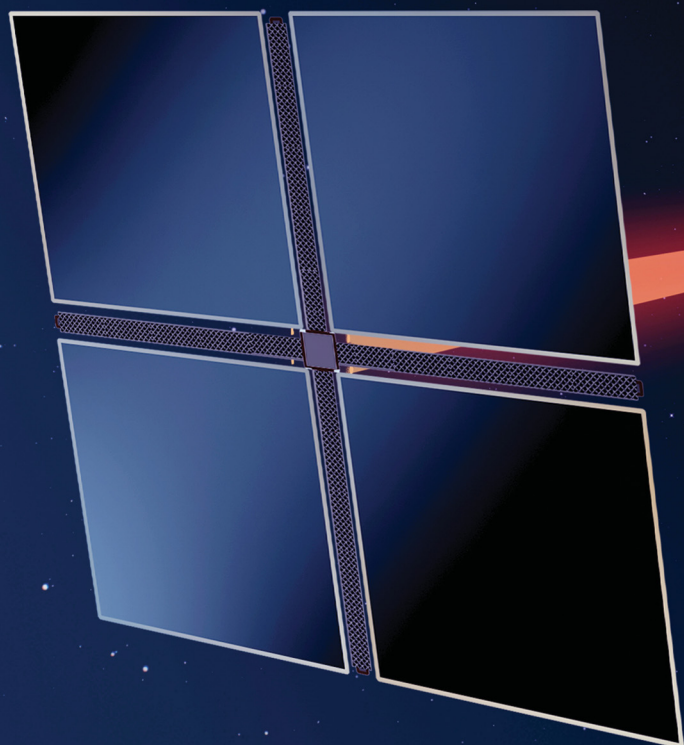


Energy Advances

Volume 2
Number 2
February 2023
Pages 229–348

rsc.li/energy-advances



ISSN 2753-1457

PAPER

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Cite this: *Energy Adv.*, 2023,
2, 298

Received 16th August 2022,
Accepted 1st December 2022

DOI: 10.1039/d2ya00218c

rsc.li/energy-advances

Low-intensity low-temperature analysis of perovskite solar cells for deep space applications†

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High specific power (power per mass) ultralight solar arrays made of perovskite solar cells (PSCs) are being considered to power spacecraft in deep space conditions as far as Neptune (30 AU). To understand how PSCs perform and respond in deep space, we characterize PSCs under low-intensity low-temperature (LILT) conditions before and after low-energy proton irradiation. PSCs show promising performance characteristics under most LILT conditions even after exposure to low-energy protons. However, the measured cell efficiency tends to decrease at extreme lower temperatures, suggesting that further research into cell architectures and materials could improve PSCs for deep space applications.

Introduction

Solar cells are an important power source for enabling non-nuclear exploration of the solar system. Solar cells supply constant, renewable power to a space mission vehicle and its payload when in sight of the sun, allowing for sustained travel to the outer reaches of the solar system. State-of-the-art (SOA) space solar arrays use triple-junction solar cells based on III–V compounds and germanium. However, these cells significantly degrade with radiation exposure, requiring heavy coverglass for shielding. Since perovskites were first identified as a viable photovoltaic material, considerable research efforts have been made regarding perovskite solar cells (PSCs) for space applications.^{1–12} Preliminary studies suggest that emerging ultrathin, flexible, and lightweight perovskite solar cells are naturally radiation hardened,^{11–17} potentially enabling high specific power¹⁸ solar arrays to be designed for power generation in high radiation and deep space environments. Additionally, the low cost of required materials means that PSCs are cost-effective.¹⁹

Here, we present two different PSC structures and demonstrate experimental current density–voltage (*J*–*V*) and external

quantum efficiency measurements (EQE) of PSCs operating in low-intensity low-temperature (LILT) conditions mimicking a deep space environment. The temperatures are as low as $-170\text{ }^{\circ}\text{C}$ and light intensities as low as 1.52 W m^{-2} , which corresponds to solar irradiance at Neptune, 30 astronomical units (AU) from the Sun. In this study, we label the light intensity of a given measurement based on distance from the Sun in AU, since solar intensity drops off as the square of distance. 1.0 AU is the distance between the Earth and the Sun and thus corresponds to the Air Mass Zero (AM0) spectrum.

Previous LILT studies^{1,2} looking at different PSC device structures have observed similar performance trends with respect to temperature and light intensity as we report here. Brown *et al.* demonstrated temperature-dependent EQE and LILT *J*–*V* measurements in Mars, Jupiter, and Saturn environments for a triple-cation PSC device fabricated using an n–i–p architecture, meaning that the cell is illuminated from the electron transport layer (ETL) side.¹ They reported temperature-dependent EQE and photoluminescence (PL) measurements that show an increase in bandgap energy with increasing temperature for perovskites and the absence of low-temperature phase transitions in the triple-cation PSC.¹ In our study, we observe these same effects in a double-cation PSC device fabricated using a p–i–n architecture, meaning that the cell is illuminated from the hole transport layer (HTL) side. Furthermore, Brown *et al.* presented *J*–*V* curves for PSCs operating in Earth, Mars, Jupiter, and Saturn conditions, which demonstrate that the low-temperature device performance improves at lower light intensities, ideal for deep space applications.¹ Our study confirms these trends for two different PSC device architectures, while extending the LILT

