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PSS: Photolysis of cometary organic dust analogs on the Expose R2 mission at the International Space Station

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ABSTRACT

We describe the results obtained on a set of organic samples that have been part of the experiment “Photochemistry on the Space Station (PSS)” on the EXPOSE-R2 mission conducted on the EXPOSE-R facility situated outside the International Space Station (ISS). The organic samples were prepared in the Catania laboratory by 200 keV He⁺ irradiation of N₂:CH₄:CO icy mixtures deposited at 17 K, on VUV transparent MgF₂ windows. This organic material contains different chemical groups, including triple CN bonds, that are thought to be of interest for astrobiology. It is widely accepted that materials similar to that produced in the laboratory by ion irradiation of frozen ices could be present in some astrophysical environments such as comets. Once expelled from comets, these materials are exposed to solar radiation during their interplanetary journey. In the young Solar System, some of these processed materials could have reached the early Earth and contributed to its chemical and pre-biotic evolution. The samples were exposed for 16 months to the unshielded solar UV photons. It was found that, if an Interplanetary Dust Particle (IDP) containing organic material (50% vol) is large enough (>20-30 μm), relevant chemical groups, such as those containing the CN triple bond, can survive for many years (>10⁴ years) in the interplanetary medium.

1. INTRODUCTION

Solar UV radiation is fundamental to organic chemistry and indeed drives most chemical evolution processes, via photochemical reactions, in the Solar System (Cottin *et al.*, 2015). In recent years, many experimental studies have been conducted with the intent to discern the effects of cosmic UV photons on organics in different astrophysical environments. These studies have used varying sources that emit photons in the range of $100\text{ nm} < \lambda < 200\text{ nm}$. It should be noted, however, that none of these sources have the capability to accurately reproduce the full solar UV spectrum (Cottin *et al.*, 2015; Cottin *et al.*, 2017). In particular, differences as large as a factor >100 in the photolysis kinetic can be found between the extrapolation of laboratory results to space conditions and measurements of organic samples exposed to low Earth orbit conditions (Guan *et al.*, 2010). Earth's atmosphere blocks all ionizing ($\lambda < 125\text{ nm}$) and most of the harmful ($\lambda < 310\text{ nm}$) UV solar photons. The International Space Station (ISS) offers the unique opportunity to expose, for relatively long periods of time, samples to the real, unfiltered solar electromagnetic spectrum. Here, we present results obtained from a set of organic samples that were part of the Photochemistry at the Space Station (PSS) experiment on the Expose-R2 mission (Cottin *et al.* This Volume, In prep). The Expose R2 mission used the ESA Expose-R facility, which is situated outside the ISS on the URIM-D platform of

the Russian module *Zvezda*. Expose-R accommodated three European astrobiological experiments (BIOMEX, BOSS, PSS) and one Russian (BIODIVERSITY) astrobiological experiment, all of which comprised several chemical and biological samples that were exposed to cosmic ionizing radiation and solar UV photons.

Our organic residues were prepared by 200 keV He⁺ irradiation of frozen N₂:CH₄:CO ice mixtures deposited, at 17 K, on MgF₂ Vacuum UV (VUV) transparent substrates. It is generally understood that materials similar to those produced by energetic processing of ices may be present in some astrophysical environments such as interstellar grains (Pendleton and Allamandola, 2002).

We chose an ice mixture that well represents ices that occur at the surface of Trans Neptunian Objects (TNOs) and comets in the Oort cloud. Indeed, the spectral signatures of solid nitrogen, methane, and carbon monoxide are seen in TNOs (e.g., Pluto; see Protopapa *et al.*, 2017). Furthermore, a number of studies have obtained evidence of dark organic materials that may have been produced by cosmic ion irradiation of TNOs surface ices (e.g., Grundy and Buie, 2002; Brunetto *et al.*, 2006). The N₂:CH₄:CO ice mixture we used in our study is also representative of comets in the Oort cloud. It has been proposed that, as a consequence of Galactic Cosmic Rays (GCR) irradiation, a refractory crust, under which water ice is the dominant species, occurs on comets in the Oort cloud (Strazzulla and Johnson, 1991). In particular, the formation of a stable

crust, due to the conversion of low molecular weight hydrocarbons into a stable organic residue, requires an irradiation dose on the order of 100 eV/16u (Strazzulla and Johnson, 1991). Owing to the solar modulation, maximum irradiation doses are reached far from the Sun in the local interstellar medium (LISM) (Cooper *et al.*, 2003). For comets in the Oort cloud, 100 eV/16u have penetrated to a depth of 1-2 meters throughout the age of the Solar System. Lower doses of 30 eV/16u penetrate to a significantly greater depth of 5-10 meters, (see figure 12 of Strazzulla and Johnson 1991). Thanks to Rosetta measurements, a complex macromolecular material has been detected in comet 67P. In particular, it was found that 50% in mass of dust particles ejected from the nucleus of the comet are composed of this organic material (Fray *et al.*, 2016; Bardyn *et al.*, 2017). When a comet from the Oort cloud or from the Kuiper belt (i.e., the TNOs region) enters the inner Solar System, the water ice-dominated sublimation at ~3 au (astronomical units) or the sublimation of other more volatile species such as CO, CO₂, N₂, CH₄, at much longer heliocentric distance (up to ~25 au; see Szabó *et al.*, 2011) ejects gas and dust. Cometary dust constitutes a fraction of those Interplanetary Dust Particles (IDPs) that have been collected in Earth's atmosphere by airborne impacting collectors flown at an altitude of 18-20 km and of those micrometeorites collected in Antarctica. The remaining fraction of IDPs and of the micrometeorites come from asteroids (Genge *et al.*, 2008). The Ultra Carbonaceous Antarctic MicroMeteorites

(UCAMMs) constitute a particular class of micrometeorites (Dobrica *et al.*, 2009,2012; Dartois *et al.*, 2013, 2018). It has been shown that UCAMMs have a carbonaceous phase with IR signatures similar to those of the organic residues described in this work (Baratta *et al.*, 2015). It is believed that these high carbon content (> 50% vol.) and high N/C ratio (e.g., 0.12 in DC65) micrometeorites originated in the outer Solar System beyond the nitrogen snow-line, for example in an Oort cloud comet by cosmic ray irradiation (Dartois *et al.*, 2013, 2018). In general, the origin of organic materials present in IDPs and micrometeorites is still under debate, though various processes may be a contributing factor, such as UV photolysis, cosmic ion irradiation, and grain-surface chemistry, all of which may act at different evolutionary phases. Comets accreted in the cold outer regions of the protoplanetary disk, and hence cometary materials may have preserved those original components which formed the Solar System (Sandford *et al.*, 2008). Hence, UV and GCR processed interstellar grains could have been incorporated into comets. It is important to note that hot-spots in some IDPs have a D/H ratio that reaches the value of interstellar molecules, which suggests that interstellar molecular-cloud organic materials from the early Solar System may have survived (Messenger, 2000). Organic matter in IDPs and micrometeorites could also have originated in the protoplanetary disk. In the cold part of the disk, in particular, ices would have covered small (from sub-micrometer to mm) refractory grains. During this early phase, the icy mantles

would have received large amounts of energy from cosmic rays, solar particles, and UV photons that could have transformed a fraction of the ice into organics (Dartois *et al.*, 2013). In particular, it has been found that icy grains that originated in the outer disk ($T < 30\text{K}$) experienced ultraviolet irradiation exposures and thermal warming similar to those that have been shown to produce complex organics in laboratory experiments (Ciesla and Sandford, 2012). Carbonaceous matter that formed in these small grains could then have been incorporated into icy proto-/cometary objects and eventually delivered to the inner Solar System by comets. Hence, IDPs and micrometeorites could contain materials similar to the PSS organic residues if emitted from the external (ion-processed) layers of a comet or as a heritage of irradiation that occurred in the protoplanetary disk phase. During their journey in the interplanetary medium, these materials would have been exposed to GCRs, solar UV photons, and the solar ion population before reaching Earth or other objects in the Solar System. Hence, materials similar to those discussed here could have reached the pre-biotic Earth. As is shown in the next sections, the organic residues we have produced contain astrobiological relevant species such as nitriles (Palumbo *et al.*, 2000a; Palumbo *et al.*, 2000b; Kaiser and Balucani, 2001). One of the key points for the synthesis of the first bio-monomers is the availability of suitable precursor organics under primitive Earth conditions. According to Ruiz-Mirazo *et al.* (2014), the role of extraterrestrial organics in providing precursor materials

was significant. The aim of the present study was to analyze the destruction/modification, induced in space by solar UV photons, of the relevant chemical species contained in the organic residue.

2. MATERIALS AND METHODS

2.1 Sample preparation and characterization

Sample preparation and characterization was described previously in the work of Baratta *et al.* (2015) to which the reader is referred for further details. Here, we offer a brief description of the procedures and experimental set up that was adopted to prepare and characterize the samples. A schematic view of the experimental apparatus can be seen in Fig. 1(a).

The experimental apparatus used to irradiate frozen ices consists of a stainless steel ultra high vacuum (UHV) chamber with a base pressure lower than 10^{-9} mbar. Gas mixtures are prepared in a separate mixing chamber that is pre-evacuated to a pressure of $\sim 10^{-7}$ mbar by using a turbo-molecular pump. Once prepared, the gas mixture is admitted through a flow regulated valve in the UHV main chamber, where it freezes on an IR transparent substrate that is placed in thermal contact with the tail section of a close-cycle CTI Helium cryostat ($T=16-300$ K). The thickness and refractive index of the deposited ices can be measured, during accretion, by looking at the

interference pattern (intensity versus time) given by a He-Ne laser beam ($\lambda=543.5$ nm) reflected at near-normal incidence ($i=2.9^\circ$) by the ice-vacuum and ice-substrate interfaces (see Urso *et al.*, 2016 and references therein).

