

# Example 1

Determine the time needed to grow 0.35  $\mu\text{m}$  of oxide on a bare silicon wafer (i.e. no initial oxide) if the oxidation temperature is 1000°C. The wafer has (100) orientation. Find the times for both wet and dry oxidations.

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Do the wet case first. The first step is to find the linear and parabolic coefficients at 1000°C = 1273 K.

$$\begin{aligned}\left(\frac{B}{A}\right)_{wet} &= \left(9.7 \times 10^7 \frac{\mu\text{m}}{\text{hr}}\right) \exp\left(-\frac{2.05\text{eV}}{kT}\right) \\ &= \left(9.7 \times 10^7 \frac{\mu\text{m}}{\text{hr}}\right) \exp\left(-\frac{2.05\text{eV}}{\left(8.617 \times 10^{-5} \frac{\text{eV}}{\text{K}}\right) (1273\text{K})}\right) = 0.742 \frac{\mu\text{m}}{\text{hr}}\end{aligned}$$

$$\begin{aligned}B_{wet} &= \left(386 \frac{\mu\text{m}^2}{\text{hr}}\right) \exp\left(-\frac{0.78\text{eV}}{kT}\right) \\ &= \left(386 \frac{\mu\text{m}^2}{\text{hr}}\right) \exp\left(-\frac{0.78\text{eV}}{\left(8.617 \times 10^{-5} \frac{\text{eV}}{\text{K}}\right) (1273\text{K})}\right) = 0.315 \frac{\mu\text{m}^2}{\text{hr}}\end{aligned}$$

Now calculate the required oxidation time. Note that since there was no initial oxide,  $\tau = 0$ .

$$t = \frac{t_{ox}}{B/A} + \frac{t_{ox}^2}{B}$$
$$= \frac{0.35 \mu m}{0.74 \frac{\mu m}{hr}} + \frac{(0.35 \mu m)^2}{0.32 \frac{\mu m^2}{hr}} = 0.86 hr$$

The wet oxidation should last 51 min, 36 sec.

The dry oxidation time is calculated in a similar manner.

$$\left(\frac{B}{A}\right)_{dry} = \left(3.71 \times 10^6 \frac{\mu m}{hr}\right) \exp\left(-\frac{2.00 eV}{\left(8.617 \times 10^{-5} \frac{eV}{K}\right) (1273K)}\right) = 0.045 \frac{\mu m}{hr}$$

$$B_{dry} = \left(772 \frac{\mu m^2}{hr}\right) \exp\left(-\frac{1.23 eV}{\left(8.617 \times 10^{-5} \frac{eV}{K}\right) (1273K)}\right) = 0.0104 \frac{\mu m^2}{hr}$$

$$t = \frac{0.35 \mu m}{0.045 \frac{\mu m}{hr}} + \frac{(0.35 \mu m)^2}{0.0104 \frac{\mu m^2}{hr}} = 19.57 hr \quad \rightarrow \text{significantly longer}$$

## Example 2

A (100)-oriented silicon wafer has 250 nm of oxide on it. It is run through a wet oxidation process at a temperature of 1100°C for 60 minutes. What is the oxide thickness at the end of the oxidation?

As in the previous example, we need the linear and parabolic coefficients first. For a wet oxidation at 1100°C (=1373 K):

$$\left(\frac{B}{A}\right)_{wet} = \left(9.7 \times 10^7 \frac{\mu m}{hr}\right) \exp\left(-\frac{2.05eV}{\left(8.617 \times 10^{-5} \frac{eV}{K}\right) (1373K)}\right) = 2.90 \frac{\mu m}{hr}$$

$$B_{wet} = \left(386 \frac{\mu m^2}{hr}\right) \exp\left(-\frac{0.78eV}{\left(8.617 \times 10^{-5} \frac{eV}{K}\right) (1373K)}\right) = 0.529 \frac{\mu m^2}{hr}$$

Because there is an initial oxide, we need to include that in the form of the equivalent time.

$$\tau = \frac{t_{oxi}}{B/A} + \frac{t_{oxi}^2}{B} = \frac{0.25 \mu m}{2.9 \frac{\mu m}{hr}} + \frac{(0.25 \mu m)^2}{0.53 \frac{\mu m^2}{hr}} = 0.20 hr$$

Then we can use the thickness equation to get the final thickness.

$$\begin{aligned}t_{ox} &= \frac{B}{2 \left(\frac{B}{A}\right)} \left[ \sqrt{1 + \frac{4 \left(\frac{B}{A}\right)^2}{B} (t + \tau)} - 1 \right] \\&= \frac{0.53 \frac{\mu m^2}{hr}}{2 \left(2.90 \frac{\mu m}{hr}\right)} \left[ \sqrt{1 + \frac{4 \left(2.90 \frac{\mu m}{hr}\right)^2}{0.53 \frac{\mu m^2}{hr}} (1hr + 0.2hr)} - 1 \right] \\&= 0.71 \mu m\end{aligned}$$

## Example 3

A (100) silicon wafer has 0.375  $\mu\text{m}$  of oxide. Determine the time needed to grow an additional 0.375  $\mu\text{m}$  of oxide using wet oxidation at temperature of 1050°C. (The wafer will have a total of 0.75  $\mu\text{m}$  of oxide when finished.)

