

Seminar in Microbiology

Monday, 6th June, 2016

Salle de séminaire, E07.3347.a, CMU

11:30 – 12:30



Prof. Dr. Regine Hengge

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Physiological stratification and c-di-GMP signaling in the control of *E. coli* biofilm architecture and morphogenesis.

Regine Hengge's group elucidates the signal transduction pathways and regulatory networks that underlie biofilm formation and architecture in the model organism *Escherichia coli*, including the important pathogen *E. coli* O104:H4. Specifically, the Hengge group is exploring how the bacterial second messenger cyclic-di-GMP controls biofilm formation through a complex and cooperating cyclic-di-GMP regulatory network via of 29 enzymes that control its levels and the c-di-GMP-binding effector proteins. As cyclic-di-GMP is used by most bacteria, understanding its production and function holds promise for the development of anti-biofilm drugs.

Recent publications:

- The green tea polyphenol EGCG inhibits *E. coli* biofilm formation by impairing amyloid curli fibre assembly and down-regulating the biofilm regulator CsgD via the σ^E -dependent sRNA RybB. *Mol Microbiol*, (2016).
- Genome-Based Comparison of Cyclic Di-GMP Signaling in Pathogenic and Commensal *Escherichia coli* Strains. *J Bacteriol*, (2015).
- C-di-GMP signaling and biofilm-related properties of the Shiga toxin-producing 2011 German outbreak *Escherichia coli* O104:H4. *EMBO Mol. Med*, (2014).
- The EAL protein YciR is a trigger enzyme in a c-di-GMP signaling cascade in *E. coli* biofilm control. *EMBO J*, (2013).
- Cellulose as an architectural element in spatially structured *Escherichia coli* biofilms. *J. Bacteriol.*, (2013).
- Principles of cyclic-di-GMP signaling. *Nature Rev. Microbiol*, (2009).
- The BLUF-EAL protein YcgF acts as a direct anti-repressor in a blue light stress response of *E. coli*. *Genes Dev.* (2009).
- Inverse regulatory coordination of motility and curli-mediated adhesion in *Escherichia coli*. *Genes Dev.* (2008).