

Experimental determination of power losses and heat generation in solar cells for photovoltaic-thermal applications

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Solar cell thermal recovery has recently attracted more and more attention as a viable solution to increase photovoltaic efficiency. However the convenience of the implementation of such a strategy is bound to the precise evaluation of the recoverable thermal power, and to a proper definition of the losses occurring within the solar device. In this work we establish a framework in which all solar cell losses are defined and described. Aim is to determine the components of the thermal fraction. We therefore describe an experimental method to precisely compute these components from the measurement of the external quantum efficiency, the current-voltage characteristics, and the reflectivity of the solar cell. Applying this method to three different types of devices (bulk, thin film, and multi-junction) we could exploit the relationships among losses for the main three generations of PV cells available nowadays. In addition, since the model is explicitly wavelength-dependent, we could show how thermal losses in all cells occur over the whole solar spectrum, and not only in the infrared region. This demonstrates that profitable thermal harvesting technologies should enable heat recovery over the whole solar spectral range.

I. INTRODUCTION

Photovoltaic (PV) technologies play a dominant role in electric power generation using renewable resources, with PV market expansion and PV conversion efficiency improvements sustaining each other [1]. Enhancements of the solar conversion efficiency are therefore highly desirable to promote further diffusion of solar converters [2]. A possible way to improve solar energy conversion comes from technologies combining PV devices with systems able to recover the heat unavoidably produced within solar cells. Co-generation of warm water or the use of thermoelectric generators (TEGs) provide typical examples [3–8]. In all cases, the profitability of hybrid solar harvesters is limited by the requirement of keeping PV cells at the lowest possible temperature, as their efficiency decreases with temperature at a rate depending on the specific PV material. This is a very well-known hurdle in the making of effective hybrid solar cells, as reported in previous papers by the present authors [9] and by other groups [10]. Reusing heat (to warm up air/water or to further convert it into electricity) may be then from completely counterproductive to quite profitable depending on the PV cell.

Power loss analysis of PV systems is a florid and popular field. Especially regarding silicon solar cells, literature is plenty of methods and tools to evaluate and predict the origin of losses [11–14]. Smaller number of studies focused instead their attention on thermal losses, especially regarding their experimental evaluation [15, 16]. This paper aims at providing a practical, experimental tool to enabling a detailed evaluation of the thermal power fraction (hereafter ξ_u) available in solar cells. The method aims at providing a practical, experimental tool to assess the convenience of hybridization in various types of PV cells. With no need to refer to any specific use of the heat released by the PV cell, it will be shown that such a heat originates from the whole solar spectrum through the many mechanisms responsible for thermal losses occurring in the PV conversion process. This point is of utmost relevance, and may provide suitable guidance to strategies based on the solar-split approach and, more in general, to hybridization schemes using optical (radiative) coupling between the PV and the thermal stage of the harvester.

The experimental method just requires measurements of the external quantum efficiency (EQE), of the current-voltage (IV) characteristics, and of the reflectivity of the solar cell. Data are then elaborated in the framework of a model returning ξ_u along with an evaluation of other (non-recoverable) losses.

The method is validated on three types of solar cells, covering the current range of available PV technologies: a commercial silicon-based bulk solar cell, a lab-made thin-film solar cell made of Copper Indium Gallium Selenide (CIGS), and a commercial triple-junction solar cell (by Spectrolab).

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II. THEORETICAL FRAMEWORK

In a solar cell the unconverted fraction (ϕ_{loss}) of the incoming solar power is the complement to one of the power conversion efficiency η_{pv} , which is defined as the ratio between the output power P_{el} and of the input solar power GA_{abs}

$$\phi_{\text{loss}} = 1 - \eta_{\text{pv}} = 1 - \frac{P_{\text{el}}}{GA_{\text{abs}}} \quad (1)$$

with G the solar irradiance, and A_{abs} the cell area. The power loss fraction is the sum of different kinds of losses.

We can sort them in four main classes:

optical losses (L_1), namely reflection losses ($L_{1\text{R}}$), transmission losses ($L_{1\text{T}}$), contact grid shadowing ($L_{1\text{sh}}$), and absorptions which cannot generate charge carriers ($L_{1\text{abs}}$)

source-absorber mismatch losses (L_2) due to the under-gap portion of the solar spectrum ($L_{2\text{a}}$), and carrier thermalization ($L_{2\text{b}}$) accounting for the voltage drop to the conduction band edge

electron-hole recombination current losses (L_3) which can be either radiative ($L_{3\text{rad}}$) or non-radiative ($L_{3\text{Nrad-J}}$), or due to electrical shunts $L_{3\text{sh}}$

electron-hole recombination voltage losses (L_4) which accounts for the voltage loss associated to the L_3 class

Actually every L_3 loss has a voltage drop counterpart (cf. Appendix A for further details). These voltage drops are why solar cells exhibit voltages smaller than E_{g}/q , and their sum actually accounts for the difference between E_{g}/q and voltage at maximum power V_{mp} .

All the losses listed above contribute to set the cell conversion ratio:

$$\eta_{\text{pv}} = 1 - \sum_{i=1}^4 L_i \equiv 1 - \phi_{\text{loss}} \quad (2)$$

A pictorial view of the loss mechanisms is reported in Fig. 1, where thermal losses are encircled in solid red square.

Note that not all losses are converted into heat within the device. Therefore, the usable thermal fraction ξ_u is smaller than ϕ_{loss} . Specifically, $L_{1\text{R}}$ and $L_{1\text{T}}$ are portions of the solar spectrum which are totally not absorbed, and thus do not contribute to ξ_u . In addition the contact grid can either absorb or reflect light, thus a portion of $L_{1\text{sh}}$ can contribute to ξ_u , while the remaining should be added to $L_{1\text{R}}$. Considering the small contribution of the grid shadowing on the total device area, in this work we will make the assumption that all the light hitting the contacts will contribute to set the total reflection $L_{1\text{R-tot}}$ (Eq. 3).

