# DAMAGE AND FAILURE EFFECTS ON RESIDUAL STRESS ANALYSIS OF ARECA FIBREREINFORCED EPOXY POLYMER COMPOSITE MATERIALS

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**ABSTRACT**

*Process-induced residual stress in fiber reinforced thermoset polymer-matrix composite materials wereanalyzed using a unit cell model, with the consideration of chemical shrinkage of the epoxy resin and thermal cooling contraction of the whole fiber and resin system. The constitutive behaviour of the epoxy matrix was described by a cure and temperature dependent viscoelastic material model. Calculated residual stress shows strong dependency on the fiber volume fraction and fiber packing. The effect of residual stress on damage and failure of the model was also studied using the maximum stress failure criterion combined with a post-failure stiffness reduction technique. Both initial and final failure envelopes, predicted for biaxial normal (longitudinal and transverse) loading, were shown to be shifted and contracted by the inclusion of residual stress.*

**Keywords:** Residual stress, Thermo-viscoelastic, Unit cell model, Failure envelope.

## 1 Introduction

Process-induced residual stress in polymer matrix composites is a direct consequence of the chemical shrinkage of the matrix during polymerization and the mismatch of thermal contraction between the fiber and the matrix during cooling. The formation of residual stresses can have significant effects on the mechanical performance of composite structures by causing warpage[1] or initiating pre-load damage such as interface de-bonding and matrix micro-cracking[2,3]. Both war page and initial damage can reduce the stiffness and the strength of the material, as well as acting as sites for nucleation of macro-cracks and environmental degradation.

For thermoset polymer composites, a typical curing process consists of two steps: curing at a constant elevated temperature and thermal cooling from the curing temperature to room temperature. During curing, the polymer shrinks as a result of thepurely chemical reaction (polymerization) and its material characteristics change dramatically through the transition from a liquid state to a solid statewhile the reinforcement remains unchanged[4,5].

For thermal cooling, both polymer and reinforcement contract but by different amounts and in addition the polymer may change its stiffness significantly. Therefore, the build-up of residual stress over time is governed by the change of volume and material properties during the complete cure process.

Effects of residual stress on mechanical behaviour of the composite materials have been studied primarily by analytical methods. Nimmer[6] and Wisnom[7] showed that the presence of compressive residual stress at the interface of fiber and matrix is beneficial for the transverse behaviour of a composite with low interfacial strength. It was also shown that residual stress can result in changing and movement of the yielding surfaces for metal matrix composites[8]. For polymer-matrix composites, process-induced residual stress may introduce contraction and shifting of biaxial failure envelopes for transverse loading[9].

Residual stress is one of the major concerns in the manufacturing of polymer matrix composites, especially the effects of residual stress on damage and failure behaviour. In the

present work, unit cell analysis was used to study residual stress and its effect on damage and failure of fiber-reinforced thermoset polymer matrix composites using a micromechanical unit cell model. The residual stress introduced during curing was determined by considering both chemical shrinkage of resin and thermal cooling contraction of fiber and resin. A cure and temperature dependent viscoelastic material model was adopted to describe the constitutive relationship of the polymer matrix. Effects of fiber volume fraction and packing on residual stress were investigated. In addition, effects of residual stress on damage and failure of the unit cell subjected to mechanical loading after curing were predicted using the maximum stress failure criterion and a post-failure stiffness reduction technique.

## 2. Materials

### 2.1 Cure Kinetics

Thermoset resin undergoes chemical reactionsduring the curing process at an elevated temperature. Curing of a resin occurs over time and the degree of cure depends on the temperature history. The chemical kinetics of thermoset resin defines the dependency of the degree of cure on temperature history. In order to capture the conversion-dependent material properties, the resin cure kinetics need to be characterized accurately. For epoxy resins, the autocatalytic model has been used widely to describe cure kinetics and this is also adopted here. The model can be expressed as follows[10].

$$\frac{d\alpha}{dt} = (K_1 + K_2\alpha^{n_1})(1-\alpha)^{n_2} \qquad (2.1)$$

$$K_1 = K_{01} exp\left(\frac{E_{01}}{RT}\right) \qquad (2.2a)$$

$$K_2 = K_{02} exp\left(\frac{E_{02}}{RT}\right) \qquad (2.2b)$$

where $\alpha$ is the degree of cure, $t$the time, $K_1$and $K_2$the rate constants, $n_1$ and $n_2$ the exponents, $K_{01}$ and $K_{02}$ the temperature-independent factors, $E_{01}$ and $E_{02}$the activation energies, and $R$and $T$the universe gas constant and the temperature, respectively. The values of the parameters are given in Eom *et al.* [10]. As shown in Eom *et al*[10], the predicted degree of cure from equations (1) and (2) agrees with the experimental measurements very well for cure temperatures between 150°C and 170°C.

