



OPEN

Enhanced photovoltaics inspired by the fovea centralis

Gil Shalev^{1*}, Sebastian W. Schmitt^{1*}, Heidemarie Embrechts¹, Gerald Brönstrup^{1,2} & Silke Christiansen^{1,2}¹Max Planck Institute for the Science of Light, Günther-Scharowsky-Straße 1, 91058 Erlangen, Germany, ²Helmholtz-Zentrum Berlin für Materialien und Energie, Kekulestrasse 5, 12489 Berlin, Germany.

The fovea centralis is a closely-packed vertical array of inverted-cone photoreceptor cells located in the retina that is responsible for high acuity binocular vision. The cones are operational in well-lit environments and are responsible for trapping the impinging illumination. We present the vertical light-funnel silicon array as a light-trapping technique for photovoltaic applications that is bio-inspired by the properties of the fovea centralis. We use opto-electronic simulations to evaluate the performance of light-funnel solar cell arrays. Light-funnel arrays present ~65% absorption enhancement compared to a silicon film of identical thickness and exhibit power conversion efficiencies that are 60% higher than those of optimized nanowire arrays of the same thickness although nanowire arrays consist of more than 2.3 times the amount of silicon. We demonstrate the superior absorption of the light-funnel arrays as compared with recent advancements in the field. Fabrication of silicon light-funnel arrays using low-cost processing techniques is demonstrated.

In order for silicon-based solar energy to become a primary energy source it is essential to render it cost effective as compared with current carbon-based technologies¹. The high costs stem from both the high grade silicon needed in order to provide the required electrical performance and sufficient bulk material to maximize light absorption to a level acceptable for sufficiently high efficiencies (~20% are required to compete with existing bulk and thin film technologies)². For cost efficiency in novel crystalline silicon thin film technologies it is therefore mandatory to at least reduce the amount of the silicon used while maintaining power conversion efficiencies (PCE) of current wafer based technologies. Recently, crystalline silicon thin film solar cell technologies reached PCE of 11.5% and promising open circuit voltages (V_{oc}) of 650 mV. However the cell design still suffers from insufficient trapping of visible and near infrared light³.

Nanostructuring the surface of a solar cells allows for a more efficient harvesting of solar radiation and also the collection of infrared radiation well below the absorption edge⁴. Silicon photovoltaics based on arrays of vertically-aligned cylindrical silicon nanowires (NWs) of several hundred nanometers in diameter with either radial or axial junction were demonstrated as potential candidates to address this challenge. These arrays exhibit appreciated absorption due to enhanced light trapping and support efficient carrier collection over short distances, and hence permit reduced thickness and the utilization of electrically low grade silicon (=low cost) with short minority carrier diffusion lengths (or lifetimes)^{5–8}. Various arrays of silicon NWs were produced and demonstrated to potentially provide the required light absorption enhancement in thin layers, support tolerating short minority diffusion lengths, and are produced by means of low cost fabrication techniques^{9–12}.

Another promising family of structures that are currently under examination for photovoltaics are either periodic or randomized vertical cone arrays. Cone arrays are reminiscent of the anti-reflective surfaces found on the transparent wings of hawkmoths and on corneas of moth and butterfly eyes, for example^{13,14}. These surfaces consist of hexagonal arrays with periodicity of approximately 240 nm that are composed of subwavelength structures with rounded tips. Effectively, the subwavelength structures introduce a gradual refractive index profile and form a favorable optical impedance matching between the organic tissues and the surrounding air. Huang et al demonstrated silicon nanotip arrays mimicking the ‘moth eye’ array and reported sub 1% reflection with a tip height of 16 μm ¹⁵. Recently, Jeong et al reported a record thin silicon solar cell efficiency of 13.7% attributed to enhanced absorption due to surface nanocone arrays, where an absorption of >95% for the wavelength range of 400–800 nm was shown for a 10 μm thick substrate¹⁶. Spinelli et al demonstrated a similar concept with arrays composed of nanocylinders residing on a silicon substrate for which the high levels of absorption are attributed to forward scattering by the cylinders into the substrate¹⁷.

In this paper, we introduce the light-funnel (LF) array which is a new light trapping scheme for photovoltaics that mimics the structure and arrangement of the cone photoreceptors in the fovea centralis region of the retina. The human retina, for example, has approximately 6 million cone photoreceptors and 120 million rod photo-

SUBJECT AREAS:

SOLAR ENERGY AND
PHOTOVOLTAIC
TECHNOLOGY

ELECTRONIC DEVICES

NANOPHOTONICS AND
PLASMONICSELECTRICAL AND ELECTRONIC
ENGINEERING

Received

3 November 2014

Accepted

23 January 2015

Published

24 February 2015

Correspondence and
requests for materials
should be addressed to
G.S. (gshalev@gmail.
com)

* These authors
contributed equally to
this work.



receptors¹⁸. Their optical properties are determined by their size, morphological structure, as well as the media constituting and surrounding them^{19,20}. Various papers described the wave guiding properties of these photoreceptors specifically with respect to mode formation^{21–23}. Currently, open questions include micro anatomical and molecular differences between rods and cones²³. Cones are operational in relatively bright light and are responsible for high acuity binocular vision, whereas rods photoreceptors trade acuity with sensitivity and are triggered in dim conditions. In that sense the bright light conditions required for the functionality of cones is reminiscent of the bright-light conditions required for the functionality of a solar cell while rods act more as sensitive light detectors. The fovea centralis is a distinct pit in the central retina containing only cones in a densely-packed mosaic that is recognized as the region responsible for high acuity binocular vision²⁴. Moreover, the packing density of the foveal cones accurately correlates with the human visual acuity^{18,25}. The geometrical difference between the retinal cones and the ‘moth eye’ cones is acute: the retinal cones are inverted with the large base facing the incoming illumination and can reach several microns in diameter and dozens of microns in length^{19,26} in contrast with the much smaller upright ‘moth eye’ cones.

We use three-dimensional (3D) Technology Computer Aided Design (TCAD) optical and electrical simulations to model periodic square-tiled arrays composed of LFs in the shape of inverted frustum cones. We demonstrate their superior absorption relative to both continuous thin films and NW arrays of identical thickness across a wide range of angles of incidence. Also, we present extensive electrical simulations showing that the overall PCE of the LF solar cell is 60% higher than the PCE of an optimized NW-based solar cell of the same thickness although the NW solar cell consists of more than 2.3 the amount of silicon. Furthermore, we demonstrate the superiority of the LF array against recent advancements in the field. Finally, fabrication of silicon LF arrays using low-cost fabrication techniques is demonstrated showing the technological feasibility of the proposed concept.

