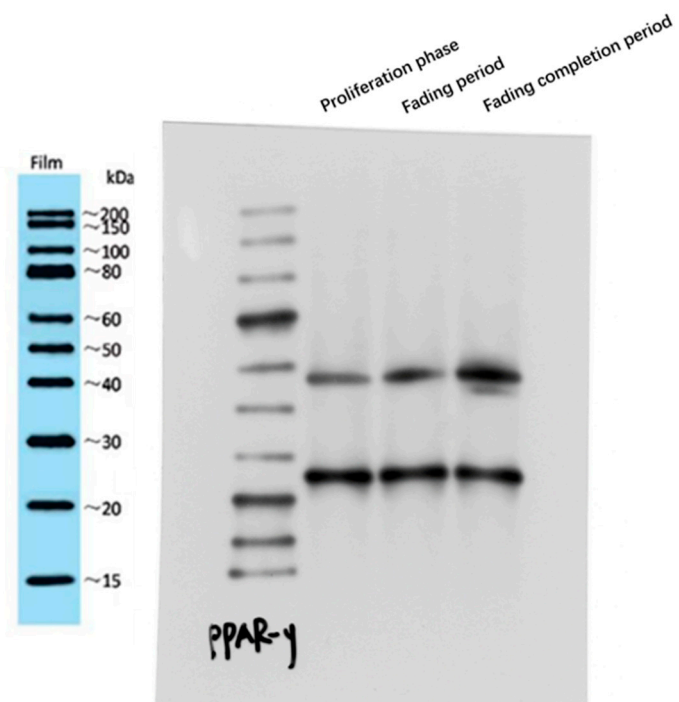
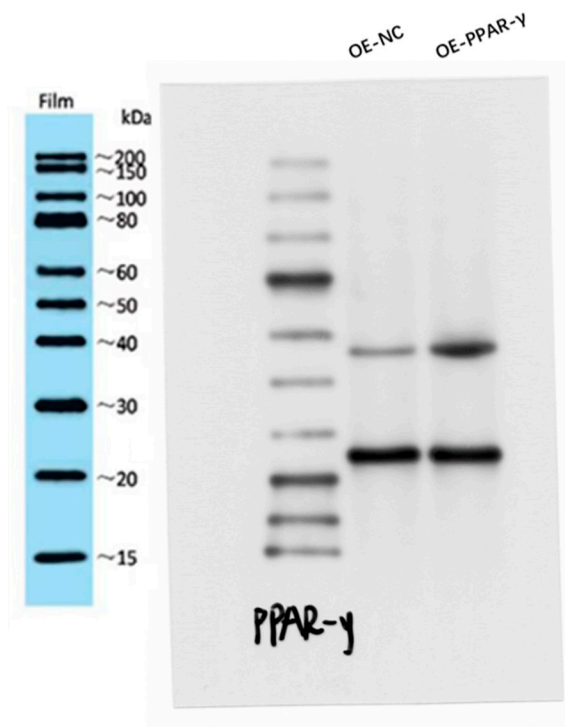


