



- Clustering and Alignment of ChIP profiles

User Manual

Revised November 16, 2010
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What is it?

CATCH is a cool new tool for clustering of ChIP profile patterns

Why would I use it?

To unravel the mysteries of epigenetics, you first have to identify the epigenetic patterns!



How do I use it?

1. Collect or generate **ChIP profiling data**

You probably already did this...

2. Identify your **genomic positions of interest**

Which sites do you want to analyse?

3. **CATCH** your ChIP profiles!

And let CATCH do alignment and clustering for you



The CATCH program

- **Unsupervised clustering** of ChIP profiles
- Input
 - A dataset of profiles, i.e. specific genome regions from your ChIP profiling experiment
 - ChIP signal (wiggle format)
 - Selected positions (bed format)
- Output
 - A clustering+alignment of all profiles from the dataset



The CATCH workflow

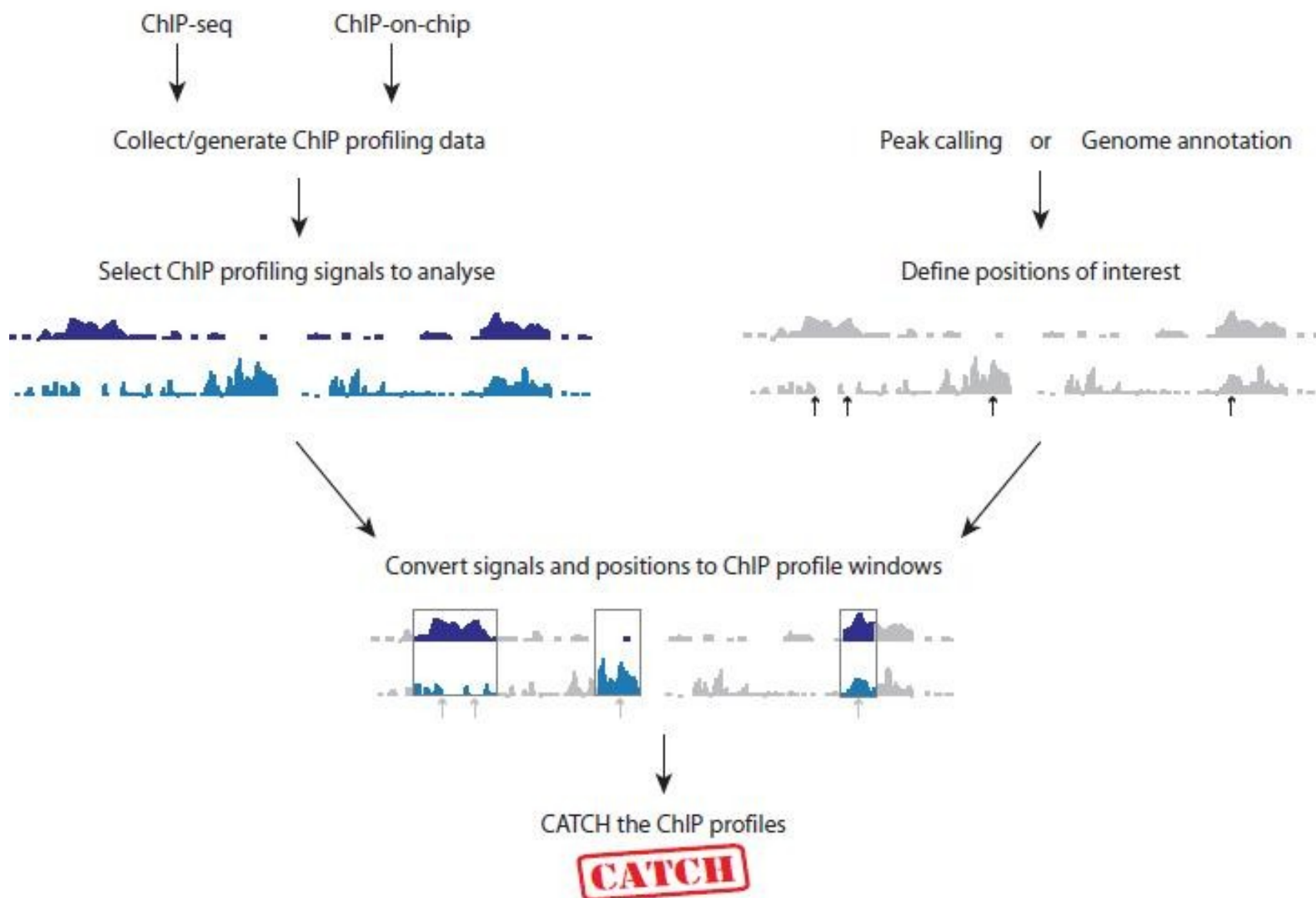
(small datasets)

