

## APPENDIX A

### Error analysis

#### **A1. Standard Deviation**

$$(\Delta S) = \left( \frac{\sum_{i=1}^N (x_i - \bar{x})^2}{N(N-1)} \right)^{1/2}$$

where  $\bar{x}$ ,  $x_i$  and  $N$  are mean data, data and number of data respectively.

#### **A2. Standard Error**

Consider  $S$  as a function of certain variable  $x, y, \dots$ ,

where,  $S = f(x, y, \dots)$ ,

$$\left( \frac{\Delta S}{S} \right) = \left[ \left( \frac{\Delta x}{x} \right)^2 + \left( \frac{\Delta y}{y} \right)^2 + \dots \right]^{1/2},$$

where  $x, y, \dots$  is various variable.

#### **A3. Error of film thickness, $\Delta d$**

The film thickness,  $d$

$$d = \left( \frac{m\lambda}{2n(\lambda)} \right)$$

The  $\Delta d$  is,  $\left( \frac{\Delta d}{d} \right)^2 = \left( \frac{\Delta \lambda}{\lambda} \right)^2 + \left( \frac{\Delta n(\lambda)}{n(\lambda)} \right)^2$ ,

where,  $\Delta \lambda = 2 \text{ nm}$ ,  $\Delta n(\lambda) = 0.01$ ,  $\Delta d = 5 \text{ nm}$ .

#### **A4. Error of optical band gap, $\Delta E_g$**

The optical band gap is determined by,

$$(\alpha h\nu) = B (h\nu - E_g)^2,$$

where,  $E_g = \left( \frac{c}{m} \right)$ ,

c and m are the intercept and slope for the plot of  $(\alpha h\nu)$  versus E.

The  $\Delta E_g$  is,  $\left( \frac{\Delta E_g}{E_g} \right)^2 = \left( \frac{\Delta c}{c} \right)^2 + \left( \frac{\Delta m}{m} \right)^2$

$$\Delta E_g = 0.05 \text{ eV}$$

The error of c and m is obtained using regression in data analysis program in Microsoft Excel.

#### **A5. Error of Hydrogen Content, $\Delta C_H$**

Error in the hydrogen content was obtained from the FTIR analysis. The equation for integrated intensity, I and area under the curve, S is given. The hydrogen concentration,  $C_H$  or H%,

$$C_H = \frac{N_H}{N_C} \times 100\%$$

$$\left( \frac{\Delta C_H}{C_H} \right)^2 = \left( \frac{\Delta N_H}{N_H} \right)^2 + \left( \frac{\Delta N_C}{N_C} \right)^2$$

$$\Delta C_H = 5\%, \text{ where, } \Delta N_H = 0.01, \Delta N_C = 0.01 \times 10^{23}$$

