An Original Approach of Tensile Behaviour and Elastic Properties of Multiphase Pre-Impregnated Composite Materials

HORATIU TEODORESCU, SORIN VLASE, LUMINITA SCUTARU, FLORIN TEODORESCU
Department of Mechanical Engineering
Transilvania University of Brasov
29 Eroilor Blvd., 500036 Brasov
Mechanical Engineering Research Institute SC ICTCM SA
103 Oltenitei Blvd., 041303 Bucharest
ROMANIA
hteodorescu@yahoo.com, svlase@unitbv.ro, lscutaru@unitbv.ro, fteodorescu@ictcm.ro

Abstract: - An original approach to compute the longitudinal tensile break stress of multiphase composite materials with short fibers reinforcement is presented. The most obvious mechanical model which reflects a multiphase composite material is a pre-impregnated material, known as prepreg. In the class of prepregs the most known are Sheet- and Bulk Molding Compounds (SMCs and BMCs). The model is seen as consisting of three phase compounds: resin, filler and fibers, model that is reduced to two phase compounds: substitute matrix and fibers. The Sheet Moulding Compounds reinforced with discontinuous and almost parallel fibers, subjected to longitudinal tensile loads, presents a specific note by the existence of a shear mechanism between fibers and matrix. This shear mechanism transfers the tensile load through the fibers. The Young’s moduli for the substitute matrix and for the entire composite are computed and a comparison between the theoretical approach and the experimental data is accomplished. The paper presents also an original homogenization method to predict the elastic properties of these materials. The upper and lower limits of the homogenized coefficients for a 27% fibers volume fraction SMC are computed. It is presented a comparison between the upper and lower limits of the homogenized elastic coefficients of a SMC material and the experimental data. The estimation model used as a homogenization method of these heterogeneous composite materials, gave emphasis to a good agreement between this theoretical approach and experimental data.


1 Introduction
The most obvious mechanical model which features a multiphase composite material is a pre-impregnated material, known as prepreg. In the wide range of prepregs the most common used are Sheet- and Bulk Molding Compounds. A Sheet Molding Compound (SMC) is a pre-impregnated material, chemically thickened, manufactured as a continuous mat of chopped glass fibers, resin (known as matrix), filler and additives, from which blanks can be cut and placed into a press for hot press moulding. The result of this combination of chemical compounds is a heterogeneous, anisotropic composite material, reinforced with discontinuous reinforcement [1], [2], [3].

A typical SMC material is composed of the following chemical compounds: calcium carbonate (36.8% weight fraction); chopped glass fibers rovings (30% weight fraction); unsaturated polyester resin (18.4% weight fraction); low-shrink additive (7.9% weight fraction); styrene (1.5% weight fraction); different additives (1.3% weight fraction); pigmented paste (1.3% weight fraction); release agent (1.2% weight fraction); magnesium oxide paste (1.1% weight fraction); organic peroxide (0.4% weight fraction); inhibitors (0.1% weight fraction). The matrix (resin) system play a significant role within a SMC, acting as compounds binder and being “embedded material” for the reinforcement. To decrease the shrinkage during the cure of a SMC prepreg, filler (calcium carbonate) have to be added in order to improve the flow capabilities and the uniform fibers transport in the mold. For the materials that contain many compounds, an authentic, general method of dimensioning is hard to find. In a succession of hypotheses, some authors tried to describe the elastic properties of SMCs based on ply models and
on material compounds [4], [5], [6]. The glass fibers represent the basic element of SMC prepreg reinforcement. The quantity and rovings’ orientation determine, in a decisive manner, the subsequent profile of the SMC structure’s properties. There are different grades of SMC prepregs: R-SMC (with randomly oriented reinforcement), D-SMC (with unidirectional orientation of the chopped fibers), C-SMC (with unidirectional oriented continuous fibers) and a combination between R-SMC and C-SMC, known as C/R-SMC.

The following informations are essential for the development of any model to describe the composite materials behaviour [7]: the thermo-elastic properties of every single compound and the volume fraction concentration of each compound.

Theoretical researches regarding the behaviour of heterogeneous materials lead to the elaboration of some homogenization methods that try to replace a heterogeneous material with a homogeneous one [8], [9], [10]. The aim is to obtain a computing model which takes into account the microstructure or the local heterogeneity of a material. The homogenization theory is a computing method to study the differential operators’ convergence with periodic coefficients [11], [12]. This method is indicated in the study of media with periodic structure like SMCs and BMCs. The matrix- and fillers elastic coefficients are very different but periodical in spatial variables. This periodicity or frequency is suitable to apply the homogenization theory to the study of heterogeneous materials.

