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On the mechanical behavior of fiber-reinforced composites

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In this paper, the derivation of the state potential is presented to model the mechanical behavior of fiber-reinforced composites. It allows matrix-cracking, interfacial debonding and sliding to be accounted for in the framework of Continuum Damage Mechanics. An application is performed on a unidirectional SiC/SiC composite. © 1998 Elsevier Science Ltd. All rights reserved.

INTRODUCTION

It is known that the formation of matrix cracks and the subsequent matrix/fiber interface debonding and sliding are the source of the non-linear stress-strain curves observed when loading Ceramic-Matrix Composites (CMCs) at room temperature. Matrix-cracking, debonding and interfacial sliding reduce the tangent Young's modulus, induce inelastic strains and hysteresis loops [1,2]. It is proposed to study these mechanisms in the framework of Continuum Damage Mechanics and to derive constitutive laws suited for structural applications. In this connection an attempt is made to base the formulation on the results of studies of the microstructural mechanisms. These studies are supported by detailed and complex experiments which are beyond the interest or abilities of those concerned with the properties of mechanical components. Therefore, the approach adopted in this study is to use the results of the micromechanical studies to develop constitutive equations which can be formulated from the results of rather simple mechanical tests.

For simplicity reasons, only experimental data on unidirectional architectures subjected to uniaxial stresses are considered but the means of generalization to multiaxial states of stresses has already been demonstrated elsewhere [3]. Special emphasis is put on the sensitivity of the identified micromechanical parameters.

A Continuum Damage Mechanics formulation applied to fiber-reinforced composites is written within the framework of the Thermodynamics of Irreversible Processes [4,5]. The first step in establishing such a model is to identify the state variables which define the *state* of the material. The second step is to derive the expression of the *state potential* $\bar{\psi}$ in terms of the state variables and the third one is to determine the *evolution laws* of the internal variables. The state potential $\bar{\psi}$ is made up of the sum of two terms: viz. the elastic energy density $\bar{\psi}_e$ and the stored energy density $\bar{\psi}_s$. The elastic part is directly related to the applied load. The stored energy density is concerned with residual stress fields giving rise to macroscopic strains with no applied load. When the local behavior is elastic, the stored energy density is expressed as the total energy density associated with the residual stress field. Therefore, by considering two elastic steps, the total free energy density can be evaluated following a so-called 'cut and paste' technique introduced by Volterra [6]. This approach will be used to model the mechanical behavior of unidirectional as well as 0/90 layered fiber-reinforced CMCs.

Loading a composite consisting of a brittle matrix supported by stronger fibers, usually causes multiple matrix-cracking which is accompanied by interfacial debonding and sliding. In the following, we assume that the whole matrix-cracking process occurs at load levels lower

than the fiber breakage mechanism, and therefore only the former mechanism will be analyzed. Using the same framework, the analysis of fiber breakage and pull-out can be found in Refs [7] and [8]. The aim of this paper is to derive a state potential and to introduce the relevant internal variables to model matrix-cracking, debonding and sliding. In the section 'Microscopic description of the degradation mechanisms', the relevant microscopic quantities are introduced to analyze the degradation mechanisms of CMCs. Based upon the results of this section, the following section, 'Macroscopic description of the degradation mechanisms' deals with a macroscopic description of constitutive laws applicable to CMCs. The section 'Micro-macro relationships for unidirectional composites' is devoted to the relationship between microscopic quantities and internal state variables. In the section 'Application to a unidirectional SiC/SiC composite', the previous results are applied to model a unidirectional fiber-reinforced SiC/SiC composite.

MICROSCOPIC DESCRIPTION OF THE DEGRADATION MECHANISMS

The matrix cracks, which are assumed to be perpendicular to the fiber direction, cause a stiffness reduction when the stress is tensile. Furthermore it is the closure of the cracks which indicates the onset of increased stiffness when the specimen is subsequently loaded in compression. In the following, we will use elementary cells of length $2L$, characterizing the crack spacing, and consisting of two different materials (m) and (f) as shown in Fig. 1 [9], where E is the Young's modulus of the unbroken composite, E_m that of part (m), and E_f that of part (f). There is a matrix crack at the center of the cell. Because the matrix contains initial flaws of random strength, there is a distribution of matrix cracks characterized by a probability density function $F(L)$ depending on the applied stress $\bar{\sigma}$. The presence of matrix cracks implies a potential energy density change $\Delta\bar{\varphi}_e$ [10]

$$\Delta\bar{\varphi}_e = \frac{1}{\bar{L}} \int_0^{+\infty} \Delta\varphi_e(L) L F(L) dL \quad (1)$$

