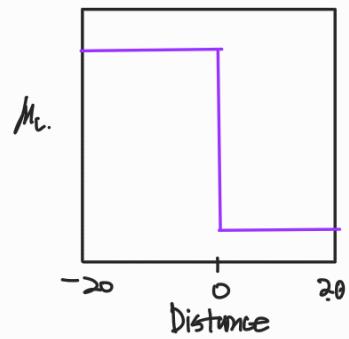
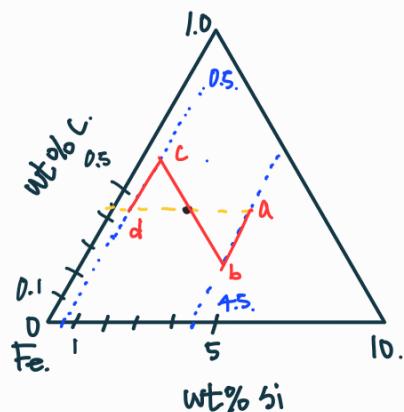
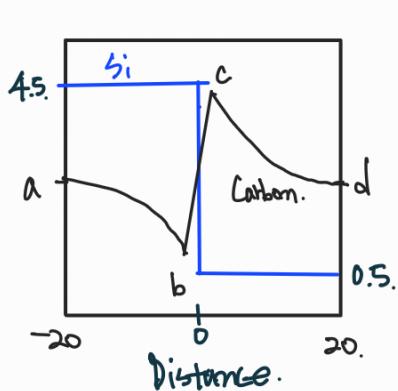


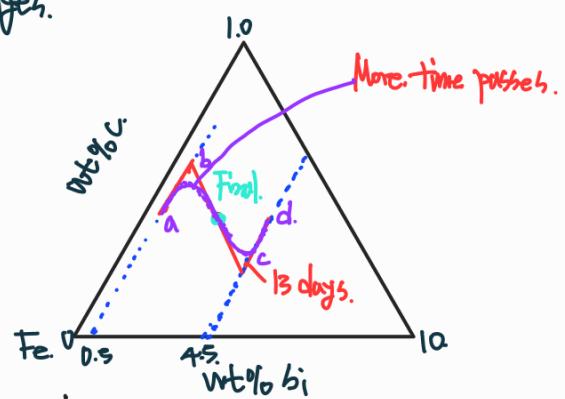
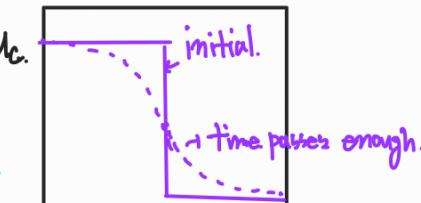
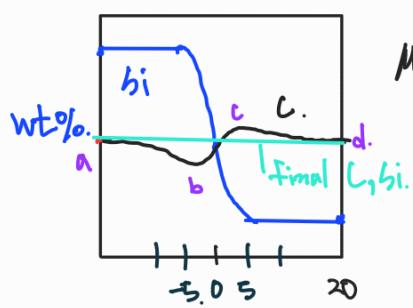
1. For the 13 days annealing time Si doesn't diffuse much so that abrupt Si concentration change would be observed at the interface.



(Assuming initial carbon concentration 0.45 wt%, silicon concentration for the site a. 4.5 wt% and site b. 0.5%.)

initially, Si-rich region (-20 to 0 mm) carbon has higher chemical potential than Si-poor region. The Mr gradient drives diffusion of carbon from a to d.

If time passes enough for Si to diffuse, Mr gradient changes.



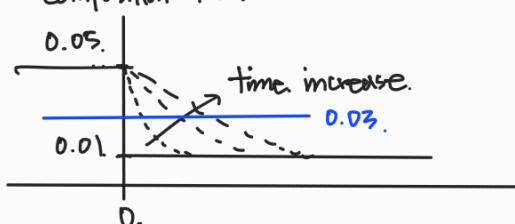
Initially sharp Mr becomes smooth and wt% C gradient reduces.

The diffusion profile no more keeps constant 5mm from interface both sides.