- 1) Import profiles
- 2) View profiles
- 3) Start CATCH algorithm
- 4) Browse patterns in clustering tree
- 5) Export interesting profiles for further analysis

The import procedure has more steps when working with large datasets (1000+ profile regions) ... →



The CATCH workflow





The CATCH workflow

(large datasets)

- 1) Generate profile wiggle files (pwig)
 - Load and view dataset in CATCHprofiles
- 2) Make json input file from pwig files
- 3) Start CATCH algorithm from commandline
 - Generating json output file
- 4) Load json result in CATCHprofiles

Example workflow

1. make a bed file of your regions of interest

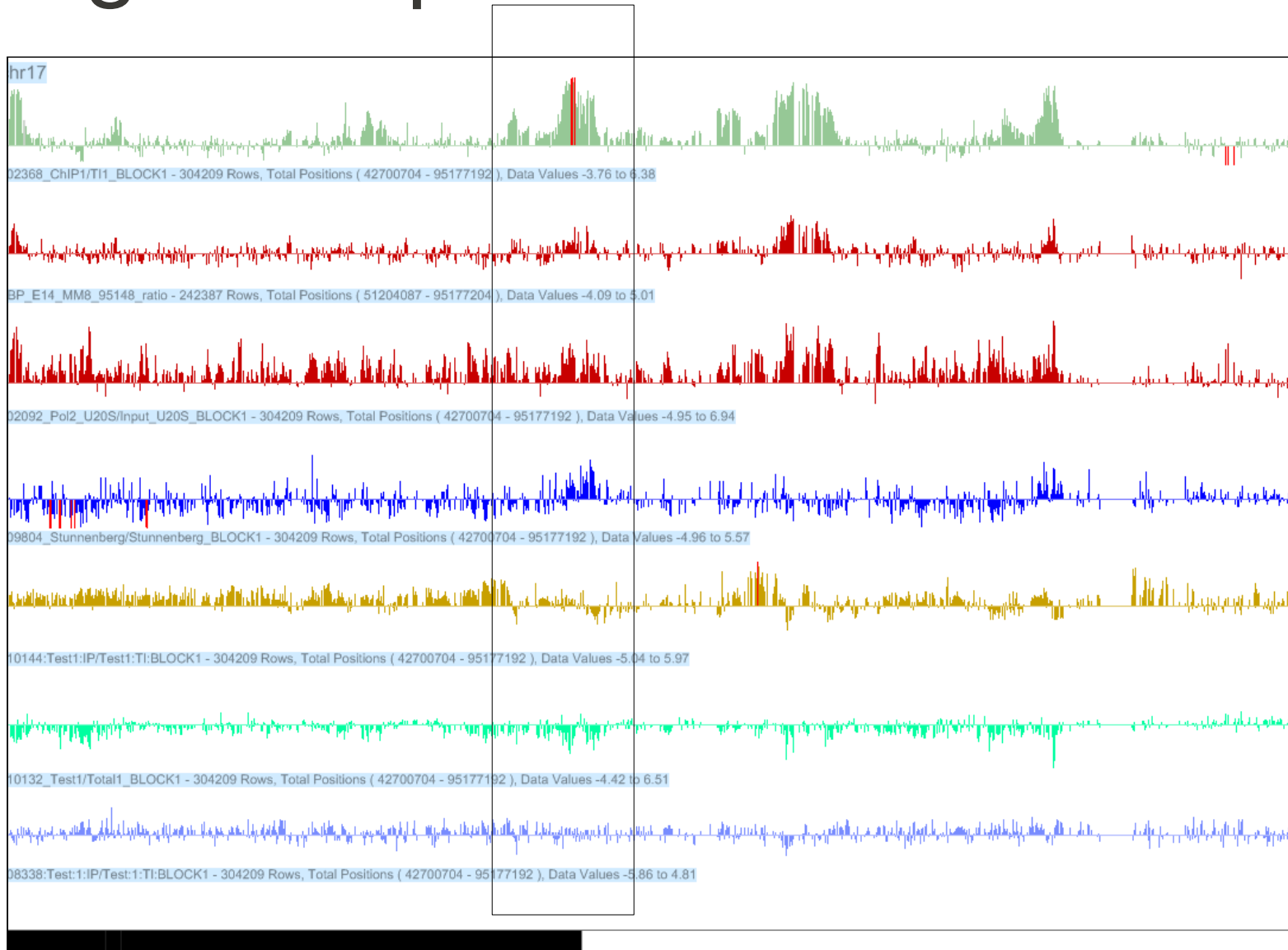
Profile regions

- You decide which areas are interesting to you:
 - Regions of annotation (TSS, CPG islands etc)
 - Peak regions
 - Other

Example workflow

1. make a bed file of your regions of interest

e.g. TSS profiles in ChIP-on-chip



TSS

Import to CATCH

2. import your wiggle files and bed file using File → 'Import Profiles'

File View

Signal Files (.wig)

Position Files (.bed)

Convert and add in CATCH

☒ Width of profile window (BP)

2000

☐ Profile window specified in ...

☐ Use merge position (BP)

