

10th June, 2018:

→ Checked on the cells.

↳ One plate is about 70% confluent, while the other is closer to ~~80~~ 85% confluent.

→ Both can be split tomorrow.

11th June, 2018:

→ P8 cells have been split-managed to save cells from both plates today!

↳ ~~Usual~~ Usual trypsinization, spin down, ^{and} resuspension in 1 mL

- 300 μ L in ~~5~~ 5 mL of media - 2.5 mL in each 100 mm dish

- Remaining 700 μ L spun down again, resuspended in 700 μ L media + 200 μ L FBS + 100 μ L DMSO

⇒ 2 HEK293 P9 plates (100 mm) & 1 P8 stock

→ Aliquots prepared:

↳ ~~5 mL~~ compound media - 45 mL DMEM + 5 mL FBS + 1 mL β /s - 1 tube

↳ Trypsin-EDTA (0.25%) - ^{~12} ~~5~~ mL - 1 tube.

→ For tomorrow -

↳ Check on cells.