Study of Raw and Recycled Polyethylene Terephthalate by Meaning of TGA and Computer Simulation

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1. Introduction

Decomposition of polymers is an important process [1]. The study of the pyrolysis process of polymers by using nonisothermal thermogravimetric analysis (TGA) was widely known in the last years [2–4]. Compared to other methods, pyrolysis procedure has more strong points: (i) requires the lowest temperature, (ii) without the need of oxygen, and (iii) fabrication of outstanding quality oil [5]. A good knowledge of the pyrolysis process will be useful and helpful element to the design and operation of pyrolysis reactors with developed efficiency [6]. To find the kinetic parameters of pyrolysis process, different models have been found [7]. Among them, we can mention the distributed activation energy model (DAEM) that is used in this paper.

Miura [8] and Miura and Maki [9] found a new procedure based on the modification of the classic DAEM method to get the activation energy \(E_a\) and the frequency factor \(k_0\) by following simple steps [10].

Recycled polymers have many advantages, such as the preserving of the environment, saving energy, and its low cost. However, due to what is common about recycled plastics that it is not good and loses its physical properties, we have compared some properties of the raw PET with the properties of recycled PET, using both of the experimental and the computer simulation. In the best of our knowledge, these elements have been not studied in other paper with the same procedure, which makes our work new in the field of polymer study.

Polyethylene terephthalate (PET) is a thermoplastic polymer vastly used in the form of fiber, sheet, and film [11]. Polyethylene terephthalate (PET) has good mechanical strength, great stability with body fluids, and strong radiation resistance, which make it appropriate for sterilization [12]. PET is widely used in many fields such as electrical engineering, water and soft drink bottle industries, and transportation and building industries, due to its physical and chemical properties [13, 14].

Many TGA tests of our polymers under five heating rates (10, 15, 20, 25, and 30 K/min) were conducted, and the found results were used to find the values of pyrolysis kinetic parameters (the activation energy \(E_a\) and the dependent frequency factor \(k_0\)) by using the distributed activation energy model (DAEM) modified by Miura and Maki.

In order to improve our work, the glass transition temperature (Tg) of PET was studied by meaning of molecular...
dynamic (MD) simulation. The results obtained by MD simulation were compared with the experimental results in other papers, the simulated values of other works, and with the results calculated by using the semiempirical method Synthia [15]. In addition, the role of some energy components (the potential energy, the kinetic energy, and the total energy components) in the glass transition temperature (Tg) process was studied, and the predicted elastic properties by the MD simulations of PET at 300°K were computed.

Many previous works have proved that molecular dynamic simulation gives results which are near to the experimental values in the study of the properties of materials [16–18].

2. Experimental

In this work, the raw polyethylene terephthalate (PET) and the recycled PET were studied by using nonisothermal thermogravimetric procedure for five different heating rates (10, 15, 20, 25, and 30 K/min) for each dry sample of 1 g. The pyrolysis experiences were conducted in thermogravimetric analyzer TGA Q500 from TA Instruments. In the goal to maintain the raw PET and recycled PET in inert atmosphere and make sure that there is no oxygen during all pyrolysis process, a flow of nitrogen of 60 ml/min is applied. The temperature profile fixed to the TGA contains two basic steps: (i) a drying process of the element at 105°C for a duration of 20 min and (ii) the pyrolysis process happen when increasing the temperature of the polymer in the oven up to 750°C at a fixed heating rate in inert atmosphere. In this study, uncertainty in the degree of temperature of our devices is ±1°C, the temperature precision is ±0.1°C, the weighing precision of the nonisothermal thermogravimetric analysis is ±0.01%, and its mass measurement sensitivity is 0.1 μg [6, 7].

