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# Nutritional vitamin B12 regulates RAS/ MAPK-mediated cell fate decisions through one-carbon metabolism

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Vitamin B12 is an essential nutritional co-factor for the folate and methionine cycles, which together constitute one-carbon metabolism. Here, we show that dietary uptake of vitamin B12 modulates cell fate decisions controlled by the conserved RAS/MAPK signaling pathway in *C. elegans*. A bacterial diet rich in vitamin B12 increases vulval induction, germ cell apoptosis and oocyte differentiation. These effects are mediated by different one-carbon metabolites in a tissue-specific manner. Vitamin B12 enhances via the choline/phosphatidylcholine metabolism vulval induction by down-regulating fat biosynthesis genes and increasing H3K4 tri-methylation, which results in increased expression of RAS/MAPK target genes. Furthermore, the nucleoside metabolism and H3K4 tri-methylation positively regulate germ cell apoptosis and oocyte production. Using mammalian cells carrying different activated KRAS and BRAF alleles, we show that the effects of methionine on RAS/MAPK-regulated phenotype are conserved in mammals. Our findings suggest that the vitamin B12-dependent one-carbon metabolism is a limiting factor for diverse RAS/MAPK-induced cellular responses.

The cell metabolism regulates physiological processes and cancer cell growth by controlling catabolic and anabolic reactions and transcriptional activity<sup>1</sup>. Diet influences metabolism directly and, as a result, cells must constantly adapt to variations in diet. One example is vitamin B12 (cobalamin), an essential micronutrient that requires dietary intake. Vitamin B12 is an important metabolic co-factor in two metabolic pathways, the mitochondrial propionate breakdown pathway and one-carbon metabolism formed by the methionine and folate cycles (Fig. 1a). In the canonical propionate breakdown pathway, vitamin B12 acts as a co-factor for methyl malonyl-CoA mutase (MUT, MMCM-1 in *C. elegans*)<sup>2–4</sup>, and in the methionine cycle, it serves as a co-factor for methionine synthase (MS, METR-1 in *C. elegans*) (Fig. 1a)<sup>2–4</sup>.

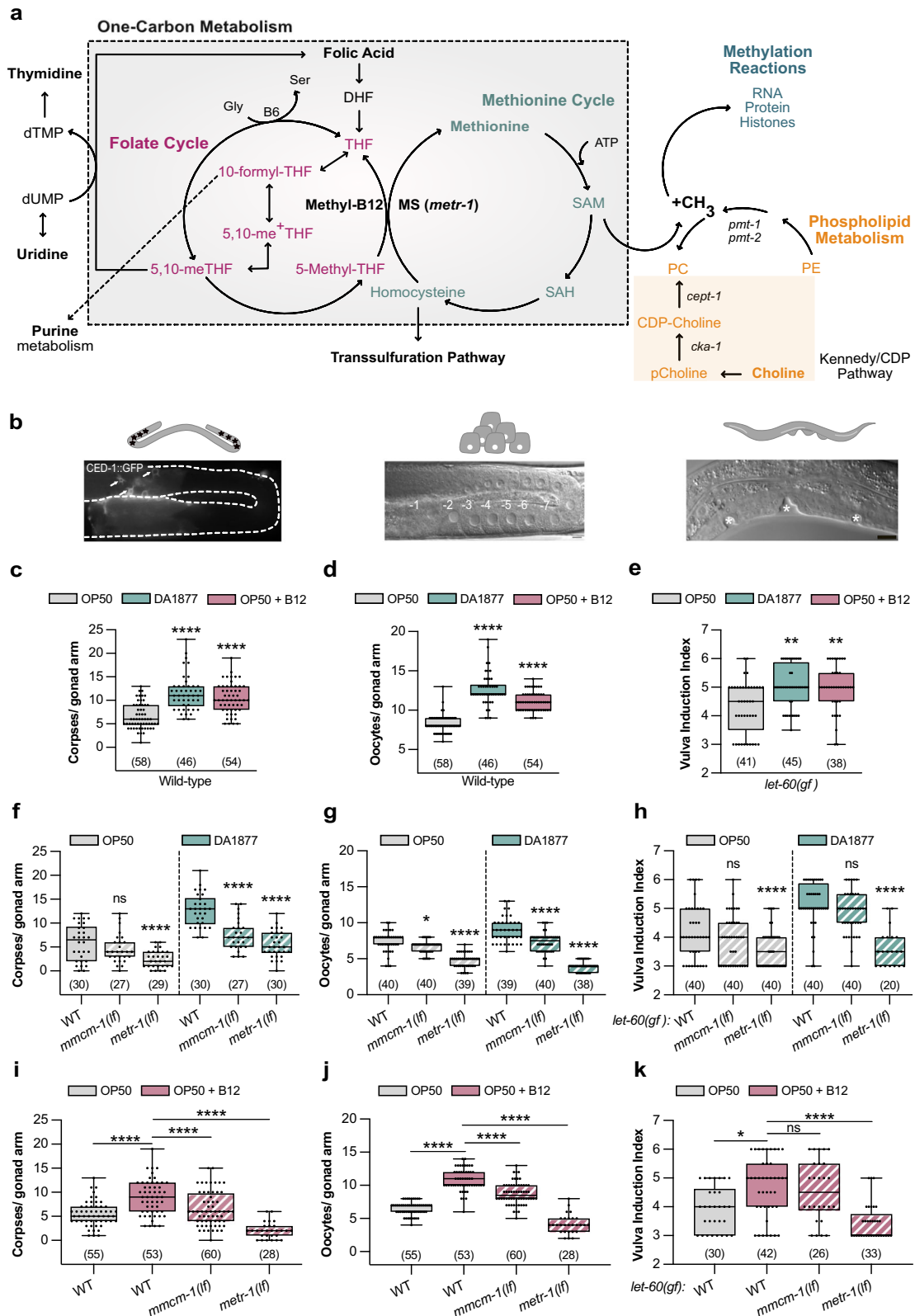
One-carbon metabolism transfers single carbon units necessary for nucleotide biosynthesis through the folate cycle and produces S-adenosyl-methionine (SAM), the main methyl donor in cells. In the methionine cycle, homocysteine is methylated by MS to form

methionine, which is then converted into SAM<sup>5,6</sup>. SAM is required for phosphatidylcholine (PC) synthesis (Fig. 1a)<sup>5</sup>. PC is a polyunsaturated fatty acid that can be produced via the Kennedy/CDP-choline pathway using choline as a precursor or by the sequential methylation pathway, which depends on SAM-methyl groups (Fig. 1a)<sup>7,8</sup>. In addition, SAM provides the methyl donors necessary for DNA, RNA, histone and protein methylation. Histone methylation, especially histone H3 tri-methylation at lysine 4 (H3K4me3), which is associated with active transcription at the promoter region, is particularly sensitive to SAM levels<sup>9,10</sup>.

In this study, we have used the nematode *C. elegans* to investigate the nutritional impact of a bacterial diet on development. *E. coli* OP50 is the commonly used bacterial diet, which is low in vitamin B12. *Comamonas aquatica* DA1877, on the other hand, is a vitamin B12-rich diet<sup>3</sup>. A *Comamonas* DA1877 diet affects different aspects of *C. elegans* behavior by modulating gene expression, reproduction, and

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**Fig. 1 | The *Comamonas* DA1877 diet promotes VPC and germ cell differentiation through the vitamin B12 metabolism. **a** Schematic representation of one-carbon metabolism. **b** Illustrations of the observed phenotypes: germ cell apoptosis (corpses), oocyte differentiation, or vulval induction. Scale bar: 10 μm. Number of corpses (**c**) or oocytes (**d**) in wild-type animals. **e** Vulval induction index of *let-60(gf)* animals. **c–e** Animals were fed with *E. coli* OP50 (gray), *Comamonas* DA1877 (blue), or *E. coli* OP50 supplemented with 64 nM B12 (pink). Number of corpses (**f**), oocytes (**g**) and vulval induction index (**h**) for indicated genotypes fed either with *E. coli* OP50 (gray) or *Comamonas* DA1877 (blue). Number of corpses (**i**), oocytes (**j**)**

and vulval induction index (**k**) for indicated genotypes fed with *E. coli* OP50 and supplemented with 64 nM B12. Dots (in **c**, **d**, **f**, **g**, **i**, **j**) represent individual corpses/oocytes or animals (in **e**, **h**, **k**) from two independent biological replicates; number of animals in brackets. Boxplots show 25-75 percentiles and median; whiskers represent min and max values. \*\**P* < 0.005, \*\*\**P* < 0.0005, \*\*\*\**P* < 0.0001 using one-way ANOVA followed by Dunnett’s multiple comparison tests in **c**, **d**, **i** and **j**, two-way ANOVA followed by a Tukey’s multiple comparison test in **f**, **g** and **h** and a Kruskal-Wallis test for non-parametric data followed by a Dunn’s multiple comparison test in **e** and **k**. Source data are provided as a Source Data file.

longevity<sup>11,12</sup>. Vitamin B12 metabolism also regulates H3K4 methylation<sup>10</sup>, affecting *C. elegans* fertility and lipid accumulation, which is linked to longevity<sup>13,14</sup>.

Here, we focused on how the vitamin B12-dependent one-carbon metabolism modulates cell fate decisions mediated by the conserved RAS/Mitogen-activated protein kinase (RAS/MAPK) pathway. The RAS/MAPK pathway is highly conserved between *C. elegans* and mammals, and it is a prevalent oncogenic signaling pathway in humans, being hyper-activated in around one-third of all tumors. Gain-of-function (*gf*) mutations in RAS genes (mostly in KRAS) occur in 30% of all cancers, and mutations in BRAF (mostly the V600E substitution) are found in 8% of all human cancers<sup>15,16</sup>. Gain-of-function mutations in *let-60*, the single *C. elegans* *ras* ortholog, hyperactivate RAS/MAPK signaling and cause enhanced germ cell death, oocyte over-production, and excess vulval induction<sup>17–19</sup>.

During larval development, the growth factor LIN-3 (an EGF-like ligand) is secreted by the gonadal anchor cell (AC). LIN-3 activates RAS/MAPK signaling via the LET-23 EGF receptor in the vulval precursor cells (VPCs) to induce their differentiation<sup>20,21</sup>. P6.p, the VPC closest to the AC, receives the highest concentration of LIN-3 and adopts the primary (1°) vulval cell fate, while P5.p and P7.p receive less LIN-3 signal and adopt the secondary (2°) cell fate in response to a lateral LIN-12 Notch signal. These three induced VPCs then start to proliferate and differentiate into 22 vulval cells forming the vulva<sup>20,21</sup>.

During adulthood, RAS/MAPK regulates germ cell progression and death in the gonads. Hermaphrodites have two symmetric, U-shaped tubular gonad arms that are connected to a common uterus. In the distal region of each gonad arm, mitotic stem cells constantly proliferate to supply new germ cells. Germ cells in the late pachytene region, near the gonad loop, activate the RAS/MAPK pathway, which is required for the pachytene to diplotene transition, cell membrane integrity, chromosomal synapsis, apoptosis, and oocyte differentiation<sup>17,18,22,23</sup>. Around half of the germ cells enter the proximal gonad arm and begin to differentiate into oocytes, while the remaining germ cells undergo apoptosis. Germ cell apoptosis is a physiological and stochastic process that eliminates half of all germ cells to maintain gonad homeostasis<sup>24,25</sup>. Dying germ cells activate the core apoptotic machinery triggering the CED-3 caspase<sup>24,25</sup> and are engulfed by the sheath cells of the somatic gonad<sup>26</sup>.

By examining three RAS/MAPK-mediated phenotypes, vulval development, germ cell death, and oocyte differentiation, we show that the activity of the vitamin B12-dependent one-carbon metabolism is a limiting factor for RAS/MAPK-induced cell fates. Switching animals to a vitamin B12-rich *Comamonas* DA1877 diet enhances all phenotypes caused by hyperactive RAS/MAPK signaling via the methionine synthetase METR-1. One-carbon metabolism affects the different RAS/MAPK-mediated cell fate decisions through distinct, tissue-specific mechanisms. The folate cycle promotes germ cell differentiation and death through nucleotide biosynthesis, while the methionine cycle affects vulval induction by repressing fatty acid synthesis. In addition, the methionine cycle regulates the PC metabolism and globally increases H3K4me3 levels. Finally, using mammalian cells carrying different activating KRAS alleles, we show that the methionine-dependency of RAS/MAPK-induced phenotypes is conserved.

## Results

### Dietary vitamin B12 enhances germ cell and vulval development

To investigate how the vitamin B12 metabolism affects RAS/MAPK-controlled cell fate decisions, we fed animals with the *Comamonas* DA1877 diet, rich in vitamin B12, and quantified germ cell death, oocyte and VPC differentiation (Fig. 1b). Using a CED-1::GFP reporter to label engulfed apoptotic cells<sup>27</sup>, we found that wild-type worms fed with the *Comamonas* DA1877 diet or *E. coli* OP50 supplemented with vitamin B12 had a 2 to 3-fold increase in apoptotic germ cells (Fig. 1c). Although animals fed with the *Comamonas* DA1877 diet have a smaller brood

size<sup>11</sup>, a B12-rich diet increases the number of oocytes in the proximal arm of one-day-old adults (Fig. 1d).

