All Eggs Are Not Equal: The Maternal Environment Affects Progeny Reproduction and Developmental Fate in Caenorhabditis elegans

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Abstract

Background: Maternal effects on progeny traits are common and these can profoundly alter progeny life history. Maternal effects can be adaptive, representing attempts to appropriately match offspring phenotype to the expected environment and are often mediated via trade-offs between progeny number and quality. Here we have investigated the effect of maternal food availability on progeny life history in the free-living nematode Caenorhabditis elegans.

Methodology/Principal Findings: The maternal environment affects both reproductive traits and progeny development. Comparisons of the progeny of worms from high and low maternal food environments indicates that low maternal food availability reduces progeny reproduction in good environments, increases progeny reproduction in poor environments and decreases the likelihood that progeny will develop as dauer larvae. These analyses also indicate that the effects on progeny are not a simple consequence of changes in maternal body size, but are associated with an increase in the size of eggs produced by worms at low maternal food availabilities.

Conclusions/Significance: These results indicate that the maternal environment affects both progeny reproduction and development in C. elegans and therefore that all progeny are not equal. The observed effects are consistent with changes to egg provisioning, which are beneficial in harsh environments, and of changes to progeny development, which are beneficial in harsh environments and detrimental in benign environments. These changes in progeny life history suggest that mothers in poor quality environments may be producing larger eggs that are better suited to poor conditions.

Introduction

Maternal effects are widespread in natural and experimental populations [1,2] and can have pervasive effects on population dynamics [3]. Many maternal effects are a response to unfavorable conditions and are often mediated via trade-offs between progeny number and aspects of progeny quality [4]. These trade-offs can represent attempts to appropriately match progeny phenotype to the expected progeny environment [1,4,5]. For example, in Daphnia, poor maternal conditions often result in the production of a smaller number of larger eggs, with progeny subsequently performing better under a range of stressful conditions (e.g. [6–10]). Other effects can be more complex, producing changes to offspring life history that are not directly related to resource provisioning (e.g. [11]) or that interact in complex ways with the direct effects of conditions on the mother [12–14].

Here, the effects of the maternal environment on progeny reproduction and development are investigated in the free-living nematode Caenorhabditis elegans. In the wild, C. elegans is associated with nutrient- and bacteria-rich substrates [15–17]. Dispersal between these ephemeral resource patches is undertaken by an alternative third larval stage, the developmentally arrested and long lived dauer larvae [18–20]. Growing populations can reach very large sizes, and it is likely that C. elegans routinely experiences very high population densities and has therefore evolved to optimize fitness, i.e. population growth and/or the production of dispersal stages, under such conditions.

Extensive studies of individual worms have defined development and reproduction in C. elegans (e.g. [21–25]). However, population sizes predicted from individual worm data significantly overestimate observed population sizes, indicating that there are strong density dependent effects on population growth in C. elegans [24]. The observed food concentration-dependent reduction in fecundity seen in C. elegans partly underlies this discrepancy [24], but does not fully explain it. One possible explanation is that, in addition to the direct effects on worms that are seen under poor conditions, there are also changes in the progeny produced by worms in poor conditions. Understanding how C. elegans responds to poor conditions is important in understanding the ecology and evolutionary biology of the species, and will also allow the molecular and genetic bases of such effects to be investigated in this important model species.

Here we show, for the first time, that maternal resource limitation affects progeny reproduction and development in C.
elegans. These effects are shown to be not directly related to changes in maternal body size, but are associated with changes in egg size. These changes are consistent with an adaptive explanation, whereby low maternal food availabilities result in fewer larger eggs and progeny that are better adapted to poor conditions.

