Dehydrated *Caenorhabditis elegans* Stocks Are Resistant to Multiple Freeze-Thaw Cycles

Patrick D. McClanahan,* Richard J. McCloskey,*  † Melanie Ng Tung Hing,* David M. Raizen,† and Christopher Fang-Yen*‡

*Department of Bioengineering, School of Engineering and Applied Science, †Department of Neurology, Perelman School of Medicine, and ‡Department of Neuroscience, Perelman School of Medicine, University of Pennsylvania, Philadelphia, Pennsylvania 19104

ORCID IDs: 0000-0003-2817-5948 (P.D.M.); 0000-0001-5935-0476 (D.M.R.); 0000-0002-4568-3218 (C.F.-Y.)

**ABSTRACT**

Ultracold preservation is widely used for storage of genetic stocks of *Caenorhabditis elegans*. Current cryopreservation protocols are vulnerable to refrigeration failures, which can result in the loss of stock viability due to damage during re-freezing. Here we present a method for preserving worms in a dehydrated and frozen form that retains viability after multiple freeze-thaw cycles. After dehydration in the presence of trehalose or glycerol, *C. elegans* stocks can be frozen and thawed multiple times while maintaining viability. While both dauer and non-dauer larvae survive desiccation and freezing, the dauer defective mutant *daf-16* does not survive desiccation. Our technique is useful for storing stocks in a manner robust to freezer failures, and potentially for shipping strains between laboratories.

Cryopreservation enables collections of biological specimens to be stored for extended periods with little maintenance. Organisms capable of maintaining viability after recovery from long-term cryopreservation include bacterial strains, eukaryotic cell lines, embryos, and small animals such as the genetic model organism *Caenorhabditis elegans* (Brenner 1974). However, during the freezing process, the specimen may be damaged directly by the formation of ice crystals and by osmotic stress and/or toxicity resulting from the increased concentration of aqueous solutions left as ice forms (Mazur et al. 1972; Muldrew and McGann 1990, 1994; Pegg 2009). For this reason, cryopreservation methods rely on manipulations that reduce the damaging effects of ice crystals.

Most contemporary cryopreservation protocols, including those for *C. elegans* (Brenner 1974; Siomi and Kuroki 2003), minimize freezing damage by slowly chilling the specimen in a buffer containing a cryoprotective agent. As the specimen is cooled, ice forms in the freezing buffer, concentrating the cryoprotectant. The slow rate of cooling allows the specimen to maintain a safe osmotic balance with the surrounding buffer (Muldrew and McGann 1994) while the increasingly concentrated cryoprotective agent inhibits formation of intracellular ice crystals (Morris et al. 2006) and may stabilize lipid membranes and proteins (Rudolph and Crowe 1985; Crowe et al. 1992; Burkewitz et al. 2012; Julca et al. 2012).

Standard preservation techniques in *C. elegans* rely on glycerol added to freezing media as a cryoprotectant (Brenner 1974; Siomi and Kuroki 2003), but minimal freezing damage by slowly chilling the specimen in a buffer containing a cryoprotective agent. As the specimen is cooled, ice forms in the freezing buffer, concentrating the cryoprotectant.
agent that protects against dehydration damage) bear some resemblance to those of glycerol in cryopreservation and include its ability to stabilize lipid bilayers as well as proteins (Carpenter and Crowe 1988; Crowe et al. 1998). Indeed, trehalose can be used as a laboratory cryoprotectant (Bhandal et al. 1985; De Antoni et al. 1989; Beattie et al. 1997), and glycerol can be used as a xeroprotectant (Jin et al. 1996; Puhlev et al. 2001). We therefore hypothesized that *C. elegans* stocks dried in the presence of protective agents might be capable of maintaining viability through multiple freeze-thaw cycles.

In this report we demonstrate that the protective agents trehalose or glycerol allow *C. elegans* to retain viability after dehydration, and that stocks dehydrated in this manner are resistant to freeze-thaw damage. We present a simple method for dehydrating and freezing worms that results in stocks that can survive multiple freeze-thaw cycles. This method is practical for the long-term preservation of *C. elegans* genetic stocks in a manner robust to refrigeration failure.

