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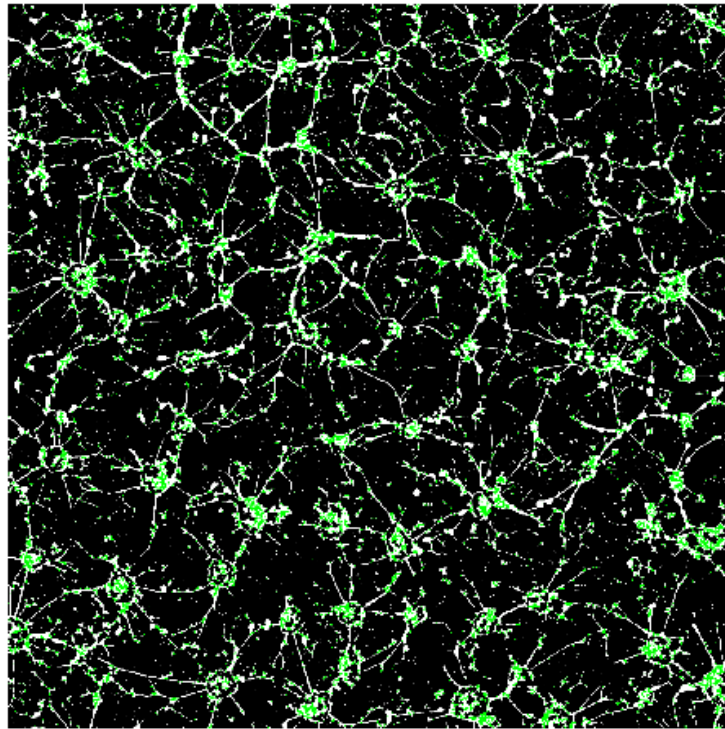
read the image

```
img = imread('resized/preprocessed-t0.tif');  
  
image = img(3001:5000, 6001:8000);
```

preprocess

```
binarize  
  
BW_p = image > 40;  
  
% dilate and erode to connect parts of neurons  
se = strel('disk', 4);  
BW_p = imclose(BW_p, se);  
  
% fill holes in neurons which caused when removing dots  
BW_filled = imfill(BW_p, 'holes');  
fill_differ = BW_filled & ~BW_p;  
% preventing from fill very large holes  
min_area = 200;  
fill = fill_differ & ~filterRegions_area(fill_differ, min_area);  
  
BW_p = BW_p | fill;  
figure;imshowpair(BW_p, image > 40);  
title('fill holes');  
  
Warning: Image is too big to fit on screen; displaying at 25%
```

fill holes



potential neurons

radius range

```
R_center = 65;
R_range = 35;
Neurons = zeros(0, 2);

parfor r = R_center - R_range:R_center + R_range
    % circular kernel: pixels inside circle(r) are 1, outside ones are
    0.
    kernel = circle_kernel(r, 0);

    % conv2
    density = conv2(double(BW_p), kernel, 'same');

    % pixels where conv2 result are larger, which means bright areas
    neu = density > 0.45 * sum(kernel(:));

    % find centroids of these bright areas
    stats = regionprops(neu, 'Centroid');
    len = length(stats);
    Centroids = zeros(len, 2);
    for i = 1:len
        Centroids(i, :) = stats(i).Centroid;
```

