


 Cite this: *RSC Adv.*, 2017, 7, 33757

 Received 8th June 2017
Accepted 27th June 2017

DOI: 10.1039/c7ra06444f

rsc.li/rsc-advances

Biomimetic nano/micro double-textured silicon with outstanding antireflective and super-hydrophilic surfaces for high optical performance

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We report the fabrication of nano/micro double-textured silicon (NMDT-Si) and its structural, optical, and surface wetting properties. The micro-pyramidal textured (MPTs) are formed on the Si surface by a simple potassium hydroxide-based wet etching process. On the other hand, for pillar-arrayed nano-textures (NTs), the thermally-dewetted gold (Au) nanoparticles are employed on the surface of the MPT-Si as an etch mask and the inductively coupled plasma etching is followed. The optical reflectance of the NMDT-Si is strongly dependent on the period and height of NTs on the surface of the MPT-Si, which can be controlled by the Au film thickness and etching time, respectively. Compared with the planar nano-textured Si, the NMDT-Si shows superior antireflection (or higher light absorption) and light-scattered propagation behaviors, which are verified from a finite-difference time-domain simulation, over wide ranges of wavelengths (350–1100 nm) and incident angles (0–70°), resulting in the average reflectance of ~2.1% and the solar weighted absorption of ~98.5% at normal incidence, respectively. In addition, it has a super-hydrophilic surface with water contact angles of <5°.

Introduction

Light management such as light scattering, antireflection, and light trapping has been widely researched for high performance of silicon (Si)-based optical and optoelectronic devices including solar cells, photodetectors, and image sensors.¹ For high optical performance, the surface modification of Si with high refractive indices of >3.5 at visible and near-infrared wavelengths is essential to suppress the high natural reflectivity (>30%) with strong spectral dependence for its planar surface. Generally, thin-film stacked antireflection coatings (ARCs) with a quarter-wavelength optical thickness and surface texturing technologies are widely used in industrial production of Si-based optical and optoelectronic devices. However, although the thin-film stacked ARC can reduce the surface reflection below 3–5%, it works only in narrow spectral bandwidth and limited incident angle, including additional drawbacks of stress caused by thermal expansion mismatch, adhesion to substrate, and material selection.² On the other hand, texturing easily forms micro-scale pyramidal arrays on the Si surface using an acid or alkaline (H₂SO₄, HNO₃:H₂O, NaOH, KOH, *etc.*) wet etching process,^{3,4} but their average reflectivity is kept above 10% though the micro-textures can lead to the strong light scattering, which is insufficient for high-efficiency Si-based optical and optoelectronic devices, especially, solar cells.⁵ As one of effective approaches, hierarchical double-scale architectures (*i.e.*, nano- and micro-

structures), like ‘moth eye’, create superior antireflection in wide ranges of wavelengths and incident angles as well as self-cleaning ability.^{6–8} However, the reflection of nano/micro double-textured surfaces is mainly dependent on the shape, period, and height of nano-structures.⁹ Thus, it is necessary to optimize the period and size of nano-patterns and the etching conditions for efficient antireflection. The nano-structures on micro-pyramidal arrays have been realized using a variety of nano-texturing techniques, such as cesium chloride (CsCl) self-assembly lithography (*i.e.*, nano-islands), metal (*e.g.*, silver, gold, nickel, *etc.*)-assisted chemical etching (catalyst nanoparticles), mask-free plasma (SF₆/O₂) surface processing, and nano-porous aluminum etch mask.^{6,10} Recently, a thermal-dewetting process of metal films as cost-effective and facile lithography has been widely employed to form the nano-patterns (*i.e.*, metallic nano-sized cluster or particles) as a dry-etch mask.^{11,12} In this nano-patterning, the period and size of metallic nano-patterns are determined by the thickness of metal films.^{11,13}

Meanwhile, the surface wettability is also important in outdoor applications since a super-hydrophobic or super-hydrophilic structure can remove dust particles or any contaminants on its surface by water droplets (*i.e.*, self-cleaning), which can maintain the device efficiency by preventing the degradation of optical performance.¹⁴ The super-hydrophobic or super-hydrophilic surface can be obtained by introducing the high roughness into the planar surface.^{15,16}

In this work, we fabricated the nano/micro-structured Si using double-texturing processes. To form the micro-pyramidal

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textures (MPTs) on the Si surface, the chemical wet etching was carried out using a mixture solution of potassium hydroxide (KOH) and isopropyl alcohol (IPA), forming micro-scale pyramidal arrays, because of its good control and reproducibility. The nano-textures (NTs) were fabricated by the Au film thermal-dewetting and dry etching processes, subsequently. For comparison, the nano-texturization was also treated on the planar Si surface from the same nano-texturing fabrication process. For all the samples, the structural and optical properties were systematically studied by adjusting the Au film thickness and etching time. Angle-dependent reflectance characteristics were explored under s-, p-, and un-polarized lights. To theoretically analyze the optical light-scattering propagation inside Si, we conducted finite-difference time-domain (FDTD) simulations. Moreover, the surface wettability was also investigated.

