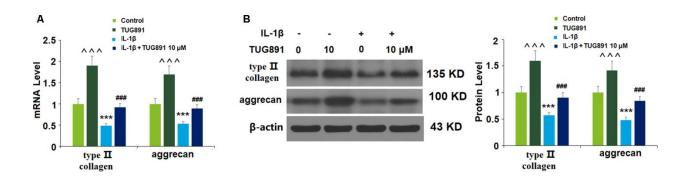
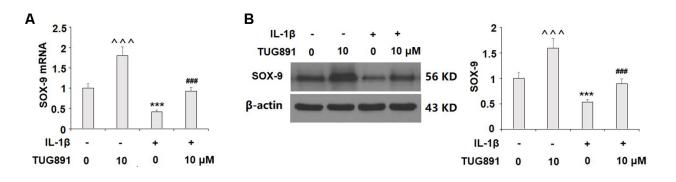
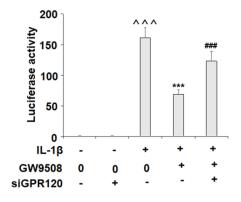
SUPPLEMENTARY FIGURES



Supplementary Figure 1. Treatment with TUG891 prevented IL-1 β -induced reduction of type II collagen and aggrecan in ATDC5 chondrocytes. Cells were treated with IL-1 β (10 ng/ml) with or without TUG891 (10 μ M) for 24 h. (A) mRNA of type II collagen and aggrecan; (B) Protein of type II collagen and aggrecan (^^, ***, P<0.0001 vs. control group; ###, P<0.0001 vs. IL-1 β treatment group, n=4-5).



Supplementary Figure 2. Treatment with TUG891 prevented IL-1 β -induced reduction of SOX-9 in ATDC5 chondrocytes. Cells were treated with IL-1 β (10 ng/ml) with or without TUG891 (10 μ M) for 24 h. (A) mRNA of SOX-9; (B) Protein of SOX-9 (^^, ***, P<0.0001 vs. control group; ####, P<0.0001 vs. IL-1 β treatment group, n=4-5).



Supplementary Figure 3. GW9508 suppressed the activation of NF-\kappaB. ATDC5 chondrocytes were transfected with GPR120 siRNA for 24 h, followed by stimulation with IL-1 β (10 ng/ml) with or without GW9508 (50 μ M) for 24 h. (**A**) Levels of nuclear NF- κ B p65 were measured; (**B**) Luciferase activity of NF- κ B promoter was measured (^^^, P<0.0001 vs. control group; ***, P<0.0001 vs. IL-1 β treatment group, ###, P<0.0001 vs. IL-1 β + GW9508 treatment, n=4-5).