# Broadband angular colour stability of dielectric thin film-coated pyramidal textured Si for photovoltaics

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In this work we demonstrate the angular colour stability of textured c-Si substrates coloured by single layer thin film coatings of SiN<sub>x</sub>. These coatings show higher angular colour stability on substrates with a random upright pyramidal surface texture compared to identical coatings on planar silicon substrates. Angle dependent reflectance measurements, supported by a modeling framework, display that the reflectance peaks originating from thin film interference of coated textured substrates only shift about 15 nm with an increasing angle of incidence from 10 to 80°, while the reflectance peaks of planar substrates with identical coatings shift about 120 nm at these angles. More specifically, reflectance peaks of planar substrates shift to shorter wavelengths, leading to a blue shift of the colour appearance. The stable peak position of the textured samples is explained by a 2D representation of their surface texture and the primarily double interference interaction on it. While it is well known that a wide range of colours can be realized exhibiting low optical losses with thin film coatings, angular colour stability was often not taken into account. However, for building integrated photovoltaics applications, a high angular colour stability is desired, underlining the importance of using these textures. In most installed c-Si photovoltaics, similar substrate surface textures and dielectric thin film layers are already used. Therefore, this work envisions a route to facilitate large scale production of coloured solar cells on textured c-Si substrates, coloured by thin film SiN<sub>x</sub> layers, with minimized optical losses and improved angular colour stability.

## I. INTRODUCTION

Two key aspects of building-integrated photovoltaic (BIPV) elements are their aesthetics and power output. Ideally, BIPV elements are desired components of a building from an aesthetic point of view, while also producing as much power as possible. In the case of coloured BIPV, these two aspects are often conflicting, as light in the visible range needs to be reflected to create colour. Since this light would otherwise be transmitted to the active layers of the cell, colouring a BIPV product decreases its power output. It is thus an important challenge to create solar cells with a wide range of possible colours at minimized optical losses.

As both, the PV and BIPV market, are dominated by crystalline silicon (c-Si) technologies,<sup>1</sup> it is especially important to create inexpensive and efficient ways to colour c-Si based solar cells. Most c-Si based coloured BIPV products that are currently present on the market are based on printing techniques.<sup>1</sup> Although inexpensive and easily integrated into the PV manufacturing process, these techniques come with high optical losses.<sup>2</sup> A more efficient way to colour c-Si cells is via deposition of thin film coatings on top of the silicon substrate. Such coatings have spectrally selective reflectance spectra due to interference. By varying the thickness and refractive index of this film, a wide variety of colours with high transmission into the active layers of the cell can be achieved.<sup>3–8</sup> In particular, films made of silicon nitride (a-SiN $_x$ :H, also called SiN $_x$ ), deposited by plasma-enhanced chemical vapour deposition (PECVD), allow for immediate large scale production of coloured solar cells, as they are already deposited onto c-Si solar cells in standard manufacturing as the anti-reflection coating (ARC). This is a major benefit of this technique over a variety of other novel solar cell colouring techniques based on for example scatterers,<sup>9,10</sup> plasmonic structures,<sup>11</sup> and other nanostructures,<sup>12–14</sup> which have also shown to colour solar cells with little optical losses, but are not yet ready to be implemented on a factory scale.

There exists extensive research on colouring with thin films, but little focus has been dedicated to the angular colour stability of these coatings, which is important to the BIPV market. When using this colouring method on planar substrates, the colour appearance of the film is strongly influenced by the angle of incidence of incoming light, as this determines the effective thickness of the thin film.

The great majority of installed c-Si PV is based on substrates that are not planar, but have some surface texture. In the case of monocrystalline silicon, a random upright pyramid texture, created by wet alkaline etching, is most commonly used.<sup>15–18</sup>

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This texture reduces reflectance and increases light trapping, and thus contributes to an increase in the current output of the solar cell<sup>15,19–21</sup> and is also cheaper to realize than a perfectly planar substrate surface.

