Lunar Dust and Lunar Simulant Activation and W.T. Wallace¹, D.K. Hammond², and A.S. Jeevarajan³

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Introduction

Prior to returning to the moon, understanding the effects of lunar dust on both human physiology and mechanical equipment is a pressing concern, as problems related to lunar dust during the Apollo missions have been well documented (J.R. Gaier, The Effects of Lunar Dust on EVA Systems During the Apollo Missions. 2005, NASA-Glenn Research Center. p. 65). While efforts were made to remove the dust before reentering the lunar module, via brushing of the suits or vacuuming, a significant amount of dust was returned to the spacecraft, causing various problems. For instance, astronaut Harrison Schmitt complained of "hay fever" effects caused by the dust, and the abrasive nature of the material was found to cause problems with various joints and seals of the spacecraft and suits. It is clear that, in order to avoid potential health and performance problems while on the lunar surface, the reactive properties of lunar dust must be auenched.

It is likely that soil on the lunar surface is in an "activated" form, i.e. capable of producing oxygen-based radicals in a humidified air environment, due to constant exposure to meteorite impacts, UV radiation, and elements of the solar wind An activated silica surface serves as a good example. An oxygen-based radical species arises from the breaking of Si-O Si bonds. This system is comparable to that expected for the lunar dust system due to the large amounts of agglutinic glass and silicate vapor deposits present in lunar soil. Unfortunately, exposure to the Earth's atmosphere has passivated the active species on lunar dust, leading to efforts to reactivate the dust in order to understand the true effects that will be experienced by astronauts and equipment on the moon.

Electron spin resonance (ESR) spectroscopy is commonly used for the study of radical species, and has been used previously to study silicon- and oxygen-based radicals, as well as the hydroxyl radicals produced by these species in solution (V. Vallyathan, et al., Am. Rev. Respir. Dis. 138 (1988) 1213-1219). The size and cost of these instruments makes them unattractive for the monitoring of lunar dust activity. A more suitable technique is based on the change in fluorescence of a molecule upon reaction with a hydroxyl radical (or other radical species). Fluorescence instruments are much less costly and bulky than ESR spectrometers, and small fluorescence sensors for space missions have already been developed (F. Gao, et al., J. Biomed. Opt. 10 (2005) 054005).

For the current fluorescence studies, the terephthalate molecule has been chosen for monitoring the production of hydroxyl radicals in solution. As shown in Scheme 1, the reaction between the non-fluorescent terephthalate molecule and a hydroxyl radical produces the highly-fluorescent 2-hydroxyterephthalate molecule. This reaction has been shown to





Figure 1: Fluorescen wavelength - 324 nr ce of Synthetic 2-Hydroxyterephthalate. Excitation

be an excellent choice for monitoring hydroxyl radical prodcution (N. Soh, Anal. Bioanal. Chem. 386 (2006) 532-543). The major advantage of this reaction is the fact that attack of the hydroxyl radical at any of the open ring positions results in the same product, which provides ease in interpreting the fluorescence spectra.

The results presented here focus on our studies of the abilities of different methods to activate lunar simulant, lunar dust, and quartz, and the ability of fluorescence to monitor this activity using the terephthalate molecule. Additionally, we show initial results on the in situ toxicity of lunar simulant on human lung and skin cells.

Materials and Methods

• Test materials: Crystalline silica (Min-U-Sil 15) was provided by U.S. Silica, Mill Creek, OK. This material has a mean diameter of 15 µm. Lunar simulant (JSC-1A-vf) was obtained from Dr. James Carter at the University of Texas at Dallas This simulant was designed to be similar to low-titanium, mature lunar mare regolith with 90% of the particles less than 13 µm. The lunar dust used for this study was an Apollo 16 soil (62241) provided by Dr. Larry Taylor at the University of Tennessee-Knoxville. This particular sample is a mature highland soil with a size distribution between 3 µm - 450 µm.

· Preparation of terephthalate solutions: Terephthalate solutions were prepared by dissolving disodium terephthalate (99%, Alfa Aesar) in phosphate-buffered saline (PBS, Sigma Aldrich) in order to produce a final terephthalate concentration of 10 mM.

