## Unit 2-6: Entropy of Mixing and the Gibbs Paradox

Consider two different gases (red gas and blue gas) at the same temperature and pressure, separated by a partition. The total volume $V$, total number of particles $N$, and total energy $E$ are all fixed to be constant.

$$
\begin{array}{lll}
E_{1} & E_{2}  \tag{2.6.1}\\
V_{1} & N_{2} \\
N_{1} & V_{2} \\
\hline
\end{array}
$$

$$
\left.\begin{array}{l}
V=V_{1}+V_{2} \\
N=N_{1}+N_{2} \\
E=E_{1}+E_{2}
\end{array}\right\} \text { constant }
$$

Since both gases are at the same $T$ and $p$ we have,

$$
\begin{array}{ll}
E_{1}=\frac{3}{2} N_{1} k_{B} T, & V_{1}=\frac{N_{1} k_{B} T}{p} \\
E_{2}=\frac{3}{2} N_{2} k_{B} T, & V_{2}=\frac{N_{2} k_{B} T}{p} \tag{2.6.3}
\end{array}
$$

With the partition in place, the total entropy is initially,

$$
\begin{equation*}
S_{\mathrm{init}}=k_{B} \ln \left[\Omega_{1}\left(E_{1}, V_{1}, N_{1}\right) \Omega_{2}\left(E_{2}, V_{2}, N_{1}\right)\right]=S_{1}\left(E_{1}, V_{1}, N_{1}\right)+S_{2}\left(E_{2}, V_{2}, N_{2}\right) \tag{2.6.4}
\end{equation*}
$$

Now we remove the partition and let the two gases mix. The temperature $T$ and the number of particles in each gas $N_{1}$ and $N_{2}$ should not change $\Rightarrow \quad E_{1}$ and $E_{2}$ remain constant. The only changes are $V_{1} \rightarrow V$ and $V_{2} \rightarrow V$.

With the partition removed, when the system comes into equilibrium the final entropy is,

$$
\begin{equation*}
S_{\text {final }}\left(E, V, N_{1}, N_{1}\right)=k_{B} \ln \left[\Omega_{1}\left(E_{1}, V, N_{1}\right) \Omega_{2}\left(E_{2}, V, N_{2}\right)\right]=S_{1}\left(E_{1}, V, N_{1}\right)+S_{2}\left(E_{2}, V, N_{2}\right) \tag{2.6.5}
\end{equation*}
$$

The entropy of mixing is

$$
\begin{equation*}
\Delta S=S_{\mathrm{final}}-S_{\mathrm{init}} \tag{2.6.6}
\end{equation*}
$$

If we use our result for the entropy of the ideal gas, computed from the microcanonical ensemble of the last section, we get,

$$
\begin{equation*}
S_{\mathrm{init}}=\frac{3}{2} k_{B} N_{1}+k_{B} N_{1} \ln \left[\frac{V_{1}}{h^{3}}\left(\frac{4 \pi m_{1} E_{1}}{3 N_{1}}\right)^{3 / 2}\right]+\frac{3}{2} k_{B} N_{2}+k_{B} N_{2} \ln \left[\frac{V_{2}}{h^{3}}\left(\frac{4 \pi m_{2} E_{2}}{3 N_{2}}\right)^{3 / 2}\right] \tag{2.6.7}
\end{equation*}
$$

and

$$
\begin{equation*}
S_{\text {final }}=\frac{3}{2} k_{B} N_{1}+k_{B} N_{1} \ln \left[\frac{V}{h^{3}}\left(\frac{4 \pi m_{1} E_{1}}{3 N_{1}}\right)^{3 / 2}\right]+\frac{3}{2} k_{B} N_{2}+k_{B} N_{2} \ln \left[\frac{V}{h^{3}}\left(\frac{4 \pi m_{2} E_{2}}{3 N_{2}}\right)^{3 / 2}\right] \tag{2.6.8}
\end{equation*}
$$

We thus get

$$
\begin{equation*}
\Delta S=k_{B} N_{1} \ln \left(\frac{V}{V_{1}}\right)+k_{B} N_{2} \ln \left(\frac{V}{V_{2}}\right) \tag{2.6.9}
\end{equation*}
$$

Now since $V_{1}=N_{1} k_{B} T / p, V_{2}=N_{2} k_{B} T / p$, and so $V=V_{1}+V_{2}=\left(N_{1}+N_{2}\right) k_{B} T / p$, this can be written as,

$$
\begin{equation*}
\Delta S=k_{B} N_{1} \ln \left(\frac{N_{1}+N_{2}}{N_{1}}\right)+k_{B} N_{2} \ln \left(\frac{N_{1}+N_{2}}{N_{2}}\right)>0 \tag{2.6.10}
\end{equation*}
$$

We expect $\Delta S>0$ since the entropy always increases when a constraint is removed.
When the red gas mixes with the blue gas we get purple gas! The process is irreversible - there is no thermodynamic process that will separate the mixture back into separate volumes of red and blue gas. In irreversible processes, the entropy always increases.