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Find the linear and parabolic coefficients for wet oxidation at 1050°C (= 1323K).

$$\left(\frac{B}{A}\right)_{wet} = \left(9.7 \times 10^7 \frac{\mu\text{m}}{\text{hr}}\right) \exp\left(-\frac{2.05\text{eV}}{\left(8.617 \times 10^{-5} \frac{\text{eV}}{\text{K}}\right) (1323\text{K})}\right) = 1.50 \frac{\mu\text{m}}{\text{hr}}$$

$$B_{wet} = \left(386 \frac{\mu\text{m}^2}{\text{hr}}\right) \exp\left(-\frac{0.78\text{eV}}{\left(8.617 \times 10^{-5} \frac{\text{eV}}{\text{K}}\right) (1323\text{K})}\right) = 0.41 \frac{\mu\text{m}^2}{\text{hr}}$$

The time corresponding to the initial oxide is

$$\tau = \frac{t_{oxi}}{B/A} + \frac{t_{oxi}^2}{B} = \frac{0.375 \mu m}{1.50 \frac{\mu m}{hr}} + \frac{(0.375 \mu m)^2}{0.41 \frac{\mu m^2}{hr}} = 0.59 hr$$

Then, the time to grow the extra 0.375  $\mu m$  is

$$t = \frac{t_{ox}}{B/A} + \frac{t_{ox}^2}{B} - \tau$$
$$= \frac{0.75 \mu m}{1.50 \frac{\mu m}{hr}} + \frac{(0.75 \mu m)^2}{0.41 \frac{\mu m^2}{hr}} - 0.59 hr = 1.28 hr$$

## Example 4

An initially bare (100) silicon wafer is oxidized in two steps. The first step uses dry oxidation at 1150°C for 120 min. The second step uses wet oxidation at 1000°C for 30 min. Find the final thickness of the oxide.

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Find the linear and parabolic coefficients for dry oxidation at 1150°C (= 1423K).

$$\left(\frac{B}{A}\right)_{dry} = \left(3.71 \times 10^6 \frac{\mu m}{hr}\right) \exp\left(-\frac{2.00eV}{\left(8.617 \times 10^{-5} \frac{eV}{K}\right) (1423K)}\right) = 0.306 \frac{\mu m}{hr}$$

$$B_{dry} = \left(772 \frac{\mu m^2}{hr}\right) \exp\left(-\frac{1.23eV}{\left(8.617 \times 10^{-5} \frac{eV}{K}\right) (1423K)}\right) = 0.034 \frac{\mu m^2}{hr}$$

The oxide thickness at the end of the dry step is

$$t_{ox1} = \frac{0.034 \frac{\mu m^2}{hr}}{2 \left(0.306 \frac{\mu m}{hr}\right)} \left[ \sqrt{1 + \frac{4 \left(0.306 \frac{\mu m}{hr}\right)^2}{0.034 \frac{\mu m^2}{hr}} (2hr)} - 1 \right] = 0.21 \mu m$$

Now find the coefficients for wet oxidation at 1000°C (=1273K). (Same as in Example 1.)

$$\left(\frac{B}{A}\right)_{wet} = \left(9.7 \times 10^7 \frac{\mu m}{hr}\right) \exp\left(-\frac{2.05eV}{\left(8.617 \times 10^{-5} \frac{eV}{K}\right) (1273K)}\right) = 0.742 \frac{\mu m}{hr}$$

$$B_{wet} = \left(386 \frac{\mu m^2}{hr}\right) \exp\left(-\frac{0.78eV}{\left(8.617 \times 10^{-5} \frac{eV}{K}\right) (1273K)}\right) = 0.315 \frac{\mu m^2}{hr}$$

Now find the *wet oxidation time* corresponding to the thickness grown in the dry step.

$$\tau = \frac{t_{oxi}}{B/A} + \frac{t_{oxi}^2}{B} = \frac{0.21 \mu m}{0.742 \frac{\mu m}{hr}} + \frac{(0.21 \mu m)^2}{0.315 \frac{\mu m^2}{hr}} = 0.42 hr$$

To finish up, find the final thickness after the wet oxidation step.

$$t_{ox2} = \frac{0.315 \frac{\mu m^2}{hr}}{2 \left(0.742 \frac{\mu m}{hr}\right)} \left[ \sqrt{1 + \frac{4 \left(0.742 \frac{\mu m}{hr}\right)^2}{0.315 \frac{\mu m^2}{hr}} (0.5hr + 0.42hr)} - 1 \right] = 0.367 \mu m$$