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Article

# Optical Study of Structural/Electronic Property Change in Thin Polyethylene Terephthalate Film By Stretching

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## Abstract

Optical spectroscopy provides several useful information about polymeric ultrathin films by combining interferometric and optical absorption data contained in the UV-Vis-NIR spectra. In particular, the UV-Vis-NIR spectrum of an ultrathin polymeric film contains information about the film thickness, structural disorder, bandgap energy, type of electron transition model (direct/indirect, allowed/forbidden), cutoff wavelength (i.e., the opaque/transparent switching wavelength), etc. Here, these properties have been determined for a model semi-crystalline polymer (polyethylene terephthalate, PET) in form of ultrathin film before and after a mild mechanical deformation treatment (manual stretching). It has been found that  $E_U$  and  $E_g$  parameters are not strictly depending on mechanical deformation due to their main dependence on chemical composition/constitution of the polymer; consequently  $E_g$  can be used for polymer identification in the case it has a dielectric nature.

**Keywords:** Mylar; polyethylene terephthalate; ultrathin film; interferometry; optical absorption coefficient; optical bandgap; Tauc plot; Urbach energy; chemical analysis

## 1. Introduction

Absorption optical spectroscopy (UV-Vis-NIR spectroscopy) provides important physical information about dielectric polymers at solid state [1,2] and can allow their identification. The electronic structure of amorphous or semi-crystalline dielectric polymers is approximately described by a model of band-structure characterized by a high content of interstitial levels (i.e., localized tail states usually named trap states) [3-5]. Bandgap energy ( $E_g$ ) is the fundamental feature of band-structures (it is equivalent to Lewis shell excitation energies in atoms) and it can be determined with high accuracy by using absorption optical spectroscopy data [6]. Yet, Urbach energy,  $E_U$ , which measures electronic disorder in solids with a band-structure can also be determined by the same optical spectroscopy approach [6]. Important polymer behaviors can be understood on the basis of these two physical quantities deriving from optical spectra. For example,  $E_g$  is correlated to relevant polymer characteristics like photostability, thermal stability, dielectric strength, electrical polarizability [7] and various optical parameters like the cutoff wavelength,  $\lambda_g=hc/E_g$ . On the other hand,  $E_U$  measures structural disorder in polymers (e.g., crystallinity degree, level of chain entanglement in the amorphous phase, impurity content, porosity, surface roughness, etc.). Both parameters are obtained by analyzing the behavior of the polymer absorption coefficient with photon energy in the fundamental absorption edge spectral region. This absorption coefficient is a determining factor for optical material applications (e.g., optical limiters) and has a key role in the optical analysis of polymers and other optical-grade solid media (e.g., silicon glass, semiconductor oxides), since it measures how strongly a material absorbs photons of a specific energy. According to the Lambert exponential law for homogeneous optical solids (i.e.,  $I(d)=I_0 \cdot \exp(-\alpha \cdot d)=I_{inc} \cdot (1-R)^2 \cdot \exp(-\alpha \cdot d)$ , where  $I_{inc}$  is the incident light intensity,  $R$  the reflectance (Fresnel loss),  $I(d)$  is the light intensity