Since *Comamonas* DA1877-fed worms reach adulthood on average 12 h earlier (Supplementary Fig. 1a)<sup>11</sup>, we tested whether these effects were due to changes in developmental timing. Sixty-two hours after L1 arrest, *Comamonas* DA1877-fed animals were approximately at the same developmental stage as animals fed on *E. coli* OP50 for 72 h. However, *Comamonas* DA1877-fed worms had more apoptotic corpses and oocytes than *E. coli* OP50-fed animals at the matching developmental stages (i.e., 62 h vs. 72 h and 72 h vs. 82 h, Supplementary Fig. 1b, c), suggesting that the *Comamonas* DA1877 effects are independent of developmental time.

Germ cell apoptosis can either be physiological or a response to DNA damage<sup>24,28</sup>. To distinguish between these two possibilities, we used *cep-1* loss-of-function (*lf*) mutants, which do not induce DNA damage-induced apoptosis<sup>27</sup>. The effect of the *Comamonas* DA1877 diet on germ cell death was still present in *cep-1(lf)* mutants (Supplementary Fig. 1d). We also examined *pch-2(lf)* mutants<sup>29</sup> to exclude that the enhanced germ cell apoptosis in *Comamonas* DA1877-fed animals was due to defects in synaptonemal complex assembly (Supplementary Fig. 1e). In both mutant backgrounds fed with the *Comamonas* DA1877 diet, the levels of germ cell death were only slightly decreased compared to the wild-type, suggesting that the *Comamonas* DA1877 diet predominantly induces physiological germ cell death and causes only low levels of genotoxic stress. In contrast, a *ced-3(lf)* mutation completely suppressed the effect of the *Comamonas* DA1877 diet, indicating that the canonical apoptotic caspase pathway is activated by the diet (Supplementary Fig. 1f).

To study the effect of the *Comamonas* DA1877 diet on VPC induction during vulval development, we used a *let-60 ras* gain-of-function mutation (*let-60(n1046)*, abbreviated *let-60(gf)*), as a sensitized genetic background. *let-60(gf)* animals grown on *E. coli* OP50 contained on average  $4.3 \pm 0.9$  (SD) induced VPCs per animal (vulval induction index). *let-60(gf)* larvae fed with *Comamonas* DA1877 or supplemented with vitamin B12 showed an increase in vulval induction to  $5.0 \pm 0.7$  and  $5.0 \pm 0.8$ , respectively (Fig. 1e). Wild-type larvae always had three induced VPCs irrespective of the bacterial diet.

To test if the effect of the *Comamonas* DA1877 diet was caused by bacterially produced B12, we used the B12 synthesis-deficient *Comamonas* DA1877 mutants  $\Delta cbiA/cobB$  and  $\Delta cbiB^3$ . Germ cell apoptosis in animals fed with *Comamonas* DA1877  $\Delta cbiA/cobB$  or  $\Delta cbiB$  mutants was comparable to *E. coli* OP50-fed animals, indicating that the increase in dietary B12 is the main cause for the enhanced apoptosis (Supplementary Fig. 1g). To determine, which metabolic pathway (i.e., the propionate breakdown or one-carbon metabolism) mediates the effects of the *Comamonas* DA1877 diet or B12 supplementation, we examined *metr-1(lf)* and *mmcm-1(lf)* mutants. Both mutations suppressed the effect of the *Comamonas* DA1877 diet and vitamin B12 on germ cell apoptosis and oocyte differentiation, though the suppression by *mmcm-1(lf)* was weaker (Fig. 1f, g, i, j). Only *metr-1(lf)* mutants showed a reduction in VPC induction in animals fed with the *Comamonas* DA1877 diet or supplemented with vitamin B12 (Fig. 1h, k). To determine if the effects of vitamin B12 on germ cell differentiation are cell-autonomous, we expressed *metr-1* and *mmcm-1* in the soma using extra-chromosomal arrays, which are silenced in the germline. The somatic expression of *metr-1* rescued the reduced vulval induction in *let-60(gf)* double mutants (Supplementary Fig. 1h), but the somatic expression of *metr-1* or *mmcm-1* did not affect germ cell death on the *Comamonas* DA1877 diet (Supplementary Fig. 1i).

Furthermore, we examined duct cell specification during larval development as a fourth phenotype controlled by RAS/MAPK signaling<sup>19</sup>. Eighty-one percent of *let-60(gf)* animals fed with *E. coli* OP50 showed a duplication of the duct cell, whereas 98% of the animals fed with the *Comamonas* DA1877 diet had two duct cells (Supplementary Fig. 1j, j').



In conclusion, the vitamin B12-dependent one-carbon metabolism is a major factor in enhancing multiple RAS/MAPK-induced cell fates on the *Comamonas* DA1877 diet.

### Vitamin B12-rich diet accelerates germ cell differentiation

Given that the *Comamonas* DA1877 diet affected both germ cell death and oocyte differentiation, we tried to decouple the two phenotypes. Using apoptosis-deficient *ced-3(lf)* mutants, we found that the increase in oocyte number in *Comamonas* DA1877-fed worms was independent of apoptosis (Fig. 2a). However, we were not able to determine if the reverse was also true.

The mitotic proliferation of germline stem cells in the distal gonad region depends on RAS/MAPK signaling in the soma<sup>30</sup>. We, therefore, tested if the *Comamonas* DA1877 diet up-regulates germ cell proliferation, which would increase germ cell number and potentially apoptosis. The mitotic index (i.e., the fraction of M-phase germ cells in the mitotic region) in adult animals was not changed by the DA1877 diet, by somatic hyper-activation of RAS/MAPK, or by loss of *metr-1* (Fig. 2b). However, *Comamonas* DA1877-fed worms contained a longer mitotic zone with more mitotic cells and an increased absolute number of cells in M-phase (Supplementary Fig. 2a), which is consistent with the overall increase in gonad size observed in *Comamonas* DA1877-fed animals. By contrast, *let-60(gf)* and *metr-1(lf)* mutants contained a shorter mitotic zone (Supplementary Fig. 2a). This suggests that the *Comamonas* DA1877 diet may affect the germline by expanding the pool of mitotic cells earlier during larval development without increasing the mitotic rate during adulthood. Thus, accelerated germ cell proliferation is unlikely to be responsible for the *Comamonas* DA1877-induced increase in apoptotic corpse and oocyte numbers.

We next focused on the pachytene region, where RAS/MAPK activation regulates both germ cell apoptosis and pachytene exit<sup>17,18,24</sup>. We used an H2B::GFP marker to label the germ cell nuclei and determine the border between the pachytene and diplotene zones (P-D) (Fig. 2c). We counted the average number of pachytene cells over 15-cell rows distal to the P-D border. *Comamonas* DA1877-fed animals contained more pachytene germ cells (Fig. 2d), consistent with an enlarged gonad. To hyper-activate RAS/MAPK signaling in the germline, we examined temperature-sensitive *let-60(ga89)* mutants, abbreviated *let-60(gf<sup>ts</sup>)*, grown for 4 h at 25 °C<sup>31</sup>. The *let-60(gf<sup>ts</sup>)* and *metr-1(lf)* mutations did not significantly alter germ cell number (Fig. 2d). *Comamonas* DA1877-fed worm and *let-60(gf<sup>ts</sup>)* mutants also showed an increased density of pachytene cells (i.e., shorter 15-cell rows), but cell density was not changed in *metr-1(lf)* mutants (Fig. 2e). Thus, neither cell number nor density seems to correlate with germ cell apoptosis or oocyte differentiation. However, *Comamonas* DA1877-fed worms showed a distal shift of the P-D border before the loop, whereas *let-60(gf<sup>ts</sup>)* mutants showed a proximal shift after the loop (Fig. 2c, f).

The distal shift of the P-D border in *Comamonas* DA1877-fed worms might be due to a change in germline length, a shift of the borders between the different meiotic zones or altered dynamics of the germ cells progressing through the gonads. Immunohistochemical staining of dissected gonads with markers for the different meiotic zones indicated that the boundary between the transition zone and early pachytene was shifted distally in *Comamonas* DA1877-fed animals (when corrected for different gonad length), but the proximal border of the late pachytene zone was not changed (Supplementary Fig. 2b). By contrast, the nuclear morphology visualized with the H2B::GFP marker pointed to a distal shift of the P-D border (Fig. 2c, f). In agreement with previous results<sup>22</sup>, *let-60(gf<sup>ts</sup>)* worms had a shorter mid-pachytene zone and no change in the proximal border of the late pachytene zone, despite the proximal shift of the P-D border determined with the H2B::GFP marker (Supplementary Fig. 2b and Fig. 2c, f). This suggests that the different meiotic zones determined by molecular markers do not always match the meiotic stage of individual germ cells scored by their nuclear morphology.

We, therefore, hypothesized that the changes in germ cell death and differentiation caused by the *Comamonas* DA1877 diet or the *let-60(gf<sup>ts</sup>)* mutation might be caused by an altered speed at which individual germ cells move through the different meiotic zones. We therefore tracked individual germ cell nuclei progressing from pachytene to diplotene by live imaging<sup>31</sup>. Germ cells progressed faster through the pachytene region in *Comamonas* DA1877-fed animals and *let-60(gf<sup>ts</sup>)* mutants (Fig. 2g and Supplementary Fig. 3a, b for the individual trackings). Moreover, *Comamonas* DA1877-fed worms ovulated more frequently (Fig. 2h and Supplementary Fig. 3c for the individual animals), indicating that the higher oocyte number is not due to reduced ovulation.

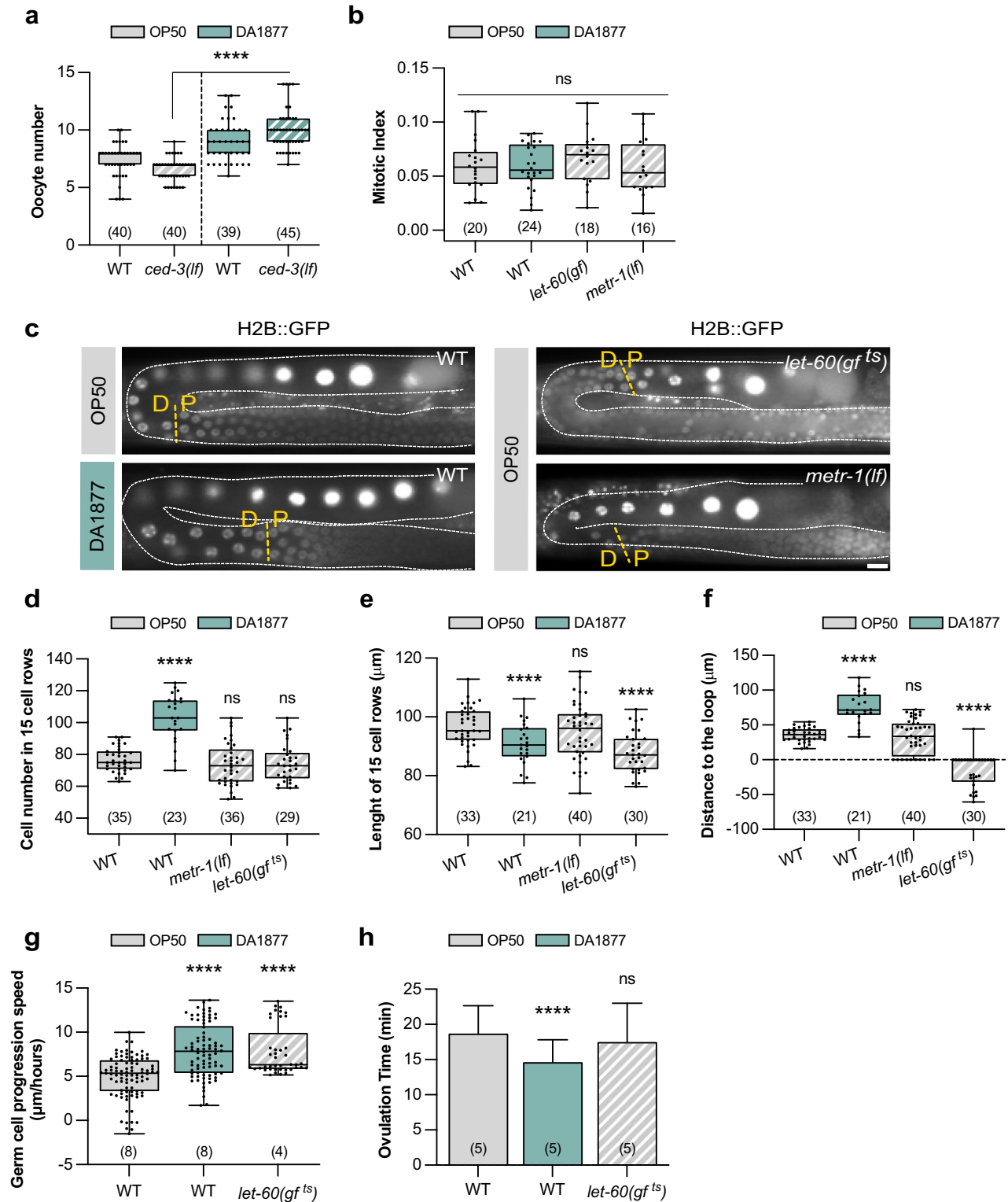
In summary, the *Comamonas* DA1877 diet or hyper-activation of the RAS/MAPK pathway resulted in a faster progression of germ cells through pachytene, which could explain the increased number of oocytes in the proximal gonads. Whether the accelerated progression through pachytene might also increase the rate of germ cell apoptosis remains unclear. It seems likely that the combination of the changes in gonad morphology and dynamics caused by the *Comamonas* DA1877 diet accounts for the increase in germ cell death.

