

Q6. (a) At 1200 °C, the Oxidation of a 1500 Å gate oxide is grown in two steps by dry oxidation. In step one- 1000 Å is grown first and in second step- wafer is re-oxidized to a total thickness of 1500 Å. Find the time required in the first and second process.

(b) In another experiment 1500 Å gate oxide is grown in one single step by dry oxidation at 1200 °C. Find the time required to grow the oxide. [13]

You may use $A = 0.04 \mu\text{m}$ and $B = 0.045 \mu\text{m}^2/\text{hr}$ at 1200 °C.

$$t = 0.027 \text{ h}$$

You may need: Atomic weight of boron = 10.81 g/mol, $N_A = 6.02 \times 10^{23}$, $K_B = 1.38 \times 10^{-23} \text{ J/K}$, Atomic weight of Si = 28.9 g/mol, Atomic weight of Ge = 72.64 g/mol. Atomic radius of Si = 0.118 nm, density of Si = 2.33 g/cm³, atomic weight of SiO₂ = 60.08 g/mol. density of SiO₂ = 2.21 g/cm³ gas constant R = 8.31 J/mol.K

$$\int_0^u \text{erfc } z \, dz = u \text{erfc } u + \frac{1}{\sqrt{\pi}} (1 - e^{-u^2})$$

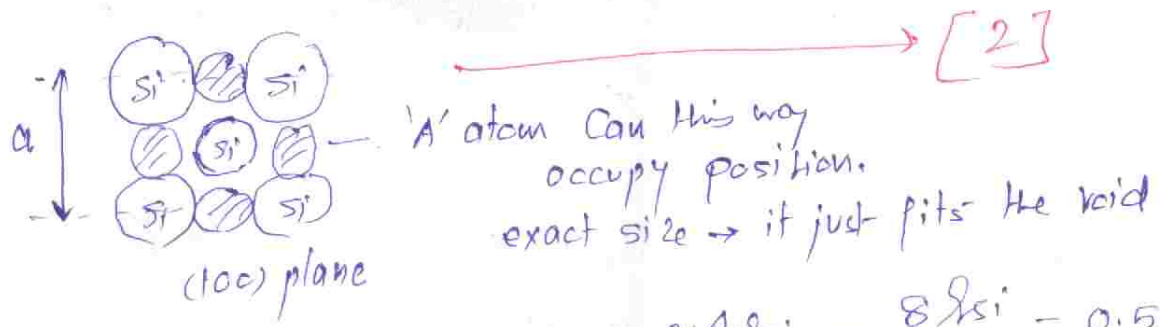
Table 5.1 Tabulation of Error Function Values

z	$\text{erf}(z)$	z	$\text{erf}(z)$	z	$\text{erf}(z)$
0	0	0.55	0.5633	1.3	0.9340
0.025	0.0282	0.60	0.6039	1.4	0.9523
0.05	0.0564	0.65	0.6420	1.5	0.9661
0.10	0.1125	0.70	0.6778	1.6	0.9763
0.15	0.1680	0.75	0.7112	1.7	0.9838
0.20	0.2227	0.80	0.7421	1.8	0.9891
0.25	0.2763	0.85	0.7707	1.9	0.9928
0.30	0.3286	0.90	0.7970	2.0	0.9953
0.35	0.3794	0.95	0.8209	2.2	0.9981
0.40	0.4284	1.0	0.8427	2.4	0.9993
0.45	0.4755	1.1	0.8802	2.6	0.9998
0.50	0.5205	1.2	0.9103	2.8	0.9999

Q1: 20 Marks
Total

Model Solutions
Mid Sem 2016
MEL 6611

Q1 (i) A Silicon unit cell will have 4 octahedral voids at edge centres and at center of unit cell.