Now consider what happens if the two gases on either side of the partition are both of the same type, say both are red gas. Before the partition is removed $S_{\text {init }}$ is the same as in Eq. (2.6.7) except with $m_{1}=m_{2}=m$.

With the partition removed, the system is a single gas of $N=N_{1}+N_{2}$ particles, with total energy $E=E_{1}+E_{2}$, confined to a volume $V$. The final entropy with the partition removed is then,

$$
\begin{align*}
S_{\text {final }}=S(E, V, N) & =\frac{3}{2} k_{B} N+k_{B} N \ln \left[\frac{V}{h^{3}}\left(\frac{4 \pi m E}{3 N}\right)^{3 / 2}\right]  \tag{2.6.11}\\
& =\frac{3}{2} k_{B}\left(N_{1}+N_{2}\right)+k_{B}\left(N_{1}+N_{2}\right) \ln \left[\frac{V}{h^{3}}\left(\frac{4 \pi m}{3} \frac{3}{2} k_{B} T\right)^{3 / 2}\right] \tag{2.6.12}
\end{align*}
$$

where we used $E / N=\frac{3}{2} k_{B} T$. Similarly using $E_{1} / N_{1}=E_{1} / N_{2}=\frac{3}{2} k_{B} T$ (since both sides were initially at the same temperature $T$ ), we can write the above as,

$$
\begin{align*}
S_{\text {final }} & =\frac{3}{2} k_{B} N_{1}+k_{B} N_{1} \ln \left[\frac{V}{h^{3}}\left(\frac{4 \pi m}{3} \frac{E_{1}}{N_{1}}\right)^{3 / 2}\right]+\frac{3}{2} k_{B} N_{2}+k_{B} N_{2} \ln \left[\frac{V}{h^{3}}\left(\frac{4 \pi m}{3} \frac{E_{2}}{N_{2}}\right)^{3 / 2}\right]  \tag{2.6.13}\\
& =S\left(E_{1}, V, N_{1}\right)+S\left(E_{2}, V, N_{2}\right) \tag{2.6.14}
\end{align*}
$$

We thus conclude that

$$
\begin{equation*}
S(E, V, N)=S\left(E_{1}, V, N_{1}\right)+S\left(E_{2}, V, N_{2}\right) \tag{2.6.15}
\end{equation*}
$$

The above result is equivalent to our earlier observation that the entropy, as we've computed in the microcanonical ensemble so far, is not extensive. Rather than obeying the extensivity condition (i) $\lambda S(E, V, N)=S(\lambda E, \lambda V, \lambda N)$, we found that the entropy computed from the microcanonical ensemble obeyed (ii) $\lambda S(E, V, N)=S(\lambda E, V, \lambda N)$. If we took our initial gases so that $E_{1}=E_{2}=E / 2$ and $N_{1}=N_{2}=N / 2$, then Eq. (2.6.15) is equivalent to (ii) with $\lambda=1 / 2$.

Comparing the above Eq. (2.6.13) with the previous Eq. (2.6.8), we see that $S_{\text {final }}$ has exactly the same form when both gases are the same as when they are different, except with $m_{1}=m_{2}=m$.

Hence we find the same entropy of mixing $\Delta S>0$ regardless of whether the two gases are the same or whether they are different! But this should not be - when the gases are the same, removing the partition is a reversible process. We can always reinsert the partition and return to a situation indistinguishable from the initial state. In such a reversible process we should have $\Delta S=0$ ! This contradiction is known as the Gibbs paradox for the entropy of mixing.

The source of the problem lies in whether or not one should regard the particles of the gas as distinguishable. If we can distinguish each and every particle of the gas from one another, then when we mix the two gases of the same type, we do not really have a reversible process. After the partition is reinserted, we have not returned to the initial state because we now have different particles on each side as compared to what was the case initially.

Think of each particle as being a different unique color and the point should be clear. If each particle is a different color (i.e. the particles are distinguishable) it is no longer clear that the entropy should be extensive (or equivalently, additive over subsystems). If we double the volume, energy, and number of particles, we have not just made a second copy of the original system - this is because all the new particles must now come in new colors!

It was Gibbs who realized that to resolve this paradox of the mixing entropy, as well as to make the entropy computed in the microcanonical ensemble extensive, it was necessary to regard the particles of a gas as being indistinguishable from one another. This assumption was later confirmed by our quantum mechanical understanding of atoms and molecules.

