

New Tools, New Datasources

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Overview

- Tools
- Datasources
- External Display Applications

The Difference?

- Tools
 - Any command-line executable
- Datasources
 - Getting Data from an external source
- External Display Applications
 - Sending Data to an external web application

Overview

- Tools
- Datasources
- External Display Applications

Recent Additions

[NCBI BLAST+](#)
[NGS: QC and manipulation](#)
[NGS: Picard \(beta\)](#)
[NGS: Mapping](#)
[NGS: Indel Analysis](#)
[NGS: RNA Analysis](#)
[NGS: SAM Tools](#)
[NGS: GATK Tools](#)
[NGS: Peak Calling](#)
[NGS: Simulation](#)
[SNP/WGA: Data; Filters](#)
[SNP/WGA: QC; LD; Plots](#)
[SNP/WGA: Statistical Models](#)
[Human Genome Variation](#)
[VCF Tools](#)

- Status: Stable (pretty well tested) --> Alpha

Recent Additions

NGS: RNA Analysis

RNA-SEQ

- [Tophat](#) Find splice junctions using RNA-seq data
- [Cufflinks](#) transcript assembly and FPKM (RPKM) estimates for RNA-Seq data
- [Cuffcompare](#) compare assembled transcripts to a reference annotation and track Cufflinks transcripts across multiple experiments
- [Cuffdiff](#) find significant changes in transcript expression, splicing, and promoter use

FILTERING

- [Filter Combined Transcripts](#) using tracking file

VCF Tools

- [Intersect](#) Generate the intersection of two VCF files
- [Annotate](#) a VCF file (dbSNP, hapmap)
- [Filter](#) a VCF file
- [Extract](#) reads from a specified region

NGS: Picard (beta)

QC/METRICS FOR SAM/BAM

- [BAM Index Statistics](#)
- [Sam/bam Alignment Summary Metrics](#)
- [Sam/bam GC Bias Metrics](#)
- [Estimate Library Complexity](#)
- [Insertion size metrics](#) for PAIRED data
- [Sam/bam Hybrid Selection Metrics](#) For (eg exome) targeted data

BAM/SAM CLEANING

- [Add or Replace Groups](#)
- [Reorder SAM](#)
- [Replace Sam Header](#)
- [Paired Read Mate Fixer](#) for paired data
- [Mark Duplicate reads](#)

FASTQC: FASTQ/SAM/BAM

- [Fastqc](#): Fastqc QC using FastQC from Babraham

NGS: GATK Tools

Alpha

REALIGNMENT

- [Realigner Target Creator](#) for use in local realignment
- [Indel Realigner](#) – perform local realignment

BASE RECALIBRATION

- [Count Covariates](#) on BAM files
- [Table Recalibration](#) on BAM files
- [Analyze Covariates](#) – perform local realignment

GENOTYPING

- [Unified Genotyper](#) SNP and indel caller

Report Bugs, please

Adding your Own

- Write or download a command-line executable
- Determine number and kind of
 - Input and Output Datasets
 - Input Parameters
- Construct a descriptive tool configuration XML file
 - Write a wrapper script, only if required

Tool Configuration

- Tool Action - Default tool action should be adequate (Upload tool uses custom tool action)
- Tool Command
- Inputs
 - Action - Used by datasource tools
 - Parameters
- Outputs
- Help
- Tests

A Basic Tool

```
<tool id="fa_gc_content_1" name="Compute GC content">
  <description>for each sequence in a file</description>
  <command interpreter="perl">toolExample.pl $input $output</command>
  <inputs>
    <param format="fasta" name="input" type="data" label="Source file">
  </inputs>
  <outputs>
    <data format="tabular" name="output" />
  </outputs>

  <tests>
    <test>
      <param name="input" value="fa_gc_content_input.fa"/>
      <output name="out_file1" file="fa_gc_content_output.txt"/>
    </test>
  </tests>

  <help>
This tool computes GC content from a FASTA file.
  </help>
</tool>
```

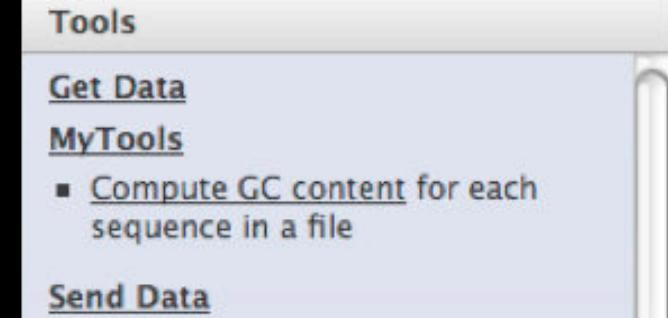
Compute GC content

Source file:

1: Uploaded FASTA File

Execute

This tool computes GC content from a FASTA file.



```
<section name="MyTools" id="mTools">
  <tool file="myTools/toolExample.xml" />
</section>
```

tool_conf.xml

Cluster

Cluster intervals of: 6: UCSC Main on Human: knownGene

max distance between intervals: 1 (bp)

min number of intervals per cluster: 2

Return type: Merge clusters into single intervals

TIP: If your query does not appear in the pulldown menu -> it is not in interval format. Use "edit attributes" to set chromosome, start, end, and strand columns

Screencasts!

See Galaxy Interval Operation [Screencasts](#) (right click to open this link in another window).

