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Particle Flux Effects on Physicochemical Polymer Degradations

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Particle flux effects on polymer physicochemical degradations were investigated. Different polymers were studied depending on application: silicone, fluoropolymer, acrylic, epoxy, polyimide, polyurethane, etc. Polymers with surface functional properties (paint, coating, etc.) were dissociated from polymers with bulk properties (potting, cable, etc.). Surface materials were exposed to protons, and bulk materials were exposed to electron radiations. Two fluxes were used: $5 \times 10^{11}$ and $7.4 \times 10^{12}$ cm$^{-2}$ s$^{-1}$ for electrons and $8.7 \times 10^{10}$ and $2.2 \times 10^{11}$ cm$^{-2}$ s$^{-1}$ for protons. The total dose was fixed at 500 Mrad for electron radiations and 1 Grad for proton radiations. Functional properties (thermo-optical and mechanical) depending on material application were measured before and after irradiation exposure. Chemical structure modifications were also investigated. From all combined results, no global tendency was evidenced. The study shows that low or high irradiation flux induces more or less degradations according to polymer type, the composition of the polymer, and functional properties of concern. Flux effects can also depend on particle nature (electron or proton). The last point was highlighted by silicone polymer degradations; high proton flux caused more damage than low flux. On the contrary, low electron flux induced more degradations than high flux.

Nomenclature

\[ T_g = \text{glass-transition temperature, °C} \]
\[ \alpha_s = \text{solar absorbance} \]
\[ \varepsilon_{irr} = \text{infrared emissivity} \]

I. Introduction

In space applications, materials are exposed to specific environments: charged particles (from the radiation belts), the solar spectrum, atomic oxygen (in low orbit, <1000 km), and micrometeorites and debris. Vacuum conditions and temperature are added to this.

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Among these different environmental parameters, the charged particles are constituted by electrons or protons trapped in the radiation belts, rash protons (also x rays on the sunny side) and solar wind. Materials on external parts of the satellite are directly exposed to this harsh environment. Charged particles mainly generate an ionizing absorbed dose profile into these materials. Because of their energy spectrum and the mode of interaction, the protons are responsible for the surface dose (first micrometers) and electrons for the bulk dose.

The interaction of particle radiation with an organic material causes the phenomena of chain scission and/or crosslinking via ionization mechanisms. The effects are linked to the accumulation over time of the ionizing dose (cumulative effect) causing surface and/or volume changes in the material. The induced macroscopic effects consist mainly of color change (absorbance increase and transmittance decrease) and the decrease of mechanical properties performance (mass loss, brittleness, reduced stress and strain at the break, etc.).

A classification of polymer materials by particle sensitivity and degradation type is given in the literature survey depending on polymer structures (e.g., the presence of the aromatic group leads to better stability) and the chemical composition of its monomer [1]. Additives can also have a high influence on degradation. An index radiation indicator is given by Tavler et al. [European Laboratory for Particle Physics (CERN)] for prioritizing materials per family according to their mechanical behavior (ultimate strength and deformation) [2]. However, these classifications are based on data obtained under specific conditions, generally not representative of the space environment (data from nuclear industry or from neutrons and a gamma accelerator, with exposure to air). They only could be used as indicative.

Therefore, evaluation of material degradation during space missions is estimated by performing ground tests that closer simulate the space environment in order to select and qualify the appropriate materials [3]. For this, test facilities are used to replicate specific degrading components of the space environment. For many reasons, particularly to reduce the aging time and due to specifications of the particle accelerator, these tests are not performed under the exact same conditions as the environment the material will see over the
lifetime of the mission. Accelerated tests, especially important for particle irradiation, are performed, according to specific standards [4–6]. The possible acceleration impact on the material degradation is currently not evaluated.

Dose rate effects have been observed in the literature, with potentially more degradation observed for low dose rates [7–9]. These dose rate effects may be due to the oxygen environment and the oxidation of radiolysis species created during irradiation. Again, many data come from nuclear applications (neutrons/in air) or CERN (γ and neutrons/in air). In this case, because of the oxygen involvement, surface effects are dominant, but low particle flux tests (for long durations) on thin layers can be critical to bulk properties. We can minimize the effect of oxygen by working under a stream of neutral gas (nitrogen and argon) or under vacuum, provided that there is no adsorbed oxygen. The effect of dose rates can be seen in vacuum particle irradiation. Briskman et al. report data on Polytetrafluoroethylene (PTFE) and low-density polyethylene [10], including the gas emission rate measurement, for which significant differences were observed for different dose rates. An increase of degradations of tensile strength and elongation at rupture was observed for PTFE at low dose rates. In contrast, Hardgrove presents other electrons (e−) + protons (p+) dose data at different acceleration rates that show for white paints that the worst case is observed at high acceleration rates [11]. A comparison with in-flight data leads him to the conclusion that the dose rate closest to space arrival provides the most realistic degradation value.

