



国外优秀化学教学用书

PEARSON  
Education

# 普通化学 原理与应用

第8版

影印版

# General Chemistry

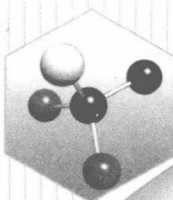
*Principles and Modern Applications*

Eighth Edition

- Ralph H. Petrucci
- William S. Harwood
- F. Geoffrey Herring



高等教育出版社  
Higher Education Press



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# Periodic Table of the Elements

1 1A	2 2A											13 3A	14 4A	15 5A	16 6A	17 7A	18 8A
1 H 1.00794	2 He 4.00260											5 B 10.811	6 C 12.011	7 N 14.0067	8 O 15.9994	9 F 18.9984	10 Ne 20.1797
3 Li 6.941	4 Be 9.01218											13 Al 26.9815	14 Si 28.0855	15 P 30.9738	16 S 32.066	17 Cl 35.4527	18 Ar 39.948
11 Na 22.9898	12 Mg 24.3050											31 Ga 69.723	32 Ge 72.61	33 As 74.9216	34 Se 78.96	35 Br 79.904	36 Kr 83.80
19 K 39.0983	20 Ca 40.078											31 Ga 69.723	32 Ge 72.61	33 As 74.9216	34 Se 78.96	35 Br 79.904	36 Kr 83.80
37 Rb 85.4678	38 Sr 87.62											49 In 114.818	50 Sn 118.710	51 Sb 121.757	52 Te 127.60	53 I 126.904	54 Xe 131.29
55 Cs 132.905	56 Ba 137.327											81 Tl 204.383	82 Pb 207.2	83 Bi 208.980	84 Po (209)	85 At (210)	86 Rn (222)
87 Fr (223)	88 Ra 226.025												114 (287)		116 (289)		118 (293)
*Lanthanide series		58 Ce 140.115	59 Pr 140.908	60 Nd 144.24	61 Pm (145)	62 Sm 150.36	63 Eu 151.965	64 Gd 157.25	65 Tb 158.925	66 Dy 162.50	67 Ho 164.930	68 Er 167.26	69 Tm 168.934	70 Yb 173.04	71 Lu 174.967		
Actinide series		90 Th 232.038	91 Pa 231.036	92 U 238.029	93 Np 237.048	94 Pu (244)	95 Am (243)	96 Cm (247)	97 Bk (247)	98 Cf (251)	99 Es (252)	100 Fm (257)	101 Md (258)	102 No (259)	103 Lr (260)		

Atomic masses are relative to carbon-12. For certain radioactive elements, the numbers listed (in parentheses) are the mass numbers of the most stable isotopes. The scheme for numbering of groups is explained on page 50. The metals are ■ and the nonmetals are ■. Metalloids are indicated by ■. The noble gases are Elements 110, 111, and 112 have not yet been named.

# The Elements

Name	Symbol	Atomic Number	Relative Atomic Weight	Name	Symbol	Atomic Number	Relative Atomic Weight
Actinium	Ac	89	227.028	Mendelevium	Md	101	(258)
Aluminum	Al	13	26.9815	Mercury	Hg	80	200.59
Americium	Am	95	(243)	Molybdenum	Mo	42	95.94
Antimony	Sb	51	121.757	Neodymium	Nd	60	144.24
Argon	Ar	18	39.948	Neon	Ne	10	20.1797
Arsenic	As	33	74.9216	Neptunium	Np	93	237.048
Astatine	At	85	(210)	Nickel	Ni	28	58.693
Barium	Ba	56	137.327	Niobium	Nb	41	92.9064
Berkelium	Bk	97	(247)	Nitrogen	N	7	14.0067
Beryllium	Be	4	9.01218	Nobelium	No	102	(259)
Bismuth	Bi	83	208.980	Osmium	Os	76	190.23
Bohrium	Bh	107	(262)	Oxygen	O	8	15.9994
Boron	B	5	10.811	Palladium	Pd	46	106.42
Bromine	Br	35	79.904	Phosphorus	P	15	30.9738
Cadmium	Cd	48	112.411	Platinum	Pt	78	195.08
Calcium	Ca	20	40.078	Plutonium	Pu	94	(244)
Californium	Cf	98	(251)	Polonium	Po	84	(209)
Carbon	C	6	12.011	Potassium	K	19	39.0983
Cerium	Ce	58	140.115	Praseodymium	Pr	59	140.908
Cesium	Cs	55	132.905	Promethium	Pm	61	(145)
Chlorine	Cl	17	35.4527	Protactinium	Pa	91	231.036
Chromium	Cr	24	51.9961	Radium	Ra	88	226.025
Cobalt	Co	27	58.9332	Radon	Rn	86	(222)
Copper	Cu	29	63.546	Rhenium	Re	75	186.207
Curium	Cm	96	(247)	Rhodium	Rh	45	102.906
Dubnium	Db	105	(262)	Rubidium	Rb	37	85.4678
Dysprosium	Dy	66	162.50	Ruthenium	Ru	44	101.07
Einsteinium	Es	99	(252)	Rutherfordium	Rf	104	(261)
Erbium	Er	68	167.26	Samarium	Sm	62	150.36
Europium	Eu	63	151.965	Scandium	Sc	21	44.9559
Fermium	Fm	100	(257)	Seaborgium	Sg	106	(263)
Fluorine	F	9	18.9984	Selenium	Se	34	78.96
Francium	Fr	87	(223)	Silicon	Si	14	28.0855
Gadolinium	Gd	64	157.25	Silver	Ag	47	107.868
Gallium	Ga	31	69.723	Sodium	Na	11	22.9898
Germanium	Ge	32	72.61	Strontium	Sr	38	87.62
Gold	Au	79	196.967	Sulfur	S	16	32.066
Hafnium	Hf	72	178.49	Tantalum	Ta	73	180.948
Hassium	Hs	108	(265)	Technetium	Tc	43	(98)
Helium	He	2	4.00260	Tellurium	Te	52	127.60
Holmium	Ho	67	164.930	Terbium	Tb	65	158.925
Hydrogen	H	1	1.00794	Thallium	Tl	81	204.383
Indium	In	49	114.818	Thorium	Th	90	232.038
Iodine	I	53	126.904	Thulium	Tm	69	168.934
Iridium	Ir	77	192.22	Tin	Sn	50	118.710
Iron	Fe	26	55.847	Titanium	Ti	22	47.88
Krypton	Kr	36	83.80	Tungsten	W	74	183.84
Lanthanum	La	57	138.906	Uranium	U	92	238.029
Lawrencium	Lr	103	(260)	Vanadium	V	23	50.9415
Lead	Pb	82	207.2	Xenon	Xe	54	131.29
Lithium	Li	3	6.941	Ytterbium	Yb	70	173.04
Lutetium	Lu	71	174.967	Yttrium	Y	39	88.9059
Magnesium	Mg	12	24.3050	Zinc	Zn	30	65.39
Manganese	Mn	25	54.9381	Zirconium	Zr	40	91.224
Meitnerium	Mt	109	(266)				

