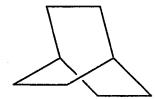
类	類組:化學類 科目:無機化學(1003)	共_	_7	頁	第 <u>_1</u>	頁
	選擇題:單選題(只有一個答案對,答錯不倒扣,每一題兩分	分,總分	分100	))		
	<ol> <li>Modern periodic table is arranged in a sequence based on the prop</li> <li>(A) Electronegativity (B) Ionization Energy (C) Atomic Weight</li> <li>(D) Atomic number (E) Atomic radius</li> </ol>	erty of	:			
	2. Which element is in liquid form at T = 50 °C (A) Antinomy (B) Gallium (C) Sulfur (D) Carbon (E) Sodium					
	<ul> <li>3. What is the scientific contribution for the 2011 Nobel Laureate Dr (A) Discovered the icosahedral phase of quasiperiodic crystals</li> <li>(B) Pioneering work on graphene.</li> <li>(C) Developed the cathode material of Li-ion battery</li> <li>(D) Discover the buckminsterfullerene</li> <li>(E) For the discovery of new superconducting materials</li> </ul>	. Dan S	shecht	tmaı	1?	
	<ul> <li>4. Which of the following molecules is the strongest Lewis acid?</li> <li>(A) NF<sub>3</sub> (B) SbF<sub>5</sub> (C) KCl (D) PCl<sub>3</sub> (E) SnCl<sub>2</sub></li> </ul>					
	<ul> <li>5. Identify the point group of SbF<sub>4</sub><sup>-</sup></li> <li>(A) T<sub>d</sub> (B) C<sub>2h</sub> (C) C<sub>2v</sub> (D) D<sub>2d</sub> (E) D<sub>4h</sub></li> </ul>					
	6. Identify the point point group of SeF <sub>3</sub> <sup>+</sup> (A) D <sub>3h</sub> (B) C <sub>3v</sub> (C) C <sub>2v</sub> (D) C <sub>2h</sub>	,				
	7. Which of the following molecule in liquid form exhibits paramagn (A) $NH_3$ (B) $CO_2$ (C) $O_2$ (D) $N_2$ (E) $H_2$	netic pro	operty	y?		
	8. The crystals of CeO <sub>2</sub> exhibit a fluorite structure with a coordination What is the coordination number of the cation? (A)4 (B)6 (C)8 (D)10 (E) 12	n numb	er of	'4 fo	or the	anion.
	<ul><li>9. Which of the following element/compound does NOT exhibit a lastate?</li><li>(A) Cadmium Iodide (B) Titanium Dioxide (C) Graphite</li><li>(D) Molybdenum disulfide (E) Boron nitride</li></ul>	yer stru	cture	in tl	ne sol	id
	10. Identify the crystal system with a $\neq$ b $\neq$ c, $\alpha = \gamma = 90^{\circ}$ , (A) Cubic (B) Monoclinic (C) Orthorhombic (D) Hexagonal (E) Triclin		·.			
	The questions 11 and 12 are for the molecule Buckminsterfullerene C6 11. Identify the point group of Buckminsterfullerene. (A) $D_{5h}$ (B) $T_d$ (C) $I_h$ (D) $O_h$ (E) $C_{5v}$	0 molec	cule.			
	12. How many symmetrically independent carbon atom(s) for a C <sub>60</sub> m (A) 1 (B) 2 (C) 4 (D) 5 (E) 10	olecule	?			
		<del></del>	<del>-</del> -			-

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Questions 13 and 14 are related to the molecule C<sub>8</sub>H<sub>14</sub> bicyclo[2.2.2]octane shown below:



bicyclo[2.2.2]octane

- 13. Identify the point group of bicyclo[2.2.2]octane
- (A)  $C_3$  (B)  $C_{3v}$  (C)  $C_{3h}$  (D)  $D_{3d}$  (E)  $D_{3h}$
- 14. How many peaks will you expected from the <sup>1</sup>H NMR spectrum for the molecule bicyclo[2.2.2] octane?
- (A) 1 (B) 2 (C) 3 (D) 4 (E) 5

Questions 15 and 17 are related to the molecule XeF<sub>4</sub> and the corresponding character table with hidden point group symbol is shown below:

