CHEMISTRY ATOMS FIRST **5th Edition**





Fundamental Constants	
Avogadro's number (N_A)	6.0221413×10^{23}
Electron charge (e)	$1.6022 \times 10^{-19} \text{ C}$
Electron mass	$9.109387 \times 10^{-28} \text{ g}$
Faraday constant (F)	96,485.3 C/mol e ⁻
Gas constant (R)	0.08206 L \cdot atm/K \cdot mol
	8.314 J/K · mol
	62.36 L \cdot torr/K \cdot mol
	1.987 cal/K · mol
Planck's constant (h)	$6.6256 \times 10^{-34} \text{ J} \cdot \text{s}$
Proton mass	$1.672623 \times 10^{-24} \text{ g}$
Neutron mass	$1.674928 \times 10^{-24} \text{ g}$
Speed of light in a vacuum (c)	2.99792458×10^8 m/s

Some Prefixes Used with SI Units								
tera (T)	10^{12}	centi (c)	10^{-2}					
giga (G)	10^{9}	milli (m)	10^{-3}					
mega (M)	10^{6}	micro (µ)	10^{-6}					
kilo (k)	10^{3}	nano (n)	10 ⁻⁹					
deci (d)	10^{-1}	pico (p)	10^{-12}					

Useful Conversion Factors and Relationships						
1 lb = 453.6 g						
1 in = 2.54 cm (exactly)						
1 mi = 1.609 km						
1 km = 0.6215 mi						
$1 \text{ pm} = 1 \times 10^{-12} \text{ m} = 1 \times 10^{-10} \text{ cm}$						
1 atm = 760 mmHg = 760 torr = 101,325 N/m ² = 101,325 Pa						
1 cal = 4.184 J (exactly)						
$1 L \cdot atm = 101.325 J$						
$1 J = 1 C \times 1 V$						
$?^{\circ}C = (^{\circ}F - 32^{\circ}F) \times \frac{5^{\circ}C}{9^{\circ}F}$						
$?^{\circ}F = \frac{9^{\circ}F}{5^{\circ}C} \times (^{\circ}C) + 32^{\circ}F$						
$?\mathbf{K} = (^{\circ}\mathbf{C} + 273.15^{\circ}\mathbf{C})\left(\frac{1\mathbf{K}}{1^{\circ}\mathbf{C}}\right)$						



	г				-	ر	1	~	د د	4	-	v	2	9	>	L.	·		9		٢	-
			18		Helium 4.003	Ne	Neon 20.18	Ar^{18}	Argon 39.95	${ m Kr}^{36}$	Krypton 83.80	Xe ⁵⁴	Xenon 131.3	⁸⁶ Rn	Radon (222)	0 8 0	Oganesson (294)		Lu	Lutetium 175.0	L^{103}	Lawrencium (262)
					17	٥ĹT	Fluorine 19.00	CI 1	Chlorine 35.45	Br 33	Bromine 79.90	I	Iodine 126.9	⁸⁵ At	Astatine (210)	T_{S}^{117}	Tennessine (293)		Yb	Ytterbium 173.0	N0	Nobelium (259)
	group				16	∞O	Oxygen 16.00	°5 N≊	Sulfur 32.07	$\mathbf{S}^{34}_{\mathbf{C}}$	Selenium 78.96	Te	Tellurium 127.6	Po Po	Polonium (209)	$L_{\rm V}^{116}$	Livermorium (291)		Tm	Thulium 168.9	Md	Mendelevium (258)
	Main §				15	۲Z	Nitrogen 14.01	₽ 5	Phosphorus 30.97	³³ AS	Arsenic 74.92	Sb Sb	Antimony 121.8	83 Bi	Bismuth 209.0	M_{c}^{115}	Moscovium (289)		Er 88	Erbium 167.3	Fm ¹⁰⁰	Fermium (257)
					14	٥C	Carbon 12.01	Si ¹⁴	Silicon 28.09	Ge ³²	Germanium 72.64	Sn Sn	Tin 118.7	Pb Pb	Lead 207.2	H ¹¹⁴	Flerovium (287)		Ho ⁶⁷	Holmium 164.9	ES %	Einsteinium (252)
					13	s B ∞	Boron 10.81	${\rm AI}^{13}$	Aluminum 26.98	Ga ³¹	Gallium 69.72	49 In	Indium 114.8	Π^{81}	Thallium 204.4	Nh ¹¹³	Nihonium (285)		Dy ⁶⁶	Dysprosium 162.5	Cf ³⁸	Californium (252)
	L							,	12	Zn ³⁰	Zinc 65.41	Cd ⁴⁸	Cadmium 112.4	Hg ⁸⁰	Mercury 200.6	Cn	Copernicium (285)		Tb	Terbium 158.9	⁹⁷ Bk	Berkelium (249)
									11	Cu Cu	Copper 63.55	${\rm Ag}^{47}$	Silver 107.9	Au	Gold 197.0	Rg	Roentgenium (280)		64 Gd	Gadolinium 157.3	Cm %	Curium (247)
nents	nents								10	${ m Ni}^{28}$	Nickel 58.69	\mathbf{Pd}^{46}	Palladium 106.4	\mathbf{Pt}^{78}	Platinum 195.1	$\mathbf{D}^{110}_{\mathbf{S}}$	Darmstadtium (281)		Eu	Europium 152.0	Am	Americium (243)
e Eler					ymbol	verage omic mass			6	C_0^{27}	Cobalt 58.93	45 Rh	Rhodium 102.9	TT Ir	Iridium 192.2	${ m Mt}^{109}$	Meitnerium (276)		Sm	Samarium 150.4	Pu	Plutonium (239)
of the				Key	C el	2.01 + A	element	n metals	∞	$\mathrm{F}^{26}_{\mathbf{C}}$	Iron 55.85	⁴⁴ Ru	Ruthenium 101.1	\mathbf{OS}^{76}	Osmium 190.2	HS ¹⁰⁸	Hassium (277)		Pm	Promethium (145)	Np ⁹³	Neptunium (237)
Table					lber	ame	An e	Transitic	7	Mn	Manganese 54.94	T^{43}_{C}	Technetium (98)	Re 75	Rhenium 186.2	Bh	Bohrium (272)		Nd ⁶⁰	Neodymium 144.2	C ²²	Uranium 238.0
odic 7					Atomic num	Ż			9	${\rm Cr}^{24}$	Chromium 52.00	${ m Mo}^{42}$	Molybdenum 95.94	74 W	Tungsten 183.8	No 60	Seaborgium (266)		\mathbf{Pr}^{59}	Praseodymium 140.9	Pa	Protactinium 231.0
Peri					P				5	V ²³	Vanadium 50.94	${\rm Nb}^{41}$	Niobium 92.91	Ta	Tantalum 180.9	Db Db	Dubnium (262)		Ce Ce	Cerium 140.1	[%] Th	Thorium 232.0
									4	Ti	Titanium 47.87	Z_{r}^{40}	Zirconium 91.22	Hf ⁷²	Hafnium 178.5	$\mathbf{R}^{104}_{\mathbf{f}}$	Rutherfordium (261)		hanoids 6		otinoide 7	
			120111						ю	Sc 21	Scandium 44.96	39 66	Yttrium 88.91	⁵⁷ La	Lanthanum 138.9		Actinium (227)		Lant			<u> </u>
	group	Groun nu	I drone		7	Be	Beryllium 9.012	M_g^{12}	Magnesium 24.31	Ca^{20}	Calcium 40.08	³⁸ Sr	Strontium 87.62	56 Ba	Barium 137.3	⁸⁸ Ra	Radium (226)			-10 -		spic
	Main	- -		Η	Hydrogen 1.008	Li Li	Lithium 6.941	Na	Sodium 22.99	61 K	Potassium 39.10	³⁷ Rb	Rubidium 85.47	Cs Cs	Cesium 132.9	$\mathbf{F}^{87}_{\mathbf{\Gamma}}$	Francium (223)		Metals	Nomo		Metall
	-	Period	E]	(4			4	-	v		9)							

List of the Elements with Their Symbols and Atomic Masses*

Element	Symbol	Atomic Number	Atomic Mass ⁺	Element	Symbol	Atomic Number	Atomic Mass ⁺
Actinium	Ac	89	(227)	Mendelevium	Md	101	(258)
Aluminum	Al	13	26.9815384	Mercury	Hg	80	200.592
Americium	Am	95	(243)	Molybdenum	Mo	42	95.95
Antimony	Sb	51	121.76	Moscovium	Mc	115	(289)
Argon	Ar	18	39.948	Neodymium	Nd	60	144.242
Arsenic	As	33	74.921595	Neon	Ne	10	20.1797
Astatine	At	85	(210)	Neptunium	Np	93	(237)
Barium	Ba	56	137.327	Nickel	Ni	28	58.6934
Berkelium	Bk	97	(249)	Niobium	Nb	41	92.90637
Bervllium	Be	4	9.0121831	Nihonium	Nh	113	(285)
Bismuth	Bi	83	208,9804	Nitrogen	N	7	14.00643
Bohrium	Bh	107	(272)	Nobelium	No	102	(259)
Boron	B	5	10.806	Oganesson	Og	118	(294)
Bromine	Br	35	79 904	Osmium	05	76	190.23
Cadmium	Cd	48	112.414	Oxygen	0	8	15 9994
Calcium	Ca	20	40.078	Palladium	Pd	46	106.42
Californium	Cf	98	(252)	Phosphorus	P	15	30.973762
Carbon	C	6	12 0096	Platinum	Pt	78	195 084
Cerium	C	58	1/0 116	Plutonium	Du	94	(230)
Cesium	C	55	132 005/152	Polonium	Po	94 84	(239) (200)
Chloring	Cl	17	25 446	Potossium	FU V	10	20,0082
Chromium	Cr	17	51.0061	Potassium	Г. Dr	19 50	39.0903
Chiomin	C	24	59 022104	Dramathium	Γ1 Den	59	(145)
Cobalt	C0	27	30.933194	Prometinium	PIII D.	01	(143)
Copernicium	Cn	112	(285)	Protactinium	Pa	91	231.03588
Copper	Cu	29	03.340	Radium	Ra	88	(220)
Curium	Cm	96	(247)	Radon	Rn	86	(222)
Darmstadtium	Ds	110	(281)	Rhenium	Re	/5	186.207
Dubnium	Db	105	(262)	Rhodium	Rh	45	102.90549
Dysprosium	Dy	66	162.5	Roentgenium	Rg	111	(280)
Einsteinium	Es	99	(252)	Rubidium	Rb	37	85.4678
Erbium	Er	68	167.259	Ruthenium	Ru	44	101.07
Europium	Eu	63	151.964	Rutherfordium	Rf	104	(261)
Fermium	Fm	100	(257)	Samarium	Sm	62	150.36
Flerovium	Fl	114	(287)	Scandium	Sc	21	44.955908
Fluorine	F	9	18.99840316	Seaborgium	Sg	106	(266)
Francium	Fr	87	(223)	Selenium	Se	34	78.96
Gadolinium	Gd	64	157.25	Silicon	Si	14	28.0855
Gallium	Ga	31	69.723	Silver	Ag	47	107.8682
Germanium	Ge	32	72.64	Sodium	Na	11	22.9876928
Gold	Au	79	196.96657	Strontium	Sr	38	87.62
Hafnium	Hf	72	178.486	Sulfur	S	16	32.065
Hassium	Hs	108	(277)	Tantalum	Та	73	180.94788
Helium	He	2	4.002602	Technetium	Tc	43	(98)
Holmium	Но	67	164.930328	Tellurium	Te	52	127.6
Hydrogen	Н	1	1.00784	Tennessine	Ts	117	(293)
Indium	In	49	114.818	Terbium	Tb	65	158.925354
Iodine	Ι	53	126.90447	Thallium	Tl	81	204.382
Iridium	Ir	77	192.217	Thorium	Th	90	232.0377
Iron	Fe	26	55.845	Thulium	Tm	69	168.934218
Krypton	Kr	36	83.798	Tin	Sn	50	118.71
Lanthanum	La	57	138.90547	Titanium	Ti	22	47.867
Lawrencium	Lr	103	(262)	Tungsten	W	74	183.84
Lead	Pb	82	207.2	Uranium	U	92	238.02891
Lithium	Li	3	6.941	Vanadium	V	23	50.9415
Livermorium	Lv	116	(291)	Xenon	Xe	54	131.293
Lutetium	Lu	71	174.9668	Ytterbium	Yb	70	173.045
Magnesium	Mg	12	24.3050	Yttrium	Y	39	88.90584
Manganese	Mn	25	54.938043	Zinc	Zn	30	65.409
Meitnerium	Mt	109	(276)	Zirconium	Zr	40	91.224

*These atomic masses show as many significant figures as are known for each element. The atomic masses in the periodic table are shown to four significant figures, which is sufficient for solving the problems in this book.

