

Données d'imagerie en microscopie: quelques exemples et besoins en visualisation

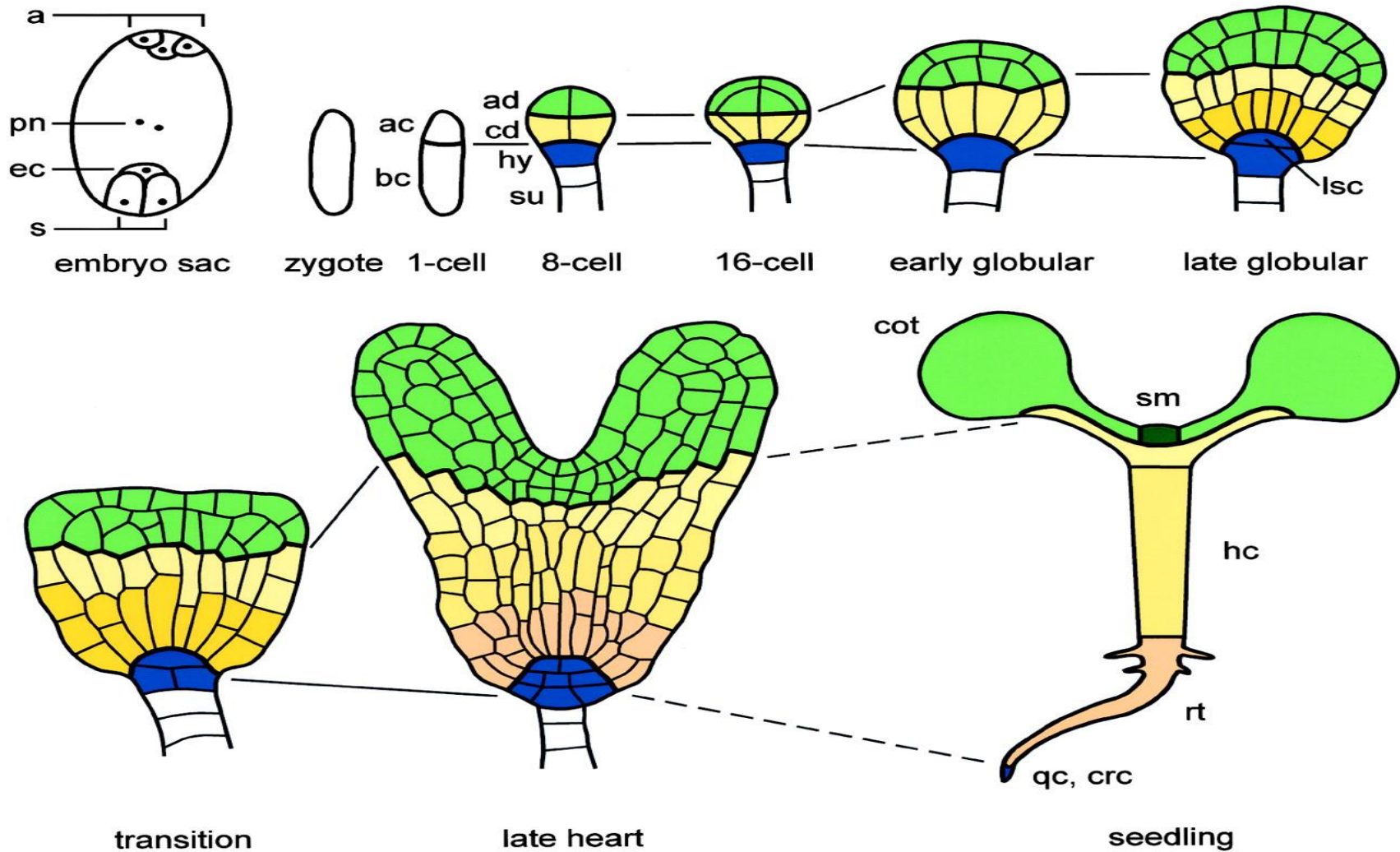
A. Trubuil
INRA/MaIAGE

1. **Besoin d'IHM pour le lignage cellulaire**

2. Besoin de navigation pour des observations 2D +T ou 3D+T

✓ **Movies** have been removed from the pdf file due to file size limitations

From a single cell to an embryo

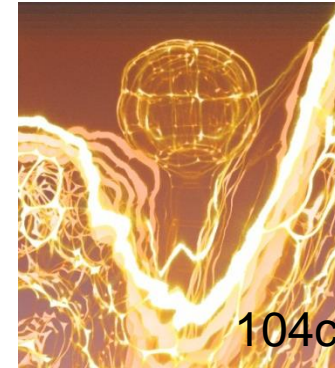
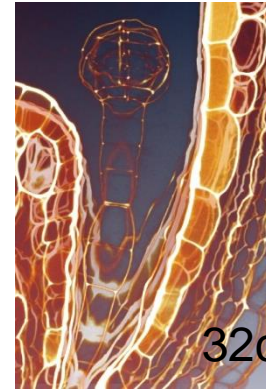
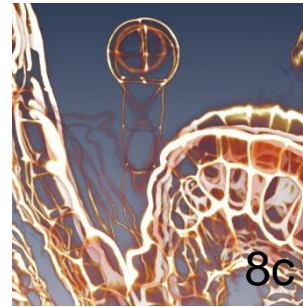


Laux T., et al., *Plant Cell*, **16**, 190 – 202 (2004)

Q1: what is the sequence of events?



