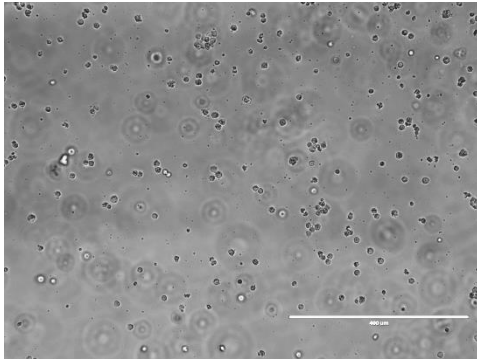
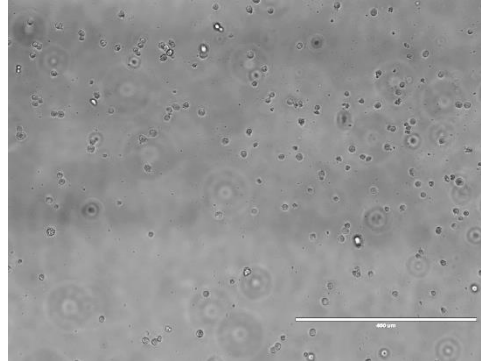


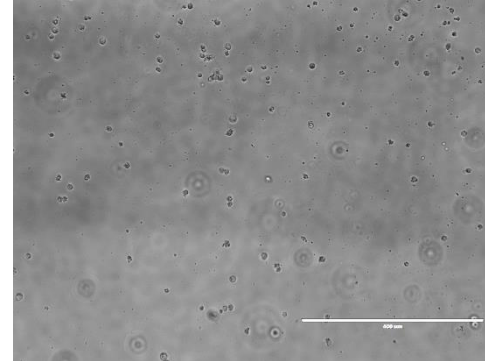
(fig 6b extended) Wider field view of FoP MK iPS cells trapped in the collagen support (4 million cells used)



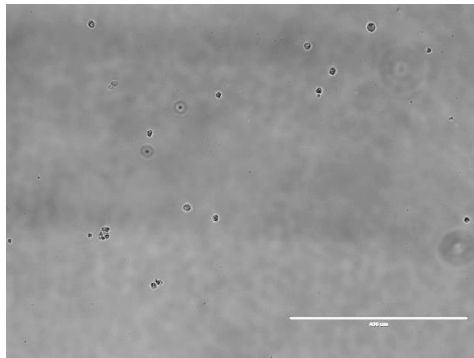
IN



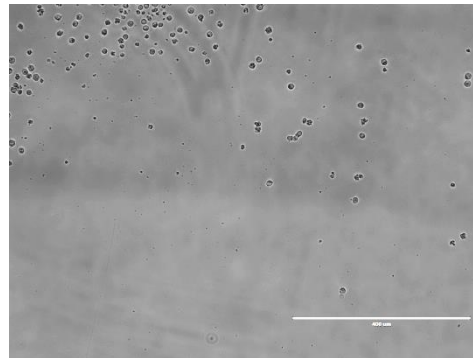
OUT



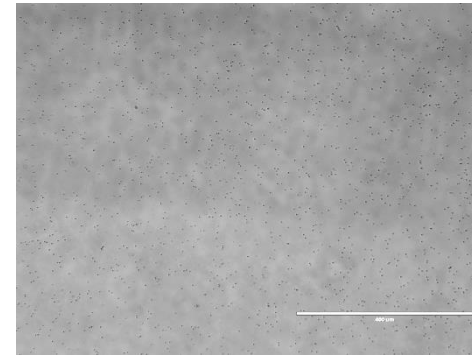
W1



W2



W3



OUTON

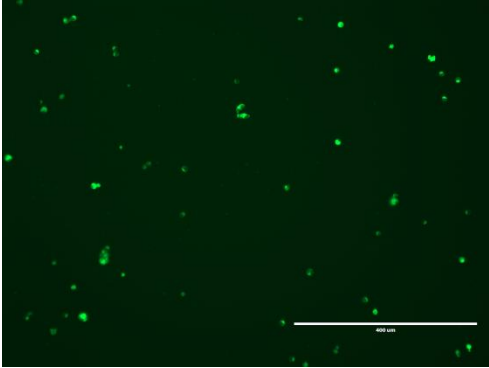
**Figure 6 extra info on outflow (filtrate)**

