

ASTR 620: Planetary Processes  
Professor Eric Nielsen

Lecture 10: Spectra



# Logistics

- Masks are encouraged
- No laptops, phones, or other electronic devices during class (I'll let you know in advance if we'll need laptops for an activity) **You may use a tablet to take notes if prefer, but please only use it for note-taking.**
- Remember to bring you response card to class
- Midterm in 2 weeks: Wednesday, October 5th (here in class)
- Homework 3 due Wednesday, September 28 at 11:59pm

# Review of the last class

- A surface is in equilibrium if:
  - (A) — Every square inch of the surface is covered by a crater
  - (B) — Craters no longer form
  - (C) — Each new crater is accompanied by the destruction of an existing crater
  - (D) — Lava flows wipe out all previous craters

# Review of the last class

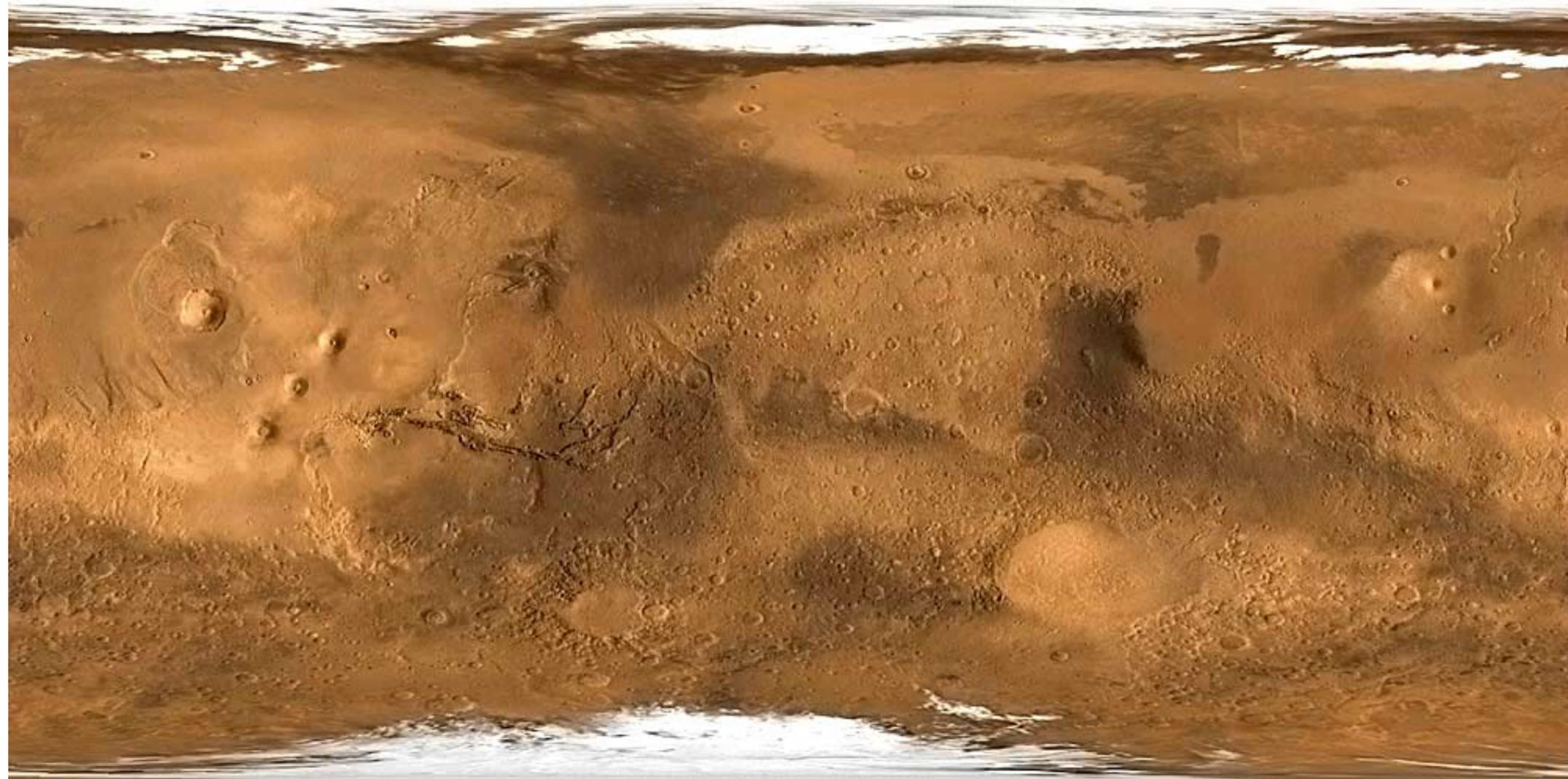
- Which best describes the distributions of craters and impactors in the Solar System?
  - (A) — There are more large craters than small craters, there are more small impactors than large impactors
  - (B) — There are more small craters than large craters, there are more small impactors than large impactors
  - (C) — There are more large craters than small craters, there are more large impactors than small impactors
  - (D) — There are more large craters than small craters, there are more small impactors than large impactors

# Review of the last class

• Which hemisphere of Mars has an older surface?

- (A) — North
- (B) — South
- (C) — They're equally old

North



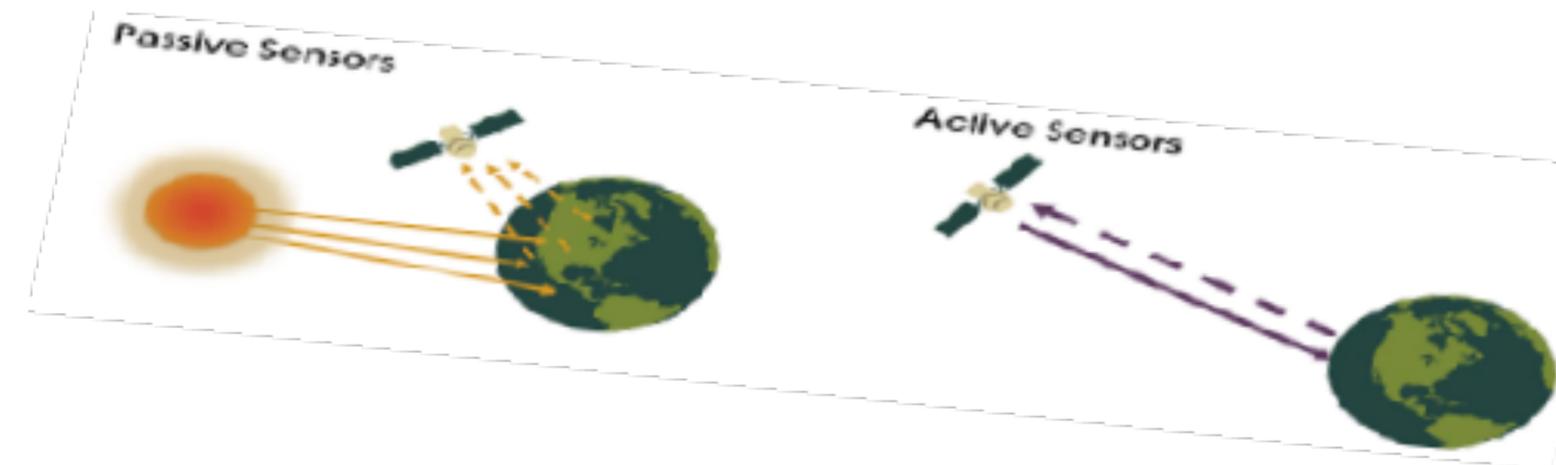
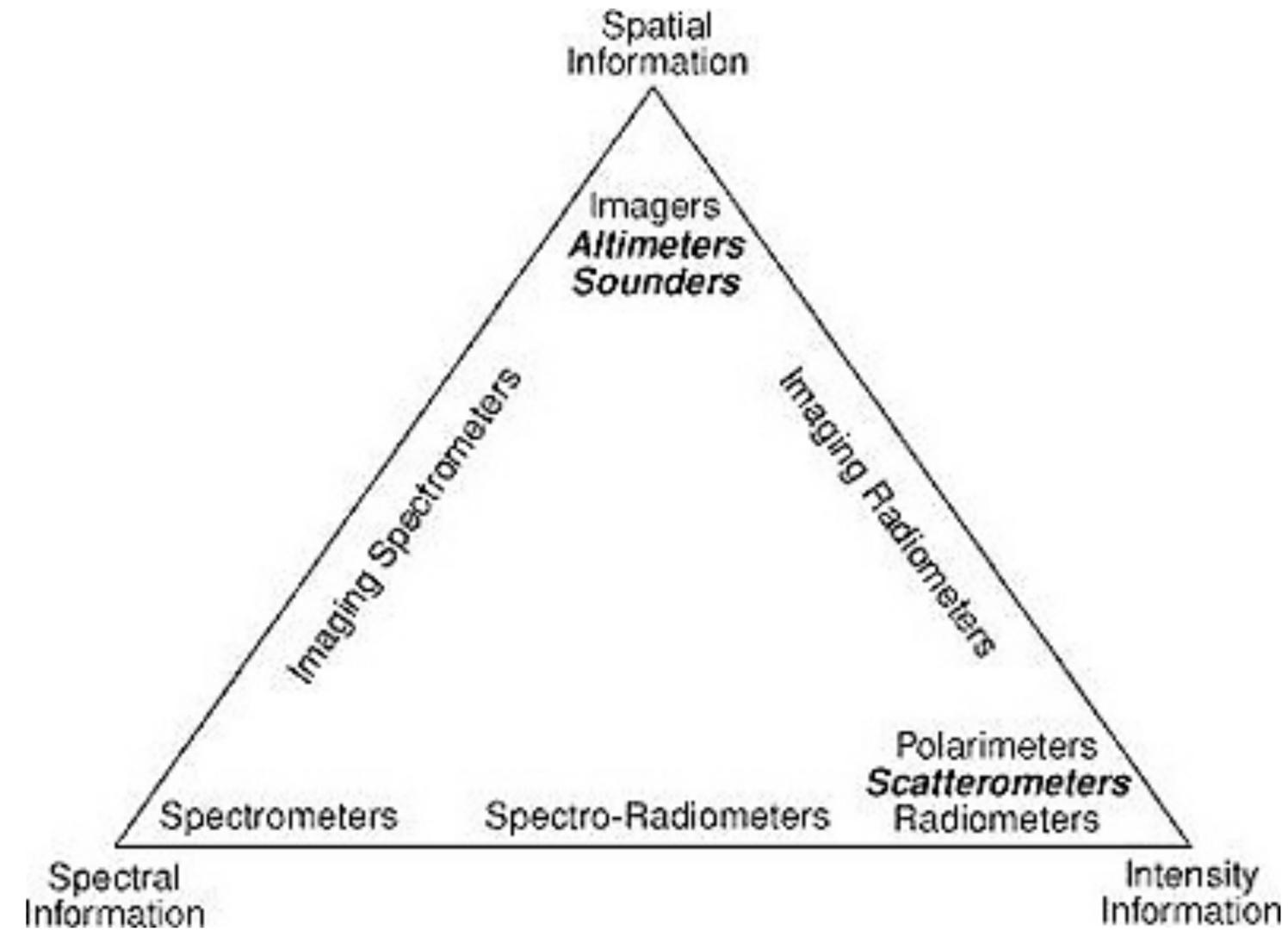
South

