



Sergiy Kyrylenko  
kyrylenk@gmail.com  
098 024 6482

---- supported by the Erasmus+ JM program ----

---- supported by the Horizon 2020 program ----





# КУЛЬТУРА КЛІТИН ЯК ІНСТРУМЕНТ ДЛЯ НАУКОВИХ ДОСЛІДЖЕНЬ ТА ПРАКТИЧНОГО ВИКОРИСТАННЯ



<https://www.shimadzu.eu/cell-culture-analysis>



Jean Monnet  
Programme



HORIZON 2020

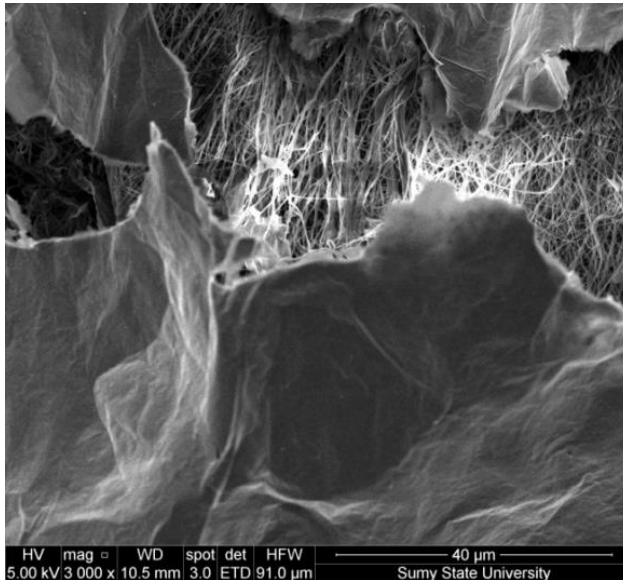
---- supported by the Erasmus+ JM program ----

---- supported by the Horizon 2020 program ----

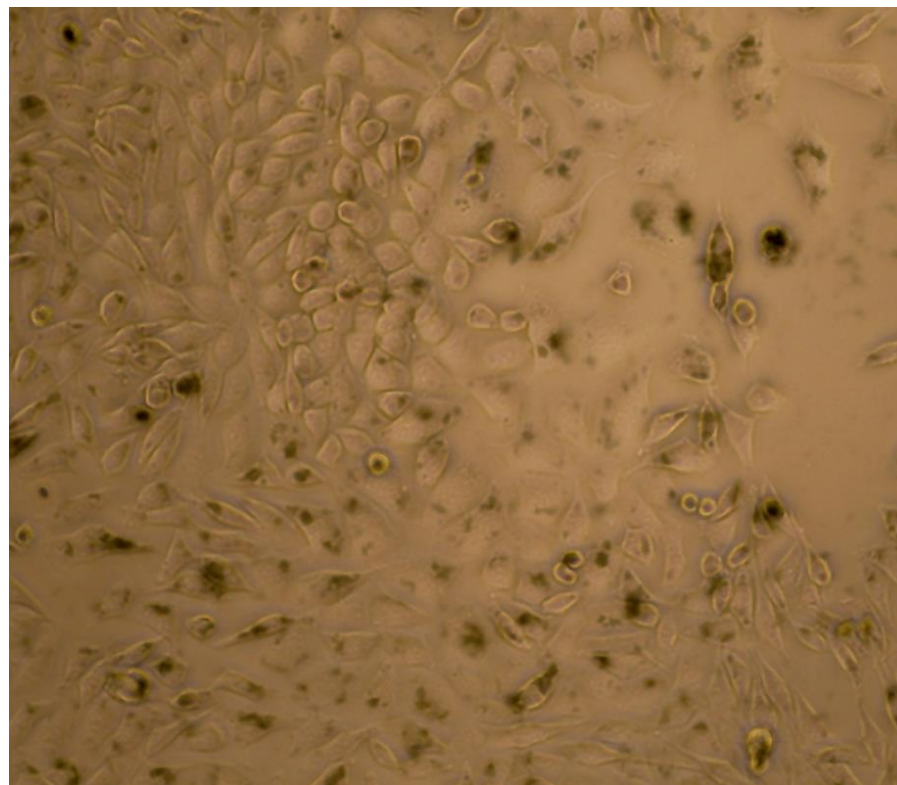
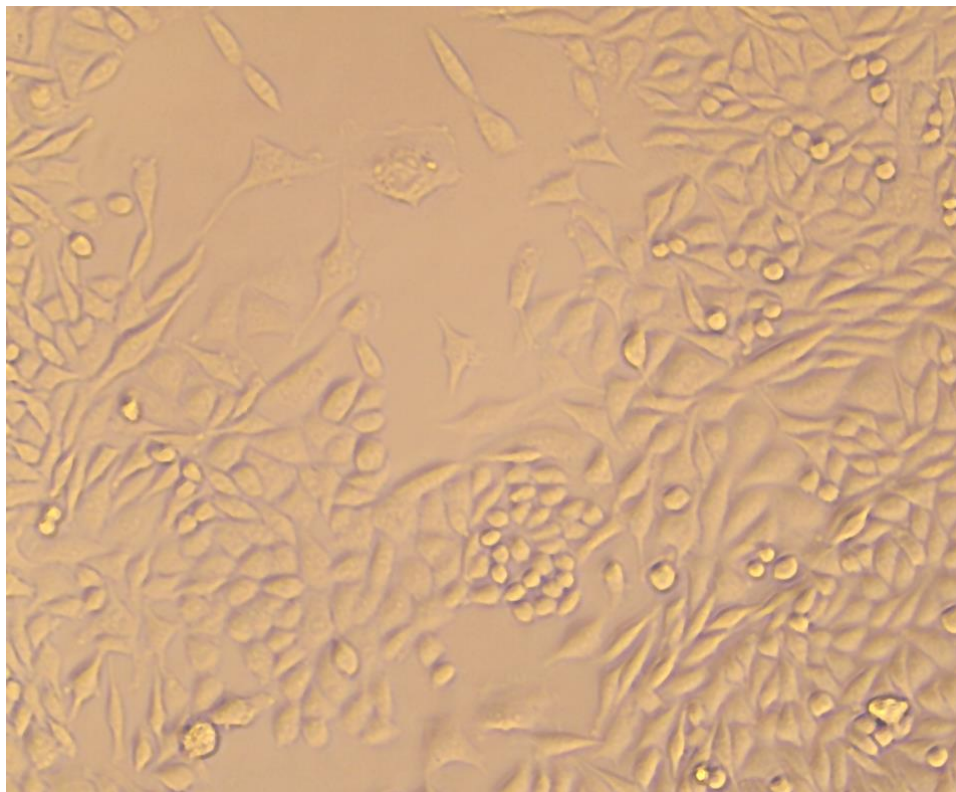
# Biomedical Research Center

## Medical Institute of Sumy State University

- Cell culture laboratory
- Microbiological laboratory
- ELISA laboratory
- Chemical and toxicological laboratory







**ZMB**

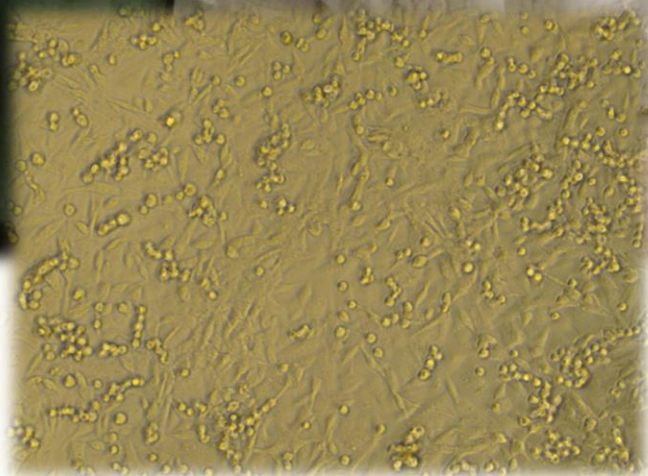
Zentrum für Medizinische Biotechnologie  
Center of Medical Biotechnology



