3O_4(s) + 4H_2(g)]$ since the ratio $\frac{p_{H_2}}{p_{H_2}O}$ is constant at a given temperature (active mass of solids is taken as constant).
 - (B) CaCO₃ dissociates as CaCO₃(s) \(\) CaO(s) + CO₂(g). The equilibrium constant \(k_p = p_{CO_2} \) at a given temperature. \(p_{CO_2} \) is called dissociation pressure. At every temperature the system attains the new equilibrium and corresponds to its new dissociation pressure.

In lime kiln the evolved gas is continuously removed so that pressure of CO₂ falls below dissociation pressure at that temperature.

- (C) The substances for which triple point pressure exceed one atmosphere the condition for sublimation is automatically maintained, hence they directly pass into vapour state on heating.
- (D) In sulphur system both rhombic and monoclinic sulphur can co-exist along with their vapour at 151°C, hence a triple point.

- 13. Which of the following statements is not correct ?
 - (A) The triple point pressure is 4.6 mm for water. If the partial pressure of water vapour in the atmosphere falls below this value, water vapour will be condensed to the solid state when the atmosphere gets sufficiently cooled (frost).
 - (B) If a solid is heated and vapour pressure above it is not allowed to exceed the triple point pressure the solid will be converted into liquid.
 - (C) The lowest possible freezing point of a solution at which solution freezes 'enmasse' is called eutectic temperature and the solid phase which separates out during freezing at the eutectic point is called eutectic mixture.
 - (D) A dilute solution is cooled, ice separates with progressive lowering of freezing point until it freezes as a whole. A solution stronger than eutectic composition is cooled the salts separate out on cooling until the eutectic composition is reached when it freezes as a whole.

If a solution of eutectic composition is cooled, the solution freezes unchanged in composition at the eutectic temperature.

- 14. Which of the following statements is not correct ?
 - (A) Lyophobic colloids are reversible and have marked affinity for dispersion medium while reverse is true for lyophilic colloids.
 - (B) Lyophobic sols are easily coagulated with a small concentration of electrolyte but lyophilic sols require relatively large concentration of electrolyte.
 - (C) Lyophilic colloids are positively charged in strongly acid solution and negatively charged in alkaline solution. There is a concentration of H⁺ ion at which it moves neither to the cathode nor to the anode. This concentration of hydrogen ion is called isoelectric point of the colloid.
 - (D) Colloidal properties of soap (for eg. sodium palmitate) are due to aggregate of palmitate ions associated with some undissociated molecules, called micelles.

- 15. Which of the following statements is not correct?
 - (A) Lyophobic sol is stable due to electrical charge present on the colloidal particle whereas stability of the lyophilic colloids depends upon both the electrical charge and solvation.
 - (B) In a lyophobic sol the charge on all the colloidal particle is of the same sign the repulsive forces prevent the particles from approaching sufficiently close to one another and to coalesce and coagulate.
 - (C) The zeta potential is the difference in potential between the surface of the tightly bound layer and the electroneutral region of the solution. If zeta potential is small, the resultant potential energy is negative so that van der Waal attraction predominates over the electrostatic repulsion and the sol coagulates rapidly.
 - (D) In a lyophobic sol solvation also plays an important role. The colloidal particles are enclosed in a solvent cage, the cage serves as a barrier preventing the particles from coalescing to form aggregates.

- 16. Which of the following statements is not correct?
 - (A) Surfactants are the substances which when added to water reduces its surface tension.
 - (B) The effect of surface tension is to increase the area of the surface to maximum.
 - (C) The vapour pressure is a measure of the escaping tendency of molecules from the surface of the liquid.
 - (D) The molar heat of vapourisation of a liquid divided by its boiling point is nearly 88.0 J mol⁻¹ K⁻¹. The quantity $\frac{\Delta H_{vap}}{T_b}$ is also called entropy of vapourisation.
 - 17. Which of the following expressions is a thermodynamic equation of state?

(A)
$$\Delta A = \Delta U + T \left(\frac{\delta(\Delta A)}{\delta T} \right)_V$$

(B)
$$n_1 d\mu_1 + n_2 d\mu_2 + \dots + n_i d\mu_i = 0$$

(C)
$$V = T \left(\frac{\partial V}{\partial T} \right)_P + \left(\frac{\partial H}{\partial P} \right)_T$$

(D)
$$\Delta G = \Delta H + T \left[\frac{\delta(\Delta G)}{\delta T} \right]_P$$

- 18. Which of the following statements is not correct?
 - (A) Thermodynamics can indicate whether a particular process is feasible or not under a given set of circumstances for eg. temperature, concentration of reactant and product.
 - (B) The methods of thermodynamics are dependent on atomic and molecular structure.
 - (C) Thermodynamics does not take into account the rate of approach to the equilibrium.
 - (D) Thermodynamics deals quantitatively with equilibrium conditions i.e. condition which do not change with time.
- 19. Which of the following statements is not correct or a mathematical expression has not been applied correctly to a situation?
 - (A) In an exothermic reaction the bonds in the products are stronger than reactants. The stronger is the bond formation, the greater is the energy release.
 - (B) Since q_p = q_v + ΔnRT where q_p and q_v are enthalpy of reactions at constant pressure and at constant volume respectively and Δn is difference in number of moles of products and reactants. Thus for Haber's process in the synthesis of NH₃ Δn = 2 - (3 + 1) = -2, and the relation becomes q_p = q_v - 2RT.
 - (C) In an endothermic reaction, energy is used up in breaking the bonds in reactants, heat does not come out and has no chance to appear as heat and raise the temperature.
 - (D) A student performed an experiment on Bomb calorimeter for determining enthalpy of combustion. He uses the relation ΔH = ΔV + PdV and found ΔH = ΔV at constant volume.

