

# Atomic Layer Deposition of Cobalt at Low Temperatures

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The conformal deposition of cobalt is still an ongoing topic of research. Applications are manifold and include interconnects<sup>1</sup>, seed layers for electroplating<sup>2</sup>, magnetic sensor systems<sup>3</sup>, and antibacterial coatings<sup>4</sup>. The thermal budget during film deposition is a crucial parameter which has to be taken into account. Here we present the development of a low temperature Atomic Layer Deposition (ALD) process for the formation of metallic cobalt thin films.

The professorship of Inorganic Chemistry at the Chemnitz University of Technology developed a set of cobalt precursors on the base of dicobalt-hexacarbonyl-alkynes as  $[\text{Co}_2(\text{CO})_6(\text{RC}\equiv\text{CR}')]^5$ . We evaluated a set of these precursors for chemical deposition methods using density functional theory (DFT) calculations. According to these calculations the precursor  $[\text{Co}_2(\text{CO})_6((\text{CH}_3)_3\text{SiC}\equiv\text{CSi}(\text{CH}_3)_3)]$  is the most favourable precursor for Chemical Vapour Deposition (CVD) as it thermally decomposes completely on the substrate surface. In contrast, precursor  $[\text{Co}_2(\text{CO})_6(\text{HC}\equiv\text{CC}_5\text{H}_{11})]$  was identified as the most favourable precursor for deposition via ALD as it adsorbs with adjoined  $(\text{HC}\equiv\text{CC}_5\text{H}_{11})$  group which can be easily removed by use of mobile hydrogen. A simplified scheme for the surface reaction and release of the hydrocarbon group is shown in Figure 1.

We developed an ALD process based on the DFT results by use of  $[\text{Co}_2(\text{CO})_6(\text{HC}\equiv\text{CC}_5\text{H}_{11})]$  as the cobalt precursor and hydrogen plasma.<sup>6</sup> The process development was done on a *scia Atol 200* reactor, which was designed and fabricated by *scia systems GmbH* in cooperation with *Fraunhofer ENAS* and the *Chemnitz University of Technology*. The processes took place on standardised 200 mm Si wafers with a preliminary  $\text{SiO}_2$  layer of 100 nm thickness. The precursor was evaporated via bubbling method. A full ALD cycle consists of cobalt precursor pulse, Ar purge,  $\text{H}_2$  plasma pulse, and a second Ar purge.

The deposited cobalt films were analysed by *in-vacuo* ellipsometry to determine in-line the film growth rates. Figure 2 shows the deposition rates in the temperature range from 35 °C to 125 °C showing the ALD window for this process within the range of 50 °C to 110 °C. *Ex-situ* measurements with spectroscopic ellipsometry were done to determine the thin film homogeneity on wafer level using a spiral measurement pattern with 5 mm edge exclusion. The optimised process had a film thickness deviation of about 1.5% relative standard deviation.

Additional measurements with X-ray photoelectron spectroscopy confirmed that the deposited films consist of cobalt in metallic state. Sole contaminations were identified as oxygen and carbon which are expected as the wafers were measured *ex situ* and may oxidise during transport.