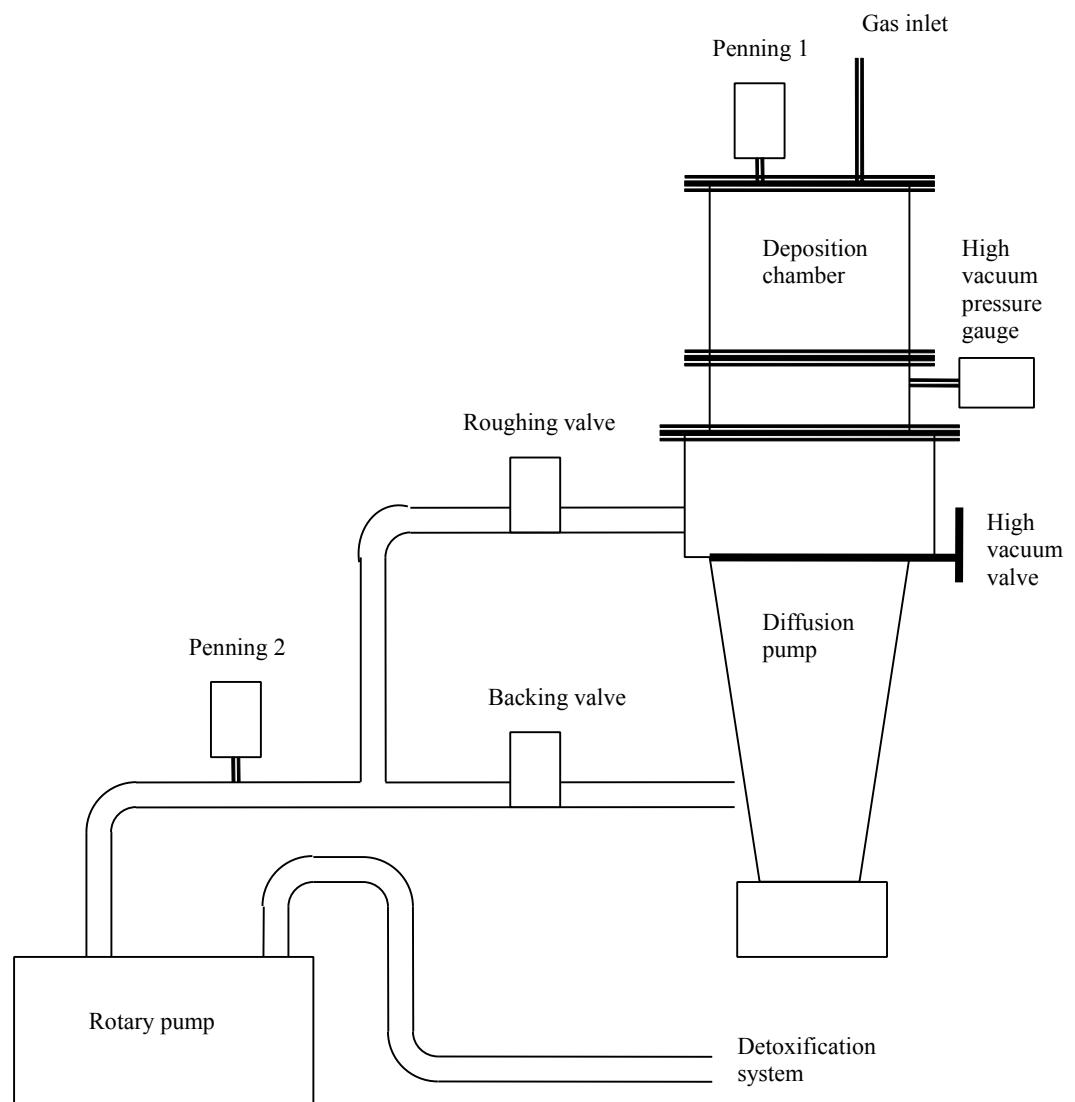
## **APPENDIX B**

### **Deposition System**

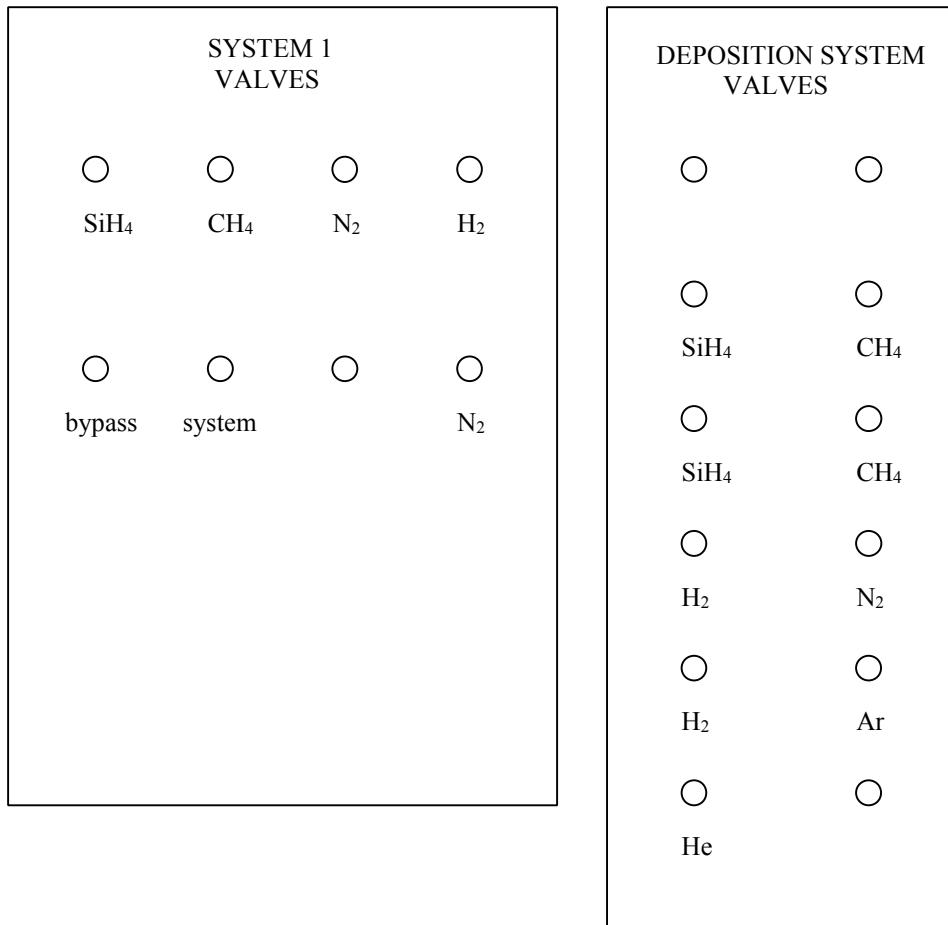
#### **B1. Deposition procedure.**

##### **VACUUMING THE DEPOSITION CHAMBER**

1. CLOSE all valves.
2. ON Rotary Pump, Penning and Pirani pressure meter.
3. OPEN roughing valve, wait until the chamber pressure or Penning meter,  $P_e$  is  $\sim 1 \times 10^{-3}$  mbar.
4. OPEN backing valve, wait until  $P_e$  is  $\sim 1 \times 10^{-3}$  mbar.
5. Checking the gas lines:
  - CLOSE backing valve.
  - OPEN the gas lines involved to vacuum the gas lines.
  - Make sure  $P_e$  is  $\sim 1 \times 10^{-3}$  mbar.
6. Flow water through the coil around the diffusion pump. Make sure there is water flowing at the end of the line. Switch on the fan.
7. OPEN backing valve.
8. ON diffusion pump, wait for about 15 minutes to heat.
9. CLOSE roughing valve.
10. OPEN high vacuum valve, wait until HV display at S1 is  $\sim 1 \times 10^{-5}$  mbar.
11. ON the substrate heater, wait for the pre-set temperature.
12. Check the base pressure, CLOSE high vacuum valve.
13. OFF diffusion pump.
14. OPEN roughing valve. (backing valve is kept open, close only when diffusion pump has cooled down.)
15. ON OSK, check voltage (zero).
16. FLOW the deposition gas, pressure goes down,  $P_e \sim 1 \times 10^{-1}$  mbar. Adjust the roughing valve to control the deposition pressure.
17. READY to deposit. ON power, deposition starts.



## **DEPOSITION USING SILANE GAS**



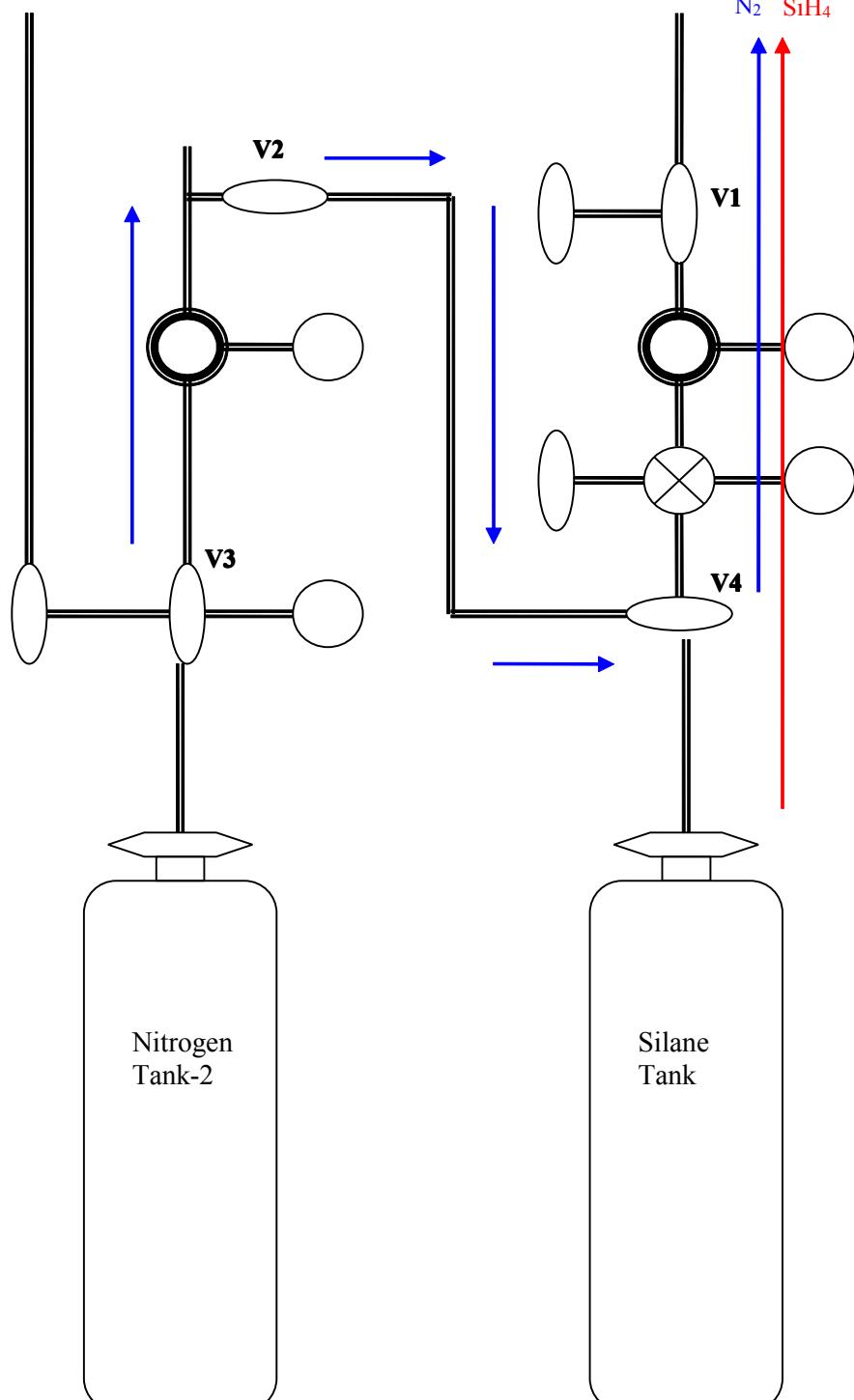
### **1. Vacuuming the gas lines**

1. Turn on the metering valve. Wait until pressure is  $10^{-3}$ .
2. On the control panel of System-1:-
  1. Turn on the system valve. Wait until pressure is  $10^{-3}$ .
  2. Turn on the SiH<sub>4</sub> valve. Wait until pressure is  $10^{-3}$ .
3. On the control panel of Deposition System:-
  1. Turn on the SiH<sub>4</sub> valve. Wait until pressure is  $10^{-3}$ .

