## **Supplementary Information for:**

## TPPU inhibits inflammation-induced excessive autophagy to restore the osteogenic differentiation potential of stem cells and improves alveolar ridge preservation

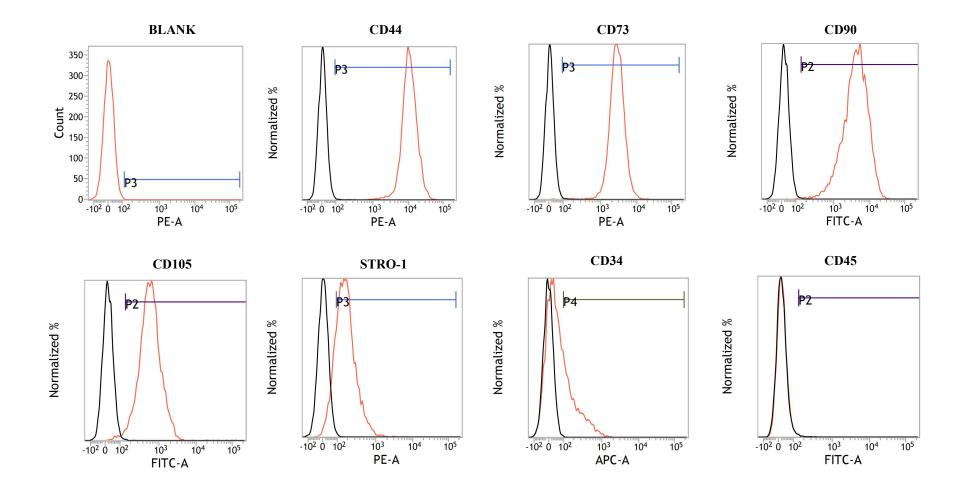
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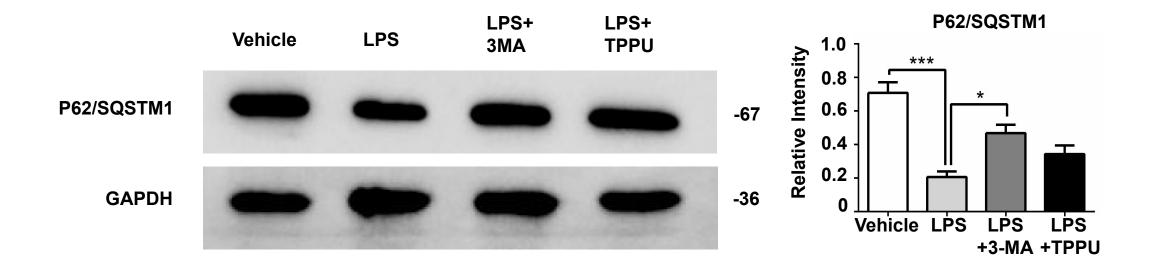
<sup>2</sup>School of Stomatology, Dalian Medical University, Dalian, 116044, China.

<sup>3</sup>Academician Laboratory of Immune and Oral Development & Regeneration, Dalian Medical University, Dalian, 116044, China.

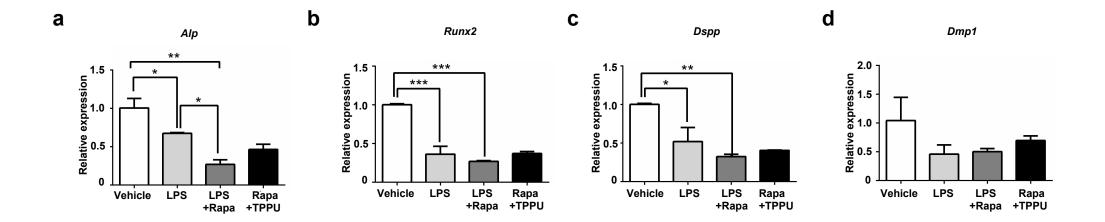
<sup>4</sup>The Affiliated Stomatological Hospital of Dalian Medical University School of Stomatology, Dalian, 116086, China.



