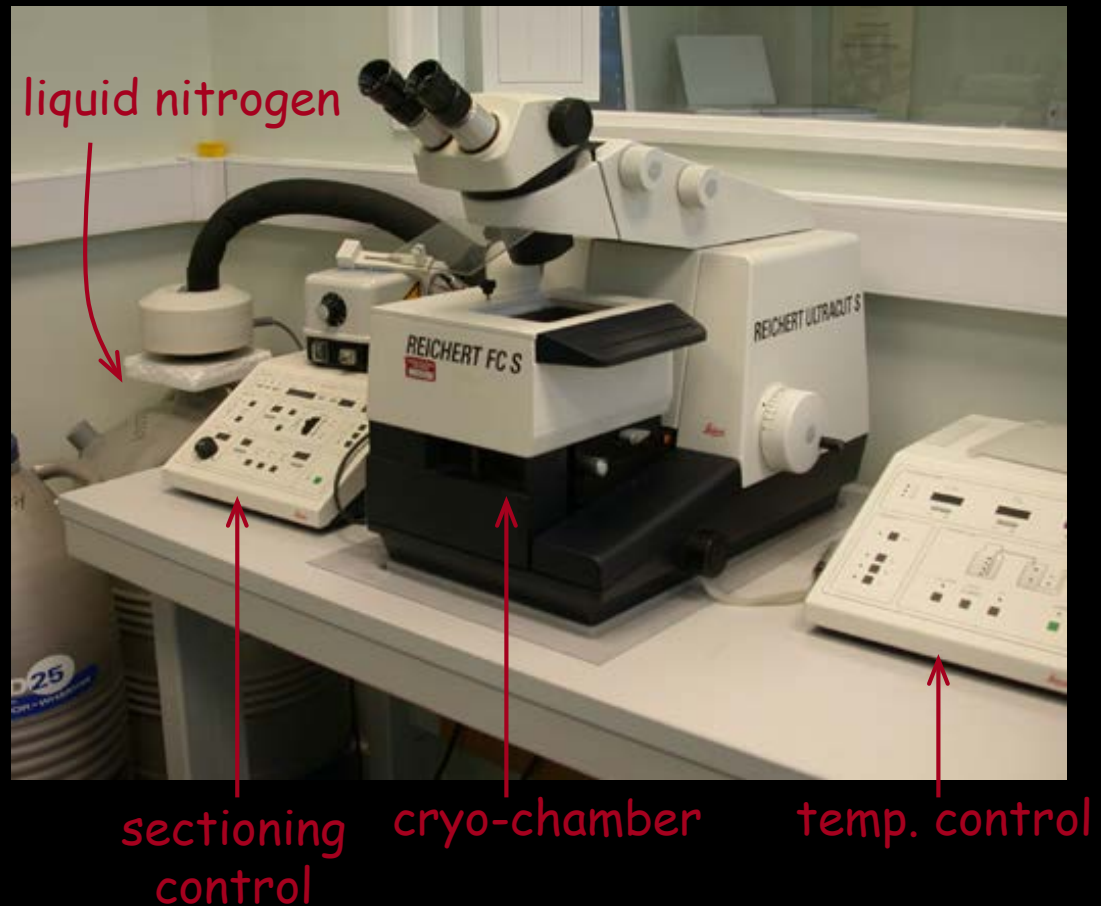


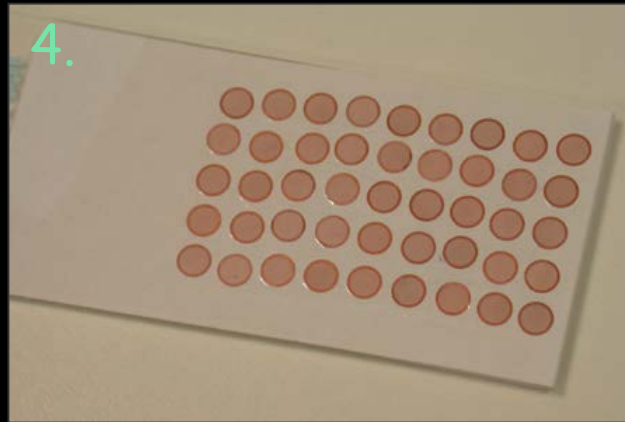
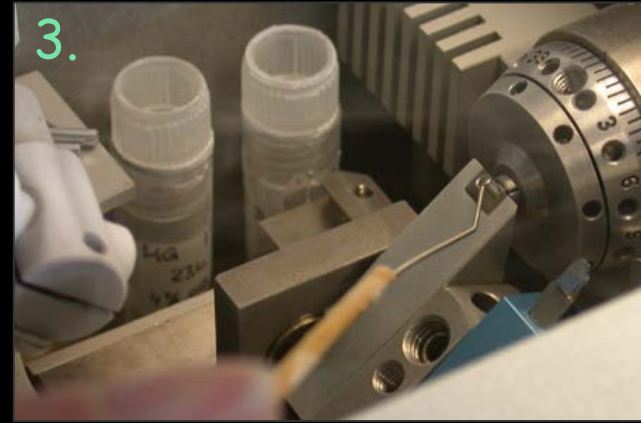
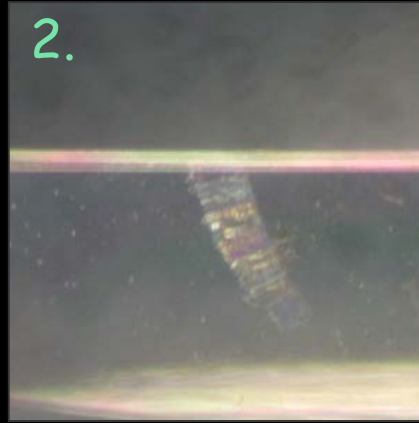
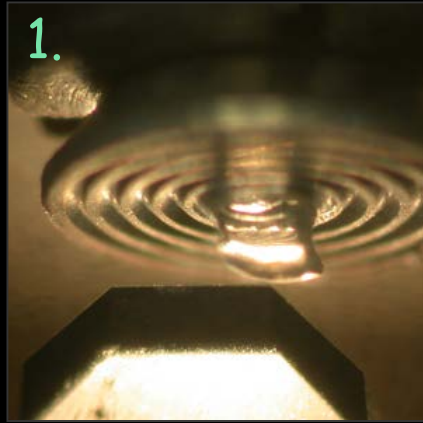
# Immuno-labelling for TEM

## Sample processing

1. Fix 4% PFA
2. Wash/Quench
3. Harvest cells/tissue
- 3a. *Gelatin embedding*
4. 2.3M sucrose infusion
5. Cut  $\leq 1\text{mm}^3$  pieces
6. Mount on cryo-stub
7. Snap freeze liquid N<sub>2</sub>