## SAFETY CABINET

To gas release system

To deposition system

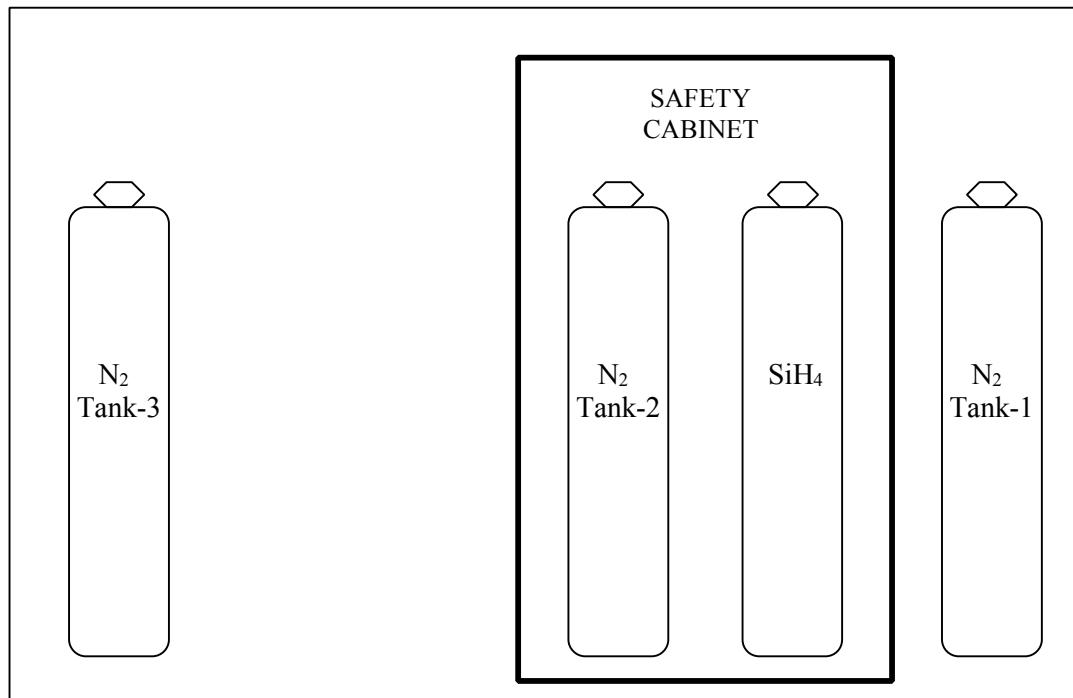


## **2. Purging with N<sub>2</sub> gas**

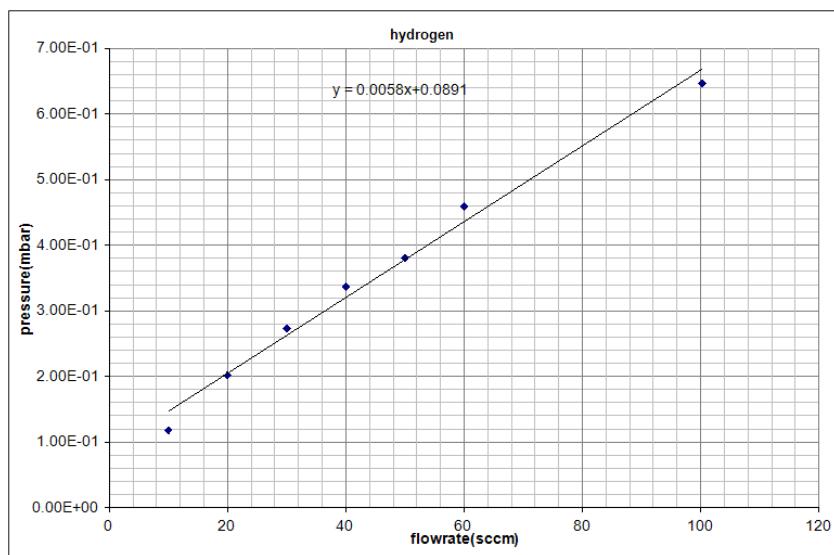
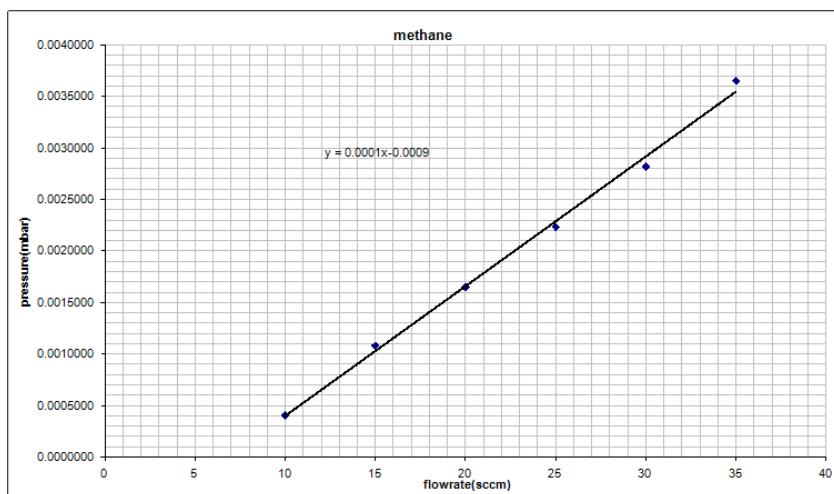
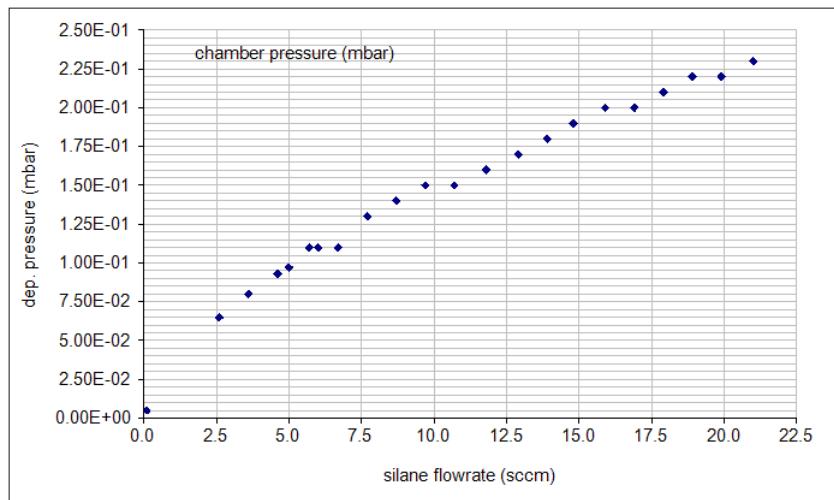
1. Switch on the main switch.
2. Switch on the switch on the switch board, including the emergency button. Make sure the green light turns on.
3. Turn on the N<sub>2</sub> tank outside the cabinet (Nitrogen tank-1)
4. Turn on V1, V2 and V3.
5. Wait until pressure is 10<sup>-3</sup>.
  
