The tricarboxylic acid (TCA) cycle (or citric acid cycle) is responsible for the complete oxidation of acetyl-CoA and formation of intermediates required for ATP production and other anabolic pathways, such as amino acid synthesis. Here, we uncovered an additional mechanism that may help explain the essential role of the TCA cycle in the early embryogenesis of Caenorhabditis elegans. We found that knockdown of citrate synthase (cts-1), the initial and rate-limiting enzyme of the TCA cycle, results in early embryonic arrest, but that this phenotype is not because of ATP and amino acid depletions. As a possible alternative mechanism explaining this developmental deficiency, we observed that cts-1 RNAi embryos had elevated levels of intracellular acetyl-CoA, the starting metabolite of the TCA cycle. Of note, we further discovered that these embryos exhibit hyperacetylation of mitochondrial proteins. We found that supplementation with acetylase-inhibiting polyamines, including spermidine and putrescine, counteracted the protein hyperacetylation and developmental arrest in the cts-1 RNAi embryos. Contrary to the hypothesis that spermidine acts as an acetyl sink for elevated acetyl-CoA, the levels of three forms of spermidine and putrescine, counteracted the protein hyperacetylation and developmental arrest in the cts-1 RNAi embryos. We found that supplementation with acetylase-inhibiting polyamines, including spermidine and putrescine, counteracted the protein hyperacetylation and developmental arrest in the cts-1 RNAi embryos. Instead, we demonstrated that the mitochondrial deacetylase sirtuin 4 (encoded by the sir-2.2 gene) is required for spermidine's suppression of protein hyperacetylation and developmental arrest in the cts-1 RNAi embryos. Taken together, these results suggest the possibility that during early embryogenesis, acetyl-CoA consumption by the TCA cycle in C. elegans prevents protein hyperacetylation and thereby protects mitochondrial function.
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**Figure 1. Citrate is decreased from the early to late embryonic stage in C. elegans.** A, the relative values of the TCA cycle metabolites between the early and late embryonic stages. The levels of each TCA cycle metabolite were normalized to the amounts of total TCA cycle metabolites at each developmental stage and indicated as relative value (early embryo versus late embryo). Data are mean ± S.E.; n = 5; *, p < 0.05; N.S., not significant. α-KG indicates α-ketoglutarate. B, the levels of the TCA metabolites in mock and cts-1 RNAi embryos with CE-TOF/MS. Data are mean ± S.E.; n = 3; **, p < 0.01; ***, p < 0.001; N.S., not significant.

thought to be largely elicited by nonenzymatic reaction of reactive lysine residues with acetyl-CoA and removed by the mitochondrial NAD⁺-dependent deacetylase SIRT3 (8–11). It should be noted that protein acetylation in mitochondria typically results in dysfunction of various major aspects of mitochondrial biology.

It was previously reported that acetyl-CoA levels were elevated and proteins were acetylated in mitochondria when fuel was increased (12). Furthermore, current evidence suggests that acetyl-CoA is a key factor determining the levels of protein acetylation during early embryonic development in Xenopus laevis (13). Given that elevated acetyl-CoA levels could induce protein lysine acetylation in mitochondria, we hypothesized that the consumption of acetyl-CoA through the TCA cycle is an indispensable process for maintaining normal mitochondrial function. Here we provide a functional relationship between the TCA cycle and mitochondrial protein hyperacetylation during early embryonic development in C. elegans.

**Results**

**Citrate is consumed predominantly from the early to late embryonic stage in C. elegans**

To understand the role of the TCA cycle during embryogenesis, we first measured the levels of metabolites of the TCA cycle at both the early and late embryonic stages in C. elegans. LC-MS analysis showed that the relative ratio of citrate, but not other metabolites of the TCA cycle, is significantly decreased in the late embryos (Fig. 1A). Given that citrate is consumed and then regenerated by the sequence of reactions to complete the TCA cycle, the reduction of citrate at the late embryonic stage implies a contribution of the TCA cycle to the process of embryogenesis. To investigate whether the consumption of citrate is an essential process involved in embryogenesis, we focused on citrate synthase encoded by cts-1 gene that is the initial and rate-limiting enzyme of the TCA cycle. As shown in Fig. 1B, the levels of all TCA metabolites in early embryos were markedly decreased when cts-1 was silenced by feeding RNAi. Thus, cts-1 knockdown approach has allowed us to clarify the role of citrate consumption in early embryogenesis.