Jean Monnet  
Programme

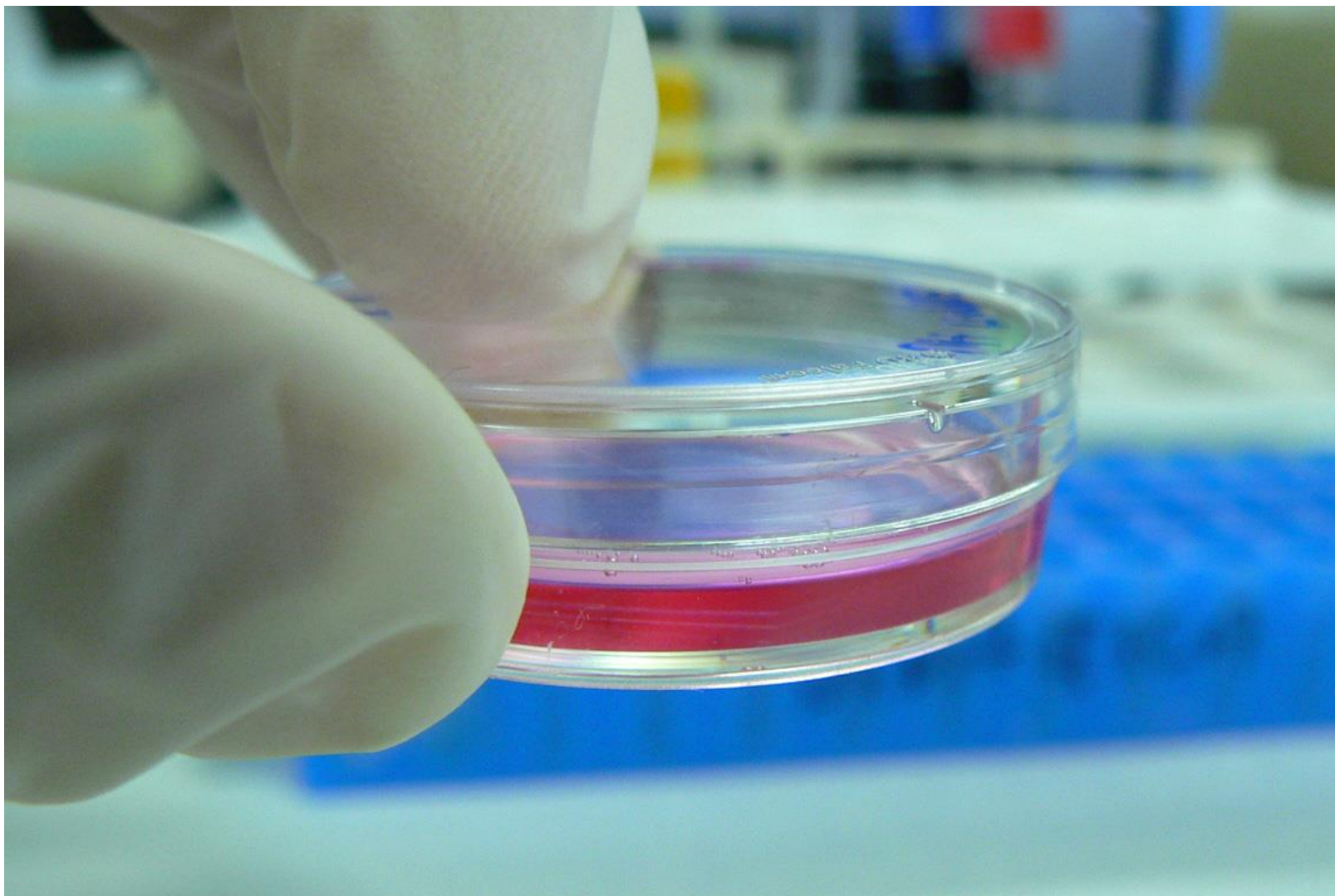






Jean Monnet  
Programme

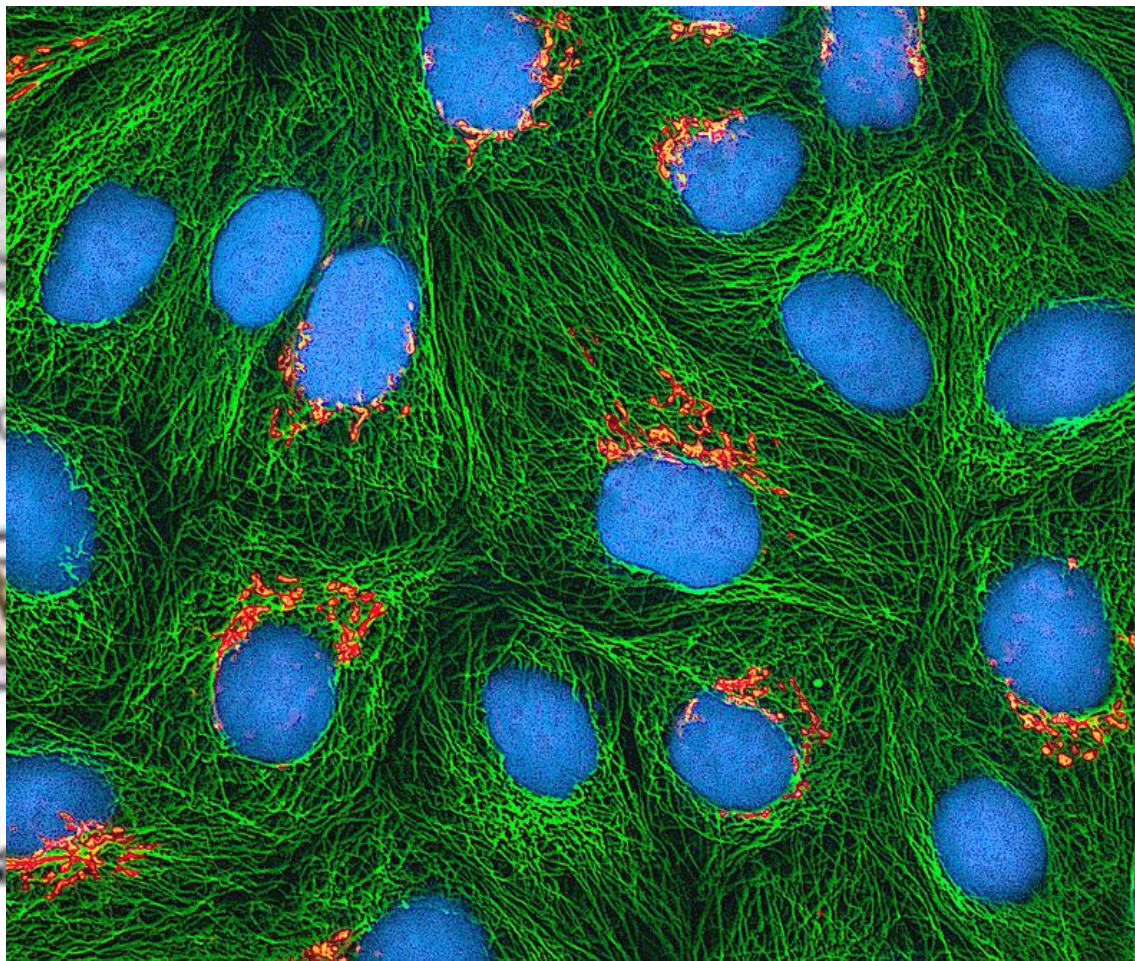
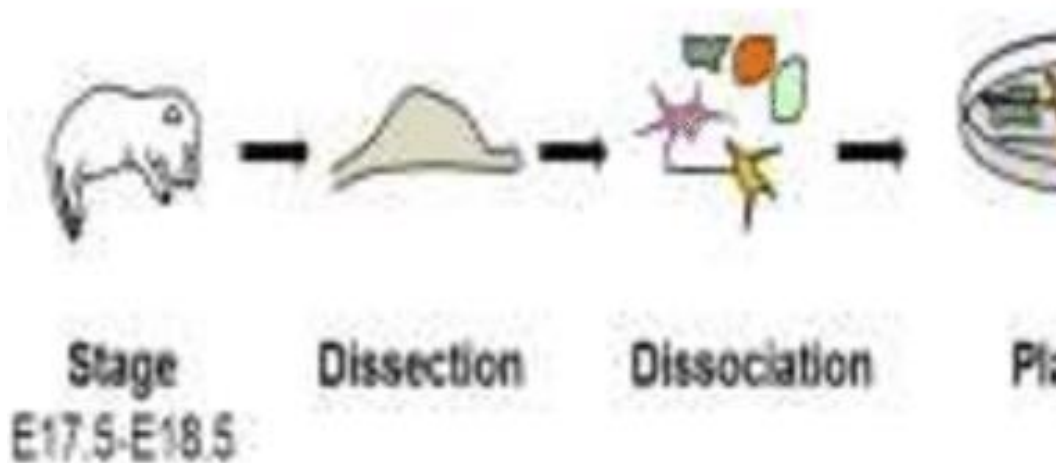
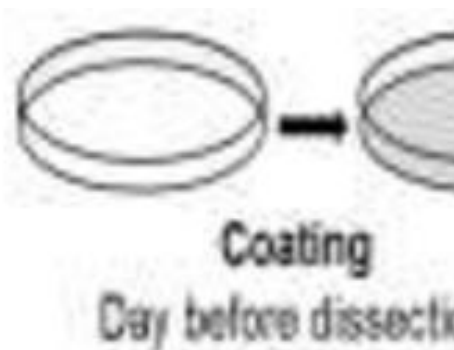






