Enhanced Microbial Survivability in Subzero Brines

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Abstract

It is well known that dissolved salts can significantly lower the freezing point of water and thus extend habitability to subzero conditions. However, most investigations thus far have focused on sodium chloride as a solute. In this study, we report on the survivability of the bacterial strain Planococcus halocryophilus in sodium, magnesium, and calcium chloride or perchlorate solutions at temperatures ranging from +25°C to −30°C. In addition, we determined the survival rates of P. halocryophilus when subjected to multiple freeze/thaw cycles. We found that cells suspended in chloride-containing samples have markedly increased survival rates compared with those in perchlorate-containing samples. In both cases, the survival rates increase with lower temperatures; however, this effect is more pronounced in chloride-containing samples. Furthermore, we found that higher salt concentrations increase survival rates when cells are subjected to freeze/thaw cycles. Our findings have important implications not only for the habitability of cold environments on Earth but also for extraterrestrial environments such as that of Mars, where cold brines might exist in the subsurface and perhaps even appear temporarily at the surface such as at recurring slope lineae. Key Words: Brines—Halophile—Mars—Perchlorate—Subzero—Survival. Astrobiology 18, xxx–xxx.

1. Introduction

LIFE AS WE KNOW it requires liquid water as the principal solvent for its biochemistry, but most planetary surfaces in our Solar System never reach temperatures above the freezing point of pure water, rendering these localities as likely uninhabitable compared with the benign climate conditions on Earth. However, the presence of salts can lead to a substantial freezing point depression down to the eutectic temperature of a given salt/water mixture (e.g., −50°C for a 31 wt% CaCl2 solution) and, thus, greatly expand the temperature range for potential habitats (Möhlmann and Thomsen, 2011). Hence, the question arises as to whether microorganisms can thrive or at least survive in such subzero brines.

On Earth, microbial organisms such as yeast can tolerate water activities (aw) down to 0.61 (Rummel et al., 2014). However, the lowest salt-induced water activity that halophilic microorganisms can tolerate is that of a saturated NaCl solution (aw = 0.75), while other salts (e.g., those containing Ca2+ and Mg2+ ions) are more inhibitory to cell metabolism (Rummel et al., 2014). Furthermore, it has been reported that certain cyanobacterial species embedded in hygroscopic sodium chloride (NaCl) deposits found in the hyperarid soils of the Atacama Desert are able to utilize water condensed from the atmosphere via deliquescence (Davila et al., 2008; Davila and Schulze-Makuch, 2016).

In addition, many halophilic microorganisms can also be psychrophilic or psychrotolerant (Gounot, 1986; Hoover and Pikuta, 2010). To date, the lowest reported temperature for microbial growth is −18°C for yeast on frozen surfaces (Collins and Buick, 1989). Metabolic ammonia oxidation has been detected down to −32°C (Miteva et al., 2007), and finally, there are indications for photosynthetic activity of lichens at −40°C (de Vera et al., 2014).

It has been argued that low-temperature and high salt tolerances are closely linked, given that at subzero temperatures, water ice forms, which increases the solute concentration of the remaining liquid water (Bakermans, 2012). Moreover, chaotropic agents such as magnesium chloride (MgCl2), that is, substances that destroy the bulk water structure and therefore reduce hydrophobic interactions (Gerba, 1984) at moderate concentrations, can decrease the minimal temperature at which cell division can occur for certain microorganisms and increase their growth rates at...
low temperatures, presumably by increasing the macromolecular flexibility (Chin et al., 2010). Furthermore, some microorganisms shift their salinity optimum for growth to higher salt concentrations if exposed to lower temperatures (Gilichinsky et al., 2003).

Organisms have evolved several adaptations for thriving and/or surviving in cold saline environments. These include production of antifreeze or ice-binding proteins, cryoprotectants, or extracellular polymeric substances (Jia et al., 1996; Gilbert et al., 2005; Kuhlmann et al., 2011), an increase of fatty acids that branch and maintain membrane fluidity (Dénich et al., 2003), a higher antioxidant defense against reactive oxygen species (Chattopadhyay et al., 2011), the expression of isozymes adapted to low temperatures and high salinities (Maki et al., 2006), or the exclusion of inhibitory ions by accumulating intercellular compatible solutes (Csonka, 1989).

