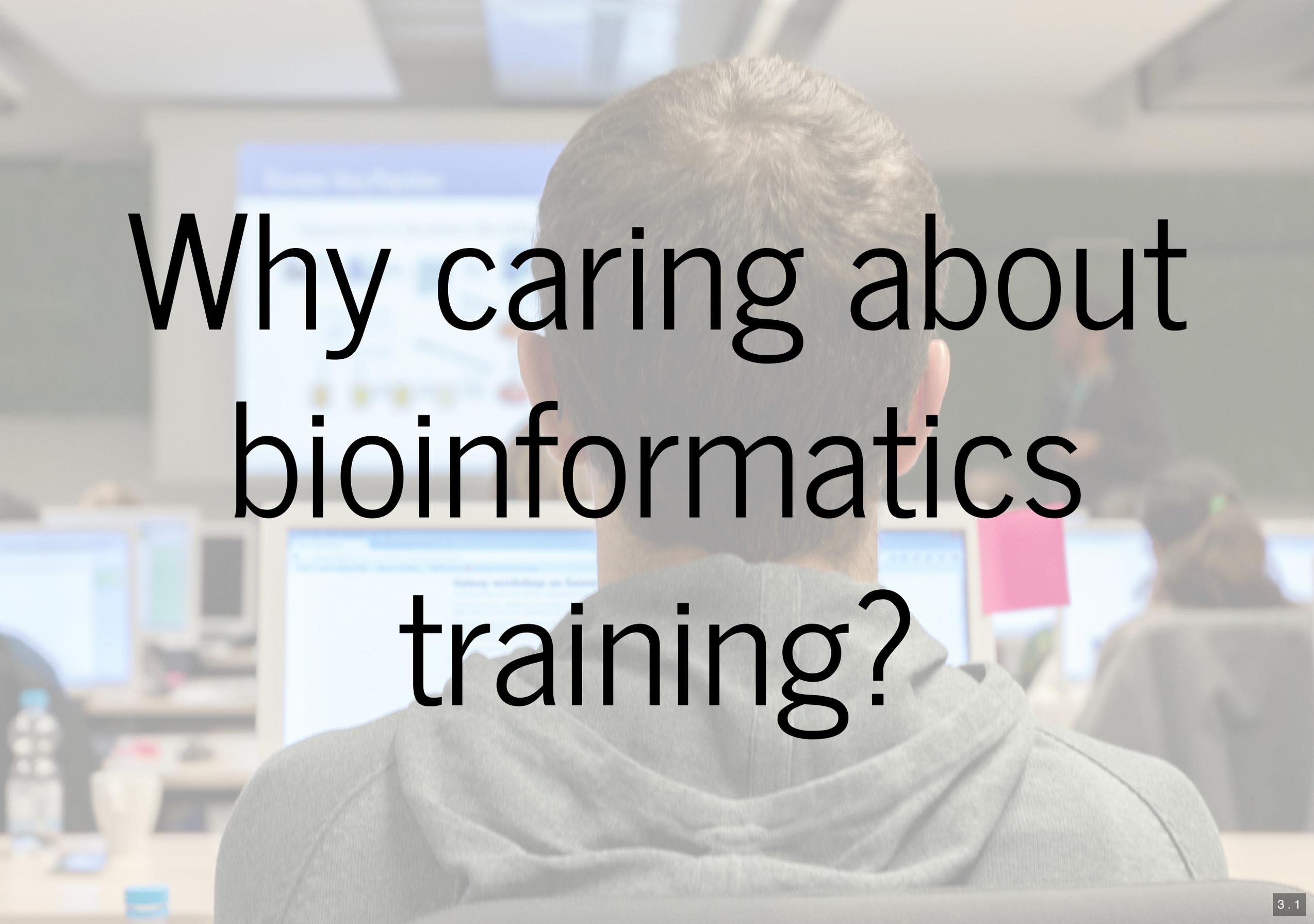


Building an open, collaborative, online infrastructure for bioinformatics training



B er enice Batut

33rd TBI Winterseminar
Februar 2018



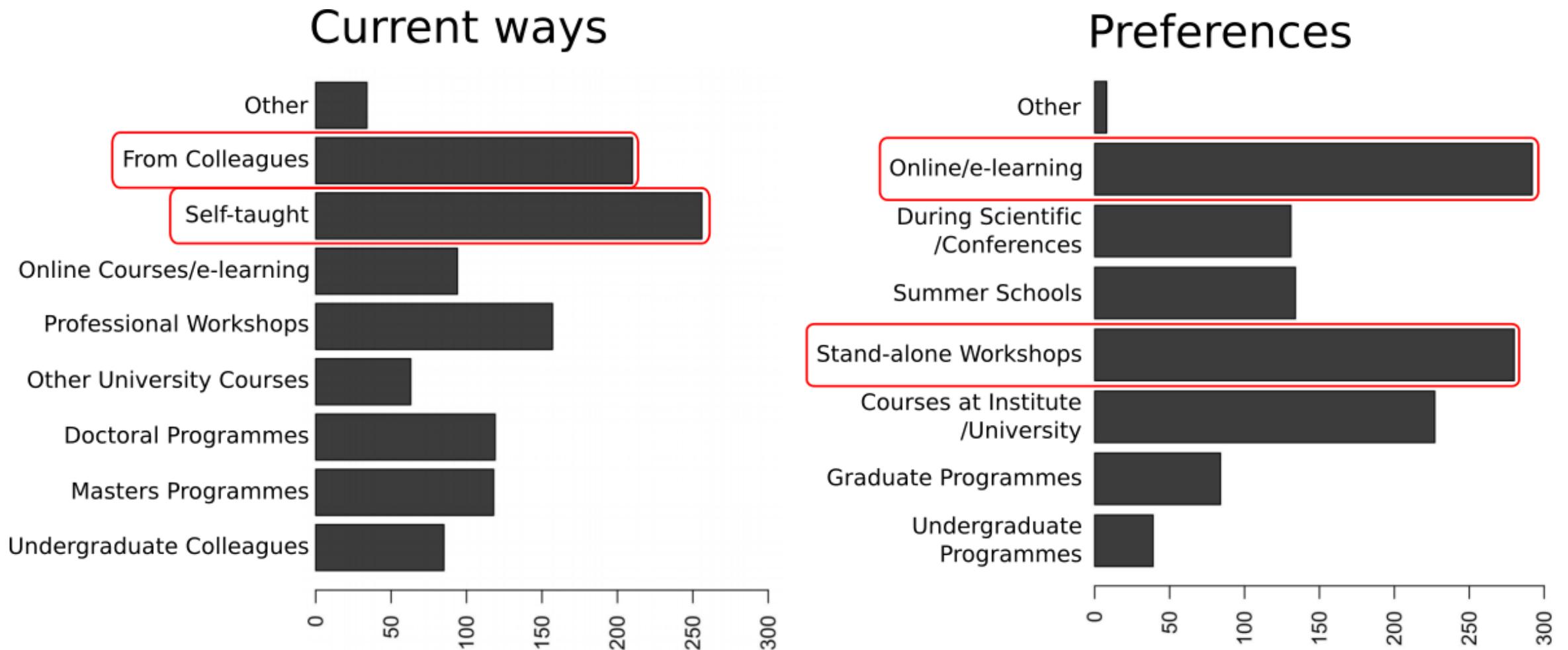
Why caring about
bioinformatics
training?