- 20. Which of the following statements is not correct?
 - (A) First law of thermodynamics puts firmly the relation between heat absorbed and work performed by the system. It puts no restriction for the direction about the flow of heat.
 - (B) The energy of an isolated system remains constant during a specified change of state. It does not tell whether a specified change or a process is spontaneous or whether it is feasible.
 - (C) The energy of one form can be converted into an equivalent amount of energy of another form. It does not tell that heat energy cannot be completely converted into an equivalent amount of work.
 - (D) Thermodynamics in the form of second law tells us that how fast a reaction will actually occur.
- 21. Which of the following statements is not correct?
 - (A) The quantum theory assumes quantisation as postulate at the start.
 - (B) Each wave function ψ_n itself has no readily interpretable physical meaning, the square of the wave function ψ²_n gives the intensity of the electron wave or the probability of finding the electron at that point in space about the nucleus.
 - (C) The quantum theory due to Schrödinger does not predict the position of the particle since it defines precisely the energy of the electron. There becomes large uncertainty in electron position in reference to Heisenberg's uncertainty principle.
 - (D) Three quantum numbers are necessary for solving Schrödinger equation for an electron in three-dimensional world.

- 22. Which of the following statements is not correct?
 - (A) The value of n gives the most probable distance of the electron from the nucleus. 'n' is a measure of orbital radial size. The probability approaches zero but never zero even at a large distance.
 - (B) The angular momentum quantum number gives the shape of electron orbitals. The number of values of l for a given value of n tells us about the different orbital types.
 - (C) The number of values of n limits the values of l which in turn limits the values of m.
 - (D) A fourth quantum number becomes necessary to determine electron energies in a multielectron atoms. The quantisation of electron spin means that there are more than two possible orientations of an electron in a magnetic field.
- 23. Which of the following statements is not correct?
 - (A) Chemisorption has a very slow rate at low temperature i.e. it has a fairly large energy of activation but physical adsorption predominates at low temperature for the same adsorbent-adsorbate system.
 - (B) Physical adsorption involves a high degree of specificity whereas chemical adsorption is non-specific.
 - (C) The forces responsible for chemisorption are much stronger and so chemical adsorption is irreversible or reversible with great difficulty.
 - (D) Physical adsorption involves weak van der Waal forces and has a heat of adsorption of the order of 10,000 cals whereas chemical adsorption involves chemical forces and has heat of adsorption 20,000 to 60,000 cals.

24. The kinetics of the photochemical reactions of hydrogen-chlorine, and hydrogen-bromine reaction have been studied and found that the quantum yield for (A) hydrogen-chlorine reaction is very high and that of (B) hydrogen-bromine reaction is too low. Mechanism for both reactions are given below:

(A) hydrogen-chlorine reaction (B) hydrogen-bromine reaction

(i)
$$Cl_2 + h\nu \longrightarrow 2Cl$$

(i)
$$Br_2 + h\nu \longrightarrow 2Br$$
.

(ii)
$$Cl + H_2 \longrightarrow HCl + H$$

$$(ii)$$
 Br + H₂ \longrightarrow HBr + H

$$(iii) \ \ \mathbf{H} \ + \ \mathbf{Cl_2} \ \longrightarrow \ \ \mathbf{HCl} \ + \ \mathbf{Cl}$$

$$(iii)$$
 H + Br₂ \longrightarrow HBr + Br

(iv) H + HBr
$$\longrightarrow$$
 H₂ + Br

$$(v)$$
 2Br \longrightarrow Br₂

The important and chief reason for the difference in quantum yield is :

- (A) Both (A) and (B) are chain reactions
- (B) Reaction (ii) in (A) is exothermic and takes place spontaneously while reaction (ii) in (B) is endothermic and takes place slowly
- (C) There is no inhibition step in (A) while there is an inhibition step (iv) in (B)
- (D) The quantum yield in both (A) and (B) depends upon the number of times reaction steps (ii) and (iii) in each case are repeated before the final termination step

- 25. Which of the following statements is not correct?
 - The reason for the low quantum yield in a photochemical reactions are
 - (A) The excited molecule may simply emit the radiation of frequency which it absorbed.
 - (B) The excited molecule is deactivated through phosphorescence and fluorescence.
 - (C) The excited molecule is deactivated by converting its energy into the kinetic energy of the other molecule.
 - (D) The secondary process may involve a step which produces the reactant molecule as one of the products.
- 26. Which of the following statements is correct for canonical ensemble :
 - A system of specified volume, composition and temperature and think of it as being reproduced N times. (This N is different from N where it means the number of particles in the actual system). An ensemble is a collection of imaginary replication of the system.
 - (A) Consider an ensemble consisting of identical closed systems in thermal contact with each other. This thermal contact ensures that they are all at the same temperature but allows them to exchange energy with each other. The total energy of the ensemble is constant [N, V and T are common.
 - (B) In the ensemble the condition of constant temperature is replaced by the requirement that all the systems should have exactly the same energy, each system is individually isolated [N, V and E are common].
 - (C) In the ensemble the volume and temperature of each system is the same, but they are open. Thus matter can be imagined as able to pass between the systems, the composition of each one may fluctuate but the chemical potential (μ) is the same in each system [μ, V and T are common].
 - (D) An ensemble is stationary if $\frac{\partial p}{\partial t} = 0$ and p does not depend explicitly on time. This equation is called Liouville theorem.

- 27. Which of the following statements is not correct ?
 - (A) NaCl crystal consists of two interpenetrating cubic lattices, one of Na⁺ ions and the other of Cl⁻ ions. Na⁺ ion has fewer electrons than Cl⁻ ion. The scattering power of Na⁺ is less than Cl⁻ ion.
 - (B) Some reflections from Na⁺ ion lattice are out of phase with some Cl⁻ ion reflections. For other orientations the two sets of reflections are in phase and intense lines appear.
 - (C) Cancellation is complete only when the ions have identical scattering power as in case of KCl where K⁺ and Cl⁻ are almost identical and have the same scattering power. This is not so in case of NaCl where cancellation is incomplete and some intense lines and some faint lines appear.
 - (D) X-ray diffraction pattern for KCl appears to be that of simple cubic lattice whereas diffraction pattern for NaCl is that of face centred cubic lattice. Therefore both KCl and NaCl do not have same crystal structure.
 - 28. Nitric oxide has the electronic configuration $kk \sigma(2s)^2 \sigma^*(2s)^2 \sigma(2p_z)^2 \pi(2p_x)^2 \pi(2p_x)^2 \pi^*(2p_y)^2 \pi^*(2p_z)^2$. Which of the following statements is *not* correct ?
 - (A) Bond order is $\frac{1}{2}(8-3)$ i.e. 2.5
 - (B) Bond dissociation energy of nitric oxide 667.8 kJ/mol, is less than nitrogen which is 945.6 kJ/mol
 - (C) Bond length in NO is shorter than in nitrogen
 - (D) Bonding M.O. would have greater characteristics of atomic orbital of oxygen and the antibonding M.O. will have greater characteristics of atomic orbitals of nitrogen