Results and Discussion

3D finite-difference time-domain (FDTD) optical simulations.

Figure 1a presents an illustration of a 3D vertical LF square-tiled array where the color coded individual LFs reflect the absorbed photon density. Figures 1b,c show scanning electron microscopy images of LF arrays that were fabricated using low-cost fabrication techniques²⁷ (see methods and Supplementary 1). Figure 2a presents 3D finite-difference time-domain (FDTD) optical simulations of the relative absorption spectra under normal incidence (Fig. 1) of two representative LF arrays with a period of 500 nm where the top diameter (D_t) is held constant at 400 nm and the bottom diameter (D_b) is fixed at 100 nm and 300 nm. The 500 nm period was selected as it couples best to the solar spectrum that peaks around wavelength of ~ 500 nm¹⁰ and therefore in the current work the considered array periods of both LF arrays and NW arrays are always set to 500 nm. For reference, the spectra of silicon thin film and of a NW array with NW diameter (D) of 400 nm, are shown as well. The juxtaposition of the LF array with a NW array not only serves for benchmarking but also allows a more acute understanding of the governing physics of the LF array. Unless otherwise specified the height (h) of all arrays and thin films is set to 2 μm . Thin film Fabry-Perot oscillations are not evident as the boundary condition at the bottom edge of the LF and the NW arrays and the thin film is set to perfectly matching layers (see methods). Note, that LF arrays exhibit the highest absorption. Furthermore, the LF arrays exhibit a broadband absorption similar to the NW array^{12,27–30} only superimposed with additional strong resonances at specific wavelengths. The ultimate absorption efficiency is the averaged and weighted (Air Mass 1.5) relative absorption, and it is assumed that each above bandgap photon generates an electron-hole pair that is collected at the

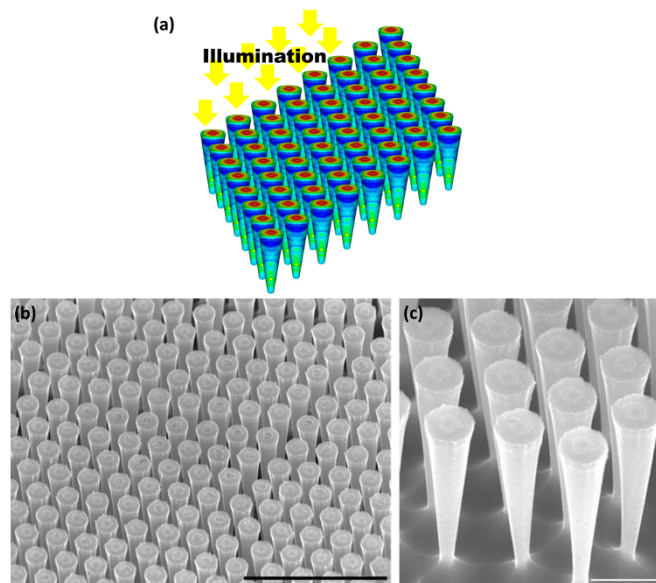


Figure 1 | The light-funnel array. (a), 3D representation of a free-standing infinite square-tiled LF array under normal incidence where the color coded individual LFs represent the absorbed photon density. (b), A scanning electron microscope (SEM) image of a silicon LF array fabricated using low-cost fabrication techniques combining nano-sphere lithography and Langmuir-Blodgett deposition of 1 μm polystyrene spheres. Scale bar is 5 μm . (c), A patterned silicon wafer was cracked in order to image LFs from the side in order to better present their inverted frustum cone structure. Scale bar is 1 μm . See methods section for LF array fabrication.

electrodes. Figure 2b presents the ultimate absorption efficiency increase relative to a continuous silicon film for a LF array with D_t of 400 nm vs. D_b . The minimum considered D_b is 100 nm, and the maximum considered D_b is 400 nm which reflects the convergence of the LF into a vertical NW. Note that LF arrays exhibit an absorption enhancement of $\sim 65\%$ (for $D_t = 400$ nm and $D_b = 140$ nm) relative to the continuous film in comparison with NW arrays that present an absorption enhancement of 36.6%. Furthermore, the absorption of the LF array is superior despite the smaller filling ratio (defined as the volume of the LF divided by the volume of a unit cell) where 22% is calculated for the LF array with $D_t = 400$ nm and $D_b = 100$ nm and 50% for the NW array. The absorption enhancement of the LF array increases with the increase in cone angle (α) from 0° to 8° , which entails a decrease in filling ratio.

The relative absorption is a figure of merit describing the absorption performance of an array, while the absorption efficiency factor (Q_{abs}) is a figure of merit describing the ability of a single particle (e.g. LF or NW) to concentrate or to couple light into itself. Q_{abs} is the absorption cross-section (C_{abs} - the ratio between the total absorbed photons and the incident photon flux) normalized by the LF top area³¹ (for normal incidence). Here Q_{abs} is given by: $Q_{abs} = C_{abs} / (\pi/4 D_t^2)$. The relative absorption and the Q_{abs} of a LF or a NW nested inside an array relate to one another by the filling ratio. Figure 2b presents also the average of Q_{abs} (averaged over the wavelength range of 400 nm–1100 nm) under normal incidence of a single LF in the array as a function of D_b . Figure 2b shows that Q_{abs} is directly correlated with the ultimate absorption efficiency and increases with a decrease in D_b . Note that Q_{abs} increase is due to increase in C_{abs} as the area of the LF top base (i.e. $\pi/4 D_t^2$) that is used to scale C_{abs} is not changed for all data points in Fig. 2b. The importance of this behavior becomes evident when considering the relocation of an isolated NW into a NW array; a single isolated subwavelength NW presents a relatively high Q_{abs} ^{32–37}, however Q_{abs} decreases substantially once