6. Turn off the metering valve.
7. Turn on(and off) the N<sub>2</sub> gas inside the cabinet (Nitrogen tank-2).
8. Turn on the metering valve slowly. Silane gas line is now being purged with nitrogen gas.
9. Wait until pressure is 10<sup>-3</sup>.
10. Wait until the released N<sub>2</sub> gas is finished (indicated at the volume meter).

## **3. Flowing the SiH<sub>4</sub> gas into the deposition chamber.**

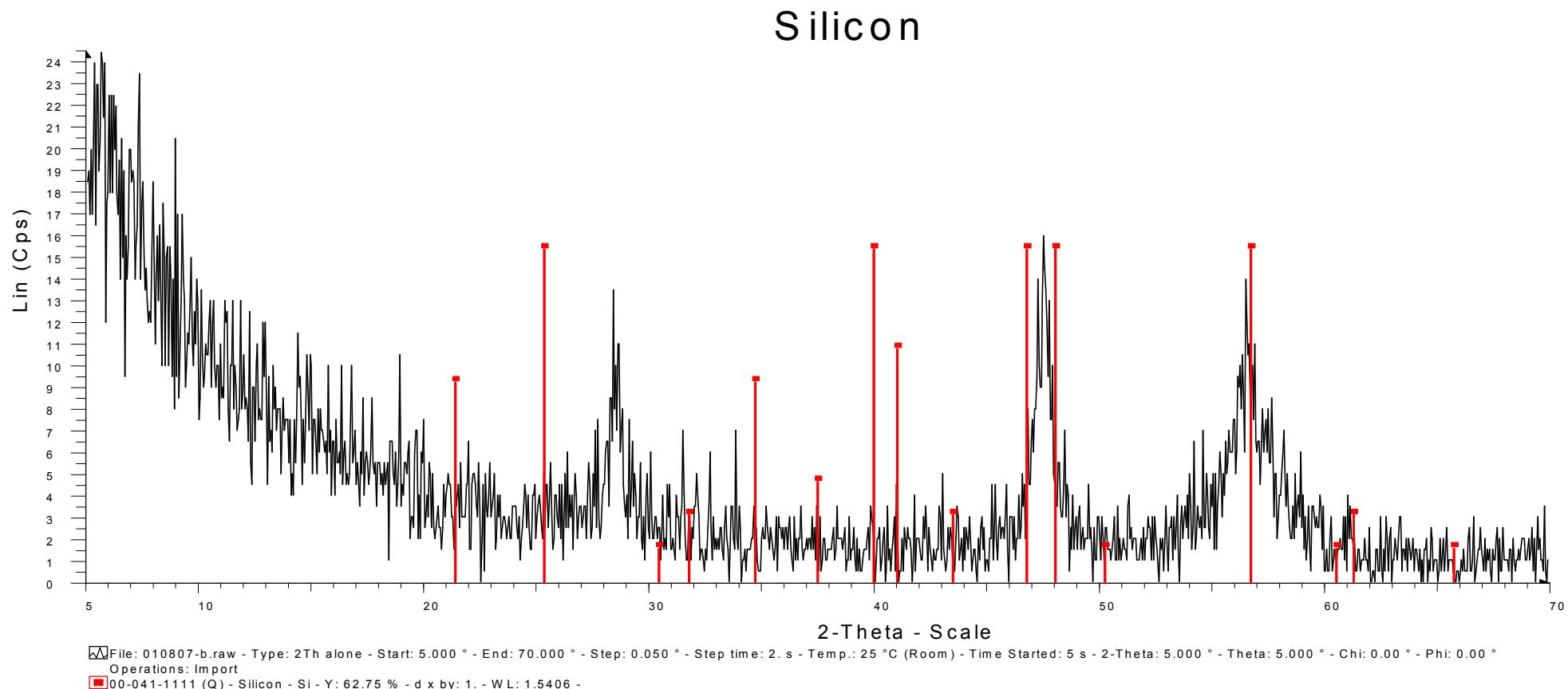
1. Close the backing valve.
2. Turn off the metering valve.
3. Turn on the N<sub>2</sub> gas for the deposition chamber (Nitrogen tank-3)
4. Turn on the N<sub>2</sub> valve at control panel of Deposition System.
5. Turn on the lower N<sub>2</sub> valve at control panel of System-1.
6. Turn on the by-pass valve at control panel of System-1.
7. Turn off V2 and V3.
8. Turn on V4.
9. Turn on(and off) the SiH<sub>4</sub> valve until the pressure meter at the tank is ~400 Psi.  
Now silane gas is in the line.
10. Turn on the metering valve until desired pressure is reached.
11. Switch on the current and voltage. DEPOSITION STARTS.



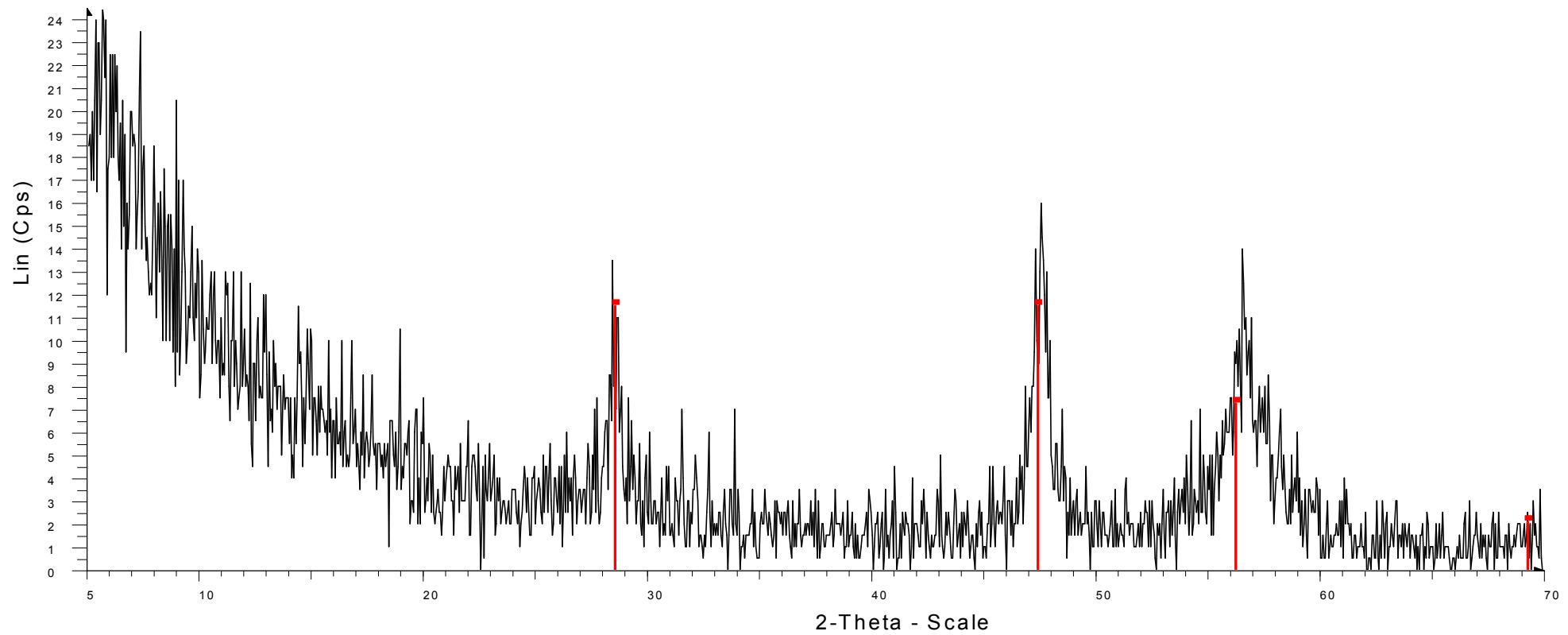
## B2. Performance chart for deposition chamber.



