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On-site evaluation of a modified pultrusion process: Fibre spreading and resin injection-based impregnation



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Abstract

The focus of the current research was to demonstrate a modified pultrusion technique at an industrial site using a commercial production machine. In this instance, the conventional resin bath was replaced by a custom-designed compact impregnation unit. The dimensions of the impregnator were $310 \text{ mm} \times 400 \text{ mm} \times 9 \text{ mm}$. The relatively short length of the impregnator, when compared to a 5 litre resin bath, meant that it had to be efficient in impregnating the reinforcing fibre bundles. This was achieved using a fibre spreading unit and a facility to inject the resin under low-pressure. The design basis for the fibre spreading unit and the impregnator are presented along with the methodology that was used to select the pultrusion speed and die temperature. The pultrusion experiments were performed using filled and unfilled vinyl ester resin and E-glass fibres. The profile of the 0.5 metre-long die was rectangular with dimensions of $32 \text{ mm} \times 2.2 \text{ mm}$. The fibre spreading unit and the impregnator were retrofitted to the pultrusion line with ease. The physical, mechanical and thermo-mechanical properties were determined for the pultruded composites using the resin bath and the modified technique. The properties of the latter were found to be equivalent or marginally superior when compared to the resin bath-based production method. However, in the modified pultrusion technique, when compared to clean the equipment after production was reduced by 90%.

Keywords

Pultrusion, glass fibres, polymer-matrix composites, impregnation, mechanical properties

Introduction

Pultrusion is an established manufacturing technique that is used to produce continuous fibre reinforced composite profiles of a constant cross-section. Examples of typical pultruded profiles include rods, hollow tubes, flat sheets and beams. The term "fibre spreading" is used here in the context of spreading the filaments in a bundle prior to impregnation.¹ Fibre spreading results in a decrease in the thickness of the bundle with a concomitant increase in the width.² This enhances the through-thickness impregnation rate.³ A schematic illustration of a typical resin bath-based pultrusion production process is shown in Figure 1(a) and a magnified view of the resin bath is presented in Figure 1(b).

The following section presents a brief discussion on the key components highlighted in Figure 1(a) and (b). The creels or spools of fibre-(i) are stacked on a creel stand-(ii) and the individual bundles are unwound from the bore (this is generally referred to as centre-pull payout). The individual bundles are guided through a series of fibre guides-(iii) to configure them spatially to the required trajectory. The arrays of fibres-(iv) are fed through a further set of guides-(v) into the resin bath-(vi) via a "plunger"-(vii). The plunger permits the array of fibres to be lowered or raised into a resin bath-(vi) where they soak up the resin system-(vii) when immersed. The fibres at this stage are effectively over-

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Figure 1. (a) Schematic illustration of a conventional resin bath-based pultrusion process. The presence of reinforcing mats that are fed into the resin bath (to enhance the transverse properties of the composite) is not shown. (b) Magnified view of the resin bath.

impregnated. The impregnated fibre bundles-(ix) are directed to sets of pre-die guides-(x) that resemble the geometry of the die entrance but with slightly larger dimensions. A fair volume of the excess resin on the fibre bundles is squeezed out as they traverse these predie guides and enter the die. The resin that is squeezed out at the pre-die guides-(x) and the entrance to the die-(xii) is channelled back to the resin bath via a drip-tray-(xi). The impregnated fibres are directed into a split die-(xiii) with surface-mounted external heaters-(xiv). The external heaters are used to create a temperature profile along the length of the die. The impregnated fibre bundles are cross-linked or cured within the heated die and the composite-(xv) with the profile of the die is created. The solid composite profile is hauled off-(xvi) and sectioned to the required length using a mechanical cutter-(xvii).

According to a report by Lucintel, the global composite materials market is predicted to reach U\$41.8 billion by 2025. The global end-product sector is estimated to attain U\$116.6 billion by 2025.⁴ Considering the global pultrusion market, it is predicted to reach U\$3 billion by 2023.⁵ The predicted growth for another manufacturing process that involves the impregnation of continuous fibres is filament winding. Two key products manufactured using filament winding are pipes and pressure vessels. The composites pipe market is forecast to expand at a compound annual growth rate of 3.2% from 2019 to 2024 and it is predicted to reach U\$4,363 million by 2024.6 According to a report by Zion Market Research, the composite liquid petroleum gas cylinder market itself was stated to be worth U\$131 million in 2018 and it is forecast to grow to U\$249 million by 2025.⁷ With reference to pultrusion, technologies that can potentially have a positive impact on the market share include. (a) innovations to enable an increase in the pultrusion rate, (b) integrated sensor systems for on-line process monitoring and structural integrity assessment and (c) fast-curable resins. Since the patent by Goldsworthy in 1959,8 the continued economic importance of pultrusion over the last three decades has resulted in significant attention being paid to modification^{9–16} and process optimisation via modelling.9,17-31

The current authors reported previously on a modified pultrusion process³² where the resin bath was replaced with a purpose-designed low-pressure resin impregnation unit. In summary, with reference to the mode of operation of the modified pultrusion technique, the resin and hardener are stored separately and pumped on demand via a pair of precision gear pumps to a static mixer. The throughput volume from each pump is pre-calibrated to deliver the required stoichiometric ratio for the resin system. The mixed resin system from the static mixer is directed to the resin impregnation unit where the partially spread bundles are impregnated. The positions of the custom-designed fibre spreading and impregnation units are shown in Figure 2. The current paper reports on the evaluation of this modified pultrusion technique on a commercial pultrusion machine at an industrial site using a filled resin system.