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† Electronic supplementary information (ESI) available. See DOI: <https://doi.org/10.1039/d2ya00218c>

condition range to light intensities matching Uranus (19.2 AU) and Neptune (30.0 AU). Sun *et al.* performed a LILT study on a triple-cation PSC device fabricated using a p-i-n architecture, which looked at simulated Martian surface conditions.² Their work showed long-term stability of PSCs operating in Martian surface conditions and noted that device performance was actually improved slightly after a simulated Martian day.² Sun *et al.* also confirmed that low-temperature induced phase transitions could contribute to the self-elimination of intrinsic defect states, improving the device efficiency.^{2,7}

Both studies by Brown *et al.* and Sun *et al.* state the promise of PSCs for space applications, but continued research is required before PSCs become a viable technology. To the best of our knowledge, our study presents the most comprehensive LILT condition study thus far, testing two different PSC architectures under a combination of five different light intensities and six different temperatures. Due to the large size of our data set, we can present cell efficiency charts that demonstrate clear trends with respect to light intensity and temperature, which highlight areas for future improvement of PSCs for deep space applications.

Furthermore, we complement our LILT study by considering the effect of low-energy proton radiation. While the radiation hardness of PSCs has been heavily documented,^{11–17} recent studies have emphasized that low-energy protons are the most appropriate radiation source for creating atomic vacancies and probing radiation effects in perovskites representative of space conditions.^{4,16} Higher energy particles, which many previous PSC tests have used due to their use in radiation tests for III–V and silicon cells,²⁰ could actually create localized heating and make tests less reliable.⁴ Thus, Kirmani *et al.* call for new low-energy proton radiation tests, such as the ones performed in this study. We do observe some degradation in device performance after radiation, but analysis of a champion cell shows that it is possible for a PSC to retain a substantial portion of its pre-radiation efficiency compared to SOA space solar cells.

PSCs have the potential to outperform SOA space solar cells in two key metrics that are important for space applications: radiation resistance and specific power. However, it remains unclear whether this would be true in extreme LILT conditions because very low temperatures at distances like Uranus (19.2 AU) are difficult to measure experimentally due to temperatures lower than what can be achieved using liquid nitrogen. Thus, while initial measurements of PSC performance in less severe LILT conditions look promising, our measurements also suggest that performance would decrease at very low temperatures, and improvements to cell architectures and materials would likely be needed to improve PSC performance.

Results and discussion

Specific power of a perovskite solar array

The specific power of a solar cell is especially important for space applications where the weight of payloads is a serious consideration. Gdoutos *et al.*²¹ from Caltech have developed a

lightweight 1.7 m × 1.7 m prototype solar array structure with areal density of 150 g m^{−2}. This array structure is designed to be scalable up to 60 m × 60 m. At this size, it is estimated to have an areal density of 50 g m^{−2} because areal density decreases for larger structures.²¹ Because perovskite solar cells are a thin-film technology that can be deposited *via* low-temperature processes, they can be fabricated on a range of thin, lightweight, and flexible substrates or foils. At the cell level (*i.e.*, not encapsulated or integrated into a module), flexible perovskite solar cells have been demonstrated with specific power values as high as 26 kW kg^{−1}, compared to values of <2.4 kW kg^{−1} for current high-efficiency multi-junction cells used in most space applications.^{22,23} Fig. 1 presents the mass of different solar array combinations using either SOA space solar cells or perovskite solar cells. We calculate the areal density if these cells were to be mounted on a standard structure²⁴ or on an ultralight structure currently being developed at Caltech.²¹ Standard SOA solar cells on a standard structure have an areal density of 2.13 kg m^{−2}, whereas PSCs on the ultralight Caltech structure would have an areal density of 0.25 kg m^{−2}.^{21,22} The drastic reduction in areal density enables a high specific power for the PSCs mounted on the Caltech structure. Large ultralight and high specific power solar arrays could enable a new class of deep space mission and a non-nuclear exploration of the solar system.

Perovskite solar cell structure

Perovskites are a group of compounds that have the same ABX₃ crystal structure as the mineral perovskite (calcium titanate – CaTiO₃). In a typical perovskite solar cell, the A site has a large cation such as methylammonium (MA), formamidinium (FA), cesium (Cs), rubidium (Rb), or a combination of these. The B site is a smaller cation such as lead (Pb) or tin (Sn). The X site balances the charge with one or more of the halides iodine (I), bromine (Br), or chlorine (Cl).¹⁹ Since 2012, when the first solid state heterojunction methylammonium lead iodide (MAPbI₃) perovskite cell was introduced,²⁵ perovskite solar cell technology has been one of the fastest growing solar technologies, with efficiencies now reaching 25%.²⁶

Fig. 2(a) and (b) show the perovskite solar cell structure for cells from The Australian National University (ANU) and Caelux

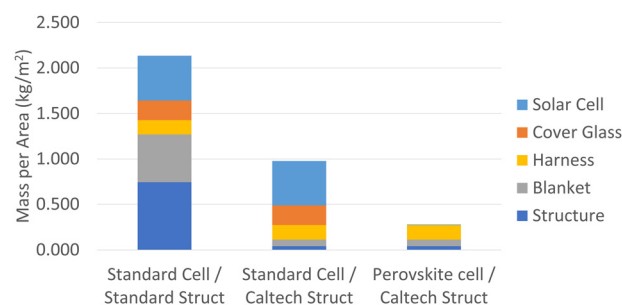


Fig. 1 Comparison of the mass per area of current state-of-the-art solar cells with perovskite solar cells. The perovskite solar cell structure assumes no coverglass (due to the radiation hardness of perovskites) and a value of 0.0056 kg m^{−2} for the PSC, which is too small of a contribution to be visible in the figure.