Fast ions, up to an energy of 200 keV (400 keV for double ionization), are obtained by a 200 kV Danfysik implanter. The ion beam is electrostatically swept over the substrate to obtain a uniform coverage of the irradiated ices. The total ion fluence found in different astrophysical scenarios can be easily reproduced in the laboratory. Nevertheless, it should be noted that this occurs on very different time scales. In general, since it is impossible to reproduce, in laboratory, the dose rate found in astrophysical scenarios and obtain measurable effects in human lifetime, the results can not depend on the dose rate; any experimental result that depends on the dose rate, such as the beam-induced heating of the target, can not be extrapolated to any astrophysical scenario. The ion beam current was kept below $1 \mu\text{A}/\text{cm}^2$ to prevent a macroscopic heating of the frozen film. It should be noted that a temperature increase of 15 K would have caused the complete sublimation of CO and N₂ from the mixture. We have verified many times that the effects induced by fast ions on frozen mixtures do not depend on the dose rate for such low ion currents. In any case, the extrapolations of the laboratory results to astrophysical scenarios implicitly assume that other processes that could interfere with the cosmic ion processing (such as resurfacing and diffusion)

do not take place on the astrophysical object. The ion fluence was measured by integrating the ion current monitored during irradiation. The irradiation dose was calculated by multiplying the ion fluence (ions/cm²) and the energy loss per unit path length (stopping power; eV cm²/molecule) given by the SRIM code (Ziegler *et al.*, 2008).

A FTIR spectrometer Vertex 70 by Bruker (10000-400 cm⁻¹=1-25 μm) is interfaced to the UHV chamber through two KBr IR transparent windows. The spectrometer is placed on a moveable optical bench that allows a precise alignment of the IR beam with respect to the substrate. For this purpose, a hole is made on the substrate holder that allows the IR beam of the spectrometer, transmitted by the ice film deposited on the substrate, to reach the detector. The substrate holder is inclined by 45° with respect to both the ion beam and IR beam directions, and hence IR transmission spectra of the irradiated ices can be acquired in situ before, during, and after irradiation.

To prepare the organic samples for the PSS experiment, we used a C, N, and O-bearing ice mixture: N₂:CH₄:CO (1:1:1). In particular, we used Aldrich chemical CH₄, CO, and Air Liquide N₂. Three fill-pump down sequences were performed before introducing the mixture into the UHV chamber. This was done to prevent contamination of the mixture by the residual gas (mostly water vapor) present in the mixing chamber. Once deposited on a VUV

transparent MgF₂ substrate (furnished by ESA) at 17 K, the ice mixtures were irradiated by 200 keV He⁺ ions at a dose of 110 eV/16u. Under UHV condition, almost all the residual gas is made of molecular hydrogen that does not freeze on the substrate at 17 K. Hence, contamination, if any, could only come from the mixing chamber. Nevertheless, we were not able to recognize in the infrared spectrum of the deposited ices any significant amount of possible contaminants such as H₂O or CO₂. We prepared a total number of 30 samples over three different thicknesses (10 samples for each thickness). The average values of the three thicknesses of the ice mixtures were 1.322 ± 0.02 , 0.889 ± 0.01 , and 0.456 ± 0.01 μm , which correspond, under our experimental conditions, to interference curves of about 6, 4, and 2 fringes, respectively. The given uncertainties are the root mean squares on the corresponding 10 samples; their low values testify to the good reproducibility of the deposited ice mixtures thickness. At the end of deposition, an FTIR spectrum of the ice mixture was acquired. Unless indicated otherwise, all the transmittance spectra reported in this article are given in optical depth scale. The optical depth (τ) was calculated by using the Beer-Lambert law: $I=I_0e^{-\tau}$ where I_0 is the normalization continuum. An example of the IR spectra of the deposited ice mixtures for the three different thicknesses can be found in the work of Baratta *et al.* (2015) (see their Fig. 2). The presence of the CO (2140 cm^{-1}) and CH₄ (3010 and 1300 cm^{-1}) features is evident in the spectra. Molecular

nitrogen is a homo-nuclear symmetric species, and hence it has no IR active vibration. For this reason, no feature due to molecular nitrogen could be seen in the deposited ice film. The column density n (molecules/cm²) of the observed species can be calculated by using the relation:

$$n = \left(\int_{\nu} \tau(\nu) d\nu \right) / A \quad (1)$$

where A represents the band strength (cm/molecule) and $\int_{\nu} \tau(\nu) d\nu$ is the band area (cm⁻¹) of a given band. The column density of CO and CH₄ for all the deposited ice mixtures was evaluated by Baratta *et al.* (2015). By supposing that the column density of N₂ was equal to the column density of both CO and CH₄ in the 1:1:1 (N₂:CH₄:CO) mixtures and by considering the measured thicknesses, Baratta *et al.* (2015) computed the density of the deposited ices. In particular, they found the following values: 0.68 g/cm³ (6 fringes), 0.67 g/cm³ (4 fringes) and 0.72 g/cm³ (2 fringes). These values can be considered the same within the experimental uncertainties that are mainly due to the procedure used to compute the column density.

An irradiation dose of 110 eV/16u is representative of the dose that occurs in the external layers of comets and TNOs on time scales on the order of the age of the Solar System (Strazzulla *et al.*, 2003). This dose is within the dosage required for the conversion of low molecular weight hydrocarbons into

refractory organic residues stable at room temperature (Strazzulla and Johnson, 1991). It is well known that ion irradiation induces several effects, including chemical changes (Hudson *et al.*, 2008; Palumbo *et al.*, 2008). The IR spectra of the ice mixtures after irradiation for the three considered thicknesses are reported in Fig. 2. The presence of many newly formed molecular species is evident. The strongest band of HCN (ν_3 , $\sim 3130\text{ cm}^{-1}$) (Hudson and Moore, 2003) is superimposed over a multi-component broad band observed between 3600 and 2600 cm^{-1} (see bottom panel of Fig. 2). This broad feature is distinctive of many N-H, O-H, and C-H bands. The spectral region between 2400 and 2000 cm^{-1} (top panel of Fig. 2) shows the presence of several new features, in particular $\sim 2340\text{ cm}^{-1}$ (CO_2), $\sim 2338\text{ cm}^{-1}$ (N_2), $\sim 2260\text{ cm}^{-1}$ (HNCO), $\sim 2235\text{ cm}^{-1}$ (N_2O), $\sim 2165\text{ cm}^{-1}$ (OCN^-), $\sim 2101\text{ cm}^{-1}$ (ν_1 HCN), $\sim 2092\text{ cm}^{-1}$ (CH_2N_2) (see, e.g., Grim and Greenberg, 1987; Hudson *et al.*, 2001; Hudson and Moore, 2003; Palumbo *et al.*, 2004; Sicilia *et al.*, 2012). This attests to the complex chemistry induced by ion irradiation on simple molecular ices. For further details on the chemistry of the irradiated $\text{N}_2:\text{CH}_4:\text{CO}$ ice mixture, the reader is referred to the work of Baratta *et al.* (2015).

Once irradiated, the frozen target was allowed to gently warm overnight and under vacuum to room temperature by switching off the cryostat. The day after, the organic residue was extracted from the chamber and a blank MgF_2

window was mounted. We let the pumping system evacuate the chamber for at least 24h before the MgF₂ window was cooled down to 17 K and another mixture was deposited and irradiated. This procedure was repeated for all the 30 samples prepared. After extraction from the chamber, an infrared (IR) spectrum, from 8000 cm⁻¹ down to the cut-off limit of the MgF₂ window at 1000 cm⁻¹, was obtained for each sample with a Bruker Equinox 55 FTIR spectrometer. No significant differences (standard deviation less than 2%) were observed between the IR spectrum acquired in air and the IR spectrum acquired in the chamber with the sample still under vacuum before extraction. In Fig. 3, the IR spectra of the organic residues obtained for the three considered thicknesses are reported. The labels 6f, 4f, and 2f in the figure indicate the organic residues obtained from irradiation of 6, 4, and 2 fringes “thick” ice mixtures respectively. In the figure, the thicknesses of the organic residues are also indicated. It should be considered that the thicknesses of 0.050 μm (2f), 0.135 μm (4f) and 0.180 μm (6f) of the organic residues are average thicknesses over all the 30 PSS samples. To obtain a precise measurement of the thickness of each individual flight sample, we used the optical constants derived by Baratta *et al.* (2015) to generate theoretical spectra that have been compared with the spectra of the corresponding organic residues acquired just after their production. For each organic residue, the thickness was measured as that which provides the best match between the

theoretical and the experimental spectra over an extended range (from the UV to the mid-IR). All the volatile species such as CO, CO₂, N₂ sublimated during the slow warm up to room temperature, and hence the thicknesses of the residues are considerably reduced with respect to the thicknesses of the ice mixtures. Many features are present in the residues. We note that the infrared spectra of the residues are quite similar to the infrared spectra of many tholins prepared by radiofrequency (RF) plasma discharge of N₂:CH₄ gas mixtures (Mutsukura and Akita, 1999; Imanaka *et al.*, 2004; Quirico *et al.*, 2008). The broad feature extending from 3500 to 2500 cm⁻¹ displays many sub-peaks that can be assigned to N-H, O-H, and C-H stretching vibrations. This broad band is consistent with the presence of -NH₂ and possibly -NH- functional groups of primary and secondary amine functions respectively. In particular, features extending from 3500 cm⁻¹ to 3300 cm⁻¹ are consistent with NH₂ groups connected to either aromatic and/or aliphatic functions, while the peak at ~3220 cm⁻¹ could be consistent with either ali-NH₂ or a -NH- functional group (Quirico, *et al.*, 2008). Sub-peaks at ~2980 cm⁻¹ and ~2940 cm⁻¹ indicate the presence of -CH₃ and -CH₂- stretching vibrations. The position of these peaks, shifted at higher wavenumbers with respect to that of alkyl chains, indicates the presence of CH₂ and CH₃ groups that are connected to nitrogen atoms, although the specific chemical groups involved cannot be identified (Quirico *et al.*, 2008).

