植物石蜡切片荧光探针原位杂交（FISH）实验报告

1.实验器材及试剂

1.1.实验器材

名称	厂家	型号
脱水机	DIAPATH	Donatello
载玻片	Wanwu	
盖玻片	江苏世泰实验器材有限公司	10212432C
病理切片机	上海莱卡仪器有限公司	RM2016
无酶离心管	Wanwu	EP-150-M
摇床(钟摆式)	Wanwu	TSY-B
涡旋混匀器	Wanwu	MX-F
移液枪	Dragon	KE0003087/KA0056573
Gene tech pen	Gene tech	GT1001
冰箱	青岛海尔股份有限公司	BCD-192TGN
显微镜	CIC	XSP-C204
正置荧光显微镜	日本尼康	NIKON ECLIPSE CI
成像系统	日本尼康	NIKON DS-U3
恒温箱	LABOTERY	GSP-70
高压灭菌锅	松下健康医疗	MLS-3751L-PC
包埋机	武汉俊杰电子有限公司	JB-P5
冻台	武汉俊杰电子有限公司	JB-L5
组织摊片	浙江省金华市科迪仪器设备有限公司	KD-P

1.2.主要实验试剂

试剂	厂家	货号	稀释比
DEPC	Amresco	E174	
FAA 固定液（DEPC 水）	Wanwu	G1112	
石蜡	Sakura		
无水乙醇	国药集团化学试剂有限公司	100092683	
二甲苯	国药集团化学试剂有限公司	10023418	
PBS 缓冲液（DEPC）	Wanwu	G0020	
20×SSC 洗脱液	Wanwu	G3016-4	
BSA	Wanwu	G5001	
蛋白酶 K	Wanwu	G1205	

抗荧光淬灭封片剂	Wanwu	G1401
杂交缓冲液	Wanwu	G3016-3
DAPI	Wanwu	

2.植物石蜡切片荧光探针原位杂交实验步骤

2.1.组织固定：组织取出洗净后立即放入固定液（DEPC 水配制）中固定 12h 以上。真空泵抽气。

2.2.脱水：组织固定完成后经梯度酒精脱水后浸蜡，包埋。脱水过程真空泵抽气。

2.3.切片：石蜡经切片机切片，摊片机捞片，62°烤箱烤片 2h。

2.4.石蜡切片脱蜡至水：依次将切片放入二甲苯I15min-二甲苯II15min-无水乙醇I5min-无水乙醇II5min,自然晾干，DEPC 水浸泡。

2.5.消化：基因笔画圈，根据不同组织不同指标特性，滴加蛋白酶 K(20ug/ml) 37°消化 min。纯水冲洗后 PBS 洗 3 次×5min。

2.6.预杂交：滴加预杂交液，37°C 孵育 1h。

2.7.杂交：倾去预杂交液，滴加含探针_____杂交液, 浓度_6ng/ul___，恒温箱_37____度杂交过夜。

2.8.杂交后洗涤：洗去杂交液，2×SSC，37°C 洗 10min，1×SSC，37°C 洗 2×5min，0.5×SSC 室温洗 10min。若非特异杂交体较多，可以增加甲酰胺洗涤。

2.9.DAPI 复染核：切片滴加 DAPI 染液，避光孵育 8min，冲洗后滴加抗荧光淬灭封片剂封片。

2.10.镜检拍照：切片于尼康正置荧光显微镜下观察并采集图像。（紫外激发波长 330-380nm，发射波长 420nm,发蓝光；FAM(488)绿光激发波长 465-495nm，发射波长 515-555 nm，发绿光；CY3 红光激发波长 510-560，发射波长 590nm，发红光。

3.植物石蜡切片荧光探针原位杂交实验结果判读

DAPI 染出来的细胞核在紫外的激发下为蓝色，阳性表达为相应荧光素（488）标记的绿光。mRNA 原位杂交显示结果理论为胞浆阳性，少数核阳性属正常。micRNA 与 lncRNA 不同指标表达定位不同。根据表达量不同荧光亮度有强弱。

