

STRUCTURAL DETERMINATION – PART 1B

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I. Definitions

Wave-particle duality (S3) Quantum (S4) Infrared spectroscopy (S5-7) Ultraviolet spectroscopy (S23-27)
Absorbance and molar absorptivity (S33) β -Carotene (S30) Vitamer (S31) Visual cycle (S31-32)

II. Infrared spectroscopy

A. The result: infrared spectrum (S7)

B. Properties determined in infrared spectroscopy (S5-7)

How are wavelength and frequency of electromagnetic radiations related to each other? (S3)

$$\text{Speed} - C = \lambda \cdot \nu = 3.0 \times 10^8 \text{ (m/sec)}$$

What is the energy of a photon? (S4)

$$\epsilon = h \cdot \nu \quad (\text{Planck constant: } h = 6.6 \times 10^{-34} \text{ J}\cdot\text{s})$$

What is wavenumber? (S5)

$$\tilde{\nu} = 1/\lambda = 33 \cdot \nu / 10^{12} \text{ (cm}^{-1}\text{)}$$

C. Interpretation of infrared spectra (S7-21)

IV. Ultraviolet spectroscopy

A. The result: ultraviolet spectrum (S23)

B. Properties determined in ultraviolet spectroscopy (S24-29)

C. Interpretation of ultraviolet spectra (S26-29)

What compounds have a peak in its UV spectrum (200 - 400 nm)?

D. Quantitative analyses based on ultraviolet spectroscopy (S33)

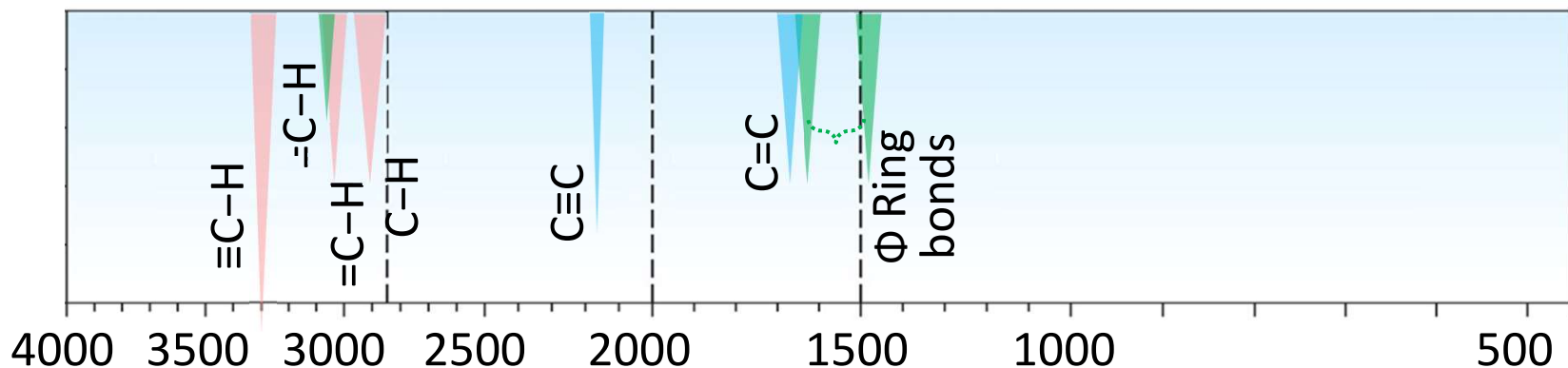
What is absorbance?

$$A = \log(I_0/I)$$

What is molar absorptivity?

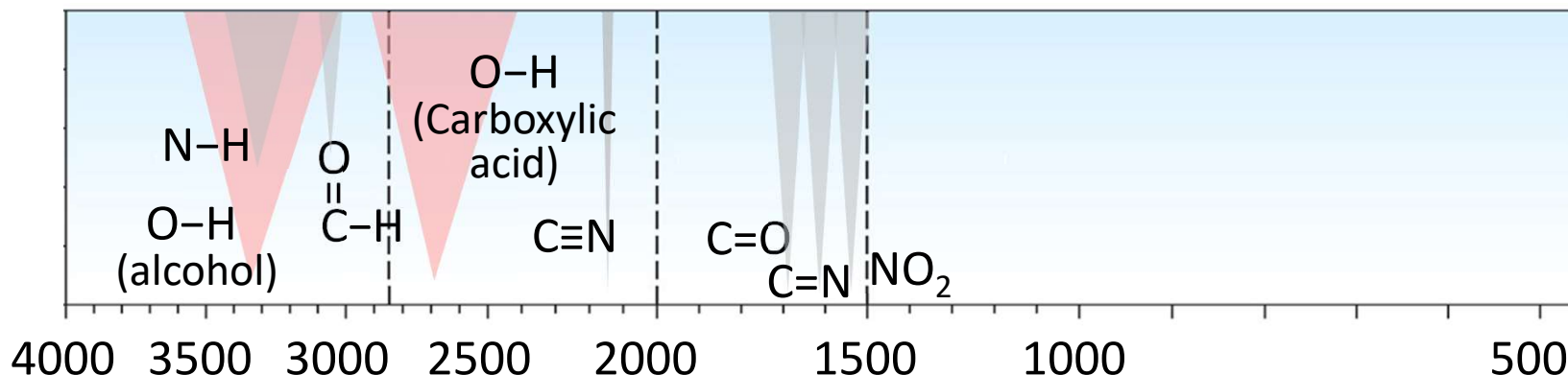
$$\epsilon = A/cL$$

Infrared absorption frequencies of hydrocarbons



Bond	Class	Frequency, cm^{-1}	Signal	Bond	Class	Frequency (cm^{-1})	Signal
C-H	Alkane	2850-2960	M	$\equiv\text{C-H}$	Alkyne	3300	S
$=\text{C-H}$	Alkene	3020-3100	M	$\text{C}\equiv\text{C}$ ³	Alkyne	2100-2260	M
$\text{C}=\text{C}$ ²	Alkene	1640-1680	M	$=\text{C-H}$	Arene	3030	W
W: weak, M: medium, S: strong				C Ring bonds ^{1 1/2}	Arene	1450-1600	M

Infrared absorption frequencies of oxygen-containing substances



Bond	Functional class	Frequency cm^{-1}	Signal	Bond	Class	Frequency cm^{-1}	Signal
O-H	Alcohol	3400-3650	S&B	N-H	Amine	3300-3500	M
O-H	Carboxylic acid	2500-3100	S&B	C=N	Imine	1610-1690	S
=C-H	Aldehyde	3020-3100	M	C≡N	Nitrile	2210-2260	M
C=O	Aldehyde, ketone, ester, amide and carboxylic acid	1670-1780	S	NO ₂	Nitro	1540	S
				W: weak, M: medium, S: strong			

Structural determination – 1(B)

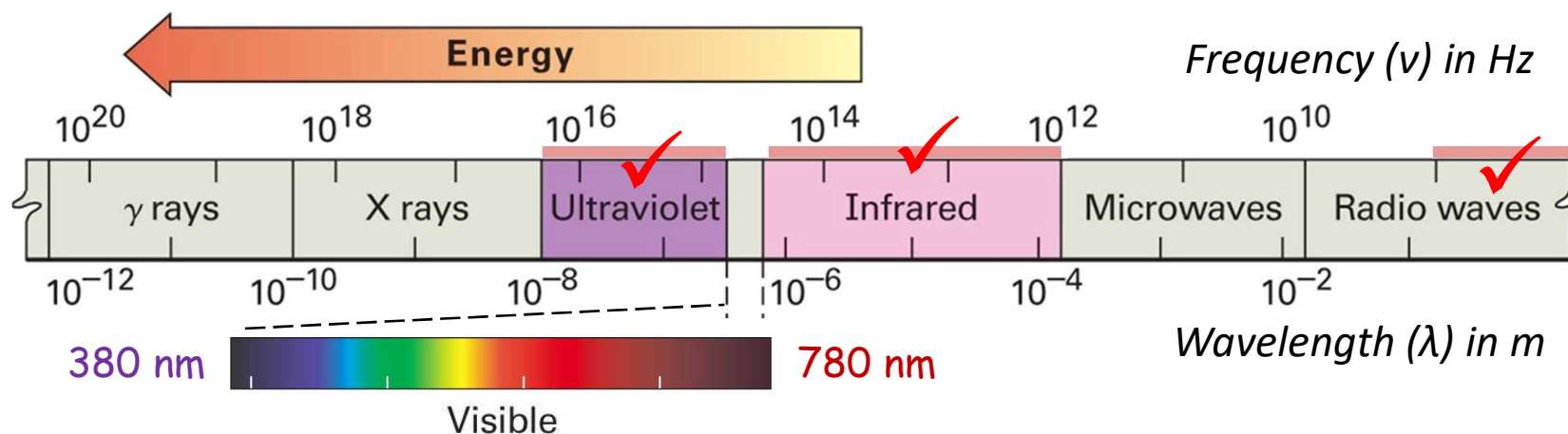
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Biomedical Industrial Ph.D. Program
Biomedical EMBA Program**

What properties are examined in IR?

Spectroscopic analyses of organic compounds

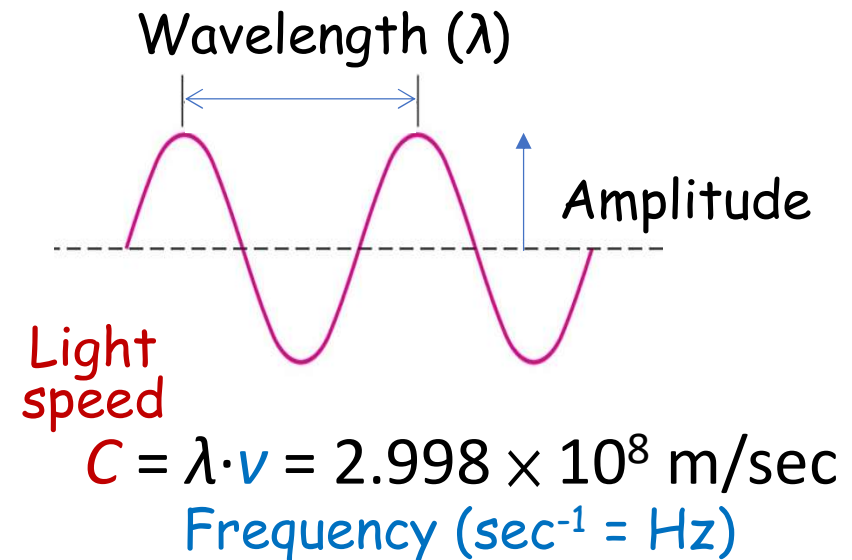
Technique	Electromagnetic radiation	Structure
Infrared spectrometry	Infrared light	Covalent bonds
Ultraviolet spectroscopy	Ultraviolet light	$\sigma-\sigma^*$, $n-\sigma^*$, $\pi-\pi^*$, $n-\pi^*$ transitions
Nuclear magnetic resonance spectroscopy	Radio wave	Atomic nuclei in magnetic field



Properties of electromagnetic radiation – Wave-particle duality

Electromagnetic radiation has dual properties.

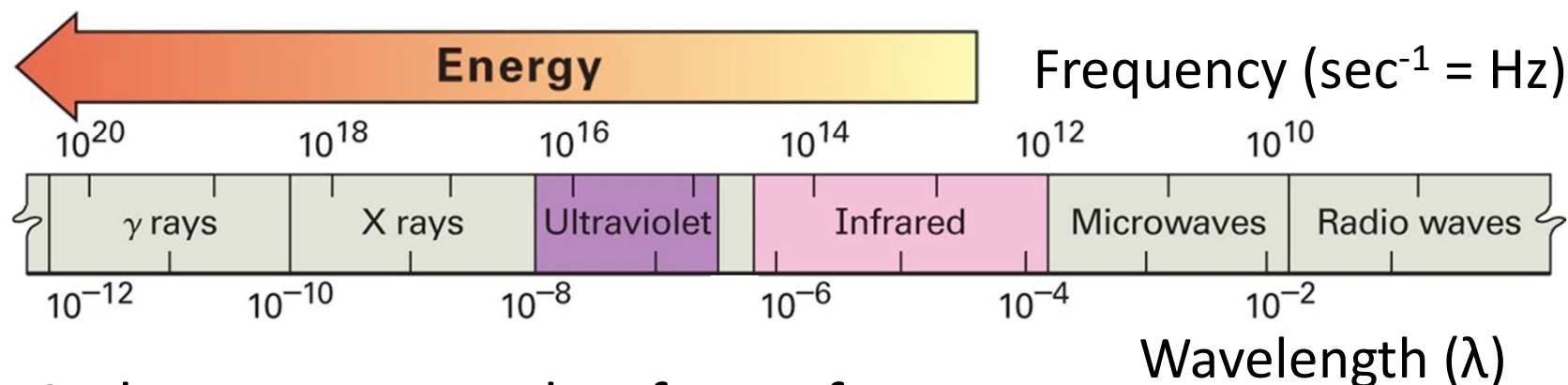
- ✓ **Wave** properties - In homogeneous media, electromagnetic radiation is a transverse wave, meaning that its **oscillations** are **perpendicular** to the direction of energy transfer and travel



- ✓ **Particle** properties - The particle of light has the name **photon**, like other particles such as the electron and proton. Thus, a photon is a single **quantum** of light

Quantum (plural: quanta)

The minimum amount of any physical property involved in an interaction. This means that the magnitude of the physical property can take on only discrete values consisting of **integer multiples** of one quantum.