Atomic masses in this table are relative to carbon-12 and limited to six significant figures, although some atomic masses are known more precisely. For certain radioactive elements the numbers listed (in parentheses) are the mass numbers of the most stable isotopes.

# About the Authors

## Ralph H. Petrucci

Ralph Petrucci received his B.S. in Chemistry from Union College and his Ph.D. from the University of Wisconsin–Madison. Following several years of teaching, research, consulting, and directing the NSF Institutes for Secondary School Science Teachers at Case Western Reserve University, Dr. Petrucci joined the planning staff of the new California State University campus at San Bernardino in 1964. There, in addition to his faculty appointment, he served as Chairman of the Natural Sciences Division and Dean of Academic Planning. Professor Petrucci, now retired from teaching, is the author of several books, including *General Chemistry* with John W. Hill.

## William S. Harwood

Bill Harwood received his B.Sc. from the University of Massachusetts, Amherst and his Ph.D. in Inorganic Chemistry from Purdue University in 1986. He is currently a Professor of Science Education at Indiana University, Bloomington. Previously, Dr. Harwood was at the Department of Chemistry and Biochemistry at the University of Maryland, College Park. In his current role, Dr. Harwood continues to teach chemistry and conduct research in chemical education. He has received several awards for teaching excellence. Dr. Harwood is also active in the American Chemical Society and the Division of Chemical Education and was a consultant to AAAS project 2061. He is involved in the science reform efforts at both the pre-college and college levels. His research focuses on how best to use technology to improve learning in chemistry.

## F. Geoffrey Herring

Geoff Herring received his B.Sc. and his Ph.D. in Physical Chemistry, both from the University of London. He is currently a Professor in the Department of Chemistry of the University of British Columbia, Vancouver. Dr. Herring has research interests in the area of biophysical chemistry and has published over 100 papers in the area of physical chemistry and chemical physics. Recently, Dr. Herring has undertaken studies in the use of information technology and interactive engagement methods in teaching general chemistry with a view to improving student comprehension and learning. Dr. Herring has taught chemistry from undergraduate to graduate levels for 30 years and has been the recipient of the Killam Prize for Excellence in Teaching.

# Preface

**“Know your audience.”** For this new edition, we have tried to follow this important advice to writers by attending more to the needs of those students who are taking a serious journey through the material. We also know that most general chemistry students have career interests not in chemistry, but in biology, medicine, engineering, environmental and agricultural sciences, and so on. And we understand that general chemistry will be the only college chemistry course for some students, and thus their only opportunity to learn some practical applications of chemistry. We have designed this book for all these students.

Students of this text should have already studied some chemistry. But those with no prior background and those who could use a refresher will find that the early chapters develop fundamental concepts from the most elementary ideas. Students who do plan to become professional chemists will also find opportunities in the text to pursue their own special interests.

The typical student may need help identifying and applying principles and visualizing their physical significance. The pedagogical features of this text are designed to provide this help. At the same time, we hope the text serves to sharpen student skills in problem solving and critical thinking. Thus, we have tried to strike the proper balances between principles and applications, qualitative and quantitative discussions, and rigor and simplification.

