

## Maize Kernels – Fixation in FAA, Embedding, Sectioning and Feulgen Staining

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**[Abstract]** The protocol describes preparation of young developing maize kernels for microscopical analysis of nuclei in tissue sections. The fixative FAA (formaldehyde, acetic acid and ethanol) is suitable for preservation of nuclear morphology and allows for quantitative staining of DNA with Schiff reagent in Feulgen staining. The fixation and embedding protocol may be used also for various other histology staining procedures, but care must be taken as the cytoplasm usually shrinks a bit using this procedure. The protocol was used for analysis of seed development of various maize lines, mutants and maize relatives (Vilhar *et al.*, 2002; Kladnik *et al.*, 2006; Dermastia *et al.*, 2009; Bernardi *et al.*, 2012).

### Materials and Reagents

#### 1. Maize Kernels

At different developmental stages; kernels up to 20 days after pollination (DAP) usually fix and section well, at later stages the endosperm becomes progressively dry and hard, causing the sections to crumble and tear while sectioning on the microtome

2. Paraplast Plus (Pelco, catalog number: 18393; Sherwood Medical, catalog number: 8889-502005; or Sigma-Aldrich, catalog number: P3683)
3. Glacial acetic acid (Sigma-Aldrich, catalog number: A6283)
4. Formaldehyde solution (histological grade, 37 wt% in H<sub>2</sub>O) (Sigma-Aldrich, catalog number: 533998)
5. Ethanol (96% and absolute)
6. Tert-butanol (TBA) (Sigma-Aldrich, catalog number: B85927)
7. Pararosaniline hydrochloride (Sigma-Aldrich, catalog number: P3750)
8. Potassium metabisulfite (K<sub>2</sub>S<sub>2</sub>O<sub>5</sub>) (Sigma-Aldrich, catalog number: 31268)
9. Xylene (Xylenes, histological grade) (Sigma-Aldrich, catalog number: 534056)
10. 5 M HCl (Merck-Millipore, catalog number: 1099110001)
11. Decolorizing charcoal (BDH, GB) (Sigma-Aldrich, catalog number: 161551)
12. DPX mounting medium (Sigma-Aldrich, catalog number: 06522)
13. Formaldehyde – acetic acid – ethanol (FAA) (see Recipes)
14. Dehydrating solutions (see Recipes)

15. Schiff reagent (see Recipes)

### **Equipment**

1. Glass vials (volume ~20 ml)
2. Glass fiber filters (Microfibre filters, Whatman GF/C)
3. Scalpels, razor blades
4. Metal forceps (use dedicated forceps for wax work, they are difficult to clean)
5. Disposable plastic Pasteur pipettes
6. Vacuum desiccator (Bel-Art, model: 420100000)
7. Staining dishes (for melting Paraplast Plus) (Electron Microscopy Sciences, catalog number: 70312-21)
8. Hot plate (set on low – around 68 °C), with aluminum foil tent to keep the heat inside
9. Alcohol burner
10. Plastic or metal molds
  - Disposable Base Mold (Pelco, catalog number: 27147)
  - Tissue-Tek Base Mold (Sakura, catalog number: 4123)
11. Tissue embedding cassette bases (Pelco, catalog number: 27168-1) or embedding rings (Pelco, catalog number: 27169-1) or Tissue-Tek Embedding Rings (Sakura, catalog number: 4151)
12. Small beaker with molten Paraplast Plus (dedicated for wax work)
13. Tray with a thin layer of cold water (cool it by adding ice)
14. Ice bucket filled with ice
15. Flat ice packet (frozen cooling pack) or brass plate (optional)
16. Rotary microtome
17. Disposable microtome blades
18. Small brush
19. Forceps with sharp ends
20. Single-edged razorblade
21. Dark cardboard plate
22. Stereomicroscope
23. Coplin jars (Electron Microscopy Sciences, catalog number: 70316-02)

### **Procedure**

#### A. Fixation in FAA

1. Aliquot FAA fixative in glass vials (~20 ml vials, 10 ml fixative). Keep vials on ice, fixative

should be cold when you put tissue in. Cut tissue with a scalpel or razor blade into pieces, that have at least one dimension smaller than 2 or 3 mm, and immediately immerse in cold fixative.