with

$$\bar{L} = \int_0^{+\infty} L F(L) dL$$

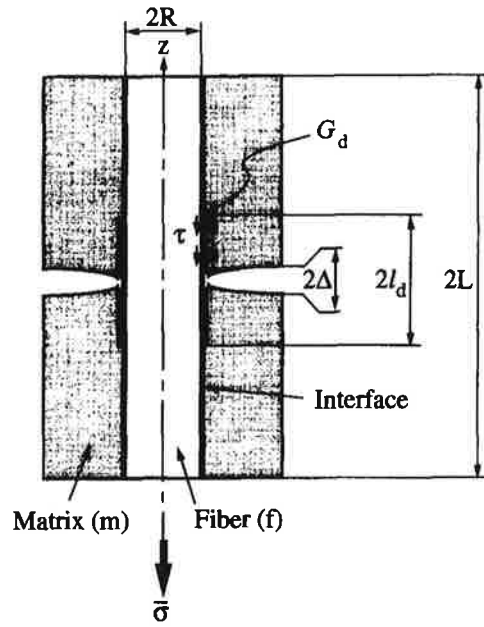


Fig. 1. Elementary cell with crack spacing $2L$ and debond length $2l_d$.

where $\Delta\varphi_e(L)$ is the average potential energy density change due to the presence of a crack in a cell of length $2L$, and $2\bar{L}$ the average crack spacing. In the framework of Linear Elastic Fracture Mechanics, it is known that the potential energy change $\Delta\varphi_e(L)$ is equal to $\bar{\sigma}^2 \omega_c / E$, where ω_c is dependent on the crack density as well as on the fact that cracks interact or not. This first basic mechanism is purely dissipative and therefore only influences the reversible part of the free energy density $\bar{\psi}_e$.

Debonding followed by sliding gives rise to inelastic strains and hysteresis loops. To describe these phenomena, different models have been proposed [11,9,12]. They all consider a friction length $2l_f$ here assumed to be equal to the debond length $2l_d$ (Fig. 1). Similarly to cracking, which is a mode I mechanism, debonding *per se* is a purely dissipative mechanism as well. When debonding and slip occur simultaneously, a self-balanced microscopic stress field is involved. The corresponding strains in the matrix and in the fiber are denoted by $\Delta\varepsilon_m(z,L)$ and $\Delta\varepsilon_f(z,L)$, respectively, where z is the current coordinate. These strain differences result from the 'cut and paste' procedure alluded to in the Introduction. By application of the principle of virtual work, the overall inelastic strain $\bar{\varepsilon}_{in}$ [13] is expressed as

$$\bar{\varepsilon}_{in} = \frac{1}{2\bar{L}} \int_0^{+\infty} \left[\int_{-L}^L \Delta\varepsilon_f(z,L) dz \right] F(L) dL. \quad (2)$$

Since the additional stress field is self-balanced the debonding and sliding processes are able to store energy. This result shows that to fully characterize both mechanisms the knowledge of the corresponding stored energy density $\bar{\psi}_{sd}$ is crucial. It is expressed as the total elastic energy density associated with the residual stress field due to debonding and sliding

$$\bar{\psi}_{sd} = \frac{1}{2\bar{L}} \int_0^{+\infty} \left[\int_{-L}^L \frac{1}{2} \{ f E_f \Delta \varepsilon_f^2(z, L) + (1-f) E_m \Delta \varepsilon_m^2(z, L) \} dz \right] F(L) dL \quad (3)$$

where f is the fiber volume fraction. It is worth noting that the expression of the stored energy is *a priori* only valid if the residual stress field is associated with debonding and sliding. Because of the way these materials are obtained, another residual stress field due to processing is likely to occur. If the residual stress in the matrix is denoted by σ_{pm} , the total stored energy density $\bar{\psi}_s$ becomes

$$\bar{\psi}_s = \bar{\psi}_{sd} - \bar{\sigma}_{pm} \bar{\varepsilon}_{in} + \bar{\psi}_{sp} \quad (4)$$

with

$$\bar{\sigma}_{pm} = \sigma_{pm} \frac{E}{E_m} \quad (5)$$

The coupling term $-\bar{\sigma}_{pm} \bar{\varepsilon}_{in}$ is due to the presence of two residual stresses of different nature. Lastly, the term $\bar{\psi}_{sp}$ is due to the residual stress field induced by processing. This term is not explicitly given since it is constant. In the following it will be omitted because the free energy density is a potential defined up to an additive constant.

From the present analysis, three different quantities are obtained. A potential energy change due to matrix-cracking, inelastic strains and a corresponding stored energy related to debonding and sliding. Furthermore, a coupling term in the stored energy density is found when the residual stresses due to processing are explicitly accounted for. These results are now used to describe macroscopically the degradation mechanisms.