# Review of the last class

- Small meteors (10 cm) that enter Earth's atmosphere at high speeds:
  - (A) — Bounce back into space
  - (B) — Form very small craters on the ground
  - (C) — Completely evaporate before they hit the ground
  - (D) — Break up, with the fragments falling at terminal velocity

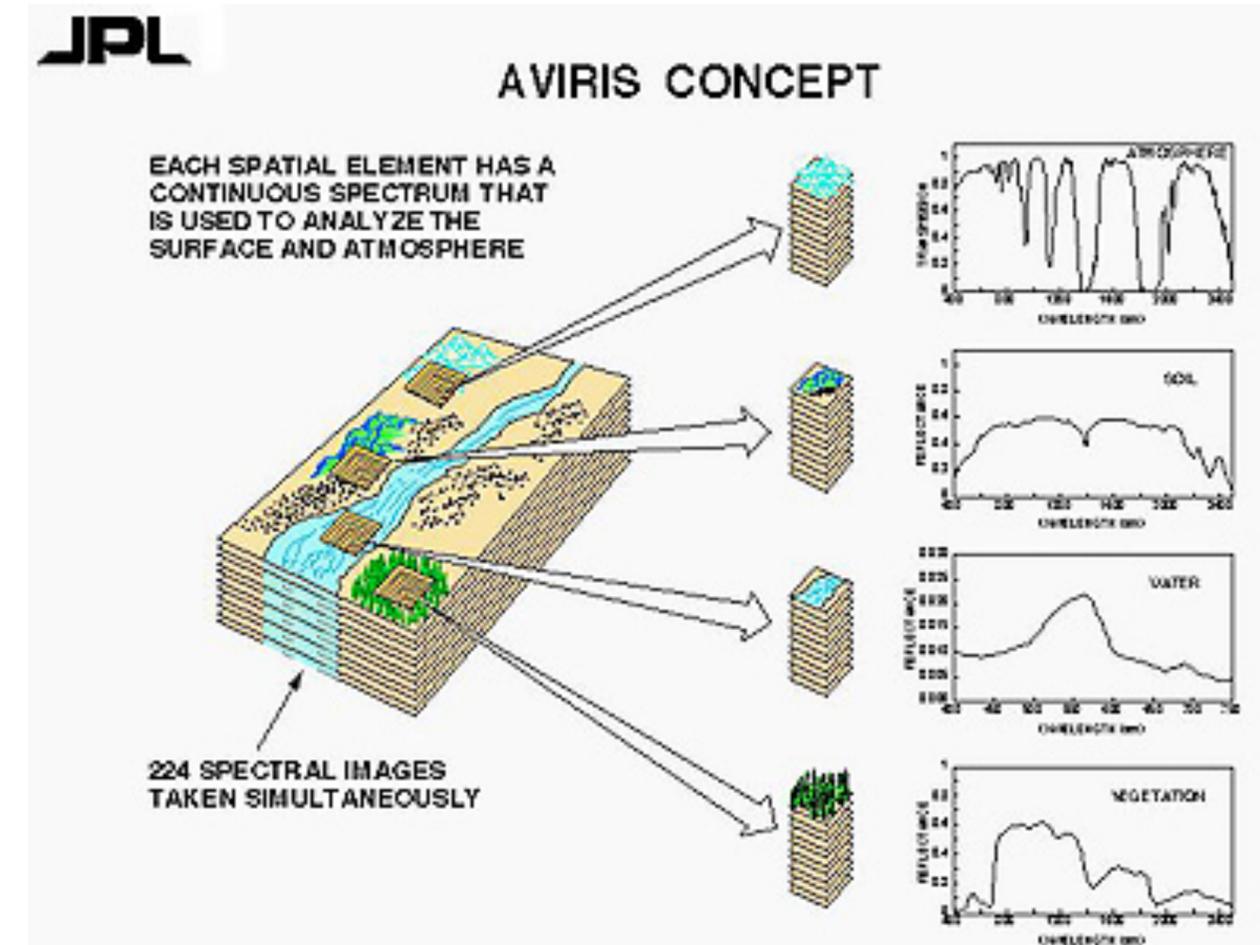
# Remote Sensing

- In Planetary Science (and astronomy) we collect information on distant objects through light
- Photometry (intensity): total brightness
- Spectra: light broken up by wavelength
- Spatially resolved images: light broken up by angular position