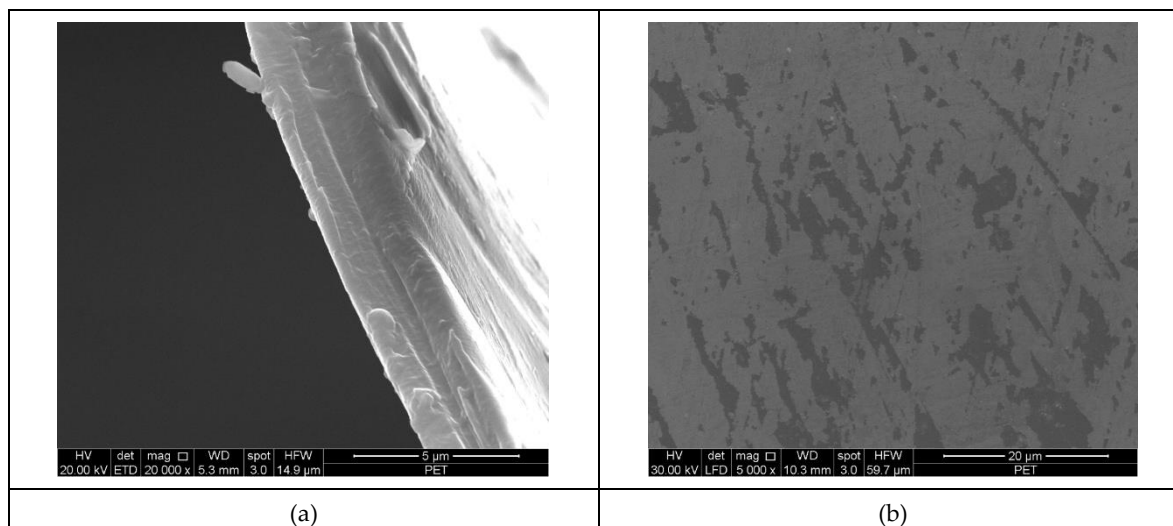
for a  $d$ -thick film, and  $\alpha$  is the absorption coefficient [6,8]), the absorption coefficient quantifies the fraction of light intensity lost per distance unit, as radiation propagates through the material. Since the beam positioning inside an UV-Vis spectrophotometer always is perpendicular to the sample (thin film) surface, reflectance  $R$  is given by the Fresnel equation  $R=(n_1-n_2/n_1+n_2)^2$ , that provides for example for an air-PET interface a Fresnel loss value of 0.04 (with  $n_{\text{air}}=1$  and  $n_{\text{PET}}=1.575$ ). Therefore, the quantity,  $1-R$ , is close to one ( $1-R=0.95$ ) and the Lambert law can be simplified (also scattering is assumed as negligible). In particular, introducing a physical parameter named total absorbance,  $A_{\text{tot}}$ , defined as  $A_{\text{tot}}=\text{Log}(I_{\text{inc}}/I)$ , the absorption coefficient at a specific wavelength is given by the following logarithmic relationship:  $\alpha(\lambda)=(\ln 10) \cdot [A_{\text{tot}}/d]$ . Both  $A_{\text{tot}}$  and  $d$  can be obtained from optical data. Indeed, total absorbance data are directly recorded in UV-Vis spectral measurements or can be calculated from the transmittance values (i.e.,  $A=-\text{Log}(T\%/100)$ , where  $T\%$  is the transmittance percentage) also provided as spectrophotometer output. In addition, interferometric analysis of spectral oscillations contained in the transparency region of the spectrum allows very accurate film thickness ( $d$ ) calculation (in the case the polymer refractive index is known). These optical data are combined together and photon wavelength is converted to energy by the modified Planck equation (i.e.,  $E(\text{eV})=1,240/\lambda(\text{nm})$ , where  $\lambda$  is the wavelength of the radiation) to achieve the  $\alpha(E)$  graph. In the  $E_u$  and  $E_g$  determination, the behavior of  $\alpha(E)$  is analyzed by using some special types of graphical representations, named Urbach and Tauc plots, respectively.

Here, as an example of this general analysis method, a model system of ultrathin polymeric film consisting in optical-grade Mylar (polyethylene terephthalate, PET) has been investigated. Bi-axially oriented ultrathin PET films allows to fabricate highly transparent and mechanically robust plastic devices with very low X-ray fluorescence, that are widely used as windows in disposable sample cups for X-ray fluorescence spectroscopy (XRF), substrates for electrodes/sensors in wearable electronics, insulator tapes, packaging material for optoelectronic devices, etc. This type of polymer has quite high refractive index (1.575) and Mylar films for XRF applications have enough flat/smooth surfaces, which increases visibility of spectral oscillations. The stretching of these PET films generates thinner layers of uniform thickness that can be analyzed by the same approaches. The optical characterization of these thin films by UV-Vis-NIR absorbance spectroscopy before and after the stretching treatment allows obtaining useful information on the possible film changes in geometric, electronic and structural properties as a consequence of this mechanical deformation.

