SRB in the BioEmergences project

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The project The architecture SRB usage iRODS?

- The project
- 2 The architecture
- SRB usage
- 4 iRODS?



Partners

- Framework Program 6 project
- Consortium:
 - CNRS Centre De Recherche en Epistémologie Appliquée (CREA) (FRANCE)
 - Institut Curie (France)
 - Slovenska Technicka Univerzita V Bratislave (Slovakia)
 - Universidad de Málaga (Spain)
 - Denali Consulting S.A. (Belgium)
 - European Molecular Biology Laboratory (Germany)
 - University of Bologna (Italy)
 - CNRS CC-IN2P3 (France)
- Project fact sheet on CORDIS:http://tinyurl.com/5yc42k



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Project goals

What?

With the BioEMERGENCES project, we aim at providing an **experimental platform** to observe **in vivo** emergent patterns at various scales and **measure their variability between different individuals** of the same species. This is a strategy towards the measurement of the individual susceptibility to genetic diseases or response to treatments.

. . .

The main result expected from BioEMERGENCES is the specification of a European platform to achieve high throughput measurement of individual differences and screening of drugs combinations such as bi or tri-therapies.



Goals

Team

 Multi-disciplinary team : biologists, mathematicians, engineers, computer scientists

Research

- **Observe:** Using high definition microscopes, capture 4D sets of images of living embryos (Zebra Fish, Sea Urchin,...)
- **Transform:** Invent methods to go from images to symbolic representations (lineage trees, contours)
- Compare: Invent methods for efficient and meaningful comparisons

Industrialize

• Platform for high throughput execution of the processes



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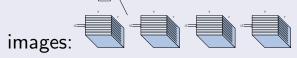
Some details

Gather observations

- Biologists place an embryo under microscope for a number of hours
- a stack of horizontal images of size x*y, separated in time by δt and space by δz are captured
- ullet a new stack is captured every ΔT
- Repeated for many individuals under different conditions

Output

A large set_of large files containing raw



A set of metadata describing the experiment



Some details

Reconstruct cell lineage tree

- Invent different algorithms to:
 - filter images (remove noise)
 - detect centers of cell nuclei ((x, y, z) position)
 - determine membrane contours (set of 3-D polygons)
 - determine nucleus contours (set of 3-D polygons)
 - identify mytosis (cell divisions)
 - track individual cell from step T_i to step T_{i+1} and build lineage tree
 - compare lineage trees, infer new results
- visualize reconstructions
- correct and annotate datasets



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The project
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Some figures

Image sizes: 512*512*8 to 1024*1024*8 pixels, $0.5\mu < \delta x, \delta y < 1.5\mu$, but soon: 2048*2048*24,

Number of images in stack: between 50 and 200,

Number of time steps: ΔT typically between 1 and 10 minutes, a few tens to a few hundreds of time intervals captured.

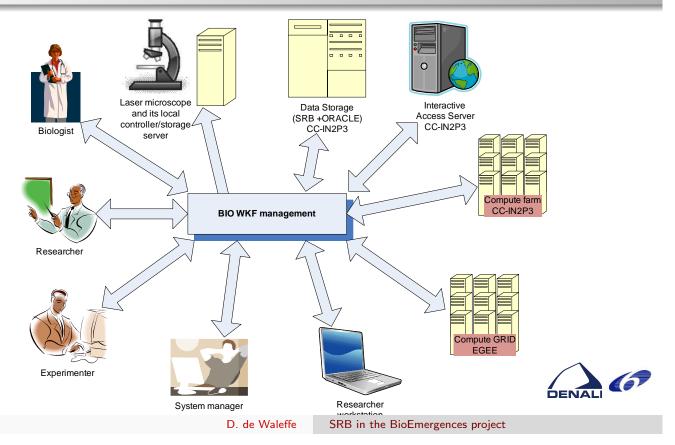
Raw data volumes: 50 to 60 Gigabytes of raw image files per experiment (size: 512) but will soon be 1/2 Terabytes with new microscope.

Number of cells: lineage trees contains several million cells.

Current storage used (SRB): in excess of 8 TB.

