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## Chapter 1

# Fabrication of Nanodiamonds

### Remark:

die section mit den silicon implanted nanodiamonds muss von dir ueberarbeitet werden. Die hab ich deshalb vorerst ausgelassen. Es gibt 3 teile (einfuehrung, preliminary tests und final procedure). Ich hab den eindruck, dass sich da konzeptionell vieles wiederholt und das man das ganze vielleicht in einer section unterbringen kann und die feineren unterscheidungen einfach beilaeufig erwaehnt.

“Diamond forms under high temperature and pressure conditions that exist only about 100 miles beneath the earth’s surface.”

Gemological Institute of America Inc.

Due to its extraordinary properties, diamond has transcended its sole purpose as a rare gem and developed into an important tool enabling various applications in industry and research. This change has been driven largely by the development of methods allowing to cheaply produce synthetic diamond. While synthetic diamonds are typically small and without the splendeur generally associated with diamond, they can be produced on-demand and are thus arguably more useful than rare gemstones. In this chapter we introduce the two most common methods for the fabrication of diamonds in a laboratory setting: The high-pressure, high-temperature and the chemical vapor deposition method. The aptly named high-pressure, high-temperature (HPHT) process mimics the conditions under which diamond is formed in nature and is widely used to synthetically produce diamonds for industrial applications such as utilization of diamond as a abrasive. While HPHT diamonds are utilized in this work, most reported measurements are based on diamond produced with the CVD method in which diamonds are grown using a hydrocarbon gas mixture. Both processes have incommon that defects and impurities are a naturally occurring. For a more extensive list of diamond production processes refer to [1]. In the context of this thesis nano-sized dimonds are required. They can be obtained by milling larger sized HPHT or CVD diamonds down to the desired grain size in a vibrational mill. The obtained nanodiamonds are small enough that individual specimen have a chance of hosting a singleton SiV center. These can be identified and used for further exploration.

## 1.1 High-Pressure High-Temperature Diamond

The HPHT method was the first process to successfully synthesize diamond in 1879. Today, it is still widely used due to its relatively cheap production costs for small diamonds [2]. In the HPHT process, diamond is synthesized from graphite under temperatures of up to  $1.5 \times 10^3$  °C pressures between  $5 \times 10^4$  bar and  $10^6$  bar [1]. Under these extreme conditions, carbon transitions from its graphite to its diamond phase because the latter becomes energetically favorable [?, ?, ?, ?]. The machine used for this kind of synthesis is a press. For some forms of this method, a metallic catalyst solvent is added lowering the required pressures and temperatures by causing graphite to dissolve earlier. At the same time the catalyst promotes the crystallisation process. Several press designs exist, all of which relying on creating and maintaining high pressures and a high temperatures. While it is possible to grow big ( $> 10$  carat) high-quality diamonds with the HPHT process, its cost quickly increases and thus becomes unfavorable.

In this thesis, HPHT nanodiamonds produced by Davydov et al. [3] are spectroscopically investigated.

## 1.2 Chemical Vapor Deposition Diamond

In contrast to the HPHT method, diamond is crystallized from carbon available in the gas phase in the CVD method. The process still requires respectable temperatures in the range of 700 °C to 1300 °C but makes due with the low pressures of less than 1 bar available in a vacuum growth chamber [?].

The chamber contains a vapour consisting of a mixture of atomic hydrogen and methane. The gas can be forced into the plasma phase using strong microwaves or hot filaments [?, ?, ?]. While the hot filament is easy to implement, it has the disadvantage that atoms which are etched from the filament during the growth process are likely to contaminate the diamond. To minimize the introduction of defects other than SiV centers in the diamond, growing diamonds in a microwave plasma is preferred. In it methane molecules dissociate and release carbon. In the presence of a substrate such as iridium containing suitable seeds, carbon can crystalize forming diamond. Figure 1.1 illustrates the setup. In a plasma containing atomic hydrogen, the formation of diamond is favored over the formation of graphite. This is due to the fact that the atomic hydrogen preferentially etches *sp*<sup>2</sup> bonded carbon, i.e. graphite.

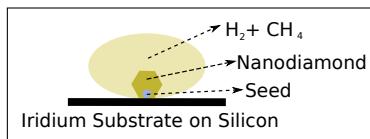


Figure 1.1: Sketch of the CVD method. A hydrogen-carbon plasma acts as a donor for carbon atoms which may crystallize to form diamond onto a suitable iridium substrate. An initial diamond seed facilitates the process.

