

# Spatial organization of neuroepithelial cells during embryo development

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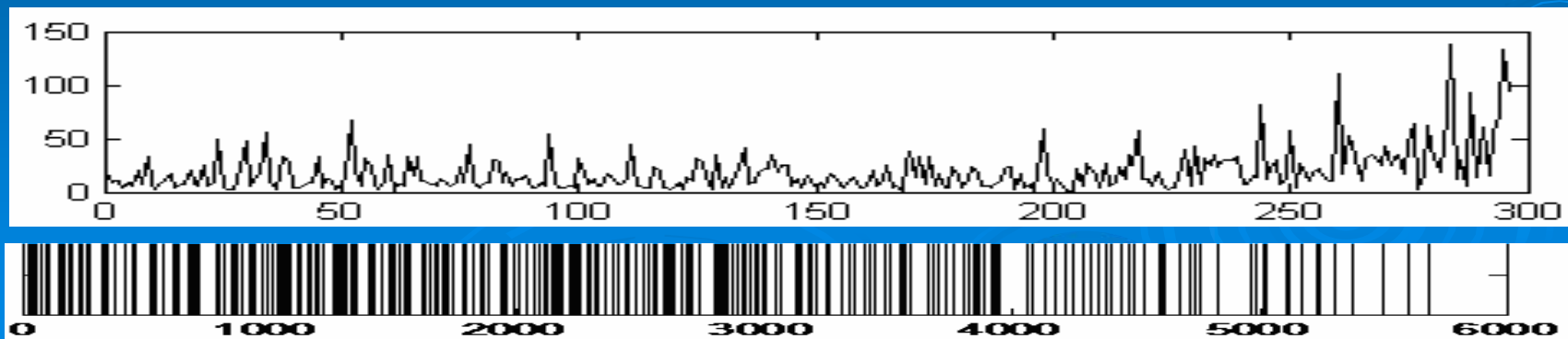
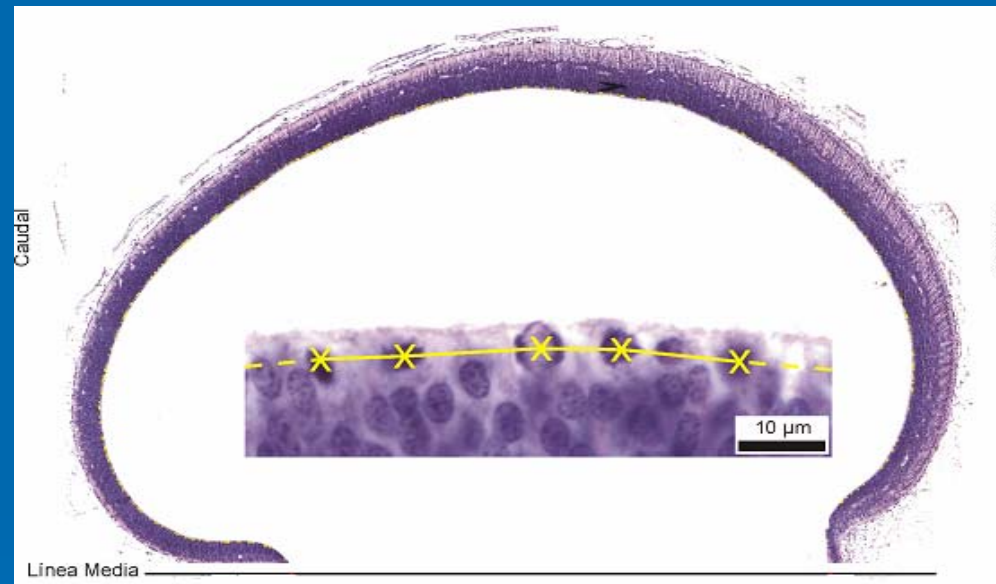
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# Embryonic development

- A few **simple** cell behaviors  
... **organized in time** and **organized in space**
- One these behaviors is proliferation
- Proliferation not only increases volume but may also generate complex shapes

