# Analyses

Bioviz Connect utilizes CyVerse's cloud computing to perform powerful analyses on genomic data.

### To Analyze:

To analyze a file right click and choose 'Analyse'. A panel will open on the right with the available analyses, if there are any. Click the appropriate button for the analysis you want to run.

Analysis ×	Analysis ×
SRR1171888.bam	fastal fas
Make scaled coverage graph	Jasanjas
Make a scaled coverage graph file in bigWig format from a bam sequence alignment file using deepTools bamCoverage version 3.3.0 with options normalizeUsing CPMbinSize 1. See	Find enriched sequence motifs
deeptools.readthedocs.io. Make coverage graph	Find over-represented sequences in a fasta file using DREME from meme-suite.org. DREME discovers short. ungapped motifs that are relatively
Wake a coverage graph file in bigWig format from a bam sequence alignment file using deepTools bamCoverage version 3.3.0 with optionsbinSize	enriched in your sequences compared with shuffled sequences.

Once you have chosen a type of analysis, a new form will appear in the same window. Enter your analysis name. An output file name will be generated for you. This can be edited to any name you like.

When you have completed entering your information click 'Run Analysis'.

Make scaled coverage graph	🗲 Bacl
Analysis Name 🔞	
$C_{sativa} Scaled Coverage Graph$	
Input File (BAM) 🔞	
SRR1171888.bam	
Input File Index (.bai) 🕑	
SRR1171888.bam.bai	
Output file name 😧	
SRR1171888.bigwig	
Run Analysis	

# Analysis Status

You can check the analyses' status by using the 'Analyses History' tab, where analyses are listed as Queued (waiting to run), Running, Failed, or Completed. The length of the time to complete a job is dependent on the size of the queue, the analysis being carried out, and the size of the file. It will likely take several minutes for the analysis to complete. CyVerse will send an email to you when the analysis is finished.

+ New>	Name	App Name	Start Date	End Date	Status	
	SRR10060912ScaledCoverag	Make scaled coverage graph	06-17-2020 14:57:21	06-17-2020 15:01:16	Completed	
🖀 Home	SRR10060911ScaledCoverag	Make scaled coverage graph	06-17-2020 14:35:39	06-17-2020 14:40:37	Completed	
Less Shared with me	TestWednesdayScaled	Make scaled coverage graph	06-17-2020 14:34:36	06-17-2020 14:39:43	Completed	
🖀 Community	TestWednesday	Make coverage graph	06-17-2020 14:33:20	06-17-2020 14:38:58	Completed	
<ol> <li>Analysis History</li> </ol>	Page 1					

#### Analysis Results

When the analysis is finished, any files or folders created through the process will appear in the 'Analyses' folder in your 'Home' directory, or in the same location as the input files if those are stored in a location where you have permission to modify or add to the folder.

BioViz Connect		Search Home	+ News	🕷 / analyses 🛛 🖗			
			😤 Home	Visualization Tools	Name ^	Size ^	Last Modified A
+ New>			44 Shared with me	View in IG8	SRR10060911ScaledCoverageGraph.bedgraph	337 140	Jun 17, 2020
🛠 Home	Visualization Tools	Name ^	de Community	View in IG8	SRR100609125caledCoverageGraph.bedgraph	135 MB	Jun 17, 2020
L Shared with me		<ul> <li>analyses</li> </ul>	Analysis History	View in IG8	C TestWednesday.bedgraph	287.548	Jun 17, 2028
🖶 Community		LocalQuickload		View in IG8	D TestWednesdayScaled.bedgraph	348.548	Jun 17, 2020
Analysis History		<ul> <li>my_quickload</li> </ul>		Page 1			
		🗅 bam.txt					
	Page 1						

To quickly navigate to the analysis results, you can click the analysis name in the 'Analyses History' tab, opening the folder where the output data files are stored.

Note that running analyses uses Compute Units which may be limited by your account's subscription status. See cyverse.org/ for more information.

## Supported File Types

Currently BioViz Connect supports analyses of:

- bam files
  - genome wide coverage graph
    - Make a coverage graph file in bigWig format from a bam sequence alignment file using deepTools bamCoverage version 3.3.0 with options --binSize 1. See deeptools.readthedocs.io.
  - genome wide scaled coverage graph
    - Make a scaled coverage graph file in bigWig format from a bam sequence alignment file using deepTools bamCoverage version 3.3.0 with options --normalizeUsing CPM --binSize 1. See deeptools.readthedocs.io.
- fasta files
  - Find enriched sequence motifs
    - Find overrepresented sequences in a fasta file using DREME from meme-suite.org. DREME discovers short, ungapped motifs that are relatively enriched in your sequences compared with shuffled sequences.