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Perfusion Live Microscopy Using Zeiss LSM 780 and Ibidi Perfusion Sets V.1

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protocol .

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Step by step protocol for setting up live microscopy experiments with Ibidi perfusion sets

Emir Bora Akmeriç 2021. Perfusion Live Microscopy Using Zeiss LSM 780 and Ibidi Perfusion Sets. **protocols.io** https://protocols.io/view/perfusion-live-microscopy-using-zeiss-lsm-780-and-b2exqbfn

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Cell Seeding 1h

1 Check whether HUVECs in T25/T75 are confluent

2 Gelatinize 2 or 3 Ibidi 0.4 luer u-slides with 0.2% gelatin in water 5m

3 Bring trypsin, PBS, media and FBS to to 37C inside cell culture incubator 25m

4 Trypsinize dish and count cells. A minimum of 500k cells are needed for 2 slides 15m

5 Seed slides with HUVECs at a density of 2.2 million/mL, with 100 uL volume 10m

6 Add 120 uL of EGM2 media with antibiotics 20 to 30 minutes after seeding. 5m

7 Put one pair of male luer couplers, 2 sets of syringes and 1 set of male extenders(for live microscopy, there are some on my bench. Basically 2 tubes inside a pipette tip box) inside the incubator for overnight degassing

Live microscopy prep 1h 10m

8 In a 2 mL eppi, dilute 1:1000 sir-DNA(in -20C, there should be two tubes in 2 seperate boxes. One box has 2 tubed labeled as sirDNA but only one of them is the real one(not sure why). Just make sure to get a tube with a colored liquid in it) in CO2 independent medium(transparent, should be a falcon in 4C. If not, there is also a stock you can add Gentamicin to) 10m

9 Prepare one set of ibidi flow unit by adding 14 mL of CO2 independent medium. Extend the male coupling with the tubes in the extra coupling box. You should have slightly elongates tubes in a tube>male adaptor>female joiner>extra tube>male coupler>luer female coupler order 30m

10 Aspirate medium from the slides and add 250 uL of previous mix into each slide. Incubate at 37C for 1 hours 1h

11 Bring the perfusion pump, laptop, 2 medium sized pipette tip boxes and the flow unit to the microscopy room(use the carts)

12 Set up the flow unit inside the incubator while setting the pump and the incubator outside. The best way to do this is putting two pipette tip boxes on the bottom right part of the chamber and then setting up the flow unit on top of these boxes, making sure that the setup is stable. Continue with connecting the air tube and electric cable, there is a stage exit for such cables. Turn on microscope and incubator and open ZEN. Calibrate, pinch test and start running at 37C but **without** CO2 25m

13 Aspirate medium from slides again and then add 250 uL CO2 independent media without sirDNA. 10m

14 Bring one slide and a plastic clamp to the microscopy room. While clamped carefully connect the slide to the unit. 15m

15 Wipe both sides of the slide with isopropanol as well as the objective 5m

On ZEN, withdraw the imaging setup from an image in Anna/210709_Wt_flow 5m

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17 Turn on Definite Focus. Click Find Surface/Focus 5m

18 Check focus on Live and make necessary manual focusing adjustments and click store focus 5m

19 Check whether definite focus is on for focus strategy, enable definite focus for every tile scan 5m

20 Check that Tile scan is set to 3x3 and that a 250+ frame timed capture will be done, change autosave to stream 5m

21 Start experiment. If possible, check whether everything is in focus after the first 10 minutes 10m

Post microscopy(next day) 1d

22 After 24 hours, dismantle the flow unit. You can bring the unit with the it to the 4C room and I can take it apart and clean afterwards. 1d