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# **Positive-unlabeled convolutional neural networks for particle picking in cryo-electron micrographs**

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## **Background**

Structure determination with cryoEM involves reconstructing a 3D molecule from 2D projections. This process often requires tens to hundreds of thousands of experimental projections, or particles. Locating these particles in cryoEM micrographs, referred to as particle picking, is a major bottleneck in the current protein structure determination pipeline. This pipeline generally consists of sample and EM grid preparation, imaging, particle picking, and eventually structure determination. Labeling a sufficient number of particles to determine a high resolution structure can require months of effort – even with the use of existing methods designed to automate the process. Limitations of these tools include high false positive rates, requiring many hand-labeled training examples, and poor performance on non-globular proteins.

In order to better automate particle picking, and thus accelerate structure determination, we newly frame the particle picking problem as an instance of positive-unlabeled classification. In our framework, for a set of micrographs containing particles of interest with a small number labeled for training, we learn a convolutional neural network (CNN) to classify particles from background using a novel generalized-expectation criteria [1] to regularize the model's posterior over the unlabeled micrograph regions. This advance allows us to achieve state-of-the-art particle detection results with minimal hand-labeling required.

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#### **Methods**

We develop Topaz, the first particle picking pipeline to use CNNs trained using only positive and unlabeled examples and GE-binomial, a general objective function for learning classifier parameters from positive and unlabeled data. The GE-binomial objective penalizes the negative log-likelihood of the labeled data points while regularizing the classifier's posterior over the unlabeled data to match a binomial distribution prior on the number of unlabeled positives. Denoting the set of labeled positive data points by  $P$ , the probabilistic classifier as  $g$ , the classifier's posterior over the number of unlabeled positives as  $q$ , and the binomial prior as p, the GE-binomial objective function is:  $- \mathbb{E} [\log g(x)] + KL(q||p)$ , where KL is the *x* ∈ *P*

Kullback-Leibler divergence.

In the Topaz pipeline, CNN classifiers are fit to labeled particles and the remaining unlabeled micrograph regions using minibatched stochastic gradient descent to minimize the GE-binomial objective. Predicted particle coordinates are next extracted by scoring each micrograph region with the trained classifier and then using the non-maximum suppression algorithm to greedily select candidate particle coordinates.

#### **Results**

We show that the Topaz pipeline is able to accurately detect particles when trained with very few labeled example particles. On the EMPIAR-10096 cryoEM data set [2], Topaz achieves 46% precision at 90% recall with only 1000 labeled particles. In contrast, at the same recall level, EMAN2's byRef method [3] only reaches 33% precision with the same set of labeled particles – corresponding to 71% more false positives than Topaz. Remarkably, Topaz still achieves better precision than EMAN2 at 90% recall with 1/10th and even 1/100th the number of labeled particles. At all numbers of labeled particles tested, we improve substantially over EMAN2's byRef method in area under the precision-recall curve. The relative improvement in particle detection provided by Topaz is even greater on a second, unpublished dataset provided by the Shapiro lab, containing stick-like particles with low signal-to-noise ratio. Furthermore, we show that combining a convolutional decoder with the convolutional feature extractor and classifier learned with GE-binomial to form a hybrid classifier+autoencoder can further improve generalization when very few labeled data points are available. Finally, we demonstrate that our GE-binomial objective function outperforms other positive-unlabeled learning methods never before applied to particle picking. Topaz runs efficiently, training in hours and predicting in seconds with a single consumer grade GPU. We expect Topaz to become an essential component of single particle cryoEM analysis and our GE-binomial objective function to be widely applicable to positiveunlabeled classification problems.

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## **References**

- 1. Mann GS, McCallum A. Generalized Expectation Criteria for Semi-Supervised Learning with Weakly Labeled Data. J Mach Learn Res. 2010; 11:955–984.
- 2. Tan YZ, Baldwin PR, Davis JH, Williamson JR, Potter CS, Carragher B, Lyumkis D. Addressing preferred specimen orientation in single-particle cryo-EM through tilting. Nat Methods. 2017; 14:793–796. DOI: 10.1038/nmeth.4347 [PubMed: 28671674]
- 3. Tang G, Peng L, Baldwin PR, Mann DS, Jiang W, Rees I, Ludtke SJ. EMAN2: An extensible image processing suite for electron microscopy. J Struct Biol. 2007; 166:205–213. DOI: 10.1016/j.jsb. 2006.05.009