

Supplementary Materials

Supplemental Tables

Table S1. Composition of the standard food and the obesity induction food

Calories(kcal%)	CHOW	HFD
Protein(%)	24.52	20
Fat(%)	12.41	60
Carbohydrate(%)	63.07	20

CHOW: Purina Lab. Rodent Chow (38057)

HFD: Research Diets. DIO Series Diets (D12492)

Table S2. CHOW diet (CHOW; Purina Laboratory Rodent Diet 38057)

Components*	Calories(%): Fat 12.41%, Protein 24.52%, Carbohydrate 63.07%	
Nutrients	Protein 20%	Arginine 1.26%, Cystine 0.37%, Glycine 0.87%, Histidine 0.5%, Isoleucine 0.82%, Leucine 1.47%, Lysine 1.01%, Methionine 0.33%, Phenylalanine 0.98%, Tyrosine 0.63%, Threonine 0.72%, Tryptophan 0.25%, Valine 0.91%
	Fat 4.5%	Linoleic Acid 1.10%, Linolenic Acid 0.12%, Arachidonic Acid 0.02%, Omega-3 Fatty Acids 1.11%
	Fiber 3.7%	
Minerals	Ash 7.25%, Calcium 1.2%, Phosphorus 0.62%, Phosphorus (non-phytate) 0.4%, Potassium 0.82%, Magnesium 0.16%, Sulfur 0.22%, Sodium 0.34%, Chlorine 0.47%, Fluorine 21.38 ppm, Iron 112.93 ppm, Zinc 128.85 ppm, Manganese 95.49 ppm, Copper 22.74 ppm, Cobalt 0.76 ppm, Iodine 1.42 ppm, Chromium 0 ppm, Selenium 0.32 ppm	
Vitamins	Vitamin K 6.69 ppm, Thiamin Hydrochloride 11.02 ppm, Riboflavin 11.57 ppm, Niacin 217.7 ppm, Pantothenic Acid 88.72 ppm, Choline Chloride 3447.96 ppm, Folic Acid 13.6 ppm, Pyridoxine 11ppm, Biotin 0.15 ppm, B12 41 ppm, Vitamin A 28.03 IU/kg, Vitamin D3 (added) 4 IU/kg, Vitamin E 100 IU/kg	

Table S3. High fat diet (HFD; Research Diets D12492) (Research Diets Inc., USA)

Components*	Calories(%): Fat 60%, Protein 20%, Carbohydrate 20%				kcal/gm
	Fat	Protein	Carbohydrate	Total	
gm%	34.9	26.2	26.3		5.24
kcal%	60	20	20	100	
Ingredient		gm		kcal	
Casein, 80 Mesh		200		800	
L-Cystine		3		12	
Maltodextrin 10		125		500	
Sucrose		68.8		275.2	
Cellulose, BW200		50		0	
Soybean Oil		25		225	
Lard		245		2205	
Mineral Mix S10026		10		0	
DiCalcium Phosphate		13		0	
Calcium Carbonate		5.5		0	
Potassium Citrate, 1 H ₂ O		16.5		0	
Vitamin Mix V10001		10		40	
Choline Bitartrate		2		0	
FD&C Blue Dye #1		0.05		0	
Total		773.85		4057	