time ~~X~~ lapse



Several embryos at different stages of development




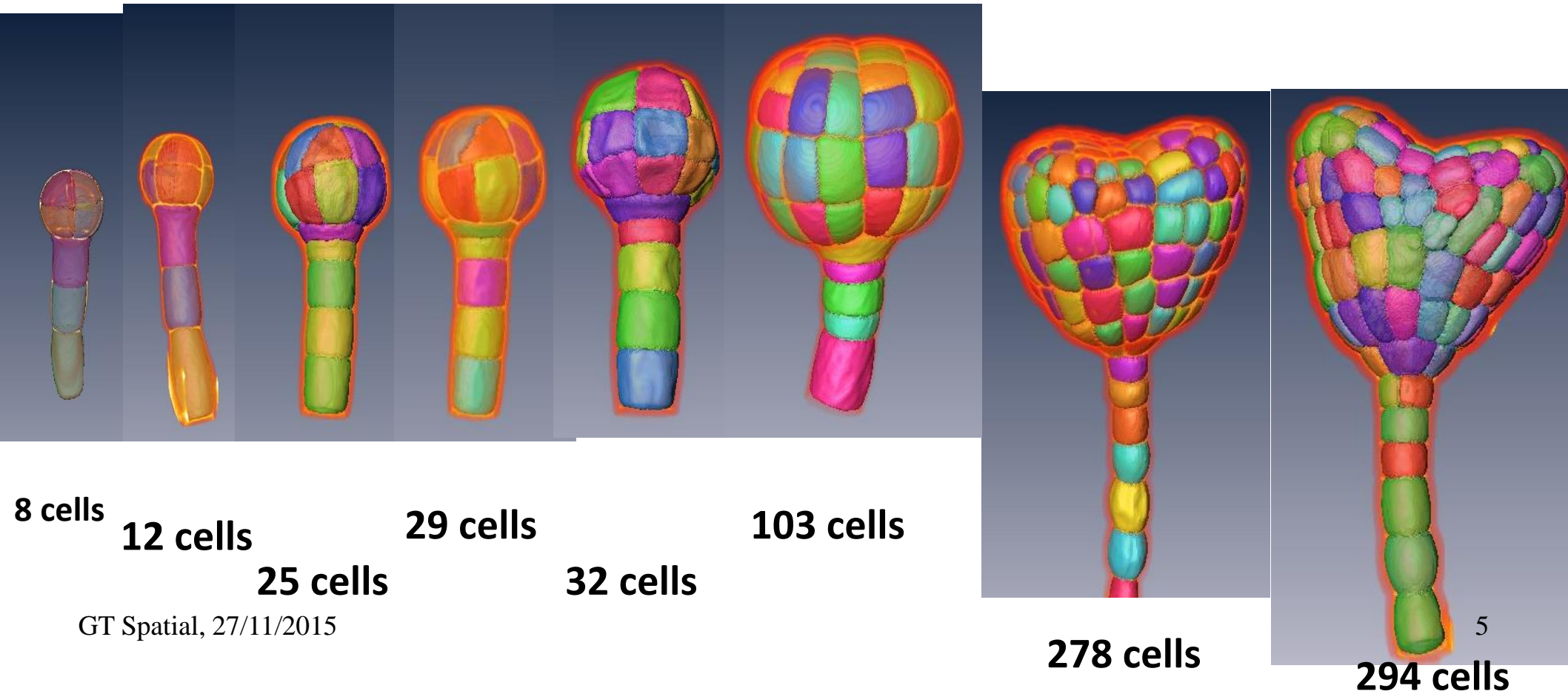
Characterize events: cells, walls, [stiffness],...

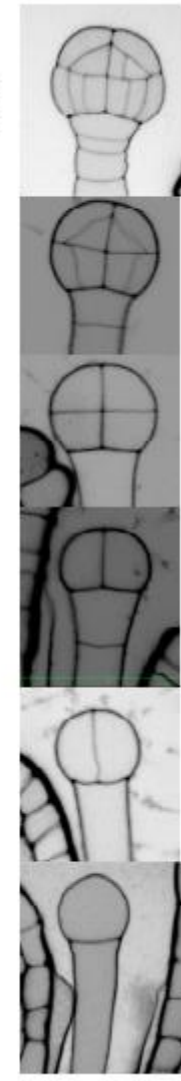
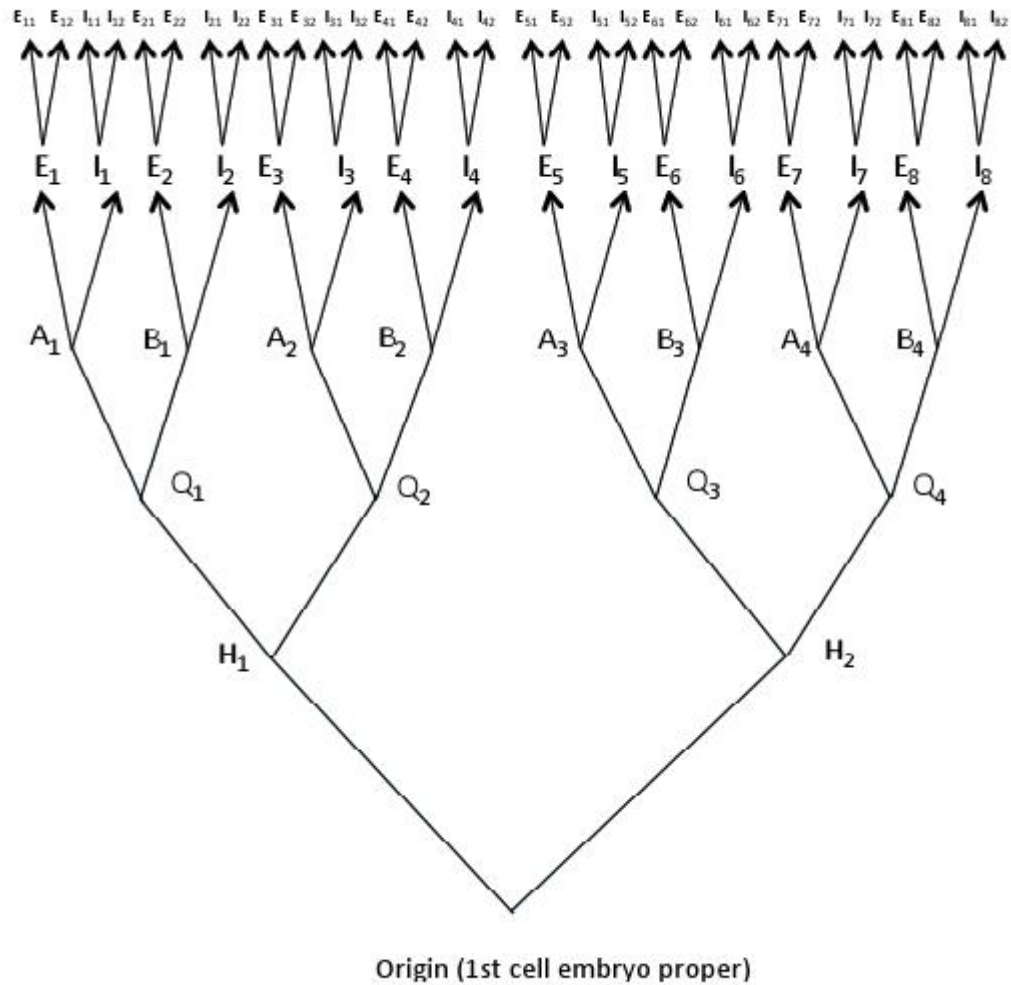
1. 3D segmentation (number of cells, volumes,...)
2. Reconstruction of the dynamics
3. Mechanical properties ?