Regarding radiative recombination ($L_{3\text{rad}}$) the photon generated by the recombination process either leaves the system or are re-absorbed, and eventually generate an electron-hole pair that is involved in a heat generation process. In this work we will consider all the photons generated by radiative recombination as emitted by the device and not re-absorbed. Thus $L_{3\text{rad}}$ will contribute to the light reflected back by the device, setting $L_{1\text{R-tot}}$ (Eq. 3).

Considering photon recycling negligible can be a source of error in evaluating thermal losses especially in the case of stacked multi-junction solar cells [17–19]. However in this work we will show that radiative recombination accounts for a very small fraction of the whole loss (1-3%) showing how this assumption leads to marginal inaccuracies only. In addition this approximation can be easily relaxed following Dupré et al. [16] considering a ratio for any of the recycling mechanisms that the emitted photon could encounter (leaving the cell, being absorbed by a process generating heat, or being absorbed by a process generating carriers). The problem with this approach is however to determine exact values for these ratios.

As of $L_{2\text{a}}$, instead, since it cannot be absorbed by the absorber layer it is generally lost by three mechanisms. It may be reflected (and thus contributes to $L_{1\text{R}}$), or it is transmitted through the solar cell without interacting with it (and thus contributes to $L_{1\text{T}}$), or it is absorbed by other cell layers (e.g. the window layers or the back contact) or by defects and traps, thus contributing to $L_{1\text{abs}}$. Hereafter we will refer to these three mechanisms respectively as $L_{2\text{a-R}}$, $L_{2\text{a-T}}$ and $L_{2\text{a-abs}}$. Thus, the total reflection and absorption losses can be written as

$$L_{1\text{R-tot}} = L_{1\text{R}} + L_{1\text{sh}} + L_{3\text{rad}} + L_{2\text{a-R}} \quad (3)$$

while

$$L_{\text{abs-tot}} = L_{1\text{abs}} + L_{2\text{a-abs}} \quad (4)$$

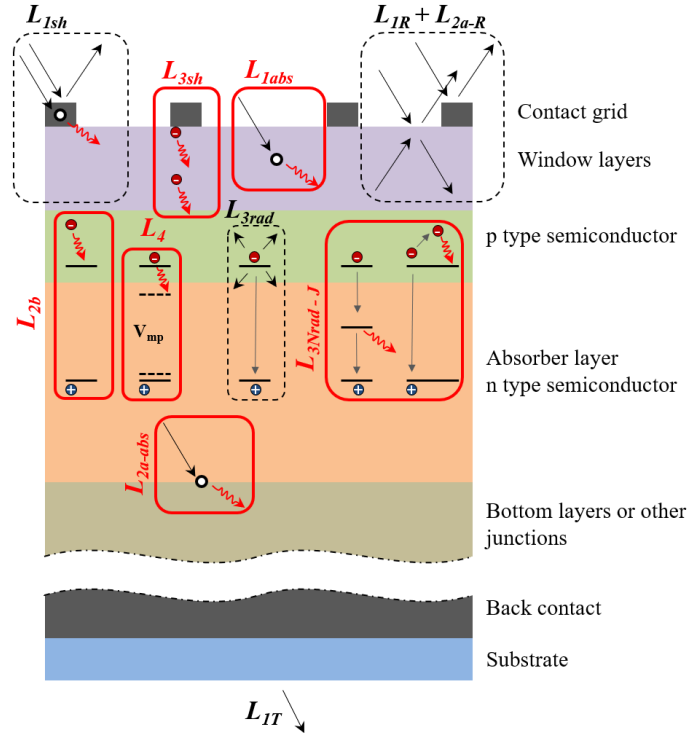


FIG. 1: Pictorial view of a general solar cell structure, with the losses occurring in it. Red solid squares marks thermal losses, black arrows the incoming radiation, and red undulating arrows the losses.

Thus the usable thermal power fraction reads

$$\xi_u = L_{\text{abs-tot}} + L_{2b} + L_{3sh} + L_{3\text{Nrad-J}} + L_4 \quad (5)$$

or, alternatively,

$$\xi_u = 1 - (\eta_{\text{pv}} + L_{1\text{R-tot}} + L_{1\text{T}} + L_{2\text{a-T}}) \quad (6)$$

In the following we will show how to quantify terms in Eq. 5, and the other losses as well, from the spectral analysis of the EQE, the reflectivity R , and the IV characteristics of the device.

A. Quantum Efficiency

In the field of photovoltaics the EQE is defined as the ratio between the number of photons reaching the PV device and the number of electrons contributing to the output electrical current produced by the device. Experimentally, EQE can be obtained as

$$EQE(\lambda) = \frac{I_{\text{out}}(\lambda)}{I_{\text{ph}}(\lambda)} \quad (7)$$

where $I_{\text{out}}(\lambda)$ is the device output current generated by a monochromatic radiation of wavelength λ , and $I_{\text{ph}}(\lambda)$ is the current that the device would produce if all the incoming photons contributed to the device current. Knowing the spectral dependency of the incident solar power, $I_{\text{ph}}(\lambda)$ can be written as

$$I_{\text{ph}}(\lambda) = \frac{qA_{\text{abs}}G(\lambda)}{hc/\lambda} \quad (8)$$

where $-q$ is the electron charge, $G(\lambda)$ is the spectral solar power density, h is the Planck constant, and c is the speed of light.

The internal quantum efficiency $\text{IQE}(\lambda)$ is instead the quantum efficiency without considering reflection losses, and can be written as

$$\text{IQE}(\lambda) = \frac{\text{EQE}(\lambda)}{(1 - R(\lambda))(1 - T(\lambda))} \quad (9)$$

where $R(\lambda)$ and $T(\lambda)$ are respectively the spectral device reflectivity and transmittance. In this work we consider only solar cells with opaque back contacts so that hereafter we will take $T(\lambda) = 0$. However, the method may be easily extended to transparent back contacts (as often found in organic solar cells) by adding a measurement of $T(\lambda)$ to the characterization.