Typically, single crystal diamond substrates are required to grow single crystal diamond. HPHT substrates are suitable to start a crystallization process. This approach is referred to as homoepitaxial growth. An alternative approach is to utilize non-diamond substrates such as iridium or platinum to trigger heteroepitaxial growth [?, ?]. This method is utilized for

all the CVD nanodiamonds grown and investigated in this thesis. It relies on small diamond crystals deposited in the substrate acting as seeds for the diamond crystallization process<sup>1</sup>. Seed diamond crystals are commercially available, and are usually particles produced by a so-called detonation process. In a detonation process, the high pressure produced by shock-waves of a detonation is used to create very small diamond particles of a size down to a few nanometers.

Growth on a substrate is favored, if the lattice constant of the substrate and the diamond to be grown are similar. The lattice constant of iridium is 0.384 nm [4, 5] and thus close to the lattice constant of diamond with 0.356 nm [6]. Therefore, the diamond was grown on a stratified substrate topped with an iridium layer of 60 nm to 150 nm thickness<sup>2</sup>. The iridium layers themselves were grown onto an yttria-stabilized zirconia (YSZ) buffer layer, which in turn was grown on a silicon wafer [7]. If the lattice constant of the substrate and the diamond are not matched, stress in the diamond lattice is induced. Therefore, the iridium substrate not only facilitates diamond growth, but also reduces unfavorable stress in the nanodiamonds. The detrimental effects of stress on the diamond lattice and its implications with respect to hosted SiV centers are briefly discussed in ??.

To produce nanodiamonds of controlled size, the growth process is stopped when the diamonds grown onto the seed crystals are large enough. If the growth process is continued, the individual crystals grow together to form diamond films. Such diamond films are used as starting material for the wet-milling process described in section 1.3.

One of the advantages of the CVD process is that silicon is incorporated automatically into diamonds, SiV centers can thus be formed *in-situ* as the diamond is grown. The presence of silicon atoms required to be absorbed into the diamond lattice is explained by the plasma etching the edges of the silicon waver underneath the growth substrate. To further increase the silicon content in the chamber, sacrificial silicon can be introduced.

After nanodiamond growth, the nanodiamonds are either investigated directly on the growth substrate or transferred to an ultrasonic bath to obtain a solution which is coated onto other substrates for further exaporation.

In this thesis, two types of nanodiamond samples were investigated. The first batch, henceforth referred to as CVD samples, was grown by the group of M. Schreck, Augsburg University using detonation diamond seeds of a size smaller than 3 nm (produced by the company Microdiamant, product Liquid Diamond monocrystalline, MSY 0-0.03 micron GAF). For the growth process, 1 % of methane was added to the hydrogen environment in the growth chamber. The growth process was performed with a pressure of 30 hPa for 30 min to 60 min, yielding nanodiamonds of a diameter of about 100 nm to 200 nm. A sample of the produced diamonds is given in Figure 1.2.

The second type of samples consist of CVD nanodiamonds grown onto molecular analogs of diamond crystals. A subgroup of these molecular diamonds are called diamondoids and are carbon crystals based on the carbon cage molecule adamantane  $C_{10}H_{16}$ . The molecular diamonds used for this work are adamantane in cyclohexane, mercapto adamantane in cyclohexane, and cyclohexane. Each of these seed crystals was used in different growth processes. During the growth process, either 1 % or 3 % methane was added to the hydrogen plasma and either silicon (Si) or silicon dioxide ( $SiO_2$ ) was injected to promote the formation of *in-situ* incorporated SiV centers, see Table 1.1.

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<sup>1</sup>CVD nanodiamond growth performed by S. Gsell, group of M. Schreck, Augsburg University

<sup>2</sup>Production of the stratified substrate performed by S. Gsell, group of M. Schreck, Augsburg University