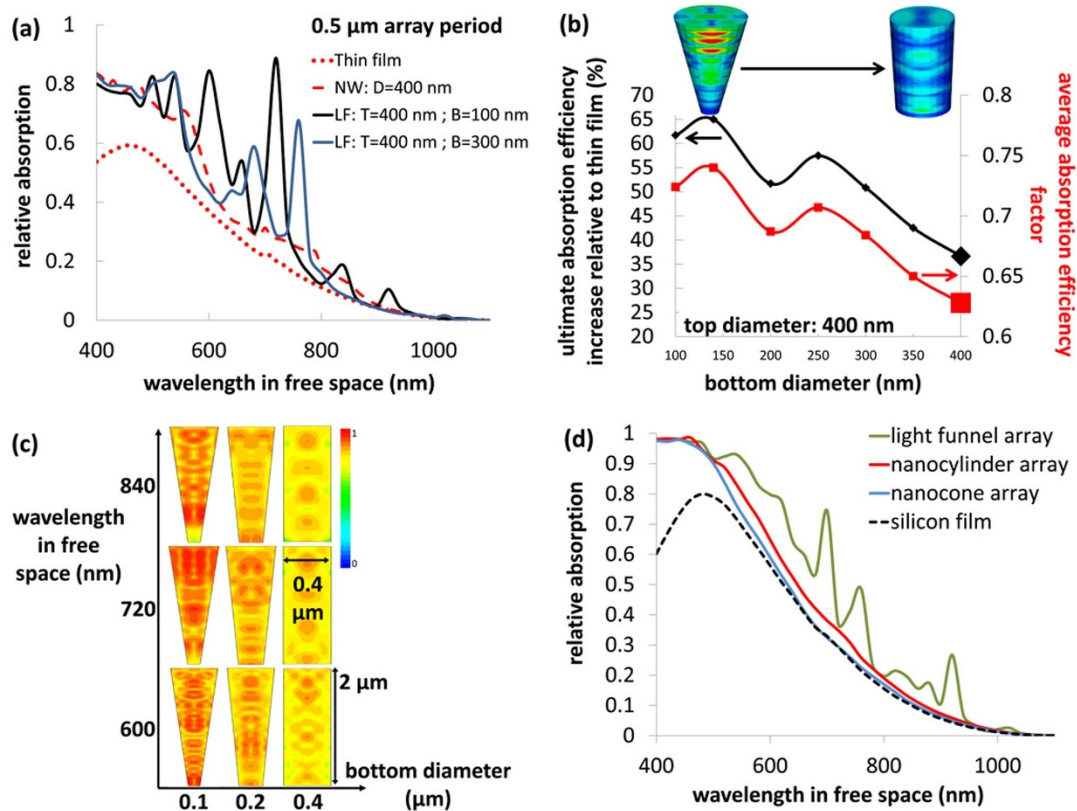


Figure 2 | Optical absorption of a light-funnel array. (a), Relative absorption of LF arrays, a NW array and a continuous silicon thin film. The height is 2 μm. Note the absorption enhancement of the LF arrays. **Legend:** T: top diameter, B: bottom diameter. (b), Increase in ultimate absorption efficiency relative to a continuous silicon film and the absorption efficiency factor of a single LF (Q_{abs}) in the array as a function of LF bottom diameter for a constant top diameter of 400 nm. Both the LF array absorption and the absorption efficiency factor of a single LF increase with decrease in bottom diameter despite the decrease in filling ratio. The period of the arrays is 500 nm and the heights are 2 μm. The large markers at bottom diameter = 400 nm reflect the values of a NW with the corresponding diameter. (c), The absorbed photon density of NWs and LFs at wavelengths 600 nm, 720 nm, 840 nm and for LFs bottom diameter of 100 nm, 200 nm, 400 nm. Note the increase in the absorbed photon density once deforming the 400 nm NW into a LF. (d), **LF array:** square-tiled array with a 520 nm period, $D_b = 100$ nm, $D_t = 400$ nm, 2 μ height and 50 nm Si_3N_4 anti-reflection coating. The geometries of the nanocone and the nanocylinder arrays are taken from the respective publications. **Nanocone array:** square-tiled array with a period of 500 nm, nanocone bottom diameter of 450 nm, nanocone height of 400 nm and 80 nm coating of SiO_2 . **Nanocylinder array:** square-tiled array with period of 450 nm, nanocylinder width of 250 nm, nanocylinder height of 150 nm, and 50 nm Si_3N_4 anti-reflection coating. **Thin silicon film:** 2 μ thickness with 80 nm standard anti-reflection coating. The thicknesses of the underlying substrates are selected to ensure that all three technologies share the same amount of silicon. Hence the thicknesses of the substrates are 1.12, 1.56 and 2 μm for the LF array, nanocone array, and the nanocylinder array, respectively.

the isolated NW is introduced into an array as adjacent NWs are now optically-coupled (Supplementary S2). This loss in Q_{abs} of a single array-nested NW can be compensated by transforming the NW into a LF, decreasing the bottom edge of it and hence increasing Q_{abs} and the overall array absorption as demonstrated in Fig. 2b where the Q_{abs} of the 400 nm NW is increased from 0.63 to 0.72 for a light-funnel with $D_b = 100$. Again, an increase in Q_{abs} is not due to scaling of C_{abs} but is due to increase in C_{abs} itself which concludes an increase in the absorption of a single LF. Still, this ~15% increase of Q_{abs} is not sufficient to account for the ~50% absorption enhancement of the light-funnel array with $D_b = 100$ nm as compared with the 400 nm NW array. Hence, the absorption enhancement in LF array is a superposition of various mechanisms; namely, light trapping associated with the array and possible scattering capabilities of a single LF^{6,38}. The contribution of each mechanism is yet to be unveiled. Interestingly, the superior absorption of a single LF as compared with single NW can also be described when considering solely geometrical optics as was shown by Quinn³⁹ who demonstrated the direct correlation between absorption and cone angle (α) with $A = I - T^n$ where A is the relative absorption, T is the relative reflectance of the cone wall and n is the number of the reflections a ray undergoes before it is reflected out of the cone. For incoming rays parallel to the

axis of the cone $n = \text{Int}(\pi/2\alpha)$ where $\text{Int}(\pi/2\alpha)$ is the greatest integer less or equal to $(\pi/2\alpha)$.

Figure 2c presents the simulated normalized absorbed photon density at wavelengths 600 nm, 720 nm and 840 nm along the vertical axis of a LF for a constant $D_t = 400$ nm and $D_b = 100$ nm and 200 nm and of a 400 nm NW for reference (all structures are nested in an infinite square-tiled arrays). Note that the NW exhibits low order modes, whereas the LF exhibit complex 3D modes. One can establish an intuitive understanding of this acute difference between the two types of structures by hypothesizing the swift transition from a NW geometry to the geometry of a LF; at a specific wavelength the hypothetical deformation of the NW into a LF modifies the constellation of internal reflections inside the cavity in such a way that renders the formation of complex 3D modes. Now, the presence of multitude complex 3D modes provides the strong coupling of light to the LF which is manifested in distinct absorption peaks.

