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Design and analysis of a plasmonic-nanorod-enhanced lead-free inorganic perovskite/silicon heterojunction tandem solar cell exceeding the Shockley–Queisser limit

Md. Sad Abdullah Sami,  Arpan Sur  and Ehsanur Rahman *

The pursuit of sustainable and highly efficient energy conversion necessitates a transition from toxic and unstable materials to environmentally friendly alternatives. This work presents a simulation-based numerical investigation of a fully inorganic, lead-free tandem solar cell that employs cesium tin-germanium tri-iodide (CsSnGeI_3) as the top cell absorber and crystalline silicon (c-Si) as the bottom cell absorber, configured in a silicon heterojunction (SHJ) arrangement. Utilizing CsSnGeI_3 as a lead-free perovskite presents a promising solution to the toxicity concerns associated with conventional lead-based perovskites. To further increase near-infrared absorption and reduce the required thickness of the c-Si layer, an ultra-thin gallium antimonide auxiliary absorber is integrated into the SHJ bottom cell. Optical and electrical simulations, conducted using finite-difference time-domain and drift-diffusion modelling, demonstrate that the optimized tandem structure attains a power conversion efficiency of 34.93%, surpassing the Shockley–Queisser limit established for single-junction Si cells. Furthermore, the optimized device showcases an open-circuit voltage of 1.93 V, a short-circuit current density of 21.30 mA cm^{-2} , and a fill factor of 84.74%. Performance is additionally enhanced by incorporating cylindrical gold nanorods within a Si_3N_4 dielectric medium positioned at the rear of the bottom cell, thus amplifying light absorption through plasmonic effects. Notably, the tandem cell sustains high efficiency even without the plasmonic structure, thereby providing flexibility for cost-effective fabrication. This work underscores the viability of all-inorganic, lead-free tandem cells for next-generation photovoltaics, guided by simulated results that pave the way for high-efficiency, non-toxic solar energy solutions and further experimental validation.

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1. Introduction

The world population has increased exponentially since the 19th century, reaching around 8 billion in 2022, and is projected to exceed 9 billion by 2040.¹ Due to this rise in population and lifestyle changes, energy demand is increasing daily. Global energy demand is expected to surpass 35 000 terawatt hours (TWh) by 2040.² The energy produced today remains predominantly dependent on fossil fuels, but the availability of fossil fuel sources is expected to diminish around 2112.³ A possible solution to this issue is the use of renewable energy. Among renewable energy sources, solar energy has the most potential.⁴ Historically, semiconductor materials have played a crucial role in photovoltaic (PV) cells, with silicon (Si) being utilized in PV cells for about 70 years. Due to this extended development period, Si-based PV cells dominate the current solar cell

industry, accounting for 97% of total PV cell production in 2023.⁵ Si is abundant, non-toxic, stable, and has a bandgap of 1.12 eV, which makes it a suitable candidate for absorbing a wide range of the solar spectrum. The fabricated crystalline Si (c-Si) single-junction solar cell has achieved a remarkable power conversion efficiency (PCE) of 27.09%.⁶ The maximum possible PCE for a single-junction Si solar cell is restricted to 29.43%.⁷ Si has an indirect bandgap, which contributes to its low photocurrent generation. Its thickness must be significantly increased to produce a sufficient photocurrent with the current industry standard for c-Si absorber layer thickness at $180 \mu\text{m}$.⁸

Single-junction solar cells are constrained by the Shockley–Queisser (SQ) limit, which defines the maximum theoretical efficiency achievable by a single p–n junction photovoltaic device under standard illumination conditions.⁹ This limit is approximately 33% under standard AM 1.5G solar spectrum conditions. Over the years, different approaches have been taken to increase the PCE of Si PV cells. Some prominent ones include passivated emitter and rear contact (PERC) cells and Si heterojunction (SHJ) solar cells. PERC solar cells have achieved

Department of Electrical and Electronic Engineering, Bangladesh University of Engineering and Technology, Dhaka, 1000, Bangladesh. E-mail: ehsaneee@eee.buet.ac.bd



a maximum recorded efficiency of 24.5%, which was produced in 2022 by Trina Solar.¹⁰ In PERC solar cells, the rear side is passivated so that unabsorbed light can be reflected from the back of the cell. The top layer of a PERC solar cell is textured to enhance the absorption of incident light.¹¹ In 2023, LONGi achieved a significant advancement in Si solar cell technology by developing a solar cell with an efficiency of 27.09%, utilizing a Si heterojunction (SHJ).⁶ Here, a thin layer of a-Si was used with a 110 μm thick c-Si layer to produce a SHJ solar cell. Thus, a higher PCE was achieved. However, this efficiency remains below the theoretical limit, prompting the exploration of various strategies to further enhance the PCE of Si solar cells. Using a multi-junction tandem structure is a prominent one among them. In Si-based tandem cells, a suitable top cell is chosen, and Si is kept as the absorber layer for the bottom cell. The tandem cell's top cell absorber layer has a bandgap typically around 1.5 to 1.9 eV. This ensures that both the top and bottom cells can absorb an adequate solar spectrum, which is crucial for the tandem structure to function correctly.¹²

Various materials have been explored as potential top cell absorbers in Si-based tandem solar cells. Among them, group III–V materials,¹³ kesterites,¹⁴ and perovskites,¹⁵ have received considerable attention. Due to their direct bandgap, these materials exhibit strong absorption and photocurrent generation capabilities. Specifically, the perovskite crystals demonstrate unique properties, including a tunable bandgap, low Auger recombination loss, high carrier mobility, high absorption coefficient, and a long charge diffusion length.^{15,16} The general chemical formula of perovskite compounds is ABX_3 , where A typically denotes a larger organic or inorganic cation, B represents a smaller metallic cation, and X corresponds to an anion, commonly a halide.^{17,18} Moreover, perovskite materials feature a tunable bandgap of 1.3 to 2.2 eV, allowing seamless integration into tandem structures for targeted solar spectrum absorption. Additionally, perovskites exhibit greater resistance to crystallographic defects than conventional semiconductor materials.^{19,20} Perovskite-on-Si tandem solar cells have recently garnered significant attention within the PV industry. As of September 2024, LONGi has reportedly fabricated a perovskite/Si tandem solar cell with an impressive efficiency of 34.85%.²¹ The quest for an optimal perovskite material that can effectively complement existing Si technology in tandem architectures remains ongoing. While a variety of perovskite compositions were initially proposed, lead-based perovskites have emerged as the primary focus of continued investigation, with methylammonium lead iodide (MAPbI_3) being the most extensively studied.^{22–26} It possesses a direct bandgap of 1.55 eV, enabling it to absorb a significant portion of the solar spectrum with a thickness of only a few hundred nanometers, compared to 100–200 micrometers in the case of Si. It also has a lower fabrication cost compared to Si,²⁷ making it an industry favourite. Moreover, it is efficient for transporting positive and negative charge carriers over long distances. Under the AM 1.5G standard solar spectrum, the electron and hole diffusion lengths have been shown to exceed 175 μm .¹⁶ Thus, MAPbI_3 can help build a tandem cell that can potentially achieve a PCE beyond 41%.²³

Despite their numerous advantages, lead-based perovskites present significant drawbacks, particularly concerning environmental and health-related issues. The inherent toxicity of lead compounds poses a major obstacle to the long-term sustainability of lead-based perovskite solar cells.^{28–30} Lead ions have the potential to contaminate soil and water resources, resulting in adverse effects on human health, as well as on animal and plant life.^{31,32} For instance, MAPbI_3 , commonly used as the light-absorbing layer in perovskite solar cells, typically has a thickness of around 500 nm, which corresponds to a lead content of approximately 0.6 g m^{-2} .^{33,34} Consequently, the development of lead-free alternatives has become a critical research priority for advancing the commercial viability and environmental safety of perovskite solar technologies. Lead ion in the perovskite structure has been replaced by alternative ions such as Sn^{2+} , Cu^{2+} , Bi^{3+} , Mn^{2+} , and Ge^{2+} , which are less toxic than Pb^{2+} , and could be utilized to form a lead-free perovskite structure. Tin-based perovskites, for instance, offer benefits like reduced toxicity, enhanced stability, and improved environmental compatibility.^{18,35,36} Chen *et al.*³⁷ has shown that partial substitution of Sn(II) with Ge(II) in CsSnI_3 , forming the mixed-cation perovskite $\text{CsSn}_{0.5}\text{Ge}_{0.5}\text{I}_3$, significantly enhances film stability. This improvement in structural robustness is attributed to the favourable Goldschmidt tolerance factor (0.94) and octahedral factor (0.4) associated with the alloy. Additionally, the high oxidative reactivity of Ge(II) facilitates the rapid formation of a uniform, ultrathin (<5 nm) native oxide layer on the surface, effectively passivating the material and conferring superior environmental stability compared to the benchmark MAPbI_3 perovskite. These characteristics not only enhance film-level chemical stability but also indicate improved device-level operational reliability under exposure to heat, moisture, and light stressors. Therefore, CsSnGeI_3 is a promising candidate for stable long-term photovoltaic performance, especially when combined with inorganic charge transport layers like TiO_2 and Cu_2O , which are also known for their chemical durability.^{38,39}

Due to these advantages, CsSnGeI_3 has recently been explored as a lead-free alternative to MAPbI_3 . Moreover, CsSnGeI_3 has a similar bandgap to MAPbI_3 , making it easy to replace MAPbI_3 in a tandem-based structure with CsSnGeI_3 as the absorber layer.^{40–43} Lead-based organic perovskites are typically used in perovskite/Si tandem cells to ensure high efficiency.^{44,45} At times, these lead-based organic perovskite/Si tandem cells can also surpass the SQ limit.⁴⁵ Sarker *et al.*⁴⁶ has exceeded the SQ limit by using a lead-free perovskite/Si tandem cell. Moreover, the perovskite used in that study was an organic one, which can potentially cause instability in the structure. However, utilizing both lead-free and inorganic perovskite materials in a perovskite/Si tandem structure typically results in a lower PCE compared to their lead-based organic counterparts.^{42,47} Moreover, previously reported perovskite/Si tandem structures neglected the issue of significant c-Si layer thickness (usually exceeding 100 μm).^{44–46,48}