*Note: When cutting the tissue take into consideration the type of sections you wish to obtain.*

2. Place open vials with fixed material in the vacuum desiccator and expose them to moderate vacuum for 15 min to pull the air out of the tissue. If the tissue doesn't sink when you release vacuum, apply and release the vacuum one more time. After vacuuming, replace fixative in vials with fresh fixative.
3. Fix overnight at 4 °C (refrigerator).

#### B. Dehydration in a series of TBA

1. Keep pure TBA (1 L) in a warm place before use (e.g. on the top of an incubation oven), it freezes below 25 °C. Prepare dehydration solutions and store solutions a – e on room temperature (RT) and f – h on the incubating oven.
2. Dehydrate tissue in each step for an hour to one day, depending on tissue size (6 DAP maize kernels for half a day, 12 DAP maize kernels for one day each step). Steps with dehydration solution a – e should be on RT, steps f – h in the incubation oven at 56-60 °C. Use a plastic Pasteur pipette to pull the old solution from the vial, then replace with equal amount of next solution (you need a relatively much larger volume of dehydrating solution compared to your sample). Mark dehydration progress on the label on vial. Alternatively, you can pour the old solution out of the vial and replace with a new one (you don't need to completely drain the vial). Work in a fume hood. In the last change of TBA, fill the vial only to the half of the vial volume, so that you will leave enough room for the adding of the Paraplast Plus in the next step.

#### C. Embedding in paraffin (Paraplast Plus)

1. Keep molten Paraplast Plus and plastic pipettes in the incubation oven at 56-60 °C. You need a large stock of molten paraffin, because it takes a long time to melt (e.g. two rectangular histology staining dishes holding ~300 ml).
2. Add pellets of Paraplast Plus wax in vials half filled with TBA. Alternatively, you can add equal volume of molten wax to the TBA in the vials. Paraplast Plus has a fine structure and additives (DMSO), which help in better penetration into the tissue, which is very important for trouble-free sectioning. Keep in 56-60 °C oven, with the vial cap attached and mix a few times, when the wax is completely melted. Leave in the oven overnight.
3. The next day, pour or pipette away the TBA/wax mixture in a waste container in a fume hood (you can make a small waste container out of aluminum foil). Add fresh molten wax,

and leave in oven overnight with vials open, so that all the TBA evaporates. The next day change the wax with fresh one several times (use a warm transfer pipette or pour away), for example twice a day for two days.

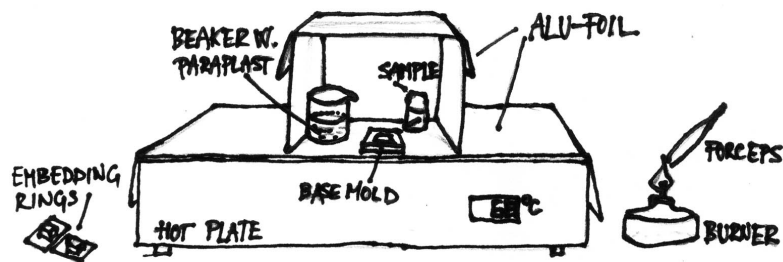
#### D. Casting wax blocks

1. Set air condition in the lab on warmer (if it is too cold, the wax will solidify too quickly).
2. Equipment:
  - a. Hot plate (set on low – around 68 °C), with aluminum foil tent to keep the heat inside
  - b. Metal forceps (use dedicated forceps for wax work, they are difficult to clean)
  - c. Alcohol burner
  - d. Plastic or metal molds
  - e. Embedding rings or embedding cassette bases
  - f. A small beaker with molten Paraplast Plus (also has to be dedicated for wax work)
  - g. A tray with a thin layer of cold water (cool it by adding ice)
  - h. An ice buckets filled with ice
  - i. Flat ice packet (frozen cooling pack) or brass plate (optional)
3. Keep a small amount of molten wax in a beaker in the aluminium foil tent in the hot plate. We also protect the hot plate with a layer of aluminium foil to keep it clean from wax. If the plate is not precisely thermo-regulated check the plate temperature all the time – if it gets too hot to touch, you have to turn it off and then turn it on when the wax starts to solidify. Prolonged temperatures over 62 °C will damage Paraplast Plus. Also, if using metal molds, keep them on a hot plate before using them.
4. Take one vial with samples out of the oven at a time, and place it on the hot plate. Label the same number of embedding rings as the number of samples with a pencil or waterproof marker. Pour some wax in the mold, take one sample out of the vial with forceps warmed in a flame of alcohol burner.

