

BV-BRC Test Report

A13. Service – Variation Analysis - Bacteria

Item to test	Variation Analysis Service using bacterial read files and SRA accessions
URL	https://www.bv-brc.org/app/Variation
Prerequisites	Bacterial Fasta contig files in Workspace
References	https://www.bv-brc.org/docs/quick_references/services/variation_analysis_service.html https://www.bv-brc.org/docs/tutorial/variation_analysis/variation_analysis.html
Tester(s)	Rebecca Wattam, Maulik Shukla
Test date	08-May-2022 (follow-up from original test)
Test result	Passed

Overview

- Test the Variation Analysis Service using exemplar bacterial datasets.
- Test input options, i.e., read files in FASTQ format and SRA accessions.
- Test different alignment strategies, i.e. BWA-mem, BWA-mem-strict, Bowtie2 and LAST. Test different SNP callers, such as FreeBayes and SAMtools.
- For each job submitted, verify successful completion of the job, presence of output files, and the quality of results.
- Review SNPs and alignment files in interactive the genome browser.

Test Data

Dataset	Rational	Input Format	Input
Escherichia coli genomes	Workshop dataset	Fastq files and SRA accessions	SRR396518.fastq.gz SRR396519.fastq.gz SRR396520.fastq.gz

- All test datasets and corresponding job results are available in the following public workspace:
<https://www.bv-brc.org/workspace/BVBRC@patricbrc.org/BVBRC%20Tests/Variation%20Analysis>

Test Results

- All variation analysis jobs completed successfully, without any errors.
- All jobs resulted in expected output files in corresponding job output directory, including BAM files, VCF files, and variation report.
- The variation report was informative and provided list of SNPs and indels as an HTML table. The table shown SNPs, their position, corresponding gene, NA and AA position, NA/AA change, and annotation of SNP effect and its significance.

- The link to genome browser worked as expected. It loaded all BAM and VCF files as separate tracks and allowed zooming in to a single SNP level. Clicking on a SNP provided all the details and annotation about that SNP.
- All test datasets and corresponding job results are available in the following public workspace: <https://www.bv-brc.org/workspace/BVBRC@patricbrc.org/BVBRC%20Tests/Variation%20Analysis>
- Below are a series of screenshots showing successful completion of the jobs, availability of the result files in the workspace, the variation table and genome browser views.

Paired read library ⓘ

READ FILE 1
↓ ↕

READ FILE 2
↓ ↕

Single read library ⓘ

READ FILE
↓ ↕ SRR396520.fastq.gz

SRA run accession ⓘ

SRR ACCESSION
SRR396520

Parameters ⓘ

ALIGNER
BWA-mem

SNP CALLER
FreeBayes

TARGET GENOME
Escherichia coli str. K-12 substr. MG1655

OUTPUT FOLDER
↓ ↕ Variation Analysis

OUTPUT NAME
Ecoli variation - SRA accessions

Selected libraries ⓘ

Place read files here using the arrow buttons.

SRR396520	i x
SRR396519	i x
SRR396518	i x

Your job has been submitted successfully. Please visit your [Jobs List](#) to check the status of your job and access the results.

Status	ID	Service	Output Name	Submit	Start	Completed
completed	7747607	Variation Analysis	Ecoli variation - fastq files	5/8/22, 11:35 AM	5/8/22, 11:35 AM	5/8/22, 12:03 PM
completed	7747608	Variation Analysis	Ecoli variation - SRA accessions	5/8/22, 11:36 AM	5/8/22, 11:36 AM	5/8/22, 12:00 PM
completed	7747609	Variation Analysis	Ecoli variation - bwa-mem-strict	5/8/22, 11:37 AM	5/8/22, 11:37 AM	5/8/22, 11:49 AM
completed	7747610	Variation Analysis	Ecoli variation - bowtie2	5/8/22, 11:37 AM	5/8/22, 11:37 AM	5/8/22, 11:53 AM
completed	7747615	Variation Analysis	Ecoli variation - last	5/8/22, 11:40 AM	5/8/22, 11:40 AM	5/8/22, 12:17 PM
completed	7747616	Variation Analysis	Ecoli variation - samtools	5/8/22, 11:40 AM	5/8/22, 11:40 AM	5/8/22, 11:57 AM

BVBRC / BVBRC Tests / Variation Analysis (9 items)

