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## Chemistry: Atoms First 2e Release Notes 2022

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## Page Count Difference:

Changes in formatting and design have allowed the page count to be reduced from 1327 in the previous revision to 1211 in the new revision.

## Errata:

Below is a table containing submitted errata and the resolutions that OpenStax has provided for this latest text. Beyond errata and related updates, this release includes significant additions of more diverse scientists and increased discussions of discoveries and scientific processes. New narratives include historical and recent contributors such as Alma Levant Hayden, Percy Lavon Julian, Lise Meitner, Alice Ball, Susan Kwolek, Reatha Clark King, and many others. These expansions are aimed at engaging students in the personal and practical impacts and opportunities related to chemical principles.

| Location | Detail | Resolution Notes | Error Type |
| :--- | :--- | :--- | :--- |
| Preface | Image in Chem 2e and Atoms First 2e prefaces <br> appears only there and nowhere else in the <br> book, so there is no alt text available to assign <br> to it. CNX_Chem_06_03_3dOrbitals.jpg | Alt text will be added to this figure. | Other |
| Chapter 1 Essential <br> Ideas: Section 1.1 <br> Chemistry in <br> Context | Exercise 51(f) In the lesson you say "if the <br> dropped digit is 5, round up or down, <br> whichever yields an even value for the retained <br> digit". But in the answer to exercise 51 f, you <br> round 0.445 with two sig. dig. to 0.45 instead of <br> 0.44 | Revise the answer to "0.44". | Incorrect <br> answer, <br> calculation, or <br> solution |
| Chapter 1 Essential <br> Ideas: Section 1.2 <br> Phases and <br> Classification of <br> Matter | The author wrote that "Pure substances that <br> can be broken down by chemical changes are <br> called compounds.", however, this definition <br> seems inaccurate. For example, we won't view <br> H2 and O2 as compounds, that's because they <br> aren't consist of two or more than two kinds of <br> elements. Nevertheless, if we apply the <br> author's definition in this case, we can find out <br> that H2 + O2 = H2O, and in this reaction, both <br> of the reactants are actually broken down by <br> chemical changes while both of which aren't <br> categorized as compounds. Therefore, I suggest <br> that we should change the definition into "a | Revise "Pure substances that can be <br> broken down by chemical changes <br> are called compounds. This <br> breakdown may produce either <br> elements or other compounds, or <br> both." to "Pure substances that are <br> comprised of two or more elements <br> are called compounds. Compounds <br> may be broken down by chemical <br> changes to yield either elements or <br> other compounds, or both." | Other factual <br> inaccuracy in <br> content |


|  | pure substance consist of two or more than two kinds of elements." |  |  |
| :---: | :---: | :---: | :---: |
| Chapter 1 Essential <br> Ideas: Section 1.4 <br> Measurements | Example 1.2 There is a minus sign "-" where there should be an equal sign "=". " 29.5 mL $25.5 \mathrm{~mL}-4.0 \mathrm{~mL}$ " should be " $29.5 \mathrm{~mL}-25.5 \mathrm{~mL}$ $=4.0 \mathrm{~mL}{ }^{\prime \prime}$ | Our reviewers accepted this change, and it will be included in the next print cycle. | Typo |
| Chapter 1 Essential Ideas: Section 1.4 Measurements | In para. 2 under Density, the phrase "this as an inconvenient unit" should be "this is an inconvenient unit". | Our reviewers accepted this change, and it will be included in the next print cycle. | Typo |
| Chapter 1 Essential <br> Ideas: Section 1.4 <br> Measurements | Exercise 41 has 5 questions. Answer (d) is the answer of question (e), and there's no answer to question (d). Answer to (d) should be: red < green < blue < yellow. | Add "red < green < blue < yellow;" as the answer to part (d), and add "(e)" before "If the volumes are..." | Incorrect answer, calculation, or solution |
| Chapter 1 Essential <br> Ideas: Section 1.4 <br> Measurements | End of chapter Qs c) asking for base SI units so speed of sound should be $\mathrm{m} / \mathrm{s}$ not $\mathrm{km} / \mathrm{s}$ | Revise from kilometers/second to meters/second. | Other factual inaccuracy in content |
| Chapter 1 Essential <br> Ideas: Section 1.4 <br> Measurements | example 1.2 <br> http://openstax.org/l/16phetmasvolden Flash is no longer supported. | This link and exercise, as well as the related questions (41-43) will be updated. | Broken link |
| Chapter 1 Essential Ideas: Section 1.5 Measurement Uncertainty, Accuracy, and Precision | The rule about rounding " 5 " to an even digit (list item 3 under "Significant Figures in Calculations) is not explained correctly. This rule only applies when 5 is the digit being dropped *and is the last nonzero digit*. The later qualifier is important because if the unrounded number has nonzero digits following that 5 , then it is closer to the rounded up value than the rounded down value. This error is reinforced by the third bullet point example, which gives an incorrect explanation: " 6.8752 rounds "up" to 6.88 (the dropped digit is 5 , and the retained digit is even)". The correct reason to round up is because 6.8752 is closer to 6.88 (abs. difference $=0.0048$ ) than to 6.87 (abs. difference $=$ 0.0052 ). 6.7852 should still be rounded up to 6.79 and not to the even digit 6.78 for this same reason. The "bias" argument for rounding to the even applies to a single non-zero digit being dropped because there are only four digits that would go down (1-4) and five that would go up (5-9) using "normal" or "high school math" rounding rules. If the dropped digit is assumed to be uniformly distributed, this is a pretty significant bias towards increasing the value. The round to the even rule fixes this by having 5 go up as often as down (also assumes odd is as likely as even for the last retained digit). With two digits that get rounded away, the bias is only 50 up to 49 down, and is similarly removed if the only two digit case that is rounded to the even is 50 . Rounding 51 through 59 to the nearest even while still always rounding 41-49 actually creates a downward bias (there are now 49 always round down pairs of digits to 40 | Revise rule 3 as follows: <br> "If the digit to be dropped (the one immediately to the right of the digit to be retained) is less than 5, "round down" and leave the retained digit unchanged; if it is more than 5, "round up" and increase the retained digit by 1 . If the dropped digit is 5 , and it's either the last digit in the number or it's followed only by zeros, round up or down, whichever yields an even value for the retained digit. If any nonzero digits follow the dropped 5, round up. (The last part of this rule may strike you as a bit odd, but it's based on reliable statistics and is aimed at avoiding any bias when dropping the digit "5," since it is equally close to both possible values of the retained digit.)" <br> Revise the third bullet below this to the following: <br> " 6.8752 rounds "up" to 6.88 (the dropped digit is 5, and a nonzero digit follows it)" |  |


|  | always round up pairs). Same argument applies no matter how many digits are being dropped. |  |  |
| :---: | :---: | :---: | :---: |
| Chapter 1 Essential Ideas: Section 1.5 <br> Measurement Uncertainty, <br> Accuracy, and Precision | Example 1.5, Solution of (b). is presented as $0.86728 \ldots \mathrm{~g} / \mathrm{mL}$ when the actual correct answer should be $0,866728 \ldots \mathrm{~g} / \mathrm{mL}$. | Revise " 0.86728 " to "0.866728". | Typo |
| Chapter 1 Essential Ideas: Section 1.6 Mathematical Treatment of Measurement Results | I think Question 71 is intended to link to table 1.2 instead of 1.3 | Our reviewers accepted this change. | Broken link |
| Chapter 1 Essential Ideas: Section 1.6 Mathematical Treatment of Measurement Results | Under "Learning objectives" and before "Conversion factors and dimensional analysis" It says $(\mathrm{m} / \mathrm{m} / \mathrm{s})=\mathrm{s}$. Based on order of operations, this would be $m / m=1$, then $1 / s=s^{\wedge}(-1)$. It needs to use $(\mathrm{m} /(\mathrm{m} / \mathrm{s}))=$ s because then if you are dividing by a fraction, you are multiplying by the reciprocal, so $\mathrm{m}^{*}(\mathrm{~s} / \mathrm{m})=\mathrm{s}$ | Revise " $(\mathrm{m} / \mathrm{m} / \mathrm{s}=\mathrm{s})$ " to " $(\mathrm{m} /(\mathrm{m} / \mathrm{s})=$ s)". | Incorrect answer, calculation, or solution |
| Chapter 1 Essential Ideas: Section 1.6 Mathematical Treatment of Measurement Results | It says $(\mathrm{m} / \mathrm{m} / \mathrm{s})=\mathrm{s}$. Based on order of operations, this would be $m / m=1$, then $1 / s=s^{\wedge}(-1)$. It needs to use $(\mathrm{m} /(\mathrm{m} / \mathrm{s}))=$ s because then if you are dividing by a fraction, you are multiplying by the reciprocal, so $\mathrm{m} *(\mathrm{~s} / \mathrm{m})=\mathrm{s}$ | Revise " $(\mathrm{m} / \mathrm{m} / \mathrm{s}=\mathrm{s})$ " to " $(\mathrm{m} /(\mathrm{m} / \mathrm{s})=$ s)". | Incorrect answer, calculation, or solution |
| Chapter 1 Essential Ideas: Section 1.6 Mathematical Treatment of Measurement Results | The time period should be listed as the late 19th Century rather than the late 1800s because late 1800s implies 1807-1809, which is not the time when the cathode ray tube experiments were performed. | Revise the sentence "The mercury or alcohol in a common..." to "The liquid in a common glass thermometer changes its volume as the temperature changes, and the position of the trapped liquid's surface along a printed scale may be used as a measure of temperature." | Other factual inaccuracy in content |
| Chapter 1 Essential Ideas: Section 1.6 Mathematical Treatment of Measurement Results | In example 1.9, before it says the first two steps for volume converison (how to get from quarts to millileter), it says "Volume may be converted from quarts to millimeters via two steps," when it should say millileters. | Revise "millimeters" to "milliliters". | Typo |
| Chapter 1 Essential Ideas: Section 1.6 Mathematical Treatment of Measurement Results | In the equation relating the temperature ( T ) scales converting from Celsius to Fahrenheit the book states incorrectly +32 C and it should be corrected to 32 F. Otherwise, you are teaching to add apples and oranges to produce apples. | Revise C to F. | Other factual inaccuracy in content |


| Chapter 1 PowerPoint, Slide 32 | Instructor PowerPoint material, Chapter 1, Slide 32 - it should read as 'most elements exist as individual atoms 'only a few exist as molecules' | This slide will be deleted from the PowerPoint. | Other factual inaccuracy in content |
| :---: | :---: | :---: | :---: |
| Chapter 1, Chapter <br> 3, Appendix D | Update to revise the definitions of several fundamental physical constants that were adopted May 2019. | Revise as indicated. | General/pedago gical suggestion or question |
| Chapter 2 Atoms, Molecules, and Ions: Section 2.1 Early Ideas in Atomic Theory | Example 2.2 In the answer, the book reads " In compound Y , the mass ratio of carbon to oxygen is" when it should be " In compound Y , the mass ratio of carbon to hydrogen is" | Revise oxygen to hydrogen. | Typo |
| Chapter 2 Atoms, Molecules, and Ions: Section 2.1 Early Ideas in Atomic Theory | In the molecules represented at the right of the figure, the angle between the aldehyde function and the phenyl ring in the benzaldehyde differs from the ideal 120 degrees. | This figure will be updated. | Other factual inaccuracy in content |
| Chapter 2 Atoms, Molecules, and Ions: Section 2.1 Early Ideas in Atomic Theory | Example 2.2's solution explanation states "...with A having one-half as much carbon per amount of oxygen (or twice as much carbon per amount of oxygen) as B." Carbon and oxygen should be switched in the first part. It should read: with A having one-half as much oxygen per amount of carbon... | Revise "carbon" to "oxygen" and vice versa. | Typo |
| Chapter 2 Atoms, Molecules, and Ions: Section 2.3 Atomic Structure and Symbolism | Exercise 25 The atomic mass listed in Ch 2 problem 25 for B-11 is incorrect. It should be 11.00931 amu, not 11.0931 amu. | This will be updated in the Answer Key and solution manual. | Other factual inaccuracy in content |
| Chapter 2 Atoms, <br> Molecules, and lons: <br> Section 2.3 Atomic <br> Structure and <br> Symbolism | typo in one occurrence of Oxygen-18 (e.g., <br> "Oxyen-18" should be "Oxygen-18") | Our reviewers accepted this change, and it will be included in the next print cycle. | Typo |
| Chapter 2 Atoms, <br> Molecules, and Ions: <br> Section 2.4 <br> Chemical Formulas | At the end of Chapter 2, right before the Key Terms, there's a "How Sciences Interconnect" section with a picture of the brain on the left and neuron cells on the right. Immediately above the picture, the last sentence in that paragraph links to a White House article from the Obama administration that's since been archived. (see first screenshot) I did a little digging and found the original linked article on an archived web page, but think this link would be better, since it's the BRAIN Initiative website: https://braininitiative.nih.gov/ | This link will be updated. | Broken link |
| Chapter 2 Atoms, <br> Molecules, and Ions: <br> Section 2.4 <br> Chemical Formulas | Value of the mole in text $6.02214179 \times 1023$. New value is $6.02214076 \times 1023$ | Our reviewers accepted this change. | General/pedago gical suggestion or question |


| Chapter 2 Atoms, Molecules, and Ions: <br> Section 2.4 <br> Chemical Formulas | Chapter 2 Problem 55 Answer in the back of the book. The problems asks for greatest mass; however the answer provided is in moles. | Revise the solution to exercise 55 as follows: <br> AIPO4: 1.000 mol or 26.98 g Al Al2CI6: 1.994 mol or 53.74 g Al Al2S3: 3.00 mol or 80.94 g Al The Al2S3 sample thus contains the greatest mass of Al. | Typo |
| :---: | :---: | :---: | :---: |
| Chapter 2 Atoms, Molecules, and Ions: Section 2.4 Chemical Formulas | Chapter 2 Exercise 31, you define the molecular formula of fructose as C 12 H 22 O 11 . However, this is incorrect: C 12 H 22 O 11 is actually the formula for table sugar, while the formula for fructose is C 6 H 12 O 6. | Revise "fructose" to "sucrose". | Other factual inaccuracy in content |
| Chapter 2 Atoms, Molecules, and Ions: <br> Section 2.4 <br> Chemical Formulas | \#48 c) last item listed should be $6.9 \times 102$ moles of OXYGEN | The solution will be updated. | Incorrect answer, calculation, or solution |
| Chapter 2 Atoms, Molecules, and Ions: Section 2.4 Chemical Formulas | Human cells do not contain cell walls. Instead, "cell membrane" should be printed instead of "cell wall". | Revise "cell wall" to "cell membrane". | Other factual inaccuracy in content |
| Chapter 2 Atoms, Molecules, and Ions: Section 2.4 Chemical Formulas | Ch 2 Problem \#61 asks for the lowest number of molecules in an assortment of compounds. It wants students to calculate the number of molecules in each compound. By these calculations, the answer is correct, however technically, the answer is the CaH 2 (Calcium hydride) compound, regardless of quantity. Because CaH 2 is ionic, it contains zero molecules. Suggest that either that compound is removed or problem is changed to ask for formula units. | Revise "CaH_2 $(42.09 \mathrm{~g} / \mathrm{mol})$ " to "C_3H_6 (42.08 g/mol)." | Incorrect answer, calculation, or solution |
| Chapter 2 Atoms, Molecules, and Ions: Section 2.4 Chemical Formulas | "Determine which of the following contains the greatest mass of aluminum: 122 g of AIPO4, 266 g of Al2C16, or 225 g of Al2S3." The "1" after the "C" in Al2C16 should be an "I" (lower case ell): Al2Cl6 | Revise "1" to lowercase "I". | Typo |
| Chapter 2 Atoms, Molecules, and Ions: Section 2.7 <br> Molecular and Ionic Compounds | In Problem 98, revise $\mathrm{MgC2H} 3 \mathrm{O} 2$ to $\mathrm{Mg}(\mathrm{C} 2 \mathrm{H} 3 \mathrm{O} 2) 2$ | In exercise 98 part b, revise "MgC2H3O2" to "Mg(C2H3O2)2". | Other factual inaccuracy in content |
| Chapter 3 Electronic Structure and Periodic Properties of Elements: Section 3.1 Electromagnetic Energy | I'm troubled that men get mentioned winning the noble prize throughout the textbook... but Dorothy Hodgkin's 1964 Nobel Prize in Chemistry isn't mentioned in her "Portrait of a Chemist". Seems a bit sexist and doesn't set a inclusive tone to skip her awards -- especially when awards for men are mentioned often in the text. | The Portrait of a Chemist box on Dorothy Hodgkin will be updated. | Other |
| Chapter 3 Electronic Structure and Periodic Properties | \#26 show work! Not just the answer. | The full solution will be added. | General/pedago gical suggestion or question |


