

Supplementary Information for:

May, A.I., Prescott, M. and Ohsumi, Y.,  
Autophagy facilitates adaptation of budding yeast to respiratory growth by  
recycling serine for one-carbon metabolism

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Supplementary Figures 1 - 12

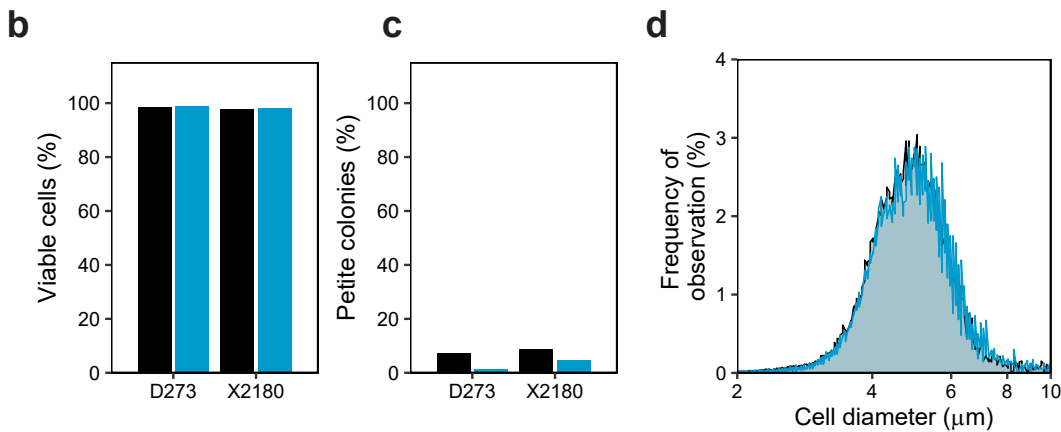
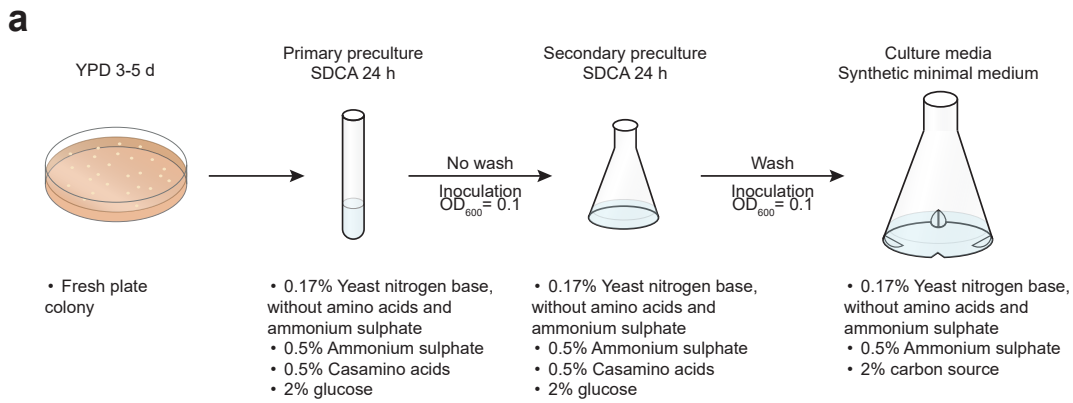
Supplementary Table 1: Antibodies used in this study

Supplementary Table 2: Reagents used in this study

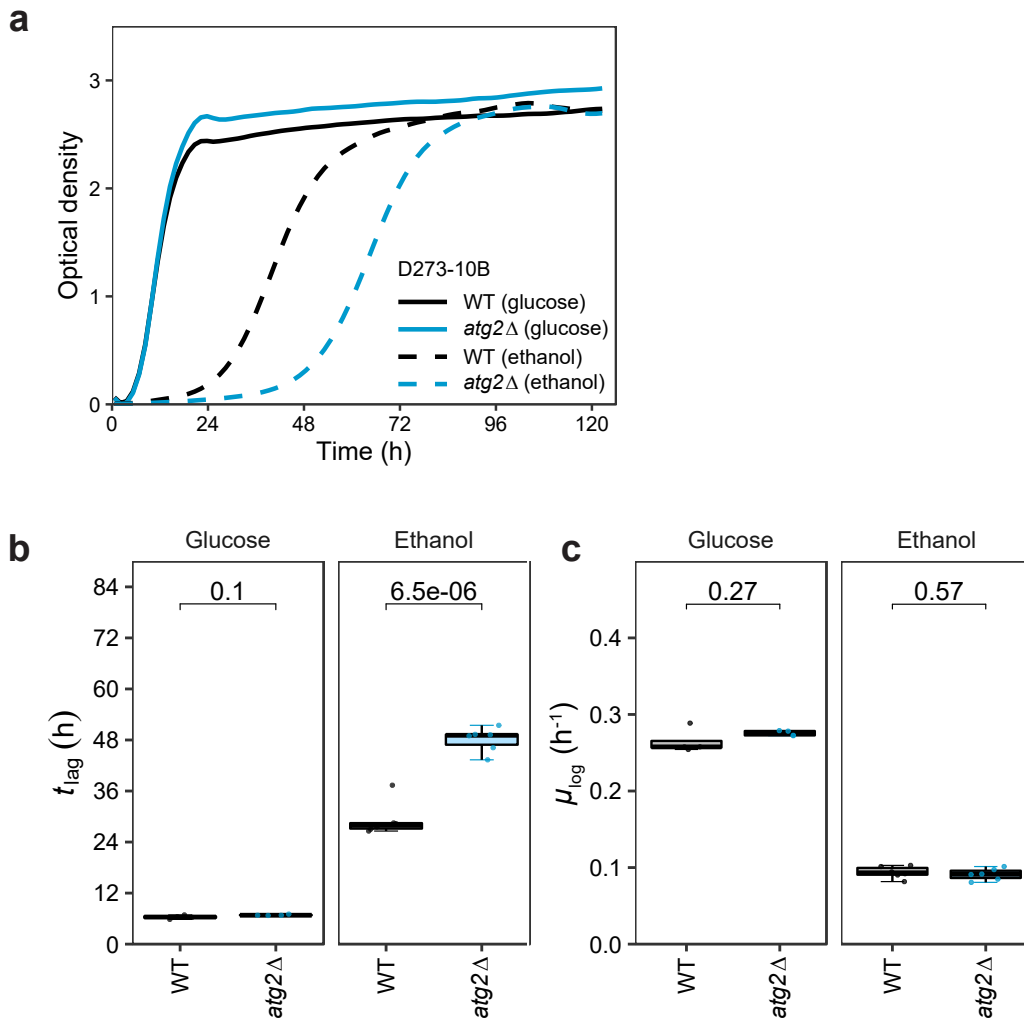
Supplementary Table 3: Strains used in this study

Supplementary Table 4: Plasmids used in this study

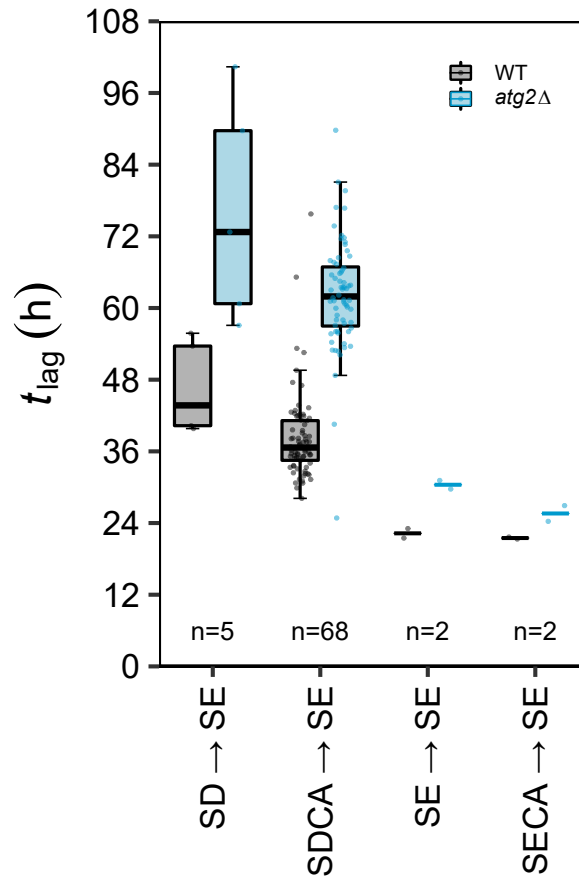
Supplementary Table 5: Oligonucleotides used in this study



