

## Cluster Seminar

# Prof. Dr. med. Martin Aepfelbacher

Medical specialist in Microbiology, Virology and Infectious Disease Epidemiology



Martin Aepfelbacher

## “Suppression of the innate immune response by a relative of the plague bacterium”

Millions of years of coevolution explain why many pathogens could develop intricate molecular mechanisms to suppress the human immune response. Central to the immunosuppressive activities of the plague agent *Y. pestis* and its enteropathogenic relatives *Y. pseudotuberculosis* and *Y. enterocolitica* is a molecular machine that injects bacterial effectors into immune cells. One group of these effectors subverts crucial immune cell functions such as downstream signaling of toll-like-receptors. However, sometimes bacterial effector activities conflict with the intended immunosuppres-

sion, i.e. because they downregulate the immunosuppressive Interleukin-10 or activate the inflammasome. To counteract these unwanted “side effects” of their own effectors, yersiniae have developed a second group of effectors, which i.e. “rescue” Interleukin-10 production.

I am a medical and cellular microbiologist heading an Institute at the University Medical Centre Hamburg-Eppendorf. In my talk I will address the cell biological effects and structural features of the *Y. enterocolitica* “rescue” effector YopM and how it modulates the inflammatory response in human macrophages.

### WHERE

Lecture Hall 2, BMZ (Building 13)  
Venusberg-Campus 1, 53127 Bonn

### WHEN

Tuesday, January 14, 2020 at 12:30 p.m.

### HOST

Dr. Gregor Hagelueken, Institute of Structural Biology