A photon, or any other form of electromagnetic radiation, has the quantum of **energy** (ϵ) that is in direct proportional to its **frequency** (ν)

$$\epsilon = h \cdot \nu$$

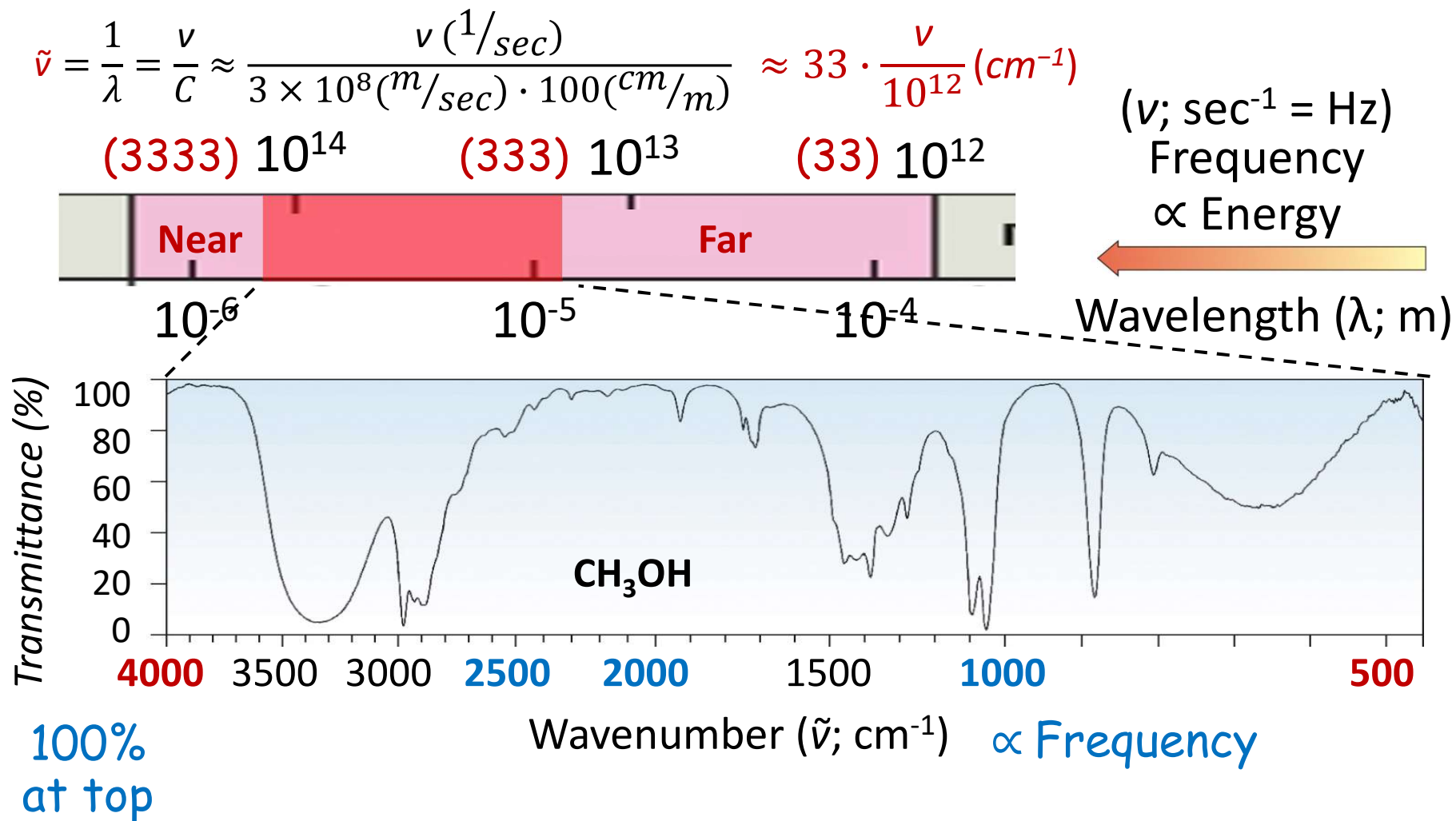
Planck constant

$$6.62607015 \times 10^{-34} \text{ J}\cdot\text{s}$$

$$E = N_A \times \epsilon = 4 \times 10^{-10} \cdot \nu \text{ (J}\cdot\text{s/mol)}$$



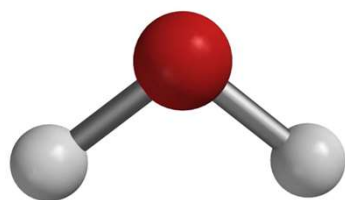
Infrared spectroscopy



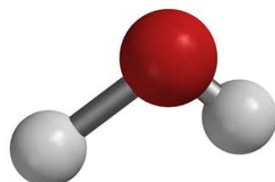
What structures absorb the energy in IR?

The absorption of infrared quanta causes vibration of covalent bonds

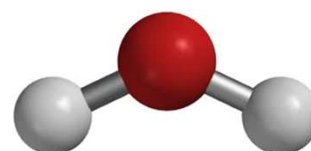
Upon absorption of the infrared radiation of a specific frequency, molecules **stretch** or **bend** if the frequency of the radiation matches the frequency of the **vibration**



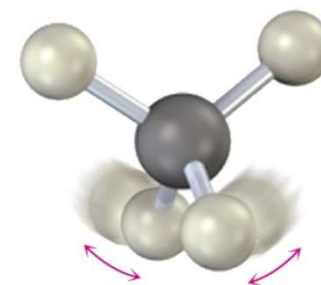
Symmetric stretching



Antisymmetric stretching



In-plane bending

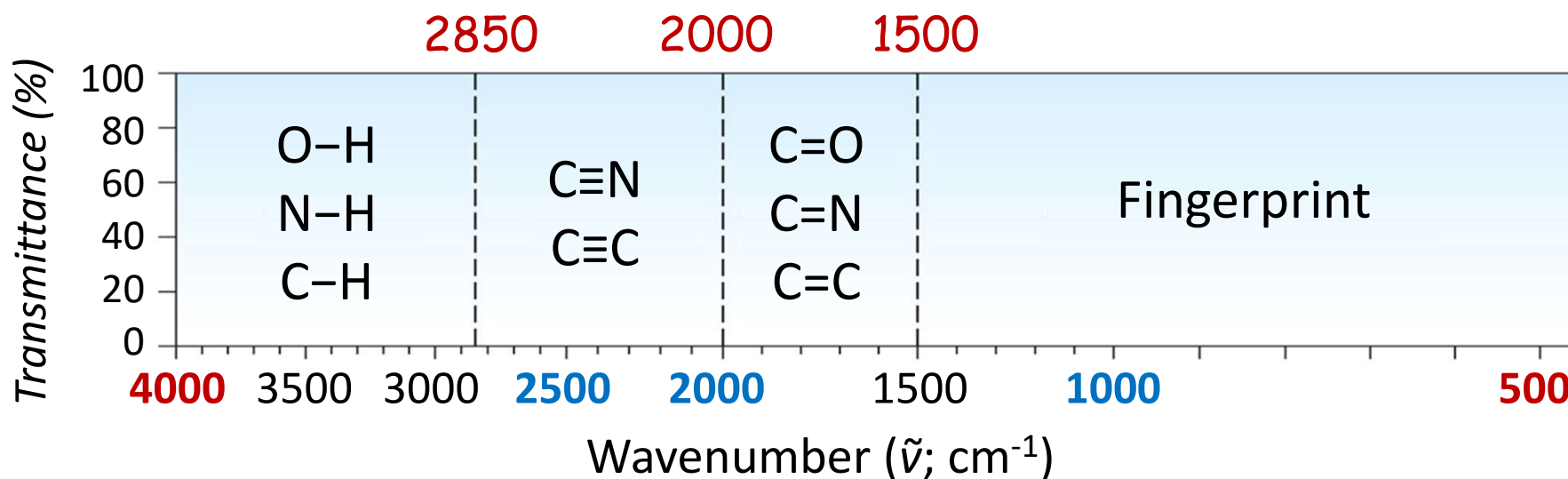


Out-of-plane bending

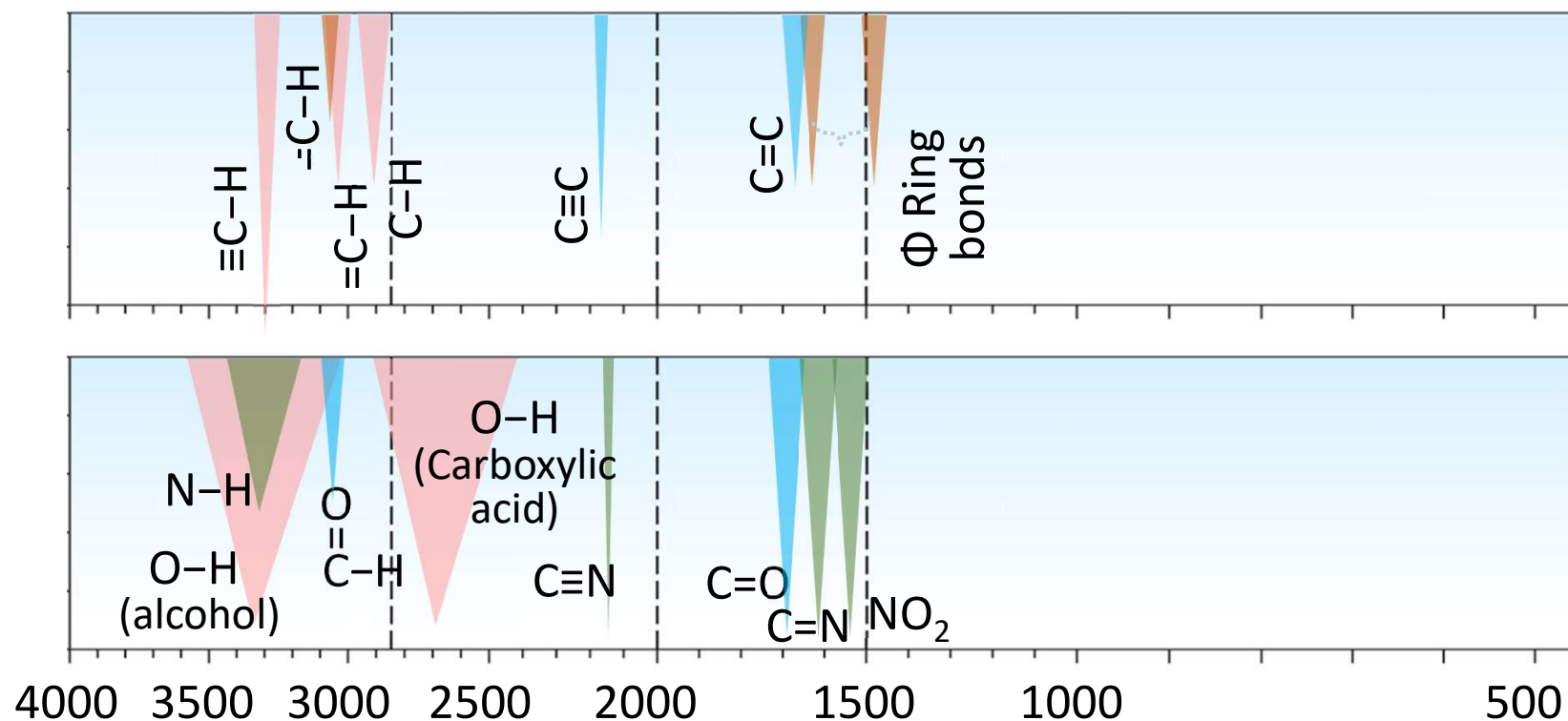
How do we interpret the IR spectra?

