

## In-depth investigation of the synthesis mechanism of novel NO oxidation catalyst $\text{Co}/\text{K}_x\text{Ti}_2\text{O}_5$

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$\text{NO}_x$  abatement is one of the key problems for lean-burn engines, and there is increasing acceptance of the view that the oxidation of NO to  $\text{NO}_2$  is an important pre-requisite step for treating emission gases.

In the present work,  $\text{Co}/\text{K}_x\text{Ti}_2\text{O}_5$  catalyst with well-dispersed nano- $\text{Co}_3\text{O}_4$  particles, synthesized by an ion exchange method from  $\text{K}_2\text{Ti}_2\text{O}_5$  precursor, showed extremely high NO oxidation activity and good stability. Morphological changes and microcosmic processes during the course of catalyst preparation were examined using AAS, XRD and SEM-EDX. During the ion exchange of  $\text{K}_2\text{Ti}_2\text{O}_5$  with Co precursor solution,  $\text{K}^+$  ions were replaced by  $\text{H}^+$  ( $\text{H}_3\text{O}^+$ ). After calcination at  $500^\circ\text{C}$ , a partially collapsed structure was obtained when the exchange was controlled to a certain degree. Fortunately, the Co that precipitated on this collapsed structure formed highly stable nano-particles of  $\text{Co}_3\text{O}_4$ . Catalytic activity for the NO oxidation was found to be highly dependent on the loading and particle size of Co and K remaining in the  $\text{K}_x\text{Ti}_2\text{O}_5$  support.