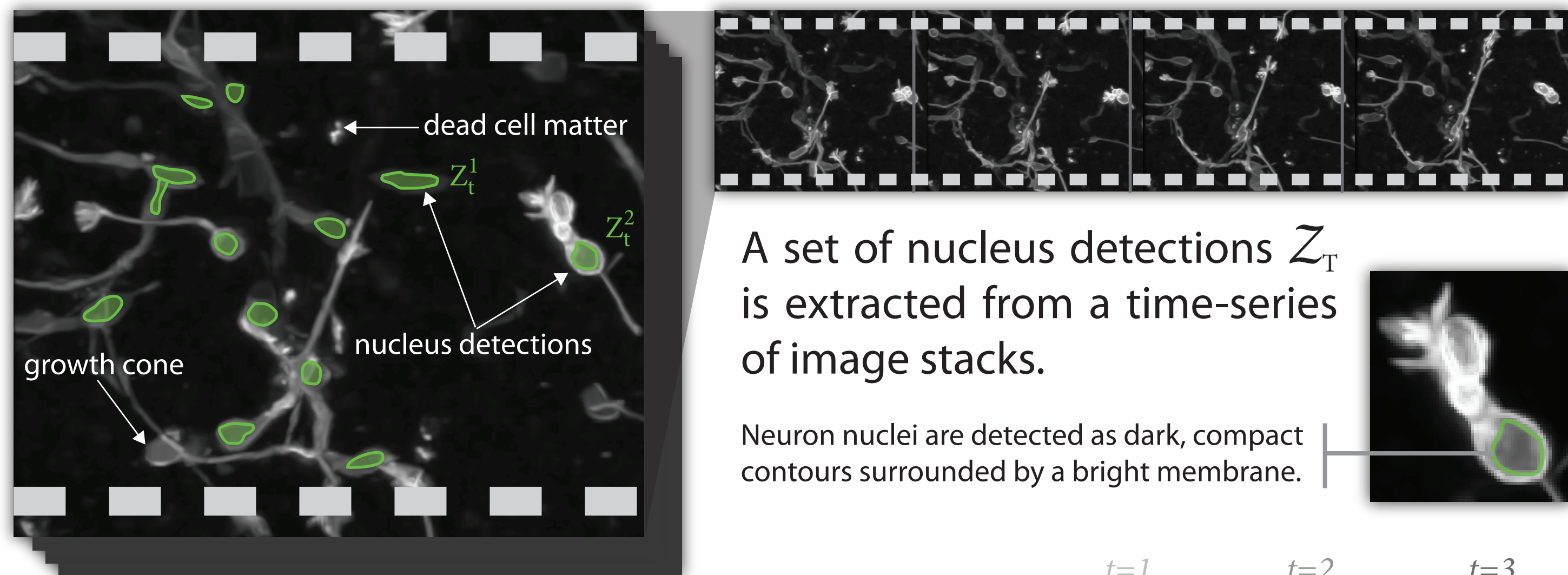


General Constraints for Batch Multiple-Target Tracking Applied to Large-Scale Videomicroscopy

Introduction

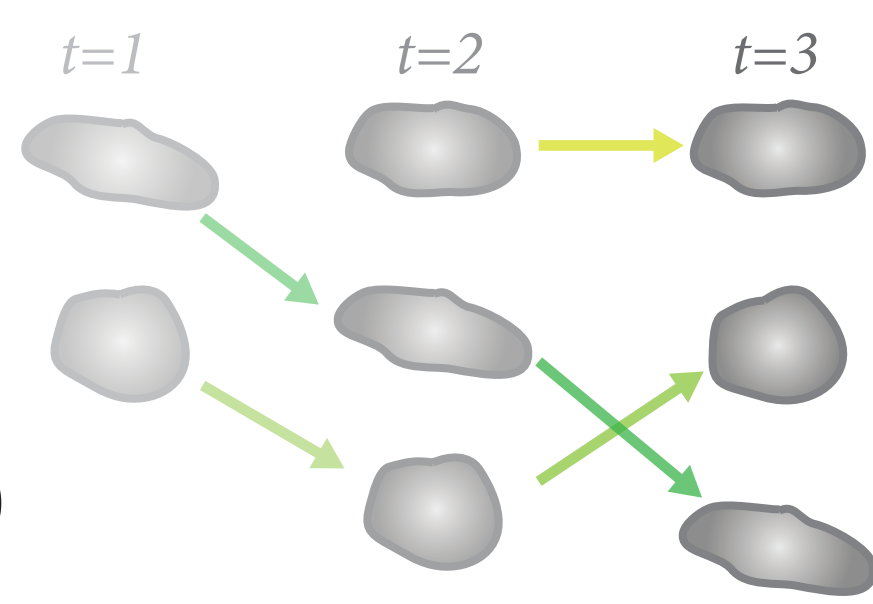
We present a principled probabilistic formalization of batch MTT introducing two general constraints to the tracking problem. The first constraint enforces a learned correlation between a target's appearance and its motion. The second constraint encourages proposed target paths to enter and exit near scene boundaries.