In this paper, it is shown that such substrates coloured with single layer thin film coatings possess better angular colour stability than planar substrates coated with identical coatings. Experimental measurements of angle dependent reflectance of the samples are supported by modeled spectra, and the angular stability of the textured surfaces is explained using a 2D representation of the surface texture. In total, four different thin film coatings, each with a different colour, were deposited and compared. As the conclusions are the same for all four films, only the full analysis of one of these, a film with a green colour appearance, called sample G1, will be shown here. More information on the other samples can be found in the Supplementary Material (SM).

# **II. THEORETICAL METHODS**

Because reflected light from a single thin film on top of a substrate consists of two beams, interference of light occurs. By changing thickness or index of the thin film, the wavelengths at which this happens can be tuned and the colour appearance of the film can be modified. For constructive interference to occur at wavelength  $\lambda$ , the optical path difference between the two reflected beams should be equal to  $m_1 \cdot \lambda$ , with  $m_1 \in \mathbb{N}$ . Therefore, one can easily derive that for a planar thin film of thickness *d*, constructive interference occurs at wavelengths

$$\lambda_{max} = \frac{2n_1 d}{m_1} \sqrt{1 - \left(\frac{n_0}{n_1} \sin \theta\right)^2} \tag{1}$$

where  $n_0$  and  $n_1$  are the refractive indices of the surrounding medium, in this work air, and the thin film layer, respectively, and  $\theta$  is the angle of incidence on the thin film. To determine at which wavelengths destructive interference occurs, one can simply substitute  $m_1$  for  $(m_2 + \frac{1}{2})$  with  $m_2 \in \mathbb{N}_0$ . From this follows that as  $\theta$  increases, the positions of the reflectance minima and maxima shift towards bluer wavelengths, impacting the colour appearance of the thin film. For substrates with a random upright pyramid texture, reflected light is made up of beams following a large amount of discrete paths, which each have one or multiple interactions with the substrate under different angles of incidence.<sup>15</sup> A change in the angle of incidence does therefore not necessarily result in a shift in the wavelengths of minimum and maximum reflectance.

In this work, the measured reflectance spectra of planar thin films are compared to modeled reflectance spectra, based on the model created by Haug et al.<sup>22</sup>, which uses the transfer matrix method. As the reflectance of textured substrates cannot be modeled in this way, the online ray tracing tool OPAL  $2^{23,24}$  was used to model the reflectance spectra of the textured substrates. It is not expected that the reflectance modeled by ray tracing will be exactly the same as what is measured, as such models often assume an ideal upright pyramid surface texture, where all pyramids have the same base angle. It has been observed, however, that the pyramid base angles often have a distribution of values below the theoretical angle of 54.7° as a result of non-ideal etching.<sup>16–18,25</sup> Furthermore, it has been shown that the surface texture can have features of hillocks with an orthogonal base, rather than of pyramids, making ray tracing approximations less exact.<sup>25</sup> Lastly, scattering and/or diffraction effects at pyramids with a size around the wavelength of light, or at the edges and tips of larger pyramids are not taken into account in most traditional ray tracing.<sup>26,27</sup> Despite these shortcomings, modeled reflectance of textured substrates with ray tracing might still give insight into the expected behaviour of ideal pyramid textures.

In addition to the modeling work, the colour of all samples is calculated and expressed in both RGB coordinates and the chromaticity at the measured angles of incidence using the CIE 1931 2 degrees standard observer.<sup>28</sup>

### III. EXPERIMENTAL METHODS

As planar substrates, p-type (100) double side polished monocrystalline silicon wafers grown by the Czochralski method were used. The wafers have a thickness of 275  $\pm$ 20  $\mu m$  and a resistivity of 1-10  $\Omega \times cm$  and are cut into 30x30 mm pieces. To create substrates with a random upright pyramid texture on both sides of the wafer, the polished wafers were etched for 30 minutes in a KOH based solution. This solution was created by mixing 11.5 L of water with 517 mL of a 48% KOH solution, which was consecutively heated up to 80 °C. Once this temperature was reached, 180 mL of a proprietary buffering agent and 18 mL of a texturizer additive were added. A nitrogen bubble flow was present in the solution during the entire etching process to promote chemical mixing. After etching, wafers were consecutively rinsed in baths of deionized (DI) water, hydrochloric acid solution and DI water, and dried with a nitrogen gas flow. The textured samples shown in this work have a quarter circle shape with a radius of 30 mm. Scanning electron microscopy (SEM) measurements were carried out to ensure that no planar fractions were present on the surface. An example of this is shown in Figure SM1. To remove the native oxide layer on the samples before depositions, all samples were put in a 5% hydrogen fluoride (HF) bath for two minutes and subsequently rinsed with DI water and then dried with a nitrogen gas flow.