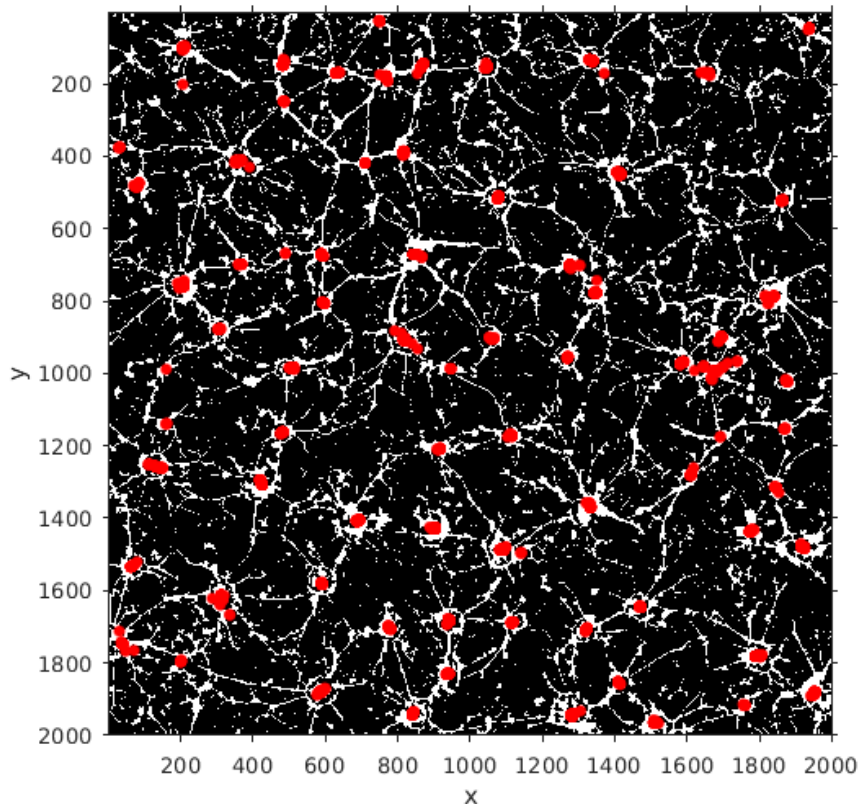
```

end
Neurons = [Neurons; Centroids];
end

% centroids of bright areas are potential neurons
draw_neurons(BW_p, Neurons);

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```



get neurons from centroids

This section tries to select neurons from points we got from lines. The main idea is to calculate numbers of bright angles in an annulus. For each degree x in $0:360$, if there are bright pixels in the annulus whose angles from the center of the annulus equal to x , we say x is a bright degree. For a point p , $A(p, R, \text{annulus})$ draws a annulus around p with radius = R and width = $\text{annulus} * 2$. If number of bright degrees in $A(p, R, \text{annulus})$ is more than $360 * \text{threshold_angle}$, we say point p is a neuron and R is the neuron's size. For every point, we try R in dimension $[R - R_range, R + R_range]$ (that is $45:105$, if using parameters below).

```

% Besides, we use another R_around to determine whether there is a
% bright
% area around the center of neuron, because most neurons are bright in
% the
% center.

threshold_angle = 0.55;
R_center = 70;

```

```

R_range = 45;
R_around = 25;
threshold_around = 0.25;

[final_Neurons, grades, R, around] = IsNeurons_new_4(BW_p,
    Neurons, ...
        'threshold_angle', threshold_angle, ...
        'merge_dis', 2, ...
        'R', R_center, 'R_range', R_range, 'annulus',
    2, ...
        'R_around', R_around, 'threshold_around',
    threshold_around);

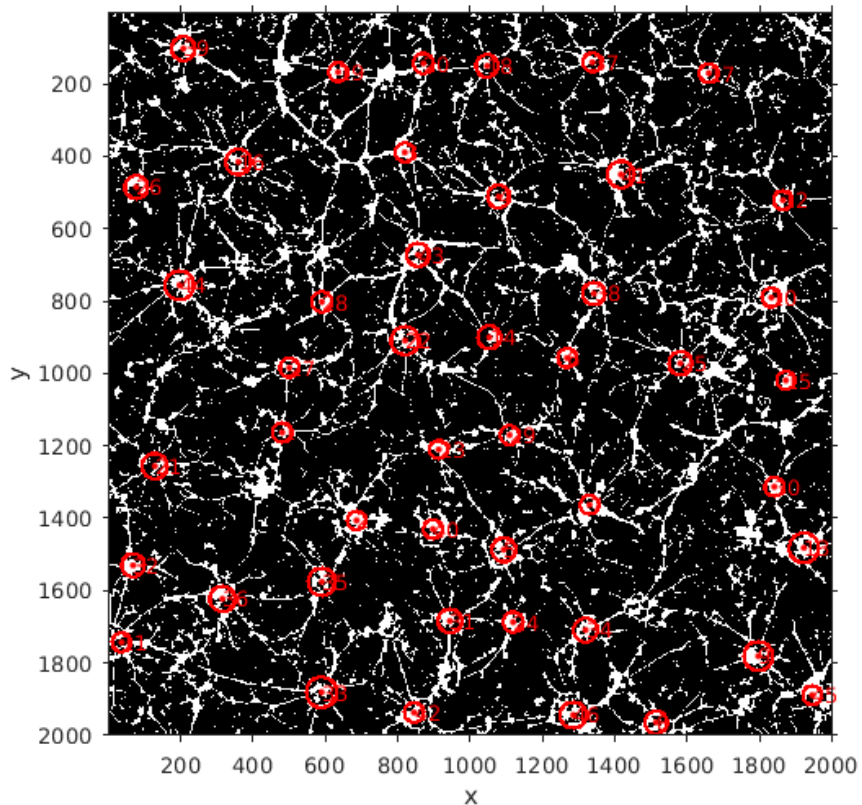
draw_circles(final_Neurons, R, BW_p);

% number neurons
for i = 1:length(grades)

    text(final_Neurons(i,1),final_Neurons(i,2),int2str(i),'FontSize',10,'Color','red')
end

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```



Published with MATLAB® R2018a