

Original next flow summary

1. Raw read QC ([FastQC](#))
2. Adapter trimming ([Trim Galore!](#))
 1. Insert Size calculation
 2. Collapse reads ([seqcluster](#))
3. Alignment against miRBase mature miRNA ([Bowtie1](#))
4. Alignment against miRBase hairpin
 1. Unaligned reads from step 3 ([Bowtie1](#))
 2. Collapsed reads from step 2.2 ([Bowtie1](#))
5. Post-alignment processing of miRBase hairpin
 1. Basic statistics from step 3 and step 4.1 ([SAMtools](#))
 2. Analysis on miRBase hairpin counts ([edgeR](#))
 - TMM normalization and a table of top expression hairpin
 - MDS plot clustering samples
 - Heatmap of sample similarities
 3. miRNA and isomiR annotation from step 4.1 ([mirtop](#))
6. Alignment against host reference genome ([Bowtie1](#))
 1. Post-alignment processing of alignment against host reference genome ([SAMtools](#))
7. Novel miRNAs and known miRNAs discovery ([MiRDeep2](#))
 1. Mapping against reference genome with the mapper module
 2. Known and novel miRNA discovery with the mirdeep2 module
8. miRNA quality control ([mirtrace](#))
9. Present QC for raw read, alignment, and expression results ([MultiQC](#))

