STAT3 Undergoes Acetylation-dependent Mitochondrial Translocation to Regulate Pyruvate Metabolism

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Figure S1. (a) Peptide slot blot analysis of the antibodies prepared for STAT3 acetylation on K87, K707 and K709. (b) y and b numbers of the mass spectrum in Fig. 2b. (c) IgG controls were detedted for Immunoprecipitation and co-immunoprecipitation experiments involved in this text.



Figure S2. (a) INSR and CBP were overexpressed in 293T cells. HA-CBP was immunoprecipitated followed by Western blotting analysis with anti-INSR and anti-HA. The bottom panel gave the input of INSR and HA-CBP. (b) 293T cells were transfected with EV, STAT3, or STAT3 and CBP. STAT3 acetylation on lysine 87 was detected with a specific antibody via Western blotting. (c) 293T cells were transfected with EV, wild-type STAT3 and STAT3 with the N-terminus, C-terminus, or both N- and C-termini deleted (i.e., 1-738, 28-770, and 28-738). Mitochondria were isolated from the 293T transfectants and STAT3 and Porin levels were shown by Western blotting. (d) STAT3-K87R mutant or STAT3-WT was transiently transfected into 293T cells. Cytoplasmic and mitochondrial fractions were prepared for STAT3 localization analysis via Western blotting with anti-Myc. (e) Mitochondrial lysates were prepared for STAT3 acetylation analysis from 293T cells transfected with EV, STAT3 or Cox4-STAT3.



Figure S3. Full scans of western blotting data in all figures.





















Fig. 2e

β-actin





STAT3

170 -130 -

95

72 • 55 • 43 •



Fig. 2f



Fig. 2g











Fig. 2j















34 🗕

130 95 72

55



STAT3











Myc-PDC E1

Myc-PDC E1

Fig. 3a

Fig. 4b



Fig. 4c





Flag-STAT3



CBP-HA























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Fig. 4g

aK709











STAT3



And A Line

SIRT3/SIRT5



Fig. 7e

STAT3



170 -130 -95 -

72 -

55 -

43 🗕

170 130 95

72

















Supplementary Fig. 1c



Supplementary Fig. 2a

