## Development of Cobalt-Catalyzed Ammonia Oxidation under Ambient Reaction Conditions

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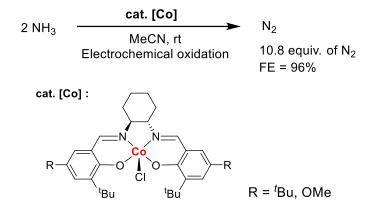
To realize the direct use of ammonia as an energy carrier, the development of catalytic ammonia oxidation reaction into dinitrogen is necessary. Homogeneous molecular catalysts for ammonia oxidation, especially with transition-metal complexes, are one of the attractive targets from viewpoints of fundamental mechanistic studies and the development of highly active and robust electrocatalysts (Scheme 1).<sup>1</sup>

## Scheme 1

$$2 \text{ NH}_3 \xrightarrow{\text{cat.}} \text{N}_2 + 6 \text{ e}^- + 6 \text{ H}^+$$

Previously, our group reported ammonia oxidation using manganese-salen complexes as catalysts.<sup>2</sup> As an extensive study, we have investigated the catalytic activity of a series of cobalt-complexes bearing various salen ligands toward ammonia oxidation under ambient reaction conditions. As a result, we have found that these cobalt salen complexes worked as efficient catalysts for ammonia oxidation under electrochemical conditions with lower overpotential ( $\eta = 1.26$  V) than the manganese complexes ( $\eta = 1.6$  V) according to the CV measurements. Bulk electrolysis of ammonia solution containing the cobalt complexes as catalysts provided 10.8 equiv. of N<sub>2</sub> based on the cobalt atom with high faradaic efficiency (FE) of 96% (Scheme 2).

## Scheme 2



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