Recently, two outstanding advancements in the field were published: the nanocylinder array^{17,40} and the nanocone array¹⁶ (reviewed above). We conduct comparative absorption simulations of the LF array against these two recent advancements in order to assess the contribution of the LF array to the state of the art. The dimensions and geometry of both the nanocylinder array and the



nanocone array are taken directly from the publications (see methods). The simulated LF array is of the following geometry: square-tiled array with a 520 nm period, $D_b = 100$ nm, $D_t = 400$ nm, 2 μ height and 50 nm Si_3N_4 anti-reflection coating. Also, for reference a 2 μ thin silicon film with 80 nm standard anti-reflection coating is simulated. The aforementioned nanocylinder and the nanocones are relatively small features that require the presence of a substrate in order to maximize their performance and therefore the arrays are simulated on top of underlying substrates. The substrate thickness of the nanocylinder array is 2 μ to match the thickness of the 2 μ thin film. The substrates of the nanocone and the LF arrays are selected to ensure that all three technologies share the same amount of silicon, i.e. the height of the underlying substrates are 1.56 μ m and 1.12 μ m for the nanocone and the LF arrays, respectively. In order to crystalize the performance of the different arrays we applied perfectly matching layers boundary condition at the bottom of the substrates in order to exclude the contribution of the coupling of Fabry-Perot resonances with the waveguide resonances to the overall absorption. Henceforth, a reference to any array refers to the array together with the underlying substrate. Figure 2d compares the simulated relative absorption spectra of the LF array, the nanocylinder array, the nanocone array and the thin silicon film. First note that the thin film absorption is considerably higher than the absorption of the thin film in Fig. 2a due to the presence of the 80 nm Si_3N_4 anti-reflective coating. Also note the increase in absorption once the thin film is decorated with nanocylinder array (increase in ultimate absorption efficiency from 16% to 20%). The nanocone array exhibits a lower absorption as compared with nanocylinder array (ultimate absorption efficiency of 19%). However, the LF array presents a significantly superior broadband absorption with an ultimate absorption efficiency of 24.3% which reflects 52% improvement as compared with the thin film (this relative improvement differs from that in Fig. 2a due to the higher thin-film absorption on account of the 80 nm Si_3N_4 anti-reflective coating).

Photovoltaics require absorption enhancement over a wide range of angles of incidence (AOI). Figure 3a, b presents the AOI dependence of the relative absorption at wavelengths of 540 nm and 600 nm for a thin silicon film, a thin silicon film with a standard 50 nm Si_3N_4 anti-reflecting coating and for a LF array with $D_t = 400$ nm, $D_b = 100$ nm also with a standard 50 nm Si_3N_4 anti-reflecting coating (shown is the average of both polarizations). These two representative wavelengths are chosen as these are located close to the peak power of the solar spectrum. Note that the LF array presents an enhanced AOI dependency in both 540 nm and 600 nm wavelengths for the AOI range of 0–80 deg. The LF array maintains a relative absorption of $\sim 85\%$ whereas the coated thin film exhibits an absorption of $\sim 70\%$ for wavelength 540 nm in the AOI range of 0–50°. For

AOI above 50° the absorption decreases for the LF array but is still substantially greater than the coated continuous silicon film.

Electrical modeling. 3D TCAD simulations are also utilized in order to model the electrical response of the LF solar cell device. In the current investigation we also adapt the methodology employed in the optical section and perform an electrical comparison between the LF array and the NW array in order to better understand and describe the electrical behavior of the LF array. For this purpose the LF array of $D_t = 400$, $D_b = 100$ nm is selected and compared with a NW array with $D = 400$ nm. The calculations pertain to the opto-electronic performance of a single LF and a single NW nested in their respective arrays as the above optical modeling is performed for an infinite periodic square-tiled arrays. The main concern relating to the electrical performance of the LF is surface effects as the surface-to-volume ratio (S/V) of the LF is greater (by 75% for the selected LF and NW) and hence expected to be more sensitive to surface recombination velocity (SRV). In order to address specifically this matter we selected an axial junction configuration as it is more susceptible to surface effects than a radial structure^{9,12,41} (see methods). Figure 4a presents the solution of the Poisson equation (electrostatic potential distribution) and the distribution of Shockley-Read-Hall recombination (SRH) for both structures. The white and pink lines mark the depletion regions and the metallurgical junction, respectively, where the metallurgical junction is defined as the interface between the n and p region or, in other words, the interface where the concentration of the acceptors and donors is equal. Note the high recombination at the degenerated emitter and the low recombination at the depletion areas.

Figure 4b presents the open circuit voltage (V_{oc}) and short circuit current (I_{sc}) of a NW and a LF for a range of base doping levels and the respective SRH bulk lifetimes as the doping-dependent SRH recombination implies that bulk SRH minority carrier lifetimes (τ_{SRH}) are coupled to the doping level^{42–45}, and Fig. 4c shows the current vs. voltage (I–V) curves for base doping level of 10^{18} cm^{-3} . For both devices I_{sc} decreases with increase in base doping due to increase in SRH recombination (see Supplementary S3). Also in both cases V_{oc} exhibits a maximum where it increases for base doping of 10^{16} cm^{-3} – 10^{18} cm^{-3} despite the decrease in I_{sc} ; the increase in base doping entails a decrease in saturation current and increase in V_{oc} while the simultaneously decrease in τ_{SRH} concludes an increase in saturation current and a decrease in V_{oc} ⁴⁶. Note, that for base doping level of 10^{19} cm^{-3} the effect of recombination overcomes the gain due to an increase in doping and an overall decrease in both I_{sc} and V_{oc} is observed. The power $I_{sc} * V_{oc}$ is maximized for base doping of 10^{18} cm^{-3} for both structures. At this base doping level the LF exhibits an increase of $\sim 55\%$ in I_{sc} and $\sim 3\%$ in V_{oc} relative to the NW device, where the relative superiority of the LF I_{sc} over the NW stems

