

Fachbereich 08 - Biologie und Chemie  
Institut für Anorganische und Analytische Chemie  
der Justus-Liebig-Universität Giessen

# **Label-free electrical biosensing based on electrochemically functionalized carbon nanostructures**

DISSERTATION

to apply for the degree of „Doctor rerum naturalium“  
(Dr. rer. nat.)

submitted by

**Tetiana Kurkina**  
born March 01, 1984  
in Brovary, Kyiv obl. (Ukraine)

Gießen, February 2012

REFEREES:           1. PROF. DR. BERNHARD SPENGLER <sup>1</sup>  
                          2. PROF. DR. JÜRGEN JANEK <sup>2</sup>  
                          3. PROF. DR. KLAUS KERN <sup>3,4</sup>

THE FOLLOWING WORK WAS CARRIED OUT BETWEEN DECEMBER 2008 AND DECEMBER 2011  
AT THE MAX PLANCK INSTITUTE FOR SOLID STATE RESEARCH, STUTT GART, GERMANY.

---

<sup>1</sup> Institut für Anorganische und Analytische Chemie, Justus-Liebig-Universität Gießen

<sup>2</sup> Physikalisch-Chemisches Institut, Justus-Liebig-Universität Gießen

<sup>3</sup> Max-Planck-Institut für Festkörperforschung, Stuttgart

<sup>4</sup> Institut de Physique de la Matière Condensée, Ecole Polytechnique Fédérale de Lausanne,  
Switzerland

# Abstract

---

The development of new analytical approaches in the diagnostics of the diseases with higher throughput, smaller sample and set-up sizes, lower cost and easier disposal is one of the major needs of modern medicine. Miniaturization and simplification of biomedical assays are required for point of care diagnostics and lab-on-a-chip systems. In this thesis the concept of electrical biosensors based on single wall carbon nanotubes (SWCNTs) and graphene is presented. The detection of saccharides and DNA was realized using field effect transistor (FET) - based sensors where carbon nanostructures play the role of a transducing component.

The theoretical part of the thesis explains the concept of biosensing and the role of nanomaterials in the development of the next generation of bioassays. Furthermore, the structure and properties of SWCNTs and graphene and their advantages for electrical biosensing are described.

The experimental section starts with a detailed description of the carbon nanotube (CNT) biosensor fabrication process. This includes carbon nanotube solution preparation, assembly of CNTs into devices, passivation of metal electrodes and modification of the CNT surface with receptors. The advantages of using impedance spectroscopy measurements at different liquid gate voltages for electrical detection of biomolecules are pointed out here.

The next chapter is dedicated to affinity-based glucose sensing. Using boronic acid functionalized carbon nanotubes the detection of glucose was demonstrated. The sensing mechanism was investigated in detail. The sensor signal was shown to be different depending on the way the CNTs were modified - covalently or non-covalently.

The biosensing setup was then utilized for the detection of DNA. It was possible to achieve very low limit of detection for oligonucleotides using CNTs non-covalently modified with a complementary DNA sequence. The sensor was shown to be highly selective as well.

Finally, the possibility of using 2D-carbon nanomaterial, namely graphene, for electrical biosensing is outlined. The approaches towards large scale preparation of graphene devices were investigated during this work. The wafer-scale

fabrication of reduced graphene oxide devices was realized using a novel chemical route.

The final part of the thesis summarizes the results obtained while conducting this work. The designed biosensing platforms based on carbon nanostructures show a great promise for application in chemical analysis and medical diagnostics. Therefore the developed biosensor is planned to be adapted for the detection of analytes from biological liquids.

*Keywords:* biosensor, label-free detection, carbon nanotubes, graphene, electrical detection

# Zusammenfassung

---

Die Entwicklung neuer analytischer Ansätze mit höherem Durchsatz, geringeren Probenmengen und kleineren Geräten, niedrigeren Kosten und einfacherer Entsorgung ist eine der wichtigsten Bedürfnisse der modernen Medizin für die Diagnostik von Erkrankungen. Die Miniaturisierung und die Vereinfachung biomedizinischer Tests sind für die „Point of Care“ Diagnostik und für „Lab-on-a-Chip“ Systeme erforderlich. In dieser Arbeit wird das Konzept elektrischer Biosensoren basierend auf einwandigen Kohlenstoff-Nanoröhrchen (SWCNTs) und Graphen präsentiert. Der Nachweis von Zuckern und DNS wurde in Form von Feldeffekt-Transistor (FET) - Sensoren realisiert, wobei die elektronischen Eigenschaften von Kohlenstoff Nanostrukturen durch an deren Oberfläche stattfindenden chemische Reaktionen beeinflusst werden.

Der theoretische Teil der Arbeit erläutert das Konzept der Biosensorik, insbesondere die Rolle von Nanomaterialien in der Entwicklung der nächsten Generation von biologischen Assays. Des Weiteren sind die Struktur und die Eigenschaften von SWCNTs und Graphen und deren Vorteile für die elektrische Bioanalytik beschrieben.

Der experimentelle Teil beginnt mit einer ausführlichen Beschreibung der Herstellung von SWCNT-Biosensoren. Dazu gehören die Dispergierung von Kohlenstoff-Nanoröhrchen (CNT) in Lösung, die Integration von CNTs in elektronische Bauteile, die Passivierung von Metallelektroden und die Modifizierung der CNT Oberfläche mit Rezeptoren. Außerdem enthält dieses Kapitel eine Erklärung der Detektionsmethode und weist auf die Vorteile des gewählten Ansatzes für die elektrische Detektion von Biomolekülen hin.

Das nächste Kapitel befasst sich mit der Affinität-basierten Detektion von Zucker. Mittels Boronsäure-Rezeptoren wurde Glucose nachgewiesen. Der Detektionsmechanismus wurde im Detail untersucht. Es konnte gezeigt werden, dass sich je nach Modifizierungsart (kovalente oder nicht-kovalente Bindung von Rezeptoren) das Sensorsignal unterschiedlich verhält.

Derselbe Biosensorik Aufbau wurde dann für den Nachweis von DNS verwendet. Es war möglich, eine sehr niedrige Nachweisgrenze für Oligonukleotide mit Hilfe von CNTs zu erreichen, die nicht-kovalent mit einer komplementären DNS-Sequenz modifiziert waren. Zusätzlich war der Sensor selektiv für die Zielsubstanz.

Des Weiteren wurde der Einsatz von 2D-Kohlenstoff-Nanomaterial, nämlich Graphen, für die elektrische Biosensorik erforscht. In diesem Kapitel werden Ansätze zur großtechnischen Herstellung von Graphen Bauteilen beschrieben. Die „Wafer-Scale“-Fertigung von reduzierten Graphenoxid Bauteilen unter Verwendung einer neuartigen chemischen Methode wurde realisiert.

Der letzte Teil fasst die Ergebnisse dieser Arbeit zusammen. Die entworfenen Biosensorik Konzepte basierend auf Kohlenstoff-Nanostrukturen sind vielversprechend für die Anwendung in der chemischen Analytik und der medizinischen Diagnostik. Daher ist es geplant, den entwickelten Biosensor für den Nachweis von Analyten aus biologischen Flüssigkeiten anzupassen.

*Schlagwörter:* Biosensor, Marker-freie Erkennung, Kohlenstoff-Nanoröhrchen, Graphen, elektrische Detektion

# List of used acronyms

---

1D	-	one dimensional
2D	-	two dimensional
ABA	-	aminobenzoic acid
AFM	-	atomic force microscopy
APBA	-	aminophenylboronic acid
BA	-	boronic acid
CE	-	counter electrode
CNT	-	carbon nanotube
CVD	-	chemical vapor deposition
DEP	-	dielectrophoresis
DMF	-	dimethylformamide
DWCNT	-	double wall carbon nanotube
ECM	-	electrochemical modification
EDC	-	ethylenediaminechloride
FET	-	field-effect transistor
GO	-	graphene oxide
GOx	-	glucose oxidase
ISFET	-	ion-selective field-effect transistor
LoC	-	lab on a chip
$\mu$ TAS	-	micro total analytical system
MWCNT	-	multi wall carbon nanotube
ncDNA	-	non-complementary DNA
NHS	-	N-hydroxysuccinimide
PDMS	-	polydimethylsiloxane
PoC	-	point of care
pTy	-	polytyramine
RB	-	Rhodamin B
RE	-	reference electrode
RGO	-	reduced graphene oxide
RT-PCR	-	real-time polymerase chain reaction
SDBS	-	sodium dodecyl benzene sulfonate
SDS	-	sodium dodecyl sulfate
SWCNT	-	single wall carbon nanotube
WE	-	working electrode

# List of figures

Figure 1.1. Schematic of a back-gated (a) and a liquid gated FET.....	- 17 -
Figure 1.2. Representation of the phase of impedance .....	- 18 -
Figure 2.1. Physical structure of CNTs. ....	- 20 -
Figure 2.2. The structure of double walled carbon nanotube (DWCNT) and multi-walled carbon nanotube (MWCNT) .....	- 21 -
Figure 2.3. Overview of different CNT-based biosensing strategies. ....	- 22 -
Figure 2.4. The structure of graphene. ....	- 23 -
Figure 2.5. Non-covalent functionalization of carbon nanostructures with pyrene-derivatives via $\pi$ - $\pi$ stacking. 1-pyrenebutanoic acid succinimidyl ester (PBASE) attached to a) graphene and b) CNT.....	- 25 -
Figure 2.6. Various covalent functionalization schemes for graphene oxide. ....	- 26 -
Figure 2.7. Addition reactions for the functionalization of the carbon nanotube sidewall .....	- 27 -
Figure 2.8. Schemes for (a) oxidative ECM with 4-aminobenzylamine and (b) reductive ECM with an aryl diazonium salt. ....	- 28 -
Figure 3.1. Visualization of the same SWCNT device with AFM and confocal microscopy.....	- 31 -
Figure 3.2. Schematic of the dielectrophoretic manipulation of 1D-nanostructures.....	- 32 -
Figure 3.3. Schematic of the dielectrophoretic trapping setup .....	- 33 -
Figure 3.4. Different electrode geometries used in this study. ....	- 34 -
Figure 3.5. One of the possible ways to upscale the DEP trapping of CNTs. ....	- 35 -
Figure 3.6. Monitoring the dielectrophoretic trapping of the nanotubes.....	- 36 -
Figure 3.7. Passivation of Pt electrodes on Si/SiO <sub>2</sub> surface.....	- 38 -
Figure 3.8. Scheme showing the functionalization strategy to obtain the BA-CNT-sensors.....	- 40 -
Figure 3.9. Gate dependence of the devices during (a) covalent and (b) non-covalent electrochemical functionalization of CNTs with APBA. ....	- 41 -
Figure 3.10. Electrochemical functionalization of CNTs. ....	- 42 -
Figure 3.11. Detection strategy .....	- 43 -
Figure 4.1. Sensing in the open system without applying liquid gate voltage.....	- 46 -
Figure 4.2. (a) An AFM image of the electrode region of a BA-CNT sensor after the sensing trials. (b) A schematic representation of the microfluidic set up with the location of the nanotube device. (c) a photograph of the final assembled device.....	- 47 -
Figure 4.3. Affinity sensing of saccharides.....	- 48 -
Figure 4.4. 2D-impedance (Z-) maps showing the impedance as a function of frequency and liquid gate voltage. Magnitude [Ohms] maps of the impedance of covalently functionalized BA-CNT devices are shown in buffer (a) and in 10 mM glucose (b).. ....	- 49 -
Figure 4.5. Reversibility of the binding of glucose in covalent-BA-CNT-sensors. ....	- 50 -
Figure 4.6. Concentration dependence of the covalent-BA-CNT-sensors .....	- 51 -
Figure 4.7. Control experiment with unmodified CNT devices. ....	- 52 -
Figure 4.8. Selectivity of the covalent BA-CNT-sensors.....	- 53 -
Figure 4.9. Charge distribution on the surface of BA-functionalized CNTs calculated using the Henderson-Hasselbach relationship.....	- 54 -
Figure 4.10. Sensor response for non-covalent-BA-CNT devices showing the gate dependence of the magnitude of impedance at low frequencies.....	- 56 -
Figure 5.1. Overview of the assembled carbon nanotube DNA sensor.....	- 59 -
Figure 5.2. The controlled functionalization of the carbon nanotube surface with probe DNA. ....	- 61 -
Figure 5.3. Evidence for the coupling of probe DNA: schematic of the experiment (a), AFM images taken before (b) and after (c) the coupling protocol.....	- 62 -



Figure 5.4. Evidence for DNA hybridization during the sensing trial .....	- 64 -
Figure 5.5. Specificity of CNT-DNA-sensors .....	- 65 -
Figure 5.6. Control experiment with non-functionalized CNTs. ....	- 66 -
Figure 5.7. Attomolar detection limit of CNT-DNA-sensors. ....	- 67 -
Figure 5.8. Sensitivity of CNT-DNA-sensors. ....	- 68 -
Figure 5.9. Attomolar target differentiation in a heterogeneous DNA mixture. ....	- 69 -
Figure 6.1. Liquid phase exfoliation of graphene from graphite. ....	- 72 -
Figure 6.2. Preparation of graphene oxide solution .....	- 73 -
Figure 6.3. An RGO device with passivated contacts for biosensing applications. ....	- 74 -
Figure 6.4. Liquid gating of RGO device.....	- 75 -
Figure 6.5. Scheme of the chemical anchoring protocol.....	- 76 -
Figure 6.6. Surface characterization by AFM.....	- 77 -
Figure 6.7. Electronic properties of the anchored GO devices.....	- 78 -
Figure 6.8. Scaling up the chemical anchoring. ....	- 80 -
Figure 6.9. Wafer scale RGO devices with high yield.....	- 81 -
Figure 6.10. AFM image (phase mode) of graphene oxide coupled to electrochemically modified electrodes on flexible kapton substrate. ....	- 82 -
Figure 6.11. Commercially available graphene oxide coupled to the electrodes. ....	- 82 -
Figure 7.1. Sensing of let-7a-miRNA in buffer (a) and serum (b).....	- 85 -
Figure 7.2. AFM images of the device before (a) and after (b) sensing trials in serum. ....	- 86 -
Figure 7.3. Schematic of RGO-FET- immunosensor. ....	- 86 -
Figure 7.4. Sensing of beta-amyloid peptide using RGO immunosensor.....	- 87 -

# Table of Contents

---

<b>Theoretical background .....</b>	<b>- 12 -</b>
<b>1. Biosensors in medical diagnostics.....</b>	<b>- 12 -</b>
1.1. Introduction.....	- 12 -
1.2. Operating principle and types of biosensors.....	- 13 -
1.3. Nanomaterials for the detection of biomolecules.....	- 14 -
1.4. Electrical/electrochemical sensing.....	- 15 -
<b>2. Carbon nanostructures for biosensing.....</b>	<b>- 19 -</b>
2.1. Introduction.....	- 19 -
2.2. Carbon nanotubes.....	- 19 -
2.3. Graphene.....	- 23 -
2.4. Functionalization of carbon nanostructures.....	- 24 -
<b>Experimental section.....</b>	<b>- 29 -</b>
<b>3. Fabrication of CNT-based sensors.....</b>	<b>- 29 -</b>
3.1. Introduction.....	- 29 -
3.2. Preparation of CNT dispersions.....	- 29 -
3.3. Visualization of CNT devices.....	- 30 -
3.4. Dielectrophoretic manipulation of CNTs.....	- 32 -
3.5. Passivation of electrodes.....	- 37 -
3.6. Chemical functionalization of CNTs.....	- 39 -
3.7. Sensing strategy.....	- 42 -
<b>4. Sensing of saccharides .....</b>	<b>- 45 -</b>
4.1. Introduction.....	- 45 -
4.2. Device fabrication.....	- 46 -
4.3. Sensing results.....	- 48 -
4.4. Conclusions.....	- 57 -
<b>5. Label-free detection of few copies of DNA.....</b>	<b>- 58 -</b>

5.1. Introduction.....	- 58 -
5.2. Sensing setup. ....	- 59 -
5.3. Functionalization of CNTs with probe DNA.....	- 60 -
5.4. Sensing trials.....	- 63 -
5.5. Sensing results.....	- 64 -
5.6. Sensitivity and detection limit.....	- 67 -
5.7. Detection in complex media.....	- 69 -
5.8. Conclusions. ....	- 70 -
<b>6. Towards graphene-based electrical sensors - 71 -</b>	
6.1. Introduction.....	- 71 -
6.2. DEP trapping of graphene. ....	- 72 -
6.3. DEP trapping of graphene oxide.....	- 72 -
6.4. Chemical anchoring of graphene oxide for preparation of the devices.....	- 75 -
6.5. Up-scaling the fabrication of RGO devices.....	- 79 -
6.6. Conclusions. ....	- 82 -
<b>Summary and outlook ..... - 84 -</b>	
7.1. CNT based electrical sensors. ....	- 84 -
7.2. Graphene devices for biosensing.....	- 86 -
<b>Bibliography..... - 89 -</b>	
<b>Acknowledgements ..... - 103 -</b>	

# Theoretical background

---

## 1. Biosensors in medical diagnostics

**1.1. Introduction.** Early and correct diagnosis plays a very important role in modern medicine. Medical diagnostics evolved far beyond the evaluation of external symptoms. Doctors rely on the results of bio-/chemical tests some of which were unknown thirty years ago. For decades researchers have been working on the discovery of *biomarkers* - molecular indicators of different biological states of the organism. As a result, the number of biomarkers for various diseases at the different stages of pathology increased dramatically. Because of this in many cases the disease can be counteracted before the appearance of the symptoms [1]. This has a huge impact on the economic and the social life. Increasing number of biomarkers as well as the demand for fast and reliable medical tests are driving the development of new analytical approaches that can be used in clinical biochemistry. Medical diagnostics is enriched by advances in biochemistry, molecular biology, physiology, chemistry and physics (optics, mechanics, electronics) [2]. Integrated knowledge from different scientific fields is required to satisfy the need of modern society for low-cost, fast, and reliable methods for the detection of different biomarkers, the number of which is constantly growing [3]. The concept of Point of care (PoC) diagnostics adds miniaturization, simplification and disposability to the above mentioned desirable characteristics of new analytical techniques in medicine [4, 5]. One of the approaches that would fit perfectly for this purpose is biosensing [6]. Arrays of biosensors with different specificities can become a part of Micro Total Analytical Systems ( $\mu$ TAS) or lab-on-a-chip devices. In such systems biosensor arrays are integrated on a specialized platform that includes processing for biological liquids.  $\mu$ TAS can fulfill the dream of cheap and fast medical tests and increase the scope for PoC examinations [7]. Biosensors are already widely used, for example, for self-monitoring of glucose or as pregnancy tests. However there is still a need for the development of cheap and ultrasensitive biosensing platform with the possibility of multiplex detection [8].

## 1.2. Operating principle and types of biosensors. A

*biosensor* is an analytical device which uses a biological or a biologically derived recognition system associated with a physico-chemical transducer to estimate the presence and/or concentration of the target substance by translating biological reactions into a quantifiable and a processible physical signal [9]. The first biosensor was introduced by Clark in 1956 and exemplified by Clark and Lyons in 1962 [10]. It was an amperometric oxygen electrode for detecting glucose that utilized Glucose oxidase (GOx) as the recognition element. The term “biosensor” appeared after shortening the term “bioselective sensor” proposed by Rechnitz in 1977 for arginine-selective electrode that used living organisms as recognition elements [9, 11].

The two most important characteristics of biosensor are its sensitivity and selectivity towards a target molecule (*analyte*). Selectivity depends only on properties of receptor element of the biosensor since that is where the analyte interacts with biosensor. Sensitivity is determined by both the biological compound and the transducer. For high sensitivity it is important to have an excellent recognition of the analyte by the receptor element as well as a very efficient transduction of the signal to the output system. Biosensors can be classified in different ways, e.g. by the type of recognition elements, by the type of transducer or by application [9].

*The biorecognition element* determines both selectivity and sensitivity of the device. The part of biosensor that contains bioreceptors is often called a biosensitive layer. The choice and amount of bioreceptors depend on the nature of the analyte, the sample matrix, interfering substances and conditions of sensor utilization. *Affinity biosensors* use ligand-receptor interactions. Antibodies, nucleic acids, aptamers, cell receptors can be utilized as biorecognition elements of such sensors. *Catalytic biosensors* use the ability of biological system to react in the presence of the specific molecules by catalyzing the corresponding reactions. In this case enzymes, abzymes, microorganisms, plant or animal cells or tissue slices can be used as the biorecognition element. *Biomimetic receptor-based biosensors* use synthetic binding and/or catalytic systems [9, 12].

*The transducer* converts a biological effect into, for instance, an electrical signal with high sensitivity and minimum disturbance. To achieve this, biorecognition events in a sensor should take place on the surface of transducer or in a very high proximity to it. Depending on the type of transducer the following kinds of biosensors are under development: optical (luminescence, fluorescence, absorption, Raman, Surface Plasmon Resonance), electrical and electrochemical (amperometric, potentiometric, impedance-based etc.), mass-sensitive (surface acoustic wave, microbalance, etc) and thermal. Biosensors and biosensor test formats, respectively, can be classified into *labeled* and *label-free* types depending on whether

the analyte is labeled or not. Common labels include enzymes, radionuclides, nanoparticles, and fluorescent or electrochemiluminescent probes. Biosensors can be also classified depending on the way they are used. For instance, those used in medical diagnostics can be noninvasive, contacting, indwelling (minimally invasive) or implantable [9, 14].

It is important to mention some specific problems one has to overcome in order to use biosensors in medical diagnostics. For example, implantable biosensors interact with patients' body tissues to the greatest extent. The interface of such sensors becomes extremely important since the organism usually tries to eliminate any foreign material from the body. Thus, the environment around the sensor might change completely and the measurement result may not be relevant at all. Biological systems can also affect the performance of the sensor by degrading its structure or damaging the biosensitive-layer due to the presence of corrosive ions and enzymes. This can, of course, lead to mis-calibration and failure of the measurements. It is especially important to overcome these problems in case of *in vivo* sensors. Once the device is implanted, the access to it is very limited. So this kind of sensors should be highly stable and maintain their calibration for long time. In this thesis biosensors for *in vitro* diagnostics are developed. High stability and reproducibility are also required for *in vitro* biosensors. But they can be fabricated in such a way that they are disposable or partially reusable. Biosensors are used not only in medical diagnostics but also in environmental monitoring, research laboratories, process industries, food and drug monitoring, bio-defense applications, etc [14, 15].

**1.3. Nanomaterials for the detection of biomolecules.** As it was mentioned before (*paragraph 1.2*), the sensitivity of the biosensors depends a lot on the characteristics of transducer (shape, type of the material, size). High surface-to-volume ratio of the transducing element can increase the efficiency of the signal transfer. Thus, nanomaterials are ideal candidates to be used as components of transducers. The first and foremost feature that nanostructures offer is their size (a few Angstroms to 100 nm), which is in the range of the size of various biomolecules such as nucleic acids, small proteins and viruses. The small size brings a tremendous increase in the surface-to-volume ratio that is essential for maximizing sensitivity. A major advantage of using nanostructures in a label-free sensing is that the amount of receptors immobilized on the detector surface can be as low as a single molecule [16 - 19]. As a result the amount of analyte required to generate a measurable signal could be just a few providing for very low

limits of detection (LOD). The lower the LOD the earlier a biomarker characteristic of a disease state can be identified [20].

Secondly nanostructures exhibit specialized physical and chemical properties that are generally not available in the bulk. For example, the binding of analytes to receptors immobilized on carbon nanotubes brings in significant changes in the resistance of the CNT that is used as a sensor signal [21]. Such a phenomenon does not occur in bulk metals. Furthermore, surface plasmons in thin metal films (2D) and nanoparticles (0D) [22] are characteristic properties at the nanoscale that are otherwise not available in bulk microscale materials. These unusual properties arise due to confinement in one to three dimensions. Another related aspect is the improvement in robustness of a certain physical property, for example, fluorescence of quantum dots due to their 0D nanostructure [23, 24]. While nanostructures exhibit favorable properties in a standalone manner, they also help augment transduction characteristics of bulk materials. A typical example is the improvement in electrochemical detection achievable by nanostructuring of the electrode surface [25-29] due to increase in surface area [30].

A key aspect for the success of nanostructures is the tunable fabrication of the materials or the ability to tailor the chemical and physical properties. For example, the emission wavelength of quantum dots can be tuned by just varying their size [23, 24]. Through various chemical and bio functionalization protocols, the surface of the nanostructures can be easily optimized for the detection of desired analytes. Other advantages of using nanostructures include the ability to miniaturize the diagnostic tool, increase the speed of detection and reduce reagent and sample consumption [17, 19]. Biochips or arrays of biosensors on one substrate can be built using nanobiosensors. Using such devices, measurements of different analytes in the same drop of liquid can be performed simultaneously. Delivery and control of liquids in  $\mu$ TAS can be achieved using a microfluidic platform. It gives a possibility to decrease the volumes of reagent solutions, avoid evaporation, shorten reaction time and perform parallel operations [31 - 33].

**1.4. Electrical/electrochemical sensing.** Currently most of the diagnostic assays are based on optical measurements. Utilization of nanomaterials (e.g. quantum dots and nanoparticles) can significantly improve limits of detection of such techniques. However, complicated readout instrumentation, long duration for sample preprocessing and the need for the labeling make optical methods expensive, time-consuming and non-portable. In contrast, electrical

detection methods rely on much simpler instrumentation that ensures lower cost and power consumption. Electrical methods are ideally suitable for implementation of label-free detection approaches, which give a number of the advantages for the biomedical assays, such as elimination of the need for modification of the molecule of interest, avoidance of the influence of the label on the binding properties, possibility of real-time monitoring of binding interactions, etc. Using nanomaterials and modern microfabrication techniques gives the possibility of miniaturization and multiplex sensing. These make electrical methods more promising for applications in the point-of-care [30, 34, 35].

Electrochemical methods of detection are based on electrochemical processes that take place on the electrode surface. There are different modes of electrochemical detection. Voltametric and amperometric assays are based on the measurement of current at the working electrode resulting from the application of a voltage at the electrode-solution interface. In case of voltammetry the current is measured as a function of changing potential that can be applied in different ways (linear, cyclic, anodic stripping voltammetry). Amperometric measurements are performed by maintaining a constant potential at the working electrode with respect to a reference electrode and measuring the generated current. In potentiometric detection the potential between measuring and reference electrodes gives an idea about the concentration of certain ions in the solution. Current flowing through the electrode is equal to or near zero [36, 37].

Another class of electrical methods used for the detection of analytes is based on the utilization of transistors. In contrast to electrochemical detection no oxidation or reduction process is required. Therefore, even without the use of an electroactive mediator the range of possible analyte molecules can be broadened. Field-effect transistors (FET) are the most common among such biosensors. An FET contains source and drain electrodes, a semiconducting channel and a gate electrode [38, 39]. Electrical transport through the semiconductor channel is modulated by an applied gate voltage ( $V_G$ ). The gate regulates the electrical field generated perpendicular to the surface of the channel. This field influences the amount of the mobile and trapped carriers in the channel and therefore the electrical conductance of the semiconductor. This electrical field can be generated, for example, in a solid state dielectric, like silicon oxide (back gate - Figure 1.1a), in a polymer, like polyethylene oxide (polymer electrolyte gate) or in a liquid (liquid gate - Figure 1.1b).



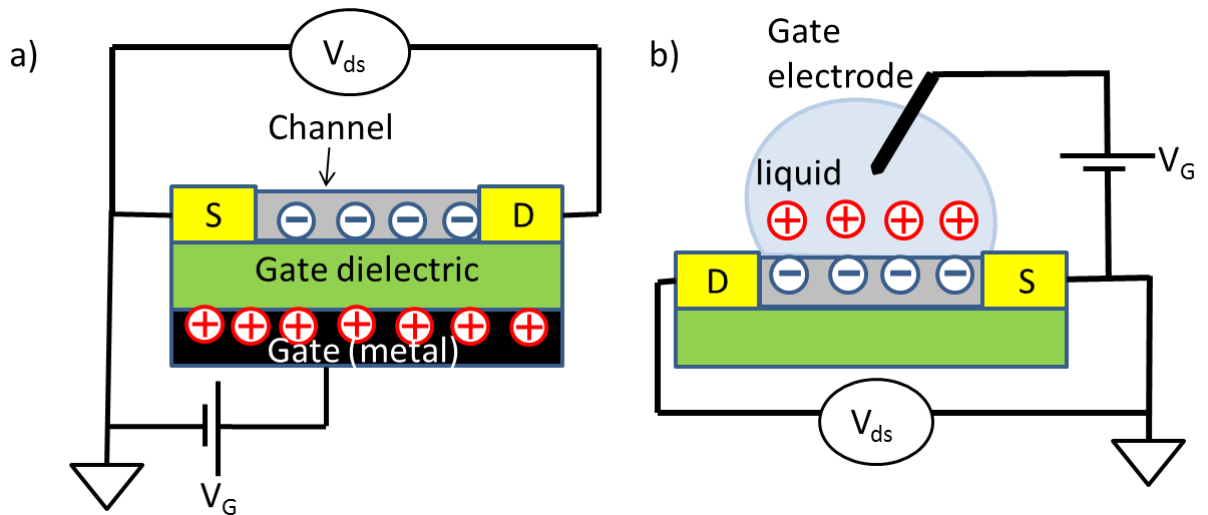


Figure 1.1. Schematic of a back-gated (a) and a liquid gated FET (b).

Liquid (electrochemical) gated configuration is more promising for medical diagnostics since it enables analyte detection directly in biological liquids. In this case the whole device is immersed into liquid and the reference electrode is used to apply gate voltage (Figure 1.1b). The basis for the electrochemically gated device is the measurement of the conductance of the system as a function of the charge-carrier concentration, controlled by the electrochemical potential. The gating effect is achieved through the formation of an electrical double layer (EDL) on the surface of the semiconductor. A change of the potential of the gate electrode leads to a change of the electrostatic potential drop over the EDL. Charges that form the EDL influence the interfacial potential and thereby the source-drain current [40]. The adsorption of additional charges due to biorecognition reactions on the surface of semiconductor will lead to changes in transport characteristics of the device. These changes can be detected by measuring source-drain current [41].

The method of detecting the changes in the transport characteristics of the FET is another important aspect in electrical biosensors. It is known, that impedance measurements compared to DC offer better signal to noise ratio as well as the possibility to change one more input parameter - frequency. *Impedance* ( $Z$ ) - is the equivalent of resistance for an AC circuit [42, 43]. It describes not only the voltage to current ratio but also the phase difference between these two parameters. The Ohm's law for AC circuits is given by:

$$V = I|Z|e^{j\varphi}$$

The magnitude of impedance is the resistance  $|Z|$  - is a drop of voltage amplitude  $V$  for a given current  $I$ . The phase factor  $e^{j\varphi}$  describes the delay of the

voltage with respect to the current by a phase of  $\varphi$  (Figure 1.2.),  $j = \sqrt{-1}$ . The phase shift is attributed to the capacitive or inductive effects that may occur in the system. At frequencies below 1 MHz impedance measurements give an idea about the resistive and capacitive behavior of the device. When the phase is  $0^\circ$  the device acts as a resistor ( $Z = R$ ). When phase is  $90^\circ$  device acts as a capacitor:  $Z_C = \frac{1}{j\omega C}$ , where  $\omega$  - is angular frequency.

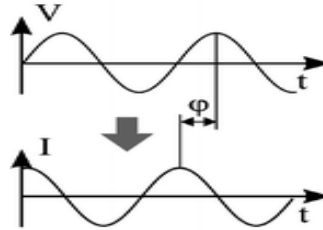


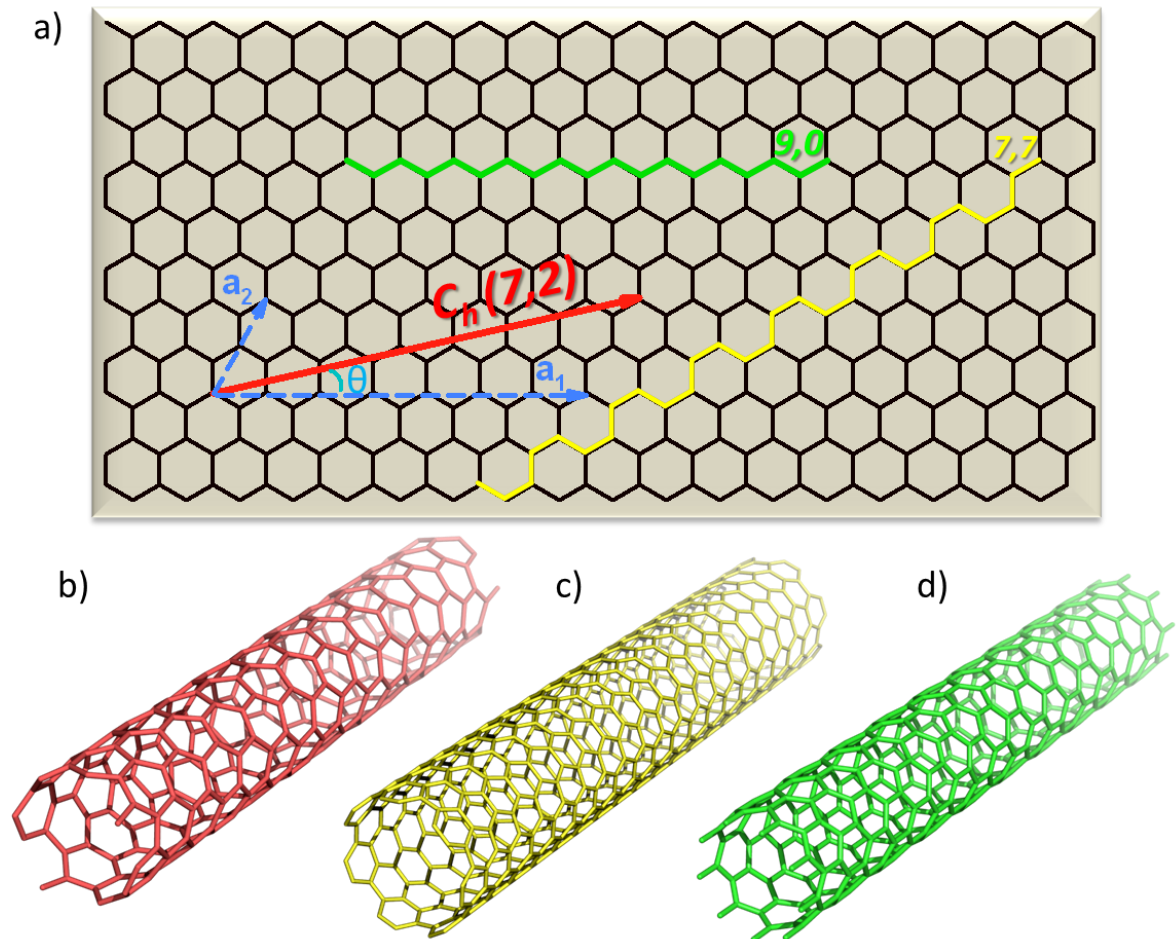
Figure 1.2. Representation of the phase of impedance [44]

Measurement of impedance at different AC frequencies is called impedance spectroscopy. When impedance measurements are taken in a FET device configuration it is possible to obtain complete electrical characteristics of the system - both the resistive and capacitive effects at different gate voltages. This kind of measurements can be crucial for understanding the mechanism of sensing and for identifying the best set of parameters of the measurements for obtaining the highest sensitivity and selectivity.

## 2. Carbon nanostructures for biosensing

**2.1. Introduction.** The electronic configuration of carbon in ground state is  $1s^2 2s^2 2p^2$ . Carbon atoms can exist in three states corresponding to  $sp^3$ -,  $sp^2$ -, and  $sp$ - hybridization of their valence orbitals. Based on the type of hybridization different allotropic forms of carbon can be distinguished [45]. Among them graphitic and diamond forms deserve particular attention. Graphite has a layered structure. Each layer of graphite is a hexagonal lattice of  $sp^2$  hybridized carbon. Graphite shows metallic behavior in plane and poor electrical conductivity along perpendicular axis. Diamond is a wide-band gap semiconductor. It has tetrahedral  $sp^3$  bonding and shows nearly isotropic properties. Graphite is the stiffest material in nature and diamond is the hardest one. In the field of nanotechnology carbon is very important due to existence of 0D (fullerenes, nanodiamonds), 1D (carbon nanotubes) and 2D (graphene) nanomaterials which have unique physical and chemical properties and are very promising in a number of applications. Carbon-derived nanomaterials can be used at wide range of temperatures [46]. This thesis concentrates mostly on Single Wall Carbon Nanotubes (SWCNTs) and graphene. The atomic structure of both materials is essentially graphitic.