2 Tensile Behaviour Model of a SMC Material

A SMC material can be regarded as a system of three basic compounds: resin, filler and reinforcement (fibers). We can consider the resin–filler system as a distinct phase compound called substitute matrix, so a SMC can be regarded as a two phase compound material (fig. 1). This substitute matrix presents the virtual volume fractions for resin and for filler. These virtual volume fractions are connected to the real volume fractions and through the relations:

\[ V_r' = \frac{V_r}{V_r + V_f}; \quad V_f' = \frac{V_f}{V_r + V_f}, \]  

so that \( V_r' + V_f' = 1 \).

It is known that during the manufacturing process of a SMC, there is dependence between the production line speed and the fibers plane orientation on its advance direction. So, this material can be assumed to have the fibers oriented almost parallel to the production line of the SMC. Due to the longitudinal tensile loading, the SMC strain \( (\varepsilon_C) \) is identical with the substitute matrix strain \( (\varepsilon_{SM}) \) and fibers strain \( (\varepsilon_F) \), see fig. 2.

\[ \varepsilon_C = \varepsilon_F = \varepsilon_{SM}, \]

Assuming the fact that both fibers and substitute matrix present an elastic linear behaviour, the respective longitudinal stresses are:

\[ \sigma_f = E_f \cdot \varepsilon_f = E_f \cdot \varepsilon_C, \]  
\[ \sigma_{SM} = E_{SM} \cdot \varepsilon_{SM} = E_{SM} \cdot \varepsilon_C. \]

The tensile force applied to the entire composite is taken over by both fibers and substitute matrix:

\[ P = P_F + P_{SM}, \]

or:

\[ \sigma_C \cdot A_C = \sigma_f \cdot A_f + \sigma_{SM} \cdot A_{SM}, \]  
\[ \sigma_C = \frac{A_f \cdot \sigma_f}{A_C} + \frac{A_{SM} \cdot \sigma_{SM}}{A_C}, \]

where \( \sigma_C \) is the medium tensile stress in the composite, \( A_f \) is the net area of the fibers transverse surface, \( A_{SM} \) represents the net area of the substitute matrix transverse surface and \( A_C = A_f + A_{SM} \).

The ratio: \( \frac{A_f}{A_C} = V_f \) is the fibers volume fraction and \( \frac{A_{SM}}{A_C} = V_{SM} = 1 - V_f \) represents the substitute matrix volume fraction, so that (5) becomes:

\[ \sigma_C = \sigma_f \cdot V_f + \sigma_{SM} \cdot (1 - V_f). \]

Taking into account (2) and (3) and dividing both terms of (6) through \( \varepsilon_C \), the longitudinal elasticity modulus for the composite is:

\[ E_C = E_f \cdot V_f + E_{SM} \cdot (1 - V_f). \]
Equation (7) shows that the value of the longitudinal elasticity modulus of the composite is situated between the values of the fibers- and substitute matrix longitudinal elasticity moduli. In general, the fibers' break strain is lower than the matrix's break strain, so assuming that all fibers present the same strength, their break lead inevitable to the composite break. According to equation (6), the composite's break strength at longitudinal tensile loads is:

$$\sigma_{BC} = \sigma_{SF} \cdot V_F + E_{SM} \cdot \epsilon_{SF} \cdot (1 - V_F)$$

(9)

The estimation of the substitute matrix longitudinal elasticity modulus in case of a heterogeneous material like SMC, obtained by mixing some materials with well-defined properties, depends both on the basic elastic properties of the isotropic compounds and the volume fraction of each compound. If we note down $E_r$ the basic elastic property of the resin, $E_f$ the basic elastic property of the filler, $V_r$ the resin volume fraction and $V_f$ the filler volume fraction, the substitute matrix longitudinal elasticity modulus can be estimated computing the harmonic media of the basic elastic properties of the isotropic compounds, as follows:

$$E_{SM} = \frac{2}{\frac{1}{E_r \cdot V_r} + \frac{1}{E_f \cdot V_f}}$$

(10)

A SMC material reinforced with discontinuous almost parallel fibers, subjected to longitudinal tensile loads, presents a particularity by the existence of a shear mechanism between fibers and matrix, mechanism that transfers the tensile load to the fibers. Due to a difference between the substitute matrix longitudinal strain and the fibers strain, a shear stress along the fiber-substitute matrix interface occurs.