To address these challenges, this study has thoroughly investigated the CsSnGeI_3 /Si tandem solar structure. This study has achieved a PCE beyond the SQ limit comparable to lead-based organic perovskite/Si tandem cells by using a lead-free



inorganic perovskite. Moreover, this study has accomplished this while also minimizing the thickness of the c-Si absorber layer ($2\ \mu\text{m}$). Gallium antimonide (GaSb) is added to the bottom SHJ cell as an auxiliary absorber layer, improving absorption specially in the near-infrared (NIR) range.⁴⁹ Furthermore, plasmonic gold (Au) nanorods are incorporated at the rear side of the tandem cell to improve its performance.^{50–52} Although the incorporation of nanorods on the rear side of solar cells has been previously investigated, their integration into a $\text{CsSnGeI}_3/\text{Si}$ tandem architecture has yet to be reported. Consequently, this study represents the inaugural demonstration of this approach, with the objective of enhancing light trapping and overall device performance within lead-free tandem configurations. Initially, optimization of the thicknesses of different layers has been conducted. The thicknesses of CsSnGeI_3 and c-Si absorber layers, along with the p-doped and n-doped a-Si layers, titanium dioxide (TiO_2) and cuprous oxide (Cu_2O) transport layers, have been optimized by analyzing generated contour plots of PCE, open-circuit voltage (V_{oc}), short-circuit current density (J_{sc}), and fill factor (FF). Parametric sweeps

have also been performed to determine the optimal thicknesses of the anti-reflecting coating (ARC), GaSb, and rear passivation layers. This study further explores an advanced tandem solar cell architecture by incorporating cylindrical-shaped Au nanoparticles embedded in a dielectric medium at the rear side of the bottom silicon cell. A comprehensive investigation was conducted to determine the optimal dielectric material, nanoparticle geometry, and particle radius to enhance light management and improve device performance. In addition, the doping concentrations of all layers within the tandem structure were systematically optimized. The combined effect of these enhancements led to a significant improvement in the PCE of the tandem cell, enabling it to exceed the SQ efficiency limit.

2. Device structure and simulation methodology

2.1. Device structure

Fig. 1 shows the tandem solar cell structure used in this study, where CsSnGeI_3 and Si are the primary absorbers of the top and

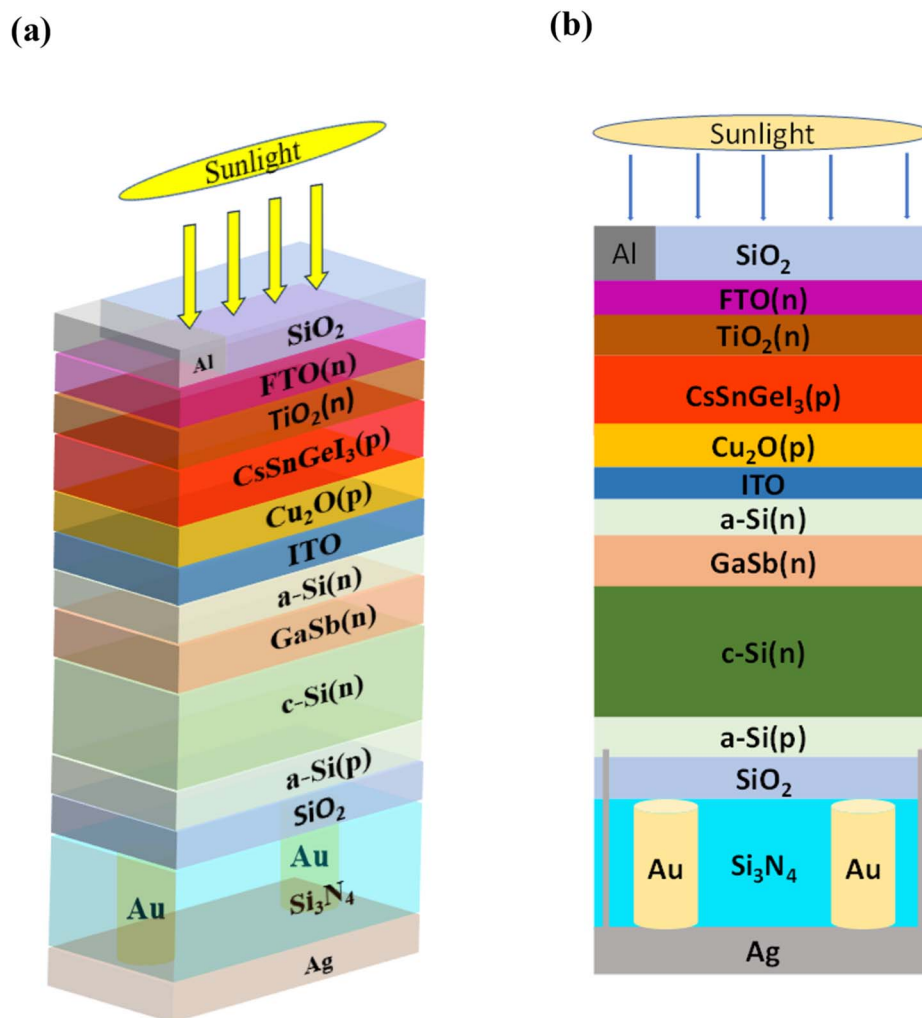


Fig. 1 Cross-sectional schematic of (a) 3D and (b) 2D views of the $\text{CsSnGeI}_3/\text{Si}$ tandem solar cell incorporating a plasmonic back reflector, where cylindrical Au nanorods are embedded in Si_3N_4 dielectric medium. An ITO recombination layer connects the perovskite top cell and the SHJ bottom cell. The SiO_2 layer on the top serves as an anti-reflective coating (ARC) layer, while the rear SiO_2 layer functions as a passivation layer.



bottom cells, respectively. At the top of the structure, there is an ARC layer made of SiO_2 . Aluminium (Al) serves as the finger electrode for the tandem solar cell. The top cell features a perovskite absorber with the following layer configuration: $\text{FTO}/\text{TiO}_2/\text{CsSnGeI}_3/\text{Cu}_2\text{O}$. Fluorine-doped Tin Oxide (FTO), a transparent conductive oxide (TCO) layer, serves as the front electrode in this tandem structure, enabling light transmission and efficient electron collection.⁵³ The electron transport layer (ETL) consists of a thin layer of TiO_2 , while the hole transport layer (HTL) is made of Cu_2O . The perovskite absorber layer is positioned between the ETL and the HTL layers. The ETL and HTL materials were selected based on the calculation and comparison of the conduction band offset (CBO) and valence band offset (VBO) of various materials.⁴⁵

In Fig. 2, various ETLs and HTLs were compared. The hole transport layers (HTLs) include Cu_2O , poly[bis(4-phenyl)(2,4,6-trimethylphenyl)amine] (PTAA), nickel(II) oxide (NiO), and cuprous iodide (CuI), with their valence band offsets being -0.03 eV, -0.14 eV, -0.14 eV, and -0.32 eV, respectively. Materials with a highly negative VBO in the HTL are undesirable because they increase the likelihood of interface recombination, which can lead to a reduction in V_{oc} .³⁸ Additionally, inorganic materials typically exhibit higher hole mobility than organic HTL materials, resulting in higher J_{sc} .⁵⁴ Cu_2O was selected as the HTL layer in the top cell due to its lowest VBO and its classification as an inorganic HTL. The ETLs illustrated in Fig. 2 include zinc oxide (ZnO), zinc selenide (ZnSe), TiO_2 , and [6,6]-phenyl- C_{61} -butyric acid methyl ester (PCBM), with their conduction band offsets being -0.36 eV, -0.19 eV, -0.1 eV, and -0.1 eV, respectively. A negative CBO is suitable for ETLs, as a positive CBO creates a barrier to the flow of photo-generated electrons.⁵⁵ An ETL with a low negative CBO is advantageous, resulting in a higher V_{oc} . Among the ETLs depicted in Fig. 2, TiO_2 and PCBM have the lowest negative CBO

of -0.1 eV. However, since PCBM is an organic ETL and organic ETLs generally have lower electron mobility than inorganic ones,⁵⁶ TiO_2 is chosen as the ETL for the top cell. An Indium Tin Oxide (ITO) layer connects the top and bottom cells by serving as a recombination layer. This layer is important to ensure continuous current flow in the tandem structure as without it, carriers would build up at the interface, and current continuity between top and bottom cells would break.⁵⁷ The bottom cell features a SHJ structure, utilizing two ultra-thin amorphous Si (a-Si) layers. The n-doped a-Si layer functions as the emitter, while the p-doped a-Si layer acts as the back surface field. Additionally, a thin layer of GaSb is used as an auxiliary absorber layer in the bottom cell, alongside a c-Si layer. On the rear side of the bottom cell, a SiO_2 layer is employed as the rear passivation layer. Beneath it lies a Si_3N_4 dielectric layer, which contains an array of cylindrical Au nanorods with a periodicity of 750 nm. At the very bottom of the structure, a Ag mirror functions as both a reflective layer and a back contact. Small contact windows are opened on the Si_3N_4 and SiO_2 layers to connect the Ag back contact with the p-doped a-Si layer.

