七座シッフ塩基配位子を有する 4 価セリウムエノラート錯体と Nーオキシルおよびフェノキシラジカルの反応

(阪大院基礎工¹・阪大院薬²)○千賀 大輔¹・真島 和志²・劍 隼人¹ Reactivity of Heptadentated Schiff-base Ligated Cerium(IV) Enolate Complexes with N-Oxyl and Phenoxy Radicals (¹Graduate School of Engineering Science, Osaka University, ²Graduate School of Pharmaceutical Sciences, Osaka University) ○Daisuke Senga,¹ Kazushi Mashima,² Hayato Tsurugi¹

Enolate ligands on redox-active metals serve as non-innocent redox-active ligands to form their valence tautomers, metal-coordinated α-carbonyl radicals. We recently found the synthesis of cerium(IV) enolate complexes 1-L having a trianionic heptadentate Schiff-base ligand L (LH₃ = tris[2-(2-salicylideneimino)ethyl]amine) from LCe(O'Bu) with alkylketones *via* deprotonation of the α-hydrogen atom of the carbonyl group, and 1,4-dicarbonyl compound 2 was obtained *via* homo-coupling of the α-carbonyl radicals under blue LED irradiation of 1-L¹. Herein, we report that addition of 2,2,6,6-tetramethylpiperidine-N-oxyl (TEMPO) to the cerium(IV) enolate complexes produced the C–O bond forming product 3 at the α-position of the carbonyl group along with trivalent LCe(THF)_n complexes: cerium(IV) enolate complex 1-L² having a sterically less-hindered ligand gave 3 in quantitative yield. Electronic structure of the cerium enolate complexes 1-L and their reactions with other stable radicals are disclosed. *Keywords: Cerium complex; Enolate complex; Schiff base ligand; N-Oxyl radical; Radical coupling*

1) Heras, C.; Gómez-Palomino, A.; Romea, P.; Urpí, F.; Bofill, J. M.; Moreira, I. de P. R. *J. Org. Chem.* **2017**, *82*, 8909.