Problem Formulation



Our goal is to link detections Z_T to form the most likely set of target paths \mathcal{X}_T : $\arg \max_{\mathcal{X}_T} p(\mathcal{X}_T | Z_T)$

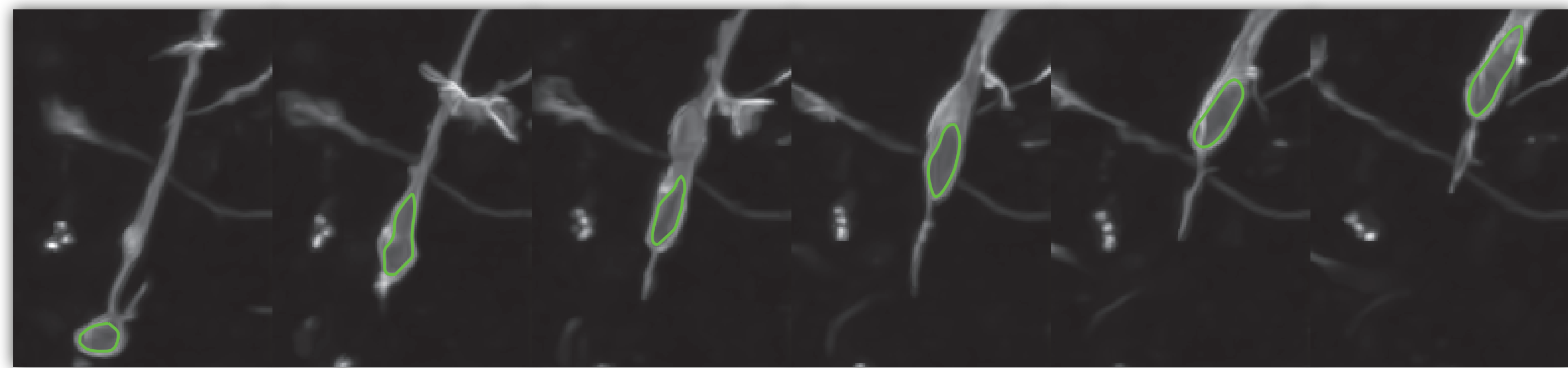
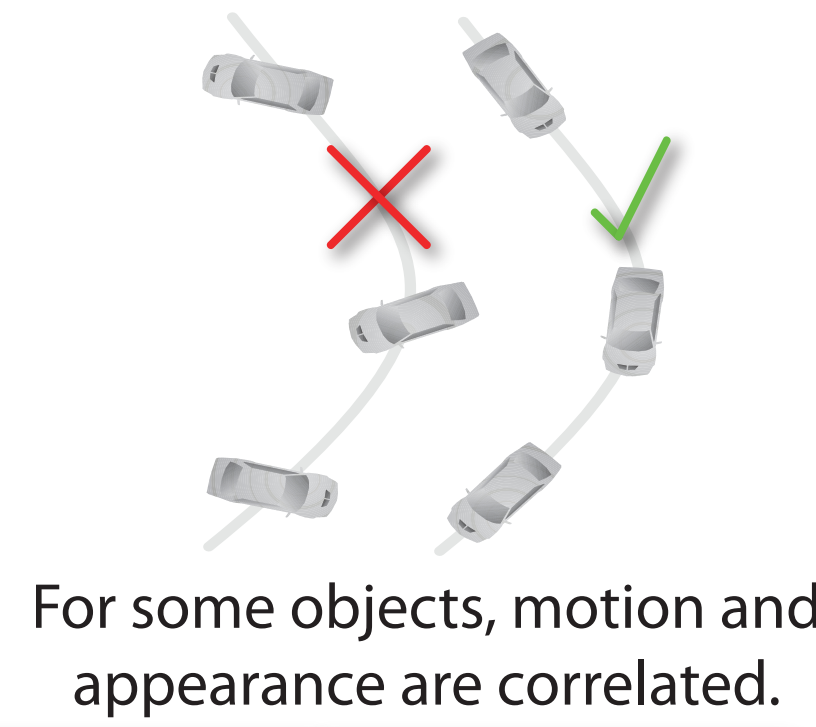
$$p(\mathcal{X}_T | Z_T) \propto p(Z_1 | X_1) p(X_1) \prod_{t=2..T} p(Z_t | X_t) p(X_t | X_{t-1}) \quad (1)$$



