



a 0.3 mm sieve. The sieved polymer was dried in the oven at 120°C for 4 h and set aside. The prepared fibres and matrices were blended in the proportions of 45% Kenaf, 5% carbon, 3% polyester, 45% PP, and 2% maleic anhydride additive. The materials were blended using a Brabender blending machine run for 10 min at 170°C and 50 rpm. Afterwards, the compounds were powdered into 0.05 mm particles by cryogenic grinding. Several batches of the prepared powder mixture were set aside to produce 15 samples (15 cm x 15 cm).

### Composite compression moulding sample preparation

A mould was precoated with a non-stick mould release agent, and the machine was preheated to 160°C (Limited, 2005). The powdered compound was then moulded carefully into the compartment, preheated for 7 min, pressurised under 50 MPa for 10 min, cooled for 3 min, and then finally removed from the machine to allow cooling at room temperature for 24 h.

### Property measurements for compressed moulded composites

Adherence to specific standards and associated parameters was required during the testing proper. American Society for Testing and Materials (ASTM) standards were followed in the current study. The prepared composites were removed from the mould and cut according to ASTM D638 for tensile strength testing and ASTM D3846 for shear strength testing. In total, three to four specimens were cut from three composite plates for various mechanical tests and measurements. For each test, 10 samples were provided and measured. The measurements were conducted in two directions of longitudinal (0°) and transverse (90°) orientation. The Archimedes method, in which deionised water is used as the immersion and infiltration liquid, was applied to measure

the bulk density of each test specimen (Table 1).

### Results and Discussion

The failure mode in the tensile tests occurred through fibre and matrix breakage. Fibre pull-out was also observed in the present study. When failure of the specimen occurred, the test was stopped and the related data were obtained (Figures 1 and 2).

The specific Young's modulus and specific strength of the composites constituted the comparison criteria, while the critical factor for the composites was the tensile property. As shown in Table 2, the specific strength of the biocomposite is greater than the reported strength of Kenaf/PP fibre composites obtained by previous researchers. Such an improvement is attributed to the incorporation of carbon fibres into the composite.

The figures illustrate that the composite exhibited properties in good agreement with common structural materials, such as carbon fibre-reinforced plastic, and even better than those of Kenaf/PP composites (Figure 3A-3C). The results also show satisfactory properties compared with concrete as a general structural material. The composite can produce curvilinear forms without imposing extremely high costs. Hence, it can be claimed that the composite can be used for a wide range of structural purposes provided that the total distributed stress and strain loads do not exceed its design allowances.

### Conclusion

In the current study, we aimed to evaluate the mechanical properties of Kenaf/carbon/PP fibre composites fabricated from randomly scattered fibres.

Property	Description	Value Gr/cm <sup>3</sup>	Standard deviation	Coefficient of variation	Design allowable
	Density	1.14	----		----
E <sub>11</sub>	Longitudinal Young's modulus (GPa)	30.2	8.28	0.27	10.7
E <sub>22</sub>	Transverse Young's mModulus (GPa)	6.9	0.45	0.065	5.84
X <sub>1</sub>	Longitudinal tensile strength (MPa)	72.58	16.22	0.22	34.38
Y <sub>1</sub>	Transverse tensile strength (MPa)	40.1	11.34	0.28	13.39
G12	Longitudinal in-plane shear modulus (GPa)	9.92	1.19	0.12	7.11
G23	Transverse in-plane shear modulus (GPa)	8.46	6.8	0.807	0.69
Sc	Shear strength (Mpa)	40	4.1	0.1025	30.34

Table 1: Mechanical properties of the composite material.



Figure 1: A) Failure mode of tensile strength test and B) ultimate tensile strength.

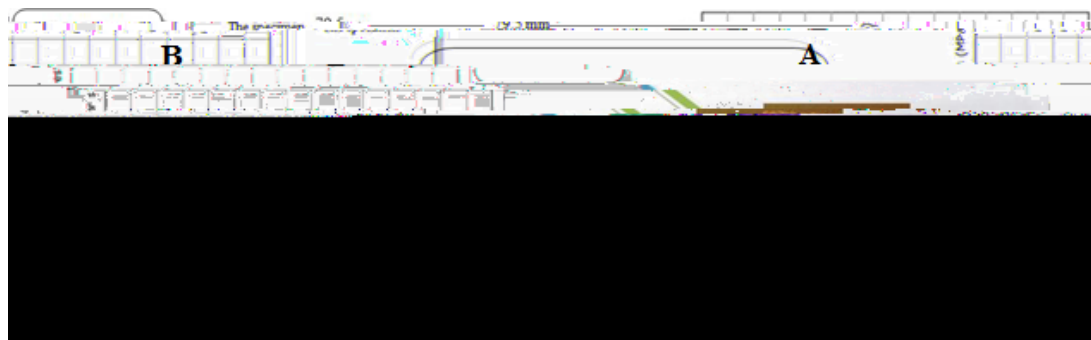


Figure 2: A) In plane-shear property failure and B) Shear stress-strain behaviour (up to 5% strain).

