

FEDERAL PUBLIC SERVICE COMMISSION COMPETITIVE EXAMINATION-2025 FOR RECRUITMENT TO POSTS IN BS-17 UNDER THE FEDERAL GOVERNMENT

CHEMISTRY, PAPER-I

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TIME ALLOWED: THREE I PART-I (MCQs) : MAXIM	HOURS UM 30 MINUTES	(PA (PA	ART-I M ART-II)	1CQ	Qs) MAXIM MAXIM	UM I UM N	MARKS: 20 ARKS: 80
NOTE: (i) First attempt PAR	Γ-I (MCQs) on sepa	rate (OMR A	nsw	ver Sheet which s	shall 1	be taken back
after 30 minutes .			_				
(ii) Overwriting/cutting	ng of the options/ans	swers	will not	t be	given credit.		
(iii) There is no negative	ve marking. All MCQ)s mu	st be atte	emp	ted.		
¥	PART-I (MCQs)	(CO I	MPULS	OR	<u>Y)</u>		
Q.1. (i) Select the best option/ans (ii) Answers given anywhere	wer and fill in the app e else, other than OM	propri R An	iate Box swer She	eet,	on the OMR Answ will not be conside	wer S ered.	heet.(20x1=20
1. Which of the following does n	ot have the ground-st	tate c	onfigura	ntior	$1s^22s^22p^6?$		
(A) Ne	(B) Na^+		((C)	Cl	(D)	None of these
2. A reaction is exothermic if:			,				
(A) ΔH>0	(B) ΔH<0		((C)	$\Delta G>0$	(D)	$\Delta G < 0$
3. In chemometrics, multivariate	e calibration involves	:					
 (A) Using multiple independer (B) Using regression only on 2 4. Langmuir isotherm assumes: 	nt variables to predict of X-data (C) Perform	depen ing co	dent vari	iable 1 wit	es hout regression	(D)	None of these
(A) Multilayer adsorption	(B) Uniform sur	face v	vithout ir	ntera	ctions between ads	orbed	molecules
(C) Gas behaves as ideal	(D) None of thes	se					
5. A zero-order reaction rate do	es not depend on:						
(A) Temperature (E	B) Pressure	(C)	Reactan	t co	ncentration	(D)	None of these
6. Enzymes act as catalysts by:		. /				. /	
(A) Increasing activation energy	gy	(B)	Lowerin	ng ac	ctivation energy		
(C) Changing equilibrium posi	ition	(D)	None of	f the	se		
7. In an electrolytic cell:							
(A) Electrons move from cathe	ode to anode	(B)	The ano	de i	s negative		
(C) Energy is consumed to driv	ve the reaction	(D)	None of	f the	se		
8. Silicones are primarily:		. .					
 (A) Organic polymers (B) Inor 9. The Nernst equation relates elements 	rganic polymers (C) H lectrode potential to:	lybrid	l organic	-ino	rganic polymers	(D)	None of these
(A) Pressure (B) Ten	mperature and concent	tratior	n ((C)	Surface area	(D)	None of these
10. In solvent extraction, distribu	tion coefficient is:	(D)	D-+' (۲_ 1			
(A) Ratio of solute concentrati	on in two phases	(B)	Ratio of	SOI'	vents used		
(C) Kauo of extractant to samp	pie volume	(D)	inone of	i ine	50		
(A) Liquid	$(\mathbf{B}) C_{\mathbf{a}\mathbf{c}}$		((\mathbf{C})	Solid	(D)	None of these
(A) Liquiu 12 Solubility of a solt is doorcess	(D) Uas d in presence of		(U)	Solid	(D)	mone of these
(A) Common ion	(B) High temper	ature	((\mathbf{C})	High Ph	(D)	None of these
13. The solubility of a gas in a light	uid is directly proper	tions) I to the r	U) nart	ial pressure of the	(U) gas a	hove the
liquid. This statement is based	l upon:		inc]		Pressure of the	545 C	
(A) Raoult's law	(B) Henry's law		(C)	Kohlrausch's law	(D)	None of these
14. The heat capacity of a calorin	neter is determined us	sing:	((-)			
(A) A bomb calorimeter (B) A l	known standard materi	ial (C) Adiabat	tic te	emperature rise	(D)	None of these
15. Fugacity is used to measure:			/		1		
(A) Real gas behavior (B) Ide	eal gas compressibility	v (C)	Equilibr	ium	constant	(D)	None of these
16. Lattice enthalpy is always:			1			· /	0
(A) Positive and represents end	ergy absorption		(B) Pe	ositi	ve and represents e	nergy	release
(C) Negative and represents er	nergy absorption		(D) N	legat	tive and represents	energ	y release
17. According to Hess's law:	. 1		. /	0	×	0	-
(A) Enthalpy change depends	on reaction pathway		(B) E	ntha	lpy change is path	indep	endent
(C) Enthalpy depends on final	state only		(D) E	ntha	lpy change include	s entr	opy changes
18. Debye-Hückel theory is used t	to calculate:						
(A) Solubility of salts (E	B) Ionic strength of sol	utions	s (C) Gi	ibbs	free energy of ions	(D)) None of these
19. Potentiometric titration involved	ves measurement of:						
(A) Current in the reaction			(B) V	olur	ne at equivalence		
(C) Electrode potential as a fun	nction of titrant added		(D) N	lone	of these		
20. According to molecular orbita	al theory the bond or	der of	f 0 ₂ ⁻ is:				
(A) 1.0 (E	3) 1.5		((C)	2.5	(D)	None of these Page 1 of 2