2.2. Proposed fabrication process

Ag contact can be created by screen-printing Ag paste for the back contact, then co-firing it in a belt furnace. A thermal SiO_2 layer can be grown for surface passivation before metallization, and a Si_3N_4 dielectric layer can be applied using the Plasma Enhanced Chemical Vapor Deposition (PECVD) method.⁵⁸ Laser openings can expose the a-Si layer locally through the dielectric and rear passivation layers. This setup improves carrier selectivity and internal reflectance, which supports the high efficiency of PERC devices. Au cylindrical nanorods can be produced by refining the seed-mediated synthesis of gold nanorods,⁵⁹ which can then be embedded in the dielectric layer via a wet-coating process.⁵² Next, a thin p-type a-Si layer can be

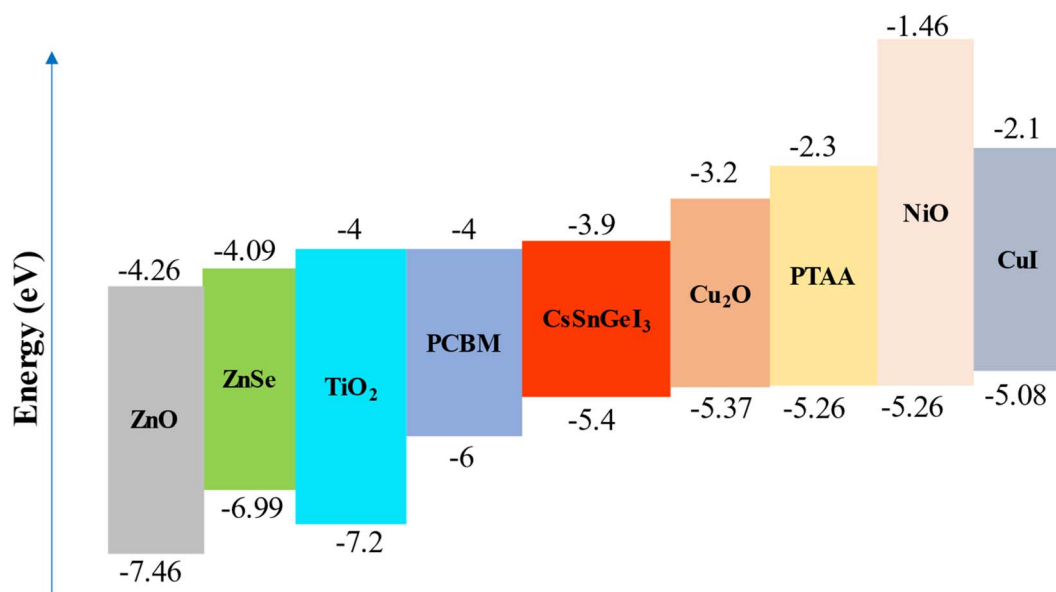


Fig. 2 Energy level diagram of CsSnGeI_3 with various ETLs and HTLs.



deposited through PECVD above the rear passivation layer. Following this, a high-quality c-Si layer can be bonded or integrated epitaxially on top of it.^{60,61} A GaSb layer can then be grown on the c-Si using Molecular Beam Epitaxy (MBE), beginning with an AlSb nucleation layer at about 450 °C.^{62–64} An n-type a-Si layer is deposited above GaSb using low-temperature PECVD. ITO on a-Si can be created through a two-step radio frequency (RF) sputtering method.⁶⁵ At first, a low-damage ITO seed layer can be applied at 50 W to minimize interface degradation, followed by a high-power deposition at 100 W to enhance conductivity and transparency. Cu₂O can be formed on ITO substrates *via* electro-deposition using an alkaline solution containing Cu²⁺ ions and a chelating agent like citric acid.³⁹ CsSnGeI₃ can be created on Cu₂O by spin-coating a precursor of CsSnGeI₃ onto the Cu₂O layer, followed by a carefully controlled annealing process.^{66,67} TiO₂ thin films can be deposited *via* a sol-gel spin-coating technique, enabling uniform coverage with controlled thickness.⁶⁸ FTO thin films can be deposited over the TiO₂ layer *via* RF magnetron sputtering at room temperature.⁶⁹ Then, a SiO₂ antireflection coating layer can be fabricated through a Physical Vapor Deposition (PVD) technique, which is based on the thermal decomposition of polydimethylsiloxane (PDMS).⁷⁰ Finally, Al can be deposited *via* electron beam evaporation under high vacuum conditions, with controlled deposition rates.⁷¹

2.3. Research methodology

The simulation consists of two parts, specifically optical and electrical simulations. Optical simulations were utilized to compute the carrier generation rate. This data served as the input for subsequent electrical simulations, from which the characteristic PV parameters, such as PCE, J_{sc} , V_{oc} , and FF, were extracted. In the optical simulations, the refractive index (n) and extinction coefficient (k) values as a function of wavelength were used to characterize the materials, enabling a rigorous exploration of their optical properties.^{72–82} Maxwell's curl equation was used to determine the light absorption and carrier generation rates through finite difference time domain (FDTD) analysis.^{83,84} This type of FDTD analysis is often used for validating experimental works.^{85,86} For the input radiation source, AM 1.5G standard solar spectrum was used. Perfectly matched layer (PML) boundary conditions were employed at the top and bottom boundaries in the vertical (Y) direction, while periodic boundary conditions were applied along the horizontal (X) axis. In the FDTD analysis, the optical electric field distribution ($\vec{E}_{op}$) was calculated inside different layers. Using the optical electric field and the imaginary part of the complex dielectric constant, the absorbed power (P_{abs}) was calculated.⁸⁷

$$P_{abs} = -\frac{1}{2} \omega \left| \vec{E}_{op}(\vec{r}, \omega) \right|^2 \text{im}\{\varepsilon(\vec{r}, \omega)\}, \quad (1)$$

where ω is the angular frequency, $\vec{E}_{op}$ is the optical electric field distribution, and $\text{im}\{\varepsilon(\vec{r}, \omega)\}$ is the imaginary part of the complex dielectric constant.

$$g(\vec{r}, \omega) = -\frac{\pi}{h} \left| \vec{E}_{op}(\vec{r}, \omega) \right|^2 \text{im}\{\varepsilon(\vec{r}, \omega)\}, \quad (2)$$

where $g(\vec{r}, \omega)$ is the electron-hole pair generation rate at position $\vec{r}$ and angular frequency ω , and h is Planck's constant.

$$G(\vec{r}) = \int g(\vec{r}, \omega) d\omega, \quad (3)$$

where $G(\vec{r})$ represents the total electron-hole pair generation rate at position $\vec{r}$ obtained by integrating $g(\vec{r}, \omega)$ over all photon frequencies above the cutoff frequency, corresponding to the optical bandgap of the absorber material.

The electrical simulation follows the optical part, with the generation rate calculated from the optical simulation used as input. At first, the top cell and bottom cell electrical simulations were done separately. Poisson's equation, drift-diffusion equations, and continuity equations were used to find the electrical characteristic metrics of the top cell and bottom cell separately.

$$-\nabla(\varepsilon_{dc} \nabla V) = q\rho, \quad (4)$$

where ε_{dc} is the DC dielectric permittivity, V is the electrostatic potential, and ρ is the net charge density.

$$J_n = q\mu_n nE + qD_n \nabla n, \quad (5)$$

where J_n is electron current density, μ_n is electron mobility, n is electron concentration, E is the electric field, and D_n is the diffusion coefficient for electrons.

$$J_p = q\mu_p pE - qD_p \nabla p, \quad (6)$$

where J_p is hole current density, μ_p is hole mobility, p is hole concentration, E is the electric field, and D_p is the diffusion coefficient for holes.

$$\frac{\partial n}{\partial t} = \frac{1}{q} \nabla J_n + G_n - R_n, \quad (7)$$

where G_n is the electron generation rate, and R_n is the electron recombination rate.

$$\frac{\partial p}{\partial t} = -\frac{1}{q} \nabla J_p + G_p - R_p, \quad (8)$$

where G_p is the hole generation rate, and R_p is the hole recombination rate.

The eqn (4) to (8) were solved self-consistently to obtain the electrical characteristic metrics of the top and bottom cells, such as J - V characteristics, PCE, V_{oc} , J_{sc} , and FF. After performing the individual electrical simulation for the top cell and bottom cell separately, the tandem cell electrical simulation was performed. For the tandem cell simulation, the series circuit rules were applied.

$$V_{tandem} = V_{top} + V_{bottom}, \quad (9)$$

where V_{top} , V_{bottom} , and V_{tandem} represent the applied bias voltages for the top, bottom, and tandem cells at the matched current of the tandem cell, respectively.

$$J_{tandem} = \min(J_{top}, J_{bottom}), \quad (10)$$



Table 1 Initial thickness and doping concentration of different layers

Layer	Initial thickness (nm)	Initial acceptor concentration (cm^{-3})	Initial donor concentration (cm^{-3})
ARC	80	—	—
FTO	30	—	10^{19}
TiO ₂	30	—	10^{18}
CsSnGeI ₃	—	10^{15}	—
Cu ₂ O	30	10^{18}	—
a-Si (n-doped)	15	—	10^{19}
GaSb	150	—	10^{16}
c-Si	—	—	10^{16}
a-Si (p-doped)	15	10^{19}	—
SiO ₂	20	—	—
Si ₃ N ₄	100	—	—

where, J_{top} , J_{bottom} , and J_{tandem} indicate the current densities for the top, bottom, and tandem cells, respectively. The PV parameters of the tandem cell, such as PCE, J_{sc} , V_{oc} , and FF, were calculated using the V_{tandem} and J_{tandem} values. The material data used in the electrical simulation, including doping concentration, carrier mobility, electron affinity, bandgap, recombination-related parameters, *etc.*, are provided in the Section S1.

3. Results and discussion

3.1. Initial thickness optimization of CsSnGeI₃ and c-Si layers

Initially, a thickness sweep was performed on the absorber layers CsSnGeI₃ and c-Si. The thicknesses of the other layers were kept fixed, and the thicknesses of these two layers were varied. The radius of cylindrical Au nanorods was kept at

