

mapDDX: Descriptive and Diagnostic plots, and Xcorrelation analysis of activity maps

Nov. 17 (Updated May 2018)

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Introduction

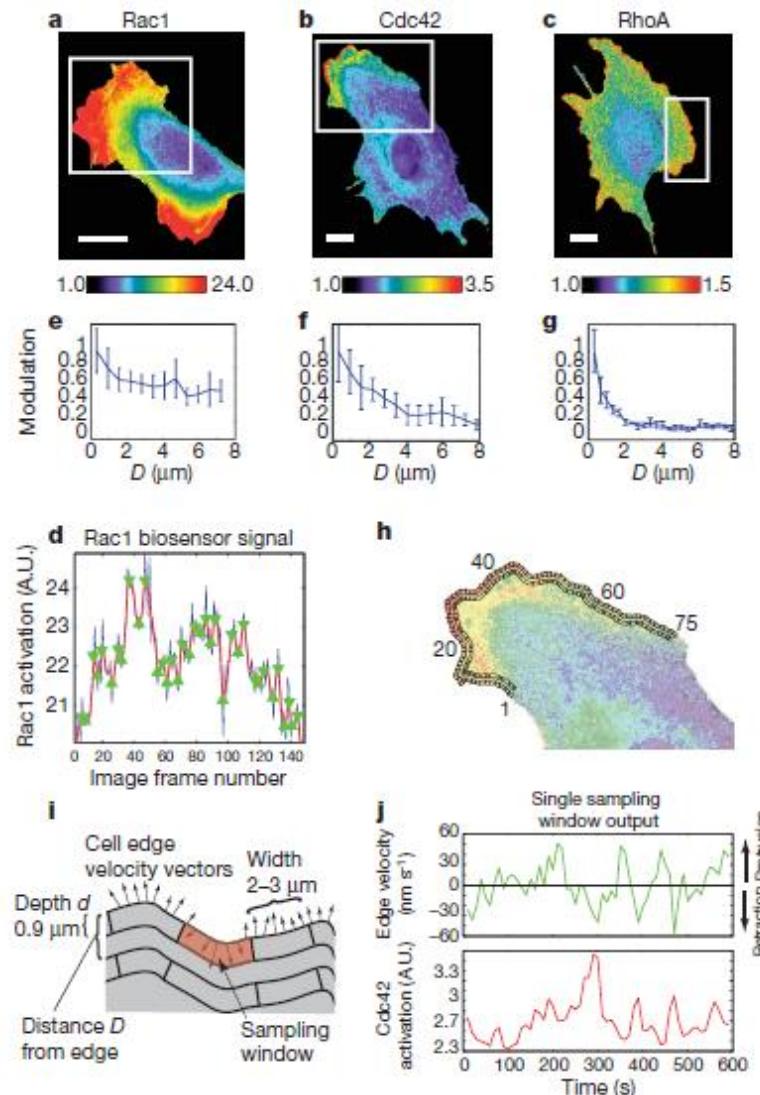


Figure 1 | Activation of Rho GTPases in migrating MEFs. a–c, Rac1 (a), Cdc42 (b) and RhoA (c) activation reported by biosensors. White box indicates region of interest selected for analysis. Scale bar, 20 μm . Colour scale indicates activation levels.

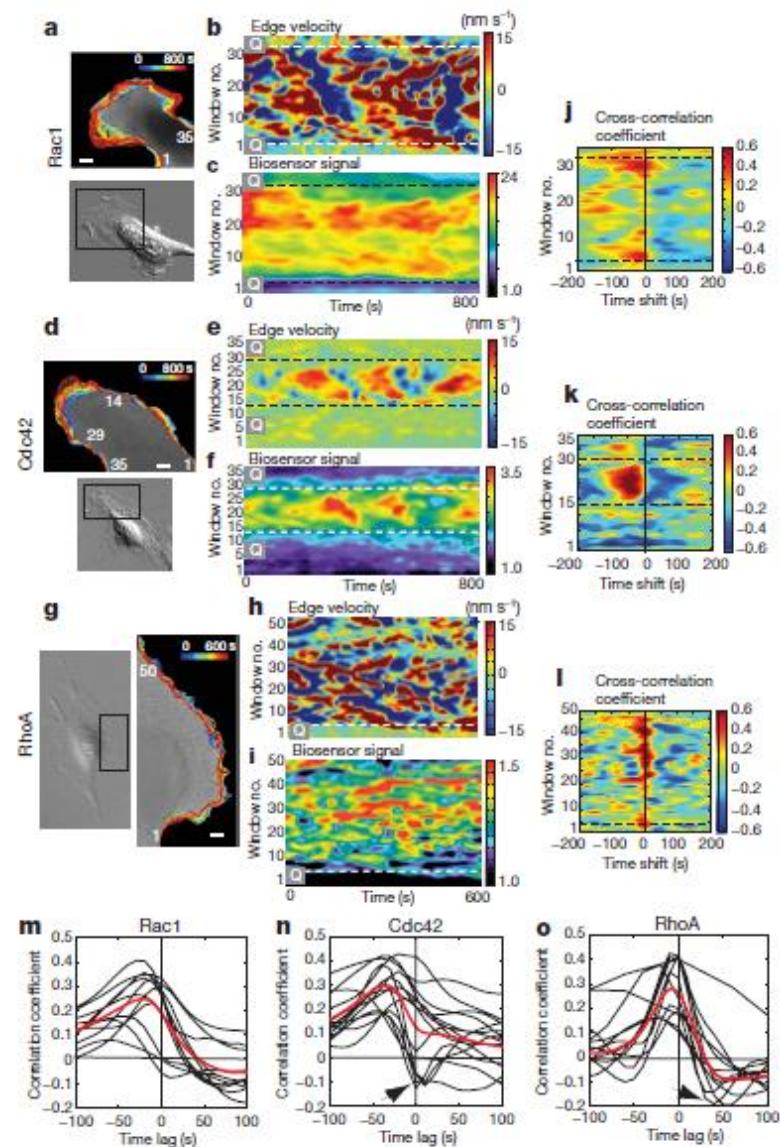


Figure 2 | Dynamics of cell edge morphology and GTPase activation.

Applications > morphodynamics > mapDiagnosticsXcorrCurves

List of .m functions for mapDDX

1. mapDescriptives_OneChan

1. mapOutlierImputation > myknnimpute
2. topographMD
3. TimeSpaceAutoCorPlot
 1. autoCorrMap > nanXcorrelation
4. autoCorrCurvePermTest > autoCorrMap
5. nanAdfTestMap
6. nanZscore
7. checkWindowJump

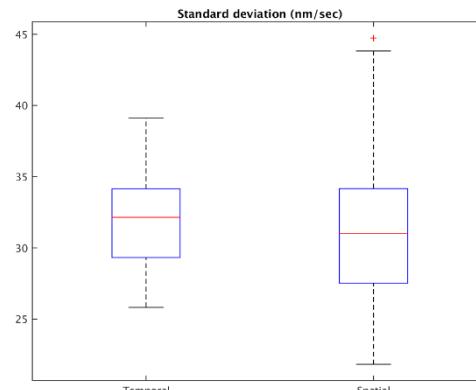
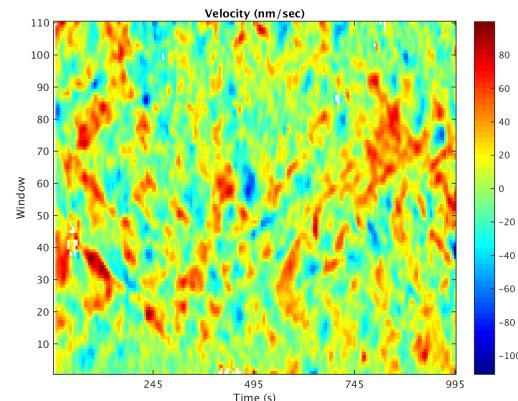
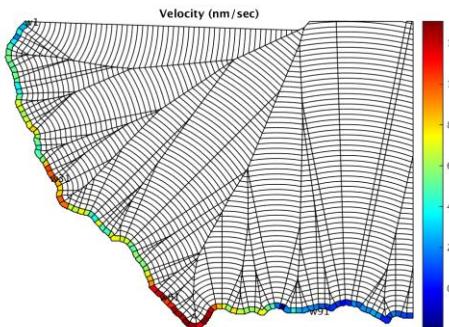
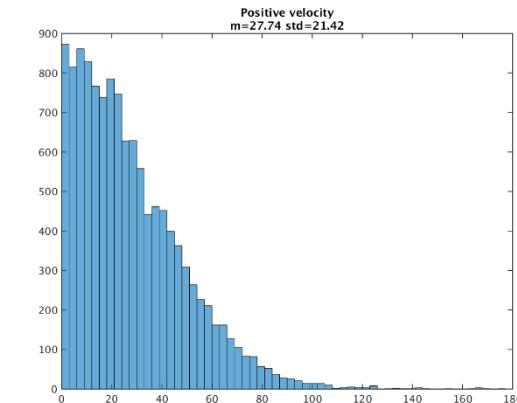
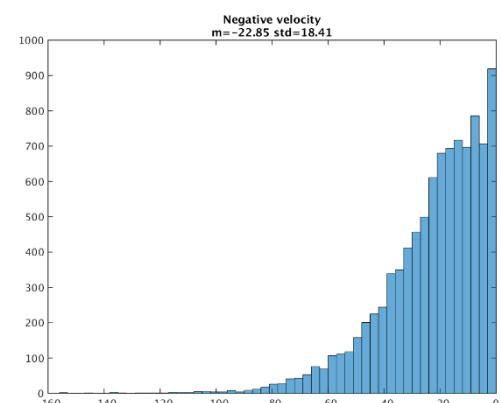
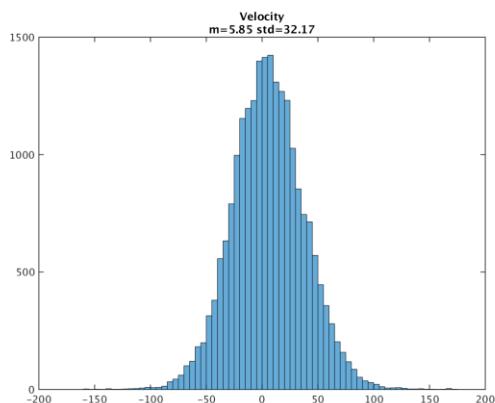
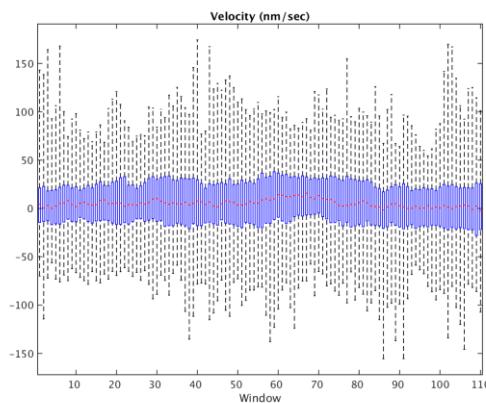
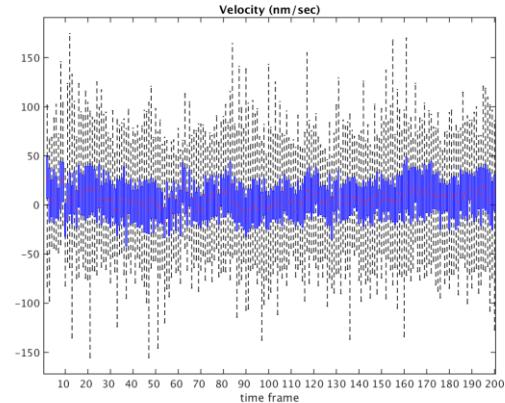
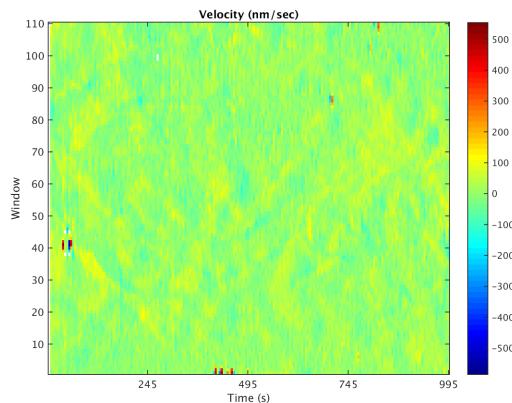
2. mapXcorrCurvePermutation, mapXcorrCurvePermutation_Vel