†Approximate values of atomic masses for radioactive elements are given in parentheses.



ISTUDY

Chemistry

ATOMS FIRST

FIFTH EDITION

Julia Burdge COLLEGE OF WESTERN IDAHO

> Jason Overby COLLEGE OF CHARLESTON





CHEMISTRY: ATOMS FIRST

Published by McGraw Hill LLC, 1325 Avenue of the Americas, New York, NY 10019. Copyright ©2024 by McGraw Hill LLC. All rights reserved. Printed in the United States of America. No part of this publication may be reproduced or distributed in any form or by any means, or stored in a database or retrieval system, without the prior written consent of McGraw Hill LLC, including, but not limited to, in any network or other electronic storage or transmission, or broadcast for distance learning.

° ° ° °

•

Some ancillaries, including electronic and print components, may not be available to customers outside the United States.

This book is printed on acid-free paper.

1 2 3 4 5 6 7 8 9 LWI 27 26 25 24 23

ISBN 978-1-266-13678-8 MHID 1-266-13678-9

Cover Image: noLimit46/iStock/Getty Images

All credits appearing on page or at the end of the book are considered to be an extension of the copyright page.

The Internet addresses listed in the text were accurate at the time of publication. The inclusion of a website does not indicate an endorsement by the authors or McGraw Hill LLC, and McGraw Hill LLC does not guarantee the accuracy of the information presented at these sites.





About the Authors



McGraw Hill

Julia Burdge received her Ph.D. (1994) from the University of Idaho in Moscow, Idaho. Her research and dissertation focused on instrument development for analysis of trace sulfur compounds in air and the statistical evaluation of data near the detection limit.

In 1994 she accepted a position at The University of Akron in Akron, Ohio, as an assistant professor and director of the Introductory Chemistry program. In the year 2000, she was tenured and promoted to associate professor at The University of Akron on the merits of her teaching, service, and research in chemistry education. In addition to directing the general chemistry program and supervising the teaching activities of graduate students, she helped establish a future-faculty development program and served as a mentor for graduate students and post-doctoral associates. In 2008, Julia relocated back to the northwest to be near family. She lives in Boise, Idaho; and she holds an affiliate faculty position as associate professor in the Chemistry Department at the University of Idaho and teaches general chemistry at the College of Western Idaho.

In her free time, Julia enjoys horseback riding, precious time with her three children, and quiet time at home with Erik Nelson, her husband and best friend.



McGraw Hill

Jason Overby received his B.S. degree in chemistry and political science from the University of Tennessee at Martin. He then received his Ph.D. in inorganic chemistry from Vanderbilt University (1997) studying main group and transition metal metallocenes and related compounds. Afterwards, Jason conducted postdoctoral research in transition metal organometallic chemistry at Dartmouth College.

Jason began his academic career at the College of Charleston in 1999 as an assistant professor. Currently, he is an associate professor with teaching interests in general and inorganic chemistry. He is also interested in the integration of technology into the classroom, with a particular focus on adaptive learning. Additionally, he conducts research with undergraduates in inorganic and organic synthetic chemistry as well as computational organometallic chemistry.

In his free time, Jason enjoys boating, bowling, and cooking. On many weekends throughout the year, he can often be found on the deck of a pool working as a nationally certified USA Swimming official. He lives in South Carolina with his wife Robin and two daughters, Emma and Sarah.

Brief Contents

- 1 Chemistry: The Science of Change 2
- 2 Atoms and the Periodic Table 38
- **3** Quantum Theory and the Electronic Structure of Atoms 66
- 4 Periodic Trends of the Elements 114
- 5 Ionic and Covalent Compounds 162
- 6 Representing Molecules 210
- 7 Molecular Geometry, Intermolecular Forces, and Bonding Theories 246
- 8 Chemical Reactions 308
- 9 Chemical Reactions in Aqueous Solutions 352
- **10** Energy Changes in Chemical Reactions 416
- **11** Gases 472
- **12** Liquids and Solids 532
- 13 Physical Properties of Solutions 576
- 14 Chemical Kinetics 622
- **15** Entropy and Gibbs Energy 686
- 16 Chemical Equilibrium 724
- **17** Acids, Bases, and Salts 788
- **18** Acid-Base Equilibria and Solubility Equilibria 850
- **19** Electrochemistry 906
- 20 Nuclear Chemistry 954
- 21 Environmental Chemistry 988
- 22 Coordination Chemistry 1016
- 23 Organic Chemistry 1040
- 24 Modern Materials 1094
- 25 Online Only Chapter: Nonmetallic Elements and Their Compounds
- 26 Online Only Chapter: Metallurgy and the Chemistry of Metals
 - Appendix 1 Mathematical Operations A-1
 - Appendix 2 Thermodynamic Data at 1 atm and 25°C A-6
 - Appendix 3 Solubility Product Constants at 25°C A-13
 - Appendix 4 Dissociation Constants for Weak Acids and Bases at 25°C A-15

viii

Contents

List of Applications xxii Preface xxiii Instructor and Student Resources xxxii Acknowledgments xxxiv

1 CHEMISTRY: THE SCIENCE OF CHANGE 2

1.1 The Study of Chemistry 3

- Chemistry You May Already Know 3 The Scientific Method 3
- Thinking Outside the Box: Tips for Success in Chemistry Class 5

1.2 Scientific Measurement 6

- SI Base Units 7 Mass 8 Temperature 8 Derived Units: Volume and Density 10 Environmental Aspects: Global Climate Change 12
- **1.3 Uncertainty in Measurement 13**
 - Significant Figures 13 Calculations with Measured Numbers 15
 Accuracy and Precision 17
- 1.4 Using Units and Solving Problems 19
 Conversion Factors 19 Dimensional Analysis—Tracking Units 20

1.5 Classification of Matter 23

- States of Matter 23 Mixtures 24 Thinking Outside the Box: How Many States of Matter Are There? 25
- 1.6 The Properties of Matter 26
 Physical Properties 26 Chemical Properties 26 Extensive and Intensive Properties 26

2 ATOMS AND THE PERIODIC TABLE 38

2.1 Atoms First 39

Subatomic Particles and Atomic Structure 40 Discovery of the Electron 40 • Radioactivity 42 • The Proton and the Nuclear Model of the Atom 43 • The Neutron 44

- 2.3 Atomic Number, Mass Number, and Isotopes 46
- 2.4 Nuclear Stability 48
 - Patterns of Nuclear Stability 48

2.5 Average Atomic Mass 50

• Thinking Outside the Box: Measuring Atomic Mass 51



Precision Graphics/McGraw Hill



Little Hand Images/Moment/Getty Images



Reprint Courtesy of IBM Corporation \bigcirc 2013

W IRM

2.6 The Periodic Table 52

• Environmental Aspects: Isolated Atoms and Stratospheric Ozone Depletion 54

2.7 The Mole and Molar Mass 54

 \bullet The Mole 54 \bullet Molar Mass 55 \bullet Interconverting Mass, Moles, and Numbers of Atoms 57

QUANTUM THEORY AND THE ELECTRONIC STRUCTURE OF ATOMS 66

.1 Energy and Energy Changes 67

• Forms of Energy 67 • Units of Energy 68

2 The Nature of Light 70

• Properties of Waves 70 • The Electromagnetic Spectrum 71 • The Double-Slit Experiment 72 • Environmental Aspects: Solar Energy 72

3.3 Quantum Theory 74

• Quantization of Energy 74 • Photons and the Photoelectric Effect 75 • Thinking Outside the Box: Everyday Occurrences of the Photoelectric Effect 76

3.4 Bohr's Theory of the Hydrogen Atom 79

• Atomic Line Spectra 80 • The Line Spectrum of Hydrogen 81

3.5 Wave Properties of Matter 87

• The de Broglie Hypothesis 87 • Diffraction of Electrons 89

3.6 Quantum Mechanics 90

- The Uncertainty Principle 90 The Schrödinger Equation 91
- The Quantum Mechanical Description of the Hydrogen Atom 92

3.7 Quantum Numbers 92

• Principal Quantum Number (*n*) 92 • Angular Momentum Quantum Number (*P*) 93 • Magnetic Quantum Number (m_{ℓ}) 93 • Electron Spin Quantum Number (m_s) 94

3.8 Atomic Orbitals 96

4

• *s* Orbitals 96 • *p* Orbitals 96 • *d* Orbitals and Other Higher-Energy Orbitals 97 • Energies of Orbitals 99

PERIODIC TRENDS OF THE ELEMENTS 114

4.1 Development of the Periodic Table 115

4.2 Electron Configurations 117

- Energies of Atomic Orbitals in Many-Electron Systems 117
- The Pauli Exclusion Principle 118 The Aufbau Principle 119
- Hund's Rule 119 General Rules for Writing Electron

Configurations 121 • Thinking Outside the Box: How do we know that *ns* orbitals fill before (n - 1)d orbitals? 122



Olga Popova/iStockphoto/Getty Images

х

- 4.3 Electron Configurations and the Modern Periodic Table 123
- 4.4 Effective Nuclear Charge 129
- 4.5 Periodic Trends in Properties of Elements 130
 - Atomic Radius 130 Ionization Energy 132 Electron Affinity 135
 - Metallic Character 138
- 4.6 Electron Configuration of lons 141
 - Ions of Main Group Elements 141 Ions of *d*-Block Elements 143
- 4.7 Ionic Radius 145

• Comparing Ionic Radius with Atomic Radius 145 • Isoelectronic Series 145 • Environmental Aspects: Lead in the Water in Flint, Michigan 147 • Thinking Outside the Box: Mistaking Strontium for Calcium 148

