From Actin Dynamics to Membrane Trafficking: Intracellular Mechanism Controlling Astrocyte Reactivity and Scar Formation in the Brain

## June 28

Tuesday, 12:30pm

**Online Webinar** 

For Researchers



Speaker: Britta J. Eickholt, Ph.D. Professor Institute of Molecular Biology and Biochemistry Director of Centrum and Group Leader Signaling Mechanisms in Brain Development and Disease Charité - Universitätsmedizin Berlin

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## Abstract

The nervous system lacks the ability to regenerate nerve cells and is therefore particularly vulnerable to injury. Following brain injury or infection, various cells have to work together in a coordinated manner in order to limit damage and enable recovery. 'Astrocytes', the most common type of glial cell found in the central nervous system, play a key role in the protection of surrounding tissues. They form part of a defence mechanism known as 'reactive astrogliosis', which facilitates scar formation, thereby helping to contain inflammation and control tissue damage. In my seminar, I will present our work describing how the actin cytoskeleton and intracellular membrane trafficking controls essential aspect of astrogliosis responses and how these cellular machineries are essential during scar formation. Our work encompasses biochemistry, cell biology (lots of live cell imaging!!) and in vivo manipulation in rodent models to test our mechanisms.



1. Schiweck, J., Murk, K., Ledderose, J., Munster-Wandowski, A., Ornaghi, M., Vida, I., Eickholt, B.J., 2021. Drebrin controls scar formation and astrocyte reactivity upon traumatic brain injury by regulating membrane trafficking. Nat Commun 12, 1490.

2. Kreis, P., Gallrein, C., Rojas-Puente, E., Mack, T.G.A., Kroon, C., Dinkel, V., Willmes, C., Murk, K., Tom-Dieck, S., Schuman, E.M., Kirstei, J., Eickholt, B.J., 2019. ATM phosphorylation of the actin-binding protein drebrin controls oxidation stressresistance in mammalian neurons and C. elegans. Nat Commun 10.

**3.** Schiweck, J., Eickholt, B.J.\*, Murk, K.\*, 2018. Important Shapeshifter: Mechanisms Allowing Astrocytes to Respond to the Changing Nervous System During Development, Injury and Disease. Front Cell Neurosci 12, 261.