Syntax

- **Maximum distance** is greatest distance in base pairs allowed between intervals that will be considered "clustered". **Negative** values for distance are allowed, and are useful for clustering intervals that overlap.
- **Minimum intervals per cluster** allow a threshold to be set on the minimum number of intervals to be considered a cluster. Any area with less than this minimum will not be included in the output.
- **Merge clusters into single intervals** outputs intervals that span the entire cluster.
- **Find cluster intervals; preserve comments and order** filters out non-cluster intervals while maintaining the original ordering and comments in the file.
- **Find cluster intervals; output grouped by clusters** filters out non-cluster intervals, but outputs the cluster intervals so that they are grouped together. Comments and original ordering in the file are lost.

Example



```
cluster.xml
1 <tool id="gops_cluster_1" name="Cluster">
2   <description>[[Cluster]] the intervals of a query</description>
3   <command interpreter="python2.4">
4     gops_cluster.py $input1 $output -1 $input1_chromCol,$input1_startC
5       -d $distance -m $minregions -o $returntype
6   </command>
7   <inputs>
8     <param format="interval" name="input1" type="data">
9       <label>Cluster intervals of</label>
10      </param>
11      <param name="distance" size="5" type="integer" value="1" help="(bp)">
12        <label>max distance between intervals</label>
13      </param>
14      <param name="minregions" size="5" type="integer" value="2">
15        <label>min number of intervals per cluster</label>
16      </param>
17      <param name="returntype" type="select" label="Return type">
18        <option value="1">Merge clusters into single intervals</option>
19        <option value="2">Find cluster intervals; preserve comments and
20          order</option>
21        <option value="3">Find cluster intervals; output grouped by clus
22        <option value="4">Find the smallest interval in each cluster</op
23        <option value="5">Find the largest interval in each cluster</opt
24      </param>
25    </inputs>
26    <help>
27      ... class:: infomark ...
28
29      **TIP:** If your query does not appear in the pulldown menu -> it is n
30
31      -----
32
33      **Screencasts!**
34
35      See Galaxy Interval Operation Screencasts (right click to open this l
36
37      ... _Screencasts: http://www.bx.psu.edu/cgi-bin/trac.cgi/wiki/GopsDesc
38
39      -----
40
41      **Syntax**
42
43      - **Maximum distance** is greatest distance in base pairs allowed betw
44      - **Minimum intervals per cluster** allow a threshold to be set on the
45      - **Merge clusters into single intervals** outputs intervals that span
46      - **Find cluster intervals; preserve comments and order** filters out
47      - **Find cluster intervals; output grouped by clusters** filters out n
```

Input Parameter types

Basic

- Text
- Integer
- Float
- Select
 - Static
 - Dynamic
- Boolean
- Genome build
- Data column
- Data
- Hidden
- Base URL
- File
- Drill down
- Grouping
- Conditional
- Repeat
- Config Files

Configfile Inputs

- Temporary file is available as input to the tool
- Cheetah templating language is used to define the contents of a temporary file
- Examples
 - Formatted list of input values
 - R (statistics package) scripts
 - Gnuplot Commands
 - Dynamically generated python scripts

Advanced Topics

- Datasets and Datatypes
- Metadata
- Dynamic Select lists
- Tool Output actions
 - changing formats, setting metadata based upon user inputs
- Filtering outputs
- Parameter Validation

Datasets and Datatypes

- All datasets are associated with a Datatype
 - File format
 - Type of Data: genomic intervals, sequence, alignment
 - Hierarchical structure allows accepting more specific datatypes without conversion (tabular <- interval <- bed)
 - Datatype Converters allow use of non-subclassed datatypes as direct input for tools (MAF to FASTA converter allows selection of MAF file as input when FASTA is required)
 - Metadata
 - datatypes_conf.xml and lib/galaxy/datatypes

[https://bitbucket.org/galaxy/galaxy-central/wiki/
AddingDatatypes](https://bitbucket.org/galaxy/galaxy-central/wiki/AddingDatatypes)

Datatype Metadata

- Information that describes the contents of the Dataset
 - Line / Sequence Counts
 - Count and type of tabular columns
 - Column assignments: start, stop, strand
Genome build(s): dbkey, species
 - Files: indexes
- Used by tools to customize interface and results

Dynamic Select Lists

- Use species **metadata** value from MAF file to dynamically generate a list of checkboxes

The image shows a screenshot of a bioinformatics application interface. On the left, there is a panel titled "1: 5-way MAF" which displays metadata about a uploaded MAF file, including the number of blocks (9), format (maf), database (hg17), and info (uploaded maf file). Below this, a "Species" section lists "hg17 panTrol mm5 rn3 canFam1". A code snippet below the species list shows the MAF header and some sequence data. On the right, a modal dialog titled "Choose species:" is open, containing a "Select All" button, an "Unselect All" button, and five checkboxes corresponding to the species listed in the MAF file: hg17, panTrol, mm5, rn3, and canFam1. Below the checkboxes is the instruction "Select species to be included in the final alignment".

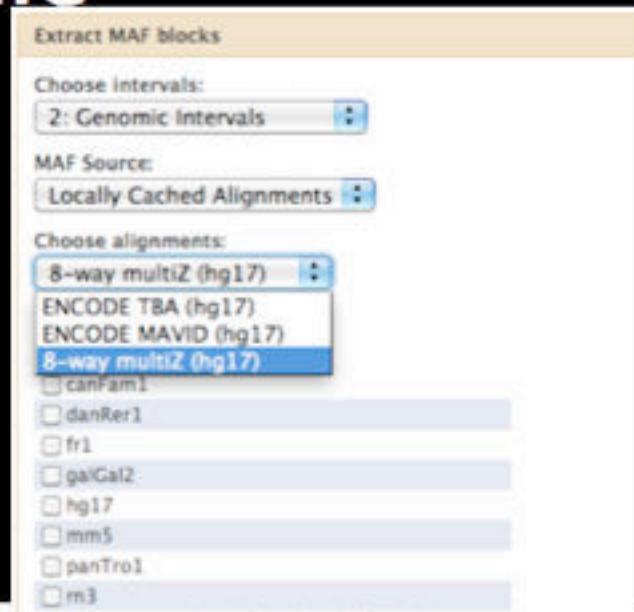
```
<param name="species" type="select" display="checkboxes" multiple="true" label="Choose species" help="Select s
<options>
  <filter type="data_meta" ref="mafFile" key="species" />
</options>
</param>
```

