

# Barnacle Counting Protocol

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## Abstract

**Citation:** Rebecca Maher Barnacle Counting Protocol. **protocols.io**

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## Protocol

### Step 1.

Open ImageJ in Chevron Visualization Lab computer --> Plugins --> Cell Counter.

### Step 2.

In ImageJ window, click File Open to open raw image file.

### Step 3.

In the Cell Counter click Initialize.

### Step 4.

Review the image to identify *Orbicella franksi* colonies by selecting the Magnifying Glass tool from the ImageJ window to zoom in or out.

### Step 5.

Zoom in on the first colony to 200%.

### Step 6.

Select Text tool from ImageJ window.

### Step 7.

In the Cell Counter window, select Type 2 from Counters. Counter Type 1 is not used because the color is not visible on Viz Lab Screen. Counters can be enlarged from initial settings to be visible on Viz Lab Screen.

### Step 8.

Mark all barnacle holes on a single colony.

### Step 9.

Use the Magnifying Glass to zoom in or out if necessary and use the Scrolling tool to scroll around.

### Step 10.

Once all the barnacles on a single colony have been marked, select Counter Type 3 and scroll to the next

colony.

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**Step 11.**

Repeat with all remaining colonies. If the Counter color is not visible select Type 4 or Type 7 (deep reds).

a. We recommend changing counters between colonies to avoid large stretches of barnacle counts (>1000) that can increase the chances of accidental data loss.

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**Step 12.**

For each image, record the barnacle count on paper.

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**Step 13.**

Click Results in Cell Counter window. In Results window, click File --> Save as marker count. Label with PhotoIDresults to export results.

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**Step 14.**

Click Export Image in Cell Counter window. In ImageJ window, click File --> Save as --> JPEG. Label with PhotoIDvizphoto.

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