

# High pressure freezing and Freeze Substitution of Nematodes

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**High Pressure Freezing: EMPact (Leica) followed by AFS: Automatic Freeze Substitution (Leica)**

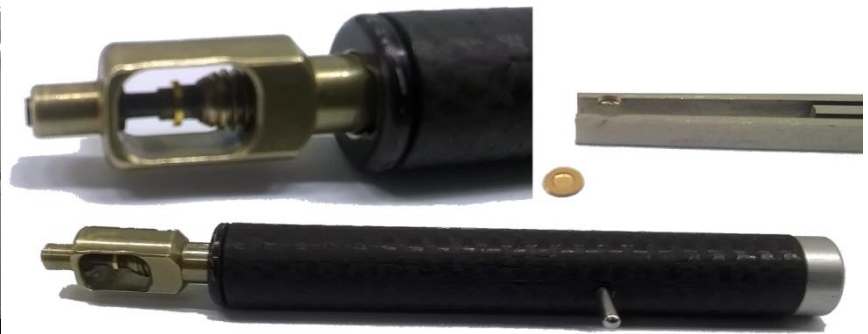
## **EMPact (identical for immunology and morphology)**

20% BSA as cryoprotectant, membrane carrier, egg lecithin(20µg/ml)

- Prepare carrier with egg-lecithin
- Fill carrier with BSA 20%
- Bring nematodes to carrier
- EMPact



EMPact Leica



EMPact carrier

## **AFS: Automatic Freeze Substitution (Leica)**

### **A. AFS cocktail for morphology:**

**1% OsO<sub>4</sub> in 10 ml dried acetone+1% H<sub>2</sub>O+0,5% GA**

9400 µl dry acetone

0,1g OsO<sub>4</sub> (crystal)

100 µl H<sub>2</sub>O

500 µl glutaraldehyde (10 %solution in acetone)

AFS: -90°C

27h

2°C/h (9h)  
 -60°C 12h  
 2°C/h (15h)  
 -30°C 32h  
 2°C/h (17h)  
 4°C

Rinse fixation cocktail with dry acetone. Impregnate with Spurr > imbedding

Polymerase 8h 70°C

### ***B. AFS: cocktail for immuno***

**10ml dried acetone+0,1% GA (10% GA in acetone)+2% water**

9700 µl dry acetone

100 µl glutaraldehyde (10 % solution in acetone)

200 µl H<sub>2</sub>O

AFS: -90°C 27h  
 2°C/h (15h)  
 -60°C 12h  
 2°C/h (15h)  
 -30°C 32h  
 2°C/h (17h)  
 4°C

Rinse fixation cocktail and impregnate with LRwhite.

Polymerisation in AFS with UV

0°C 24h  
 2°C/h 10h  
 20°C 24h  
 2°C/h 8,5h



37°C

72h

*Leica EM-AFS*

### ***Immunolocalization***

Example: localisation of Major Sperm Protein (MSP)

1. 5' PBS
  2. 30min BS (bovine serum 5%; Aurion) Blocking solution
  3. 5x5' washing in IB (incubation buffer):10ml PBS + 100µl BSA-c (10% Aurion)+2 drops HCl (pH 7!)
  4. 60 min Ab prim in IB
  5. 5x5' washing in IB
  6. 30 min Ab bridge. (RAMs; 1:100) DAKO
  7. Washing 5x5' IB
  8. 30 min PAG (protein A 10nmgold)
  9. Washing 2x5' IB
  10. 3x5' PBS
- rinse in bidi distilled water (3x)