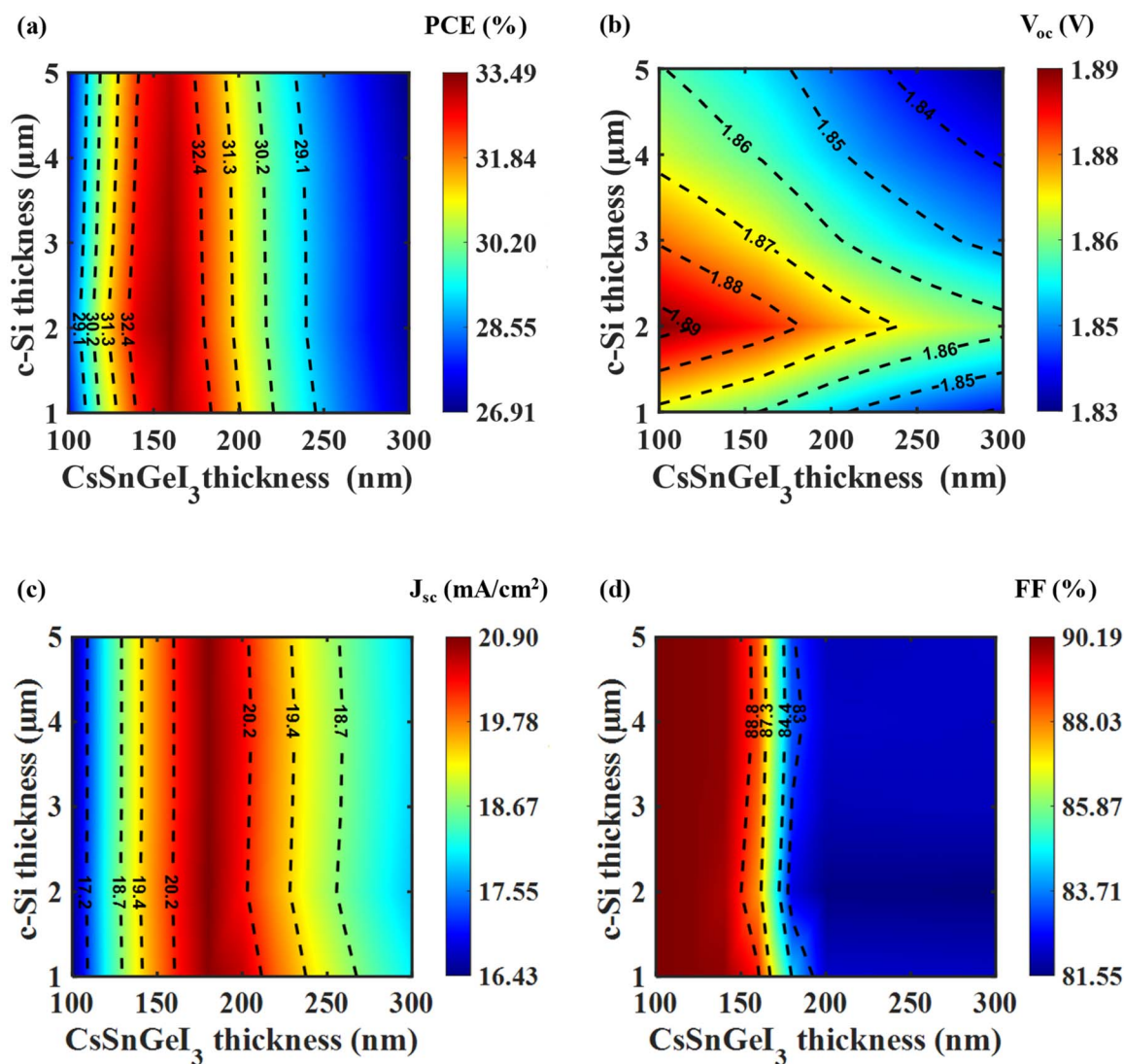


Fig. 3 Variation of (a) PCE, (b) V_{oc} , (c) J_{sc} , and (d) FF of tandem cell as a function of CsSnGeI₃ and c-Si absorber layer thicknesses. Optimal performance occurs at a CsSnGeI₃ thickness of 160 nm and a c-Si thickness of 2 μm. The efficiency is mainly determined by the thickness of the CsSnGeI₃ layer, with only a slight effect from changes in the c-Si thickness.



130 nm. The thicknesses and doping concentrations of different layers for this sweep are given in Table 1.

Fig. 3 depicts the variation of key performance metrics, namely PCE, V_{oc} , J_{sc} , and FF, as functions of the thicknesses of CsSnGeI₃ and c-Si absorber layers in a tandem solar cell setup. The peak PCE, illustrated in Fig. 3(a), occurs when the CsSnGeI₃ layer is 160 nm thick and the c-Si layer measures 2 μm , achieving a value of nearly 33.5%. PCE strongly depends on the thickness of the CsSnGeI₃ layer, but shows little sensitivity to changes in the c-Si thickness beyond 2 μm . As illustrated in Fig. 3(b), the V_{oc} of the tandem structure initially increases with the thickness of the c-Si layer up to 2 μm . Beyond this thickness, a slight decrease is observed, indicating that the maximum V_{oc} is attained at this particular thickness. Furthermore, the V_{oc} progressively decreases with the increasing thickness of the perovskite layer, implying that a thinner perovskite layer is advantageous for achieving a higher V_{oc} . In Fig. 3(c), J_{sc} is mainly influenced by the CsSnGeI₃ layer, with peak values found in the 150–200 nm range.

The J_{sc} variation with c-Si thickness is minimal, indicating that the top absorber primarily controls series circuit current. In contrast, Fig. 3(d) shows that FF remains fairly constant with c-Si thickness variation, but significantly declines as the CsSnGeI₃ layer thickens. A high FF, around 90%, is recorded when the CsSnGeI₃ layer is about 170 nm thick, while values fall below 85% when the thickness exceeds 200 nm. These results suggest that CsSnGeI₃ thickness is crucial for overall device performance, with c-Si thickness playing a limited role beyond 2 μm . Therefore, the optimal configuration for the tandem cell comprises a CsSnGeI₃ thickness of 160 nm in conjunction with a c-Si thickness of 2 μm .

3.2. Thickness optimization of ETL and HTL layers

As discussed above, the initial optimized thicknesses for perovskite and c-Si layers are 160 nm and 2 μm , respectively. Moreover, both a-Si layers have an optimum thickness of 15 nm

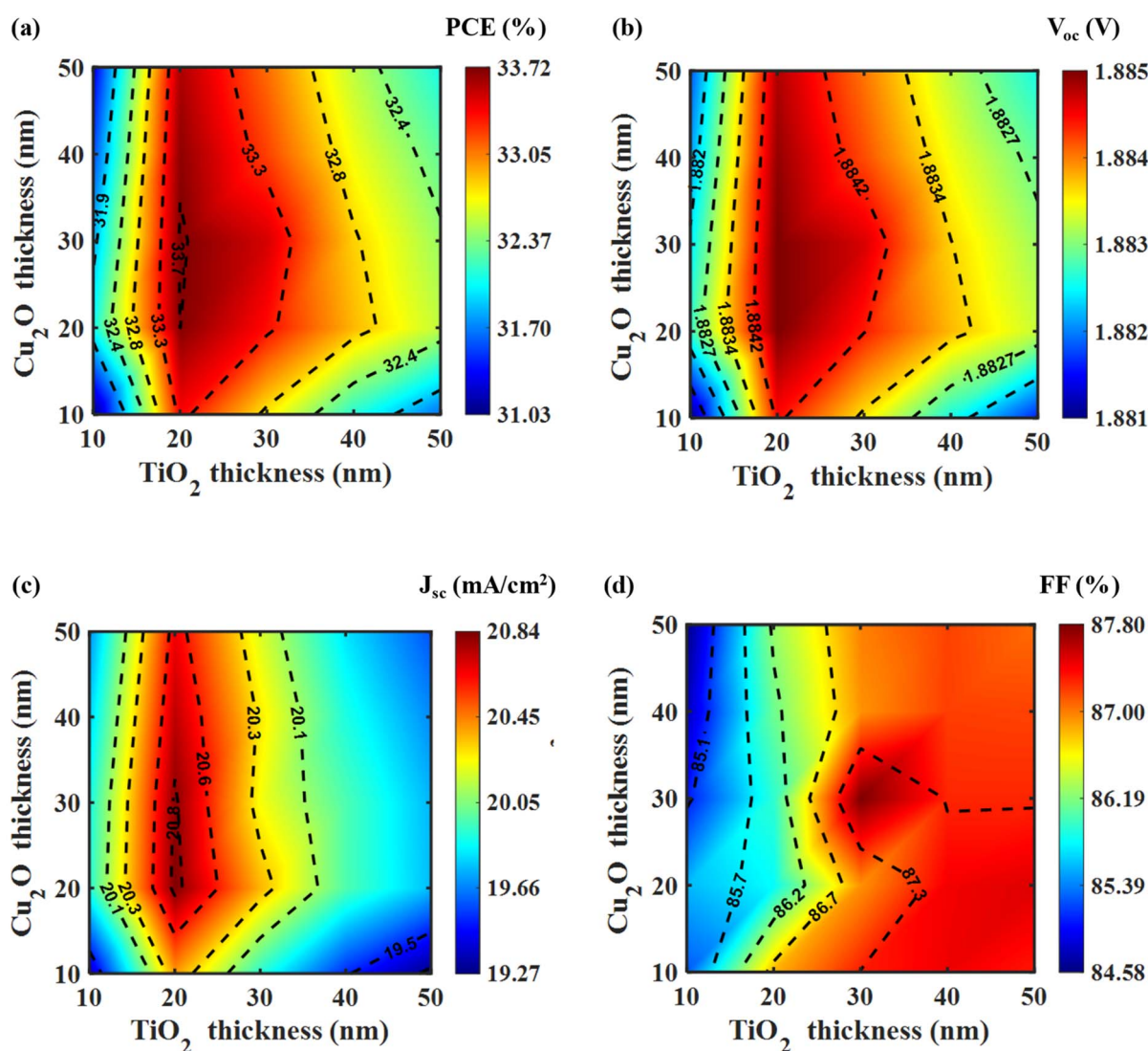


Fig. 4 Variation of (a) PCE, (b) V_{oc} , (c) J_{sc} , and (d) FF of tandem cell as a function of Cu_2O and TiO_2 transport layer thicknesses. Optimal performance is observed at a TiO_2 thickness of 20 nm and Cu_2O thickness of 30 nm, corresponding to the highest PCE and V_{oc} . While the maximum J_{sc} occurs at a Cu_2O thickness of 20 nm, the highest FF is achieved when both layers are 30 nm thick.



(see Section S2 of SI). By applying these updated thicknesses for perovskite, c-Si, and a-Si layers, while maintaining the thickness of the other layers as before, the thickness variation was performed for the ETL and HTL layers of the top cell of the tandem structure.

Fig. 4 illustrates the influence of varying layer thicknesses of the TiO_2 ETL layer and Cu_2O HTL layer on the performance metrics of the tandem solar cell. Fig. 4(a) shows that the optimum PCE is attained with a TiO_2 layer thickness of 20 nm and a Cu_2O layer thickness of 30 nm. This configuration also yields the peak value for V_{oc} as illustrated in Fig. 4(b). However, Fig. 4(c) shows a slight variation, revealing that the maximum J_{sc} occurs with a 20 nm thick Cu_2O layer instead of 30 nm, suggesting that thinner Cu_2O layers boost current generation. In case of FF, the highest value is achieved using 30 nm thickness for both layers, as seen in Fig. 4(d). These results emphasize the complex interaction between the transport layers in tandem cells and the necessity to optimize each layer thickness separately to achieve maximum PV efficiency. In summary, the tandem cell reaches optimal overall performance with a TiO_2 thickness of 20 nm and a Cu_2O thickness of 30 nm. These ultra-thin carrier transport layers enable efficient charge extraction, minimize recombination losses, and improve overall device performance. Then, the thickness optimization of GaSb, ARC, and rear passivation layers was done (see Sections S3, S4, and S5 of the SI). The optimal thicknesses of GaSb, ARC, and rear passivation layers are 150 nm, 100 nm, and 30 nm, respectively.

