

Supplemental data

Fruit of *Gardenia jasminoides* induces mitochondrial activation and non-shivering thermogenesis through regulation of PPAR γ

Park et al.

Figure S1

Figure S2

Figure S3

Figure S4

Figure S5

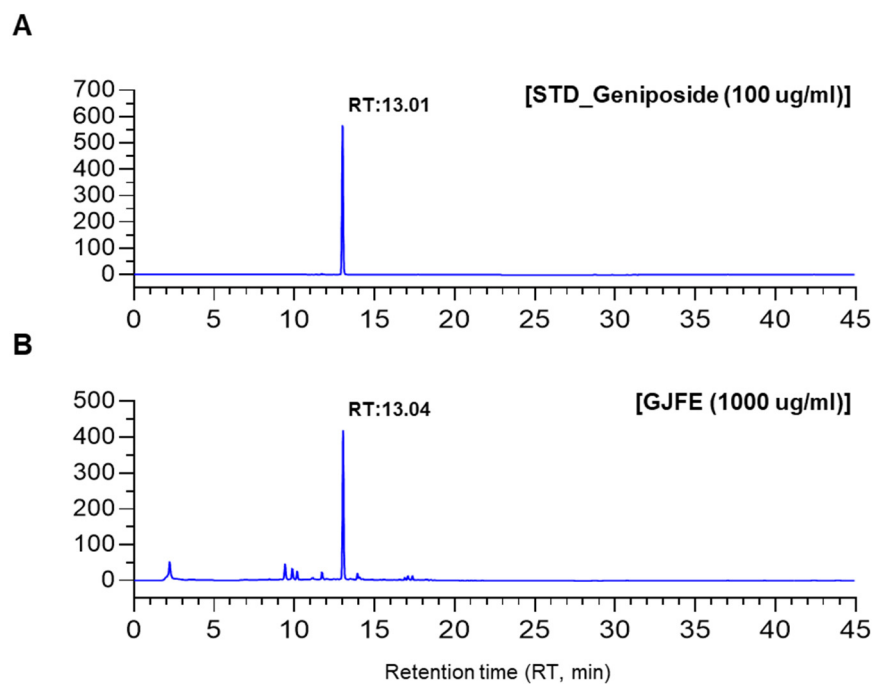


Figure S1. HPLC analysis of geniposide in GJFE. Chromatograms of (A) geniposide and (B) GJFE were exhibited by HPLC analysis. GJFE, *Gardenia jasminoides* fruit extract.

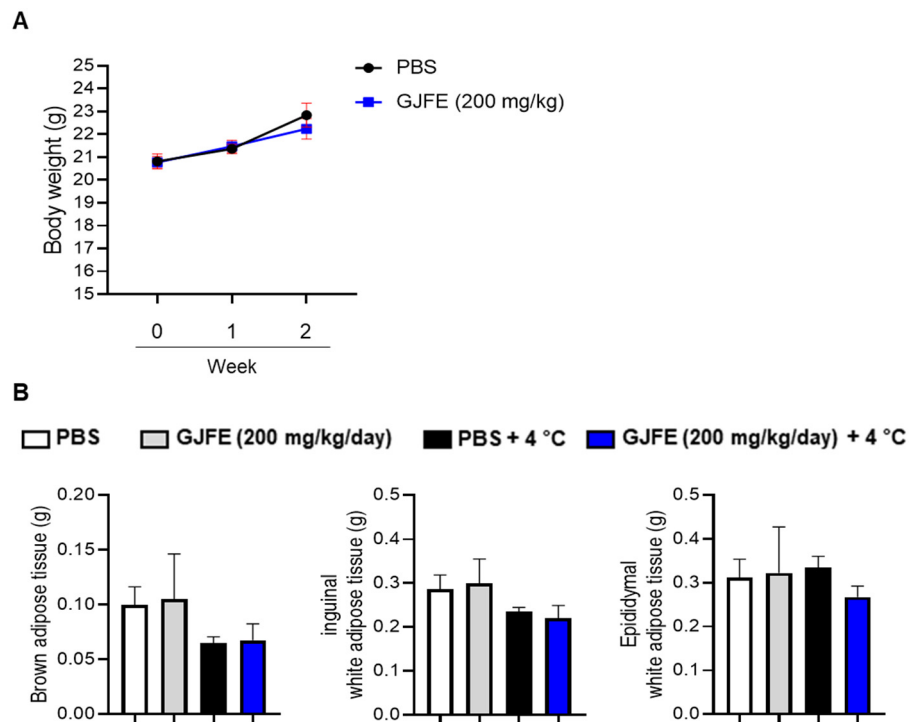


Figure S2. Effect of GJFE on body and tissue weight in cold-exposed mice. (A) Body weight of PBS-fed or GJFE-fed mice ($n = 10$) was measured at indicated time points. (B) Weight of BAT, iWAT, and eWAT was measured ($n = 5$). GJFE, *Gardenia jasminoides* fruit extract; BAT, brown adipose tissue; iWAT, inguinal white adipose tissue; eWAT, epididymal white adipose tissue.