Experimental and simulation modeling details

Fabrication of the NMDT-Si

Fig. 1 shows the schematic illustration of process steps for the fabrication of double-textured Si consisting of nano-structures on micro-pyramidal arrays (NMDT-Si) using a combination of wet and dry etching processes. The scanning electron microscope (SEM) image of the micro-pyramidal textured Si (MPT-Si) is also shown in Fig. 1. In this experiment, single-side polished Si substrates with a square area of $2 \times 2 \text{ cm}^2$ were cleaned with acetone, methanol, and de-ionized water for 10 min, subsequently. And then, we further performed a plasma (ozone) cleaning process to remove the impurities and contaminants on the Si surface for 10 min. To fabricate the MPT-Si, we simply soaked the Si substrates in a mixed solution of KOH : IPA (10 : 3 vol%) at a temperature of 75°C for 40 min, followed by dipping the samples into a $\text{HCl} : \text{H}_2\text{O}_2 : \text{H}_2\text{O}$ (1 : 1 : 5) mixed solution at 80°C for 10 min for removal of the remaining potassium impurities on the surface of wet-etched samples. As shown in the SEM image of Fig. 1, the randomly-distributed micro-pyramidal arrays are well formed on the Si surface, indicating the height and (bottom) size in ranges of $\sim 5.5 \pm 3$ and $\sim 5 \pm 3 \mu\text{m}$ (mean \pm standard deviation), respectively. For the formation of NTs with pillar arrays, we deposited Au films with different thicknesses of 5, 7, and 9 nm on the surface of MPT-Si *via* a thermal evaporator, respectively. During the deposition, the thickness of Au films was controlled with material-related calibration by using a quartz crystal thickness monitoring.¹⁷ And then, the samples were thermally treated to form dot-like Au nanoparticles (Au-NPs) at temperatures of 500, 600, and 700°C for 2 min in a rapid thermal annealing system, respectively. Using the Au-NPs as an etch mask, the double-textured Si (*i.e.*, NMDT-Si) with a combination of nano-pillars and micro-pyramidal arrays was fabricated by an inductively coupled plasma (ICP) etching process at radio frequency (RF) power of 50 W and additional ICP power of 200 W under a process pressure of 5 mTorr in $\text{CF}_4 : \text{Ar}$ (20 : 50 sccm) plasma. After the ICP etching, the remaining Au-NPs were removed using an Au etchant solution mixed by potassium iodide and iodine (KI/I_2).

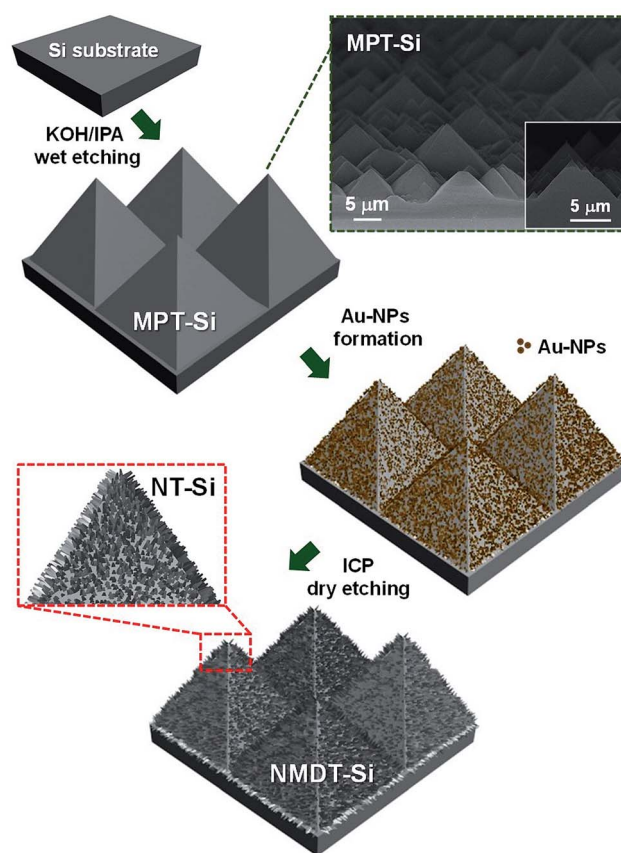


Fig. 1 Schematic illustration of process steps for the fabrication of double-textured Si consisting of NMDT-Si using a combination of wet and dry etching processes. SEM image of the MPT-Si is also shown. In this experiment, the KOH/IPA-mixed solution for the MPT-Si and the thermally-dewetted Au-NPs/ICP etching for the nano-textured Si (NT-Si) are used, respectively.

Characterization

The etched profiles and surface morphologies of samples were observed from a SEM (LEO SUPRA 55, Carl Zeiss). The optical total (specular + diffuse) reflectance was measured by using a UV-vis-NIR spectrophotometer (Cary 5000, Varian) with an integrating sphere at near normal incidence. Spectroscopic ellipsometry (V-VASE, J. A. Woollam Co. Inc.) was used to examine the angle-dependent specular reflectance at incident angles (θ_{in}) of $20\text{--}70^\circ$ for s-, p-, and un-polarized lights. The surface wetting behaviors were characterized by measuring contact angles at five different positions on the surface of samples *via* a contact angle measurement system (Phoenix-300, SEO Co., Ltd.) and the values were averaged.