# Primary cell cultures

# Established cell lines



HeLa - cervical cancer cells



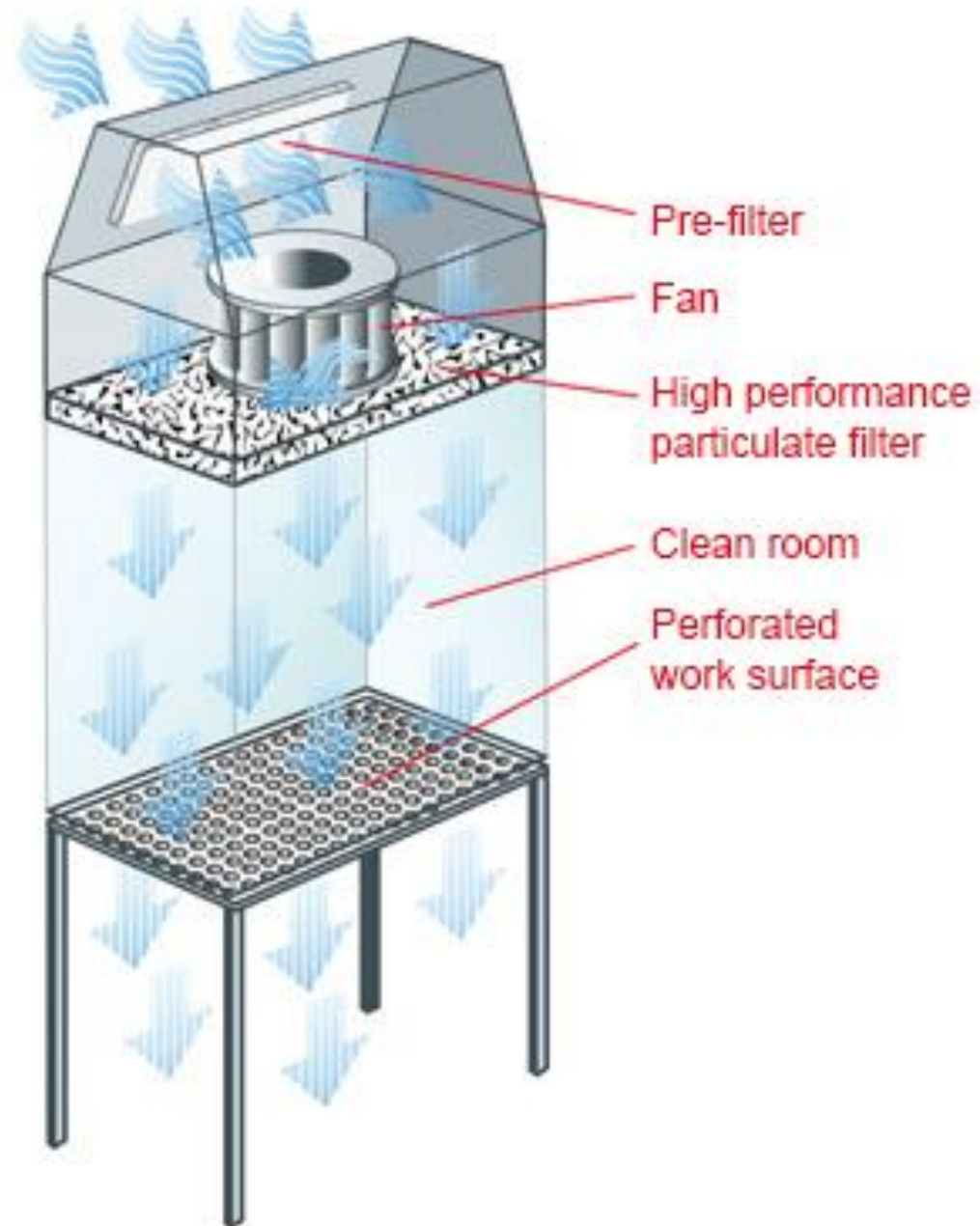














Jean Monnet  
Programme















Jean Monnet  
Programme













Jean Monnet  
Programme













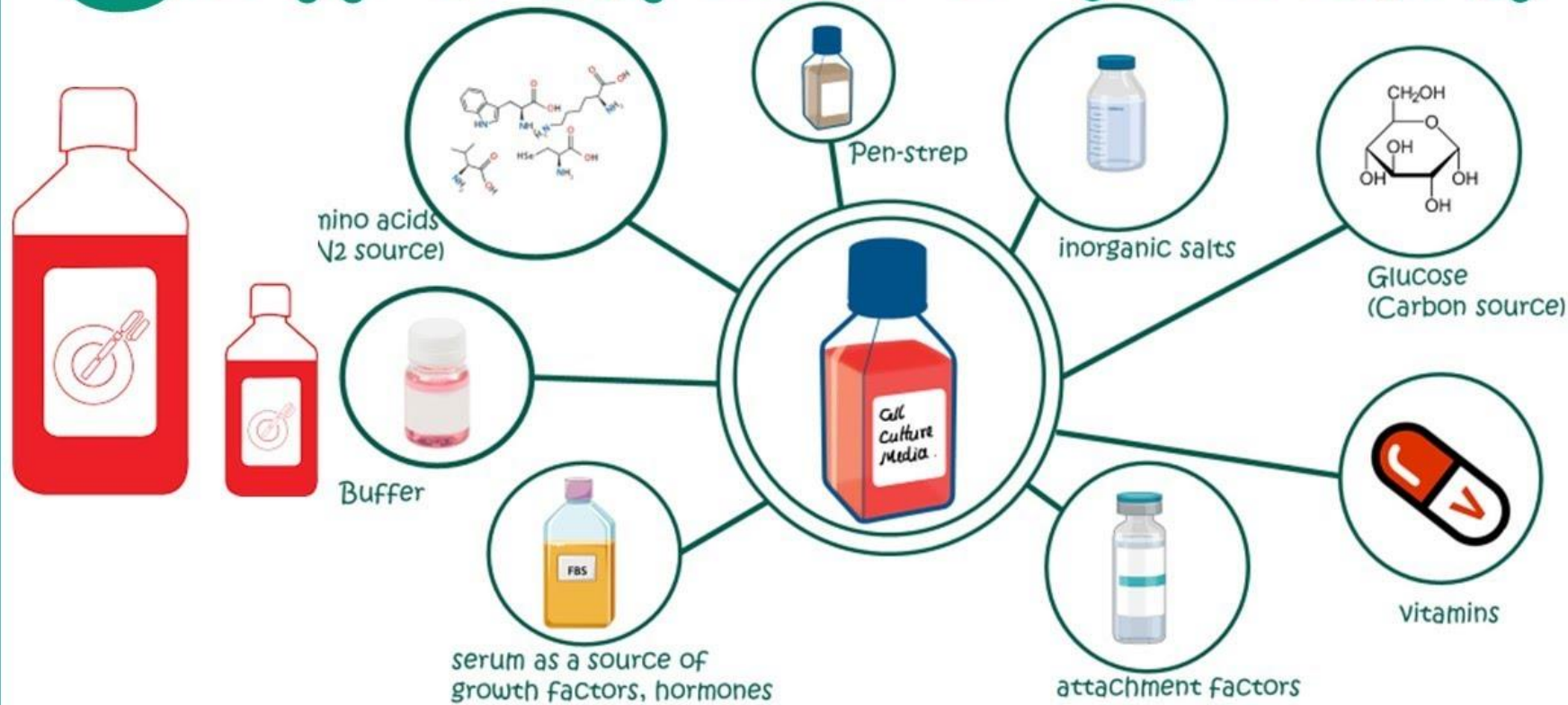








# Cell culture media

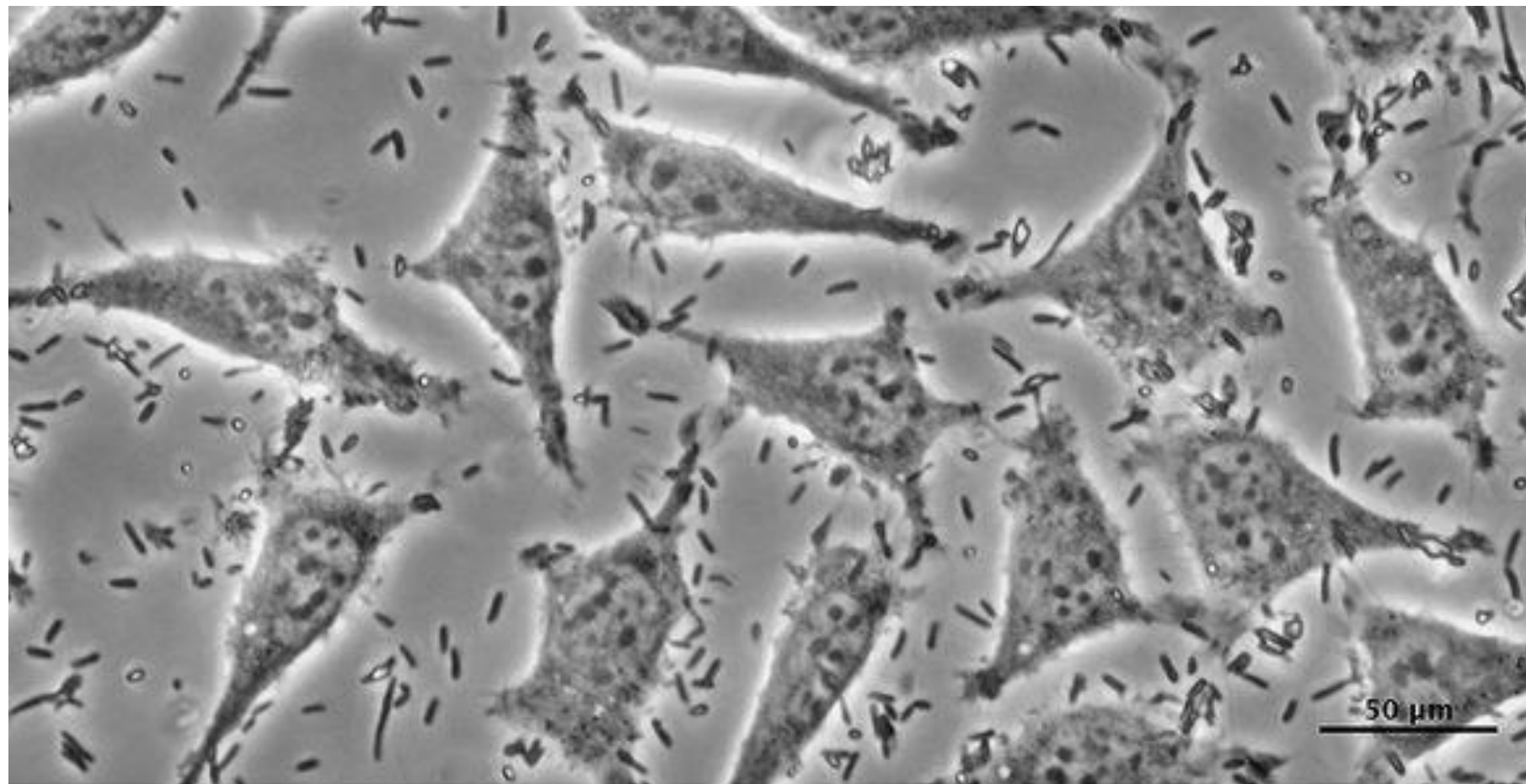






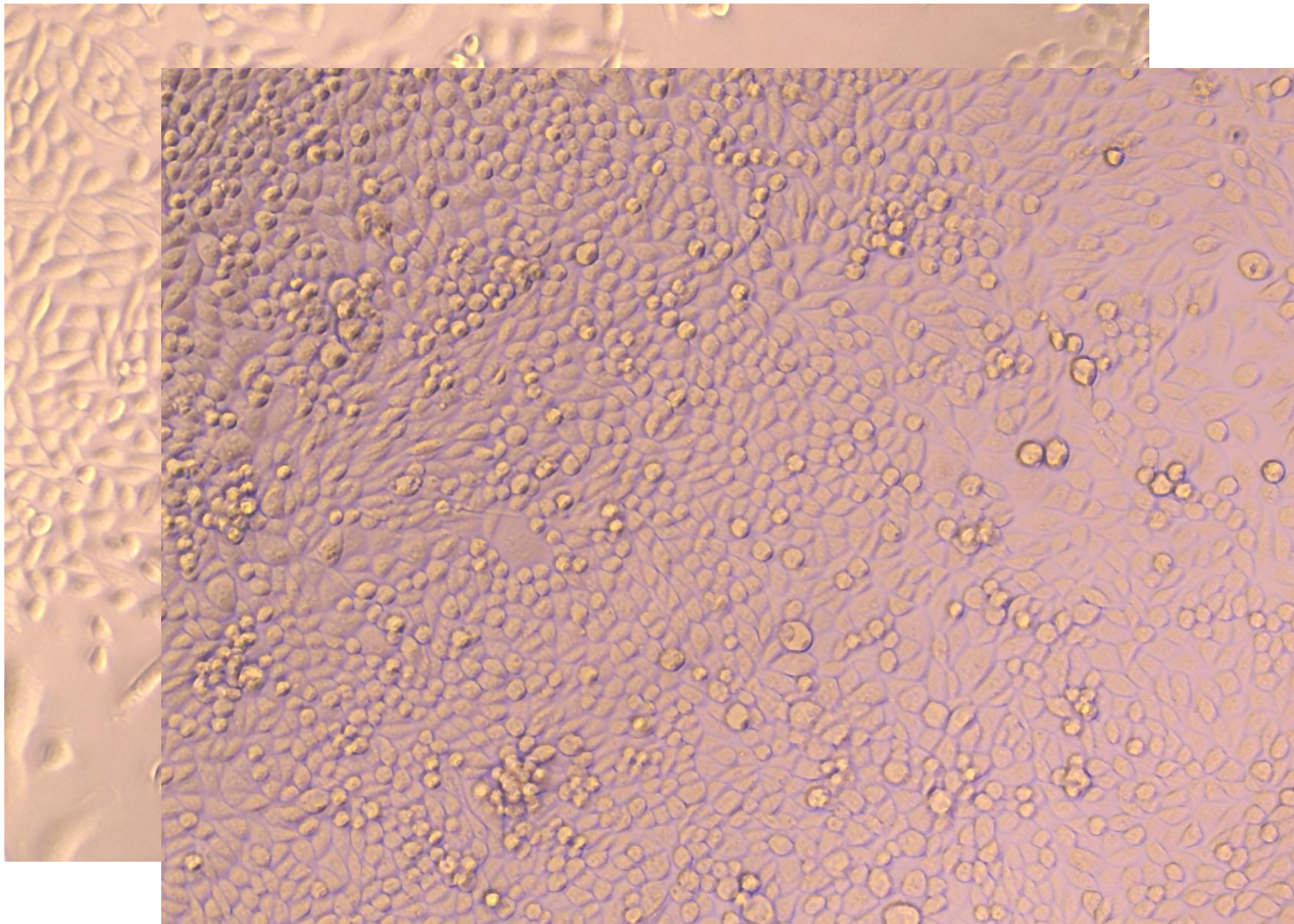






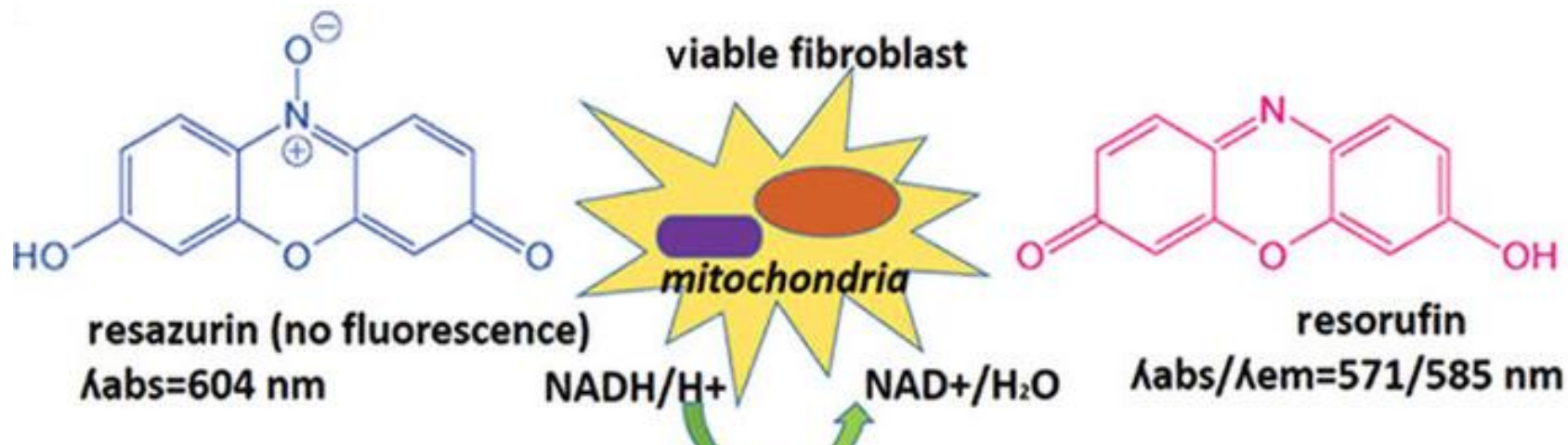








