Abstract: The aims and objectives of this paper are two-fold: first to investigate the production of a green composite manufactured with renewable materials (i.e., jute fabric as reinforcement, acrylate epoxidase soybean oil (AESO) as matrix and sisal particles (SP) as filler), by the wet layup method; second, to propose a recycling procedure to recover the individual materials and reuse them in the production of a second-life composite. In the first part, different combinations of SP (0, 5 and 10 wt%) and hardener (2, 5 and 10 wt%) were mixed with AESO resin, poured into a mould, cured and submitted to mechanical and physicochemical characterizations to identify the best conditions for the composite production. Virgin green composites with 10 wt% SP, 5 wt% hardener and 5 layers of jute fabric, capable of assuring 91 HA of hardness and 10.6 MPa of tensile strength, were fabricated. The second part describes the recycling process of the composites with acetone, an organic solvent recommended by the safety, health and environmental criteria, to breakdown the resin matrix and recover the jute fabric reinforcement and resin particles, which were then reused to fabricate a second-life composite. Although the hardness values of the second-life composite were smaller (4%) and the tensile strength varied with composition, the absorption of water was considerably reduced (in the range of 22 to 51%). This last result mitigates one of the green composite’s limitations and fosters circular economy by assuring the applications of the second-life composite in the field of transportations, packing and furniture, among others.

Keywords: green composite; recycling; second-life composite; natural fibres; soybean oil; mechanical properties; water absorption

1. Introduction

Nowadays, conventional manufacturing processes are being pushed to their limits to produce increasingly intricate parts at lower cost. If, on the one hand, this stimulates the development of new processes with conventional materials, on the other it promotes the extension of the application domain of existing processes to new types of materials. Composites are in the front line of ongoing developments to replace conventional materials in many different fields of applications, due to
their capability of offering enhanced mechanical, physical and chemical properties with significant reductions in material weight [1,2].

Since most composites are made from petroleum-based polymers and synthetic fibres, their use comes with a price of difficult recyclability and are out of alignment with the current environmental sustainability trends and requirements. In fact, the end-of-life disposal of composites contributes to increase the serious global pollution problem, particularly with regard to greenhouse emissions and toxic gases, as an outcome from incineration and permanent landfill accumulation [2–4]. All these concerns, together with the growing of environmental legislation restriction trends, results in an inevitable pressure towards the use of composites from renewable and natural resources and the development of effective recycling technologies [5,6].

Although some synthetic fibre reinforcements can be directly replaced by alternative sustainable natural fibres, it has been shown that the greatest environmental footprint of composites when they replace metals is due to the use of petroleum-based matrix polymers, rather than the use of reinforcement fibres [7]. The growing interest to develop bio-based resins is, therefore, aligned with the solution of all these problems.

Meanwhile, since the use of composites with petroleum-based polymers cannot yet be replaced in many industrial applications, the solution is to maximize its utilization to reduce end-of-life disposal as plastic waste. Hodzic [4], for example, proposes the following solutions: (i) re-create virgin material for other applications using the extracted monomers; (ii) develop cost-effective recycling processes that justify their re-use; (iii) develop biodegradable and naturally sourced polymers; (iv) develop biochemical degradation processes to shorten polymeric chains.

Regarding recyclability and reutilization, the past few years have seen the development of several technologies capable of handling the problems posed by the recovery of the individual constituent materials from the typical heterogeneous hybrid structure of composites, as shown in Figure 1a. Mechanical processes, such as grinding, as shown in Figure 1b, are relatively cheap and allow for the recovery of the composites as small granulates or powder to be used as reinforcements for polymers or cement applications. Thermal pyrolysis, as shown in Figure 1c, which is more expensive, allows for the production of particles with smaller or longer sizes and shapes at the expense of some loss in mechanical behaviour. It is also possible to recover the fibres by burning the composite, but the resulting mechanical properties will be significantly lower. Chemical processes, as shown in Figure 1d, are also able to produce small and long particles (300 µm to 100 mm) with lower losses in mechanical performance than pyrolysis [8–19].

Although the abovementioned technologies allow for the recovery of some material and are theoretically good for global sustainability, they still have a negative environmental impact due to the use of organic solvents, energy consumption for achieving the required temperatures and pressures and, due to the investments in equipment, R&D and advertisement [20].