MACROSCOPIC DESCRIPTION OF THE DEGRADATION MECHANISMS

The matrix-cracking process described by an internal damage variable D complies with the

requirement of a fully dissipative mechanism. In a Continuum Damage Mechanics framework, the presence of a crack results in a stiffness reduction defined by an internal damage variable D [14,15]. The Gibbs' elastic energy density of a damaged material is written as [16]

$$\bar{\varphi}_e(D) = \frac{\bar{\sigma}^2}{2E(1-D)} \quad (6)$$

so that the potential energy density change can be expressed as

$$\Delta \bar{\varphi}_e = \bar{\varphi}_e(D) - \bar{\varphi}_e(D=0) = \frac{\bar{\sigma}^2}{2E} \frac{D}{1-D} \quad (7)$$

Equation (1) shows that the correct internal variable modeling cracking is the damage variable D whereas the corresponding potential energy density change depends upon the damage variable D as well as the applied stress level $\bar{\sigma}$, and therefore is not an acceptable measure of the degradation due to matrix-cracking. Equations (1) and (7) show that the damage variable D is a function of all statistical moments of the crack spacing distribution. A 3D formulation requires an anisotropic damage description [13]. In the case of cracking perpendicular to the fiber direction, the generalization is straight-forward since the only compliance change is given in the fiber direction, and therefore only one scalar anisotropic damage variable is needed and the present 1D analysis is still relevant. For a 0/90 layered composite, the Helmholtz free energy density, which is the Legendre transformation of the Gibbs' specific enthalpy, depends upon the damage variables modeling matrix-cracking in the 0 and 90-degree layers

$$\bar{\psi}_e = \frac{1}{2} \underline{\varepsilon} : E(D_m^0, D_m^{90}) : \underline{\varepsilon} \quad (8)$$

where $\underline{\varepsilon}$ denotes the total strain tensor, $\cdot\cdot$ the contraction with respect to two indices, E the elastic stiffness tensor which is a function of the matrix-cracking damage variables D_m^0 and D_m^{90} [13].

To fully characterize debonding and sliding two variables are needed. The first one is the inelastic strain $\bar{\varepsilon}_{in}$, and the second one, denoted by d , is introduced to define the stored energy

density level associated with debonding and sliding alone

$$\bar{\psi}_s = \frac{E}{2} \frac{\bar{\epsilon}_{in}^2}{d} - \bar{\sigma}_{pm} \bar{\epsilon}_{in} \quad (9)$$

This expression can be found in Refs [17,18] to model concrete and rocks for which there is one damage variable modeling the stiffness reduction when the crack is open and sliding in mode II when the *same* crack is closed. However, in the study of CMCs there is a mode I matrix crack as well as a mode II debond crack (Fig. 1), therefore two damage variables are needed. Furthermore, there is a non-uniform stress field along the debond crack, thus eqn (9) does not necessarily allow to relate the damage variable d to a debond crack density. In a 3D formulation, a second order tensor is needed to model the inelastic strains due to debonding and slip. In the case of a layered 0/90 composite, a first order approximation only requires the in-plane components (i.e. $\bar{\epsilon}_{in11}$, $\bar{\epsilon}_{in22}$ and $\bar{\epsilon}_{in12}$), of the inelastic strain tensor $\bar{\epsilon}_{in}$. Since each operative slip system can be integrated separately in terms of energetic contributions, the internal damage variables can be defined separately for each inelastic strain term. Therefore, there are as many debond damage terms as non-vanishing inelastic strain components [13]

$$\begin{aligned} \bar{\psi}_s = & \frac{E_{11}}{2} \frac{\bar{\epsilon}_{in11}^2}{d_{11}} + \frac{E_{22}}{2} \frac{\bar{\epsilon}_{in22}^2}{d_{22}} \\ & + \frac{G_{12}}{2} \frac{\bar{\epsilon}_{in12}^2}{d_{12}} - \bar{\sigma}_{mp} : \bar{\epsilon}_{in} \end{aligned} \quad (10)$$

where E_{11} , E_{22} , are the Young's moduli along the 1-direction and the 2-direction, respectively, and G_{12} is the shear modulus in the 1–2 plane. It is worth noting that the residual stress tensor $\bar{\sigma}_{mp}$ is an average tensor in the composite to be computed for each specific architecture. In the case of cracking perpendicular to the fiber direction, only one inelastic strain component is different from zero, viz. the normal component along the fiber direction. Similarly, only one scalar debond damage variable is needed.

