

## Drug reduces risk of pneumonia in newborn mice

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The underdeveloped lungs and immune systems of premature babies put them at high risk of potentially deadly pneumonia. Now, a study from Washington University School of Medicine in St. Louis has found, in

mice, that an inhaled drug promotes the development of lung immunity and reduces the risk of pneumonia.

The findings, published Aug. 14 in *Science Advances*, suggest that the drug—an inhalable form of a normal immune protein called GM-CSF—could reduce the risk of life-threatening lung infections common among [premature infants](#). Senior author S. Celeste Morley, MD, Ph.D., an associate professor of pediatrics, and of pathology and immunology, discussed the study.

## **Why did you start exploring whether GM-CSF could prevent pneumonia?**

Immune cells called macrophages are critical for controlling lung infections, and they appear in the lungs in the first day or two after birth. We know that newborn [mice](#) naturally experience a burst of GM-CSF in their lungs in the first 24 to 48 hours after birth that triggers the development of specialized lung macrophages. We don't know for sure whether the same thing happens in babies. But we think that full-term babies also get a burst of GM-CSF, and preemies might not. If so, the macrophages in the lungs of premature [infants](#) would not develop normally, which could explain why preemies are at such high risk of [pneumonia](#).

## **What happened when you gave mice GM-CSF?**

We used mice with a [genetic defect](#) that prevents their lung macrophages from maturing normally. We couldn't study premature mice—they're just too small to work with. But full-term newborn mice with this genetic defect are very susceptible to bacterial infection and pneumonia, like human preemies, so we used them instead. We gave these newborn mice GM-CSF intranasally for the first 24 hours after birth. The number of

lung macrophages increased immediately, and since they are very long-lived cells, the mice were protected from pneumonia even into adulthood.

## **GM-CSF has been tried before as a way to protect preemies from pneumonia. Why didn't it work then, and why do you think it will work now?**

People have tried giving premature infants GM-CSF intravenously or subcutaneously. When you give GM-CSF this way, it mobilizes neutrophils, a different immune cell that is also very important for fighting off infections in general, but is not the crucial immune cell type in the lungs. So it wasn't a bad idea, but the clinical trials have been disappointing. GM-CSF injections did not lower the risk of pneumonia. What we did differently is we administered the drug to the lungs directly, by inhalation, which triggered the development of [lung](#) macrophages instead of neutrophils. And then we saw long-lasting protection against pneumonia.

## **Could inhaled GM-CSF help protect premature infants from pneumonia?**

I hope so. The key point is that it has to be given within the first day after birth. These specialized macrophages are only made in the first couple of days after [birth](#). After that, the maturation window is over and you've missed your chance to intervene. And you have to give it through inhalation so it gets directly to the lungs. GM-CSF is an FDA-approved drug, and it has been used safely in previous clinical trials in premature infants. We still have some questions to answer, but I hope we can take this into clinical trials soon.

**More information:** Elizabeth M. Todd et al. Inhaled GM-CSF in

neonatal mice provides durable protection against bacterial pneumonia, *Science Advances* (2019). [DOI: 10.1126/sciadv.aax3387](https://doi.org/10.1126/sciadv.aax3387)

Provided by Washington University School of Medicine in St. Louis

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