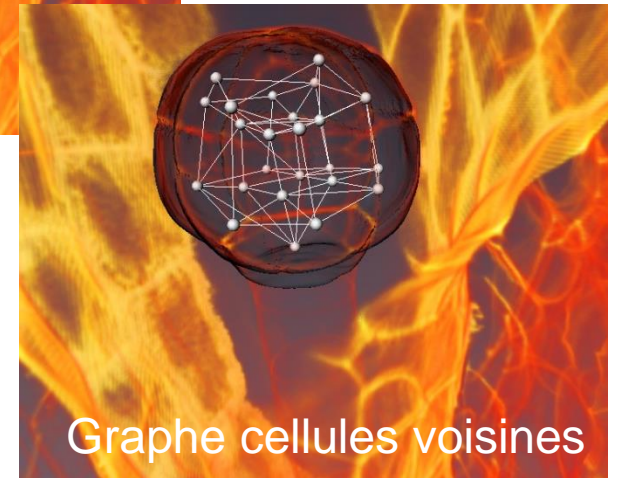
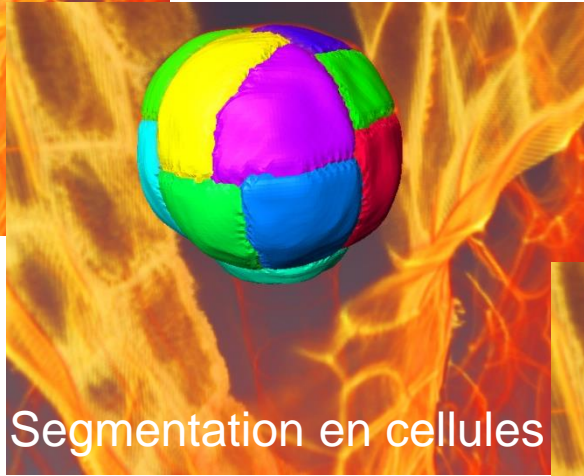
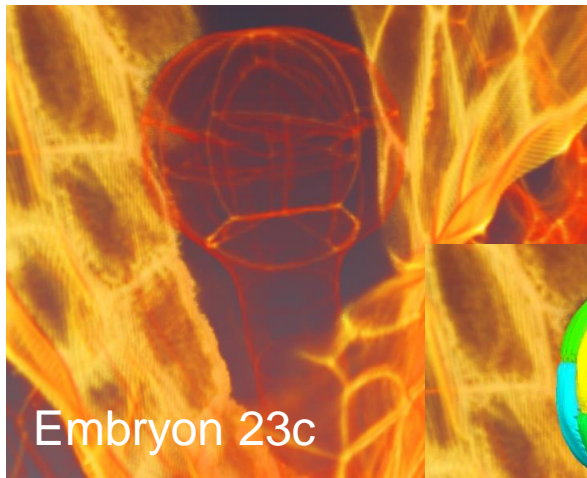
Q1: 3D segmentation

- Embryo not alone
- Observation of a continuous process (walls under construction)
- 3D
- Complexity (from 1-300 cells)
- Artifact due to experiment
- Validation

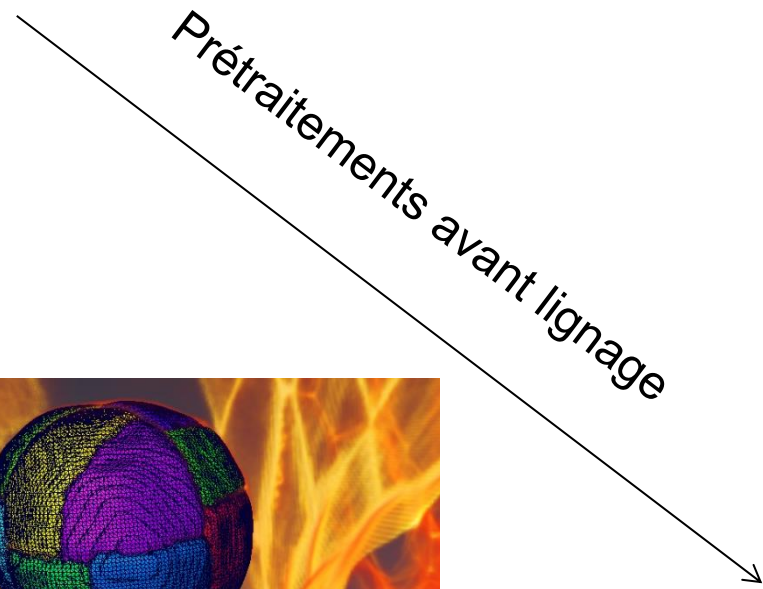
 pipeline (C, Matlab, Avizo)