UPL
 ADD FOLDER
 SHOW HIDDEN

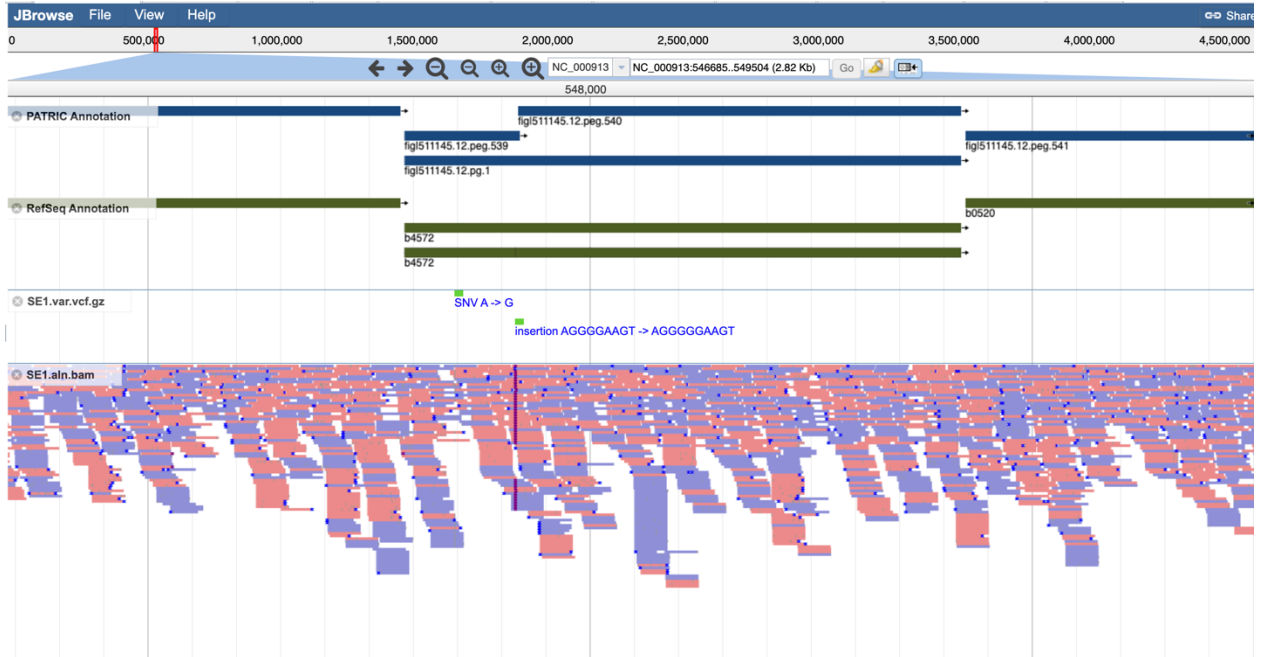
Name	Size	Owner	Members	Created
Parent folder	-	-	-	-
SRR396520.fastq.gz	228.6 MB	me	Public	5/8/22, 11:33 AM
SRR396519.fastq.gz	276.7 MB	me	Public	5/8/22, 11:33 AM
SRR396518.fastq.gz	248.2 MB	me	Public	5/8/22, 11:33 AM
Ecoli variation - bwa-mem-strict	10.1 kB	me	Public	5/8/22, 11:49 AM
Ecoli variation - bowtie2	9.9 kB	me	Public	5/8/22, 11:53 AM
Ecoli variation - samtools	9.9 kB	me	Public	5/8/22, 11:57 AM
Ecoli variation - SRA accessions	10.1 kB	me	Public	5/8/22, 12:00 PM
Ecoli variation - fastq files	10.3 kB	me	Public	5/8/22, 12:03 PM
Ecoli variation - last	9.8 kB	me	Public	5/8/22, 12:17 PM

