



Revisiting PIN1 polarity interpretation at the SAM based on automated multi-level reconstruction of PIN1 polarities from confocal images

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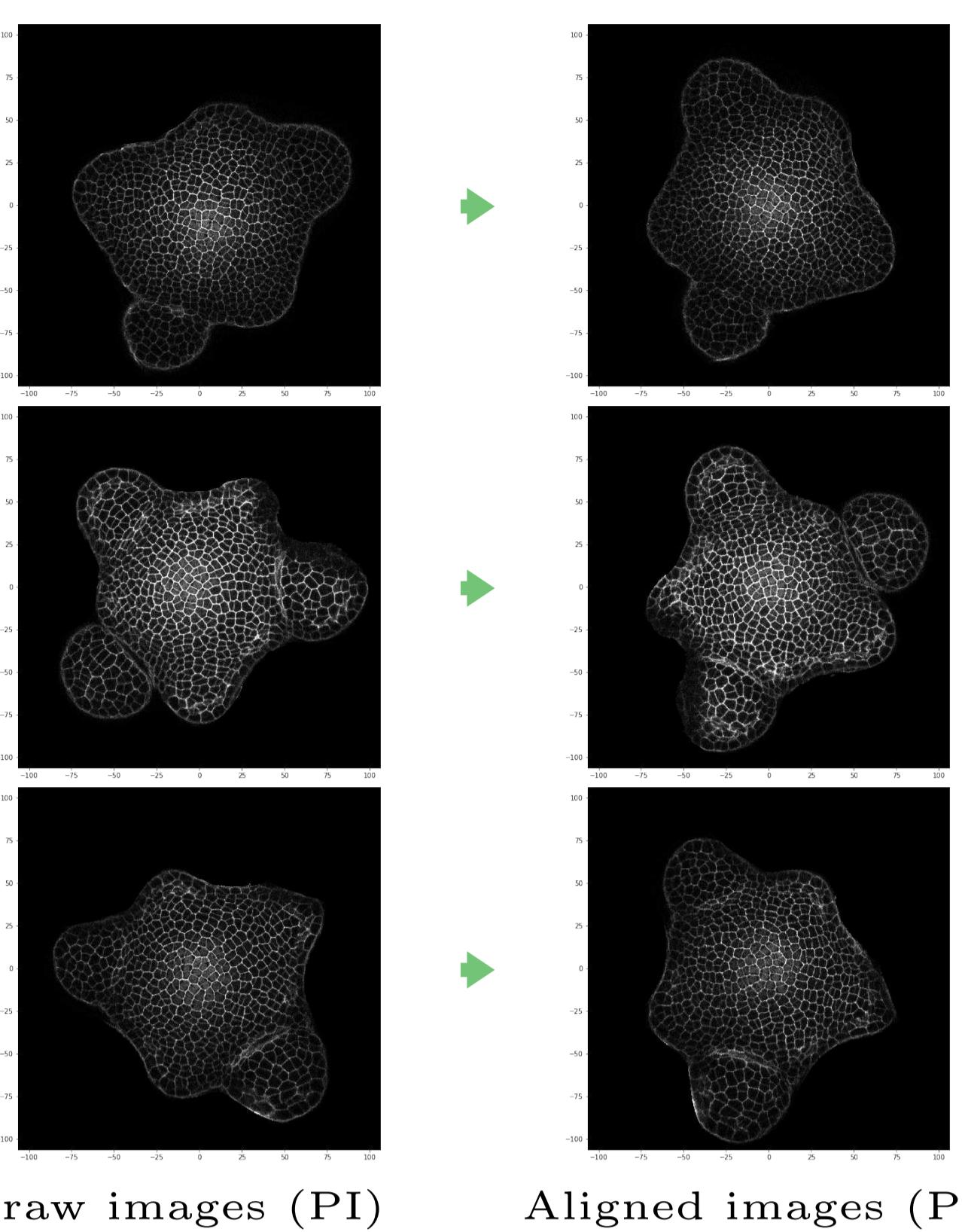
Revisiting PIN1 polarity interpretation at the SAM based on automated multi-level reconstruction of PIN1 polarities from confocal images

