Composition Analysis of a Thin Gate Oxide Layer using Model-Based EELS Quantification

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The reductions of device dimensions in MOSFET technologies demand the introduction of thin gate oxides with thicknesses of a few nanometers. The electrical properties of such layers vary strongly with the structure and composition near the $\mathrm{SiO}_x/\mathrm{Si}$ interface. It is therefore of critical importance to be able to quantify the gate oxide composition with a high spatial resolution. Electron energy-loss spectroscopy (EELS) is often used for this purpose. However, the interaction of incident electrons with both SiO_x and Si makes conventional EELS quantification [1] impossible near the $\mathrm{SiO}_x/\mathrm{Si}$ interface. To separate this overlapping effect, model-based quantification of EEL spectra has to be employed [2, 3]. The results obtained on a thin gate oxide layer making use of this technique are presented in this paper.

The TEM sample was prepared by focused ion beam (FIB) milling, with the final thickness of the specimen around 100 nm. A cross-sectional high resolution TEM image of the gate oxide is shown in Fig. 1a. EELS spectra with an energy resolution of 1.5 eV were collected simultaneously with a high-resolution Z-contrast STEM (scanning transmission electron microscopy) image (Fig.1b). The estimated probe size was 0.2 nm and the convergence and collection half-angle were respectively 18 mrad and 47 mrad. The conventional composition analysis uses the core-loss edges of O and Si with the concentration ratio proportional to the edge integral ratio. This method only works for locations certain distances away from the SiO_x/Si interface where no edge overlapping occurs. As shown in Fig.2c, near the SiO_x/Si interface, both Si and SiO_x contribute to the Si signal which is essentially an overlap between two distinct Si-L_{2,3} edges (Figs. 2a and 2b) with an energy shift of about 5eV. Accurate composition analysis requires determination of the Si signal contributed by the gate oxide only. This can be achieved using the so-called model-based quantification method [2, 3]. Si and O were analyzed independently using two different models. In Si analysis, the model includes two Si-L₂₃ edges from Si and SiO_x respectively and their corresponding fine structures, which are determined experimentally from locations well away from the SiO_x/Si interface. Multiple scattering effect is taken into consideration through convolution of the single-scattering core-loss spectrum with a low-loss spectrum. The cross-section ratio of O-K/Si-L_{2,3} was calibrated using a standard SiO₂ sample. In Fig.3, the composition profile obtained using the model-based method is compared with that obtained from the conventional method where the total Si signal is used. Significant differences were found between the two methods near the SiO_x/Si interface. It should be noted that the accuracy of the model-based quantification is governed by the statistical quality of the spectrum and the accuracy of the cross-section ratio. In this work, the accuracy is estimated to be about 5%. It is suggested that the model-based method should be used for EELS quantification in case that the ionization edges of interest are overlapped and the conventional method fails to provide accurate results.

References

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- [2] J. Verbeeck, S. Van Avert, Ultramicroscopy 101(2004) 207.

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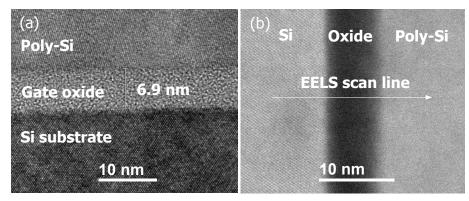


FIG.1. (a) Cross-section high-resolution TEM image of the gate oxide, (b) High-resolution STEM image of the gate oxide and the position of the EELS line scan.

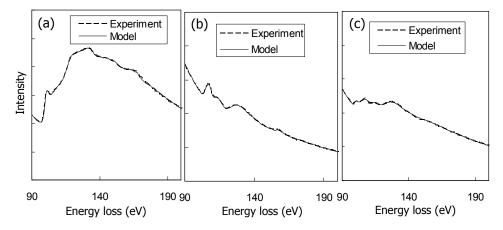


FIG. 2. Typical EELS spectrum of Si- L_{23} edge (dashed line) obtained in Si (a), SiO_x (b) and near the SiO_x/Si interface (c). The simulated spectrum (solid line) is also shown for comparison.

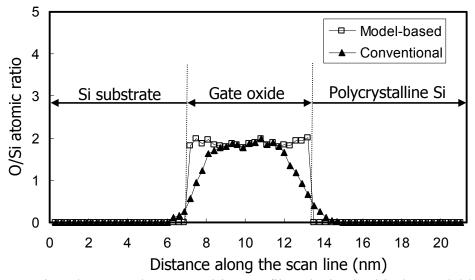


FIG.3. Comparison between the composition profiles obtained with the model-based method and the conventional method. In the latter case, the total Si signal from both Si and SiO_x is involved in O/Si ratio determination near the SiO_x/Si interface.