SUPPLEMENTAL INFORMATION

Supplemental Table S1. Primers for quantitative real-time PCR.

Supplemental Table S2. Primary antibodies used in this study.

Supplemental Table S3. Effects of dietary quercetin on the body weight gain, liver weight and adipose tissue weight in C57BL/6 mice.

Supplemental Figure S1. Effect of quercetin on ACC phosphorylation and *Cpt1a* expression in BMDMs.

Supplemental Figure S2. Effect of quercetin on GLUT4 translocation in 3T3-L1 adipocytes.

Supplemental Figure S3. Mechanism of quercetin anti-inflammatory effects in adipose tissue macrophages.

SUPPLEMENTAL INFORMATION

 $\label{thm:conditional} \textbf{Table S1. Primers for quantitative real-time PCR} \\$

Gene	Sequence of forward primers (5' to 3')	Sequence of reverse primers (5' to 3')
F4/80	CTTTGGCTATGGGCTTCCAGTC	GCAAGGAGGACAGAGTTTATCGTG
тМср-4	CTTCTGACTTTATCAAGCCGGG	CACTCCAGTTCGCCCCC
Cd11c	CTGGATAGCCTTTCTTCTGCTG	GCACACTGTGTCCGAACTC
Nos2	CCAAGCCCTCACCTACTTCC	CTCTGAGGGCTGACACAAGG
Chi3l3	AGAAGGGAGTTTCAAACCTGGT	GTCTTGCTCATGTGTGTAAGTGA
Mgl2	TTAGCCAATGTGCTTAGCTGG	GGCCTCCAATTCTTCTTGAAACCT
Adiponectin	GCAGAGATGGCACTCCTGGA	CCCTTCAGCTCCTGTCATTCC
GLUT4	GTGACTGGAACACTGGTCCTA	CCAGCCACGTTGCATTGTAG
Mcp-1	CCCCAAGAAGGAATGGGTCC	GGTTGTGGAAAAGGTAGTGG
IL-6	CCTTCCTACCCCAATTTCCAA	AGATGAATTGGATGGTCTTGGTC
Tnf-α	ACGGCATGGATCTCAAAGAC	AGATAGCAAATCGGCTGACG
IL-1β	GCAACTGTTCCTGAACTCAACT	TCTTTTGGGGTCCGTCAACT
IL-10	ACTGCACCCACTTCCCAGT	TGTCCAGCTGGTCCTTTGTT
Cpt1a	CTATGCGCTACTCGCTGAAGG	GGCTTTCGACCCGAGAAGA
Ucp1	CACTCAGGATTGGCCTCTACG	GGGGTTTGATCCCATGCAGA
GAPDH	TGTCATACTTGGCAGGTTTCT	CGTGTTCCTACCCCCAATGT

Table S2. Primary antibodies used in this study

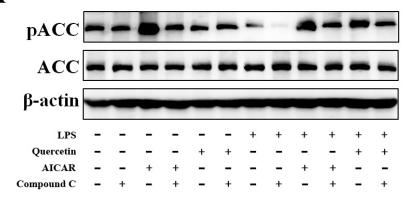
Antibody name	Supplier	Catalog No.	Dilution
AKT rabbit mAb	Cell Signaling Technology	4691	1:1000
Phospho-AKT (Ser473) antibody	Cell Signaling Technology	9271	1:1000
AMPKα antibody	Cell Signaling Technology	2532	1:1000
Phospho-AMPKα (Thr172) antibody	Cell Signaling Technology	2531	1:1000
SIRT1 rabbit mAb	Cell Signaling Technology	9475	1:1000
Acetyl-CoA Carboxylase rabbit mAb	Cell Signaling Technology	3676	1:1000
Phospho-Acetyl-CoA Carboxylase (Ser79)	Cell Signaling Technology	3661	1:1000
antibody			
LKB1 rabbit mAb	Cell Signaling Technology	3047	1:1000
Phospho-LKB1 (Ser428) rabbit mAb	Cell Signaling Technology	3482	1:1000
Anti-GLUT-4 polyclonal antibody	Millipore	07-1404	1:500
Anti-Actin	Sigma-Aldrich	A2066	1:100
Purified anti-mouse Mac-2	Biolegend	125401	1:800

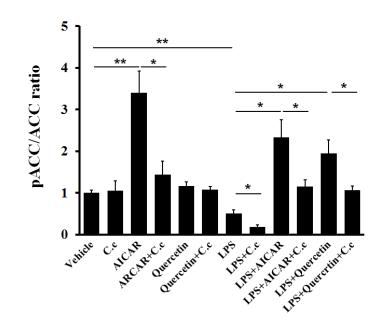
Table S3. Effects of dietary quercetin on the body weight gain, liver weight and adipose tissue weight in C57BL/6 mice.