### B3. Example of XRD spectrum for Silicon and Silicon Carbide

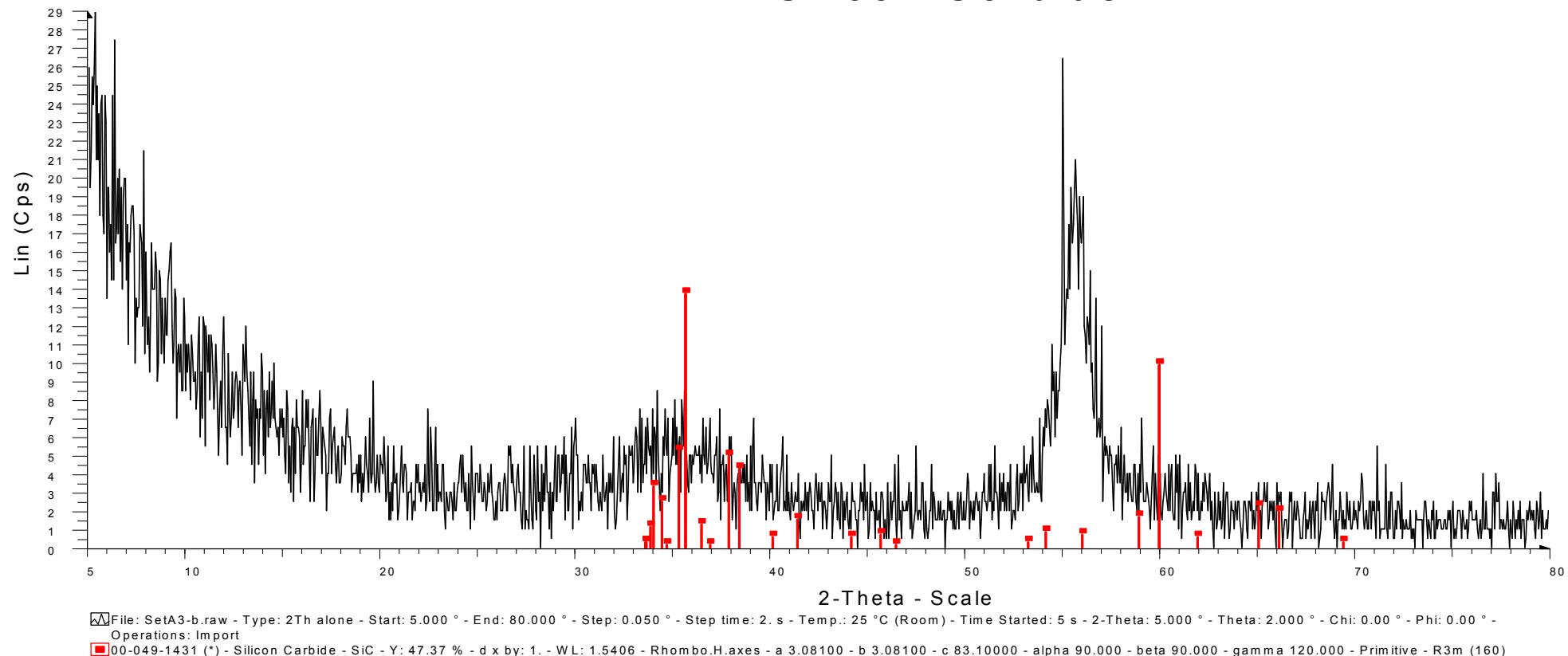


# Silicon

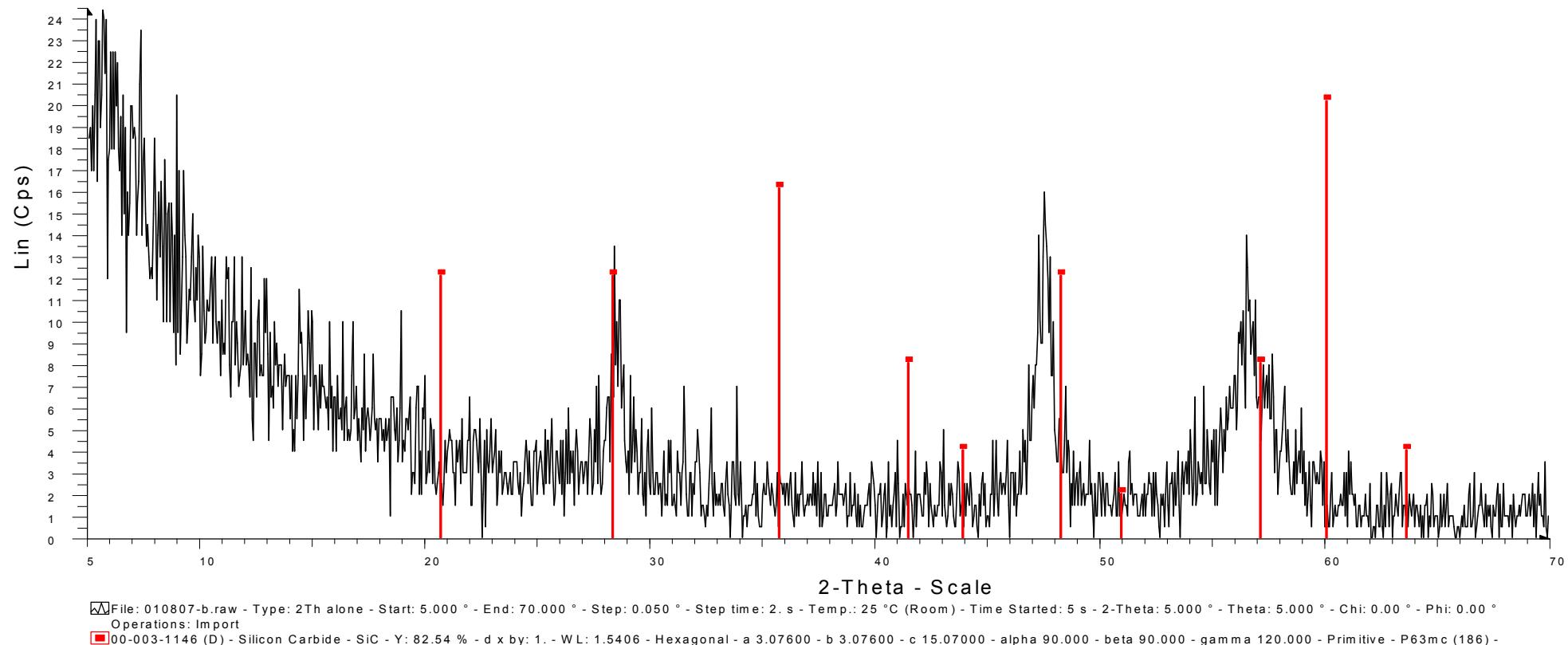


File: 010807-b.raw - Type: 2Th alone - Start: 5.000 ° - End: 70.000 ° - Step: 0.050 ° - Step time: 2. s - Temp.: 25 °C (Room) - Time Started: 5 s - 2-Theta: 5.000 ° - Theta: 5.000 ° - Chi: 0.00 ° - Phi: 0.00 °  
Operations: Import  
00-001-0791 (D) - Silicon - Si - Y: 46.96 % - d x by: 1. - WL: 1.5406 - Cubic - a 5.42000 - b 5.42000 - c 5.42000 - alpha 90.000 - beta 90.000 - gamma 90.000 - Face-centered - Fd-3m (227) - 8 - 159.220