# Biological model

- ❖ Chicken central nervous system (Optic Tectum)

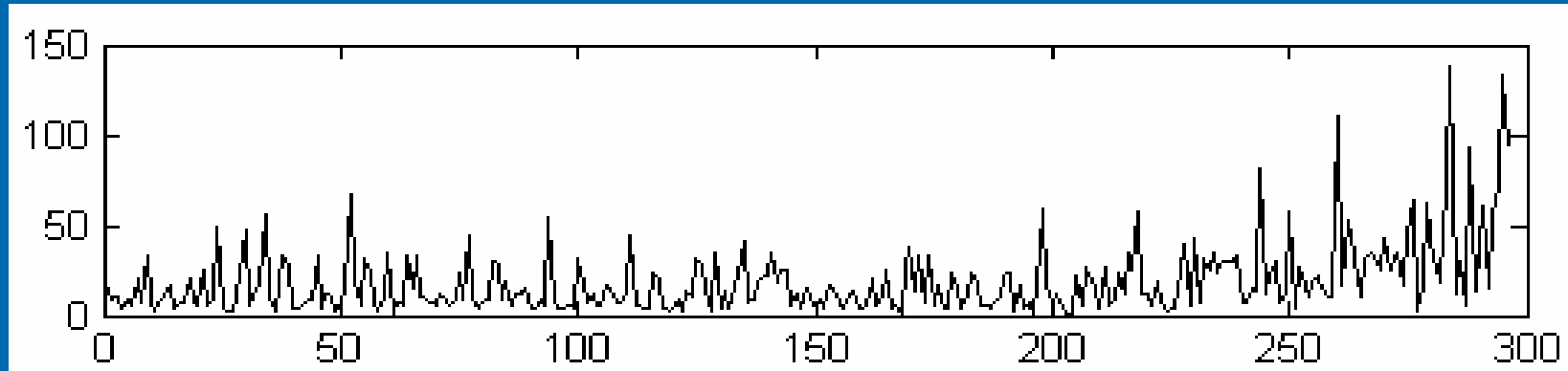


# What are we searching in these records?

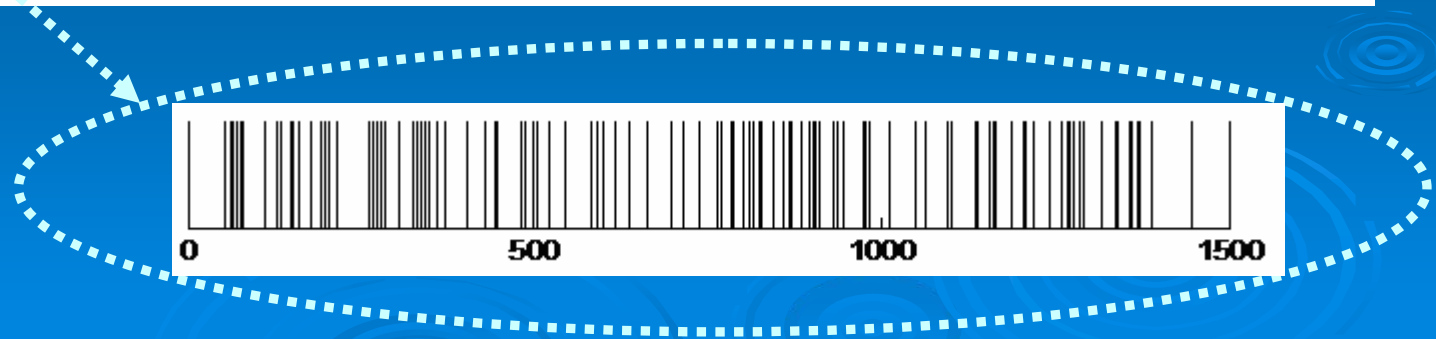
- In general:
  - assess organization in space
- In particular:
  - Are cells distributed just at random?
  - Do correlation exist between the intervals?
  - What kind of correlations?
  - Characterize non-stationarity
- ✓ **Correlations tells us about communications between cells**