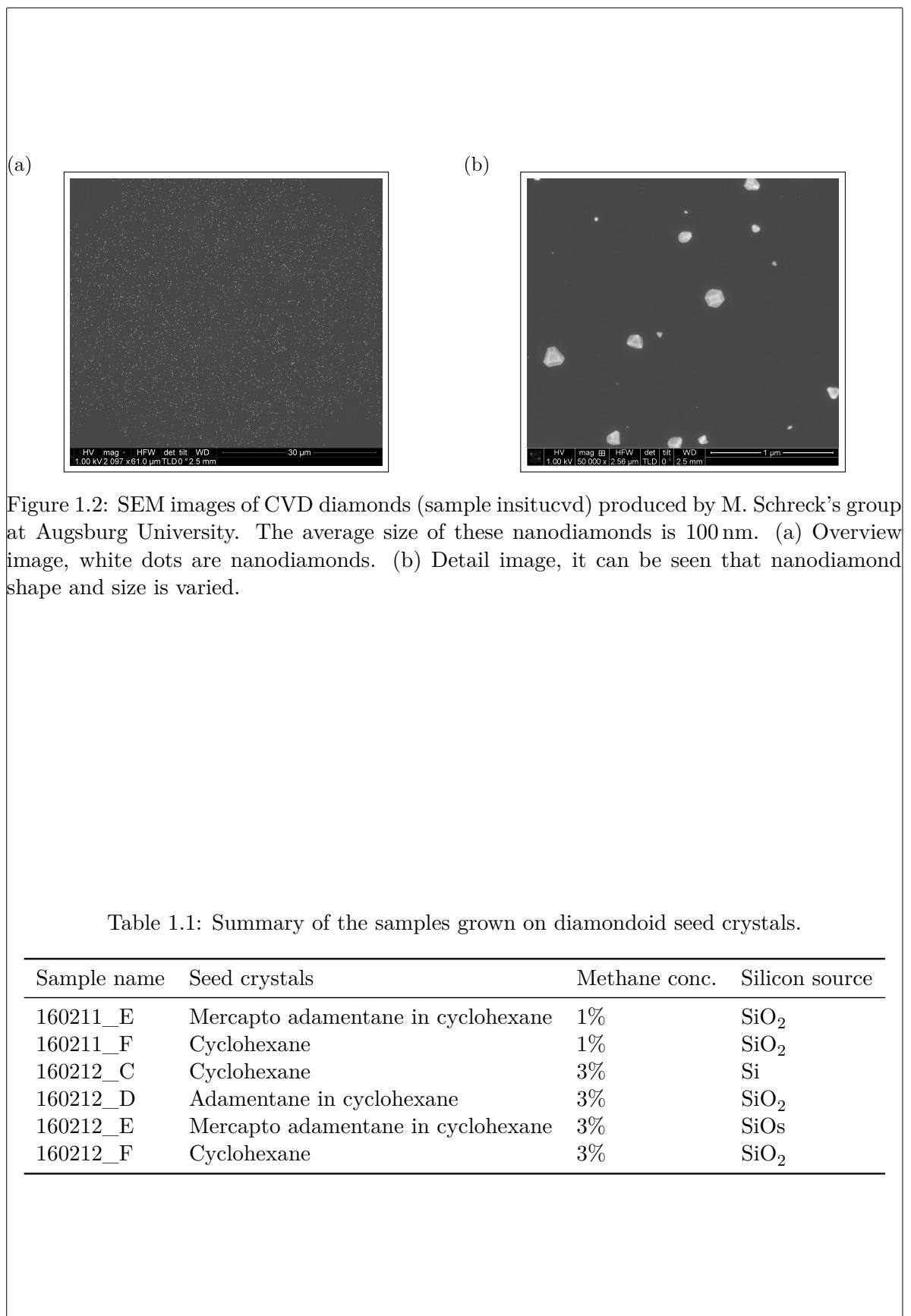


Figure 1.2: SEM images of CVD diamonds (sample insituvcvd) produced by M. Schreck's group at Augsburg University. The average size of these nanodiamonds is 100 nm. (a) Overview image, white dots are nanodiamonds. (b) Detail image, it can be seen that nanodiamond shape and size is varied.

Table 1.1: Summary of the samples grown on diamondoid seed crystals.

Sample name	Seed crystals	Methane conc.	Silicon source
160211_E	Mercapto adamentane in cyclohexane	1%	SiO <sub>2</sub>
160211_F	Cyclohexane	1%	SiO <sub>2</sub>
160212_C	Cyclohexane	3%	Si
160212_D	Adamentane in cyclohexane	3%	SiO <sub>2</sub>
160212_E	Mercapto adamentane in cyclohexane	3%	SiOs
160212_F	Cyclohexane	3%	SiO <sub>2</sub>

### 1.3 Wet-Milled Nanodiamonds

In addition to growing nanodiamonds of a specific size directly via a CVD process, macroscopic diamond starting material can be crushed to obtain small diamond particles. In contrast to nanodiamonds directly grown by a CVD process, the process is divided into two sub-processes: At first a macroscopic diamond is produced using one of the methods introduced in this chapter. Then macroscopic diamond is continuously milled into smaller diamond particles.

