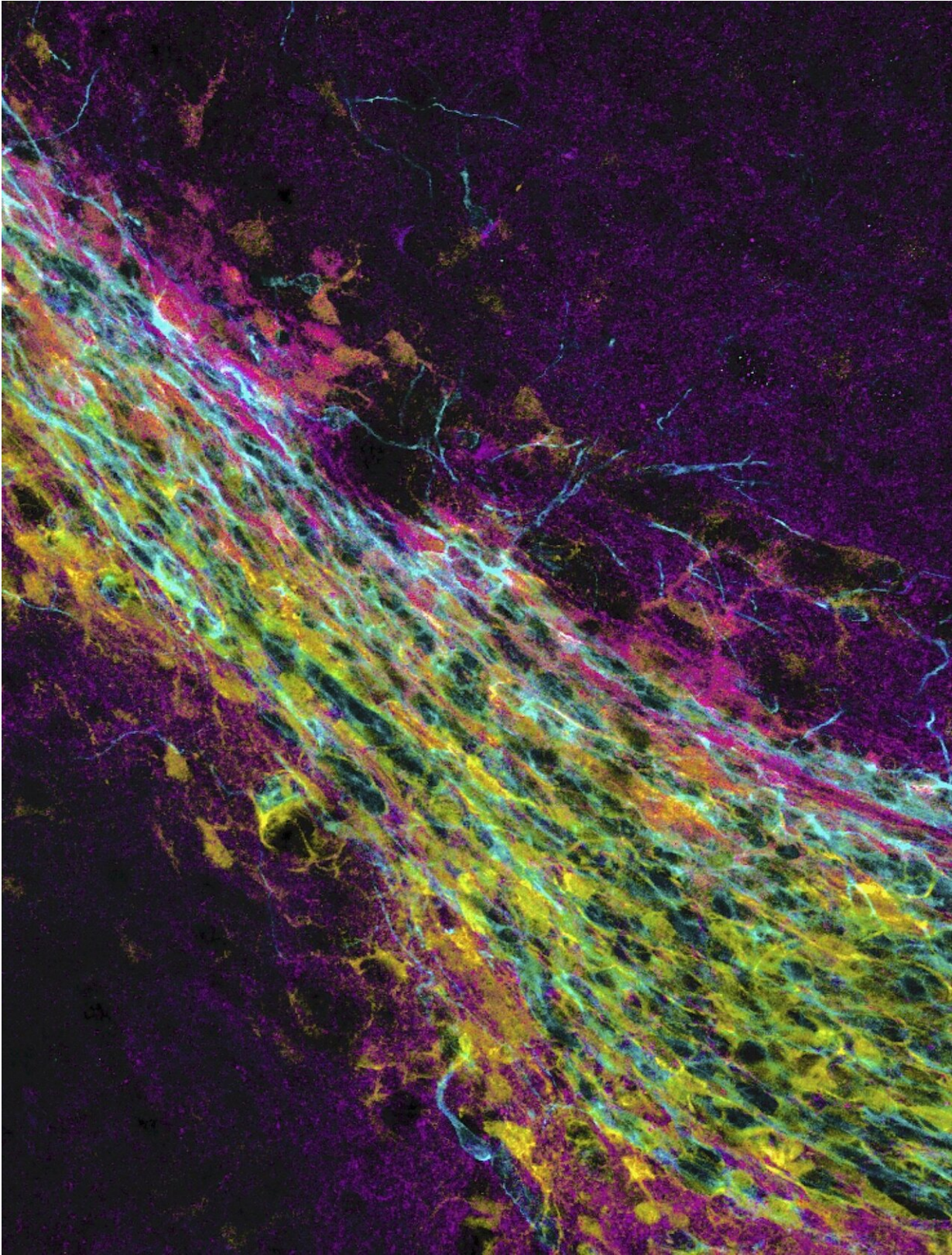


Hard times are coming: Brain tissue stiffness is crucial for neurogenesis

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Neuronal progenitor cells travel from the subependymal zone to the olfactory bulb. Credit: Helmholtz Zentrum München / Jacob Kjell

In mammalian adult brains, neural stem cells are only present in few specific regions, so-called niches. Only these niches are capable of generating new neurons. For the first time, researchers have defined the proteome of these niches, the entire set of expressed proteins, and compared it to other regions of the brain. The findings help to identify key regulators for neurogenesis, an important step towards activating neurogenesis after brain injuries.

What distinguishes neural stem cell niches from other parts of the [brain](#)? Why do only these niches contain [neural stem cells](#) and the rest of the brain does not? So far, we did not know what gives the niches their special function. Also, little was known about factors allowing the integration of new [neurons](#) into existing networks in the adult brain. A comparison of the proteome of neural stem cell niches and the [region](#) in which new neurons integrate with the remaining part of the brain, aimed to shed light about the unique niche environment allowing neurogenesis in the otherwise non-permissive adult mammalian brain.

Defining the niche characteristics

In the first step, Magdalena Götz and her team characterized the proteome of these specific brain regions. They used the largest neural stem cell niche in the subependymal zone of the brain and the olfactory bulb as this is the region where newly-generated neurons from the subependymal zone migrate, differentiate and integrate. They then compared this proteome to the cerebral cortex, an area where neither neurogenesis nor integration of new neurons occurs as in most parts of the adult mammalian brain. As a result, the team found that the

proteome of the neurogenic niche evokes a niche-specific extracellular matrix architecture. One of the most important characteristics was the high solubility of the extracellular matrix. Other characteristic proteins however, such as those of the multifunctional enzyme transglutaminase 2, are hardly soluble meaning strongly cross-linked. Using pharmacological inhibitors as well as genetic experiments, the team was able to show that transglutaminase 2 is crucial for the regulation of neurogenesis. Furthermore, due to its cross-linking properties, the enzyme may contribute to the unique stiffness of the neural stem cell niche in otherwise soft brain tissue.

Turning the injured brain region into a neural stem cell niche

Having established the differences between the [proteome](#) of a neurogenic and a non-neurogenic site is a key step. It could help to find ways to convert the one to the other. Equally important for future cell replacement therapies is the generation of a beneficial environment for the integration of new neurons in the [cerebral cortex](#) by activating factors from the [olfactory bulb](#), an area where new neurons are constantly being integrated. Next, the researchers want to compare these proteomes to the environment after brain injury. Their aim is to create [new neurons](#) after brain injury by turning the injured region into a neurogenic stem cell niche. Injured tissue demonstrates alternative phenotypes, that may create additional barriers for activating neurogenesis: "One of our collaborators at the University of Cambridge showed that scar tissue in the brain is particularly soft—a trait that is hostile to the function of neurogenesis. Overcoming this hurdle and generating a beneficial environment for repair will be the next steps of our research," says Götz.

The study is published in *Cell Stem Cell*.

More information: *Cell Stem Cell*, [DOI: 10.1016/j.stem.2020.01.002](https://doi.org/10.1016/j.stem.2020.01.002)

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