| of Elements: Section 3.2 The Bohr Model |  |  |  |
| :---: | :---: | :---: | :---: |
| Chapter 3 Electronic Structure and Periodic Properties of Elements: Section 3.2 The Bohr Model | Question 28 describes multiple hydrogens with electrons in $n=1,2,3,4$ *orbitals*. This is confusing because the quantum number $n$ refers to the shells of the atom, which contain multiple orbitals. The quantum number m_l would refer to the orbitals. | Revise "orbital" to "orbit" and "orbitals" to "orbits" in section 6.2 and problem 28. | Other factual inaccuracy in content |
| Chapter 3 Electronic Structure and Periodic Properties of Elements: Section 3.2 The Bohr Model | The text reads "The sizes of the circular orbits for hydrogen-like atoms are given in terms of their radii by the following expression, in which $\alpha 0$ is a constant called the Bohr radius, with a value of $5.292 \times 10-11 \mathrm{~m}$ :" The equation listed uses a0, rather than $\alpha 0$. | Revise " $\alpha$ " to "a" in the text before the equation. | Typo |
| Chapter 3 Electronic Structure and Periodic Properties of Elements: Section 3.2 The Bohr Model | In the teacher solutions to chapter 3 \#27, Joules are converted to eV, though the question asks for Joules. Was the question intended to ask for eV? | This solution will be updated. | Incorrect answer, calculation, or solution |
| Chapter 3 Electronic Structure and Periodic Properties of Elements: Section 3.3 Development of Quantum Theory | The highlighted value should be $\hbar$, not $h$. $h$ is the Planck's constant with value $6.626 \times 10^{\wedge}$ 34 Js . $\hbar=\mathrm{h} /(2 \mathrm{pi})$, which is $1.055 \times 10^{\wedge}$ - 34 | Our reviewers accepted this change, and it will be included in the next print cycle. | Incorrect answer, calculation, or solution |
| Chapter 3 Electronic Structure and Periodic Properties of Elements: Section 3.3 Development of Quantum Theory | That h (highlighted one) should be a ' $\dagger$ '. | Revise "h" to " $\dagger$ ". | Typo |
| Chapter 3 Electronic <br> Structure and <br> Periodic Properties <br> of Elements: Section <br> 3.4 Electronic <br> Structure of Atoms <br> (Electron <br> Configurations) | Within the key terms, discussing transition elements, a sentence reads as "The valence electrons (those added after the last noble gas configuration) in these elements include the ns and ( $n-1$ ) d electrons." The electrons being referred to are the 'outer' electrons and not the 'valence' electrons. Then, within the instructor's powerpoint slides, where it says the same incorrect sentence, there is an example for Vanadium, discussing it has 5 valences electrons. Again, this should be corrected to say 5 outer electrons and followed up with 2 valence electrons (being the 4 s 2 as they can extend furthest form the nucleus). | While our reviewers determined that the current text is consistent with the widespread usage of the term "valence," revisions will be made to the definitions of valance electrons and valance shell in the Key Terms section. <br> Revise the definition of valance electron to "electrons in the high energy outer shell(s) of an atom" and valance shell to "high energy outer shell(s) of an atom". | Other factual inaccuracy in content |
| Chapter 3 Electronic Structure and Periodic Properties of Elements: Section 3.4 Electronic Structure of Atoms (Electron Configurations) | At exercise 47, in the answer key, you write the wrong electron configuration for $\mathrm{Ca} 2+$ | $\begin{aligned} & \text { Revise "Ca^2+: } 1 s^{\wedge} 22 s^{\wedge} 22 p^{\wedge} 6 \text { " to } \\ & " C a^{\wedge} 2+: 1 s^{\wedge} 22 s^{\wedge} 22 p^{\wedge} 63 s^{\wedge} 23 p^{\wedge} 6 " \text { " } \end{aligned}$ | Incorrect answer, calculation, or solution |


| Chapter 3 Electronic <br> Structure and <br> Periodic Properties <br> of Elements: Section <br> 3.4 Electronic <br> Structure of Atoms <br> (Electron <br> Configurations) | Figure 3.27 is supposed to show the electron configuration for each subshell at top right of each element. As such, it gives a $4 f \wedge 1$ configuration for $\mathrm{Ce}, 4 f \wedge 2$ for $\mathrm{Pr}, 4 \mathrm{f}^{\wedge} 3$ for Nd etc. This is wrong. The correct electron configurations are shown on figure 6.29: 4f^3 for $\operatorname{Pr}, 4 f \wedge 4$ for $N d$ etc. With the exception of La, Ce and Gd for the lanthanides (more exceptions in the case of the actinides), they follow the rule represented figure 6.26, and correspond to electronic configurations $6 d^{\wedge} 24 f^{\wedge} n$ for La-Yb and $7 s^{\wedge} 25 f^{\wedge} n$ for Ac-No, with no electrons in the $d$ shell. In short, for the purpose of figure 6.27, La and Ac should be seen as part of the $f$ block, and Lu and Lr as part of the $d$ block (irrespective of the larger debate about where to place them in general). Maybe showing the table as in appendix A (with the spaces under Y occupied by La-Lu and Ac-Lr place markers) would provide a more general representation. | This figure and caption will be updated. | Other factual inaccuracy in content |
| :---: | :---: | :---: | :---: |
| Chapter 3 Electronic <br> Structure and <br> Periodic Properties <br> of Elements: Section <br> 3.4 Electronic <br> Structure of Atoms <br> (Electron <br> Configurations) | Shouldn't the electron configuration "1s2 2s2 2p6 3s2 3p6 3d10 4s2 4p6 4d5" be written in this order? "1s2 2s2 2p6 3s2 3p6 4s2 3d10 4p6 4 d 5 " (order change between 3 d 10 and 4 s 2 ) | $\begin{aligned} & \text { Revise } \\ & \text { "1s22s22p63s23p63d104s24p64d5" } \\ & \text { to } \\ & \text { "1s22s22p63s23p64s23d104p64d5". } \end{aligned}$ | Typo |
| Chapter 3 Electronic <br> Structure and Periodic Properties of Elements: Section 3.4 Electronic Structure of Atoms (Electron Configurations) | \#52 a) $2 p$ needs 3 orbitals shown, c) 3d needs 5 orbitals shown | The solution will be updated. | Incorrect answer, calculation, or solution |
| Chapter 3 Electronic Structure and Periodic Properties of Elements: Section 3.4 Electronic Structure of Atoms (Electron Configurations) | Figure 3.27 The table is supposed to show the electron configuration for each subshell at top right of each element. As such, it gives a $4 \mathrm{f}^{\wedge} 1$ configuration for $\mathrm{Ce}, 4 f \wedge 2$ for $\mathrm{Pr}, 4 \mathrm{f} \wedge 3$ for Nd etc. This is wrong. The correct electron configurations are shown on figure 6.29: 4f^3 for $\mathrm{Pr}, 4 \mathrm{f}^{\wedge} 4$ for Nd etc. With the exception of La, Ce and Gd for the lanthanides (more exceptions in the case of the actinides), they follow the rule represented figure 6.26, and correspond to electronic configurations $6 d^{\wedge} 24 f^{\wedge} n$ for La-Yb and $7 s^{\wedge} 25 f^{\wedge} n$ for Ac-No, with no electrons in the $d$ shell. In short, for the purpose of figure 6.27, La and Ac should be seen as part of the $f$ block, and Lu and Lr as part of the $d$ block (irrespective of the larger debate about where to place them in general). Maybe showing the table as in appendix A (with the spaces under Y occupied by La-Lu and Ac-Lr | This figure and caption will be updated. | Other factual inaccuracy in content |


|  | place markers) would provide a more general representation. |  |  |
| :---: | :---: | :---: | :---: |
| Chapter 3 Electronic <br> Structure and Periodic Properties of Elements: Section 3.4 Electronic Structure of Atoms (Electron Configurations) | Under the header "Electron Configurations and the Periodic Table" and under 3. Inner Transition Metals, it says <br> The lanthanide series: lanthanide (La) through lutetium (Lu) <br> The actinide series: actinide (Ac) through lawrencium (Lr) <br> when it should say (stars indicate the error) The lanthanide series: *lanthanum* (La) through lutetium (Lu) <br> The actinide series: *actinium* (Ac) through lawrencium (Lr) | Revise "lanthanide" to "lanthanum" and revise "actinide" to "actinium". | Typo |
| Chapter 3 Electronic <br> Structure and Periodic Properties of Elements: Section 3.4 Electronic Structure of Atoms (Electron Configurations) | The electron configurations sometimes have the wrong order, for example, the solution to 63 gives the order as ... $3 \mathrm{~s}, 3 \mathrm{p}, 3 \mathrm{~d}, 4 \mathrm{~s}, 4 \mathrm{p}, 4 \mathrm{~d}, 5 \mathrm{~s}$ ... when it should be ... $3 \mathrm{~s}, 3 \mathrm{p}, 4 \mathrm{~s}, 3 \mathrm{~d}, 4 \mathrm{p}, 5 \mathrm{~s}, 4 \mathrm{~d}$ ... following Madelung's rule. | Revise the solution to $1 s^{\wedge} 22 s^{\wedge} 22 p^{\wedge} 63 s^{\wedge} 23 p^{\wedge} 64 s^{\wedge} 23 d^{\wedge} 104$ $p^{\wedge} 65 s^{\wedge} 24 d^{\wedge} 105 p^{\wedge} 66 s^{\wedge} 24 f^{\wedge} 145 d^{\wedge} 10$ | Incorrect answer, calculation, or solution |
| Chapter 3 Electronic Structure and Periodic Properties of Elements: Section 3.5 Periodic Variation in Element Properties | The label for Mg is floating away from where it should be and it is unclear which data point it is associated with. It should be in a similar position to the Be label, directly above the data point associated with it. | This figure will be updated to remove the "Mg" label because there is no unambiguous location to place it. | Other |
| Chapter 3 Electronic Structure and Periodic Properties of Elements: Section 3.5 Periodic Variation in Element Properties | Figure 3.36. The electron affinity in the table should be -200 (-20 is currently listed in the table). | This figure will be updated. | Typo |
| Chapter 3 Electronic Structure and Periodic Properties of Elements: Section 3.6 The Periodic Table | \#92 a) Answer $=\mathrm{Cl}$ | The solution will be updated. | Incorrect answer, calculation, or solution |
| Chapter 3 Electronic Structure and Periodic Properties of Elements: Section 3.7 Molecular and Ionic Compounds | In the table of Common Polyatomic Ions the formula for chromic acid is incorrect in both the on-line and paper copies of OpenStax Chemistry. Currently, OpenStax has the formula as H 2 Cr 2 O 4 (forgive the lack of ability to insert the subscripts correctly but I did not see any word processor on-line assistant; this is incorrect. The correct formula for chromic acid is H 2 CrO 4 . One of the themes of Section 2.6 is to teach the students the correct formulas for common compounds and ions. | This formula will be corrected. | Incorrect answer, calculation, or solution |


| Chapter 3 Electronic <br> Structure and Periodic Properties of Elements: Section 3.7 Molecular and Ionic Compounds | Table 3.4 The acid for sulfate is misspelled. It should be sulfuric acid, not sulfiric acid. | Revise to "sulfuric". | Typo |
| :---: | :---: | :---: | :---: |
| Chapter 3 Electronic <br> Structure and Periodic Properties of Elements: Section 3.7 Molecular and Ionic Compounds | The title of this section is "Molecular and Ionic Compounds", but the text describes ionic compounds first and then molecular. Consider changing the order in the title to better reflect the contents. | Revise the title to "lonic and Molecular Compounds". | General/pedago gical suggestion or question |
| Chapter 4 Chemical <br> Bonding and <br> Molecular <br> Geometry: Section <br> 4.1 Ionic Bonding | There is a small typo in the section on Electronic Structures of Cations. When discussing the formation of Fe2+ it should say "....by the loss of the 4 s electrons..." The word electron should be plural. | Revise as indicated. | Typo |
| Chapter 4 Chemical <br> Bonding and <br> Molecular <br> Geometry: Section <br> 4.2 Covalent <br> Bonding | Figure 4.4 provides the $\mathrm{H}-\mathrm{H}$ bond length as 0.74 which is the Internuclear Distance in angstrom but the Internuclear Distance is labeled as picometers (pm). | This figure will be updated. | Typo |
| Chapter 4 Chemical <br> Bonding and <br> Molecular <br> Geometry: Section <br> 4.3 Chemical <br> Nomenclature | In Problem 31, part b), revise titanium dioxide to titanium(IV) oxide | In exercise 31, revise part b "titanium dioxide" to "titanium(IV) oxide". | Typo |
| Chapter 4 Chemical <br> Bonding and <br> Molecular <br> Geometry: Section <br> 4.3 Chemical <br> Nomenclature | "Naming lonic Compounds" -- question (d) lists MgSO4.7H2O, but the solution discusses Cl and Cr , neither of which are in (d) | Delete "Cl-" and revise "Cr^3+" to "Mg^2+". | Incorrect answer, calculation, or solution |
| Chapter 4 Chemical <br> Bonding and <br> Molecular <br> Geometry: Section <br> 4.4 Lewis Symbols <br> and Structures | Ch. 4 End of Chapter Questions \#37 E needs to specify if the solution is acidic or basic. | Add the following to the end of the answer: "(in basic solution)". | Other factual inaccuracy in content |
| Chapter 4 Chemical <br> Bonding and <br> Molecular <br> Geometry: Section <br> 4.4 Lewis Symbols <br> and Structures | \#43 a) phosphate ion needs one $\mathrm{P}=\mathrm{O}$ bond. <br> c) sulfite ion needs one $\mathrm{S}=\mathrm{O}$ double bond | The solution will be updated. | Incorrect answer, calculation, or solution |
| Chapter 4 Chemical <br> Bonding and <br> Molecular <br> Geometry: Section 4.5 Formal Charge and Resonance | \#69 in SOO structure, the formal charge of the Oxygen on the far right should $=0$ | The solution will be updated. | Incorrect answer, calculation, or solution |


| Chapter 4 Chemical <br> Bonding and <br> Molecular <br> Geometry: Section <br> 4.5 Formal Charge <br> and Resonance | shortlink http://openstax.org///16LewisMake needs new target, as http://firstyear.chem.usyd.edu.au/iChem/lewis. shtml times out | This link will be updated. | Broken link |
| :---: | :---: | :---: | :---: |
| Chapter 4 Chemical <br> Bonding and <br> Molecular <br> Geometry: Section 4.5 Formal Charge and Resonance | Example 4.10 The Lewis structure that shows a nitrogen-nitrogen triple bond and a nitrogenoxygen single bond (nitrogen in the center) should also be shown and compared to the others. This one, which puts the negative formal charge on the oxygen, is the most stable (point \#4 in the preceding text). | This example will be updated. | General/pedago gical suggestion or question |
| Chapter 4 Chemical <br> Bonding and Molecular Geometry: Section 4.5 Formal Charge and Resonance | When the resonance structures of nitrite are shown, they are properly shown with one $\mathrm{N}=\mathrm{O}$ and one N-O. However, my question pertains to the blended structure. If the bond lengths are the "same," shouldn't the number of lone pairs on each $O$ be the same? | The section on resonance will be updated. | Other factual inaccuracy in content |
| Chapter 4 Chemical <br> Bonding and Molecular Geometry: Section 4.6 Molecular Structure and Polarity | \#96 b) need a lone pair of electrons on the N. c) REMOVE the lone pair of electrons on the S. d) should be trigonal pyramid, not tetrahedral; Also, one S-O single bond should be a $\mathrm{S}=\mathrm{O}$ double bond | The solution will be updated. | Incorrect answer, calculation, or solution |
| Chapter 4 Chemical <br> Bonding and Molecular Geometry: Section 4.6 Molecular Structure and Polarity | \#86 b) molecular shape should be tetrahedral | The solution will be updated. -WW SE | Incorrect answer, calculation, or solution |
| Chapter 4 Chemical <br> Bonding and Molecular Geometry: Section 4.6 Molecular Structure and Polarity | \#84 b) molecular shape should be see-saw. c) $\mathrm{SnCl} 3+$ needs to be a positive ion with a +1 charge to have the properties listed. e) molecular shape should be T-shape | In part C), revise "SnCl_3^-" to "SnCl_3^+". The rest of the solution will also be updated. | Incorrect answer, calculation, or solution |
| Chapter 4 Chemical <br> Bonding and <br> Molecular <br> Geometry: Section <br> 4.6 Molecular <br> Structure and <br> Polarity | Figure 4.16 and 4.19 The left column is labeled "Number of electron pairs", but actually describes the "Number of Electron Groups". It is possible, and often occurs that an electron group consists of two or more electron pairs. | Figures 7.16 and 7.19 will be updated to use "region" instead of "pair". | Other factual inaccuracy in content |
| Chapter 4: Key Equations | in the equation for formal charge instead of "\# one pair electrons" it should be "\# lone pair electrons" | Revise "one" to "lone". | Typo |
| Chapter 5 Advanced <br> Theories of Bonding: Section 5.1 | Example 5.2 In the worked-out solution for HCl , the charge on an electron has an exponent of +19 (should be -19). | In the solution to Example 5.2 Finding the Partial Ionic Character, | Typo |


| Valence Bond Theory |  | $\begin{aligned} & \text { revise "(1.60218×10^19 C)" to } \\ & \text { "(1.60218 } \left.\times 10^{\wedge}-19 C\right) \text { ". } \end{aligned}$ |  |
| :---: | :---: | :---: | :---: |
| Chapter 5 Advanced <br> Theories of <br> Bonding: Section 5.1 <br> Valence Bond <br> Theory | Example 5.2 "To get the partial ionic character, we divide This means the bond is about $17 \%$ ionic" The calculation to get partial ionic character is incomplete | In the solution after "we divide," add the following: the experimentally measured bond moment by this limiting value: <br> \%ionic character $=\mu \_\exp / \mu \_\lim \times$ $100 \%=(1.03 \mathrm{D}) /(6.09 \mathrm{D}) \times 100 \%=$ 17\% | Typo |
| Chapter 5 Advanced Theories of Bonding: Section 5.2 Hybrid Atomic Orbitals | Figure 5.10 The intermediate structure (to the right of the "hybridization" arrow and above the "gives a trigonal planar arrangement" arrow) looks to me like those sp2 hybrid orbitals are oriented along the $+x,-x$ and -y axes, at 90 degrees to each other. Then somehow they go to 120 degrees in the final structure. Compare this to the analogous figure 8.15 , where it is clear in the intermediate structure that the hybridized orbitals are NOT aligned with the axes. | This figure will be updated. | Other factual inaccuracy in content |
| Chapter 5 Advanced Theories of Bonding: Section 5.4 Molecular Orbital Theory | In the example box, charge is given as $10^{\wedge} 19 \mathrm{C}$ but should be 10^-19C | Our reviewers accepted this change. | Typo |
| Chapter 5 Advanced <br> Theories of <br> Bonding: Section 5.4 <br> Molecular Orbital <br> Theory | The explanation of how to get to the solution is cut off mid-sentence: "To get the partial ionic character, we divide This means...". It should probably say "by the mass of the molecule" or something similar. | Our reviewers accepted this change. | Other |
| Chapter 5 Advanced <br> Theories of <br> Bonding: Section 5.4 <br> Molecular Orbital <br> Theory | Link to Orbitron website is broken. Message says "account suspended." Is there an updated link? Or even a new source? | This link will be updated. | Broken link |
| Chapter 5 Advanced <br> Theories of <br> Bonding: Section 5.4 <br> Molecular Orbital <br> Theory | The description for Figure 5.38 in the hardback OpenStax 2016 and the e-book on-line (where it is Chapter 8 Figure 12) both have a typo that is confusing my students. The description states, "Without mixing, the MO pattern occurs as expected, with the op (sorry that the " $P$ " is not a subscript - I created the sigma symbol with a "P" subscript in Word but after copying and pasting into your errata description box, your box removed the subscript function)orbital energy lower in energy than the $\sigma p$ orbitals...." The only way this sentence can make sense is for the second $\sigma p$ to be $\pi p$ so that the description matches the figure." Please make this correction because in its current form it makes no sense and the students are confused. Thanks | Revise the caption as indicated. | Typo |