Prétraitements avant lignage



Matlab, Avizo

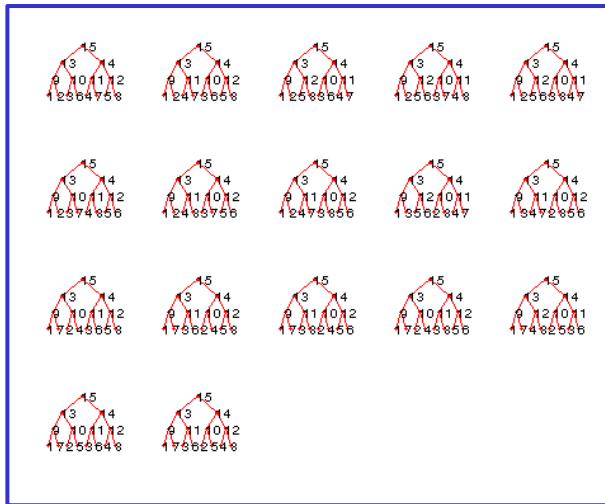
Q1: Cell lineage, constrained trees enumeration (5)

Input : neighbour graph of cells

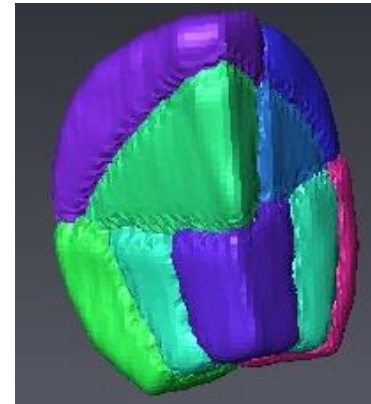
Constraints :

1. daughter nodes are neighbours,
2. maximal depth difference of branches

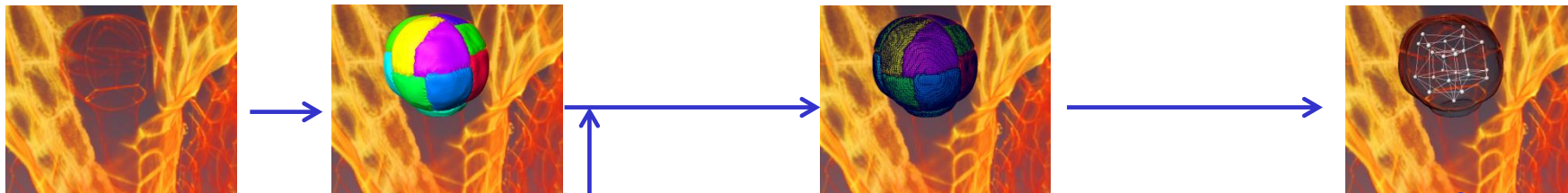
Algorithm : recursive



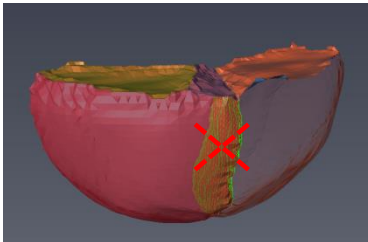
8 cells: 17 trees



Quarter of 28 cells embryo



Merge pairs



$$\phi_1(d) \stackrel{\text{def}}{=} \oint_{\partial d} \delta(s) ds$$

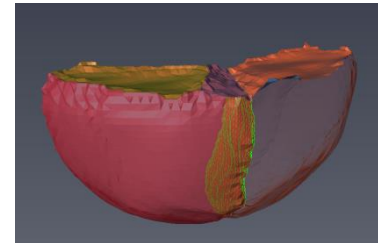
Interactive constraints:
 a ----- b or a ~~-----~~ b

- Propagate constraints
- Feed DB of pairs

Maximal Matchings (MM)

Maximal cliques in the complement dual graph (Uno et al)

Evaluate and Filter MM



Visualize partial lineages

- Display shared pairs
- Display non shared pairs
- Display new pairs
- Display known pairs

