INTRODUCTION

Barium titanate (BaTiO₃; BTO) is a ferroelectric material, which has been used as multilayer ceramic capacitors for long period (Tsurumi, 2007), and is a promising candidate for ferroelectric random access memory (Varghese et al., 2013). Because there has been a strong demand to downsize these devices in recent years, ferroelectric materials of smaller scale such as nanoparticles (NPs), has been intensively studied (Arlt et al., 1985; Frey & Payne, 1996; Imanaka et al., 2013; Mimura & Kato, 2014; Tsurumi et al., 2006; Varghese et al., 2013; Zhao et al., 2004). Although it is well known that electrical property of nanoparticles depends on the atomistic structure. Although surface may possibly have an impact on the atomistic structure, reconstructed structure at the surface has not been widely investigated. In the present study, Ba-ions position near surface in a BaTiO₃ nanoparticle has been quantitatively characterized by scanning transmission electron microscopy. It was found that some Ba ions at the surface were greatly displaced in non-uniform directions.

Key Words: Nanoparticle, BaTiO₃, Scanning transmission electron microscopy, Surface, Ferroelectrics
of the particle size, which might be contradictory to the conventional picture that Curie temperature decreases as the particle size decreases (Hoshina et al., 2008; Huan et al., 2014; Imanaka et al., 2013; Smith et al., 2008; Tsurumi et al., 2006; Varghese et al., 2013; Zhao et al., 2004). On the other hand, Hoshina et al. (2008) reported that crystal structure varies within a particle. There is three-layer structure composed of surface (paraelectric), intermediate, and inner (ferroelectric) regions for bigger particle, while the whole particle becomes paraelectric for smaller particle. In analogy to the dielectric/ferroelectric properties, these examples showed that clear understanding between the particle size and the crystal structure has not been obtained. This also implies that local structural characterization is required in addition to averaged crystal structure characterization.

For the sake of local structural characterization, electron microscopy such as high-resolution transmission electron microscopy or scanning TEM (STEM) is a powerful method, since atomic arrangement can directly be observed. In particular, precision in the atomic position determination has been improved to picometer level in recent years (Borisevich et al., 2012; Jia et al., 2008; Polking et al., 2012; Yadav et al., 2016; Yankovich et al., 2014). Such structural characterization has been reported for example by Polking et al. (2012) with high-resolution TEM and electron holography and by Li et al. (2014) with high-resolution TEM under in-situ heating. Polking et al. (2012) has visualized the polarization distribution within NP and reported that BTO NP with the size below 10 nm exhibited ferroelectricity at room temperature. On the other hand, Li et al. (2014) has reported that BTO NPs with the sizes ranging from 2.5 nm to 10 nm composed of multiple phases and the increment of Curie temperature up to 600°C. Even though these studies have demonstrated structural distribution in NPs, structure of NP with the size larger than 20 nm may be needed since such larger NP was not investigated in the previous studies (Li et al., 2014; Polking et al., 2012). Also, both of the studies have suggested that ferroelectric order or crystal phase is affected by the surface. However, the detailed reconstructed structure at the surface has not been clarified. Therefore, in the present study, we report a structural investigation near surface in a BTO NP studied by STEM observation. In particular, displacement of Ba ion near surface was quantitatively studied by STEM observation. In particular, precision in the atomic position determination has been improved to picometer level in recent years (Borisevich et al., 2012; Jia et al., 2008; Polking et al., 2012; Yadav et al., 2016; Yankovich et al., 2014). Such structural characterization has been reported for example by Polking et al. (2012) and high-resolution TEM and electron holography and by Li et al. (2014) with high-resolution TEM under in-situ heating. Polking et al. (2012) has visualized the polarization distribution within NP and reported that BTO NP with the size below 10 nm exhibited ferroelectricity at room temperature. On the other hand, Li et al. (2014) has reported that BTO NPs with the sizes ranging from 2.5 nm to 10 nm composed of multiple phases and the increment of Curie temperature up to 600°C. Even though these studies have demonstrated structural distribution in NPs, structure of NP with the size larger than 20 nm may be needed since such larger NP was not investigated in the previous studies (Li et al., 2014; Polking et al., 2012). Also, both of the studies have suggested that ferroelectric order or crystal phase is affected by the surface. However, the detailed reconstructed structure at the surface has not been clarified. Therefore, in the present study, we report a structural investigation near surface in a BTO NP studied by STEM observation. In particular, displacement of Ba ion near surface was quantitatively analyzed. Commercially available BTO NP with nominal size of 50 nm was studied as a specimen.