| Chapter 5 Advanced Theories of Bonding: Section 5.4 Molecular Orbital Theory | Figure 5.29 The two s orbitals that are being combined have different phases. When they are added, it should result in an antibonding orbital, and when they are subtracted it should result in a bonding orbital. The figure is inaccurate in its current form. The easy solution is to give both s orbitals the same phase. | This figure will be updated. | Other factual inaccuracy in content |
| :---: | :---: | :---: | :---: |
| Chapter 5 Advanced <br> Theories of <br> Bonding: Section 5.4 <br> Molecular Orbital <br> Theory | Figure 5.29 contradicts what you discuss in the section on sigma bonding with p-orbitals as well as standard MO representations. As you say, "We indicate the phases by shading the orbital lobes different colors." In figure 8.29, you show s orbitals with two different phases interacting by representing the s-orbitals with different colors. Both should be the same color and the sigma(bonding) should be shaded with a single color. To help students understand this representation, you would need to mention the difference in colors earlier in the text. <br> There is an additional typo; you say that "The plus (+) signs indicate the locations of nuclei." The image has dots, not + . | In the caption, revise "The plus (+) signs" to "The dots (•)". The figure colors were addressed in a previous report and are correct in webview. | Other factual inaccuracy in content |
| Chapter 5 Advanced <br> Theories of <br> Bonding: Section 5.4 <br> Molecular Orbital <br> Theory | shortlink openstax.org/l/16labelorbital needs new target, as http://firstyear.chem.usyd.edu.au/calculators/ mo_diagrams.shtml times out | Delete this link to learning feature box. | Broken link |
| Chapter 5 Advanced <br> Theories of Bonding: Section 5.4 Molecular Orbital Theory | According to the New York Times article "Walter Kohn, Who Won Nobel in Chemistry, Dies at 93", Walter Kohn died in 2016, so the sentence "Although he is now an emeritus professor, he is still actively working on projects involving global warming and renewable energy" cannot be correct. | Revise the sentence beginning "Although he is now..." to "Dr. Kohn passed away in 2016 at the age of 93." | Other factual inaccuracy in content |
| Chapter 5 Advanced <br> Theories of Bonding: Section 5.4 Molecular Orbital Theory | The image in figure 8 shows the MO diagram for Be2 + ion having the 2 p MO's in the sigma, pi, pi filling (energy) order, but later in the chapter (image 11) the filling (energy) order is pi, pi, sigma. The image could be fixed by changing the bonding order or by changing the molecule to Ne 2 . | This figure will be updated. | Other factual inaccuracy in content |
| Chapter 6 Composition of Substances and Solutions: Section 6.1 Formula Mass and the Mole Concept | In exercise \#43(a), the molar mass of 58 is given as $256.528 \mathrm{~g} / \mathrm{mol}$. This is correct when the molar of mass of $S$ is taken to the 3rd decimal place. However, the periodic table provided in Appendix A provides the atomic/molar mass of S to only 2 decimal places. This inconsistency in presentation may cause confusion among students. | Revise " $256.528 \mathrm{~g} / \mathrm{mol}$ " to " 256.48 g/mol". | Other |
| Chapter 6 Composition of Substances and Solutions: Section | Examples 2.7, 2.8, 2.9, 2.10, 2.11, 2.12 cancel units but also should cancel unit (say g ) and substance (say CO). some have unit (g) where it is better to show unit and substance ( g CO ) | This will be updated throughout the chapter. | General/pedago gical suggestion or question |


| 6.1 Formula Mass <br> and the Mole <br> Concept |  |  |  |
| :--- | :--- | :--- | :--- |
| Chapter 6 <br> Composition of <br> Substances and <br> Solutions: Section <br> 6.2 Determining <br> Empirical and <br> Molecular Formulas | Problem 8 The phrase: '(...) photographic <br> "hypo" ' is perhaps not the best - or at the very <br> least requires a few additional words of <br> explanation. I developed film in the day - and I <br> didn't readily pick up on what this <br> meant. Many of my students think that it is a <br> 'typo'. Maybe: 'the percent composition of <br> sodium thiosulfate' | In part b of exercise 8, revise as <br> follows: <br> 8. Calculate the following to four <br> significant figures: <br> (b) the percent composition of <br> photographic fixer solution ("hypo"), <br> Na2S2O3 | General/pedago <br> gical suggestion <br> or question |
| Chapter 6 <br> Composition of <br> Substances and <br> Solutions: Section <br> 6.2 Determining <br> Empirical and <br> Molecular Formulas | Change "Hydrogen" to "Oxygen". | Revise as indicated. |  |
| Chapter 6 <br> Composition of <br> Substances and <br> Solutions: Section <br> 6.2 Determining <br> Empirical and <br> Molecular Formulas | In the sentence that leads into the first worked <br> problem, it states, "Consider a sample of <br> compound determined to contain 1.71g <br> C....." However, in the worked problem the <br> mass used is not 1.71g C but the mass 1.17g C is <br> used. Please correct this error. I have already <br> had my students complaining about this. The <br> error is in the current version on-line. Thanks <br> for your help, Dr. Palamara | Change "1.17 g" to "1.71 g" in the <br> first equation. | Typo |


| Chapter 6 <br> Composition of Substances and Solutions: Section 6.2 Determining Empirical and Molecular Formulas | Problem 10asks to find the percent composition of ammonia of a certain chemical formula containing no calcium; yet, I find that in the detailed solution guide that calcium's atomic mass is used in the calculations to find the answer. I believe the answer is 69.2\% rather than $38.2 \%$. Additionally, the provided answer is unreasonable based on the given information in the problem. Please go over this problem in case my answer is incorrect as I cannot check that due to the error in the key. | The solution manual will be updated. | Incorrect answer, calculation, or solution |
| :---: | :---: | :---: | :---: |
| Chapter 6 <br> Composition of Substances and Solutions: Section 6.2 Determining Empirical and Molecular Formulas | Right after "Consider as another example a sample of compound determined to contain 5.31 g Cl and 8.40 g O . Following the same approach yields a tentative empirical formula of:" <br> In line 7 of this code you will see C1 but it should be Cl. | Revise "C1" to "CI" (letter I). | Typo |
| Chapter 6 Composition of Substances and Solutions: Section 6.3 Molarity | Exercise \#26 when calculating this, it does not match the answer key. the problem should be $10^{\wedge}-5$ or the answer key as 10^-7 | In part (b), revise " $10^{\wedge}-6$ " to "10^-7" and "10^-4" to "10^-5". | Incorrect answer, calculation, or solution |
| Chapter 6 Composition of Substances and Solutions: Section 6.3 Molarity | Problem \#47 - the molar mass should be 74.09 instead of 79.09. Problem \#51 - should read b) should be $3.8 \times 10^{\wedge}-6$, not $10^{\wedge}-5$. | Revise problem 51 question stem part (b) to $3.8 \times 10^{\wedge}-6$. The problem 47 solution will be updated in the solution manual. | Incorrect answer, calculation, or solution |
| Chapter 6 Composition of Substances and Solutions: Section 6.3 Molarity | Example 6.9 references "sweetened tea" from example 6.8, but there is now previous mention of sweetened tea. | This issue has been fixed previously. | Other |
| Chapter 7 <br> Stoichiometry of Chemical Reactions: <br> Section 7.2 <br> Classifying Chemical <br> Reactions | It seems like there's an error because in the instruction it's said ClO4-, but in the answer it's Cl03- <br> The correct answer for the problem is therefore: $4 \mathrm{Cl} 2+8 \mathrm{OH}-=>7 \mathrm{Cl}-+\mathrm{ClO} 4-+4 \mathrm{H} 2 \mathrm{O}$ | Right before the "HINT" revise "ClO4-" to "ClO3-". The answer is then correct. | Incorrect answer, calculation, or solution |
| Chapter 7 <br> Stoichiometry of Chemical Reactions: <br> Section 7.2 <br> Classifying Chemical <br> Reactions | When demonstrating on how to use the solubility guidelines (Table 4.1), the textbook gives an example of mixing solutions of silver nitrate and sodium fluoride. One of the products in this chemical reaction is silver fluoride, AgF. According to the textbook, "The solubility guidelines indicate... AgF is one of the exceptions to the general solubility of fluoride salts." However, in Table 4.1, the only exceptions to the solubility rule written for fluoride are "compounds with group 2 metal cations, $\mathrm{Pb}(2+$ charge $)$ and $\mathrm{Fe}(3+$ charge)." | Revise the line of Table 4.1 beginning "compounds with group 2 " to "compounds with group 2 metal cations, $\mathrm{Pb}^{\wedge} 2+, \mathrm{Fe}^{\wedge} 3+$, and $\mathrm{Ag}^{\wedge}+$ ". | Other factual inaccuracy in content |


| Chapter 7 <br> Stoichiometry of Chemical Reactions: <br> Section 7.2 <br> Classifying Chemical <br> Reactions | Table 7.1 has been updated to list all of the group I cations as soluble, but ammonium ( $\mathrm{NH} 4+$ ), which is also always soluble, has been omitted. Please modify the table to contain all group I ions plus ammonium in the 'always soluble' section. | Revise to add "NH4+" before "group I cations:". | Other factual inaccuracy in content |
| :---: | :---: | :---: | :---: |
| Chapter 7 <br> Stoichiometry of Chemical Reactions: <br> Section 7.2 <br> Classifying Chemical Reactions | Example 7.5 Part (a) should be referring to guideline 3 for the oxidation number of H , not guideline 1 | Revise "guideline 1" to "guideline 3". | Typo |
| Chapter 7 <br> Stoichiometry of Chemical Reactions: <br> Section 7.2 <br> Classifying Chemical <br> Reactions | \#20 c) C is oxidized from -2 to +4 | The solution will be updated. | Incorrect answer, calculation, or solution |
| Chapter 7 <br> Stoichiometry of Chemical Reactions: <br> Section 7.2 <br> Classifying Chemical <br> Reactions | AgF is soluble in water, and should not form a precipitate. The text says that it would form a precipitate. This does not agree with the information provided in the solubility table on page 183. | In Table 4.1, revise $\mathrm{Pbb}^{\wedge} 2+, \mathrm{Fe}^{\wedge} 3+$, and $\mathrm{Ag}^{\wedge}+$ " to " $\mathrm{Pb}^{\wedge} 2+$ and $\mathrm{Fe}^{\wedge} 3+$ ". Also revise "fluoride" to "chloride" and "F" to "Cl" in the text and equation below the table. Replace "one of the exceptions to the general solubility of fluoride salts" with "insoluble". | Incorrect answer, calculation, or solution |
| Chapter 7 <br> Stoichiometry of Chemical Reactions: <br> Section 7.2 <br> Classifying Chemical <br> Reactions | Chapter 4 Question 34 stated $\mathrm{SO} 3+\mathrm{H} 2 \mathrm{O}$ produces "liquid hydrogen sulfate" as the only product, "liquid hydrogen sulfate" should be "sulfuric acid". | Revise "hydrogen sulfate" to "dihydrogen sulfate". | Typo |
| Chapter 7 <br> Stoichiometry of Chemical Reactions: Section 7.3 Reaction Stoichiometry | Answer key at the back of the book for Chapter 4, question 45 (d). The correct answer should be 769 mol H 2 O and 13.8 kg of H 2 O . <br> This question requires students to answer question 44 first for this chapter. In the instructors solution manual, the molar mass of C 2 H 2 is written as $28.054 \mathrm{~g} / \mathrm{mol}$. Instead, the molar mass should be $26 \mathrm{~g} / \mathrm{mol}$. Hence the wrong numbers in the key. | Revise "713 mol" to "768 mol" and "12.8 kg" to "13.8 kg". | Incorrect answer, calculation, or solution |
| Chapter 7 <br> Stoichiometry of <br> Chemical Reactions: <br> Section 7.5 <br> Quantitative <br> Chemical Analysis | \#94 Formula for KHP is KHC8H4O4 and product is $\mathrm{KNaC8H} 4 \mathrm{O} 4$. Incorrect formulas written are KHC8H5O4 and KNaC8H5O4. | Revise "H_5" to "H_4". | Incorrect answer, calculation, or solution |
| Chapter 7 <br> Stoichiometry of <br> Chemical Reactions: <br> Section 7.5 | Example 7.15 The problem involves MgSO 4 , but the associated flow chart indicates CaSO 4 . | The image in the solution will be updated. | Typo |


| Quantitative Chemical Analysis |  |  |  |
| :---: | :---: | :---: | :---: |
| Chapter 7 <br> Stoichiometry of Chemical Reactions: <br> Section 7.5 <br> Quantitative <br> Chemical Analysis | Exercises \#89 in the balanced equation the product H 2 PO 4 should be H 3 PO 4 . | This will be updated in the solution manual. |  |
| Chapter 7 <br> Stoichiometry of Chemical Reactions: <br> Section 7.5 <br> Quantitative <br> Chemical Analysis | Example 7.14, Check Your Learning Question The question states the reaction containing potassium permanganate, KMnO , however the molecular equation for the reaction in the question has only permanganate. | Add the following after "KMnO_4": "(see net ionic equation below)". | Other factual inaccuracy in content |
| Chapter 7 <br> Stoichiometry of Chemical Reactions: <br> Section 7.5 <br> Quantitative <br> Chemical Analysis | The answer for the question was actually for another similar question with slight tweaks on the figures. The 3 figures in Question 79 are 18, 72, 36 The answer in the student manual was meant for $24,96,48$. | Revise the answer to "XeF_4". The full solution will also be updated in the solution manual. | Incorrect answer, calculation, or solution |
| Chapter 8 Gases: <br> Section 8.1 Valence <br> Bond Theory | In an example demonstrating pressure as $\mathrm{P}=$ F/A, the text determines that a figure skater is more likely to fall through ice than an elephant. The text states the pressure of a figure skater on one skate is higher than the pressure of an elephant standing on its four feet. While this is true, it misleads the reader since it implies that average pressure is the only consideration to take to determine if an object will break on ice. In the case of average sheer stress (assuming that the ice has a maximum allowable shear stress of ice is 600 kPa or $87 \mathrm{lb} / \mathrm{in}^{\wedge} 2$ ), a 14,000 lb elephant would need ${ }^{\sim} 160$ in^2 of cross sectional ice for support while the 120 lb skater would need $\sim 1.4 \mathrm{in}^{\wedge} 2$ of cross sectional ice for support. Assuming that the circumference of an elephant's foot is $\sim 48$ inches, it would take ~192 inches to enclose all four feet in perimeter and require the ice to be about 160 in^2/192 in $=0.83$ inches thick. Assuming that a skate blade is ~12 inches long, it would take $\sim 24$ inches to enclose one skate in perimeter and require the ice to be about 1.4 in^2/24in $=0.058$ inches thick. Given these calculations, it is more likely that the elephant will break the ice by average shear stress than the ice skater. Regardless of the relevance of average shear stress to chemistry, there should be a better and more direct example/application of pressure than falling through ice, such that it takes less force to puncture a balloon with a sharp object than a blunt object. | Revise the sentence "Let's apply this concept to determine..." to "Let’s apply this concept to determine which exerts a greater pressure in Figure 9.3-the elephant or the figure skater?" <br> Also revise the sentence "Even though the elephant is more than..." to " Even though the elephant is more than one hundred-times heavier than the skater, it exerts less than one-half of the pressure." | Other factual inaccuracy in content |
| Chapter 8 Gases: <br> Section 8.1 Valence <br> Bond Theory | In the worked-out solution for HCl , the charge on an electron has an exponent of +19 (should be -19). | In the solution to Example 5.2 Finding the Partial Ionic Character, | Typo |


|  |  | $\begin{aligned} & \text { revise "(1.60218×10^19 C)" to } \\ & \text { "(1.60218 } \left.\times 10^{\wedge}-19 C\right) \text { ". } \end{aligned}$ |  |
| :---: | :---: | :---: | :---: |
| Chapter 8 Gases: <br> Section 8.2 Relating <br> Pressure, Volume, <br> Amount, and <br> Temperature: The Ideal Gas Law | question 45 describes a balloon going over Mount Crumpet in British Colombia. The correct spelling for this location is Mount Crumpit | Our reviewers accepted this change. | Typo |
| Chapter 8 Gases: <br> Section 8.2 Relating <br> Pressure, Volume, <br> Amount, and <br> Temperature: The Ideal Gas Law | \#45: The number 278.24 is listed with units of atm. It should be 278.24 K . | The solution manual will be updated. | Incorrect answer, calculation, or solution |
| Chapter 8 Gases: <br> Section 8.2 Relating <br> Pressure, Volume, <br> Amount, and <br> Temperature: The Ideal Gas Law | \#29: The answer lists the temperature as -193. It should be -196 as stated in the problem | The solution manual will be updated. | Incorrect answer, calculation, or solution |
| Chapter 8 Gases: <br> Section 8.2 Relating Pressure, Volume, <br> Amount, and <br> Temperature: The Ideal Gas Law | \#28 wrong answer entirely. Answer should solve for Temperature | The solution will be updated. | Incorrect answer, calculation, or solution |
| Chapter 8 Gases: <br> Section 8.3 <br> Stoichiometry of Gaseous Substances, Mixtures, and Reactions | Figure 8.22 Y -axis (vapor pressure) is missing its unit. If this graph is to match the data in Table 9.2, the unit should be torr. | This figure will be updated. | Typo |
| Chapter 8 Gases: <br> Section 8.3 <br> Stoichiometry of Gaseous Substances, Mixtures, and Reactions | Figure 8.20 The middle cylinder (red/purple) has $\mathrm{P}=600 \mathrm{kPa}$ and the right-hand cylinder (yellow) has P $=450 \mathrm{kPa}$. These should be switched, as it appears that there are more gas particles in the yellow cylinder. Also, the caption refers to gas A, gas B and gas C, but these letters do not appear in the figure. | This figure will be updated. Also update the caption to "If equalvolume cylinders containing gasses at pressures of $300 \mathrm{kPa}, 450 \mathrm{kPa}$, and 600 kPa are all combined in the same-size cylinder, the total pressure of the gas mixture is 1350 kPa." | Other factual inaccuracy in content |
| Chapter 8 Gases: <br> Section 8.3 <br> Stoichiometry of Gaseous Substances, Mixtures, and Reactions | Ch 8, ex69, temperature should be in Kelvin not unspecified degreesTemp is given as "875" which implies Celsius but the answer is only valid if 875 Kelvin. | Revise "875" to "875 K". | Typo |
| Chapter 8 Gases: <br> Section 8.3 <br> Stoichiometry of Gaseous Substances, | https://openstax.org///16SusanSolomon redirect is broken. Needs new link. | This link will be updated. | Broken link |