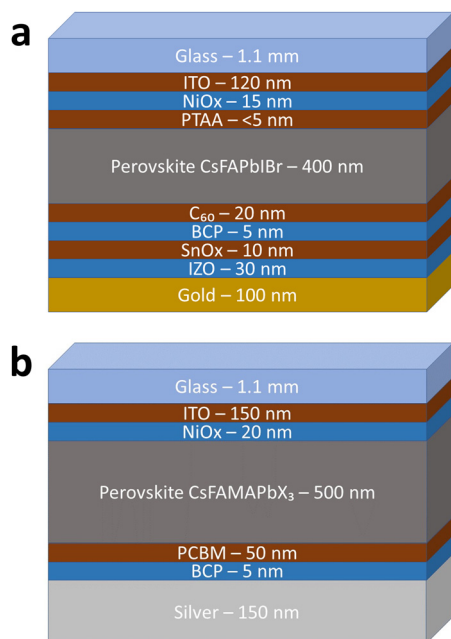


Fig. 2 (a) ANU perovskite solar cell structure. (b) Caelux perovskite solar cell structure.

Corporation, respectively. The ANU PSCs had an active area of 1 cm^2 and utilized a double-cation, mixed halide perovskite with composition $\text{Cs}_{0.22}\text{FA}_{0.78}\text{PbI}_{2.55}\text{Br}_{0.45}$ for the active layer. The Caelux PSCs had an active area of 0.13 cm^2 and utilized a similar triple-cation perovskite with composition $\text{Cs}_{0.09}\text{FA}_{0.8}\text{MA}_{0.11}\text{PbI}_{2.85}\text{Br}_{0.15}$ for the active layer. In both cells, the top layer of glass serves as a rigid superstrate during the fabrication process and provides some physical and environmental protection. However, preliminary studies on perovskite solar cells show radiation tolerance surpassing other known photovoltaic materials, even when radiated from the rear (non-glass) side.^{11–17} While further studies are required to fully understand and verify these results, they suggest that perovskite cells may not require a heavy coverglass for radiation protection, further reducing mass and volume, assuming that perovskite solar cells have sufficient mechanical stability without glass encapsulation, which is an active research topic.²⁷ Thus, in our specific power calculations, we omit the structural glass layer and assume that the same cell structure could be deposited on an ultra-lightweight polyimide film or foil.²³

Low-intensity low-temperature (LILT) measurement conditions

To simulate the solar cell operation in deep space, we tested PSCs at various mission-relevant temperatures ranging from 28°C to -170°C and light intensities based on the solar irradiance from 1.0 AU to 30.0 AU. Table 1 details the solar light intensities, temperatures, and radiation dose rates at the distances of Earth and the outer planets. Solar light intensity is inversely proportional to the square of distance and is calculated by dividing the value of the solar constant, 1367 W m^{-2} , by the square of the distance (in AU). Approximate solar array operating temperatures at these distances in space are

Table 1 Relationship between distance, solar intensity, and temperature at five different planets in the solar system. The lowest temperature that can be tested in the liquid nitrogen (LN_2)-cooled photovoltaic lab setup at JPL is -170°C

Planet	Distance (AU)	Solar intensity (W m^{-2})	Temperature ($^\circ\text{C}$)	Radiation dose rate (rad s^{-1})
Earth	1.0	1367	60	3×10^{-4}
Jupiter	5.5	45.2	-140	8×10^{-1}
Saturn	9.5	15.2	-165	6×10^{-5}
Uranus	19.2	3.71	-200	8×10^{-4}
Neptune	30.0	1.52	-230	3×10^{-4}

calculated by balancing heat flow through a solar array in thermal equilibrium using the Stefan-Boltzmann law. In determining heat flow, we assume the same solar absorptivity and emissivity for SOA cells and PSCs,²⁸ yielding a close approximation to commonly used values.^{1,29,30} In actual space missions, these temperature values could vary slightly if the cell absorptivity, emissivity, or efficiency differs or if the solar intensity changes due to a particular mission path or distance from the sun. Therefore, we use the values in Table 1 as an approximate flight envelope. Lastly, the radiation dose rates are calculated at $L = 7$ of each planet under 100 mil aluminum shielding.³¹ $L = 7$ means 7 times a planet's radius from the planet's center on the equatorial plane and line with the same magnetic field at different magnetic latitudes.

