
博士論文

**Rapid vapor deposition of micrometer-thick silicon-base
porous anodes for lithium secondary batteries**

(リチウム二次電池に向けたマイクロメートル厚さのシリコン系多孔質負極の
急速蒸着技術の開発)

Jungho LEE

李 重昊

A thesis presented in partial fulfilment of
the requirements for the degree of
DOCTOR OF PHILASOPHY

東京大学大学院工学系研究科化学システム工学専攻

Department of Chemical System Engineering

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Dedication

This thesis is dedicated to all my dearly beloved family.

For their endless love, support and encouragement

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Chapter 1 – Introduction

1.1 Electrical energy storage

Over the past decades, current manufacturing and transportation activities has been derived from the discovery and exploitation of primary fossil fuels, such as coals, oils and natural gases. The Energy Information Administration (EIA) recently reported that world's energy consumption will increase by 56% between 2010 and 2040, from 524 quadrillion British thermal units (Btu) to 820 quadrillion Btu [1]. However, global warming, caused by CO₂ emission of burning carbonaceous fuels, and exhaustion of resources have demanded alternative renewable energies like solar and wind power [2-4]. The development of new renewable resources, such as wind and solar power, brought out the research of energy storage because they inherently fluctuate over time [5]. Moreover, new technologies for suppressing emission of CO₂, offer a huge potential of commercializing plug-in-hybrid electric vehicles (PHEV) and electric vehicles (EVs) as new transportation methods [6-8], together with energy storage system (ESS) [9]. Electrochemical energy storage in the form of batteries is fundamental in those energy systems and devices, used as not only national electric smart grids [10], but also the power source from portable devices to medical and aerospace applications. Batteries and electrochemical capacitors are representative of common electrical energy storages in meeting growing electricity demand in large scale market [11, 12]. Among them, the electrochemical capacitors, also known as ultracapacitors, with high specific power, long cycle stability and life time

can be a good choice for applications [13], but suffer from low specific energy [14, 15]. In contrast, batteries with high energy densities attract public attention of realizing a global electric transportation industry due to their great specific energy in spite of low specific power and cycle degradation issues [16, 17]. Therefore, improvement of energy and power densities of batteries for long cycles with low cost is needed for penetration into the global market.

1.2 Rechargeable batteries

A rechargeable battery, also called secondary battery, is a type of batteries, which consists of one or more electrochemical cells operating reversible electrochemical reactions. The first electrochemical cell was discovered by Alessandro Volta in 1800. He repeated and checked Galvani's experiment using dead frogs' legs [18], and demonstrated that electrical energy was derived from spontaneous redox reaction occurring on two different metals, zinc and copper, dipped in an acidic electrolyte. The discovery brought about innovation of development of rechargeable batteries, composed of lead-acid with aqueous electrolytes in 1859, and nickel-cadmium in a potassium hydroxide solution, known as first alkaline battery, was invented by Waldemar Jungner in 1899, and further Thomas Edison patented nickel-iron battery design in 1903. He also tried to commercialize electric vehicles, but the era of electric car came to an end soon because of short battery life and appearance of inexpensive gasoline automobiles [19]. However, his achievements were followed about a century later by introducing first commercial hybrid-electric vehicle (HEV, Prius, Toyota, Aichi, Japan) in 1989, which uses nickel-metal hydride batteries (NH-MH; nickel oxyhydroxide (NiOOH) as cathode and hydrogen-absorbing alloy (MH) as anode) [20]. Moreover, two sodium-base batteries, operated at high temperature, were developed for meeting higher voltage and energy densities in 1970s, but were at a risk of safety concern, and thus limited stationary energy storages [21, 22]. On the other hand, it was reported that the lithium metal would be theoretical ideal material for batteries due to the

highest specific capacity per weight and the highest oxidation electrochemical potential of all known elements, and rechargeable lithium batteries was firstly introduced by G.N. Lewis in 1912, followed by launching first generation Lithium ion batteries (LiBs) by Sony corporation (Tokyo, Japan) in 1991[23]. Figure 1-1 shows that LiBs offer high energy and power densities among various batteries. LiBs have a good balance among the energy density, long cycle life, and stability, thus are promising cell for electric devices [24].

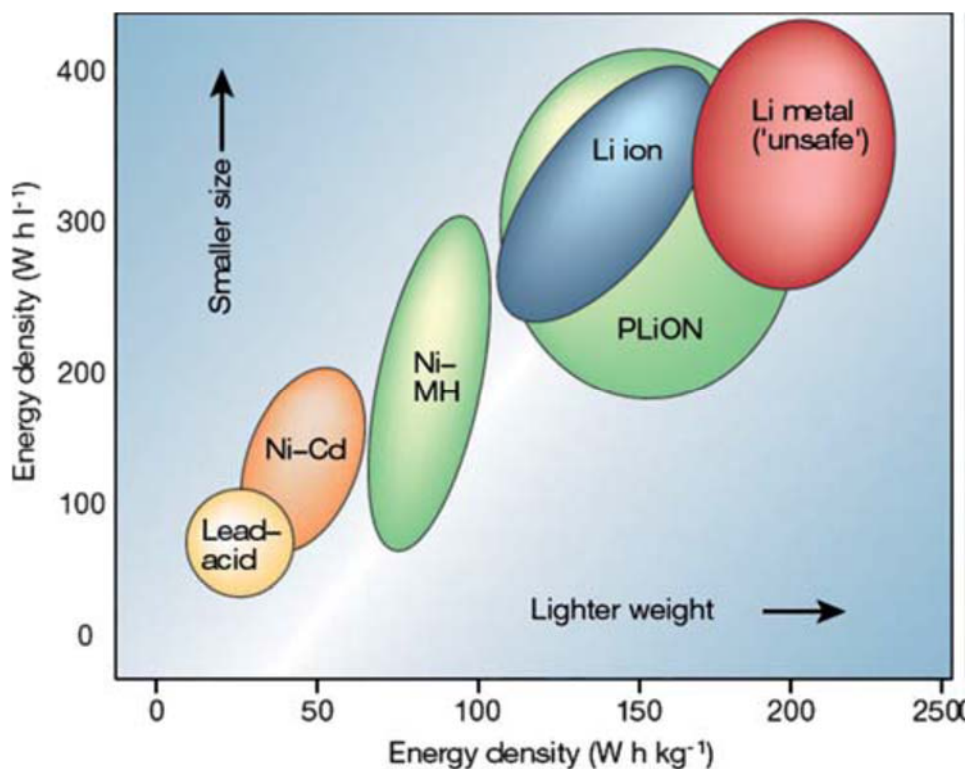


Figure 1-1 Comparison of the different type of batteries in terms of volumetric and gravimetric energy density. Currently, the market share of Ni-Cd, Ni-MH and LiBs is 23, 14 and 63%, respectively *[24].

* Reprinted from Nature, 414, M. Tarascon and M. Armand, Issues and challenges facing rechargeable lithium batteries, 2001, Copyright (2001), with permission from Nature Publishing Group (see Appendix).

1.3 Lithium ion batteries (LiBs)

1.3.1 Fundamentals of LiBs

Lithium ion batteries (LiBs) are one of the rechargeable batteries, base in redox reaction including the intercalation of lithium ions between the cathode and anode as shown in Figure 1-2 [25]. The performance of LiBs can be determined by energy storage density, lithium ion diffusion rate and cycle ability. Energy storage density of LiBs, E , is proportional to the lithium ion storage capacity, C and the difference of electrochemical potential, V , as below [26];

$$E = \int CV \quad (1-1)$$

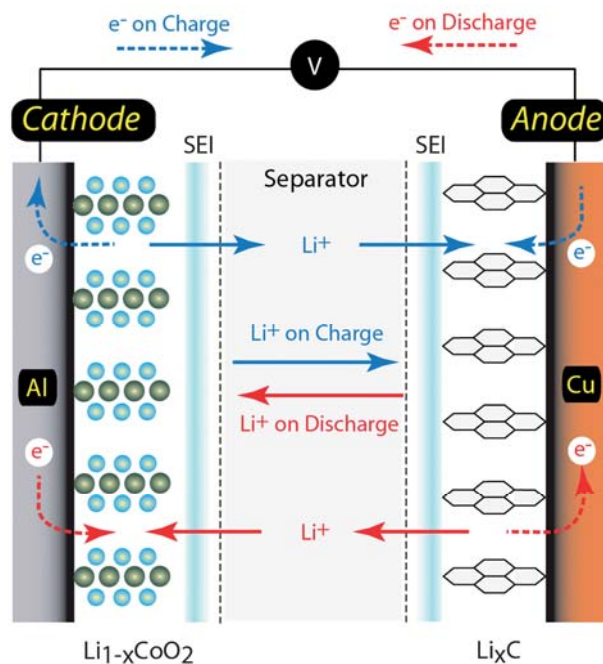


Figure 1-2 Schematic illustration of charge/discharge reaction mechanisms in a conventional lithium ion battery between lithium metal oxide as cathode and graphite as anode. Electron and Li^+ transport during charge/discharge was stated by blue and red arrows, respectively. The electrons travel through the external circuit *[25].

* Reprinted from Energy & Environmental Science, 2, B. J. Landi, M. J. Ganter, C. D. Cress, R. A. DiLeo and R. P. Raffaele, Carbon nanotubes for lithium ion batteries, 2009, Copyright (2009), with permission from Royal Society of Chemistry (see Appendix)

From the equation (1-1), it is clear that high energy storage density can be achieved by either increasing storage capacity or electrochemical potential difference of electrodes, which demands developing next generation electrode materials. Electrochemical potential as a function of storage capacities of Li-base cells is plotted in Figure 1-3 [25]. The anode materials have been explored having a wide range of theoretical capacities depending on materials from conventional Li^+ intercalated graphite LiC_6 of $\sim 372 \text{ mAh g}^{-1}$ to silicon of $\sim 4200 \text{ mAh g}^{-1}$, while the highest capacity of oxides cathode materials is below 500 mAh g^{-1} . Therefore, new generation LiBs with enhanced energy density demand more research effort of the cathode compounds mixture as well as practical use of anode materials with high theoretical capacities. Rate capability of LiBs can also be achieved

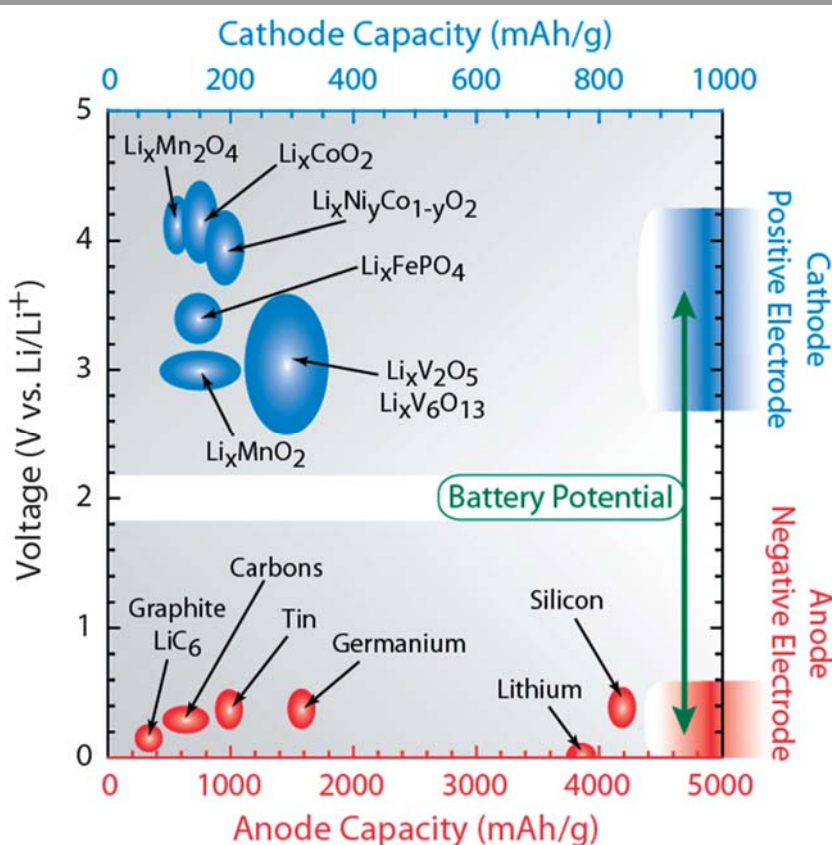


Figure 1-3 Schematic illustration of electrochemical potentials of conventional cathode (blue) and anode (red) as a function of storage capacities presently used for under consideration of Li-based cells *[25].

* Reprinted from Energy & Environmental Science, 2, B. J. Landi, M. J. Ganter, C. D. Cress, R. A. DiLeo and R. P. Raffaele, Carbon nanotubes for lithium ion batteries, 2009, Copyright (2009), with permission from Royal Society of Chemistry (see Appendix).

by increasing lithium ion diffusion rate to electrode during intercalation process, corresponding with conductivity and structure of electrodes [27, 28]. And cycle ability is also highly connected with the structure of electrode material because lithium intercalation process accompanies with a huge volume change, which motivated researchers to study various micro- and nano-structures for electrodes.

1.3.2 Current status of LiBs

The breakthrough in lithium ion battery technology is that lithium metal with safety concern as an anode was substituted by graphite, which accommodates one lithium per three C₆ hexagons, and formed LiC₆ without forming dendrite. The discovery of LiCoO₂ [29, 30] as a cathode and stable liquid organic carbonate solvent enabled the reversible operation at high voltage of ~4.2 V by Sony corporation in 1991 [23]. More recently, several cathode materials, such as LiNiO₂, LiMnO₂ [31], spinel LiMn₂O₄ [32] and LiFePO₄ [33], have been investigated, but even though individual advantages, such as improved safety, reduced cost, and enhanced power density, the energy density has not been improved compared with LiCoO₂ yet. The chemical reaction of typical LiCoO₂ cathodes is described as below;



In case of the anode material, the various carbonaceous materials with a low potential versus lithium were introduced, and graphite, carbon fibers and mesocarbon microbeads (MCMB) were a commercial success in LiBs market [34]. Graphite provides several advantages in the view point of cost, abundance, non-toxicity and structural stability for repeated cycles [35]. In addition, lithium intercalation to graphite causes small volume change (9–10%), with the typical reaction expressed as;



However, graphite anodes possess the disadvantages of low capacity ($\sim 375 \text{ mAh g}^{-1}$) with a maximum configuration of one lithium atom to every six carbon atoms of LiC_6 and safety issues related to Li deposition [36, 37], which has limited its feasibility meeting with the growing demand in high power technology for electric vehicles (EVs) or hybrid electric vehicles (HEVs). Moreover, a low operating voltage around 100 mV (vs. Li^+/Li) causes the reaction of graphite with electrolyte, resulting in reducing battery performances as well as serious safety concerns such as thermal runaway.

Carbonaceous materials have been extensively researched, such as one-dimensional (1-D), two-dimensional (2-D) and porous carbon, trying to create more surface areas for lithium storage and to increase the energy and power densities [38]. One-dimensional structured carbon materials were well known as carbon nanotubes (CNTs) and carbon nanofibers (CNFs). The anode of CNTs achieved $\sim 1116 \text{ mAh g}^{-1}$ of reversible capacity using various treatments, such as ball-milling [39], acid oxidation [16], and metal oxide cutting [40], while the reversible capacities of CNFs-anodes by electrospinning/carbonization process combined with chemical treatments was found to be $\sim 430 \text{ mAh g}^{-1}$ at 50 mA g^{-1} . [41-43]

Graphene, on the other hand, is representative 2-D structured carbonaceous material with high surface area including electrical and thermal conductivity. Graphene or graphene-base composited material with active metal (Sn [44], Sn-Sb [45] and Si [46]) and/or metal oxide (Co_3O_4 [47, 48], TiO_2 [49], Fe_3O_4 [50, 51], Mn_3O_4 [52], CuO [53] and SnO_2 [54, 55]) has been researched. The graphene as anode material was currently reported having $\sim 500 \text{ mAh g}^{-1}$ after 30 cycles [56], and $\text{SnO}_2/\text{graphene}$ anodes showed more enhanced performance high reversible capacity of $\sim 800 \text{ mAh g}^{-1}$ at 50 mA g^{-1} , and $\sim 570 \text{ mAh g}^{-1}$ after 30 cycles with 70% capacity retention, while high irreversible capacity and unstable performance still existed [57], which needs to further researches. Meso-porous carbon anodes with different pore sizes from nano- to micro-scale have been also researched with advantages of suppressing volume change by high surface area, which achieved reversible capacity of $\sim 800 \text{ mAh g}^{-1}$

g^{-1} at 100 mA g^{-1} after 20 cycles because of minimized ion transport resistance around the pores with ordered porous structures [58, 59].

In summary, carbonaceous anode materials with different nanostructure, such as 1-D, 2-D and ordered porous carbon) exhibited a good electrochemical performance with individual their advantages, but poor rate performance (except for graphene), low volumetric energy density by SEI, and low capacity was still in common challenge for practical usage suitable for high-powered-applications.

1.4 Silicon (Si) anodes in LiBs

1.4.1 Advantages of Si for anode material

Silicon (Si) has been currently considered to be a promising anode material, substituted for conventional carbonaceous materials like graphite, and thus probably most studied anode material. Silicon exhibits low lithiation potential (vs. Li/Li^+) below $\sim 300 \text{ mV}$, which makes it relatively stable and safe as an anode material compared to graphite with a high energy density material (up to 120

Table 1-1 Different properties of various anode materials (Th. Sp. Cap. Indicated theoretical specific capacity) [64]

Materials	Li	C	$\text{Li}_4\text{Ti}_5\text{O}_{12}$	Si	Sn	Sb	Al	Mg
Density (g cm^{-3})	0.53	2.25	3.5	2.3	7.3	6.7	2.7	1.3
Lithiated phase	Li	LiC_6	$\text{Li}_7\text{Ti}_5\text{O}_{12}$	$\text{Li}_{4.4}\text{Si}$	$\text{Li}_{4.4}\text{Sn}$	Li_3Sb	LiAl	Li_3Mg
Th.Sp.Cap. (mAh g^{-1})	3862	372	175	4200	994	660	993	3350
Volume change (%)	100	12	1	420	260	200	96	100
Potential vs. Li (V)	0	0.05	1.6	0.4	0.6	0.9	0.3	0.2

Wh kg^{-1} [60]), Besides, Si can accommodate up to 4.4 lithium ions per unit (the stoichiometry of the lithiated to $\text{Li}_{4.4}\text{Si}$ alloy), corresponding with the highest theoretical gravimetric capacity of 4200 mAh g^{-1} , which is ten times larger than that of conventional graphite [61-63], and high volumetric capacity of 9786 mAh cm^{-3} , derived from the calculation on the initial volume of Si [64]. Because of this, Si anodes possess an enough potential to apply high-powered storage system, such as electric vehicles (EVs). In addition, Si is abundant material on earth with a low cost and environment-friendly, and thus has been used for semiconductor devices, and also the delithiation potential of Si occurs at a relatively low voltage of ~ 0.4 V [65], as shown in Table 1-1 [64]. When lithium is inserted into Si anode, Si experiences a series of phase transformations from electrochemical lithiation/delithiation

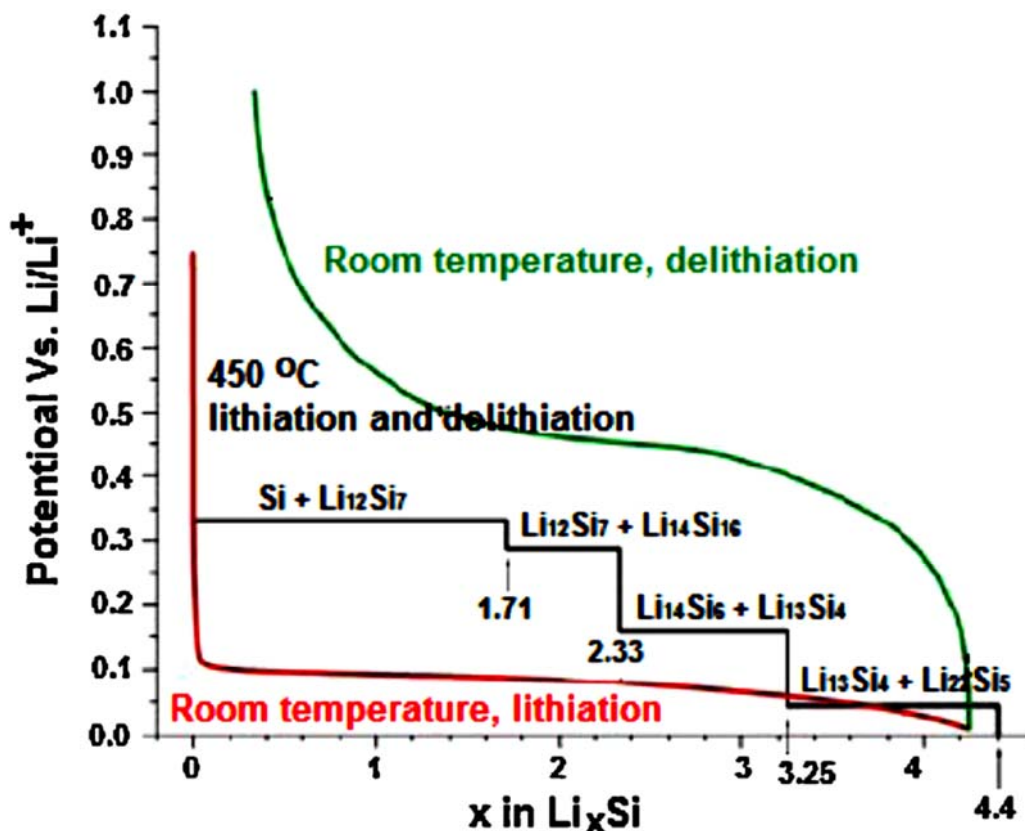


Figure 1-4 Electrochemical lithiation/delithiation curves of Si anodes at high temperature of 450 °C and room temperature. Multiple step curves (black) indicated at 450 °C, while lithiation/delithiation of crystalline Si at room temperature is plotted, as red and green line, respectively * [64].

* Reprinted from Nano Today, 7, H. Wu and Y. Cui, Designing nanostructured Si anodes for high energy lithium ion batteries, 2012, Copyright (2012), with permission from Elsevier (see Appendix).

diagram of Si, as shown in Figure 1-4 [64], which is reported multiple step curve of voltage plateau at high temperature of 450 °C [65], but crystalline Si (cr-Si) transformed into amorphous phase, and further amorphous or crystalline lithium silicide phase depending on lithiated potential at initial lithium insertion, while amorphous Si (a-Si) goes through amorphous or crystalline lithium silicide phase at room temperature.

1.4.2 Challenges for Si anodes

As I stated above, Si is more attractive material than any other materials as anode for LiBs. However, this high specific capacity is realized by inserting a large amount of Li^+ into Si materials, which accompanied with large volume change (300–400%, depending on the state of Li_xSi), supported by various microstructural analysis methods, such as X-ray diffraction [66, 67] and nuclear magnetic resonance [68], and subsequently Si electrodes do not withstand the heavy strain of such volumetric expansion and break down, resulting in pulverization and delamination of the whole structure [69-71]. Further capacity losses are caused by solid electrolyte interphase (SEI) formation, while delamination results in loss of electrical contact with the current collector. The fundamental three issues of Si failure phenomena, as an anode for LiBs, were summarized in illustrated Figure 1-5 [64]. The large volume change induces large stress and strain, which causes cracking and pulverization of the Si anode, results in loss of electrical contact. Moreover, such a large volume change causes disconnection of Si nano- or micro-particles in case of nanostructured Si materials or assembled materials of Si nanoparticles. Another issue is the formation of passivating film before lithium insertion to Si at the surface, called to SEI, induced by decomposition of the organic electrolyte because the potential of lithium insertion is lower than reduction potential ≥ 1 (versus Li/Li^+) of

organic solvent in commercial electrolyte [72], such as ethylene carbonate (EC), propylene carbonate (PC), dimethyl carbonate (DMC), diethyl carbonate (DEC), and so on. The SEI films were reported to consist of lithium carbonate (Li_2CO_3), various lithium alkylcarbonates (ROCO_2Li), mainly, including Li_2O , LiF or LiCl , and nonconductive polymers [73-78].

Consequently, various approaches based on control of Si-base material structure or composition as well as on the operation control of lithiation/delithiation cutoff voltage, were carried out to accommodate and buffer the effects of volume change. Moreover, conductive materials for enhanced lithium diffusion rate, electrolyte additives for improvement of SEI and polymer binder for enhanced adhesion between electrodes and current collectors have been researched continuously.

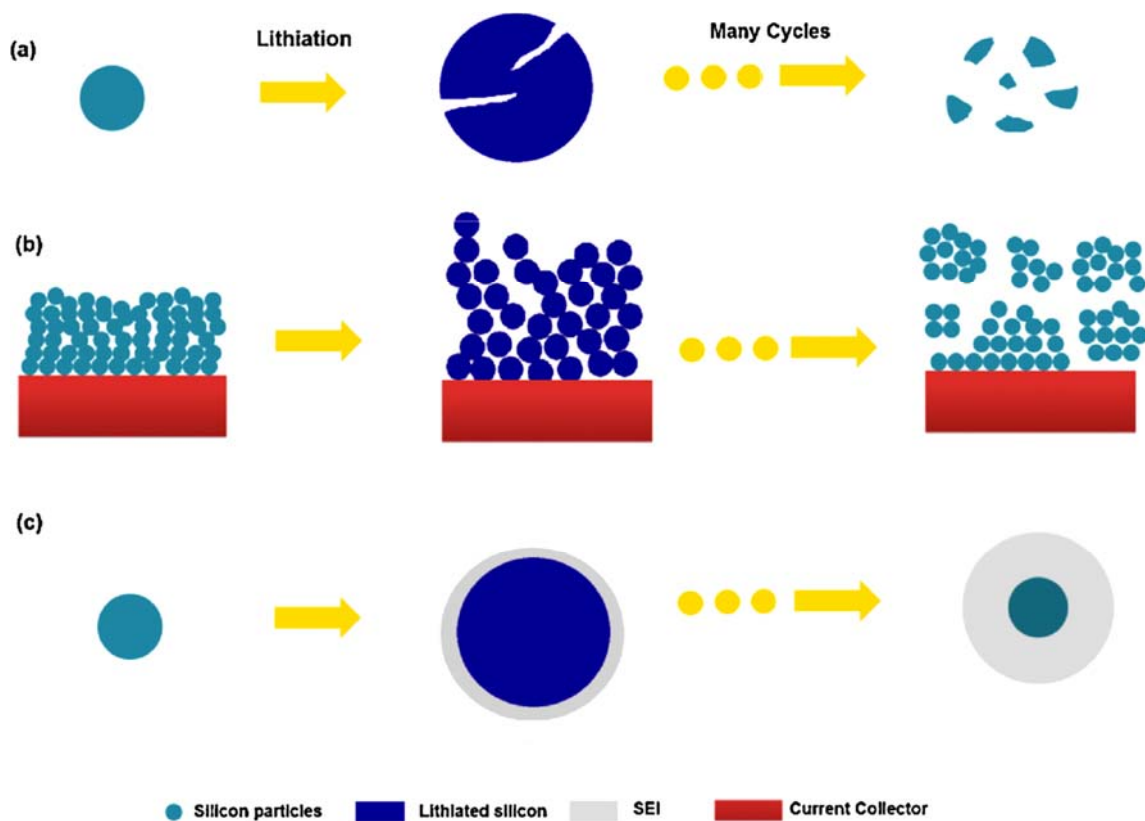


Figure 1-5 Illustration of Si-electrode failure mechanism (a) Material pulverization. (b) Disconnection of Si nano- and micro-particles. (c) SEI regeneration *[64].

* Reprinted from Nano Today, 7, H. Wu and Y. Cui, Designing nanostructured Si anodes for high energy lithium ion batteries, 2012, Copyright (2012), with permission from Elsevier (see Appendix).

The various structures of Si-base materials, such as thin film, micro-/nano-particles, nanowires (SiNWs), and composite materials of Si with carbon and/or silicides, were introduced and charged issues. One-dimensional SiNWs with various structures, introduced by Yi Cui group [79], were currently charged issues, achieving excellent electrochemical performance for long cycles. They suggested one dimensional structure of SiNWs relax the stress during volume change, and hollow structured double-walled Si nanotubes with outside SiO_x surface layer also make stronger surface film, resulting in no significant capacity fade [80]. Another issue of Si-base materials is composite of Si with carbon and/or silicides, such as electrodeposited Si-O-C composite film [81, 82] and metal silicide-coated Si films [83], which enhanced electrochemical performance for long cycles because of their unique structure and control of SEI.

1.5 Research scope and objectives

The previously presented studies mainly focused on the electrochemical performance of Si materials, and thus various structures and compositions, focusing on realizing accommodates stresses induced by volume change, were reported. As a result, high gravimetric capacities of Si-base anode materials were achieved successfully, but the weight of heavy current collector was, sometimes, neglected for very thin active layers. Commercialized graphite anode, Sony 18650 cell as currently reported, consists of LiCoO₂ as cathode and graphite as anode. The physical properties of the graphite anode of the cell were the mass of 5.7 g, 11 mg cm⁻² of areal density, 193 μm in thickness (including current collector), and dimensions of the anode is 52.9 × 5.7 cm, and areal capacity is ~4 mAh cm_{anode}⁻². Even though theoretical capacity of Si is ten times higher than graphite, the thickness of Si has to approach ~10 μm for practical use in LiBs market, supposing that conventional Cu current collector has ~100 μm in thickenss. However, previously reported Si nanomaterials have still low areal density of ~250–300 μg cm_{anode}⁻², showing low areal capacity of 0.2–0.4 mAh cm_{anode}⁻², even though high

gravimetric capacity of 4000 mAh g_{Si}⁻¹. Furthermore, various fabrication methods of Si anode were reported, such as high-energy mechanical milling as top-down approach, and chemical vapour deposition (CVD), sol-gel synthesis, hydrothermal synthesis and electron-beam deposition (EB) as bottom-up approach. Especially, silane (SiH₄) gas and silicon tetrachloride (SiCl₄) have been used as a precursor by CVD and EB method, which was time-consuming process with low yield.

In summary, the researches of various structured Si-base materials have successfully demonstrated the possibility of Si for usage of next generation anode material in LiBs, while further researches are also needed in some respects. The structure design of Si was the most progressive in qualitative level, but quantitative understanding, such as size-dependence of particles and pores, has been still investigated quantitatively. Furthermore, although various approaches were reported to understand and improve of passivating surface film (SEI), high irreversible capacity of Si anodes demands further studies. Finally, the development of large-scale, low-cost fabrication and enhanced mass production for Si materials with desirable electrochemical performance is the most important challenge of all to commercialize Si anode for LiBs.

