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Supplementary figure



Fig. S1 γ-GC inhibits D-gal-induced senescence in PC12 cells and aging in BALB/c mice. (A)

Images of the cell cycle in each group and percentages of cells in G0/G1, S, and G2/M phases. (B, C) Protein levels of p-p53 (S15), p53 and β -actin in PC12 cells and BALB/c mice by western blotting. Data are presented as mean ± SD of three independent experiments in each group. **p < 0.01, ***p < 0.001 as compared with the control group; ###p < 0.001 as compared with the D-gal group; ns: no significant difference.



Fig. S2 γ -GC prevents D-gal-induced PC12 cell senescence via the activation of AMPK. PC12 cells were treated with D-gal, CC (2 μ M), or D-gal + CC (2 μ M) and then incubated in a complete medium with γ -GC (80 μ M) or Met (80 μ M) for 48 h. Immunoblotting analysis of p53, and p-p53 (S15) levels in each group. Data are presented as mean \pm SD of three independent experiments in each group. **p < 0.01, ***p < 0.001 as compared with the control group; ###p < 0.001 as compared with the D-gal group; &&p < 0.01, &&p < 0.001 as compared with the D-gal and CC combined treatment group; ns: no significant difference.



Fig. S3 γ -GC prevents the increase of Ace-p53 in the nucleus via activating AMPK. (A) D-gal treated PC12 cells were incubated in a complete medium with γ -GC (20, 40, 80 μ M) for 48 h. Immunoblotting analysis the protein level of Ace-p53 and SIRT1 in whole-cell protein in each group. Data are presented as mean \pm SD of three independent experiments in each group. (B) Cells were treated with D-gal or γ -GC (80 μ M) in the presence or absence of CC (2 μ M); immunoblotting analysis of the protein levels of Ace-p53 in the whole-cell protein in each group. Data are presented as mean \pm SD of three independent experiments in each group. Data are presented as mean \pm SD of three independent experiments in each group. Data are presented as mean \pm SD of three independent experiments in each group. Data are presented as mean \pm SD of three independent experiments in each group. (C) Cells were treated with D-gal or γ -GC (80 μ M) in the presence of EX527 (1 μ M); immunoblotting analysis of the protein levels of p53, p-p53 in each group. Data are presented as mean \pm SD of three independent experiments in each group; ###p < 0.001 as compared with the Control group; ###p < 0.01 as compared with the EX527 group; ^{§§}p < 0.01 as compared with the D-gal and EX527 combined treatment group; ns: no significant difference.