1. mapOutlierImputation
2. topographMD
3. smoothingSplineCorMap
4. xcorrCurvePermutationTest
 1. xcorrMapPlot > nanXcorrMaps > nanXcorrelation
 2. timePermDistXcorrMean > nanXcorrMaps

3. mapDescriptives_Vel_LB

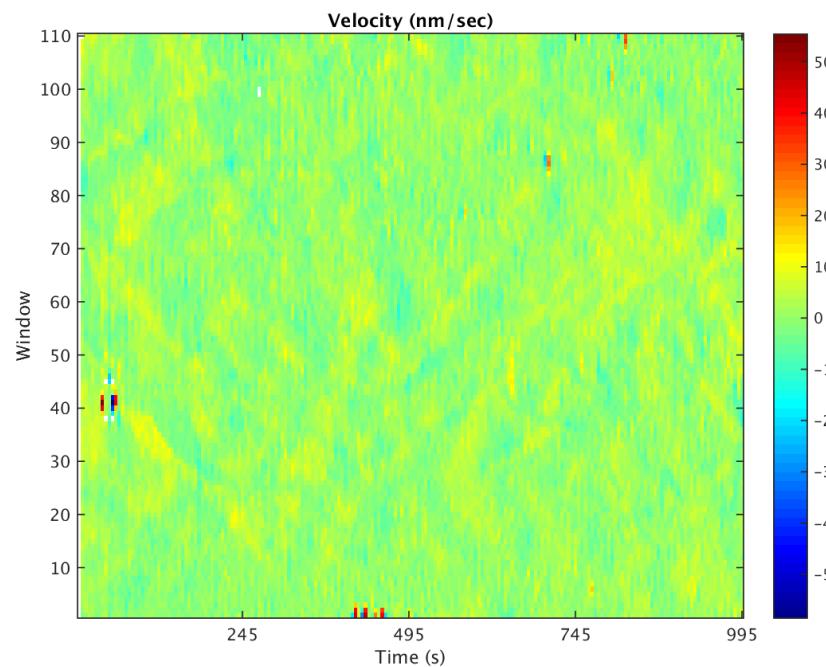
4. MLsummary_XcorrCurvesVelAcf

Descriptives: edge velocity

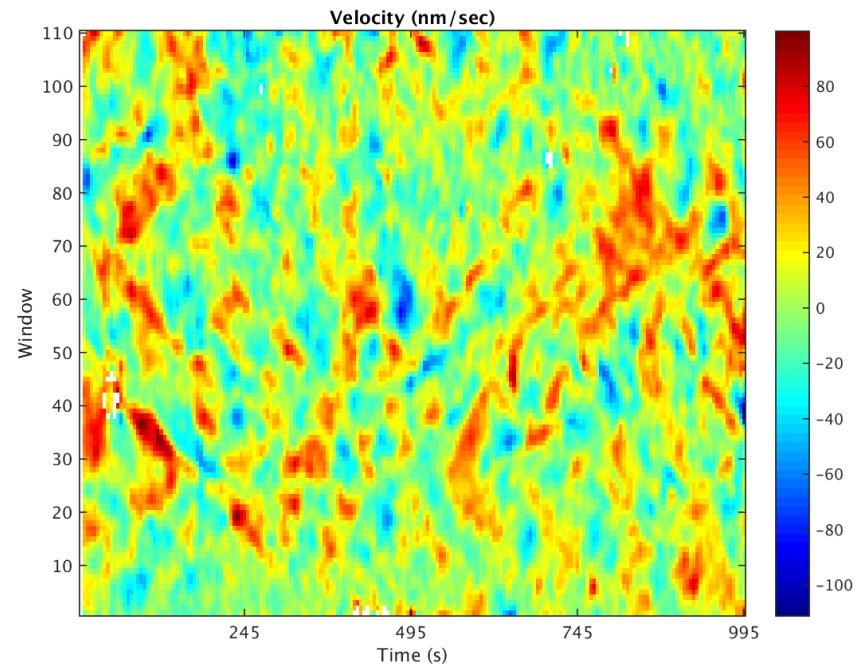


Ptk1 Dec.-1 (cell#5)

Raw activity map

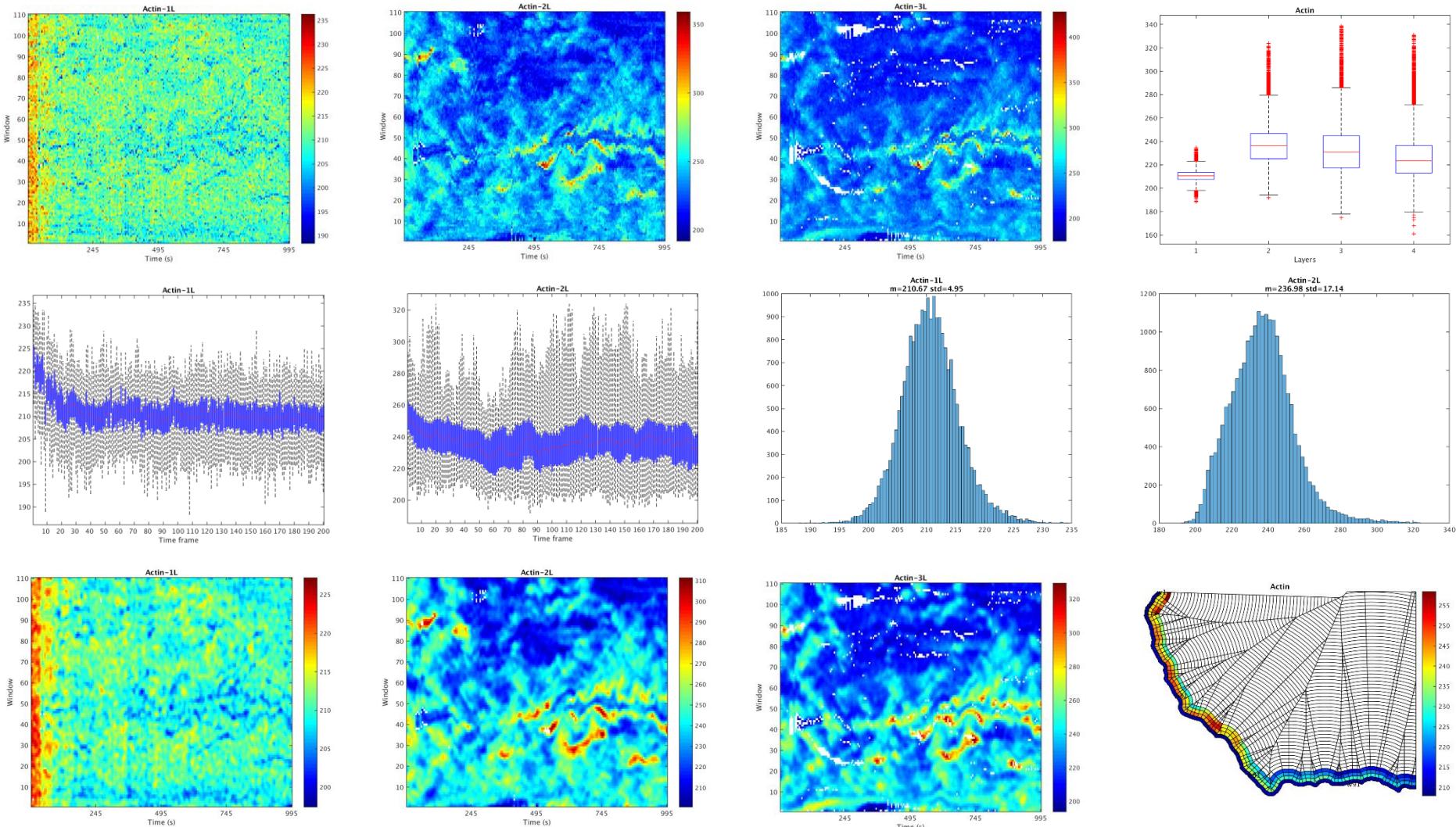


Outlier adj./smoothed activity map



- Velocity map = $\{V(t, w): t \geq 2 \text{ (frames)}, w \geq 1 \text{ (windows)}\}$
- $V(t, w)$: displacement (nm) between the frames at (t-1) and t divided by time interval (sec)
- Z-scores of all the $\{V(t, w)\}$ are computed
- $|Z\text{-scores}| > 5 \Rightarrow$ detected outliers
- Outliers and missing windows can be imputed by using the most similar windows (`knnimpute.m` function)