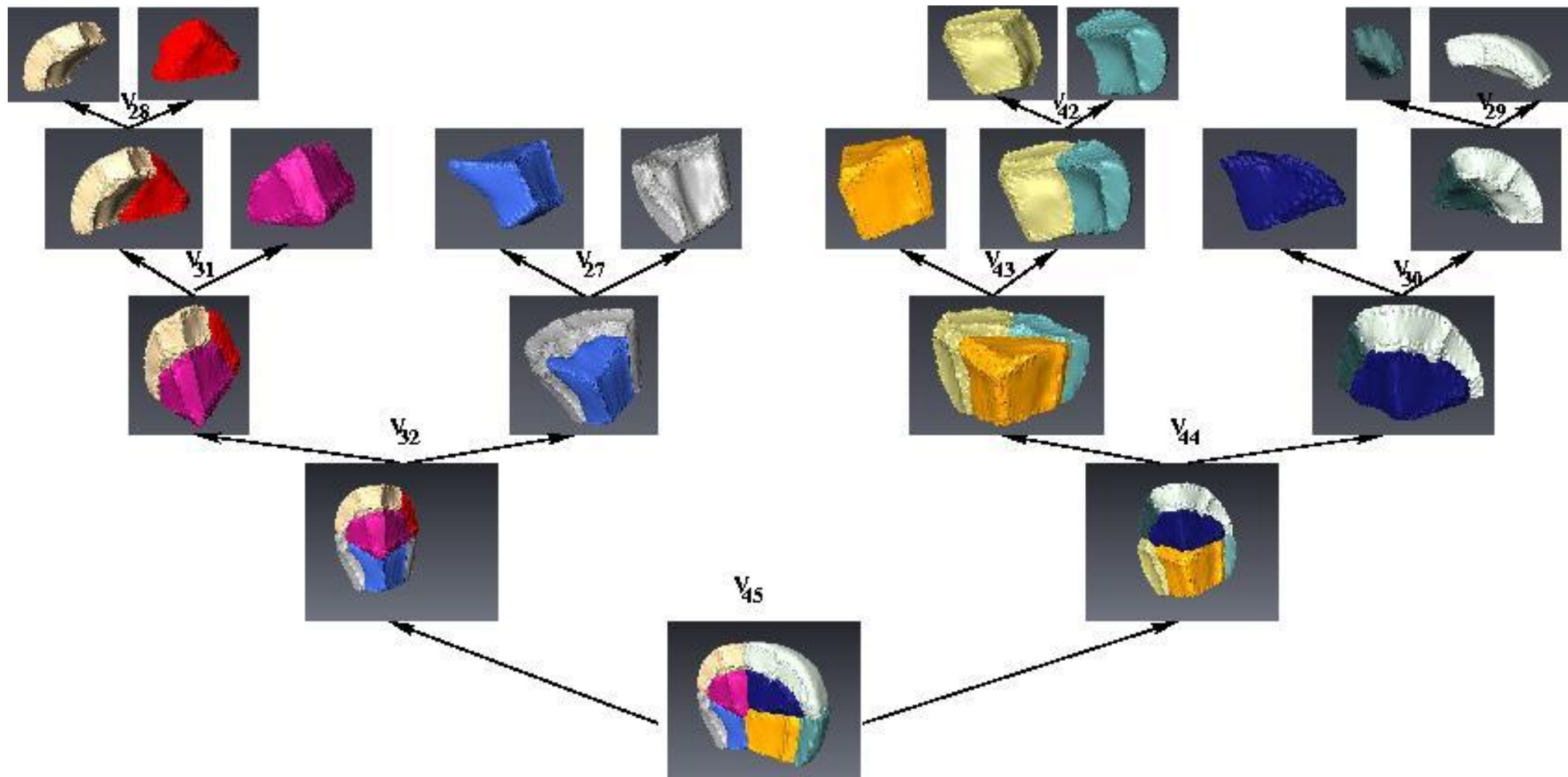
Next round?

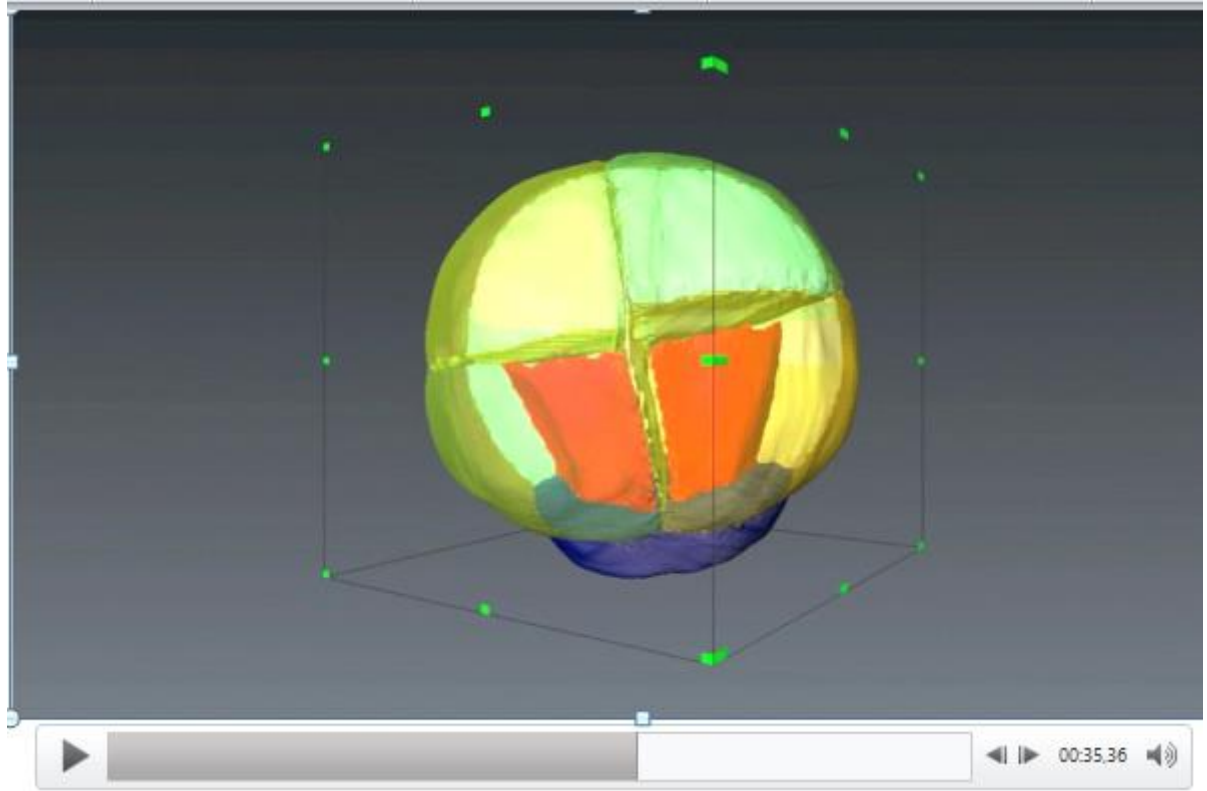
Feed DB of matchings

Feed DB of pairs

Visualize lineages

Lignage cellulaire (embryon 23 cellules [moitié])





