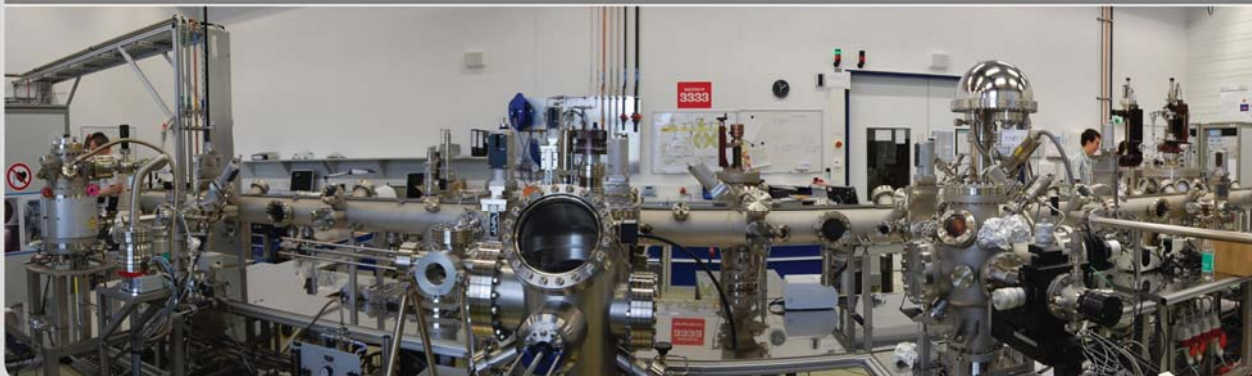


In situ X-Ray Reflectivity measurements during Sputtering of Vanadium Carbide thin films

M.Kaufholz¹, B. Krause¹, S. Kotapati¹, M.Stüber², S.Ulrich² and T. Baumbach^{1,3}

¹ Institut für Synchrotronstrahlung, Karlsruher Institut für Technologie (KIT), ² Institut für Angewandte Materialien - Angewandte Werkstoffphysik, Karlsruher Institut für Technologie (KIT), ³ ANKA, Karlsruher Institut für Technologie (KIT)

ANKA / Institut für Synchrotronstrahlung (ISS)



KIT – University of the State of Baden-Württemberg and
National Large-scale Research Center of the Helmholtz Association

www.kit.edu

Content

■ Motivation

■ *In situ* X-Ray Reflectivity

■ Three Examples:

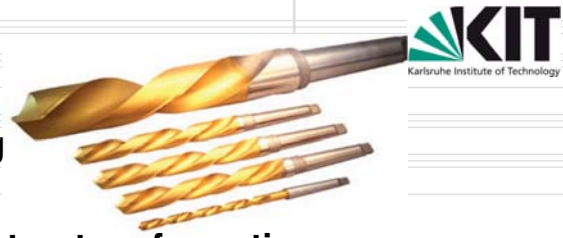
- *In situ* XRR at different DC Power
- *In situ* XRR at different Growth Temperatures
- Interruption of Deposition

■ Summary & Outlook

Motivation

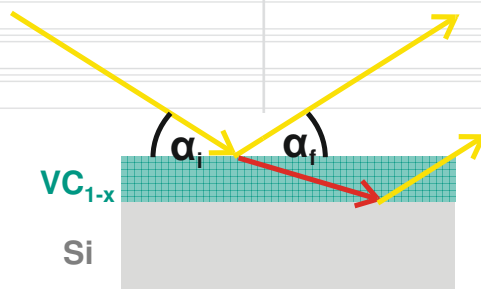
Vanadium Carbide (VC_{1-x})

- Growth of thin films by Sputtering
 - Hard coating material for tools
- deposition conditions** and **microstructure formation** define **mechanical properties**
- **Understand growth process depending on sputtering conditions**
- **Investigation needs suitable methods**
- **nondestructive** monitoring of growth process
 - resolution in **sub-nanometer scale**
 - compatibility with the **gas atmosphere**
 - investigation of
 - **polycrystalline** material
 - **high deposition rates** (0.22 nm/s @ DC Power 200 W)