The wet-milled nanodiamonds investigated in this thesis were kindly provided by A. Muzha, group of A. Krueger, Julius-Maximilians Universität Würzburg. During the wet-milling process small metal beads in an aqueous solution are driven by the vibrations of a vibrational mill. The moving beads continuously collide with the present diamonds and thus keep breaking them into smaller and smaller particles. A sketch of the process is shown in ???. Steel contamination introduced by the beads can be eliminated in a post-processing treatment with acid.

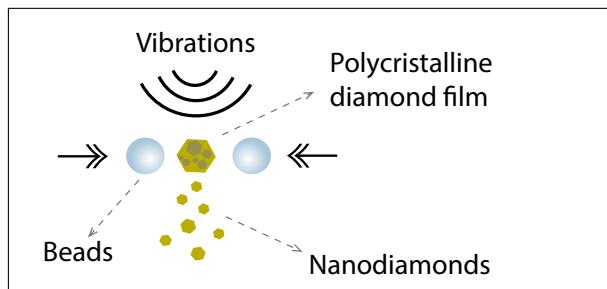


Figure 1.3: Sketch of a wet-milling process in a vibrational mill. A macroscopic polycrystalline nanodiamond is progressively broken down into smaller polycrystalline nanodiamonds by metal beads.

The big advantage of the milling process is that it enables the production of a large quantity of diamond nanoparticles. When producing nanodiamonds directly via a CVD process, the number of produced nanodiamonds in one process scales with the surface of the substrate on which the nanodiamonds are grown. In contrast, the quantity of milled nanodiamonds scales with the volume of the starting material. Another advantage of the wet-milling process is that the nanodiamonds are already in an aqueous solution after milling. Therefore, it can be used to spin-coat the nanodiamonds directly onto any substrate. Samples created in this fashion can directly be used for further investigation and do not require additional processing.

If the starting material for the wet-milling process is a polycrystalline diamond film, it is likely to break along distinct crystal boundaries. However, due to imperfections in the growth process, particles may break down such that progressively smaller milled diamond grain remain polycrystalline. As a result, the final nanodiamonds contain crystal boundaries themselves, indicative of reduced crystal quality. To improve diamond quality nanodiamonds are treated with post-processing steps including annealing in vacuum and oxidation in air. A detailed description of these processes and their effects is given in ??.

In general any diamond, independent of production method, can be used as starting material for the milling process. In the following sections, available samples are distinguished by the respective starting material and milling method.

### 1.3.1 Wet-Milled HPHT Nanodiamonds

We investigated nanodiamonds wet-milled from a HPHT starting material to median sizes of about 45 nm, 80 nm and 260 nm. They were then drop-cast onto an iridium substrate and implanted with  $^{28}\text{Si}^{1+}$ . The implantation itself was performed by D. Rogalla, Ruhr-Universität Bochum (RUBION - Zentrale Einrichtung für Ionenstrahlen und Radionuklide). All HPHT nanodiamonds were oxidized in air at 450 °C for 3 h. Resulting samples available for explorations are designated hphtimp45, hphtimp80, hphtimp260 and listed in Table 1.2.

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### 1.3.2 Wet-Milled CVD Nanodiamonds

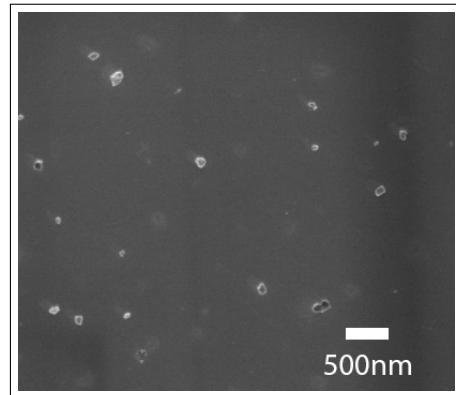


Figure 1.4: Pictures of milled nanodiamonds (sample insitu100) showing the distribution of the nanodiamond crystals on the iridium substrate.