Most of the studies dealing with brines at subzero temperatures have focused on NaCl as a solute, the most common salt found in saline environments on Earth. However, certain environments on Earth are dominated by high concentrations of other salts such as calcium chloride (CaCl₂) in Don Juan Pond, Antarctica (Cameron et al., 1972; Dickson et al., 2013), or sodium and magnesium sulfates in Spotted Lake, Canada (Ponterfract et al., 2017). Furthermore, martian soils are known to contain various chloride (Cl⁻) and perchlorate (ClO₄⁻) salts (Hecht et al., 2009; Kounaves et al., 2010), emphasizing the importance of research in the field of non-NaCl briny habitats at subzero temperatures.

In this study, we used the halo- and cryotolerant bacterial strain Planococcus halocryophilus Or1 (DSM 24743T), which was isolated from the active layer of permafrost soil in the Canadian High Arctic (Mykytczuk et al., 2012). This organism grows at temperatures between −15°C and +37°C and under NaCl concentrations of up to 19 wt% at which metabolic activity has been detected at temperatures down to −25°C (Mykytczuk et al., 2013).

This bacterial strain shows many cold and osmotic stress responses such as the expression of cold-adapted proteins, the expression of various osmolyte transporters, a high lipid turnover rate, a high resource efficiency at cold temperatures with an accumulation of carbohydrates as a energy resource (Mykytczuk et al., 2013), and complex changes in protein abundances (Raymond-Bouchard et al., 2017). Furthermore, under cold growth conditions, P. halocryophilus develops a nodular sheet-like crust around the cells (Ronholm et al., 2015; Mykytczuk et al., 2016). The above-described ability of P. halocryophilus to cope with low temperatures and high salt concentrations makes it an ideal organism for studying whether, and how well, terrestrial life might be able to survive or even thrive in martian environments. In particular, we have investigated how well P. halocryophilus can survive repeated freezing/thawing cycles and in subzero chloride and perchlorate brines, since such conditions may be temporarily present on Mars (Martínez and Renno, 2013).

2. Materials and Methods

2.1. Strain and culture conditions

We used the bacterial strain Planococcus halocryophilus Or1 (DSM 24743T), which was obtained from the DSMZ (Leibniz Institute DSMZ—German Collection of Microorganisms and Cell Cultures). P. halocryophilus was grown in DSMZ growth medium #92 containing additional 10 wt% NaCl. Its growth curve at 25°C was determined via colony forming units (CFUs), and cell suspensions used for inoculating the experiments were either retrieved after 4 days (sample type ST 1) or 7 days (sample type ST 2) of growth (Fig. 1).
**Table 1. Eutectic Concentrations and Temperatures, Ionic Strength, and Water Activities at 25°C for Salt Solutions Used in This Study**

| Eutectic concentration | Ionic strength | Water activity at 25°C | Eutectic temperature |
|------------------------|----------------|-----------------------|----------------------|
| NaCl                   | 23.3 a         | 5.20                  | −22                  | 251 a                 |
| MgCl₂                  | 21 a           | 2.79                  | 0.80                 | −33.5                  |
| CaCl₂                  | 30.2 a         | 3.90                  | 11.70                | 0.52                   |
| NaClO₄                 | 52.6 a         | 9.06                  | 0.68                 | 0.56                   |
| Mg(ClO₄)₂              | 44 a           | 5.32                  | 10.56                | 0.56                   |
| Ca(ClO₄)₂              | 50.1 a         | 4.20                  | 12.60                | 0.52                   |

aMöhlmann and Thomsen (2011).
Hennings et al. (2013).
Stillman and Grimm (2011).
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2.2. Experiments in eutectic salt solutions

In all experiments, 2 mL of the cell suspension (prepared as described in Section 2.1) was mixed with 8 mL of a salt solution resulting in 10 mL of sample solution with a eutectic salt concentration. The eutectic compositions of the investigated salts are listed in Table 1, together with the ionic strength, the water activity at 25°C calculated from the Pitzer equation (Pitzer, 1991) with Pitzer parameters taken from the work of Toner et al. (2015), and the eutectic temperature. All samples were prepared and analyzed as biological duplicates.