Name	Size	Owner	Members	Created
Parent folder	-	-	-	-
SE1.aln.bam	191.8 MB	me	Public	5/8/22, 12:00 PM
SE1.aln.bam.bai	14.2 kB	me	Public	5/8/22, 12:00 PM
SE1.consensus.fa	4.7 MB	me	Public	5/8/22, 12:00 PM
SE1.var.annotated.tsv	3.0 kB	me	Public	5/8/22, 12:00 PM
SE1.var.snpEff.vcf	15.2 kB	me	Public	5/8/22, 12:00 PM
SE1.var.vcf	12.9 kB	me	Public	5/8/22, 12:00 PM
SE1.var.vcf.gz	3.4 kB	me	Public	5/8/22, 12:00 PM
SE1.var.vcf.gz.tbi	269 B	me	Public	5/8/22, 12:00 PM
SE2.aln.bam	209.2 MB	me	Public	5/8/22, 12:00 PM
SE2.aln.bam.bai	14.3 kB	me	Public	5/8/22, 12:00 PM
SE2.consensus.fa	4.7 MB	me	Public	5/8/22, 12:00 PM
SE2.var.annotated.tsv	5.8 kB	me	Public	5/8/22, 12:00 PM
SE2.var.snpEff.vcf	22.9 kB	me	Public	5/8/22, 12:00 PM
SE2.var.vcf	19.3 kB	me	Public	5/8/22, 12:00 PM
SE2.var.vcf.gz	4.6 kB	me	Public	5/8/22, 12:00 PM
SE2.var.vcf.gz.tbi	329 B	me	Public	5/8/22, 12:00 PM
SE3.aln.bam	178.2 MB	me	Public	5/8/22, 12:00 PM
SE3.aln.bam.bai	14.1 kB	me	Public	5/8/22, 12:00 PM
SE3.consensus.fa	4.7 MB	me	Public	5/8/22, 12:00 PM

BVBRC / BVBRC Tests / Variation Analysis / Ecoli variation - SRA accessions / all.var.html

html file: all.var.html

Samples	Contig	Pos	Ref	Var	Score	Var_cov	Var_frac	Type	Ref_nt	Var_nt	Ref_nt_pos_change	Ref_aa_pos_change	Frameshift	Gene_ID	Locus_tag	Gene_name
1:SE2	NC_000913	1049100	C	T	11.4092	2.0	0.09									
1:SE2	NC_000913	1049669	A	G	49.2323	3.0	0.13	Synon	gga	ggG	294A>G	Gly98Gly		fgl511145.12.peg.1024	b0988	insB
1:SE2	NC_000913	1049673	A	G	51.2844	2.0	0.10	Nonsyn	aag	Gag	298A>G	Lys100Glu		fgl511145.12.peg.1024	b0988	insB
1:SE2	NC_000913	1049680	T	C	39.3747	2.0	0.11	Nonsyn	ctg	cCg	305T>C	Leu102Pro		fgl511145.12.peg.1024	b0988	insB
1:SE2	NC_000913	1049682	T	G	43.4713	2.0	0.12	Nonsyn	tcg	Gcg	307T>G	Ser103Ala		fgl511145.12.peg.1024	b0988	insB
1:SE2	NC_000913	1049685	T	G	42.3477	2.0	0.17	Nonsyn	ttc	Gtc	310T>G	Phe104Val		fgl511145.12.peg.1024	b0988	insB
1:SE3	NC_000913	1568836	C	G	57.3728	2.0	0.13	Nonsyn	ggc	Cgc	1234G>C	Gly412Arg		fgl511145.12.peg.1560	b1493	gadB
1:SE1	NC_000913	1568846	G	A	19.4249	2.0	0.09	Synon	gcc	gcT	1224C>T	Ala408Ala		fgl511145.12.peg.1560	b1493	gadB
1:SE3	NC_000913	1617114	TGCAACTAATTACTTGCCAGGG	TG	528.949	24.0	0.96	Deletion								
2:SE1,SE2	NC_000913	1617423	C	T	1011.0	50.5	0.96	Nonsyn	cgc	Tgc	280C>T	Arg94Cys		fgl511145.12.peg.1599	b1530	marR
1:SE3	NC_000913	1633932	C	A	58.363	2.0	0.12	Nonsyn	gat	Tat	460G>T	Asp154Tyr		fgl511145.12.peg.1618		
1:SE3	NC_000913	1718907	GCAATTGATTGCTAAAC	GC	341.821	16.0	0.94	Deletion								
3:SE1,SE2,SE3	NC_000913	1903785	G	A	1667.44	72.3	0.99	Nonsyn	ggt	gAt	74G>A	Gly25Asp		fgl511145.12.peg.1898	b1821	yebN



Primary Data

Type	SNV
Score	562.275
Description	SNV A -> G
Position	NC_000913:547694..547694
Length	1 bp

Attributes

AB	0
ABP	0
AC	1
AF	1
AN	1
AQ	28
CIGAR	1X

SRF	0
SRP	0
SRR	0
alternative_alleles	G
description	SNV A -> G
reference_allele	A
seq_id	NC_000913

Genotypes (1)

variant	1	100%
homozygous	1	100%
G variant	1	100%
Total	1	100%

Name	GT	DP	AD	RO	QR	AO	QA	GL
unknown	G	28	0 28	0	0	28	679	-61.336 0

References

- [Variation Analysis Service Quick Reference Guide](#)
- [Variation Analysis Service Tutorial](#)