Chapter 2 – Micrometer-thick porous Si films by rapid vapour deposition (RVD)

2.1 Introduction

The current anode material of commercial LiBs is usually graphite because of its long cycle life, abundance and relatively low cost [84]. However, graphite anodes possess the disadvantages of low capacity (375 mAh gc^{-1}) and safety issues related to Li deposition [37]. Thus, there has been a growing interest in developing alternative anode materials with low cost, good safety, high energy density and long cycle life. Si is an attractive alloy-type anode material with a theoretical specific capacity ($4200 \text{ mAh g}_{\text{Si}}^{-1}$) based on the stoichiometry of the alloy $\text{Li}_{22}\text{Si}_5$ [61-63]. However, this high specific capacity is realized by inserting a large amount of Li^+ into the active material, causing volume expansion of $\sim 300\text{--}400\%$. Si electrodes easily break down by the heavy strain of such volumetric expansion, which triggers pulverization and delamination of the whole structure [69-71], resulting in a loss of electrical contact with the current collector.

To overcome pulverization of electrodes during expansion and shrinkage, many groups, starting with the study by Huggins and coworkers [61], have developed various approaches to control electrode structures, initially thin films [85, 86], micro-particles [87], nanowires [79, 88-91], nanoparticles [92, 93], pillar formation [94], $\text{NiSi}_x\text{-Si}$ core-shell nanowires [83], and other structures [95, 96]. Full cells containing cr-Si core-shell nanowires [97] have been reported. High capacities were reported for such

structures; however, sometimes the weight of heavy current collectors was neglected for very thin active layers, and the studies about large-scale, mass production and low-cost fabrication are currently insufficient compared with structure design reports.

In this chapter, I rapidly deposited 3–14 μm -thick porous Si films in 1 min or less by a physical vapour deposition method called rapid vapour deposition (RVD), focusing on developing low-cost fabrication method of Si films with large-scale. Such a high deposition rate is achieved by heating the source Si to 2000–2400 $^{\circ}\text{C}$, well above its melting point of 1414 $^{\circ}\text{C}$. Control over the amorphous/crystalline structure, film porosity, structure, surface roughness, and Si/Cu interface is realized by maintaining the temperature of the Cu substrate at 100–500 $^{\circ}\text{C}$ during and after deposition. In particular, control over the roughness of the growing Si films caused by the shadowing effect [98, 99], is important to tailor the film microstructure. The electrochemical behavior of the Si films was investigated by a half-cell test using Li as the counter electrode and discussed in relation to their microstructure.

2.2 Materials and methods

2.2.1 Si film fabrication

Circular Cu plates (15 mm in diameter, 0.5 mm in thickness) instead of thin Cu foils were used as substrates for porous Si films to make the handling (adjusting the size to fit the coin cells) easier. Before Si deposition, the Cu plates were sonicated in isopropanol for 10 min and then exposed to UV- O_3 for 3 min to remove organic contaminants on their surface. Then, the Cu plates were annealed under hydrogen (5 vol% H_2 / Ar, 1 atm) at 800 $^{\circ}\text{C}$ for 10 min to reduce the oxide layer on the Cu surface. Finally, Si was deposited on the Cu substrates by RVD.

Figure 2-1a shows a schematic diagram of the RVD system. The Si source for RVD was prepared by first immersing a Si wafer (CZ p-type, resistivity of 10–20 Ω cm) in 5 wt% hydrogen fluoride (HF) solution for 1 min, and then rinsing it with purified water for 1 min. The wafer was ground into Si powder using mortar and pestle and then loaded into a carbon boat. The Si source was heated by resistive heating of the carbon boat under 0.1 Torr Ar to 2000–2400 $^{\circ}$ C (boat temperature, T_{boat}) to increase its vapor pressure and thus the deposition rate. A Cu substrate was positioned in the chamber and kept at 100–500 $^{\circ}$ C (substrate temperature, T_{sub}) during RVD to suppress the surface diffusion of

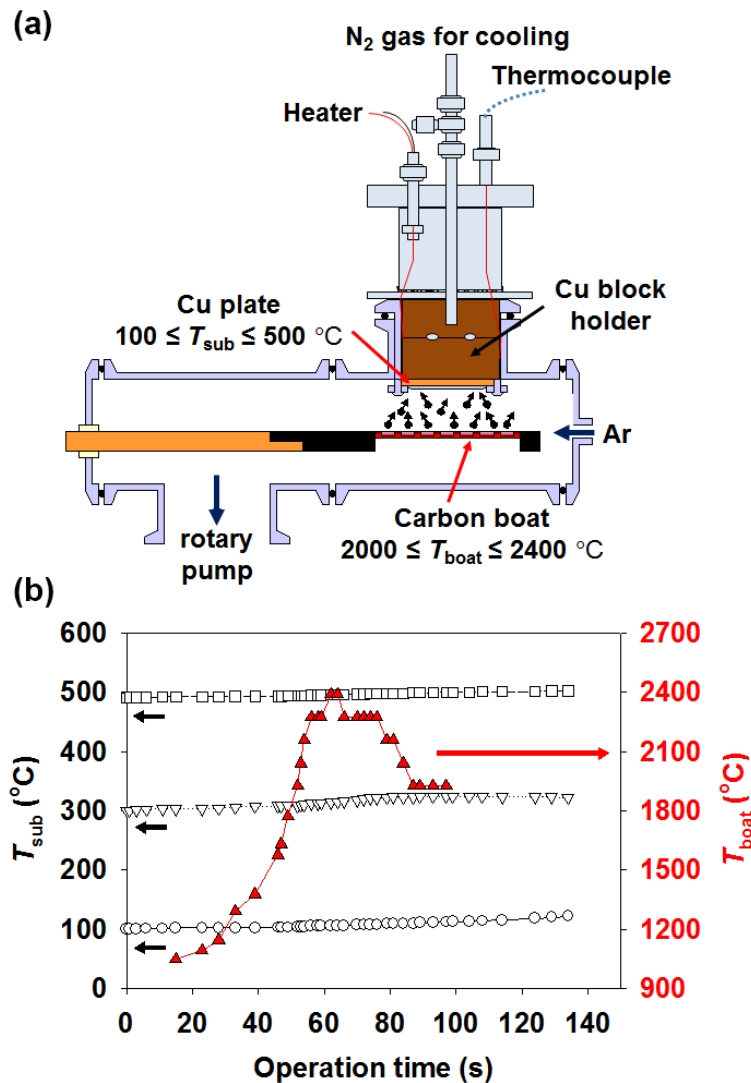


Figure 2-1 (a) Schematic diagram of the RVD system. (b) Typical time profiles of the temperatures of the Si source and Cu plate during RVD at $P_{\text{boat}} = 1600$ W. T_{boat} is shown for a run with $T_{\text{sub}} = 300$ $^{\circ}$ C and is similar for different T_{sub} .

Si, which induces rough and porous structure in the deposited Si films [83, 98]. To maintain the Cu substrate at constant temperature under the strong thermal radiation from the Si source, we designed a substrate holder made of a block of Cu with a large heat capacity that contained an embedded ceramic heater and cooling line for N₂ gas. After removing the sample from the RVD system, some of the samples were further annealed under 4 vol% H₂/Ar at ambient pressure at 200–600 °C (annealing temperature, T_{an}) for 10 min to form a copper silicide (CuSi_x) intermixed layer to improve the adhesion between the Cu substrate and Si film.

Figure 2-1b shows temperature profiles of the Si source and Cu substrate as a function of operation time during deposition. The boat heating power, P_{boat} , was increased manually, and when the source temperature was well above the melting point of Si, deposition began and was then completed after about 1 min. Because of the Cu block holder, the substrate temperature was able to be held at 100, 300 or 500 °C as desired (Figure 2-1b). Before and after Si deposition, each sample was weighed by a microbalance with a sensitivity and precision of 10 µg. Effective thickness t_{eff} was calculated using a value of 2.33 g cm⁻³ for bulk Si crystal. By changing the substrate temperature, a series of Si films with different morphology and electrochemical performance were fabricated.

2.2.2 Characterization

X-ray photoelectron spectroscopy (XPS; JPS9010 TR; JEOL, Akishima, Japan) with a monochromatised Mg K α X-ray source was used to investigate the surface condition of treated and untreated Cu substrates. Microstructural analysis of the fabricated Si films was carried out using X-ray diffraction (XRD; ATX-G; Rigaku, Akishima, Japan) and laser micro-Raman spectroscopy (HR-800; Horiba, Kyoto, Japan). The microstructure and composition distribution of the Si/Cu samples were analyzed by scanning electron microscopy (SEM; S-4800; Hitachi, Tokyo, Japan) with energy-

dispersive X-ray spectroscopy (EDS, EDAX Genesis; AMETEK, Elancourt, France).

2.2.3 Electrochemical characterization

Capacity and cycle performance measurements were performed by collaborative researchers, Mr. Kenji Nakane and Mr. Shingo Matsumoto at Sumitomo Chemical Co. Ltd., using the Si films on Cu substrates as a working electrode with Li metal (0.5-mm-thick foil) as the counter electrode for R2032-type coin-shaped half cells. LiPF₆ solution (1 M) in a 1:1:1 (v/v) mixture of ethylene carbonate (EC), dimethyl carbonate (DMC), and ethyl methyl carbonate (EMC) was used as the electrolyte. Half-cell tests were carried out in the range from 0.005 to 1.000, 1.5000 or 2.000 V vs. Li/Li⁺ with different charge rates of 0.05C–0.2C at a constant temperature of 25 °C. C-rate was determined using the weight of Si and a theoretical capacity of 4200 mAh g_{Si}⁻¹. After different numbers of cycles, electrochemical impedance spectroscopy (EIS) measurements were performed using a Solartron[®]1287 electrochemical interface coupled to a Solartron[®]1260 frequency response analyzer (Ametek, Elancourt, France) in the 10⁶ to 10⁻² Hz frequency range.

2.3 Structure on Si film deposited on Cu substrate by RVD

Figure 2-2a shows a top-view SEM image of an as-purchased Cu plate, which clearly contains lines patterned at intervals of about 2 μm on its surface. Figure 2-2b and c show top-view and cross-sectional SEM images of a 14- μm -thick Si film deposited on a Cu substrate. The Si film contained

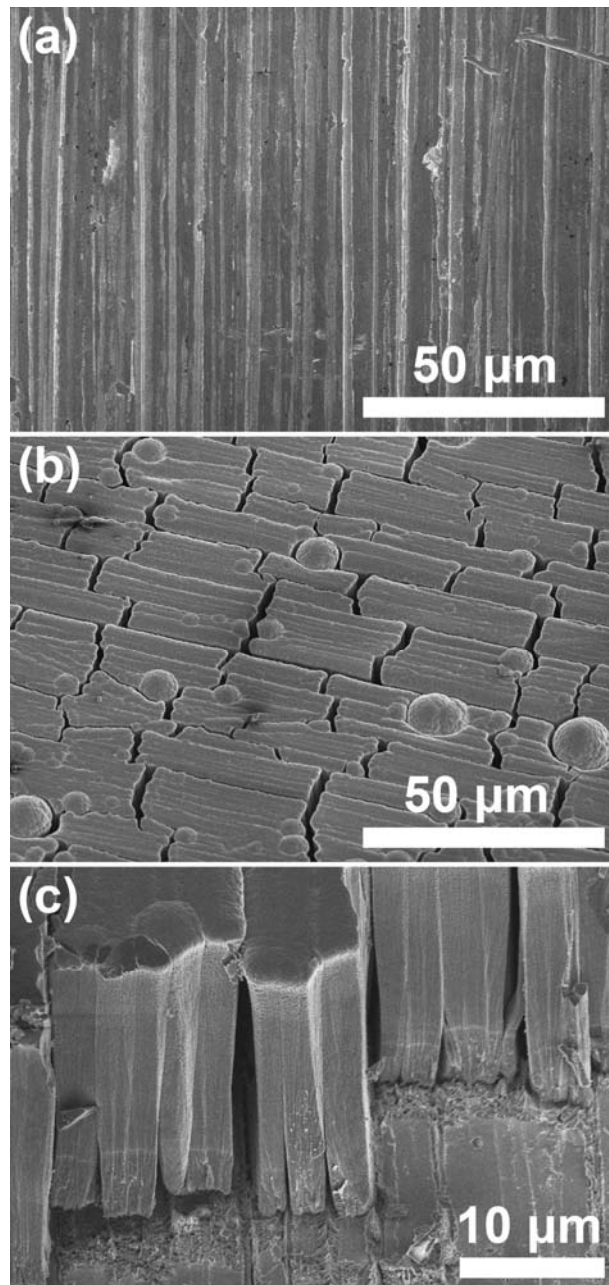


Figure 2-2 (a) A top-view SEM image of an as-purchased Cu plate. (b) Top-view and (c) tilted-view SEM images of a typical Si film ($t_{\text{act}} = 14 \mu\text{m}$) prepared by RVD in 10 s at a high P_{boat} of 1600 W.

many 2- μm -thick stripes that were aligned in one direction. It is clear that the striped structure originates from the line patterns of the as-purchased Cu plates. The Si film also contained cracks in the in-plane and vertical directions to the alignment of the stripes. These cracks were at rather random positions but appeared at intervals of around 20 μm . The surface of the Si film formed with a high P_{boat} of 1600 W contained numerous protrusions with a height of a few μm . If the source heating power was too high, although Si deposition occurred within 10 s, it caused the Si source to boil and spread Si droplets on the Cu substrate, resulting in protrusions in the Si film (Figure 2-3a and c). The cycle performance of such films was very poor regardless of T_{an} for post-annealing treatment; maximum initial charge and discharge capacities were ~ 2000 and ~ 600 $\text{mAh g}_{\text{Si}}^{-1}$, respectively, and

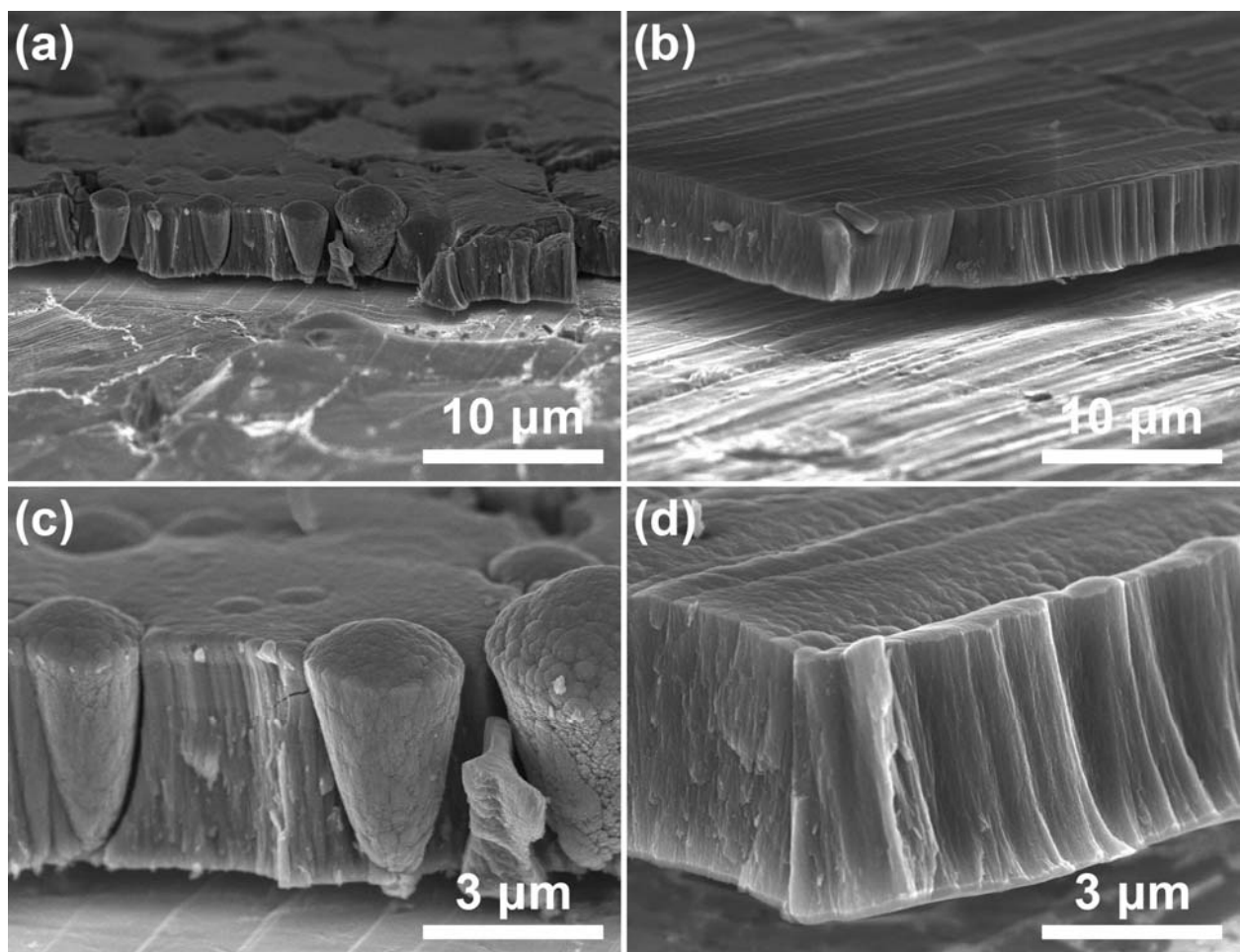


Figure 2-3 Low-magnification cross-sectional SEM images of Si films ($t_{\text{act}} = 3\text{--}5$ μm) deposited with (a) high P_{boat} of 1600 W and (b) moderate P_{boat} of 1300 W. (c, d) High-magnification cross-sectional SEM images of (a) and (b), respectively.

capacity decreased within a few cycles (Figure 2-4). At a moderate P_{boat} of 1300 W, porous Si films without protrusions were formed although the deposition time was a little longer (1 min) (Figure 2-3b and d). The weight of Si deposited at $P_{\text{boat}} = 1300$ W was about 1 mg on average for a circular film with a diameter of 14 mm. This areal weight ($0.6\text{--}0.7$ g cm $^{-2}$) corresponds to $t_{\text{eff}} \sim 3$ μm if the film has the same mass density as bulk crystalline Si (2.33 g cm $^{-3}$), and the actual thickness t_{act} is larger because of the porous structure of the film. Such films were showed some improvement of cycle performance compared with those deposited at higher P_{boat} ; maximum initial charge and discharge capacities were ~ 2800 and ~ 1300 mAh g $_{\text{Si}}^{-1}$, respectively, but capacity reduced within a few cycles

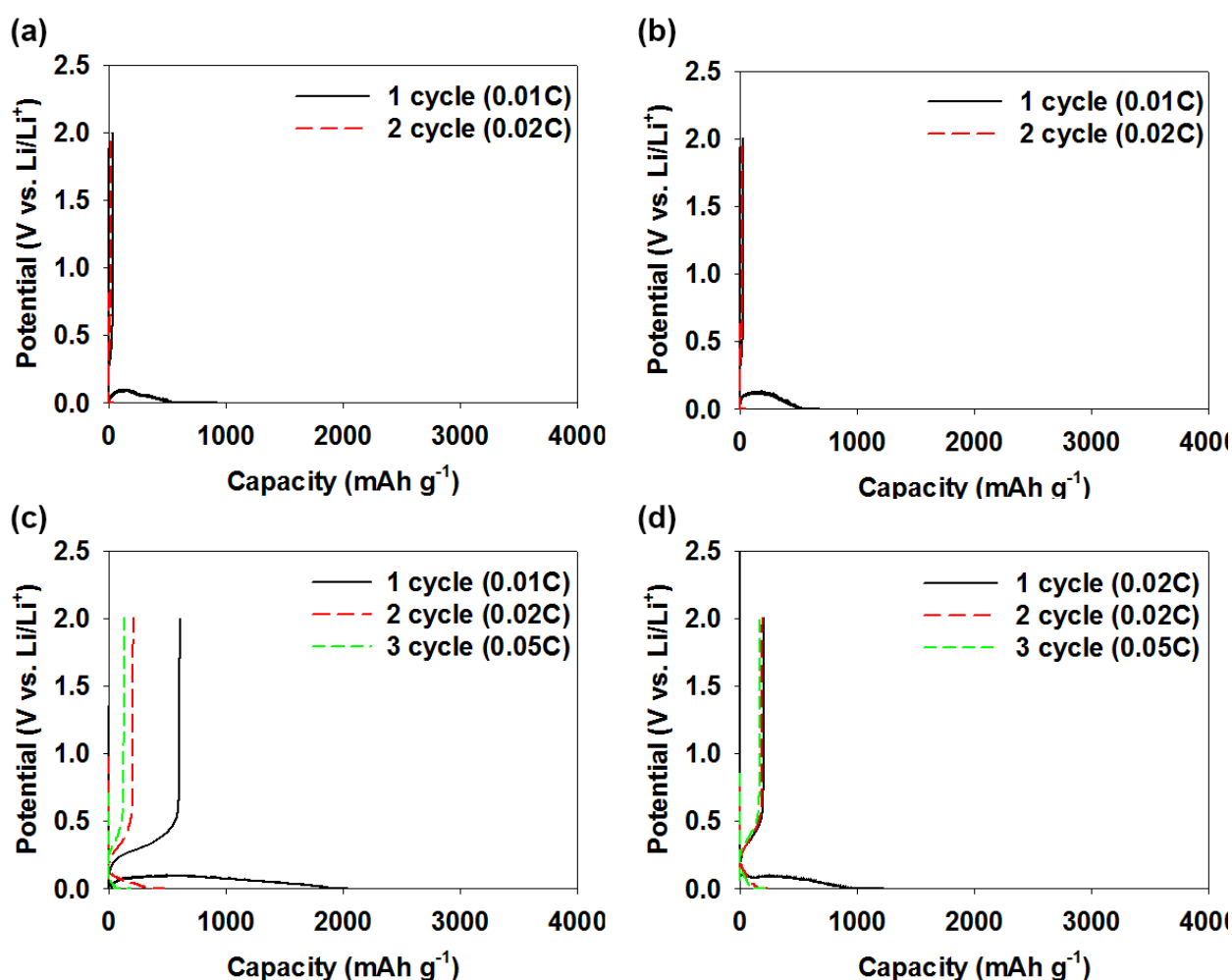


Figure 2-4 Voltage-capacity curves of thick Si films (~ 10 μm) deposited at high $P_{\text{boat}} = 1600$ W and low $T_{\text{sub}} = RT$ on as-purchased Cu substrates. (a) An as-deposited Si film, and Si films post-annealed at (b) 200, (c) 400, and (d) 600 $^{\circ}\text{C}$. Effective Si thicknesses were $t_{\text{eff}} =$ (a) 9.6, (b) 11.0, (c) 10.8 and (d) 10.3 μm .

(Figure 2-5). Such rapid capacity fade can be attributed to the insufficient adhesion of the Si films to the Cu substrates, as can be seen in the SEM image (Figure 2-3) that show the Si films detaching from the Cu substrates (note that the cross-sections were prepared by vending the 0.5-mm-thick Cu substrate, resulting in such detachment). Post-annealing treatment was used to improve electrode performance by forming a CuSi_x intermixed layer to prevent detachment of the Si film from the Cu substrate. Because it is known that solid-state crystallisation of amorphous Si takes about 10 min at 700 °C [100], we annealed the samples for 10 min at $T_{\text{an}} = 200, 400,$ and 600 °C to avoid crystallization of the Si films (Figure 2-5b–d). The Si film post-annealed at $T_{\text{an}} = 600$ °C (Figure 2-

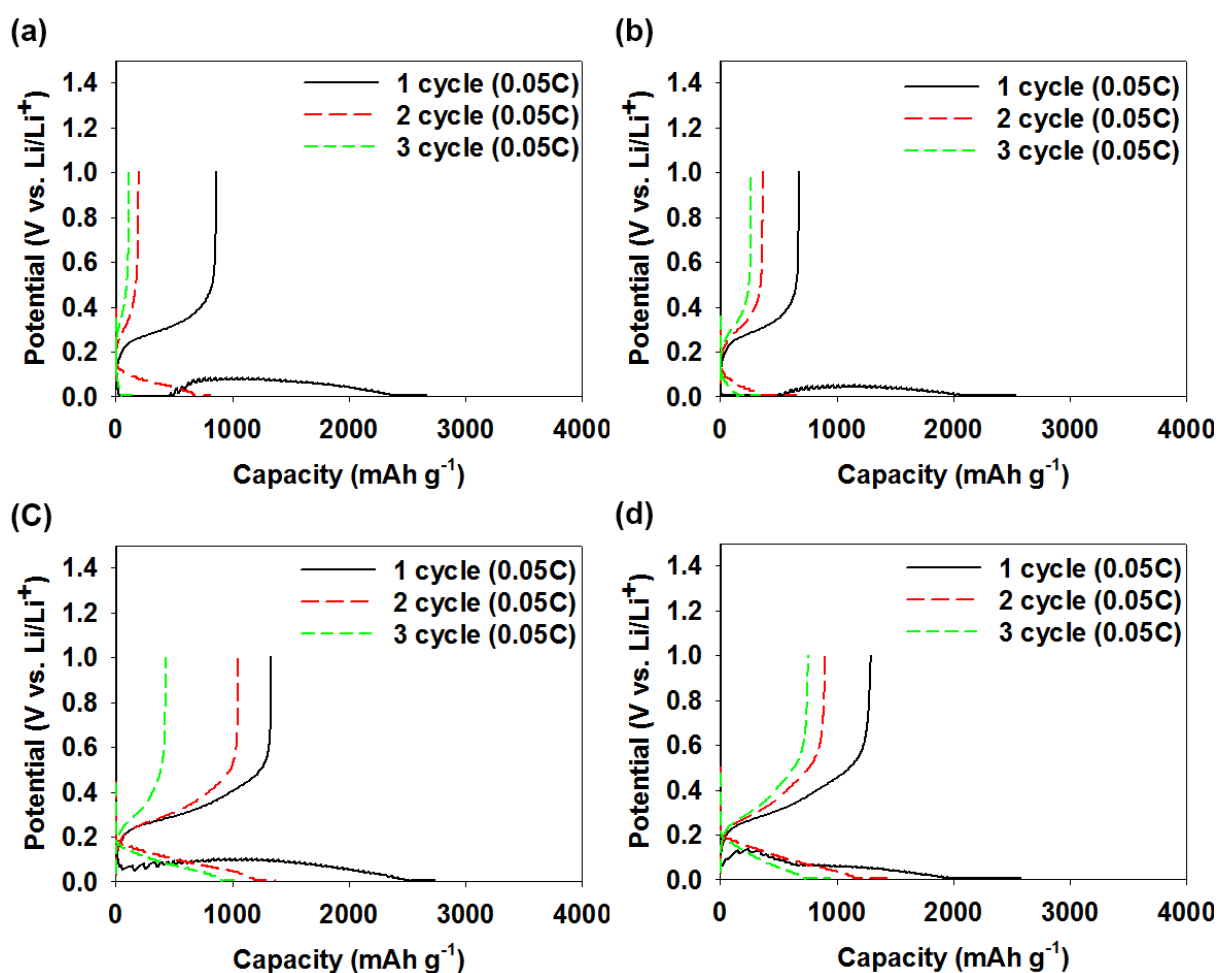


Figure 2-5 Voltage-capacity curves of thick Si films (t_{eff} of ~ 3 μm) deposited at high $P_{\text{boat}} = 1300$ W and low $T_{\text{sub}} = RT$ on as-purchased Cu substrates. (a) An as-deposited Si film, and Si films post-annealed at (b) 200, (c) 400, and (d) 600 °C. Effective Si thicknesses were $t_{\text{eff}} =$ (a) 3.2, (b) 3.1 (c) 2.8 and (d) 3.4 μm .

5d) showed the best initial capacity and capacity retention, so we used $T_{\text{an}} = 600$ °C as the standard post-annealing temperature.

Figure 2-6a shows a top-view SEM image of a Si film with t_{eff} of ~ 3 μm deposited at a moderate P_{boat} of 1300 W. Compared with the 14 μm -thick Si film deposited at a high P_{boat} of 1600 W (Figure 2-2), a similar striped structure was realized but it did not contain any Si protrusions. The morphology of the Si films with t_{eff} of ~ 3 μm depended on T_{sub} during deposition (Figure 2-6b–d). A low T_{sub} of 100 °C suppressed the surface diffusion of deposited Si atoms on the surface of the Cu substrate and growing Si film, which made Si film have a porous structure. These Si films were composed of

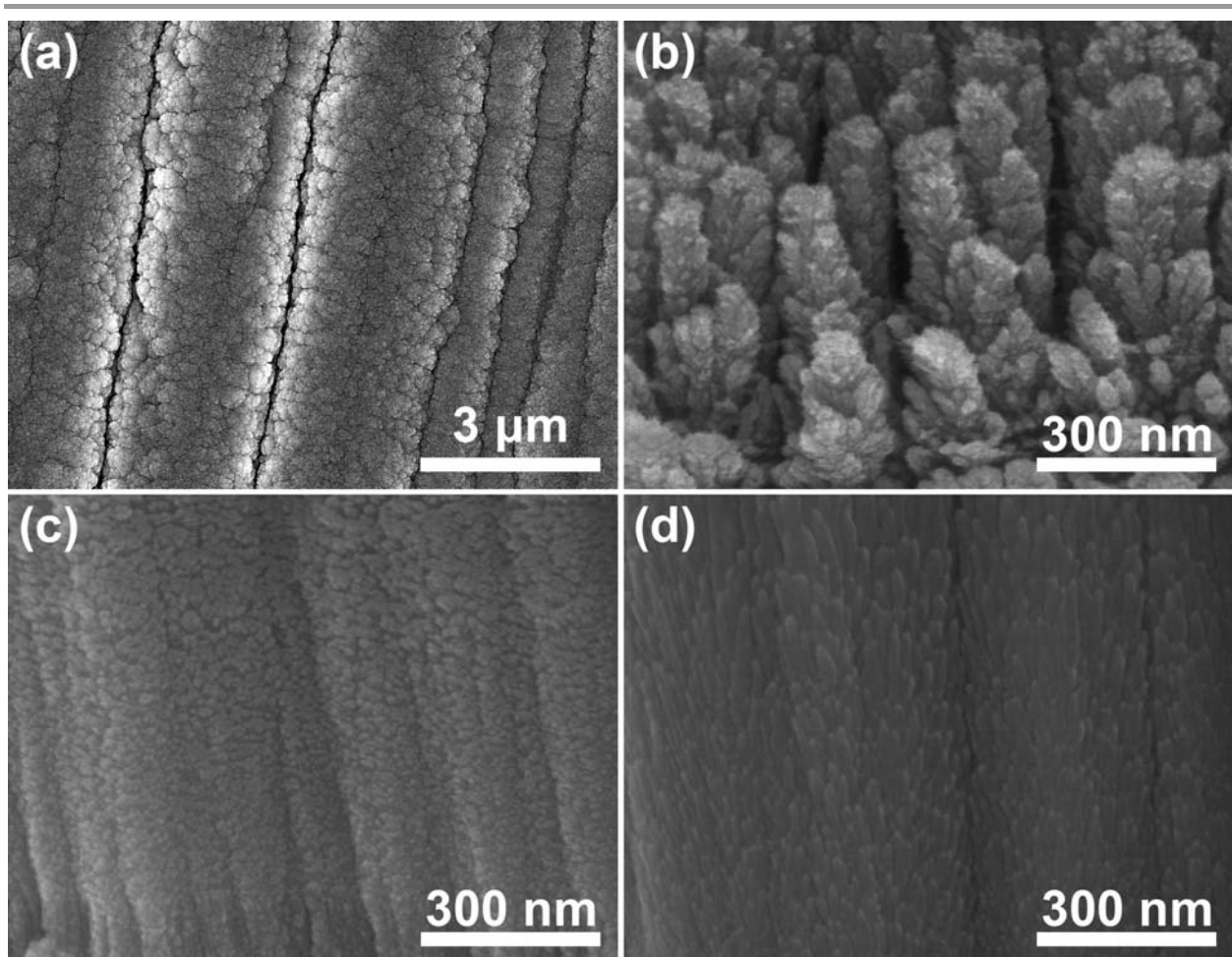


Figure 2-6 (a) Top-view SEM image of a Si film deposited at $T_{\text{sub}} = 300$ °C. Cross-sectional SEM images of Si films deposited at T_{sub} of (b) 100, (c) 300, and (d) 500 °C. All of the films were deposited by RVD in 1 min at a moderate P_{boat} of 1300 W.

numerous Si pillars with a lateral size ~ 50 nm and many pores between Si pillars, which were obviously induced by a shadowing effect (Figure 2-6b). In contrast, T_{sub} of 300 and 500 °C during deposition yielded denser porous Si films (Figure 2-6c and d, respectively). The striped, porous structures will not only facilitate rapid diffusion of Li^+ , but also relax the stress caused by volume changes during cycling. The electrochemical performance of these Si films should be modulated by their different porous structure.

