EUROPEAN AND MEDITERRANEAN PLANT PROTECTION ORGANIZATION ORGANISATION EUROPEENNE ET MEDITERRANEENNE POUR LA PROTECTION DES PLANTES Summary sheet of validation data for a diagnostic test

The EPPO Standard PM 7/98 Specific requirements for laboratories preparing accreditation for a plant pest diagnostic activity describes how validation should be conducted. It also includes definitions of performance criteria.

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Laboratory contact details	Bacteriology. Instituto Valenciano de Investigaciones Agrarias CV-315, km. 10.7, 46113 Moncada, Spain	
Short description of the test	Isolation of Erwinia amylovora from plant material in medium King's B	
Date, reference of the validation report	2012-03-01 - Not specified	
Validation process according to EPPO Standard PM7/98?	yes	
Is the lab accredited for this test?	no	
Was the validated data generated in the framework of a project?		
Description of the test		
Organism(s)	Erwinia amylovora (ERWIAM)	
Detection / identification	detection	
Method(s)	Isolation	
Method: Isolation		
Reference of the test description		
As or adapted from an EPPO diagnostic protocol	yes	
EPPO Diagnostic Protocol name	PM 7/020 Erwinia amylovora (version 1)	
Name of the test	Isolation on King's B medium (King et al., 1954)	
Is the test modified compared to the reference test	no	
Other information		
Are the performance characteristics included in the EPPO diagnostic protocol?	yes	
Performance Criteria :		
Organism 1.:	Erwinia amylovora(ERWIAM)	
Analytical sensitivity		
What is smallest amount of target that can be detected reliably?	10^3 CFU/mL plant extract	
<u>Diagnostic sensitivity</u>		

Proportion of infected/infested samples tested positive compared to results from the standard test, see appendix 2 of PM 7/98	Proportion of true positives/total number of samples: 0.63 (in samples from 1 to 10^6 CFU/mL of plant extract and healthy samples in ring test 2010)	
Standard test(s)	Not specified	
Analytical specificity - inclusivity		
Number of strains/populations of target organisms tested	Contact lab for details	
Specificity value		
Diagnostic Specificity		
Proportion of uninfected/uninfested samples (true negatives) testing negative compared to results from a standard test	Proportion of true negatives/total number of samples:0.98 (in samples from 1 to 10^6 CFU/mL of plant extract and healthy samples in ring test 2010)	
Reproducibility		
Provide the calculated % of agreement for a given level of the pest (see PM 7/98)	100% when tested with different operators in IVIA assays	
Repeatability		
Provide the calculated % of agreement for a given level of the pest (see PM 7/98)	100% in IVIA assays	
Test performance study		
Test performance study?	yes	
Brief details of the test performance study and its output.It available, link to published article/report	14 laboratories from Europe, Morocco, USA and New Zealand) analysed 12 samples each (from 1 to 10^6 CFU/mL plant extract and healthy samples). Details about ring test protocol are available.	

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