Before mixing cell suspensions and salt solutions, the suspensions were cooled to 4°C and the salt solutions to the respective experimental temperature. In addition, for testing whether ClO₄⁻ preconditioning of the cells has a positive effect on their survival in ClO₄⁻ containing samples, cell suspensions with either up to 10 wt% NaClO₄ or 5 wt% NaClO₄ + 10 wt% NaCl were prepared and incubated for 7 days at 25°C.

2.3. Cell number quantification

The concentration of viable cells in the samples was determined after specific time intervals via CFU counts, and, where necessary, samples were diluted in phosphate-buffered saline (PBS) containing 21 wt% NaCl or MgCl₂ to avoid osmotic bursting of cells. Highest values of CFU mL⁻¹ were achieved when dilution was done with NaCl-enriched PBS for samples containing NaCl, MgCl₂, NaClO₄, or magnesium perchlorate (Mg(ClO₄)₂), and MgCl₂-enriched PBS for samples containing CaCl₂ or calcium perchlorate (Ca(ClO₄)₂). Because cell death occurred at higher temperatures during plating, especially in CaCl₂ containing samples, plating for all experiments described in this study was carried out rapidly at cold temperatures. The NaCl/MgCl₂-enriched PBS was precooled to −15°C/−30°C and agar plates to 4°C.

2.4. Freeze/thaw cycle experiments

For investigating the effect of dissolved salts on cell survival when subjected to multiple freeze/thaw cycles, we incubated P. halocryophilus at 25°C for 1 week in six individual vials. Three of them contained 10 mL of DMSZ growth medium #92 (with no additional NaCl), while the other three samples contained additionally 10 wt% NaCl. After incubation, all samples where repeatedly frozen at −50°C, stored at this temperature between 1 and 3 days, and thawed at room temperature until the unfrozen sample reached 20°C, which took ~ 2 h. After taking an aliquot from each sample for CFU determination, the samples were frozen again. These freeze/thaw cycles were repeated up to 70 times, and the survival was tested intermittently. The results for samples with the same growth media composition were averaged, and the standard deviation was calculated.

3. Results

3.1. Microbial survival rates in chloride brines

The survival rates of P. halocryophilus in eutectic Cl⁻ samples were significantly increased when the samples were kept at lower temperatures (Fig. 2). For example, if P. halocryophilus was left in NaCl containing samples at room temperature, all cells died within 2 weeks, while their survival was substantially increased at 4°C, and nearly no CFU reduction occurred at −15°C. Two samples of ST 2 were investigated for the NaCl system to confirm reproducibility. Samples of ST 2 had slightly higher starting cell numbers in all cases studied. However, survival rates of ST 1 and ST 2 samples were similar, although the curve for the NaCl ST 1 sample at 4°C had a steeper slope during the first 40 days but approached the slope of the ST 2 curves afterward.

The cell survival results for the MgCl₂ containing samples were very similar to those of NaCl, but at −15°C and −30°C there appears to be a slow reduction of surviving cells. The survival rates of P. halocryophilus in CaCl₂ containing samples at 25°C and 4°C were significantly lower than those containing NaCl or MgCl₂. In contrast, survival rates at subzero temperatures were comparable to the MgCl₂ system, that is, cells were dying slower at these lower temperatures.

3.2. Microbial survival rates in perchlorate brines

The survival rates of P. halocryophilus in eutectic ClO₄⁻ samples (Fig. 3) were orders of magnitude lower than in Cl⁻ samples (Fig. 2). Although survivability at lower temperatures in NaClO₄ samples increased, the survival rate was generally so low that even at −30°C, few cells survived for only 1 day (Fig. 3A, B). For Mg(ClO₄)₂ and Ca(ClO₄)₂ containing samples, survival was even lower, where CFU detection was only possible for samples stored at −30°C, and none was detected for samples kept at higher temperatures.

We increased the NaClO₄ concentration in the growth media to determine whether ClO₄⁻ preconditioning of P. halocryophilus could enhance survival in eutectic ClO₄⁻ samples. It was found that P. halocryophilus could grow in the presence of up to at least 10 wt% NaClO₄ (with no additional NaCl in the growth medium) or up to 10 wt% NaCl + 5 wt% NaClO₄. However, cell growth under these conditions was markedly slower than in ClO₄⁻ free medium. Thus, for the preconditioning experiments we used cells preconditioned with 8 wt% NaClO₄ or with 10 wt% NaCl + 3 wt% NaClO₄ (Fig. 3B). Nevertheless, in these cases, cells grew slower than in the experiments with 10 wt% NaCl in the growth media, which resulted in a lower starting cell number.

