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# Mesoporous silica-based abrasion resistant antireflective (AR)-*cum*-hydrophobic coatings on textured solar cover glasses by a spray coating technique†

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The present work describes the fabrication of functionalized mesoporous silica based antireflective-*cum*-hydrophobic coatings on textured solar cover glasses *via* a spray-coating technique; the fabricated coatings exhibit good abrasion resistance. The thicknesses of the AR and hydrophobic layers have been varied in the ranges of 350–500 nm and 30–45 nm, respectively. The application of a bilayer coating on textured solar cover glass resulted in a significant enhancement of  $\geq 4\%$  in the average solar transmission in the wavelength range of 400–1100 nm. A hydrophobic coating with self-cleaning properties was designed on the mesoporous silica coating, and it exhibited an enhancement of the photo-current density ( $\text{mA cm}^{-2}$ ) of 4.65% under simulated solar light of 1 sun with a maximum output power enhancement of  $\geq 4\%$  based on  $I$ - $V$  (ampere-volt) characterization. After 50 cycles of sandpaper (80 mesh) abrasion testing of the coated solar cover glass, its water contact angle value was reduced from  $130^\circ$  to  $105^\circ$ , and 50 h UV radiation exposure testing confirmed that there were no significant changes in the coating quality. These results indicate the good durability, abrasion resistance, and AR/hydrophobic properties of the coatings.

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## 1. Introduction

The generation of solar power as a renewable energy source is highly important in society for a variety of applications.<sup>1–8</sup> In the current scenario, the development of Si solar cells or photovoltaic cells that convert direct sunlight into electric energy is drawing increasing attention because it prevents environmental pollution. Solar cell modules are normally fabricated by sandwiching an Si solar cell between a back sheet and a cover glass (front sheet) and sealing them with a resin material. The cover glass of the solar cell that protects the power-generating elements contains silicon or other related materials in its interior. In order to improve the conversion efficiency of the solar cell, the cell should take in a greater amount of sunlight. Accordingly, the cover glass must have excellent transparency and low reflectivity. For this reason, the use of mesoporous silica, which has thick pore walls with excellent hardness and a low refractive index, as a coating material could enhance both the normal and diffuse transmission of light

through the solar cover glass (antireflective (AR) effect) to the solar cell, resulting in an increase in photocurrent. Presently, ‘millimeter scale textured solar cover glass’ (MSTSCG), which is the third-generation solar cover glass, is capable of trapping extra light, and has the advantages of antidazzle and positive temperature effects, is being used. Still, there exists ample scope for research to further improve the light-harvesting properties of such textured solar cover glasses. The lifetime of cover glasses for solar cell applications is also an important factor in fabricating solar cell modules. As the cover glass is exposed to open air, dust particles may adhere to it, reducing its transparency and thus decreasing the conversion efficiency of the solar cell module.<sup>1–3</sup> To maintain the conversion efficiency of the solar cell at a desirable level, the cover glass must be coated with a protective, hard and hydrophobic layer with low reflective properties. The use of mesoporous silica with a relatively ordered pore structure as a coating layer has been proven to achieve better light conversion efficiency with easy cleaning and maintenance.<sup>4–8</sup>

Various approaches have been used to apply thin coatings on ‘solar cover glasses’ (SCGs) and solar cells to increase the efficiency and durability of the coatings deposited on solar panels or modules. Silica-based AR coatings with disordered pore structures have been reported for photovoltaic cells,<sup>9–14</sup> solar glass and ophthalmic lenses. To improve their moisture resistance or hydrophobicity, fluorosilane<sup>4–6</sup> and hexamethyldisilazane

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(HMDS)<sup>9</sup> based precursors have been used. Some of the reported coatings have been applied on solar cover glass or solar cells and cured at high temperature with a low solar photo-current.<sup>15,16</sup> R. Ding *et al.*<sup>17</sup> developed a broadband double-layer AR coating *via* a solvent evaporation self-assembly method using two different mesoporous films. The broadband AR coating increases the power conversion efficiency of solar cell by 1.23% for quartz, 1.31% for borosilicate glass and 1.37% for K9 glass. Attempts were made to develop AR coatings *via* a sol-gel route on glass and other substrates using a single- or double-layer deposition technique.<sup>17–19</sup> Furthermore, single-, double- or multilayer AR coatings, some with hydrophobic surfaces, on glass and solar cover glass substrates using mesoporous or low-refractive-index materials to enhance transmission or the photo-current efficiency of the solar cell have also been reported.<sup>20–33</sup> However, these coatings showed relatively low photo-current efficiency (<3%). Gundulohelsch *et al.*<sup>29</sup> prepared an AR coating with self-cleaning and photo-catalytic properties on solar cover glass using a SiO<sub>2</sub>-TiO<sub>2</sub> sol *via* a dip-coating technique. Vipul Jain *et al.* reported a silica-based nano-coating on solar glass with 3% transmission and a water contact angle range of 109–112° using a dip-coating technique.<sup>33</sup>