The middle of the spectral range reveals a structured feature that peaks at $\sim 2200\text{ cm}^{-1}$. This feature is due to $-\text{C}\equiv\text{N}$ (nitrile) and $-\text{N}\equiv\text{C}$ (iso-nitrile) terminal groups (Mutsukura and Akita, 1999; Imanaka *et al.* 2004); the carbodiimide function ($-\text{N}=\text{C}=\text{N}-$) has also been proposed to give a contribution to this feature (Imanaka *et al.*, 2004). This structured feature also exhibits a well-resolved shoulder that peaks at $\sim 2150\text{ cm}^{-1}$. The shoulder is better resolved with respect to the other sub-peaks observed, and its width is similar to that of the OCN^- feature detected in the irradiated ice mixture before warming up. Although the OCN^- feature peaks at 2165 cm^{-1} at 17 K, its peak position shifts to lower wavenumbers, and its intensity significantly decreases during warm up toward room temperature (Palumbo *et al.*, 2004, Accolla *et al.*, 2018). Hence, the shoulder seen at $\sim 2150\text{ cm}^{-1}$ could be due to residual OCN^- trapped in the organic residue.

At lower wavenumbers, a structured broad asymmetric feature, peaking near 1600 cm^{-1} and extending down to the cut-off limit of the MgF_2 window, is present. In analogy to some tholins, the two main peaks at $1650\text{-}1630\text{ cm}^{-1}$ and 1560 cm^{-1} could be indicative of the presence of the following functional groups: $\text{C}=\text{C}$ aromatics, $\text{C}=\text{N}$ heteroaromatics, aromatic- and aliphatic- NH_2 bend (Imanaka *et al.*, 2004; Quirico *et al.*, 2008). A contribution to the 1630 cm^{-1} feature could also be due to O-H bending vibrations. We noted the

presence of a weak shoulder at 1710 cm^{-1} that is due to C=O stretching vibrations. Three weak sub-peaks are present at 1450 cm^{-1} , 1415 cm^{-1} , and 1380 cm^{-1} ; the first two are probably due to the bending modes of the -CH₃ and -CH₂- groups, which give rise to the stretching modes seen at $\sim 2980\text{ cm}^{-1}$ and $\sim 2940\text{ cm}^{-1}$, while the peak at 1380 cm^{-1} points to the presence of C-CH₃ or C-(CH₃)₂ groups (Quirico *et al.*, 2008). With regard to the remaining sub-peaks, the feature that peaks at 1339 cm^{-1} can be assigned to C-N (aromatic) stretching vibration (Imanaka *et al.*, 2004), while we were not able to identify the feature seen at 1240 cm^{-1} . A summary of the IR features seen in the organic residue is reported in Table 1.

From this analysis, the chemical complexity exhibited by the organic residues becomes evident; in particular, many chemical groups have been assigned, although no specific molecular species have been recognized. Furthermore, we know that the residues probably contain complex species that have not been detected by IR spectroscopy. As an example, complex molecules such as succinonitrile 'NCCH₂CH₂CN', 3-amminocrotonitrile '(CH₃)C(NH₂)CHCN' etc., have been identified in the organic residues obtained by ion irradiation of nitrile (CH₃CN) containing ice; these astrobiologically relevant molecules have a very low abundance and have been identified by gas chromatography-mass spectroscopy (GC-MS), while

they are difficult or even impossible to identify by IR spectroscopy (Hudson *et al.* 2008). Furthermore, by comparing the results obtained on residues produced by UV photolysis and ion irradiation of different ice mixtures, Hudson *et al.* (2008) suggested that energetic processing of almost any ice mixture containing C, H, N, and O atoms probably results in the formation of aminoacid precursors that, if hydrolyzed, give rise to the aminoacids themselves. Similar conclusions have been drawn by Materese *et al.* (2014), who found, in the organic residue of a far-UV irradiated N₂:CH₄:CO (100:1:1) ice mixture, complex molecules such as glyceric acid, lactid acid, urea, etc. Hence, even if IR spectroscopy does not reveal them due to its relatively low sensitivity (Hudson *et al.*, 2008; Materese *et al.*, 2014), complex molecules of astrobiological interest are probably present in the organic residues described in this work.

In addition to FTIR spectroscopy, the 30 organic samples were characterized by vacuum-UV and UV-Vis-NIR transmission spectroscopy. In particular, on the basis of the transmission spectra acquired on the 30 samples over the three different thicknesses, Baratta *et al.*, (2015) derived the optical constants (*n* and *k*) of the organic residue in the range of 500-84000 cm⁻¹ by adopting the Kramers-Kronig constrained variational dielectric function (KKCVDF) fitting procedure (Kuzmenko, 2005). In particular, Baratta *et al.* (2015) obtained a Kramers-Kronig constrained set of optical constants as the best fit solution to

the experimental transmission spectra together with the three average best fit thicknesses of the 6f, 4f, and 2f residues. The optical constants of the organic residue are available on line at the link: <http://www.oact.inaf.it/weboac/labsp/ORPSS.html>.

2.2 Exposure on the International Space Station

The samples of all the experiments of the Exposer-R2 mission were placed in special carriers accommodated in three different trays, in particular all the PSS samples were placed in tray number 3 (see Cottin et al. in prep for further details). PSS samples, placed in individual sample containers called cells, were integrated in two stacked carrier layers each in three compartments of tray 3. Only the samples in the top carriers were exposed to the solar UV radiation, while the samples in the underlying carriers were completely shaded, providing the respective “in-flight” dark controls. Three temperature sensors were attached to the bottom side of each tray. In addition, tray 3 was equipped with the active dosimetry R3D-R2, a radiation sensor package for ionizing radiation and UV measurements. Environmental data of the Expose R2 mission (temperatures, UV exposure, etc.) were collected every 10 s and stored and down linked as telemetry.

On the night of 2014 July 23, the cargo 56P progress carrying the three trays of the Expose R2 mission was launched from Baikonur (Kazakhstan) to the

ISS. After docking on 2014 July 24, the three trays were stored inside the ISS. On August 6, the trays were integrated in the monoblock Expose-R2 that remained inside the ISS until it was installed outside on the URM-D platform, during the EVA (Extravehicular Activity) -39 on 2014 August 18. The vacuum exposure period of the mission began on 2014 August 20, when the venting valves, of the still covered Expose-R2, were opened remotely by the Mission Control Center in Moscow (MCC-M). During the next 62 days, Expose-R2 was subjected to an out gassing period while shielded from solar UV irradiation. On 2014 October 22, the Expose-R2 cover was removed during EVA-40, starting the UV exposure of the experiments. Solar UV exposure ended on 2016 February 3, when Expose-R2 was returned to the ISS interior during EVA-42. On 2016 March 2, tray 3 landed at Karaganda (Kazakhstan), on-board the Soyuz 44S return capsule. The exposure of the samples to solar UV radiation in low Earth orbit outside ISS lasted 469 days, while the total exposure to space vacuum was 531 days.

In parallel to the space experiment, a Mission Ground Reference (MGR) experiment was performed at the German Aerospace Center (DLR), Cologne, Germany. The experiment used a set of trays and carriers similar to those exposed outside ISS. In particular, three ground trays identical to the flight trays, with respect to the sample interface, were loaded with a full set of samples identical to the flight ones. The trays were placed in a temperature-

controlled interface and connected to the vacuum facility of the PSI (Planetary and Space Simulation Facilities) at DLR. Automated temperature simulation of the MGR trays started with application of transport temperatures. As soon as the Expose-R2 temperatures in orbit became available by telemetry, they were applied and updated every 10 seconds, with a maximum delay of two months due to the reformatting and analysis process of data. The highest and lowest temperatures experienced by Expose-R2 trays in orbit were 58° C (tray 3, compartment 4) and -20.9° C (tray 3, compartment 2), respectively. In orbit, temperature fluctuations follow warm periods (temperatures up to 50° C on average), which are followed by cold periods (temperatures down to -20° C on average) with a slow periodicity of about 1 month due to the variation of the ISS orbital plane. Superimposed to this slow variation, Expose-R2 trays experienced a fast variation of temperature, with a periodicity of nearly 90 minutes due to the ISS day/night times of each orbit. Day/night temperature fluctuations were approximately 10° C during warm periods and a few degrees during cold periods.

As in space, only the top carriers of the Expose-R2 MGR were irradiated with a SOL2000 metal halide lamp-based solar simulator (by Dr. Hönle GmbH), which provided a continuum spectrum with wavelength >200 nm; the spectral irradiance of the solar simulator can be found in figure 3 in the work of Onofri *et al.*, (2008). The MGR tray compartments were irradiated at ground

individually with the same UV fluences accumulated by the Expose-R2 trays outside ISS during the entire mission. Unfortunately, due to a malfunction of the system, irradiation was not performed under vacuum. The UV fluences were provided by RedShift Design and Engineering BVBA, Belgium, contracted by ESA (calculated ultraviolet radiation fluence data). It should be noted that the samples were not always facing the Sun because ISS orbits around Earth and changes its orientations relative to the Sun. In addition, shadows by solar panels and the ISS environments in the vicinity of the Expose-R2 facility (radiators, etc.) significantly decrease the effective insolation. The orientation of the ISS has been computed and the UV fluence at each sample position has been evaluated taking into account the daily variation of insolation along the whole mission duration. The UV sensor data have been used to verify the adopted models. The variation of the solar irradiance due to the eccentricity of the Earth orbit and to the solar activity have also been taken into account. This was done by RedShift considering the actual solar spectrum, averaged along the mission duration, measured by the satellite SORCE. The transmittance of the windows, weighted by the solar spectrum, has also been considered. Hence the mean UV fluence for samples in each compartment has been determined. Fluences, in the wavelength range (200-400 nm), have been applied individually on the corresponding compartment and experiment during the MGR. In order to find out any possible modification induced by the thermal cycles experienced in orbit and

at ground, in the case of the PSS experiment, an additional ground reference full set of samples has been stored under vacuum at a constant temperature of 5° C during the MGR for comparison. The scheme showing the experimental breakdown of samples between the ISS and DLR Cologne, and the environmental conditions to which they have been exposed is reported in Fig. 1(b).