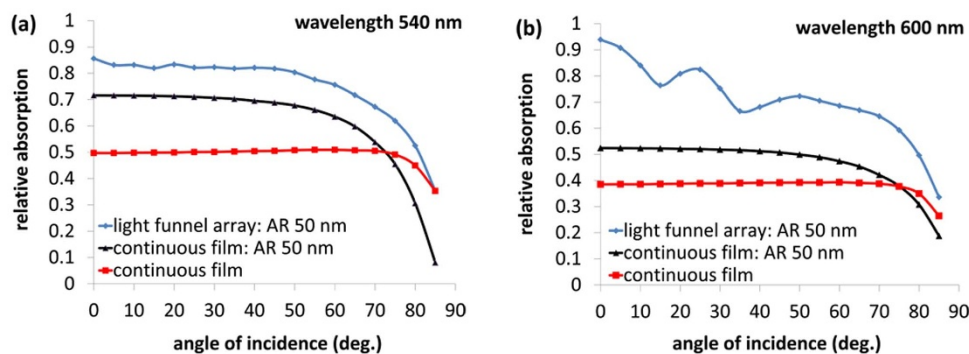


Figure 3 | The dependency of the relative absorption on the angle of incidence for a continuous silicon film, a continuous silicon film with 50 nm Si_3N_4 antireflection coating and for a LF array with a period of 500 nm, $D_t = 400$ nm and $D_b = 100$ nm and with a 50 nm Si_3N_4 antireflection coating (all heights are 2 μ m). These wavelengths are chosen as these are located at the vicinity of the peak power of the solar spectrum. (a), wavelength 540 nm. (b), wavelength 600 nm.

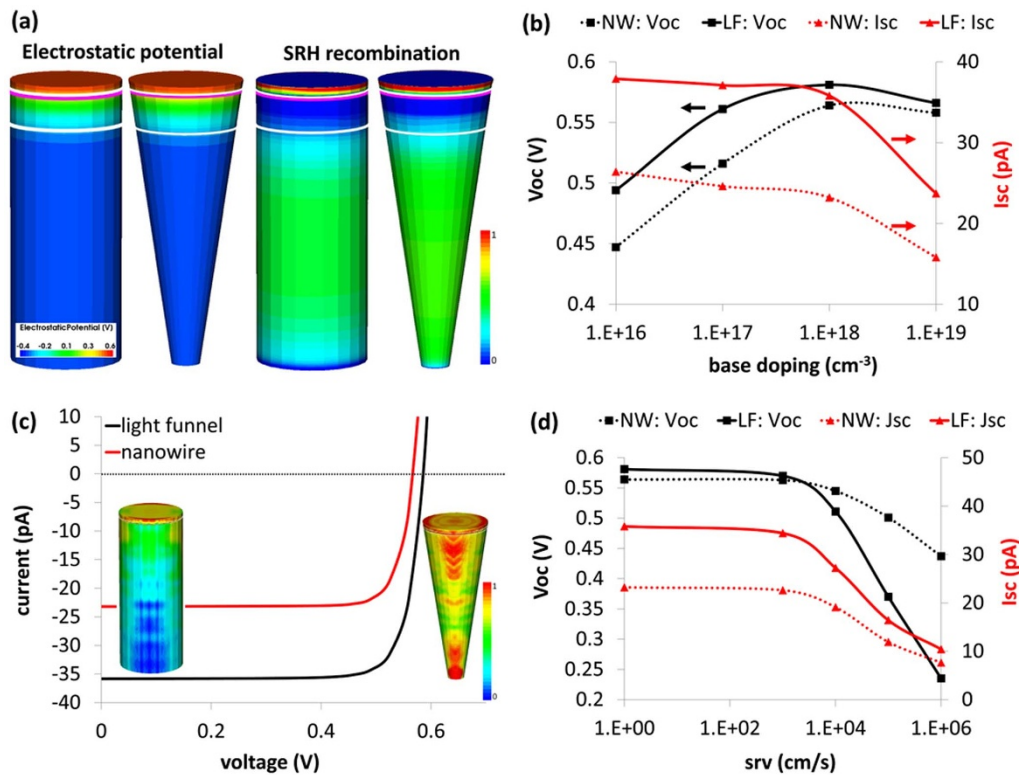


Figure 4 | 3D electrical simulations comparing the performance of a single LF with a single NW in their respective arrays under Air Mass 1.5 illumination. (a), Left. The electrostatic potentials inside a NW and a LF for 10^{20} cm^{-3} emitter (phosphorus) and 10^{16} cm^{-3} p-type base. Right. Normalized SRH recombination. (b), V_{oc} and I_{sc} vs. base doping for NW and LF solar cells. The main advantage of LF over NW is the increased I_{sc} which is a consequence of the enhanced LF absorption. (c), Current vs. voltage of NW and LF for base doping of 10^{18} cm^{-3} . The 3D NW and LF images presents the superior carrier generation in LF. (d), V_{oc} and I_{sc} of LF and NW vs. surface recombination velocity. Both structures are relatively insensitive to SRV in the range of 0– 10^3 cm/s . However for $\text{SRV} > 10^3 \text{ cm/s}$ a sharper decrease in LF performance is recorded. This decrease is attributed to S/V effects.

from the superior light absorption. Finally, at base doping of 10^{18} cm^{-3} the LF presents an overall increase in $I_{sc} * V_{oc}$ of 60% relative to the NW which reflects 60% increase in the PCE of the LF solar cell device over the NW-based device.

Figure 4d presents the dependencies of V_{oc} and I_{sc} for both devices on SRV (base doping of 10^{18} cm^{-3}). V_{oc} and I_{sc} are insensitive to surface for both NW and LF in the range of $\text{SRV} = 0-1 * 10^3 \text{ cm/s}$. However, for $\text{SRV} > 10^4 \text{ cm/s}$ a substantial decrease in LF V_{oc} is shown. This is expected as the S/V ratio of the LF is higher than the S/V ratio of the nanowire by 75%. This is specifically evident when considering the model formulated by Allen et al.⁴⁷ for carrier lifetimes in NWs. Here, we easily translate this model to an oversimplified but insightful description of the LF where we slice the LF along the vertical axis and the following describes the relationship between the various lifetimes in each slice: $1/\tau_{eff} = 1/\tau_b + 4 * \text{SRV}/d_{LF}(z)$ where τ_{eff} is the effective carrier lifetime, τ_b is the bulk lifetime and $d_{LF}(z)$ is the diameter of the LF at a certain location along its vertical z-axis. As d_{LF} decreases toward the lower base of the LF so is τ_{eff} which reflects the fact that at the bottom of the LF the surface plays a more pronounced role and hence the overall τ_{eff} of the LF is smaller than that of the NW for $\text{SRV} > 10^4 \text{ cm/s}$. Specifically, τ_{eff} of the NW and at the very top of the LF is 4 times greater than τ_{eff} at the bottom of the LF (10^{-9} sec and $2.5 * 10^{-10} \text{ sec}$ at the top and bottom of the LF, respectively). Finally, one should keep in mind that in order to avoid detrimental surface effects one could utilize radial junction configurations or apply appropriate surface passivation in order to suppress SRV as, for example, was already demonstrated with Al_2O_3 passivation where $\text{SRV} < 10 \text{ cm/s}$ was measured⁴⁸.