Because the residual stresses due to processing are assumed to be constant over the whole composite length, there is no need to introduce another internal variable to describe the coupling term in the expression of the stored energy

density $\bar{\psi}_s$. In addition to the total strain $\bar{\epsilon}$, the total free energy density $\bar{\psi}$ is dependent on three internal variables: one damage variable D modeling matrix-cracking and related to the crack spacing distribution, two variables $\bar{\epsilon}_{in}$ and d describing debonding and sliding, and related to the debond length distribution and to the crack opening distribution

$$\bar{\psi} = \frac{E(1-D)}{2} (\bar{\epsilon} - \bar{\epsilon}_{in})^2 + \frac{E}{2} \frac{\bar{\epsilon}_{in}^2}{d} - \bar{\sigma}_{pm} \bar{\epsilon}_{in}. \quad (11)$$

The associated forces are defined as follows

$$\bar{\sigma} = \frac{\partial \bar{\psi}}{\partial \bar{\epsilon}} = E(1-D)(\bar{\epsilon} - \bar{\epsilon}_{in}) \quad (12)$$

$$Y = - \frac{\partial \bar{\psi}}{\partial D} = \frac{\bar{\sigma}^2}{2E(1-D)^2} \quad (13)$$

$$\bar{X} = - \frac{\partial \bar{\psi}}{\partial \bar{\epsilon}_{in}} = \bar{\sigma} + \bar{\sigma}_{pm} - E \frac{\bar{\epsilon}_{in}}{d} \quad (14)$$

$$\begin{aligned} y = - \frac{\partial \bar{\psi}}{\partial d} &= \frac{E}{2} \left(\frac{\bar{\epsilon}_{in}}{d} \right)^2 \\ &= \frac{(\bar{\sigma} + \bar{\sigma}_{pm} - \bar{X})^2}{2E} \end{aligned} \quad (15)$$

Equation (12) defines the macroscopic stress $\bar{\sigma}$. To properly characterize the overall stress–strain behavior, this result shows that the reversible part of the free energy density is crucial. The following equations define the associated forces to each dissipative mechanism studied previously. Equation (13) defines the energy release rate density Y playing a similar role as the energy release rate \mathcal{G} in Linear Elastic Fracture Mechanics. Combining eqns (12) and (13), one shows that the energy release rate density Y is proportional to the square of the ‘effective stress’ $\bar{\sigma}/(1-D)$ [16]. Equation [14] defines the back stress \bar{X} related to debonding and sliding. Its exact value depends upon the interfacial behavior. Equation [15] defines the stored energy density y released during debonding and sliding. Since the variable d depends upon the details of the interfacial behavior, the definition of its associated force is also dependent upon the interfacial behavior. The damage variable d and its associated force y are

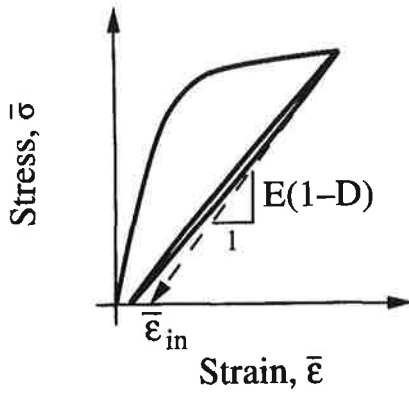


Fig. 2. Schematic stress/strain curve.

not important in terms of overall composite behavior. However in terms of energy contributions these variables are relevant. To determine the evolution laws of the internal variables, two ways can be followed. The first one is using simulations of the complete micromechanical model along the lines developed for instance by Feillard *et al.* [19] to get the distributions of crack spacings $2L$ and debond lengths $2l_d$, and then to compute the state potential. The second one is by performing experiments. The damage variable D is obtained by measuring the initial unloading modulus and the corresponding inelastic strain is $\bar{\epsilon}_{in}$ (Fig. 2). To measure the variable d , the stored energy $\bar{\psi}_s$ has to be evaluated, for instance by using methods developed by Chrysochoos [20] or Cho *et al.* [21].

For a 0/90 layered composite, the Helmholtz free energy density depends upon the damage variables modeling matrix-cracking in the 0 and 90-degree layers, the inelastic strain tensor as well as the damage variables modeling debonding and sliding

$$\begin{aligned} \bar{\psi} = & \frac{1}{2} \bar{\epsilon} : E(D_m^0, D_m^{90}) : \bar{\epsilon} + \frac{E_{11}}{2} \frac{\bar{\epsilon}_{in11}^2}{d_{11}} \\ & + \frac{E_{22}}{2} \frac{\bar{\epsilon}_{in22}^2}{d_{22}} + \frac{G_{12}}{2} \frac{\bar{\epsilon}_{in12}^2}{d_{12}} - \bar{\sigma}_{mp} : \bar{\epsilon}_{in} \end{aligned} \quad (16)$$

The associated forces are defined by

$$\bar{\sigma} = \frac{\partial \bar{\psi}}{\partial \bar{\epsilon}} \quad (17)$$

$$Y_m^0 = - \frac{\partial \bar{\psi}}{\partial D_m^0}, \quad Y_m^{90} = - \frac{\partial \bar{\psi}}{\partial D_m^{90}} \quad (18)$$

$$y_{11} = - \frac{\partial \bar{\psi}}{\partial d_{11}}, \quad y_{22} = - \frac{\partial \bar{\psi}}{\partial d_{22}},$$

$$y_{12} = - \frac{\partial \bar{\psi}}{\partial d_{12}} \quad (19)$$

$$\underline{X} = - \frac{\partial \bar{\psi}}{\partial \bar{\epsilon}_{in}}. \quad (20)$$

Equations (17)–(20) constitute a generalization of eqns (12)–(15). A description of the identification procedure can be found in Ref. [3].