Landry Duguet¹, Guillaume Cerutti¹, Carlos Galvan-Ampudia¹, Etienne Farcot², Teva Vernoux¹, and Christophe Godin¹

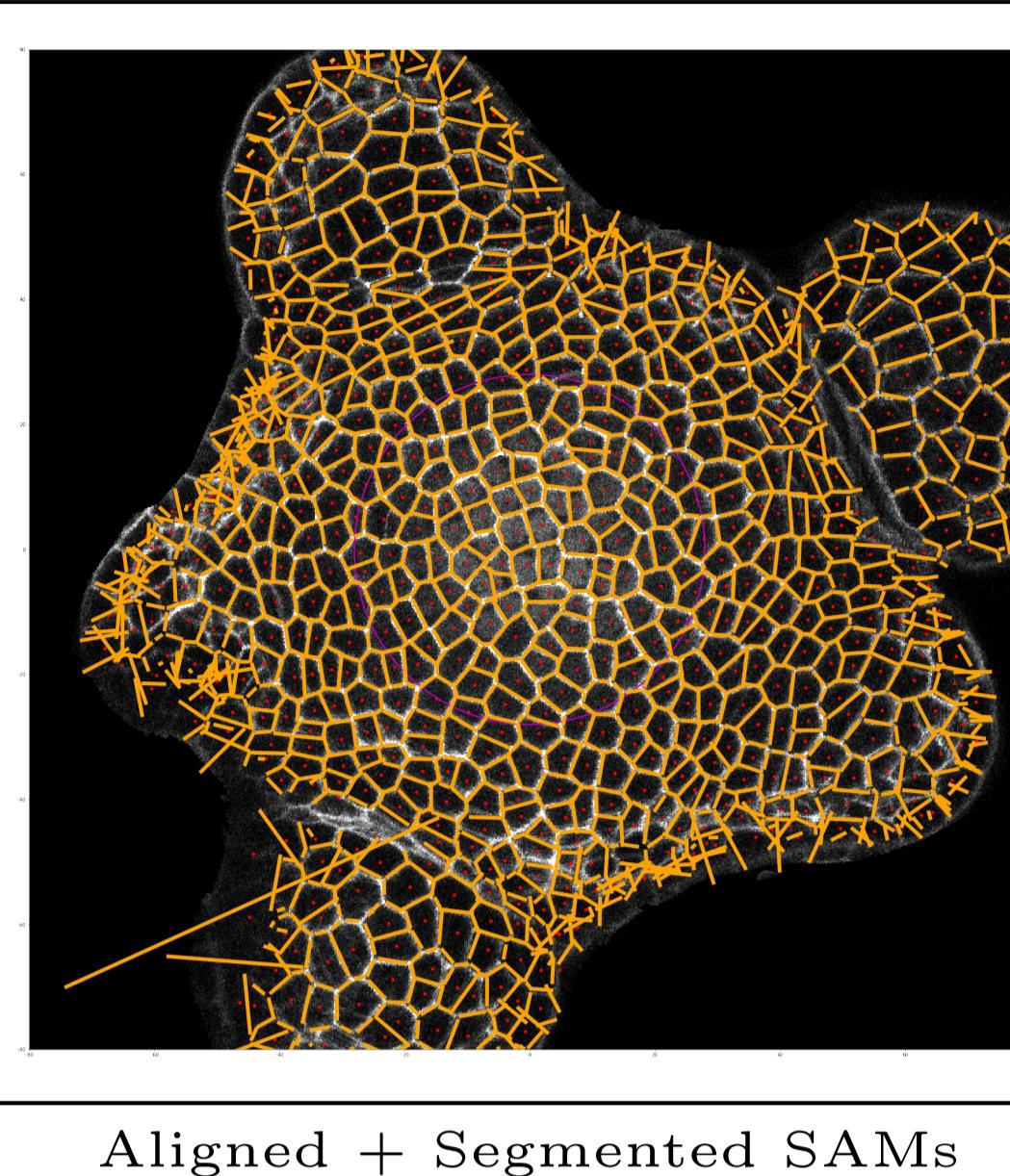
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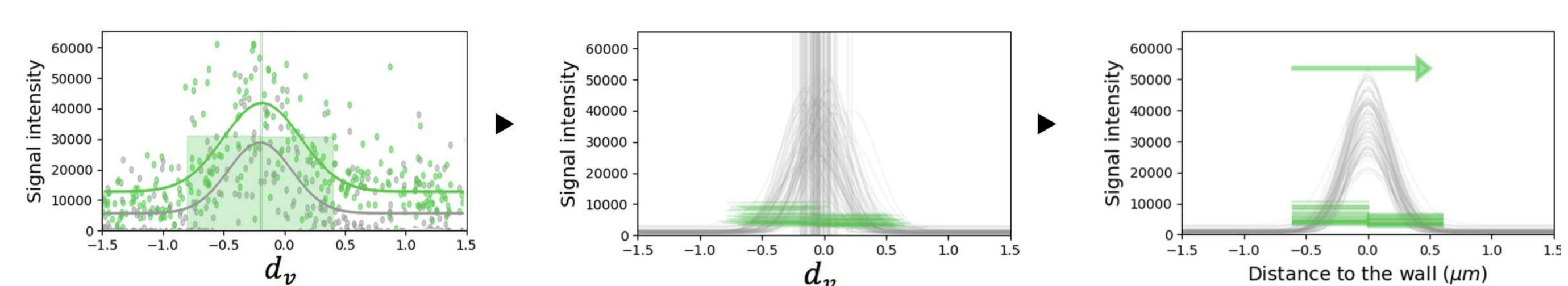
SAM alignment based on *CLV3* and *qDII* markers