1000

Ref point interval

38

Cancel

Add Signal Files

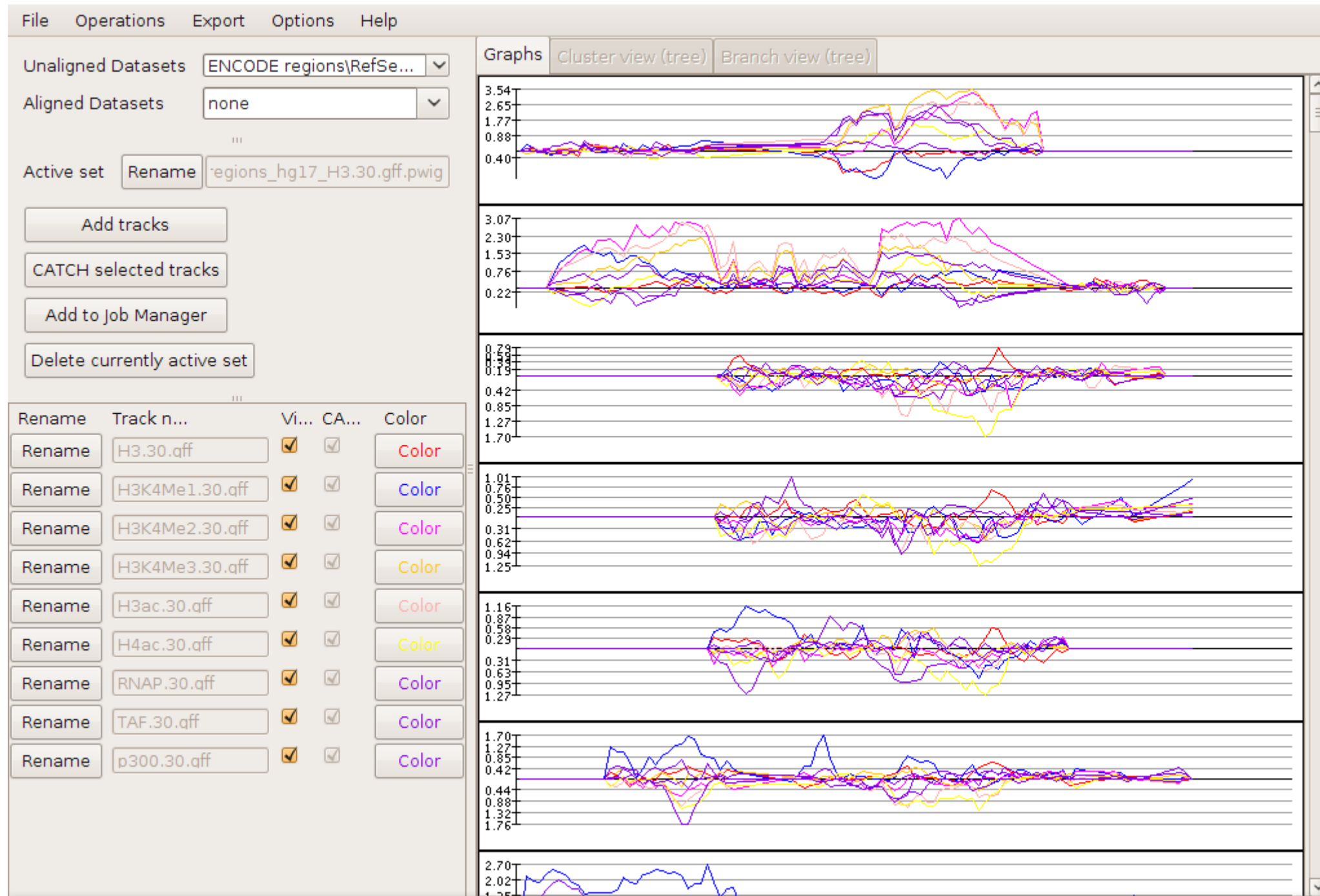
Remove selected Signal Files

Add Position Files

Remove selected Position Files

View profiles

3. Examine your data



