

Supplemental Figures

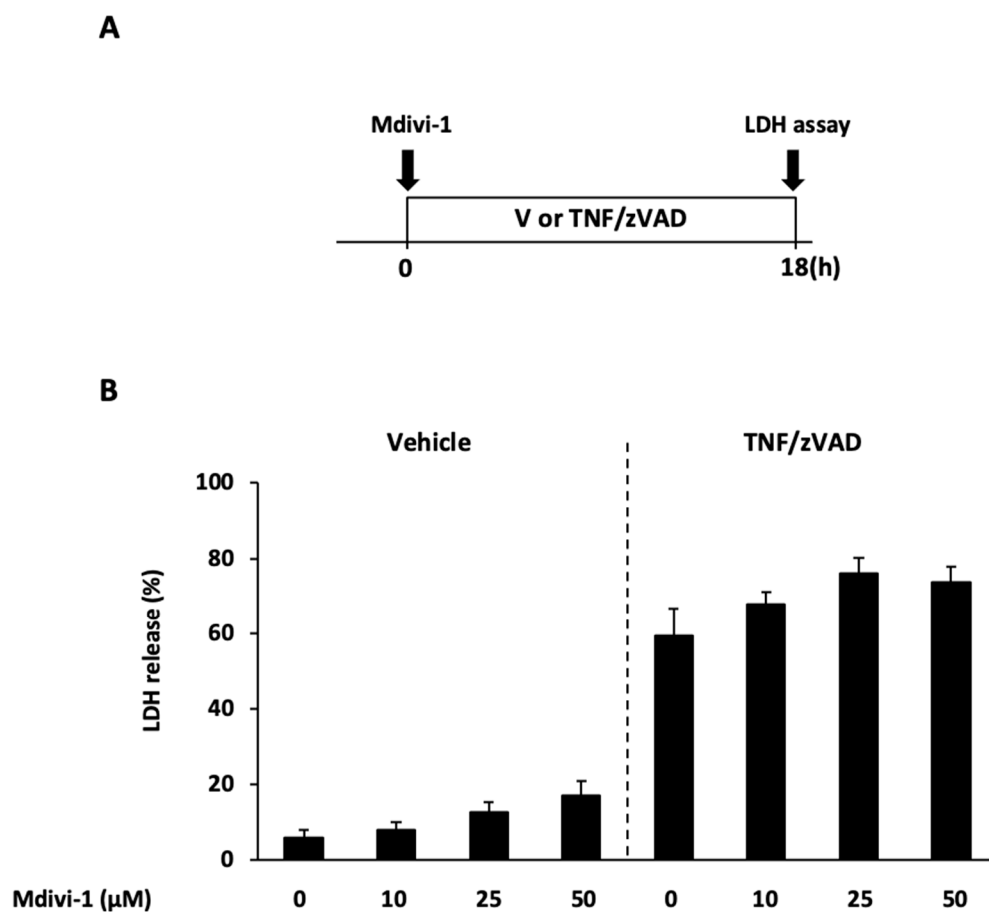


Figure S1. (A, B) Effects of Mdivi-1, a Drp1 inhibitor, on TNF/zVAD-induced LDH release. Experimental protocol (A) and results of quantitative analyses (B) are shown. H9c2 cells were treated with vehicle or various concentration of Mdivi-1 at the same time as the addition of vehicle or TNF- α and zVAD (TNF/zVAD; TNF- α , 50 ng/ml; zVAD, 20 μ M). .

