

## Density of distribution in phase space :-

The use of ensembles in st. mechanics is guided by the following factors :-

- ① There is no need to maintain distinction between individual systems as we are only interested in number of systems at any time which would be found in different state that corresponding different phase space.
- ② The no. of elements in an ensemble is so large that there is a continuous change in this number in passing from one region of phase to another.

The condition of an ensemble at any time can be specified by the density of with the phase points are distributed over the phase space named as density of distribution or Probability Density of Distribution Function.

The density of distribution  $f$  or  $\rho$  is the function of  $f$  atoms position coordinates and  $f$ -momentum co-ordinate i.e.  $(q_1, q_2, \dots, q_f)$  and  $(P_1, P_2, \dots, P_f)$  corresponding to  $2f$  axes in phase space.

The density of distribution is also a function of time because at any time  $t$

$$f = f(q_1, \dots, q_f; p_1, p_2, \dots, p_f; t)$$

$$\Rightarrow f = f(q, p, t)$$

The small volume of the phase space named as Hyper-volume  $\delta\Gamma$  of the phase points

$$\delta\Gamma = \delta q_1 \delta q_2 \dots \delta q_f; \delta p_1 \dots \delta p_f$$

The number of systems  $\delta N$  lying in the specified region can be obtained by multiplying the density of distribution and the Hyper-volume in the phase space region.

$$\delta N = f(q, p, t) \delta q_1 \dots \delta q_f, \delta p_1 \dots \delta p_f$$

$$\delta N = f \delta q_1 \dots \delta q_f \delta p_1 \dots \delta p_f$$

In brief

$$\delta N = f \prod_{i=1}^f \delta q_i \delta p_i$$

By integrating over the whole of the phase space

$$N = \underbrace{\int f dq_1 \dots dq_f dp_1 \dots dp_f}_{X}$$

General discussion of Mean values:-

by

The average value of variable  $u$  in an ensemble is defined by multiplying each possible  $u_i$  by the number  $w_i$  of the systems in the ensemble which exhibit this value, adding the resultant product for all possible values of the variable  $u$  and then dividing this sum by the total number of systems in the ensemble.

Let  $u$  be the variable which can assume any of the  $M$  discrete values.

$$u_1, u_2, \dots, u_M$$

with respective probabilities

$$w(u_1), \dots, w(u_M)$$

Then mean (or average) value of  $u$  is denoted by  $\bar{u}$  and is defined by

$$\bar{u} = \frac{w(u_1)u_1 + w(u_2)u_2 + \dots + w(u_M)u_M}{w(u_1) + w(u_2) + \dots + w(u_M)}$$

$$\bar{u} = \frac{\sum_{i=1}^M w(u_i) u_i}{\sum_{i=1}^M w(u_i)}$$

More generally, if  $f(u)$  is any function of  $u$  then the mean value  $f(u)$  is denoted

by

$$\bar{f}(u) = \frac{\sum_{i=1}^M w(u_i) f(u_i)}{\sum_{i=1}^M w(u_i)}$$

This expression can be simplified as unity,

$$\sum_{i=1}^M w(u_i) = w(u_1) + w(u_2) + \dots + w(u_M) = 1$$

$$\therefore \bar{f}(u) = \sum_{i=1}^M w(u_i) f(u_i)$$

As the probability distribution function is continuous in all position and momentum co-ordinates the above eqn +

$$\bar{w} = \frac{\int u(a, p) w(a, p) d\Gamma}{\int w(a, p) d\Gamma}$$

where  $d\Gamma = da_1 \dots da_M, dp_1 \dots dp_M$ .