The focus of the research was four-fold. Firstly, it was to demonstrate that the modified pultrusion equipment could be retrofitted on an industrial pultrusion machine without requiring any modification to the original machinery. Secondly, since unfilled resins were used in the laboratory trials,³² it was necessary to demonstrate that filled resins could be used with this modified pultrusion technique in an industrial setting. Thirdly, it was necessary to demonstrate that the physical and mechanical properties of composites manufactured using the modified pultrusion technique did not result in any degradation of the physical, mechanical and thermo-mechanical properties. Finally, data were

gathered on-site to verify the "green credentials" of the modified pultrusion method when compared to the conventional resin bath-based production technique.

Description of the modified low-pressure impregnation pultrusion process

When compared to the conventional pultrusion technique (see Figure 1(a)), items (i) to (v) and (xii) to (xvii) in the modified method (as illustrated in Figure 2(a)) remain the same. In the modified pultrusion method, the resin bath shown in Figure 1 is replaced with a compact fibre spreading unit and a low-pressure resin impregnator or a pressure-pot (containing the mixed resin and hardener - not shown in Figure 2).

With reference to Figure 2(b), the arrays of fibre bundles are guided through eyelets to the fibre spreading unit-(a). The primary function of the fibre spreading unit is to initiate partial de-cohesion of the binder and to enable the lateral spreading of the filaments with a concomitant decrease in the thickness.^{1,2} The details



Figure 2. (a) Schematic illustration of the modified low-pressure injection pultrusion technique. (b) Magnified view of the resin impregnator.

of this unit were presented previously.³² The array of fibres are directed under the resin impregnation unit-(b) where the partially spread fibres are impregnated at the resin delivery head-(c). In the modified pultrusion technique, the drip tray that is used with conventional pultrusion is replaced by an inclined and integrated platform that serves as the resin-drip tray and the fibre compaction zone-(d), before the fibres are directed to the heated die. The integrated platform is inclined to permit the resin that is squeezed out from the bundles at the fibre pre-guide-(e) to flow back under gravity to the impregnation zone. In effect, this serves as a miniature reservoir and it aids resin pick-up by the fibre bundles below the impregnator. Hence, under ideal processing conditions, the dripping of resin from the impregnated fibre bundles as they enter the die is minimised. The impregnated and aligned fibre bundles enter the die and the items (xii-xvii) mentioned previously are identical. However, in this instance, a feedback control system-(f) is used to synchronise the haul-off unit to the pumping speed of two precision gear pumps-(g) where each independently services a reservoir containing the resin-(h) and the hardener-(h'). The two pumps are controlled independently by a master control unit-(g'). Since the pumps serving the resin and hardener reservoir are controlled independently, it is possible to set the required stoichiometry and the throughput rates. The feedback control system ensures that the pumping of the resin and hardener is synchronised to the haul-of unit. The resin and hardener are pumped on demand to a manifold-(j) and the two streams are fed to a static mixer-(k). The mixed resin system is supplied to the resin delivery head-(c).

The fibre spreading unit and the resin impregnator were designed to fulfil the following requirements.

Fibre spreading unit: This unit was designed to be integrated into a pultrusion line where the fibre bundles are spread partially by passing them in a serpentine manner through a series of rollers. This had the effect of partially breaking up the binder and permitting the lateral spreading of the filaments in the bundle; at the same time, the thickness of each bundle is reduced. This enables the through-thickness impregnation of the resin to be accelerated.³ The fibre spreading unit also served

to present a ribbon of fibre bundles, with minimal segmentation, to the resin impregnation unit. This was important because the impregnator was designed to be operated with a ribbon of fibre bundles.

Impregnation unit: The main functions of the resin delivery-head were to: (a) accelerate the throughthickness impregnation of the fibre bundles by enabling further spreading; (b) offer control over the volume of resin that is deposited on the spread fibres via a resin dispensing unit; (c) enable a number of modes of impregnation to be enacted (pin, injection, compaction and capillary-impregnation); (d) facilitate easy threading of the fibre bundles; (e) enable easy dismantling and cleaning; (f) ensure minimal dead-space to prevent the mixed resin and hardener from stagnating; and (g) facilitate the application of minimal tension on the bundles prior to them entering the heated die. The resin impregnation unit was designed to accommodate thirty 4800 tex or sixty 2400 tex E-glass fibre bundles. The number of rovings was selected on the basis of the dimensions of the profile of the available die $(32 \text{ mm} \times 2.2 \text{ mm})$.

With reference to Figures 2 and 3, a ribbon of the spread rovings from the fibre spreading unit is directed to the resin-applicator unit. The resin-applicator was designed to inject the resin directly into the spread fibre bundles at the same time facilitating the impregnation via pin-impregnation. From this point, the trajectory of the fibre bundles is changed as the partially impregnated fibres are passed over a pin and under two further pins. The fibre arrays are then converged in the "converging section" of the impregnation unit. The resin that was squeezed out from the fibres within the converging section and the pre-former profile flows back towards the resin-applicator. This was possible because this section of the impregnation unit was inclined at 5°. This accumulation of resin represents a miniature reservoir. With reference to Figure 3, the rig was designed to enable the positions of pins-(i) and (ii) to be secured within the vertical channel. Additional pins can be introduced at pins-(iii) and (iv) to provide more contact points for the fibre bundles. The position of the impregnator head-(v) could also be adjusted. As stated previously, the fibre preform guide was integrated and secured within the impregnation rig.