**MATERIALS AND METHODS**

**C. elegans strains and maintenance**

The following strains were used in our study: Bristol N2 (WT) and CF1038 daf-16(mu86). We cultured *C. elegans* on OP50 *E. coli* food bacteria on standard NGM agar (Stiernagle 2006) or standard NGM agar with an additional 200 mM NaCl (salt conditioning experiments only) in 6-cm plates. To prepare worms, we picked five gravid adult hermaphrodites onto each plate and allowed the population to grow at ambient temperature (RT, 16–24°C) for three weeks prior to conducting experiments. The food bacteria became depleted about five days after picking, so these animals were without food for about two weeks. With the exception of freezing and long-term storage experiments, culturing and experiments were performed at RT.

**Buffers and solutions**

M9 buffer, S buffer, and soft agar freezing media were made according to standard methods (Stiernagle 2006). Alternative freezing (AF) media consisted of 15.1 g trehalose, 17.7 mL DMSO, and M9 buffer to a total volume of 500 mL (Kevin O’Connell, personal communication). All cryo/xeroprotectants were prepared immediately prior to use as solutions in M9 at 30% concentration (v/v for glycerol and DMSO, w/v for trehalose).

**Liquid freezing (LF)**

The LF procedure was previously described (Stiernagle 2006) except that starved animals from three week old plates were used instead of freshly starved animals, and M9 buffer was used instead of S buffer (Brenner 1974). *C. elegans* were washed from growth plates into a 15 mL conical centrifuge tube using 0.7 mL M9 per plate. An equal volume of M9 with 30% (v/v for trehalose, v/v for glycerol and DMSO, or M9 alone) xeroprotectant solution in M9 for a final concentration of 15%. We added 50 μL of well-mixed worm-xeroprotectant mixture to each cryovial and 1 μL was pipetted onto a food plate for baseline counting (see counting section). The uncapped cryovials were placed in an airtight box also containing approximately 5 g anhydrous calcium sulfate desiccant (Hammond Drierite 23001, 8 mesh) per cryovial and allowed to dry for 48 h at RT. Trehalose-containing buffer dried to a soft solid, and glycerol-containing buffer dried to a viscous liquid. See also Supplemental Protocol (File S2).

**Soft agar freezing (SA)**

The SA procedure was as previously described (Stiernagle 2006) except that starved animals from three week old plates were used instead of freshly starved animals. Worms were washed from growth plates into a 15 mL centrifuge tube using 0.7 mL S buffer per plate and placed on ice. After 15 min, an equal volume of molten, 50% soft agar freezing solution was added to the conical and mixed by gentle vortexing. Freezing and recovery was identical to that described for the LF procedure.

**Trehalose-DMSO alternative freezing (AF)**

The trehalose-DMSO freezing protocol was similar to one in use elsewhere (K. O’Connell, personal communication). Worms starved for 3 weeks were washed from their growth plates with M9, pelleted at 700 rcf in a clinical centrifuge for 3 min, washed once in trehalose-DMSO freezing buffer, and resuspended in trehalose-DMSO freezing buffer to make a worm freezing mixture. We added 1 mL of worm freezing mixture into each cryovial and placed them in a -78.5°C dry ice-liquid cooler in a foaming container.

**Desiccation with xeroprotectants**

Animals were transferred from several plates into a 15 mL centrifuge tube by floating them in M9 buffer using a transfer pipette (USA Scientific), then pelleted (5 min at 700 rcf), washed in 10 mL of fresh M9, pelleted again, and transferred to a 1.5 mL microcentrifuge tube. We added M9 to the tube to adjust the volume to approximately 50 μL per worm plate used. After brief vortexing, this worm mixture was combined with equal parts of 30% (w/v for trehalose, v/v for glycerol and DMSO, or M9 alone) xeroprotectant solution in M9 for a final concentration of 15%. We added 50 μL of well-mixed worm-xeroprotectant mixture to each cryovial and 1 μL was pipetted onto a food plate for baseline counting (see counting section). The uncapped cryovials were placed in an airtight box also containing approximately 5 g anhydrous calcium sulfate desiccant (Hammond Drierite 23001, 8 mesh) per cryovial and allowed to dry for 48 h at RT. Trehalose-containing buffer dried to a soft solid, and glycerol-containing buffer dried to a viscous liquid. See also Supplemental Protocol (File S2).

