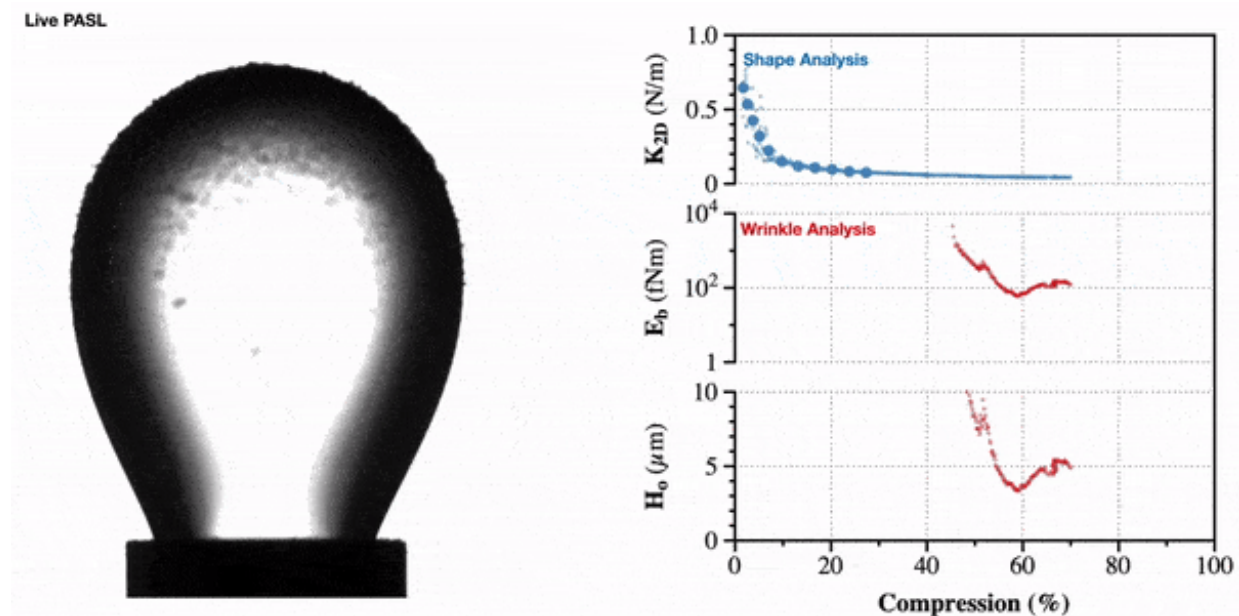


New research examines mechanics of mucus in cystic fibrosis patients

October 28 2020



Researchers used pendant drop elastometry to compress and expand the biofilm that the PASL cells formed. Credit: Tagbo Niepa / University of Pittsburgh Swanson School of Engineering

If you've ever had a cold, you know that too much mucus can be an annoyance, but mucus plays a very important role in the body. The respiratory system creates mucus as part of the immune system, meant to trap inhaled bacteria, viruses, and dirt so they can be removed before causing infection. However, for people with the genetic disorder cystic

fibrosis (CF), the mucus that their bodies produce is thicker and stickier, leading to an increased risk from infection and decreased ability to breathe over time.

New research led by the University of Pittsburgh's Swanson School of Engineering examines the properties of the [mucus](#) of CF patients and the role it plays in a pathogens' ability to survive. The new information could have important implications for CF treatment.

The researchers examined nonmucoid (PANT) and mucoid (PASL) strains of *P. aeruginosa*, a common pathogen that infects the lungs. *P. aeruginosa* adapts to the host environment mutating from a non-mucoid phenotype (PANT) to a mucoid phenotype (PASL). This mutation in *P. aeruginosa* creates a protective film of mucus around the bacteria thereby forming a more hydrated and slimy biofilm in the mucus.

"Think of the [cells](#) like grains of rice. PANT cells are like basmati rice, while PASL cells are like sushi rice: coated in such a way that they stick together when they're compressed," explained Tagbo Niepa, assistant professor of chemical and petroleum engineering, whose lab led the study. Niepa also has appointments in the Departments of Bioengineering, Civil and Environmental Engineering, and Mechanical Engineering and Materials Science. "We can measure how investigational drugs can alter the sticky nature of the coating that pathogens such as *P. aeruginosa* create upon mutation."

This mutation gives the mucus [unique properties](#) that contribute to increased antibiotic resistance. It also shields them against phagocytic cells, which help the immune system clear out dead or harmful cells by ingesting them.

In order to study these properties, the researchers used pendant drop elastometry to compress and expand the biofilm that the cells formed.

They also assessed the transcriptional profile of the cells to correlate the film's mechanics to cell physiology.

"This is the first time that the pendant drop elastometry technique has been used to study the mechanics of these cells. We demonstrate that these techniques can be used to investigate the efficacy of mucolytic drugs—drugs that are used to break down the film of mucus that the cells are making," noted Niepa. "This technique could be powerful for investigating those agents, to see if they have the anticipated effect."

More information: Sricharani Rao Balmuri et al, Material properties of interfacial films of mucoid and nonmucoid *Pseudomonas aeruginosa* isolates, *Acta Biomaterialia* (2020). [DOI: 10.1016/j.actbio.2020.10.010](https://doi.org/10.1016/j.actbio.2020.10.010)

Provided by University of Pittsburgh

Citation: New research examines mechanics of mucus in cystic fibrosis patients (2020, October 28) retrieved 3 May 2024 from <https://medicalxpress.com/news/2020-10-mechanics-mucus-cystic-fibrosis-patients.html>

<p>This document is subject to copyright. Apart from any fair dealing for the purpose of private study or research, no part may be reproduced without the written permission. The content is provided for information purposes only.</p>
--