3.3. Optimization of plasmonic nanorods

Optical simulation was employed to compare various dielectric materials, and the resulting $J_{\text{ph_top}}$ (top cell photocurrent density) and $J_{\text{ph_bottom}}$ (bottom cell photocurrent density) were utilized to identify a suitable dielectric medium. The thickness of the dielectric layer was kept at 100 nm.

Among the dielectric materials evaluated in Table 2, Si_3N_4 emerges as a particularly advantageous option. While it

demonstrates competitive photocurrent densities in both the top and bottom cells, its broader significance resides in its established utility within Si-based PERC and SHJ solar cell architectures. Notably, its role in passivating layers within SHJ and PERC structures demonstrates its ability to reduce surface recombination losses, thus improving overall device efficiency.⁸⁸ Since the bottom cell in the tandem setup is Si-based, incorporating Si_3N_4 on the rear side enhances optical performance while also providing rear passivation. Therefore, Si_3N_4 stands out as the most advantageous dielectric medium for the nanorod-embedded plasmonic reflector layer in the proposed tandem device.

The Ag nanoparticles used in Jamil *et al.*⁸³ had a hemispherical shape with a 130 nm radius. In this study, cylindrical Au nanorods were used instead of hemispherical Ag nanoparticles. This change resulted in a slightly better photocurrent density in this tandem structure, as shown in Table 3. $J_{\text{ph_top}}$ and $J_{\text{ph_bottom}}$ were calculated through optical simulation and were compared. Both nanoparticles had a 130 nm radius for the comparison in Table 3.

Table 3 confirms that the cylindrical shape is optimal for the nanoparticles in this tandem structure. The optimal radius for these cylindrical nanorods is 150 nm (see Section S6 of the SI).

The Au nanorods with a cylindrical shape used here are based on structures proven in experiments. For example, Park *et al.* successfully created high-aspect-ratio Au nanorods and verified their crystallinity and shape using Transmission Electron Microscopy (TEM).⁵⁹ Kim *et al.* provided detailed TEM analyses of nanorods embedded in solar cells, emphasizing their size control and optical advantages.⁸⁹ Mendes *et al.* showed how plasmonic nanoparticles can be effectively embedded into dielectric layers to enhance rear-side light trapping in silicon solar cells.⁵² These findings support the structural assumptions in our simulations and show that incorporating such nanostructures into practical devices is feasible.

3.4. Optimization of doping concentration

To enhance understanding of doping effects in various layers of the top and bottom cells, sweeping was conducted with different doping concentrations during electrical simulations.

Fig. 5(a) shows that the highest PCE is achieved with an n-doped TiO_2 layer at a doping concentration of 10^{17} cm^{-3} . Fig. 5(a) and (b) reveal a consistent trend among them. The PCE, V_{oc} , and J_{sc} values of the top cell initially increase with the ETL layer's doping concentration up to 10^{17} cm^{-3} , but start to decrease if the doping concentration is further increased. Additionally, Fig. 5(a) indicates that the top cell maintains a good FF value at this concentration. Fig. 5(c) illustrates that the highest PCE was achieved with a p-doped CsSnGeI_3 layer at a doping concentration of 10^{14} cm^{-3} . Beyond this concentration, the PCE value decreases gradually. As shown in Fig. 5(d), the V_{oc} value remains relatively constant within the 10^{13} to 10^{15} cm^{-3} range. In contrast, the J_{sc} value in Fig. 5(d) peaks at 10^{14} cm^{-3} , significantly outperforming the values observed at other concentrations. Even though the highest FF occurs at 10^{17} cm^{-3}

Table 2 Photocurrent density data for various dielectric materials in the tandem structure

Dielectric material	$J_{\text{ph_top}}$ (mA cm^{-2})	$J_{\text{ph_bottom}}$ (mA cm^{-2})
MgF_2	21.2388	22.4974
Si_3N_4	21.2105	22.5704
AZO	21.2054	21.8701
ZnO	21.1697	22.3405
Al_2O_3	21.1828	22.5374

Table 3 Comparison between nanoparticle material and shape

Nanoparticle material	Nanoparticle shape	$J_{\text{ph_top}}$ (mA cm^{-2})	$J_{\text{ph_bottom}}$ (mA cm^{-2})
Ag	Hemispherical	21.174	22.3254
Au	Cylinder	21.186	22.5704



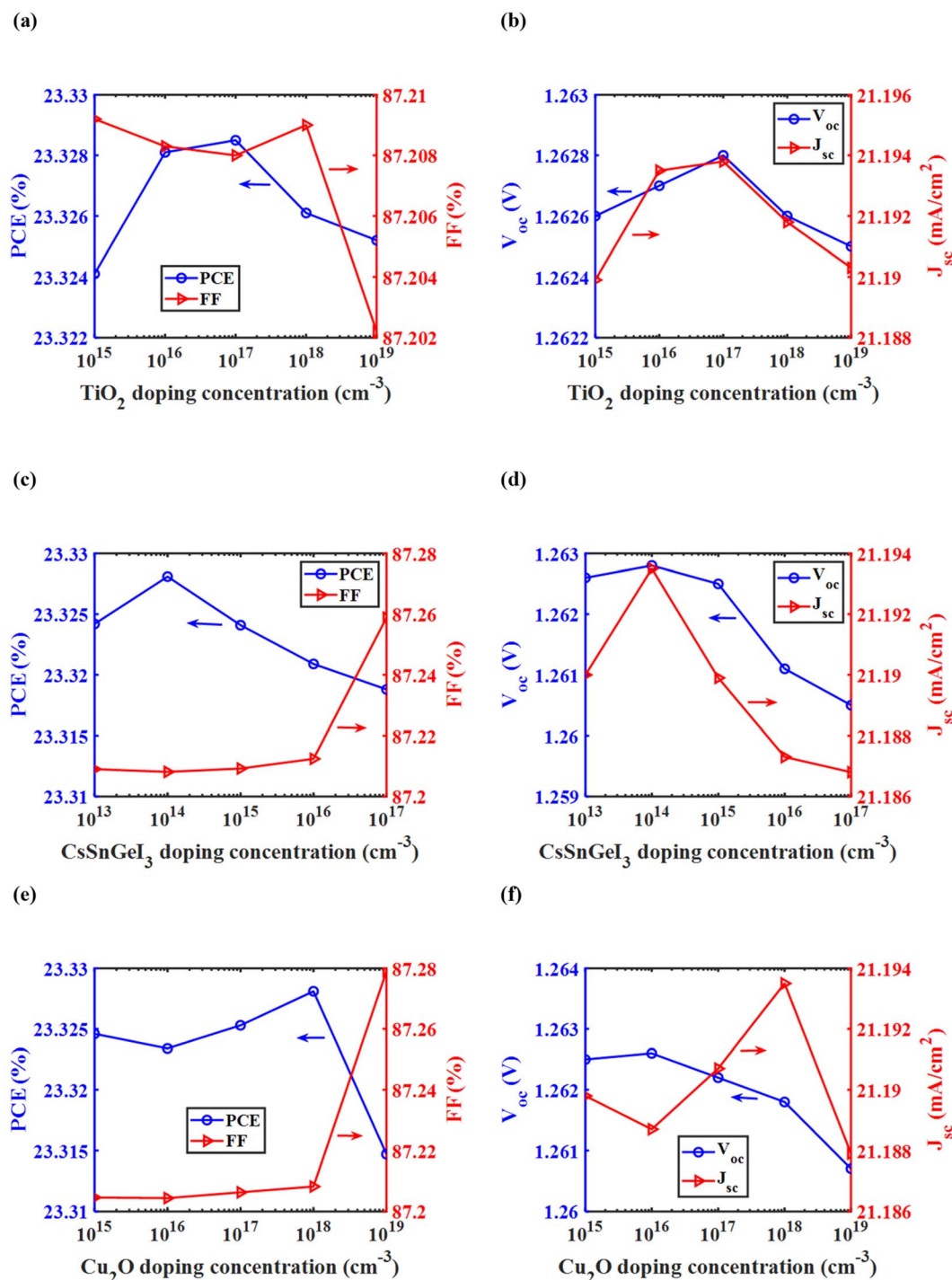


Fig. 5 PV performance metrics of the top cell as a function of doping concentration for critical layers. (a) PCE and FF, and (b) V_{oc} and J_{sc} for different doping concentration of the n-doped TiO₂ layer; (c) PCE and FF, and (d) V_{oc} and J_{sc} for different doping concentration of the p-doped CsSnGeI₃ layer; (e) PCE and FF, and (f) V_{oc} and J_{sc} for different doping concentration of the p-doped Cu₂O layer.

in Fig. 5(c), the other parameters indicate that the optimal doping concentration for the perovskite absorber is 10^{14} cm⁻³. Fig. 5(e) illustrates that the peak PCE occurs at a doping concentration of 10^{18} cm⁻³ for the p-doped Cu₂O layer. A fairly high FF is also obtained at this doping concentration of the HTL layer. Fig. 5(f) reveals that the V_{oc} declines gradually after reaching a doping concentration of 10^{16} cm⁻³. Nevertheless, the

J_{sc} is the highest at the 10^{18} cm⁻³ doping concentration, establishing it as the optimal value. Thus, based on the trends observed in Fig. 5, the optimized doping concentrations for TiO₂, CsSnGeI₃, and Cu₂O layers are 10^{17} cm⁻³, 10^{14} cm⁻³, and 10^{18} cm⁻³, respectively. These optimized values of doping concentration are crucial in improving charge transport and enhancing the overall efficiency of the device.