**Simulated refrigeration failure and freeze-thaw cycling**

Cryovials containing *C. elegans* were placed in an expanded polystyrene foam shipping container (same as used for conventional freezing) which was placed in an insulated container lined with dry ice slabs on the bottom and two sides. The following day, cryovials subjected to a single freeze-thaw (1X) were removed, thawed, and rehydrated (see rehydration and recovery). The remaining vials were kept in the shipping container, which was stored at RT for approximately 8 h and then returned to -78.5°C to simulate a single refrigeration failure (2X). The protocol was repeated the next day to simulate a second refrigeration failure (3X). On the last day, cryovials subject to an additional three rounds of freeze-thaw cycling (6X) were placed in a cardboard cryostorage box and switched between RT and -78.5°C at 1 h intervals.

**Counting animals before and after freezing and/or desiccation**

For all counting, we pipetted some or all of the volume of worm suspension onto an NGM plate that had been seeded with a lawn of OP50 bacteria. To minimize underestimation due to worms sticking to the inside of the pipette, we pipetted the worm suspension up and down once to pre-coat the inside of the pipette tip with worms before...
transferring worms. We counted animals that moved spontaneously or in response to either tapping the plate or to touching them with a worm pick made of platinum-iridium wire (Tritech Research PT-9010, 254 μm diameter). Counted animals were removed from the plate.

To estimate the number of worms prior to freezing and/or desiccation, we counted animals in a small sample of the worm suspension (5 μL for SA, LF, and AF, 1 μL for desiccation). Sample volumes were chosen in order to maintain the number of counted worms in roughly the range of 100 - 500. Dauer, L4, and adult animals were counted within 1 h after pipetting the worm sample on the agar surface, and the remaining larvae counted the following day, allowing L1s to reach a more easily visible size. The number of worms placed in each tube for freezing and/or desiccation ranged from 3800 to 59200 (mean 13221).

To count the number of animals surviving after recovery, we used a similar procedure, except counting was performed a few hours after recovery and then repeated daily for six days to allow animals additional time to recover. For LF, SA, and AF experiments, we allowed the vials to thaw for 15 min, vortexed the contents briefly, and pipetted 10 μL (single freeze-thaw samples) or the full 1 mL (one or more simulated refrigeration failures) of the thawed suspension onto a food plate. For desiccation experiments, we allowed vials to thaw for 10 min (if frozen or chilled), rehydrated the worms in 50 μL of M9 at RT for 10 min, mixed the worm suspension by tapping the side of the cryovial, and pipetted 5 – 6.5 μL (WT in trehalose or glycerol buffer except 1 or 3.5 month storage at 20°C) or the full 50 μL (all daf-16, WT in plain M9, WT in DMSO buffer, and WT in trehalose or glycerol buffer stored for 1 or 3.5 months at 20°C) of the worm suspension onto a food plate.

Since some worms took several days to recover, and plates were assayed once per day, there was some uncertainty in the identification of developmental stages. During counting, we classified surviving worms as L1-L3 larvae, dauer larvae, or L4 / adult. Because dauers develop into L4s, recovered L1-L3 larvae were assumed to have survived freezing and/or drying as non-dauer larvae. Therefore, L1-L3 survival rate was estimated by dividing the number of L1-L3 larvae recovered by the number of L1-L3 larvae desiccated / frozen. Similarly, because the food-rich, low population density recovery plate conditions favor dauer exit rather than dauer entry, recovered dauers were assumed to have survived as dauers as opposed to having entered the dauer state after recovery. Therefore, dauer survival rate was estimated by dividing the number of dauers recovered by the number of dauers desiccated / frozen. In contrast, recovered L4 / adults could have developed from surviving dauer larvae or L1-L3 larvae, or survived as L4 / adults. Therefore, these animals were not used in estimations of survival by type, but did contribute to calculations of overall survival. The type-specific survival rates may be underestimated because some surviving L1-L3s and dauers could have developed into L4s or adults after recovery but before being counted.