**A reduction in the levels of glutamate, aspartate, and citrate is not a cause of early embryonic arrest by cts-1 knockdown**

Several amino acids are synthesized from TCA metabolites, including α-ketoglutarate and oxaloacetate. Thus, homeostatic imbalance of amino acids could be a possible explanation for early embryonic arrest by cts-1 knockdown. To test this hypothesis, we quantified the relative amounts of amino acids using CE-TOF/MS and found that glutamate and aspartate, which are derived from α-ketoglutarate and oxaloacetate, respectively, were decreased in cts-1 RNAi embryos as compared with mock control (Fig. S1, B and C). Next, to determine whether reduction of glutamate and aspartate is sufficient to account for early embryonic arrest by cts-1 knockdown, we supplemented cts-1 RNAi embryos with an excess dose (30 mM) of glutamate and aspartate and examined hatching rate. Although the levels of glutamate and aspartate in cts-1 RNAi embryos were completely restored by the supplementation, this treatment failed to rescue the reduced hatching rate even when both glutamate and aspartate were added (Fig. 2, D–F). These observations

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**cts-1 knockdown causes developmental arrest at the early embryonic stage without lack of intracellular ATP levels**

The TCA cycle is an oxidative device coupled to the electron transport chain for the production of ATP and the genes responsible for the electron transport chain are known to be required for early embryogenesis (4). Because cts-1 knockdown led to dysfunction of the TCA cycle, we hypothesized that cts-1 RNAi animals exhibit early developmental defects. As expected, cts-1 knockdown dramatically reduced hatching rate and also caused developmental arrest at the early embryonic stage (Fig. 2, A and B). Next, to test whether these phenotypes result from a failure of ATP production by cts-1 knockdown, we measured ATP levels in mock or cts-1 RNAi embryos using CE-TOF/MS. Surprisingly, however, the ATP levels of cts-1 RNAi embryos were comparable with those of mock RNAi embryos (Fig. 2C). Similar result was obtained using the luciferase-luciferin system (Fig. S1A). These data suggest that energy shortage is not a cause of early embryonic arrest in cts-1 RNAi animals.

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Figure 2. cts-1 knockdown results in early embryonic arrest without depletion of intracellular ATP levels. A, the hatching rate in mock and cts-1 RNAi embryos (data are mean ± S.E.; n = 3; ***,$p < 0.001$). B, percentage of animals at each embryonic stage in cts-1 RNAi embryos (data are mean ± S.E.; *, $p < 0.05$). C, ATP levels in mock and cts-1 RNAi embryos with CE-TOF/MS (N.S., not significant). D, the levels of glutamate in mock RNAi embryos with vehicle (−), cts-1 RNAi embryos with vehicle (−), and 30 mM glutamate (Glu) with LC-MS/MS (data are mean ± S.E.; n = 3; N.S., not significant). E, the levels of aspartate in mock RNAi embryos with vehicle (−), cts-1 RNAi embryos with vehicle (−), and 30 mM aspartate (Asp) with LC-MS/MS (data are mean ± S.E.; n = 3; N.S., not significant). F, the hatching rate of mock RNAi embryos with vehicle (−), cts-1 RNAi embryos with vehicle (−), and 30 mM Glu and/or Asp (data are mean ± S.E.; n = 3; ***,$p < 0.001$). G and H, citrate levels (G) and the hatching rate (H) in mock RNAi embryos with vehicle (−) and cts-1 RNAi embryos with vehicle (−), and 30 mM citrate (data are mean ± S.E.; n = 3; ***, $p < 0.001$; N.S., not significant).

Implies that decreased levels of glutamate and aspartate do not account for early embryonic arrest by cts-1 knockdown.