With these motivations, in this work, the main objective was to develop a silica-based transparent, protective, hard and hydrophobic mesoporous coating on the millimeter scale for textured solar cover glasses (MSTSCG) that would increase the transmission to >3% and consequently enhance the output photocurrent, as well as exhibit self-cleaning properties, *via* the spray-coating technique. To obtain a suitable AR effect using a monolayer AR coating, the refractive index (RI) of the AR coating layer should be the geometric mean value of the refractive indices of air and the substrate ( $n_a n_s$ )<sup>1/2</sup>. The optical glass of transparent components generally has a RI of 1.47–1.92. Therefore, the RI of the AR film must be in the range of 1.21–1.38. Here, the solution for the AR coating was prepared using a silica precursor in the presence of pluronic F127 as co-block polymer *via* a sol-gel technique. The refractive index value of the thin films was optimized by varying the concentrations of the precursors and surfactants. The coated samples were characterized using FTIR, AFM, TEM, TG and GIXRD studies, along with evaluation of the adhesion, abrasion, optical transmission, durability and *I-V* (ampere-volt) characteristics of the coatings.

## 2. Experimental details

### 2.1 Materials

All chemicals were used as received. Tetraethyl orthosilicate (TEOS), methyltriethoxysilane (MTES), the surfactant Pluronic F127 (EO<sub>100</sub>PO<sub>70</sub>EO<sub>100</sub>; Mol. Wt = 12 600 g mol<sup>-1</sup>) and 1,1,1,3,3,3-hexamethyl disilazane (HMDS) were supplied by Sigma-Aldrich, while ethanol, 1-propanol, 1-butanol, *N*-methyl-2-pyrrolidone, and HCl (35.4%) were obtained from MERCK Specialties Pvt. Milli-Q (Millipore) water (18.2 MΩ) was used throughout the study.

**Table 1** Variation of the RI values of the thermally cured coatings derived from base coating sols having different F127 concentrations (with 6.5 wt% of SiO<sub>2</sub> equivalent)

Sample name	Concentration of F127 present in the base coating sol (wt%)	RI value of the thermally cured coating (measured at 633 nm)
SF-3	3	1.355 ± 0.005
SF-5	5	1.328 ± 0.005
SF-6	6	1.292 ± 0.005
SF-7	7	1.288 ± 0.005
SF-8	8	1.269 ± 0.005

### 2.2 Preparation of sol for AR coating

For the preparation of nanocomposite AR coating sol, MTES (0.0913 mol) and TEOS (0.1875 mol) were mixed with 1-butanol (0.2844 mol), followed by adding a mixed solution of HCl (0.00122 mol), water (0.202 mol), 1-butanol (0.036 mol) and *N*-methyl-2-pyrrolidone (0.10 mol) to the former solution. After stirring for 10–15 min at room temperature (RT) (25 ± 2 °C), the solution was then heated at ~60 °C for 60 min followed by cooling at RT to obtain solution A. The surfactant F127 (0.0012328 mol) was mixed with ethanol (3.41 mol) under stirring for 1.5 h to obtain solution B. Solution A was added to solution B, maintaining the optimized 6.5 wt% SiO<sub>2</sub> equivalent and 6 wt% F127 (designated as SF-6), under stirring for 60–70 min to obtain a sol suitable for AR coating. In a similar way, by varying the surfactant concentration to 3, 5, 6, 7 and 8 wt%, different sols were prepared and designated as SF-3, SF-5, SF-6, SF-7 and SF-8, respectively (Table 1).

### 2.3 Preparation of sol for hydrophobic coating

For the preparation of a sol suitable for use as the hydrophobic coating, a mixed solution of TEOS (1.23 mol), MTES (1.23 mol), 1-propanol (94.2 mol), water (7.9 mol), and ammonia (0.33 mol) was prepared under stirring at RT for 2 h, followed by ageing for 4 days. To this, hexamethyldisilazane (HMDS) (0.9 mol) was added for surface modification of the SiO<sub>2</sub> generated in the above sol and aged at RT for another 3 days. The acid-based 'Solution-A' prepared for AR coating was added to the above solution, maintaining the molar ratio of (TEOS + MTES): 1-propanol : water : ammonia : HMDS as 7.64 : 29 : 24.47 : 1 : 2.72 to obtain a 1.5 wt% SiO<sub>2</sub> equivalent (considering SiO<sub>2</sub> equivalents from TEOS + MTES + HMDS) sol for the hydrophobic coating sol.

### 2.4 Coating preparation

Prior to the deposition of coatings, the glass substrates were first cleaned with detergent, followed by washing with tap water and rinsing with distilled water, and then finally cleaned with ethanol. In the first step, an AR coating was prepared on the cleaned glass substrates using the AR coating sol *via* the spray-coating technique (USI, Ultrasonic System Inc.). The deposited coating was dried at 70–80 °C for 12 h and heat-treated at 450 °C for 1 h in an air oven (ramp rate 1.5 °C min<sup>-1</sup>). The thicknesses of the thus-obtained coatings were in the range of 350–500 nm through XY movement of the spray head twice with

