

Researchers' method holds promise for brain study, better tests for viruses

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Dr. Haihang Ye, research associate, synthesizes nanoparticles that will be used in a new infectious disease testing method that he and fellow researchers are developing. Credit: University of Texas at Dallas

University of Texas at Dallas researchers have developed a promising method for remotely stimulating activity in deep brain regions, advancing understanding of how molecules act in the brain and paving the way for better cancer treatments and therapies for other diseases.

The approach is based on the powerful combination of gold nanoparticles and lasers, which also plays a critical role in another UT Dallas research project aimed at developing a rapid diagnostic test for influenza and, possibly, the COVID-19 virus.

Gold for Neuromodulation

Light is an important tool to modulate [biological systems](#), but absorption and scattering in biological tissues significantly limit its penetration. The system developed by researchers in the Erik Jonsson School of Engineering and Computer Science and the School of Behavioral and Brain Sciences packages molecules inside microscopic gold-coated capsules, or nanovesicles, that can be very sensitive to near-[infrared light](#)

The system could solve challenges in treating diseases, such as ensuring that medication is delivered to difficult-to-reach tumors in deep brain regions while reducing damage to healthy tissue. Using that example, the nanovesicles and their cargo are injected into the brain tissue. External near-infrared lasers that penetrate the tissue cause the capsules to open and release the drug. The researchers describe the approach and results of tests in an [animal model](#) in an article published online in the chemistry journal *Angewandte Chemie*.

"Our system converts light to a mechanical wave that shakes the vesicle open," said Dr. Zhenpeng Qin, assistant professor of mechanical engineering at UT Dallas and corresponding author of the study.

Other researchers have used near-infrared light to trigger drug-carrying nanoparticles, such as phospholipid liposomes, which release their cargo when heated by the laser, but Qin's approach with gold-coated nanovesicles uses about 40 times less laser energy.

In tests in animal models, Qin and his colleagues found that the near-infrared light penetrated 4 millimeters in the brain, which was enough to reach most targeted brain regions. Qin said he anticipates the laser penetrates far enough to reach targets deep in the rodent brain that will help answer important questions in neuromodulation.

While the nanovesicle system must undergo more development and testing before it could be used in [clinical care](#), Qin said the approach eventually could be applied to neurological disorders or other cancers. Dr. Hejian Xiong, research associate in Qin's lab and co-author of the journal article, received a new postdoctoral fellowship from the Phospholipid Research Center in Germany to study the use of gold-coated nanovesicles and ultra-short near-infrared lasers to target and relieve pain in patients after surgery. The project aims to provide an adjustable pain management system that could reduce the need for opioids.

Infectious Disease Testing

In a separate research project, Qin recently received a \$293,000 grant from the Congressionally Directed Medical Research Programs to develop a rapid, accurate and less expensive test for infectious diseases, including influenza, that could be conducted in doctors' offices. The testing principle could also be applied to diagnose COVID-19.

While many doctors conduct rapid flu tests on site, the tests can miss influenza in 30% to 50% of cases, according to the Centers for Disease Control and Prevention. Samples must be sent to a lab for an accurate

diagnosis, which can take days.

"We want to improve the tests' sensitivity so that doctors can make the judgment call right in front of the patient, to be able to say either you have it or you don't have it," Qin said.

In the testing method, gold nanoparticles are attached to antibody molecules that can recognize and bind with protein molecules found on the surfaces of viruses. Researchers apply short laser pulses to activate the nanoparticles to generate nanoscale bubbles, or nanobubbles. The accumulation of the nanobubbles signals the presence of a virus.

"By using optics to detect and count the nanobubbles, we can sensitively and quickly detect the presence of specific respiratory viruses," Qin said.

One of the advantages of the approach is that it would not require extensive sample preparation, Qin said. The method could help doctors diagnose viruses much faster and reduce health care costs by eliminating the need for expensive lab visits. The approach could be used to detect a single virus or multiple viruses.

Ultimately, researchers envision the test being broadly used in hospitals and clinics that do not have labs; however, the diagnostic method will need to be tested further before it can be made widely available.

Qin's group is not working with the live coronavirus, only with viral genes, proteins and antibodies. Qin has previously obtained such patient samples for his research on respiratory syncytial virus and influenza.

More information: Hejian Xiong et al. Near-infrared Light Triggered-release in Deep Brain Regions Using Ultra-photosensitive Nanovesicles, *Angewandte Chemie* (2020). [DOI: 10.1002/ange.201915296](https://doi.org/10.1002/ange.201915296)

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