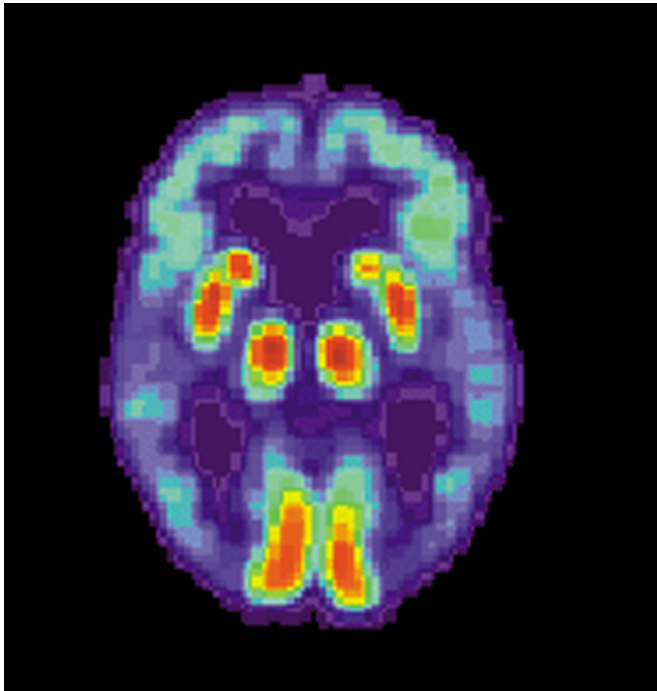


# Team uses copper to image Alzheimer's aggregates in the brain

24 November 2020, by Diana Yates



PET scan of a human brain with Alzheimer's disease.  
Credit: public domain

A proof-of-concept study conducted in a mouse model of Alzheimer's disease offers new evidence that copper isotopes can be used to detect the amyloid-beta protein deposits that form in the brains of people living with—or at risk of developing—Alzheimer's.

Several types of isotopes give off positively charged particles called positrons that are detectable by positron emission tomography scanners. The copper isotope used in the study, Cu-64, lasts much longer than the carbon or fluorine isotopes currently approved for use in human subjects, researchers report. Having access to longer-lasting diagnostic agents would make the process of diagnosing Alzheimer's more accessible to people who live far from major

medical centers. Any clinic with a PET scanner could have the agents shipped to it in time to use the compounds in [brain](#) scans of patients living nearby.

University of Illinois at Urbana-Champaign researchers report their findings in the *Proceedings of the National Academy of Sciences*.

The effort to develop copper-based compounds to detect Alzheimer's disease in living patients is a complicated affair, said Liviu Mirica, a chemistry professor who led the new study with research scientist Hong-Jun Cho. Any diagnostic agent created in the lab must meet several criteria.

"There is a part that binds copper and another part that binds to these amyloid peptides," Mirica said.

In tests with compounds created in Mirica's lab, the team discovered that the copper-binding region of the molecule interfered with the amyloid-binding fragment. To overcome this problem, the researchers introduced a tiny chemical spacer between the two components. This improved their molecule's affinity for the amyloid peptides.

To be effective, however, such [diagnostic agents](#) also must be able to cross the blood-brain barrier.

"They have to be small enough and greasy enough to make it into the brain," Mirica said. "But they can't be too greasy, because then they might not be bioavailable."

The [imaging agent](#) must last long enough for imaging but ultimately decay, leaving no potentially problematic radioactive metals in the body or the brain.

When they first tested their compounds in mouse brain tissue, the researchers saw that their agents' affinity for amyloid deposits was limited. Adding a second amyloid-binding component to the molecule

enhanced its binding and improved its ability to pass through the blood-brain barrier.

"If we do live PET imaging of mice with and without Alzheimer's disease pathologies, we see a statistically significant difference in signal intensity," Mirica said.

**More information:** Design of a multivalent bifunctional chelator for diagnostic  $^{64}\text{Cu}$  PET imaging in Alzheimer's disease, *Proceedings of the National Academy of Sciences* (2020). DOI: [10.1073/pnas.2014058117](https://doi.org/10.1073/pnas.2014058117) , [www.pnas.org/content/early/2020/11/23/2014058117](https://www.pnas.org/content/early/2020/11/23/2014058117)

Provided by University of Illinois at Urbana-Champaign

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