Low-intensity low-temperature (LILT) current density–voltage (J – V) measurements

We first analyze LILT J – V measurements taken on PSCs from ANU and Caelux. These measurements were taken using an X-25 solar simulator at temperatures and light intensities determined by Table 1. The solar simulator was calibrated using InGaP and GaAs calibration standards. To properly measure the PSC test cells, a spectral mismatch correction factor³² was calculated for our experimental setup as:

$$M = \frac{\int_{\lambda_1}^{\lambda_2} E_{\text{AM0}}(\lambda) S_{\text{Ref}}(\lambda) d\lambda}{\int_{\lambda_1}^{\lambda_2} E_{\text{AM0}}(\lambda) S_{\text{TestCell}}(\lambda) d\lambda} \frac{\int_{\lambda_1}^{\lambda_2} E_{\text{Lamp}}(\lambda) S_{\text{TestCell}}(\lambda) d\lambda}{\int_{\lambda_1}^{\lambda_2} E_{\text{Lamp}}(\lambda) S_{\text{Ref}}(\lambda) d\lambda}$$

$$= \frac{195.4}{274.6} \frac{291.2}{249.7} = 0.83,$$

and applied to the PSC experimental data by dividing the measured PSC test cell current by M .

Fig. 3 shows the J – V curves of a selected ANU PSC labeled ANU9 and a selected Caelux PSC labeled CA3 at 1.0 AU and 19.2 AU. As intensity decreases from 1.0 AU to 19.2 AU, the measured current decreases, which follows from the decrease in the number of incident photons. The J – V curves of better performing cells have a higher fill factor, the ratio of the maximum power to the product of the open-circuit voltage and short-circuit current. In Fig. 3, we visually observe that fill factor tends to decrease as temperature decreases, although this trend is not necessarily monotonic at temperatures $> -100^\circ\text{C}$. For the ANU cell at 1.0 AU in Fig. 3(a), the highest fill factor is achieved at 28°C with a value of 0.50. As temperature decreases to -170°C , the fill factor decreases monotonically to 0.164. However, for the ANU cell at



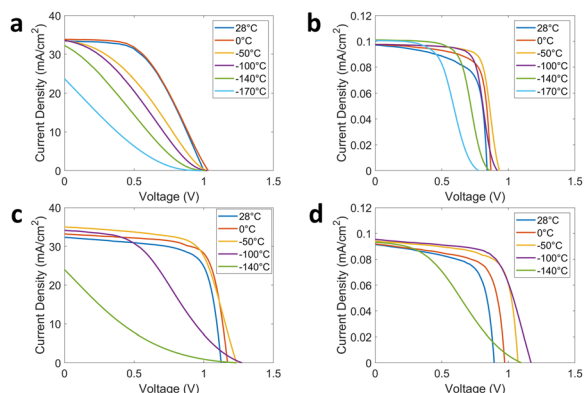


Fig. 3 Pre-radiation LILT J - V curves for (a) ANU9 cell at 1.0 AU, (b) ANU9 cell at 19.2 AU, (c) Caelux CA3 cell at 1.0 AU, and (d) Caelux CA3 cell at 19.2 AU.

19.2 AU in Fig. 3(b), the highest fill factor is achieved in the middle of the temperature range at $-50\text{ }^{\circ}\text{C}$ with a value of 0.75. As temperature increases to $28\text{ }^{\circ}\text{C}$ or decreases to $-170\text{ }^{\circ}\text{C}$, we observe a decrease in fill factors to values of 0.67 and 0.53, respectively. For the Caelux cell at 1.0 AU in Fig. 3(c), the highest fill factor is achieved at $0\text{ }^{\circ}\text{C}$ with a value of 0.69, while at temperatures of $-100\text{ }^{\circ}\text{C}$ and $-140\text{ }^{\circ}\text{C}$, we observe decreasing fill factors of 0.38 and 0.14, respectively. For the Caelux cell at 19.2 AU in Fig. 3(d), the highest fill factor is achieved at $-50\text{ }^{\circ}\text{C}$ with a value of 0.68, and again we see a decrease in fill factor at $-100\text{ }^{\circ}\text{C}$ and $-140\text{ }^{\circ}\text{C}$ with values of 0.65 and 0.34, respectively.

To further explore the effects of temperature and incident light intensity, we look at the average cell efficiency for the PSCs as a function of temperature at each intensity, shown in Fig. 4. We observe a non-monotonic temperature dependence, which has also been observed for a NiOx/triple-cation PSC device in a recent study of the temperature coefficients of PSCs.³³ Notably, Moot *et al.* found that these temperature-dependent trends were different for four different PSC device structures,³³ suggesting that the ideal PSC device architecture may differ depending on the specific LILT conditions. Importantly, Moot *et al.* also highlight reversible changes in device efficiency with respect to temperature, ruling out degradation as a mechanism for performance change.³³ To consider the effects of temperature-induced or light-induced degradation in our study, we performed the same $28\text{ }^{\circ}\text{C}$ room temperature measurement before and after subjecting the PSCs to LILT conditions. As shown in Fig. S6–S8 (ESI[†]), no signs of degradation are present, suggesting that the observed trends are indeed reversible. Furthermore, we observe that as light intensity decreases, the temperature at which maximum efficiency is measured decreases. This leads to a trend at lower temperatures, where the device efficiency improves as light intensity decreases. This is attributed to a more efficient extraction rate for the minority electrons under illumination, as shown previously for a different PSC structure by Brown *et al.*¹ We confirm these previous results from Brown *et al.* and Moot *et al.* and further explain the trends by considering the effects of non-radiative recombination, series resistance, and material conductivity.