### One-carbon metabolism enhances MAPK target gene expression

To characterize the interactions between one-carbon metabolism and RAS/MAPK signaling, we combined the *metr-1(lf)* and *let-60(gf)* mutations. *metr-1(lf)* rescued the *let-60(gf<sup>ts</sup>)*-dependent increase in germ cell apoptosis and oocyte number in animals fed with *E. coli* OP50, even in germline-specific *metr-1(lf)* mutants (Fig. 3a, b and Supplementary Fig. 4a, b). Furthermore, the gonads in *let-60(gf<sup>ts</sup>)* animals grown for 18 h at 25 °C shrank, resulting in a reduced rachis diameter<sup>18,29</sup>, which was suppressed by *metr-1(lf)* (Fig. 3c). Also, the increased VPC differentiation in *let-60(gf)* mutants fed with *E. coli* OP50 was suppressed by *metr-1(lf)* (Fig. 1h and 3d). Together, these results show that inhibiting the methionine cycle reduces the penetrance of phenotypes caused by RAS/MAPK hyper-activation in different tissues. On the other hand, the *Comamonas* DA1877 diet enhanced the effects of *let-60(gf<sup>ts</sup>)* on germ cell apoptosis and oocyte differentiation (Fig. 3e, f). To further examine the influence of the *Comamonas* DA1877 diet on RAS/MAPK signaling, we used the temperature-sensitive *mpk-1(ga111<sup>ts</sup>)* allele<sup>32</sup>, rescued in the soma with an extra-chromosomal *mpk-1(+)* array to inhibit RAS/MAPK specifically in the germline<sup>29</sup>. The reduced germ cell apoptosis and oocyte number in germline-specific *mpk-1(lf<sup>ts</sup>)* mutants were partially suppressed by the *Comamonas* DA1877 diet (Fig. 3g, h). This suggested that the RAS/MAPK pathway and the methionine cycle have synergistic effects, or that the methionine cycle performs a limiting function downstream of the RAS/MAPK pathway.

Next, we examined whether the methionine cycle directly regulates RAS/MAPK pathway activity. Neither immunostaining of dissected gonads nor Western blot analysis of total animal extracts with a di-phospho-ERK-specific antibody showed an increase in MAPK phosphorylation in animals fed with the *Comamonas* DA1877 diet (Fig. 3i, j and Supplementary Fig. 4c for the Western blots used for quantification). Accordingly, *metr-1(lf)* did not cause a decrease in MAPK phosphorylation (Supplementary Fig. 4d, e). Also, using an ERK biosensor to quantify RAS/MAPK pathway activity in the VPCs<sup>33</sup> we observed no significant difference in activity between worms fed with the *E. coli* OP50 or *Comamonas* DA1877 diet (Fig. 3k). Together, these results indicate that the *Comamonas* DA1877 diet does not increase RAS/MAPK pathway activity but rather the cellular responses to MAPK activation.

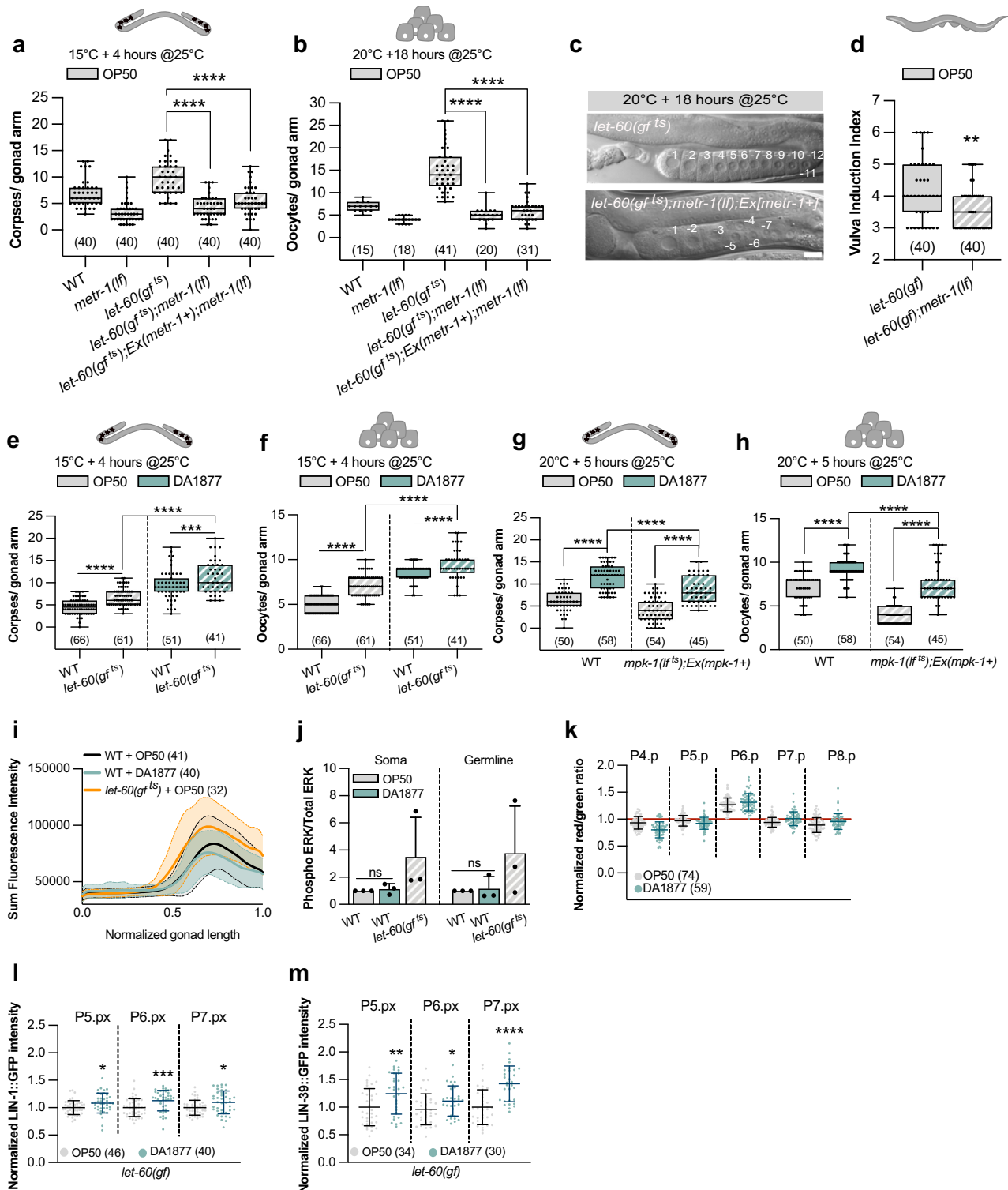
We thus tested whether the *Comamonas* DA1877 diet changes the expression of RAS/MAPK signaling target genes during vulval development. We used endogenous GFP reporters for *lin-1*, which encodes an ETS-family transcription factor phosphorylated by MPK-1, and *lin-39*, which encodes a *hox* gene regulated by LIN-1<sup>34</sup>. The *Comamonas* DA1877 diet increased LIN-1::GFP and LIN-39::GFP expression in the



**Fig. 2 | The *Comamonas* DA1877 diet changes germ cell number and dynamics.**

**a** Oocyte number for indicated genotypes. Dots represent individual oocytes. **b** Mitotic index for indicated genotypes. Dots represent individual animals. **c** Fluorescence images showing the pachytene-to-diplotene (P-D) transition (yellow dashed lines) for indicated genotypes; germ cell nuclei are marked with H2B::GFP. Gonads are outlined by white dashed lines. Scale bar: 10  $\mu$ m. **d** Number of cells inside an area starting from the P-D and ending 15 cell-rows distally of indicated genotypes; P-D was determined based on the nuclear marker shown in **c**; in *let-60(gf)* animals the loop region was used as the starting point. **e** Length of 15-cell rows for indicated genotypes. **f** Distance from the P-D to the loop region for

indicated genotypes. **d-f** dots represent individual animals. **g** Speed of germ cell progression through the pachytene region for indicated genotypes; for individual tracking see Supplementary Fig. 2. Dots represent the speed of individual cells. **h** Ovulation time for indicated genotypes. Dots represent individual oocytes. Bars represent mean  $\pm$  SD. **a-h** Animals were fed with *E. coli* OP50 (gray) or *Comamonas* DA1877 (blue). Number of animals in brackets. Boxplots show 25-75 percentiles and median; whiskers represent min and max values. \*\*\*\* $P < 0.0001$  using two-way ANOVA followed by uncorrected Fisher's LSD multiple comparison test in **a**, one-way ANOVA followed by Dunnett's multiple comparison tests in **b, d-h**. Source data are provided as a Source Data file.



proximal VPCs of *let-60(gf)* larvae (Fig. 3l, m), but it had no significant effect in wild-type worms (Supplementary Fig. 4f, g).

Thus, the interaction between the methionine cycle and the RAS/MAPK pathway most likely occurs at the level of the target genes. Furthermore, the methionine cycle appears to limit vulval induction only in the context of a hyper-activated RAS/MAPK pathway.

### Tissue-specific effects of one-carbon metabolites

Since loss of *metr-1* function suppressed the effects of the *Comamonas* DA1877 diet (Fig. 1f, g, i, j), we supplemented the growth medium with

different one-carbon metabolites to test if they would mimic the dietary effects on RAS/MAPK-induced cell fates. Specifically, we tested folinic acid, a derivative of folic acid that is converted into the different THF metabolites to enhance the folate cycle, as well as L-methionine and choline to enhance the methionine cycle and phospholipid metabolism, respectively (Fig. 1a).

Wild-type animals fed with *E. coli* OP50 and exposed to one of the three metabolites had more apoptotic corpses and oocytes (Fig. 4a, b). Moreover, each of the three metabolites was sufficient to bypass the inhibitory effects of *metr-1(lf)* on germ cell differentiation (Fig. 4a, b).

**Fig. 3 | Interaction between one-carbon metabolism and the RAS/MAPK pathway.** **a, b** Corpse and oocyte numbers for indicated genotypes grown at 15 °C and transferred to 25 °C 4 h before quantification, or grown at 20 °C and transferred to 25 °C 18 h. **c** DIC images of gonads of indicated genotypes grown at 20 °C and transferred to 25 °C 18 h before quantification. Numbers indicate mature oocytes. Scale bar: 10 μm. **d** Vulval induction index for indicated genotypes. **a–d** Animals were fed with *E. coli* OP50. **e, f** Corpse and oocyte numbers for indicated genotypes grown at 15 °C and transferred to 25 °C 4 h before quantification. **g, h** Corpse and oocyte numbers for indicated genotypes grown at 20 °C and transferred to 25 °C 5 h before quantification. **e–h** Animals were fed with *E. coli* OP50 (gray) or *Comamonas* DA1877 (blue). **i** dpERK intensity profiles in gonads of wild-type fed with *E. coli* OP50 (gray) or *Comamonas* DA1877 (blue), and *let-60(gf<sup>5</sup>)* fed with *E. coli* OP50 (orange). x-axis represents normalized gonad length. Average intensities ± SD (shaded) are shown. Data from three independent biological replicates. **j** Western blot quantification of the somatic and germline isoform of

dpERK for indicated genotypes. Bars represent mean ± SD normalized to the control (wild-type *E. coli* OP50) from three independent biological and technical replicates. See Supplementary Fig. 4 for the individual WB. **k** ERK-nKTR biosensor quantification. Dots represent individual animals from 2 independent biological replicates. Data show median ± min and max. **l–m** Normalized intensity of LIN-1::GFP or LIN-39::GFP at the Pn.px stage in 1° and 2° VPCs of *let-60(gf<sup>5</sup>)* animals. Data were normalized to the control (*E. coli* OP50). Dots represent individual animals from two independent biological replicates. **j–m** Animals were fed with *E. coli* OP50 (gray) or *Comamonas* DA1877 (blue). Number of animals is in brackets. In **a–h, j, l, m**, boxplots show 25–75 percentiles and median; whiskers represent min and max values. \**P* < 0.05, \*\**P* < 0.005, \*\*\**P* < 0.0005, \*\*\*\**P* < 0.0001 using one-way ANOVA followed by Dunnett's multiple comparison tests in **a** and **b**, a Kruskal-Wallis test for non-parametric data in **d**, two-way ANOVA followed by an uncorrected Fisher's LSD multiple comparison tests in **e–h** and **j** and unpaired, two-tailed t-tests in **l** and **m**. Source data are provided as a Source Data file.

Only choline significantly enhanced VPC induction in *let-60(gf)* single mutants, but methionine and choline both reversed the effect of *metr-1(lf)* on vulval induction (Fig. 4c). In contrast, folinic acid did not affect vulval induction (Fig. 4c).

Since the folate cycle is involved in nucleotide production, we tested two bacterial strains that produce more nucleotides, *E. coli* OP50 *cytR* and *E. coli* HT115<sup>35</sup>. In the wild-type background, both strains enhanced germ cell apoptosis and oocyte differentiation (Supplementary Fig. 5a, b). Since *metr-1(lf)* worms showed an abnormal gonad development when grown on these two bacterial strains, we examined somatically rescued *metr-1(lf)* mutants, which showed an increase in apoptotic corpse and oocyte numbers when fed with *E. coli* *cytR* or *E. coli* HT115 (Supplementary Fig. 5a, b), suggesting that nucleotide levels affect both processes. Vulval induction, on the other hand, was not enhanced by the two nucleotide-rich bacterial strains (Supplementary Fig. 5c). Furthermore, supplementing the growth medium with nucleosides caused an increase in apoptotic corpses and oocyte numbers in both wild-type animals and somatically rescued *metr-1(lf)* mutants (Fig. 4d, e), but did not affect vulval induction (Fig. 4f).