Supplemental Figures

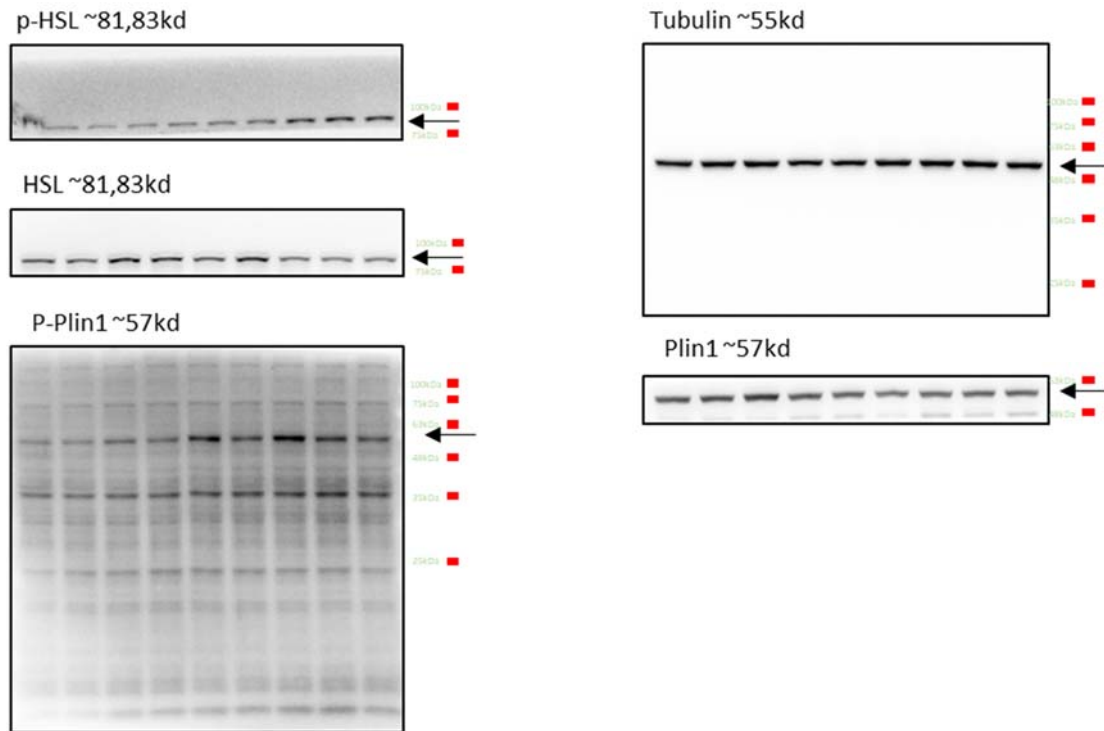


Figure S1. Membrane images of western blot analysis used for Figure 1.