Start CATCH

4. Click 'CATCH selected tracks' and wait..

File Operations Export Options Help

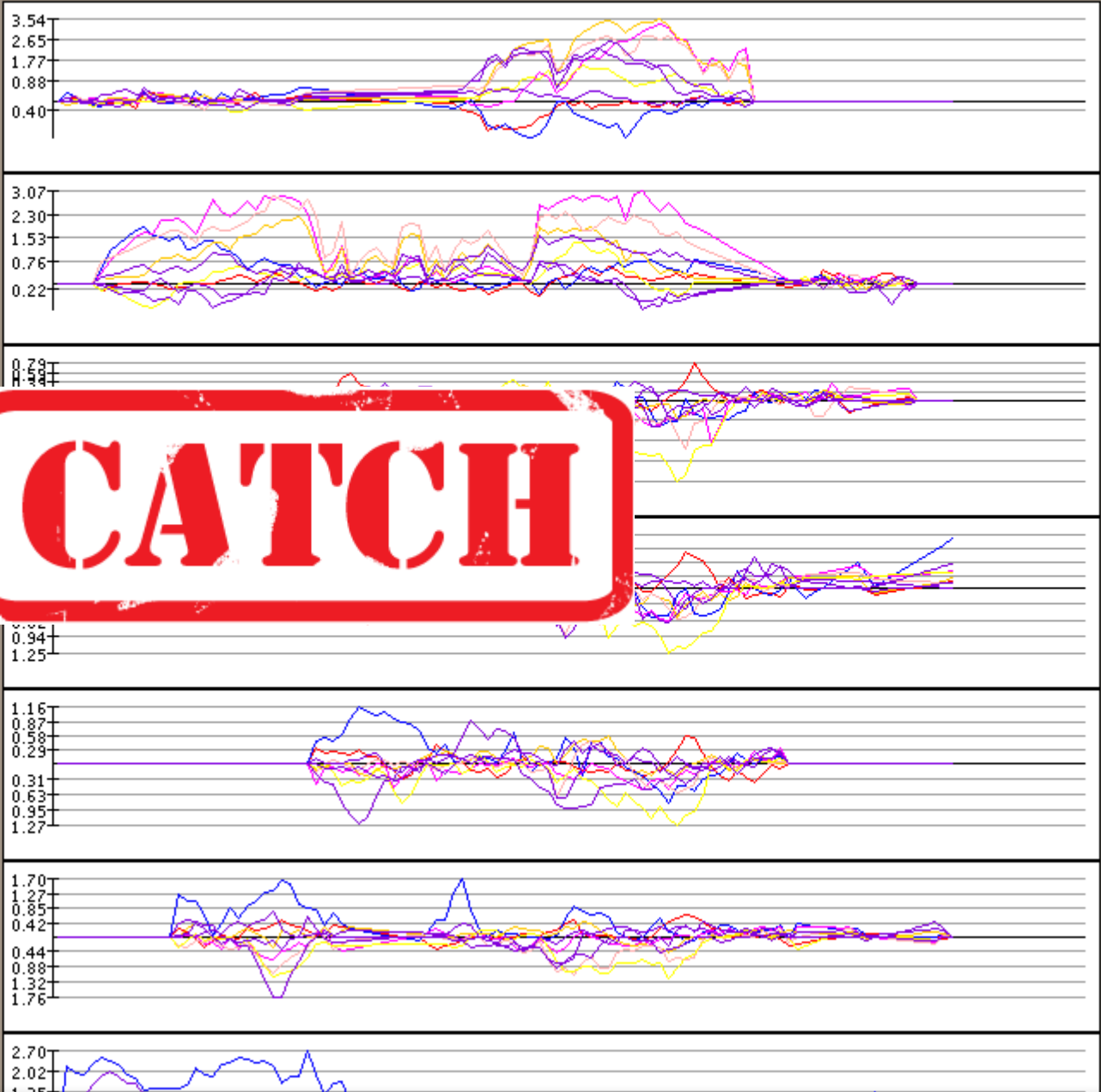
Unaligned Datasets ENCODE regions\RefSe...
Aligned Datasets none