But according to normalisation condition

$$\int w(a, p) d\Gamma = 1$$

$$\bar{w} = \int u(a, p) w(a, p) d\Gamma$$

As  $f(u)$  and  $g(u)$  are any two functions of  $u$ , then

$$\begin{aligned}\bar{f(u)+g(u)} &= \sum_{i=1}^M w(u_i) [f(u_i) + g(u_i)] \\ &= \sum_{i=1}^M w(u_i) f(u_i) + \sum_{i=1}^M w(u_i) g(u_i)\end{aligned}$$

Suppose we have to determine the average distance from the origin with total points  $N$

$$\bar{x} = \frac{\sum_{i=1}^N x(i)}{N}$$

Now, if the line is divided into cell, and the distribution is given in terms of the number of points in each cell, then

$$\bar{x} = \frac{\sum_i N_i x(i)}{N}$$

$$\Rightarrow \bar{x} = \frac{\int_{-\alpha}^{\alpha} x n(x) dx}{N}, \text{ noting that}$$

$$\int_{-\alpha}^{\alpha} n(x) dx = N$$

$$\bar{x} = \frac{\int_{-\alpha}^{\alpha} x n(x) dx}{\int_{-\alpha}^{\alpha} n(x) dx}$$

Macroscopic behaviour as an average over microscopic

behaviour :- In macroscopic point of view the

thermodynamic probabilitis remain constant but in microscopic point of view we never stay constant. one can only determine the probability for the set of all possible microscopic states of the system.

The microscopic value of  $x$  is represented by  $x(a, p) = x(p)$  where  $p$  is the phase point

The observed value  $x_{\text{obs}}$  in macroscopic sense should be the average value of microscopic  $x$ .

$$x_{\text{obs}} = \bar{x}$$

Let  $M$  is the set of all microscopic states which can be realized by the system under a given macroscopic condition and probability  $d\omega$  over the volume element  $d\Gamma$

$$d\omega = \int p(p) d\Gamma, \quad \text{where } d\Gamma = da_1 \dots da_n dr_1 \dots dr_n$$

$d\Gamma$  belongs to  $M$  and  $p$  is probability density

and whole phase

$$\int p d\Gamma = 1$$

$$\therefore \bar{x} = \int_M x(p) p(p) d\Gamma$$

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## LOUVILLE'S THEOREM

In dynamical state of a system the point in phase space is not stationary but it will move along a definite trajectory path which is determined from the equation of motion,

$$\dot{q}_i = \frac{\partial H}{\partial P_i} \quad \text{and} \quad \dot{P}_i = -\frac{\partial H}{\partial q_i}$$

Where  $H = H(q_1, \dots, q_8, P_1, \dots, P_8)$  is the Hamiltonian of the system.

As a result of this motion the density  $\rho$  of system in phase space changes with time.

In finding  $\frac{d\rho}{dt}$  at a given point in phase

Liouville's made theorem -

This theorem is primarily concerned with defining a fundamental property of the phase space i.e. space for position & momentum coordinate and hence consists of two part

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$$\frac{dp}{dt} = 0$$

2nd Part

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First Point :- The first point states the conservation of density in phase space ie

$$\frac{dp}{dt} = 0.$$

2nd Point :- The second point gives the conservation of extension in phase space ie

$$\frac{d(S_f)}{dt} = 0$$

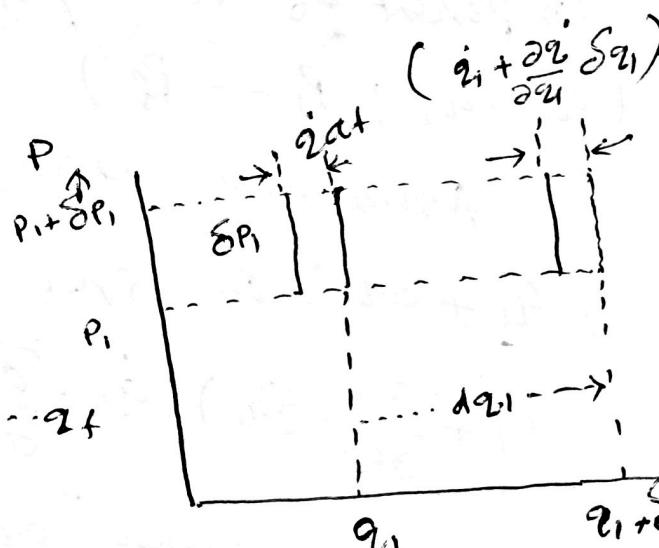
or the volume at the disposal of a particular number of phase points is conserved through the phase space.

