NANYANG TECHNOLOGICAL UNIVERSITY SCHOOL OF ELECTRICAL & ELECTRONIC ENGINEERING ACADEMIC YEAR 2022-2023 SEMESTER 1

EE3013 SEMINCONDUCTOR DEVICES AND PROCESSING

Ion-Implantation

Q1. Boron is implanted with an energy of 60 keV through a 0.25µm layer of silicon dioxide. The implanted dose is $1x10^{14}$ cm⁻². Find the boron concentration at the silicon-silicon dioxide interface if the straggle range ΔR_p is 0.09µm and the projected range R_p is 0.19µm.

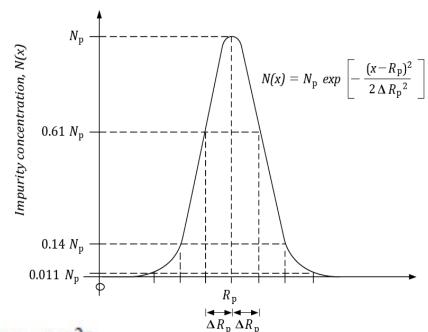
Given $R_p = 0.19 \ \mu m$ and $\Delta R_p = 0.09 \ \mu m$. For the Gaussian implant,

$$N(x) = N_p \exp[-(x-R_p)^2/2\Delta R_p^2]$$

$$N_p = Q/[\Delta Rp \sqrt{(2\pi)}]$$
= $(1x10^{14})/[(0.09x10^{-4})\sqrt{(2\pi)}]$
= $4.43x10^{18} \text{ cm}^{-3}$

$$N(0.25 \mu m) = 4.43 \times 10^{18} exp [-(0.25-0.19)^2/2(0.09)^2]$$

= 3.5 x 10¹⁸/em³



Distance into material, x

- **Q2**. A 200 mm-silicon wafer is implanted with P⁺ ions at 100 KeV for 5 seconds. If the average current measured is 10 μA, find
- (a) the dose
- (b) the maximum dopant concentration, and
- (c) the depth at which the maximum dopant concentration occurs
- (d) If a p-n junction is located at 0.3 μ m, what is the background dopant concentration of the wafer? Sketch the dopant profile, and verify if there is one more p-n junction!

Use the charts (*Lecture Notes*) to determine the approximate values of R_p and ΔR_{p} .

Wafer diameter =200 mm=20 cm,

$$A = \pi r^2 = 3.1416 \times 10^2 = 314.16 \text{ cm}^2$$

P⁺ ion implanted at $V = 100 \text{KV}$
E=100 KeV, t=5 Sec, I=10 μ A

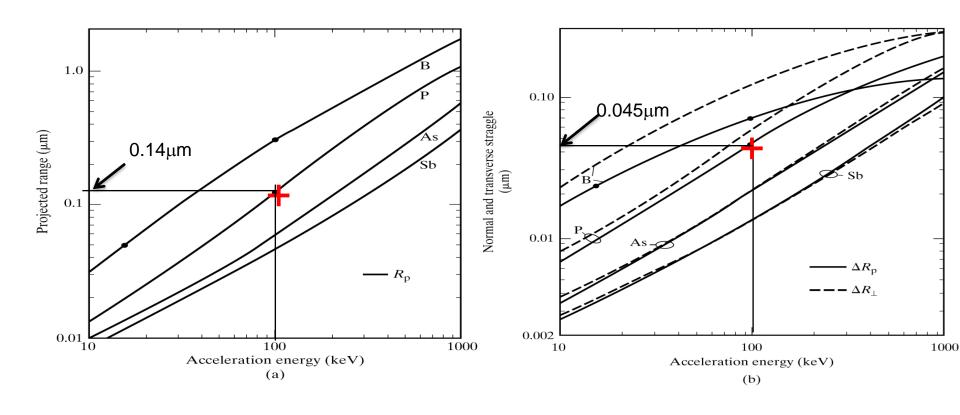
a) **Dose**

$$Q = \frac{It}{qA} = 0.995 \times 10^{12} \text{ cm}^{-2} \approx 1 \times 10^{12} \text{ cm}^{-2}$$