In the following paragraphs, details of the production processes of nanodiamond produced by wet-milling a CVD diamond film in a vibrational mill are given. For an overview of the samples refer to Table 1.2. The starting material for the wet-milled nanodiamonds was a nanocrystalline diamond film [8] directly grown on a silicon wafer by CVD. A microwave hydrogen plasma containing 1% methane was used to grow on purified 5 nm nanodiamond seeds (produced by PlasmaChem). To induce *in-situ* SiV center creation, sacrificial Silicon pieces are placed in the growth chamber. During diamond growth the silicon pieces are etched by the plasma leading to individual atoms being incorporated into the diamond lattice. The diamond film is then wet-milled in a vibrational mill with steel beads. The high amount of steel containment due to the steel beads is removed by extensive acid treatment. We also investigated nanodiamonds milled with silicon nitride beads, and found that the choice of material of the beads does not result in any noticeable difference in dependent measurements. The median diameters of the diamonds are 50 nm, 70 nm and 100 nm as determined by laser diffraction spectroscopy. ?? shows milled nanodiamonds ontop of an iridium substrate. The aqueous solution containing the nanodiamonds is drop cast onto an iridium film on a Silicon substrate. The iridium film of a thickness of 130 nm is grown onto a buffer layer of yttria-stabilized zirconia (YSZ) which in turn is grown onto a Silicon wafer. The iridium surface has the advantage that it acts as an antenna and therefore enhances the collection efficiency of fluorescence light [9]. For a discussion on the properties of the substrate see section 1.4. Post-process treatment consist of annealing in vacuum at 900 °C or consecutive oxidation in air at a temperature of 450 °C, or a combination of the two. The duration for either treatment

method was 3 h to 6 h.

### 1.3.3 Doubly Wet-Milled Implanted Nanodiamonds Implanted With Silicon

In addition to SiV centers that were implanted during diamond growth, we investigated nanodiamonds with SiV centers implanted after diamond growth. We developed a procedure composed of consecutive milling steps to obtain nanodiamonds with an optimal concentration and homogeneous distribution of SiV centers in the host nanodiamonds. Suboptimal concentration of SiV centers in nanodiamonds arises from the fact, that silicon implantation causes the SiV centers to form in a specific depth dependent on the implantation energy. As a consequence, when implanting bulk material most of the diamond is left vacant of SiV centers. Hence, a big portion of nanodiamonds milled from such a bulk material would not host any SiV centers.

To obtain diamond particles with a homogeneous distribution of SiV centers, the process of fabricating implanted nanodiamonds is the following: First, a polycrystalline diamond film (produced by Element Six, electronic grade) is milled to diamond particles a few microns in size. Next, these microdiamonds are spin-coated onto iridium substrates and implanted with  $^{28}\text{Si}^{1+}$ . To eliminate lattice damage and unnecessary vacancies resulting from the implantation process, diamonds were annealed in vacuum and subsequently oxidized. At last, the micrometer sized diamond particles are milled to the desired smaller sizes.

For the milling process in a vibrational mill, a minimum amount of diamond material is necessary. When starting with a diamond film, this threshold is easily reached, however, a big quantity of microdiamonds is needed to meet the requirements. To develop an optimal production procedure of implanted nanodiamonds, we first ran preliminary tests with a small amount of diamond material. After determination of the optimal specifications of the procedure, we production at a larger scale.

For our first experiments, we coated a small amount of microdiamonds milled from the diamond film onto a silicon substrate without an iridium layer. The microdiamonds were then implanted with  $^{28}\text{Si}^{1+}$  at an implantation energy of 1.7 MeV, and fluences of  $10^9 \text{ cm}^{-2}$  to  $10^{12} \text{ cm}^{-2}$ <sup>3</sup>. After implantation, the microdiamonds on the silicon substrate were annealed for 2 h at 900 °C and subsequently oxidized in air for 2 h at 450 °C (for a detailed explanation of the effect of annealing and oxidation refer to ??). Using this procedure, we encountered the problem that the silicon from the substrate sublimated and re-nucleated during annealing. This process caused the diamonds to sink into the silicon surface (Figure 1.5a).

Iridium is less prone to damage by high temperatures and withstands annealing procedures up to our standard annealing temperature of 900 °C without problems. Hence, we repeated the production process in the same manner as described above, but this time, we used an iridium coated silicon wafer as substrate. Thus we circumvented the reported problems of microdiamonds sinking into the substrate surface.

A further technical challenge was the limited space in the vacuum oven used for annealing the microdiamonds. This circumstance capped the size of individual substrates for the microdiamonds to a side length of 2 cm.

