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Mesenchymal stromal cell-derived extracellular vesicles promote myeloid-biased multipotent hematopoietic progenitor expansion via Toll-like receptor engagement.

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There were several errors in the original figure legend to Fig. 4. The figure legend should be replaced with the following.

FIGURE 4. MSC-derived EVs activate NF- κ B and Notch signaling to promote HSPC differentiation. *A*, diagrammatic representation of the lentiviral vector design used to generate the NF- κ B enhancer-luciferase reporter cell line. Stably transduced and sorted SIM-A9 microglial cells were used for EV exposure followed by luciferase assay. The SIM-A9-NF- κ B-Luc cells were exposed with MSC EVs and incubated for 48 h followed by luciferase assay. Non-treated SIM-A9-NF- κ B-Luc and LPS-treated SIM-A9-NF- κ B-Luc cells were used as negative and positive controls, respectively. Luciferase activity was measured as relative fluorescence units (*RFU*). Data represent $n = 4$ independent experiments; *error bars* depict S.D. All p values have been calculated using two-tailed Student's t test. *B*, quantitative mRNA expression of NF- κ B and Notch-1 signaling and downstream targets involved in cell proliferation in MSC EV-exposed HSPCs. Data represent $n = 3$ independent experiments; *error bars* are S.D. *C*, we next tested a candidate panel of canonical TLR4 responsive cytokines for both transcriptional activation and secretion, either after EV or S100 exposure. Transcriptional analysis of EV-exposed HSPCs reveals an up-regulation of several known TLR4-responsive cytokine genes (*IL6*, *TNFA*, *Stat1*, and *EGFR*) in WT but not *MyD88*^{-/-} HSPCs. Data represent $n = 3$ independent experiments; *error bars* are S.D. *D*, TLR4 signaling was specifically inhibited in WT MSC-EV-exposed HSPCs using TAK-242. Transcriptional analysis of EV-exposed HSPCs with and without TLR4 inhibitor showed down-regulation of the same TLR-responsive genes described above. Data represent $n = 2$ independent experiments; *error bars* represent S.D. *E*, cytokines released by HSPCs after 48 h co-culture, with MSC EVs compared with S100, were measured using a Luminex assay, whereby *error bars* are S.D. and p values are an indication of the variance across three technical replicates in this screening panel.

Authors are urged to introduce these corrections into any reprints they distribute. Secondary (abstract) services are urged to carry notice of these corrections as prominently as they carried the original abstracts.