The normal stress distribution in a discontinuous fiber can be computed, considering an infinitely small portion $dx$ at the distance $x$ from one fiber end (fig. 3) [13]:

$$\left(\frac{\pi}{4} \cdot d_F^2\right) \cdot (\sigma_F + d\sigma_F) \cdot \left(\frac{\pi}{4} \cdot d_F^2 \cdot \sigma_F\right) = \frac{\pi}{2} \cdot d_F^2 \cdot \sigma_F \cdot dx \cdot \tau_i = 0$$

(11)

or:

$$\frac{d\sigma_F}{dx} = \frac{4\tau_i}{d_F}$$

(12)

where: $\sigma_F$ is the fiber longitudinal stress at the distance $x$ from one of its end, $d_F$ is the fiber diameter and $\tau_i$ represents the shear stress at the fiber-substitute matrix interface. Assuming $\tau_i$ constant, $\sigma_F = 0$ at the distance $x = 0$ and integrating (12), we get:

$$\sigma_F = \frac{4}{d_F} \cdot \int_0^x \tau_i \cdot dx = \frac{4\tau_i}{d_F} \cdot x$$

(13)

The maximum fiber stress can be reached at a distance $x = \frac{l_T}{2}$ from both fiber ends, $l_T$ being the load transfer length and represents the fiber minimum length in which fiber maximum stress is
from (14) we may compute a critical fiber length for given \( d_F \) and \( \tau_i \):

\[
I_{\text{critical}} = \frac{\sigma_{bF}}{2\tau_i} d_F. 
\]  

(15)

Taking into account the normal stress distributions also near the fiber ends (for \( x < \frac{l_F}{2} \)) then a medium stress in fiber can be computed:

\[
\sigma_F = \frac{1}{l_F} \int_0^{l_F} \sigma_F \cdot dx, \quad (16)
\]

or:

\[
\sigma_F = \sigma_{\text{max}} F \left( 1 - \frac{l_T}{2l_F} \right). \quad (17)
\]

If the fiber length is greater than its critical length \( (l_F > I_{\text{critical}}) \), replacing \( \sigma_{\text{max}} F = \sigma_{bF} \) and \( I_T = I_{\text{critical}} \), the longitudinal break strength of a SMC material can be computed as follows:

\[
\sigma_{bC} = \sigma_{bF} V_F + \sigma_{SMY} (1-V_F) = \frac{1}{2l_F} \left( 1 - \frac{l_{\text{critical}}}{2l_F} \right) V_F + \sigma_{SMY} (1-V_F). \quad (18)
\]

3 A Homogenization Method

We consider \( \Omega \) a domain from \( R^3 \) space, in coordinates \( x_i \), domain considered a SMC composite material, in which a so called substitute matrix (resin and filler) is represented by the field \( Y_1 \) and the reinforcement occupies the field \( Y_2 \) seen as a bundle of glass fibers, (fig. 4).

Let us consider the following equation [14]:

\[
\frac{\partial}{\partial x_i} \left[ a_{ij} \frac{\partial u}{\partial x_j} \right] + a_{ij} = 0, \quad (19)
\]

or under the equivalent form:

\[
f = \frac{\partial p_i}{\partial x_i}; \quad p_i = a_{ij} \frac{\partial u}{\partial x_j}. \quad (20)
\]

In the case of SMC materials that present a periodic structure containing inclusions, \( a_{ij}(x) \) is a function of \( x \). If the period’s dimensions are small in comparison with the dimensions of the whole domain then the solution \( u \) of the equation (19) can be considered equal with the solution suitable for a homogenized material, where the coefficients \( a_{ij} \) are constants.