Spectroscopic measurements revealed, that microdiamonds implanted with a fluence of fluence

<sup>3</sup>Implantation performed by D. Rogalla, Ruhr-Universität Bochum (RUBION - Zentrale Einrichtung für Ionenstrahlen und Radionuklide).

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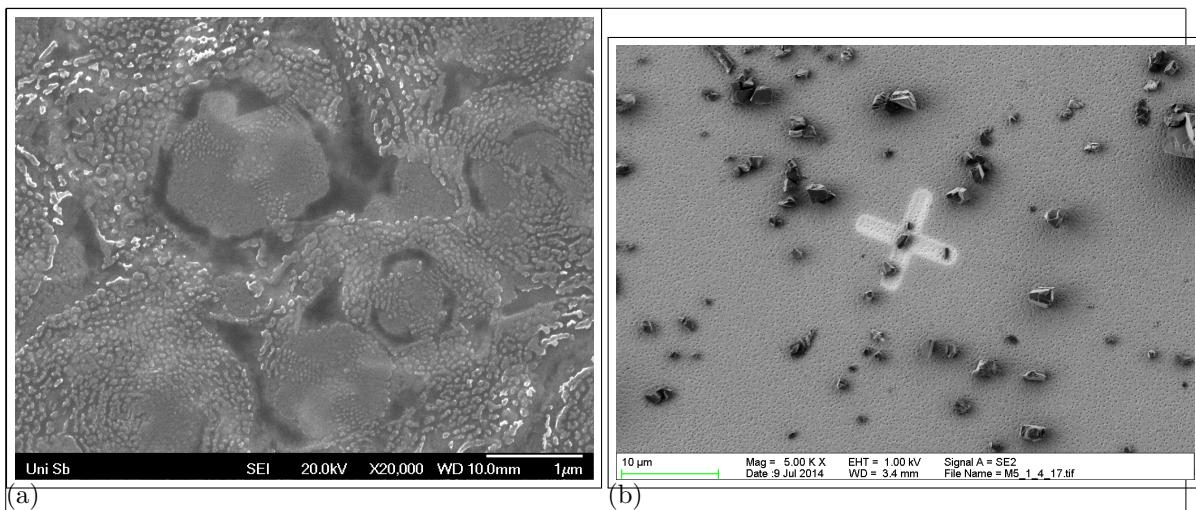


Figure 1.5: (a) SEM picture of nanodiamonds sunken into a silicon substrate after annealing at 900 °C for 3 h. Magnification 20 000. (b) SEM image of the microdiamonds after milling on an iridium substrate, but before implantation. In the middle of the picture, there is a reference cross milled into the iridium substrate with a focused ion beam. Its size amounts to 10  $\mu\text{m}$ . It can be seen, that the microdiamonds exhibit sizes of a few micrometer.

$10^{11} \text{ cm}^{-2}$  was best suited to form a high concentration of SiV centers within the microdiamonds.

Following our results from the preliminary tests, we initiated mass production. The microdiamonds were directly spin-coated onto 20 iridium substrates of a side length of 2 cm which yielded densely coated samples(??). After spin-coating the samples were implanted with  $^{28}\text{Si}^{1+}$  (implantation energy 900 keV, fluence  $10^{11} \text{ cm}^{-2}$ )<sup>4</sup>. The microdiamonds were then annealed in vacuum for 3 h at 900 °C and oxidized in air for 3 h at 450 °C. As mentioned, a bottleneck of the production was the limited space of the vaccuum oven. Only 3 samples fit into it at once, hence it was necessary to recursively repeat the annealing process until all samples were annealed.

At last, the microdiamonds were milled again to sizes of about 40 nm, 45 nm, 240 nm and 260 nm. The diamonds of sizes 40 nm and 240 nm were directly annealed in vacuum at 1200 °C<sup>5</sup> without prior spin-coating.

## 1.4 Iridium Substrate

As mentioned in section 1.2, growth on a substrate is favored, if the lattice constant of the substrate and the diamond to be grown are similar. Therefore, for CVD nanodiamond growth a multilayered substrate was prepared in a heteroepitaxial growth process. This substrate consists of a silicon wafer on the bottom, a yttria-stabilized zirconia buffer layer in the middle and an iridium layer on top. Initially, whole 4 in wafers were prepared with a multilayered structure. After the heteroepitaxial growth processes, the wafer is cut to squares with a side

<sup>4</sup>Implantation performed by J. Klug at rubitec - Gesellschaft für Innovation und Technologie der Ruhr-Universität Bochum mbH.