$$Q = \frac{It}{qA} = \frac{(C/s)(s)}{C. \text{ cm}^2} = \text{cm}^{-2}$$

From the charts, for P+ at 100 KeV, we have

$$R_{p} = 0.14 \mu m; \quad \Delta R_{p} = 0.045 \mu m$$



b) & c), Maximum dopant concentration occurs at a depth $(x = R_p)$

Maximum dopant concentration is obtained from

$$N(x) = \frac{Q}{\sqrt{2\pi}\Delta R_{p}} \cdot \exp(-\frac{(x - R_{p})^{2}}{2\Delta R_{p}^{2}})$$

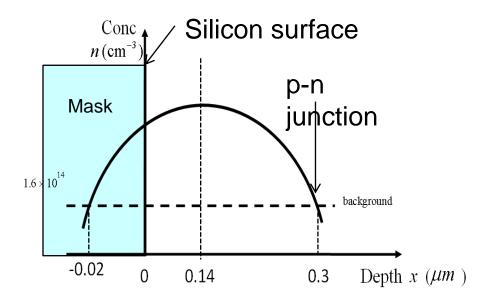
$$N_{\text{max}} = \frac{Q}{\sqrt{2\pi}\Delta R_{p}} \approx 8.87 \times 10^{16} \text{ cm}^{-3}$$

d) If a p-n junction is located at 0.3µm, the background p-type concentration must equal to the implanted n-type profile at

$$x = x_j = 0.3 \times 10^{-4} \text{ cm}$$

$$N(x_j) = 8.87 \times 10^{16} \cdot \exp(-\frac{(0.3 - 0.14)^2}{2 \times 0.045^2}) = 1.6 \times 10^{14} \,\mathrm{cm}^{-3}$$

To sketch the dopant profile:



Since the Gaussian curve is symmetrical about R_p , two junction depths will be obtained such that $(x_{j2}-R_p)=(R_p-x_{j1})$

Where $x_{j2} > x_{j1}$ In this case,

$$x_{j2} - R_p = 0.3 - 0.14 = 0.16 \ \mu m \text{ and } x_{j1} = -0.02 \ \mu m \text{ (outside Si)}$$

One p-n junction only!

- **Q3**. We wish to implant double-ionized boron into 0.25 μ m oxide-covered n-type <100> silicon wafer with a background concentration of 1x10¹⁵ atoms/cm³. The dose is 2x10¹³cm⁻², and the acceleration voltage used is 150KV.
 - (a) Locate the p-n junctions and sketch the doping profile, if any.
- (b) Determine the amount of boron atoms that ended up inside the silicon wafer.

You may assume that, for boron: R_p in $Si = R_p$ in SiO_2 , and ΔR_p in $Si = \Delta R_p$ in SiO_2 . Values of R_p and ΔR_p can be taken from the graphs provided in the *Lecture Notes*.

V=150 KV; doubly ionized B (B²⁺); E = 300 KeV (2e×150KV), and Q = 2×10^{13} cm⁻².

From R_p and ΔR_p charts for B, $R_p = 0.7 \mu m$ and $\Delta R_p = 0.1 \mu m$.

$$N_p = \frac{Q}{\sqrt{2\pi} \cdot \Delta R_p} = 7.98 \times 10^{17} \text{ cm}^{-3}$$

Backgroud n-type conc. = 1×10^{15} cm⁻³.