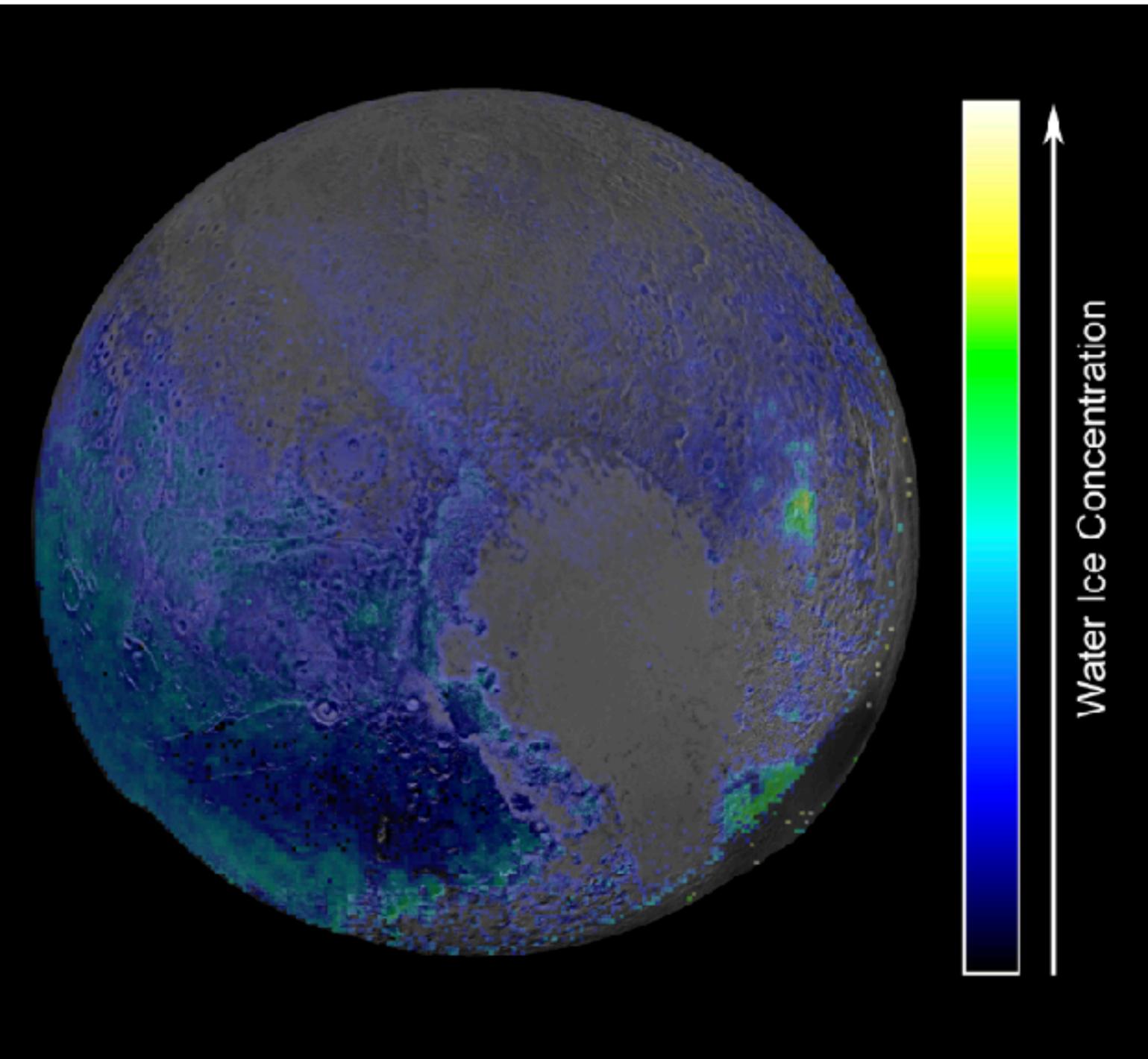


# Hyperspectral Imaging (Planetary Science)

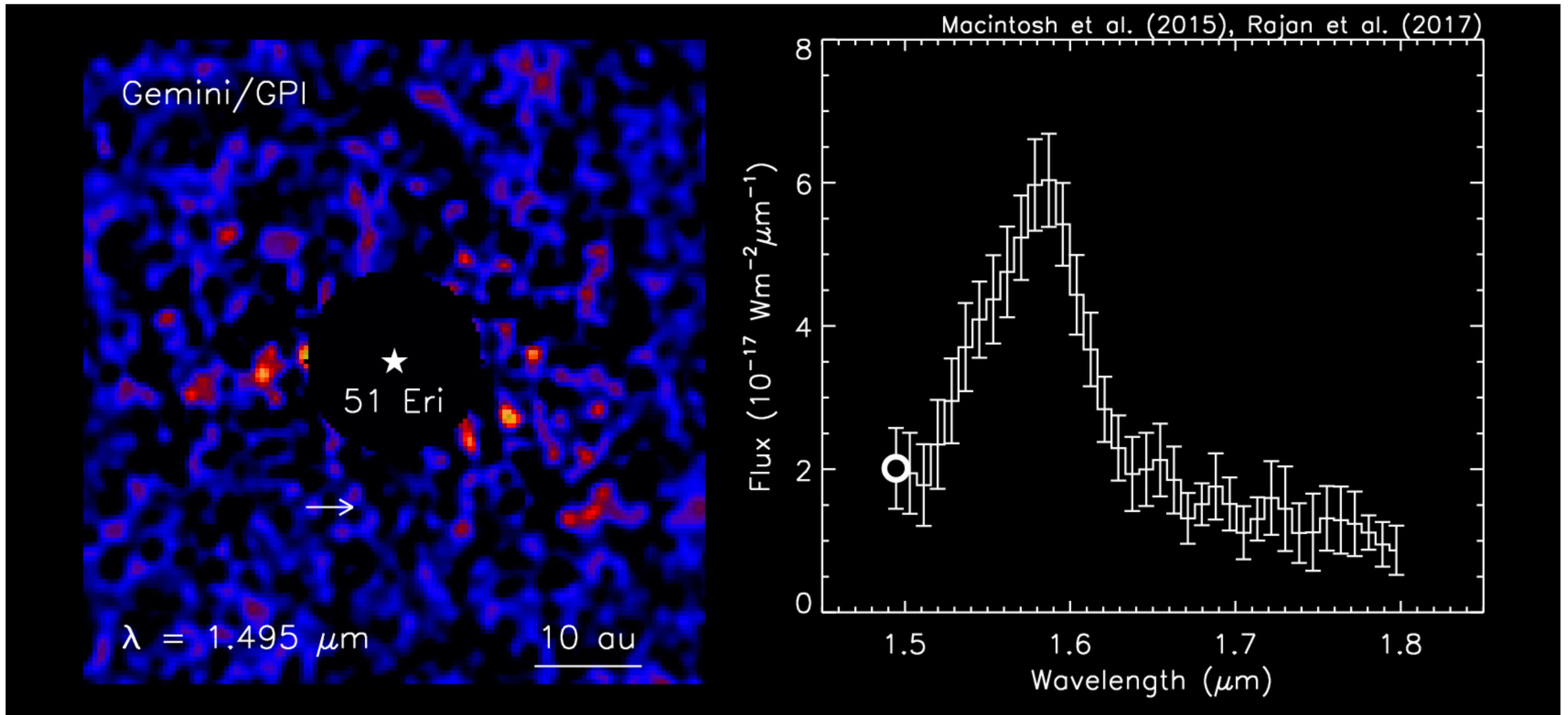
- Combination of imaging and spectroscopy:
  - Spatially resolved in the x (east-west, say) and y (north-south, say) directions
  - Spectrally resolved in the wavelength direction
  - Data “cube” is x vs y vs lambda



# Hyperspectral Imaging (Planetary Science)

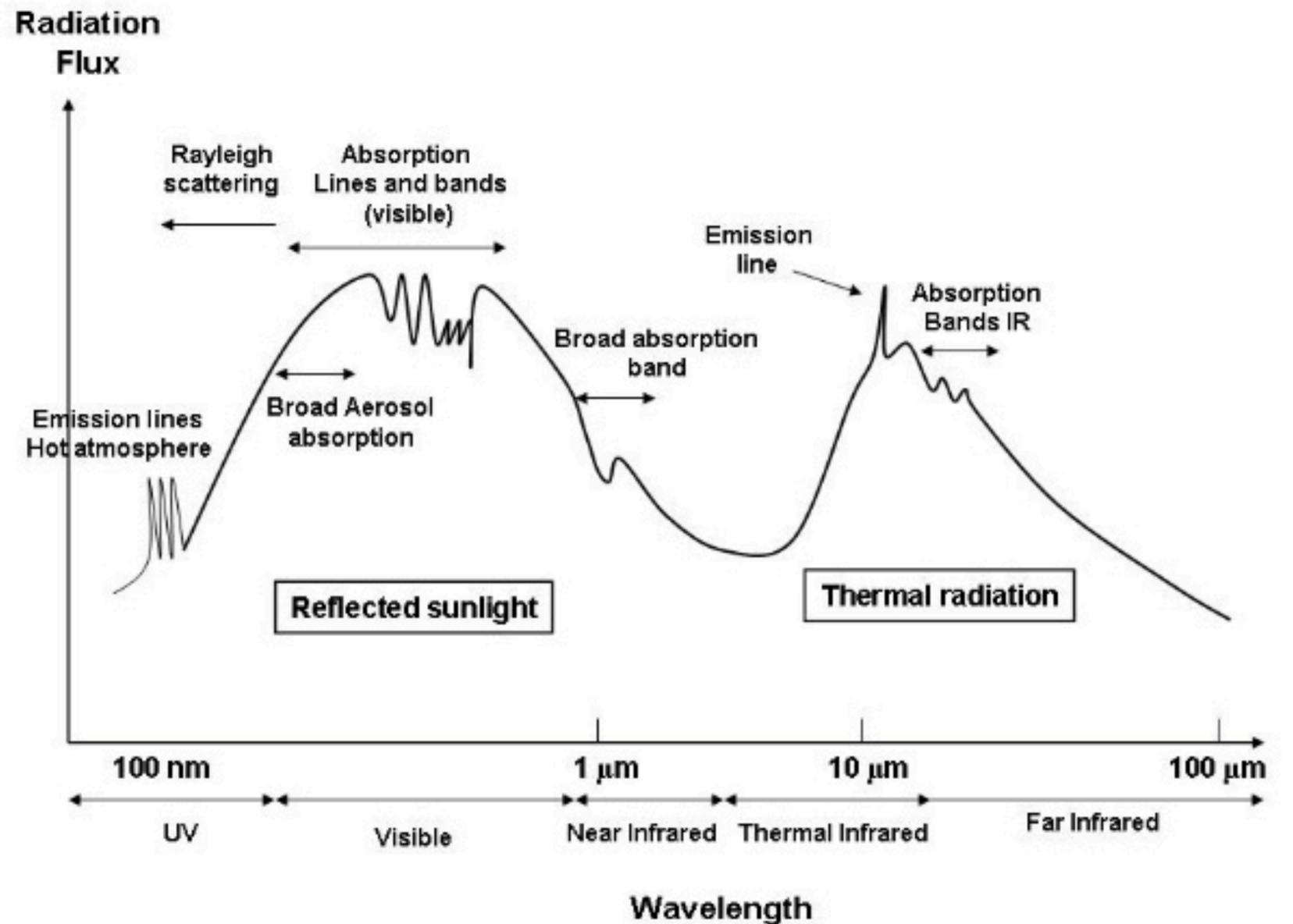


# Integral Field Spectroscopy (Astronomy)



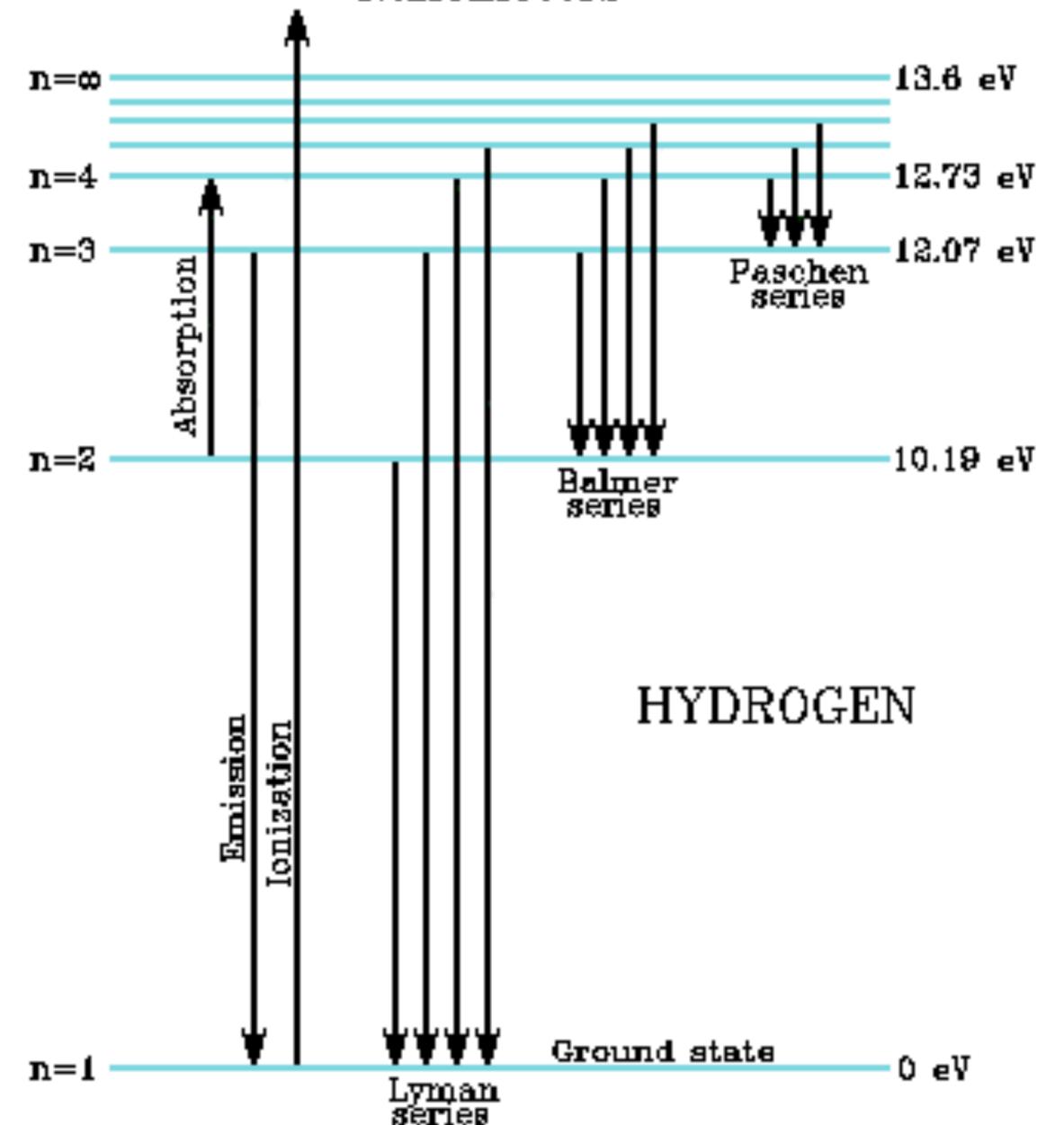
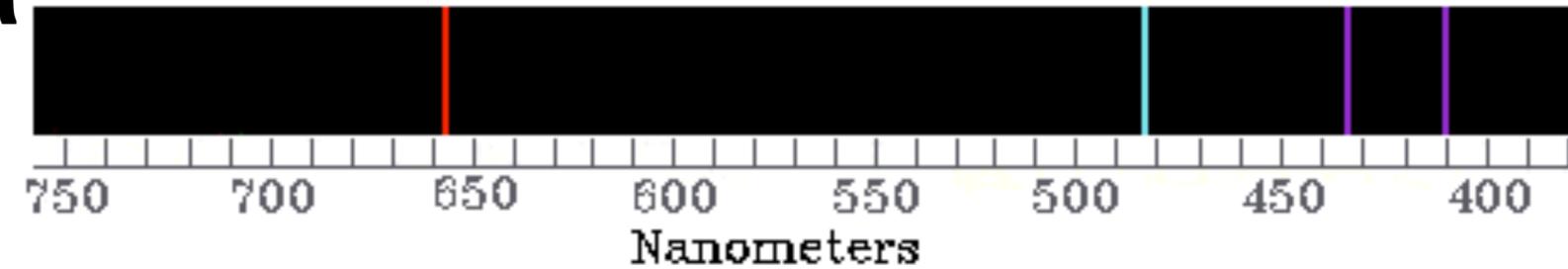
# Planetary Spectra