Alternatively, because it has been reported that citrate allosterically modulates metabolic enzymes, such as phosphofructokinase, pyruvate kinase, pyruvate dehydrogenase, and acetyl-CoA carboxylase, thereby controlling glycolysis and fatty acid synthesis (14), we assessed a direct effect of citrate depletion on the embryogenesis by adding back citrate to cts-1 RNAi embryos. However, consistent with the results of amino acid supplementation (Fig. 2, D–F), addition of citrate had no effect on the reduced hatching rate in cts-1 RNAi embryos, despite the fact that citrate levels were recovered to those in mock RNAi embryos (Fig. 2, G and H). Taken together, these results suggest that early embryonic arrest by cts-1 knockdown is not attributed to reduced levels of glutamate, aspartate, and citrate in C. elegans.

crs-1 RNAi embryos show an increase in the levels of acetyl-CoA and acetylation of mitochondrial proteins

As referred to above, our results demonstrated that at least production of ATP, amino acids, and citrate through the TCA cycle does not play a critical role in the hatching process of C. elegans. If so, what is the role of the TCA cycle in early embryogenesis? To find a clue to address this question, we performed LC-MS/MS analysis with mock and cts-1 RNAi embryos and found that acetyl-CoA, the starting point for the TCA cycle, is significantly increased by cts-1 knockdown (Fig. 3A). This finding raised the possible involvement of acetyl-CoA...
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Figure 3. cts-1 knockdown causes an increase in acetyl-CoA levels and mitochondrial protein hyperacetylation in early embryos. A, the levels of acetyl-CoA in mock and cts-1 RNAi embryos with LC-MS/MS. Data are mean ± S.E.; n = 3; *, p < 0.03. B, left, acetylated proteins of mock RNAi embryos and cts-1 RNAi embryos (AcK, acetylated proteins; CTS-1, citrate synthase-1). Right, the levels of total acetylated proteins larger than 28 kDa in mock and cts-1 RNAi embryos (data are mean ± S.E.; n = 3; ***, p < 0.001). C, acetylated proteins of total lysate (TL), post-mitochondrial supernatant (PMS), or mitochondrial pellets (MP) in mock and cts-1 RNAi embryos (ATPB, ATP synthase β; H3, histone H3).

Spermidine does not act as a sink for the elevated acetyl-CoA levels even when treated with high dose

Although the rescue effect of polyamine in embryogenesis would be achieved through inhibition of hyperacetylation because of overproduction of acetyl-CoA, the detailed mechanism how polyamine reduces protein acetylation remains unclear. Meanwhile, we noticed key findings from the CE-TOF/MS data demonstrating that cts-1 knockdown augments the levels of acetylated polyamines, namely, acetylpolyamine and acetylperimidine (Fig. 5A). These emerging data prompted us to consider that excess concentrations of polyamines might be sinks for the elevated acetyl-CoA levels as acetylation acceptors, thereby siphoning off the amount of acetyl-CoA for protein acetylation. To test this hypothesis, we set up to determine the levels of acetylspermidine in mock and cts-1 RNAi embryos treated with or without spermidine. Because amino groups of spermidine are acceptable for acetylation at both its ends (Fig. 5B), the two distinct mono- (N1- and N8-acetylperimidine) and the diacetylated (N4,N8-diacetylperimidine) forms of spermidine were measured specifically using LC-MS/MS. Unexpectedly, however, we found that the levels of all forms of acetylspermidine in cts-1 RNAi embryos are not significantly increased when treated with exogenous spermidine (Fig. 5, C–E). A similar result was observed in the levels of acetylpolyamine (data not shown). These data indicate that polyamine-mediated inhibition of protein hyperacetylation is not attributed to the function of spermidine as a sink for acetyl groups under overproduction of acetyl-CoA.

sir-2.2 is required for recovery effect of spermidine on protein hyperacetylation and developmental arrest in cts-1 RNAi embryos