List of available alignments should be filtered based upon the dbkey of the input interval file

- Locally cached MAFs
- Don't want to modify the tool each time a new MAF set is added to the cache
- Dynamic Select: Use a separate file to describe available cached MAFs

```
<param name="mafType" type="select" label="Choose alignments">
<options from_file="maf_index.loc">
  <column name="name" index="0"/>
  <column name="value" index="1"/>
  <column name="dbkey" index="2"/>
  <column name="species" index="3"/>
  <filter type="data_meta" ref="input1" key="dbkey" column="2" multiple="True" separator=","/>
  <validator type="no_options" message="No alignments are available for the build associated with the selected inter
</options>
</param>
```

```
$ cat maf_index.loc
#Display name UID
#Indexed for:build1,build2,build3      exists in maf:build1,build2,build3      Comma Separated Full Paths To Files
ENCODE TBA (hg17)          ENCODE_TBA_hg17_armadillo,baboon,galGal2,panTro1,colobus_monkey,cow,canFam1,dusky_titi,elephant,fri,galago,hedgehog,hg
ENCODE MAVID (hg17)        ENCODE_MAVID_hg17      armadillo,baboon,galGal2,panTro1,colobus_monkey,cow,canFam1,dusky_titi,elephant,fri,galago,he
ENCODE TBA (hg16)          ENCODE_TBA_hg16_armadillo,baboon,galGal2,panTro1,colobus_monkey,cow,canFam1,dusky_titi,elephant=elephant,fri,galago,he
8-way multiZ (hg17)        8_WAY_MULTI2_hg17      canFam1,danRer1,fri,galGal2,hg17,mm5,panTro1,m3      canFam1,danRer1,fri,galGal2,hg17,mm5,p
17-way multiZ (hg18)       17_WAY_MULTI2_hg18     hg18,panTro1,bosTau2,rheMac2,mm8,mm4,canFam2,echTel1,loxAfr1,oryCun1,danRer3,monDom4,dasNov1,g
3-way multiZ (hg18,panTro2,rheMac2)  3_WAY_MULTI2_hg18   hg18,panTro2,rheMac2      hg18,panTro2,rheMac2      /depot/data2/galaxy/hg18/align
```



See the Split MAF blocks by Species tool for more information.

Execute

Tool Output Actions

- Allow run-time attributes (parameter settings, input metadata, datatype) to change attributes of a created output file
- Examples:
 - Cut tool
 - Default output is tabular
 - What if input was BED and all required interval columns are kept?
 - NGS mappers (e.g. bowtie)
 - Set dbkey of output when using pre-built indices