In the \( R^3 \) space of \( y_i \) coordinates, a parallelepiped with \( y_i^0 \) sides (fig. 4) is considered, as well as parallelepipeds obtained by translation \( n_i y_i^0 \) (\( n_i \) integer) in axes directions. The functions:

\[
a_{ij}^\eta (x) = a_{ij} \left( \frac{x}{\eta} \right), \quad (21)
\]

can be defined, where \( \eta \) is a real, positive parameter. Notice that the functions \( a_{ij}(x) \) are \( \eta Y \)-periodical in variable \( x \) (\( \eta Y \) being the parallelepiped with \( \eta Y^0 \) sides). If the function \( f(x) \) is in \( \Omega \) defined, the problem at limit can be considered:

\[
f(x) = \frac{\partial}{\partial x_i} \left[ a_{ij}^\eta (x) \frac{\partial u^\eta}{\partial x_j} \right], \quad (22)
\]

\[
u^\eta \big|_{\partial \Omega} = 0.
\]

Similar with equation (20), the vector \( \bar{p}^\eta \) can be defined with the elements:

\[
p_{ij}^\eta (x) = a_{ij}^\eta (x) \frac{\partial u^\eta}{\partial x_j}. \quad (23)
\]

For the function \( u^\eta(x) \) an asymptotic development will be looking for, under the form:

\[
u^\eta(x,y) = u^\eta(x,y) + \xi \eta u^\eta(x,y) + \xi^2 \eta^2 u^\eta(x,y) + \ldots; \quad y = \frac{y}{\eta}, \quad (24)
\]

where \( u^\eta(x,y) \) are \( \eta \)-periodical in \( y \) variable. The functions \( u^\eta(x,y) \) are defined on \( \Omega \times R^3 \) so that the derivatives behave in the following manner:
If the values of \( u(x, y) \) are compared in two homologous points \( P_1 \) and \( P_2 \), homologous through periodicity in neighbour periods, it can be notice that the dependence in \( x \) is the same and the dependence in \( y \) is almost the same since the distance \( P_1P_2 \) is small (fig. 5). Let us consider \( P_3 \) a point homologous to \( P_1 \) through periodicity, situated far from \( P_1 \). The dependence of \( u(x, y) \) is the same but the dependence in \( x \) is very different since \( P_1 \) and \( P_3 \) are far away. For instance, in the case of two points \( P_1 \) and \( P_4 \) situated in the same period, the dependence in \( x \) is almost the same since \( P_1 \) and \( P_4 \) are very close, but the dependence in \( y \) is very different since \( P_1 \) and \( P_4 \) are not homologous through periodicity. The function \( u^0 \) depends on the periodic coefficients \( a_{ij} \), on the function \( f(x) \) and on the boundary \( \partial\Omega \). The development (24) is valid at the inner of the boundary \( \partial\Omega \), where the periodic phenomena are prevalent but near and on the boundary, the non-periodic phenomena prevail [15], [16].

Using the development (24), the expressions \( \frac{\partial u^0}{\partial x_i} \) and \( p^0 \) can be computed as following [14]:

\[
\frac{\partial u^0}{\partial x_i} = \left( \frac{\partial}{\partial x_i} + \frac{1}{\eta} \frac{\partial}{\partial y_i} \right) \left( u^0 + \eta \cdot u^1 + \ldots \right) = \\
\frac{\partial u^0}{\partial x_i} + \frac{\partial u^1}{\partial y_i} + \eta \left( \frac{\partial u^1}{\partial x_i} + \frac{\partial u^2}{\partial y_i} \right) + \ldots, \quad (26)
\]

\[
p^0 (x, y) = p^0 (x, y) + \eta \cdot p^1 (x, y) + \eta \cdot p^2 (x, y) + \ldots. \quad (27)
\]

where:

\[
p^0 (x, y) = a_{ij} (y) \left( \frac{\partial u^0}{\partial x_j} + \frac{\partial u^1}{\partial y_j} \right), \quad (28)
\]

\[
p^1 (x, y) = a_{ij} (y) \left( \frac{\partial u^1}{\partial x_j} + \frac{\partial u^2}{\partial y_j} \right) + \ldots
\]

The function \( f(x) \) presented in equation (22) can be written in the following manner:

\[
f(x) = \left( - \frac{\partial}{\partial x_i} - \frac{1}{\eta} \frac{\partial}{\partial y_i} \right) \left( p^0 + \eta \cdot p^1 + \ldots \right). \quad (29)
\]

The terms \( \eta \cdot p^1 \) and \( \eta \cdot p^0 \) will be:

\[
\frac{\partial p^0}{\partial y_i} = 0, \quad (30)
\]
\[ f(x) = -\frac{\partial p^0}{\partial x_i} + \frac{\partial p^1}{\partial y_i}. \] (31)

