Name	 	
Signature	 	

T.A.

This exam contains 23 questions on 11 numbered pages. Check now to make sure you have a complete exam. You have two hours to complete the exam. Determine the best answer to the first 20 questions and enter these on the special answer sheet. Also, circle your responses in this exam booklet.

Show all of your work and provide complete answers to questions 21, 22 and 23.

Useful Information:

$$N_A = 6.022 \times 10^{23}$$

$$E = -2.178 \times 10^{-18} \text{ J} (Z^2/n^2) = -1312 \text{ kJ/mol} (Z^2/n^2)$$

$$c = 2.998 \times 10^8 \text{ m/s}$$

$$h = 6.62608 \times 10^{-34} \text{ Js}$$

 $\Delta x \Delta p = h$

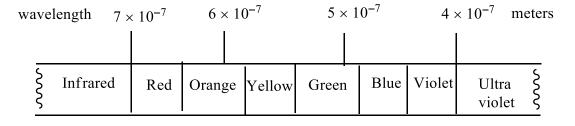
$$m_e = 9.10939 \ x \ 10^{-31} \ kg$$

$$E_n = \frac{n^2 h^2}{8mL^2}$$

$$E = \frac{h^2}{8m} \left[\frac{n_x^2}{L_x^2} + \frac{n_y^2}{L_y^2} + \frac{n_z^2}{L_z^2} \right]$$

$$1 \text{ pm} = 10^{-12} \text{ m}$$
 $1 \text{ nm} = 10^{-9} \text{ m}$

$$1 \text{ nm} = 10^{-9} \text{ m}$$



a) -1312

electron on this planet?

b) 0

n

e) 1312

Hour Exam I

2.	Con	sider a hydroge	en atom. How m	nany of	the following	are true?	
	I. II.	transition from	n = 4 to n = 2.			olves the same energeneers emit radiation of a	
		wavelength th	nan one from $n =$	= 5 to <i>n</i>	= 1.		
	III.	The wavelength of radiation required to cause the electron to go from energy level to energy level $n + 1$ decreases with increasing n .					
	IV.		s among the first			volve radiation in th	ne visible
	a) 0)	b) 1	c) 2		d) 3	e) 4
3.	Hov	w many of the fo	ollowing statem	ents are	true?		
	I. II.		2s electron in un			energy than the $3d$ of than the Z_{eff} for a $2s$	
	III.	•	of the lithium (Li) ator	n is higher in	energy than the 1s o	orbital of the
	IV.		fred light is grea	ater tha	n the energy o	f violet light.	
	a) 0)	b) 1	c) 2		d) 3	e) 4
4.	Whi	Thich of the following has the shortest wavelength ?					
	a) b)		raveling at 1% th	-	_		
	c)	An electron traveling at 10% the speed of light. A baseball traveling at 50 mph.					
	d) e)		veling at 100 m ame wavelength				
5.			ving set of quan d hydrogen aton		mbers describe	es an electron with t	he highest
	,	n = 3	l = 0		$m_1 = 0$		
	,		l=1				
			l = 1 $l = 2$				
	a) e)				$m_1 = 1$	with the same energy	
	-)		(a. a.) a				,

Imagine a planet for which the temperature is high enough so that the ground state for a hydrogen electron is n = 3. What is the ionization energy (in kJ/mol) for the hydrogen

d) 437

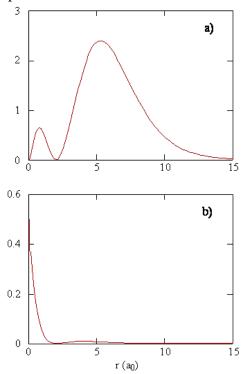
c) 146

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6. Consider a ground state quantum particle trapped in a one-dimensional box of length L. The most probable position for the quantum particle is at x =

- a) L
- b) L/2
- c) L/3
- d) 0
- e) The particle is equally likely to be found anywhere in the box.

7. The plots below are for the same orbital in the hydrogen atom. Which of the following statements correctly describes these plots?



- a) The plots are for the 1s orbital in hydrogen. The top plot is the electron radial probability distribution $(4\pi r^2 R^2 \text{ vs distance from the nucleus})$ and the bottom plot is the probability density of the electron (R^2 vs. distance from the nucleus).
- b) The plots are for the 2s orbital in hydrogen. The top plot is the electron radial probability distribution $(4\pi r^2 R^2)$ vs distance from the nucleus and the bottom plot is the probability density of the electron (R^2 vs. distance from the nucleus).
- c) The plots are for the 2s orbital in hydrogen. The top plot is the probability density of the electron (R^2 vs. distance from the nucleus) and the bottom plot is the electron radial probability distribution ($4\pi r^2 R^2$ vs distance from the nucleus).
- d) The plots are for a 2p orbital in hydrogen. The top plot is the electron radial probability distribution ($4\pi r^2 R^2$ vs distance from the nucleus) and the bottom plot is the probability density of the electron (R^2 vs. distance from the nucleus).
- e) The plots are for a 2p orbital in hydrogen. The top plot is the probability density of the electron (R^2 vs. distance from the nucleus) and the bottom plot is the electron radial probability distribution ($4\pi r^2 R^2$ vs distance from the nucleus).