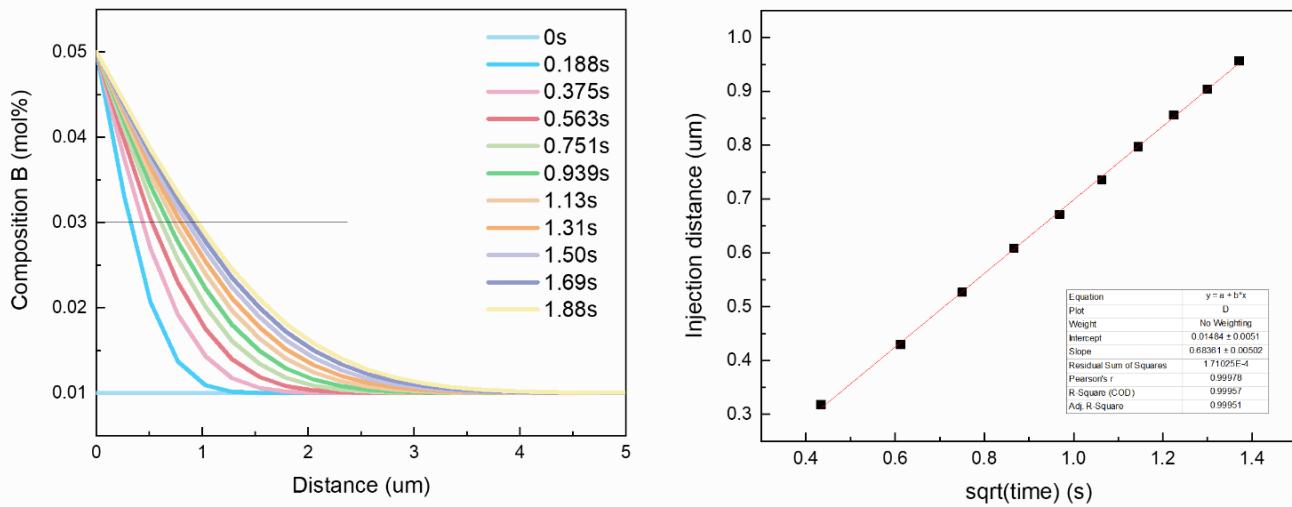
When equilibrium is reached, the composition is uniform at whole region of the sample.

2. $C_{B0} = 0.01$, $L_{B0} = 0.05$, $D_B = 5.25 \times 10^{-13} \text{ m}^2/\text{s.} = 5.25 \times 10^{-9} \text{ cm}^2/\text{s.}$ at 1300K.

Composition of B.



(a).



It is well known that injection distance l is proportional to \sqrt{Dt} .

$$\left(\begin{array}{l} \text{Diagram: A graph of concentration } C_s - C_0 \text{ vs. distance. The initial concentration } C_0 \text{ is at the origin. At distance } l, \text{ the concentration is } C_s - C_0. \text{ The area under the curve from } 0 \text{ to } l \text{ is shaded.} \\ \frac{C_s - C_0}{C_s - C_0} = 1 - \operatorname{erf}\left(\frac{l}{\sqrt{4Dt}}\right), \quad l(Dt) = l_0 + (C_s - C_0) \operatorname{erf}\left(\frac{l}{\sqrt{4Dt}}\right). \\ \operatorname{erf}\left(\frac{l}{\sqrt{4Dt}}\right) \sim \frac{1}{2}, \quad l\left(\frac{1}{2}, t\right) = l_0 + (C_s - C_0) \cdot \frac{1}{2} = \frac{C_0 + C_s}{2}, \quad \therefore l \sim \sqrt{Dt}. \end{array} \right)$$

At constant temperature, the diffusion coefficient is constant.

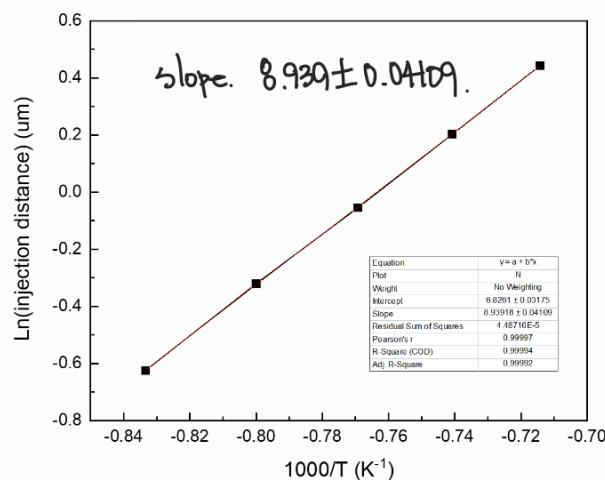
The injection distance l has linear relationship with \sqrt{Dt} .

(b). Diffusion coefficient. $D = D_0 \exp\left(-\frac{Q}{RT}\right) \propto l^2/t$

injection distance l is proportional to \sqrt{D} for some diffusion duration.

$$\sqrt{D} = \sqrt{D_0} \exp\left(-\frac{Q}{2RT}\right), \text{ in other expression. } \ln l = \frac{1}{2} \ln D_0 - \frac{Q}{2RT}$$

$\ln l$ vs. $(-\frac{1}{T})$ graph. should be linear.



(c). When the reaction is plotted as $\ln l$ vs. $(-\frac{1}{T})$ graph, slope is $-\frac{Q}{2R}$.

If the graph shows linearity, activation energy $Q = -2R \times (\text{slope})$.

$$Q = 1000 \times 8.94 \times 2 \times 8.3144 = 148 kJ$$