Active set Rename regions_hg17_H3.30.gff.pwig

Add tracks
CATCH selected tracks
Add to Job Manager
Delete currently active set

Rename	Track n...	Vi...	CA...	Color
Rename	H3.30.gff	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	Color
Rename	H3K4Me1.30.gff	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	Color
Rename	H3K4Me2.30.gff	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	Color
Rename	H3K4Me3.30.gff	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	Color
Rename	H3ac.30.gff	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	Color
Rename	H4ac.30.gff	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	Color
Rename	RNAP.30.gff	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	Color
Rename	TAF.30.gff	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	Color
Rename	p300.30.gff	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	Color

Graphs Cluster view (tree) Branch view (tree)



CATCH

Examine profile clusters 4. Browse results interactively

File Operations Export Options Help

Unaligned Datasets none

Aligned Datasets TSS5kb_merge2kb

Active set Rename TSS5kb_merge2kb

Add tracks

CATCH selected tracks

Add to job Manager

Delete currently active set

Rename	Track n...	Vi...	CA...	Color
Rename	H3.30.qff	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	Color
Rename	H3K4Me1.30.qff	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	Color
Rename	H3K4Me2.30.qff	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	Color
Rename	H3K4Me3.30.qff	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	Color
Rename	H3ac.30.qff	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	Color
Rename	H4ac.30.qff	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	Color
Rename	RNAP.30.qff	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	Color
Rename	TAF.30.qff	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	Color
Rename	p300.30.qff	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	Color

Graphs Cluster view (tree) Branch view (tree)

Mouse-over clusters
to view profile below



interactive browsing

- Browse clustering tree:
 - Zoom: + and –
 - Collapse subtree: ctrl + leftclick
 - Highlight: leftclick
- Right-click to:
 - Open subtree
 - Export results



Save & Export

- CATCH clustering result (.catch)
 - To browse later or share results
- Use the export menu or right click to export:
 - Profile patterns (.csv)
 - Positions (.bed)
 - Cluster tree (.newick)



Quick Start Guide

1. Ensure JAVA 1.6 is installed
(follow link from <http://catch.cmbi.ru.nl>)
2. Prepare your data
 - ChIP profiling tracks in wiggle format
 - Your chosen peaks/positions in .bed format
3. Start CATCH
4. Import files or load dataset
5. CATCH your ChIP profiles!



Java memory issues

- Run CATCH from the commandline:
 - `Java -Xmx1000m -Xss2000k -jar CATCH3v513.jar`
- If you get errors of insufficient Java memory:
 - Heapsize: increase -Xmx value
 - Stacksize: increase -Xss value
- If 'Import files' fails on big files:
 - Use script to generate datasets (.pwig)
 - Use script to generate CATCH input file (.json)



Big datasets?