MICRO-MACRO RELATIONSHIPS FOR UNIDIRECTIONAL COMPOSITES

In this section, expressions of the three internal variables D , $\bar{\epsilon}_{in}$ and d are derived in some particular cases dealing with unidirectional composites. Equation (1) can be written in terms of an equivalent cracking length \bar{L}_c defined by

$$\bar{L} \Delta \bar{\varphi}_e = \bar{L}_c \Delta \varphi_e(\bar{L}_c). \quad (21)$$

It is worth noting that when $\Delta \varphi_e(\bar{L}_c)$ is inversely proportional to \bar{L}_c , then $\Delta \varphi_e(L)$ is inversely proportional to L for any value of L and any distribution F . The relevant length to consider is the average crack spacing \bar{L} . In that case a local damage theory is applicable since the matrix cracks are not interacting with each other. Otherwise, choosing the equivalent cracking length equal to the average crack spacing is *a priori* only a zero-th order approximation. The damage variable then depends upon higher order moments of the crack spacing distribution and a non-local damage description seems more appropriate.

Using a shear lag analysis [22], eqns (7) and (21) allow us to link the damage variable D to crack spacings \bar{L}_c and \bar{L} by [10]

$$\frac{D}{1-D} = \frac{(1-f)E_m}{fE_f} \frac{\tanh(\beta \bar{L}_c)}{\beta \bar{L}} \quad (22)$$

where the constant β is a function of the elastic and geometric properties of the fiber and the matrix. When $\beta \bar{L}_c > 2.5$ (i.e. $\tanh(\beta \bar{L}_c) \cong 1$), $D/(1-D)$ is inversely proportional to $\beta \bar{L}$. This last result can also be found in the framework

of Linear Elastic Fracture Mechanics for non interacting cracks.

The strain distribution $\Delta\varepsilon_f(z, L)$ in the friction zone in the part (f) is assumed to be linear and characterized by a constant interfacial sliding strength τ_0 [23, 11]. The inelastic strain $\bar{\varepsilon}_{in}$ is given by [12]

$$\bar{\varepsilon}_{in} = \frac{\tau_0 \bar{l}_d^2}{E_f R \bar{L}_c} \left(1 + \frac{\sigma_d R}{\tau_0 \bar{l}_d} \right) \quad (23)$$

where R is the fiber radius, and \bar{l}_d is the equivalent debond length defined the same way as the equivalent cracking length \bar{L}_c . The debond strength σ_d is related to the interfacial debond toughness \mathcal{G}_d by

$$\sigma_d = 2 \sqrt{\frac{(1-f)E_f E_m \mathcal{G}_d}{RE}} \quad (24)$$

These results are a 1D simplification of the axisymmetric calculations given in Ref. [12]. Equations (2) and (23) show that the inelastic strain is a function of the crack spacing distribution as well as the debond length distribution. If the crack spacing distribution is such that the smallest spacing is greater than the debond length, there is a *unique* debond length value. The definition of the variable d is directly obtained from the evolution of the debond length \bar{l}_d as well as its definition in the expression of the stored energy ψ_s given in eqn (4). Using the debond strength σ_d and the interfacial shear strength τ_0 , the damage variable can be expressed as

$$d = \frac{(1-f)E_m}{fE_f} \frac{\bar{l}_d}{\bar{L}_c} \times \left[1 + \frac{1}{3} \left(1 + \frac{\sigma_d R}{\tau_0 \bar{l}_d} \right)^{-2} \right]^{-1} \quad (25)$$

The damage variable d is proportional to the size of the slip zone related to the crack spacing length \bar{L}_d/\bar{L}_c . When the ratio $\sigma_d R/(\tau_0 \bar{L}_d)$ approaches zero, the damage variable d reaches its lower bound. In that case, the material behaves in a low debond energy regime [24]. Conversely, when the ratio $\sigma_d R/(\tau_0 \bar{L}_d)$

approaches infinity, the damage variable d attains its upper bound and the material behaves in a high debond energy regime [24]

$$\frac{3(1-f)E_m}{4fE_f} \frac{\bar{L}_d}{\bar{L}_c} \leq d \leq \frac{(1-f)E_m}{fE_f} \frac{\bar{L}_d}{\bar{L}_c} \quad (26)$$

The present analysis shows that the key parameter discriminating low versus high debond energy regimes is the ratio $\sigma_d R/(\tau_0 \bar{L}_d)$ measuring the relative stress influence due to debonding (σ_d) and friction ($\tau_0 \bar{L}_d/R$). If the stress redistribution is more complex [19], the previous results do not apply: the damage variable d is proportional to the ratio \bar{L}_d/\bar{L}_c only if the stress field ahead of the debond crack tip is assumed to be unaltered by the presence of sliding and cracking.