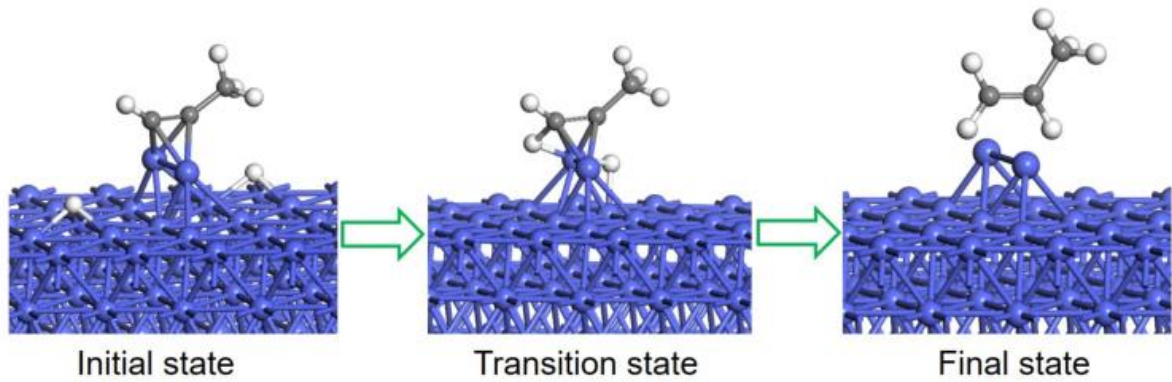
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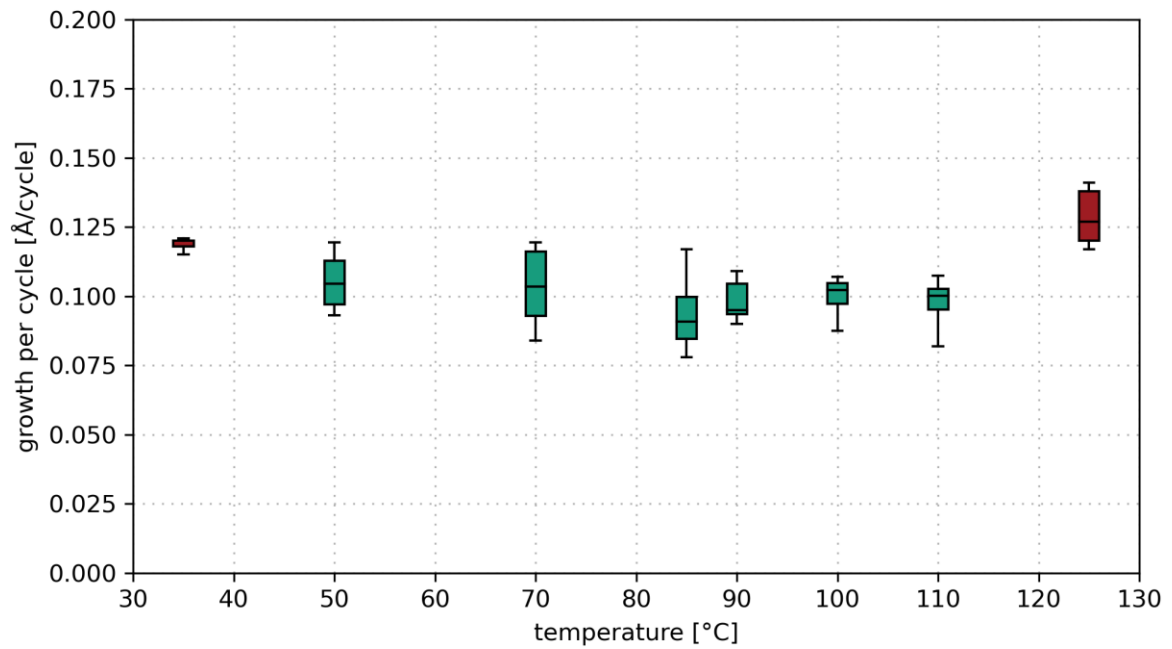
## References

1. Lanzillo, N. A., Chu, A., Bhosale, P. & Dechene, D. *IEEE Trans. VLSI Syst.* **30**, 60–67 (2022).
2. Liu, Y. *et al. J. Electrochem. Soc.* **169**, 082508 (2022).
3. Sharma, A. *et al. J. Mater. Chem. C* **8**, 11822–11829 (2020).
4. Jeong, Y.-M., Lee, J.-K., Ha, S.-C. & Kim, S. H. *Thin Solid Films* **517**, 2855–2858 (2009).
5. Georgi, C. *et al. J. Mater. Chem. C* **2**, 4676–4682 (2014).
6. Franz, M. *et al. Beilstein J. Nanotechnol.* **14**, 951–963 (2023).

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**Figure 1: density functional theory simulations showing the surface reactions between hydrocarbon ligand and hydrogen.**



**Figure 2: Temperature dependence of growth rate.**