INTRODUCTION

As Li-ion batteries have been popular as the next-generation energy source, various promising electrode materials have been investigated to improve battery performance. Especially, silicon appears as one of the most prominent anode materials for Li-ion batteries due to higher theoretical capacity and lower working potential. However, silicon experiences a huge volume expansion during lithiation and delithiation process, which causes rapid capacity degradation and very poor cycling performance (Chan et al., 2008; Obrovac & Christensen, 2004; Park et al., 2010; Winter et al., 1998). In order to solve these problems, material researches like nanostructures and amorphous Si and Si-alloy materials combined with matrix and have been investigated (Jung et al., 2011; Lee & Lee, 2002; Mao et al., 1999). In order to understand the reaction mechanism of Li and silicon crystal structures, obvious and direct observation method is very important. In this paper, we investigated the change of microstructure of Si alloy anode materials composed of active Si and inactive TiFeSi2 matrix after Li charging by using transmission electron microscope (TEM).

A successful transmission electron microscope (TEM) analysis is closely related to the preparation of the TEM specimen and should be followed by the suitable TEM specimen preparation depending on the purpose of analysis and the subject materials. In the case of the Si-based anode material, lithium atoms of formed Li silicide were removed due to ion beam and electron beam during TEM specimen preparation and TEM observation. To overcome the problem, we proposed a new technique to make a TEM specimen without the ion beam damage. In this study, two types of test specimens from the Si-based anode material of Li-ion battery were prepared by respectively adopting the only focused ion beam (FIB) method and the new FIB-ultramicrotome method. TEM analyses of two samples were conducted to compare the Ga ion damage of the test specimen.

**Key Words:** Li ion battery, Si-based anode material, Focused ion beam, Ultramicorome, Transmission electron microscopy
TEM observation. However, in the case of the Si-based anode material, lithium atoms of formed Li silicide were removed due to ion beam and electron beam during TEM specimen preparation and TEM observation (Danet et al., 2010). To overcome the problem, TEM specimens need to be prepared using mechanical techniques without ion beam damage. Although the specimen made by FIB has damages induced by Ga ions, the one made by ultramicrotome has artifacts such as plastic deformation and tears of TEM specimen due to mechanical force. Since the thickness of the TEM specimen depends on the size, material and shape of the material (Muto et al., 2009; Sasaki et al., 2009), we prepared a cube specimen of about 10 µm × 10 µm × 10 µm in a specific region using FIB and then prepared a TEM specimen preparation technique using ultramicrotome to obtain lamella. This technique (FIB-ultramicrotome technique) is expected to possess the advantages of both the FIB, which is favorable for sampling a specific area, and the ultramicrotome, which enables the preparation of thin specimens without Ga ion. In this study, two types of test specimens from the Si-based anode material of Li-ion battery was prepared by respectively adopting the FIB method only and the new FIB-ultramicrotome method. TEM analyses of two samples were conducted to compare the Ga ion damage of the test specimen.

MATERIALS AND METHODS

Battery Manufacturing
The manufacturing process of Si-based anode materials and battery, and the charge-discharge cycle testing method are the same as previous study (Chae et al., 2013). After 20 cycle testing, the battery was dismantled inside a glove box filled with inert Ar gas.

The lithium salts remaining on the surface was removed by using an organic solvent, dimethyl carbonate. The remaining materials were dried for approximately a day.

TEM Specimen Preparation
The prepared material was cracked and broken into shards through the volume expansion after undergoing the process as shown in Fig. 1. For TEM analysis, two types of specimen were prepared by using two kinds of preparation methods. The first method is a conventional TEM sampling method using FIB. Firstly, the FIB was used to select a specific area of interest from the shards of the charged materials. The selected area was thinned by FIB milling. In general, the thinner the thickness of the specimen, the better. But in this study, when the thickness of the specimen was too thin (<70 nm), the material properties were changed by ion-beam damage. The optimum thickness of our cross-sectional TEM specimen was 70 to 100 nm. The second method is a new method using FIB and ultramicrotome, which is firstly proposed in this study. Initially, FIB was used to extract a cube sample of about 10 µm × 10 µm × 10 µm from a specific area of interest. The extracted sample was attached to the top of the “B” post of lift-out grid for FIB in Fig. 2A. The grid was fixed to a support stick with glue, then placed in a plastic cap tube and filled with epoxy resin to make a block, as shown in Fig. 2A. The epoxy resin was prepared at a ratio of 4:1 resin to hardener. With a block being made, cautions should be taken to prevent bubble formation. The unintentionally formed microbubbles were removed through a vacuum defoamer followed by drying in a vacuum oven at 40°C for 24 hours. The block was then cut with a razor blade, in order to fit into a fixing holder. The block was smoothly trimmed into the size of about 50 µm × 50 µm and located at the centered on the grid “B” post, to which the test piece was attached using a trimming blade (Fig. 2B). During the cutting in using the ultramicrotome, cutting rate was set as 0.8 mm/s and as the thickness 70 nm. A diamond blade (3 mm, 35°) was used to cut the sample trimmed in the prior step. Water, anhydrous ethanol and organic solvent were used as the solvents to float the severed specimen. In the case of anhydrous ethanol, the specimen was sank due to low specific gravity, and the organic solvent were solidified along the diamond blade and block in the boat. These caused difficulties in cutting and extracting the specimen. Therefore, water was used in this experiment. In order to minimize the reaction with Li, the severed specimens were immediately

Fig. 1. Schematic illustration showing volume change process of Si-TiFeSi₂ active materials.
dried and used for TEM analysis.