We next evaluated the film density by measuring the weight and volume of the films. Typical thickness distribution is shown in Figure 2-7, from which we determined film volumes. The film density and porosity obtained using different conditions are summarized in Table 2-1. Here we define the porosity p_{film} as (1),

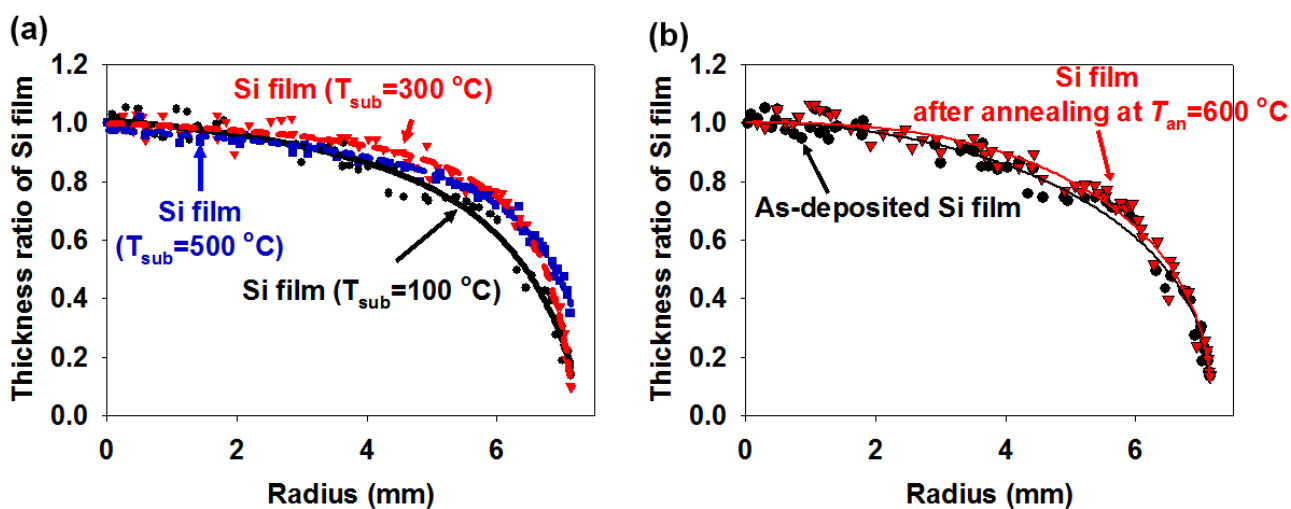


Figure 2-7 Thickness profiles of typical porous Si films deposited on Cu substrates by RVD. (a) As-deposited films at different T_{sub} . (b) As-deposited and post-annealed films with $T_{\text{sub}} = 300$ °C.

Table 2-1 Mass density and porosity of Si films formed at different temperatures

Sample conditions	ρ_{film} (g cm^{-3})	p_{film} (a.u.)
As-deposited at $T_{\text{sub}} = 100$ °C	1.54	0.34
Deposited at $T_{\text{sub}} = 100$ °C and annealed at $T_{\text{an}} = 600$ °C	1.57	0.33
As-deposited at $T_{\text{sub}} = 300$ °C	1.74	0.25
As-deposited at $T_{\text{sub}} = 500$ °C	1.98	0.15

$$p_{\text{film}} \equiv \frac{\rho_{\text{Si}} - \rho_{\text{film}}}{\rho_{\text{Si}}} \quad (1)$$

where ρ_{Si} and ρ_{film} are the mass densities of bulk Si and porous Si film, respectively. The Si film showed the lowest $\rho_{\text{film}} = 1.54 \text{ g cm}^{-3}$ and highest $p_{\text{film}} = 0.34$ when deposited at $T_{\text{sub}} = 100 \text{ }^{\circ}\text{C}$, moderate $\rho_{\text{film}} = 1.74 \text{ g cm}^{-3}$ and $p_{\text{film}} = 0.25$ when deposited at $T_{\text{sub}} = 300 \text{ }^{\circ}\text{C}$, and highest $\rho_{\text{film}} = 1.98 \text{ g cm}^{-3}$ and lowest $p_{\text{film}} = 0.15$ when deposited at $T_{\text{sub}} = 500 \text{ }^{\circ}\text{C}$. In contrast, post-annealing at $T_{\text{an}} = 600 \text{ }^{\circ}\text{C}$, had little influence on the density and porosity of the films.

2.4 Effects of the surface condition of the Cu substrates on the cycle performance

The surface condition of the Cu substrate strongly affects the adhesion between the Si films and Cu substrates, and thus was evaluated by XPS. Only O and C were observed as contaminants in addition to Cu, as shown in Figure 2-8 and Table 2-2. The as-purchased Cu substrate contained a large amount of C (69.9 at%) on its surface, possibly from an oily contaminant used during mechanical processing, which resulted in Si films being easily detached from the Cu substrate in a "Scotch-tape" test. Sonication in isopropanol followed by UV-O₃ treatment reduced the surface C content to 29.8 at%, but increased the O content to 50.9 at%. The Cu 2p peaks became evident with the 2p_{3/2} peak centred

Table 2-2 Elemental compositions of Cu plate surfaces after different treatments

Surface conditions	Cu (2p _{3/2}) (at%)	O (1s) (at%)	C (1s) (at%)
As-purchased	6.5	23.6	69.9
UV-O ₃ treated	19.3	50.9	29.8
UV-O ₃ & H ₂ annealed	63.6	24.6	11.8

at 933.8 eV (Figure 2-8b), which is chemically shifted to higher binding energy from the Cu⁰ position (932.4 eV) because of surface oxidation. The full-width at half maximum (FWHM) of the Cu 2p_{3/2} peak was large (3.84 eV), which was attributed to multiplet splitting caused by Cu²⁺ [101, 102]. The UV-O₃-treated surface improved the adhesion between Cu and as-deposited Si (stable against the Scotch-tape test), but corrosion of CuO occurred at the Cu-Si interface during Li⁺ intercalation [103-105], resulting in detachment of the Si film. H₂ annealing following UV-O₃ exposure resulted in considerable reductions in both C (11.8 at%) and O (24.6 at%) surface contents (Table 2-2). The Cu 2p peaks were more intense, sharper (FWHM = 1.31 eV for Cu 2p_{3/2}), and the binding energy of the Cu 2p_{3/2} peak was centred at 932.4 eV, consistent with the Cu substrate having a metallic surface.

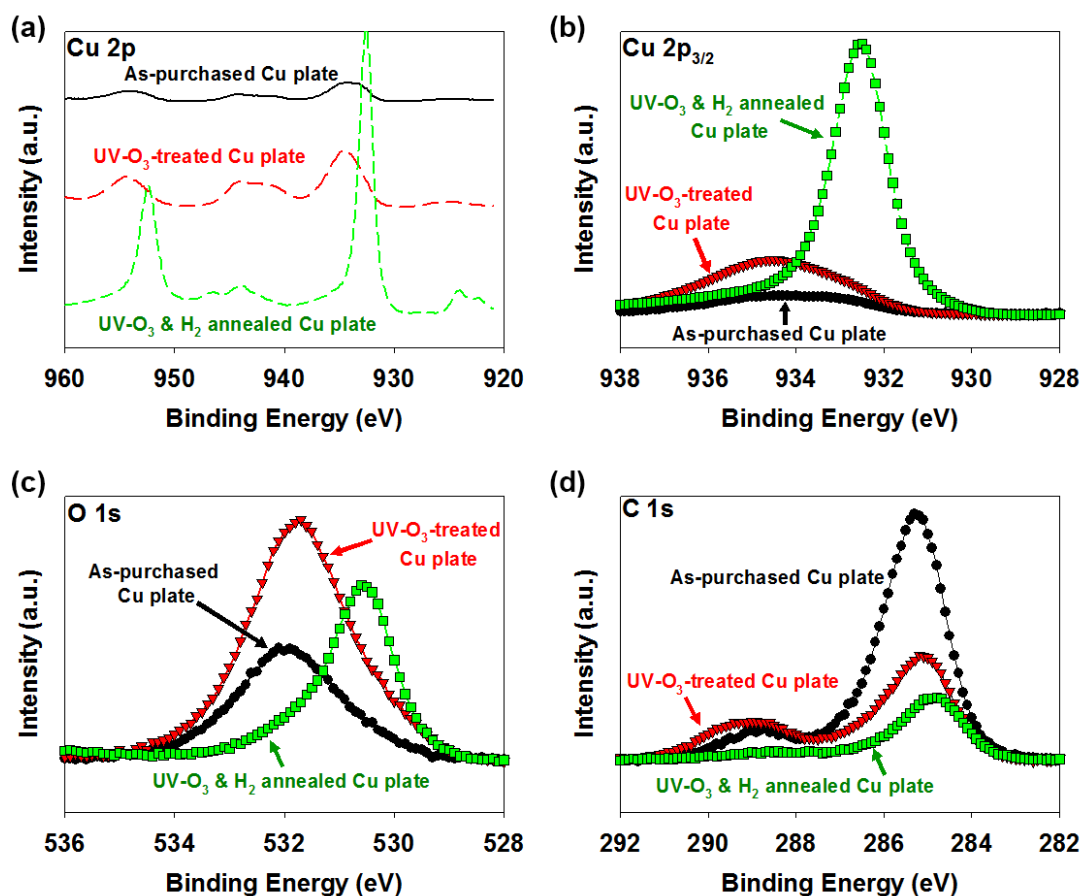


Figure 2-8 XPS analysis of the surfaces of Cu plates after different treatments; as purchased, after UV-O₃ treatment for 3 min, and after UV-O₃ treatment for 3 min followed by annealing under 5 vol% H₂ at 1 atm and 800 °C for 10 min. (a) Cu 2p, (b) Cu 2p_{3/2}, (c) O 1s, and (d) C 1s.

Such a metallic surface allowed good adhesion of the Si films to the Cu substrates without the problem of corrosion at the Cu-Si interface during Li^+ insertion, as shown later. Thus, we used UV- O_3 and H_2 annealing as the standard procedure for treating the Cu substrates in the following experiments.

2.5 Heat treatment to enhance interfacial adhesion while retaining porous structure

The temperatures of the Cu substrate during RVD (T_{sub}) and post-annealing (T_{an}) are the most important factors that determine the crystallinity of deposited Si films as well as the thickness of the intermixed layer between the Si film and Cu substrate. With regard to the crystallinity of Si films, the amorphous phase is preferred because of its isotropic expansion behavior rather than the crystalline phase, which shows anisotropic expansion during Li^+ insertion [106]. Moreover, the diffusion of Li^+ into amorphous Si is much faster than that into crystalline Si [107]. Figure 2-9a shows the Raman spectra of a series of Si films formed at different Cu substrate temperatures. These films possess different degrees of crystallinity from a pure amorphous phase to a mixture of amorphous and microcrystalline phases depending on the temperature of the Cu substrate. The crystal structure of Si is diamond cubic, which is characterized by one intense sharp peak around 520 cm^{-1} in Raman spectra. This peak broadens and/or shifts to lower frequency when the crystal size is \leq several tens of nanometers. $T_{\text{sub}} = 500\text{ }^\circ\text{C}$ resulted in a mixture of microcrystalline and amorphous phases, while lower T_{sub} (100–300 $^\circ\text{C}$) gave the pure amorphous phase with a broad band at around 480 cm^{-1} . Intermixing between Cu and Si is also important to enhance the adhesion between Si films and Cu substrates and suppress the lifting of the Si films from the Cu substrates. In addition to the effect of T_{sub} during Si deposition, we examined the effect of T_{an} on the depth profile of the elemental composition of the films. SEM-EDS analysis showed little intermixing in the as-deposited Si film

prepared at low $T_{\text{sub}} = 100\text{ }^{\circ}\text{C}$, and more intermixing for higher T_{sub} of $300\text{--}500\text{ }^{\circ}\text{C}$ (Figure 2-9b). Intermixing was also enhanced by post-annealing at $T_{\text{an}} = 600\text{ }^{\circ}\text{C}$ the Si films deposited at $T_{\text{sub}} = 300\text{ }^{\circ}\text{C}$. Deposition at high Cu substrate temperature (T_{sub}) increased both the crystallisation of Si films

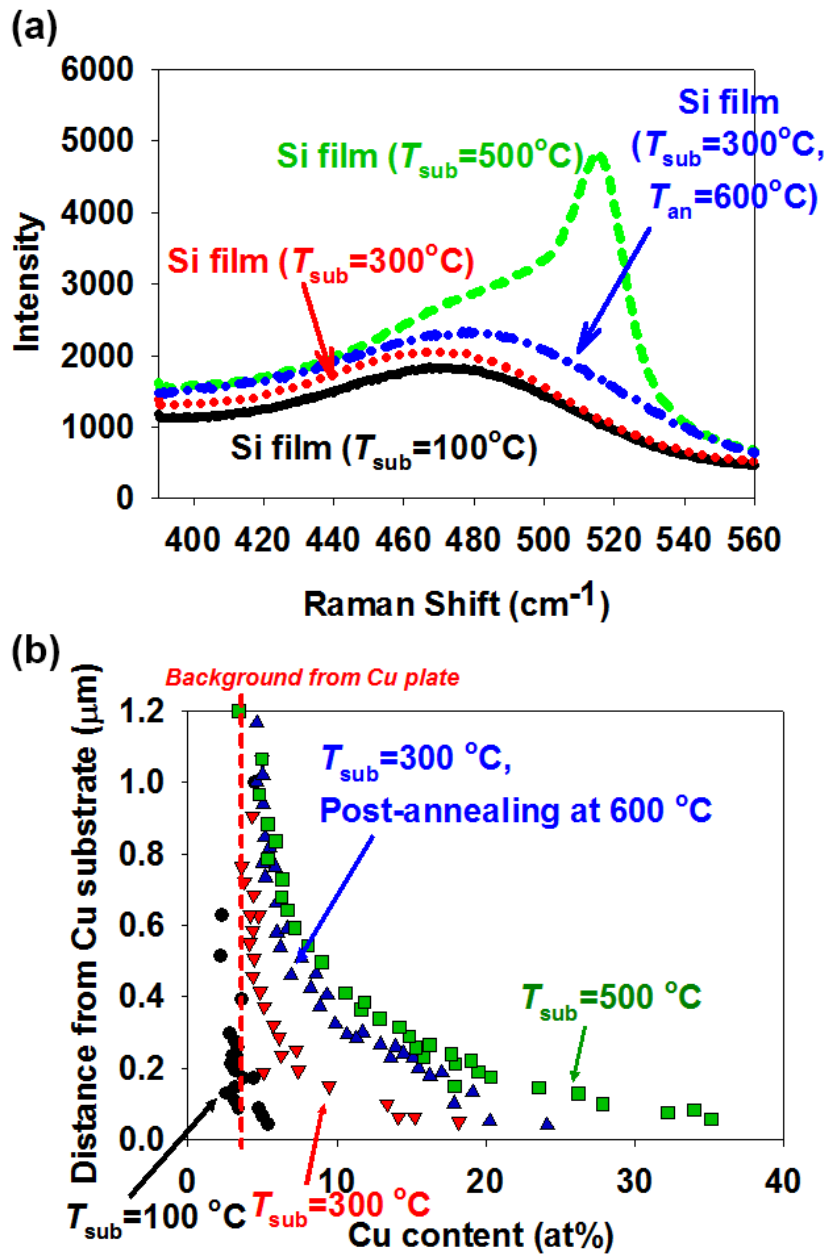


Figure 2-9 (a) Raman spectra of Si films ($t_{\text{eff}} = 3\text{--}4\text{ }\mu\text{m}$) deposited at $T_{\text{sub}} = 100, 300,$ and $500\text{ }^{\circ}\text{C}$ on Cu plates. (b) Depth profiles of the elemental compositions of Si films ($t_{\text{eff}} = 3\text{--}4\text{ }\mu\text{m}$) deposited at $T_{\text{sub}} = 100, 300,$ and $500\text{ }^{\circ}\text{C}$ without post annealing and that deposited at $T_{\text{sub}} = 300\text{ }^{\circ}\text{C}$ with post-annealing at $T_{\text{an}} = 600\text{ }^{\circ}\text{C}$ measured by SEM-EDS. The Cu signal of $\sim 3\text{ at}\%$ is a background signal from the Cu plate for all measurements.

and intermixing between the Si films and Cu substrates while post-annealing at an appropriate temperature ($T_{\text{an}} = 600\text{ }^{\circ}\text{C}$) enhanced intermixing only. Through such temperature control, we can obtain 3–4- μm -thick Si films that contain both amorphous and microcrystalline phases with an intermixed layer that is 10% of the total thickness of the Si films.

2.6 Cycle performance and impedance analysis of porous Si films

The electrochemical behavior of Si films ($t_{\text{eff}} = 3\text{--}4\text{ }\mu\text{m}$) deposited at different T_{sub} on Cu substrates was investigated by galvanostatic charge-discharge measurements using half cells (Figure 2-10). Figure 2-10a shows the initial charge/discharge capacities of the Si films. In the initial lithiation process, the voltage profile of all films coincided with that of previous Si films with a plateau region at a potential below 0.2 V vs. Li/Li⁺, which indicates that amorphous Si reacted with Li⁺ to form amorphous Li_xSi alloy [61]. In the case of lower T_{sub} of 100 and 300 °C, charge capacities reached 2759 and 2471 mAh g_{Si}⁻¹ while discharge capacities were 1151 and 1494 mAh g_{Si}⁻¹, for 0.05C for $T_{\text{sub}} = 100\text{ }^{\circ}\text{C}$ and 0.1C for $T_{\text{sub}} = 300\text{ }^{\circ}\text{C}$, respectively. The large irreversible capacity is mainly attributed to the consumption of Li⁺ in the formation of an SEI layer [108]. In particular, the large exposed surface area of the porous structure formed at low T_{sub} (Figure 2-6b, c) consumed a large amount of Li⁺ during SEI formation, resulting in low first Coulombic efficiencies of 41% ($T_{\text{sub}} = 100\text{ }^{\circ}\text{C}$) and 60% ($T_{\text{sub}} = 300\text{ }^{\circ}\text{C}$). Moreover, the Si films are vulnerable to oxidation, particularly amorphous Si compared with crystalline Si. Oxidation of the Si films during the time (2 days or more) between Si deposition and electrochemical testing should have a considerable effect on cycling performance. In contrast, the Si film deposited on a Cu substrate at $T_{\text{sub}} = 500\text{ }^{\circ}\text{C}$ exhibited the highest initial charging (Li⁺ insertion) capacity of 4045 mAh g_{Si}⁻¹ at a rate of 0.05C with a discharge capacity

of 3328 mAh g_{Si}⁻¹. The charge capacity is nearly equivalent to the theoretical value for Li₂₂Si₅ alloy (4200 mAh g_{Si}⁻¹) [61]. This is because the high T_{sub} yielded dense Si films with reduced surface area (Figure 2-6d) and partial crystallization of the Si film (Figure 2-9a), which can reduce SEI formation

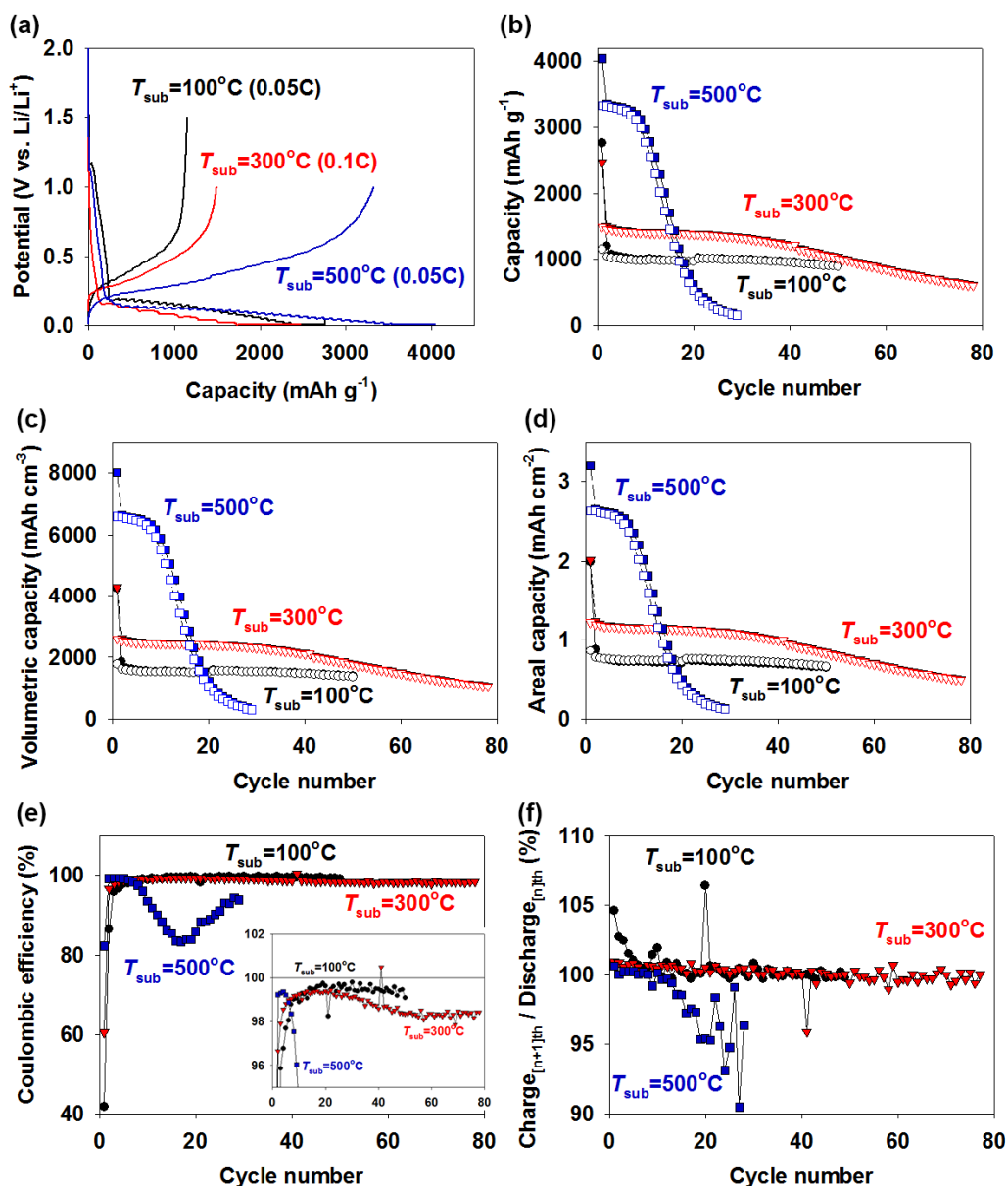


Figure 2-10 Electrochemical performance of Si films deposited at $T_{\text{sub}} = 100, 300, \text{ and } 500\text{ }^{\circ}\text{C}$ on Cu plates ($t_{\text{eff}} = 3.1, 3.5, \text{ and } 3.4\text{ }\mu\text{m}$ respectively). (a) Voltage-capacity curves for the first charge/discharge cycles. Cycle performance at 0.1C expressed as (b) gravimetric capacity, (c) volumetric capacity, and (d) areal capacity. (e) Coulombic efficiency. (f) The capacity ratio of charge at the $[n+1]^{\text{th}}$ cycle to discharge at the $[n]^{\text{th}}$ cycle, which provides information about the reaction of the newly used Si through cycles. The cycle test was made at 0.1C for all cycles (1–80) for $T_{\text{sub}} = 300\text{ }^{\circ}\text{C}$, and at 0.05C for the first cycle and 0.1C for later cycles (≥ 2) for $T_{\text{sub}} = 100 \text{ and } 500\text{ }^{\circ}\text{C}$.

and oxidation of the Si film, compared with the films formed at lower T_{sub} . However, the Si film deposited at $T_{\text{sub}} = 500\text{ }^{\circ}\text{C}$ showed more rapid degradation than those deposited at lower T_{sub} , resulting in a poor capacity retention of 15% after 20 cycles (Figure 2-10b). This is possibly because of the small porosity ($p_{\text{film}} = 0.15$, Table 2-1) and mixed amorphous and microcrystalline phases (Figure 2-9a) in this film, which resulted in large stress upon volume change during cycling. Another possible explanation is that Li^+ reached the Si/Cu interface at around the 10th cycle and corroded the interface for this film having the thickest CuSi_x intermixed layer (Figure 2-9b). The previous works reported the absence at room temperature [109] but existence at $\sim 120\text{ }^{\circ}\text{C}$ [110] of the reaction of Cu_3Si alloy with Li^+ . When we consider that the Si film deposited at $T_{\text{sub}} = 300\text{ }^{\circ}\text{C}$ had a similarly thick Cu_3Si intermixed layer but did not show such abrupt capacity fade, the latter explanation seems less probable. In contrast, the Si film deposited at $T_{\text{sub}} = 300\text{ }^{\circ}\text{C}$ showed much higher capacity retention of 81% with a discharge capacity of $1210\text{ mAh g}_{\text{Si}}^{-1}$ after 40 cycles, which decreased to 40% and $\sim 600\text{ mAh g}_{\text{Si}}^{-1}$, respectively, after 80 cycles.

The Si film deposited at $T_{\text{sub}} = 100\text{ }^{\circ}\text{C}$ showed interesting behavior; it exhibited the smallest discharge capacity of the films of $1151\text{ mAh g}_{\text{Si}}^{-1}$ for the first cycle but a good capacity retention of 70% after 50 cycles. The high porosity of this film of $p_{\text{film}} = 0.34$ (Table 2-1) as well as the thick SEI layer formed in the first cycle should contribute to the stable charge/discharge capacity. The volumetric capacity ($\text{mAh cm}_{\text{film}}^{-3}$) and areal capacity ($\text{mAh cm}_{\text{anode}}^{-2}$) of films are critically important for practical battery devices. Thus, the cycle performance based on gravimetric capacity ($\text{mAh g}_{\text{Si}}^{-1}$) in Figure 2-10b is plotted as volumetric capacity and areal capacity in Figure 2-10c and d, respectively. The volumetric capacity of the Si film formed at $T_{\text{sub}} = 100\text{ }^{\circ}\text{C}$ (Figure 2-10c) was about 1500 mAh cm^{-3} after 50 cycles, which is nearly four times the capacity of commercial graphite anodes [111]. It should be noted that there is still potential to increase the capacity of later cycles by optimising the Si film porosity and operating conditions such as electrolyte additives and cut-off potential. The areal

capacity of the Si film formed at $T_{\text{sub}} = 100\text{ }^{\circ}\text{C}$ (Figure 2-10d) was also higher (0.66 mAh cm^{-2}) than the previous values reported for Si nanomaterials ($0.2\text{--}0.4\text{ mAh cm}^{-2}$) [79, 88] and approaching that of commercial graphite anodes ($\sim 4\text{ mAh cm}^{-2}$, 18650 Li-ion cells, Sony, Japan) [112] and the recent Si-base hybrid materials (2 mAh cm^{-2}) [113].