5 IONIC AND COVALENT COMPOUNDS 162

- 5.1 Compounds 163
- 5.2 Lewis Dot Symbols 163
- 5.3 Ionic Compounds and Bonding 165
- 5.4 Naming lons and lonic Compounds 169
 Formulas of lonic Compounds 170 Naming lonic Compounds 171
- 5.5 Covalent Bonding and Molecules 172
 Molecules 173 Molecular Formulas 175 Empirical Formulas 176
- 5.6 Naming Molecular Compounds 179

 Specifying Numbers of Atoms 179 • Compounds Containing Hydrogen 181 • Organic Compounds 182 • Environmental Aspects: CFC Nomenclature 183 • Thinking Outside the Box: Functional Groups 183

5.7 Covalent Bonding in Ionic Species 184

- Polyatomic Ions 184 Oxoacids 186 Hydrates 188 Familiar Inorganic Compounds 188
- 5.8 Molecular and Formula Masses 190
- 5.9 Percent Composition of Compounds 192
- 5.10 Molar Mass 193
 - Interconverting Mass, Moles, and Numbers of Particles 194

• Determination of Empirical Formula and Molecular Formula from Percent Composition 196

6 REPRESENTING MOLECULES 210

- 6.1 The Octet Rule 211
 - Lewis Structures 211 Multiple Bonds 214
- 6.2 Electronegativity and Polarity 215

• Electronegativity 216 • Dipole Moment, Partial Charges, and Percent Ionic Character 218



Keith Corrigan/Alamy Stock Photo



Ken Welsh/Pixtal/age fotostock

xi



- 6.4 Lewis Structures and Formal Charge 225
- 6.5 Resonance 228
- 6.6 Exceptions to the Octet Rule 230
 - Incomplete Octets 230 Odd Numbers of Electrons 231
 - Thinking Outside the Box: Species with Unpaired Electrons 232
 - Environmental Aspects: Tropospheric Ozone 233
 - Expanded Octets 233
- 7 7. 7. 7.

Jason Overby



7.1 Molecular Geometry 247

• The VSEPR Model 248 • Electron-Domain Geometry and Molecular Geometry 249 • Deviation from Ideal Bond Angles 253 • Geometry of Molecules with More Than One Central Atom 253

- 7.2 Molecular Geometry and Polarity 255
 - Environmental Aspects: Polarity and the Greenhouse Effect 259
- 7.3 Intermolecular Forces 260
 - Dipole-Dipole Interactions 260 Hydrogen Bonding 260
 - Dispersion Forces 262 Ion-Dipole Interactions 263
- 7.4 Valence Bond Theory 265
- 7.5 Hybridization of Atomic Orbitals 268

• Hybridization of s and p Orbitals 269 • Hybridization of s, p, and d Orbitals 272 • Profiles in Chemistry: Linus Pauling 273

- 7.6 Hybridization in Molecules Containing Multiple Bonds 276
- 7.7 Molecular Orbital Theory 283

• Bonding and Antibonding Molecular Orbitals 284 • σ Molecular Orbitals 284 • Thinking Outside the Box: Phases 285 • Bond Order 286 • π Molecular Orbitals 286 • Molecular Orbital Diagrams 288 • Thinking Outside the Box: Molecular Orbitals in Heteronuclear Diatomic Species 289

7.8 Bonding Theories and Descriptions of Molecules with Delocalized Bonding 291

CHEMICAL REACTIONS 308

8.1 Chemical Equations 309

8

Interpreting and Writing Chemical Equations 309 • Balancing
 Chemical Equations 311 • Patterns of Chemical Reactivity 314

8.2 Combustion Analysis 317

• Determination of Empirical Formula 318 • Profiles in Chemistry: Marie-Anne Paulze Lavoisier 319



LWA/Photodisc/Getty Images

xiii

8.3 Calculations with Balanced Chemical Equations 320

- Moles of Reactants and Products $\ 320$ - Mass of Reactants and Products $\ 322$

- 8.4 Limiting Reactants 323
 - Determining the Limiting Reactant 324 Reaction Yield 326
 Atom Economy 330 Environmental Aspects: Green Chemistry Metrics 331

8.5 Periodic Trends in Reactivity of the Main Group Elements 333
 General Trends in Reactivity 333 • Hydrogen (1s¹) 334 • Reactions of the Active Metals 334 • Reactions of Other Main Group Elements 335

Comparison of Group 1 and Group 11 Elements 339

9 CHEMICAL REACTIONS IN AQUEOUS SOLUTIONS 352

9.1 General Properties of Aqueous Solutions 353

• Electrolytes and Nonelectrolytes 353 • Strong Electrolytes and Weak Electrolytes 354

9.2 Precipitation Reactions 359

Solubility Guidelines for Ionic Compounds in Water 359 • Molecular Equations 361 • Ionic Equations 362 • Net Ionic Equations 362
Environmental Aspects: Softening Hard Water 364

9.3 Acid-Base Reactions 365

- Strong Acids and Bases 365 Brønsted Acids and Bases 365
- Acid-Base Neutralization 368

9.4 Oxidation-Reduction Reactions 370

- Oxidation Numbers 371 Oxidation of Metals in Aqueous Solutions 374 Balancing Simple Redox Equations 376
- Other Types of Redox Reactions 378

9.5 Concentration of Solutions 381

• Molarity 382 • Dilution 383 • Serial Dilution 387 • Thinking Outside the Box: Visible Spectrophotometry 388

- The pH Scale 390 Solution Stoichiometry 392
- 9.6 Aqueous Reactions and Chemical Analysis 394
 - Gravimetric Analysis 394 Acid-Base Titrations 396

10 ENERGY CHANGES IN CHEMICAL REACTIONS 416

- **10.1 Energy and Energy Changes 417**
- 10.2 Introduction to Thermodynamics 419
 - States and State Functions 420 The First Law of Thermodynamics 420
 - Work and Heat 421



liseykina/Shutterstock



Steve Gschmeissner/Science Source



- Reactions Carried Out at Constant Volume or at Constant Pressure 424
- Enthalpy and Enthalpy Changes 426 Thermochemical Equations 427
- 10.4 Calorimetry 429

• Specific Heat and Heat Capacity 430 • Constant-Pressure Calorimetry 431 • Constant-Volume Calorimetry 435 • Thinking Outside the Box: Heat Capacity of Calorimeters 438 • Environmental Aspects: Energy Density and Specific Energy 439

- 10.5 Hess's Law 440
- **10.6 Standard Enthalpies of Formation 442**
- 10.7 Bond Enthalpy and the Stability of Covalent Molecules 446
- 10.8 Lattice Energy and the Stability of Ionic Solids 450

 The Born-Haber Cycle 450 • Comparison of Ionic and Covalent Compounds 454

11 GASES 472

- 11.1 Properties of Gases 473
- 11.2 The Kinetic Molecular Theory of Gases 474
 - Molecular Speed 475 Diffusion and Effusion 478

11.3 Gas Pressure 479

Definition and Units of Pressure 479 • Calculation of Pressure 480
 Measurement of Pressure 480

11.4 The Gas Laws 482

• Boyle's Law: The Pressure-Volume Relationship 482 • Charles's and Gay-Lussac's Law: The Temperature-Volume Relationship 485

- Avogadro's Law: The Amount-Volume Relationship 487
- The Gas Laws and Kinetic Molecular Theory 489 The Combined Gas Law: The Pressure-Temperature-Amount-Volume Relationship 491

11.5 The Ideal Gas Equation 493

• Applications of the Ideal Gas Equation 495 • Environmental Aspects: Lead in Gasoline 497

11.6 Real Gases 498

• Factors That Cause Deviation from Ideal Behavior 498 • The van der Waals Equation 498 • van der Waals Constants 500

11.7 Gas Mixtures 502

- Dalton's Law of Partial Pressures 502 Mole Fractions 504
- Thinking Outside the Box: Decompression Injury 505

11.8 Reactions with Gaseous Reactants and Products 507

- Calculating the Required Volume of a Gaseous Reactant 507
- Determining the Amount of Reactant Consumed Using Change
- in Pressure 509 Using Partial Pressures to Solve Problems 509
- Profiles in Chemistry: Fritz Haber 514



Mike Lyvers/Moment/Getty Images

xν

12 LIQUIDS AND SOLIDS 532

12.1 The Condensed Phases 533

12.2 Properties of Liquids 534

• Surface Tension 534 • Viscosity 535 • Vapor Pressure of Liquids 535 • Boiling Point 539

12.3 Properties of Solids 540

• Melting Point 540 • Vapor Pressure of Solids 540 • Amorphous Solids 541 • Crystalline Solids 542 • Thinking Outside the Box: X-Ray Diffraction 546

12.4 Types of Crystalline Solids **549**

Ionic Crystals 549 • Covalent Crystals 551 • Molecular Crystals 552 • Metallic Crystals 553

12.5 Phase Changes 554

- Liquid to Vapor 554 Solid to Liquid 556 Solid to Vapor 558
- Environmental Aspects: Water Vapor in the Atmosphere 560
- 12.6 Phase Diagrams 561

13 PHYSICAL PROPERTIES OF SOLUTIONS 576

13.1 Types of Solutions 577

- 13.2 A Molecular View of the Solution Process 578
 - The Importance of Intermolecular Forces 578 Energy and Entropy in Solution Formation 580 Profiles in Chemistry: Alice Ball 582

13.3 Concentration Units 583

Molality 583 • Percent by Mass 583 • Comparison of Concentration Units 585

13.4 Factors That Affect Solubility 587

Temperature 588 • Pressure 588

13.5 Colligative Properties 590

- Vapor-Pressure Lowering 591 Boiling-Point Elevation 593
- Freezing-Point Depression 594 Osmotic Pressure 596 Electrolyte
- Solutions 597 Thinking Outside the Box: Intravenous Fluids 599
- Thinking Outside the Box: Fluoride Poisoning 600

13.6 Calculations Using Colligative Properties 601

13.7 Colloids 605

• Environmental Aspects: EPS Foam Bans 606

14 CHEMICAL KINETICS 622

- 14.1 Reaction Rates 623
- 14.2 Collision Theory of Chemical Reactions 623



Courtesy of Paul C. Canfield and Ian R. Fisher/ Ames Laboratory/U.S. Department of Energy



Pam Walker/Shutterstock



Jonathan Nourok/Getty Images



- Average Reaction Rate 625 Instantaneous Rate 630
- Stoichiometry and Reaction Rate 632
- 14.4 Dependence of Reaction Rate on Reactant Concentration 636The Rate Law 636 Experimental Determination of the Rate Law 636
- 14.5 Dependence of Reactant Concentration on Time 641
 - First-Order Reactions 642 Second-Order Reactions 647
- 14.6 Dependence of Reaction Rate on Temperature 650The Arrhenius Equation 651 Thinking Outside the Box: Surface
 - Area 655
- 14.7 Reaction Mechanisms 656
 - Elementary Reactions 657 Rate-Determining Step 658
 - Mechanisms with a Fast First Step 662 Experimental Support for
 - Reaction Mechanisms 664
- 14.8 Catalysis 665
 - Heterogeneous Catalysis 666 Homogeneous Catalysis 667
 - Enzymes: Biological Catalysts 667 Environmental Aspects: Catalytic Converters 668



Saran Poroong/Alamy Stock Photo



15.1 Spontaneous Processes 687

15.2 Entropy 688

• A Qualitative Description of Entropy 688 • A Quantitative Definition of Entropy 688

