Pressure Effect on Humidity Diffusion through Polyester/Glass Fiber Composite in Sea Water

Mohamed Ounaies*and Hachmi Ben Daly
Laboratory of Mechanics of Sousse, National Engineering School of Sousse, University of Sousse, Tunisia

Abstract
The hygrothermal behavior of the polyester/glass fiber composite is studied under the effect of the pressure in sea water. Fillers additives Aluminum Silicate Pigments are inserted to the composite material. Humidity absorption is made through one lateral side, under different temperature. The humidity concentration is higher on the surface and it continuously decreases toward the core of the material. At high temperature, the humidity absorption curve falls. It is caused by the coalescence of cracks propagating through the composite material, which are created by the stress concentration at a point of the free volume existing in the material. The rise of the percentage of the fillers additives and the pressure induces an augmentation of the diffusion coefficient and the amount of absorbed humidity. The finite element method is utilized to present a numerical model of the humidity absorption process. The comparison between numerical and experimental results confirms that the hygrothermal behavior of the polyester/glass fiber composite could be numerically predicted.

Keywords: Pressure; Sea water; Humidity; Diffusion; Composite

Introduction

Composite material has been more and more utilized in the industrial fields, due to the advantages that offers like high resistance and light weight. However, when it is utilized, composite material is exposed to extreme environmental conditions like temperature and humidity, which have negative effects on the material proprieties. It causes a decrease of the material proprieties and its life durations, which can be fatal in the safety side. Therefore, the utility of studying the hygrothermal behavior of a composite material.

Humidity absorption is explained by two theories: the molecular approach linked to the hydrophilic character of the material. The hydrophilic sites linked to the diffusing molecules through hydrogen bonds [1]. Also, it is explained by the free volume theory, related to the number and size of the free volumes existing in the material. It determines the amount of absorbed humidity [1,2]. The humidity diffusion through composite material causes a swelling and a plasticization of the matrix [2]. Moreover, it induces a decrease of the fibers resistances, a debonding of the matrix/fiber interface [3,4] and a decrease of the transition temperature (Tg) [3,5].

The humidity diffusion is influenced by different parameters like the temperature; [6,7] proved that the augmentation of the temperature induces an increase of the diffusion coefficient and the amount of absorbed humidity, as it provides more agitation to the diffusing molecules [8]. Likewise, the pressure effect in the hygrothermal behavior of a composite material is investigated: [9] demonstrated that the rise of the hydrostatic pressure accelerates the diffusion phenomena, as a result the diffusion parameters increases. These results are confirmed in the case of a polyester/glass fiber composite [10,11] showed that the elevation of the pressure generates an augmentation of the diffusion coefficient and the amount of absorbed humidity.

Furthermore, the humidity absorption kinetics is analyzed: [12] showed that humidity diffusion through the polyester/glass fiber composite follows the Fick’s Law. Likewise, [13] proved that that humidity diffusion through a glass fiber-reinforced plastic composite presents a Fickian behavior at low temperature. Also, [4,13] affirmed that humidity absorption follows the Fick’s law.
Theoretical aspects on absorption in composite material

The humidity diffusion parameters are the diffusion coefficient (D) and the amount of absorbed humidity (M). These parameters are influenced by the temperature [14,15]. The diffusion coefficient is calculated using the following formula [16]:

$$D = \pi^2 \left( \frac{h}{4 \cdot M_m} \right)^2 \left( \frac{M_1 - M_2}{\sqrt{t_1} - \sqrt{t_2}} \right)^2$$ (1)

Where h: the thickness, M_m: the amount of absorbed humidity at saturation and M_1, M_2 are the amount of absorbed humidity at t_1, t_2 respectively.

The humidity diffusion process is described the one dimensional Fick’s second law [17]:

$$\frac{\partial C}{\partial t} - \frac{\partial^2 (D \cdot C)}{\partial x^2} = 0$$ (2)

Where t is the time, D is the diffusion coefficient, C is the concentration of the diffusing and x is the distance through the sample thickness. The diffusion absorption should respect the boundary conditions expressed in equation (3):

$$C = C_i; 0 < x < h; t \leq 0$$
$$C = C_m; x = 0, x = h, t > 0$$ (3)

Where C_i: the initial concentration, h: the sample thickness and C_m: the concentration at the saturation. Jost [18] presents the solution of equation (2):

$$C - C_i = 1 - \sum_{n=1}^{\infty} \frac{1}{2n+1} \sin \left( \frac{(2n+1)\pi x}{h} \right) e^{-\left( \frac{(2n+1)\pi D}{h^2} \right)t}$$ (4)

Where C_i: the initial concentration, h: the sample thickness and C_m: the concentration at the saturation. The concentration gradient from each side of the material surface generates a proportional flow. The flow can be expressed by equation (5) [19]:

$$J = -D \cdot C \cdot \left( \frac{C_i}{s} \cdot V(\theta) + K_s \cdot V(P) \right)$$ (5)

$$J = -D \cdot C \cdot \left( \frac{C_i}{s} \cdot V(\theta) + K_s \cdot V(P) \right)$$ (6)

Where D: the diffusion coefficient, C: the concentration, \( \theta \): the temperature, s: the solubility, P: the pressure, K_s: the pressure stress factor and K_s: the soret effect.

