## 05. (May) 2019

Project: iGEM\_Munich2019 Shared Project

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## **Exosome Precipitation**

- 1. Centrifugation: 3,500 x g for 20 min, remove as much supernatant as possible
- 2. transfer supernatant to new eppi
- 3. repeat centrifugation
- 4. prepare PEG solutions (stock 20 % PEG (2.0 g) and 150 mM NaCl (0.08766 g) stir and heat up to 40 °C
  - a. PEG 8000 + Protamine (final concentration 0.25 mg)
  - b. PEG 35,000 + Protamine (final concentration 0.25 mg)
- 5. Make 200 µl aliquots from the supernatant form step 3 and add 200 µl of the solutions form step 4
- 6. Incubate samples over night at 4°C
- 7. Centrifuge at 2,000 x g for 30 min, RT
- 8. Discard supernatant
- 9. Resuspend in 50 µl filterd PBS
- 10. Store at 4 °C in the fridge up to one week

Exosome precipation from the supernatant of the 4 wells left form the 6 well plate from 27.05.19

file://tmp/tmpPklOgw.html