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Project 11410AA-NGS948 data delivery, 23 Oct 2017:

Samples:

Sequencing amount: 20 million read pairs

300 bp paired-end read (Illumina MiSeq V3) Sequencing type:

Delivery contents:

'RAW': raw sequencing data after basecalling in compressed FASTQ format

'AdapterClipped': compressed FASTQ files containing sequencing adapter clipped reads

FastQC reports (https://www.bioinformatics.babraham.ac.uk/projects/fastqc/), containing read quality metrics, are stored along with the FASTQ files.

Data analysis overview:

- Demultiplexing of all libraries for each sequencing lane using the Illumina bcl2fastq 1.8.4 software (folder 'RAW'):
 - 1 or 2 mismatches or Ns were allowed in the barcode read when the barcode distances between all libraries on the lane allowed for it
- Clipping of sequencing adapter remnants from all raw reads (folder 'AdapterClipped'):
 - reads with final length < 20 bases were discarded
- Creation of FastQC reports for all FASTQ files
- Generation of read_counts.xlsx, containing all read counts for all samples at a glance

If you have any questions related to your data or some steps of the data analysis, do not hesitate to contact me directly:

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