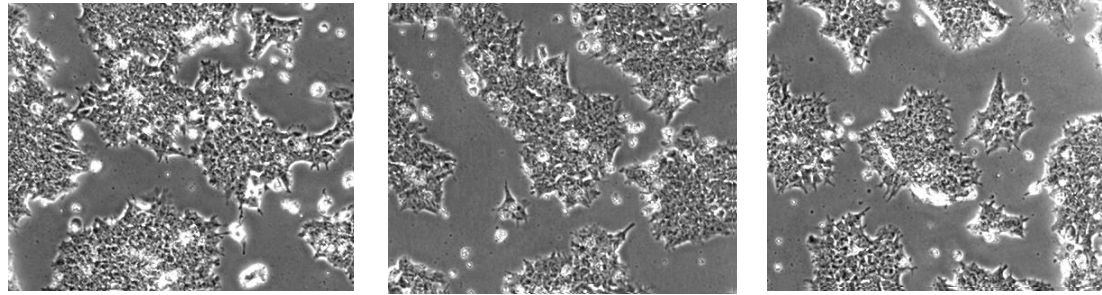
# Resazurin reduction assay



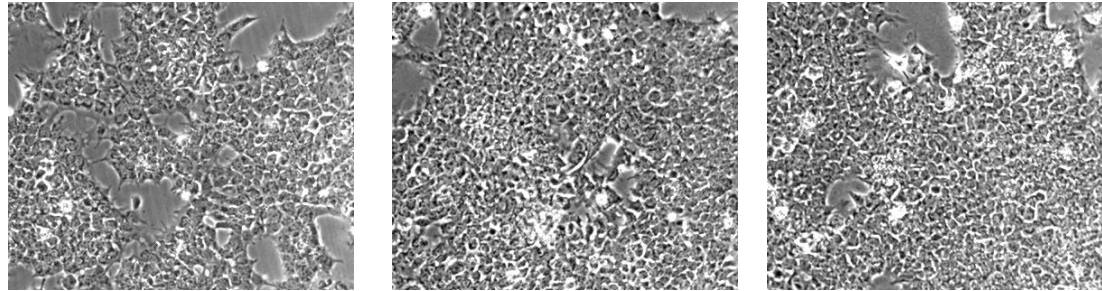


# Застосування

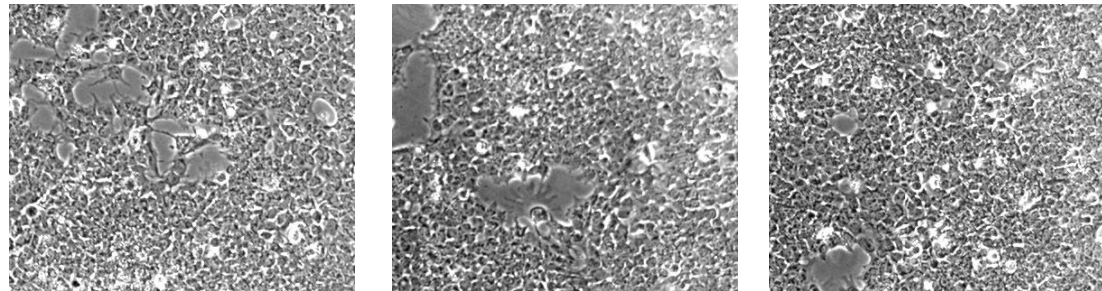
- Доклінічні дослідження нових лікарських препаратів
- Накопичення вірусів
- Дослідження біохімії живих систем
- Виробництво рекомбінантних білків
- Виробництво антитіл
- Дослідження покриттів імплантатів на біосумісність
- Токсикологічні дослідження