**Results**

Maternal food availability affects progeny reproductive traits

To investigate the effects of maternal food availability on progeny reproductive traits, eggs were isolated from early adult worms grown at differing food availabilities and progeny were allowed to develop at *ad libitum* food conditions. In these conditions, progeny lifetime fecundity (Figure 1A: \(F_5, 148 = 2.69, p = 0.017\)) and reproductive schedule was affected by maternal food availability (Figure 1B & C), with progeny from high and very low maternal food availabilities having higher lifetime fecundity than progeny from mothers at intermediate food levels. Progeny from the highest maternal food concentrations also showed the greatest early reproduction (Figure 1B & C), while progeny from very low maternal food concentrations showed greater late fecundity (Figure 1C). As only limited numbers of eggs could be isolated from mothers at the lowest food concentrations used here, these changes could not be analyzed in greater detail. However, the changes in the reproductive schedule of the progeny from very low maternal food concentrations would slow their expected rate of population growth. These data indicate that maternal conditions can affect progeny reproductive traits in *C. elegans*.

Maternal size does not fully explain variation in progeny reproductive traits

Fecundity is often found to increase with body size and in the nematoda there is generally a close linkage between adult female body size and fecundity [26]. Therefore, the influence of maternal body size on progeny reproduction was investigated. Progeny were isolated from individual adult worms from three maternal food treatments at each of two maternal age groups. Here, maternal food treatments were selected at the point of progeny isolation from a wider range of food concentrations. This was done to account for variation between experiments in batches of *E. coli* and in the maternal *per capita* food availability. High maternal food availability represented *ad libitum* food and excess food remained on these plates. Low maternal food availability represented the lowest food concentration from which sufficient progeny could be isolated and no food remained on these plates. An intermediate food concentration, where small quantities of food remained on the plates, was then selected to represent Medium maternal food availability. Analysis of maternal size indicated that, as expected, older mothers were larger (AGE: \(F_{1, 41} = 144.6, p < 0.001\)) and that decreased food availability reduced maternal size (FOOD: \(F_{2, 41} = 64.7, p < 0.001\)). There was no interaction between maternal food availability and maternal age (FOOD×AGE: \(F_{2, 41} = 0.7, p = 0.53\)). This indicates that the maternal size did differ across the High, Medium and Low food concentrations chosen and that it did so in a similar way in the two maternal age groups.

There was a positive relationship between maternal size and mean progeny lifetime fecundity for five day old mothers (Figure 2B, Pearson product-moment correlation: \(r = 0.63, p = 0.002\)), but not for four day old mothers (Figure 2A, Pearson product-moment correlation: \(r = 0.32, p = 0.10\)). Analysis of early reproduction indicated that there was a positive relationship between maternal size and mean progeny fecundity on the first day of reproduction for both the 4 and 5 day old worms (Pearson product-moment correlation: \(r = 0.45, p = 0.03\) and \(r = 0.44, p = 0.02\), respectively). These data indicate that there is a link between maternal size and offspring reproduction, with larger mothers producing progeny that start reproduction earlier and tend to have higher lifetime fecundity. However, with maternal size fitted as a covariate, analysis of progeny lifetime fecundity indicated that, while maternal age did not alter progeny fecundity (AGE: \(F_{1, 365} = 0.06, p = 0.80\)), decreased maternal food availability did reduce progeny lifetime fecundity (FOOD×AGE: \(F_{1, 365} = 3.32, p = 0.02\), respectively). These data indicate that there is a link between maternal size and offspring reproduction, with larger mothers producing progeny that start reproduction earlier and tend to have higher lifetime fecundity. However, with maternal size fitted as a covariate, analysis of progeny lifetime fecundity indicated that, while maternal age did not alter progeny fecundity (AGE: \(F_{1, 365} = 0.06, p = 0.80\)), decreased maternal food availability did reduce progeny lifetime fecundity (FOOD×AGE: \(F_{1, 365} = 3.32, p = 0.02\), respectively).