Figure 2.A

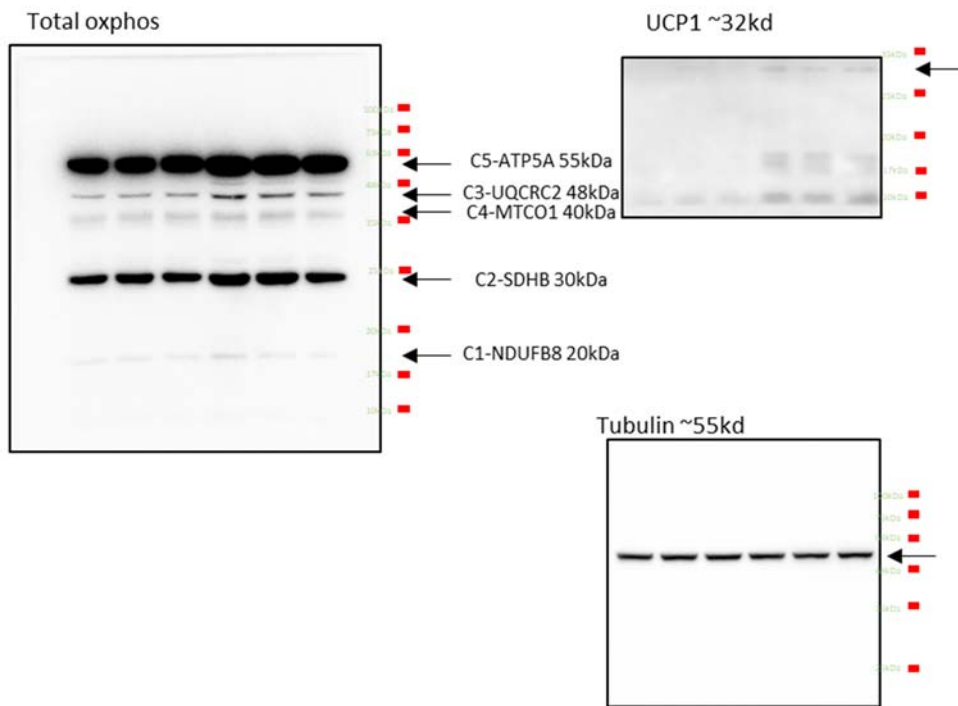


Figure 2.E

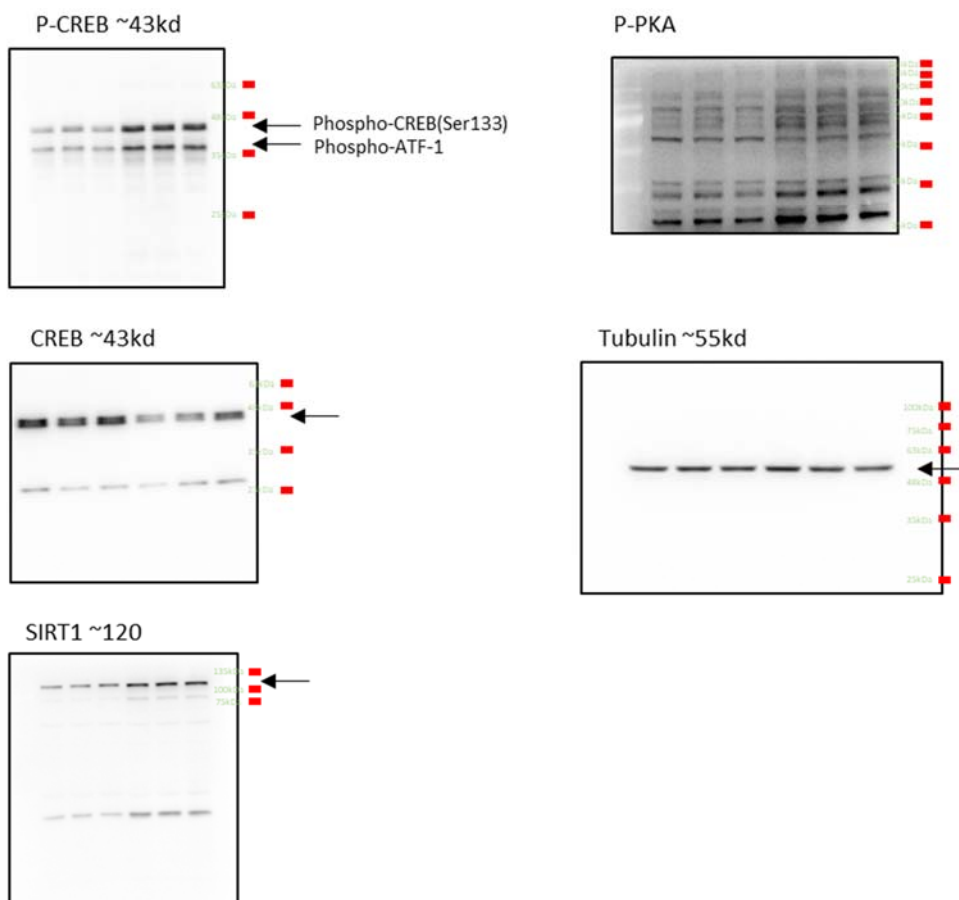


Figure S2. Membrane images of western blot analysis used for Figure 2.