Simulation modeling

To design the numerical models, the nano-pillars and micro-pyramids on the Si surface were roughly represented by a periodic geometry in the Cartesian coordinate system by a scalar-valued function of two variables, $f(x, z)$, for simplicity. The light, which is normalized at a wavelength (λ) of 530 nm for un-



polarization, entered from air into the textured Si surface at normal incidence. The amplitude of y-polarized electric field (*i.e.*, E_y) was calculated for the incident plane wave with a slab mode beam profile under a perfectly-matched-layer boundary condition at a grid size of 5 nm. The height and period of models were kept at 300 and 100 nm for nano-pillars and 5.5 and 5 μm for micro-pyramids, respectively. The optical parameters (*i.e.*, refractive index and extinction coefficient) of the Si used in these calculations were acquired from the refractive index Web site.¹⁸

Results and discussion

Thermally-dewetted Au-NPs on the MPT-Si

Fig. 2 shows (a–c) the SEM images of the thermally-dewetted Au-NPs on the surface of MPT-Si for different thicknesses (5, 7, and 9 nm) of Au films and (d) the estimated average period, size, and density of Au-NPs for the corresponding Au films. As shown in the SEM images of Fig. 2(a–c), the dot-like Au-NPs are uniformly formed on the MPT-Si surface by the thermal-dewetting process. During the dewetting process, the Au films with very thin thicknesses are agglomerated into the nano-sized clusters or particles due to the concurrent effects of surface diffusion, surface energy minimization, coalescence, and Ostwald's ripening.¹⁹ In Fig. 2(d), distributions of the Au-NPs are considerably changed as the thickness of Au film is increased, indicating average densities of ~ 60 , 52, and 44% for 5, 7, and 9 nm, respectively, which are characterized using a public domain image processing program (ImageJ 1.42q, NIH). On the other hand, the correlation distance (*i.e.*, period) between the adjacent Au-NPs and their size are roughly estimated to be $\sim 50 \pm 15$, 90 ± 45 , and 200 ± 50 nm and $\sim 30 \pm 10$, 50 ± 15 , and 90 ± 40 nm for 5, 7, and 9 nm from the corresponding SEM images of

Fig. 2(a–c), respectively. Here, we measured the period and size of Au-NPs at least 15 different positions on the SEM image and calculated the mean (*i.e.*, $\mu = \sum x/n$) and standard deviation (*i.e.*, $\sigma = [\sum (x - \mu)^2/n]^{1/2}$) values (rounded off) for each Au film.

Structural and optical properties of the NMDT-Si

Fig. 3 shows (a–c) the SEM images of NMDT-Si samples for 5, 7, and 9 nm-thick Au films at an etching time of 10 min and (d) the total reflectance spectra of the corresponding samples. For comparison, the reflectance spectra of bare Si and MPT-Si are also shown in the inset of Fig. 3(d). It can be evidently observed that the patterns of Au-NPs are directly transferred onto the surface of MPT-Si by dry etching, which creates the NMDT-Si, as can be seen in Fig. 3(a–c). With increasing the thickness of Au film, the morphology of NTs on the surface of the MPT-Si is changed from the truncated conical pillars to the cylindrical ones, while maintaining the height of $\sim 200 \pm 20$ nm without period variation for each Au film. This is the reason why the smaller-sized NPs generated from thinner Au films (*e.g.*, 5 nm) are further quickly eliminated at the edge of NPs compared to the thicker and larger ones during the dry etching. To investigate the effect of NTs with different periods on the reflectance of NMDT-Si samples, their optical reflectance properties were measured in the wavelength range of 350–1100 nm. As shown in Fig. 3(d), the bare Si shows a high surface reflectivity of $>30\%$ at wavelengths of 350–1100 nm owing to the large difference in refractive indices between air and the Si, indicating the average reflectance (R_{avg}) of $\sim 38.5\%$. However, the MPTs decrease the reflectance of the bare Si to be $R_{\text{avg}} \sim 20.6\%$ due to the linearly effective graded-refractive-index (GRIN) profile from air to the Si and enhanced light-path lengths caused by the pyramidal micro-structure.^{20,21} Clearly, an introduction of subwavelength-scale textures (*i.e.*, NTs) into the surface of the MPT-Si leads to the further

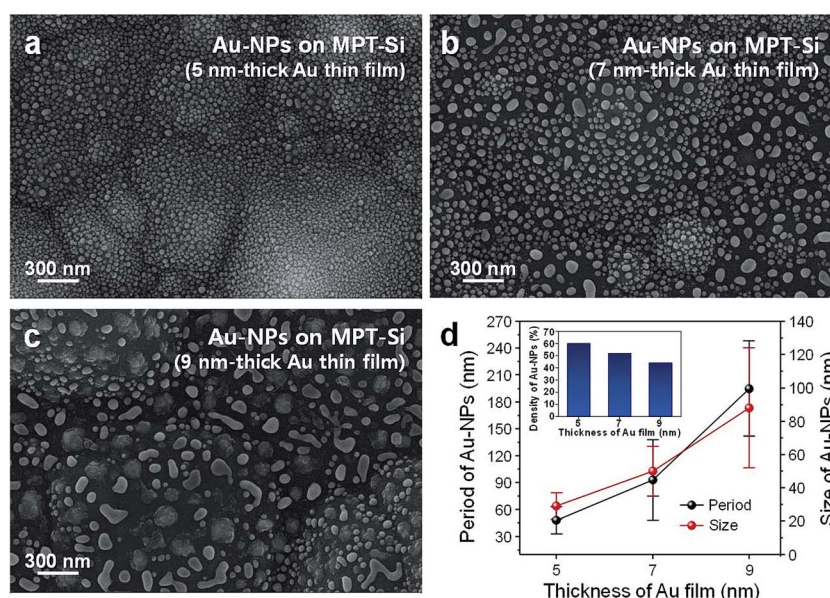


Fig. 2 (a–c) SEM images of the thermally-dewetted Au-NPs on the surface of MPT-Si samples for different thicknesses (5, 7, and 9 nm) of Au films and (d) estimated average period, size, and density of Au-NPs for the corresponding Au films.