Considering these different results, the study presented in this paper aims to evaluate the particles flux effects on the physico-chemical properties of different polymers. Indeed, as previously indicated, the irradiation aging tests performed on the ground in order to simulate the materials aging in orbit cannot be fully representative of the in-flight environment. In particular, the particle flux on the ground is highly accelerated compared to that in the space environment. In the first part of this paper, the experimental process is described with a presentation of the materials tested, the different irradiation conditions, and the associated analysis methods.

In the second part, a global synthesis on the flux effects observed on the different polymers tested is presented.

Finally, in the last part, a focus on the flux effects on silicone rubber is discussed.

II. Materials and Experimental Methods

A. Materials

Various polymers used on satellites were studied including surface materials, such as paints and coatings, and bulk materials, such as cables, fixing, and structural bonding.

The following surface materials that were of concern that were tested include room temperature vulcanized (RTV) silicone, polyurethane (PU), polyetherimide (PEI), and polyester film. The investigated bulk materials were acrylic adhesive polyimide tape, RTV silicone, epoxy, fluorinated ethylene propylene (FEP), and acetal.

Two types of irradiations tests were performed depending on the function of the materials studied; surface materials were exposed to proton flux, and bulk materials were exposed to electron flux.

B. Irradiations

1. Proton Flux Irradiation

Proton irradiation was performed in the SEMIRAMIS facility (ONERA), using a 2.5 MeV Van de Graaff accelerator. The proton energy was 240 keV, and the total dose was 1 x 107 Gy (penetration at about 3 μm). Two different fluxes were selected: 8.7 x 109 p+ · cm−2 · s−1 and 2.2 x 1010 p+ · cm−2 · s−1. A vacuum lower than 10 μmbar was obtained after a 1 day pump down period. The sample holder temperature was maintained at 25°C for the duration of the test.

2. Electron Flux Irradiation

The electron irradiation was performed by TRAD using a Van de Graaff accelerator able to produce high-energy electrons (1–3 MeV). A fixed electron current is delivered by the horn. The samples are horizontally disposed on the table. The dose rate is directly correlated to the average amount of electrons received by the samples under the beam.

The irradiation table is cooled down to 20°C. A thermocouple is put on the table next to the samples, to check with a remote controller the eventual temperature increase during the experiment.

A display screen in the control room provides a direct view of the table and the samples.

The irradiation is performed under a nitrogen environment, thanks to a flowing nitrogen setup containing all the samples that are irradiated.

Before every irradiation, a preliminary dosimetry step is performed in order to determine the dose rate.

The dosimetry is performed using thin film radiographic dosimeters at a given beam current and by measuring the deposited dose. The dose rate is calculated as the needed beam current to perform the irradiation at the target dose rate.

In the frame of this study, irradiations are performed at 3 MeV and at standard and high dose rates, respectively, 5 x 1011 and 7.4 x 1012 e− · cm−2 · s−1.

C. Analysis Techniques

1. Ultraviolet Visible Near-Infrared Spectroscopy

Different Ultraviolet-Visible-Near-Infrared (UV-Vis-NIR) analyses were performed at ONERA:

1) For measurements in air, the UV VIS NIR measurement system was composed of a Perkin Elmer Lambda 1050 spectrophotometer associated with a 150 mm diameter integrating sphere in air. The calibration was performed using a diffuser Spectralon SRS-99-010 standard. The diffuse transmission spectra in the range 200–2500 nm were acquired at intervals of 1 nm, with the lamp change at 319.2 nm.

2) For measurements under vacuum, the SEMIRAMIS facility is equipped with a Perkin Elmer Lambda 1050 spectrophotometer connected to a mobile integrating sphere under vacuum. Reflectance spectra in the range 250–2500 nm can be obtained, without breaking vacuum, before and after irradiation.

2. X-Ray Photoelectron Spectroscopy

Atomic concentration and chemical bonding information were obtained from the specimen surfaces by x-ray photoelectron spectroscopy (XPS). XPS measurements were realized with a Thermo Scientific K alpha spectrometer with a monochromated Al Kα radiation (hv = 1486.6 eV). Spectra were collected using an x-ray spot size of 400 μm.