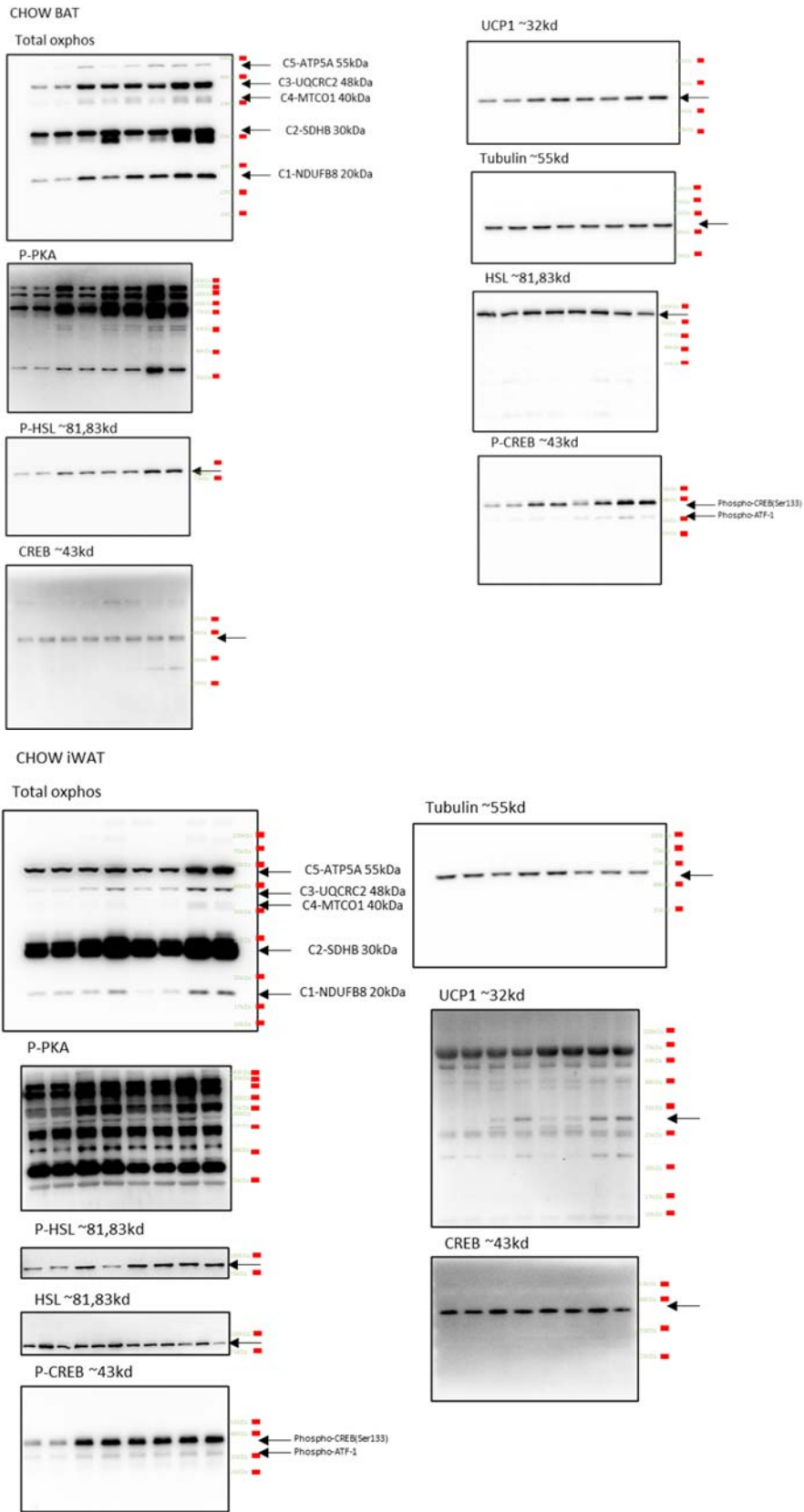


Figure S3. Membrane images of western blot analysis used for Figure 3.

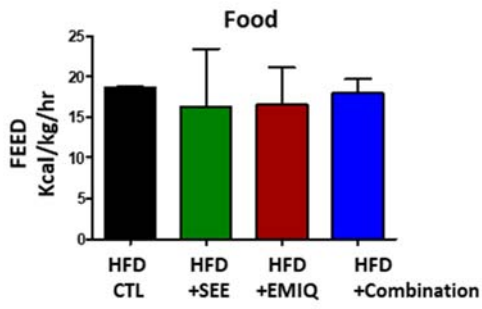


Figure S4. Effects of SEE and EMIQ on food consumption

Mice fed with 8 weeks of HFD were treated with SEE (400mg/kg/day), EMIQ (100mg/kg/day), or SEE/EMIQ for 2 weeks. Food consumption was monitored for 48 hours after the last dose by using TSE PhenoMaster system.