注：上述涉及到的所有试剂，仪器等在 RNA 原位杂交实验时都需使用 DEPC 处理后的 RNAse free 环境。

附表 1 探针信息

Fluorescence probe-FISH assay protocol (plant paraffin setion)

1. Apparatus and reagents

1.1 Apparatus

Name	Producer	Model
Dehydrator	DIAPATH	Donatello
RNase-free Glass microscope slides	Wanwu	
Coverslip	Citotest	10212432C
Pathologic microtome	Leica	RM2016
Enzyme-free centrifuge tubes	Wanwu	EP-150-M
Shaker	Wanwu	TSY-B
Vortex	Wanwu	MX-F
Pipettor	Dragon	KE0003087/KA0056573
Liquid blocker pen	Gene tech	GT1001
Refrigerator	HAIER	BCD-192TGN
Microscopy	CIC	XSP-C204
Fluorescent Microscopy	NIKON	Nikon Eclipse ci
Imaging system	NIKON	Nikon DS-U3
Incubator	LABOTERY	GSP-70
autoclave	PANASONIC	MLS-3751L-PC
Paraffin embedding machine	WHJJ	JB-P5
Frozen flat	WHJJ	JB-L5
Tissue spreader	Kedee	KD-P

1.2 Major reagents

reagent	manufacturer	article number
Ethanol	SCRC	100092683
Xylene	SCRC	10023418
PBS solution	Wanwu	G0002
Proteinase K	Wanwu	G1205
4% of paraformaldehyde（DEPC water）	Wanwu	G1113
DEPC	Amresco	E174
20×SSC solution	Wanwu	G3016-4

BSA	Wanwu	G5001
Nuclear fast red dye	Wanwu	G1035
hybridization buffer	Wanwu	G3016-3

2. Procedure

2.1. **Organization fixed**: Take out the organization, wash clean, then Immediately put in the fixed fluid (DEPC) to fix above 12h.

2.2. **Dehydration**: The tissue is dehydrated by gradient alcohol, paraffin, embedding and Vacuum pumping in dehydration process.

2.3. **Section**: The paraffin is sliced through the slicer, the piece of the slice machine and the 62 - degree oven roast for 2 hours.

2.4. **Dewaxing and dehydration:** Soak sections in 2 changes of xylene, 15 minutes each. Dehydrate in 2 changes of pure ethanol for 5 minutes each. Then, followed respectively by dehydrating in gradient ethanol of 85% and 75% ethanol 5 minutes each. Wash in DEPC dilution.

2.5. **Digestion**: Mark the objective tissue with liquid blocker pen, according to the characteristics of tissues, add proteinase K(20 ug/ml) working solution to cover objectives and incubate at 37℃ for___min. Washing with pure water, then wash three times with PBS (pH 7.4) in a r0ocker device, 5 min each.

2.6. **Pre-hybridization**: Add Pre-hybridization solution to each section and incubate for 1 h at 37℃.

2.7. **Hybridization**: Discard the pre-hybridization solution, add the probe hybridization solution ,concentration_____, and incubate the section in a humidity chamber and hybridize overnight at___℃.

2.8. **Washing**: Remove the hybridization solution. Wash sections with 2×SSC for 5 min at 37℃, Wash sections in 1×SSC two times for 5 min each at 37℃, and wash in 0.5×SSC for 10 min at room temperature. Formamide washing can be added if there are more non-specific hybrids.

2.9 **Stain cell nuclei (counter stain)**: incubate with DAPI for 8min in the dark, and then mounting.

2.10 **Microscopic examination and photography**: to take photos with positive fluorescence microscope. DAPI glows blue by UV excitation wavelength 330-380 nm and emission wavelength 420 nm; FAM glows green by excitation wavelength 465-495 nm and emission wavelength 515-555 nm; CY3 glows red by excitation wavelength 510-560 nm and emission wavelength 590 nm.

3. Interpretation of the results

The positive expression of BCIP/NBT was blue-purple. The results of mRNA in situ hybridization showed that cytoplasm was positive and a few nuclear positive were normal. The expression of micRNA and lncRNA was different. According to the expression, the color is very

light.

Attached table 1 probe information.