| Mixtures, and Reactions |  |  |  |
| :---: | :---: | :---: | :---: |
| Chapter 8 Gases: <br> Section 8.3 <br> Stoichiometry of Gaseous <br> Substances, <br> Mixtures, and <br> Reactions | \#66 it would be preferable to use the stoichiometric ratio $4 / 4$ rather than $1 / 1$ so students know where the numbers come from | The solution will be updated. | Incorrect answer, calculation, or solution |
| Chapter 8 Gases: <br> Section 8.3 <br> Stoichiometry of <br> Gaseous <br> Substances, <br> Mixtures, and <br> Reactions | it incorrectly says "and the definition of molarity" when it should say "and the definition of molar mass". | Revise "definition of molarity" to "definition of molar mass". | Typo |
| Chapter 8 Gases: <br> Section 8.3 <br> Stoichiometry of <br> Gaseous <br> Substances, <br> Mixtures, and <br> Reactions | broken link http://openstax.org///16GlobalWarming | This link will be updated. | Broken link |
| Chapter 8 Gases: <br> Section 8.3 <br> Stoichiometry of Gaseous Substances, Mixtures, and Reactions | Example 8.15 Question asks for N2O mole fraction and partial pressure but the solution is written as N2. | $\begin{aligned} & \text { Revise "N_2" to "N_2O" and "143.6" } \\ & \text { to "144". } \end{aligned}$ | Other factual inaccuracy in content |
| Chapter 8 Gases: <br> Section 8.5 The <br> Kinetic-Molecular <br> Theory | This chapter makes many references to molecular velocities, but all reasoning relies upon using the Maxwell-Boltzman speed distribution. It looks like a previous errata (ID 6109) addressed this in several figures, but the chapter text and figure captions still refer to velocities. The online version of the text currently uses the word "velocity" or "velocities" 11 times. In all cases, replacing with the words "speed" or "speeds" would be appropriate and improve the accuracy of the arguments. | Replace "velocity" and "velocities" with "speed" and "speeds" throughout the chapter. | Other factual inaccuracy in content |
| Chapter 8 Gases: <br> Section 8.5 The <br> Kinetic-Molecular <br> Theory | The label for the scenario of decreased volume says "Volume decreased Wall area decreased = Pressure increased". The pressure increases whenever the volume is decreased even if the wall area remains constant (e.g. half sphere capping a cylinder, invert and push the half sphere inside the cylinder: same wall area, lower volume -> pressure increased). On the other hand, if you increase or decrease the wall area while keeping the volume constant (for example by going from "zig-zag" wall to straight wall), the pressure will not change. | Revise Boyle's law bullet point to "Boyle's law. If the volume of a given amount of gas at a given temperature is decreased (that is, if the gas is compressed), the molecules will be exposed to a decreased container wall area. Collisions with the container wall will therefore occur more frequently and the pressure exerted by the gas will increase (Figure 9.31)." | Other factual inaccuracy in content |


|  | In short, the wall area does not influence the pressure. |  |  |
| :---: | :---: | :---: | :---: |
| Chapter 9 <br> Thermochemistry: Section 9.1 Energy Basics | Example 9.1 In the "check your learning" question in example 9.1 (found on p 240 of pdf), the letter $x$ is used instead of the multiplication symbol. | Revise Example 9.1 Measuring Heat as follows: <br> Ensure multiplication symbol is used throughout, not the letter x . | Typo |
| Chapter 9 <br> Thermochemistry: Section 9.1 Energy Basics | Example 9.1 - "Check your learning" - The question reads - How much heat, in joules, must be added to a $5.07 \times 10^{\wedge} 4 \mathrm{~J}$ iron skillet.... The answer is $5.07 \times 10^{\wedge} 4 \mathrm{~J}$. According to the calculation, the question should read "How much heat, in Joules, must be added to a 502 g iron skillet...." | Revise " $5.07 \times 10^{\wedge} 4 \mathrm{~J}$ iron skillet" to "502 g iron skillet". | Incorrect answer, calculation, or solution |
| Chapter 9 <br> Thermochemistry: Section 9.1 Energy Basics | Check Your Learning: How much heat, in joules, must be added to a $5.07 \times 10^{\wedge} 4 \mathrm{~J}$ iron skillet to increase its temperature from $25^{\circ} \mathrm{C}$ to $250^{\circ} \mathrm{C}$ ? The specific heat of iron is $0.449 \mathrm{~J} / \mathrm{g}^{\circ} \mathrm{C}$. <br> Solution: $5.07 \times 10^{\wedge} 4 \mathrm{~J}$ <br> The unit for $5.07 \times 10^{\wedge} 4$ should be measured in $g$, not J. the answer should be closer to $5.12 \times 10^{\wedge} 6$ if the number above is in g . | Revise the unit for $5.07 \times 10^{\wedge} 4$ to $g$ and the answer to $5.12 \times 10^{\wedge} 6 \mathrm{~J}$. | Incorrect answer, calculation, or solution |
| Chapter 9 <br> Thermochemistry: <br> Section 9.2 <br> Calorimetry | Check Your Learning of Exercise 9.7 The fuel burned it Benzene and not Glucose as the question asks. Replace Glucose by Benzene. | Revise "glucose" to "benzene". | Typo |
| Chapter 9 <br> Thermochemistry: <br> Section 9.2 <br> Calorimetry | Example 9.7 - Bomb Calorimetry - Check Your Learning Answer should be -39.0 kJ indicating it's an exothermic reaction. | Revise the answer to "q_rx = -39.0 kJ (the reaction produced 39.0 kJ of heat)". | Incorrect answer, calculation, or solution |
| Chapter 9 <br> Thermochemistry: <br> Section 9.2 <br> Calorimetry | Example 9.5-Check Your Learning The answer should be -1.34 X 10^3 J indicating it's an exothermic reaction. | Revise the answer to "1.3 kJ". | Incorrect answer, calculation, or solution |
| Chapter 9 <br> Thermochemistry: <br> Section 9.2 <br> Calorimetry | \#28 - states 6.9 degrees C, should use the symbol for degrees: $6.0^{\circ} \mathrm{C}$ | In exercise 28, revise "increased the temperature of the solution by 6.9 degrees" to "increased the temperature of the solution by 6.9 ${ }^{\circ} \mathrm{C}$ ". | Typo |
| Chapter 9 <br> Thermochemistry: Section 9.3 Enthalpy | Question 50 states that the heat capacity of the products is $4.19 \mathrm{~J} / \mathrm{g}$ C. This should be described as the specific heat instead. Solving the question while treating the number as heat capacity provides an unreasonable temp change of -346 C. Solving it while treating it as specific heat gives a much more reasonable 1.46 C temperature change. | Revise "heat capacity" to "specific heat". | Typo |
| Chapter 9 <br> Thermochemistry: Section 9.3 Enthalpy | Exercise 75 When I do the calculation using Appendix G, I get -66.4 kJ The solution in the back of the book is +67.1 kJ | Update the answer to "-66.4 kJ". | Incorrect answer, calculation, or solution |


| Chapter 9 Thermochemistry: Section 9.3 Enthalpy | Exercise 83 Ethylene should be C2H4 | Revise C_2H_2 to C_2H_4. | Typo |
| :---: | :---: | :---: | :---: |
| Chapter 9 Thermochemistry: Section 9.3 Enthalpy | Example 9.13 shouldn't $\mathrm{Cl} 2+1 / 2 \mathrm{Cl} 2$ be $3 / 2 \mathrm{Cl} 2$ ? | Revise 1/2 Cl_2 to 3/2 Cl_2. | Typo |
| Chapter 9 Thermochemistry: Section 9.3 Enthalpy | Example 9.10 The second line of " $692 \mathrm{~g} \mathrm{C8H18"}$ should be changed to " $6.07 \mathrm{~mol} \mathrm{C8H18"}$. | Revise "692g" to "6.07 mol". | Typo |
| Chapter 9 <br> Thermochemistry: Section 9.3 Enthalpy | Chapter 5: Exercise 85 part b states: <br> (b) Calculate the volume of air at $25^{\circ} \mathrm{C}$ and 1.00 atmosphere that is needed to completely combust 25.0 grams of propane. Assume that air is 21.0 percent O 2 by volume. (Hint: We will see how to do this calculation in a later chapter on gases-for now use the information that 1.00 L of air at $25^{\circ} \mathrm{C}$ and 1.00 atm contains 0.275 g of O 2 per liter.) <br> IT should state that " 1.00 L of oxygen at $25^{\circ} \mathrm{C}$ and 1.00 atm contains 0.275 g of 02. .") | Delete "per liter" from the end of part b. | Typo |
| Chapter 9 <br> Thermochemistry: Section 9.3 Enthalpy | Question \#85, answer (b) given in the book is for the volume of O2, but the question was asking the volume of AIR, so the answer should be 330Liters/0.21 = 1571 Liters | Revise the answer for part (b) to "1570 L air". | Incorrect answer, calculation, or solution |
| Chapter 9 <br> Thermochemistry: <br> Section 9.4 <br> Strengths of Ionic and Covalent Bonds | The calculation following table 7.4 is not correct, and the Hlattice listed does not match that show in figure 7.13. <br> Hlattice=(553.5+76.5+79.4+375.7+328.2) kJ/mol $=1413.3 \mathrm{~kJ} / \mathrm{mol}$ <br> Correction that matches what is listed in figure 7.13 <br> Hlattice $=(553.5+76.5+79.4+375.7-$ <br> $328.2) \mathrm{kJ} / \mathrm{mol}=756.9 \mathrm{~kJ} / \mathrm{mol}$ | Table 7.4 will be updated, as well as the Hlattice calculation. | Incorrect answer, calculation, or solution |
| Chapter 10 Liquids and Solids: Introduction | Sentences starting with "Aside from the symptoms..." is repeated twice | Our reviewers accepted this change, and it will be included in the next print cycle. | Typo |
| Chapter 10 Liquids and Solids: Introduction | The entire body is repeated twice, i.e. it starts out "Leprosy has been .." and then in the middle it comes back again: "... treatment for leprosy for decades.Leprosy has been ...". <br> Also there is a typo with "her" being written "hr" | corrected typo | Typo |
| Chapter 10 Liquids and Solids: Section 10.1 Intermolecular Forces | Figure 10.14 The base labeled as thymine is uracil (no -CH3 group). I'll upload a suggested replacement for this figure as several bonds are awkwardly drawn as well or would be happy to be contacted. | This figure will be updated. | Other factual inaccuracy in content |


| Chapter 10 Liquids and Solids: Section 10.2 Properties of Liquids | Problem \# 24 It looks like for the shown 3-d structure of acetone the carbon atom geometry is tetrahedral. It should be optimized or rearranged to be trigonal planar. | This figure will be updated. | Other factual inaccuracy in content |
| :---: | :---: | :---: | :---: |
| Chapter 10 Liquids and Solids: Section 10.3 Phase Transitions | Figure 10.22 In the series of pictures, the pressure gauge indicates an increase in pressure. However, the mercury level in the manometer does not change. This seems inconsistent. | The figure will be updated. | Other factual inaccuracy in content |
| Chapter 10 Liquids and Solids: Section 10.4 Phase Diagrams | example 10.12 the answer is for 50 k Pa not 500 k Pa | Revise "500" to "50". | Typo |
| Chapter 11 <br> Solutions and Colloids: Section 11.1 The Dissolution Process | The sentence "Ammonium nitrate ( NaNO 3 ) is one such example and is used to make instant cold packs for treating injuries like the one pictured in Figure 11.5" is an example of a misplaced modifier. Figure 11.5 doesn't illustrate an injury, it illustrates a cold pack. | Revise "instant cold packs for treating injuries like the one pictured in Figure 11.5." to "instant cold packs like the one pictured in Figure 11.5, which are used for treating injuries. | General/pedago gical suggestion or question |
| Chapter 11 <br> Solutions and Colloids: Section 11.3 Solubility | Example 11.2 requires students to used Figure 11.8. But this information isn't mentioned until the solution. The question needs to refer students to this graph so that they don't have to read the solution to find out. Not that this change would mean the solution to the problem would no longer need to mention Figure 11.8. | Add the following to the end of the question: "Hint: Use the data in Figure 11.8 to estimate a value for the Henry's law constant at this temperature." | General/pedago gical suggestion or question |
| Chapter 11 <br> Solutions and Colloids: Section 11.3 Solubility | Example 11.2 asks students to use values read from Figure 11.8 to solve the problem. The figure is currently only marked at 1.0 and 2.0. Additional horizontal lines (perhaps in a lighter grey, but not labeled on the $y$-axis. "minor horizontal gridlines") would help students estimate the values, especially at the precision referred to in the given solution. Gridlines at least every 0.2 , and perhaps every 0.1 , would allow all students to make the estimate near 1.15. | Revise the sentence "First, estimate the Henry's law..." to "First, estimate the Henry's law constant for oxygen in water at the specified temperature of $30.0^{\circ} \mathrm{C}$ (Figure 11.8 indicates the solubility at this temperature is approximately $\sim 1.2$ mol/L)." | General/pedago gical suggestion or question |
| Chapter 11 <br> Solutions and Colloids: Section 11.3 Solubility | In the first step of the solution to Example 11.2, it says "First, compute the Henry's law constant for oxygen in water at the specified temperature of $30.0^{\circ} \mathrm{C}$ (see Figure 11.19)." <br> 1.) Just from the perspective of clarity, nothing in the reading to his point actually explains how to do this. <br> 2.) The referenced Figure 11.19, has nothing to do with the Henry's Law. It is a picture of a laboratory distillation unit. <br> 3.) There is a graph in Figure 11.8 (which would probably be the more appropriate figure to reference) that contains solubilities of gases in water. However, your Y-axis could probably stand to be presented differently if students are expected to use that. Going from a value of 1.0 | Revise "compute" to "estimate". Replace the three occurrences of " 1.15 " with " 1.2 ". Replace the two occurrences of 1.95 " with " 2.0 ". Replace " 6.2 " with " 6.4 ". | Other factual inaccuracy in content |


|  | $\times 10-3 \mathrm{~mol} / \mathrm{L}$ to $2.0 \times 10-3 \mathrm{~mol} / \mathrm{L}$ makes it very difficult come back with a figure as precise as $1.15 \times 10-3 \mathrm{~mol} / \mathrm{L}$, which is what was used in the solution. |  |  |
| :---: | :---: | :---: | :---: |
| Chapter 11 <br> Solutions and <br> Colloids: Section <br> 11.4 Colligative <br> Properties | Example 11.5 it is 1.04 kg soln and not 1.02 kg solution | Replace " 1.02 " with " 1.04 ", replace " 0.92 " with " 0.94 ", and replace " 0.60 " with " 0.59 ". | Typo |
| Chapter 11 <br> Solutions and Colloids: Section 11.4 Colligative Properties | Colligative Properties of Electrolytes section there are two equations after the first paragraph. These have the charges for the ions on the product side of each as subscripts. These should be superscripts. | Revise the indicated subscripts to superscripts. | Typo |
| Chapter 11 <br> Solutions and <br> Colloids: Section <br> 11.4 Colligative <br> Properties | Exercise 47b 0.0363 M and not 0.363 M | In the solution, "0.363" will be updated to "0.0363". | Typo |
| Chapter 11 <br> Solutions and Colloids: Section 11.4 Colligative Properties | In example 11.10 the 0.30 M solution of glucose in water became $0.03 \mathrm{~mol} / \mathrm{L}$ in the solution to the problem. | Revise "0.03" to "0.30". | Typo |
| Chapter 11 <br> Solutions and Colloids: Section 11.4 Colligative Properties | Problem 57. The Kb in the problem (2.43) and the $\mathrm{Kb}(2.34)$ in the answer are reversed. | Revise the value of K_b in the problem from "2.43" to "2.34". | Incorrect answer, calculation, or solution |
| Chapter 11 <br> Solutions and Colloids: Section 11.4 Colligative Properties | Near the end of Example 11.3 on page 619 in the PDF version the problem says "Finally, calculate molarity per its definition" when in this example we are calculating molality. The remainder of the problem correctly says molality, but there is an incorrect part that says we are looking for molarity. | Revise "calculate molarity" to "calculate molality". | Typo |
| Chapter 11 <br> Solutions and Colloids: Section 11.4 Colligative Properties | Chapter 11 End-Of-Chapter 48 The solution has the incorrect power of 10 and the incorrect sig figs. <br> The molar mass is given as 5700 (2sf), so the answer is limited to 2 sig figs. The correct answer is $4.3 \times 10^{\wedge} 03 \mathrm{~atm}$ (not -4 ). thanks SB | The solution will be updated. | Incorrect answer, calculation, or solution |
| Chapter 11 <br> Solutions and <br> Colloids: Section <br> 11.4 Colligative <br> Properties | Example 11.10. It should say $0.30 \mathrm{~mol} / \mathrm{L}$, but it instead says $0.03 \mathrm{~mol} / \mathrm{L}$. | Revise "0.03" to "0.30". | Typo |
| Chapter 12 <br> Thermodynamics: <br> Section 12.1 <br> Spontaneity | Chapter 12 problem 2. Ozone decomposes... 2 $\mathrm{O} 3(\mathrm{~g})-->3 \mathrm{O} 2(\mathrm{~g})$. The rate expression in the answer key is missing the $1 / 3$ for the O 2 portion. | The solution manual will be updated. | Incorrect answer, calculation, or solution |