Figure 3. Schematic illustration of the impregnator showing generalised impregnation modes at specified regions.

Design calculations for the impregnation unit

The theoretical basis for the design of the impregnator unit, including an assessment of the different modes of impregnation as a function of specified processing parameters, was detailed in reference.³² Hence, only a summary is presented here. The modes of impregnation at the zones indicated in Figure 3 are summarised in Table 1 along with the relevant equations that were used to calculate the degree of impregnation.³²

The degree of impregnation (I_{imp}) was determined using the following relationship:

$$I_{imp} = \frac{\left(I_{inj} + I_{pin} + I_{cap} + I_{comp}\right)}{100} \tag{1}$$

where

 I_{inj} = impregnation via injection pressure; I_{pin} = pin-base impregnation; I_{cap} = capillary impregnation; and I_{comp} = compaction-based impregnation.

In Figure 4, the histogram shows the individual contribution of specific modes of impregnation (capillary, compaction, pin, injection and the total (a sum of the four modes of impregnation considered)). The scatterplot above the histogram represents the observed trend for the total modes of impregnation. The values above "1" (unity) imply that the fibre bundles are overimpregnated. It can be seen from Figure 4 that the degree of impregnation for the impregnator designed in this study (shown in Figure 3) is greater than unity for the three pultrusion speeds considered. This implies that complete impregnation of the fibres should take place at the pultrusion speeds considered. A decreasing trend for the degree of impregnation with an increase in the pultrusion speed is apparent. The data required for estimating the degree of impregnation I_{imp} are summarised in Tables 2 and 3.

Materials, selection of pultrusion parameters and test methods

Resin and fibres

The details of the vinyl ester resin formulation used in the pultrusion experiments are presented in Table 4. The E-glass fibres (386 Direct Roving, 4800 tex) with an average fibre diameter of 24 μ m were used in this study.

Selection of the die-temperature and pultrusion speeds

A power-compensated differential scanning calorimeter (Diamond DSC, Perkin Elmer, UK) was used to study the cross-linking behaviour of the resin system. This was required to enable the selection of suitable temperatures for the die and to define the pultrusion speeds. The procedures reported in reference 42 were adopted. The conversion-temperature curve for the unfilled vinyl ester resin is shown in Figure 5. It can be seen from Figure 5 that at 140°C, the conversion of the reaction is nearly 90% complete. Hence, this temperature was selected as the die-temperature.

In the second set of experiments, the resin system was cross-linked isothermally at 140°C. The objective of this experiment was to establish the pultrusion speed which would ensure the complete cross-linking of the resin within the 0.5 m long die. From an isothermal experiment that was conducted at 140°C, a conversion curve as a function of time was generated as shown in Figure 5. A pultrusion speed of 0.4 m minute⁻¹ was chosen for the experiments because the degree of conversion at 75-seconds (corresponding to a pultrusion speed of 0.4 m minute⁻¹) for the 0.5 m die was approximately 90%. The average glass transition temperature (T_{σ}) from three experiments for the unfilled vinyl ester resin was 117.9 ± 1.8 °C. The T_g was measured at the mid-point of the change in the heat capacity trace in accordance with ASTM E1356-08.43 The heating rate used for the dynamic heating experiments in the DSC was 5 K/minute.

Pultrusion experiments

The pultrusion machine used in the industrial trial was designed and built by Pultrex Ltd (Model PX500-6T) with a maximum pulling force of 6,000 kg for each of the two grippers. It was equipped with reciprocating pullers and a programmable automatic cutting device. The cross-section of the die profile used in this study was $32 \text{ mm} \times 2.2 \text{ mm}$. The conventional pultrusion experiments were performed with a dip-type resin bath with a capacity of five litres. The details of the resin systems, fibre reinforcements, die temperatures and pultrusion speeds are provided in Table 5. At the end of the pultrusion experiments, the resin that remained in the bath was transferred into small containers and cross-linked prior to disposal. The components that came into contact with the mixed resin (resin bath, profile guides, plunger and feed-back tray) were removed at the end of the trial and cleaned using acetone. The resin impregnator, designated as "PUL-I" was used for the modified pultrusion experiments. The resin system was premixed and transferred to a two-litre pressure-pot that was operated using compressed air. The details of the pultrusion experiments using the modified technique are provided in Table 5.

