

Grafting and Incorporation of Transition metals

to Unsaturated Silicon Clusters (Siliconoids)

N. Poitiers,^[a] L. Giarrana^[a] and <u>D. Scheschkewitz^[a]</u>

^[a] Saarland University, 66123 Saarbrücken, Germany, E-Mail: scheschkewitz@mx.uni-saarland.de

Unsaturated silicon clusters (siliconoids) are intermediates of the gas phase deposition of silicon, plausible constituents of amorphous silicon, and suitable model systems for bare surfaces of crystalline silicon.^[1] In combination with transition metals, silicon is central to an increasing number of heterogenous catalytic processes.^[2] Based on the ensuing hypothesis that silicon-rich ligands may also be advantageous in a homogenous setting, we became interested in the interaction of the by now well-established stable siliconoids with transitions metal centers. As direct grafting through salt metathesis only worked in case of Group 4 metallocene chlorides, we resorted to the attachment of pending auxiliary ligand moieties, namely tetrylenes of the Roesky-type.^[3]



Figure 1. Emblematic illustration of the coordination of iridium and rhodium to silylene-substitued siliconoids. [4]

Depending on the nature of the introduced transition metal center either exohedral coordination or partial incorporation into the cluster scaffold produced unique silicon-rich coordination environments.^[4] Some of the resulting complexes show promising catalytic activity in alkene isomerization (Figure 1).

References

- [1] H. Neergaard Waltenburg, J. T.Yates, J. T. Chem. Rev. 1995, 95, 1589-1673.
- [2] X. Chen, C. Liang, *Catal. Sci. Technol.* **2019**, *9*, 4785-4820.
- [3] N. E. Poitiers, N. E.; L. Giarrana, K. I. Leszczyńska, K. I.; V. Huch, M. Zimmer, D. Scheschkewitz, *Angew. Chem. Int. Ed.* **2020**, *59*, 8532-8536.
- [4] N. E. Poitiers, L. Giarrana, V. Huch, M. Zimmer, D. Scheschkewitz, Chem. Sci. 2020, 11, 7782-7788.