Due to the slower growth rates in ClO₄⁻ containing media, the cells should still be in the exponential growth phase after 7 days of incubation. We found that changing the preconditioning salt from NaCl to NaClO₄ did not increase the survivability in ClO₄⁻ containing samples. However, increasing the total salt concentration by adding 3 wt% NaClO₄ on 10 wt% NaCl resulted in a slight increase in survival. Cells in these
samples doubled their maximum survival time from ~1 day in samples containing either 8 wt% NaClO₄ or 10 wt% NaCl to 2 days in samples containing 3 wt% NaClO₄ + 10 wt% NaCl.

3.3. Arrhenius plot

For a better comparison of the temperature dependences of cell survival in different Cl⁻ and ClO₄⁻ containing samples, the data were plotted as an Arrhenius-type graph, with the slopes of the survival rate-fitted lines for all Cl⁻ and NaClO₄ containing samples (values for same salt/temperature combinations were averaged) plotted logarithmically against the temperature of the sample (Fig. 4A). As the slope (S) of these curves is the crucial parameter for evaluating the extent to which survival is increased with lowering temperature, the slope values for each curve were plotted as well (Fig. 4B).

The slopes for the Cl⁻ containing samples, especially for MgCl₂ and CaCl₂, flatten below death rate constants of about 0.1 day⁻¹. However, it has to be kept in mind that the death rates are on a logarithmic scale, and therefore, the
flattening might only be the result of approaching a non-lethal state, that is, a death rate of zero. Therefore, only the steeper slopes of the curves toward higher temperatures were compared, as given in Figure 4B.

3.4. Microbial survival rates during freeze/thaw cycles

*P. halocryophilus* survived repeated freeze/thaw cycles more readily if the growth medium contained additional NaCl. Without NaCl, the CFU reduction is 20% per freeze/thaw cycle, whereas an addition of 10 wt% NaCl lowered the death rate to 7% per freeze/thaw cycle (Fig. 5). Cells in the salt-free samples survived up to 70 freeze/thaw cycles, while extrapolation of the death rate curve for the samples containing 10 wt% NaCl reveals that cells in these samples could survive up to ~200 freeze/thaw cycles.

4. Discussion

We have shown that survival of *P. halocryophilus* is significantly lower in eutectic ClO₄⁻ samples than in Cl⁻ containing samples at all investigated temperatures, although ionic strength
and water activities at 25°C are similar, for example, for CaCl₂ and NaClO₄ samples (Table 1 and Fig. 4A). Moreover, the water activity should not change markedly when lowering the temperature since it has been shown to remain reasonably constant at subzero temperatures for solutions containing Cl⁻ (Fontan and Chirife, 1981) and ClO₄⁻ (Toner and Catling, 2016). Furthermore, the oxidizing ability of ClO₄⁻ is negligible in solutions at these low temperatures (Brown and Gu, 2006). Thus, other ion-specific properties must be responsible for the differences in the inhibitory effects of the ClO₄⁻ and Cl⁻ containing samples.

In addition, we demonstrated that the survival of P. halocryophilus cells in eutectic Cl⁻ and ClO₄⁻ samples increases systematically with decreasing temperatures. The Arrhenius plot (Fig. 4) indicates that this correlation is more significant in Cl⁻ containing samples. The slope for the CaCl₂ containing samples (0.225°C⁻¹) is more than 2.5-fold steeper than for the NaClO₄ containing samples (0.079°C⁻¹), which means that survivability in the CaCl₂ samples is increased by lowering temperature to a significantly higher extent than in the NaClO₄ samples. The slopes for MgCl₂ (0.152°C⁻¹) and NaCl (0.135°C⁻¹) containing samples lie between those of NaClO₄ and CaCl₂.