MATERIALS AND METHODS

Commercially available BTO NPs (model no. 745952; Sigma-Aldrich, USA; purity: 99.9%, nominal particle size: 50 nm) were investigated in the present study. For comparison, commercially available SrTiO₃ single crystals (Shinkosha, Japan) were also studied. BTO NP powder was homogeneously dispersed in ethanol using ultrasonic cleaner. Subsequently, a small amount of the solution was dropped onto a commercially available TEM grid (NS-C15; Okenshoji, Japan) and the grid was dried in air. On the other hand, a STEM thin foil of the SrTiO₃ single crystal was prepared using an FB-2100 focused-ion-beam system (Hitachi High-Technologies, Japan) with the beam energy of 40 keV. Final milling was conducted to remove the damaged layer from the surface by argon-ion-beam milling system (PIPS; Gatan, USA).

STEM observation was carried out using a JEM-ARM200F (JEOL, Japan) equipped with a spherical aberration corrector for the electron probe (Rose, 1994). The observation was carried out under an acceleration voltage of 200 kV. TEM images were acquired with an Orius 200D charge coupled device camera (Gatan). Annular dark-field (ADF) STEM images containing 512×512 pixels were recorded with a detection angle range of 90 and 370 mrad and with a pixel dwell time of 2 μs. Approximately twenty images were acquired from a single region of interest, and the image series was processed using a non-rigid registration method (Jones et al., 2015) (SmartAlign; HREM Research, Japan) to improve the signal-to-noise ratio. This procedure is useful to reduce the influence of image distortion (Jones et al., 2015). The positions of Ba and Ti ions in the image were identified by the peak fitting method. The peak fitting was carried out using the two-dimensional Gaussian function in equation (1), as was done in a previous study (Yankovich et al., 2014),

$$Z = Z_0 + A \exp \left[ \frac{-1}{2 (w_x^2 + w_y^2)} \left( \frac{x - x_c}{w_x} \right)^2 + \left( \frac{y - y_c}{w_y} \right)^2 + \left( \frac{2 \sigma (x - x_c) (y - y_c)}{w_x w_y} \right) \right]$$

where $Z$ and $Z_0$ are the total and background intensity, respectively, $A$ is the peak intensity, $x_c$ and $y_c$ are the peak positions in the $x$ and $y$ directions, respectively, $w_x$ and $w_y$ are the width of the Gaussian function in the $x$ and $y$ directions, respectively, and $c$ is a fitting parameter. Differences between the intensities of the experimental image and the Gaussian function ($Z$) were minimized by the least-squares method by optimizing the parameters of $Z, Z_0, A, x_c, y_c, w_x, w_y$, and $c$. The optimization was performed by the Marquart method (Marquardt, 1963) using a script coded using Visual Basic 6 (Microsoft, USA). It should be noticed here that analysis of atomic-column position was conducted both for Ba and Ti ions. However, it was found that the analysis became less accurate in the case of Ti columns near surface for some images, which might be partly due to lower image contrast of Ti column near the surface. Therefore, for clear justification of the conclusion, the present study focuses on the analysis of Ba column positions.
RESULTS AND DISCUSSION