Alignment & Segmentation

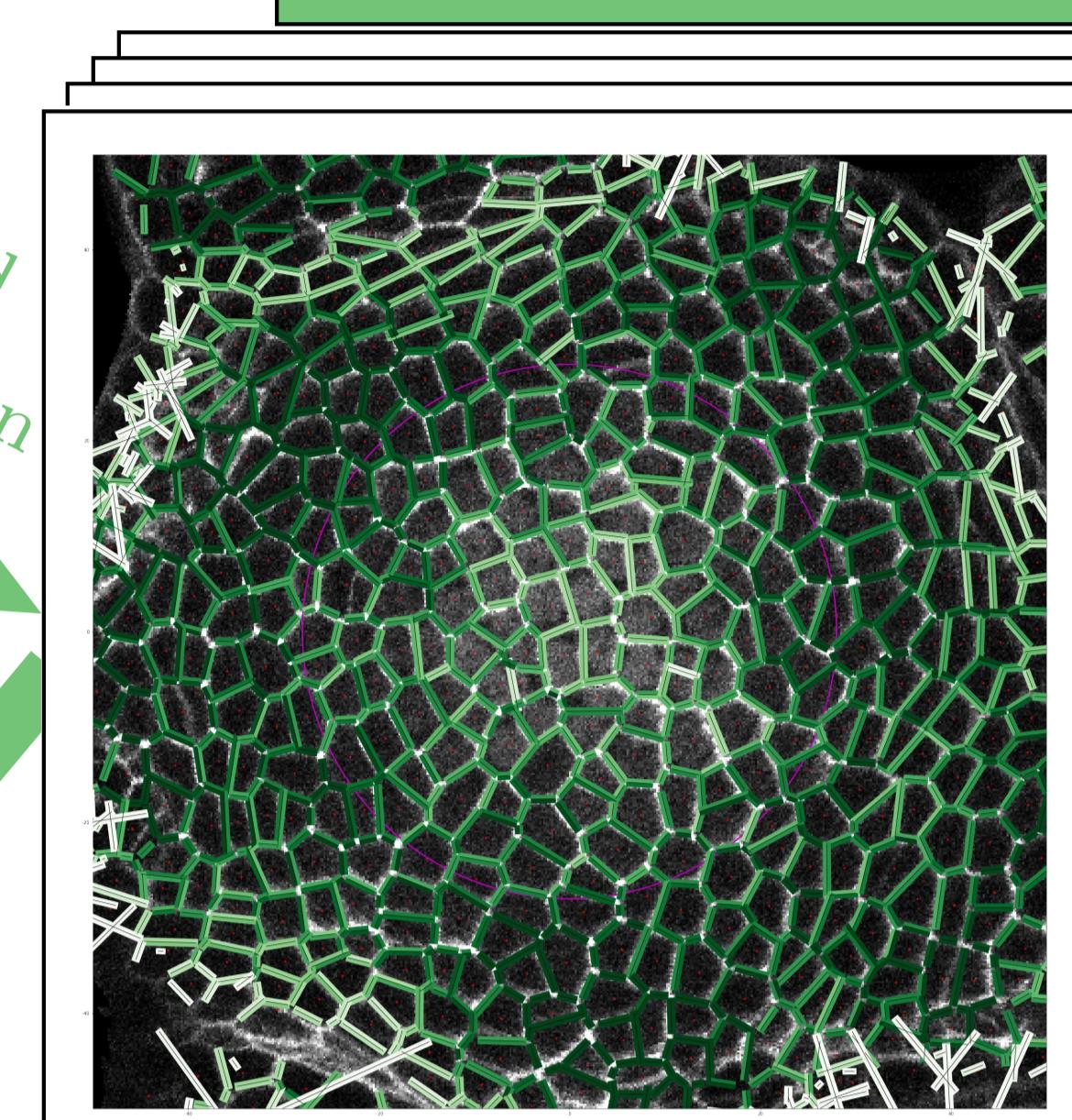


Aligned + Segmented SAMs



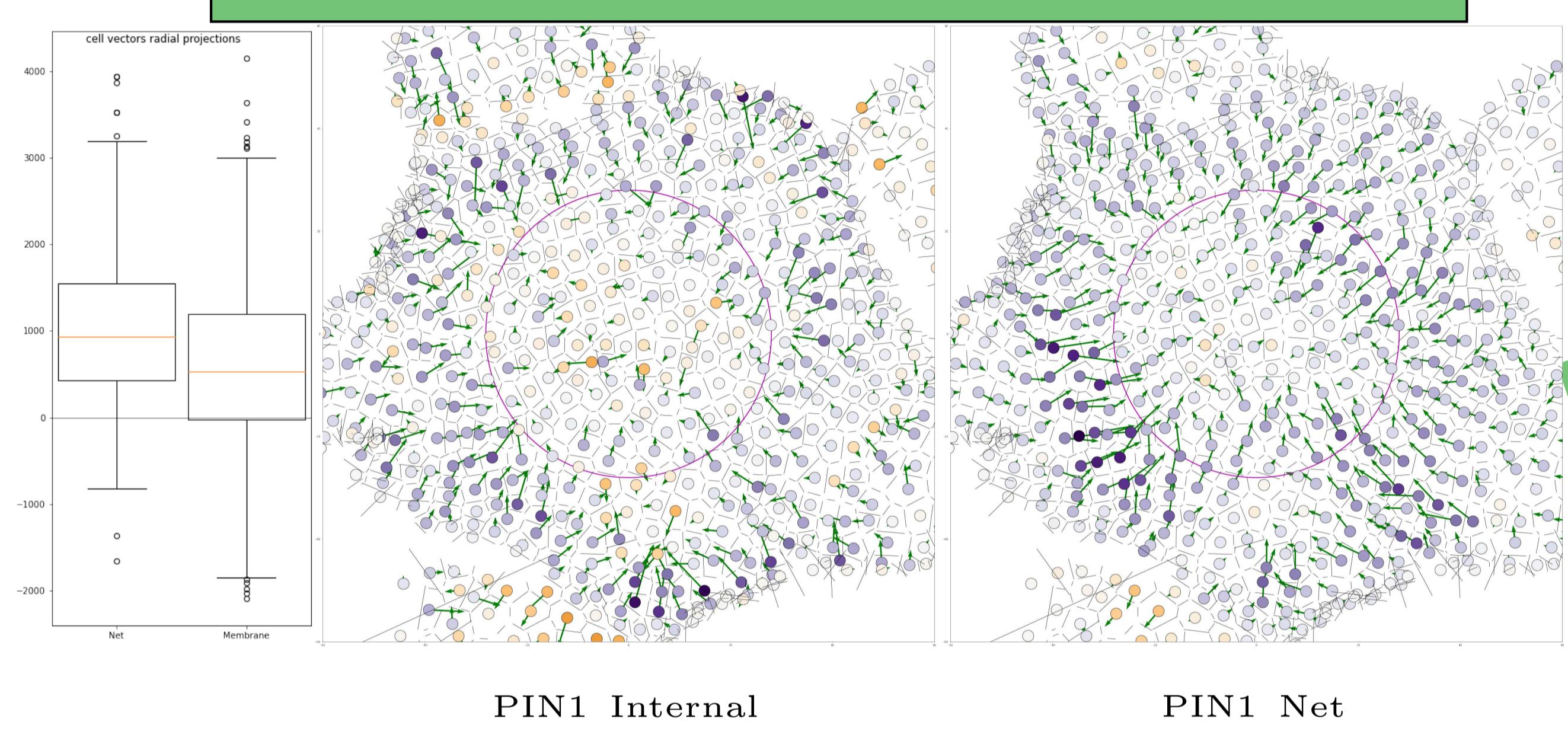
Automated PIN1 extraction method [Galvan-Ampudia et al. (2020)] based on pPIN1::PIN1-GFP and PI (cell wall) markers. The method infers a PIN1 level to both sides of the cell wall.