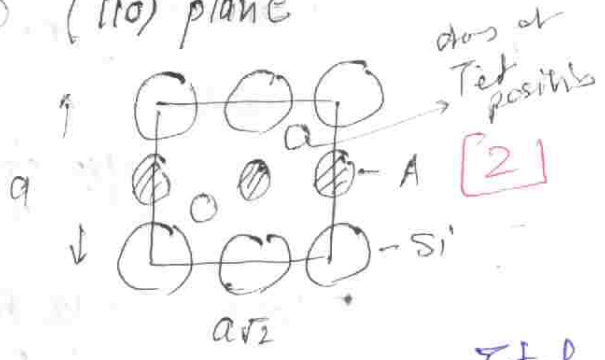
$$a = 2r_{Si} + 2r_A = \frac{2 \times 4 r_{Si}}{\sqrt{3}} = \frac{8 r_{Si}}{\sqrt{3}} = 0.5450 \text{ nm}$$

given $r_{Si} = 0.118 \text{ nm}$

$$\Rightarrow r_A = 0.1545 \text{ nm}$$

This problem is of course hypothetical! Dop is substituted nearly

(ii) (110) plane



$$PD = \frac{(4+2)}{\sqrt{2} a^2} = 14.26 / \text{nm}^2$$

(iii) packing fraction

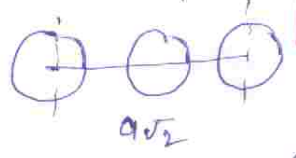
$$\text{Total APF} = \frac{4 \times \frac{4}{3} \pi r_A^3 + 8 \times \frac{4}{3} \pi r_{Si}^3}{a^3}$$

Thus APF =

$$= \frac{4 \times \frac{4}{3} \pi r_A^3}{a^3} + \frac{8 \times \frac{4}{3} \pi r_{Si}^3}{a^3} = 0.38$$

by 'A' atoms

(iv) linear density of [110]



$$LD = \frac{2}{a\sqrt{2}} = 2.595 / \text{nm} \quad [3]$$

(v) Number density: $4/a^3$

$$= \frac{24.70}{\text{nm}^3} \quad [4] \rightarrow [2+2]$$

$\frac{4}{a^3} = \text{Total} = \dots$

Q2
10 Marks

(a) wt% to atom% Conversion

$$C_1 = \frac{m_1}{m_1 + m_2} \times 100 \quad (\text{wt}\%)$$

$$C_1' = \frac{n_1}{n_1 + n_2} \times 100 \quad (\text{atom}\%)$$

$$C_1' = \frac{\frac{m_1}{A_1}}{\frac{m_1}{A_1} + \frac{m_2}{A_2}} \times 100 = \frac{\left(\frac{m_1}{m_1 + m_2}\right) / A_1}{\frac{m_1}{(m_1 + m_2) A_1} + \frac{m_2}{(m_1 + m_2) A_2}} \times 100$$

$$C_1' = \left(\frac{C_1/A_1}{C_1/A_1 + C_2/A_2} \right) \times 100$$

$$C_1' = \frac{C_1 A_2}{C_1 A_2 + C_2 A_1} \times 100 \quad [2]$$

Given $A_{Si} = 28.9 \text{ g/mol}$ $A_{Ge} = 72.64 \text{ g/mol}$

Substituti- finally gives

$$C_{Si}' = \frac{90x}{90x + 10x} \times 100 = 95.76\%$$

$$= 95.76\% \quad [3]$$

$$C_{Ge}' = 100 - C_{Si}' = 4.23\%$$

$$4.23\%$$

(b) Surface energy =

$$\text{bond energy per atom} \times \frac{\text{no of broken bonds}}{\text{Total bonds}} \times PD$$

Ge (110) plane

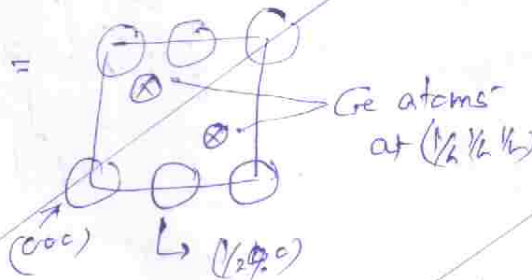
planar density =

Total atoms = 4

$$PD = \frac{4}{\sqrt{2}a^2}$$

$$a = \frac{8 \rho_{Ge}}{\sqrt{3}} = 0.5634 \text{ nm}$$

$$PD = 8.91 / \text{nm}^2$$



PD: planar density.