- 29. The valence bond wave function for the HF molecule assuming that it is formed from 1s orbital of H i.e. A (electron 1) and 2p_z orbital of F i.e. B (electron 2). Which of the following statements or mathematical expressions is not correct?
 - (A) For purely covalent structure of HF, $\psi_{covalent} = \phi_A(1) \phi_B(2) + \phi_A(2) \phi_B(1)$
 - (B) For purely ionic structure H^+F^- , $\psi_{ionic} = \phi_B(1) \phi_A(2)$
 - (C) HF is a resonance hybrid shown as H − F ←→ H+F-
 - (D) HF is 80% covalent and 20% ionic

$$\psi = c_1 \ \psi_{\text{covalent}} + c_2 \ \psi_{\text{ionic}} \ \text{with} \ c_1^2 + c_2^2 = 1; \ c_1^2 = 80 \ \text{i.e.} \ c_1 = 0.89 \ \text{and}$$
 $c_2^2 = 20 \ \text{i.e.} \ c_2 = 0.45$

$$\psi = 0.89 \ [\phi_A(1) \ \phi_B(2) + \phi_A(2) \ \phi_B(1)] + 0.45 \ [\phi_B(1) \ \phi_B(2)]$$

- 30. Which of the following represents the correct electronic configuration of CO molecule ?
 - (A) $kk \sigma(2s)^2 \sigma^*(2s)^2 \pi(2p_x)^2 \pi(2p_y)^2 \pi^*(2p_x)^1 \pi^*(2p_y)^1$
 - (B) $kk \ \sigma(2s)^2 \ \sigma^*(2s)^2 \ \sigma(2p_x)^2 \ \pi(2p_y)^1 \ \pi(2p_z)^1 \ \sigma^*(2p_x)^2$
 - (C) $kk \sigma(2s)^2 \sigma^*(2s)^2 \sigma(2p_x)^2 \pi(2p_x)^2 \pi(2p_y)^2 \pi^*(2p_x)^1$
 - (D) $kk \sigma(2s)^2 \sigma^*(2s)^2 \pi(2p_x)^2 \pi(2p_y)^2 \sigma(2p_z)^2$

- 31. Which of the following statements is not correct?
 (A) Electronic configuration of He₂ is σ(1s)²σ*(1s)² and bond order is zero
 (B) Electronic configuration of He[±] is σ(1s)²σ*(1s)¹ and bond order is
 - (B) Electronic configuration of He_2^+ is $\sigma(1s)^2 \, \sigma^*(1s)^1$ and bond order is $\frac{1}{2}(2-1)$
 - (C) Bond order of He2 is zero and that of He2 is 0.5
 - (D) He₂ exists but He₂ does not exist
 - 32. In a hypothetical nuclear reaction $^{238}_{92}U + 17\,^{1}_{0}n \rightarrow X + 8\,^{0}_{-1}e$, the element X is :
 - (A) Nobelium

(B) Fermium

(C) Einstenium

- (D) Mendelevium
- 33. Which of the following is an example of linkage isomerism?
 - (A) [Co(en)₂ (NCS)₂]Cl and [Co(en)₂ (NCS)Cl] NCS
 - (B) $[(NH_3)_5 \text{ Ir} NO_2]Cl_2$ and $[(NH_3)_5 \text{ Ir} ONO]Cl_2$
 - (C) $[Pt(NH_3)_4]$ $[PtCl_6]$ and $[Pt(NH_3)_4Cl_2]$ $[PtCl_4]$
- (D) Cis-diglycinato platinum (II) and trans-diglycinato platinum (II)

 AP(AS&H)CHEMISTRY—2016 20

- 34. Which of the following is a natural order of stability of divalent complexes of transition metal ions with nitrogen ligands?
 - (A) Mn < Fe < Co < Ni < Cu > Zn
 - (B) Mn < Fe < Ni < Co > Cu > Zn
 - (C) Mn < Ni < Co < Fe > Cu < Zn
 - (D) Mn < Cu < Zn < Ni < Co < Fe
- 35. Which of the following rules is not correct for atomic term symbol:

The term atomic term symbol is designated as $2(\sum m_s) + 1$ L_J where the super script represents the multiplicity of the term 5 i.e. $\sum m_s$ and subscript the value of J. S is the sum of the spin quantum number $(m_s = \pm \frac{1}{2})$ for each unpaired electron. L is indicated by letters S, P, D, F for L values 0, 1, 2, 3:

- (A) Maximise the spin multiplicity (S) i.e. electrons occupy degenerate orbitals so as to retain parallel spins as long as possible
- (B) Maximise orbital angular momentum (L) i.e. fill the orbitals with the highest positive m values first
- (C) Select the minimum J value for the ground state of the subshell is more than half filled and the maximum value if the subshell is less than half filled
- (D) The complete subshell do not contribute to L or S because sum of the m_s and m_l values for a filled s, three filled p or five filled d-orbitals is zero