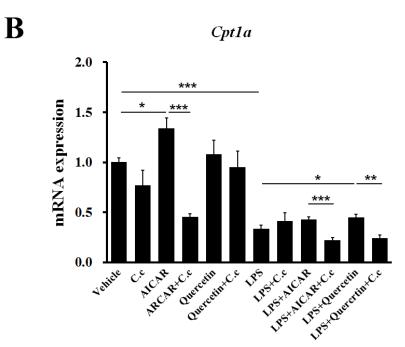
	HFD	HFD+0.1%Qu	LFD
Initial body weight (g)	19.660±0.252 a	19.588±0.472 a	19.300±0.407 a
Final body weight (g)	41.320±0.950 ^a	37.244±1.455 ^b	30.455±0.592°
Liver weight (g)	1.359±0.060 a	1.341±0.079 a	1.494±0.068 ^a
EAT weight (g)	1.861 ±0.061 ^a	1.302±0.175 ^b	0.361±0.063°
SAT weight (g)	1.335±0.109 a	0.783±0.121 ^b	0.263±0.029°
BAT weight (g)	0.168±0.015 ^a	0.18±0.028 ^a	0.082±0.005 ^b

Values are mean \pm SEM, n=8. Values within a row with different superscript letters in each experiment are significantly different from each other (p<0.05) by Mann-Whitney test. LFD, low fat diet; HFD, high fat diet; HFD+0.1% Qu, high fat diet containing 0.1% quercetin; EAT, epididymal adipose tissue; SAT, subcutaneous adipose tissue; BAT, brown adipose tissue.

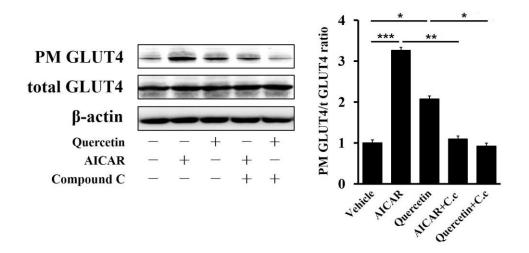




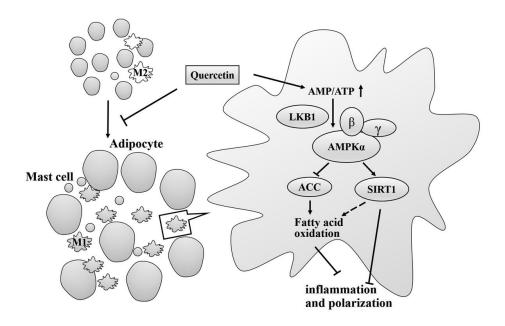




Supplemental Figure S1. Effect of quercetin on ACC phosphorylation and *Cpt1a* expression in BMDMs. (A) Immunoblots for phospho-ACC (Ser 79) and ACC in BMDMs. Quantification of protein expression was described as ratio of phosphorylated ACC to total ACC. (B) Real-time PCR quantitative mRNA expression of *Cpt1a* in BMDMs. *GAPDH* was used as reference gene in qPCR analysis. Data are represented as fold changes compared to vehicle. C.c represented Compound C. Statistical difference between groups was shown using a Student's t test. n=4 per group. *p<0.05, **p<0.01, ***p<0.001. All data are mean ± SEM.



Supplemental Figure S2. Effect of quercetin on GLUT4 translocation in 3T3-L1 adipocytes. Differentiated 3T3-L1 adipocytes were starved for 2 h in serum-free medium before starting the experiment. Thereafter, 2 mM AICAR or 20 μM quercetin were added into medium with or without 10 μM Compound C for 30 min. Quantification of protein expression was described as ratio of plasma membrane protein to total protein. Data are represented as fold changes compared to vehicle. C.c represented Compound C. Statistical difference between groups was shown using a Student's t test. n=4 per group. *p<0.05, **p<0.01, ***p<0.001. All data are mean ± SEM.



Supplemental Figure S3. Mechanism of quercetin anti-inflammatory effects in adipose tissue macrophages.