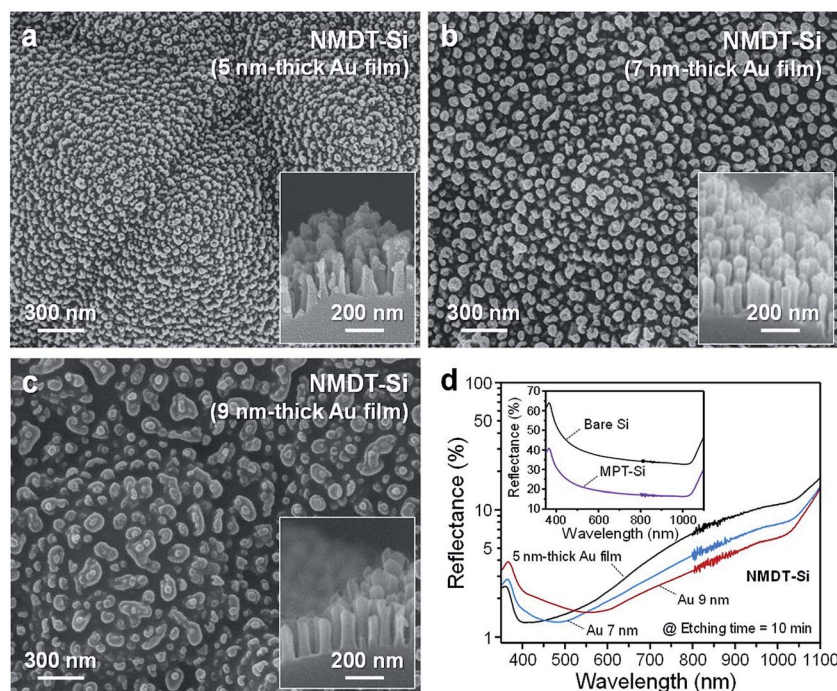


Fig. 3 (a–c) SEM images of NMDT-Si samples for 5, 7, and 9 nm-thick Au films at an etching time of 10 min and (d) total reflectance spectra of the corresponding samples. For comparison, the reflectance spectra of bare Si and MPT-Si are also shown in the inset of (d).

suppressed reflectivity over a broad wavelength range of 350–1100 nm since they cause not only the additional linearly effective GRIN distribution between air and the surface of MPT-Si, but also allow the only zeroth-order diffracted wave to propagate through the surface of NTs, known as ‘moth eye effect’.²² As the thickness of Au film is increased from 5 to 9 nm, the low reflectance band of the NMDT-Si is shifted towards the longer wavelength region because the period of the NTs becomes larger (*i.e.*, $\sim 50 \pm 15$, 90 ± 45 , and 200 ± 50 nm for 5, 7, and 9 nm, respectively). Generally, when the grating period becomes much smaller than the wavelength of the incident light, a zeroth-order diffracted light wave is only allowed to reflect (or penetrate) on the grating structure and all the higher orders of diffracted light waves become evanescent, leading to the superior antireflection effect.^{23,24} This result implies that the low reflectance in a specific wavelength range can be obtained by adjusting the period of NTs (*i.e.*, Au film thickness). For the NMDT-Si at 7 nm-thick Au film, the lowest R_{avg} of $\sim 4\%$ is obtained compared to the other samples ($R_{\text{avg}} \sim 5.9$ and 4.5% for 5 and 9 nm, respectively).

We also investigate the influence of NTs with different heights on the reflectance of NMDT-Si samples. Fig. 4 shows (a–c) the SEM images of NMDT-Si samples at different etching times of 5, 15, and 20 min for 7 nm-thick Au film and (d) the total reflectance spectra of the corresponding samples. For comparison, the SEM image and total reflectance spectrum of the planar NT-Si at 15 min are also shown. As shown in Fig. 4(a–c), the height of NTs on the MPT-Si surface is increased from 100 ± 20 nm at 5 min to 300 ± 30 nm at 15 min while it is contrastively decreased to be 280 ± 40 nm at 20 min, showing the partly collapsed nano-pillars with uneven distributions. This is attributed to the absence of Au-NPs (*i.e.*, complete

removal), which are used as an etch mask, at longer etching times. Additionally, the shape of NTs on the MPT-Si surface is varied from the truncated cone at 5 min to the cone at 15 min. As a result, the NMDT-Si consisting of conical NTs with larger heights at 15 min shows the lowest reflectance spectrum ($R_{\text{avg}} \sim 2.1\%$) compared to the other samples ($R_{\text{avg}} \sim 10$ and 2.7% at 5 and 20 min) over a wide wavelength range of 350–1100 nm, as shown in Fig. 4(d). Furthermore, it also exhibits the superior antireflection property in comparison with the planar NT-Si ($R_{\text{avg}} \sim 6\%$). The outstanding antireflection property can be also confirmed from the photograph in the inset of Fig. 4(d). The fluorescent white light is strongly reflected at the surface of the bare Si. For the planar NT-Si, the faint reflected fluorescent white light is also seen on its surface in spite of a black surface. However, there is no any reflected fluorescent white light on the NMDT-Si with a dark black surface.