*Note: Be careful, just pass forceps through the flame for a short time, if smoke is coming off the forceps, they are too hot.*

Orient the sample in the mold on the hotplate considering the type of sections you wish to obtain later. Carefully transfer the mold on the brass-plate/ice-packet or just on the table, so that the bottom layer of wax solidifies and fixes the sample in place. If the sample is difficult to position on the hot plate, you can position it now, when the bottom layer of wax is solid. Fill the mold with molten wax to the edge, and cover with the labeled embedding ring. Then transfer the mold into the tray with cold water and ice and leave it there for about 10 min so that wax solidifies completely. Then put the mold on the ice, wait for a few minutes and then remove the wax block from the mold (the cooling with ice helps the block getting loose from the mold). Put the mold back on a hotplate to re-use it. We usually

process 5 to 10 blocks at a time. Store the blocks in the refrigerator at 4 °C (see Figure 1).



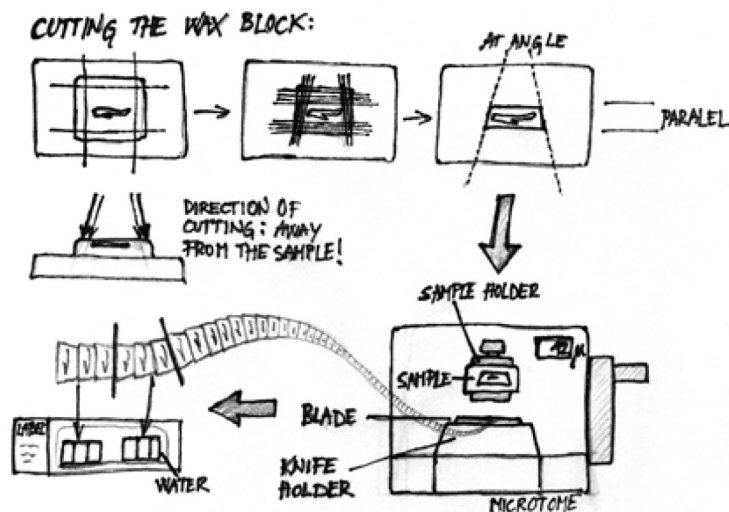
**Figure 1. Schematic drawing of workplace organization for embedding tissue samples in paraffin**

## E. Sectioning

### 1. Equipment:

- a. Rotary microtome
- b. Disposable microtome blades
- c. A small brush
- d. A forceps with sharp ends
- e. A single-edged razorblade
- f. Dark cardboard plate (for putting the sections on and observing them)
- g. A stereomicroscope (optional but recommended)
- h. Hot plate set on 40 °C
- i. A small jar with distilled water
- j. A plastic pipette
- k. Cleaned objective microscope slides, frosted on one edge
- l. A pencil

2. Trim the wax block around the sample with a single-edged razor blade so that the upper and lower edges are parallel, and the left and right edges are at an angle, so that the lower edge of the block is longer than the upper. Also the sides of the block should not be vertical but inclined – always cut the block with the razor blade away from the sample in clean cuts from the top to the bottom of the block (see Figure 2).



**Figure 2. Sectioning paraffin-embedded tissue samples.** Top: Directions of trimming the paraffin around the sample. Bottom: sectioning on the microtome and arranging the sections on a slide.

