# **Chapter 2 On Few Electronic Properties of Nanowires of Heavily Doped Biosensing Materials**



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**Abstract In this chapter, we study the** effective electron mass (EEM), the **Einstein relation for the diffusivity–mobility ratio (ER)**, **the Einstein's photoemission (EP)**, the field emission (FE) **and** the thermo-electric power (TP) in **heavily doped nanowires (HDNWs) of** different **biosensing materials** together with the relative comparison of the said transport features with that of the HDNW compounds. The EEM is an important transport quantity which is used in the analysis of different devices of low-dimensional electronics. The ER is useful in the characterizations of various types of hetero-structures and occupies a central position in the field of materials science. The EP is a physical phenomenon which finds extensive application in modern opto-electronics, and the FE is a quantum mechanical process. Besides, with the advent of quantum Hall effect, there has been considerable interest in studying the TP for various low-dimensional compounds. Although biosensing materials find wide applications and many physical properties have already been studied, nevertheless the investigations of the said electronic quantities for nanowires (NWs) of heavily doped (HD) biosensing materials are becoming increasingly important. Keeping this in mind in this chapter, an attempt is made to study the aforesaid quantities, **talking HDNWs of** various **biosensing material**s. We observe that the EEM is quantum number dependent. The ER oscillates with the electron statistics  $(n_0)$  and the magni**tude and nature of oscillations are totally different as compared with the ER in**

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**HDNWs of other materials talking HDNW of InSb as an example.** T**he Einstein's photo current from HDNWs of** different biosensing materials **also oscillates with**   $n_0$  in radically different fashion as found from HDNWs of other materials. The **field emitted current oscillates with increase in electric field due to van Hove**  singularities and the TP increases with increasing  $n_0$  in oscillatory ways. The **most important realization is that the quantum signatures in all the cases are not only totally different, but also the variations of the said electronic quantities as compared with that of HDNWs different compounds excluding biomaterials are also different**.

#### **2.1 Introduction**

The EEM  $[1-4]$  $[1-4]$ , ER  $[5-8]$  $[5-8]$ , EP  $[9-12]$  $[9-12]$ , FE  $[13-16]$  $[13-16]$  and TP  $[17-20]$  $[17-20]$  have extensively been investigated in the recent literature, and they have important contributions in controlling control the transport phenomena in biosensing materials. Although biosensing materials find wide applications and many physical properties have already been studied  $[21-37]$  $[21-37]$  $[21-37]$ , nevertheless it appears from the literature that the study of the said electronic properties has yet to be made. In this chapter, they are being investigated in HDNWs of biosensing materials. It may be noted that HDNWs are also being studied by various workers [[38–](#page-8-1)[40\]](#page-8-2). The theoretical background is described in Sects. [2.2](#page-1-0), and [2.3](#page-2-0) contains the results and discussion in this context.

#### <span id="page-1-0"></span>**2.2 Theoretical Background**

The  $E - k_x$  relation assumes the form [\[37](#page-8-0)]

$$
k_x^2 = A_{11}(E, \eta_g, n_y) \tag{2.1}
$$

where

$$
A_{11}(E, \eta_g, n_y)
$$
  
=  $\left[\frac{2}{\sqrt{3}}\cos^{-1}\left[\left[f\gamma(E, \eta_g) + g\right]^2 - 3 - D - 2\cos\left(\frac{n_y\pi}{d_y}\right)\right]\right] \left(4\cos\left(\frac{3\pi n_y}{2d_y}\right)\right)\right]^2$ 

and the other notations are defined in [\[37](#page-8-0)]

The use of (2.1) leads to the expression of EEM as

<span id="page-1-1"></span>
$$
m^*(E_F, \eta_g, n_y) = \frac{\hbar^2}{2} A'_{11}(E_F, \eta_g, n_y)
$$
 (2.2)

where the notations have their usual significances.

The  $n_0$  can be written as

$$
n_{y} = \frac{2g_{v}}{\pi} \sum_{n_{y}=1}^{n_{y_{\text{max}}}} \left[ \sqrt{A_{11}(E_{F}, \eta_{g}, n_{y})} + \sum_{r=1}^{r=n} 2(1 - 2^{1-2r})\xi(2r) \frac{\partial^{2r}}{\partial E_{F}^{2r}} \left[ \sqrt{A_{11}(E_{F}, \eta_{g}, n_{y})} \right] \right]
$$
(2.3)

where the notations have their usual significances.