Throughout the text we provide real-world examples to enhance the discussion. Examples relevant to the biological sciences, engineering, and the environmental sciences will be found in numerous places. This should help to bring the chemistry alive for these students, and help them understand its relevance to their career interests. It also, in most cases, should help them master core concepts.

## Organization

In this edition we retain the core organization of the sixth and seventh editions of this text, but with additional coverage of material, in depth and breadth, in a number of chapters. After a brief overview of core concepts in Chapter 1, we introduce atomic theory, including the periodic table, in Chapter 2. The periodic table is an extraordinarily useful tool, and presenting it early allows us to use the periodic table in new ways throughout the early chapters of the text. In Chapter 3 we introduce chemical compounds and their stoichiometry. Organic compounds are included in this presentation. The early introduction of organic compounds allows us to use organic examples throughout the book. Chapters 4 and 5 introduce chemical reactions. We discuss gases in Chapter 6, partly because they are familiar to students (which helps them build confidence), but also because some instructors prefer to cover this material early to better integrate their lecture and lab programs. Note that Chapter 6 can easily be deferred for coverage with the other states of matter, in Chapter 13. In Chapter 9 we delve more deeply into wave mechanics than in earlier editions, although we do so in a way that allows excision of this material at the instructor's discretion. As with previous editions, we have emphasized real-world chemistry in the final chapters that cover descriptive chemistry (Chapters 22–25), and we have tried to make this material easy to bring forward into earlier parts of the text. Moreover, many topics in these chapters can be covered selectively, without requiring the study of entire chapters. The text ends with heavily revised, comprehensive chapters on organic chemistry (Chapter 27) and biochemistry (Chapter 28).

## Changes to This Edition

We have made a number of smaller organizational changes to improve the flow of information to the student and to reflect contemporary thoughts about how best to teach general chemistry. In Chapter 7 (Thermochemistry), the order in which heat and work are presented has been reversed from that of the 7<sup>th</sup> edition. Also, the concept of standard states is introduced earlier, so that most data in the chapter can be standard-state data. In Chapter 20 (Thermodynamics), the sections on entropy have been reorganized so that all of them precede the introduction to free energy.

► **Increased level of detail and much more problem-solving pedagogy this edition.**

Major changes in this edition have focused on increasing the depth of coverage and adding some more challenging end-of-chapter exercises. Specifically, in Chapter 3 (Chemical Compounds) we have added a section introducing organic compounds, including nomenclature. This allows more reference to organic compounds throughout the book and also suits those who chose to introduce organic chemistry earlier in the course. In Chapter 6 (Gases) there is greater emphasis on the use of SI units and more detail on the kinetic-molecular theory of gases. In Chapter 7 (Thermochemistry) the calculation of quantities of work and the discussion of state functions and path-dependent functions are more extensive than in the previous edition. Chapter 9 (Electrons in Atoms) has been significantly revised to include new sections dealing with wave mechanics and more information on the treatment of wave functions, quantum numbers, orbitals, and radial probability distributions. Chapter 10 (The Periodic Table and Some Atomic Properties) draws more on ideas from Chapter 9 than in earlier editions, permitting a fuller discussion of screening, penetration, and  $Z_{\text{eff}}$ . Chapter 11 (Basic Concepts of Chemical Bonding) has been rearranged to provide a clearer exposition of the general strategy for writing Lewis structures. In Chapter 12 (Additional Aspects of Chemical Bonding), the new ideas developed in Chapter 9 are applied to the hybridization of atomic orbitals and the treatment of molecular orbitals. Molecular orbital theory is extended to cover heteronuclear molecules.

Chapter 13 (Liquids, Solids, and Intermolecular Forces) features an expanded section on crystal structures. In Chapter 15 (Chemical Kinetics) the IUPAC-recommended definition of a general rate of reaction is used in the treatment of reaction rates. Also, reaction mechanisms and enzyme catalysis are presented in more detail.

A new feature in Chapter 17 (Acids and Bases) is a discussion of a general method for equilibrium calculations based on equilibrium constant expressions, material balances, and electroneutrality. In Chapter 20 (Thermodynamics), the concept of entropy is introduced in a new way, and the relationship between  $\Delta G$  and  $\Delta G^\circ$  is developed and explained more fully. The chapters on descriptive inorganic chemistry (Chapters 22–24) have been updated and now include electrode potential (Latimer) diagrams. Chapter 27 (Organic Chemistry) now includes an introductory discussion of  $S_N1$  and  $S_N2$  reactions and other topics of interest to those covering more organic chemistry in this course. Discussions of metabolism and enzyme reactions have been added to Chapter 28 (Chemistry of the Living State).

## In-Text Learning Aids for Students

As with previous editions, we have tried to create the most useful possible text for students. Here are some of the things that should make this so:

**Important Expressions.** The most significant equations, concepts, and rules are highlighted with colored panels so that students can readily find them.

**Summary/Key Terms/Glossary.** Each chapter concludes with a comprehensive verbal *Summary* of important concepts and factual information. The *Summary* is followed by a list of *Key Terms*—terms that appear in boldface type in the text and are defined again in the *Glossary* (Appendix E). Students can use *Key Terms* lists and the *Glossary* to help them master the terminology of general chemistry.