Under these circumstances, the development of bio-composites with at least one constituent derived from a natural source seems to be a viable answer to the actual impasse in terms of the environmental impact of composite production versus the impact of their recyclability. Green composites, which are totally made from natural fibres embedded in a plant-based matrix or other natural resin, assume special relevance in the solution of the abovementioned impasse [21–23].

Nowadays it is possible to find a variety of bio-composites produced from polymeric resins from renewable resources, such as vegetable oils [24], poly-L-lactic acid [6] and furan resins [25]. Within plant oils as feedstock for polymeric biomaterial, the utilization of soybean oil in the development of bio-based thermosetting matrices is gaining greater attention [26,27]. Particularly, acrylate epoxidase soybean oil (AESO), which is synthesized from soybean oil via epoxidation and acrylation, has attracted great research interest [27–29].

Several natural fibre reinforcements, such as jute, sisal, bamboo, kenaf and flax have been used to fabricate bio-composites for different applications. [30,31]. Natural fibres have several advantages over traditional reinforcing fibres, such as accessibility from renewable resources, biological degradation,
non-abrasive processing, good adhesion, especially after treatment, and low density and cost. However, they often suffer from moisture absorption, low thermal and UV resistance and quality variations [26].

Research in the field of recovering natural fibre composites is a relatively unexplored topic when compared to the recyclability of synthetic fibre reinforced composites. For the latter, different strategies have been proposed, mainly due to the relatively high cost and energy consumption to produce synthetic fibres [15–19,32,33]. Although the number of studies in the recyclability of natural fibre reinforcement composites is scarce, some works show that their mechanical and physical properties remain practically unchanged and can even be enhanced, due to the new generated surfaces after the recycling process. This justifies the investment in their recovery and reutilization [34,35].

Hence, by focusing on green polymers, research efforts should be directed to the characterization of the pros and cons of the different materials, as well as on their ability to provide good interface performance. It is authors’ conviction that this is the pathway to the enhance sustainability of the composites industry and to cope with the new trends in recyclability and reuse in circular economy [36].

This paper is driven by this conviction and aims to produce a green composite from renewable materials (acrylate epoxidase soybean oil, as resin, and sisal and jute as reinforcement materials), and to investigate its chemical, physical and mechanical performances, as well as its recyclability and reuse as a second-life composite. The recovering of reinforcement and resin materials was performed at room temperature and atmospheric pressure and these resources were reused in the manufacturing of a second-life composite, whose properties were also investigated.

Jute fabric and sisal particles were chosen for the reinforcement and filler because they are natural materials with good mechanical properties. Besides their biodegradability and low price, they offer...
significant processing advantages, such as non-abrasive and high specific strength, due to their high lignocellulosic content [7,23]. Acrylated epoxidized soybean oil (AESO) was chosen as matrix because it is obtained from soybean vegetable oil, one of the most promising starting materials to produce thermosetting polymers, due to its high availability, low toxicity, sustainability and relatively low cost [27–29].

2. Experimentation

2.1. Materials and Methods

The work used a commercial acrylate epoxidase soybean oil (AESO), \( d = 1.04 \text{ g/mL at } 25 ^\circ \text{C} \) with inhibitor (4000 ppm monomethyl ether hydroquinone) as matrix material and tert-butyl peroxybenzoate (TBPB, 98%) as hardener, both supplied by Sigma-Aldrich. Reinforcement fibres of sisal rope, crushed to small particles (±1 mm), were used as filler. A dark brown biodegradable jute fabric produced by a binary system of simple interlacing—where a warp yarn is over or under a weft yarn at the crossover areas—was also used in the production of virgin and second-life composites. Both sisal rope and jute fabric were purchased from a local supplier and used with no further treatment in order to avoid increasing the overall manufacturing costs.

The resin specimens were prepared with different contents (0, 5 and 10 wt% based on AESO) of sisal particles (SP) added to the viscous liquid AESO. The particles were mechanically stirred at 40 °C for 5 min, to ensure uniform dispersion and the resulting mixture was subsequently subjected to the addition of different contents (2, 5 and 10 wt%) of hardener. After further stirring for 2 min, full homogeneity was achieved and the resulting mixtures were poured into an aluminium test specimen mould. The mould was manufactured on the Cincinnati Arrow VMC-500 CNC milling machine (Cincinnati, OH, USA), according to ISO D638-03 testing standards, as shown in Figure 2a.