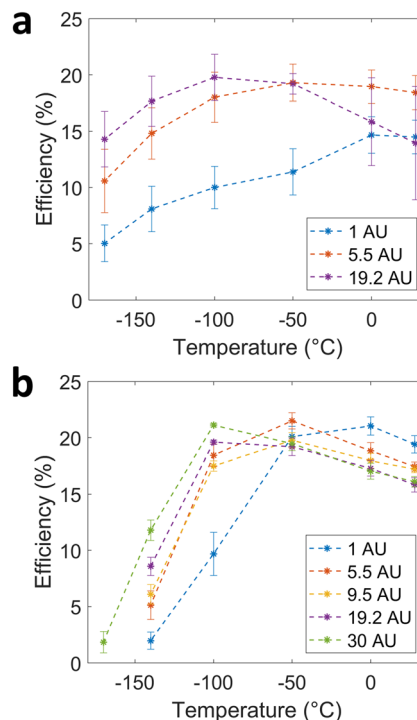
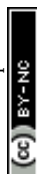


Fig. 4 Plots of average efficiency with respect to temperature at various light intensities for (a) ANU perovskite solar cells and (b) Caelux perovskite solar cells, both before radiation. The error bars denote the standard deviation in the set of measurements ($n = 3$ cells in each case).

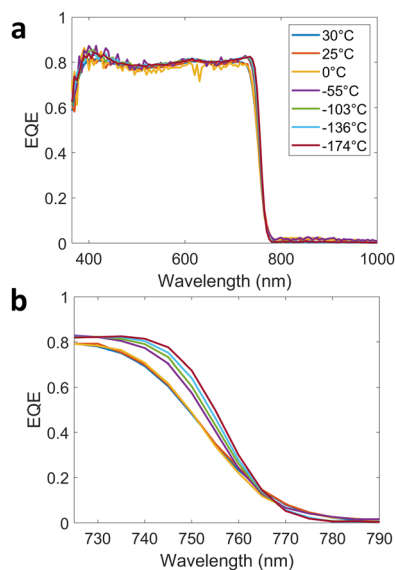
Non-radiative recombination is an unwanted process in solar cells because it reduces the charge carrier density in the solar cell, impacting the voltage, and to a lesser extent, the current.³⁴ Typically, the effects of non-radiative recombination are lessened as temperature decreases³⁵ which could explain the increase in efficiency as temperature drops from $28\text{ }^{\circ}\text{C}$ to $-50\text{ }^{\circ}\text{C}$. The initial increase in open-circuit voltage with a decrease in temperature (which is particularly obvious for the Caelux cells in Fig. 3(c) and (d)) is the result of a reduction in non-radiative recombination *via* thermally activated traps, and thus an increase in charge carrier density in the active layer of the cell. This is also observed in steady-state PL measurements of perovskite films of the type used in the ANU cells (Fig. S1, ESI[†]), which show a monotonic increase in emission intensity with decreasing temperature. We also observe a small red-shift in the PL emission peak. Similar temperature-dependent trends in emission intensity and wavelength have also been reported by others.^{1,7,36}

Fig. 4 also demonstrates increased efficiencies at lower light intensities, which can likely be attributed to the effect of series resistance. Series resistance losses can come from anywhere in the cell and they cause a voltage loss proportional to the current. Thus, power losses due to series resistance increase as light intensity increases because of the higher maximum power point current. Fig. 3(a) shows a J - V curve indicative of the presence of series resistance, leading to a significant voltage loss in the cell at the maximum power point, which explains the poor fill factor at higher intensities. But, towards 19.2 AU, the



The effects of non-radiative recombination and series resistance together create an optimal temperature at which the device efficiency is a maximum. At 1.0 AU in Fig. 4(b), series resistance at lower temperatures plays a large role, so the maximum efficiency occurs at a relatively high temperature. At 9.5 AU, resistive losses are proportionally smaller, so cell performance initially benefits from reduced non-radiative recombination at lower temperature; hence, cell performance peaks at a lower temperature compared to 1.0 AU. As the light intensity is further reduced to 19.2 AU and 30 AU, series resistance losses become very low, so the optimum efficiency temperature drops further. However, we still observe a drop in

We look at temperature-dependent EQE measurements of the PSCs. One ANU PSC was tested at various temperatures ranging from 30 °C down to approximately -170 °C, and the resulting EQE curves are shown in Fig. 5(a). The EQE plateaus around 0.8 and the bandgap is shown to be around 760 nm or 1.63 eV. We observe little change due to temperature, which is unlike