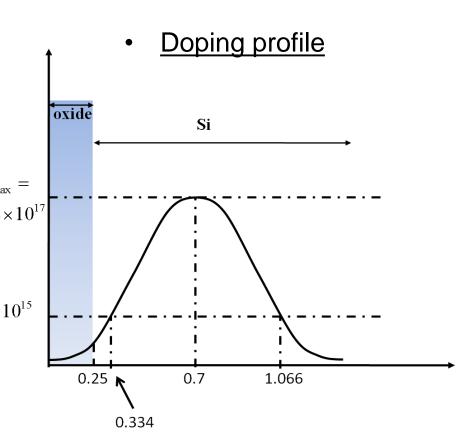
(a) To solve for junction depth:

$$N_{B} = \frac{Q}{\sqrt{2\pi}\Delta R_{p}} \cdot \exp(-\frac{(x_{j} - R_{p})^{2}}{2\Delta R_{p}^{2}}) \qquad n_{\text{max}} = \frac{1 \times 10^{15}}{7.98 \times 10^{17}}$$

$$1 \times 10^{15} = 7.98 \times 10^{17} \exp(-\frac{(x_{j} - 0.7)^{2}}{2 \times 0.1^{2}}) \qquad 1 \times 10^{15}$$

$$x_{j2} = 0.7 + 0.366 = 1.066 \ \mu m$$

$$x_{j1} = 0.7 - 0.366 = 0.334 \ \mu m$$



Since oxide is 0.25 µm thick, both junctions are formed inside the semiconductor.

Boron inside Si

$$Q(Si) = \frac{Q}{\sqrt{2\pi}\Delta R_{p}} \cdot \int_{d}^{\infty} \exp(-\frac{(x - R_{p})^{2}}{2\Delta R_{p}^{2}}) dx$$

See lecture notes or Highlight slide
$$\frac{Q(Si)}{Q} = \frac{1}{\sqrt{2}} \operatorname{erfc} \left(\frac{d - R_{p}}{\sqrt{2}\Delta R_{p}} \right) = \frac{1}{2} \operatorname{erfc} \left(\frac{0.25 - 0.7}{\sqrt{2}(0.1)} \right) = \frac{1}{2} \operatorname{erfc} \left(-3.182 \right)$$

$$\operatorname{erfc}(-z) = 1 - \operatorname{erf}(-z) = 1 - [-\operatorname{erf}(z)] = 1 + \operatorname{erf}(z)$$

$$erf(-z) = -erf(z)$$

$$Erfc(-3.182) = 1 + erf(3.182) = 1 + 0.999993$$

$$\therefore \frac{Q(Si)}{O} = \frac{1}{2} [1 + 0.9999993] = 0.99999965$$

Thus
$$Q(Si) \approx Q = 2 \times 10^{13} \text{ cm}^{-2}$$

Supplementary

The complementary error function is defined as

$$erfc(x) = \frac{2}{\sqrt{\pi}} \int_{z}^{\alpha} \exp(-t^2) dt.$$

Hence,

$$Q(Si) = \frac{Q}{\sqrt{2\pi}\Delta R_{p}} \cdot \int_{d}^{\infty} \exp(-\frac{(x-R_{p})^{2}}{2\Delta R_{p}^{2}}) dx$$

$$\frac{Q(Si)}{Q} = \frac{1}{\sqrt{2\pi}\Delta R_{p}} \cdot \int_{d}^{\infty} \exp\left(-\frac{x-R_{p}}{\sqrt{2}\Delta R_{p}}\right)^{2} dx$$

$$= \frac{1}{\sqrt{2\pi}\Delta R_{p}} \cdot \int_{d}^{\infty} \exp\left(-\frac{x-R_{p}}{\sqrt{2}\Delta R_{p}}\right)^{2} d\left(\frac{x-R_{p}}{\sqrt{2}\Delta R_{p}}\right) (\sqrt{2\Delta}R_{p})$$

$$= \frac{\sqrt{2\Delta}R_{p}}{\sqrt{2\pi}\Delta R_{p}} \cdot \int_{v}^{\infty} \exp(-u^{2}) du \quad \text{[where } (u = \frac{x-R_{p}}{\sqrt{2}\Delta R_{p}}) \text{ and } v = \frac{d-R_{p}}{\sqrt{2}\Delta R_{p}}]$$

