**Introduction**

The Fe-ZSM-5 catalysts have been widely investigated for such significant reactions as the selective catalytic reduction (SCR) reaction of NO, one-step oxidation of benzene to phenol. Recently, it was found that Fe-ZSM-5 exhibits high photocatalytic reactivity for the decomposition of N₂O into N₂ and O₂. The present study deals with the effect of the different Si/Al ratio on the local structures of the Fe-oxides within Fe-ZSM-5 by means of XAFS (EXAFS and XANES) spectroscopy.

**Experimental**

The Fe-ZSM-5 samples (1.0 Fe wt%) were prepared by the solid-state reaction of H-ZSM-5 having different Si/Al ratios (20, 34, 950) supplied by Tosoh Co with FeCl₃-6H₂O. The samples were calcined at 823 K for 9 h in air. XAFS (XANES and EXAFS) spectra were obtained in the fluorescence mode at 295 K at the BL-9A facility of the High Energy Acceleration Research Organization (KEK) in Tsukuba. The EXAFS data were examined by the analysis program Rigaku EXAFS (REX).

**Results and Discussion**

Figure 1 shows the Fe K-edge XANES and EXAFS spectra of Fe-ZSM-5 (20, 34, 950). The characteristic feature of the XANES spectra of Fe-ZSM-5 is the appearance of a pre-edge peak assigned to a 1s-to-3d-like transition, which is forbidden in pure octahedral symmetry. It was found that the shape of the XANES spectrum of Fe-ZSM-5 (Si/Al=20) is quite similar to that of the Fe₂O₃ compound, indicating that this sample consists of Fe-oxides with an octahedral coordination. The pre-edge peak intensity is the highest in Fe-ZSM-5 (Si/Al=950), however, it decreases slightly and the shape of the XANES spectra changes when the Si/Al ratio is decreased. From the EXAFS radial structure function of Fe-ZSM-5 (Si/Al=950) in the 3-10.5 Å⁻¹ region, only one peak due to the presence of neighboring oxygen atoms (Fe-O) was observed at ca. 1.0-2.0 Å (without phase-shift correction), while an additional peak due to the Fe-O-Fe bond was observed at ca. 3.0 Å with the averaged Fe-oxide clusters between two and three Fe ions in Fe-ZSM-5 (Si/Al=20).

![Fig. 1](image-url)