

19 WGS datasets
~30x mean coverage, 7×10^9 reads

fastq

bwatrim

read quality trimmer
BaseQ ≥ 15 , min length 35bp or toss

fastq

BWA

NGS read read alignment
(aln and sampe)

BAM

SAMtools

sort, remove duplicates, calculate BAQ,
and index
(sort, rmdp, calmd, and index)

BAM

GATK

Call variants and annotate
(UnifiedGenotyper and
VariantAnnotator)
maxAltAlleles 5,
BAQ re-calculated as necessary,
BAQ gap-open 30,
minCallConf 50,
minStandConf 20,
min BaseQ 17

VCF ~24M variants (raw, unfiltered)

Variant filtering

multiple tools
Criteria:
HWE, strand bias, call quality,
genotype quality, depth, repeat

VCF