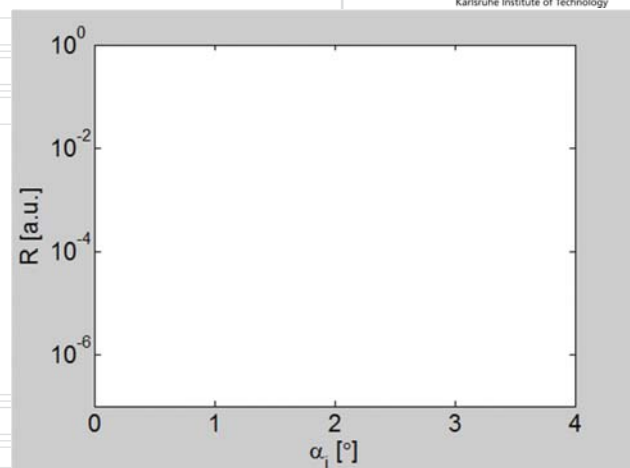


➔ **In situ X-Ray Reflectivity**

Basics of X-Ray Reflectivity



- **Electron density** ('Critical Angle')
- **Thickness** ('Kiessig fringes')
- **Roughness** ('Slope') [1]
- Description by **Parratt-Algorithm** [2]
 - Fully dynamical description of XRR



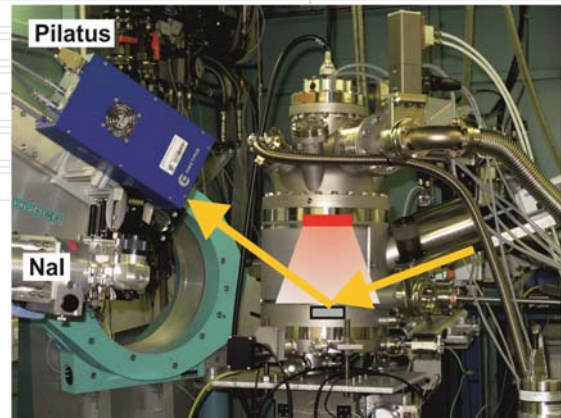
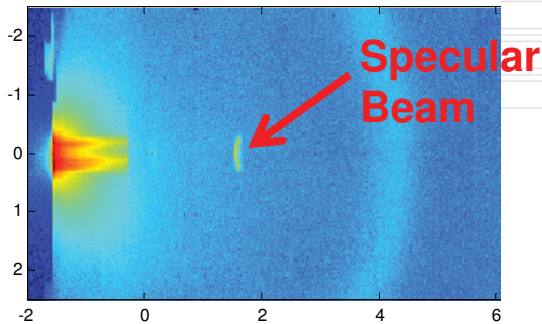
➔ **Two options** to measure *in situ* XRR

1. Full angular range XRR
2. XRR at a fixed angular position

[1] Pietsch, Holy, Baumbach, *High Resolution X-Ray Scattering from thin films and lateral Nanostructures*, Springer 2004

[4] Parratt, *Phys. Rev.* 95, 2, p. 359-369,(1954)

Experimental Setup



■ Setup @ MPI-Beamline:

- Energy: 10 keV
- Beamsize: 300 μ m x 200 μ m
- Optics
 - Resolution in q_z : $\sim 0.005 \text{ \AA}^{-1}$
- Detector: Pilatus 1K
 - Resolution in time: $\sim 1.1\text{-}2.3 \text{ s}$

■ Sputter conditions [1]:

- Target: VC_{1-x}
- Substrate: Si(100) with natural oxide
- Target-substrate Distance: 10 cm
- Argon Pressure: $2 \times 10^{-3} \text{ mbar}$
- Deposition rate 0.22 nm/s@ 200 W