Analogous to the *Comamonas* DA1877 diet, supplementing choline or nucleosides partially suppressed the effects of germline-specific loss of *mpk-1* on germ cell apoptosis and oocyte development (Supplementary Fig. 5d, e), indicating that these metabolites enhance RAS/MAPK signaling. Taken together, folate and methionine cycle metabolites affect germ cell differentiation and death, whereas vulval induction is only sensitive to methionine cycle metabolites.

Phosphatidylcholine (PC) biosynthesis depends on choline and methyl groups provided by the methionine cycle to methylate phosphatidylethanolamine (PE) (Fig. 1a). Since choline supplementation had a strong effect on all processes examined (Fig. 4a–c), we hypothesize that PC levels might regulate germ cell and VPC differentiation. PC produced via the SAM-dependent pathway requires the *pmt-1* and *pmt-2* methyltransferases, while the choline-dependent pathway involves *cka-1* and *cept-1* activity<sup>8,36</sup> (Fig. 1a). We therefore used RNAi to test if the two PC synthesis pathways are necessary for the vitamin B12 effect on RAS/MAPK-controlled cell fates. Down-regulation of either *cka-1* or *cept-1* suppressed the vitamin B12-dependent increase in oocyte number and vulval induction (Fig. 4g, h). Due to the loss of the CED-1::GFP reporter signal in RNAi-treated animals, we could not score germ cell apoptosis. Since *pmt-2* RNAi induced larval arrest, we treated larvae with *pmt-2* RNAi bacteria diluted with empty vector (EV) starting at the L1 stage or exposed L4 larvae to undiluted *pmt-2* RNAi bacteria. In both treatments, *pmt-2* RNAi caused a reduction in oocyte number of wild-type animals supplemented with vitamin B12 (Fig. 4i). *pmt-2* RNAi also reduced vulval induction in *let-60(gf)* animals supplemented with vitamin B12 (Fig. 4j).

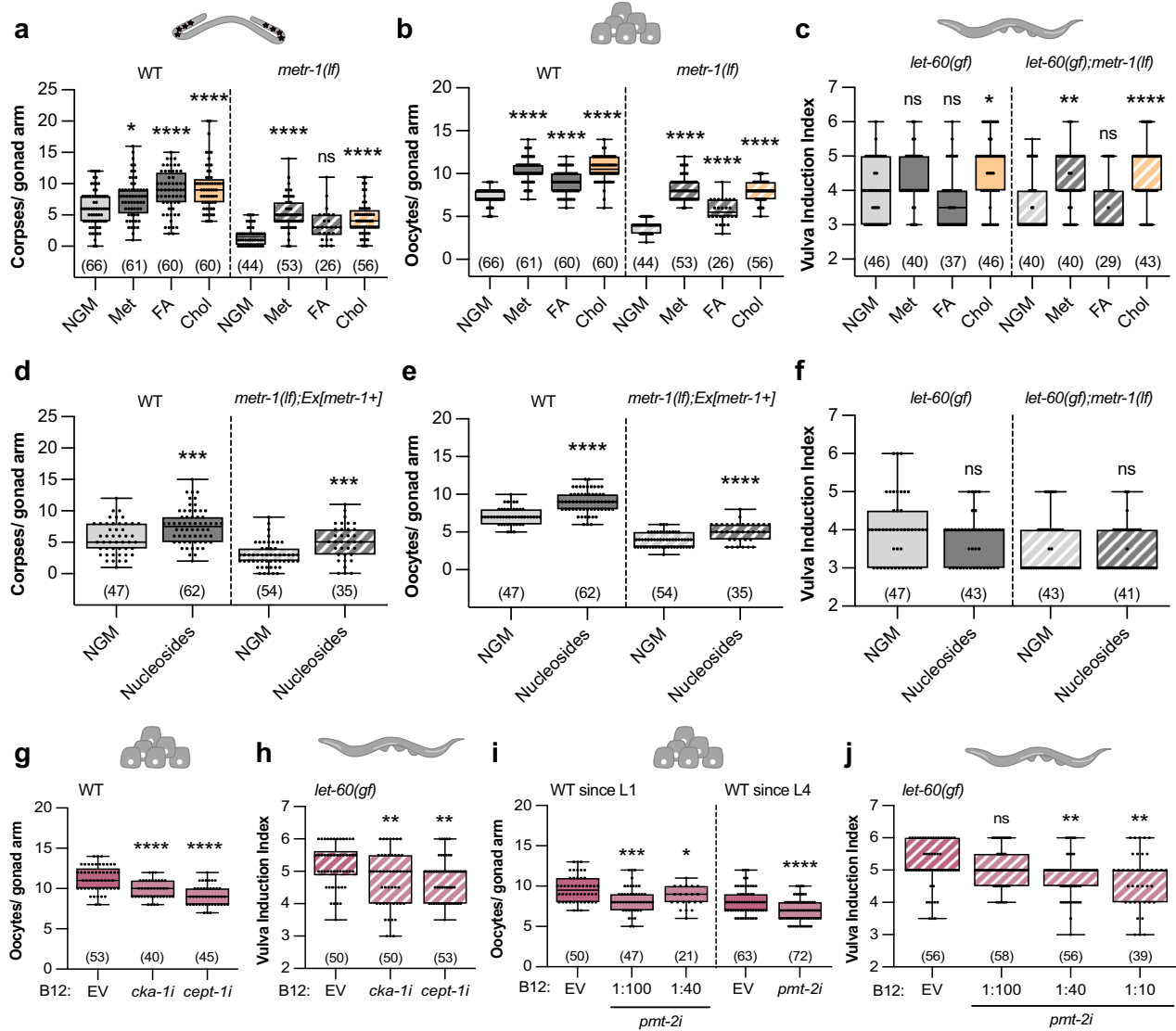
Together, these findings indicate that PC biosynthesis mediates the effects of vitamin B12 supplementation on oocyte and VPC differentiation, suggesting that the effects of choline are mediated by PC.

### The lipid metabolism represses vulval differentiation

Since one-carbon metabolism positively regulates germline and vulval development, and the *Comamonas* DA1877 diet alters gene expression<sup>11,37,38</sup>, we hypothesized that the *Comamonas* DA1877 diet enhances germ cell and VPC differentiation by changing the expression of genes that modulate the cellular responses to RAS/MAPK signaling. RNAseq analysis of adult wild-type and *let-60(gf<sup>5</sup>)* animals fed with the *E. coli* OP50 or *Comamonas* DA1877 diet identified in total 389 significantly down-regulated and 394 significantly up-regulated genes in *Comamonas* DA1877-fed animals (Fig. 5a, b and Supplementary Data 1). The *Comamonas* DA1877 diet caused similar changes in the transcriptome of wild-type and *let-60(gf<sup>5</sup>)* worms (Fig. 5b). Specifically, vitamin B12-, methionine cycle- and PC biosynthesis-related genes (*nhr-114*, *pmp-5*, *metr-1*, *sams-1*, *pmt-1* and *pmt-2*) were down-regulated in *Comamonas* DA1877-fed worms (Fig. 5a, b and Supplementary Fig. 6a), suggesting a negative feedback response to increased levels of dietary vitamin B12<sup>11,37,38</sup>. Furthermore, we did not detect significant changes in the expression levels of RAS/MAPK pathway genes (Supplementary Fig. 6a). Pathway network analysis showed that the *Comamonas* DA1877 diet regulates several genes involved in innate immune response and fatty acid biosynthesis (*fat-5*, *fat-6* and *fat-7*) (Supplementary Fig. 6b). Previous work has shown that diet-induced changes in gene expression can differ between young and gravid adult worms<sup>11</sup>. We, therefore, tested if the *Comamonas* DA1877 diet might influence gene expression in an age-dependent manner. We performed transcriptomic analysis of wild-type and *let-60(gf)* larvae at the mid-L3 stage, the stage when vulval induction occurs. The larval transcriptome showed smaller but similar changes in gene expression as observed in adults, as SAM, PC, and fatty acid metabolism were also among the most strongly affected pathways in L3 larvae (Supplementary Fig. 7a–c and Supplementary Data 2). We did not observe significant changes in *lin-1* and *lin-39* expression (Supplementary Fig. 7c), possibly because bulk RNAseq may not be sensitive enough to detect tissue-specific changes.

Previous work established a connection between vitamin B12, the methionine cycle and fat metabolism<sup>39</sup>. Given that genes controlling the lipid metabolism (i.e., the lipid Δ9-desaturases *fat-5*, *fat-6* and *fat-7*) were among the most strongly down-regulated genes, we explored the effect of the *Comamonas* DA1877 diet on the lipid metabolism and its consequences on RAS/MAPK-induced cell fates. Quantification of lipid droplet size using the *dhs-3::gfp* reporter<sup>40</sup> indicated that the intestines of *Comamonas* DA1877-fed animals contained smaller lipid droplets than *E. coli* OP50-fed animals (Fig. 5c, c'). Choline supplementation had a similar effect (Fig. 5c, c'), suggesting the *Comamonas* DA1877 diet's effect on lipid droplet size is choline-dependent. The *fat* genes are involved in different fatty acid desaturation and elongation pathways. *fat-5* is responsible for palmitoleic acid (C16:1n7) production, whereas *fat-6* and *fat-7* are responsible for oleic acid (C18:1n9)<sup>41</sup>. We tested if *fat-5*, *fat-6* or *fat-7* RNAi affected vulval induction and oocyte





**Fig. 4 | One-carbon metabolites regulate VPC and germline differentiation.** Number of corpses (**a**, **d**), oocytes (**b**, **e**), and vulval induction index (**c**, **f**) for indicated genotypes fed with *E. coli* OP50 and supplemented with the following metabolites; Met – 5 mM methionine; FA – 100  $\mu$ M folic acid; Chol – 40 mM choline; 1 mM nucleosides. Oocyte number of wild-type animals (**g**, **i**) and vulval induction index of *let-60(gf)* animals (**h**, **j**) fed with indicated RNAi clones or control (EV – empty vector). NGM was supplemented with 64 nM of B12; animals were exposed to RNAi from L1 except where indicated (L4). **a–j** Dots represent individual corpses/oocytes or animals from two independent biological replicates. Boxplots

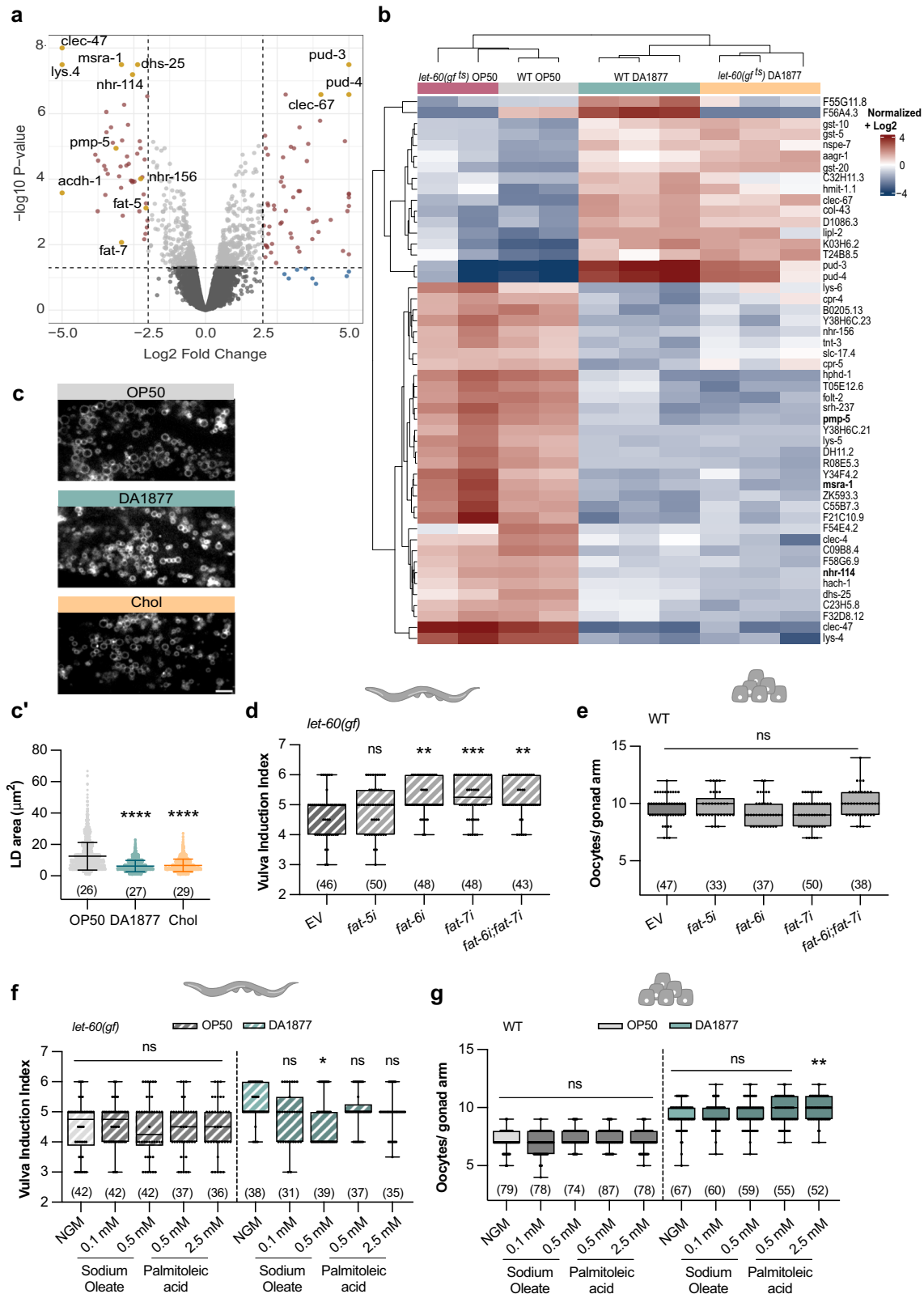
show 25–75 percentiles and median; whiskers represent min and max values. Number of animals in brackets. \* $P < 0.05$ , \*\* $P < 0.005$ , \*\*\* $P < 0.0005$ , \*\*\*\* $P < 0.0001$  using two-way ANOVA followed by a Tukey’s multiple comparison test in **a–c**, two-way ANOVA followed by uncorrected Fisher’s LSD multiple comparison test in **d**, **e**, and **f**, one-way ANOVA followed by Dunnett’s multiple comparison test in **g** and **i** (WT since L1), an unpaired, two-tailed *t*-test in **i** (since L4) and a Kruskal-Wallis test for non-parametric data followed by a Dunn’s multiple comparison test in **h** and **j**. Source data are provided as a Source Data file.

differentiation. Single or double knock-down of *fat-6* and *fat-7* increased vulval induction in *let-60(gf)* animals, but *fat-5* knock-down had no significant effect (Fig. 5d). Oocyte development was not affected by *fat-5*, *fat-6* or *fat-7* RNAi (Fig. 5e). Finally, we supplemented palmitoleic acid and sodium oleate to animals fed with *E. coli* OP50 or *Comamonas* DA1877. Supplementing sodium oleate to *let-60(gf)* animals fed with the *Comamonas* DA1877 diet reduced vulval induction to the levels observed in *E. coli* OP50-fed animals (Fig. 5f). However, oocyte differentiation was not changed by fatty acid supplementation, irrespective of the diet (Fig. 5g).

