

### Doping. Epitaxy

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Chapters 6, 14, 15



### **Previous lectures**

- Oxidation
- Lab device

#### **Next lecture**

• Bonding and CMP



### **Outline I**

- Selective doping
  - -Diffusion
  - -Implantation
- Epitaxy



### **Outline II**





### Types of doping

- Blank doping (wafer, ingot)
  - during crystal growth (doped raw Si)
  - during epitaxy (gas containing dopant)
- Selective doping
  - by ion implantation (ions of dopant)
  - by diffusion (gas or solid dopant)

### Selective doping by phosphorus



Gas phase doping Oxide as a mask POCl<sub>3</sub> as a source 1000°C Lateral spread = depth

Solid-source doping Oxide as a mask  $P_2O_5$  as a source 1000°C Lateral spread = depth Ion implantation Photoresist as a mask Accelerated P<sup>+</sup> as a source Room temperature? Lateral spread = 1/3 depth





Diffusion is the movement of foreign, or impurity atoms with respect to the atoms of the host crystal along concentration gradient



![](_page_7_Picture_0.jpeg)

#### Diffusion mechanisms in perfect lattice

![](_page_7_Figure_2.jpeg)

![](_page_8_Picture_0.jpeg)

#### Fick's first law

• Diffusion flux, atoms/(s·cm<sup>2</sup>)

$$j = -D\left(\frac{\partial N}{\partial x}\right)$$

- where *D* is the diffusion coefficient  $(cm^2/s)$ , *N* is concentration  $(cm^{-3})$ .
- Diffusion coefficient can be presented by

$$D = D_0 e^{-\frac{E_a}{kT}}$$

- *D<sub>o</sub>* is the frequency factor
- $E_a$  is the activation energy
- k is Boltzman's constant,  $k = 1.38*10^{-23}$  J/K
- *T* is temperature in Kelvin

#### At 1050 °C

Boron

**Phosphorous** 

For boron in Si at 950 °C $D = 4.3 \times 10^{-15} \text{ cm}^2/\text{s}$	$D_o$ (cm <sup>2</sup> /s)	0.76 3.46	3.85 3.66
	$D(cm^{2}/s)$ x (µm), 1h	5.2 × 10 <sup>-14</sup> 0.27	4.5 × 10 <sup>-14</sup> 0.25

![](_page_9_Picture_0.jpeg)

#### **Characteristic diffusion length**

For infinite dopant source distribution depends fully on characteristic length X at which impurity concentration is  $C \sim C_s/2$ , where  $C_s$  is surface concentration of dopant

#### $x \approx \sqrt{Dt}$

# Diffusion profiles: dopant concentration

#### Infinite source

#### Limited source or drive-in

![](_page_10_Figure_3.jpeg)

![](_page_11_Picture_0.jpeg)

# Time evolution of diffusion depth: $x \approx \sqrt{Dt}$

![](_page_11_Figure_2.jpeg)

![](_page_12_Picture_0.jpeg)

- Diffusion is done in oxidation furnaces
- Always O<sub>2</sub> is added to a doping gas, i.e diffusion is connected with Si oxidation
- In case of several diffusions, the 1-st one must be with highest temperature (the deepest one)
- Diffusion areas are invisible
- Sheet resistance decreases after doping
- 4-point probe, SIMS, SRP

![](_page_13_Picture_0.jpeg)

#### Mask thickness for selective diffusion

#### Usually, dopant transmission through mask is below 0.0001%

![](_page_13_Figure_3.jpeg)

#### The mask is thinner for B than for P at the same conditions

![](_page_14_Picture_0.jpeg)

#### Diffusion in solar cell

![](_page_14_Figure_2.jpeg)

#### Diffusion profiles in bipolar transistor

![](_page_15_Figure_1.jpeg)

### **OED** in LOCOS

OED -oxidation enhanced diffusion

![](_page_16_Figure_3.jpeg)

### **Multiple diffusions**

![](_page_17_Figure_1.jpeg)

- 1. Take n-type silicon wafer
- 2. Thermal oxidation
- 3. Lithography
- 4. Oxide mask etching +strip
- 5. Perform *p*-diffusion
- 6. Etch oxide away
- 7. Thermal oxidation
- 8. Lithography
- 9. Oxide etching + strip
- 10. n-diffusion;

*p*-diffusion becomes deeper

*n*-concentration (step 10) must be higher than *p*; otherwise dopant type does not change.

### Selective implantation and dopant profile

#### Room temperature

![](_page_18_Figure_3.jpeg)

Maximum concentration is below the surface Compare with diffusion!

![](_page_19_Picture_0.jpeg)

#### Projected range and straggle

![](_page_19_Figure_2.jpeg)

![](_page_20_Picture_0.jpeg)

### Projected range (R<sub>p</sub>) in Si

![](_page_20_Figure_2.jpeg)

#### Implantation damage

Can be removed by anneal at 1000°C during 30s

![](_page_21_Figure_3.jpeg)

![](_page_22_Picture_0.jpeg)

### Mask thickness for implantation

![](_page_22_Figure_2.jpeg)

Figure 4-61. Masking Thickness Required, Boron and Antimony Implants.

![](_page_23_Picture_0.jpeg)

#### Simulated implantation profiles

![](_page_23_Figure_2.jpeg)

![](_page_24_Picture_0.jpeg)

#### Lateral straggle $\Delta R_{\rm t}$

![](_page_24_Figure_2.jpeg)

Lateral standard deviation of boron, phosphorus, arsenic and antimony in silicon'

http://www-inst.eecs.berkeley.edu/~ee143/fa16/lectures/Lecture07-Ion%20Implantation.pdf

### Measured implantation profiles

#### Boron

Phosphorus

![](_page_25_Figure_3.jpeg)

**Fig. 6.3** Ion concentration. (*a*) Boron in silicon, 250-keV ions, annealed at 850°C for 30 min. Adapted from Moline [5]. (*b*) Phosphorus in silicon, 300-keV ions, annealed at 800°C for 30 min. Adapted from Dearnaley et al. [3].