- Use scripts to generate input files in JSON format
- Execute CATCH from the commandline
 - `./execute.sh catch jobin.json jobout.json`
 - Where second argument is the name of the executable
 - `win: catch`
 - `linux: catch-linux-static-x86`
 - `mac: catch-macOSX`
- Load json result in CATCHprofiles

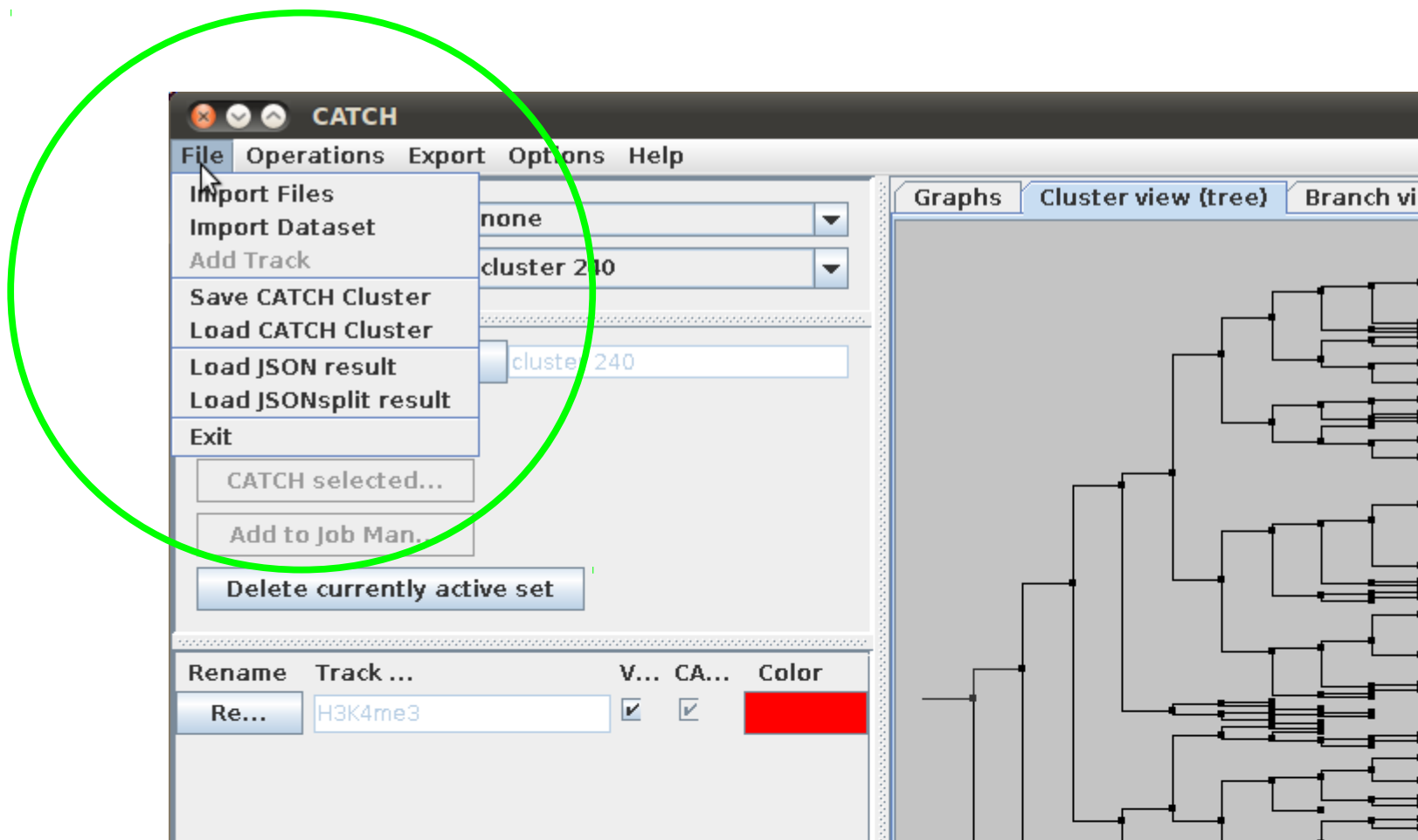


More information

- Go to the CATCH homepage:
<http://catch.cmbi.ru.nl>
 - User manual, examples and links
- Updates?
 - Subscribe to: CATCH-updates@bioinformatics.org
- Questions?
 - Send them to: CATCH-users@bioinformatics.org