APPLICATION TO A UNIDIRECTIONAL SIC/SIC COMPOSITE

In the following, we will analyze experiments performed on a unidirectional SiC/SiC composite [25]. For this material no matrix-cracking saturation is observed. This assumption is consistent with experimental observations of the evolution of the average crack spacing $2\bar{L}$ (Fig. 3) as well as that of the damage parameter ratio $D/(1-D)$. This fact allows us to make the hypothesis that the average crack spacing \bar{L} is sufficient to characterize matrix-cracking ($\bar{L}_c = \bar{L}$). By combining measurements and micromechanics, eqn (22) shows that there is a way of identifying the parameter β . Furthermore, the evolution of the average crack density

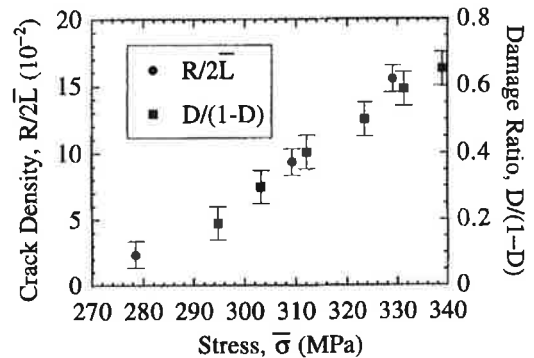


Fig. 3. Evolution of the crack density $R/2\bar{L}$ and the damage ratio $D/(1-D)$ as a function of the stress level $\bar{\sigma}$ for a unidirectional SiC/SiC composite (after Domergue [25]).

$R/2\bar{L}$ as a function of applied stress is shown to be given by [25]

$$\frac{R}{\bar{L}} = \frac{R}{\bar{L}_F} \frac{\bar{\sigma} - \bar{\sigma}_{mc}}{\bar{\sigma}_F - \bar{\sigma}_{mc}} \quad (27)$$

where $\bar{\sigma}_{mc}$ is the macroscopic stress level at which matrix-cracking starts ($\bar{\sigma}_{mc} \cong 275$ MPa), $\bar{\sigma}_F$ is the failure stress ($\bar{\sigma}_F \cong 340$ MPa), and $2\bar{L}_F$ is the average crack spacing at failure ($2\bar{L}_F \cong 40$ μm). Equations (22) and (27) can be used to identify the value of β : $R\beta = 1.09 \pm 0.07$ so that the dimensionless number $\beta\bar{L}_F = 2.8$ is in agreement with the hypothesis that $\bar{L}_c = \bar{L}$ (i.e. $\beta\bar{L}_F > 2.5$).

From a practical stand point, the evolution law can be expressed in a more tractable manner by relating the damage variable D to its associated force Y

$$D = D_\infty \left[1 - \exp \left\{ - \left(\frac{\sqrt{Y} - \sqrt{Y_{mc}}}{\sqrt{Y_n}} \right) \right\} \right] \quad (28)$$

where Y_n and Y_{mc} are material parameters, Y_{mc} is a threshold value of the energy release rate density below which no damage occurs. At the onset of matrix-cracking, the parameter Y_{mc} is related to the matrix-cracking stress by (see eqn (13) when $D = 0$)

$$Y_{mc} = \frac{\bar{\sigma}_{mc}^2}{2E} \quad (29)$$

It is worth noting that instead of eqn (28) a more complicated relationship can be obtained between the damage variable D and its associated force Y using eqns (13), (22), (27) and (29). The result of the derivation shows that \sqrt{Y} is the right quantity to consider. Therefore, eqn (28) was written in terms of \sqrt{Y} rather than Y itself. For the unidirectional SiC/SiC composite, the following values are identified (Fig. 4)

$$D_\infty = 0.67 \pm 0.05, Y_{mc} = 147 \pm 0.7 \text{ kPa},$$

$$Y_n = 9.6 \pm 0.4 \text{ kPa}. \quad (30)$$

The identified value of Y_{mc} is in agreement with the value of the matrix-cracking stress $\bar{\sigma}_{mc}$ according to eqn (29). If eqn (29) is considered (i.e. the value of Y_{mc} can be obtained from the knowledge of $\bar{\sigma}_{mc}$), the only parameter to identify is Y_n , since the value of D_∞ is obtained by inspecting eqn (22) when $\bar{L} \rightarrow 0$ and $\bar{L}_c \rightarrow 0$, $D \rightarrow D_\infty = (1-f) E_m/E$. The value of D_∞ is also