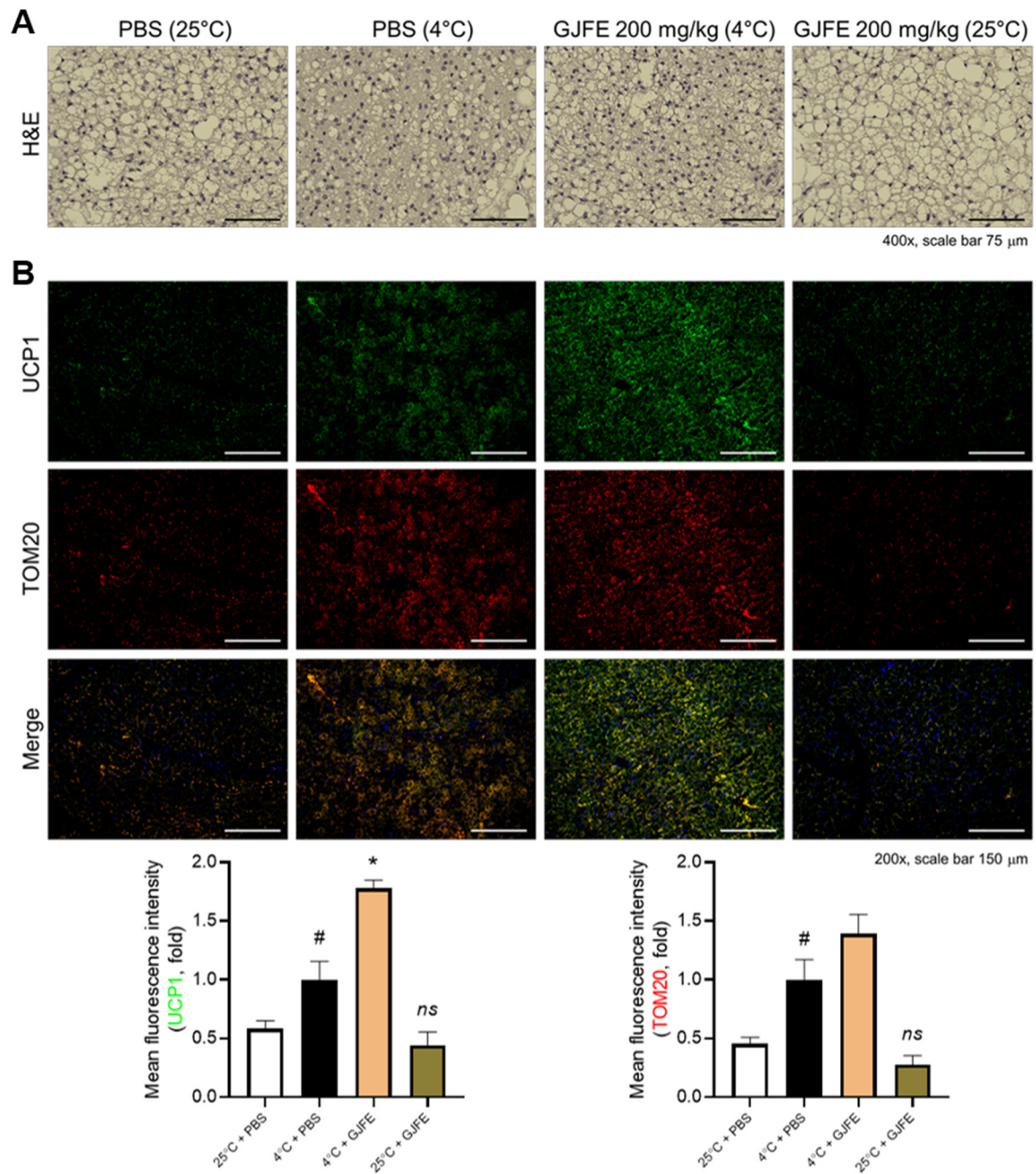


Figure S3. Effect of GJFE on BAT in cold-exposed mice. (A) Paraffin-embedded iBAT were stained with H&E (magnification 400 \times , scale bar 75 μ m). (B) UCP1 (green) and TOM20 (red), and nuclei (blue) were detected in the BAT of the mice by immunofluorescence staining (magnification 200 \times , scale bar 150 μ m). Intensity of the proteins was quantified using the ImageJ software. All data are expressed as the mean \pm S.E.M. of the data from three or more separate experiments. # p < 0.05 vs. BAT of PBS-fed mice, * p < 0.05 vs. BAT of PBS-fed and cold-exposed mice. ns, no significant difference. GJFE, *Gardenia jasminoides* fruit extract.

