



# Fluorescent Focus Unit Assay using LICOR Imaging System

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#### **Abstract**

Indirect immunofluorescence infectivity assay for reovirus using LICOR Imaging System

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### **Guidelines**

This assay is standardized to for 96-well plates using reovirus polyclonal antiserum. New batches of polyclonal antiserum or monoclonal antibodies should be tested to determine optimal dilution.

### **Before start**

Plates and reagents:

Corning Black with clear bottom TC treated 96 well plates (Corning #3904)

DRAQ5 #4084 from Cell Signaling

Sapphire 700 #928-40022 from LICOR

Antibody 800 from LICOR (Life Technologies ones don't work for this)

### **Protocol**

#### Step 1.

Remove media from wells.

#### Step 2.

Wash 1x with Phosphate Buffered Saline (PBS).

#### Step 3.

Add 100 uL ice-cold methanol per well.

### Step 4.

Store plate at -20°C for at least 30 min (can remain at -20°C for several weeks).

### Step 5.

Remove methanol, allow plate to come to room temperature, and excess ethanol to evaporate.

### Step 6.

Wash wells with 150 ul PBS

#### Step 7.

Add 150 uL Dulbecco's PBS (DPBS) with 0.5% Tween-20 (DPBS-T) to each well.

### Step 8.

Remove immediately.

### Step 9.

Add 50 uL rabbit anti-reovirus polyclonal antiserum (1:1000 dilution) in DPBS with 1% BSA (DPBS-BSA) to experimental wells.

Add 50 uL DPBS-BSA with no antibody to background control.

### Step 10.

Incubate for 1 h at 37°C.

# Step 11.

Remove DPBS-BSA.

### Step 12.

Wash 3x with DPBS-T for 5 min each wash while shaking.

### **Step 13.**

Add 50 uL of DPBS-BSA to all wells.

### **Step 14.**

Incubate 1 h at 37°C.

### Step 15.

Wash plates 3x with DPBS-T for 5 min each wash.

### **Step 16.**

Prepare cell staining solution in DPBS-BSA:

Secondary antibody (e.g. Goat Anti Rabbit Alexa 488) - 1:1000

Draq5 - 1:10,000

Sapphire 700 - 1:1000

### Step 17.

Add 50 uL cell staining solution to all wells that were treated with primary.

Add 50 uL of DPBS-BSA with secondary antibody (1:1000) only to background control wells.

### **Step 18.**

Remove solution.

### Step 19.

Incubate 1 h at 37°C.

### Step 20.

Wash 3x with 150 uL DPBS-T.

# Step 21.

Add 50 uL of water.

### Step 22.

Scan plates on LICOR Odyssey Imaging System.

- a. Focus offset = 3.0 (depends on the plates)
- b. 700nm Intensity = 6.5
- c. 800nm Intensity = 7.5

### Step 23.

### **Plates and reagents:**

Corning Black with clear bottom TC treated 96 well plates (Corning #3904)

DRAQ5 #4084 from Cell Signaling

Sapphire 700 #928-40022 from LICORAntibody 800 from LICOR (Life Technologies secondary antibodies give extra background and should be avoided)