<sup>5</sup>performed by A. Muzha, group of A. Krueger, Julius-Maximilians Universität Würzburg

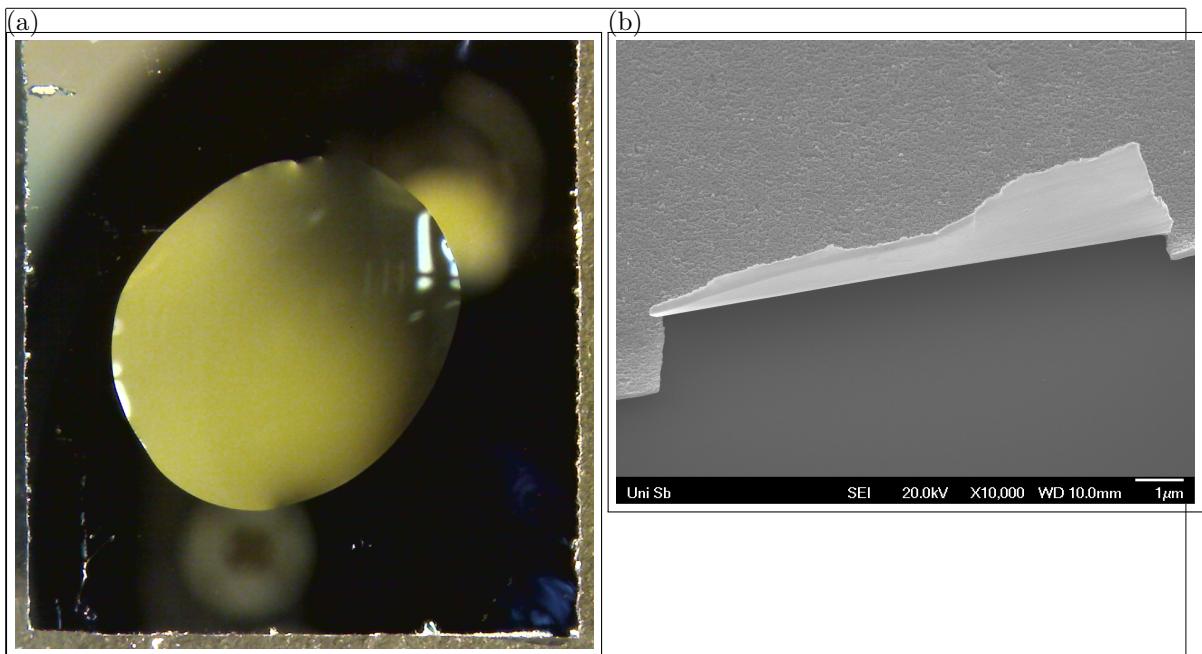


Figure 1.6: (a) Microscope image of a drop of water on an iridium substrate cleaned with Piranha etch. This picture was used to estimate the contact angle. (b) SEM picture of an 60 nm iridium layer that peeled off the substrate after an ultrasonic bath.

length of 1 cm to fit into all machinery used during experiments, such as the spin-coater, the confocal setup, the focussed ion beam and the SEM.

Apart from the favorable lattice constant, iridium exhibits further advantages which lead us to coating the wet-milled diamonds onto these same multilayered substrates:

- The high hydrophilicity of iridium enhances the homogeneity of the nanodiamonds on the substrate after spin-coating or drop-casting. The hydrophilicity is further improved by treating the substrate in a Piranha etch (50:50 mixture of sulfuric acid  $H_2SO_4$  and hydrogen peroxide  $H_2O_2$ ) by removing oxide layers on the surface. The treatment with Piranha etch also has the advantage that all organic contamination is removed. Measurements after applying the Piranha cleaning yielded an estimation of the contact angle of slightly more than one degree. To determine the contact angle with water, the volume of a water drop is compared to the surface it covers after dropping it onto an iridium substrate, see Figure 1.6a. From that an estimation of the contact angle is deduced.
- During the post-processing steps, it is of major importance, that the substrate can withstand high temperatures. As described in ??, during the preliminary tests with implanted nanodiamonds on a silicon substrate we encountered difficulties with diamonds on a silicon substrate as they sunk into the surface after annealing. In contrast, iridium withstands the high temperatures required for annealing without damage.
- Iridium acts both as a mirror and as an antenna for the fluorescence light emitted by the SiV center [9]. Therefore, the collection efficiency of the fluorescence light is enhanced.

