NCBI SRA Filter output of Fetch data rRNA contamination database search initial search for analysis / chimera analysis Filtering conditions: Search parameters/queries: Requirements: Analysis steps: 1) BWA mapping of each RNA-seq • Virome & viral Number of reads SILVA SSU + LSU rRNA FASTQ on the SILVA rRNA ref FASTA metagenomics studies (>=100000 & <=15000000) database (removed rRNAs 2) BWA mapping of each RNA-seq • Related to Homo sapiens or Average read size/length with >1% ambiguous N's) FASTQ on the corresponding virus NO human organism (>=100 & <=500) (SILVA rRNA ref FASTA) ref FASTA RNA-seq strategy only Context-based search of • RNA-seq data per sample 3) Detect rRNA-virus chimeric reads as · Single-end & Paired-end virus-related samples (RNA-seg FASTQ) found in common between the • Exclusion of Retro-, DNA-· Studied viral genomes per layout alignments of step 1 & 2 · Illumina & Ion torrent and overrepresented viruses sample (virus ref FASTA) 4) Calculate rRNA contamination & platform in the examined samples • QC of RNA-seg data by fastp rRNA-virus chimera statistics rRNA-virus YES chimeras present? Legend: De novo assembly Implementation of the Individual file operation compared methods statistics / analysis Per RNA-seq sample operation Compared methods: Analysis steps: · RAW method i.e. the raw 1) MEGAHIT de novo assembly on Per compared the RNA-seg FASTQ files treated unprocessed reads method operation • BWA method i.e. the unmapped by each compared method reads after BWA alignment on the 2) Construction of a local BLASTn SILVA rRNA ref FASTA database based on the • SORTMERNA method i.e. the corresponding virus ref FASTA unmapped reads after SortMeRNA 3) BLASTn mapping of the alignment on the SILVA rRNA ref generated MEGAHIT contigs on **FASTA** the local viral database Decision ViRAE method i.e. the clean reads 4) BWA mapping of the generated node after ViRAE mapping and trimming MEGAHIT contigs on the SILVA based on the SILVA rRNA ref FASTA rRNA ref FASTA for chimeric Analysis steps: identification **Process** 1) Implement the compared methods 5) Calculation of *de novo* assembly separately on each RNA-seg FASTQ statistics based on BLASTn output **Process** 2) Measure performance of each report & virus-specific assembly notes method metrics