Cut Tool

```
<outputs>
  <data format="tabular" name="out_file1" >
    <actions>
      <conditional name="delimiter">
        <when value="T">
          <conditional name="input">
            <when datatype_isinstance="interval">
              <action type="format" default="tabular">
                <option type="from_param" name="columnList" column="0" offset="0"> <!-- chromCol is 1-->
                  <filter type="insert_column" column="0" value="interval"/>

                  <filter type="insert_column" ref="columnList" /> <!-- startCol -->
                  <filter type="insert_column" ref="columnList" /> <!-- endCol -->

                  <filter type="multiple_splitter" column="1" separator=","/>
                  <filter type="column_strip" column="1"/> <!-- get rid of all external whitespace -->
                  <filter type="string_function" column="1" name="lower" />
                  <filter type="param_value" column="1" value="^c\d{1,} $" compare="re_search" keep="True"/>
                  <filter type="column_strip" column="1" strip="c"/> <!-- get rid of c's -->
                  <filter type="boolean" column="1" cast="int" />

                  <filter type="multiple_splitter" column="2" separator=","/>
                  <filter type="column_strip" column="2"/> <!-- get rid of all external whitespace -->
                  <filter type="string_function" column="2" name="lower" />
                  <filter type="param_value" column="2" value="^c\d{1,} $" compare="re_search" keep="True"/>
                  <filter type="column_strip" column="2" strip="c"/> <!-- get rid of c's -->
                  <filter type="boolean" column="2" cast="int" />

                  <filter type="multiple_splitter" column="3" separator=","/>
                  <filter type="column_strip" column="3"/> <!-- get rid of all external whitespace -->
                  <filter type="string_function" column="3" name="lower" />
                  <filter type="param_value" column="3" value="^c\d{1,} $" compare="re_search" keep="True"/>
                  <filter type="column_strip" column="3" strip="c"/> <!-- get rid of c's -->
                  <filter type="boolean" column="3" cast="int" />

                  <filter type="metadata_value" ref="input" name="chromCol" column="1" />
                  <filter type="metadata_value" ref="input" name="startCol" column="2" />
                  <filter type="metadata_value" ref="input" name="endCol" column="3" />
                </option>
              </action>
            </when>
          </conditional>
        </when>
      </conditional>
    </actions>
  </data>
</outputs>
```

```
<conditional name="out_file1">
  <when datatype_isinstance="interval">
    <action type="metadata" name="chromCol">
      <option type="from_param" name="columnList" column="0" offset="0"> <!-- chromCol is 0-->
        <filter type="multiple_splitter" column="0" separator=","/>
        <filter type="column_strip" column="0"/> <!-- get rid of all external whitespace -->
        <filter type="string_function" column="0" name="lower" />
        <filter type="param_value" column="0" value="^c\d{1,} $" compare="re_search" keep="True"/>
        <filter type="column_strip" column="0" strip="c"/> <!-- get rid of c's -->
        <filter type="insert_column" value="1" iterate="True" column="0"/>
        <filter type="boolean" column="1" cast="int" />
        <filter type="metadata_value" ref="input" name="chromCol" column="1" />
      </option>
    </action>

    <action type="metadata" name="startCol">
      <option type="from_param" name="columnList" column="0" offset="0"> <!-- startCol is 0-->
        <filter type="multiple_splitter" column="0" separator=","/>
        <filter type="column_strip" column="0"/> <!-- get rid of all external whitespace -->
        <filter type="string_function" column="0" name="lower" />
        <filter type="param_value" column="0" value="^c\d{1,} $" compare="re_search" keep="True"/>
        <filter type="column_strip" column="0" strip="c"/> <!-- get rid of c's -->
        <filter type="insert_column" value="1" iterate="True" column="0"/>
        <filter type="boolean" column="1" cast="int" />
        <filter type="metadata_value" ref="input" name="startCol" column="1" />
      </option>
    </action>

    <action type="metadata" name="endCol">
      <option type="from_param" name="columnList" column="0" offset="0"> <!-- endCol is 0-->
        <filter type="multiple_splitter" column="0" separator=","/>
        <filter type="column_strip" column="0"/> <!-- get rid of all external whitespace -->
        <filter type="string_function" column="0" name="lower" />
        <filter type="param_value" column="0" value="^c\d{1,} $" compare="re_search" keep="True"/>
        <filter type="column_strip" column="0" strip="c"/> <!-- get rid of c's -->
        <filter type="insert_column" value="1" iterate="True" column="0"/>
        <filter type="boolean" column="1" cast="int" />
        <filter type="metadata_value" ref="input" name="endCol" column="1" />
      </option>
    </action>

    <action type="metadata" name="nameCol" default="0">
      <option type="from_param" name="columnList" column="0" offset="0"> <!-- nameCol is 0-->
        <filter type="multiple_splitter" column="0" separator=","/>
        <filter type="column_strip" column="0"/> <!-- get rid of all external whitespace -->
        <filter type="string_function" column="0" name="lower" />
        <filter type="param_value" column="0" value="^c\d{1,} $" compare="re_search" keep="True"/>
        <filter type="column_strip" column="0" strip="c"/> <!-- get rid of c's -->
        <filter type="insert_column" value="1" iterate="True" column="0"/>
        <filter type="boolean" column="1" cast="int" />
        <filter type="metadata_value" ref="input" name="nameCol" column="1" />
      </option>
    </action>
```

Cut Tool

```
<action type="metadata" name="strandCol" default="0">
    <option type="from_param" name="columnList" column="0" offset="0"> <!-- strandCol is 0-->
        <filter type="multiple_splitter" column="0" separator=","/>
        <filter type="column_strip" column="0"/> <!-- get rid of all external whitespace -->
        <filter type="string_function" column="0" name="lower" />
        <filter type="param_value" column="0" value="^c\d{1,} $" compare="re_search" keep="True"/>
        <filter type="column_strip" column="0" strip="c"/> <!-- get rid of c's -->
        <filter type="insert_column" value="1" iterate="True" column="0"/>
        <filter type="boolean" column="1" cast="int" />
        <filter type="metadata_value" ref="input" name="strandCol" column="1" />
    </option>
</action>
</when>
</conditional>

</when>
</conditional>
</when>
</conditional>
</actions>
</data>
```

tools/filters/cutWrapper.xml

Filtering Outputs

- Parameter settings can be used to change the outputs that are created

```
<outputs>
  <data name="output_bed_file" format="bed" label="${tool.name} on ${on_string} (peaks: bed)"/>
  <data name="output.xls_to_interval_peaks_file" format="interval" label="${tool.name} on ${on_string}
(peaks: interval)">
    <filter>xls_to_interval is True</filter>
  </data>
  <data name="output.xls_to_interval_negative_peaks_file" format="interval" label="${tool.name} on $
{on_string} (negative peaks: interval)">
    <filter>xls_to_interval is True</filter>
    <filter>input_type['input_control_file1'] is not None</filter>
  </data>
  <data name="output_treatment_wig_file" format="wig" label="${tool.name} on ${on_string} (treatment:
wig)">
    <filter>wig_type['wig_type_selector']=='wig'</filter>
  </data>
  <data name="output_control_wig_file" format="wig" label="${tool.name} on ${on_string} (control: wig)">
    <filter>wig_type['wig_type_selector'] == 'wig'</filter>
    <filter>input_type['input_control_file1'] is not None</filter>
  </data>
  <data name="output_extra_files" format="html" label="${tool.name} on ${on_string} (html report)"/>
</outputs>
```

Parameter Validation

Choose intervals:

- 1: UCSC Main on Human (genome) ▾
- 1: UCSC Main on Human (genome)
- 2: (as interval) Extract MAF blocks on data 1

```
<param format="interval" name="input1" type="data" label="Choose intervals">↓  
  <validator type="unspecified_build" />↓  
</param>↓
```

Choose intervals:

11: Gene from unknown Species ▾

✖ Unspecified genome build, click the pencil icon in the history item to set the genome build

11: Gene from
unknown Species

1 regions, format: bed,
database: ?