3. Attenuated Total Reflection Fourier Transform Infrared Spectroscopy

Attenuated total reflection (ATR) Fourier transform infrared (FTIR) spectroscopy was performed at ONERA to characterize surface chemical degradations. The ATR FTIR spectra were recorded using a Nicolet 800 FT-IR spectrometer equipped with a deuterated triglyceride sulfate detector in the range 400 – 4000 cm−1, with a resolution of 4 cm−1. A diamond ATR accessory was used to investigate the first micrometers of the samples.

4. Nuclear Magnetic Resonance Spectroscopy

28Si solid-state nuclear magnetic resonance spectra were recorded on a Bruker Avance 400 spectrometer equipped with a 4 mm probe operating at 9.39 MHz. Samples were spun at 12 kHz at the magic angle using ZrO2 rotors. 29Si Magic Angle Spinning (MAS) spectra were obtained under high-power proton decoupling conditions, with a small flip angle (30 deg) and a recycle delay of 60 s.

5. Differential Scanning Calorimetry

At Thales Alenia Space, Differential Scanning Calorimetry (DSC) measurements were carried out using a NETZSCH DSC 204 apparatus for RTV silicone and a DSC823e from Mettler Toledo for other polymers. The ramp was fixed at 10°C/min. The analysis temperature range was fixed depending on polymer characteristics.
6. Peel Test
Peeling tests were performed in accordance with ASTM D1000 standard at Airbus Industries. Peelings were performed with a 180 deg angle at 400 mm/min and at ambient temperature on a Zwick Roell facility.

7. Adherence Test
Adherence tests performed on coatings according to ISO EN 2409. P224 tape was used at Airbus Industries.

8. Shear Test
Shear tests were performed according ISO specification (ISO.4587:2003). Tests were performed at room temperature on a Zwick Roell facility at Airbus Industries. Shear Tests were performed on adhesives (epoxy and silicone) at Thales Alenia Space. Tests were performed at room temperature on a Zwick 100 kN apparatus.

9. Nanoindentation
At Centre National d’Études Spatiales (CNES), nanoindentation was carried out with a nanoindenter AX (MTS), using a Berkovich point and a depth penetration of 5 μm. Twelve indentation points were performed on each sample. A sample Poisson ratio was fixed at 0.4.

10. Dynamic Mechanical Analysis
Three-point flexure tests were performed on a Dynamic Mechanical Analysis (DMA) 242 NETZSCH at CNES. The temperature range of measurement was fixed according to polymer characteristics. Measurement was performed with a ramp of 5°C/min and a frequency of 1 Hz. The distortion amplitude was 30 μm, dynamic strength limit was fixed at 7.2 N, and the constant part of static strength was fixed at 0 N. Each sample was tested two or three times.

III. Flux Effects on Polymers: General Results
A. Surface Irradiation
Table 1 summarizes all the results obtained on surface materials during the aging tests.
No flux impact was evidenced except for RTV silicone. RTV silicone showed higher degradation at high proton flux.

B. Bulk Irradiation
Table 2 summarizes all the results obtained on bulk materials during the aging tests.
At this dose level and with the considered fluxes, no flux impact was evidenced except for RTV silicone, the second epoxy, and FEP. It has to be noted that in the same material family the degradation could be different (epoxy adhesive).

Table 1  Flux effects on surface polymers

<table>
<thead>
<tr>
<th>Materials</th>
<th>1 Grad (p⁺)</th>
<th>1 Grad (p⁺)</th>
</tr>
</thead>
<tbody>
<tr>
<td>RTV silicone</td>
<td>Increase of ( \alpha_p ), decrease of stress at break and modification of the polymer structure</td>
<td></td>
</tr>
<tr>
<td>PEI</td>
<td>1) No significant degradation (variation of ( \alpha_s, \epsilon_{irr} ), and adherence test lower than the uncertainty of the measurement)</td>
<td>2) No flux significant impact</td>
</tr>
</tbody>
</table>