Prob. of 1st Part :-

Consider any fixed element of volume of phase space  $P_i$  located with  $q_1 \in q_1 + \delta q_1 \dots q_f$  and  $P_i \in P_i + \delta P_i, \dots, P_f$

The no. of volume

$$(\delta q_1 \dots \delta q_f; \delta P_1, \dots \delta P_f)$$



The change in number of systems within volume of phase space is given by

$$\left( \frac{\delta q}{\delta t} \right) dt (\delta p_i - \delta p_f)$$

- (P)

This change is due to the number of system entering and leaving this volume in time  $\delta t$ . Let two faces of hyper volume normal to the  $q_1$  axis with  $q_1$  and  $q_1 + \delta q_1$ . Hence the no. of phase points entering the first face in time will be

$$p_i q_1 dt (\delta q_2 \dots \delta q_f) (\delta p_1 - \delta p_f)$$

where  $p$  and  $q_i$  are the density and indicated component of velocity for representative points  $(q_1 - q_f)$ .

$$(q_1 - q_f; p_1 - p_f)$$

Again, the phase points leaving the face  $(q_1 + \delta q_1)$  in time  $\delta t$  will be

$$(p + \frac{\partial p}{\partial q_1} \delta q_1) (q_1 + \frac{\partial q_1}{\partial p} \delta p_1) \delta q_2 \dots \delta q_f \delta p_1 - \delta p_f - ①$$

Neglecting higher order terms, we have

$$\left[ p q_1 + \left( p \frac{\partial q_1}{\partial q_1} + q_1 \frac{\partial p}{\partial q_1} \right) \delta q_1 \right] dt \delta q_2 \dots \delta q_f \delta p_1 - \delta p_f - ②$$

Substituting ② from eqn ①, we have

The

$$-(P \frac{\partial q_i}{\partial p_1} + q_i \frac{\partial P}{\partial q_1}) dt \delta q_1 \dots \delta q_f \delta p_1 \dots \delta p_f$$

likewise for  $p_i$  coordinate, we have

$$-(P \frac{\partial p_i}{\partial p_1} + p_i \frac{\partial P}{\partial p_1}) dt \delta q_1 \dots \delta q_f \delta p_1 \dots \delta p_f$$

The net increase in time  $dt$  of number of system in this volume of phase space is then obtained by summing the net number of system entering the volume through all the faces labelled by  $q_1 \dots q_f$  and  $p_1 \dots p_f$ . Then

$$\frac{d(\delta N)}{dt} = - \sum_{i=1}^f \left\{ p \left( \frac{\partial q_i}{\partial p_1} + \frac{\partial p_i}{\partial p_1} \right) + \left( \frac{\partial P}{\partial q_i} q_i + \frac{\partial P}{\partial p_i} p_i \right) \right\} dt \delta q_1 \dots \delta q_f \delta p_1 \dots \delta p_f \quad (3)$$

$$\text{Now } \frac{d(\delta N)}{dt} = \frac{\partial P}{\partial t} dt \delta q_1 \dots \delta q_f \delta p_1 \dots \delta p_f$$

$$\frac{\partial P}{\partial t} dt \delta q_1 \dots \delta q_f \delta p_1 \dots \delta p_f \\ = - \sum_{i=1}^f \left\{ p \left( \frac{\partial q_i}{\partial p_1} + \frac{\partial p_i}{\partial p_1} \right) + \left( \frac{\partial P}{\partial q_i} q_i + \frac{\partial P}{\partial p_i} p_i \right) \right\} dt \delta q_1 \dots \delta q_f \delta p_1 \dots \delta p_f$$

$$\frac{\partial P}{\partial t} = - \sum_{i=1}^f \left\{ p \left( \frac{\partial p_i}{\partial p_1} + \frac{\partial q_i}{\partial p_1} \right) + \left( \frac{\partial P}{\partial q_i} q_i + \frac{\partial P}{\partial p_i} p_i \right) \right\} \quad (3)$$