Using Eqs. 7 and Eq. 9 (with $T(\lambda) = 0$) one immediately obtains

$$\text{IQE}(\lambda) = \frac{I_{\text{out}}(\lambda)}{I_{\text{ph}}(\lambda)} \frac{1}{(1 - R(\lambda))} = \frac{I_{\text{out}}(\lambda)}{I_{\text{gen}}(\lambda)} \quad (10)$$

where $I_{\text{gen}}(\lambda)$ is the current that would be generated by the device if all photons actually entering the PV cell (thus those photons which are not reflected) will contribute to the device current.

Using Eqs. 8, and 10 an explicit expression for $I_{\text{gen}}(\lambda)$ is obtained:

$$I_{\text{gen}}(\lambda) = \frac{qG(\lambda)A_{\text{abs}}(1 - R(\lambda))}{hc/\lambda} \quad (11)$$

Finally, using Eq. 11 one can define the fraction of solar power actually entering the solar cell as

$$G_{\text{gen}}(\lambda) = G(\lambda)(1 - R(\lambda)) \quad (12)$$

B. Determination of Losses

For the sake of clarity, it is useful to summarize the main assumptions made in the model.

1. the model neglects the photons that could be absorbed by the metallic contact grid and contribute to ξ_u , assuming that all photons hitting the contacts are reflected
2. the model neglects photon recycling for radiative recombination, considering all these photons as emitted
3. the model takes into account only solar cells with opaque back contact, namely $T(\lambda) = 0$, and thus $L_{1\text{T}} = L_{2\text{a-T}} = 0$

Losses may be now related to measurable quantities.

Since $R(\lambda)$ is defined as the whole device spectral reflectivity (thus accounting also for the contributions from $L_{1\text{sh}}$, $L_{2\text{a-R}}$, and $L_{3\text{rad}}$) its relationship with the (integral) loss $L_{1\text{R-tot}}$ is immediate, namely

$$L_{1\text{R-tot}} = L_{1\text{R}} + L_{1\text{sh}} + L_{3\text{rad}} + L_{2\text{a-R}} = \frac{\int G(\lambda)R(\lambda)d\lambda}{\int G(\lambda)d\lambda} \quad (13)$$

In addition the spectral dependency of $L_{1\text{R-tot}}$ is simply given by

$$L_{1\text{R-tot}}(\lambda) \propto R(\lambda) \quad (14)$$

Likely conversions of spectral into integral quantities (and viceversa) may be carried out for all losses and wavelength-dependent parameters.

Thus, using Eq. 13 for $L_{1\text{R-tot}}$, and Eq. 12 for $G_{\text{gen}}(\lambda)$ one can actually calculate all remaining losses as follows.

The under-gap fraction $L_{2\text{a}}$ which contributes to $L_{\text{abs-tot}}$ reads

$$L_{2\text{a-abs}}(\lambda) = \frac{G_{\text{gen}}(\lambda)}{G(\lambda)} H(\lambda - \lambda_g) \quad (15)$$

where $H(z)$ is the Heaviside step function

$$H(z) = \begin{cases} 1 & \text{for } z > 0 \\ 0 & \text{otherwise} \end{cases} \quad (16)$$

and $\lambda_g = hc/E_g$, with E_g the energy gap of the absorber material. This is clearly an approximation. Actually, the absorbance of a semiconductor, especially for indirect energy gaps, is not a step function. This leads to an underestimation of the thermal components coming from losses that involve the part of the solar spectrum with energy higher than the absorber material E_g (namely L_{2b} , $L_{3\text{Nrad-J}}$, $L_{3\text{sh}}$, and L_4), and an overestimation of L_{2a} , that depends upon the absorption of photons with energy lower than E_g .

The carrier thermalization fraction L_{2b} , accounting for the electron-hole relaxation to the band edge, is instead

$$L_{2b}(\lambda) = \frac{G_{\text{gen}}(\lambda)IQE(\lambda)}{G(\lambda)} \left(\frac{\lambda_g}{\lambda} - 1 \right) H(\lambda_g - \lambda) \quad (17)$$

A likely equation is valid for the sum of all the L_4 losses accounting for the relaxation between the band edge and the energy corresponding to the voltage at maximum power V_{mp} , at which the solar cell is supposed to work:

$$L_4(\lambda) = L_{4\text{carnot}} + L_{4\text{boltz}} + L_{4\text{Nrad-V}} + L_{4s} = \frac{G_{\text{gen}}(\lambda)IQE(\lambda)}{G(\lambda)} \left(1 - \frac{qV_{\text{mp}}}{E_g} \right) H(\lambda_g - \lambda) \quad (18)$$

In Appendix A we show how to split the non-spectral contributions of every L_4 component.

The remaining losses can be only cumulatively estimated. Therefore we conveniently group them under the generic name of thermal losses L_{therm} , computable as

$$L_{\text{therm}}(\lambda) = L_{3\text{Nrad-J}}(\lambda) + L_{1\text{abs}}(\lambda) + L_{3\text{sh}}(\lambda) = \frac{G_{\text{gen}}(\lambda)[1 - IQE(\lambda)]}{G(\lambda)} \quad (19)$$

Using Eq. 4 and 5, along with Eqs. 17–19 one can determine the thermal fraction as a function of the wavelength (or in its integral form) by

$$\xi_u(\lambda) = L_{2a\text{-abs}}(\lambda) + L_{2b}(\lambda) + L_4(\lambda) + L_{\text{therm}}(\lambda) \quad (20)$$

For the sake of clarity we want to point out that since EQE measurements were performed at short-circuit (SC) conditions, instead of at maximum power point (MPP), our results are effected by a small underestimation of L_{therm} in favour of L_{2b} and L_4 losses. This is due to the well known effect of carrier lifetime dependence on injection, for which EQE values change at different applied bias [20]. In this framework it worth to clarify that L_3 losses defined in Sect. II, in our model are essentially seen as short-circuit current losses.

However since L_{therm} , L_{2b} , and L_4 contribute to the overall thermal power, their sum does not change. Thus the evaluation of ξ_u is essentially correct.