FoP Mk iPScells / platlet particles phase contrast microscopy

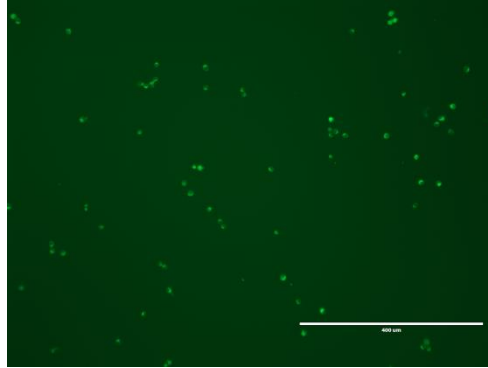
O/N 100 sample 500 PFA 24 well plate

Key: Cells IN, solution OUT filtrate, 3 washes through sponge, OUTON out overnight collection.

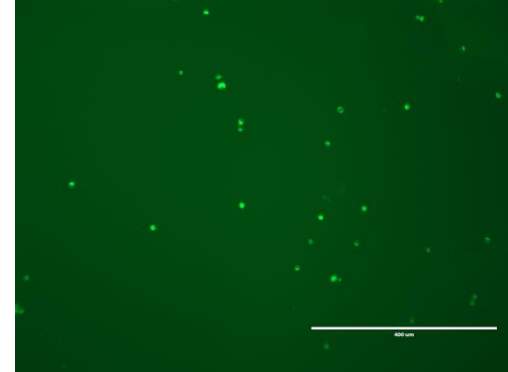
# Bioreactor run, 400K Mk Fop IPSC



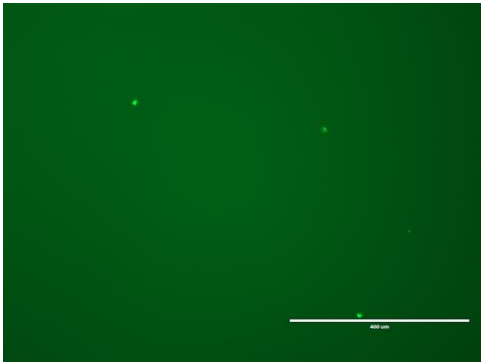
IN



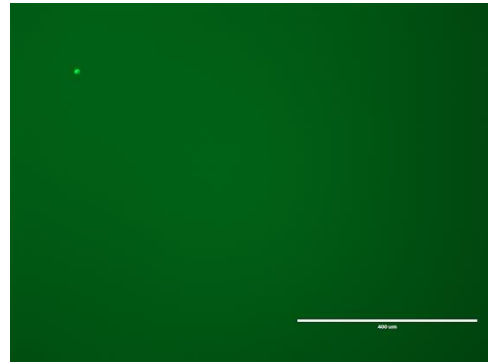
OUT



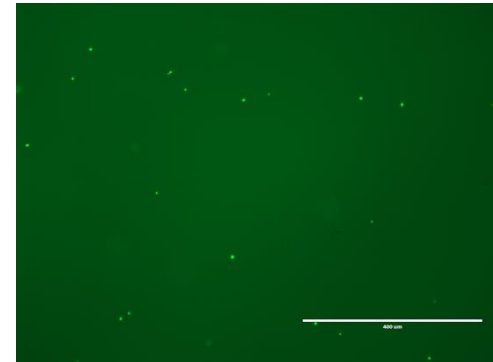
W1



W2



W3

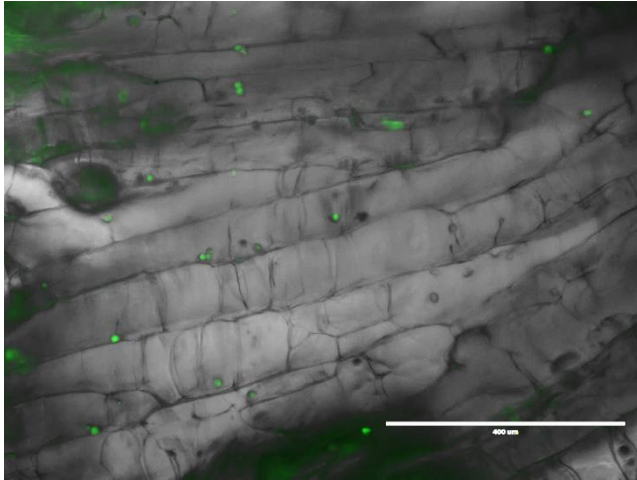