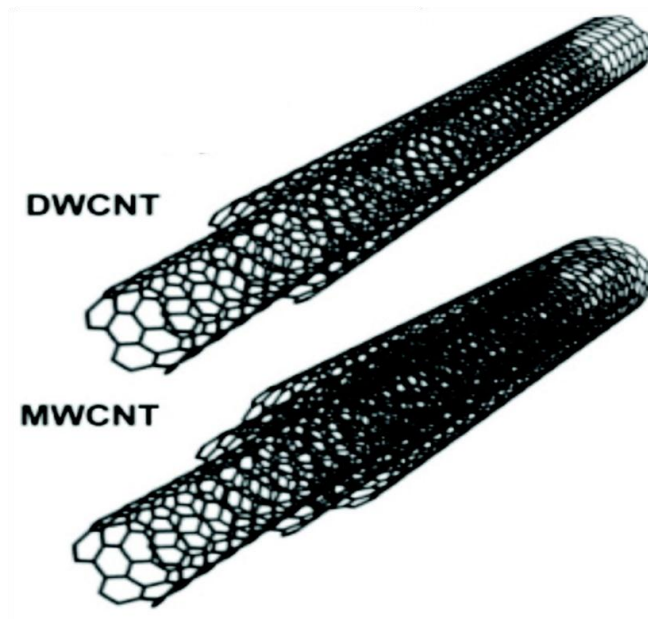
**2.2. Carbon nanotubes.** Carbon nanotubes were discovered by Iijima in 1991 [47]. Each SWCNT (single walled carbon nanotube) can be viewed as a cylinder that has a single sheet of graphite (graphene) as its wall. The way the graphene sheet is rolled up into a tube defines the electrical and optical properties of SWCNT. To describe the geometry of SWCNT one can use lattice vectors  $a_1$  and  $a_2$  and indices  $n, m$  (Figure 2.1 a). The lattice vector  $C_h$  is defined as  $C_h = na_1 + ma_2 = (n, m)$ .  $0 \leq |m| \leq n$ . The length of the chiral vector  $C_h$  is directly related to the diameter of the nanotube. The chiral angle  $\theta$  between  $C_h$  and zigzag direction of the lattice  $(n, 0)$  is related to indices  $n, m$  [48].



**Figure 2.1.** Physical structure of CNTs. Chiral vector  $C_h$  is defined by unit vectors  $a_1$  and  $a_2$  and the angle  $\theta$ . Yellow and green lines represent armchair and zigzag configurations respectively. b) (7,2) - chiral SWCNT, c) (7,7) - armchair configuration d) (9,0) - zigzag configuration. The CNT structures were simulated using CNTbands2.7

CNTs with  $n=m$  ( $\theta = 30^\circ$ ) are called armchair nanotubes due to the characteristic shape of their cross-section (Figure 2.1). CNTs with  $m=0$  ( $\theta=0$ ) are called zigzag nanotubes. The remaining configurations of CNTs ( $n \neq m$ ,  $0 < \theta < 30^\circ$ ) are called chiral. If  $n=m$  or  $n-m=3i$  ( $i$  - integer) the CNT is metallic. If  $n-m=3i \pm 1$  then the CNT is a semiconductor.

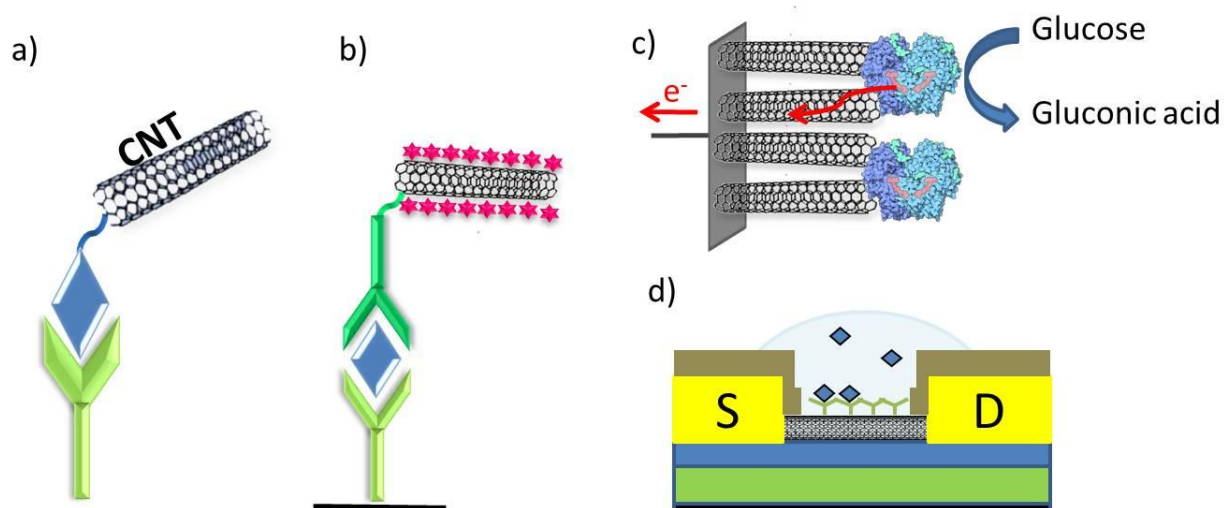
It is also possible to obtain double-wall and multi-wall CNTs. In this case, the CNTs of different chiralities can form two or more walls of the nanotube (Figure 2.2). The variety of this kind of nanotubes is enormous due to the large amount of possible combinations of the nanotubes.



**Figure 2.2.** The structure of double walled carbon nanotube (DWCNT) and multi-walled carbon nanotube (MWCNT) [49]

Carbon nanotubes can be produced using chemical vapor deposition, laser ablation and arc discharge [50]. The resulting product contains a mixture of CNTs, catalyst particles and amorphous carbon. Moreover the nanotubes in the mixture are not identical. They have different lengths and chiralities and thus different properties. It is still challenging to obtain long CNTs of a specific structure. This is the reason why the large-scale manufacturing of identical CNT devices has been difficult. Nevertheless, the unique properties of CNTs as a 1D nanomaterial make them advantageous even though it is not yet possible to separate efficiently enough different types of nanotubes. For example, this thesis stipulates the idea of using SWCNTs as a channel in FET. As it was previously discussed (*paragraph 1.4*), an FET requires a semiconducting material to be used as a channel. However, it is possible to achieve gating of electrochemically modified metallic nanotubes [51]. In the current work the devices that contained a mixture of metallic and semiconducting nanotubes showed sufficient gate-dependence to be used as FET-based sensors.

Chemically, SWCNTs are not very reactive due to their highly graphitized nature. Oxidation was first shown at high temperatures (above 750°C) in the gas phase resulting in the formation of three functional groups: carboxylic (-COOH), carbonyl (-CO) and hydroxyl (-COH) with the ratio 4 to 2 to 1 respectively [52]. Oxidation significantly increases the reactivity of nanotubes and also modifies their wetting properties. Young's module of carbon nanotubes is 2 terapascal [53]. Mechanically, SWCNTs are very flexible and elastic [54, 55].



**Figure 2.3.** Overview of different CNT-based biosensing strategies. a) CNT is used as a label b) CNT is used as a support for loading tags, c) electrochemical CNT sensor, d) CNT-FET sensor [30].

The chemical and physical properties of CNTs make them well suited material for sensing applications. Numerous results have been reported on the utilization of CNTs in pressure, flow, thermal, gas, optical, mass, stress, strain, chemical and biological sensors [56]. With CNT-based biosensors both labeled and label-free detection schemes have been demonstrated. Carbon nanotubes have favorable optical properties such as a characteristic Raman signal and can be used as labels (Figure 2.3 a) [57]. Moreover, they act as a support to carry a payload of labels (Figure 2.3 b) [58]. They can also be used as supports functioning as label-free electrical detectors in heterogeneous assays [59, 60]. In electrochemical sensors excellent transport properties of CNTs and the possibility to increase the sensitive surface of the electrode are utilized. In this case nanotube forests on platinum electrodes have been used for immobilization of enzymes that catalyze certain electrochemical reactions. The changes of electrical current due to the electrochemical reaction are efficiently transferred through the nanotubes to the metal surface and can be easily detected using voltammetry or amperometry (Figure 2.3 c) [61]. In CNT-FET based sensors individual nanotubes or their networks can be used as a channel providing high-sensitivity for this kind of measurements (Figure 2.3 d). FETs based on networks of nanotubes offer better reproducibility and manufacturability but they show lower sensitivity compared to FETs based on single CNT [60]. The first FET chemical sensors utilizing CNTs were demonstrated for the detection of gases [62, 63]. Later CNT-FET platforms were applied for the detection of chemical species in liquids [64, 65]. Using both catalytic and affinity based CNT-FETs the sensing of biological molecules, such as metabolites, proteins and nucleic acid, was

demonstrated [40, 66, 67]. The reported sensors have good sensitivity however in most of the cases their fabrication cannot be upscaled. Reproducibility and reliability of reported CNT based sensors have been their major drawbacks. In this thesis a scalable route to fabricate CNT-based sensors is presented. The fabrication is performed in a way to ensure robustness, high sensitivity and selectivity of the devices. Using glucose as an analyte for affinity-based saccharides sensing the mechanism of sensing using designed platform has been investigated. Following this, high sensitivity of the developed biosensor was demonstrated for the detection of nucleic acids.

**2.3. Graphene.** Graphene - is a single layer of graphite (Figure 2.4). Due to its atomically thin structure it can be considered as an ideal 2D-material. In recent years graphene has attracted massive attention of the researchers in science and industry due to its very interesting physical properties. Ambipolar field effect along with ballistic conduction of charge carriers [68] makes it extremely attractive for electronics applications. Graphene has been used for the development of various nanodevices, such as field-effect transistors, capacitors, mass, optical and electrical sensors, etc [69-72]. For the field of sensing graphene is very attractive due to its very high surface to volume ratio. Estimated specific surface area of single layer graphene is  $2630 \text{ m}^2/\text{g}$  (SWCNTs or graphene on substrate -  $1315 \text{ m}^2/\text{g}$ ) [73].

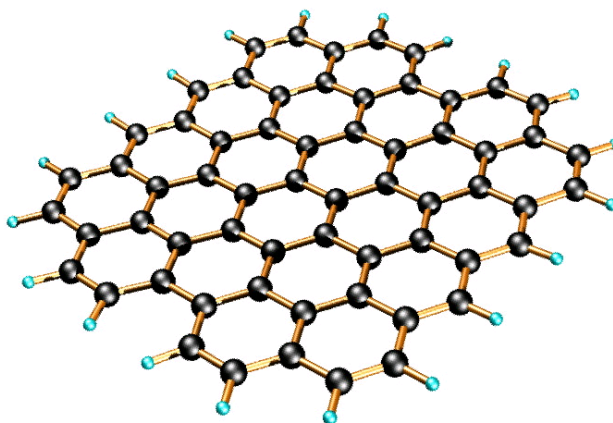


Figure 2.4. The structure of graphene.

For optical bioassays graphene oxide is very attractive due to its optical properties, solubility in water and the presence of functional groups for binding biomolecules. Fluorescence resonance energy transfer (FRET) induces quenching of



the fluorescence of fluorophore by graphene oxide. Graphene oxide shows very good quenching ability to a variety of organic dyes and quantum dots. FRET - based assays utilizing graphene oxide have been demonstrated for the detection of DNA and proteins both in the solution and on the surface. [74, 75, 76]

Application of graphene in electrical biosensors has been demonstrated as well. Graphene and its derivatives or composites have been used for the modification of electrodes in electrochemical sensors. Electrochemical detection of glucose, dopamine, proteins, etc has been demonstrated using graphene-modified electrodes [77-81]. Some results were reported on the utilization of graphene-FET for sensing. As in the case of CNTs, first graphene-FET sensors have been demonstrated for the detection of gases [82, 83] and determination of pH [84]. Later graphene-FETs for the detection of cells and proteins were demonstrated [85-87]. Graphene-FET sensors are expected to take an advantage of better reproducibility compared to CNT-FETs that suffer from inhomogeneity of the carbon nanotubes [88]. However, the absence of the reliable technique for the large-scale fabrication of identical graphene devices is an important factor that is currently limiting the development of this field.

**2.4. Functionalization of carbon nanostructures.** Most of the possible applications of carbon nanomaterials (like sensors, microelectronics, optoelectronics, molecular filters and other) require the modification their surface in order to improve stability, biocompatibility, electrical or optical properties, solubility or selectivity of the nanostructures [89, 90]. The process of surface modification in order to obtain certain functionality is called chemical functionalization. In the case of biosensing applications, functionalization of CNTs and graphene is important for rendering them selective to specific analyte molecules. This is usually achieved by attaching the receptor molecules to their surface. The receptors can be attached either covalently or non-covalently. Functionalization of carbon nanostructures can play another important role in biosensors. Specifically, surface modification is often used in order to reduce non-specific binding of the various biomolecules to the surface of the transducing component [91]. In this work, for example, polyaminobenzoic acid non-covalently attached to CNTs was used for subsequent immobilization of the receptor DNA as well as for the protection of the CNT surface against non-specific binding of the oligonucleotides.

Due to the high importance of CNT and graphene functionalization for biosensing applications, possible modification routes deserve special attention here. From the chemical point of view the surface of CNTs and graphene is essentially the same. Therefore, similar strategies can be used for their modifications. However, the



chemical reactions of pristine graphene are much less explored [92, 93]. It is known, that the degree of functionalization of different carbon nanomaterials and their different areas varies significantly. For example, the edges and defect sites of CNTs and graphene are generally more reactive [90, 93, 94]. Nanotubes with smaller diameter are chemically more reactive due to their increased curvature [95]. In the graphene lattice the most reactive sites are the geometrically strained regions [96].

**2.4.1. Non-covalent functionalization.** Highly hydrophobic surfaces of graphene and CNTs adsorb molecules with aromatic residues by  $\pi$ - $\pi$  stacking. Therefore, some receptor molecules, like nucleic acids or hydrophobic proteins, can be easily attached to carbon nanostructures [96 - 98]. The same molecular interactions occur when pyrene-like molecules or other aromatic compounds are used as anchors for the receptor attachment (Figure 2.5) [93, 99]. Often amphiphilic molecules are used to prepare and stabilize the aqueous dispersions of CNTs or graphene [100, 101]. As an example, in this work the CNTs were dispersed in Triton X-100 solution. Electrochemical polymerization can be also utilized for the non-covalent wrapping of the nanostructure surface by the polymer of interest. This strategy was used in this work and therefore will be discussed in detail later.

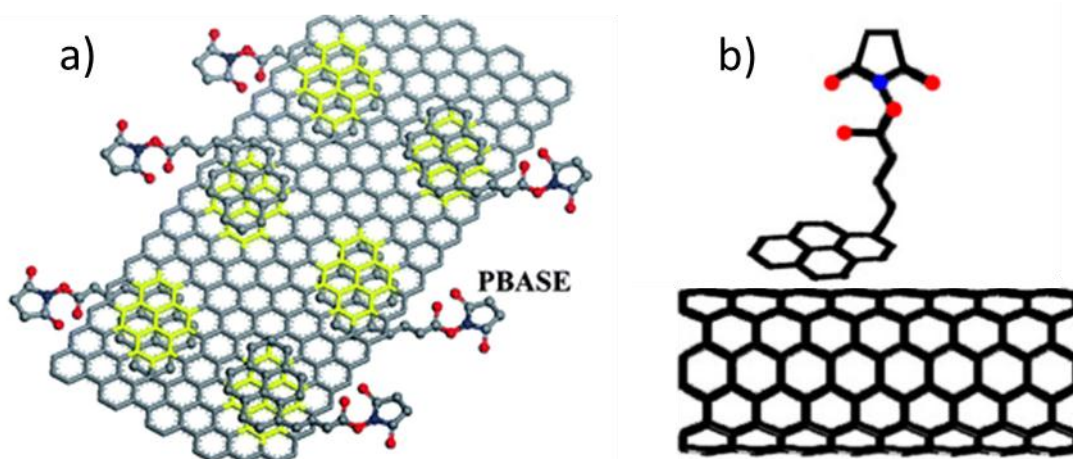
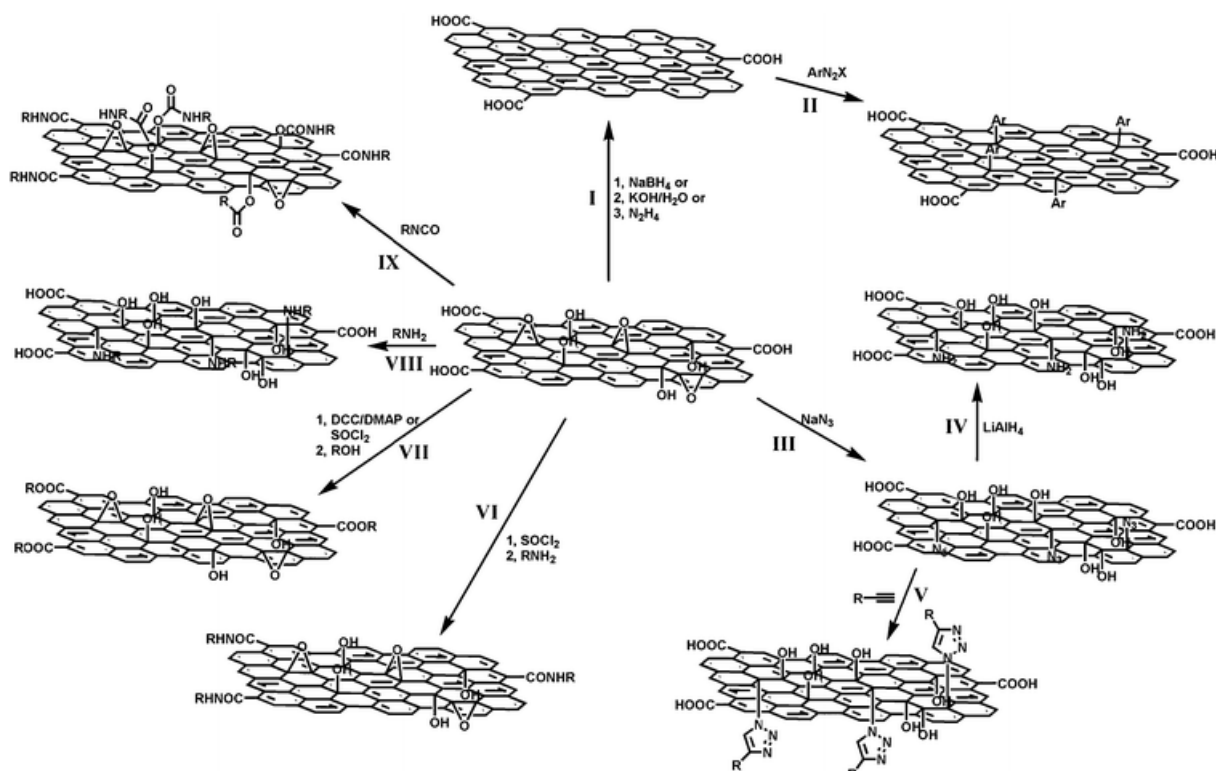


Figure 2.5. Non-covalent functionalization of carbon nanostructures with pyrene-derivatives via  $\pi$ - $\pi$  stacking. 1-pyrenebutanoic acid succinimidyl ester (PBASE) attached to a) graphene and b) CNT [93, 99].

**2.4.2 Covalent functionalization.** A wide range of reactions, including thermally activated, photochemical and electrochemical, can be used for covalent modification of carbon nanostructures. One of the examples of *thermally activated reactions* is the oxidation of CNTs or graphene. It results in the formation of the reactive carboxyl, epoxide or hydroxyl groups [52, 102]. With the help of various chemical reactions these oxygen-containing groups can act as sites for the subsequent attachment of the receptor. For example, molecule of interest can be coupled to

carboxyl groups via esterification or amidation reactions [93, 102, 103]. Different reactions of the oxygen-containing groups in oxidized graphene are shown in Figure 2.6.



**Figure 2.6. Various covalent functionalization schemes for graphene oxide. I: Reduction of GO II: Covalent surface functionalization of reduced graphene via diazonium reaction. III: Functionalization of GO by the reaction between GO and sodium azide. IV: Reduction azide-GO with  $\text{LiAlH}_4$  resulting in the amino-functionalized GO. V: Functionalization of azide-GO through click chemistry. VI: Modification of GO with long alkyl chains by the acylation reaction between the carboxyl acid groups of GO and alkylamine. VII: Esterification of GO by DCC chemistry or the acylation reaction between the carboxyl acid groups of GO and ROH alkylamine. VIII: Nucleophilic ring-opening reaction between the epoxy groups of GO and the amine groups of an amine-terminated organic molecules ( $\text{RNH}_2$ ). IX: The treatment of GO with organic isocyanates leading to the derivatization of both the edge carboxyl and surface hydroxyl functional groups via formation of amides or carbamate esters ( $\text{RNCO}$ ) [93].**

Various thermally activated addition reactions for graphene and the sidewalls of CNTs are available. Figure 2.7 collects different routes for the covalent modification of carbon surface using addition reactions [102].

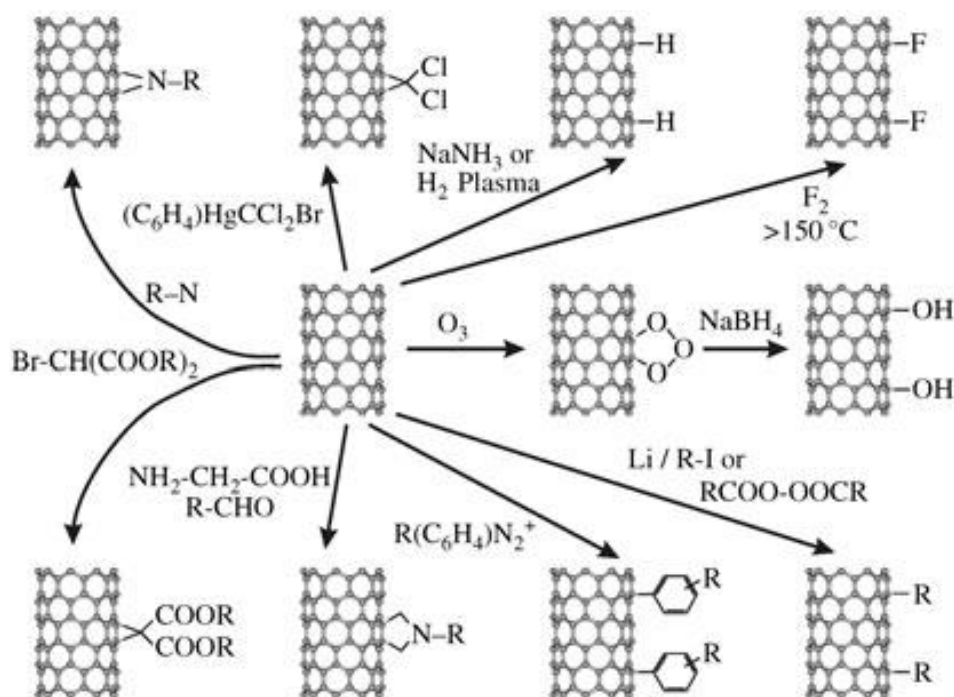


Figure 2.7. Addition reactions for the functionalization of the carbon nanotube sidewall [102].

As in the case of oxidized carbon nanostructures the products of the addition reactions are often used for subsequent attachment of the receptor molecule. For instance, fluorine in fluorinated CNTs can be replaced by nucleophilic substitution using alcohols, amines, Grignard reagents or alkyl lithium compounds [102].

*Photochemical activation* was also implemented for modification of CNTs and graphene. For instance, CNTs have been osmylated using osmium tetroxide ( $OsO_4$ ) under UV light irradiation [104]. Similarly, under intense UV-irradiation graphene can react with benzoyl peroxide [105].

*Electrochemical functionalization* provides the attachment of biomolecules on to carbon surfaces in a very versatile manner in a covalent or non-covalent fashion [106, 107]. For performing electrochemical reactions an electrochemical cell is used. A typical electrochemical cell consists of a working electrode (WE), a reference electrode (RE) and a counter electrode (CE). The working electrode provides the surface for electrochemical reactions. When a positive overpotential is applied, oxidation of an electrochemically active component in the solution can be achieved. Negative overpotential may induce reduction processes. RE is used to set the reference potential and CE is important for monitoring the electrical current generated in the cell during the electrochemical process. Both covalent and non-covalent modification can be realized, for instance, by using receptor molecules

containing aminophenyl groups (Figure 2.8) [108]. If CNT is used as a working electrode, application of positive voltages leads to polymerization of the molecule on the CNT surface forming a uniform functional layer. In order to functionalize nanotubes covalently aminophenyl moieties should be first converted into its diazonium salt which is very reactive. Electrochemical reduction causes the formation of aryl radical which can be coupled to the nanotube wall.

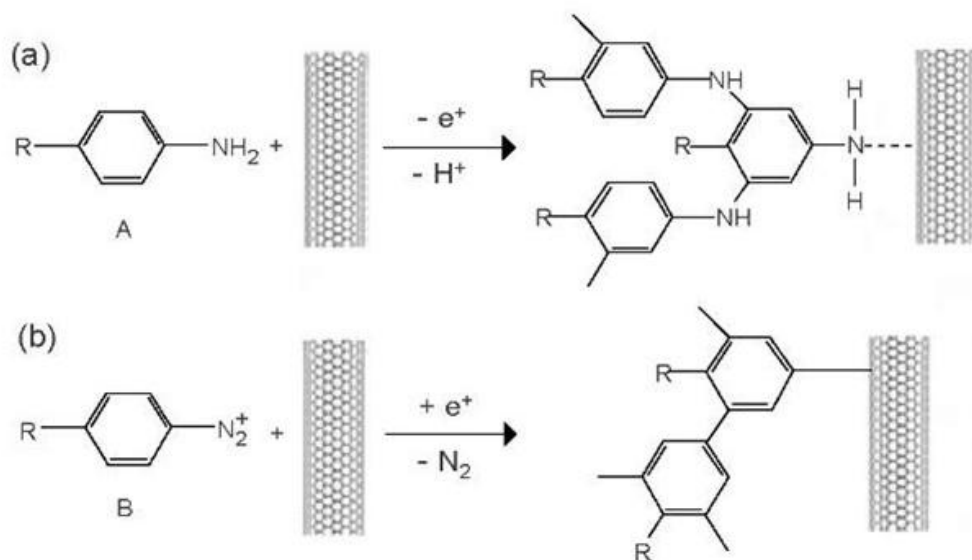


Figure 2.8. Schemes for (a) oxidative ECM with 4-aminobenzyl-R and (b) reductive ECM with an aryl diazonium salt. In the former case, the modified SWNT is covered by a polymeric layer of aminobenzyl groups without the formation of a chemical bond, whereas in the latter case, polyphenyl groups covalently attached to the nanotube are formed [108].

Electrochemical modification (ECM) is a very fast and efficient method for the modification of carbon nanostructures. It can be easily upscaled and automated. ECM ensures the coupling of the receptor selectively on the surface of transducing component. This is essential for achieving low limit of detection. Due to these reasons ECM was chosen as a strategy for the functionalization of CNTs in this work.

# Experimental section

---

## 3. Fabrication of CNT-based sensors

**3.1. Introduction.** CNTs are shown to be promising for various fields of use. A lot of possible applications including electrical sensing require integration of CNTs into an electronic circuit. However, the fabrication of CNT devices at the industrial level is still challenging [49, 109]. One of the most common ways to integrate CNTs into a circuit is the writing of the metal contacts on randomly deposited CNTs using e-beam lithography. However, this method is time-consuming and requires manual operation for fabricating individual devices. CNTs can be grown directly from catalyst particles deposited on the pre-patterned electrodes [110]. This approach requires high temperatures and positioning of the catalysts at the exact point. Another drawback of this method is the non-uniform growth of the CNTs from each catalyst. Therefore up-scaling of mentioned technique seems very difficult.

It is also very important to define the exact location of the final device on the substrate from the beginning of fabrication process. This makes it possible to assemble other parts of the sensor in a modular way. We used the dielectrophoresis (DEP) technique to position the nanotubes at the desired location, for example, in between platinum electrodes patterned on a substrate. This method offers considerable advantages: it is fast, does not require expensive and bulky equipment and can be automated and upscaled [111, 112]. To perform dielectrophoretic trapping it is necessary to solubilize CNTs. After the DEP trapping and visualization, the electrodes are passivated with an insulating layer and subsequently the nanotubes are modified with receptors specific to certain analytes. The details of fabrication are outlined in this chapter.

**3.2. Preparation of CNT dispersions.** Hydrophobic and strong van der Waals interactions between the carbon nanotubes make it difficult to

suspend them in water or other polar solutions. Some organic solvents (N,N-dimethylformamide (DMF), dichloroethane), surfactants (sodium dodecyl sulfate (SDS), sodium dodecyl benzene sulfonate (SDBS), etc) and polymers (nafion, DNA) were shown to be efficient for nanotube solubilization [113-115].

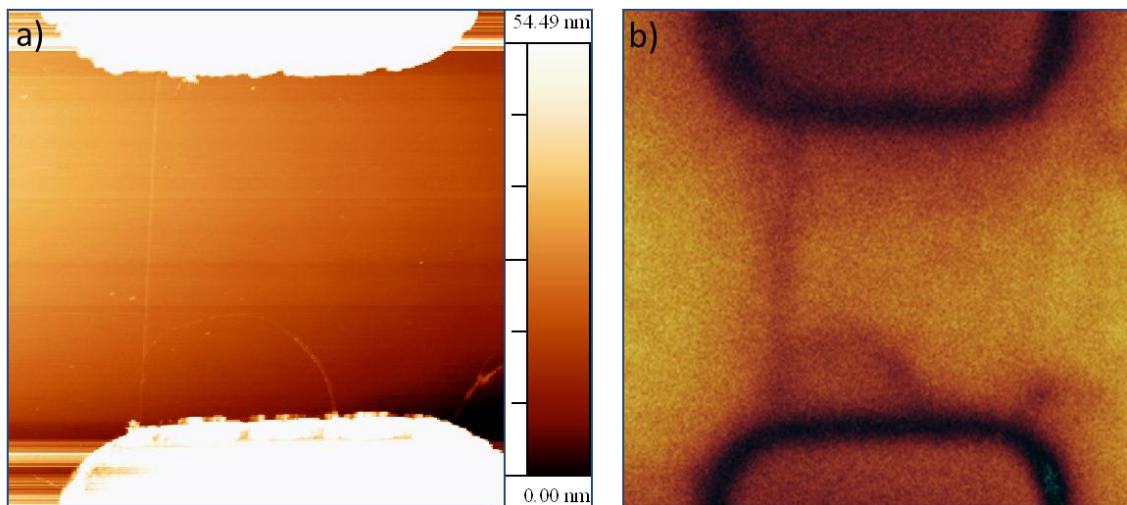
Purification is another important aspect during the preparation of CNT dispersion. SWCNTs powder contains a number of impurities (amorphous carbon, catalysts, nanoparticles, etc) that might affect the quality of devices [116]. Gas phase oxidation is often used to improve the homogeneity of CNTs [117]. To purify the CNT solution standard purification techniques, like centrifugation, filtering, etc can be used [116, 118]. Density gradient ultracentrifugation and gel-chromatography are shown to be efficient techniques to separate SWCNTs with different chiralities from their dispersions in surfactant. The solutions prepared using these approaches are enriched with the nanotubes of only a certain physical structure. However the length of the nanotubes in the resulting suspensions is less than 1  $\mu\text{m}$  [119-122]. Therefore these solutions are not very suitable starting material for the fabrication of devices based on individual nanotubes.

We used 0.1% Triton X-100 aqueous solution for dispersing of the SWCNTs. A small amount of CNT powder (HipCO oxidized) was added to surfactant solution and dispersed with the help of ultrasonic agitation (Tip sonicator HD-3100) for 40 sec with 1 sec impulse and 2 sec rest intervals. Ultrasonic cavitation helps to overcome the bonding forces between the nanotubes, and separate them. After that the SWCNTs dispersion was centrifuged (Rotanta 460 RS, Hettich) at 4637 g (4600 rpm) for 30 min, and the supernatant was filtered using filter paper [123]. Hexane extraction was used for further purification of the solution. This approach allows the removal of hydrophobic impurities and undispersed CNTs from the solution. Well-dispersed nanotubes due to the surfactant covering their surface remain in aqueous phase. For the extraction equal volume of hexane was added to CNT solution, mixed well and centrifuged for 5 min at 4600 pm. Upper hexane fraction containing impurities is carefully separated from the clear CNT solution. The CNT suspension prepared in this manner is not stable for long time, therefore some processing steps, like short sonication and centrifugation, of older solution (more than 1 week) are required before usage.

**3.3. Visualization of CNT devices.** Visualization of the carbon nanotube devices is important for the optimisation of fabrication steps and sensing conditions. Atomic force microscopy (AFM) is the most commonly used technique for visualization of CNT devices. It gives an idea about the geometry of the

electrodes and nanostructures, diameter and number of the trapped nanotubes or bundles and the cleanliness of the surface. For the same purpose electron microscopy can also be used. However imaging of SiO<sub>2</sub> surface with scanning electron microscope (SEM) requires its coating with a conducting layer to prevent accumulation of electrostatic charge during electron irradiation. Therefore the devices cannot be used after imaging. The resolution of optical microscope does not allow the imaging of single CNTs or small bundles.

We designed a new method for the CNT visualization using fluorescent dyes. CNTs have the ability to quench fluorescence of fluorescent dyes [124,125]. By covering the sample with a thin layer of Rhodamin B (RB) it was possible to image the CNTs on the SiO<sub>2</sub> surface using a confocal microscope (TCS SP2, Leica). It is apparent that the thin nanotube bundle (Figure 3.1.) can be clearly visualized due to the efficient quenching of the dye fluorescence by complex formation with the nanotubes [105]. Since confocal microscope imaging is much faster than AFM, this approach can be used for a quick characterization of the sample. However it is not possible to obtain information about the CNT diameter as it can be done using AFM. Therefore all the devices used for this work were characterized using AFM - (Digital Instruments Dimension IV, Veeco).



**Figure 3.1. Visualization of the same SWCNT device with AFM and confocal microscopy: AFM (a) and confocal fluorescence (b) images of the same sample showing a tube trapped across electrodes. The confocal image was recorded after leaving the sample in a solution of RB in ethanol. RB complexes to the nanotubes and hence the fluorescence is quenched along the tubes. This helps in the clear visualization of the tubes on a substrate [123].**

### 3.4. Dielectrophoretic manipulation of CNTs.

Dielectrophoresis enables the positioning of CNTs at the desired location. It can also be used for parallel assembly of 1D-nanostructures for fabrication of an array of devices over a large wafer. By varying parameters of trapping it is possible to control the density of nanotubes on the substrate surface. To understand what the variable dielectrophoresis parameters are, it is important to know the nature of dielectrophoretic force and factors that influence it.

**3.4.1. The theory of DEP.** Dielectrophoresis is an electrokinetic motion of dielectrically polarized materials in non-uniform electric fields [126]. The polarized object can move toward or away from the high field region depending on the complex dielectric permittivity of the object with respect to its surrounding medium. The CNT assembly process is conceptually illustrated in Figure 3.2. When nanotube in the solution is located within the AC electric field, it polarizes. The dielectric constant of the nanotube is larger than dielectric constant of the solvent. This gives an origin to a positive DEP force. The CNT at kHz to MHz frequencies gets attracted to the place where field is the strongest – to the tip of the electrode. The CNT assembly process depends on the relative balance of the dielectrophoretic, hydrodynamic and electrostatic double-layer interactions between the nanotube and surface [127].

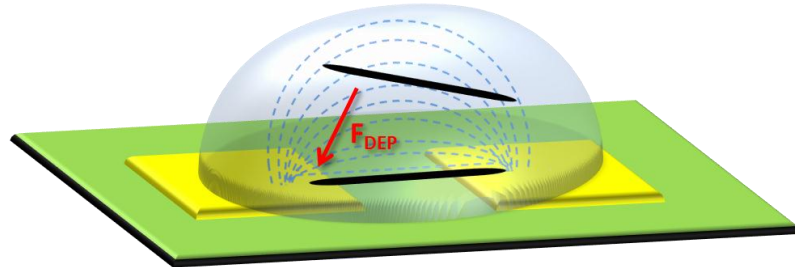


Figure 3.2. Schematic of the dielectrophoretic manipulation of 1D-nanostructures.