?	Ε	$2C_{4}$	$C_2$	$2C_2'$	$2C_2''$	i	2 <i>S</i> <sub>4</sub>	$\sigma_{h}$	$2\sigma_{\nu}$	$2\sigma_{\sf d}$
$\overline{A_{1g}}$	1	1	1	1	1	1	1	1	1	1
$A_{2g}$	1	1	1	-1	-1	1	1	1	<b>—1</b>	-1
$B_{1g}$	1	-1	1	1	<b>-1</b>	. 1	_1	1	1	-1
$B_{2g}^{\circ}$	1	-1	1	-1	1	1	-1	1	-1	1
$E_{\rm g}$	2	0	-2	0	0	2	0	-2	O	0
$\ddot{A_{1u}}$	1	1	1	1	1	-1	<b>-1</b>	<b>-1</b>	-1	-1
$A_{2u}$	1	1	1	<b>-1</b>	-1	<b>-1</b>	-1	<b>-1</b>	1	1
$B_{1u}$	1	-1	1	1	-1	-1	1	<b>-1</b>	-1	1
$B_{2u}$	1	-1	1	-1	1	$_{1}-1$	1	-1	1	-1
$E_{u}$	2	0	-2	0	0	-2	0	2	O	0

- 15. Identify the point group of XeF<sub>4</sub>
- (A)  $C_4$  (B)  $C_{4h}$  (C)  $C_{4v}$  (D)  $D_{4h}$  (E)  $D_{4d}$
- 16. Identify the irreducible representation of  $d_{xy}$  orbital.
- (A)  $A_{1g}$  (B)  $A_{2g}$  (C)  $B_{1g}$  (D)  $B_{2g}$  (E)  $A_{1u}$
- 17. What is the order for the point group of XeF<sub>4</sub>
- (A) 10 (B) 12 (C) 14 (D) 16 (E) 18

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- 18. Which of the following element/compound is an p-type semiconductor?
- (A) Silicon Carbide (B) Diamond (C) Boron Nitride
- (D) Arsenic-doped Germanium (E) Gallium-doped Germanium
- 19. Which of the following species is paramagnetic in its ground state?
- (A)  $O_2^{2-}$  (B) CO (C)  $F_2$  (D) CN<sup>-</sup> (E) NO

The questions 20 and 21 are related to the molecule [Fe(CO)<sub>5</sub>] with D<sub>3h</sub> point group.

$D_{3h}$	E	2C <sub>3</sub>	$3C_2$	$\sigma_{h}$	2S <sub>3</sub>	$3\sigma_{\rm v}$		
$A_1{}'$	1	1	1	1	1	1		$x^2 + y^2$ , $z^2$
$A_2{'}$	1	. 1	-1	1	1	-1	$R_z$	
E'	2	-1	0	2	<b>—1</b>	0	(x, y)	$(x^2-y^2,xy)$
$A_1''$	1	1	1	<b>—</b> 1	-1	-1	(x, y)	, , , ,
$A_2^{"}$	1	1	-1	<b>-1</b>	<b>-1</b>	1	Z	
E''	2	-1	0	<b>-</b> 2	1	0	$(R_x, R_y)$	( <i>xz</i> , <i>yz</i> )

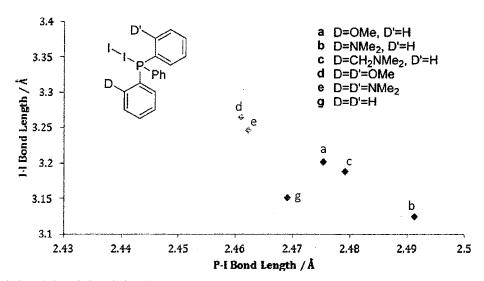
- 20. How many vibration modes for [Fe(CO)<sub>5</sub>] molecule?
- (A) 12 (B) 14 (C) 16 (D) 27 (E) 40
- 21. Identify the following irreducible representation for CO stretches that is IR active and Raman inactive.
- (A) A<sub>1</sub>'(B)A<sub>2</sub>'(C)E'(D) A<sub>1</sub>"(E) A<sub>2</sub>"
- 22. How many unpaired electrons are there in a Co<sup>3+</sup> ion?
- (A) 2 (B) 3 (C) 4 (D) 5 (E) 6
- 23. Which of the following compounds is the strongest base in water?
- (A) B<sub>2</sub>O<sub>3</sub> (B) CO<sub>2</sub> (C) Cl<sub>2</sub>O (D) K<sub>2</sub>O (E) P<sub>4</sub>O<sub>10</sub>
- 24. Which rotation symmetry operation is NOT allowed for the crystalline solid with 3D structure (A) 2 fold rotation (B) 3 fold rotation (C) 4 fold rotation (D) 5 fold rotation (E) 6 fold rotation
- 25. How many Na+ cations in an unit cell of NaCl crystal structure?
- (A) 1 (B) 2 (C) 3 (D) 4 (E) 5