### 2.2 Viscoelastic Model

As shown in Xia *et al*[11], Visco-elastic material behaviour for epoxy resin can be well described by a combination of Kelvin elements connected in series in a uniaxial representation. As a result, viscous strain rate can be obtained as the sum of the strain rate of each Kelvin element, i.e.,

$$\dot{C}_{ij} = \sum_{m=1}^{n} \dot{C}_{ij}^{m} = \sum_{m=1}^{n} \left(\frac{S_{ijkl}}{E_m \tau_m}\sigma_{kl} - \frac{1}{\tau_m} C_{ij}^{m}\right) \qquad (2.3)$$

Where $S_{ijkl}$ is the compliance components and $\tau_m = {}^{\eta_m}/_{E_M} (m = 1, 2, \ldots, n)$ denotes the retardation time with $E_M$being the spring stiffness and $\mu_m$being the dashpot viscosity for the $m$-th Kelvin element, respectively.

The retardation time $\tau_m$ in equation (2.3) determines a time duration after which the contribution from the individual Kelvin element becomes negligible. Therefore, the number of Kelvin elements adopted in the constitutive equation depends on the required time range. For simplicity, a time scale factor b was introduced in Xia *et al*[11], and it was assumed that

$$\tau_m = (\beta)^{m-1}\tau_1. \qquad (2.4)$$

The description of the nonlinear behaviour inthe above model was achieved in Xia *et al*[11], by letting $E_M$be a function of the current equivalent stress, thus,

$$E_M = E_1(\sigma_{eq}) = 1.055 * 10^5 exp\left(-\frac{\sigma_{eq} - 22.764}{18.0}\right) \qquad (2.5)$$

For epoxy resin, it was shown that the material shows different behaviour inuniaxial tension and compression[12]. To account for this, the equivalent stress in equation (5) was defined as

$$\sigma_{eq} = \frac{(R-1)I_1 + \sqrt{(R-1)^2 I_1^2 + 12J_2}}{2R} \qquad (2.6)$$

Where $I_1 = \sigma_1 + \sigma_2 + \sigma_3$ is the first invariant of the stress tensor, $J_2 = {}^{S_{ij}S_{ij}}/_2$ is the second

invariant of the deviatoric stress $S_{ij}$ , and $R$ is the ratio of the tensile to compressive "yield stress".

### 2.3 Cure and Temperature Dependent Material Properties

The composite constituents considered here areglass fiber and epoxy resin. The properties of glass fiber are assumed to remain constant and independent of temperature, with the Young's modulus $E$ = 72.5 GPa, the Poisson's ratio v= 0.22 and the coefficient of thermal expansion$\chi = 5.0 * 10^{-6} / {}^{0}\text{C}$ [13].

For epoxy resin, the Young's modulus is afunction of the temperature $T$ and the degree of cure a, which is expressed as[14]

$$E(T,\alpha) = \alpha \frac{E_0}{\cosh(\alpha T)^b} \qquad (2.7)$$

Where $E_0$ is the Young's modulus for fully curedresin at room temperature, $a$ and $b$ are constants. The function (2.7) is also applied for the spring stiffness $E_M$of each Kelvin element to simulate the dependency of spring stiffness on the temperature and the degree of cure, i.e.,

$$E_m = \alpha \frac{E_1(\sigma_{eq})}{\cosh(\alpha T)^b} \qquad (2.8)$$

The thermal expansion coefficient $\chi$ of the epoxy resin is expressed as a linear function of the temperature, with $\chi$ = 63*10-6 / °C for 23°C and $\chi$= 139*10-6 / °C for 110°C [13].

In addition, the Poisson's ratio for epoxy resinis taken to be constant and assumed to be independent of degree of cure and temperature. Values of the material properties and the model parameters required by equations (2.3) to (2.8) are all given in [11] and [14].