## Silicon Carbide



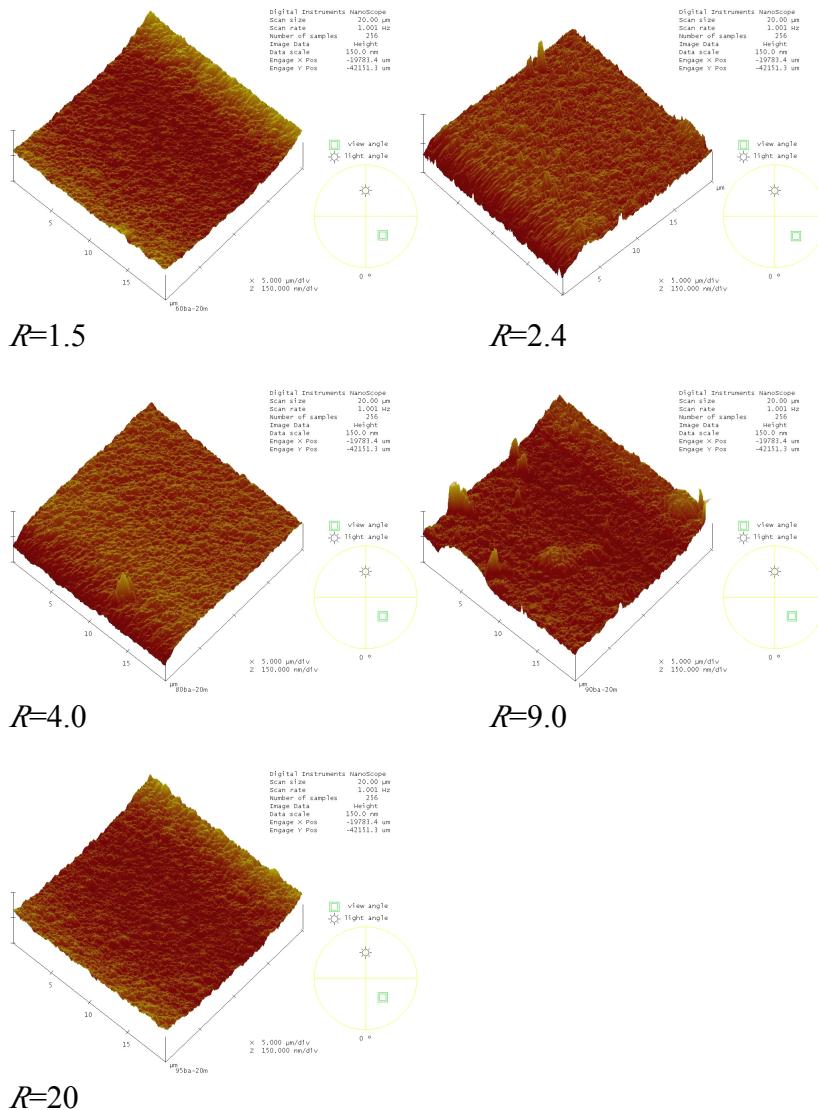
## Silicon Carbide



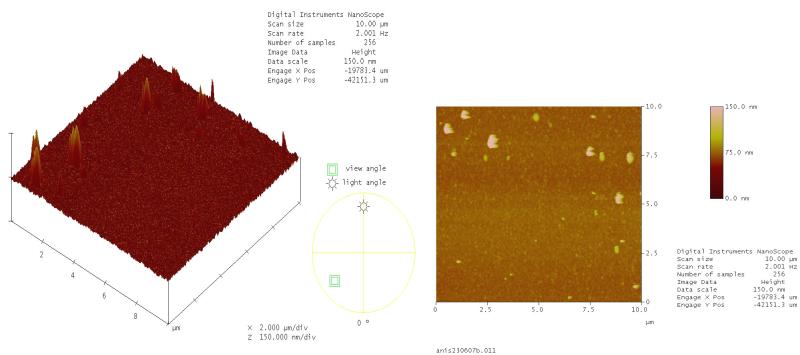
## APPENDIX C

### Morphology studies

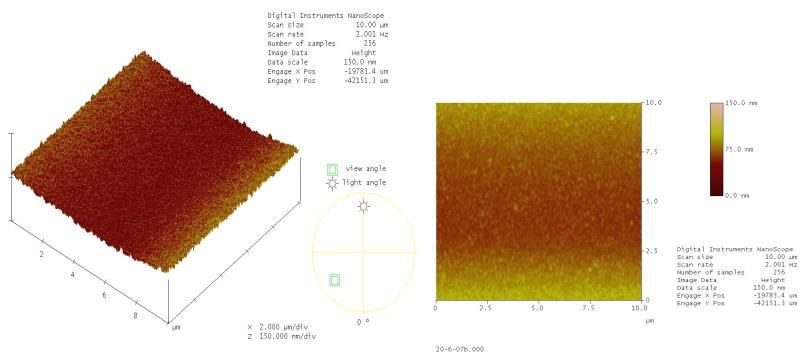
#### C1. AFM images of SiC thin films prepared by RF-PECVD technique.



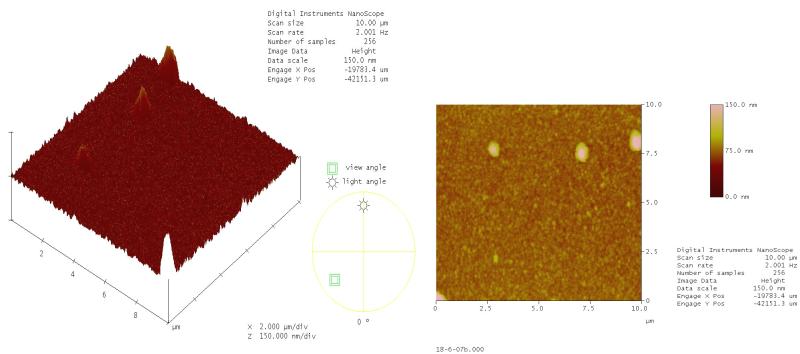
## C2. (a)AFM images of SiC thin films prepared by DC-PECVD technique.