The dielectrophoretic force on a carbon nanotube assuming it to be a cylindrical particle can be described by the following equation:

$$F_{DEP} = \frac{\pi}{2} r^2 l \epsilon_m \text{Re}(f_{CM}) \nabla |E|^2$$

Where  $\epsilon_m$  – relative permittivity of medium,  $r, l$  – radius and length of the nanotube respectively,  $E$  – the local electric field,  $\text{Re}(f_{CM})$  – real part the dipolar



Clausius-Mossotti factor. For an elongated object with the long axis aligned with the field  $f_{CM}$  is given by:

$$f_{CM} = \left[ \frac{\varepsilon_p^* - \varepsilon_m^*}{\varepsilon_m^*} \right]$$

where  $\varepsilon_p^*$  and  $\varepsilon_m^*$  are the complex permittivity of the CNT and surrounding medium, respectively [128]. The complex permittivity is defined as  $\varepsilon^* = \varepsilon - j(\sigma/\omega)$ , where  $\varepsilon$  is the permittivity,  $\sigma$  is the conductivity, and  $\omega$  is the angular frequency of the applied electric field,  $j = \sqrt{-1}$ .

In agreement with the abovementioned equations for a certain type of particle in a given medium one can tune the dielectrophoretic force by varying the amplitude and the frequency of the applied electric field. The amount of nanotubes trapped at the electrodes will depend also on the quality and the concentration of solution, the design of the electrodes and the deposition time.

**3.4.2. CNT trapping results.** In our case, the aim was to trap carbon nanotubes between platinum electrodes on silicon/silicon oxide samples. Pt electrodes were written using standard photolithography on 4mm x 4 mm or 6 mm x 30 mm Si/SiO<sub>2</sub> substrates. Dielectrophoretic trapping was performed by applying a field between the electrode and backplate of the substrate while a drop of the carbon nanotube solution was placed on the surface around the electrode (Figure 3.3). After a required time (typically 15 sec) the substrate was washed with acetone and isopropanol and blow-dried. Subsequently the substrate was annealed at 250°C for 2 hours. After that the devices can be visualized and characterized electrically.

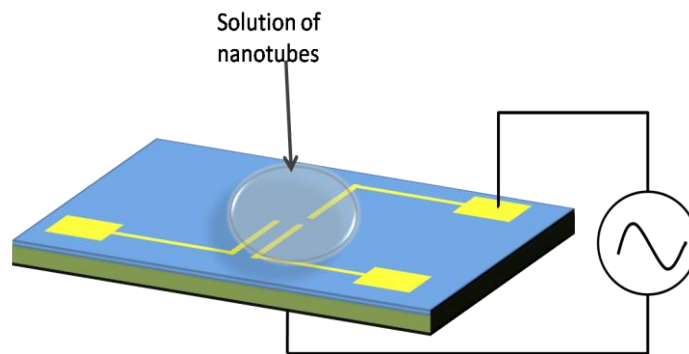


Figure 3.3. Schematic of the dielectrophoretic trapping setup. The trap waveform is applied between one of the electrodes and the backplate.

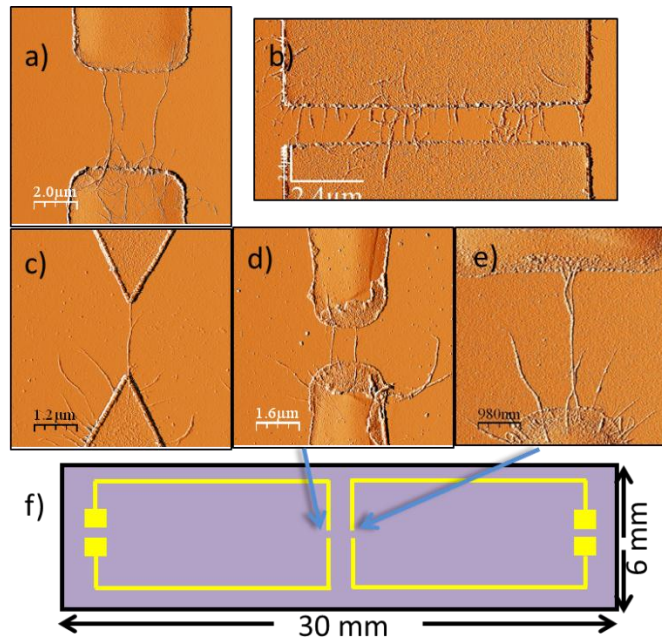
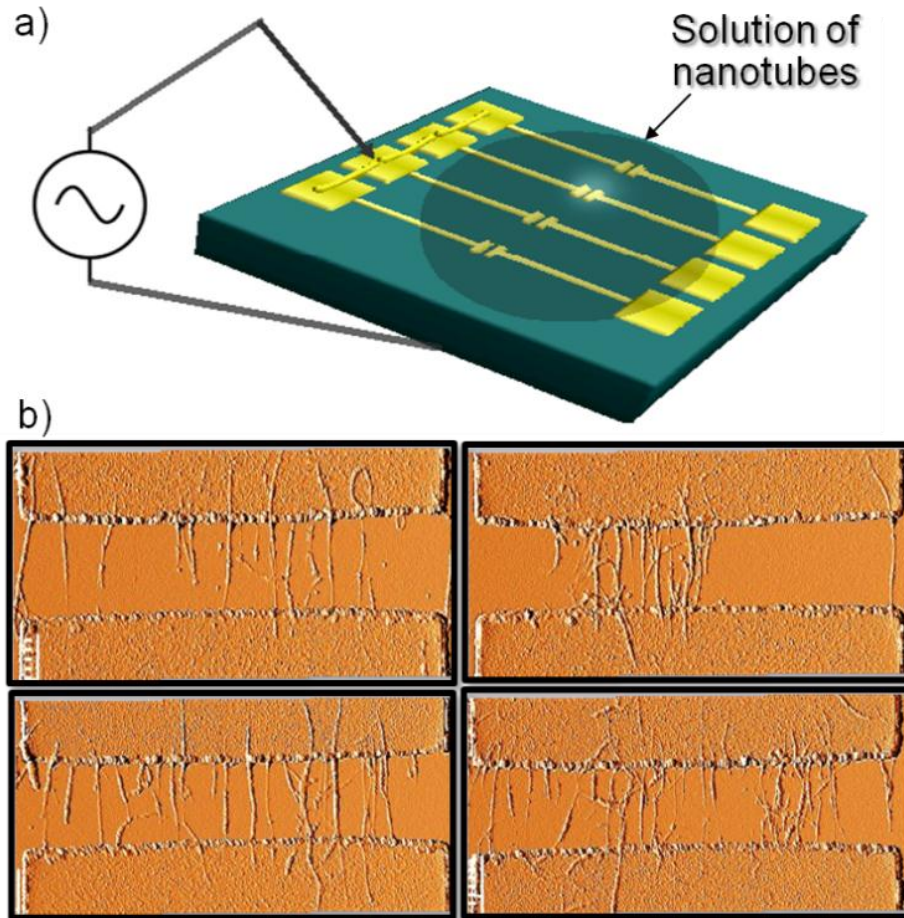


Figure 3.4. Different electrode geometries used in this study. Optimal parameters for CNT trapping: a) 25Mhz, 10V, b) 10 MHz, 3V, c) 25 MHz, 10V, d) and e) 10 Mhz, 10 V. Schematic of the substrate chosen as the most suitable for sensing trials (f). The AFM images of the electrode gaps with trapped CNTs are shown on (d) and (e).

Different shapes and sizes of electrode gaps were tested for trapping the CNTs and subsequent biosensing trials (Figure 3.4). Geometry of the electrodes as well as properties of the substrate strongly influence the dielectrophoretic trapping of CNTs. Therefore the parameters had to be optimized in every case. Among various electrode and sample designs the most suitable was chosen during the initial sensing trials. Most of the sensing results described in this thesis were obtained on the 6 mm x 30 mm Si/SiO<sub>2</sub> samples with two pairs of platinum electrodes. The geometry of the gaps for this kind of samples is shown in Figure 3.4 (d, e). In contrast to samples of smaller sizes (4mm x 4 mm), liquid manipulation is much easier on the chosen larger size samples since it makes it possible to integrate microfluidic channel or reservoir for analyte solutions on top of the substrate. The DEP parameters used for chosen samples were the following: 10 MHz, 10V, 15 sec.

**3.4.3. Scalability of DEP.** The possibility to scale-up dielectrophoretic trapping of CNTs was subsequently investigated. For this purpose parallel connection of the electrodes and common backplate were used for applying the AC voltage. In this manner the dielectrophoretic force was generated simultaneously at

all connected electrodes and the nanotubes were trapped concomitantly at several positions (Figure 3.5).



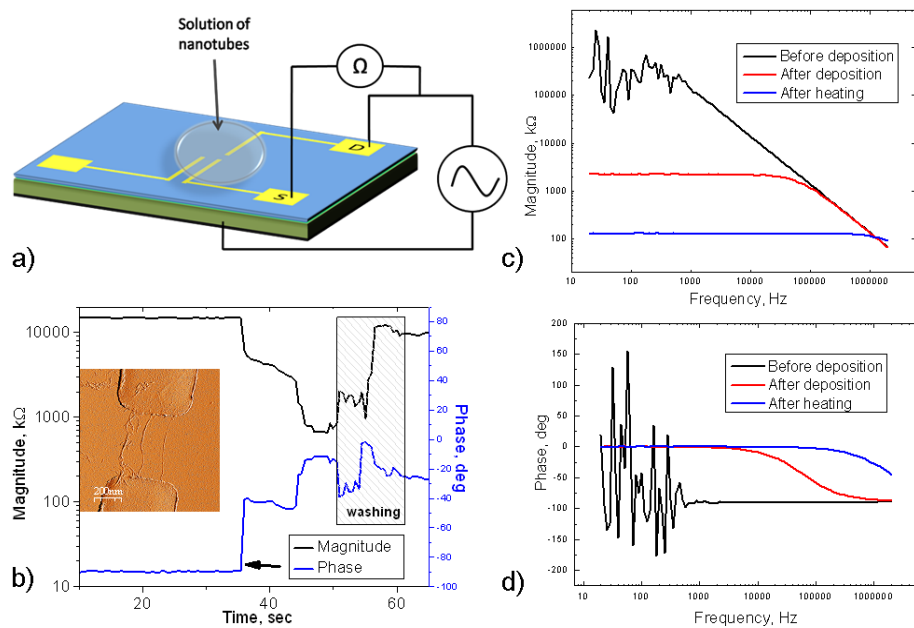
**Figure 3.5.** One of the possible ways to upscale the DEP trapping of CNTs. a) Schematic of the trapping procedure. b) AFM images of four devices on the chip after a single trapping step.

Another way to scale up the trapping is by using a different configuration of the electrodes. In this case the DEP force can be applied between the connected microelectrodes on the substrate and a (quasi)-reference electrode (for instance platinum wire). The advantage of this approach is that metalized backplate as well as connecting additional electrodes on the surface are not required. Therefore it is possible to fabricate CNT devices not only on Si/SiO<sub>2</sub> samples, but also on glass and flexible polymer substrates.

**3.4.4. Monitoring the trapping of CNTs during DEP.** For optimizing the parameters of DEP it is reasonable to monitor the trapping by measuring the impedance between the electrodes. It helps to control the resistance of the device and thereby the density of nanotubes during the trapping itself. This

configuration allows disconnecting the trapping field as soon as a trapping event occurs. The value of the impedance directly provides us with a measure of the density of nanotubes.

The impedance monitoring technique is implemented by superimposing two AC waveforms: a *trap waveform* (10 V at 25 MHz) between one Pt electrode and the backplane of the substrate and a *probe waveform* (10 mV at a lower frequency 2623 Hz) between the two Pt electrodes to monitor the trapping event (Figure 3.6 a). The trap waveform is supplied by a function generator (Agilent 33250A) and the impedance is recorded at the frequency of the probe waveform using an Agilent E4980A LCR Meter.



**Figure 3.6. Monitoring the dielectrophoretic trapping of the nanotubes. a)** Schematic of the experimental set-up used to monitor the dielectrophoretic trapping of nanotubes. The trap waveform is applied between the drain (D) electrode and the backplane, while the impedance is monitored across the drain and the source (S) electrode using the probe waveform. **b)** In situ impedance monitoring during DEP trapping: the magnitude and phase of the gap impedance are plotted as a function of time. A CNT suspension is first introduced (arrow) and ~10s afterwards clear steps can be seen indicating trapping events. (Inset: an AFM picture showing the trapped nanotubes across the gap). **c)** the magnitude and **d)** the phase of the impedance spectrum of the electrode gap in air before and after deposition, as well as after annealing.

Figure 3.6b shows a typical deposition sequence. Initially, the substrate is in air and the system is purely capacitive with the impedance higher than 10 M $\Omega$  and a phase of  $-90^\circ$ . A drop of CNTs is placed above the electrode gap. This leads to a decrease in impedance to a few MOhm and a phase of  $-40^\circ$ , due to the conductivity

of the solution. After several seconds, trapping events are registered, which appear as discrete steps both in the magnitude and phase signals. At this instance, the impedance is just a few hundreds of kOhm and the phase close to zero suggesting that CNTs are trapped (see inset AFM image). The sample is then washed with acetone during which the trap waveform is still active. The washing procedure introduces some changes in the impedance. Finally the sample is removed and washed thoroughly with isopropanol.

A frequency of 2623 Hz of the probe waveform was chosen so that the change in the signal between the empty situation and the trapped condition is maximized with a good detection sensitivity and fast response time. The contact resistances of the devices could be reduced by around an order of magnitude by a subsequent rapid thermal annealing step at 650°C (45s) or a longer heating period (2h) at 250°C in argon (Figures 3.6c and 3.6d). Online impedance monitoring can be used either to actively control the amount of nanotubes trapped at a specific location or as a way to calibrate the deposition time needed for a given nanotube suspension.

**3.5. Passivation of electrodes.** The chosen device architecture is designed in such a way that it can be used in a liquid environment – for sensing different analytes in serum or other biological liquids. These fluids contain a significant amount of electrolytes that cause a background electrical current between electrodes through the liquid. This ionic current is strong enough to affect sensing results. Besides, electrodes can provide large surface for electrochemical reactions. This can reduce reaction rate on the CNT surface during electrochemical functionalization and increase the limit of detection. Some components of analyte solution may also sediment non-specifically on the metal electrodes during the sensing trials and cause changes in electrical response of the device. All these factors make it important to ensure that CNT is the only conductive element that is exposed to solution. To achieve this passivation of electrodes is required. The passivation process should ensure robust and stable electrical isolation of electrodes on the areas of substrates coming in contact with the liquid. It should have a minimal effect on the electrical properties of CNTs. Furthermore, surface contamination during this process has to be avoided.

In this work we have used a second lithography step to deposit 100 nm titanium covered with 200 nm silicon oxide to passivate Pt microelectrodes near the gap. For this purpose the substrate was spin-coated with a photoresist (ma-P 1215, micro-resist technology GmbH) and baked for 2 min at 90°C to obtain a 1.5 µm thick



layer. Absorption maximum of ma-P1215 is close to 365 nm and 415 nm. Exposure of this resist to UV light causes chemical changes that increase the solubility of the exposed areas. Illumination of this photoresist with visible light of up to 580 nm can induce chemical changes. Using a confocal microscope (TCS SP2, Leica) the desired areas were first localized with a high level of precision using low power illumination by a 633 nm laser (imaging laser). Then the required areas were exposed using a high power 476 nm write-laser [129]. Writing was performed by using a piezoelectric stage with precise computer control for positioning and moving the sample. Alternatively, a well aligned mask for exposing desired areas was also experimented. In this case a mask-aligner is used for the precise positioning of the mask and exposure of the required areas.

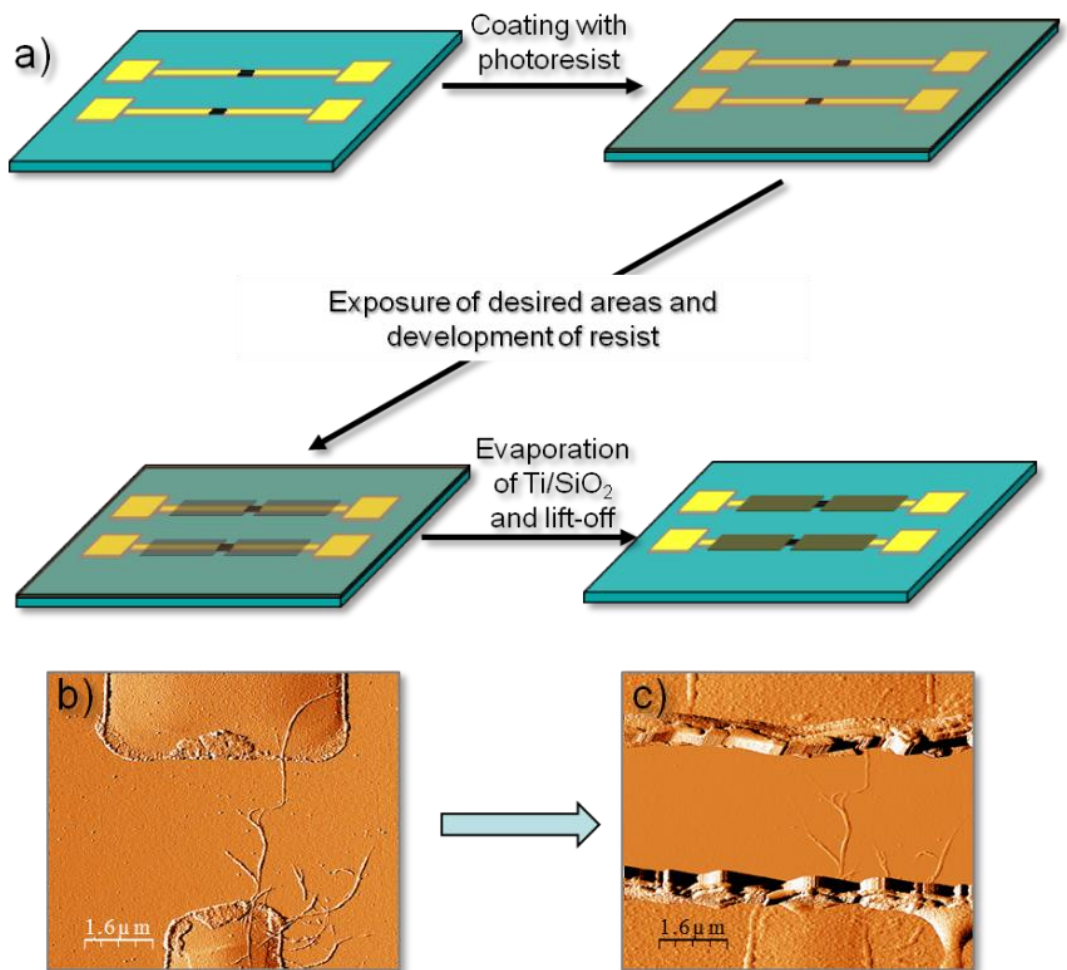


Figure 3.7. Passivation of Pt electrodes on Si/SiO<sub>2</sub> surface. a) schematic of passivation, b) and c) AFM image of CNT device before and after passivation respectively.

Using a developer solution ma-D 331, the resist on exposed areas was dissolved. After the washing with water and drying in nitrogen flow,

100nmTi/200nmSiO<sub>2</sub> was evaporated on the substrate. Such a combination of metallic (Ti) and insulating (SiO<sub>2</sub>) layers gave the best results for keeping conductivity of CNTs intact. Without the use of Ti a significant number of devices lost their conductivity, which can be explained by the mechanical breakage of CNTs upon direct deposition of the silicon oxide layer. For removing evaporated layers from the unexposed areas a lift-off in 1-methyl-2-pyrrolidone (55°C for 3 h) was performed. After this, samples were washed with acetone and isopropanol and blow-dried. After the lift-off a thermal annealing of the samples (2 hours at 250°C or 45 sec at 600°C in argon) was also carried out.

**3.6. Chemical functionalization of CNTs.** The discussed above fabrication steps were aimed to obtain devices with high sensitivity. Another important aspect for biosensors is selectivity. A biosensor should be sensitive mainly to the molecule of interest. Other components of analyte solution should not cause any significant sensor signal changes. As it was mentioned before (*paragraph 1.2*), selectivity of the biosensor is provided by the biorecognition component of the sensor which should be in high vicinity to transducing element. In our case, receptor molecules should be immobilized onto the carbon nanotube.

Ideally the functionalization strategy should ensure that the receptor is immobilized on the surface of CNT and not on the surrounding silicon oxide. This is important for the efficient recognition and transduction processes and proper interpretation of the signal changes. For example, using a spotting technique for receptor immobilization can lead to significant amount of biorecognition events happening on the surrounding silicon oxide surface but not on the CNT surface itself. In this case, the limit of detection may be high. Electrochemical modification leads to the attachment of the receptor to CNT surface only providing the basis for highly efficient biosensor functioning. It is also possible to perform electrochemical reactions inside the microfluidic channel when the biosensing system is fully assembled. This strategy was used, for example, for the preparation of glucose-sensors.

As it was explained *in paragraph 2.4* using electrochemistry it is possible to modify CNT covalently and non-covalently. Covalent ECM has advantages due to good homogeneity and stability of ligand distribution attainable on the nanotube surface, coupled with a high signal to noise ratio. Non-covalent functionalization does not affect the underlying electronic structure of the carbon nanotubes. Two ways of electrochemical modification - covalent and non-covalent - affect the electronic properties of CNTs differently. It was demonstrated using modification of CNTs with boronic acid (BA) as an example. Boronic acid was used as a receptor for

glucose sensing. 3-aminophenylboronic acid (APBA) was used as a precursor for the modification. The functionalization is carried out according to Figure 3.8 [130].

In order to covalently attach the phenylboronic acid receptor to the nanotube, a highly reactive diazonium radical is created. It binds to the nanotube via reductive coupling. For this purpose, 5 mg of APBA was dissolved in 5 ml of 20mM  $\text{H}_2\text{SO}_4$ . After this the beaker was put into an ice bath. 5 ml of 8mM  $\text{HNO}_2$  were added slowly with continuous stirring to the reactive mixture. Shortly after the components are mixed the solution starts to become yellow indicating the formation of diazonium radicals [130].

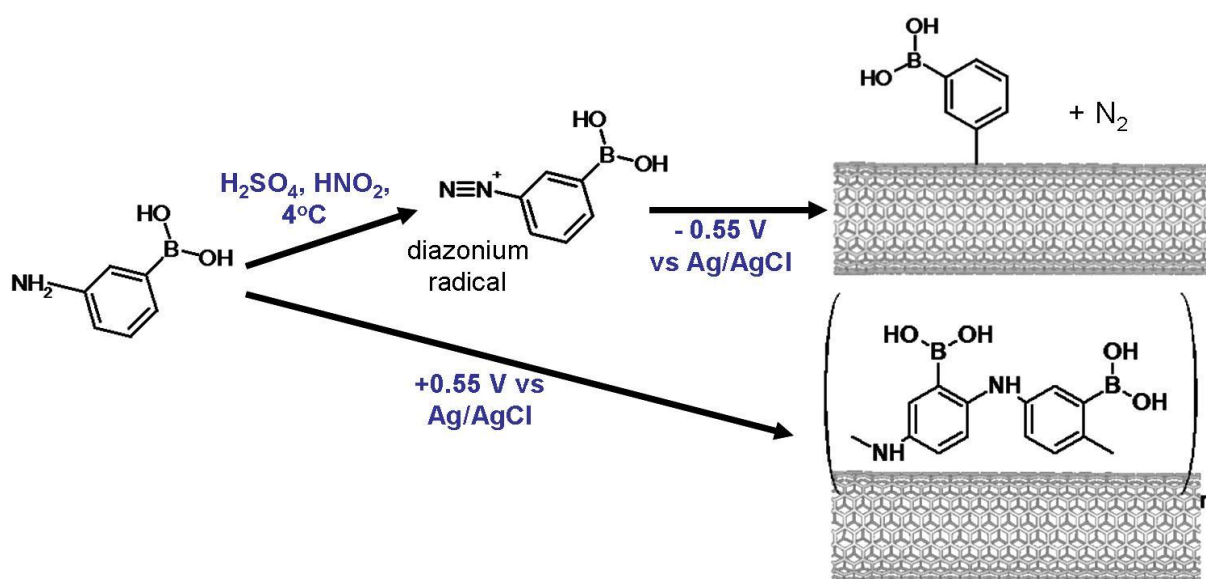
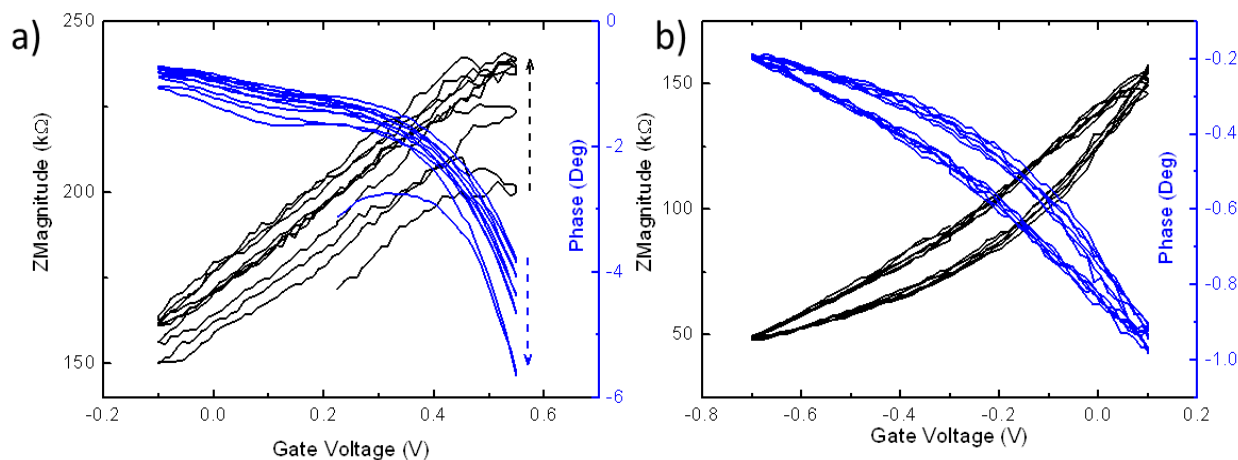


Figure 3.8. Scheme showing the functionalization strategy to obtain the BA-CNT-sensors. For covalent attachment (top route), the phenyl boronic acid precursor is converted into a reactive diazonium radical through diazotization at  $4^\circ\text{C}$ . This cold solution is flushed through the microchannel and the electrochemical functionalization of the nanotube carried out under reductive conditions ( $-0.55\text{ V vs. Ag/AgCl}$ ). For the non-covalent attachment (bottom route), the precursor is directly electropolymerized to obtain a polymer wrapping on the nanotube surface under oxidative conditions at room temperature [130].

The microchannel made of polydimethylsiloxane (PDMS) placed on the chip is filled with a fresh diazonium solution and the voltage applied at the Ag/AgCl reference electrode is swept between  $-0.1\text{ V}$  and  $+0.55\text{ V}$ . Correspondingly, the voltage on the nanotube varies between  $+0.1\text{ V}$  and  $-0.55\text{ V}$ . The impedance of nanotube device is monitored during the ECM. Figure 3.9 shows the changes in magnitude and phase of impedance during the modification. During the five cycles of voltage sweep the magnitude of impedance increases and the phase of impedance decreases. The resistance value normally increased by a factor ranging from 1.5 to 2.



Excessive covalent modification of CNT can lead to a complete loss of conductivity due to damage of the nanotube structure [130].



**Figure 3.9.** Gate dependence of the devices during (a) covalent and (b) non-covalent electrochemical functionalization of CNTs with APBA. Arrows indicate the increase of resistance (black) and decrease of phase (blue) values during reductive coupling of boronic acid.

The diameter of the nanotubes also increases after the ECM. This was observed in AFM images of the device before and after covalent functionalization. The changes of CNT properties were also monitored using Raman spectroscopy. The D-band of CNTs (which corresponds to Raman intensity peak at  $1300\text{ cm}^{-1}$  frequency) is associated with defects in the carbon nanotube structure. It is clear that during the covalent attachment of the diazonium radical to the nanotube some of the native  $sp^2$  bonds of the CNT wall are destroyed to form a new bond. This can be observed as an increase of D-peak in Raman spectra as it is seen from Figure 3.10 d.

Non-covalent attachment of the receptor was achieved via direct electropolymerization of the APBA under oxidative conditions, by sweeping the voltage at the nanotube from  $-0.1\text{ V}$  to  $+0.7\text{ V}$  versus Ag/AgCl in the ethanol solution of  $5\text{ mM}$  APBA containing  $100\text{ mM}$   $\text{LiClO}_4$ . In this case the resistance increased only slightly during the sweep (Figure 3.9 b). After functionalization,  $10\text{ mM}$   $\text{H}_2\text{SO}_4$  and water were flushed through the microchannel. The non-covalent coupling results in a height increase of  $3\text{--}5\text{ nm}$  as inferred from AFM measurements [130].

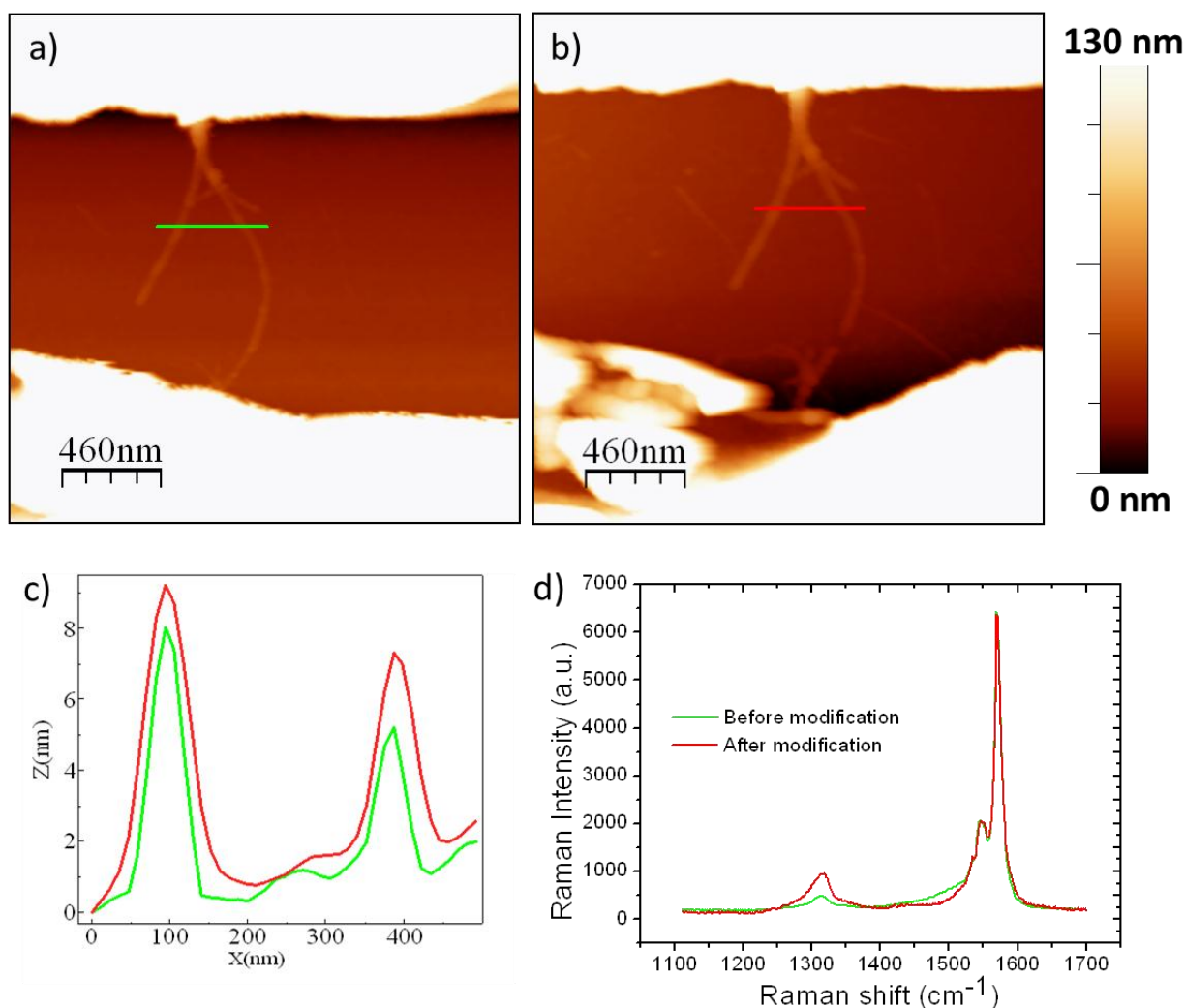
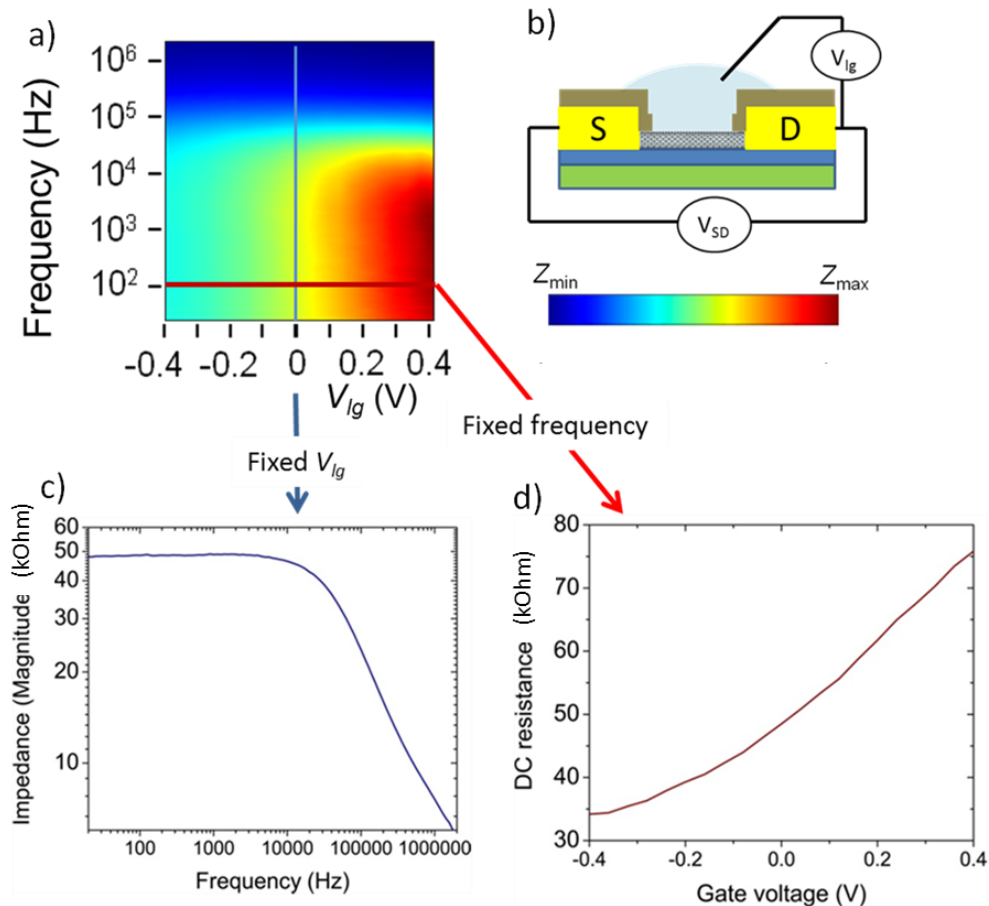


Figure 3.10. Electrochemical functionalization of CNTs. AFM images of the device before (a) and after (b) covalent modification of CNTs with APBA and corresponding height profiles (c) show increase of the nanotube diameter. Raman spectra before and after modification (d) showing a clear increase in the D-band intensity signifying the formation of covalent bonds with the BA moieties.

**3.7. Sensing strategy.** Impedance measurements at different gate voltages were taken for the detection of biomolecules. The impedance ( $Z$ , a complex quantity with magnitude and phase) was measured using an impedance analyzer (Agilent 33250A LCR Meter) in a frequency range of 20 Hz to 2 MHz. Using an AC signal to record the sensor characteristics improves the signal-to-noise ratio and reproducibility compared to a DC measurement while enabling acquisition of both the magnitude and phase spectra of the sensor. This frequency response is measured at varying gate voltages to characterize the field-effect behavior. The gate voltages are maintained in the range of -0.4 V to 0.4 V (at the Ag/AgCl reference) [130].

The resulting dataset can be visualized in the form of a magnitude map as shown in Figure 3.11. The X-axis corresponds to the liquid gate voltage, while the Y-axis indicates frequency. Every point in the image corresponds to the magnitude of impedance at a certain frequency and gate voltage. It is possible to derive both frequency- and gate-dependence from such kind of measurement. Therefore 2D Z-maps contain the complete electrical characteristics of the sensor and can be used to probe their transduction mechanism in detail.