• Grinding procedure: 70 mg of quartz, lunar simulant, or lunar dust were placed in a mortar and ground with a pestle for 10 minutes, stopping every 2 minutes to scrape the sides in order to ensure even grinding. It is important to start with the same amount of material for every test, as grinding different amounts will lead to different activities

 Fluorescence testing procedure: Ground and unground material were added to 15 mL centrifuge tubes containing 2.5 mL of 10 mM terephthalate dissolved in PBS. The mixtures were allowed to interact for 30 minutes before being filtered using 0.22 µm syringe filters. Two mL of the filtered solution was added to a quartz fluorescence cuvette, and the fluorescene spectrum was obtained using a Perkin-Elmer LS-50B spectrometer

• Calibration procedure: The spectra and calibration curve shown in Figure 1 were produced using 2-hydroxyterephthalate synthesized using a method from the literature (L. Field and P.R. Engelhardt, J. Org. Chem. 35 (1970) 3647-3655). The material was then purified using recrystallization until the fluorescence spectra were maximized. Mass spectrometry also confirmed that the only species present following recrystallization was 2-hydroxyterephthalate. A 1 mM solution was prepared by dissolving the product in PBS. Further dilutions were made from this stock solution. For concentrations above 10 nM, the fluorescence saturated the detector. Dilutions below this threshold varied linearly with concentration, as shown in the figure. The excitation wavelength was 324 nm.



Figure 2: Activity Comparison of Freshly Ground and Unground Apollo 16 Soil (62241).

Results

The addition of a small amount of unactivated materials to a 10 mM fluorescence (not shown). However, it can be seen that the general or in the order lunar dust > lunar simulant > quartz. Grinding of the mater instance, grinding of Apollo 16 soil (62241) produces the spectrum in F unground material also shown, and a comparison to the calibration curv lunar dust produces a large amount of hydroxyl radicals. The amount of lunar dust, lunar simulant, and quartz are compared. In Figure 3, it ca conditions, lunar soil produces 2-3 times more hydroxyl radicals than lu

UV Exposure Time (min)

Figure 4: UV Activation of Unground Lunar Simulant.



Figure 5: Effect of Unactivated Lunar Simulant on Lung Cells. (courtesy M.J. Cunningham)



Figure 6: Effect of Unactivated Lunar Simulant on Skin Cells. (courtesy M.J. Cunningham)

Future work

Future work will focus on a variety of tests involving lunar dust acti and cellular toxicity

- Proton beam and further LIV exposure experiments will be conduexposures experienced by dust on the lunar surface.
- . The deactivation kinetics of activated lunar dust will be monitored the development of dust mitigation programs.
- Dissolution studies will be carried out to determine if any potential and to determine their concentrations.
- · Further cellular toxicity experiments will be conducted using differ of lunar dust activation in vitro and to determine if different cells are

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Figure 3: Activit

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e solution produces very little xyl radical production decreases wn to have a large effect. For th the spectrum produced by the to see that simply grinding the even more striking when the t, under the same grinding and 10 times more than quartz.

g changes the particle size ods would be used to reactivate mic the ultraviolet radiation experiments have been carried ant under moderate vacuum to in Figure 4, after performing the be seen that exposure to UV ease in reactivity.

numan body is of major interest. en performed. Figures 5 and 6 an lung (NHBE) and skin (HEK) ntrations of unground lunar ry large concentrations of dust bility. However, cell death is not o lunar dust. It is possible that use the cells to produce other eroxide, or to damage the cell ust may also cause damage or er studies are needed in order to t on cells.

nods of activating quartz, lunar ng the levels of activation using a duction of hydroxyl radicals in cinogen, was shown to be much oxyl radicals than ground lunar tionally, the same assay has simulant to a high-powered diation obtained by soil on the tion. Concurrent studies on the tems has shown that extremely required in order to cause cell evidence of the need for further urning to the lunar surface

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radley, Ph.D., M.J. Cunningham,

determine the full effects Conclusions This work has demor simulant and lunar soil fluorescence assay base

solution. Ground quartz, less active for the produc simulant and ground luna shown that exposing ung ultraviolet source (to min lunar surface) can also le toxicity of lunar simulant high doses of unground death. Each of these res studies on these materia