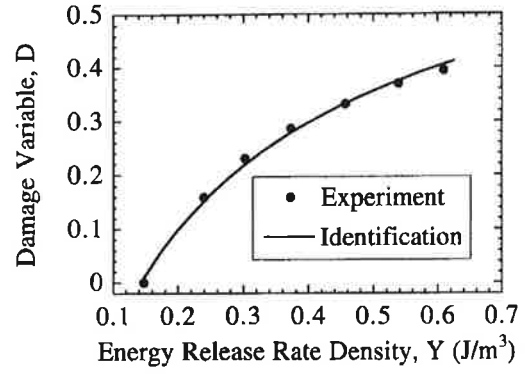


Fig. 4. Evolution of the damage variable D as a function of the energy release rate density Y for a unidirectional SiC/SiC composite.

in agreement with that predicted by using the elastic properties of the SiC/SiC composite ($E_m = 300$ GPa, $E_r = 200$ GPa, $f = 0.43$).

To completely identify the evolution of the inelastic strains, one needs to determine the residual stress $\bar{\sigma}_{pm}$, the debond strength σ_d and the interfacial shear strength τ_0 . The difference in coefficients of thermal expansion is assumed to be on the order of $\Delta\alpha \cong 10^{-6} K^{-1}$ and the temperature variation on the order of $\Delta T = 1000$ K. Consequently, the residual stress level can be approximated by $\bar{\sigma}_{pm} \cong 80$ MPa. This value is in reasonable agreement with other studies [25]. In the following, it is assumed that only one debond length is needed to fully characterize the debond regions. The other parameters are obtained by identification of the evolution of the inelastic strains $\bar{\epsilon}_{in}$ as a function of the stress level $\bar{\sigma}$

$$\bar{\epsilon}_{in} = \frac{(1-f)^2 E_m^2 R}{4f^2 E_r \tau_0 \bar{L}} \times \left[\left(\frac{\bar{\sigma} + \bar{\sigma}_{pm}}{E} \right)^2 - \left\{ \frac{f\sigma_d}{(1-f)E_m} \right\}^2 \right]. \quad (31)$$

The following values are obtained (see Fig. 5)

$$\sigma_d = 550 \pm 10 \text{ MPa}, \tau_0 = 46 \pm 5 \text{ MPa}. \quad (32)$$

The value of the interfacial shear strength τ_0 is in good agreement with other identifications [25] as well as the debond strength (the corresponding interfacial toughness \mathcal{G}_d is on the order of 4.2 ± 0.1 J/m²).

If one assumes that the onset of matrix-cracking coincides with the onset of debonding, the value of the debond strength can be easily iden-

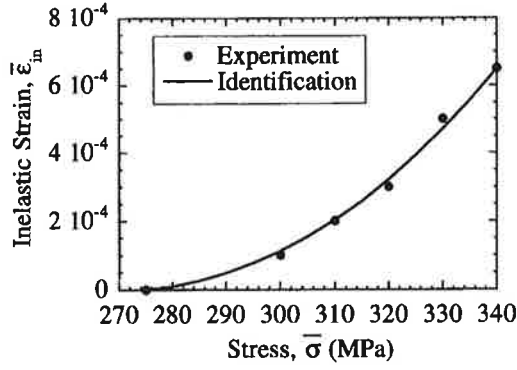


Fig. 5. Evolution of the inelastic strain $\bar{\varepsilon}_{in}$ as a function of the stress level $\bar{\sigma}$ for a unidirectional SiC/SiC composite.

tified, provided that the residual stresses are known

$$\frac{f\sigma_d}{(1-f)E_m} = \frac{\bar{\sigma}_{mc} + \bar{\sigma}_{pm}}{E} \quad (33)$$

This hypothesis can be made in the present case. Using eqn (33), a relative variation of the residual stress $\Delta\bar{\sigma}_{pm}/|\bar{\sigma}_{pm}|$ leads to a relative variation of $\Delta\sigma_d/\sigma_d$ expressed as

$$\frac{\Delta\sigma_d}{\bar{\sigma}_d} = \left(1 + \frac{\bar{\sigma}_{mc}}{\sigma_d}\right)^{-1} \frac{\Delta\bar{\sigma}_{pm}}{|\bar{\sigma}_{pm}|} \quad (34)$$

Equation (34) shows that the higher the dimensionless group $(1 + \bar{\sigma}_{mc}/\sigma_d)$, the lower the sensitivity of the debond strength σ_d to the residual stress $\bar{\sigma}_{pm}$. In the present case eqn (34) becomes

$$\frac{\Delta\sigma_d}{\bar{\sigma}_d} = 0.23 \frac{\Delta\bar{\sigma}_{pm}}{|\bar{\sigma}_{pm}|} \quad (35)$$

This result indicates that there is a weak influence of the parameter $\bar{\sigma}_{pm}$ on the identification of the debond strength $\bar{\sigma}_d$.