An alternative possible mechanism for polyamine-mediated inhibition of hyperacetylation could be the activation of protein deacetylase. Considering that polyamine treatment inhibited protein hyperacetylation occurring primarily in mitochondria, potential targets of polyamine should be the mitochondrial deacetylases, including three sirtuin isoforms, SIRT3, SIRT4, and SIRT5. Among them, mammalian SIRT3 is the major protein deacetylase in mitochondria (10), but there are no genes with similarity to SIRT3 and also SIRT5 in C. elegans. Instead,
C. elegans possesses two genes, sir-2.2 and sir-2.3, both of which are orthologs of SIRT4 and their products are localized in mitochondria (17). However, because C. elegans SIR-2.2 and SIR-2.3 proteins have not been demonstrated to catalyze deacetylation on a variety of substrates, we first examined broad deacetylase activity of SIR-2.2 in mock and cts-1 RNAi embryos. As shown in Fig. 6A, we found that although endogenous protein acetylation levels of early embryos are comparable between WT N2 and sir-2.2(tm2648) mutant animals, cts-1 knockdown-induced hyperacetylation of proteins larger than 28 kDa is slightly augmented in sir-2.2(tm2648) mutant, suggesting that SIR-2.2 contributes to protein deacetylation, at least, when acetyl-CoA levels are elevated.

We next investigated the role of sir-2.2 in spermidine-mediated inhibition of protein hyperacetylation under cts-1 knockdown condition. Notably, the inhibitory effect of spermidine on protein hyperacetylation in cts-1 RNAi embryos was completely abolished by deletion of sir-2.2 gene (Fig. 6B). In contrast, spermidine treatment substantially decreased the cts-1 knockdown-induced hyperacetylation in sir-2.3(ok444) mutants (Fig. 6C), indicating that not sir-2.3, but sir-2.2 is involved in the spermidine-mediated inhibition of protein hyperacetylation. Finally, we examined hatching rate of WT N2 and deletion mutants of sir-2.1 (the SIRT1 ortholog), sir-2.2, and sir-2.3 to ascertain whether sir-2.2 indeed plays a role in the spermidine-mediated recovery of embryonic development. As expected, the reduced hatching rates by cts-1 knockdown were observed in all strains tested, whereas the recovery effect of spermidine was partially abrogated in the sir-2.2(tm2468) mutant compared with N2 and other mutants (Fig. 6D). Taken together, these results imply that SIR-2.2 is a target of spermidine for preventing hyperacetylation when the TCA cycle is impaired during early embryogenesis.

Discussion

The fundamental question in this study is what the role of the TCA cycle during embryogenesis in C. elegans is. We demonstrated that the TCA cycle is indispensable for early embryonic development by RNAi depletion of CTS-1 that functions as the initial and rate-limiting enzyme of the TCA cycle (Fig. 1, A and B). Although ATP deprivation appeared to be a cause of early developmental arrest in cts-1 RNAi embryos, there was no difference in intracellular ATP levels between control and cts-1 RNAi embryos (Figs. 1C and S1A). This finding indicates that intracellular ATP levels are sustained during early embryogenesis even without ATP production through the TCA cycle. If so, what pathways are used for supplying ATP? We consider two possibilities: (1) through glycolysis and (2) through amino acid metabolism. To address this issue, we treated early embryos with 2-deoxyglucose, a glycolysis inhibitor, but no decrease was observed in ATP levels (data not shown), suggesting that glycolysis is not likely to participate in ATP supply during early embryogenesis. Another possibility is the use of amino acid as a substitute for acetyl-CoA to produce ATP. An interesting model has been proposed in which the TCA cycle reactions are split into complementary mini-cycles: namely, the first mini-cycle converts α-ketoglutarate to oxaloacetate and the second mini-cycle converts oxaloacetate to α-ketoglutarate (18). Based on this model, the first mini-cycle would allow metabolizing glutamate for producing ATP even when the second mini-cycle does not function by cts-1 knockdown. Supporting this notion, we found that the levels of glutamate are considerably decreased in cts-1 RNAi embryos (Fig. S1B). Instead, the existence of multiple metabolic pathways to ensure the level of intracellular ATP implies other roles of the TCA cycle in early embryonic development.

