

**OL76** 

## Controlling the activity of bacterial sialidases with bioorthogonal chemical reporters

Frederic FRISCOURT [1]

[1] Université de Bordeaux, IECB, ISM CNRS UMR5255, Pessac, France f.friscourt@iecb.u-bordeaux.fr

Sialic acids are anionic nine-carbon carbohydrates generally found as terminal sugars of mammalian cell-surface glycoproteins and glycolipids. Because of their distinct cellular location, sialo-glycoconjugates (also known as sialosides) are often key mediators of physiological and pathological events, including cell adhesion, host–pathogen interactions, and cancer progression.<sup>[1]</sup>

The bioorthogonal chemical reporter strategy, which elegantly combines the use of metabolically labeled azido-sugars and highly reactive cyclooctyne probes, is emerging as a versatile technology for labeling and visualizing sialosides.<sup>[2]</sup> This strategy relies on the fact that bioorthogonal chemical reporters are highly reactive species while being biologically noninvasive.

During this talk, I will present our recent efforts to show that chemical bioorthogonal reporters may actually impact sialosides processing enzymes activity.<sup>[3]</sup> More specifically, I will describe how bacterial sialidases may be significantly affected by the presence of bioorthogonal reporters on mammalian cell-surface sialosides, providing us with novel, more selective, chemical biology tools for studying the biological roles of cell-surface glycans.



Factors that can influence bacterial sialidases activity



Chemical (glyco)biology and bioorthogonal chemistry