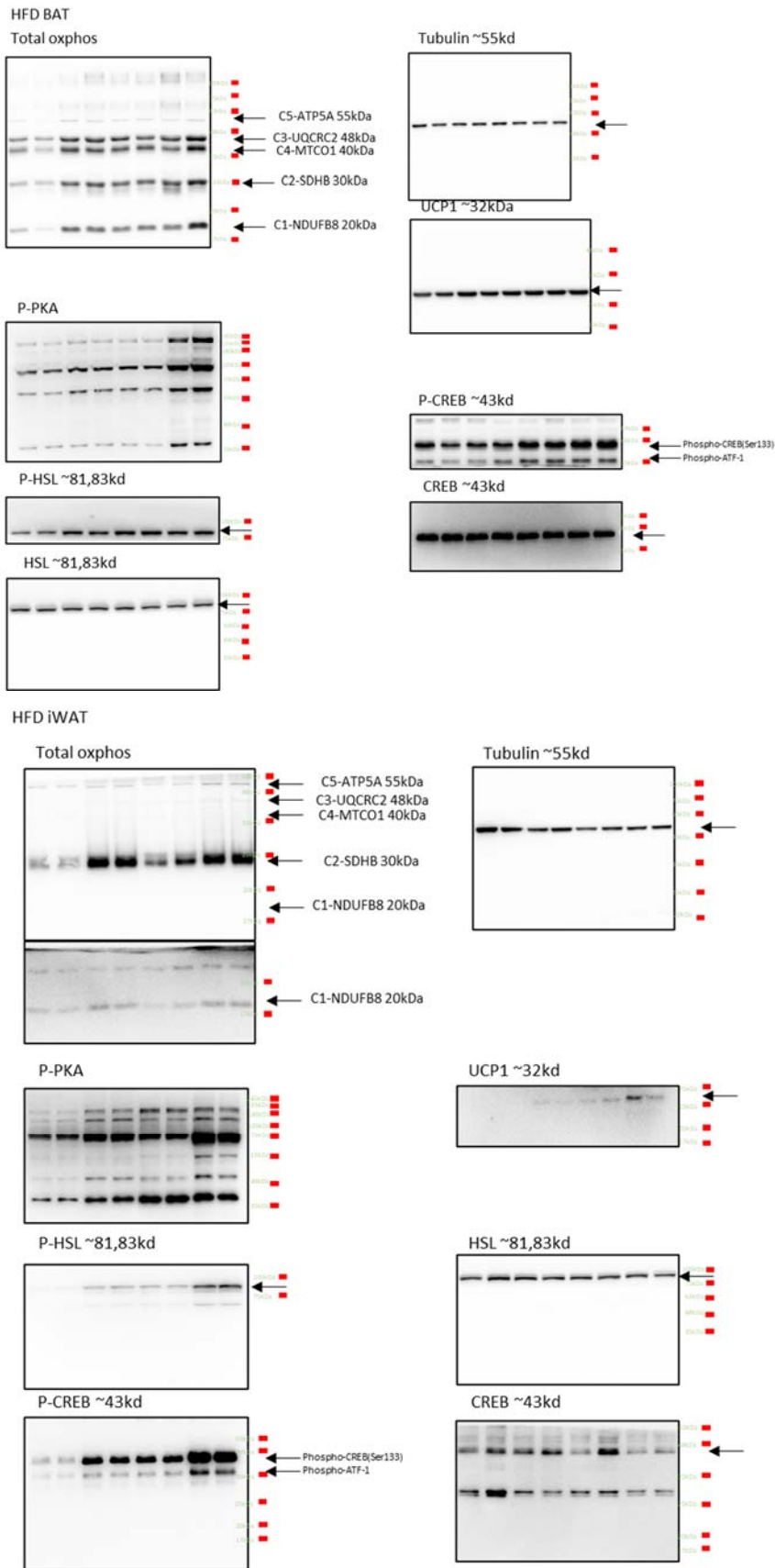


Figure S5. Membrane images of western blot analysis used for Figure 5.