Equation (31) leads to the homogenized- or macroscopic equation. For this, the medium operator is introduced, defined for any function \( \Psi(y) \), Y-periodical:

\[ \langle \Psi \rangle_Y = \frac{1}{|Y|} \int_Y \Psi(y) \, dy, \] (32)

where \(|Y|\) represents the periodicity cell volume. To obtain the homogenized equation, the operator (32) \( f(x) = -\frac{\partial (p^1)}{\partial x_i} + \frac{\partial p^1}{\partial y_i} \) is applied to equation (31):

\[ f(x) = -\frac{\partial (p^1)}{\partial x_i} + \frac{\partial p^1}{\partial y_i}. \] (33)

According to the operator (32), the second term of the left side of the equation (33) becomes:

\[ \langle \Psi \rangle_Y = \frac{1}{|Y|} \int_Y \frac{\partial p^1}{\partial y_i} \, dy = \frac{1}{|Y|} \int_Y p^1 n_i ds = 0. \] (34)

Due to Y-periodicity of \( p^1 \) and the fact that \( n_i \) is the normal vector at the boundary of \( Y \), the relation (34) is equal with zero. So, the equation (33) becomes:

\[ f(x) = -\frac{\partial (p^1)}{\partial x_i}. \] (35)

With help of relation (28), the equation (30) can be written as follows:

\[ \frac{\partial}{\partial y_i} \left[ a_y(y) \left( \frac{\partial u^0 + \partial u^1}{\partial y_j} \right) \right] = 0, \] (36)

to therefore:

\[ \frac{\partial}{\partial y_i} \left[ a_y(y) \cdot \frac{\partial u^0}{\partial y_j} \right] = \frac{\partial u^0}{\partial y_j} \cdot \frac{\partial a_y}{\partial y_j}. \] (37)

The solution \( u^1(y) \) of equation (37) is Y-periodical and to determine its periodicity cell of a SMC composite material which occupies the domain \( Y_1 \) and presents the coefficient \( a^{1}_y \) and the inclusion occupies the domain \( Y_2 \) with the coefficient \( a^{1}_y \) separated by a surface \( \Gamma \), the equation (21) must be seen as a distribution.

In the case of a SMC composite material which behaves macroscopically as a homogeneous elastic environment, it is important the knowledge of the elastic coefficients. Unfortunately, a precise calculus of the homogenized coefficients can be achieved only in two cases: the one-dimensional case and the case in which the matrix- and inclusion coefficients are functions of only one variable. For a SMC material is preferable to estimate these homogenized coefficients between an upper and a lower limit.

Since the fibers volume fraction of common SMCs is 27%, to lighten the calculus, an ellipsoidal inclusion of area 0.27 situated in a square of side 1 is considered. The plane problem will be considered and the homogenized coefficients will be 1 in matrix and 10 in the ellipsoidal inclusion. In fig. 6, the structure’s periodicity cell of a SMC composite

Knowing the expression of \( u^1 \) as a function of \( u^0 \), from the expressions (28) with (40), the homogenized coefficients can be computed:

\[ p^0(x,y) = a_y(y) \left( \frac{\partial u^0}{\partial x_j} + \frac{\partial u^1}{\partial y_j} \right) \]

\[ a_y(y) \left( \frac{\partial u^0}{\partial x_j} + \frac{\partial u^1}{\partial y_j} \right) = \frac{\partial u^0}{\partial y_j} \cdot \frac{\partial a_y}{\partial y_j}. \] (41)

Applying the medium operator (32), the relation (41) can be written:

\[ p^0(x) = a^0_y \frac{\partial u^0}{\partial x_k}, \]

\[ a^0_y = a^0_y(y) + a^0_y(y) \left( \delta_{jk} + \frac{\partial \chi^k}{\partial y_j} \right) = \left( a^0_y \right) + \left( a^0_y \frac{\partial \chi^k}{\partial y_j} \right). \] (43)

Therefore, the relation (33) becomes an equation in \( u^0 \) with constant coefficients:

\[ f = -\frac{\partial}{\partial x_i} \left( a^0_y \frac{\partial u^0}{\partial x_k} \right). \] (44)

### 4 Problem Solution for a SMC Material

For a composite material in which the matrix occupies the domain \( Y_1 \) and presents the coefficient \( a^1_y \) and the inclusion occupies the domain \( Y_2 \) with the coefficient \( a^1_y \) separated by a surface \( \Gamma \), the equation (21) must be seen as a distribution.
material is presented, where the fibers bundle is seen as an ellipsoidal inclusion.