Without using mathematic, we can obtain the very proper kinetic parameters (the distribution curve of activation energy f(E) and the activation energy dependent frequency factor (k0)) of polymer pyrolysis by applying the distributed activation energy model (DAEM) modified by Miura [8] and Miura and Maki [9]. This procedure is based on the modification of the classic DAEM method to get the activation energy (Ea) and the frequency factor (k0) by following a few points:

(a) Finding the devolatilization rate for different heating rates
(b) Compute and trace the curve ln (a/T^2) versus 1/T at each rate of devolatilization
(c) Finding the activation energy (Ea) and the frequency factor (k0) from the Arrhenius plots at each rate of devolatilization
(d) Trace the values of activation energy (Ea) versus the rate of devolatilization to obtain the function f(Ea)
(e) Predict other values of devolatilization rate curves for other heating rates. [10]

3. Simulations Methods

In this paper, all molecular dynamic (MD) simulations were computed by using the Materials Studio6.0 software package of Accelrys, Inc. [19].

Molecular mechanics (MM) correspond to a set of empirical procedure for calculation of molecular structures and different physical properties using atoms as the basic elements. It analyzes the progress of a force field (ff) in which polymer chain can be relaxed (energy minimization or geometry optimization) or in which they can be in movement according to classical motional equations (MD) [20]. The choice of the force field plays an important role in influencing on the value of the final results. In our study, COMPASS has been chosen as a force field.

The PET polymer structure with 20 repeat units is embedded in a cell with periodic boundary conditions (Figure 1). To relax our polymer chain, a drastic minimization is applied by using Conjugate Gradient with 20000 iterations to confirm that our polymer chain does not contain any undesirable interactions.

To ensure the molecular relaxation of PET polymer chain, a series of molecular dynamics of 100 ps (using Verlet-leapfrog algorithm) with the canonical ensemble NPT were applied [21]. First, a NVT dynamic is applied in order to stabilize our polymer chain. Then, the dilatometric procedure is applied to get the glass transition temperature (Tg), applying the ensemble NPT by fixing the pressure to 0.0001 GPa and the temperature at 300°K, to find the density of equilibrium that it is employed to get the specific volume. This last can be employed to compute the glass transition temperature (Tg) of PET. Our structure will overheat from 300°K to 800°K and decreases from 800°K to 200°K with a step of 50°K during 100 ps for each temperature.

Furthermore, the elastic properties of the raw polyethylene terephthalate (PET) were studied using the molecular dynamic MD simulation at 300°K. The Smart algorithm was used, and COMPASS was chosen as a force field. The bulk modulus and shear modulus were simulated by using the Reuss, Voigt, and Hill methods. The internal stress tensor is calculated by using the relation:

$$\sigma = -\frac{1}{V} \left[ \sum_{i=1}^{n} a_i (v_i v_i^T) + \sum_{i<k} p_{ik} w_{ik}^T \right],$$

(1)

where i is the number of particle; a_i, v_i, and w_i are the mass, velocity of the particle, and force acting on the particle, respectively; V is the volume of the structure [18, 22].

4. Results and Discussion

4.1. Pyrolysis Process. Figure 2 shows the devolatilization curves versus temperature of both raw PET (a) and recycled PET (b) for the five different heating rates (10, 15, 20, 25, and 30 K/min).

From the curve obtained in Figure 2, a ln (a/T^2) versus 1/T plot of structures: raw PET (a) and recycled PET (b) for each heating rate, at different values of the devolatilization
Figure 1: Molecular model of PET before (a) and after construction (b).

Figure 2: Devolatilization rate of the PET (a) and recycle PET (b) samples.

Figure 3: Arrhenius plot obtained using conversion rate intervals of 5% for the PET (a) and recycled PET (b) samples.
rate was plotted in Figure 3 by using conversion rate range of 5% (5%-95%) to give a good visualization. The Arrhenius plots of raw PET and recycled PET are almost similar.

The activation energy ($E_a$) and the frequency factor ($k_0$) of our structures can be given from the slope and intercept in the Arrhenius plots at each rate of devolatilization using the linear fitting and with the two relations:

$$E_a - mR, \quad (2)$$

$$k_0 = -m \exp (n - 0.6075), \quad (3)$$

where $T$ is the temperature and $R$ the universal constant.

The activation energy ($E_a$) (a) and the frequency factor ($k_0$) (b) obtained from the raw PET and recycled PET samples are shown in Figure 4. We can remark that the activation energy ($E_a$) found for the raw PET is in the range of 120–150 kJ/mol, and the frequency factor ($k_0$) was in the range of $10^7$–$10^9$ s$^{-1}$, which is in accordance with the results obtained in other studies [23, 24]. For recycled PET, the activation energy ($E_a$) found is in the range of 159–185 kJ/mol, and the frequency factor ($k_0$) was in the range of $10^{10}$–$10^{11}$ s$^{-1}$.

