## Comparative analysis of Fastqc using Nextflow vs Bash script

1. Bash script

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{ CODE BLOCK
#!/bin/bash
# Define input and output directories
INPUT_DIR="/home/Desktop/Prcatise/Nexflow_prac/my_pipelines/fastq_files"
OUTPUT DIR="/home/Desktop/Prcatise/Nexflow prac/my pipelines/fastq files
/output_bash_fastqc"
# Create the output directory if it doesn't exist
mkdir -p "$OUTPUT DIR"
# Loop through all .fastq.gz files in the input directory
for file in "$INPUT DIR"/*.fastq.gz
do
      echo "Running FastQC on $file..."
      fastqc "$file" --outdir "$OUTPUT_DIR"
done
echo "All FastQC analyses completed!"
Approx 55% complete for SRR33317478.fastq.gz
```

```
Approx 55% complete for SRR33317478.fastq.gz
Approx 60% complete for SRR33317478.fastq.gz
Approx 65% complete for SRR33317478.fastq.gz
Approx 70% complete for SRR33317478.fastq.gz
Approx 75% complete for SRR33317478.fastq.gz
Approx 80% complete for SRR33317478.fastq.gz
Approx 85% complete for SRR33317478.fastq.gz
Approx 90% complete for SRR33317478.fastq.gz
Approx 95% complete for SRR33317478.fastq.gz
Approx 95% complete for SRR33317478.fastq.gz
Analysis complete for SRR33317478.fastq.gz
All FastQC analyses completed!
[ble: elapsed 42.864s (CPU 117.5%)] bash fastqc_bash.sh
```

Bash takes 42 seconds to complete analysis

## 2. Nextflow script [CODE BLOCK #!/usr/bin/env nextflow params.reads = '/home/Desktop/Prcatise/Nexflow\_prac/my\_pipelines/fastq\_files/\*.fastq.gz' params.outdir = '/home/Desktop/Prcatise/Nexflow\_prac/my\_pipelines/fastq\_files/output\_fastqc' process FastQC { publishDir params.outdir, mode: 'copy' input: path read\_file output: path "\*.html" path "\*.zip" script: ..... fastqc \$read file } workflow { Channel .fromPath(params.reads) .set { read\_files } FastQC(read\_files)

Nexflow take only 21 seconds to finish task

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