## CORRECTION

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sepsis-induced lung injury by regulating oxidative stress and inflammation

Correction: Extracellular vesicles derived

from CD4<sup>+</sup>T cells carry DGKK to promote

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The original article can be found online at https://doi.org/10.1186/ s11658-023-00435-y.

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Following publication of the original article [1], the authors identified a similarity between the image labeled "CLP+shNC" in Fig. 6C of their article and an image in another publication from their hospital (PubMed: 34616481, Figure 5a). Upon careful investigation, the authors discovered that the image in question had been incorrectly labeled. The error occurred unintentionally due to the fact that two research teams from hospital, both studying acute lung injury, were working in the same laboratory. This resulted in an accidental mislabeling of the H&E histopathology slides because of an incorrect slide number in the shared platform.



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**Fig. 6** DGKK knockdown attenuated tissue damage, oxidative stress, and inflammation in mice with CLP-induced lung injury. Mice with CLP were injected with virus carrying plasmid expressing shRNA control (shNC) and shRNA targeting DGKK (shDgkk). The elevation of Dgkk abundance at the **A** mRNA and **B** protein levels in CLP model mice was restored by DGKK RNAi. **C** H and E staining indicated that the severe lung injury in CLP model mice was restored by Dgkk RNAi (scale bar, 100  $\mu$ m). **D** The severity of histological injury and plasma levels of **E** ALT, **F** AST, and **G** LDH in CLP model mice was restored by DGKK RNAi. The elevation of **H** ROS levels and **I** MDA content in CLP model mice was restored by DGKK RNAi. The reduction of **J** SOD and **K** GPX activities in CLP model mice was restored by Dgkk RNAi. The elevation of **B** ALT, **F** AST, and **G** LDH in CLP model mice was restored by DGKK RNAi. The reduction of **J** SOD and **K** GPX activities in CLP model mice was restored by Dgkk RNAi. The elevation of **L** DAG content and **M** PKC activity in CLP model mice was restored by Dgkk RNAi. **O** The shortened survival in CLP model mice was restored by Dgkk RNAi. **O** The shortened survival in CLP model mice was restored by Dgkk RNAi. **D** the shortened survival in CLP model mice was restored by Dgkk RNAi. **D** The shortened survival in CLP model mice was restored by Dgkk RNAi. **D** The shortened survival in CLP model mice was restored by Dgkk RNAi. **D** The shortened survival in CLP model mice was restored by Dgkk RNAi. **D** The shortened survival in CLP model mice was restored by Dgkk RNAi. **D** The shortened survival in CLP model mice was restored by Dgkk RNAi. **D** The shortened survival in CLP model mice was restored by Dgkk RNAi. **D** The shortened survival in CLP model mice was restored by Dgkk RNAi. **D** The shortened survival in CLP model mice was restored by Dgkk RNAi. **D** The shortened survival in CLP model mice was restored by Dgkk RNAi. **D** The shortened survival in CLP model mice was restored by Dgkk RNAi. **D** The sho

The authors re-examined the original samples and thoroughly reviewed the data, and they are confident that this mistake does not alter the overall conclusions of the article.

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## Reference

 Tu GW, Zhang Y, Ma J, Hou J, Hao G, Su Y, Luo J, Sheng L, Luo Z. Extracellular vesicles derived from CD4+ T cells carry DGKK to promote sepsis-induced lung injury by regulating oxidative stress and inflammation. Cell Mol Biol Lett. 2023;28:24. https://doi.org/10.1186/s11658-023-00435-y.

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