4.2. Molecular Dynamic Simulation

4.2.1. Glass Transition Temperature (Tg). The glass transition temperature (Tg) of a polymer is an essential phenomenon which can have different values. There is no exact method to find the glass transition temperature [25]. Currently, the most used procedure is the free volume theory proposed by Fox and Flory [26].

Figure 5 shows the curve of the specific volume versus the temperature of the raw PET. The specific volume (sp) is obtained from the NPT trajectory file. By the analysis of the curve of specific volumes vs. the temperature, we detect the simulated glass transition temperature of PET (about 358.34 K) that is little higher than the experimental value (346.6 K [27]) and smaller than the simulated value of the glass transition temperature (Tg) of PET in other papers (about 362 K [28]).

In addition, the glass transition temperature (Tg) of PET was computed by using the semiempirical method Synthia. Table 1 shows the simulated glass transition temperature of PET by using the molecular dynamic (MD) simulation, Synthia and the simulated value of other study, and experimental glass transition temperatures of PET.

4.2.2. Roles of the Energy Components in the Tg Process. In order to study the roles of the different elements in the glass transition temperature process, some energy components versus the temperature are shown in Figure 6. Many previous works have studied the role of the energy components in the transition glass temperature process for different structures [30, 31].

In this paper, the potential energy, the kinetic energy, and the total energy components decrease linearly with temperature decreasing, and there are no any breaks at any point in the curves of the energies versus the temperature. This shows that the potential energy and the kinetic energy play important roles during the glass transition process.
4.2.3. Mechanical Properties. The predicted elastic properties by the molecular dynamic (MD) simulations with COMPASS force field of PET at 300 °K are given in Table 2.

By the analysis of the table, we can remark that our value of Young’s modulus ($E$) of PET is near to the available result with a difference of just 0.03 GPa. Both of bulk modulus and Poisson’s ratio are very near to the values obtained in other works. The shear modulus ($G$) simulated in our study is a little different to the found shear modulus in other paper.

5. Conclusion

Thermogravimetric curves were used to study the pyrolysis process of raw PET and recycled PET by using DAEM simplified by Miura and Maki. Devolatilization rate of the PET and recycle PET samples is almost similar. The activation energy ($E_a$) found for the raw PET is in the range of 120–150 kJ/mol, and the frequency factor ($k_0$) was in the range of $10^7$–$10^9$ s$^{-1}$, which is in accordance with the results.
Table 2: Young’s modulus (E), shear modulus (G), bulk modulus (K), and Poisson’s ratio (υ) at 300 K of PET.

| Polymer | E   | G   | K   | υ   | E   | G   | K   | υ   |
|---------|-----|-----|-----|-----|-----|-----|-----|-----|
| PET     | 2.7 | 0.95| 6.43| 0.41| 3   | 0.02| 6.3 | 0.2 |

aValue from Ref [32]. bValue from Ref [33]. cValue from Ref [34]. dValue from Ref [35].

obtained in other studies. For recycled PET, the activation energy (E_a) found is in the range of 159–185 kJ/mol, and the frequency factor (k_0) was in the range of 10^{10}–10^{11} s^{-1}. From all above results, it can be concluded that the recycled PET is more cohesive than the raw PET contrary to popular belief.

Furthermore, the glass transition temperature (T_g) of PET was studied by means of molecular dynamic (MD) simulation. By the analysis of the curve of specific volumes vs. the temperature, we find that the T_g of PET is 358.34 K that is between the experimental value (346.6 K) and the simulated value of T_g in other study (about 362 K). In the addition, the role of some energy components was analyzed. The potential energy and the kinetic energy play an important role during the glass transition process. Our simulated results of mechanical properties of PET were very near to the available values in literature.

From the comparison of our simulated results with other values in different papers, we can conclude that our methods of simulation give good results.

Data Availability

No data were used to support this study.

Conflicts of Interest

All the authors declare no conflict of interest.

Acknowledgments

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