[1] Krause, Kaufholz et al., J. Synchrotron Rad. (2012), **19**, 216-222

5

29.03.2012

DPG- Frühjahrstagung Berlin 2012, M. Kaufholz

In situ X-Ray Reflectivity measurements during Sputtering of Vanadium Carbide thin films

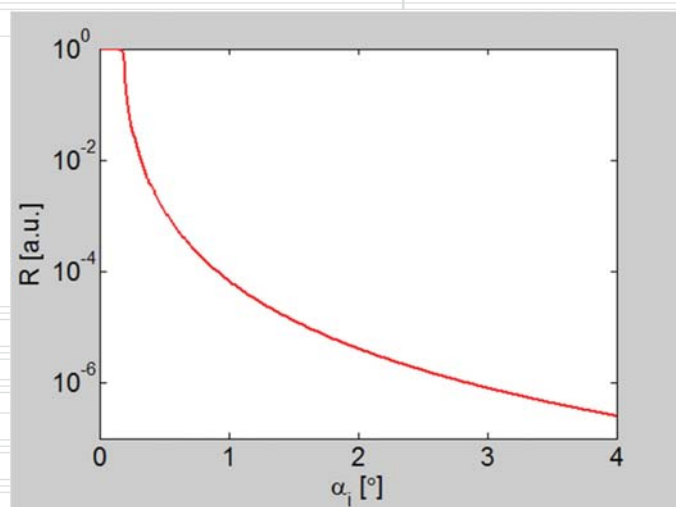
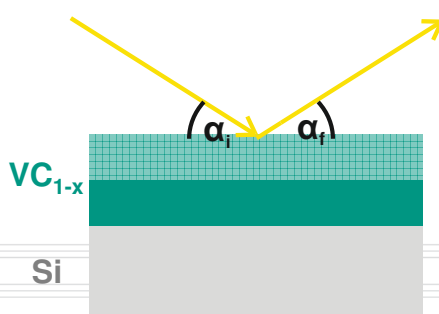


Institut für
Synchrotronstrahlung (ISS)

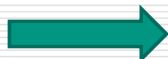
In situ X-Ray Reflectivity: “full angular range”



■ Measure full angular range



- High deposition rate of 0.22 nm @ 200 W \rightarrow $\sim 90\text{nm}$ deposition/XRR
- Possible electron density and roughness changes



Interpretation of XRR curve difficult

6

29.03.2012

DPG- Frühjahrstagung Berlin 2012, M. Kaufholz

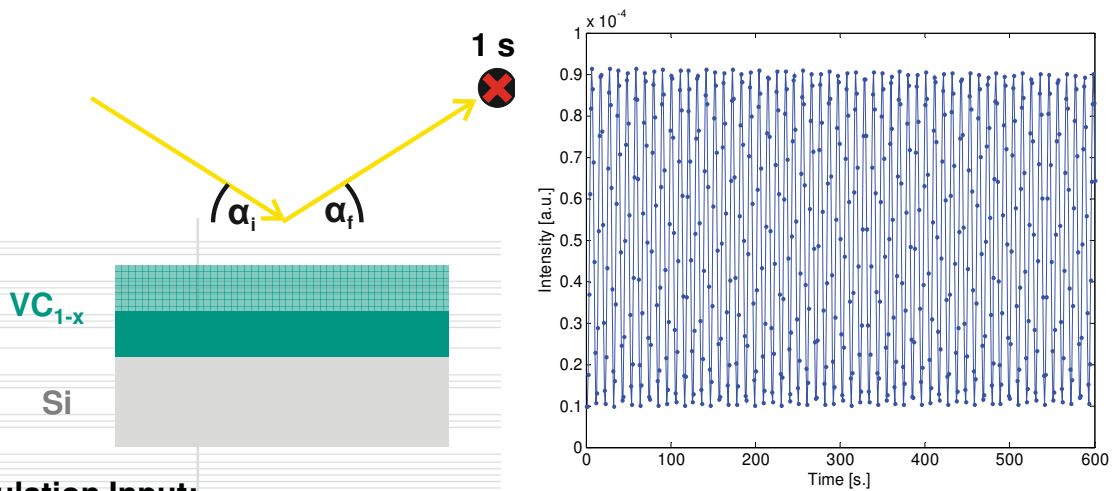
In situ X-Ray Reflectivity measurements during Sputtering of Vanadium Carbide thin films