The films deposited at $T_{\text{sub}} = 100$ and $300\text{ }^{\circ}\text{C}$ also showed rather good Coulombic efficiency (Figure 2-10e). After the 5th cycle, Coulombic efficiency was above 98%, and remained at $\sim 99.5\%$ for 50 cycles for $T_{\text{sub}} = 100\text{ }^{\circ}\text{C}$ and 98–99.5% for 80 cycles for $T_{\text{sub}} = 300\text{ }^{\circ}\text{C}$. The capacity ratio of the $[n+1]$ th charge over the $[n]$ th discharge for the films is plotted in Figure 2-10f. The ratio was above 100% for the initial 30 cycles, suggesting that the Si anode did not fully react initially and unreacted Si gradually contributed to the reaction with increasing cycle number, compensating for the decrease in discharge capacity (99% Coulombic efficiency corresponds to 1% loss in effective Si for each cycle) and maintaining the capacity retention. The quick capacity fade observed for the Si film with $T_{\text{sub}} = 300\text{ }^{\circ}\text{C}$ after the 30th cycle (Figure 2-10b) suggests the depletion of the unreacted Si. The Si film deposited at $T_{\text{sub}} = 300\text{ }^{\circ}\text{C}$ and post-annealed at $T_{\text{an}} = 600\text{ }^{\circ}\text{C}$ ($t_{\text{eff}} = 3.5\text{ }\mu\text{m}$) was investigated by EIS after charging for different numbers of cycles. In the Nyquist plot (Figure 2-11), a pronounced

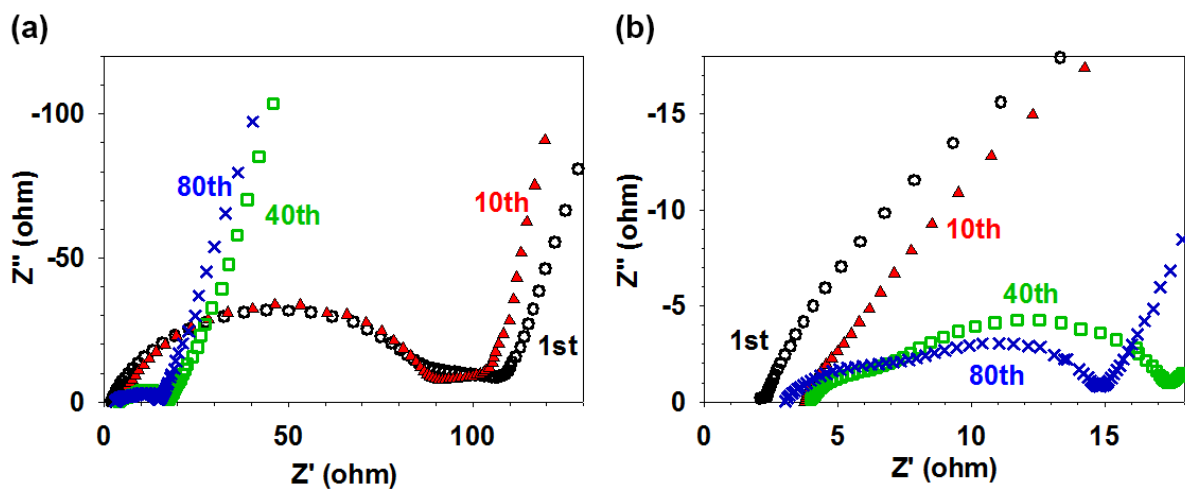


Figure 2-11 Nyquist plots of a thick Si film anode ($t_{\text{eff}} = 3.5\text{ }\mu\text{m}$) deposited at $T_{\text{sub}} = 300\text{ }^{\circ}\text{C}$ and post-annealed at $T_{\text{an}} = 600\text{ }^{\circ}\text{C}$ after charging for different numbers of cycles.

semicircle appeared at the 1st cycle and this semicircle did not change for the 10th cycle. We attribute this to the large reaction resistance for the unreacted Si remaining in the charged Si film, which was evidenced by the capacity ratio of the $[n+1]$ th charge over the $[n]$ th discharge exceeding 100% for the initial 30 cycles (Figure 2-10f). This semicircle got much smaller after the 40th cycle (Figure 2-11), which we attribute to the depletion of the unreacted Si in the charged Si film, which was observed in the capacity ratio of the $[n+1]$ th charge over the $[n]$ th discharge of less than 100% for >30 cycles

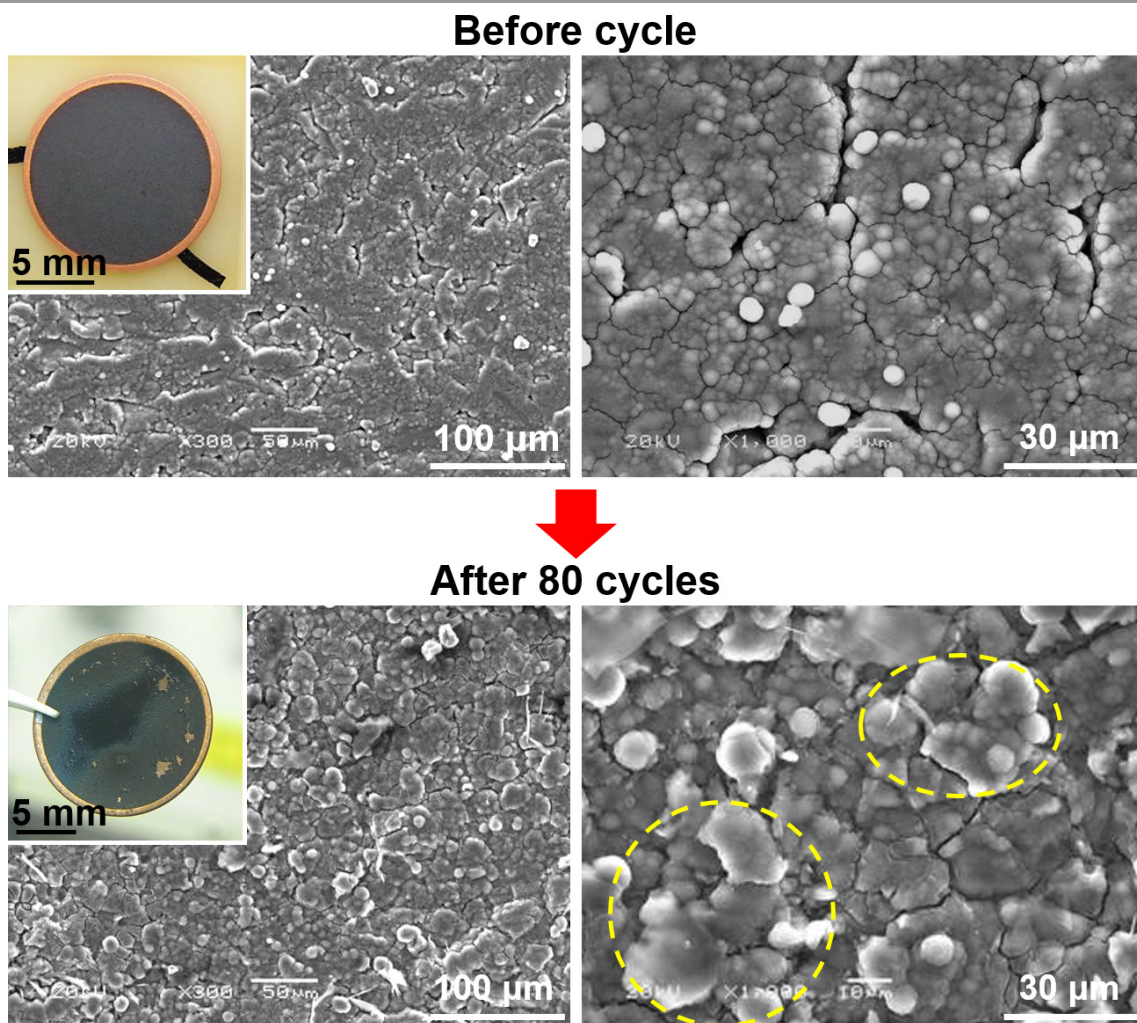


Figure 2-12 Photographs and SEM images of the Si film deposited at $T_{\text{sub}} = 300 \text{ }^\circ\text{C}$ and post-annealed at $T_{\text{an}} = 600 \text{ }^\circ\text{C}$ before and after the charge-discharge cycle. The sample is the same as that in Fig. 7. Before the cycle, the Si film was uniform over the Cu substrate and show porous surface structure. While it was detached from the substrate at some regions to show the Cu surface in the photograph and had several-tens- μm large protrusions in the SEM images after 80 cycles.

(Figure 2-10f). It is reasonable to consider that the reaction resistance became much smaller once the porous Si film was lithiated [114]. We also see a slight increase of contact resistance between the 1st and 10th cycles (Figure 2-11). We attribute this to some detachment of the Si film from the Cu substrate despite the improved Si/Cu interface with a composition gradient fabricated by thermal treatment at $T_{\text{sub}} = 300\text{ }^{\circ}\text{C}$ and post-annealing at $T_{\text{an}} = 600\text{ }^{\circ}\text{C}$. We also characterized the same Si film

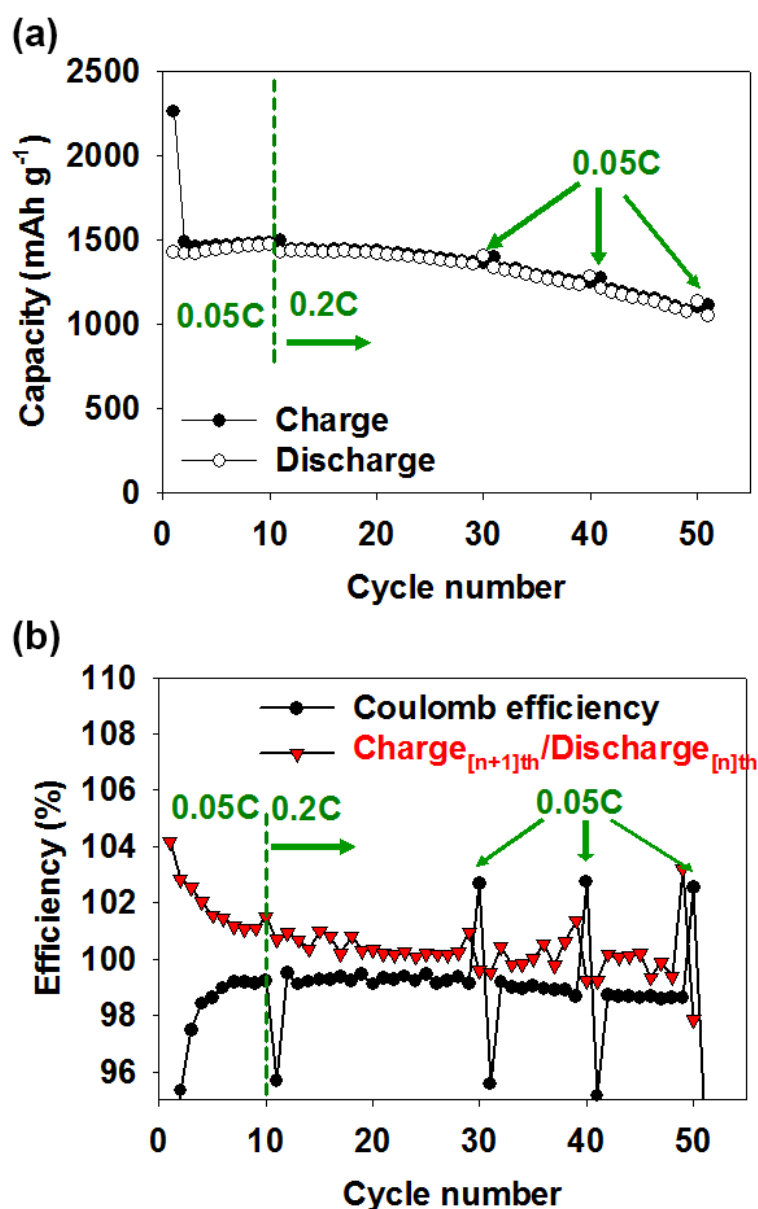


Figure 2-13 (a) Rate capability of a thick Si film ($t_{\text{eff}} = 3.1\text{ }\mu\text{m}$) deposited on a Cu plate at $T_{\text{sub}} = 300\text{ }^{\circ}\text{C}$ and post-annealed at $T_{\text{an}} = 600\text{ }^{\circ}\text{C}$ for 50 cycles. (b) Coulombic efficiency and the capacity ratio of charge at the $(n+1)^{\text{th}}$ cycle to discharge at the $(n)^{\text{th}}$ cycle as a function of cycle number n of the same film as in (a).

before and after the cycles. The photographs and SEM images (Figure 2-12) shows the uniform Si film over the Cu substrate with porous surface structure before the cycle. On the other hand, the Si film was detached from the substrate at some regions to show the Cu surface in the photograph and had several-tens- μm large protrusions in the SEM images after 80 cycles. Some countermeasure such as using nano-/micro-structured substrate would be needed to enhance the interfacial adhesion further.

The rate capability of a representative 3.1- μm -thick Si anode deposited at $T_{\text{sub}} = 300\text{ }^{\circ}\text{C}$ and post-annealed at $T_{\text{an}} = 600\text{ }^{\circ}\text{C}$ is shown in Figure 2-13. Starting from 0.05C at the 1st cycle, the Si film achieved a discharge capacity of 1426 $\text{mAh g}_{\text{Si}}^{-1}$, which increased to 1472 $\text{mAh g}_{\text{Si}}^{-1}$ after 10 cycles. This increase is because unreacted Si gradually reacted with Li^{+} and contributed to the discharge capacity as can be seen in the $\text{charge}_{[n+1]\text{th}}/\text{discharge}_{[n]\text{th}}$ capacity ratio of $> 100\%$ with smaller loss of reacted Si, which is evidenced by the Coulombic efficiency approaching 100% (Figure 2-13b). When the current density was quadrupled to 0.2C from the 11th cycle, Coulombic efficiency dropped from above 99% to 95%, but recovered immediately at the 12th cycle. Moreover, the discharge capacity showed only a slight decrease to 1428 $\text{mAh g}_{\text{Si}}^{-1}$ at 0.2C. After the 20th cycle, both the charge and discharge capacity of the film started to decrease gradually. When the charge/discharge rate were set back to 0.05C at the 30th, 40th and 50th cycles, both the charge and discharge capacity increased slightly by a few tens of $\text{mAh g}_{\text{Si}}^{-1}$. This small capacity difference of a few percent confirms the sufficient rate performance of the Si anode for 0.05C–0.2C. At the 50th cycle, a reversible capacity of 1050 $\text{mAh g}_{\text{Si}}^{-1}$ was achieved at 0.05C. The Coulombic efficiency was above 98% for cycle 5–50 regardless of the current density (0.05C or 0.2C).

2.7 Conclusions

We developed the RVD method, in which a Si source is heated to well above its melting point (T_{boat}

= 2000–2400 °C) while a Cu substrate is kept at much lower temperature ($T_{\text{sub}} = 100\text{--}500$ °C), and rapidly deposited 3–14- μm -thick porous Si films directly on Cu substrates in 10 s to 1 min. Such deposition is several orders of magnitude faster than the conventional physical vapour deposition methods using Si source near its melting point (6–90 nm min⁻¹ by thermal evaporation) or the sputtering (8.6 nm and 2.8–6.1 nm min⁻¹ by radio frequency magnetron sputtering). And such fast deposition eliminates the need for ultra-high vacuum systems because the contaminant oxygen is diluted by the rapidly depositing Si. Compared with chemical vapour deposition methods, RVD uses the safe source instead of the explosive/toxic silane/chlorosilane sources, and moreover the Si vapour enables deposition at low temperatures, leading to the spontaneous roughening of the Si film. The Si films had striped, porous structure, with different porosity ($p_{\text{film}} = 0.15\text{--}0.34$) and ratio of amorphous to microcrystalline phases depending on T_{sub} . Rapid deposition of 14 μm of Si in 10 s resulted in protrusions in the films, while moderately rapid deposition of 3–4 μm in 1 min gave films without protrusions. Pretreatment of the Cu substrates by sonication in isopropanol followed by UV-O₃ treatment and finally annealing in H₂/Ar at 800 °C was effective to remove C and O contaminants from their surfaces. The as-deposited porous Si films showed very poor charge-discharge cycle performance that was improved considerably once the films were annealed at 600 °C for 10 min because a sub-micrometer-thick CuSi_x intermixed layer formed at the Si/Cu interface without the Si films crystallizing. The rather dense ($p_{\text{film}} = 0.15$), thick Si films ($t_{\text{eff}} = 3.4$ μm) of mixed amorphous-microcrystalline phase deposited at $T_{\text{sub}} = 500$ °C showed fairly high initial charge/discharge capacities of 4045 and 3328 mAh g_{Si}⁻¹ at 0.05C, respectively, and kept a high capacity of >3000 mAh g_{Si}⁻¹ for the first 10 cycles but their capacity decreased to below 500 mAh g_{Si}⁻¹ after 20 cycles. The dense structure of these films possibly suppressed their oxidation in air and excess SEI layer formation, resulting in good initial performance but large stress caused by volume changes during cycling, resulting in the rapid capacity fade within 20 cycles. In contrast, the low-density ($p_{\text{film}} = 0.33$) thick amorphous Si films ($t_{\text{eff}} = 3.1$ μm) deposited at $T_{\text{sub}} = 100$ °C showed a rather small discharge

capacity of 1151 mAh g_{Si}⁻¹ for the first cycle but good capacity retention of 70% after 50 cycles. The porous structure of these films possibly facilitated their oxidation in air and excess SEI layer formation, resulting in a small initial discharge capacity but suppressed the stress caused by volume changes during cycling and yielded a stable SEI layer, resulting in rather good cycle stability. The reversible capacity of ~1000 mAh g_{Si}⁻¹ after 50 cycles of the Si film deposited at low T_{sub} (100 °C) corresponds to a high volumetric capacity of ~1500 mAh cm_{film}⁻³ and areal capacity of ~0.5 mAh cm_{anode}⁻², which suggest Si anodes may be suited for practical use. Toward the future goal having porous Si films of $t_{\text{eff}} \sim 10 \mu\text{m}$ on both sides of a 15- μm -thick Cu foil and operating it for longer cycles of ~1,000 or more, we are now examining the charge limitation to reduce the volume expansion and additives to electrolyte to make more stable SEI. Although further improvement is needed in thickness and cycle performance, the RVD method yielding micrometer-thick Si films rapidly from an inexpensive, safe Si source, and that allows control over porosity, crystallinity, and interface with the Cu collector, and that can be applied to various substrates including nano-/micro-structured substrates, is a promising route to fabricate practical Si anodes for LiBs.

Chapter 3 – Rapidly deposited, porous Si–Cu anodes with compositional gradients

3.1 Introduction

For the efficient use of electrical energy and particularly renewable energy, electrical energy storage systems are very important. Rechargeable lithium ion batteries have been used successfully in portable electronic device and electric vehicle applications, but there is considerable demand for further enhancements of the energy and power densities of these batteries, along with the need for cost reduction. Si is considered to be a promising anode material for lithium secondary batteries [65]. Si can accommodate up to 4.4 lithium atoms per Si atom ($\text{Li}_{4.4}\text{Si}$ alloy), thus yielding huge theoretical gravimetric and volumetric capacities of $4200 \text{ mAh g}_{\text{Si}}^{-1}$ and $9786 \text{ mAh cm}_{\text{Si}}^{-3}$, respectively, which are approximately ten times larger than the corresponding values for conventional graphite [61-64]. However, this high specific capacity is realized by holding large amounts of Li with Si, and is accompanied by large volume changes (300–400%, depending on the state of Li_xSi [66-68]). Si anodes cannot withstand the heavy strain of this expansion and thus break down, resulting in the pulverization and delamination of entire structures [69-71]. Various structures, including porous thin films [85, 86], hollow-structured double-walled Si nanotubes [80] and composites of Si with carbon [82, 115], carbon fibres [60], and silicides [83], have previously been reported that can accommodate the volume changes. As a result, the high gravimetric capacities of Si-base anode materials were achieved successfully, but sometimes with small Si loads compared with the heavy current collectors,

and also required complicated and time-consuming processing using expensive raw materials. Large-scale and low-cost fabrication of Si anodes with stable electrochemical performance is essential for practical use of these anodes in lithium secondary batteries.

We recently reported a rapid vapour deposition (RVD) method that enables simple fabrication of 3–14- μm -thick porous Si films rapidly in 10–60 s that are deposited directly on the Cu current collectors using inexpensive Si powder sources [116]. Rapid vapour deposition is a physical vapour deposition method in which a source material is heated to much higher temperatures than the melting point of the source (2000–2400 °C for the case of Si), leading to deposition rates that are several orders of magnitude higher than conventional physical vapour deposition methods [85]. Post-annealing at 600 °C enhanced the adhesion of the Si films to the Cu current collectors by yielding a sub- μm -thick intermixed layer, and produced anode capacitances of 1000 mAh $\text{g}_{\text{Si}}^{-1}$ and 0.66 mAh $\text{cm}_{\text{anode}}^{-2}$ for the 50th cycle of a moderately thick Si film. The film had an effective thickness of 3–4 μm (t_{eff} , calculated by dividing the areal Si mass by the bulk Si density of $\rho_{\text{Si}} = 2.33 \text{ g cm}^{-3}$) and porosity of 15–30%. However, delamination failures ultimately occurred because of the high stress at the Si/Cu interface. Strain-graded C-Al-Si multilayers (C, Al and Si experience volume changes of 10, 100 and 400%, respectively, upon lithiation) produced by sputter deposition [94] and Si-base nanowall arrays fabricated by oblique angle deposition [117] are effective approaches for stress relaxation during lithiation/delithiation. Furthermore, well-aligned CuSi nanorod arrays with gradient profiles in Cu and Si compositions were realized by co-depositing Cu and Si with independent dynamic tuning of their deposition rates at 0–0.5 nm s^{-1} (0–0.03 $\mu\text{m min}^{-1}$) at an oblique angle of 88° with respect to substrate normal although their electrochemical performance was not reported [118]. These processes are attractive in engineering the Si anode structure, however, require high vacuums and long processing times.

Here we propose porous and amorphous Si–Cu films with gradient composition profiles that change

continuously from Cu to Si as ideal structures for defocusing stress during lithiation/delithiation processes. This type of structure corresponds to a strain-graded multilayer with an infinite number of layers, and can be fabricated easily and spontaneously by RVD in 1 min. Si and Cu are co-deposited on the Cu current collector by heating Si and Cu powders together in a carbon boat to ~ 2000 °C. Cu has a lower melting point (1085 °C) and a higher vapour pressure than Si (melting point of 1414 °C), and vaporizes and is deposited preferentially at the initial stage, whereas Si vaporizes and is deposited preferentially at the later stage. Also, the porous and amorphous structures of the films are realized by keeping the Cu current collector at a low temperature of ~ 100 °C. I compared the microstructures and electrochemical performances of the Si–Cu films with pure Si films and discussed the effects of the composition gradient and diffuse interface in the Si–Cu films in enhancing the cycle performances.

3.2 Materials and methods

3.2.1 Si–Cu film fabrication

Circular Cu plates (15-mm diameter, 0.5-mm thickness) were used as substrates for the Si–Cu films. The Cu substrates were pretreated by bath sonication in isopropanol for 10 min followed by UV-O₃ exposure for 3 min to remove organic contaminants from their surfaces. The Cu substrates were then annealed under a 4 vol% H₂/Ar flow at ambient pressure and 800 °C for 10 min to remove any organic contaminants and reduce the Cu surface [116]. The Si wafer (p-type, resistivity of 10–20 Ω cm) was immersed in 5 wt% hydrofluoric acid for 1 min and then ground into powder using a mortar and pestle. The Cu powder was used as purchased (~ 75 - μ m diameter, 99.9%, Wako Pure Chemicals, Osaka, Japan). These Si and Cu powders were loaded together (with weight ratios of 100:0, 95:5, 90:10, and 80:20) in a carbon boat, heated in the resistively heated carbon boat to ~ 2000 °C (boat temperature, T_{boat}) in a 0.1-Torr Ar atmosphere, and vaporized rapidly in 1 min. A Cu substrate was

held ~40 mm from the boat and was kept at 100–500 °C (substrate temperature, T_{sub} , 100 °C in most experiments) during deposition to suppress the surface diffusion of the Si and Cu adatoms and yield rough and porous films through the shadowing effect [98, 99]. The Cu substrate was maintained at constant temperature under the strong thermal radiation from the carbon boat by using a substrate holder made of a block of Cu with a large heat capacity that contained an embedded ceramic heater and N₂-gas cooling line [116]. The co-deposited Si–Cu films were post-annealed under a 4 vol% H₂/Ar flow at ambient pressure at 600 °C (annealing temperature, T_{an}) for 10 min to improve the adhesion between the Cu substrate and the Si–Cu film. The areal weight of the Si–Cu film for electrochemical measurements was in the 0.7–0.8 mg cm⁻² range, corresponding to $t_{\text{eff}} = 3\text{--}3.5 \mu\text{m}$. Note that the Si content of the Si–Cu films is less than 100% and thus the actual Si content in the films is lower than that in bulk Si films with the same t_{eff} .

3.2.2 Characterization

The microstructures and the elemental composition distributions of the Si–Cu films were characterized by scanning electron microscopy (SEM; Hitachi S-4800, Tokyo, Japan) equipped with energy-dispersive X-ray spectroscopy (EDX; EDAX Genesis, AMETEK, Elancourt, France). The specific surface area and pore volume of the Si–Cu films were obtained by Brunauer-Emmett-Teller (BET) analysis of nitrogen adsorption isotherms measured at 77 K (Quantachrome Instruments Autosorb 1C, Boynton Beach, FL, USA). The crystal structures of the Si–Cu films were characterized by X-ray diffraction (XRD; Rigaku RINT Ultima III, Akishima, Japan) using Cu K α radiation ($\lambda = 1.54 \text{ \AA}$). Following the charge–discharge cycles, some samples were also characterized after delithiation and rinsing with dimethyl carbonate (DMC).

3.2.3 Electrochemical characterization

Electrochemical measurements were performed using three-electrode beaker cells. Either a Si film or a Si–Cu film on a Cu substrate as a working electrode, Li metal (15 mm × 30 mm) as the counter electrode, and a Li reference electrode was set up in an Ar glove box with a dew point that was lower than -90 °C. Lithium perchlorate (LiClO_4 , 1 M) in a 1:1 (v/v) mixture of ethylene carbonate (EC) and propylene carbonate (PC) with H_2O content of less than 20 ppm (Kishida Chemical, Osaka, Japan) was used as the electrolyte. Charge/discharge measurements were carried out in a potential range of 0.005–1.200 V vs. Li/Li^+ in constant-current (CC) mode with different rates of 0.1C–1C using a battery charge/discharge system (Hokuto denko HJ1020mSD8, Tokyo, Japan). The C-rate was determined using the sum weight of Si and Cu and a value of $4200 \text{ mAh g}_{\text{film}}^{-1}$ for the theoretical capacity regardless of the Cu content. For the Si–Cu films, the Cu contributes to the weight but not to the capacity, making the theoretical capacity smaller than this value, and thus the actual C-rate should be higher. In this work, I define lithiation as “charge” and delithiation as “discharge”. Three-electrode cells were also used to perform electrochemical impedance spectroscopy with a potentiostat (Bio-Logic VMP3, Claix, France).

3.3 Microstructure and composition profile of Si–Cu films

Figure 3-1 shows a typical Si–Cu film with actual thickness of $2.7 \mu\text{m}$ (t_{act}). The film had a wall-shaped structure that was induced by stripe patterns on the Cu substrate surface (Figure 3-1a), which was quite similar to the pure Si films in our previous report [116]. The porous structure is visible in the high-resolution cross-sectional images (Figure 3-1b and c), and is induced by suppressed surface diffusion of the deposited Si and Cu at the low T_{sub} of 100 °C. The Si–Cu films deposited at higher T_{sub} are also shown in Figure 3-1d and e. The film got denser for the higher T_{sub} . Among the Si–Cu

films deposited at $T_{\text{sub}} = 100, 300,$ and $500\text{ }^{\circ}\text{C}$, those deposited at $100\text{ }^{\circ}\text{C}$ showed the best electrochemical performance (Figure 3-2), and thus T_{sub} was fixed at $100\text{ }^{\circ}\text{C}$ in the following

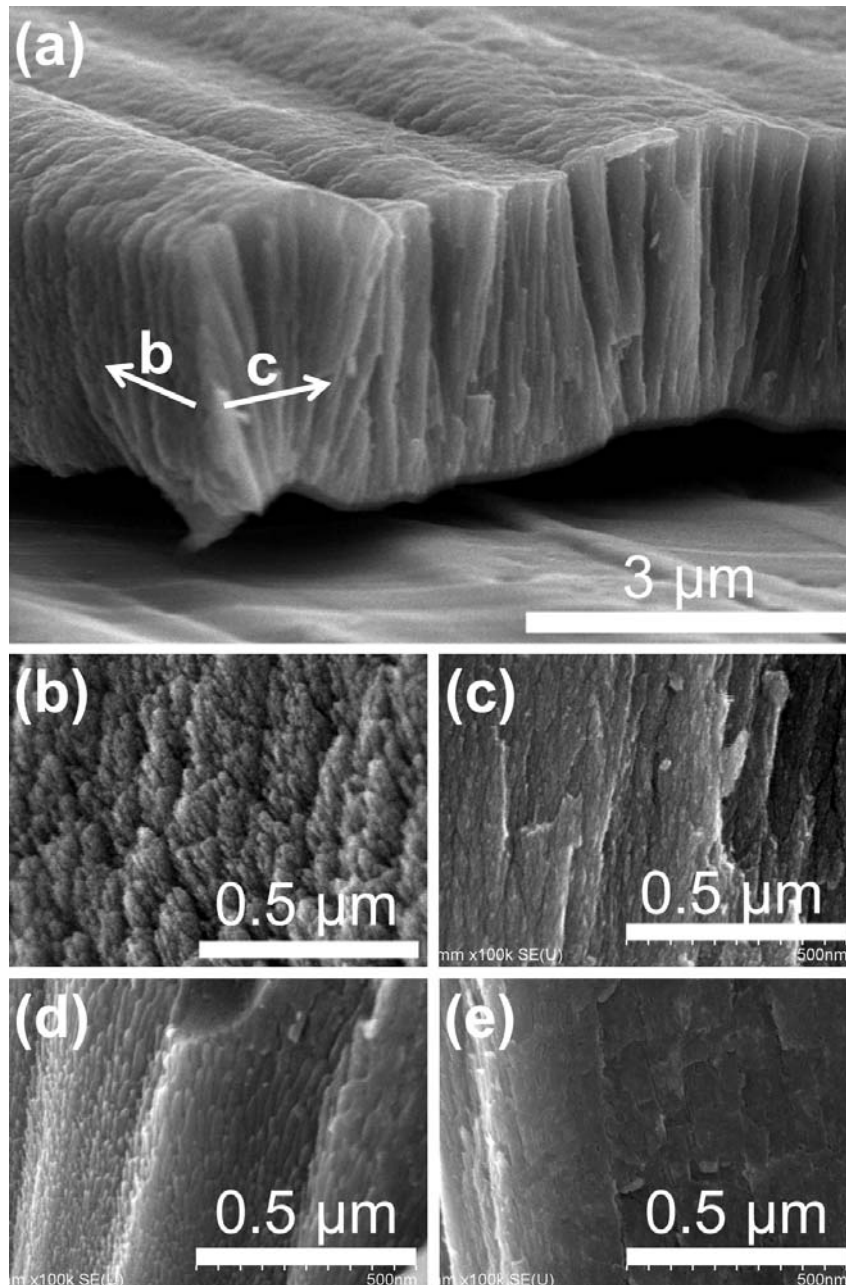


Figure 3-1 SEM images of typical Si–Cu films prepared by RVD using a 5 wt% Cu source in 1 min without post-annealing. The Si–Cu film deposited at $T_{\text{sub}} = 100\text{ }^{\circ}\text{C}$ ($t_{\text{act}} \approx 2.7\text{ }\mu\text{m}$, $\rho_{\text{film}} \approx 1.54\text{ g cm}^{-3}$); (a) the tilted view, (b) the cross-sectional view in the longitudinal direction, and (c) the cross-sectional view in the transverse direction of the stripe patterns on the Cu substrate surface. The Si–Cu films deposited at $T_{\text{sub}} = 300\text{ }^{\circ}\text{C}$ (d) ($t_{\text{act}} \approx 2.8\text{ }\mu\text{m}$, $\rho_{\text{film}} \approx 1.87\text{ g cm}^{-3}$) and at $T_{\text{sub}} = 500\text{ }^{\circ}\text{C}$ (e) ($t_{\text{act}} \approx 2.6\text{ }\mu\text{m}$, $\rho_{\text{film}} \approx 2.01\text{ g cm}^{-3}$)

experiments. The cross-sectional images were acquired by bending the 0.5-mm-thick Cu substrate, which resulted in detachment of the Si–Cu film. Post-annealing at $T_{\text{an}} = 600\text{ }^{\circ}\text{C}$ for 10 min was thus performed on the films that were used for electrochemical measurements to enhance the adhesion of these films to the Cu substrates.