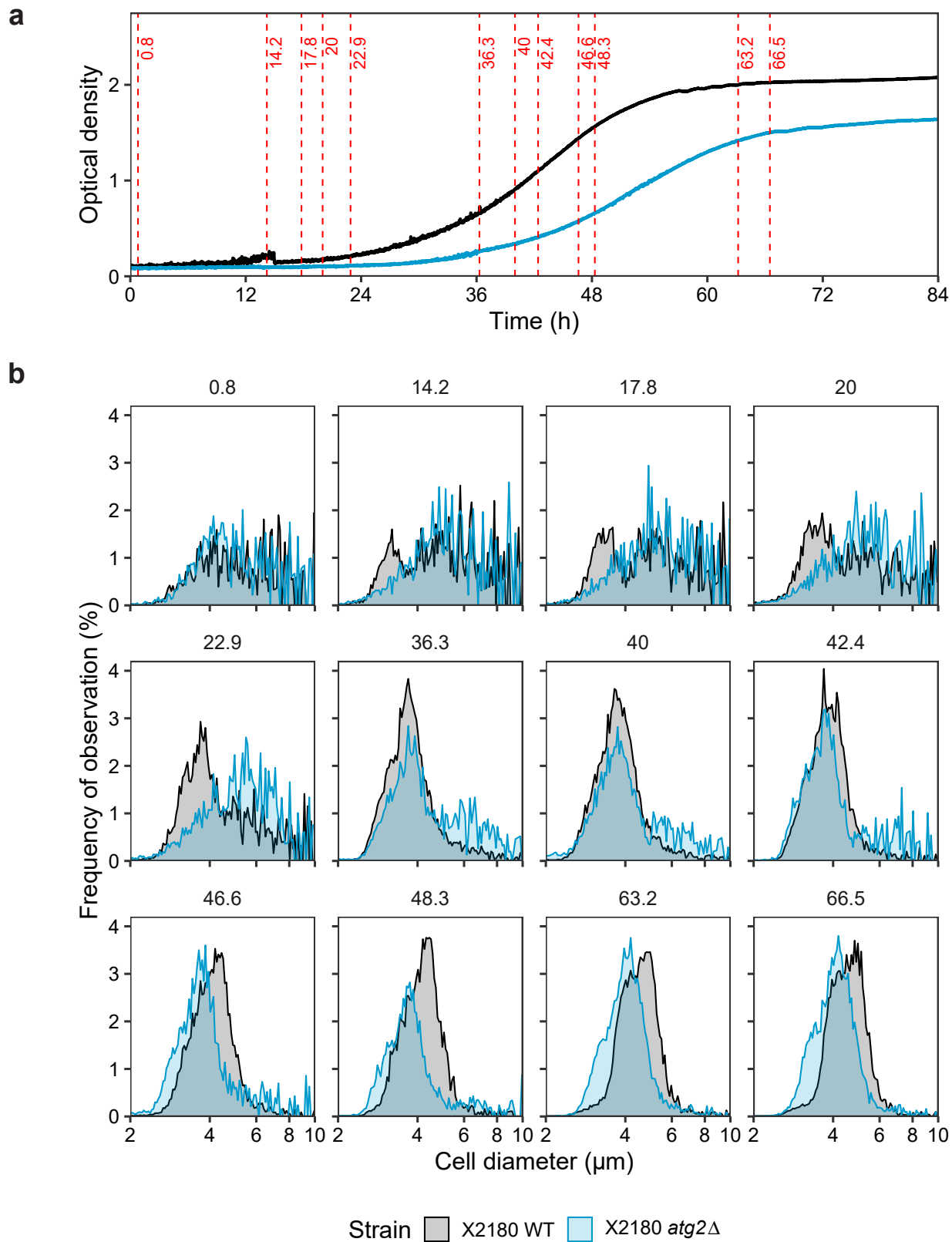
**Supplementary Figure 1.** Overview of the culture regime employed in this study. (a) Cells were taken from fresh (no older than 5-day-old) YPD plates and inoculated into SDCA medium. Following 24 h of culture, cells were diluted to  $OD_{600} = 0.1$  in fresh SDCA media. After a second 24 h incubation in SDCA, cells were washed in and inoculated to subsequent culture media at  $OD_{600} = 0.1$ . (b) Viability and (c) petite frequency of wild-type (black) and *atg2Δ* (blue) cells at the point of inoculation to culture media. (d) Histogram of cell size of wild-type (black) and *atg2Δ* (blue) cells at the point of inoculation to culture media. Error bars = 1 standard deviation. Data are from one (b, d) or two (c) independent experiments.



**Supplementary Figure 2.** Prolonged autophagy mutant respiratory growth  $t_{lag}$  is confirmed in an alternative yeast background strain of distinct pedigree. (a) Growth of WT (black lines) and *atg2Δ* (blue lines) strains from the alternative background D273-10B on synthetic media supplemented with glucose (solid lines) or ethanol (broken lines) as carbon sources. Data are a single representative determination of data presented in (b) and (c). (b) Statistical analysis of  $t_{lag}$  and (c)  $\mu_{log}$  for wild-type and *atg2Δ* cells. Data are from at least five independent experiments Mean growth curves from 3 independent determinations of WT (black) and *atg2Δ* (blue) cell growth on a range of fermentative and respiratory carbon sources. Boxplots in (b) and (c) are presented as median (middle bar), 25th and 75th percentiles (upper and lower limits of the box) and 1.5 \* interquartile range (whiskers). Sample sizes:  $n = 4$  (glucose) or  $n = 6$  (ethanol). Indicated p-values were calculated using the two-sided Student's t-test with Welch modification. Error bars = 1 standard deviation.



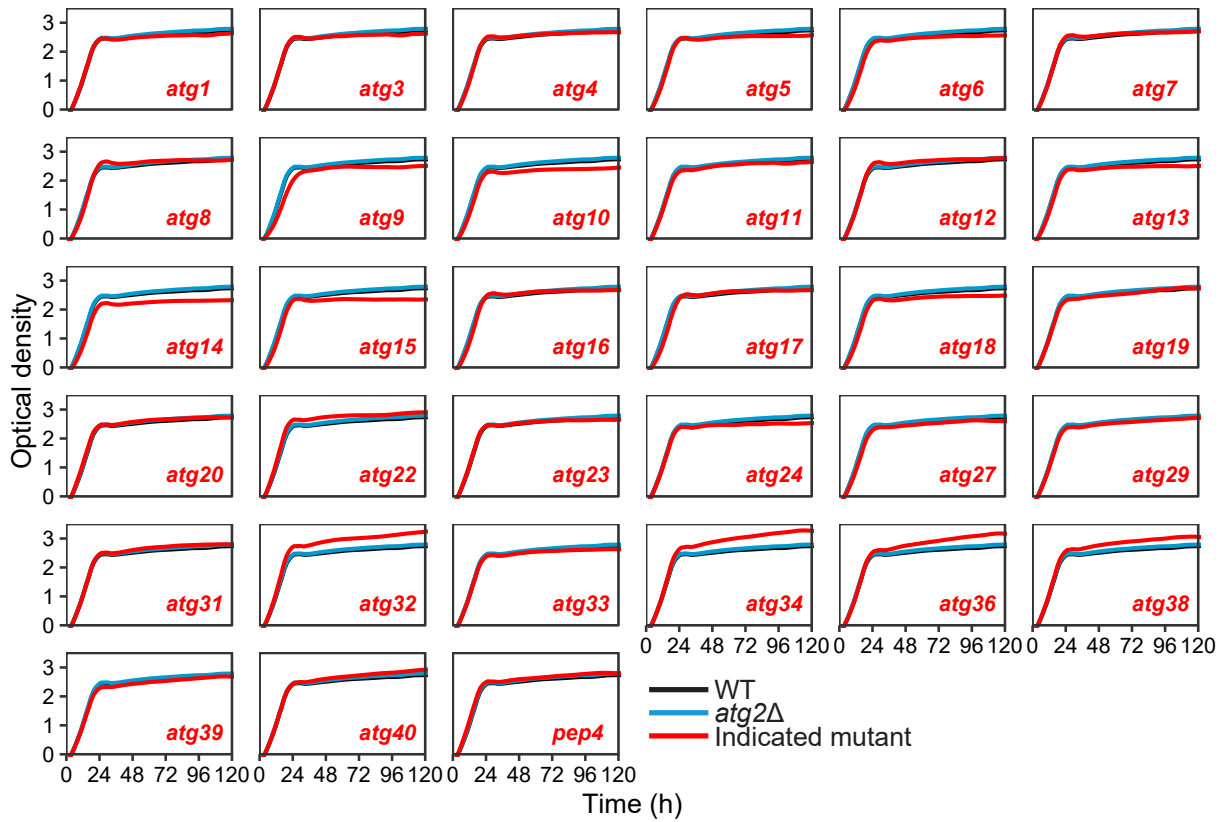
**Supplementary Figure 3.** The shift from fermentative to non-fermentative media, and not removal of casamino acids, is implicated in the prolonged  $t_{lag}$  of *atg2Δ* cells. Shown are boxplots of WT (black) and *atg2Δ* (blue)  $t_{lag}$  for cells cultured under the indicated conditions. As  $n < 3$  for cells pre-grown on ethanol media, these data are presented as individual determinations with a bar indicating the mean  $t_{lag}$ . SD, synthetic glucose media without casamino acids, SDCA, synthetic glucose media with casamino acids, SE, synthetic ethanol media without casamino acids, SECA, synthetic ethanol media with casamino acids. Data from at least three ( $n = 3$ ) independent experiments are shown. Error bars = 1 standard deviation. Boxplots are shown as median (middle bar), 25th and 75th percentiles (upper and lower limits of the box) and  $1.5 \times$  interquartile range (whiskers).



