

# ECE 4813

# Semiconductor Device and Material Characterization

#### Dr. Alan Doolittle School of Electrical and Computer Engineering Georgia Institute of Technology

As with all of these lecture slides, I am indebted to Dr. Dieter Schroder from Arizona State University for his generous contributions and freely given resources. Most of (>80%) the figures/slides in this lecture came from Dieter. Some of these figures are copyrighted and can be found within the class text, *Semiconductor Device and Materials Characterization*. <u>Every serious</u> *microelectronics student should have a copy of this book!* 



# Doping

**Plasma Resonance Free Carrier Absorption Infrared Spectroscopy Photoluminescence** Hall Measurements Magnetoresistance Time of Flight



- Electron/hole plasmas are ensembles of free carriers that can, at certain frequencies, oscillate in concert as a group.
- Such a plasma resonance exists whose frequency / wavelength (v=c/λ) is determined by the free carrier density.

$$\lambda_{plasma} = \frac{2\pi c}{q} \sqrt{\frac{K_s \varepsilon_0 m^*}{n}}$$

- Good for p (or n) > ~10<sup>18</sup> cm<sup>-3</sup>
- Since plasma resonances are hard to detect in practice, most of the time, free carrier densities are determined by empirical reflectivity minimums...

$$n = \left(A\lambda_{plasma} + B\right)^{C}$$

... or free carrier absorption



## **Free Carrier Absorption**

- Free carrier absorption occurs within the conduction or valance band (not between).
  - For example a conduction electron is absorbs an IR photon and is promoted into a higher energy state still inside the conduction band



In practice, empirical fitting is used

$$\alpha_{fc} = A\lambda^2 p$$

- Generally measured using Fourier Transmission Infrared (FTIR) Spectrometer
- Good for p (or n) > ~10<sup>17</sup> cm<sup>-3</sup>



## Infrared Spectroscopy

- Measures dopant concentrations (low temperature) and can be used at room temperature to measure free carrier absorption.
- Cryogenic temperatures are used to freeze carriers into their dopant ground state
- IR (very small energy) light is used to excite electrons/holes into their dopant "excited state" creating sharp absorption lines
- Absorption line intensity is calibrated to dopant density
- **Good for**  $N_A$  **(or**  $N_D$ **) > ~10<sup>11</sup> cm<sup>-3</sup>**







## Interferometer

- Let source be cos2π*fx* 
  - f: frequency of light
  - x: movable mirror location
- $L_1 = L_2$ 
  - Constructive interference
  - Maximum detector output
- $L_1 = L_2 + \lambda/4$ 
  - Destructive interference
  - Zero detector output





## **Fourier Transform Infrared Spectroscopy**

#### Fourier transform infrared spectroscopy (FTIR)

 $I(x) = B(f)[1 + \cos 2\pi x f] \qquad I(x) = \int_0^f B(f)[1 + \cos 2\pi x f] df$  $I(x) = \int_0^{f_1} A \cos 2\pi x f \, df = A f_1 \frac{\sin 2\pi x f_1}{2\pi x f_1}$ 



In a Fourier spectrum, frequency bandwidth determines resolution.

Interferogram

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## Interferogram - Spectrum



www.chem.orst.edu/ch361-464/ch362/irinstrs.htm



## **FTIR Applications**

 Determine oxygen and carbon density by transmission dip





## **Mobilities**

- **Conductivity** Mobility:  $\mu_p = 1/qp\rho$ 
  - Majority carrier mobility; need carrier concentration and resistivity
- *Drift* Mobility:  $\mu_p = v_d/\varepsilon$ 
  - Minority carrier mobility
  - Need drift velocity and electric field (Haynes-Shockley experiment)
- Hall Mobility:  $\mu_H = R_H / \rho$ 
  - Need Hall measurement
  - Hall mobility does not necessarily equal conductivity mobility
- *MOSFET* Mobility:
  - MOSFET mobility lowest, carriers are scattered at the Si-SiO<sub>2</sub> interface
  - Interface is microscopically rough