Need for bioinformatic training

*Bioinformatics has become too central to biology
to be left to specialist bioinformaticians*

- Explosion of data to analyze
- Access to computational power
- Thousand of possible tools for specialized analyses

An increasing demand for learning bioinformatics



Graphs of [Brazas et al, 2017](#)

Galaxy - Mozilla Firefox

Galaxy / Uni Freiburg

Tools

deepTools

computeMatrix has two main output options

scale-regions

Distance in bases to which all regions are going to be stretched or shrunk to

500

Set distance up- and downstream of each genomic region

no

Show advanced output settings

no

Show advanced options

no

History

ChIPseq_sept2016_afternoon

31: plotProfile on data 28: Underlying data

empty

format: tabular, database: hg18

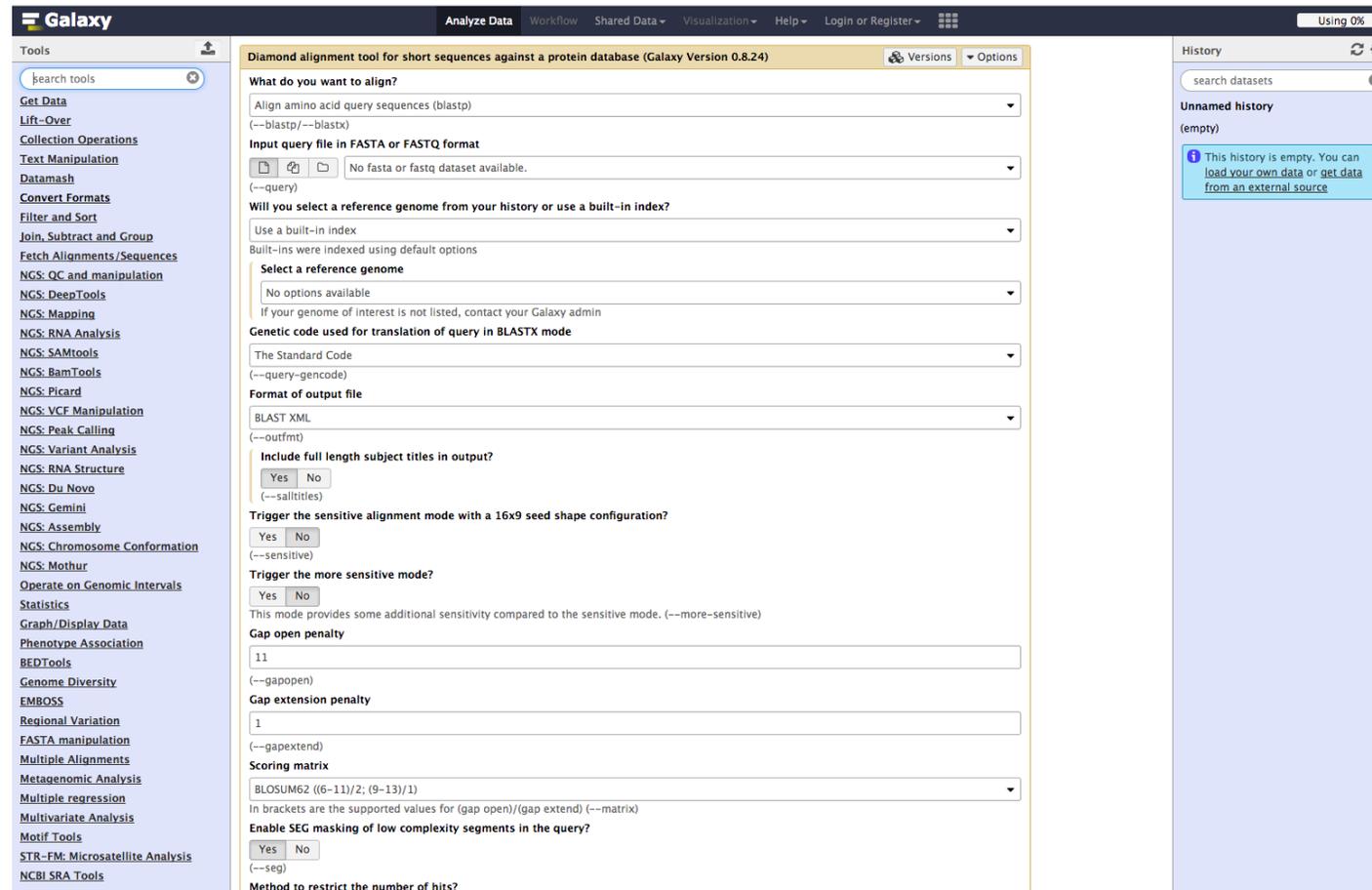
30: plotProfile on data 28: Underlying data

format: png, database: hg18

Galaxy

a great solution!

Computational knowledge: Not required!