Info: uploaded file
[save](#) | [rerun](#)

1. Chrom	2. Start	3. End	4. Name
chrX	152643515	152663410	NM_000



Available Validators

Validator	Use
expression	Evaluates a python expression using the parameter value
regex	Checks if parameter value matches the specified regular expression
in_range	Ensures a numerical parameter value falls between a min and max
length	Ensures the length of a text parameter value falls between a min and max
metadata	Checks if specified metadata is missing for a dataset
unspecified_build	Checks if the dbkey for a dataset has been set
no_options	Ensures at least one option has been set for select parameters
empty_field	Ensures a text field has not been left empty
dataset_metadata_in_file	Checks if a particular metadata value for a dataset exists in a File
dataset_ok	Ensures an input dataset is in the OK state

I created a tool...now what

- Share it!
- Galaxy Tool Shed
 - coming up next
 - Talk by Greg Von Kuster

Overview

- Tools
- Datasources
- External Display Applications

Recent Additions

Galaxy

Analyze Data Workflow Shared Data Visualization Admin Help User

Tools Options ▾

Get Data

- Upload File from your computer
- UCSC Main table browser
- UCSC Test table browser
- UCSC Archaea table browser
- BX main browser
- Get Microbial Data
- BioMart Central server
- BioMart Test server
- CBI Rice Mart rice mart
- GrameneMart Central server
- modENCODE fly server
- Flymine server
- Flymine test server
- modENCODE modMine server
- Ratmine server
- YeastMine server
- metabolicMine server
- modENCODE worm server
- Wormbase server
- Wormbase test server
- EuPathDB server
- EncodeDB at NHGRI
- EpiGRAPH server
- EpiGRAPH test server
- HbVar Human Hemoglobin Variants and Thalassemia

SGD YeastMine v 2011-05-22 Saccharomyces Genome Database provides *S. cerevisiae* data powered by InterMine 0.96

Home Templates Lists QueryBuilder Tools Data API MyMine Search: e.g. act1 Take a tour

Welcome Back!

YeastMine provides *S.cerev*. You can run flexible querie results and analyse lists of how YeastMine works (using an example).

Search

Search YeastMine. Enter names, identifiers or keywords for genes, proteins, ontology terms, authors, abstract etc. (e.g. rad54, Act1p, DNA binding, Betel D).

e.g. act1, rad51, dmcl

SEARCH

Analyse

Enter a list of identifiers.

Gene

rad51; rad52; rad53; ddcl; rad55; rad57; spo11; dmcl; rad17; rad9; rad24; msh1; msh5; mre11; vre2; nrd1Rn;

advanced ANALYSE

GENES PROTEINS GENE ONTOLOGY PHENOTYPES LITERATURE INTERACTIONS PATHWAYS GENE EX

Read more

Query for genes:

- Chromosome ➔ All genes.
- All genes in organism ➔ All chromosomal locations
- FeatureType ➔ Genes
- All genes in organism ➔ All genes with introns
- All genes of a selected Feature Type ➔ Genes with introns

Adding your Own

- **Synchronous**
 - data ready immediately
- **Asynchronous**
 - data needs to be generated by external source first

<https://bitbucket.org/galaxy/galaxy-central/wiki/DataSources>

Synchronous Datasource Tools

- When the user has finished at the external site, the external site directs the user back to Galaxy
- Parameters and a retrieval URL are provided
- In the background, Galaxy creates a new dataset and then uses provided URL and parameters to retrieve the new dataset content

UCSC Table Browser Example

Asynchronous Datasource Tools

- When the user has finished at the external site, the external site directs the user back to Galaxy
- Parameters and an URL is provided
- In the background, Galaxy creates a new dataset and then creates a new GALAXY_URL (uniquely identifying this new dataset) and sends this and provided parameters to the data source
- External data source then runs processes to generate dataset contents and sends a URL back to GALAXY_URL

UCSC Table Browser Example

A

Chromosome	Start	End	Name
chr1	11973	14409	uc001aaa.2.0+
chr1	11973	14409	uc001aaq.1.0+
chr1	11973	14409	uc001aav.1.0-
chr1	14363	14565	uc001aay.2.0-
chr1	16897	17770	uc001aay.1.0-
chr1	15792	18063	uc001ayd.2.0-

To reset all user cart settings (including custom tracks), [click here](#).

B

A name, description and unique tool id are assigned

The default data_source.py script is used to retrieve the data from UCSC

The URL where Galaxy will forward the user when this tool is accessed

Default parameters provided by Galaxy to the data resource, including the automatic selection of the "Send results to Galaxy" option

A set of parameters provided by the Table Browser are translated into names and values that are known to Galaxy

A single output dataset is defined

```

<tool name="UCSC Main" id="ucsc_table_direct1" tool_type="data_source">
    <description>table browser</description>
    <command_interpreter>python</command_interpreter>
    <data_source>py Output $__app__.config.output_size_limit</data_source>
    <inputs action="http://genome.ucsc.edu/cgi-bin/hgTables" check_values="false" method="get">
        <param name="sendToGalaxy" type="hidden" value="1"/>
        <param name="hgta_compressType" type="hidden" value="none"/>
        <param name="hgta_outputType" type="hidden" value="bed"/>
    </inputs>
    <request_param_translation>
        <request_param galaxy_name="URL_method" remote_name="URL_method" missing="post"/>
        <request_param galaxy_name="dbkey" remote_name="db" missing="?"/>
        <request_param galaxy_name="data_type" remote_name="hgta_outputType" missing="tabular">
            <value_translation>
                <value galaxy_value="tabular" remote_value="primaryTable"/>
                <value galaxy_value="tabular" remote_value="selectedFields"/>
                <value galaxy_value="wig" remote_value="wigData"/>
                <value galaxy_value="interval" remote_value="tab"/>
                <value galaxy_value="html" remote_value="hyperlinks"/>
                <value galaxy_value="fasta" remote_value="sequence"/>
                <value galaxy_value="gtf" remote_value="gtf"/>
            </value_translation>
        </request_param>
        <request_param_translation>
        <outputs>
            <data name="output" format="tabular"/>
        </outputs>
        <options sanitize="False" refresh="True"/>
    </request_param_translation>
    <outputs>
        <data name="output" format="tabular"/>
    </outputs>
</tool>
```

