Sample Preparation for Flow Cytometry

Introduction

Materials

- > Versene
 - > Clear liquid on Brian's side of drawer
- > Trypsin
 - > Remember to put Trypsin and cell media into dry bath to warm them up before starting
- > HEK cell media
- Flow cytometry tubes
 - > Tubes with Blue filters on top
 - > One for each well
- > Ice Bucket
- > Glass pipette tips
- > P200 and P1000 pipette tips

Procedure

Preparing Samples

- 1. Retrieve cells from incubator
- 2. Look at the wells under the EVOS microscope by the Lu Bench to check the cells and look for fluorescence
- 3. For each well:
 - Aspirate the media (make sure the tip does not touch the bottom of the well)
 - Pipette 200 uL of versene slowly down the side of the well
- 4. Again, for each well:
 - Aspirate the versene
 - Pipette 200 uL of trypsin and wait three minutes
- 5. Add 500 uL of media to each well
- 6. Label flow cytometry tubes and put in an ice bucket

| 7. Pipette up and down in circles, and then add entire mixture of media and trypsin to a flow cytometry tube |
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| through the filter 8. Go find Brian, and he will load your samples into the cytometer |
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