![Fig. 6. Structure’s periodicity cell of a SMC material with 27% fibers volume fraction](image)

Let us consider the function \( f(x_1, x_2) = 10 \) in inclusion and \( 1 \) in matrix. To determine the upper and the lower limit of the homogenized coefficients, first the arithmetic mean as a function of \( x_2 \)-axis followed by the harmonic mean as a function of \( x_1 \)-axis must be computed.

The lower limit is obtained computing first the harmonic mean as a function of \( x_1 \)-axis and then the arithmetic mean as a function of \( x_2 \)-axis. If we denote with \((x_1)\) the arithmetic mean against \( x_2 \)-axis of the function \( f(x_1, x_2) \), it follows:

\[
(x_1) = \frac{\int_{x_2=0.5}^{x_2=0.5} f(x_1, x_2) \, dx_2}{0.5},
\]

for \( x_1 = 0.5; 0.45; 0.45; 0.5 \).

\[
(x_1) = \frac{\int_{x_2=0.5}^{x_2=0.5} f(x_1, x_2) \, dx_2}{0.5},
\]

for \( x_1 = 0.45; 0.45 \).

The upper limit is obtained computing the harmonic mean of the function \((x_2)\):

\[
a = \frac{\int_{x_1=0.5}^{x_1=0.5} 1}{0.5} \frac{\int_{x_2=0.5}^{x_2=0.5} f(x_1, x_2) \, dx_2}{0.5},
\]

for \( x_2 = 0.19; 0.19 \).

The lower limit will be given by the arithmetic mean of the function \((x_2)\):

\[
a = \frac{\int_{x_1=0.5}^{x_1=0.5} dx_2}{0.5},
\]

Since the ellipsoidal inclusion of the SMC structure may vary angular against the axes’ centre, the upper and lower limits of the homogenized coefficients will vary as a function of the intersection points coordinates of the ellipses, with the axes \( x_1 \) and \( x_2 \) of the periodicity cell (fig. 7).

![Fig. 7. ± 30° angular variation of the ellipsoidal inclusion](image)

The following micrographs (fig. 8) make obvious this angular variation of the fibers’ bundles and the extreme heterogeneity and the layered structure of these materials as well as the glass fibers and fillers distribution. The micrographs show that there are areas between 100 – 200 m in which the glass fibers are missing and areas where the fibers distribution is very high.
5 Results

Typical elasticity properties of the SMC isotropic compounds and the composite structural features are presented in table 1.

Table 1.

<table>
<thead>
<tr>
<th>Property</th>
<th>UP resin</th>
<th>Fiber (E-glass)</th>
<th>Filler (CaCO₃)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Young modulus $E$ [GPa]</td>
<td>3.52</td>
<td>73</td>
<td>47.8</td>
</tr>
<tr>
<td>Shear modulus $G$ [GPa]</td>
<td>1.38</td>
<td>27.8</td>
<td>18.1</td>
</tr>
<tr>
<td>Volume fraction [%]</td>
<td>30</td>
<td>27</td>
<td>43</td>
</tr>
</tbody>
</table>

According to equations (10) and (7), the longitudinal elasticity moduli $E_{SM}$ (for the substitute matrix) and $E_C$ (for the entire composite) can be computed. A comparison between these moduli and experimental data is presented in fig. 9.

In practice, due to technical reasons, the fraction of each isotropic compound is expressed as percent of weight, so that the dependence between volume- and weight fraction can be determined:

$$\varphi = \frac{1}{1 + \frac{1 - \psi}{\psi} \cdot \frac{\rho_F}{\rho_{SM}}}$$

where $\varphi$ and $\psi$ are the volume- respective the weight fraction, $\rho_F$ as well as $\rho_{SM}$ are the fibers- respective the substitute matrix density.

---

Fig. 8. Micrographs of various SMCs taken in-plane and perpendicular to their thickness [15]

Fig. 9. Young moduli $E_{SM}$ and $E_C$ for a 27% fibers volume fraction SMC material
From fig. 9, it can be noticed that the Young modulus for the entire composite is closer to the experimental value unlike the Young modulus for the substitute matrix. This means that the rule of mixture used in equation (7) give better results than the inverse rule of mixture presented in equation (10), in which the basic elastic property of the filler and the filler volume fraction can be replaced with fibers Young modulus and fibers volume fraction, appropriate for a good comparison.

