

ML methods for Edge detection and Tracking for Drosophila Dorsal Closure

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We have developed fast and efficient methods to analyse Dorsal closure in Drosophila that perform better than traditional methods in performance as well. We first analyse the change in the area of the cells in Drosophila using Structured decision forests to make accurate and fast edge detection. Next, we track the vertices of the cells in the amnioserosa using the Kalman filter.

1 Introduction and Related Work

Biologists have analysed the change in the area of cells as it closes, or Dorsal closure for the larger understanding of [1, 2] the forces that drive cell shape changes. Kinematic models and dynamic models[1] have been developed to this end. Recently [2] well-understood tools in Vision have been used to analyse the problem.

However, these methods need the detection and tracking to be optimized over several variables and need explicit point correspondences between consecutive frames. Such methods are not only computationally intense but also slow.

We analyse the system from a purely visual perspective, using machine learning techniques such as structured decision forests and Kalman filter equations (which in fact bear a striking resemblance to biologically motivated state equations) that are very fast and cheap to compute. We hope that this intuitive framework can enable biologists to infer useful information about the system.

2 Method

We detect edges on the Drosophila amnioserosa using Structured Random Forests that are more efficient than traditional edge detectors such as Canny. The Canny detector typically yields threshold-dependent noisy edges, where detecting faint and thick edges would require manual specification of thresholds. Structured Random Forests can be trained to yield edges required for the application at hand. They are also faster, more robust and capture the relationship between neighbouring pixels.

Once we detect edges, we detect the vertices connecting the edges using a corner detector such as Harris, and track them across frames in the video, using the Kalman Filter. Since the area of each cell in the amnioserosa changes to different extents, a separate velocity model is built for each vertex. The cells in the remaining frames can be automatically tracked using this model. A modest set of training images (first 30 frames of the video) is used to obtain the parameters of the velocity model.

2.1 Structured Random Forests

A binary decision tree enables one to test on an attribute or feature at each one of its nodes, and branch to one of two outcomes depending on the value of the input and the test performed at the node. The test could be as simple as a threshold operation.

Random forests are a collection of decision trees, trained using a different parameters for each tree[3]. A typical decision tree, as opposed to a structured decision tree, takes scalar inputs and contains a scalar value at its leaf. For example, the leaf of a typical decision tree that performs edge detection contains the probability that the input pixel is an edge. A structured decision tree takes in

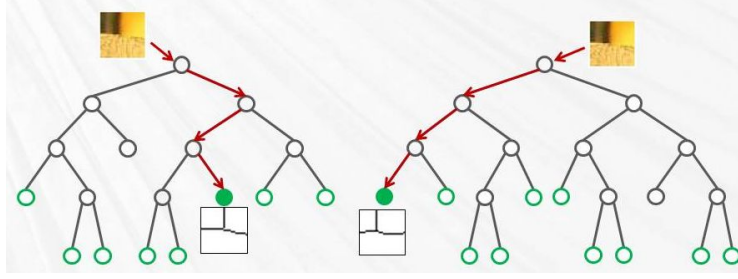


Figure 1: The relationship between pixels is captured in an image patch in a structured decision tree.

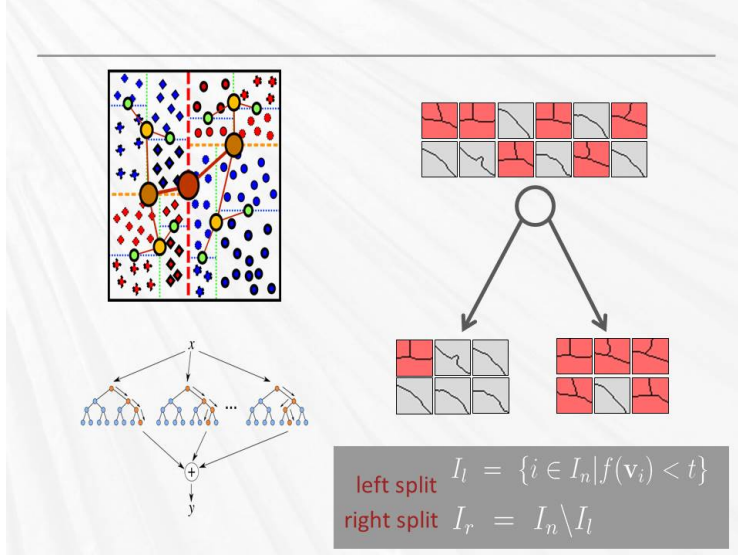


Figure 2: (left) Training and Testing modules of a decision tree (right) The information gain at each node

structured input (patch of an image) and contains structured output (edge patch) at the leaf as shown in Figure1[3]. This method also minimizes computations for each pixel since each leaf contains edge values of a collection of pixels (in the patch) and is thus much faster than its traditional version.

During training, we find parameters of the split function at each node that result in a good split of the data. This corresponds to maximizing the difference in values on either branch of the node. This is defined using the information gain criterion of the form given in Eq1.

During training, the threshold value associated with the split function is determined at each node according to the information gain criteria[3] in Eq1.

$$\Delta E = -\frac{|I_l|}{|I_n|} E(I_l) - \frac{|I_r|}{|I_n|} E(I_r) \quad (1)$$

where $|I_l|$ and $|I_r|$ correspond to the number of elements in the left and right branches respectively from the input set of elements in $|I_n|$. $E(I_l)$ and $E(I_r)$ correspond to the entropy of the elements in the left and right branches.

Computing the information gain in the structured input scenario involves computations over high dimensions, ie 256 dimensions for a 16X16 patch. To perform this operation efficiently, a 2-means clustering of the input patches is done at each node as shown in Figure2.1. Thus, the input image at each node is sent to the left or right branch depending on the cluster it belongs to.

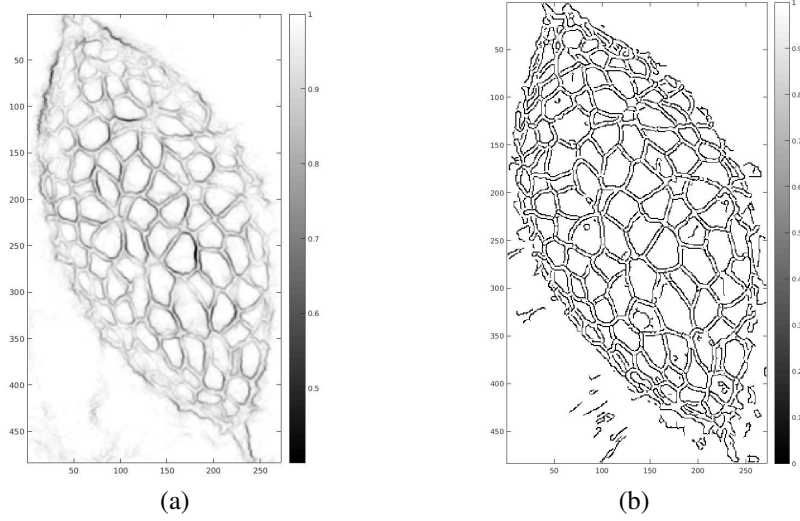


Figure 3: (a) General structured forest (b) Canny Edge detector

2.2 Kalman Filter

After obtaining the edge, we detect vertices on the first 30 frames using the Harris corner detector and associate them with the corresponding points to form the tracks or the measured values. Using the Kalman filter equations below, we build a separate velocity model for each point.

$$x_{t+1} = Ax_t + Bu_t + w_t \quad (2)$$

$$y_t = Cx_t + v_t \quad (3)$$

where $w_t \sim N(0, \Sigma_w)$, $v_t \sim N(0, \Sigma_v)$, and $x_0 \sim N(x_{0|-1}, P_{0|-1})$. Note that $x \sim N(\mu, \Sigma)$ means

$$P(x) = \frac{1}{(2\pi)^{1/2} |\Sigma|^{1/2}} e^{-\frac{1}{2}(x-\mu)\Sigma^{-1}(x-\mu)}. \quad (4)$$

We also have: $Ex = \mu$ and $E(x - \mu)(x - \mu) = \Sigma$.

After obtaining the parameters above, the velocity and therefore location of any point in the *Drosophila* cell can be predicted.

3 Results

4 Discussion

Figure 3 shows the comparison between the output edges obtained with training using Structured forests and that of the Canny edge detector. We observe that the edges are cleaner and more adapted to the *Drosophila* training images. Figure 4 shows edges detected on two frames after a time lag. This shows that our edge-detection is robust to the shrinkage in the cell and therefore conducive to tracking. Figure 5 shows a variation of our method where the edges are detected using smaller window sizes in the training and testing. This enhances the output in a manner similar to contrast normalisation.

We observe from these figures that the edges in *Drosophila* outside the amnioserosa can be detected effectively with low noise and high accuracy at very high speed by modifying the Structured Random forests.

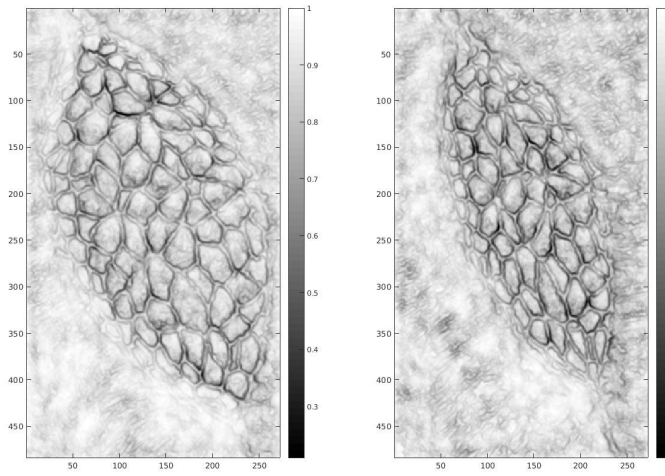


Figure 4: Edge detection on frames 5 and 100. The edges outside the amnioserosa overlap to a good extent and are trackable.

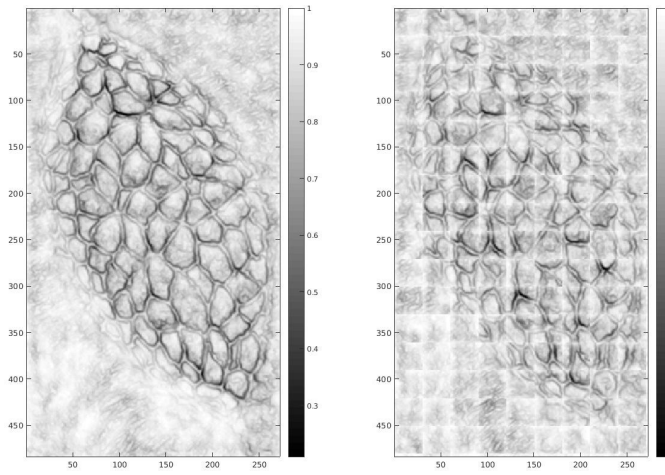


Figure 5: Two variations of our method, where even faint edges appear visible

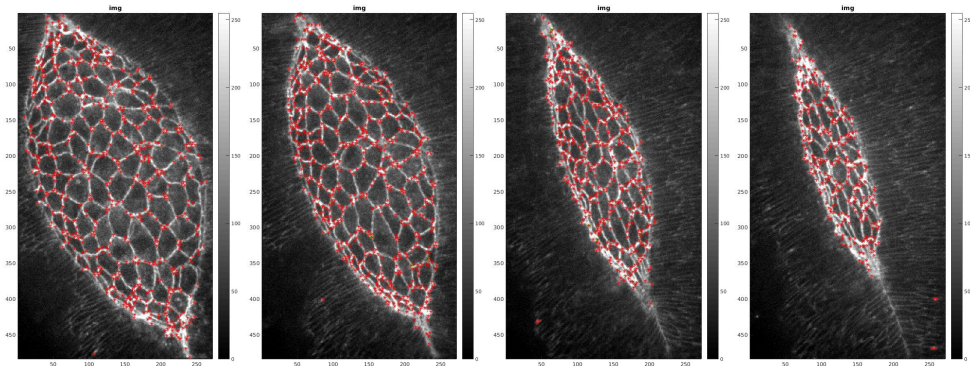


Figure 6: Tracking results at frames 1, 50, 180 and 280 as the cell closes

We observe from Figure6 that the Kalman filter can be used to predict the location of vertices inside the amnioserosa and track them efficiently.

5 Conclusion

We have developed faster methods to enable biologists to infer the kinematics of Dorsal closure in *Drosophila* using efficient and accurate image analysis aided by machine learning tools.

References

References

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