The First Constraint

Constraint 1: The movement of a target and its appearance are not necessarily independent.

We relax the conventional assumption that motion is independent from appearance and learn a motion-appearance correlation model to assist tracking.



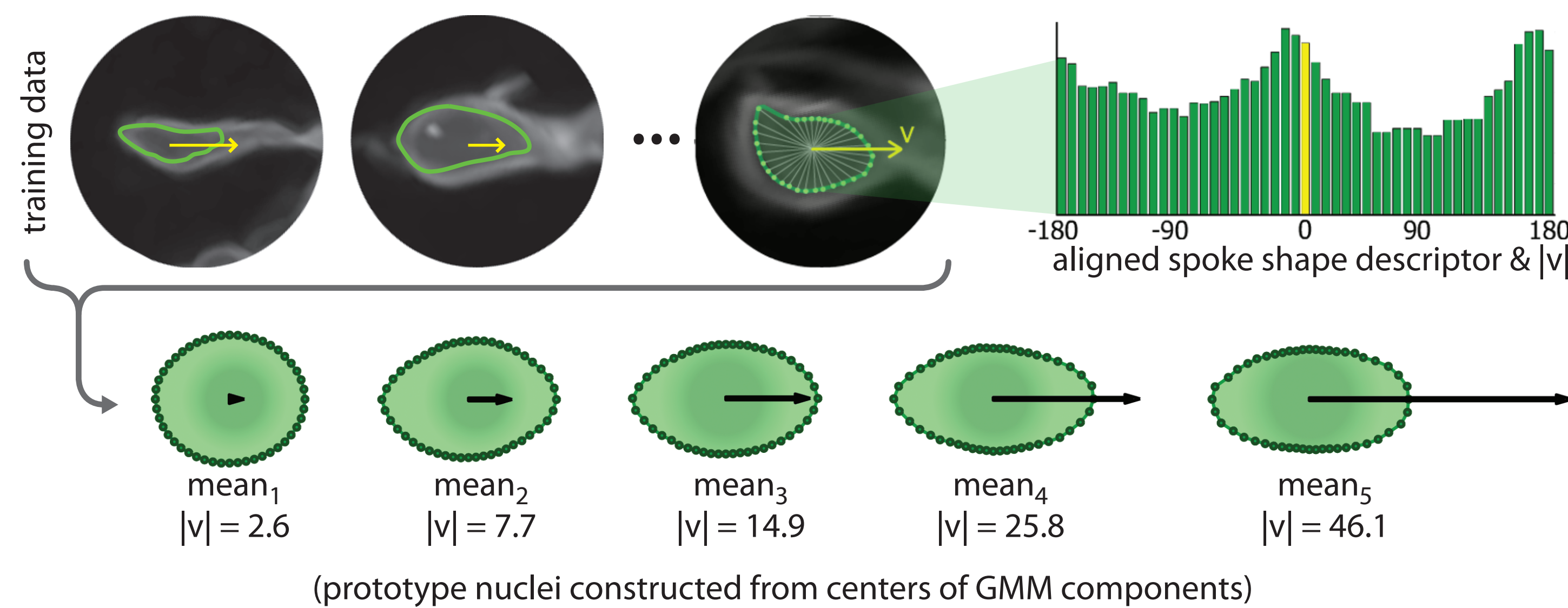
Constraint 1 gives rise to a new term in the observation model from $p(\mathcal{X}_T | Z_T)$