Interpret infrared spectra

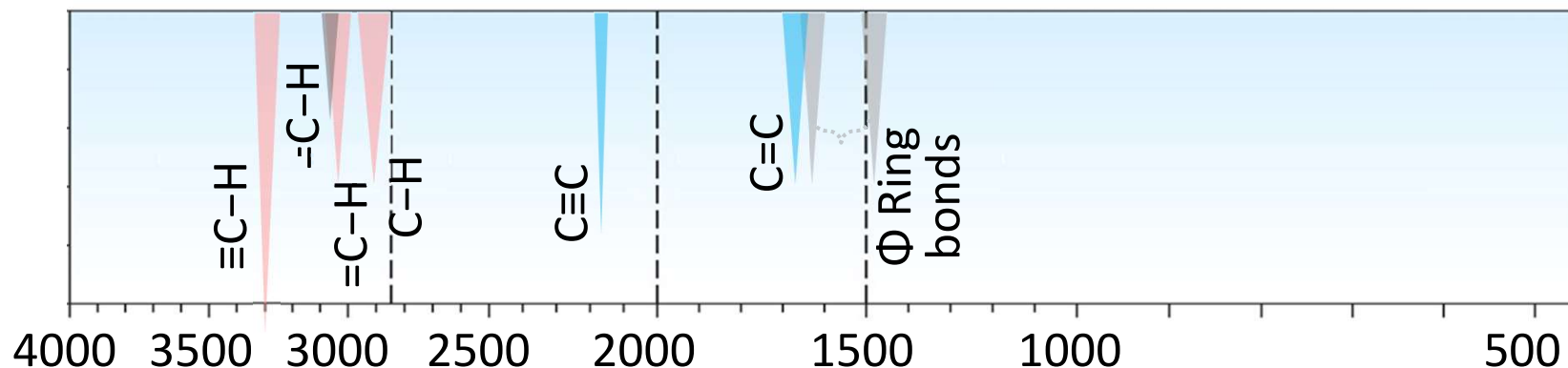
Most functional groups have characteristic *IR* absorption bands that do not change from one compound to another



Infrared absorption frequencies in organic chemistry

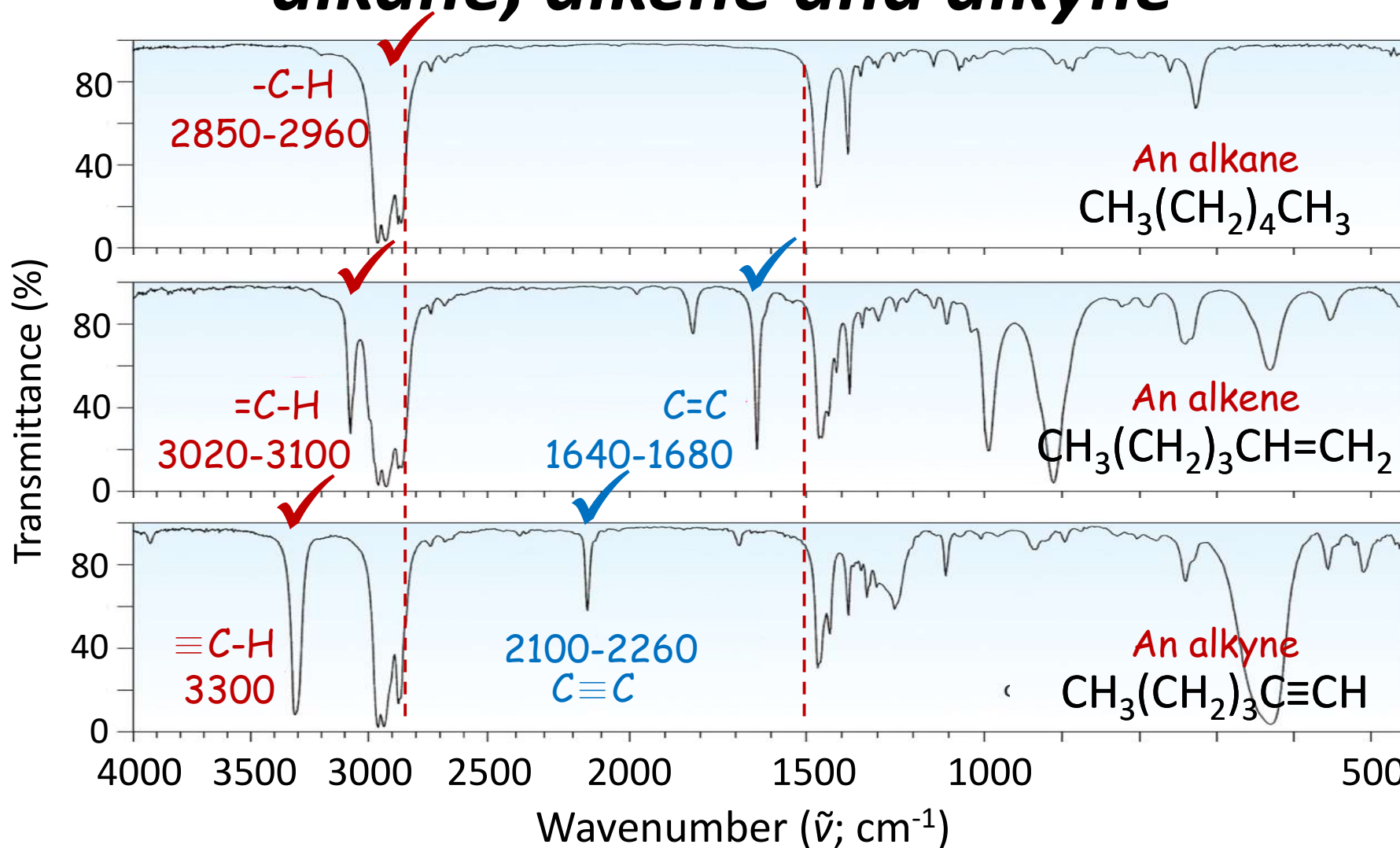


Infrared absorption frequencies of hydrocarbons

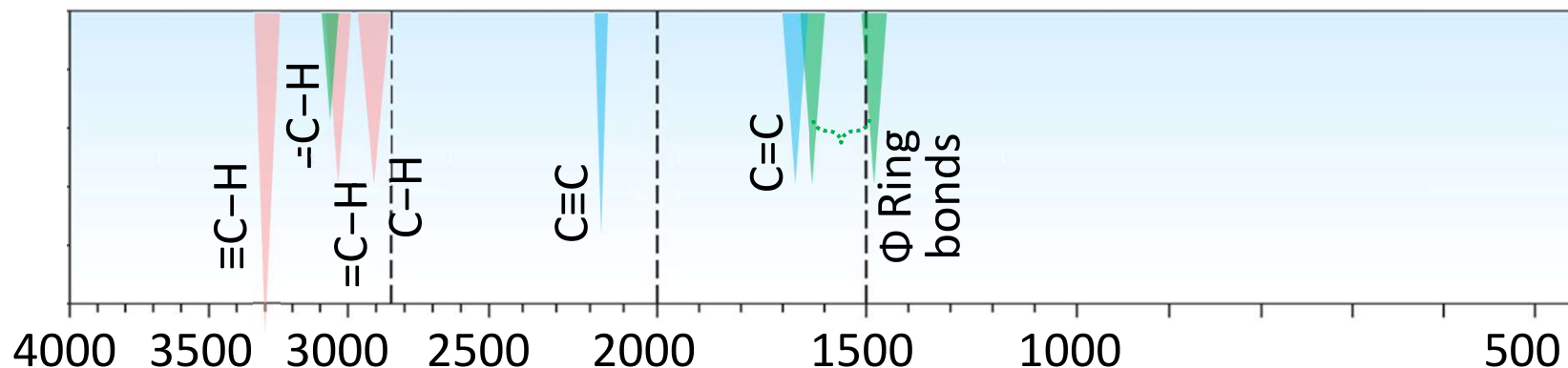


Group	Class	Frequency, cm^{-1}	Signal	Group	Class	Frequency (cm^{-1})	Signal
C-H	Alkane	2850-2960	M	$\equiv\text{C-H}$	Alkyne	3300	S
$=\text{C-H}$	Alkene	3020-3100	M	$\text{C}\equiv\text{C}$	Alkyne	2100-2260	M
\checkmark C=C 2	Alkene	1640-1680	M	$=\text{C-H}$	Arene	3030	W
W: weak, M: medium, S: strong				Φ Ring bonds	Arene	1450-1600	M

IR characteristics in alkane, alkene and alkyne

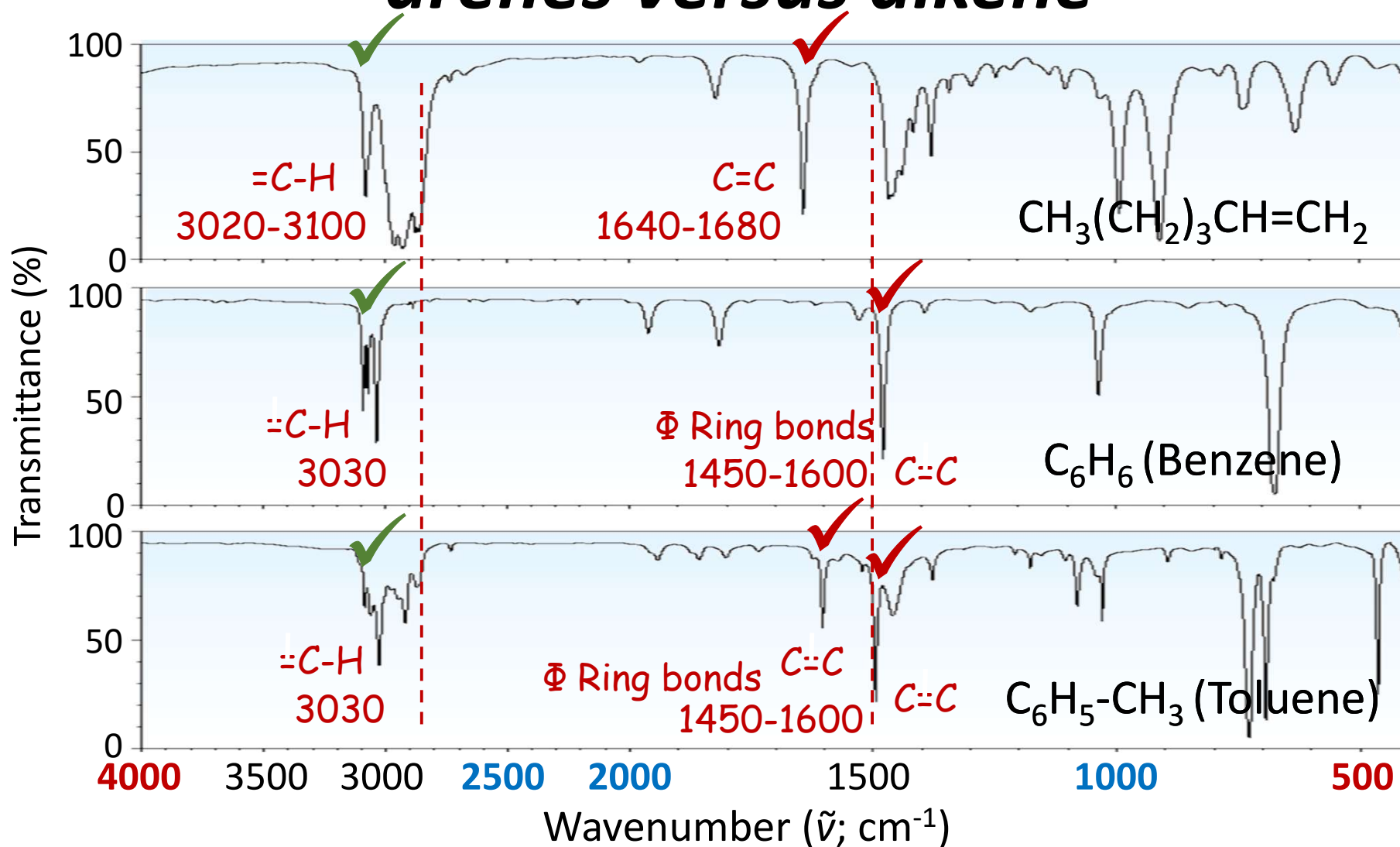


Infrared absorption frequencies of hydrocarbons

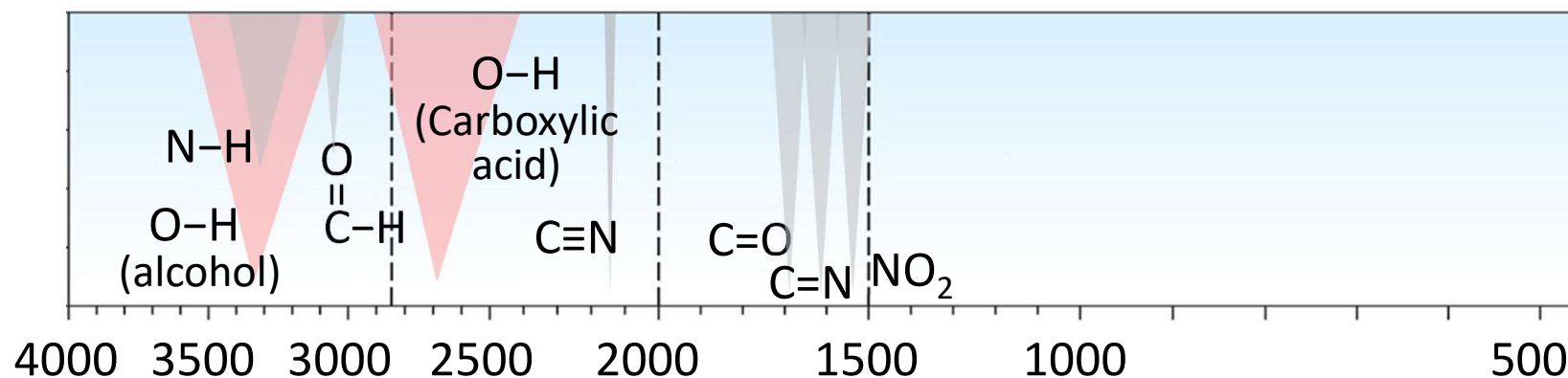


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IR characteristics in arenes versus alkene