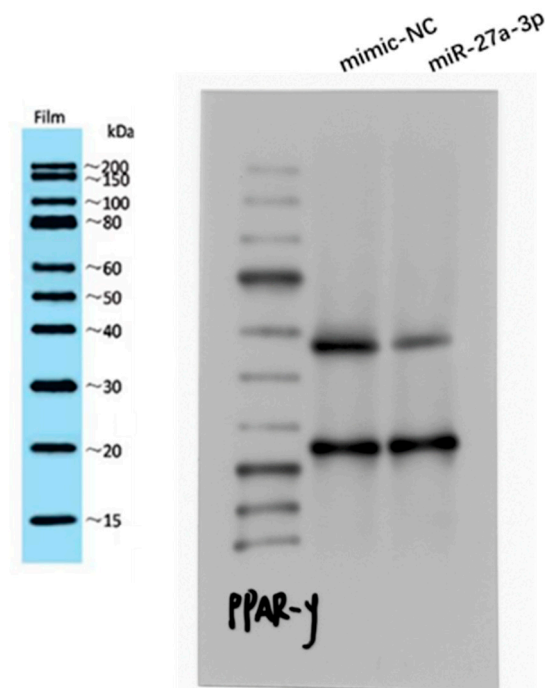
Supplementary Materials



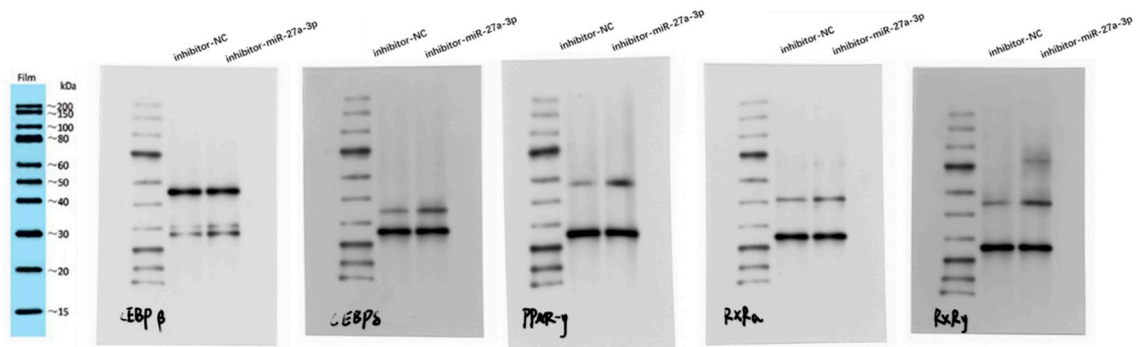
Supplementary Figure S1: The original protein blots of Figure 2B. The protein levels of PPAR- γ in the proliferation, regression, and completion stages of IH were determined using Western Blot.



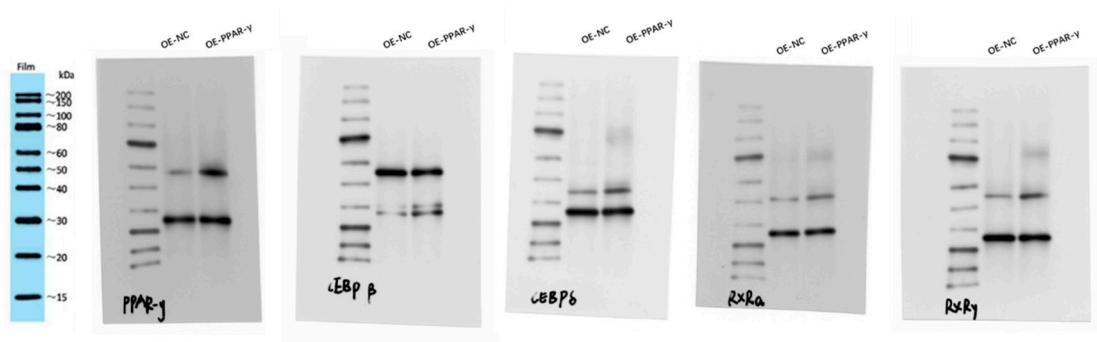
Supplementary Figure S2: The original protein blots of Figure 4A. The transfection efficiency of OE-PPAR- γ was assessed by Western Blot.



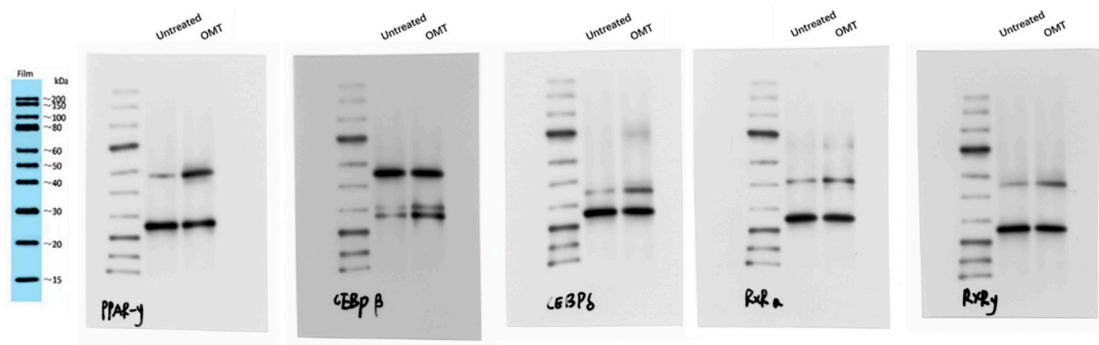
Supplementary Figure S3: The original protein blots of Figure 5E. Western Blot analysis were employed to examine the effects of miR-27a-3p on the expression and protein levels of PPAR- γ .



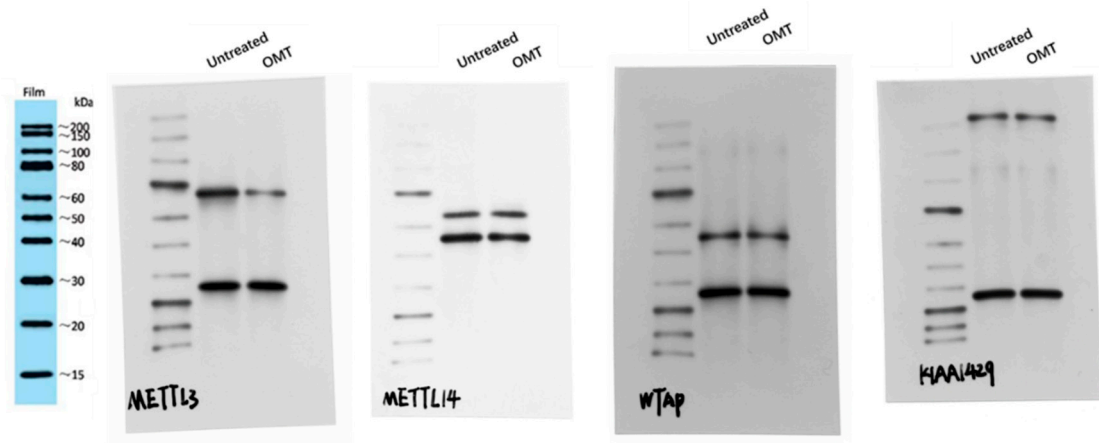
Supplementary Figure S4: The original protein blots of Figure 7C. Western Blot was used to detect the effect of miR-27a-3p-inhibitor.



