

software	purpose	command
picard	regenerate fastq files from BAM file aligned to hg18	java -d64 -Xmx4g -jar SamToFastq.jar I=\$pfx.bam F=\$pfx.1.fastq F2=\$pfx.2.fastq 2>&1
bwa	align fastq files to hg19	bwa aln -q 30 -t 8 \$hgReference \$fastq > \$fastq.aln.sai
bwa, samtools	convert aligned fastq files into new BAM file	bwa sampe -a 600 -P -r "\$RG" \$hgReference \$fastq1.aln.sai \$fastq2.aln.sai \$fastq1 \$fastq2 samtools view -bSh -o \$outprefix.bam -
samtools	sort and index new BAM file	samtools sort -@ 16 \$outprefix.bam \$outprefix.sorted 2, samtools index \$outprefix.sorted.bam 2
samtools	remove duplicate reads from BAM files	samtools rmdup ../\$tumorpfx/\$tumorpfx.out.sorted.bam \$tumorpfx.dedup.bam
GATK	indel realignment	java -d64 -jar \$gatkJar -R \$hgReference -T IndelRealigner -rf BadCigar -I \$tumorpfx.dedup.bam -known \$G1000.Mills -known \$G1000.Phase1.Indels -targetIntervals \$tumorpfx.intervals -o \$tumorpfx.realn.bam
GATK	base recalibration	java -d64 -jar \$gatkJar -nct 8 -T BaseRecalibrator -rf BadCigar -I \$tumorpfx.realn.bam -R \$hgReference -knownSites \$dbSNP -o \$tumorpfx.recal.grp
samtools	index recalibrated BAM file	samtools index \$tumorpfx.realn.recal.bam
SomaticSniper	call somatic mutations, generate VCF	bam-somaticsniper -q 40 -Q 40 -J -s 0.001 -F vcf -f \$hgReference \$tumorbam \$normalbam \$tumorpfx.SS.vcf