Infrared absorption frequencies of oxygen-containing substances

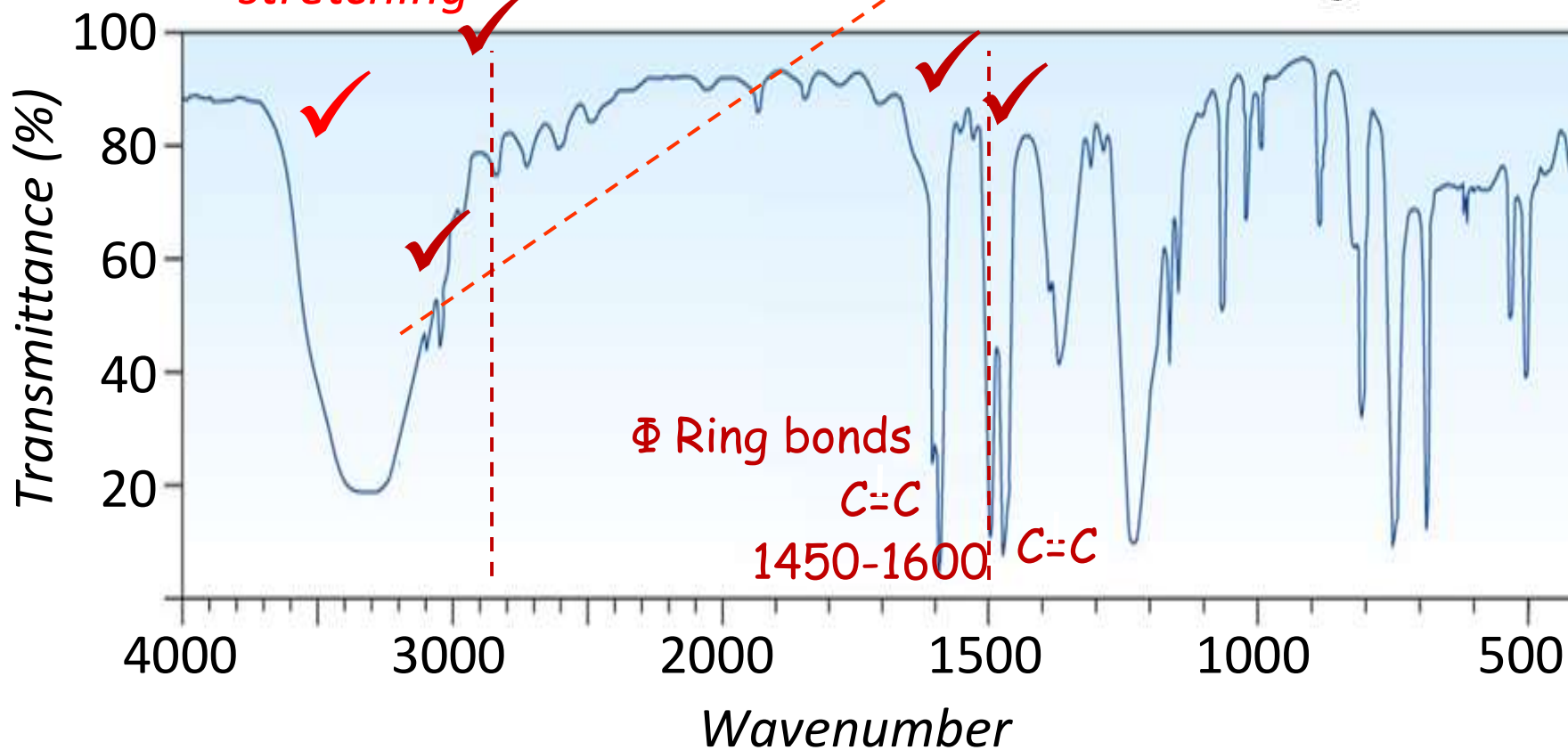
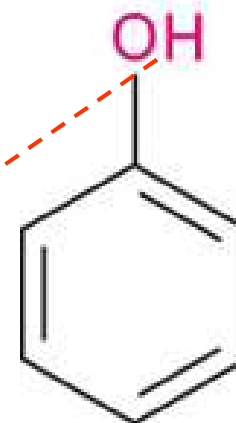


Bond	Functional class	Frequency cm^{-1}	Signal	Bond	Class	Frequency cm^{-1}	Signal
O-H	Alcohol	3400-3650	S&B	N-H	Amine	3300-3500	M
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C=O	Aldehyde, ketone, ester, amide and carboxylic acid	1670-1780	S	NO ₂	Nitro	1540	S
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Infrared spectroscopy – phenol

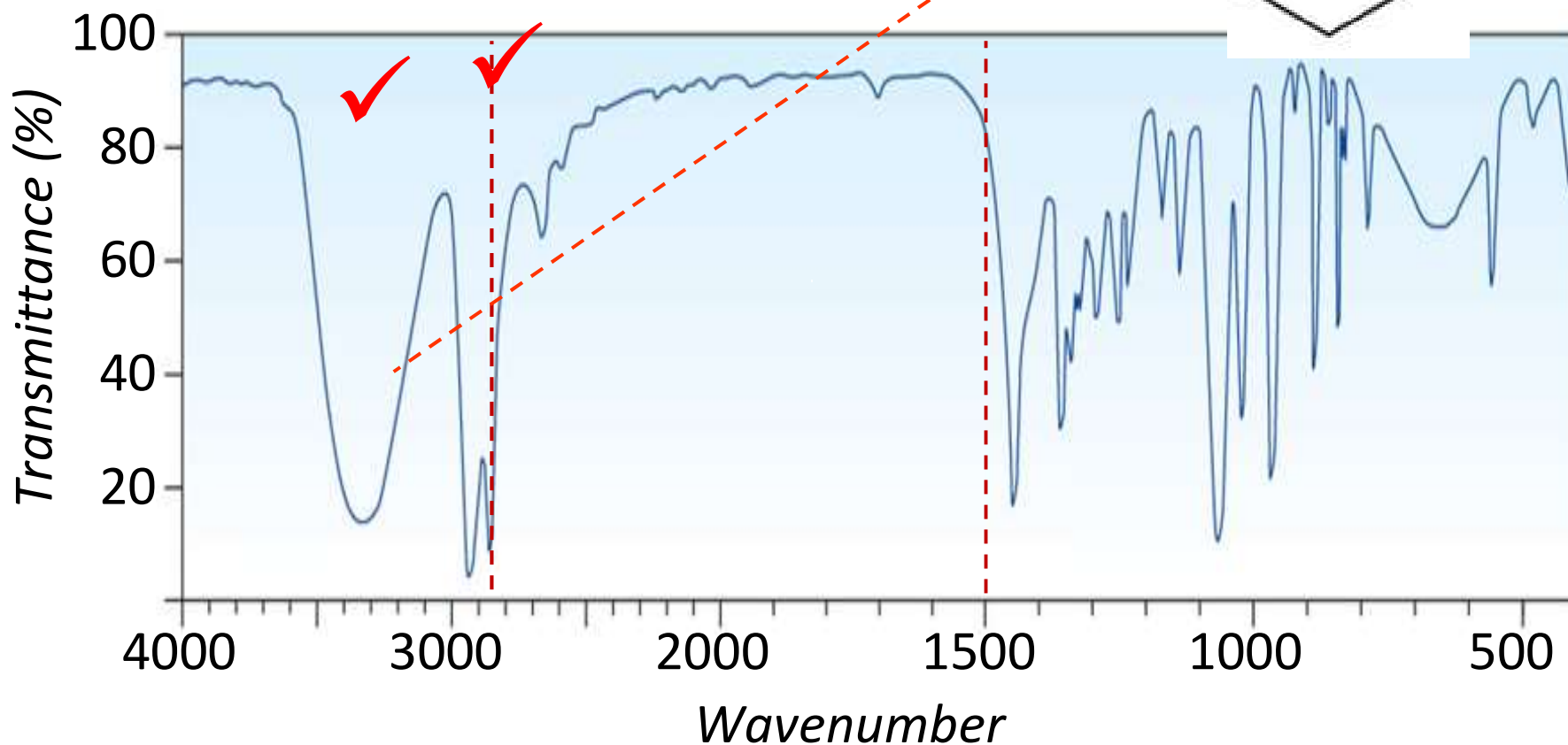
3400~3650 cm^{-1}
(Strong, broad)

H-O
stretching

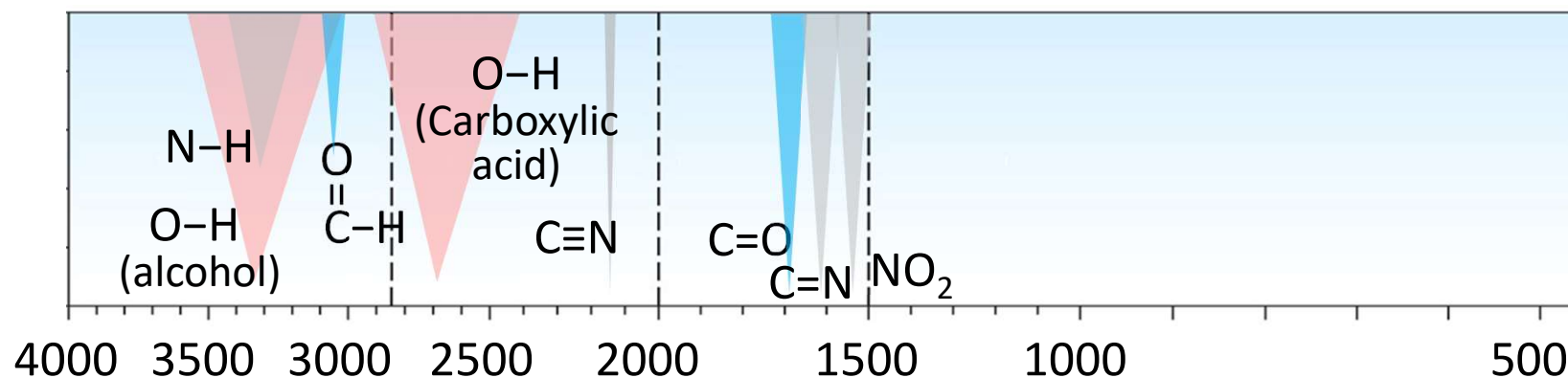


Infrared spectroscopy

H-O stretching – alcohol
3400~3650 cm⁻¹
(Strong, broad)

