Fast and Nondestructive Detection on the EVA Gel Content in Photovoltaic Modules by Optical Reflection

Heng-Yu Li, Yun Luo, Christophe Ballif, and Laure-Emmanuelle Perret-Aebi

Abstract—Poly(ethylene-co-vinyl acetate) (EVA) has been the dominating material in the photovoltaic (PV) encapsulant market for decades, owing to its superior cost-performance balance. To achieve its desired material properties, EVA undergoes a curing reaction during the module encapsulation process. The resulting EVA gel content after encapsulation is an important criterion for the module encapsulation quality control. Normally, the determination of gel content is achieved using a tedious solvent extraction method. In this paper, a fast and nondestructive detection method on the EVA gel content based on the optical reflection is explored. First, the homogeneity of the EVA gel content distribution after the standard EVA encapsulation process is studied. Then, the feasibility of the proposed optical approach applied to transparent modules is investigated. After that, a method is developed to apply it to opaque modules by incorporating a mirror into the module construction. It was found that the haze factor of the reflected light correlates well with the EVA gel content in the opaque modules. This proof-of-concept work could lead to the development of a fast and nondestructive tool for detecting the EVA gel content in both transparent and opaque PV modules, which is promising for integration as an inline diagnostic tool in the module manufacturing line.

Index Terms—Curing, optical reflection, packaging, photovoltaic cells, polymer films.

I. INTRODUCTION

CLEAN and renewable electricity resources are of utmost importance to the sustainable development of human society. As a promising candidate, photovoltaic (PV) technology and industry have both experienced tremendous growth over the past decades [1]. However, in order to further compete in utility scale with the traditional fossil fuel-based electricity generation, the levelized cost of electricity of PV technology still needs to decline further [2], [3]. Toward that aim, improvements in the reliability, i.e., the life time, of PV modules are critical.

The PV modules deployed in the natural environment are exposed to the combination of multiple stresses including UV radiation, heat, chemical ingress, mechanical impact, etc., [4]–[6]. To protect the active energy conversion unit, i.e., the solar cells from the harsh operational conditions, it is necessary to encapsulate them using a qualified packaging scheme. In fact, nearly all the failure modes discovered in the field-degraded modules are due to the failure of the module packaging [7], [8]. Thus, the encapsulation materials and the quality of the encapsulation process are decisive for the PV module reliability and, hence, the future of PV technology.

Typical crystalline Si-based PV modules are composed of a front glass superstrate, a glass or polymeric back substrate with the stringed solar cells are sandwiched in the middle [9]. To achieve structural integrity, two layers of encapsulants are utilized right above and below the cells. The encapsulant not only functions as the adhesive, but also as the optical index matching medium, as a barrier for chemical ingress (moisture, oxygen, etc.), as a heat conductor, and as a dielectric medium, etc. Currently, ethylene-co-vinyl acetate (EVA) occupies the majority of the PV encapsulant market. EVA encapsulant is normally crosslinked during the module encapsulation process in order to achieve the desired material properties. EVA reaches certain curing level at the end of the encapsulation process. In the PV industry, this is frequently represented by a value called gel content, which equals the weight percentage of the gel network (cross-linked portion) in the overall EVA.

Due to its importance, the EVA gel content is often adopted as criteria for judging the encapsulation quality of the PV modules. In the module manufacturing process, a number of modules have to be removed at a certain frequency for monitoring the EVA gel content inside. Historically, EVA gel content is measured by the solvent extraction method, where an organic solvent like toluene or xylene are used to dissolve the soluble portion and extract the cross-linked portion [10], [11]. The gel content is calculated as the weight percentage of the nonsoluble part of the EVA. Typically manufacturers require that the EVA gel content in the encapsulated modules should exceed 80%. The solvent extraction method is not only time consuming (usually requiring about 24 h), but a piece of EVA has to be taken from the PV module as well. It is destructive and implies an economic loss.

In today’s high-output PV module manufacturing lines, a much faster and nondestructive quality control method is in demand to ensure a high production yield.