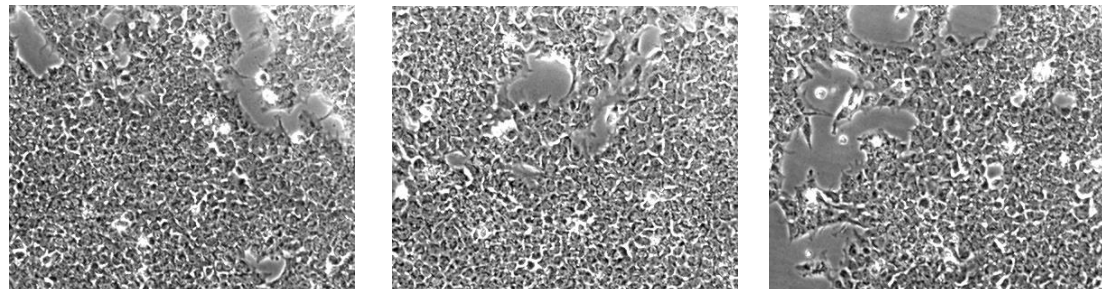
No FGF2



FGF2-133 cleaved



FGF2-133 fused

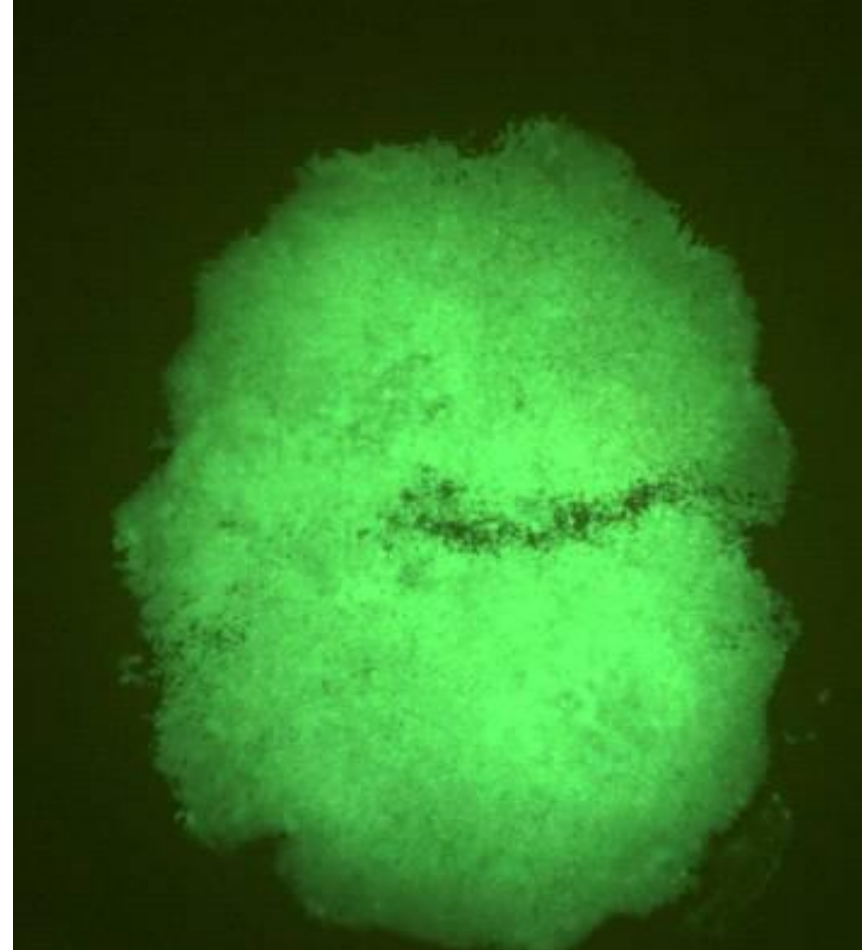
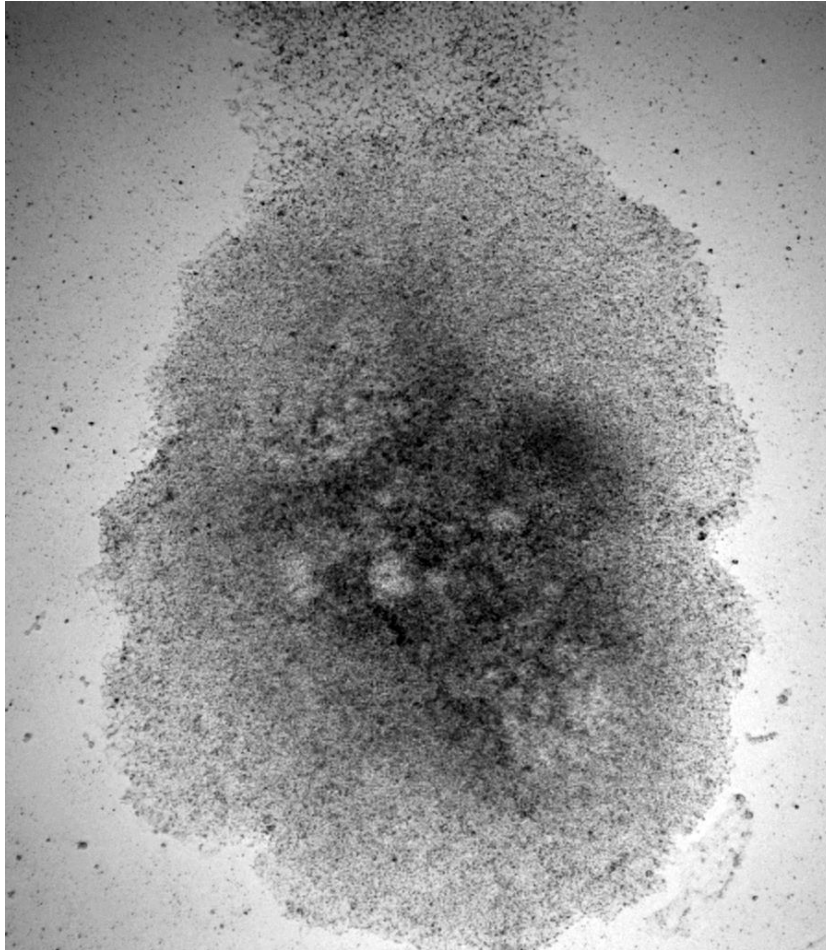


FGF2 standard

Cells plated at density of 20 000/cm<sup>2</sup>, grown for 3 days  
3 representative pictures shown for each treatment