In summary, the down-regulation of unsaturated fatty acid biosynthesis by the *Comamonas* DA1877 diet, especially oleic acid, positively regulated vulval induction, but it did not affect germ cell differentiation.

### A vitamin B12-rich diet increases histone H3K4 methylation

Since the methionine cycle is the main methyl donor in cells, we investigated whether changes in histone methylation might regulate germ cell and VPC differentiation. We focused on histone H3K4me3 because this type of histone modification is sensitive to diet, particularly a high-fat diet<sup>39</sup>, as well as changes in methionine cycle activity<sup>10,14,42</sup>. Western blot analysis showed higher global H3K4me3 levels in worms fed with *Comamonas* DA1877 or supplemented with choline and slightly reduced levels in *metr-1(lf)* mutants fed with *E. coli* OP50 (Fig. 6a and Supplementary Fig. 8a for the Western blots used for quantification). Since the *Comamonas* DA1877 diet down-regulated fat biosynthesis (Fig. 5c), we tested if inhibiting *fat* genes affected histone methylation. H3K4me3 levels did not increase after *fat-6* or *fat-7* single or double RNAi, indicating that the oleic acid metabolism does not



regulate H3K4me3 levels (Fig. 6b and Supplementary Fig. 8b for the Western blots used for quantification).

To test if the increase in H3K4me3 levels caused by the *Comamonas* DA1877 diet contributed to vulval induction and germ cell differentiation, we inhibited by RNAi the four H3K4 demethylases *amx-1*, *lsd-1*, *rbr-2* and *spr-5* (which might mimic the *Comamonas* DA1877 diet in *E. coli* OP50-fed animals) and the four H3K4

methylases *ash-2*, *set-2*, *set-16* and *wdr-5.1* (which might suppress the vitamin B12 effect). RNAi of the demethylases *amx-1* and *spr-5* increased, and RNAi of the methylase *wdr-5.1* decreased vulval induction in *let-60(gf)* animals supplemented with vitamin B12 (Fig. 6c). The effect of *wdr-5.1* RNAi appeared to be specific to enhanced one-carbon metabolism, since *wdr-5.1* RNAi caused no reduction in vulval induction without vitamin B12 supplementation

**Fig. 5 | Dietary effects on gene expression and lipid metabolism.** **a** Differentially expressed genes in adult wild-type animals fed with *E. coli* OP50 or *Comamonas* DA1877. Genes with a *p*-value < 0.05 and a log<sub>2</sub>-fold change > 2.5 are highlighted. *p*-values were calculated as described in the methods. **b** Top 50 differentially expressed genes for the indicated conditions. **c** Fluorescence images of *dsh-3::gfp* marking intestinal lipid droplets (LD) in animals fed with *E. coli* OP50 (gray), *Comamonas* DA1877 (blue), or *E. coli* OP50 supplemented with 40 mM choline (yellow). Scale bar: 10 μm. **c'** Area of intestinal LD; dots represent individual droplet areas of 100 droplets per animal, from two independent biological replicates. Bars represent mean with SD. vulval induction index of *let-60(gf)* (**d**) or oocyte number of wild-type animals (**e**) fed with indicated RNAi clones and control (EV – empty

vector). **f** Vulval induction index of *let-60(gf)* and **g**, oocyte number of wild-type animals fed with *E. coli* OP50 (gray) or *Comamonas* DA1877 (blue), supplemented with indicated fatty acids. **d–g** dots represent individual gonads (**e, g**) or animals (**d, f**) from two independent biological replicates. Boxplots show 25–75 percentiles and median; whiskers represent min and max values. Number of animals in brackets. \*\**P* < 0.005, \*\*\**P* < 0.0005, \*\*\*\**P* < 0.0001 using one-way ANOVA followed by Dunnett's multiple comparison test in **c'** and **e**, two-way ANOVA followed by a Tukey's multiple comparison test in **f** and **g** and a Kruskal-Wallis test for non-parametric data followed by a Dunn's multiple comparison test in **d, h, i**. Source data are provided as a Source Data file.

(Supplementary Fig. S8c). Oocyte differentiation was also increased by *amx-1*, *lsd-1* and *spr-5* RNAi and decreased by *set-2* and *wdr-5.1* RNAi (Fig. 6d). Both, *set-2* and *wdr-5.1* are important for oocyte development, since knock-down of both genes reduced oocyte numbers without vitamin B12 supplementation (Supplementary Fig. S8d).

Finally, we tested if changes in H3K4 methylation might alter RAS/MAPK target gene expression by quantifying LIN-1::GFP levels after RNAi knock-down of *amx-1*, *spr-5* and *wdr-5.1* (Supplementary Fig. S8e, f). We observed no significant changes in LIN-1::GFP expression, suggesting that altering H3K4 methylation alone is not sufficient to change RAS/MAPK target gene expression.

We conclude that a vitamin B12-rich diet causes a global increase in H3K4me3 levels, which together with one-carbon metabolites contribute to enhanced vulval induction and oocyte differentiation.

### Methionine-dependency of KRAS-transformed mammalian cells

To investigate if the role of the methionine cycle in regulating RAS/MAPK-induced cell fates is conserved in mammals, we cultured human cancer cell lines carrying activating mutations in KRAS in growth medium without methionine. We first performed collective cell migration assays with A549 lung cancer cells that carry a KRAS G12S mutation, to test if methionine restriction (MR) interfered with cell migration. To exclude possible effects of reduced cell proliferation, we performed these assays with cells arrested in the G1 phase by double thymidine blockade<sup>43</sup>. A549 cells grown in MR (without methionine but supplemented with its precursor homocysteine) migrated at a lower speed than cells without MR (Fig. 7a–a"). The addition of the MEK inhibitors MEK162 and LGX818 without MR reversibly blocked cell migration, indicating that RAS/MAPK signaling is necessary for the migration of A549 cells, and removal of the MEK inhibitors 8 h after starting the assay allowed the cells to resume migration (Fig. 7a–a"), indicating that the reduced migration was not caused by increased cell death.

We next used A431 epidermoid carcinoma cells, which over-express the EGFR and strongly respond to EGF stimulation by forming filopodia in a MAPK-dependent manner<sup>43</sup>. Since filopodia formation is difficult to assess quantitatively, we categorized cellular morphologies into three classes; 0 for cells without filopodia, 1 for cells with small filopodia covering part of the cortex, and 2 for cells with long filopodia covering most of the cortex (Supplementary Fig. 9a). Class 1 and 2 cells were scored as stimulated and class 0 cells as unstimulated. Without EGF stimulation, most cells did not form filopodia, while 90% of the cells stimulated with EGF in the presence of methionine formed filopodia (Fig. 7b, b' and Supplementary Fig. 9b). By contrast, 55% of cells with MR and stimulated with EGF did not form filopodia (Fig. 7b, b' and Supplementary Fig. 9b), indicating that methionine is required for A431 cells to fully respond to EGF stimulation. Treatment with the MEK inhibitors in the presence of methionine strongly reduced EGF-induced filopodia formation (Fig. 7b, b').

Since human cancer cell lines such as A459 carry many additional mutations besides the KRAS mutations, we used primary mouse embryonic fibroblasts (MEFs), in which the three RAS genes had been

deleted and replaced with vectors expressing wild-type, G12C, G12V KRAS, or V600E BRAF in the same genetic background<sup>43</sup>. Without MR, the KRAS mutant MEFs showed the same migration speed as KRAS wild-type control cells, suggesting that hyper-activation of the RAS/MAPK pathway alone is not sufficient to accelerate collective cell migration (Fig. 7c, c'). However, KRAS G12V and G12V cells migrated significantly slower with MR, whereas KRAS wild-type MEFs migrated at the same speed with or without MR (Fig. 7d, d'), suggesting that methionine levels are limiting in the context of hyper-active RAS/MAPK signaling. V600E BRAF mutant MEFs showed a similar effect, though the reduction in migration caused by MR was smaller.

Taken together, these results suggested that the methionine-dependency of RAS/MAPK-induced phenotypes may be conserved in mammalian cells.

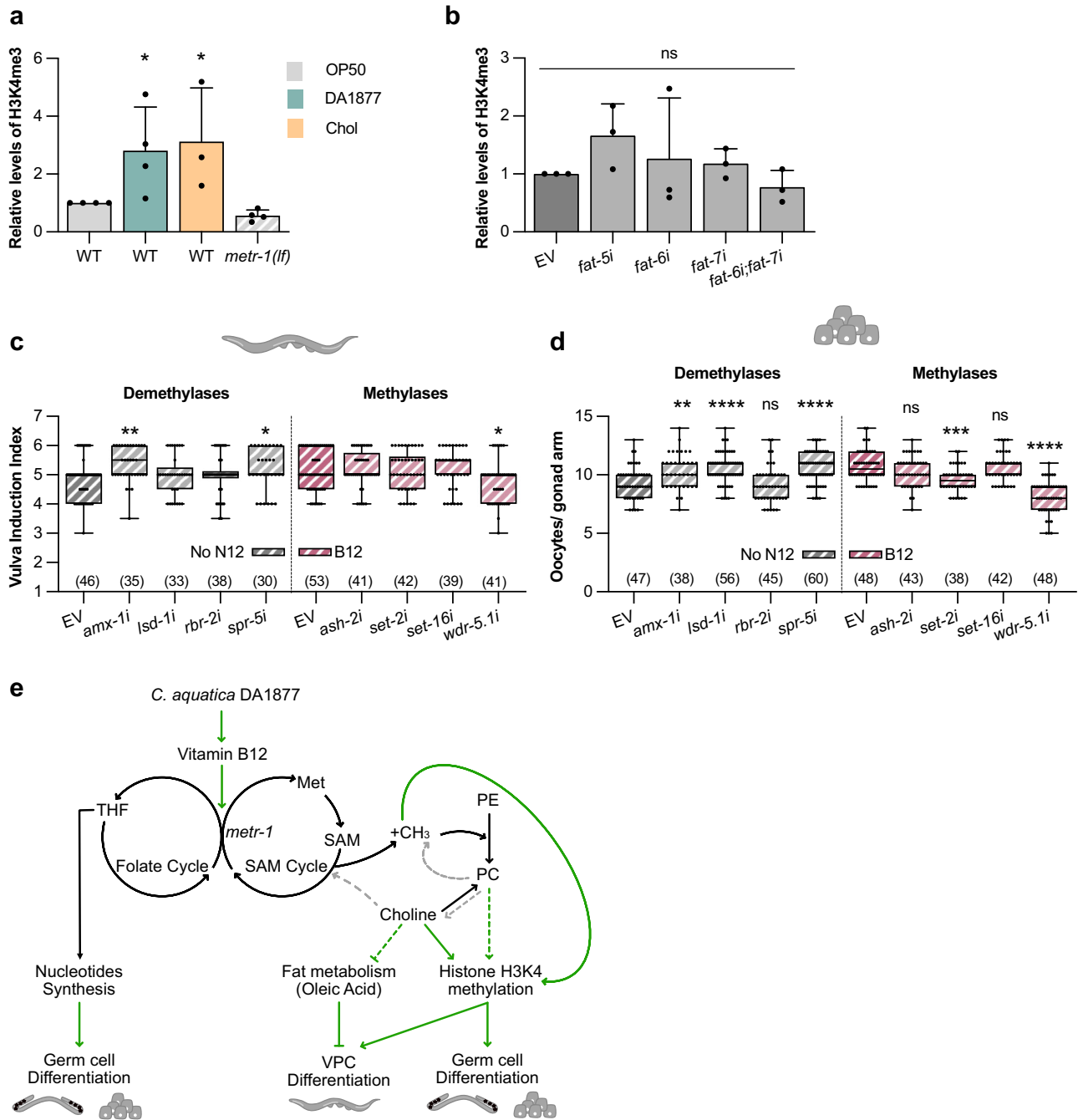
### Discussion

Diet has a major impact on the cellular metabolism. Therefore, cells must adapt their metabolism to the available nutrition to ensure survival in changing environments. Here, we focused on the vitamin B12-dependent one-carbon metabolism and its effects on cell fate decisions controlled by the oncogenic RAS/MAPK signaling pathway (Fig. 6e). A bacterial diet rich in vitamin B12 (*Comamonas* DA1877) or direct vitamin B12 supplementation enhanced germ cell apoptosis, oocyte differentiation and VPC induction, which are all controlled by the RAS/MAPK pathway. However, it should be noted that our data do not exclude potential vitamin B12-independent effects that may be caused by feeding a *Comamonas* DA1877 diet.

