

1. Introduction	<ul style="list-style-type: none"> - Objectives - Computing requirements - Data 	<ul style="list-style-type: none"> ❖ Linux ❖ R ❖ wget
2. Data Pre-processing	<ul style="list-style-type: none"> - Merge technical replicate - Quality Control 	<ul style="list-style-type: none"> ❖ FastQC
3. Alignment	<ul style="list-style-type: none"> - Bowtie2 alignment - Duplicates - Fragment size 	<ul style="list-style-type: none"> ❖ Bowtie2 ❖ Samtools ❖ Picard
4. Filtering and Conversion	<ul style="list-style-type: none"> - Sam to bam - Bam to bed 	<ul style="list-style-type: none"> ❖ Samtools ❖ Bedtools
5. Spike-in Calibration	<ul style="list-style-type: none"> - Spike-in alignment - Scaling factor - ChIPseqSpikeInFree 	<ul style="list-style-type: none"> ❖ Bowtie2 ❖ Bedtools ❖ R: ChIPseqSpikeInFree
6. Peak Calling	<ul style="list-style-type: none"> - SEACR peak calling - MACS2 peak calling - % fragment in peaks 	<ul style="list-style-type: none"> ❖ SEACR ❖ MACS2 ❖ R: chromVAR
7. Visualization	<ul style="list-style-type: none"> - Genome browser - Heatmap 	<ul style="list-style-type: none"> ❖ Integrative Genome Viewer ❖ UCSC Genome Browser ❖ Deeptools
8. Differential Analysis	<ul style="list-style-type: none"> - DESeq2 - Peak x Sample matrix - Normalization - Differential detection 	<ul style="list-style-type: none"> ❖ R: DESeq2 ❖ R: GenomicRanges