A check of the impact of the approximations introduced in the model is achievable by computing $L_{3\text{rad}}$. Actually, considering that radiative recombination is basically the reverse of the optical absorption process, one may estimate the rate of the latter event, obtaining [8]

$$R_{\text{rad}} = RR_0 \left[\exp\left(\frac{eV}{k_B T}\right) - 1 \right] \quad (21)$$

where V is the external voltage, k_B the Boltzmann constant, T the device temperature and

$$RR_0 = \frac{2\pi}{c^2 h^3} \int_{E_g}^{\infty} \frac{E^2 dE}{\exp[E/k_B T] - 1} \quad (22)$$

In this work we will consider solar cells working at room temperature (300 K), but R_{rad} can be found at any temperature using Eq. 21. The radiative recombination rate R_{rad} sets in turn the recombination current I_{rad} . This leads to express $L_{3\text{rad}}$ as

$$L_{3\text{rad}} = \frac{I_{\text{rad}}}{I_{\text{gen}}} = \frac{qR_{\text{rad}}A_{\text{abs}}}{I_{\text{gen}}} \quad (23)$$

that, in view of Eqs. 11 and 12, becomes

$$L_{3\text{rad}}(\lambda) = \frac{hcR_{\text{rad}}}{\lambda G_{\text{gen}}(\lambda)} \quad (24)$$

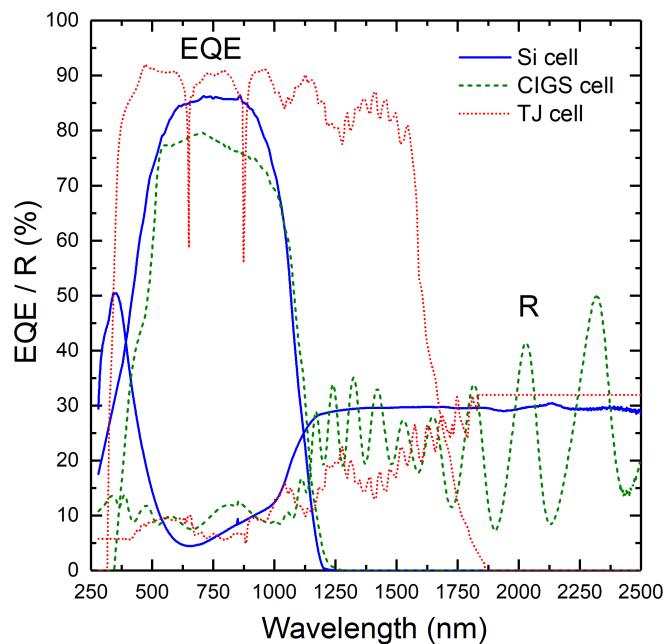


FIG. 2: $EQE(\lambda)$ and $R(\lambda)$ for the three solar cells analysed in this work. Data for the TJ solar cell were obtained from literature [21].

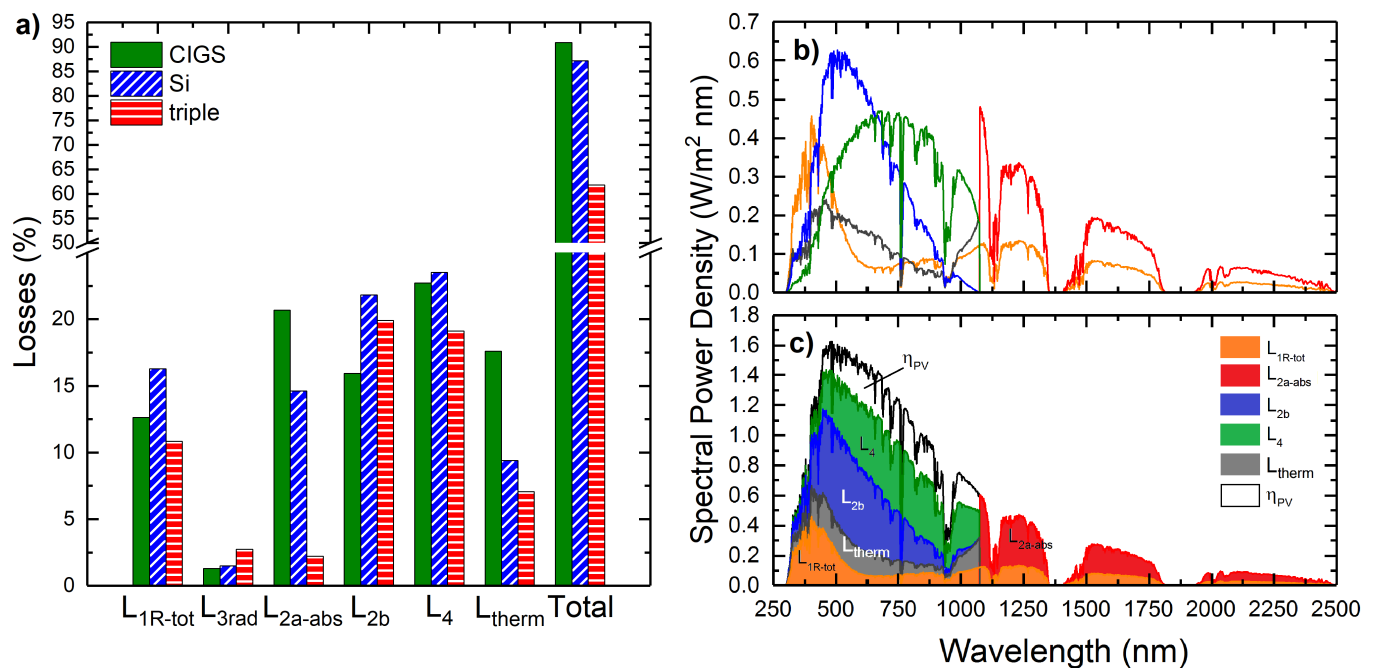


FIG. 3: (a) Computed losses for the three solar cells. (b) Spectral dependency of losses in the Si cell. (c) Cumulative spectral dependency of losses for the Si cell compared to the solar spectrum.