| Chapter 12 <br> Thermodynamics: <br> Section 12.2 <br> Entropy | 17a While it is generally true that precipitation is an example of a loss of entropy, care needs to be taken with aqueous ions, whose presence decreases the entropy of water, as is often the case with insoluble ionic solids. If you look at the standard entropy values in Appendix G, you can see that $\mathrm{PbS}(\mathrm{s})$ has more entropy ( 91.2 $\mathrm{J} / \mathrm{mol} \mathrm{K})$ than the sum of $\mathrm{Pb} 2+(\mathrm{aq})(10.5 \mathrm{~J} / \mathrm{mol}$ $\mathrm{K})+\mathrm{S} 2-(\mathrm{aq})(22 \mathrm{~J} / \mathrm{mol} \mathrm{K})$. As I understand it this is not uncommon with insoluble ionic compounds | The equation in part (a) will be updated. | Other factual inaccuracy in content |
| :---: | :---: | :---: | :---: |
| Chapter 12 <br> Thermodynamics: <br> Section 12.2 <br> Entropy | In Check Your Learning, problem (d), the reactant is given as CaCO . It should be CaCO 3 . | Revise "CaCO" to "CaCO_3". | Typo |
| Chapter 12 <br> Thermodynamics: <br> Section 12.3 The <br> Second and Third <br> Laws of <br> Thermodynamics | For Ch. 12, Problem \#25. The questions should say values listed in Appendix G, calculate delta S degrees 298 for the following changes: | In exercise 25, add the delta symbol in front of "S degrees 298". | Typo |
| Chapter 12 <br> Thermodynamics: <br> Section 12.3 The <br> Second and Third <br> Laws of <br> Thermodynamics | Problem 12.28 asks the student to determine the change in entropy for several chemical reactions using appendix $G$. Problem 12.29 states the following: "Use the standard entropy data in Appendix G to determine the change in entropy for each of the reactions listed in Exercise 12.28. All the processes occur at the standard conditions and $25^{\circ} \mathrm{C}$." <br> The first issue is that problem 12.29, as it is currently written, is asking the student to re-do exactly what they did in problem 12.28 . The second issue is that if one compares their answers from problem 16.28 and the solution to problem 16.29 in the back of the book, they are different. They should not be since, as it currently stands, problems 12.28 and 12.29 are identical. <br> The answers in the answer key for problem 12.29 are ACTUALLY the correct answers to the chemical reactions found in problem 12.34. If one goes through and calculates the standard entropy (not free energy as requested in 12.34) for all the chemical reactions in 12.34, they would get exactly the answers found as the solution for problem 12.29. | In the question stem, update the exercise link to "Exercise 16.34". | Incorrect answer, calculation, or solution |
| Chapter 12 <br> Thermodynamics: <br> Section 12.3 The <br> Second and Third <br> Laws of <br> Thermodynamics | In the part of Example 16.6 labelled "Check Your Learning" instead of the answer being 24.7 $\mathrm{J} \mathrm{K}^{\wedge}-1 \mathrm{~mol}{ }^{\wedge}-1$ the answer should be 24.7 J/K (no moles) | Update the answer to "24.7 J/K". | Typo |


| Chapter 12 <br> Thermodynamics: <br> Section 12.4 Free Energy | Question 40 part c, there are no units on Kp. | Delete "atm" from the answer. | Other factual inaccuracy in content |
| :---: | :---: | :---: | :---: |
| Chapter 12 <br> Thermodynamics: <br> Section 12.4 Free Energy | Chapter 12 question 33 (f) has an answer of - 30 kJ spontaneous when the answer is actually 154.25 kJ spontaneous. I have done the problem about 30 times to see if I could even get near -30 but if your appendix $G$ values are correct there is no possibility of -30 kJ spontaneous being the correct answer. Hope someone can correct this as it was a homework problem for my chemistry class and it is incorrect. Thank you and all the best. | Revise the answer to " -154.3 kJ spontaneous". | Incorrect answer, calculation, or solution |
| Chapter 12 <br> Thermodynamics: <br> Section 12.4 Free Energy | The answer for number 42 part b is incorrect. The correct answer should be $98.9 \mathrm{~kJ} / \mathrm{mol}$. The textbook says the answer is $61.5 \mathrm{~kJ} / \mathrm{mol}$, but this is wrong because - <br> (8.31425)(975)ln(0.000005) $=98.9 \mathrm{~kJ} / \mathrm{mol}$. The person that solved this problem accidentally did (0.00250)^2/0.0125 for K, instead of $(0.000250)^{\wedge} 2 /(0.0125)$. This mistake is also in the student solutions manual. The student solutions manual says the molarity for Cl is 0.000250 M , but then proceeds to use the number 0.002550 in the calculations, leading to an incorrect answer. | Revise the answer to part (b) to " $98.9 \mathrm{~kJ} / \mathrm{mol}$ ". The solution will also be updated. | Incorrect answer, calculation, or solution |
| Chapter 12 <br> Thermodynamics: Section 12.4 Free Energy | On page 674 the answer for the 12.8 check your learning the answer is given negative and then nonspontaneous. Was that right because it is kind of confusing | In Example 12.8 Calculation of delta G 298, revise Check Your Learning answer as follows: <br> (a) $140.8 \mathrm{~kJ} / \mathrm{mol}$, nonspontaneous <br> (b) $141.5 \mathrm{~kJ} / \mathrm{mol}$, nonspontaneous | Typo |
| Chapter 12 <br> Thermodynamics: Section 12.4 Free Energy | In the student solution guide for Chapter 12 $(\bmod 4) \# 33 \mathrm{c}$ lists the free energy of $\mathrm{S}(\mathrm{g})$ as zero when it should actually be $238.25 \mathrm{~kJ} / \mathrm{mol}$. Therefore, the solution should be "[1mol(-53.6 $\mathrm{kJ} / \mathrm{mol})]$ - [1 mol $(0 \mathrm{~kJ} / \mathrm{mol})+1 \mathrm{~mol}(238.25$ $\mathrm{kJ} / \mathrm{mol})]=-291.9 \mathrm{~kJ} "$. <br> In the student solution guide Chapter 12 (mod 4) \#33f, the incorrect standard free energy value is used for carbon tetrachloride. The value used is for the liquid, rather than for the gas, which is used in the original problem. Therefore, the solution should read "[1mol($58.2 \mathrm{~kJ} / \mathrm{mol})+1 \mathrm{~mol}(-29.25 \mathrm{~kJ} / \mathrm{mol})]$ - <br> $[1 \mathrm{~mol}(66.8 \mathrm{~kJ} / \mathrm{mol})+3 \mathrm{~mol}(0 \mathrm{~kJ} / \mathrm{mol})]=-154.3$ kJ". | Revise the solution to part (c) of exercise 33 from "-53.6 kJ" to "$291.9 \mathrm{~kJ} "$. | Incorrect answer, calculation, or solution |
| Chapter 12 <br> Thermodynamics: Section 12.4 Free Energy | Ch. 12, Problem \#45. Should be " $77.1 \mathrm{~kJ} /$ mole". The "J" is missing in "kJ". | In exercise 45, revise " $k$ /mole" to "kJ/mole". | Typo |
| Chapter 12 Thermodynamics: | In the solution to Example 12.8 (Calculation of delta $G$ zero of yellow mercury (II) oxide), in the | Delete the $S$ that appears in the first equation in the solution. | Typo |


| Section 12.4 Free <br> Energy | part (a) of the solution, in the first equation, <br> there is a letter "S" that does not belong. |  |  |
| :--- | :--- | :--- | :--- |
| Chapter 12 <br> Thermodynamics: <br> Section 12.4 Free <br> Energy | In the student solution guide for chapter 12 <br> (mod 4) \#43b, the value substituted for $[\mathrm{Cl} 2]$ <br> should be 0.0124 M, rather than 0.0125 M <br> since $1 \%$ of the original quantity dissociates. <br> This changes the value of K to 0.000504, thus <br> making the value of delta G = 61,500 J/mol, or <br> 61.5 kJ/mol. | Revise the solution for exercise 43 <br> part b to "61.5 kJ/mol". | Incorrect <br> answer, <br> calculation, or |
| solution |  |  |  |


| Chapter 13 <br> Fundamental <br> Equilibrium <br> Concepts: Key <br> Equations | There are equations with delta-G listed in the key equations that aren't covered until later in the book, and do not appear in the PDF. | Delete the two equations beginning with " $\Delta \mathrm{G}$ ". | Typo |
| :---: | :---: | :---: | :---: |
| Chapter 13 <br> Fundamental <br> Equilibrium <br> Concepts: Section <br> 13.1 Chemical <br> Equilibria | in the rate of $\mathrm{f}(\mathrm{kf})$ is written as N -2O4 and it should be N2O4 | Revise this to N2O4. | Typo |
| Chapter 13 <br> Fundamental <br> Equilibrium <br> Concepts: Section <br> 13.1 Chemical <br> Equilibria | The chemical formula for dinitrogen tetroxide is represented by N -2O4 in two places. It should be N2O4. | Revise "N_-2" to "N_2". | Typo |
| Chapter 13 <br> Fundamental <br> Equilibrium <br> Concepts: Section <br> 13.2 Equilibrium <br> Constants | Exercise 29 The question asks students to write the Q expression for the ionization of NH 3 in water. The answer in the key is: $\mathrm{Q}=[\mathrm{NH} 4+][\mathrm{OH}-$ ]/[HN3]. The denominator is wrong and the expression should be: $\mathrm{Q}=[\mathrm{NH} 4+][\mathrm{OH}-] /[\mathrm{NH} 3]$ | Revise the denominator to [ NH 3$]$. | Typo |
| Chapter 13 <br> Fundamental <br> Equilibrium <br> Concepts: Section <br> 13.2 Equilibrium <br> Constants | In "example 13.3", and the subsection "Check your learning", in part (a), it says: <br> ...mol of $\mathrm{Cl} 2(\mathrm{~g}), \ldots$. The 2 should be in the subscript, not written like it is. | Revise "Cl2" to "Cl_2". | Typo |
| Chapter 13 <br> Fundamental <br> Equilibrium <br> Concepts: Section <br> 13.2 Equilibrium <br> Constants | The expression for $Q$ in terms of partial pressures lists the superscripts from the stoichiometry in the wrong place. The superscripts should be on each of the "P"s. Highlighted in the screenshot. | This expression will be updated. | Typo |
| Chapter 13 <br> Fundamental <br> Equilibrium <br> Concepts: Section <br> 13.2 Equilibrium <br> Constants | There are no learning objectives for chapter 13.2. | Add: "By the end of this section, you will be able to: <br> Derive reaction quotients from chemical equations representing homogeneous and heterogeneous reactions <br> Calculate values of reaction quotients and equilibrium constants, using concentrations and pressures Relate the magnitude of an equilibrium constant to properties of the chemical system" | General/pedago gical suggestion or question |
| Chapter 13 <br> Fundamental <br> Equilibrium <br> Concepts: Section <br> 13.3 Shifting | Exercise 49 The prompt states that alpha- and beta-analine of the same concentration will have different freezing points, implying that these stereo-isomers have different dissociation constants. This is factually incorrect in the absence of other chiral | This question will be replaced. | Other factual inaccuracy in content |


| Equilibria: Le Châtelier's Principle | compounds. (If this is correct, I would like to see the source data.) I suggest removing the question, or replacing the alanine with two different compounds that actually do have different freezing points at the same concentration. |  |  |
| :---: | :---: | :---: | :---: |
| Chapter 13 <br> Fundamental <br> Equilibrium <br> Concepts: Section <br> 13.3 Shifting <br> Equilibria: Le <br> Châtelier's Principle | 13.48 links to the EOC answer section and has a listed answer of (a). The question is an essay questions, not a multiple choice. The correct answer to 13.49 is (a), so either the link should be moved to 13.49 and the answer label changed to 13.49 , or the answer to 13.48 should be included:(Saturated solutions are an example of dynamic equilibrium. Dissociation and precipitation do not completely halt in a saturated solution; they just occur at the same rate, so individual $\mathrm{Ag}+$ ions (radioactive or not) can move back and forth between the solid and solution. | The answer for question 48 will be updated to "Though the solution is saturated, the dynamic nature of the solubility equilibrium means the opposing processes of solid dissolution and precipitation continue to occur (just at equal rates, meaning the dissolved ion concentrations and the amount of undissolved solid remain constant). The radioactive $\mathrm{Ag}^{\wedge}+$ ions detected in the solution phase come from dissolution of the added solid, and their presence is countered by precipitation of nonradioactive Ag^+." | Incorrect answer, calculation, or solution |
| Chapter 13 <br> Fundamental <br> Equilibrium <br> Concepts: Section <br> 13.3 Shifting <br> Equilibria: Le <br> Châtelier's Principle | Chapter 13 end-of-chapter problem 35 should have an equilibrium arrow rather than a "one way" arrow as it is an equilibrium question. | The chemical equation will be updated. | Typo |
| Chapter 13 <br> Fundamental <br> Equilibrium <br> Concepts: Section <br> 13.3 Shifting <br> Equilibria: Le <br> Châtelier's Principle | CHEMISTRY IN EVERYDAY LIFE Box carbon dioxidey | Replace "dioxidey" with "dioxide". | Typo |
| Chapter 13 <br> Fundamental <br> Equilibrium <br> Concepts: Section <br> 13.3 Shifting <br> Equilibria: Le <br> Châtelier's Principle | Where is the following link supposed to go? It doesn't link to a PHET simulation. Link to Learning direct links to http://openstax.org///16chatelier But it redirects to Learnerstv.com \| PPU Library library.ppu.edu | This Link to Learning box will be deleted. | Broken link |
| Chapter 13 <br> Fundamental <br> Equilibrium <br> Concepts: Section <br> 13.4 Equilibrium <br> Calculations | Solutions for exercise 90 (a and c) from Chapter 13 For $a$, the enthalpy value for the reaction should be positive and slightly larger according to the enthalpy value for F 2 O in the appendix. In the end, the value for K comes out to 1.1 x 10E-13. <br> For c , the enthalpy value comes to -89.3 and the entropy value should be -20.43 with the values in the appendix. In the end, the value for K comes out to $2.73 \times 10$ E4. | Revise the answers as follows: <br> (a) $K=1.07 \times 10^{\wedge}-13$ <br> (b) $K=2.42 \times 10^{\wedge}-3$ <br> (c) $K=2.73 \times 10^{\wedge} 4$ <br> (d) $K=0.229$ | Incorrect answer, calculation, or solution |


| Chapter 13 <br> Fundamental <br> Equilibrium <br> Concepts: Section <br> 13.4 Equilibrium <br> Calculations | Example 13.9, check your learning. Acetic acid plus ethanol to give the ester and water. The answer you provide for the 4 components gives a Kc of 0.24 but the given Kc is 4.0. The error is that you reversed the correct equilibrium amounts: 0.17 M concentrations are the two reactants and the 0.36 M concentrations are the 2 products of the reaction as written. As given, the reaction proceeds from right to left to react equilibrium. I suggest you switch the initial concentrations so that the reaction proceeds in the direction written. | Revise " 0.37 " to " 0.18 " and " 0.18 " to "0.37". |  |
| :---: | :---: | :---: | :---: |
| Chapter 13 <br> Fundamental <br> Equilibrium <br> Concepts: Section <br> 13.4 Equilibrium <br> Calculations | Exercise \#60 and \#61 It is unclear what the rows are below each reaction initial (I), change (C), or equilibrium (E). This should be added to clarify. | Revise the question stem to "Complete the partial ICE tables below." Each table for each part will be revised to show a column on the left with "change" and "equilibrium" at the beginning or rows 2 and 3 . | General/pedago gical suggestion or question |
| Chapter 13 <br> Fundamental <br> Equilibrium <br> Concepts: Section <br> 13.4 Equilibrium <br> Calculations | It seems really flawed to omit the initial row of an ICE table on principle (they're not called CE tables), and especially so given that equilibrium concentrations are found by adding change to initial. Presumably the answers shown in the answer key were derived by assuming all initial concentrations/pressures were 0 , but this leads to all answers having the impossible condition of negative concentration or negative pressure at equilibrium. That concentrations and partial pressures cannot be negative is an important concept since it is the only way to rule out the incorrect root(s) of a quadratic (polynomial) equation used to find the changes to reach equilibrium. | The ICE tables will be updated for questions 60 and 61. The answers will also be updated. | Other factual inaccuracy in content |
| Chapter 13 <br> Fundamental <br> Equilibrium <br> Concepts: Section <br> 13.4 Equilibrium <br> Calculations | The "check your learning" example problem associated with Example 13.13 is an equilibrium constant determination associated with the synthesis of nitrogen dioxide to dinitrogen tetroxide. A student and two instructors can't recreate the answer in the textbook of 6.9. Based on a delta-G of formation for NO2 of 99.8 $\mathrm{kJ} / \mathrm{mol}$ and for N 2 O 4 of $51.3 \mathrm{~kJ} / \mathrm{mol}$, we determine a delta-G of reaction of $-2.8 \mathrm{~kJ} / \mathrm{mol}$ and an equilibrium constant K of 3.1. | Revise the equation right before the answer to "N_2O_4 $(\mathrm{g}) \rightleftharpoons 2 \mathrm{NO}_{2} 2(\mathrm{~g})$ ". Revise the answer to "0.32". | Incorrect answer, calculation, or solution |
| Chapter 13 <br> Fundamental Equilibrium Concepts: Section 13.4 Equilibrium Calculations | (4th example in section, title of first problem is "Calculation of Concentration Changes as a Reaction Goes to Equilibrium") I believe the answers were added/subtracted to the wrong reverse values. <br> When solving this expression, $Q=7.1$, which is greater than $K$ therefore the reaction moves towards the reactants. When solving for x (after taking the square root of both sides) $x=0.033$ M . When adding to both reactants, acetic acid and ethanol, it is $0.15 \mathrm{M}+0.033 \mathrm{M}=0.18 \mathrm{M}$, and subtracting from both products, ethyl | Update the answer below Step 4 to $\begin{aligned} & {[\mathrm{CH} 3 \mathrm{CO} 2 \mathrm{H}]=0.37 \mathrm{M},[\mathrm{C} 2 \mathrm{H} 5 \mathrm{OH}]=} \\ & 0.37 \mathrm{M},[\mathrm{CH} 3 \mathrm{CO} 2 \mathrm{C} 2 \mathrm{H} 5]=0.18 \mathrm{M}, \\ & {[\mathrm{H} 2 \mathrm{O}]=0.18 \mathrm{M} .} \end{aligned}$ | Incorrect answer, calculation, or solution |