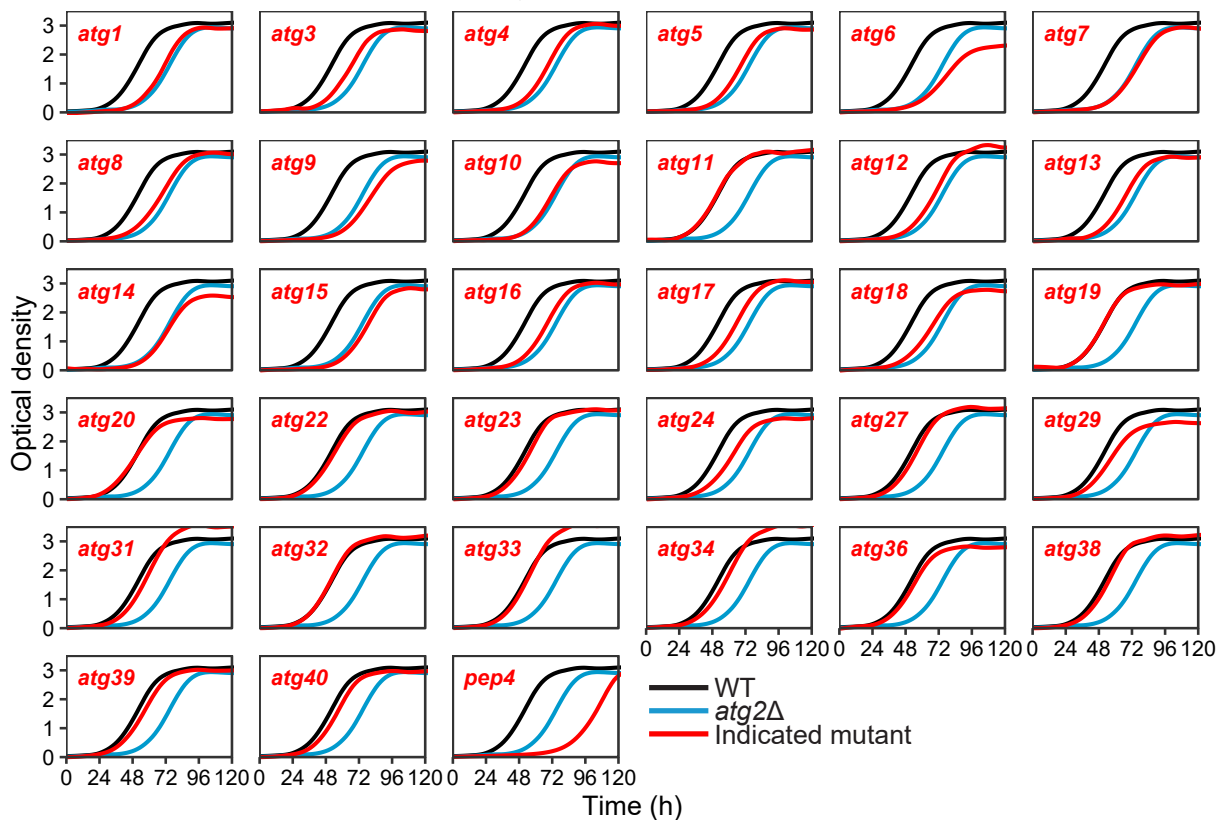
**Supplementary Figure 4.** Changes in cell size reflect delayed onset of autophagy mutant growth on ethanol media. (a) Growth of WT (black) and *atg2* $\Delta$  (blue) cells on ethanol media. Sampling points are indicated by dotted red lines. (b) Cell size was assessed by particle counter at time points indicated in (a) and are presented as histograms of particle size frequency. Black, wild-type cells, blue, *atg2* $\Delta$  cells.

**a**

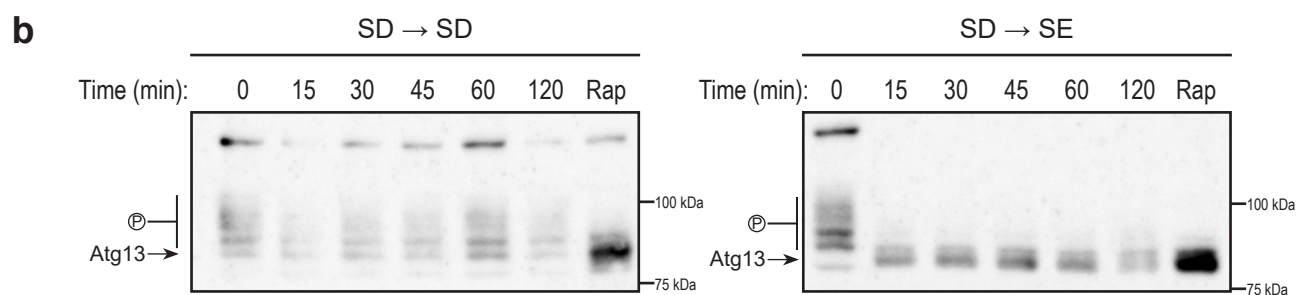
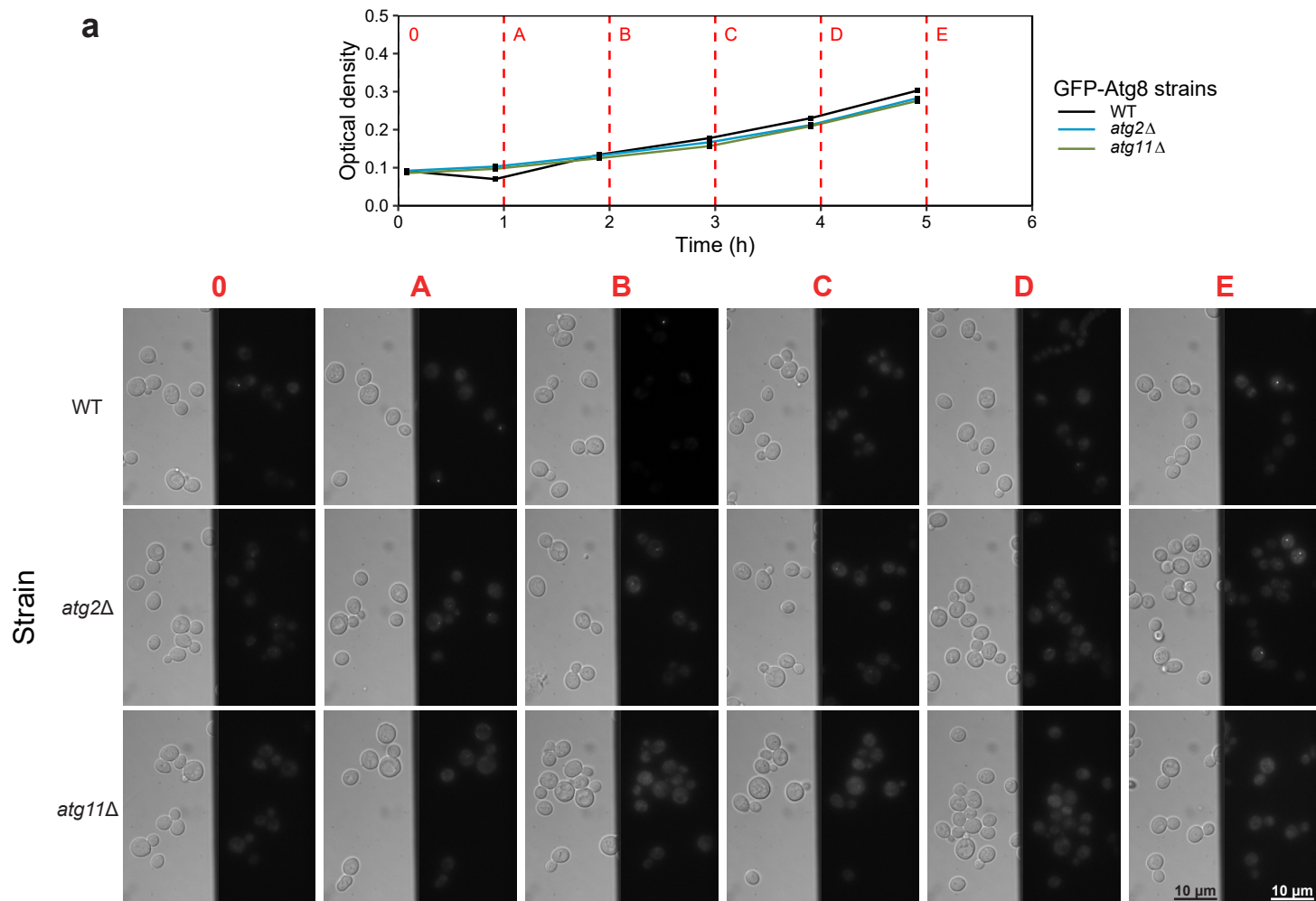
## Glucose growth of deletion strains