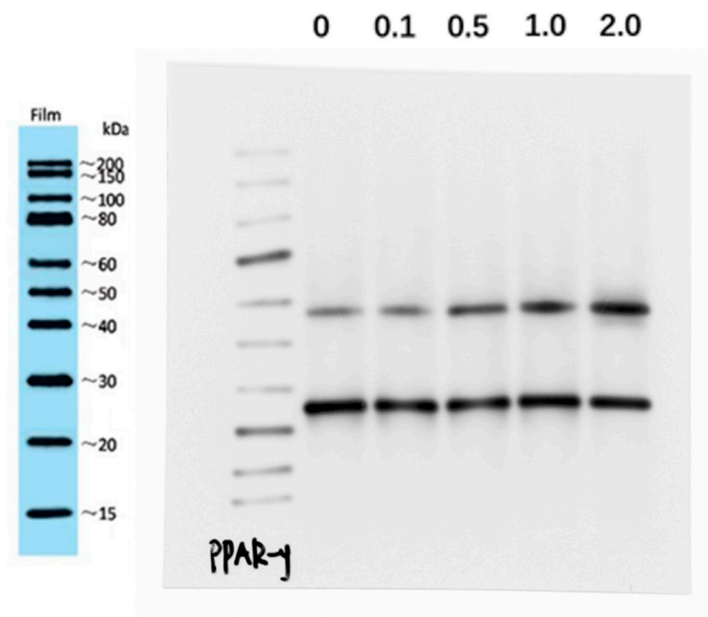
Supplementary Figure S5: The original protein blots of Figure 7D. OE-PPAR- γ and its effect on the expression of proteins related to adipogenesis in HemSCs, including PPAR- γ , C/EBP β , C/EBP δ , RXR α , and RXR γ .



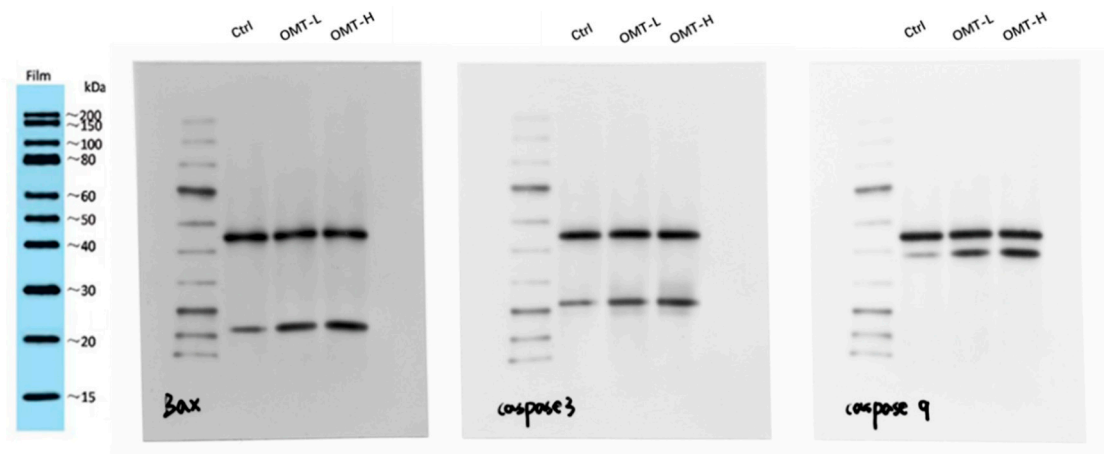
Supplementary Figure S6: The original protein blots of Figure 9B. Western Blot detected the effect of OMT on HemSCs cell fat differentiation-related proteins.



Supplementary Figure S7: The original protein blots of Figure 10A. Western Blot detects the effect of OMT treatment on the protein levels of HemSCs/PPNL drug-resistant cells METL3, METL14, WTAP and KIAA1429.



Supplementary Figure S8: The original protein blots of Figure 10F. Western Blot detection of the effect of OMT on HemSCs cell PPAR-γ protein level.



Supplementary Figure S9: The original protein blots of Figure 11D. Western Blot detection of apoptosis-related proteins under different treatments the protein expression of Bax, Caspase 3 and Caspase 9.