21

- 36. Which of the following statements is not correct?
 - (A) Term symbols are used to indicate both the electronic configuration and the resultant angular momentum of an atomic state
 - (B) The term symbol for carbon in the ground state is 3P0
 - (C) The term symbol for V³⁺ in the ground state 3F₂ (L = 3, S = 1, J = 4, 3, 2)
 - (D) With nitrogen the term 4S_{5/2} results in the ground state
 - 37. Crystal field theory could not explain satisfactorily :
 - (A) Stereochemistry
 - (B) Favourable coordination number
 - (C) The possibility of double bonding between metal and ligand
 - (D) Reaction path for substitution reactions
 - 38. Valence bond theory when applied to metal complexes provides a satisfactory explanation for :
 - (A) Limited qualitative aspects
 - (B) Interpretation or prediction of spectra
 - (C) Prediction of detailed magnetic properties
 - (D) Splitting of d-energy levels

- 39. Which of the following statements is not correct ?
 - (A) π-acid ligands have the ability to accept electron density into low lying empty π-orbitals, the word acid being used in the Lewis sense.
 - (B) π-acid ligands can stabilise low oxidation states.
 - (C) These ligands possess vacant π-orbitals in addition to lone pairs. These vacant orbitals can accept electron density from filled metal orbitals to form a type of π-bond that supplements the σ-bonding arising from lone pair donation.
 - (D) High electron density on the metal atom do not delocalise on to the ligand.
- There are unsaturated organic molecules and ions which form stable transition metal, π-complexes, metal ion being in low oxidation state.

Which of the following statements is not correct?

- (A) There is a donation of π-bonding electrons from the highest filled olefine orbitals into metal orbitals of suitable symmetry and there is a donation of electrons from filled metal orbitals of suitable symmetry back into the lowest lying π-antibonding orbitals of the olefine. The two components are synergically related.
- (B) The donation of π-bonding electron to the metal σ-orbital and the introduction of electron into π-antibonding orbital both strengthen the π-bonding in olefine. There is a significant shortening of the olefine C-C bond.
- (C) bis (penta hapto-cyclopentadienyl) iron, (h⁵ C₅H₅)₂Fe, has sand-wich structure confirmed by X-ray studies, and other physical and chemical evidence.
- (D) [PtCl₃(C₂H₄)⁻, anion Zeise salt : the plane of the olefine and C = C axis are itself perpendicular to one of the expected bond direction from the central metal atom. The expected line of a bond orbital from the metal strikes the C = C bond at its mid-point.

- 41. Which of the following statements is not correct:
 - (A) The π-acceptor ligands form bonds to the metal using σ-orbital and exercise their π-acidity by using π-orbitals whose nodal planes include the axis of the σ-bond.
 - (B) The metal atom in π-acceptor ligand complex lies out of the molecular plane of the ligand.
 - (C) The π-acceptor ligands include CO, dinitrogen (N₂), NO and isocyanide.
 - (D) CO and N₂ are isoelectronic and dinitrogen form metal complexes [Ru(NH₃)₅ N₂|Cl₂. [Ru(NH₃)₅N₂|⁺² cation can be obtained by the reaction of nitrogen (100 atm.) with [Ru(NH₃)₅ H₂O]⁺³. An important aspect is dinitrogen replaces water molecule. The chief difference is that for CO the σ-donor orbital is weakly antibonding whereas corresponding orbital for N₂ is of bonding character.
 - 42. Which of the following complexes of lanthanides has coordination number 12?
 - (A) $[Er(NCS)_6]^{-3}$

(B) $(NH_4)_2[Ce(NO_3)_6]$

(C) La[(ac ac)₃ (H₂O)₂]

(D) $(NH_4)_2CeF_6$

[ac ac = acetyl acetone]

- 43. Which of the following statements is not correct?
 - (A) Eu^{+2} , Sm^{+2} and Yb^{+2} cannot prepared by electrolytic reduction in aqueous solution.
 - (B) Eu^{+2} ($E^{\circ} = -0.43$ V) reduces V^{+3} more slowly than does than Cr^{+2} ($E^{\circ} = -0.41$ V).
 - (C) Eu^{+2} (E° = -0.43 V) and Cr^{+2} (E° = -0.41 V) will not reduce ClO_4^- but a weaker reductant V^{+2} (E° = -0.25 V) will do so.
 - (D) Sm⁺², Yb⁺² and Eu⁺² give blue solution in liquid NH₃. The blue solution reacts with 2, 2'-bipyridine (L) to form ML₄ type of compounds.
- 44. Which of the following statements is not correct?
 - (A) The magnetic moments in actinides are usually lower than the values calculated by Russell Saunders coupling. This is due to ligand field effects and to inadequacy of the coupling scheme.
 - (B) For f^q configuration (Cm⁺³) the lowest excited state lies about 4 eV above the ground level so that these ions show only charge transfer absorption band in UV region.
 - (C) Md⁺³ cannot be reduced to Md⁺² by Zn amalgam Cf, Es, Fm and Md do not exist as M⁺² ion in solution.
 - (D) 5f-orbitals can participate to some extent in covalent bonding, ligand effects are therefore expected.

45. Which of the following is an acid-base neutralisation reaction in liquid NH₃?

(A) Co +
$$2NH_4NO_3 \xrightarrow{liq NH_3} Co(NO_3)_2 + 2NH_3 + H_2$$

(B)
$$NH_4Cl + KNH_2 \xrightarrow{Liquid} KCl + 2NH_3$$

(C)
$$\operatorname{Zn}(\operatorname{NH}_2)_2 + 2\operatorname{NaNH}_2 \xrightarrow{\operatorname{Liquid}} \operatorname{Na}_2\operatorname{Zn}(\operatorname{NH}_2)_2 + 2\operatorname{NH}_3$$

(D)
$$H_2N(SO_2) NH_2 + NH_3 \xrightarrow{Liquid} H_2N(SO_2) \overline{N}H + NH_4^+$$

- 46. Which of the following statements is not correct?
 - (A) Low temperature or high pressure is necessary to work with liquid NH₃.
 - (B) The alkali metals in liquid NH₃ are powerful reducing agents.
 - (C) Ammonium salts dissolved in liquid NH₃ finds application in preparative chemistry.
 - (D) The tendency of solvolyis is more in liquid NH3 than in water
- 47. In which of the following reactions the non-aqueous solvent donates a proton so that the substance act as a base in it?