1. Besoin d'IHM pour le lignage cellulaire

2. Besoin de navigation pour des observations 2D +T ou 3D+T



**Exploration of movies and Visualization of
content :
Examples**

- ✓ A collection of 2D+T or 3D+T movies
 - ❑ Biofilms (GreenSwimmers project)
 - ❑ Lipid Droplets
 - ❑ Vesicles

- ✓ Detect, classify, show events

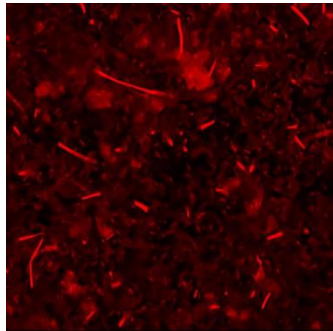
- ✓ Conclusion

EXAMPLE 1 : VIDEOS FROM BACTERIA SWIMMING INSIDE BIOFILMS

125 strains x 2-5 videos for 30 seconds

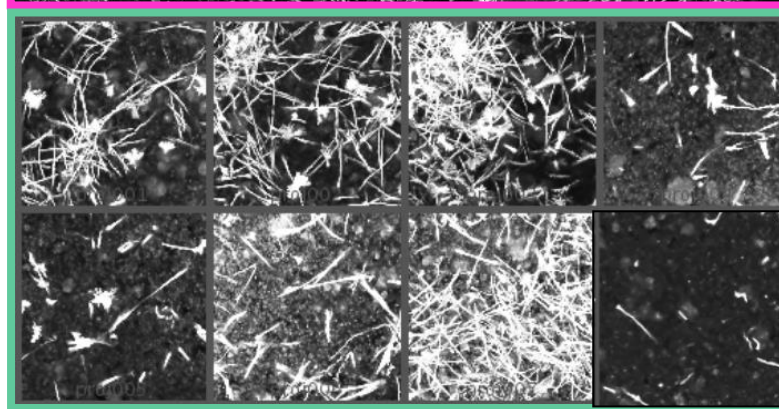
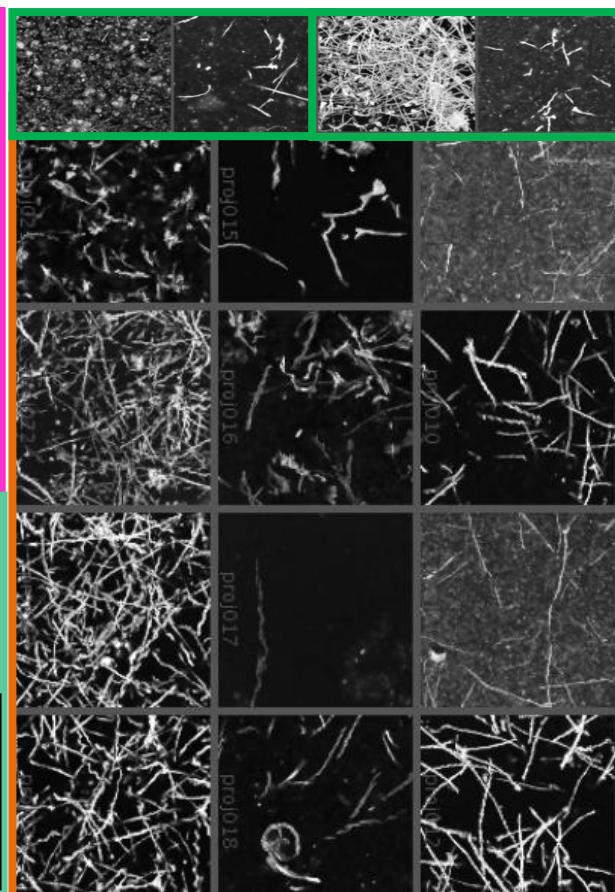
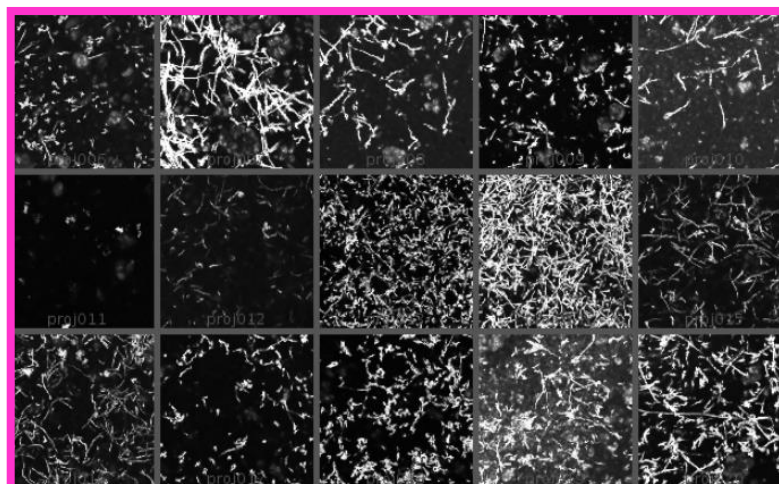
- Speed of swimmers
- Persistence of swimmers
- Surface coverage of swimmers