686

15.3 Entropy Changes in a System 690

• Calculating $\Delta S_{\rm sys}~690$ • Standard Entropy, S° 691 • Qualitatively Predicting the Sign of $\Delta S_{\rm sys}^\circ~694$

15.4 Entropy Changes in the Universe 699

- Calculating ΔS_{surr} 700 The Second Law of Thermodynamics 700
- Thinking Outside the Box: Thermodynamics and Living Systems 703
- Environmental Aspects: Thermodynamic Limits on Energy
- Conversion 703 The Third Law of Thermodynamics 704
- 15.5 Predicting Spontaneity 706
 - Gibbs Energy Change, $\Delta {\it G}~$ 706 Standard Gibbs Energy Changes,
 - ΔG° 709 Using ΔG and ΔG° to Solve Problems 710
- 15.6 Thermodynamics in Living Systems 713

16 CHEMICAL EQUILIBRIUM 724

- 16.1 The Concept of Equilibrium 725
- 16.2 The Equilibrium Constant 727
 - Calculating Equilibrium Constants 728 Magnitude of the Equilibrium Constant 730



Harry Kikstra/Moment/Getty Images

16.3 Equilibrium Expressions 733

• Heterogeneous Equilibria 733 • Manipulating Equilibrium Expressions 734 • Gaseous Equilibria 738

- 16.4 Chemical Equilibrium and Gibbs Energy 741
 - Using Q and K to Predict the Direction of Reaction 741

• Relationship Between ΔG and ΔG° 743 • Relationship Between ΔG° and K 745

16.5 Calculating Equilibrium Concentrations 749

16.6 Le Châtelier's Principle: Factors That Affect Equilibrium 758

Addition or Removal of a Substance 758 · Changes in Volume and Pressure 761 · Changes in Temperature 762 · Catalysis 764
Environmental Aspects: Bioaccumulation of Toxins 764 · Thinking Outside the Box: Biological Equilibria 765

17 ACIDS, BASES, AND SALTS 788

17.1 Brønsted Acids and Bases 789

- 17.2 Molecular Structure and Acid Strength 792
 Hydrohalic Acids 793 Oxoacids 793 Carboxylic Acids 794
 Profiles in Chemistry: St. Elmo Brady 795 Thinking Outside the Box: Substituent Electronegativity and Acid Strength 795
- 17.3 The Acid-Base Properties of Water 797

17.4 The pH and pOH Scales 799

17.5 Strong Acids and Bases 801

Strong Acids 801 • Strong Bases 803

17.6 Weak Acids and Acid Ionization Constants 806

- The Ionization Constant, K_a 806 Calculating pH from K_a 806
- Percent Ionization 812 Environmental Aspects: Acid Rain 812
- Using pH to Determine K_a 814

17.7 Weak Bases and Base Ionization Constants 816

- The Ionization Constant, $K_{\rm b}$ 816 Calculating pH from $K_{\rm b}$ 816
- Using pH to Determine $K_{\rm b}$ 818

17.8 Conjugate Acid-Base Pairs 819

• The Strength of a Conjugate Acid or Base 819 • The Relationship Between $K_{\rm a}$ and $K_{\rm b}$ of a Conjugate Acid-Base Pair 820

17.9 Diprotic and Polyprotic Acids 823

17.10 Acid-Base Properties of Salt Solutions 826

• Basic Salt Solutions 826 • Acidic Salt Solutions 827 • Neutral Salt Solutions 829 • Salts in Which Both the Cation and the Anion Hydrolyze 831

17.11 Acid-Base Properties of Oxides and Hydroxides 832

• Oxides of Metals and Nonmetals 832 • Basic and Amphoteric Hydroxides 833

17.12 Lewis Acids and Bases 834



Aleksandra H. Kossowska/Shutterstock



Lisa Stokes/Moment Open/Getty Images

18 ACID-BASE EQUILIBRIA AND SOLUBILITY EQUILIBRIA 850

18.1 The Common Ion Effect 851

18.2 Buffer Solutions 853

• Calculating the pH of a Buffer 853 • Preparing a Buffer Solution with a Specific pH 859 • Environmental Aspects: Ocean Acidification 860

18.3 Acid-Base Titrations 861

• Strong Acid–Strong Base Titrations 861 • Weak Acid–Strong Base Titrations 862 • Strong Acid–Weak Base Titrations 867 • Acid-Base Indicators 869

18.4 Solubility Equilibria 872

- Solubility Product Expression and K_{sp} 872
- Calculations Involving K_{sp} and Solubility 872
- Predicting Precipitation Reactions 876

18.5 Factors Affecting Solubility 878

• The Common Ion Effect 878 • pH 879 • Complex Ion Formation 883 • Thinking Outside the Box: Equilibrium and Tooth Decay 884

18.6 Separation of Ions Using Differences in Solubility 888

• Fractional Precipitation 889 • Qualitative Analysis of Metal lons in Solution 890

19 ELECTROCHEMISTRY 906

- 19.1 Balancing Redox Reactions 907
- 19.2 Galvanic Cells 911
- 19.3 Standard Reduction Potentials 914
- 19.4 Spontaneity of Redox Reactions Under Standard State Conditions 922
- 19.5 Spontaneity of Redox Reactions Under Conditions Other Than Standard State 926
 - The Nernst Equation 926 Concentration Cells 928

19.6 Batteries 931

- Dry Cells and Alkaline Batteries 931 Lead Storage Batteries 932
- Lithium-Ion Batteries 932 Fuel Cells 933 Thinking Outside the Box: Molten Carbonate Fuel Cells 934

19.7 Electrolysis 934

• Electrolysis of Molten Sodium Chloride 935 • Electrolysis of Water 935 • Environmental Aspects: Artificial Leaves as a Renewable Energy Source 936 • Electrolysis of an Aqueous Sodium Chloride Solution 937 • Quantitative Applications of Electrolysis 937

19.8 Corrosion 940



National Geographic Image Collection/Alamy Stock Photo

xix

20 NUCLEAR CHEMISTRY 954

- 20.1 Nuclei and Nuclear Reactions 955
- 20.2 Nuclear Stability 957
 - Types of Nuclear Decay 957 Nuclear Binding Energy 957
- 20.3 Natural Radioactivity 961
 Kinetics of Radioactive Decay 961 Dating Based on Radioactive Decay 962
- 20.4 Nuclear Transmutation 965
- 20.5 Nuclear Fission 967
 - Environmental Aspects: Nuclear Power, Waste, and Recycling 974
- 20.6 Nuclear Fusion 974
- 20.7 Uses of Isotopes 976
 - Chemical Analysis 976 Thinking Outside the Box: Nuclear Medicine 977 Isotopes in Medicine 977
- 20.8 Biological Effects of Radiation 979

21 ENVIRONMENTAL CHEMISTRY 988

- 21.1 Earth's Atmosphere 989
- 21.2 Phenomena in the Outer Layers of the Atmosphere 992
 Aurora Borealis and Aurora Australis 992
 The Mystery Glow of Space Shuttles 993
- 21.3 Depletion of Ozone in the Stratosphere 994Polar Ozone Holes 996
- 21.4 Volcanoes 998
- 21.5 The Greenhouse Effect 999
- 21.6 Acid Rain 1003
- 21.7 Photochemical Smog 1006
- 21.8 Indoor Pollution 1007

 The Risk from Radon 1007 • Carbon Dioxide and Carbon Monoxide 1009 • Formaldehyde 1010

22 COORDINATION CHEMISTRY 1016

- 22.1 Coordination Compounds 1017
 - Properties of Transition Metals 1017 Ligands 1019
 - Nomenclature of Coordination Compounds 1021
 - Thinking Outside the Box: Chelation Therapy 1023
- 22.2 Structure of Coordination Compounds 1024
- 22.3 Bonding in Coordination Compounds: Crystal Field Theory 1027
 - Crystal Field Splitting in Octahedral Complexes 1027 Color 1028
 - Magnetic Properties 1030 Tetrahedral and Square-Planar Complexes 1032



Wellcome Trust



Chaleephoto/Shutterstock



Best View Stock/Alamy Stock Photo



XX



Vlad61/Shutterstock



- 22.4 Reactions of Coordination Compounds 1033
- 22.5 Applications of Coordination Compounds 1033

23 ORGANIC CHEMISTRY 1040

23.1 Why Carbon Is Different 1041

23.2 Classes of Organic Compounds 1043

• Basic Nomenclature 1047 • Molecules with Multiple

Substituents 1050 • Molecules with Specific Functional Groups 1051

23.3 Representing Organic Molecules 1054

- Condensed Structural Formulas 1054 Kekulé Structures 1055
- Bond-Line Structures 1056 Resonance 1057

23.4 Isomerism 1061

- Constitutional Isomerism 1061 Stereoisomerism 1061
- Thinking Outside the Box: Thalidomide Analogues 1065

23.5 Organic Reactions 1066

- Addition Reactions 1066 Substitution Reactions 1068
- Other Types of Organic Reactions 1072

23.6 Organic Polymers 1074

- Addition Polymers 1075 Condensation Polymers 1076
- Biological Polymers 1077

24 MODERN MATERIALS 1094

24.1 Polymers 1095

Addition Polymers 1095 • Condensation Polymers 1101 • Thinking
 Outside the Box: Electrically Conducting Polymers 1104

24.2 Ceramics and Composite Materials 1104

Ceramics 1104 · Composite Materials 1106

24.3 Liquid Crystals 1106

24.4 Biomedical Materials 1109

Dental Implants 1110 • Soft Tissue Materials 1111 • Artificial Joints 1112

24.5 Nanotechnology 1112

- Graphite, Buckyballs, and Nanotubes 1113
- 24.6 Semiconductors 1115
- 24.7 Superconductors 1119

25 NONMETALLIC ELEMENTS AND THEIR COMPOUNDS (ONLINE ONLY)

25.1 General Properties of Nonmetals 1127

25.2 Hydrogen 1128



Zoran Karapancev/Shutterstock



Lorelle Mansfield/NIST/Science Source



xxi

- Binary Hydrides 1128 Isotopes of Hydrogen 1130
- Hydrogenation 1131 The Hydrogen Economy 1131
- 25.3 Carbon 1132
- 25.4 Nitrogen and Phosphorus 1133
 - Nitrogen 1133 Phosphorus 1136
- **Oxygen and Sulfur 1139**Oxygen 1139 Sulfur 1141 Thinking Outside the Box:
 - Arsenic 1145
- 25.6 The Halogens 1145
 - Preparation and General Properties of the Halogens 1146
 - Compounds of the Halogens 1148 Uses of the Halogens 1150

26 METALLURGY AND THE CHEMISTRY OF METALS (ONLINE ONLY)

26.1 Occurrence of Metals 1159

26.2 Metallurgical Processes 1160

- Preparation of the Ore 1160 Production of Metals 1160
 The Metallurgy of Iron 1161 Steelmaking 1162 Purification
- of Metals 1164 Thinking Outside the Box: Copper 1166
- 26.3 Band Theory of Conductivity 1166 • Conductors 1166 • Semiconductors 1167
- 26.4 Periodic Trends in Metallic Properties 1169
- 26.5 The Alkali Metals 1169
- 26.6 The Alkaline Earth Metals 1172• Magnesium 1172 Calcium 1173
- 26.7 Aluminum 1174
- Appendix 1 Mathematical Operations A-1
- Appendix 2 Thermodynamic Data at 1 atm and 25°C A-6
- Appendix 3 Solubility Product Constants at 25°C A-13
- Appendix 4 Dissociation Constants for Weak Acids and Bases at 25°C A-15
- Glossary G-1
- Answers to Odd-Numbered Problems AP-1
- Index I-1



