

## Electrochemical Synthesis

### Electrochemical Preparation of Silicon and Its Alloys from Solid Oxides in Molten Calcium Chloride\*\*

Xianbo Jin, Pei Gao, Dihua Wang, Xiaohong Hu, and George Z. Chen\*

Silicon plays essential roles in the fabrication of solar cells, silicon chips, optical fibres, silicones, and is important as an element in lighter and stronger alloys, as well as hundreds of other advanced applications.<sup>[1,2]</sup> The industrial production of silicon is at present mainly through the carbothermic reduction of SiO<sub>2</sub> at 1700 °C, in which the oxygen is removed by the generation of CO<sub>2</sub>.<sup>[3,4]</sup> This old-fashioned charcoal technology should be replaced by a more advanced process from the viewpoint of an environmentalist, in that the earth's climate is gradually becoming worse owing to the emission of greenhouse gases.<sup>[5]</sup> The world production of silicon in 2002 was

about  $4.1 \times 10^6$  tonnes,<sup>[6]</sup> equivalent to about  $6.5 \times 10^6$  tonnes of CO<sub>2</sub> entering the atmosphere. A desired method of producing silicon may be the electrochemical reduction of SiO<sub>2</sub> in which the use of carbon reductant can be avoided and hence less environmental damage. In fact, electrolytic production of silicon began in 1854,<sup>[7]</sup> and silicon of 99.999 % in purity was later claimed<sup>[8]</sup> upon electrolysis of fluorosilicates in molten fluorides. DeMattei et al. in 1982 suggested<sup>[9]</sup> that the ideal raw material for silicon production should be SiO<sub>2</sub>, and silicon metal with a purity of 99.97 % was produced in the BaO/SiO<sub>2</sub>/BaF<sub>2</sub> system with the cell temperature being around 1450 °C. Recently, in molten CaCl<sub>2</sub> at about 900 °C, electrochemical deoxygenation of metals was investigated.<sup>[10–12]</sup> It was reported more recently<sup>[13–17]</sup> that when solid metal oxides are made into an electrode, regardless of their electron conductivities, they can be electroreduced (or electrodeoxidised) directly to the respective metals or alloys. This leads to a great possibility for the production of silicon directly from solid SiO<sub>2</sub>. However, until now, there is only one relevant report that concerns the electroreduction of solid SiO<sub>2</sub>, and only partial reduction of the SiO<sub>2</sub> electrode (quartz plate) was achieved.<sup>[18]</sup>

Herein, we report the fast, complete, and low energy electroreduction of a porous electrode prepared from SiO<sub>2</sub> powder. We also report an experimental observation that might indicate the existence of an optimal thickness of an insulating solid oxide, for example, SiO<sub>2</sub>, through which the electroreduction can progress “quickly” to completion. Furthermore, two examples of silicon alloys have also been produced by the same electroreduction method.

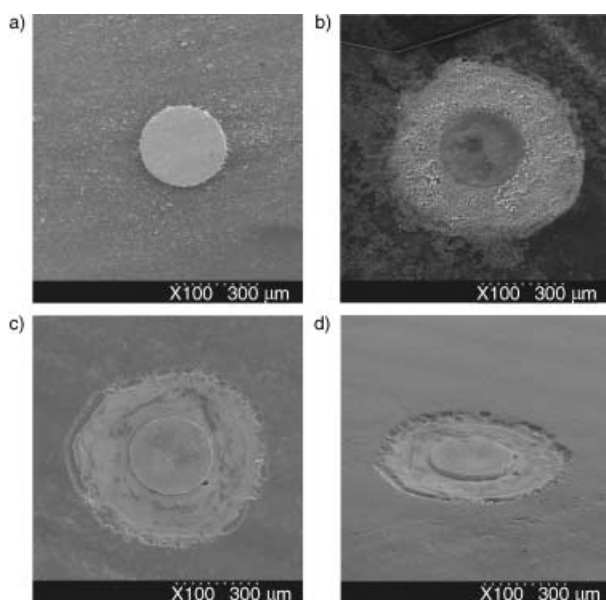
Recently, Nohira et al. succeeded in removing oxygen from the surfaces of solid SiO<sub>2</sub> plates (quartz) in a molten CaCl<sub>2</sub> electrolyte at 850 °C.<sup>[18]</sup> Similar results were also obtained in our laboratory by using a different electrode design. In the experiments, a tungsten wire of 300 µm in diameter was sealed in a quartz tube by using a gas flame. The end face of the tungsten wire was revealed by grinding. The SEM image showed a very intimate contact between the quartz and the W wire (Figure 1a). This W–SiO<sub>2</sub> electrode was then inserted into molten CaCl<sub>2</sub> at 850 °C and cyclic voltammetry was carried out by using a Pt wire and a graphite rod as the pseudoreference and counter electrodes, respectively. As shown in Figure 2a, the reduction of SiO<sub>2</sub> began at –0.85 V, which led to a sharp increase in the current that reached a peak of about –12.5 mA at –1.0 V. The current then went through a slightly inclined plateau on which a couple of small peaks can be seen (Figure 2a). These peaks are thought to correspond to the formation of the calcium and silicon compounds.<sup>[18,19]</sup> Upon reversing the potential sweep the current formed a typical stripping peak. When the sweep reached more positive potentials, a large poorly defined anodic peak occurred at about –0.7 V, thus suggesting that the reoxidation process may involve both pure and compounded forms of Si. A further study of the reaction mechanism is ongoing.

Figure 1a–d shows the SEM images of the W–SiO<sub>2</sub> electrode before and after the first potential sweep cycle. After this sweep the central W disc was surrounded by a ring of porous product of about 200 µm in breadth (Figure 1b).

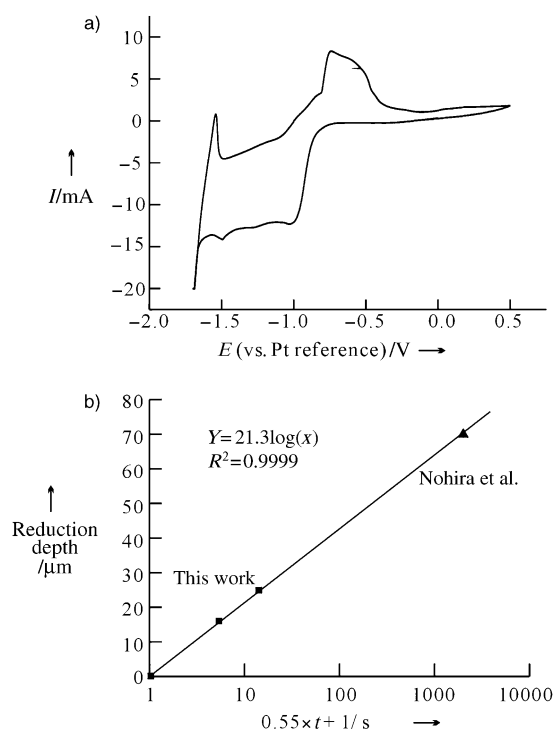
[\*] Dr. X. B. Jin, P. Gao, Dr. D. H. Wang, Dr. X. H. Hu  
College of Chemistry and Molecular Science  
Wuhan University, Wuhan, 430072 (P. R. China)  
Fax: (+86) 27-87210319  
E-mail: mel@chem.whu.edu.cn

Dr. G. Z. Chen  
Specially Invited Professor  
College of Chemistry and Molecular Science  
Wuhan University, Wuhan, 430 072 (P.R. China)  
and  
School of Chemical, Environmental and Mining Engineering  
University of Nottingham  
University Park, Nottingham NG7 2RD (UK)  
Fax: (+44) 115-9514171  
E-mail: george.chen@nottingham.ac.uk

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**Figure 1.** SEM images of the W-SiO<sub>2</sub> electrode a) before and b) after one cycle of potential sweep between 0.5 and -1.7 V, and c) after washing the electrode shown in (b) in an ultrasonic water bath. d) A side view of the situation in (c).



**Figure 2.** a) Cyclic voltammogram of the W-SiO<sub>2</sub> electrode in molten CaCl<sub>2</sub> (200 mVs<sup>-1</sup>, 850 °C), b) the relationship between the depth and time of reduction of solid SiO<sub>2</sub>.

The product was removed by immersing the electrode in an ultrasonic water bath, thus resulting in a circular cave in the shape of a flat-bottom pan. By using the depth (16  $\mu\text{m}$ ), the top (200  $\mu\text{m}$ ) and bottom breadths (150  $\mu\text{m}$ ) of the cave (Figure 1 c, d), the volume of the reacted SiO<sub>2</sub> was estimated

to be about  $4.2 \times 10^{-6} \text{ cm}^3$ . Considering the density of quartz ( $2.2 \text{ g cm}^{-3}$ )<sup>[19]</sup> and the molecular mass of SiO<sub>2</sub> (60 g/mol), we estimate that about  $1.54 \times 10^{-7}$  moles of SiO<sub>2</sub> were reduced, which is equivalent to a total charge of about 59.4 mC if four electrons were transferred. This value can be compared with the total charge of about 61.5 mC passed during the negative and positive sweep between -0.85 V and -1.7 V (Figure 2 a). The good agreement between the two charge values is strong evidence that the reduction product between -0.85 and -1.7 V was mainly silicon.

By using the average current of the plateau between -1.0 V and -1.7 V, the reaction rate was estimated to be higher than  $80 \text{ A cm}^{-2}$  with reference to the revealed side surface of the tungsten wire, which is  $1.41 \times 10^{-4} \text{ cm}^2$  (Figure 1 d). This value is even higher than that usually observed in the electrodeposition of metals from liquid electrolytes. Such a high current density suggests a great potential of the electroreduction of solid SiO<sub>2</sub> for the industrial production of silicon. However, it should be pointed out that this high reaction rate was observed at the very beginning of the electroreduction, and the affected volume of the quartz was very small. The question is whether this high reaction rate could be achieved if the volume of quartz were significantly increased.

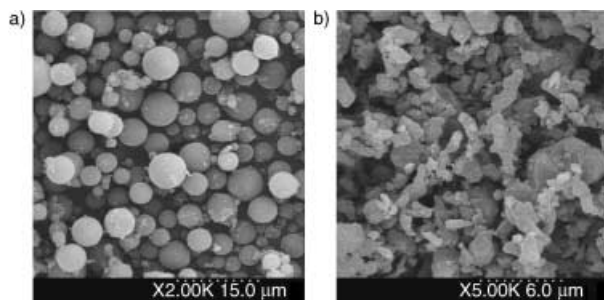
It was proposed that the electroreduction of insulating solid oxides could proceed at the three-phase boundary linking the conductor, oxide and electrolyte.<sup>[14]</sup> This principle was advocated and demonstrated with an elegantly designed experiment for the reduction of a quartz plate.<sup>[18]</sup> In our experiment, before reduction had occurred, there was a W/SiO<sub>2</sub>/CaCl<sub>2</sub> three-phase boundary at the edge of the central W disc (end of wire). Upon electron transfer, the SiO<sub>2</sub> next to the W disc was converted into silicon which has a reasonably high electronic conductivity ( $50 \Omega^{-1} \text{ cm}^{-1}$ ).<sup>[20]</sup> Therefore, further reduction may occur at the newly formed Si/SiO<sub>2</sub>/CaCl<sub>2</sub> three-phase boundary. In theory, the continuation of this change should lead to the propagation of the three-phase boundary in a 3D manner. However, Figure 1 c, d tells a different story and shows a transverse reaction width of 200  $\mu\text{m}$ , but a reaction depth of only 16  $\mu\text{m}$ . While this is an indication of a greater reaction resistance along the depth direction than the transverse direction, the phenomenon may be broadly understood by the accessibility of the molten salt to the reaction sites that are directly in touch with the molten salt along the transverse direction, but in a different situation along the depth direction.

However, if the volume remains the same, the removal of oxygen from SiO<sub>2</sub> should result in a porosity of about 56.2 % if the density difference between SiO<sub>2</sub> and Si is considered (2.2 and  $2.33 \text{ g cm}^{-3}$ ,<sup>[19]</sup> respectively). Therefore, the transverse propagation of the Si/SiO<sub>2</sub>/CaCl<sub>2</sub> three-phase boundary opens up the solid phase by creating pores/channels for the molten salt to access the SiO<sub>2</sub> phase behind the porous Si layer.<sup>[18]</sup> This explanation agrees well with the porous product shown in Figure 1 b. Nonetheless, it can be reasonably considered that the mass transfer within the formed porous silicon layer is slower than that at the SiO<sub>2</sub>/molten salt interface, thus leading to a slower reaction rate along the depth direction than the transverse direction.

The observation of a flat bottom in the cave (Figure 1c) suggests that the electroreduction of the insulating  $\text{SiO}_2$  might be limited to a certain distance in the depth direction, which is less dependent on the time of the applied potential. This depth limit as shown in Figure 1d may be mainly attributed to the limit of mass transfer through the porous silicon layer formed on the surface.

We would like to highlight the practical significance of this finding of different reaction rates along the transverse and depth directions. For industrial production, the use of bulk solid quartz as the electrode is obviously not practical. Instead, a process is likely to involve the use of readily available  $\text{SiO}_2$  powder to fabricate a porous  $\text{SiO}_2$  electrode, on the analogy of the reports by Chen and co-workers.<sup>[13,14]</sup> However, it can be anticipated that, if the oxide particles are too large, the same situation would occur to the individual oxide particles, as observed in our electroreduction of the bulk solid quartz electrode. That is, within a practically tolerable time, the reduction would take place only on the surface of the oxide particle with the inner part being unaffected. To the best of our knowledge, there has not been any report in the literature giving an accurate account, either experimental or theoretical, of the effect of powder particle size on the efficiency of the electro-reduction. On the basis of our findings and that of Nohira et al.,<sup>[18]</sup> a logarithm relation could be approximated between the reduction depth and time (see Figure 2b). It can then be estimated that, for example, the complete reduction takes place within 0.1 s, 1.3 s and 403 s of three  $\text{SiO}_2$  particles of 1  $\mu\text{m}$ , 10  $\mu\text{m}$  and 100  $\mu\text{m}$  in diameter, respectively.

The above estimation was found to agree broadly with the result from the electrolysis of the  $\text{SiO}_2$  powder, with particle sizes ranging from 2 to 7  $\mu\text{m}$  (Figure 3a). The powder was manually pressed into a porous pellet (1.3 cm in diameter,

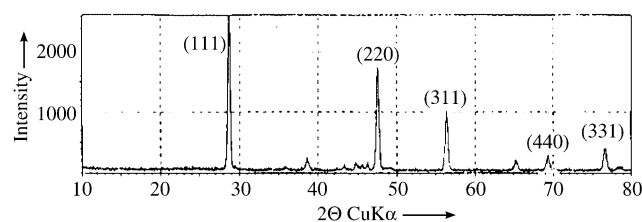


**Figure 3.** SEM images showing a) the  $\text{SiO}_2$  powder electroreduced in molten  $\text{CaCl}_2$  (2.8 V, 850 °C, 4 hrs) into b) the Si powder.

0.5 cm in thickness, 1.5 g in weight). A manually operated press was used for making the pellets and hence the actual pressure was unknown but was sufficient to form a strong pellet. After sintering at 1000 °C, the pellet was pressed and sandwiched between two porous nickel foils. Then, constant-voltage (2.8 V) electrolysis was performed, under argon, on the assembled cathode in molten  $\text{CaCl}_2$  at 850 °C with a graphite rod as the anode. The recorded current was very high (about 4 A) during the first few minutes but declined thereafter (within 2 h) to the background value,  $\approx 0.2$  A.

After electrolysis, which usually lasted for 4 h to ensure complete reduction, the cathode was removed from the molten salt. Upon washing the cathode (pellets) in distilled water in an ultrasonic bath, a grey powder was collected.

The obtained powder was dried and analyzed by XRD (Figure 4), thus confirming the product to be dominated by pure silicon. In addition to the pure silicon phase, the XRD spectrum also exhibited some small but distinguishable sundry peaks which are likely due to  $\text{MgSiO}_3$ . This was tracked down to an  $\text{MgCl}_2$  impurity (about 0.3 %) contained in the  $\text{CaCl}_2$  used in this work.



**Figure 4.** The XRD spectrum of the Si powder obtained from the electroreduction of pellets of the  $\text{SiO}_2$  powder in  $\text{CaCl}_2$  molten salts.

When inspected under SEM (Figure 3b), the silicon powder particles were observed to be 1–3  $\mu\text{m}$  in diameter, which is in accordance with the theoretical prediction from the density difference between Si and  $\text{SiO}_2$ . Because both  $\text{SiO}_2$  and Si are almost insoluble in  $\text{CaCl}_2$ , the output rate should be very high. These results prove that the silicon powder can be mass produced from the  $\text{SiO}_2$  powder by the electroreduction method. It is worth mentioning that in our experiments complete reduction of the  $\text{SiO}_2$  pellets took a much shorter time than the reduction of  $\text{TiO}_2$  ( $\approx 15$  h) under similar conditions as reported in literature.<sup>[13]</sup> This difference may be due to the fact that oxygen is highly soluble in solid titanium but not in solid silicon.

By mixing powders of different metal oxides, alloys can be produced by the electroreduction method.<sup>[13,14,21]</sup> Because fine powders of oxides can be prepared easily and mixed uniformly, the electroreduction of such a mixture leads to an alloy, the composition of which is precisely controlled. Indeed, in our experiments, Si–Fe and Si–Cr alloy powders with a particle size of 2–3  $\mu\text{m}$  were also prepared.

In conclusion, we have demonstrated, for the first time, that porous pellets of the  $\text{SiO}_2$  powder or its mixture with other metal oxide powders can be electroreduced to pure silicon or the respective silicon alloy powders in molten  $\text{CaCl}_2$ . Furthermore, cyclic voltammetry revealed that the electroreduction of  $\text{SiO}_2$  could proceed very quickly, with the current density reaching beyond  $80 \text{ A cm}^{-2}$  with reference to the surface area of the conducting substrate. Although such a high current density may be difficult to achieve on an industrial scale, it may be used as a fundamental reference for the design of an industrial process for the mass production of silicon powder by electrolysis. Furthermore, we found that the relation between the reduction depth and time of solid  $\text{SiO}_2$  may be approximated by the logarithm law. This finding may be used to select the particles sizes of the  $\text{SiO}_2$  powder so

that complete reduction of the electrode prepared from the  $\text{SiO}_2$  powder can be achieved within a practically tolerable time. This work shows that when the particle size is in the range of 2–7  $\mu\text{m}$ , complete reduction of a 0.5 mm thick porous pellet at 2.8 V and 850 °C could be less than 4 h with the energy consumption being within 13 kWh (kg of Si) $^{-1}$  which is expected to decrease further upon optimization. This can be compared with the energy consumption, 13–16 kWh (kg of Si) $^{-1}$ , of the carbothermic method used by the industry in China.<sup>[22]</sup>

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