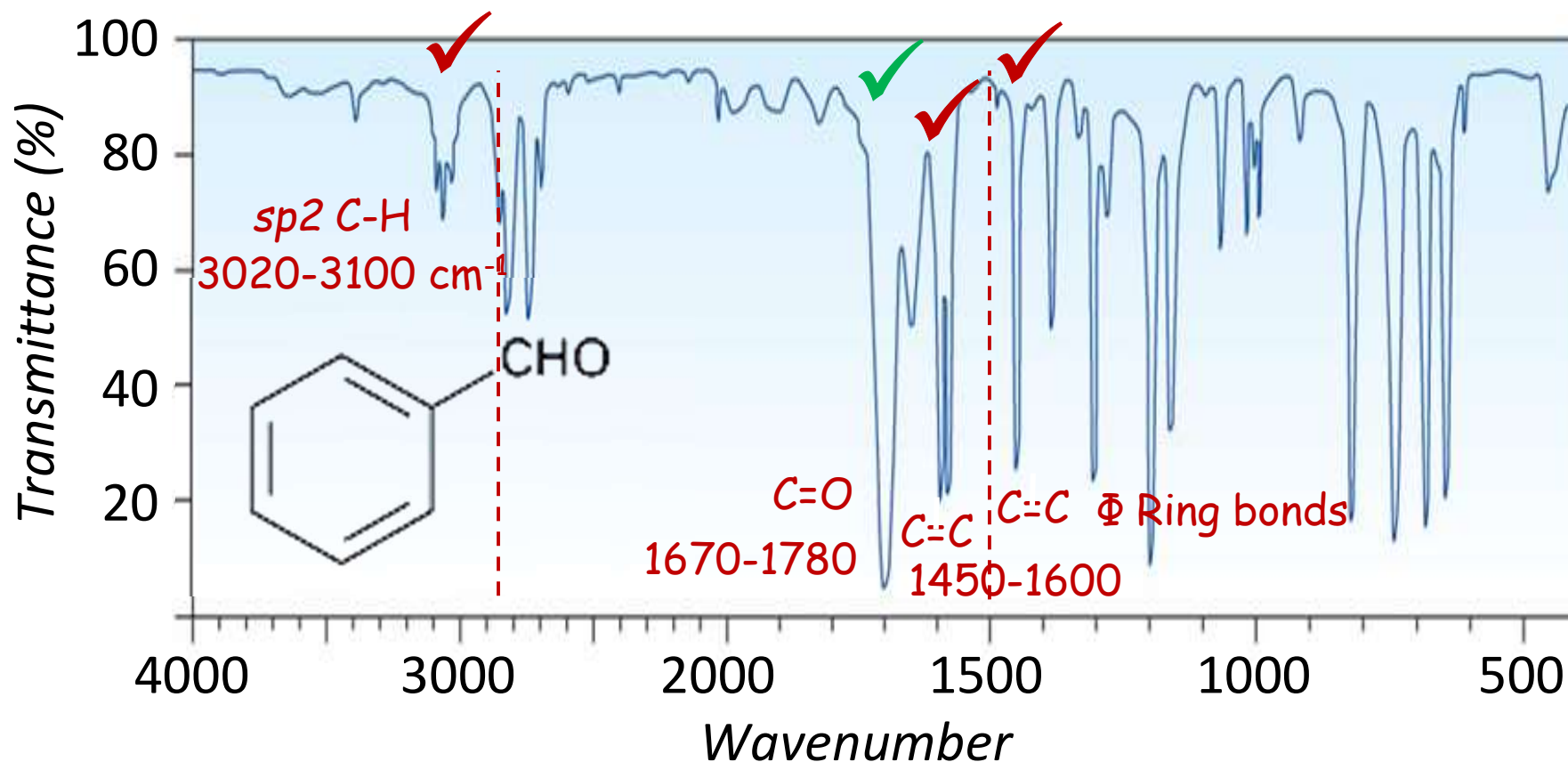


Infrared absorption frequencies of oxygen-containing substances

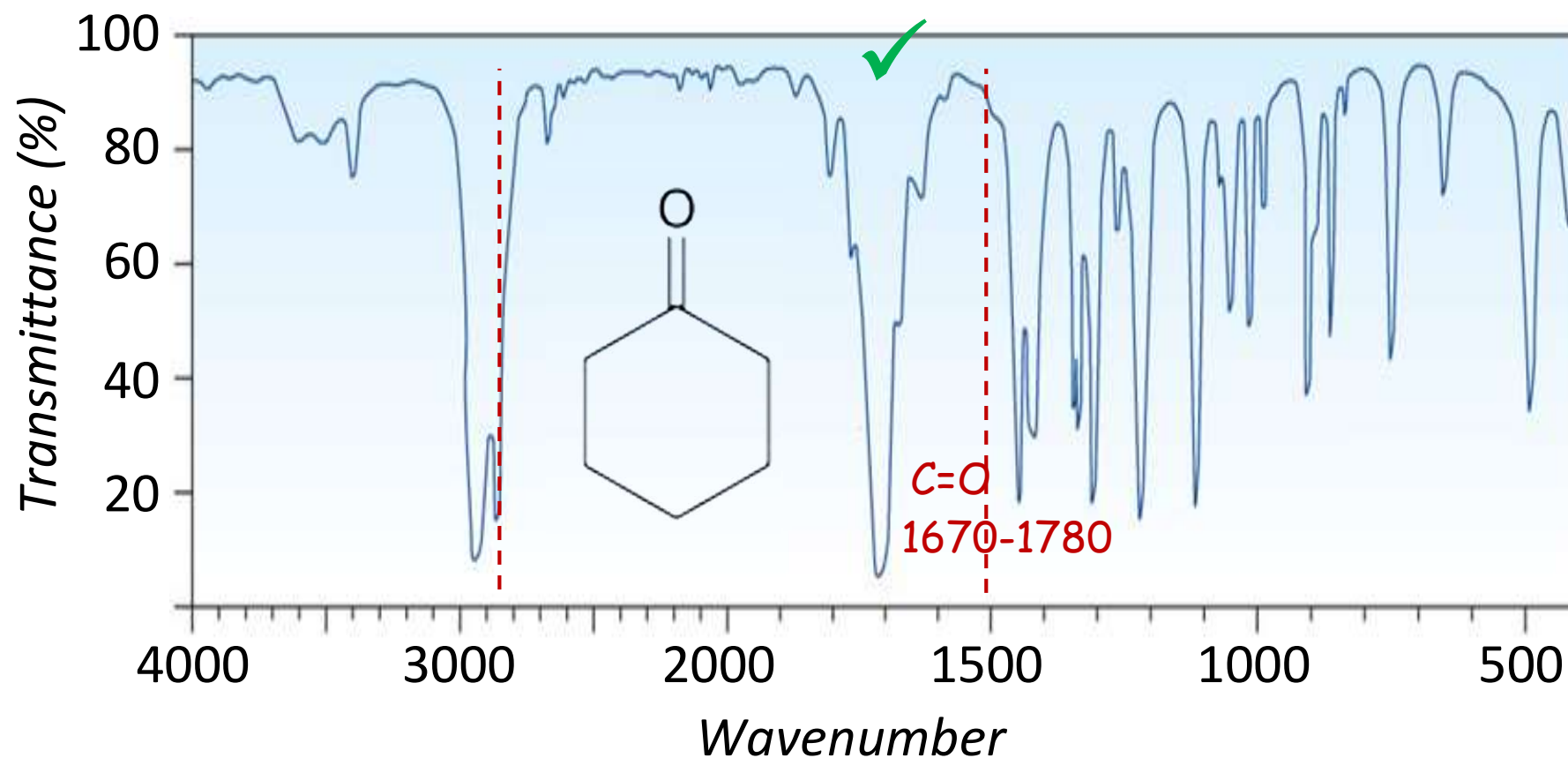


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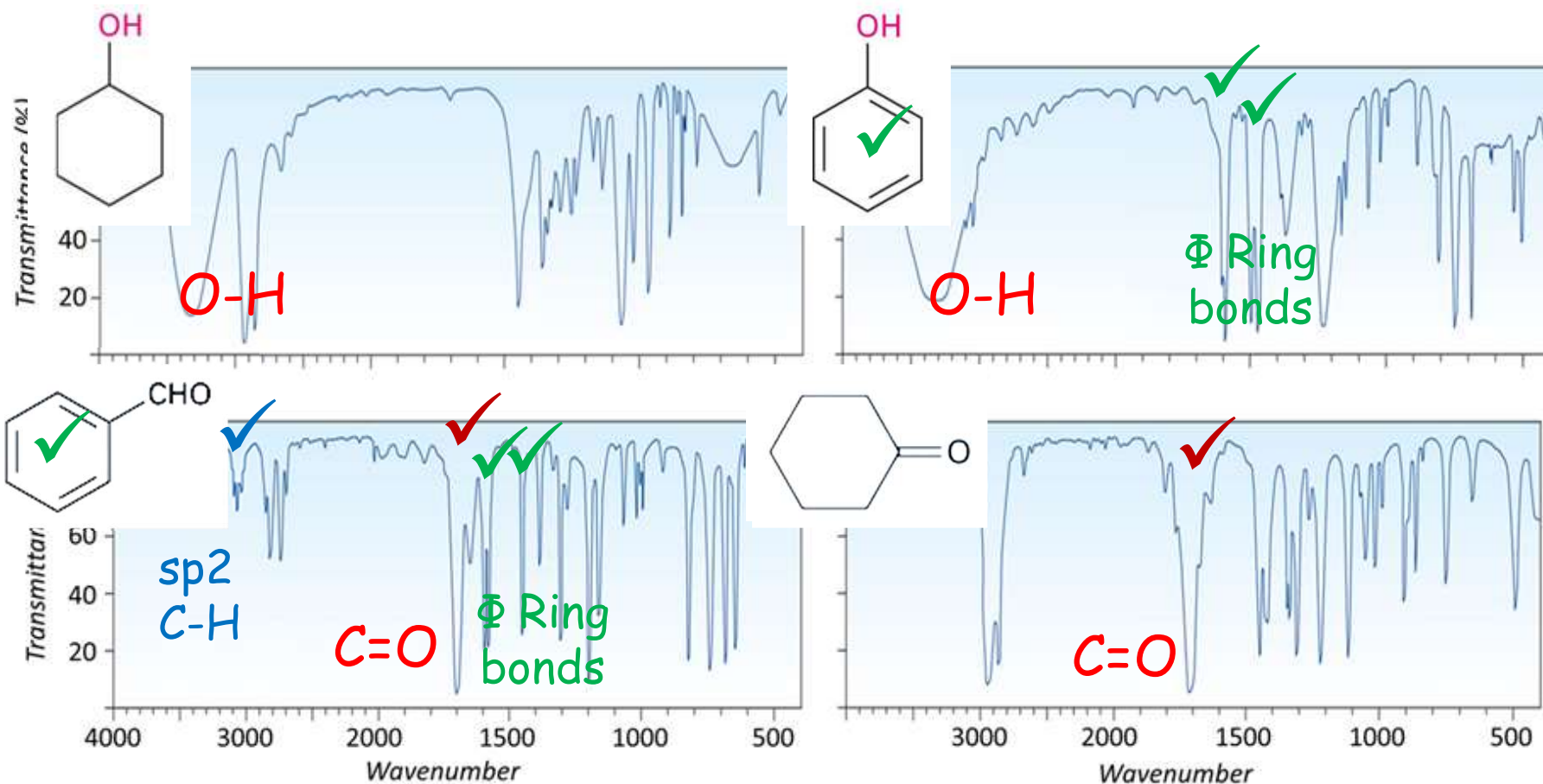
Infrared spectroscopy - aldehyde



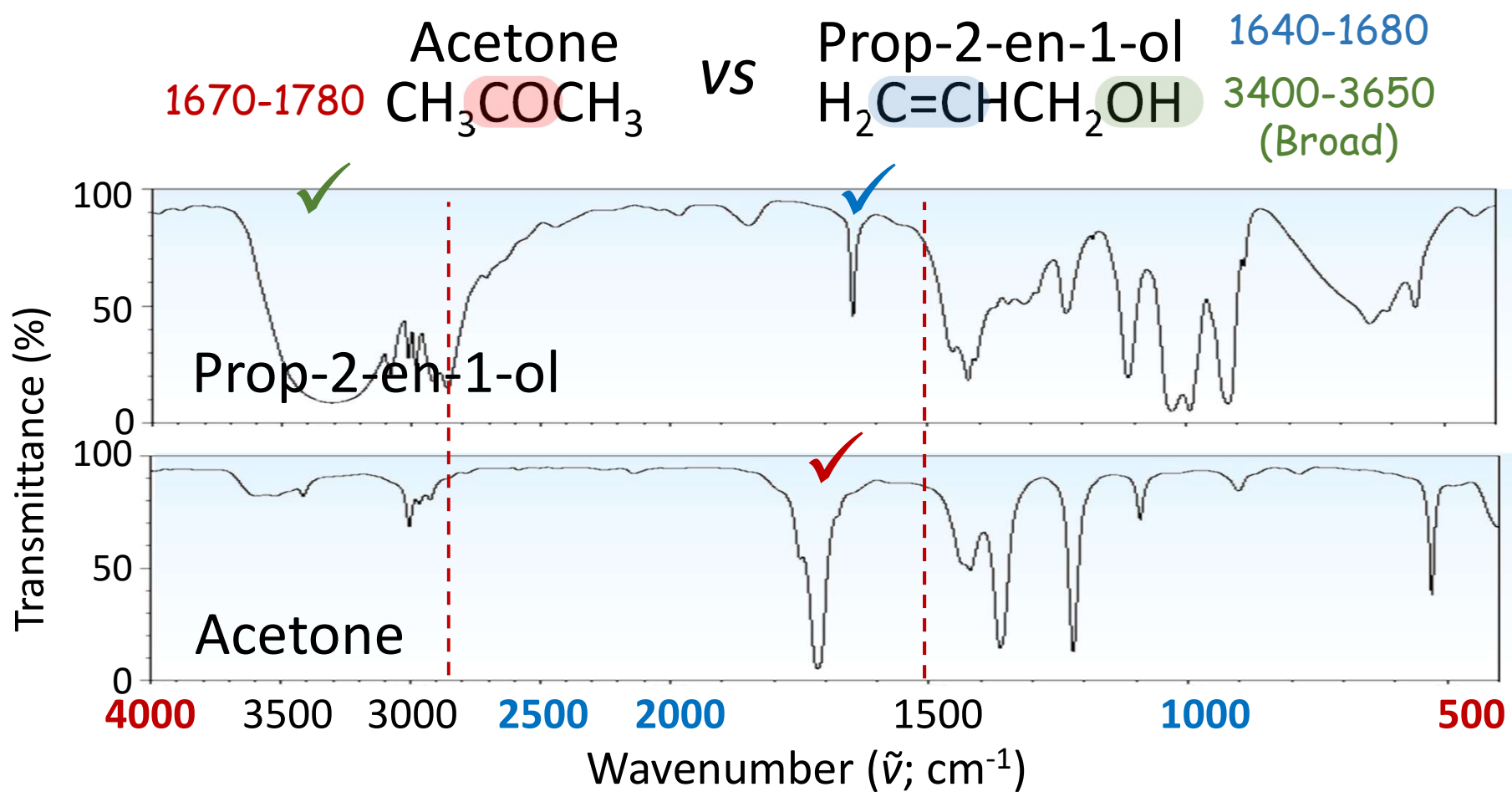
Infrared spectroscopy - ketone



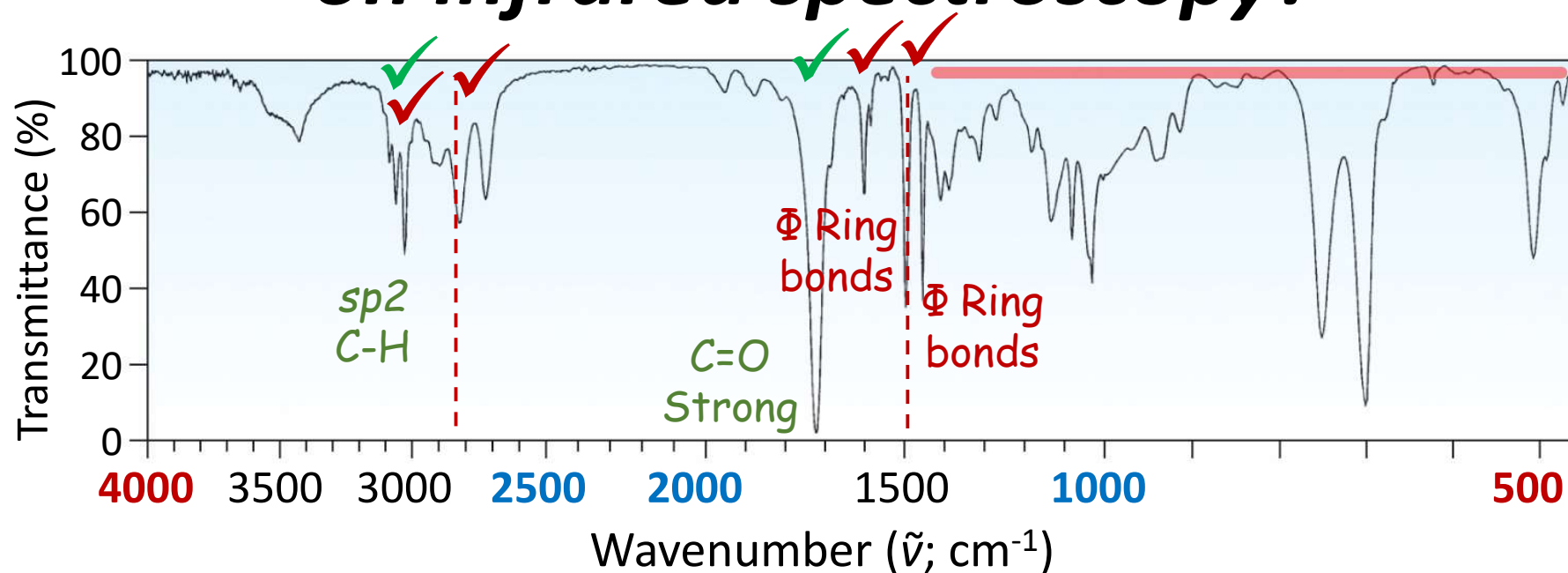
Infrared spectroscopy of alcohols, phenols, aldehydes and ketones