## 2. Materials and Methods

XRF grade Mylar films (PANalytical B.V., X-Ray film Polyesterpetp, 943050007191) have been investigated as a model of ultrathin polymer film. This type of films are made of biaxially oriented polyethylene terephthalate (PET) and are used as windows in disposable sample cups for X-Ray Fluorescence Spectroscopy (XRF), since they allow maximum transmission of low energy X-rays, high mechanical strength, low X-ray fluorescence, etc. The optical spectra were acquired at room temperature using a double beam UV/Visible spectrophotometer (VWR, UV-6300PC Spectrophotometer, VWR International Europe bvba, Leuven, Belgium). Spectra were recorded in the 190-1100 nm wavelength range, using a low scan speed and a scan step (resolution) of 0.5 nm (2mm slit). The optical spectra were processed by a devoted analysis software (UV-Vis Analyst, Version 5.44), which allowed automatic reading of peaks and valley wavelengths of interferometric fringes (spectral oscillations) present in the spectrum (high transparency region) and the corresponding full-width at half-maximum (FWHM) values. The software also allowed smoothing operations for cleaning the spectrum from these interference fringes and measure the average absorbance value in such part of the optical spectrum. In particular, the PET film was optically characterized without blanking and both reflectance (Fresnel loss) and scattering phenomena were assumed as negligible because of the normal incidence of the spectrophotometer beam on the film surface and low amount of surface defects, respectively. In particular, in order to have a perfectly perpendicular incidence of the spectrophotometer light beam, the films were accurately fixed to the instrument cuvette holder in a planar manner by using adhesive tape.

The ultrathin PET film cross-section and surface have been morphologically characterized by using a scanning electron microscope (SEM, FEI Quanta 200 FEG). According to the SEM characterization of ultrathin Mylar films, they showed uniform thickness and the presence of some surface defects, probably generated during post-processing treatments like rolling and cutting (see Figure 1a,b).



**Figure 1.** SEM micrographs of ultrathin PET film showing cross-section (a) and surface (b).

The semi-crystalline nature of ultrathin PET film has been established by X-ray diffraction (XRD, X'Expert PRO, PANalytical, Oxford, UK) and crystallinity degree has been measured by differential scanning calorimetry (Discovery DSC 2500, Waters TA Instrument).

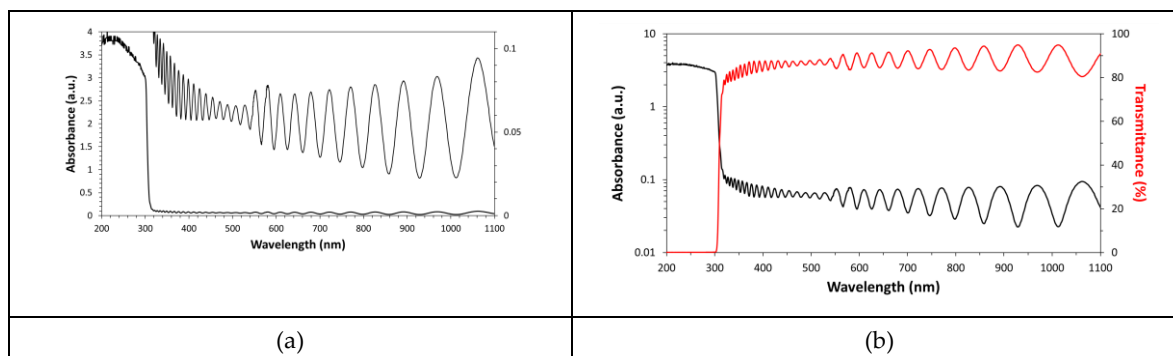
### 3. Results

The interference phenomenon can be clearly detected in the high transparence region of the UV-Vis-NIR spectrum of ultrathin Mylar windows. This physical phenomenon can be used for accurately measuring the film thickness ( $d$  is of the same order as the light wavelength). In particular, the following equation, derived from the interference maxima condition (i.e.,  $2n \cdot d = m \cdot \lambda_m$ ) [9] in the case of a normally incident light beam, can be adopted [10]:

$$d = \frac{\Delta m}{2n} \left( \frac{1}{\lambda_1} - \frac{1}{\lambda_2} \right) \quad (1)$$

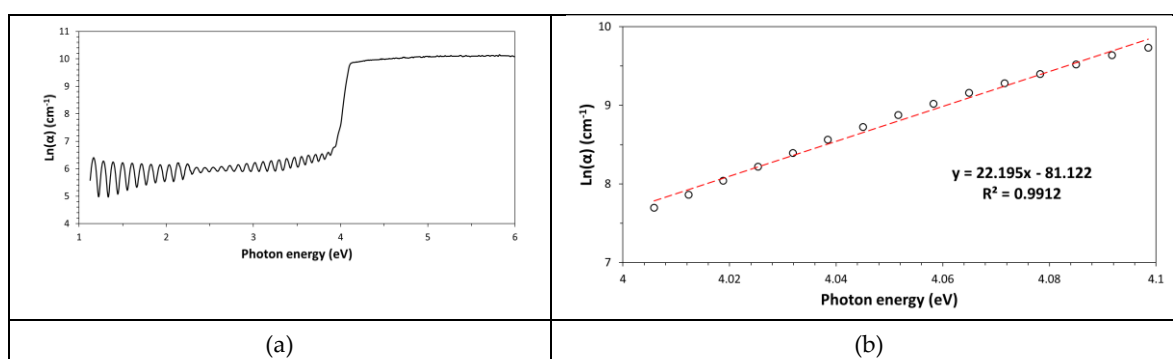
where  $d$  is the film thickness (in nm),  $\Delta m$  is the number of fringes (oscillations maxima) contained in the measurement interval,  $n$  is the average polymer refractive index, and  $\lambda_1$ ,  $\lambda_2$  are the wavelength limits of the measurement interval. According to the spectral oscillations in the ultrathin PET film optical spectrum shown in Figure 2, the film thickness obtained by using the above equation is  $3.665 \mu\text{m}$  (an average refractive index value of 1.575 has been used for calculation [11]).

Since the PET films have a regular shape (i.e., thin circular dishes), this optically measured thickness value has been confirmed by a simple geometrical calculation based on density [12]. In particular, the following expression has been used:  $d = m / (\pi \cdot R^2 \cdot \rho)$ , where  $m$  is the film weight,  $R$  the dish radius and  $\rho$  the PET density. A value of  $3.59 \mu\text{m}$  has been found by using a PET density value of  $1.3 \text{g/cm}^3$  [13]. This thickness value perfectly agrees with value obtained by the interferometric measurement.



**Figure 2.** Optical spectrum of PET thin film expressed as absorbance (a) and in comparison with transmittance (b).

The interferometrically measured film thickness has been used for calculating the absorption coefficient values. It must be pointed out that the absorbance data in the fundamental absorption edge (required for both Urbach and Tauc analysis) are readily accessible for the absorption coefficient calculation because spectral oscillations are present only in the high transparency region of the UV-Vis spectrum. In addition, the UV-Vis Analyst software allows also the application of spectral smoothing and such operation can be conveniently used to reduce fringes in the spectrum and obtain the average absorbance in this spectral region [14]. A graph of  $\ln(\alpha)$  vs. the photon energy is named Urbach plot (see Figure 3a) and this special graph can be used to know the amount of structural disorder in this ultrathin PET film [15]. Structural disorder is proportional to the Urbach energy,  $E_U$ , which is given by the inverse of the slope of the linear part of curve in the Urbach plot. In particular, this quantity can be obtained by linear regression analysis of this straight part of the graph (see Figure 3b). In general, the structural disorder obtained by the Urbach plot depends on all types of defects contained in this thin PET film like for example air bubbles (porosity), surface roughness, chemical impurities (molecular additives like plasticizers, UV-stabilizers, etc.) dissolved in the polymer, percentage of the amorphous phase, etc. However, with respect to perfectly processed films, most contributions can be neglected and the measured structural disorder corresponds exactly to the content of amorphous phase and to the degree of chain entanglement characterizing this amorphous phase. In the case of the ultrathin PET film such calculation led to a value of 45.0meV (with  $R^2=0.9912$ ), which indicates a quite low disorder, probably due to the semi-crystalline nature of this polymer.