Furthermore, by using eqns (27) and (31), the evolution of the inelastic strain can be rewritten as a function of the applied stress $\bar{\sigma}$

$$\bar{\varepsilon}_{in} = B \frac{(\bar{\sigma} - \bar{\sigma}_{mc})^2 (\bar{\sigma} + 2\bar{\sigma}_{pm} + \bar{\sigma}_{mc})}{E^2 (\bar{\sigma}_F - \bar{\sigma}_{pm})} \quad (36)$$

with

$$B = \frac{D_F}{1 - D_F} \frac{(1-f)E_m \beta R}{f\tau_0} \quad (37)$$

Equation (37) shows that the identification of τ_0 is directly related to the value of β . There-

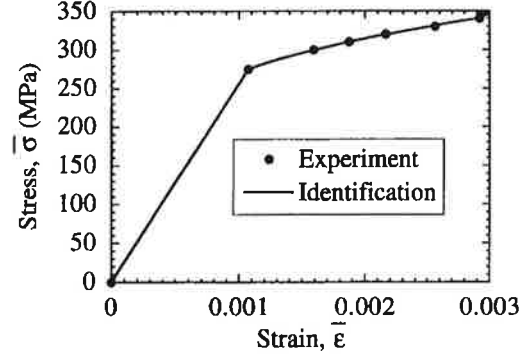


Fig. 6. Evolution of the stress $\bar{\sigma}$ as a function of the strain $\bar{\varepsilon}$ for a unidirectional SiC/SiC composite.

fore, a relative variation of the parameter β , $\Delta\beta/\beta$, leads to the same relative variation of the shear strength τ_0 , $\Delta\tau_0/\tau_0$

$$\frac{\Delta\tau_0}{\tau_0} = \frac{\Delta\beta}{\beta} \quad (38)$$

This influence is significantly higher than that exhibited by the residual stress $\bar{\sigma}_{pm}$ in relation with the debond strength $\bar{\sigma}_d$. These features can be observed when the scatter in values for τ_0 and \mathcal{G}_d are summarized by Evans *et al.* [24]. The identified stress/strain curve is compared with the experimental one in Fig. 6: there is a good agreement.

To fully characterize the model, one needs to study the evolution of the damage parameter d defined in eqn (25). Figure 7 shows the evolution of the damage parameter d as a function of stress level $\bar{\sigma}$. The prediction is compared with the upper and lower bounds of the damage parameter given in eqn (26). In the particular case of a unidirectional SiC/SiC composite, a

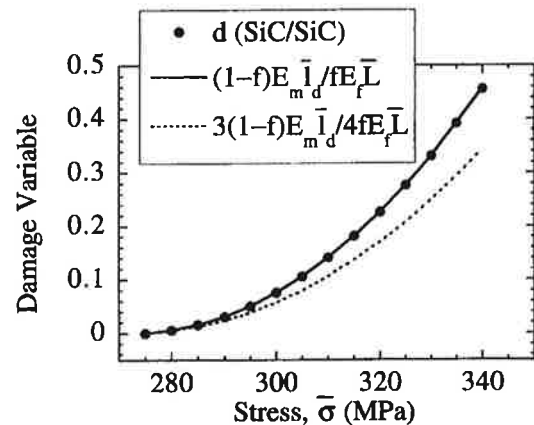


Fig. 7. Evolution of the damage variable d and the corresponding upper bound (solid line) and lower bound (dashed line) as a function of the stress level $\bar{\sigma}$ for a unidirectional SiC/SiC composite.

large debond energy regime is relevant for the whole stress range since the damage parameter coincides with its upper bound. Other composite materials such as layered alumina with carbon/epoxy prepregs are low debond energy materials for which the debond strength is negligible with respect to the average shear stress [26].

CONCLUSIONS

A Continuum Damage Mechanics formulation has been applied to fiber-reinforced composites. In addition to the total strain, the internal variables which define the state of the material have been identified. Matrix-cracking is described by one damage variable, debonding and sliding are modeled by an inelastic strain and another damage variable measuring the amount of stored energy. These variables are related to microscopic quantities introduced to analyze the degradation mechanisms of CMCs.

Micromechanical parameters are exhibited to model matrix-cracking, interfacial debonding and sliding. Their identification is discussed by analyzing the mechanical behavior of a unidirectional SiC/SiC composite. This composite behaves in a large debond energy regime in which the stress levels related to the debond strength are significantly higher than those related to sliding. In the present case, a sensitivity analysis shows that the identification of the interfacial shear strength is more sensitive than that of the debond strength.

The framework presented in this paper can be extended to model layered as well as woven fiber-reinforced composites. The nature of the different internal variables have been discussed. The same formalism can also be used to model high temperature applications in which the change of residual stresses as well as creep mechanisms need to be incorporated. This work is still in progress.

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