- Web interface for numerous bioinformatics tools
- Scalable
- No issue with computer configuration during training

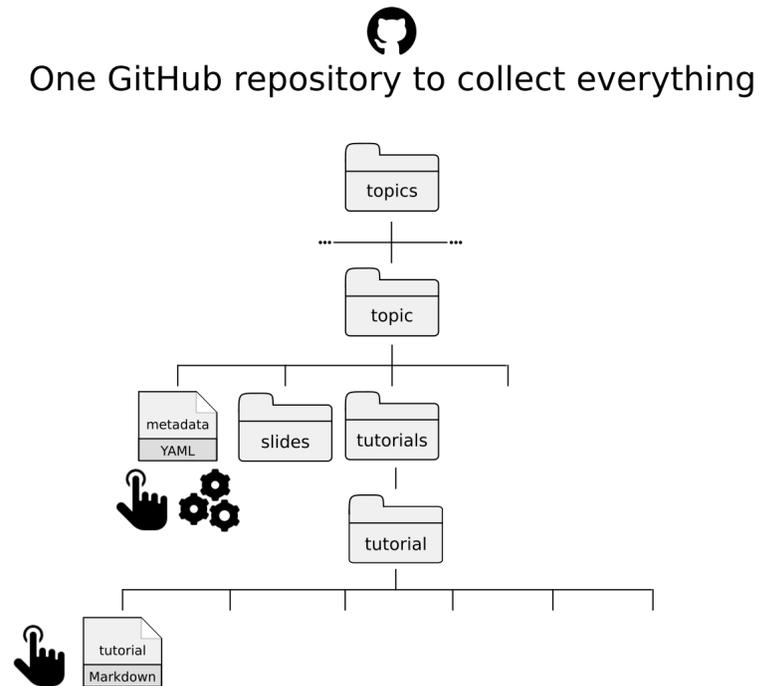


Building a new **open, collaborative** and **FAIR**
model for bioinformatics training

Requirements

- Easy to use
- Support for effective training for
 - Individual users
 - Instructors
- Definition of technological infrastructure
- Limited redundancy

An open, collaborative, FAIR, online infrastructure



-  Findable
-  Accessible
-  Interoperable
-  Reusable

Separation between content and format

Here treatment is the primary factor which we are interested in. The sequencing type is some further information that we know about the data that might affect the analysis. This particular multi-factor analysis allows us to assess the effect of the treatment, while taking the sequencing type into account, too.

```
> ### {% icon comment %} Comment
>
> We recommend you to add as many factors as you think may affect gene expression
in your experiment. It can be the sequencing type like here, but it can also be the
manipulation (if different persons are involved in the library preparation), ...
{: .comment}

> ### {% icon hands_on %} Hands-on: Analysis of the differential gene expression
(1)
>
> 1. Create a new history
> 2. Import the seven count files from [Zenodo]
(https://dx.doi.org/10.5281/zenodo.290221)
> - `GSM461176_untreat_single.deseq.counts`
> - `GSM461177_untreat_paired.deseq.counts`
> - `GSM461178_untreat_paired.deseq.counts`
> - `GSM461179_treat_single.deseq.counts`
> - `GSM461180_treat_paired.deseq.counts`
> - `GSM461181_treat_paired.deseq.counts`
> - `GSM461182_untreat_single.deseq.counts`
>
> 3. **DESeq2** {% icon tool %}: Run **DESeq2** with:
> - "Treatment" as first factor with "treated" and "untreated" as levels and
selection of count files corresponding to both levels
>
> > ### {% icon tip %} Tip
> >
> > You can select several files by keeping the CTRL (or COMMAND) key pressed
and clicking on the interesting files
> {: .tip}
>
```

Markdown



Here treatment is the primary factor which we are interested in. The sequencing type is some further information that we know about the data that might affect the analysis. This particular multi-factor analysis allows us to assess the effect of the treatment, while taking the sequencing type into account, too.

Comment

We recommend you to add as many factors as you think may affect gene expression in your experiment. It can be the sequencing type like here, but it can also be the manipulation (if different persons are involved in the library preparation), ...

Hands-on: Analysis of the differential gene expression (1)

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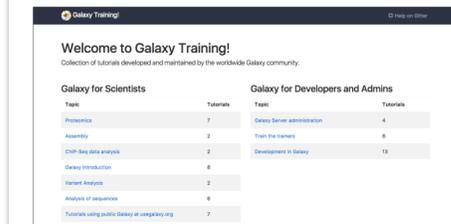
Tip

You can select several files by keeping the CTRL (or COMMAND) key pressed and clicking on the interesting files

User-friendly HTML

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Separation between content and form

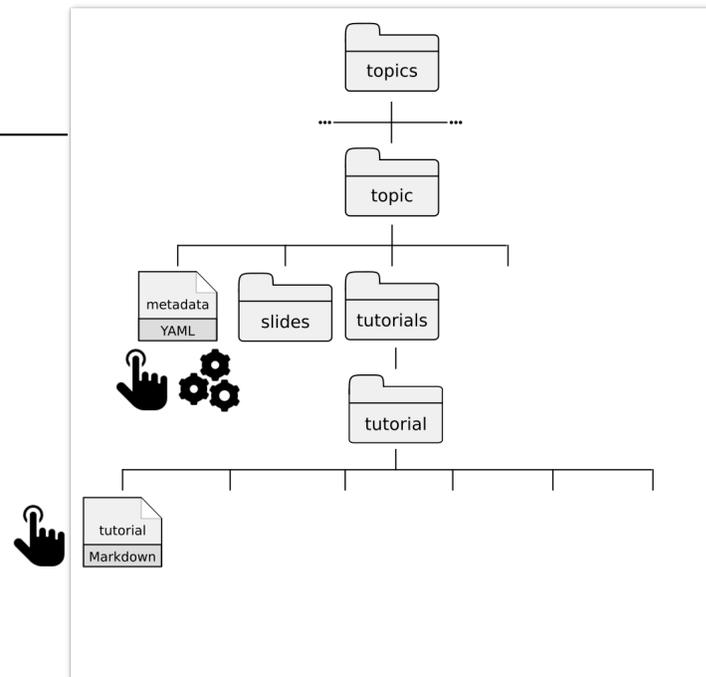


Welcome to Galaxy Training!
Collection of tutorials developed and maintained by the worldwide Galaxy community.

Galaxy for Scientists		Galaxy for Developers and Admins	
Topic	Tutorials	Topic	Tutorials
Proteomics	7	Galaxy Server administration	4
Assembly	2	Train the trainers	6
CHIP-seq data analysis	2	Development in Galaxy	15
Galaxy Introduction	8		
Variant Analysis	2		
Analysis of sequences	6		
Tutorials using public Galaxy at usegalaxy.org	7		