Cross-sectional SEM images of the pure Si and Si–Cu films that were deposited using Si sources with various Cu contents are shown in Figure 3-3a–d. The films that were deposited in 1 min had t_{act} of 2.3–6.7 μm , showing that such rapid deposition is possible, regardless of the Cu content of the source. The pure Si film showed no contrast in its cross-section from bottom to top (Figure 3-3a), while the Si–Cu film showed a clear change in brightness from bottom to top (Figure 3-3b–d). This change in

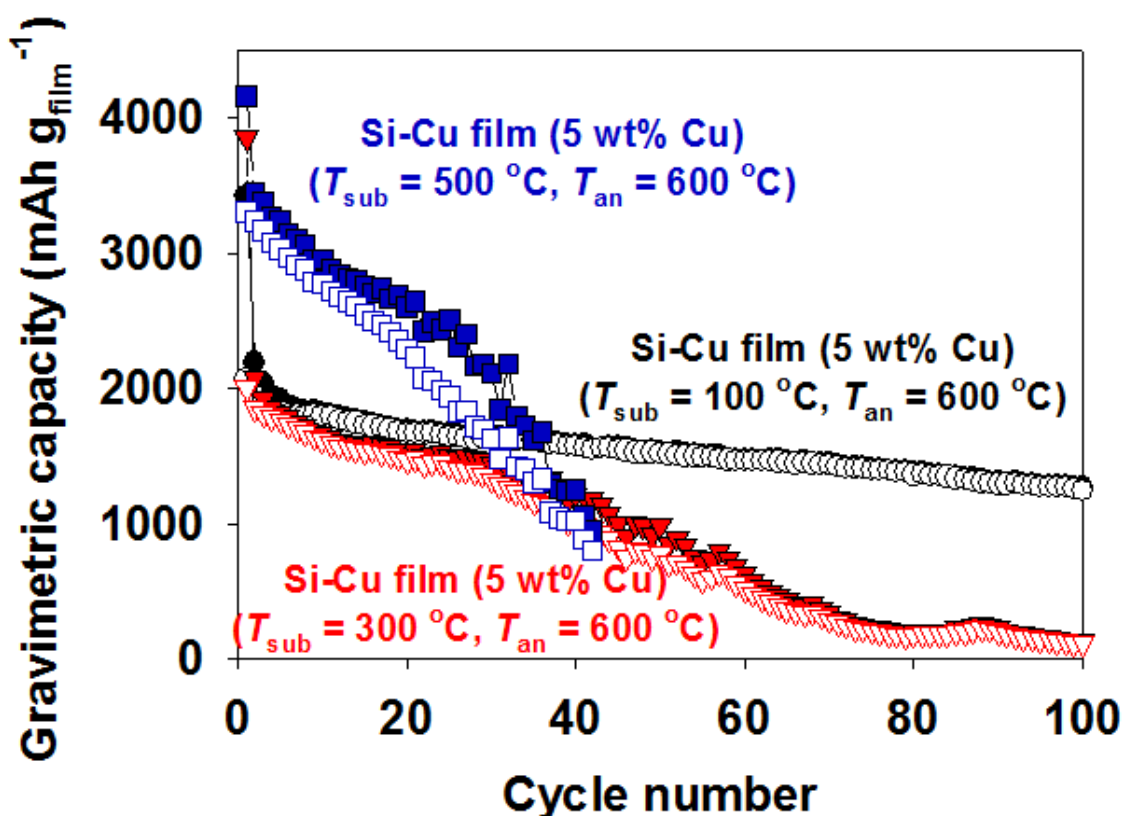


Figure 3-2 Electrochemical performance of the Si–Cu films deposited using the 5 wt% Cu source at various T_{sub} of 100, 300, and 500 $^{\circ}\text{C}$. All samples were post-annealed at $T_{\text{an}} = 600\text{ }^{\circ}\text{C}$ and measurements were carried out using 1 M LiClO₄ in a 1:1 (v/v) mixture of EC and PC as the electrolyte

brightness indicates the change in the Cu content of the film. I also evaluated the mass densities of the Si and Si–Cu films by measuring the masses and volumes of these films [116]. The pure Si film

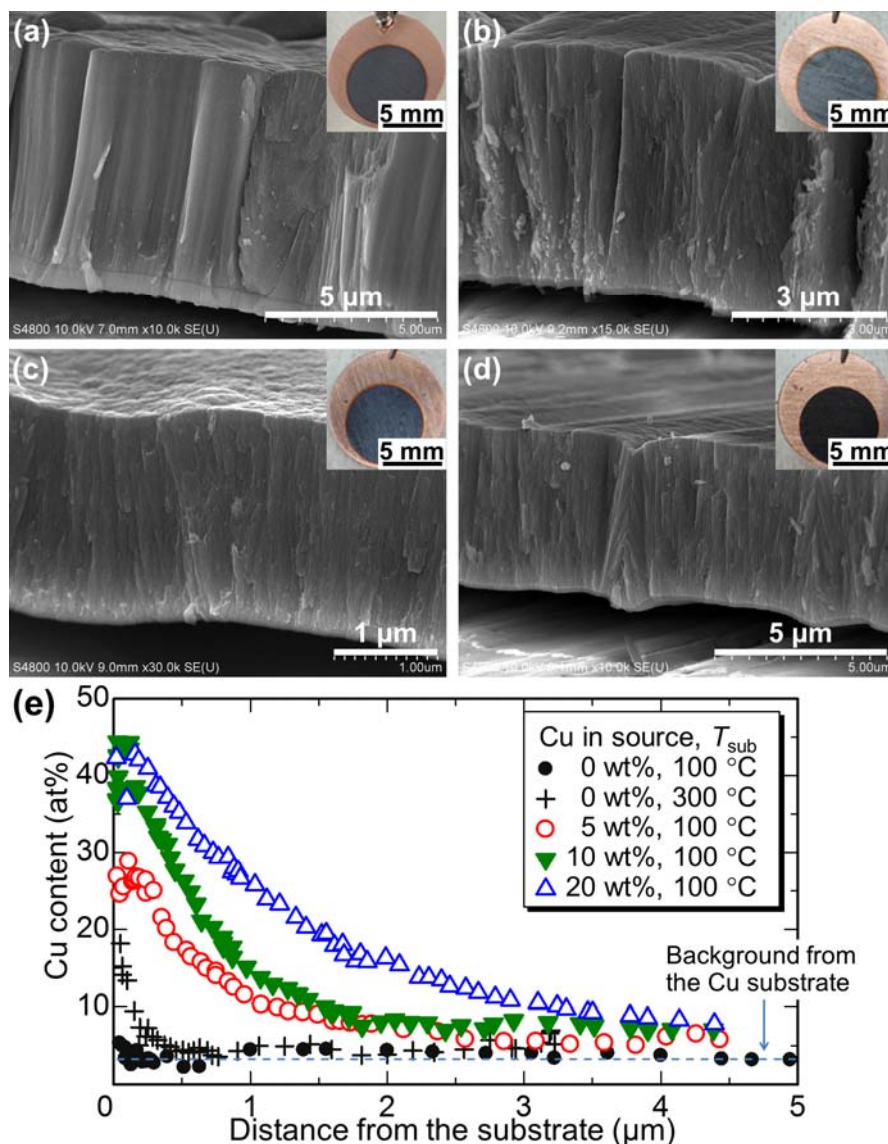


Figure 3-3 Cross-sectional SEM images and photographs (inset) of (a) a Si film ($t_{\text{act}} \approx 6.7 \mu\text{m}$ at the centre of the film, $\rho_{\text{film}} \approx 1.54 \text{ g cm}^{-3}$) prepared with a pure Si source, and of Si–Cu films prepared using Si sources containing (b) 5 wt% Cu ($t_{\text{act}} \approx 4.9 \mu\text{m}$, $\rho_{\text{film}} \approx 1.60 \text{ g cm}^{-3}$), (c) 10 wt% Cu ($t_{\text{act}} \approx 2.3 \mu\text{m}$, $\rho_{\text{film}} \approx 1.67 \text{ g cm}^{-3}$), and (d) 20 wt% Cu ($t_{\text{act}} \approx 4.5 \mu\text{m}$, $\rho_{\text{film}} \approx 1.82 \text{ g cm}^{-3}$). All samples were prepared at $T_{\text{sub}} = 100 \text{ }^\circ\text{C}$ without post-annealing. (e) Depth profiles of the elemental compositions of the pure Si films and the Si–Cu films as measured by SEM-EDX. The standard deviation was 1–3% of the intensity (Cu content value) and 0.2–0.4 at% as the Cu content value (see Figure 3-4). These films showed detachment from the Cu substrate upon bending because post-annealing treatments were not applied to the samples. The ~ 3 at% Cu signal is a background signal from the Cu plate for all measurements (see Figure 3-5).

had a mass density (ρ_{film}) of 1.54 g cm^{-3} , and its porosity (P_{film}) was calculated to be 0.33 using the following equation: $P_{\text{film}} = 1 - \rho_{\text{film}}/\rho_{\text{Si}}$. The mass densities of the films increased from 1.60, to 1.67, to 1.82 g cm^{-3} with increasing Cu content (5, 10, and 20 wt%, respectively) in the Si source. Using the mass density of bulk Si ($\rho_{\text{Si}} = 2.33 \text{ g cm}^{-3}$), the respective porosities of these films were calculated to be 0.31, 0.28, and 0.21; however, their actual porosities should be higher because the bulk Si–Cu alloys have increasing mass densities with increasing Cu content.

Elemental composition of these films were analysed by SEM-EDX at ~ 50 points for each sample as shown in Figure 3-4 and summarized in a graph of the composition profiles (Figure 3-3e). The graph

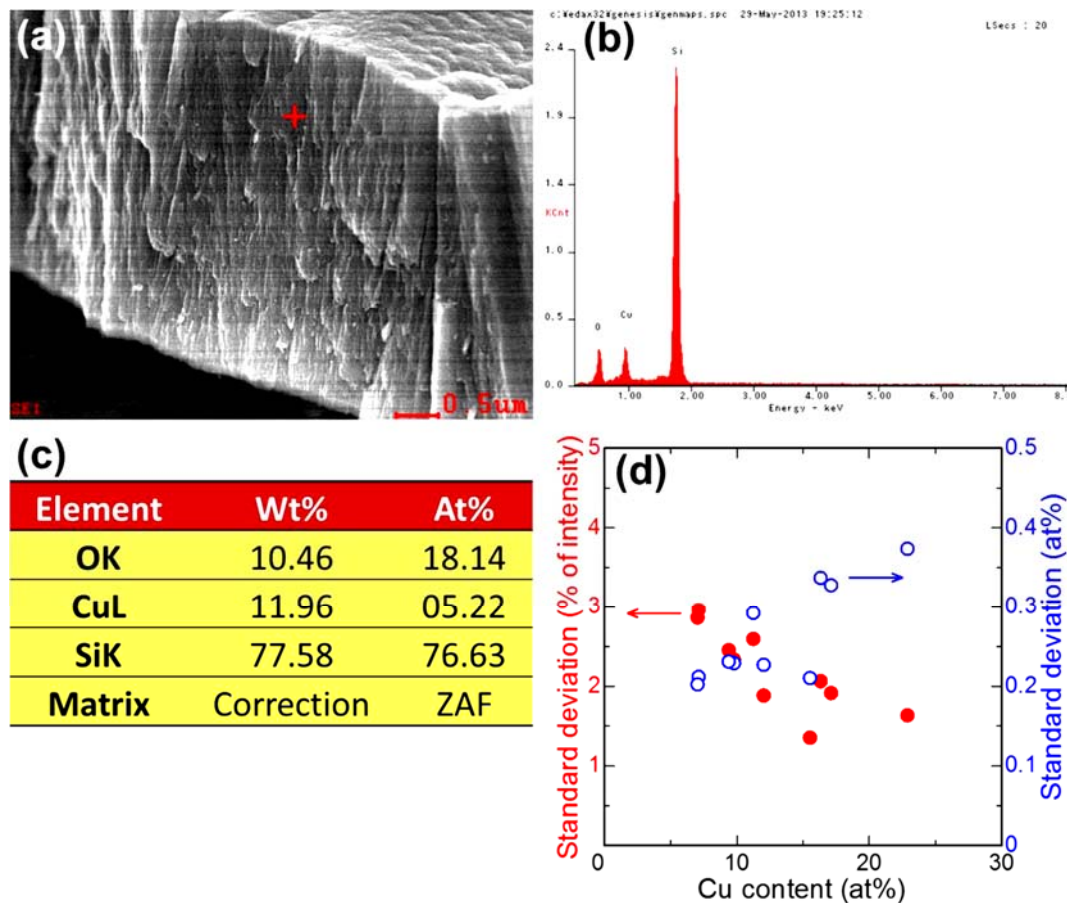
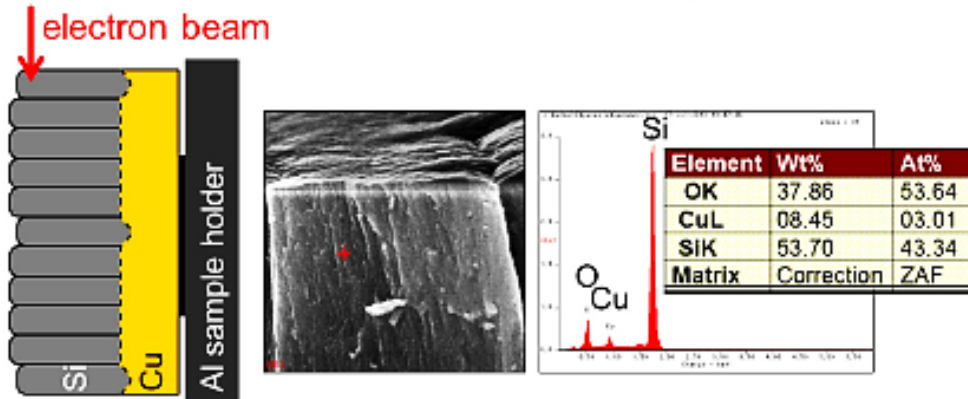


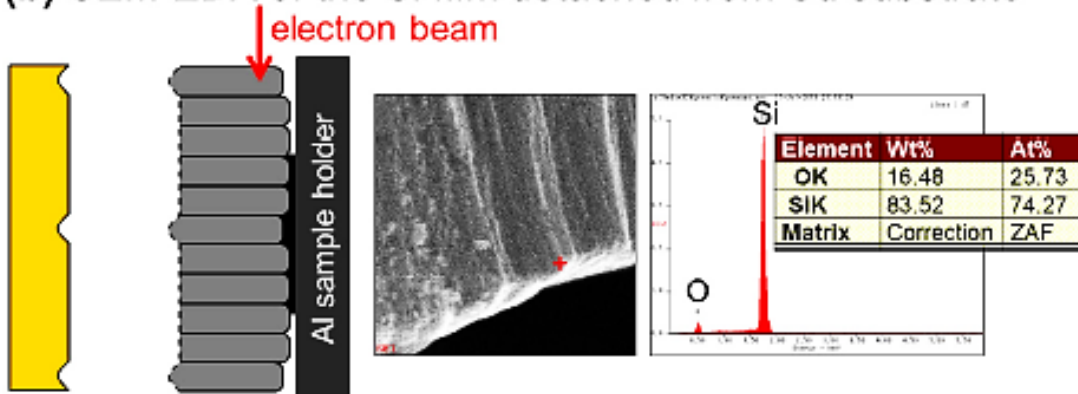
Figure 3-4 Typical SEM-EDX result for a point-measurement of the cross-section of the Si–Cu film (5 wt% Cu source, $T_{\text{sub}} = 100 \text{ }^\circ\text{C}$, without post-annealing). This kind of measurement was made for ~ 50 different points at various distances from the interface to make a profile for each sample in Figure 3-3e. Standard deviation of the Cu content values is shown in (d) for the film from 5 wt% Cu source.

clearly demonstrated the Cu concentration gradient in the direction perpendicular to the Cu substrate. The Si film that was deposited at $T_{\text{sub}} = 100\text{ }^{\circ}\text{C}$ showed minimal intermixing of Si with Cu, while the

(a) Normal SEM-EDX of the Si film on Cu substrate



(b) SEM-EDX of the Si film detached from Cu substrate



(c) SEM-EDX of the Si film on Cu from the top surface

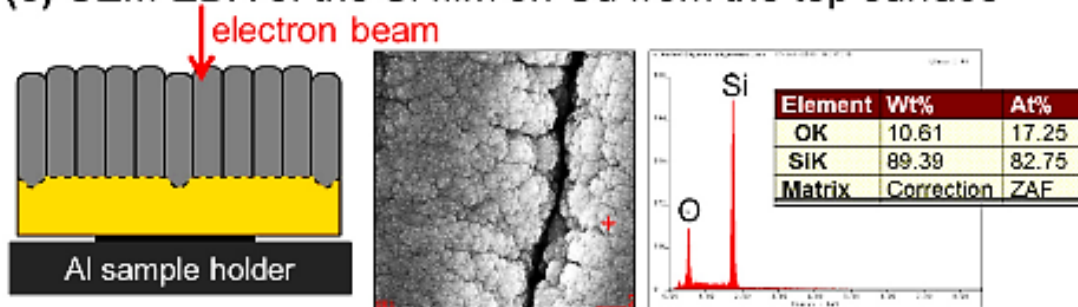


Figure 3-5 SEM-EDX measurements examining the background signal from the Cu substrate. The sample was a pure Si film deposited at $T_{\text{sub}} = 100\text{ }^{\circ}\text{C}$ without post-annealing. The point measurement detected Cu at ~ 3 at% for the top position of the cross-section of the Si film on the Cu substrate. Whereas no Cu was detected for the top position of the cross-section of the Si film detached from the Cu substrate and for the top surface of the Si film on the Cu substrate.

film that was deposited at $T_{\text{sub}} = 300$ °C showed a thin intermixed layer of 0.1–0.2 μm , which is consistent with the results of our previous report [116]. A higher deposition temperature ($T_{\text{sub}} = 500$ °C) can yield a thicker intermixed layer but containing a mixed amorphous-microcrystalline phase with small porosity, which resulted in poor cycle performance [116]. In contrast, the co-deposited Si–Cu films showed a much thicker compositional gradient layer; ≥ 5 at% for ≤ 2 μm from the interface and 2–3 at% for the entire thickness range of 4.5- μm -thick films without the need to increase T_{sub} . The Cu content of the gradient layer can also easily be increased to ~ 40 at% by increasing the Cu content of the source.

3.4 Crystallinity and lithiation behavior of porous Si–Cu films

To enhance the adhesion of the Si and Si–Cu films with the Cu substrate, post-annealing was performed at $T_{\text{an}} = 600$ °C. The high-magnification cross-sectional SEM images of the films at the interface with the Cu substrates clearly show the significant improvements in the adhesion compared with the as-deposited films (Figure 3-3); the gap was very small for the pure Si film (Figure 3-6a) and was eliminated for the Si–Cu films (figure 3-6b–d). Nitrogen adsorption measurement was also performed for the Si–Cu film deposited using a 5 wt% Cu source. It showed that the Si–Cu film had a specific surface area as high as $95.8 \text{ m}^2 \text{ g}_{\text{film}}^{-1}$ (Figure 3-6f) and pore volume as high as $0.237 \text{ cm}^3 \text{ g}_{\text{film}}^{-1}$ (Figure 3-6g). When we use the mass density of Si (2.33 g cm^{-3}), 1 g of the Si–Cu film has 0.429 cm^3 solid with 0.237 cm^3 pore, resulting in porosity of 0.356, which is closed to the porosity value (0.31) estimated using the mass density of the film (1.60 g cm^{-3}) for the as-deposited film.

The crystallinity of the Si-base films is one of the most significant factors to affect the electrochemical performance, and causes different lithiation behaviour at the initial cycle. We previously reported a

change in the crystallinity of pure Si films with T_{sub} [116]; deposition at low T_{sub} values of 100 and 300 °C yielded Si films with a purely amorphous phase that did not crystallize upon post-annealing

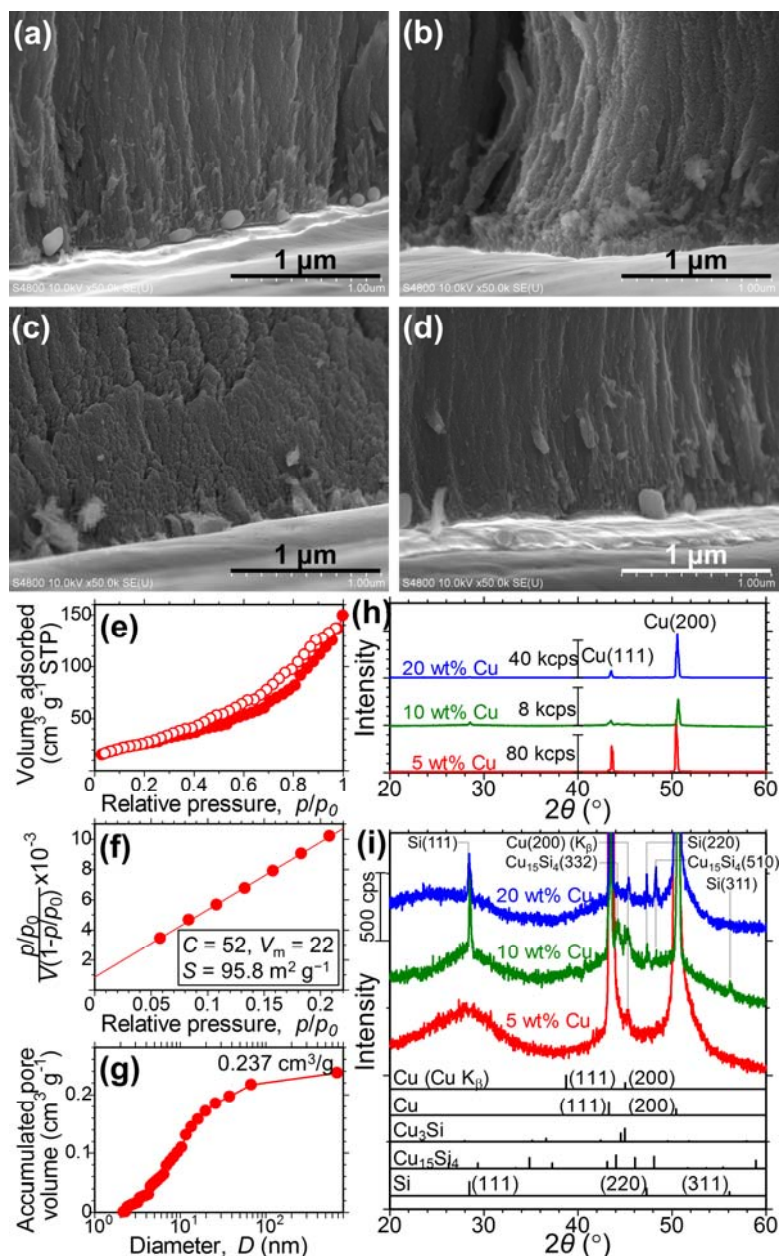


Figure 3-6 Structure analysis of the Si–Cu films deposited on Cu substrates at $T_{\text{sub}} = 100$ °C and post-annealed at $T_{\text{an}} = 600$ °C. Cross-sectional SEM images at the interface between the Cu substrates and Si–Cu films made with (a) pure Si source, (b) 5 wt% Cu source, (c) 10 wt% Cu source, and (d) 20 wt% Cu source. BET analysis of the Si–Cu films deposited on a 10 μm-thick Cu foil at $T_{\text{sub}} = 100$ °C and post-annealed at $T_{\text{an}} = 600$ °C; (e) N₂ adsorption isotherm, (f) BET plot, and (g) pore size distribution. XRD patterns of Si–Cu films deposited at $T_{\text{sub}} = 100$ °C and post-annealed at $T_{\text{an}} = 600$ °C; in full intensity scales (h) and a magnified intensity scale (i).

at $T_{\text{an}} = 600\text{ }^{\circ}\text{C}$, while deposition at the high T_{sub} of $500\text{ }^{\circ}\text{C}$ yielded Si films with a mixed amorphous and microcrystalline phase. The solid phase crystallization of amorphous Si (a-Si) is known to occur upon annealing at temperatures $\geq 700\text{ }^{\circ}\text{C}$ for 10 min or more [100], but it does so more easily in coexistence with a metal, which is known as metal-induced crystallization [119-123]. I therefore analysed Si–Cu samples that had been post-annealed at $T_{\text{an}} = 600\text{ }^{\circ}\text{C}$ by XRD (Figure 3-6h and i). From the spectra for full y-axis range (Figure 3-6h), we can see that the Cu substrates had (200) out-of-plane orientation. The film from 5 wt% Cu source showed a broad peak of a-Si at $\sim 29^{\circ}$ [124], in addition to the intense Cu (111) and (200) peaks at 43.3° and 50.4° , respectively. Because of the strong (200) orientation of the Cu substrate, we can also see its diffraction at 45.0° due to the weak Cu K_{β} line. The film from 10 wt% Cu source showed the sharpened peak of a-Si at $\sim 29^{\circ}$ with a strong Si (111) peak, showing the partial crystallization of a-Si. In addition to the above-mentioned peaks from the Cu substrate at 43.3° , 45.0° and 50.4° , the film had peaks of $\text{Cu}_{15}\text{Si}_4$ (332) at 44.1° [123, 125], Si (220) at 47.3° and Si (311) at 56.1° [124]. The film from 20 wt% Cu source showed similar spectrum with that from 10 wt% Cu source with some differences; the broad peak of a-Si at $\sim 29^{\circ}$ got weakened, the $\text{Cu}_{15}\text{Si}_4$ (510) peak at 48.1° got pronounced, and the Si (311) peak at 56.1° got less obvious. In the standard powder pattern, Cu_3Si has intense diffractions by the (012) and (300) planes at 44.6° and 45.0° with relative intensities of 64 and 100, respectively [123, 125]. We attribute the peak at 45.0° to the Cu (200) diffraction of the Cu K_{β} line for the film from 5 wt% Cu source, which show no other diffraction except for those from the Cu substrate, but the Cu_3Si (012) and (300) peaks may overlap with it for the films from 10 to 20 wt% Cu sources. From these analysis, we can conclude that the film from 5 wt% Cu source was fully amorphous and that the films from 10 wt% and 20 wt% Cu sources were partially crystallized with c-Si and crystalline copper silicide phases (with c- $\text{Cu}_{15}\text{Si}_4$ phase, at least). Next, I examined the electrochemical performances of the Si–Cu films via galvanostatic charge–discharge measurements using three-electrode beaker cells. First, the

performance was evaluated using an electrolyte of 1 M LiPF₆ in EC/diethyl carbonate (1:1 v/v); however, all the Si and Si–Cu films showed poor cycle performances (Figure 3-7). Hydrofluoric acid possibly formed because of insufficient removal of oxygen/water and the excess amounts of the electrolyte used in the beaker cells, which resulted in the etching and delamination of the films. Therefore, I subsequently used 1 M LiClO₄ in EC/PC (1:1 v/v) as the electrolyte.

Figure 3-8 shows the lithiation/delithiation behaviour of these films over the first two cycles measured at a rate of 0.1C. Note that the rate is calculated by assuming 100% Si content, and thus the actual rate is higher than 0.1C for the Si–Cu films. In the case of the Si–Cu film with the lowest Cu content (5 wt% Cu source), the first plateau appeared at ~200 mV vs. Li/Li⁺ during lithiation, which

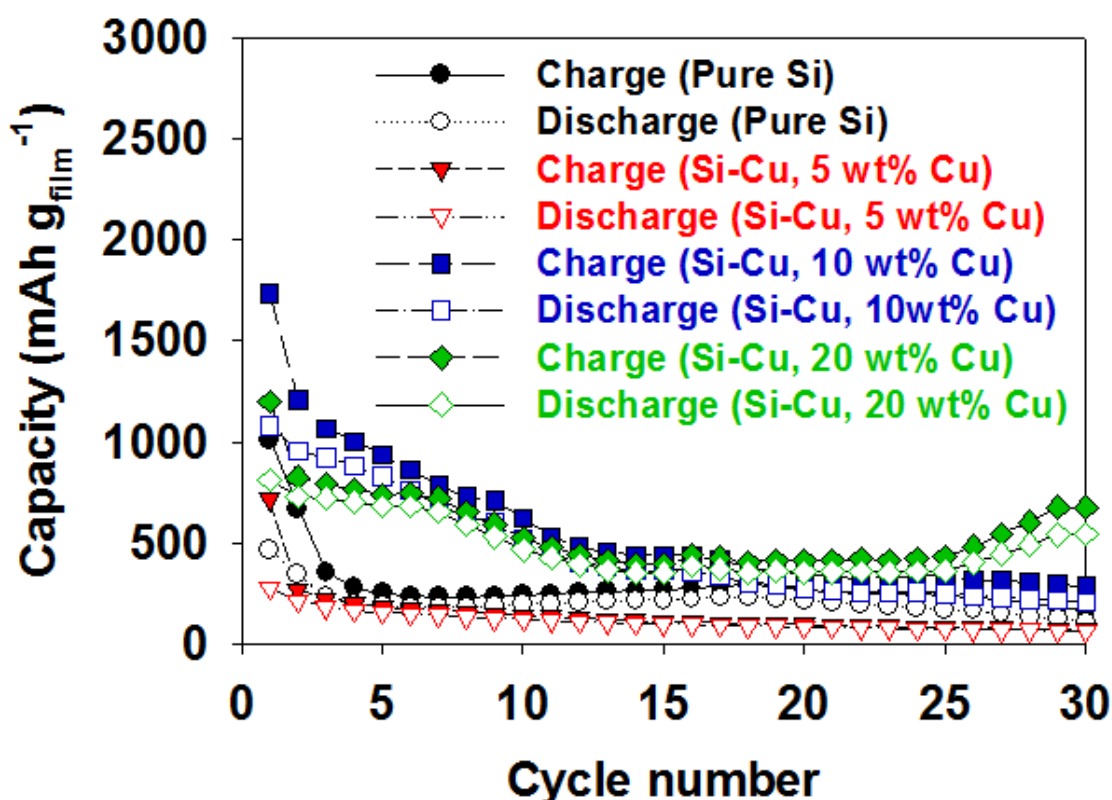


Figure 3-7 Electrochemical performance of pure Si and Si–Cu films using lithium hexafluorophosphate (LiPF₆, 1 M) in a 1:1 (v/v) mixture of EC and diethyl carbonate as the electrolyte with H₂O content of less than 20 ppm (Kishida Chemical, Osaka, Japan)

corresponds to the lithiation potential of a-Si (Figure 3-8a), while the Si–Cu films with higher Cu content (10 and 20 wt% Cu sources) showed plateaus at ~ 150 mV (Figure 3-8b) and ~ 100 mV (Figure 3-8c), respectively, because of partial crystallization of the Si–Cu films (Figure 3-6i). It is known that the initial lithiation of c-Si generally occurs at ~ 100 mV, which corresponds to the conversion of c-Si to a-Li_xSi ($x \sim 3.5$) [66, 68, 126], while a-Si converts into a-Li_xSi ($x = 0–2.0$) at ~ 200 mV, and then converts further into a-Li_xSi ($x = 2.0–3.5$) at ~ 100 mV [66, 67, 127–130].