(A)
$$H_2S_2O_7 + H_2SO_4 \longrightarrow H_3SO_4^+ + HS_2O_7^-$$

solvent

(B)
$$SbF_5 + 2HF \longrightarrow SbF_6^- + H_2^+F$$

solvent

(C)
$$HNO_3 + H_2SO_4 \longrightarrow NO_2^+ + 2HSO_4^- + H_3O_4^+$$

solvent

(D)
$$Ag_2SO_4 + 2HCN \longrightarrow AgCN + H_2SO_4$$

solvent

- 48. Which of the following statements is not correct?
 - (A) Diamagnetism is a property of all forms of matter. All substances contain at least some electrons in closed shells. In closed shells the electron spin moments and orbital moments of individual electrons balance one another so that there is no net magnetic moment. Diamagnetic substances are repelled from magnetic field.
 - (B) Diamagnetism does not vary with temperature. Paramagnetism is inversely proportional to temperature and independent of field strength.
 - (C) The magnetic permeability, ratio of $\frac{B}{H}$ is given as:

$$\mu = \frac{B}{H} = 1 + 4\pi \begin{pmatrix} I \\ H \end{pmatrix} \text{ i.e. } \frac{B}{H} = 1 + 4\pi \hbar$$

where k is magnetic susceptibility per unit volume.

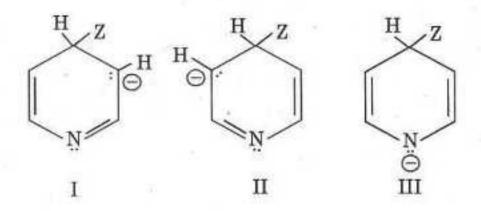
(D) For ferromgnetic substances above Curie's temperature a simple paramagnetic behaviour is seen but below Curie's temperature it is field strength dependent and different behaviour is seen.

For antiferromagnetic substances above Neel's temperature the susceptibility increases with increase in temperature and simple paramagnetic behaviour is not seen but below this temperature simple paramagnetic behaviour is observed. Both ferromagnetism and antiferromagnetism are independent of field strength.

49.	Whi	ch of the fol	lowing alloys	of copp	er has high	est percents	ige of
	copp	er ?				1	
	(A)	Gun metal		(B)	Brass	2.1	3 × 3
	(C)	Bronze		(D)	Bell metal		
50.	The	element, ato	mic number 11	.8, as a	nd when dis	covered will	belong
	to:					963	, S
,	(A)	s-block		(B)	p-block		
	(C)	d-block		(D)	f-block		
51.	The	structures of	diborane (B ₂ H ₆) and et	hane (C ₂ H ₆)	are not alike.	Which
	of t	the following s	statements is no	t correct	: ?		
٠.	(A)	Diborane is	diamagnetic an	d colour	less liquid	SY.	
4	(B)	There is a	free rotation ab	out B-B	bond	**	
X	(C)	Tetramethy	diborane on hy	drolysis	gives (CH ₃) ₂	B-OH only. T	wo CH ₃
		groups are a	attached per bore	on atom	in diborane a	nd that four h	ydrogen
98		atoms are a	alike and other	two are	different		
	(D)	Diborane ha	as two kinds of l	B-H bon	d. C ₂ H ₆ has 1	14 electrons ar	nd B ₂ H ₆
		has 12. elec	trons		8.8	186	
AP	AS&I	H) CHEMISTRY	7—2016	28		*	

- 52. Which of the following statements is not correct?
 - (A) Molecules such as CO and CN⁻ provide the large crystal field and are able to form π-bonds with the central metal ion. The π-bonding can markedly increase the magnitude of crystal field splitting.
 - (B) The crystal field splitting is strongly influenced by the oxidation state of a central metal ion and the type of d-electrons.
 - (C) The higher oxidation state of the metal ion, larger will be the crystal field splitting.
 - (D) Crystal field splitting in [Rh(NH₃)₆]⁺³, [Ir(NH₃)₆]⁺³ is much less than in [Co(NH₃)₆]⁺³ Crystal field splitting is least for complexes containing 5d electrons and greatest for those containing 3d electrons
- 53. Which of the following statements is not correct?
 - (A) The differences between lanthanides (4f series) and actinides (5f series) arise because of lower binding energies of 5f electrons in actinides than 4f electrons in lanthanides.
 - (B) 4f orbitals provide less shielding than 5f orbitals.
 - (C) Actinides have a variety of oxidation states ranging from +7 in Np and Pu (atomic numbers 93 and 94), +6 in Pa (At. No. 91) to Pu, + 5 in Th (at. no. 90) to Pu, +4 in Ac (at. no. 89) to Cm (at. no. 96). However the dominant oxidation state in actinides is +3 like in lanthanides.
 - (D) Ac⁺³ and Th⁺⁴ are colourless in aqueous solution and so also Cm⁺³ ions. These do not contain 5f electrons except Cm⁺³ which has half filled 5f subshell. Rest trivalent and tetravalent ions of U, Np, Pu, Am are coloured. Cm⁺⁴ is pale yellow in colour.

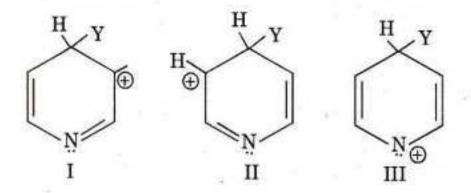
54. Nucleophilic attack at the 4-position yields a carbanion that is a hybrid of structure I, II and III ?



Which of the following statements is not correct?

- (A) Structure III is especially stable since the negative charge is located on nitrogen atom.
- (B) Nucleophilic substitution occurs more rapidly at 2-, and 4-positions than at 3-position.
- (C) Nucleophilic substitution occurs more rapidly on the pyridine ring than on benzene ring.
- (D) The same electronegativity of nitrogen that makes pyridine reactive towards electrophilic substitution makes pyridine less reactive towards nucleophilic substitution.

55. Electrophilic attack at the 4-position yields a carbocation that is a hybride of structures I, II and III:



Which of the following statements is not correct?