The curing process was performed in an atmospheric pressure oven at 120 °C during 2 h, followed by an additional 4 h stage at 160 °C. After being cooled to room temperature, the test samples were removed from the moulds.

The different resin combinations led to the production of nine different samples that were identified with the acronyms HxSPx, where H and SP refer to hardener and filler and x represents the respective content (wt%). Figure 2b,c show the final visual aspect of the test specimen produced with AESO and 5 wt% of hardener with 0 wt% (H5SP0) and 10 wt% (H5SP10) SP, respectively.
Once the resin best performance conditions were determined (5 wt% hardener and 0 wt% or 10 wt% sisal particles as filler), green composite testing samples were produced via a wet layup method with five layers of bi-directional jute fabric as reinforcement (5 g per layer) and 50 g and 40 g of AESO for virgin and second-life composites, respectively. The production process was similar to the one described above.

After recycle assessment, two second-life composites were produced using recycled jute fibres and a third was produced using 40% of the recovered resin and 60% AESO. The same nomenclature was followed, and the samples were identified as CoHxSPxFR, where Co and FR refer to composite and recycled jute fabric (or RR recycled resin), respectively.

Figure 3 summarizes the methodology and different characterization tests that support the objectives of the work. Once the resin best performance conditions were determined (5 wt% hardener and 10 wt% sisal particles as filler or without filler), green composite testing samples were produced, and all tests were performed at least in triplicate.

**Figure 3.** Schematic diagram of research methodology for manufacturing a recyclable green composite.

### 2.2. Testing Procedures

#### 2.2.1. Chemical Characterization

Chemical characterization was performed by attenuated total reflectance Fourier transform infrared (ATR FT-IR) spectroscopy whose spectra were recorded in IRAffinity-1S Shimadzu apparatus (Kyoto, Japan), equipped with an ATR ZnSe Performance Crystal Plate accessory (4000–400 cm\(^{-1}\) range with 2.0 cm\(^{-1}\) resolution). Proton nuclear magnetic resonance (\(^1\)H NMR) spectrometry was performed on a Bruker AdvancedII+ Spectrometer (300 MHz) with 5mm probe (Rheinstetten, Germany), in CD\(_2\)Cl\(_2\) as solvent.

#### 2.2.2. Physical Characterization

Physical characterization consisted of water absorption tests that were carried out with samples (~19 × 24 × 2 mm) that were dried at 55 °C for 24 h to remove residual moisture that diffuse along the interfaces. Higher temperatures (>66 °C) were not used because they can affect the structure of the samples and promote microcracking [37]. Dry samples were weighed in a semi-analytical balance (±0.001 g) and immersed in containers with 40 mL of pure water. After 24 h, at room temperature,
the samples were removed, softly wiped with absorbent paper and weighed again. The water uptake rate (%) was calculated according to the following expression:

\[
W_{up} (%) = \frac{W_f - W_i}{W_i} \times 100
\]

where \( W_i \) and \( W_f \) are the initial and final weights of the samples, respectively.

2.2.3. Mechanical Characterization

Tensile and hardness tests were performed to characterize the mechanical properties of the composites. The tensile tests were carried out in a Shimadzu (Kyoto, Japan) AG IS 20 kN testing machine equipped with a 10 kN load cell and the hardness tests were done in a Zwick (Ulm, Germany) durometer that allowed measurements on the Shore A scale.

2.3. Recyclability

The recycling process was performed in rectangular cured AESO resin samples (~19 × 24 × 2 mm) using seven organic solvents, namely ethanol, 1-buthanol, acetone, methyl acetate, ethyl acetate, dichloromethane and chloroform. Samples were immersed in containers with 20 mL pure solvents with mechanical stirring at room temperature and dissolution and matrix breakdown were monitored over time (24 h). The same procedure was performed with selected solvents with composite samples.

3. Results and Discussion

3.1. Chemical Characterization

Sisal fibre, one of the most widely used natural fibres, is fully biodegradable and 65% of its composition is cellulose, with the remaining 35% composed of hemicellulose, lignin and waxes [38]. Cellulose is a polysaccharide consisting of a linear chain of over 500 \( \beta \)-1,4 linked d-glucose units, as presented in Figure 4. AESO is produced by the epoxidation of fatty acid double bonds followed by epoxy ring acrylation and its chemical structure is composed of fatty acid long aliphatic chains with various functional groups, such as the acrylate, epoxy and hydroxy, as schematized in Figure 4.