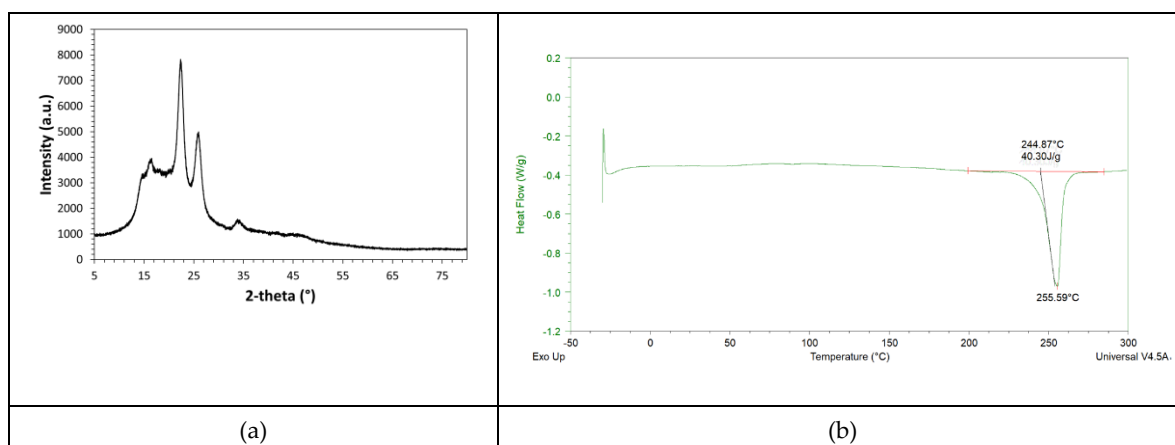


**Figure 3.** PET ultrathin film Urbach plot (a) and best fitting of the linear portion of curve (b).

The small value of  $E_U$  that has been found by applying the Urbach approach to the spectral data of PET films can be well justified on the basis of the semi-crystalline nature of this thermoplastic polymer. Indeed, according to the XRD diffractogram shown in Figure 4a, the polymer contains an amorphous fraction, which causes the presence of a broad diffuse halo in the diffractogram, and a crystalline part, which causes the presence of three main peaks overlapped to the diffuse halo. These peaks are located at  $2\theta$  values of 16.5°, 22.4°, and 25.8°; such diffraction peak positioning is in good