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Energy Adv., 2023, 2, 298–307 | 303

performing radiation tests on the same cells to contribute to the limited literature on low-energy proton radiation tests for PSCs.^{4,16} With the resulting large data set of measured PSC efficiencies, we predict how the ANU PSCs would perform relative to SOA cells in the context of a space mission to Jupiter (5.5 AU, -140°C from Table 1). Measured without the effects of radiation, SOA cells have demonstrated efficiencies exceeding 35% at 5.5 AU and -140°C .⁴⁹ With this efficiency, a standard²⁴ SOA space solar array has an estimated specific power of 6.5 W kg^{-1} . This value could be increased using the ultralight Caltech structure,²¹ yielding a projected specific power of 12 W kg^{-1} . We compare these values to the projected specific power of ANU PSCs mounted without coverglass on an ultralight Caltech structure,²¹ using both pre-radiation and post-radiation efficiencies. Pre-radiation efficiencies are an average of 3 measured cells, as shown in Fig. 4(a), and post-radiation efficiencies are from the champion cell as shown in Fig. 6(b). Then, at 5.5 AU and -140°C , the projected ultralight PSC array specific power is around 24 W kg^{-1} before radiation and 7.3 W kg^{-1} after radiation. The projected ultralight PSC array specific power clearly exceeds estimated standard SOA space solar array values. However, if the SOA cells were to be similarly mounted on an ultralight Caltech structure, then the PSC array does not have a clear advantage, particularly when comparing with the post-radiation PSC value. Although the SOA values are assuming no radiation, the SOA cells typically employ a protective coverglass layer to prevent severe radiation degradation. Thus, because the SOA values assume coverglass and the PSC values do not assume coverglass, we compare the beginning-of-life SOA value (6.5 W kg^{-1} , 12 W kg^{-1} with ultralight structure) and the post-radiation PSC value (7.3 W kg^{-1}). From this comparison, PSCs show promise to enable higher specific power arrays than currently available. However, if improvements were made to SOA cell arrays that would likewise enable the SOA cells to be mounted on ultralight structures, the PSCs tested here would not offer a conclusive advantage over SOA cells, especially if we consider more extreme LILT conditions where the PSC performance degraded.

We cannot make a direct comparison at more extreme LILT conditions such as Uranus (19.2 AU, -200°C) because this requires a temperature of -200°C , and the lowest temperature that can be tested in the LN_2 -cooled photovoltaic setup at JPL is -170°C . However, both Fig. 4 and 6(b) suggest that efficiencies would likely decrease at lower temperatures like -200°C , and thus PSCs may not outperform SOA cells. Because these low temperatures are not currently achievable, further research must be done to make the comparison between PSCs and SOA cells at these extreme conditions. Regardless, the measured PSCs suffer significant reduction in cell efficiency at extreme low temperatures, suggesting that cell architectures and materials may have to be optimized for LILT conditions.

A comparison of the temperature and illumination dependent performance of the ANU and Caelux cells in Fig. 3 and 4 reveals significant differences between the two cell types which may provide guidance on future optimization for LILT conditions. Apart from the active areas (1 cm^2 versus 0.13 cm^2), which could

explain the relatively high series resistance, and hence low fill factor of the ANU cells under 1.0 AU illumination, other notable differences between the cells are the perovskite composition and the rear electron transport layer stack. It is likely that both may contribute to the differences in LILT performance, but further LILT studies that compare different types of PSC architectures and material choices are required to identify the specific contributions of each layer and the optimal material combinations. In this context, we propose that future studies and optimization for LILT performance should focus on three distinct aspects of the cell design: (i) tuning of the perovskite composition to take advantage of low-temperature phase transitions that can improve material quality;⁷ (ii) selecting charge transport materials with a high carrier concentration and/or intentionally increasing the doping level so a reasonable conductivity can be maintained at low temperatures, and (iii) optimizing the device structure and fabrication to maximize shunt resistance, as low-light performance is severely impacted by low shunt resistance.⁵⁰ To support this, detailed characterization of the fundamental semiconducting properties (e.g. bandgap, energy levels, mobility, carrier density, etc.) of different perovskite compositions and charge transport layers at deep-space temperatures below -170°C is required, as much of this information is currently unavailable. Then, on the device level, along with light J - V measurements and EQE measurements such as the ones in this study, other solar cell characterization techniques such as dark J - V measurements or power-dependent measurements could be employed to further investigate device performance in LILT and radiation conditions. Lastly, *in situ* characterization of PSCs in radiation and LILT conditions simultaneously would help to eliminate uncertainty introduced by the handling of devices and delay between irradiation and characterization of devices. While much work has gone into researching PSCs for space applications, continued research that compares specific materials and device structures³³ in full LILT and radiation conditions is vital to enabling viable PSCs for deep space missions.