Impregnation zones in Figure 3	Mode of impregnation	Relevant equations	Ref	Calculations for each zone at V $=$ 0.3 m/min	Comment
1, 2, 3, 5	Pin	$I_{pin} = \frac{Z_{pin}}{Z} \times 100 \text{ equation (A-1)}$ $Z_{pin} = \sqrt{\frac{2I_{pin}KP_{pin}}{\eta}} = \frac{\sqrt{144W^4K^2 + 12KW^2h_o^4 + 12W^2K}}{h_o^3}$	32 41	$\begin{array}{l} \textbf{4.99}+\textbf{2.5}+\textbf{2.5}+\textbf{2.5}=\textbf{12.49\%}\\ \textbf{12.48}+\textbf{6.23}+\textbf{6.23}\\ +\textbf{6.23}=\textbf{31.5}\ \mu\text{m} \end{array}$	The extent of impregnation via pin-impregna- tion was expressed via an impregnation number (I_{pin}). The depth of the fibre thick- ness that is impregnated via this mode (Z_{pin})
		equation (A-2) $h_{o} = \left(\frac{\pi^{2}}{128}\right)^{1/3} R \left(\frac{6\eta V}{T_{o}}\right)^{2/3}$	35	12.47 μm	was estimated from equation A-2 from Wang et al. ^{33,34,41} The parameters employed in equation A-2 were calculated using equa- tions A-3 and A-4.
		equation (A-3) $K = \frac{16}{9\pi\sqrt{6}} \left(\sqrt{\frac{V_{fmax}}{V_f}} - I \right)^{5/2} R^2$	37	$\textbf{2.18}\times \textbf{10}^{-13}~m^2$	
I	Injection	equation (A-4) $I_{inj} = rac{Z_{inj}}{Z} imes$ 100 equation (B-1)	32	12.41%	The depth of the impregnation in the roving (7) are a result of injection based impress
		$Z_{inj} = \sqrt{rac{2 t_{inj} KP_{inj}}{\eta}}$ equation (B-2)	3	31.03 μm	nation was estimated from the Darcy's
	Coolillana	$t_{inj} = rac{L_{inj}}{V}$ equation (B-3)	3	0.08 s	equation. The P _{inj} is the pressure used to inject the resin onto the spread rovings.
4,6	Capillary	$I_{capillary} = \frac{Z_c}{Z} \times 100$ equation (C-1)	32	100.78%	capillary-impregnation was estimated by
		$Z_{c}=\sqrt{rac{2t_{c}KP_{c}}{\eta}}$ equation (C-2)	3	252 μm	eters to be used were estimated from Ahn
		$t_{c} = \frac{L_{c}}{V}$ equation (C-3)	3	22 s	
		$P_{c} = rac{4\sigma cos \theta}{D_{E}}$ equation (C-4)	36	3.3 kPa	
_	-	$D_E = \frac{8r_f}{F_Z} \frac{1 - V_f}{V_f}$ equation (C-5)	37 32	2.06×10^{-5} m	
7	Compaction	$I_{comp} = \frac{-comp}{Z} \times 100 \text{equation (D-1)}$	52	50.69%	The depth of the impregnation in the roving (Z_{comp}) as a result of impregnation due to
		$Z_{comp} = \sqrt{\frac{2t_{comp}KP_{comp}}{\eta}}$ equation (D-2)	3	Ι26.7 μm	compaction at the exit point of the resin
		$t_{comp} = \frac{L_{comp}}{V}$ equation (D-3)	32	0.1 s	of Darcy's equation. Here the pressure was selected from the compaction chambers of similar dimensions reported in the literature
l to 7	Total	$I_{imp} = \frac{(I_{inj} + I_{cap} + I_{comp})}{100}$	32	1.76	An impregnation number (l_{imp}) of I suggests complete impregnation of the fibre tow.
		$I_{imp} = \frac{12.49 + 12.41 + 100.78 + 50.69}{100}$			

Table I. A summary of the relevant modes of impregnation for impregnator shown in Figure 3.

The relevant terms in Equations A-I and D-3 are as follows. I_{pin} = impregnation number related to impregnation taking place via pin-impregnation mode, $I_{capillary}$ = impregnation number related to impregnation taking place via injection mode, $I_{capillary}$ = impregnation number related to impregnation number related to impregnation taking place via capillary mode, I_{camp} = impregnation number related to impregnation taking place via capillary mode, I_{camp} = thickness of the fibre bundle impregnated due to pin-impregnation mode, Z_{inj} = thickness of the fibre bundle impregnated due to capillary mode, Z_{camp} = thickness of the fibre bundle impregnated due to capillary mode, Z_{camp} = thickness of the fibre bundle impregnated due to capillary mode, Z_{camp} = thickness of the fibre bundle impregnated due to capillary mode, Z_{camp} = thickness of the fibre bundle impregnated due to capillary mode, Z_{camp} = thickness of the fibre bundle impregnated due to capillary mode, I_{camp} = time for which the fibre bundle experiences impregnation via capillary mode, t_{camp} = time for which the fibre bundle experiences impregnation via capillary mode, t_{comp} = time for which the fibre bundle experiences impregnation via capillary mode, t_{comp} = time for which the fibre bundle experiences impregnation via capillary mode, P_{camp} = pressure available for the resin impregnation via injection mode, P_{pin} = pressure available for the resin impregnation via compaction mode, L_{pin} = distance for which the fibre bundle experiences impregnation via capillary mode, L_{comp} = distance for which the fibre bundle experiences impregnation via capillary mode, K_{camp} = distance for which the fibre bundle experiences impregnation via compaction mode, K_{e} = transverse permeability coefficient, η = resin viscosity, W = width of the fibre bundle experiences impregnation via compaction mode, K_{e} = transverse permeability coefficient, η = resin viscosity, W = width

The bold numbers represent the final result of the calculations in this section.

Test methods

Density, fibre and void volume fraction: The densities of the pultruded composites were determined using an

OHAUS density measurement kit (Model: AP110S). The fibre volume fraction of the pultruded samples was determined by the ignition-loss method in accordance with ASTM D 2584-08.⁴⁴ The void fraction (V_v)



Figure 4. Simulation of the degree of impregnation as a function of the pultrusion speed.

Table 2.	Summary	of the	parameters	that	were	used	to	calcu-
late the e	xtent of ir	npregn	ation.					