![Figure 4. The chemical structure of cellulose and acrylate epoxidized soybean oil.](image)

Chemical structures were confirmed by the ATR FT-IR spectra, which were recorded for initial materials (AESO and SP) and the nine exploratory samples produced (data not shown). This technique allowed us to identify the chemical functional groups and investigate the possible interactions and chemical bonding established between the AESO resin and SP fibres. Since the results were very similar for all the cured samples, a representative example is presented for the cured samples H5SP0 and H5SP10. Figure 5 shows the comparative analysis of the ATR FT-IR spectra of pure sisal particles, pure AESO and cured samples manufactured with 5 wt% hardener, H5SP0 (without SP) and H5SP10 (with 10 wt% SP), respectively.
In the spectrum of SP, as shown in Figure 5A the lignocellulosic matrix is recognised by the presence of a broad band at 3328 cm\(^{-1}\), characteristic of the stretching vibration of hydroxyl (–OH) groups and by the characteristic signals at, 2921 cm\(^{-1}\), 2850 cm\(^{-1}\) and 1018 cm\(^{-1}\) assigned to the stretching vibrations of C–H and C–O–C, respectively. In the pure AESO spectrum, as shown in Figure 5B, the –OH stretch was shifted to 3543 cm\(^{-1}\) and the characteristic ester (C=O) and (C–O) stretching bands were observed at 1723 cm\(^{-1}\) and 1265 cm\(^{-1}\), respectively. The characteristic C–H stretching bands of the CH\(_2\) and CH\(_3\) groups of saturated chains were observed at 2918 and 2848 cm\(^{-1}\) and the C–H bending band was identified at 1458 cm\(^{-1}\). Moreover, the bands at 1186 cm\(^{-1}\) and 1046 cm\(^{-1}\) can be identified due to the C–O–C stretching vibration of ester and also at 809 cm\(^{-1}\) in the epoxy band. The two weak bands at 1636 and 1617 cm\(^{-1}\), characteristic of the C=C stretch, disappear, as expected, after the curing process, as shown in Figure 5C,D. Comparative analysis between the spectra of cured AESO and H5SP10 allows us to conclude that they follow the same trend, with a slight change in the peak at 1031 cm\(^{-1}\) due to the cumulative effect of the sisal particles. This similarity shows that the added SP has no covalent bonds with AESO, as also reported by Liu et al. [28] with microcrystalline cellulose.

AESO structural characterization after the curing and recycling processes were performed by \(^1\)H-NMR. Spectra of origin AESO, cured AESO and AESO recovery from the dissolution process, obtained in CD\(_2\)Cl\(_2\), are presented in Figure 6.

For pure AESO, as shown in Figure 6A, the spectrum analysis allows us to identify the signals for the different protons in the expected areas. In the region between 0.75 and 1.00 ppm (4), the signal
corresponding to the terminal CH$_3$ groups of the chains is identified and the CH$_2$ groups, in $\alpha$ position relative to the carbonyl group (C=O), are identified by the presence of the triplet at 2.3 ppm (3). The three signals in the range between 5.75 and 6.50 ppm are attributed to the presence of the acrylate groups (5) and the central protons of the structure (1 and 2) are attributed to the signals between 4.10 and 4.50 ppm and 5, 2.5 and 5.50 ppm, respectively. The spectra of cured AESO, as shown in Figure 6B, and recovery AESO from the recycling process, as shown in Figure 6C, present the same trend, showing that no structural changes occurred during the recycling process. The absence of the signals of acrylate groups (5) in both cured AESO samples, as shown in Figure 6B,C, confirms the ATR FT–IR results.

3.2. Physical Characterization

Water absorption rates for the specimens produced with AESO resin and different hardener (2, 5 and 10 wt%) and SP (0, 5 and 10 wt%) contents were evaluated after 24 h submersion in water, as shown in Figure 7. The results followed a common trend with the incorporation of SP producing an increase in the amount of water uptake, reaching values between 2.0–2.7 wt% and 3.9–4.8 wt% for 5 and 10 wt% SP, respectively.

![Figure 7](image)

**Figure 7.** Effect of sisal fibre and hardener (refer to the percentage of hardener inside each column) on water absorption rate.

