

Barbara Florkowska, Wiesław Syrek, Romuald Włodek

EFFECTS OF VOLTAGE AGEING OF POLYESTER FOIL  
DETECTED BY TSDC METHOD

1. Introduction

The application of polyethylene-terephthalate foil in insulating systems results in changes of partial discharge characteristics because of the charge trapping in solid dielectric [1,2]. Such a result is important for the test-and measurement procedure of insulating system, the results of which can be missinterpreted. Particularly the p.d. inception voltage and  $\text{tg } \delta$  vs. test voltage are influenced by the formation of space charge in solid dielectric [3].

In the research presented in this paper the time characteristics of overvoltage acting were investigated using the method of thermally stimulated depolarisation current (TSDC).

2. Program and results

As the investigated material the polyethylene-terephthalate foil, Estrofol, 10  $\mu\text{m}$  thickness have been used [4]. The scope of the research program is as follows:

- the acting of special determined periodically repeated overvoltages in gas space contacting to the tested foil; their values and time of activity were the parameters of this program [4],

- the TSDC-measurements to detect the electrical effects of overvoltages.

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Dr. Barbara Florkowska, Eng. Wiesław Syrek, Dr. hab. Romuald Włodek, -  
Academy of Mining and Metallurgy, Electrical Power Institute, Kraków,  
Poland

The parameters of overvoltages were as follows: overvoltage duration - 1 s, normal voltage operation - 10 s; overvoltage value was taken as triple of the initial p.d. inception voltage  $U_0$  (at  $t = 0$ ) while the normal operating voltage was taken lower than  $U_0$ .

The TSDC-measurements were carried out in the open circuit, 0,3 mm air space, with the rate of temperature increase of 8 deg per minute, in the temperature range from the ambient temperature up to 410 K [5,8].

The typical form of the TSDC thermograms on the material before the overvoltage acting is presented in the Fig 1. The depolarisation peaks being the consequence of the natural polarisation after the technological process are seen.

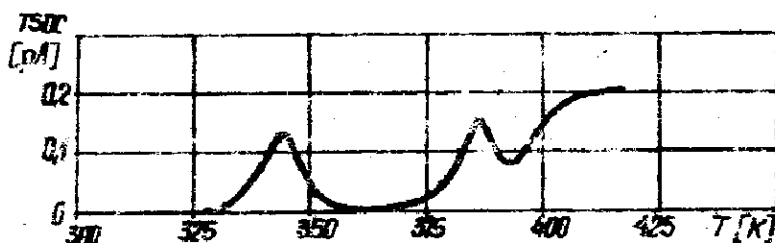


Fig.1. TSDC thermogram of PET foil in natural stage

The typical TSDC-thermograms measured after different time of voltage duration are presented in the Figures 2, 3 and 4.

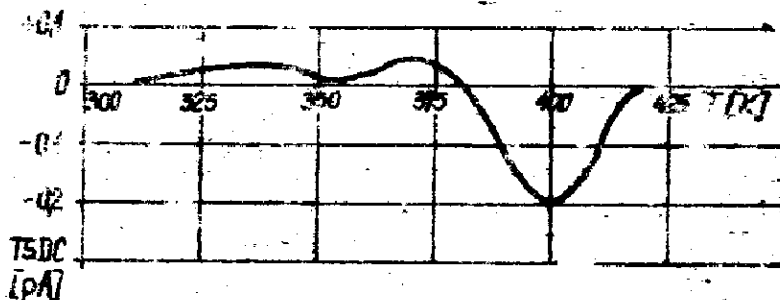


Fig.2. TSDC thermogram after 10 min. of voltage ageing

The measured results are the basis for calculation of the depolarisation charge values (DC). If we consider only this part of the thermogram which depends on the influence of p.d. acting i.e. in the temperature range from about 350 up to 400 K, the calculated depolarisation charges measured after different ageing duration can be presented in the form of