As was performed in the above optical section, we compare the electrical performance of the LF array with recent advancements in the field, namely the nanowire and the nanocylinder arrays.

Provided the LF array, the nanowire array and the nanocylinder array are transformed into solar cells with identical electrical configurations (i.e. same type of junction, same doping...) than all three cells will exhibit identical V_{oc} and fill-factor. Therefore the increase in ultimate absorption efficiency is directly translated into an increase in PCE. This implies almost 30% PCE increase of the LF solar cell as compared with nanowire array that holds the record PCE for thin film solar cell. Remarkably, the LF array is superior to the nanocylinder array although the enhanced performance of the nanocylinder array as compared with a thin film is most significant when coupled to a thin substrate such as demonstrated here⁴⁹. S/V ratio is another paramount parameter that needs consideration. For the optimized nanowire and nanocylinder geometries used in the optical section the S/V ratio of the nanowire and the LF are identical (15%) while the S/V ratio of the nanocylinder is higher (22%) (see methods), and therefore we expect the LF array to be less susceptible to SRV. Also, both the nanowire and the nanocylinder geometries cannot support the formation of a diffused homo-junction inside the nanostructures as these are too small, whereas the larger LF could also potentially accommodate this electrical configuration. Lastly, further improvements in the performance of the LF array could take place as the LF geometry used here is not fully optimized.

Conclusions

We present here a new light trapping technique for photovoltaics based on LF arrays that mimics the structure and arrangement of the foveal cones. The absorption of the LF array is demonstrated for an array period of $0.5 \mu\text{m}$ where absorption enhancement of $\sim 65\%$ in comparison with a thin continuous silicon film of the same thickness is shown. The increase in absorption, as compared to a corresponding NW array, is attributed partly to superior light trapping coupled



with an increase in the absorption efficiency factor of a single LF. Also, it is shown that the PCE of a single LF solar cell nested in an array is ~60% higher than the efficiency of the corresponding NW solar cell. The PCE increase is attributed to the increase in absorption with the consequential increase in I_{sc} although the NW solar cell consists of more than 2.3 times the amount of silicon. We also demonstrate the superior absorption of the LF arrays as compared with recent advancements in the field. Finally, the fabrication of silicon LF arrays is exemplified using low-cost processing techniques.

Methods

Fabrication of light-funnel arrays. Silicon wafers were patterned using nano-sphere lithography. This process entails the formation of self-assembled monolayers of polystyrene nanospheres (PSS) on top of the wafers using the Langmuir-Blodgett technique. In the present case PSS of 1 μm in diameter were used. The resulting monolayers exhibit densely-packed hexagonal tiling that spans areas of several square centimeters. After deposition the size of the PSS was reduced with O_2 plasma using Oxford Instruments Plasmalab 100 inductively coupled reactive ion etching. The silicon etch was performed with Oxford Instruments Plasmalab 100 inductively coupled plasma reactive ion etching system run in a cryogenic mode using a gas mixture of SF_6 , C_4F_8 and O_2 . Finally, the PSS were removed in an ultrasonic bath. See Supplementary S1 for more details.

Numerical calculations. The optical and the electrical simulations were performed with Synopsys TCAD Sentaurus, Mountain View, CA, USA.

Finite-difference time-domain (FDTD) electromagnetic simulations. The relative absorption, absorption efficiency factor and the field distributions were calculated using a three-dimensional finite-difference time-domain simulations. The simulation box size was set to the size of the unit cell with periodic boundary condition along the lateral dimensions and perfectly matching layers along the vertical axis (A perfectly matching layer is an artificial absorbing layer for wave equations that is used especially in FDTD to truncate computational regions in open boundaries problems). For each run the absorption, reflection and transmission were calculated using sensors that were located above the device (reflection) and below the device (transmission). In addition, for each simulation run the absorbed photon density and the power flux density were calculated at each mesh point. The calculations were performed for a plane wave excitation and a spectrum range of 400–1100 nm in 10 nm steps. The material optical constants were taken from the literature⁵⁰. The details of the nanocone and the nanocylinder arrays were taken directly from the publications^{16,17}. The nanocylinder array: square-tiled array with period of 450 nm, nanocylinder width of 250 nm, nanocylinder height of 150 nm, and 50 nm Si_3N_4 anti-reflection coating. The nanocone array was simulated with the following: square-tiled array with a period of 500 nm, nanocone bottom diameter of 450 nm, nanocone height of 400 nm and 80 nm coating of SiO_2 (note that Jeong et al measured hexagonal array¹⁶).

Electrical simulations. The Poisson and the Continuity equations were solved for each mesh vertex in conjunction with the carrier generation file that was calculated from the absorbed photon density. Top and bottom contacts were defined and forced the respective biasing as boundary conditions. The simulated axial devices are composed of a top degenerately phosphorus-doped emitter (n-type) and a boron-doped base (p-type) (Fig. 4a). The emitter is 200 nm–300 nm in height with an error function dopant distribution to account for a standard implant process⁵¹. The degenerated emitter prohibits the generation of photocurrent due to the high levels of recombination and hence serves solely for the formation of a pn-junction and as a top ohmic contact and therefore confined to a small volume as realistic implant/diffusion processes render possible. The absorbed photon density reflects the weighted absorption once the devices are subjected to solar spectral irradiance of Air Mass 1.5 (as performed in the optical part). The optical generation is calculated in the following manner: for excitation energy greater than or equal to the bandgap energy, the quantum efficiency is set to one and otherwise, it is set to zero. For each simulation mesh point the Poisson and the Continuity equations are solved where doping dependent Shockley-Read-Hall (SRH) recombination, Auger recombination, surface recombination, bandgap renormalization for degenerately doped silicon and doping dependent mobility, are all accounted for. It is assumed that the starting material is standard Czochralski silicon which is reflected in the selection of the following parameters. The rate of SRH is calculated in the following manner⁴⁶:

$$R_{\text{net}}^{\text{SRH}} = \frac{np - n_i^2}{\tau_p(n + n_1) + \tau_n(p + p_1)} \quad (1)$$

where n and p are the respective electron and hole carrier densities, n_i is the effective intrinsic density (accounting for bandgap narrowing), τ_n and τ_p are the respective electron and hole lifetimes and n_1 and p_1 are:

$$n_1 = n_{i,\text{eff}} \exp\left(\frac{E_{\text{trap}}}{kT}\right) \quad (2)$$

$$p_1 = n_{i,\text{eff}} \exp\left(\frac{-E_{\text{trap}}}{kT}\right) \quad (3)$$

where E_{trap} is the difference between the defect level and the intrinsic level and is set to zero in the current case. The dependency of SRH on doping is reflected in the dependency of carrier lifetime on doping:

$$\tau_{\text{dop}}(N_{A,0} + N_{D,0}) = \tau_{\text{min}} + \frac{\tau_{\text{max}} - \tau_{\text{min}}}{1 + \left(\frac{N_{A,0} + N_{D,0}}{N_{\text{ref}}}\right)} \quad (4)$$

where $N_{A,0}$ and $N_{D,0}$ are the chemically active acceptor and donor concentrations, respectively. In the present case $\gamma = 1$, $\tau_{\text{min}} = 0$, $N_{\text{ref}} = 10^{16} \text{ cm}^{-3}$ for both holes and electrons and $\tau_{\text{min}} = 10^{-5} \text{ sec}$ and 3×10^{-6} for electrons and holes, respectively.

The rate of Auger recombination is calculated in the following manner⁴⁶:

$$R_{\text{net}}^{\text{A}} = (C_n n + C_p p) \left(np - n_{i,\text{eff}}^2 \right) \quad (5)$$

where C_n and C_p are set to $2.9 \times 10^{-31} \text{ cm}^6\text{s}^{-1}$ and $1 \times 10^{-31} \text{ cm}^6\text{s}^{-1}$, respectively⁵².

- Lewis, N. S. Toward cost-effective solar energy use. *Science* **315**, 798–801 (2007).
- Green, M. A. The path to 25% silicon solar cell efficiency: History of silicon cell evolution. *Prog. Photovoltaics Res. Appl.* **17**, 183–189 (2009).
- Haschke, J., Amkreutz, D., Korte, L., Ruske, F. & Rech, B. Towards wafer quality crystalline silicon thin-film solar cells on glass. *Sol. Energy Mater. Sol. Cells* **128**, 190–197 (2014).
- Garin, M. et al. All-silicon spherical-Mie-resonator photodiode with spectral response in the infrared region. *Nat. Commun.* DOI:10.1038/ncomms4440 (2014).
- Kayes, B. M., Atwater, H. A. & Lewis, N. S. Comparison of the device physics principles of planar and radial p-n junction nanorod solar cells. *J. Appl. Phys.* DOI:10.1063/1.1901835 (2005).
- Kosten, E. D., Warren, E. L. & Atwater, H. A. Ray optical light trapping in silicon microwires: exceeding the $2n^2$ intensity limit. *Opt. Express* **19**, 3316–3331 (2011).
- Gharghi, M. On the design and applicability of nanowire solar cells using low-grade semiconductors. *J. Appl. Phys.* DOI:10.1063/1.3679134 (2012).
- Voigt, F., Stelzner, T. & Christiansen, S. Geometrical optimization and contact configuration in radial pn junction silicon nanorod and microrod solar cells. *Prog. Photovoltaics Res. Appl.* **21**, 1567–1579 (2013).
- Kelzenberg, M. D. et al. High-performance Si microwire photovoltaics. *Energy Environ. Sci.* **4**, 866–871 (2011).
- Li, J. et al. Si nanopillar array optimization on Si thin films for solar energy harvesting. *Appl. Phys. Lett.* DOI:10.1063/1.3186046 (2009).
- Garnett, E. C. & Yang, P. Silicon nanowire radial p-n junction solar cells. *J. Am. Chem. Soc.* **130**, 9224–9225 (2008).
- Garnett, E. & Yang, P. Light trapping in silicon nanowire solar cells. *Nano Lett.* **10**, 1082–1087 (2010).
- Yoshida, A., Motoyama, M., Kosaku, A. & Miyamoto, K. Antireflective Nanoprotuberance Array in the Transparent Wing of a Hawkmoth, *Cephonodes hylas*. *Zoological Science* **14**, 737–741 (1997).
- Parker, A. R. & Townley, H. E. Biomimetics of photonic nanostructures. *Nat. Nanotechnol.* **2**, 347–353 (2007).
- Huang, Y.-F. et al. Improved broadband and quasi-omnidirectional anti-reflection properties with biomimetic silicon nanostructures. *Nat. Nanotechnol.* **2**, 770–774 (2007).
- Jeong, S., McGehee, M. D. & Cui, Y. All-back-contact ultra-thin silicon nanocone solar cells with 13.7% power conversion efficiency. *Nat. Commun.* DOI:10.1038/ncomms3950 (2013).
- Spinelli, P., Verschuuren, M. A. & Polman, A. Broadband omnidirectional antireflection coating based on subwavelength surface Mie resonators. *Nat. Commun.* DOI:10.1038/ncomms1691 (2012).
- Curcio, C. A., Sloan, K. R., Kalina, R. E. & Hendrickson, a. E. Human photoreceptor topography. *J. Comp. Neurol.* **292**, 497–523 (1990).
- Borwein, B., Borwein, D., Medeiros, J. & McGowan, J. W. The ultrastructure of monkey foveal photoreceptors, with special reference to the structure, shape, size, and spacing of the foveal cones. *Am. J. Anat.* **159**, 125–146 (1980).
- Westheimer, G. Directional sensitivity of the retina: 75 years of Stiles-Crawford effect. *Proc. Biol. Sci.* **275**, 2777–2786 (2008).
- Stiles, W. S. The directional sensitivity of the retina and the spectral sensitivities of the rods and cones. *Proc. R. Soc. London. Ser. B, Biol. Sci.* **127**, 64–105 (1939).
- Enoch, J. M. Optical Properties of the Retinal Receptors. *Journal of the Optical Society of America* **53**, 71–85 (1963).
- Provis, J. M., Dubis, A. M., Maddess, T. & Carroll, J. Adaptation of the central retina for high acuity vision: cones, the fovea and the avascular zone. *Prog. Retin. Eye Res.* **35**, 63–81 (2013).
- Masland, R. H. The fundamental plan of the retina. *Nat. Neurosci.* **4**, 877–886 (2001).
- Williams, D. R. Seeing through the photoreceptor mosaic. *Trends Neurosci.* **9**, 193–198 (1986).
- Vukusic, P. & Sambles, J. R. Photonic structures in biology. *Nature* **424**, 852–855 (2003).