|  | acetate and water, it is $0.40 \mathrm{M}-0.03 \mathrm{M}=0.37$ M . The answer stated is acetic acid and methanol 0.37 M , and ethyl acetate and water 0.17 M . |  |  |
| :---: | :---: | :---: | :---: |
| Chapter 13 <br> Fundamental Equilibrium Concepts: Section 13.4 Equilibrium Calculations | The answer in the book and on-line are incorrect. If you plug the values into the equation for the equilibrium constant ( $\mathrm{K}=$ [products]/[reactants]) they do not give the value of 4.0. <br> The correct equilibrium concentrations of the reactants CH 3 CO 2 H and C 2 H 5 OH are 0.18 M , and of the products CH 3 CO 2 C 2 H 5 and H 2 O are 0.37 M . | Revise the Answer line below the Check Your Learning after Step 4 in this example to $[\mathrm{CH} 3 \mathrm{CO} 2 \mathrm{H}]=0.37$ <br> $\mathrm{M},[\mathrm{C} 2 \mathrm{H} 5 \mathrm{OH}]=0.37 \mathrm{M}$, $[\mathrm{CH} 3 \mathrm{CO} 2 \mathrm{C} 2 \mathrm{H} 5]=0.18 \mathrm{M},[\mathrm{H} 2 \mathrm{O}]=$ 0.18 M . | Incorrect answer, calculation, or solution |
| Chapter 13 <br> Fundamental Equilibrium Concepts: Section 13.4 Equilibrium Calculations | Exercise 13.52 answer keyThe first part of the equation shows the concentration values with the exponents. The second part of the solution has squared and cubed the relevant reagents, but the square and cube superscripts are still there. These need to be removed. | This solution will be updated. | Typo |
| Chapter 13 <br> Fundamental Equilibrium Concepts: Section 13.4 Equilibrium Calculations | Paired with Example 13.8 is a "Check your Learning" problem: reaction of acetic acid with ethanol to produce ethyl acetate and water. Based on the balanced equation mole ratios and the given equilibrium constant ( $\mathrm{K}=4$ ), the equilibrium concentrations of products should be .37 M and reactants .18 M . The change in concentration is 0.03333333333333 which rounds to 0.03 with sig figs making the substituted K $.37^{\wedge} 2 / .18^{\wedge} 2=4.225$. However, when the nonterminal decimal is carried through the calculations substituted K is (0.3666666666666)^2/(0.18333333333333)^2 <br> = 4. The text shows the equilibrium concentration of reactants (left side of equation) as 0.37 M and products (right side of equation) as 0.18 M which seems to be simple mix up of which side has which values at equilibrium. Incidentally, evaluating these concentrations yields an equilibrium constant of $K=(0.18)^{\wedge} 2 /(0.37)^{\wedge} 2=0.2366691015$ which seems to be the reciprocal of the given $K$ value of 4 . | Revise " 0.37 " terms to " 0.18 " and " 0.18 " terms to " 0.37 ". | Incorrect answer, calculation, or solution |
| Chapter 13 <br> Fundamental Equilibrium Concepts: Section 13.4 Equilibrium Calculations | example 13.5 Check your learning letter C shows a reversible reaction of ammonia with water. The reaction is not balanced as it has 26 hydrogen atoms on the left side but only 12 hydrogen atoms on the right side. Also, there are 7 oxygen atoms on the left side, but a total of 14 oxygen atoms on the right side. | Revise "7H_2O" to "70_2". | Other factual inaccuracy in content |
| Chapter 13 <br> Fundamental Equilibrium Concepts: Section 13.4 Equilibrium Calculations | In the short paragraph between Example 13.13 and the Figure 13.14, it refers to a plot of the "free energy change" versus extent of reaction. Of course it is "free energy", not the change which is plotted here. | Delete the word "change" in the sentence "As may be shown by plotting the free energy change versus the extent...." | General/pedago gical suggestion or question |


| Chapter 13 <br> Fundamental <br> Equilibrium <br> Concepts: Section 13.4 Equilibrium Calculations | Q90 I am pretty sure I have been doing these thermodynamic problems correctly but when I check the student solution manual it says that I am wrong. I don't think the solution manual that I have downloaded from your website includes the correct values being used. I am under the impression that question \# 43 (as well as other previous questions) is asking for the student to use Appendix G in order to answer the problem using the constants found for delta H and delta S in order to find delta G to use in another formula. However, when you use those constants to do the problem, it seems that the solution manual is using a different constant that inn't anywhere in the appendix. | The solution for question 90 will be updated as follows: (a) $\mathrm{K}=1.07 \times$ $10^{\wedge}-13$; (b) $K=2.42 \times 10^{\wedge}-3$; (c) $K=$ $2.73 \times 10^{\wedge} 4$; (d) $K=0.229$ (e) $K=$ 16.1. | Incorrect answer, calculation, or solution |
| :---: | :---: | :---: | :---: |
| Chapter 13.2-13.4 | There are several superscripting and subscripting errors throughout the chapter. | These superscripting and subscripting errors will be corrected. | Typo |
| Chapter 14 Acid- <br> Base Equilibria: 14.3 Relative Strength of Conjugate Acids and Bases | On the Kb constant appears [OH], it should be [OH-]. | Revise to $\left[\mathrm{OH}^{\wedge}-\right]$. | Typo |
| Chapter 14 AcidBase Equilibria: Introduction | On the first sentence of the paragraph, there is a typo on OH . It should be $\mathrm{OH}^{\wedge}$-, not $\mathrm{OH}^{\wedge}$ a, | Revise to "OH^-". | Typo |
| Chapter 14 Acid- <br> Base Equilibria: <br> Section 14.1 <br> Brønsted-Lowry <br> Acids and Bases | In the first equation of example 14.1, Kw should be equal to $x^{*} x$ instead of $x$. | Revise x to $(\mathrm{x})(\mathrm{x})=\mathrm{x}^{\wedge} 2$. | Incorrect answer, calculation, or solution |
| Chapter 14 Acid- <br> Base Equilibria: <br> Section 14.2 pH and pOH | Exercise 19 Calculate the pH and pOH of b) 0.21 M NaOH and d) 2.5 M KOH . Both questions have answers for which the pH and pOH are reversed. (These strong bases have pH's over 7, not under 7.) | For part b) and d), revise "pH" to "pOH" and vice versa. | Incorrect answer, calculation, or solution |
| Chapter 14 Acid- <br> Base Equilibria: <br> Section 14.3 <br> Relative Strengths <br> of Acids and Bases | $60 a$ and $b$ The first several lines of the solutions are correct, but at the final concentrations things fell apart. <br> a) $[\mathrm{H3O}+]=[\mathrm{ClO}-]=1.6 \times 10^{\wedge}-5$ (missing charge on ion, wrong conc) $[\mathrm{HClO}]=0.0092 \mathrm{M}$ (an extra zero in current answers) <br> b) wrong ion [C6H5NH3+] (not CH3CO2-) and wrong conc [C6H5NH2] $=0.0784 \mathrm{M}$ | The answer will be updated. | Incorrect answer, calculation, or solution |
| Chapter 14 AcidBase Equilibria: Section 14.3 Relative Strengths of Acids and Bases | 58b, c, e Answers have the wrong powers of 10. b) Should be Ka= $1.6 \times 10^{\wedge}-10$ c) Should be $\mathrm{Kb}=5.9 \times 10^{\wedge}-8$ (And ion should have been HAsO4^2-). e) Should be Kb $=2.3 \times 10^{\wedge}-5$ | In the question stem, revise part (c) to "HAsO_4^2- (as a base)". <br> Revise the answers to: <br> b) $K \_a=1.6 \times 10^{\wedge}-10$ <br> c) K _ $b=5.9 \times 10^{\wedge}-8$ <br> e) K _ $b=2.3 \times 10^{\wedge}-5$ | Incorrect answer, calculation, or solution |


| Chapter 14 AcidBase Equilibria: Section 14.3 Relative Strengths of Acids and Bases | 54c The answer in the back of the book is incorrect. It should be $6.4 \times 10^{\wedge}-5$ and not $7.4 \times$ 10^-5. SB | Revise the answer to "6.4 $\times 10^{\wedge}-5$ ". | Incorrect answer, calculation, or solution |
| :---: | :---: | :---: | :---: |
| Chapter 14 AcidBase Equilibria: Section 14.3 Relative Strengths of Acids and Bases | \# 28 I assume the intent of part b) and c) is to determine the weak acids and weak bases from the list in the problem prompt. As written this is not clear since in part a) the strong acids and bases are identified, it would seem to indicate from the set of strong acids and bases identify which are weak, which is not possible. <br> Currently: <br> (a) Identify the strong Brønsted-Lowry acids and strong Brønsted-Lowry bases. <br> (b) List those compounds in (a) that can behave as Brønsted-Lowry acids with strengths lying between those of $\mathrm{H} 3 \mathrm{O}+$ and H 2 O . <br> (c) List those compounds in (a) that can behave as Brønsted-Lowry bases with strengths lying between those of H 2 O and OH -. <br> Proposed correction: <br> (a) Identify the strong Brønsted-Lowry acids and strong Brønsted-Lowry bases. <br> (b) List the compounds, not already in (a), that can behave as Brønsted-Lowry acids with strengths lying between those of $\mathrm{H} 3 \mathrm{O}+$ and H2O. <br> (c) List the compounds, not already in (a), that can behave as Brønsted-Lowry bases with strengths lying between those of H 2 O and $\mathrm{OH}-$. | In the question stem, delete "CaO" and "CO_2". In parts (b) and (c), revise "List those compounds in (a)..." to "Identify the compounds...". The solution will also be updated. | General/pedago gical suggestion or question |
| Chapter 14 AcidBase Equilibria: Section 14.3 Relative Strengths of Acids and Bases | 14.33 This problem asks for a balanced equation and to identify the conjugate pairs. In the answers at the back of the book the CA and CB are misidentified. Perhaps it is just a spacing issue and you need to move the labels under the Cl - and H 2 O . | The alignment will be updated. | Incorrect answer, calculation, or solution |
| Chapter 14 Acid- <br> Base Equilibria: <br> Section 14.3 <br> Relative Strengths <br> of Acids and Bases | Exercise 58c The original ion had the incorrect " 3 -" charge. It has been updated online incorrectly to "2" rather than to "2-" SB | Revise the superscript "2" to superscript "2-". | Typo |
| Chapter 14 AcidBase Equilibria: Section 14.3 Relative Strengths of Acids and Bases | Chapter 14 End-of-Chapter Exercise 58c The original ion had the incorrect "3-" charge. It has been updated online incorrectly to "2" rather than to "2-" SB | Revise the superscript "2" to superscript "2-". | Typo |
| Chapter 14 AcidBase Equilibria: <br> Section 14.4 <br> Hydrolysis of Salts | Example 14.7, Solution Kb of F - should be $1.6 \times 10^{\wedge}-11$ instead of $1.4 x^{\wedge}-11$, since $1 \times 10^{\wedge}$ $14 / 6.4 \times 10^{\wedge}-4=1.6 \times 10^{\wedge}-11$. The Ka of HF is $6.4 \times 10^{\wedge}-4$. | $\begin{aligned} & \text { Revise " } 1.4 \times 10^{\wedge}-11 \text { " to } " 1.6 \times \\ & 10^{\wedge}-11 \text { ". } \end{aligned}$ | Incorrect answer, calculation, or solution |
| Chapter 14 AcidBase Equilibria: | The equation is written without reaction arrow and the charge for the sodium is not written as a superscript. | Revise so the "+" charge after Na is set as a superscript. | Other |


| Section 14.4 Hydrolysis of Salts |  |  |  |
| :---: | :---: | :---: | :---: |
| Chapter 14 AcidBase Equilibria: <br> Section 14.4 Hydrolysis of Salts | In Example 14.15 the formula of anilinium chloride in the first line of the problem either needs to have the positive charge within the brackets removed or the negative sign added after the Cl . The compound is not an ion as is currently shown. SB | Remove the superscripted "+". | Typo |
| Chapter 14 AcidBase Equilibria: Section 14.4 Hydrolysis of Salts | After the 3rd equation in this section, "ammonia, NH3;" the 3 must be subscripted. SB | Set the "3" as subscript. | Typo |
| Chapter 14 AcidBase Equilibria: Section 14.4 Hydrolysis of Salts | The first two paragraphs in the Salts with Acidic Ions subsection (the explanation of acid hydrolysis) are repeated successively. | The repeated content will be deleted. | Typo |
| Chapter 14 AcidBase Equilibria: Section 14.5 Polyprotic Acids | Example 14.19 The question and solution use the Ka 2 of H 2 CO 3 value given in Appendix H $\left(4.7 \times 10^{\wedge}-11\right)$, which is also the answer for the [CO3^2-]. <br> The summary section at the end then lists the [CO3^2-] as $5.6 \times 10^{\wedge}-11$, which is a value for the Ka2 of H2CO3 given in other textbooks (like McMurry). | $\begin{aligned} & \text { Revise " } 5.6 \times 10^{\wedge}-11 \mathrm{M} \text { " to " } 4.7 \times \\ & 10^{\wedge}-11 \mathrm{M} \text { ". } \end{aligned}$ | Incorrect answer, calculation, or solution |
| Chapter 14 AcidBase Equilibria: Section 14.5 Polyprotic Acids | In the last equation Bicarbonate ion is written as $\mathrm{HCO}^{\wedge} 3$-, it should be $\mathrm{HCO}^{\wedge}$-1 | Revise "HCO^3-" to "HCO_3^-". | Typo |
| Chapter 14 AcidBase Equilibria: Section 14.6 Buffers | Figure 14.14 caption: The (un)buffered solution on the left and the ()buffered solution on the right | Revise "The buffered solution on the left and the unbuffered solution on the right..." to "The unbuffered solution on the left and the buffered solution on the right..." | Typo |
| Chapter 14 AcidBase Equilibria: Section 14.6 Buffers | Example 14.20 Calculation (c), the equation between the hidronium and hydroxide is missing the arrows. | Add arrows after "(aq)". | Typo |
| Chapter 14 AcidBase Equilibria: Section 14.6 Buffers | Example 14.20 Calculation (c), says "hydrogen ion", it should say "hidronium" ion. | Revise "hydrogen ion" to "hydronium ion". | Typo |
| Chapter 14 AcidBase Equilibria: Section 14.6 Buffers | The solution to part c of 14.90 makes no sense. There is a calculation for NH 4 Cl which is not in the problem at all. The volume changes from 0.200 liters in the problem to 0.5 . It is just a mess. <br> Also, there is really no reason at his level not to use the Henderson Hasselbalch Equation to solve this problem. <br> The whole point of this exercise is that adding acid doesn't really change the pH significantly. That gets lost in the mess. | This solution will be updated. | Incorrect answer, calculation, or solution |


| Chapter 14 AcidBase Equilibria: Section 14.6 Buffers | Solution to Exercise 80 In the ICE table, the concentration of acetate at equilibrium is listed as 0.030-x it should be 0.030+x This doesn't change the results since we ignore $x$. But it would be confusing to students. | This solution will be updated. | Typo |
| :---: | :---: | :---: | :---: |
| Chapter 14 Acid- <br> Base Equilibria: <br> Section 14.7 Acid- <br> Base Titrations | In Acid-Base Indicators after explaining the use of the Henderson-Hasselbach equation for a methyl orange solution, when explaining that the log term of the equation should be negative if $\mathrm{pH}<\mathrm{pKa}$ is wrongly written as $\mathrm{pH}>\mathrm{pKa}$. This is written correctly in the first sentence where the log term is positive, thus, $\mathrm{pH}>\mathrm{pKa}$. | Revise "pH > pK_a" to "pH < pK_a". | Typo |
| Chapter 14 AcidBase Equilibria: Section 14.7 AcidBase Titrations | part D of Example 14.21. During the equation a 35.70 was put in instead of 37.50 which caused the answer to be incorrect. | Revise as indicated. | Incorrect answer, calculation, or solution |
| Chapter 15 <br> Equilibria of Other <br> Reaction Classes: <br> Section 15.1 <br> Precipitation and Dissolution | Ch15, Q7 This is a question about which of a series of salts would likely be affected by hydrolysis of the anion. The candidates include both PbS and ZnS , but only ZnS is indicated in the answer key as such a salt. I'm no inorganic chemist, but I can't see why the anions would differ in this case. I imagine that lead(II) might be able to form hydroxide complexes that zinc maybe doesn't, but as far as the sulfide goes, I think they would be treated similarly. My feeling is that the question writer intended to write either ZnS or PbS , but not both. More generally, I'm not sure where in the chapter this topic is discussed. | Delete ", PbS" from the end of question 7. | Incorrect answer, calculation, or solution |
| Chapter 15 <br> Equilibria of Other <br> Reaction Classes: <br> Section 15.1 <br> Precipitation and Dissolution | The answer key gives the hydroxide concentration here as $4.5 \mathrm{e}-5$. I think it's 4.5e-6. Appendix J gives the $\mathrm{Al}(\mathrm{OH}) 3 \mathrm{Ksp}$ as $2 \mathrm{e}-32$. The molar solubility of the compound is reported as 2.2e-20, which doesn't appear to work for that Ksp with either $4.5 \mathrm{e}-5$ or $4.5 \mathrm{e}-6$. I think it's 2.2e-16. | Revise "10^-5" to "10^-6" and "10^-20" to "10^-16". | Incorrect answer, calculation, or solution |
| Chapter 15 <br> Equilibria of Other <br> Reaction Classes: <br> Section 15.1 <br> Precipitation and Dissolution | Chapter 15, Exercise 34 requires the use of Ksp for strontium fluoride, SrF2, but there is no value for the Ksp for strontium fluoride in Appendix J | Revise SrF_2 to SrCrO_4 and revise F_- to $\mathrm{CrO}_{-} 4^{\wedge} 2-$. The solution will be updated. | Other |
| Chapter 15 <br> Equilibria of Other <br> Reaction Classes: <br> Section 15.1 <br> Precipitation and Dissolution | End of Chapter question 59. (1) The listed Ksp values for the five salts are all different from the values listed in Appendix J. (2) Based on the given values, the answer should be e, but the given answer key was c. | Revise "MnCO3" to "MgCO3". | Incorrect answer, calculation, or solution |