**b**

## Ethanol growth of deletion strains



**Supplementary Figure 5.** Bulk autophagy is implicated in prolonged  $t_{lag}$  phenotype. (a) Growth of core and non-core *ATG* strains on glucose media. Growth curves from  $n = 1$  experiment are shown. (b) Growth of core and non-core *ATG* strains on ethanol media. These data were quantified and are shown in Fig. 2. Black lines, wild-type, blue lines, *atg2Δ*, red lines, indicated mutant strain.



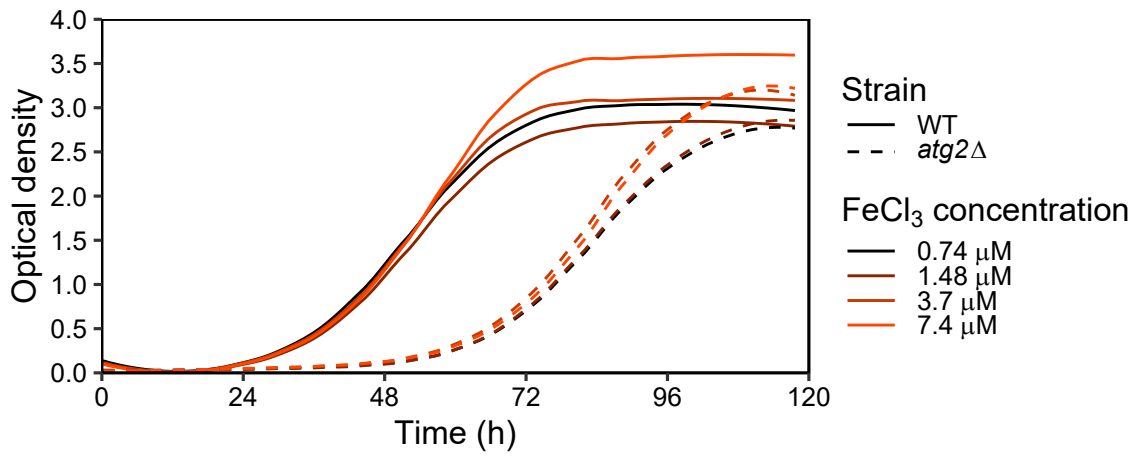
**Supplementary Figure 6.** Autophagy is not induced during the lag phase of fermentative growth.

(a) WT (black), *atg2Δ* (blue) and *atg11Δ* (green) cells expressing GFP-Atg8 were inoculated to SD media under the same conditions as shown in Fig. 3b and observed by microscopy at the indicated time points (broken red lines). Data are from a single experiment and were reproduced twice.

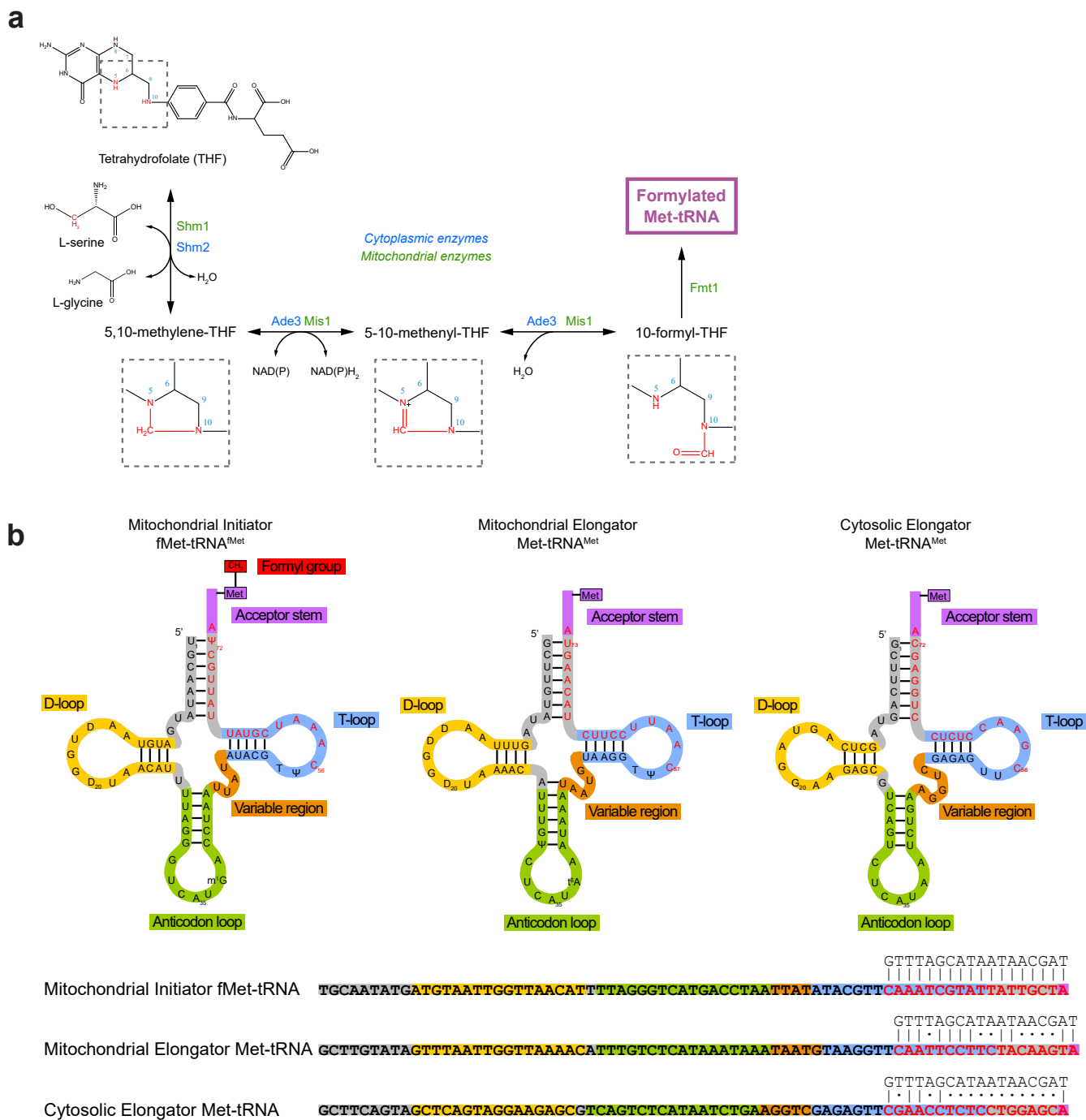
(b) The phosphorylation of Atg13 was determined by western blotting in WT cells shifted from synthetic glucose media not containing casamino acids (SD) to SD or synthetic ethanol media not containing casamino acids (SE). Data are from a single experiment and were reproduced three times.





