
[EE] Evening Poster | B (Biogeosciences) | B-AO Astrobiology & the Origin of Life

[B-AO01]Astrobiology

convener:Hikaru Yabuta(Hiroshima University, Department of Earth and Planetary Systems Science), Seiji Sugita(Department of Earth and Planetary Science, Graduate School of Science, The University of Tokyo), Misato Fukagawa(名古屋大学, 共同), Fujishima Kosuke(Tokyo Institute of Technology, Earth-Life Science Institute)

Tue. May 22, 2018 5:15 PM - 6:30 PM Poster Hall (International Exhibition Hall7, Makuhari Messe)

Twenty years have passed since when the field of Astrobiology, which aims to unveil the origins, evolution, and habitability of life by integrating multidisciplinary fields, was established. Origins of Life are currently being re-conceptualized via expansion of prebiotic chemistry to systems chemistry and chemical space. Besides their relationship to life's building blocks, it is expected to demonstrate the significant roles of organic molecules in the history of planetary formation. The linkages among the variations in chemical compositions of deep-sea hydrothermal environments, geological settings, and ecological systems were systematically investigated. Cassini, which accomplished in the long-term explorations of the planets bearing liquid, had "Grand Finale" this year. Discoveries of extrasolar planets have been dramatically increased to date.

Originally, Astrobiology does not need a specific science category. We therefore aim to make this session so that Earth and Planetary scientists from all the categories join for discussing 'where we came from and where we are going' and for making novel integrated researches.

For the next stage of Astrobiology, presentations on the instrument development in space explorations, comparative studies of solar system and exoplanets, etc, will be very much welcome.

[BAO01-P13]Mutation analysis of the *rpoB* gene in the radiation-resistant bacterium, *Deinococcus radiodurans* R1 exposed to space

*Daisuke Fujiwara¹, Yuko Kawaguchi¹, Yuka Togashi¹, Iori Kinoshita¹, Jun Yatabe¹, Issay Narumi², Hirofumi Hashimoto³, Shin-ichi Yokobori¹, Akihiko Yamagishi¹ (1.Tokyo University of Pharmacy and Life Sciences, 2.Toyo University, 3.ISAS/JAXA)

Keywords:DNA damage, *Deinococcus* spp., International Space Station, Mutation, *rpoB* gene, Space

To investigate the microbial viability and their DNA damage, the radiation-resistant bacteria *Deinococcus* spp. have been exposed at the Exposure Facility of the International Space Station (ISS) in Tanpopo mission since May 2015 [1,2]. The Exposure Panels (EPs) harboring dried-deinococcal cells returned to the ground on October 2016 after about one-year exposure. We analyze the survival rate and DNA damage of dried deinococcal cells using pulsed-field gel electrophoresis, quantitative-PCR and mutation assay. The antibiotic rifampicin binds the RNA polymerase β -subunit, which is encoded by the *rpoB* gene, and inhibits the initial step of transcription. Certain mutations in the *rpoB* gene confer rifampicin resistance [3]. Based on these characteristics of the *rpoB* gene, we determined mutant frequency and the mutation spectrum in the *D. radiodurans* *rpoB* gene exposure to space. From these mutation data, we estimated major DNA damage induced by the space environment.

D. radiodurans R1 cell-suspension was dropped in the wells of aluminum plates and was dried under vacuum (vacuum-dried). The dried cells were exposed to space, stored in ISS cabin or in the ground laboratory. After exposure experiment, the cells recovered from each well were used to inoculate 10 ml of mTGE medium and cultured until OD_{590 nm} reached between 0.7 and 3.0. The cell suspension was plated

on mTGE agar containing 50 µg/ml rifampicin to determine the number of rifampicin resistant cells (Rif^R), and on mTGE agar without rifampicin to determine the total number of viable cells.

We also determined the sequences of the *rpoB* gene extracted from Rif^R. The rifampicin resistant mutation frequencies of the space exposed *D. radiodurans* R1 cells and those of the ground control were comparable (Fig. 1). The result suggested that the effect of UV for dried-cells exposure to space was less. Further, we will report and discuss the rifampicin-resistant spectra in the *rpoB* gene in rifampicin-resistant cells exposure to space, stored in ISS or in ground laboratory.

- [1] Yamagishi, A. et al., (2007) *Bio. Sci. Space* 21: 67-75.
- [2] Kawaguchi et al., 2016, *Astrobiology* 16 : 363-376.
- [3] Campbell, E. A. et al., (2001) *Cell* 104: 901-912