Elenathewise/iStockphoto/Getty Images

List of Applications

Thinking Outside the Box

Tips for Success in Chemistry Class 5 How Many States of Matter Are There? 25 Measuring Atomic Mass 51 Everyday Occurrences of the Photoelectric Effect 76 How Do We Know that ns Orbitals Fill Before (*n* – 1) *d* Orbitals? 122 Mistaking Strontium for Calcium 148 Functional Groups 183 Species with Unpaired Electrons 232 Phases 285 Molecular Orbitals in Heteronuclear Diatomic Species 289 Visible Spectrophotometry 388 Heat Capacity of Calorimeters 438 Decompression Injury 505 X-Ray Diffraction 546 Intravenous Fluids 599 Fluoride Poisoning 600 Surface Area 655 Thermodynamics and Living Systems 703 Biological Equilibria 765 Substituent Electronegativity and Acid Strength 795 Equilibrium and Tooth Decay 884 Molten Carbonate Fuel Cells 934 Nuclear Medicine 977 Chelation Therapy 1023 Thalidomide Analogues 1065 Electrically Conducting Polymers 1104 Arsenic 1145 Copper 1166

Environmental Aspects

Global Climate Change 12 Isolated Atoms and Stratospheric Ozone Depletion 54 Solar Energy 72 Lead in the Water in Flint, Michigan 147 CFC Nomenclature 183 Tropospheric Ozone 233 Polarity and the Greenhouse Effect 259 Green Chemistry Metrics 331 Softening Hard Water 364 Energy Density and Specific Energy 439 Lead in Gasoline 497 Water Vapor in the Atmosphere 560 EPS Foam Bans 606 Catalytic Converters 668 Thermodynamic Limits on Energy Conversion 703 Bioaccumulation of Toxins 764 Acid Rain 812 Ocean Acidification 860 Artificial Leaves as a Renewable Energy Source 936 Nuclear Power, Waste, and Recycling 974

Key Skills

Dimensional Analysis 36 Interconversion Among Mass, Moles, and Numbers of Atoms 64 The Importance of Orbital Size 112 Determining Ground-State Valence Electron Configurations Using the Periodic Table 160 Ionic Compounds: Nomenclature and Molar Mass Determination 208 Drawing Lewis Structures 244 Molecular Shape and Polarity 306 Limiting Reactant 350 Net Ionic Equations 414 Enthalpy of Reaction 470 Mole Fractions 530 Intermolecular Forces 574 Entropy as a Driving Force 620 First-Order Kinetics 684 Determining ΔG° 722 Equilibrium Problems 786 Salt Hydrolysis 848 Buffers 904 Electrolysis of Metals 952

Profiles in Chemistry

Linus Pauling 273 Marie-Anne Paulze Lavoisier 319 Fritz Haber 514 Alice Ball 582 St. Elmo Brady 795

Preface

The fifth edition of *Chemistry: Atoms First* by Burdge and Overby builds further on the success of the first four editions. Changes to this edition focus on new additions to the pedagogy, refinement of the current approach, and other innovations driven by feedback from instructors and students alike.

NEW! Environmental Aspects

Given the current climate of environmental awareness in both the classroom and the public in general, we have added a new series of vignettes in the form of boxed features titled Environmental Aspects. Each of the first twenty chapters of the text contains one of these boxes, which provides instructors an opportunity to include timely, environmentally focused material within the context of each chapter. To encourage student engagement with the Environmental Aspects material, many of the boxes have accompanying end-of-chapter problems associated with them. These problems are designated with the Environmental Aspects icon.

Environmental Aspects

Global Climate Change

Those who describe themselves as "skeptical" about climate change sometimes posit that global temperature change is normal, and that any observed increase in temperature is simply the result of natural processes— outside the control of humans. However, there is an enormous body of climate research that clearly demonstrates otherwise. One line of inquiry that has helped to established the connection between human activity and so-called "global warming" involves what is known as *vertical structure of temperature*.

Earth's atmosphere is divided into a series of altitudinal layers: the troposphere (ground-level to 8-14.5 km), the stratosphere (top of the troposphere–50 km), the mesosphere (50–80 km), the thermosphere (80–700 km), and the exosphere (700–10,000 km). The troposphere is where we live, where weather events occur, and where nearly all human activity takes place. When we burn fossil fuels, we increase the amount of CO₂ in the *troposphere*.

In 1988, atmospheric scientist V. Ramanathan, now of the Scripps Institution of Oceanography at the University of California, San Diego, proposed that global temperature change caused by the anthropogenic increase in atmospheric CO₂ could be readily distinguished from that caused by *natural* events, such as increased solar activity. Global temperature increase caused by the Sun, he reasoned, would occur in both the troposphere *and* the stratosphere. Conversely, changes caused by the enhanced greenhouse effect (the result of increased atmospheric CO₂ concentration) would cause warming of the troposphere; but *cooling* of the stratosphere—because more of the heat radiating from Earth's surface would be trapped by greenhouse gases in the troposphere, thus never reaching the stratosphere. Indeed, temperature monitoring over several decades has demonstrated an *increase* in tropospheric temperature, and a *decrease* in stratospheric temperature. This is one of the observations that climate scientists refer to as a *human fingerprint* on global climate change.

Reflecting the Diverse World Around

McGraw Hill believes in unlocking the potential of every learner at every stage of life. To accomplish that, we are dedicated to creating products that reflect, and are accessible to, all the diverse, global customers we serve. Within McGraw Hill, we

foster a culture of belonging, and we work with partners who share our commitment to equity, inclusion, and diversity in all forms. In McGraw Hill Higher Education, this includes, but is not limited to, the following:

- Refreshing and implementing inclusive content guidelines around topics including generalizations and stereotypes, gender, abilities/disabilities, race/ethnicity, sexual orientation, diversity of names, and age.
- Enhancing best practices in assessment creation to eliminate cultural, cognitive, and affective bias.
- Maintaining and continually updating a robust photo library of diverse images that reflect our student populations.
- Including more diverse voices in the development and review of our content.
- Strengthening art guidelines to improve accessibility by ensuring meaningful text and images are distinguishable and perceivable by users with limited color vision and moderately low vision.

NEW! Profiles in Chemistry

Many important discoveries and contributions have been made by people whose names may not be well known in the history of chemistry. Our Profiles in Chemistry boxes feature the names and accomplishments of some of those people.

Profiles in Chemistry

St. Elmo Brady

St. Elmo Brady (1916–1966) was the first African American to earn a Ph.D. in chemistry in the United States. He attended Fisk University in Nashville, Tennessee, and then did his graduate work at the University of Illinois Urbana-Champaign. There he did exhaustive studies on the impact that various substituents have on the strength of carboxylic acids. He went on to teach at the Tuskegee Normal and Industrial Institute, now Tuskegee University. He later became chair of the chemistry department at Howard University in Washington, DC. Howard University, founded in 1867, is the oldest HBCU (historically Black colleges and universities) in the United States. Later, Brady would return to lead the chemistry department at Fisk University, his undergraduate alma mater. There he designed the nation's first HBCU graduate program in chemistry. Eventually, he would go on to develop such



programs at several other HBCUs. In 2019, Brady was honored posthumously with a National Historic Chemical Landmark by the American Chemical Society.

Updated Pedagogy

To refresh student self-assessments, we have updated all Section Review questions to reimagined or completely new questions. Students report benefiting from these self-evaluation questions as they assess their level of mastery of the material in one section before proceeding to the next. They also report using them to review for quizzes and exams. In addition, there is a significant number of new or revised endof-chapter problems.

All numerical values in tables and appendices have been updated to match the values that students will encounter in ALEKS. Figures and tables have been refined for accessibility and ADA compliance.



New and Updated Chapter Content

Chapter 1—A new Student Annotation has been added to note the recent redefinition of the meter and of the kilogram.

Chapter 2—A new chapter-opening photo and caption detail the first observation of the radioactive decay of bismuth-209. We have also updated the value of Avogadro's number.

Chapter 5—A new chapter-opening photo and caption describe non-stoichiometric compounds.

Chapter 6—Electronegativity values have been updated to reflect the most recent determinations.

Chapter 7—New Profiles in Chemistry box features the work of Linus Pauling. Chapter 8—A new Profiles in Chemistry box features the work of Marie-Anne Paulze Lavoisier.

Chapter 11—A new Profiles in Chemistry box features the work of Fritz Haber. **Chapter 13**—A new Profiles in Chemistry box features the work of Alice Ball.

Chapter 14—We have refined the introduction of catalysis and have updated the reaction pathway figure to more accurately reflect the function of a catalyst.



Figure 14.7 From left to right: The decrease in bromine concentration as time elapses is indicated by the loss of color.

Ken Karp/McGraw Hill

Chapter 15—A new chapter-opening photo and caption allude to the statistical treatment of entropy and a new conceptual end-of-chapter problem queries students' understanding of the description of entropy as "time's arrow."

Chapter 17—The description of Brønsted acid-base reactions has been enhanced by the addition of curved arrows to illustrate the movement of electrons. A new Profiles in Chemistry box features the work of St. Elmo Brady.

Chapter 19—A new Thinking Outside the Box features the use of molten carbonate fuel cells for carbon capture.

ARTWORK

A popular feature of our book is the use of thought-provoking chapter-opening photos that may not be immediately obvious in their connection to the content of chapter. The accompanying captions elucidate the important connections. We continue this with new photos and updated captions in many of the chapters.

The Construction of a Learning System

Writing a textbook and its supporting learning tools is a multifaceted process. McGraw Hill's 360° Development Process is an ongoing, market-oriented approach to building accurate and innovative learning systems. It is dedicated to continual large scale and incremental improvement, driven by multiple customer feedback loops and checkpoints.



Figure 3.16 (a) Clockwise and (b) counterclockwise spins of an electron. The magnetic fields generated by these two spinning motions are analogous to those from the two magnets. The upward and downward arrows are used to denote the direction of spin. This is initiated during the early planning stages of new products and intensifies during the development and production stages. The 360° Development Process then begins again upon publication, in anticipation of the next version of each print and digital product. This process is designed to provide a broad, comprehensive spectrum of feedback for refinement and innovation of learning tools for both student and instructor. The 360° Development Process includes market research, content reviews, faculty and student focus groups, course- and product-specific symposia, accuracy checks, and art reviews, all guided by carefully selected Content Advisors.

The Learning System Used in Chemistry: Atoms First

Building Problem-Solving Skills. The entirety of the text emphasizes the importance of problem solving as a crucial element in the study of chemistry. Beginning with Chapter 1, a basic guide fosters a consistent approach to solving problems throughout the text. Each **Worked Example** is divided into four consistently applied steps: *Strategy* lays the basic framework for the problem; *Setup* gathers the necessary information for solving the problem; *Solution* takes us through the steps and calculations; *Think About It* makes us consider the feasibility of the answer or information illustrating the relevance of the problem.