All thin films were deposited with an Oxford instruments plasmalab system 133 under a radio frequency (RF) power of 40 W, a reaction chamber pressure of 800 mTorr and a temperature of 350 °C. SiN<sub>x</sub> films were created using SiH<sub>4</sub> and NH<sub>3</sub> gas flows. A N<sub>2</sub> background flow of 980 sccm was used for all depositions. The other gas flows as well as the deposition time varied per sample, depending on the required refractive index and thin film thickness.

The thickness, as well as its accuracy, and the optical data of the thin films on planar substrates were determined by ellipsometry, using a Variable Angle Spectroscopic Ellipsometer (VASE) from J. A. Woollam. All samples were characterised from 300 to 1000 nm with 5 nm intervals using the same measurement parameters, which can be found in the SM.

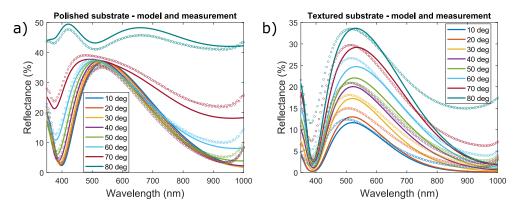


FIG. 1. Measured (symbols) and modeled (lines) angle dependent reflectance of sample G1 on polished silicon (a) and on textured silicon (b).

Ellipsometric data were fitted using the WVASE32 software from Woollam, using Cauchy or Tauc-Lorentz oscillators, depending on whether the thin films were light absorbant below 600 nm or not. The resulting optical data of the green sample are shown in Figure SM2.

The angle dependent reflectance was measured with an integrating sphere setup. A Spectral Products QTH 30 W ASB-W-030 high stability tungsten-halogen light in combination with an Oriel Cornerstone 260 monochromator from Newport was used as light source. To modulate the intensity of the light beam and improve the signal-to-noise ratio, the beam was sent through a chopper and lock-in amplifier. The beam was then focused by a series of lenses and depolarized by a depolarizer before entering a 6 inch RTC-060-SF integrating sphere from Labsphere, in which samples were placed inside via a clip-style center mount sample holder. Reflected light was detected with a S1336-5BQ silicon detector from Hamamatsu. All measurements were performed between 350 and 1000 nm, with 5 nm intervals. The reflectance was therefore modeled with the same wavelength domain and intervals. It was observed that from 950 nm, measured reflectance increased for all samples as the substrates became partially transparent. All samples were measured from the zenith angle  $\theta = 10$  to  $80^{\circ}$ with 10° intervals and results are shown in Figure 1 and 2 and discussed in the next section.

The thickness of the thin films on the textured substrates could not be determined with ellipsometry, as the reflected light missed the detector due to the surface texture. An estimation of the thickness was therefore made by measuring the reflectance of the samples at the mentioned angles and using Equation 1 to calculate the thickness. Although this equation is only valid for planar films, it has been demonstrated<sup>29</sup> that this is also a good approximation to determine thin film thickness on textured substrates, as the reflectance minima of textured and polished substrates with identical thin films are typically not far off. Transmission electron microscopy (TEM) measurements on other samples were carried and confirmed that this method determines the thickness of the thin film accurately within 5 nm. These measurements were carried out along multiple cross sections of the samples to confirm uniform thin film layer thickness.

