1	Supplementary Information
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3	Mechanism of Pentagalloyl Glucose in Alleviating Fat Accumulation in
4	Caenorhabditis elegans
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8	1. The properties of the mutants used in the study were provided as followed:
9	(1) ZXW618 <i>hkdIs618</i> [<i>dhs-3p::dhs-3::GFP</i>] strain expressed green fluorescence
10	protein-fused lipid droplet marker protein DHS-3 (DHS-3::GFP);
11	(2) XA7702 [mdt-15 (tm2182) III] is a short-lived and toxin sensitive strain with
12	altered fat storage and low brood size;
13	(3) DG2179 [tub-1 (nr2044) II] is deficient in lipolysis and results in accumulating
14	fatty acid;
15	(4) BX106 [fat-6 (tm331) IV] lacks the gene encodes stearoyl-CoA desaturases in fatty
16	acid biosynthesis;
17	(5) BX153 [fat-7 (wa36) IV] lacks the gene encodes stearoyl-CoA desaturases in fatty
18	acid biosynthesis.
19	(6) RB754 [aak-2 (ok524) X] lacks the AMP-activated protein kinase gene.
20	(7) EU1 [skn-1 (zu67) IV] is sensitive to oxidative
21	stress and have shortened lifespans.
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24 **2. Supplementary Figure Captions**

25 Figure S1: The illustrative diagram indicating the treatment methods of PGG. Figure S2: The efficacy of this strain ZXW618. (A) Representative fluorescence 26 27 photographs of ZXW618 worms induced with 10 mM glucose for 3 days from eggs. (B) Distribution of lipid droplets in high-fat strain ZXW618. (C) Average number of 28 lipid droplets of high-fat strain ZXW618. (D) Average size of lipid droplets of high-fat 29 strain ZXW618. (E-F) Distribution of lipid droplets in normal and high-fat strain 30 ZXW618 stained with Nile red, respectively. (G) Average number of lipid droplets of 31 strain ZXW618 stained with Nile red. (H) Average size of lipid droplets of strains 32 ZXW618 stained with Nile red. 33 34 Figure S3: Effects of PGG on morphology and physiological function in *C. elegans*. (A) Body length of C. elegans cultured with PGG for 4 days from eggs. (B) Body 35 width of C. elegans cultured with PGG for 4 days from eggs. More than 15 worms for 36 each assay were measured. (C-F) Pharyngeal pumping rate, body bending frequency, 37 head thrash frequency and locomotivity of worms cultured with or without PGG at 800 38

- μ M from eggs. More than 10 worms for each assay were measured on days 4, 8, and
- 40 12 respectively. Experiments were repeated 3 times.
- Figure S4: (A-B) GC traces of fatty acids extract of normal wild-type worms. (C-D)
 GC traces of fatty acids extract of high-fat wild-type worms.



Figure S1









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Figure S3





Gene	Primer
mdt-15	AACATCAGCTCAGGCAAGAAA (F)
	TCTGTCCACCTGGACGAATAC (R)
nhr-49	AGGCTCGTGTCAATCAAGAGA (F)
	CTTCCATCGAAAGATCCATGA (R)
sbp-1	CTACTCGCACCATTCTTCTCG (F)
	CCAAATCTCAACTGCTTCTGC (R)
fat-5	TGTCGCTTTATTTGTGGCTCT (F)
	GAAAGCGATCGAGTTGATGAG (R)
fat-6	CTTGTGCTGCTTCATTCTTCC (F)
	GAAGTTGTGACCTCCCTCTCC (R)
fat-7	ACCCGTGGATTCTTCTTCACT (F)
	AACGGGGATAATTGTTGGAAG (R)
fasn-1	ACTGTCGGATCAGCTGAGAAA (F)
	GACGAGCCAAACATCTGAGAG (R)
pod-2	AAGACGATCCTCGAGAAGGAG (F)
	CATCATGCTGAGAGACACGAA (R)
elo-2	ATGCCCTCACATTTGTCTACG (F)
	CCAGGAACAGAATCAGCAGAC (R)
acs-2	GGCTGAACAACAACGCATATT (F)
	CGGAACATGGTAGGACTTTGA (R)
vit-2	CTGAGCAAATCCAAGAAGTCG (F)
	AGTGGGAGAATGTCCTCGTTT (R)
lipl-4	ATGGCCGAGAAGTTCCTACAT (F)
	ATACATGTTTCCGGCTGTACG (R)
aak-2	CATATCATCCGCCTCTACCAA (F)
	GTCCGCAATCTTCACATTGTT (R)
tub-1	AGAGTCCACAATGGAACGATG (F)
	GTTTTCCGTGGAAACTTGTCA (R)
act-1	TCCAAGAGAGGTATCCTTAC (F)
	CGGTTAGCCTTTGGATTGAG (R)

3. Primer sequences for qRT-PCR analysis