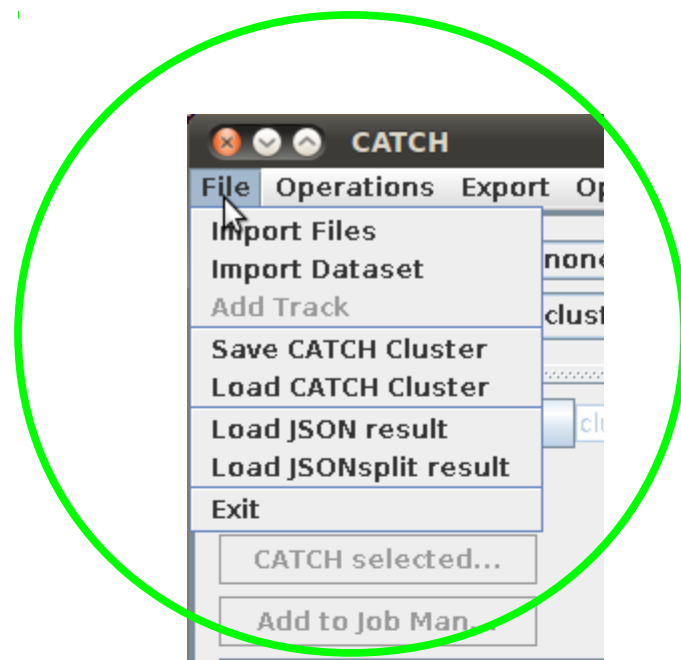
Load files





File formats

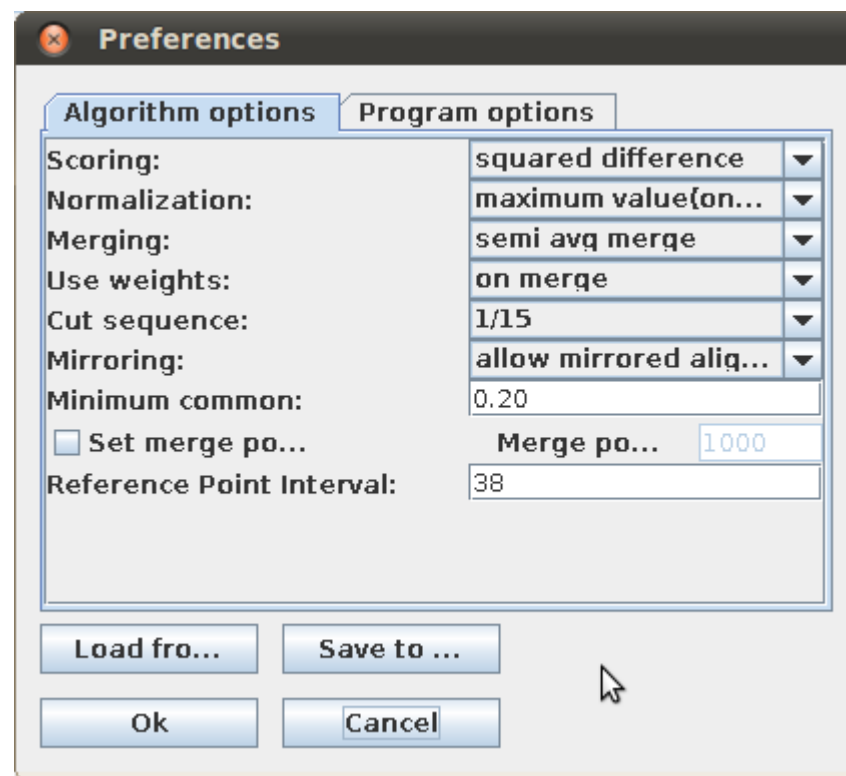
- Import files: **wiggle** and **bed**
- Import dataset: **pwig**
- CATCH cluster: **catch**
- JSON result: **json**





Change settings

- Check and save your default settings before starting the CATCH clustering
- Settings are written to the json-formatted input file before executing the CATCH core algorithm



Acknowledgements

IMADA, Odense:

- **Kasper Markus**
first implementation
of CATCH
- **Lene Favrholt**

Nijmegen:

- **Moniek Riemersma**
- **GiPCATCH** team
- **Maarten Kooyman**
The PinkThing

Supervisors:

Henk Stunnenberg

Molecular Biology

Martijn Huynen

CMBI

Acknowledgements

CATCH users:

Hendrik Marks

Rasa Jurgelenaite

Copenhagen:

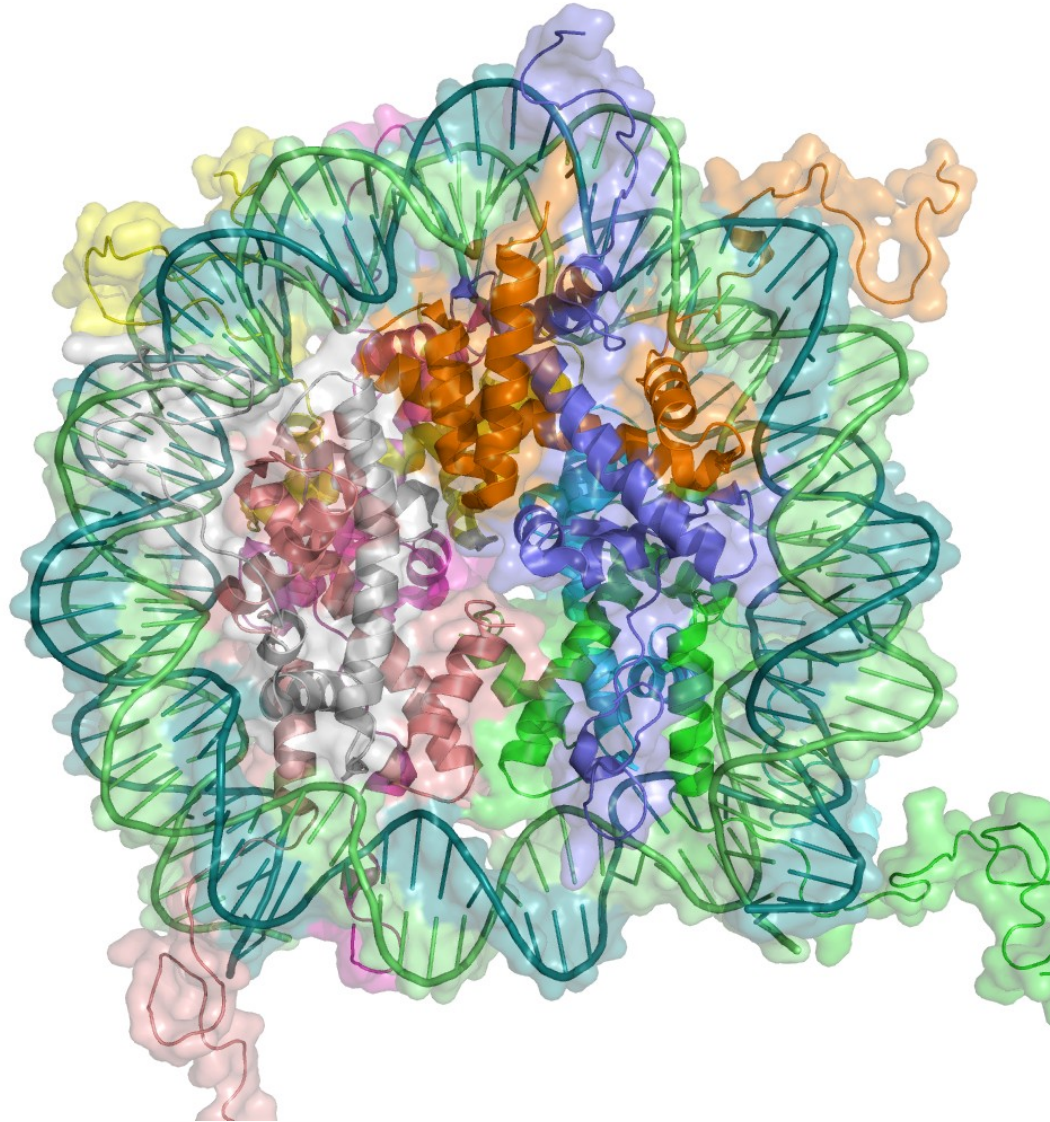
Rune Friborg



Funding:

HEROIC – mouse epigenetics

Thank you for using CATCHprofiles



<http://catch.cmbi.ru.nl>

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