- Most planetary spectra have two components:
  - Reflected sunlight (starlight) spectrum, usually in the visible
    - Depends on angle between sun, planet, and observer
  - Thermal emission, mainly in the infrared, approximately a blackbody given the object's temperature. Internal heat comes from:
    - Reprocessed sunlight
    - Internal heat sources (e.g. radioactive decay)
    - Residual heat from formation



# Absorption and Emission Spectra

- Atoms absorb (or emit) a photon when an electron moves between two energy levels
- Ground state ( $n=1$ ) is the “closest” to the nucleus, and the lowest energy state for an electron
- Moving up to a higher energy level absorbs a photon
- Moving down to a higher energy level emits a photon
- The energy spacing between higher energy levels ( $n=\text{large}$  to  $n=\text{large} + 1$ ) is smaller than the spacing between lower energy levels

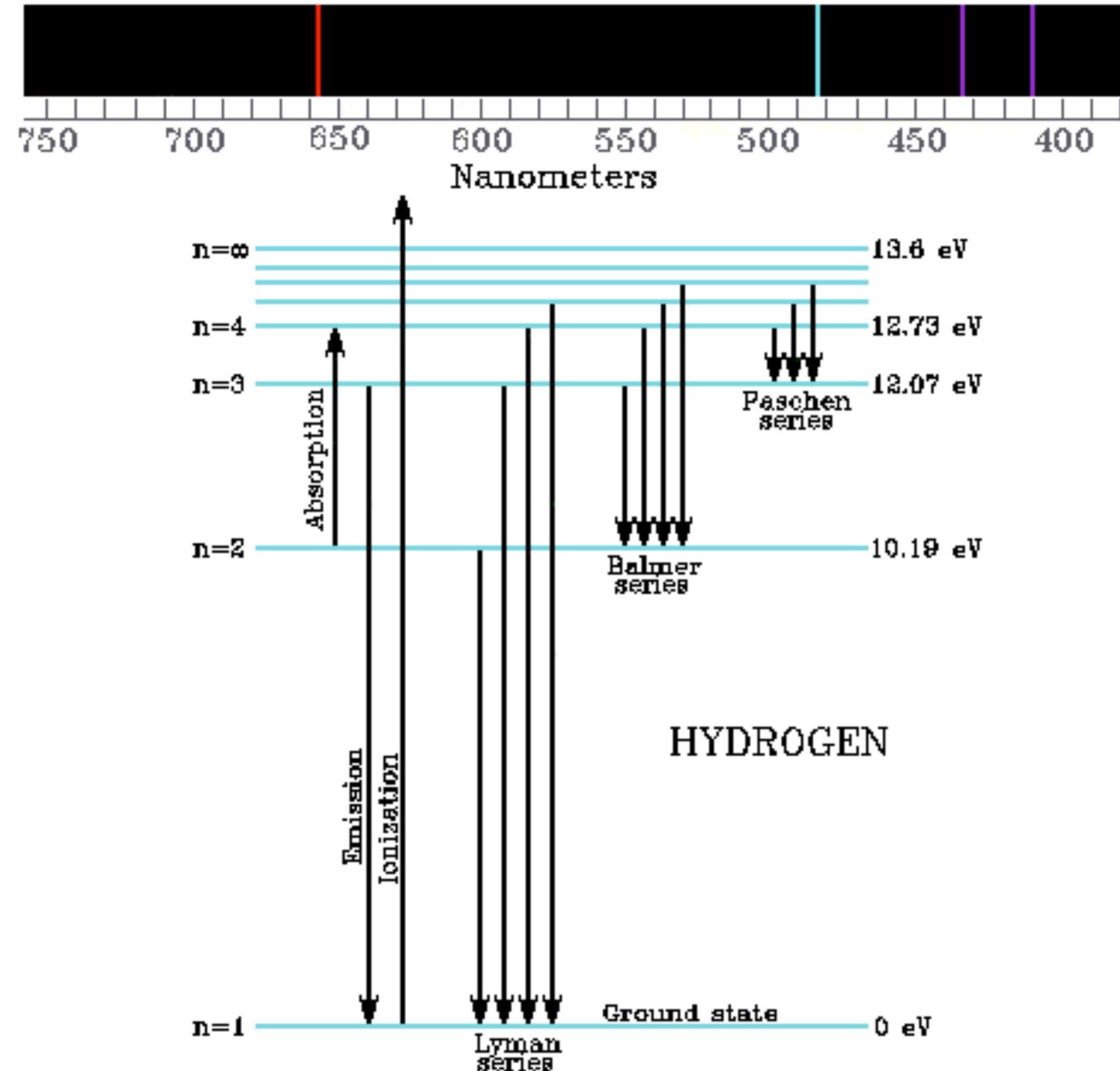


# Emission

- Requires atom starts in an excited (not ground) state
- Can spontaneously move to a lower state and emit a photon
- Intensity of emission spectrum from level  $j$  to level  $i$  is given by:

$$N_j h \nu_{ji} A_{ji}$$

- $N_j$ : Number of atoms in state  $j$  per unit volume
- $h \nu_{ji}$ : energy difference between the two levels
- $A_{ji}$ : Einstein A coefficient for spontaneous emissions (number of transitions per second of atom from excited state  $j$  to state  $i$ )

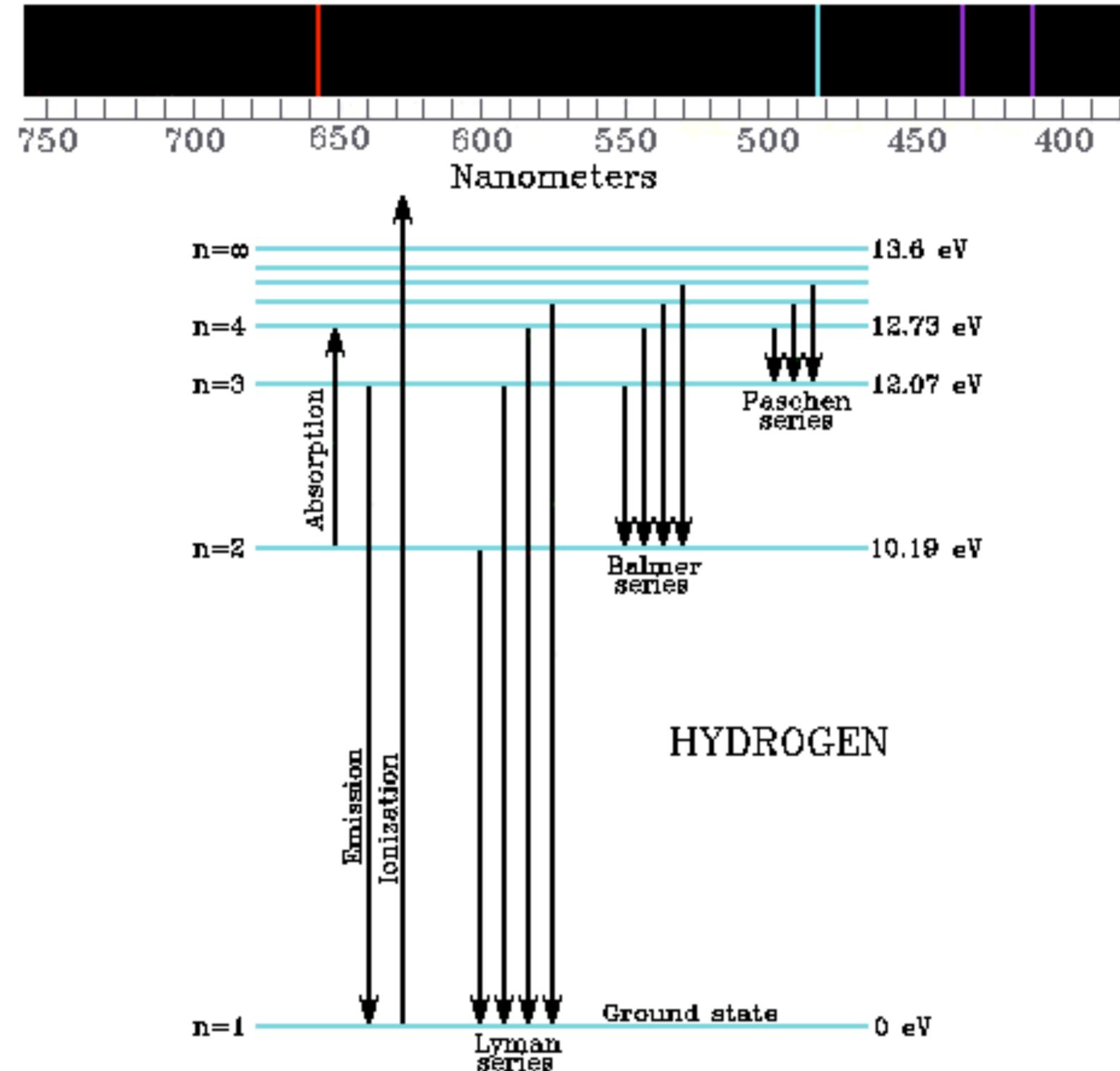


# Absorption

- Intensity of absorbed light resulting from electrons initially in state  $i$  move up to level  $j$ :

$$N_i h\nu_{ij} B_{ij} \rho_\nu$$

- $N_i$ : Number of atoms in state  $i$  per unit volume
- $h\nu_{ij}$ : energy difference between the two levels
- $B_{ij}$ : Einstein B coefficient for absorption
- $\rho_\nu$ : density of radiation with frequency  $\nu$