# Data

*intervals*

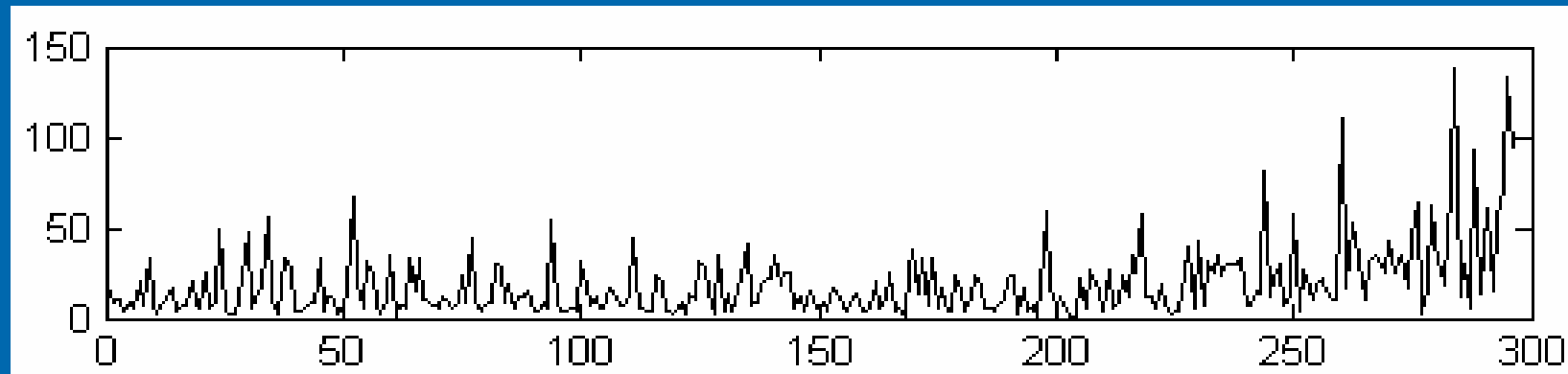


*Location of cells*



# Data

*intervals*



*Location of cells*



Some of the characteristics are:

- around 300 samples long
- non-stationary
- PDF of intervals is far from being Gaussian

# Methodology

- Stochastic point process
- **Fractal** characteristics assessed by estimating Hurst index, scaling index



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“Size”



