

Sequentially reinforced additive coating for transparent and durable superhydrophobic glass

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ABSTRACT

Now that there are various routes to prepare superhydrophobic surfaces for self-cleaning, anti-icing, liquid collecting and etc., attentions are moving towards low-cost upscaling route and increasing the reliability for actual applications. However, the required micro-nano structures for the superhydrophobicity are light scattering and very vulnerable to abrasion, which intrinsically conflicts with the transparency and durability, existing as the major barrier for commercialization of superhydrophobic glass. In this study, we present a novel sequentially reinforced additive coating (SRAC) process to realize robust and transparent micro-nano structured film with tough intergranular sintering. A benign aqueous-based ink with poly(furfuryl alcohol) (PFA) and silica species are carefully designed and sprayed on glass to enable self-phase separation and morphology construction. The coatings reach the static contact angle (SCA) for water over 166° and withstand a 6H pencil scratching, the cross-cut test and sands abrasion. Moreover, we also perform 90 days out-door performance test and the glass maintains superhydrophobicity with SCA of 154° . These results provide a low cost waterborne ink formula, and the high throughput and up-scalable SRAC process could be a convenient technology for fabrication large area robust superhydrophobic coatings.

Keywords: superhydrophobic glass; furfuryl alcohol; sequential reinforcement; durable; transparent

INTRODUCTION

Self-cleanable surfaces that enable anti-external contamination from either nature or humanity activities are extremely appealing for the upcoming intelligent buildings or vehicles, which could save billions of dollars in personnel and accessories costs to maintain the aesthetic transparency for windows.¹⁻⁵ One of the most desirable measure to realize this function is to construct the superhydrophobic surface coating that water droplets in-combine with the dirt could roll off.⁶⁻⁷ Resembling to the lotus leaf,⁸ which is admired as the model for biomimic synthesizing of superhydrophobic coating, fabrications of multi-leveled micro-nano structure are required.⁹⁻¹² The roughness (on the micro and nano-scales) enables the trapping of air under water droplets, thereby contributing to repelling of water droplet to form a static contact angle (SAC) over 150°. However, as-regarding to superhydrophobic glass, the required rough structures, intrinsically conflicts with the transparency and durability, because the textured part is light scattering and vulnerable to abrasion. This challenge inhibits obtaining high performance superhydrophobic glass and causes substantial barrier for commercial applications.¹³⁻¹⁴

Current reported methods for fabrication roughness to realize superhydrophobicity can be generally classified into two categories, the “top-down” and the “bottom-up” measures.¹⁵ The “top-down” techniques including laser ablation,¹⁶⁻¹⁷ etching,¹⁸⁻¹⁹ galvanic corrosion,²⁰ photolithography²¹⁻²² and etc.²³⁻²⁴ construct rough structures as the integrated parts of the substrate, contributing to the relatively high mechanical durability. While processability of these measures are typically dependent on

substrates materials, and their relatively low throughput is not accommodated to the square kilo-meters scale applications cost-effectively.²⁵⁻²⁷ On the other hand, the “bottom-up”^{9, 28-30} processes utilized pre-prepared nano/micro building blocks³¹⁻³² with various shapes³³⁻³⁶ such as rods, spheres, core-shells, random agglomerates and etc. to pack and establish the textured surfaces, showing considerable adaptability to the current low-cost industrial painting technologies. The “bottom-up” processing, in contrast to the “top-down” methods, requires additional design to construct tough connections between the substrate and the topographic structure to maintain the water repelling ability. Forming hybrid organic-inorganic bonded composites,^{30, 33, 37-41} such as polydimethylsiloxane (PDMS), epoxy-siloxane, thiol-ene resins, polymethyl methacrylate (PMMA), fluorinated-decyl polyhedral oligomeric silsequioxane (FD-POSS) and fluorinated alkyl silane (FAS), have emerged as effective measures to enhance durability for various surface coatings. However, organic reinforcement films are required to increase the bonding strength between the particles and substrate. Moreover, because of the low crystallinity, multi-components complex layer and varied refractive indexes, those coatings inevitably result in a problem of blocking of light. Transparency is normally sacrificed for toughness and superhydrophobicity in organic-inorganic coating.

Thus, a simple, economical and effective way to realize a large-scale production of superhydrophobic glass is crucial for commercial interest. We notice that 3-D printing (additive manufacturing) technology using discrete particles as the building blocks have been successfully applied to fabricate ceramics parts with various shapes and

microstructures.⁴²⁻⁴⁴ Most importantly, the inter-particles bonding can be enhanced after heat processing and able to endure extensive mechanical friction, tensile and pressure.⁴⁵⁻⁴⁷ In this report, we present a novel route to enhance mechanical strength of the textured film by mimic the additive manufacturing of ceramics, i.e. a sequentially reinforced additive coating (SRAC) process. Different from the additive manufactured parts with computer aided design macrostructures, the textured film was directly constructed in the additive coating of SRAC process with careful design of precursor. We realized a durable and transparent textured layer for superhydrophobic glass with the high toughness up to 6H and sustainable to ~500 g sand abrasion at height of 20 cm. Moreover, we performed a 90 days out-door performance test. The glass maintained superhydrophobic properties with contact angle of 154°. Moreover, this textured structure can be deposited on the glass by spraying the waterborne ink directly, which can be convenient for potential large-scale environmentally friendly and cost-effective application.