8. What is the expected ground state electron configuration for the most stable ion of bromine in an ionic compound?

- a) $[Ar] 4s^2 4d^{10} 4p^5$
- b) $[Ar] 4s^2 3d^{10} 4p^5$
- c) $[Ar] 4s^2 4d^{10}4p^6$
- d) $[Ar] 4s^2 3d^{10} 4f^{14} 4p^5$
- e) $[Ar] 4s^2 3d^{10} 4p^6$

9. The hybridization of the **central nitrogen atom** in the N₂O molecule is

- a) sp
- b) sp^2
- c) sp^3
- d) dsp^3
- e) The hybridization can vary depending on the chosen resonance structure of the molecule.

10. Cyanogen chloride is a highly toxic chemical with the formula NCCl (the carbon atom is central). Predict the hybridization for each of the atoms in the molecule.

	N	C	C1
a)	sp^2	sp	sp^3
a) b)	sp	sp^2	sp^2
c) d)	sp^3	sp^2	sp^2
d)	sp^3	sp^3	sp^3
e)	sp	sp	sp^3

11. Compare the ionization energy of the nitrogen (N) atom to the oxygen (O) atom, nitrogen gas (N_2) and the N_2 ion. In each case, which species has the **highest ionization energy**?

	N vs. O	$N \text{ vs. } N_2$	$N \text{ vs } N_2^-$
a)	N	N_2	N
b)	O	N_2	N
c)	N	N	N
d)	O	N	N_2^-
e)	N	N_2	N_2^-

12. The F-F bond in $F_2(g)$ is much weaker than the B-B bond in $B_2(g)$. Why is this?

- a) The fluorine atom is much more electronegative than the boron atom.
- b) The boron atom is smaller than the fluorine atom.
- c) According to MO theory, the bond order for F_2 is less than that for B_2 .
- d) The electron-electron repulsions are greater in F_2 than in B_2 .
- e) The B-B bond in $B_2(g)$ is nonpolar.

13. Which of the following bonds gets **stronger** if an electron is **added**?

- a) B_2
- b) O₂⁻
- c) N_2
- d) F_2^+
- e) All the bonds get weaker.

14. Consider the description: "Two wave functions of similar energy in a given atom overlap to give a new wavefunction that solves the Schrödinger equation. This new wave function is able to overlap with the wavefunction in a different atom to form a molecule."

This description best describes:

- Simple valence bond theory a)
- Localized electron model b)
- Hybridization c)
- Molecular orbital theory d)
- Crystal field theory e)
- Which of the following compounds requires the *cis/trans* designation when naming?
 - pentaamminebromocobalt(II) chloride a)
 - diamminedichlorocobalt(II) (tetrahedral) **b**)
 - diamminedichloroplatinum(II) (square planar) c)
 - $[Co(en)_3]Cl_2$ d)
 - At least two of the above (a-d) require the *cis/trans* designation when naming. e)
- 16. Which of the following coordination compounds will form a precipitate when treated with an aqueous solution of AgNO₃?
 - a) $[Cr(NH_3)_3Cl_3]$
 - b) [Cr(NH₃)Cl]NO₃
 - c) Na₃[CrCl₆]
 - d) $[Cr(NH_3)_6]Cl_3$
 - e) At least two of the above will form a precipitate when treated with an aqueous solution of $AgNO_3$.
- Consider the complex ion $[Co(en)_2Cl_2]^+$, where en = ethylenediamine = NH₂CH₂CH₂NH₂. How many geometric isomers and optical isomers does this complex ion exhibit?

	Geometric isomers	Optical isomers
a)	0	0
b)	2	2
c)	2	3
d)	3	2
e)	4	4

- 18. You discover a new ligand (congratulations!) and wish to place it on the spectrochemical series. Which of the following would be a good choice to determine if the ligand were a weak-field or strong-field ligand by determining the number of unpaired electrons?
 - a) Cr²⁺
- b) V^{2+} c) Ni^{2+}
- d) Zn^{2+}
- e) Cu⁺

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19. The complex ions of Zn^{2+} are colorless in solution. The most likely explanation for this is

- a) the Zn^{2+} complex ion is paramagnetic.
- b) the *d*-orbital splitting in such complexes is outside of the visible region in the spectrum.
- c) the *d*-orbital splitting in such complexes allows for the absorbing of all wavelengths in the visible region.
- d) the Zn^{2+} ion is a d^{10} ion.
- e) only strong-field ligands can bind with the Zn^{2+} ion.
- 20. Consider the complex ions $[Co(A)_6]^{2+}$, $[Co(A)_6]^{3+}$, $[Co(B)_6]^{2+}$, and $[Co(C)_6]^{3+}$. The ligands A, B, and C are all neutral. The number of unpaired electrons for each of the complex ions (respectively) are found to be 3, 0, 1, 4. Rank the ligands from **strongest field to weakest field.**
 - a) A, B, C
 - b) C, B, A
 - c) B, C, A
 - d) A, C, B
 - e) B, A, C