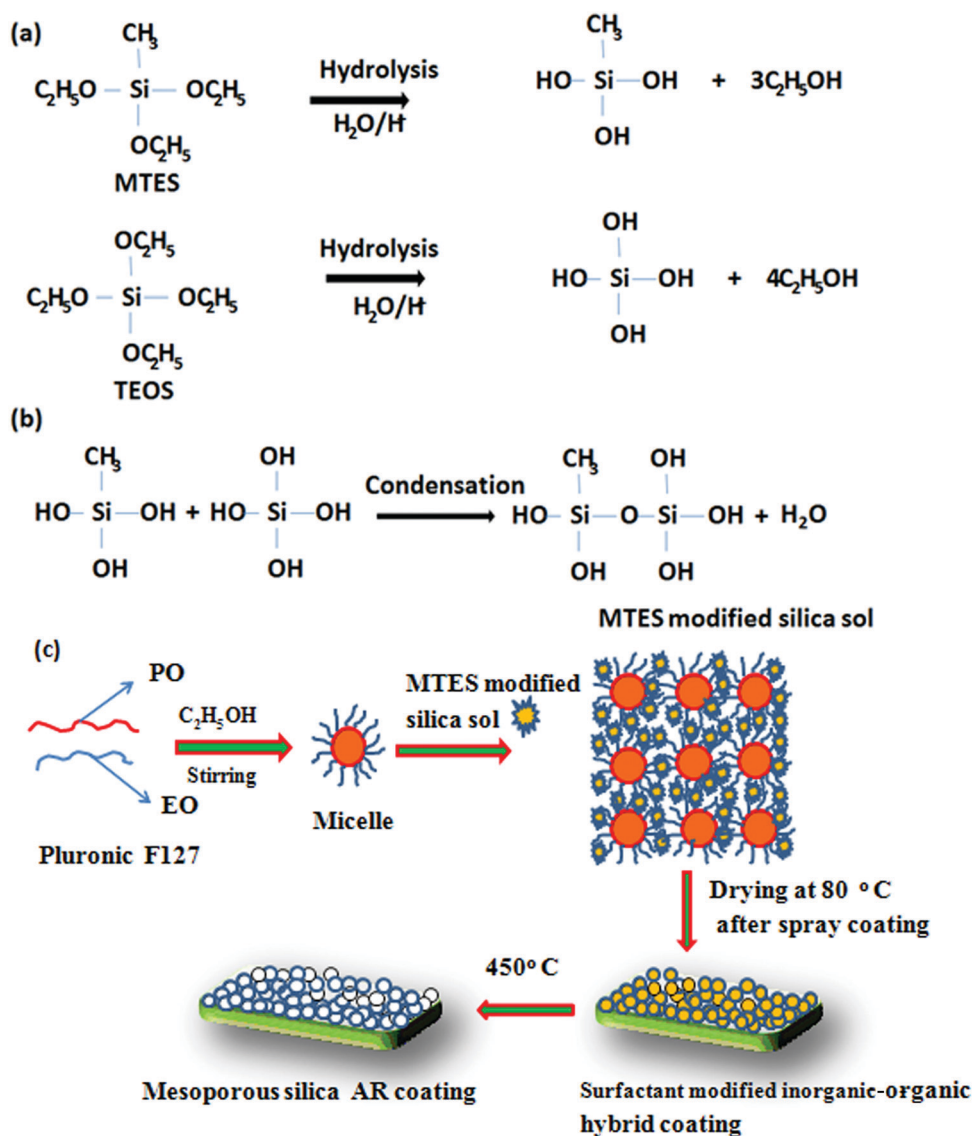


the optimized spray head operational parameters, *i.e.*, flow rate  $0.5 \text{ mL min}^{-1}$ , head speed  $100 \text{ mm s}^{-1}$ , spray width  $15 \text{ mm}$  and air pressure  $20 \text{ psi}$ . In the second step, the hydrophobic coating was developed on the AR-coated substrate using the hydrophobic coating sol *via* the above technique with a single XY movement of the spray head with the following operational parameters: flow rate  $0.5 \text{ mL min}^{-1}$ , head speed  $200 \text{ mm s}^{-1}$ , spray width  $15 \text{ mm}$  and air pressure  $20 \text{ psi}$ . The two-layered (AR and hydrophobic) coated substrates were then dried at  $60 \text{ }^\circ\text{C}$  for  $1 \text{ h}$  followed by heating at  $180 \text{ }^\circ\text{C}$  for  $2 \text{ h}$ . The thicknesses of the hydrophobic coating were designed to be  $30\text{--}45 \text{ nm}$  ( $\leq 50 \text{ nm}$ ) to avoid compromising the optical properties of the two layers.