- (A) All these structures are less stable than the corresponding one for attack on benzene because of electron withdrawal by the nitrogen atom.
- (B) Pyridine undergoes substitution more slowly than C₆H₆.
- (C) Structure III is especially unstable because nitrogen atom bears a positive charge and has only sextet of electrons.
- (D) Electrophilic substitution occurs predominantly at 2- and 4-position.
- 56. Compounds containing two or more OH groups or = 0 groups on adjacent carbon atom undergo oxidative cleavage of carbon-carbon bond by periodic acid.

Which of the following compounds does not undergo oxidative cleavage of C-C bond by periodic acid?

- (A) CH₂OH.CH(OCH₃).CH₂OH
- (B) RCHOH.COR'

(C) R-CO.COR'

(D) R₂C(OH).CH(OH)R'

57. Which of the following reactions is not an example of Cannizzaro's reaction or crossed Cannizzaro's reaction?

(A)
$$C_6H_5CHO + HCHO \longrightarrow C_6H_5CH_2OH + HCOONa$$

(B)
$$CHO$$
 + HCHO $CONCEN$ + HCOONa OCH_3 + HCOONa

(C) CHO
$$\longrightarrow$$
 CHO \longrightarrow C

58. Which of the following is an example of Claisen condensation?

(A)
$$\left\langle \bigcirc \right\rangle$$
 $-\text{COOC}_2\text{H}_5$ + $\text{CH}_3\text{COOC}_2\text{H}_5$

$$\xrightarrow[]{C_2H_5ONa} \left(\bigcirc \right) \xrightarrow[]{C-CH_2COOC_2H_5} + C_2H_5OH$$

(B) 2CH₃CH₂COOC₂H₅ + C₂H₅ONa EtOH

$$\begin{array}{c} \text{CH}_3\text{CH}_2\text{CO-C-COOC}_2\text{H}_5 + 2\text{EtOH} \\ \downarrow \text{CH}_3 \\ \downarrow \text{H}^+ \\ \text{CH}_3\text{CH}_2\text{-C-C-COOC}_2\text{H}_5 \\ \mid \mid \quad \mid \\ \text{O} \quad \text{CH}_3 \end{array}$$

$$(C) \quad \stackrel{COOC_2H_5}{\vdash} + CH_3COOC_2H_5 \xrightarrow{C_2H_5ONa} \quad CH_3CH_2OOC - CH_2COOC_2H_5 \\ COOC_2H_5 \quad O$$

+ C₂H₅OH

(D)
$$CH_3CH_2$$
- C - $OC_2H_5 + C_6H_5CH_2COOC_2H_5 \xrightarrow{C_2H_5ONa} C_2H_5O$ - C - II
 O

C₆H₅ CHCOOC₂H₅ +EtOH

- 59. Which of the following statements is not correct?
 - (A) The B-complex vitamin deficiencies are generally multiple rather than individual with overlapping symptoms.
 - (B) Vitamin B₁₂, anti-pernicious anemia vitamin, is cyanocobalamine complex. Methyl cobalamine and 5'-deoxyadenosyl cobalamine are co-enzymes. Food for animal origin is the only source of B₁₂.
 - (C) Vitamin E is an oxidant. It helps in the non-enzymatic oxidation of various cell components (unsaturated fatty acids) by molecular oxygen, free radicals, hydrogen peroxide
 - (D) Vitamin A is collectively used many structurally related and biologically active compounds, retinol, retinal, retinoic acid, β-carotine. Its deficiency causes night blindness. Its severe deficiency leads to xerophthalmia (dryness in conjunctiva and cornea and keratinisation of epithelial cell).
- 60. Which of the following statements is not correct with regard to Friedel-Craft alkylation reaction?
 - (A) Aryl halides cannot take the place of alkyl halides.
 - (B) An aromatic ring less reactive than that of halobenzene can only undergo Friedel-Craft reaction.
 - (C) Aromatic rings containing the -NH₂, -NHR, or NR₂ do not undergo Friedel-Craft alkylation reaction.
 - (D) There is a danger of poly-substitution.

61. Which of the following statements is not correct?

A carbocation may:

- (A) rearrange to a more stable carbocation
- (B) eliminate hydrogen ion to form alkene
- (C) eliminate an alkene to form a larger carbocation
- (D) alkylate an aromatic ring
- 62. Which of the following is an example of stereospecific eletrophilic addition reaction ?

(B)
$$CH_2: + C=C \subset \longrightarrow [-C-C-] \longrightarrow -C-C-C CH_2: + CH_2$$

$$(\mathbf{C}) \quad -\overset{1}{\mathbf{C}} - \mathbf{H} + \mathbf{C} \mathbf{H}_2 \ \longrightarrow \ -\overset{1}{\mathbf{C}} - \mathbf{C} \mathbf{H}_2 - \mathbf{H}$$

(D)
$$CH_2 + C=C \longrightarrow -\stackrel{1}{C}-\stackrel{$$

63. Which of the following reactions gives sec-butyl benzene?

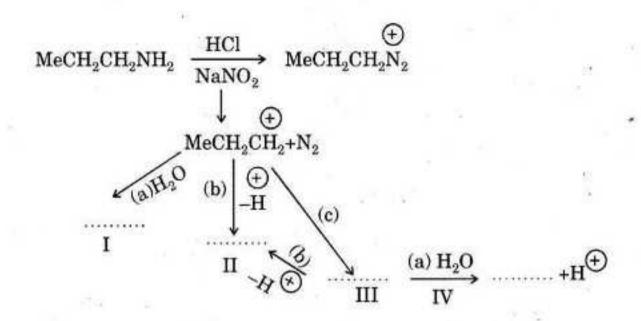
(A)
$$C_6H_6 + CH_3 - CHCH_2C1 \xrightarrow{AlCI_3} -18 \text{ to } 80^\circ$$