![Figure 1. Maternal food availability affects progeny reproduction.](image-url)
Variation in maternal size does not therefore fully explain the differences in progeny reproductive traits when progeny developed in a poor quality environment, worms were recovered from a dauer development assay and their reproductive traits then investigated. Three different maternal food availabilities were investigated and three progeny treatments were analyzed: dauer and non-dauer larvae recovered from a dauer larvae development assay and control worms that had developed at ad libitum food. As dauer development assays are performed at 25°C, fecundity was also analyzed at this temperature and this experiment therefore included two separate stresses. Firstly, all progeny treatments were grown and assayed at 25°C, a temperature at upper limit of the thermal niche that greatly reduces lifetime fecundity [21,23]. Secondly, the dauer and non-dauer larvae recovered from the dauer development assay developed with limited food and high dauer pheromone levels, conditions that are similar to what would be expected in late stage growing populations of C. elegans. This therefore represents the conditions that the progeny of food restricted mothers might be expected to experience in the wild. Lifetime fecundity was reduced in worms recovered from the dauer assays in comparison to that in the control worms (Figure 4), indicating that the conditions in the dauer development assays imposed a further stress above that of development at 25°C. Maternal food availability did not affect lifetime fecundity in control worms, recovered dauer larvae or in recovered non-dauer larvae (Figure 4: F₂, 70 = 1.05, p = 0.36, F₂, 56 = 0.20, p = 0.83, F₂, 75 = 2.32, p = 0.11, respectively). Similarly, there was no effect of maternal food availability on early reproduction in the control worms and recovered dauer larvae (Figure 4A & B: F₂, 70 = 0.66, p = 0.52 and F₂, 56 = 0.24, p = 0.79, respectively). This indicates that the reproductive advantages seen in the progeny of high food mother at 20°C is not seen when progeny are grown at 25°C. However, maternal food availabilities did affect early reproduction in recovered non-dauer larvae (F₂, 75 = 5.02, p = 0.009), with progeny from high maternal food concentrations showing lower early reproduction (Figure 4C). The comparison of Figure 4A with Figures 4B and C indicates that the additional stress of having gone through a dauer development assay does affect the non-dauer worms (reduced early fecundity, Fig. 4C), but not the recovered larvae [18–20]. To investigate if maternal food availability affects the likelihood that progeny develop as dauer larvae, the arrested L1 progeny from differing maternal food availabilities were assayed for dauer larvae development (Figure 3). Analysis of these data indicated that dauer larvae development is affected by maternal food availability (H = 21.31, d.f. = 2, p < 0.001, H = 9.99, d.f. = 2, p = 0.007 and H = 7.88, d.f. = 2, p = 0.019 for assays 1, 2 and 3, respectively), with decreased maternal food availability reducing the proportion of progeny that developed into dauer larvae (Figure 3).

A lower likelihood of dauer larvae development in the progeny of worms from a poor environment could imply that these worms are better suited to growth and reproduction in poor environments. This would imply that changes in progeny life history are adaptive, representing an attempt to increase progeny fitness in poor environments. Alternatively, if progeny are low quality and dauer larvae development is affected by body condition, a common observation in dispersal decisions [27], then this could explain the reduced likelihood of dauer larvae formation. To distinguish between these possibilities, the effects of maternal food availability on progeny reproductive traits were investigated when progeny were allowed to develop in poor conditions.