► **Are You Wondering ...?**, probing questions asked by good students, are enhanced in this edition.

► **Keep in Mind** margin notes are new to this edition.

► Detailed applications of chemistry are covered at the end of the chapter in a non-distracting way.

► Many worked examples, carefully developed, step-by-step.

► **Two** practice examples after every in-text example.

► **Integrative Examples**, designed to help students learn how to solve these more complicated problems, are new to this edition.

**Are You Wondering ...?** To help clarify matters that often puzzle students, we pose and then answer questions under this special heading. For obvious pedagogical reasons, these questions are cast in the form in which students typically ask them. Some are designed to help students avoid common misconceptions; others provide analogies or alternative explanations of a concept; still others address apparent inconsistencies in the material they are learning. In response to reviewer and student comments and suggestions, these have been expanded considerably in this edition. Specifically, the *Are You Wondering ...?* format is used in a number of instances to introduce material directed at the better-prepared students. Some of these topics are pursued further in end-of-chapter exercises. These topics can be assigned or omitted at the discretion of the instructor.

**Keep in Mind margin notes.** To help students appreciate the significance of earlier ideas, or to warn them about common pitfalls, we have added to this edition *Keep in Mind* margin notes. As the name suggests, these notes ask students to recall key information about concepts and problem-solving skills. At times, we use these in conjunction with worked examples to forewarn students about common mistakes.

**Focus On boxes.** We believe that relevant applications should be an integral part of the text and that asides should be limited to margin notes and *Are You Wondering ...?* features. With this in mind, we have concluded the text of each chapter with a short essay on a practical topic appropriate to the chapter content. These essays, which may be considered optional reading, focus on ideas introduced in the chapter.

### The Strongest Available Problem-Solving Focus

You probably won't become a better golfer just by watching Tiger Woods play; you have to get onto the course yourself, and often. To give students the support they need to develop strong problem-solving skills, we offer extensive in-text examples that cover all the key concepts introduced in the book, each accompanied by two practice examples. We also provide integrative problems as concluding in-text examples in each chapter and a very large selection of end-of-chapter exercises, including a set that integrate the student media:

**In-Text Illustrative Examples.** In each chapter, most concepts—especially those that students will be expected to apply in homework assignments and examinations—are illustrated with worked-out examples. To aid visual learners and to emphasize abstract concepts, in many cases a line drawing or photograph accompanies an example to help students visualize what is going on in the problem.

**Practice Examples.** These are designed to give students immediate practice in applying the principle(s) illustrated in the example. We offer two for every illustrative example. The first, Practice Example A, provides immediate practice in a problem very similar to the illustrative example. The second, Practice Example B, generally takes the student one step further than the illustrative example. This combination helps students to integrate and extend their knowledge and problem-solving skills. Answers to all Practice Examples are given in Appendix F. Complete solutions are given in the *Selected Solutions Manual*.

**Integrative Examples.** The text includes a special category of problems that requires students to link various important problem types introduced in the chapter—with each other and with problem types from earlier chapters. These problems are meant to be challenging for students, and to help them learn how to solve such problems. Each chapter concludes with a multi-part *Integrative Example*, sometimes of a practical nature. In each case, the problem is broken down into parts, each part is solved, and intermediate results are combined into a final solution and answer.

► Integrative and Advanced Exercises are enhanced this edition.

► Feature Problems, the most challenging in the book, are expanded this edition.

► eMedia Exercises are new to this edition.

► New instructor's supplement.

► New instructor's supplement. Includes prebuilt MS PowerPoint® slides.

► 50% more overhead transparencies this edition.

**End-of-Chapter Exercises.** Each chapter ends with five categories of exercises. *Review Questions* require straightforward application of principles introduced in the chapter, each generally involving a single concept, and either a numerical, symbolic, or short written (or verbal) answer. *Exercises* are grouped by categories related to the text sections, and they are of a broader nature than the *Review Questions*. The *Exercises* are paired, so that there are two problems of the same type. The *Integrative and Advanced Exercises* are not grouped by type. These are generally more difficult than those in the previous sections. They tend to integrate material from multiple sections, or multiple chapters, and they may introduce new ideas or pursue certain ideas further than is done in the text. *Feature Problems* are of special interest. These problems generally require the highest level of cognitive skill on the part of students to solve. Some of these problems retrace aspects of the history of chemistry; a few deal with classic experiments; others require students to interpret data or graphs; some present new material; some suggest alternative techniques for problem solving; and a few summarize main points of the chapter in a comprehensive manner. The *Feature Problems* are a resource that can be used in several ways: as discussion points in class, as assigned homework for individuals, or for collaborative group work. Finally, the *eMedia Exercises*, new to this edition, are questions that can only be solved using the interactive media accompanying this text. This permits the instructor to mandate the use of the media by simply assigning one or more of these problems.

Answers to all red-numbered problems are given in Appendix F. Full solutions to all red-numbered problems are found in the *Selected Solutions Manual*.