	One type of laser used in the treatment of vascular skin lesions is a neodymium-doped yttrium aluminum garnet, or Nd:YAG, laser. The wavelength commonly used in these treatments is 532 nm. What is the frequency of this radiation?				
	Strategy We must convert the wavelength to meters and solve for frequency using Equation 3.3 ($c = \lambda \nu$).				
Setup Rearranging Equation 3.3 to solve for frequency gives $\nu = \frac{c}{\lambda}$. The speed of light, c, is 3.00×10^8 m/s. λ (in meters) = $532 \text{ nm} \times \frac{1 \times 10^9 \text{ m}}{1 \text{ nm}} = 5.32 \times 10^{-7} \text{ m}.$					
	Solution				
	$\nu = \frac{3.00 \times 10^8 \text{ m/s}}{5.32 \times 10^{-7} \text{ m}} = 5.64 \times 10^{14} \text{ s}^{-1}$				
	Think About It				
	Make sure your units cancel properly. A common error in this type of problem is neglecting to convert wavelength to meters.				
	Practice Problem ATTEMPT What is the wavelength (in meters) of an electromagnetic wave whose frequency is 1.61 × 10 ¹² s ⁻¹ ?				
	Practice Problem BUILD What is the frequency (in reciprocal seconds) of electromagnetic radiation with a wavelength of 1.03 cm?				
	Practice Problem CONCEPTUALIZE Which of the following sets of waves best represents the relative wavelengths/				

After working through this problem-solving approach in the Worked Examples, there are three Practice Problems for students to solve. *Practice Problem A* (Attempt) is always very similar to the Worked Example and can be solved using the same strategy and approach.

Although *Practice Problem B* (Build) probes comprehension of the same concept as Practice Problem A, it generally is sufficiently different in that it cannot be solved using the exact approach used in the Worked Example. Practice Problem B takes problem solving to another level by requiring students to develop a strategy independently. *Practice Problem C* (Conceptualize) provides an exercise that further probes the student's conceptual understanding of the material and many employ concept and molecular art. The regular use of the Worked Example and Practice Problems in this text will help students develop a robust and versatile set of problem-solving skills.

Section Review. Every section of the book that contains Worked Examples and Practice Problems ends with a Section Review. The Section Review enables the student to evaluate whether they understand the concepts presented in the section. **Key Skills.** Newly located immediately before end-of-chapter problems, Key Skills are easy to find review modules where students can return to refresh and hone specific skills that the authors know are vital to success in later chapters. The answers to the Key Skills can be found in the Answer Appendix in the back of the book.



Student Hot Spots. In the text, we have identified areas of particularly difficult content as "Student Hot Spots"—and use them to direct students to a variety of learning resources specific to that content. Students will be able to access over 1,000 digital learning resources throughout this text's eBook. These learning resources present summaries of concepts and worked examples, including over 200 videos of chemistry faculty solving problems or modeling concepts which students can view over and over again.

Applications. Each chapter offers a variety of tools designed to help facilitate learning. *Student Annotations* provide helpful hints and simple suggestions to the student.

The nomenclature of molecular compounds follows in a similar manner to that of ionic compounds. Most molecular compounds are composed of two nonmetals (see [If Section 2.6, Figure 2.10]). To name such a compound, we first name the element that appears first in the formula. For HCl that would be hydrogen. We then name the second element, changing the ending of its name to -ide. For HCl, the second element is chlorine, so we would change chlorine to chloride. Thus, the systematic name of HCl is *hydrogen chloride*. Similarly, HI is hydrogen iodide (iod*ine* \longrightarrow iod*ide*) and SiC is silicon carbide (carbon \longrightarrow carbide).

Student Hot Spot

Student data indicate you may struggle with VSEPR. Access your eBook to view additional Learning Resources on this topic.

Student Annotation: Recall that compounds composed of two elements are called *binary* compounds.

Thinking Outside the Box is an application providing a more in-depth look into a specific topic. *Learning Outcomes* provide a brief overview of the concepts the student should understand after reading the chapter. It's an opportunity to review areas that the student does not feel confident about upon reflection.

Iany organic compounds are derivatives of alkar atoms has been replaced by a group of atoms i orop. The functional group determines many of f a compound because it typically is where a che able 5.9 lists the names and provides ball-and-si- nportant functional groups. Ethanol, for example, the alcohol in alcoholi 2,H ₀ with one of the hydrogen atoms replaced b roup. Its name is derived from that of <i>ethane</i> , in wo carbon atoms. TABLE 5.9 Organic Function	tes in which one of the rnown as a <i>functional</i> the chemical properties smical reaction occurs. tick models of several c beverages, is ethane by an alcohol ($-OH$) dicating that it contains al Groups	Ethanol Ethanol can also be written C ₂ H ₆ O, but e information about the structure of the molecule. Ind several functional groups are discussed in greater
Name	Functional Group	Model
Alcohol	—ОН	_
Aldehyde	—СНО	
Carboxylic acid	—СООН	

Visualization. This text seeks to enhance student understanding through a variety of both unique and conventional visual techniques. A truly unique element in this text is the inclusion of a distinctive feature entitled **Visualizing Chemistry.** These two-page spreads appear as needed to emphasize fundamental, vitally important principles of chemistry. Setting them apart visually makes them easier to find and revisit as needed throughout the course term. Each Visualizing Chemistry feature concludes with a "What's the Point?" box that emphasizes the correct take-away message.

There is a series of conceptual end-of-chapter problems for each Visualizing Chemistry piece. The answers to the Visualizing Chemistry problems, Key Skills problems, and all odd-numbered end of chapter Problems can be found in the Answer Appendix at the end of the text.

Flow Charts and a variety of inter-textual materials such as *Rewind* and *Fast Forward Buttons* and *Section Review* are meant to enhance student understanding and comprehension by reinforcing current concepts and connecting new concepts to those covered in other parts of the text.



Media. Many Visualizing Chemistry pieces have been made into captivating and pedagogically-effective *animations* for additional reinforcement of subject matter first encountered in the textbook. Each Visualizing Chemistry animation is noted by an icon.

Integration of Electronic Homework. You will find the *electronic homework* integrated into the text in numerous places. A large number of the end-of-chapter problems are in the electronic homework system ready to assign to students.

For us, this text will always remain a work in progress. We encourage you to contact us with any comments or questions.

Julia Burdge juliaburdge@gmail.com

Jason Overby overbyj@cofc.edu







Break down barriers and build student knowledge

Students start your course with varying levels of preparedness. Some will get it quickly. Some won't. ALEKS is a course assistant that helps you meet each student where they are and provide the necessary building blocks to get them where they need to go. You determine the assignments and the content, and ALEKS will deliver customized practice until they truly get it.

Experience The ALEKS Difference



Easily Identify Knowledge Gaps



- ALEKS's "Initial Knowledge Check" helps accurately evaluate student levels and gaps on day one, so you know precisely where students are at and where they need to go when they start your course.
- You know when students are at risk of falling behind through ALEKS Insights so, you can remediate be it through prep modules, practice questions, or written explanations of video tutorials.
- Students always know where they are, how they are doing, and can track their own progress easily.



Gain More Flexibility and Engagement

Teach your course your way, with best-in-class content and tools to immerse students and keep them on track.

- **ALEKS gives you flexibility** to assign homework, share a vast library of curated content including videos, review progress and provide student support, anytime anywhere.
- You save time otherwise spent performing tedious tasks while having more control over and impact on your students' learning process.
- **Students gain a deeper level of understanding** through interactive and hands-on assignments that go beyond multiple-choice questions.

with ALEKS[®] Constructive Learning Paths.



Narrow the Equity Gap

Efficiently and effectively create individual pathways for students without leaving anyone behind.

- ALEKS creates an equitable experience for all students, making sure they get the support they need to successfully finish the courses they start.
- You help reduce attrition, falling enrollment, and further widening of the learning gap.
- Student success rates improve not just better grades, but better learning.



Count on Hands-on Support

A dedicated Implementation Manager will work with you to build your course exactly the way you want it and your students need it.

- An ALEKS Implementation Manager is with you every step of the way through the life of your course.
- You never have to figure it out on your own or be your student's customer service. We believe in a consultative approach and take care of all of that for you, so you can focus on your class.
- Your students benefit from more meaningful in moments with you, while ALEKS—directed by you—does the rest.



Already benefitting from ALEKS?

Check out our New Enhancements: mheducation.com/highered/aleks/new-releases.html

Instructor and Student Resources



ALEKS (Assessment and LEarning in Knowledge Spaces) is a web-based system for individualized assessment and learning available 24/7 over the Internet. ALEKS uses artificial intelligence to accurately determine a students' knowledge and then guides them to the material that they are most ready to learn. ALEKS offers immediate feedback and access to ALEKSPedia—an interactive text that contains concise entries on chemistry topics. ALEKS is also a full-featured course management system with rich reporting features that allow instructors to monitor individual and class performance, set student goals, assign/grade online quizzes, and more. ALEKS allows instructors to spend more time on concepts while ALEKS teaches students practical problem-solving skills. And with ALEKS 360, your student also has access to this text's eBook. Learn more at www.aleks.com/highered/science

McGraw Hill Virtual Labs is a must-see, outcomes-based lab simulation. It assesses a student's knowledge and adaptively corrects deficiencies, allowing the student to learn faster and retain more knowledge with greater success. First, a student's knowledge is adaptively leveled on core learning outcomes: Questioning reveals knowledge deficiencies that are corrected by the delivery of content that is conditional on a student's response. Then, a simulated lab experience requires the student to think and act like a scientist: recording, interpreting, and analyzing data using simulated equipment found in labs and clinics. The student is allowed to make mistakes—a powerful part of the learning experience! A virtual coach provides subtle hints when needed, asks questions about the student's choices, and allows the student to reflect on and correct those mistakes. Whether your need is to overcome the logistical challenges of a traditional lab, provide better lab prep, improve student performance, or make your online experience one that rivals the real world, McGraw Hill Virtual Labs accomplishes it all.



Instructors have access to the following instructor resources:

- Art Full-color digital files of all illustrations, photos, and tables in the book can be readily incorporated into lecture presentations, exams, or custom-made class-room materials. In addition, all files have been inserted into PowerPoint slides for ease of lecture preparation.
- Animations Numerous full-color animations illustrating important processes are also provided. Harness the visual impact of concepts in motion by importing these files into classroom presentations or online course materials.

xxxii

- Accessible PowerPoint Lecture Outlines Ready-made presentations that combine art and lecture notes are provided for each chapter of the text.
- **Instructor's Solutions Manual** This supplement contains complete, worked-out solutions for the Practice Problem C questions, Key Skills questions, and *all* the end-of-chapter problems in the text.
- **Computerized Test Bank** Also among the instructor resources is a computerized test bank. The testing software TestGen[®] can be used to create customized exams quickly. Hundreds of text-specific, open-ended, and multiple-choice questions are included in the question bank.

Create

Your Book, Your Way McGraw Hill's Content Collections Powered by Create[®] is a self-service website that enables instructors to create custom course materials—print and eBooks—by drawing upon McGraw Hill's comprehensive, cross-disciplinary content. Choose what you want from our high-quality textbooks, articles, and cases. Combine it with your own content quickly and easily, and tap into other rights-secured, third-party content such as readings, cases, and articles. Content can be arranged in a way that makes the most sense for your course and you can include the course name and information as well. Choose the best format for your course: color print, black-and-white print, or eBook. The eBook can be included in your Connect course and is available on the free ReadAnywhere app for smartphone or tablet access as well. When you are finished customizing, you will receive a free digital copy to review in just minutes! Visit McGraw Hill Create[®]—www.mcgrawhillcreate.com—today and begin building!