Changes in progeny life history are consistent with an adaptive explanation

To investigate the effects of maternal food availability on progeny reproductive traits when progeny developed in a poor quality environment, worms were recovered from a dauer development assay and their reproductive traits then investigated. Three different maternal food availabilities were investigated and three progeny treatments were analyzed: dauer and non-dauer larvae recovered from a dauer larvae development assay and control worms that had developed at ad libitum food. As dauer development assays are performed at 25°C, fecundity was also analyzed at this temperature and this experiment therefore included two separate stresses. Firstly, all progeny treatments were grown and assayed at 25°C, a temperature at upper limit of the thermal niche that greatly reduces lifetime fecundity [21,23]. Secondly, the dauer and non-dauer larvae recovered from the dauer development assay developed with limited food and high dauer pheromone levels, conditions that are similar to what would be expected in late stage growing populations of C. elegans. This therefore represents the conditions that the progeny of food restricted mothers might be expected to experience in the wild. Lifetime fecundity was reduced in worms recovered from the dauer assays in comparison to that in the control worms (Figure 4), indicating that the conditions in the dauer development assays imposed a further stress above that of development at 25°C. Maternal food availability did not affect lifetime fecundity in control worms, recovered dauer larvae or in recovered non-dauer larvae (Figure 4: F₂, 70 = 1.05, p = 0.36, F₂, 56 = 0.20, p = 0.83, F₂, 75 = 2.32, p = 0.11, respectively). Similarly, there was no effect of maternal food availability on early reproduction in the control worms and recovered dauer larvae (Figure 4A & B: F₂, 70 = 0.66, p = 0.52 and F₂, 56 = 0.24, p = 0.79, respectively). This indicates that the reproductive advantages seen in the progeny of high food mother at 20°C is not seen when progeny are grown at 25°C. However, maternal food availabilities did affect early reproduction in recovered non-dauer larvae (F₂, 75 = 5.02, p = 0.009), with progeny from high maternal food concentrations showing lower early reproduction (Figure 4C). The comparison of Figure 4A with Figures 4B and C indicates that the additional stress of having gone through a dauer development assay does affect the non-dauer worms (reduced early fecundity, Fig. 4C), but not the recovered
Mean proportions were calculated from 8–20 plates/treatment (median combination of 1.25% and 0.625% w/v maternal food availability was 1.25% w/v combination of 2.5% and 5% w/v and 3, respectively), with eggs from low maternal food availabilities developed as dauer larvae is also shown for each treatment. In all cases mean proportion (Figures 1 & 2 vs. Figure 4). This supports the idea that the observed changes in progeny life history are adaptive. To further investigate this, the effect of maternal food availability on egg size was investigated. These assays indicated that maternal food availability did affect egg size (Figure 5, W = 4693.5, p = 0.016 and W = 7746.5, p = 0.005, respectively), with eggs from low maternal food availabilities being larger than those from high maternal food availabilities. These data indicate that egg size differs across maternal food environments, with low food mothers producing eggs that are, on average, larger than those of high-food mothers. However, variation in egg size within maternal food treatments is high, and eggs from low food mothers are also more variable in size than those from high food mothers (Figure 5).

### Discussion

Here we show that maternal environment changes progeny reproduction and development in *C. elegans* (Figures 1, 2, 3, 4 and Table 1). Many maternal effect genes, primarily identified by their ability to rescue embryonic lethal phenotypes, are known in *C. elegans* [28], and some maternal effect mutations alter progeny reproduction and developmental timing (e.g. [29]). This however represents the first demonstration of maternal effects on reproduction and development in non-mutant *C. elegans*, a finding with important implications for understanding and estimating fitness in *C. elegans*.

These data indicate that low maternal food availability can reduce progeny lifetime fecundity (Figure 1A & Figure 2) and delay progeny reproduction (Figure 1B & C) when progeny are allowed to develop in good environments. In contrast, when progeny are allowed to develop in poor environments, it is high maternal food availability that decreases early reproduction in progeny (Figure 4). Further, low maternal food availability is shown to decrease the likelihood that progeny will develop as dauer larvae (Figure 3). These observed effects of maternal conditions on progeny life history in *C. elegans* could represent a switch to the production of fewer, better provisioned eggs, in poor conditions and would mirror observations in *Daphnia* [6–10]. As such they would represent attempts to match progeny development to the expected environmental conditions and are therefore likely to be adaptive [1,4].

Mechanistically it is not clear how the observed changes in progeny life history are produced. While maternal size is related to progeny reproduction, progeny lifetime fecundity is affected by the maternal environment when maternal size is controlled (Figure 2) and maternal body size and progeny reproductive traits can vary independently (Table 1). It is therefore clear that there is not a simple relationship between progeny quality and parental size. The observed increases in egg size at low maternal food availabilities (Figure 5) suggest that, assuming a direct link between egg size and quality, there could be differences in resource provisioning. However, maternal food availability also changes the number of eggs in utero and the developmental stage at which eggs are laid, which in turn will slightly change the duration of L1 arrest experienced by larvae from different maternal food treatments. It is therefore possible that these differences may affect progeny development and this is the reverse of the pattern observed in Figures 1 and 2. This therefore indicates that, under non-optimal, stressful conditions, the progeny of low food mothers do not perform worse than the progeny of high food mothers (Figure 4A & B) and that under certain conditions they perform better (Figure 4C).