For Si-based solar cell applications, it is necessary to explore the percentage of the incoming solar energy, which is absorbed by the Si, by obtaining the solar weighted absorption (A_{SW} , ratio of the usable photons absorbed to the total usable photons). The A_{SW} can be estimated by normalizing the absorption spectrum with the solar spectral photon flux (*i.e.*, air mass 1.5 global) integrated over a wavelength range of 350–1000 nm.²⁵ The A_{SW} is given by the following equation:

$$A_{\text{SW}} = \frac{\int_{350 \text{ nm}}^{1000 \text{ nm}} S(\lambda) a(\lambda) d\lambda}{\int_{350 \text{ nm}}^{1000 \text{ nm}} S(\lambda) d\lambda}, \quad (1)$$

where $S(\lambda)$ is the spectral photon flux and $a(\lambda)$ is the absorption (100 reflectance–transmittance) of samples at normal



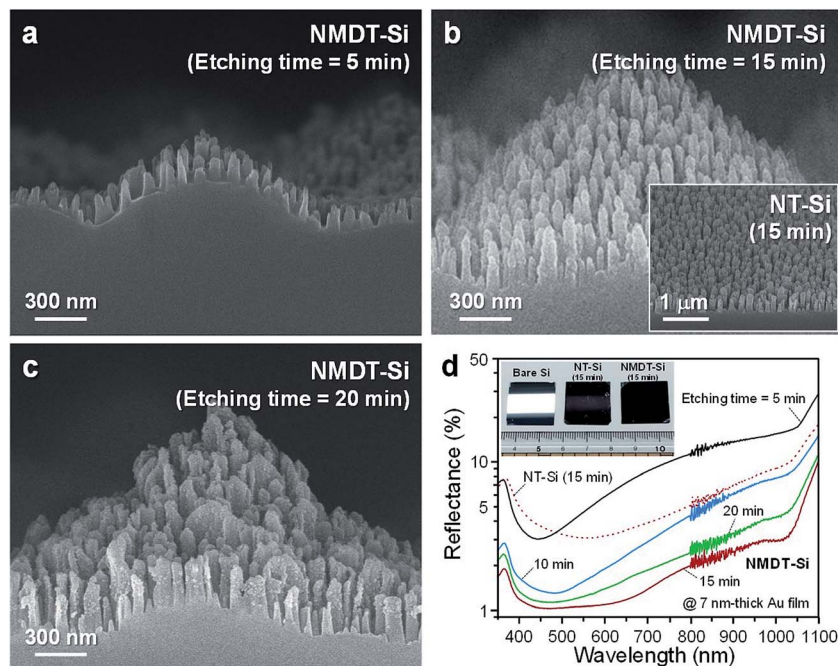


Fig. 4 (a–c) SEM images of NMDT-Si samples at different etching times of 5, 15, and 20 min for 7 nm-thick Au film and (d) total reflectance spectra of the corresponding samples. For comparison, SEM image and total reflectance spectrum of the planar NT-Si at 15 min are also shown in (b) and (d), respectively. The inset of (d) shows the photographic image of bare Si, NT-Si, and NMDT-Si samples.

incidence. Here, absorption data are not shown and transmittance is zero. As expected, for the NMDT-Si at 15 min, the highest A_{SW} of $\sim 98.5\%$ is obtained compared to the other NMDT-Si samples ($A_{SW} \sim 92.4, 97.2$, and 98% at 5, 10, and 20 min, respectively) as well as the bare Si ($A_{SW} \sim 61.3\%$), the MPT-Si ($A_{SW} \sim 79.6\%$), and the NT-Si ($A_{SW} \sim 95.7\%$). These results suggest that the NMDT-Si with the superior antireflection ability can be useful to realize high-performance Si-based photovoltaic systems.

Light-scattering property of the NMDT-Si

To study the light-scattering propagation of the NMDT-Si with micro-features, the numerical optical analysis was performed using the FDTD method. Fig. 5 shows the contour plots of calculated electric-field intensities for optical light-propagation properties of bare Si, NT-Si, and NMDT-Si *via* FDTD simulations at $\lambda = 530$ nm. As shown in Fig. 5, for both the bare Si and NT-Si, there are no scattered propagating lights inside Si. On the other hand, the NMDT-Si with micro-pyramidal arrays has a strong light scattering property with the wide angular spread due to the extended effective optical path lengths caused by the rebound lights between the micro-pyramids. This implies that the probability of photons transmitted through the surface of the NMDT-Si is increased instead of escaping back to air.^{6,21}

Angle-dependent reflectance characteristics of the NMDT-Si

For high-performance devices in optical and optoelectronic applications, the good antireflective characteristics should be obtained in wide angles of incident light for both s- and p-polarized lights. Fig. 6 shows the angle-dependent specular

reflectance characteristics of bare Si, NT-Si, and NMDT-Si samples at $\theta_{in} = 20\text{--}70^\circ$ for (a) s-, (b) p-, and (c) un-polarized lights. Here, the NT-Si and NMDT-Si samples fabricated at 7