Experimental section

Materials

Tetraethyl orthosilicate (TEOS), concentrated sulfuric acid, hydrogen peroxide (30% AR), ethanol (99.7%), ammonia solution (25–28%), furfuryl alcohol (FA, 98.5%), polyvinylpyrrolidone (PVP) were obtained from Sinopharm Chemical Reagent Co. Ltd. Polydimethylsiloxane (PDMS) terminated by OH group were purchased from Aladdin Co. All of the mentioned reagents were used as received.

Silica Sol

10 ml deionized water was added into 12 ml ethanol. Then 2 ml TEOS was quickly dripped in the mixture of ethanol and water. This solution would be stirred at 600 rpm for 15 s. After that, 0.2 ml ammonia solution was dripped into the mixture under continuous stirring. Then the stirring rate would be raised to 900 rpm. The final mixture is kept at 25 °C for 24 h.

PFA Ink

Typically, we added 80 ml deionized water into a flask. Then 2 g PVP was added into the flask. The mixture should be stirred at 600 rpm until PVP completely dissolved. Then 1 ml FA was added into the flask. 10 copies of this solution should be prepared. The samples were reacted at different times (6-18 h) at 80 °C in oil bath to investigate the reaction time. The temperatures are also varied for stirring (60-180 °C) for 12 h in oil bath. When the heating process started, the stirring speed is raised to 1000 rpm. The liquids should be cooled to room temperature when these reactions ended, then they were poured into centrifuge tubes for separation. After 20 min centrifugation at 9000 rpm, the brown deposit left at the bottom of each tube was washed with the deionized water. Finally, 6 ml ethanol was poured into each tube. It would be through ultrasonic treatment (200W 30min) until the brown dope was dispersed.

Superhydrophobic Films

The obtained SiO₂ sol was mixed with different amounts of PFA. Typically, 1 ml silica sol was mixed with 0.5-2.5 ml PFA. Soda-lime-silica glass was used as substrates and was washed with ultrapure water and ethanol. The cleaned substrates

were heated to 220 °C, and then mixture of SiO₂ and PFA was sprayed onto glass. The spraying rate was 4 ml/min. The SRCA glass substrates were annealed in air at 520 °C for 1 h and cooled naturally. The coated substrates after sintering were soaked into Piranha solution (volume ratio of concentrated sulfuric acid to hydrogen peroxide (30% AR) is 7:3) for 3 min. After this, the films should be washed by deionized water and dried. Then an ethanol solution containing 1 wt% PDMS would be dipped onto the films to cover them, dried naturally in air for 30 s and then placed on a heating platform at 320 °C for 4 min. Among the obtained films, the samples that were produced by the PFA reacting at 150 °C for 12 h were denoted as F1, F2, F3, F4 and F5, according to the different amounts of PFA (0.5 ml, 1 ml, 1.5 ml, 2 ml and 2.5 ml).

Tempered Superhydrophobic Glass

0.002 g NaCl was added in the each F1, F2, F3, F4 and F5 inks. They were sprayed on auto-glass at 220 °C. After it, the temperature was raised up to 670 °C in 40 min and kept for 100 s in air. Then the glass with the coating was quenched and cooled naturally in the air. 0.25 ml hexane solution containing 1 wt% PDMS would be sprayed on it for 30 s. Then the glass was heated at 280 °C for 10 minutes. The obtained films were denoted as S1, S2, S3, S4 and S5.

Characterizations

The morphologies of the particles and the films were examined via transmission electron microscopy (TEM, JEM-2100F STEM/ EDS), scanning electron microscopy (SEM, ULTRA PLUS-43-13) and optical microscopy (Olympus BX51). The contact angles of different films were measured using a contact angle tester (Theta Lite,

Biolin Scientific) in the sessile mode at room temperature. The stylus profiler was the DektakXT model. The transmittance of the films was measured on an ultraviolet and visible spectrophotometer (UV-vis, PerkinElmer 750). The FT-IR was Nexus model from Thermo Nicolet. The TG/DTA test was measured on integrated thermal analyzer from Netzsch STA449F3. The pencil-scratching test was carried out using a pencil hardness tester according to State Standard Testing Method (GB/T6739-1996), equivalent of the American Society for Testing and Materials (ASTM D3363). A pencil was held firmly against the film at a 45° and pushed forward by the tester at a speed of 0.5 mm·s⁻¹. Cross-cut tape adhesion test is according to the American Society for Testing and Materials (ASTM D3359), equivalent to the ISO 2409. The particle sizes and Zeta potentials results were obtained by Malvin analyzer (Mastersizer 3000).

RESULTS AND DISCUSSIONS

To obtain the textured film, we conceive an organic-inorganic hybrid waterborne ink composed of silica nanocrystal and poly(furfuryl alcohol). Furfuryl alcohol is a type low cost renewable monomer and can be polymerized into PFA at various conditions such as acid and heat inclusion. Moreover, by employing the low-cost and water-ethanol ink materials, it promises environment-friendly manufacturing. After a simple sequentially reinforced additive coating process the superhydrophobic glass was obtained, shown in Figure 1. It contains 4 major steps of ink preparation, deposition, annealing and low surface energy treatment to achieve required properties.