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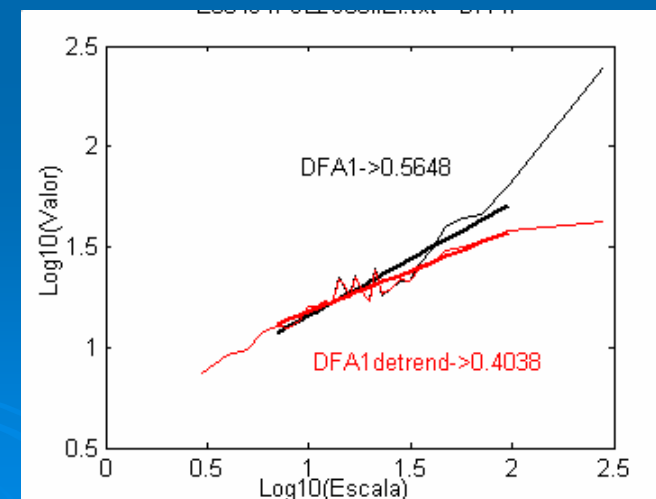
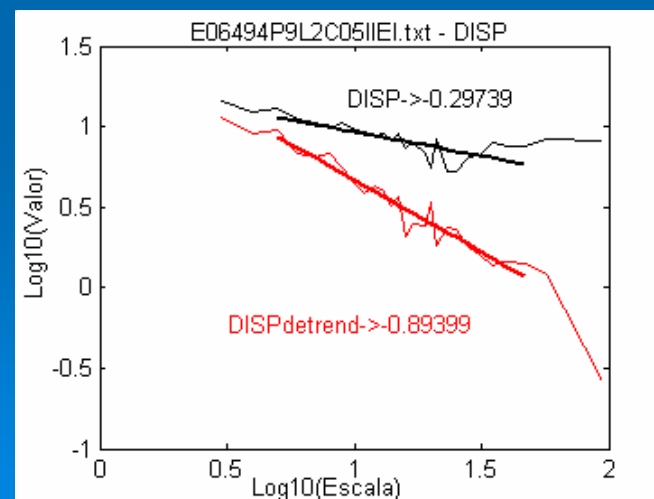
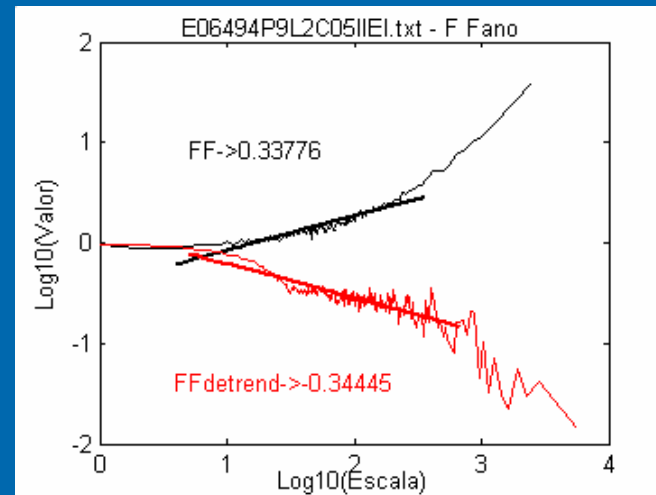
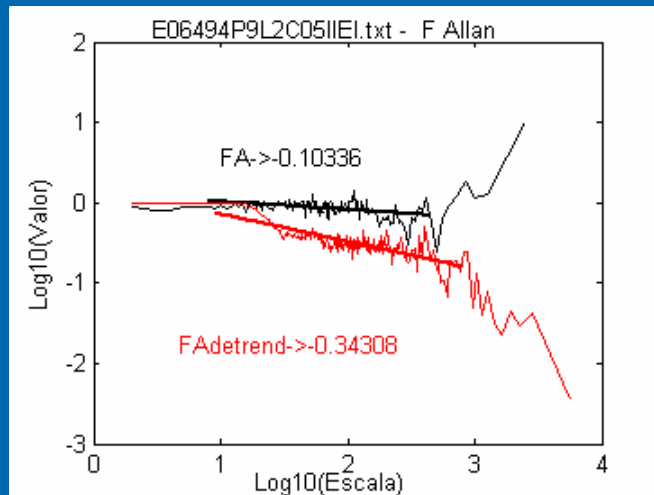
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- **Fractal** characteristics assessed by estimating Hurst index, scaling index



—  
“Size”

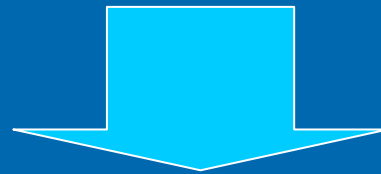
- ✓ If  $f(\text{size})$  follows  $\approx a^H$   $\Rightarrow$   $H$  is the fractal exponent
- ✓ *Trend estimation and removal before processing*

# Results



# Interpretation of results

- Negative correlations in a wide range of scales
- Global trend



- ✓ *proliferating cells communicate with each other*
- ✓ *their interaction is inhibitory*
- ✓ *Effect of “external control” generates non-stationarity*