Parameter	Units	Values
Viscosity of the resin (η)	Pa∙s	0.5 38
Fibre volume fraction (V_f)	%	70
Thickness of the roving (Z)	μ m	125
Width of the roving (W)	μ m	8
Radius of the fibre (r_f)	μ m	12
Pre-tension in the roving (T)	N	10
Maximum fibre packing (V_A)	%	90.7
Roving architecture	-	Hexagonal
Coefficient of transverse permeability (K)	m ²	2.18×10^{-13} 37

Table 3. Summary of the parameters that were used tocalculate the impregnation-zone length assumingcapillary-impregnation.

Parameter	Units	Value
Surface tension of the resin (ζ)	N/m	0.034 ³⁹
Contact angle between	Degrees	60 ⁴⁰
the fibre and the resin (°)		
Form factor (F)	_	2
Maximum fibre packing (V_A)	%	90.7
Fibre architecture	_	Hexagonal
Calculated values		
Wetted diameter (D_E)	μm	20.57
Transverse capillary pressure (P_c)	kPa	3.3

was determined in accordance with ASTM D 2734-16.⁴⁵ The total number of specimens used was 20.

Image analysis: The void fractions of the composites were calculated from micrographs using ImageJ image analysis processing software.⁴⁶ Standard metallographic techniques were used for potting and polishing the samples for subsequent inspection using an optical microscope (Zeiss Axioskop 2).

Thermo-mechanical properties: A dynamic mechanical analyser (DMA 242 C, NETZSCH) was used to obtain the thermo-mechanical properties of the neat resin system and the pultruded composites. Each sample was clamped in the machine using a dual-cantilever fixture and heated from 25°C to 180°C at 3 K minute⁻¹. The experiments were conducted at 1 Hz. A minimum of three samples were tested for each of the specimen types.

Inter-laminar shear and flexural properties: The interlaminar shear strength (ILSS) of the pultruded composites was measured in accordance with BS EN ISO 14130:1998 and the flexural strength and modulus of the composites were measured in compliance with ASTM D 6272-08. Five samples each were used for the ILSS and flexural tests.

Results and discussion

Site trial

The advantages and the future potential of the modified pultrusion method over the conventional resin bath-based technique observed during the site trial were as follows.

(i) Ease of retrofitting: The retrofitting of the impregnation unit was relatively straightforward as it involved securing the impregnation unit on the top of the empty resin bath. It did not require

Commercial name	Chemical family	Parts by weight
Atlac 430 (Palatal A430)	Vinyl ester resin	100
Styrene	Reactive solvent	2
Perkadox 16	Peroxide	I
t-butyl peroxybenzoate	Peroxide	0.5
Tert-Butyl peroxy-2-ethylhexanoate	Peroxide	0.5
INT-PUL-24	Mould release	I
Dolomite Microdol [®] H Extra	Talc filler	15

Table 4. Details of the filled and un-filled vinyl ester resin systems used in this study.



Figure 5. Combined plots show: (i) conversion versus temperature curve for the unfilled vinyl ester resin. The conversion curve was generated from a dynamic DSC experiment conducted at 5 K/minute; and (ii) degree of conversion at 140°C for vinyl ester resin obtained via an isothermal DSC experiment. The corresponding residence time of the fibre bundles in the die for a pultrusion speed of 0.4 m/minute used in this study is also indicated.

any modification to the original equipment. A photograph of the experimental setup that was used on site is shown in Figure 6(a) to (d): (a) illustrates the fibre creels on a creel stand; (b) shows the resin impregnation unit; (c) illustrates the impregnated fibre bundles from the impregnation unit entering the die; and (d) shows the haul-off unit. The rectangular strip (indicated by the arrow) in Figure 6(d) represents the composite made using the modified pultrusion process.

(ii) Control over the extent of impregnation and potlife of the resin: In the modified pultrusion method, the required volume of resin is delivered to the partially spread fibres in a consistent basis. During conventional pultrusion, it was observed that the fibres emanating from the resin bath were over impregnated. A significant volume of the excess resin was squeezed out from the bundles at the pre-die guides and they were directed back to the resin bath via the drip tray. The dimensions of the pre-die guides are engineered to remove the majority of the excess resin from the fibre bundles because at the entry point to the die, the excess resin that is expelled back to the resin bath is heated to some extent by the die. This can have implications to the pot-life of the resin and its viscosity in the resin bath. The effect of some of the processing parameters on the pultrusion speed was reported previously by the authors.³² A detailed study on the effect of process variables (viscosity, applied injection pressure, tow thickness and hauloff rate) on impregnation was reported previously for filament winding.³

- (iii) Non-unidirectional reinforcements and complex die profiles: In order to undertake a comparative study between the conventional resin bath and modified pultrusion techniques, a rectangular strip die geometry was chosen as it permitted the standard range of properties to be evaluated as outlined in "Test methods" section. This also meant that the impregnator in the modified technique was relatively simple as shown in Figure 2(b). However, it was capable of accommodating continuous strips (facings) of woven fabrics. This strip was generated by a weaver of E-glass fabrics where the edges are trimmed and disposed. This so-called "waste slitting"47 was spooled and used as the facings during pultrusion. The integration of woven mats and chopped strand mats as additional reinforcements during pultrusion was not investigated. However, the experience on site with the waste slittings suggests that the integration of off-axis reinforcements should not cause any practical difficulties when using the modified technique. With regard to the pultrusion of more complex shapes such as C-sections or I-beam profiles, it is proposed that the impregnator can be designed to inject the mixed resin system and then, the impregnated fibre can be separated into the different fibres trajectories using a series of pre-die profilers.
- (iv) Synchronising the pultrusion speed to the resin delivery rate: A notable advantage of the modified pultrusion technique is that the resin dispensing unit is

Resin system	Resin impregnation system	Fibres	Die temperature (°C)	Resin delivery method	Pultrusion speed (m/minute)
	Conventional pultrusion				
Unfilled vinyl ester	Resin bath	386 Direct Roving	140	Manual Pouring	0.4
Filled vinyl ester	Resin bath Modified pultrusion	386 Direct Roving	140	Manual Pouring	0.4
Unfilled vinyl ester	PUL-I	386 Direct Roving	140	Pressure-pot	0.4
Filled vinyl ester	PUL-I	386 Direct Roving	140	Pressure-pot	0.4

Table 5. Detail of the conventional resin bath and modified pultrusion experiments.