## 2.5 Coating characterization

A spectroscopic ellipsometer (J. A. Woollam Co., Inc., USA) was used to measure the thickness and RI of the coatings deposited on single-side-polished Si-wafers. Coatings were developed on

the substrates using an automatic ultrasonic spray-coating technique (BT Prism 400, USI, USA). The attenuated total reflection Fourier transform infrared (ATR-FTIR) spectra of the dried and heat-treated coatings were recorded using a Bruker ECO (ALPHA) spectrometer in the range of  $600\text{--}4000 \text{ cm}^{-1}$  with a resolution of  $4 \text{ cm}^{-1}$  and 200 scans. A UV/visible/NIR spectrometer (PerkinElmer Lambda 900) was used to measure the transmission spectra of the coatings. An X-ray diffractometer (Rigaku SmartLab) operating at  $9 \text{ kW}$  ( $200 \text{ mA}$ ,  $45 \text{ kV}$ ) using  $\text{Cu-K}\alpha$  ( $\lambda = 1.54059 \text{ \AA}$ ) radiation was used for low-angle XRD of the coatings; a grazing angle of  $0.3^\circ$  was maintained with a scanning speed of  $1^\circ \text{ min}^{-1}$  using a step size of  $0.02^\circ$ . Thermogravimetric (TG) measurements were performed using a Netzsch STA model 449F3 thermal analyzer at a dynamic heating rate of  $1.5 \text{ }^\circ\text{C min}^{-1}$ . Transmission electron microscopy (TEM) measurements were carried out using a JEOL 2010 transmission electron microscope equipped with an EDX



Scheme 1 The proposed reaction pathway during the preparation of the AR coating.





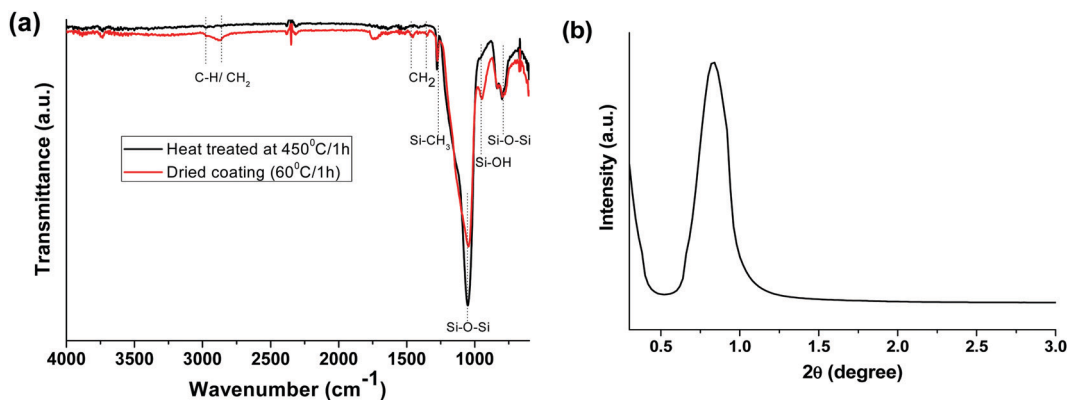


Fig. 1 (a) ATR-FTIR spectra of as-prepared and heat-treated (450 °C/1 h) AR coatings and (b) low-angle XRD patterns of the AR coated glass heated at 450 °C for 1 h.

### 3.2 FTIR and GIXRD

Fig. 1a presents the FTIR spectra of the as-dried (60 °C/1 h) and heat-treated (450 °C/1 h) AR coatings. The absorption bands at 2856–1961  $\text{cm}^{-1}$  and 1350–1450  $\text{cm}^{-1}$  are due to C–H/CH<sub>2</sub> stretching and bending vibrations, respectively. The peak at 1276  $\text{cm}^{-1}$  is attributed to Si–CH<sub>3</sub>.<sup>32</sup> The characteristic asymmetric and symmetric stretching vibrations of Si–O–Si were indicated by the absorption bands at 1046 and 796  $\text{cm}^{-1}$ , respectively. However, the as-dried AR layer showed the characteristic band of Si–OH at 946  $\text{cm}^{-1}$ .<sup>35,36</sup> Interestingly, the peaks originating from organic moieties (C–H/CH<sub>2</sub>) disappeared after heat treatment, indicating the complete removal of organics from SiO<sub>2</sub>, except the Si–CH<sub>3</sub> peak (1276  $\text{cm}^{-1}$ ), which was unaffected at this temperature.<sup>37</sup> Thermal treatment of the coating resulted in a decrease in the Si–OH peak (946  $\text{cm}^{-1}$ ) intensity and a shift of the Si–O–Si peak

from 1046 to 1051  $\text{cm}^{-1}$ , indicating some silanol condensation, which helps to further strengthen the silica network.

A low-angle XRD study was carried out to analyse the mesostructural properties of the heat-treated AR coating at 450 °C (Fig. 1b). The broad reflection peak at around 0.7 (2 theta) indicates the porous nature of the silica layer.<sup>38,39</sup> The mesoporous silica coating was formed *via* the spray coating technique due to the fast evaporation of the solvents, as well as the heat treatment of the coating.

### 3.3 TEM analysis

To investigate the microstructural features of the coating, a TEM study was performed (Fig. 2). Fig. 2a and b show the TEM images of the scraped SF-6 coating at different magnifications, along with the selected area diffraction (SAED) pattern (Fig. 2c) and EDX spectrum (d). Low- and high-magnification TEM images (Fig. 2a and b) confirm the mesoporous nature of the SiO<sub>2</sub> coating with an average pore diameter of 6 nm. The volume fraction porosity (*P*) of the coating was evaluated using the Lorenz–Lorentz equation<sup>35</sup> and was found to be ~25% with a pore size of ~6 nm. The SAED pattern (Fig. 2c) indicates the amorphous nature of the coating, while the EDX spectrum (Fig. 2d) reveals the presence of C, O and Si in the coated sample.