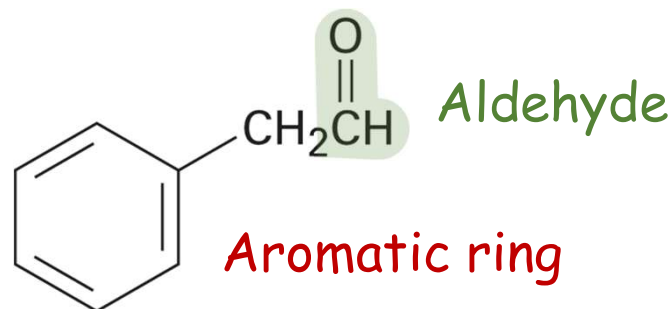
How to use infrared spectroscopy to distinguish two compounds with the same chemical formula C_3H_6O ?



How to predict the structure based on infrared spectroscopy?



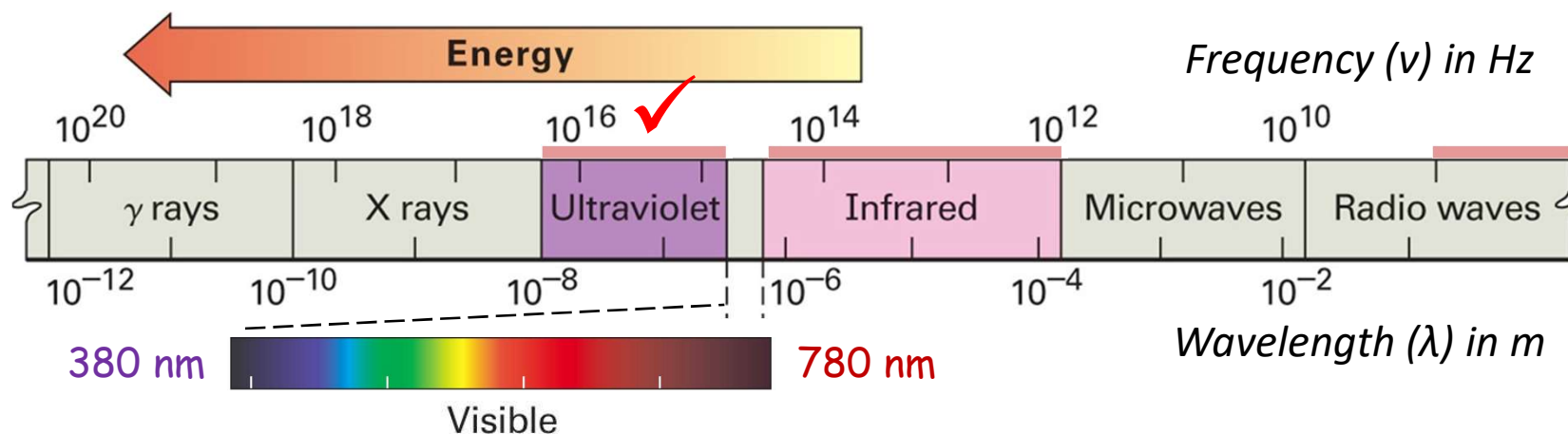
The table will be given in my test. Learn how to find the needed info from the table



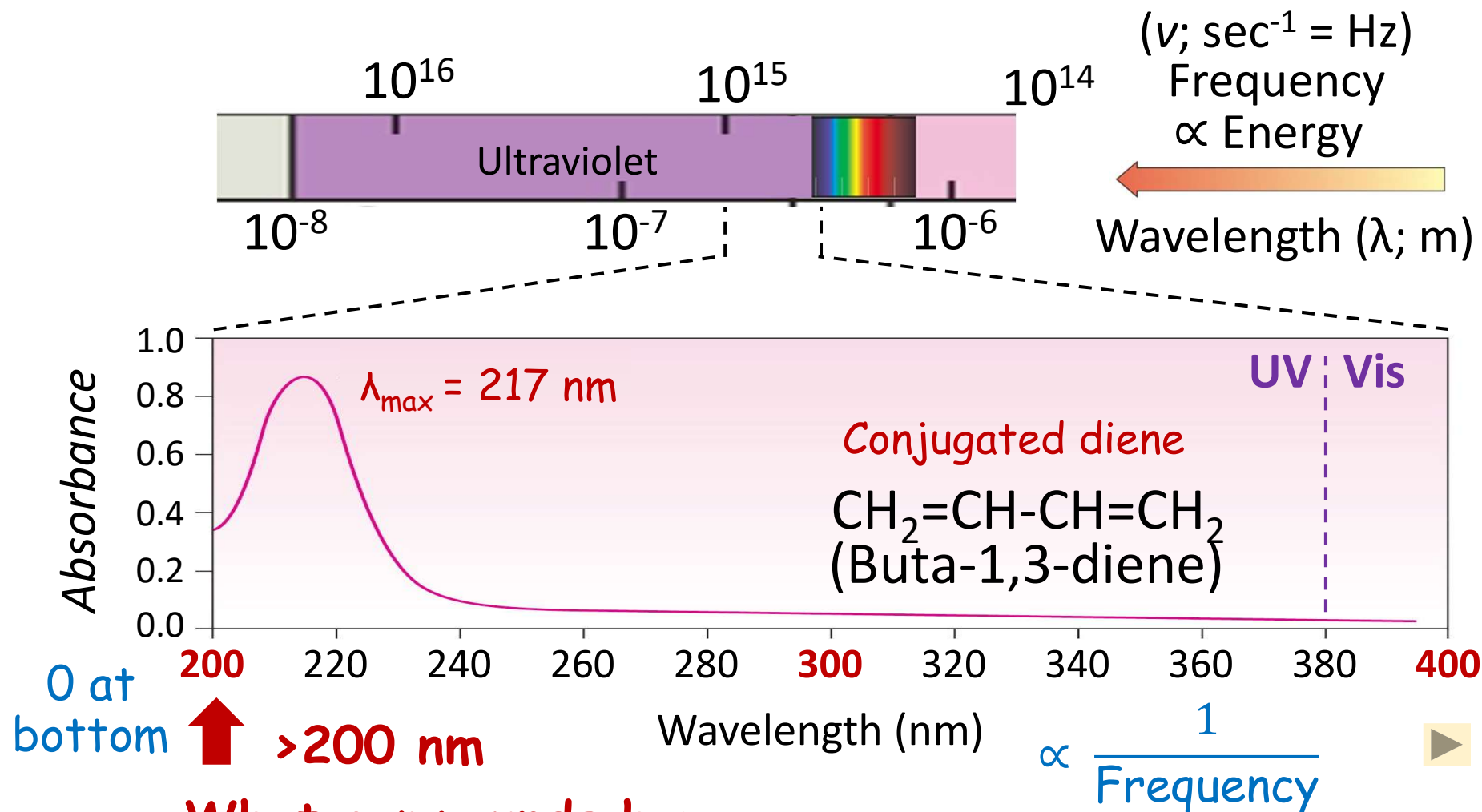
= S2

Spectroscopic analyses of organic compounds

Technique	Electromagnetic radiation	Structure
Infrared spectrometry	Infrared light	Covalent bonds
Ultraviolet spectroscopy	Ultraviolet light	$\sigma-\sigma^*$, $n-\sigma^*$, $\pi-\pi^*$, $n-\pi^*$ transitions
Nuclear magnetic resonance spectroscopy	Radio wave	Atomic nuclei in magnetic field



Ultraviolet spectroscopy



What compounds have a peak in the UV spectrum?

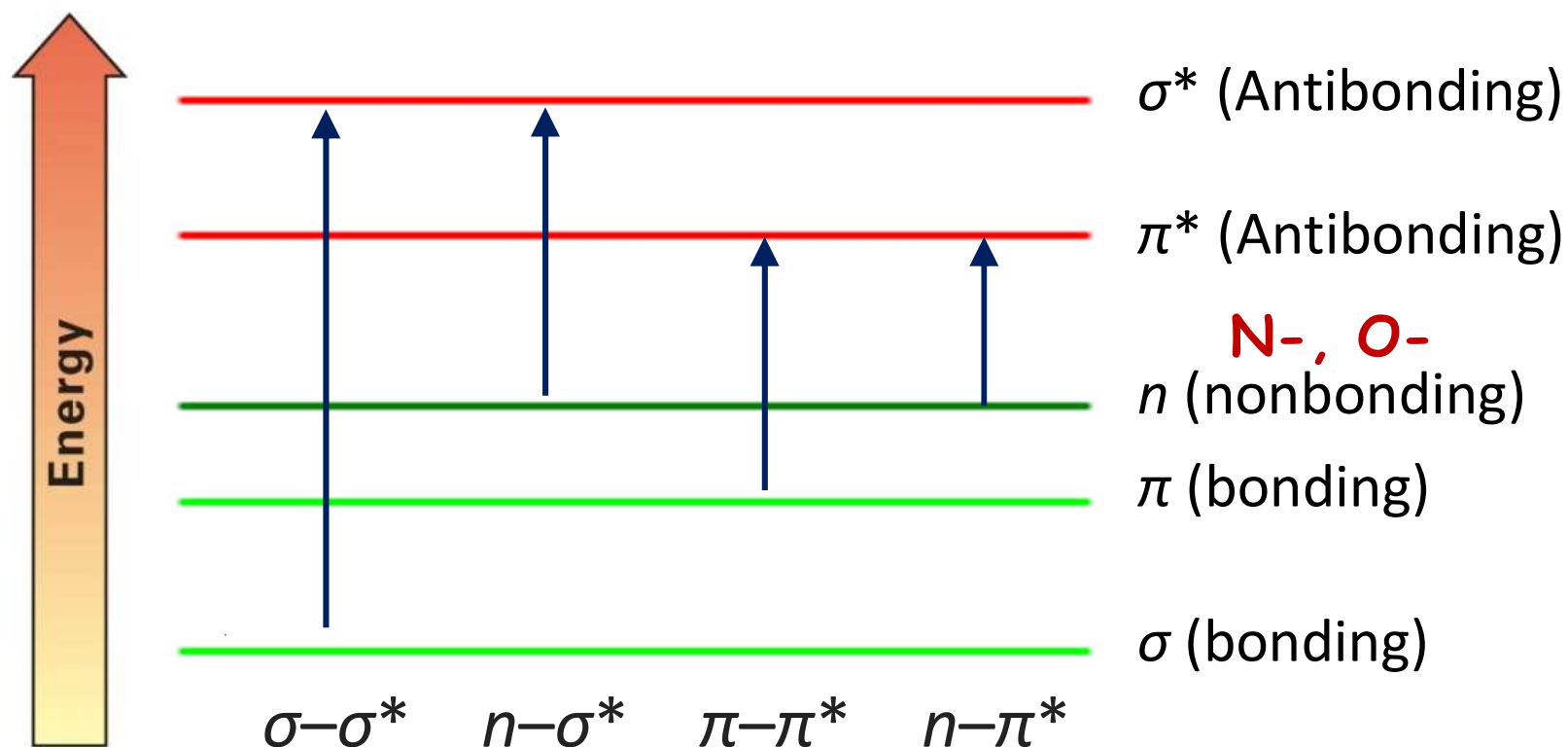
Ultraviolet spectroscopy

- ✓ Ultraviolet spectroscopy refers to absorption spectroscopy or reflectance spectroscopy in part of the ultraviolet spectral regions. In this region of the **electromagnetic** spectrum, atoms and molecules undergo **electronic transitions**. Absorption spectroscopy measures transitions from the _____ **state** to the _____ **state**.
- ✓ Molecules containing **bonding** (__ and __ electrons) or **non-bonding electrons** (__-electrons) can absorb energy in the form of ultraviolet to excite these electrons to higher anti-bonding molecular orbitals.
Highest Occupied MO
- ✓ The more easily excited the electrons, (*i.e.* lower energy gap between the **HOMO** and the **LUMO**), the longer the wavelength of light it can absorb **Lowest Unoccupied MO**

