# Exam. Code : 107405 <br> Subject Code : 1860 

## B.Sc. Biotechnology $5^{\text {th }}$ Semester PHYSICAL, ORGANIC \& INORGANIC ASPECTS OF SPECTROSCOPY-A

## Paper-BT-7

## Time Allowed-Three Hours] [Maximum Marks-40

## SECTION-A

Note :-ALL questions in this section are compulsory and each question is of $\mathbf{1}$ mark.

1. Calculate the frequency of radiations having an energy of $1.6 \times 10^{-19}$ Joules.
2. Using IR spectroscopy how will you distinguish between an aliphatic aldehyde and an aliphatic ketone ? 1
3. Why is absorption and not emission spectroscopy used to study the spectra of organic compounds ? 1
4. What is the effect of hydrogen bonding on ultraviolet absorption ?
5. What is force-constant? $\quad 1$
6. Using IR spectroscopy how will you distinguish between a cis olefin and a trans olefin ?

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7. What is fluorescence ? How it is different from Phosphorescence ?
8. Describe the effect of ring size on the IR absorption of carbonyl group of cyclo-ketones.

## SECTION-B

Note :-Attempt any FIVE questions from this section. Each question is of 4 marks.
9. How many fundamental vibrational frequencies would you expect to observe in the IR spectrum of $\mathrm{CO}_{2}$ ?

## 10. Distinguish between the following pairs of compounds

 with the help of IR spectroscopy :(i) Propanol from Propanone 1
(ii) Ethanol from Ethyl amine 1
(iii) Aniline from N -methyl aniline 1
(iv) Acetone from acetylene. 1
11. What is Born-Oppenheimer approximation? How it is different from Frank-Condon principle ? 4
12. Several ionic crystals such as KBr and NaCl etc. are transparent to ultraviolet regions, but are never used in these regions. Why ?

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13. Aniline absorbs at $280 \mathrm{~nm}\left(\varepsilon_{\max } 8600\right)$ but in acidic solution the main absorption band is seen at 203 nm ( $\varepsilon_{\max } 7500$ ) which is comparable to benzene. Explain.
14. (i) Why water can not be used as a solvent in IR spectroscopy ?2
(ii) Why methyl alcohol is a good solvent for UV and not for IR determination? 2
15. What type of transitions are observed in case of $\alpha, \beta$-unsaturated carbonyl compounds ? How absorption maximum and intensity are shifted when carbonyl group is not conjugated ?
16. What do you mean by a good solvent in UV spectroscopy ? What is the effect of solvent on absorption maximum in case of $\pi$ to $\pi^{*}$ transitions of conjugate olefins as well as conjugated carBonyl compounds ?

## SECTION-C

Note :-Do any TWO questions from this section. Each question is of $\mathbf{6}$ marks.
17. A conjugated diene absorbs at a higher wavelength with higher value of extinction co-efficient as compared to a diene in which double bonds are isolated. Explain giving examples and the chemistry involved.

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18. Using Woodward Fieser rules calculate the $\lambda_{\max }$ for the following :




$1.5+1.5+1.5+1.5$
19. Discuss at least three types of groups for which the study of fingerprint region is most essential in IR spectroscopy.
20. (i) Discuss with examples the inductive and the mesomeric effects influencing the carbonyl absorption frequency.
(ii) Write the expected IR peaks for p-nitrophenol and p-nitrobenzoic acid.
