Heat affected zone in unburned, antimony trioxide containing plasticised PVC

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Abstract
What are the structural changes in the PVC layer located directly under the flame zone which is exposed to high temperatures without burning? We will try to explain in this article what actually happens in this layer, which we called heat affected zone. The layers were tested both with and without antimony trioxide (ATO or Sb2O3) flame retardant additive. Results obtained from Scanning electron microscopy (SEM) for the heat affected zone of the samples after Limited oxygen index (L.O.I.) test and compared with the samples before this test showed that there was a significant decrease in the chlorine content in the PVC structure accompanied by a significant increase in the carbon content. We suppose that this early dehydrochlorination improves the efficiency of Sb2O3.

Keywords: Plasticised PVC, HAZ, L.O.I., SEM, EDS

1. Introduction
The process of flame retardancy doesn't relate to the material's melting temperature itself, but depending only on the metaphase transformations that grow inside the material, as well as depends on compounds consisting after exposure retardant material to high temperatures. For instance, ceramic materials make a full thermal insulation due to the poor low-lying conductive materials, but at high flame is a good heat insulators in addition to immunity to flame spread [1]. Antimony trioxide is not considered a flame retardant when it is added to low-flame-resistant materials, but it is used to increase the flame retardant effect of halogen flame retardants if added to different proportions and as needed to improve its flame resistance performance [2-7]. When the polymer contains halogen flame retardants with antimony trioxide is exposed to flame, the antimony trioxide starts by interacting with the chlorine from the polymer, and chemical compounds such as SbCl, SbOCl will be formed which neutralize the free radicals, kill the flame and prevent its spread. This reaction mechanism is described in the equations as follows [8-12]:

\[ RCH_2 - CH_2Cl \rightarrow R - CH = CH_2 + HCl \]  \hspace{1cm} (1)
\[ 2HCl + Sb_2O_3 \rightarrow 2SbOCl + H_2O \]  \hspace{1cm} (2)
\[ 5SbOCl(s) \rightarrow Sb_2O_3Cl_2(s) + SbCl_3(g) \]  \hspace{1cm} (3)
\[ 4Sb_2O_3Cl_2(s) \rightarrow 5Sb_2O_3Cl(s) + SbCl_4(g) \]  \hspace{1cm} (4)
\[ 3Sb_2O_4Cl(s) \rightarrow 4Sb_2O_3(s) + SbCl_4(g) \]  \hspace{1cm} (5)

Eq. (1) was extended for the better representation of the reaction. However, when the antimony trioxide is added to PVC it becomes very effective to flame retarding, because it acts as a synergist for the halogen contained inherently in the PVC (56.7%). This situation is similar to that obtained from the stimulation of halogen flame retardants and the same compounds will be formed when ATO reacts with chlorine released from the PVC after it is burned [13-16].

2. Materials and methods
a. Materials: PVC S-5070 (OngroviII) from BorsodChem Zrt., Hungary; Dioctyl Phthalate (DOP) as plasticizer; Newstab 50 as stabilizer; Wax-E as lubricant; and antimony trioxide (ATO) with chemical formula Sb2O3.

b. Sample preparation: four batches were prepared with a laboratory extruder (Göttfert Extrusiometer G20) at 170 ºC. The barrel diameter is 20 mm, the length 20D, i.e. 400 mm. A screw of comparison rate was used with long comparison zone (PVC screw). The die was 10×4 mm flat die suitable for extruding rods for L. O. I. test without further forming.

c. Thermal tests: Limiting oxygen index test (L. O. I.) was done by the instrument Stanton Redcroft FTA flammability unit with oxygen and nitrogen cylinders gas and accurate pressure control system found in BorsodChem Zrt. The results obtained from this test shown in Fig. 1. As expected, we can observed from this figure that the limiting oxygen index of PVC increased with ATO additions, This increase is direct, as the percentage of ATO increased the value of L. O. I. will increase. Also, the percentage of oxygen which the PVC samples needs to burn will increase. This behavior due to the ATO absorbed heat and there will be phase transformations in its internal structure. This represent endothermic process which decreased surface temperature which will prevent fire propagation [4,13].
d. Scanning electron microscopy FESEM: FESEM was used for the structural analysis of plasticised PVC. This test was done by using Carl Zeiss EVO MA10 SEM.

e. Thermal gradient: thermal imaging camera (FLIR Systems) was used in order to determine the thermal gradient in L. O. I. test samples as shown in Fig. 2. The Emissivity was 0.5, reflected apparent temperature 20°C, and Object distance 1.0 m

3. Results and discussion

The images obtained from SEM show that the chlorine percentage decreased associated with an increase in carbon percentage after L. O. I. test, when comparing with PVC SEM images before the test, as shown in Fig. 3 and Fig. 4. This state as a result of PVC dehydrochlorination, where it begins to release chlorine in form of HCl from its internal structure at high temperatures, i.e. begins to decompsion in these tempreature. The loss of chlorine from the internal structure of PVC involves not only burnt and directly exposed layers of fire, but also extends to the under layers far from the combustion zone, which we called heat affected zone (HAZ) as shown in figures.

In addition, we noticed that the under layers of the test samples ignite faster when cutting the burned area and continue the L. O. I. test, which means the low flammable resistance for heat affected layer because less chlorine. However, this situation soon changes and PVC resistance to ignition improves after the addition of the antimony trioxide as shown in Figs. 5-7, where the released chlorine will be reduced from the heat affected zone. It is true that the antimony trioxide alone is not a flame retardant but it improves and increases flame retardation if it is added to halogen-containing flame retardant material containing chlorine. Antimony trioxide and chlorine combine compounds that have the ability to inhibit flame and improve flame resistance. While the structure of PVC contains a chlorine, it begins to release as a result of combustion and thermal decomposition. This chlorine reacts with the antimony trioxide forming SbCl$_3$ compound [17], which help to reduce PVC combustion and increase heat resistance of the heat affected zone. After the addition of antimony trioxide, the heat affected layer ignites relatively slowly when the burnt area is cut and the L. O. I. test continues, which means high flame resistance of this layer. The efficient flame retardant is the SbCl$_3$, is already prepared in the heat affected zone, therefore the extinguishing of the falme starts sooner.
Fig. 4. SEM - energy dispersive X-ray microanalysis for HAZ in burned plasticised PVC L. O. I. sample

Fig. 5. SEM - energy dispersive X-ray microanalysis for HAZ in burned plasticised PVC + 1% Sb$_2$O$_3$ LOI sample

Fig. 6. SEM - energy dispersive X-ray microanalysis for HAZ in burned plasticised PVC + 3% Sb$_2$O$_3$ LOI sample

Fig. 7. SEM - energy dispersive X-ray microanalysis for HAZ in burned plasticised PVC + 5% Sb$_2$O$_3$ LOI sample
4. Conclusions

The PVC combustion is not confined to the flame exposed layer but extends beyond the burning layer. Experiments have shown a decrease in chlorine content in the non burning heat affected zone. Without Sb₂O₃ ease of ignition of heat affected zone was observed due to structural changes. The heat affected zone has a low inclination of ignition and is immediately flame retardant if the compound contains Sb₂O₃ because the SbCl₃ is already present in this layer.

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