III. MATERIALS AND EXPERIMENTAL

In this work the losses of three different types of solar cells were evaluated. The first solar cell was a commercial, single-junction, bulk solar cell made of multicrystalline silicon (hereafter Si cell). The second solar cell was a lab-made single-junction thin film CIGS solar cell (hereafter CIGS cell). This cell was manufactured following a well-established

	CIGS	Si	TJ 1	TJ 2	TJ 3
η_{pv} (%)	10.03	12.89	18.82	14.94	7.30
V_{mp} (eV)	0.39	0.44	1.31	1.07	0.40
E_g (eV)	1.25	1.16	1.89	1.41	0.67

TABLE I: Values of η_{pv} , V_{mp} , and E_g for the three types of solar cells analysed in this work.

Losses class	Loss name	Description	Measurable quantity	Equation
L_1	L_{1R}	Reflection loss	$L_{1R-tot} = L_{1R} + L_{1sh} + L_{3rad} + L_{2a-R}$	Eq. 13
	L_{1T}	Transmission loss	Neglected in the model	none
	L_{1sh}	Contact grid shadowing loss	$L_{1R-tot} = L_{1R} + L_{1sh} + L_{3rad} + L_{2a-R}$	Eq. 13
	L_{1abs}	Absorption not generating carriers	$L_{therm} = L_{3Nrad-J} + L_{1abs} + L_{3sh}$	Eq. 19
L_2	L_{2a-R}	Under-gap contribution to reflection	$L_{1R-tot} = L_{1R} + L_{1sh} + L_{3rad} + L_{2a-R}$	Eq. 13
	L_{2a-T}	Under-gap contribution to transmission	Neglected in the model	none
	L_{2a-abs}	Under-gap contribution to L_{1abs}	L_{2a-abs}	Eq. 15
	L_{2b}	Carrier thermalization	L_{2b}	Eq. 17
L_3	L_{3rad}	Radiative recombination current loss	L_{3rad}	Eq. 24
	$L_{3Nrad-J}$	Non-radiative recombination current loss	$L_{therm} = L_{3Nrad-J} + L_{1abs} + L_{3sh}$	Eq. 19
	L_{3sh}	Current loss due to shunt resistance	$L_{therm} = L_{3Nrad-J} + L_{1abs} + L_{3sh}$	Eq. 19
L_4	$L_{4carnot}$	Radiative recombination voltage loss - same solid angle than absorption	$L_4 = L_{4carnot} + L_{4boltz} + L_{4Nrad-V} + L_{4s}$	Eq. 18
	L_{4boltz}	Radiative recombination voltage loss - difference between solid angles	$L_4 = L_{4carnot} + L_{4boltz} + L_{4Nrad-V} + L_{4s}$	Eq. 18
	$L_{4Nrad-V}$	Non-radiative recombination voltage loss	$L_4 = L_{4carnot} + L_{4boltz} + L_{4Nrad-V} + L_{4s}$	Eq. 18
	L_{4s}	Voltage loss due to series resistance	$L_4 = L_{4carnot} + L_{4boltz} + L_{4Nrad-V} + L_{4s}$	Eq. 18

TABLE II: Summary of the losses, the measurable quantities, and the related equations.

procedure reported in a previous work [22]. Both cells were measured using the same procedure and the same experimental setup. A SpeQuest Lot-Oriel quantum efficiency system was used to measure EQEs. Spectral response curves of PV devices were measured from 350 nm to 1800 nm with a 10 nm wavelength increment. Current-voltage (I - V) characteristics were recorded under 1 Sun (100 mW/cm^2) illumination in Air Mass 1.5G conditions as generated by a Thermo Oriel Solar simulator. Finally, $R(\lambda)$ was measured using a Jasco V-570 spectrometer equipped with an integrating sphere with a diameter of 60 mm between 250 and 2500 nm.

The last solar cell was instead a commercial triple-junction GaInP/GaInAs/Ge solar cell (hereafter TJ cell) developed by Spectrolab, and the data needed for loss evaluation were found in literature [21].

Figure 2 reports $EQE(\lambda)$ and $R(\lambda)$ data for Si, and CIGS cell, along with the data available for the TJ cell. Table I shows instead the efficiencies and the voltage at maximum power (obtained from I-V characteristics) along with the E_g values obtained from EQE measurements following a method reported in a previous publication [23].

A summary of the losses defined, and the related measurable quantities is reported in Tab. II. The procedure to access all loss terms is instead summarized for reader's convenience as follows:

1. inputting $R(\lambda)$ into Eq. 12 and making use of standard $G(\lambda)$ data one computes $G_{gen}(\lambda)$
2. L_{1R-tot} , is computed from Eq. 13
3. L_{2a-abs} is then obtained from Eq. 15
4. L_{2b} follows from Eq. 17
5. L_4 is computed from Eq. 18
6. L_{therm} is found from Eq. 19.
7. the last contribution, namely L_{3rad} is calculated from Eq. 24.