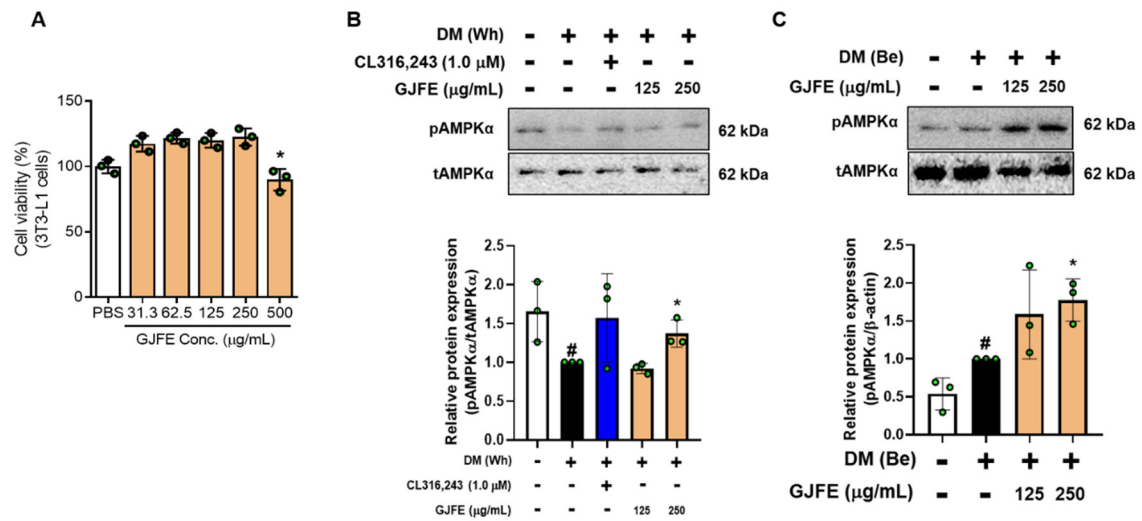


Figure S4. Effect of GJFE on cytotoxicity and protein levels of AMPK in 3T3-L1 adipocytes. (A) Cytotoxicity of GJFE was measured by WST-1 analysis. Phosphorylation of AMPK in (B) white-induced 3T3-L1 cells and (C) beige-induced 3T3-L1 cells was analyzed by Western blot analysis, normalized by levels of total AMPK, and quantified using the ImageJ software. All data are expressed as the mean \pm S.E.M. of the data from three or more separate experiments. $^{\#}p < 0.05$ vs. DM-untreated 3T3-L1 cells, $^*p < 0.05$ vs. PBS-treated 3T3-L1 cell (A) or DM-treated 3T3-L1 cells (B and C). GJFE, *Gardenia jasminoides* fruit extract.

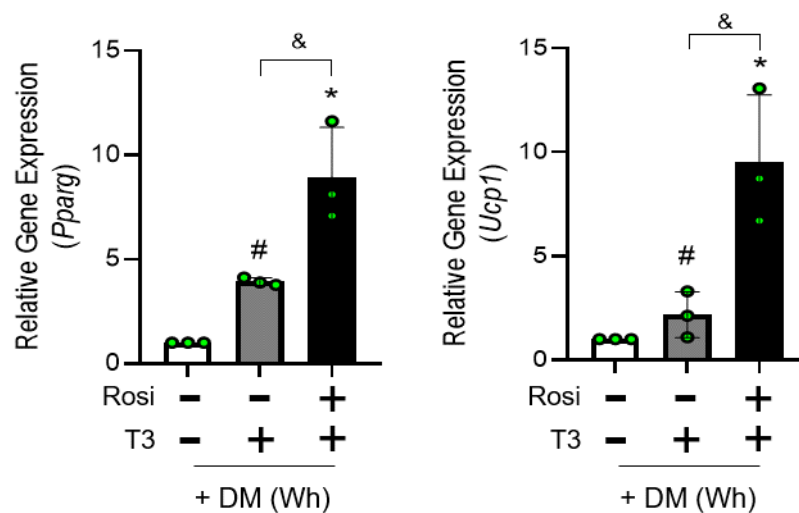


Figure S5. Effect of PPAR γ agonist on mRNA expression of UCP1 in beige adipocytes. mRNA expression of *Pparg* and *Ucp1* was measured by RT-PCR analysis and normalized by *Gapdh*. All data are expressed as the mean \pm S.E.M. of the data from three or more separate experiments. # $p < 0.05$ vs. DM (Wh)-treated 3T3-L1 cells, * $p < 0.05$ vs. DM (Wh) and T3-treated 3T3-L1 cells. GJFE, *Gardenia jasminoides* fruit extract. Rosi; rosiglitazone.