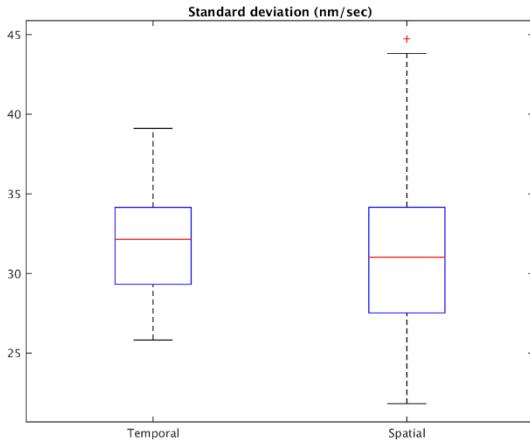
Descriptives: Actin 1-3 layers



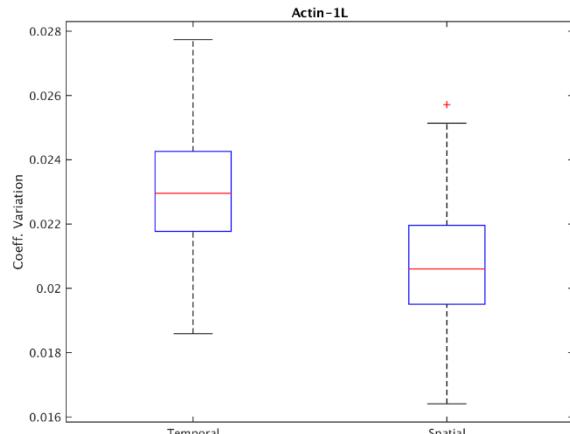
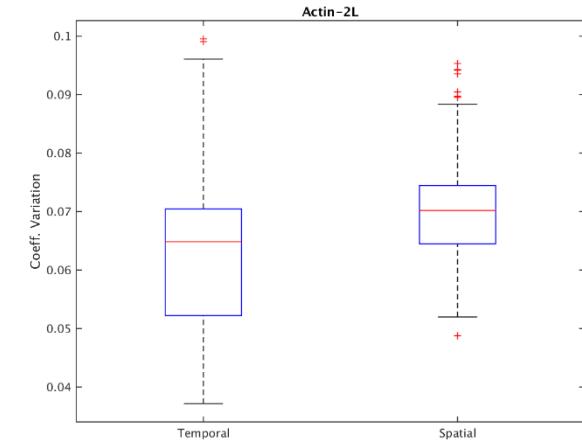
- Check layer-wise heterogeneity.
- Use the boxplot over time to check if there is a trend over time which can mislead temporal correlation analysis.
- A topograph shows spatial distribution of activities averaged over time.

Diagnostics: coefficient of variation (CV)

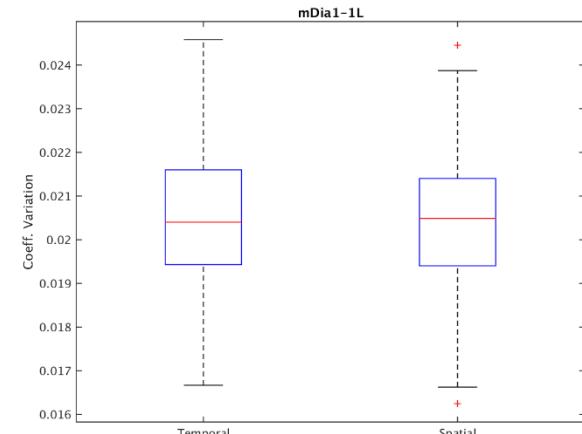
- $CV(X) = SD(X)/E(X)$
- CV is a statistical measure of dispersion of a probability distribution.
- CV has no unit and more suitable for non-negative observations.



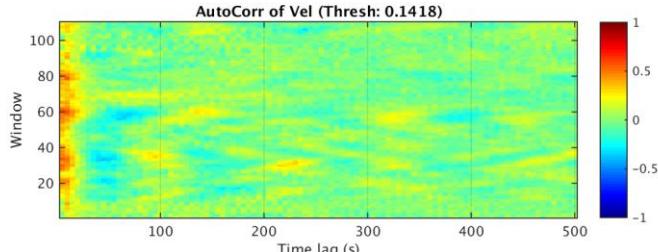
- Temporal/spatial variations of velocity are similar for the above cell.
- Common case



- Temporal CV is bigger than spatial CV, which suggests temporal heterogeneity.
- Such heterogeneity may not be relevant to underlying biological questions.

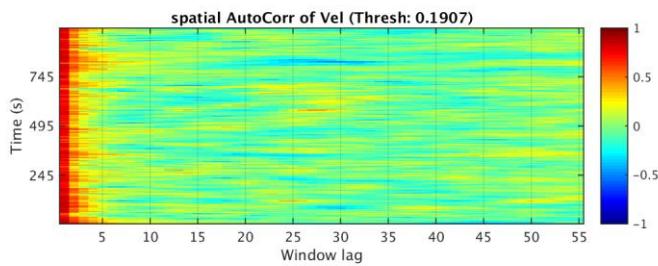


Diagnostics: temporal/spatial autocorrelations



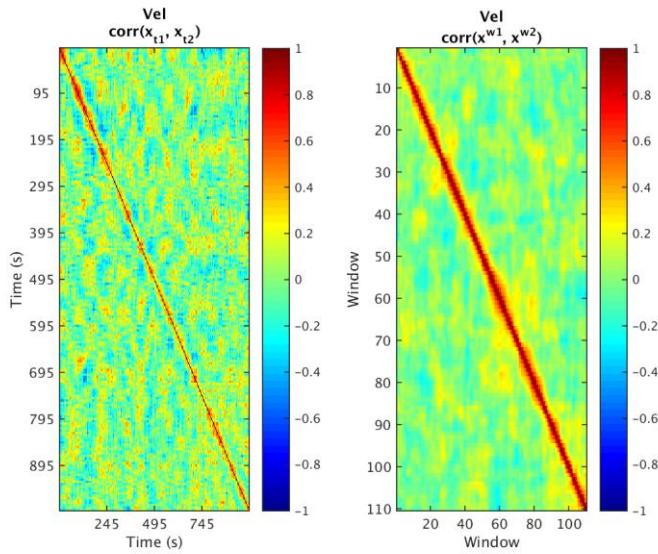
Temporal
autocorr.

$$\rho_w(h) = \frac{1}{t_m} \sum_{t=1}^{t_{max}-h} \dot{X}(t+h, w) \dot{X}(t, w)$$



Spatial
autocorr.

$$\rho_t(h) = \frac{1}{w_{max}} \sum_{w=1}^{w_{max}} \dot{X}(t, w+h) \dot{X}(t, w)$$



Corr. between
two time frames
(two columns)
- Left

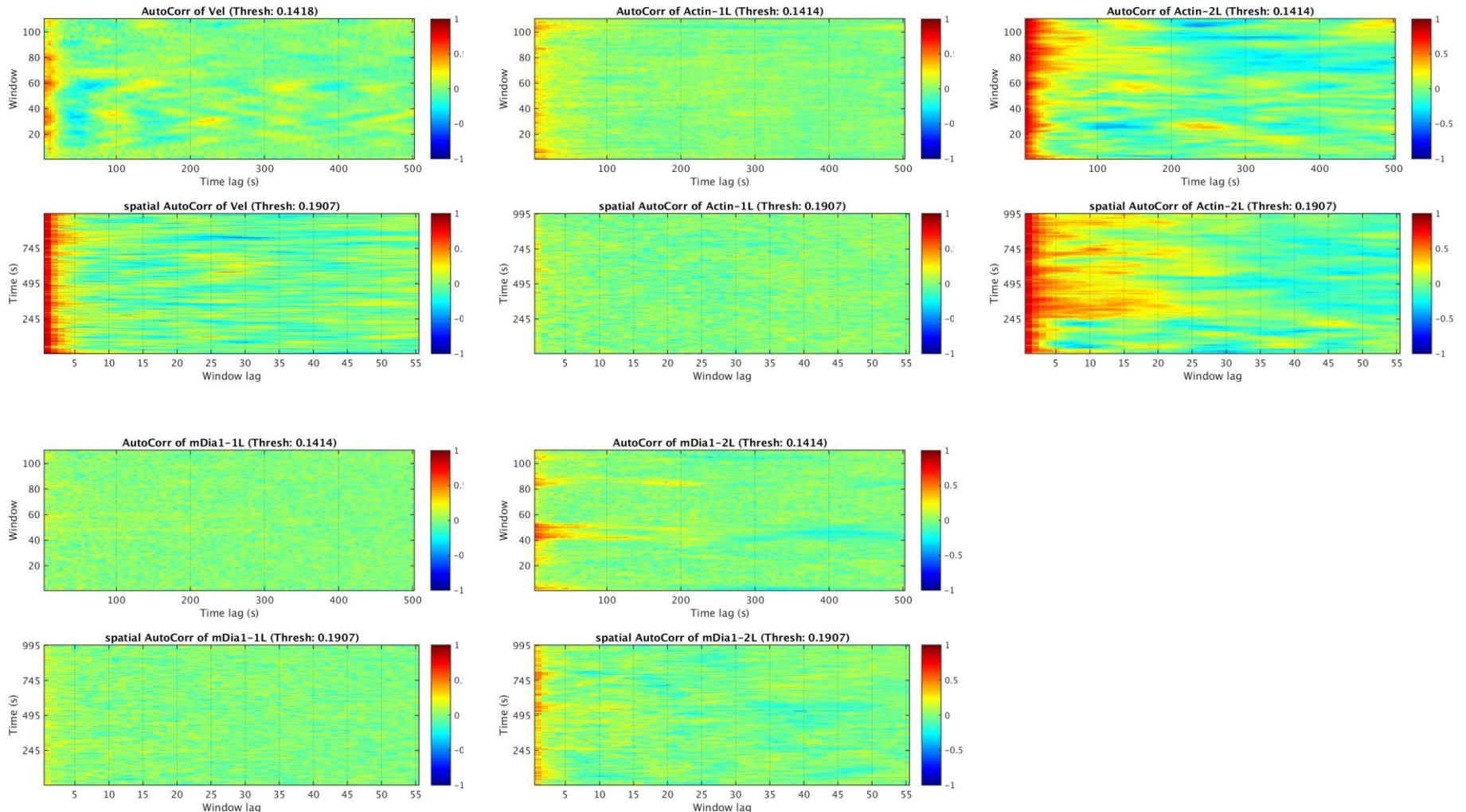
Corr. between
two windows
(two rows)
- Right

$$\rho(t_1, t_2) = \frac{1}{w_{max}} \sum_{w=1}^{w_{max}} \dot{X}(t_1, w) \dot{X}(t_2, w)$$

$$\rho(w_1, w_2) = \frac{1}{t_{max}} \sum_{t=1}^{t_{max}} \dot{X}(t, w_1) \dot{X}(t, w_2)$$

