

**Figure S1.** The effects of chronic i.c.v. infusion of RFRP-3 on the mRNA expression of feeding and energy metabolism-related genes. mRNA expression levels for (**A**) hypothalamus and (**B**) pituitary. We chose neuropeptide Y (NPY), agouti-related peptide (AgRP), and proopiomelanocortin (POMC) as neuropeptides related to feeding; thyrotropin-releasing hormone (TRH), growth hormone (GH), and thyroid-stimulating hormone  $\beta$  (TSH $\beta$ ) as neuropeptides related to energy metabolism; gonadotropin-releasing hormone (GnRH), prolactin (PRL), luteinizing hormone  $\beta$  (LH $\beta$ ), and follicle-stimulating hormone  $\beta$  (FSH $\beta$ ) as neuropeptides related to reproduction; RFRP and G protein-coupled receptor 147 (GPR147), the receptor for RFRP-3. Each value represents the mean ± standard error of the mean. *n* = 7–8.

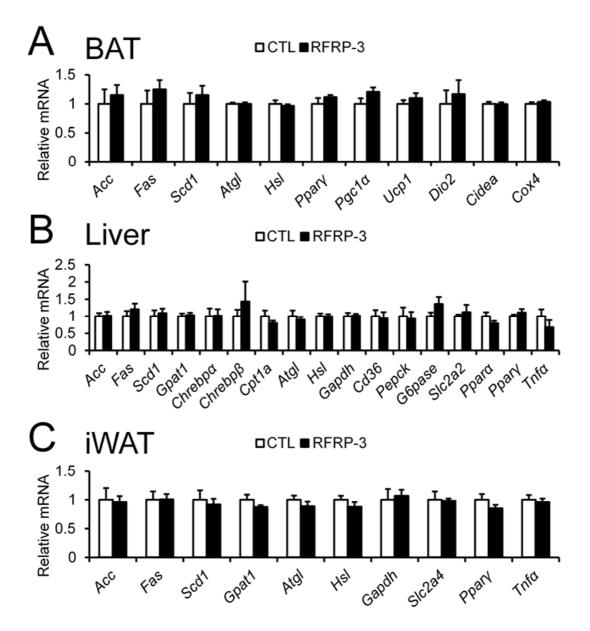


Figure S2. The effects of chronic i.c.v. infusion of RFRP-3 on the mRNA expression of lipid metabolism-related genes in the adipose tissues and liver, and thermogenesis-related genes in the BAT. mRNA expression levels for (A) BAT, (B) liver, and (C) inguinal WAT (iWAT). We chose acetyl-CoA carboxylase (ACC), fatty acid synthase (FAS), stearoyl-CoA desaturase 1 (SCD1), and glycerol-3-phosphate acyltransferase 1 (GPAT1) as lipogenic enzymes; carbohydrate-responsive elementbinding protein  $\alpha$ ,  $\beta$  (ChREBP $\alpha$ ,  $\beta$ ) as lipogenic transcription factors; carnitine palmitoyltransferase 1a (CPT1a), adipose triglyceride lipase (ATGL), and hormone-sensitive lipase (HSL) as lipolytic enzymes; peroxisome proliferator-activated receptor  $\alpha$ ,  $\gamma$  (PPAR $\alpha$ ,  $\gamma$ ) as lipid-activated transcription factors; glyceraldehyde-3-phosphate dehydrogenase (GAPDH) as a carbohydrate metabolism gene; cluster of differentiation 36 (CD36) as a fatty acid transporter; phosphoenolpyruvate carboxykinase (PEPCK) and glucose-6-phosphatase (G6Pase) as gluconeogenesis enzymes; solute carrier family 2 member 2 (SLC2A2) and solute carrier family 2 member 4 (SLC2A4) as glucose transporters; tumor necrosis factor  $\alpha$  (TNF $\alpha$ ) as an inflammatory cytokine; peroxisome proliferator-activated receptor  $\gamma$ coactivator  $1\alpha$  (PGC1 $\alpha$ ), uncoupling protein 1 (UCP1), type II iodothyronine deiodinase (DIO2), cell death-inducing DNA fragmentation factor-like effector A (CIDEA), and cytochrome c oxidase subunit 4 (COX4) as genes related to thermogenic function. Each value represents the mean ± standard error of the mean. n = 7-8.