$$= \frac{1}{\sqrt{\pi}} \cdot \frac{2}{2} \int_{v}^{\infty} \exp(-u^{2}) du = \frac{1}{2} \cdot \frac{2}{\sqrt{\pi}} \int_{v}^{\infty} \exp(-u^{2}) du$$

$$= \frac{1}{2} \operatorname{erfc}(v) = \frac{1}{2} \operatorname{erfc}(\frac{d-R_{p}}{\sqrt{2\Delta}R_{p}})$$

- **Q4**. Boron was implanted into an n-type oxide-covered silicon wafer with a background concentration of $1x10^{15}cm^{-3}$. The oxide thickness was 0.6 μ m and the ion dose used was $2x10^{13}$ atoms/cm². After implantation the wafer was annealed at 1050 °C for 30 minutes to activate the implanted ions. Assume that the values of projected range and straggle are same for B in Si and the oxide. Given R_p =700 nm and ΔR_p =100 nm, Diffusion Coefficient of B =5x10⁻¹⁴ cm²/s at 1050 °C.
- (a) Calculate the peak concentration of the implanted profile and the junction depths from the wafer surface. Draw the schematic of the dopant profile.
- (b) Calculate the peak concentration and the projected straggle after annealing. Does the peak location change after annealing?
- (c) Find the required thickness of the oxide before and after annealing if the maximum concentration of B in Si is at most 10% of the background concentration.

(a) Peak concentration, junction depth and dopant profile before annealing:

$$N(x) = \frac{Q}{\sqrt{2\pi}\Delta R_{p}} \cdot \exp(-\frac{(x - R_{p})^{2}}{2\Delta R_{p}^{2}}),$$

$$R_p = 0.7 \,\mu\text{m}, \ \Delta R_p = 0.1 \,\mu\text{m}, \ Q = 2 \times 10^{13} \text{ cm}^{-2}$$

Peak concentration of implanted profile:

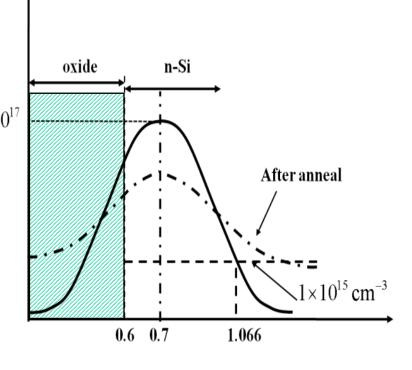
$$N_p = \frac{2 \times 10^{13}}{\sqrt{2\pi} \times 0.1 \times 10^{-4}} \,\text{cm}^{-3} = 7.98 \times 10^{17} \,\text{cm}^{-3}$$

$$7.89 \times 10^{17}$$

• Junction depth: $1 \times 10^{15} = 7.98 \times 10^{17} \times \exp(-\frac{(x-0.7)^2}{2 \times 0.1^2})$

$$\Rightarrow x_{j2} = 1.066 \mu \text{m},$$

 $x_{j1} = 0.334 \mu \text{m} \text{ (Invalid as it is in oxide)}$



(b) Peak concentration, straggle and peak location after annealing

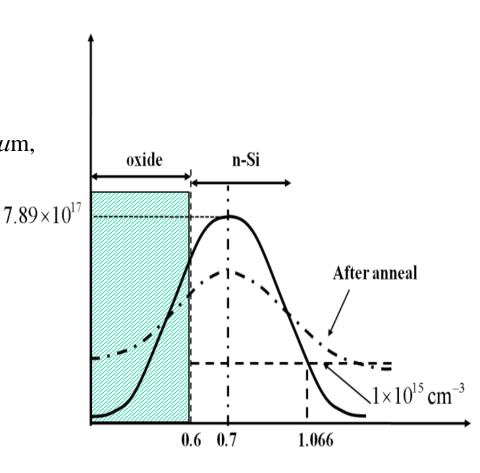
$$N'_{\text{max}} = \frac{Q}{\sqrt{2\pi} \left(\Delta R_{p}^{2} + 2Dt\right)^{1/2}}$$