With regard to our samples, as previously indicated, we prepared 30 organic residues that had three different thicknesses (10 samples for each thickness). Twelve of them were accommodated in Expose-R2 outside ISS. In particular, two samples for each thickness (six in total) were exposed in the top carrier of tray 3 to solar UV photons, ionizing cosmic radiation, and temperature cycling. The remaining flight samples (two for each thickness) were placed in the underlying carrier where they were exposed only to ionizing cosmic radiation and temperature cycling, providing the in-flight dark controls. The remaining 18 samples were part of the ground MGR simulation experiment. Referring to the ground samples, two samples for each thickness were exposed to the PSI solar simulator and temperature cycling in the top carrier of Expose-R2 MGR facilities, whereas the other two dark control samples for each thickness, were exposed only to temperature cycling in the underlying carrier. Lastly, the remaining two ground samples for each thickness were stored under vacuum at 5° C.

Following the work of Baratta *et al.* (2015), we evaluate here the irradiation doses suffered by the flight exposed organic residues. UV photons (i.e., $E > 3.1$ eV, $\lambda < 400$ nm) have enough energy to break chemical bonds and initiate substantial chemical changes (Stern, 2003), where a photon energy of 3 eV corresponds to the dissociation threshold in the gas phase of relatively weak bounded species such as CH₂ (Heays *et al.*, 2017).

By considering the absorption coefficient of the organic residue (Baratta *et al.*, 2015) and the ASTM E-490 standard solar spectrum for space application (that is given down to $\lambda=119.5$ nm) over the range ($119.5 < \lambda < 400$ nm), the UV irradiation dose suffered by the 2f (50 nm) organic residue over the 2111 hours of equivalent to perpendicular solar irradiation, as determined by RedShift, is $4.7 \times 10^5 \pm 5\%$ eV/16u. If we consider only the more energetic, vacuum UV, solar photons ($E > 6.2$ eV, $\lambda < 200$ nm) the irradiation dose over the range ($119.5 < \lambda < 200$ nm) is $1.2 \times 10^3 \pm 14\%$ eV/16u. The errors given derive from the uncertainties associated with the equivalent to perpendicular solar irradiation exposure time; they were provided by RedShift and depend on the particular frequency band considered. The main sources of uncertainties are as follows: the geometrical model (depends on samples position), window transmission (depends on the frequency band), variation in solar irradiance (depends on the frequency band), and reflections (depend on the samples position and the frequency band). For the Expose-R2 mission, the total contribution of ionizing radiation given by GCRs,

South Atlantic Anomaly (SAA) protons and Outer Radiation Belt (ORB) electrons, reached dose values up to 1 Gray (Gy) (Berger *et al.*, 2018 this volume). One Gy corresponds to $1.7 \cdot 10^{-7}$ eV/16u, and hence the effects given by ionizing radiation are negligible with respect to those given by solar UV photons, even by considering only the more energetic ($E > 6.2$ eV) solar photons.

An accurate description of the Expose-R2 and MGR hardware and of the different mission phases can be found in the work of Rabbow *et al.* (2017).

3. RESULTS

3.1 Comparison between the sample controls

The post-flight FTIR, UV-Vis-NIR, and VUV spectra of the organic residues were acquired by using the same conditions (i.e., resolution, sampling, etc.) that were adopted for the pre-flight samples. In this article, we focus on the results obtained by FTIR spectroscopy. In particular, FTIR spectra were acquired in air at normal incidence, with a resolution of 1 cm^{-1} and a sampling of 0.25 cm^{-1} . Every sample on the ISS and at ground was realized in duplicate. We observed no significant differences in the profile of the IR features between the duplicates, and only a small variation in the intensity of the IR features was observed. In particular, with regard to all the duplicates, we found an average percentage

variation of the intensities of the 3220, 2200, and 1600 cm^{-1} features of $10\% \pm 6\%$, $10\% \pm 8\%$, and $12\% \pm 9\%$ respectively. To improve the signal to noise ratio of the IR spectra of the organic residues, in the following for any thickness and typology, we use the average spectrum obtained from the duplicates.

In Fig. 4, a comparison between the in-flight dark controls and the DLR (MGR) controls is reported for the three different thicknesses. From Fig. 4, it is evident that, for each thickness, no systematic differences can be found between the controls. In particular, the differences observed between the IR spectra of different controls are similar to the differences found between the IR spectra of duplicates. Hence, the organic residues are very stable with respect to the temperature cycles experienced in orbit. On the other hand, by comparing Fig. 4 and Fig. 3, a decreased intensity of the $\text{-C}\equiv\text{N}$ feature at $\sim 2200 \text{ cm}^{-1}$ with respect to the other two main broad features at $\sim 3220 \text{ cm}^{-1}$ and $\sim 1600 \text{ cm}^{-1}$ and the disappearance of the $\sim 2150 \text{ cm}^{-1}$ shoulder is evident in the post-flight samples. To discern any temporal modification of the FTIR spectra of the organic residues that we prepared for the PSS experiment, we also prepared an additional set of samples identical to the flight ones that we monitored by FTIR spectroscopy during the Expose-R2 mission. We found that the two main broad features displayed very small changes, while the $\text{-C}\equiv\text{N}$ feature decreased at a rate that was fast at the beginning but slower and slower with

time. We verified that the fast decrease of the $\text{-C}\equiv\text{N}$ feature, observed for several days after the production of the residue, occurred as well for the organic residues kept under UHV condition. In Fig. 5, the band area of the $\text{-C}\equiv\text{N}$ feature of the lab-control organic residues is reported versus the time elapsed since their production. In the same figure, the band area obtained for the flight control PSS residues is also reported. Note that in order to minimize any possible damage or contamination, only the band area obtained just after the production of the organic residue was measured for the flight samples. Indeed, just after their production and spectra acquisition, all the PSS organic residues were stored in special teflon containers thought to preserve the samples from moisture, contamination, and damage. All our samples are, at present, still stored in these containers. After the end of the mission, two additional spectra were acquired. For each thickness, the lab-control and flight-control samples showed a common trend in Fig. 5. In particular, after a fast decrease at the beginning, the $\text{-C}\equiv\text{N}$ feature band area almost stabilizes after 200 days and did not vary further, within uncertainties, after 700 days. The two vertical dashed lines in the figure indicate the start and the end of exposure outside the ISS; hence, the organic residues were already stabilized during exposure to the solar UV photons. The reason of the decrease of the 2200 cm^{-1} feature is not clear. It could be either due to some volatile, that is, $\text{C}\equiv\text{N}$ bearing species that leave the residue, or to the reaction and

modification of a $\text{-C}\equiv\text{N}$ bearing species with other species in the matrix. If the shoulder seen at $\sim 2150\text{ cm}^{-1}$ in the pre-flight organic residues is actually due to OCN^- , the reduction of the intensity of the 2200 cm^{-1} feature could have an alternative explanation. In particular, we know that ion irradiation of a different mixture ($\text{N}_2:\text{CH}_4:\text{H}_2\text{O}$, 1:1:1) leads to the formation of an organic residue that exhibits a feature peaking at 2158 cm^{-1} , with a different profile with respect to the 2200 cm^{-1} feature seen in the PSS organic residues that is attributed to OCN^- (Accolla, *et al.*, 2018). The authors observed that, in those experiments, the OCN^- feature exhibits a gradual reduction in intensity over time, decreasing by 60% after two weeks and almost by 100% after 150 days, that is, with similar a time scale of the reduction of the 2200 cm^{-1} feature. Hence, it is possible that the decrease of the 2200 cm^{-1} feature seen in the PSS organic residues is due to the decrease of the OCN^- ion/salt either desorbing or reacting with time. After the complete disappearance of the OCN^- from the matrix, the 2200 cm^{-1} does not change further, which indicates that the nitriles/isonitriles terminal groups are stable in the organic residue.

3.2 Space exposed organic residues

In Fig. 6, a comparison between the IR spectra of the flight solar UV exposed organic residues (solid lines) and the corresponding in-flight dark controls (dotted lines) is reported for the three considered thicknesses. The presence of

relatively strong -CH_3 and $\text{-CH}_2\text{-}$ stretching features (2950, 2930, and 2860 cm^{-1}) and bending features (1458 and 1380 cm^{-1}) is evident in the exposed samples. We notice that the intensity of these features is inversely correlated with the thickness of the samples, and the features are not observed in the in-flight dark controls. As was already observed for the former Expose-R mission (Demets *et al.*, 2015), the aliphatic features are due to a contamination layer deposited on the inner side of the solar exposed MgF_2 windows. It is thought that the contamination is due to the release of outgassing products from the facility itself and its samples, which were subsequently photo-processed by solar UV at the inner window surfaces into a thin contamination layer (Rabbow *et al.*, 2017). In general, the contamination observed on Expose-R2 seems to be lesser than after the Expose-R mission. This improvement was due substantially to three reasons: 1) Expose-R2 used the Expose-R module that has been already outgassing in orbit during the Expose-R mission, 2) the duration of exposure to solar UV was shorter for Expose-R2, 3) Expose-R2 has been subjected to a 62 days outgassing period while shielded from solar UV irradiation; this procedure has not been previously adopted for Expose-R (Rabbow *et al.*, 2017). It should be noted that the organic residues are on the internal surfaces of the MgF_2 windows, and thus the contamination layer is accreted on the organic residues from the opposite side with respect to that exposed to solar UV photons. Hence, the

contamination layer did not interfere with the solar UV exposure. To “clean” the spectra from the contaminant features, we acquired a transmission spectrum of a flight exposed MgF₂ clean (blank) window on which the contamination layer was formed. Once normalized to the continuum and converted in optical depth units, the contamination spectrum was subtracted from each sample thickness by using a different scaling factor that was chosen to minimize the contribution of the contaminant CH features on the residual spectra. The corrected flight exposed sample spectra are reported in Fig. 6 dashed lines. We notice that the contaminant features almost disappear in the corrected spectra. Furthermore, thicker residues correspond to thinner contamination layers. On the other hand, thicker organic residues absorb more UV light, and hence the contamination layers can be considered to have a common composition and different thicknesses that are directly related to the UV fluences transmitted by the MgF₂ plus organic residues assemblies.