(B)
$$C_6H_6 + (CH_3)_3C-OH \xrightarrow{H_2SO_4}$$

(C)
$$C_6H_6 + (CH_3)_2C = CH_2 \xrightarrow{H_2SO_4}$$

(D)
$$C_6H_6 + CH_3(CH_2)_3Cl \xrightarrow{AlCl_3}$$

64. Carbonium ion can undergo three main reactions (a) combination with a nucleophile (b) elimination of a proton and (c) rearrangement of structure.
This is illustrated in the following reaction:



The species obtained are : iso-propyl cation, propylene, n-propanol and isopropanol. Identify the *correct* sequence of the species I to IV :

- (A) n-propanol, propylene, iso-propyl cation, isopropanol
- (B) iso-propanol, iso-propyl cation, propylene, n-propanol
- (C) iso-propyl cation, iso-propanol, n-propanol, propylene
- (D) propylene, n-propanol iso-propanol, iso-propyl cation
- 65. Which of the following is an example of pyridine alkaloid?
 - (A) Cocaine

(B) adrenaline

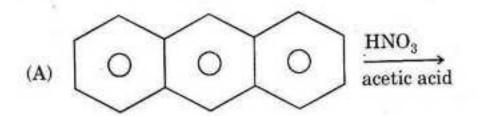
(C) nicotine

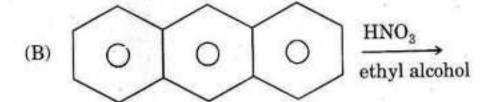
(D) cinchonine

AP(AS&H)CHEMISTRY-2016

36

66. In which example a substituent (-NO₂) occupies 9 position in anthracene to give 9-nitroanthracene?





67. Which of the following groups containing compound cannot react with Grignard reagent ?

The first three reactions are given for the preparation of alcohol from alkenes using oxymercuration-demercuration method and the last one is also used for the preparation of alcohol from alkenes using hydroboration-oxidation method.

Which of the following reactions is not correct as per Markownikov rule (addition of H_2O to carbon-carbon double bond) ?

$$(A) \quad \mathrm{CH_3(CH_2)_3CH=CH_2} \xrightarrow{\mathrm{Hg(OAc)_2}} H_2\mathrm{O} \xrightarrow{\mathrm{NaBO_4}} \mathrm{CH_3(CH_2)_3-C-CH_3}$$

$$(B) \quad CH_3CH_2-C=CH_2 \xrightarrow{Hg(OAc)_2} H_2O \xrightarrow{NaBO_4} CH_3CH_2-C-CH_3$$

$$(C) \quad CH_{3} \xrightarrow{\mid C-CH=CH_{2}} \quad \frac{Hg(OAc)_{2}}{H_{2}O}, \quad \stackrel{NaBO_{4}}{\longrightarrow} \quad CH_{3} \xrightarrow{\mid C-C-CH_{3}} \quad CH_{3}$$

(D)
$$CH_3$$
— C — CH = CH_2 $\xrightarrow{(BH_3)_2}$ $\xrightarrow{H_2O_2}$ $\xrightarrow{CH_3}$ CH_3 — C — CH_2CH_2OH CH_3

68.

69. In which of the following example rearrangement of carbocation is not possible giving S_N2 mechanism for the reaction ?

$$(D) \quad \begin{array}{cccc} CH_3 & CH_3 \\ & & & \\ I & & & \\ CH_3-C-CH_2X & Z \end{array} & CH_3-C-CH_3 & Z : entering group \\ & & & \\ I & & \\ I & & & \\ I & & \\$$

- 70. Which of the following disaccharides neither forms osazone nor shows mutarotation ?
 - (A) Lactose

(B) Sucrose

(C) Maltose

(D) Cellobiose

- 71. Which of the following statements is not correct?
 - (A) α D (+) glucose and β D (+) glucose are diastereomers differ in configuration Cl.
 - (B) D (+) glucose gives a positive test with Schiff's reagent and forms bisulphite addition product.
 - (C) D (+) glucose exists in two isomeric forms (I mp 146° and II mp 150°) which undergo mutarotation, the specific rotation drops from an initial +112° to +52.7° and in other case the value rises from 19° to 52°.
 I is called α D (+) glucose and II is called β D (+) glucose.
 - (D) D-(+)-glucose forms two isomeric methyl D-glucosides. D-(+)-glucose on treating with CH_3OH and HCl gives a compound methyl -D-glucoside which contains only one CH_3 group, hemiacetal $-C\frac{H}{OH}OCH_3$ but resembles with properties of acetal $-C\frac{H}{OCH_3}OCH_3$
 - 72. Which of the following oxidising agents oxidise glucose to glucaric acid?
 - (A) Bromine water
 - (B) Alkaline solution of Cu(II) complexed with tartrate ions
 - (C) HNO₃
 - (D) Ammonical solution of AgNO₃

- 73. How many chiral centres are there in fructose and how many stereoisomeric 2-ketohexoses are there?
 - (A) 4 and 16

(B) 3 and 8

(C) 5 and 32

- (D) 4 and 8
- 74. In glyceraldehyde the sequence for R, S-configuration is as :
 - (A) H, CHO, OH, CH2OH
- (B) CH₂OH, OH, H, CHO
- (C) CHO, CH2OH, OH, H
- (D) OH, CHO, CH2OH, H
- 75. Which of the following compounds reacts with Grignard's reagent (CH₃MgI) to give acetic acid under appropriate conditions?
 - (A) H—C

(B) CO₂

(C) CH₃C≡N

- (D) H₂O
- 76. Which of the following reactions gives ketene?
 - (A) $CH_3Li + N_2O \longrightarrow CH_3N = N -OLi \longrightarrow$ ethereal solution
 - (B) $CH_2=C-CH_2Br \xrightarrow{Zn \text{ dust}}$ Br
 - (C) C_2H_2 $\xrightarrow{pass C_2H_2 \text{ gas}}$ $\xrightarrow{at \ 0^{\circ}\text{C in NaOCl}}$ solution in the absence of light and air
 - (D) CH₃COCH₃ pass vapours of

 acetone through
 an iron free copper tube
 at 700°C

P.T.O.