## 3. Analysis of Residual Stress

Residual stress caused by constrainedshrinkage and thermal cooling contraction of the resin can be expressed as

$$d\sigma_{ij} = C_{ijkl}\{d\varepsilon_{ij} - dc_{ij} - \delta_{ij}\,ds - \delta_{ij}\,\alpha(T)dT\} \qquad (2.9)$$

where $d\sigma_{ij}$ are the stress increments, $d\varepsilon_{ij}$ is the total strain increments, $dc_{ij}$ is the viscous strain increments, $ds$ the shrinkage strain increment, $\alpha(T)$ the thermal expansion coefficient, $\delta_{ij}$ is the Kronecker delta, $dT$ is the temperature change, and $C_{ijkl}$ the stiffness components, related to the Young's modulus $E$ and the Poisson's ratio n of the material.

In Equation (2.9), the viscous strain can beobtained from the viscoelastic model in Section 2.2 and the thermal strain can be obtained from the assigned thermal expansion coefficients and temperature history. The chemical shrinkage strain needs to be determined from the volumetric shrinkage of the resin associated with the reaction process. For a given incremental change in the degree of cure during reaction$d\alpha$, the associatedchange in specific volume of the resin $dV$can beexpressed as[15,16]

$$dV = d\alpha V_{sh} \qquad (2.10)$$

Where $V_{sh}$ is the total volume change in the resin atthe end of cure. The isotropic resin shrinkage strainof a unit volume element of resin, resulting from incremental volume resin shrinkage, is then given by[15,16]

$$ds = \sqrt[3]{1 + dV} - 1 \qquad (2.11)$$

## 4. Damage and Failure Prediction

The unit cell model consists of fiber reinforcement and resin matrix. Depending on the failure mechanism, the following criteria for damage onset prediction were used.

For transverse failure, damage tends to occurwithin the resin matrix and is related to the stress state within the ($x$, $y$) plane. In this case, failure is predicted from the maximum principal stress criterion in the ($x$, $y$) plane, i.e.

$$\sigma_{max}^{(x,y)} \geq \sigma_u^t \; or \; \sigma_{min}^{(x,y)} \leq \sigma_u^c \qquad (2.12)$$

Where $\sigma_{max}^{(x,y)}$and $\sigma_{min}^{(x,y)}$are the maximum and minimum principal stresses in the (x, y) plane and $\sigma_u^t$ and $\sigma_u^c$are the tensile and compressive strengths of the resin, respectively.

For longitudinal failure, damage might occur inboth the fiber and the resin. Obviously, longitudinal failure is due to the normal stress in the fiber direction and the failure criterion can be expressed as

$$\sigma_{zz} \geq \sigma_u^t \; or \; \sigma_{zz} \leq \sigma_u^c \qquad (2.13)$$

Where $\sigma_{zz}$ is the normal stress in the z-direction (fiber-direction) and $\sigma_u^t$ and $\sigma_u^c$are the tensile

and compressive strengths of the fiber or the resin, respectively.

Material strengths used for failure prediction are taken in line with Soden *et al*[17], with tensile strength of 2150N/mm$^2$ and compressive strength of 1450N/mm$^2$for the areca fiber and tensile strength of 80N/mm$^2$and compressive strength of 120N/mm$^2$ for the epoxy resin

In simulating material damage, it is common practice to reduce the stiffness (or stiffness in a certain direction) to a near zero value following the onset of damage. Selective and non-selective stiffness reduction schemes are often used. Selective schemes are typically applied for composites wherethe load-carrying nature is dependent on the damage orientation[18]. In the present study, under normal loading, damage is distinguished by the transverse and longitudinal failure for matrix and thus aselective scheme is used. Specifically, for transverse failure only the modulus in the transverse direction was reduced to a near zero value (0.01 times the original value) after damage onset, while for longitudinal failure, only the modulus in the longitudinal direction was reduced (also 0.01 times the original value).

The stiffness degradation scheme, togetherwith the residual stress analysis from equation (2.9), was programmed into a user-defined material subroutine (UMAT) interfaced with the commercial standard[19]. During the analysis, the stress level was calculated at the Gauss integration points for each time increment and examined for damage detection using the above failure criteria. Once the failure criterion was satisfied, the stiffness reduction was applied for further analysis until final failure of the model.