PIN1 at membranes



- P_{ij} : PIN1 level in the membrane of cell i facing cell j .
- PIN1 total wall values $P_{ij}^T = P_{ij} + P_{ji}$.
- PIN1 net wall values $\Delta P_{ij} = P_{ij} - P_{ji}$.
- PIN1 net wall vectors $\Delta \mathbf{P}_{ij} = \Delta P_{ij} \hat{\mathbf{n}}_{ij}$

PIN1 cell vectors

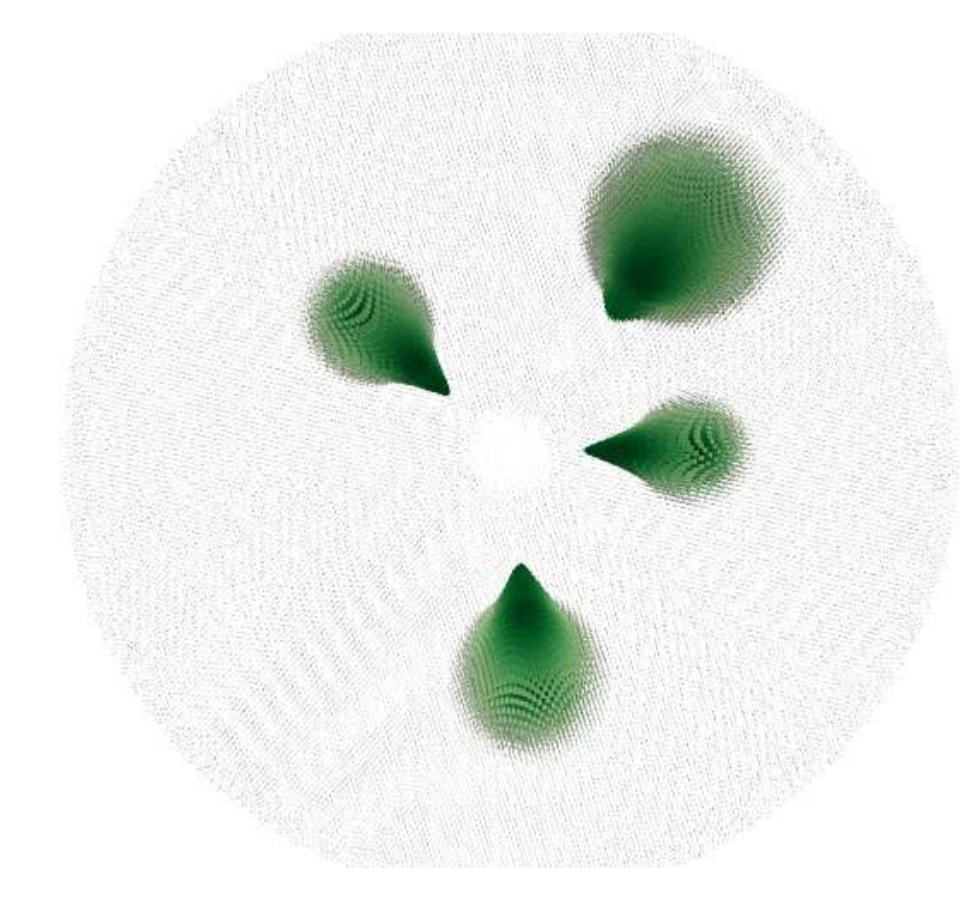
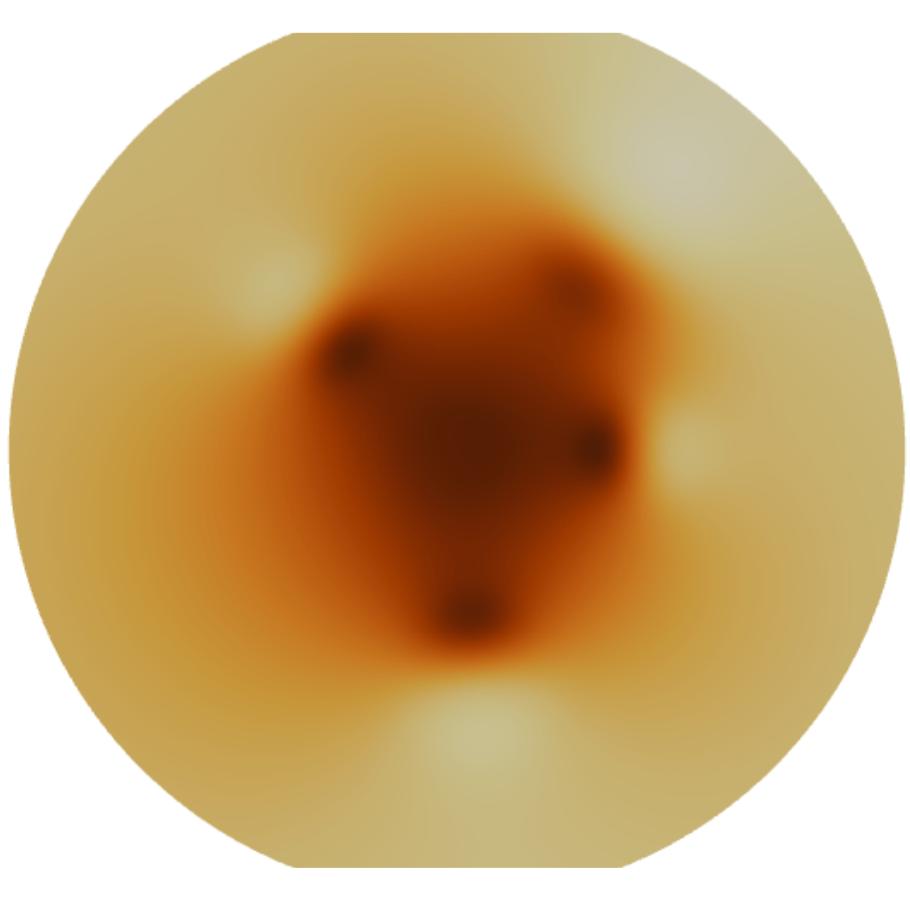


- 97.8% of PIN1 net cell vectors are oriented toward the center (in the peripheral zone).
- Only 47.8% of PIN1 internal cell vectors are oriented toward the center (in the CZ).
- PIN1 net cell vectors are more oriented toward the center than PIN1 internal cell vectors regarding both number and strength.
- Both cell vectors have a low intensity in the central zone, and a global convergence toward the center.