Figure 3-8d–g shows differential capacity curves of the Si–Cu films that were derived from the

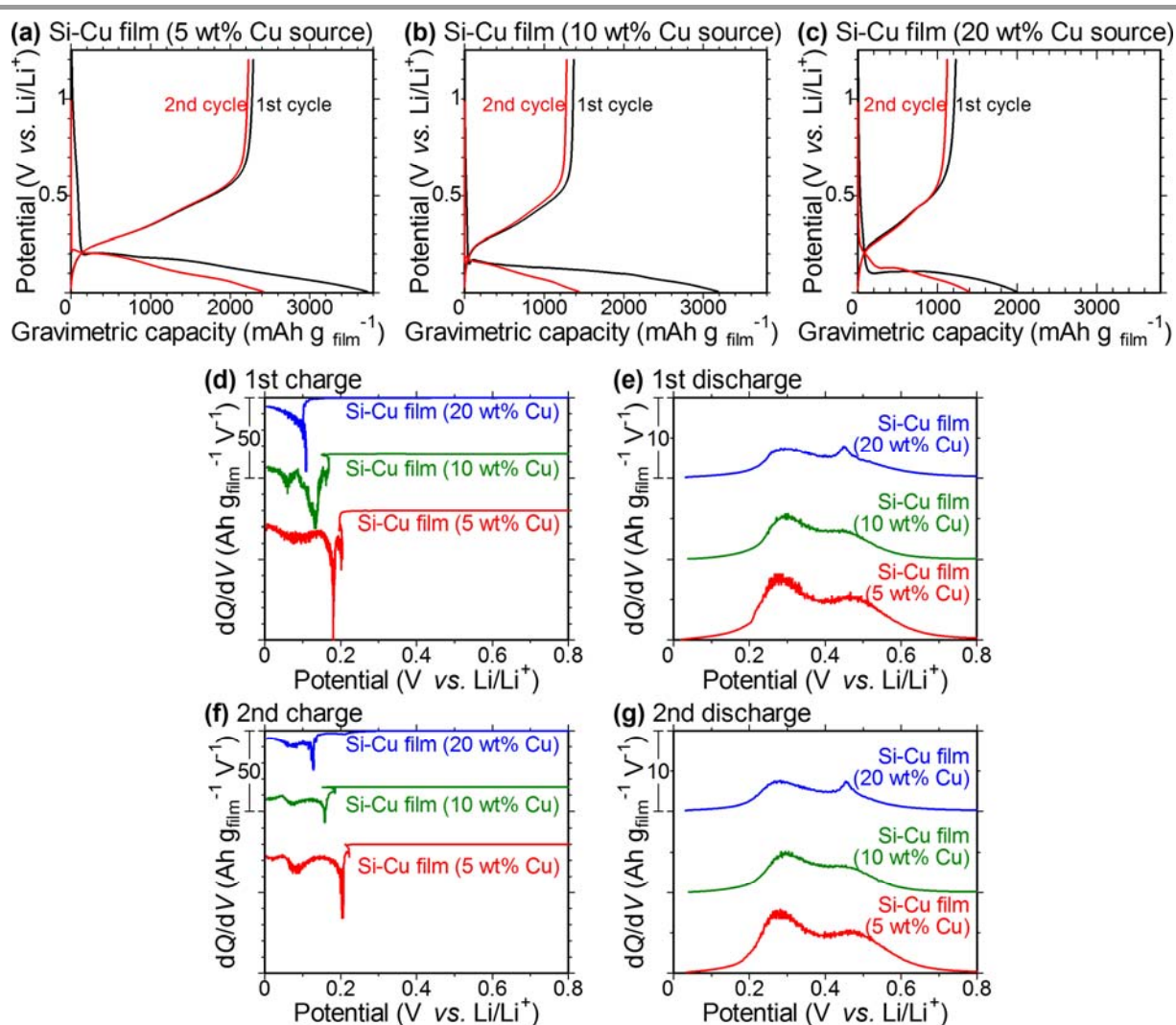


Figure 3-8 Voltage-capacity profiles of Si–Cu films with (a) 5 wt% Cu source ($t_{\text{eff}} = 3.3 \mu\text{m}$), (b) 10 wt% Cu source ($t_{\text{eff}} = 3.4 \mu\text{m}$), and (c) 20 wt% Cu source ($t_{\text{eff}} = 3.1 \mu\text{m}$) for the first and second cycles measured at a rate of 0.1C. (d–g) Differential capacity curves derived from the voltage-capacity profiles shown in parts (a–c).

voltage-capacity profiles (Figure 3-8a–c). During the first charging process (Figure 3-8d), the Si–Cu film with the low Cu content (5 wt% Cu source) showed three peaks; small and dominant sharp peaks at ~200 and ~180 mV, respectively, and another broad peak at ~90 mV, which correspond to potentials forming a-Li_{2.0}Si, a-Li_{3.5}Si, and a-Li_{3.75}Si, respectively [70, 87, 130, 131]. The Si–Cu film with moderate Cu content (10 wt% Cu source) also showed three different peaks, which shifted to lower potentials of ~160, ~130 and ~70 mV. In contrast, the Si–Cu film with the highest Cu content (20 wt% Cu source) showed one dominant peak at ~100 mV, which was similar to the lithiation behaviour of c-Si [87]. During the first discharge process (Figure 3-8e), all the Si–Cu films exhibited the same initial delithiation peak at ~270 mV, which corresponds to the transformation of a-Li_{3.5}Si into a-Li_{2.0}Si. A difference was found at the second delithiation peak, where the Si–Cu films with the lowest and moderate Cu contents (5 and 10 wt% Cu sources) both exhibited a broad peak in the 400–500 mV range, corresponding to the transformation of a-Li_{2.0}Si into a-Si [68, 70], while that with the highest Cu content (20 wt% Cu source) showed a sharp peak at ~450 mV, corresponding to the transformation of c-Li_{3.75+δ}Si into a-Li_xSi [68, 131]. In the second charging process (Figure 3-8f), the lithiation potentials shifted to higher potentials in all the Si–Cu films, which is possibly due in part to the conversion of c-Si to a-Si during the first charge–discharge process. Little change was found in the second discharge process (Figure 3-8g) when compared with the first discharge process (Figure 3-8e). The dominant first peak at ~270 mV appeared for all films, with the broad second peak at 400–500 mV for the Si–Cu films with the lowest and moderate Cu contents (5 and 10 wt% Cu sources) and the sharp second peak occurred at ~450 mV for the Si–Cu film with the highest Cu content (20 wt% Cu source). The high Cu content of the film may have inhibited complete lithiation of Si in the initial cycles, and thus c-Si content possibly remained in this film for at least the first two cycles.

3.5 Cycle performance of porous Si–Cu anodes

Figure 3-9 shows the electrochemical behaviour of the Si and Si–Cu films. The initial charge and discharge capacities of the pure Si film reached 3625 and 2520 mAh g_{film}⁻¹, respectively (Figure 3-9a). The large irreversible capacity is attributed to Si consumption during the formation of the solid electrolyte interphase (SEI), which was significant because of the large exposed surface of the porous Si film. The Coulombic efficiency increased to 98% after 20 cycles, but then dropped abruptly at the 30th cycle (Figure 3-9b). This occurs because the Si film begins to peel off the Cu substrate because of the large volume change during the charge–discharge cycles. Finally, the discharge capacity at the 50th cycle decreased to ~500 mAh g_{film}⁻¹. These performances were worse when compared with the results of our previous report [116], which were a Coulombic efficiency of ~99.5% and a discharge capacity of ~1000 mAh g_{film}⁻¹ at the 50th cycle. This difference originated from the different cell structure (coin cells) and electrolyte (1 M LiPF₆ in a 1:1:1 (v/v) mixture of EC, DMC, and ethyl methyl carbonate) that were used in the previous work. In contrast, the small addition of 5 wt% Cu to the Si source yielded a Si–Cu film with remarkably enhanced cycle performance. The film exhibited a high initial charge capacity of 3425 mAh g_{film}⁻¹ with a discharge capacity of 2073 mAh g_{film}⁻¹ (Figure 3-9a). The capacity retention of this film was the highest among all the films examined, at 73% and 60% with discharge capacities of 1518 and 1250 mAh g_{film}⁻¹ at the 50th and 100th cycles, respectively. The Coulombic efficiency exceeded 98% at the 10th cycle and remained stable at ~99% for 100 cycles. These values are much higher than that of the pure Si film, possibly because of the gradient volume expansion of the Si–Cu film in the direction perpendicular to the Cu substrate. Further increases in the Cu content of the Si source to 10 and 20 wt%, however, resulted in poorer performance levels with lower initial charge and discharge capacities, Coulombic efficiencies, and capacity retention. This is because of the partial crystallization of the Si–Cu films (Figure 3-6i) and the inhibited lithiation (Figure 3-8) for those with higher Cu contents.

The volumetric capacity ($\text{mAh cm}_{\text{film}}^{-3}$) and the areal capacity ($\text{mAh cm}_{\text{anode}}^{-2}$) (shown as inset of Figure 3-9a) of the films are highly important factors for practical battery devices. The volumetric

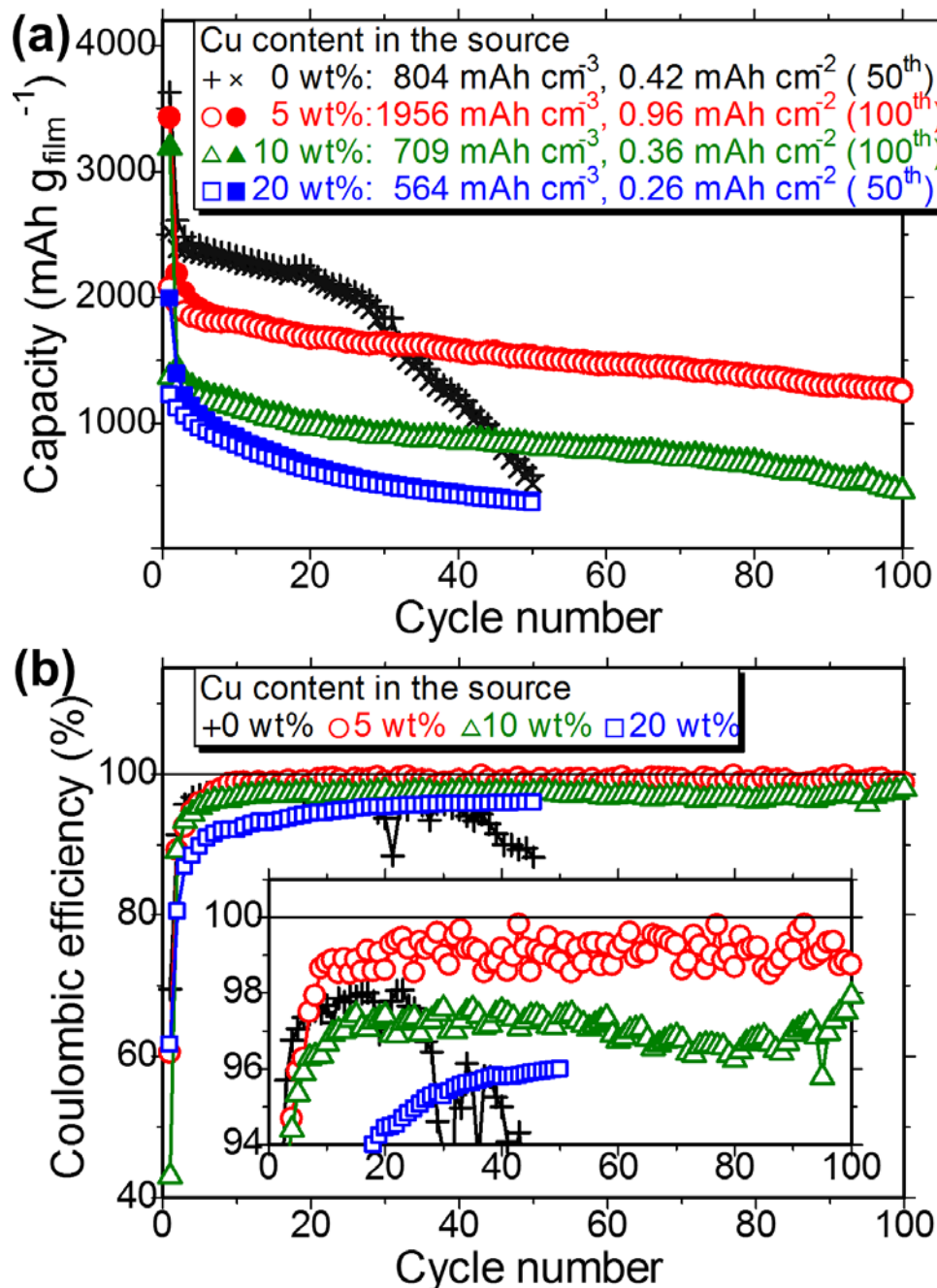


Figure 3-9 Electrochemical performance of pure Si and Si–Cu films deposited at $T_{\text{sub}} = 100$ °C and post-annealed at $T_{\text{an}} = 600$ °C. $t_{\text{eff}} = 3.5, 3.3, 3.4,$ and 3.1 μm for films from Si sources with 0, 5, 10, and 20 wt% of Cu, respectively. The cycle performance at 0.1C was expressed as gravimetric capacity (a) with volumetric capacity and areal capacity values as inset. (b) Coulombic efficiency of the films. Cycle tests were performed at 0.1C for all cycles (1–100).

capacity of the Si–Cu film (5 wt% Cu source) was $1956 \text{ mAh cm}_{\text{film}}^{-3}$ after 100 cycles, which is much higher than that of commercial graphite anodes. The areal capacity of the Si–Cu film (5 wt% Cu source) was $0.96 \text{ mAh cm}_{\text{anode}}^{-2}$, which is higher than previously reported values for Si nanomaterials ($0.2\text{--}0.4 \text{ mAh cm}_{\text{anode}}^{-2}$ [79, 88]) and porous Si films by ourselves ($0.66 \text{ mAh cm}_{\text{anode}}^{-2}$ [116]), but is still smaller than that of commercial graphite anodes ($\sim 4 \text{ mAh cm}_{\text{anode}}^{-2}$, 18650 Li-ion cells, Sony, Japan) [112] and Si–carbon nanotube hybrid anodes ($2 \text{ mAh cm}_{\text{anode}}^{-2}$) [113]. The cycle performance of the pure Si film in this work is smaller than that in our previous work [116], which is possibly because of differences in the measurement method, and thus the Si–Cu film from the 5 wt% Cu source may perform better in a coin cell test and/or electrolyte of 1 M LiPF₆ in a 1:1:1 (v/v) mixture of EC, DMC, and ethyl methyl carbonate. The Si–Cu film (5 wt% Cu source) showed very stable Coulombic efficiency of $\sim 99\%$ for 10–100 cycles, indicating the formation of stable SEI layer and/or efficient use of Si in the film from the early cycles. The Si–Cu film (10 wt% Cu source) showed lower values of 97% for the first 50 cycles, and fluctuating values in the 96%–98% range for the later cycles. The Si–Cu film (20wt% Cu source) showed a very low initial value, which increased to $>94\%$ after 20 cycles. These values lower than 100% indicate the continuous formation of SEI layer due to the incomplete formation of stable SEI layer in the early cycles. And some fraction of the Si in the films may not have reacted in the early cycles, as can be seen in the small initial charge capacity of $1992 \text{ mAh g}_{\text{film}}^{-1}$ (Figure 3-9a) and in the low potential for lithiation (Figure 3-7), and then gradually began to react with the increasing number of cycles. High Cu contents inhibited the formation of stable SEI layer and/or the reaction of Si with Li⁺.

The Si–Cu film made using 5 wt% Cu in the Si source showed the best cycle performance and I therefore examined its rate performance (Figure 3-10). Beginning with the 0.1C rate, the Si–Cu film showed an initial discharge capacity of $2065 \text{ mAh g}_{\text{film}}^{-1}$, and a reduced capacity of $1755 \text{ mAh g}_{\text{film}}^{-1}$ after 10 cycles with a Coulombic efficiency of $\sim 97\%$. When the charge–discharge rate was doubled

to $0.2C$ from the beginning of the 11th cycle, the Coulombic efficiency dropped slightly to 95%, but immediately increased to $\sim 98\%$ at the 12th cycle, and that efficiency was maintained with a reduced discharge capacity of $1485 \text{ mAh g}_{\text{film}}^{-1}$ at the 20th cycle. Similar changes were observed for later cycles, where the Coulombic efficiency dropped slightly upon an increase in the charge–discharge

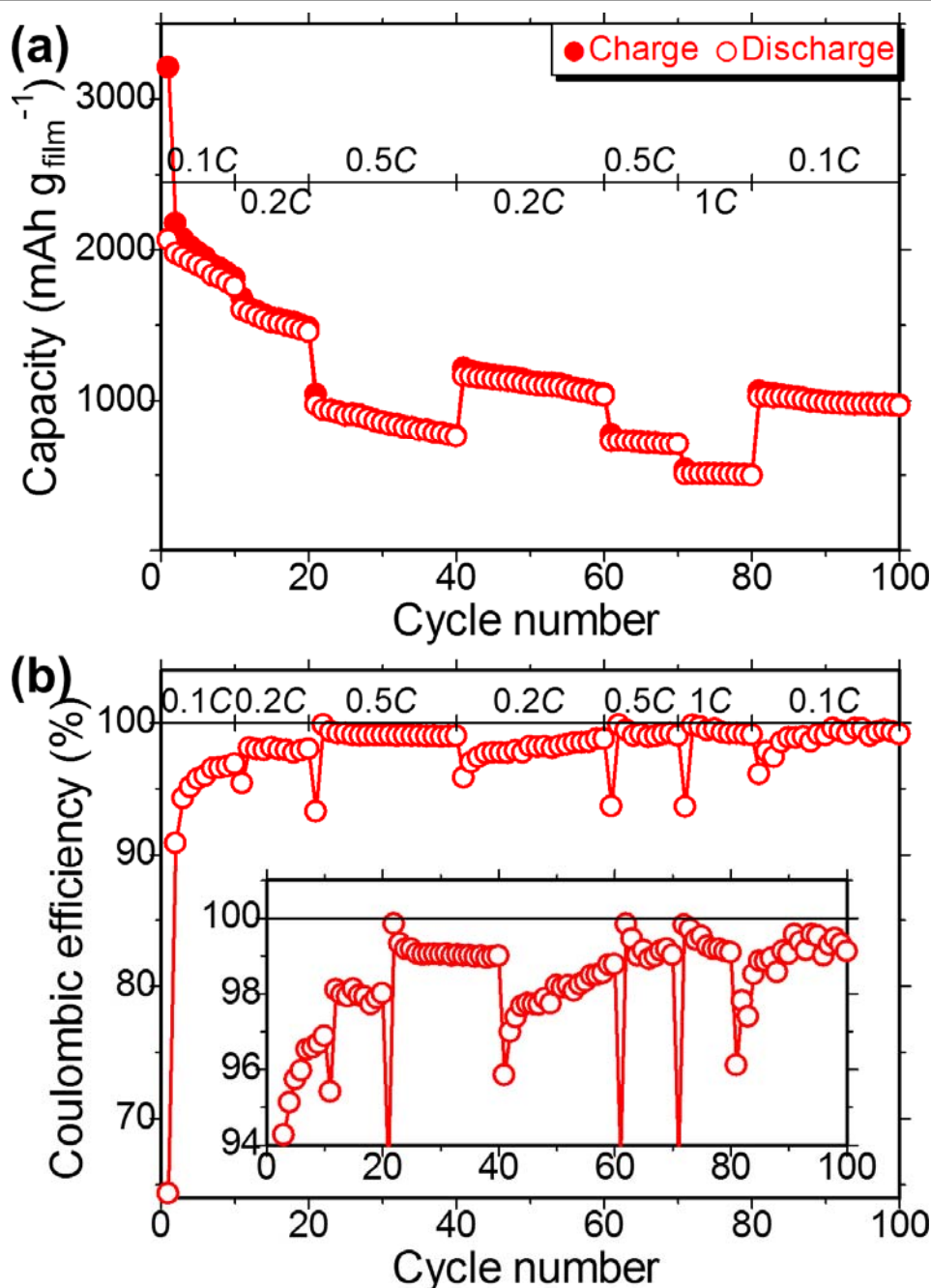


Figure 3-10 (a) Rate capability of representative Si–Cu film ($t_{\text{eff}} = 3.3 \mu\text{m}$, 5 wt% Cu source, $T_{\text{sub}} = 100 \text{ }^\circ\text{C}$, $T_{\text{an}} = 600 \text{ }^\circ\text{C}$). (b) Coulombic efficiency and (c) capacity ratio of charge at the $[n+1]$ th cycle to discharge at the $[n]$ th cycle as a function of cycle number n .

rate but immediately recovered in the next cycle, and maintained a steady value of 98–100% at 0.1C–1C rates for 100 cycles (Figure 3-10b). The discharge capacity decreased/increased upon any increase/decrease in the charge–discharge rate, with continuous gradual decay with increasing numbers of cycles. The discharge capacity decreased to 505 mAh g_{film}⁻¹ at 1C at the 80th cycle and recovered to 964 mAh g_{film}⁻¹ at 0.1C at the 100th cycle (Figure 3-10b). These results show that the stable SEI layer formed on this Si–Cu film and the Si in this film reacted efficiently with the Li⁺ from the early cycles.

3.6 Cycle performance of porous Si–Cu anodes with different cut-off potentials for charge

Figure 3-11 shows the effects of the different cut-off potentials for charge on the electrochemical performances of the Si–Cu films at a rate of 0.1C for 50 cycles. When compared with the 5 mV cut-off, all films with the 100 mV cut-off showed reduced values for the initial charge capacities and reduced or similar values for the initial discharge capacities (Figure 3-11a–c, Figure 3-12a). The Si–Cu film with the lowest Cu content (5 wt% Cu source) showed little change in capacity retention, whereas the Si–Cu films with moderate and highest Cu contents (10 and 20 wt% Cu sources) showed remarkable increases in capacity retention of 6% and 28%, respectively (Figure 3-11d). These results show that the cut-off operation for charge (lithiation) helps the Si–Cu anodes that contain c-Si to increase their capacity retention during cycling, possibly by suppressing the phase transformation, while the cut-off operation does not help the a-Si–Cu anode without c-Si, possibly because of the absence of such a phase transformation, regardless of the cut-off potential. The areal capacities of the Si–Cu films (5, 10 and 20 wt% Cu sources) were 0.8, 0.7 and 0.5 mAh cm_{anode}⁻², respectively (Figure

3-12b), at the 50th cycle. The Coulombic efficiencies of the Si–Cu films (10 and 20 wt% Cu sources)

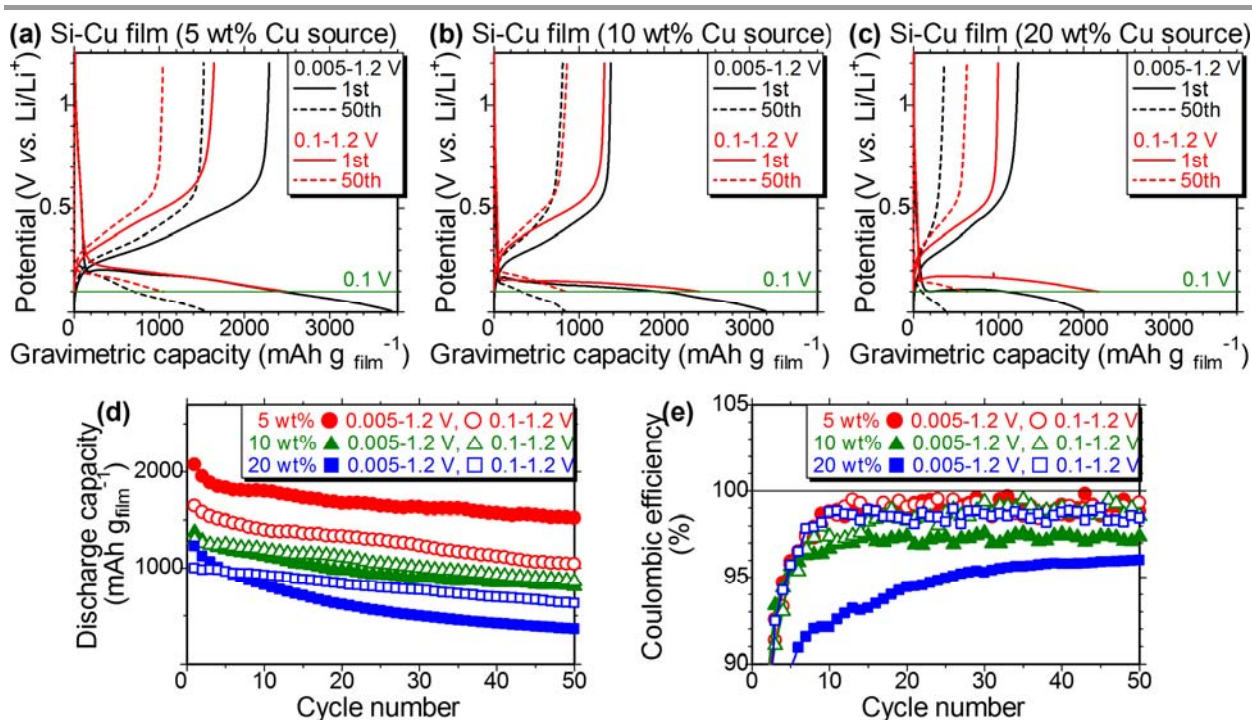


Figure 3-11 Voltage-capacity curves for the first and 50th charge/discharge cycles of Si–Cu films deposited using sources with (a) 5 wt% Cu ($t_{\text{eff}} = 3.3 \mu\text{m}$), (b) 10 wt% Cu ($t_{\text{eff}} = 3.5 \mu\text{m}$), and (c) 20 wt% Cu ($t_{\text{eff}} = 3.2 \mu\text{m}$) with different cut-off operations at 5 and 100 mV for charge. (d) Cycle performance and (e) Coulombic efficiency of the Si–Cu films. The Si–Cu films were deposited at $T_{\text{sub}} = 100 \text{ }^\circ\text{C}$ and post-annealed at $T_{\text{an}} = 600 \text{ }^\circ\text{C}$.

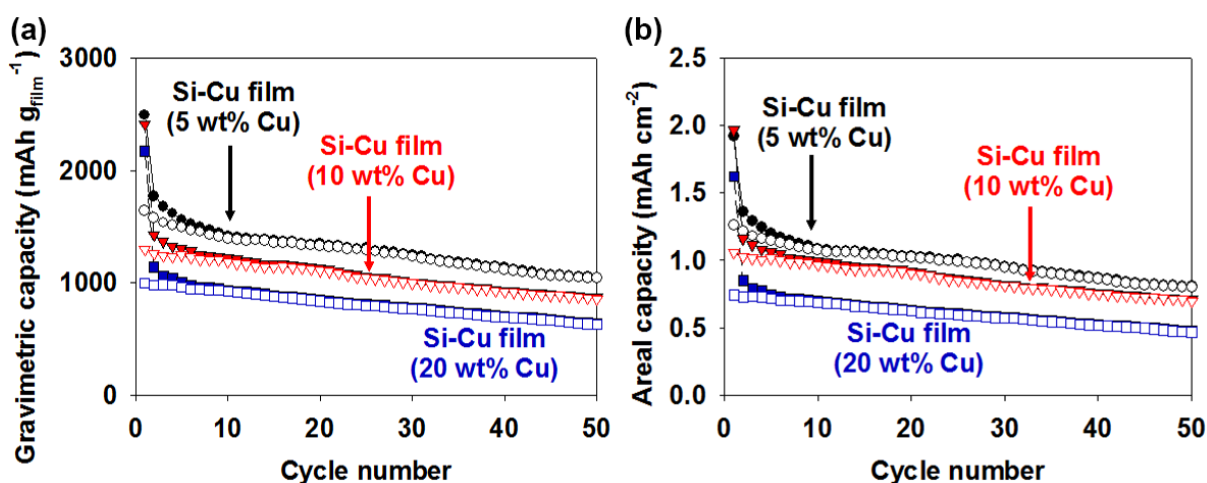


Figure 3-12 Electrochemical performance of Si–Cu films with 100 mV cut-off potential for charge. The Si–Cu films were deposited using Si sources with different Cu contents (5, 10 and 20 wt%) at $T_{\text{sub}} = 100 \text{ }^\circ\text{C}$, post-annealed at $T_{\text{an}} = 600 \text{ }^\circ\text{C}$, and had $t_{\text{eff}} = 3.3, 3.5$ and $3.2 \mu\text{m}$, respectively. (a) Gravimetric capacity and (b) areal capacity of the deposited Si–Cu films.

increased to more than 95% after the fifth cycle, and remained at ~99% for 50 cycles, which are similar values to the efficiencies of the Si–Cu films with the lowest Cu content (5 wt% Cu source) with both 5 and 100 mV cut-offs (Figure 3-11e).