## 5 Results and Discussion

### 5.1 Residual Stress

The cure process considered here has twostages: curing at 150°C for 3 hours and thermal cooling from 150°C to 23°C (room temperature) at a cooling rate of 2°C/min. Therefore finite element analysis was performed in two discrete steps, wherestep one is the chemical shrinkage stress analysis at 150°C and step two is the thermal cooling stress analysis from 150°C to 23°C. For the epoxy resin considered here, the total volume shrinkage $V_{sh}$ was chosen to be 3% [20-22], which corresponds to a shrinkage strain of 0.99% (about 1%).

A contour plot of the maximum principal residual stress is shown in Fig.1 for square fiberpacking with a fiber volume fraction of 60%. It can be seen that, as expected, the resin experiences a tensile maximum principal residual stress while the fiber has a compressive maximum principal residual stress. The greatest value (48.8N/mm$^2$) occurs in the central region (at $\theta$ = 45°) of the fiber-matrix interface of the quarter unit-cell model, within the resin. According to the maximum stress failure criterion, the residual stress can introduce resin failure along the interface of the fiber and the resin, which agrees with the experimental observation of interface micro-cracking shown in Gentz*et al*[2,3].

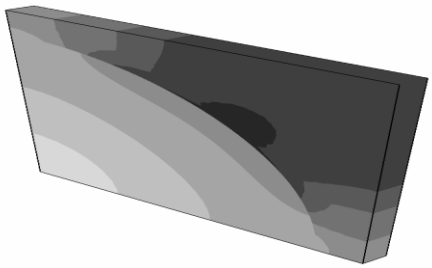

Figure 1: Contour plot of the maximum principalresidual stress (N/mm$^2$) for *Vf*= 60%.

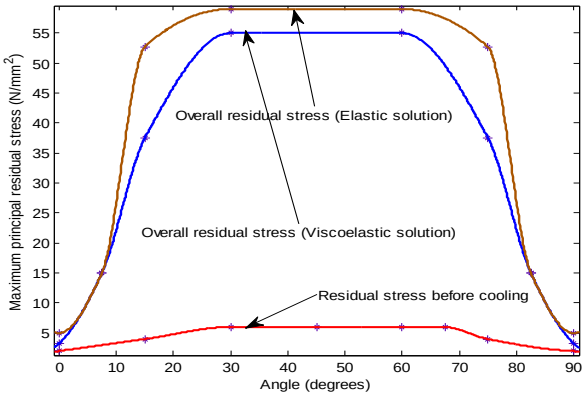


Figure 2: Distribution of the maximum principal residualstress in the resin along the fiber/matrix interface for *Vf*= 60%.

Table 1: Analysis of distribution for the max principal residualstress in resin along the fiber/matrix interface ($Vf$= 60%).

| $X_i$ | $df\ (X_i)/dX$ | $d^2f(X_i)/dX^2$ | $integral\ f(X_i)$ |
|---|---|---|---|
| 0 | 0.133333 | 0 | 0 |
| 9 | 0.133333 | 0 | 23.4 |
| 18 | 0.170667 | 0.00711111 | 57.668 |
| 27 | 0.0906667 | -0.0248889 | 104.648 |
| 36 | 0 | 0 | 158.5 |
| 45 | 0 | 0 | 212.5 |
| 54 | 0 | 0 | 266.5 |
| 63 | 0 | 0 | 320.5 |
| 72 | -0.361846 | 0.00328205 | 372.643 |
| 81 | -0.155624 | 0.00784046 | 406.662 |
| 90 | -0.044444 | 0.0168661 | 428.237 |

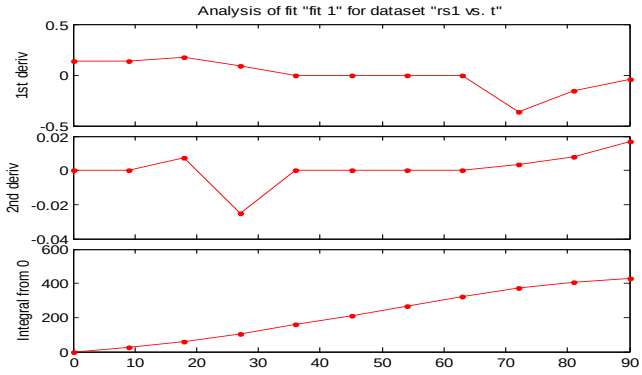


Figure2a: Analysis fit for distribution of the max principal residual stress in resin along fiber/matrix interface ($Vf$= 60%).