Here, we demonstrated the relationship between the TCA cycle and protein/metabolite acetylation through acetyl-CoA. The primary function of the TCA cycle is to release stored energy through the oxidation of acetyl-CoA into carbon dioxide and chemical energy in the form of ATP. As mentioned above, however, we revealed that ATP levels are not influenced by cts-1 knockdown and rather successive consumption of acetyl-CoA via the TCA cycle plays a crucial role in preventing accumulation of acetyl-CoA in mitochondria (Fig. 3A). Given that mitochondrial proteins possessing intrinsic acetyltransferase activity have not been identified so far (9), excessive accumulation of acetyl-CoA exclusively in mitochondria seems to cause aberrant acetylation of proteins and metabolite through
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the nonenzymatic reaction, thereby resulting in mitochondrial dysfunction related to early embryonic arrest. Moreover, it is noted that a large number of mitochondrial proteins involved in metabolic pathways have been shown to be regulated by reversible acetylation (7). Taken together, we suggest the notion that homeostatic modulation of mitochondrial protein acetylation could be an alternative function of the TCA cycle at least during embryogenesis. Meanwhile, because blocking citrate synthase would have a number of effects beyond producing an excess amount of acetyl-CoA, it is important to provide evidence that increasing acetylation of mitochondrial proteins by a method except cts-1 knockdown also leads to mitochondrial dysfunction and early embryonic lethality.

To prevent hyperacetylation of mitochondrial proteins, we used natural polyamines, such as spermidine, which has been reported to inhibit histone acetyltransferases and thereby promote longevity through up-regulation of autophagy-related transcripts (15). In addition to the fact that a definitive mitochondrial acetyltransferase has not been identified, our findings from metabolome analysis led to the hypothesis that excess concentrations of polyamines might be sinks for the elevated acetyl-CoA levels by being acetylated themselves and thus siphon off the amount of acetyl-CoA for protein acetylation. However, because no significant increase in the levels of acetyl forms of spermidine, including N¹-acetylspermidine, N⁸-acetylspermidine, and N¹,N⁸-diacetylspermidine, was observed even when the TCA cycle was compromised by cts-1 knockdown (Fig. 5), this hypothesis appears to be incorrect at least in our experimental conditions. Instead, we discovered that the nematode mitochondrial sirtuin, SIR-2.2, has deacetylase activity, albeit at a lower level, and could be a target of spermidine-mediated inhibition of hyperacetylation (Fig. 6). Although we are unable to uncover the molecular mechanism underlying the action of polyamine on SIR-2.2 function, our findings provide

![Figure 5. Spermidine does not act as an acetyl sink even when treated with high-dose.](image-url)
new insight into the physiological significance of polyamine as an acetylation inhibitor.

Interestingly, it was reported that \(N_1,N_8\)-diacetylspermidine is a potential cancer marker for early detection because its levels in the urine from patients are increased along with a progression of cancer (20). However, to date, there is no report describing the existence of \(N_1,N_8\)-diacetylspermidine in animals other than human and it remains unknown how \(N_1,N_8\)-diacetylspermidine is produced in cancer cells. In this study, we first discovered the endogenous production of \(N_1,N_8\)-diacetylspermidine in nematode when the TCA cycle is compromised by \(cts-1\) knockdown (Fig. 5E). Considering that \(N_1,N_8\)-diacetylspermidine is predicted to arise simultaneously with the accumulation of mitochondrial acetyl-CoA, this diacetylpolyamine could be a biomarker that identifies the metabolic balance between glycolysis and the TCA cycle.

Our present study revealed a possible link among the TCA cycle, acetylation, and polyamine at the stage of early embryonic development in \(C.\ elegans\). On the other hand, it is likely that this mutual relationship plays an important role in various
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biological phenomena throughout life. In particular, considering that spermidine is a longevity drug that can increase life span in yeast, nematodes, and flies (15), the acetylation status mediated by the TCA cycle is emerging as a promising regulatory mode of chronological aging. Consistent with this notion, a glucose-enriched diet has been shown to shorten life span by down-regulating the DAF-16 and HSF-1 transcription factors in *C. elegans* (21). Further investigations of the relationship between the TCA cycle and protein/metabolite acetylation in view of metabolic change with age will shed light on how spermidine increases longevity and also contributes to identify therapies to slow down aging and delay age-associated diseases.

**Experimental procedures**

**Strains and the maintenance**

Bristol N2 WT, *sir-2.1(ok434)*, and *sir-2.3(ok444)* were provided by the Caenorhabditis Genetics Center (CGC). *sir-2.2(tm2468)* was obtained from National BioResource Project (NBRP). All mutants were backcrossed five times to N2. All strains were maintained on Nematode Growth Medium plates at 20 °C.