More than ten types of different approaches have been proposed to develop fast quality control on the EVA gel content. Proposed techniques like Raman spectroscopy [11], scanning acoustic microscopy [12], differential scanning...
II. EXPERIMENTAL SETUP

A. Encapsulation Process

Various commercial grades of EVA encapsulants have been tested for the encapsulation, DSC, and optical studies. Due to the similarity of the results, only one representative EVA grade is presented here. The encapsulation and, thus, the curing of EVA are performed in a controlled manner in a flat-bed vacuum bag laminator (3S S1815 laminator). The temperature and pressure profiles of a typical lamination process are shown in Fig. 1. After a preheating step on pins for approximately 5 min (plate temperature at 140 °C unless otherwise mentioned), the pins are retracted downward and 1 bar pressure is immediately applied to the laminates. The pressure is applied starting at the onset of the curing time, which occurred under vacuum at below 1 mbar. After a preset curing time between 0 and 1300 s, the pressure is removed and the sample is transferred out of the lamination chamber and cooled naturally in ambient environment [9].

B. Differential Scanning Calorimetry Study

EVA samples with different curing times are analyzed using a DSC setup (Mettler Toledo DSC1). The DSC samples are prepared as 1-mm thick circular discs of 5–10 mg in an aluminum crucible. Thermograms are recorded under constant nitrogen flow from 30 to 200 °C at a heating rate of 15 °C·min⁻¹ (unless otherwise indicated).

C. Measurement on the Temperature of Heating Plate

The temperature measurement is performed with a PT100 T-sensor connected to a data logger. Measurements are performed on 20 locations on the heating plate. The distribution of the measurement sites are shown in Fig. 2, where the number displayed is the distance in millimeters. The temperature of the heating plate is set to be 140 °C and dwell for 1 h. At each measurement site, the T-sensor is pressed firmly onto the heating plate for 5 min till the temperature displayed in the data logger is stabilized, and then, the temperature is recorded.

D. UV/VIS Spectroscopy

The optical characterization on the encapsulants is carried out with a UV/Vis/NIR spectrophotometer (Perkin Elmer, Lambda 900). The measured wavelength range is set to be 320–2000 nm. Total transmission and total reflection (TR) are obtained using an integrating sphere. From the UV/Vis spectroscopy, the haze factor of the transmitted or reflected light is calculated. The haze factor is defined as the percentage of the diffused light in the total transmitted or reflected light, which is often used to describe the extent to which light is diffused after traveling through a medium.

III. RESULTS AND DISCUSSIONS

A. Evolution of Homogeneity of Ethylene-Co-Vinyl Acetate Gel Content Distribution

Normally, the EVA gel content in the PV module is measured at one or a few spots in the module. To validate the representativeness of the measurement results, it is necessary to assess the homogeneity of the EVA curing level across the whole module.
As the curing of EVA is a thermally activated process following an Arrhenius-type temperature dependence, the temperature uniformity of the heating plate can greatly influence the homogeneity of the resulting EVA curing level. In this paper, the uniformity of the heating plate in the laminator is studied first.

The result of the temperature mapping is visualized in the contour plot in Fig. 3. As one can see, the heating plate exhibits a uniform temperature distribution with only a deviation of ±1 °C. This result allows us to ignore the influence of the temperature uniformity of heating plate on the homogeneity of the EVA gel content distribution after the curing step.

Besides the temperature, the initial concentration of the curing agent (peroxides) in the uncured EVA encapsulant can also greatly affect the EVA conversion rate. Thus, it is equally important to know the homogeneity of initial additive (mainly peroxides) concentration distribution in the EVA. Here, DSC is used as the tool. As described in [13], the exothermic band in the DSC thermograph of the EVA encapsulant correlates well with the amount of residual peroxides in the EVA. For the DSC sample preparation, 16 samples are cut from 16 evenly spaced locations on a 670 × 710 mm² uncured fresh EVA encapsulant, as drawn in Fig. 4(a).

After the DSC measurements, the enthalpy of the exothermic band in the uncured fresh EVA encapsulant. The result in Fig. 4(b) shows that the average enthalpy is 16.0 ± 1.0 J/g. The small variation suggests that the initial peroxide distribution is rather homogeneous in this commercial-grade EVA encapsulant.

In the discussion above, the two major factors that could lead to the inhomogeneous distribution of the EVA gel content in the encapsulated module are analyzed, which are the uniformity of the plate temperature and the homogeneity of the initial peroxides distribution in the EVA encapsulant. During the module encapsulation process, other issues may also result in the inhomogeneity of the final EVA gel content, like uneven hot press throughout the module layup, glass warping during the preheating step, and location-dependent nonuniform presence of peroxides in EVA at elevated temperature, etc. Therefore, it is very useful to track the evolution of the homogeneity of the EVA gel content in the PV module throughout the encapsulation process.