The silica and PFA precursors were first prepared separately and then mixed as the hybrid ink (see details in the experimental session). The organic-inorganic species were additively coated on the glass substrate and self-assembled into required textures. The sacrificial pattern was constructed by crossed linked PFA, which induced self-phase separation from matrix and is removed by annealing in atmosphere. We prepared a 10*10 cm² sized superhydrophobic glass after low surface energy treatment of the annealed glass with good transparency (Figure 1).

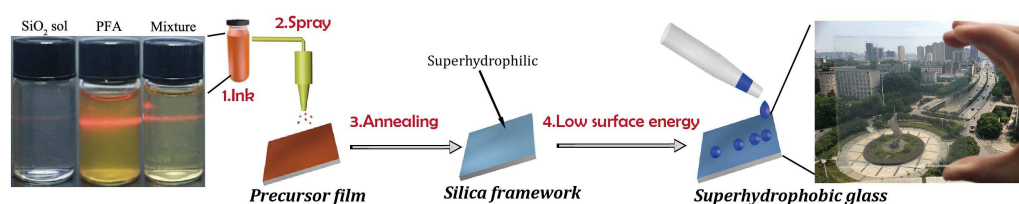


Figure 1. Schematic diagram of the sequentially reinforced additive coating (SRAC). Inserted: (left) The silica sol, poly(furfuryl alcohol) and the mixture ink demonstrating the Tyndall effects; (right) The transparent superhydrophobic glass prepared from the SRAC process.

The key issue during up-mentioned process is how to enable superhydrophobic repelling and also maintain high transparency and robustness. As can be noted in Figure 1, silica, PFA and the hybrid ink demonstrated obvious Tyndall effects, indicating colloidal dispersion. This is confirmed by the dynamic laser scattering (DLS) test. The particle sizes are around 10 nm and 300 nm for silica and PFA, respectively. However, the particles size in the mixture increased, with an average increasing to ~800 nm. This resulted from the interaction between PFA and silica, forming larger agglomerates. The peaked Zeta potential of the mixture (-6.28 mV) is just located between silica (-8.15mV) and PFA (-3.02 mV) indicating certain

connection happen in two species. However, no new bonds for silica and PFA mixture were suggested in the FTIR curves (Figure 2c), indicating possible hydrogen bonding and static electrical attraction for PFA and silica particles. Figure 2d shows the TEM image of PFA ink. The diameters of PFA spheres varied from tens of nanometers to micrometers with bottleneck connections. These perfect PFA spheres were disappeared in mixture (Figure 2e) and covered with the nano sized silica sol (inserted in Figure 2e). These observations suggest PFA and silica have been homogenously mixed and forming nano-micro scale multi-level architectures. This is a benefit to form required porous channel for superhydrophobicity after PFA being removed.

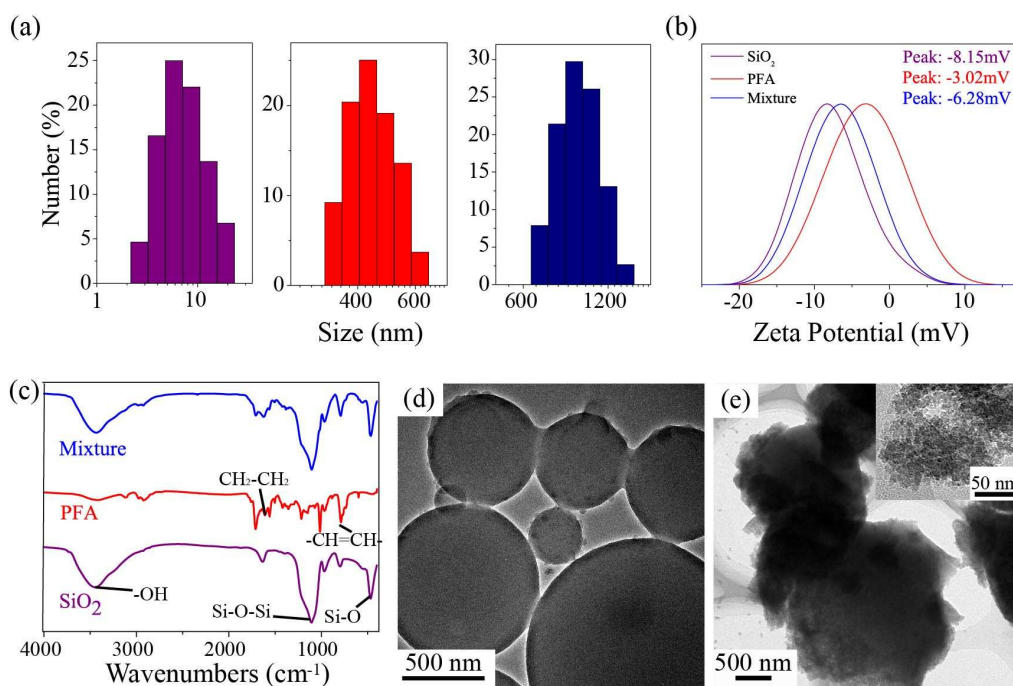


Figure 2. (a) The DLS results of SiO₂ sol, PFA and the mixture ink. The vertical axis represents the proportions of each particle size. The Zeta potential patterns (b) and FTIR curves (c) of SiO₂ sol, PFA and the mixture. The TEM images of PFA particles (d) and the mixture (e). The image of silica species is inserted.