**Imaging dehydration and rehydration of worms in a single droplet**

We recorded the process of dehydration and recovery using a Leica M165 FC stereo microscope with a 5 megapixel CMOS camera (Imaging Source DMK 23GP031) attached to the trinocular port. We prepared 3-week starved N2 animals in M9 solution containing 15% trehalose, then pipetted a 10 μL droplet of the suspension onto a polydimethylsiloxane (PDMS, Dow Corning Sylgard 184) surface in a clear polystyrene dish. We used PDMS for its optical clarity, allowing the entire process to be recorded, and its non-stick nature, enabling us to easily remove the dehydrated flake. After pipetting, we added approximately 1 g of anhydrous calcium sulfate to the polystyrene dish as a desiccant and sealed the dish with Parafilm. The next day, we used forceps to transfer the sample to an OP50-seeded plate. We then recorded images for 5 d as animals recovered and reproduced (Supplemental Video File S1).

**Plotting and statistics**

All statistical tests and plotting were performed in MATLAB. All bar graphs show mean ± SEM. The number of replicates for each experiment is given in the figure captions or text. Circles represent data from individual replicates. We used α = 0.05 for determining statistical significance. Statistical tests are described in Results.

**Data availability**

All data required for confirming the conclusions of this article are represented fully within its text and figures. Supplemental material available at figshare: https://doi.org/10.25387/g3.13060790.

**RESULTS**

**C. elegans frozen by established methods do not survive repeated freeze-thaw cycles**

It has been reported that a power failure can result in loss of all *C. elegans* stocks stored in a -80°C freezer (Stiernagle 2006). To confirm...
this observation, and to obtain baseline data against which to compare subsequent results, we quantified the viability of *C. elegans* frozen by standard methods after single or repeated freeze-thaw cycles.

Two *C. elegans* freezing protocols are in common use (Stiernagle 2006). In liquid freezing (LF), starved worms are frozen in liquid buffer supplemented with 15% glycerol as cryoprotectant. In soft agar freezing (SA), a small concentration of low melting temperature agar is added to the freezing medium; the agar keeps worms suspended throughout the medium, such that repeated small portions of the frozen medium can be removed and thawed instead of thawing the entire tube. We froze WT animals using both the LF and SA protocols, then measured their survival rates upon thawing with or without one or two simulated refrigeration failures. In a simulated refrigeration failure, tubes housed in an insulated expanded polystyrene box were thawed at RT for 8 h (see Methods) and then refrozen.

Survival after a single freeze and thaw, corresponding to the normal procedure for cryopreservation, was $36 \pm 4\%$ for LF and $41 \pm 11\%$ for SA (Figure 1a-b), consistent with the previously reported range of 35–45% (Stiernagle 2006). After a second freeze-thaw cycle, survival dropped precipitously to $0.037 \pm 0.032\%$ for LF and $0.020 \pm 0.001\%$ for SA, although a few individuals from each of the three vials tested per condition survived. After three freeze-thaw cycles, there were no survivors out of over 200,000 worms frozen. Interestingly, dauer larvae, which are resistant to a wide range of stressors including cold (Savory et al. 2011), survived at a rate similar to non-dauer larvae, confirming prior observations that dauer larvae are not more resistant to freezing than non-dauer larvae (Stiernagle 2006).

While conventional freezing buffers include glycerol as a cryoprotectant, some laboratories have developed *C. elegans* freezing buffers with alternative cryoprotectants, for example the disaccharide trehalose (Mitani 2009). To determine if a different type of cryoprotectant might better protect *C. elegans* over multiple freeze-thaw cycles, we also tested an alternative freezing (AF) buffer including both trehalose and DMSO (see Methods). This buffer performed similarly to LF and SA (Figure 1c).