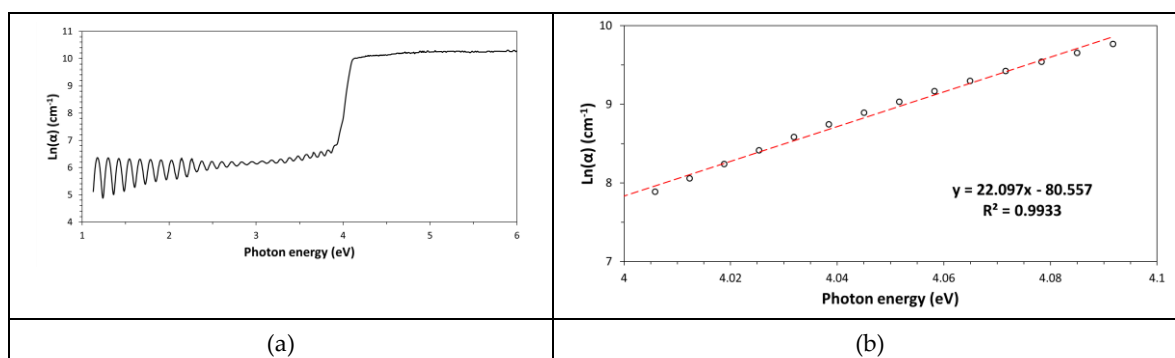
accordance with literature information [16]. The FWHM of diffraction peaks is inversely proportional to the crystallite size (Scherrer equation [16]); consequently, the observed broad diffraction peaks suggest the presence of very small sized crystallites in the polymer, which justifies to observed high optical transparency in the visible spectral region.

The semi-crystalline nature of the ultrathin PET films has been further confirmed by Differential Scanning Calorimetry (DSC). According to the DSC-thermogram shown in Figure 4b, the polymer presents an endothermic signal due to the melting of the crystalline phase and such melting signal appears above the glass transition temperature ( $T_g$ ), which is located at ca. 80.91°C. The PET crystallites melting signal has an onset temperature at ca. 240°C and minimum at 255.59°C. The DSC characterization of the polymer allows to measure the melting enthalpy, which resulted of ca. 40.30J/g. The calorimetric crystallinity degree,  $X_c$ , of the polymeric sample can be determined by comparing the specimen melting enthalpy with the theoretical melting enthalpy of an hypothetical fully crystalline PET sample (ca. 140J/g [17]) and such ratio resulted of ca. 28.8%. This crystallinity value agrees with the value characteristic of bi-axially oriented PET films [18]. Therefore, the small  $E_U$  value experimentally found for the PET sample by the Urbach approach seems to be completely justified by its semi-crystalline nature.



**Figure 4.** XRD diffractogram (a) and DSC thermogram (b) of the ultrathin PET film.

The same type of analysis (Urbach plot) has been applied to a slightly mechanically deformed (manually stretched) ultrathin PET film (see Figure 5a,b). Although, mechanical deformation had the effect to modify the film thickness, the resulting Urbach energy value ( $E_U=45.2\text{meV}$ , with  $R^2=0.9933$ ) remained practically unchanged (see Figure 5b). Therefore, structural disorder did not increase appreciably in this system as a result of the applied stress.



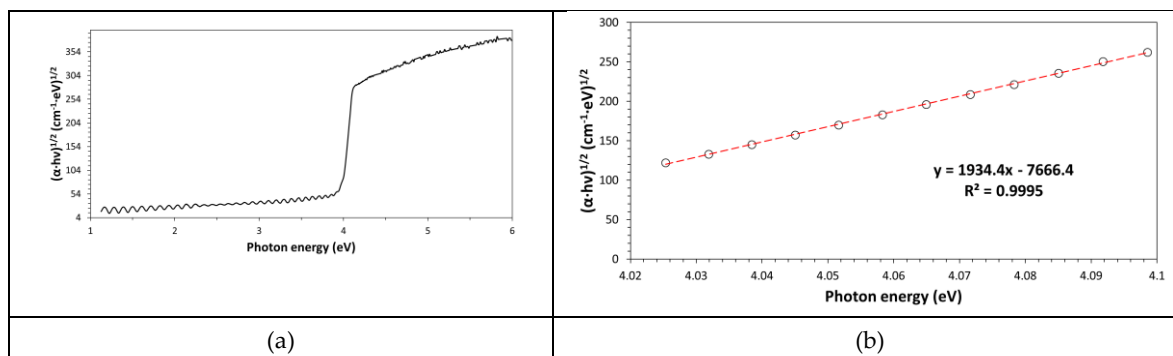
**Figure 5.** Urbach plot (a) and Urbach energy evaluation by linear regression analysis (b) for uniaxially stretched ultrathin PET film.