As a kind of polymerizable monomer, with different synthetic conditions, such as the temperature and aging time, FA will transform into PFA with varied molecular weight in the solution. The polymerization process would affect the hydrophobicity and cause varied self-phase separation⁴⁸ and morphology construction, and thusly the superhydrophobicity and robustness of the SRCA films. The silica is mixed and bonded with PFA molecules in the nanometer scales. This phase separation is basically due to the PFA agglomeration and silica grain growth during solvent evaporation and temperature elevation. The morphology of porous architecture of silica film indicates the removing of PFA phases which are separated from the homogeneous silica-PFA solution precursor. All of those variables are crucial effects to the SRAC process for the micro-nano structure construction, and thus to the hydrophobicity of the final products. Figure 3 shows the hydrophobicity performances of samples with PFA prepared at different aging temperature and time. The samples are named as PFA-X, where X stands for the volume ratio of PFA to silica sol. For example, PFA-1.5 means that the proportion of PFA to SiO₂ is 1.5. With the increasing of aging time (Figure 3a) at 80 °C, the samples demonstrated similar trend of rapid increasing hydrophobicity and reached a plateau at around 12 h. All samples aged over 12 h are superhydrophobic except PFA-0.5. The average static contact angles increase with more PFA inclusion and reach a highest value of 170° for PFA-2.5 aging for 18 h. The similar result of enhancing the hydrophobicity with PFA concentration was observed for the samples prepared at different temperatures. The optimized aging temperature is 150 °C when the majority samples are

superhydrophobic. The corresponding photographs of the static contact angles for water droplets are inserted in Figure 3.

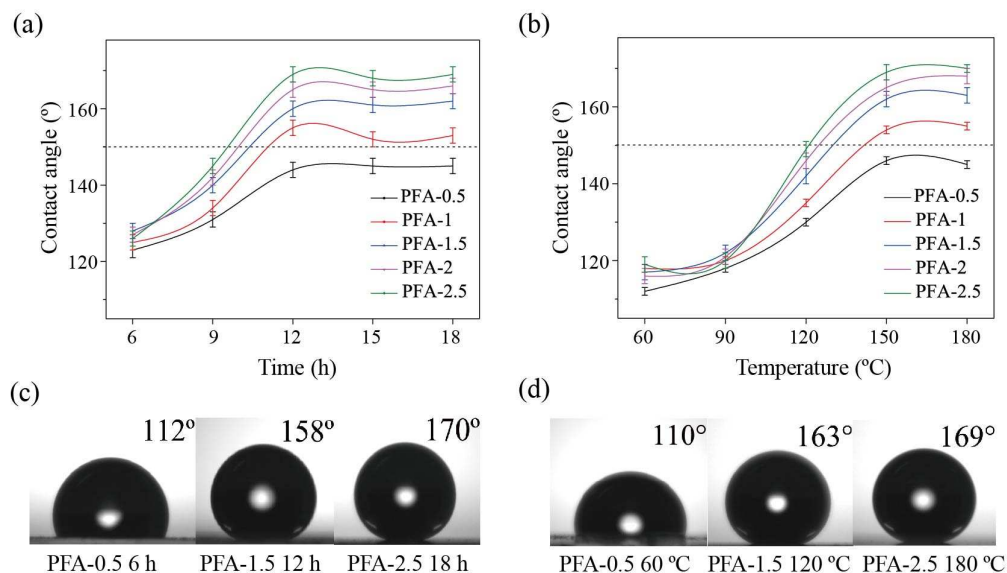


Figure 3. The effect of the polymerization conditions of PFA (aging time and temperature) on the hydrophobic performances of SRAC processed films. The PFA-0.5, PFA-1, PFA-1.5, PFA-2 and PFA-2.5 represent the different volume ratios of PFA/silica of 0.5, 1, 1.5, 2 and 2.5, respectively. The SCA results of at (a) the different aging time and (b) the different aging temperatures of PFA. Optical images of the contact angle of water droplets (c) varied aging time and (d) varied aging temperature.

We fixed the heating temperature at 150 °C and the aging time for 12 h, and the samples were named as F1 to F5 with the changing of PFA/SiO₂ volume ratio from 0.5 to 2.5 as shown in Figure 4a. With the increasing PFA/SiO₂ volume ratio, the hydrophobicity of the SRAC processed film was promoted, similar to Figure 3. The water contact angles of the samples appear almost linearly increasing from below 150° of F1 to ~170° of F5. However, from the UV-vis tests, the average transmittance shows opposite trend that decreases from over 90% to 70% (Figure 4b). This phenomenon is ascribed to the micro-nano structure construction with more PFA involved. The rougher surface was obtained, which induces more light-scattering. We

also confirmed this proposal by increasing SRAC volume of precursor. As shown in Figure 4c, when SRAC volume of the ink was increased (comparing to F3), the dramatic decreasing of transparency (at 600 nm wavelength) was observed with slightly enhanced SCA. It seems that the SCA reaches the highest plateau around the value of 170° as the continuous increasing of ink volume ratio to F3, whereas the transmittance continues dropping rapidly as the increasing of the ink volume. In the premise of high transmittance and contact angle, 1 to 1.25 times of F3 is a proper range, with the transparency over 80% and the SCA over 160° .