For this study, four identical module layups are encapsulated with the same encapsulation process. Each of them is taken out at different stages of the process, namely 0 s cured (taken out immediately after preheating for 300 s), 300 s cured (taken out at 300 s after the curing step starts), 600 s cured (taken out at 600 s after the curing step starts), and 900 s cured (taken out at 900 s after the curing step starts). For each module, 16 DSC samples are taken out in the same manner shown in Fig. 4(a). The encapsulation process of all the modules are as follows: $T_{\text{plate}} = 150$ °C; preheating for 300 s; hot press = 1 bar. After the encapsulation, all modules are cooled down with uniform cooling across the module surface to eliminate the influence of the postcuring during cooling. Then, the DSC samples are cut out from the back side of the modules. The procedure of the DSC measurements follows the one for the fresh EVA reported above.

Fig. 5(a) shows the contour plots of the enthalpy distribution at all four stages of the curing steps. One can clearly see the good homogeneity and the development of the enthalpy in the encapsulation process. This development is due to the evolution of the peroxide consumption in the encapsulation process. Fig. 5(b) shows the enthalpies measured in the DSC plotted against the curing time. As the encapsulation progresses, the enthalpy measured in the EVA decreases, indicating the ascent of the EVA curing level. The error bars represent the standard deviation from the average enthalpy of the 16 samples. Notably, in the worst...
case, the standard deviation is less than 8% of the average value. Therefore, this leads to the conclusion that the distribution of the EVA curing level in the encapsulated PV module is sufficiently homogenous throughout the encapsulation process. It validates the representativeness of the results obtained by the technique to be developed here for EVA gel content detection in this paper.

B. Optical Detection in Reflection Mode

The optical detection in transmission mode on the EVA gel content in PV modules has been well documented in [17]. However, the optical detection in transmission mode only works for transparent PV modules. As the majority of crystalline PV modules currently use opaque backsheet instead of glass for the sake of lightweight, it limits greatly the application scale of our optical detection technique. Moreover, from the view point of the device development, it is also believed to be more convenient to operate in reflection mode than in transmission mode.

Here in this section, the development on the optical detection in reflection mode is presented. The front architecture of the PV module is designated to exhibit low reflection. Toward the development of the technique in reflection, first it is necessary to increase the reflection signal from the PV module. One idea is to use a mirror element to enhance the reflectivity. The paramount requirement on the material properties of this mirror material is the low scattering effect at the surface for the light of about 400 nm. Ideally, the haze effect of the mirror itself should be much lower than the hazing effect induced by the EVA layer, in order to increase the sensitivity of this technique. The next requirement is the compatibility of the mirror material with the other module components and the encapsulation process. This will be addressed in the future work on the realization and commercialization of the actual device.

1) Optical Detection in Reflection Mode for Transparent Photovoltaic Modules: For transparent modules, the mirror needs not to be embedded into the module layup, as illustrated in Fig. 6(a). Instead, the mirror can be applied at the exterior of the module. Thus, the application of the mirror does not have to take place prior to the encapsulation process. For the transparent module, a mirror deposited on a plastic substrate (Reflectec) is selected for further test.

In Fig. 7, the TR and diffuse reflection (DR) measured on the Reflectec mirror are shown. At between 400 to 500 nm, the TR of this mirror lies well above 80%. The haze factor of the reflected light is lower than 2% in the studied wavelength range, with the maximum reached at about 400 nm. The hazing effect fulfills the requirement. The main disadvantage of this mirror is its low working temperature at merely 60 °C, which is considerably lower than the typical module encapsulation temperature. Experimentally, we have exposed this mirror to the actual encapsulation condition at 140 °C. After the process, the mirror surface is severely deformed and forms numerous wrinkles, resulting in high hazing effect.

The procedure of the optical detection in reflection mode for transparent modules is proposed as follows. After the module encapsulation, the Reflectec mirror is applied onto the module surface, where no opaque objects like cells and metallizations present on top. To achieve better optical coupling, a layer of index matching liquid is used as the medium layer. The optical detection takes place on the other side of the module. To test this procedure, a series of minimodules are prepared under standard encapsulation process with the curing time of 100, 300, 500, 700, 900, and 1100 s. Total transmission and diffuse transmission are measured on these laminates. The haze factor of the transmitted beam versus the EVA curing time is plotted in Fig. 8(a). The error bar represents the reproducibility of the data measured at five different locations on each sample. It is clearly seen that the haze factor of the transmitted light correlates well with the EVA curing time.