**Figure 3.11. Detection strategy: a) ZMagnitude-2D-map, b) schematic of a liquid gated CNT-FET, c) frequency dependence of magnitude of impedance at fixed gate voltage 0V, d) gate-dependence of magnitude of impedance at fixed frequency 100 Hz.**

Most of the devices obtained in described way show a semi-metallic transport behavior with a gate modulation of less than an order of magnitude, signifying the predominant presence of metallic tubes. Due to the low density of states around the Fermi level, metallic nanotubes show a slight gate modulation [131] and are hence suited for field-effect based detection [107]. Unlike other FET-based sensors, which require semiconducting materials, our AC measurement strategy coupled with the use of a stable reference electrode enables the use of metallic tubes

as detectors. Another advantage of using metallic nanotubes lies in their low resistance in the  $k\Omega$  range, which facilitates improved sensor signals in comparison to their semiconducting counterparts [130].

## 4. Sensing of saccharides

**4.1. Introduction.** Carbohydrates are important biomarkers for different diseases. The most common example of them is glucose as a marker for diabetes [132]. Glucose sensors mostly rely on the use of an enzyme such as glucose oxidase (GOx) or glucose dehydrogenase to generate redox mediator species, which are thereafter detected electrochemically [133]. These sensors require the coupling of the enzyme in close proximity to the transducing electrode to obtain high sensitivity. CNTs have also been employed as electrode material in electrochemical sensors for glucose [61, 134]. While providing good selectivity to glucose, such sensors rely on the rate of reaction between the enzyme and glucose and hence are inherently sensitive to factors influencing enzyme activity. In addition, the limited stability of the immobilized enzyme may restrict the shelf life of such sensor systems and their application for *in vivo* glucose monitoring [135]. Apart from electrochemical transduction, field-effect based detection has also been presented using a CNT-GOx sensor [40]. However, the sensing mechanism and concentration dependence were not discussed there [130].

During the last decade there is a tremendous amount of research directed towards the realization of non-enzymatic glucose sensors [136]. One major approach involves the direct oxidation of glucose at nanostructured electrodes [137-139]. The nanostructuring allows the detection of glucose at a lower overpotential [140]. However, such electrodes are very sensitive to adsorbed interferents such as the amount of chloride and suffer from surface fouling over longer times [130, 141, 142].

An alternative enzymeless approach for glucose detection involves the use of fully synthetic receptors such as boronic acid (BA) containing compounds. BA binds reversibly with *cis*-1,2- and *cis*-1,3-diols to form five- and six-membered cyclic esters respectively [143, 144]. Based on this, fluorescence sensors, [145, 146], potentiometric / amperometric sensors [147, 148] and recently a field-effect based sensor [149] have been demonstrated. The chemical structure of the boronic acid receptor can be optimized to obtain highly selective and sensitive affinity-based glucose sensors [145]. In addition to glucose, free carbohydrates as well as their conjugates such as liposaccharides and glycoproteins are also important in the diagnosis of pathological states such as cancer [150]. While enzyme-based sensors are not directly amenable for the detection of bound saccharides, an affinity-based sensor

can potentially be deployed for the detection of both free and bound forms of these molecules [130, 151, 152].

In contrast to enzymatic sensors, the detection principle here is based on complexation, which is a reversible, equilibrium-based reaction without the consumption of the analyte. So the present sensor does not rely on enzymes or electrochemical redox mediators to function but rather on a combination of affinity sensing and the field effect transistor (FET) effect. Glucose detection with BA-CNT device was chosen to be a model for investigation of the sensing mechanism in presented biosensor configuration [130].

**4.2. Device fabrication.** The details of device fabrication were described in previous chapter. As it was mentioned there, the control of the kind of chemical coupling (covalent or non-covalent) between the boronic acid receptor and the nanotube was achieved using electrochemical modification. The first attempts to detect glucose using boronic acid functionalized CNTs were performed in an open system without the application of liquid gate voltage, by measuring the impedance spectra of the device before and after its exposure to analyte solution (Figure 4.1a). This configuration was discarded due to significant amount of aggregates on the surface of the sensor as observed using an AFM (Figures 4.1b and 4.1c). These adsorbates affect the quality of sensing and as an outcome no reproducible and reliable results were achieved. In later investigations the contamination of the SiO<sub>2</sub> surface could be eliminated when the measurement was performed using continuously applied liquid gate voltage. The sample dimensions and design were therefore changed in order to allow better manipulation of liquids and easier integration of the gate electrode.

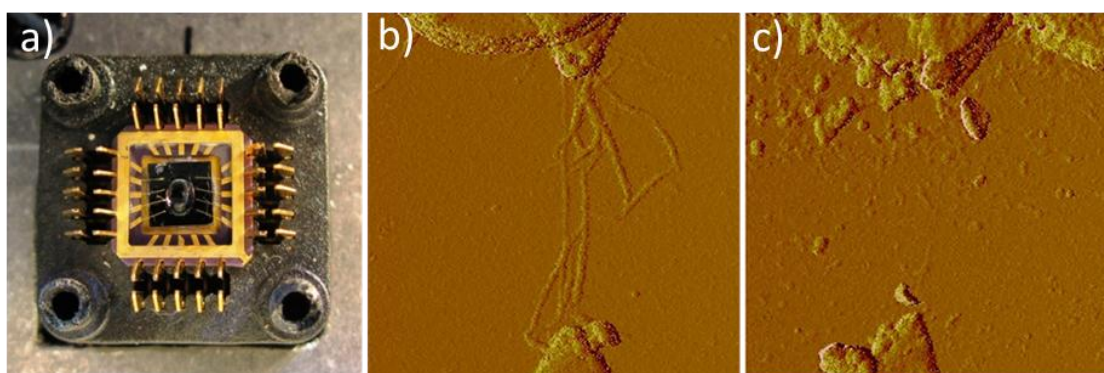


Figure 4.1. Sensing in the open system without applying liquid gate voltage: photo of the integrated chip with the drop of analyte solution (a), AFM image of the device before (b) and after (c) sensing trials.



Liquid handling was improved by using a microfluidic channel connected to a flow-control system. For the microfluidic fabrication polydimethylsiloxane (PDMS) was used to prepare 2 mm thick plates with a 30  $\mu\text{m}$  channel. The degassed mixture of PDMS and curing agent (PDMS: Sylgard 184, 10:1 ratio) was poured on a SU-8 mold and baked at 65°C for 1 h. The sample was then placed on the chip-carrier, wire-bonded and covered with the PDMS and glass plates, so that the nanotube devices were integrated into the microchannel (Figures 4.2b and 4.2c). The flow of the solutions was regulated using a computerized pressure-based flow control system (MFCS-4C, Fluigent). An Ag/AgCl reference electrode placed in contact with the liquid in the microfluidic circuit allows us to set the potential precisely on the surface of the nanotube. This electrode serves two purposes namely as a reference electrode for electrochemical functionalization and as a gate electrode for measuring the FET characteristics. The Ag/AgCl reference that acts as a gate electrode is placed in the waste reservoir that is connected to the outlet [130].

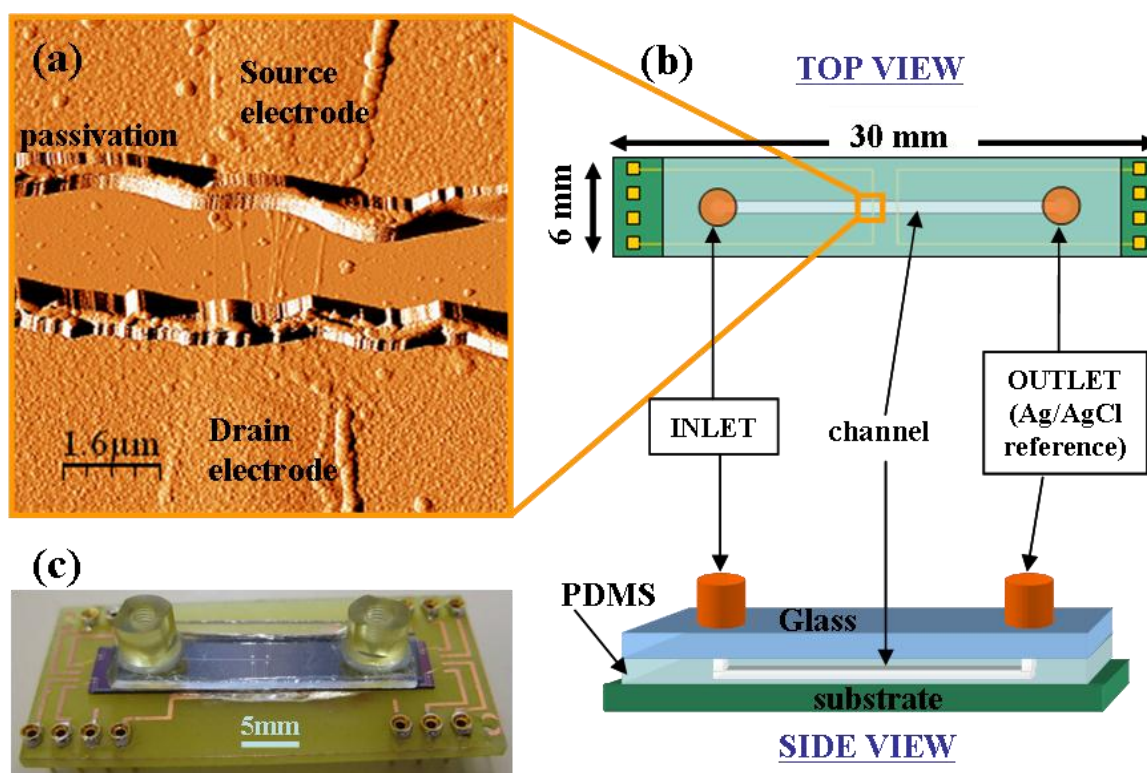
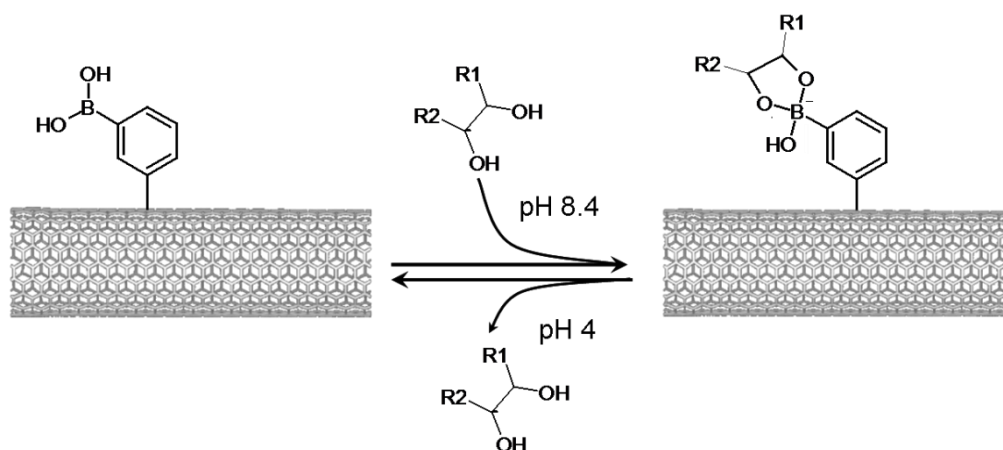


Figure 4.2. (a) An AFM image of the electrode region of a BA-CNT sensor after the sensing trials. The SiO<sub>2</sub> passivation layer can be seen extending above both electrodes. (b) A schematic representation of the microfluidic set up with the location of the nanotube device. The channel is carved out in the PDMS layer and the nanotube device is directly below the channel on the substrate. (c) a photograph of the final assembled device [130].

### 4.3. Sensing results.

The sensing trials were performed using the measurement strategy described in the previous chapter on more than 10 devices using varying concentrations of glucose solutions in 10 mM phosphate buffer (pH 8.4). First the sensing response of the covalently coupled BA-CNT devices will be discussed.



**Figure 4.3.** Affinity sensing of saccharides: binding of diol to boronic acid-functionalized carbon nanotube at a pH of 8.4. The formation of the ester leads to an increase in the relative negative charge at the surface of the nanotube which can be detected electrically. The reaction is reversible at low pH. In an acid solution at a pH of around 4, the attached saccharide can be released going back to the initial state [130].

**4.3.1. Sensitivity to glucose.** The general idea behind the functioning of the sensor is shown in Figure 4.3. The binding of glucose to the attached BA receptors leads to a change in the net charge distribution on the nanotube surface, which can be sensed through changes in the impedance of the nanotube. The Z-maps of a typical sensor in buffer and in glucose are shown in Figures 4.4a and 4.4b respectively. The magnitude plots show subtle differences in the low frequency region. These variations are further apparent in the cross-section profiles collected in Figures 4.5a and 4.5b. These section profiles are extracted from the 2D Z-maps by plotting either the frequency response at a fixed gate voltage or the gate dependence at specific frequencies. From the magnitude spectrum in Figure 4.5a it is apparent that at low frequencies, the device acts as a resistor, while capacitive effects become dominant as the frequency increases. The resistance of the device shows nominally a 5% increase upon binding of glucose, as is apparent in Figure 4.5a. The increase in resistance can be attributed to stronger scattering centers due to the increase in net charge when glucose binds to the receptor sites on the nanotube (Figure 4.3). This mechanism is consistent with the occurrence of analyte dependent carrier scattering [41]. Further support for the scattering based sensor response is obtained from the



gate voltage dependence in Figure 4.5 (b), where it is apparent that the higher resistance persists over almost the entire gate voltage range.

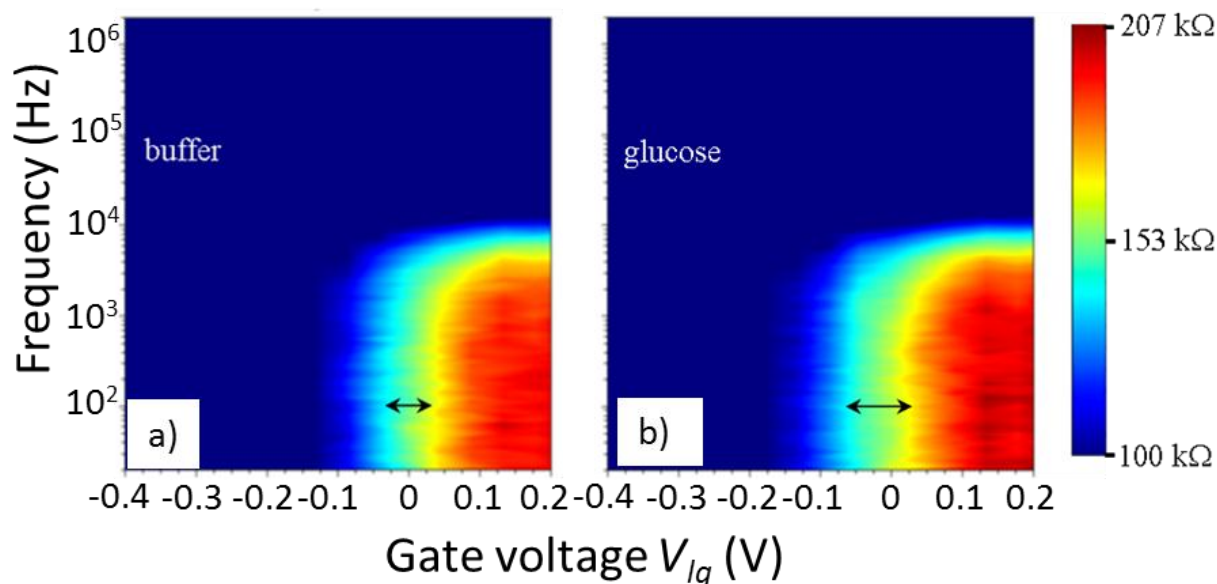
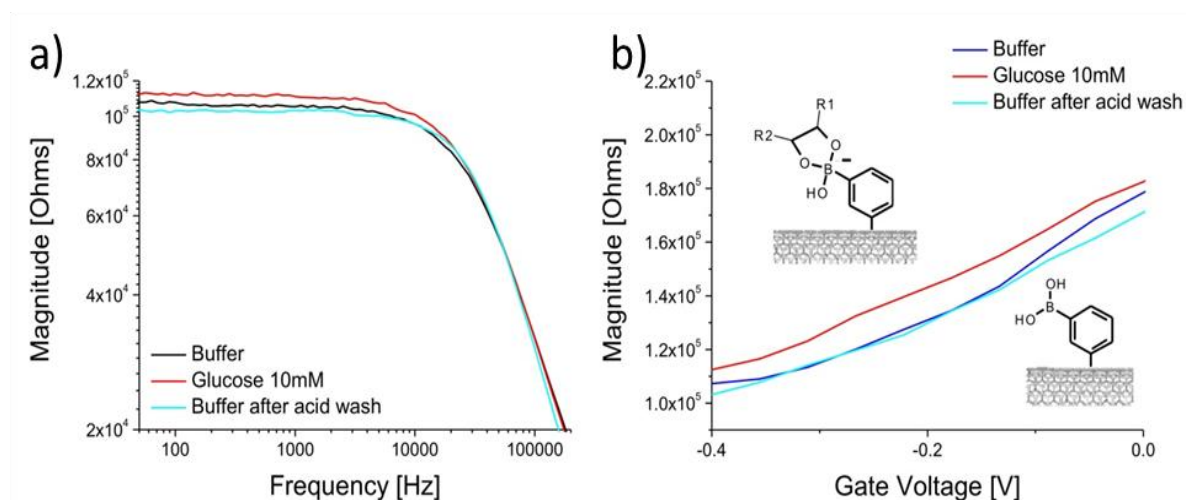


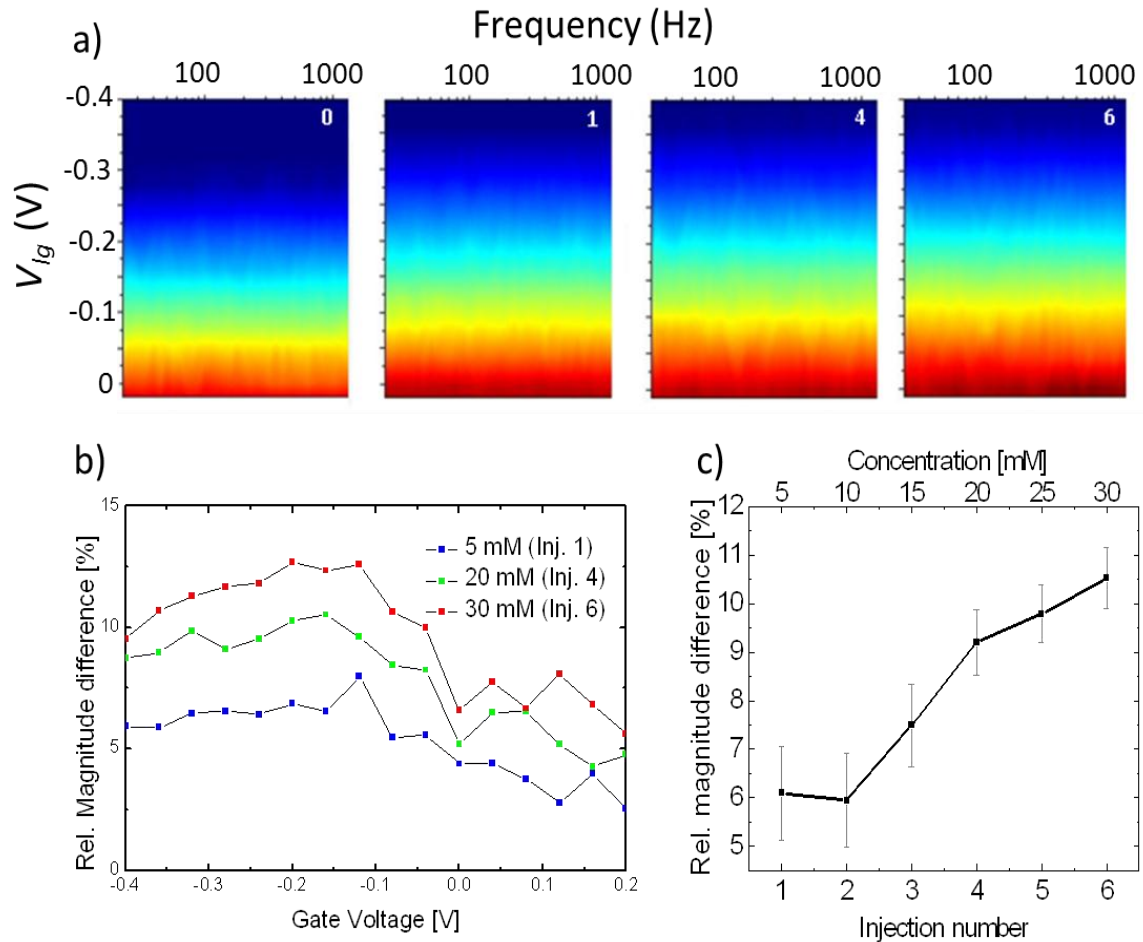
Figure 4.4. 2D-impedance ( $Z$ -) maps showing the impedance as a function of frequency and liquid gate voltage. Magnitude [Ohms] maps of the impedance of covalently functionalized BA-CNT devices are shown in buffer (a) and in 10 mM glucose (b). Subtle changes in the impedance response can be clearly seen upon exposure of the sensor to glucose as indicated by the twin-headed black arrows [130].

To measure the concentration response of covalently modified BA-CNT, the devices were exposed to successive aliquots of a 5 mM glucose solution. A 1  $\mu$ L aliquot of the analyte solution was flushed over the sensor at a precise speed, followed by 9  $\mu$ L of buffer solution at which point the flow was stopped and the impedance measured. This cycle was repeated several times. The  $Z$ -maps for the various injections are collected in Figure 4.6a.



**Figure 4.5. Reversibility of the binding of glucose in covalent-BA-CNT-sensors.** Glucose binds to phenyl boronic acid at a  $\text{pH} > 5.8$ . Hence the washing step to dissociate glucose from BA involves the flushing of an acid solution ( $\text{pH} = 4$ ). (a) Frequency response and (b) gate dependence of the sensor at low frequencies in buffer, in glucose and after the acid wash. The impedance response of the sensor recovers after this washing step. The magnitude of impedance at low frequencies increases by around 5% in the presence of 10 mM glucose [130].

It is apparent that with every injection the response to glucose becomes stronger. This is further clear from the line profiles in Figure 4.6b, which show the relative increase in the magnitude of impedance for various injections. The calibration plot shown in Figure 4.6c displays a clear trend of the sensor signal as a function of glucose exposure. It can be seen that with every exposure, the sensor signal increases, however it tends to saturate for larger injections. This behavior is expected due to the finite number of BA moieties that can complex with the incoming glucose molecules. It is noteworthy that by using this multiple injection procedure one can adjust the volume flushed through the channel to operate the sensor in a desired concentration range [130].



**Figure 4.6. Concentration dependence of the covalent-BA-CNT-sensors.** a) Z-maps in the absence of glucose (0) and after 1, 4 and 6 injections of 5 mM glucose. Scale: 20 kOhm - 40 kOhm. Every injection comprises of a 1  $\mu$ L aliquot of 5 mM glucose followed by a 9  $\mu$ L aliquot of buffer solution, at which point the impedance response is acquired. b) The plot shows the relative change in the magnitude of impedance glucose injections. c) Calibration curve plotting the relative % change as a function of glucose exposure showing clear concentration dependence. The actual amount of glucose can be read from the top Y-axis [130].

Control experiments were performed with CNTs that were not functionalized. Figure 4.7 shows the gate- and frequency dependences of such devices initially and after exposure to glucose. No significant changes were observed in this case [130].

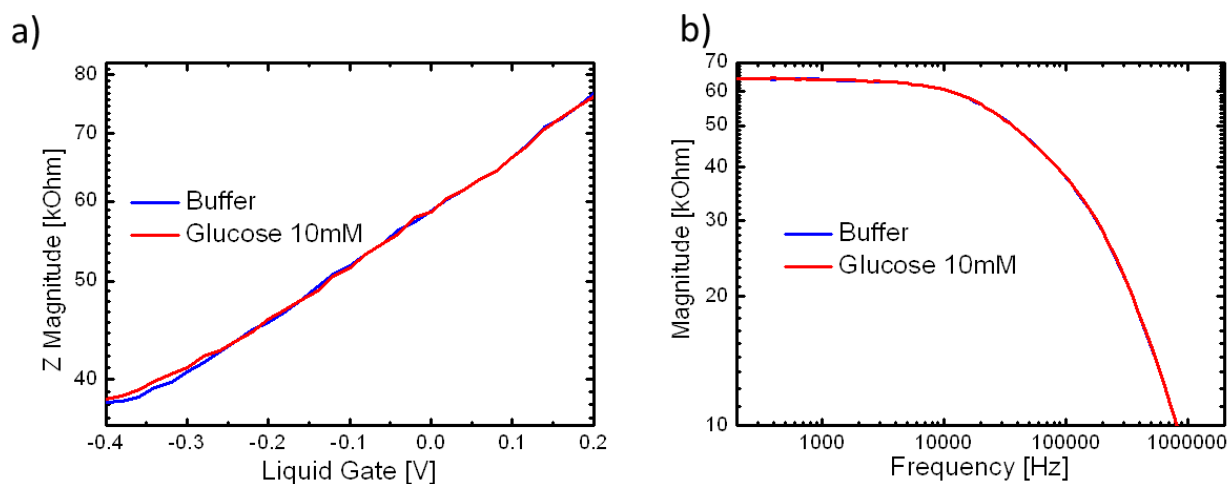
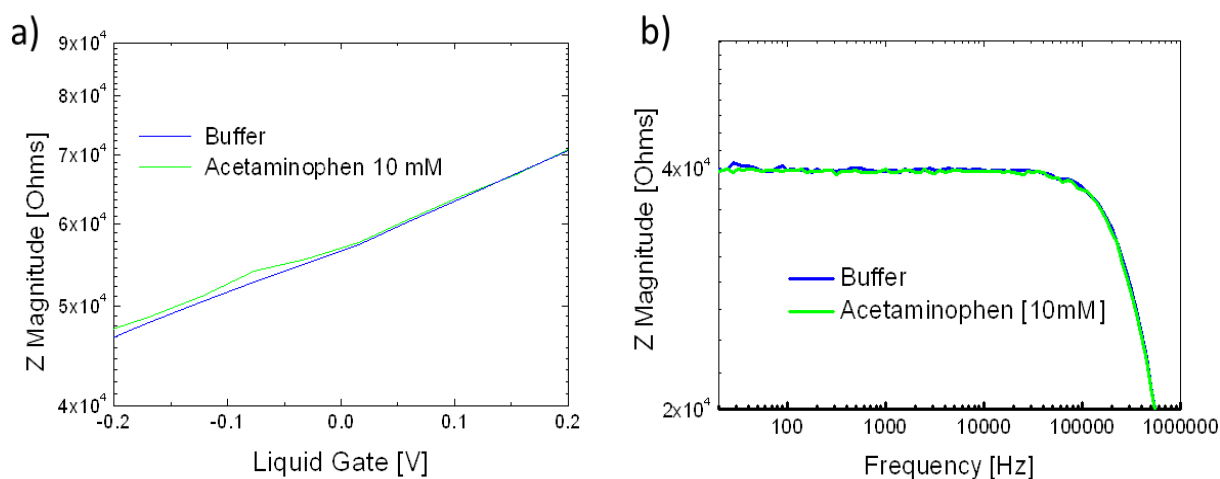


Figure 4.7. Control experiment with unmodified CNT devices. A non-functionalized CNT shows negligible changes in its response when exposed to glucose as can be seen from the (a) gate voltage scans and (b) frequency responses [130].

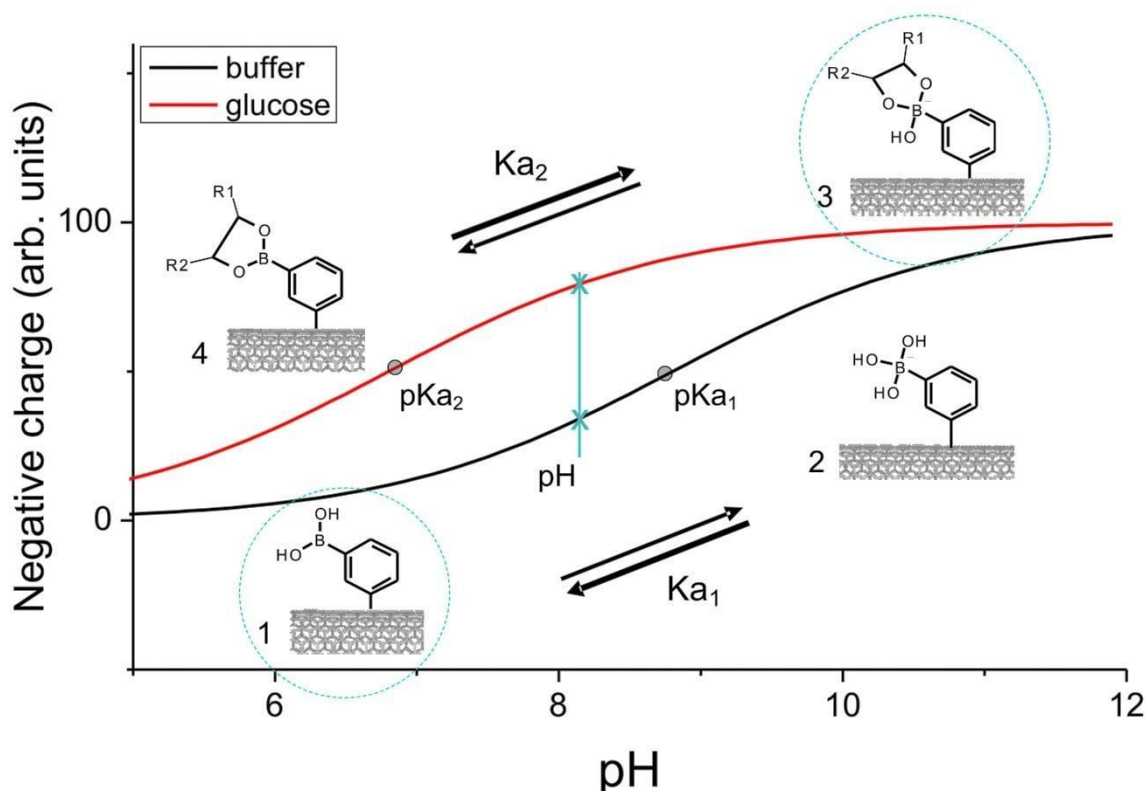
**4.3.2. Response to other monosaccharides and interferences.** As it was mentioned before (*paragraph 4.1*), boronic acid is not a specific receptor for glucose. It can bind other saccharides as well. The detection of fructose binding was demonstrated using the presented BA-CNT sensor. As it is seen from Figure 4.8, the response of the sensor to fructose was similar to the glucose case. Furthermore, the response was reversible as is apparent in the recovery of the signal after the acid wash.

Electrochemical glucose sensors suffer from the interference of other substances that may be present in body fluids. Among such interfering molecules is acetaminophen [133]. The sensor response on the presence of acetaminophen was evaluated here (Figure 4.8). It is apparent that acetaminophen did not cause signal change of BA-CNT sensor for glucose. This is expected since the BA moieties exhibit a high degree of selectivity to saccharides [130].



**Figure 4.8. Selectivity of the covalent BA-CNT-sensors.** The BA-CNT sensors presented here are not sensitive to interferents that are common for enzymatic sensors. Here the case of acetaminophen is presented. The BA-CNT sensor shows negligible response when exposed to a solution of acetaminophen (even at a high concentration of 10mM) as can be seen from the (a) gate voltage scans and (b) frequency response [130].

**4.3.3. Sensing mechanism.** In order to elucidate the sensing mechanism it is important to understand the physico-chemical phenomena responsible for the impedance changes brought by exposure to glucose. For this purpose, the reversible binding of boronic acid to glucose on the surface of CNTs must be presented first. A schematic of this equilibrium can be seen in Figure 4.9. In the absence of glucose, the neutral (1) and negatively charged (2) forms of the attached phenyl boronic acid exist in equilibrium with a certain pKa ( $pK_{a1} \approx 8.8$ ) [153]. cis-1,2- and cis-1,3-diols complex with boronic acid at  $pH > 5.8$  to form cyclic esters. The complexed forms (3 and 4) have a pKa ( $pK_{a2} \approx 6.8$ ) that is lower than that of the free BA [153, 155]. Thus all the four species are in equilibrium leading to a net charge on the complexed or non-complexed BA species depending on the pH (Figure 4.9). In order for the sensor to detect the binding of glucose, there should be some difference in the net charge before and after binding [130].



**Figure 4.9.** Charge distribution on the surface of BA-functionalized CNTs calculated using the Henderson-Hasselbach relationship. The black curve corresponds to the case without glucose, while the red curve represents the situation with glucose. ( $pK_{a1} \sim 8.8$ ,  $pK_{a2} \sim 6.8$ ). The cyan-colored vertical line corresponds to the working pH of 8.4. Refer to text for further details [130].

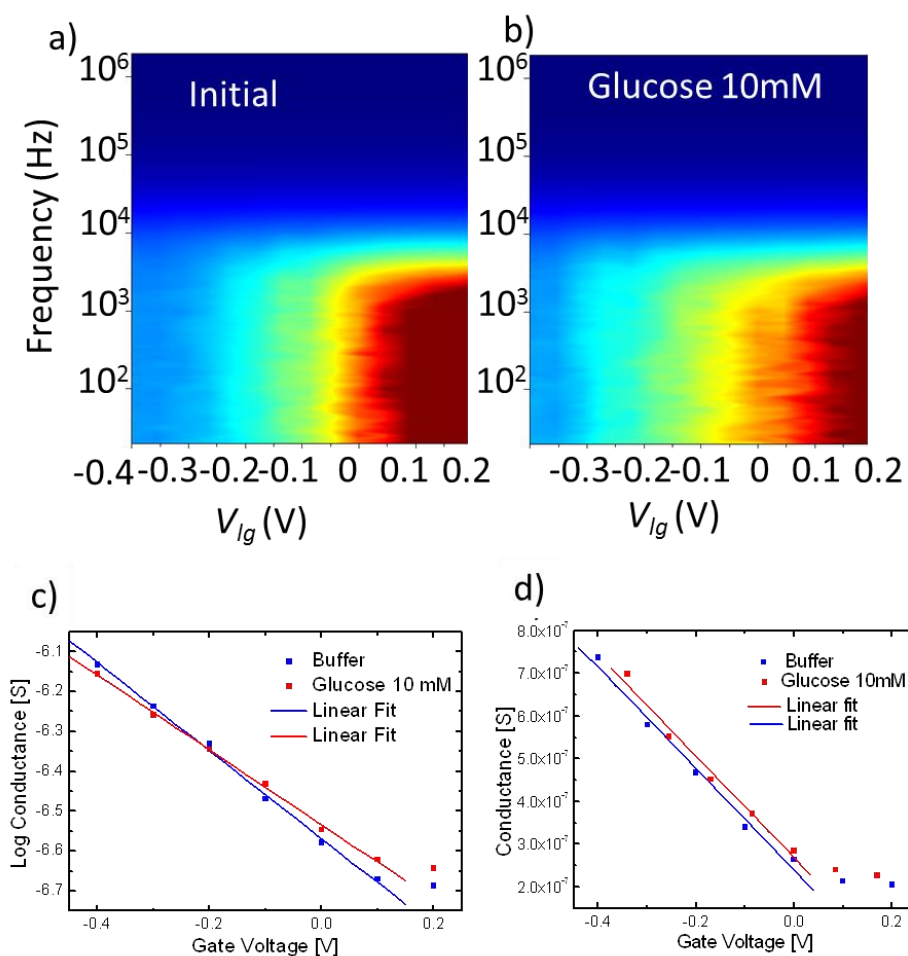
Based on this, the optimal working pH would be the one that has a value between  $pK_{a1}$  and  $pK_{a2}$ . Due to this reason, the working pH was chosen to be 8.4. At this pH (marked by the vertical line), the relative charge on the complexed species is maximized with respect to the charge on the unbound species. In other words, at this pH, 1 and 3 are expected to be the dominant species before and after complexation respectively. As a result, the net charge on the surface of the CNT is relatively higher in the presence of glucose in comparison to the situation without glucose. It is noteworthy mentioning here that the exact values of  $pK_{a1}$  and  $pK_{a2}$  of the BA-CNT assembly might be slightly different from the known values. However, this would only shift the black and red curves slightly reducing the difference in the net charge by a small amount. Based on this picture, it can be concluded that the increased charge on the covalently attached BA centers in the presence of glucose leads to a stronger scattering of charge carriers resulting in a higher resistance [130].

