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#!/bin/bash

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# COPYRIGHT="Copyright (C) 2016-2018 $AUTHOR"
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# VERSION="2018.04.03.1800"
# USAGE="bash Script6.sh $INF_MAPPED_R1 $INF_MAPPED_R2 $INF_SAM $LOG"

#####
# SUPPLEMENTARY FILE 6
# Bash script to automatically generate assembly quality statistics
# (i.e., the mean read length of mapped reads and the number of
# nucleotides with coverage depth equal or greater than 20, 50 and
# 100).
#####

# Check if sufficient commandline parameters
numArgmts=$#
if [ ! $numArgmts -eq 4 ]; then
    echo "ERROR | Incorrect number of commandline parameters" >&2
    exit 1
fi

# Check if input files exist
for v in "$@"; do
    if [ ! -f "$v" ]; then
        echo "ERROR | File not found: $v" >&2
        exit 1
    fi
done

# Check if dependencies exist
DEPS=(bowtie2 samtools bedtools)
for d in "${DEPS[@]"; do
    if [ ! -x "$(command -v $d)" ]; then
        echo "Error: $d is not installed" >&2
        exit 1
    fi
done

# Assigning commandline arguments
INF_MAPPED_R1=$1
INF_MAPPED_R2=$2
INF_SAM=$3
LOG=$4

#####

## Calculating mean read length

# Defining outfiles
OUF1=${INF_MAPPED_R1%.fastq*}_readLen.txt
OUF2=${INF_MAPPED_R2%.fastq*}_readLen.txt

# Calculate read lengths of R1 and R2
grep --no-group-separator -A1 "^@M" $INF_MAPPED_R1 | grep -v "^@M" | awk '{print len
gth($0)}' > $OUF1
grep --no-group-separator -A1 "^@M" $INF_MAPPED_R2 | grep -v "^@M" | awk '{print len
gth($0)}' > $OUF2

# Logging results
echo -e "\n# MEAN READ LENGTH (R1 PLUS R2)" >> $LOG
paste -d '\t' $OUF1 $OUF2 | awk '{ total += $1+$2 } END { print total/NR }' >> $LOG

#####

## Calculating N of bases with coverage depth equal or greater than to x, where x={2
0,50,100}

# Defining temporary files and outfiles
INF_BAM=${INF_SAM%.sam*}.bam
TMP_SRTD=${INF_BAM%.bam*}.sorted.bam
TMP_GNM=${INF_BAM%.bam*}.genome
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OUF3=${INF_BAM%.bam*}.genomecov

# Generating genome dummy file
touch $TMP_GNM
REF_LEN=$(head $INF_SAM | grep "^@SQ" | awk -F'LN:' '{print $2}')
echo -e "plastid\t$REF_LEN" >> $TMP_GNM

# Convert SAM to BAM
samtools view -bS $INF_SAM > $INF_BAM

# Sorting BAM file by position
samtools sort $INF_BAM > $TMP_SRTD

# Report per-base genome coverage
bedtools genomecov -ibam $TMP_SRTD -g $TMP_GNM -d > $OUF3

# Logging results
echo -e "\n# COVERAGE DEPTH STATISTICS" >> $LOG

echo -ne "Genome size (bp): " >> $LOG
GNM_SIZE=$(awk '{print $2}' $TMP_GNM)
LC_ALL=C printf "%d\n" $GNM_SIZE >> $LOG

echo -ne "N of bases with coverage depth greater than 20-fold: " >> $LOG
COV_DEP_20=$(awk '$3>20' $OUF3 | wc -l)
LC_ALL=C printf "%d\n" $COV_DEP_20 >> $LOG
echo -ne "P of bases with coverage depth greater than 20-fold: " >> $LOG
LC_ALL=C printf "%.2f\n" "$(echo "scale=4; ($COV_DEP_20/$GNM_SIZE)*100" | bc)" >> $LOG

echo -ne "N of bases with coverage depth greater than 50-fold: " >> $LOG
COV_DEP_50=$(awk '$3>50' $OUF3 | wc -l)
LC_ALL=C printf "%d\n" $COV_DEP_50 >> $LOG
echo -ne "P of bases with coverage depth greater than 50-fold: " >> $LOG
LC_ALL=C printf "%.2f\n" "$(echo "scale=4; ($COV_DEP_50/$GNM_SIZE)*100" | bc)" >> $LOG

echo -ne "N of bases with coverage depth greater than 100-fold: " >> $LOG
COV_DEP_100=$(awk '$3>100' $OUF3 | wc -l)
LC_ALL=C printf "%d\n" $COV_DEP_100 >> $LOG
echo -ne "P of bases with coverage depth greater than 100-fold: " >> $LOG
LC_ALL=C printf "%.2f\n" "$(echo "scale=4; ($COV_DEP_100/$GNM_SIZE)*100" | bc)" >> $LOG

# File hygiene
rm $TMP_SRTD
rm $TMP_GNM

#EOF

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