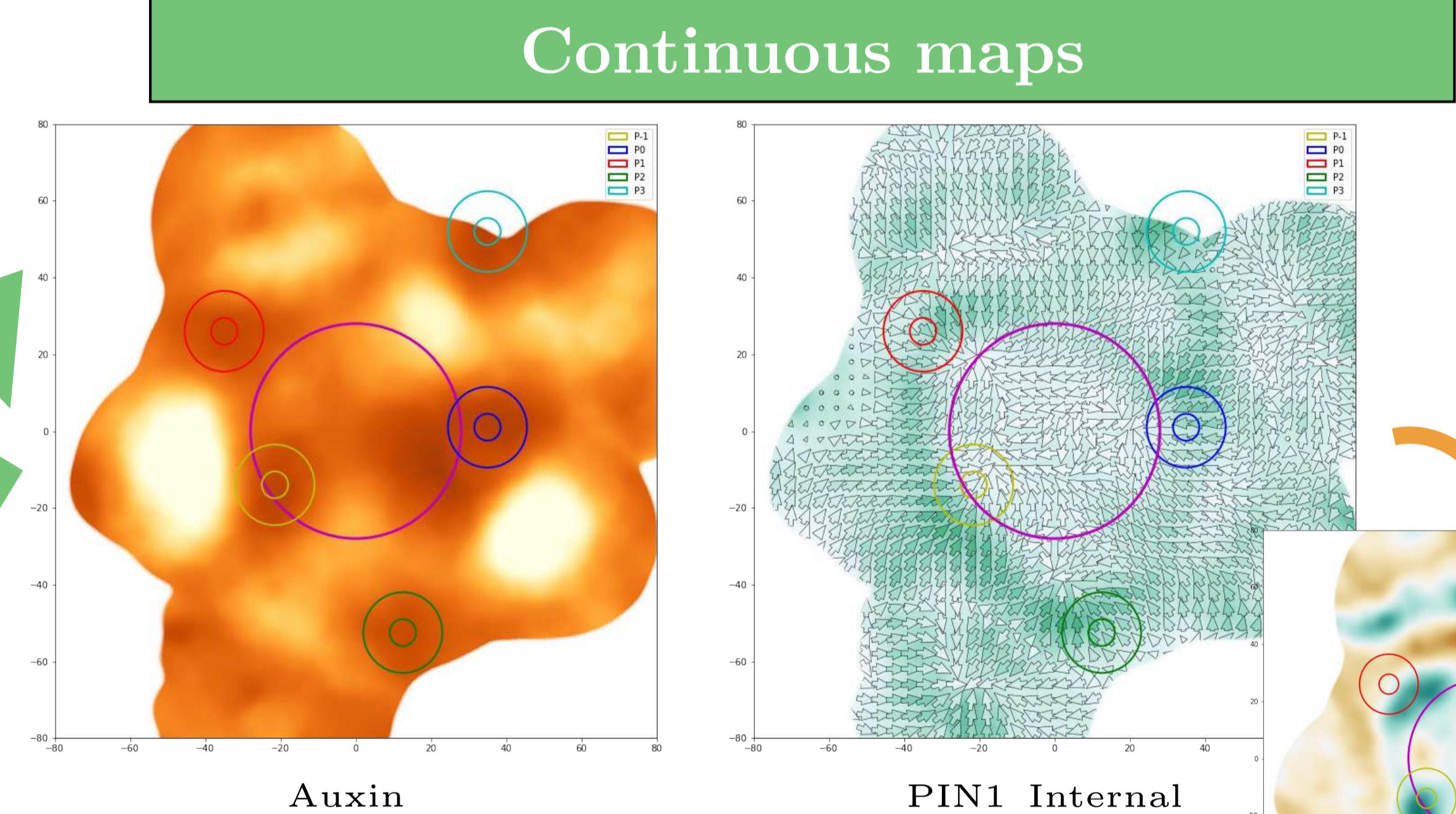
Modeling PIN1 polarity fields

- a : auxin continuous concentration.
- \mathbf{v} : PIN1 abstracted vector field.
- Dynamics: a diffuses and is transported by \mathbf{v} .