## Mobility

- Electron/hole mobility is a measure of carrier scattering in the semiconductor
  - Lattice scattering
    - Silicon atoms
  - Ionized impurity scattering
    - Dopant atoms
  - Interface scattering
    - Surface roughness at SiO<sub>2</sub>/Si interface
    - Polar scattering
- Silicon bulk

• 
$$\mu_n \approx 3 \ \mu_p$$
  
• MOSFET mobility (effective mobility)  $\approx 0.3$  bulk mobility  
 $\frac{1}{\mu} = \frac{1}{\mu_l} + \frac{1}{\mu_i} + \frac{1}{\mu_s}$ 



#### Ionized Impurity Scattering:





Courtesy of M.A. Gribelyuk, IBM.

### **Mobilities**

 For bulk semiconductors, lattice and ionized impurity scattering dominate the mobility





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## "Simple" Hall Effect

- Hall effect is commonly used during the development of new semiconductor material
- Resistivity, carrier concentration AND mobility can all be determined simultaneously
- Lorentz Force deflection of free carriers by an applied magnetic field
- Temperature dependent Hall is very powerful and can elucidate scattering mechanisms (plotting mobility vs T<sup>a</sup>), and determine dopant activation energies
  - Compensated Dopant Freeze out regime Arrhenius slope results in E<sub>A</sub>
  - Uncompensated Dopant Freeze out regime Arrhenius slope results in  $E_A/2$
  - At moderate temperatures,  $p \sim (N_A N_D)$
  - At elevated temperatures, p ~ n<sub>i</sub>

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## **"Simple" Hall Effect**





## **"Simple" Hall Effect**

 Resistivity is simply found from the voltage drop along the length (no magnetic field),

$$\rho = \frac{dw \, V_{\rho}}{s \, I}$$

**Carrier density**  $p = \frac{r}{qR_{H}}; n = -\frac{r}{qR_{H}};$ 

(r ~ 1 - 2, Hall scattering factor)



$$p = \frac{r}{qR_H}$$

$$q\mu_p p = \sigma = \frac{r\mu_p}{R_H} = \frac{\mu_H}{R_H}$$
$$\mu_H = |\sigma R_H| = r\mu_p$$



## **Detailed Hall Effect**

The Hall Coefficient for both electrons and holes present in the same material is in general:

$$R_{H} = r \left( \frac{\left( p - \left(\frac{\mu_{n}}{\mu_{p}}\right)^{2} n \right) + (\mu_{n}B)^{2}(p-n)}{q \left[ \left( p + \left(\frac{\mu_{n}}{\mu_{p}}\right) n \right)^{2} + (\mu_{n}B)^{2}(p-n)^{2} \right]} \right)$$



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### **Detailed Hall Effect**

#### The Hall Coefficient is in general:

$$R_{H} = r \left( \frac{\left( p - \left(\frac{\mu_{n}}{\mu_{p}}\right)^{2} n \right) + (\mu_{n}B)^{2}(p-n)}{q \left[ \left( p + \left(\frac{\mu_{n}}{\mu_{p}}\right) n \right)^{2} + (\mu_{n}B)^{2}(p-n)^{2} \right]} \right)$$

• At low fields (B<<1/ $\mu_n$ )...

$$R_{H} = \frac{r\left(p - \left(\frac{\mu_{n}}{\mu_{p}}\right)n\right)}{q\left(p + \left(\frac{\mu_{n}}{\mu_{p}}\right)n\right)^{2}} \approx \frac{r}{qp} \quad or \quad -\frac{r}{qn}$$
$$r = \frac{\langle \tau^{2} \rangle}{\langle \tau \rangle^{2}} \quad \text{where}$$