Total bonds = 12 For FCC
but only 4 for Ge
or even Si

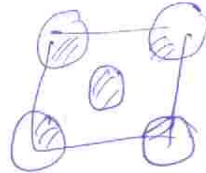
b (4) $SE = \text{bond energy per atom} \times \frac{\text{Total broken bonds}}{\text{Total bonds}} \times PD$ [1]

(100) plane of Ge

Total atoms about any Ge = 4 = Total bonds

For (100) you must appreciate that 2 bonds are broken. [2]

$PD_{100} \text{ Ge} = \frac{2}{a^2}$



$a = \frac{8}{\sqrt{3}} r_{\text{Ge}}$
 $= 0.5634 \text{ nm}$

$PD = 6.30 / \text{nm}^2$ [2]

Bond energy = $\frac{300 \text{ kJ}}{\text{mol}} = \frac{300 \times 10^3 \text{ J}}{6.02 \times 10^{23} \times 1.6 \times 10^{-19}} \text{ eV/particle}$
 $= 3.11 \text{ eV/atom}$

$SE = 3.11 \text{ eV} \times \frac{2}{4} \times 6.30 / \text{nm}^2$

$SE = 9.79 \text{ eV/nm}^2$ [2]

$n = 3.6 \times 10^{23} / \text{m}^3 = 3.6 \times 10^{17} / \text{cm}^3$

Q2C8

atom density

$n = N e^{-E/KT}$

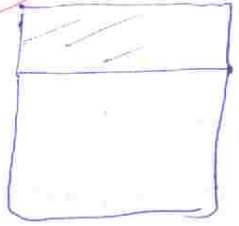
$N = \frac{8 N_A}{A} = \frac{8 \times 6.02 \times 10^{23}}{107.9} = 5.30 \times 10^{22} / \text{cm}^3$ [3]

$E = KT \ln\left(\frac{N}{n}\right) = 1.38 \times 10^{-23} \times 1273 \cdot \ln\left(\frac{5.30 \times 10^{22}}{3.6 \times 10^{17}}\right)$
 $= 1.30 \text{ eV/particle}$ [2]

Fraction at 1300K = $\frac{n}{N} = e^{-E/KT} = e^{-11.5942} = 9.21 \times 10^{-6}$
 $\Rightarrow 9.21 \times 10^{-6} \%$ [3]

~~Q1~~ Q3
15

$C_s = 10^{21} / \text{cm}^3$
 $100 \text{ \AA} = 10^{-6} \text{ cm}$



$$\Phi_T = \Phi_{SiO_2} + \Phi_{Si}$$

$$\Phi_T = 2C_s \sqrt{\frac{Dt}{\pi}} \quad [1]$$

$D = D_0 e^{-E_a/RT} = 3.26 \times 10^{-18} \text{ cm}^2/\text{s}$
Diffusion length

$$\sqrt{Dt} = 2.15 \times 10^{-7} \text{ cm} \quad [2]$$

Then $\Phi_T = 2.43 \times 10^{14} / \text{cm}^2 \quad [2]$

$$\Phi_T = \int_0^{10^{-6}} C(x) dx + \Phi_{Si}$$

$$\Phi_{Si} = \Phi_T - \int_0^{10^{-6}} C(x) dx \quad [2]$$

$$\Phi_{SiO_2} = \int_0^{10^{-6}} C_s e^{-\frac{x}{2\sqrt{Dt}}} dx$$

$$\frac{x}{2\sqrt{Dt}} = z$$

$$dx = 2\sqrt{Dt} dz$$

upper limit changes to $\frac{10^{-6}}{2\sqrt{Dt}} = 2.3255$

$$\Phi_{SiO_2} = 2C_s \sqrt{Dt} \int_0^{2.3255} e^{-z^2} dz \quad [2]$$

$$\Phi_{SiO_2} = 2C_s \sqrt{Dt} \int_0^{2.3255} e^{-z^2} dz$$

$$= 2C_s \sqrt{Dt} \left[2.3255 (e^{-2.3255}) + \frac{1}{\sqrt{\pi}} (1 - e^{-2.3255^2}) \right]$$