$$= \frac{2 \times 10^{13}}{\sqrt{2\pi} \left(0.1^{2} + 2 \times 5 \times 10^{-14} \times 30 \times 60\right)^{1/2}}$$

$$= 4.77 \times 10^{17} \text{ cm}^{-3}$$

$$R'_{p} = 0.7 \,\mu\text{m}, \quad \Delta R'_{p} = \left(\Delta R_{p}^{2} + 2Dt\right)^{1/2} = 0.167 \,\mu\text{m},$$

 Peak position remains unchanged although junction goes deeper after annealing,

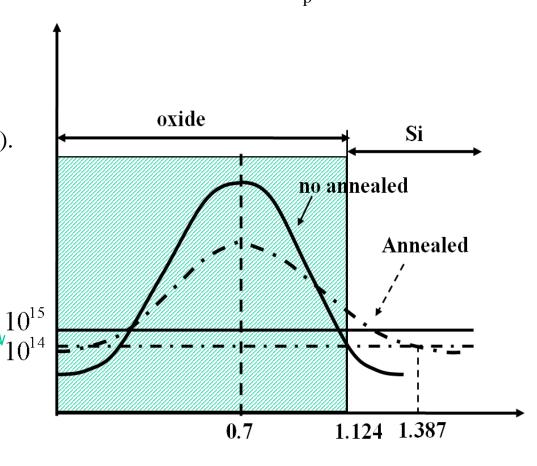


- (c) Required thickness of oxide with implanted B of 10% of background (1x10¹⁵cm⁻³) at the surface of Si.
- Before annealing, $N(x) = 7.98 \times 10^{17} \cdot \exp(-\frac{(x R_p)^2}{2\Delta R_p^2})$

When x=d,
$$N(d) = N_B / 10 = 10^{14} \text{ cm}^{-3}$$

 $10^{14} = 7.98 \times 10^{17} \cdot \exp(-\frac{(d-0.7)^2}{2 \times 0.1^2})$
 $\therefore d_2 = 1.124 \, \mu\text{m}, d_1 = 0.276 \, \mu\text{m} \text{ (invalid)}.$

If we have an oxide of 1.124 μm, the max conc. of the implanted dopant in the Si surface will be 10% of background concentration. The oxide should be > 1.124 μm for mask purpose.



· After annealing,

$$\Delta R'_{p} = (\Delta R_{p}^{2} + 2Dt)^{1/2} = 0.167 \,\mu\text{m},$$

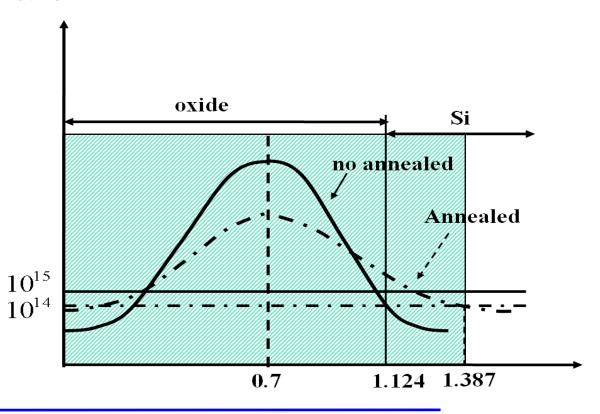
$$N'_{\text{max}} = \frac{Q}{\sqrt{2\pi} \cdot \Delta R'_{p}} = 4.77 \times 10^{17} \text{ cm}^{-3};$$

$$\therefore 10^{14} = 4.77 \times 10^{17} \cdot \exp(-\frac{(d-0.7)^2}{2 \times 0.167^2})$$

$$\Rightarrow d_2 = 1.387 \,\mu\text{m},$$

$$d_1 = 0.013 \,\mu\text{m (invalid)})$$

 Oxide thickness required in this case is at least 1.387 µm.