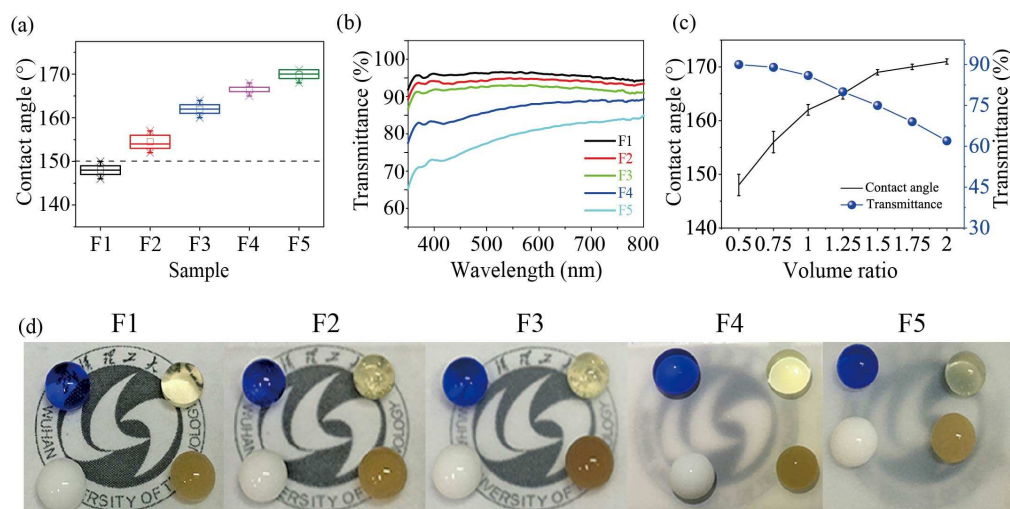


Figure 4. (a) The SAC box plots and (b) the transmittance of samples with varied PFA/silica volume ratio in the mixture ink. The samples with volume ratios of 0.5, 1, 1.5, 2 to 2.5 are named as F1, F2, F3, F4 and F5, respectively. (c) The effect of SRCA ink volume (ratio to F3) on hydrophobic performances and transmittances of samples. The transmittances are obtained at 600 nm wavelength. (d) The photographs of F1, F2, F3, F4 and F5 with different water solutions droplets such as fountain pen ink (blue), milk (white), coffee (brown) and juice (yellow).

The photographs of superhydrophobic glass F1 to F5, demonstrating in Figure 4d, suggest well performed superrepelling to the common water solutions such as fountain pen ink, milk, coffee and juice. The transparency of the samples decreased

from F1 to F5, and the printed logo of Wuhan University of Technology became vaguing as the increasing of PFA contains.

The SEM morphologies of as-derived films show porous structures (Figure 5a of F1, F2, F3, F4 and F5) with the voids derived from carbon removal. According to the Cassie model, the formula can be rewritten as follow:

$$\cos \theta^{\text{app}} = rf \cos \theta - 1 + f. \quad (1)$$

θ^{app} is the apparent contact angle on the surface, f is the fraction of the projected area of the solid surface in contact with the liquid, r is the roughness of the solid surface that is in contact with water and θ is the equilibrium contact angle on a smooth surface. A small value of f is the key to the construction of superhydrophobic films with continuous pore structure. As shown in Figure 5a, with the amount of PFA increasing, there are more and more pores on the surface and the silica structure is relatively slimmer. Such phenomena make more air remain, then lead to the significant decrease of projected area of the solid surface in contact with the liquid. So, the contact angle was increasing from F1 to F5.

The separated pores gradually evolve to interconnected channels as the increasing of PFA from F1 to F3. And this structure became finer when PFA was further increased for F5. This unique channeled structure facilitates air flowing when liquid droplets cover the pore and also promotes the sintering to obtain reinforced silica framework. The micrographs of the samples after scratching with 4H pencils are demonstrated in Figure 5b, suggesting obviously enhanced mechanical strength of F3 with no observable scratch mark even at 4H scratching (the scratching by different are shown

in Figure S1). Moreover, the superhydrophobic glass prepared from SRAC is also sustainable to 500 g sand abrasion⁴⁹⁻⁵¹ with the CA maintains over 150° (Figure S2 and S3). The hardness and CA for samples is shown in Figure 5c. Although the hardness varied with composition, the stylus profiling suggested only F1 film was fully stripped by pencil scratching. As showing in Figure 5a, the special structure, coffee ring, makes main contribution to the high hardness. More silica species are concentrated on the edges of the coffee rings, which make the walls of rings much tougher. Moreover, the higher thickness of the walls could protect the relatively brittle structure inside the rings from abrasion. The thickness of the SRAC films increased with the addition of PFA from ~2 μm to ~7 μm , indicating smaller pores and thinner framework walls were formed at high PFA inclusion in accord with the SEM observation.

