

## Information pest:

### *Candidatus phytoplasma solani*

*Candidatus phytoplasma solani*, belonging to the 16SrXII-A ribosomal subgroup, has been found to cause a range of plant diseases in different agro-ecosystems in many countries in Europe and the eastern Mediterranean area and a number of others all over the world. Diseases caused include Bois noir in grapevines, stolbur in tomatoes, potatoes and other wild and cultivated plants, maize redness, lavender decline, and yellowing, reddening, decline, dwarfism, leaf malformation and degeneration diseases of other plants. *Candidatus phytoplasma solani* is usually transmitted from plant to plant by the insect vector *Hyalesthes obsoletus*, but not only. It is also transmitted by parasitic plants and by grafting and vegetative propagation of infected host plants.

This phytoplasma is included in European legislation as a quarantine pest.

Disease symptoms develop mainly in summer. Leaves turn yellow or red depending on the cultivar. They roll down-ward and become brittle; the interveinal area of leaves may become necrotic. Shoots show incomplete lignification.

Bois noir's symptoms are similar to those caused by other yellows diseases of grapevine, in particular Flavescence dorée; molecular diagnostic is the only way to differentiate these two phytoplasmas.

## Introduction

The qPCR BN set is based on an analysis method developed by the company International Plant Analysis and Diagnostics ([www.ipadlab.eu](http://www.ipadlab.eu)). The primers and probes were designed on the *rpl22-rps3* BN gene. The primers and probes sequences and their use in diagnostic tests are the subject of a PCT patent application by IpadLab (PCT/IB2010/053563). This Duplex Real-Time PCR set offers a specific and sensitive method to detect 16SrXII phytoplasmas including BN, as well as an endogenous control designed on the COX gene.

Validation data of the method are available from a test performance study realized in 2013 (Euphresco Grafdepi project - report available from this link: [http://www.euphresco.net/media/project\\_reports/grafdepi\\_final\\_report.pdf](http://www.euphresco.net/media/project_reports/grafdepi_final_report.pdf)). This method is also referred in the Appendix 6 of the PM7/079 (2) Grapevine Flavescence dorée phytoplasma, European and Mediterranean Plant Protection Organization Bulletin (2016) 46 (1), 78-93 available from this link: <https://onlinelibrary.wiley.com/doi/epdf/10.1111/epp.12280>.

*This product should be used only for research purposes.*

## Intended use

The qPCR set is validated for the simultaneous detection of Bois noir and Internal Control (IC) in Duplex Real-Time PCR. Suitable tissues are grapevine leaves, preferably primary veins and petioles. Other parts of grapevine tissues can be analyzed, for example vine stocks, or insects' vectors, using specific nucleic acids extraction's methods.

It is preferable to test plant tissues with vine yellows symptoms, sampled from an early stage of the veraison until the beginning of the senescence (from end of August to end of October in the Europe zone).

## Set format and content

Two sets are available for 24 and 96 tests.

Article N°	Product name	
7BN---q2	qPCR BN set 24	
7BN---q9	qPCR BN set 96	

Content	set 24	set 96
Direct Master Mix	24 tests 7BN---q2-DM-	2x48 tests 7BN---q9-DM-
Positive Control	3 tests 7BN---q2-PC-	8 tests 7BN---q9-PC-
Negative Control	3 tests 7BN---q2-NC-	8 tests 7BN---q9-NC-

## Storage conditions

This set can be shipped at room temperature but upon receipt it should be stored immediately at the recommended storage temperature: **from -30 ° C to -10 ° C**.

Avoid prolonged exposure to light and repeated freeze and thaw cycles.

## Shelf life

If the set is correctly stored, at constant-temperature freezer, its performance is guaranteed until the expiration date indicated on the tubes label.

## Materials and equipment (not provided)

- DNA extraction tools and reagents
- Nuclease-free filter tips and micropipettes
- Optical grade nuclease-free tubes/plate
- Disposable latex or vinyl gloves
- Thermal cycler for Real-Time PCR with filters calibrated for VIC<sup>®</sup> and Cy5<sup>®</sup>

## Nucleic acids extraction

Extract DNA from samples according to your usual protocol. Upon request, Qualiplante can recommend you an extraction method.

## Reaction set-up

- Slowly thaw **Direct Master Mix** by placing it on ice or at 4°C.
- Shake briefly **Direct Master Mix** and spin down the liquid.
- Add 18 µl of **Direct Master Mix** (without DNA template) to each PCR tubes or wells of an optical-grade PCR plate.
- Add 2 µl of DNA template to the **Direct Master Mix**. Do not forget to prepare a PCR tube or well of an optical-grade PCR plate for the **Positive Control** and the **Negative Control**.