From the comparison between the flight exposed and the in-flight dark control samples we notice the following: 1) An overall decrease in the exposed samples of the two main broad features and of the -C≡N feature for all the thicknesses except for the thickest 6f sample. In particular, an increase in intensity of the broad 1600 cm⁻¹ feature was observed in the region between 1600 and 1240 cm⁻¹, for the 6f sample. 2) Thinner is the sample, stronger is the reduction of intensity for the two main broad feature and for

the $\text{-C}\equiv\text{N}$ feature. 3) The reduction in intensity of the features is accompanied by more or less evident changes of the profile of the bands. All these effects are particularly evident in the 2200 cm^{-1} $\text{-C}\equiv\text{N}$ feature.

It should be noted that the solar UV flux is exponentially attenuated into the film, and hence thinner residues received higher average UV doses. The higher UV doses absorbed by thinner residues explains why the relative reduction of intensity of the features increases from the thickest 6f to the thinnest 2f organic residue. In Fig. 7, the normalized spectra of the in-flight dark controls and the flight exposed samples are presented. Very small differences, having standard deviation with respect to the average normalized spectrum of about 2%, are evident in the profile of the main features for the three considered thicknesses in the flight control samples; this points out that the organic residue is quite homogeneous versus the thickness as already observed by Baratta *et al.* (2015). On the other hand, a variation of profile of the 3220 , 2200 , and 1600 cm^{-1} features is evident in the exposed samples with respect to the dark controls. Nevertheless, for the 3220 and 2200 cm^{-1} features, the profile is very similar for the three different thicknesses in the flight exposed samples. Hence, the relative abundance of the species that contribute to the 3220 cm^{-1} (-NH_2 , -NH- , etc.) and 2200 cm^{-1} ($\text{-C}\equiv\text{N}$, $\text{-N}\equiv\text{C}$, etc.) features changes at low UV irradiation doses but does not change further at higher doses from the thickest 6f to the thinnest 2f flight exposed organic

residue. On the other hand, the profile of the 1600 cm^{-1} (C=C, C=N, etc.) feature still changes up to the highest doses corresponding to the 2f thinnest sample. A possible explanation for this trend is given in the Discussion section.

3.3 VUV, UV-Vis-NIR Spectroscopy

The transmission spectra of the flight exposed samples in the UV-Vis-NIR and VUV range were also acquired. Although the contaminant layer deposited on the inner side of the flight exposed samples did not interfere with the solar UV exposure, it strongly affects the transmission spectra in the UV-Vis range. This was evident when we acquired the transmission spectra of the contaminated flight exposed blank MgF_2 window (see section 3.2) in the UV-Vis-NIR and IR range. To clean the UV-Vis-NIR spectra, we followed the same procedure described in section 3.2 using the same scaling factors determined for the IR spectra. We found that, in the Vis-NIR range between 400 and 1200 nm, the absorption coefficient increases with the dose from the thickest (6f) to the thinnest (2f) sample. Only small differences of the absorption coefficient (within 7%) were found, between the different thicknesses, in the UV range (200-400 nm). Nevertheless, unlike the IR case, we could not perform any normalization to the continuum to correct the spectra. This is because the variations of transmittance due to the variation of

absorption and continuum (dispersion and interference effects) cannot be easily disentangled in the UV-Vis range. Hence, the conclusions given above on the variation of the absorption coefficient must be considered indicative.

3.4 The 2200 cm^{-1} feature

Since the 2200 cm^{-1} feature does not overlap with any other feature, we can try to obtain semi-quantitative information on the relative abundances of $\text{-C}\equiv\text{N}$ and $\text{-N}\equiv\text{C}$ species by adopting a fit procedure (Quirico *et al.*, 2008). We were able to obtain a good fit by using 5 Gaussian components with peak positions similar to those used by Mutsukura and Akita (1999). In Fig. 8, the best fit to the 2200 cm^{-1} feature for the 6f flight control and flight exposed samples is reported. The best fit output parameters and the corresponding uncertainties are reported in Table 2. Following the work of Mutsukura and Akita (1999), the peaks at 2105, 2145, and 2190 cm^{-1} are associated to terminal $\text{-N}\equiv\text{C}$ iso-nitrile units, while the peaks at 2220 and 2252 cm^{-1} are associated to terminal $\text{-C}\equiv\text{N}$ nitrile units. By assuming that the band strength of the five sub-components is the same, we obtain a relative amount of nitrile over iso-nitrile terminal units ($\text{-C}\equiv\text{N}/\text{-N}\equiv\text{C}$) of 0.23 and 0.9 before and after UV space exposure respectively. Nevertheless, these relative amounts should be considered as indicative for two reasons: 1) according to Quirico *et al.*, (2008), conjugation can lower the nitrile band position down to 2180 cm^{-1} , in

this case the 2190 cm^{-1} sub-component would be due to a nitrile terminal unit,
2) the band strength can be different for the five sub-components.

By using the spectra obtained for the three different thicknesses, it is possible to discern the effect of solar UV photons at different depths in the organic residue. In particular, following the work of Baratta *et al.* (2002), we indicate with τ_{2f} , τ_{4f} , and τ_{6f} the optical depths of the 2f, 4f, and 6f organic residues respectively. Since the residues have been exposed to the same UV fluences, the optical depth of the slab at a depth between 50 nm (2f) and 135 nm (4f) is given by the relation: $\tau_{4f-2f} = \tau_{4f} - \tau_{2f}$, and the optical depth of the slab between 135 nm and 180 nm (6f) is given by: $\tau_{6f-4f} = \tau_{6f} - \tau_{4f}$. In Fig. 9, the spectra of the slabs at different depths for the flight exposed and in-flight dark control organic residues are reported. From Fig. 9, it is evident that the decrease in intensity of the 2200 cm^{-1} feature is larger at lower depths into the organic residue, and hence deeper layers received a lower UV dose.

To evaluate the lifetime at 1 au of the $\text{-C}\equiv\text{N}$ plus $\text{-N}\equiv\text{C}$ bearing species versus the depth in the organic residue, we assume that the cumulative band area is proportional to the number of $\text{-C}\equiv\text{N}$ plus $\text{-N}\equiv\text{C}$ terminal units and the band strength is $4.0 \times 10^{-17}\text{ cm/molecule}$ (Demyk *et al.*, 1998). In the following, we also use the suffix CN to mean the cumulative number of $\text{-C}\equiv\text{N}$

and -N≡C terminal units. We assume that the decrease of the number of CN can be described by an exponential decay, in particular:

$$dN_{CN} = N_{CN}^0 e^{-\Phi_{UV}^{eff}(x)t\sigma(CN)} dx \quad (2)$$

where: dN_{CN} is the number of CN per square centimeter contained in a layer of thickness dx after an exposure time t at a depth x in the organic residue; $\Phi_{UV}^{eff}(x)$ is the effective UV flux at a depth x ; $\sigma(CN)$ is the destruction cross section and N_{CN}^0 is the number density of CN (cm^{-3}) before exposure. We can also write:

$$\Phi_{UV}^{eff}(x) = \Phi_{UV}^{eff}(0) e^{-\alpha(UV)_{eff} \cdot x} \quad (3)$$

where $\Phi_{UV}^{eff}(0)$ is the incoming effective UV flux and $\alpha(UV)_{eff}$ is the effective UV absorption coefficient. If we indicate the decay rate with $\Gamma = \Phi_{UV}^{eff}(0) \cdot \sigma(CN)$, by integrating (Equation 2) we obtain:

$$n_{CN} = \int_{x_1}^{x_2} dN_{CN} = N_{CN}^0 \int_{x_1}^{x_2} e^{-\Gamma \cdot t \cdot e^{-\alpha(UV)_{eff} \cdot x}} dx \quad (4)$$

where n_{CN} is the column density (cm^{-2}) of CN contained in the slab at a depth between x_1 and x_2 after an exposure time t .

The computed CN column densities of the flight exposed and flight dark control samples versus the thickness are reported in Fig. 10. As expected, the

control samples stay along a straight line that passes through the origin. We compute the initial number density of CN as the slope of the linear best fit to the flight dark controls; in particular, we found: $N_{CN}^0 = 1.94 \times 10^{21} \text{ cm}^{-3}$ that, for a density $\rho = 1.17 \text{ g cm}^{-3}$ (Baratta *et al.*, 2015) of the organic residue, corresponds to an abundance of the CN units of nearly 7% in weight.

We use a non-linear best fit routine, written in IDL language, to fit the column densities of the flight exposed residues with Equation (4). In Equation (4), we used the N_{CN}^0 value determined above and an exposure time t , over the whole duration of the mission, of 2111 hours of equivalent to perpendicular solar irradiation at 1 au. The equivalent exposure time was provided by RedShift and has been corrected for the transmission losses of the MgF_2 windows. In Fig. 10, the good agreement between the theoretical fit (solid curve) and the experimental points is evident; the corresponding best fit values of the free parameters $\alpha(UV)_{eff}$ and Γ are $5.1 \text{ } \mu\text{m}^{-1}$ and 10.6 yr^{-1} respectively. If we consider a thin ($< 50 \text{ nm}$) organic residue, the lifetime τ of the CN units within the film is quite short, in particular $\tau = \frac{1}{\Gamma} = 0.1 \text{ yr}$. On the other hand, the UV solar flux is exponentially attenuated within the organic residue, and hence CN units at greater depths in thicker organic residues can survive much longer times. By using Equation (4) and the determined $\alpha(UV)_{eff}$ and Γ values, we can predict the survival time of the CN units versus the depth at 1

au. In particular, we compute the CN column density, within a 50 nm thick slab, at variable depth and at different exposure times. The calculated survival curves are reported in Fig. 11. The first curve on the left corresponds to the PSS equivalent to perpendicular solar irradiation exposure, and the solid black circles are the interpolated column densities corresponding to the 2f, 4f, and 6f exposed organic residues. From Fig. 11, it is evident that the CN bearing species can survive many years in organic residues thicker than 1 μm .