| Chapter 15 <br> Equilibria of Other <br> Reaction Classes: <br> Section 15.1 <br> Precipitation and Dissolution | The answer to Chapter 15 \#59 states "MnCO3 will form first, since it has the smallest Ksp value it is the least soluble. MnCO 3 will be the last to precipitate, it has the largest Ksp value." Obviously MnCO3 cannot have simultaneously the largest and smallest value! BaCO 3 will be the last to precipitate since it has the largest Ksp value. | Revise the question to: The carbonate ion concentration is gradually increased in a solution containing equal concentrations of the divalent cations of magnesium, calcium, strontium, barium, and manganese. Which of the following carbonates will precipitate first? Which will precipitate last? Explain. <br> (a) $\mathrm{MgCO} 3 \cdot 3 \mathrm{H} \_2 \mathrm{O} \mathrm{K} \_\mathrm{sp}=1 \times$ 10^-5 <br> (b) CaCO_3 K_sp $=8.7 \times 10^{\wedge}-9$ <br> (c) SrCO_3 K_sp $=7 \times 10^{\wedge}-10$ <br> (d) $\mathrm{BaCO}_{3}$ K_sp $=1.6 \times 10^{\wedge}-9$ <br> (e) MnCO 3 K_sp $=8.8 \times 10^{\wedge}-11$ <br> Revise the answer to "MnCO_3 will form first since it has the smallest K_sp value among these homologous compounds and is therefore the least soluble. <br> $\mathrm{MnCO} 3 \cdot 3 \mathrm{H} \_2 \mathrm{O}$ will be the last to precipitate since it has the largest K_sp value and is the most soluble." | Incorrect answer, calculation, or solution |
| :---: | :---: | :---: | :---: |
| Chapter 15 <br> Equilibria of Other <br> Reaction Classes: <br> Section 15.1 <br> Precipitation and <br> Dissolution | 15.1 Exercise 1d Formatting error in that " $x$ " is not under $3 \mathrm{Mg}^{\wedge} 2+$ but is with the blank line under 2PO4^3-. Additionally, the solutions use $3 x$ as the "answer" and $2 x$ as the given. | The alignment will be updated. Also update the answer to " $x$ " and " $2 / 3 x$ ". | Typo |
| Chapter 15 <br> Equilibria of Other <br> Reaction Classes: <br> Section 15.1 <br> Precipitation and Dissolution | The error is in the worked out solution to Precipitation of Silver Halides towards the end of 15.1 (using the online text version). The concentration substituted for iodide ions should be 0.0010 M and not 0.10 M . | While the example itself is correct, there are a few typos that will be corrected. Revise Q to Q_sp = K_sp. | Typo |
| Chapter 15 <br> Equilibria of Other <br> Reaction Classes: <br> Section 15.1 <br> Precipitation and Dissolution | Example 15.13 Answer given is $1.10 \times 10^{\wedge}-10 \mathrm{M}$. This is actually the value for the concentration of OH - at which $\mathrm{Al}(\mathrm{OH}) 3$ begins to precipitate, but the question is asking for the molar solubility of $\mathrm{Al}(\mathrm{OH}) 3$. The correct answer should be $3.67 \times 10^{\wedge}-11 \mathrm{M}$. | Revise the answer to " $4 \times 10^{\wedge}-11$ M". | Incorrect answer, calculation, or solution |
| Chapter 15 <br> Equilibria of Other <br> Reaction Classes: <br> Section 15.1 <br> Precipitation and Dissolution | Example 15.5 The answer of the solubility is written as $2.08 \times 10^{\wedge}-4$, whereas it should be $1.7 \times 10^{\wedge}-4$ since the molarity of any of both ions is $1.3 \times 10^{\wedge}-2 .\left(1.3 \times 10^{\wedge}-2\right)\left(1.3 \times 10^{\wedge}-2\right)=1.7 \times 10^{\wedge}-4$ | Revise the answer to "1.69 $\times 10^{\wedge}-4$ ". | Incorrect answer, calculation, or solution |
| Chapter 15 <br> Equilibria of Other <br> Reaction Classes: <br> Section 15.1 <br> Precipitation and Dissolution | Example 15.12 There are 7 total answers to this example, whereas there should be only 4 . There is (a), (b), (c) and on the last one the last sentence is interrupted, then continues another (a), then another (b) and another (c). It finishes with one (d). | Delete the first (a), (b), and (c) and associated text in the solution. | Other |