PART-II

NOTE	2: (i) (ii)	Part-II is to be attempted on the separate Answer Book . Attempt ONLY FOUR questions from PART-II by selecting TWO questions from PART-II by se	om EACH				
	(iii)	SECTION . ALL questions carry EQUAL marks. All the parts (if any) of each Question must be attempted at one place instead of at different					
	(iv)	Write Ω No in the Answer Book in accordance with Ω No in the Ω Paper					
	(\mathbf{v})	No Page/Space be left blank between the answers All the blank pages of At	nswer Book				
	(•)	must be crossed.	ISWCI DOOK				
	(vi)	Extra attempt of any question or any part of the question will not be considered.					
	(vii)	Use of Calculator is allowed.					
Q. 2.	(a)	Define adsorption and explain its types. Derive an expression for the Langmuir adsorption isotherm and discuss factors influencing adsorption on solids	(08)				
	(b)	Why are transition metal complexes often colored? Explain using crystal field	(07)				
	<i>.</i>	theory with an example.					
	(c)	Compare and contrast the Lewis concept and the Bronsted-Lowry theory of acids and bases. Include relevant examples and limitations.	(05) (20)				
Q. 3.	(a)	What is lanthanide contraction? Analyze its impact on the atomic radii and chemical properties of lanthanides and post-lanthanides.	(08)				
	(b)	Analyze the quantum numbers associated with atomic orbitals and describe their	(07)				
	(c)	Describe the catalytic activity of transition metals with reference to their variable	(05) (20)				
	(0)	oxidation states. Support your explanation with one example.	(00) (20)				
Q. 4.	(a)	Derive the Gibbs free energy equation from the first and second laws of	(10)				
-		thermodynamics and discuss its importance in predicting the spontaneity of chemical					
	(b)	Explain the principle of fractional distillation. Compare it with simple distillation	(05)				
		using a diagram and examples of their applications.					
	(c)	Adsorption isotherms fail at high pressure. Why? Explain with reference to the limitations of the Freundlich isotherm.	(05) (20)				
			(1.0)				
Q. 5.	(a)	Why are Lewis's acids and bases more versatile in their applications than Bronsted I owry acids and bases? Provide two examples to support your answer	(10)				
	(b)	Derive the integrated rate law for a first-order reaction and explain why the half-life of					
		such a reaction is independent of its initial concentration.	(05)				
	(c)	State and explain the Heisenberg Uncertainty Principle. How does it limit our ability	(05) (30)				
		to precisely determine the position and momentum of particles?	(05) (20)				
Q. 6.	(a)	Discuss the Freundlich adsorption isotherm. Derive the equation and highlight its limitations.	(07)				
	(b)	Using a diagram, explain the principle and working of a galvanic cell. Discuss	(07)				
	(c)	Why does an exothermic reaction sometimes become non-spontaneous at high	(06) (20)				
	(-)	temperatures?					
Q. 7.	(a)	Explain the relationship between entropy, enthalpy, and Gibbs free energy. Derive	(08)				
	(b)	an expression for equilibrium using these thermodynamic quantities.	(07)				
	(u)	energy and enthalpy.	(07)				
	(c)	Define the concept of activation energy and illustrate its significance in reaction kinetics using the Arrhenius equation.	(05) (20)				
0.0	Daf-	the following: (5 seek)	(20)				
Q. 0.	Delli	(a) Difference between Homogenous and Heterogenous Catalysis	(20)				
		(b) Define Molecular Orbital Theory (MOT)					
		(c) Difference between ANOVA and Gaussian Distribution					
		(d) Define Chromatography and write key differences between Paper Chromatography and Gos Chromatography					
		Chromatography and Gas Chromatography *******					