**Collection of animals**

To suppress the loss of the collected embryos and metabolites through the nonspecific adsorption with tubes and tips, we used platinum tubes and tips (BM-Bio Inc.) in all experiments.

**Reagents and antibodies**

Reagents were purchased from Sigma-Aldrich, Wako, Tokyo-INC, Tokyo Chemical Industry, Cambridge Isotope Laboratories, Inc., and Cayman Chemical Co. Antibodies were purchased from Santa Cruz Biotechnology, Cell Signaling Technology, MBL International, and Abcam. *N*<sup>1</sup>,*N*<sup>8</sup>-diacetylsperrmidine was chemically synthesized and purified. The detailed information on reagents and antibodies is shown in Table S1.

**Sample preparation for metabolite measurement**

Early and late embryos were collected immediately and 11–12 h after bleaching of 5000 gravid adults, respectively. Embryos of mock RNAi or *cts-1* RNAi were collected after thorough bleaching ~3000 or 4000 gravid adults, respectively. The embryonic pellets were boiled in 50 μl of 1 × SDS-PAGE sample buffer (50 mM Tris-HCl, pH 7.5, 1.8% mercaptoethanol, 1% SDS, 0.01% bromphenol blue, 10% glycerol). Then, 5–10 μl of the lysates were separated on 15% SDS-PAGE, and then the proteins were transferred to PVDF membrane. The membrane was treated with the blocking buffer, and then incubated with primary antibodies followed by HRP-conjugated secondary antibodies (Table S1). Densitometric analyses of anti–acetylated-lysine Western blotting bands were performed by ImageJ software. Briefly, areas of Western blot bands larger than 28 kDa were framed as acetylated protein signals and then converted into gray values. After subtracting background values, the intensity profiles were determined and presented as a relative ratio. The acetylated protein signals larger than 28 kDa were divided into three sizes, namely: upper (> 75 kDa), middle (48 –75 kDa), and lower (28 –48 kDa) and then the linearity of signal intensities was validated by quantification of densitometric values of each area around 3-s exposure.

**Measurement of hatching rate**

Hatching rate was measured in reference to previous research (3). In brief, early L4 animals were bred on RNAi plates for 45 h, and then 6 –8 gravid adults were transferred to new RNAi plates. These animals were removed 6 h thereafter, and then hatchlings and eggs were counted 24 h later.

**Mitochondria/cytosol fractionation**

In this fractionation, we referred to the protocol shown by Jonassen *et al.* (19). Large numbers of early embryos were collected by bleaching 35,000 gravid adults. The embryos were washed four times in M9 buffer, then resuspended in 2 ml of isolation buffer (IB) (210 mM mannitol, 70 mM sucrose, 0.1 mM EDTA, 5 mM Tris, pH 7.4, 1 mM PMSF). The samples were kept cold on ice throughout the fractionation procedure. The ani-
mals were homogenized with 15 up-and-down strokes with a Dounce homogenizer. The homogenates were centrifuged at 1000 × g for 10 min. The supernatants were saved, and 2 ml of IB was further added to the pellets. The pellets were suspended and homogenized again with additional 15 strokes, and centrifuged at 1000 × g for 10 min. An aliquot of the combined supernatants was saved as total lysate (TL). The supernatants were centrifuged at 5000 × g for 10 min. The supernatants were saved with IB, and then centrifuged at 2000 × g for 10 min. An aliquot of the combined supernatants was centrifuged at 14,000 × g for 10 min, and the mitochondrial pellet (MP) was obtained. The supernatants centrifuged at 1000 × g for 10 min. An aliquot of the resulting supernatant was saved as post-mitochondrial supernatant (PMS). Protein pellets of TL and PMS were collected by chloroform precipitation. SDS sample buffer was added to all pellets of TL, PMS, and MP.

Statistical analysis

Results were presented as mean ± S.E. Statistical significances were determined by two-tailed Student’s t test using GraphPad Prism software. A p value < 0.05 was considered statistically significant.

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