

Galaxy

Tools

Get Data

Get ENCODE Data

ENCODE Tools

Text Manipulation

Filter and Sort

Join, Subtract and Group

Convert Formats

Extract Features

Fetch Sequences

Fetch Alignments

Get Genomic Scores

Operate on Genomic Intervals

Intersect the intervals of two queries

Subtract the intervals of two queries

Merge the overlapping intervals of a query

Concatenate two queries into one query

Base Coverage of all intervals

Coverage of a set of intervals on second set of intervals

Complement intervals of a query

Cluster the intervals of a query

Join the intervals of two queries side-by-side

Get flanks returns flanking region/s for every gene

Statistics

Graph/Display Data

Evolution: HYPHY

EMBOSS

Join

Join:
1: RefSeq coding exons
First query

with:
2: SNPs
Second query

with min overlap:
1
(bp)

Return:
Only records that are joined (INNER JOIN)

Execute

7: Human/Chimp/Dog alignments for 957 sequences, format: fasta, database: hg18
Info: 319 regions were extracted successfully.
8886

>hg18_chr8(+)1:18301793-18302666
ATGGACATTGGACGATATTTTGAAGAAATTTGGCTATAGAACTCT7
>panTro2
ATGGACATTGGACGATATTTTGAAGAAATTTGGCTATAGAACTCT7
>canFam2

6: Coordinates of Exons with at least 5 SNPs

5: IDs Exons with at least 5 SNPs

4: Number of SNPs per Exon

3: Join of Exons and SNPs

2: SNPs

1: RefSeq coding exons

259,080 regions, format: bed, database: hg18
Info: Uploaded from UCSC
8886 | display at UCSC main

1 2 3 4

chr1 67052400 67052451 NM_024763_cds_0_6_chr1_ chr1 67066931 67067088 NM_024763_cds_1_0_chr1_ chr1 67065090 67065317 NM_024763_cds_2_0_chr1_ chr1 67064082 67064681 NM_024763_cds_3_0_chr1_ chr1 67071855 67071977 NM_024763_cds_4_0_chr1_ chr1 67072261 67072419 NM_024763_cds_5_0_chr1_

1: RefSeq coding exons

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chr1 67052400 67052451 NM_024763_cds_0_6_chr1_ chr1 67066931 67067088 NM_024763_cds_1_0_chr1_ chr1 67065090 67065317 NM_024763_cds_2_0_chr1_ chr1 67064082 67064681 NM_024763_cds_3_0_chr1_ chr1 67071855 67071977 NM_024763_cds_4_0_chr1_ chr1 67072261 67072419 NM_024763_cds_5_0_chr1_

Info: report bugs | wiki | screencasts | blog

Logged in as [sun1@psu.edu](#): manage | logout

Tools (left)

Tools (currently over 100) are organized in categories. Clicking on a category expands and shows available tools. For example, here the *Operate on Genomic Intervals* category is expanded.

Interfaces (center)

If you click a tool, its interface will appear here. For example, clicking on *Join* (left pane) will bring up its interface (shown in the center pane). If this case we are joining RefSeq exons with SNPs. The names of datasets in the two dropdowns ("1:RefSeq coding exons" and "2:SNPs") correspond to items listed in the right (History) pane. The tool interface also provides help information.

History (right)

This is where all your data and analyses results go. The output of every tool creates another history item, so that your original data is never changed. For example, we started with two datasets imported from UCSC: "1:RefSeq coding exons" and "2:SNPs". Joining the two datasets created a new History Item called "3:Join on Exons and SNPs". And so on... Thus the History serves as a complete record of every analyses. Clicking "options" (just above the right pane) allows saving and sharing (!) of histories.



VIVA LA EVOLUCIÓN

Introducing **GALAXY**, a revolutionary tool for reproducible genomic research!

<http://g2.bx.psu.edu>

(since you cannot click the paper - flip it!)

See what **GALAXY** can do for you:

Biologists

Use our public site (<http://g2.bx.psu.edu>) to access popular sources of data like the UCSC Table Browser. Run analyses right on the spot using a variety of integrated tools. Your results are always available and can be easily shared with others.

