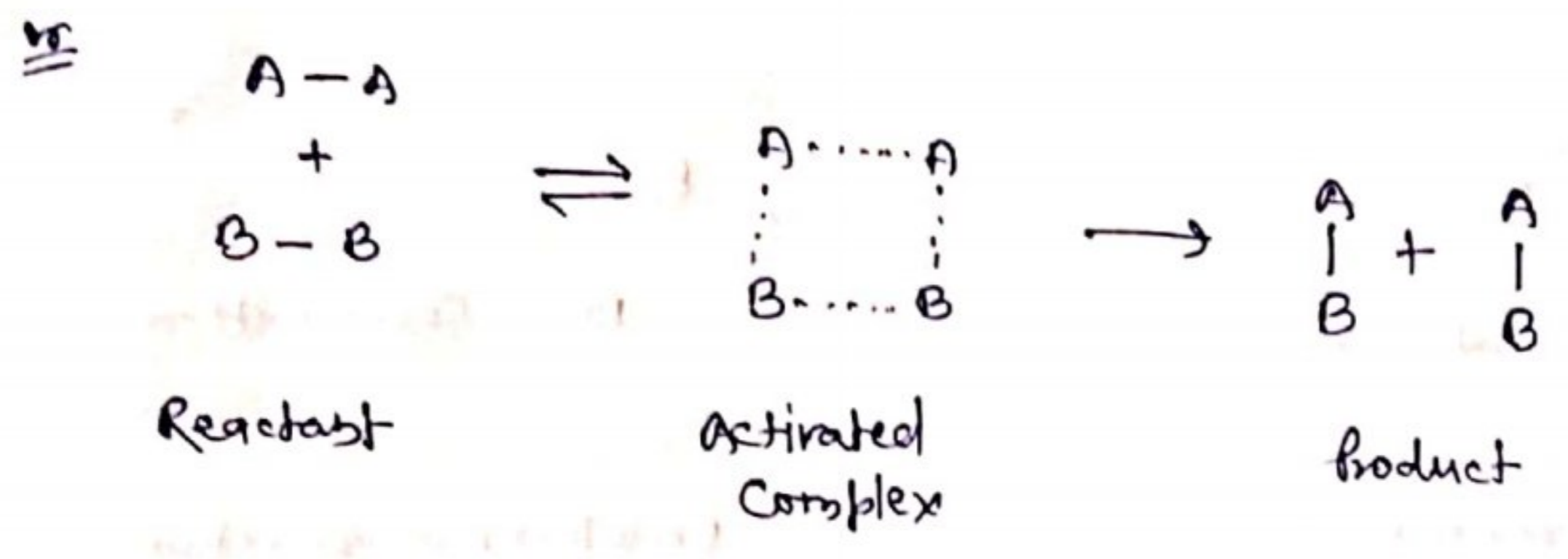
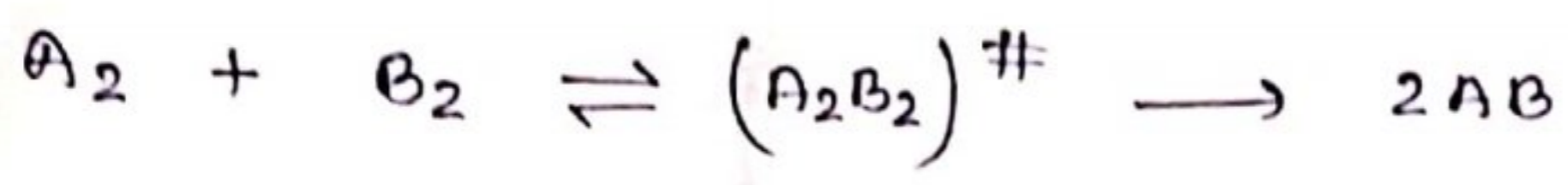


\* Potential Energy diagram and Concept of activated complex :-

A/c to activated Complex theory (ACT), the bimolecular reaction between two molecules A<sub>2</sub> & B<sub>2</sub> progresses through the formation of the so-called activated complex which then decomposes to yield the product AB -



Threshold Energy :-

The minimum energy which the reactant molecules should possess so that they may react to produce the product is called threshold energy.