Different one-carbon metabolites have tissue-specific effects, as the folate cycle only affected germline development, while methionine cycle metabolites enhanced all RAS/MAPK-induced phenotypes. Choline and phosphatidylcholine (PC) are key mediators of vitamin B12-dependent effects by regulating histone H3K4me3 methylation and repressing fatty acid biosynthesis (Fig. 6e). We thus propose that methionine cycle activity is a limiting factor for different phenotypes caused by RAS/MAPK hyper-activation, by modulating the expression of downstream RAS/MAPK target genes. Finally, we show that methionine cycle activity is also a limiting factor in mammalian cells carrying activating mutations in KRAS, pointing to a conserved role of one-carbon metabolism in regulating cell RAS/MAPK-induced cell fates.

Folic acid and nucleoside supplementation positively regulated germ cell death and oocyte differentiation, but not vulval induction. Since the purine and pyrimidine metabolisms are important for cell proliferation the *E. coli* HT115 or *Comamonas* DA1877 diets may promote germline proliferation by increasing folate levels<sup>35</sup>. Accordingly, *Comamonas* DA1877-fed animals contained more M-phase cells. However, the *Comamonas* DA1877 diet did not accelerate cell cycle progression because the number of germ cells in M-phase relative to the total number of mitotic cells remained constant in a vitamin B12-rich diet. However, the vitamin B12-rich diet accelerated germ cell transition through pachytene into diakinesis.

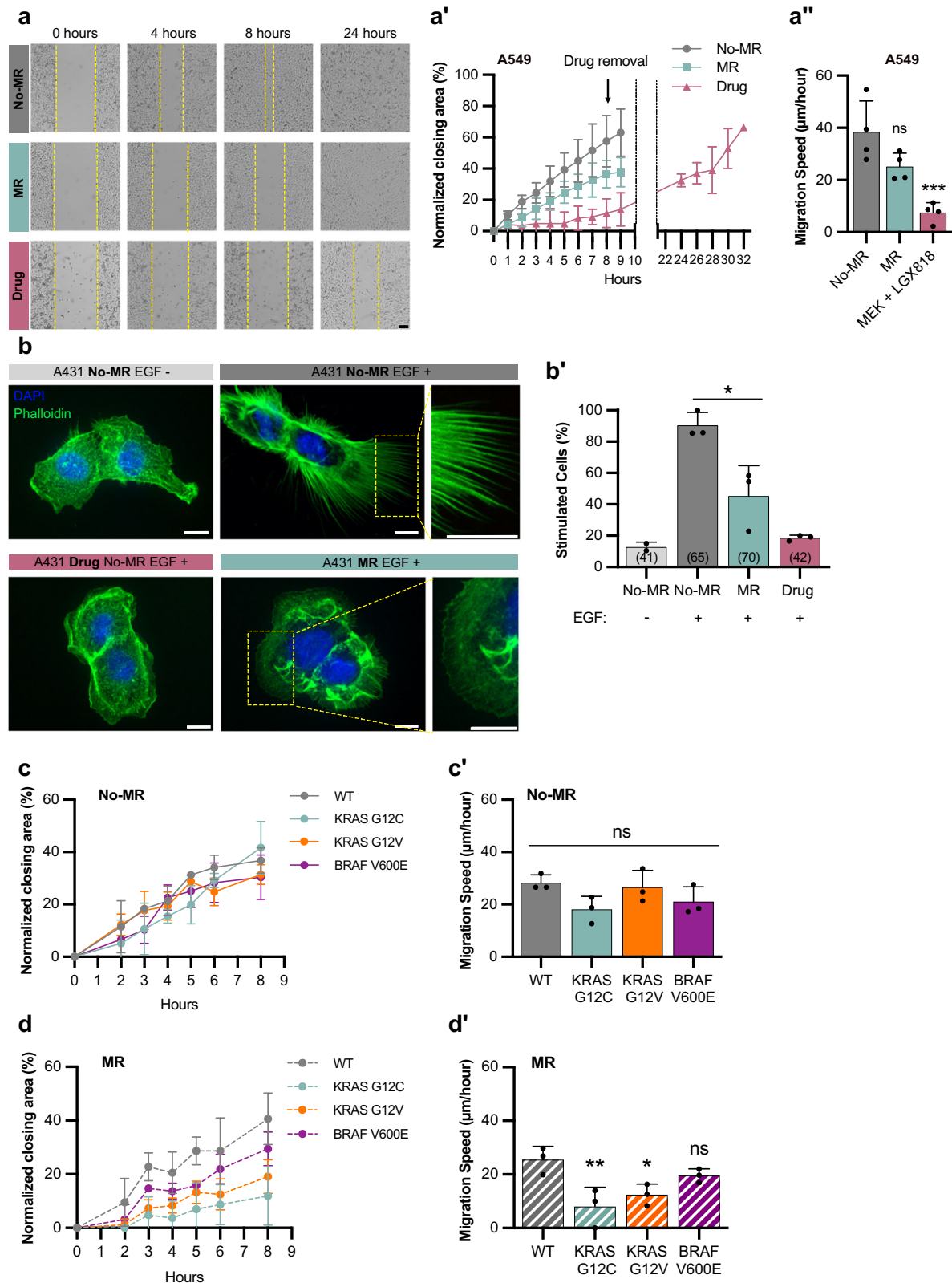
The increase in oocyte numbers is not linked to enhanced germ cell apoptosis since the apoptosis-deficient *ced-3(lf)* caspase mutants



**Fig. 6 | The *Comamonas* DA1877 diet increases H3K4 methylation to regulate VPC and oocyte differentiation.** **a** Western blot quantification of tri-methylated H3K4 for indicated genotypes fed with *E. coli* OP50 (gray), *Comamonas* DA1877 (blue), or *E. coli* OP50 supplemented with 40 mM choline (yellow); levels of tri-methylated H3K4 were normalized to total H3 levels. Bars represent mean  $\pm$  SD, normalized to the control (wild-type *E. coli* OP50), from four independent biological and technical replicates; see Supplementary Fig. 8a for individual Western blots. **b** Western blot quantification of tri-methylated H3K4 in wild-type animals fed with indicated RNAi clones; levels of tri-methylated H3K4 were normalized to total H3 levels. Bars represent mean  $\pm$  SD, normalized to the control (wild-type EV), from three independent biological and technical replicates; see Supplementary Fig. 8b for individual Western blots. Vulval induction of *let-60(gf)* animals (c) and oocyte number of wild-type animals (d) fed with indicated RNAi clones or control (EV –

empty vector); non-supplemented NGM (gray) or supplemented with 64 nM B12 (pink). Dots represent individual animals (c) or oocytes (d) from two independent biological replicates. Boxplots show 25–75 percentiles and median; whiskers represent min and max values. Number of animals in brackets. **e** Model depicting the tissue-specific effects of one-carbon metabolism on nucleosides synthesis, fat metabolism and histone methylation and its effects on germ cell and VPCs differentiation; green arrows are interactions supported by this work; dashed arrows are hypothesized from literature (gray) or conclusions made in this work (green). \* $P < 0.05$ , \*\* $P < 0.005$ , \*\*\* $P < 0.0005$ , \*\*\*\* $P < 0.0001$  using a Kruskal-Wallis test for non-parametric data followed by a Dunn’s multiple comparison test in a–c and one-way ANOVA followed by Dunnett’s multiple comparison test in d. Source data are provided as a Source Data file.





still showed the increase in oocyte number caused by the *Comamonas* DA1877 diet. While the mechanism by which nucleosides enhance germ cell apoptosis is not clear, we hypothesize that increasing nucleoside levels could promote oocyte differentiation by enhancing maternal mRNA production<sup>44</sup>. Oocyte differentiation is controlled by a meiotic checkpoint established by the HORMA complex. RAS/MAPK signaling regulates this checkpoint by controlling HTP-1

phosphorylation<sup>22</sup>. It is therefore possible that the B12 metabolism enhances HTP-1 phosphorylation and thereby increases the rate of oogenesis.

The effects of the vitamin B12-rich diet are associated with increased choline and PC levels. PC biosynthesis has previously been implicated in RAS/MAPK signaling in the survival of the *uv1* cells under excitotoxic cell death<sup>45</sup>. Hyper-activation of the RAS/MAPK pathway by

**Fig. 7 | Methionine dependency of RAS/MAPK-induced phenotypes in mammalian cells.** **a** Collective migration of A459 cells grown with MR (0  $\mu$ M Met; 400  $\mu$ M Hcy) (blue), without MR (200  $\mu$ M Met; 400  $\mu$ M Hcy) (gray), or treated with 1  $\mu$ M MEK162 and 1  $\mu$ M LGX818 MEK inhibitors (pink); yellow dashed lines represent the migrating front over time. **a'** Normalized percentage of the area closed over time. Symbols indicate mean  $\pm$  SD from four independent biological replicates. **a''**, Migration speed was calculated as the slope of graphs shown in **a'**. Bars represent the mean  $\pm$  SD. **b** Staining of A431 cells with DAPI (blue) and phalloidin (green), with and without EGF stimulation or MR. Scale bar: 10  $\mu$ m. For additional examples and the scoring criteria, see Fig. S9. **b'** Percentage of stimulated cells with filopodia

under indicated conditions. Bars represent mean  $\pm$  SD from three independent biological replicates; total number of cells scored in brackets. **c-d'** Quantification of collective migration in MEFs expressing different KRAS and BRAF mutations grown without **c** and with MR **d**. **c, d** Normalized percentage of area closed over time. Data represents mean  $\pm$  SD from three independent biological replicates. **c', d'** Migration speed was calculated as the slope of graphs shown in **c** and **d**, respectively. Bars represent the mean  $\pm$  SD. \* $P < 0.05$ , \*\* $P < 0.005$ , \*\*\* $P < 0.0005$  using one-way ANOVA followed by Dunnett's multiple comparison test in **a''**, **c'** and **d'**. Source data are provided as a Source Data file.

LET-23 EGFR leads to uv1 cell survival, which requires PC biosynthesis. Choline supplementation not only increased vulval induction but also enhanced physiological germ cell death and oocyte differentiation. Low choline or PC levels have previously been associated with stress- or radiation-induced apoptosis, which is mediated by ceramide signaling<sup>46,47</sup>. In this context, our results point to a distinct regulatory mechanism of physiological germ cell apoptosis controlled by the choline metabolism. Given that PC is one of the most abundant phospholipids in cell membranes and that RAS/MAPK signaling is required for membrane integrity during oogenesis<sup>23</sup>, choline could also promote oocyte production by facilitating membrane production. This is consistent with our observation that germ cells in animals fed with the *Comamonas* DA1877 diet exited pachytene faster, which may reflect a higher availability of membrane components.

A vitamin B12-rich diet resulted in the global downregulation of the *fat* genes that are necessary for the biosynthesis of mono-unsaturated fatty acids (MUFAs). The *fat* genes are regulated by different transcription factors such as SBP-1, NHR-49, NHR-80, and MDT-15. Interestingly, the methionine cycle and choline metabolism were previously reported to affect lipogenesis controlled by SBP-1 (a homolog of the mammalian sterol regulatory element-binding protein, SREBP)<sup>10,36,48</sup>. Moreover, in human cancer cells, RAS/MAPK signaling can regulate SREBP activity<sup>49</sup>. These interactions are consistent with our data indicating that a vitamin B12-rich diet increases PC levels and reduces the size of lipid droplets in the intestine, probably by down-regulating *fat-5*, *fat-6*, and *fat-7* expression. The relationship between PC and fat metabolism is not exclusive to *C. elegans*, as PC supplementation suppresses obesity-related phenotypes in mice fed with a high-fat diet<sup>50</sup>, and a choline-deficient diet enhances liver fat levels in rats<sup>51</sup>. The fatty acid metabolism influences *C. elegans* at the organismic level by regulating survival, developmental time, reproduction, ferroptosis, and modulating drug effects<sup>52,53</sup>. However, our data indicate that fatty acid levels also affect developmental cell fate decisions specifically, as only oleic acid inhibited VPC differentiation. Since fatty acids can act as signaling molecules<sup>41</sup>, we hypothesize that oleic acid may regulate vulval induction via a specific signaling pathway. Especially, oleic acid was reported to activate stress-induced signaling pathways in mammalian immune cells<sup>54</sup>.