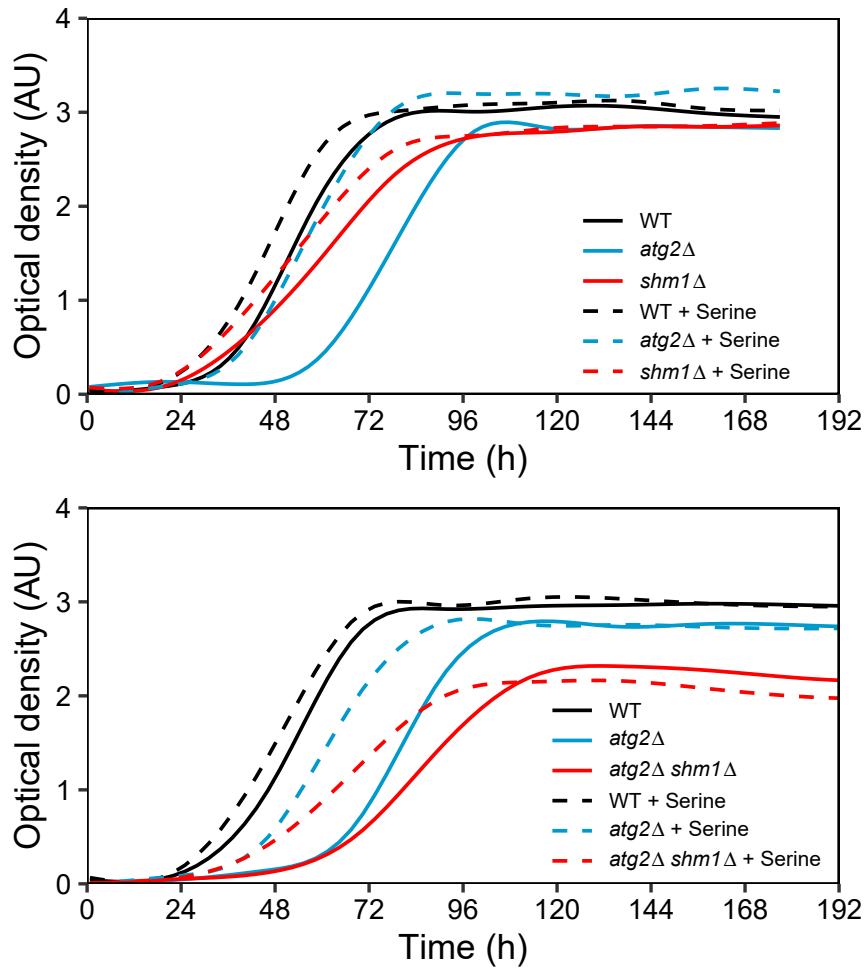


**Supplementary Figure 8.** Prolonged *atg2Δ*  $t_{lag}$  is not alleviated by the supplementation of iron. Ferric chloride was added to respiratory media at the indicated concentrations and WT (solid lines) and *atg2Δ* (broken lines) growth were determined. The average of  $n = 3$  growth experiments is shown.

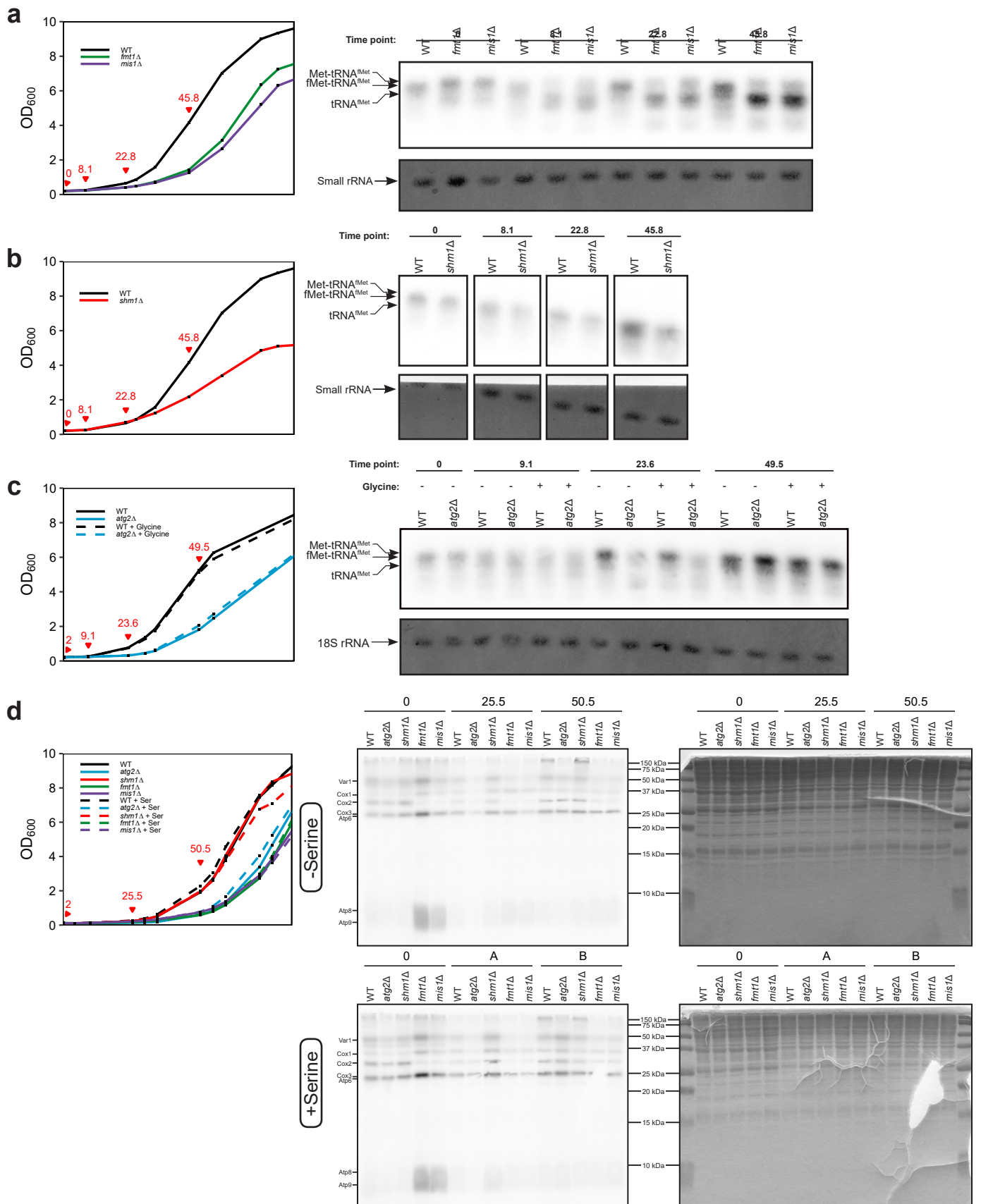


**Supplementary Figure 9.** Overview of the strategy used to determine formylation of Met-tRNA<sup>Met</sup>.

(a) detailed overview of the fate of serine in the reactions of one-carbon metabolism. The phenomenon addressed in this paper concerns the mitochondrial reactions (enzymes catalysing these reactions are indicated in green). (b) A comparison of Met-tRNA species found in yeast. The sequence of initiator tRNA in mitochondria is sufficiently different as to allow for its specific probing by northern blotting.

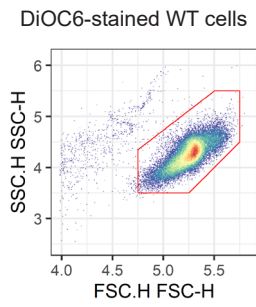


**Supplementary Figure 10.** The deletion of *SHM1* strongly affects  $\mu_{\log}$  but not the  $t_{\text{lag}}$  of ethanol-grown cells. WT (black), *atg2Δ* (blue) and *shm1Δ* (red) cells were grown on ethanol media and growth in the absence (solid lines) or presence (broken lines) of serine determined. The average of  $n = 2$  growth experiments is shown.