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26. Given phosphine-I<sub>2</sub> (Ar<sub>3</sub>P-I<sub>2</sub>) adducts and the correlations between their P–I and I–I bond lengths are shown below. How many of following statement(s) is/are correct?

- (i) Phosphines forms Lewis acid-base adducts with  $I_2$ , in which phosphines are Lewis base and  $I_2$  is Lewis acidic.
- (ii) Generally, stronger phosphine donors result in the Ar<sub>3</sub>P-I<sub>2</sub> adduct with shorter I–I bond length.
- (iii) The I–I bond lengthening is attributed to the donation from phosphine ligands into  $\pi^*$  orbitals of I<sub>2</sub>.
- (iv) The I–I bond lengths of Ar<sub>3</sub>P-I<sub>2</sub> adducts are longer than that of I<sub>2</sub>.
- (v) Steric hindrance caused by the ortho-substituents does not influence the magnitude of the P-I and I-I bond lengths.



- (A) 1 (B) 2 (C) 3 (D) 4 (E) 5
- 27. For ditungsten W<sub>2</sub> molecule, what is the possible maximum bond order for it? (A) 3 (B) 4 (C) 5 (D) 6 (E) 7
- 28. Following the Q27, what is the electron count for W in  $W_2$  molecule?
- (A) 10 (B) 12 (C) 14 (D) 16 (E) 18
- 29. Assume that Hund's rule holds for all the systems considered in this problem, and L-S coupling is applicable for all systems. Which of the following has  ${}^{7}S_{3}$  as its ground term? (A) V (B) Cr (C) Mn (D) Fe (E) Co
- 30. Which of the following ions or molecules do you expect to be paramagnetic?
- (A)  $[Fe(CO)_5]$  (B)  $[Fe(CN)_6]^{2+}$  (C)  $[Cu(CH_3CN)_4]^+$  (D)  $[CoF_6]^{3-}$  (E)



31. Which of the following ions or molecules do you expect to be colorless?

(A)  $[Fe(CO)_5]$  (B)  $[Fe(CN)_6]^{2+}$  (C)  $[Cu(CH_3CN)_4]^+$  (D)  $[CoF_6]^{3-}$  (E)