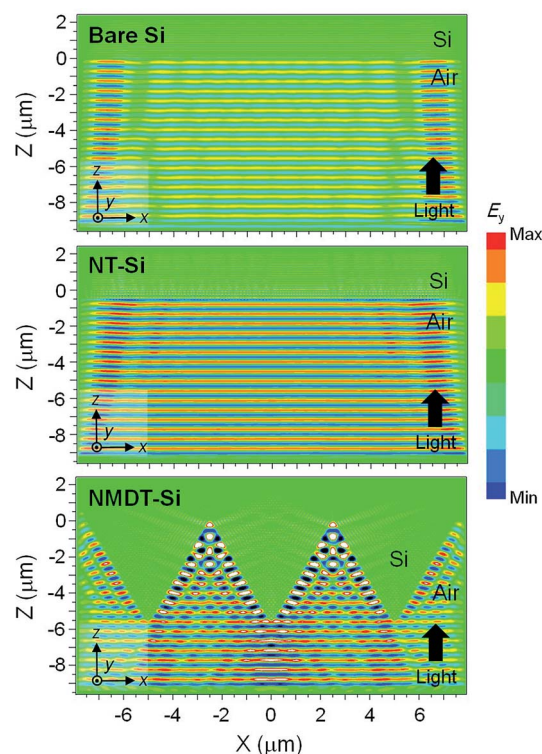


Fig. 5 Contour plots of calculated electric-field intensities for optical light-propagation properties of bare Si, NT-Si, and NMDT-Si *via* numerical FDTD calculations at $\lambda = 530$ nm.



nm-thick Au film and 15 min-etching time were examined. The bare Si shows strong variations with angles of light incidence for both s- and p-polarized lights. On the other hand, the NT-Si is not relatively sensitive to polarization at θ_{in} values lower than 50° , while exhibiting the increased reflectivity with fluctuations

at $\theta_{\text{in}} \geq 60^\circ$. However, the NMDT-Si has much less angle-dependent reflectance characteristics over wide ranges of wavelengths (350–1100 nm) and incident angles (20–70°), maintaining the reflectance values below 1% for both s- and p-polarized lights. For the un-polarized light in Fig. 6(c), it shows the much lower R_{avg} of $\sim 0.06\%$ compared to the bare Si ($R_{\text{avg}} \sim 36.07\%$) and the NT-Si ($R_{\text{avg}} \sim 2.22\%$) in the wavelength range of 350–1100 nm at $\theta_{\text{in}} = 20\text{--}70^\circ$.

Surface wettability

Finally, the surface wetting behaviors of the highly-textured NMDT-Si were studied, compared with the bare Si, MPT-Si, and NT-Si samples. Fig. 7 shows the photographic images of a droplet on the surface of (a) bare Si, (b) MPT-Si, (c) NT-Si, and (d) NMDT-Si samples with each static contact angle value. For both the NT-Si and NMDT-Si at 7 nm-thick Au film and 15 min-etching time, the time-sequential photographic images are also shown. The surface water contact angle (θ_{CA}) of MPT-Si is slightly reduced to be $\sim 32.5^\circ$ compared to that ($\theta_{\text{CA}} \sim 41.2^\circ$) of bare Si, exhibiting the hydrophilic surface. On the other hand, both the NT-Si and NMDT-Si show strong hydrophilic properties with θ_{CA} values of $< 5^\circ$. As shown in Fig. 7(c) and (d), a water droplet is immediately spread on the surface like a water sheet film after dropping within 1 s. This complete wetting is attributed to the considerably increased surface roughness due to the NTs, which can be explained by the Wenzel's equation though it

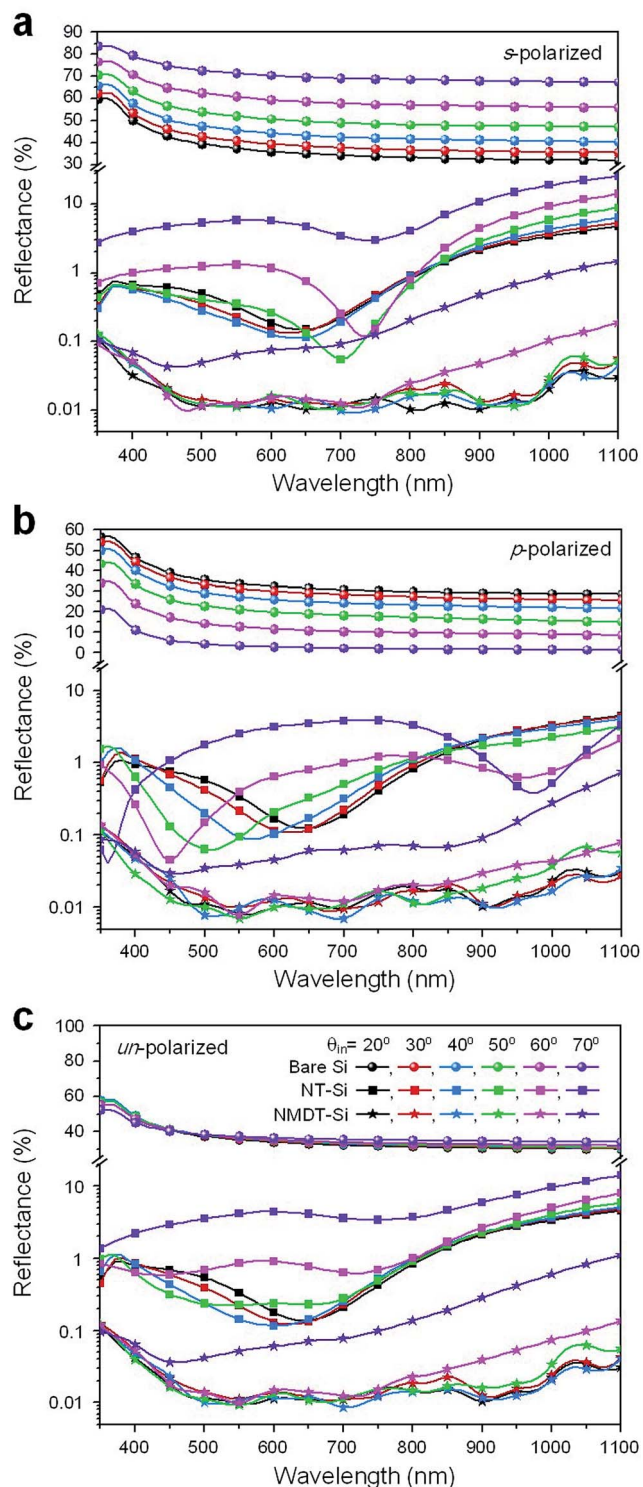


Fig. 6 Angle-dependent specular reflectance characteristics of bare Si, NT-Si, and NMDT-Si samples at $\theta_{\text{in}} = 20\text{--}70^\circ$ for (a) s-, (b) p-, and (c) un-polarized lights.