3.7 Failure behavior of pure Si and Si–Cu films after charge–discharge cycles

Figure 3-13 shows top-view SEM images of the pure Si and Si–Cu films before and after 100 cycles. Both films initially had similar morphologies with wall-shaped structures (Figure 3-13a and c), but the pure Si film was completely pulverized into a few μm -sized particles, and peeled off after 100 cycles. This occurred because of large-scale volume change of the pure Si film and its poor adhesion to the Cu substrate. At the region where the Si film was peeling off, SEM-EDX analysis detected Si and Cu at 1.1 and 74.4 at%, respectively, indicating the delamination at the interface between the Si film and the Cu substrate (Figure 3-13e). In contrast, the Si–Cu film (5 wt% Cu source) had square domains with ordered domain boundaries at a pitch of 10–20 μm in one direction, with random domain boundaries in a direction that was roughly perpendicular to the former direction. The ordered domain boundaries apparently originated from the wall-shaped structure found in the as-deposited film (Figure 3-1a), which in turn originated from the stripe patterns on the as-received Cu substrates. Also, the domains contained numerous small cracks. Such a structure could possibly show reversible expansion and shrinkage for 100 cycles without pulverization. The similar results were obtained for the Si–Cu films using 10 and 20 wt% Cu source (Figure 3-14); the films retained mostly attached to the Cu substrate with square domains. The Si–Cu films were delaminated in some areas but mostly remained attached to the Cu substrate (Figure 3-13d), possibly because of the gradient volume expansion of the Si–Cu films with gradient composition profiles and because of stress defocusing on the diffuse interface between the Si–Cu film and the Cu substrate. In the region where the Si–Cu film

was being delaminated, SEM-EDX detected Si and Cu at 15.6 and 40.7 at%, respectively, which indicated that the delamination occurred not at the interface but in the film itself (Figure 3-13f).

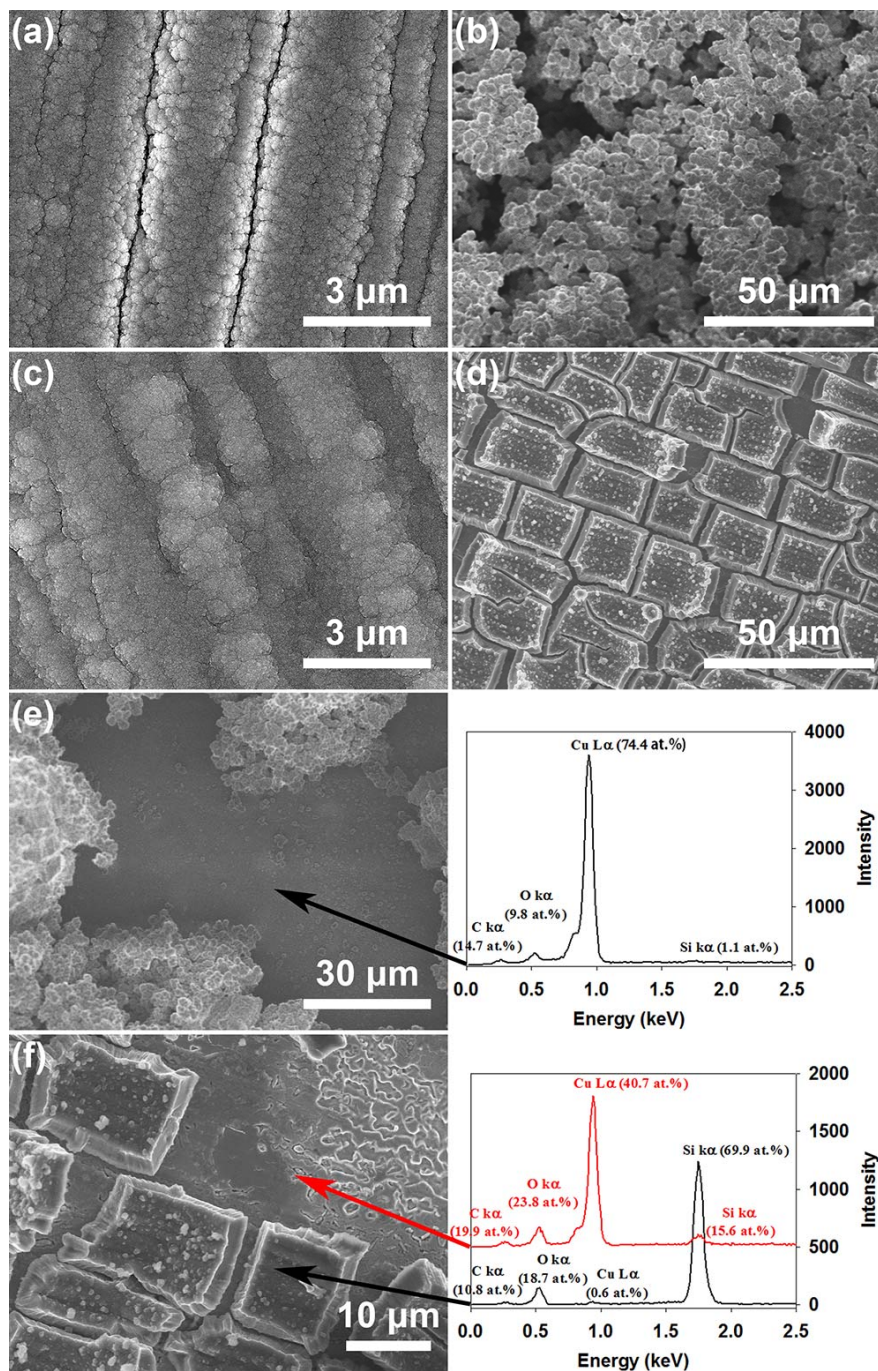


Figure 3-13 Top-view SEM images of the pure Si film ($t_{\text{eff}} = 3.3 \mu\text{m}$) (a,b) and the Si-Cu film (5 wt% Cu source, $t_{\text{eff}} = 3.3 \mu\text{m}$) (c,d) before cycles (a,c) and after (b,d) the 100th discharge. Cycles were performed at 0.1C charge/discharge rates. Both films were deposited at $T_{\text{sub}} = 100 \text{ }^\circ\text{C}$ and post-annealed at $T_{\text{an}} = 600 \text{ }^\circ\text{C}$. SEM-EDX analysis results for the pure Si film (e) and the Si-Cu film (f) after the 100th discharge.

In order to understand the underlying mechanism for the cycle performances, electrochemical impedance spectroscopy was performed for a pure Si film and Si–Cu films (Figure 3-15). The very high frequency region (above 126 kHz) corresponds to the sum of the solution resistance, resistance at the film/Cu interface and external circuits [132], and the degradation of electrolyte can be neglected here due to the huge amount of electrolyte in the beaker cell. For the pure Si film, this resistance increased continuously during cycles, corresponding to the continuous detachment of the film from the Cu substrate. For the Si–Cu films (5 and 20 wt% Cu), on the other hand, this resistance showed very small increase due to the better adhesion at the film/Cu interface as shown in Figure 3-

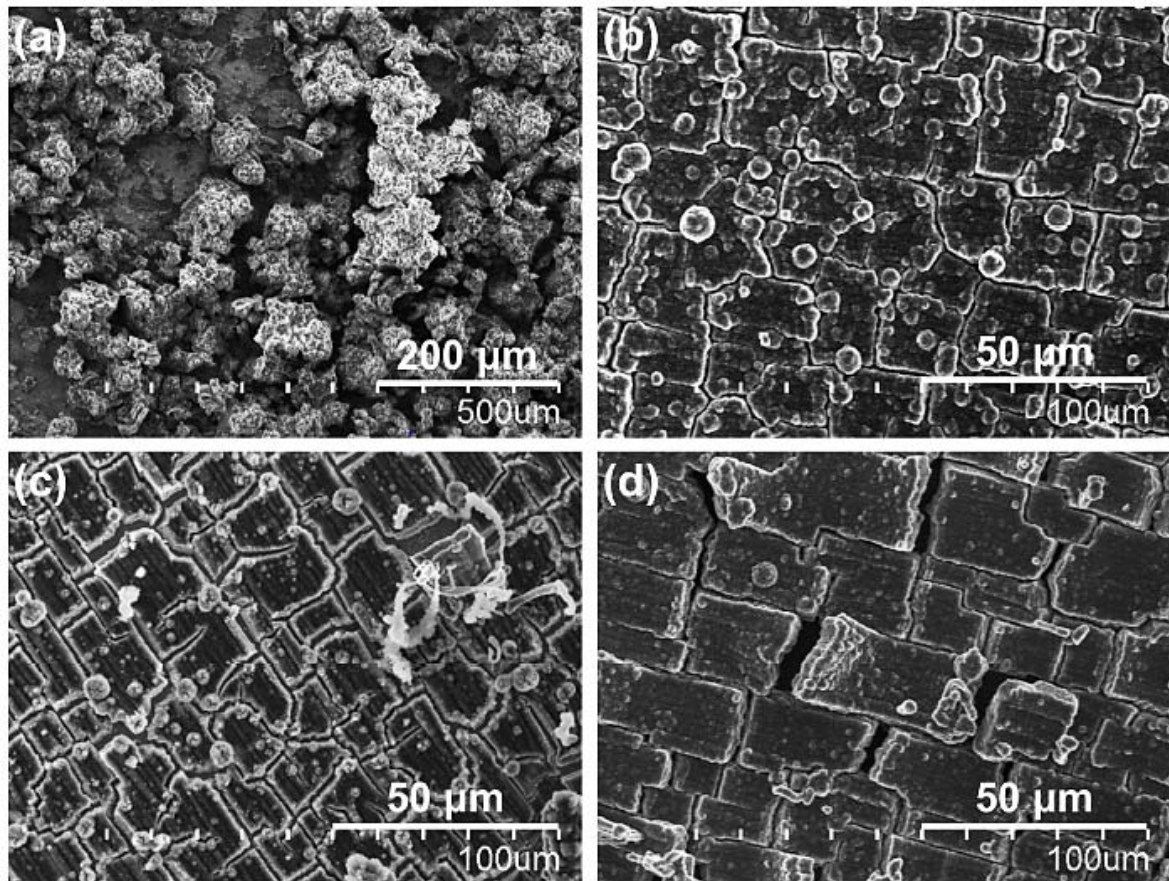


Figure 3-14 Top-view SEM images of the Si and Si–Cu films after the 100th discharge. The films were deposited at $T_{\text{sub}} = 100\text{ }^{\circ}\text{C}$ using (a) a pure Si source and (b) 5 wt%, (c) 10 wt%, and (d) 20 wt% Cu sources and post annealed at $T_{\text{an}} = 600\text{ }^{\circ}\text{C}$. The cycles were performed at 0.3C except for the 1st, 25th, 75th and 100th cycles. The films (a, b, d) were charged at 0.1C, measured by impedance test, and discharged at 0.1C at the 1st, 25th, 75th and 100th cycles.

13 and Figure 3-14. Two semicircles at high and medium frequency range (126 kHz–0.5 Hz) were clearly observed. The small semicircle at high frequency and the large semicircle at middle frequency are attributed to the surface film resistance and charge transfer resistance, respectively [132]. The

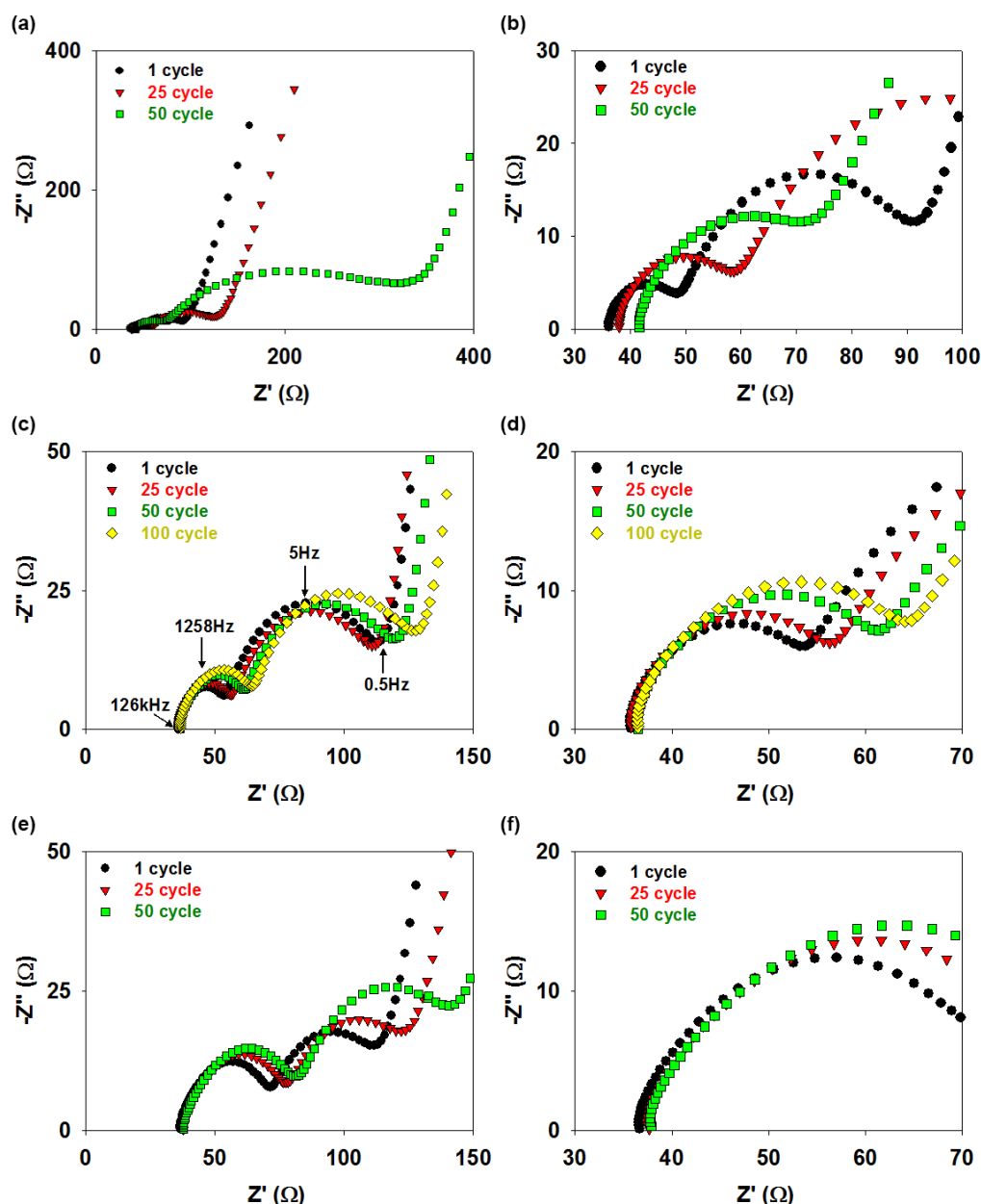


Figure 3-15 Nyquist plots of the (a, b) pure Si films ($t_{\text{eff}} \sim 3.3 \mu\text{m}$) and the Si-Cu films deposited using (c, d) 5 wt% Cu ($t_{\text{eff}} \sim 3.4 \mu\text{m}$) and (e, f) 20 wt% Cu ($t_{\text{eff}} \sim 3.4 \mu\text{m}$) at different cycles. All the films were deposited at $T_{\text{sub}} = 100 \text{ }^\circ\text{C}$ and post-annealed at $T_{\text{an}} = 600 \text{ }^\circ\text{C}$. Cycles were performed at 0.3C except for the 1st, 25th, 50th, and 100th cycles. Impedance measurements were made after the 1st, 25th, 50th, and 100th charge; as soon as potential reached 5 mV during lithiation, the films were left at open circuit voltage for 30 min, and then impedance measurement was performed between 1 MHz and 5 mHz.

surface film resistance of the pure Si films increased quickly, suggesting the continuous formation of the SEI layer possibly due to its fracture (as can be seen in the pulverized Si film at the 100th cycle in Figure 3-13b) and its regeneration at the newly exposed Si surface. The surface film resistance remained smallest for the Si–Cu film (5 wt%) while somewhat larger for the Si–Cu film (20 wt%). As the dQ/dV analysis (Figure 3-8) showed, lithiation proceeded at lower potential for the Si–Cu film (20 wt%) than the Si–Cu film (5 wt%), resulting in the longer period at low potential and thus in the possible enhancement in the SEI layer formation in the former. The charge transfer resistance of the pure Si film increased significantly in 50 cycles, indicating that the Si film got more and more resistive during the cycles, which will be related with the pulverization of the pure Si film (Figure 3-13b). The Si–Cu film (20 wt%) had the smallest charge transfer resistance at the early cycles, possibly due to the highest Cu content and lowest film resistivity, but the resistance increased continuously and approached to that of the Si–Cu film (5 wt%).

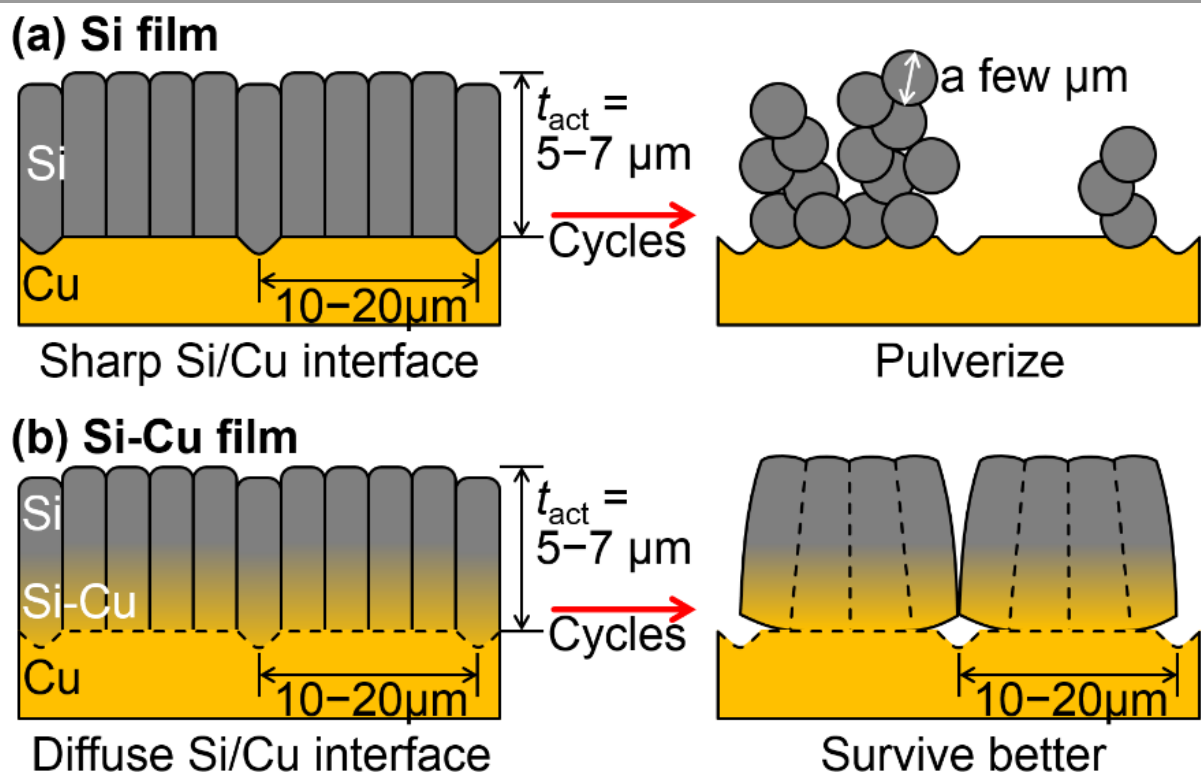


Figure 3-16 Schematic summarizing the initial structures of the porous Si and Si–Cu films and the final structures of these films after charge–discharge cycles.

Figure 3-16 schematically summarizes the initial structures of the pure Si and Si–Cu films and final structures after the charge–discharge cycles. The pure Si films showed pulverization and delamination from the Cu substrates due to the brittle nature of Si and clear difference in the volumetric change between the Si films (up to 400%) and Cu (0%) even with the porous and amorphous structure. They quickly lost electrical contact with the Cu substrates, resulting in rapid capacity fade and short lifetime. In contrast, the Si–Cu films had gradient composition profiles and diffuse interfaces with the Cu substrates, resulting in gradient volumetric expansion in the films and defocused stress at the interface. These Si–Cu films showed no pulverization and suppressed delamination, enabling them to survive for more cycles than the pure Si films. Although further improvements are needed, the gradient composition profile and the diffuse interface that formed spontaneously in the μm -thick Si–Cu film on the Cu current collector certainly enhanced the cycle performance of these Si-base anodes.

3.8 Conclusions

I have realized rapid deposition of 5–7- μm -thick porous Si–Cu films in 1 min on Cu current collectors by RVD using a mixed source of Si and Cu powders. Because of the preferential vaporization and deposition of Cu at the initial stage, which has a higher vapour pressure than Si, these films had 2–4.5- μm -thick composition gradient that changed from a Cu-rich region at the bottom to a Si-rich region at the top. The Cu content of the gradient layer was changed easily by varying the Cu content in the source. A porous structure was built into the Si–Cu films by keeping the substrate at a low temperature ($T_{\text{sub}} = 100\text{ }^{\circ}\text{C}$), while the adhesion of the films to their substrates was improved by post-annealing at a higher temperature ($T_{\text{an}} = 600\text{ }^{\circ}\text{C}$).

The resulting films were mostly amorphous in the low Cu content (5 wt% Cu source) case, whereas they were partially crystallized in the higher Cu content cases (10 and 20 wt% Cu source). The pure

Si film showed poor cycling performance with a discharge capacity of $\sim 500 \text{ mAh g}_{\text{film}}^{-1}$ at $0.1C$ at the 50th cycle, while Cu addition at even the smallest content (5 wt% Cu source) enhanced the film performance remarkably to 1518 and 1250 $\text{mAh g}_{\text{film}}^{-1}$ at the 50th and 100th cycles, respectively. The Cu composition gradient in the film and the diffuse interface between the film and the Cu substrate possibly yielded the gradient volume expansion and the stress defocusing at the interface. The film had a 10–20- μm -sized square domain structure during cycling, which showed no pulverization and suppressed the delamination from the substrate. However, excess addition of Cu (10 and 20 wt% Cu sources) reduced the cycle performances of the Si–Cu films due to partial crystallization, inhibited Si lithiation, and inhibited formation of stable SEI layer. The increased cut-off potential for charge from 5 to 100 mV vs. Li/Li^+ , improved the cycle performances of the Si–Cu films with higher Cu contents (10 and 20 wt% Cu sources) that contained some crystalline phases, but did not do so for the Si–Cu film with the lowest Cu content (5 wt% Cu source) that had a mostly amorphous phase.

The Si–Cu film with the lowest Cu content (5 wt% Cu source) showed a gravimetric capacity of 1250 $\text{mAh g}_{\text{film}}^{-1}$, a volumetric capacity of 1956 $\text{mAh cm}_{\text{film}}^{-3}$, and an areal capacity of 0.96 $\text{mAh cm}_{\text{anode}}^{-2}$ at the 100th cycle. Although further improvement is required, this performance is encouraging for practical use of these Si-base anodes in lithium secondary batteries when we consider the quick and simple fabrication process for these Si–Cu films using inexpensive Si and Cu powder sources, along with the applicability of the RVD method to various substrates.

Chapter 4 – Carbon nanotubes (CNTs) and Si hybrid films as anodes in LiBs

4.1 Introduction

The micrometer-thick Si-Cu film, fabricated rapidly by Cu co-deposition with Si in RVD system, demonstrated that gradient Si-Cu layer can enhance the adhesion at the interface between Si(-Cu) layer and Cu current collector in spite of fairly large thickness ($t_{\text{eff}} = 3\text{--}4\ \mu\text{m}$) with high packing density of $\sim 1.5\ \text{g cm}^{-3}$. The films achieved enhanced electrochemical performance for 100 cycles compared with the pure Si film, as stated in Chapter 3. However, cycle stability for long cycles still remains a challenge because of highly dense Si-Cu films. It was reported that one-dimensional structure of Si, such as Si nanowires, on the other hand, was close to an ideal structure for improvement of electrochemical performance, but complicated process with expensive/explosive silane source used for nanowire makes the low-cost production difficult.

Since the discovery of CNTs [133], lots of efforts improving the performance of energy storage devices, such as electrochemical capacitors (ECs) and lithium ion batteries (LiBs), by using CNTs have been reported [134, 135]. Especially, CNTs have been used as an additive either in the anode or cathode for LiBs due to high theoretical conductivity over $5 \times 10^5\ \text{S cm}^{-1}$ at room temperature [136] and high aspect ratio, which enable CNTs to make a role of electron/hole pathway. The incorporation of CNTs as a conductive additive at a lower weight loading than conventional carbon black and

graphite presents that the use of CNTs could represent more than an order of magnitude reduction in additive mass [25, 137]. In addition, CNTs have the capability to be assembled free-standing carbon electrode in electrochemical application without any binder or current collector, or a physical support and sufficient electronic transport for high capacity anode materials like silicon or germanium [25].

Although further research is needed to solve the issues including high irreversible capacity [39, 40, 138], the use of CNTs in LiBs is attractive due to their flexibility as well as the effective diffusion of Li^+ ions into their porous matrix.

In this chapter, I designed the core-shell wall structure of CNT–Si hybrid film by depositing Si onto the standing walls of shrunk CNT arrays on Cu substrate. I designed such a hybrid film to make the best combination of rapid vapor deposition for Si fabrication and self-organized CNT structure. In Chapter 2, dense Si films deposited at higher substrate temperatures showed good initial performances due to their stability against oxidation in air and suppressed SEI formation but poor long-cycle performances due to the volumetric change during charge–discharge cycles. Thus I used the CNT walls as templates and deposited Si on them at high temperature to realize dense Si films on CNT walls while spaces for volume expansion between the walls. Moreover, in order to avoid the side reaction of CNTs against electrolyte, the vertical aligned CNTs were covered completely with Si by RVD.

4.2 Materials and methods

4.2.1 Synthesis of vertical aligned CNTs

Cu plates with 15 mm diameter and 0.5 mm thickness were used as substrates, cleaned by sonication in isopropanol for 10 min, and further exposed to UV- O_3 for 3 min to remove organic contaminants

on their surface, followed by annealing treatment under 4 vol% H₂/Ar at ambient pressure and 800 °C for 10 min. Synthesis of CNT arrays on Cu substrates were conducted by the collaborative researcher Mr. Nuri Na. The combination of 10 nm-thick Ta and 15 nm-thick TiN as diffusion barriers were deposited, followed by 4 nm-thick Fe or Co thin layers using a magnetron sputtering apparatus (MPS-2000HC2S, ULVAC, Chigasaki, Japan). Vertical-aligned CNT arrays were grown from Fe or Co catalyst particles and 0.2 Torr C₂H₂ at 700 °C for 5–10 min using chemical vapour deposition (CVD) method. The wall-shape structure of the vertical aligned CNTs was developed by a drop of ethanol (EtOH), which cause shrinkage of CNTs by surface tension during drying.

4.2.2 Fabrication of CNT–Si hybrid films

The CNT–Si hybrid films were fabricated by Si deposition on ethanol (EtOH)-treated and untreated CNTs on Cu substrates using RVD in 2 min. Si sources were prepared using the same procedures as previously stated in Chapter 2. The Si sources were heated by resistive heating of carbon boat under 0.1 Torr Ar to T_{boat} of 2000–2400 °C, while substrate was heated to T_{sub} of 400 °C. After Si deposition, the CNT–Si hybrid films were further annealed at T_{an} of 600 °C under 4 vol% H₂/Ar at ambient pressure to improve adhesion between Si and Cu substrate. Before and after Si deposition, each sample was weighed by microbalance with a precision of 1 µg.

4.2.3 Characterization

The analysis of microstructure of ethanol-treated and untreated CNTs-Si core-shell wall films was carried out using field-emission scanning electron microscopy (SEM; S-4800, Hitachi, Tokyo, Japan). To evaluate electrochemical performances of CNT-Si hybrid films, three-electrode beaker cells were

assembled in Ar glove box, which had a CNT-Si hybrid film as the anode, Li metal as the cathode and reference electrode in ~40 mL of electrolyte (1 M LiClO₄ in a 1:1 (v/v) mixture of EC and PC with H₂O content of less than 20 ppm, Kishida Chemical, Osaka, Japan). Charge/discharge measurements by constant-current (CC) mode were carried out in the potential range between 0.005–1.200 V vs. Li/Li⁺ at the rate of 0.4 A g_{Si}⁻¹ (corresponding to ~0.1C) using galvanostatic cycling with potential limitation system (Bio-Logic VMP3, Claix, France). C-rate was determined using the weight of Si and a theoretical value of 4200 mAh g_{Si}⁻¹. In this work, I define the Li⁺ insertion to CNT-Si films as “charge” and Li⁺ extraction from CNT-Si films as “discharge”.

4.3 Structure change in CNT arrays by capillary action

Figure 4-1 shows photographs the Cu substrates with CNT arrays grown by Fe catalyst (deposited in a circular area of 8 mm in diameter), without and with ethanol treatment and without and with Si deposition. The untreated CNT arrays had a smooth surface, resulting in a CNT-Si hybrid film with plain surface. While ethanol-treated CNT arrays had clear patterns on the surface, resulting in a CNT-Si hybrid film with patterns after Si deposition at T_{sub} of 400 °C.

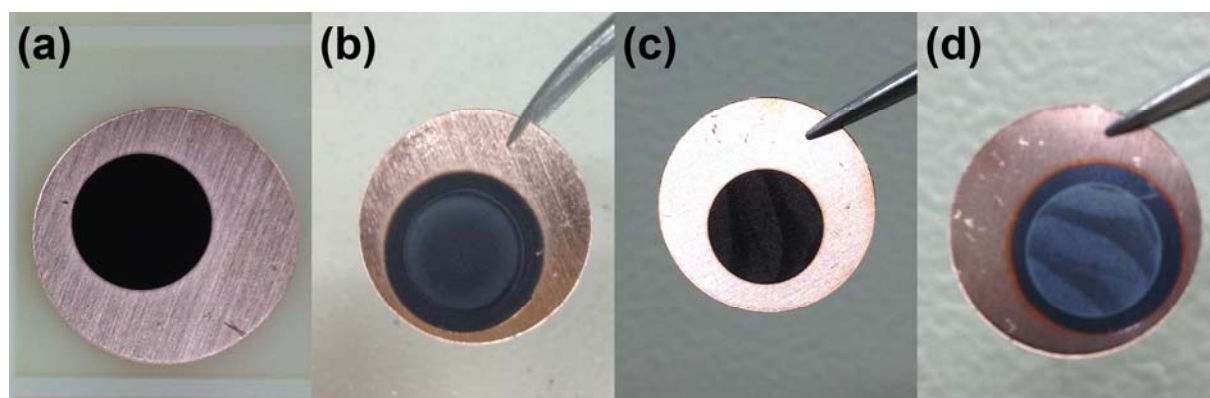


Figure 4-1 Photographs of (a) vertical aligned CNTs grown on Cu substrate, (b) CNTs-Si hybrid film deposited on as-grown CNTs, (c) ethanol-treated CNTs and (d) CNT-Si hybrid film deposited on ethanol-treated CNTs.

The thickness of as grown CNTs was characterized by SEM. For the synthesis of vertical aligned CNTs on Cu substrates, 10-nm-thick Ta was used as diffusion barrier to suppress Cu diffusion [139, 140], and 15-nm-thick TiN was also used as underlayer, which enables Fe or Co to form particles by dewetting [141-143].