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32. What d orbital splitting diagram would you expect for Fe(CO) <sub>5</sub>	molecule?								
(A) — — (B) — — (C) — — (D) — —	(E) — — —								
33. The electronic configuration of the boron atom is $1s^22s^22p^1$ . You expect the electron probability about a boron atom in free space is found to be  (A) tetrahedral (B) spherically symmetric (C) dumbbell shaped (D) double-dumbbell shaped (E) octahedral									
34. Which of the following ions or molecules do you expect to have the shortest S-N bond?  (A) PhSO <sub>2</sub> (NH <sub>2</sub> )  (B) NS <sub>2</sub> <sup>+</sup> (where N is the central atom)  (C) S <sub>2</sub> N <sub>2</sub> (with a square planar structure having only one type of S-N bond)  (D) N <sub>2</sub> S (where N is the central atom)  (E) NSF <sub>3</sub> (where S is the central atom)									
35. Both CrO <sub>4</sub> <sup>2-</sup> and MnO <sub>4</sub> <sup>-</sup> are very colorful. What type of electronic transitions is responsible for the colors of the complexes?  (A) d-d transitions  (B) Interligand electron transfer  (C) Metal-metal-to-ligand charge transfer  (D) Metal-to-ligand charge transfer  (E) Ligand-to-metal charge transfer									
36. The absorption spectra of $[Ni(H_2O)_6]^{2+}$ and $[Ni(en)_3]^{2+}$ (en = eth the figure below, along with a color wheel. How many of the following	36. The absorption spectra of $[Ni(H_2O)_6]^{2+}$ and $[Ni(en)_3]^{2+}$ (en = ethylenediamine) are shown in the figure below, along with a color wheel. How many of the following statement(s) is/are correct?								
(i) The dotted-curve belongs to $[Ni(H_2O)_6]^{2+}$ (ii) The color of $[Ni(H_2O)_6]^{2+}$ is orange (iii) The color of $[Ni(en)_3]^{2+}$ is purple (iv) Only $[Ni(en)_3]^{2+}$ exhibits absorption in near-infrared region (v) Both $[Ni(H_2O)_6]^{2+}$ and $[Ni(en)_3]^{2+}$ have an inversion center and th intense.	eir absorption bands should be								
Wavelength (Å)	<b>_</b>								
2000 4000 6000 8000 10000 12000  THE COLOR WHEEL  R = RED  \$\lambda = 6100 \text{ Å} \text{ O = ORANGE}\$  \$\lambda = 6800 \text{ Å}  \lambda = 5800 \text{ Å} \text{ Y = YELLOW}\$  \$\lambda = 6800 \text{ Å}  \lambda = 5300 \text{ Å} \text{ B = BLUE}\$									
Frequency (cm <sup>-1</sup> ) $\lambda = 4800 \text{ Å}$ V = VIOLET									
(A) 1 (B) 2 (C) 3 (D) 4 (E) 5	· <b>-</b>								
37. Which of the following fragments do you expect to be isolable w (A) CpCu (Cp = C <sub>5</sub> H <sub>5</sub> ) (B) Mn(CO) <sub>5</sub> (C) Cr(CO) <sub>4</sub> (D) Ni(CO) <sub>4</sub> (E) Cp	vith CH <sub>2</sub> ? pCo								

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- 38. Which of the following ions do you expect to exhibit the greatest observed magnetic moment?
- (A)  $[Ni(NH_3)_6]^{2+}$  (B)  $[Co(H_2O)_6]^{2+}$  (C)  $[Cu(H_2O)_6]^{2+}$  (D)  $[FeF_6]^{3-}$  (E)  $[Fe(CN)_6]^{3-}$
- 39. Which ion is mostly kinetically labile?
- (A)  $Co^{3+}$  (B)  $Ru^{2+}$  (C)  $Ti^{3+}$  (D)  $Cr^{3+}$  (E)  $Rh^{3+}$
- 40. The reactions of  $Cu^{2+}(aq)$  ion with en (en = ethylenediamine) feature the equilibrium constants K shown below:

$$[Cu(H_2O)_6]^{2+}$$
 + en  $\frac{-2 H_2O}{}$   $[Cu(en)(H_2O)_4]^{2+}$   $logK_1 = 10.72$ 

$$[Cu(en)(H_2O)_4]^{2+}$$
 + en  $\frac{-2 H_2O}{}$   $[Cu(en)_2(H_2O)_2]^{2+}$   $logK_2 = 9.31$ 

$$[Cu(en)_2(H_2O)_2]^{2+}$$
 + en  $\frac{-2 H_2O}{}$   $[Cu(en)_3]^{2+}$   $logK_3 = -1.0$ 

Why is the  $K_3$  significantly smaller than  $K_1$  and  $K_2$ ?

