Supplementary Figure Legends

Supplementary Fig. I. Immunoblotting of various tissue extracts from WT and Tg mice using anti-Flag antibody. 30µg of protein were fractionated in each lane. Flag immunoblotting demonstrated the kidney specificity of transgene expression.

<u>Supplementary Fig. II.</u> RNA was isolated from the kidneys of WT or Tg mice with or without cisplatin, and analyzed for the levels of PPAR α , PPAR γ , PGC1 α target genes. Data are expressed relative to the mRNA of each gene in WT mice without cisplatin and are the mean±S.E. (n=4): *p<0.05 vs. saline-infused WT mice, †p<0.05 vs. saline-infused TG mice.

Supplementary Fig. III. RNA was isolated from the kidneys of NC or CR mice with or without cisplatin, and analyzed for the levels of PPAR α , PPAR γ , PGC1 α genes. Data are expressed relative to the mRNA of each gene in NC mice without cisplatin and are the mean±S.E. (n=4): *p<0.05 vs. saline-infused NC mice, †p<0.05 vs. saline-infused CR mice.

Supplementary Fig. IV. RNA was isolated from the kidneys of NC or CR mice with or without cisplatin, and analyzed for the levels of ACOX1 and MCAD which are also PPAR α target genes and play significant roles in FAO. Data are expressed relative to the mRNA of each gene in NC mice without cisplatin and are the mean±S.E. (n=4): *p<0.05 vs. saline-infused NC mice, p<0.05 vs. saline-infused CR mice, p<0.05 vs. cisplatin-infused NC mice.

Supplementary Fig. V. A, RNA was isolated from the kidneys of WT or Tg mice with or without cisplatin, and analyzed for the levels of CPT-1 α and CPT-1 β genes. B, RNA was isolated from the kidneys of WT or Tg mice with or without I/R, and analyzed for the levels of CPT-1 α and CPT-1 β genes. Data are expressed relative to the mRNA of each gene in WT mice without cisplatin and are the mean±S.E. (n=4): *p<0.05 vs. saline-infused or sham-operated WT mice, **p<0.01 vs. sham-operated WT mice, †p<0.05 vs. saline-infused or sham-operated TG mice, ††p<0.01 vs. sham-operated TG mice, §p<0.05 vs. cisplatin-infused WT mice.



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ACOX1

MCAD

A.

CPT1 a



CPT1 β







Supplementary Figure V, Hasegawa et al.

B.

Supplementary Table I. Primer sequences designed for the probe of Southern blotting

	Product Size	Sequences
Forward	459	TTTAATCAGGTAGTTCCTCGATGTC
Reverse		TGAACTTGAGTCTTCTGAAACATGA

Supplementary Table II. Primer sequences for real-time PCR

		Sequences (5'-3')		
		Forward	Reverse	
28S rRNA	Conrol	AACGGCGGGGGGGAGTAACTATGA	TAGGGACAGTGGGAATCTCG	
FATP-2	PPAR alpha target genes	ATGCCGTGTCCGTCTTTTAC	CTTCAGACCTCCACGACTCC	
CYP4A10		CTCATTCCTGCCCTTCTCAG	GTAGTTCGAAGCGGAGCAGT	
ACOX1		ATGGTTTTCGTAAGGTCCTTC CT	GGCTCGCTTCTCTTGATTTCA	
MCAD		ATGCCCTGGATAGGAAGACA	CATAGCCTCCGAAAATCTGC	
GlyK	PPAR gamma target genes	TGAAGTCAATTGGTTGGGTT ACA	ATGCAGCCAGTGGCTTATGAA	
GPAT		CAACACCATCCCCGACATC	GTGACCTTCGATTATGCGATCA	
Atp5g1	PGC-1 alpha target genes	AGTTGGTGTGGGCTGGATCA	GCTGCTTGAGAGATGGGTTC	
Cox5a		GGGTCACACGAGACAGATGA	GGAACCAGATCATAGCCAACA	

Supplementary Table III. Primer sequences for real-time PCR

		Forward	Reverse	
28S rRNA	Conrol	AACGGCGGGGAGTAACTATGA	TAGGGACAGTGGGAATCTCG	
CPT1a	PPAR alpha target genes	GATGTGGACCTGCATTCCTT	TCCTTGTAATGTGCGAGCTG	
CPT1β		CCCATGTGCTCCTACCAGAT	CCTTGAAGAAGCGACCTTTG	