Experimental

ANU cell fabrication

The indium tin oxide (ITO) substrates were sequentially cleaned in detergent, acetone, isopropyl alcohol and ethyl alcohol for 20 min in an ultrasonic bath, and then dried with nitrogen. Cleaned ITO substrates were exposed to approximately 30 min of UV-O₃ treatment before the subsequent deposition step. $\sim 15\text{ nm}$ nickel oxide thin films were deposited on the pre-cleaned ITO substrates by radiofrequency magnetron sputtering at 150 W and 18 sccm Ar and 2 sccm O₂ gas using a nickel oxide target under a pressure of 1.5 mTorr. The pre-fabricated nickel oxide thin films were then transferred to a furnace, and annealed 30 min under a controlled oxygen atmosphere. For the $\sim 5\text{ nm}$ PTAA (a poly(triaryl amine) semiconductor) thin film deposition, $\sim 50\text{ uL}$ PTAA solution (2 mg ml^{-1} in chlorobenzene) was dropped onto the ITO/NiOx substrates and deposited by spin-coating at 4000 rpm for 20 s. The perovskite film ($\text{Cs}_{0.22}\text{FA}_{0.78}\text{Pb}(\text{Cl}_{0.03}\text{Br}_{0.15}\text{I}_{0.85})_3$, 1.2 mmol



(68.6 mg CsI, 99.1 mg PbBr₂, 161 mg FAI, 429 mg PbI₂) in 0.8 ml dimethylformamide (DMF) and 0.2 ml dimethyl sulfoxide (DMSO) with 5 mol% MACl and 3 mol% PbCl₂ additive) was deposited by one-step spin-coating at 3000 rpm with a ramp of 100 rpm s⁻¹ for 12 s. After spinning, the substrate was transferred immediately into a vacuum chamber and pumped down to 120 mTorr for 20 s, then 1.5 torr for 15 s before releasing the pressure to ambient, followed by annealing on the hot plate at 120 °C for 15 min. After the substrate had cooled down, GuBr solution (2.5 mg ml⁻¹ in 2-propanol) was spun on at 4000 rpm with a ramp of 1000 rpm for 15 s, followed by annealing on the hot plate at 120 °C for 15 min. The C₆₀ (20 nm) and bathocuproine (BCP) (~5 nm) electron transport layers were sequentially deposited *via* thermal evaporation at 0.1 A s⁻¹. Next, a thin sputter-protection layer of SnOx (~10 nm) was deposited by atomic layer deposition at 83 °C (Beneq TFS200). The hot source vessel (55 °C) was charged with N₂ for 200 ms before a 100 ms pulse of the TDMASn, followed by 12 s purge with N₂ at 300 sccm, water pulse for 200 ms and 12 s purge with N₂ at 300 sccm. Then a 30 nm indium zinc oxide (IZO) layer was deposited by sputtering. Finally, the ~100 nm gold electrode was deposited through a shadow mask (cell area 1.0 cm²).

Caelux cell fabrication

Patterned ITO glass substrates with sheet resistance of 15 Ohm sq⁻¹ were cleaned in soap water, DI water, Acetone and then isopropyl alcohol for 15 min respectively, followed by 15 min treatment in UV ozone. NiO of 20 nm were deposited by a RF sputter at 120 W and then annealed in N₂/H₂ (95/5) at 350 °C for 10 min. After cooling down to room temperature, NiO-coated substrates were transferred into N₂-filled glovebox. Perovskite precursor of 1.3 M Cs_{0.09}FA_{0.8}MA_{0.11}Pb(I_{0.95}Br_{0.05})₃ was dissolved in DMF/DMSO (v/v 4/1). 50 µl precursor was deposited on NiO and spun at 4500 rpm for 40 s, and antisolvent DEE of 200 µl was dripped at 7 s before the end of spinning. Then samples were annealed at 110 °C for 10 min. The thickness of perovskite is around 500 nm. PCBM (20 mg ml⁻¹ in chlorobenzene) was deposited on perovskite layer at 1000 rpm for 60 s and followed by BCP (0.5 mg ml⁻¹ in isopropyl alcohol) at 4000 rpm for 20 s. Silver of 120 nm was thermally evaporated under vacuum at 1 × 10⁻⁶ Torr. The active area is 0.13 cm².

Light current density–voltage measurements

Light *J*–*V* measurements were taken at JPL using the Spectrolab X-25 Mk II Solar Simulator. The measured cells were held in vacuum and light *J*–*V* measurements were taken while varying light intensity and temperature. To record the operating temperature of each individual cell tested, a thermocouple was attached to the back of each cell. The bulb light intensity was adjusted for different measurements to correspond to the solar intensity at distances from the sun of 1.0 AU, 5.5 AU, 9.5 AU, 19.2 AU, and 30 AU. These solar intensities were calibrated using InGaP and GaAs standard calibration cells. The cell temperature was lowered using LN₂, and the lowest achievable cell temperature in the lab is –170 °C. Starting at 1.0 AU and then going to lower intensities, light *J*–*V* measurements were taken at 28 °C, 0 °C, –50 °C, –100 °C, –140 °C, –170 °C with a

tolerance of ±5 °C. For each light intensity and temperature condition, the voltage of each cell is swept three times in each the forward and reverse direction, and the resulting current is measured. The standard deviation between the three sweeps was typically <1%, so our results present each cell as the average of the three forward sweeps for consistency and brevity. Additionally, in the presentation of *J*–*V* curves, the figures depicting *J*–*V* curves show the first forward sweep of each case. After measuring at –170 °C, the system was brought back up to 28 °C and a new measurement was taken and compared to the initial 28 °C measurement to verify the reversibility of trends and lamp and cell stability in LILT conditions. In all our experiments, we found that the measurements taken at 28 °C both before and after cooling and heating were consistent, indicating cell and lamp stability. The same exact procedure, equipment, and cells were used for both the pre-radiation and post-radiation measurements.