The slow decrease of the death rate constant in the NaClO₄ containing samples with decreasing temperature is caused by the normal temperature dependence of all

![FIG. 4. (A) Arrhenius-type plot for all Cl⁻ samples and NaClO₄, including slopes (S) for linear parts of the curves and molar concentrations (c), water activities (aw), and ionic strengths (I) for all samples. (B) Slopes (S) of the steeper curve parts plotted as bar charts.](image)
chemical reactions (including cell damaging reactions) described by the Arrhenius equation. The steeper slopes for the Cl\textsuperscript{−} samples indicate an additional effect on the decrease of death rates with lowering the temperature.

We propose that the main reason for this difference in the temperature dependence of the cell survival in Cl\textsuperscript{−} and ClO\textsubscript{4}\textsuperscript{−} containing samples is the increase of size and stability of hydration spheres around the ions in the Cl\textsuperscript{−} brines at lower temperatures. Previous studies have shown that with decreasing temperatures the hydration number around cations such as Ca\textsuperscript{2+} increases (Zavitsas, 2005) and that the first hydration sphere around Na\textsuperscript{+} in NaCl solutions becomes more rigid (Gallo et al., 2011). Furthermore, X-ray and neutron diffraction studies have shown that a decrease in temperature results in the first hydration shell of Cl\textsuperscript{−} ions becoming gradually more structured and a second hydration sphere forming (Yamaguchi et al., 1995). Data from the method of integral equations reveal a strengthening of the hydrogen bonding between Cl\textsuperscript{−} and water molecules in the first hydration shell at lower temperatures (Oparin et al., 2002).

These results demonstrate that lowering the temperature in Cl\textsuperscript{−} containing samples increases the stability and size of hydration spheres around the dissolved ions, known to reduce the permeability of ions through cell membranes (Degrève et al., 1996; Jahnen-Dechent and Ketteler, 2012). Hence, we conclude that a reduced ion permeability caused by larger and more stable hydration spheres minimizes the toxicity of the extracellular high ion concentration. Therefore, cell survivability in low-temperature Cl\textsuperscript{−} brines is increased over the extent of the normal Arrhenius-like temperature dependence.

In contrast to cations and Cl\textsuperscript{−} ions, ClO\textsubscript{4}\textsuperscript{−} ions do not tend to form stable hydration shells (Neilson et al., 1985; Lindqvist-Reis et al., 1998). The reason for the small size and low stability of hydration shells around ClO\textsubscript{4}\textsuperscript{−} is its large ionic radius and its low electrical charge with an even distribution over the entire anion (Brown and Gu, 2006) resulting in weak hydrogen bonds and one of the lowest hydration energies of common inorganic anions (Moyer and Bonnesen, 1979). The low tendency of ClO\textsubscript{4}\textsuperscript{−} ions to form stable hydration spheres at any temperature presumably correlates with a constant cell membrane permeability and is a reasonable explanation for the observed low survival rate increase in ClO\textsubscript{4}\textsuperscript{−} containing samples with lower temperatures.

Finally, the higher membrane permeability at all investigated temperatures explains the general low survivability of cells in ClO\textsubscript{4}\textsuperscript{−} versus Cl\textsuperscript{−} containing samples. However, several other structural factors may play a significant role as well, for example, the formation of chloro-complexes in CaCl\textsubscript{2}.

FIG. 5. Survivability of *P. halocryophilus* during freeze/thaw cycles. Cells were incubated for 7 days at 25°C in growth medium containing either no additional salt (black circles) or 10 wt% NaCl (gray triangles) before subjecting them to freeze/thaw cycles. CFUs obtained from biological triplicates.
containing samples (Phutela and Pitzer, 1983; Wang et al., 2016), ion pair formations (Fleissner et al., 1993; Smirnov et al., 1998), molecular mimicry (Cianchetta et al., 2010), or the reported formation of a crust around \( P. \) halocryophilus cells at low temperatures, consisting of peptidoglycan, choline, and calcium carbonate (Mykytczuk et al., 2016) that might provide protection against Cl\(^-\) but not ClO\(_4^-\).

The freeze/thaw experiments have shown that the survivability of cells during freezing and thawing processes increases when NaCl is present. Studies have argued that the formation of large water crystals during freezing might be destructive to cell membranes and might even grow larger during thawing due to migratory recrystallization (Mazur, 1960). Greater amounts of large water crystals should only form in the salt-free samples, because in the salt-rich samples, pure water crystals are formed during freezing only until the solution under the ice layer reaches the eutectic composition. After that point, eutectic freezing results in very small water ice and salt hydrate particles, which potentially could be physically less harmful to the cells.