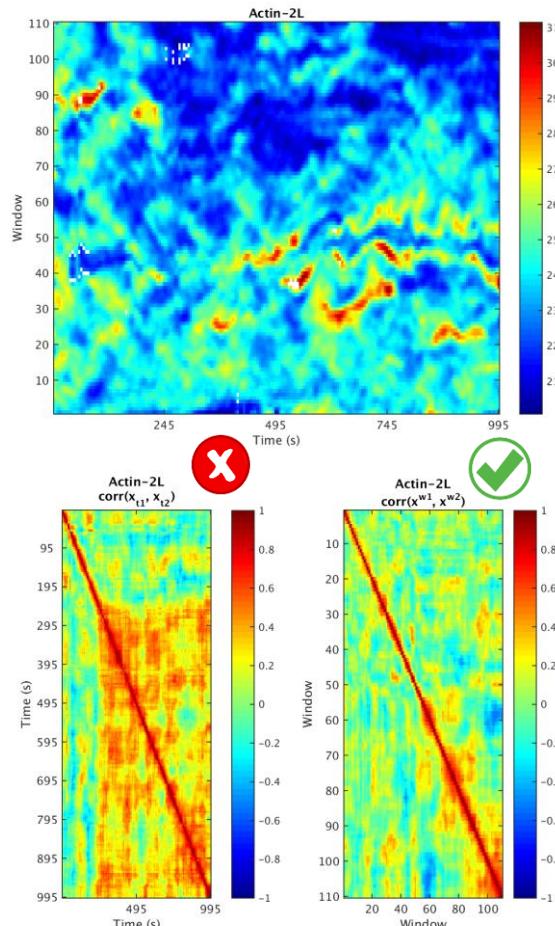
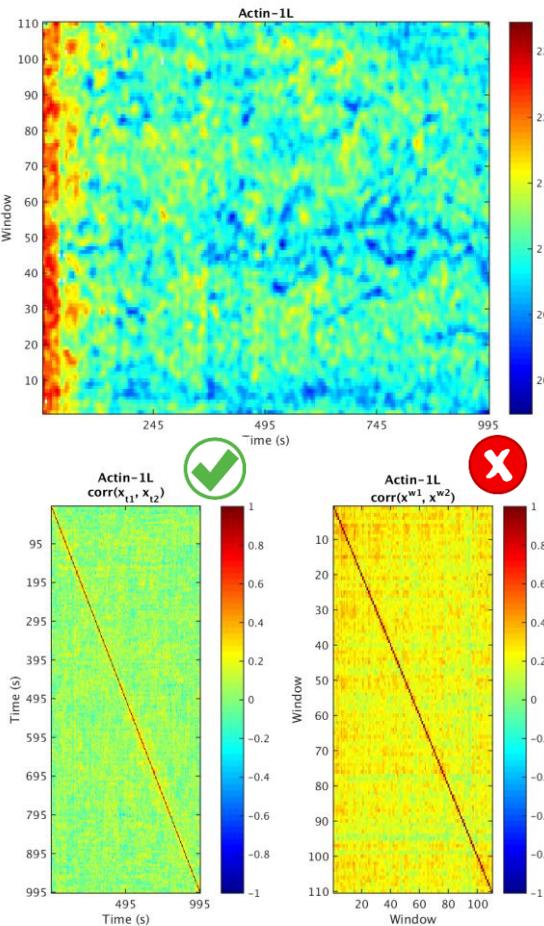
- Here, $\dot{X}(t, w)$ denotes a standardized variable.
- The standardization can be different in different context.

Autocorrelation maps for vel, actin, mDia1



- The quality of 1st layer is not good.
- Autocorrelations of actin are higher than mDia1.

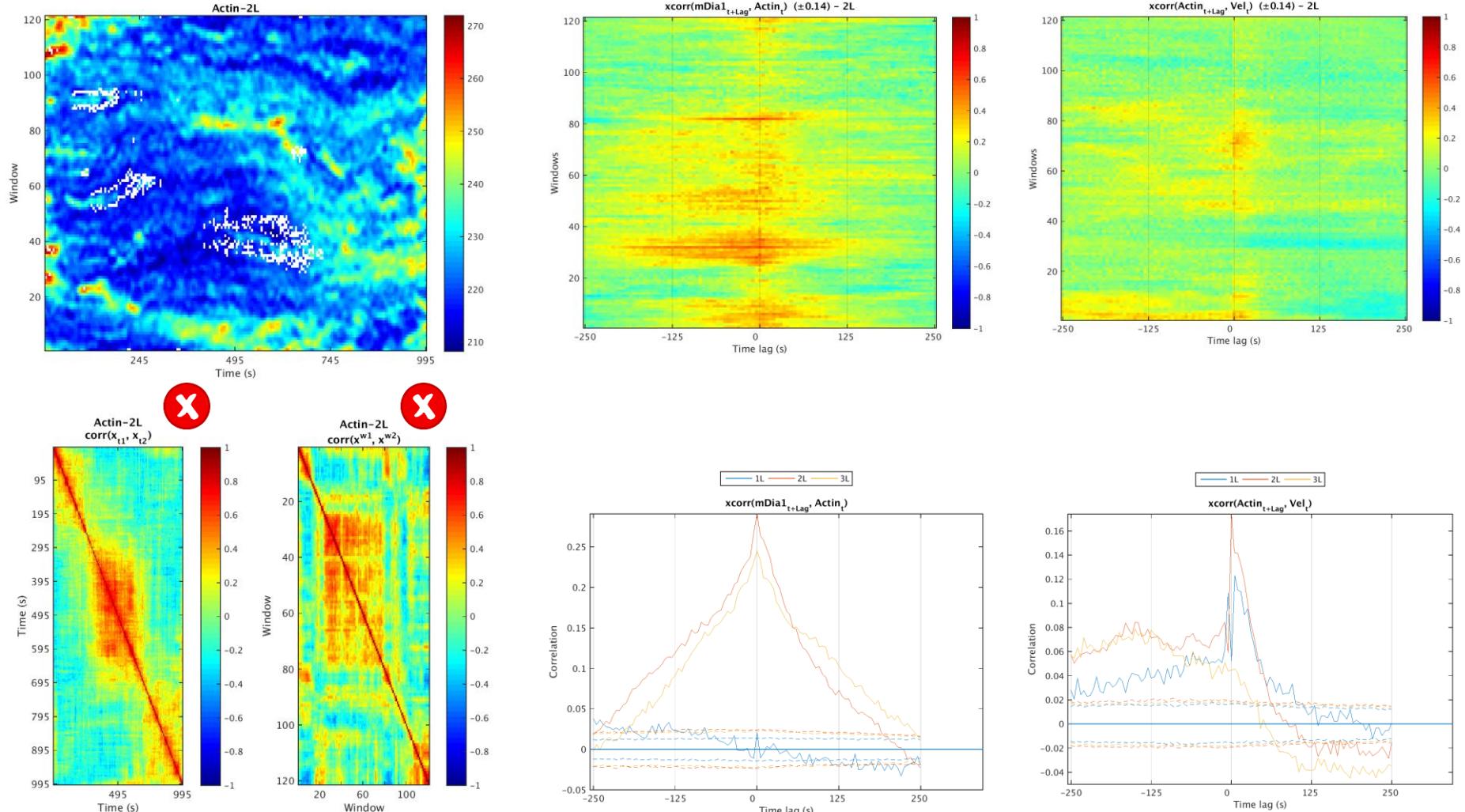
Correlations between column or row vectors



- High correlations between windows indicate that row vectors share common temporal pattern (Activities are high at the beginning).
- In such case, temporal analysis can be misleading.

- High correlations between time frames indicate that column vectors share common spatial pattern (Activities are low in win80~110 after 250 sec) .
- In such case, spatial analysis can be misleading.

Ptk1 Cell#1



- The above cross correlations are not typical.
- The common temporal pattern in win20~60 may result in spurious high correlations.

Augmented Dickey-Fuller (ADF) test

- `adftest.m` tests whether a TS is stationary or non-stationary.
- Matlab manual

Autoregressive model variant, which specifies a test of the null model

$$y_t = y_{t-1} + \beta_1 \Delta y_{t-1} + \beta_2 \Delta y_{t-2} + \dots + \beta_p \Delta y_{t-p} + \varepsilon_t$$

against the alternative model

$$y_t = \phi y_{t-1} + \beta_1 \Delta y_{t-1} + \beta_2 \Delta y_{t-2} + \dots + \beta_p \Delta y_{t-p} + \varepsilon_t,$$

with AR(1) coefficient, $\phi < 1$.

$$(\Delta y_t = y_t - y_{t-1})$$

- Idea:

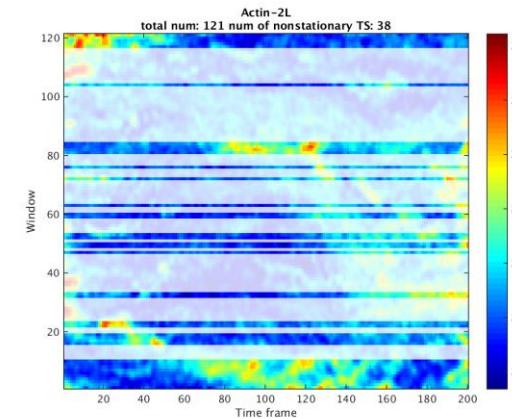
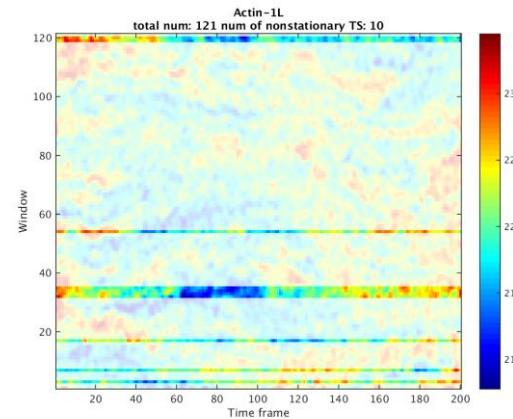
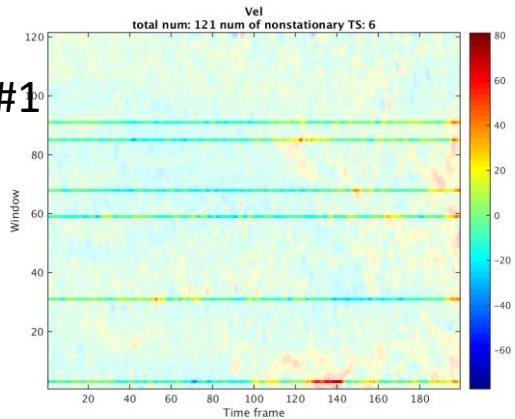
- A stationary TS has the property that $E(Y_t) = \text{constant}$ which is a long-term equilibrium level.
- Consider a TS regression model:

$$\Delta y_t = \alpha + \gamma y_{t-1} + \beta_1 \Delta y_{t-1} + \varepsilon_t$$

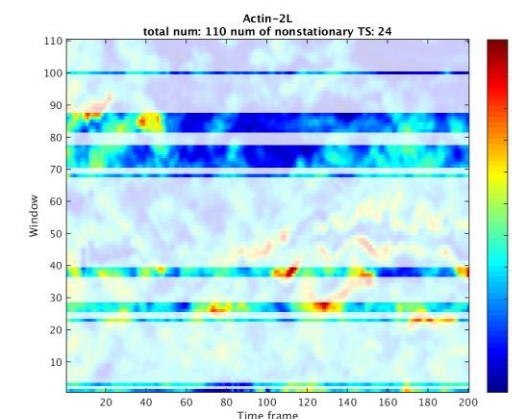
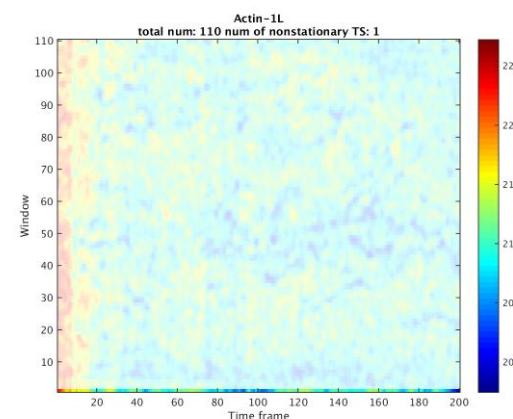
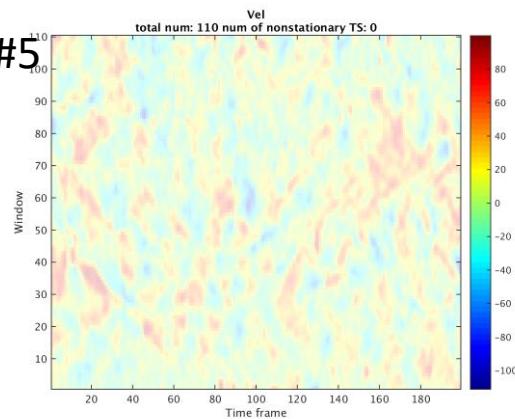
- For the stationarity, y_{t-1} needs to be negatively correlated with $\Delta y_t = y_t - y_{t-1}$.
- We can test the non-stationarity ($H_0: \gamma = 0$) vs. stationarity ($H_1: \gamma < 0$).
- If H_0 is true, then $\{\Delta y_t: t \geq 1\}$ is an AR(1) process, and $\{y_t: t \geq 1\}$ is an integrated process like a random walk process which does not have a constant long-term equilibrium level.

- Non-transparency (alpha=1) indicates windows tested to be non-stationary
(Transparent, alpha=.8: stationary).

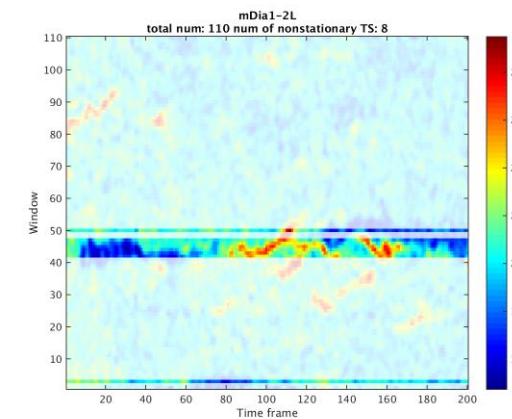
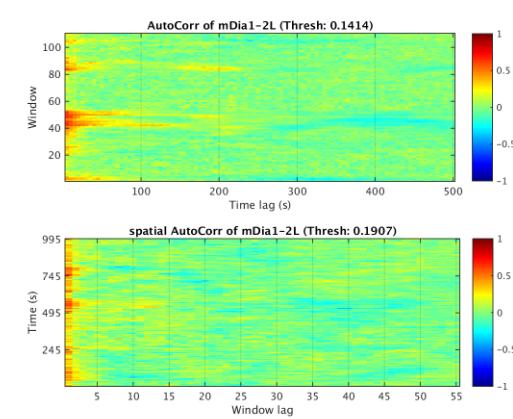
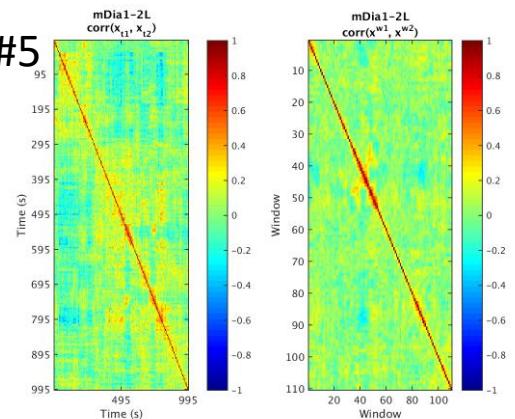
Ptk1 Cell#1



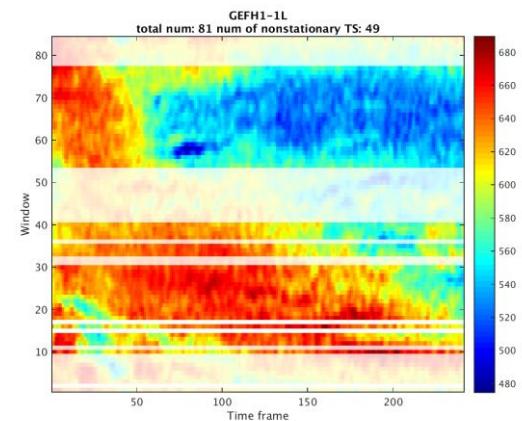
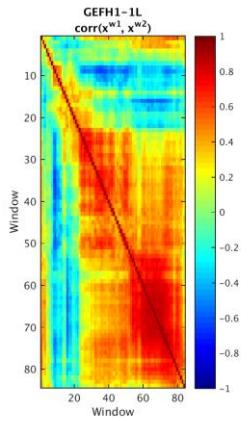
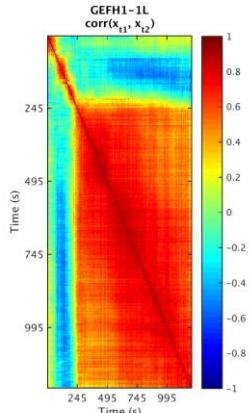
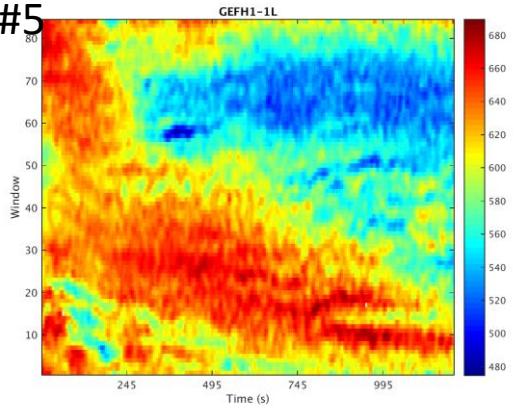
Ptk1 Cell#5



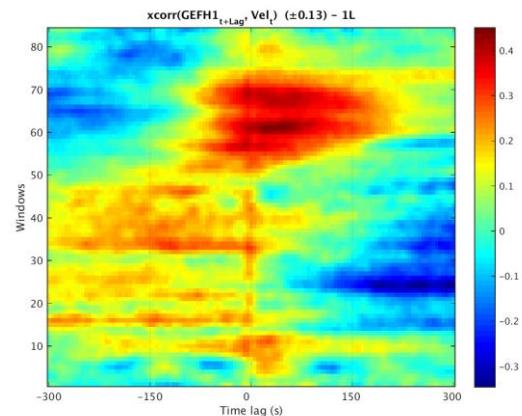
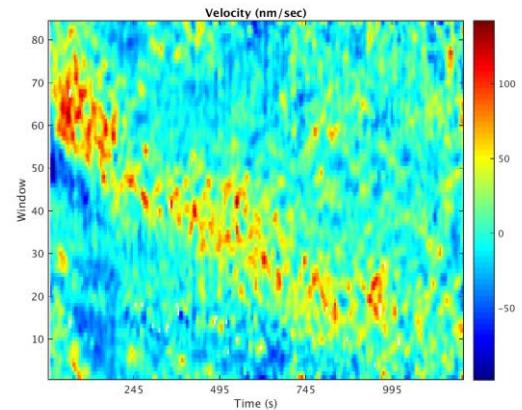
Ptk1 Cell#5
mDia1