Student Solutions Manual

Students will find answers to the Visualizing Chemistry and Key Skills questions and detailed solutions and explanations for the odd-numbered problems from the text in the solutions manual.

Laboratory Manual

Laboratory Manual to Accompany Chemistry: Atoms First by Gregg Dieckmann and John Sibert from the University of Texas at Dallas. This laboratory manual presents a lab curriculum that is organized around an atoms-first approach to general chemistry. The philosophy behind this manual is to (1) provide engaging experiments that tap into student curiosity, (2) emphasize topics that students find challenging in the general chemistry lecture course, and (3) create a laboratory environment that encourages students to "solve puzzles" or "play" with course content and not just "follow recipes." The laboratory manual represents a terrific opportunity to get students turned on to science while creating an environment that connects the relevance of the experiments to a greater understanding of their world. This manual has been written to provide instructors with tools that engage students, while providing important connections to the material covered in an atoms-first lecture course.

Important features of this laboratory manual:

- Early experiments focus on topics introduced early in an atoms-first course properties of light and the use of light to study nanomaterials, line spectra and the structure of atoms, periodic trends, etc.
- Prelab or *foundation* exercises encourage students to understand the important concepts/calculations/procedures in the experiment through working together.
- Postlab or *reflection* exercises put the lab content in the context of a larger chemistry/science picture.
- Instructor's resources (found in the Instructor Resources on Connect[®]) provided with each experiment outline variations that can be incorporated to enrich the student experience or tailor the lab to the resources/equipment available at the institution.

Acknowledgments

We wish to thank the many people—past and present—who have contributed to the development of this text but especially the reviewers of the fourth edition text for this new fifth edition text:

William Alexander, University of Memphis	Joseph M. Lanzafame, Rochester Institute of Technology
Ramesh Arasasingham, University of California, Irvine	Jason McAfee, University of North Texas
Samar Ayesh, Harold Washington College	Tom McGrath, Baylor University
Anne Distler, Cuyahoga Community College	Kevin McGregor, United States Air Force Academy Prepa-
Steven B. Dulaney, Trine University	ratory School
Shiv Halasyamani, University of Houston	Dr. Vicki Moravec, Trine University
Adam Hoffman, University of Dubuque	Andrew Petit, California State University, Fullerton
Jeffry Kelber, University of North Texas	Prasad Polavarapu, Vanderbilt University
Rizalia M. Klausmeyer, Baylor University	Shiv K. Sharma, University of Miami
Jeremy Kua, University of San Diego	Ashley Corrigan Steffey, University of San Diego

Raymond Chang's contributions have been invaluable. His unfaltering diligence and legendary attention to detail have added immeasurably to the quality of this book.

We both thank and acknowledge our families for their continued and devoted support.

Finally, we must acknowledge our McGraw Hill family for their inspiration, excitement, and support of this project: Managing Director Kathleen McMahon; Executive Portfolio Manager Ian Townsend; Senior Product Developer Mary Hurley, Senior Content Project Manager Laura Bies; Senior Director of Digital Content Shirley Hino, and Senior Marketing Manager Cassie Cloutier.



Chemistry

ATOMS FIRST

FIFTH EDITION

Julia Burdge COLLEGE OF WESTERN IDAHO

> Jason Overby COLLEGE OF CHARLESTON



Chapter

Chemistry: The Science of Change

1.1 The Study of Chemistry

- Chemistry You May Already Know
- The Scientific Method

1.2 Scientific Measurement

SI Base Units • Mass • Temperature
Derived Units: Volume and Density

1.3 Uncertainty in Measurement

- Significant Figures
- Calculations with Measured Numbers
- Accuracy and Precision

1.4 Using Units and Solving Problems

- Conversion Factors
- Dimensional Analysis—Tracking Units

1.5 Classification of Matter • States of Matter • Mixtures

1.6

The Properties of Matter

• Physical Properties • Chemical Properties • Extensive and Intensive Properties



AT ONLY a billionth of a degree above absolute zero, ultra-cold atomic gases are used to study the behaviors exhibited by simple quantum particles when they interact with one another. These quantum mechanical interactions ultimately give rise to observable phenomena such as high-temperature superconductivity and quantum magnetism. Research on atomic behavior under these conditions may facilitate new discoveries in fields ranging from materials science to quantum computing.

Before You Begin, Review These Skills

- Basic algebra
- Scientific notation [▶] Appendix 1]

1.1 THE STUDY OF CHEMISTRY

Chemistry often is called the *central science* because knowledge of the principles of chemistry can facilitate understanding of other sciences, including physics, biology, geology, astronomy, oceanography, engineering, and medicine. *Chemistry* is the study of *matter* and the *changes* that matter undergoes. Matter is what makes up our bodies, our belongings, our physical environment, and in fact our entire universe. *Matter* is anything that has mass and occupies space.

Chemistry You May Already Know

You may already be familiar with some of the terms used in chemistry. Even if this is your first chemistry course, you may have heard of *molecules* and know them to be tiny pieces of a substance—much too tiny to see. Further, you may know that molecules are made up of *atoms*, even smaller pieces of matter. And even if you don't know what a *chemical formula* is, you probably know that H_2O is water. You may have used, or at least heard, the term *chemical reaction;* and you are undoubtedly familiar with a variety of common processes that are chemical reactions, such as those shown in Figure 1.1. Don't worry if you are not familiar with these terms; they are defined in the early chapters of this book.

The processes illustrated in Figure 1.1 are all things that you can observe at the *macroscopic level*. In other words, these processes and their results are visible to the human eye. In studying chemistry, you will learn to visualize and understand these same processes at the *submicroscopic* or *molecular level*.

The Scientific Method

Advances in our understanding of chemistry (and other sciences) are the result of scientific experiments. Although scientists do not all take the same approach to experimentation, they must follow a set of guidelines known as the scientific method to have their results added to the larger body of knowledge within a given field. The flowchart in Figure 1.2 illustrates this basic process. The method begins with the gathering of data via observations and experiments. Scientists study these data and try to identify patterns or trends. When they find a pattern or trend, they may summarize their findings with a law, a concise verbal or mathematical statement of a reliable relationship between phenomena. Scientists may then formulate a *hypothesis*, a tentative explanation for their observations. Further experiments are designed to test the hypothesis. If experiments indicate that the hypothesis is incorrect, the scientists go back to the drawing board, try to come up with a different interpretation of their data, and formulate a new hypothesis. The new hypothesis will then be tested by experiment. When a hypothesis stands the test of extensive experimentation, it may evolve into a theory. A theory is a unifying principle that explains a body of experimental observations and the laws that are based on them. Theories can also be used to predict related phenomena, so theories are constantly being tested. If a theory is disproved by experiment, then it must be discarded or modified so that it becomes consistent with experimental observations.

Student Annotation: Macroscopic means large enough to be seen with the unaided eye

Student Annotation: Submicroscopic means too small to be seen, even with a microscope. Atoms and molecules are submicroscopic.



(a)



(c)



(b)



Figure 1.1 Many familiar processes are chemical reactions: (a) The flame of a gas stove is the combustion of natural gas, which is primarily methane. (b) The bubbles produced when Alka-Seltzer dissolves in water are carbon dioxide, produced by a chemical reaction between two ingredients in the tablets. (c) The formation of rust is a chemical reaction that occurs when iron, water, and oxygen are all present. (d) Many baked goods "rise" as the result of a chemical reaction that produces carbon dioxide.

(a): Caspar Benson/fStop/Getty Images; (b): Colin Anderson Productions pty Itd/Stockbyte/Brand X Pictures/Getty Images; (c): Anthony Grote/Getty Images; (d): Sharon Dominick/iStock Exclusive/Getty Images



Figure 1.2 Flowchart of the scientific method.

A fascinating example of the use of the scientific method is the story of how smallpox was eradicated. Late in the eighteenth century, an English doctor named Edward Jenner observed that even during outbreaks of smallpox in Europe, milkmaids seldom contracted the disease. He reasoned that when people who had frequent contact with cows contracted *cowpox*, a similar but far less harmful disease, they developed a natural immunity to smallpox. He predicted that intentional exposure to the cowpox virus would produce the same immunity. In 1796, Jenner exposed an 8-year-old boy to the cowpox virus using pus from the cowpox lesions of an infected milkmaid. Six weeks later, he exposed the boy to the *smallpox* virus and, as Jenner had predicted, the boy did *not* contract the disease. Subsequent experiments using the same technique (later dubbed *vaccination* from the Latin *vacca* meaning *cow*) confirmed that immunity to smallpox could be induced.

A superbly coordinated international effort on the part of healthcare workers was successful in eliminating smallpox worldwide. In 1980, the World Health Organization declared smallpox officially eradicated. This historic triumph over a dreadful disease, one of the greatest medical advances of the twentieth century, began with Jenner's astute observations, inductive reasoning, and careful experimentation—the essential elements of the *scientific method*.



Until recently, almost everyone had a smallpox vaccine scar—usually on the upper arm. Chris Livingston/Getty Images

enne zinngeten, eetty innagee

Student Annotation: The last naturally occurring case was in 1977 in Somalia.

Thinking Outside the Box

Tips for Success in Chemistry Class

Success in a chemistry class depends largely on problem-solving ability. The Worked Examples throughout the text are designed to help you develop problem-solving skills. Each is divided into four steps: Strategy, Setup, Solution, and Think About It.

Strategy: Read the problem carefully and determine what is being asked and what information is provided. The Strategy step is where you should think about what skills are required and lay out a plan for solving the problem. Give some thought to what you expect the result to be. If you are asked to determine the number of atoms in a sample of matter, for example, you should expect the answer to be a whole number. Determine what, if any, units should be associated with the result. When possible, make a ballpark estimate of the magnitude of the correct result, and make a note of your estimate.

Setup: Next, gather the information necessary to solve the problem. Some of the information will have been given in the problem itself. Other information, such as equations, constants, and tabulated data (including atomic masses) should also be brought together in this step. Write down and label clearly all of the information you will use to solve the problem. Be sure to write appropriate units with each piece of information. Solution: Using the necessary equations, constants, and other information, calculate the answer to the problem. Pay particular attention to the units associated with each number, tracking and canceling units carefully throughout the calculation. In the event that multiple calculations are required, label any intermediate results. If rounding is necessary, do it only after the last step in the calculation.

Think About It: Consider your calculated result and ask yourself whether or not it makes sense. Compare the units and the magnitude of your result with your ballpark estimate from the Strategy step. If your result does not have the appropriate units, or if its magnitude or sign is not reasonable, check your solution for possible errors. A very important part of problem solving is being able to judge whether the answer is reasonable. It is relatively easy to spot a wrong sign or incorrect units, but you should also develop a sense of *magnitude* and be able to tell when an answer is either way too big or way too small. For example, if a problem asks you to determine the length of a sheet of paper in millimeters and you calculate a number that is less than 1, you should know that it cannot be correct.

