

New chemical reagent turns mouse brain transparent

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Figure 1

Two mouse embryos, one (right) incubated in ScaleA2 solution. Credit: RIKEN

Japanese researchers at RIKEN have developed a ground-breaking new aqueous reagent which literally turns biological tissue transparent. Experiments using fluorescence microscopy on samples treated with the reagent, published this week in *Nature Neuroscience*, have produced vivid 3D images of neurons and blood vessels deep inside the mouse brain. Highly effective and cheap to produce, the reagent offers an ideal means for analyzing the complex organs and networks that sustain living systems.

Our understanding of [biological organisms](#) and how they function is intrinsically tied to the limits of what we can actually see. Even today's

most promising techniques for visualizing biological tissue face this limitation: mechanical methods require that samples be sectioned into smaller pieces for visualization, while optical methods are prevented by the scattering property of light from probing deeper than 1mm into tissue. Either way, the full scope and detail of the biological sample is lost.

The new [reagent](#), referred to as Scale and developed by Atsushi Miyawaki and his team at the RIKEN Brain Science Institute (BSI), gets around these problems by doing two things together that no earlier technique has managed to do. The first is to render biological tissue transparent. Scale does this significantly better than other clearing reagents and without altering the overall shape or proportions of the sample. The second is to avoid decreasing the intensity of signals emitted by genetically-encoded fluorescent proteins in the tissue, which are used as markers to label specific cell types.

This combination makes possible a revolution in optical imaging, enabling researchers to visualize fluorescently-labeled brain samples at a depth of several millimeters and reconstruct neural networks at sub-cellular resolution. Already, Miyawaki and his team have used Scale to study neurons in the mouse brain at an unprecedented depth and level of resolution, shedding light onto the intricate networks of the cerebral cortex, hippocampus and white matter. Initial experiments exploit Scale's unique properties to visualize the axons connecting left and right hemispheres and blood vessels in the postnatal hippocampus in greater detail than ever before.

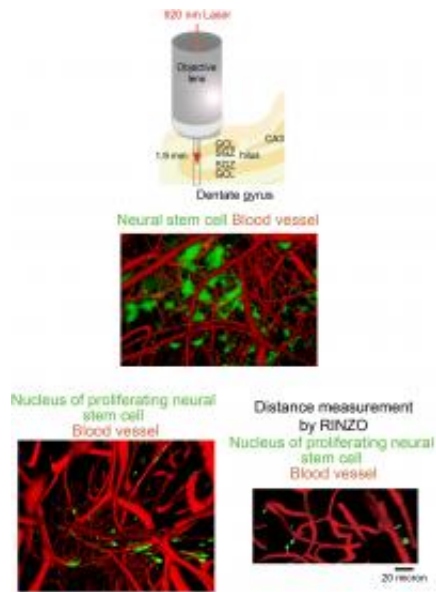


Figure 7

Top: Schematic diagram showing the approach of TPEFM imaging (red arrow) to a cleared excised hippocampus. Middle: Visualization of neural stem cells (NSCs) labeled with green fluorescent protein (GFP) and Texas Red-labeled blood vessels in the adult mouse hippocampus. Bottom: An image of nuclei of proliferating neural stem cells (green) and blood vessels (red) when tunneling into the transparent hippocampus. The green signal comes from the Fucci S/G2/M marker. Credit: RIKEN

But the potential of Scale goes much further. “Our current experiments are focused on the [mouse brain](#), but applications are neither limited to mice, nor to the brain,” Miyawaki explains. “We envision using Scale on other organs such as the heart, muscles and kidneys, and on tissues from primate and human biopsy samples.”

Looking ahead, Miyawaki’s team has set its sights on an ambitious goal. “We are currently investigating another, milder candidate reagent which would allow us to study live tissue in the same way, at somewhat lower levels of transparency. This would open the door to experiments that have simply never been possible before.”

More information: Hiroshi Hama, et al. "Scale: a chemical approach for fluorescence imaging and reconstruction of transparent mouse brain." *Nature Neuroscience*, 2011, [DOI: 10.1038/nn.2928](https://doi.org/10.1038/nn.2928)

Provided by RIKEN

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