Institut für
Synchrotronstrahlung (ISS)

In situ X-Ray Reflectivity: "fixed angular position"

- Detector and sample are at a **fixed angular position**
- Measuring Pre- and Post-growth full angular range XRR



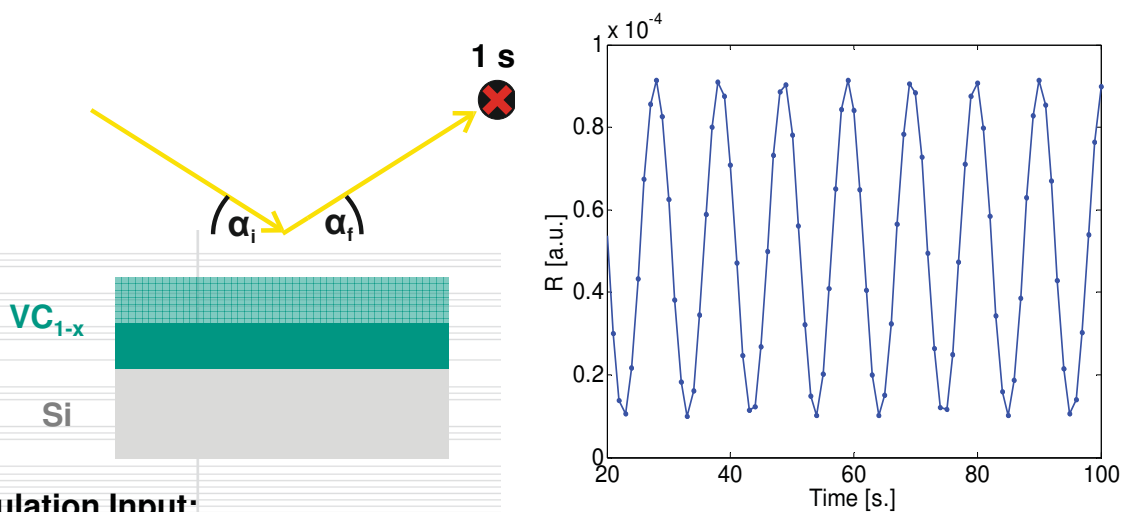
Simulation Input:

DC Power: 200 W \rightarrow Deposition Rate: 0.217 nm/s

$$\alpha_i = 1.6^\circ$$

In situ X-Ray Reflectivity: "fixed angular position"

- Detector and sample are at a **fixed angular position**
- Measuring Pre- and Post-growth full angular range XRR



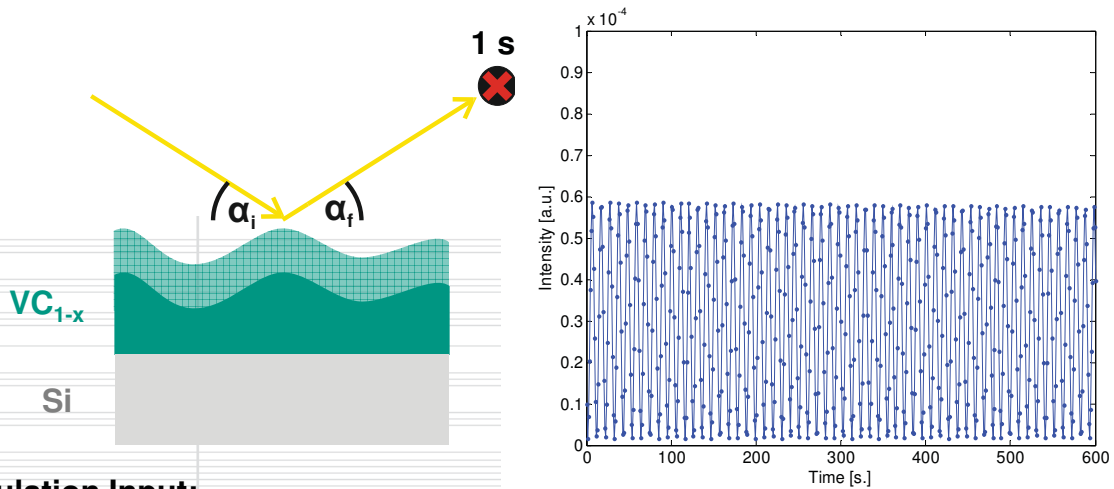
Simulation Input:

DC Power: 200 W \rightarrow Deposition Rate: 0.217 nm/s

$$\alpha_i = 1.6^\circ$$

In situ X-Ray Reflectivity: "fixed angular position"

- Detector and sample are at a **fixed angular position**
- Measuring Pre- and Post-growth full angular range XRR



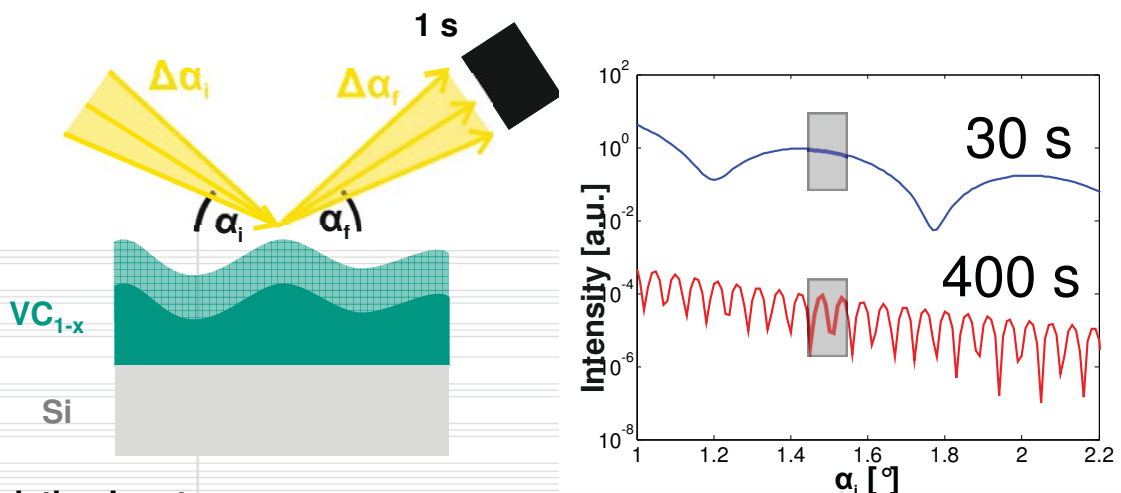
Simulation Input:

DC Power: 200 W → Deposition Rate: 0.217 nm/s

$$\alpha_i = 1.6^\circ$$

In situ X-Ray Reflectivity: "fixed angular position"

- Detector and sample are at a **fixed angular position**
- Measuring Pre- and Post-growth full angular range XRR



