The key feature of Bohr's theory of spectrum of hydrogen atom is the quantization of angular momentum when an electron is revolving around a proton. We will extend this to a general rotational motion to find quantized rotational energy of a diatomic molecule assuming it to be rigid. The rule to be applied is Bohr's quantization condition. (IIT JEE, 2010)

4. A diatomic molecule has moment of inertia *I*. By Bohr's quantization condition, its rotational energy in the *n*th level (n = 0 is not allowed) is

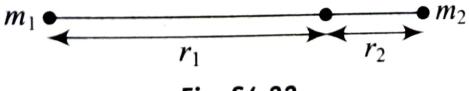
**a.** 
$$\frac{1}{n^2} \left( \frac{h^2}{8\pi^2 I} \right)$$
**b.** 
$$\frac{1}{n} \left( \frac{h^2}{8\pi^2 I} \right)$$
**c.** 
$$n \left( \frac{h^2}{8\pi^2 I} \right)$$
**d.** 
$$n^2 \left( \frac{h^2}{8\pi^2 I} \right)$$

- 5. It is found that the excitation frequency from ground to the first excited state of rotation for the CO molecule is close to  $4/\pi \times 10^{11}$  Hz. Then the moment of inertia of CO molecule about its center of mass is close to (Take  $h = 2\pi \times 10^{-34}$  J s)
  - **a.**  $2.76 \times 10^{-46} \text{ kg m}^2$  **b.**  $1.87 \times 10^{-46} \text{ kg m}^2$  **c.**  $4.67 \times 10^{-47} \text{ kg m}^2$ **b.**  $1.17 \times 10^{-47} \text{ kg m}^2$
- 6. In a CO molecule, the distance between C (mass = 12 a.m.u) and O (mass = 16 a.m.u.), where 1 a.m.u  $5/3 \times 10^{-27}$  kg, is close to
  - **a.**  $2.4 \times 10^{-10}$  m **b.**  $1.9 \times 10^{-10}$  m **c.**  $1.3 \times 10^{-10}$  m **d.**  $4.4 \times 10^{-11}$  m

 $4. d. L = \frac{nh}{2\pi}$ 

K.E. = 
$$\frac{L^2}{2I} = \left(\frac{nh}{2\pi}\right)^2 \frac{1}{2I}$$

5. a. 
$$hv = KE_{n=2} - kE_{n=1}$$
  
 $I = 1.87 \times 10^{-46} \text{ kg m}^2$   
6. c.  $r_1 = \frac{m_2 d}{m_1 + m_2}$  and  $r_2 = \frac{m_1 d}{m_1 + m_2}$ 





$$I = m_1 r_1^2 + m_2 r_2^2$$
  
$$\therefore \quad d = 1.3 \times 10^{-10} \text{ m}$$