A Vitamin B12-rich diet or choline supplementation increased global histone H3K4me3 levels, consistent with previous reports in worms and mammals indicating that the methionine cycle promotes H3K4 methylation<sup>9,55</sup>. Choline can be oxidized to betaine, functioning as a methyl donor to regenerate SAM<sup>56</sup>. However, the *C. elegans* genome does not encode an ortholog of choline oxidase<sup>57</sup>, and choline supplementation does not bypass the reduction in H3K4me3 levels in intestinal cells caused by *sams-1* knock-down<sup>10</sup>. Since phosphatidylethanolamine (PE) is a major consumer of methyl groups<sup>6</sup>, SAM and thereby also H3K4me3 levels may increase if PC production is shifted to the Kennedy/CDP-choline pathway (Fig. 1a). Thus, choline supplementation may promote histone methylation indirectly by reducing the competition for available methyl donors between histones and PE-dependent PC biosynthesis. Alternatively, choline could act directly as a methyl donor for H3K4 methylation.

The H3K4 methyltransferases SET-16 and WDR-5.1 were previously reported to regulate VPC differentiation<sup>58</sup>. Here, we show that only WDR-5.1 is required to mediate the effects of dietary vitamin B12, while the demethylases the AMX-1 and SPR-5 appear to counteract WDR-5.1. Enhancing methionine cycle activity may modulate chromatin accessibility of RAS/MAPK target genes by increasing H3K4me3 methylation of these loci by WDR-5.1, and thereby enhancing the transcriptional responses to MAPK activation.

In summary, we propose that the methionine cycle is a limiting factor for RAS/MAPK-induced cell fate decisions in several tissues of *C. elegans*.

The importance of RAS/MAPK signaling during oncogenic transformation is well documented, stimulating tumor cell proliferation, migration, and invasion<sup>15,16</sup>. Also, the methionine-dependency of cancer cells has been known since 1959<sup>59,60</sup>. The exact reason why cancer cells depend on methionine is not completely understood. One possible explanation is that cancer cells have a reduced ability to use endogenous methionine for SAM synthesis due to a reallocation of homocysteine to the trans-sulfuration pathway<sup>61</sup>. It is also possible that cancer cells have an increased demand for methionine and SAM due to more methylation reactions or to a SAM-dependent methylation check-point that regulates cell cycle progression through interactions with MCM, Cdc6, or Cdk2<sup>61</sup>. Moreover, MR globally decreases histone methylation, particularly H3K4me3 levels, which could attenuate the expression of RAS/MAPK target genes<sup>9,62</sup>. The relationship between one-carbon metabolism and cancer is also used for cancer therapy, as multiple chemotherapeutic drugs such as 5-fluorouracil (5-FU) or methotrexate inhibit one-carbon metabolism<sup>5</sup>.

Here, we show that different RAS/MAPK-regulated cellular processes depend on methionine. Using MEFs expressing different KRAS mutations as their only source of RAS activity, we observed remarkable differences in their sensitivity to MR. In the presence of methionine, KRAS mutant MEFs were indistinguishable from wild-type MEFs, but with MR only MEFs carrying activating KRAS mutations exhibited reduced cell migration. These findings support the idea of an increased dependency of RAS-transformed cancer cells on methionine cycle metabolites and mirror our findings with *C. elegans let-60 ras(gf)* mutants, which indicated that reducing methionine cycle activity limits the strength of the different phenotypes caused by RAS/MAPK hyperactivation. MR in cancer cells may attenuate RAS/MAPK signaling through one or several of the mechanisms we identified in the *C. elegans* model, i.e. by decreasing H3K4me3 methylation, increasing oleic acid biosynthesis or reducing nucleoside production. Taken together, these findings suggest that the mechanisms regulating this key carcinogenic signaling pathway by one-carbon metabolism may be conserved between worms and mammals.

## Methods

### *C. elegans* culture and bacterial strains used

*C. elegans* strains were maintained at 20 °C on standard nematode growth medium (NGM) plates seeded with *Escherichia coli* OP50. *C. elegans* N2 (Bristol) strain was used as wild-type<sup>63</sup>. Before an experiment, worms were synchronized by bleaching gravid adults and letting the embryos hatch without food. *Comamonas aquatica* DA1877, *E. coli*

OP50, *E. coli* HT115, and *E. coli* cytR<sup>-</sup> were obtained from CGC. *C. aquatica* DA1877 cIA/conB<sup>-</sup> and *C. aquatica* DA1877 cbiB<sup>3</sup> were obtained from the Walhout lab. All bacterial strains were grown overnight at 37 °C in liquid culture containing streptomycin (100 µg/ml) for *Comamonas* DA1877, kanamycin (50 µg/ml) for cytR<sup>-</sup> and streptomycin (100 µg/ml) and gentamycin (20 µg/ml) for both *Comamonas* DA1877 mutants. Supplementary Data 3 contains a list of all *C. elegans* strains used in this study.

### Microscopy

Live *C. elegans* were mounted either on 3% agarose pads or in custom-made microfluidic devices for high-throughput and long-term imaging<sup>31,64</sup>. Nomarski (DIC) and epifluorescence images were acquired on one of three microscope systems: a DMRA2 (Leica) microscope equipped with a sCMOS camera (Prime BSI, Photometrics), a multicolor fluorescence light source (Spectra, Lumencor) and a piezo objective drive (MIPOS 100 SG, Piezosystems Jena); a DMRA (Leica) microscope equipped with two sCMOS cameras (C11440-42U30 Hamamatsu), an image splitter (TwinCam, Cairn Research), a multicolor fluorescence light source (Spectra, Lumencor) and a piezo objective drive (MIPOS 100 SG, Piezosystems Jena); or an inverted Ti-U (Nikon) equipped with a sCMOS camera (Prime 95B, Photometrics), a multicolor fluorescence light source (LedHub, Omicron Laserage Laserprodukt GmbH) and a piezo objective drive (Nano-F100, Mad City Labs). Images were acquired using a 20x air objective (HC PL APO 20x/0.80, Leica), a 40x oil objective (HCX PL APO 40x/1.32, Leica; CFI Plan Fluor 40X Oil, Nikon) or a 63x oil objective (HCX PL APO 63x/1.32-0.60, Leica). Confocal images were acquired on a spinning disk confocal (SDC) system consisting of a BX61 (Olympus) microscope equipped with an X-light V2 spinning disc unit, an EMCCD camera (iXon Ultra 888, Andor), two high-power LEDs (UHP-T-460-DI and UHP-T-560-DI, Prizmatix) for GFP and RFP excitation, a mercury vapor lamp (X-Cite exacter, Excelitas Technologies Corp) for DAPI, CFP and YFP excitation and a piezo objective drive (MIPOS 100 SG, Piezosystems Jena). Images were acquired using a 40x oil objective (UPlanFL N 40x/1.30, Olympus) or a 60x oil objective (UPlanAPO 60x/1.40, Olympus).

### Germ cell apoptosis and oocyte numbers

Unless stated otherwise, all experiments were performed in one-day-old adult animals (72 h post L1 starvation) grown at 20 °C. To score germ cell apoptosis in temperature-sensitive *let-60(ga89)* (*gf<sup>ts</sup>*) animals, all worms were initially grown at 15 °C and transferred to 25 °C for 4 h before corpse scoring. Germ cell apoptosis was scored using the *bcls39[Plim-7::ced-1::gfp]* reporter<sup>27</sup> and quantified in one gonad arm by counting CED-1::GFP positive cells. To count oocytes with the *let-60(ga89)* (*gf<sup>ts</sup>*) allele, worms were initially grown at 20 °C and transferred to 25 °C for 18 h before counting oocyte numbers. Oocytes were scored in DIC images or using the SYN-4::GFP membrane marker, counting the number of completely cellularized oocytes in the proximal gonad arm.

### Vulval induction

All vulval induction experiments were conducted in the *let-60(n1046<sup>ts</sup>)* background. Vulval induction was scored in L4 animals (46-48 h post-L1) using DIC images<sup>65</sup>. The vulval induction index, the average number of induced VPCs per animal, was calculated by assigning a score of 1 to a VPC that had undergone three rounds of cell divisions, and a score of 0.5 when only one of the two VPC descendants had differentiated.

### Metabolites, nucleoside and fatty acid supplementation

Vitamin B12, metabolites, and nucleosides were added to freshly prepared NGM plates. Vitamin B12 (#V2876, Sigma-Aldrich) was dissolved in water at 1 mM stock concentration and added at a final concentration of 64 nM unless indicated otherwise. All of the metabolites were

directly added to the NGM plates at the following final concentrations: 40 mM for choline (#401757, Sigma-Aldrich), 5 mM for L-methionine (#1.05707, Sigma-Aldrich), and 100 µM for folinic acid (#F7878, Sigma-Aldrich). Uridine (#U3003-50G, Sigma-Aldrich), thymidine #T9250-25G, Sigma-Aldrich), guanosine (#G6264-25G, Sigma-Aldrich), and cytidine (#C4654-5G, Sigma-Aldrich) were also added directly at a final concentration of 1 mM each. Sodium oleate (#O3880, Merk) was likewise added directly at a final concentration of 0.1 and 0.5 mM, and palmitoleic acid (#P9417-100, Merk) at a final concentration of 0.5 and 2.5 mM. For fatty acid supplementation, 1 µM tergitol (NP40S) (#MFCD01779855, Merk) was included in the final mix for solubilization.

### Long-term image acquisition

Long-term imaging experiments were performed as described by Berger et al.<sup>31</sup>. Briefly, *E. coli* OP50 or *Comamonas* DA1877 bacteria were grown overnight at 37 °C in LB medium. Bacterial cultures were then washed three times with S-Basal buffer and concentrated to 1 ml. For *E. coli* OP50, the bacterial culture was re-suspended in 650 µl of Optiprep (#D1556, Sigma-Aldrich) and 332 µl of S-Basal supplemented with 1% Pluronic F127 (#D1556, Sigma-Aldrich) and for *Comamonas* DA1877 in 600 µl of Optiprep and 380 µl of S-Basal supplemented with 1% Pluronic F127. The bacterial suspension was filtered using a 10 µm strainer (pluriS-trainer Mini 10 µm, PluriSelect). Worms were washed three times with S-Basal before loading on the chip. Animals were loaded as described, and immobilized by the microfluidic device channel and two on-chip hydraulic valves<sup>31</sup>. Epifluorescence images were acquired using a 40x oil objective, at an interval of 10 min for germ cell tracking, and 5 min to measure ovulation rates. Wild-type animals were imaged at 20 °C, and *let-60(ga89)* animals at 25 °C, with animals shifted from 20 °C to 25 °C two hours before the start of the experiment. The temperature was controlled (±0.5 °C) by either the room's air conditioning system or a microscope cage incubator (H201-T-UNIT-BL-CRYO and H201-ENCLOSURE-CRYO, Okolab).

### Germ cell tracking

Worm movement during long-term imaging acquisition was corrected using a custom-built MATLAB script. Briefly, sample drift in the z-direction was compensated for each time point assuming linear sample drift within the device channel. Image registration was initially performed on the channel outlining the germ cell membrane, with the derived transformation applied to all other channels. Sample shift in between time points was compensated by visually identifying the gonad's loop region and cropping the original image to a final size of 750×250 pixels from the loop. For each time point, a maximum intensity z-projection of 5-10 slices was created.

Germ cell migration was tracked over 2 h, manually identifying the position of individual germ cells in each projection. Germ cells were tracked using the SYN-4::GFP membrane marker alone for *let-60(ga89)* (*gf<sup>ts</sup>*) or combined with the mCherry::H2B marker for the wild-type. Since germ cells migrate collectively, at least 10 individual cells distributed over the distal gonad arm were tracked per worm, and eight worms per genotype were analyzed.

### RNA interference

RNAi plates were prepared by adding ampicillin (100 µg/ml) and IPTG (1 mM) to the NGM mixture. *E. coli* HT115 RNAi cultures were grown overnight at 37 °C with shaking in LB medium supplemented with ampicillin (100 µg/ml) and tetracycline (10 µg/ml). Before plating the bacteria on the RNAi plates, the overnight liquid culture was diluted 1:1 with fresh LB (supplemented with ampicillin, tetracycline and IPTG) and bacteria were allowed to grow for 3-5 h at 37 °C with shaking (except for the *pmt-2* RNAi clone). 400 µl of bacterial culture were



seeded on each plate, and plates were incubated at 37 °C overnight. RNAi clones were obtained from either the Ahringer RNAi library or the ORFeome RNAi library (both from Source BioScience). All RNAi clones were sequenced before the experiments.

### ERK-nKTR biosensor quantification

MPK-1 activity in the VPCs was measured in mid-L2 larvae (1-cell-stage) using the ERK-nKTR biosensor described in de la Cova et al.<sup>33</sup>. Worms fed on *E. coli* OP50 were imaged 20–23 h post L1 and worms fed on *Comamonas* DA1877 were imaged 19–21 h post L1. The developmental stage was determined by measuring gonad length (35–75 µm in mid-L2). Epifluorescence images were acquired with a 0.25 µm z-step, using a 63x oil objective described as above.

Image analysis was performed using a custom-built MATLAB script. Acquired images were flat-field corrected, after which the in-focus slice was identified using the mCherry::H2B nuclear signal. The images were then z-projected (sum intensity) around the in-focus slice (two slices above and below) and individual VPCs were identified. The red/green fluorescence intensity ratio was calculated for each VPC except P3.p and normalized to the average intensity ratio of all VPCs for each animal.

### Mitotic Index of germ cells

The mitotic zone of dissected gonads was identified by HIM-3 antibody staining (see Gonad dissection and immunostaining for sample preparation). Mitotic cells were counted using Fiji software<sup>66</sup> based on nuclear DAPI staining. The number of pH3-positive M phase nuclei was divided by the total number of mitotic cells to calculate the mitotic index. Images were acquired using a confocal microscope with a 60x objective.

