1. Introduction

Nowadays polyethylene terephthalate (PET) is one of the most common plastics, because of its widespread use. There are certain places where recycled PET is already used and produced. Our goal is to increase the quantity of the recycled PET. From the recycled materials the new bottles, products can be made. This requires a development path and a more sophisticated technology since it is not easy to determine for once used bottle what was stored in it, how it was used previously [1]. PET is a hygroscopic thermoplastic polymer, therefore drying process is very important, since it affects the rheological properties of the material, the subsequent processing, and the product characteristics [2]. The rheological analysis mentioned in this paper is of polyethylene terephthalate (PET) base material, in which a specific raw material - a melt viscosity of structural, elastic and viscoelastic behavior was investigated. The study was carried out by ARES-G2 type oscillational rheometer, and Goettfert Rheograph 25 capillary rheometer with which the material was examined at low and high shear rate. The original granule were analyzed between 270-290 °C. Then, the original raw material was processed BT injection molding machine and then it was granulated again. There was a full investigation at the same temperatures. The measuring values are presented on diagrams and from these the conclusions were drawn.

2. Material

RAMAPET N180 is a general purpose, non-reheat PET resin for bottles, film, thermoformed containers and other transparent applications. It has good optical properties, the mechanical properties are representative for PET resin. The PET is hygroscopic in little extent, and since moisture adversely affects the IV during the melt processing of the polymer, it must be dried prior to molding. Undried PET pellets can contain up to 0.2 % (2000 ppm) of moisture. After drying, the moisture content is typically below 50 ppm [3].

PET has good surface hardness, stiffness, and dimensional stability. It has good gas barrier properties and good chemical resistance except for alkaline materials (which hydrolyze it). Its structure varies from amorphous to fairly high crystalline; it can be highly transparent and colorless but thicker sections are usually opaque and off - white. The PET material is heat - resistant until 180 °C, it does not decrease in strength, its melting temperature is around 260 °C. Above 250 °C the crystallite section is continuously disappearing, so the melting process of the material could be observed. Usually, the material is processed in the molten state, so it is worth to study the rheological properties only above this temperature [4]. The measurements were performed at 270 °C, 280 °C and 290 °C.
3. Method

3.1 The rotational viscometry

Our measurements were carried out by a shear viscosity investigation method; that means a low range of shear rate and temperature steps near the processing temperature. This was the reason why the rotational viscometry was measured in the low shear rate range (0.05-20 \(1/s\)). With the rotational viscometry we could measure torsion flow in a polymer melt between parallel plates. The measuring technique conceptual layout is shown in Figure 1. The CMT means combined motor and transducer concept [5].

From the marking system the stress, deformation and deformation rate components can be calculated with the equations described below:

\[
\sigma = \frac{2}{\pi r} \times M \tag{1}
\]

\[
\gamma = \frac{L}{h} \times \theta \tag{2}
\]

\[
\dot{\gamma} = \frac{r}{L} \times \Omega \tag{3}
\]

where \(r\) is the circular plate radius; \(h\) is the gap between the plates; \(\theta\) is the driving engine rotation (rad); \(\Omega\) is the driving engine angular speed (rad/s) and \(M\) is the torque (\(\mu\)Nm) [5].

The stress and deformation rate can be calculated from Equations (1) to (3) and the shear viscosity can be defined. It is a frequent measuring technique to determine shear viscosity for small molecular liquids, or oils and thermoplastics at low shear rate. ARES G2 is equipped with Forced Convention Oven (FCO). With this accessory materials can be examined at higher temperature (to 600 °C) and the degradation process can be monitored. For our testing TA Ares G2 type rotation viscometer was used with cone - plate geometry in 25 mm diameter. The trim gap was 0.09 mm, while the measuring gap was 0.04 mm. SMT (Separate motor and transducer) layout was used. In this type of rotation viscometer, the drive engine and the transducer are separated [6].