OUTON

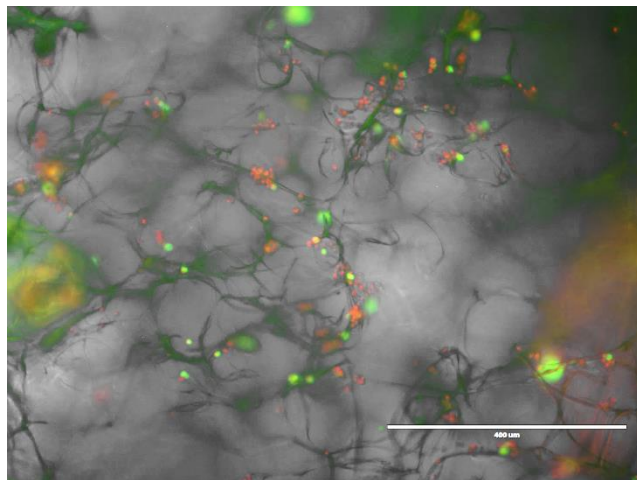
100ul of sample to 500ul of filtered 4% PFA, 24 well plate, left to settle before imaging  
DIOC6

# Bioreactor run, layered collagen filter, 400K BOBC

Note: not sectioned, image taken *after* bioreactor run



Upper sponge layer  
Calcein stain plus PI

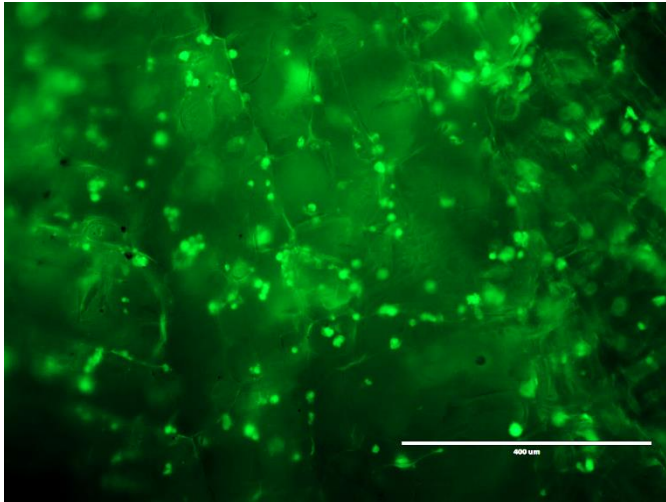


lower sponge layer  
Calcein stain plus PI

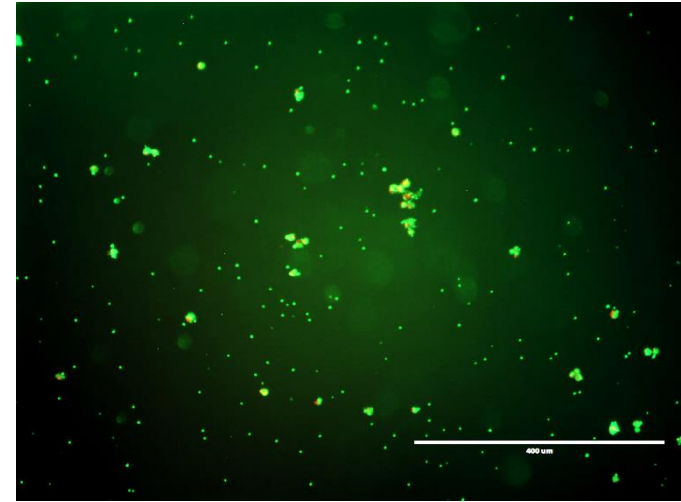
extra images showing MK FoP iPS cell trapping in collagen supports