### 3.4 AFM study

The hydrophobicity of a surface is dependent on both its chemical composition and topography. An AFM study of one representative hydrophobic film with a thickness of ~400 nm was performed to examine the surface topography. Fig. 3 shows an AFM image of the film covering an area of 20 × 20  $\mu\text{m}^2$ . The surface of the AR with the hydrophobic coating was found to be spiky or micro/nano in nature with a root-mean square (rms) roughness value of about ~64 nm. This type of surface topography favours a mesoporous-silica-based AR-*cum*-hydrophobic coating surface. The photovoltaic performance will completely depend on the AR-*cum*-hydrophobic mesoporous silica, which could improve the light trapping mechanism and reduce the surface reflection of incident light.<sup>40</sup> The light scattering of the

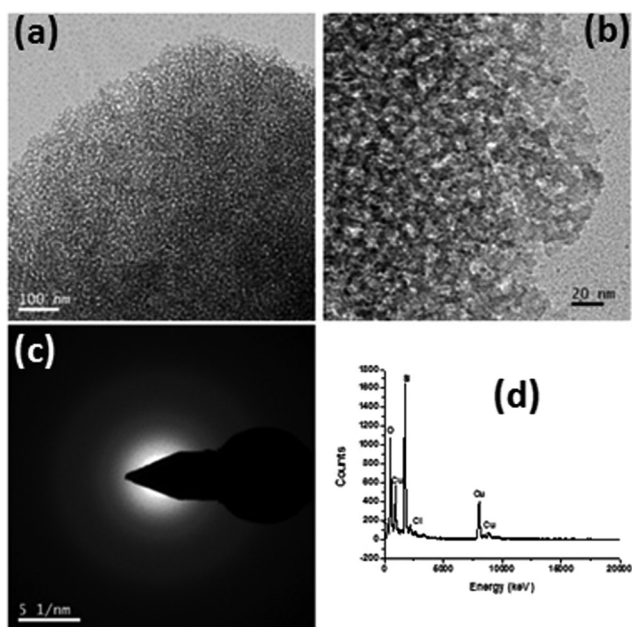


Fig. 2 (a and b) TEM images of the AR nanostructured mesoporous hydrophobic coating at different magnifications along with the (c) SAED pattern (d) and EDX spectrum.



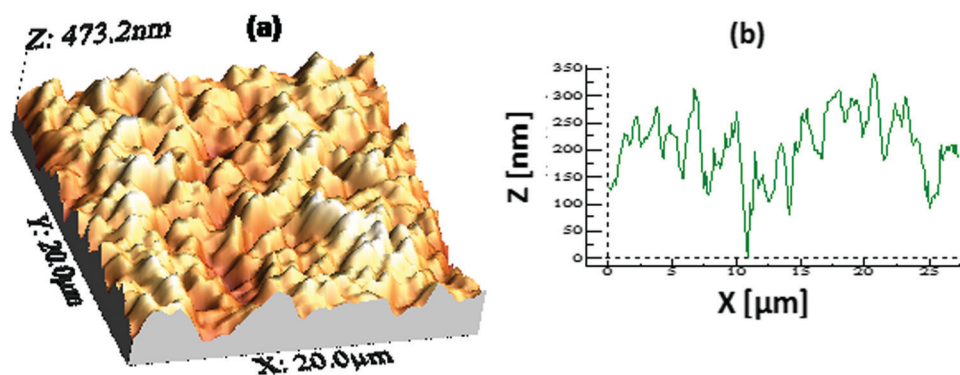


Fig. 3 (a) An AFM image ( $20 \times 20 \mu\text{m}^2$  area) showing the surface topography of a representative coating with a thickness of  $\sim 400$  nm and (b) the peak height profile of the coating. The rms surface roughness value obtained from the image is  $\sim 64$  nm.

mesoporous hydrophobic surface is strongly related to the surface roughness, which was characterized using AFM. Therefore, it increases the diffuse transmitted light from the incident beam scattered on the rough surface with a reduction of Fresnel reflection in the polycrystalline silicon solar cell.

### 3.5 Properties of the coatings

Fig. 4 shows the spray-coating technique for the preparation of the AR with a hydrophobic layer onto a  $>300 \times 300 \text{ mm}^2$  solar cover glass. The optimized thickness of the coating (350 to 500 nm) shows an RI value of  $1.295 \pm 0.005$  during the thermal decomposition of the surfactant, enhancing the porosity of the thin film. Hence, lowering the refractive index of the film, which is optically transparent and crack-free, helps to improve the optical performance of the polycrystalline silicon solar cell. The cross-cut scotch tape test (following the DIN 53151 or ASTM D 3359 specification) of the coating showed no damage, indicating good adhesion of the coated material to the substrate. The surface hardness of the AR coating was  $\sim 5\text{H}$  (ASTM D 3363), and after the deposition of the hydrophobic coating on it, a hardness value of  $\sim 3\text{H}$  was obtained. Table 2 shows the properties of the coatings derived from specific sols. After the application of hydrophobic coatings of thicknesses of 30–45 nm on top of the AR coating, the water contact angle (WCA) was in the range of  $127 \pm 3^\circ$  with a hysteresis value of