Figure 6. Photographs showing the on-site pultrusion trials: (a) Fibre creel and stand; (b) impregnator; (c) impregnated fibre bundles entering the heated die; and (d) haul-off unit (the sample in the front represents the composite that was manufactured using the modified pultrusion technique.

synchronised to the pultrusion speed. Therefore, as the pultrusion speed is increased, the resin delivery rate can be increased in proportion. There is potential to increase the pultrusion speed further in the modified method but it will need ancillary equipment to enable a higher degree of fibre spreading.²

- (v) Solvent usage and the time required for cleaning: The time required to clean the resin bath and the resin impregnation units were 20 and 8 minutes respectively. The volume of solvent that was required to clean the conventional resin bath was \sim 2000 cc where as that for the resin impregnator was \sim 200 cc.
- (vi) Fast curable resins: There is also scope to introduce fast-curing resins using the modified pultrusion technique.

As with any modification to an established commercially relevant manufacturing method, it was necessary to demonstrate that the relevant properties of the pultruded composite were not compromised. In the current study, the fibre and void volume fractions were used in conjunction with the density, mechanical and thermo-mechanical properties as indicators of the quality of the composites manufactured via the conventional and modified methods. The retention of fibre orientation (as opposed to meandering) and the extent of compaction were assessed via image analysis.

Fibre volume and void fractions and density

Fibre volume fraction and density: The fibre volume fractions and densities for the unfilled and filled vinyl



Figure 7. Fibre volume fraction and density for the pultruded vinyl ester/E-glass composites produced on-site at 0.4 m/minute.



Figure 8. Void fraction data for the unfilled and filled vinyl ester/E-glass pultruded composites produced on-site at 0.4 m/minute. The void fractions were determined in accordance with ASTM D 2584-08.

ester/E-glass composites are shown in Figure 7. In the case of the unfilled resin system, the fibre volume fractions and the densities for the conventional and modified pultruded composites are similar. When the unfilled resin was used, it was possible to feed twenty eight rovings of 4800 tex E-glass into the die. However, only twenty-four rovings were used with the filled resin system. The number of fibres for the filled-resin was reduced as it was not possible to add more than twenty-four rovings into the die. The conventional and modified techniques show similar values for the fibre volume fractions and density.

Void Fraction: With reference to the unfilled vinyl ester resin system (see Figure 8), the average void fractions for the composites produced via the conventional and modified pultrusion processes were $1.40 \pm 0.29\%$ and $0.27 \pm 0.25\%$ respectively. The average void fraction for the filled resin system for the conventional and modified pultrusion methods were $0.45 \pm 0.53\%$ and $0.64 \pm 0.78\%$ respectively. However, significant scatter is observed in all the data sets. Possible reasons for the relatively large scatter may be due to: (i) the variability in the binder content in the E-glass fibre bundles; (ii) the fibre

spreading unit breaking up areas of excess binder that could have been present on the bundles; or (iii) the homogeneity of the filler in the composite. A detailed study on this subject will be presented in a subsequent publication. Experimentally derived numerical data for the densities, fibre volume and void fractions for the two manufacturing techniques, with and without the filler are provided in Table 6.

Table 6. Experimentally derived data for the fibre volume fraction, void fraction and densities of pultruded composites using the conventional and modified pultrusion techniques; with and without the filler.

Resin system	Pultrusion method	Fiber volume fraction (%)	Void fraction (%)	Density (kg/cm ³)
Un-filled vinyl ester	Resin bath	69.4 ± 0.77	$\textbf{1.403} \pm \textbf{0.287}$	2153.31 ± 12.91
,	Resin impregnator PUL-I	$\textbf{69.1} \pm \textbf{0.77}$	$\textbf{0.268} \pm \textbf{0.251}$	$\textbf{2184.04} \pm \textbf{2.46}$
Filled vinyl ester	Resin bath	60.1 ± 1.16	$\textbf{0.459} \pm \textbf{0.229}$	$\textbf{2060.59} \pm \textbf{18.60}$
	Resin impregnator PUL-I	$\textbf{61.1} \pm \textbf{0.11}$	$\textbf{0.642} \pm \textbf{0.484}$	${\bf 2052.12} \pm {\bf 4.19}$



Figure 9. Micrographs of unfilled and filled vinyl ester/E-glass pultruded samples: (a) resin bath using unfilled resin (low-void sample); (b) resin bath using unfilled resin (high-void sample); (c) impregnator PUL-I using unfilled resin (low-void sample); (d) impregnator PUL-I using unfilled resin (high-void sample); (e) resin bath using filled resin (low-void sample); (f) resin bath using filled resin (high-void sample); (g) impregnator PUL-I using filled resin (low-void sample); (g) impregnator PUL-I using filled resin (high-void sample); (d) impregnator PUL-I using filled resin (high-void sample).