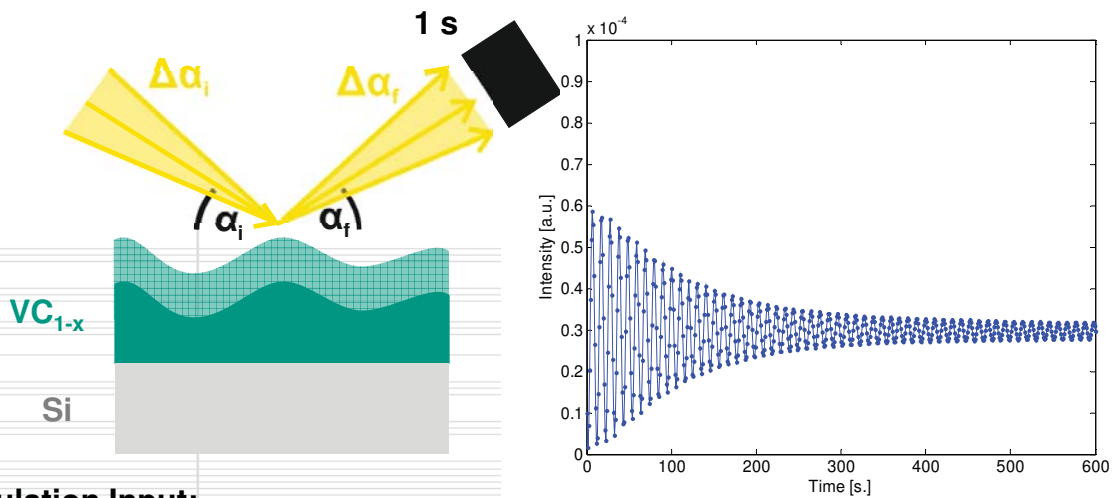
Simulation Input:

DC Power: 200 W → Deposition Rate: 0.217 nm/s

$$\alpha_i = 1.6^\circ$$

In situ X-Ray Reflectivity: "fixed angular position"

- Detector and sample are at a **fixed angular position**
- Measuring Pre- and Post-growth full angular range XRR



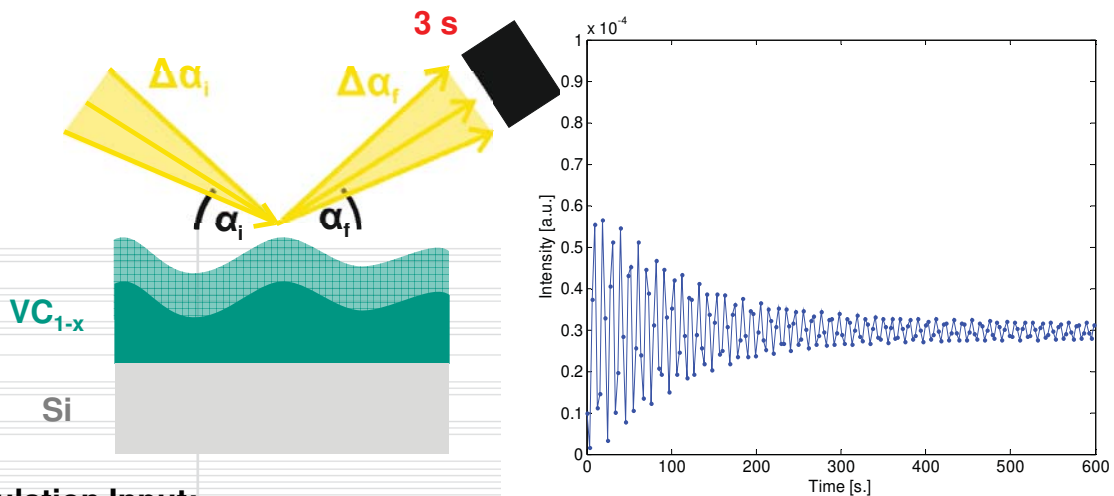
Simulation Input:

DC Power: 200 W → Deposition Rate: 0.217 nm/s

$$\alpha_i = 1.6^\circ$$

In situ X-Ray Reflectivity: "fixed angular position"

- Detector and sample are at a **fixed angular position**
- Measuring Pre- and Post-growth full angular range XRR