Description	Kenaf/PP 50/50	Carbon/PP 50/50	K/C/PP 50/50
Tensile strength (MPa)	62	2495	72.58
Tensile modulus (GPa)	7.7	125-150	30.2
Flexural modulus (MPa)	3.6	37.92	19.88
Shear strength (MPa)	6.36	310	40
Density (Kg/m <sup>3</sup> )	1400–1500	1070	1140
Cost \$/kg	6	7.5	6.2

Table 2: Comparison of the properties of Kenaf/PP and carbon/PP composites.

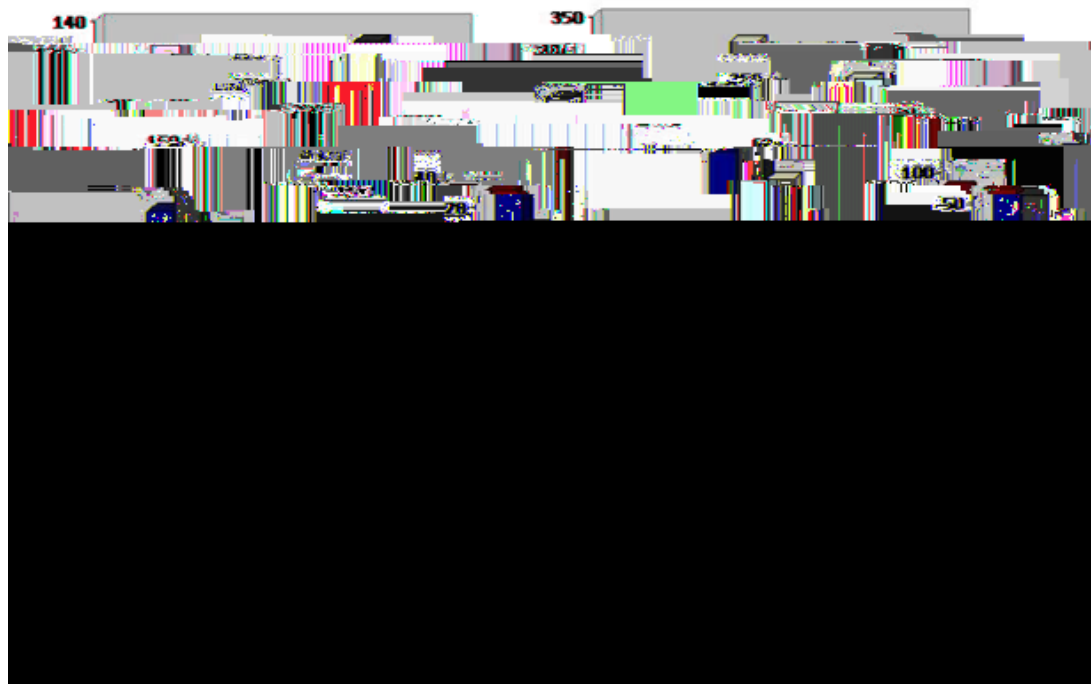


Figure 3: A complex, multi-layered image showing various components and structures, possibly related to the manufacturing process or the composite materials.

First, the process (heating and compressing) was performed well below 180°C to avoid fibre degradation. Control of the moulding process (time, temperature, and press) was crucial to reduce fibre damage. During determination of the processing parameters that needed to be adjusted to each thermoplastic polymer, the rheological and thermal properties of neat polymers must be considered. The second part of the study focused on the mechanical behaviour of the polymeric matrix composites reinforced by the hybrid fibres. With volume fractions of

45% Kenaf and 5% carbon fibres, satisfactory performance levels were obtained for the Kenaf/carbon/PP fibre biocomposites.

The substitution of PP by Kenaf/carbon/PP fibre biocomposites for RM may yield a stronger product. The results are very encouraging for the development of biocomposites for structural applications. Complementary studies, such as the application of the material under an RM procedure and adoption of different fibre designs and matrices, will be conducted in the future. The standard deviation and allowable