$10 \pm 2^\circ$  without affecting the AR, output power or photocurrent. The surface hydrophobicity increased due to the functionalization and micro/nanostructure of the AR coating fabricated on the millimeter-scale textured solar glass. Measuring the transmission spectra in the solar-sensitive wavelength range of 400–1100 nm on the double-layer AR-*cum*-hydrophobic coated SCGs for different sets of coating compositions (SF-3–SF-8), it is clear that the average solar transmission is increased  $\geq 4\%$  with a decrease in the reflection (average  $\sim 4\%$ ) in SF-6 (Fig. 5 and Table 3). Lower light transmission values were observed in case of the SF-3 and SF-5 coatings compared to SF-6 due to the higher refractive index values and lower percentages of pores on the thin film surface. On the other hand, SF-7 and SF-8 coated glasses were found to show little adverse effect due to the higher percentages of surfactants present in their compositions. Fig. 5 shows the transmission spectra of the different compositions, along with that of uncoated SCGs for comparison.

To determine their long-term durability in atmospheric conditions, the mechanical properties of the AR-*cum*-hydrophobic coating were evaluated *via* an abrasion test followed by measurement of the contact angle of the coating. For this test, 80 mesh sandpaper and a 100 g weight are required.<sup>41</sup> First, the 80 mesh sandpaper was placed on the coated substrate and the 100 g weight was loaded on the sandpaper. The sandpaper was then pulled slowly over a distance of up to 6 cm on the coating area for 50 cycles. After 50 such cycles, the static WCA value

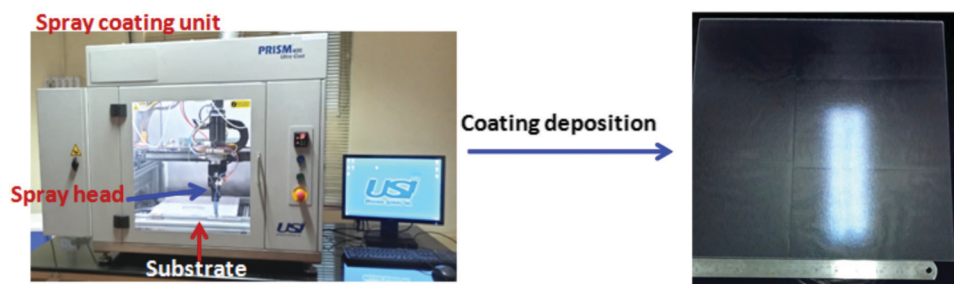


Fig. 4 Spray-coating technique for the fabrication of the AR coating with a hydrophobic layer onto  $>300 \times 300 \text{ mm}^2$  solar cover glass.



Table 2 Detailed test results for the mesoporous-silica-based AR and hydrophobic coatings on solar glass

Name of the test/physical property	Specifications	Results
Visual appearance	—	Optically clear
Average transmission (% <i>T</i> )	—	Overall % <i>T</i> increases about 4–5% (in the wavelength range of 400–1100 nm) as compared to bare solar glasses
Thickness	—	AR coating: 350–500 nm Hydrophobic coating: 20–45 nm
Refractive index (RI)	—	1.295 ± 0.005
Adhesion	DIN 53151 or ASTM D 3359	ASTM class 5B (highest standard)
Pencil hardness	ASTM D 3363	≈ 5H; after hydrophobic coating
Volume fraction porosity and pore size	—	23–27% with a pore size of 5–7 nm
Water contact angle (WCA)	—	127 ± 3° and hysteresis of 10 ± 2°
Maximum photo-current density ( <i>J</i> <sub>sc</sub> )	—	Increases > 3 to 4.5% compared to solar panels available on the market

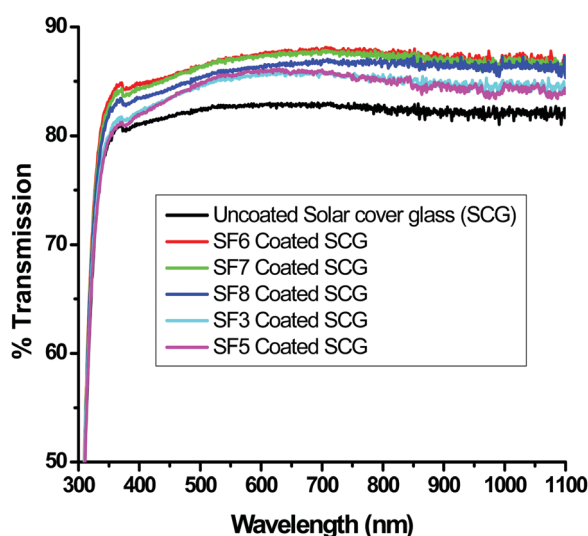


Fig. 5 Transmission spectra of bare (uncoated) solar cover glass (SCG) and those of AR-coated SCGs using different coating compositions.

(average data of ~10 points) was reduced to 105° from its initial value of 130°, indicating the good abrasion-resistance properties of the coatings (Fig. 6). This indicates that the AR/hydrophobic properties remained unchanged after the abrasion test without any damage to the coating.