In prior to the observation of BTO NP, STEM observation of SrTiO3 single crystals was carried out to calibrate the non-squareness of the scan pixel (Yankovich et al., 2014). Fig. 1A shows the ADF STEM image used for the evaluation. Because the contrast of the ADF STEM image is dependent on the atomic number of constituent ions in the atomic column (Pennycook & Jesson, 1990), the brightest spot and second brightest spot indicate Sr and Ti ions, respectively, whereas the O ion is not visible. Interatomic spacing between neighboring Sr ions along the [100] or [010] directions, \( d_{Sr[100]} \) and \( d_{Sr[010]} \) (Fig. 1B) was investigated using peak fitting results, which is similar to the process used by Bals et al. (2006). Because it was found that the left-hand side of the image was considerably distorted, these regions were intentionally excluded from the analysis hereafter. As SrTiO3 has a cubic crystal structure with a lattice constant \( a \) of 390.5 pm (Mitchell et al., 2000) the ratio between \( d_{Sr[100]} \) and \( d_{Sr[010]} \) (\( d_{Sr[100]}/d_{Sr[010]} \)) should ideally be 1.000. On the other hand, it was found in our measurement that the average \( d_{Sr[100]} \) and \( d_{Sr[010]} \) was 17.45 pixels and 17.19 pixels long, resulting in \( d_{Sr[100]}/d_{Sr[010]} \) of approximately 1.015. Hereafter, the results were calibrated with this ratio. As a result of calibration, the average \( d_{Sr[100]} \) was determined as 390.5 pm with the standard deviation (SD) of 5.8 pm and the average \( d_{Sr[010]} \) was determined as 390.5 pm with the SD of 5.2 pm. Fig. 2A shows a representative TEM image of the BTO NP studied. The particle size was approximately 50 nm, having \{100\} and \{110\} habit planes on the edge and rounded corners at the atomic scale. It is considered from the image contrast that the particle was composed of multiple domains (Fig. 2B). Presence of multiple domains was also supported by the result that the diffraction spot measured from this particle (Fig. 2C) exhibited splitting (Fig. 2D). A typical ADF STEM image is shown in Fig. 2E, which was taken from the region (I) in Fig. 2B. Image contrast is higher in the grain interior (left top side) and lower near and at the surface (right bottom side), which suggests that the particle is thicker in the grain interior and thinner near and at the surface. For simplicity, the thicker grain-interior region, several unit cells from the surface, and
the outermost layer of the surface will be called the “grain interior”, “near surface”, and “outermost surface” hereafter, as schematically shown in Fig. 2F.

ADF STEM images (Fig. 2E and Fig. 3) taken from different regions (I-V) as shown in Fig. 2B were analyzed to measure the Ba ion position quantitatively. In particular, Ba ion displacement from the regular lattice position was determined for the near surface and the outermost surface regions (Fig. 3), wherein the reference regular lattice in the near surface and the outermost surface regions was obtained by extending the reference lattice in the grain interior region toward the surface. It was found that some Ba ions were displaced by larger than 20 pm near the surface, although the trends were different for different regions. Ba ions were displaced toward the right-bottom side (outward) in the top half, whereas the displacement in the bottom half was smaller in the region (I) (Fig. 4A). Ba ions were mostly displaced rightward in the outermost surface region in the region (II) (Fig. 4B), while the displacement was smaller and irregular in the region (III) (Fig. 4C). On the other hand, Ba ions in the outermost surface region were mostly displaced rightward (inward) in the region (IV) (Fig. 4D), which was a different trend from the other cases. Finally, in the region (V) (Fig. 4E), Ba ions were displaced upward (outward) in the right half of the outermost surface region, while some ions in the left half of the outermost surface region were displaced leftward nearly along the surface. Thus, some Ba ions were greatly displaced in non-uniform directions. In particular, outward displacement as observed for many Ba ions in Fig. 4A and B possibly leads to expansion of unit cell near surface, and the influence of the surface would be greater for smaller particle. This may explain by a trend reported by Smith et al. (2008) that unit cell volume increased as the particle size decreased, which was obtained from Rietveld and pair distribution function analysis for the X-ray diffraction measurement. It was also suggested that coherence between the neighboring unit cells was reduced for the smaller NP. Such a reduction of coherence may be induced by Ba-ion displacement toward non-uniform direction (Smith et al., 2008).

In summary, the atomic-scale STEM observation was carried out for near-surface region in a BTO NP with nominal size of 50 nm. Ba-ion position observed in the STEM images was

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**Fig. 3.** (A-D) Atomic scale annular dark-field scanning transmission electron microscopy image taken from the region (II)–(V).
quantitatively analyzed. It was found that some Ba ions at the outermost surface were greatly displaced in non-uniform directions.

**CONFLICT OF INTEREST**

No potential conflict of interest relevant to this article was reported.

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