- (A) Jahn-Teller effect
- (B) Trans effect
- (C) Cotton effect
- (D) Hard-Soft Acid-Base Principle
- (E) 18-electron rule
- 41. For [Os<sub>2</sub>Cl<sub>8</sub>]<sup>2-</sup> anion, which of the following statements is correct?
- (A) This species should be paramagnetic.
- (B) There is a metal-metal quintuple bond present in this species.
- (C) This ion adopts a staggered conformation.
- (D) The electron count for each Os is 18.
- (E) The oxidation state for Os is +4.
- 42. Me<sub>2</sub>CHMgBr reacts with Ir(I)ClL<sub>3</sub> to give Ir(I)HL<sub>3</sub>. How many of following statement(s) is/are correct?
- (i) The reaction also produces propene as a by-product.
- (ii) The possible intermediate is Ir(CHMe<sub>2</sub>)L<sub>3</sub>.
- (iii) β-H elimination likely involves in this process.
- (iv) The reaction of Me<sub>2</sub>CHMgBr with IrClL<sub>3</sub> likely proceeds through associative mechanism.
- (A) 0 (B) 1 (C) 2 (D) 3 (E) 4
- 43. The reactions of  $[PtCl_4]^{2-}$  with  $NH_3$  (reaction I) and of  $[PtCl_4]^{2-}$  with  $[NO_2]^-$  (reaction II) followed by  $NH_3$  are ways of preparing:
- (A) I: trans- $[PtCl_2(NH_3)_2]$ ; II: trans- $[PtCl_2(NH_3)(NO_2)]$
- (B) I: cis-[PtCl<sub>2</sub>(NH<sub>3</sub>)<sub>2</sub>]; II: trans-[PtCl<sub>2</sub>(NH<sub>3</sub>)(NO<sub>2</sub>)]
- (C) I:  $\operatorname{cis-[PtCl_2(NH_3)_2]}$ ; II:  $\operatorname{cis-[PtCl_2(NH_3)(NO_2)]^{-1}}$
- (D) I: trans- $[PtCl_2(NH_3)_2]$ ; II: cis- $[PtCl_2(NH_3)(NO_2)]$

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- 44. The kinetics of the isomerization of cis to trans-[Mo(CO)<sub>4</sub>(PEt<sub>3</sub>)<sub>2</sub>] can be followed by IR spectroscopy. Which of the following regions of the IR spectrum would you focus on to monitor the reaction?
- (A) 800-1000 cm<sup>-1</sup>
- (B) 1400-1600 cm<sup>-1</sup>
- (C) 1800-2000 cm<sup>-1</sup>
- (D) 2400-2600 cm<sup>-1</sup>
- (E) 3000-3500 cm<sup>-1</sup>
- 45. Which experimental technique operates on the slowest timescale?
- (A) NMR spectroscopy
- (B) UV-vis spectroscopy
- (C) IR spectroscopy
- (D) Raman spectroscopy
- (E) Photonelectron spectroscopy
- 46. Which of the following complex has a highest oxidation state of metal?
- (A)  $(\eta^6 C_6 H_6)_2 Cr$
- (B)  $Na_2[Fe(CO)_4]$
- (C) Mn(CO)<sub>5</sub>Cl
- (D)  $Na[(\eta^5-C_5H_5)Fe(CO)_2]$
- (E) Ni(CO)<sub>4</sub>
- 47. If complex [W(Cp)<sub>2</sub>(CO)<sub>2</sub>] follows 18e- rule. What is Hapticity of Cp?
- (A) 5 and 5 (B) 3 and 5 (C) 3 and 3 (D) 1 and 5 (E) 1 and 3
- 48. Which of the following is mostly stable?
- (A)  $[AuF_2]^-$  (B)  $[AuCl_2]^-$  (C)  $[AuBr_2]^-$  (D)  $[AuI_2]^-$  (E)  $[Au(CN)_2]^-$
- 49. The intense red of an [Fe(phen)<sub>3</sub>]<sup>2+</sup> (shown below) solution is replaced by pale blue when cerium(IV) sulfate is added to it. Which of the following statements about this observation is correct?



- (A) The intense coloration of [Fe(phen)<sub>3</sub>]<sup>2+</sup> is due to inter-ligand charge transfer.
- (B) The electron count of  $[Fe(phen)_3]^{2+}$  is 20.
- (C) Addition of cerium(IV) sulfate causes the reduction of [Fe(phen)<sub>3</sub>]<sup>2+</sup> to [Fe(phen)<sub>3</sub>]<sup>+</sup>
- (D) The change of coloration upon the addition of cerium(IV) sulfate is due to lack of ligand to metal charge transfer transfer transfer.
- (E) None of statements are correct.
- 50. Which of the following is *closo*?
- (A) C<sub>2</sub>B<sub>3</sub>H<sub>5</sub>(CoCp)<sub>2</sub> (B) B<sub>4</sub>H<sub>10</sub> (C) C<sub>2</sub>B<sub>8</sub>H<sub>12</sub> (D) SB<sub>9</sub>H<sub>11</sub> (E) P<sub>4</sub>