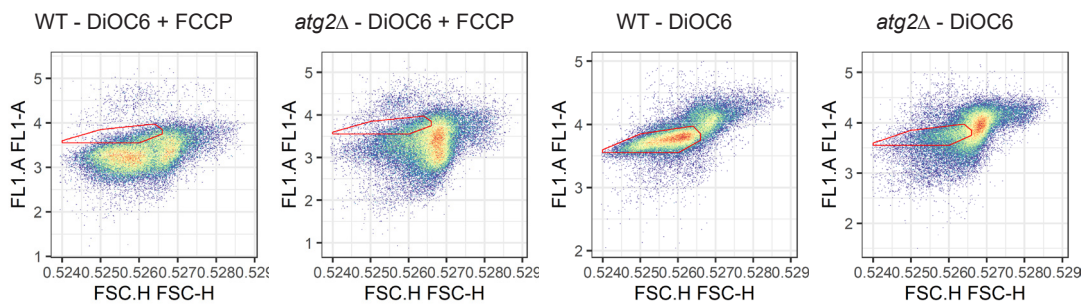


**Supplementary Figure 11.**  $t_{lag}$  duration correlates with initiator tRNA formylation and mitochondrial protein expression in a range of one carbon metabolism mutant strains. (a) Formylation of WT, *fmt1* $\Delta$  and *mis1* $\Delta$  initiator tRNA under the conditions shown in Fig. 6c. (b) Formylation of WT and *shm1* $\Delta$  initiator tRNA under the conditions shown in Fig. 6c. (c) Formylation of WT and *atg2* $\Delta$  initiator tRNA under the conditions shown in Fig. 6c, but with glycine supplemented to media instead of serine. (d) Mitochondrial protein expression, as determined by the incorporation of  $^{35}\text{S}$ -labelled cysteine and methionine, was determined by autoradiography under the same conditions as shown in Fig. 6d. Samples were collected as indicated in the growth curves at the left of each figure. Data are single determinations from individual experiments.

Gating for cells:



Gating for cells with high  $\Delta\Psi_m$ :



**Supplementary Figure S12.** Gating strategy used to determine cells with high mitochondrial membrane potential ( $\Delta\Psi_m$ ). First, non-cell debris were removed by gating for cells by forward and side scatter. Following this, the proportion of highly fluorescent newly-emerged cells was identified by gating for the indicated region in the FL1-A channel. The proportion of cells falling within this region are shown in Fig. 5b. The logicle transformation was used for forward and side scatter data. The data shown in this figure are from the  $t = 26$  h time point of the experiment shown in Fig. 5.

SUPPLEMENTARY INFORMATION

Supplementary Table 1: Antibodies used in this study

Resource	Supplier	Reference number
Mouse monoclonal Anti-GFP, clones 7.1 and 13.1	Roche	11814460001
Rabbit polyclonal Anti-Ape1	Ohsumi lab stock	NA
Mouse monoclonal Anti- $\beta$ -actin	Wako	011-24554
Rabbit polyclonal Anti-Atg13	Ohsumi lab stock	NA

Supplementary Table 2: Reagents used in this study

Reagent	Supplier	Reference number
Difco Yeast Nitrogen Base w/o Amino Acids and Ammonium Sulfate	BD	233520
Ammonium sulfate	Nacalai	02620-04
Bacto Casamino acids	BD	223050
Bacto Yeast extract	BD	212750
Bacto Peptone	BD	211677
Glucose	Nacalai	16806-54
Ethanol	Wako	14713-53
Galactose	Wako	075-00035
Sucrose	Wako	193-09545
Glycerol	Wako	075-00616
Sodium acetate	Wako	190-01075
Raffinose	Wako	184-00015
Lactate	Wako	195-02305
L-Glutamic Acid	Wako	070-00502
L(+)-Glutamine	Wako	074-00522
Glycine	Wako	073-00732
L-Serine	Wako	199-00402
D-Serine	Wako	191-08821
Adecanol	Adeka	LG-109
FCCP	Sigma	C2920
Urea	Sigma	U6504
TRIzol	Life Tech	15596026
Methyl blue	Sigma	M6900
DiOC6	Molecular probes	D273
Triphenyltetrazolium chloride	Sigma	T8877
Adenosine 5'-triphosphate, [ $\gamma$ - $^{32}$ P]-	PerkinElmer	NEG502Z
Met- $^{35}$ S label (stabilized)	ARC Inc.	ARS0110A
T4 Polynucleotide Kinase	Toyobo	PNK-111
KOD -plus- Ver. 2	Toyobo	KOD-211
PerfectHyb Plus	Sigma	H7033

Supplementary Table 3: Strains used in this study

Strain	Source	Reference number
<i>S. cerevisiae</i> X2180-1B ( <i>MAT<math>\alpha</math> SUC2 mal mel gal2 CUP1</i> )	Lab stock	ATCC#204505
<i>S. cerevisiae</i> X2180-1B <i>atg1</i> $\Delta$ ( <i>MAT<math>\alpha</math> SUC2 mal mel gal2 CUP1 atg1<math>\Delta</math>::KANMX</i> )	This study	yAM150
<i>S. cerevisiae</i> X2180-1B <i>atg2</i> $\Delta$ ( <i>MAT<math>\alpha</math> SUC2 mal mel gal2 CUP1 atg1<math>\Delta</math>::KANMX</i> )	This study	yAM151
<i>S. cerevisiae</i> X2180-1B <i>atg3</i> $\Delta$ ( <i>MAT<math>\alpha</math> SUC2 mal mel gal2 CUP1 atg1<math>\Delta</math>::KANMX</i> )	This study	yAM306
<i>S. cerevisiae</i> X2180-1B <i>atg4</i> $\Delta$ ( <i>MAT<math>\alpha</math> SUC2 mal mel gal2 CUP1 atg1<math>\Delta</math>::KANMX</i> )	This study	yAM308
<i>S. cerevisiae</i> X2180-1B <i>atg5</i> $\Delta$ ( <i>MAT<math>\alpha</math> SUC2 mal mel gal2 CUP1 atg1<math>\Delta</math>::KANMX</i> )	This study	yAM310
<i>S. cerevisiae</i> X2180-1B <i>atg6</i> $\Delta$ ( <i>MAT<math>\alpha</math> SUC2 mal mel gal2 CUP1 atg1<math>\Delta</math>::KANMX</i> )	This study	yAM312
<i>S. cerevisiae</i> X2180-1B <i>atg7</i> $\Delta$ ( <i>MAT<math>\alpha</math> SUC2 mal mel gal2 CUP1 atg1<math>\Delta</math>::KANMX</i> )	This study	yAM314
<i>S. cerevisiae</i> X2180-1B <i>atg8</i> $\Delta$ ( <i>MAT<math>\alpha</math> SUC2 mal mel gal2 CUP1 atg1<math>\Delta</math>::KANMX</i> )	This study	yAM316
<i>S. cerevisiae</i> X2180-1B <i>atg9</i> $\Delta$ ( <i>MAT<math>\alpha</math> SUC2 mal mel gal2 CUP1 atg1<math>\Delta</math>::KANMX</i> )	This study	yAM443
<i>S. cerevisiae</i> X2180-1B <i>atg10</i> $\Delta$ ( <i>MAT<math>\alpha</math> SUC2 mal mel gal2 CUP1 atg1<math>\Delta</math>::KANMX</i> )	This study	yAM318
<i>S. cerevisiae</i> X2180-1B <i>atg11</i> $\Delta$ ( <i>MAT<math>\alpha</math> SUC2 mal mel gal2 CUP1 atg1<math>\Delta</math>::KANMX</i> )	This study	yAM153
<i>S. cerevisiae</i> X2180-1B <i>atg12</i> $\Delta$ ( <i>MAT<math>\alpha</math> SUC2 mal mel gal2 CUP1 atg1<math>\Delta</math>::KANMX</i> )	This study	yAM320
<i>S. cerevisiae</i> X2180-1B <i>atg13</i> $\Delta$ ( <i>MAT<math>\alpha</math> SUC2 mal mel gal2 CUP1 atg1<math>\Delta</math>::KANMX</i> )	This study	yAM322
<i>S. cerevisiae</i> X2180-1B <i>atg14</i> $\Delta$ ( <i>MAT<math>\alpha</math> SUC2 mal mel gal2 CUP1 atg1<math>\Delta</math>::KANMX</i> )	This study	yAM340
<i>S. cerevisiae</i> X2180-1B <i>atg15</i> $\Delta$ ( <i>MAT<math>\alpha</math> SUC2 mal mel gal2 CUP1 atg1<math>\Delta</math>::KANMX</i> )	This study	yAM342