The ER can be expressed as

$$
\frac{D}{\mu} = \left(\frac{n_0}{e}\right) \left[\frac{\partial n_0}{\partial (E_F - Z)}\right]^{-1} \tag{2.4}
$$

where *Z* is given by

<span id="page-2-1"></span>
$$
A_{11}(Z, \eta_g, n_y) = 0 \tag{2.5}
$$

Thus by using  $(2.2)$  $(2.2)$  $(2.2)$ – $(2.4)$  $(2.4)$  $(2.4)$ , we can study the DMR numerically. Incidentally, the photo current  $I$  can be written as

$$
I = \frac{\alpha_0 e g_v k_B T}{\pi \hbar} \sum_{n_y=1}^{n_{\text{ymax}}} \ln(1 + \exp[(E_F - (Z + W - h\upsilon)) (k_B T)^{-1}] \tag{2.6}
$$

where  $\alpha_0$  is the probability of photoemission

The field emitted current  $(i_f)$  assumes the form

$$
I = \frac{2eg_vk_BT}{h} \sum_{n_y=1}^{n_{\text{ymax}}} [\ln(1 + \exp[(E_F - Z)(k_BT)^{-1}] \exp(-Q)] \tag{2.7}
$$

where

$$
Q = \frac{4[A_{11}(V_0, \eta_g, n_y)]^{3/2}}{3eF_x[A'_{11}(V_0, \eta_g, n_y)]}, \ V_0 = E_F + \phi_w
$$

## <span id="page-2-0"></span>**2.3 Results and Discussion**

The plot of the normalized EEM in HDNWs of  $MOS_2$  versus  $n_0$  for three sub-bands is given in Fig. [2.1.](#page-3-0) The plots of normalized DMR  $(\overline{D})$  in HDNWs of MOS<sub>2</sub> versus

*n*0 are given in Figs. [2.2](#page-3-1), and [2.3](#page-4-0) shows the same for HDNWs of InSb for the purpose of relative comparison. Figures [2.4](#page-4-1) and [2.5](#page-4-2) explore the normalized photo current  $(\overline{I})$ from HDNWs of  $MOS_2$  versus  $n_0$  and the same for HDNWs of InSb respectively. Figures [2.6](#page-5-0) and [2.7](#page-5-1) exhibit the plots of the normalized FE and TP for different HDNW biomaterials versus  $n_0$  respectively.

The salient features are given below:

- 1. In Fig.  $2.1$ , the EEM increases with increasing  $n_0$  where the value of EEM for  $n<sub>y</sub> = 1$  is the greatest.
- 2. In Fig. [2.2](#page-3-1), the  $\overline{D}$  in HDNWs of MOS<sub>2</sub> oscillates with enhanced  $n_0$ , and the magnitude and nature of oscillations are totally different as compared with the  $\overline{D}$

<span id="page-3-1"></span><span id="page-3-0"></span>

 $n_0(10^{10}m^{-1})$ 



<span id="page-4-0"></span>**Fig. 2.3** Plot of the  $\overline{D}$  versus *n*<sub>0</sub> for the NWs of *n* − InSb with two different values of  $d_y$ 



<span id="page-4-2"></span><span id="page-4-1"></span>**Fig. 2.4** Plot of the  $\overline{I}$  from NWs of MOS<sub>2</sub> versus  $n_0$  for four different values of film thickness

**Fig. 2.5** Plot of the  $\overline{I}$  versus *n*<sup>0</sup> for the NWs of *n* − InSb





<span id="page-5-0"></span>**Fig. 2.6** Plot of the normalized field emitted current versus electric field for three different HDNWs of biomaterials



<span id="page-5-1"></span>**Fig. 2.7** Plot of the normalized TP versus  $n_0$  for four different HDNWs of biomaterials as shown by a, b, c and d, respectively

in HDNWs of other material as given in Fig. [2.3](#page-4-0). The quantum signatures of two different types of 1D motion can be assessed by comparing Figs. [2.2](#page-3-1) and [2.3](#page-4-0).

- 3. From Figs. [2.4](#page-4-1) and [2.5](#page-4-2), it appears that the  $\overline{I}$  HDNWs of MOS<sub>2</sub> oscillates with  $n_0$  in radically different manner as compared with that from HDNWs of other materials.
- 4. From Fig. [2.6](#page-5-0), we note that the field emitted current oscillates with increase in electric field due to Van Hove singularities
- 5. From Fig. [2.7](#page-5-1), we note that the TP increases with increasing  $n_0$  in oscillatory ways.

**Most important to realize is that the quantum signatures in all the cases are not only totally different, but also the variations of the said electronic quantities as compared with that of HDNWs different materials excluding biocompounds are also different**.

#### **2.4 Conclusion**

**In this chapter, we study the** EEM, **ER, EP**, FE **and** the TP in **heavily doped nanowires (HDNWs) of** different **biosensing materials** together with the relative comparison of the said transport features with that of the HDNW compounds. We observe that the EEM is quantum number dependent. T**he ER oscillates with** the electron statistics  $(n_0)$ , and the magnitude and nature of oscillations are totally **different as compared with the ER in HDNWs of other materials talking HDNW of InSb as an example.** T**he Einstein's photo current from HDNWs of** different biosensing materials **also oscillates with**  $n_0$  **in radically different fashion as found from HDNWs of other materials.** The field emitted current oscillates with increase in electric field due to Van Hove singularities, and the TP increases with increasing  $n_0$  in oscillatory ways. **The most important realization is that the quantum signatures in all the cases are not only totally different, but also the variations of the said electronic quantities as compared with that of HDNWs different compounds excluding biomaterials are also different.** 

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