where  $\tau$  is the mean time between collisions

• And at high fields (B>>1/ $\mu_n$ )...  $R_H = \frac{r}{q(p-n)}$ 



## **Two Layer Hall Effect**

- Sometimes, a semiconductor has two different conduction layers (surface inversion, fermi-level pinning, substrate layers, n-p junctions or p+/n or n+/n layers)
- The Hall coefficient is then a weighted sum of both layers and can be either positive or negative leading to confusion (shown for the low B field limit):

$$R_{H} = R_{H1} \left( \frac{t_{1}}{t_{total}} \right) \left( \frac{\sigma_{1}}{\sigma} \right)^{2} + R_{H2} \left( \frac{t_{2}}{t_{total}} \right) \left( \frac{\sigma_{2}}{\sigma} \right)^{2}$$
  
where  $\sigma = \left( \frac{t_{1}}{t_{total}} \right) \sigma_{1} + \left( \frac{t_{2}}{t_{total}} \right) \sigma_{2}$ 

and  $t_{total} = t_1 + t_2$ 





# Two Layer Hall Effect (more detail)

The Hall coefficient is then a weighted sum of both layers and can be either positive or negative leading to confusion:

$$R_{H} = \begin{pmatrix} \frac{t_{total} \left[ \left( R_{H1} \sigma_{1}^{2} t_{1} + R_{H2} \sigma_{2}^{2} t_{2} \right) + R_{H1} \sigma_{1}^{2} R_{H2} \sigma_{2}^{2} \left( R_{H1} t_{2} + R_{H2} t_{1} \right) B^{2} \right] \\ \hline \left( \sigma_{1} t_{1} + \sigma_{2} t_{2} \right)^{2} + \sigma_{1}^{2} \sigma_{2}^{2} \left( R_{H1} t_{2} + R_{H2} t_{1} \right) B^{2} \\ \end{pmatrix} \\ Low B \\ Field \\ R_{H} = R_{H1} \left( \frac{t_{1}}{t_{total}} \right) \left( \frac{\sigma_{1}}{\sigma} \right)^{2} + R_{H2} \left( \frac{t_{2}}{t_{total}} \right) \left( \frac{\sigma_{2}}{\sigma} \right)^{2} \\ where \quad \sigma = \left( \frac{t_{1}}{t_{total}} \right) \sigma_{1} + \left( \frac{t_{2}}{t_{total}} \right) \sigma_{2} \\ and \quad t_{total} = t_{1} + t_{2} \end{cases}$$



# Two Layer Hall Effect (more detail)

The Hall coefficient is then a weighted sum of both layers and can be either positive or negative leading to confusion (generally):

$$R_{H} = \left(\frac{t_{total}\left[\left(R_{H1}\sigma_{1}^{2}t_{1}+R_{H2}\sigma_{2}^{2}t_{2}\right)+R_{H1}\sigma_{1}^{2}R_{H2}\sigma_{2}^{2}\left(R_{H1}t_{2}+R_{H2}t_{1}\right)B^{2}\right]}{\left(\sigma_{1}t_{1}+\sigma_{2}t_{2}\right)^{2}+\sigma_{1}^{2}\sigma_{2}^{2}\left(R_{H1}t_{2}+R_{H2}t_{1}\right)B^{2}}\right)$$





## Hall Effect Measurements

#### Two approaches:

#### Hall Bar (5 or 6 contacts)



Van der Pauw configuration

- Based on Conformal mapping theory
- Contacts assumed point sources
- Uniform thickness
- Cannot contain isolated (interior) holes





## Hall Effect Measurements



#### Van der Pauw configuration

#### • Measure resistivity first by "perimeter measurements"...

( ( ) )

Example: determine R<sub>12,34</sub> where current goes in 1 and leaves 2 and voltage is measured between terminals 3 and 4. Next determine R<sub>23,14</sub> where current goes in 2 and leaves 3 and voltage is measured between terminals 1 and 4.