Ultraviolet radiation interacts with dielectric polymers causing photoexcitation (i.e., optical absorption due to electronic transitions). For some polymers like Kapton-H and polyetherimide (PEI),

even visible photons may cause photoexcitations [19]. The cutoff wavelength has been observed for these ultrathin PET films at ca. 300nm. Consequently, photons with a wavelength lower than 300nm are completely absorbed by the film, while transparency is close to 87.4% at wavelengths higher than 300nm. The type of photoexcitation model and bandgap energy value ( $E_g$ ) can be established optically by using the Tauc plot method.  $E_g$  is an important parameter for polymers since it is related to other physical properties like for example the permittivity ( $E_g$  is inversely proportional to material permittivity, which is the capability of electric field instauration in a material and corresponds to the material polarizability [7]). In addition, accurate  $E_g$  determination allows dielectric polymer identifying (however, identification is not possible with conjugated polymers like polyacetylene because of the  $E_g$  dependence on the conjugation extension and therefore on molecular weight). The bandgap energy has been easily and accurately calculated by using the Tauc plot (see Figure 6a,b), which is a special graphical representation obtained by graphing the quantity  $(\alpha \cdot E)^n$  as a function of the photon energy,  $E$ . The  $n$  value must be selected to allow the linearization of the  $(\alpha \cdot E)^n$  function and  $n$  is 0.5 for an indirect electron transition model and 2 for a direct electron transition model (both models are for allowed electron transitions). Depending on the most convenient value found for  $n$  (the model with best correlation factor), the type of electron transition model can be established. According to the following equation:

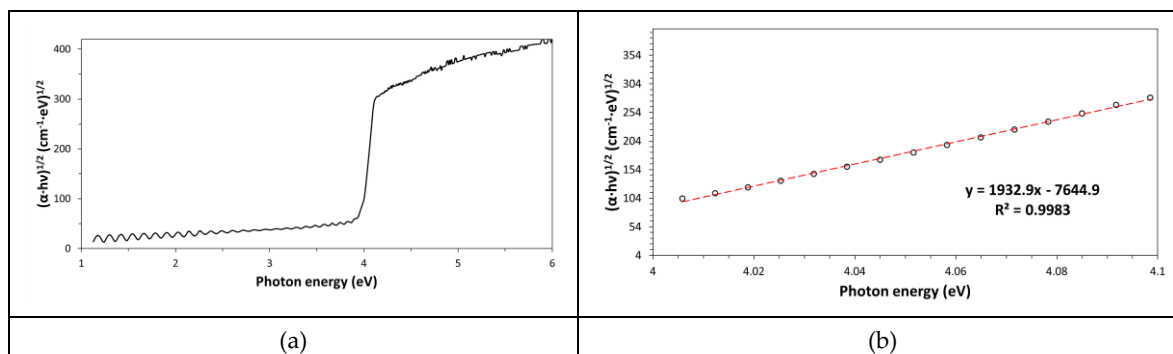
$$(\alpha \cdot E)^n = B \cdot (E - E_g) \quad (2)$$

the value of the intercept with the energy axis corresponds to  $E_g$  (indeed, for  $(\alpha \cdot E)^n = 0$ , it results  $E = E_g$ ). The intercept value is obtained from the best fitting equation of the linear portion of curve by using the following expression:  $-\text{intercept}/\text{slope}$ . In our case, an  $E_g$  value of 3.96eV (with  $R^2=0.9995$ ) has been found for an indirect allowed electron transition model ( $n=0.5$ ), which perfectly agrees with information provided in the literature [20, 21].



**Figure 6.** PET ultrathin film Tauc plot (a) and best fitting of the linear section of the curve (b).

Manual stretching allowed simulating mild uniaxial mechanical stresses that polymeric films typically suffer in service or during industrial uses. This investigation has shown that  $E_g$  is a polymer physical characteristic, which remains practically unchanged after mild mechanical damage undergone by this material (see Figure 7a,b). Indeed,  $E_g$  depends mainly on atom types (composition) and the way atoms are chemically bonded together (constitution) in the repeating unit.