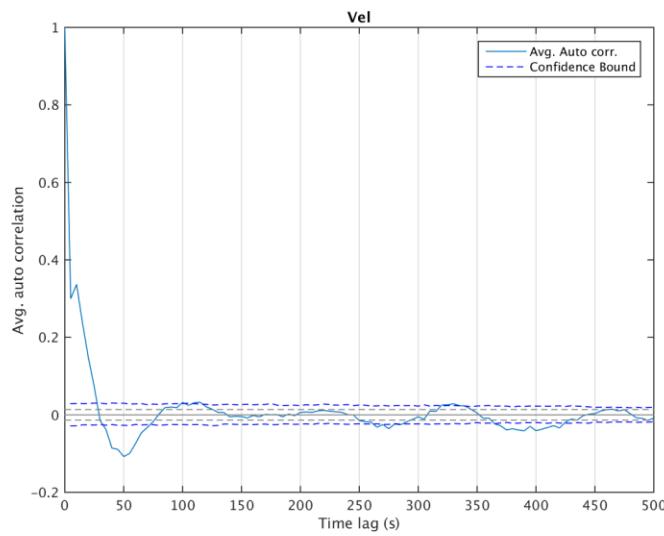
MDA Cell#5



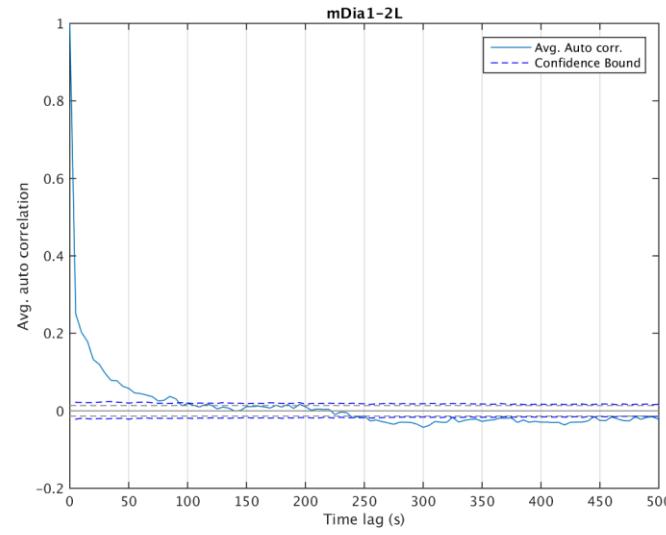
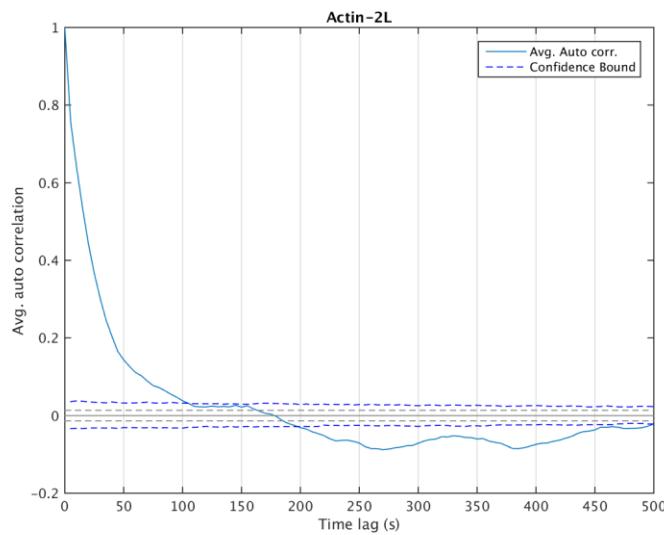
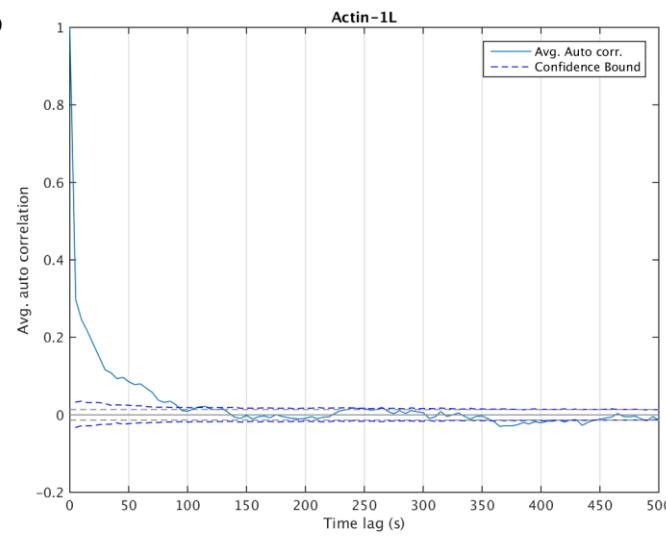
- Highly non-stationary case
- Correlations can be spurious.
- Methods for non-stationary TS are recommended.



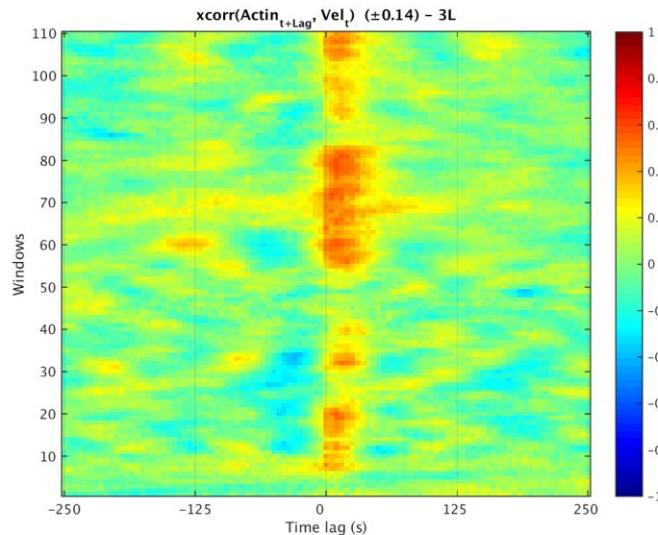
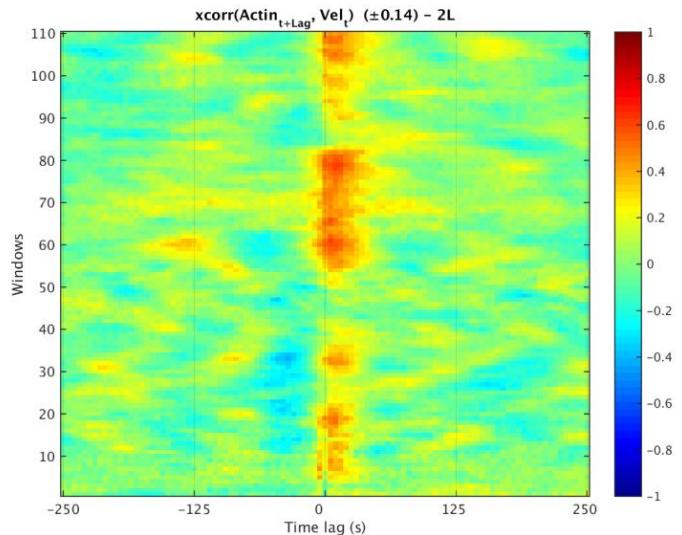
Autocorrelation curves (temporal)



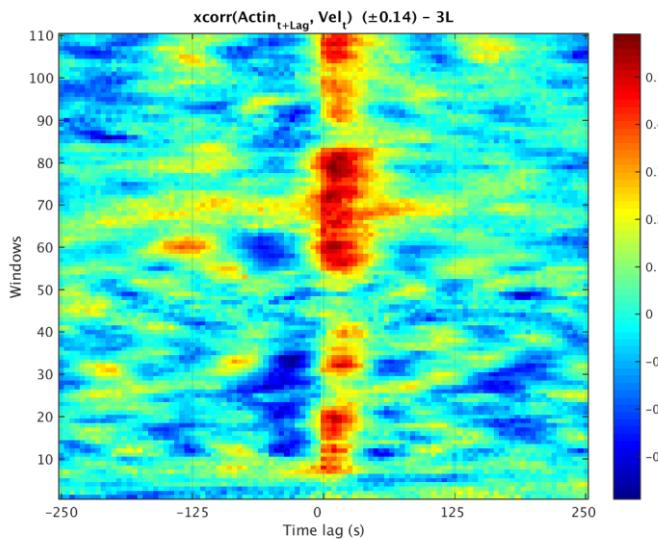
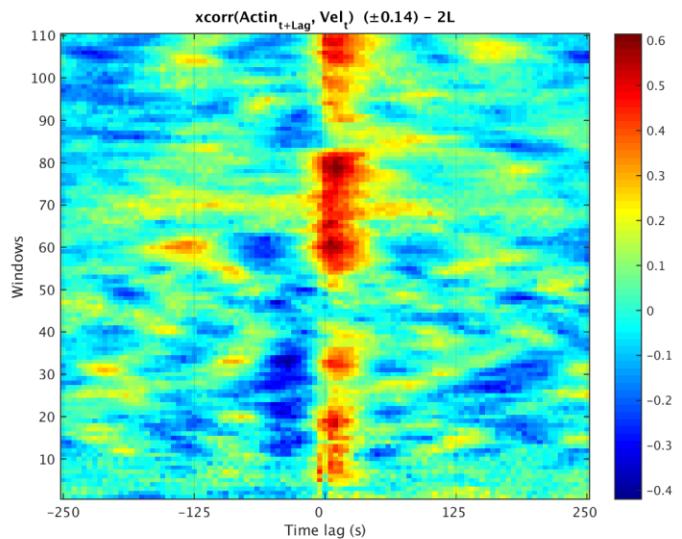
Ptk1 Cell#5