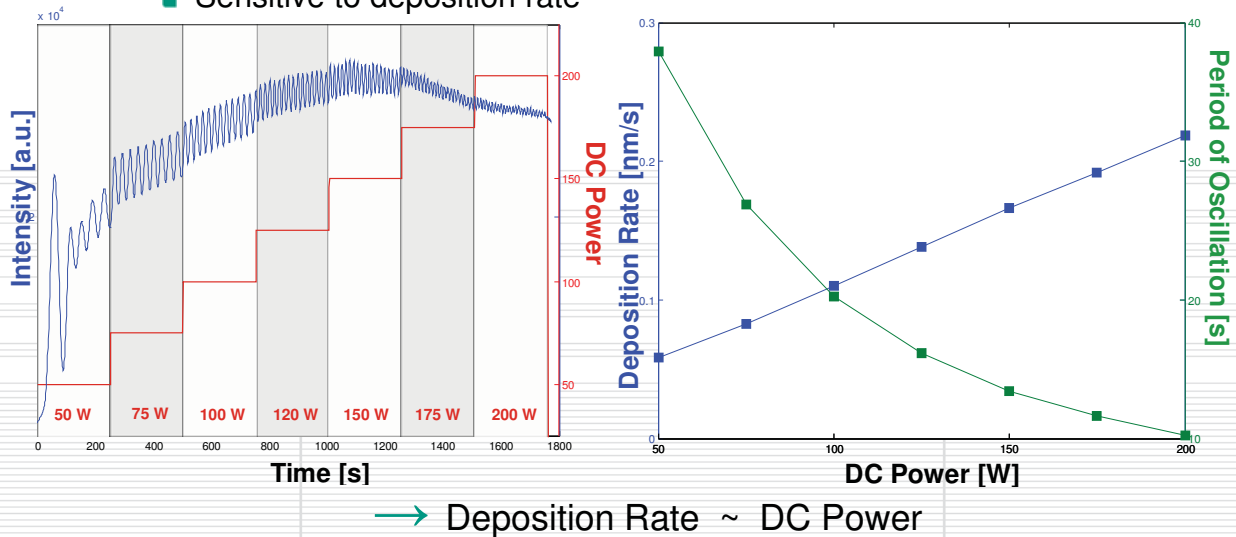
Simulation Input:

DC Power: 200 W → Deposition Rate: 0.217 nm/s

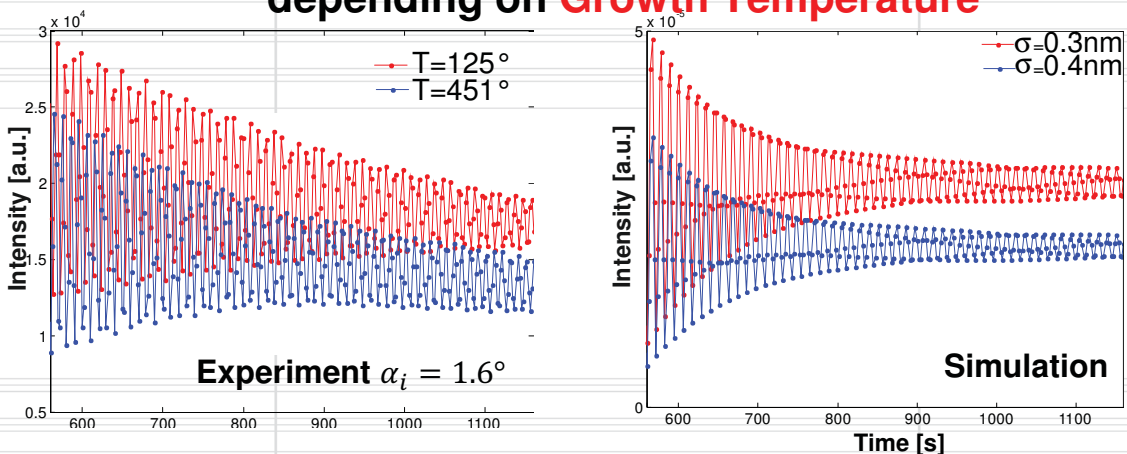
$$\alpha_i = 1.6^\circ$$

Example 1: Determination of Deposition Rate depending on DC Power at RT

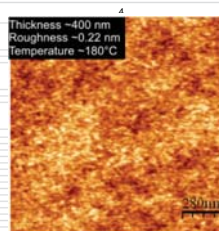
- Increase of DC Power by $\Delta P = 25\text{W}$ every 250s
- $\alpha_i = 1.6^\circ$:
 - Error due to changes in electron density <1%
 - Sensitive to deposition rate