Components	Volume/PCR tube or well
DNA template or <b>Positive control</b> or <b>Negative control</b>	2 µl
<b>Direct Master Mix</b>	18 µl
Total Volume / PCR tube or well	20 µl

In order to confirm the absence of any reagent's contamination, we strongly recommend including a no-template control (e.g. DEPC water) in the assay.

## Run and thermal cycling

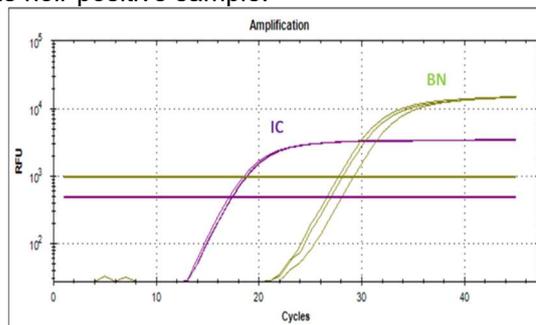
- Seal carefully the PCR tubes or PCR plate. Centrifuge briefly to collect components at the bottom of the PCR tubes or wells of the plate. Protect from light before thermocycling.
- Load the PCR tubes or plate into the thermal-cycler and follow the thermal cycling below:

Steps	Temp (°C)	Time	Cycle(s)
Enzyme activation	95°C	10 min	1
Denaturation	95°C	15 sec	45
Annealing and elongation	60°C	60 sec	

## Results analysis

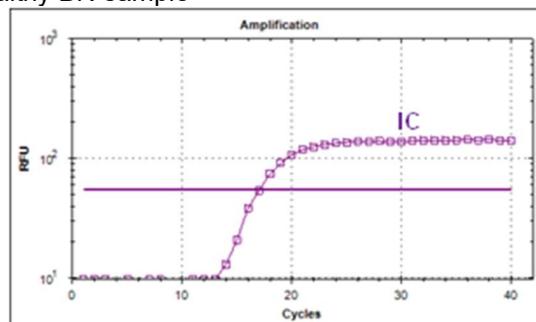
The reaction for Bois noir will generate a specific VIC®-labeled amplification curve.  
 The reaction for Internal Control will generate a specific Cy5®-labeled amplification curve in all the wells except those that contains the no-template control.

**Fig.1:** Example of an amplification curve relative to a Bois noir positive sample.



**fig.1** shows the amplification curves associated to a **BN-infected sample** (green curve) and the relative **Internal Control** (violet curve). This figure shows the **Positive Control** amplification curves of the set.

**Fig 2:** Example of amplification curve relative to a healthy BN sample



**fig. 2** shows the amplification curve associated to a healthy sample and to the **Negative Control** of the set. Only the **Internal Control** curve must appear (violet curve).

## ANALYSIS VALIDATION

The PCR plate is validated only when:

- ✓ the **Positive Control** generates an amplification curve for the two fluorophores VIC® and Cy5® higher than the respective threshold lines (**fig. 1**).
- ✓ the **Negative Control** does not generate any curve associated to the fluorophore VIC®, but it generates an amplification curve associated to the fluorophore Cy5® for the **Internal Control** (**fig. 2**).
- ✓ the amplification plot associated to the **Internal Control** is present in all the wells, except in the no-template control(\*).
- ✓ the no-template control does not generate any curve for the two fluorophores VIC® and Cy5® (\*).

(\* An amplification curve may sometimes appear for the fluorophore Cy5® in the no-template control. In this case, the experiment is validated only if the Ct value is higher than 28.

## RESULTS INTERPRETATION

When all the previous conditions are performed, the amplification results are interpreted as indicated in the **tab. 1**, by considering for each sample the Ct of the curve generated by the VIC® fluorophore specific to BN and the Cy5® fluorophore specific to the IC.

The columns of the **tab. 1** refer to the Ct of the IC amplification and the lines refer to the Ct of BN-specific amplifications.

Fluorophore	Cy5 fluorophore	
	Ct IC < 22	Ct IC ≥ 22
Ct BN ≤ 40	Positive	Positive
40 < Ct BN < 45	Uninterpretable(*)	Unreliable (**)
No Ct	Negative	Unreliable (**)

**tab.1** shows the results interpretation

(\*) The concentration of phytoplasma could be very low in this sample, approaching the limit of detection. The results may not be reproducible. We recommend you to use another method to analyze this sample (PCR.FD BN set for example).