The distribution of the maximum principalresidual stress in the resin along the interface is presented in Figure 2, where the elastic solution and the chemical shrinkage contribution are also included. Compared to the purely elastic solution, a reduction in residual stress was predicted due to the stress relaxation caused by the viscoelastic behaviour of the epoxy matrix. Also the results show that the chemical shrinkage of resin makes only a relatively small contribution to the overall residual stress due to the relatively low modulus at the high curetemperature (150°C). At the central position in the interface of the quarter unit cell (q = 45°), the maximum principal residual stress due to chemical shrinkage has a value of 5.9 N/mm², about 12% of the overall maximum principal residual stress 48.8N/mm². This indicates that the curing shrinkage still makes a reasonable contribution to the overall residual stress, and should be included for stress analysis in polymer composites.

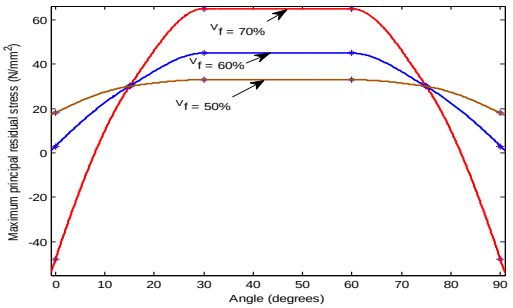


Figure3: Distribution of max principal residual stress in resin along fiber/matrix interface for three different fiber volume fractions.

Table 2: Analysis table of distribution of the max principal residual stress in resin along fiber/matrix interface for three different fiber volume fractions

| $X_i$ | $df\ (X_i)/dX$ | $d^2f(X_i)/dX^2$ | $integral\ f(X_i)$ |
|---|---|---|---|
| 0 | 6.63333 | -0.118387 | 0 |
| 9 | 4.97878 | -0.24929 | -181.71 |
| 18 | 3.2708 | -0.0412979 | 33.6768 |
| 27 | 1.33805 | -0.388201 | 509.012 |
| 36 | 0 | 0 | 1091.87 |
| 45 | 0 | 0 | 1676.87 |
| 54 | 0 | 0 | 2261.87 |
| 63 | -1.33805 | -0.388201 | 2844.74 |
| 72 | -3.2708 | -0.0412979 | 4420.07 |
| 81 | -4.97878 | -0.24929 | 3535.46 |
| 90 | -6.63333 | -0.118387 | 3353.75 |

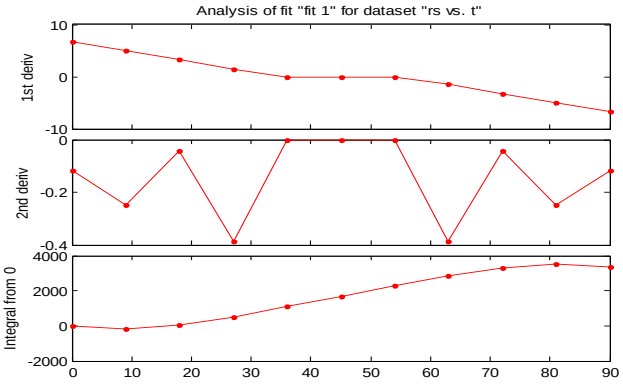


Figure3a: Analysis fit for distribution of max principal residual stress in resin along fiber/matrix interface for three different fiber volume fractions

The effects of fiber volume fraction on process-induced residual stress were studied for

square packing arrangement. Figure3 shows the maximum principal residual stress in the resin along the fiber-matrix interface for three different volume fractions, i.e., $V_f$= 50%, 60% and 70%, respectively. For all three volume fractions, the distribution of maximum principal residual stress follows a similar pattern, with the greatest value at $\theta$ = 45° and the lowest values at $\theta$ = 0° and 90°. The magnitude of residual stress was seen to increase with the increase in fiber content, since higher fiber content tends to prevent free shrinkage of the resin more significantly and causes increased residual stress. With the increase of fiber volume fraction, the maximum principal residual stress always remains tensile at $\theta$= 45°. However, at $\theta$ = 0° and 90°, the maximum principal residual stress tends to become compressive with the increase of fiber volumefraction. For $V_f$ = 70%, it is noticed that acompressive maximum principal residual stress, up to -50.2N/mm$^2$, was developed at $\theta$ = 0° and 90°. Increased tensile residual stress at $\theta$ = 45° might facilitate crack initiation or interface de-bonding in these areas, while developed compressive residualstress at $\theta$ = 0° and 90° might be beneficial inpreventing interface de-bonding in those areas[6,7].

