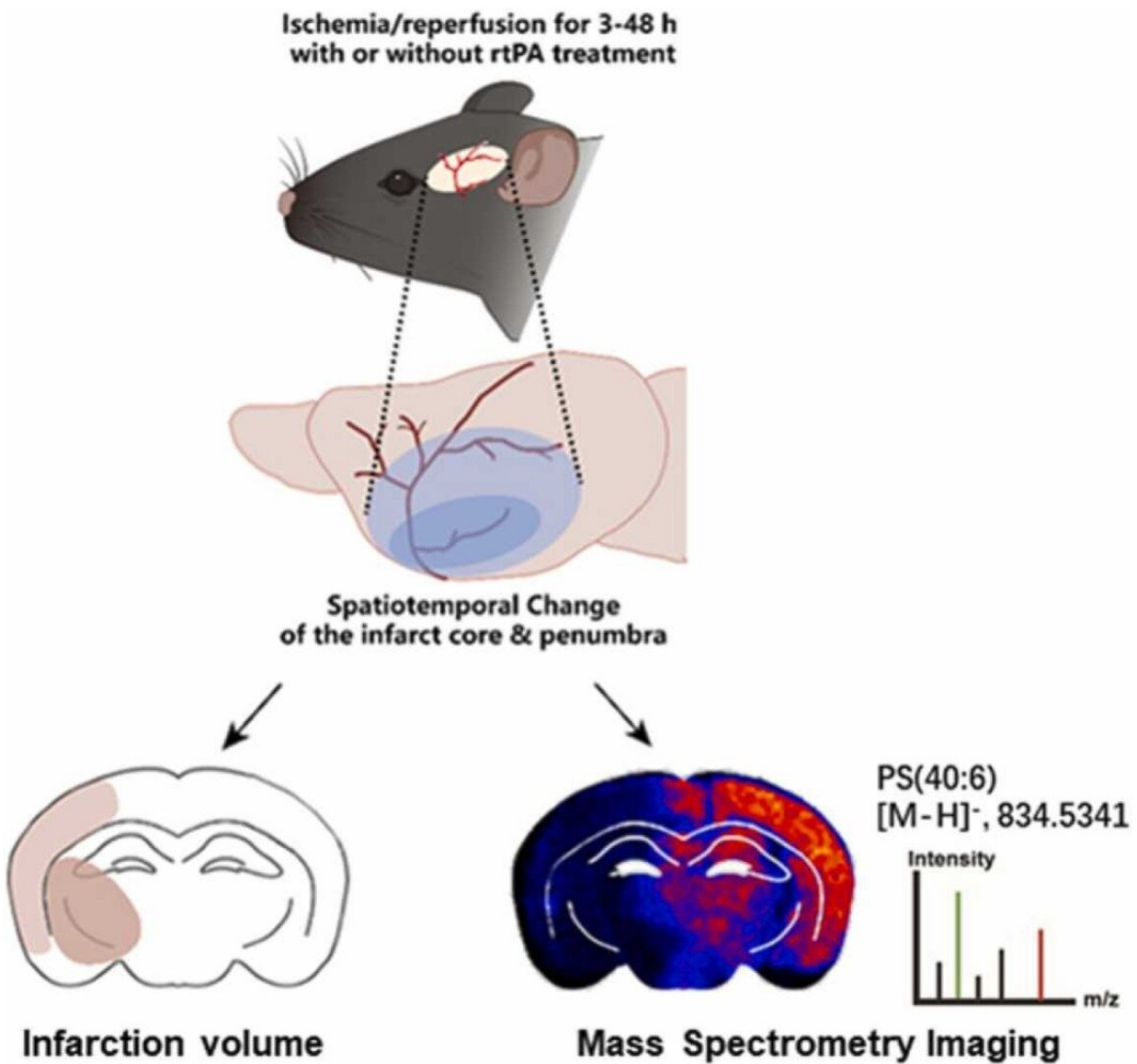


# Researchers demonstrate spatiotemporal dynamic changes in brain lipids

November 1 2022, by Li Yuan



Graphical abstract. Credit: *Pharmacological Research* (2022). DOI:

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Reperfusion therapy, including intravenous recombinant tissue plasminogen activator (rtPA) and/or endovascular thrombectomy, is the most effective treatment for ischemic stroke. However, there is uncertainty regarding the benefits and risks for recanalization since a large proportion of stroke patients remain severely disabled even after receiving timely reperfusion therapy.

It remains unclear how [reperfusion therapy](#) results in secondary injury to the brain tissue and whether different reperfusion therapies induce different effects.

Recently, a joint research group led by Prof. Chang Junlei and Prof. Luo Qian from the Shenzhen Institute of Advanced Technology (SIAT), Chinese Academy of Sciences (CAS) explored the dynamic process of cerebral ischemia/[reperfusion injury](#) using desorption electrospray ionization (DESI)-mass spectrometry imaging (MSI).

They experimentally determined the quantitative and qualitative alterations in brain lipids during the occurrence and progression of ischemia/reperfusion injury in a mouse model of transient middle cerebral artery occlusion treated with rtPA.

Their study was published in *Pharmacological Research* on Oct. 3.

Lipids are the most common biomolecules found in the central nervous system, and comprise up to 50% of its dry weight. Lipids have a variety of biological functions in which they can not only provide structural integrity for [cell membrane](#) and energy reservoir, but also as precursors for various second messengers for cell signal transduction.

The team discovered that the [cerebral blood flow](#) in the ischemic hemisphere restored slowly within nine hours after reperfusion, and then dropped sharply at nine hours to 12 hours, temporally coinciding with the rapid expansion of the infarction core and decreased lipid subspecies, such as phosphatidylcholine [PC(36:4)] and [PC(38:4)], phosphatidylethanolamine [PE(40:5)], phosphatidylinositol [PI(34:1)], phosphatidylserine [PS(40:6)], and sphingomyelin [SM(36:1)]. These lipids could potentially be used as biomarkers of reperfusion injury. Interestingly, some PCs, such as PC(38:7), PC(40:5), showed the opposite trend compared with the others. The regulatory mechanism of any single PC molecules are still open questions.

Moreover, the team discovered that rtPA treatment following reperfusion caused increased brain infarction, cerebral edema, and neurological deficits. Consistently, rtPA treatment caused extensive brain lipid alterations by facilitating brain-wide changes in [lipid metabolism](#) and inducing ischemic region-specific lipid changes.

These results provide novel insights into how reperfusion therapy affects [brain tissue](#) and the outcome of stroke patients, which may facilitate the optimization of the treatment of [ischemic stroke](#).

**More information:** Yinzhong Ma et al, Spatiotemporal lipidomics reveals key features of brain lipid dynamic changes after cerebral ischemia and reperfusion therapy, *Pharmacological Research* (2022). [DOI: 10.1016/j.phrs.2022.106482](https://doi.org/10.1016/j.phrs.2022.106482)

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