SL pH Titration Microscopy

Calibration of pH and chloride in Hek cells

Materials

Ensure the following materials are available before starting the procedure:

- **Effectene**: Transfection kit
- **Plasmid DNA**: Midi or mini transfection-grade

Procedure

- **Day 1**: Cell Seeding

Seed 25,000 HEK cells per well in a Labtec chamber with 0.5 mL of culture medium.

- **Day 2**: Transfection (6-hour protocol, using Qiagen Effectene)
- Mix 0.3 µg of DNA with 50 µL EC buffer in a clean microcentrifuge tube.
- Add 2.4 µL of Enhancer, vortex gently, and incubate for 5 minutes at room temperature.
- Add 4 µL of Effectene, mix thoroughly, and incubate for 10 minutes at room temperature.
- Add the transfection mixture to the cells dropwise and incubate for 6 hours.

	1 well	**… wells**
Plasmid DNA	0.3 µg	
Enhancer	2.4 µL	
EC buffer	50 µL	
Effectene	4 µL	

- **Days 3 & 4**: Chloride and pH Calibration
- Remove medium from one well carefully.
- Add 1 mL of the desired buffer containing ionophores (prepared fresh).
- Incubate the cells for 2 minutes, then change the buffer. Repeat this process 4 times.