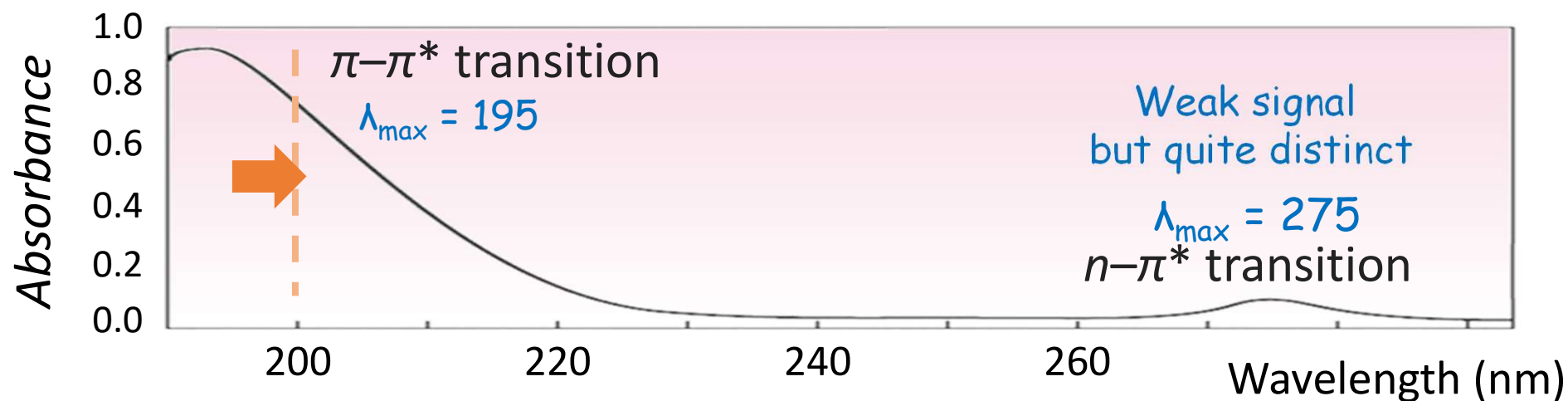
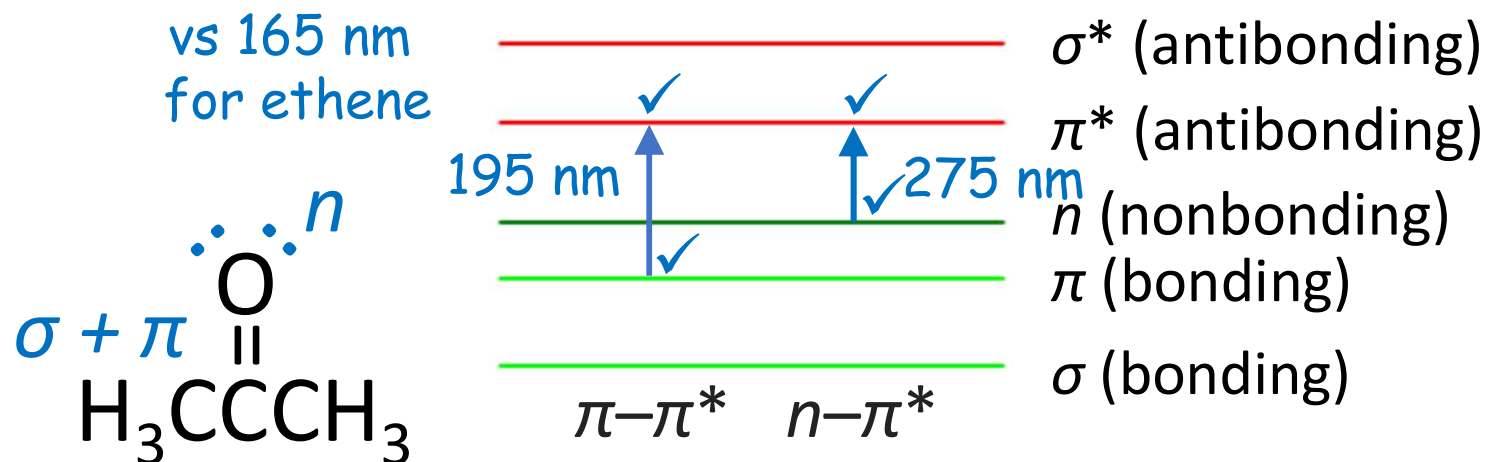
What properties are examined in UV-Vis?

Energy gap between HOMO and LUMO

Various HOMO-LUMO transitions leading to ultraviolet absorption

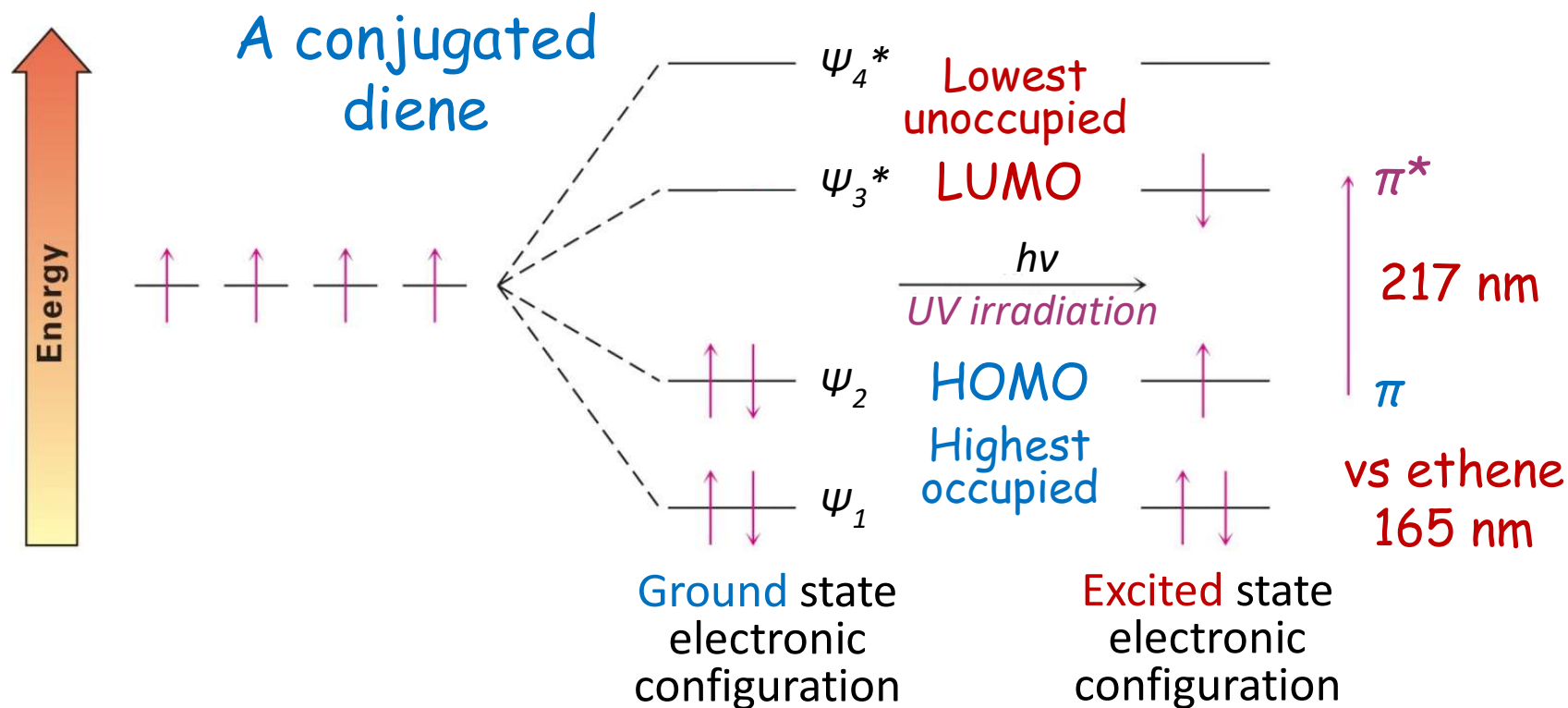


Ultraviolet spectroscopy of acetone



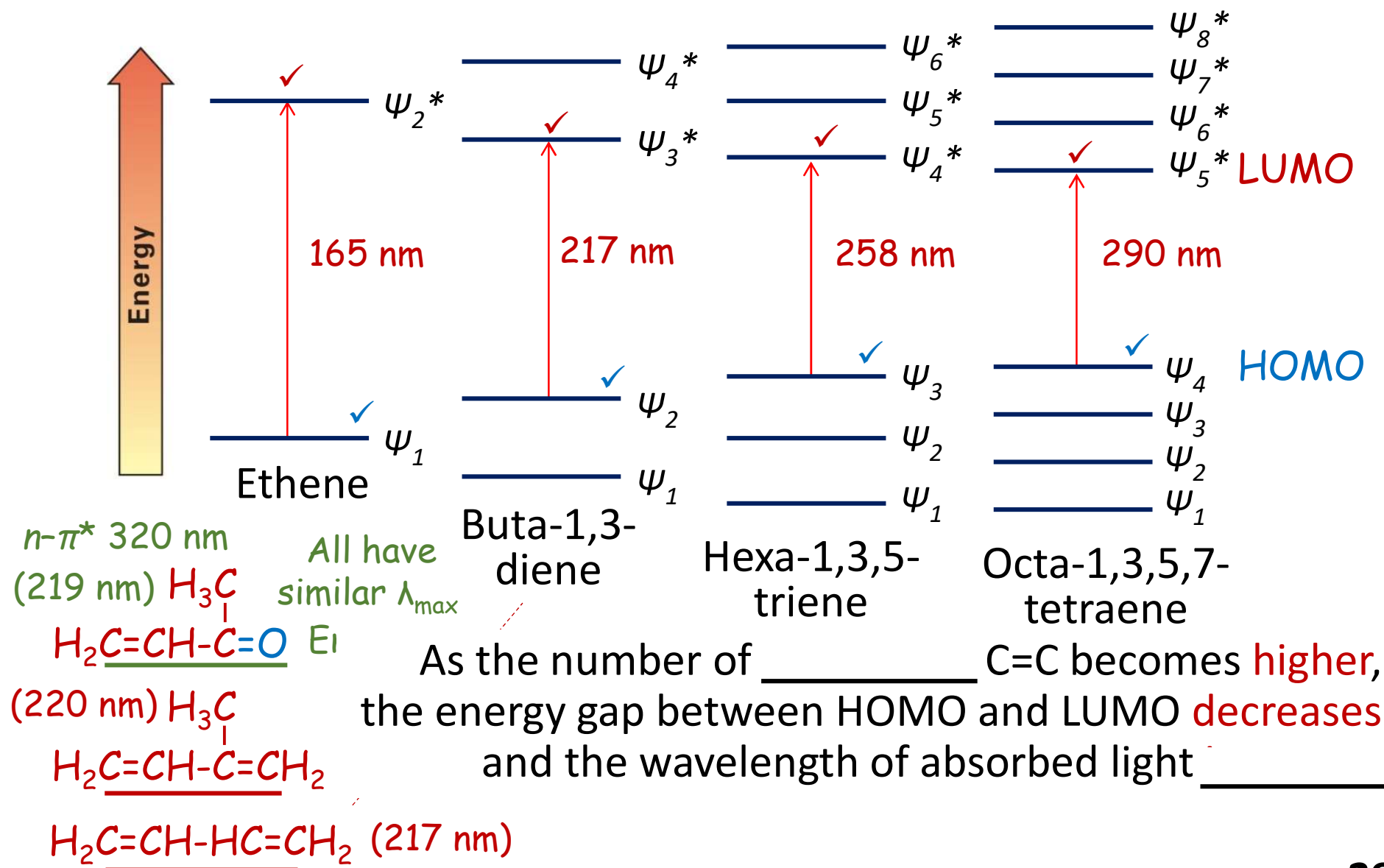
**What structure can narrow the energy gap
between HOMO and LUMO?**