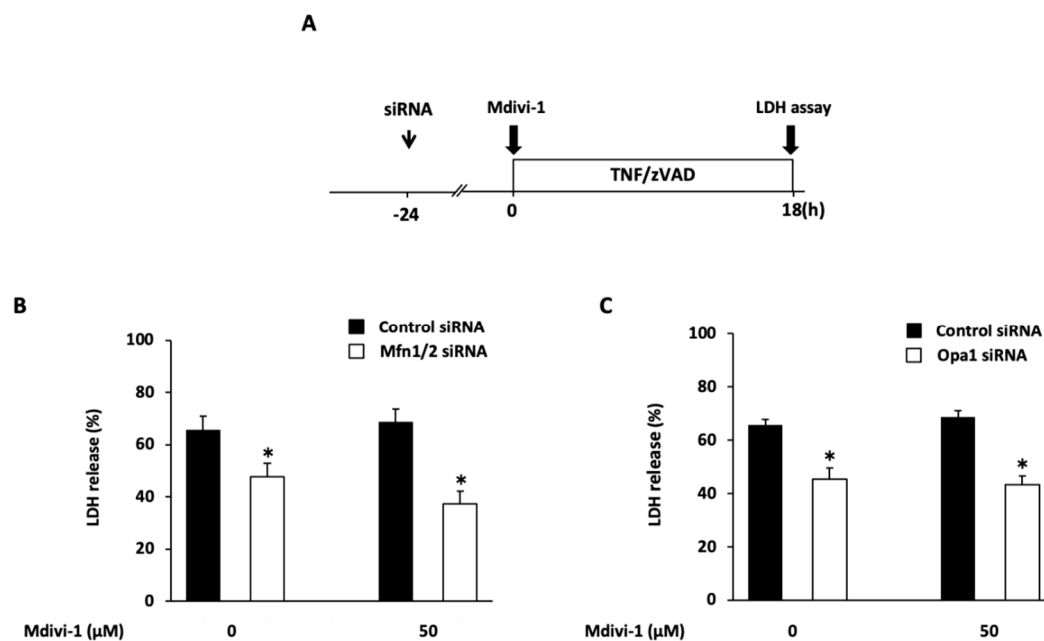


Figure S2. (A–C) Effects of Mdivi-1, a Drp1 inhibitor, on protection from TNF/zVAD-induced cell death by knockdown of Mfn1/2 and Opa1. Experimental protocol (A) and results of quantitative analyses (B and C) are shown. The siRNAs were transfected into H9c2 cells 24 h before the addition of TNF- α and zVAD (TNF/zVAD; TNF- α , 50 ng/ml; zVAD, 20 μ M). Mfn1/2 = mitofusin 1/2, Opa1 = optic atrophy-1. N = 4 in each group. *p<0.05 vs. cells transfected with control siRNA. .