# Immuno-labelling for TEM

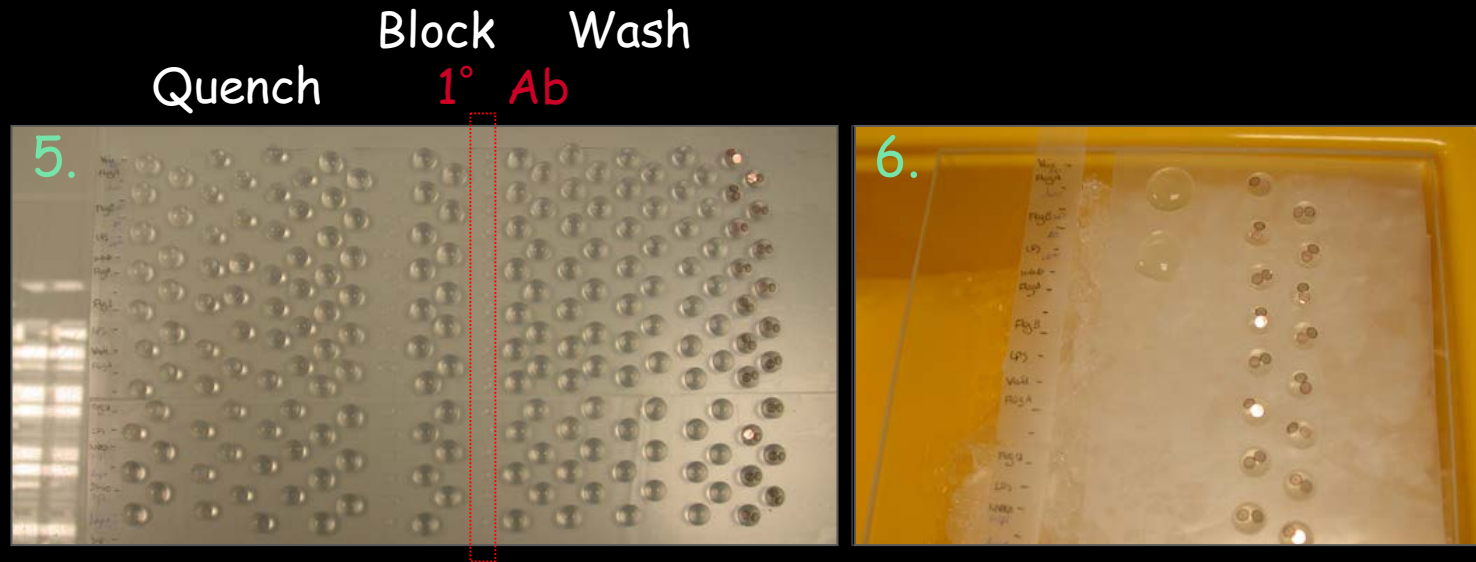


1. Trim
2. Section
3. Pick-up
4. Add to EM grids

5. Immuno-labelling
6. MC/UA contrasting

7. Grid looping
8. Drain excess
9. Dry

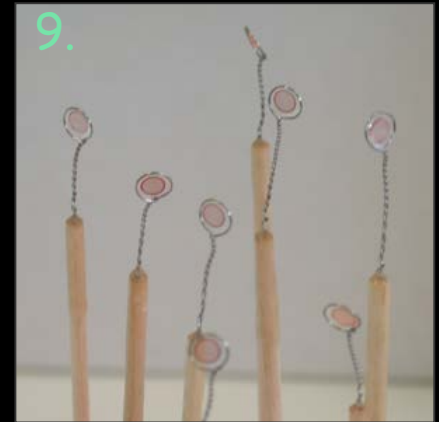
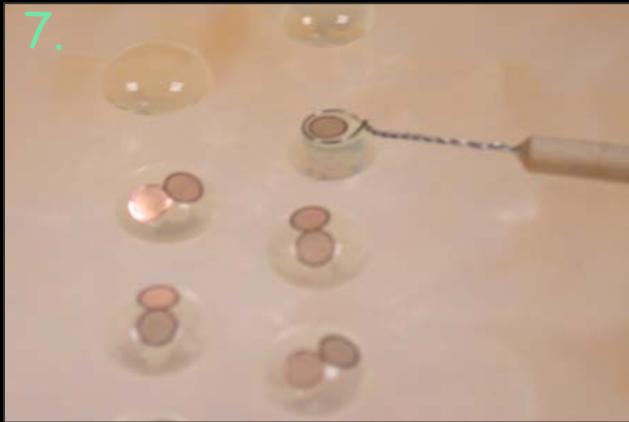
# Immuno-labelling for TEM



50 $\mu$ l washes  
5 $\mu$ l/grid Ab incubations

1. Trim
2. Section
3. Pick-up
4. Add to EM grids
5. Immuno-labelling
6. MC/UA contrasting
7. Grid looping
8. Drain excess
9. Dry

# Immuno-labelling for TEM



1. Trim
2. Section
3. Pick-up
4. Add to EM grids

5. Immuno-labelling
6. MC/UA contrasting

7. Grid looping
8. Drain excess
9. Dry