3.4 MGR exposed organic residues

In the case of the MGR samples that were irradiated with the solar simulator SOL2000, only wavelengths greater than 200 nm have been used. On the other hand, for Expose-R2 in space, the MgF_2 windows let photons with wavelength down to 110 nm to reach the samples. One of our objectives was to compare the effects on space exposed samples and ground exposed ones in order to reveal the contribution of the more energetic (below 200 nm) vacuum UV solar photons to modify the samples. Unfortunately, our MGR samples exposed to the solar simulator were contaminated during the ground exposure by a relatively thick film accreted on the MgF_2 windows from the opposite side to that where the organic residues were deposited and, hence, from the same side of UV exposure. We noted that this contaminant layer was present, with variable amount, on all our MGR samples exposed to the solar simulator but

not in any other unexposed MGR sample. From the infrared analysis, we know that the contaminant MGR layer spectrum presents a broad strong feature peaking $\sim 3450 \text{ cm}^{-1}$, indicating the presence of -NH- stretching vibrations (Imanaka *et al.*, 2004). This feature strongly affects the 3220 cm^{-1} band of the organic residues. Furthermore, since the contaminant layer is on the same side of the incoming UV photons, it interfered with the exposure to the solar simulator. For this reason we decided not to use the spectra of the MGR UV exposed samples for any quantitative comparison with the flight exposed samples. Due to a technical problem, irradiation was not performed under vacuum conditions. In these circumstances, the contamination may have an atmospheric origin. Nevertheless, after cleaning an MGR exposed sample (4f) from the contaminant layer, by polishing the corresponding MgF_2 window face, we saw that the effects produced by the solar simulator on the IR spectrum were similar, from a qualitative point of view, to the effects seen on the spectrum of the corresponding flight exposed organic residue. This should imply that, even less energetic solar UV photons with wavelength greater than 200 nm, were effective in modifying the chemical structure of the organic residues.

4. DISCUSSION

4.1 Solar UV modification of the organic residue

We saw in section 3.2 that one general effect of the exposure to the UV solar photons is the decreased intensity of the main infrared features except for the 1600 cm^{-1} feature that, in the region between 1600 and 1240 cm^{-1} , gets more intense at low exposure doses (6f). The reduction in intensity of the features is accompanied by a more or less evident variation of the profile. For the 3220 cm^{-1} and 2200 cm^{-1} features, the profile does not change after the initial variation corresponding to the thickest (6f) organic residue. This circumstance seems to indicate that functional groups that give rise to the 3220 cm^{-1} ($-\text{NH}_2$, $-\text{NH}-$, etc) and 2200 cm^{-1} ($-\text{C}\equiv\text{N}$, $-\text{N}\equiv\text{C}$, etc.) features decrease in number at higher UV exposure without changing their relative amounts. On the other hand, the profile of the 1600 cm^{-1} feature still changes up to the highest dose (2f sample). In particular, while the peak at 1590 cm^{-1} decreases more in thinner (higher UV doses) exposed samples (Fig. 6), the wing at lower wavenumbers does not significantly decrease, with respect to the dark control samples, for the three considered thicknesses. Except for the CH bending, according to Ferrari *et al.*, (2003), C=C or mixed C=N both in sp^2 chains and rings are responsible for the broad feature seen in the $1000\text{-}2000\text{ cm}^{-1}$ region for N rich hydrogenated amorphous carbon nitride (a-C:N:H). We tentatively attribute the initial increased intensity of the wing of the 1600 cm^{-1} feature to the possible partial conversion of the $-\text{C}\equiv\text{N}/-\text{N}\equiv\text{C}$ terminal units lost by the 2200 cm^{-1} band into C=N units added to the C=C skeleton. Indeed it is

generally found that nitrogen increases the intensity of the 1600 cm^{-1} feature seen in a-C:N:H (Ferrari *et al.*, 2003). This can be understood if it is taken into account that nitrogen polarizes all the sp^2 bonds of the conjugated system to which nitrogen is a terminal atom. Thus, it increases the dipole moment of each C=C bond giving rise to an increase in the IR intensity that is larger for longer conjugation. Furthermore, N addition also increases the clustering of the sp^2 phase, and this also increases the IR intensity. In particular, the integrated absorbance in the $1000\text{-}1800\text{ cm}^{-1}$ region can be used as an indicator of the N content (Ferrari *et al.* 2003).

Sharp $1600\text{-}1580\text{ cm}^{-1}$ peaks are typical of a-C:N:H with high hydrogen content (Ferrari *et al.*, 2003). In particular, these authors observed, in a high H content a-C:N:H film, a sharp peak at $1580\text{-}1600\text{ cm}^{-1}$ that was as intense as twice the broad wing at 1300 cm^{-1} . We interpret the decrease of the peak at 1590 cm^{-1} with respect to the wing at 1300 cm^{-1} , observed in the flight exposed residues, as due to a progressive loss of hydrogen during solar UV exposure. The loss of hydrogen is also evidenced by the progressive decrease of the broad feature at 3220 cm^{-1} due to -NH-, -NH₂, -CH₃, -CH₂-, and possibly -OH functional groups.

Hence, as a consequence of the solar UV exposure, the organic residue is stabilized into a C=C/C=N amorphous carbon network, while losing -NH-, -NH₂ (amine) and -C≡N/-N≡C (nitrile/iso-nitrile) functional groups.

To determine $\alpha(UV)_{eff}$ and Γ , we implicitly assumed that the thickness of the flight exposed samples did not change as a consequence of exposure to solar UV photons. On the other hand, we know that the number of functional groups such as amines and nitriles decreased in the flight exposed samples, and hence it is plausible that the thickness decreased accordingly if the missing/destroyed species, or some fragments of them, left the UV exposed samples. If we indicate with x and x^{exp} the thickness before and after UV exposure respectively, in first approximation we can assume that: $x^{exp} = c \cdot x$, where the scale factor c is a constant ≤ 1 .

By substituting x^{exp} in equation (4), the best fit effective absorption coefficient becomes $\alpha(UV)_{eff}/c$ and the destruction rate Γ does not change. Since the effective absorption coefficient would increase ($c \leq 1$), we assume, by continuing to suppose that the thickness after solar UV exposure does not change, a conservative point of view with respect to the nitriles survival ability at a given depth in the organic residue. Furthermore, although in principle the thickness of the exposed samples could decrease greatly, we observed that the wing of the 1600 cm⁻¹ feature at low wavenumbers does not

significantly decrease in the flight exposed samples, with respect to the dark controls, for the three considered thicknesses. In particular, we observe that, due to the solar UV exposure, the organic residue stabilizes into a C=C/C=N amorphous carbon network, and hence the scaling factor c is not too far from 1. On the other hand, the scaling factor itself could depend on the sample thickness; nevertheless the good fit obtained in Fig. 10 should indicate that the scaling factor c is nearly constant. So far we have implicitly assumed that $\alpha(UV)_{eff}$ does not vary as well with the solar UV exposure, and thus it is constant versus the thickness. On the other hand, the conversion of $-C\equiv N$ into C=N/C=C units probably determined an increase of the absorption coefficient in the UV-Vis range. Hence, we would expect an increase in the average (versus thickness) absorption coefficient. In this case, the scaling factor c defined before should decrease with the sample thickness. Nevertheless, we already noted that the scaling factor c is not too far from 1. Hence, we do not expect a dramatic variation of the absorption coefficient in the range of the UV photons that are effective in destroying the -CN units. This confirms the small variation of the absorption coefficient versus the thickness found in the UV range (see section 3.3). In any case, in addition to the wavelength of the solar UV photons, the $\alpha(UV)_{eff}$ coefficient obtained by the fit procedure should be regarded as effective also with respect to the UV exposure. If we suppose, in first approximation, that the absorption coefficient of the pre-

flight organic residue is equal to the absorption coefficient of the organic residue after solar UV exposure, we can obtain some additional information on the wavelength of the solar UV photons that were active in modifying the chemical structure. The absorption coefficient α is related to the imaginary part of the complex index of refraction k through the relation: $\alpha = 4\pi k/\lambda$, where λ is the wavelength. By considering the optical constants of the pre-flight organic residues given by Baratta *et al.* (2015), and the effective absorption coefficient derived by the best fit procedure $\alpha(UV)_{eff} = 5.1 \mu\text{m}^{-1}$, we obtain an effective wavelength $\lambda_{eff} = 0.342 \mu\text{m}$, confirming that less energetic solar UV photons, with wavelength greater than 200 nm, were effective in modifying the organic residues chemical structure. One could wonder why less energetic UV photons significantly contributed to the destruction of the nitriles units. This can be easily understood if it is noted that the number of absorbed solar UV photons is much larger in the 200-400 nm range. In particular, the UV dose absorbed by the organic residue in the 200-400 nm range was 390 times larger than the dose absorbed in the 119.5-200 nm range (see section 2.2).

4.2 Survival in the Interplanetary Medium

Once a comet enters the inner Solar System and the coma develops, a large quantity of dust is lost; it is thought that this dust constitutes a significant fraction of those IDPs that reach the Earth atmosphere (Sandford, 1987).