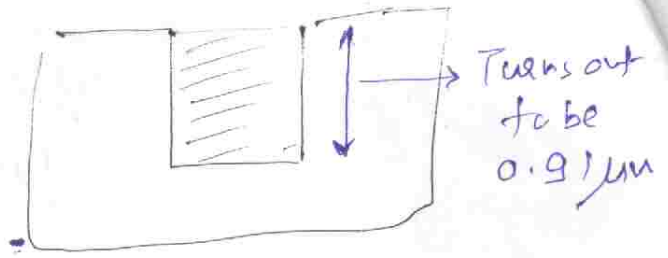
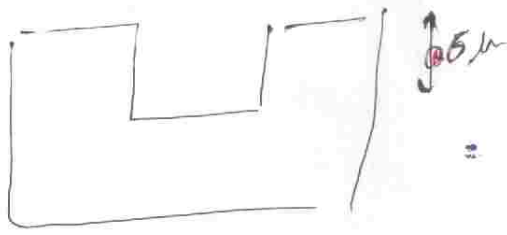
$$\Phi_{SiO_2} = 2C_s \sqrt{Dt} \left[2.3255 \times 10^{-6} + \frac{1}{\sqrt{\pi}} 0.5618 \right]$$

$$= 2.4257 \times 10^{14} / \text{cm}^2 \quad [2]$$

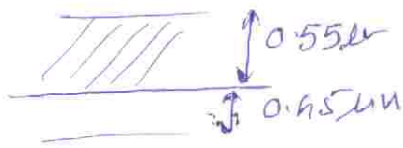
$$\Phi_{Si} = \Phi_T - \Phi_{SiO_2}$$

$$\approx 4.3 \times 10^{11} / \text{cm}^2 \quad [4]$$

Q5 12



Groove is of $0.5 \mu\text{m}$.



During oxidation vol. exp.

takes place in such a way that for a bare Si surface $0.55 \mu\text{m}$ oxide grows above, and $0.45 \mu\text{m}$ grows below. This is if we oxidize Total $1 \mu\text{m}$

We need to fill groove by oxidizing Si

$\therefore 0.55 \mu\text{m}$ is grown ^{above surface} if total $1 \mu\text{m}$

$\therefore 0.6 \mu\text{m}$ to grow we need to grow $1.09 \mu\text{m}$

$$\text{Total } x = \frac{0.6}{0.55} = \boxed{6} \mu\text{m}$$

Thus $x^2 + Ax = B(t + \epsilon)$ [2]

given $A = 0.421 \mu\text{m}$ $B = 0.316 \frac{\mu\text{m}^2}{\text{h}}$

$\epsilon = 0$
for wet oxidation \uparrow

This gives

$$t = \frac{x^2 + Ax}{B}$$

$$= \frac{(0.6)^2 + 0.421 \times 0.6}{0.316}$$

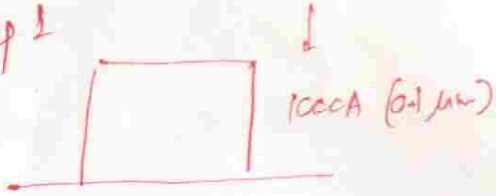
$$= 5.21 \text{ h}$$

$t = \quad \text{h}$

4

26 (13)

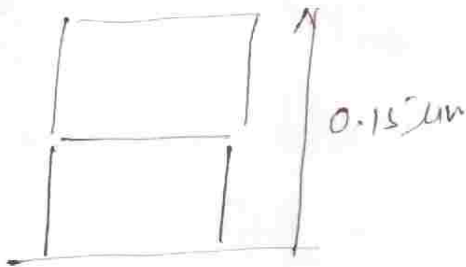
step 1



$$x^2 + Ax = B(t + E) \quad [1]$$

$$t_1 = \frac{(0.1)^2 + (0.045)(0.1) - 0.027}{0.045}$$

$$t_1 = \frac{0.284}{0.045} = 0.284 \text{ h} \quad [3]$$



$$E_{\text{new}} = 0.284 + 0.027 = 0.311 \text{ h} \quad [2]$$

$$x^2 + Ax = B(t + E)$$

$$t_2 = \frac{(0.15)^2 + (0.15) \times 0.045 - 0.027}{0.045}$$

$$= 0.322 \text{ h} \quad [3]$$

Now, to grow

0.15 μm oxide

in one single

day oxidation $E = 0.027 \text{ h}$

process: $x = 0.15 \mu\text{m}$

$$t = \frac{(0.15)^2 + (0.045)(0.15) - 0.027}{0.045}$$

$$= 0.322 \text{ h} \quad [4]$$

$$t_1 = 0.284 \text{ h}$$

$$t_2 = 0.322 \text{ h}$$

$$t = 0.322 \text{ h} \quad [4]$$

Thus $t = t_1 + t_2$

as expected!