While the consistent increase in resistance for a broad gate voltage range and its clear concentration dependence strongly support the scattering mechanism related to changes of charge carrier mobility, two other mechanisms also deserve attention as has been postulated for semiconducting CNTs [41, 155]. The first one concerns an alternative effect of the analyte dependent change in charge on the nanotube surface namely a threshold voltage shift, analogous to the working principle of an ion-selective field-effect transistor (ISFET) [156]. In a second scenario, the docking of analytes on the nanotube surface could lead to an increased screening of the gate potential by an increase in the thickness or a change in dielectric constant of the receptor coating. This would alter the effective gate capacitance, which would result in a change in the slope of transconductance characteristics, signifying a modulation of the gate coupling coefficient [41, 130].

Non-covalent attachment of the receptor does not alter the electronic structure of the CNT. Therefore this kind of functionalization is preferred for evaluation of these two mechanisms of sensing - threshold shift and change of capacitive coupling. As it was described previously (*paragraph 3.6*), through electropolymerization phenyl boronic acid moieties were attached in a non-covalent manner to obtain coatings of 3 nm thickness. Figures 4.9a and 4.9b show the ZMagnitude-2D-maps for the non-covalently functionalized BA-CNT device in buffer and in a 10 mM glucose solution. It is apparent that the behavior in glucose is quite different from that of the covalent-BA-CNT case, with the device having a higher resistance at negative gate voltage and a lower resistance at positive gate voltage and thus a markedly different slope of the gate-dependence of the conductance (Figure 4.9c). The reduction in slope signifies a less efficient coupling of the liquid gate. This can be explained by change in the polymer layer - increased dielectric shielding arising from bound glucose moieties on the BA layer.

Furthermore, at working pH of 8.4, the bound glucose moieties on the thick BA layer must have an average net charge higher than in the free BA condition, as discussed in Figure 4.9. That is indeed the case as can be inferred from the curve shown in Figure 4.10d (obtained after the correction for slope change), where clear thresholds shift can be extracted. For the sample in Figure 4.10, we calculated this shift to be around 24 mV. Similar values (in the range of 10 - 30 mV) have been obtained on other devices [130].





**Figure 4.10.** Sensor response for non-covalent-BA-CNT devices showing the gate dependence of the magnitude of impedance at low frequencies. ZMagnitude 2D-maps before (a) and after (b) glucose exposure. c) coupling efficiency (characterized by the slope of transconductance) of the sensor before and after exposure to a 10 mM glucose solution. d) threshold voltage calculation after correction of the coupling efficiency (c and d are derived from ZMagnitude-2D-maps). In (c), it can be seen that in the presence of glucose the coupling (slope) is lower, while in (d) the binding of glucose to BA moieties shifts the threshold voltage [130].

It could be argued that the sensor responses for the covalent case could also be explained using these two mechanisms similar to the non-covalent scenario. However, there are a number of arguments, which speak against this possibility. First, the moieties that are covalently attached are very and hence the screening effect if present is expected to be very minimal. Furthermore, since we have more than one tube across the electrodes, it is unlikely that the sparsely distributed charge centers can bring in a significant threshold voltage shift in the gate characteristics. Finally, the gate dependence of resistance in the presence of glucose shows an increase in



resistance for all gate voltages in a number of covalent-BA-CNT samples, strongly supporting the previous arguments [130].

**4.4. Conclusions.** In conclusion, using a novel type of non-enzymatic glucose sensor based on carbon nanotube devices site-selectively positioned in microfluidic channels the mechanisms of sensing were investigated. By a control of the nature of coupling between the receptor and the nanotube, it was possible to elaborate on the various transduction mechanisms with the help of the impedance detection technique. The sensitivity can be further increased by the use of appropriate receptors that provide a higher binding constant for glucose. The presented sensors are compact and show promise for applications such as in vivo glucose monitoring.

## 5. Label-free detection of few copies of DNA

**5.1. Introduction.** The detection of specific nucleic acid sequences plays a vital role in environmental, food and clinical monitoring and in forensic screening. Deoxyribonucleic Acid (DNA) hybridization is the basis for the detection of various infections and genetic disorders [157, 158]. On the other hand, the detection of messenger RNA (mRNA) is important for applications such as in the estimation of the gene expression level [159]. The introduction of real-time polymerase chain reaction (RT-PCR) [160] and microarrays [161] have revolutionized the way in which nucleic acid assays are implemented, enabling the highly sensitive detection of various biomarkers. In all these methods, the target DNA is amplified, which is a prerequisite for obtaining high sensitivity [162]. Furthermore, microarray-based detection requires a labeling step, which affects the time, efficiency and the total costs involved in the complete detection protocol. The ability to detect few copies of unlabeled DNA without the need for a bulky optical reading instrument will pave way for the facile entry of nanoanalytical devices for point-of-care diagnostics [163]. Sensors based on hybridization of nucleic acids constitute the basis for today's molecular diagnostics [164]. Such sensors find application in the detection of various diseases, depending on the presence or absence of specific nucleotide sequences. Current methods are able to identify few copies by amplifying the sample using PCR to obtain a detectable amount of DNA [165]. For applications involving comparison of gene expression levels, microarrays are used [166]. This requires the labeling of the target sequences to enable their subsequent detection using a fluorescence microscope [167].

Label-free electrical detection of DNA has been demonstrated in a variety of configurations [168]. The majority of them are based either on field-effect detection [18, 60] or on electrochemical detection [169]. 1D-nanostructures were used for improving the limit of detection of DNA assays. For example, electrochemical GaN nanowire-based DNA detection was performed by monitoring electrochemical peak of guanidine oxidation. In this case the authors could reach sub-pM level of DNA detection [170]. Gold nanowires functionalized with PNA were used for electrochemical detection of mRNA with the help of catalytic reporting. This sensor

showed sensitivity up to 100 fM RNA [171]. The best detection limit reported until now based on silicon nanowires is 10 fM [172, 173]. However, silicon nanowires suffer from extremely high resistances (in the gigaohm range), which sets a limitation on the achievable sensitivity [156]. Almost all of the silicon nanowire sensors are based on resistive detection without the use of a reference electrode. While this is useful in the demonstration of prototype sensors, the use of a reference electrode is unavoidable when one requires stability and reproducibility of the sensors [18, 174, 175]. Therefore using a CNT as sensitive element and a reference electrode to set a stable potential can bring significant improvement in the field of DNA electrical sensors. Some results on CNT-based electrical DNA detection were already demonstrated by other groups. However a significant improvement of the limit of detection was not achieved there. In comparison to state-of-the-art field-effect sensors based on carbon nanotube networks [66], the sensor presented here shows up to 5 orders of magnitude improvement in the detection limit.

**5.2. Sensing setup.** For DNA sensing the sample with CNT device prepared as described previously (*Chapter 3*) was attached to chip-carrier and wire-bonded.

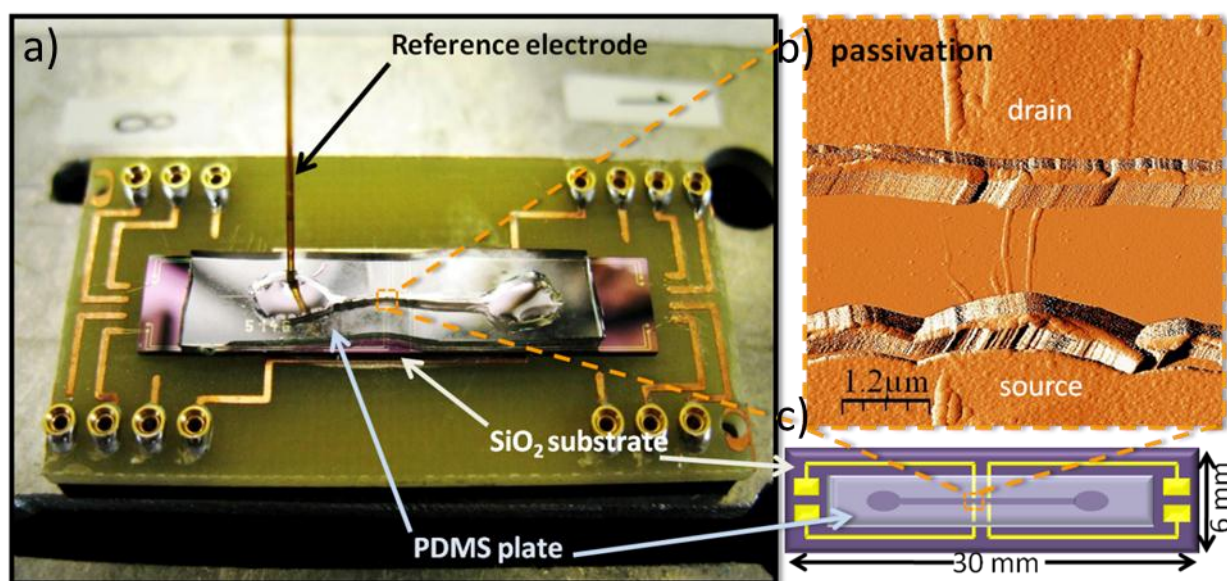


Figure 5.1. Overview of the assembled carbon nanotube DNA sensor. (a) Photo showing the chip-carrier (printed circuit board) with the wire-bonded sensor chip covered with a PDMS layer. The sensor chip contains Pt electrode gaps (b), where the nanotubes are trapped. The PDMS layer comprises two reservoirs connected by a microchannel and is shown filled with the buffer solution. An Ag/AgCl reference electrode placed in one of the reservoirs is also visible. (b) AFM image of the nanotubes dielectrophoretically trapped between Pt electrodes and subsequent passivation with SiO<sub>2</sub> (c) Schematic of the sensor chip showing the layout of the Pt electrode lines and the electrode gap [167].

The well, shaped as 150  $\mu\text{m}$  wide channel connecting two reservoirs, was cut in the PDMS plate. This PDMS plate was placed on the sample in a way that the channel was exactly above the gap between the electrodes and unpassivated parts of the electrodes were covered with PDMS (Figure 5.1a). The hole in PDMS plate was used as a microwell for liquid during functionalization and sensing. Ag/AgCl reference electrode was inserted into liquid [167].

**5.3. Functionalization of CNTs with probe DNA.** The functionalization protocol to attach the probe sequence is shown in Figure 5.2. First, 4-aminobenzoic acid (ABA) is electropolymerized on to the nanotube surface. For this purpose, the mixture of 10mM 4-aminobenzoic acid and 10mM  $\text{LiClO}_4$  in ethanol was added to the microwell and the voltage at the nanotubes was swept from -0.1 to +0.7V against Ag/AgCl reference electrode. After 5 sweeps, the chip was carefully washed with ethanol and water. This results in the non-covalent [106] wrapping of the nanotubes with poly-ABA containing free  $-\text{COOH}$  groups. In a second step amino-functionalized probe DNA is covalently coupled to the  $-\text{COOH}$  groups by carbodiimide coupling [176]. For this purpose, the well was filled for 15 min with a mixture of ethylenediaminechloride (EDC) and N-hydroxysuccinimide (NHS) 0.5 mM (1:1 ratio) in 10mM phosphate buffer pH 6 for activation of the carboxyl groups. After that most of the solution was taken out from the well (avoiding the drying of the coupling mixture on the substrate). Then the well was filled with 3'- $\text{NH}_2$ -DNA (MWG) in the 10mM phosphate buffer pH 7.4 for 30 min. 20mM ethanolamine in 10mM phosphate buffer pH 7.4 was used to block remaining activated carboxyl groups. The well was subsequently washed with 10mM phosphate buffer pH 7.4 [167].

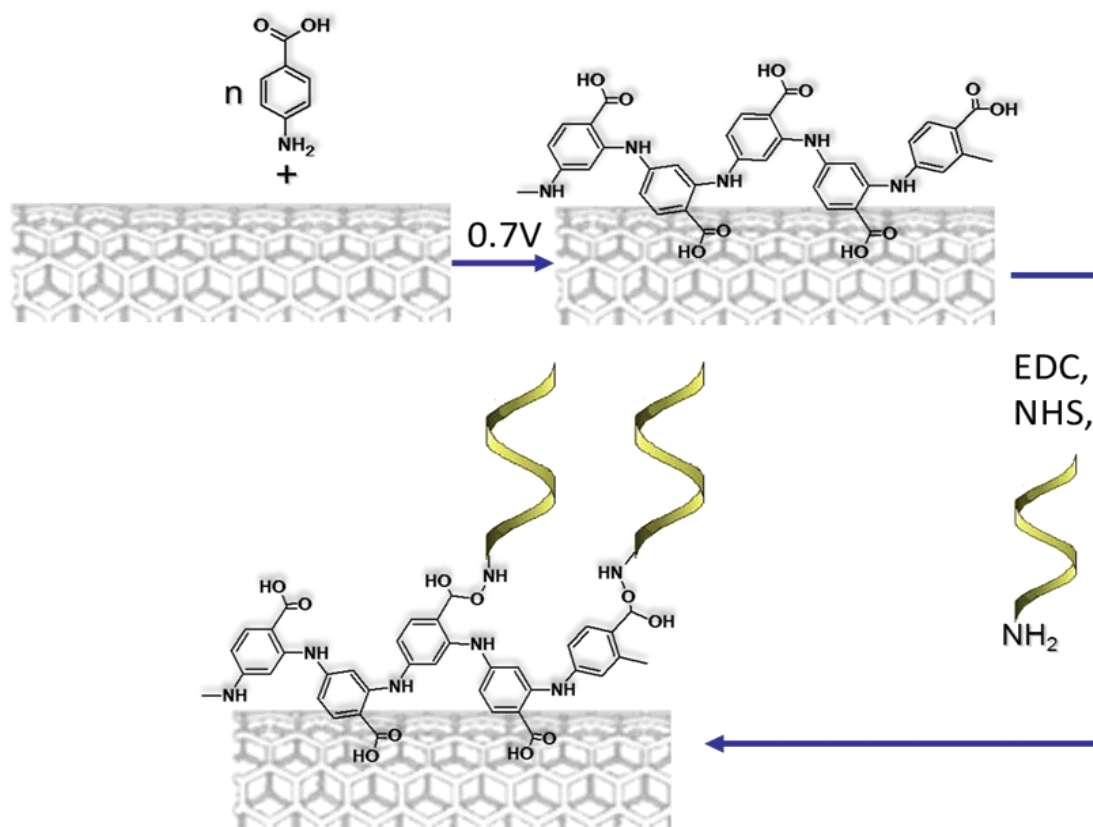
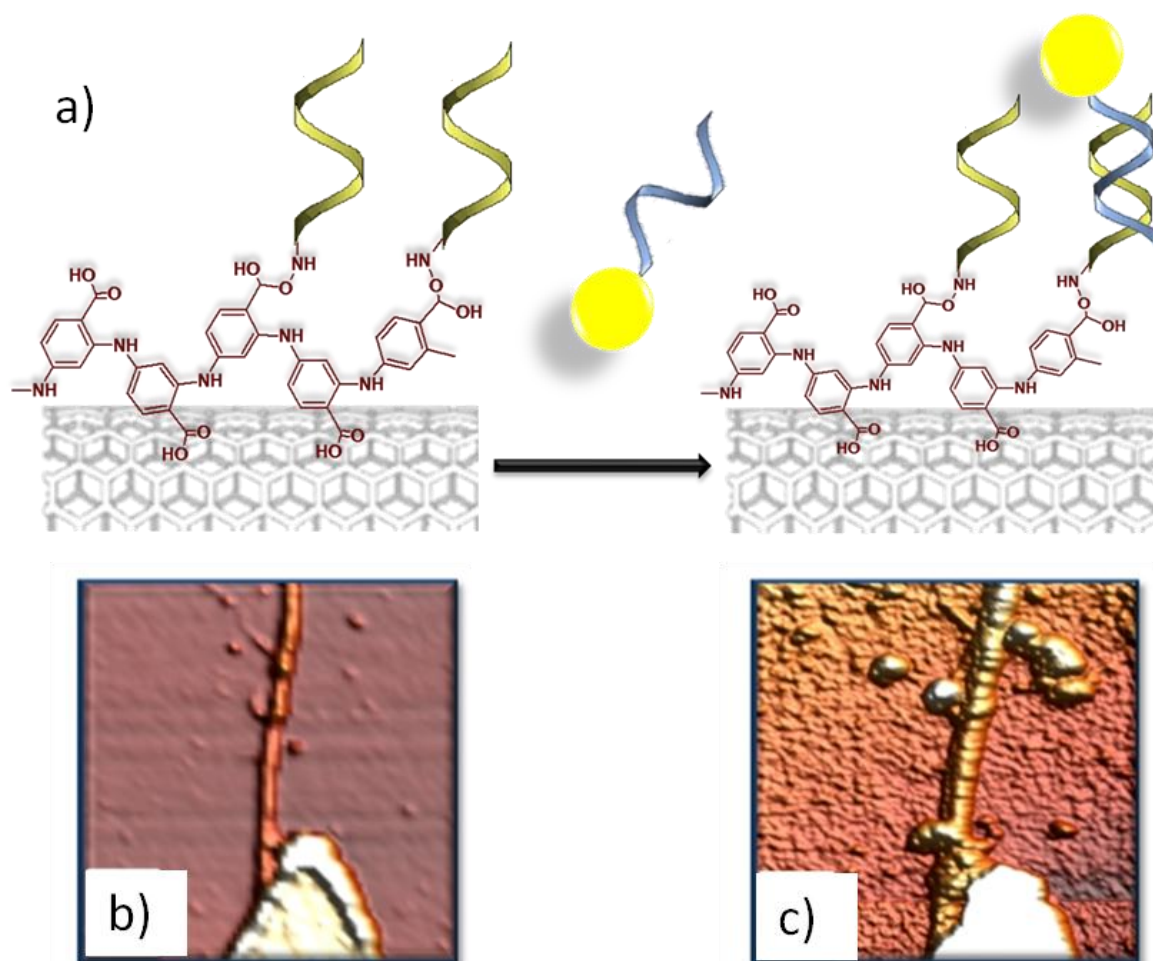


Figure 5.2. The controlled functionalization of the carbon nanotube surface with probe DNA [167].

The height increase upon electropolymerization was estimated to be around 2-3 nm as obtained from AFM images. As it was explained before (*paragraph 4.6*), electrochemical functionalization is site-specific, i.e. only the nanotubes addressed by the electrochemical modification are preferentially decorated with probe DNA. This also ensures that there is no DNA in the vicinity of the nanotube on the chip surface. This is in contrary to situations such as spotting, where the DNA can be immobilized anywhere on the spotting area [66, 177], which could lead to a high level of background noise commonly observed in microarray detection [161, 166]. Furthermore, the negatively charged carboxylic groups are expected to minimize direct non-specific binding of DNA on the nanotube surface via hydrophobic interactions [178].



**Figure 5.3. Evidence for the coupling of probe DNA: schematic of the experiment (a), AFM images taken before (b) and after (c) the coupling protocol [167].**

To prove the applicability of functionalization scheme it was important to show the presence of probe DNA on the surface of the CNT after the coupling. This task was complicated by the ability of nanotubes to weaken the fluorescence of the attached molecules [125, 179]. Therefore, the use of fluorescently labeled oligonucleotides for their detection on the CNT surface was discarded. Instead, for this purpose nanoparticles decorated with complementary DNA were used. In details, the samples with modified CNTs were incubated with splint strand DNA for 40 min in 10mM phosphate buffer (pH 7.0) containing 0.3M NaCl. Following this, the samples were left in a solution of 60 nm nanoparticles decorated with the complementary DNA sequence. AFM images (Figure 5.3) taken before (b) and after (c) the coupling protocol show the presence of attached nanoparticles confirming that the probe DNA was indeed attached to the surface of the CNTs. CNT devices without poly-ABA and with poly-ABA but without probe DNA sequence were used as control. In these cases no nanoparticles were coupled to nanotube surface after the same procedure.

**5.4. Sensing trials.** The sensing trials were performed at varying concentrations of target DNA in 10 mM potassium phosphate buffer containing 0.1M NaCl. Before the sensing trials, several measurements were taken to ensure stable electrical properties of device in buffer solution and also to limit the amount of possible non-specifically adsorbed probe DNA. Impedance measurements were taken twice in target DNA solution, in buffer after washing non-specifically adsorbed DNA and in buffer after DNA melting. For melting of hybridized DNA the microwell was filled with hot water (80°C) several times and washed with buffer afterwards.

The DNA sequences used in this study: probe DNA: 5-ggcctcagtcacactctccgcgc-3, target DNA: 5-gcgcggagagtgtgacgtgaggcc-3, 3-basepair-mismatched DNA: 5-gcgagcagagggtgacgtgaggcc-3.

To prove the suitability of sensing protocol, in other words, to show that the target oligonucleotide is indeed binding to the probe DNA during sensing a target sequence with biotin functionality on 3'-end was used. The functionalization and sensing were carried out in the same way as it was described above. After exposure to biotin labeled target DNA, washing of non-specifically adsorbed DNA with buffer and impedance measurements, the sample was incubated in a solution of 10 nm diameter streptavidin-gold nanoparticles (Sigma).

The AFM and SEM images show the nanoparticles adsorbed to the surface of the CNT after this procedure. This proves that biotin-labeled DNA is present on the surface of the CNT. In control experiments with nanotubes that were not functionalized or were functionalized only with polyABA no nanoparticles were adsorbed to the CNTs, however few nanoparticles were still present on the Si/SiO<sub>2</sub> surface which can be explained as occurring due to non-specific binding. As in the experiment described in *paragraph 5.3* the use of fluorescently labeled biomolecules was also avoided here, due to the ability of CNTs to quench fluorescence [125, 179].



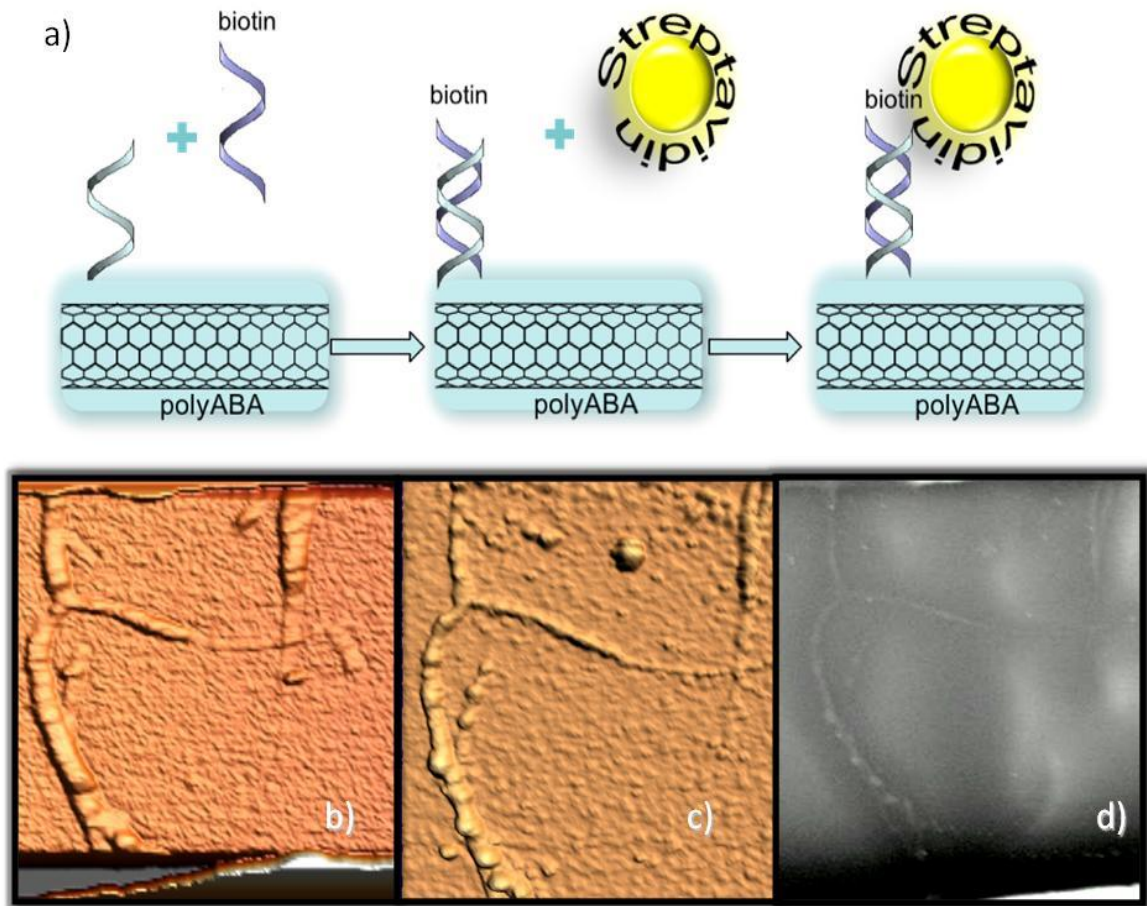
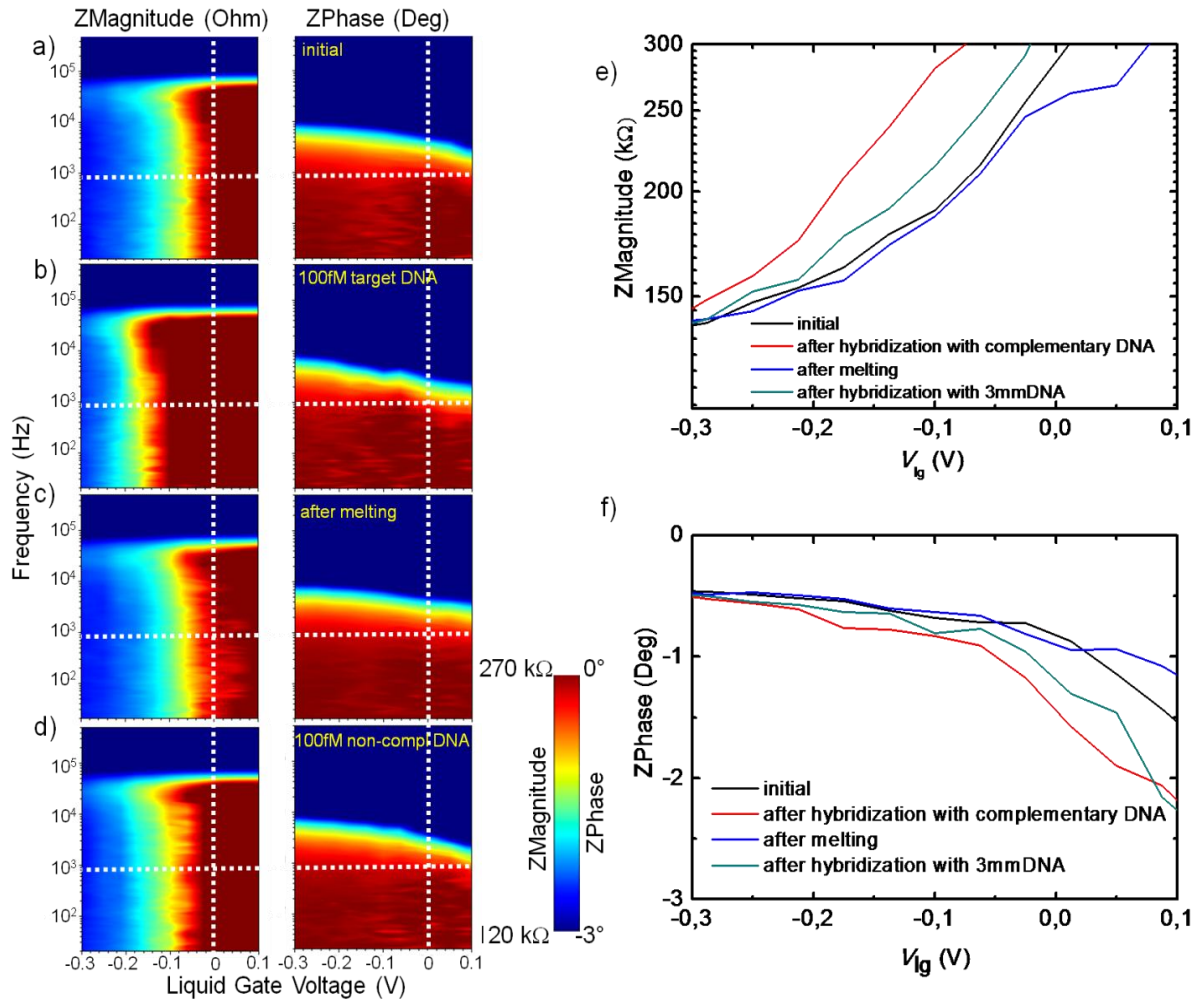


Figure 5.4. Evidence for DNA hybridization during the sensing trial: a) schematic of the experiment, b) AFM image before, c) AFM image after and d) SEM image after the sensing trial. It is apparent that the nanoparticles decorate the nanotube surface confirming that hybridization indeed takes place during the sensing trial. When the nanotubes were not functionalized with probe DNA, no particles were observed after incubation with streptavidin-gold [167].

**5.5. Sensing results.** First the sensor response of the fabricated devices to 100fM of target DNA in buffer will be discussed (Figure 5.5). To perform these measurements the Z-magnitude and Z-phase maps are first recorded in the buffer without any target DNA. The resistance of the device was around 100-500 kOhm, which is dominant at low frequencies. In this range, the device impedance exhibits a low gate modulation. At high frequencies capacitive components arising both from the electrical double layer and the substrate dominate (Figure 5.5a). Upon introduction of complementary target DNA in the microwell, the devices show a clearly different response (Figure 5.5b). It is apparent that the Z-magnitude response shifts to the left along the gate voltage scale. The Z-phase response shows differences



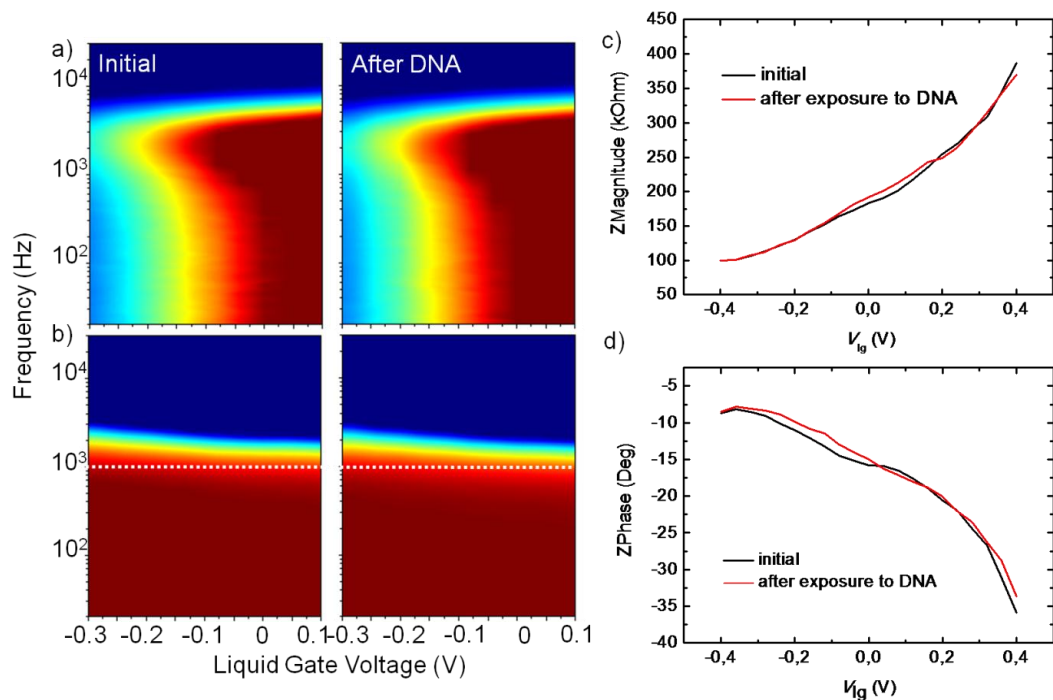
in a frequency range of 1 to 10 kHz. After melting of the hybridized strand and subsequent washing, the initial response is recovered, as shown in Figure 5.5c. In order to confirm that the signal is indeed due to the specific hybridization of the complementary target strand, the sensor response (Figure 5.5d) was measured upon introduction of a 100fm 3-bp-mismatched complementary sequence. The Z-maps remain almost unaffected in this situation showing the threshold voltage shift of less than 20 mV [167].



**Figure 5.5.** Specificity of CNT-DNA-sensors. The plots show magnitude Z-maps (left column) and phase Z-maps (right column) in different solutions. The data was acquired in (a) buffer without target DNA (b) with complementary target DNA (c) after melting and washing and (d) with non-complementary DNA. It is clearly discernible in (b) that for the complementary target DNA the sensor response shifts to the left. This change is minimal in the presence of non-complementary DNA (d). e, f - sensor response (e - magnitude and f - phase) at a fixed frequency (as extracted from the section profiles from the data in figure 5.5 a,b,c,d). The data shows the gate dependence at a frequency of 1kHz. It is apparent that the gate response is reversible after melting and washing.  $V_{lg}$  is liquid gate voltage at the Ag/AgCl reference electrode [167].

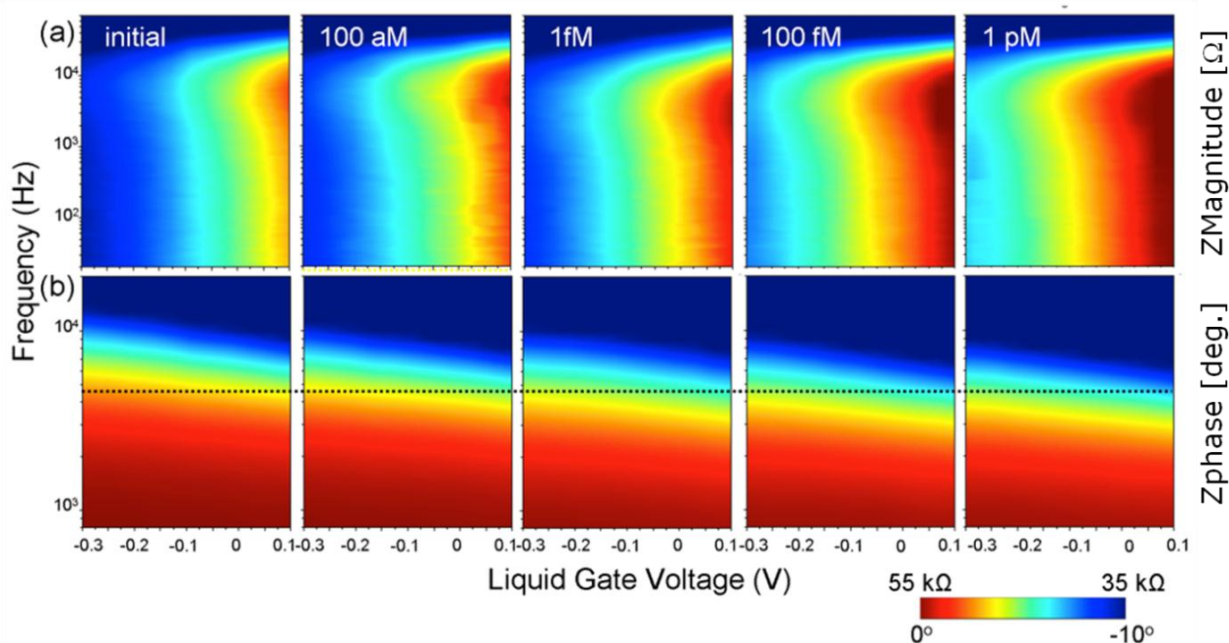
In order to understand the sensor response it is worthwhile looking at the section profiles along the horizontal dotted lines in the Z-maps of Figure 5.5. Figures 5.5e and 5.5f depict these section profiles, which show the gate dependence of the impedance at a frequency of 1 kHz, where the reversibility of the sensor response and the specificity towards the complementary target is clearly discernible. In the presence of the complementary target DNA the magnitude (Figure 5.5e) as well as the phase responses (Figure 5.5f) are shifted to the negative gate voltages. This threshold shift of around 150 mV can be attributed to the accumulation of negative charges on the nanotube surface upon hybridization. The sign of the threshold shift is consistent with data reported on DNA sensors based on nanotube networks [66].

Upon melting and subsequent washing of the sensor surface, the negative charges of the complementary strand are removed and the gate response returns to its initial scenario. The gate response for the 3-bp-mismatched DNA shows only a comparatively negligible shift (less than 20 mV) to the left signifying a much lower degree of hybridization as is normally expected for mismatched DNA sequences [161].



**Figure 5.6. Control experiment with non-functionalized CNTs: magnitude Z-maps (a) and phase Z-maps (b) in different solutions. No change in transport characteristics was observed after exposure to 100 fM DNA solution. Curves showing the magnitude (c) and phase (d) of impedance during different stages of the sensing trial. In this case, the nanotubes were not functionalized with the probe DNA. It is apparent that the transport characteristics do not show any difference with or without the complementary target DNA [167].**

Nanotubes that were not functionalized with ABA with or without subsequent coupling of probe DNA did not show any significant response towards complementary sequence even at the highest concentration used here (Figure 5.6) [167].