A major concern regarding the synthesis of biological monomers is the availability of precursor materials under primitive Earth conditions (Ruiz-Mirazo *et al.*, 2014). In this context, two possible sources are commonly considered: endogenous (terrestrial) synthesis and exogenous delivery by comets, meteorites, and IDPs. Among the endogenous processes that could have sown the early Earth with suitable precursor materials there are (i) the synthesis of organic aerosols in a neutral or more reducing atmosphere than today (Trainer, 2013; Ferus *et al.*, 2017) or (ii) hydrothermal synthesis (Lemke *et al.*, 2009). The relative amount of organic matter brought by endogenous versus exogenous sources is unknown.

It has been estimated that the delivery of organic carbon (mostly by IDPs, comets, and meteorites) on the early Earth was in the order of 10^8 kg yr⁻¹, or 10^{16} kg per 100 million years (Chyba and Sagan, 1992). Since there is about 6×10^{14} kg of organic matter in the present biosphere, it is likely that the role of extraterrestrial organics was significant (Ruiz-Mirazo *et al.*, 2014). Micrometeorites represent the dominant source of extraterrestrial material

accreted by Earth today (Taylor, *et al.* 1998). Carbonaceous matter in Antarctic micrometeorites (except UCAMMs) is up to a few % in weight (Matrajt *et al.*, 2003). UCAMMs are rare. They represent a few percent of the micrometeorites in the CONCORDIA collection. Although UCAMMs have an exceptional high carbonaceous matter content (from ~35% to ~55% in weight; Dobrica *et al.*, 2009), they may not be representative of the majority of the carbonaceous mass delivered to early Earth. Nevertheless, even for very low abundances (a few parts per billion), it is thought that nitriles could be key intermediates to form biologically relevant molecules (Kaiser and Balucani, 2001); indeed nitriles can be hydrolyzed and react via multistep synthesis ultimately to amino acids.

When not part of an active meteor stream, IDPs spend up to 10^5 - 10^6 years spiraling toward the Sun (Rietmeijer, 2009). In active meteor streams, the orbital age is shorter but not more than a few millennia (McNaught and Asher, 1999). From Fig. 11, it is evident that a significant fraction of the contained nitriles, provided that there is enough organic residue within the IDPs, has a chance to survive in the interplanetary medium for a long time. Note that, in Fig. 11, we are implicitly assuming that the nitriles present in the unexposed organic residue will be stable for very long periods. From Fig. 5, we are confident that nitriles will be present for many years in residues that

are not exposed to UV, but we cannot say whether they continue to be present after 10^2 - 10^3 years or more. Nevertheless, we already know that a few UCAMMs (Dartois *et al.*, 2013, 2018) have a carbonaceous phase with IR signatures similar to those of the PSS organic residues (Baratta *et al.*, 2015). In particular, these micrometeorites present a $\text{-C}\equiv\text{N}$ band peaking at around 2220 cm^{-1} (Dartois *et al.*, 2013, 2018), which is close to the peak position of the nitrile feature observed in the organic residues after solar UV exposure. It is thought that this high organic carbon content (ranging from $\sim 50\%$ to $\sim 85\%$ in volume; Dobrica *et al.*, 2009) and high N/C ratio micrometeorites originated in the outer Solar System beyond the nitrogen snow-line, for example in an Oort cloud comet in 100 millions to billions of years by cosmic ray irradiation (Dartois *et al.*, 2013, 2018). Hence, nitriles can survive in UCAMMs for a very long time. On the other hand, according to the results presented in this work, nitriles can be destroyed by exposure to solar UV. In the following, we try to discern the fate of the nitriles contained in an IDP, flying inward from long heliocentric distance, in the hypothesis that the same volume fraction of organic matter presents in the UCAMMs previously discussed is made of the PSS organic residue. The objective of this assumption is to take into account, in some way, that in a real IDP other minerals that may have a low absorption efficiency in the UV range are present; this increases the UV field within the IDP with a consequent decrease

of the nitriles lifetime. In the following, we assume that 50% of the IDP volume is occupied by the organic residue, while the remaining 50% is occupied by crystalline forsterite that has a very low absorption coefficient at $\lambda_{eff} = 0.342 \mu\text{m}$. To describe the absorption, reflection, and refraction of light in the composite material, we use the Maxwell-Garnet effective medium formula. In particular, the effective (average) dielectric constant of the composite medium $\epsilon_{av} = \tilde{N}_{av}^2$ (where \tilde{N} is the complex index of refraction) is given by the relation:

$$\epsilon_{av} = \epsilon_m \left[1 + \frac{3F_v \frac{\epsilon - \epsilon_m}{\epsilon + 2\epsilon_m}}{1 - F_v \frac{\epsilon - \epsilon_m}{\epsilon + 2\epsilon_m}} \right] \quad (5)$$

where ϵ_m is the dielectric constant of the matrix, ϵ and F_v are the dielectric constant and the volume fraction of the inclusions respectively. We assume that the organic material is the matrix and the inclusions are made of crystalline forsterite in equation 5. By considering the absorption coefficient $\alpha(UV)_{eff} = 5.1 \mu\text{m}^{-1}$ and the optical constants of the organic residue (Baratta *et al.* 2015), we obtain $\tilde{N}_m = 1.72 - i0.139$ at $\lambda_{eff} = 0.342 \mu\text{m}$. For crystalline forsterite, $\tilde{N}(0.342 \mu\text{m}) = 1.66 - i4 \cdot 10^{-4}$ (Pitman *et al.*, 2013). At last, by considering a volume fraction $F_v = 0.5$, we obtain from equation (5): $\tilde{N}_{av} = 1.69 - i0.069$ at $\lambda_{eff} = 0.342 \mu\text{m}$. The Maxwell-Garnett formula reported above was derived in the hypothesis of an inhomogeneous medium of a two-

component mixture composed of inclusions embedded in an otherwise homogeneous matrix. The inclusions are spherical and small compared with the wavelength. On the other hand, the UCAMMs consist of sub-micrometer sized minerals embedded in carbonaceous material, where most of the minerals are present in the form of assemblages with sizes in the range from 100 nm up to 4 μm (Dobrica *et al.*, 2012). Hence, inclusions are not always small compared with the wavelength. Thus, the average complex index of refraction obtained above should be regarded as just an approximation to take into account that the minerals present in the UCAMM, may absorb less than the organic material. At any rate, since other minerals found in the UCAMMs may absorb efficiently in the UV (i.e., iron-sulfides), the average complex index of refraction obtained by using equation 5 should be considered as a conservative estimation.

Here, we consider IDPs of spherical shape and different diameters. We suppose that during their journeys in the interplanetary medium, IDPs assume random orientation with respect to the Sun; hence, the radiation field can be considered isotropic with an intensity that is proportional to R^{-2} where R is the heliocentric distance. We consider also IDPs with size significantly larger than the wavelength, and hence, in first approximation, geometrical optics can be used to evaluate the field within the particles. In particular, we adopt a ray trace method developed by Liu *et al.* (2002). In this method, the authors

derive the internal differential absorption in a sphere of radius R_{sph} of complex index of refraction $\tilde{N} = n - ik$ at a wavelength λ , illuminated by an isotropic field. We indicate with $F(R_{sph}, \tilde{N}_{av}, r)$ the fraction of the incident radiation field transmitted within a sphere of radius R_{sph} down to a distance r from the center. For a sphere equation (4) can be modified in:

$$n_{CN} = \int_0^{R_{sph}} dN_{CN} = V_{or} N_{CN}^0 4\pi \int_0^{R_{sph}} e^{-\frac{1}{4}\Gamma \cdot t \cdot F(R_{sph}, \tilde{N}_{av}, r)} r^2 dr \quad (6)$$

where the quantity $F(R_{sph}, \tilde{N}_{av}, r)$ is computed by using the ray trace method and V_{or} is the volume fraction of the organic residue contained in the IDP. In the hypothesis that the spherical grain is uniformly irradiated, the 1/4 factor in the exponent takes into account the ratio between the geometrical cross section and the whole surface. At last, the fraction F_{CN} of survived -C≡N units, within a sphere of radius R_{sph} at 1 au after a time t , is given by:

$$F_{CN} = \frac{n_{CN}}{V_{or} N_{CN}^0 \frac{4}{3} \pi R_{sph}^3} = \frac{3}{R_{sph}^3} \int_0^{R_{sph}} e^{-\frac{1}{4}\Gamma \cdot t \cdot F(R_{sph}, \tilde{N}_{av}, r)} r^2 dr \quad (7)$$

The resulting survival curves versus the size (diameter) of the IDP are reported in Fig. 12.

The time of flight of an IDP evolving inward (spiraling) under Poynting-Robertson (P-R) drag force from a heliocentric distance R down to 1 au is given in the work of Pepin *et al.*, (2000):

$$\tau = 1.1 \cdot 10^{14} \delta \rho (R/R_0)^2 \text{ (seconds)} \quad (8)$$

where δ and ρ are the diameter and the density of the particle in c.g.s units respectively, R is the starting heliocentric distance, and R_0 is the distance that corresponds to 1 au. Especially for IDPs coming from great heliocentric distances, it should be noted that they spend most of the time far from the Sun and, hence, at much lower UV flux with respect to the flux experienced at 1 au. Baratta *et al.* (2004) computed the fluence, due to solar wind ions and solar energetic particles, suffered by IDPs under P-R drag force, in the hypothesis that the solar particles flux is proportional to R^{-2} . In particular, by using equation (8), Baratta *et al.*, (2004) found that the fluence suffered by an IDP during its P-R drag from a heliocentric distance R down to 1 au is the same as that the IDP would suffer at 1 au in a χ times shorter period, where $\chi = 2 \cdot \ln(R/R_0) \cdot (R/R_0)^{-2}$. This relation can be used to evaluate the equivalent exposure time to the solar UV photons at 1 au since the solar photon flux is also proportional to R^{-2} .