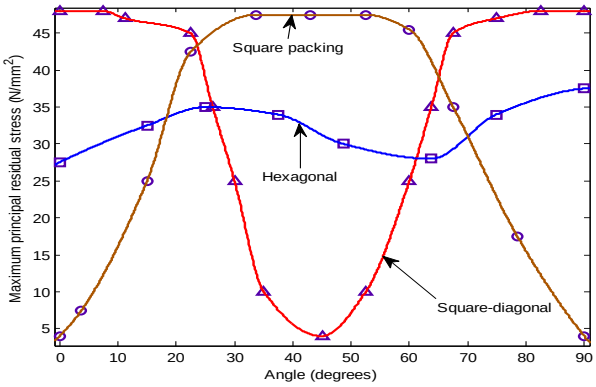


Figure 4: Distribution of the maximum principal residual stress in the resin along the fiber/matrix interface for three different fiber packing (fiber volume fraction $V_f$= 60%).

Table 3: Analysis of distribution of the maximum principal residual stress in the resin along the fiber/matrix interface for three different fiber packing (fiber volume fraction $V_f$= 60%).

| $X_i$ | $df\ (X_i)/dX$ | $d^2f(X_i)/dX^2$ | $integral\ f(X_i)$ |
|---|---|---|---|
| 0 | 0 | 0 | 0 |
| 9 | -0.313379 | -0.108874 | 431.854 |
| 18 | -0.13855 | -0.0286916 | 852.277 |
| 27 | -2.62495 | 0.0317815 | 1235.84 |
| 36 | -1.0044 | 0.0792 | 1416.37 |
| 45 | 0 | 0.335238 | 1467.9 |
| 54 | 1.63539 | 0.285591 | 1531.52 |
| 63 | 2.76301 | -0.073405 | 1727.37 |
| 72 | 0.192 | -0.0331852 | 2111.78 |
| 81 | 0.0782222 | -0.045037 | 2537.14 |
| 90 | 0 | 0 | 2969.11 |

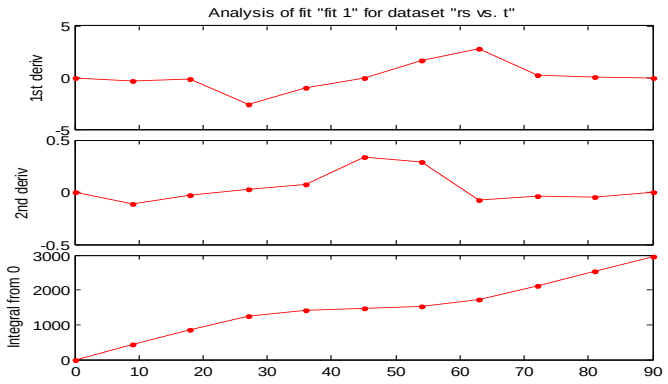


Figure 4a: Analysis fit for distribution of the maximum principal residual stress in the resin along the fiber/matrix interface for three different fiber packing (fiber volume fraction $V_f$= 60%).

In addition to the square packing, hexagonaland square-diagonal packing were also considered to study the effects of fiber packing on residual stress (fiber volume fraction was 60% for both packing arrangements). Distributions of the maximum principal residual stress of the resin along the fiber/matrix interface are presented in Figure 4 for the three different fiber-packing arrangements. The magnitude of residual stress depends on the fiber arrays, with higher values for square and square diagonal packing and lower values for hexagonal packing. The magnitude of maximum principal residual stress for hexagonal packing is about 36.8N/mm$^2$ at $\theta$ = 45°, about 25% lower than those seen in square and square diagonal

packing. This may be as a result of the more uniform distribution of resin as seen in hexagonal packing.

### 5.2 Effects of Residual Stress on Failure Envelopes

To study the influence of residual stress on theresponse of the unit cell model, the damage onset and final failure were examined under mechanical loading for a square packing model with a fiber volume fraction of 60%. After curing and thermal cooling analysis, distributed normal traction was applied to the model surfaces. At each time increment of the analysis, the initiation and evolution of damage are monitored using the maximum stress failure criterion and stiffness reduction technique described in Section 5.