C

```

<toolbox>
    <section name="Get Data" id="getdata">
        <tool file="data_source/upload.xml"/>
        <tool file="data_source/ucsc_tablebrowser.xml"/>
        <tool file="data_source/ucsc_tablebrowser_test.xml"/>
        <tool file="data_source/ucsc_tablebrowser_archaea.xml"/>
        <tool file="data_source/bx_browser.xml"/>
        <tool file="data_source/microbial_import.xml"/>
        <tool file="data_source/biomart.xml"/>
        <tool file="data_source/biomart_test.xml"/>
        <tool file="data_source/cbi_ncic_mart.xml"/>
        <tool file="data_source/gramene_mart.xml"/>
        <tool file="data_source/fly_modencode.xml"/>
        <tool file="data_source/flymine.xml"/>
        <tool file="data_source/flymine_test.xml"/>
        <tool file="data_source/modmine.xml"/>
        <tool file="data_source/ratmine.xml"/>
        <tool file="data_source/worm_modencode.xml"/>
        <tool file="data_source/wormbase.xml"/>
        <tool file="data_source/wormbase_test.xml"/>
        <tool file="data_source/epopathdb.xml"/>
        <tool file="data_source/encode_db.xml"/>
        <tool file="data_source/epigraph_import.xml"/>
        <tool file="data_source/epigraph_import_test.xml"/>
        <tool file="data_source/fbvar.xml"/>
        <tool file="validation/fix_errors.xml"/>
    </section>
    <section name="Send Data" id="send">
        <tool file="data_destination/epigraph.xml"/>
    </section>
</toolbox>
```

Setting Attributes of Newly obtained Datasets

Parameter name	Usage
Name	The external resource can provide a descriptive name for the retrieved data set. If not provided, a default name based upon the name provided in the XML tool configuration is used.
Info	A free-form text string that a resource can use to provide additional information about the data set.
data_type	The type of data returned to Galaxy. Examples include bed, sam, gff and maf.
Dbkey	If the data belongs to a single reference genome, this string is used to store this information. Examples include hg18, mm9 and canFam2.
URL	The user's history will be populated with a new data set containing the results returned by submitting all provided parameters to this URL.