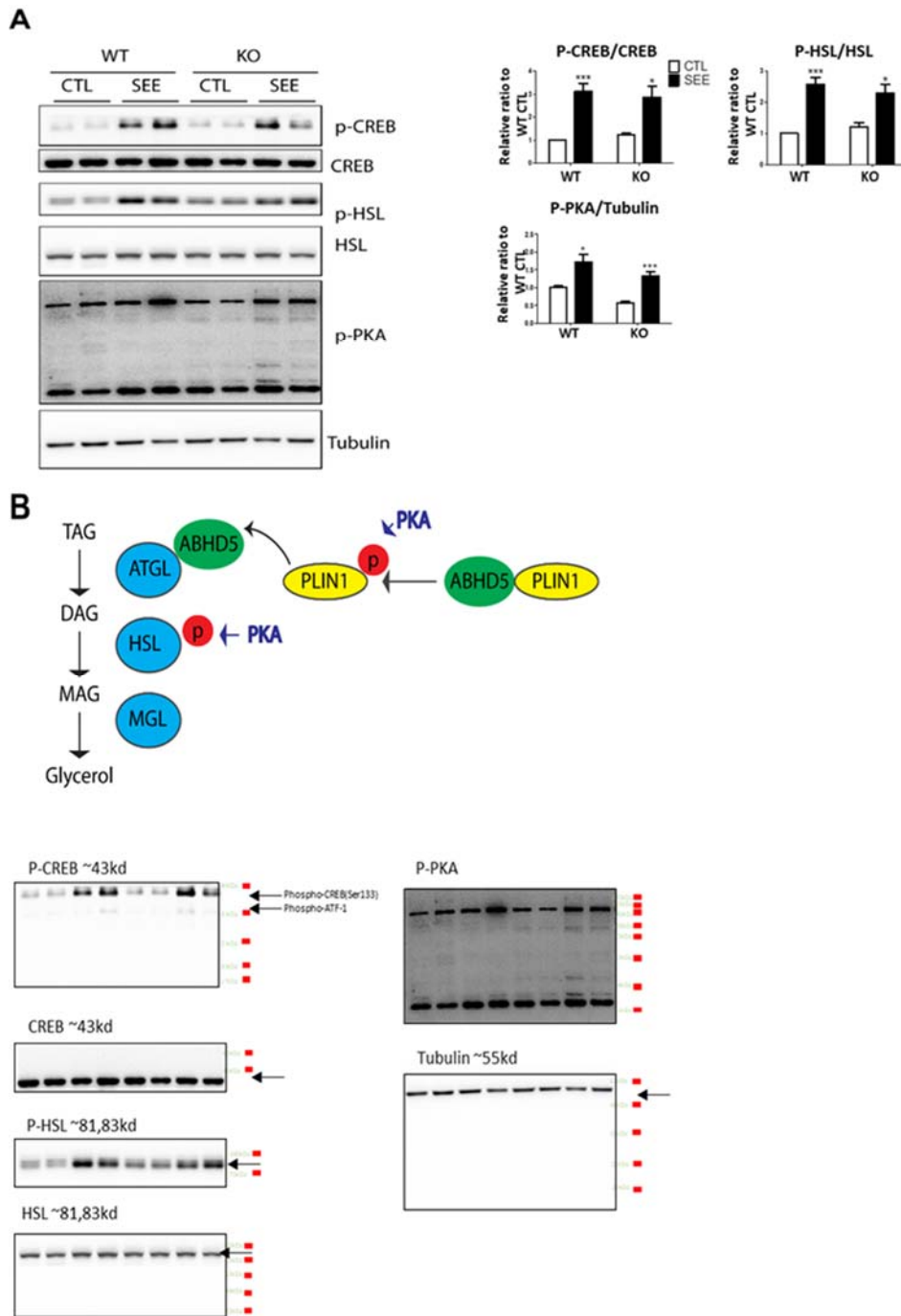


Figure S6. Effects of SEE on phosphorylation levels of CREB and HSL in BAT of adipocyte-specific ATGL knockout mice.

(A) Immunoblot analysis of proteins involved in PKA signaling (CREB, HSL and PKA substrates) in BAT of WT controls and adipocyte-specific ATGL KO mice, fed HFD for 8 weeks and treated with vehicle (CTL) and SEE (400mg/kg) for 2 weeks. (n = 6, means \pm SEM, * p < 0.05, ** p < 0.01, *** p < 0.001) with membrane images of western blot analysis used for Figure S6. (n=6, means \pm SEM, * p < 0.05, ** p < 0.01, *** p < 0.001).

(B) Diagram of PKA signaling.

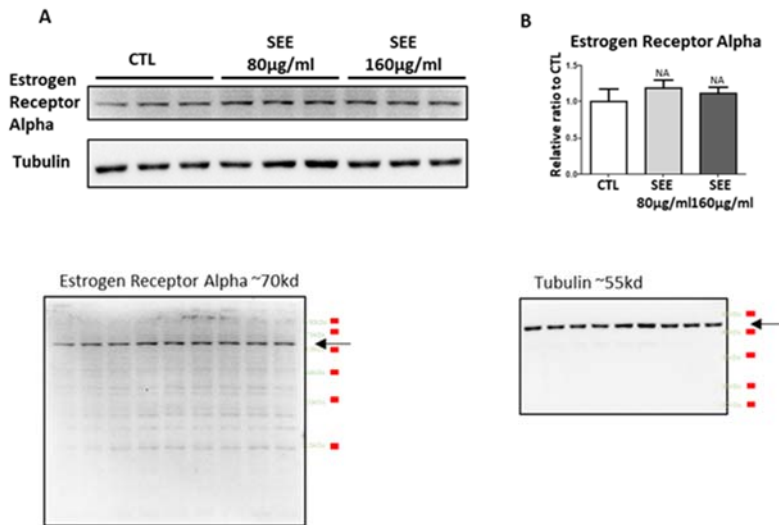


Figure S7. (A,B) Immunoblot analysis of Estrogen Receptor Alpha in adipocytes differentiated from C3H10T1/2 cells treated with Control vehicle (CTL), SEE (80µg/ml) and SEE (160µg/ml) for 24 h. (n=3, means ± SEM, * p < 0.05, ** p < 0.01, * p < 0.001) and Membrane images of western blot analysis used for Figure S7.**