Together, these results confirm that standard frozen *C. elegans* stocks are highly vulnerable to freeze-thaw cycles.

**Exogenous trehalose or glycerol can serve as *C. elegans* xeroprotectants**

Since damage during freezing is thought to be partly caused by the formation of ice crystals (Mazur 1963; Pegg 2009), we hypothesized that desiccation prior to freezing might improve *C. elegans* survival after repeated freeze-thaw cycles. However, like freezing, desiccation also causes damage to *C. elegans* (Ohba and Ishibashi 1981; Gal et al. 2004; Erkut et al. 2011). We therefore first sought to develop a simple method for dehydrating worms while preserving viability.

Some compounds, such as trehalose and glycerol, can serve as both cryoprotectants and xeroprotectants (Julca et al. 2012). *C. elegans* adaptation to osmotic stress involves the production of endogenous glycerol (Lamitina et al. 2004), and the ability of dauer larvae to survive desiccation depends on genes required for production of trehalose (Erkut et al. 2011).

We asked whether the presence of exogenous protective agents might improve desiccation tolerance in various developmental stages of *C. elegans*. In addition to trehalose and glycerol, we also tested dimethylsulfoxide (DMSO), which has been used as a cryoprotectant (Pegg 2009), including for the nematodes *C. briggsae* (Hwang 1970) and *C. elegans* (K. O’Connell, personal communication). We desiccated starved *C. elegans* in droplets of M9 buffer alone or M9 supplemented with 15% trehalose (Figure 2-3, Supplemental Video File S1), 15% glycerol, or 15% DMSO for 48 h in uncapped cryotubes placed in sealed containers containing a desiccant. We then rehydrated and transferred the desiccated worms to the surface of NGM plates seeded with bacteria (Figure 2-3) and tracked their recovery.

Addition of either trehalose or glycerol led to a robust increase in survival after desiccation: overall survival was $5.9 \pm 1.2\%$ for worms...
Having established that exogenous trehalose and glycerol promote freeze-thaw cycles with little reduction in viability, we next asked whether the presence of trehalose or glycerol is similarly higher for dauer and non-dauer larvae. Surviving L4s and adults were excluded because they could have survived desiccation as either dauer or non-dauer larvae and molted prior to being counted. With either trehalose or glycerol, dauer survival was several times higher. Survival of non-dauer larvae also improved (Figure 3).

These results show that exogenous trehalose and glycerol can serve as xeroprotectants for both dauer and non-dauer larvae.

Desiccated C. elegans stocks withstand multiple freeze-thaw cycles with little reduction in viability

Having established that exogenous trehalose and glycerol promote tolerance of desiccation, we next asked whether C. elegans stocks desiccated in the presence of these agents can maintain viability through freeze-thaw cycles. We subjected worms dried with exogenous trehalose and glycerol to one, two, or three freeze-thaw cycles, as before. We then subjected some samples to an additional three rapid freeze-thaw cycles with little reduction in viability. Desiccated C. elegans stocks withstand multiple freeze-thaw cycles. We then subjected some samples to an additional three rapid freeze-thaw cycles before rehydration and recovery.

We observed no significant correlation between number of freeze-thaw cycles and overall survival for samples dried in trehalose buffer (coefficient of correlation $\rho = -0.079$ and $P = 0.78$) or glycerol buffer ($\rho = -0.23$ and $P = 0.41$) (Figure 4), indicating that dehydrated stocks of C. elegans do not lose viability when repeatedly frozen and thawed up to at least six cycles.

These results show that C. elegans desiccated with exogenous trehalose and glycerol are robust to multiple freeze-thaw cycles.

Desiccated C. elegans retain viability indefinitely at low temperature

The ability to survive multiple freeze-thaw cycles represents a major advantage of desiccated C. elegans stocks over conventional C. elegans freezing protocols. To further evaluate the suitability of our method for long-term storage, we asked how long C. elegans dehydrated with a protective agent remain viable at various temperatures. We dehydrated worms in trehalose or glycerol media and stored sealed tubes at either -80°C, -20°C, +4°C, or +20°C. We then assayed survival after one day, one week, one month, and 3.5 months (Figure 5).