Table 2  Flux effects on bulk polymers

<table>
<thead>
<tr>
<th>Materials</th>
<th>500 Mrad (e⁻)</th>
<th>500 Mrad (e⁻)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Acrylic tape</td>
<td>1) High adherence degradation due to outgassing under irradiation for high and low flux; bubble creation (adherence test: ~95% after irradiation)</td>
<td>2) No significant flux effect at this dose level</td>
</tr>
<tr>
<td>2nd Epoxy adhesive</td>
<td>1) Degradation of mechanical properties</td>
<td>2) Degradation of mechanical properties (stress at break: ~15% after irradiation) even if physicochemical property changes observed [( T_g ) decrease (~10°C after irradiation) and observation of a new pic in ATR]</td>
</tr>
<tr>
<td>FEP</td>
<td>1) No significant flux impact</td>
<td>2) No significant flux impact</td>
</tr>
<tr>
<td>Delrin®</td>
<td>1) Very strong degradation of the material at a dose level of 500 Mrad</td>
<td>2) No significant impact of the irradiation flux</td>
</tr>
</tbody>
</table>

Fig. 1 Optical micrograph of the high-flux proton irradiated sample (cracks on the left side).

Fig. 2 Reflectance from 250 to 800 nm of pristine and protons irradiated silicone rubber.
IV. Flux Effects on Silicone Rubbers

To illustrate the process used and the results obtained in this study, particular attention is paid to the material considered as a surface and bulk sample during this testing: RTV silicone rubbers.

A. Surface Irradiation

A slight yellowing of samples was observed after proton irradiations. Moreover, inhomogeneous degradation on the high flux sample was noted, as shown on the optical micrograph (Fig. 1). An important surface damage was observed with the formation of many cracks localized in one-half of the sample. This phenomenon was previously observed on silicone under proton irradiation [12,13].

1. Thermo-Optical Properties

The SEMIRAMIS facility enables measurement of thermo-optical properties in situ (under vacuum). Thermo-optical properties were measured by reflectance under air and vacuum conditions. Results are presented for the two different irradiation fluxes in Fig. 2. A UV cutoff shift to higher wavelengths was observed under vacuum after irradiation, showing absorbent species formation. This shift is more important after the high flux irradiation. After the replacement of the samples under ambient air, new measurements showed that there were recoveries of their reflectance spectra. This recovery had been previously observed on different materials [14,15]. This phenomenon is particularly significant for the high flux irradiated sample. The low and high flux final spectra (measurement in air) were closed.

Thermo-optical coefficients (IR emissivity and solar absorptances) have been calculated. IR emissivity changes were not significant. Solar absorptances are reported in Table 3. Recovery phenomenon was also observed in the change in $\alpha_s$, particularly for the high-flux irradiated sample. Accordingly, the $\alpha_s$ end of test values (measurement in the air environment) for both aged samples were similar, even though the interim $\alpha_s$ (measurement under vacuum just after irradiation) was higher when proton flux was higher.

2. X-Ray Photoelectron Spectroscopy

XPS measurements show that the atomic percentages obtained for the different samples are similar (C, O, and Si). However, a shoulder corresponding to the SiO$_x$ bond is observed for irradiated samples. This is more evident for the high-flux sample (Fig. 3). This result indicates an evolution of the surface nature of the silicone rubbers into silica, which is a function of the total dose and the irradiation flux.

3. Attenuation Total Reflection Fourier Transform Infrared Spectroscopy

ATR FTIR spectra obtained before and after proton irradiation are presented in Fig. 4. A distinction can be seen on the ATR FTIR measurements between the high-flux cracked sample and the other irradiated samples (low- and high-flux noncracked):

1) For the high-flux cracked sample, ATR FTIR measurement confirms silica formation on the high-flux irradiated sample, with an increase of the SiOH bands absorption (3100 – 3600 cm$^{-1}$) [16,17]. The appearance of a new absorption band (1700 cm$^{-1}$), attributed to the CO groups, responsible for the sample yellowing, is also observed.

2) For the other irradiated samples, spectra obtained after irradiation are closer to that of the pristine sample. However, an increase of the SiH and a decrease of the SiCH3 and SiPhenyl groups are observed. A SiOSi characteristic peak (1070 cm$^{-1}$) decreases after irradiation, for both high-flux irradiated samples, while it is unchanged for the low-flux irradiated sample.

The formation of a silica-type layer localized on the surface can explain why cracks appear: the density of silica is higher than that of silicone rubber.

4. $^{29}$Si Solid-State Nuclear Magnetic Resonance Spectroscopy

$^{29}$Si solid-state nuclear magnetic resonance (NMR) spectra of all silicone rubber samples are presented in Fig. 5 ($e^-$ and $p^+$ irradiation). Pristine and irradiated spectra show a high signal at $-22$ ppm attributed to $D$ siloxane units within the chain. Considering proton irradiation, a slight peak corresponding to SiCH$_2$Si bonds appears.