Another lethal effect during the freezing process might be the osmotic shock resulting from the increasing solute concentration in the remaining liquid solution (Harrison, 1956). It is reasonable to assume that the decreased water activity, as a result of the enhanced solute concentration in the growth media during the freezing process, is less harmful to bacteria that were already preconditioned with 10 wt% NaCl during incubation. Furthermore, studies have shown that a heat or cold shock treatment of \( D. \) radiodurans cells increases their survivability against freeze/thaw cycles (Airo et al., 2004), hence an exposure to higher salt concentrations may result in a similar stress response in \( P. \) halocryophilus and a higher tolerance against freeze/thaw cycling. The beneficial effect of NaCl during the freeze/thaw process has been described in previous studies (e.g., Calcott and Rose, 1982).

In contrast, other studies have shown the opposite trend, that the presence of NaCl decreases the percent of surviving bacteria during freeze/thaw cycles (Postgate and Hunter, 1961; Nelson and Parkin-son, 1979). However, these bacteria are not known to be halotolerant and therefore might suffer more under increased osmotic stress than \( P. \) halocryophilus does. In the future, freeze/thaw experiments with halophilic microorganisms such as \( P. \) halocryophilus should also include other types of salts in the growth medium to test their influence on cell survivability in comparison with NaCl.

On Mars, NaCl has been detected globally, and especially at high levels by remote sensing in the Southern Highlands (Osterloo et al., 2008). Perchlorates have been detected at the Phoenix Lander and the Curiosity Rover sites and are likely global in extent (Clark and Kounaves, 2016). These salts have also been suggested to be part of the brines associated with the recurrent slope lineae (RSL) (Ojha et al., 2015). However, more recent studies have argued that only small amounts of water might be present within the RSL (Edwards and Pi-queux, 2016) and that the darkening of the RSL might only be a result of a rewetting process of former flows of salty water (Heinz et al., 2016). Furthermore, it has also been suggested that RSL may be the result of granular flows where water plays no, or only a subordinate, role (Dundas et al., 2017).

However, in general, the ubiquitous presence of hygroscopic salts and of water in the form of ice on the poles or in the subsurface or as gas in the atmosphere makes the existence of cold, highly concentrated brines conceivable. Such brines could develop through deliquescence or at salt-ice contacts (Fischer et al., 2014), being temporally stable at the surface of Mars and perhaps permanently stable in the subsurface as briny groundwater (Burt and Knauth, 2003; Martinez and Renno, 2013).

Our data reveal that microorganisms resident in such brines could survive significantly longer at subzero temperatures than previously thought, and they might even thrive in slightly diluted brines as has been shown for \( P. \) halocryophilus in a 19 wt% NaCl solution (Mykytczuk et al., 2012). As temperatures on Mars change throughout the day and the seasons, it is conceivable that temperatures drop temporally below the eutectic temperature of the brine. Our freeze/thaw experiments demonstrate that the freezing and thawing of cells in eutectic brines would be less lethal than freezing and thawing in salt-free water.

5. Conclusion

We have shown enhanced microbial survival in subzero eutectic Cl\(^-\) brines compared with their warmer analogues. Based on the results, the best hypothesis is that the increase in size and stability of hydration shells around ions at lower temperatures reduces osmotic and chaotropic stress factors for microbial organisms.

Although \( P. \) halocryophilus grew even in the presence of 10 wt% NaClO\(_4\), higher ClO\(_4^-\) concentrations lower survival rates significantly even at subzero temperatures. It appears that the decreased capability of ClO\(_4^-\) ions to form stable hydration spheres causes the high toxicity of eutectic ClO\(_4^-\) solutions and the lower temperature dependence of cell survival compared with Cl\(^-\) brines. Furthermore, we have shown that the presence of salts such as NaCl increases the survivability during freeze/thaw processes. This has broad implications for the habitability of some extreme environments on Earth and the potential habitability of Mars.

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Author Disclosure Statement

No competing financial interests exist.

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**Abbreviations Used**

CFUs = colony forming units
PBS = phosphate-buffered saline
RSL = recurrent slope lineae
ST = sample type