<i>S. cerevisiae</i> X2180-1B <i>atg16</i> $\Delta$ ( <i>MAT<math>\alpha</math> SUC2 mal mel gal2 CUP1 atg1<math>\Delta</math>::KANMX</i> )	This study	yAM344
<i>S. cerevisiae</i> X2180-1B <i>atg17</i> $\Delta$ ( <i>MAT<math>\alpha</math> SUC2 mal mel gal2 CUP1 atg1<math>\Delta</math>::KANMX</i> )	This study	yAM346
<i>S. cerevisiae</i> X2180-1B <i>atg18</i> $\Delta$ ( <i>MAT<math>\alpha</math> SUC2 mal mel gal2 CUP1 atg1<math>\Delta</math>::KANMX</i> )	This study	yAM348
<i>S. cerevisiae</i> X2180-1B <i>atg19</i> $\Delta$ ( <i>MAT<math>\alpha</math> SUC2 mal mel gal2 CUP1 atg1<math>\Delta</math>::KANMX</i> )	This study	yAM350
<i>S. cerevisiae</i> X2180-1B <i>atg20</i> $\Delta$ ( <i>MAT<math>\alpha</math> SUC2 mal mel gal2 CUP1 atg1<math>\Delta</math>::KANMX</i> )	This study	yAM352
<i>S. cerevisiae</i> X2180-1B <i>atg22</i> $\Delta$ ( <i>MAT<math>\alpha</math> SUC2 mal mel gal2 CUP1 atg1<math>\Delta</math>::KANMX</i> )	This study	yAM358
<i>S. cerevisiae</i> X2180-1B <i>atg23</i> $\Delta$ ( <i>MAT<math>\alpha</math> SUC2 mal mel gal2 CUP1 atg1<math>\Delta</math>::KANMX</i> )	This study	yAM356
<i>S. cerevisiae</i> X2180-1B <i>atg24</i> $\Delta$ ( <i>MAT<math>\alpha</math> SUC2 mal mel gal2 CUP1 atg1<math>\Delta</math>::KANMX</i> )	This study	yAM360
<i>S. cerevisiae</i> X2180-1B <i>atg27</i> $\Delta$ ( <i>MAT<math>\alpha</math> SUC2 mal mel gal2 CUP1 atg1<math>\Delta</math>::KANMX</i> )	This study	yAM362
<i>S. cerevisiae</i> X2180-1B <i>atg29</i> $\Delta$ ( <i>MAT<math>\alpha</math> SUC2 mal mel gal2 CUP1 atg1<math>\Delta</math>::KANMX</i> )	This study	yAM364
<i>S. cerevisiae</i> X2180-1B <i>atg31</i> $\Delta$ ( <i>MAT<math>\alpha</math> SUC2 mal mel gal2 CUP1 atg1<math>\Delta</math>::KANMX</i> )	This study	yAM366
<i>S. cerevisiae</i> X2180-1B <i>atg32</i> $\Delta$ ( <i>MAT<math>\alpha</math> SUC2 mal mel gal2 CUP1 atg1<math>\Delta</math>::KANMX</i> )	This study	yAM162
<i>S. cerevisiae</i> X2180-1B <i>atg33</i> $\Delta$ ( <i>MAT<math>\alpha</math> SUC2 mal mel gal2 CUP1 atg1<math>\Delta</math>::KANMX</i> )	This study	yAM368
<i>S. cerevisiae</i> X2180-1B <i>atg34</i> $\Delta$ ( <i>MAT<math>\alpha</math> SUC2 mal mel gal2 CUP1 atg1<math>\Delta</math>::KANMX</i> )	This study	yAM370
<i>S. cerevisiae</i> X2180-1B <i>atg36</i> $\Delta$ ( <i>MAT<math>\alpha</math> SUC2 mal mel gal2 CUP1 atg1<math>\Delta</math>::KANMX</i> )	This study	yAM372
<i>S. cerevisiae</i> X2180-1B <i>atg38</i> $\Delta$ ( <i>MAT<math>\alpha</math> SUC2 mal mel gal2 CUP1 atg1<math>\Delta</math>::KANMX</i> )	This study	yAM374
<i>S. cerevisiae</i> X2180-1B <i>atg39</i> $\Delta$ ( <i>MAT<math>\alpha</math> SUC2 mal mel gal2 CUP1 atg1<math>\Delta</math>::KANMX</i> )	This study	yAM376



<i>S. cerevisiae</i> X2180-1B <i>atg40</i> Δ ( <i>MATα SUC2 mal mel gal2 CUP1 atg1Δ::KANMX</i> )	This study	yAM378
<i>S. cerevisiae</i> X2180-1B <i>pep4</i> Δ ( <i>MATα SUC2 mal mel gal2 CUP1 pep4Δ::KANMX</i> )	This study	yAM154
<i>S. cerevisiae</i> X2180-1B <i>fnt1</i> Δ <i>atg2</i> Δ ( <i>MATα SUC2 mal mel gal2 CUP1 fnt1Δ::natNT2 atg2Δ::KANMX</i> )	This study	yAM237
<i>S. cerevisiae</i> X2180-1B <i>mis1</i> Δ <i>atg2</i> Δ ( <i>MATα SUC2 mal mel gal2 CUP1 mis1Δ::natNT2 atg2Δ::KANMX</i> )	This study	yAM440
<i>S. cerevisiae</i> X2180-1B <i>mis1</i> Δ ( <i>MATα SUC2 mal mel gal2 CUP1 mis1Δ::natNT2</i> )	This study	yAM461
<i>S. cerevisiae</i> X2180-1B <i>fnt1</i> Δ ( <i>MATα SUC2 mal mel gal2 CUP1 fnt1Δ::natNT2</i> )	This study	yAM463
<i>S. cerevisiae</i> X2180-1B GFP-Atg8 ( <i>MATα SUC2 mal mel gal2 CUP1 atg8Δ::GFP - ATG8::hphNT1</i> )	This study	yAM158
<i>S. cerevisiae</i> X2180-1B <i>atg2</i> Δ GFP-Atg8 ( <i>MATα SUC2 mal mel gal2 CUP1 atg8Δ::GFP - ATG8::hphNT1 atg2Δ::KANMX</i> )	This study	yAM159
<i>S. cerevisiae</i> X2180-1B <i>atg11</i> Δ GFP-Atg8 ( <i>MATα SUC2 mal mel gal2 CUP1 atg8Δ::GFP - ATG8::hphNT1 atg11Δ::KANMX</i> )	This study	yAM160
<i>S. cerevisiae</i> D273-10B ( <i>MATα mal</i> )	Trevor Lithgow	yAM086
<i>S. cerevisiae</i> D273-10B <i>atg2</i> Δ ( <i>MATα mal atg2Δ::KANMX</i> )	This study	yAM096
<i>S. cerevisiae</i> D273-10B <i>atg11</i> Δ ( <i>MATα mal atg11Δ::KANMXMX</i> )	This study	yAM100

Supplementary Table 4: Plasmids used in this study

Name	Type	Source	Description
pFA6a-kanMX6	Plasmid	Janke et al. <sup>51</sup>	Generic plasmid for PCR-mediated generation of targeted deletion cassettes
pFA6a-natNT2	Plasmid	Janke et al. <sup>51</sup>	Generic plasmid for PCR-mediated generation of targeted deletion cassettes
pRS303-GFP-ATG8	Plasmid	Suzuki, et al. <sup>50</sup>	C-terminally tagged Atg8 for genomic integration following amplification of cassette by PCR.