3.2 The capillary rheometry

The capillary rheometer is a measuring device in which the shear viscosity of thermoplastics can be measured. The shear rate ranges were between 100 and 10,000 \(1/s\) near to the processing temperature [7]. The Goettfert Rheograph 25 capillary rheometer was used for the measurements. The measuring method is based on continuous melt flow in known geometry capillary (dies), while the pressure drops are measured between the capillary inlet side and the atmospheric pressure (Figure 2).

The variable parameters were the barrel temperature and the piston speed (which means the deformation rate). In this case, the shear viscosity depends on temperature and deformation (shear) rate. In our tests, the shear rate was changed between 100-10,000 \(1/s\) and the barrel temperatures were changed in three steps around processing temperature, that is for the material of the supplier recommended, at 270-290 °C. The measuring method used two other geometry capillaries because of the entrance and exit effects (pressure drop). With this equipment in one step the Bagley correction can be done, which corrects the pressure drop and gives the real shear viscosity [8].

4. Results

4.1 The rotational viscometry

The viscosity curves were determined in function of the shear rate. The measurements were performed with original material and with recycled material. The viscosity of the original material is shown in Figure 3 at different temperatures.

The curves measured at different temperatures are not the same. It can be observed that the viscosity is the highest at 270 °C. The higher is the temperature, the lower is the viscosity. From the differences of the curves, the transformation of the molecular structure can be determined. The viscosity of the recycled materials is shown in Figure 4.
The measurements were performed with original material and with recycled material. The viscosities of the original and recycled materials are shown in Figure 6. Comparing the data to the 270 °C viscosity curve in the lower shear range: There is a ~30-35 % reduction at 280 °C, and at 290 °C we can see a 40-50 % reduction. As we approach the larger shear ranges (10000-18000 1/s) the differences are reduced: at 280 °C ~15-18 %, while at 290 °C ~7-3 %. In the initial shear speed range, the recycled material shows a 70 % reduction at 270 °C, while in the higher shear range this value shows a 15-30 % reduction. We could observe the following viscosity reductions in the results: at 280 °C, ~ 65 % and at 290 °C ~ 50 %. Although the initial viscosity values at 290 °C show a different tendency compared to the curve measured at the previous two temperatures, this can be attributed to a measurement error with a great certainty. From these values we can conclude that we have to consider the viscosity change of the material during the recycling.

4.2 The capillary rheometer

The measurements were performed with original material and with recycled material. The viscosities of the original and recycled materials are shown in Figure 6.

Increasing the temperature the viscosity decreases. During the recycling and grinding processes, the molecules in the materials can easily degrade. This is caused by the rising thermal and mechanical strain during the processes. The degradation makes the molecular weight lower, so the molecular density dispersion widens. The polymer with shorter chains can move easier. Above the glass transition temperature the segments of the polymer move increasingly. The viscosities of the original and recycled materials are shown in Figure 5.

In each diagram the viscosity curves show a typical form (Figure 5). In the curves of newtonian fluids it could be recognised that the curves are almost linear and do not depend on the shear rate. In our measurements, the viscosity curves approach the horizontal line, but they decrease within a small range. It was confirmed that these materials behave like non newtonian fluids.
because with such a degree of deviation we could assume that the material will not be able to withstand the requirements, which are set by design.

5. Conclusion

In our investigation one type of material was measured, but at different temperatures and in different conditions. An original and a recycled material were examined. The measurements were performed by an ARES G2 rotational rheometer and a Goettfert Rheograph 25 capillary rheometer at 270 °C, at 280 °C and at 290 °C. The shear rate was changed from 0.05 to 20 1/s and 100-10 000 1/s. It was determined that the viscosity of the recycled material is lower than the value of the original material at each temperature. The molecules of the recycled material were supposed to shorten during the grinding process. If the temperature increases, the shear viscosity decreases, but there was no linear correlation between these two variables. The structure viscous behavior of the polymer melt was revealed. This area offers lot of possibilities for further research, so we intend to continue investigations.

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