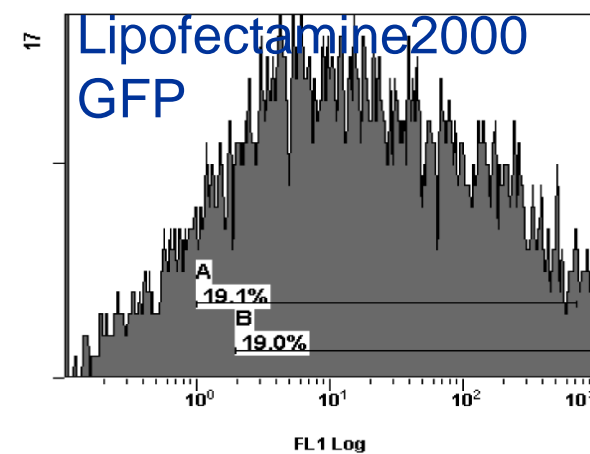
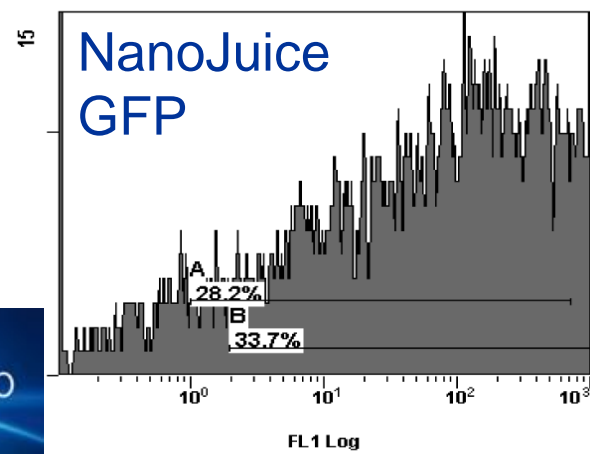
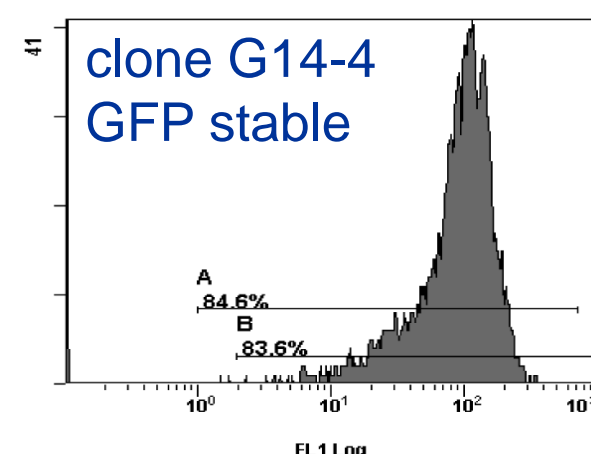
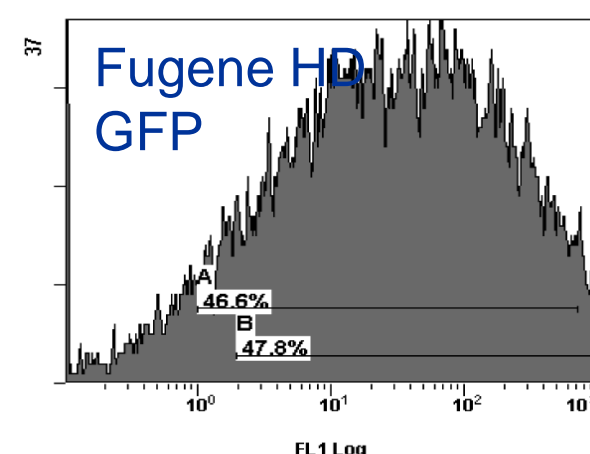
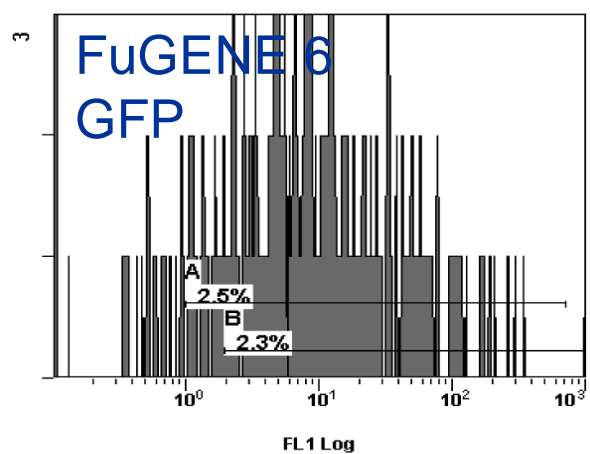
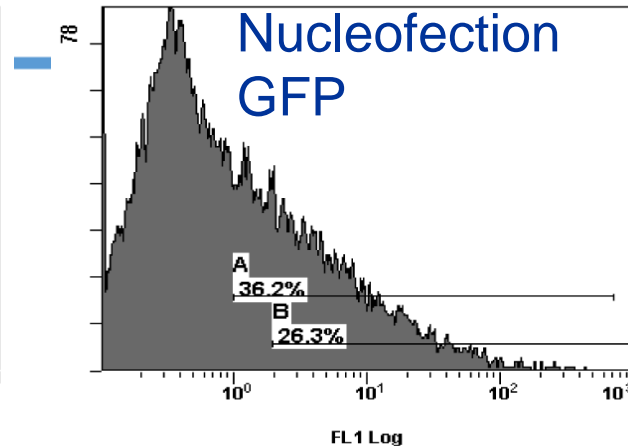
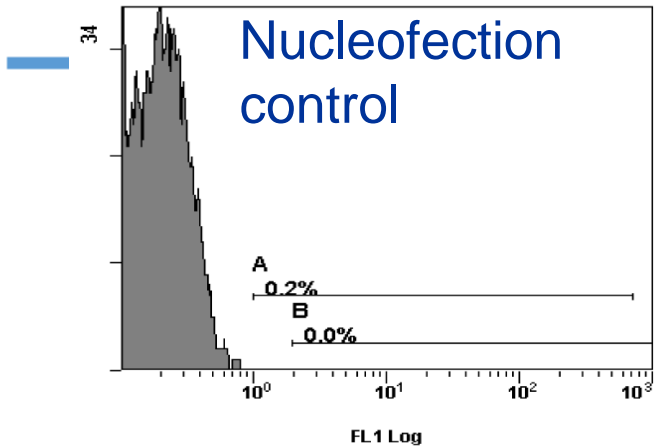
# Stable hESC clone overexpressing GFP



Line CCTL14cp25/6 on Matrigel

Nucleofection – pEGFP-N1, program A-023, solution HSC1

Selection for 15 days with G418 120 ug/ml

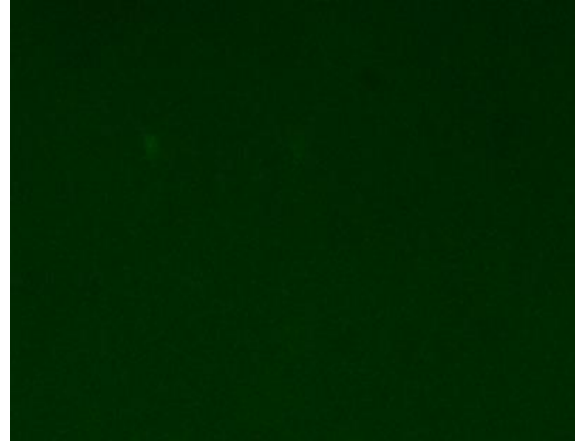
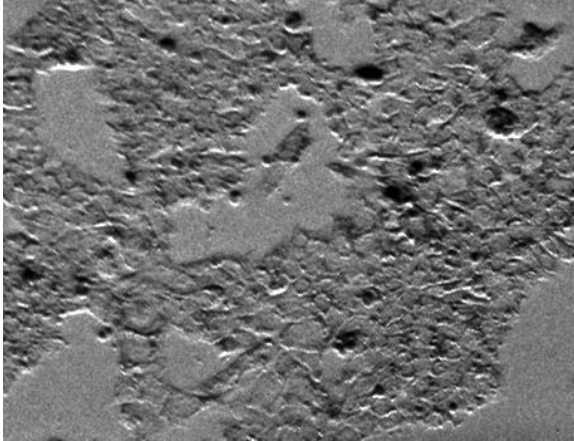


S. Kyrylenko, unpublished

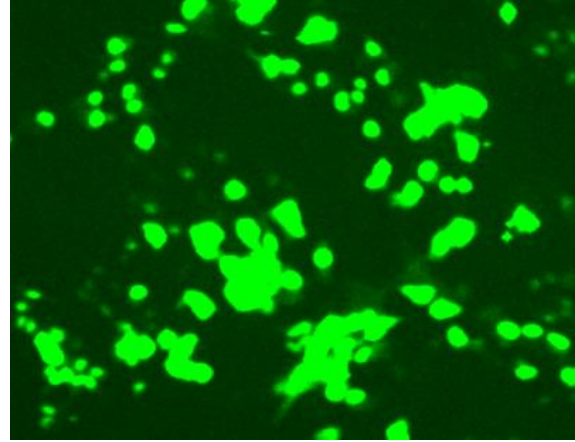
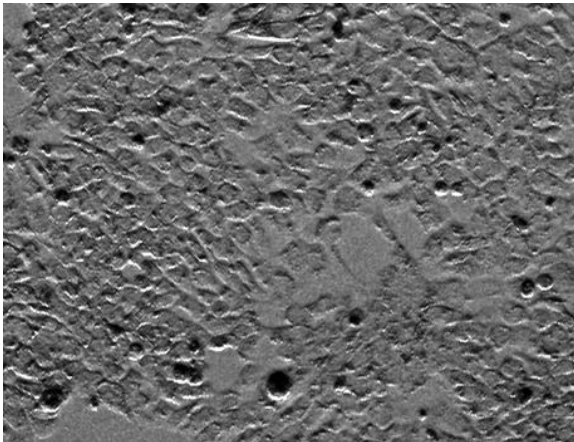