By fitting with an exponential function $s(t) = s_0 \exp(-t/\tau)$ we estimated the characteristic decay time $\tau$ for overall survival at each temperature. As expected, survival decayed more rapidly at higher temperature. Estimated decay times were $\tau = 267 \pm 6$ d (95% confidence interval 82-\infty d) at -20°C, $\tau = 61 \pm 1 \text{ d (95\% CI 18-\infty d) at +4\°C, and } \tau = 3.2 \pm 0.3 \text{ d (95\% CI 3.0-3.4 d) at +20\°C for worms dried in trehalose buffer, }$ and $\tau = 51 \pm 5 \text{ d (95\% CI 28-237 d) at +20\°C for worms dried in glycerol buffer.}$

We noted that the proportion of dauers surviving in this set of experiments was lower than in previous experiments. This may have been due to fluctuations in cultivation temperature which ranged from 16-24°C (see Methods).

These results suggest that desiccated tocks can be stored indefinitely under ultracold temperatures, but also remain viable for extended periods under ordinary refrigeration and several days at room temperature. These findings may be useful for managing storage of desiccated stocks after a freezer failure.

Starvation time affects survival after desiccation

For the results reported above, we used animals from plates that were initially populated with five gravid adults and then incubated at 20°C for three weeks. Under these conditions, the growing population depletes the bacterial food lawn after about five days, and thus these animals have starved for approximately 16 days. In contrast, the classical C. elegans freezing protocols call for freshly starved animals (Stiernagle 2006).

Since starvation of C. elegans activates stress resistance pathways (Weinkove et al. 2006; Larance et al. 2015), and has been shown to upregulate endogenous trehalose production (Hibshman et al. 2017), we ased if starvation time may affect the rate of survival after desiccation.

We compared survival after drying in trehalose buffer of worms incubated for three weeks vs. worms incubated for one week. We found that overall survival was significantly higher for worms incubated for three weeks ($\pm 0.6\%$ vs. $1.1 \pm 0.4\%$, $P = 0.009$, Wilcoxon rank-sum test, 6 replicates with an average of 7,700 and 12,500 worms per replicate for three week and one week worms, respectively). However, we did not find a significant difference between survival rates of non-dauer (L1-L3) larval stages in the two groups ($1.8 \pm 0.5\%$ for three week plates vs. $0.95 \pm 0.28\%$ for one week plates, $P = 0.22$, Wilcoxon rank-sum test). While dauers were 7.6% of the total population on the three week plates, they made up only 0.5% of the population in the one week plates. Dauers from three week plates survived desiccation at an average rate of 32%; we did not
measure the survival rate of dauer larvae from one week plates due to their low numbers.

These results suggest that prolonged starvation increases survival after desiccation primarily by increasing the abundance of dauer larvae.

The dauer defective mutant daf-16 does not survive desiccation

Some C. elegans mutants have a lower rate of survival than wild-type following freezing using conventional methods. One example is mutants for the forkhead box (FOXO) homolog DAF-16 (Hu et al. 2015). DAF-16 is involved in the response to cellular stressors, and daf-16 loss-of-function mutants are dauer defective (cannot enter the dauer stage). To determine if daf-16 mutants can be stored in desiccated form, we desiccated daf-16(mu86) mutants in our trehalose and glycerol supplemented buffers. Upon rehydration, no survivors were recovered from four replicates using trehalose buffer, and a single survivor was recovered from four replicates using glycerol buffer.