To determine the thermal stability of the films, the final AR sol was evaporated to dryness *via* heat-treatment at 80 °C for 12 h and subjected to a thermogravimetric (TG) study.

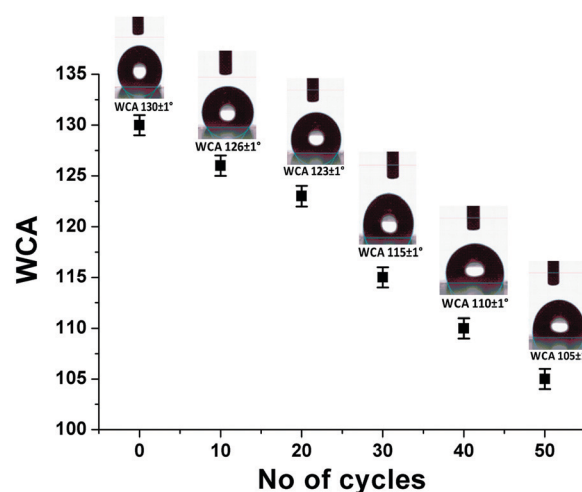


Fig. 6 Water contact angle (WCA) value (SF-6) after different numbers of abrasion cycles.

The results are shown in Fig. 7. It was observed that the weight loss up to 300 °C corresponded to the liberation of moisture, trapped solvents and other volatile components present in the material. The TG curve shows an approximately 22.5% weight loss up to about 300 °C; above this temperature, the decomposition of organic components began, and beyond 450 °C, no noticeable weight loss was observed.

We also evaluated the stability of the coated films with respect to UV radiation exposure (in a UV chamber) by measuring

Table 3 Percent transmission data of uncoated, AR-coated and AR-*cum*-hydrophobic coated millimeter-scale textured solar cover glasses (MSTSCG)

Sample	Average % transmission in the wavelength region		Increase in % transmission due to AR- and AR- <i>cum</i> -hydrophobic coatings in comparison to the uncoated solar glass in the wavelength region	
	400–800 nm	800–1100 nm	400–800 nm	800–1100 nm
Millimeter-scale textured solar cover glass (SCG) (uncoated)	82.59	82.15	—	—
SCG with SF-3 AR and hydrophobic coating	84.95	84.72	2.36	2.57
SCG with SF-5 AR and hydrophobic coating	85.07	84.38	2.48	2.23
SCG with SF-6 AR and hydrophobic coating	87.01	87.12	4.42	4.97
SCG with SF-7 AR and hydrophobic coating	86.82	86.83	4.23	4.68
SCG with SF-8 AR and hydrophobic coating	85.90	86.41	3.31	4.26



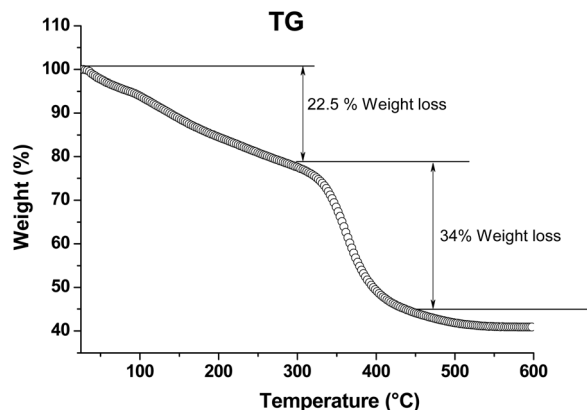


Fig. 7 TG analysis of an AR powder sample. Measurements were carried out at a heating rate of  $1.5\text{ }^{\circ}\text{C min}^{-1}$  in air.

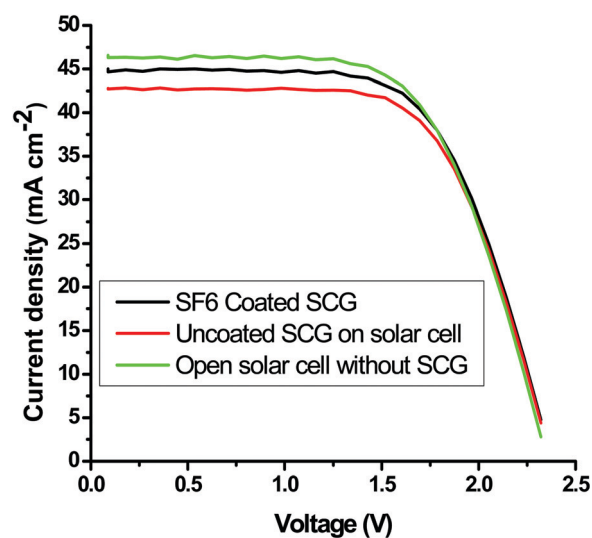


Fig. 8 Evaluation of the maximum output photo-power and short circuit current of the mesoporous-coated and uncoated textured solar cover glass.

