## Brillouin and Raman scattering studies of the isotopically induced ferroelectric phase transition of SrTi<sup>18</sup>O<sub>3</sub>

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The quantum paraelectric SrTiO<sub>3</sub> has attracted a lot of interest of many researchers since the quantum-mechanical stabilization of the paraelectric phase below 4K was pointed out as a quantum effect on the polarization fluctuation.<sup>1)</sup> Recently a ferroelectric phase transition has been reported in the SrTiO<sub>3</sub> crystal exchanged of isotope <sup>18</sup>O for <sup>16</sup>O.<sup>2)</sup> The difference of mass between <sup>16</sup>O and <sup>18</sup>O seems to suppress the quantum fluctuations and to bring divergence of the spatial correlation of the fluctuating polarization at the paraelectric-ferroelectric phase transition. In the present study, the dynamical mechanism of the ferroelectric phase transition of the isotopically exchanged SrTiO<sub>3</sub> has been studied by both of the Brillouin- and Raman scatterings.<sup>3-5)</sup> The exchange rate of the isotope <sup>18</sup>O is 87% which gives the ferroelectric transition temperature Tc=24K.<sup>2)</sup>

Brillouin scattering study found anomalous temperature dependence of the acoustic mode near Tc; a transverse mode shows a softening behavior in contrast to no anomaly of the longitudinal c<sub>33</sub> mode. The rather complicated behavior of the acoustic modes near Tc has been elucidated. A strong central component appeared in the Brillouin scattering spectra suggests existence of the dynamical cluster near Tc. The isotope effect on the doublet problem is discussed.

Raman scattering study elucidates a broad spectrum characteristic for the ferroelectric microregion (FMR) reported previously for  $SrTi^{16}O_3$ .<sup>5,6)</sup> The size of FMR increases anomalously when temperature approaches to Tc in the paraelectric phase. The soft mode spectra are not clear below Tc, though every soft-ferroelectric mode should be Raman active in the ferroelectric phase.

The ferroelectric phase transition induced isotopically in SrTiO<sub>3</sub> seems to be caused by somewhat complicated cluster dynamics rather than by homogeneous fluctuation of polarization.

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