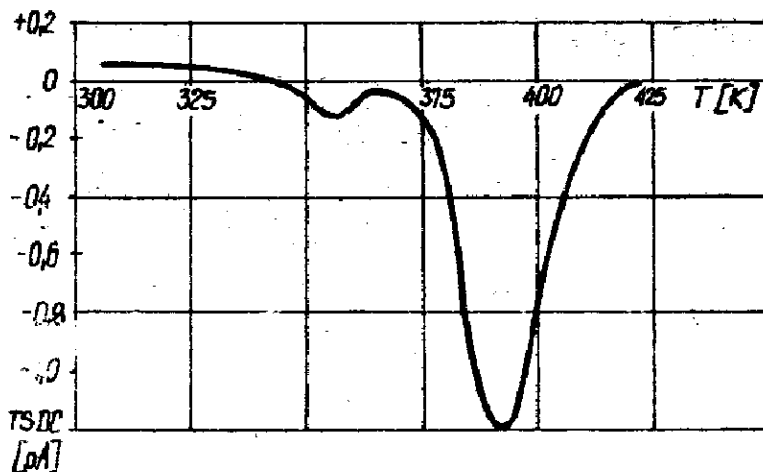


Fig. 3. TSDC thermogram after 2,5 h of voltage ageing

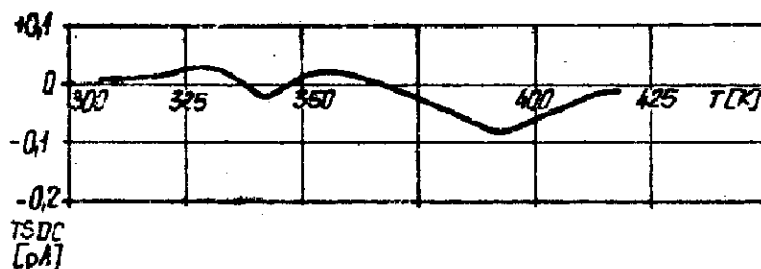


Fig. 4. TSDC thermogram after 28 h of voltage ageing

the relation of charge value vs. time. The typical form of such a time relation is given in the Fig. 5.

The initial increase of the depolarisation charge value followed by its further decrease can be observed as the typical relation obtained during the p.d. ageing in overvoltage cycles on PST foil.

### 3. Interpretation and conclusions

The p.d. acting in the gaseous phase can be treated as the source of

charges which are injected into the solid dielectric. The charges are trapped within it. The depths of traps in polymeric materials differ in relatively wide range 0,1 - 0,5 eV and more, even up to some eV [6,7] depending on the material and its crystal and amorphous structure.

Such a time relationship of the measured depolarisation charge is the result of space charge in solid dielectric. Assuming that the charge transport into the dielectric is limited by this space charge, the charged layer of the depth  $r$  is formed within it.

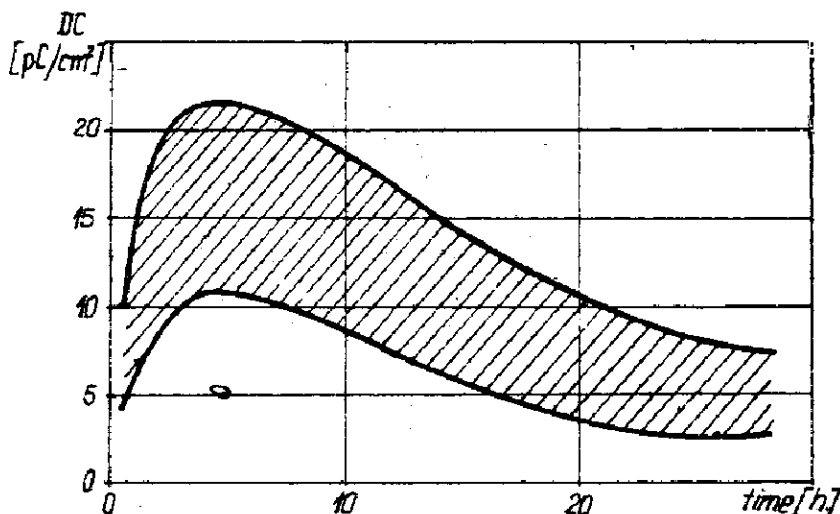


Fig.5. Typical change of depolarisation charge density

The  $r$  value depends on the duration of charge injection time, i.e. on the effective time of partial discharge action:  $r = \hat{r}(t)$ . The depolarisation charge  $DC$  which is detected and measured in the TSDC is thus influenced by the depth of the charged layer related to the thickness of dielectric. Neglecting the conductive and diffusion components of the current across the dielectric the following formula for the relative  $DC$  value can be taken [6]:

$$\frac{DC}{Q_0} = - \frac{r}{2d} \left\{ 1 - \frac{d^2}{r^2} \cdot \exp \left[ 2 \left( 1 - \frac{d}{r} \right) \right] \right\} \quad (1)$$

where:  $r$  - thickness of the charged layer,

$d$  - dielectric thickness,

$$Q_0 = \rho_0 \cdot r$$

$\rho_0$  - charge density in the  $r$ -layer.

The theoretical relation  $\frac{DC}{Q_0} = f(r)$  reaches its maximum at  $r \approx 0,6$  . d.

The real conditions existing in the dielectric as intrinsic conductivity, diffusion current component and AC field influence the resulting time relation  $r = f(t)$ . Thus the formula for the effective  $\frac{DC}{Q_0}$  value is more complicated than (1) and is now the object of investigations but its experimentally obtained results can be interpreted qualitatively as above. It should be noted that the relationship  $\frac{DC}{Q_0} = f(t)$ , where  $t$  is the ageing time, has the similar form as (1).

The described characteristics of TSDC during the process of overvoltage operation present the electrical effects of p.d. acting in polyethylene terephthalate foil. They can be used to explain the observed change of p.d. characteristics [4] which are thus treated as the result of charge trapping in the dielectric.

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