From the measured reflectance spectra, the colours and chro-

maticities of the samples were calculated as shown in Figure 3 and discussed below. This calculation is based on the overlap of the reflectance spectra and the colour matching functions of the RGB and XYZ colour spaces, respectively. This allows the colour belonging to any reflectance spectrum to be expressed in the tristimulus values of the respective colour spaces, which are used to quantify colour and chromaticity.<sup>28,30</sup>

## IV. RESULTS AND DISCUSSION

As can be seen in Table I the thin film thickness of the samples is almost identical for the planar and textured substrates. Since the thin films are deposited under the same deposition parameters, it is assumed that the optical data are equal.

TABLE I. Thickness of thin film G1 on polished and textured substrate, determined by ellipsometry and angle dependent reflectance measurements, respectively.

Sample Name	Substrate	Film thickness (nm)
Green (G1)	Polished	$125.3 \pm 0.1$
	Textured	$123.0\pm5.0$

In Figure 1, one can see the measured and modeled angle dependent reflectance from both the polished and textured samples for incoming angles from  $\theta = 10^{\circ}$  to  $\theta = 80^{\circ}$ . Figure 1a) shows that the modeled and measured reflectance of the polished sample are generally in good agreement, with some exceptions for higher angles of incidence. The positions of the reflectance minima and maxima shift around 40 nm to shorter wavelengths when the incoming angle increases from 10 to  $60^{\circ}$ , and an additional 80 nm when the incoming angle increases from 60 to  $80^{\circ}$ .

The measured and modeled reflectance of the textured sample are shown in Figure 1b), and the positions of the reflectance minima and maxima are in reasonable agreement. However, there are differences in the intensities of the modeled and measured reflectance spectra, with the modeled reflectance generally being higher than the measured reflectance outside the reflectance maxima. These discrepancies can be caused by

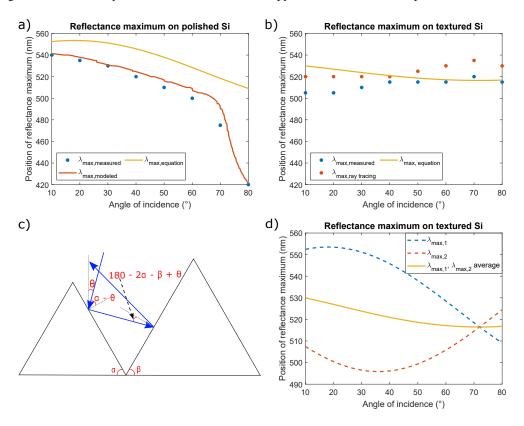


FIG. 2. (a) Position of reflectance maxima of G1 on polished silicon as measured (blue dots), modeled (red dots) and calculated (yellow solid line) using Equation 1, (b) Position of reflectance maxima of G1 on textured silicon as measured (blue dots), modeled (red solide line) and calculated (yellow solid line), (c) path of reflected light under incoming angle  $\theta$  on a textured substrate and (d) calculated reflectance maxima of G1 on textured silicon 3 (red dashed line), and the average of both equations (yellow solid line).

the shortcomings of the ray tracing model, which have been discussed before. Most importantly, one can see that in both cases the positions of the reflectance minima and maxima are stable under an increase in the angle of incidence, with a total shift of only about 15 nm under an increase from 10 to  $80^{\circ}$ .

One can also see that in general the reflectance of the textured sample is much lower than that of the polished one. As a result, identical thin films will have a darker colour appearance on textured substrates than on polished ones, which might be a disadvantage when trying to reach bright colours. Figure 1 also shows that the reflectance of the textured sample increases much more with an increase in the angle of incidence than that of the polished sample. This can be explained by the fact that the double bounce effect that usually decreases reflectance from textured samples is not that effective anymore for large angles of incidence.<sup>31,32</sup>

Figures 2a) and 2b) show the measured (blue dots), modeled (red solid line and dots) and calculated (yellow line) positions of the reflectance maxima with changing incoming angles for the planar and textured samples. The measured positions were extracted from the measurements shown in Figure 1. For the planar sample the peak positions were modeled with a precision of 0.1 nm at angles of incidence 10 to  $80^{\circ}$  at  $0.1^{\circ}$  intervals using the model based on the transfer matrix method, and for the textured sample the peaks were extracted from the mod-

eled data plotted in Figure 1b), which were determined by the OPAL model that was described above. Figures 2a) and 2b) show that the modeled and measured reflectance maxima are in good agreement for both substrates.