Material and Experimental Methods

Material

The polyester/glass fiber composite is composed of an unsaturated polyester resin reinforced by unidirectional E-glass fiber. The glass fibers are provided by the Owens Corning Fiberglass Company and the polyester is supplied by the Reichold Chemicals Inc Company. Fillers additives, Aluminum Silicate Pigments (ASP400), are inserted to the composite material. It is a clay powder with an average size of 4.8 µm. It is added in order to increase the productivity and decrease the formation of rich resin zones [20]. The composite material is manufactured by the pultrusion technique. It is a continuous composite material manufacturing process [21]. The samples present a dimension of 20*8*6.3mm. Table 1 exposes the volume percentage of the components of the polyester/glass fiber composite. The composition of two varieties (A and B) is presented in (Table 1).

| Material | % ASP400 | % Fiber |
|----------|----------|---------|
| A        | 20%      | 51.5    |
| B        | 40%      | 51.6    |

Experimental methods

Water absorption test: The experimental device is presented in Figure 1, the sample is fixed in the experimental assembly, and the container is filled with sea water, heated to the desired temperature thanks to a heating resistance. The humidity diffuses through one lateral side of the sample, the exposed pressure is fixed by the handle and is displayed on the manometer. Periodically, the sample is removed from the experimental device, dried with an absorbent paper and weighted with a Gibertini E-42-B electronic scale with an accuracy of 0.1mg. After that, the sample is put back in the experimental device. The humidity diffusion process is examined under 2, 3, 5, 9, 20, 30, 40 and 50 bars.

Equation (7) presents the humidity content:

$$M(\%) = \left( \frac{M_t - M_0}{M_0} \right) \times 100$$ (7)

Where M_t is the material weight at instant t and M_0 is the initial material weight.

Simulation of the water absorption: The Fick’s law described the humidity absorption process [19]:

$$\frac{\partial C}{\partial t} = D \cdot \frac{\partial^2 C}{\partial x^2}$$ (8)

Where D is the diffusion coefficient, x is the direction of diffusion and C is the humidity concentration in the material. The
diffusion coefficient \( (D) \) is expressed at each temperature through the Arrhenius law \([22]\):

\[
D = D_0 e^{\frac{E}{RT}} \tag{9}
\]

where \( D_0 \) is the Arrhenius constant, \( T \) is the temperature, \( R \) is the perfect gas constant and \( E \) is the activation energy. In order to simulate the hygrothermal behavior of the polyester glass fiber composite, the finite element method is employed. The humidity diffusion in made through one lateral side and the initial humidity concentration inside the material is equal to zero. A DC3D8, a three-dimensional solid contained in the heat transfer/mass diffusion element library \([23]\), is used. The UTEMP subroutine ensured the thermo-diffusion coupling. The results are normalized as regards to the thickness.

**Discussion**

In this section, we present a comparison of the experimental and numerical results of the humidity diffusion through the polyester/glass fiber composite. (Figures 2-7) present a comparison between experimental and numerical results of the humidity diffusion through the material (A) and material (B), respectively.

**Figure 2:** Humidity absorption curve over the time, normalized to the thickness of the material sample (A) at 25°C:
- a. 2/20 bar,
- b. 3/30 bar,
- c. 5/40 bar,
- d. 9/50 bar.
Figure 3: Humidity absorption curve over the time, normalized to the thickness of the material sample (A) at 45°C:

a. 2/20 bar,
b. 3/30 bar,
c. 5/40 bar,
d. 9/50 bar.
Figure 4: Humidity absorption curve over the time, normalized to the thickness of the material sample (A) at 65°C:

a. 2/20 bar,
b. 3/30 bar,
c. 5/40 bar,
d. 9/50 bar.
Figure 5: Humidity absorption curve over the time, normalized to the thickness of the materials sample (B) at 25°C:

a. 2/20 bar,

b. 3/30 bar,

c. 5/40 bar,

d. 9/50 bar.
Figure 6: Humidity absorption curve over the time, normalized to the thickness of the material sample (B) at 45°C:

a. 2/20 bar,
b. 3/30 bar,
c. 5/40 bar,
d. 9/50 bar.
Figure 7: Humidity absorption curve over the time, normalized to the thickness of the materials sample (B) at 65°C:

a. 2/20 bar,
b. 3/30 bar,
c. 5/40 bar,
d. 9/50 bar.
The humidity absorption presents a two-parts behavior: a first linear part, during which an equilibration of the concentration gradient takes place on each side of the material surface. The second part is concave to the axis of the abscissas. It is a phase of physical and chemical equilibrium inside the material. At a temperature equal or superior to 65 °C, we note a fall of the absorption curve. In fact, when osmotic pressure exceeds a critical value, the stress concentration at a point of the free volume induces the propagation of cracks inside the material. When the cracks coalesce, certain molecular chains get loose. Therefore, the absorption curve falls [24].

The rise in the pressure improves the diffusion coefficient and the amount of absorbed humidity [25]. The pressure elevates the humidity flow diffusing inside the material. Consequently, the humidity diffuses rapidly [26]. Also, the augmentation of the temperature induces an increase of the diffusion parameters, as the temperature offers more agitation to the diffusing molecules [27].

Material (B), having a higher percentage of fillers additives, absorbs a higher amount of humidity than material (A). In fact, the fillers additives reduce the rigidity of the interface matrix/fiber, facilitating the absorption process.

Conclusion

This research presents an analysis of the humidity diffusion through polyester/glass fiber composite. The pressure effect on the hygrothermal behavior of the composite material was examined. At high temperature, the humidity absorption presents a fall; it is due to the coalescence of cracks, caused by the stress concentration at the free volume. As a result, some molecular chains get loose. The augmentation of the pressure increases the diffusion coefficient and the amount of absorbed humidity. Also, the rise of the temperature and the percentage of fillers additives induce a rise of the diffusion parameters.

The finite element method was used to numerically reproduce the experimental results of the diffusion process. The comparison between experimental and numerical results showed that the humidity diffusion behavior through the polyester glass fiber composite can be predicted. As a perspective, we are investigating the pressure direction on humidity absorption.

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