$$p(Z_t | X_t) = \prod_j p(Z_t^j | X_t) = \prod_j p(Z_t^j \text{ is false alarm}) + \sum_i p(Z_t^j | X_t^i) p(X_t^i \text{ created } Z_t^j) + \text{higher order...}$$

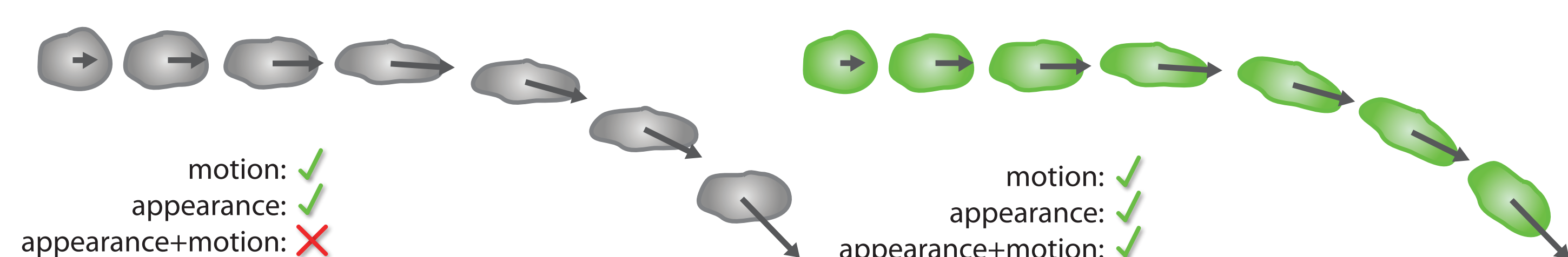
$$p(Z_t^j | X_t^i) = p(L_t^j, A_t^j | M_t^i, O_t^i, R_t^i) \propto p(L_t^j | \text{pos}(M_t^i)) p(A_t^j | O_t^i) p(A_t^j | v(M_t^i))$$

Models the probability of an observed appearance given a motion hypothesis.

$p(A_t | v(M_t))$ is modeled as a 5-component GMM learned from nucleus motion and appearance training data.