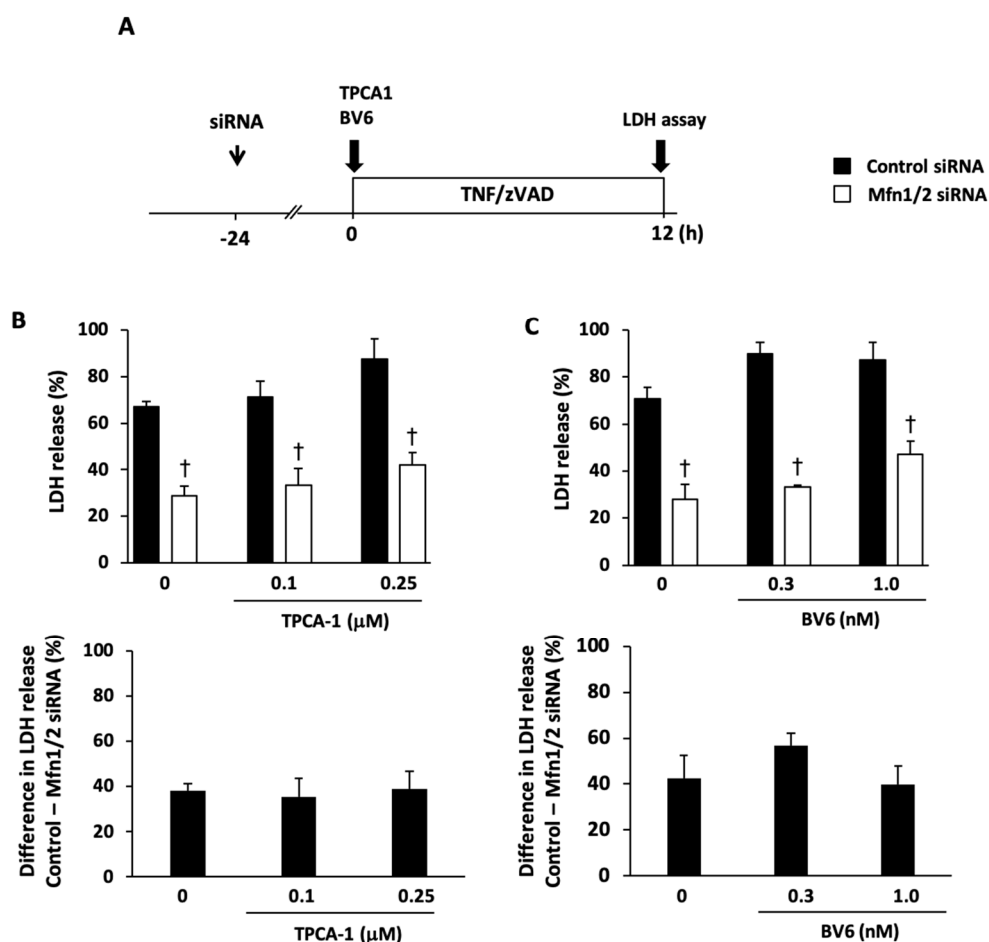


Figure S3. (A, B) Effects of TPCA-1, an IKK α/β inhibitor, and BV6, an inhibitor of cellular inhibitor of apoptosis proteins 1, on protection from TNF/zVAD-induced cell death by knockdown of mitochondrial fusion-regulating proteins. Experimental protocol (A) and results of quantitative analyses (B and C) are shown. The siRNAs were transfected into H9c2 cells 24 h before the addition of TNF- α and zVAD (TNF/zVAD; TNF- α , 50 ng/ml; zVAD, 20 μ M). Mfn1/2 = mitofusin 1/2, Opa1 = optic atrophy-1. N = 6 in each group. †p<0.05 vs cells transfected with control siRNA and treated with similar concentration of inhibitors.

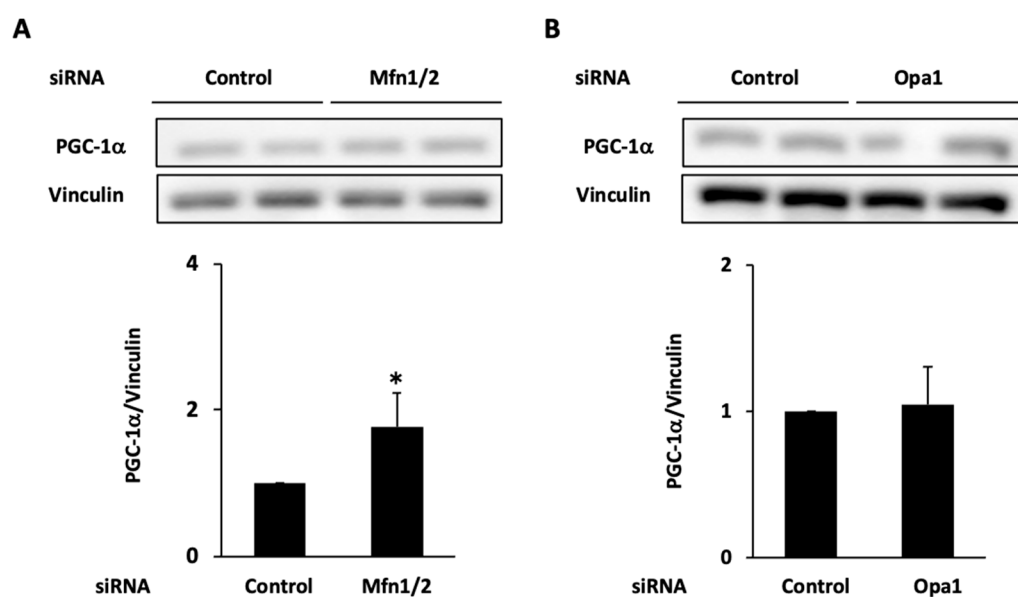


Figure S4. (A, B) Effects of knockdown of mitofusin (Mfn) 1/2 (A) and optic atrophy-1 (Opa1, B) on levels of peroxisome proliferator-activated receptor gamma coactivator-1 α (PGC-1 α). Representative Western blots and results of densitometric analyses are shown. The siRNAs were transfected into H9c2 cells and cell lysates were collected 24 h after the transfection of siRNAs. N = 4 in each group. * $p < 0.05$ vs. cells transfected with control siRNA.