| Chapter 15 <br> Equilibria of Other <br> Reaction Classes: <br> Section 15.1 <br> Precipitation and Dissolution | After the introduction Common Ion Effect it provides two "links to learning" that take you to the same page. | Delete the second Link to Learning box. | Other |
| :---: | :---: | :---: | :---: |
| Chapter 15 <br> Equilibria of Other <br> Reaction Classes: <br> Section 15.1 <br> Precipitation and Dissolution | After calculating the Q , it says "Agl", it should say AgBr . | Revise "Agl" to "AgBr". | Typo |
| Chapter 15 <br> Equilibria of Other <br> Reaction Classes: <br> Section 15.1 <br> Precipitation and Dissolution | In the calcium carbonate equilibrium reaction, the carbonate ion superscript isn`t written as a superscript, just as an ordinary number. | Revise so that "2-" is set as a superscript. | Typo |
| Chapter 15 <br> Equilibria of Other <br> Reaction Classes: <br> Section 15.1 <br> Precipitation and Dissolution | Example 15.15 There are a couple of typo: 1. Ca should be Al. 2. molar solubility in water should be 1/4. 3. [Al2+] should be [Al3+] | $\begin{aligned} & \text { Revise "Ca" to "Al", " } 1 / 3 \text { " to " } 1 / 4 \text { ", } \\ & \text { " } 9 \times 10^{\wedge}-12 \mathrm{M}^{\prime \prime} \text { to " } 5 \times 10^{\wedge}-9 \mathrm{M}^{\prime \prime} \text {, } \\ & \text { and "Al^2+" to "Al^3+". } \end{aligned}$ | Typo |
| Chapter 15 <br> Equilibria of Other <br> Reaction Classes: <br> Section 15.1 <br> Precipitation and Dissolution | There are numerous typos in examples 15.15 and 15.16. In example 15.15, a solubility calculation is performed for aluminum hydroxide, but in the first line, this compound is written $\mathrm{Ca}(\mathrm{OH}) 2$, and two lines later, is called calcium fluoride. In example 15.16, in line 6 of the solution, the bromide ion is written incorrectly with the negative charge as a subscript rather than a superscript. Also, that line gives a number of moles of 0.00532 mol , which is incorrectly written as 0.00521 mol on the following lines. Also, the thiosulfate ion in several places and silver thiosulfate in one place is written incorrectly, again confusing subscripts and superscripts. | These examples will be updated. | Typo |
| Chapter 15 <br> Equilibria of Other <br> Reaction Classes: <br> Section 15.1 <br> Precipitation and Dissolution | Example 15.12 Check your learning the solubility of $\mathrm{Al}(\mathrm{OH}) 3$ is calculated incorrectly | Revise the answer to " $1.10 \times 10^{\wedge}$ 10". | Incorrect answer, calculation, or solution |
| Chapter 15 <br> Equilibria of Other <br> Reaction Classes: <br> Section 15.2 Lewis <br> Acids and Bases | Answer to question 64 The concentration of 0.001 mol of [Cd2+] ions in the resulting 1.250 L solution should be $8 \times 10^{\wedge}-4$ (not $8 \times 10^{\wedge}-5$ ). | Revise "1.150 L" to "0.150 L" in the question. Revise the answer to "1.8 $\times 10^{\wedge}-5 \mathrm{M}$ ". | Incorrect answer, calculation, or solution |
| Chapter 15 <br> Equilibria of Other <br> Reaction Classes: | The "check your learning" portion of Example 15.14 The answer is listed as $2.5 \mathrm{e}-22 \mathrm{M}$, however the correct answer should be $2.9 \mathrm{e}-$ | Revise "2.5" to "2.9" in the answer. | Incorrect answer, |
| Section 15.2 Lewis <br> Acids and Bases | 22 M . Assuming AgNO3 is the limiting reagent and the reaction goes to completion, you would need to consume a total of 0.012 M of KCN . Since you start with 0.154 M of KCN , the leftover KCN should be 0.142 M of KCN . Setting up the ICE table and going through the calculations yields a silver ion concentration of $2.9 \mathrm{e}-22 \mathrm{M}$. The answer listed in the solution is $2.5 \mathrm{e}-22 \mathrm{M}$. I believe that the book authors arrived at this solution by using 0.154 M of KCN in the ice table instead of 0.142 M of KCN (ie it did not subtract the amount of CN used to make $\mathrm{Ag}(\mathrm{CN}) 2$-. If you set up the ICE table and use 0.154 M of KCN , you would get exactly 2.5 e 22M. |  | calculation, or solution |
| :---: | :---: | :---: | :---: |
| Chapter 15 Equilibria of Other Reaction Classes: Section 15.3 Coupled Equilibria | Exercise 105 HPO4- should be HPO4^2- | Revise "HPO_4^-" to "HPO_4^2-". | Typo |
| Chapter 15 <br> Equilibria of Other <br> Reaction Classes: <br> Section 15.3 <br> Coupled Equilibria | \#99 In the problem Ksp for MnS should be listed as $2.3 \times 10^{\wedge}(-13)$ as per Appendix J. In the problem K should be listed as $8.9 \times 10^{\wedge}(-27)$ if calculated from from K_a1 and K_a2 for hydrogen sulfide, as given in Appendix H | $\begin{aligned} & \text { Revise the } \mathrm{Ksp}=\text { value to } 2.3 \times \\ & 10^{\wedge}-13 " . \end{aligned}$ | Incorrect answer, calculation, or solution |
| Chapter 15 <br> Equilibria of Other <br> Reaction Classes: <br> Section 15.3 <br> Coupled Equilibria | There are some serious problems with Example 15.16. In the first part, they calculate the Br concentration of the fully dissolved AgBr . That is fine and 0.0532 is the correct result. <br> The second part is mathematically correct but would only be chemically correct if there were still some AgBr solid present. That means that there is still some film that has not been fixed at this point. The problem is unclear since it sounds like you are dissolving all the AgBr . The problem only works as they solve it, if there is still some undissolved AgBr present As soon as the precipitate is completely dissolved, then the free Ag+ ion concentration is only controlled by the thiosulfate equilibrium. Since the Br concentration is based on the total amount of AgBr dissolving, this is inconsistent. I suppose they are assuming that there is still some AgBr in the film that has not dissolved. <br> The next step is wierd becaue the math doesn't work. $0.00532-9.4 \times 10^{\wedge}-11$ is still 0.00532 । have no idea where the value 000521 comes from. If there is still AgBr undissolved on the film, then the free $\mathrm{Ag}+$ must come from the AgBr solubility and will be the $9.4 \times 10^{\wedge}-11$ that is calculated here. One can then put this into the formation constant equilibrium expression and determine the free thiosulfate that is not complexed with silver. Using the correct 0.00532 for the silver complex, the math gives $1.09 \times 10^{\wedge}-3=1.1 \times 10^{\wedge}-3 \mathrm{M}$ thiosulfate | This example will be updated. | Incorrect answer, calculation, or solution |
|  | uncomplexed. The problem in the text incorrectly has $1.15 \times 10^{\wedge}-3$ instead because they use 0.00521 for the silver complex. Both end up with $1.1 \times 10^{\wedge}-3$. <br> The final step for the problem is incomplete. The final solution that has dissolved the 1.0 g AgBr must contain $1.1 \times 10^{\wedge}-3$ molar thiosulfate that is free, not complexed with silver. The formation constant equilibrium requires that to keep the AgBr dissolved. It takes 1.7 g of sodium thiosulfate to reach this concentration in the absence of AgBr . however, it takes another $0.00532 \times 2$ moles of thiosulfate to form the complex. So that requires 0.0106 moles of additional thiosulfate or 1.68 g . So the total thiosulfate required is twice the amount in the problem. It takes 1.7 g to form the complex and an excess of 1.7 g to reach the free thiosulfate concentration required. |  |  |
| :---: | :---: | :---: | :---: |
| Chapter 15 <br> Equilibria of Other <br> Reaction Classes: <br> Section 15.3 <br> Coupled Equilibria | The first error is a typo in the molarity of Silver ion (also the molarity of Silver Thiosulfide complex ion). The third step of the calculation shows the $\mathrm{Ag}(\mathrm{S} 2 \mathrm{O} 3) 2$ 3- concentration as 0.00532 M . The fourth step and fifth step uses 0.00521 M with no explanation given for the change in molarity. The overall answer is also incorrect -1.00 L of $1.1 \times 10^{\wedge}-3 \mathrm{M}$ Sodium Thiosulfate should have a mass of 0.17 grams. The answer says 1.7 grams which seems like $1.1 \times 10^{\wedge}-2 \mathrm{M}$ solution was present. | Replace two occurrences of "0.00521" with "0.00532". Replace " $1.15 \times 10^{\wedge}-6$ " with " $1.18 \times 10^{\wedge}-6$ ". Replace two occurrences of " 1.1 x $10^{\wedge}-3^{\prime \prime}$ with " $1.08 \times 10^{\wedge}-3^{\prime \prime}$. Replace two occurrences of " 1.7 " with "0.17". | Incorrect answer, calculation, or solution |
| Chapter 15 <br> Equilibria of Other <br> Reaction Classes: <br> Section 15.3 <br> Coupled Equilibria | 15.91 At the end of the fourth line (solving for Kf ), the denominator has the term ( 0.025 + 6Delta). This needs to be raised to the 6th power as it is shown earlier in the same line in the Kf expression. Unfortunately, when solved using the $x$-is-small approximation, there is no valid solution. So students would have to expand a 6th-order polynomial, and then carry out the remaining operations on all those terms in order to solve this properly. Keeping similar numbers, use 0.0100 mol of the complex ion salt. Use 0.25 M ammonia. the solution will be [Co2+] = $3.2 \times 10^{\wedge-4}$ | In the question, revise " 0.100 " to " 0.010 " and " 0.025 " to " 0.25 ". The solution will also be updated. | Incorrect answer, calculation, or solution |
| Chapter 15 <br> Equilibria of Other <br> Reaction Classes: <br> Section 15.3 <br> Coupled Equilibria | Q99 Ksp should be $4.3 \times 10^{\wedge}-22$ and not $2.3 \times$ $10^{\wedge}-13$. k should be $1 \times 10^{\wedge}-26$ and not $8.9 \times$ $10^{\wedge}-27$, answer should be calculated based on the Ksp and $k$ values above | The values will be updated in webview to match the current solution manual. | Typo |
| Chapter 16 <br> Electrochemistry: <br> Key Equations | the fifth bullet is only the right side of the equation. there should be an $E(c e l l)=$ to the left of the text that is included. | Add "E_cell =" before the current equation. | Typo |
| Chapter 16 Electrochemistry: Section 16.1 Review of Redox Chemistry | Example 16.2 There's a missing plus sign after the chromate ion. | Add "+" before "H+(aq)". | Tуро |
| Chapter 16 <br> Electrochemistry: <br> Section 16.1 Review <br> of Redox Chemistry | Example 16.1 Step 7: After the nitrogen monoxide gas, there is a plus sign missing before the 4 molecules of water. | Add "+" before "4H2O(I)". | Typo |
| :---: | :---: | :---: | :---: |
| Chapter 16 <br> Electrochemistry: <br> Section 16.1 Review of Redox Chemistry | Exercise 3 solution to part d. MnO4 ion is missing its charge ( -1 ) | Revise "MnO4" to "MnO_4^-". | Typo |
| Chapter 16 <br> Electrochemistry: <br> Section 16.2 <br> Galvanic Cells | Example 16.3 In the Solution, in the halfreactions, the cathode is mislabeled as "(oxidation)", whereas it should be "(reduction". | Revise "cathode (oxidation)" to "cathode (reduction)". | Typo |
| Chapter 16 <br> Electrochemistry: <br> Section 16.3 <br> Electrode and Cell <br> Potentials | example 16.5 Check Your Learning The first equation (from top to bottom) should be nonspontaneous since the bromide ion experiments an oxidation to bromine (anode). Hence the negative sign. The second should be corrected too following the explanation. | Revise "+0.5518 V (spontaneous)" to " -0.5518 V (nonspontaneous)" and <br> "-0.5518 V (nonspontaneous)" to <br> "+0.5518 V (spontaneous)". | Incorrect answer, calculation, or solution |
| Chapter 16 <br> Electrochemistry: <br> Section 16.3 <br> Electrode and Cell <br> Potentials | Question 17.25 labels both half cells as anodes. The solution assumes the aluminum is the cathode. The question should either state the reaction will be spontaneous/nonspontaneous. Or it should combine one each of a cathode and anode. | Revise "an anode" to "a cathode". | Typo |
| Chapter 16 <br> Electrochemistry: <br> Section 16.4 Free <br> Energy | Tthe answer for the 16.8 check your learning the answer is given negative and then nonspontaneous. Was that right because it is kind of confusing | In Example 16.8 Calculation of delta G 298, revise Check Your Learning answer as follows: (a) $140.8 \mathrm{~kJ} / \mathrm{mol}$, nonspontaneous (b) $141.5 \mathrm{~kJ} / \mathrm{mol}$, nonspontaneous | Typo |
| Chapter 16 <br> Electrochemistry: <br> Section 16.4 Free <br> Energy | In the student solution guide for chapter 12(mod 04) \#62, the original problems says the concentration of G6P is 120 micromolar, however 128 was substituted into the equation. Also, the temperature for 37 C should be 310 K , not 335 K . The solution should read "delta $\mathrm{G}=$ $1.7 \times 10^{\wedge} 3+(8.314 \times 310 \times \ln 28 / 210)=-2.1 \mathrm{~kJ}{ }^{\prime \prime}$ | Revise the solution for exercise 62 as follows: "delta G = $1.7 \times 10^{\wedge} 3+$ $(8.314 \times 310 \times \ln 28 / 210)=-2.1 \mathrm{~kJ} "$ | Incorrect answer, calculation, or solution |
| Chapter 16 <br> Electrochemistry: <br> Section 16.4 Free <br> Energy | before example 16.7 The last equation is written with an minus sign for the normal potential of the cell, this should be positive. | Revise "InQ" to "logQ" in the equation right before Example 16.7. | Typo |
| Chapter 16 <br> Electrochemistry: <br> Section 16.4 Free <br> Energy | Under "Potentials at Nonstandard Conditions: <br> The Nernst Equation", where the base-10 logarithm version is presented. Somehow a minus sign has snuck in, in front of \$E_\{cell\}^\{0\}\$ on the right side of the equation. The equation should use \$log_\{10\} Q\$ instead of $\$ \ln Q \$$ to avoid confusion. I'd also like to add that the text doesn't specify that the 0.0592 V "constant" is derived at the absolute temperature 298.15 K ( $=25$ degrees C ). This should be included in the edit to avoid erroneous use at other temperatures. | Revise the sentence before the equation to "A convenient form of the Nernst equation for most work is one in which values for the fundamental constants (R and F) and standard temperature (273.15 K), along with a factor converting from natural to base-10 logarithms, have been included:" <br> The equation will be updated in errata 11706. | Other factual inaccuracy in content |
$\left.\begin{array}{|l|l|l|l|}\hline \begin{array}{l}\text { Chapter 16 } \\ \text { Electrochemistry: } \\ \text { Section } 16.4 \text { Free } \\ \text { Energy }\end{array} & \begin{array}{l}\text { EOC problem \#29(c). The words "bromine ion" } \\ \text { are confusing. It says "aqueous bromide is } \\ \text { oxidized to 0.11 M bromine ion" - bromine is } \\ \text { not the ion in this case and it is } \\ \text { confusing. Leaving out "ion" would be } \\ \text { sufficient to correct this error. }\end{array} & \begin{array}{l}\text { Revise the answer to part (c) to "The } \\ \text { cell comprised of a half-cell in which } \\ \text { aqueous bromine (1.0 M) is being } \\ \text { oxidized to bromide ion (0.11 M) } \\ \text { and a half-cell in which Al^3+ (0.023 } \\ \text { M) is being reduced to aluminum } \\ \text { metal." }\end{array} & \\ \hline \begin{array}{l}\text { Chapter 16 } \\ \text { Electrochemistry: } \\ \text { Section 16.5 } \\ \text { Batteries and Fuel } \\ \text { Cells }\end{array} & \begin{array}{l}\text { Under the section that is discussing the lead } \\ \text { acid battery, the set of reactions shown pertain } \\ \text { to the lithium ion battery. The resolution is to } \\ \text { replace those reactions with the lead/lead } \\ \text { oxide reaction with sulfuric acid. I believe this is } \\ \text { a "chemical typo." }\end{array} & \text { This set of reactions will be updated. }\end{array} \begin{array}{l}\text { Other factual } \\ \text { inaccuracy in } \\ \text { content }\end{array}\right\}$
|  | and not "What is the half-life (ms) for the butadiene dimerization reaction described in Example 17.8?" |  |  |
| :---: | :---: | :---: | :---: |
| Chapter 17 Kinetics: <br> Section 17.4 <br> Integrated Rate <br> Laws | Answer for Chapter 17 \#44 for half life is given as: $1.67 \times 10^{\wedge} 3 \mathrm{~s}$. The answer in the instructors solution manual is give as: $1.16 \times 10^{\wedge} 3$. The instructors solution manual is correct, the answer in the text book is incorrect. | Revise the answer from "1.67" to "1.16". | Incorrect answer, calculation, or solution |
| Chapter 17 Kinetics: <br> Section 17.4 <br> Integrated Rate <br> Laws | Example 12.12 There is an arithmetical error. $1 /(0.0576 \mathrm{~L} / \mathrm{mol} / \mathrm{min} * 0.200 \mathrm{~mol} / \mathrm{L})=86.8 \mathrm{~min}$. The example erroneously shows 0.0115 min , which is the reciprocal of the correct solution. The example then converts the incorrect solution to ms, which doesn't make sense given that the correct solution is more convenient in minutes. | Revise " 0.0115 min" to " 86.8 min". Delete the second equation in the solution. |  |
| Chapter 17 Kinetics: <br> Section 17.4 <br> Integrated Rate <br> Laws | The answer given for the Check Your Learning problem after Example 17.2 is incorrect. The answer should be 18 minutes (not 86.8 minutes). | Revise the answer to "18 min". | Incorrect <br> answer, calculation, or solution |
| Chapter 17 Kinetics: <br> Section 17.6 <br> Reaction <br> Mechanisms | Chapter 12 End-of-Chapter 71c The equation given is not a balanced equation which makes the question moot and the answer provided incorrect. SB | Revise "O" to "O_2". | Incorrect answer, calculation, or solution |
| Chapter 17 Kinetics: <br> Section 17.6 <br> Reaction <br> Mechanisms | Example 17.14 The reaction mechanism sums to the overall reaction in this example correctly. But the slow step will impart a one headed arrow on the overall reaction, so the equilibrium arrow should be switched for a oneheaded arrow in the overall reaction. | Revise " $\rightleftharpoons$ ' to " $\rightarrow$ ". | Typo |
| Chapter 17 Kinetics: <br> Section 17.6 <br> Reaction <br> Mechanisms | The example includes 3 places where reaction arrows are incorrect. Each is highlighted in the attached pdf. In two places, equal signs should be replaced with equilibrium reaction arrows. In one place, the reaction arrow is missing completely. The reactions are: <br> $\mathrm{NO}(\mathrm{g})+\mathrm{Cl} 2(\mathrm{~g})$ [EQUILIBRIUM ARROWS] NOCl 2 $(\mathrm{g})$, then $2 \mathrm{NO}(\mathrm{g})+\mathrm{Cl} 2(\mathrm{~g})$ [ONE-HEADED ARROW] 2NOCl(g), finally F2(g) [EQUILIBRIUM ARROWS] 2F(g) | Replace the equal sign (=) with equilibrium arrows $(\rightleftharpoons)$ as indicated and add the missing equilibrium arrows. | Typo |
| Chapter 17 Kinetics: <br> Section 17.7 <br> Catalysis | Figure 17.25 One of the arrows in figure b, says "Ethylene absorbed" where it should say "Ethylene adsorbed" (with a "d" instead of a "b"). I guess it is a typo, but it changes the meaning and can cause misunderstandings. | This figure will be updated. | Typo |
| Chapter 18 Representative Metals, Metalloids, and Nonmetals: Section 18.3 Structure and | In q41 The students are referred to Appendix I instead of to Appendix $G$, which contains the thermodynamic values needed to solve the problem. | This link will be updated to direct to Appendix G. | Typo |
| General Properties of the Metalloids |  |  |  |
| :---: | :---: | :---: | :---: |
| Chapter 19 <br> Transition Metals and Coordination Chemistry: Section 19.2 Coordination Chemistry of Transition Metals | Example 19.5, Figure 19.22 This example is describing cis and trans isomerism in coordination complexes. The trans[ $\mathrm{Pt}(\mathrm{NH} 3) 2 \mathrm{Cl} 2]$ should contain wedges and dashes - equivalent to how the cis isomer is shown in Figure 19.20. If the wedges and dashes are omitted, then the angles should be corrected to 90 degrees. (Wedges and dashes are preferable.) | This figure will be updated. | Typo |
| Chapter 19 <br> Transition Metals and Coordination Chemistry: Section 19.2 Coordination Chemistry of Transition Metals | Table 19.5 [ $\mathrm{Ni}(\mathrm{Cl}) 4] 2-$ is TETRAHEDRAL, not square planar. Cl - is a weak field ligand | Revise "NiCl_4" to "Ni(CN)_4". | Incorrect answer, calculation, or solution |
| Chapter 19 <br> Transition Metals and Coordination Chemistry: Section 19.2 Coordination Chemistry of Transition Metals | In Table 19.1, the last entry reads that $\mathrm{CO}_{2}{ }^{\wedge} 2$ is an anionic ligand with the name carbonato. The number of oxygens in the entry is incorrect. The carbonate ion (which is the carbonato ligand) is $\mathrm{CO}^{\wedge} \mathbf{2 -}^{-}$ | Revise CO_2 to CO_3. | Typo |
| Chapter 19 <br> Transition Metals and Coordination Chemistry: Section 19.2 Coordination Chemistry of Transition Metals | On pg 1051 the text states that "Transition metals often form geometric isomers, in which...etc." The text should read "Coordination complexes" rather than "Transition metals". | Revise "Transition metals often form..." to "Transition metal complexes often exist as..." | Other factual inaccuracy in content |
| Chapter 19 <br> Transition Metals and Coordination Chemistry: Section 19.3 Spectroscopic and Magnetic Properties of Coordination Compounds | In Example 19.9 the frequency is represented as the greek letter nu. In the next line, the frequency is represented as "nu". | Revise "nu" to "v". | Typo |
| Chapter 19 <br> Transition Metals and Coordination Chemistry: Section 19.3 Spectroscopic and Magnetic Properties of Coordination Compounds | In the second line of text, the symbol for the crystal field splitting energy is given. The 'oct' should be subscript. In the spectrochemical series, the caption reads '...in order of increasing field trength...'. It should read '...in order of increasing field strength...' | Revise to set "oct" as subscript. The typo in strength is correct in webview. | Typo |
| Chapter 19 <br> Transition Metals and Coordination | When the Example 19.8 box continues from pg 1063 to pg 1064, the gray box for the example stops before the end of the example. The | Revise the text before the image to "....more complicated pattern, as | Other |
| Chemistry: Section 19.3 Spectroscopic and Magnetic Properties of Coordination Compounds | content from the top of the page until (and not including) the "Magnetic Moments of Molecules and lons" should be part of the example. The figure for the example is missing the ligand locations, and missing the $x$ and $y$ axes labels for the $\mathrm{dz}^{\wedge} 2$ diagram. It appears the authors meant to show where the ligands would be located in this geometry but failed to do so. The text reads "This results in the octahedral t 2 g and the eg sets splitting and gives a more complicated pattern with no simple (delta)oct." This text is ambiguous because it's unclear why the splitting energy would be called (delta)oct rather than (delta)sqp. Furthermore, the "pattern" referred to is this sentence is not the "pattern" referred to in the next sentence that refers to the dorbital diagrams. | depicted below:". The image will also be updated. |  |
| :---: | :---: | :---: | :---: |
| Chapter 20 Nuclear Chemistry: Section 20.1 Nuclear Structure and Stability | Example 20.1 The solution for part (b) of this example compares the density of a neutron star to the density of a U-235 nucleus. The calculations appear correct, however, in summing up the answer by stating the comparison, your text says "These values are fairly similar (same order of magnitude), but the nucleus is more than twice as dense as the neutron star". According to your calculations, it should be the other way around, and it should state .."the neutron star is more than twice as dense than a U-235 nucleus." | Revise the last sentence before "Check Your Learning" to "These values are fairly similar (same order of magnitude), but the neutron star is more than twice as dense as the U-235 nucleus." | Incorrect answer, calculation, or solution |
| Chapter 20 Nuclear Chemistry: Section 20.2 Nuclear Equations | The incorrect table was uploaded to the website for Chapter 20.2.Tables 20.4 and 20.7 are identical on the web, though different in the hard copy and in their table description. Table 20.4 should show the names of the particles, their symbols, etc. This is what the hardcopy of the book and the Figure 20.4 text indicate. However, the table shown is that from section 20.3, Table 20.7 of nuclear equations. | Figure 20.4 will be updated. | Incorrect answer, calculation, or solution |
| Chapter 20 Nuclear Chemistry: Section 20.2 Nuclear Equations | In figure 20.7 none of the --> arrows are present in the print version. <br> In the online version this has been partially corrected. Unfortunately, for Electron capture the arrow has been incorrectly placed. Electron capture should have the electron on the reactant side of the equation. As is, it is whowing Beta decay. | This figure will be updated. | Typo |
| Chapter 20 Nuclear Chemistry: Section 20.3 Radioactive Decay | At the Dead Sea Scrolls, the calculations are correct, but the equation is not, due to the fact that it should be $\ln ((N O / N(t))$ instead of $\ln$ $(\mathrm{N}(\mathrm{t}) / \mathrm{NO})$. In terms of answer, the right equation was taken into account, but it was written wrong. Please accept my humble opinion | While our reviewers determined there is no error with the equation or the computed result, a revision is needed to the related text narrative. Revise "-kt" to "- $\lambda t$ ". | General/pedago gical suggestion or question |
| Chapter 20 Nuclear Chemistry: Section 20.3 Radioactive Decay | Figure 20.7 In the second column of the table all of the arrows are missing. In addition, in the bottom two rows the species " Y " is listed on the right hand side with mass number A and charge $\mathrm{Y}-1$. The charge should be $\mathrm{Z}-1$ rather than Y -1. | This figure will be updated. | Typo |
| :---: | :---: | :---: | :---: |
| Chapter 20 Nuclear Chemistry: Section 20.4 Transmutation and Nuclear Energy | On page 1151, I find this phrase: <br> "Plutonium is now mostly formed in nuclear reactors as a byproduct during the decay of uranium. Some of the neutrons that are released during U-235 decay combine with U238 nuclei to form uranium-239...." <br> In fact, neutrons are released during fission of uranium, not during radioactive decay. I suggest replacing both instances of the word "decay" with "fission". | Revise the paragraph beginning "Plutonium is now mostly formed..." to "Plutonium is now mostly formed in nuclear reactors as a byproduct during the fission of U-235. <br> Additional neutrons are released during this fission process (see the next section), some of which combine with U-238 nuclei to form uranium-239; this undergoes $\beta$ decay to form neptunium-239, which in turn undergoes $\beta$ decay to form plutonium-239 as illustrated in the preceding three equations. These processes are summarized in the equation:". | Other factual inaccuracy in content |
| Chapter 21 Organic Chemistry: Section 21.1 Hydrocarbons | 11 g exercise. Incorrect answer key 4-methyl-1pentene instead of 5-methyl-1-pentene | Revise " 5 -methyl-1-pentene" to "4-methyl-1-pentene". | Incorrect answer, calculation, or solution |
| Chapter 21 Organic Chemistry: Section 21.3 Aldehydes, Ketones, Carboxylic Acids, and Esters | Figure 21.15 The structures for butyl acetate and propyl isobutyrate are not correct (the chains are switched around to make methyl butyrate and isobutyl propionate, respectively). | This figure will be updated. | Other factual inaccuracy in content |
| Chapter 21 Organic Chemistry: Section 21.4 Amines and Amides | Problem 57 Geometry about the nitrogen in pyridine is bent, not trigonal planar. Or you could ask what the electron geometry is in the problem. | Revise the second sentence in the question stem to "What are the hybridizations, electron domain geometries, and molecular geometries about the nitrogen atoms in pyridine and in the pyridinium ion?" The figure in the answer will also be updated. | Incorrect answer, calculation, or solution |
| Throughout | 141 Efina <br> 4011 atm used as standard ? <br> 409 Fig.8.11 Celsius to Kelvin conversion <br> 441 Fig. 8.32 axis label: 'speed u' instead of 'velocity v', peak at Up, not Vp. <br> 442 Fig. 8.33 axis label: 'speed $u$ ' instead of 'velocity v'. <br> 443 Fig. 8.34 axis label: 'speed u' instead of 'velocity v'. <br> 466 Fig. 9.4 Hot/cold liquid rather than water due to anomalies <br> 471 Tfina <br> 472 2x Tfina <br> 505 Example $9.162 x$ units of kJ missing 510 Tfina <br> 583 Fig. 10.64 for destructive interference case, rays of diffracted beam are in phase. <br> 670 Table 12.2 standard at 1 atm ? <br> 671 Example 12.6 units of J K-1 mol-1 missing 675 Key Equations last two: Q and K not in this chapter 12 <br> 736 Ch. 14.1 Arrhenius acid is based on work by Swante Arrhenius, not Carl Axel Arrhenius <br> 742 Fig. 14.2 caption [H3O+] instead of H3O+ <br> 757 Example 14.12 table in 'step 1': should be col 1 for HCO 2 H , col 2 for $\mathrm{H} 3 \mathrm{O}+$, col 3 for HCO2-. <br> 757 Example 14.122 times $\times 2+$ instead of $\times 2$ <br> 787 Fig. 14.21 caption (b) should be acetic acid, not HCl <br> 814 Example 15.4 answer for molar solubility of $\mathrm{Ca}(\mathrm{OH}) 2$ should be $6.9 \times 10-3 \mathrm{M}$, not $1.3 \times 10-2$ M. <br> 872 Example 16.6 should read Ecell $=-0.14 \mathrm{~V}$, not Ecell $=-0.014 \mathrm{~V}$ <br> 895 prob. 24(b) should be $3 \mathrm{Cu}(\mathrm{s})$, not $2 \mathrm{Cu}(\mathrm{s})$ <br> 915 third rxn scheme must be $2 \mathrm{~N} 2 \mathrm{O} 5 \rightarrow 4 \mathrm{NO} 2$ <br> + O 2 to balance. <br> 958 Exercise 46 should read ' $918 \mathrm{~F} \rightarrow 8180+$ e+', not '5189F $\rightarrow 1880$ + e-' <br> 1114 Figure 20.4 beta particle should be '-10e', not '10e' <br> 1121 first-order equ. should read 'Nt = NO e- $\lambda \mathrm{t}$ ', not 'Nt = NO e-kt ' <br> 1122 Example 20.5 (b) exponent should read (15 y), not (15/y) <br> 1123 carbon-14 decay should read '614C <br> $\rightarrow 714 \mathrm{~N}+-10 \mathrm{e}^{\prime}$, not ' $614 \mathrm{C} \rightarrow 712 \mathrm{~N}+-10 \mathrm{e}^{\prime}$ <br> 1129 first rxn scheme twice $t 1 / 2$ misplaced <br> 1140 end of page should read ' $2+10$ e' or ' 2 <br> $+10 \beta$ ', not ' $2+10$ ' <br> 1141 line 5 should read ', 12 H and triton 13 H ', <br> not ', 12 and triton 13' <br> 1161 Exercise 17 should read '919 F', not '919 F' <br> 1247 Appendix G HgSO4 : S298 = 0 ? <br> 1321 Ch.16: 5d should be MnO4-, not MnO4. <br> 1336 Sol. 13 b) should read '24 He', not He2' | Some of these revisions were implemented in the 2 e update. The remaining will be updated accordingly. | Typo |
| :---: | :---: | :---: | :---: |
| Appendix B \| <br> Essential <br> Mathematics | In the problem of the example B2 ( $3.6 \times 10^{\wedge} 5$ by $6.0 \times 10^{\wedge}-4$ ), the minus sign of the power of ten in the numerator is missing. <br> Correction: ( $3.6 \times 10^{\wedge}-5$ by $\left.6.0 \times 10^{\wedge}-4\right)$. | Our reviewers accepted this change. | Typo |
| :---: | :---: | :---: | :---: |
| Appendix B \| <br> Essential <br> Mathematics | In explaining the usefulness of the exponential notation, just before the last line at page 1199, it's written "every large" instead of "very large". | Our reviewers accepted this change. | Typo |
| Appendix B \| <br> Essential <br> Mathematics | At "EXAMPLE B4": <br> It says "Divide $3.6 \times 10^{\wedge} 5$ ", but it continues as if it said "... 10^-5" | Revise " 5 " to " 5 " | Incorrect answer, calculation, or solution |
| Appendix G \| <br> Standard <br> Thermodynamic <br> Properties for <br> Selected Substances | The Delta G of formation for liquid sulfuric acid, H2SO4 (I), is listed as positive $690.00 \mathrm{~kJ} / \mathrm{mol}$, but it should be negative, as in $-690.00 \mathrm{~kJ} / \mathrm{mol}$. | Revise " 690.00 " to "-690.00". | Typo |
| Appendix G \| <br> Standard <br> Thermodynamic <br> Properties for <br> Selected Substances | entry for HNO3 The phase is not listed. It should be HNO3(aq). This value should be listed with the compounds of nitrogen after HNO3(I) and HNO3(g). Example 5.15 is the first example using standard enthalpy of formation and Appendix G. It is important to have these values listed clearly and with their phase. | A new row for HNO3(aq) will be added to the table. | Typo |