## Supplements

### For the Instructor

**Annotated Instructor's Edition (with Guide to Media Resources)** (ISBN 0-13-017677-X). This special edition of the text provides marginal notes and information for instructors and TAs, including teaching tips, suggested lecture demonstrations, references to the chemical education literature, and icons identifying all art that appears on overhead transparencies and on the Media Portfolio CD-ROM for instructors.

**Media Portfolio: Your Presentation Resource CD-ROM (dual platform; ISBN 0-13-017686-9).** Specific to Petrucci/Harwood/Herring, this CD includes almost all art and photos from the book, over 61 short animations, 31 video demonstrations, and Java and Flash simulations from the Student iBook. All these pieces are presented in a thumbnail catalog format that allows easy porting of the files to presentation software such as MS PowerPoint®. Also included are electronic versions of suggested course outlines (which can be edited), a set of pre-built MS PowerPoint slides, and a special chemistry font that lets you quickly edit and add to the electronic files on the CD.

The **Instructor's Resource Manual** (ISBN 0-13-017678-8) by Michael L. Denniston, Georgia Perimeter College and Robert K. Wismer, Millersville University. Ideal for novice instructors or others using this text for the first time, this book integrates all ancillary material, offers Notes for the Instructor, lists key concepts, itemizes Chapter Objectives, and contains the solutions to the Advanced and Integrative Problems not found in Appendix F of the text.

**Transparencies** (ISBN 0-13-017685-0) Includes over 250 full-color images from this text. Each of these is also provided in electronic form on the MediaPortfolio Instructor's CD.

**Test Item File** (ISBN 0-13-017679-6) by C. Alton Hassell, Baylor University. This hardcopy test bank includes over 2000 unique questions, each accuracy checked for

► New testing software

this new edition and not available to students. These questions are also available in WebCT format for adopting institutions.

**Prentice Hall Test Manager** This newly updated testing software includes all 2000 questions from the Test Item File and permits easy creation and editing of quizzes. The software allows easy porting of quizzes into MS Word® format and also supports administration of quizzes over a LAN. Available in both Macintosh (*ISBN 0-13-017681-8*) and Windows (*ISBN 0-13-017670-2*) formats.

**Solutions Manual** (*ISBN 0-13-017683-4*) by Lucio Gelmini and Robert Hiltz, both of Grant MacEwan College, and Robert K. Wismer, Millersville University. Contains completely revised, step-by-step solutions to all end-of-chapter (exercises except for eMedia exercises and the Advanced and Integrative Problems found in Appendix F of text). With instructor permission, these manuals may be made available to students.

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### For the Laboratory

**Experiments in General Chemistry** (*ISBN 0-13-017688-5*) by Gerald S. Weiss, Thomas G. Greco and Lyman H. Rickard, all at Millersville University. A comprehensive laboratory manual containing 37 experiments that parallel the text, including a final group of six experiments on qualitative cation analysis. There is an accompanying instructor's manual (*ISBN 0-13-017689-3*).

### For the Student

► Powerful new student supplement—an interactive electronic version of the text.

**Student iBook** (*ISBN 0-13-017680-X*) by Scott Perry, University of Houston. This interactive version of the text includes hundreds of animations, simulations, manipulable molecular models, movies, and interactive glossary terms, all integrated in-context within an electronic version of the text. Accessed easily using a web browser, this product allows students to see and discover chemistry in ways never before possible. Each interactive exercise is followed by a self-assessment question so students can make sure they understand the key points before moving on to the next topic. Organized exactly like the book, this product is available free with every new copy of the text.

► Enhanced student website features algorithmic questions.

**The Petrucci/Harwood/Herring Companion Website** [www.prenhall.com/petrucci](http://www.prenhall.com/petrucci) by Narayan S. Hosmane, Northern Illinois University. Now in its "second edition," this innovative online resource center is designed to specifically support and enhance students use of Petrucci/Harwood/Herring 8/e. It features

- A Problem Solving Center, where student have access to more than 2000 additional problems, including algorithmically generated questions and non-multiple-choice problems—all organized by chapter, with hints and specific feedback.

- A Visualization Center, where students can view hundreds of pre-built 3-D molecular models using Chime.
- Current Topics, where recent articles from the popular press are summarized and further questions are posted for students to answer on paper or online

**Student Study Guide (ISBN 0-13-032567-8)**, by Dixie Goss of Hunter College and Robert K. Wismer of Millersville University, guides students through the text's coverage with discussion of chapter learning objectives, drill problems, self quizzes, and sample tests.

**Student Solutions Manual (ISBN 0-13-017684-2)**, by Lucio Gelmini and Robert Hilts, both of Grant MacEwan College, and Robert Wismer of Millersville University. Contains full, step-by-step solutions to the red-numbered problems from the text (those answered in Appendix F of the textbook).

► New student supplement.

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**WARNING:** Many of the compounds described or pictured in this text are hazardous, as are many of the chemical reactions. Do not attempt any experiment pictured or implied in the text except with permission, in an authorized laboratory setting, and under adequate supervision.