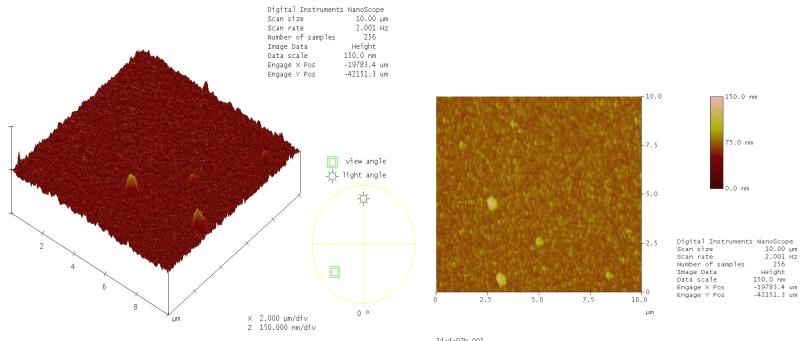
$R=2$



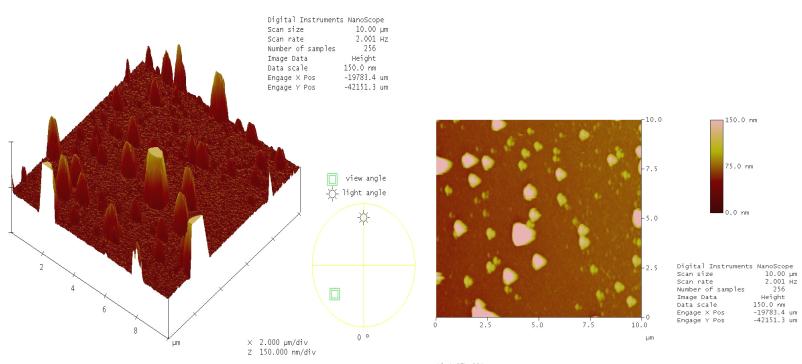
$R=4$



$R=6$

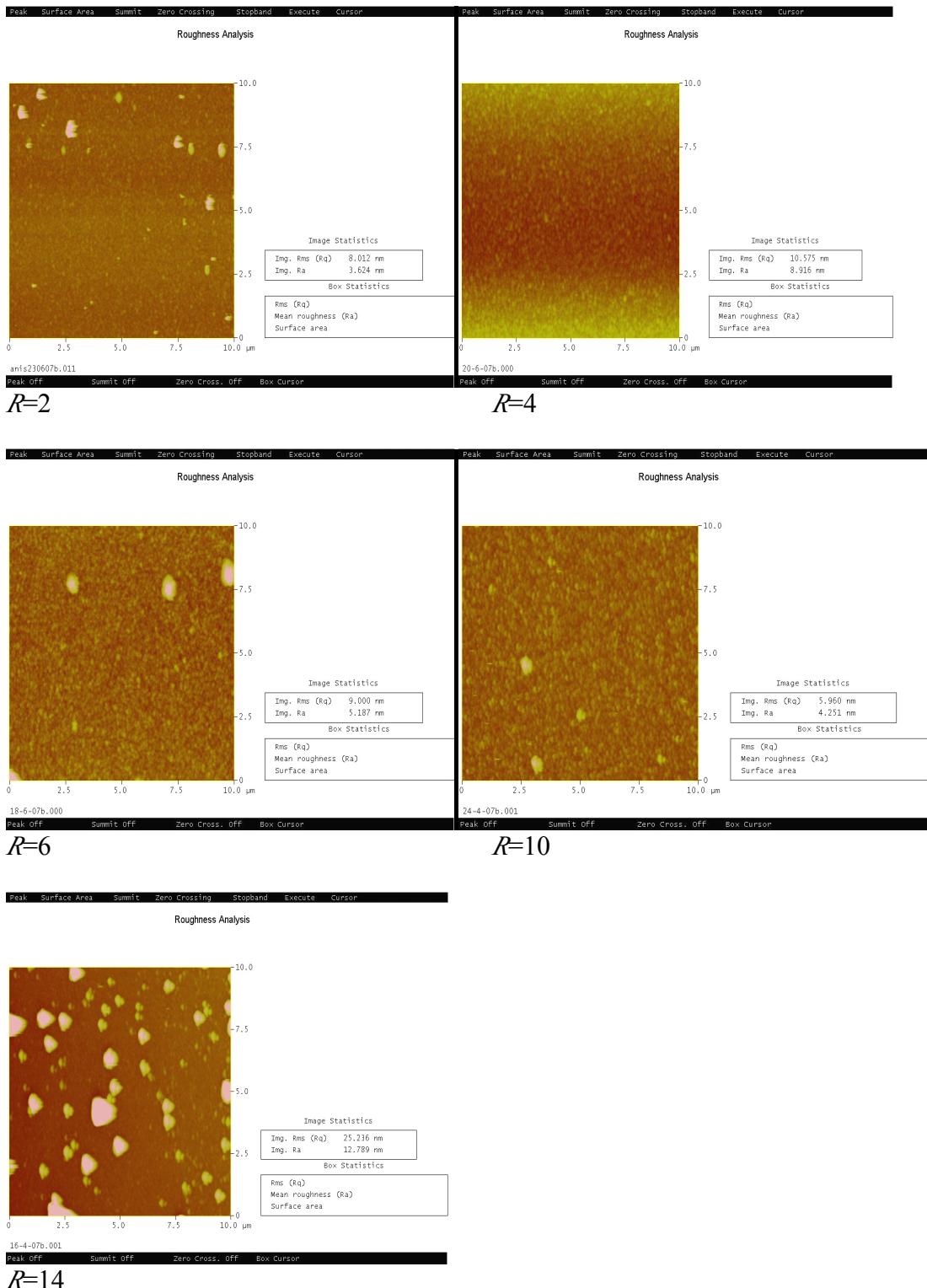


$R=10$

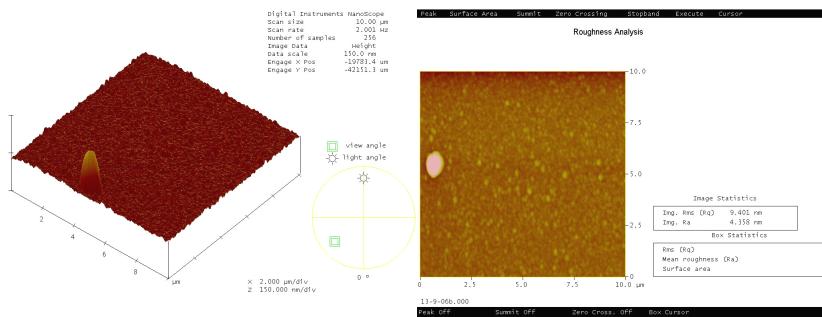


$R=14$

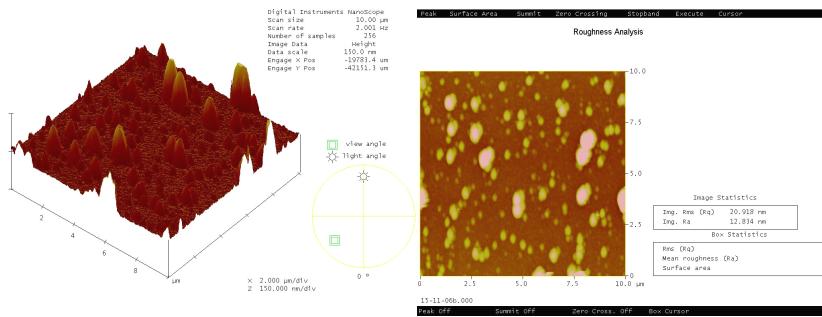
## C2. (b)Roughness analysis of SiC thin films prepared by DC-PECVD technique.



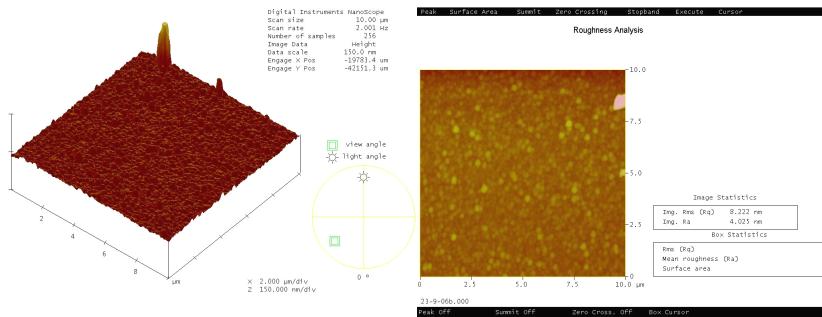
### C3. AFM images of SiC thin films prepared by HW-CVD technique.