77. Which of the following reactions of benzene diazonium chloride gives benzene?

(A)
$$C_6H_5N=N-Cl + C_6H_5NH_2 \rightarrow$$

(B)
$$C_6H_5N=N-Cl + H_3PO_2 + H_2O \rightarrow$$

(C)
$$C_6H_5N=N-Cl + C_6H_5OH \rightarrow$$

(D)
$$C_6H_5N=N-C1$$
 $\xrightarrow{H_2O, H^+}$

Which of the following statements is not correct ? 78.

- Silicon oils is a condensation product of dihydroxy silane molecules with linkage and two methyl groups on each silicon atom
- Nylon 6 is a polymer, a self-condensation product of aminocaproic acid (B)
- Dacron is a polymer, a condensation product of ethylene glycol and terphthalic acid
- Teflon is a polymer of tetrachloroethylene (D)

Which of the following compounds is strongest in acidic character? 79.

(A)
$$O_2N$$
 O_2 O_2N O_2 O_2N O_2 O_2N O_2 O_2N O_2 O_3N O_2 O_4 O_4 O_4 O_5 O_7 O_8 O_8

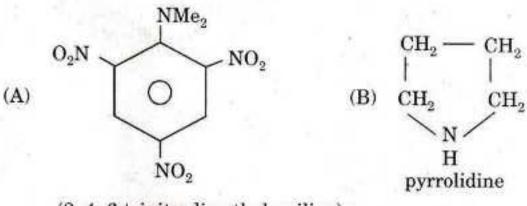
Br

(C)

(D)

OH

80. Which of the following is among the strongest organic bases known (having pKb too small to be measured accurately). Quaternary alkyl ammonium hydroxide is an exception?



- 81. Which of the following is on the bank of river Pabbar?
 - (A) Rohru

(B) Hatkoti

(C) Saraswati Nagar

- (D) All of these
- 82. Which pass joins Kangra and Bharmaur?
 - (A) Jalsu

(B) Darati

(C) Kugti

(D) Chobia

83.	Raja	of which princely sta	ate helped Jahan	gir in capturi	ing the Kangra fort?
	(A)	Shahpur	(B)	Nurpur	
,	(C)	Guler	(D)	Kutlehar	±1
84.	In w	which princely state d	id Tughlaq Khan	who rebelled	against Muhammed
	Shal	h I take shelter ?			*
	(A)	Kangra	(B)	Kutlehar	
	(C)	Handur	(D)	Sirmaur	
85.	Whe	en did Ram Singh orga	nise a force consis	sting mainly o	f Pathanias of Nurpur
	and	Katoaches of Kangr	a to oust the Br	itish ?	
117	(A)	1843-44	(B)	1846-48	
	(C)	1854-56	(D)	1857-59	
86.	Wh	ich was the first hyd	lel project to be	given to priv	ate sector ?
	(A)	Baspa	(B)	Larji	
	(C)	Binwa	(D)	Malana	
87.	In	which district of H.P	. is Talra sanctu	ary?	
	(A)	Kinnaur	(B)	Shimla	
	(C)	Kullu	. (D)	Bilaspur	
AP(AS&H) CHEMISTRY—2016	44		

88. E	Date	of which festival i	is decided by	the I	amas on the l	oasis of ast	rological
c	calcu	lations ?			te*		
((A)	Losser		(B)	Gochi		
((C)	Jidjed		(D)	Holi		
89.	To w	which Viceroy was a	a bell gifted	by the	king of Nepal	which was	used as
1	Dinr	nergong at the Vic	eregal Lodge	Shim	ala (now IIAS)	?	
	(A)	Lord Mayo		(B)	Lord Dufferin	ì	
ì	(C)	Lord Curzon		(D)	Lord Ripon		
90.	To v	which district of H	.P. does film	star	Kangna Rana	ut belong ?	
	(A)	Mandi		(B)	Shimla	,	
	(C)	Chamba		(D)	Kangra		
91.	Who	is the author of	The Chinar	Leave:	s ?		
	(A)	M.L. Fotedar		(B)	N.N. Vohra		
	(C)	Mehbooba Mufti		(D)	Omar Abdul	lah	Ŷ.
AP(AS	S&H	CHEMISTRY-201	6 4	5		26 19	P.T.O.

92.	Who won the 2015 Delhi Vidhan	Sabha election	s by biggest m	argin?		
	(A) Arvind Kejriwal	(B) Manisl	n Sisodia			
	(C) Pankaj Pushkar	(D) Mahen	idra Yadav			
93.	Through which city of Punjab is	he proposed na	tural gas pipe l	line (TAPI)		
	designed to pass?			ij*.		
	(A) Amritsar	(B) Taran	Taran			
×	(C) Fazilka	(D) Patha	nkot			
94.	4. With which of the following is Sunita Narain associated?					
	(A) Cricket	(B) Music				
	(C) Banking	(D) Enviro	onment			
95.	When was the Bureau of Indian	Standards (BI	S) set up ?			
	(A) 1987	(B) 1989				
	(C) 1992	(D) 1996				
96.	With which of the following is S	atya Nadella a	ssociated?			
	(A) Google	(B) Nokia				
	(C) Microsoft	(D) Maste	er card			
AP(A	AS&H) CHEMISTRY—2016	46				

	(A) Medicine	O	B)	Algebra		
					1.0	
	(C) Geography	a	D)	Dance		
			-		3	77
98.	To which country does S	vetlana Alexie	evic	h, who won the	2015 Nob	el prize
	for literature belong?					
				165		
	(A) Russia	(B)	Belarus		
	8.1					
	(C) Hungary	0	D)	Poland		
99.	When was new (second)	Suez canal in	nau	gurated ?		
	(A) January 2015	(B)	March 2015		
			-			
	(C) August 2015	C	D)	October 2015		
		2.		16		
100.	With which of the follow	ving is Niels l	Boh	r associated ?		
	(A) Botany	(B)	Psychology		4
	(C) Music	- (D)	Physics		
		5 6 7 CHAR				and the last of th
AP(A	S&H) CHEMISTRY—2016	47				P.T.O.

97. Besides Rubaiyats, what is Omar Khayyam known for ?