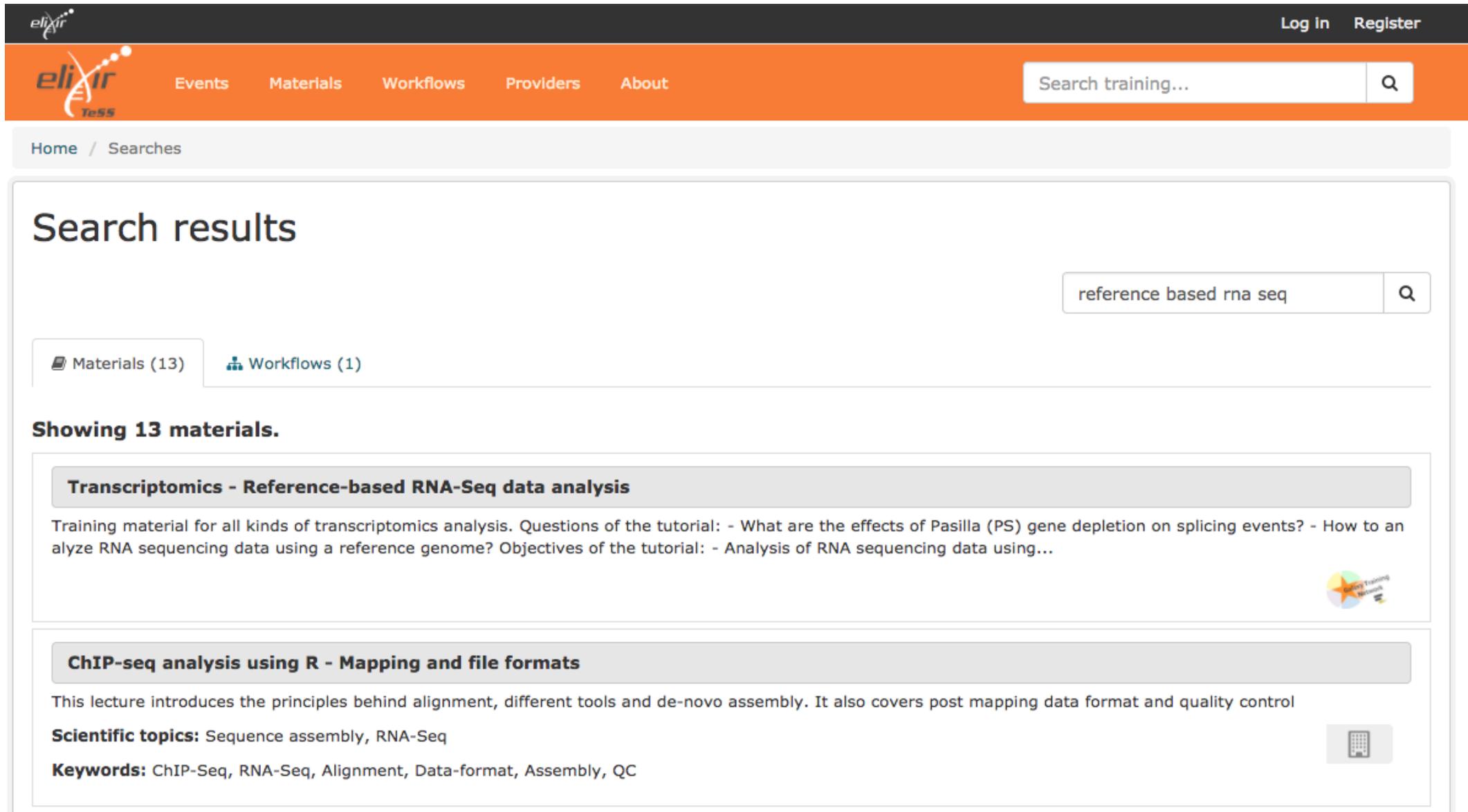


One GitHub repository to collect everything



-  Findable
-  Accessible
-  Interoperable
-  Reusable

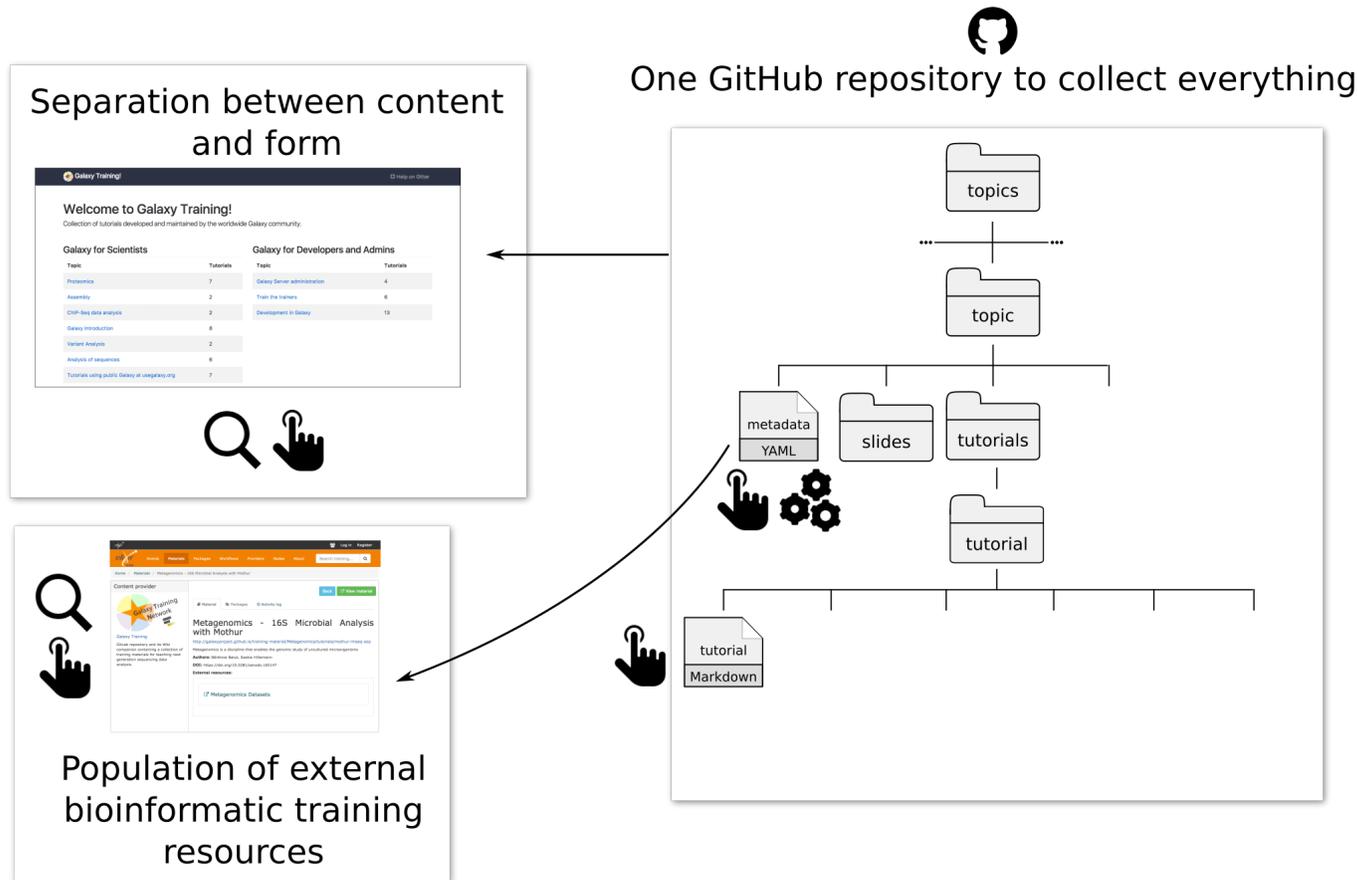
Findable material via TeSS



The screenshot shows the TeSS website interface. At the top, there is a navigation bar with the Elixir logo and the text 'TeSS'. The navigation menu includes 'Events', 'Materials', 'Workflows', 'Providers', and 'About'. A search bar is located on the right side of the navigation bar, containing the text 'Search training...'. In the top right corner, there are links for 'Log In' and 'Register'. Below the navigation bar, there is a breadcrumb trail: 'Home / Searches'. The main content area is titled 'Search results'. On the right side of this area, there is a search bar containing the text 'reference based rna seq'. Below the search bar, there are two filters: 'Materials (13)' and 'Workflows (1)'. The main content area displays two search results. The first result is titled 'Transcriptomics - Reference-based RNA-Seq data analysis'. The description for this result is: 'Training material for all kinds of transcriptomics analysis. Questions of the tutorial: - What are the effects of Pasilla (PS) gene depletion on splicing events? - How to analyze RNA sequencing data using a reference genome? Objectives of the tutorial: - Analysis of RNA sequencing data using...'. The second result is titled 'ChIP-seq analysis using R - Mapping and file formats'. The description for this result is: 'This lecture introduces the principles behind alignment, different tools and de-novo assembly. It also covers post mapping data format and quality control'. Below the description, there are two sections: 'Scientific topics: Sequence assembly, RNA-Seq' and 'Keywords: ChIP-Seq, RNA-Seq, Alignment, Data-format, Assembly, QC'. There are small icons next to each result: a star icon for the first result and a building icon for the second result.