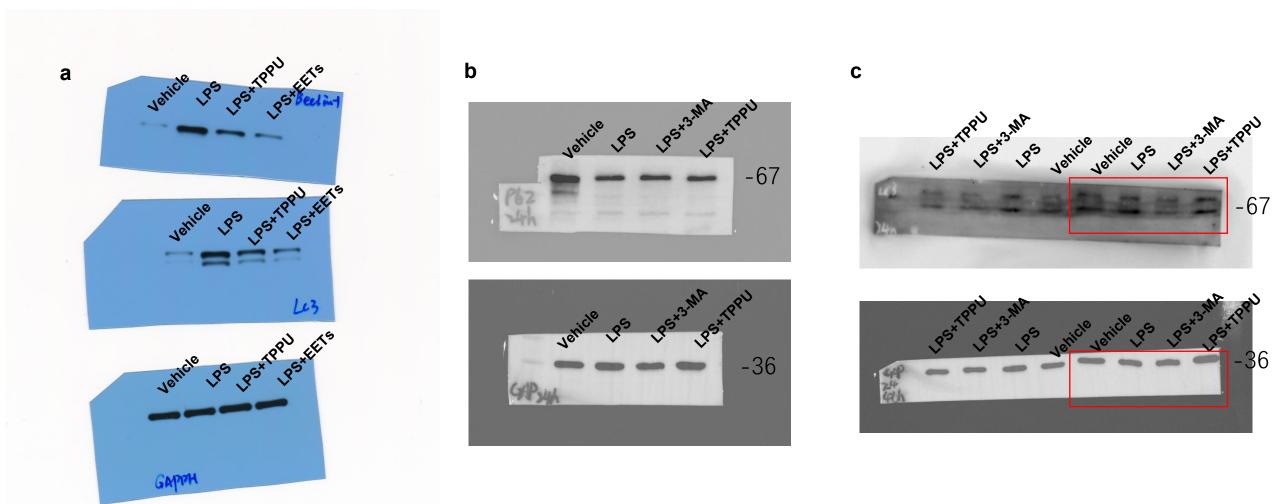
**Supplementary Fig. 1 Flow cytometry showing stem cells surface makers.** The hDPSCs are positive for CD44, CD73, CD90, CD105, and STRO-1, and negative for CD34 and CD45. The black histograms show the isotype control-stained cells and the red histograms show the antibody-stained cells. The control population was stained with matched isotype antibodies.



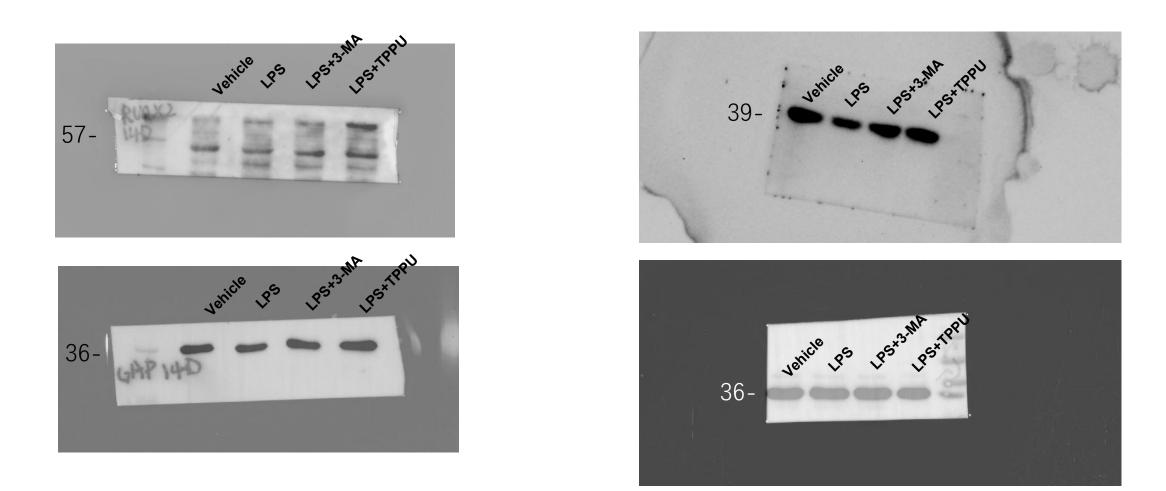
Supplementary Fig. 2 Western blotting for the P62/SQSTM1 expression in hDPSCs. hDPSCs treated with LPS, hDPSCs treated with LPS and 3MA, hDPSCs treated with LPS and TPPU following 48 h of osteogenic induction. Uncropped Western blot images are shown in Supplementary Fig. 6.



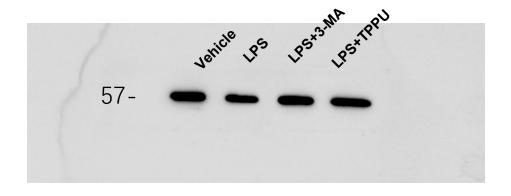
Supplementary Fig. 3 mRNA expression of *Alp, Runx2, Dspp, Dmp1* was evaluated by RT-qPCR in hDPSCs. hDPSCs treated with LPS, LPS and Rapa, Rapa and TPPU following 7 d of osteogenic induction.

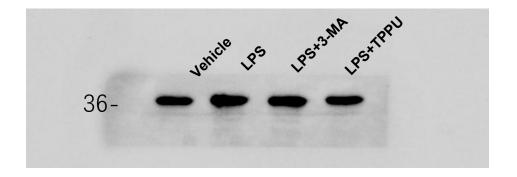


Supplementary Fig. 4 Unprocessed western blots and/or gels. Source Data for Figure 2a (a); Source Data for Figure 2c (b); Source Data for Figure 2d (c)



Supplementary Fig. 5 Unprocessed western blots and/or gels. Source Data for Figure 3c





**Supplementary Fig. 6 Unprocessed western blots and/or gels.** Source Data for Supplementary Figure 2