DAF-16 regulates production of endogenous trehalose and glycerol (Hibshman et al. 2017). However, following preconditioning on plates containing high NaCl, wild-type and daf-16 mutants survive acute osmotic shock equally well (Lamitina and Strange 2005). We hypothesized that preconditioning daf-16 mutants on high NaCl plates might enable them to survive desiccation. To test this hypothesis, we dried and attempted to recover daf-16 mutants raised on NGM plates containing an additional 200 mM NaCl. Out of three replicates for each protective agent, trehalose and glycerol, we did not recover a single survivor. This result shows that the dauer defective daf-16 mutant does not survive desiccation, even when preconditioned to withstand osmotic shock.

DISCUSSION

We have described a method for desiccating and freezing C. elegans in the presence of xeroprotective agents. Strains stored this way can survive cycles of warming and refreezing that accompany equipment failures or power outages. We provide a detailed version of our protocol in Supplemental File S2.

While C. elegans dauers (Erkut et al. 2011), and potentially other developmental stages (Ohba and Ishibashi 1981), were previously shown to be capable of anhydrobiosis, we found that the addition of the exogenous xeroprotective agents trehalose or glycerol dramatically improved survival rates. In the wild, C. elegans commonly inhabits decomposing fruit (Frézal and Félix 2015). While most fruits do not contain trehalose, other sugars often found at high concentrations in fruits, including fructose, glucose, and sucrose, have been shown to act as cryoprotectants in some organisms (Storey and Storey 1991) and may also function as xeroprotectants. Ethanol, a product of sugar fermentation in rotting fruit (McKenzie and McKechnie 1979), has been shown to increase C. elegans desiccation tolerance (Kaptan et al. 2020). Therefore it is possible that the ability to survive drying in the presence of exogenous sugar and other components of C. elegans’ natural surroundings may provide a selective advantage.

We demonstrated that dehydrated stocks can be frozen and thawed multiple times without significant loss of viability, and that they can be stored at warmer temperatures for shorter periods. The ability of desiccated stocks to retain viability at conventional refrigeration temperatures may be useful after a refrigeration failure.

A significant limitation of our method is that daf-16 mutants, and perhaps others, do not survive drying and refreezing. In conventional freezing, it is important to conduct a test thaw whenever preparing strains for storage.

Our attempts to precondition daf-16 mutants by culturing them on high-salt plates did not improve survival after desiccation. It is possible that the expression of other DAF-16-regulated products, such as genes involved in the hypertonic stress response (Lamitina and Strange 2005), is required in addition to exogenous cryoprotectants for enhanced survival after desiccation. It is also possible that the ability to form dauers is required for desiccation survival, and that other dauer defective mutants (Hu 2007) would also fail to survive desiccation.

Our method may be useful for the exchange of strains between C. elegans laboratories. Strains are normally shipped on NGM plates, which are vulnerable to cold temperatures during transport, and require the sender to subculture worms before shipping. Shipping desiccated C. elegans would have the advantages of not requiring subculture of strains at either end, and of being impervious to cold, although desiccated worms remain vulnerable to heat.

There have been relatively few descriptions of cryopreservation of desiccated organisms. In freeze-drying (lyophilization), samples are frozen first, then dehydrated, then stored at low temperature or room temperature. While lyophilization is often done with microbes (Morgan et al. 2006), few animals or animal cells can be stored this way (Katkov et al. 2012). In contrast, our method involves dry freezing, i.e., first drying and then freezing the specimen. Reports

![Freeze-thaw survival of desiccated worms](chart)

**Figure 4** Desiccated worms survive multiple freeze-thaw cycles. Survival of worms dried in M9 buffer supplemented with 15% trehalose or 15% glycerol and subjected to one, two, three, or six freeze-thaw cycles. There are three replicates per condition. Black bars indicate overall survival. Gray and white bars indicate L1-L3 and dauer survival, respectively. Error bars represent ± SEM. Circles represent individual replicates. Each replicate contains approximately n = 15000 - 22500 animals.
of dry freezing as a practical storage method have been limited to the preservation of plant seeds (Pritchard 2007) and seedlings (Sun 1958). However, survival of freezing by dehydration seems to be relatively common in nature, particularly in soil-dwelling animals (Holmstrup et al. 2002). Our method may suggest exogenous xeroprotectant-based dry freezing protocols in other systems.

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