In Figure 2a), the positions of the reflectance maxima are also calculated using Equation 1. All three curves in Figure 2a) show the expected blueshift of the reflectance maxima. However, at high angles of incidence, the positions of the reflectance maxima as calculated by Equation 1 deviate from the modeled and measured positions. This is due to the fact that at these angles, internal reflectance becomes significant. As a result, the reflectance spectra largely consist of direct reflectance from the thin film, while the effects of interference decrease. This is taken into account by the optical model, which calculates the full reflectance of the sample, but not by Equation 1.

In Figure 2b), one can see that both the measured and the modeled reflectance maxima of the textured sample are much more stable against changes in the angle of incidence than those of the polished sample. This can be explained by considering the way in which light reflection depends on the angle of incidence on the textured substrate surface. Under normal incidence, about 68% of light incident on the textured substrate surface is reflected along the same path,<sup>15</sup> which is described in Figure 2c) for an angle of incidence  $\theta$ . By treating the two

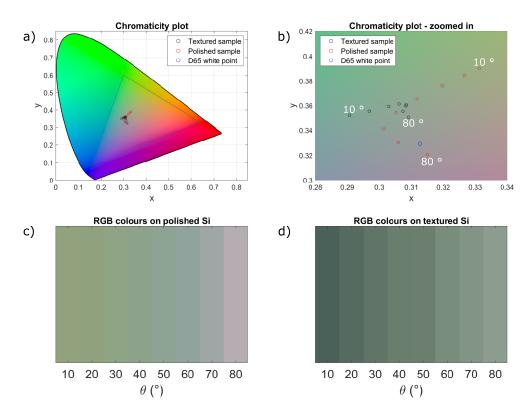


FIG. 3. Chromaticity diagram of the polished and textured samples at angles of incidence 10 to  $80^{\circ}$  (a), with additional zoom around the relevant area (b), together with colour patches of the polished (c) and textured (d) samples at angles of incidence 10 to  $80^{\circ}$ .

pyramids independently, we can describe the reflectance maximum that is caused by each pyramid using Equation 1:

$$\lambda_{max,1} = \frac{2n_1d}{m_1} \sqrt{1 - \left(\frac{n_0}{n_1}\right)^2 \sin^2(\alpha - \theta)}$$
(2)

$$\lambda_{max,2} = \frac{2n_1 d}{m_1} \sqrt{1 - \left(\frac{n_0}{n_1}\right)^2 \sin^2(180 - 2\alpha - \beta + \theta)}$$
(3)

Here,  $\lambda_{max,1}$  and  $\lambda_{max,2}$  stand for the reflectance maxima caused by pyramids 1 and 2, as indicated in Figure 2c). As both pyramids are coated with the same thin film,  $n_0$ ,  $n_1$ ,  $m_1$ and d are the same for both equations. Since  $\alpha$  and  $\beta$  are constant, we can see that under a change in the angle of incidence  $\theta$ , the two reflectance maxima shift in the opposite direction for most values of  $\alpha$  and  $\beta$ . Since we measure only one reflectance minimum and not two for these coated substrates, it is likely that the measured reflectance maximum is the average of the two maxima  $\lambda_{max,1}$  and  $\lambda_{max,2}$ . Under the assumptions that  $\alpha = \beta = 54.7^{\circ}$ , and  $n_1 = 2.25$  at all wavelengths, the positions of the two maxima  $\lambda_{max,1}$  (blue dashed line) and  $\lambda_{max,2}$  (red dashed line) and their average (yellow line) at different angles of incidence are shown in Figure 2d). In Figure SM2a), one can see that at wavelengths 500 to 550 nm, which is where the reflectance maxima occur at most angles of incidence, n = 2.23 - 2.26. The assumption that n =2.25 at all wavelengths thus does not impact the position of the calculated reflectance maximum much.