# Student's Guide to Using this Text

The next six pages walk you through some of the main features of this text and its integrated media resources. Using this text as designed will help you develop the essential knowledge and skills you need to succeed in chemistry. Good luck!

**Keep in Mind Margin Notes** ▶  
These will help to remind you of ideas introduced earlier in the text that are important to understand what's currently being discussed.

**KEEP IN MIND** ▶  
that if you know any four of the five quantities— $q$ ,  $m$ , specific heat,  $T_f$ ,  $T_i$ —you can solve equation (7.5) for the remaining one.

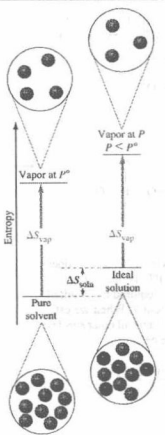


**Practice Example A:** When 1.00 kg lead (specific heat =  $100.0\text{ }^\circ\text{C}$  is added to a quantity of water at  $28.5\text{ }^\circ\text{C}$ , the final temperature of the mixture is  $35.2\text{ }^\circ\text{C}$ . What is the mass of water present?

**Practice Example B:** A 100.0-g copper sample (specific heat =  $100.0\text{ }^\circ\text{C}$  is added to 50.0 g water at  $26.5\text{ }^\circ\text{C}$ . What is the final temperature of the copper–water mixture?

**Significance of Specific-Heat Values**  
Table 7.1 lists the specific heats of several solid elements. The specific heat of aluminum compared with other metals helps to account for the “miracle thaw” products designed to thaw frozen foods rapidly. Aluminum cools only slowly as it transfers heat to the frozen food, and the

**Student iBook icon** ▶  
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the equilibrium vapor pressure  $P^*$ , entropy increases by the amount  $\Delta S_{\text{vap}}$  in Figure 20-6, because the entropy of the ideal solution is greater than that of the pure solvent, the entropy of the vapor produced by the vaporization of a pure solvent is also greater than the entropy of the vapor obtained from a solution. For the vapor above the solution to have the higher entropy, it must have a greater number of accessible microscopic energy levels. This requires a larger volume and hence a lower pressure than the vapor coming from the pure solvent. This relationship is Raoult's law:  $P_A = x_A P_A^*$ .

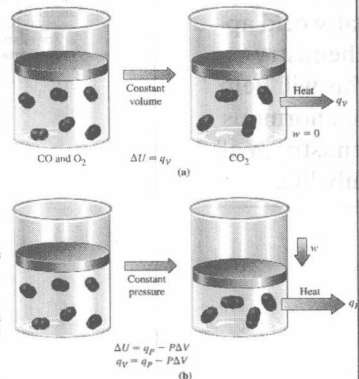
**Absolute Entropies**  
To establish an *absolute* value of the entropy of a substance, we look for a state in which the substance is in its lowest possible energy state, called the *ground state*. We take this state to have an entropy of zero. Then we evaluate changes as the substance is brought to other conditions of temperature and pressure. We add together these entropy changes and obtain a numerical value for the absolute entropy. The principle that permits this procedure is the third law of thermodynamics, which can be stated as follows:  
The entropy of a pure perfect crystal at 0 K is zero.

Figure 20-7 illustrates the method outlined in the preceding paragraph for determining absolute entropy as a function of temperature. Where phase transitions occur, equation (20.2) is used to evaluate the corresponding entropy change. In temperature ranges in which there are no transitions,  $\Delta S^\circ$  values are obtained from measurements of specific heats as a function of temperature.

The absolute entropy of one mole of a substance in its standard state is the *standard molar entropy*,  $S^\circ$ . Standard molar entropies of a number of substances at  $25\text{ }^\circ\text{C}$  are tabulated in Appendix D. To use these values to calculate the entropy change of a reaction, we use an equation with a familiar form: equation 7.21).

▲ FIGURE 20-6  
An entropy-based rationale of Raoult's law

**Molecular Art** ▼  
It is sometimes difficult to visualize molecules and processes that can't be seen directly. To help you understand what's going on at the molecular level, carefully review and make sure you understand the molecular depictions provided in the text.



► FIGURE 7-13  
Comparing heats of reaction at constant volume and constant pressure for the reaction:  $2\text{ CO(g)} + \text{O}_2\text{(g)} \rightarrow 2\text{ CO}_2\text{(g)}$   
(a) No work is performed at constant volume because the piston cannot move;  $q_v = \Delta U = -563.5\text{ kJ}$  (b) When the reaction is carried out at constant pressure, the surroundings do work on the system as the system shrinks into a smaller volume. More heat is evolved than in the constant-volume reaction;  $q_p = \Delta H = -566.0\text{ kJ}$ .



## Are You Wondering ...

Why  $\text{Na}^+(\text{aq})$  does not act as an acid in aqueous solution?

Whether an aqueous solution of a metal ion is acidic depends on two principal factors. The first is the amount of charge on the cation; the second is the size of the ion. The greater the charge on the cation, the greater is the ability of the metal ion to draw electron density away from the O—H bond in a  $\text{H}_2\text{O}$  molecule in its hydration sphere, favoring the release of a  $\text{H}^+$  ion. The smaller the cation, the more highly concentrated is the positive charge. Hence, for a given positive charge, the smaller the cation, the more acidic the solution.