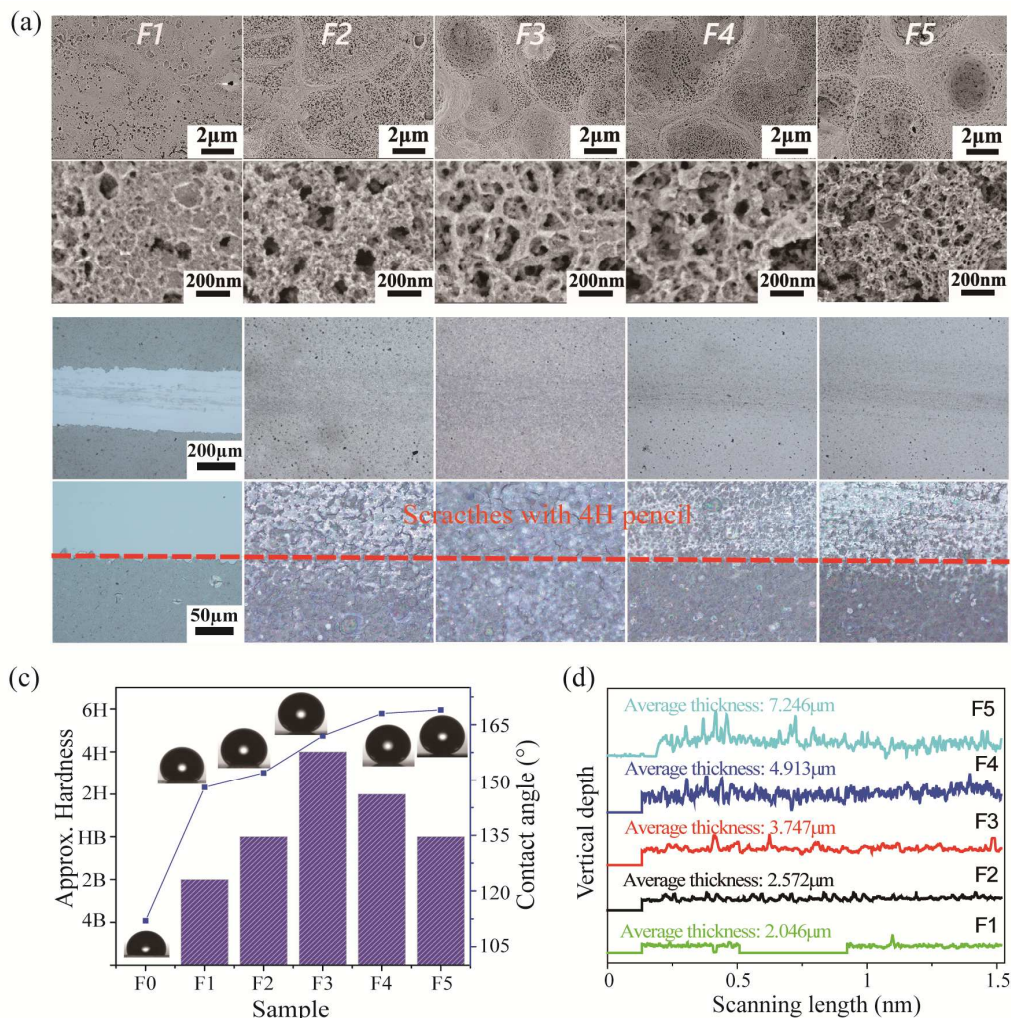


Figure 5. (a) SEM morphologies of samples F1, F2, F3, F4 and F5. The ~ 10 micrometer sized textures were obtained from spray coating. (b) The optical microscope images of 4H pencil scratches with suggesting varied the hardness of F1 to F5. (c) The approximate hardness and contact angles relations and (d) stylus profiles of the 4H pencil scratches of F1, F2, F3, F4 and F5.

No separated regions are stripped off for F3 film in a cross-cut tape adhesion test (Figure S4), indicating very tough bonding between the structured film and the bottom glass substrate. We further elevate the annealing temperature of F3 to 670 $^\circ\text{C}$ (renamed as S3), similar to the glass tempering process. The received superhydrophobic surface exhibits even higher hardness up to 6H, shown in Table 1. The improvement of mechanical strength is due to Na ion and elevated temperature,

which promoted the sintering of silica framework. The detailed results such as pencil scratching, contact angels, stylus profile, transmittance and etc. are demonstrated in Figure S5, S6, S7 and S8 in support information. The resistance to abrasion has been substantially improved with slightly decreasing of CA, indicating high potential of SRCA process to the glass production. A ratio of 1 to 1.25 times of S3 ink volume is the proper to obtain samples with transparency over 75% and the SCA over 160° (Figure S9).

In the 3D printing, means are required to support the free-standing, disconnected features and keep all the features of the part in place during the fabrication process. We consider that the interaction of PFA and silica presents the key function of supporting. The PFA polymers have branched molecular structure allowed them soften and solidify organic-inorganic film.⁵²⁻⁵⁴ This is also supported by the formation of inconsecutive film of powder like surface from silica ink alone, in contrast to the tough hybrid coating from hybrid ink (Figure S10). The thermal gravity analysis (TG) and differentiate thermal analysis (DTA) results suggest the silica precursor presents only a massive weight loss of ~20% at around 100-150 °C, which can be ascribed to the removing of residue solvent and hydroxyl condensation. No obvious weight lost was observed for silica in the following heat treatment up to 700 °C. The decomposing for PFA happened around 325 °C, and the further elevation of temperature to ~475 °C fully removed the PFA species into carbon oxides.

As mentioned above, when the PFA-silica hybrid lost H, O and partial organic groups at elevated temperatures, the residue carbon framework could still provide rigid

supporting to silica nano parts. This could be observed in the TEM images of the particles scraped from the SRAC film Figure 6b. The denoted red cycles are correlated to the vacuum of removed PFA/carbon spheres, with the pore size around 500 nm to 30 nm, forming multi-level channeled structure. This facilitates inter-granular sintering and forming high strength nano-structured film.