Vis  
↓

GFP  
↓

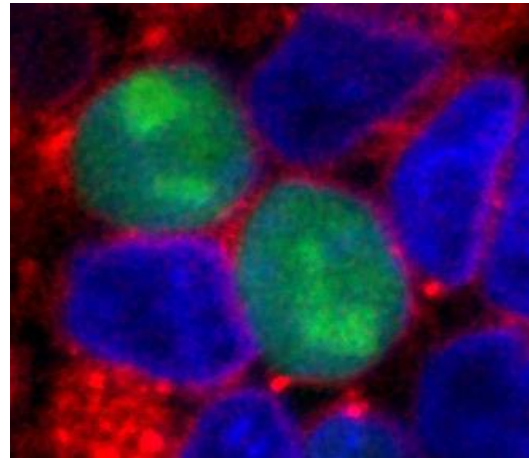


← Non-  
induced

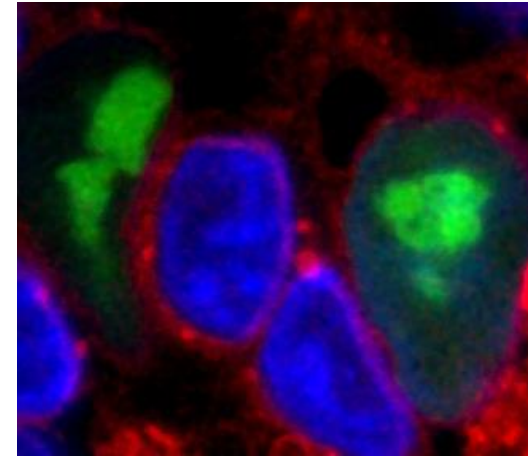


← Induced

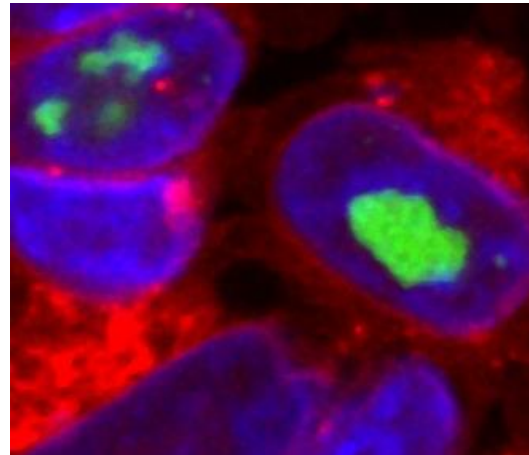
hES cells transfected with pTe105: pTRE-FGF2-GFP(22,3kDa)