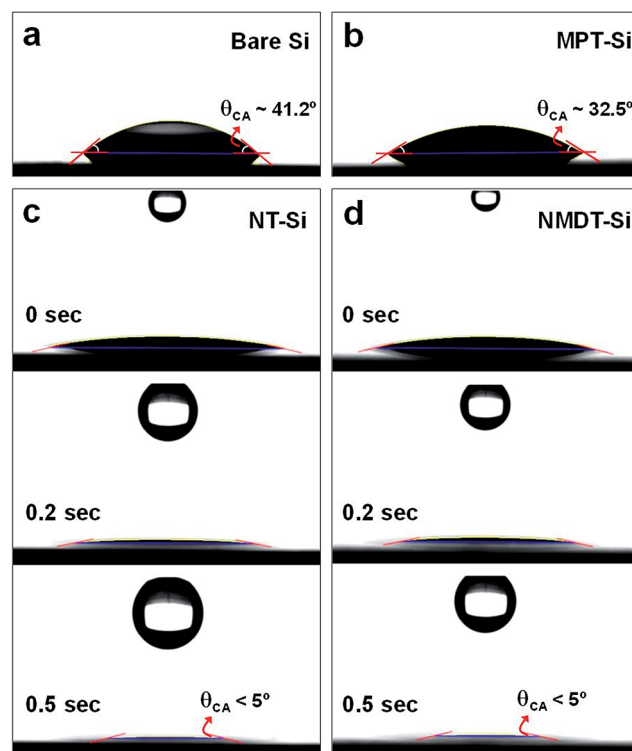


Fig. 7 Photographic images of a droplet on the surface of (a) bare Si, (b) MPT-Si, (c) NT-Si, and (d) NMDT-Si samples with each static contact angle value. For both the NT-Si and NMDT-Si at 7 nm-thick Au film and 15 min-etching time, the time-sequential photographic images are also shown.



is also related to the surface energy of materials and the plasma (ozone) treatment.^{16,26} The dust particles or contaminants on the super-hydrophilic surface can be easily removed from the surface by the flowing water droplets which can squeeze into the space between the dusts and the surface. Therefore, the NMDT-Si may self-clean the dust particles and quickly dry the rainwater on the surface of Si-based optical and optoelectronic devices in outdoor applications.

Conclusions

The outstanding antireflective and hydrophilic architected Si was fabricated by the nano/micro double-texturing processes. The NMDT-Si was realized by the KOH-based wet etching for the MPTs and the ICP etching *via* the thermally-dewetted Au-NPs for NTs and their surface morphologies and optical reflectance properties were investigated, together with surface wettability. For the NMDT-Si at 7 nm-thick Au film and 15 min-etching time, surpassing antireflection performance was obtained in broad ranges of wavelengths (350–1100 nm) and incident angles (0–70°) compared to the planar NT-Si as well as the bare Si and the MPT-Si, resulting in the R_{avg} values of $\sim 2.1\%$ and $\sim 0.06\%$ at $\theta_{\text{in}} = 0^\circ$ and $20\text{--}70^\circ$, respectively. In addition, it strongly scattered the lights penetrated through the surface of NMDT-Si, as confirmed by numerical FDTD simulations. It also had a super-hydrophilic surface with θ_{CA} values of $< 5^\circ$. From these results, the nano/micro double-textured Si with high optical light absorption (*i.e.*, efficient antireflection/light-scattering) and self-cleaning ability can be very useful for high-performance Si-based optical and optoelectronic devices.

Acknowledgements

This work was supported by the National Research Foundation of Korea (NRF) Grant funded by the Korea government (MSIP) (No. 2017R1A2B4011998).