Figure 4-2 shows the cross-sectional view SEM images of the vertical aligned CNTs, grown by Fe catalyst for different growth time with a partial pressure of 0.2 Torr C_2H_2 and 2 Torr H_2 at 700 °C. The height of CNTs varied from ~10 μm for 4.5 min, ~20 μm for 6 min, ~50 μm for 8 min, to ~110 μm for 30 min. Since the target height of CNTs was 15–20 μm , I used 6 min as the standard CVD time. Figure 4-3 shows the CNTs grown for 6 min, having a thickness of 15–20 μm and a density of

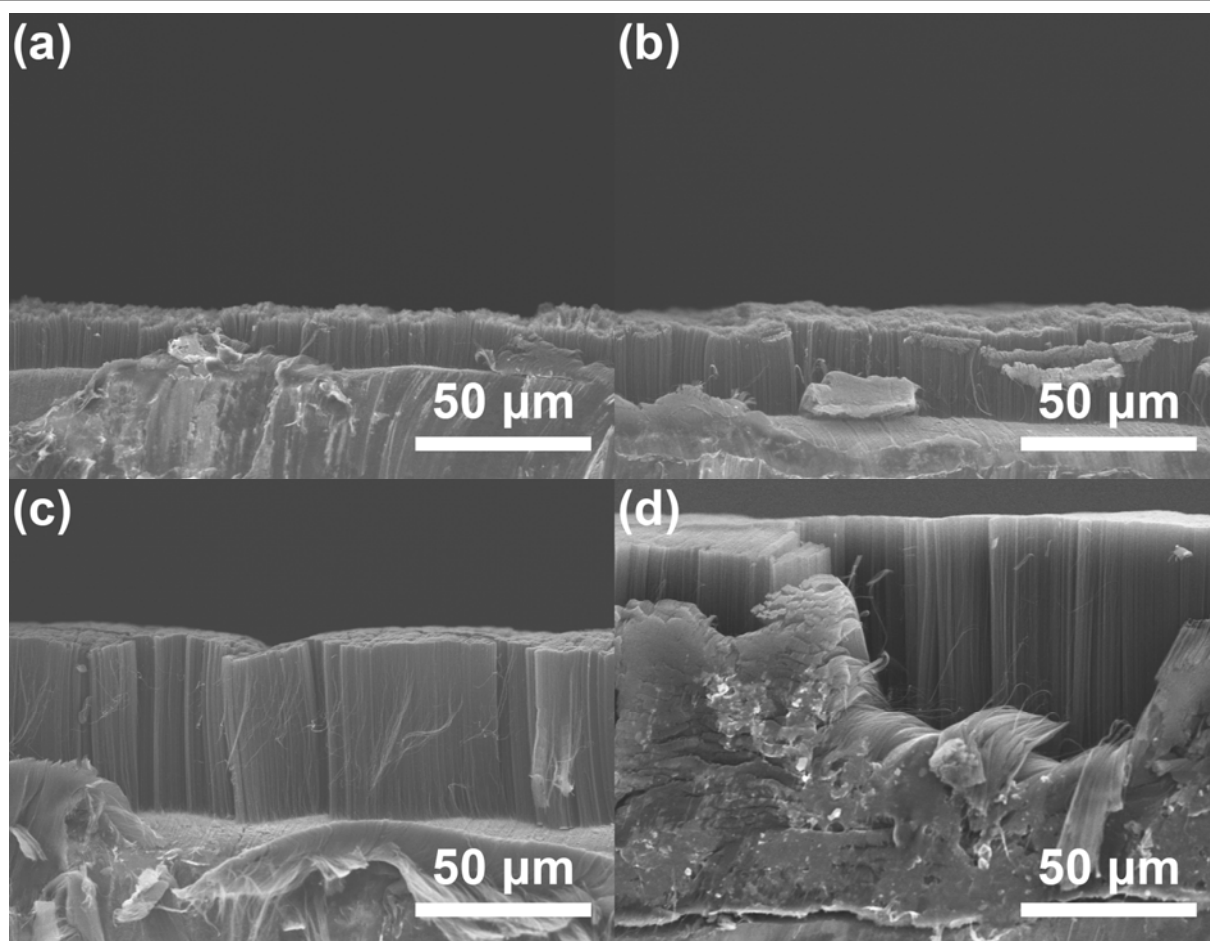


Figure 4-2 Cross-sectional view SEM images of vertical aligned CNTs grown by Fe catalyst for (a) 4.5 min, (b) 6 min, (c) 8 min and (d) 30 min.

$\sim 0.07 \text{ g cm}^{-3}$. All of the CNT arrays were aligned vertically to the substrate (Figure 4-3a), while CNTs were entangled with each other at the array top (Figure 4-3b–c).

In contrast, Figure 4-4 shows SEM images of self-organized CNTs as a result of capillary action. When liquid evaporates from CNT arrays, capillary force works on CNTs, which induces the CNT arrays to shrink, and finally results in such wall structure [144-148]. The wall structure also turns to

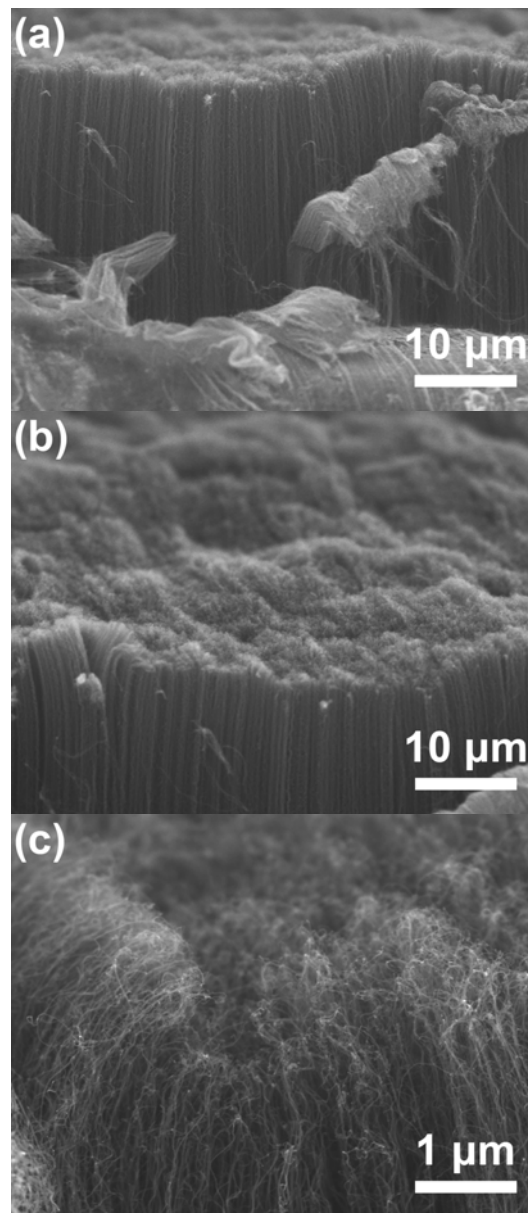


Figure 4-3 (a) Cross-sectional and (b) tilted-view SEM images of vertical aligned CNTs grown using Fe catalyst for 6 min. (c) High magnification of (b)

diverse 3-D structures by either density of CNTs or roughness of substrates. Since as-purchased Cu substrates had line-patterns of $\sim 1 \mu\text{m}$ in depth on its surface, CNT arrays had lower density near the patterns. Therefore, when the CNT arrays were treated with ethanol, the vertical aligned CNT arrays

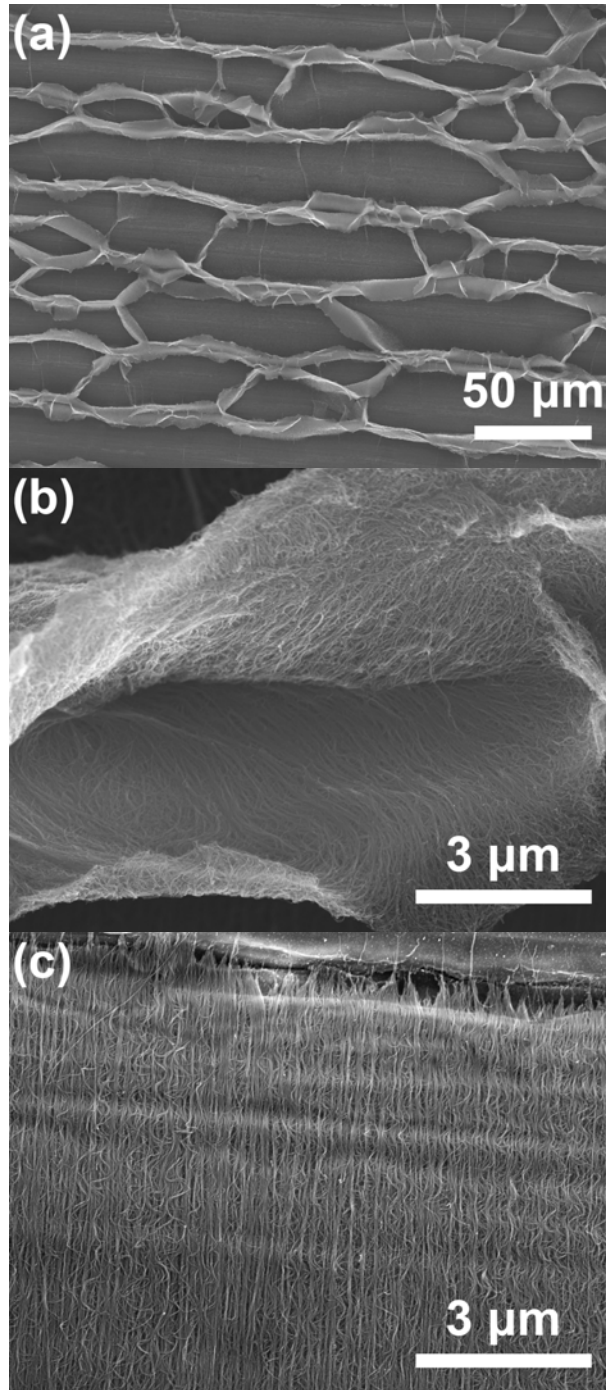


Figure 4-4 (a) A top-view SEM image of self-organized CNT arrays on Cu substrate prepared by ethanol-treatment. (b) High magnification SEM image of aligned CNT walls, and (c) cross-linked CNT walls.

aggregated by themselves to wall structures aligned parallel to the line patterns on the Cu substrate with some crosslinks at random positions.

Figure 4-5 shows the CNT-Si hybrid film deposited on as-grown CNTs by RVD in 2 min at T_{sub} of

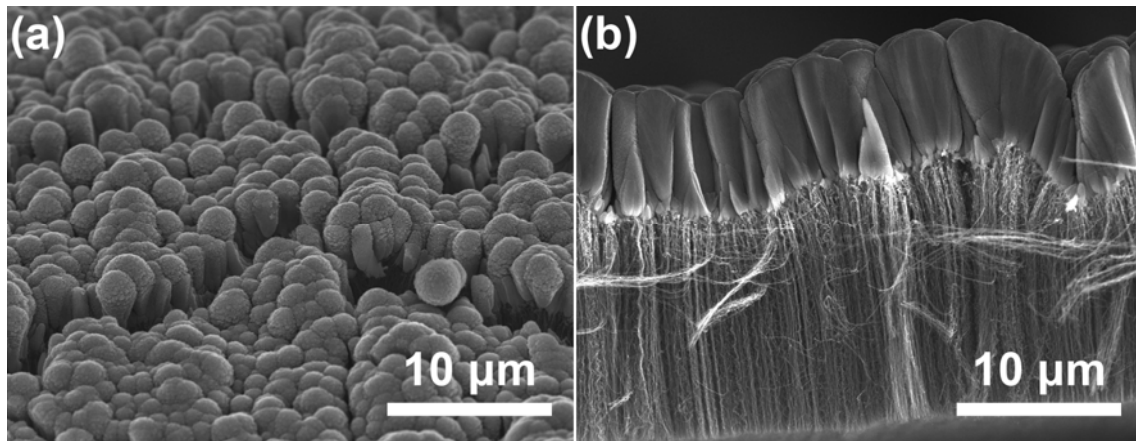


Figure 4-5 (a) Top and (b) cross-sectional view SEM images of the CNT-Si hybrid film deposited on as-grown CNTs at T_{sub} of 400 °C

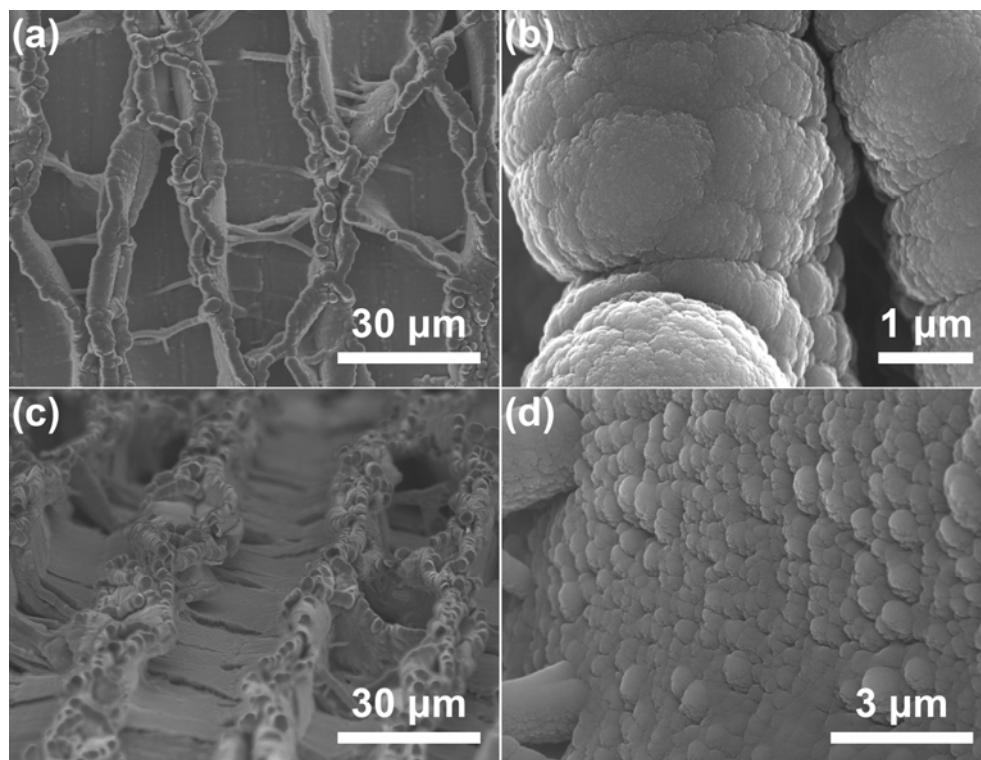


Figure 4-6 (a) A top-view SEM image of the CNT-Si core-shell wall film prepared using ethanol-treated CNT arrays by RVD at T_{sub} of 400 °C. (b) High magnification image of (a). (c) Tilted-view and (d) cross-sectional SEM images of the CNT-Si core-shell wall film.

400 °C. Si did not penetrate deep into the CNT arrays with a narrow interspace (~tens of nm) between CNT bundles, causing most Si to deposit on the array surface without covering CNTs inside the arrays. Resulting CNT–Si hybrid film looked like bean sprouts. In contrast, the Si deposition on ethanol-treated, self-organized CNT walls yielded wall-structured CNT–Si hybrid films with a wall height of ~20 μm and interspace of ~20 μm, as shown in Figure 4-6. Such 2-D microstructure of CNT–Si hybrid film supplies sufficient space for Si to change in volume during charge-discharge cycles although the interspacing should be controlled further. Furthermore, both higher areal density of 0.8–1.7 mg_{Si} cm⁻² and short-fabrication time of 2 min suggest the high potential of this Si-base film for practical use.

4.4 Cycle performance of CNT–Si hybrid film

Figure 4-7 shows the electrochemical behavior of CNT–Si hybrid films deposited at $T_{\text{sub}} = 400$ °C and post-annealed at $T_{\text{an}} = 600$ °C. The initial charge capacities of CNT-Si films with Fe or Co catalysts for CNT growth reached 3822 and 3545 mAh g_{film}⁻¹, respectively, and the initial discharge capacities were 1980 and 1232 mAh g_{film}⁻¹, respectively (Figure 4-7a). The initial charge capacities were closed to that of the pure Si film deposited at $T_{\text{sub}} = 500$ °C (Figure 2-10a), while the initial discharge capacities were much smaller. The large irreversible capacity is attributed to Si consumption during the formation of solid electrolyte interphase (SEI), which was significant because of the large exposed surface area of 2-D wall-shape structure of CNT–Si film formed by CNT arrays. The CNT–Si film using Fe catalyst for CNT growth showed poor cycle performance for 10 cycles (Figure 4-7, and Figure 4-8a). As shown in Figure 4-7a, capacity dropped continuously even after 2nd charge. Finally, the discharge capacity at the 10th cycle decreased to 293 mAh g_{film}⁻¹. The CNT–Si film (Fe catalyst for CNT growth) peeled off the Cu substrate had a similar morphology as initial

film, without being completely pulverized (Figure 4-8b and c). At the region where the CNT–Si film was peeling off, SEM-EDX analysis detected Ti $K\alpha$, but did not detect Fe $K\alpha$, indicating Fe catalyst layer corroded at interface between CNT and Ti underlayer during electrochemical reaction, causing CNT–Si films peeled off. It is known that Fe can easily corrode when it is exposed to moisture and also even more quickly if the moisture is salt water like electrolyte of electrochemical cell [152].

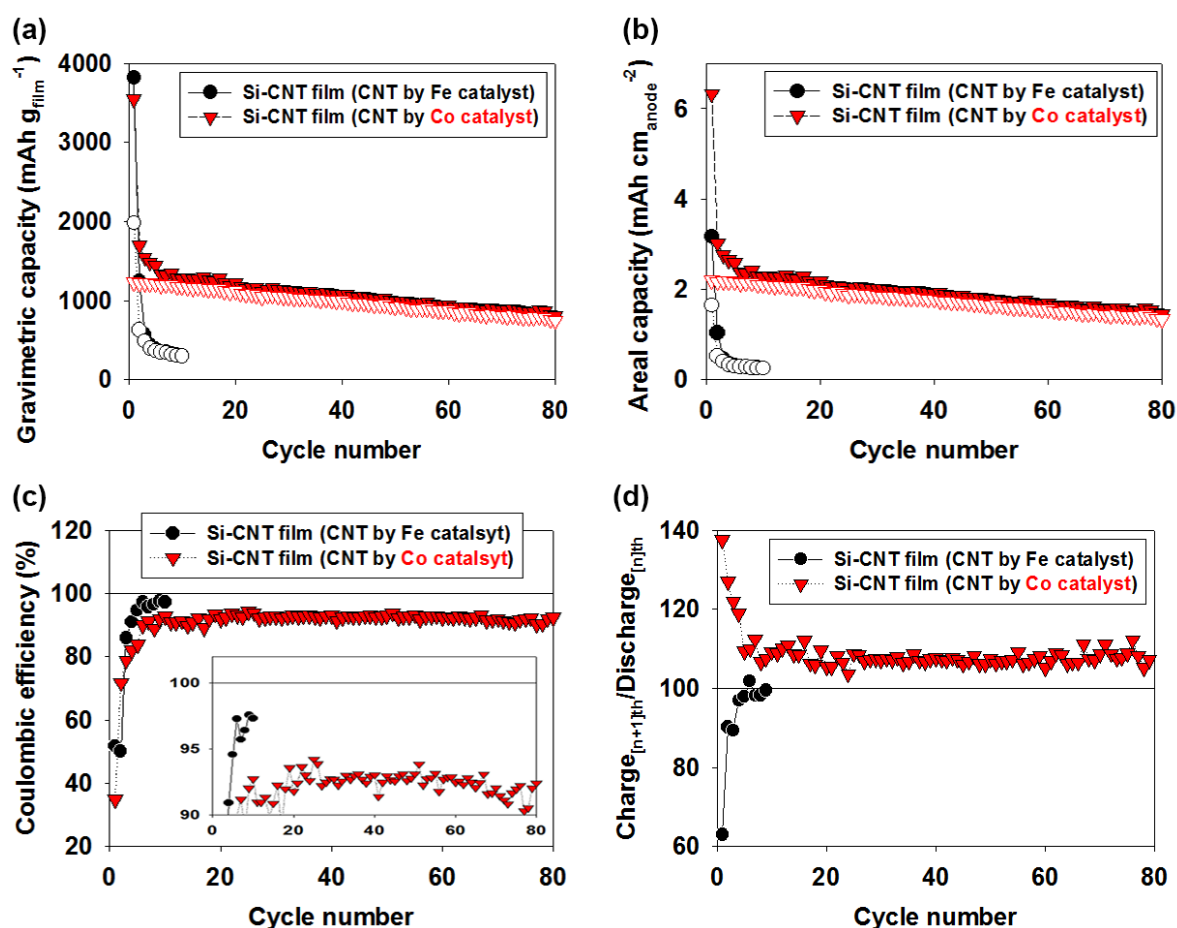


Figure 4-7 Electrochemical performance of CNT–Si films with CNT arrays grown by different catalysts (Fe and Co). Si was deposited by RVD at $T_{\text{sub}} = 400\text{ }^{\circ}\text{C}$ and post-annealed at $T_{\text{an}} = 600\text{ }^{\circ}\text{C}$. (a) Gravimetric capacity, (b) areal capacity, (c) Coulombic efficiency, and (d) capacity ratio of $\text{charge}_{[n+1]\text{th}}/\text{discharge}_{[n]\text{th}}$.

In contrast, the CNT–Si hybrid film (Co catalyst for CNT growth) showed better cycle performance with a discharge capacity of 749 mAh g_{film}⁻¹ and a capacity retention of 60% at the 80th cycle, indicating that the Co catalyst layer resisted against corrosion by the electrolyte of 1 M LiClO₄ in a

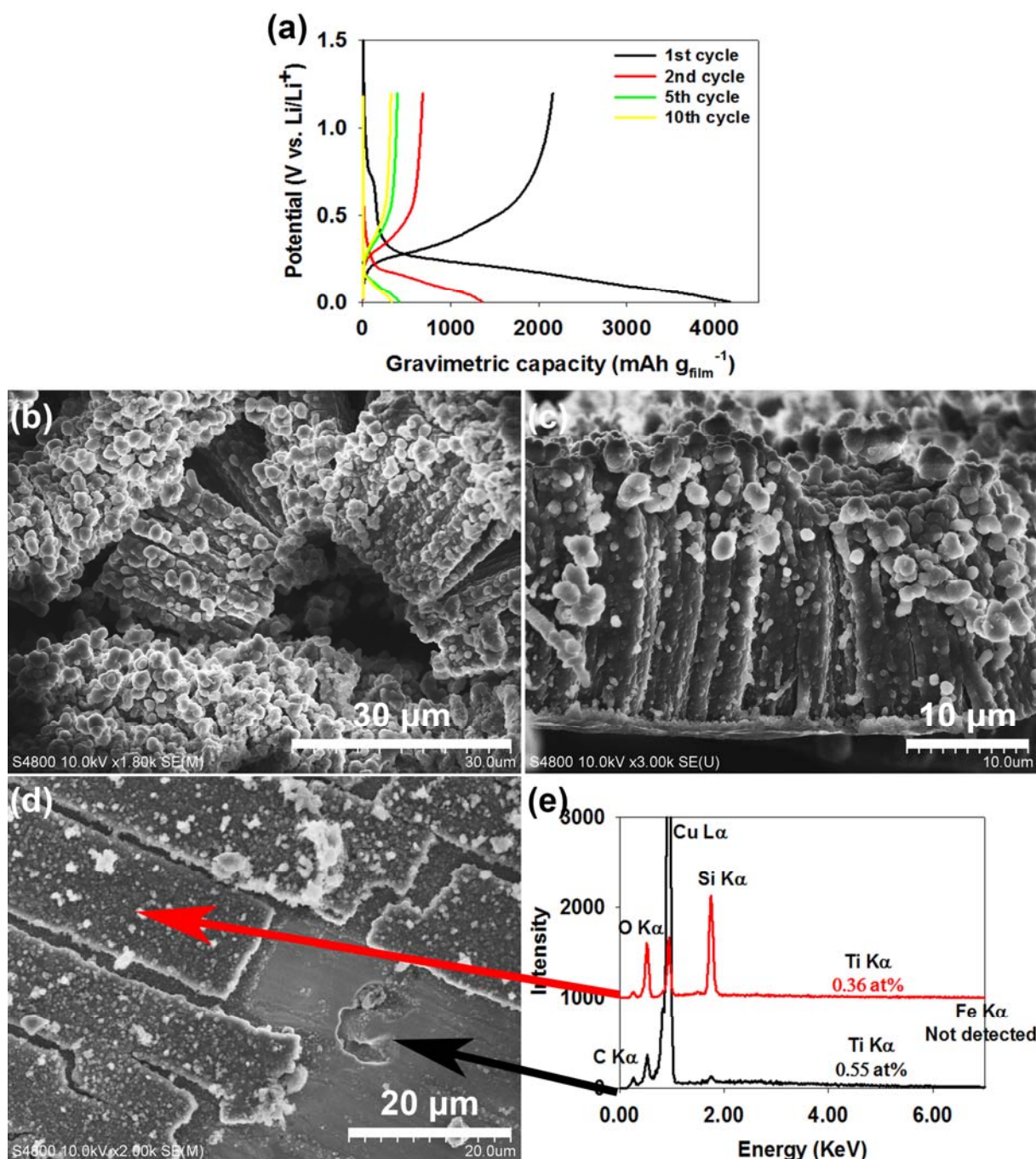


Figure 4-8 (a) Voltage-capacity profiles, (b) tilted-view, (c) cross-sectional view, and (d) top-view of SEM image of CNT–Si film with Fe catalyst for CNT growth after 80th discharge. Cycle test was performed at the rate of 0.4 A g⁻¹. (e) SEM-EDX analysis results for the CNT–Si film after 80th discharge.

1:1 (v/v) the mixture of EC and PC. It is reported that the use of PC causes exfoliation of graphene layers in graphitic carbons via solvent co-intercalation, resulting in its reduction and release of propylene gas [153, 154]. The Coulombic efficiency of CNT–Si film (Co catalyst for CNT growth) was ~92% at the 10th cycle, and remained at ~92% for 80 cycles, which is worse than previous our Si-base films. The capacity ratio of the [n+1]th charge over the [n]th discharge for the films was above 107% at the 80th cycle, suggesting that continuous SEI formation due to incomplete formation of stable SEI layer in the early cycles, possibly because of CNTs. The areal capacity ($\text{mAh cm}_{\text{anode}}^{-2}$) of the CNT–Si film is highly important factor for practical battery device. The areal capacity of the CNT–Si film (Co catalyst for CNT growth) was $1.33 \text{ mAh cm}_{\text{anode}}^{-2}$ at the 80th cycle, which is higher than previously reported values for Si nanomaterials ($0.2\text{--}0.4 \text{ mAh cm}_{\text{anode}}^{-2}$ [79, 88]) and porous Si–Cu films by ourselves ($0.96 \text{ mAh cm}_{\text{anode}}^{-2}$ [155]).

4.5 Conclusions

I successfully fabricated wall-shaped CNT–Si core-shell hybrid films by Si deposition on capillary force-assisted CNT arrays. In case of Si deposition on as-grown CNT arrays, the arrays held Si mostly on their top. Whereas ethanol-treated CNT arrays, they were well covered with Si. The areal density of CNT–Si hybrid film was $0.8\text{--}1.7 \text{ mg cm}^{-2}$. The CNT–Si hybrid film by Fe catalyst for CNT growth suffered from the corrosion of Fe during electrochemical reaction, resulting in delamination of film at the interface between the CNT–Si film and Ti underlayer. On the other hand, the CNT–Si hybrid film by Co catalyst for CNT growth were more stable with a gravimetric capacity of $749 \text{ mAh g}_{\text{film}}^{-1}$ and an areal capacity of $1.33 \text{ mAh cm}_{\text{anode}}^{-2}$ at the 80th cycle. Although further improvement such by controlling the wall distance, and by optimization of electrolyte and electrolyte additives is required, the CNT–Si films by RVD within 2 min are attractive for practical use in lithium secondary batteries when we consider their quick and simple fabrication using inexpensive source materials.

Chapter 5 – Conclusions and perspectives

Targeting at lithium secondary battery anodes, porous Si-base films were fabricated on Cu current collectors through a simple process, called RVD, without using any binder and conductive additives. The RVD process, making 3–14-micrometer-thick Si-base films, features rapid Si deposition in 1 min, which is proper to large-scale, low-cost production for practical battery devices. The porous Si-base films had wall-shaped structure along the line patterns on Cu substrate surface, and the porosity was simply controlled by changing substrate temperature, T_{sub} . The adhesion between Si-base films and Cu substrates was improved by pretreatment of Cu substrate surface through UV-O₃ exposure followed by annealing under H₂ flow and by development of gradient Si-Cu layers with sub- μm thickness by post-annealing and with several- μm thickness (entire film) by co-deposition of Si with Cu. Especially, the co-deposition methods proposed in this study remarkably enhanced the adhesion at the interface between the Si-Cu films and Cu substrates without crystallizing and densifying the Si-base films. The films deposited using a Si source with 5 wt% Cu at T_{sub} of 100 °C exhibited the highest electrochemical performance of fairly high initial charge and discharge capacities of 3425 mAh g_{film}⁻¹ and 2073 mAh g_{film}⁻¹, respectively, and capacity retention of 60% with a discharge capacity of 1250 mAh g_{film}⁻¹ at the 100th cycle. The amorphous phase with micrometer-thick Si/Cu gradient and high porosity of ~30% built in the Si-Cu film realized such high capacities for fairly thick Si-Cu films. Furthermore, the volumetric capacity of ~1900 mAh cm_{film}⁻³ and areal capacity of

$\sim 0.95 \text{ mAh cm}_{\text{anode}}^{-2}$ at the 100th cycle approaches to the value of practical lithium ion battery anodes. In addition, when the RVD method is applied to the capillary force-assisted self-organized CNT arrays, RVD yielded wall-structured CNT-Si hybrid films with an areal capacity of $1.33 \text{ mAh cm}_{\text{anode}}^{-2}$ at the 80th cycle, which suggests an alternative method for making practically thick 3-dimensional Si-base films.

Rapid vapour deposition will be compatible with practical production of Si-base anodes in terms of its high deposition rate, simple process, and inexpensive and safe source materials, but further research is needed to improve the anode performance of Si-base films such as optimum cut-off potential during lithation/delithiation and electrolyte additives like vinylene carbonate for more stable SEI layer to meet the requirements for practical devices.

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Appendix

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