Trajectories projection for 30 seconds



B. polymyxa

B. subtilis

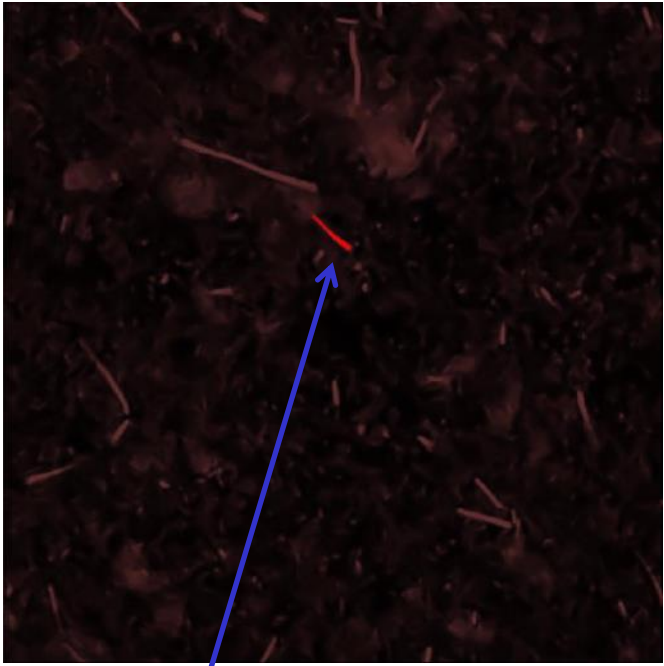


B. sphaericus

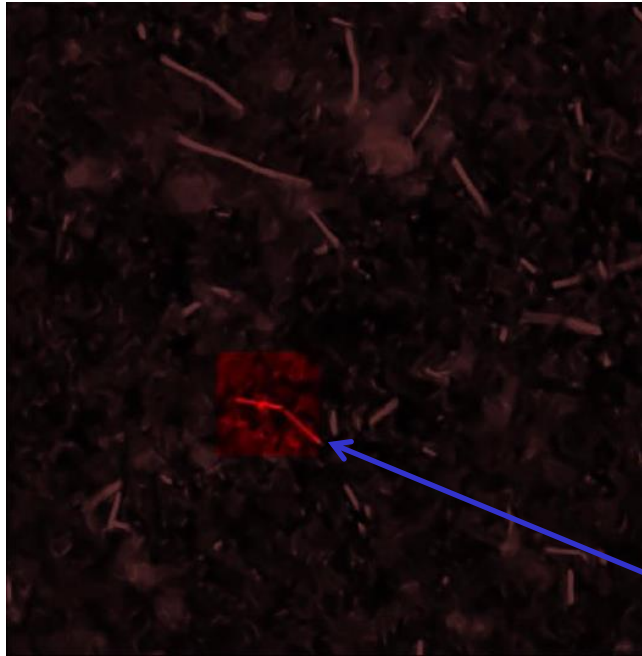
B. cereus

Species	# of strains
<i>B. polymyxa</i>	17
<i>B. subtilis</i>	18
<i>B. licheniformis</i>	31
<i>B. pumilus</i>	19
<i>B. sphaericus</i>	4
<i>B. megaterium</i>	3
<i>B. cereus</i>	30
<i>B. mycoides</i>	3
Total	125

SHOW EVENTS

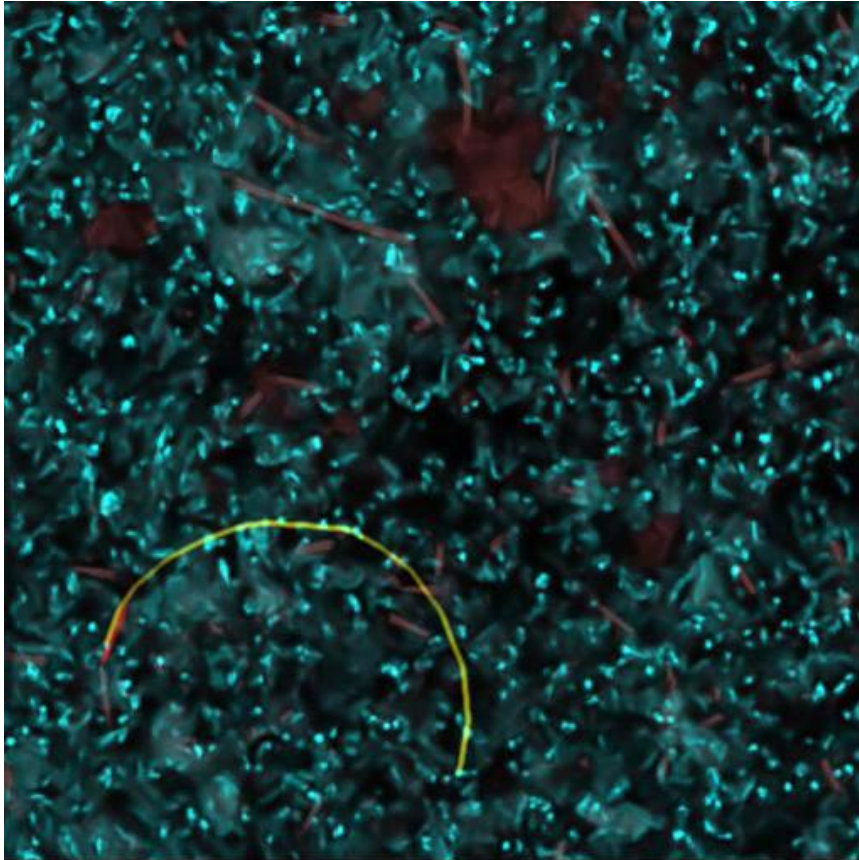


DNA pocket attraction

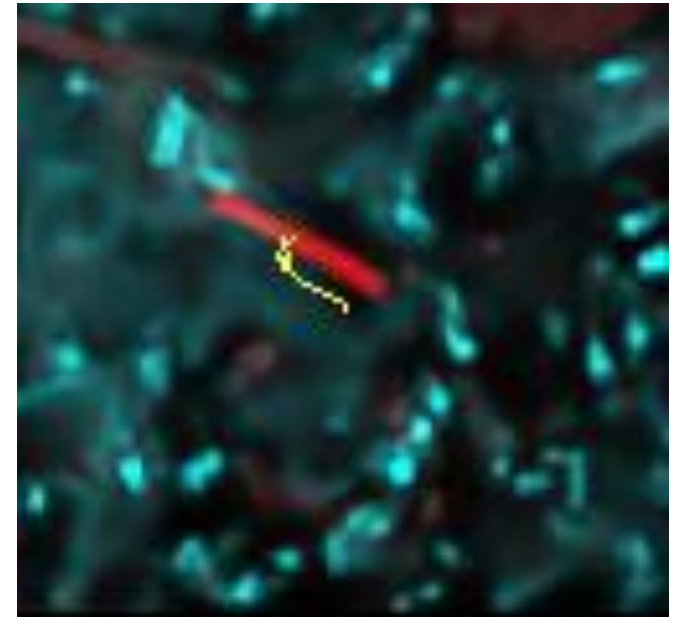


Circular motion?