(\*\*) Some inhibitors of the *Taq* polymerase might be present in this sample. Repeat the analysis by diluting the DNA of this sample to 1:5 or to 1:10. If the Ct value of IC is still  $\geq 22$  after repeating the analysis on diluted sample, the DNA extraction must be repeated.

### Special handling instructions

This set was designed to be used by laboratory staff trained to follow the usual molecular biology precautions. Always perform the tests in a nuclease-free work environment. Always wear gloves when handling samples containing DNA and the components of the set. Do not touch any set components with an ungloved hand. Use appropriate laboratory disposable parts. Use nuclease-free tubes and filter tips to avoid degradation and cross-contamination. Do not use components from sets with different batch numbers in the same test procedure. Do not interchange reagents with other sets. To avoid cross-contamination, use separate rooms for (a) nucleic acids extraction, (b) preparation of the Master Mix and (c) amplification. To avoid cross-contamination and obtain reliable results, it is essential to strictly follow the protocol in this manual. Avoid unnecessary freeze-thaw cycles of the set components. Do not use reagents after their expiration date.

### Warranty and Responsibilities

Qualiplante SAS guarantees the buyer exclusively concerning the quality of reagents and of the components used to produce the Sets. Any product not fulfilling the specifications included in the product sheet will be replaced. This warranty limits Qualiplante SAS responsibility to the replacement of the product. No other warranties, of any kind, express or implied-are provided by Qualiplante SAS.

Qualiplante SAS is not responsible and cannot anyway be considered responsible or jointly responsible for possible direct and indirect damages resulting of the use and/or the misuses of the Sets. The user consciously and under her/his own responsibilities decides for the utilization purposes of the Sets and uses it the way she/he considers most suitable in order to reach her/his goals and/or objectives. Qualiplante SAS is not responsible for the data resulting from the use of the Sets, for the utilization that the user independently decides to make of them or for the direct or indirect damages possibly resulting from the disclosure or transmission of the data themselves to third parties under any form or circumstance. This clause is automatically accepted by the user when purchasing the Sets. Some of the applications which may be performed with this product may be covered by applicable patents in certain countries. The purchase of this product does not include or provide a license to perform patented applications. Users may be required to obtain a license depending on the country and/or application. Qualiplante SAS does not encourage the unlicensed use of patented applications. The Sets may require the use of *Taq* Polymerase enzyme, DNA binding components and fluorochromes/quencher, often registered as trademark by companies. The product, equipment and information included in the Sets consist of assembled reagents. The Sets are designed for the services supply, quality control or any other application that is not exclusively an internal company's research and requires a specific license for PCR and Real-Time PCR use. The license and authorization for PCR and Real-Time PCR use are not included in the Sets. The user is responsible for setting prefixed goals, choosing whether or not to perform the PCR or Real-Time PCR reaction and to apply for register her/his own license.

The Sets have been internally tested by our quality control. Any responsibility is waived if the warranty of quality control does not refer to the specific Sets. The user is personally responsible for data that she/he will obtain and/or she/he will supply to third parties using these Sets. Once the sealed package is opened the user accepts all the conditions without fail; if the package is still sealed the set can be returned and the user can be refunded.

Sets components are intended, developed, designed, and sold for Research Purpose Only. Product claims are subject to change.

### Troubleshooting

**Post-PCR data analysis shows no amplification, or amplification plots look grossly abnormal:**

Possible causes	Corrective actions
Evaporation of the sample due to inadequate sealing of the plate	Repeat the test using the appropriate tools to seal correctly the plate
Consumables are not appropriate for the method	Repeat the test using consumables recommended by the thermal cycler supplier
The quality of nucleic acid extracted is low	Repeat the extraction step. Ensure that the method of extraction has been performed correctly. In any doubt, contact us
Abnormal amplification	Centrifuge the plate briefly to spin down the contents and eliminate any air bubbles

**No amplification reaction is observed in the positive control well, while other samples are positive:**

Possible causes	Corrective actions
The positive control provided with the set was not added into the reaction well	Repeat the test. If the problem persists, contact us

**An amplification plot is observed in the negative control well:**

Possible causes	Corrective actions
Contamination of the negative control or the Master Mix with target-positive nucleic acid	Repeat the test by applying appropriate quality procedures to prevent contamination. Seal the plate correctly