These results cope with the predictable behaviour of the green composite, due to the hydrophilic nature of the cellulosic filler, which contains a large number of hydroxyl (OH) groups that highly interact with water molecules. For the unfilled AESO samples (0% SP), the amount of hardener does not significantly influence the water uptake percentages with values of 0.27, 0.29 and 0.31%, for 2, 5 and 10 wt% hardener, respectively. In accordance with the report by Liu et al. [28], AESO contains, in their molecular chain structure, polar hydroxy and epoxy groups which, due to its hydrophilicity, are responsible for the inherent water absorption rate of unreinforced samples.

The out-of-trend results obtained for 10 wt% hardener are attributed to the curing process that influences the overall homogeneity of the specimens—namely the dispersion of the sisal particles (heavier sisal particles are deposited on the bottom of the mould). Resin contraction variations may also induce small voids and cracks in the specimens that will act as water collectors.

3.3. Mechanical Characterization

Figure 8 shows the results obtained for the hardness and tensile tests performed on nine AESO resin specimens manufactured with 2, 5 and 10 wt% hardener and 0, 5 and 10 wt% sisal particles.
which allows for the production of second-life composites, the preliminary test dissolution behaviour were excluded, despite presenting good results because, after curing, they gave rise to the deposit of fibres at the bottom of the mould and to some variation in the experimental results.

From all the organic solvents tested, those that belong to the halogenated family proved to be the most effective to dissolve the AESO resin matrix both before and after the curing process, followed by the AESO resin particles after 30 min, while acetone achieved a similar result after 2 h 30 min. The different chemical categories: alcohol; ketone; ester; halogenated.

After concluding the physical, chemical and mechanical tests, the compositions with 5 wt% hardener +0% sisal particles, H5SP0, and 5 wt% hardener +10 wt% sisal particles, H5SP10, were selected for the preparation of the virgin composites as well as for the investigation into recyclability and second-life production and performance. Specimens with 5 wt% fibre particles and 2 or 5 wt% hardener were excluded, despite presenting good results because, after curing, they gave rise to the deposit of fibres at the bottom of the mould and to some variation in the experimental results.

### 3.4. Recyclability

Since one of the objectives of the present work is the implementation of a recycling protocol, which allows for the production of second-life composites, the preliminary test dissolution behaviour of cured AESO resins were performed with solvents from different chemical categories: alcohol; ketone; ester; halogenated.

From all the organic solvents tested, those that belong to the halogenated family proved to be the most effective to dissolve the AESO resin matrix both before and after the curing process, followed by...
acetone. Alcohols and esters proved to dissolve the AESO resin before the curing process but, after curing, its dissolution was very slow (alcohols) or did not even occur (esters). Regarding the virgin composites, dichloromethane was able to separate the entire five layers of jute and recover the AESO resin particles after 30 min, while acetone achieved a similar result after 2 h 30 min. The results are shown in Figure 9.

![Figure 9](image-url) Dissolution process of virgin composite in dichloromethane or acetone. Separation of recovery jute samples and AESO polymer.

However, despite the higher dissolution rate of dichloromethane over acetone, the latter was selected due to its better performance according to the safety, health and environmental (SHE) criteria, as shown in Table 1.

| Family      | Solvent    | BP (°C) | FP (°C) | Safety Score | Health Score | Env. Score | Ranking after Discussion |
|-------------|------------|---------|---------|--------------|--------------|------------|--------------------------|
| Alcohol     | Ethanol    | 78      | 13      | 4            | 3            | 3          | Recommended              |
|             | 1-Butanol  | 118     | 29      | 3            | 4            | 3          | Recommended              |
| Ketones     | Acetone    | 56      | −18     | 5            | 3            | 5          | Recommended              |
| Ester       | Methyl Acetate | 57   | −10     | 5            | 3            | 5          | Problematic              |
|             | Ethyl Acetate | 77   | −4      | 5            | 3            | 3          | Recommended              |
| Halogenated | Dichloromethane | 40  | −       | 1            | 7            | 7          | Hazardous                |
|             | Chloroform | 61      | −       | 2            | 7            | 5          | Highly hazardous          |

BP: boiling point, FP: flash point. Higher numbers correspond to higher hazard or toxicity for safety, health and environmental hazards.

After the dissolution process, the structural integrity of the recycled/recovered AESO resin was accessed by proton NMR, as shown in see Figure 6C, and compared with the original sample, showing that no structural changes occurred during the recycling process. The recycled resin particles, as well as the recovered jute fabric, were used to produce second-life composites and their performances were evaluated together with the previous virgin composites.