Under biaxial longitudinal (z-direction) andtransverse (x-direction) normal loading, failure envelopes were constructed by considering different biaxial load ratios for initial and final failures, respectively. Failure envelopes for no residual stress, as well as the test data of final failure for unidirectional E-glass/epoxy composites from Soden *et al*[23], are also included for comparison. Initial failure level corresponds to the onset of damage. With further increase of load after damage onset, damage develops from the location of onset and spreads over the unit cell. Final failure occurs when the damage spreads across the section normal to the loading direction which makes the unit cell unable to carryany further load. It can benoticed that by considering the residual stress, both initial and final failure envelopes are shifted and contracted when compared to those derived by excluding residual stress. A similar shifting effect of residual stress on failure envelopes was also shown in Zhao *et al* [9], for transverse biaxial normal loading. Also, Aghdam and Khojeh[8] showed that residual stress caused a shifting of initial yield surfaces for unidirectional fiber-reinforced metal matrix composites under transverse loading. By comparing, it can be noticed that initial and final failure envelopes are very different in the regions with tensile longitudinal loading, since the initial failure is mainly transverse and occurs within the matrix, which is much earlier than the final matrix or fiber failure. In the regions with compressive longitudinal loading, the initial and final failure envelopes are comparable due to the rapid spread of damage over the unit cell after the damage onset

From the failure envelopes, residual stress is shown to have little effect on the load levels for longitudinal final failure due to the high fiber strengths (see the two straight edges on the left and the right), but greatly affects the transverse failure behaviour due to the relatively low resin strengths. In the transverse failure region, residual stress is mainly detrimental for transverse compression by causing earlier compressive failure. For transverse tension, residual stress has a complex effect depending on the load ratios. Residual stress is beneficial for load ratio -14.0 $<RLT<$ 4.0 and detrimental for 4.0 $<RLT<$ 48.0 ($RLT$ is the ratio of longitudinal load to transverse load).

Experimental data follow the predictions in the tension-tension region, but show a big discrepancy in the tension-compression region. There are four aspects which might contribute to the discrepancy. Firstly, test data in Soden *et al*[23] were taken from the biaxial tests performed by Al-Khalil *et al*[24] on nearly unidirectional ±85° E-glass/MY750-epoxy tubes (with a fiber volume fraction of approximately 60%), which are, strictly speaking, bi-directional composites.

Therefore, the tests could over estimate the transverse compressive strength due to the additional contributions from the fibers present in the transverse direction. Secondly, the curing procedure for the tested tubes was different, with 2 hours at 90°C followed by 1.5 hours at 130°C and 2 hours at 150°C [23], which might introduce different residual stress states in the tested tubes and affect their failure behaviour, particularly for transverse loading situations.

Thirdly, the stress based criterion for compressive damage predictionin the matrix resin used in this work might underestimate the failure level, since the test curve for compressive stress and strain shows a softening behaviour [12]. This means that compressive strength does not reflect the final failure correctly under compressive loading. Other failure criteria, such as a strain based failure criterion, may be more appropriate for detection of compressive failure in the resin matrix. Finally, non-ideal fiber distribution and concentration in real

composites could alsomake contributions to this difference [25].

## 6 Conclusions

A thermo-viscoelastic micro-mechanical model and the finite element method have been used tostudy process-induced residual stress in unidirectional fiber-reinforced polymer-matrix composites. Viscoelastic behaviour was assigned to the epoxy matrix with cure and temperature dependent material properties. From a three dimensional unit cell, residual stress was computed by considering the chemical shrinkage of the epoxy resin and the thermal cooling contraction of the whole fiber and resin system. Computed residual stress shows strong dependency on the fiber volume fraction and fiber packing. A higher fiber volume fraction will result in much greater residual stress levels due to stronger fiber constraints on the matrix contraction. Fiber packing studies suggest that evenly distributed fibers, as in hexagonal packing, could reduce the magnitude of residual stress by weakening the overall fiber constraints. Using the maximum stress failure criterion, effects of residual stress were addressed on failureenvelopes. The inclusion of residual stress results incontraction and movement of both the initial and final failure envelopes predicted for biaxial normal (longitudinal and transverse) loading. For final failure, residual stress shows little effect on the load levels for fiber-dominated longitudinal failure, but greatly affects the load levels for matrix-dominated transverse failure.

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