It can be concluded that PFA plays an important role in the construction of the micro-nanostructure in the process of coating and annealing. Due to its thermosetting property, the PFA skeletons can remain and support the silica framework until the decomposition of carbon. This effect can balance the contraction of silica in the sintering process, and then benefit the silica framework sintering. When the temperature reaches 400 °C, there are only carbon skeleton and silica left and the silica has started sintering. This could be partially supported by the tiny weight loss and exothermal peak at TG and DTA curves, respectively, indicating for further removing of organic species attached to the surface of silica particles. The EDS mapping images of films treated at different temperatures (Figure S11) also reveal the carbon removing process. We thus proposed a sequential reinforcement as following (Figure 6c): 1) The PFA-silica agglomerates are spraying and attached on the glass substrate; 2) PFA is acting as the binder for the powders during the additive deposition and form interconnected network to solidify the precursor film. 3) The hybrid film is strengthened by PFA network at the temperature up to 300 °C, and further supported by PFA-derived carbon network with temperature up to 450 °C; 4) The nano sized silica species start to sinter and be transformed into grain-bounded

silica film with required texture when PFA is removed. Please note that, reinforcement steps are sequentially taken the major responsibility to support the consecutive film, while the boundary temperatures for those mechanisms may not be defined exactly.

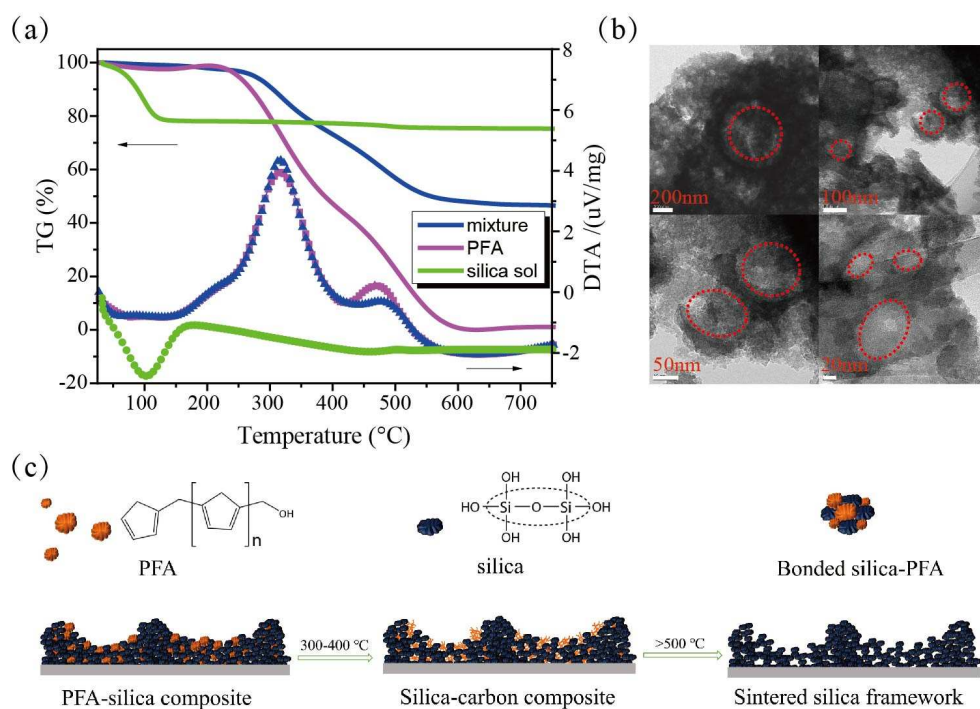


Figure 6. (a) The TG/DTA curves of silica sol, PFA and mixture. (b) The TEM images of the mixture when it has been heated to 400 °C. (c) The Schematic diagram of the strengthen process.

Table 1. The selected properties of superhydrophobic glass samples of PFA-0 F3, F5 and S3.

Samples	Contact angle (°)	Roll-off angle (°)	Pencil hardness	Transmittance (600nm)
PFA-0	100±1	>90	<10B	—
F3 (PFA-1.5, 500 °C)	162±2	3-5	4H	~87%
F5 (PFA-2.5, 500 °C)	170±1	~3	2H	~62%
S3 (PFA-1.5, 0.004g NaCl, 670 °C)	162±2	3-5	6H	86%

For the aim of application, we carried out an outdoor test demonstration. The sample was placed in open air at a 45° angle with superhydrophobic surface outwards in Wuhan, China for a continued period of 3 months. The recorded CA results are shown in Figure 7 along with the weather information. The temperature variations of this period are shown in Figure S12. According to China's National Ambient Air Standard GB3095-2012 and Technical Regulation on Ambient Air Quality Index HJ633-2012, AQI (air quality index) contains a rating on six substances, including PM2.5, PM10, SO₂, NO₂, O₃ and CO. The highest one among the values is published, which are majorly on PM2.5 in cities. When AQI value over 100 is considered as pollution and the density of PM2.5 reaches 75 µg/cm³. It seems that the high AQI weather will decrease the CA of the glass, but the rain flush could create remedy for the hydrophobicity to some extent, demonstrating self-cleaning effect. Moreover, ultraviolet light also causes damage to the performance probably due to the degradation of organic coating. The CA decreased to ~154° after 3 months continuous exposing outdoor weather, maintaining superhydrophobicity. This result suggested high application potential for SRCA film with further modifications.

