

Welcome to Clinical Cancer Genomics, vt 2023

Who are we?

Bioinformaticians @ MEB

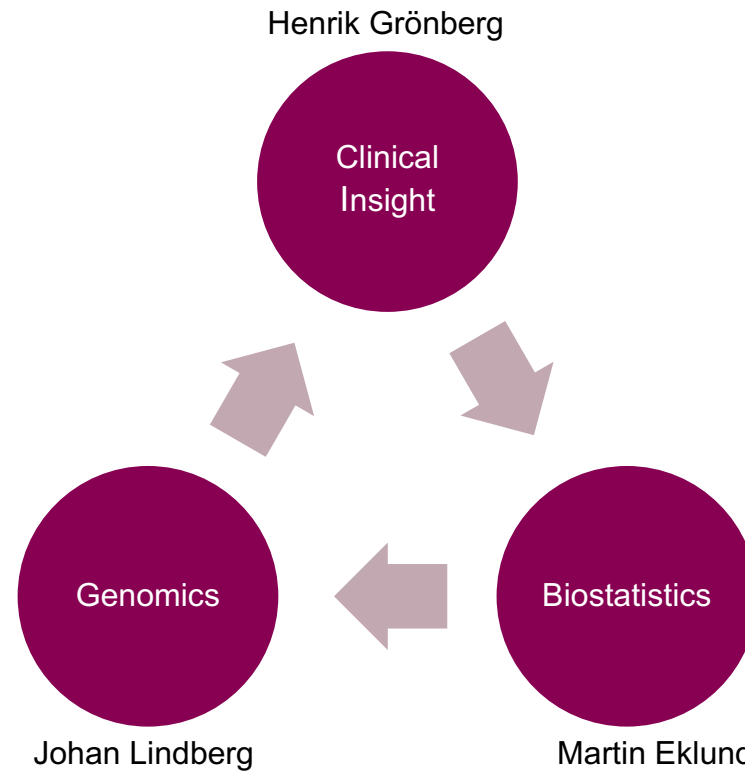
Venkatesh Chellappa
Sarath Murugan
Karthick Maniram
Rebecka Bergstrom
Markus Mayrhofer
Nawal Hamidi
Preeti Lakshman Kumar
Anupriya Sadhasivam

Lab-staff @ SciLife

Mahsan Banijamali
Anastassija Kotsalaynen

Academic postions

Konstantin Carlberg
Sinja Taavitsainen

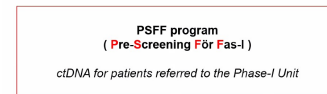
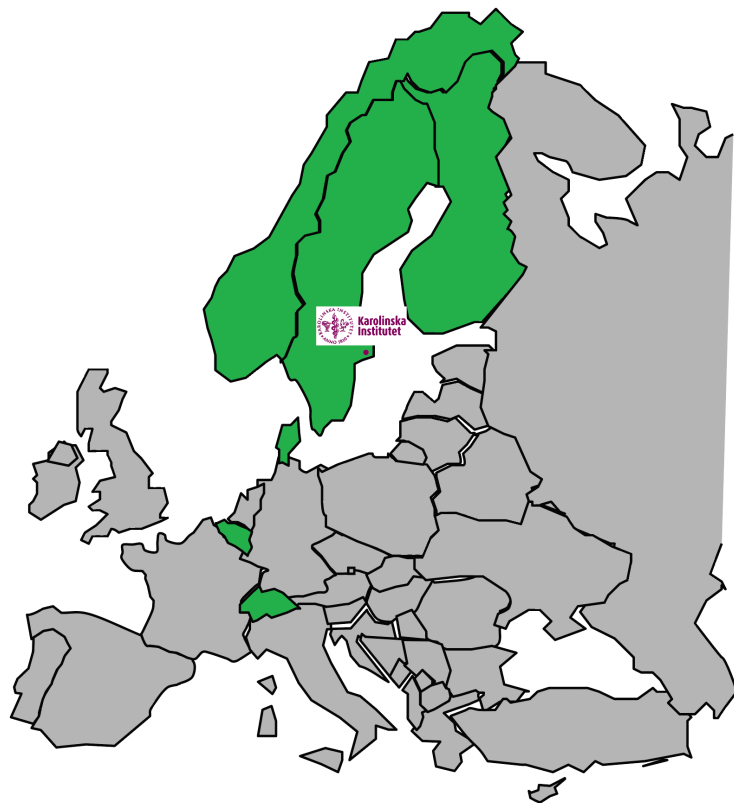


ProBio statisticians

Alessio Crippa
Andrea Discacciati

Others ...