The ratio of the charge on the cation to the volume of the cation is called the *charge density*.

$$\text{charge density} = \frac{\text{ionic charge}}{\text{ionic volume}}$$

The greater its charge density, the more effective a metal ion is at pulling electron density from the O—H bond and the more acidic is the hydrated cation. A highly concentrated positive charge on a small cation is better able to pull electron density from the O—H bond than is a less concentrated positive charge on a larger cation.

Thus the small (53 pm ionic radius), highly charged  $\text{Al}^{3+}$  ion produces acidic solutions, but the larger (99 pm)  $\text{Na}^+$  cation, with a charge of just  $1+$ , does not increase the concentration of  $\text{H}_3\text{O}^+$ . In fact, none of the group 1 cations produces appreciably acidic solutions, and only  $\text{Be}^{2+}$  of the group 2 elements is small enough to do so ( $\text{p}K_a = 5.4$ ).

## Are You Wondering?

Always ask questions about what you're learning. As you read the text, new ideas may prompt you to raise important queries. The "Are You Wondering?" boxes answer good questions that students frequently ask.

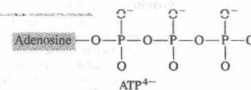
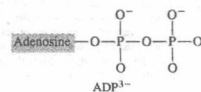
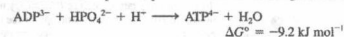
## Focus On Coupled Reactions in Biological Systems



### ▲ Mitochondria and endoplasmic reticulum

A colorized scanning electron micrograph of mitochondria (blue) and rough endoplasmic reticulum (yellow) in a pancreatic cell. Mitochondria are the powerhouses of the cell. They oxidize sugars and fats, producing energy for the conversion of ADP to ATP. Rough endoplasmic reticulum is a network of folded membranes covered with protein-synthesizing ribosomes (small dots).

An important example of a coupled reaction in living organisms is the metabolism of glucose ( $\text{C}_6\text{H}_{12}\text{O}_6$ ) that converts adenosine diphosphate (ADP) to adenosine triphosphate (ATP) in the mitochondria of cells (Fig. 20-13). The ATP is utilized in the ribosomes to produce proteins. The ATP-forming reaction



is spontaneous under standard conditions at  $37^\circ\text{C}$ , so why does the cell need to use glucose to make ATP? The answer is that cells do not operate with  $[\text{H}^+] = 1 \text{ M}$  as required by standard conditions; in fact, the pH in a cell is about 7. When we estimate  $\Delta G$  for the reaction at that pH and assume all other species are at  $1.0 \text{ M}$  (still far from actuality), we get

$$\Delta G = \Delta G^\circ + RT \ln \left( \frac{a_{\text{ATP}} a_{\text{H}_2\text{O}}}{a_{\text{ADP}} a_{\text{H}} a_{\text{HPO}_4}} \right)$$

Note that reactions (20.15) and (20.16) are not the same, even though each has  $\text{Cu}(s)$  as a product. The purpose of coupled reactions, then, is to produce a spontaneous overall reaction by combining two other processes: one nonspontaneous and one spontaneous. Many metallurgical processes employ coupled reactions, especially those that use carbon or hydrogen as reducing agents.

To sustain life, organisms must synthesize complex molecules from simpler ones. If carried out as single-step reactions, these syntheses would generally be accompanied by increases in enthalpy, decreases in entropy, and increases in free energy—in short, they would be nonspontaneous and would not occur. In living organisms, changes in temperature and electrolysis are not viable options for dealing with nonspontaneous processes. Here, coupled reactions are crucial, as described in the Focus On feature.

## "Focus On" Boxes

You'll find one of these at the end of each chapter. They detail interesting, real-world applications of the chemistry you're learning. This will help you appreciate the enormous importance of chemistry in all aspects of your daily life.



## Problem-Solving

Worked Examples are followed by A & B Practice Exercises. Read carefully through each Worked Example. They help you understand how to solve important types of problems. To help you practice, each worked example is followed by two Practice Examples. Practice Example A is another problem very similar to the Worked Example. Practice Example B is more difficult, closer to the end-of-chapter problems and those you'll find on an exam.

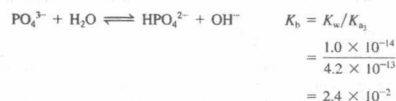
## Problem-Solving Notes

These marginal reminders warn you of common mistakes and highlight key strategies for solving particular types of problems.

► Sodium phosphate is often sold under the name trisodium phosphate (TSP). Anyone working with TSP should wear protective gloves. Fats and greases, including those in human skin, are solubilized in strongly basic solutions.

► The ratio  $M_b/K_b = 1.0/0.024 = 42$ . This value is smaller than the minimum value of 100 that we have been using as the usual criterion.