Representative micrographs of the conventional and modified pultrusion methods, with and without the filler, are shown in Figure 9. On inspecting Figure 9(a) and (b) (resin bath and unfilled resin) with Figure 9(c) and (d) (PUL-1 and unfilled resin), extent of fibre compaction is seen to be marginally higher for the modified pultrusion method. A notable feature in these samples was the significantly lower void content. Figure 9(e) to (h) represents that samples that were produced with the filled resin system using a conventional resin bath and PUL-I respectively. The observed degree of fibre compaction for the filled resin system is seen to be comparatively lower and this is assumed to be a consequence of the presence of the filler. The unfilled vinyl ester resin system shows higher shrinkage when compared to the matrix with the filler.48-50 A magnified image of de-bonding caused by the compact fibre architecture and resin-shrinkage in the unfilled vinyl ester resin system is shown in Figure 10. The effect of the higher void fraction in the unfilled composites is manifested by the lower mechanical properties (flexural and inter-laminar shear strengths).

A comparison of the void fraction data, determined using the resin burn-off and image analysis methods, is presented in Table 7. With the exception of the modified pultrusion method using the unfilled resin system, a good correlation is observed between the two methods.



Figure 10. Micrograph of an unfilled vinyl ester/E-glass sample showing the de-bonding of the resin from the fibres caused by resin shrinkage.

Flexural and inter-laminar shear properties, and dynamic mechanical thermal analysis

Flexural properties: Figure 11(a) and (b) shows the flexural strengths and flexural moduli respectively for the modified and conventional pultrusion techniques for the unfilled and filled vinyl ester/E-glass composites. The term "normalised" represents the case where the flexural strength and modulus were normalised to 60% fibre volume fraction: the average fibre volume fraction which was used for each class of pultruded composite. It can be seen that there is no significant difference between the flexural strengths of the composites produced by the conventional and modified techniques. The flexural properties are lower for the unfilled systems which may be due to de-bonding at the fibrematrix interface as seen in Figure 10. A T-test was undertaken to compare the normalised pairs of data and it was found that the data pairs are statistically insignificant for the conventional and modified pultrusion methods. On comparing the flexural strengths of the composites produced in this study with those reported in the literature,^{51–53} it can be seen from Table 8 that the data for the filled composites are similar or higher than those reported by other researchers. On the other hand, the unfilled composites exhibit lower flexural strengths. This may be related to the presence of extensive de-bonding as discussed previously.

Inter-laminar shear strength: The inter-laminar shear strengths (ILSS) for the pultruded composites are presented in Figure 11(c). As observed with the flexural strength data, the ILSS data for the composites pultruded using the conventional and modified techniques are similar for the unfilled composites. In the case of the filled composites, the ILSS values are similar for the conventional and modified techniques; however, these are higher than that observed for the unfilled composites. This is in agreement with published data.⁵⁴

Dynamic mechanical thermal analysis: As observed previously for the flexural moduli, the storage moduli (E') for the unfilled and filled resin systems are similar for the composites (normalised fibre volume fraction data) produced via conventional and modified pultrusion techniques – see Figure 11(d). The importance of normalising the mechanical properties, using the

Table 7. Comparison of the void fraction data for the pultruded vinyl ester/E-glass composites obtained by image analysis and the resin-burn off method.

Resin system	Pultrusion method	Void fraction using the resin burn-off (ASTM D 2584–08)	Void fraction using image analysis
Unfilled vinyl ester	Resin bath	$\textbf{1.40}\pm\textbf{0.29}$	$\textbf{I.48} \pm \textbf{0.46}$
	Impregnator PUL-I	0.27 ± 0.25	1.07 ± 0.29
Filled vinyl ester	Resin bath	0.45 ± 0.23	$\textbf{0.62}\pm\textbf{0.12}$
	Impregnator PUL-I	0.64 ± 0.48	$\textbf{0.62}\pm\textbf{0.12}$



Figure 11. Mechanical and thermo-mechanical properties of the pultruded vinyl ester/E-glass composites produced at 0.4 m/minute. (a) Flexural strength. (b) Flexural moduli. (c) Inter-laminar shear strength. (d) Storage moduli.

respective fibre volume fractions, to a common reference value (60% in the current case) is apparent in Figure 11 and Table 8. This approach also enables the data presented here to be compared with that reported in the literature.

Solvent usage and cleaning time

A key feature of the impregnator design was the multiple modes of impregnation of the partially spread fibre bundles (see Table 1 and Figure 3). This meant that the effective lengths of impregnation zones could be minimised to achieve full impregnation of the fibre bundles. Furthermore, the "dead volume" in the impregnator was approximately 100 cm³. The overall volume occupied by the resin during production was 965 cm³. The net effect of these factors was: (i) the volume of the resin that was retained in the impregnator at the end of a production cycle was 100 cm³; and (ii) the volume of solvent required to clean the impregnator was 200 cm³. A direct comparison with the conventional resin bath-based pultrusion process is not disclosed here due to industrial confidentiality. However, Figure 12 shows the outcome of a laboratory-based study where the conventional and modified pultrusion processes were subjected to a life cycle assessment using a commercially available LCA package. The primary benefit of the modified pultrusion technique is the significant reduction in the generation of waste resin and the lower volume of solvent required to clean the impregnator at the end of a production cycle.