Notes and references

- 1 M. L. Brongersma, Y. Cui and S. Fan, *Nat. Mater.*, 2014, **13**, 451; S. F. Leung, Q. Zhang, F. Xiu, D. Yu, J. C. Ho, D. Li and Z. Fan, *J. Phys. Chem. Lett.*, 2014, **5**, 1479; J. W. Leem, M. Choi, D. Bhaskar and J. S. Yu, *RSC Adv.*, 2016, **6**, 55159.
- 2 N. Kadakia, S. Naczas, H. Bakhru and M. Huang, *Appl. Phys. Lett.*, 2010, **97**, 191912; C. L. Tien and T. W. Lin, *Appl. Opt.*, 2012, **51**, 7229.
- 3 H. Seidel, L. Csepregi, A. Heuberger and H. Boumgärtel, *J. Electrochem. Soc.*, 1990, **137**, 3612; T. Rahman, M. Navarro-Cía and K. Robelet, *Nanotechnology*, 2014, **25**, 485202.
- 4 B. Dudem, J. H. Heo, J. W. Leem, J. S. Yu and S. H. Im, *J. Mater. Chem. A*, 2016, **4**, 7573.
- 5 D. Qi, N. Lu, H. Xu, B. Yang, C. Huang, M. Xu, L. Gao, Z. Wang and L. Chi, *Langmuir*, 2009, **25**, 7769; Y. Jiang, X. Zhang, F. Wang and Y. Zhao, *RSC Adv.*, 2015, **5**, 69629.
- 6 M. Pranaitis, L. Jaraminè, V. Čyras, A. Selskis and A. Galdikas, *J. Appl. Phys.*, 2013, **114**, 163523; Y. Song, K. Kim, K. Choi, B. Ki and J. Oh, *Sol. Energy*, 2016, **135**, 291.
- 7 W. R. Wei, M. L. Tsai, S. T. Ho, S. H. Tai, C. R. Ho, S. H. Tsai, C. W. Liu, R. J. Chung and J. H. He, *Nano Lett.*, 2013, **13**, 3658.
- 8 B. Dudem, J. W. Leem, J. H. Lim, S. H. Lee and J. S. Yu, *RSC Adv.*, 2015, **5**, 90103; B. Dudem, Y. H. Ko, J. W. Leem, J. H. Lim and J. S. Yu, *ACS Appl. Mater. Interfaces*, 2016, **8**, 30165; B. Dudem, J. W. Leem and J. S. Yu, *RSC Adv.*, 2016, **6**, 3764.
- 9 S. A. Boden and D. B. Bagnall, *Appl. Phys. Lett.*, 2008, **93**, 133108; Y. M. Song, H. J. Choi, J. S. Yu and Y. T. Lee, *Opt. Express*, 2010, **18**, 13063; J. W. Leem, S. Kim, S. H. Lee, J. A. Rogers, E. Kim and J. S. Yu, *Adv. Energy Mater.*, 2014, **4**, 1301315.
- 10 J. Liu, M. Ashmkhan, G. Dong, B. Wang and F. Yi, *Sol. Energy Mater. Sol. Cells*, 2013, **108**, 93; Y. Cao, A. Liu, H. Li, Y. Liu, F. Qiao, Z. Hu and Y. Sang, *Appl. Surf. Sci.*, 2011, **257**, 7411; J. Liu, B. Liu, S. Liu, Z. Shen, C. Li and Y. Xia, *Surf. Coat. Technol.*, 2013, **229**, 165; J. Liu, T. Zhang, G. Dong, X. Zhang, B. Wang, Y. Liao and F. Yi, *J. Phys. D: Appl. Phys.*, 2013, **46**, 375302.
- 11 J. W. Leem, J. S. Yu, Y. M. Song and Y. T. Lee, *Sol. Energy Mater. Sol. Cells*, 2011, **95**, 669.
- 12 C. I. Yeo, Y. M. Song, S. J. Jang and Y. T. Lee, *Opt. Express*, 2011, **19**, A1109; J. H. Lim, J. W. Leem and J. S. Yu, *RSC Adv.*, 2015, **5**, 61284.
- 13 G. C. Park, Y. M. Song, J. H. Ha and Y. T. Lee, *J. Nanosci. Nanotechnol.*, 2011, **11**, 6152.
- 14 Y. B. Park, H. Im, M. Im and Y. K. Choi, *J. Mater. Chem.*, 2011, **21**, 633; J. Son, S. Kundu, L. K. Verma, M. Sakhuja, A. J. Danner, C. S. Bhatia and H. Yang, *Sol. Energy Mater. Sol. Cells*, 2012, **98**, 46; J. W. Leem and J. S. Yu, *RSC Adv.*, 2015, **5**, 60804.
- 15 A. B. D. Cassie and S. Baxter, *Trans. Faraday Soc.*, 1944, **40**, 546.
- 16 R. N. Wenzel, *Ind. Eng. Chem.*, 1936, **28**, 988.
- 17 J. W. Leem, Y. M. Song and J. S. Yu, *Opt. Express*, 2011, **19**, 26308.
- 18 SOPRA, N&K Database (2017), retrieved 2017, <http://refractiveindex.info>.
- 19 P. L. Redmond, A. J. Hallock and L. E. Brus, *Nano Lett.*, 2005, **5**, 131; G. Palasantzas, T. Vystavel, S. A. Koch and J. T. M. De Hosson, *J. Appl. Phys.*, 2006, **99**, 024307; H. Chen, Y. Yu, H. L. Xin, K. A. Newton, M. E. Holtz, D. Wang, D. A. Muller, H. D. Abruña and J. DiSalvo, *Chem. Mater.*, 2013, **25**, 1436.
- 20 W. H. Southwell, *J. Opt. Soc. Am. A*, 1991, **8**, 549.
- 21 J. Zhao and M. A. Green, *IEEE Trans. Electron Devices*, 1991, **38**, 1925.
- 22 P. B. Clapham and M. C. Hutley, *Nature*, 1973, **244**, 281.
- 23 E. Hecht, *Optics*, Addison-Wesley, NY, USA, 4th edn, 2002, ch. 10.
- 24 H. Kikuta, H. Toyota and W. Yu, *Opt. Rev.*, 2003, **10**, 63.
- 25 J. W. Leem, S. Kim, C. Park, E. Kim and J. S. Yu, *ACS Appl. Mater. Interfaces*, 2015, **7**, 6706.
- 26 L. Schneider, M. Laustsen, N. Mandsberg and R. Taboryski, *Sci. Rep.*, 2016, **6**, 21400.