**Figure 5.7. Attomolar detection limit of CNT-DNA-sensors.** (a) Magnitude Z-map and (b) Phase Z-map for various concentrations of complementary target DNA. After every exposure of the sensor to target DNA, the hybrids were melted in order to obtain the initial state as shown in Figure 5.5 (a-c). It is apparent that even at a concentration of 100aM (corresponding to around 1800 molecules in our 30  $\mu$ L droplet), the sensor response can be unambiguously discerned, signifying the ultrahigh sensitivity of CNT impedance biosensors [167].

**5.6. Sensitivity and detection limit.** The use of high frequency detection ensures a very low noise of the electrical measurements. The Ag/AgCl reference electrode provides for excellent stability. This enables the repetitive use of the same sensor for a series of different DNA concentrations with minimal drifts. Figure 5.7 collects the concentration dependence of the sensor response for one of the devices, displaying the magnitude Z-maps (Figure 5.7a) and phase Z-maps (Figure 5.7b) for the various concentrations of complementary target DNA. Section profiles as extracted from the maps in Figure 5.7a are collected in Figure 5.8a. It is apparent that the gate response shifts to increasing negative gate voltages with increasing concentration of target DNA [167].

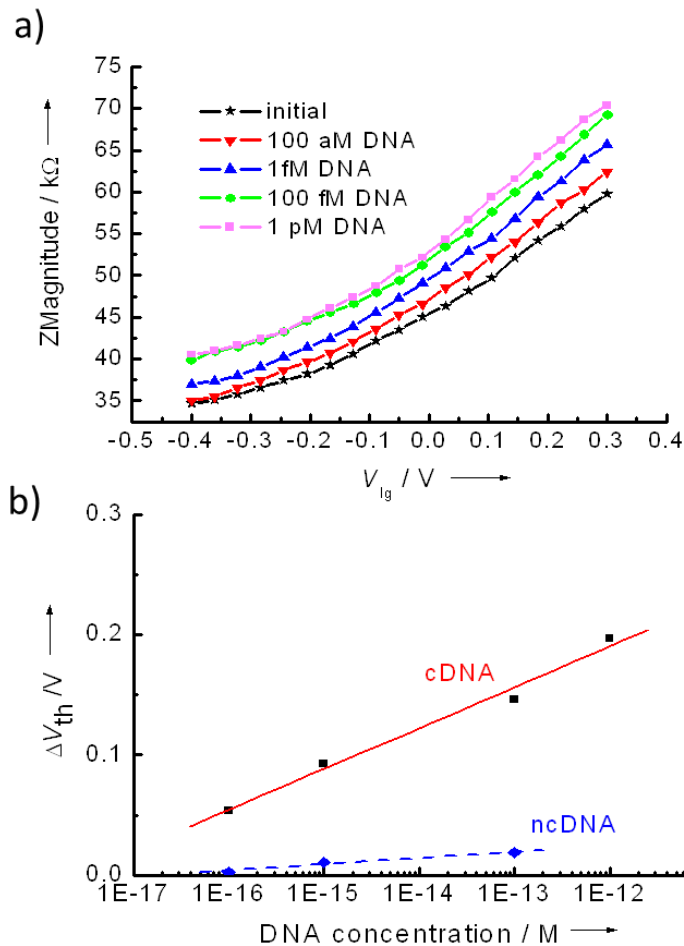
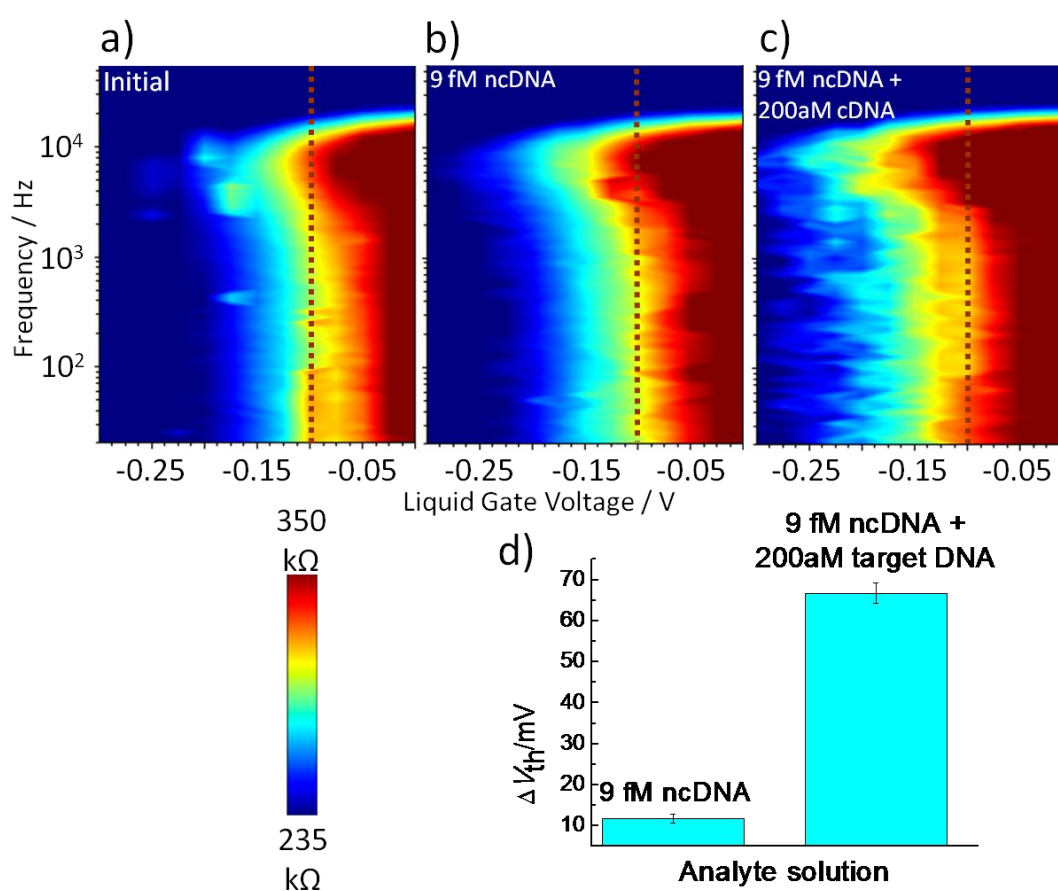


Figure 5.8. Sensitivity of CNT-DNA-sensors: (a) Gate dependence of the magnitude of impedance at a frequency of 1 kHz in buffer (black line) and for varying concentrations of complementary target DNA in buffer (extracted from Figure 5.7a). The threshold voltage shifts to more negative voltages with increasing concentrations of DNA. (b) Calibration curve showing the threshold shift as a function of complementary (cDNA) and non-complementary (ncDNA) concentrations showing a linear response over a broad concentration range [167].

Figure 5.8b) presents the calibration curve where the shift in threshold voltage is plotted as a function of target DNA concentration. It is apparent that the sensor response is linear over a broad concentration range and a concentration as low as 100 aM can be unambiguously detected. The threshold voltage shift caused by non-specific binding of non-complementary DNA (Figure 5.8b - blue line) is significantly lower for all tested concentrations. This corresponds to around 1800 molecules of target DNA in our 30  $\mu$ L sample droplet in the microwell. This is the lowest detectable concentration that has been reported using any label-free or mediator-free direct detection technique [167].

**5.7. Detection in complex media.** In order to evaluate the use of presented nanobiosensors in a realistic application scenario, we have validated our analytical strategy towards the specificity of DNA differentiation at ultralow concentrations. For this purpose, we have taken a heterogeneous mixture of three different DNA sequences (ncDNA – noncomplementary to the probe sequence) each at a concentration of 3 fM. To this we add 200 aM of complementary DNA (cDNA) giving a total DNA concentration of 9.2 fM. The amount of cDNA we are aiming to detect corresponds to just around 2% of the total DNA present in the solution mixture. As a control we use the ncDNA mixture without the cDNA at the same total DNA concentration. Figure 5.9 collects the Z-maps measured in various analyte solutions [167].



**Figure 5.9. Attomolar target differentiation in a heterogeneous DNA mixture.** Magnitude Z-maps (a) in buffer (b) in buffer with a heterogeneous mixture of three different non-complementary sequences (ncDNA) each at 3 fM giving a total DNA concentration of 9 fM (denoted as 1) and (c) in solution 1 with added 200 aM of complementary target DNA (denoted as 2). (d) Sensor signal (threshold shift) for solutions 1 and 2. In 2, the complementary target amounts to just 2% of total DNA. The CNT-DNA-sensor is capable of differentiating this small amount from the high 9 fM non-complementary background, signifying a high selectivity coupled to an ultralow detection limit [167].

It is apparent that the sensor response shifts clearly towards negative gate voltages, similar to Figure 5.5. A significant shift is only seen if the buffer contained the complementary target (Figure 5.9c). This is further clear from the threshold voltage shift collected in Figure 5.9d for the heterogeneous sample with and without the complementary DNA. It is apparent that 200 aM of target DNA comprising just 2% of the heterogeneous sample is able to bring in a significant threshold shift of around 65mV. On the other hand, the 9fM ncDNA control solution without the cDNA shows only a shift of around 12 mV, which is below the 20 mV occurring due to non-specific interactions (as mentioned earlier – *paragraph 5.5*) [167].

**5.8. Conclusions.** The power of presented nanosensor is apparent from the above described measurements and soars hopes of extending the technique to directly detect low quantities of DNA from realistic biological samples such as blood (after preprocessing), thereby showing promise for use in a clinical setting without amplification or labeling steps. As mentioned earlier, the ability to detect few copies of the DNA is made possible primarily by ensuring that the high surface-to-volume ratio is guaranteed through appropriate passivation of the electrodes, leaving exclusively the nanotube surface as the active element. Furthermore, the low resistance, the stability gained by using the impedance measurement and the site-specific electrochemical functionalization route have been crucial in achieving this ultra-low detection limit.



## 6. Towards graphene-based electrical sensors

**6.1. Introduction.** In spite of the constant efforts from the researchers, the integration of graphene into devices still remains challenging. Mechanical exfoliation of graphene from graphite crystal followed by e-beam lithography is the commonly used technique for the preparation of contacted graphene flakes [68, 180]. This approach is widely used in the research laboratories, but has a number of limitations, for example, critical manual operation steps which cannot be automated. Therefore this approach is unsuitable for device production at the industrial level. Moreover the chips produced in this manner may have differing geometry and position of contacts and flakes. This can complicate manufacturing and operation of the devices at later stages. Researchers are trying to overcome these limitations by using chemical vapor deposition (CVD) grown graphene as a starting material for device fabrication [181-183].

Solution-based approaches for the preparation of carbon-based devices offer the possibility for large-scale processing without using bulky equipment or operation at very high-temperatures. Graphene suspension can be obtained from graphite through intercalation using organic solvents, surfactants, etc [184]. Another solution-based approach is to produce an aqueous solution of oxidized form of graphene (Graphene oxide - GO) which is hydrophilic due to the presence of polar groups, such as carboxyl, hydroxyl and epoxygroups [104, 185]. Graphene oxide is insulating but it can be made conductive after a reduction process [103, 187]. The resulting material is called reduced graphene oxide (RGO). To fabricate RGO devices using this kind of suspension drop casting of the solution on the substrate with subsequent contacting of the flakes of interest can be used. However this approach does not improve the scalability of the fabrication. Therefore pre-patterning of the surface and dielectrophoresis have been used to position the flakes at the desired locations [186 - 190].

**6.2. DEP trapping of graphene.** Several approaches for solution based graphene devices fabrication were investigated here. Prior to the fabrication of graphene devices using dielectrophoresis a solution of graphene or graphene oxide must be prepared. Solutions containing graphene flakes were obtained by 10 min sonication of graphite powder in dimethylformamide (DMF) and Triton X-100. Large particles of graphite were removed using centrifugation (3000 rpm, 5 min). AFM imaging revealed the extreme inhomogeneity of resulting suspensions. Dropcasting of the solution on a substrate and AFM measurements showed the presence of large enough single or few-layer graphene flakes (Figure 6.1a). However, during the dielectrophoretic trapping (500kHz, 1V, 10 sec) which was performed to position graphene from these solutions at the electrodes, along with the single or few-layer graphene flakes, large graphite particles were trapped (Figure 6.1 b). Further attempts to purify the solution led to a decrease of the graphene flakes size, which complicated bridging of the gap between the electrodes (Figure 6.1 c). Therefore other methods were considered as alternative solution-based approaches for the fabrication of graphene-devices.

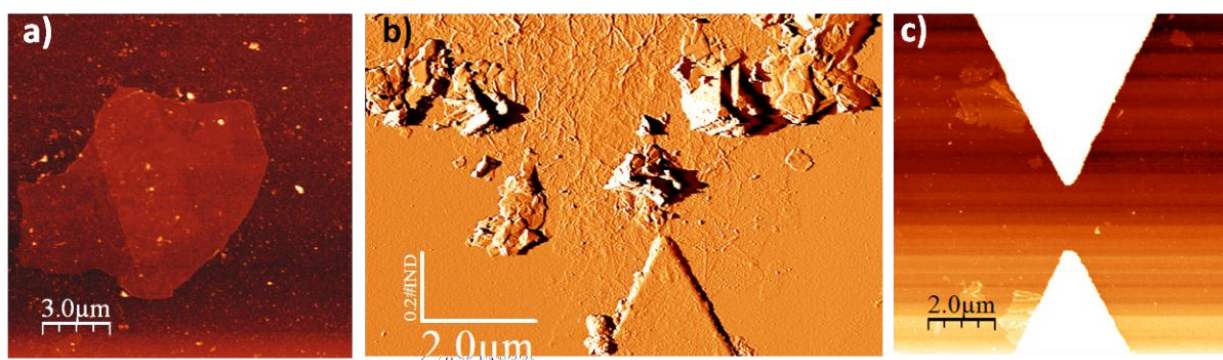
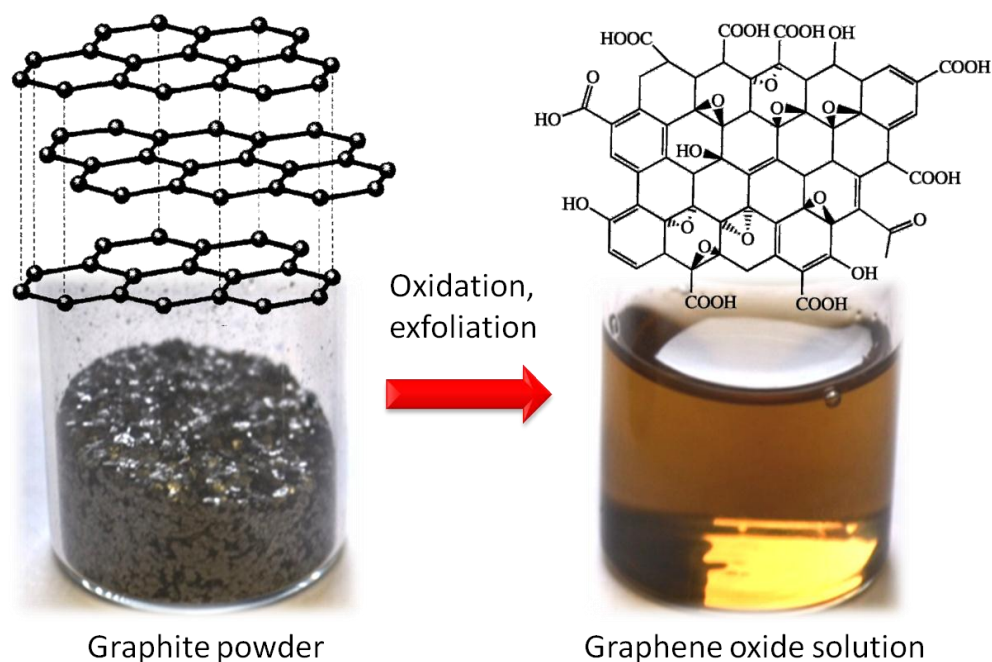


Figure 6.1. Liquid phase exfoliation of graphene from graphite. a - AFM images of few-layer graphene flake obtained by exfoliation of graphite in DMF, b - graphene flakes dielectrophoretically trapped together with large graphite particles, c - individual graphene flakes trapped at electrodes using DEP.

**6.3. DEP trapping of graphene oxide.** As it was discussed above, an alternative route was to prepare graphene oxide solution, followed by DEP trapping and reduction. GO solution was prepared using Hummers method [191]. 1g of graphite powder (200 $\mu$ m grain size) was added under constant stirring to the flask with 23 ml of 98% sulfuric acid. 0.5 g of NaNO<sub>3</sub> was added and the speed of stirring was increased. After that 3 g of KMnO<sub>4</sub> was added carefully. The above mentioned procedures were performed in the ice bath with the temperature control of the



solution. The temperature of the mixture was kept below 20°C. Upon removing the iced water bath, the temperature was increased to 35°C in 30 minutes, leaving a grey paste. 46 ml of deionized water was then added and the temperature was increased to 98°C where it was maintained for 15 minutes. Then 70 ml of water was poured into the mixture. 3% H<sub>2</sub>O<sub>2</sub> was added slowly until the solution turned bright yellow. The suspension was vacuum filtered resulting in a yellow brown filtrate which was washed several times with water. The exfoliation was performed by overnight shaking. The obtained graphite oxide suspension in water was centrifuged at 3000 rpm for 3 minutes to remove large particles.



**Figure 6.2. Preparation of graphene oxide solution**

Before DEP the GO solution was diluted 1:10 with water and sonicated for 5 seconds. DEP trapping was performed by applying 500 kHz, 10 V AC-voltage for 15 second. Afterwards the device was washed with water and isopropanol and blow-dried in nitrogen flow. Then GO was thermally reduced in the furnace at 350°C for 1 hour in an inert atmosphere (argon). The devices prepared in this manner showed resistances in the range of 200kOhm to 2 MOhm. Due to incomplete reduction the resistance of the obtained devices is higher than in the case of pristine graphene but the fabrication is rather simple and scalable. Another advantage of the GO-based approach for fabrication of devices for biosensing is that the remaining oxygen containing functionalities can be used for chemical functionalization of RGO.

The RGO devices can be further improved for the use in liquids by passivating the metal electrodes as it was done for CNT devices [130, 167]. An example of device prepared in this manner is shown in Figure 6.3

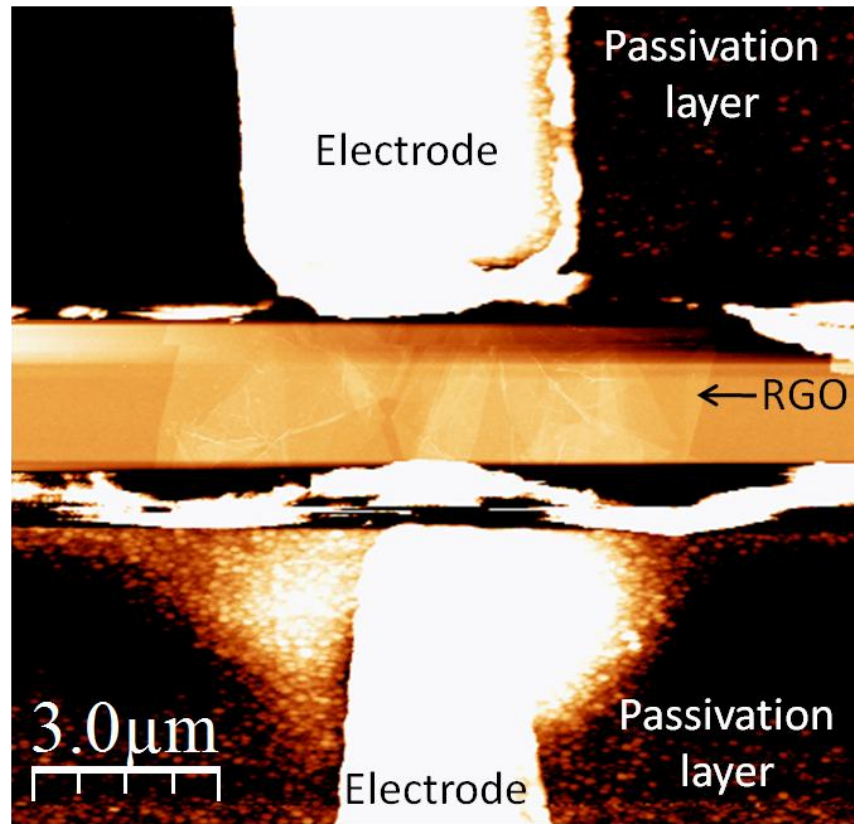


Figure 6.3. An RGO device with passivated contacts for biosensing applications.

The devices were assembled on a chip-carrier for impedance measurements in a liquid gated configuration. The reservoir for liquid was made using PDMS plate as it was described previously (*paragraph 5.2*). Impedance measurements were performed for the range of the gate voltages from -0.4 to 0.4 V. Figure 6.4. shows the typical 2D-ZMagnitude-map of RGO device (a) and its gate dependence at 1000 Hz (b) [130, 167].

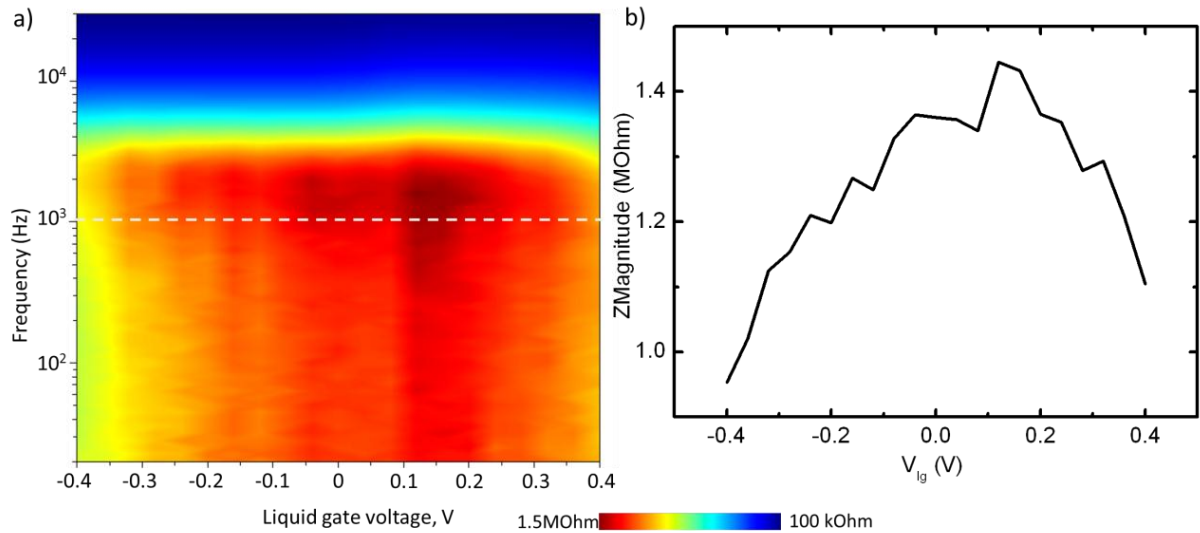


Figure 6.4. Liquid gating of RGO device. a) 2D-map of the magnitude of impedance, b) gate dependence of the device at a fixed frequency of 1000 Hz.

**6.4. Chemical anchoring of graphene oxide for preparation of the devices.** DEP of GO often results in the trapping of several flakes. The devices prepared in this manner therefore are not as reproducible as they are required to be. Moreover, the flakes often have numerous wrinkles and folds. Therefore, an alternative chemical route for positioning GO flakes at the desired location was proposed. It utilizes the presence of the various reactive functional groups in graphene oxide [103]. These functional groups can be used as hooks for anchoring the graphene flakes on the electrode surface previously functionalized with appropriate reactive groups. Selective modification of the electrodes can be achieved using electrochemical reactions. In this case, relevant functional groups for coupling the functionalities on graphene oxide will be selectively present on the surface of the electrodes [192].

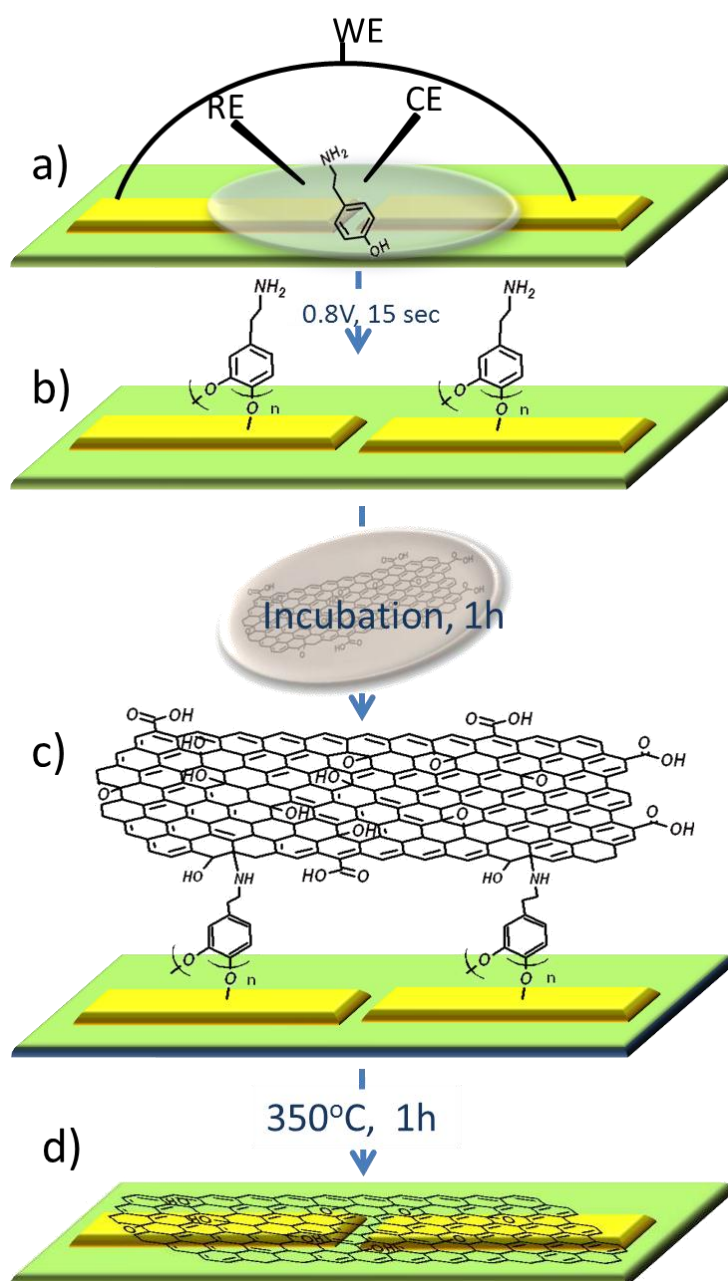


Figure 6.5. Scheme of the chemical anchoring protocol. (a) electrochemical functionalization of Pt electrodes with tyramine leading to (b) a coating of polytyramine on the electrode surface; (c) incubation of the chip in a GO solution results in the coupling of GO flakes to the polytyramine layer; (d) annealing in argon at 350°C leads to the removal of the polytyramine layer and the reduction of most of the oxygen containing groups. WE - working electrode, CE - counter electrode, RE - reference electrode [192].

The platinum electrodes were modified by oxidative polymerization of tyramine with the formation of a thin film of polytyramine (pTy) which has a high density of free aminogroups (Figure 6.5 a,b) [193]. Previously the immobilization of biomolecules on microelectrodes with the help of electropolymerized tyramine was

successfully demonstrated [194, 195]. Amine-containing polymers have been shown to be effective for cross-linking of graphene oxide sheets via amino-epoxy coupling [196]. The polymerization of tyramine was carried out by applying 0.8V for 15 sec at the platinum electrodes pre-written on the silicon substrates which were immersed into 10mM tyramine solution in 10mM H<sub>2</sub>SO<sub>4</sub> using CompactStat potentiostat (Ivium technologies). Platinum wires were used as counter and pseudo-reference electrodes. After electrochemical deposition of polytyramine the substrate was washed with ethanol and water and dried in nitrogen flow. For the coupling of pTy with graphene oxide we immersed the substrate in a suspension of graphene oxide and left it for 1h in a shaker. During the incubation of the sample in GO solution pTy reacts with graphene oxide and acts as an anchor on the electrode surface (Figure 6.5c). Afterwards, the substrate was taken out of the GO solution, immediately washed with water and isopropanol and dried in nitrogen flow. After that the sample was thermally annealed in inert atmosphere (argon) at 350°C for 1 hour (Figure 6.5d). After cooling down to the room temperature the sample was washed with isopropanol and dried [192].

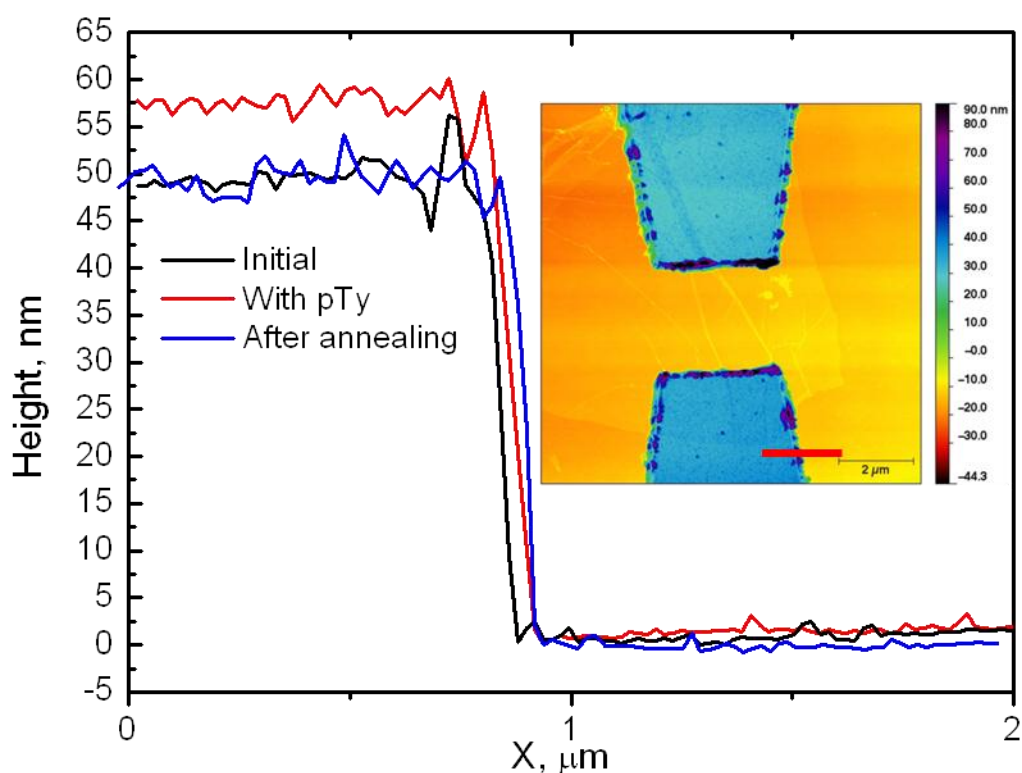
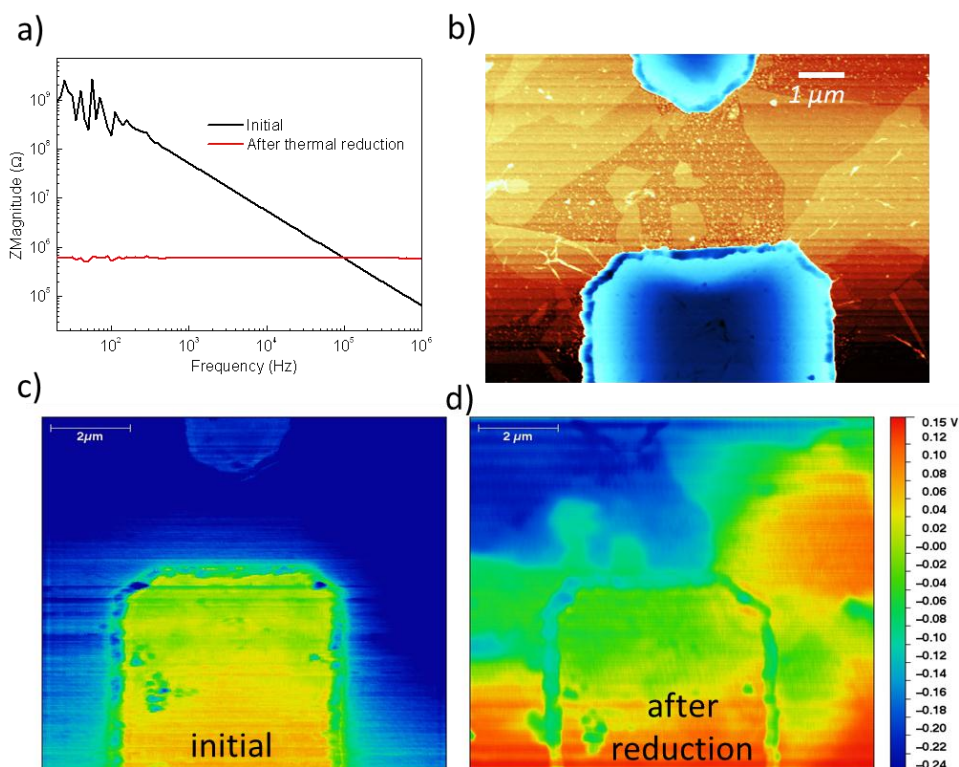


Figure 6.6. Surface characterization by AFM. Inset - AFM image of a typical sample obtained at the end of the chemical anchoring protocol. The electrode gap and the immobilized graphene oxide flake are visible here. Main plot - Height profiles (along the red line in the AFM image) taken at a GO-free region on the electrode - as extracted from AFM images recorded at various stages of the anchoring procedure. It is apparent that the pTy film after electrodeposition is 5 to 8 nm thick (red curve). Annealing leads to the reduction in electrode height (blue curve) to the initial value signifying the removal of the pTy film [192].



The AFM images of the electrodes on the substrate were taken before and after pTy formation as well as after incubation in GO solution and after annealing. Figure 6.6 (inset) shows the AFM image of the device prepared using our approach. The coupling of graphene oxide to the modified electrodes was clearly observed while there was no graphene oxide on the electrodes that were not electrochemically treated. As it is seen from height profiles (Figure 6.6) the layer of polytyramine that was formed on the surface of the electrodes during electrochemical functionalization had a thickness of around 5-8 nm. After the thermal annealing step, the height of the electrodes decreased to the initial value which indicated the removal of the polymer layer from the surface of the electrodes not covered with GO. Similar height profile analysis of the electrode regions with graphene oxide on top suggested a decrease of the polymer layer thickness below the graphene oxide sheet [192].

During thermal annealing in argon the reduction of graphene oxide also takes place and the device becomes conducting [197]. This was observed by impedance spectroscopy measurements (Figure 6.7a). The resistance improvement confirmed the removal of a majority of oxygen functionalities from the GO surface [192].

