## Nguyen TD, Shaid S, Vakhrusheva O, et al. Loss of the selective autophagy receptor p62 impairs murine myeloid leukemia progression and mitophagy. *Blood.* 2019;133(2):168-179.

Errata

On page 173 in the 10 January 2019 issue, in each pair of bars in Figure 4D, the left bar (wild type [WT]) should be red and the right bar ( $p62^{-/-}$ ) should be blue. The corrected Figure 4 is shown below. The error has been corrected in the online version of the article.

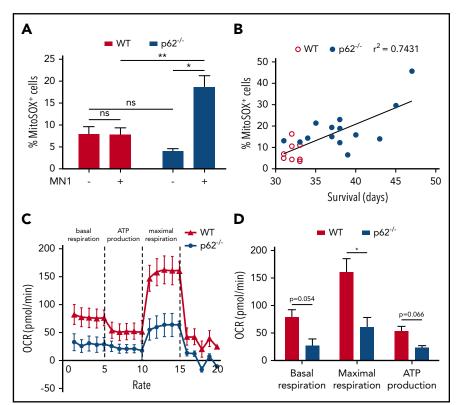


Figure 4. Mitochondrial functions in p62<sup>-/-</sup>leukemia cells were defective. (A) Mitochondrial superoxide levels of LK HSC cells from healthy mice (n = 3per group) and GFP<sup>+</sup> LK blasts from MN1 leukemic mice (n = 8 in WT and n = 14 in  $p62^{-/-}$  leukemic mice) were analyzed by flow cytometry using MitoSOX. (B) Pearson's correlation was used to determine the correlation between the proportions of MitoSOX+ leukemic cells with the survival time of leukemic mice. (C) Mitochondrial respiration of WT and  $p62^{-/-}$  MN1 leukemic mice (n = 3 per group) was determined by measuring the OCR. The experiment was performed in real time by the 96-well Seahorse Bioscience Extracellular Flux Analyzer XF96. (D) The rates of basal respiration, maximal respiration, and adenosine triphosphate (ATP) production were compared between WT and  $p62^{-/-}$  group (n = 3 per group). Values are mean  $\pm$  SEM. ns, not significant; \*P  $\leq$  .05; \*\*P  $\leq$  .01; \*\*\* $P \le .001$ .

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## Artuso I, Lidonnici MR, Altamura S, et al. Transferrin receptor 2 is a potential novel therapeutic target for $\beta$ -thalassemia: evidence from a murine model. *Blood*. 2018;132(21):2286-2297.

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