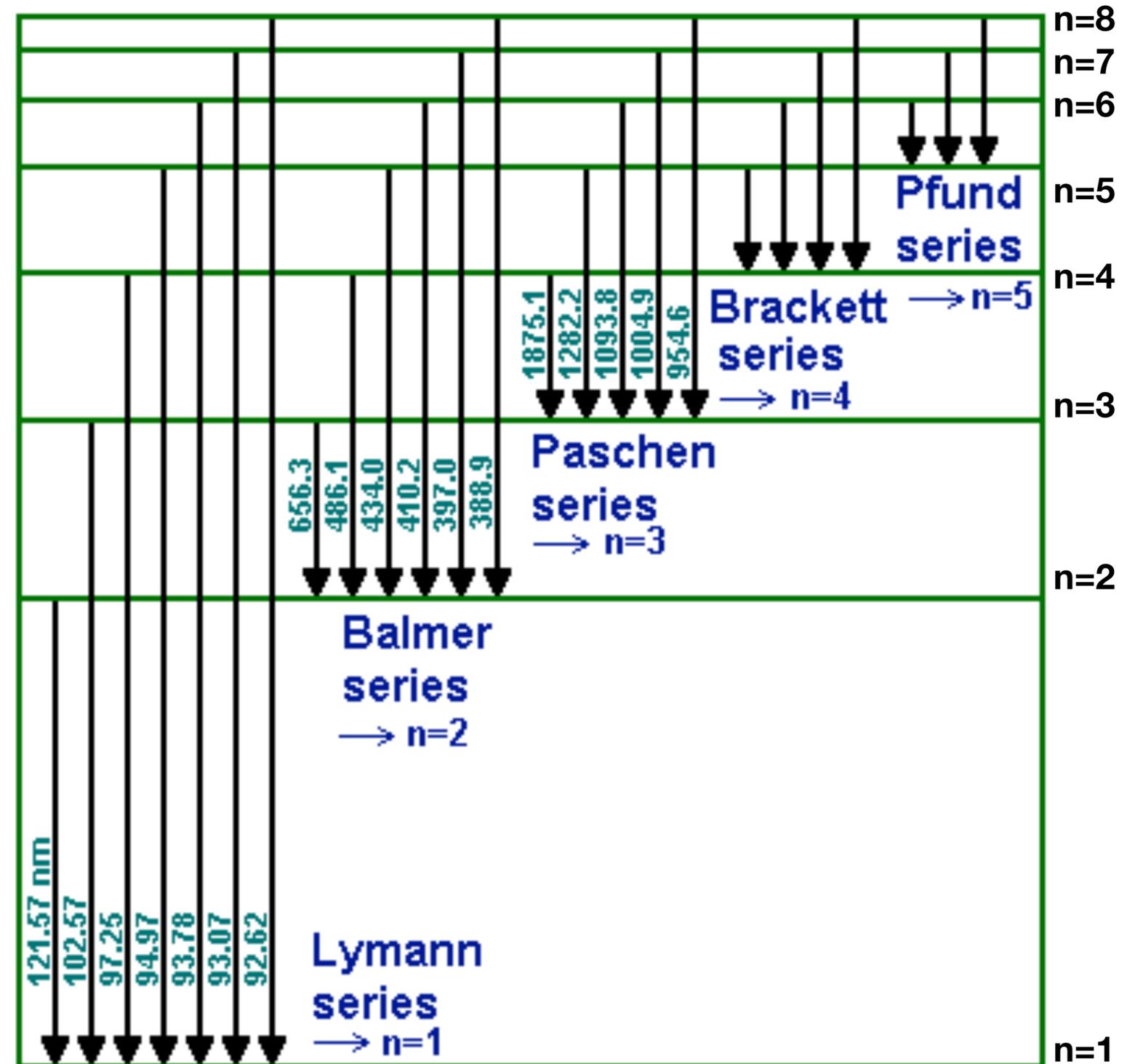


# Hydrogen

- The Schrodinger equation can be solved for hydrogen and other single-electron atoms (and not much else)
- The energy for each energy level (n) is given by:

$$E_n = -\frac{m_e Z^2 e^4}{8h^2 \epsilon_0^2 c} \frac{1}{n^2} = -R \frac{Z^2}{n^2}$$

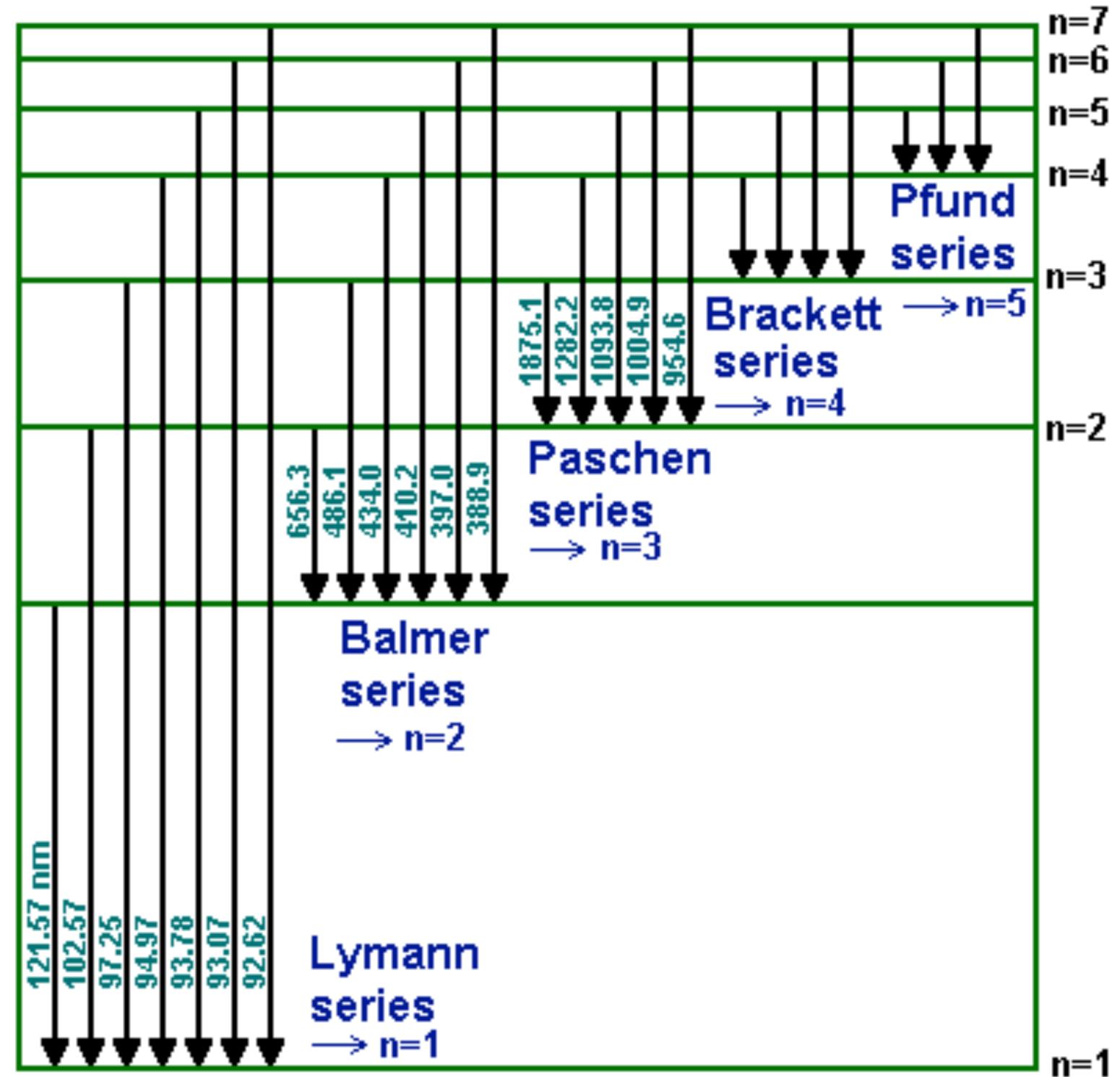
- $m_e$ : electron mass
- Z: Number of protons
- e: electron charge
- h: Planck constant
- $\epsilon_0$ : permittivity of free space
- c: speed of light
- R: Rydberg constant