### mRNA isolation and sequencing

For transcriptome analysis by RNAseq, animals were collected from plates seeded with either *E. coli* OP50 or *Comamonas* DA1877 bacteria. For adults, approximately 500 animals (wild-type or *let-60(ga89)* (*gf<sup>ts</sup>*)) were collected. For larval analysis, 1500 animals (wild-type or *let-60(n1046<sup>gf</sup>)*) were collected. Before sample collection, adult worms were transferred to 25 °C for 4 h. Three biological replicates were used per condition. Animals were washed off plates with M9 and snap-frozen in liquid nitrogen. Samples were stored at –80 °C until RNA extraction. Total mRNA extraction was performed for all samples at the same time with trizol, and samples were purified using a Rneasy Mini Kit (QIAGEN). Quality control (Agilent DNF-471 RNA kit), cDNA library preparation, and whole exome sequencing (Illumina Novaseq 6000 system) were performed by the transcriptomics service of the Functional Genomics Center Zurich (FGCZ). Between 15 and 30 million reads were obtained per sample. Bioinformatic and statistical analysis was performed using the FGCZ in-house bioinformatics pipeline SUSHI, using the open-source tools STAR for read alignment and EdgeR for differential gene expression analysis. *p*-values were calculated by paired two-tailed t-tests and corrected with the Benjamini-Hochberg method to control the false discovery rate (FDR).

### Western blot analysis

One hundred worms were washed three times in ice-cold M9 and lysed in 50 µl 2x SDS PAGE buffer (containing 200 mM β-MCE) for 5 min at 95 °C. Genomic DNA was digested by adding 2 µl of Rnase-free Dnase (QIAGEN) for 5 min at room temperature followed by a 5-minute incubation at 95 °C. Samples were kept at 4 °C overnight or –20 °C for longer periods before loading. Before loading, samples were heated to 95 °C, and 15 µl were loaded onto 4–12% gradient polyacrylamide gels.

For MAPK, histone H3, and histone H3 tri methyl K4 quantification, the membrane was treated with transfer buffer containing 20% methanol and blocked with 5% Bovine Serum Albumin (BSA) (#10735086001, Merk) for anti-MAPK and 5% milk for anti-histone

blots. Samples were incubated with anti-dephosphorylated ERK 1&2 (1:2000 dilution, #M8159, Sigma-Aldrich), anti-MAPK (1:10'000 dilution, #T6074, Sigma-Aldrich), anti-histone H3 (1:1000 dilution, #ab1791, Abcam), anti-histone H3 tri methyl K4 (1:1000 dilution, #ab8580, Abcam) and anti-alpha tubulin (1:10'000 dilution, #ab18251, Abcam) at 4 °C overnight. Secondary antibodies HRP conjugated anti-rabbit (1:2000 dilution, #111-035-144, Jackson Immuno Research) and HRP conjugated anti-mouse (1:2000 dilution, #115-035-146, Jackson Immuno Research) were incubated for 2 h at RT. The HRP was detected by incubating the membrane with a chemiluminescence assay SuperSignal West Dura Extended Duration Substrate (#34075, Thermo Fisher Scientific) or SuperSignal West Pico PLUS Chemiluminescent Substrate (#34577, Thermo Fisher Scientific) for 4 min. For MAPK quantification, the membrane was then incubated with sodium azide (1:10) for 1 h to inactivate the HRP-conjugates, and re-stained for alpha Tubulin, followed by incubation with HRP-conjugated secondary antibodies, as well as the chemiluminescence assay.

Signal levels were quantified by measuring band intensities using a built-in tool in Fiji<sup>66</sup>. For MAPK quantification, both total MAPK and phospho-MAPK were normalized to the tubulin loading control. Phospho-MAPK levels were then normalized by total MAPK levels. Uncropped blots are shown in the Western blot raw data file in the Source Data.

### Gonad dissection and immunostaining

Gonad dissection was performed in PBS containing 0.2 mM tetra-misole. Dissected gonads were transferred to siliconized 2 ml Eppendorf tubes and fixed in 4% formaldehyde for 10–60 min. For HIM-3 and pH3 staining, gonads were washed three times with PBS-T (0.05% Tween 20) and permeabilized with 0.5% Triton for 12 min. All gonads were washed three times with 0.05% PBS-T and post-fixed for 5 min at –20 °C in 100% methanol. Dissected gonads were blocked for 2–4 h at room temperature in PGB buffer (PBS-T, 1% BSA, 0.2% gelatin) or 1% BSA. Primary antibodies were diluted in PGB buffer or BSA and incubated overnight at 4 °C, except for pH3 staining, which was incubated for 4 h at room temperature after primary incubation with HIM-3 overnight at 4 °C. Gonads were stained for anti-dephosphorylated ERK 1&2 (1:2000 dilution, #M8159, Sigma-Aldrich), anti-XND-1 (1:2000 dilution, Judith Yanowitz lab), anti-SUN-1 (pS8) (1:700 dilution, Verena Jantsch lab), anti-GLD-1 (1:100 dilution, Judith Kimble lab), anti-HIM-3 (1:500 dilution, Monique Zetka lab) and anti-phospho-histone H3 (Ser10) (1:400 dilution, #05-806, Merk). Secondary antibodies anti-rabbit alexa 568 (1:1000 dilution, #A-10042, Invitrogen), anti-guinea pig alexa 594 (1:1000 dilution, #A-11076, Invitrogen), and anti-mouse alexa 488 (1:1000 dilution, #A-11001, Invitrogen) were incubated for 2 h at room temperature. After secondary incubation, gonads were washed three times with PBS-T. To stain germ cell nuclei DAPI dihydrochloride was added (1:10'000) during the second wash for 10 min. Gonads were mounted in Mowiol (pH 8.3) (#20211125, DABCO) on a glass slide and polymerized overnight at 4 °C. Epifluorescence images were acquired using a 20x air objective as described above.

Before quantification, images were flat-field corrected using a custom-built MATLAB script. To correct for differences in gonad length, the length of the distal arm was normalized such that the distal tip cell (DTC) corresponded to 0 and the loop region to 1. Intensity profiles from the DTC to the loop were measured along each gonad in sum-intensity projections using Fiji<sup>66</sup>.

### Quantification of reporter gene expression

LIN-1::GFP and LIN-39::GFP fluorescence intensity were scored in VPC descendants at the 2-cell Pnp.x stage (mid-L3). Wild-type worms fed with *E. coli* OP50 were imaged 29–30 h post L1; wild-type worms fed with *Comamonas* DA1877 were imaged 27–29 h post L1; *let-60(n1046)* (*gf*) worms fed with *E. coli* OP50 were imaged 32–33 h post-L1; and *let-60(n1046)* (*gf*) worms fed with *Comamonas* DA1877 were imaged 29 h



post LI. Images were acquired on an epifluorescence microscope using a 63x oil-immersion lens, as described above. Before quantification, images were flat-field corrected using a custom-built MATLAB script. VPC nuclei were manually selected and the average fluorescence intensity was measured on an in-focus z-slice using built-in tools in Fiji<sup>66</sup>. Fluorescence intensities for each pair of VPC descendants were averaged. The data for each VPC was normalized to the average value obtained from *E. coli* OP50-fed animals (i.e., P5.p on *Comamonas* DA1877 or *E. coli* OP50 divided by the average of P5.p on *E. coli* OP50).

### Quantification of lipid droplet size

Lipid droplet size was measured using the DHS-3::GFP reporter<sup>40</sup>. Twenty worms were analyzed per condition, and the area of 100 droplets was measured in the posterior section of the intestine of each worm. Images were acquired using a confocal microscope using a 100x oil-immersion lens as described above. The lipid droplet area was measured by manually selecting lipid droplets using Fiji<sup>66</sup>.

### Generation of transgenic *C. elegans*

Somatic rescue of *metr-1* was achieved by creating extrachromosomal arrays in the *metr-1(lf)* background. Microinjections were performed according to<sup>67</sup>. Purified PCR DNA was injected at a concentration of 20 ng/μl. Two co-injection markers were used: 2.5 ng/μl of pCFJ90 (*Pmyo-2>mCherry*)<sup>68</sup> and 100 ng/μl of pBluescript-KS. Three independent lines were generated and characterized.

### Mammalian cell culture

For cell culture experiments, we used the human lung carcinoma epithelial cell line A459, the epidermoid carcinoma A431 cell line (#85090402, Sigma-Aldrich) and four mouse embryonic fibroblast (MEFs) lines: KRAS 4B WT (#RPZ26216, National Cancer Institute), KRAS 4B G12C (#RPZ26186, National Cancer Institute), KRAS 4B G12V (#RPZ26425, National Cancer Institute) and BRAF V600E (#RPZ26275, National Cancer Institute). All cell lines were tested for mycoplasma and kept at 5% CO<sub>2</sub>, 98% relative humidity and 37 °C in a modified DMEM with high glucose and sodium pyruvate (#41966029, Thermo Fisher Scientific), containing 10% fetal calf serum (FCS) (#10500-064, Gibco).

For methionine restriction (MR) experiments, cells were kept in a modified DMEM medium with high glucose, no glutamine, no methionine, and no cysteine (#21013024, Thermo Fisher Scientific). For methionine-positive conditions (no-MR), the medium was supplemented with 200 mM L-Glutamine (#A29168-01, Thermo Fisher Scientific), 200 μM L-Methionine (#J61904.18, Thermo Fisher Scientific), 100 mM L-Cysteine dihydrochloride (#J62292.14, Thermo Fisher Scientific), 400 μM DL-Homocysteine thiolactone hydrochloride (#L09077.14, Thermo Fisher Scientific) and 100 mM sodium pyruvate (#11360070, Thermo Fisher Scientific). For methionine-restricted (MR) conditions, the medium was equally supplemented except for methionine.

### Cell migration assays

G1-arrested A549 cells and MEFs were used for migration assays. Cells were seeded at 80'000 cells/ml density in DMEM medium with or without methionine. After 32 h, cells were exposed to thymidine (1:50) to arrest the cell cycle as described in ref. 43 (first thymidine block). 16 h after, the first thymidine block was released by replacing the medium. 8 h after, cells were transferred to ibidi 2-well silicon culture inserts at a density of 400'000 cells/ml and again exposed to a medium containing thymidine (second thymidine block). Cells remained in the silicon culture inserts for 16 h after which the inserts were removed and pictures were taken every hour for 8 h. For experiments with MEK inhibitors, cells were exposed to 1 μM MEK162 (binimetinib) (#606143.89-9, Selleckchem) and 1 μM LGX818 (ecorafenib) (#1269440-17-6, MedChemExpress) during the first thymidine block. The relative migration was calculated as  $R[\%] = (RO - Rn) / RO \times 100[\%]$ ,

where RO represents the initial open area and Rn represents the remaining open area at each time point. Linear regression was used to determine migration rates shown in the bar graphs (Fig. 7a").

### EGF stimulation and immunostaining of A431 cells

The A431 cell line was used to analyze filopodia formation upon EGF stimulation. Cells were exposed to no-methionine restriction (no-MR, 200 μM methionine + 400 μM homocysteine) or MR (0 μM methionine + 400 μM homocysteine) 24 h before the experiment. Cells were then seeded on glass slides in 24-well plates at 114'000 cells/ml density in no-MR or MR conditions. After 24 h, cells were serum starved and treated with 1 μM MEK162 (binimetinib) and 1 μM LGX818 (ecorafenib). After another 16 h, cells were stimulated with 100 ng/ml of EGF for 10 min at 37 °C<sup>67</sup>, followed by 15 min fixation in 4% PFA at 37 °C. Cells were permeabilized for 5 min with 0.2% Triton X-100% and 0.5% BSA and then blocked for 1 h in 0.5% BSA and 0.2% gelatine. Fixed cells were stained with Phalloidin 568 for 40 min and 0.1 μg/ml DAPI for 5 min. Glass slides were mounted with ProLong Gold Antifade Mountant (Thermo Fisher Scientific).

### Statistical analysis and reproducibility

No statistical method was used to predetermine the sample size. Animals were randomly selected for analysis, and no data were excluded from the analyses. The Investigators were not blinded to allocation during experiments and outcome assessment. Statistical analysis was performed using GraphPad Prism 10. When comparing two samples, an unpaired t-test was used. For multiple samples, data was analyzed by one-way ANOVA followed by Dunnett's multiple comparison test, by two-way ANOVA followed by Tukey's multiple comparison test for parametric data, by two-way ANOVA followed by an uncorrected Fisher's LSD multiple comparison test for interactions between multiple conditions or a Kruskal-Wallis test for non-parametric data followed by a Dunn's test for multiple comparisons. The specific tests used and the number of biological and technical replicates obtained are specified in the figure legends.

### Reporting summary

Further information on research design is available in the Nature Portfolio Reporting Summary linked to this article.

### Data availability

The transcriptomic data generated in this study have been deposited in the European Nucleotide Archive under accession number PRJEB67437. Source data are provided in this paper.

### Code availability

Image processing scripts are available at Zenodo <https://doi.org/10.5281/zenodo.13383462> (<https://www.zenodo.org>).

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Conceptualization: A.C.L. and A.H.; Investigation: A.C.L., S.B., T.K., and A.H.; Formal Analysis: A.C.L., S.B. and N.R.G.; Writing – Original draft: A.C.L. and A.H.; Writing – Review and Editing: A.C.L. and A.H.; Funding acquisition: A.C.L. and A.H.

## Competing interests

The authors declare no competing interests.

## Additional information

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