Bioreactor run, layered collagen filter, 400K BOBC

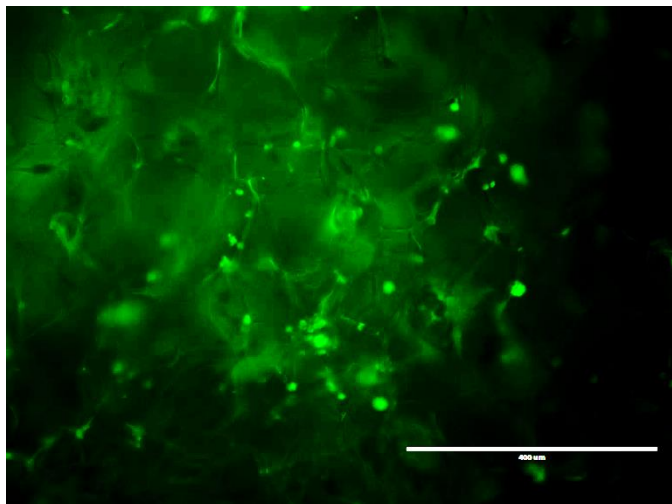
Note: not sectioned, image taken *after* bioreactor run



Upper sponge layer  
DIOC6 stain



Washed out of sponge  
after staining

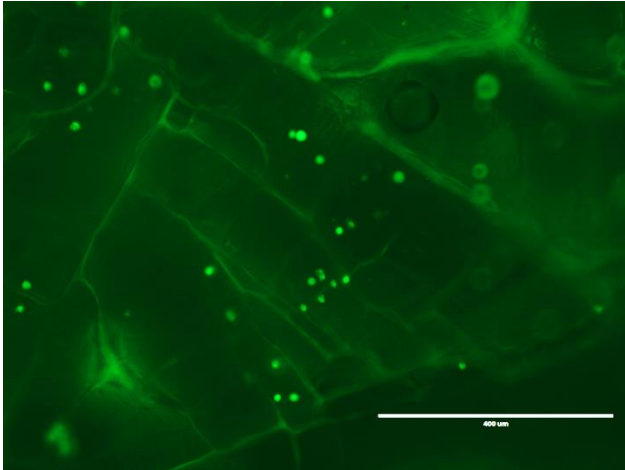


lower sponge layer  
DIOC6 stain plus PI

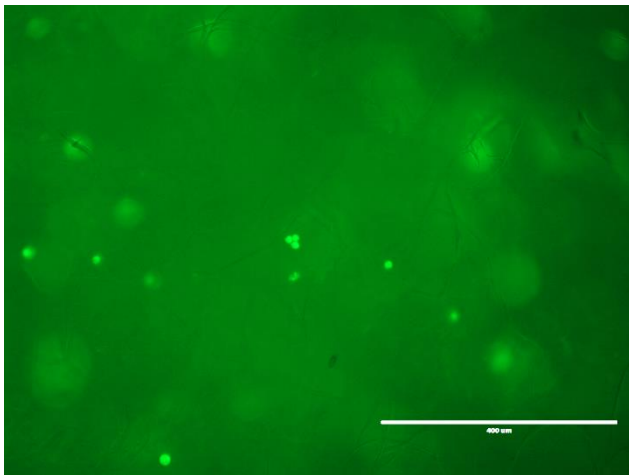
(Note: some washed out during staining)

# Bioreactor run, layered collagen filter, 400K BOBC

Note: not sectioned, image taken *after* bioreactor run



Upper sponge layer  
CAL only



lower sponge layer  
Cal only

(Note: some washed out during staining)