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Protocol status: Working

We use this protocol and it's working

A protocol of molecular detection of phytoplasmas and *Xylella* spp. in post-entry quarantine for plants. V.6

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ABSTRACT

In the STEPS, we describe TaqMan multiplex real-time PCR to universally detect phytoplasmas (PP) and *Xylella* spp. (XL) with plant internal control (IC) from crude extracts. A protocol file [Protocol-JpEnXX.pdf] shows further details of the protocol in Japanese and English.

ATTACHMENTS

[Protocol-JpEn13.pdf](#)

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18061

Keywords: phytoplasma,
xylella, real-time PCR,
conventional PCR, multiplex,
detection, dual priming
oligonucleotides, plant
quarantine

- 1 1. Extraction
 - 1.1. Crude extraction
 - 1.1.1. Put leaf petioles (< 50mg), a metal beads, and 1mL extraction buffer into a tube.
 - 1.1.2. 2,500 rpm 60 sec. (the Multi-beads shocker)
 - 1.1.3. 9,000 x g 10min 4C
 - 1.1.4. Transfer the supernatant to a new tube. Next steps, or keep it in a freezer.
- 2 1.2. Isopropanol precipitation
 - 1.2.1. Add an equal volume of cold isopropanol to the crude extract and mix.
 - 1.2.2. 20,000 x g 5 min 4C
 - 1.2.3. Discard supernatants and dry pellets.
 - 1.2.4. Suspend the pellet in one-fifth volume of TE. Next steps, or keep it in a freezer.

3	2. Real-time PCR		
	2.1. Reagent mixture		
	Reagents	1 reaction	10 reactions
	Sterile water	2.5	25
	TaqMan FAST Advanced Master Mix	5	50
	Primer mixture	1	10
	Probe mixture	1	10
	Total (µL)	9.5	95
	2.2. Dispense 9.5 µL of the reagent mixture to PCR tubes		
	2.3. Add 0.5 µL of the extract (1.2.4) to the tube.		
	2.4. Set the tubes and run the StepOnePlus with the following parameters:		
	50C 2 min. → 95C 20 sec. →		
	95C 1 sec. → 60C 20 sec. (50 cycles)		
	* Targets (Reporter/Quencher) : PP (FAM/NFQ-MGB) ,		
	XL (VIC/NFQ-MGB) , IC (TAMRA/None)		
	See the next step.		

- 4 3. Data analysis
 - 3.1. Export data
 - 3.2. Consider positives of PP/XL at Ct<45 and IC at Ct<40.
 - 3.3. Refer the PDF file (a protocol in Japanese and English) in the Absract for detail.