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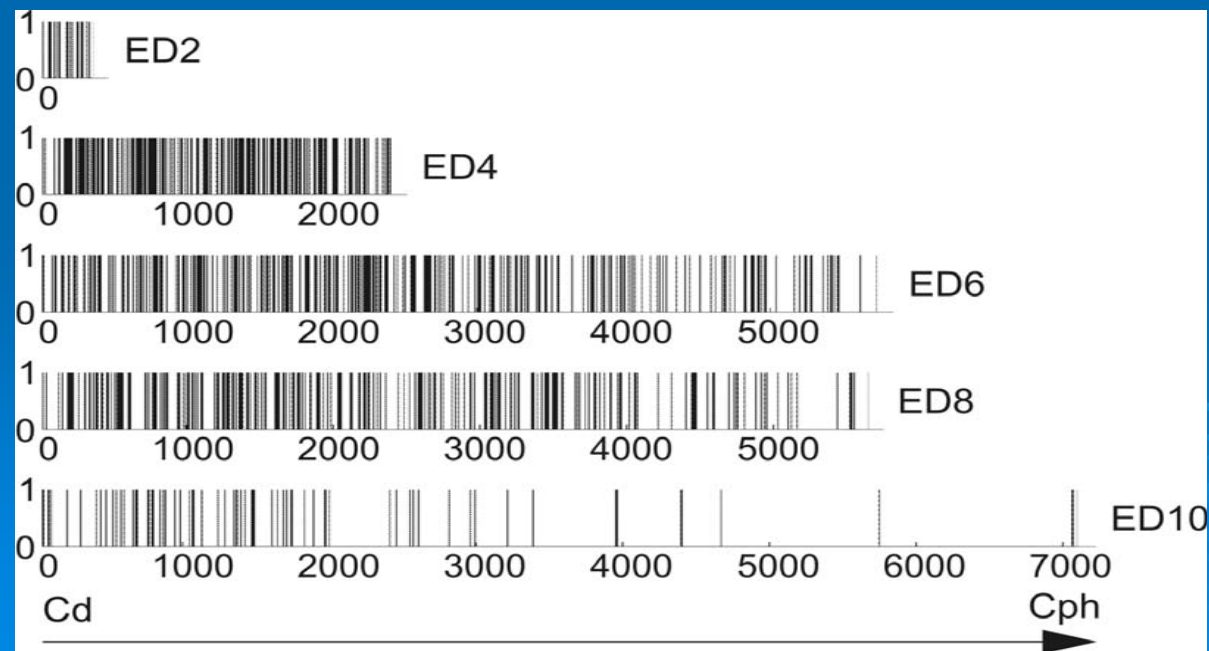
This is cell by cell approach is much more descriptive approach than the averaged data generally used in Developmental Biology

# Future work

- Alternative methods for studying this data set
- Include organization in time

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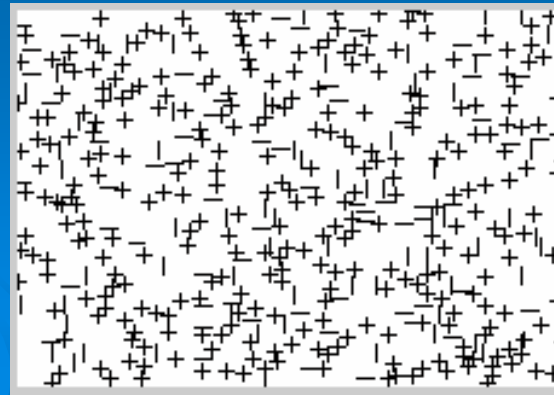
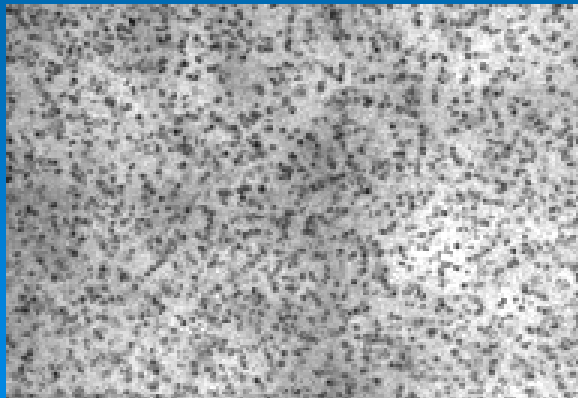


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Thank you!