# Hydrogen

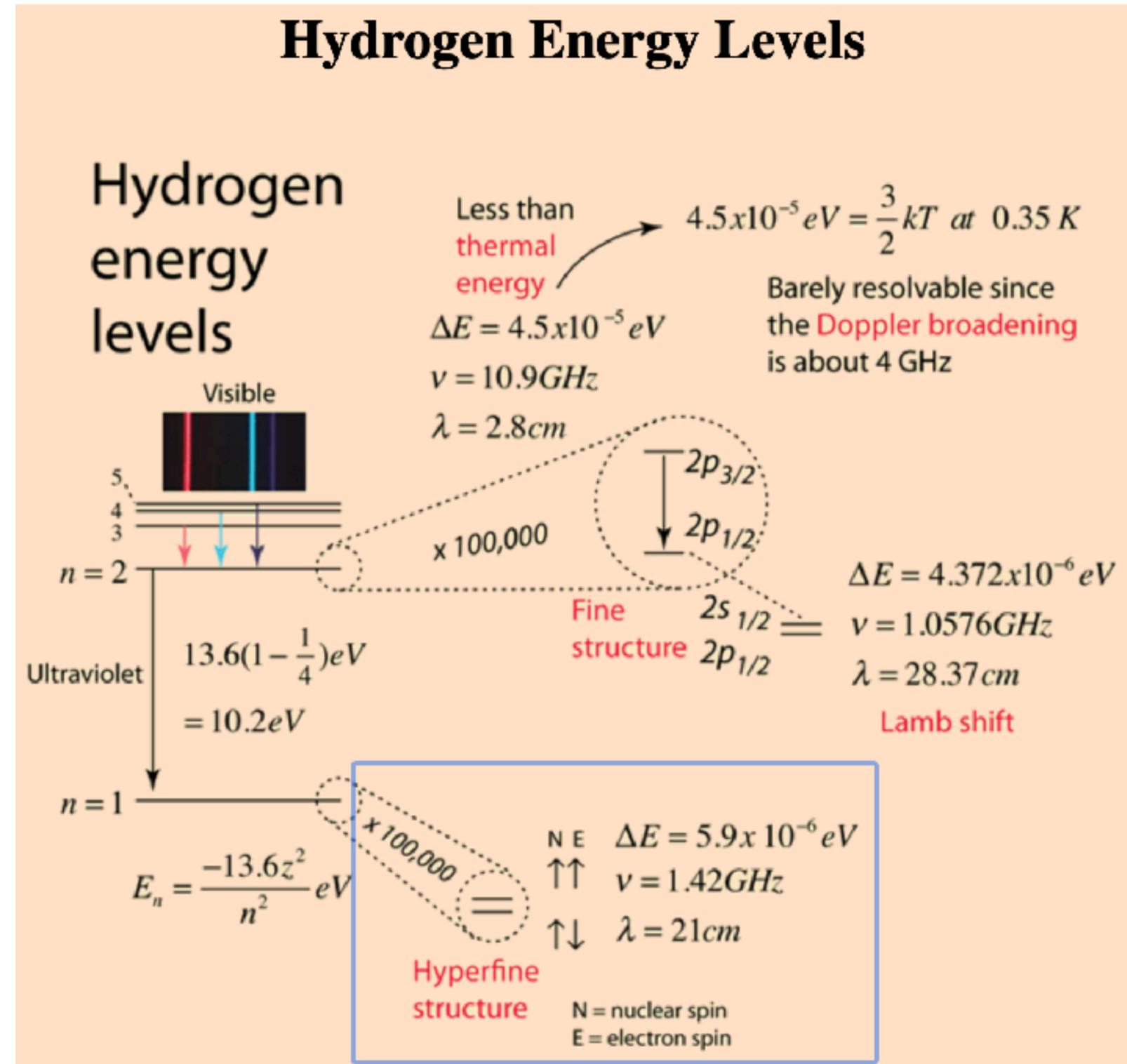
- Hydrogen spectrum comes from changes in  $n$  due to electrons jumping between different energy levels in the atom:

$$\frac{1}{\lambda} = \frac{1}{hc}(E_{n=1} - E_{n=2}) = R_H \left( \frac{1}{n_1^2} - \frac{1}{n_2^2} \right)$$



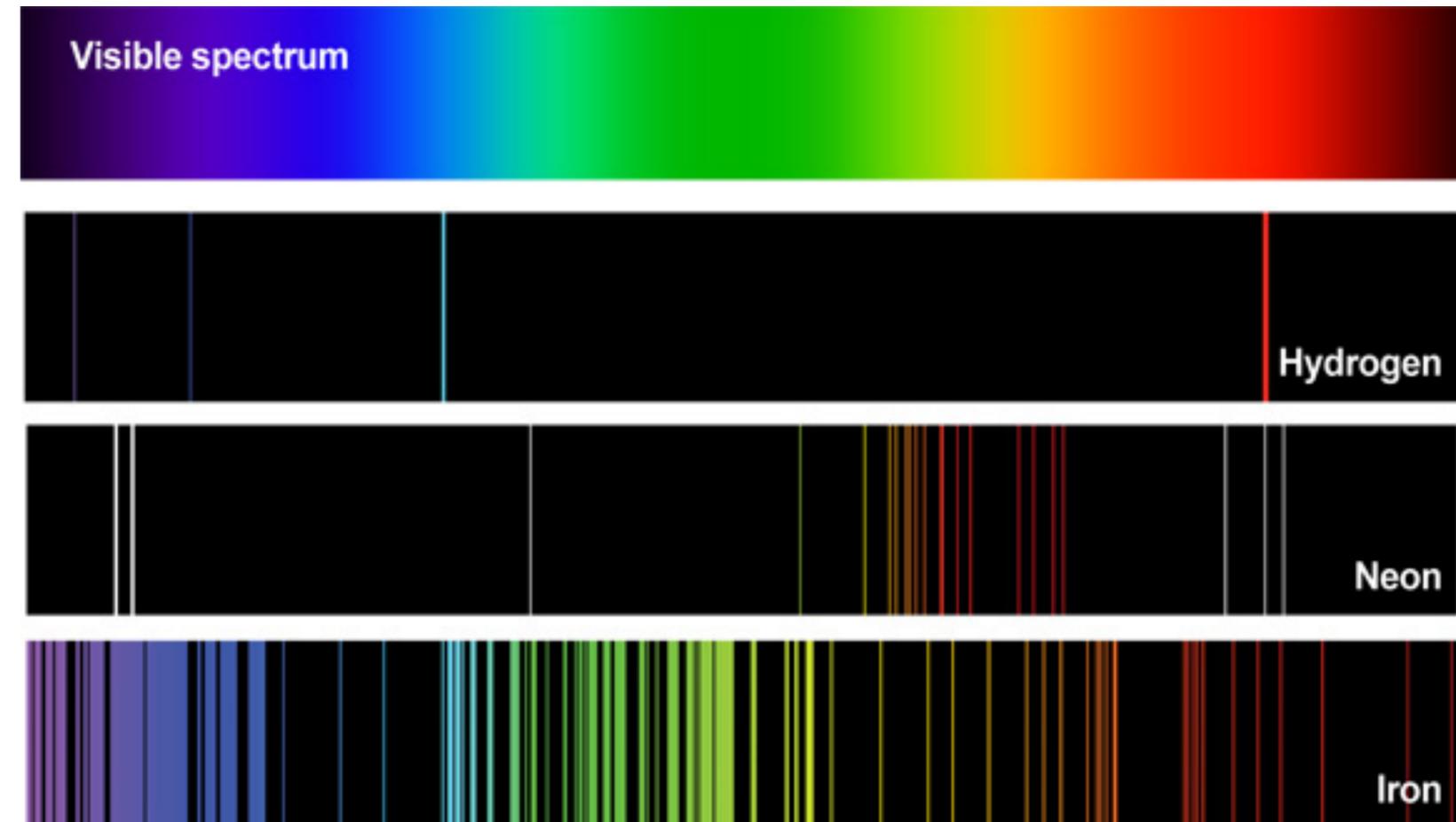
# Hydrogen

- Fine structure: interaction between electron spin and electron's orbital angular momentum
- Hyperfine structure: interaction between electron spin and nuclear spin



# Atomic Spectra

- Atomic lines dominate stellar spectra (except M stars, where you start to get molecules)
- These lines result from electron transitions between energy levels
  - Electron energy levels (and angular momentum) are quantized
- Highest energy transitions: electron energy levels
- Next-highest energy transitions: vibrational
- Smallest energy transitions: rotational

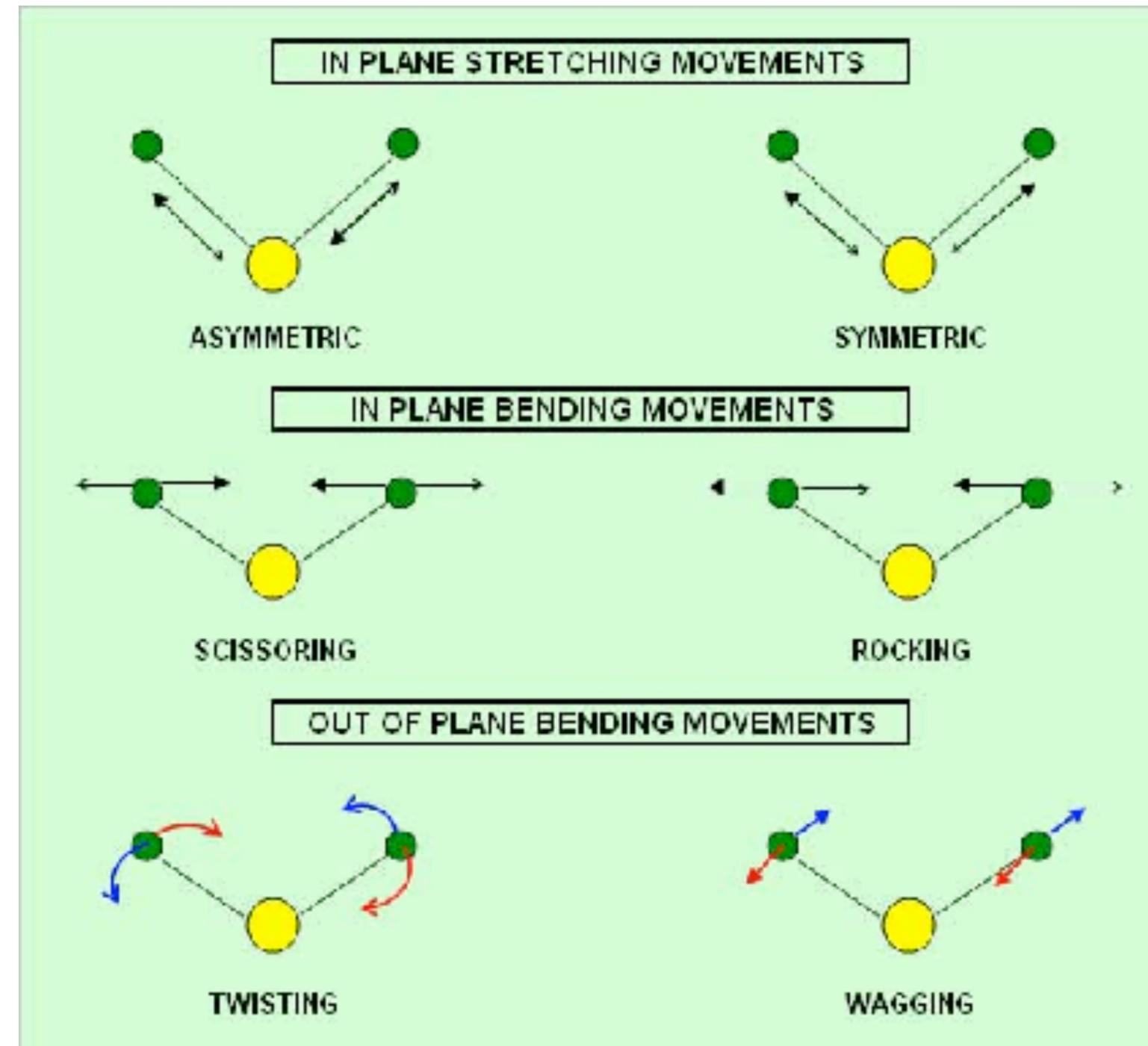


# Break

**05:00**

# Molecular Spectra

- The nuclei of molecules can move with respect to each other as well:
  - The whole molecule can rotate
  - The individual nuclei can vibrate (the bonds between them can stretch/contract and bend)
- These motions are also quantized (just like electron energy levels)
- Photons (which carry energy) are emitted or absorbed when a molecule moves from one rotation/vibration state to another