Cross correlation maps

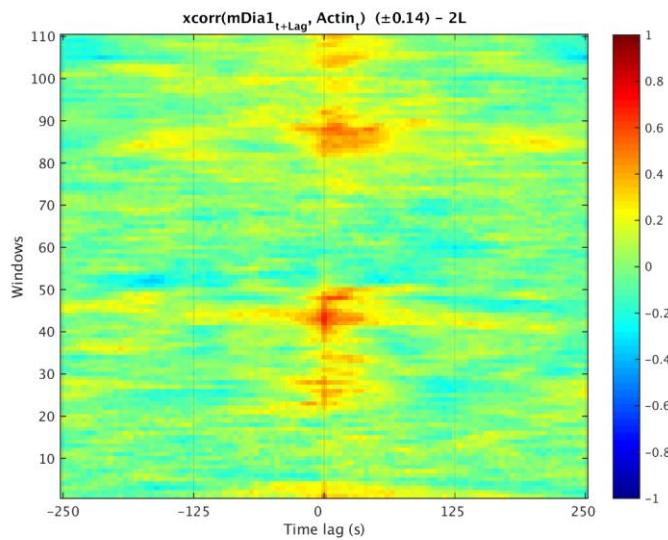
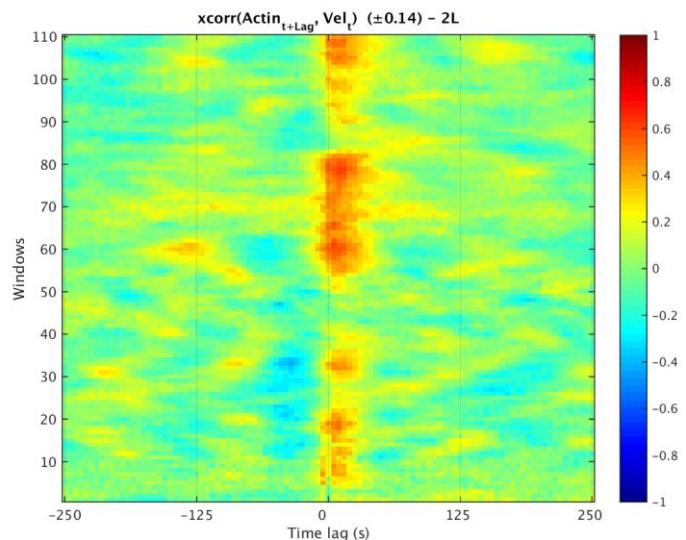
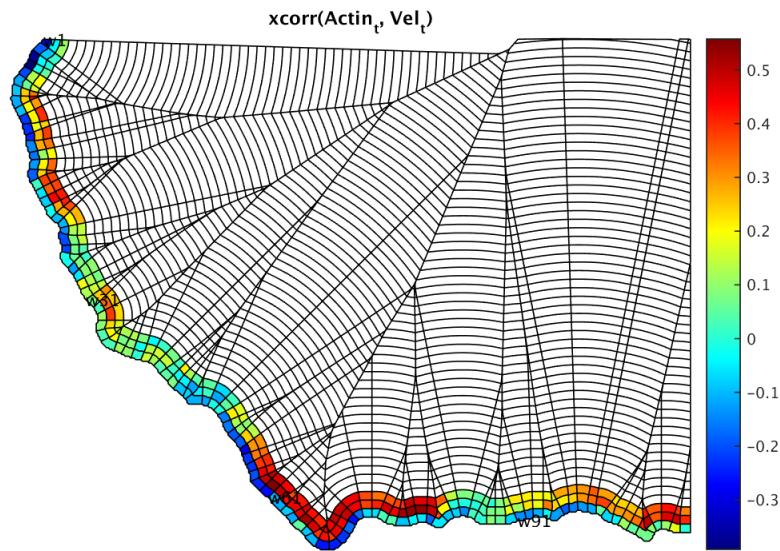
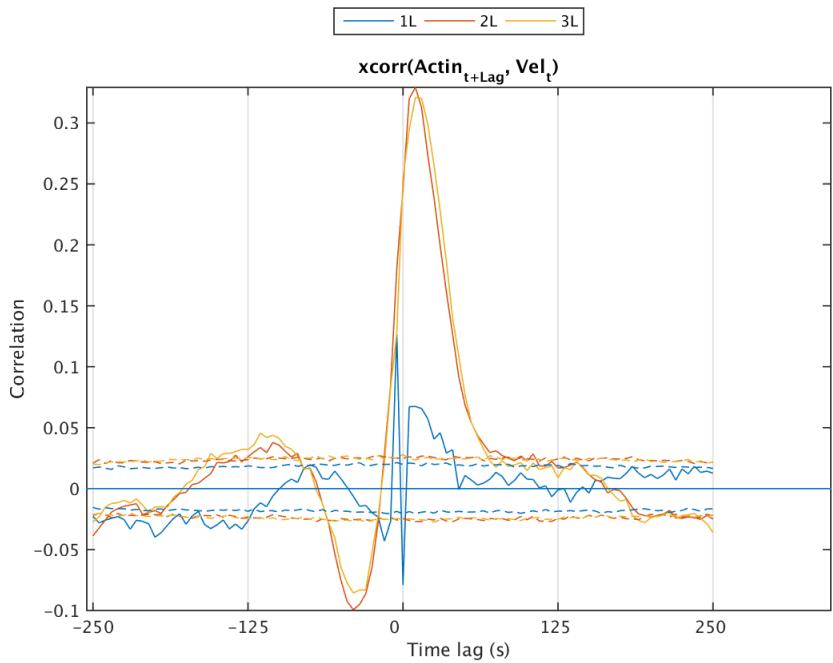


$[-1, 1]$ range



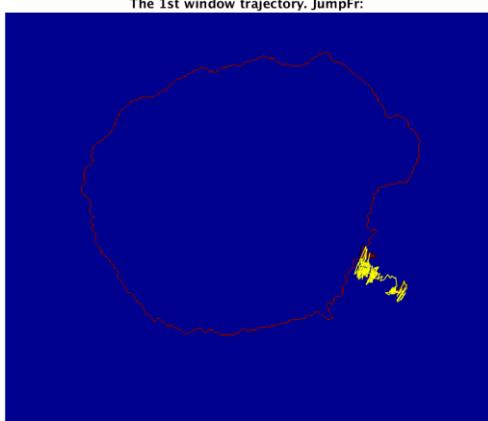
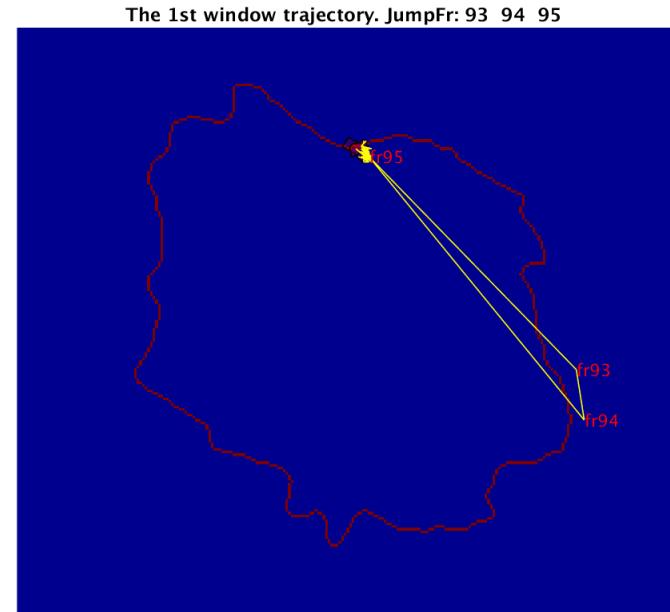
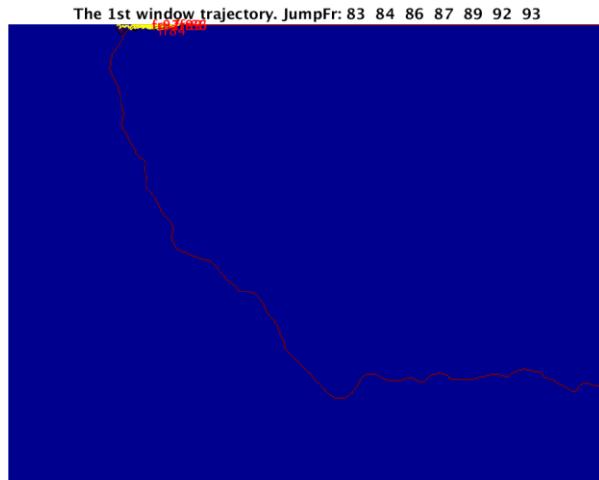
Range adjusted

Cross correlations



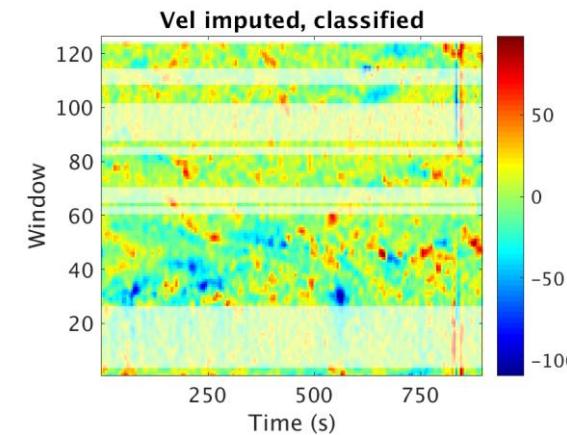
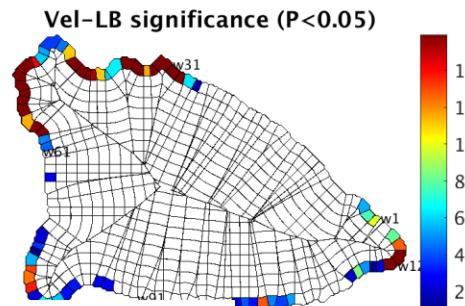
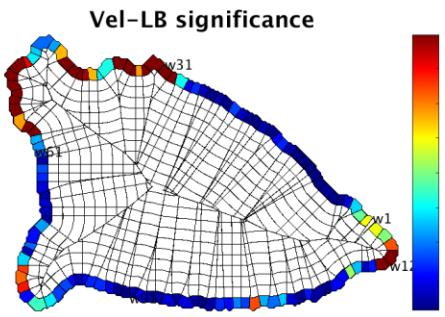
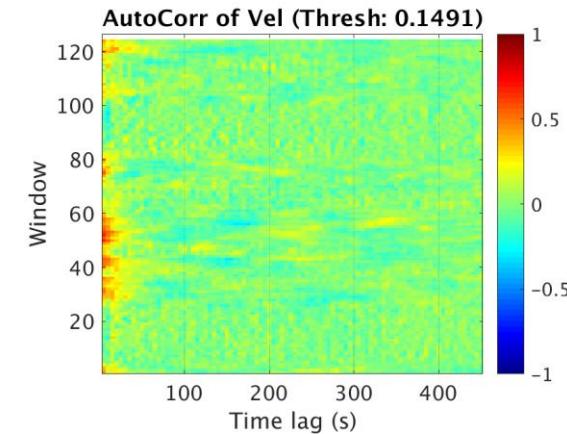
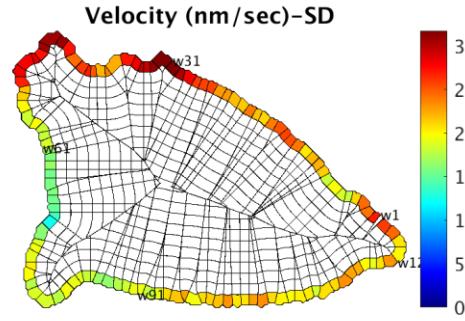
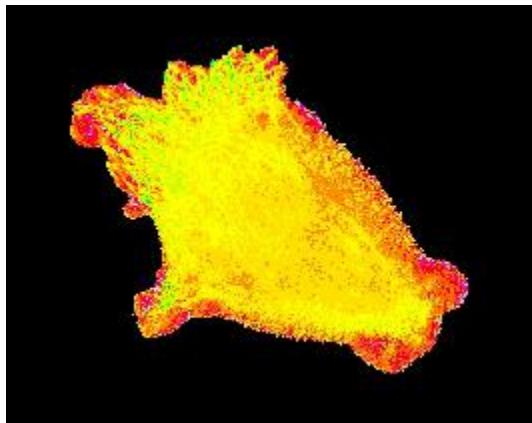
Diagnostics: Check windowing jumps