<https://tess.elixir-europe.org/>

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Storage of training datasets on Zenodo

The screenshot shows the Zenodo website interface. At the top, there is a blue navigation bar with the Zenodo logo, a search bar, and links for 'Upload' and 'Communities'. On the right side of the bar are 'Log in' and 'Sign up' buttons. Below the navigation bar, the page title is 'Reference-based RNA-seq data analysis (training data)', dated 'February 10, 2017'. The authors listed are 'Batut, Bérénice; Videm, Pavankumar; Erxleben, Anika; Houwaart, Torsten; Grüning, Björn'. A description of RNA-seq is provided. A table of files is shown with columns for 'Name', 'Size', and 'Download' buttons. On the right side, there are sections for 'Indexed in' (OpenAIRE), 'Publication date', 'DOI' (10.5281/zenodo.290221), 'Communities' (Galaxy training network), 'License' (Creative Commons Attribution 4.0), and 'Share' (Cite as).

zenodo Search Upload Communities Log in Sign up

February 10, 2017 Dataset Open Access

Reference-based RNA-seq data analysis (training data)

Batut, Bérénice; Videm, Pavankumar; Erxleben, Anika; Houwaart, Torsten; Grüning, Björn

RNA-seq (RNA sequencing) uses high-throughput (HTS) data to reveal the presence and quantity of RNA in a biological sample at a given moment in time.

In the training available at http://galaxyproject.github.io/RNA-Seq/tutorials/ref_based, we introduce the bioinformatics methods to analyze RNA-seq data using a reference genome. The toy datasets were extracted from the study of Brooks *et al.* 2011.

Name	Size	Download
dexseq.gtf md5:51df689fc40c62fe062f6338f1e1c0f6	61.7 kB	Download
Drosophila_melanogaster.BDGP5.78.gtf md5:e07b8dcbd4f4f7602feeba0d5ed1698e	160.8 MB	Download
GSM461176_untreat_single.deseq.counts md5:409bc406c787b0f652ba0778e1e3eaa9	231.3 kB	Download

