## CORRIGENDUM

## Rapid mobilization of hematopoietic progenitors by AMD3100 and catecholamines is mediated by CXCR4-dependent SDF-1 release from bone marrow stromal cells

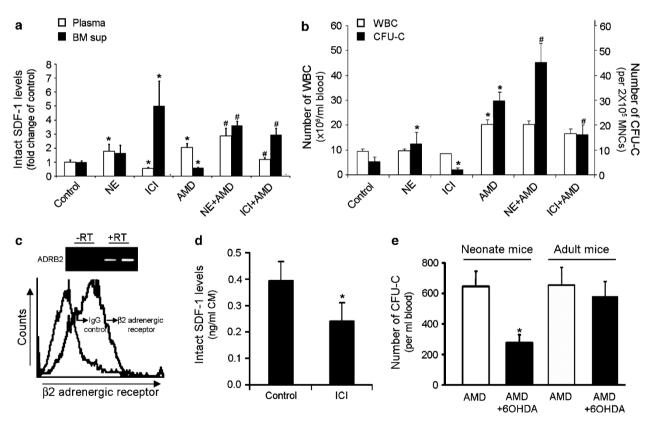
A Dar, A Schajnovitz, K Lapid, A Kalinkovich, T Itkin, A Ludin, W-M Kao, M Battista, M Tesio, O Kollet, NN Cohen, R Margalit, EC Buss, F Baleux, S Oishi, N Fujii, A Larochelle, CE Dunbar, HE Broxmeyer, PS Frenette and T Lapidot

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The authors apologize for any inconvenience caused.

Since the publication of this paper, the authors have noticed an error in Figure 5, concerning the BM SDF-1 levels in the control AMD-only treated mice. The correct version is shown below.



**Figure 5** Neurotransmitter stimulation induces functional SDF-1 release and rapid progenitor mobilization. (**a**, **b**) SDF-1 levels in the plasma and BM sup. (**a**) and circulating WBC and progenitor cells (**b**) in mice treated with NE or the  $\beta$ 2 adrenergic antagonist ICl, 1 h after administration. Control mice received injections of PBS, n = 6 mice/group. Values of plasma SDF-1 levels:  $1.1 \pm 0.17$ ,  $1.8 \pm 0.5$ ,  $0.6 \pm 0.06$ ,  $2 \pm 0.3$ ,  $2.8 \pm 0.5$  and  $1.2 \pm 0.1$  ng/ml, respectively, \*P<0.05 compared with control mice, \*P<0.05 compared with AMD3100-treated mice. (c) RT-PCR analysis (top) for mRNA expression and flow-cytometry analysis (bottom) for cell surface expression of  $\beta$ 2 adrenergic receptor on cultured primary human BMEC. -RT = cDNA was prepared without reverse transcriptase as a control. (**d**) SDF-1 release from primary human BMEC in response to stimulation with ICl (10 ng/ml), n = 3. (**e**) AMD3100-induced mobilization of progenitors in control and sympathectomized (6OHDA) of either neonate or adult mice. \**P*<0.005. n = 10-15 mice/group.