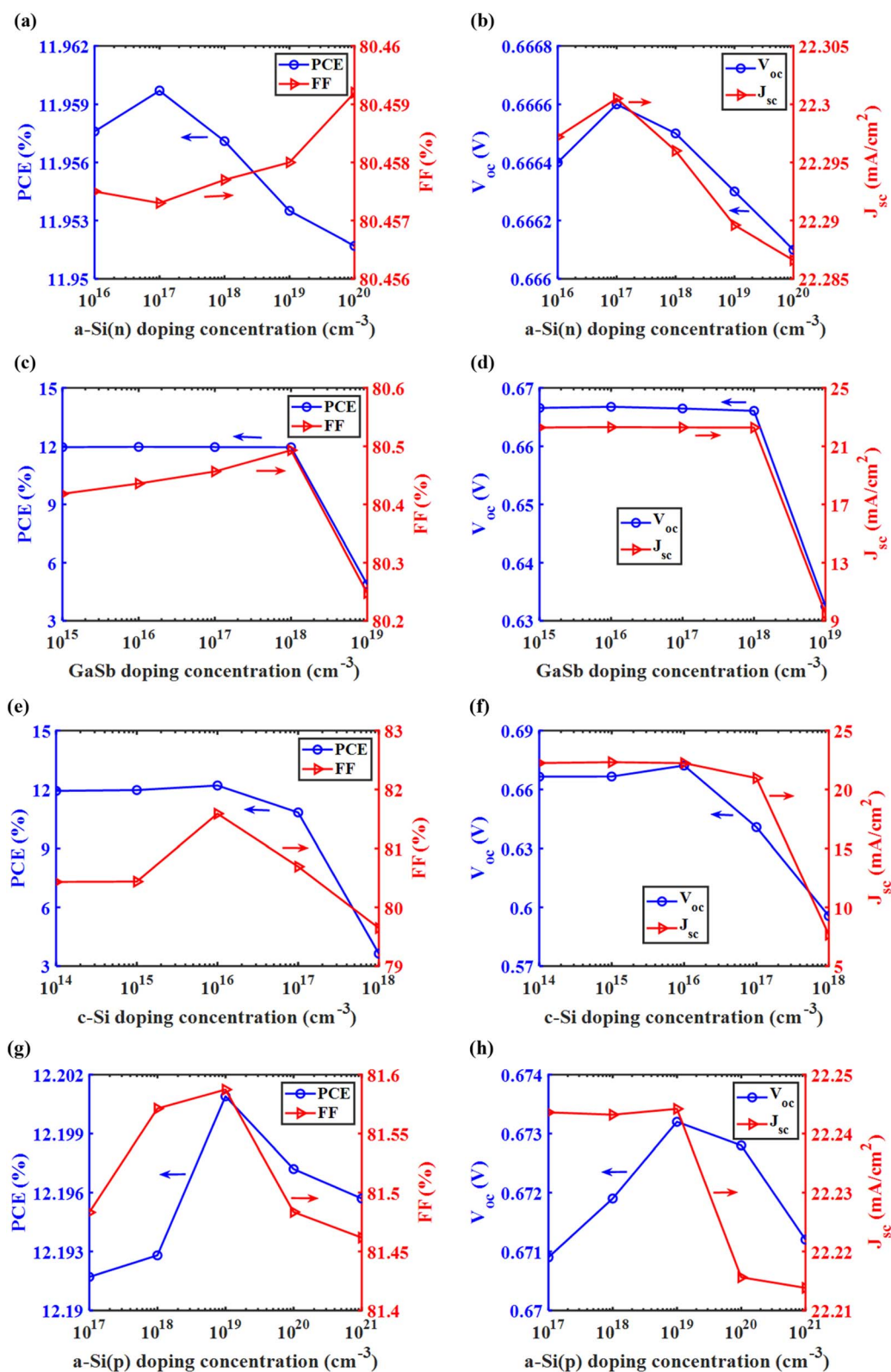


Fig. 6 PV performance metrics of the bottom cell as a function of doping concentration for critical layers. (a) PCE and FF, and (b) V_{oc} and J_{sc} for different doping concentration of the n-doped a-Si layer; (c) PCE and FF, and (d) V_{oc} and J_{sc} for different doping concentration of the n-doped GaSb layer; (e) PCE and FF, and (f) V_{oc} and J_{sc} for different doping concentration of the n-doped c-Si layer; (g) PCE and FF, and (h) V_{oc} and J_{sc} for different doping concentration of the p-doped a-Si layer.

Fig. 6(a) demonstrates that peak PCE occurs when the n-doped a-Si layer has a doping concentration of 10^{17} cm^{-3} . This peak in PCE aligns with the observation that both V_{oc} and J_{sc} reach their maximum at the same concentration, as shown in Fig. 6(b). Thus, the ideal doping concentration for the n-doped a-Si layer is 10^{17} cm^{-3} , even though FF is the lowest at this level. Fig. 6(c) and (d) show that the PCE, V_{oc} , and J_{sc} values remain almost constant across the doping concentration of the n-doped GaSb layer, which ranges from 10^{15} to 10^{18} cm^{-3} . The highest value is observed at a doping concentration of 10^{16} cm^{-3} for all three metrics, indicating that this doping concentration is optimal for this layer, even though the FF is not at its maximum at this value. Fig. 6(e) shows that PCE and FF peak at a doping concentration of 10^{16} cm^{-3} for the n-doped c-Si absorber layer. Additionally, Fig. 6(f) illustrates that J_{sc} remains nearly constant for a doping concentration ranging between 10^{14} and 10^{16} cm^{-3} , while V_{oc} achieves its maximum at 10^{16} cm^{-3} , leading to the highest PCE at this concentration. Fig. 6(g) and (h) show that PCE, FF, V_{oc} , and J_{sc} all peak at a doping concentration of 10^{19} cm^{-3} for the p-doped a-Si layer. While J_{sc} stays almost the same between 10^{17} and 10^{19} cm^{-3} , V_{oc} reaches its highest point at 10^{19} cm^{-3} , leading to the maximum PCE at this concentration. Thus, from the trends depicted in Fig. 6, the optimized doping

concentrations for n-doped a-Si, GaSb, c-Si, and p-doped a-Si layers are 10^{17} cm^{-3} , 10^{16} cm^{-3} , 10^{16} cm^{-3} , and 10^{19} cm^{-3} , respectively. These precisely adjusted doping concentrations facilitate optimal overall device efficiency.

3.5. Final thickness optimization of CsSnGeI₃ and c-Si layers

After optimization of all the layers through sweeping the thicknesses and doping levels, final thickness optimization for the CsSnGeI₃ and c-Si absorber layers was conducted. This step aimed to confirm if the prior optimizations, along with the thickness and doping variations of the various layers, affected these primary absorber layers of the tandem cell.

Fig. 7(a) demonstrates that the PCE peaks at a CsSnGeI₃ thickness of 160 nm, showing minimal variation across different c-Si thicknesses. In Fig. 7(b), the maximum V_{oc} occurs at a c-Si thickness of 2 μm , emphasizing its crucial role in influencing V_{oc} , while modifications in CsSnGeI₃ thickness result in only slight changes. Fig. 7(c) reveals that the highest J_{sc} is observed at 160 nm for the CsSnGeI₃ layer, with only minor effects from variations in the c-Si layer thickness. Lastly, Fig. 7(d) illustrates that the FF remains relatively high for the previously optimized thicknesses of both layers. These findings

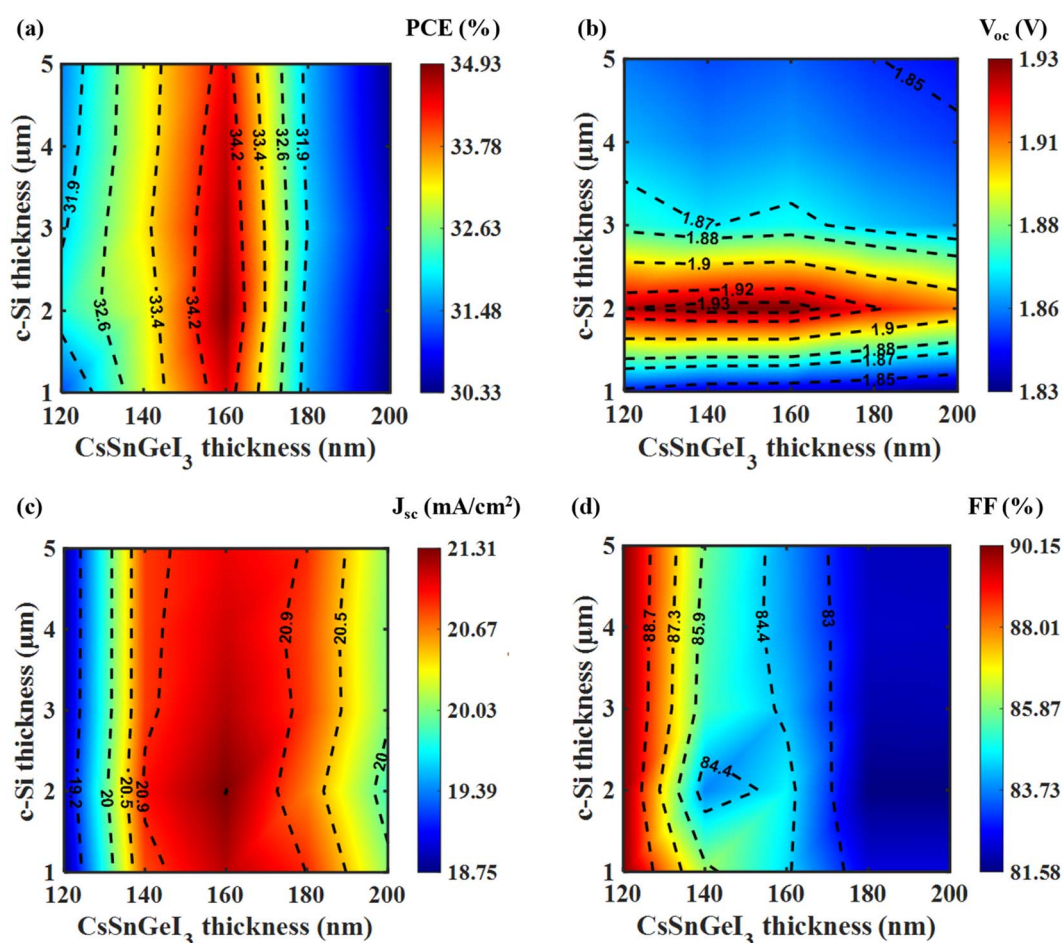


Fig. 7 Variation of (a) PCE, (b) V_{oc} , (c) J_{sc} , and (d) FF of tandem cell as a function of the thicknesses of CsSnGeI₃ and c-Si absorber layers. Optimal performance is achieved with a CsSnGeI₃ layer thickness of 160 nm and a c-Si layer thickness of 2 μm .



indicate that the ideal thicknesses for the CsSnGeI_3 and c-Si layers are 160 nm and 2 μm , respectively. This finding supports the previous thickness optimization results of these two absorber layers, confirming that changes in thickness and doping concentrations across different layers do not affect the optimal thickness of these absorber layers.