Software Developers

Galaxy is an easy-to-use, open-source, scalable framework for tool and data integration. Stop wasting time writing interfaces and get your tools used by biologists! GALAXY includes everything you need to get started, so download and start integrating!



Why?

You are an experimental biologist. You keep watching databases fill with more and more data. You keep thinking: “*even if I knew how to use Excel as a pro, it would probably not load 12,435,654 SNPs*”. So how do you perform analyses without calling somebody on the Computer Science side of campus? Suppose you want to find human exons with the highest SNP density. There is no straightforward way of doing it without learning programming first. And this is why:

Databases are not analyses tools

Databases are where you get the data. Browsers are where you visualize the results. For a bench biologist there is not much in between besides spreadsheets or Perl scripting. GALAXY fills this gap by providing a tool-rich analysis medium.

No tools for essential datatypes

Some datatypes generated by high throughput genomics are so new that there are no tools to analyze them. For example, how do you extract sequences of coding exons from the latest 28-way alignments of vertebrate genomes or analyze genome-wide disease association data? With GALAXY!

Is genomics reproducible?

The Methods section of too many papers sound like “*the data were analyzed using a collection of in-house scripts*”. How do you repeat such an analysis? GALAXY saves every step of your analysis and allows you to share these workflows with others.

Too many tools

Bioinformatics has published 166 Application Notes since January 2007. How does one know which tool to use? GALAXY integrates a multitude of different tools by giving them the same “look and feel” and linking them to data warehouses.



How?

So is it all hot air or can you actually do something with GALAXY that you cannot do otherwise? Let's try this:

find all human exons that contains at least 5 non-synonymous (amino acid changing) SNPs and extract their alignments with chimp and dog

This analysis requires 7 steps outlined below but can be completed easily and quickly with GALAXY (in fact, 90% of that time is spent downloading SNP data from the UCSC Table Browser). These steps correspond exactly to the contents of GALAXY's left pane on the opposite side of this brochure. You can watch a QuickTime screencast detailing this analysis at <http://g2.bx.psu.edu>.

Download exon and SNP coordinates

Here we just grab coordinates of all coding exons and SNPs from the UCSC Table Browser (the efforts of the UCSC Team were crucial for this integration). This creates History items #1 and #2 (right pane of Galaxy's interface on the opposite side)

Join exons with SNPs

This will find all exons that contain SNPs. This operation is performed using the *Join* function and creates History item #3. *Join* is a part of our large suite of tools for manipulation of genomic intervals that include intersection, coverage, and other functions.

Compute number of SNP per Exon

Here we first compute the number of SNPs per exon with the *Count* function (History item #4) and then use the *Filter* tool to find exons with at least 5 SNPs (History item #5).

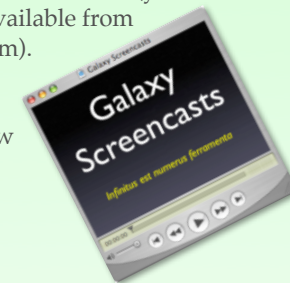
Extract human/chimp/dog alignments

Finally, we extract human/chimp/dog alignments corresponding to the exons from the previous step. These alignments are trimmed per exon boundaries. If multiple alignment blocks overlap an exon - they are stitched together (History item #6).



See it!

Still not convinced? Go to <http://g2.bx.psu.edu>. There you will see a link to screencasts (you will need a free QuickTime player available from Apple website to view them). Screencasts highlight all aspects of GALAXY's functionality and also show examples of very complex analyses.



Share!

Best of all, GALAXY's history system provides a complete analyses record that can be shared. Every history is an analysis workflow, which can be used to reproduce the entire analyses.

History is an analysis record

Look at GALAXY's interface on the opposite side. Every step of the analyses is recorded. You can have any number of histories saved. This way you can go back to your analyses anytime.

Share your analyses

Alice works at Penn State, while Bob suffers from the terrible San Diego climate. Alice wants Bob to see her analyses. Alice clicks the “*share*” link and enters Bob's e-mail address. Now Alice's history is visible to Bob (see “*Sharing history*” screencast).

Now your results are reproducible!

When publishing results, replace “*the data were analyzed using a collection of in-house scripts*” with a URL pointing to Galaxy's history. Your reviewers will have no further questions. That's reproducible genomics!