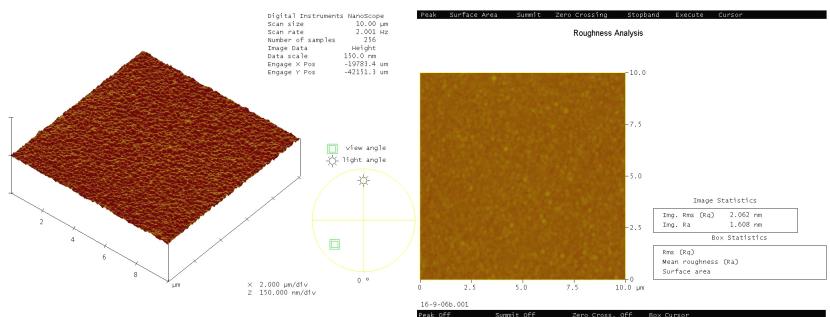
$R=2$



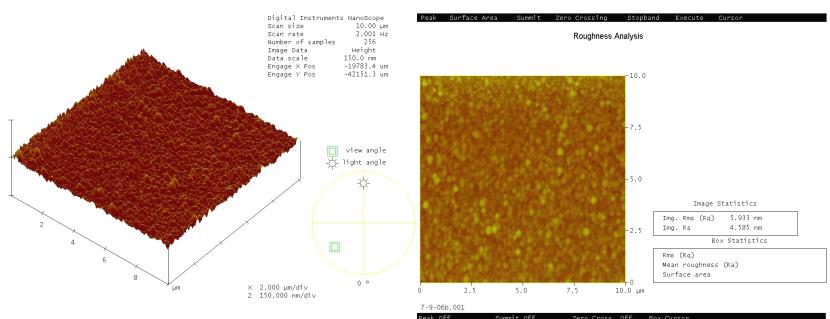
$R=4$



$R=6$

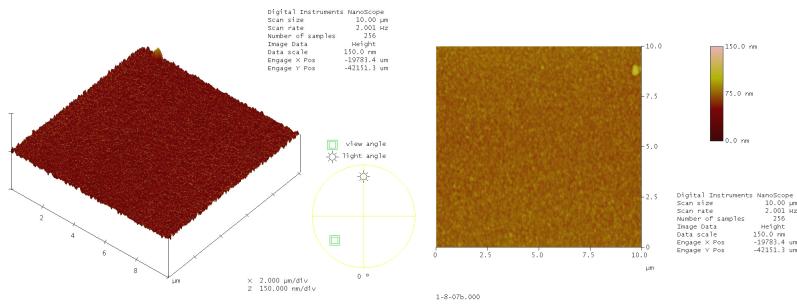


$R=10$

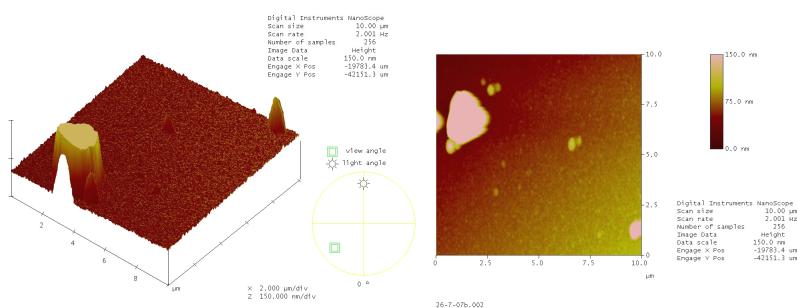


$R=14$

**C4. (a)AFM images of SiC thin films prepared by hybrid HW-PECVD technique with hydrogen dilution.**

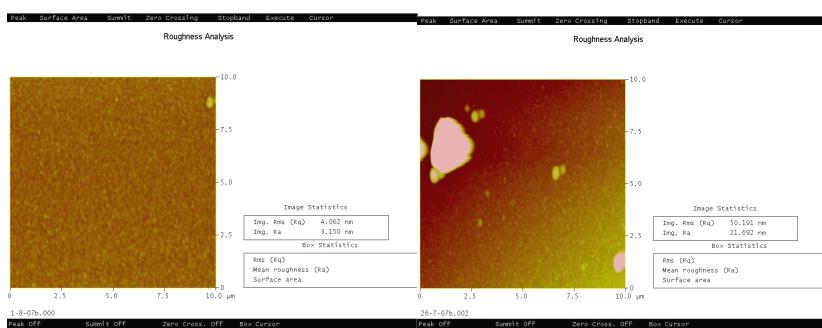


$R=2$  with hydrogen dilution.



$R=14$  with hydrogen dilution.

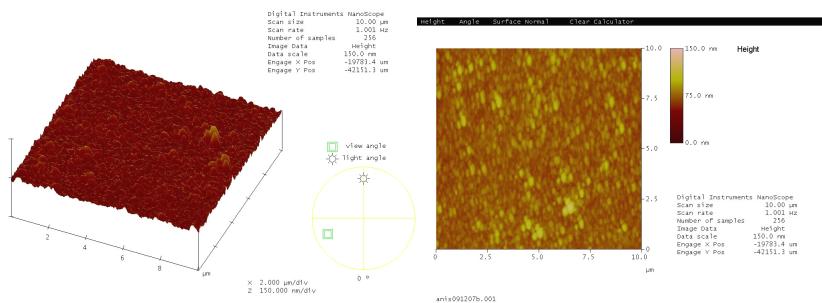
**C4. (b)Roughness analysis of SiC thin films prepared by hybrid HW-PECVD technique with hydrogen dilution.**



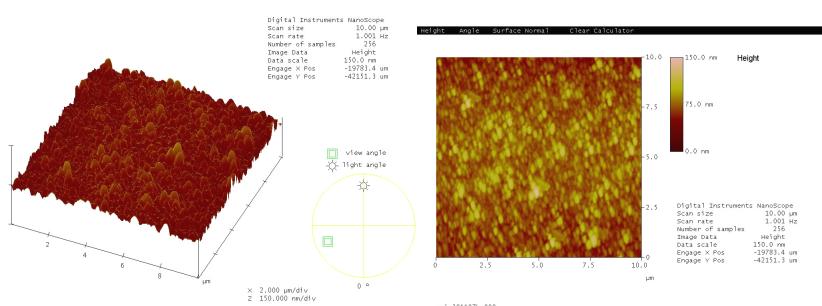
$R=2$  with hydrogen dilution.

$R=14$  with hydrogen dilution.

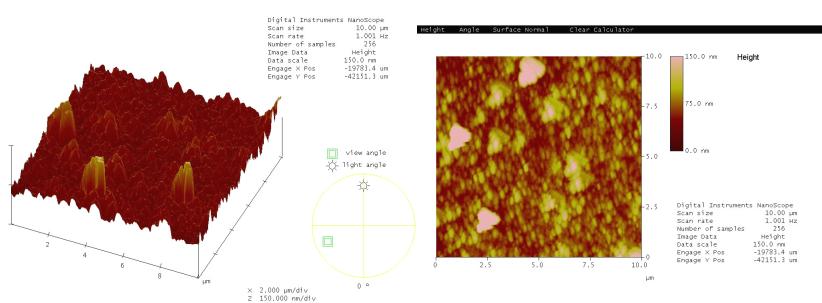
## C5. AFM images of SiC thin films prepared by hybrid HW-PECVD technique with hydrogen surface treatment.



Treatment time = 3 minutes



Treatment time = 2 minutes



Treatment time = 1 minute