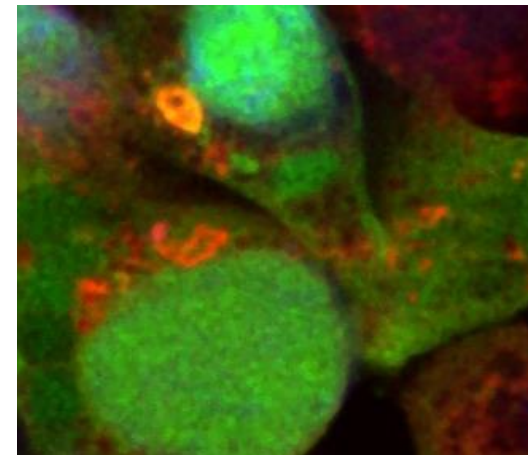
FGF2-GFP  
18 kDa



FGF2-GFP  
22,3 kDa

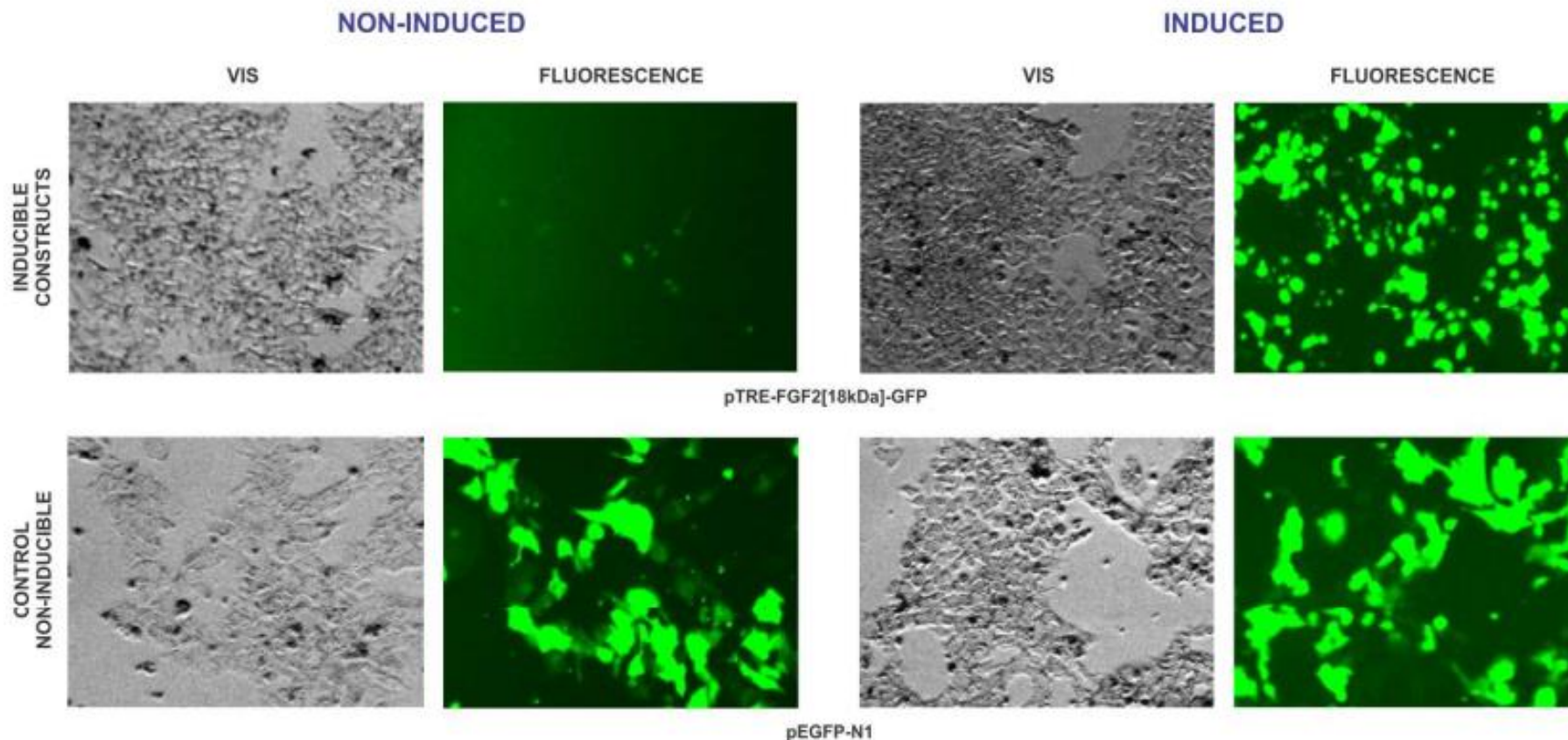


FGF2-GFP  
31 kDa



GFP only  
control





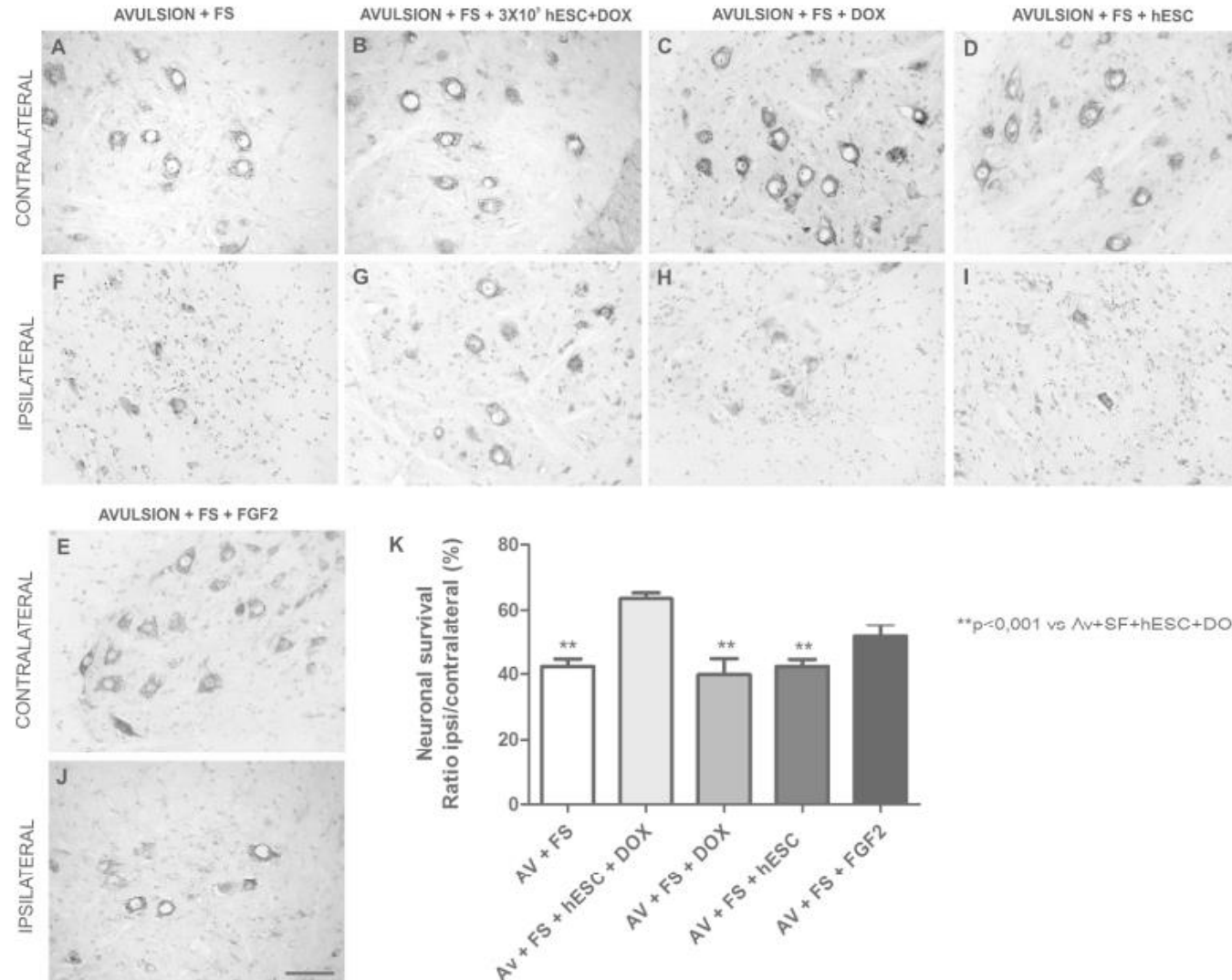
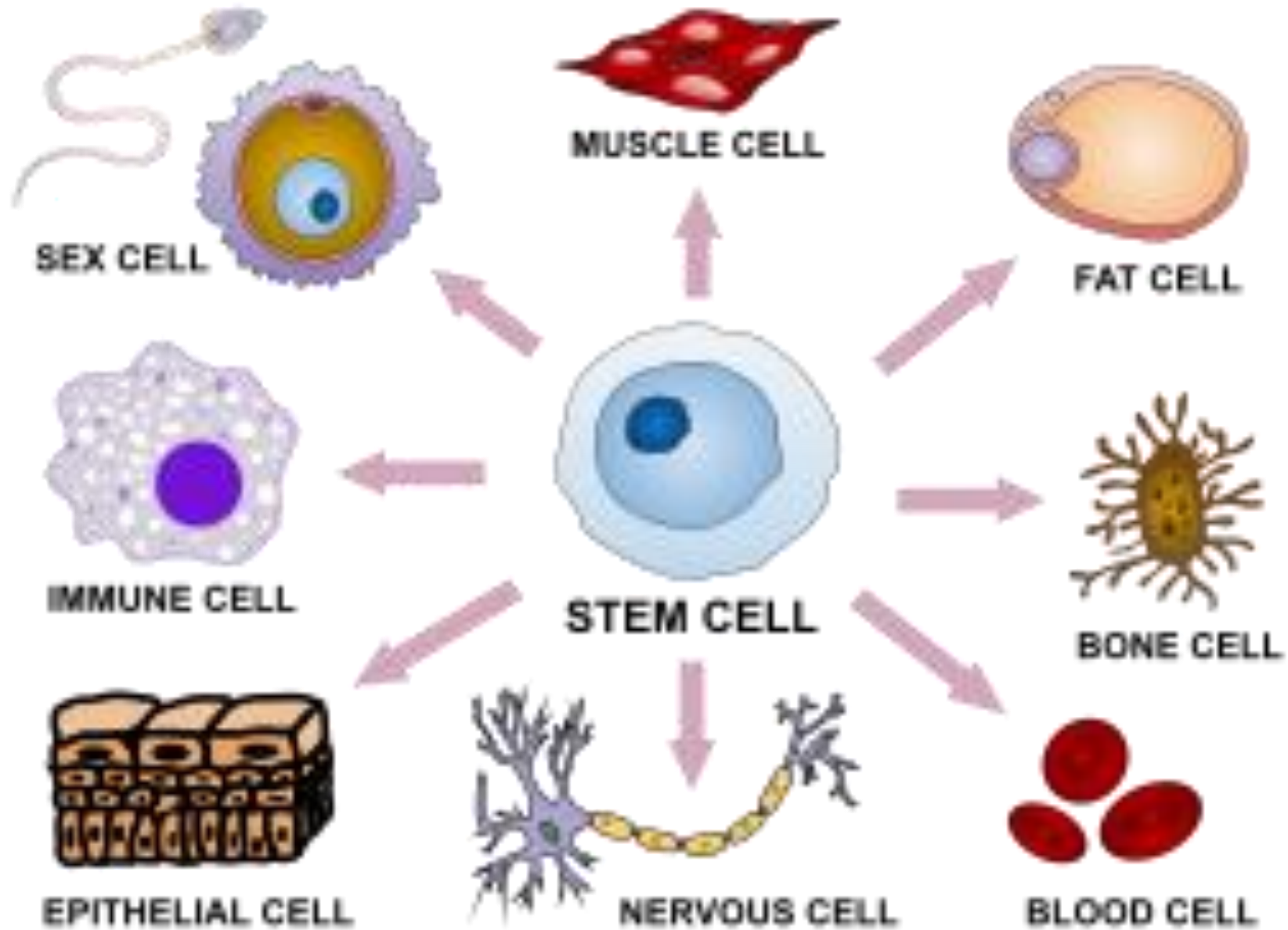
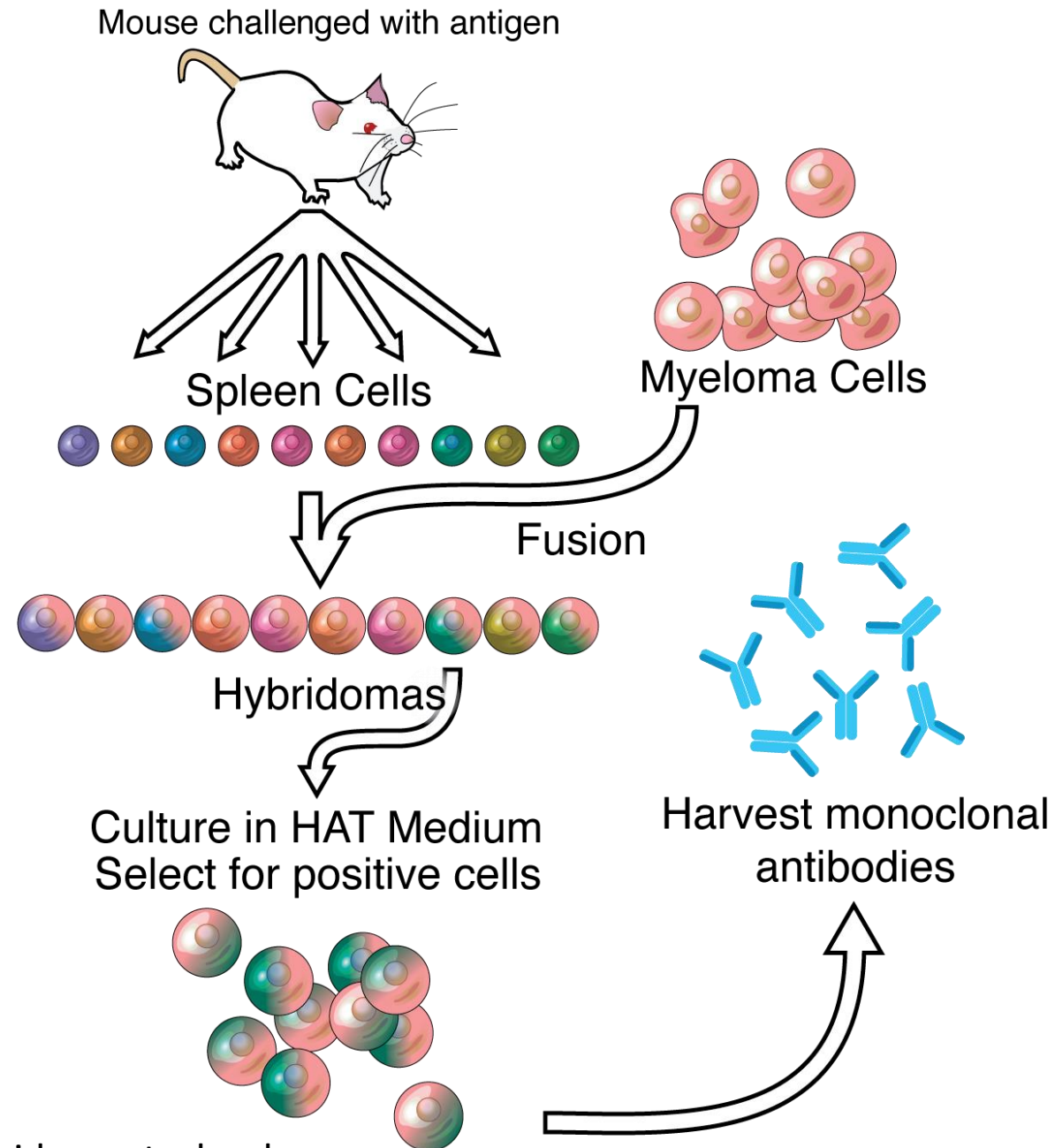


Figure 3. Motoneuron survival after VRA. (A-J) Nissl staining in cross sections of the spinal cord at lamina IX in five different experimental groups, 2 weeks following ventral root avulsion. Note the decreased number of motoneurons, ipsilateral to the lesion. Also, observe the significant improvement of neuronal survival in the group treated with hESC+DOX. (K) Average of the number of motoneurons present in the spinal cord, 2 weeks after avulsion (\*\*  $p < 0.001$  -  $n = 5$ ). Scale bar = 50  $\mu$ m. Mean  $\pm$  SE.









# Cultured meat





**Wish you  
very successful  
cell culture experiments!**



Jean Monnet  
Programme





KIITOS  
DĚKUJI  
ДЯКУЮ  
СПАСИБО  
THANK YOU  
DANKE SCHÖN  
MUITO OBRIGADO