The average of  $\lambda_{max,1}$  and  $\lambda_{max,2}$  is also plotted with the measured reflectance maxima of the textured films in Figure 2b) (yellow line). Indeed, Figure 2b) shows that the average of  $\lambda_{max,1}$  and  $\lambda_{max,2}$  is in fairly good agreement with the measured reflectance maxima of the textured sample.

The effect of the (in)stability of the reflectance extrema can clearly be seen when looking at the colours of the samples at different angles of incidence, as shown in Figure 3. Figure 3c) demonstrates that the colours of the polished sample become bluer as the angle of incidence increases. On the other hand, the colour of the textured sample, which is as expected much darker than that of the polished sample, remains green under a change in the angle of incidence, as shown in Figure 3d). The colour of this sample does become brighter with an increase in the angle of incidence as the magnitude of the reflectance increases, with Y increasing from 0.10 at 10° to 0.32 at 80°. This is caused by the fact that at higher angles of incidence, more light will take paths which are only incident on the textured substrate surface once instead of twice or more.

The difference in colour stability between the respective samples is especially apparent when one looks at their chromaticities, which are shown in Figures 3a) and 3b). While the chromaticity of the textured sample remains relatively stable, that of the polished sample changes to blue chromaticities more drastically, before changing to a more purple region at high angles of incidence, as the reflectance spectra completely change due to a decrease in the effect of interference. In the SM, the colours and chromaticities of the three samples with different colours are shown. There, too, samples on textured substrates show a stronger colour stability than those on polished ones.

This is also seen when looking at photographs of the samples taken under varying angles of incidence, as shown in Figure 4. All photographs are taken under office lighting, which influences the observed colour of the samples. This is most clearly visible in figures 4a) and 4b): the light patch on the samples is caused by direct illumination of the office lamps, while the darker regions show the colour that best agrees with what is observed with the naked eye. Furthermore, the photographs show the advantage of creating matte colours with the textured samples over shiny colours on the polished ones. This is beneficial for BIPV applications as glare effects are avoided. To image the observed colour of the samples as accurately as possible, different camera exposure times were used, which affect the colour of the background. The photographs clearly demonstrate the colour change of the samples. Figures 4a) and 4b) show that the colour of the polished sample changes from a yellow-green colour at 15 degrees to a green-blue colour at 60 degrees. Figures 4c) and 4d) show that the textured sample, on the other hand, remains green at both angles of incidence, but with different brightness. The samples show slight colour inhomogeneities around the edges, which are caused by thin film thickness variations from the PECVD depositions.

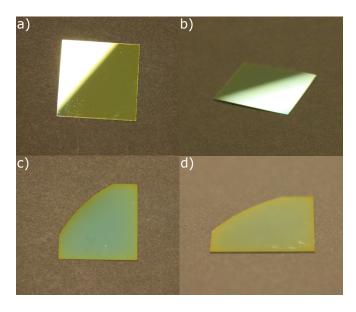


FIG. 4. Photographs of the polished (a and b) and textured (c and d) sample, taken under office lighting at 15 (a and c) and 60 (b and d) degrees. The same black paper is used as background for all samples, but seems to have a different colour in some photographs due to variations in the camera exposure time. The polished substrates are 30x30 mm, and the textured ones have a radius of 30 mm.

# V. CONCLUSION

In this work, it has been shown that dielectric thin filmcoated silicon substrates with a pyramidal surface texture have a much stronger angular colour stability than planar substrates with identical coatings. This has been shown by angle resolved reflectance measurements of fabricated planar and textured coloured c-Si substrates. It has been found that the colour stability can be explained by the primarily averaged double interference on the pyramides of the textured substrates. This is a major benefit for building integrated photovoltaic applications, as stable colours are often desired for aesthetic integration into the built environment. Considering that the textured substrates in this work are widely used in commercial c-Si PV, this thin film colouring technique on textured substrates might be ready to be applied on a large scale.

### SUPPLEMENTARY MATERIAL

See supplementary material for an SEM image of the textured substrate surface texture, as well as the ellipsometric measurement parameters, the refractive index of the sample and thin film thickness and colour plots of three additional samples.

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# DATA AVAILIBILITY

The data that support the findings of this study are available from the corresponding author upon reasonable request.

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