Indexed in
OpenAIRE

Publication date:
February 10, 2017

DOI:
DOI 10.5281/zenodo.290221

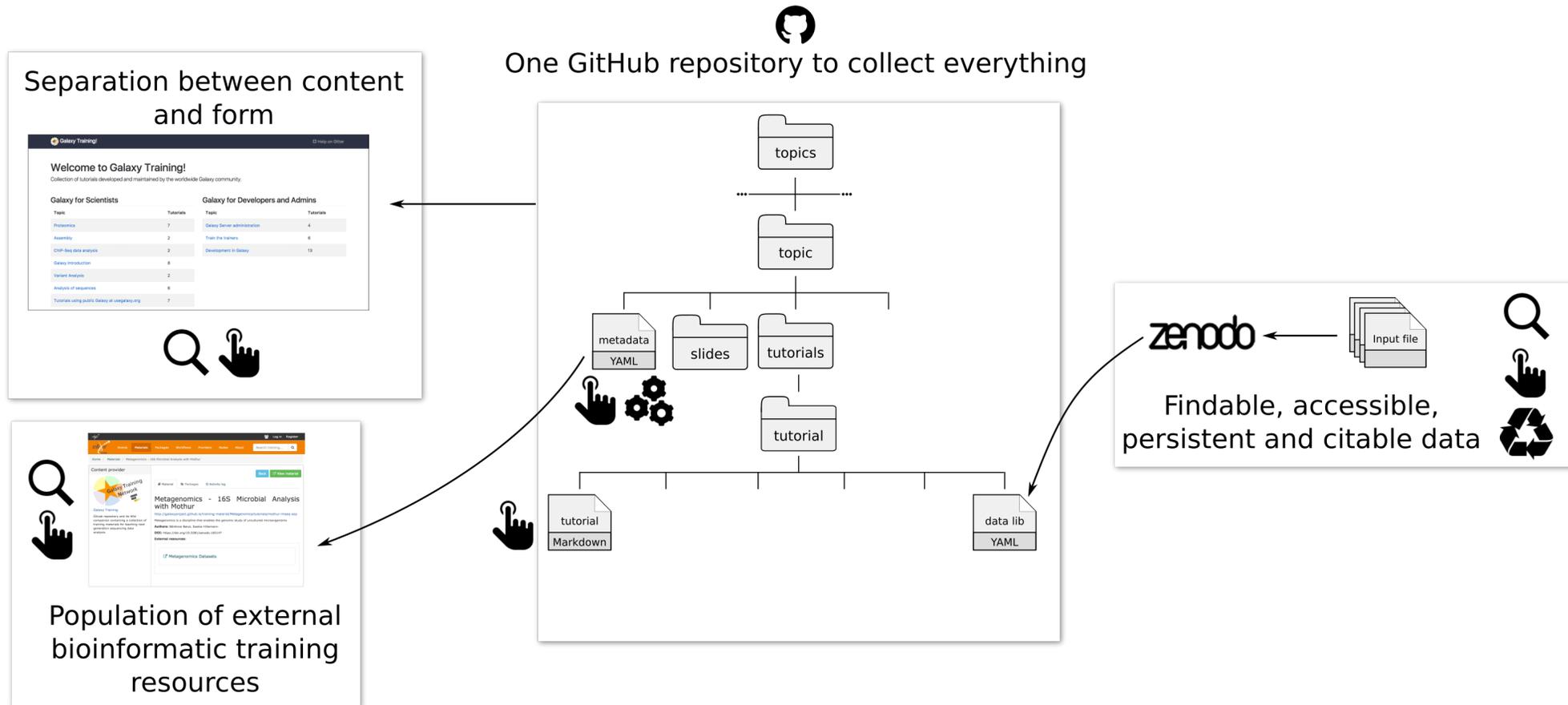
Communities:
Galaxy training network

License (for files):
Creative Commons Attribution 4.0

Share
Cite as
Batut, Bérénice, Videm, Pavankumar, Erxleben, Anika, Houwaart, Torsten, & Grüning, Björn. (2017). Reference-based RNA-seq data analysis (training data) [Data set]. Zenodo. <http://doi.org/10.5281/zenodo.290221>