Unlike conventional pultrusion where the excess resin that drips from the entrance to the hot-die is channelled back to the resin bath, in the modified pultrusion process, this volume is reduced significantly. Hence, it can be assumed that the pot-life of the resin contained within the impregnator is likely to be longer. The physical and mechanical properties obtained using the modified pultrusion technique were found to be marginally superior or equivalent to that obtained using the conventional resin bath-based method. Thus, it can be concluded that the modified pultrusion technique does not lead to any reduction in the abovementioned properties.

The main advantages of the modified pultrusion process are: (i) a 97% reduction in the generation of waste resin; and (ii) a 90% lower solvent consumption for cleaning the equipment at the end of each production shift. The cost of the implementation of the modified pultrusion technique is dictated by the choice of the resin delivery system. The cost of a pressure-pot based system will be significantly lower but it will require the resin and hardener to be weighed and

Table 8. Summary of s fraction at 60%.	elected papers to ena	ıble a comparison	of the pr	operties of tl	he UD E-glas	s composit	es. The data ir	ו square brackets ו	epresents normalised fibre volume.
Reference	Resin system	Fibres	۲ _ر [%]	Flexural strength [MPa]	Flexural modulus [GPa]	ILSS [MPa]	Storage modulus [GPa]	Manufacturing method	Other relevant information
Bogner et al. ⁵¹	Vinyl ester	E-glass	35.5	404.0 [682 8]	13.0 [72_0]	I	I	Pultrusion	Flexural: ASTM 790
Bogner et al. ⁵¹	Vinyl ester	E-glass	35.5	389.0 1467 61	[2:3 [2:3 [70 8]	I	I	Pultrusion	Flexural: ASTM 790
Johsi ⁵²	Vinyl ester	E-glass	43.5	552.6	[20.0] 25.4 ГЭС М	I	I	Pultrusion	Flexural: ASTM 790
Pultrex ⁵⁴	Vinyl ester	E-glass	I	[/ 92.2] 689.5	[0.0c] 4.14	I	I	Pultrusion	Flexural: ASTM 790
Chu and Robertson ⁵⁵	1625 series Vinyl ester	E-glass	62.0	I	I	43.0	I	Pultrusion	ILSS: ASTM D 2344
	Dow Derakane 441– 400	Hybon [®] 2100				[41.6]			
Garray et al. ⁵⁶	Vinyl ester	E-glass fiber mats	23.0	I	I	I	7.0 [18.31	RTM	DMA: Dual Cantilever
Current study									
Conventional with	Vinyl ester	E-glass	69.2	615.0	53.7	48.2	19.7	Pultrusion	Flexural: ASTM 6272
resin bath	Atlac 430	386 Direct		[533.4]	[46.5]	[41.8]	[17.1]		ILSS: ISO 14130
-	-	Roving 					-		DMA: Dual Cantilever
Modified with resin impregnator PUL-I	Vinyl ester Atlac 430	E-glass 386 Direct	c.0/	625.3 [532.5]	5.15 [43.9]	50.8 [43.2]	21.1 [17.9]	Modified Pultrusion	Flexural: ASTM 6272 ILSS: ISO 14130
0		Roving							DMA: Dual Cantilever
Conventional with	Filled vinyl ester	E-glass	61.9	785.0	47.5	61.7	18.1	Pultrusion	Flexural: ASTM 6272
resin bath	Atlac 430	386 Direct		[200.6]	[46.0]	[59.8]	[17.5]		ILSS: ISO 14130
Modified with resin	Filled vinyl ester	Koving E-glass	60.9	794.7	46.2	62.2	19.0	Modified	UMA: Dual Cantilever Flexural: ASTM 6272
impregnator PUL-I	Atlac 430	386 Direct		[782.8]	[45.5]	[61.2]	[18.71]	Pultrusion	ILSS: ISO 14130
		Roving							DMA: Dual Cantilever



Figure 12. Volume of solvent required for cleaning and the volume of waste resin generated for the conventional and modified pultrusion methods using experimental data generated in the laboratory using E-glass fibres and vinyl ester resin.

mixed manually. An automated pumping system will be more expensive but it can enable the use of resin systems with a higher reactivity and shorter pot-life. Moreover, the long-term benefits accrued from lower solvent consumption and significantly reduced cleaning time at the end of each production cycle need to be considered. The impregnator (Figures 2 and 3) can be modular and additional features such as pins, rollers, injectors, etc can be included as required.

Conclusions

A modified pultrusion process was evaluated in an industrial environment to pultrude vinyl ester/E-glass fibres using filled and unfilled resin formulations. The custom-designed impregnator was retrofitted to the pultrusion line by securing it on the top of the empty resin bath. The impregnator design used in this study was capable of handling filled resins. In this production method, the fibre bundles were partially spread via a fibre spreading rig prior to impregnation to enhance the through-thickness impregnation. Thermal analysis was used to obtain the die temperature profile and the pultrusion speeds. The physical, mechanical and thermo-mechanical properties of the pultruded composites produced, using the conventional and modified techniques, were compared. It was shown that the composites produced using the modified pultrusion technique had similar or marginally superior properties when compared to those produced using the conventional resin bath-based method. In the context of the green credentials of the modified pultrusion technique, when compared to the conventional five litre resin bath, it was demonstrated in an industrial setting that: (i) the volume of solvent required for cleaning the custom-designed impregnator, at the end of each production cycle, was reduced by 90%; and (ii) the volume of resin that is retained within the impregnator was lowered by 97%.

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