![Figure 3. Maternal food availability affects development fate.](image)

Jitter plot showing the proportions of the L1 progeny of mothers from high (H), medium (M) and low (L) maternal food concentrations that developed as dauer larvae in three independent experiments (1–3). The mean proportion (±95% confidence interval) of L1 progeny that developed as dauer larvae is also shown for each treatment. In all cases High maternal food availability was 10% w/v *E. coli*. Medium maternal food availability was 2.5% w/v *E. coli* for assays 1 and 3, and a combination of 2.5% and 5% w/v *E. coli* treatments for assay 2. Low maternal food availability was 1.25% w/v *E. coli* for assays 2 and 3, and a combination of 1.25% and 0.625% w/v *E. coli* treatments for assay 2. Mean proportions were calculated from 8–20 plates/treatment (median 12).

Superscript letters indicate groups that differ significantly in either maternal length or progeny lifetime fecundity (*p*<0.05), NS indicates no significant difference between treatments.

*The daily fecundity was higher for the progeny of the parents grown at 15°C worms on each day, but these differences were not significant.

| Temperature | Maternal length ± se (mm) | Lifetime fecundity ± se | Reproductive schedule |
|-------------|---------------------------|-------------------------|-----------------------|
| 15°C        | 0.940±0.005              | 280.36±9.02              | NS*                   |
| 20°C        | 0.95±0.004               | 248.64±8.15              | NS                    |
| Control     | 1.226±0.011              | 261.04±9.82              | NS                    |
| Heat shock  | 1.17±0.013               | 262.72±9.00              |                       |

| Superscript letters indicate groups that differ significantly in either maternal length or progeny lifetime fecundity (*p*<0.05), NS indicates no significant difference between treatments. |
reproduction and development. Further, as progeny from low maternal food conditions do not perform best under all conditions (Figures 1 & 2 vs. Figure 4), it is unlikely that there is a simple relationship between egg size and progeny quality. Indeed, the data suggests that the observed effects may be a consequence of changes to egg provisioning, which are beneficial in harsh environments, and of changes to progeny development, which are beneficial in harsh environments and detrimental in benign environments. It would also be interesting to know how differences in progeny lifetime fecundity are produced. Given that lifetime fecundity in *C. elegans* under *ad libitum* food conditions is normally constrained by the number of sperm produced during the L4 stage [30], this suggests that either the duration or rate of spermogenesis has been altered. Any such changes might then alter the onset of reproduction.

Given that very small changes (2–3 hours) in the onset of reproduction have dramatic effects on population growth rate in *C. elegans* [22] the observed changes in progeny life history will affect the properties of growing populations. Indeed, changes in early reproduction are likely to be particularly important for *C. elegans* in the wild given the greatly reduced lifespan observed under more natural conditions [31]. However, given the complex nature of the changes in progeny traits, and the changes in the likelihood that worms will develop as dauer larvae (Figure 3), it is difficult to predict how the dynamics of growing populations of *C. elegans* will be affected by the maternal effects observed here. Given this, it is a priority to determine how eggs differ in their composition, if differences in gene expression in developing larvae can be identified and to determine the mechanistic bases of changes in progeny reproductive traits.

**Materials and Methods**

**Worms**

Experiments were performed using the N2 isolate of *C. elegans*, obtained from the *Caenorhabditis* Genetics Center, which is funded by the NIH National Center for Research Resources (NCRR). Synchronized populations of L1s were produced by allowing eggs isolated from hypochlorite treated adults [32] to hatch on plates without food and to develop at 20°C for 24 hours. This causes worms to arrest development prior to the L1/L2 molt, a state in which they can survive for several weeks, and hence produces a cohort of worms at the same developmental stage. All treatments were randomized and blind coded, with plates on which worms failed to grow, burrowed into the agar or climbed the sides of the plates excluded from analyses.