![](_page_26_Picture_0.jpeg)

#### Implantation parameters

Ion energies 10-200 keV

Implantation depths 10-500 nm

Doses  $10^{11}$  to  $10^{16}$  ions/cm<sup>2</sup>, what corresponds concentrations ca.  $10^{15}$  cm<sup>-3</sup> to  $10^{20}$  cm<sup>-3</sup>.

5.10<sup>15</sup> cm<sup>-2</sup> ion implant dose and depth of ca. 200 nm translates to ca. 25 Ohm/sq sheet resistance

![](_page_27_Picture_0.jpeg)

### **Doping level**

Wafers always come doped: 10<sup>13</sup>-10<sup>20</sup> cm<sup>-3</sup> of dopant.

Diffusion and implantation can only add dopants  $\rightarrow$  doped region dopant concentration always higher than original wafer.

P-type 10 <sup>17</sup> cm <sup>-3</sup>	P-type 10 <sup>15</sup> cm <sup>-2</sup>
N-type 10 <sup>15</sup> cm <sup>-3</sup>	N-type 10 <sup>17</sup> cm <sup>-3</sup>

Possible

Impossible by diffusion/implantation

![](_page_28_Picture_0.jpeg)

### In which order are dopings made ?

![](_page_28_Figure_2.jpeg)

US 6297070 B1

p-epi layer growth Deepest first (NWELL)

#### Shallowest last (p+)

It is possible to do lithoimplant-litho-implant, and one annealing step to cure the damage and drive the dopants deeper, e.g. medium depth n+ and p+ in the figure could be combined this way.

![](_page_29_Picture_0.jpeg)

#### Implantation advantages

Implantation is:

- more accurate and uniform in dose control
- produces greater variety of profiles
- possible through oxide and nitride
- provides wide selection of mask materials
- less sensitive to surface cleaning procedures

![](_page_30_Picture_0.jpeg)

### Implantation vs. diffusion

- Sideways spreading in diffusion ≈ depth
- Sideways spreading in implantation ( $\Delta R_t$ ) is  $\approx 1/3$  depth
- Diffusion is high-temperature process → needs oxide or nitride mask
- Implantation is room temperature process → resist mask

but: damages after implantation are annealed at high temperature → both need ca. 1000°C

Diffusion is the best for high doping level, deep junctions and double side doping

### Epitaxy - "arranging upon"

![](_page_31_Figure_1.jpeg)

#### **Epitaxy conditions:**

Substrate and film are single crystalline Crystal lattices are closely matching Atomically clean substrate surface Doping can be done during film growth

#### Common epitaxy pairs (heteroepitaxy):

Si wafer –  $CaF_2$ ,  $Y_2O_3$ ,  $CoSi_2$ ,  $CeO_2$ Sapphire wafer - Si, GaN  $CeO_2$  film – YBCO (yttrium barium copper oxide) GaAs wafer – GaAlAs/GaAs

![](_page_32_Picture_0.jpeg)

# Homoepitaxy

Crystalline film A on top a crystalline wafer A

![](_page_32_Figure_3.jpeg)

Single crystal wafer

Epitaxial layer of the same material deposited on top

#### Why epitaxy of *c*-Si on *c*-Si (homoepitaxy)?

- 1. Freedom in the order of doping
- 2. Absence of O<sub>2</sub> and C contaminations

# A Preferable nucleation places: kink growth model

![](_page_33_Figure_1.jpeg)

https://www.physik.uni-kl.de/hillebrands/research/methods/molecular-beam-epitaxy/

# A Miscutting of (111) wafers for epitaxy

![](_page_34_Figure_1.jpeg)

![](_page_35_Picture_0.jpeg)

### Vapor-Phase Epitaxy (VPE)

#### VPE is modified CVD

![](_page_35_Figure_3.jpeg)

![](_page_36_Picture_0.jpeg)

![](_page_36_Figure_1.jpeg)

Lightly doped epi Heavily doped substrate

Because epitaxy is a high temperature process, dopant atoms diffuse during epitaxy.

Dopant diffusion from high concentration to low concentration.

Epi doping level is independent of substrate doping level, but the interface is not sharp due to diffusion.

![](_page_37_Picture_0.jpeg)

### Selective epitaxy

![](_page_37_Figure_2.jpeg)

![](_page_38_Picture_0.jpeg)

### Doping and epitaxy

P-type 10 <sup>16</sup> cm <sup>-3</sup>		P-type 10 <sup>18</sup> cm <sup>-3</sup>
P-type 10 <sup>18</sup> cm <sup>-3</sup>	VS.	N-type 10 <sup>15</sup> cm <sup>-3</sup>

Epitaxy is the only way to get this.

Thick doped layer or uniform dopant profile.

![](_page_39_Picture_0.jpeg)

#### Epitaxial layer vs. diffused layer

![](_page_39_Figure_2.jpeg)

![](_page_40_Picture_0.jpeg)

### Epitaxy advantages

- Epitaxy is suitable for uniform doping of monocrystalline
  Si layers from 100 nm to 100 µm
- Purity of epilayers is higher than Si substrate one
- Epitaxy is limited by monocrystalline substrate with matching lattice cells