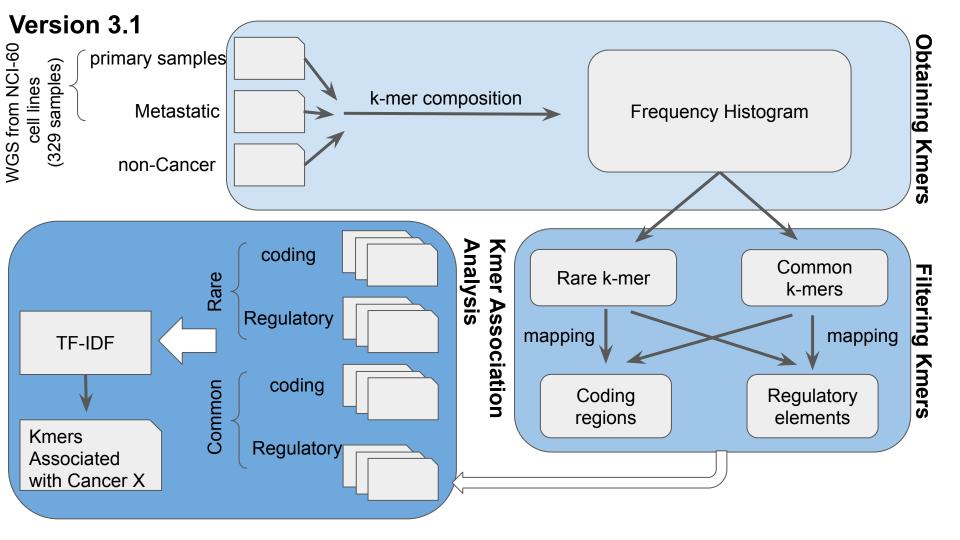


Github: https://github.com/collaborativebioinformatics/kvar



Working in Parallel (Defining Inputs/Outputs)

Obtaining Kmers (Rohit)

INPUT: Fastq file

EXAMPLE: <Standard Fastq File>

OUTPUT: Frequency (count) of each kmer

EXAMPLE: GCTGATCGAC, 2000 / AVG COV per sample

GCTAAACGAC, 0.004 (for example)

Filtering Kmers (Daniela)

INPUT: Frequency (count) of each kmer

EXAMPLE: GCTGATCGAC, 0.0005 GCTAAACGAC, 0.004

OUTPUT: Filter Freq. AND 4x < Standard Bam files>

EXAMPLE: GCTGATCGAC, 0.0004 AND <Standard Bam file>

Kmer Association Analysis (Dreycey)

INPUT: Files with kmers and counts, seperated by region/etc

EXAMPLE: GCTGATCGAC, 0.0004

OUTPUT: A matrix or CSV with TF-IDF calc. AND flat file with disease-associated kmers

EXAMPLE: GCTGATCGAC, TF-IDF score

GOALS / RESPONSIBILITIES

Team Lead - Ben Busby

Writer(s) - Rohit, Danliela

SysAdmins - Dreycey, Quek

- Download primary data -- Quek
 - Find QUICK controls -- Quek (using primary vs metastatic)
- Implement kmer counting -- Rohit
 - First look into using NibSV https://github.com/fritzsedlazeck/nibSV
 - If too cumbersome
 - meryl https://github.com/marbl/meryl
 - Jellyfish https://github.com/gmarcais/Jellyfish
 - KMC https://github.com/refresh-bio/KMC
- Find appropriate (non-cancer) controls -- Daniela (Ben can help)!
- Set up "overlap analysis" -- Daniela
- Set up "tertiary analysis" -- Dreycey