Cancer genomics team at KI



~50 sites in 6 European countries

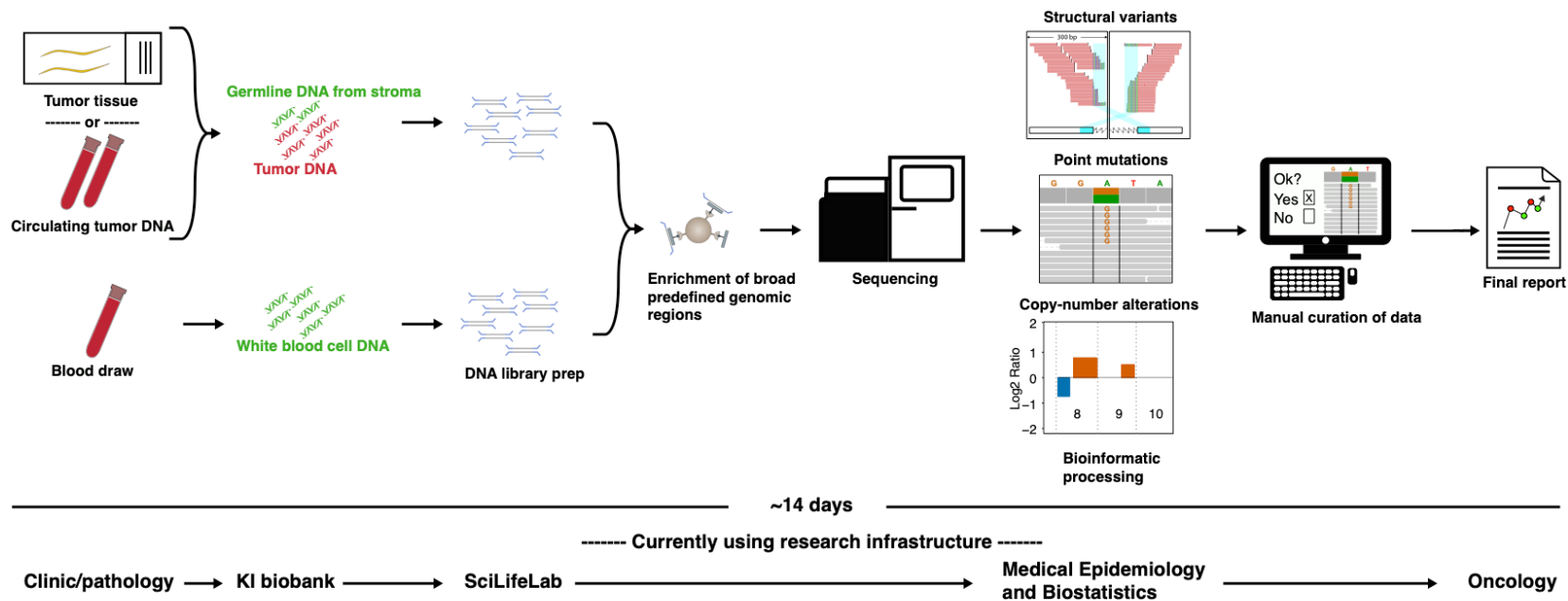


Karolinska Institutet
>5000 cases analysed

Genomics infrastructure

Experience

- > 5000 cases profiled using a validated research infrastructure/process
 - Research projects (DNA and RNA analysis)
 - Prospective clinical trials (DNA analysis)
 - Work towards accreditation and clinical implementation

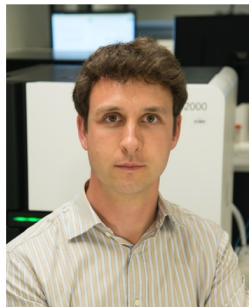


Modified version of an existing course

- <https://pmbio.org/course/>
- Course content under MIT license and Creative Commons.



Malachi Griffith, PhD
Assistant Professor of Medicine
Assistant Professor of Genetics
Assistant Director, MGI

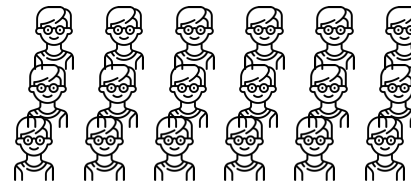


Obi Griffith, PhD
Assistant Professor of Medicine
Assistant Professor of Genetics
Assistant Director, MGI

McDonnell Genome Institute, Washington University School of Medicine

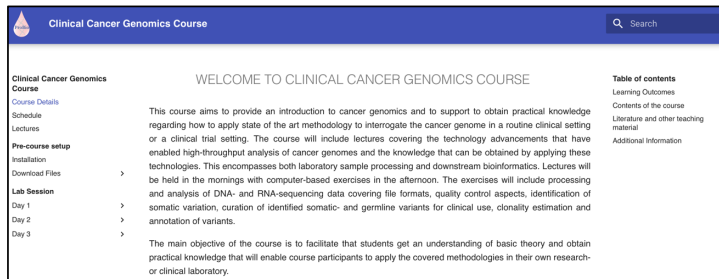
Set-up

Lectures before lunch



Practical sessions after lunch

Course webpage, info, lectures and practical sessions.



Student



AWS server



Course web page

<http://clinical-cancer-genomics.s3-website-us-east-1.amazonaws.com>

Schedule

	Mon	Tues	Wed	Thur	Fri
Lecture 1	Lectures about the cancer genome and how to interrogate it		Small variants (mutations)	Copy number alterations	Continue labwork
Lecture 2		Guest lecture, Clinical Genomics	RNA sequencing	Structural variation	How to curate somatic- and germline variation for clinical use Joint curation of variants
Lecture 3		Lab introduction	Lab introduction	Lab introduction	Curation demonstration, real samples Bioinformatics pipelines & HTC computing environments
Labwork	Tutorials, basic tools for cancer genomics	Files, tools and running a basic bioinformatic pipeline	Calling and QC of somatic/germline variants RNAsequencing analysis	Analysis of copy number data Analysis of structural rearrangements	Continue leftover labwork Genomic Alteration Interpretation Depending on time: TCGA/ICGC integrated genomic analysis.

Set-up

- Examination – show for a course instructor each day that you have completed and understood the labwork.
 - During the exercises you get questions ...
- This is our first time with the course IRL – have patience!
 - Any ideas/things you want to discuss for your own projects – let us know during the practical sessions, we can have a project brainstorm.

Questions?

Menti!