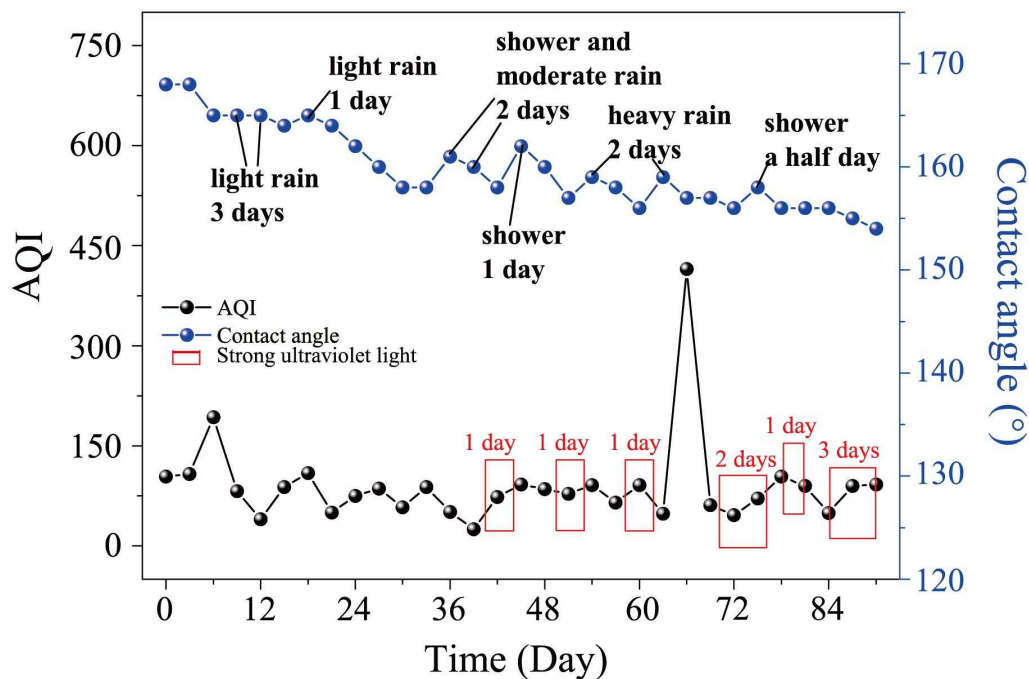


Figure 7. 90-day outdoor examination of F3 sample in Wuhan. Strong ultraviolet light weather means the ultraviolet radiation reaching the ground is 15-30 W/m².

CONCLUSION

In summary, we present a facile sequentially reinforced additive coating (SRAC) process to fabricate transparent robust superhydrophobic glass. The aqueous ink with poly furfuryl-alcohol (PFA) and silica species are carefully designed, and spray deposited on glass. Through delicate ink design and heating process, we enabled self-phase separation and sequentially reinforcement for the silica to achieve robust and transparent superhydrophobic films. The PFA aging conditions and PFA/ratio exhibit varied morphology and mechanical strength. The robust micro-nano structured film with tough intergranular sintering, which can withstand 4H pencil test, cross-cut test and sand abrasion. The SCA for water is as high as 164° and the transmittance

can be as high as 87%. This SRAC method can be combined with glass-tempered process. We realized a durable and transparent textured layer for superhydrophobic glass with the high hardness up to 6H. Moreover, we performed 90 days out-door performance test. The glass maintained superhydrophobicity with the contact angle of 154° . Thus, this convenient waterborne SRAC method is promising for up-scaling fabrication of superhydrophobic glass environmentally friendly and cost-effectively.

Associated Content

Supporting Information. Brief statement in nonsentence format listing the contents of the material supplied as Supporting Information.

Acknowledgement

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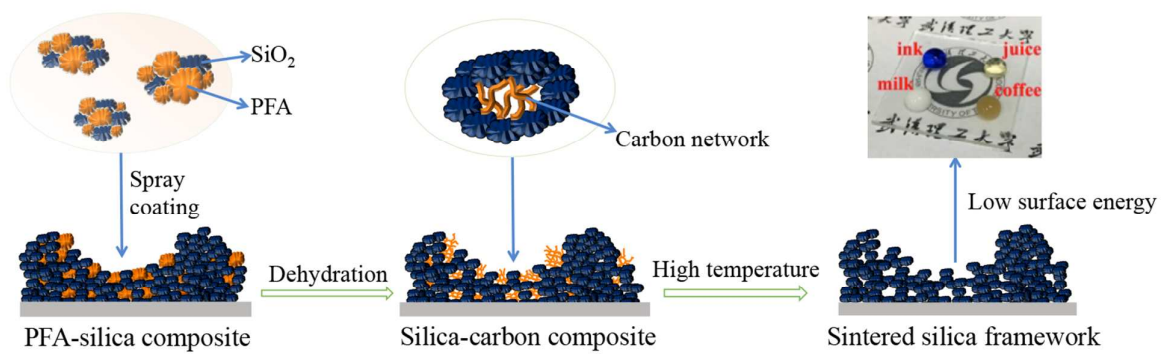
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