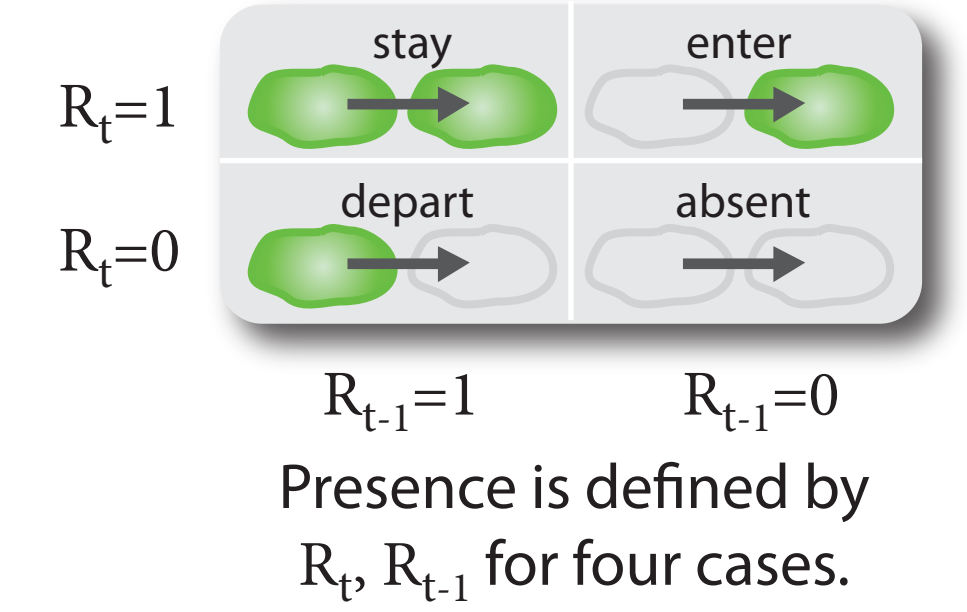


We search for good target paths by evaluating proposed paths according to the motion, appearance, and joint motion-appearance models.



The Second Constraint

Constraint 2: The entrance and departure of a target should occur near a boundary of the scene.



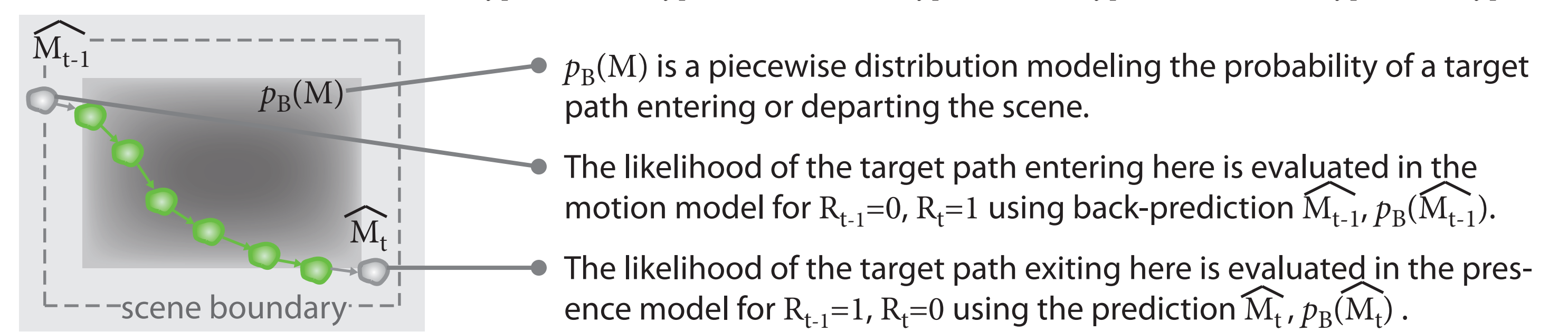
The term $p(X_t | X_{t-1})$ is traditionally limited to a motion model. To enforce Constraint 2, $p(X_t | X_{t-1})$ also depends on the presence of targets at t and $t-1$.

$$p(X_t | X_{t-1}) = \prod_i p(X_t^i | X_{t-1}^i) = p(M_t^i, O_t^i, R_t^i | M_{t-1}^i, O_{t-1}^i, R_{t-1}^i)$$

$$= p(M_t^i | M_{t-1}^i, R_{t-1}^i, R_t^i) \quad p(O_t^i | O_{t-1}^i, R_{t-1}^i, R_t^i) \quad p(R_t^i | M_{t-1}^i, R_{t-1}^i)$$

motion appearance presence

$R_{t-1}=1$	$\mathcal{N}(M_t^i \hat{M}_t^i, \Sigma_M)$	$p_B(\hat{M}_{t-1}^i)$	$\mathcal{N}(O_t^i \hat{O}_t^i, \Sigma_O)$	$\mathcal{N}(O_t^i \bar{O}, \Sigma_{\bar{O}})$	$1 - p_B(\hat{M}_t^i)$	p_e
$R_{t-1}=0$	λ_M	λ_M	λ_O	λ_O	$p_B(\hat{M}_t^i)$	$1 - p_e$
	$R_{t-1}=1$	$R_{t-1}=0$	$R_{t-1}=1$	$R_{t-1}=0$	$R_{t-1}=1$	$R_{t-1}=0$