- It plots the trajectory of the 1st window in the 1st layer in order to find frames where window jumping happened.
- It plots the trajectory, the cell boundary at the last frame and estimated jump frames.



Identifying quiescent windows (Optional)

NovCell3



The test statistic is:^[2]

$$Q = n(n+2) \sum_{k=1}^h \frac{\hat{\rho}_k^2}{n-k}$$

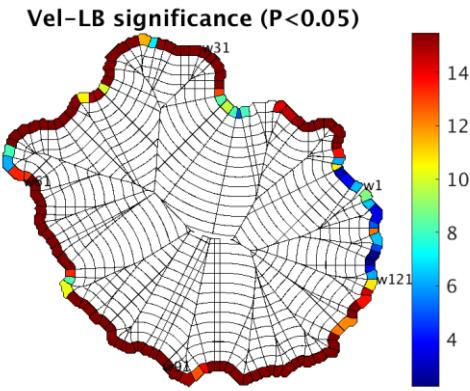
where n is the sample size, $\hat{\rho}_k$ is the sample autocorrelation at lag k , for rejection of the hypothesis of randomness is:

$$Q > \chi^2_{1-\alpha, h}$$

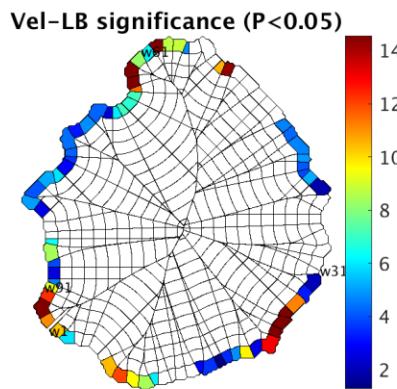
- We tested in each window whether velocity time series is auto-correlated or not by Ljung-Box (LB) test.
- Velocities are not auto-correlated in quiescent windows.

After filtering out quiescent sections

cell3



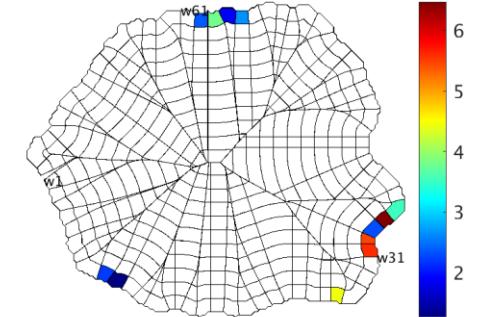
cell6



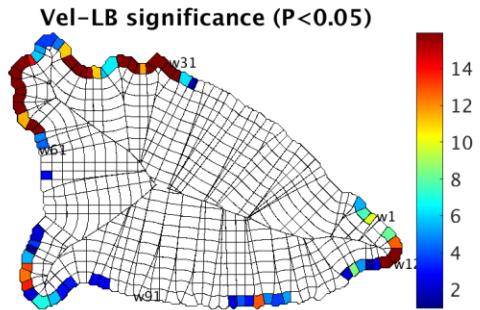
NovCell2

(The movie is quiescent)

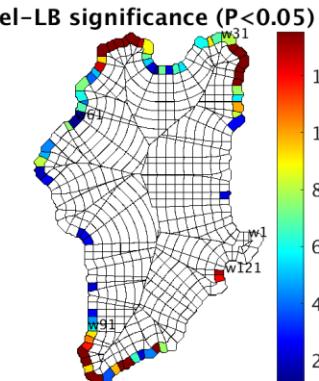
Vel-LB significance ($P < 0.05$)



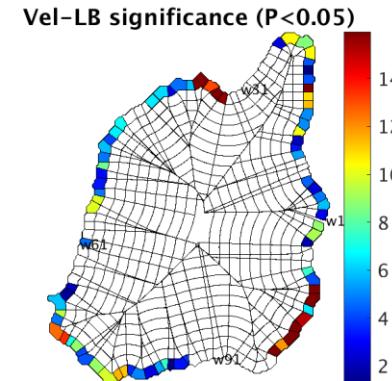
NovCell3



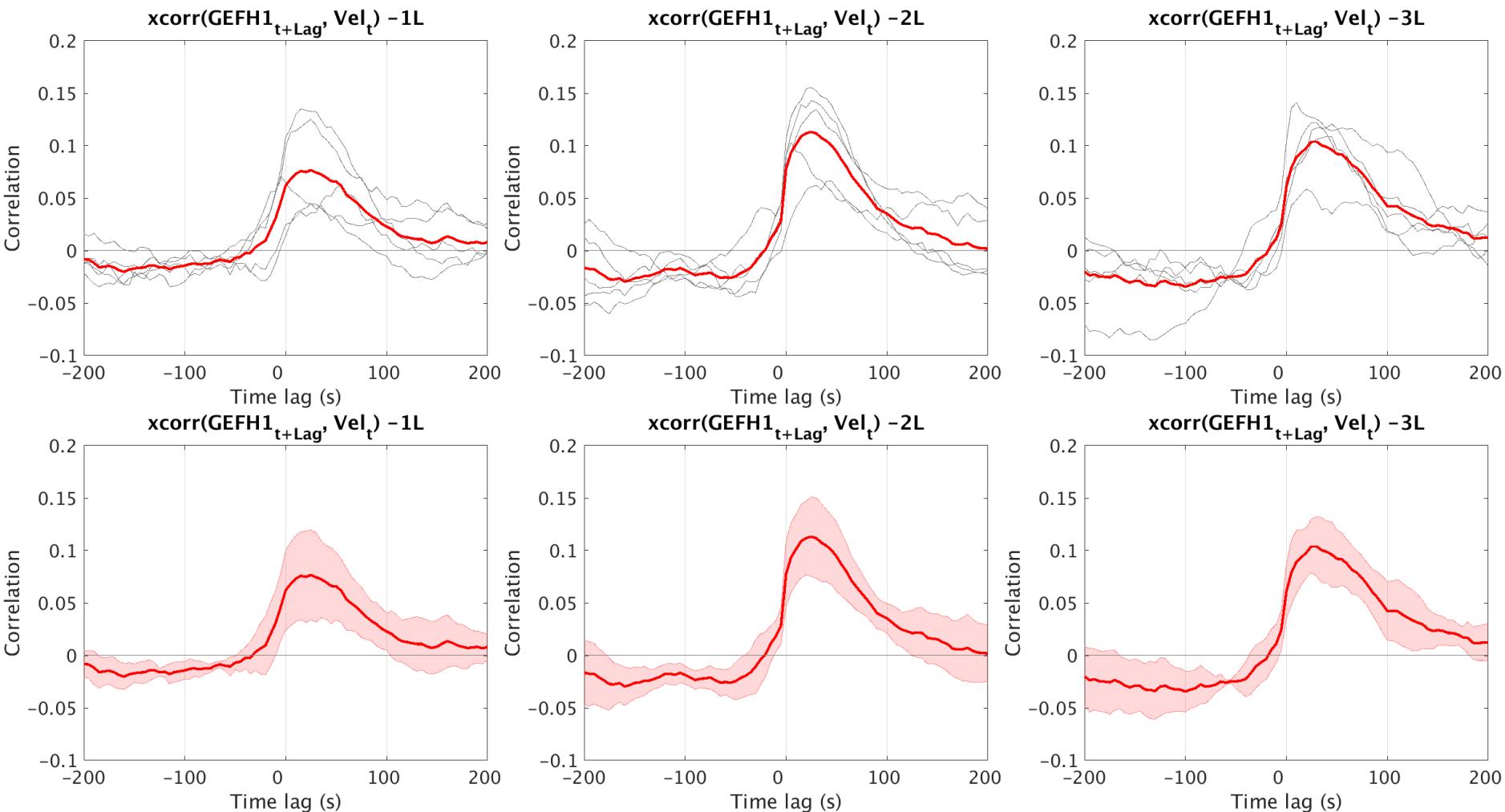
NovCell4



NovCell6



To summarize multiple movies and check cell-to-cell variation of correlation curves by using `MLsummary_XcorrCurvesVelAcf()`
 Cross correlation curves (n=5, GEFH1)



- Pos. correlation ($r \approx 0.11$) between current velocity and **25-sec** later GEFH1 activity shown in 1L~3L.
- Velocity precedes GEFH1 with a 25 sec lag.

See also the following to measure the coupling between molecular dynamics and edge motion as shown in the figure

1. Readme_FPAEME.pdf (Fluctuation Profiling Around Edge Motion Events)
2. 6 examples of pipeline codes (after movieData are processed by windowing.)
 1. example_pipeline_DX_FPAEME_1chan_actin.m
 2. example_pipeline_DX_FPAEME_2chan.m
 3. example_pipeline_DX_FPAEME_1chan_biosensor.m
 4. example_pipeline_DX_FPAEME_1chan_LB_actin.m
 5. example_pipeline_DX_FPAEME_2chan_LB.m
 6. example_pipeline_DX_FPAEME_1chan_LB_biosensor.m

