Workshop Bioinformatics analysis of (bulk) RNA-seq data

Organizer

CUBiDA (Core Unit Bioinformatik, Datenintegration und Analyse) at the Universitätsklinikum Erlangen

Instructors

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Outline

The participants will receive an overview of the biological and technical background of the bulk RNA-seq technology, get practical tips about the experimental design, and know the RNA-seq analysis pipeline as well as the downstream analysis steps of the resulting data. The workshop also includes hands-on sessions to explore and perform the different analysis pipeline steps in RStudio.

Outcome

At the end of the workshop, participants will be able to run a typical RNA-seq workflow on a small dataset and understand the essential steps of data exploration, analysis, and interpretation.

Number of participants

5 to 15

Target audience

This workshop is suitable for life and medical researchers who are new to RNA-seq. The course is beginner friendly and intended for those interested in using the command line interface for their analysis.

Prerequisites

This workshop offers a beginner-friendly introduction to RNA-seq analysis. To ensure you can effectively follow along and participate, **basic proficiency in the R statistical environment is essential**. This includes comfort with fundamental R syntax (e.g., variable assignment, function calls, logical operations), core R data structures (vectors, lists, and data frames), basic data frame manipulation and transformation techniques (dplyr, tidyr packages), and R help files. We strongly advise completing an introductory R course before registering for this workshop. In addition, you should be comfortable with **basic file system concepts** (navigating directories, understanding file paths) and **fundamental UNIX commands** (e.g., ls, cd, pwd, cp, mv, rm).

Teaching format

The workshop will comprise a mix of interactive lectures with short hands-on exercises. Computers with preinstalled software and data will be provided to all participants. During the workshop, we will also give you access to all our teaching materials.

Provisional workshop schedule

DAY 1	
8:30 am-	Introduction to RNA-Sequencing.
4:30 pm	Viewing and interpreting FASTQ files.
	Quality control of sequencing reads with FASTQC.
	Adapter and quality trimming of sequencing reads with TrimGalore!
	Mapping sequencing reads to a reference genome/transcriptome.
	Transcript quantification with Salmon.
	DAY 2
8:30 am-	Differential expression analysis with DESeq2.
4:30 pm	Basic visualization of differentially expressed genes. Functional enrichment analysis with gProfiler2.

Aggregation of reports with interactive plots for multiple bioinformatics analyses with MultiQC.

Questions

Please, contact us at mik-cubida@uk-erlangen.de.