Supplementary Table 5: Oligonucleotides used in this study

Name	Source	Sequence
ATG1_S1	This study	tttgaagctaccccatattttcaaatctcttttacaacaccagacgagaaattaagaa aatgCGTACGCTGCAGGTCGAC
ATG1_S2	This study	aagatacttgaaaatatagcaggtcattgtacttaataagaaaacatattatgcat cacATCGATGAATTCGAGCTCG
ATG1_8F	This study	taatataataaaaacataaggc
ATG1_5R	This study	caggataaaaagagctgg
ATG2_S1	This study	gcataaagattaagcaaatagaggaacccttttttttatttcgatacaatgC GTACGCTGCAGGTCGAC
ATG2_S2	This study	cggccgaataattgccacaggtgcagctctagcaacataaactgctcggcgct cggcccATCGATGAATTCGAGCTCG
ATG2_VerF-216	This study	cgcgataatgaatgcaagg
ATG2_VerR+96	This study	cattttggcttcccacc
ATG3_S1	This study	tagaagttaggaacaaagaagtacaaaggagtaaatacaattttattatcCGTA CGCTGCAGGTCGAC
ATG3_S2	This study	ctgtttttgaccacctggcttgagctaatagtgaaaaacacaaatttATCGA TGAATTCGAGCTC
ATG3_2F	This study	tagtagatttctatttttc
ATG3_3R	This study	aatattcataatttatgta
ATG4_S1	This study	ttagtagatgaagaatggacgacttcttatcacgtataggagtgatatacCGTA CGCTGCAGGTCGAC
ATG4_S2	This study	gaatatataaaacaagtatatatgcttatgaactagtgaaattccttacaATCGA TGAATTCGAGCTC
ATG4_VerF-218	This study	cacttctcctctagaaactc
ATG4_VerR+244	This study	ggcattttggactcgcac
ATG5_S1	This study	ttgttctttgggttctagaagaacggagataggaaacctatgatgtaagtCGTAC GCTGCAGGTCGAC
ATG5_S2	This study	tattttctgcgatatttgaatgacacttttaaatgcgtatataacagctcATCGAT GAATTCGAGCTC
ATG5_2F	This study	atgatcgaagttttgtccgg
ATG5_3R	This study	aaagtatcaacagctctcga
ATG6_S1	This study	gccaaacatgcaactaccctgcaactagaccatctttagaaggcttagtCG TACGCTGCAGGTCGAC
ATG6_S2	This study	gtggaaaaagaattttccctttatcacatttatgaaaaatgcatttatgaactac ATCGATGAATTCGAGCTCG
ATG6_VerF-185	This study	ggactgggtgataatcgaa
ATG6_VerR+170	This study	ctggcatatcatctgatgaaag
ATG7_S1	This study	gctacttagataactaaagttcattatatttcaacaaatataagataatcaagaataaa atgCGTACGCTGCAGGTCGAC
ATG7_S2	This study	ggaaagtggcaccacaatatgtaccaatgctattatgcaaaatattaagcaatct catcATCGATGAATTCGAGCTCG
ATG7_VerF-196	This study	ggcaagcaagatttttcatttg
ATG7_VerR+261	This study	ggtagctttaccgacacttt
ATG8_S1	This study	tgataagagaatctaataattgtaaagttgagaaaaataataaaaaataaactag agacCGTACGCTGCAGGTCGAC
ATG8_S2	This study	cctataattcgaatttagatgtaacgcttcatttctttcatataaaaagactacctgcc ATCGATGAATTCGAGCTCG
ATG8_VerF-171	This study	gcacatgatataatcacc

ATG8_VerR+207	This study	gctcacattgtctccaatac
ATG9_S1	This study	gcctgaaatatcaaaatcacggaattattaggttatggagagagatgaataccagt taccCGTACGCTGCAGGTCGAC
ATG9_S2	This study	gccttatctccgacgtcagacttcttgaataactctttaacaagtctaagacacca cccATCGATGAATTCGAGCTCG
ATG9_6F	This study	cccagttcggaaactttagggg
ATG9_5R	This study	attaggcttctcagagac
ATG10_S1	This study	gaagagaacaccatgaaaaaaaaaaaaagggctaaaaaacagaattatcag acttgatCGTACGCTGCAGGTCGAC
ATG10_S2	This study	atatatatataatttatacatagatgattgcatagtgttttaaaaagctttctaggttaag ATCGATGAATTCGAGCTCG
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ATG10_VerR	This study	ccgacacatcatcagaaga
ATG11_S1	This study	attattttagtgactgttgttgcggaaagtacttctttttttttatacatcatgC GTACGCTGCAGGTCGAC
ATG11_S2	This study	gttaaatagatacataaataaacttgcatttgtgacaaacgtttagcactgttcaa acATCGATGAATTCGAGCTCG
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ATG11_6R	This study	cctttccagagatgaggaaaattcaccagg
ATG12_S1	This study	aacgtacatccctaactgtatattctacagtagagtgaaccaatgacagtCGTA CGCTGCAGGTCGAC
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ATG12_1F	This study	GTGATTTTCTTTTTTCAGTGT
ATG12_2R	This study	TCAAACACTTTCGATGAAAT
ATG13_S1	This study	cctccaggctcaagtctgaaaagaagcagaacatacagcccgggtgaatagc atgagtcCGTACGCTGCAGGTCGAC
ATG13_S2	This study	gattattttcttagttgtgccctttaaataaaactttaccatttttaacctttagA TCGATGAATTCGAGCTCG
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ATG14_S2	This study	ctgactacatgcaactttatacacacggcaggaaaaaaagtgcgcaacttagcct accacgATCGATGAATTCGAGCTCG
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ATG17_VerR+268	This study	ccaacacagattcacagc
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ATG18_S2	This study	gtgtatcggttgacgtacggaaggcagcgagacacttccgtgatcaatcca taagATCGATGAATTCGAGCTCG
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ATG20_VerR+273	This study	gcgatggagcataatcatct
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ATG24_VerR+207	This study	ggaagaacaatgcaaaatgttt
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ATG29_VerR+249	This study	gcctcagacagcagagaata
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ATG32_S1	This study	gataagcaatattgaagtcctaatacacaagcaaaaaaatctgccaggaacagtaaacaCGTACGCTGCAGGTCGAC
ATG32_S2	This study	aacagaagtgatagtaaaaaagtgagtaggaacgtgtatgtttgtatattggaaaaaggATCGATGAATTCGAGCTCG
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ATG32_2R	This study	AATTTGCTAACAGAAGAAAT
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ATG34_S2	This study	gttaaataagtactatagccaaagaactggaagaatataaaaaagcatttaATCGATGAATTCGAGCTCG

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ATG38_S1	This study	gttcgataattcaacgacagttataactaacttgggtgatggaatgCGTACGCT GCAGGTCGAC
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ATG38_VerF-97	This study	gcgtaatggaacgctct
ATG38_VerR+247	This study	gcaccttgattccgtttc
ATG39_S1	This study	cgataatagagactagtaaaacagtcgagttgctggacctaataatgCGTACG CTGCAGGTCGAC
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ATG39_VerR+247	This study	gagatggatagcatacgttatg
ATG40_S1	This study	cctaacgtttcttctgctgtgcttccactccacatagaaaactaatgCGTACGC TGCAGGTCGAC
ATG40_S2	This study	ggtacctcatagactaccattatggtaaaatggaaaaactattctaATCGAT GAATTCGAGCTCG
ATG40_VerF-182	This study	cctaagcagcgaatacac
ATG40_VerR+201	This study	ccgatacagcatgtttaacc
PEP4_S1	This study	agcctagtacctagtatttaatccaaataaaattcaaacaaaaacaaaactaac atgttcCGTACGCTGCAGGTCGAC
PEP4_S2	This study	ctagatggcagaaaaggatagggcggagaagtaagaaaagttagctcaaattg ctttggcATCGATGAATTCGAGCTCG
PEP4_1F	This study	gggaagaataacaaaaagtatatctcacc
PEP4_2R	This study	cgggctaccgcatataatgacattatggg
FMT1_S1	This study	ggggacctaggcactcatagtgagaagattgaaaacggcgtatcaacaatg CGTACGCTGCAGGTCGAC
FMT1_S2	This study	gaaagaagggagaagaagaataactaaaaaggaatgtagaccgggttaAT CGATGAATTCGAGCTCG
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FMT1_VerR	This study	gggcttctacacttaattca
MIS1_S1	This study	tctcagctctgttcatttgcagatatttaaggctaaaaggaaatgCGTACGCT GCAGGTCGAC

MIS1_S2	This study	catcgaggtcgaattgatgccattaggagtctgccaactatttaATCGATG AATTCGAGCTCG
MIS1_VerF	This study	ctcctgcaacactcttctat
MIS1_VerR	This study	gctttattgaaggaagacgc
Mitochondrial initiator tRNA oligonucleotide probe for northern blotting	Li, et al. <sup>29</sup>	tagcaataatacgatttg