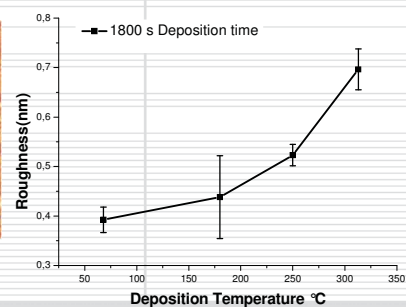
Example 2: Monitoring of Roughness depending on Growth Temperature



- Increase of Temperature leads to increase of Roughness
- Consistent with ex situ AFM

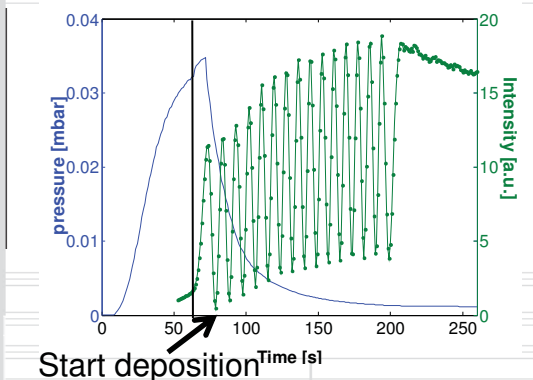


Ex situ: AFM



Example 3: Different Electron Densities due to Interruption of Deposition

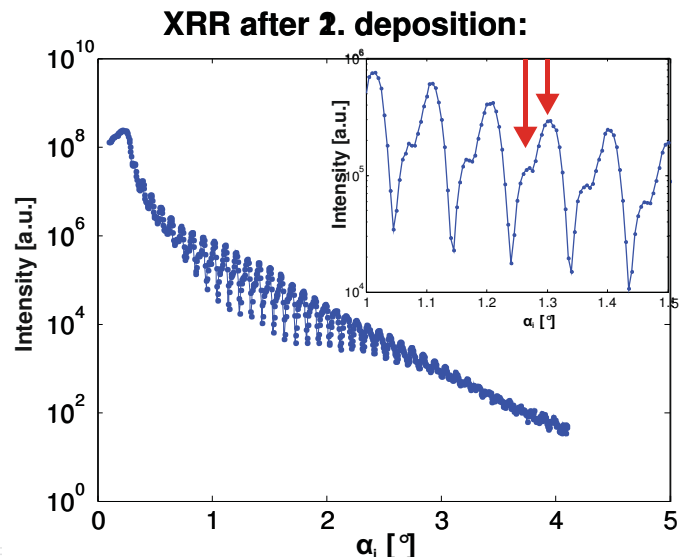
- Interruption of deposition after 200s @ RT and DC Power of 200 W



Start deposition



➔ **Multilayer of one material**



Summary

- *In situ* X-Ray Reflectivity is suitable for investigation of VC_{1-x}
 - Sensitive to
 - Deposition Rate
 - Roughness
 - Density
 - Sensitive to different sputtering conditions

Outlook

- Simulation of *in situ* XRR curves
 - Growth Model (Scaling law)
 - Include diffuse scattering
 - Limits of method
- Combining with other methods for a better understanding
 - *In situ* & *ex situ* X-Ray Diffraction and Absorption Spectroscopy
 - XPS, AFM, TEM, ... (in UHV conditions)
 - Measuring Hardness via Nano-/Microindentation

Acknowledgements

- M. Mantilla for technical support @ MPI Beamline @ ANKA
- H. Gräfe for technical support @ UHVLab @ ANKA
- S. Darma, J. Gemmler for fruitful discussion
- Financed in the framework of Excellence Initiative within the project KIT-Nanolab@ ANKA

Thank You for Your Attention !