**Figure 7.** Tauc plot (a) and bandgap energy estimation by linear regression analysis (b) for a stretched ultrathin PET film.

According to the achieved experimental results, dielectric polymers at solid-state have a band structure principally depending on the polymer chemical composition/constitution and only marginally on crystallinity, molecular weight and texture (i.e., fibrous/spherulitic morphology). Such behavior allows using the characteristic bandgap energy numerical value to establish polymer nature. Since the very dawn of analytical chemistry, the identification of inorganic compounds (e.g., elemental metals, metal salts) has been based on recognizing the electronic transitions (emission or absorption lines) involving frontier orbitals, that is the valence shell orbitals (the Lewis electron shell). One need only think about the classical 'flame-tests' used by chemists more than one hundred years ago, or the early quartz-prism spectrosopes (Kirchhoff-Bunsen spectroscope), now replaced by Atomic Absorption Spectroscopy (AAS). The yellow color imparted by sodium to the flame, the violet of potassium, the red of calcium, and so on, are well-known phenomena. Thermally excited metals emit radiations that they are capable of absorbing too in the analysis by spectrosopes. Similarly, the precise assignment of the energy of the optical absorption edge (HOMO-LUMO transition energy,  $E_g$ ) of dielectric polymers by UV-Vis spectrometric analysis can enable their reliable identification. This concept cannot be applied to inorganic semiconductors (consider, for instance, how silicon is used - through doping - to fabricate multiple types of semiconductor devices), nor to conductive/semiconductive polymers, due to their electronic configuration being dependent on the extent of conjugation (i.e.,  $E_g = f(M_w)$ ); however, it works perfectly with dielectric polymers.

This chemical analysis approach can be conveniently adopted in fields like microplastics since these types of waste belong to only few, very different plastic classes that can be easily and with safety distinguished on the basis of the  $E_g$  parameter (identifying microplastics is an important aspect, since it allows adequate selection of suitable recycling/remediation processes). UV-Vis spectrophotometers could be very convenient in this research area because they have a very small beam spot and therefore sampling requires only little polymer amounts. Furthermore, these devices usually can have small size, lightweight, and are easily transportable; they are structurally simple and consequently inexpensive, mechanically robust (diode-array devices have no moving parts) and capable of resisting in various environmental conditions. Therefore, optical spectrophotometers are suitable for building field/offshore laboratories to be placed, for example, on boats, on beaches, in landfills, etc. Yet, the chemical analysis of microplastics can be conducted directly in water (aqueous suspensions of microplastics) since this type of molecule is perfectly transparent to visible and ultraviolet light (up to approximately 100 nm).

#### 4. Conclusions

The optical absorption coefficient of dielectric polymers quantifies how much light these synthesis materials absorb per unit distance. This important optical parameter allows analyzing band structure and structural disorder in dielectric polymers and provides many other important physical insights on these materials (e.g., cutoff wavelength, transparency %, spectral opacity range, etc.). In particular, the behavior of absorption coefficient with photon energy,  $E=h\nu$ , allows measuring the optical bandgap energy,  $E_g$ , and the Urbach energy,  $E_u$ , by using the Tauc method and Urbach plot, respectively. The behavior of the  $\alpha(E)$  function can be obtained by combining interferometric measurements based on the analysis of spectral oscillations present in the high transparency region of the UV-Vis-NIR spectra with absorbance data contained in the area of fundamental absorption edge in the same spectra. This approach is quite general since spectral oscillations are of low intensity but can be always detected in the UV-Vis spectra of ultrathin films. Owing to the convenient refractive index value and high surface flatness/smoothness, ultrathin Mylar (PET) films have been selected as model polymeric material for this type of analysis and the changes in the optical bandgap and Urbach energy with stretching operation have been investigated showing a scarce dependence of these parameters on mechanical damage of films.

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