We consider now an IDP flying from a relatively large heliocentric distance $R = 7$ au, with a size similar to UCAMM-D65 ($\delta = 40 \mu m$) and a density

$\rho = 1 \text{ gcm}^{-3}$. Using equation (8), we obtain: $\tau = 6.8 \cdot 10^5$ years and $\chi = 7.9 \cdot 10^{-2}$. Finally, the resulting equivalent exposure time at 1 au is given by $\tau \cdot \chi = 5.4 \cdot 10^4$ years. From Fig. 12, it is evident that a significant fraction of nitriles can survive within the IDP. As already noted, cometary IDPs that are part of an active meteor stream may have orbital ages as short as a few millennia; in this case, the amount of nitriles that can survive within a $40 \mu\text{m}$ size IDP would be more than 50%. Nevertheless, even by considering these short exposure periods, nitriles are almost all destroyed in IDPs with sizes smaller than $6\text{-}8 \mu\text{m}$. If we consider an IDP with a diameter $\delta = 10 \mu\text{m}$, flying from 3 au, we obtain: $\tau = 3.1 \cdot 10^4$ years, $\chi = 2.44 \cdot 10^{-1}$ and $\tau \cdot \chi = 7.6 \cdot 10^3$ years. From Fig. 12, we estimate that only the 5% of the initial nitriles would survive. There could be some exceptions to the results found so far for relatively small IDPs. In particular, it has been proposed that it is possible to identify dust from comets that formed Earth-crossing dust trails (Messenger, 2002). Unlike the majority of IDPs collected in the stratosphere, which spend millennia or more in space before the Earth encounter, the dust particles may, in this case, have been in space for a few decades or less (Messenger and Walker, 2011). For such "fresh" IDPs, it is evident, from Fig. 12, that a significant fraction of nitriles could have survived even in a $10 \mu\text{m}$ particle.

It is important to stress that the results reported in Fig. 12 are strictly applicable to an IDP that contains organic matter similar to that of the organic residue studied here. It is of interest to compare the amount of the nitriles found in the PSS organic residues with the amount of nitriles found in UCAMMs. We saw that the amount of nitriles units in the PSS organic residues is 7% in weight. We do not have this information for UCAMMs, but we can estimate the relative contribution of the strength of the IR $-C\equiv N$ feature with respect to the $C=C$ feature. In particular, for the UCAMMs studied by Dartois *et al.* (2018), only three of the eight studied present a $-C\equiv N$ IR feature with intensity ratios $I(-C\equiv N)/I(C=C)$ in the range (0.055-0.027). For the PSS flight control organic residue, the intensity ratio is $I(-C\equiv N)/I(C=C)=0.18$, which is 3-7 times higher than that found in UCAMMs. At any rate, provided that the effective absorption coefficient $\alpha(UV)_{eff}$ and destruction rate Γ are the same of that found for the PSS organic residues, the survival fraction given by equation (7) for an IDP should not depend on the initial amount of nitriles units given by $V_{or}N_{CN}^0$.

5 Conclusion

In the present study, we saw that materials similar to the PSS organic residues could be present on the surface of Solar System objects, such as comets in the Oort cloud, beyond the nitrogen snow-line. When a comet enters into the inner Solar System and the coma develops, a large quantity of

dust is lost. This dust constitutes a significant fraction of the IDPs that reach the Earth atmosphere. Hence, materials similar to that we have produced in laboratory could have reached the prebiotic Earth.

According to Ruiz-Mirazo *et al.* (2014), the role of extraterrestrial organics, in providing precursor materials for the synthesis of biological monomers, was significant. Among the organics that could have had a role in providing the bricks from which life could be initiated, there are the nitriles. We know that the nitriles and amines contained in some UCAMMs can survive the heating suffered during a present-day atmospheric entry. However, modeling the interaction with the primordial terrestrial atmosphere, to determine the role of UCAMMs as carrier of pre-biotic molecules, deserves consideration (Briani *et al.*, 2013).

On the basis of the results obtained on the organic residues exposed outside the ISS, we have shown that the nitriles contained in relatively large IDPs (>20-30 μm) could have survived in the interplanetary medium, and hence they could have reached the prebiotic Earth. From Fig. 6, it is evident that the NH amine functional groups appear to be more resistant than nitriles to the solar UV photon irradiation. Hence, it is expected that a significant fraction of amines could also have reached the prebiotic Earth. However, we note that UCAMMs represent a few percent of the micrometeorites in the CONCORDIA

collection, and hence they are probably not representative of the majority of the mass delivered to early Earth. Nevertheless, even for very low abundances (a few parts per billion), nitriles are of particular importance because they are thought to be key intermediates to form amino acids, thus providing one of the basic “ingredients” for life (Kaiser and Balucani, 2001).

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Authors Disclosure Statement

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Table 1. Assignment of the IR features observed in the organic residues

Wavenumber (cm ⁻¹)	Implied functional group	Reference
3500-3300	ali-NH ₂ , aro-NH ₂ str.	(a)
3220	ali-NH ₂ , -NH- str.	(a)
2980	-CH ₃ str.	(a)
2940	-CH ₂ - str.	(a)
2200	-C≡N, -N≡C, -N=C=N-	(b, c) (c)
2150	OCN ⁻	(d)
1710	C=O str.	(a)
1650-1630, 1560	arom. C=C, he-arom. N=C ali-/arom-NH ₂ bend	(a, c)
1450	-CH ₃ bend	(a)
1415	-CH ₂ - bend	(a)
1380	C-CH ₃ , C-(CH ₃) ₂	(a)
1339	C-N arom. str.	(c)
1240	unidentified	

References: (a)= Quirico *et al.* 2008, (b)= Mutsukura and Akita, 1999,

(c)= Imanaka *et al.* 2004, (d)= Palumbo *et al.*, 2004

Table 2. Five Gaussian curves best fit parameters of the 2200 cm^{-1} feature observed in the 6f flight control and 6f flight exposed organic residues. For each Gaussian curve, the FWHM and the band area are reported. The peak positions of the curves (fixed) and the corresponding assignment (Mutsukura and Akita, 1999) are also reported in the table.

peak position (cm^{-1})	assignment	flight control area (cm^{-1})	flight control FWHM (cm^{-1})	flight exposed area (cm^{-1})	flight exposed FWHM (cm^{-1})
2105	-N \equiv C	0.0179 ± 0.0006	32.6 ± 0.6	0.0075 ± 0.0002	44.5 ± 1
2145	-N \equiv C	0.159 ± 0.003	43.7 ± 0.6	0.0228 ± 0.0006	38.5 ± 0.9
2190	-N \equiv C	0.847 ± 0.006	56.1 ± 0.4	0.111 ± 0.001	46.0 ± 0.6
2220	-C \equiv N	0.193 ± 0.004	36.9 ± 0.6	0.1165 ± 0.001	32.9 ± 0.3
2252	-C \equiv N	0.0483 ± 0.001	24.7 ± 0.3	0.0037 ± 0.0003	19.9 ± 0.6

Figure Legends

FIG.1 (a) Schematic view of the experimental apparatus used to produce the organic residues. (b) Scheme showing the experimental breakdown of samples between the ISS and DLR Cologne, and the environmental conditions to which they have been exposed.

FIG.2 Spectra after 200 keV He^+ irradiation of the ice mixtures for the three considered thicknesses.

FIG.3 Spectra of the organic residues obtained after irradiation at a dose of 110 eV/16u at 17 K followed by a slow warm up to room temperature and sublimation of the volatile species. Each displayed spectrum is the average of the 10 different samples realized for each thickness.

FIG.4 Comparison between the spectra of the ground and in-flight dark controls.

FIG.5 Temporal evolution of the 2200 cm^{-1} ($-\text{C}\equiv\text{N}$) band area of the laboratory and in-flight dark control unexposed samples. The two dashed vertical lines correspond to the start and the end of exposure to the solar UV photons experienced in orbit by the exposed samples. The best fit to the data was obtained

by using the equation: $y=sc*(y_0+A_1*exp(-t/t_1)+A_2*exp(-t/t_2))$. Once the best fit solution was found for the thickest (6f) samples with sc set to 1, the other two thicknesses were fitted by adjusting only the sc scaling factor.

FIG.6 Comparison between the in-flight dark controls (dotted lines) and the flight exposed samples before (solid lines) and after (dashed lines) the correction for the contamination spectrum.

FIG.7 Comparison of the normalized spectra, for the three different thicknesses, between the in-flight dark controls (upper curves) and the flight exposed (lower curves) samples in the three spectral regions of interest.

FIG.8 Fives Gaussian with fixed peak position (thin grey lines) best fit (thick grey lines) of the thickest 6f in-flight dark control (upper curves) and flight exposed (lower curves) samples. The 95% confidence interval of the best fit is indicated with black dashed lines. The best fit output parameters and their uncertainties are reported in Table 2.

FIG.9 Comparison of the 2200 cm^{-1} CN feature between the in-flight dark control (dotted lines) and flight exposed (solid lines) samples at different depths (see the text).

FIG.10 CN column density of the in-flight dark control (open squares) multiplied by a factor 1/2 and flight exposed (closed squares) samples versus the thickness. The solid curve is the best fit to the flight exposed column density obtained by using equation 4. The fit is for 2111 h of solar irradiation at 1 au. The dashed straight line is the linear best fit to the in-flight dark control column density values.

FIG.11 Residual CN column density contained in a 50 nm thick slab at different depths in the organic residue and different exposure time at 1 au. The horizontal dashed line is the column density value for the unexposed organic residue. The solid line correspond to the equivalent to perpendicular solar exposure suffered by PSS samples on the ISS. The full circles are the interpolated flight exposed samples data for the three different thicknesses.

FIG.12 Residual fraction of the CN units contained in spherical IDPs of different radius and different exposure time at 1 au obtained in the hypothesis that the organic residue is diluted by a factor of 50 % in volume and the remaining fraction is made of crystalline forsterite.