<table>
<thead>
<tr>
<th>Materials</th>
<th>In air</th>
<th>Under vacuum</th>
<th>Under vacuum</th>
<th>In air</th>
</tr>
</thead>
<tbody>
<tr>
<td>Low flux</td>
<td>0.19</td>
<td>0.18</td>
<td>0.20</td>
<td>0.21</td>
</tr>
<tr>
<td>High flux</td>
<td>0.17</td>
<td>0.17</td>
<td>0.27</td>
<td>0.21</td>
</tr>
</tbody>
</table>

Table 3 Evolution of solar absorptance $\alpha_s$ at each step of proton irradiation

![Fig. 4 ATR FTIR silicone rubber before and after proton irradiation.](image-url)

![Fig. 3 XPS spectra of silicone rubber before and after proton irradiation.](image-url)
after low-flux irradiation (~37 ppm). This phenomenon indicates new crosslinks. An increase of the SiO$_4$ broad signal (~110 ppm) is observed after the high-flux irradiation, particularly into the cracked zone. This indicates the creation of silica during proton high-flux irradiation.

### B. Bulk Irradiation

To assess the bulk properties of the material, different characterization techniques were used such as UV VIS NIR spectroscopy, NMR, DMA, nanoindentation, and DSC to determine the evolution of the physicochemical properties of the material submitted to the electron beam.

The ATR FTIR spectroscopy shows a significant degradation of the samples submitted to a standard dose rate, leading to the apparition of SiOH, CO, and SiH bonds. For high dose rate, the sample modification consists mainly in the creation of SiH bonds and on the decrease of the number of SiPhenyl bonds (Fig. 5).

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**Fig. 5** $^{29}$Si NMR spectra of silicone rubber before and after electron and proton irradiation.

**Fig. 6** ATR FTIR silicone rubber before and after electron irradiation.

**Fig. 7** DMA spectra of silicone rubber before and after electron irradiations.
This tendency is confirmed by the NMR analysis, which reveals the formation of crossing points (SiO₂ bonds) coupled with the diminution of the SiPhenyl bonds (Fig. 5). After irradiation at the standard dose rate only, another type of crossing point is also observed (SiCH₂Si). Complementary NMR analysis shows a stiffening of the silicone skeleton after electron irradiation.

The DMA tests focused on the mechanical behavior of the material depending on the temperature (Fig. 7). Results show an evolution of the temperature of the inflexion point for the storage modulus, with a decrease of about 10°C after electron irradiation. Additionally, no significant difference was noticed between the values after standard and high dose rate irradiation.

DSC analysis was also performed. The results reveal that the material was not fully polymerized, leading to the apparition of an exothermic peak during the first scan. These peaks were localized at the standard dose rate and the one submitted to a high dose rate. Thus, there was an evolution of this temperature after irradiation, but no effect of the dose rate was noticed.

Finally, mechanical properties of the material were assessed through the irradiation of shear test samples. A clear decrease of the tensile strength of around 40% was measured, from 4.1 to 2.3 MPa. The samples irradiated at the standard dose rate seemed slightly more degraded.

V. Conclusions

Different materials families were tested in the frame of this study. They were selected to represent some of the main organic material families used on spacecraft such as epoxies, silicones, polyurethanes, fluoropolymers, etc.

The aim of this study was to determine the influence of the irradiation flux on material degradation. It has to be noticed that, due to facility limitations, the flux as applied in the frame of this study, even the lowest ones, was still largely above real fluxes in orbit.

During this study, various analysis tests were performed on the materials in order to be able to determine the evolution of the functional properties of the tested materials but also to determine some physicochemical changes.

The general following observations can be reported:
1) There is no evidence of a similar effect of the irradiation flux on the different materials families. Indeed, for some materials families, the degradation is higher with a higher irradiation flux, while it could be the contrary for another material family.
2) The effects can also be different for two materials in the same family (different behavior observed on two epoxies).
3) The irradiation flux effect can be different on the same material depending on the type of particle irradiation (electrons or protons).

This study demonstrated that ground-testing parameters are important to the evaluation of material behavior against radiation. The current study focused on the irradiation flux effect. The investigation of other irradiation parameters such as the type of particles, the atmosphere, and the irradiation temperature is also relevant and will be investigated in future work.

References


T. K. Minton
Associate Editor