

$$\text{So } U_i = \epsilon_0 \beta \frac{1}{2} \cos 2\pi \nu t \cdot \omega_0 + \epsilon_0 \beta \frac{1}{2} \cos 2\pi \nu t (\nu - \nu_0) - \epsilon_0 \beta \frac{1}{2} \cos 2\pi \nu t (\nu + \nu_0)$$

$\epsilon_0 \beta \frac{1}{2} \cos 2\pi \nu t \cdot \omega_0 \Rightarrow$  Rayleigh ~~scattering~~ line.

$\epsilon_0 \beta \frac{1}{2} \cos 2\pi \nu t (\nu - \nu_0) \Rightarrow$  Stokes line

$\epsilon_0 \beta \frac{1}{2} \cos 2\pi \nu t (\nu + \nu_0) \Rightarrow$  anti-Stokes line.

### Quantum mechanical theory of Raman effect

- \* A quantum of radiation  $h\nu_i$  of incident radiation collide with the molecule. If it is scattered without any energy loss, the scattered light will have the same frequency as the incident light; the phenomenon is called Rayleigh scattering.
- \* If the incident light transfers its energy to the vibrational or rotational state in the form of quanta  $h\nu_m$ . The scattered light will have the slightly lower frequency than the incident light.

$$\boxed{\nu_{sc} = \nu_{in} - \nu_m} \quad \text{These are referred to as Stokes lines}$$

where  $\nu_{sc}$  = scattered frequency

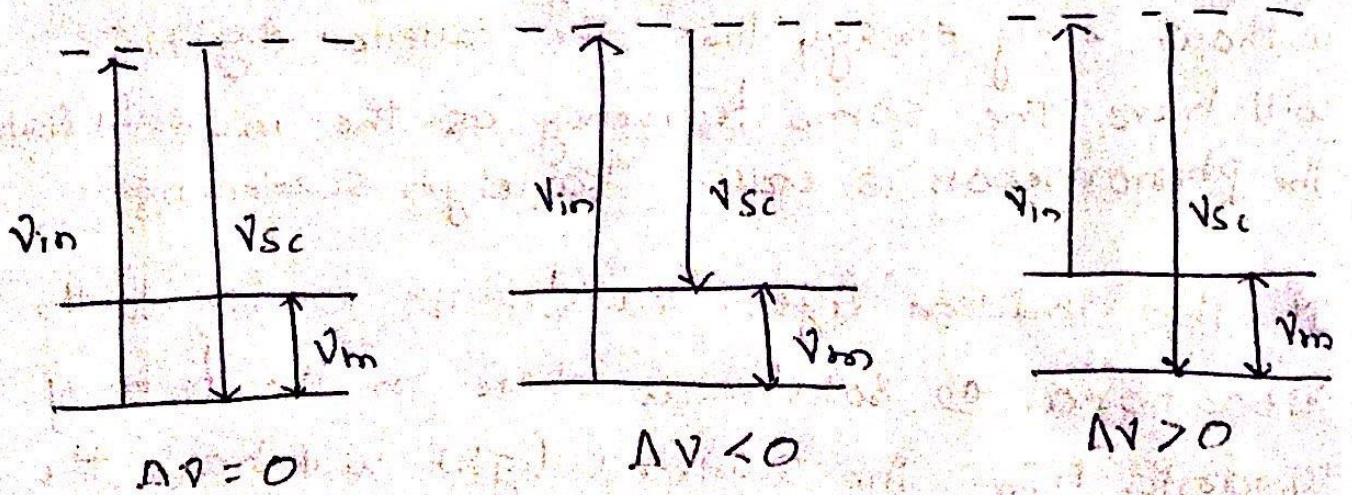
$\nu_{in}$  = Incident frequency

$\nu_m$  = Frequency equivalent to energy difference b/w two vibrational or rotational states of the molecule.

\* The incident light when goes to the excited state, It takes the energy of excited state and hence the scattered light has greater frequency than the incident light. If  $\nu_{in}$  (incident frequency) of light & scattered frequency ( $\nu_{sc}$ ) of light

then 
$$\nu_{sc} = \nu_{in} + \nu_m$$

These are referred to as anti-stokes lines. Both Stokes & anti-stokes lines are known as Raman scattering.



Rayleigh scattering

Stokes lines

Anti-stokes lines

Raman scattering.