$\pi-\pi^*$ transition for *buta-1,3-diene*

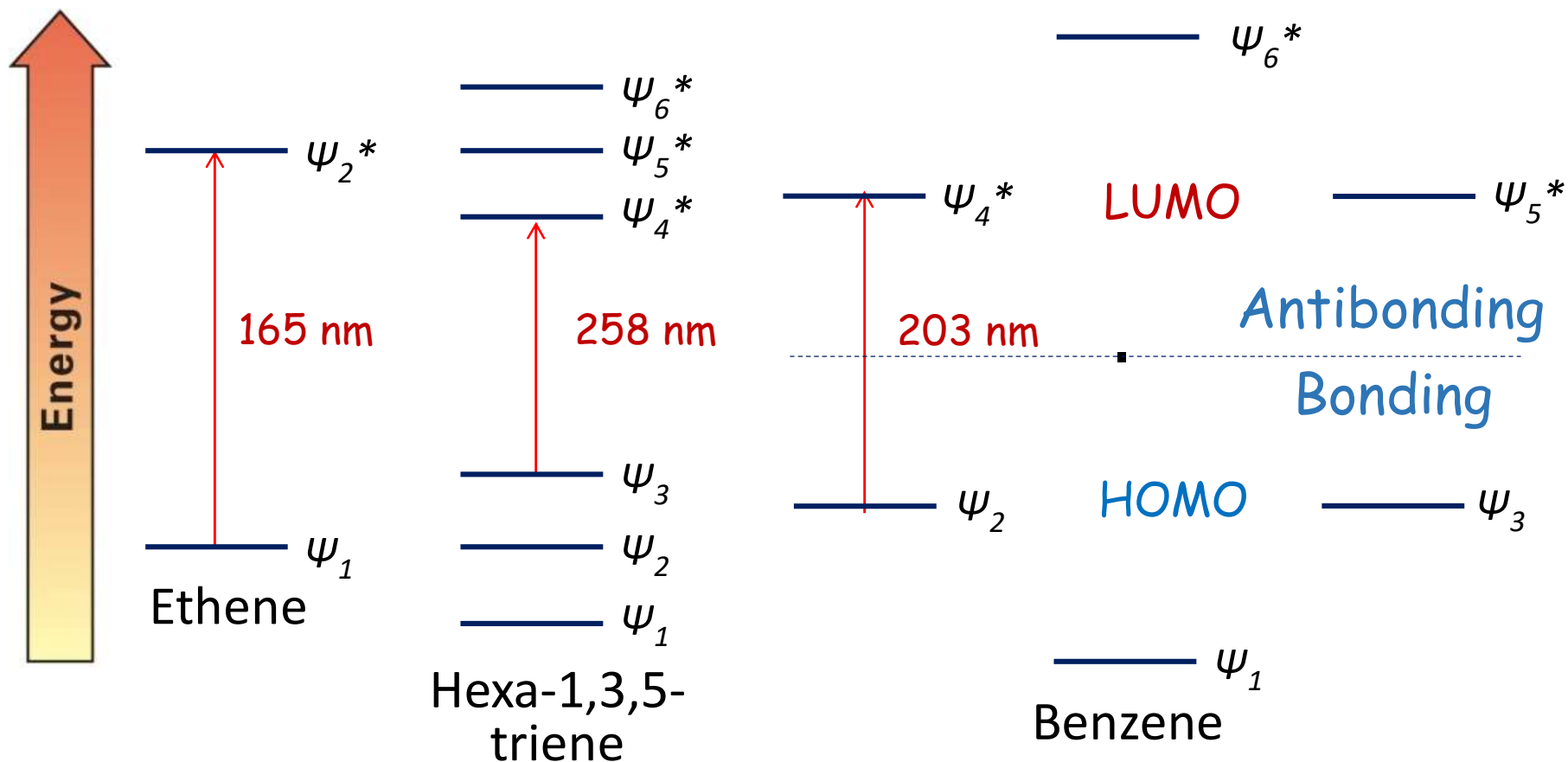


Four 2p atomic orbitals (AO) \rightarrow Four ψ molecular orbitals (MO)

UV absorption in conjugated polyenes



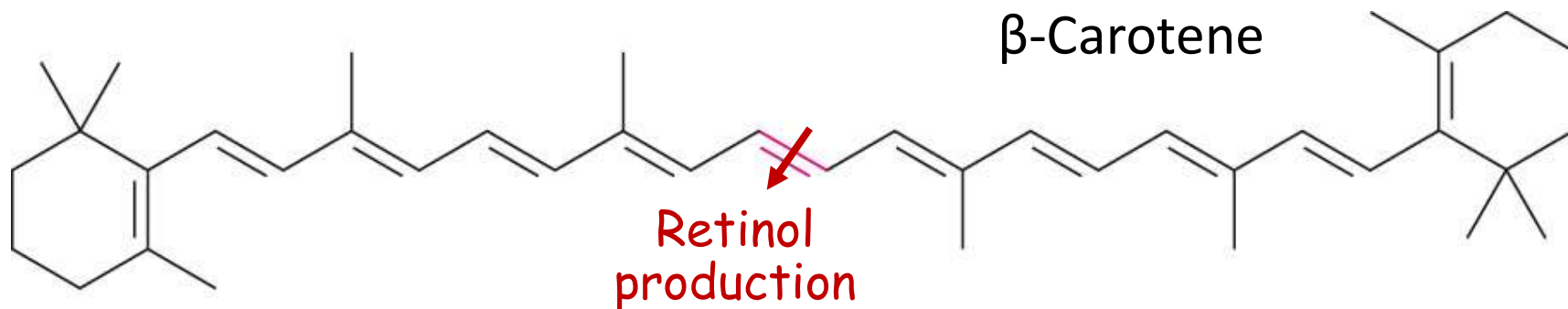
UV absorption in cyclic vs linear conjugation



Cyclic conjugation may give a peak with $\lambda_{\text{max}} > 200$ nm in UV spectra as well

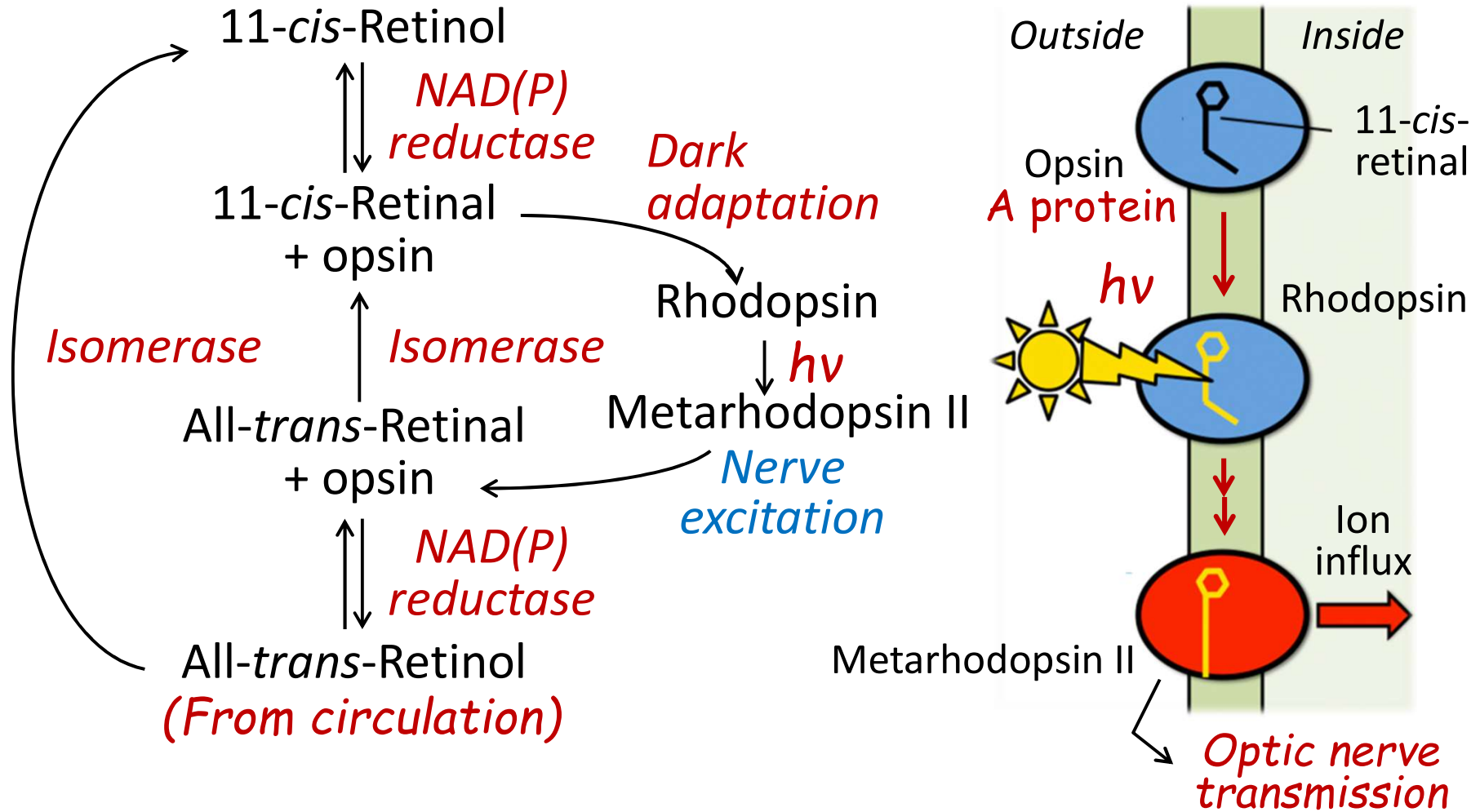
Colors of organic compounds result from extensive conjugation

- ✓ As the number of conjugated C=C becomes **higher**, the energy gap between HOMO and LUMO **decreases**. When the wavelength of absorbed light is in the range of **visible light**, the compound should become **colored**
- ✓ **β -Carotene** has $\lambda_{\text{max}} = 455 \text{ nm}$. When white light strikes β -carotene, wavelengths in the blue region are absorbed. Thus, the **yellow-orange colors** are transmitted to our eyes

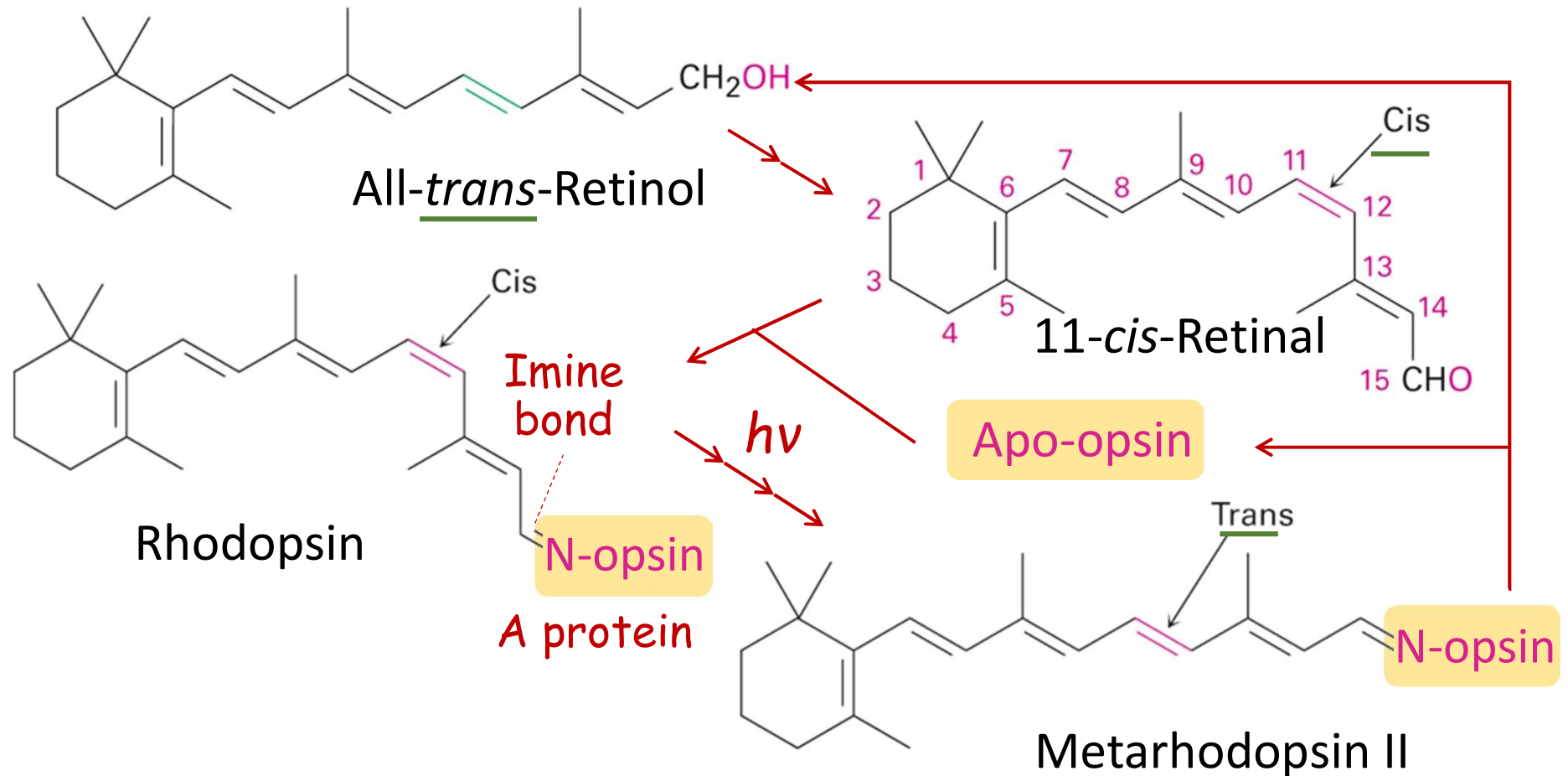


A vitamers in visual cycle

In retina



Some structures in visual cycle



Absorbance (A) and molar absorptivity (ϵ)

Absorbance (A) is the common logarithm of the ratio of **incident** (I_0) to **transmitted** (I) radiant power through a material

$$A = \log \frac{I_0}{I}$$

Molar absorptivity (ϵ), or **molar extinction coefficient**, is a measure of how strongly a chemical species or substance absorbs light at a particular wavelength

$$\epsilon = \frac{A}{cL} ; c = \frac{A}{\epsilon L}$$

Molar concentration **Sample pathlength (cm)**