

Spectroscopy & Structure Practice

INSTRUCTOR ANSWER KEY — COMPLETE EXPLANATIONS

Directions: Answers are in red. Key includes a concise rationale for each item.

1. A strong IR absorption near 1715 cm^{-1} is consistent with:

- A. C=O stretch in a simple saturated ketone**
- B. $\text{C}\equiv\text{N}$ stretch
- C. O–H stretch only
- D. C–H bending only

Reasoning: Saturated acyclic ketones commonly show a strong carbonyl stretch near 1715 cm^{-1} .

2. A broad IR band spanning roughly $3200\text{--}3600\text{ cm}^{-1}$ often indicates:

- A. Alcohol O–H**
- B. Nitrile $\text{C}\equiv\text{N}$
- C. Alkyne $\text{C}\equiv\text{C}$ only
- D. Carbonyl $\text{C}=\text{O}$

Reasoning: Hydrogen-bonded alcohol O–H stretching typically produces a broad absorption in this region.

3. A carboxylic acid O–H stretch is often characterized by:

- A. Sharp peak at 2250 cm^{-1}
- B. Very broad absorption about $2500\text{--}3300\text{ cm}^{-1}$**
- C. Single sharp peak at 1600 cm^{-1}
- D. No IR absorption

Reasoning: Carboxylic acids frequently exhibit a very broad O–H envelope overlapping C–H stretching.

4. A relatively sharp IR absorption around 2250 cm^{-1} can indicate:

- A. $\text{C}\equiv\text{N}$**
- B. C–O single bond
- C. Alcohol O–H
- D. Alkane C–H

Reasoning: Nitrile $\text{C}\equiv\text{N}$ stretching occurs near 2250 cm^{-1} , although exact values vary.

5. An aldehydic ^1H NMR signal commonly appears near:

- A. 0–1 ppm
- B. 2–3 ppm
- C. 9–10 ppm**
- D. 14–16 ppm

Reasoning: The proton directly bonded to a carbonyl carbon in an aldehyde is strongly deshielded.

6. An aromatic proton often resonates in approximately which ^1H NMR region?

- A. 0–1 ppm
- B. 6.5–8.5 ppm**
- C. 12–16 ppm
- D. 18–20 ppm

Reasoning: Many aromatic C–H signals lie around 6.5–8.5 ppm, depending on substituents.

7. An ethyl fragment often shows what integration and splitting pattern in first-order ^1H NMR?

- A. 3H triplet and 2H quartet**
- B. 3H singlet and 3H singlet
- C. 1H septet and 6H doublet
- D. 2H singlet and 2H singlet

Reasoning: An isolated $\text{CH}_3\text{--CH}_2\text{--}$ unit frequently gives a methyl triplet and a methylene quartet.

8. A 6H doublet plus 1H septet can support which fragment?

- A. Methoxy group
- B. Isopropyl group**
- C. Ethene
- D. Aldehyde

Reasoning: Two equivalent methyl groups coupled to one methine proton commonly show 6H doublet + 1H septet.

9. A TLC spot travels 2.4 cm while the solvent front travels 6.0 cm. R_f = ?

- A. 0.25
- B. 0.40**
- C. 2.5
- D. 3.6

Reasoning: $R_f = \text{spot travel} / \text{solvent-front travel} = 2.4 / 6.0 = 0.40$.

10. In simple acetone, the two methyl groups give what expected ^1H NMR pattern?

- A. Two 3H doublets
- B. One 6H singlet**
- C. One 3H triplet
- D. One 1H aldehyde singlet

Reasoning: The two equivalent methyl groups next to the carbonyl have no vicinal hydrogen splitting and give one 6H singlet.

11. A curved arrow in an organic mechanism starts at:

- A. The electron source**

- B. The atom receiving electrons only
- C. The positive charge always
- D. The reaction arrow

Reasoning: The tail of a curved arrow points to the electron pair (bond or lone pair) being moved.

12. Lower pKa generally indicates:

- A. Stronger acid**
- B. Weaker acid
- C. Stronger base only
- D. No connection to acidity

Reasoning: For comparable conditions, lower pKa corresponds to larger Ka and stronger acidity.

13. Which is a Lewis base?

- A. Electron-pair donor**
- B. Electron-pair acceptor
- C. Only a proton donor
- D. A compound with no electrons

Reasoning: A Lewis base donates an electron pair to form a bond.

14. What is the degree of unsaturation for C₆H₁₂O (neutral closed-shell)?

- A. 0
- B. 1**
- C. 2
- D. 3

Reasoning: $DBE = (2C + 2 - H)/2 = (12 + 2 - 12)/2 = 1$; oxygen is ignored.

15. What is the degree of unsaturation for C₆H₆?

- A. 2
- B. 3
- C. 4**
- D. 5

Reasoning: $DBE = (2 \times 6 + 2 - 6)/2 = 4$.