The equations of motion in canonical form

$$i = \frac{\partial H}{\partial t} \quad \text{and} \quad \dot{i} = -\frac{\partial H}{\partial \dot{t}}$$

$$\frac{\partial \dot{i}}{\partial t} = \frac{\partial^2 H}{\partial t \partial \dot{t}}$$

Since we have the initial condition

$$\therefore \sum_{i=1}^n \left( \frac{\partial \dot{i}}{\partial t} + \frac{\partial^2 H}{\partial t \partial \dot{t}} i \right) = 0 \quad \rightarrow \textcircled{4}$$

Substituting  $\dot{i} = 0$  in  $\textcircled{4}$

$$\left( \frac{\partial^2 H}{\partial t \partial \dot{t}} \right)_{\dot{t}=0} = - \sum_{i=1}^n \left( \frac{\partial^2 H}{\partial t \partial i} i \right) \rightarrow \textcircled{5}$$

This result is known as Liouville's theorem hence  
eqn  $\textcircled{5}$  can be written as

$$\left( \frac{\partial^2 H}{\partial t \partial \dot{t}} \right)_{\dot{t}=0} + \sum_i \frac{\partial^2 H}{\partial t \partial i} i + \sum_i \frac{\partial^2 H}{\partial \dot{t} \partial i} \dot{i} = 0 \rightarrow \textcircled{6}$$

and is identical with the equation of continuity in  
hydrodynamics if  $H$  is a function of  $t, p, \dot{t}$  and  
 $\partial H / \partial \dot{t} = 0$  and the function of  $H$  (then  
 $\partial H / \partial \dot{t} \neq 0$ )

$$\frac{dp}{dt} = \frac{\partial H}{\partial t} + \sum_i \frac{\partial H}{\partial i} i + \frac{\partial H}{\partial \dot{t}} \dot{t}$$

Generalising this for all  $f(t)$  or  $f(\dot{t})$  we get

$$\frac{df}{dt} = \frac{\partial f}{\partial t} + \sum_i \frac{\partial f}{\partial i} i + \frac{\partial f}{\partial \dot{t}} \dot{t} \quad \textcircled{7}$$

comparing the eqn (6) & (7) we have that

$$\frac{df}{dt} = 0$$

This form of expression may be called the principle of the conservation of density in phase space. Therefore, the density of a group of points remains constant along their trajectory in the phase space. i.e. will never have uniform density!

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2nd Part :- Since we have

$$\Sigma N = \rho \Sigma \Gamma$$

$$\Rightarrow \frac{d}{dt} (\Sigma N) = \frac{d\rho}{dt} \Sigma \Gamma + \rho \frac{d(\Sigma \Gamma)}{dt}$$

Since the number of phase points  $\Sigma N$  in a given region of the phase space must remain constant. fixed, as the system can neither be created nor destroyed, we have

Hence

$$\frac{d}{dt} (\rho \Sigma \Gamma) = 0$$

$$\Rightarrow \frac{d\rho}{dt} (\Sigma \Gamma) + \rho \frac{d(\Sigma \Gamma)}{dt} = 0$$

We have proved that  $\frac{d\rho}{dt} = 0$ , Hence it follows

that,  $\rho \neq 0$

$$\therefore \frac{d}{dt} (\rho \Sigma \Gamma) = 0$$

This equation gives the principle of conservation of extension in phase space //

## Postulate of equal A Priori Probability. —

So far the methods used in st. mechanics are chiefly based on the principle of classical mechanics. In order to make applications to situations of actual interest we must now understand a consideration of the fundamental hypothesis as to equal a "PRIORI PROBABILITIES" for equal regions in phase space. This postulate doesn't arise due to any inadequacy in the principles of classical mechanics but due to the incompleteness of our knowledge concerning the systems.

As to this postulate the probability of finding the phase point for given systems in any region of the phase space is identical with that for any other region of equal extension or volume.