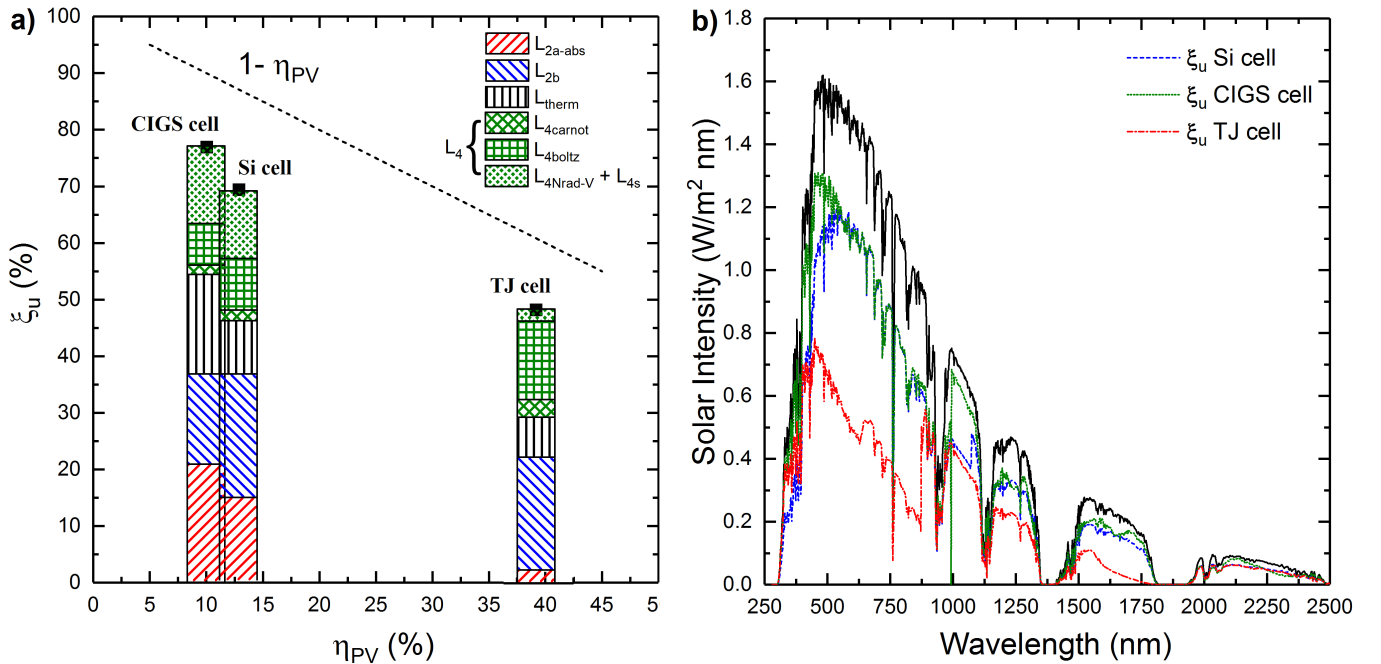


FIG. 4: (a) Bar graph of ξ_u and of its components vs. the cell efficiency for the three devices. The gap between the bars and the dashed line is the non-thermal lost power fraction. See Appendix A for the L_4 components. (b) Spectral dependency of ξ_u for the three cells.

IV. RESULTS AND DISCUSSION

Figure 3 a and Table III report the losses computed for the three solar cells. As expected, the total loss is higher for single-junction (CIGS and Si) solar cells. The sum of the total losses and of the cell efficiencies returns $\approx 100\%$ for all devices, with a maximum deviation of $\pm 1\%$. This result validates the model and the suitability of the approximations it relies upon as well.

Figure 3 a also clarifies that L_{2a-abs} and L_{therm} are mostly responsible for the loss differences among cells. Specifically, while the under-gap absorption loss L_{2a-abs} is almost negligible in TJ, in single-junction cells it is significant. This loss is found to be higher for CIGS because of its larger energy gap, and because of the presence of many layers on top of CIGS (buffer and finalization layers) [22] causing larger absorptions compared to the Si cell. The optimization of layers thickness is known to be a crucial matter in order to achieve optimized performances, especially in thin film technologies [24, 25].

Material quality rules instead L_{therm} which accounts for non-radiative recombination (L_{3Nrad}), absorptions not generating carriers (L_{1abs}), and electrical shunts (L_{3sh}) – all due to the presence of defects. Thus, the higher L_{therm} for CIGS is not surprising, and it actually witnesses the larger defectivity of the material. Silicon and TJ solar cells are instead almost comparable, as the material quality is.

No relevant differences for the upper-gap losses, namely L_{2b} and L_4 are found in the three types of solar cells (we will highlight in Fig. 4 a the differences about L_4 components for the three solar cells analysed). For single-junction

	CIGS	Si	TJ
L_{1R-tot} (%)	12.63	16.29	10.83
L_{3rad} (%)	1.30	1.48	2.73
L_{2a-abs} (%)	20.68	14.63	2.20
L_{2b} (%)	15.94	21.83	19.90
L_{therm} (%)	17.59	9.39	7.07
L_4 (%)	22.71	23.52	19.10
Total (%)	90.86	87.16	61.80

TABLE III: Values of computed losses.

cells, CIGS shows the smallest losses, once again because of its higher E_g . This is in line with what reported in previous works [8]. Interestingly enough, in the case of the TJ solar cell we found L_{2b} and L_4 values very close to that of single-junction solar cells, as the addition of junctions cannot reduce these types of losses. This is consistent with previous evidence [26] in the framework of the Shockley–Queisser limit [27].

Radiative loss L_{3rad} provides a marginal contribution, as expected. However it is interesting to note that it is larger for the TJ solar cell, as anticipated by Hirst et al. [26] who correlated such an increase to the number of junctions. The last contribution L_{1R-tot} mostly depends on the top layer roughness and on the anti-reflective coating used in the cell, so that it cannot be correlated to the absorber characteristics.

In summary, one may conclude that:

1. the material and device quality impact mainly on L_{therm} ;
2. For single-junction solar cells the energy gap set the balance between L_{2a-abs} (that increases with E_g) and L_{2b} (larger for smaller E_g);
3. Multi-junction solar cells are very effective at limiting L_{2a-abs} but cannot avoid most of the L_{2b} and L_4 contributions.

Since all the losses were computed as a function of the wavelength, one may consider their spectral dependence on the wavelength (Fig. 3 b). The reported case (Si cell) is representative of the trends observed also in the other cells. Figure 3 b reports the spectral dependency of the losses calculated for the Si cell, while Fig. 3 c shows their cumulative spectral dependency, with respect to the solar spectrum.

Concerning the thermal power loss, a plot of ξ_u vs. the cell efficiency η_{PV} (Fig. 4 a) shows that ξ_u parallels $1 - \eta_{PV}$, rescaled by $\approx 10 - 15\%$. The downshift depends on L_{1R-tot} (cf. Eq. 6). Fig. 4 a shows also the L_4 components (see Appendix A). It is interesting to note how the total L_4 loss, which is almost equal for all the cells, actually results from different combination of its components. In fact it can be seen how the higher radiative recombination in the TJ solar cell leads to a higher $L_{4carnot}$, and L_{4boltz} contributions, which compensate the smaller ($L_{4Nrad-V} + L_{4s}$) component. For CIGS and Si solar cells instead L_4 is basically equally split between ($L_{4carnot} + L_{4boltz}$) and ($L_{4Nrad-V} + L_{4s}$).