3.6. Fully optimized tandem cell characteristic metrics

After optimizing the tandem cell, the current density and power density of the tandem cell as a function of the output voltage is demonstrated in Fig. 8.

Fig. 8(a) illustrates that the short circuit current density of the optimized tandem cell is constrained by that of the top cell. Fig. 8(b) shows that the tandem structure achieves maximum power at an applied voltage of 1.625 V. A band diagram illustrating the top and bottom cell structures at the Maximum Power Point Tracking (MPPT) conditions is provided in the Section S7 of the SI, offering a visual overview of charge carrier transport throughout the tandem device.

Under MPPT conditions, a 34.93% PCE is achieved, which exceeds the SQ limit. Additional PV characteristic metrics of the optimized structure are provided in Table 4.

Table 4 summarizes the characteristic metrics for the top, bottom, and tandem cells. The PCE of the tandem cell exceeds the PCE of the individual cells. The V_{oc} of the tandem cell is particularly high, as it represents the combined V_{oc} of both the top and bottom cells. However, the J_{sc} of the tandem cell is constrained by the lower J_{sc} of the two due to the series circuit rule of the tandem structure, which in this case is the J_{sc} of the top cell.

Fig. 9(a) illustrates the excellent absorption characteristics of the tandem cell, where the top and bottom cells work harmoniously across a broad spectrum of wavelengths. The absorption spectra indicate that the tandem structure effectively captures light throughout the range, with the top cell mainly absorbing shorter wavelengths and the bottom cell focusing on longer

Table 4 Characteristic metrics of the final optimized structure under MPPT conditions

Cell type	PCE (%)	V_{oc} (V)	J_{sc} (mA cm^{-2})	FF (%)
Top cell	23.46	1.26	21.30	87.18
Bottom cell	12.13	0.67	22.12	81.61
Tandem cell	34.93	1.93	21.30	84.74

ones. This collaboration between the two cells significantly enhances the device's overall absorption, thereby boosting the efficiency of the tandem solar cell. Adding a thin GaSb layer to the SHJ bottom cell improved absorption, particularly in the NIR spectrum, and helped in reducing the thickness of the c-Si absorber layer. In Fig. 9(b), the reflection spectra of the tandem cell exhibit consistently low reflectance across the entire measured spectral range, particularly within the 450–600 nm region where solar irradiance is most intense. This suppressed reflectivity is crucial for enhancing light absorption and minimizing optical losses, thereby improving the overall photovoltaic performance. Fig. 9(c) displays the normalized AM 1.5G solar spectrum, representing the standard solar irradiance reaching Earth's surface. The spectrum peaks between 450 and 700 nm, with maximum intensity near 550 nm. This wavelength range is critical for solar energy conversion, as it encompasses the majority of available photon energy. Notably, the absorption characteristics of the tandem cell shown in Fig. 9(a) align well with this high-irradiance region, confirming that the tandem configuration is effectively optimized for efficient solar energy harvesting.

Table 5 shows that when combined with perovskites, Si-based tandem cells achieve higher PCE than other materials like kesterites. Generally, lead-based organic perovskites are associated with greater PCE values. Yet, this study reveals a significantly improved PCE using a lead-free inorganic perovskite, representing a significant advancement in perovskite-based tandem cell technology. This study achieves

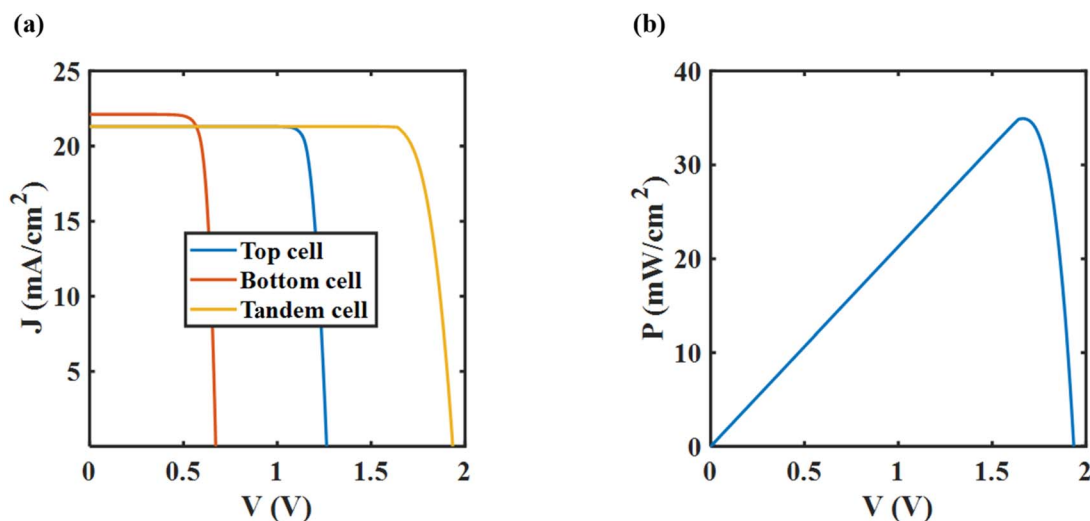


Fig. 8 (a) Current density–voltage (J – V) and (b) power density–voltage (P – V) characteristics of fully optimized $\text{CsSnGeI}_3/\text{Si}$ tandem cell.

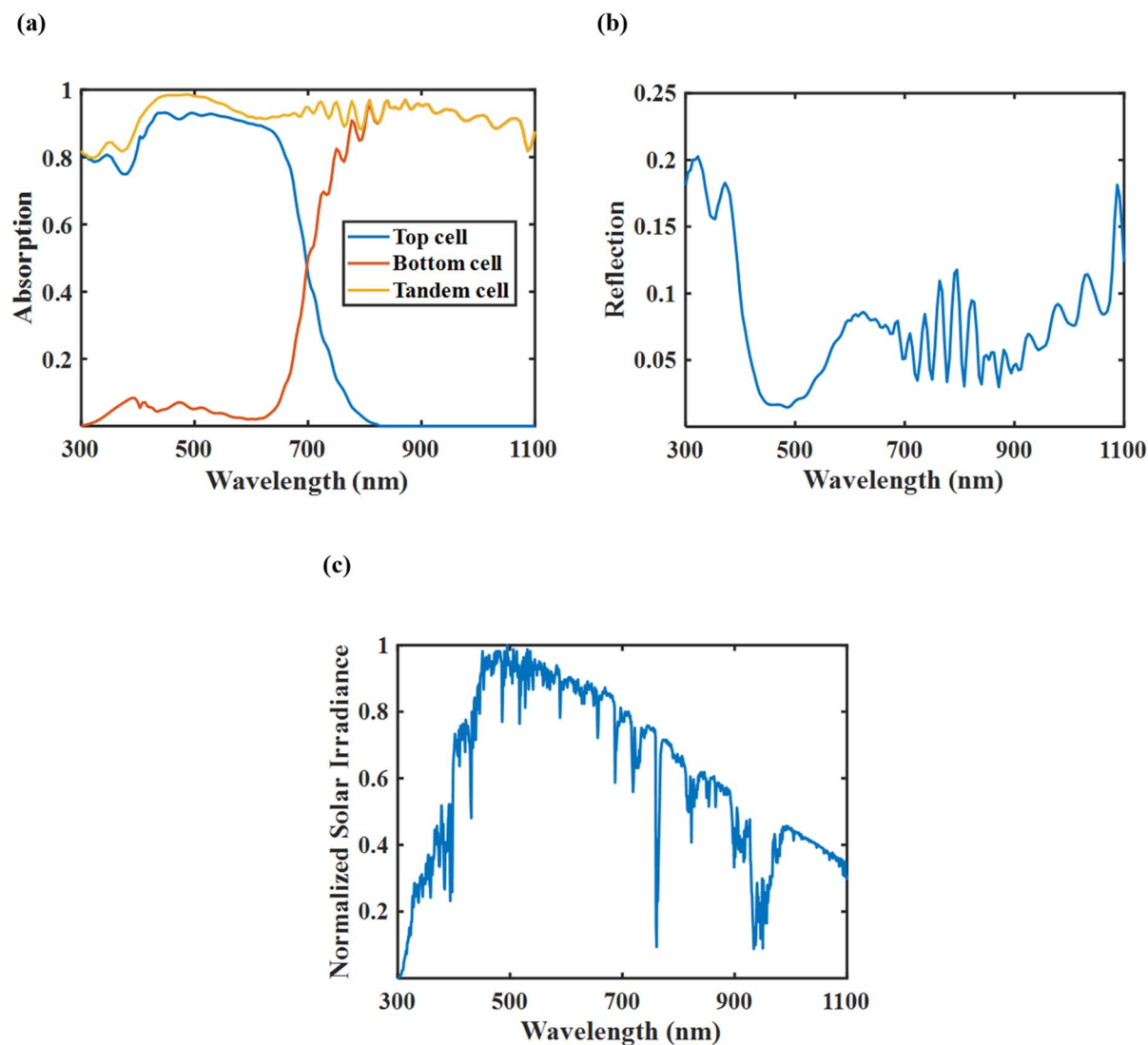


Fig. 9 (a) Normalized absorption spectra of the top, bottom, and tandem cells, (b) normalized reflection spectra of the tandem cell, and (c) normalized AM 1.5G solar spectrum. The figures indicate that the top and bottom cells complement each other effectively, enabling the tandem configuration to achieve broad and enhanced absorption across the entire spectral range.

a favourable balance between V_{oc} and J_{sc} , thereby, enhancing the efficiency. In comparison, while the AlGaAs/Si tandem cell has a higher J_{sc} ,⁹¹ it demonstrates a significantly lower V_{oc} , resulting in a decreased PCE. Some of the perovskite/Si tandem

cells shown in Table 5 exhibit V_{oc} levels comparable to those reported in this study.^{46,48} However, their J_{sc} values are lacking, negatively affecting their overall performance. Ugur *et al.* fabricated a perovskite/Si tandem solar cell with a certified PCE of 33.7% that employs a lead-based organic–inorganic hybrid perovskite, surpassing the SQ limit.⁹⁰ In contrast, the present study demonstrates superior PV performance, including PCE, V_{oc} , J_{sc} , and FF, while employing a lead-free inorganic perovskite absorber.