Activation energy :- (E<sub>a</sub>)

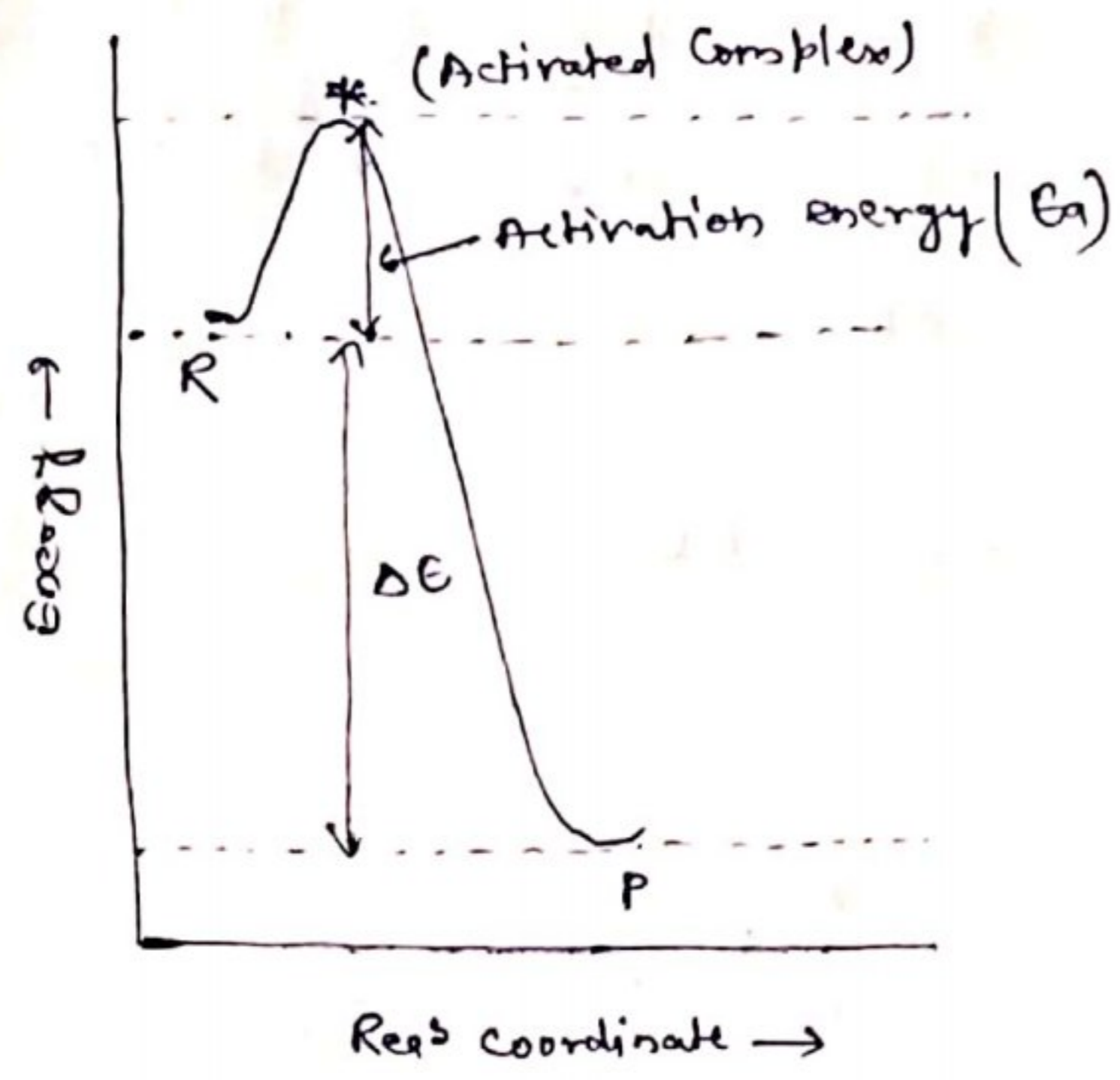
Reactant molecules possess some energy initially also. The minimum extra energy given to the reactant molecules to make their energy equal to threshold energy is called Activation energy.