Overview

- Tools
- Datasources
- External Display Applications

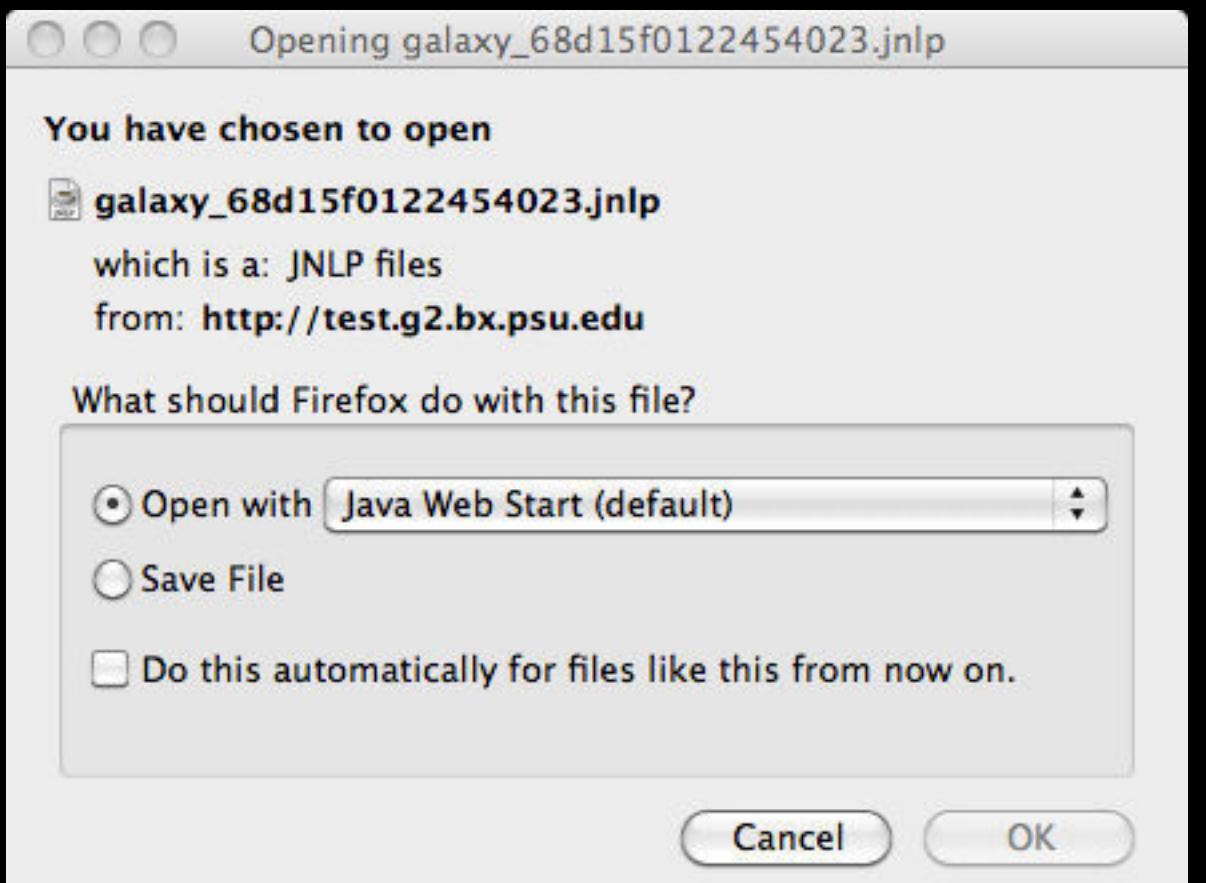
Recent Additions

1: Sample data

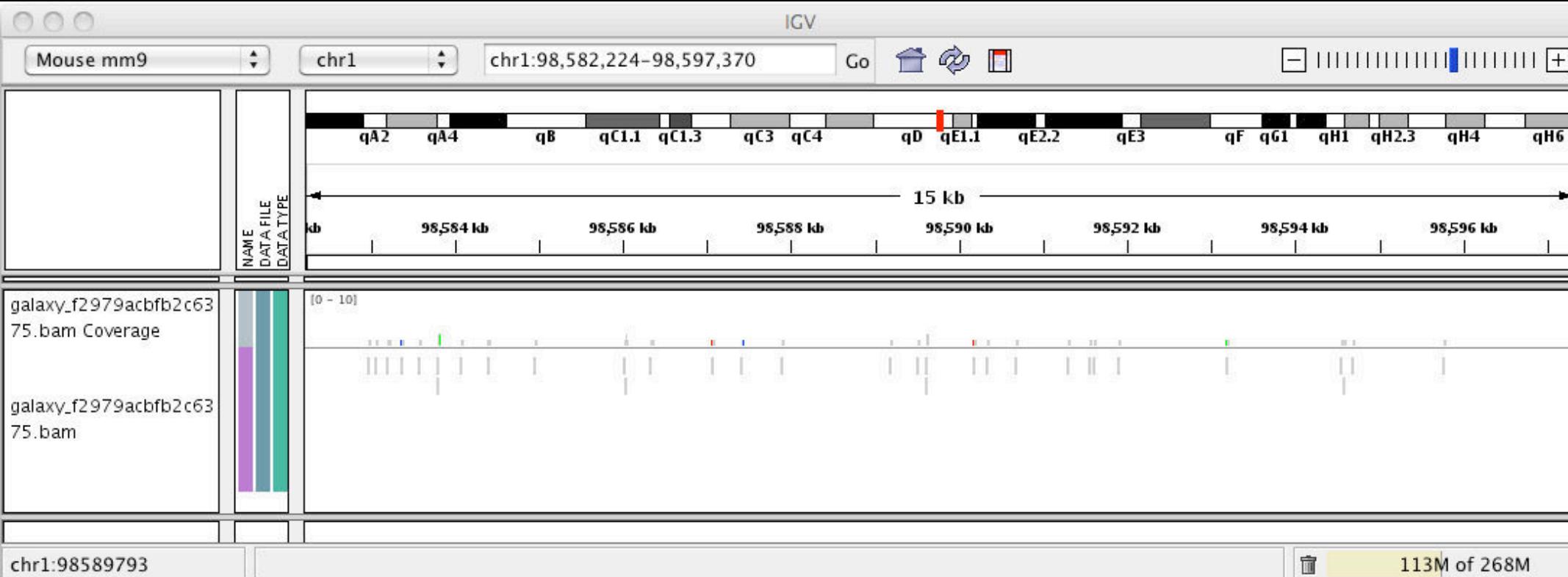
1.2 Gb
format: bam, database: mm9
Info: uploaded bam file

display at UCSC [main test](#)
display at Ensembl [Current](#)
display with IGV [web local](#)

Binary bam alignments file



Integrative Genomics Viewer (IGV)



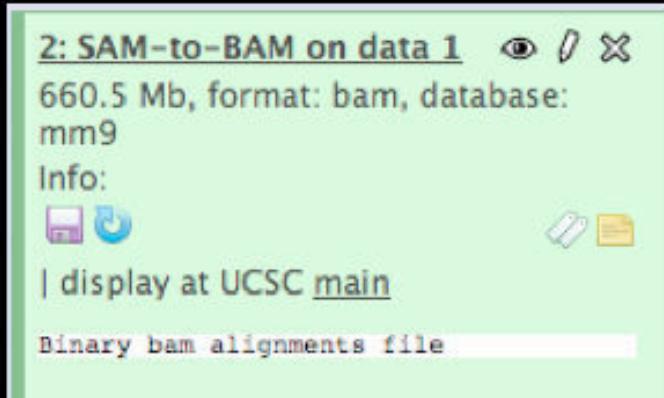
Adding your Own

- Define An XML configuration which describes how and where to present the data to the External Web Application
 - Static
 - Dynamic - display options can be loaded from a file
- Inform Galaxy about the new display by adding to the appropriate datatype in `datatypes_conf.xml`

Static External Display Application

```
<display id="ucsc_bam" version="1.0.0" name="display at UCSC">
  <link id="main" name="main">
    <url>http://genome.ucsc.edu/cgi-bin/hgTracks?db=${qp($bam_file.dbkey)}&hgt.customText=${qp($track.url)}</url>
    <param type="data" name="bam_file" url="galaxy.bam" strip_https="True" />
    <param type="data" name="bai_file" url="galaxy.bam.bai" metadata="bam_index" strip_https="True" />
    <param type="template" name="track" viewable="True" strip_https="True">
      track type=bam name="${bam_file.name}" bigDataUrl=${bam_file.url} db=${bam_file.dbkey}
    </param>
  </link>
</display>
```

```
<datatype extension="bam" type="galaxy.datatypes.binary:Bam"
  mimetype="application/octet-stream" display_in_upload="true">
  <display file="ucsc/bam.xml" />
</datatype>
```



BAM at UCSC

Home Genomes Blat Tables Gene Sorter PCR DNA Convert Ensembl PDF/PS Session Help

UCSC Genome Browser on Mouse July 2007 (NCBI37/mm9) Assembly

move <<< << < > >> zoom in 1.5x 3x 10x base zoom out 1.5x 3x 10x

position/search chr12:57,795,963-57,815,592 gene jump clear size 19,630 bp. configure

move start < 2.0 > Click on a feature for details. Click or drag in the base position track to zoom in. Click gray/blue bars on left for track options and descriptions. move end < 2.0 >

default tracks hide all manage custom tracks configure reverse refresh

collapse all expand all

Use drop-down controls below and press refresh to alter tracks displayed.
Tracks with lots of items will automatically be displayed in more compact modes.