RESULTS AND DISCUSSION

The material used in this study is a Si-based anode material comprising active material Si and TiFeSi$_2$ matrix. The material was manufactured by melt-spinning method and composed of fine and coarse grain regions (Chae et al., 2013; Raghavan, 2009; Weitzer et al., 2008). In this study, the fine grain region was selected for the preparation of TEM specimens and was observed by TEM to confirm the Ga ion damage of the specimens.

TEM specimens made by the FIB-ultramicrotome technique were continuously obtained with the slice thickness of below 100 nm. A slice thickness is affected by material, sample size, sample shape and trimmed surface area. In the conventional ultramicrotome technique, it is necessary to make many slices to secure large observation regions, because the region of interest cannot be exactly selected. However, since the FIB-ultramicrotome technique uses FIB to make the region of interest sampled, the trimming size can be reduced to achieve continuous slices that maintain thin slice thickness, as shown in Fig. 2C.

Fig. 3 shows TEM images and selected-area electron diffraction (SAED) patterns, obtained from the TEM specimens made by FIB and FIB-ultramicrotome technique, respectively (Fig. 3A and B). Fig. 3C and D show the SAED patterns obtained from Fig. 3A and B. Both patterns showed the (312), (232) plane and (140) plane of the TiFeSi$_2$ matrix. While SAED pattern of the specimen made by only FIB showed for Si to be amorphized by ion beam damage, SAED pattern of the specimen made by the FIB-ultramicrotome technique showed clearly Si (111) and Li$_{13}$Si$_5$ (620) plane. Typically, Li-silicide is very weak to electron beam. However, the Li-silicide was stable during SAED analysis because it used a very low intensity of electron beam in widely expanded state. Therefore, it is considered that both Si and Li-silicide remain crystalline in the FIB-ultramicrotome specimen.

These results show that the FIB-ultramicrotome technique for TEM analysis can prepare the TEM specimen without artifacts such as the ion beam damage caused by FIB and plastic deformation and specimen tear by the ultramicrotome. To confirm the phase distributions of Si and TiFeSi$_2$ matrix, an energy dispersive spectroscopy (EDS) mapping analysis was conducted. Fig. 4 shows the bright-field–TEM and EDS mapping images of specimens made by the only FIB and FIB-ultramicrotome technique, respectively. As a result, both samples showed two phases of Si and TiFeSi$_2$ matrix, and it was confirmed that TiFeSi$_2$ matrix was maintained even after charging.

In addition, in the specimen prepared by the only FIB technique, Ga elements were detected in the EDS analysis area, but
Fig. 3. Transmission electron microscope images (A and B) and selected-area electron diffraction patterns (C and D) obtained from specimens prepared by focused ion beam (FIB) and FIB-ultramicrotome techniques, respectively.

Fig. 4. Bright-field images (BF) and energy dispersive spectroscopy elemental mapping images of Ti, Fe, and Si for fine-grained regions. (A-D) Focused ion beam (FIB) and (E-H) FIB-ultramicrotome techniques, respectively.
not detected in specimens prepared by FIB-ultramicrotome technique (not shown in this paper). It is thought that when the region of interest is extracted using the FIB, in FIB-ultramicrotome technique the surface of cube sample is damaged by the ion beam but does not affect the inside. It means that the FIB-ultramicrotome specimen shows no ion beam damage and then may can be suitable for EDS analysis. These results as shown in Fig. 3 and 4 prove that the TEM specimen prepared by the FIB-ultramicrotome technique is more preferable to analyze the microstructure of the Si-based anode material, compared with the TEM specimen made by the only FIB technique.

CONCLUSIONS

In this paper, FIB-ultramicrotome technique was proposed as a TEM specimen preparation method for a successful TEM analysis of the Si-based anode material. The specimen prepared by FIB-ultramicrotome technique was compared with that prepared by only FIB technique.

1. From the SAED patterns and the EDS mapping analysis, we estimated the extent of ion beam damage of the two specimens. The SAED pattern of the specimen made by only FIB showed amorphized Si and Li-silicide by ion beam damage and the specimen made by new FIB-ultramicrotome technique showed clearly crystalline Si and Li-silicide. Also, in the specimen made by only FIB technique, Ga atoms were detected in EDS mapping area, but not in the specimen made by FIB-ultramicrotome technique.

2. By using the FIB-ultramicrotome technique, we can extract the specific area using FIB and cut continuous slices mechanically using an ultramicrotome without ion beam damages. In conclusion, this technique is expected to be very useful for preparing TEM specimen of not only Si-based anode materials, but also various materials which are weak to ion beams and in which specific areas need to be observed.

CONFLICT OF INTEREST

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

ACKNOWLEDGMENTS

This research was supported by the MOTIE (Ministry of Trade, Industry & Energy) (10080654) and KSRC (Korea Semiconductor Research Consortium) support program for the development of the future semiconductor device.

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