At this point we comment upon a minor complication we experienced when using iridium

surfaces. If the iridium layer is too thin, it tends to peel off the substrate, see Figure 1.6b. We encountered this problem during a cleaning procedure in the ultrasonic bath. However, this disadvantage is easily circumvented by using a thicker iridium layer. For our measurements, we used an iridium layer of a thickness of 130 nm, for which we did not encounter any adhesion problems.

#### 1.4.1 Substrate Preparation

Prior to drop-casting or spin-coating a diamond solution onto the iridium substrate, the substrate is cleaned. The standard cleaning procedure is started with treating the substrate in diverse chemicals in an ultrasonic bath. Each step is carried out for a duration of 3 min to 7 min, approximately adjusted to the amount of contamination present:

- Distilled water with a drop of dishwasher detergent: removes coarse contamination off the substrate.
- Isopropanol (purity 99.9 % p.a.): dissolves oils without leaving oil traces as compared to ethanol
- Acetone (purity 99.9 % p.a.): a heavy-duty degreaser, also dissolves many plastics
- Distilled water: Acetone heavily reacts with the later applied Piranha solution, potentially causing explosions. Therefore, any acetone residue has to be removed. As acetone is water-soluble, distilled water is used to remove the residue.

Thereafter, the substrates are put into a Piranha solution (50 % sulfuric acid H<sub>2</sub>SO<sub>4</sub>, 50 % hydrogen peroxide H<sub>2</sub>O<sub>2</sub>) to enhance the surface hydrophilicity and therefore obtain a homogeneous distribution of diamonds on the surface. They are then put again into distilled water and blow-dried with compressed air to avoid residue from the water.

The substrates were then either drop-casted or spin-coated with aqueous diamond solutions. For the former, the substrates are heated on a heating plate to a temperature of 60 °C. While the substrates are permanently heated by the heating plate, drops of a volume of about 5 µL are dropped onto the substrate surface. If substantially more than 5 µL is needed, then several drops of about 5 µL are dropped onto the substrate consecutively. A single large drop is ill-advised since the solution would flow off the substrate before drying.

For spin-coating, an on-site built spin coater was used. Therefore the spin-coater is pre-set to velocity of 2500 rpm. Then drops of 5 µL are dropped on the substrate and the spin-coater is immediately turned on. After about 3 min, the drop has dried. If more than 5 µL of diamond solution is wanted, the spin-coating is repeated until the desired amount is reached.

## 1.5 Listing of Investigated Samples

Table 1.2: Listing of investigated wet-milled samples. While this is a subset of all the samples investigated throughout the work of this thesis, these samples are the most thoroughly investigated ones. They are the basis for the results reported in ???. The first column indicates the names of the samples, the second the mean diameter of the nanodiamonds, and the third designates how the silicon was incorporated into the diamond.

Sample name	Size	Si incorporation	Post-processing
hphtimp45	45 nm	<i>implanted</i>	oxidized in air at 450 °C
hphtimp80	80 nm	<i>in-situ</i>	annealed in vacuum for 1 h at 1000 °C and at 900 °C for 3 h
hphtimp260	260 nm	<i>in-situ</i>	oxidized in air at 450 °C
insitu50	50 nm	<i>in-situ</i>	series of individual samples with diverse post-processing steps
insitu70	70 nm	<i>in-situ</i>	series of individual samples with diverse post-processing steps
insitu70n	70 nm	<i>in-situ</i>	no post-processing subset of insitu70
insitu70o	70 nm	<i>in-situ</i>	oxidized in air at 450 °C subset of insitu70
insitu100	100 nm	<i>in-situ</i>	series of individual samples with diverse post-processing steps
insitu100ao	100 nm	<i>in-situ</i>	annealed in vacuum at 900 °C, consecutively oxidized in air at 450 °C subset of insitu100
implanted250ao	250 nm	implanted	annealed in vacuum at 900 °C, consecutively oxidized in air at 450 °C
hphtimp260	40 nm	implanted	annealed in vacuum at 1200 °C, consecutively oxidized in air at 450 °C
hphtimp260	50 nm	implanted	oxidized in air at 450 °C
hphtimp260	240 nm	implanted	annealed in vacuum at 1200 °C, consecutively oxidized in air at 450 °C
hphtimp260	260 nm	implanted	oxidized in air at 450 °C

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