Each Worked Example is followed by three Practice Problems: A, B, and C. Practice Problem A, "Attempt," typically is a problem very similar to the Worked Example that can be solved using the same strategy. Practice Problems B and C, "Build" and "Conceptualize" generally test the same skills, but require approaches slightly different from the one used to solve the preceding Worked Example and Practice Problems.

Finally, each section that contains one or more Worked Examples concludes with a Section Review. These consist of multiple-choice problems that enable you to assess your comprehension of the material in the section before moving on to the next section. Answers to Practice Problems A and B and to Section Review Questions can be found at the end of each chapter.

Regular use of the Worked Examples, Practice Problems, and Section Reviews in this text can help you develop an effective set of problemsolving skills. They can also help you assess whether you are ready to move on to the next new concepts. If you struggle with the Practice Problems or Section Reviews, then you probably need to review the corresponding Worked Example and the concepts that led up to it.

5

25mL

Volumetric pipette

(b)

1.2 SCIENTIFIC MEASUREMENT

Scientists use a variety of devices to measure the properties of matter. A meterstick is used to measure length; a burette, pipette, graduated cylinder, and volumetric flask are used to measure volume (Figure 1.3); a balance is used to measure mass; and a thermometer is used to measure temperature. Properties that can be measured are called *quantitative* properties because they are expressed using numbers. When we express a measured quantity with a number, though, we must always include the appropriate unit; otherwise, the measurement is meaningless. For example, to say that the depth of a swimming pool is 3 is insufficient to distinguish between one that is 3 *feet* (0.9 meter) and one that is 3 *meters* (9.8 feet) deep. Units are essential to reporting measurements correctly.

The two systems of units with which you are probably most familiar are the *English system* (foot, gallon, pound, etc.) and the *metric system* (meter, liter, kilogram, etc.). Although there has been an increase in the use of metric units in the United States in recent years, English units still are used commonly. For many years, scientists

Figure 1.3 (a) A burette is used to measure the volume of a liquid that has been added to a container. A reading is taken before and after the liquid is delivered, and the volume delivered is determined by subtracting the first reading from the second. (b) A volumetric pipette is used to deliver a precise amount of liquid. (c) A graduated cylinder is used to measure a volume of liquid. It is less precise than the volumetric flask. (d) A volumetric flask is used to prepare a precise volume of a solution for use in the laboratory.



Graduated cylinder (c)



Volumetric flask (d)

6

recorded measurements in metric units, but in 1960, the General Conference on Weights and Measures, the international authority on units, proposed a revised metric system for universal use by scientists. We use both metric and revised metric (SI) units in this book.

SI Base Units

The revised metric system is called the *International System of Units* (abbreviated SI, from the French *Système Internationale d'Unités*). Table 1.1 lists the seven SI base units. All other units of measurement can be derived from these base units. The *SI unit* for *volume*, for instance, is derived by cubing (raising to the power 3) the SI base unit for *length*. The prefixes listed in Table 1.2 are used to denote decimal fractions and decimal multiples of SI units. The use of these prefixes enables scientists to tailor the magnitude of a unit to a particular application. For example, the meter (m) is appropriate for describing the dimensions of a classroom, but the kilometer (km), 1000 m, is more appropriate for describing the distance between two cities. Units that you will encounter frequently in the study of chemistry include those for mass, temperature, volume, and density.

Student Annotation: According to the U.S. Metric Association (USMA), the United States is "the only significant holdout" with regard to adoption of the metric system. The other countries that continue to use traditional units are Myanmar (formerly Burma) and Liberia. 7

Student Annotation: Although historically defined using physical objects, the SI base units meter and kilogram have been redefined in terms of universal physical constants. The meter is now defined in terms of the speed of light in a vacuum [▶] Section 3.2], and the kilogram is defined in terms of Planck's constant [▶] Section 3.3].

TABLE 1.1	Base S	SI Units	
Base Quant	ity	Name of Unit	Symbol
Length		meter	m
Mass		kilogram	kg
Time		second	S
Electric curr	rent	ampere	А
Temperatu	re	kelvin	K
Amount of sub	stance	mole	mol
Luminous inte	ensity	candela	cd

TABLE 1.2Prefixes Used with SI Units

Prefix	Symbol	Meaning	Example
Tera-	Т	$1 \times 10^{12} (1,000,000,000,000)$	1 teragram (Tg) = 1×10^{12} g
Giga-	G	$1 \times 10^9 $ (1,000,000,000)	1 gigawatt (GW) = 1×10^9 W
Mega-	М	$1 \times 10^{6} (1,000,000)$	1 megahertz (MHz) = 1×10^6 Hz
Kilo-	k	$1 \times 10^3 (1,000)$	1 kilometer (km) = 1×10^3 m
Deci-	d	$1 \times 10^{-1} (0.1)$	1 deciliter (dL) = 1×10^{-1} L
Centi-	с	$1 \times 10^{-2} (0.01)$	1 centimeter (cm) = 1×10^{-2} m
Milli-	m	$1 \times 10^{-3} (0.001)$	1 millimeter (mm) = 1×10^{-3} m
Micro-	μ	$1 \times 10^{-6} \ (0.000001)$	1 microliter (μ L) = 1 × 10 ⁻⁶ L
Nano-	n	$1 \times 10^{-9} (0.000000001)$	1 nanosecond (ns) = 1×10^{-9} s
Pico-	р	$1 \times 10^{-12} \ (0.000000000001)$	1 picogram (pg) = 1×10^{-12} g

Mass

Although the terms *mass* and *weight* often are used interchangeably, they do not mean the same thing. Strictly speaking, weight is the force exerted by an object or sample due to gravity. *Mass* is a measure of the amount of matter in an object or sample. Because gravity varies from location to location (gravity on the Moon is only about one-sixth that on Earth), the weight of an object varies depending on where it is measured. The mass of an object remains the same regardless of where it is measured. The SI base unit of mass is the kilogram (kg), but in chemistry the smaller gram (g) often is more convenient and is more commonly used:

$$1 \text{ kg} = 1000 \text{ g} = 1 \times 10^{3} \text{ g}$$

Occasionally, the most convenient and/or commonly used unit for a particular application is not an SI unit. One such example is the atomic mass unit. The *atomic mass unit (amu)*, as the name suggests, is used to express the masses of atoms—and other objects of similar size. In terms of SI units, the amu is equal to $1.6605378 \times 10^{-24}$ g or $1.6605378 \times 10^{-27}$ kg. Another example is the *angstrom (Å)*, a measure of length that is equal to 1×10^{-10} m.

Temperature

There are two temperature scales used in chemistry: the *Celsius* scale and the *absolute* or *Kelvin* scale. Their units are the *degree Celsius* (°C) and the *kelvin* (K), respectively. The *Celsius* scale [named after Swedish physicist Ander Celsius (1701–1744)] was originally defined using the freezing point (0°C) and the boiling point (100°C) of pure water at sea level. As Table 1.1 shows, the SI base unit of temperature is the *kelvin*. Kelvin is also known as the *absolute* temperature scale because the lowest temperature theoretically possible is 0 K, a temperature referred to as *absolute zero*. No *degree* sign (°) is used to represent a temperature on the Kelvin scale.

Units of the Celsius and Kelvin scales are equal in magnitude, so *a degree Celsius* is equivalent to a *kelvin*. Thus, if the temperature of an object increases by 5° C, it also increases by 5 K. Absolute zero on the Kelvin scale is equivalent to -273.15° C on the Celsius scale. We use the following equation to convert a temperature from units of degrees Celsius to kelvins:

Equation 1.1 $\mathbf{K} = ^{\circ}\mathbf{C} + 273.15$

Worked Example 1.1 illustrates conversions between these two temperature scales.

WORKED EXAMPLE

Normal human body temperature can range over the course of the day from about 36°C in the early morning to about 37°C in the afternoon. Express these two temperatures and the range that they span using the Kelvin scale.

Strategy Use Equation 1.1 to convert temperatures from the Celsius scale to the Kelvin scale. Then convert the range of temperatures from degrees Celsius to kelvins, keeping in mind that 1°C is equivalent to 1 K.

Setup Equation 1.1 is already set up to convert the two temperatures from degrees Celsius to kelvins. No further manipulation of the equation is needed. The range in kelvins will be the same as the range in degrees Celsius.

Solution $36^{\circ}C + 273 = 309 \text{ K}$, $37^{\circ}C + 273 = 310 \text{ K}$, and the range of $1^{\circ}C$ is equal to a range of 1 K.

1.1

Think About It

Check your math and remember that converting a temperature from degrees Celsius to kelvins is different from converting a *difference* in temperature from degrees Celsius to kelvins.

Student Annotation: There is no such thing as a negative temperature on the Kelvin scale.

Student Annotation: The theoretical basis of the Kelvin scale has to do with the behavior of gases. [**WI Chapter 11**]

Student Annotation: Depending on the precision required, the conversion from degrees Celsius to kelvins often is done simply by adding 273, rather than 273.15.

Practice Problem A TTEMPT Express the freezing point of water (0°C), the boiling point of water (100°C), and the range spanned by the two temperatures using the Kelvin scale.
Practice Problem B UILD According to the website of the National Aeronautics and Space Administration (NASA), the average temperature of the universe is 2.7 K. Convert this temperature to degrees Celsius.
Practice Problem CONCEPTUALIZE If a single degree on the Celsius scale is represented by the rectangle on the left, which of the rectangles on the right best represents a single kelvin?

Outside of scientific circles, the Fahrenheit temperature scale is the one most used in the United States. Before the work of Daniel Gabriel Fahrenheit (German physicist, 1686–1736), there were numerous different, arbitrarily defined temperature scales, none of which gave consistent measurements. Accounts of exactly how Fahrenheit devised his temperature scale vary from source to source. In one account, in 1724, Fahrenheit labeled as 0° the lowest artificially attainable temperature at the time (the temperature of a mixture of ice, water, and a substance called *ammonium chloride*). Using a traditional scale consisting of 12 degrees, he labeled the temperature of a healthy human body as the twelfth degree. On this scale, the freezing point of water occurred at the fourth degrees. For better resolution, each degree was further divided into eight smaller degrees. This convention makes the freezing point of water 32°F and normal body temperature 96°F. (Today we consider normal body temperature to be somewhat higher than 96°F.)

The boiling point of water on the Fahrenheit scale is 212°, meaning that there are 180 degrees (212°F minus 32°F) between the freezing and boiling points. This separation is considerably more degrees than the 100 between the freezing point and boiling point of water on the Celsius scale. Thus, the size of a degree on the Fahrenheit scale is only 100/180 or five-ninths of a degree on the Celsius scale. Equation 1.2 gives the relationship between temperatures on the Fahrenheit and Celsius scales.

temperature in °F =
$$\frac{9^{\circ}F}{5^{\circ}C}$$
 × (temperature in °C) + 32°F Equation 1.2

Worked Example 1.2 lets you practice converting from Celsius to Fahrenheit.

WORKED EXAMPLE (1.2

A body temperature above 39°C constitutes a high fever. Convert this temperature to the Fahrenheit scale. **Strategy** We are given a temperature in degrees Celsius and are asked to convert it to degrees Fahrenheit. **Setup** We use Equation 1.2:

temperature in Fahrenheit = $\frac{9^{\circ}F}{5^{\circ}C} \times (\text{temperature in degrees Celsius}) + 32^{\circ}F$

(Continued on next page)