# Rotation of Molecules

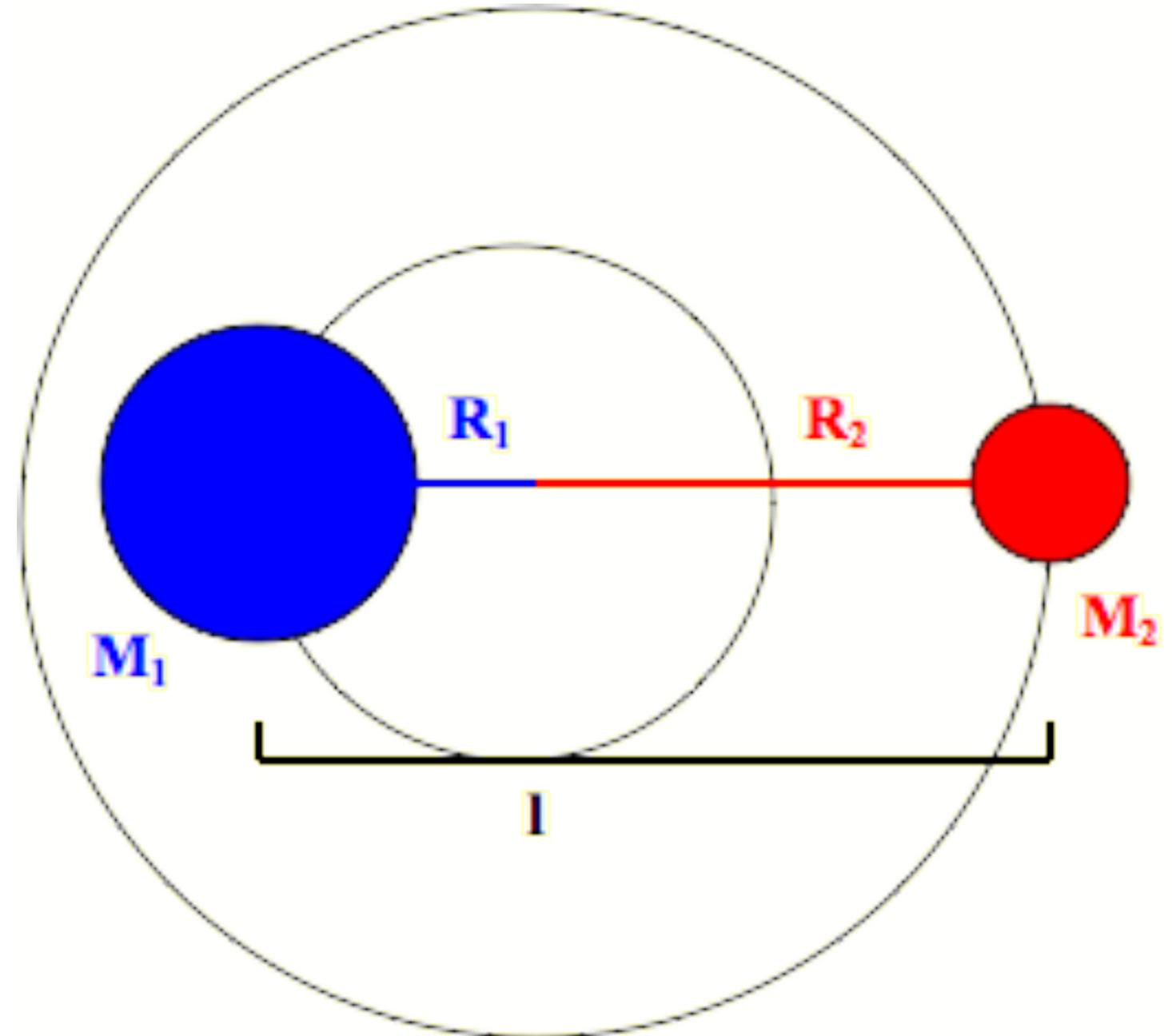
- Consider two (different) atoms in a molecule (CO, say)
- Moment of inertia for rotation about the center of mass:

$$I = M_1 R_1^2 + M_2 R_2^2$$

- Solving the Schrodinger equation, rotational energy levels are given by:

$$E_r = \frac{h}{8\pi^2 c I} J(J + 1) = B J(J + 1)$$

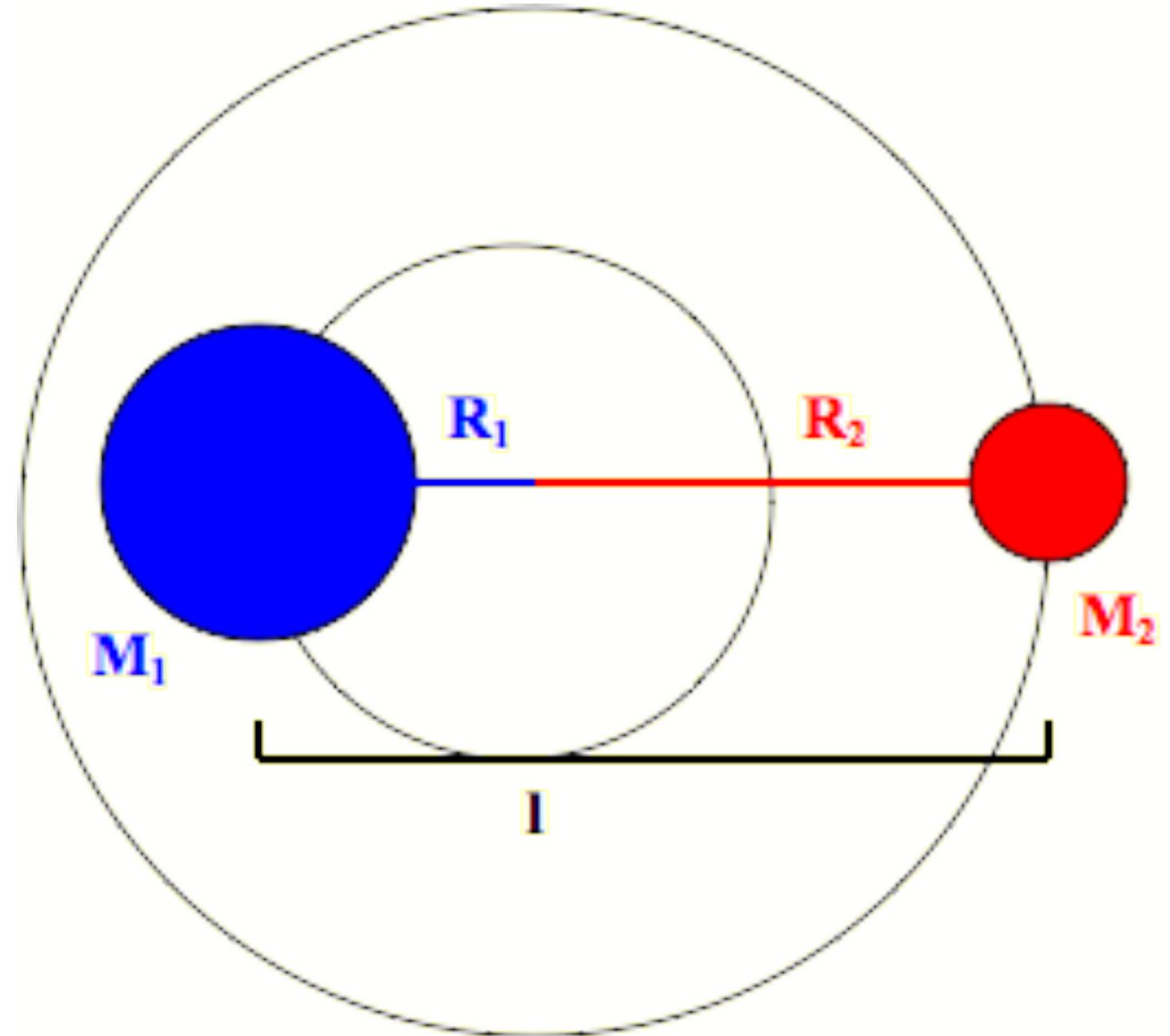
- $h$ : Planck's constant
- $J$ : rotational quantum number (an integer)
- $B$ : rotational constant
- $c$ : speed of light



# Rotation of Molecules

$$E_r = \frac{h}{8\pi^2 c I} J(J + 1) = B J(J + 1)$$

- Molecular transitions written as upper-lower (1-0, 17-16)
- If  $J=0$ , then  $E_r = 0$  (no rotational zero-point energy)
- A molecule will only show a pure rotation spectrum if it has a permanent dipole moment
- Rotational constant  $B$  has units of inverse-length, wavenumber