Dynamic External Display Application

```
<display id="ucsc_bam" version="1.0.0" name="display at UCSC">
  <!-- Load links from file: one line to one link -->
  <dynamic_links from_file="tool-data/shared/ucsc/ucsc_build_sites.txt" skip_startswith="#" id="0" name="0">

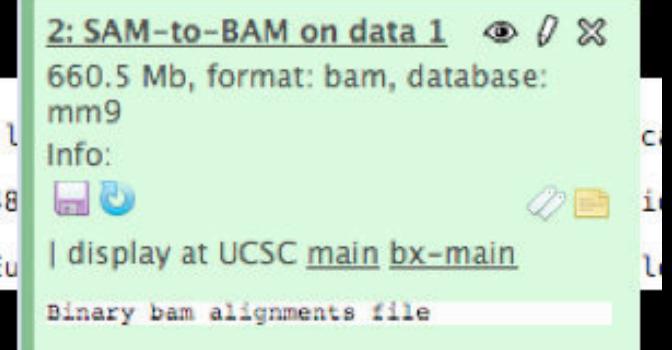
    <!-- Define parameters by column from file, allow splitting on builds -->
    <dynamic_param name="site_id" value="0"/>
    <dynamic_param name="ucsc_link" value="1"/>
    <dynamic_param name="builds" value="2" split="True" separator="," />

    <!-- Filter out some of the links based upon matching site_id to a Galaxy application configuration parameter and b
    <filter>${site_id} in $APP.config.ucsc_display_sites</filter>
    <filter>${dataset.dbkey} in $builds</filter>

    <!-- We define url and params as normal, but values defined in dynamic_param are available by specified name -->
    <url>${ucsc_link}db=${qp($bam_file.dbkey)}&hgt.customText=${qp($track.url)}</url>
    <param type="data" name="bam_file" url="galaxy_${DATASET_HASH}.bam" strip_https="True" />
    <param type="data" name="bai_file" url="galaxy_${DATASET_HASH}.bam.bai" metadata="bam_index" strip_https="True" />
    <param type="template" name="track" viewable="True" strip_https="True">
      track type=bam name="${bam_file.name}" bigDataURL=${bam_file.url} db=${bam_file.dbkey}
    </param>

  </dynamic_links>
</display>
```

```
#Harvested from http://genome.ucsc.edu/cgi-bin/das/dsn
main    http://genome.ucsc.edu/cgi-bin/hgTracks?    anoCar1,ce6,ce4,ce2,rn3,l
#Harvested from http://archaea.ucsc.edu/cgi-bin/das/dsn
archaea http://archaea.ucsc.edu/cgi-bin/hgTracks?    therSibi1,symbTher_IAM148
#Harvested from http://main.genome-browser.bx.psu.edu/cgi-bin/das/dsn
bx-main http://main.genome-browser.bx.psu.edu/cgi-bin/hgTracks?    oviAri1,eriEu
```



But I don't want to send my data to an external site

- Galaxy Trackster and Visual Analytics
 - Jeremy Goecks
 - 1:35 today

Using Galaxy

- Use public Galaxy server: [UseGalaxy.org](#)
- Download Galaxy source: [GetGalaxy.org](#)
- Galaxy Wiki: [GalaxyProject.org](#)
- Screencasts: [GalaxyCast.org](#)
- Public Mailing Lists
 - galaxy-bugs@bx.psu.edu
 - galaxy-user@bx.psu.edu
 - galaxy-dev@bx.psu.edu

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- All Members of the Galaxy Team
- Thousands of our users
- GMOD Team
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- BioMart Team
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 - Huck Institutes at Penn State
 - Pennsylvania Department of Public Health
 - Emory University

Galaxy Team



Enis Afgan | Emory



Guru Ananda | Penn State



Dannon Baker | Emory



James Taylor | Emory



Jeremy Goecks | Emory



Sergei Kosakovsky Pond | UCSD



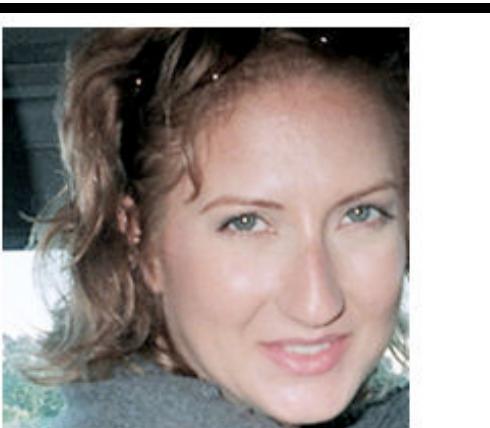
Greg von Kuster | Penn State



Dave Clements | Emory



Nate Coraor | Penn State



Jennifer Hillman Jackson | Penn State



Ross Lazarus | Harvard | BakerIDL



Kanwei Li | Emory



Anton Nekrutenko | Penn State



Kelly Vincent | Penn State