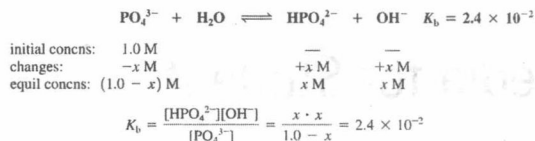
is not difficult to calculate. It corresponds to that of  $\text{Na}_2\text{PO}_4(\text{aq})$ , and  $\text{PO}_4^{3-}$  can ionize (hydrolyze) only as a base.



*Determining the pH of a Solution Containing the Anion ( $A^{n-}$ ) of a Polyprotic Acid.* Sodium phosphate,  $\text{Na}_3\text{PO}_4$ , is an ingredient of some preparations used to clean painted walls before they are repainted. What is the pH of 1.0 M  $\text{Na}_3\text{PO}_4(\text{aq})$ ?

### Solution

In the usual fashion, we can write



Because  $K_b$  is quite large, we should not expect the usual simplifying assumption to work here. That is,  $x$  is *not* very much smaller than 1.0. Solution of the quadratic equation  $x^2 + 0.024x - 0.024 = 0$  yields  $x = [\text{OH}^-] = 0.14 \text{ M}$ . Thus,

$$\begin{aligned}\text{pOH} &= -\log[\text{OH}^-] = -\log 0.14 = +0.85 \\ \text{pH} &= 14.00 - 0.85 = 13.15\end{aligned}$$

**Practice Example A:** Calculate the pH of an aqueous solution that is 1.0 M  $\text{Na}_2\text{CO}_3$ .

(Hint: Use data from Table 17.4 to establish  $K_b$  for  $\text{CO}_3^{2-}$ .)

**Practice Example B:** Calculate the pH of an aqueous solution that is 0.500 M  $\text{Na}_2\text{SO}_3$ .

(Hint: Use data from Table 17.4.)

It is more difficult to calculate the pH values of  $\text{NaH}_2\text{PO}_4(\text{aq})$  and  $\text{Na}_2\text{HPO}_4(\text{aq})$  than of  $\text{Na}_3\text{PO}_4(\text{aq})$ . This is because with both  $\text{H}_2\text{PO}_4^-$  and  $\text{HPO}_4^{2-}$ , two equilibria must be considered *simultaneously*: ionization as an acid and ionization as a base (hydrolysis). For solutions that are reasonably concentrated (say, 0.10 M or greater), the pH values prove to be *independent* of the solution concentration. Shown here (with  $\text{p}K_a$  values from Table 17.4) are general expressions, printed in blue, and their application to  $\text{H}_2\text{PO}_4^-(\text{aq})$  and  $\text{HPO}_4^{2-}(\text{aq})$ :

$$\text{for } \text{H}_2\text{PO}_4^-: \quad \text{pH} = \frac{1}{2}(\text{p}K_{a_1} + \text{p}K_{a_2}) = \frac{1}{2}(2.15 + 7.20) = 4.68 \quad (18.5)$$

$$\text{for } \text{HPO}_4^{2-}: \quad \text{pH} = \frac{1}{2}(\text{p}K_{a_2} + \text{p}K_{a_3}) = \frac{1}{2}(7.20 + 12.38) = 9.79 \quad (18.6)$$

## Integrative Example

Microwave ovens have become increasingly popular in kitchens around the world. They are also useful in the chemical laboratory, particularly in drying samples for chemical analysis. A typical microwave oven uses microwave radiation with a wavelength of 12.2 cm.

Are there any electronic transitions in the hydrogen atom that could conceivably produce microwave radiation of wavelength 12.2 cm?

1. Calculate the frequency of the microwave radiation. Microwaves are a form of electromagnetic radiation and thus travel at the speed of light,  $2.998 \times 10^8 \text{ m s}^{-1}$ . Convert the wavelength to meters, and then use the equation  $\nu = c/\lambda$ .

$$\nu = \frac{2.998 \times 10^8 \text{ m s}^{-1}}{12.2 \text{ cm} \times 1 \text{ m}/100 \text{ cm}} = 2.46 \times 10^9 \text{ Hz}$$

2. Calculate the energy associated with one photon of the microwave radiation. This is a direct application of Planck's equation.

$$E = h\nu = 6.626 \times 10^{-34} \text{ J s} \times 2.46 \times 10^9 \text{ s}^{-1} = 1.63 \times 10^{-24} \text{ J}$$

3. Determine if there are any electronic transitions in the hydrogen atom with an energy per photon of  $1.63 \times 10^{-24} \text{ J}$ . Look at Figure 9-14, the energy-level diagram for the Bohr hydrogen atom. Energy differences between the low-lying levels are of the order  $10^{-19}$  to  $10^{-20} \text{ J}$ . These are orders of magnitude ( $10^4$ – $10^5$  times) greater than the energy per photon of  $1.63 \times 10^{-24} \text{ J}$  from part 2. Note, however, that the energy differences become progressively smaller for high-numbered orbits. As  $n$  approaches  $\infty$ , the energy differences approach zero, and some transitions between high-numbered orbits should correspond to microwave radiation (see also Exercise 101).

## Integrative Examples

These conclude each chapter and serve two purposes: they summarize many of the key concepts in the chapter, and they show you how to approach more complicated, integrative exercises (like those at the end of the chapter).