

Answers to self-tests and exercises

Please note that just the final numerical solutions and short-answer solutions are supplied. For full worked solutions, please refer to the Solutions Manual by Adam Hascall (www.wiley.com/go/AdamHascall).

CHAPTER 1

Self-tests

ST1.1 $177900\text{eV}^{-1} = 1.23 \times 10^7\text{ m}^{-1}$ or 12300 nm .

ST1.2 All set of orbitals, 5 orbitals.

ST1.3 0.

ST1.4 7p_z.

ST1.5 The s electrons already present in the regular shell increasing electrons move strongly away from other p electrons already present in the regular shell increasing p electrons because the increasing p electrons give little in-bond control.

ST1.6 $\text{Na}^+/\text{Ar}/\text{F}^-/\text{He}^+$, $\text{Na}^{10+}/\text{Ar}^{10-}/\text{F}^-$

ST1.7 Period 4, Groups 2, 3, Block.

ST1.8 Conversely, going down a group the atomic radius increases and the first ionisation energy generally decreases.

ST1.9 Group 14.

ST1.10 The electrons gain positive (+) or (-) in incompatibility.

ST1.11 None.

Worked exercises

1.1 $20\text{ MeV}/(2 \times 10^6\text{ eV}) = 20 \times 10^6\text{ eV}/(2 \times 10^6\text{ eV}) = 10/2 = 5$

1.2 Get the numbers.

One thousand m^2 blocks radius, $r_0 = 100\text{ fm} = 10^{-15}\text{ m}$.

1.3 $20\text{ MeV} = 20 \times 10^6\text{ eV}$, $m = 80 = 1.6 \times 10^{-27}\text{ kg}$, $c = 3 \times 10^8\text{ m s}^{-1}$.

1.4 0.582 eV. The discrepancy is due to the relativistic effect.

1.5 $14.0\text{ GeV} = 1.4 \times 10^9\text{ eV}$

1.6 $1652.5\text{ nm}, 1.5937 \times 10^{-19}\text{ J}$

1.7 $1.5234 \times 10^{-10}\text{ m}^{-1}$, 656.3 nm

$$\frac{1}{\lambda} = 2c \left(\frac{1}{E^2} - \frac{1}{h\nu^2} \right) = 1.49974 \times 10^7\text{ m}^{-1}$$

$$\frac{1}{\lambda} = 2c \left(\frac{1}{E^2} - \frac{1}{h\nu^2} \right) = 1.02298 \times 10^7\text{ m}^{-1}$$

$$\frac{1}{\lambda} = 2c \left(\frac{1}{E^2} - \frac{1}{h\nu^2} \right) = 10.2947 \times 10^6\text{ m}^{-1}$$

$$\frac{1}{\lambda} = 2c \left(\frac{1}{E^2} - \frac{1}{h\nu^2} \right) = 18.27005 \times 10^6\text{ m}^{-1}$$

When given values of n , the angular momentum quantum number, it can assume all integer values from 0 to $n-1$.

n^2 (e.g., $n^2 = 1$ for $n=1$, $n^2 = 4$ for $n=2$, etc.)

n	l	m_l	Orbital designations	Number of orbitals
1	0	+1, 0, -1	2p	2
2	0	+1, 0, -1	3d	5
3	0	+1, 0, -1	4s	1
3	1	+1, 0, -1, -2	4p	3

When $n = 3$, $l = 1$ after the 3s orbital and $m_l = -1, 0, +1$, 3p₁, 3p₀, 3p₋₁, which represent the three orbitals that complete the 3d subshell. The 3d orbitals represent the start of the transition metals, starting with Cr and ending with Cu.

Comparing the plots for the 1s (Figure 1.11) and the off-axis 1s (Figure 1.12) orbitals, the radial distribution function for $n = 1$ is centred on a single maximum, and that for $n = 2$ consists of two maxima and a minimum (at $r = 2a_0/2$) for the higher-energy 2s subshell. The presence of the node at $r = 2a_0/2$ for 2s₀ causes the presence of the two maxima and the minimum in the 2p radial distribution function. The off-axis off- z radial mode for 2p₀ requires that the 2p radial distribution function has only a single maximum.

Solutions Manual Inorganic Chemistry Huheey

Ian Pickup

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Inorganic Chemistry: Principles of Structure and Reactivity, 4e Huheey,Medhi, Inorganic Chemistry Principles of Structure and Reactivity 4e **Inorganic Chemistry** Geoffrey Rayner-Canham,Tina Overton,2014-03-28 The Student Solution Manual includes the worked solutions to all of the odd numbered problems found in Descriptive Inorganic Chemistry sixth edition **Catalog of Copyright Entries. Third Series** Library of Congress. Copyright Office,1975 **Cumulated Index to the Books**,1972 **Scientific and Technical Books in Print**,1972 **Frontier Orbitals** Nguyen Trong Anh,2007-04-30 Written by one of the pioneers of the field Frontier Orbitals is an essential practical guide to the successes and limitations of this theory Applications are classified by chemical criteria competition between reagents sites or reaction trajectories The steps involved in solving each problem such as the choice of model the calculation of molecular orbitals and the interpretation of results are explained Numerous exercises are found throughout the text and the full solution and references are given in each case An extensive listing of MO s is also given to allow those without access to a computer to work out the exercises Practical advice is given for those wishing to do their own calculations Frontier Orbitals is aimed at experimentalists who are well versed in organic chemistry but have little or no understanding of quantum mechanics A greater emphasis is put on chemistry than on quantum mechanics and the intelligent use of the rules rather than their mathematical derivation

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