Thus,

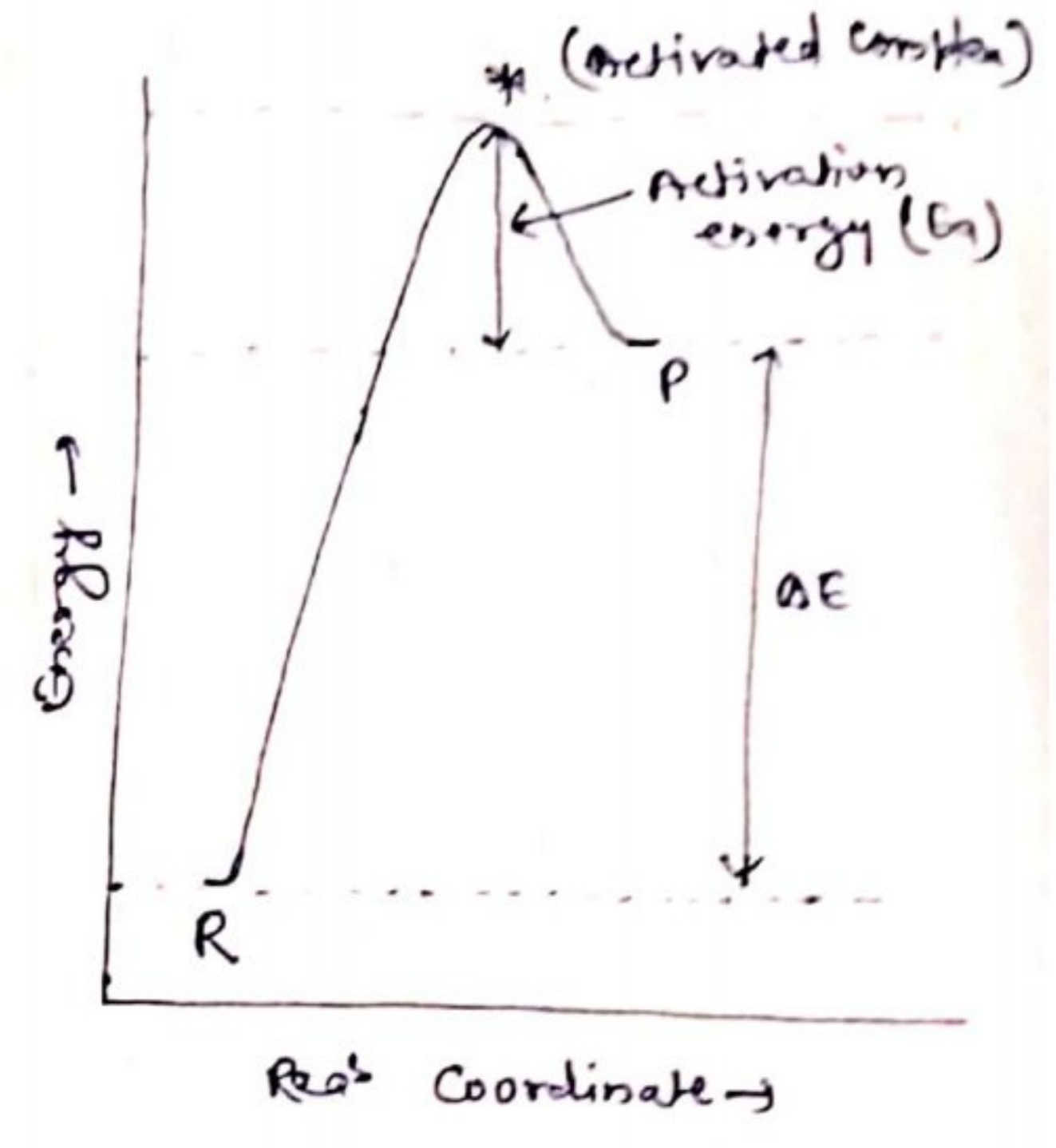
$$\text{Activation energy} = \text{Threshold energy} - \text{Energy possessed by molecule.}$$