$$\frac{\partial a(x, t)}{\partial t} = \underbrace{\Delta a(x, t)}_{\text{Diffusion}} - \underbrace{\nabla \cdot (\mathbf{v}(x)a(x, t))}_{\text{Advection at speed } \mathbf{v}}$$

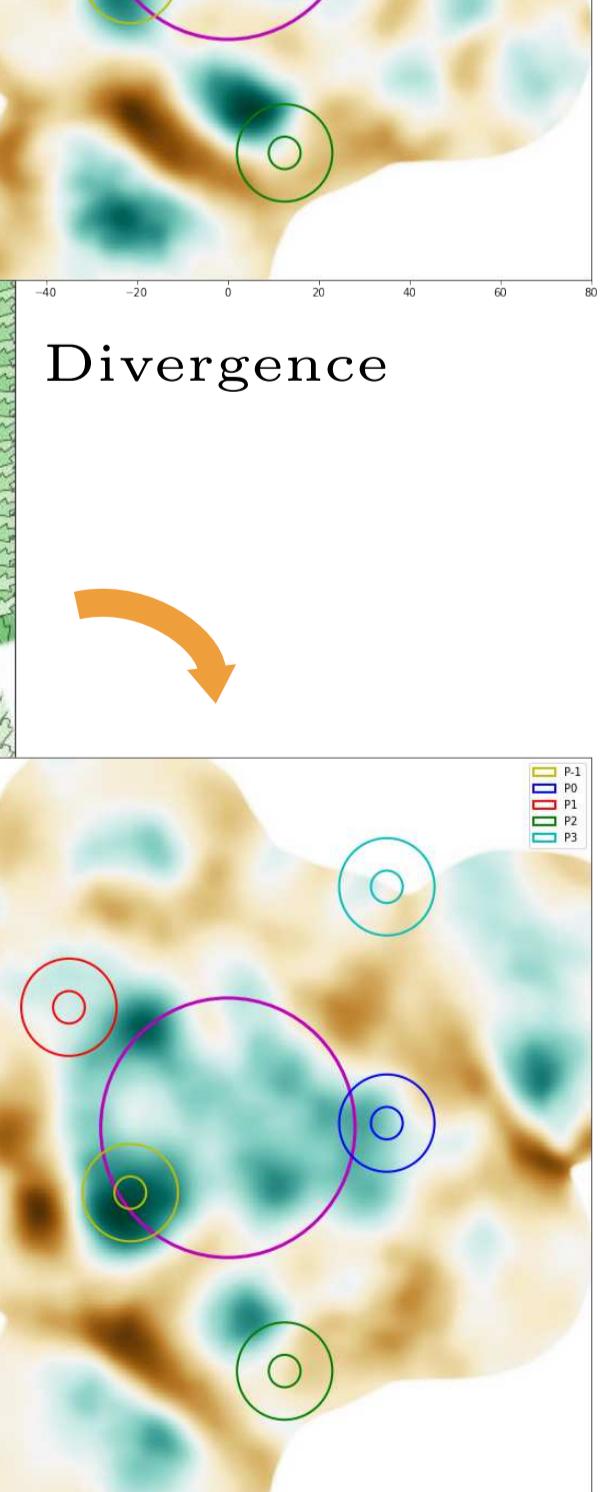


New hypothesis:
accumulations shape due
to PIN1 level regulation



- \mathbf{v} : uniformly pointing to the center, lower in the CZ.
- Localized upregulations of \mathbf{v} creates maxima and minima of a .

- Primordia are located in the highest convergence zones.
- Continuous maps allow to compare different signals, individuals, and time points.



Quantification of different PIN1 quantities at different scales leads to enrich our characterization of the SAM. PIN1 level is upregulated close to primordia. It is sufficient to make auxin accumulate according to the modeling.

Future work: track PIN1 during primordia development, include temporal aspects to the PIN1 modeling.