From the spectral dependency of ξ_u showed in Fig. 4 b, it is possible to see how the thermal fraction is quite equally distributed over the whole solar spectrum, and it is *not* peaked in the infrared region. Therefore, whichever strategy is used to recover ξ_u , it should be conceived so as to collect the widest spectral range. This leads to two rather important conclusions regarding spectrum splitting-based thermal recovery strategies, which are normally devoted to the harvesting of the infrared part of the solar spectrum [28–30]. First, the use of such solutions in conjunction with multiple-junction cells may not be effective enough to justify the additional costs and complexity of the overall converter, as the harvester and the multi-junction policies compete to each other in the conversion of the long-wavelength part of the solar spectrum. Second, they are necessarily sub-optimal, as the thermal power output is spread over the whole solar spectrum. Therefore, thermal harvesters should operate collecting heat at all wavelengths, covering also the short-wavelength region where heat resulting from carrier thermalization is larger.

It is worth stressing that these conclusions are limited to solar cells operating at room temperature. Clearly enough, at higher temperature the solar cell efficiency is expected to decrease [31] because of the increase of some losses. In particular, since the temperature sensitivity of solar cells is mainly due to a higher recombination ratio, L_{3rad} and $L_{3Nrad-J}$ are expected to increase significantly, impacting consequently on $L_{4carnot}$, L_{4boltz} , and $L_{4Nrad-V}$ losses [16]. A minor effect is instead expected on L_{2b} , and L_{4s} losses, respectively due to the slight change of the energy gap of the absorber material, and to the associated small variation of the current flowing within the device.

V. CONCLUSIONS

In this work we have reported a method to refine the evaluation of the usable thermal power released by solar cells. The method is based on a novel approach to the analysis of EQE, IV , and reflectance measurements. It has been shown to be applicable to any kind of PV devices, and it is therefore very useful for a detailed evaluation of the thermal-recovery potential of a given solar cell.

Its application to three different kinds of solar cells (bulk, thin film, and multi-junction cell) has shown that the material and device quality mostly set the thermal losses L_{therm} . Also, it proved that in single-junction solar cells the energy gap modulates the balance between L_{2a-abs} and L_{2b} . It was also shown that multi-junction cells are very effective at minimizing the L_{2a-abs} term, although they cannot significantly reduce L_{2b} and L_4 losses.

Finally, the study of the spectral dependency of all terms has shown how thermal losses are uniformly distributed over the whole solar spectrum, not only in the infrared region. This sets important constraints to viable thermal recovery strategies implementable when hybridizing PV systems.

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APPENDIX A COMPUTATION OF L_4 COMPONENTS

In this section we show how to split the L_4 components. As mentioned in Sect. II, L_4 losses are voltage drops associated to L_3 losses. Actually current losses L_3 impact the generation-recombination balance, reducing the voltage that the device can generate, and are the reason why solar cells exhibit voltages smaller than E_g/q . The sum of these voltage losses actually accounts for the difference between E_g/q and voltage at maximum power V_{mp} . Previous studies [15, 26, 32–34] showed how the sum of two of such losses corresponds to the radiative recombination L_{3rad} . The first is called Carnot loss ($L_{4carnot}$) with a corresponding voltage drop firstly derived by Landsberg and Badescu [35]:

$$\Delta V_{4carnot} \approx \frac{E_g}{q} \frac{T_c}{T_s} \quad (\text{A.1})$$

with T_c the cell temperature, and T_s the temperature of the Sun. This loss takes into account only radiative emission in the solid angle within which the device absorbs the solar spectrum. The second is instead the so-called Boltzmann voltage loss (L_{4boltz}) which takes into account the difference between the solid angle within which the solar cell absorbs the solar power, and the solid angle within which it emits. The voltage drop associated can be calculated as

$$\Delta V_{4boltz} \approx \frac{k_B T_c}{q} \ln \left(\frac{\Omega_{emit}}{\Omega_{abs}} \right) \quad (\text{A.2})$$

where k_B is the Boltzmann constant and Ω_{emit} and Ω_{abs} respectively the emission and absorption solid angles. We then define with $L_{4Nrad-V}$, and L_{4s} the voltage drops corresponding to non-radiative recombination, and to electrical shunts.

The total voltage drop due to L_4 (hereafter ΔV_4) is therefore equal to

$$\Delta V_4 = \frac{E_g}{q} - V_{mp} = \Delta V_{4carnot} + \Delta V_{4boltz} + \Delta V_{4Nrad-V} + \Delta V_{4s} \quad (\text{A.3})$$

While V_{mp} is known from the solar cell current-voltage characteristic, and the Carnot and Boltzmann contributions are known from Eqs. A.1 and A.2, one can obtain the sum of the two unknown voltage drops as

$$\Delta V_{4Nrad-V} + \Delta V_{4s} = \frac{E_g}{q} - V_{mp} - \Delta V_{4carnot} - \Delta V_{4boltz} \quad (\text{A.4})$$

Finally knowing from Eq. 18 the total L_4 , and from Eqs. A.1, A.2, and A.4, the ratio between the different components, one can sort out the loss components $L_{4carnot}$, L_{4boltz} , and $(L_{4Nrad-V} + L_{4s})$.

It is worth to point out that ΔV_{4s} can also be extracted by the determination of the solar cell series resistance as

$$\Delta V_{4s} = R_s I_{mp} \quad (\text{A.5})$$

where R_s is the series resistance which can be obtained from the solar cell IV characteristic by several methods [36], and I_{mp} the solar cell current at maximum power.

Note that the method does not allow to obtain the spectral dependency of the L_4 components.