• Use:

$$\rho = \left(\frac{\pi t}{\ln(2)}\right) \left(\frac{R_{12,34} + R_{23,14}}{2}\right) F$$



• ...where F is a symmetry term derived from conformal mapping theory

• F is determined from:

$$\left(\frac{R_r - 1}{R_r + 1}\right) = \frac{F}{\ln(2)} \cosh^{-1} \left(\frac{e^{\left(\frac{\ln(2)}{F}\right)}}{2}\right)$$

where 
$$R_r = \frac{R_{12,32}}{R_{23,12}}$$





Differs some from your text. For details see http://www.nist.gov/pml/semiconductor/hall\_resistivity.cfm



# Hall Effect Measurements

- Van der Pauw configuration
  - Now measure the Hall voltage using "Crossing configurations"
    - Example: Apply the magnetic field and determine V<sub>13,24P</sub> where current goes in 1 and leaves 3 and voltage is measured between terminals 2 and 4. Next reverse the field and determine V<sub>13,24N</sub> again.
    - To find the sheet concentration (#/cm<sup>2</sup>) use:



...where we have intentionally left out the proportionality constant

 In reality, 8 resistivity and eight hall voltage measurements are made to reduce contact related offset voltage errors resulting in an equation that is of the form:

$$\rho = \left(\frac{(8x10^{-8})IB}{q[(V_{13,24P} - V_{13,24N}) + (V_{31,42P} - V_{31,42N}) + (V_{42,13P} - V_{42,13N}) + (V_{24,31P} - V_{24,13N})]}\right)$$

See text for important sample geometry considerations (if ignored, significant error can result)

Coordinate

System

 $V_{\rm H} = V_{24\rm P}$ 

I13





## **Haynes - Shockley Experiment**

 Allows mobility, diffusion constant, and minority carrier lifetime to be determined







## **Haynes - Shockley Experiment**

 Allows mobility, diffusion constant, and minority carrier lifetime to be determined



If at least two lengths and two times are measured, the FWHM of the time plot, ∆t can be used to...

$$t_{d} = d / v_{drift}; \quad \mu_{n} = \frac{d}{\varepsilon t_{d}}; \quad D_{n} = \frac{(d\Delta t)^{2}}{16t_{d}^{3}\ln 2}; \quad \tau_{n} = \frac{t_{d2} - t_{d1}}{\ln(V_{out1} / V_{out2}) - 0.5\ln(t_{d2} / t_{d1})}$$

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## Haynes – Shockley Experiment

$$\Delta n(x,t) = \frac{N}{\sqrt{4\pi D_n t}} \exp\left(-\frac{(x-vt)^2}{4D_n t} - \frac{t}{\tau_n}\right)$$





# **MOSFET** Effective Mobility

- Effective mobility determined from drain current drain voltage characteristics
- The MOSFET drain current for small V<sub>D</sub> (50 100 mV) is

 $I_{D} = (W/L)\mu_{eff}Q_{N}V_{D} \approx (W/L)\mu_{eff}C_{ox}(V_{G} - V_{T})V_{D}$ 

• Determine  $g_d = \Delta I_D / \Delta V_D$  for low  $V_D$ 

Solve for  $\mu_{eff}$ 

$$\mu_{eff} = \frac{Lg_d}{WQ_N} \approx \frac{Lg_d}{WC_{ox}(V_G - V_T)}$$

• Need  $g_d$ ,  $Q_N$ ,  $V_T$ 



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#### **MOSFET** *Effective* Mobility

 $I_D - V_D \Rightarrow g_d; C_{GC} - V_G \Rightarrow Q_N; I_D - V_G \Rightarrow V_T$ 



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#### **MOSFET** *Effective* Mobility



 $\mu_o =$  low-field mobility;  $\theta =$  mobility degradation factor



## **Review Questions**

- What are the different mobilities?
- Why is the MOS effective mobility less than the bulk mobility?
- How is μ<sub>eff</sub> most commonly determined?
- Why does the Hall mobility differ from the conductivity mobility
- How does a Hall mobility measurement work?
- How does the Haynes-Shockley experiment work?
- What is determined with the Haynes-Shockley experiment ?
- For what is the time-of-flight technique used?