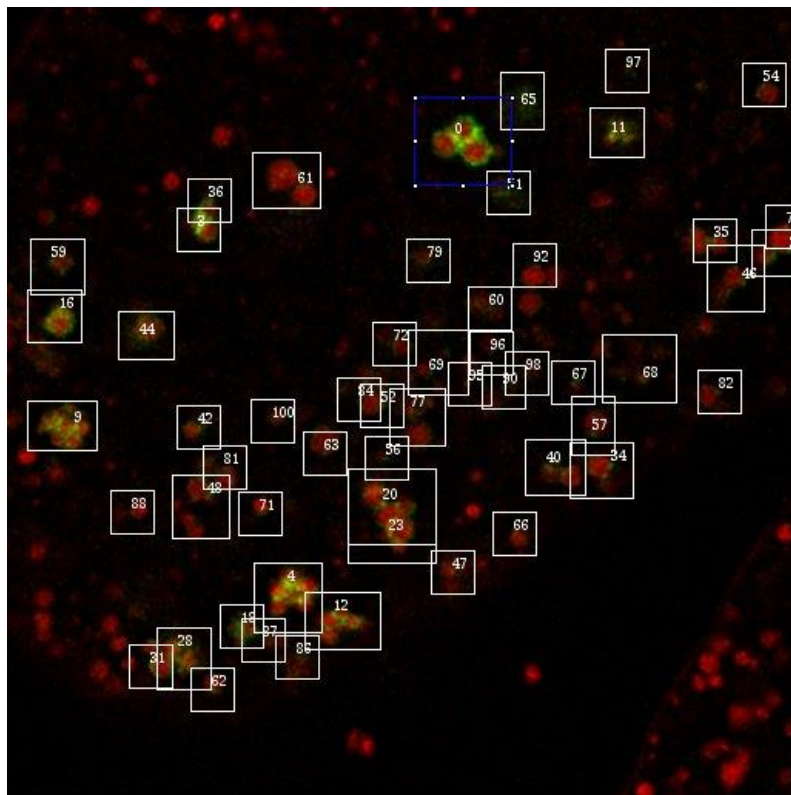
Sphaericus 10C3



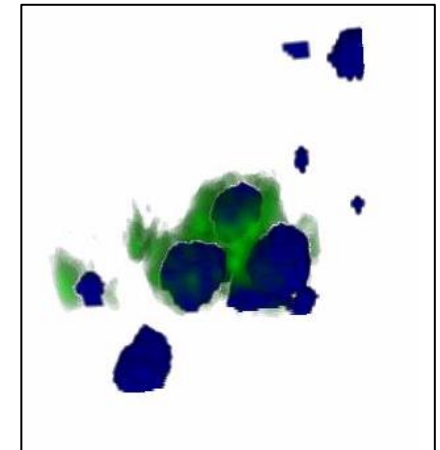
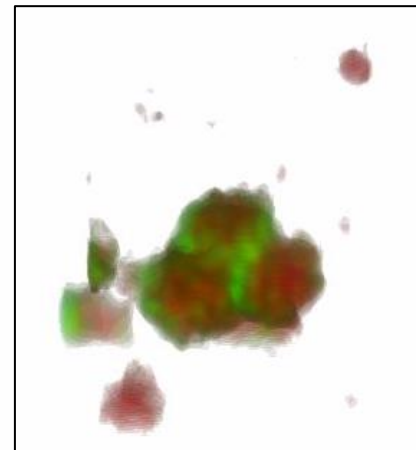
Circular motion



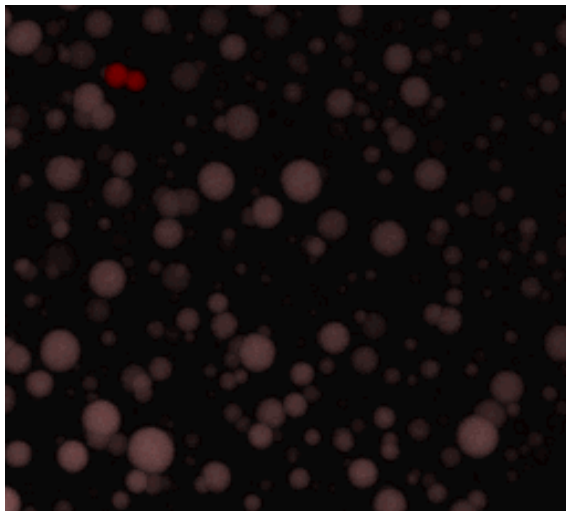
**DNA pocket attraction
or biofilm obstacle**



- **Several stages along the development**
- **Several genotypes**
- **Several instances from one or several embryos**



Selection and 3D visualization of LDs



Fusion of two LDs is highlighted

With F. Deslandes

CONCLUSIONS

1. Many advanced visualization for large audience
 - sport (see TV)
 - security
 - cinema
 - biomedical?
2. Not so much for microscopy(to check)
 - Identify class of objects that should be visualized
 - How to put forward spatio-temporal interactions
 - Video summarization for cell biology

THANK YOU

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