Compared to earlier tandem designs, the CsSnGeI₃/SHJ architecture shows significant benefits. Although lead-based hybrid perovskite/Si tandems have achieved high efficiencies,^{26,46,48} their toxicity and environmental instability limit commercial viability. Conversely, fully inorganic, lead-free perovskites like MAgE₃,²⁶ CsSnI₃,⁴² and Bi or Cu-based options^{28,36} often face challenges such as lower carrier

Table 5 Comparison of characteristic metrics with other Si-based tandem cells

Cell type	PCE (%)	V_{oc} (V)	J_{sc} (mA cm ⁻²)	FF (%)
This work	34.93	1.93	21.30	84.74
Perovskite/Si tandem cell ⁹⁰	33.7	1.985	21.02	81.6
Perovskite/Si tandem cell ⁴⁶	32.2	1.87	20.3	84.7
Perovskite/Si tandem cell ²⁶	23.8	1.76	19.2	70
Perovskite/Si tandem cell ⁴⁸	28.2	1.88	19.6	78.6
Kesterite/Si tandem cell ⁸³	28.28	1.662	19.93	85.38
AlGaAs/Si tandem cell ⁹¹	25.2	1.55	27.90	58



mobility, phase instability, or shorter diffusion lengths, resulting in modest tandem PCEs ($\sim 30\text{--}32\%$). CsSnGeI_3 , however, features a suitable bandgap, better film quality, and increased environmental stability thanks to a native oxide layer.^{40,43} Its higher carrier mobility enhances charge transport and boosts current generation. The SHJ bottom cell enhances the top absorber by providing excellent surface passivation, high V_{oc} , and compatibility with well-established manufacturing techniques. This makes SHJ a more dependable choice compared to traditional homojunction or Passivated Emitter Rear Locally diffused (PERL) Si structures, or even less efficient bottom cells like CIGS or CZTS.^{6,88} To confirm the combined effectiveness of both absorbers, we simulated each subcell separately, achieving efficiencies of 23.46% and 12.13%, as shown in Table 4. When combined in the optimized tandem design, the overall PCE increases significantly to 34.93%, demonstrating that the tandem configuration and material synergy are crucial for exceeding the SQ limit. This highlights the vital contributions of the CsSnGeI_3 absorber and the SHJ structure in attaining high efficiency while maintaining environmental sustainability.

Recent advancements in perovskite/silicon tandem photovoltaics have demonstrated that achieving efficiencies in the mid-30% range is now possible both at the industrial and laboratory scale. For example, LONGi achieved a certified 34.85% efficiency for a two-terminal perovskite/Si tandem cell, and research institutions like Helmholtz-Zentrum Berlin and King Abdullah University of Science and Technology have reported laboratory efficiencies between 32.5% and 33.7%.^{21,92} The simulated efficiency of 34.93% achieved in this study marginally exceeds previously reported state-of-the-art values, demonstrating the strong performance potential of a fully inorganic, lead-free tandem configuration and underscoring its significance for future high-efficiency photovoltaic technologies.

3.7. Impact of plasmonic nanorods

In order to assess the influence of plasmonic nanorods on device performance, simulations were executed utilizing the fully optimized tandem structure. For the purpose of comparative analysis, the cylindrical Au nanorods, along with their surrounding Si_3N_4 dielectric medium, were removed to model the structure without plasmonic enhancement. This approach facilitated a direct evaluation of the nanorods' impact on the device's optical behavior.

Fig. 10 illustrates that the presence of plasmonic nanorods leads to a marginal enhancement in absorption and a slight decrease in reflection, especially at higher wavelengths. However, the overall spectral response remained consistently high across the visible spectrum in both situations, confirming that the tandem cell demonstrates strong light-harvesting capability even without plasmonic enhancement. This implies that while the nanorods provide additional light-trapping advantages, they are not mandatory for achieving high efficiency in the optimized configuration.

Table 6 shows that the PCE of the tandem structure drops from 34.93% to 34.32% when both the cylindrical Au nanorods and the dielectric layer on the rear side of the bottom cell are removed. This decrease is primarily attributed to a slight reduction in the J_{sc} , as the V_{oc} remains nearly constant despite the removal of the plasmonic back reflector. This finding differs from the research findings by Jamil *et al.*,⁸³ where the elimination of the plasmonic back reflector resulted in a PCE drop of more than 7%. In contrast, this study reports a decline of only 0.61%, suggesting that the structure could still exceed the SQ limit, even without the plasmonic back reflector. This is accomplished by utilizing an appropriate perovskite absorber in the top cell and incorporating an additional ultra-thin GaSb absorber layer within the bottom SHJ cell. Thus, if minimizing complexity and production costs is a priority, the plasmonic

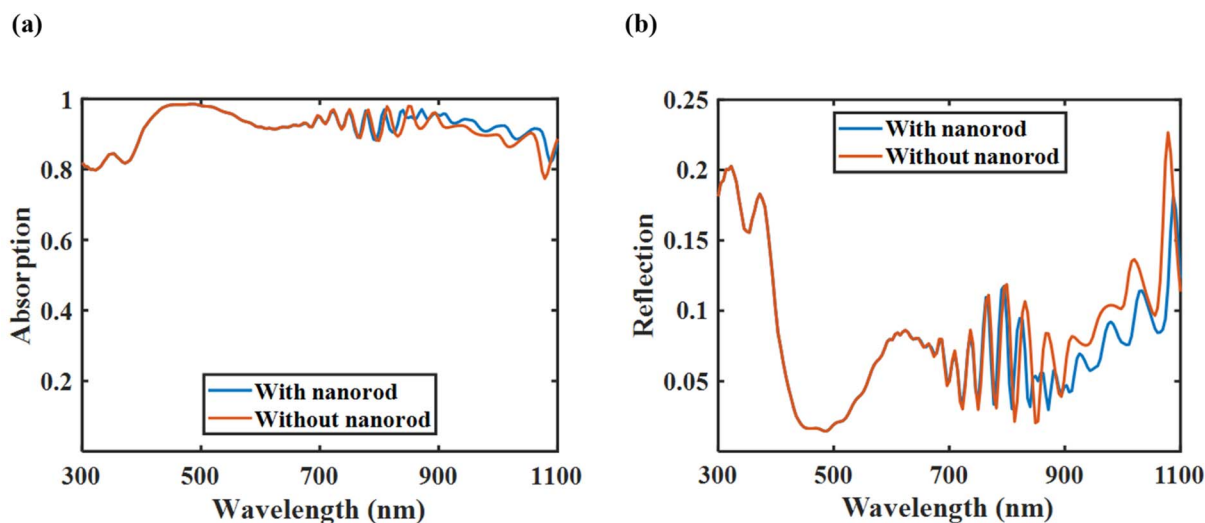


Fig. 10 (a) Normalized absorption spectra of the tandem cell with and without nanorods, and (b) normalized reflection spectra of the tandem cell with and without nanorods. The incorporation of nanorods resulted in a slight increase in optical absorption, indicating their contribution to improved light harvesting.

Table 6 Characteristic metrics of tandem structure with and without a plasmonic back reflector

Cell type	PCE (%)	V_{oc} (V)	J_{sc} (mA cm ⁻²)	FF (%)
With a plasmonic back reflector	34.93	1.9347	21.30	84.74
Without a plasmonic back reflector	34.32	1.9334	21.18	83.79

back reflector and the dielectric layer can be omitted from the design while maintaining a high efficiency.

3.8. Limitations and future outlook

This study offers a detailed numerical simulation of a lead-free perovskite/SHJ tandem solar cell with plasmonic enhancement. However, it is essential to note that the findings are based solely on FDTD and drift-diffusion models. While the study carefully considered realistic material properties, interfacial recombination (through surface recombination velocity), and fabrication practicality, no experimental validation has been conducted yet. Furthermore, issues such as the quality of interface passivation, long-term material stability, and challenges in layer deposition may influence actual performance. Future efforts will involve collaborating with experimental groups to develop and confirm the proposed design experimentally.

4. Conclusion

In summary, this study presents a fully inorganic and lead-free CsSnGeI₃/SHJ tandem solar cell that is simulated to achieve a remarkable PCE of 34.93%, surpassing the SQ limit for single-junction cells. The integration of CsSnGeI₃, a non-toxic and inorganic perovskite, directly addresses the pressing environmental concerns associated with traditional lead-based counterparts. The inclusion of a GaSb auxiliary absorber within the SHJ bottom cell enhances optical absorption, particularly in the NIR region, allowing for the utilization of a thinner c-Si layer without compromising performance. Performance is additionally enhanced by incorporating cylindrical Au nanorods within a Si₃N₄ dielectric medium positioned at the rear of the bottom cell, thus amplifying light absorption through plasmonic effects. Notably, the structure retains high efficiency even in the absence of the plasmonic back reflector, providing flexibility in fabrication and cost reduction. The design illustrates a pragmatic trade-off between complexity and performance, paving the way for low-toxicity and cost-effective photovoltaic technologies. These findings are based on comprehensive numerical modeling and await experimental validation.

Opportunities for future work include enhancing this architecture through the application of surface texturing techniques, particularly to the top cell, to further improve light trapping and reduce reflection as typically observed in PERC cells. Incorporating additional junctions with suitably engineered bandgap materials may enable multi-junction configurations that capture a broader portion of the solar spectrum, thereby further enhancing the efficiency. These avenues, coupled with ongoing advancements in lead-free materials,

establish the groundwork for next-generation high-efficiency tandem PV cells.

Conflicts of interest

The authors have no conflict of interest to declare.

Data availability

Data that supports the findings of this work is available from the corresponding author upon reasonable request.

Supplementary information is available. See DOI: <https://doi.org/10.1039/d5ra05323d>.

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