CRISPR editing using chemically-modified crRNA:tracrRNA complexes

Venue
CMU
Room B02.2526
Rue Michel-Servet 1
1206 Geneva

Speaker
Dr. Mirko Vanetti, European Application Manager - Functional Genomics (IDT)

Host
University Transgenic Core Facility
Fabrizio Thorel, Ph.D, Department of Genetic Medicine and Development (GEDEV)

Agenda
11:15 – 12:00  Seminar
12:00 – 13:00  Sandwich lunch and discussion

CRISPR/Cas is a simple and efficient method of gene editing that utilizes the Cas9 protein and RNA molecules as guides to either disrupt host genes or insert sequences of interest. The prospect of efficiently creating tailored changes to a gene of interest is revolutionizing biomedical research.

The speaker will introduce the basics of CRISPR genome editing. Then, he will discuss the development of the ALT-R CRISPR/Cas system. This system combines recombinant Cas9 protein with length-optimized and chemically modified guide RNAs to form ribonucleoprotein complexes (RNPs). You will hear about using RNPs to perform CRISPR genome editing and tools to optimized reagent delivery and screening. Finally, the presentation will conclude with an overview of novel Cas9 variants with enhanced target specificity and reduced off-target effects.

Registration
Following the seminar, a sandwich lunch will be provided. We therefore kindly ask you to register at
https://crispr_lunchseminar_geneva2019.eventbrite.de