3. Fix the embedding ring with the sample in the sample holder on the microtome. Set the thickness of sections (usually 10 to 20  $\mu\text{m}$ ) and the cutting angle (around 7 degrees). Start sectioning; the first few sections won't be good, so remove them with a small brush, always away from the blade, not towards it! When the surface of the sample evens, the ribbon will start forming. Cut enough sections to fit them on a dark cardboard, and cut the ribbon with a razor blade. Cut several ribbons, until you are sure that you have cut the region of interest, and then look at the sections with a stereomicroscope, to locate the interesting sections (e.g. median longitudinal sections of the kernel).
4. Label the objective slides with a pencil, put them on a hot plate (at 40  $^{\circ}\text{C}$ ) and apply some water on them (to almost cover them). Cut the ribbon in smaller regions with two or three sections, and apply two sets of sections on each slide (one will be used for the experiment and the other will serve as a control). The water area should be larger than the size of the ribbon, so the sections have room to spread and even out. Optionally you can drain excess water from the slides using filter paper. Leave the sections on the hot plate overnight to dry and attach to the slides. Store them in slide boxes on room temperature or in the refrigerator at 4  $^{\circ}\text{C}$ .

#### F. Feulgen staining

1. The Feulgen reaction (Feulgen and Rossenbeck, 1924) quantitatively stains DNA. The nuclei become purple, while the rest of the cell is clear. The staining is done in Coplin jars (the volume of staining solution is 40 ml). Sections on objective slides are dewaxed in xylene, rehydrated through an ethanol series to water, hydrolyzed in 5 M HCl for 75 min at

20 °C, stained with Feulgen reagent for 120 min at 20 °C, washed for 45 min in six changes of SO<sub>2</sub>-water, dehydrated in an ethanol series, then mounted in DPX mounting medium.

2. Feulgen staining protocol (adapted from Greilhuber and Ebert, 1994)

Rehydration	100% xylene	RT	Date of solution preparation: Feulgen: HCl: SO <sub>2</sub> water: Xylene: Ethanol:
	100% xylene	5 min/step	
	absolute ethanol		
	96% ethanol		
	70% ethanol		
	30% ethanol		
	distilled water		
Hydrolysis	5 M HCl (in a water bath)	20 °C ____ min	<i>75 min for FAA fixed Paraplast embedded samples</i>
	distilled water (cold water stops hydrolysis)	4 °C 5 min	
Feulgen staining	Feulgen (water bath)	20 °C 120 min	
Washing	SO <sub>2</sub> water (work in fume hood!)	RT	<u>250 ml SO<sub>2</sub> water:</u> 247.5 ml dH <sub>2</sub> O 2.5 ml 5 M HCl 1.25 g K <sub>2</sub> S <sub>2</sub> O <sub>2</sub>
		3 x 2 min	
		2 x 10 min 20 min	
Dehydration	distilled water	RT	
	30% ethanol	5 min/step	
	70% ethanol		
	96% ethanol		
	absolute ethanol		
	100% xylene		
	100% xylene		
Mounting	DPX + cover slip		

## Recipes

1. FAA (200 ml)
  - 100 ml 95% ethanol
  - 70 ml dH<sub>2</sub>O
  - 20 ml 37% formaldehyde solution
  - 10 ml glacial acetic acid
  - Store at 4 °C
2. Dehydrating solutions (for 100 ml each)
  - 10 ml TBA, 40 ml 95% ethanol, 50 ml dH<sub>2</sub>O
  - 20 ml TBA, 50 ml 95% ethanol, 30 ml dH<sub>2</sub>O
  - 35 ml TBA, 50 ml 95% ethanol, 15 ml dH<sub>2</sub>O
  - 55 ml TBA, 45 ml 95% ethanol
  - 75 ml TBA, 25 ml 95% ethanol
  - 100 ml TBA
  - 100 ml TBA
  - 100 ml TBA
3. Schiff reagent (Feulgen stain)
 

Boil 800 ml of dH<sub>2</sub>O. Add 4 g of pararosaniline hydrochloride and dissolve while mixing. Cool to 50 °C. Filter the solution through 2 glass fiber filters using vacuum. Add 120 ml 1 M HCl and 12 g K<sub>2</sub>S<sub>2</sub>O<sub>5</sub>. Leave the solution overnight in the dark at room temperature. Add 2 to 4 g decolorizing charcoal in the solution and mix. Filter the solution through 2 glass fiber filters using vacuum into a dry bottle (the stain must be clear and colorless). Store Schiff reagent in the refrigerator at 4 °C for up to 1 year (use until precipitate starts forming in the reagent)

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