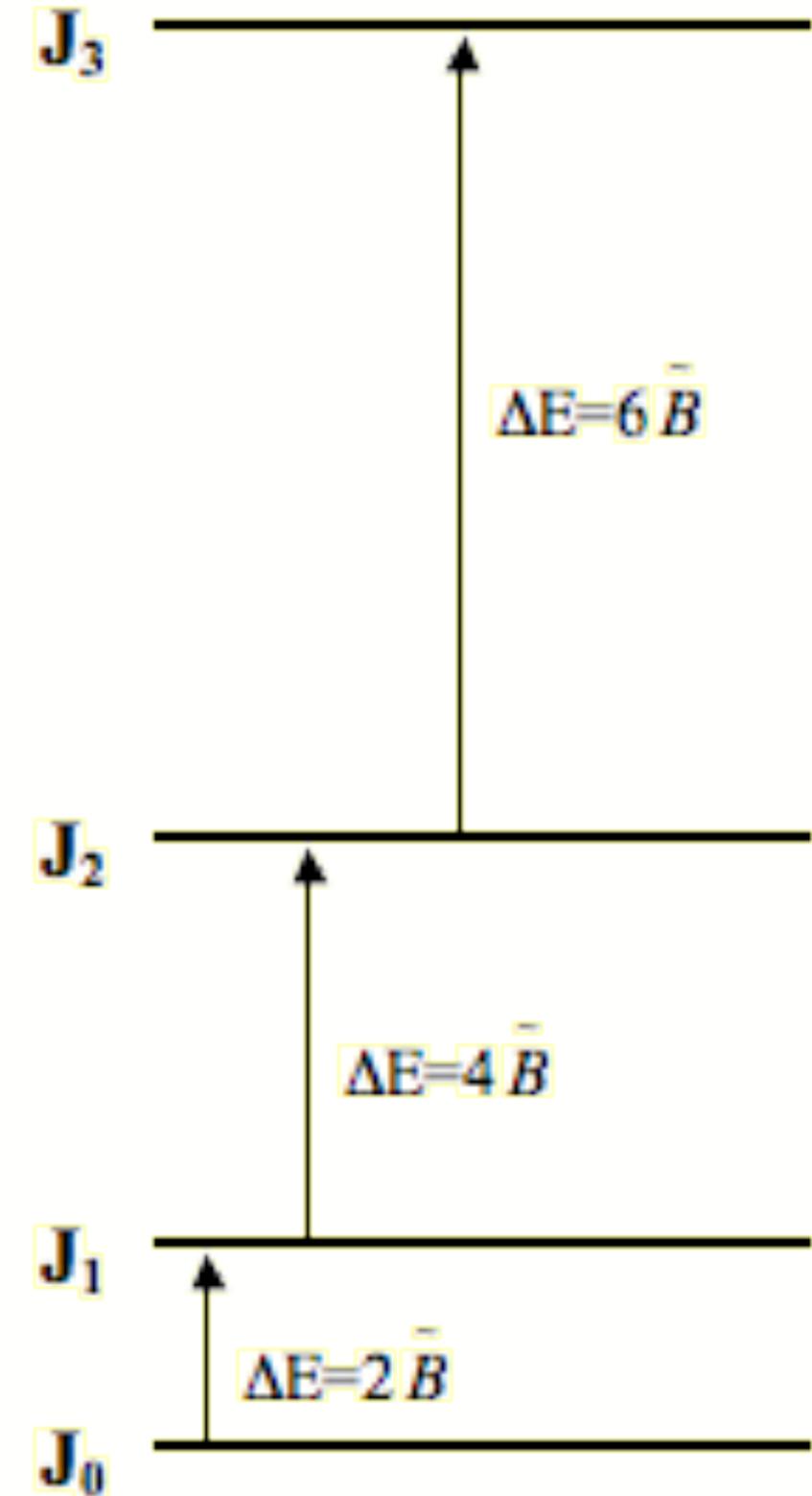
# Response Card Question

- A certain energy transition has a wavenumber of  $10^6 \text{ cm}^{-1}$ .  
What is the wavelength of the light emitted or absorbed from this transition?
  - (A) —  $3 \times 10^{16} \text{ cm}$
  - (B) —  $10^6 \text{ cm}$
  - (C) —  $1 \text{ cm}$
  - (D) —  $10^{-6} \text{ cm}$
  - (E) —  $3 \times 10^{-16} \text{ cm}$

# Rotation of Molecules

$$E_r = \frac{h}{8\pi^2 c I} J(J + 1) = B J(J + 1)$$

- Transitions are allowed when  $\Delta J = 1$
- What's the energy of the transition from state  $J+1$  to state  $J$ ?
- $\Delta E = E(J + 1) - E(J) = B(J + 1)(J + 2) - B J(J + 1)$   
 $\Delta E = B(J^2 + 3J + 2 - J^2 - J) = B(2J + 2)$   
 $\Delta E = 2B(J + 1)$
- So we expect lines at  $2B, 4B, 6B, 8B, \dots$
- Rotational lines are equally spaced in energy (and so frequency)



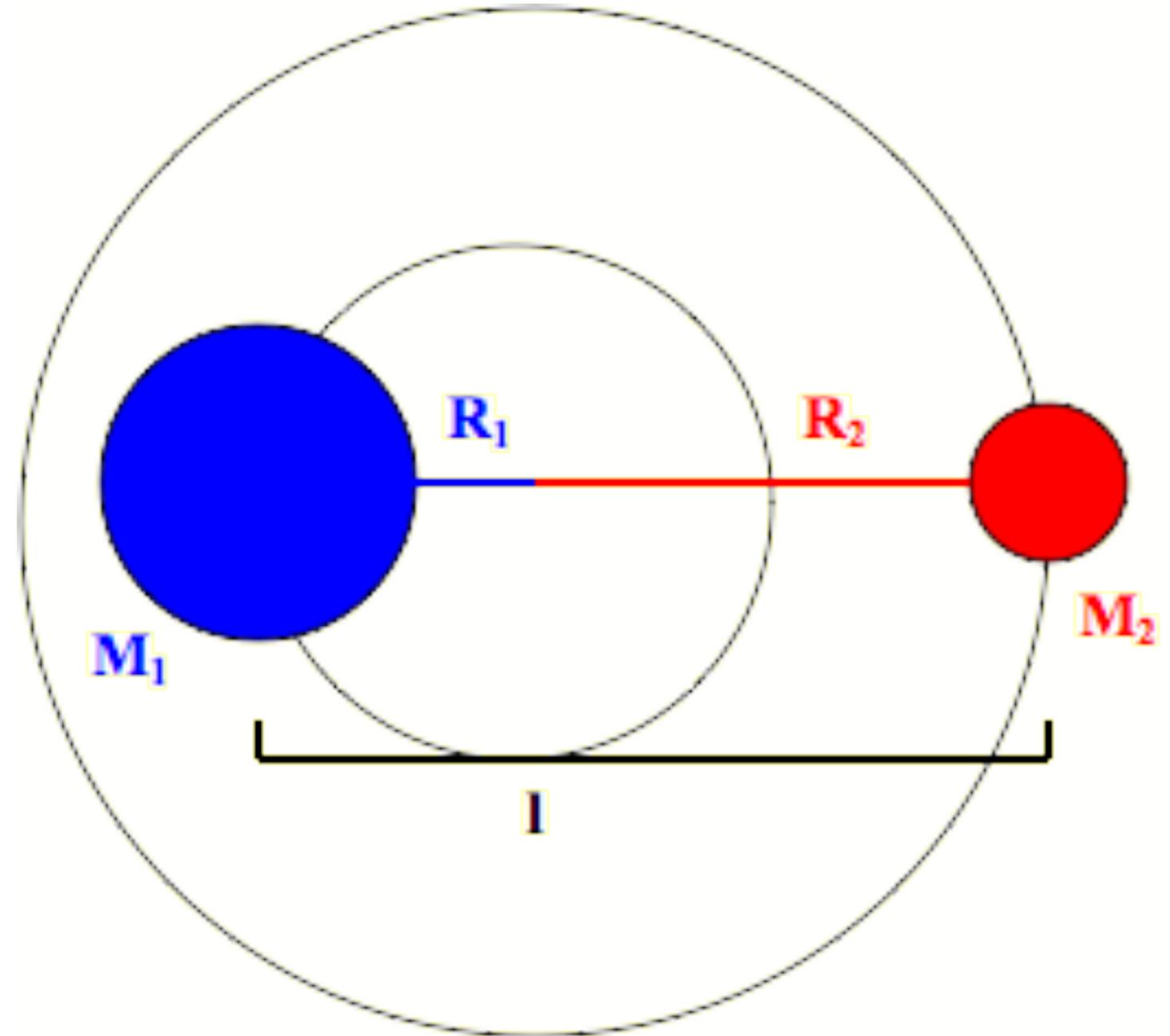
# Order of magnitude

$$I = M_1 R_1^2 + M_2 R_2^2 \quad m_p = 1.7 \times 10^{-27} \text{ kg}$$

$$E_r = \frac{h}{8\pi^2 c I} J(J+1) = B J(J+1) \quad h = 6.7 \times 10^{-34} \text{ Js}$$

$$\Delta E = 2B(J+1)$$

- Consider the molecule NO (nitrogen+oxygen)
- (1) (order-of-magnitude) what is the bond length  $R$  ( $R_1 + R_2$ ) between the two atoms? (hint: in a molecule, you can think of it as the two electron clouds touching each other)
- (2) What is the moment of inertia ( $I$ ) for NO? (hint: order of magnitude, so assume  $R_1 \approx R_2 \approx R$ )
- (3) What is  $B$  for NO?
- (4) What type of light would be emitted/absorbed for NO rotational transitions? (e.g., visible, X-ray, ...)



# Order of magnitude

- $I = M_1 R_1^2 + M_2 R_2^2$

$$E_r = \frac{h^2}{8\pi^2 c I} J(J+1) = B J(J+1)$$

$$\Delta E = 2B(J+1)$$

- Consider the molecule NO (nitrogen+oxygen)
- (1) (order-of-magnitude) what is the bond length R (R1+R2) between the two atoms? (hint: in a molecule, you can think of it as the two electron clouds touching each other)
- The typical size of an atom is about half an angstrom (round up to 1), 1 angstrom is  $10^{-10}$  m
- (2) What is the moment of inertia (I) for NO?
- Nitrogen has an atomic number of 7, Oxygen 8, assume the atomic weight is double that, 14 nucleons for N, 16 for O, or 30 total.

# Order of magnitude

- Since we assume the bond length is about the same for both:

- $I = (M_1 + M_2)R^2 = 30m_pR^2$

- Mass of the proton is  $10^{-27}$  kg, R is 1 angstrom,  $10^{-10}m$

- $I = 30(10^{-27}kg)(10^{-10}m)^2 = 3 \times 10^{-46}m^2kg$

- (3) What is B for NO?

- $$B = \frac{h}{8\pi^2cI} = \frac{(7 \times 10^{-34}m^2kg/s)}{8\pi^2(3 \times 10^8m/s)(3 \times 10^{-46}m^2kg)} = \frac{10^{-34}}{100 \times 10^{-38}}m^{-1} = \frac{1}{10^{-2}}m^{-1} = 100m^{-1}$$

- (4) Let's convert this to a wavelength by inverting:

- $$Wavelength = \frac{1}{Wavenumber} = \frac{1}{100m^{-1}} = 0.01m = 1cm \quad \text{Microwave!}$$

# For next time

- Reading: de Pater & Lissaeuer Chaper 4, section 4.1, 4.2 (intro only)
- Homework 3 due Wednesday, September 28 at 11:59pm