Subsequently, the optical measurement in reflection is performed following the procedure described above. TR and DR are measured, and from which the haze factor of the reflected light is calculated. The results for different curing times are summarized in Fig. 8(b). Evidently, identical trend is observed.
For opaque modules, it is proposed here to embed a mirror into the module layup, as shown in Fig. 6(b).

1) Optical Detection in Transmission Mode for Transparent PV Modules: The size of the mirror can be as small as $1 \times 1$ cm$^2$ in the present testing condition, which is the approximate size of the light spot in the UV/V is spectrometer. With the optimization on the device design, the area of this embedded mirror can be further reduced. In the case of opaque modules, the mirror will have to expose itself to the module encapsulation condition. The Reflectech mirror is not applicable due to its low working temperature. Instead, a type of aluminum mirror (4200 AG MIRO SILVER from Alanod) with a nominal thickness of 0.2 mm has been chosen for this study. The TR and DR of the MIRO mirror are first measured and shown in Fig. 9. Notably, MIRO mirror has reflection over 80% at 400 nm, while the hazing effect of the Reflectech mirror is not applicable due to its low working temperature. Instead, a type of aluminum mirror (4200 AG MIRO SILVER from Alanod) with a nominal thickness of 0.2 mm has been chosen for this study.

2) Optical Detection in Reflection Mode for Opaque Photovoltaic Modules: For opaque modules, it is proposed here to embed a mirror into the module layup, as shown in Fig. 6(b). The size of the mirror can be as small as $1 \times 1$ cm$^2$ in the present testing condition, which is the approximate size of the light spot in the UV/V is spectrometer. With the optimization on the device design, the area of this embedded mirror can be further reduced. In the case of opaque modules, the mirror will have to expose itself to the module encapsulation condition. The Reflectech mirror is not applicable due to its low working temperature. Instead, a type of aluminum mirror (4200 AG MIRO SILVER from Alanod) with a nominal thickness of 0.2 mm has been chosen for this study. The TR and DR of the MIRO mirror are first measured and shown in Fig. 9. Notably, MIRO mirror has reflection over 80% at 400 nm, while the hazing effect of the Reflectech mirror is not applicable due to its low working temperature. Instead, a type of aluminum mirror (4200 AG MIRO SILVER from Alanod) with a nominal thickness of 0.2 mm has been chosen for this study.

To prove the proposed concept of the optical detection on EVA gel content in opaque modules, numerous glass/EVA/glass laminates are encapsulated for different curing time of 100, 300, 500, 700, 900, and 1100 s. All of them are encapsulated with a piece of MIRO mirror. The mirror is located in between the EVA layer and the backsheet in this study. Moreover, to determine the actual EVA gel content after each encapsulation process, one additional EVA sample for DSC study is prepared in parallel.

Fig. 10(a) shows the evolution of the measured enthalpy with the EVA curing time. The small error bar shows the good reproducibility of these DSC measurements. The enthalpy is seen to decline with the increasing curing time. A correlation curve between the EVA gel content and the DSC enthalpy has been set up for the EVA used here [18]. This allows us to translate the enthalpy in Fig. 10(a) into the actual EVA gel content shown in Fig. 10(b). As expected, the EVA gel content increases as the EVA curing time is extended.

The TR and DR are measured at the embedded mirror in the modules with different curing time. From the results, the haze factors of the reflected light from the mirror at 400 nm are calculated and plotted in Fig. 10(c). As the curing time rises up, the haze factor of the reflected light through the EVA layer drops monotonically. Combining Fig. 10(b) and (c), the correlation between the EVA gel content and the measured haze factor can be constructed, as illustrated in Fig. 10(d). With the curing time longer than 500 s, the resulting EVA gel content exceeds over 80%. Fig. 10(d) shows that there exists a good correlation between the measured haze factor of the reflection from the module at the mirror site and the actual EVA gel content in the module. The measurement sensitivity of the optical detection is high, especially in the regime of low EVA gel content. There is certain difficulty in differentiating the EVA with gel content higher than 90%. This is expected to be improved if an alternative mirror with even lower hazing effect is used.

Currently, there is great interest from PV industries in the fast and nondestructive method for the determination of EVA gel content in PV modules. Several companies/institutes have also proposed other concepts based on different techniques [11], [19]. Compared to them, the advantages of the concept presented here are 1) high sensitivity in the range of high gel content (over 80%), 2) potentially applicable to both glass/glass and glass/backsheet PV modules, and 3) ideal for inline diagnosis thanks to its nondestructive nature and unrivaled detection speed (time needed can be well below 5 s). The potential problem of this concept is the embedding mirror needed in the case of the opaque modules, although its size can be minimized to be...
well below 1 cm². In the following realization of the device based on this concept proved above, alternative solutions will be sought to either utilize existing module components (e.g., ribbons) as the mirror or to combine the embedded mirror with other functionalities in the future “smart” PV modules.

