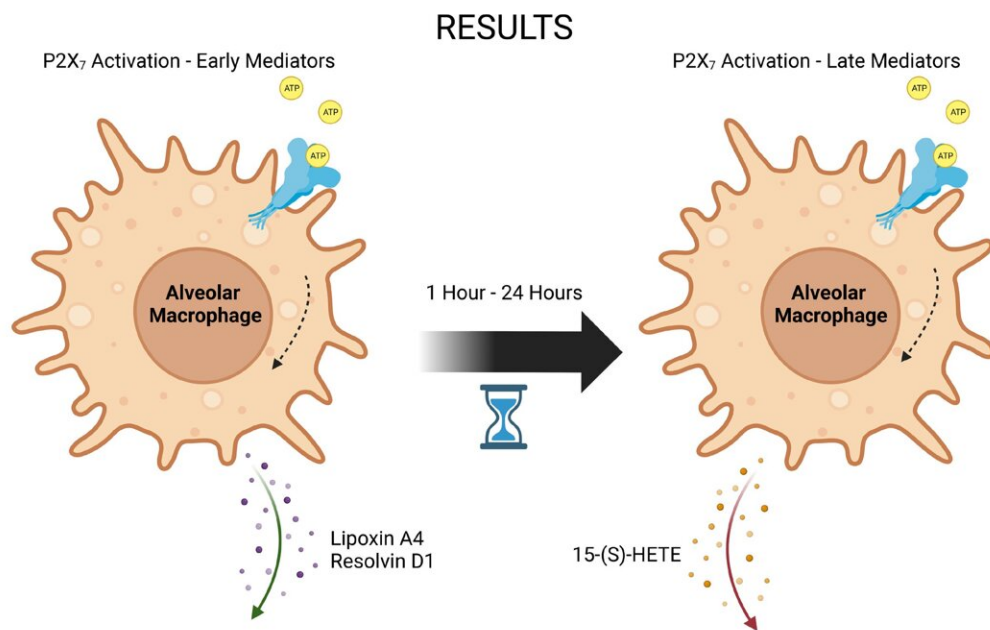


Changing dosing methods means fewer mice needed to study lung infections

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CONCLUSION: P2X₇ receptor activation in alveolar macrophages results in temporally distinct secretion of multiple soluble lipid mediators.

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Credit: *American Journal of Physiology-Lung Cellular and Molecular Physiology* (2023). DOI: 10.1152/ajplung.00070.2023

Researchers will need fewer mice to study lung infections thanks to improvements in dosing methods, according to a new study from the University of California San Francisco (UCSF). Changing how animals are anesthetized and infected with microbes allows scientists to study lung infections using smaller group sizes and without having to use invasive dosing methods. The study is published in the [*American Journal of Physiology-Lung Cellular and Molecular Physiology*](#).

Research using mouse models of [lung infections](#) is important for improving scientists' understanding of how infections lead to lung diseases and for testing vaccines and drugs. When researchers use different anesthesia and dosing approaches in their experiments, the amount of dose that reaches the lungs and the extent of lung inflammation changes.

Raising awareness of these effects to encourage harmonization of dosing methods between different groups of researchers in different laboratories has potential to improve reproducibility and reduce the number of failed experiments.

In this study, researchers compared the effects of delivering [microbes](#) to lungs of mice through three different delivery methods: intranasal (through the nostrils), by oropharyngeal aspiration (placing doses into the back of the mouth) and intratracheal (into the windpipe). "Intratracheal dosing approaches have potential for more precise [lung](#) dosing relative to intranasal and oropharyngeal aspiration dosing," the researchers wrote.

"Optimized dosing methods allow researchers to complete mouse studies faster and at lower cost," said Simon Cleary, Ph.D., a postdoctoral scholar at UCSF and a lead author of the study. "Making preclinical studies more efficient in this way has potential to speed up the development of new and improved treatments for patients. Researchers switching to less invasive methods that mean fewer [animals](#) are required

is also good news for mice."

More information: Elizabeth A. Townsend et al, P2X7 signaling influences the production of pro-resolving and pro-inflammatory lipid mediators in alveolar macrophages derived from individuals with asthma, *American Journal of Physiology-Lung Cellular and Molecular Physiology* (2023). [DOI: 10.1152/ajplung.00070.2023](https://doi.org/10.1152/ajplung.00070.2023)

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