External quantum efficiency measurements

EQE measurements were taken with JPL's test setup. A Spectral Products DK240 high resolution $\frac{1}{4}$ meter monochromator was used to generate a narrow-spectrum beam, which was directed through a vacuum chamber's optical window and onto the cell mounted within. The beam was calibrated with a bare silicon standard. Vacuum was drawn in the chamber, and EQE was measured at 28 °C, 0 °C, –50 °C, –100 °C, –140 °C, –170 °C, and then again at 28 °C, all with a tolerance of ±5 °C. Calibration and measurements were completed with room lights turned off to reduce error.

Proton radiation

Proton radiation was conducted at the Boeing Radiation Effects Laboratory (BREL). The cells were exposed to 75 keV protons with a flux of 2 × 10⁹ p+cm⁻² s⁻¹ for up to 24 hours in 6 hours increments. These exposures correspond to doses of 4.32 × 10¹³ p+cm⁻², 8.64 × 10¹³ p+cm⁻², 1.30 × 10¹⁴ p+cm⁻², and 1.73 × 10¹⁴ p+cm⁻². The glass layer used to grow the perovskite solar cells acts as an unwanted radiation protection layer for the purpose of this study. Therefore, to evaluate the true impact of proton radiation on perovskites, the solar cells were exposed with the back layer facing the proton beam. To perform the tests, the cells were placed in the radiation chamber at room temperature. The test plate had four columns of cells, with each column corresponding to the amount of time that the cells were irradiated. Each column had one ANU cell, but we could not test the Caelux cells similarly because the Caelux cells are part of a single footprint made up of many cells. Thus, they could not be separated for the radiation test. Additionally, one ANU control cell was shipped and handled like the other cells but not subject to proton radiation. Because proton radiation was conducted at a different facility than the LILT *J*–*V* measurements, shipping and handling time between the irradiation and characterization could introduce the potential for performance recovery or other changes in device performance. While some performance change is unavoidable, all cells, including the control cells, were shipped and handled with the same



protocol to preserve the validity of comparisons between cells at difference radiation fluences.

Photoluminescence characterization

Further experimental details on the low-temperature photoluminescence measurements can be found in the ESI.†

Conclusions

We have conducted a study of two different types of PSCs in lab-controlled deep space conditions across a wide temperature and light intensity range. The PSCs have different structural compositions, and thus respond differently to light intensity and temperature. The Caelux PSCs start off with a higher efficiency of 19.4% at 1.0 AU and 28 °C compared the ANU PSC efficiency of 14.5%. However, at the most extreme LILT conditions tested (19.2 AU, −170 °C), the ANU PSCs retain a relatively high efficiency of around 14.3% while the Caelux PSCs suffer a larger loss in efficiency, down to 8.6% (19.2 AU, −140 °C). While the rate at which efficiency changes as a function of light intensity and temperature differs between the cells, both PSCs demonstrate a trend where the optimum efficiency temperature decreases as intensity decreases. While this trend is promising for deep space applications, the measured optimal conditions do not correspond with actual mission-relevant conditions, and the PSCs demonstrate reduced performance at very low temperatures. We also show post-radiation LILT data for a champion ANU cell that demonstrates that PSCs can retain a substantial portion of their pre-radiation efficiency even without protective coverglass. However, as with the pre-radiation measurements, we observe reduced performance at very low temperatures. These observations prompt research looking at the cell architecture and material composition to determine if PSCs can be better optimized for LILT conditions, particularly by looking at the charge transport layers and material interfaces as mentioned in our discussion section. With further development, PSCs could potentially enable ultra-light photovoltaic structures for deep space applications.

Author contributions

T. C. contributed to conceptualization, data curation and analysis, experimental procedures, and writing. J. P., Y. W., J. H., A. D. B., D. N., and H. T. N. contributed to cell fabrication and material characterization. M. K., C. M., and R. K. contributed to cell characterization and experimental measurements. D. T., W. K., and J. B. contributed to conceptualization and planning of the experimental test conditions. H. A. A., T. P. W., and J. G. contributed to conceptualization, analysis, reviewing, and supervising the work. All authors have discussed the results and commented on the manuscript.

Conflicts of interest

The authors declare no conflicts of interest.

Acknowledgements

The author would like to thank Joel Schwartz for helpful discussions. This research was carried out at The Australian National University, Caelux Corporation, the California Institute of Technology, and the Jet Propulsion Laboratory, California Institute of Technology, under contract with the National Aeronautics and Space Administration (80NM0018D0004). This work was supported by the Australian Government through the Australian Renewable Energy Agency (ARENA) and the Australian Research Council. J. P., H. T. N., and Y. W. are recipients of postdoctoral fellowships from the Australian Centre for Advanced Photovoltaics (ACAP). T. P. W. is the recipient of an Australian Research Council Future Fellowship (project number FT180100302) funded by the Australian Government. Responsibility for the views, information, or advice expressed herein is not accepted by the Australian Government. Work performed at Caltech was supported by the Space Solar Power Project. We acknowledge the assistance of Arthur Rugtvedt and Nathaniel Williams at BREL.

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