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- [1] IEA-PVS, Tech. Rep., IEA-PVS (2015).
- [2] A. Polman, M. Knight, E. C. Garnett, B. Ehrler, and W. C. Sinke, *Science* **352** (2016).
- [3] K. Yazawa and A. Shakouri, in *InterPACK 2011, 6-8 July 2011, Portland, USA.* (2011), pp. 1–7, ISBN 9780791844618.
- [4] V. V. Tyagi, S. C. Kaushik, and S. K. Tyagi, *Renewable and Sustainable Energy Reviews* **16**, 1383 (2012), ISSN 13640321.
- [5] N. Wang, L. Han, H. He, N.-H. Park, and K. Koumoto, *Energy Environ. Sci.* **4**, 3676 (2011), ISSN 1754-5692.
- [6] K.-T. Park, S.-M. Shin, A. S. Tazebay, H.-D. Um, J.-Y. Jung, S.-W. Jee, M.-W. Oh, S.-D. Park, B. Yoo, C. Yu, et al., *Scientific reports* **3**, 422 (2013), ISSN 2045-2322.
- [7] T.-J. Hsueh, J.-M. Shieh, and Y.-M. Yeh, *Progress in Photovoltaics: Research and Applications* **23**, 507 (2015), ISSN 10627995.
- [8] B. Lorenzi, M. Acciarri, and D. Narducci, *Journal of Materials Research* **30**, 2663 (2015), ISSN 0884-2914.
- [9] D. Narducci and B. Lorenzi, *IEEE Transactions on Nanotechnology* **15**, 348 (2016), ISSN 1536125X.
- [10] T. T. Chow, *Applied Energy* **87**, 422 (2010), ISSN 03062619.
- [11] A. G. Aberle, W. Zhang, and B. Hoex, *Energy Procedia* **8**, 244 (2011), ISSN 1876-6102, URL <http://dx.doi.org/10.1016/j.egypro.2011.06.131>.
- [12] J. Greulich and S. Rein, **114**, 204504 (2013).
- [13] J. Wong, S. Duttagupta, R. Stangl, B. Hoex, and A. G. Aberle, **5**, 619 (2015).
- [14] S. Wasmer, A. V. D. Horst, P. Saint-cast, and J. Greulich, **8**, 689 (2018).
- [15] O. Dupré, R. Vaillon, and M. A. Green, *Solar Energy Materials and Solar Cells* **140**, 92 (2015), ISSN 09270248.
- [16] O. Dupré, R. Vaillon, and M. A. Green, *Solar Energy* **140**, 73 (2016), ISSN 0038092X.
- [17] M. A. Steiner, J. F. Geisz, J. Scott Ward, I. Garcia, D. J. Friedman, R. R. King, P. T. Chiu, R. M. France, A. Duda, W. J. Olavarria, et al., *IEEE Journal of Photovoltaics* **6**, 358 (2016).
- [18] X. Sheng, M. H. Yun, C. Zhang, A. M. Al-Okaily, M. Masouraki, L. Shen, S. Wang, W. L. Wilson, J. Y. Kim, P. Ferreira, et al., *Advanced Energy Materials* **5**, 1400919 (2015).
- [19] D. M. Tex, M. Imaizumi, H. Akiyama, and Y. Kanemitsu, *Scientific reports* **6**, 38297 (2016).
- [20] C. Hibberd, F. Plyta, C. Monokroussos, M. Bliss, T. Betts, and R. Gottschalg, *Solar Energy Materials and Solar Cells* **95**, 123 (2011), ISSN 0927-0248.
- [21] R. R. King, C. Fetzer, P. Chiu, E. Rehder, K. Edmondson, and N. Karam, in *ECS Transactions* (2012), vol. 50, pp. 287–295, ISBN 9781607683575, ISSN 19385862.
- [22] M. Acciarri, S. Binetti, A. Le Donne, B. Lorenzi, L. Caccamo, L. Miglio, R. Moneta, S. Marchionna, and M. Meschia, *Crystal Research and Technology* **46**, 871 (2011), ISSN 02321300.
- [23] S. Marchionna, P. Garattini, A. Le Donne, M. Acciarri, S. Tombolato, and S. Binetti, *Thin Solid Films* **542**, 114 (2013), ISSN 00406090.
- [24] S. Yaar, S. Kahraman, S. Çetinkaya, . Apaydn, . Bilican, and . Uluer, *Optik - International Journal for Light and Electron Optics* **127**, 8827 (2016), ISSN 0030-4026.
- [25] M. J. Taghavi, M. Houshmand, M. H. Zandi, and N. E. Gorji, *Superlattices and Microstructures* **97**, 424 (2016), ISSN 10963677, URL <http://dx.doi.org/10.1016/j.spmi.2016.06.031>.
- [26] L. C. Hirst and N. J. Ekins-Daukes, *Progress in Photovoltaics: Research and Applications* **19**, 286 (2011), ISSN 10627995.
- [27] W. Shockley and H. J. Queisser, *Journal of Applied Physics* **32**, 422 (1961), ISSN 00218979.
- [28] Y. Vorobiev, J. González-Hernández, P. Vorobiev, and L. Bulat, *Solar Energy* **80**, 170 (2006), ISSN 0038092X.
- [29] D. Kraemer, L. Hu, A. Muto, X. Chen, G. Chen, and M. Chiesa, *Applied Physics Letters* **92**, 243503 (2008), ISSN 00036951.
- [30] M. Mizoshiri, M. Mikami, and K. Ozaki, *Japanese Journal of Applied Physics* **51**, 06FL07 (2012), ISSN 00214922.
- [31] A. Virtuani, D. Pavanello, and G. Friesen, in *Proc. 25th EU PVSEC* (2010), pp. 4248 – 4252.
- [32] C. H. Henry, *Journal of Applied Physics* **51**, 4494 (1980), ISSN 00218979.
- [33] P. F. Baldasaro, J. E. Reynolds, G. W. Charache, D. M. DePoy, C. T. Ballinger, T. Donovan, and J. M. Borrego, *Journal of Applied Physics* **89**, 3319 (2001), ISSN 00218979.
- [34] T. Markvart, *Applied Physics Letters* **91**, 2005 (2007), ISSN 00036951.
- [35] P. T. Landsberg and V. Badescu, *Journal of Physics D: Applied Physics* **33**, 3004 (2000), ISSN 0022-3727.
- [36] D. T. Cofas, P. A. Cofas, D. Ursutiu, and C. Samoila, in *2012 13th International Conference on Optimization of Electrical and Electronic Equipment (OPTIM)* (IEEE, 2012), pp. 966–972, ISBN 978-1-4673-1653-8.