**Maternal food availability and progeny reproduction**

Arrested L1s were transferred, at 20°C, to plates containing a food source consisting of 100 μL of various w/v concentrations of OP50 *Escherichia coli* diluted in water. For these assays worms were grown on dauer agar plates [33], on which the *E. coli* food source...
cannot grow. After transfer to food, worms were allowed to develop for three days, by which time egg-laying had started, worms were then hypochlorite treated to isolate eggs [32]. The lifetime fecundity and reproductive timing of the arrested L1s that developed from these eggs was then determined by transferring worms individually to NGM agar plates with an excess of OP50 and allowing them to develop to adulthood. Once reproduction had started, worms were transferred to fresh plates daily until reproduction had ceased, with the number of larvae that developed on the plates from which the progeny had been removed counted to determine daily and lifetime fecundity [22]. Note that progeny spent the time from fertilization to egg isolation, estimated as a maximum of 2–3 hours, in the reproductive tract of mothers in differing environments. All data were analyzed using Minitab® Statistical Software (Minitab Ltd., Coventry). Lifetime fecundities were Johnson transformed and analyzed by ANOVA, with parental food concentration fitted as a factor. Progeny reproductive schedules were analyzed by comparing the daily fecundities using Kruskal-Wallis tests. Note that direct comparisons of specific % w/v food concentrations between experiments is not possible as the per capita food availability varied between experiments (i.e. the number of mothers per plate varied) and there are differences between batches of E. coli that affect worm life history.

Maternal size and progeny reproduction

Worms from a single cohort of arrested L1s (grandmothers) were fed on two consecutive days and each population allowed to develop for three days on NGM plates with excess food. Eggs were then isolated and progeny allowed to arrest as L1s. These L1s (mothers) were then transferred to dauer agar plates [33] containing 100 μL food patches of various % w/v concentrations of OP50 E. coli diluted in water and allowed to develop until they were four or five days old (the staggered feeding of the grandmothers allowed both four and five day old mothers to be analyzed on the same day). To account for variation between batches of E. coli and for variation in the per capita food availability, maternal food treatments were selected at the point of progeny isolation from a wider range of food concentrations. The highest food concentration used in the assay was defined as High maternal food availability, and excess food remained on the plates in this treatment. Low maternal food availability was selected as the lowest food concentration from which sufficient numbers of progeny could be isolated and on these plates the bacterial food had always been depleted. An intermediate concentration, where small quantities of food remained on the plates, was then selected to represent Medium maternal food availability. Assessment and classification of maternal food treatments therefore occurred prior to progeny isolation and analysis. Once food treatments had been selected, mothers were individually transferred to fresh dauer agar plates without food, photographed using a Moticam 2000 video camera (Motic, Wetzlar, Germany) and parental body size, the length from the mouth to the base of the tail, determined in ImageJ [34]. Mothers were then transferred to a drop of hypochlorite solution to isolate fertilized in utero eggs (progeny), which were allowed to hatch and arrest as L1s. Progeny therefore differed in maternal age (4 day old vs. 5 day old mothers), maternal food availability (defined as High, Medium and Low, as described above) and in the length of time their grandmothers had spent in L1 arrest (1 vs. 2 days). As above, progeny also spent the time from fertilization to egg isolation in different environments. Lifetime fecundity and reproductive timing were then determined as described above. Here, High maternal food availability was 20% w/v E. coli for both the four and five day old mothers, Low maternal food availability was 1.25% and 2.5% w/v E. coli for four and five day old worms, respectively, and Medium maternal food availability was 2.5% and 5% w/v E. coli for four and five day old worms, respectively. Note that the differences between food concentrations for the four and five day old worms are a consequence of the additional day that the five day old mothers spent on the plates.

Maternal body size was analyzed by ANOVA, with Age and Food and the Age by Food interaction tested. Progeny lifetime fecundities were Johnson transformed and analyzed by ANOVA, with maternal Body Size fitted as a covariate, and maternal Age and maternal Food fitted as factors, with maternal Food nested within maternal Age. The relationship between maternal body size and progeny reproduction was investigated by correlating parental size with the mean number of progeny produced on the first day of reproduction and the mean progeny lifetime fecundity.