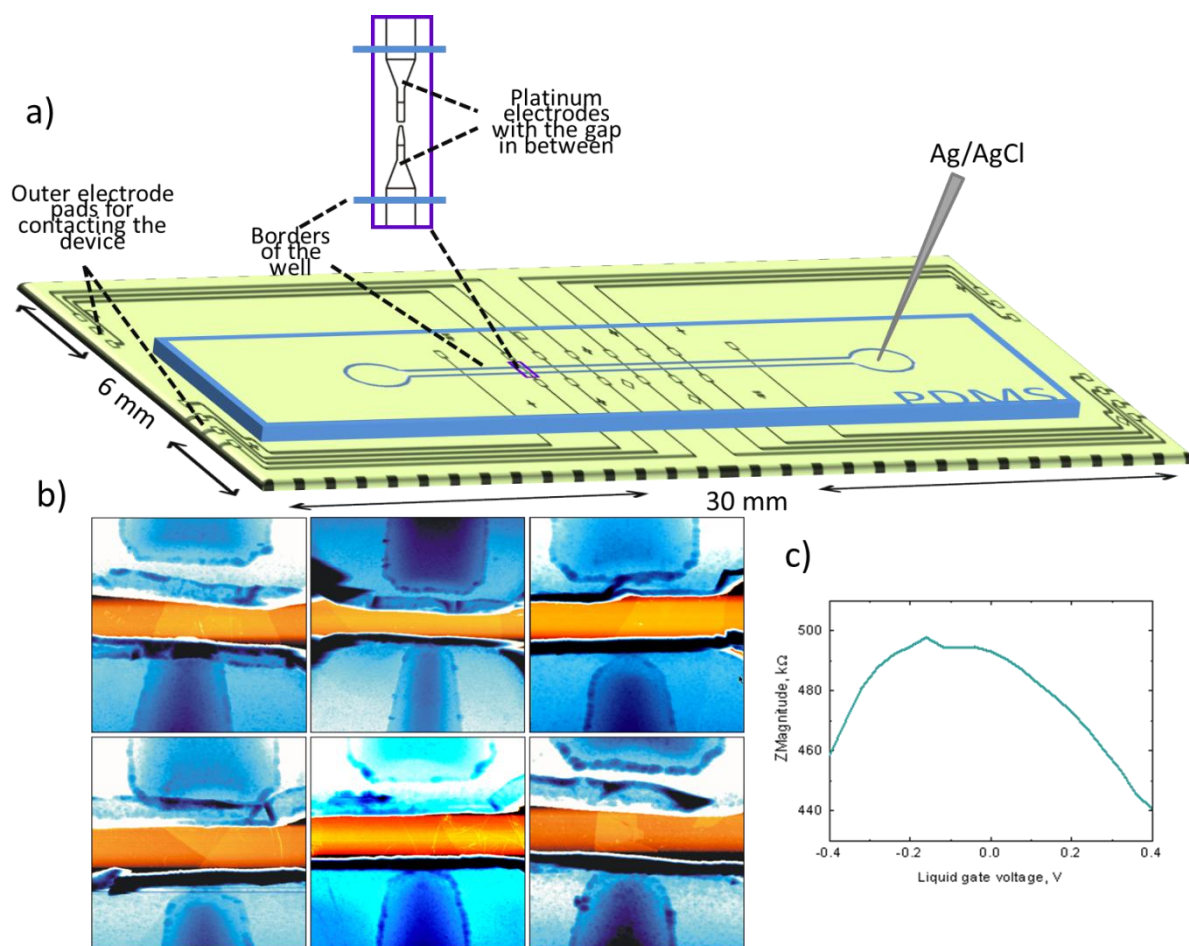


**Figure 6.7. Electronic properties of the anchored GO devices. (a) Impedance spectrum of a typical anchored GO device before and after the thermal annealing step (350°C, Argon, 1h), AC amplitude 100 mV. (b) Tapping mode AFM height image of the same device. (c) and (d) Surface potential maps of the same device before (c) and after (d) the thermal annealing step. The tip was grounded while a fixed bias (0.5 V for (c) and 0.25**

**V for (d) was applied across the gap. It can be inferred that the annealing step improves the resistance and the electronic properties of the anchored flakes [192].**

In order to gather further support for the improvement in transport upon annealing, the devices were investigated using Kelvin Probe Force Microscopy [198, 199]. The resulting surface potential maps are depicted in Figures 6.7c and 6.7d. In the former case, there is no conducting pathway measurable in the device before annealing, consistent with its insulating behavior. By contrast, after annealing the current carrying flake can be clearly identified in Figure 6.7d, analogous to recent results on individually contacted RGO flakes [200]. This leads us to conclude that the annealing procedure serves as a single processing step that results in a graphene-like backbone of the GO flakes, while simultaneously removing the polytyramine layer at the contacts [192].

**6.5. Up-scaling the fabrication of RGO devices.** The advantage of the offered approach for device fabrication is the possibility to scale-up the process. Towards this goal, we utilized a specialized layout as shown in Figure 6.8a. The substrates comprised of 6 mm x 30 mm Si/SiO<sub>2</sub> chips allowing for long leads. The long leads enable us to perform electronic transport measurements in liquids as is discussed later. Each chip comprises of 6 electrode gaps. All the 12 electrodes are initially wire-bonded in the outer region of the substrate to form a parallel connection of 12 electrodes. In this manner all the electrodes can be decorated electrochemically with polytyramine in a single step. A drop of tyramine solution was placed in the region of the gaps. Pt wires used as counter and reference electrodes were inserted into this drop. The voltage applied to one of the pads of connected electrodes led to an electrochemical reaction on all of them at the same time. Placing such a sample into the GO solution led to the coupling of graphene oxide to all of the electrodes on the surface. The contacts of the devices were passivated using an appropriate mask and a mask aligner MA-6 (SüssMicrotech). The resulting passivated devices are shown in Figure 6.8b. Liquid gate dependence was measured in water using Ag/AgCl reference electrode to apply the gate voltage and PDMS plate with the hole to form a well. The contacts to the electrodes were located outside PDMS well. The devices showed resistances in a range of 100kOhm – 1 MOhm which is comparable to the reduced graphene oxide devices obtained by dielectrophoretic trapping of GO followed by reduction. Liquid gate dependence of a typical device is shown in Figure 6.8c [192].

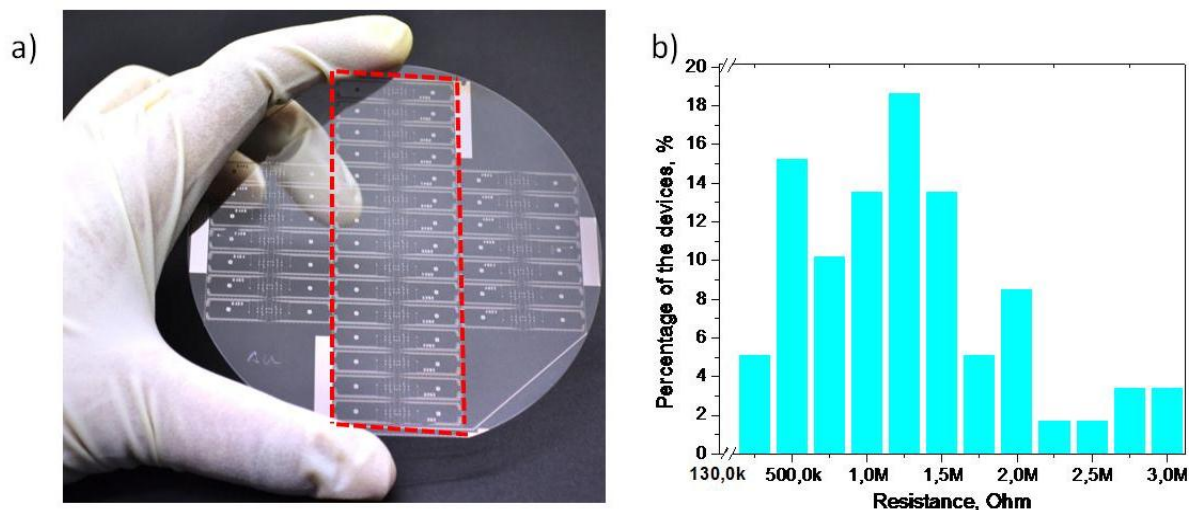


**Figure 6.8. Scaling up chemical anchoring.** (a) Device layout showing 6 electrode gaps. Initially, the 12 electrodes are linked externally through wire bonding. For liquid gating measurements, a liquid channel connected by two circular wells is created with the help of a polydimethylsiloxane (PDMS) layer. The inset shows a close-up of the electrode gap and the relative extent of the liquid channel. An Ag/AgCl reference electrode inserted in the channel serves as the gate electrode. (b) AFM images of the 6 gap positions (from one chip) at the end of the chemical anchoring protocol. In addition to the steps shown in Figure 6.5, the electrodes are passivated with SiO<sub>2</sub> using a separate photo mask. (c) Liquid gate dependence of resistance of one of the RGO devices, showing the typical ambipolar behavior [192].

From the foregoing discussions it is apparent that the chemical anchoring protocol serves as a versatile route to obtain RGO field-effect devices with a high throughput. The real advantage of such a technique in an industrial scenario requires the demonstration of such a protocol at the wafer scale. Furthermore, the feasibility of this protocol for other substrates, such as glass or polymers, is also an important factor. To evaluate this, we have fabricated electrodes on 4" glass wafers using photolithography as shown in Figure 6.9a. The wafer is segmented into chips, each of which is laid out as shown in Figure 6.8a. The layout is designed in such a way that the electrodes of the chips in the region marked with the red dashed line (Figure



6.9a) are all connected to each other through the large common pads. After carrying out the chemical anchoring protocol, the wafer is diced whereby these electrode connections are lost resulting in standalone chips with 6 individual gaps each. For the wafer in Figure 6.9a, 77 working devices (out of 90 electrode gaps) were obtained signifying a yield of 86%. A histogram of the resistances from these 77 devices is presented in Figure 6.9 b, with resistances in the range of 200 kOhm to 3 MOhm [192].



**Figure 6.9.** Wafer scale RGO devices with high yield. (a) A photograph of a 4" glass wafer with photolithographically patterned electrodes. The electrodes in the red dashed region are connected to each other through the large pads in the upper left and lower bottom. These large pads are used to deposit a polytyramine layer on all the electrodes in a single step. There are 15 chips in red dashed region, each chip containing six electrode gaps with an electrode layout as shown in Figure 6.8a. (b) Histogram of resistances of 77 out of the 90 electrode gaps at the end of the chemical anchoring protocol. 13 devices showed very high resistances [192].

These results underline the scalability of this approach demonstrating a capability to obtain RGO devices at a high yield. The devices prepared on wafers showed higher resistances on the average in comparison to those observed on individual samples. This could be improved by optimizing the conditions of electropolymerization and thermal reduction for the substrate of interest.

It is worth mentioning here that this approach was also successfully implemented on substrates with differing electrode geometries, as well as on flexible kapton foils (Figure 6.10).

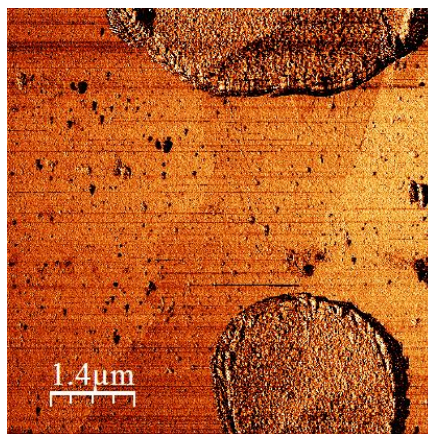


Figure 6.10. AFM image (phase mode) of graphene oxide coupled to electrochemically modified electrodes on flexible kapton substrate [192].

The method was also tested using commercially available graphene oxide (Graphene Supermarket, USA). It is noteworthy to say that the bridging of the gap between electrodes is limited by the relative sizes of the gap and graphene oxide flakes. It is understood that if flakes are smaller than the gap, coupling will still take place but the gap will not be bridged. Therefore, the yield of the preparation using commercially available graphene oxide was worse. However, by using the samples with smaller gap sizes it is still possible to improve the yield of fabrication in case when flakes are small (Figure 6.11) [192].

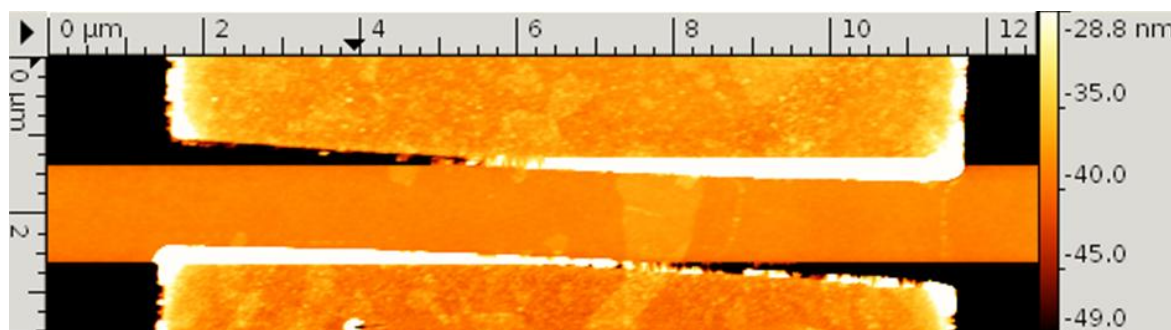


Figure 6.11. Commercially available graphene oxide (Graphene supermarket, USA) coupled to the electrodes [192].

**6.6. Conclusions.** In conclusion, several routes for the fabrication of graphene-based devices were investigated. Two main approaches were utilized for scalable fabrication - DEP trapping and chemical coupling of GO to the electrodes surface. These approaches allow preparing arrays of RGO devices in a short time without the use of serial techniques such as electron beam lithography. A field-effect was observed on these devices when operating in liquids using an electrochemical

gate. This is important for the electrical biosensing applications. The ability to realize wafer-scale RGO devices on arbitrary substrates is an important step towards their widespread use in a number of fields [192].

# Summary and outlook

---

**7.1. CNT based electrical sensors.** In this thesis the electrical biosensing platforms based on electrochemically functionalized carbon nanotubes are presented. The important aspect of this work was the optimization of the sensor fabrication procedure. The device fabrication steps were optimized to ensure high throughput of this process. Thus, Si/SiO<sub>2</sub> samples with photolithographically written Pt electrodes were used as substrates for the fabrication of nanosensors. The scalable assembly of CNTs into devices was achieved by dielectrophoretic trapping of the nanotubes from their dispersion. This ensures low-cost, reproducibility and possibility of automation of the CNT sensor fabrication. Sensitivity and robustness of the sensors was improved considerably with the help of passivation of the metal contacts with insulating layer. Electrochemical functionalization strategy, used in this work, provided site-specific modification of the nanotubes. Site-specificity of functionalization is one of the crucial requirements for achieving high sensitivity of the sensor. It also makes it possible to fabricate on the same chip several nanosensors designed to detect different analytes. Multiplex detection of several analytes can be realized in this manner. This will pave way for lab-on-a-chip systems that can be tailored for specific applications in point-of-care diagnostics.

In this work biosensing is based on the measurement of the impedance of the modified CNTs at different liquid-gate voltages. Impedance measurements in field-effect configuration of the device ensure low signal to noise. Glucose detection with boronic acid functionalized CNT devices helped to reveal mechanisms of sensing using the presented biosensing configuration. Finally, very high sensitivity and selectivity of the designed nanosensors were achieved in the case of DNA detection. Nevertheless, demonstrated sensing platform leaves a lot of room for further development and improvement.

For example, in the case of DNA sensing, the experiments were conducted with synthetic oligonucleotides. It will be important to demonstrate this high sensitivity with real nucleic acid sequences obtained from biological samples. Possible analytes for such tests can be microRNA (miRNA) molecules. MicroRNAs are small (18 to 25 nucleotides) regulatory genes [201]. They can inhibit the

expression of target protein-coding gene. The suppression of the protein expression occurs due to the complementarity of the miRNA to a part of one or more messenger RNAs [202]. This inhibits translation of the protein. It was discovered that miRNAs circulate in blood serum. The concentration of some of the circulating miRNA changes depending on the pathological states of the organism, like sepsis, autoimmune diseases, cardiac diseases and cancer [203-207]. Taking into account the stability of miRNA they can serve as potential biomarkers of various diseases [204, 208, 209]. The low concentration of miRNAs in serum requires highly sensitive and selective methods for their quantification. The biosensing platform described in this thesis may potentially be applied for this purpose.

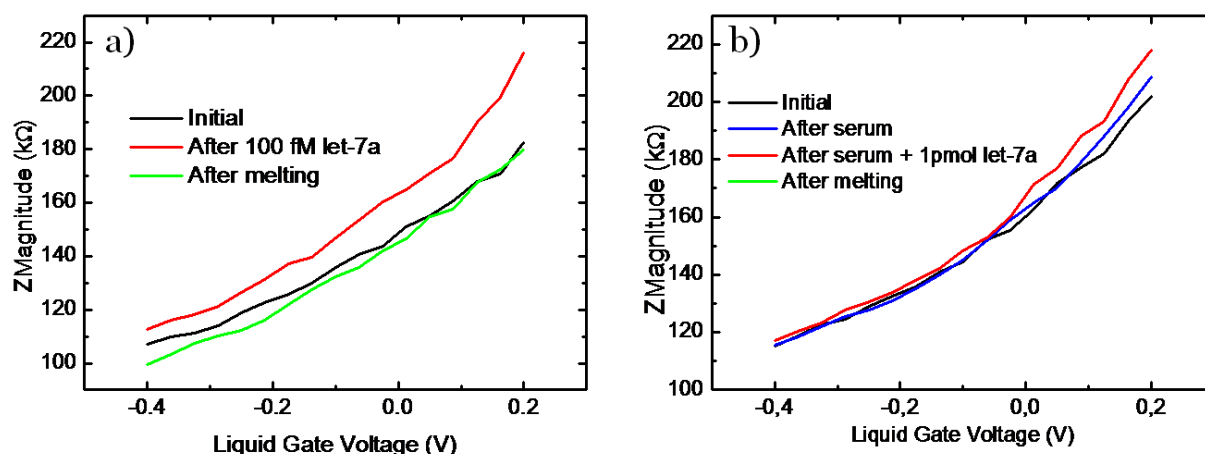


Figure 7.1. Sensing of let-7a-miRNA in buffer (a) and serum (b).

Towards this goal the first miRNA sensing trials were conducted. We used let-7a-miRNA as an analyte and a DNA sequence complementary to it as a receptor. The fabrication of the sensor, functionalization of CNTs and electrical detection were performed in the same way as in the case of the DNA sensor. The results obtained during the sensing of let-7a-miRNA in buffer were similar to that of DNA detection (Figure 7.1 a). The gate characteristic shifts to the left after the CNT-device was exposed to a 100 fM let-7a solution.

However, the detection of miRNA in serum was not straightforward. miRNA added to serum diluted with working buffer (1:10) did not cause significant changes in the gate dependence of the sensor (Figure 7.1 b). This can be explained by the presence of the proteins and metabolites in serum which may affect the access of the target RNA to the carbon nanotube. The AFM images of the device taken after sensing trials in serum showed the contamination of SiO<sub>2</sub> and CNT surface (Figure 7.2). It is clear that the sensing protocol should be optimized for the direct measurement of analytes from the serum. One possibility to improve this is to



perform a pretreatment of the serum samples, like liquid-liquid or solid-phase extraction, protein precipitation, etc [210].

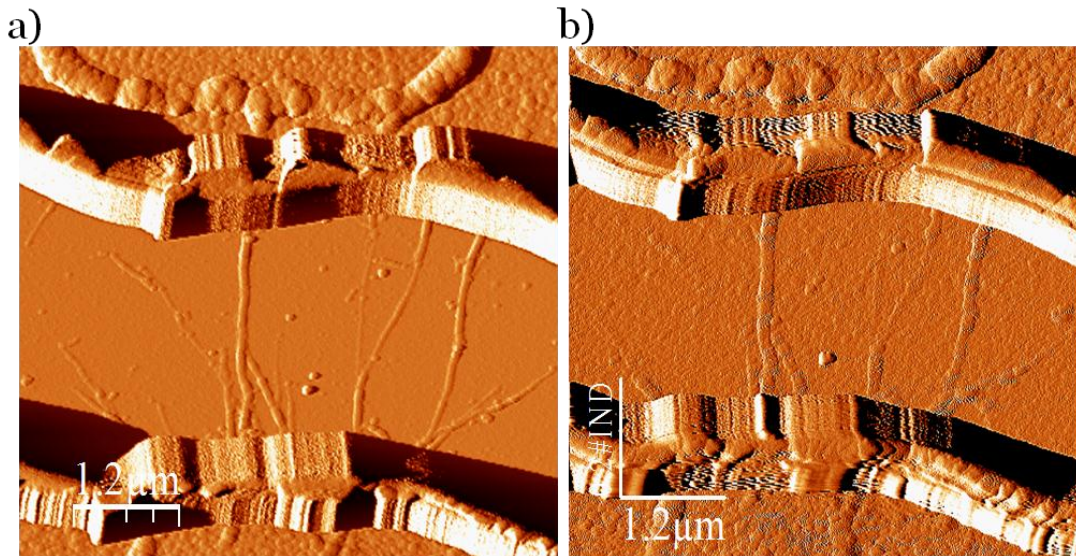


Figure 7.2. AFM images of the device before (a) and after (b) sensing trials in serum.

**7.2. Graphene devices for biosensing.** Although CNT-sensors show good sensitivity, their reproducibility suffers from the inhomogeneity of the starting material. Therefore, graphene was considered as an alternative nanomaterial to be used as transducing element of the nanosensors. Several approaches for large-scale fabrication of the graphene or reduced graphene oxide were tested here. The graphene oxide solution was used for the wafer scale fabrication of RGO devices using a novel chemical approach. This fabrication protocol is suitable for building FET-based sensors. With such sensors it is possible to utilize the same detection strategy as it was used here for CNT-FET-sensors (Figure 7.3).

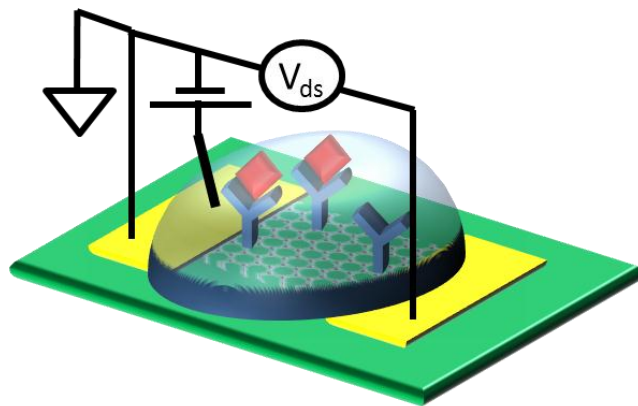
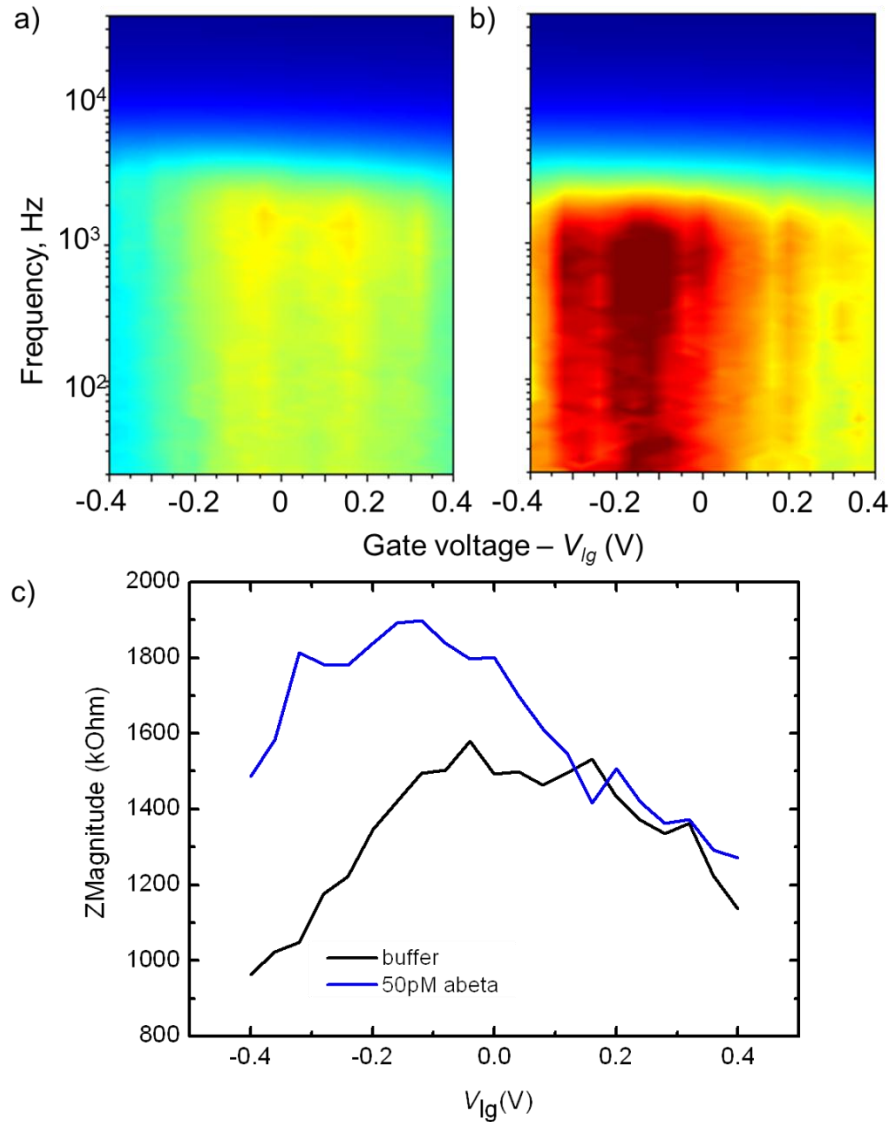


Figure 7.3. Schematic of RGO-FET-immunosensor.

Preliminary results on the detection of amyloid-beta peptide were obtained with the RGO-devices. Amyloid beta peptide is an important marker for the diagnostics of Alzheimer disease [211]. In order to detect amyloid-beta, RGO was modified with a specific monoclonal antibody. Z-Maps were obtained for gate voltages from -0.4 V to 0.4 V on the devices before and after their exposure to the solution of target peptide.



**Figure 7.4. Sensing of beta-amyloid peptide using RGO immunosensor. Z-Maps of the magnitude of impedance before (a) and after (b) exposure to 50pM beta-amyloid solution (scale - 2kOhm to 2.5 MOhm). c) gate characteristics of the device at 100 Hz.**

A clear shift of the gate-dependence curves was observed after the amyloid beta solution was introduced (Figure 7.4). This shift can be explained by the accumulation of amyloid beta on the RGO surface. Amyloid-beta peptide (pI 5.5) carries a negative charge at a working pH of 7.4. This can cause a shift in the gate dependence. The response was not completely reversible. After 90 minutes of wash with high ionic strength buffer, the gate-dependence of the device comes back but not exactly to the same level. This can be explained by a strong antigen-antibody interaction. Future steps in this work will include the optimization of the sensing protocol and the required control experiments.

Demonstrated carbon nanostructure-based sensing platforms show promise for their application due to high sensitivity and low cost. It can be also optimized for performing fundamental single-molecule and kinetic studies, which may benefit from the high sensitivity of the measurements. Both CNT and graphene devices can be used for the detection of low amounts of many other analytes relevant for the medical diagnostics, food and environmental monitoring.



# Bibliography

---

1. Atkinson AJ, Colburn WA, DeGruttola VG, Demets DL, Downing GJ, et al. Biomarkers and surrogate endpoints: preferred definitions and conceptual framework. *Clin Pharmacol Ther.* 2001, 69, 89-95.
2. Elin RJ. Instrumentation in clinical chemistry. *Science.* 1980, 210, 286-289.
3. Durner J. Clinical Chemistry: Challenges for Analytical Chemistry and the Nanosciences from Medicine. *Angew Chem Intern Ed.* 2010, 49, 1026-105.
4. Kiechle FL, Holland CA. Point-of-care testing and molecular diagnostics: miniaturization required. *Clin Lab Med.* 2009, 29, 555-560.
5. Holland CA, Kiechle FL. Point-of-care molecular diagnostic systems--past, present and future. *Curr Opin Microbiol.* 2005, 85, 504-509.
6. Rasooly A. Moving biosensors to point-of-care cancer diagnostics. *Biosensors & Bioelectronics.* 2006, 21, 1847-1850.
7. Lien K-Y, Lee G-B. Miniaturization of molecular biological techniques for gene assays. *Analyst.* 2010, 135, 1499-1518
8. Mascini M, Tombelli S. Biosensors for biomarkers in medical diagnostics. *Biomarkers.* 2008, 13, 637-657
9. Marks RS, Lowe CR, Cullen DC, Weetall HH, Karube Isao. Handbook of Biosensors and Biochips. 2007, Wiley, DOI: 10.1002/9780470061565
10. Clark LC, Lyons C. Electrode systems for continuous monitoring in cardiovascular surgery. In: Ann. N.Y. Acad. Sci. Bd, 1962, 102, 3129-3145
11. Rechnitz GA, Kobosa RK, Riechela SJ, Gebauer CR. A bio-selective membrane electrode prepared with living bacterial cells. *Analytica Chimica Acta.* 1977, 94, 357-365
12. Chambers JP, Arulanandam BP, Matta LL, Weis W, Valdes JJ. Biosensor Recognition Elements. *Curr. Issues Mol. Biol.* 10, 1-12.
13. Collings AF and Caruso F. Biosensors: recent advances. *Rep. Prog. Phys.* 1997, 60, 1397-1445.
14. Joseph D. Bronzino. Medical devices and systems. CRC Press, 2006
15. North SH, Lock EH, Taitt CR, Walton SG. Critical aspects of biointerface design and their impact on biosensor development. *Anal Bioanal Chem.* 2010, 397, 925-933.
16. Azzazy HME, Mansour MMH, Kazmierczak SC. Nanodiagnostics: A New Frontier for Clinical Laboratory Medicine. *Clinical Chemistry.* 2006, 52, 1238-1246.

17. Roy S and Gao Z. Nanostructure-based electrical biosensors. *Nano Today*. 2009, 4, 318-324.
18. Balasubramanian K. Challenges in the use of 1D nanostructures for on-chip biosensing and diagnostics. *Biosens. Bioelectron.* 2010, 26, 1195-1204.
19. Rosi NL and Mirkin CA. Nanostructures in Biodiagnostics. *Chem. Rev.* 2005, 105, 1547-1562.
20. Erickson D, Mandal S, Yang AHJ and Cordovez B. Nanobiosensors: optofluidic, electrical and mechanical approaches to biomolecular detection at the nanoscale. *Microfluid. Nanofluid.* 2008, 4, 33-52.
21. Maroto A, Balasubramanian K, Burghard M and Kern K. Functionalized metallic carbon nanotube devices for pH sensing. *ChemPhysChem*, 2007, 8, 220-223.
22. Homola J. Surface Plasmon Resonance Based Sensors. 2006, Springer.
23. Ozkan M. Quantum dots and other nanoparticles: What they can offer to drug discovery? *Drug Discov. Today*. 2004, 9, 1065-1071
24. Costa-Fernandez JM, Pereiro R and Medel SA. The use of luminescent quantum dots for optical sensing. *Tr. Anal. Chem.* 2006, 25, 207-218.
25. Xu K, Huang JR, Ye ZZ, Ying YB and Li YB. Recent Development of Nano-Materials Used in DNA Biosensors. *Sensors*. 2009, 9, 5534-5557.
26. Hu C and Hu S. Carbon Nanotube-Based Electrochemical Sensors: Principles and Applications in Biomedical Systems. *J. Sens.* 2009, 187615-187645.
27. Vaddiraju S, Tomazos I, Burgess DJ, Jain FC and Papadimitrakopoulos F. Emerging synergy between nanotechnology and implantable biosensors. *Biosens. Bioelectron.* 2010, 25, 1553-1565.
28. Yogeswaran U and Chen SM. A Review on the Electrochemical Sensors and Biosensors Composed of Nanowires as Sensing Material. *Sensors*, 2008, 8, 290-313.
29. Siqueira JR, Caseli L, Cresphilo FN, Zucolotto V and Oliviera ON. Immobilization of biomolecules on nanostructured films for biosensing. *Biosens. Bioelectron.* 2010, 25, 1254-1263.
30. Kurkina T, Balasubramanian K. Towards in vitro molecular diagnostics using nanostructures. *Cell Mol Life Sci.* 2012, 69, 373-88
31. Fiorini GS, Chiu DT. Disposable microfluidic devices: fabrication, function, and application. *Biotechniques*. 2005, 38, 429-446.
32. Haeberle S, Zengerle R. Microfluidic platforms for lab-on-a-chip applications. *Lab Chip*. 2007, 7, 1094-110.
33. Rivet C, Lee H, Hirsch A, Hamilton S, Lu H. Microfluidics for medical diagnostics and biosensors. *Chemical Engineering Science*. 2011, 66, 1490-1507.
34. Wang J. Electrochemical biosensors: Towards point-of-care next term cancer diagnostics. *Biosens Bioelectron.* 2006, 21, 1887-92.

35. Wang Y, Xu H, Zhang J and Li G. Electrochemical Sensors for Clinic Analysis. *Sensors*, 2008, 8, 2043-2081.
36. Grieshaber D, MacKenzie R, Vörös J and Reimhult E. Electrochemical Biosensors - Sensor Principles and Architectures. *Sensors*, 2008, 8, 1400-1458.
37. Pohanka M, Skládal P. Electrochemical biosensors - principles and applications. *J. Appl. Biomed.* 2008, 6, 57-64.
38. Kuriyama T, Kimura J. FET-based biosensors. *Bioprocess Technol.* 1991, 15, 139-162.
39. Yuqing M, Jianguo G, Jianrong C. Ion sensitive field effect transducer-based biosensors. *Biotechnol Adv.* 2003, 21, 527-34.
40. Besteman K, Lee J-O, Wiertz FGM, Heering HA, Dekker C. Enzyme-Coated Carbon Nanotubes as Single-Molecule Biosensors. *Nano Lett.* 2003, 3, 727-730
41. Heller I, Janssens AM, Männik J, Minot ED, Lemay SG, Dekker C. Identifying the mechanism of biosensing with carbon nanotube transistors. *Nano Lett.* 2008, 8, 591-595.
42. Lindholm-Sethson B, Nyström J, Malmsten M, Ringstad L, Nelson A, Geladi P. Electrochemical impedance spectroscopy in label-free biosensor applications: multivariate data analysis for an objective interpretation. *Anal Bioanal Chem.* 2010, 398, 2341-2349.
43. Bonanni A, del Valle M. Use of nanomaterials for impedimetric DNA sensors. *Anal Chim Acta.* 2010, 678, 7-17.
44. Lisdat F, Schafer D. The use of electrochemical impedance spectroscopy for biosensing. *Analytical and Bioanalytical Chemistry.* 2008, 391, 1555-1567
45. Heimann RB, Esvukov SE and Koga Y. Carbon allotropes: a suggested classification scheme based on valence orbital hybridization. *Carbon.* 1997, 35, 1654-1658
46. Qureshi A, Kang WP, Davidson JL, Gurbuz Y. Review on carbon-derived, solid-state, micro and nano sensors for electrochemical sensing applications. *Diamond & Related Materials.* 2009, 18, 1401-1420
47. Iijima S. Helical microtubules of graphitic carbon. *Nature.* 1991, 354, 56 - 58.
48. Dresselhaus MS, Dresselhaus G, Saito R, Physics of carbon nanotubes. *Carbon.* 1995, 33, 883-891
49. Schnorr JM, Swager TM. Emerging Applications of Carbon Nanotubes. *Chemistry of Materials.* 2011, 23, 646-657
50. Endo M, Hayashi T, Kim Y-A. Large-scale production of carbon nanotubes and their applications. *Pure Appl. Chem.* 2006, 78, 1703-1713.
51. Balasubramanian K, Burghard M, Kern K. Effect of the electronic structure of carbon nanotubes on the selectivity of electrochemical functionalization. *Phys. Chem. Chem. Phys.* 2008, 10, 2256-2262.

52. Hiura H, Ebbesen T W, Tanigaki K. Opening and purification of carbon nanotubes in high yields. *Advanced Materials*. 1995, 7, 275-276.
53. Treacy MMJ, Ebbesen TW and Gibson JM. Exceptionally high Young's modulus observed for individual carbon nanotubes. *Nature*, 1996, 381, 678-680.
54. Desprès JF, Daguerre E, Lafdi K. Flexibility of graphene layers in carbon nanotubes. *Carbon*. 1995, 33, 87-92.
55. Ajayan PM, Stephan O, Colliex C, Trauth D. Aligned Carbon Nanotube Arrays Formed by Cutting a Polymer Resin-Nanotube Composite. *Science*, 1994, 265, 1212-1214.
56. Sinha N, Ma J, Yeow JTW. Carbon Nanotube-Based Sensors. *Journal of Nanoscience and Nanotechnology*. 2006, 6, 573-590
57. Chen Z, Tabakman SM, Goodwin AP, Kattah MG, Daranciang D, Wang X, Zhang G, Li X, Liu Z, Utz PJ, Jiang K, Fan S and Dai H Protein microarrays with carbon nanotubes as multicolor Raman labels. *Nat. Biotechnol.* 2008, 26, 1285-1292.
58. Yu X, Munge B, Patel V, Jensen G, Bhirde A, Gong JD, Kim SN, Gillespie J, Gutkind JS, Papadimitrakopoulos F and Rusling JF. Carbon nanotube amplification strategies for highly sensitive immunodetection of cancer biomarkers. *J. Am. Chem. Soc.* 2006, 128, 11199-11205.
59. Balasubramanian K and Burghard M. Biosensors based on carbon nanotubes. *Anal. Bioanal. Chem.* 2006, 385, 452-468.
60. Gruner G. Carbon nanotube transistors for biosensing applications. *Anal. Bioanal. Chem.* 2006, 384, 322-335.
61. Sotiropoulou S, Chaniotakis NA, Carbon nanotube array-based biosensor. *Anal Bioanal. Chem.* 2003, 375, 103-105.
62. Collins PG, Bradley K, Ishigami M, Zettl A. Extreme oxygen sensitivity of electronic properties of carbon nanotubes. *Science*. 2000, 287, 1801-1804
63. Kong J, Franklin NR, Zhou C, Chapline MG, Peng S, Cho K, Dai H. Nanotube molecular wires as chemical sensors. *Science*. 2000, 287, 622-625
64. Star A, Han T-R, Gabriel J-CP, Bradley K, Grüner G. Interaction of aromatic compounds with carbon nanotubes. *Nano Lett.* 2003, 3, 1421-1423
65. Bradley K, Gabriel J-CP, Briman M, Star A, Grüner G. Charge transfer from aqueous ammonia absorbed on nanotube transistors. *Phys. Rev. Lett.* 2003, 91, 2183011-2183014.
66. Star A, Tu E, Niemann J, Gabriel J-CP, Joiner C.S, Valcke C. Label-free detection of DNA hybridization using carbon nanotube network field-effect transistors. *Proc. Natl. Acad. Sci. USA.* 2006, 103, 921-926.
67. Maehashi K, Matsumoto K, Takamura Y, Tamiya E. Aptamer-based label-free immunosensors using carbon nanotube field-effect transistors. *Electroanalysis*. 2009, 21, 1285-1290.