A Curve is obtained by potential energy vs reaction co-ordinate. Reaction coordinate represents the profile energy change when reactants change into products.



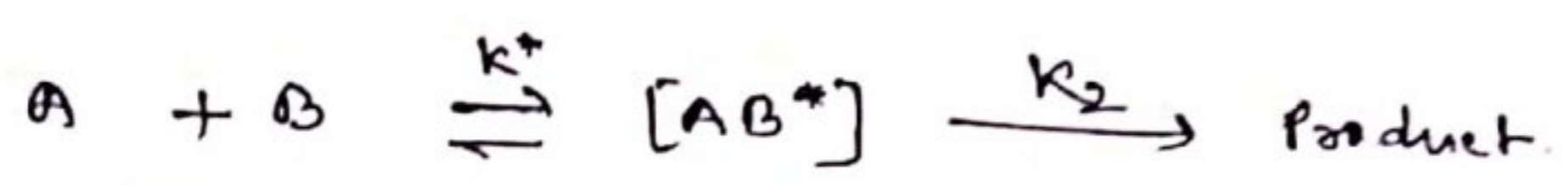
Exothermic reaction



Endothermic reaction.

\* Rate data and Mechanism of a chemical Reaction :-

For thermodynamic formulation of the activated complex theory, let us consider a simple bimolecular reaction -



Where,  $[AB^*]$  is called Activated Complex

$k^*$  is the equilibrium constant between reactant and activated complex.

From classical mechanics, the energy of vibrations is given by -  $RT/N_A$ .



and from quantum mechanics, it is given by  $h\nu$ .

Where,

$$h\nu = RT/N_A$$

$$\nu = RT/N_A h$$

Thus, the reaction rate is given by -

$$-\frac{d[A]}{dt} = k k_2 [AB^*] = k (k_B T/h) [AB^*] \quad \text{--- (1)}$$

where, the factor  $k$ , called transmission coefficient.

Now, at equilibrium -

$$K^* = \frac{[AB^*]}{[A][B]}$$

$$\text{or } [AB^*] = K^* [A][B] \quad \text{--- (2)}$$

Now putting this value in eq<sup>n</sup> (1)

$$-\frac{d[A]}{dt} = (k_B T/h) K^* [A][B] \quad \text{--- (3)}$$

Thus the rate constant  $k_2$  may be expressed as -

$$k_2 = (k_B T/h) K^* \quad \text{--- (4)}$$

The equilibrium constant  $K^*$  can be expressed in terms of  $(\Delta G^\circ)^*$  called the standard Gibbs free energy of activation.

Thus,

$$(\Delta G^\circ)^* = -RT \ln K^*$$

$$\text{or } (\Delta G^\circ)^* = (\Delta H^\circ)^* - T(\Delta S^\circ)^*$$



Thus, we obtained

$$k^* = e^{-(\Delta G^\ddagger)/RT}$$

$$= e^{(\Delta S^\ddagger)/R} e^{-(\Delta H^\ddagger)/RT}$$

Hence, Eq<sup>s</sup> - (4) becomes -

$$k_2 = \left( k_B T / h \right) e^{(\Delta S^\ddagger)/R} e^{-(\Delta H^\ddagger)/RT} \quad \text{--- (5)}$$

Eq<sup>s</sup> - (5) is well known Eyring equation.

&  $(\Delta S^\ddagger)^*$  = standard entropy of activation.

$(\Delta H^\ddagger)^*$  = standard enthalpy of activation.

From  


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