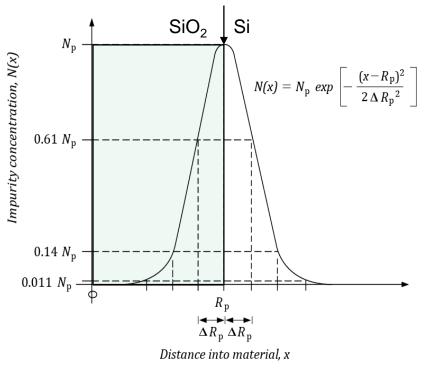


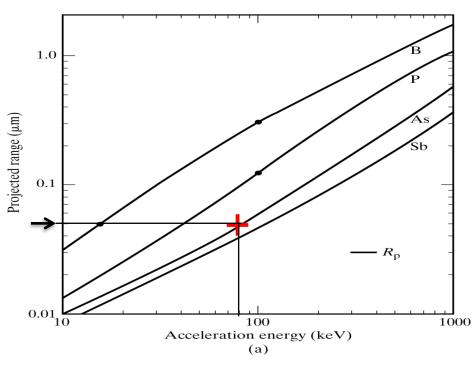
Q5. An arsenic dose of $1x10^{12}$ cm⁻³ is implanted through a 50 nm layer of silicon dioxide with the peak of the distribution at the Si-SiO₂ interface. The straggle range ΔR_p is $0.017\mu m$. A silicon nitride film on top of the silicon oxide is to be used as a barrier material in the regions where arsenic is not desired. How thick should the nitride layer be if the background concentration is $1x10^{15}$ cm⁻³? Assume silicon oxide is only 85% effective in stopping ions as compared to silicon nitride.

- First, calculate the total oxide thickness needed to ensure that the implanted impurity concentration is less than $10^{15}/10 = 10^{14}/\text{cm}^3$ at the Si-SiO₂ interface.
- We know that $R_p = 0.05 \, \mu m$ which is the thickness of the oxide. For arsenic, this requires E = 80 keV from the R_p vs Acceleration energy chart, and $\Delta R_p = 0.017 \, \mu m$.

$$N_P = Q/\Delta R_P \sqrt{2 \pi} = 2.35 \times 10^{17} / cm^3$$
.

Given that the peak of the distribution at the Si-SiO₂ interface





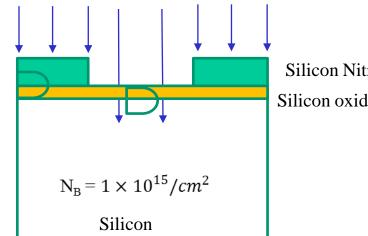
19

Recall from lecture that:

$$N(t_{ox}) = N_p \exp \left[-\frac{\left(t_{ox} - R_p\right)^2}{2\Delta R_p^2} \right] < \frac{N_B}{10}$$

and the required oxide thickness for the mask is:

$$t_{ox} \ge R_p + \Delta R_p \sqrt{2 \ln \left(\frac{10N_p}{N_B}\right)}$$



Silicon Nitride mask (t_{SiN}) Silicon oxide (50nm)

See lecture notes or Highlight slide

• Here,
$$N_P/N_B = 235$$
, $t_{ox} = R_P + 3.94 \Delta R_P$

$$\rightarrow$$
 $t_{ox} = R_P + 3.94 \Delta R_P = 0.05 + 3.94(0.017) = 0.117 \mu m of oxide$

- The additional oxide required is $3.94(0.017) = 0.067 \mu m$
- However, $t_{SiN} = 0.85 t_{ox}$ so only 0.057 µm of silicon nitride is required.