According to equations (47) and (50), the upper and lower limits of the homogenized coefficients for a 27% fibers volume fraction SMC material are computed and shown in table 2.

Table 2: Upper and lower limits of the homogenized coefficients for a 27% fibers volume fraction SMC material [17]

<table>
<thead>
<tr>
<th>Angular variation of the ellipsoidal inclusion</th>
<th>Upper limit ( a^+ )</th>
<th>Lower limit ( a_- )</th>
</tr>
</thead>
<tbody>
<tr>
<td>( 0^\circ )</td>
<td>2.52</td>
<td>0.83</td>
</tr>
<tr>
<td>( \pm 15^\circ )</td>
<td>2.37</td>
<td>0.851</td>
</tr>
<tr>
<td>( \pm 30^\circ )</td>
<td>2.17</td>
<td>0.886</td>
</tr>
</tbody>
</table>

The results presented in table 2, show that the upper limit of the homogenized coefficients decreases with the increase of angular variation of the ellipsoidal inclusion unlike the lower limit which increases with the increase of this angular variation.

The material’s coefficients estimation depends both on the basic elasticity properties of the isotropic compounds and the volume fraction of each compound. If we write \( P_M \), the basic elasticity property of the matrix, \( P_F \) and \( P_f \) the basic elasticity property of the fibers respective of the filler, \( \phi_M \) the matrix volume fraction, \( \phi_F \) and \( \phi_f \) the fibers respective the filler volume fraction, then the upper limit of the homogenized coefficients can be estimated computing the arithmetic mean of these basic elasticity properties taking into account the volume fractions of the compounds:

\[
A^+ = \frac{3}{P_M \cdot \phi_M + P_F \cdot \phi_F + P_f \cdot \phi_f}.
\]  

The lower limit of the homogenized elastic coefficients can be estimated computing the harmonic mean of the basic elasticity properties of the isotropic compounds:

\[
A_- = \frac{1}{\frac{1}{P_M \cdot \phi_M} + \frac{1}{P_F \cdot \phi_F} + \frac{1}{P_f \cdot \phi_f}},
\]

where \( P \) and \( A \) can be the Young modulus respective the shear modulus.

Fig. 10 shows the Young moduli and fig. 11 presents the shear moduli of the isotropic SMC compounds as well as the upper and lower limits of the homogenized elastic coefficients.
6 Conclusions
For the same fibers length (e.g. \( l_F = 4.75 \text{ mm} \)) but with a shear stress 10 times greater at the fiber-matrix interface, it results an increase with 18% of the longitudinal break strength of the composite. Therefore, improving the bond between fibers and matrix by using a technology that increases the fibers adhesion to matrix, an increase of composite longitudinal break strength will be achieved.

In the case of using some fibers with greater lengths (e.g. \( l_F = 25.4 \text{ mm} \)), the 10 times increase of the shear stress at the fiber-matrix interface leads to an increase with only 3% of the composite longitudinal break strength. Two SMC composite materials with same shear stress at the fiber-matrix interface (e.g. \( \tau_i = 5 \text{ MPa} \)) but with different fibers lengths, present different longitudinal break strength values, the composite with fibers length \( l_F = 25.4 \text{ mm} \) exhibit an increase with about 16% of this strength. The computing model regarding the longitudinal tensile behaviour of multiphase composite materials like SMCs shows that the composite’s Young modulus computed by help of rule of mixture is closer to experimental data than the inverse rule of mixture.

The presented results suggest that the environmental geometry given through the angular variation of the ellipsoidal domains can leads to different results for same fibers volume fraction. This fact is due to the extreme heterogeneity and anisotropy of these materials. The upper limits of the homogenized elastic coefficients are very close to experimental data, showing that the proposed homogenization method give better results than the computed composite’s Young modulus determined by help of rule of mixture.

The proposed estimation of the homogenized elastic coefficients of pre-impregnated composite materials can be extended to determine elastic properties of any multiphase, heterogeneous and anisotropic composite material.

Future researches will be carried out taking into account the filler’s particles size upon the homogenized elastic coefficients of these materials.

References: