Facile preparation of superhydrophobic surface with high adhesive forces based carbon/silica composite films

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Abstract. Glass substrates modified by carbon/silica composites are fabricated through a two-step process for the preparation of a superhydrophobic surface (water contact angle $\geq 150^{\circ}$). Carbon nanoparticles were first prepared through a deposition process on glass using a hydrothermal synthesis route, then the glass was modified by SiO₂ using the hydrolysis reaction of tetraethylorthosilicate at room temperature. It is not only a facile method to create a superhydrophobic surface, but also helps to form a multi-functional surface with high adhesive forces.

Keywords. Superhydrophobic surface; high adhesive forces; hydrothermal synthesis route; hydrolysis reaction.

1. Introduction

In recent years, bionics has attracted considerable attention for its great prospect. Shining drops of rain water forming a nearly perfect spherical shape on lotus leaves, which can be rolled off easily, is one of the most beautiful wonders of nature. This natural phenomenon is known as lotus effect (Zhai et al 2004; Cha et al 2010; Hipp et al 2010) and is considered as the basic study of superhydrophobic behaviour on a surface. Now the study of superhydrophobic behaviour of various surfaces is not only part of fundamental research, but also used in practical applications in many fields, especially in daily life (Yang et al 2007; Zhang et al 2011; Ogihara et al 2012), industry (Li et al 2008; Xu et al 2011) and agriculture (Sheng and Zhang 2009; Wang et al 2011). Cassie and Baxter (1944) as well as Wenzel (1936) proposed two mathematical models to explain the wetting phenomena on rough surfaces that leads to superhydrophobic property (Erbil and Cansoy 2009; Su et al 2009). Usually, any surface with water contact angle larger than 150° can be defined as a superhydrophobic surface (Tuberquia et al 2010). Various materials and methods were tried out for the fabrication of superhydrophobic surfaces. Generally, these approaches to obtain superhydrophobicity can be divided into two groups, one of them is to achieve superhydrophobicity by changing the structure and surface energy of substrates. For example, Meng et al (2008) created superamphiphobic surfaces on common engineering metals by immersing them into perfluorocarboxylic acid and keeping for a few hours. Ishizaki and Sakamoto (2011) prepared colour-tuned superhydrophobic magnesium alloy with corrosion resistance. Xiu *et al* (2008) obtained superhydrophobic surfaces on silicon surfaces by hierarchical etching. Shen *et al* (2010) employed a flower-like silicon particulate film by chemical etching and obtained superhydrophobicity. Another method is by coating superhydrophobic surface on the substrates such as Tsai and Lee (2007) who employed a raspberry-like particulate film prepared by layer-by-layer deposition and produced superhydrophobic surfaces. Yildirim *et al* (2011) prepared a superhydrophobic surface by coating fluorinated mesoporous silica nanoparticles. Brassard *et al* (2011) prepared superhydrophobic thin films by synthesizing monodisperse fluorinated silica nanoparticles. Manca *et al* (2009) obtained superhydrophobic surfaces by trimethylsilanized silica nanoparticles based sol-gel processing.

Recently, various methods were used to obtain superhydrophobic surfaces, such as template method, spray painting, self-assemble way and etching method. However, most of them employed just only one material to create a superhydrophobic surfaces, while few employed composites to create superhydrophobic surfaces. Here, we report the design of superhydrophobic surface using high adhesive forces by means of employing C/SiO₂ composites, which is simple and environmental friendly. Especially, the high adhesive forces enhance water-retaining capacity of the surface which is very important for long-distance transport of water. C nanoparticles were prepared through a deposition process on the glass using a hydrothermal synthesis route, then tetraethylorthosilicate (TEOS) dissolved in isopropanol (IPA) forming homogeneous solution, was dropped onto the glass modified by C nanostructures and kept in a container for a few days. C/SiO₂ composites were fabricated during the hydrolysis reaction of TEOS. In order to reduce

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surface energy, the glass was further treated with fluorosilicone. After that, the fabricated coatings exhibited superhydrophobic property because of their dual-size rough surface and low surface energy.

2. Experimental

2.1 Materials

All chemicals were of analytical grade and used as received. Glucose was purchased from Tianjin Henxing Chemical Reagent Manufacturing Co., Ltd. TEOS was obtained from Tianjin Kermal Chemical Reagent Co. Ltd. (IPA) was purchased from Hangzhou Gaojing Fine Chemicals Co. Ltd. 1H,1H,2H,2H-Perfluordecyltrimethoxysilane (>97 wt%) was obtained from Nanjing Daoning Chemicals Co. Ltd.

2.2 Methods

A quantity of 6 g of glucose was added to 60 mL of deionized water by stirring for a few minutes, forming homogeneous solution. The resulting mixture was transferred to and sealed in a Teflon-lined autoclave, and a piece of glass was also inserted into the autoclave. The autoclave was sealed and maintained at 160 °C for 6 h. After the autoclave was cooled down to room temperature naturally, the glass was collected and washed with deionized water carefully, and dried for 12 h. Then TEOS was dissolved in IPA with different volume ratios forming homogeneous solution, which was dropped onto the glass modified by C nanostructures and kept in a container for a few days. The C/SiO₂ composites were fabricated during the hydrolysis reaction of TEOS, followed by perfluorosilane deposition to obtain high water-repellance.

2.3 Characterization

Microstructures of SiO_2 and C/SiO_2 composites were investigated by ULTRA-55 field-emission scanning electron

microscopy (FE–SEM) and the Fourier transform infrared (FT–IR) spectra were recorded on a Nicolet 5700 spectrophotometer. The wettability of the as-prepared C/SiO₂ composites films was analysed by measurement of the water angles using SL200B contact angle system at ambient temperature. Water droplets were dropped carefully onto the samples. The average contact angle was obtained by measuring at three different positions of the same sample, the accuracy of measurement was $\pm 1^{\circ}$.

3. Results and discussion

Figure 1 shows FE–SEM images of the resulting surface after the two-step process. In contrast to network-like SiO_2 nanostructures which have single particle chains (figure 1b), the network-like C/SiO₂ composites (figure 1a) appear more rough.

FT-IR analyses were carried out on thin films deposited on the glass to provide a description of silica layer covering the carbon particles. The spectrum of C nanoparticles in figure 2(c) shows a broad band with a maximum breadth of around 3365 cm^{-1} , which is attributed to the O–H stretching. A small peak appearing around 2925 cm⁻¹ is attributed to the stretching vibration of C-H bond in the form of CH₂ (Wan *et al* 2008). The stretching at 1282 cm⁻¹ is attributed to the C–O–C skeleton. The spectrum of C/SiO₂ composites in figure 2(b) shows a broad band with a maximum breadth of around 3425 cm⁻¹, which is attributed to the O-H stretching. A strong peak appears at 1100 cm⁻¹ due to the presence of Si-O-Si asymmetric bond stretching. Two small peaks appearing at 950 cm⁻¹ and 2926 cm⁻¹ are attributed to the stretching vibration of Si-OH and C-H bonds, respectively. The peak at 2700 cm^{-1} is attributed to the C–H stretching vibration. The spectrum of fluorinated C/SiO₂ composites in figure 2(a) shows a small peak appearing at 960 cm⁻¹, which is attributed to the stretching vibration of the C-F bond in the form of CF₃ (Brassard *et al* 2011).



Figure 1. FE–SEM images of resulting surface after different processes: glass modified by (**a**) C/SiO₂ composites and (**b**) SiO₂. All scale bars are of 2 μ m.

Water contact angle tests are carried out on thin films to obtain better understanding of wettability. 5 μ L water is dropped on the glass modified by SiO₂, which is due to the presence of hydrophilic Si–OH bonds and exhibits



Figure 2. FT–IR spectra of thin films prepared by (a) fluorinated C/SiO_2 composites, (b) C/SiO_2 composites and (c) C nanoparticles.

hydrophilic property. After treatment with fluorosilicone, its water contact angle (WCA) reaches 133° (the inset of figure 3(a)). The wettability of glass modified by C nanoparticles is similar to that of glass modified by SiO₂. However, when the glass is modified by C/SiO₂ composites, followed by treatment with fluorosilicone, its WCA is 152° (inset of figure 3(b)), which indicates that the surface of the glass has superhydrophobic property. In addition, as IPA keeps increasing, WCA increases first, and then decreases gradually. The possible reason for this phenomenon is the changing of surface roughness at different volume ratios of TEOS and IPA. As shown in figure 3, when the volume ratios of IPA and TEOS are 2:1 or 5:1, the surface exhibits superhydrophobic properties.

Furthermore, we also find that when the drop is turned upside-down, it still adheres to the surface. As shown in figure 4, the C/SiO₂ composite film modified by TEOS and IPA with volume ratio of 1:5 has strong adhesive force that is similar to a gecko's foot (Zeng *et al* 2009; Liu *et al* 2010; Bhushan 2012). A gecko's feet possess superhydrophobicity and high adhesive ability towards water, because they are covered with millions of well-aligned microscopic keratinous hairs called setae (20–70 mm in length and 3–7 mm in diameter), which are further split into hundreds of smaller nanoscale ends called spatulae (100–200 nm in diameter)



Figure 3. FE–SEM images of glasses modified by C/SiO₂ composites with different volume ratios of IPA and TEOS: 0 (a), 2 (b), 5 (c) and 10 (d); insets are WCA images. All scale bars are of 2 μ m.



Figure 4. A 5- μ L water drop hanging on micro-fluctuant area that is upside-down; left is WCA image, right is schematic of a 5- μ L water drop hanging on rough surface.

(Liu et al 2012). The possible reason for superhydrophobic surface with high adhesive force has been studied by many researchers such as Li and Shen (2011), Lai et al (2009) and Hong et al (2007). The carbon/silica composite films reported in this paper also exhibit strong adhesive forces. We suggest that the strong adhesion can be attributed to van der Waals force and negative pressure generated by the air trapped in a closed 'liquid-solid' system when the drop is turned upside-down. The micro-fluctuant surface and hair-like fluorosilicone result in a large contact area when a droplet is placed on it, thereby increasing the van der Waals force between the water droplet and surface. When the droplet is turned upside-down, the direction of gravity changes which leads to change in the surface energy of 'liquid-solid' and 'liquid-air' systems. Generally, the variation of surface energy results in negative pressure which is an important factor for the water droplet to adhere to the surface.

4. Conclusions

In summary, we designed a two-step process for the preparation of a superhydrophobic surface. A binary structure results from the combination of C nanoparticles and network-like SiO₂ nanostructures, after treatment with fluorosilicone. The obtained surface has two functional areas that present different wettabilities and high adhesive force that enhances the water-retaining capacity of the surface. This feature can be used in places where water has to be retained for a long time. The advantages of this methodology also include its simplicity and ease of control.

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