Maternal growth temperature and progeny reproduction

Arrested L1s were fed on two consecutive days, with worms fed on day 1 maintained at 15°C and those fed on day 2 maintained at 20°C, and allowed to develop on NGM plates with excess food. Progeny were isolated and analyzed, as previously described, from all worms three days after the second batch of arrested L1s were fed (the first day of reproduction for worms grown at 15°C and 20°C) and progeny therefore differed in the maternal growth temperature and in the duration of maternal L1 arrest (1 vs. 2 days). Data were analyzed as described above for Maternal food availability and progeny reproduction.

Maternal heat stress and progeny reproduction

Arrested L1s were transferred to NGM plates with excess food at 20°C and allowed to develop for 24 hours. Worms to be heat shocked were then transferred to at 35°C for 2 hours before being returned to 20°C, a treatment that is sufficient to significantly decrease lifetime fecundity but not to cause immediate mortality (data not shown). Control and heat shock worms were then allowed to develop for a further two days before progeny were isolated and analyzed as previously described. Progeny therefore differed in maternal heat shock exposure. Data were analyzed as described above for Maternal food availability and progeny reproduction.

Maternal food availability and progeny dauer larve development

Worms from a single cohort of arrested L1s were allowed to develop at a range of food concentrations until they were four days old at which point High, Medium and Low food maternal concentrations were selected as described above. Eggs were then isolated by hypochlorite treatment [32] and progeny allowed to arrest as L1s. These arrested L1s were transferred to 3.5 cm diameter plates (40–50 L1s/plate) containing 2 mL of dauer agar [33], 60 μL of dauer pheromone extract [19] and 20 μL of 2% OP50 E. coli. Plates were incubated at 25°C for two days and the proportion of dauer larvae on each plate determined. This assay was repeated three times. In all cases High maternal food availability was 10% w/v E. coli. Medium maternal food availability was 2.5% w/v E. coli for assays 1 and 3, and a combination of 2.5% and 5% w/v E. coli treatments for assay 2. Low maternal food availability was 1.25% w/v E. coli for assays 2 and 3, and a combination of 1.25% and 0.625% w/v E. coli treatments for assay 2. The proportions of progeny that developed as dauer larvae were analyzed by Kruskal-Wallis test.
Progeny reproduction in stressful conditions

Dauer and non-dauer larvae were recovered from a dauer larvae assay (set up as described above). The lifetime fecundity and reproductive timing of these worms, was then determined, as described above, except that worms were maintained on dauer agar plates with 50 µl of 1% OP50 E. coli and maintained at 25°C. While this represents a smaller amount of food than that found on an NGM plate, it is still an excess of food for a single worm maintained on the plate for one day and other work has shown no differences between the lifetime fecundities or reproductive schedules of worms grown in these conditions in comparison to those grown on NGM plates (data not shown). Lifetime fecundity and reproductive timing of control worms, from the same cohorts of arrested L1s used in the dauer assay, was also determined on those grown on NGM plates (data not shown). Lifetime fecundity and 1.25% w/v E. coli treatments for assay 3. Egg size data were not normally distributed and were therefore analyzed by Mann Whitney U test.

Author Contributions

Conceived and designed the experiments: SCH HO. Performed the experiments: SCH HO. Analyzed the data: SCH. Wrote the paper: SCH.

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Egg size

Worms, allowed to develop from L1 arrest at a range of food concentrations, High and Low food concentrations were then selected as described above and worms were then hypochlorite treated to isolate eggs [32]. Eggs were then transferred to dauer agar plates without food, photographed using a Moticam 2000 video camera (Motic, Wetzlar, Germany) and the cross sectional area determined in ImageJ [34]. This assay was repeated three times. In all cases High maternal food availability was 10% w/v E. coli. Low maternal food availability was a combination of 2.5% and 0.125% w/v E. coli for assay 1, 0.3125% w/v E. coli for assay 2 and 1.25% w/v E. coli treatments for assay 3. Egg size data were not normally distributed and were therefore analyzed by Mann Whitney U test.

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