IV. CONCLUSION

Nowadays, the PV industry highly desires a fast and nondestructive technique that can be integrated as an inline tool for detecting EVA gel content in PV modules. Toward this goal, an optical approach working in reflection mode based on the UV/V is spectroscopy is proposed here. The EVA gel content is first shown to be uniformly distributed in the studied module configuration after the standard EVA encapsulation process. Then, the proof of the proposed concept is achieved for both transparent and opaque PV modules. It is demonstrated that the proposed approach has high sensitivity and robustness on detecting the EVA gel content in transparent modules with no modification on the conventional module configuration. As for the opaque modules, a mirror is required to be integrated into the module configuration in this paper. A good correlation is found between the EVA gel content and the haze factor of the reflected light from the opaque module. The results show that the proposed approach can easily distinguish the EVA with gel content of below or above 80%. This paper proves the feasibility of the proposed optical approach in reflection mode for detecting the EVA gel content in both transparent and opaque PV modules. It is expected that, with further optimization, it will become a strong candidate as the inline diagnostic tool for the gel content detection in the module manufacturing line.

REFERENCES

[10] Internal unpublished result.

Christophe Ballif received the Master's degree in physics and the Ph.D. degree from École polytechnique fédérale de Lausanne (EPFL), Lausanne, Switzerland, in 1994 and 1998, respectively, focusing on novel photovoltaic materials. He was a Postdoctoral Researcher with the National Renewable Energy Laboratory, Golden, CO, USA, where he was involved with compound semiconductor solar cells (CIGS and CdTe). He was then with the Fraunhofer Institute for Solar Energy Systems, Freiburg, Germany, where he was involved in mono- and multicrystalline silicon photovoltaics till 2003. He then joined the Swiss Federal Laboratories for Materials Testing and Research, Thun, Switzerland, before becoming a Full Professor and a Chair with the Institute of Microsystems, University of Neuchâtel, Neuchâtel, Switzerland, in 2004. In 2009, the Institute was transferred to EPFL. He is currently the Director of the Photovoltaics and Thin-Film Electronics Laboratory within the institute, as well as of the CSEM PV-Center. His research interests include thin-film solar cells, high-efficiency heterojunction crystalline cells, and module technology, contributing to technology transfer and industrialization of novel devices.

Heng-Yu Li received the B.E. degree in materials engineering from the Beijing University of Chemical Technology, Beijing, China, in 2004, the M.Sc. degree in nanosciences from the Delft University of Technology, Delft, The Netherlands, in 2006, and the Ph.D. degree in materials science from the Ecole Polytechnique Fédérale de Lausanne (EPFL), Lausanne, Switzerland, in 2013.

Since 2009, he has been working on the encapsulation process and the reliability of photovoltaic (PV) modules with the Photovoltaics and Thin-Film Electronics Laboratory, EPFL. In 2014, he joined the PV-Center, CSEM, Neuchâtel, Switzerland, where his focus is on the development of reliable speciality PV modules for various applications, including BIPV.
Yun Luo received the Ph.D. degree from the Max-Planck-Institut für Mikrostruktur Physik, Halle, Germany, under the supervision of Prof. U. Gösele in 2005. After three years as a Postdoctoral Fellow with the Ecole Polytechnique Fédérale de Lausanne, Switzerland, she joined Pason SA as a Project Leader and continued as a Technology Manager with 3S Swiss Solar Systems AG and Meyer Burger Technology Group. In 2011, she founded Plus MAT AG, aiming to recycle Silicon from kerf loss for applications in photovoltaics, energy storage, and additive manufacturing.

Laure-Emmanuelle Perret-Aebi received the Ph.D. degree in chemistry from the University of Fribourg, Fribourg, Switzerland, in 2004. After four years of Postdoctoral research with the University of Edinburgh, Edinburgh, U.K., and the University of Neuchâtel, Switzerland, in 2009, she joined the PVlab of the Swiss Federal Institute of Technology (EPFL), Lausanne, Switzerland, as a Group Leader of PV module back-end activities. She is currently a Section Head with the “Modules & Systems Integration” sector, CSEM PV-center, Neuchâtel, Switzerland. Her main research interest includes the development of innovative and highly reliable photovoltaic modules and systems for various applications, such as building, mobility, and land power plants.