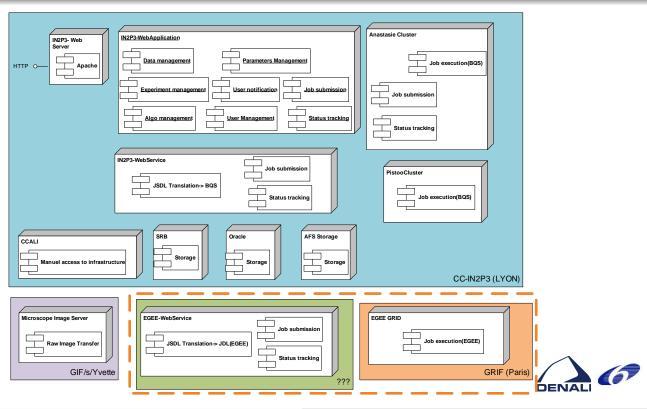


Context diagram

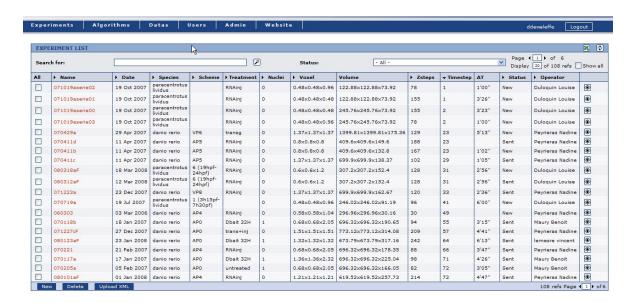


The project
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Deployment viewpoint



Application: experiment list





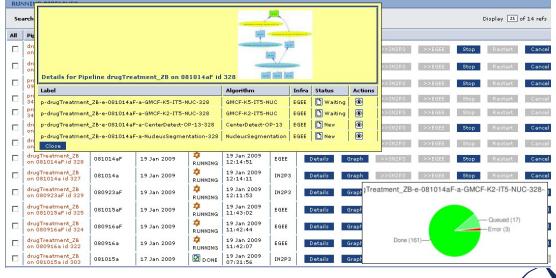
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The project
The architecture
SRB usage
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Application: processing pipelines

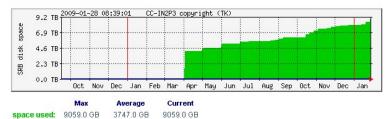
Details button: brings view below:



SRB usage: What

- Main repository: cc-in2p3 Lyon.
- Raw data storage. Data captured in Paris, format standardized, then copied to SRB. Srsync is used.
- Derived data.
 - Kept for reuse in further processing.
 - Data history kept in application's DB.
- Some additionnal files (movies of reconstructions,...)
- Stores algorithms (scripts, sources, builds procedures, executables)

Yearly Graph (1 Day Average)



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SRB usage: how

- Used in identical manner from both farms.
- Mostly used as a file system
- Command line: Smkdir, Scd, Sls, Sput, Sget, Srm, Srsync
- Jargon library used for
 - displaying raw data file lists
 - Streaming movies [todo]
- Not used in the project:
 - user meta data
 - web based browser



SRB usage: issues

- Slow access when doing operations requiring catalog access (e.g. Sls −1).
- Bizarre error messages, or no exit codes (makes it difficult in scripts)
- Locking bugs (multiple e.g. Smkdir X) which impact the whole group!
- deleted stuff is not always fully deleted



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The project
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Why iRODS?

- Post-processing on ingestion. Could trigger raw data format changes on upload.
- Workflows. Probably redundant with what we already have or can have.
- Looks like iRODS would be a good fit but:
 - we have no extra budget in current project
 - team is currently looking at using ROOT as storage framework
 - taking advantage of iRODS imply large re-architecting effort
- sources available.



Why iRODS, some risks?

- maintainability of complex rule base?
 - rule syntax (one liners, readability, choices of operators, comments?)

```
myRule|foo==1|action1(...);action2(...);...|action3(...);action4(...);...
```

Why not sligthly more verbose

```
rule myRule { //this rule is triggered when foo and does bar
  when ( foo == 1)
  do {
     /* watch that this action has side-effects */
     action1(...);
     action2(...);...
}
  on failure {
     action3(...);
     action4(...);...
}
```

• can I define new microServices as complex jobs (e.g submit job(s) to farm) without going to C programming?

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Conclusion

- BioEmergences has complex distributed data/processing needs
- Could make use of iRODS if risks are shown to be a non issue



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Questions?



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