27. Schmitt, S. W. *et al.* Nanowire arrays in multicrystalline silicon thin films on glass: a promising material for research and applications in nanotechnology. *Nano Lett.* **12**, 4050–4054 (2012).
28. Kelzenberg, M. D. *et al.* Enhanced absorption and carrier collection in Si wire arrays for photovoltaic applications. *Nat. Mater.* **9**, 239–244 (2010).
29. Hu, L. & Chen, G. Analysis of optical absorption in silicon nanowire arrays for photovoltaic applications. *Nano Lett.* **7**, 3249–3252 (2007).
30. Wang, W., Wu, S., Reinhardt, K., Lu, Y. & Chen, S. Broadband light absorption enhancement in thin-film silicon solar cells. *Nano Lett.* **10**, 2012–2018 (2010).
31. Bohren, C. F. & Huffman, D. R. *Absorption and Scattering of Light by Small Particles* (Wiley-VCH, Weinheim, 1998).
32. Kim, S. K. *et al.* Tuning light absorption in core/shell silicon nanowire photovoltaic devices through morphological design. *Nano Lett.* **12**, 4971–4976 (2012).
33. Kim, S.-K. *et al.* Design of nanowire optical cavities as efficient photon absorbers. *ACS Nano* **8**, 3707–3714 (2014).
34. Cao, L. *et al.* Engineering light absorption in semiconductor nanowire devices. *Nat. Mater.* **8**, 643–647 (2009).
35. Cao, L. *et al.* Semiconductor nanowire optical antenna solar absorbers. *Nano Lett.* **10**, 439–445 (2010).
36. Krogstrup, P. *et al.* Single-nanowire solar cells beyond the Shockley–Queisser limit. *Nat. Photonics* **7**, 306–310 (2013).
37. Brönstrup, G. *et al.* Optical Properties of Individual Silicon Nanowires for Photonic Devices. *ACS Nano* **4**, 7113–7122 (2010).
38. Brönstrup, G. *et al.* Statistical model on the optical properties of silicon nanowire mats. *Phys. Rev. B* DOI:10.1103/PhysRevB.84.125432 (2011).
39. Quinn, T. J. The absorptivity of a specularly reflecting cone for oblique angles of view. *Infrared Phys.* **21**, 123–126 (1981).
40. Kippenberg, T. J., Tchebotareva, A. L., Kalkman, J., Polman, A. & Vahala, K. J. Purcell-factor-enhanced scattering from Si nanocrystals in an optical microcavity. *Phys. Rev. Lett.* DOI:10.1103/PhysRevLett.103.027406 (2009).
41. Christesen, J. D. *et al.* Design principles for photovoltaic devices based on Si nanowires with axial or radial p–n junctions. *Nano Lett.* **12**, 6024–6029 (2012).
42. Roulston, D. J., Arora, N. D. & Chamberlain, S. G. Modeling and measurement of minority-carrier lifetime versus doping in diffused layers of n⁺–p silicon diodes. *IEEE Transactions on Electron Devices* **29**, 284–291 (1982).
43. Fossum, J. G. Computer-aided numerical analysis of silicon solar cells. *Solid-State Electronics* **19**, 269–277 (1976).
44. Fossum, J. G. & Lee, D. S. A physical model for the dependence of carrier lifetime on doping density in nondegenerate silicon. *Solid-State Electronics* **25**, 741–747 (1982).
45. Fossum, J. G., Mertens, R. P., Lee, D. S. & Nijs, J. F. Carrier recombination and lifetime in highly doped silicon. *Solid-State Electronics* **26**, 569–576 (1983).
46. Sze, S. M. *Physics of Semiconductor Devices*. (Wiley, New-York, 1981).
47. Allen, J. E. *et al.* High-resolution detection of Au catalyst atoms in Si nanowires. *Nat. Nanotechnol.* **3**, 168–173 (2008).
48. Hoex, B., Gielis, J. J. H., van de Sanden, M. C. M. & Kessels, W. M. M. On the c-Si surface passivation mechanism by the negative-charge-dielectric Al₂O₃. *J. Appl. Phys.* DOI:10.1063/1.3021091 (2008).
49. Spinelli, P. & Polman, A. Light Trapping in Thin Crystalline Si Solar Cells Using Surface Mie Scatterers. *IEEE J. Photovoltaics* **4**, 554–559 (2014).
50. Palik, E. D. *Handbook of Optical Constants of Solids*. (Academic, 1985).
51. Schmitt, S. W. *et al.* Probing photo-carrier collection efficiencies of individual silicon nanowire diodes on a wafer substrate. *Nanoscale* DOI:10.1039/c4nr01258e (2014).
52. Lochmann, W. & Haugt, A. Phonon-assisted Auger recombination in Si with direct calculation of the overlap integrals. *Solid State Commun.* **35**, 553–556 (1980).

Author contributions

G.S. and S.W.S. established the idea of light funnels. G.S. performed the simulations. S.W.S. and H.E. carried out the fabrication and the SEM micrographs of the light-funnels. G.B. and S.C. contributed discussions of experimental and theoretical results to the paper.

Additional information

Supplementary information accompanies this paper at <http://www.nature.com/scientificreports>

Competing financial interests: The authors declare no competing financial interests.

How to cite this article: Shalev, G., Schmitt, S.W., Embrechts, H., Brönstrup, G. & Christiansen, S. Enhanced photovoltaics inspired by the fovea centralis. *Sci. Rep.* **5**, 8570; DOI:10.1038/srep08570 (2015).



This work is licensed under a Creative Commons Attribution 4.0 International License. The images or other third party material in this article are included in the article's Creative Commons license, unless indicated otherwise in the credit line; if the material is not included under the Creative Commons license, users will need to obtain permission from the license holder in order to reproduce the material. To view a copy of this license, visit <http://creativecommons.org/licenses/by/4.0/>