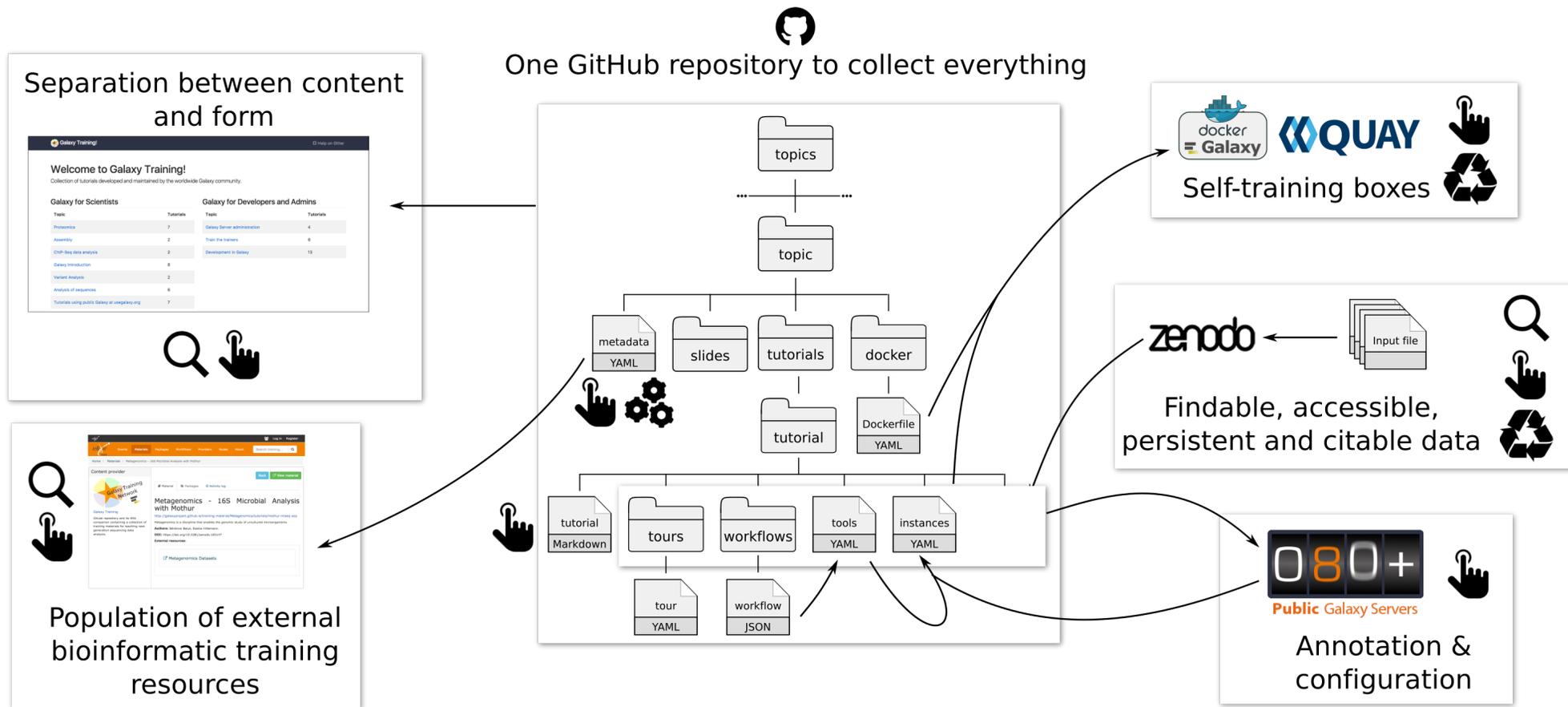
<https://zenodo.org/>

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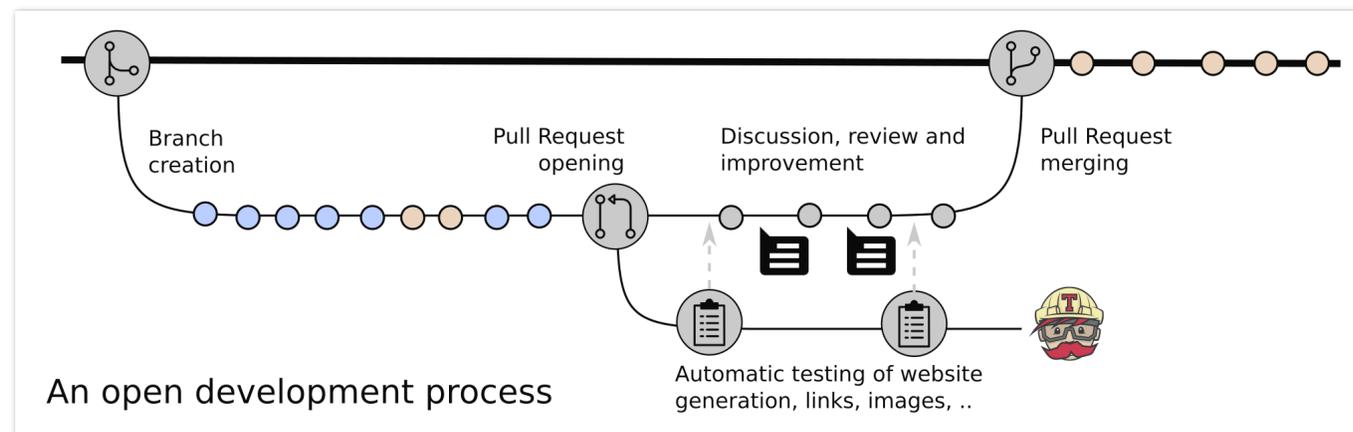
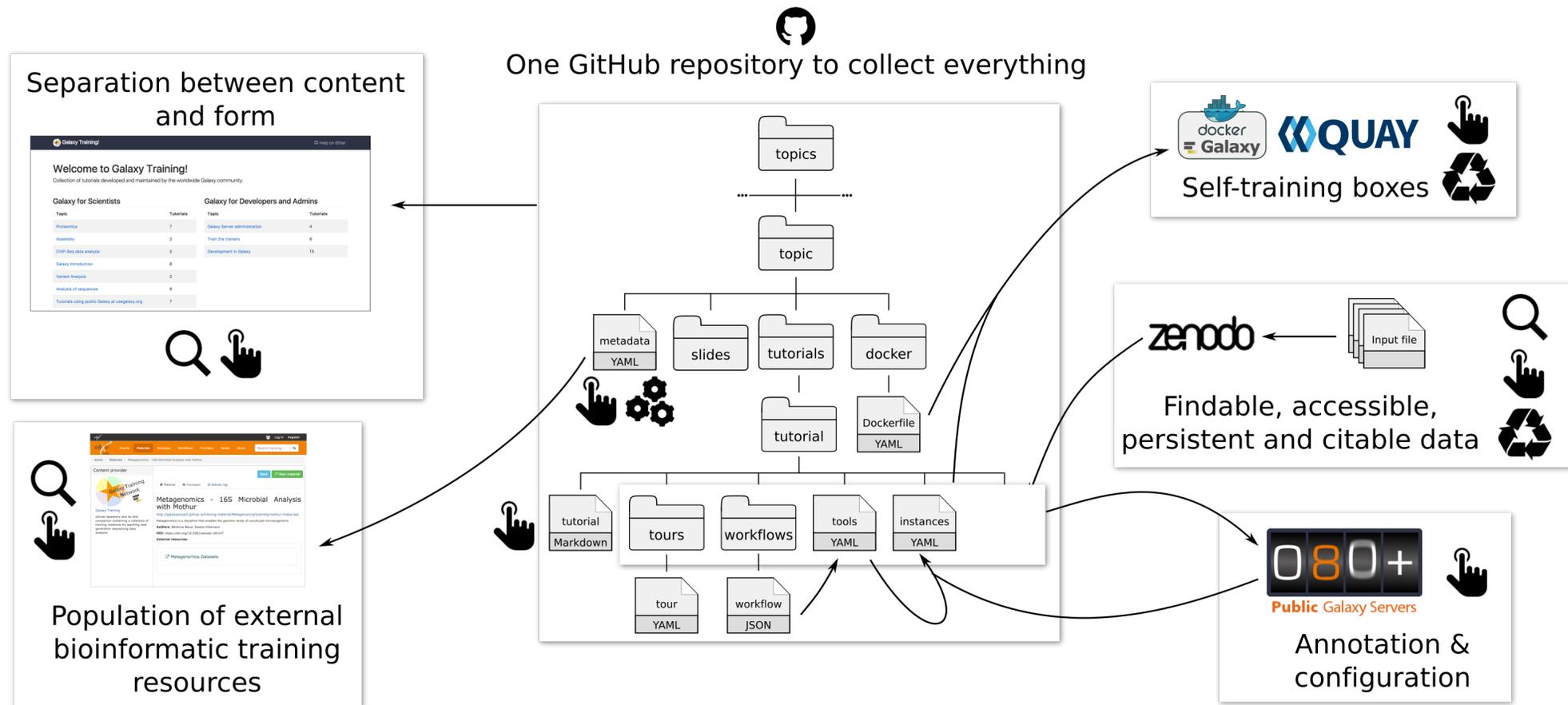
-  Findable
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Galaxy Training materials