68. Novoselov KS, Geim AK, Morozov SV, Jiang D, Zhang Y, Dubonos SV, Grigorieva IV and Firsov AA. Electric Field Effect in Atomically Thin Carbon Films. *Science*. 2004, 306, 666-669
69. Sakhaee-Pour A, Ahmadian MT, Vafai A, Applications of single-layered graphene sheets as mass sensors and atomistic dust detectors. *Solid State Communications*. 2008, 145, 168-172.
70. Stoller MD, Park S, Zhu Y, An J and Ruoff RS. Graphene-Based Ultracapacitors. *Nano Lett*, 2008, 8, 3498–3502
71. Sundaram RS, Gomez-Navarro C, Balasubramanian K, Burghard M, Kern K. Electrochemical Modification of Graphene. *Advanced Materials*. 2008, 20, 3050–3053.
72. Choi SH, Kim YL, Byun KM. Graphene-on-silver substrates for sensitive surface plasmon resonance imaging biosensors. *Opt Express*. 2011, 19, 458-466
73. Peigney A, Laurent Ch, Flahaut E, Bacsá RR, Rousset A. Specific surface area of carbon nanotubes and bundles of carbon nanotubes. *Carbon*. 2001, 39, 507-514
74. Dong H, Gao W, Yan F, Ji H, Ju H. Fluorescence resonance energy transfer between quantum dots and graphene oxide for sensing biomolecules. *Anal. Chem*. 2010, 82, 5511–5517.
75. Lu C-H, Yang H-H, Zhu C-L, Chen X, Chen GN. A graphene platform for sensing biomolecules. *Angew. Chem. Int. Ed*. 2009, 48, 4785–4787.
76. He S. et al. A graphene nanoprobe for rapid, sensitive, and multicolor fluorescent DNA analysis. *Adv. Funct. Mater*. 2010, 20, 453–459.
77. Kang X, Wang J, Wu H, Aksay IA, Liu J and Lin Y. Glucose Oxidase-graphene-chitosan modified electrode for direct electrochemistry and glucose sensing *Biosens. Bioelectron*. 2009, 25, 901-905.
78. Liu Y, Yu D, Zeng C, Miao Z and Dai L. Biocompatible Graphene Oxide-Based Glucose Biosensors. *Langmuir*, 2010, 26, 6158-6160.
79. Zhuang Z, Li J, Xu R, Dan X. Electrochemical Detection of Dopamine in the Presence of Ascorbic Acid Using Overoxidized Polypyrrole/Graphene Modified Electrodes. *J. Electrochem. Sci*. 2011, 6, 2149 - 2161.
80. Du D, Zou Z, Shin Y, Wang J, Wu H, Engelhard MH, Liu J, Aksay IA, Lin Y. Sensitive Immunosensor for Cancer Biomarker Based on Dual Signal Amplification Strategy of Graphene Sheets and Multienzyme Functionalized Carbon Nanospheres. *Anal. Chem*. 2010, 82, 2989-2995.
81. Shao Y, Wang J, Wu H, Liu J, Aksay IA, Lina Y. Graphene Based Electrochemical Sensors and Biosensors. *Electroanalysis* 2010, 22, 1027 - 1036.
82. Dan P, Lu Y, Kybert NJ, Luo ZT, Johnson ATC. Intrinsic Response of Graphene Vapor Sensors. *Nano Letters*. 2009, 9, 1472-1475.

83. Schedin F, Geim AK, Morozov SV, Hill EW, Blake P, Katsnelson MI, Novoselov KS. Detection of individual gas molecules adsorbed on graphene. *Nature Materials*. 2007, 6, 652 – 655.
84. Ang PK, Chen W, Wee ATS, Loh KP. Solution-gated epitaxial graphene as pH sensor. *J. Am. Chem. Soc.* 2008, 130, 14392–14393.
85. Dong X, Shi Y, Huang W, Chen P, Li LP. Electrical Detection of DNA Hybridization with Single-Base Specificity Using Transistors Based on CVD-Grown Graphene Sheets. *Adv. Mater.* 2010, 22, 1–5.
86. Mohanty N, Berry V. Graphene-based single-bacterium resolution biodevice and DNA transistor: interfacing graphene derivatives with nanoscale and microscale biocomponents. *Nano Letters*. 2008, 8, 4469–4476.
87. Ohno Y, Maehashi K, Yamashiro Y, Matsumoto K. Electrolyte-gated graphene field-effect transistors for detecting pH and protein adsorption. *Nano Letters*. 2009, 9, 3318–3322.
88. Yang W, Ratinac KR, Ringer SP, Thordarson P, Gooding JJ, Braet F. Carbon Nanomaterials in Biosensors: Should You Use Nanotubes or Graphene? *Angew. Chem. Int. Ed.* 2010, 49, 2114 – 2138
89. Wang Y, Li Z, Wang J, Li J, Lin Y. Graphene and graphene oxide: biofunctionalization and applications in biotechnology. *Trends in Biotechnology*. 2011, 29, 205-212
90. Hirsch A, Vostrowsky O. Functionalization of Carbon Nanotubes. *Topics in Current Chemistry*. 2005, 45, 193-237
91. Martinez MT, Tseng YC, Ormategui N, Loinaz I, Eritja R, Bokor J. Label-Free DNA Biosensors Based on Functionalized Carbon Nanotube Field Effect Transistors. *Nano Letters*. 2009, 9, 530–536
92. Geim AK. Graphene: Status and Prospects. *Science*. 2009, 324, 1530-1534
93. Loh KP, Bao Q, Ang PK, Yang . The chemistry of graphene. *J. Mater. Chem.* 2010, 20, 2277–2289
94. Bahr JL, Tour JM. Covalent chemistry of single-wall carbon nanotubes. *J. Mater. Chem.* 2002, 12, 1952-1958.
95. Malig J, Englert JM, Hirsch A, Guldi DM. Wet Chemistry of Graphene. *Interface*. 2011, 20, 53-56.
96. Jeng ES, Moll AE, Roy AC, Gastala JB, Strano MS. Detection of DNA Hybridization Using the Near-Infrared Band-Gap Fluorescence of Single-Walled Carbon Nanotubes. *Nano Letters*, 2006, 6, 371–375.
97. Lu C-H, Yang H-H, Zhu CL, Chen X, Guo-Nan Chen Prof. A Graphene Platform for Sensing Biomolecules. *Angew. Chem. Int. Ed*, 2009, 48, 4785–4787.
98. Balavoine F, Schultz P, Richard C, Mallouh V, Ebbesen TW, Mioskowski C. Helical crystallization of proteins on carbon nanotubes: a first step towards the development of new biosensors. *Angew. Chem. Int. Ed.* 1999, 38, 1912–1915.

99. Leyden MR, Schuman C, Sharf T, Kevek J, Remcho VT, Minot ED. Fabrication and Characterization of Carbon Nanotube Field-Effect Transistor Biosensors. *Proc. SPIE*, 2010, doi:10.1117/12.861329.
100. Lotya M, King PJ, Khan U, De S, Coleman JN. High-Concentration, Surfactant-Stabilized Graphene Dispersions. *ACS Nano*. 2010, 4, 3155–3162.
101. Lotya M, Hernandez Y, King PJ, Smith RJ, Nicolosi V, Karlsson LS, Blighe FM, De S, Wang Z, McGovern, Duesberg GS, Coleman JN. Liquid Phase Production of Graphene by Exfoliation of Graphite in Surfactant/Water Solutions. *J. Am. Chem. Soc.*, 2009, 131, 3611–3620.
102. Balasubramanian K, Burghard M. Chemically functionalized carbon nanotubes. *Small*, 2005, 1, 180-192.
103. Dreyer DR, Park S, Bielawski CW and Ruoff RS. The chemistry of graphene oxide. *Chem. Soc. Rev.* 2010, 39, 228-240.
104. Cui J, Burghard M, Kern K. Reversible Sidewall Osmylation of Individual Carbon Nanotubes. *Nano Letters*. 2003, 3, 613-615.
105. Liu H, Ryu S, Chen Z, Steigerwald ML, Nuckolls C, Brus LE. Photochemical Reactivity of Graphene. *J. Am. Chem. Soc.* 2009, 131, 17099–17101.
106. Balasubramanian K, Burghard M. Electrochemically functionalized carbon nanotubes for device applications. *J. Mater. Chem.* 2008, 18, 3071-3083.
107. Maroto A, Balasubramanian K, Burghard M, Kern K. Functionalized metallic carbon nanotube devices for pH sensing. *ChemPhysChem*. 2007, 8, 220-223.
108. Balasubramanian K, Friedrich M, Jiang C, Fan Y, Mews A, Burghard M, Kern K. Electrical transport and confocal Raman studies of electrochemically modified individual carbon nanotubes. *Adv. Mater.* 2003, 15, 1515-1518.
109. Allen BL, Kichambare PD, Star A. Carbon Nanotube Field-Effect-Transistor-Based Biosensors. *Adv. Mater.* 2007, 19, 1439–1451.
110. Kong J, LeRoy BJ, Lemay SG, Dekker C. Integration of a gate electrode into carbon nanotube devices for scanning tunneling microscopy. *Appl. Phys. Lett.* 2005, 86, 112106–112108.
111. Chen XQ, Saito T, Yamada H, Matsushige K. Aligning single-wall carbon nanotubes with an alternating-current electric field. *Appl. Phys. Lett.* 2001, 78, 3714- 3716.
112. Krupke R, Hennrich F, Weber HB, Kappes MM and Lohneysen HV, Simultaneous Deposition of Metallic Bundles of Single-walled Carbon Nanotubes Using AC-dielectrophoresis. *Nano Lett*, 2003, 3, 1019-1023.
113. Leonhardt AS, Hampel, et al. Synthesis, Properties, and Applications of Ferromagnetic-Filled Carbon Nanotubes. *Chemical Vapor Deposition* 2006, 12, 380-387.

114. Nakashima N. Solubilization of single-walled carbon nanotubes with condensed aromatic compounds. *Sci. Technol. Adv. Mater.* 2006, 7, 609.
115. Wang J, Musameh M, Lin Y. Solubilization of Carbon Nanotubes by Nafion toward the Preparation of Amperometric Biosensors. *Journ. Am. Chem. Soc.* 2003, 125, 2408-2409.
116. Hou P, Liu C, Cheng H. Purification of carbon nanotubes. *Carbon*, 2008, 46, 2003-2025.
117. Chiang IW, Brinson BE, Huang AY, Willis PA, Bronikowski MJ, Margrave JL, Smalley RE, and Hauge RH. Purification and Characterization of Single-Wall Carbon Nanotubes (SWNTs) Obtained from the Gas-Phase Decomposition of CO (HiPco Process). *J. Phys. Chem. B*, 2001, 105, 8297-8301
118. Yu AP, Bekyarova E, Itkis ME, Fakhрутdinov D, Webster R and Haddon RC. Application of centrifugation to the large-scale purification of electric arc-produced single-walled carbon nanotubes. *J Am Chem Soc*, 2006, 128, 9902-9908.
119. Liu H, Nishide D, Tanaka T, Kataura H. Large-scale single-chirality separation of single-wall carbon nanotubes by simple gel chromatography. *Nat. Commun.* 2011, 2, doi:10.1038/ncomms1313.
120. Arnold MS, Green AA, Hulvat JF, Stupp SI, Hersam MC. Sorting carbon nanotubes by electronic structure using density differentiation. *Nature Nanotechnology*, 2006, 1, 60 - 65.
121. Ghosh S, Bachilo SM, Weisman RB. Advanced Sorting of Single-Walled Carbon Nanotubes by Nonlinear Density Gradient Ultracentrifugation. *Nature Nanotechnology*, 2010, 5, 443-450.
122. Duesberga GS, Blaua W, Byrneb HJ, Muste J, Burghard M, Roth C. Chromatography of Carbon Nanotubes. *Synthetic Metals*, 1999, 103, 2484-2485.
123. Ahmad A, Kurkina T, Kern K, Balasubramanian K. Applications of the static quenching of rhodamine B by carbon nanotubes. *ChemPhysChem*, 2009, 10, 2251-2255.
124. Lu Q, Freedman KO, Rao R, Huang G, Lee J, Larcom LL, Rao AM, and Ke PC. Diffusion of carbon nanotubes with single-molecule fluorescence microscopy *J. Appl. Phys.* 2004, 96, 6772-6775.
125. Satishkumar BC, Brown LO, Gao Y, Wang C-C, Wang H-L, Doorn SK. Reversible fluorescence quenching in carbon nanotubes for biomolecular sensing. *Nature Nanotechnology*, 2007, 2, 560 - 564.
126. Pohl HA, Dielectrophoresis the behavior of neutral matter in nonuniform electric fields. 1978, Cambridge University Press. Cambridge.
127. Freer EM, Grachev O, Duan X, Martin S, Stumbo DP. High-yield self-limiting single-nanowire assembly with dielectrophoresis. *Nature Nanotechnology*, 2010, 5, 525-530.



128. Dimaki M, Bøggild P. Dielectrophoresis of carbon nanotubes using microelectrodes: a numerical study. *Nanotechnology*, 2004, 15, 1095-1102.
129. Sagar AS, Kern K, Balasubramanian K. Marker-free on-the-fly fabrication of graphene devices based on fluorescence quenching. *Nanotechnology*, 2010, 21, 015303.
130. Vlandas A, Kurkina T, Ahmad A, Kern K, Balasubramanian K. Enzyme-free sugar sensing in microfluidic channels with an affinity-based single-wall carbon nanotube sensor. *Anal Chem.* 2010, 15, 6090-6097.
131. Balasubramanian K, Lee EJH, Weitz RT, Burghard M, Kern K. Carbon nanotube transistors - chemical functionalization and device characterization. *Phys. Stat.Sol. A*, 2008, 205, 633-646.
132. Kernohan AFB, Perry CG, Small M. Clinical impact of the new criteria for the diagnosis of diabetes mellitus. *Clin. Chem. Lab. Med.* 2003, 41, 1239-45.
133. Wang J. Electrochemical glucose biosensors. *J. Chem. Rev.* 2008, 108, 814-825.
134. Liu G, Lin Y. Amperometric glucose biosensor based on self-assembling glucose oxidase on carbon nanotubes. *Electrochem.Commun.* 2006, 8, 251-256.
135. Wilson GS, Gifford R. Biosensors for real-time in vivo measurements. *Biosens. Bioelectron.* 2005, 20, 2388-2403
136. Park S, Boo H, Chung TD. Electrochemical non-enzymatic glucose sensors. *Anal. Chim. Acta*, 2006, 556, 46-57.
137. Yuan J, Wang K, Xia X. Highly Ordered Platinum-Nanotubule Arrays for Amperometric Glucose Sensing. *Adv. Funct. Mater.* 2005, 15, 803-809.
138. Cherevko S, Chung C-H. Gold nanowire array electrode for non-enzymatic voltammetric and amperometric glucose detection. *Sens. Actuat. B* 2009, 142, 216-223.
139. Wang G, Wei Y, Zhang W, Zhang X, Fang B, Wang L. Enzyme-free amperometric sensing of glucose using Cu-CuO nanowire composites. *Microchim. Acta*, 2010, 168, 87-92.
140. Ozcan L, Sahin Y, Tuerk H. Non-enzymatic glucose biosensor based on overoxidized polypyrrole nanofiber electrode modified with cobalt(II) phthalocyanine tetrasulfonate. *Biosens. Bioelectron.* 2008, 24, 512-517.
141. Satheesh Babu TG, Ramachandran T. Development of highly sensitive non-enzymatic sensor for the selective determination of glucose and fabrication of a working model. *Electrochim. Acta*, 2010, 55, 1612-1618.
142. Vassiliyev YB, Khazova OA, Nikolaeva NN. Kinetics and mechanism of glucose electrooxidation on different electrode-catalysts: Part I. Adsorption and oxidation on platinum. *J. Electroanal. Chem.* 1985, 196, 105-125

143. James TD, Phillips MD, Shinkai S. Boronic Acids in Saccharide Recognition. 2006, RSC Publishing.
144. Lorand JP, Edwards JO. Polyol complexes and structure of the benzenboronate ion. *J. Org. Chem.* 1959, 24, 769-774.
145. Fang H, Kaur G, Wang B. Progress in Boronic Acid-Based Fluorescent Glucose Sensors. *J. Fluoresc.* 2004, 14, 481-489.
146. Pickup JC, Hussain F, Evans ND, Rolinski OJ, Birch DJS. In vivo glucose monitoring: the clinical reality and the promise. *Biosens. Bioelectron.* 2005, 20, 2555-2565.
147. Arimori S, Ushiroda S, Peter LM, Jenkins ATA, James TD. A modular electrochemical sensor for saccharides. *Chem Commun.* 2002, 2368-2369.
148. Shoji E, Freund MS. Potentiometric Saccharide Detection Based on the pKa Changes of Poly(aniline boronic acid). *J. Am. Chem. Soc.* 2001, 123, 3383-3384.
149. Matsumoto A, Sato N, Sakata T, Kataoka K, Miyahara YJ. Glucose-sensitive field effect transistor using totally synthetic compounds. *Solid State Electrochem.* 2009, 13, 165-170.
150. Song SY, Yoon HC. Boronic acid-modified thin film interface for specific binding of glycated hemoglobin (HbA1c) and electrochemical biosensing. *Sens. Actuators B*, 2009, 140, 233-239.
151. Aime S, Botta M, Dastru W, Fasano M, Panero M, Arnelli A. Synthesis and characterization of a novel DTPA-like gadolinium(III) complex: a potential reagent for the determination of glycated proteins by water proton NMR relaxation measurements. *Inorg Chem*, 1993, 32, 2068-2071.
152. Djanashvili K, Frullano L, Peters JA. Molecular Recognition of Sialic Acid End Groups by Phenylboronates. *Chem. Eur. J.* 2005, 11, 4010-4018.
153. Yoon J, Czarnik AW. Fluorescent chemosensors of carbohydrates. A means of chemically communicating the binding of polyols in water based on chelation-enhanced quenching. *J. Am. Chem. Soc.* 1992, 114, 5874-5875.
154. Springsteen G, Wang B. A Detailed Examination of Boronic Acid-Diol Complexation. *Tetrahedron*, 2002, 58, 5291-5300.
155. Back JH, Shim M. pH-dependent electron-transport properties of carbon nanotubes, *J. Phys. Chem. B*, 2006, 110, 23736-23741.
156. Bergveld P. ISFET, theory and practice. *Sens. Actuators B*. 2003, 88, 1-20.
157. Sklar J. DNA hybridization in diagnostic pathology. *Hum Pathol.* 1985, 16, 654-658.
158. Viscidi RP, Yolken RG. Molecular diagnosis of infectious diseases by nucleic acid hybridization. *Mol Cell Probes*. 1987, 1, 3-14
159. Paul JH, John HP. Gene expression by mRNA analysis. in *Methods in Microbiology*, 30, 395 (Academic Press, 2001).

160. VanGuilder HD, Vrana KE, Freeman WM. Twenty-five years of quantitative PCR for gene expression analysis. *BioTechniques*, 2008, 44, 619-626.
161. Bubendorf L, High-throughput microarray technologies: from genomics to clinics. *Eur Urol*. 2001, 40, 231-238.
162. Campas M, Katakis I. DNA biochip arraying, detection and amplification strategies. *Trends in Analytical Chemistry*, 2004, 23, 49-62.
163. Moeller R, Fritzsche W. Chip-based electrical detection of DNA. *IEE Proc Nanobiotechnol*. 2005, 152, 47-51.
164. Mao X. et al. The Lab-on-a-Chip Approach for Molecular Diagnostics. in *Molecular Diagnostics 21* (Academic Press, San Diego, 2010).
165. Kaltenboeck B, Wang C, Gregory SM. Advances in Real Time PCR: Application to Clinical Laboratory Diagnostics. in *Advances in Clinical Chemistry*, 40, 219 (Elsevier, 2005).
166. Yee AJ et al. DNA Microarrays in Biological Discovery and Patient Care. in *Genomic and Personalized Medicine*, 157 (Academic Press, New York, 2009).
167. Kurkina T, Vlandas A, Ahmad A, Kern K, Balasubramanian K. Label-Free Detection of Few Copies of DNA with Carbon Nanotube Impedance Biosensors. *Angew Chem Int Ed*. 2011, 50, 3710-3714
168. Yan F, Tang H. Applications of thin film transistors in label-free DNA biosensors. *Exp. Rev. Molecul. Diagn*. 2010, 10, 547-549.
169. Drummond TG, Hill MG, Barton JK. Electrochemical DNA sensors. *Nature Biotechnol*. 2003, 21, 1192-1199.
170. Ganguly A, Chen C-P, Lai Y-T, Kuo C-C, Hsu C-W, Chen K-H, Chen L-C. Functionalized GaN nanowire-based electrode for direct label-free voltammetric detection of DNA hybridization. *Journ. Mat. Chem*. 2009, 19, 928-933.
171. Fang Z, Kelley SO. Direct electrocatalytic mRNA detection using PNA-nanowire sensors, *Anal. Chem*, 2009, 81, 612-617.
172. Gao Z, Agarwal A, Trigg AD, Singh N, Fang C, Tung C-H, Fan Y, Buddharaju KD, Kong, J. Silicon nanowire arrays for label-free detection of DNA. *Anal. Chem*. 2007, 79, 3291-3297.
173. Hahm J, Lieber CM. Direct ultrasensitive electrical detection of DNA and DNA sequence variations using nanowire nanosensors. *Nano Lett*. 2004, 4, 51-54.
174. Larrimore L, Nad S, Zhou X, Abruna H, McEuen PL. Probing electrostatic potentials in solution with carbon nanotube transistors. *Nano Lett*. 2006, 6, 1329-1333.

175. Minot ED, Janssens AM, Heller I, Heering HA, Dekker C, Lemay SG. Carbon nanotube biosensors: the critical role of the reference electrode. *Appl. Phys. Lett.*, 2007. 91, 093507-9.
176. Hermanson, G. T. (2008) *Bioconjugate Techniques* (2nd Ed.) Academic Press, San Diego.
177. Gui EL, Li LJ, Zhang K, Xu, Y, Dong X, Ho X, Lee PS, Kasim J, Shen ZX, Rogers JA, Mhaisalkar SG. DNA sensing by field-effect transistors based on networks of carbon nanotubes. *J. Am. Chem. Soc.* 2007, 129, 14427-14432.
178. Zheng M, Jagota A, Semke ED, Diner BA, Mclean RS, Lustig SR, Richardson RE, Tassi NG. DNA assisted dispersion and separation of carbon nanotubes. *Nature Mater.* 2003, 2, 338-342.
179. Yang R, Jin J, Chen Y, Shao N, Kang H, Xiao Z, Tang Z, Wu Y, Zhu Z, Tan W. Carbon Nanotube-Quenched Fluorescent Oligonucleotides: Probes that Fluoresce upon Hybridization. *Journ. Am. Chem. Soc.* 2008, 130, 8351-8358.
180. Inagaki M, Kim YA and Endo M. Graphene: preparation and structural perfection. *J. Mater. Chem.*, 2011, 21, 3280-3294.
181. Liu Z, Bol AA, Haensch W. Large-Scale Graphene Transistors with Enhanced Performance and Reliability Based on Interface Engineering by Phenylsilane Self-Assembled Monolayers. *Nano Letters*, 2010, 11, 523-528.
182. Kim KS, Zhao Y, Jang H, Lee SY, Kim JM, Kim KS, Ahn JH, Kim P, Choi J-Y, Hong BH, Large-scale pattern growth of graphene films for stretchable transparent electrodes. *Nature*, 2009, 457, 706-710.
183. Li X, Cai W, An J, Kim S, Nah J, Yang D, Piner R, Velamakanni A, Jung I, Tutuc E, Banerjee SK, Colombo L, Ruoff RS. Large-Area Synthesis of High-Quality and Uniform Graphene Films on Copper Foils. *Science*, 2009, 324, 1312-1314.
184. Hernandez V, Lotya NM, et al. High-yield production of graphene by liquid-phase exfoliation of graphite. *Nature Nano*, 2008, 3, 563-568.
185. Zhu Y, Murali S, Cai W, Li X, Suk JW, Potts JR, Ruoff RS. Graphene and Graphene Oxide: Synthesis, Properties, and Applications. *Adv. Mater.*, 2010, 22, 3906-3924.
186. Burg BR, Lütolf F, Schneider J, Schirmer NC, Schwamb T, Poulikakos D. High-Yield Dielectrophoretic Assembly of Two-Dimensional Graphene Nanostructures. *Appl. Phys. Lett.* 2009, 94, 053110
187. Stankovich S, Dikin DA, Piner RD, Kohlhaas KA, Kleinhammes A, Jia Y, Wu Y, Nguyen ST and Ruoff RS. Synthesis of graphene-based nanosheets via chemical reduction of exfoliated graphite oxide. *Carbon* 2007, 45, 1558-1565.
188. Vijayaraghavan A, Sciascia C, Dehm S, Lombardo A, Bonetti A, Ferrari AC, Krupke R. Dielectrophoretic Assembly of High-Density Arrays of

- Individual Graphene Devices for Rapid Screening. *ACS Nano*, 2009, 3, 1729-1734.
189. Wie Z, Barlow DE, Sheehan DE. The Assembly of Single-Layer Graphene Oxide and Graphene Using Molecular Templates. *Nano Letters*, 2008, 8, 3141-3145.
190. Joung D, Chunder A, Zhai L and Khondaker S. High yield fabrication of chemically reduced graphene oxide field effect transistors by dielectrophoresis. *Nanotechnology*, 2010, 21, 165202.
191. Hummers WS, Offeman RE. Preparation of Graphitic Oxide. *J. Am. Chem. Soc.*, 1958, 80, 1339-1339.
192. Kurkina T, Sundaram S, Sundaram RS, Kern K, Balasubramanian K. Site-specific chemical anchoring of graphene oxide for device applications. *ACS Nano*, 2012, 6, 5514-5520.
193. Losic D, Cole M, Thissen H, Voelcker NH. Ultrathin polytyramine films by electropolymerisation on highly doped p-type silicon electrodes. *Surface Science*, 2005, 584, 245-257.
194. Tran LD, Piro B, Pham MC, Ledoan T, Angiari C, Dao LH, Teston F. A polytyramine film for covalent immobilization of oligonucleotides and hybridization. *Synthetic Metals*, 2003, 139, 251-262.
195. Liu M-Q, Jiang J-H, Feng Y-L, Shen G-L, Yu R-Q. Glucose Biosensor Based on Immobilization of Glucose Oxidase in Electrochemically Polymerized Polytyramine Film and Overoxidised Polypyrrole Film on Platinized Carbon Paste Electrode. *Chinese Journal of Analytical Chemistry*, 2007, 35, 1435-1438.
196. Park S, Dikin DA, Nguyen ST, Ruoff RS. Graphene Oxide Sheets Chemically Cross-Linked by Polyallylamine. *The Journal of Physical Chemistry C*. 2009, 11, 15801-15804.
197. Kang H, Kulkarni A, Stankovich S, Ruoff RS, Baik S. Restoring electrical conductivity of dielectrophoretically assembled graphite oxide sheets by thermal and chemical reduction techniques. *Carbon*, 47, 2009, 1520-1525.
198. Melitz W, Shen J, Kummel AC, Lee S. Kelvin probe force microscopy and its application. *Surface Science Reports*, 2011, 66,1-27
199. Nonnenmacher M, O'Boyle MP, and Wickramasinghe HK. Kelvin probe force microscopy. *Appl. Phys. Lett.* 1991, 58, 2921-2923
200. Yan L, Punckt C, Aksay IA, Mertin W, Bacher G. Local Voltage Drop in a Single Functionalized Graphene Sheet Characterized by Kelvin Probe Force Microscopy. *Nano Lett.* 2011, DOI: 10.1021/nl201070c
201. Shruti K, Shrey K, Vibha R. Micro RNAs: tiny sequences with enormous potential. *Biochem Biophys Res Commun.* 2011, 407, 445-449

202. Soifer HS, Rossi JJ, Saetrom P. MicroRNAs in Disease and Potential Therapeutic Applications. *Mol Ther.* 2007, **15**, 2070-2079
203. Lawrie CH, Gal S, Dunlop HM, et al. Detection of elevated levels of tumour-associated microRNAs in serum of patients with diffuse large B-cell lymphoma. *Br J Haematol.* 2008, **141**, 672-675.
204. Wang R, Li N, Zhang Y, Ran Y, Pu J. Circulating MicroRNAs are Promising Novel Biomarkers of Acute Myocardial Infarction. *Intern Med.* 2011, **50**, 1789-95.
205. Zheng D, Haddadin S, Wang Y, Gu LQ, Perry MC, Freter CE, Wang MX. Plasma microRNAs as novel biomarkers for early detection of lung cancer. *Int J Clin Exp Pathol.* 2011, **15**, 575-86
206. Yu DC, Li QG, Ding XW, Ding YT. Circulating MicroRNAs: Potential Biomarkers for Cancer. *Int J Mol Sci.* 2011, **12**, 2055-2063
207. Ceribelli A, Nahid MA, Satoh M, Chan EK. MicroRNAs in rheumatoid arthritis. *FEBS Lett.* 2011, doi:10.1016/j.febslet.2011.05.020
208. Gilad S, Meiri E, Yogev Y, Benjamin S, Lebanony D, et al. Serum MicroRNAs Are Promising Novel Biomarkers. *PLoS ONE*, 2008, **3**, e3148
209. Chen X, Ba Y, Ma L, et al. Characterization of microRNAs in serum: a novel class of biomarkers for diagnosis of cancer and other diseases. *Cell Res.* 2008, **18**, 997-1006.
210. Kole PL, Venkatesh G, Kotechac J, Sheshalad R. Recent advances in sample preparation techniques for effective bioanalytical methods. *Biomed. Chromatography*, 2010, **25**, 199-217.
211. Barber RC. Biomarkers for early detection of Alzheimer disease. *J Am Osteopath Assoc.* 2010, **110**, 10-15.

# Acknowledgements

---

It is a pleasure to thank those who made this thesis possible. First of all, I am grateful to Prof. Klaus Kern for giving me the opportunity to work in the Nanoscale Science department of the Max Planck Institute for Solid State Research and to Prof. Bernard Spengler who agreed to supervise my work in Justus Liebig University in Giessen.

Of the many people who have been enormously helpful in the preparation of this thesis, I am especially thankful to my group leader and a good friend Dr. Kannan Balasubramanian for the guidance, encouraging discussions and great ideas. It was also very pleasant to work together with Dr. Alexis Vlandas gave me an introduction to the lab instrumentation and was always helpful in my measurements. The valuable suggestions given by Dr. Ashraf Ahmad have been a remarkable help in performing experiments during my PhD. I am particularly thankful to Dr. Vivek Pachauri with whom I shared the most amount of time and who was my greatest support. And, of course, for the sharing nice time in the lab and performing experiments together I am grateful to other members of Nadia group - Nassim Rafiefard, Divya Balakrishnan, Subramanian Sundaram and Janina Krieg, and to our special member - Ravi Shankar Sundaram.

I am particularly thankful to Armin Schulz for the introduction to Raman spectroscopy and Achim Güth for the assistance in the clean room. I really appreciate the help of Stephan Schmid and Yvonne Link from Technology Group who participated in preparation of the most of my devices. I am also grateful to Dr. K. Heimann (Karlsruhe Institute of Technology) and Prof. C. Richert (University of Stuttgart) for the help with nanoparticles.

I can't not to remember my first scientific supervisor, beloved Varetska Tamara Vladimirovna. I will never forget her wise suggestions. She was a great treasure to me. I am also very grateful to my former supervisor Verevka S.V. who was supporting me during my master thesis and gave me nice advice during my PhD.

I am very happy that during my postgraduate work I met a lot of friends. It was a great fun spending time with them in the institute, during lunches and parties, while going out in Stuttgart and during our trips to beautiful places in Europe. My thanks for all that fun during PhD goes to Ravi Shankar Sundaram,

Eduardo Lee, Liza Herrera Diez, Eberhard Ulrich Stuetzel, Adarsh Sagar, Eva Peters, Hadj Benia, Paola Acosta, Violetta Sessi, Nasiba Abdurakhmanova, Christina Gomez-Navarro, Magali Lingenfelder, Nadja Amsharov and many-many others. I would also like to thank people from other departments - Jaysen Nelayah, Hilton Borbosa de Aguiar, Burac Ozdol, Burcu Ogüt, Ivana Krkljus, Hye Jin Park, Katarina Djuris, Hyeyeon Ryu, Slava Saltykov, ..... - for the great time spent together.

Finally, I owe my deepest gratitude to my family and friends in Ukraine. I can always feel their warm support.



### **PUBLICATIONS:**

1. Ahmad A, Kurkina T, Kern K, Balasubramanian K. Applications of the static quenching of rhodamine B by carbon nanotubes. *ChemPhysChem*, 2009, 10, 2251-2255.
2. Vlandas A, Kurkina T, Ahmad A, Kern K, Balasubramanian K. Enzyme-free sugar sensing in microfluidic channels with an affinity-based single-wall carbon nanotube sensor. *Anal Chem*. 2010, 15, 6090-6097.
3. Kurkina T, Vlandas A, Ahmad A, Kern K, Balasubramanian K. Label-Free Detection of Few Copies of DNA with Carbon Nanotube Impedance Biosensors. *Angew Chem Int Ed*. 2011, 50, 3710-3714
4. Kurkina T, Balasubramanian K. Towards in vitro molecular diagnostics using nanostructures. *Cell Mol Life Sci*. 2011, DOI: 10.1007/s00018-011-0855-7
5. Balasubramanian K, Kurkina T, Ahmad A, Burghard M, Kern K. Tuning the functional interface of carbon nanotubes by electrochemistry: Towards nanoscale chemical and biosensors. (accepted to *J. Mater. Res.*)
6. Kurkina T, Sundaram S, Sundaram RS, Kern K, Balasubramanian K. Site-specific chemical anchoring of graphene oxide for device applications. *ACS Nano*, 2012, 6, 5514-5520.

### **CONFERENCE PRESENTATIONS:**

1. International Conference on Nanosciences & Nanotechnologies, 10-17 July 2010, Ouranoupolis, Greece. Poster presentation: T. Kurkina, A. Vlandas, A. Ahmad, K. Kern, K. Balasubramanian. Electrical sensing of sugars based on electrochemically functionalized single wall carbon nanotubes.
2. NT11 International Conference on the Science and Application of Nanotubes, 10 - 16 July, 2011, Cambridge, UK. Oral presentation: T. Kurkina, A. Vlandas, A. Ahmad, K. Kern, K. Balasubramanian. Label-free electrical detection of DNA based on electrochemically functionalized SWCNTs.

## DECLARATION

I declare that I have completed this dissertation without the unauthorized help of a second party and only with the assistance acknowledged therein. I have appropriately acknowledged and referenced all text passages that are derived literally from or are based on the content of published or unpublished work of others, and all information that relates to verbal communications. I have abided by the principles of good scientific conduct laid down in the charter of the Justus Liebig University of Giessen in carrying out the investigations described in the dissertation.

---

Place, Date

Signature

## VERSICHERUNG

Ich erkläre: Ich habe die vorgelegte Dissertation selbstständig und ohne unerlaubte fremde Hilfe und nur mit den Hilfen angefertigt, die ich in der Dissertation angegeben habe. Alle Textstellen, die wörtlich oder sinngemäß aus veröffentlichten Schriften entnommen sind, und alle Angaben, die auf mündlichen Auskünften beruhen, sind als solche kenntlich gemacht. Bei den von mir durchgeführten und in der Dissertation erwähnten Untersuchungen habe ich die Grundsätze guter wissenschaftlicher Praxis, wie sie in der "Satzung der Justus-Liebig-Universität Gießen zur Sicherung guter wissenschaftlicher Praxis" niedergelegt sind, eingehalten.

---

Ort, Datum