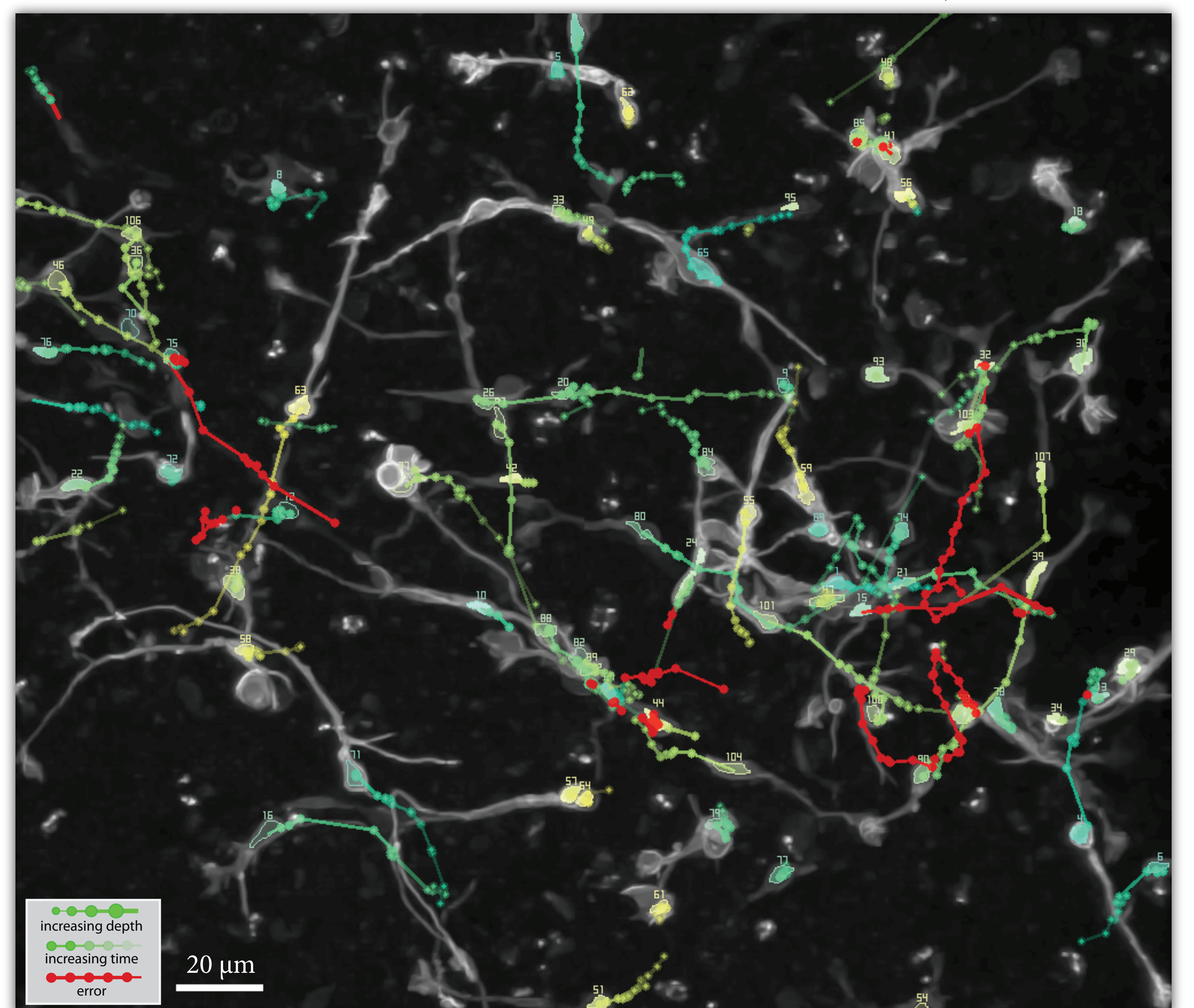
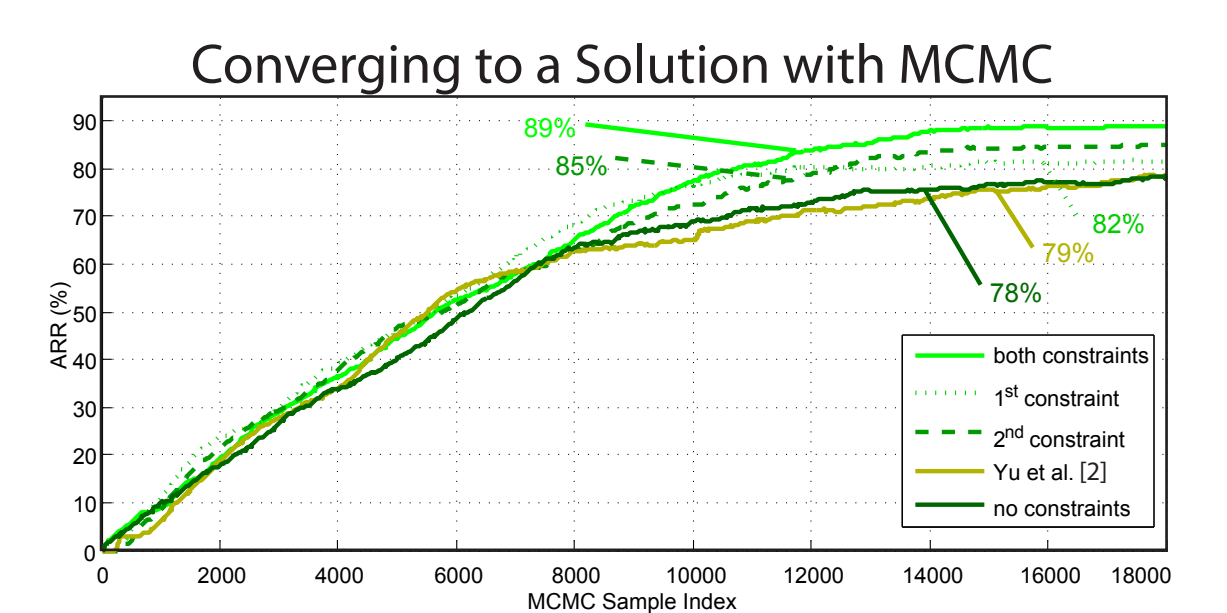
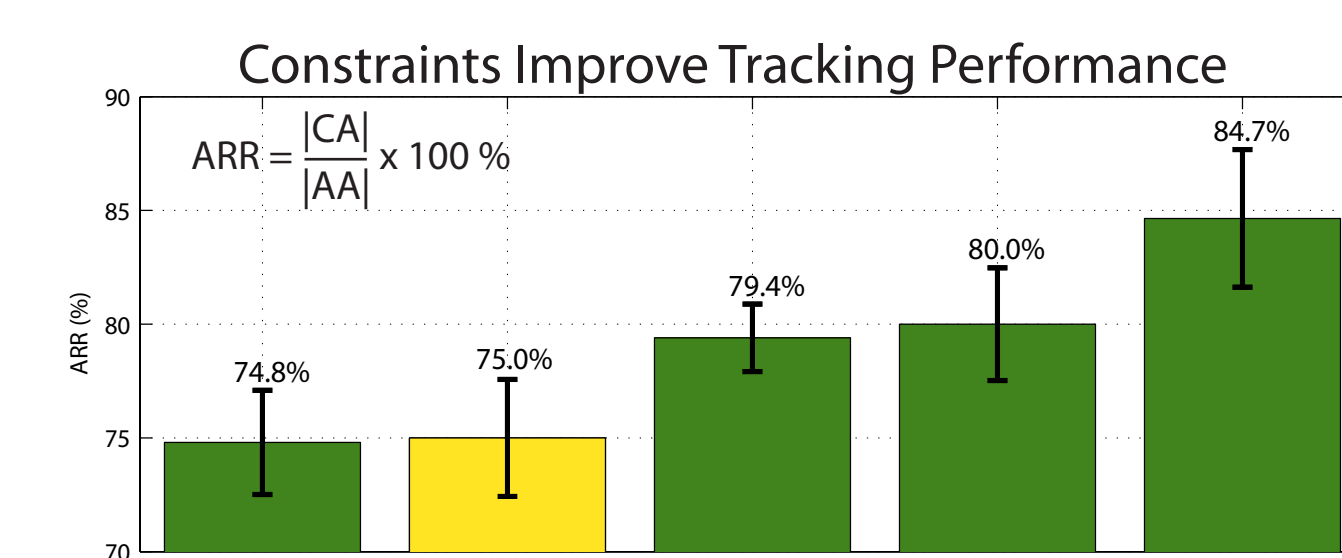
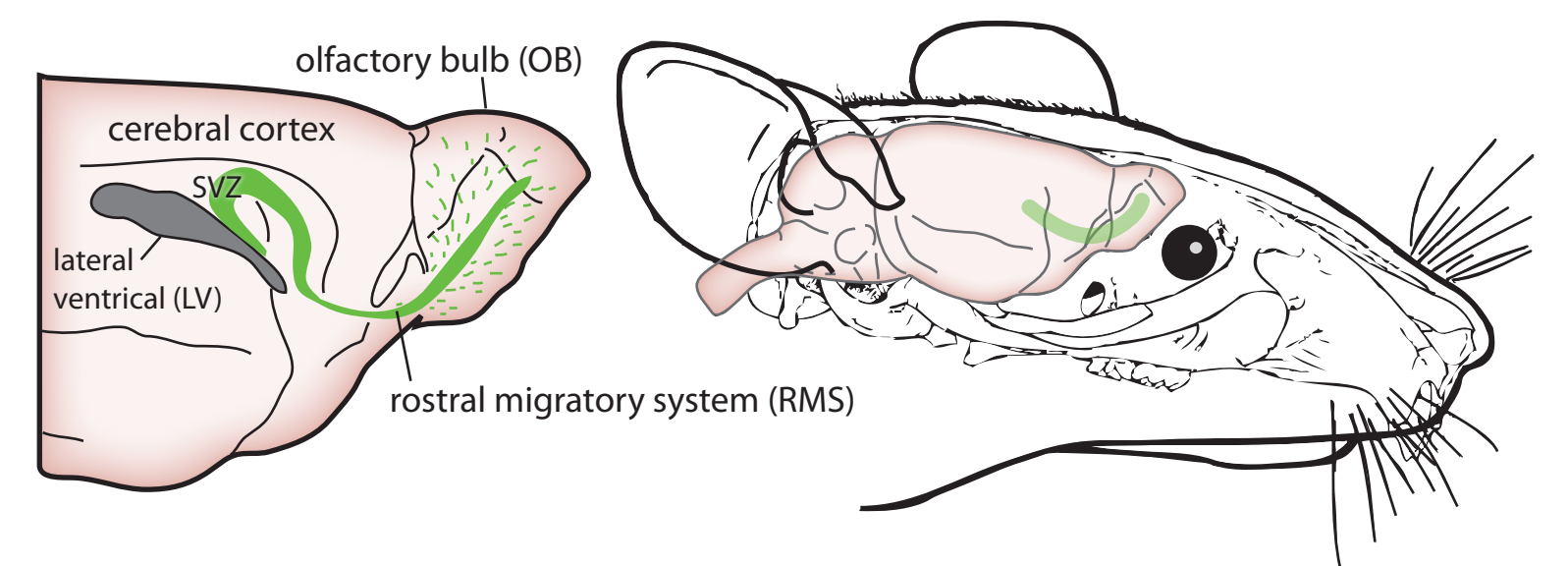


Inference

We use MCMC to efficiently estimate the MAP of (1). We initialize a Markov chain to an empty state and generate new samples by proposing changes to the previous state via a randomly selected MCMC move: *birth, death, associate, dissociate, merge, split, or swap* [1, 2]. The proposed state is added to the chain according to an acceptance probability, otherwise the previous state is added. After generating N samples, the MAP solution is given by the state with this highest posterior, \bar{X}_T .

Results for Neuron Videomicroscopy

We collaborate with neuroscientists studying neuroplasticity. A lentivirus is injected into the SVZ causing newly born neurons to express GFP. We image a $270 \times 270 \times 62 \mu\text{m}$ OB tissue sample with a 2-photon microscope.



Maximum intensity projection with recovered neuron nuclei paths. Errors are shown in red.