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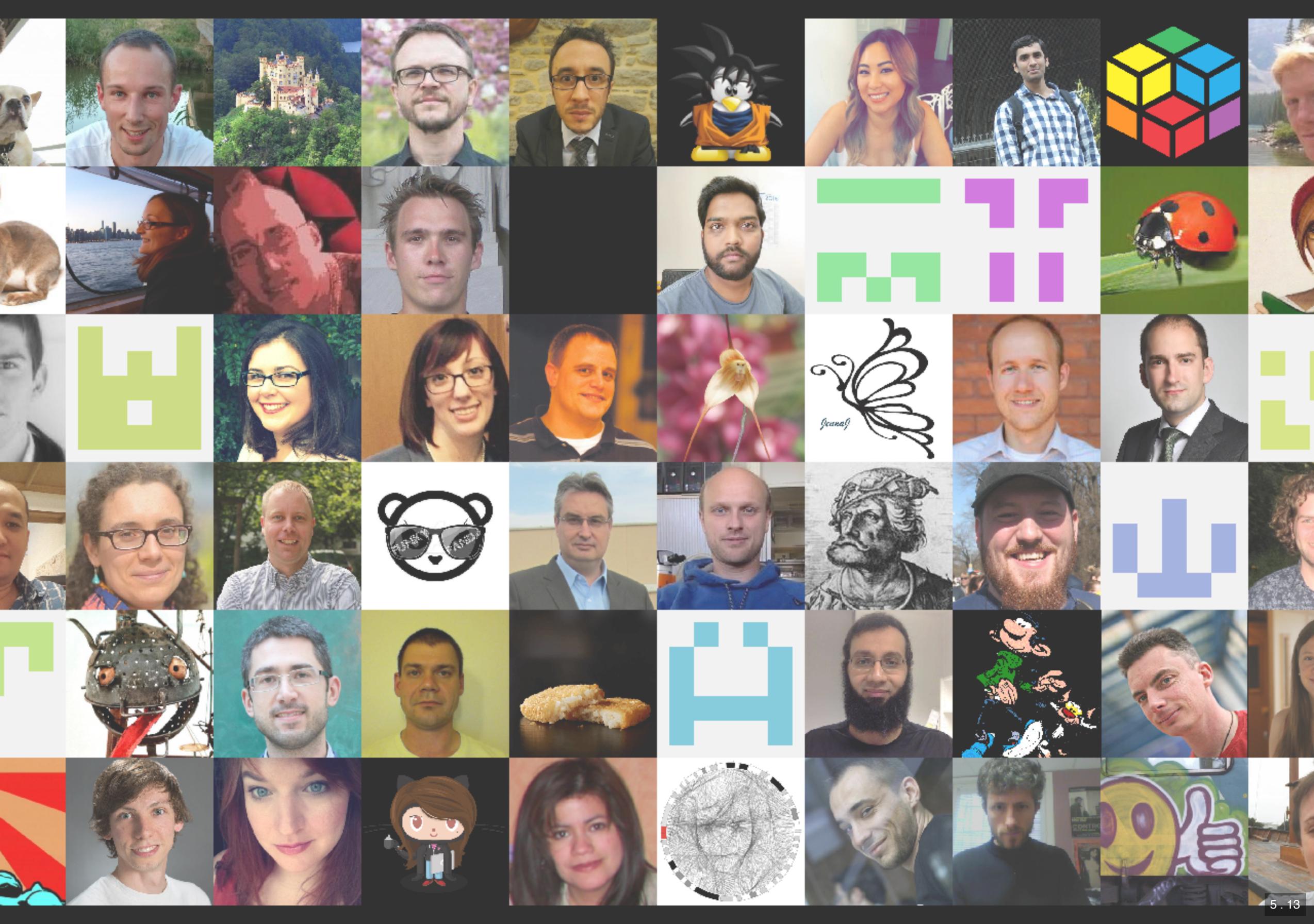
Galaxy for Scientists

Topic	Tutorials
Introduction to Galaxy	13
Assembly	3
ChIP-Seq data analysis	2
Epigenetics	1
Metagenomics	2
Proteomics	8
Sequence analysis	6
Transcriptomics	5
Variant Analysis	5

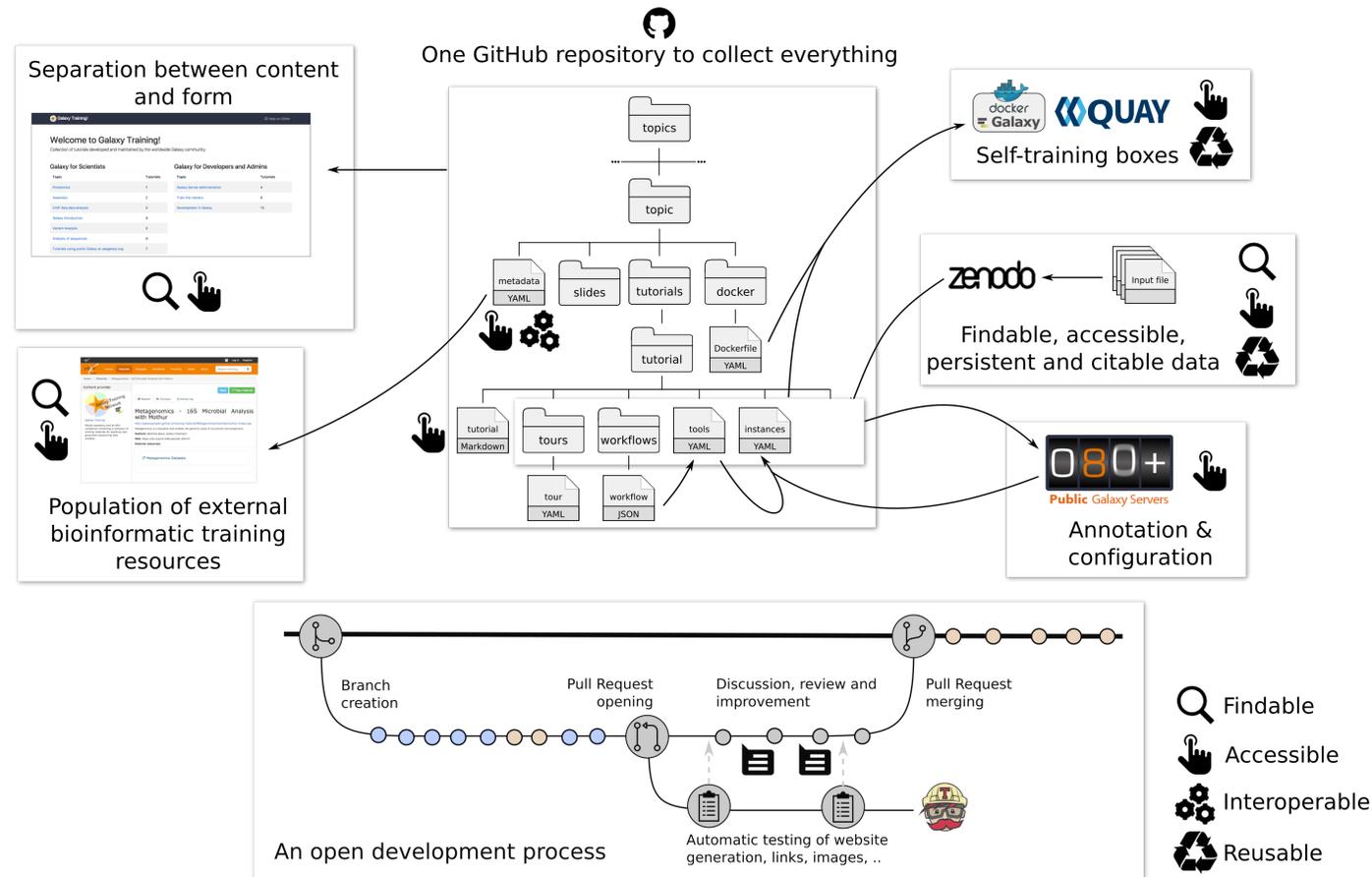
Galaxy for Developers and Admins

Topic	Tutorials
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Development in Galaxy	13
Train the trainers	6

<http://training.galaxyproject.org>



Thank you!



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 training.galaxyproject.org

 github.com/galaxyproject/training-material