3.5. Virgin and Second-Life Composites

Virgin and second-life composites were produced and their performance evaluated after selecting the resins and establishing the chemical recycling procedure.

3.5.1. Physical Characterization

Water absorption was analysed for both virgin (CoH5SP0 and CoH5SP10) and second-life composites (CoH5SP0FR, CoH5SP10FR and CoH5SP10RR). Compared to the respective resin samples, which provided a maximum absorption of 4.2 wt% for the H5SP10 sample, as shown in Figure 7 in the first part of the paper, the incorporation of the jute reinforcement in the new virgin and second-life composites resulted in a significant increase in water absorption, reaching values of 11.3% in the
The high hydrophilicity characteristics of the jute fabric, which allows for absorbing almost 110% of its weight and may even induce more voids to entrap more water, is responsible for these results.

The corresponding second-life composites produced with the recovered jute fabric (CoH5SP0FR and CoH5SP10FR) presented lower water absorption (less 56–62%) than the virgin composites (CoH5SP0 and CoH5SP10) but the overall evolution trend is similar because water absorption resistance diminishes with the addition of sisal particles. A possible explanation for the enhanced water absorption resistance of the second-life composites may be related with the saturation of the jute fabric exposed to the recycling process, and to a better uniform adhesion of the new matrix.

The second-life composite, CoH5SP10RR, produced with recovered resin particles, absorbs less than 22% of the water than the original virgin composites, but approximately 80% more water than the other second-life composites.

The abovementioned results show that the production of second-life composites helps in the reduction in water absorption, which is commonly pointed out as a negative feature of green composites. They also allow for the conclusion that the variation in the properties of the recycled resin will have less impact on water absorption when compared to second-life composites manufactured without recycled resin.

### 3.5.2. Mechanical Characterization

Figure 11 shows the hardness and tensile strengths of virgin and second-life composites fabricated with 5 wt% hardener, and 0% or 10 wt% sisal particles. A first comparison of the hardness measurements obtained with virgin composites CoH5SP0/10, as shown in Figure 11a, and resin test specimens H5SP0/10, as shown in Figure 8a, allows for the conclusion that the results are similar with a positive influence of sisal particles on the final hardness of the composites.

A similar comparison for the tensile strength values, as shown in Figure 11b, provides different conclusions. Firstly, the overall levels of tensile strength are significantly higher for the composites because the resin test specimens were produced without jute fabric. Secondly, the combined presence of jute fabric and sisal particles improves the tensile strength of the composites. This last conclusion is opposite to that obtained from the resin test specimens, probably because the heterogeneity caused by the presence of sisal particles dispersed in the resin matrix is only mechanically effective in the presence of jute fabric.

Second-life composites are more difficult to analyze because the results obtained with CoH5SP10FR seem to be out-of-trend against those obtained with CoH5SP0FR and against what was said above. A possible explanation for the out-of-trend results of CoH5SP10FR and for presenting a tensile strength 50% lower than that of the virgin composite CoH5SP10 may be related to the non-uniform distribution of sisal particles. In fact, the recycling of the jute fabric always keeps some resin residues that constrain...
sisal particles to penetrate the core of the specimens, thereby increasing its concentration at the surface. Hence, hardness increases slightly more than expected and tensile strength decreases to values close to the resin test specimens, because cracks are triggered at the centre of the specimens during tensile tests.

![Graph](image)

**Figure 11.** Mechanical tests for virgin and second-life composites: (a) Hardness; (b) Tensile strength.

Regarding the composite manufactured with reused resin (COE5F10RR), it presents the highest mechanical strength, probably because the reused resin already contains small sisal particles from previous utilization, which enables for the attainment of a composite with a uniform distribution of sisal particles throughout the thickness of the specimens.

### 4. Conclusions

This work allowed for the conclusion that is possible to manufacture green composite with renewable materials and that it is also possible to reuse the materials for a second-life composite. The experiments made use of jute fabric as reinforcement, acrylate epoxidase soybean oil (AESO) as matrix and sisal particles (SP) as filler. The recycling process was performed with acetone, an organic solvent recommend by the SHE criteria. The best mechanical performance (hardness of 91 Shore A and tensile strength of 10.6 MPa) of the virgin green composites was obtained with 10 wt% sisal particles and 5 wt% hardener. The second-life composites generally present a slight decrease in mechanical properties but have a higher resistance to water absorption, which is often considered the main problem associated with the use of green composites.

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