their WCA values. For this purpose, coatings of about 350 nm in thickness were prepared on solar cover glass substrates and their WCA values were measured as a function of the exposure time to UV energy radiation (intensity UV-A:  $83.75\text{ W m}^{-2}$ ; UV-B:  $13.75\text{ W m}^{-2}$ , sample surface temperature:  $60\text{ }^{\circ}\text{C}$ ). From the test results, it was observed that after continuous UV exposure for 50 h, the WCA of the coating changed slightly from  $130^{\circ}$  to  $124^{\circ}$ ,

resulting in no significant changes in the percent transmission, which indicates the good durability of the coatings.<sup>42</sup>

### 3.6 Solar performance

Polycrystalline solar cells were used to investigate the effect of the optimized mesoporous silica AR-cum-hydrophobic coating on the performance of the solar cell. Standard 1 sun light from a solar simulator was illuminated (illuminated area:  $2 \times 2\text{ cm}^2$ ) on a silicon solar cell with a coated textured solar cover glass and compared with an uncoated textured solar glass using a Keithley 4200-SCS unit to measure the electrical power through the  $I$ - $V$  curve. From the  $I$ - $V$  curve, it was found that maximum current density,  $J_{sc}$  ( $\text{mA cm}^{-2}$ ) of the SF-6 coated SCG substrate was increased up to 4.65%, and, interestingly, the maximum power output was also enhanced up to 4.0% compared to that of the uncoated solar cover glass substrate (Fig. 8 and Table 4). This also further supports the average increase of 4.66% in the light transmission value in the wavelength range of 400 to 1100 nm (Fig. 5) compared to that of the uncoated SCG. The AR performance and light-trapping efficiency were improved on the porous coated textured glass *via* the diffuse transmission mechanism with a reduction of the Fresnel reflection, which increases the optical path length of light to enhance the current density ( $J_{sc}$ ) in the solar cell.

Table 4 shows the details of the  $I$ - $V$  data of the samples coated with coatings of different compositions (SF-3, SF-5, SF-6, SF-7 and SF-8) along with the uncoated SCG and reference open solar cell. We achieved excellent  $I$ - $V$  results through enhanced light trapping and charge carrier density using our optimized coated solar cover glass substrate as compared to the results obtained using commercially available ARC glasses on polycrystalline silicon solar cells. Therefore, the enhancement in the performance of the polycrystalline solar cells exceeded the gain in the light transmittance of the textured solar cover glass.

## 4. Conclusions

The present work describes the fabrication of mesoporous-silica-based transparent, hard, and antireflective (AR) coatings (thickness: 350–500 nm) with hydrophobic (WCA:  $124$ – $130^{\circ}$ ) surfaces on millimeter-scale textured solar cover glasses *via* a spray-coating technique. The coatings increased the photocurrent density ( $J_{sc}$ ) or power generation efficiency of solar modules and exhibited easy maintenance and/or self-cleaning

Table 4 Maximum photocurrent ( $I_{max}$ ), maximum voltage ( $V_{max}$ ), and maximum power ( $P_{max}$ ) output from a standard solar cell without cover glass, with commercial millimeter-scale textured cover glass, and with porous silica-coated commercial textured glass substrates under 1 sun for a projected illumination area of  $5 \times 5\text{ cm}^2$

Sample	$J_{sc}$ ( $\text{mA cm}^{-2}$ )	$V_{oc}$ (V)	$I_{max}$ (mA)	$V_{max}$ (V)	$P_{max}$ (mW)
Solar cell without cover glass (SCG) (1 sun on open solar cell)	46.57	2.32	163.65	1.70	277.62
Cell with SCG (uncoated) on the solar cell	43.00	2.32	155.12	1.70	263.16
Cell with SF-3 AR- and hydrophobic-coated SCG	44.85	2.32	159.90	1.70	271.71
Cell with SF-5 AR- and hydrophobic-coated SCG	44.88	2.32	160.10	1.70	271.71
Cell with SF-6 AR- and hydrophobic-coated SCG	45.00	2.32	161.63	1.70	274.20
Cell with SF-7 AR- and hydrophobic-coated SCG	44.66	2.32	162.20	1.70	275.17
Cell with SF-8 AR- and hydrophobic-coated SCG	45.20	2.32	159.71	1.70	270.94



properties. The bilayer-coated millimeter-scale textured solar glass showed an increase in average solar transmission of  $\geq 4\%$  due to the reduced reflection (average  $\sim 4\%$ ) in the wavelength range of 400–1100 nm. The maximum output current density ( $J_{sc}$ ) of the fabricated solar cell (compared to those available on the market) increased by  $>3$  to 4.6% (under 1 sun illumination), while the surface hydrophobicity increased after functionalization due to the micro/nanostructure of the AR coating fabricated on millimeter-scale solar glass. After 50-cycle sandpaper (80 mesh) abrasion testing, the WCA value was reduced from  $130^\circ$  to  $105^\circ$ , and the coating could withstand  $>50$  h UV radiation exposure testing, indicating the durability and good abrasion resistance properties of the coatings. Therefore, these AR and hydrophobic coatings on solar cover glass could be useful to trap more sunlight in silicon solar cells for higher conversion efficiencies, and they would be applicable in harsh and rough environments. Therefore, this cost-effective, highly abrasion-resistant AR and hydrophobic coating could be effectively utilized to achieve superior photo-power performance in solar cells to fulfill global energy demands.

## Conflicts of interest

The authors declare that they have no conflicts of interest.

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