

Information pest: Tomato infectious chlorosis virus

Tomato infectious chlorosis virus (TICV) is a regulated plant pathogen included in European legislation as a quarantine pest.

TICV is a whitefly-transmitted virus from tomato originally identified in the USA in 1993. Since then, the virus has spread worldwide in all major tomato-production areas. TICV is transmitted exclusively by *Trialeurodes vaporariorum*.

Tomato plants affected by the criniviruses (TICV and ToCV) show similar symptoms, which consist of interveinal yellowing beginning on lower leaves. Yellowing develops from the base to the top but is rarely visible on younger leaves. In addition, reddening, bronzing, necrosis and rolling of older leaves may be observed. Infected plants show early senescence and less vigour.

Introduction

The qPCR TICV kit has been developed and optimized by QualiPlante based on Tiberini et al., 2011. A verification was performed by QualiPlante (data not published) and the performance characteristics of the kit are the same as the original publication.

Primers and probe have been derived from the CP gene located on RNA2 of TICV. The test of Tiberini et al., 2011 was evaluated in the framework of the ARNADIA project with the following performances:

- repeatability: 100%
- reproducibility: 99%,
- diagnostic sensitivity: 100%,
- diagnostic specificity: 100%,
- relative accuracy: 100%,
- analytical sensitivity: 10⁻⁶.

The One-Step Real-Time RT-PCR test of Tiberini et al. 2011 is referred in the Appendix 2 of the [PM7/118](#) (1) Tomato chlorosis virus and Tomato infectious chlorosis virus, European and Mediterranean Plant Protection Organization Bulletin (2013) 43 (3), 462-470 and recommended by Progetto Aron-Arnadia - Protocollo per test diagnostici Tomato infectious chlorosis virus and Tomato chlorosis virus.

This product should be used only for research purposes.

Intended use

The qPCR TICV kit is validated for the detection of Tomato infectious chlorosis virus (TICV) in One-Step Real-Time RT-PCR.

Suitable tissues are tissue from symptomatic tomato leaves.

Kit format and content

Article N°	Product name
qPCR TICV 96	qPCR Tomato infectious chlorosis virus 96 tests

Content	96 tests
Direct Master Mix	2x48 tests
RT-Enzyme	96 tests
Positive Control	8 tests
Negative Control	8 tests

Storage conditions

This kit 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 kit 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)

- RNA 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 FAM[®]

Nucleic acids extraction

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

Preparation of the TICV 1-Step master mix

- Slowly thaw **Direct Master Mix** and **RT-Enzyme** by placing it on ice or at 4°C.
- Shake briefly **Direct Master Mix** and **RT-Enzyme** and spin down the liquid.
- In a new tube called **TICV 1-Step master mix**, mix 17,5 µl of **Direct Master Mix** and 0,5 µl of **RT-Enzyme** per reaction. Do not forget to count the **Positive Control** and the **Negative Control** in the number of reactions to prepare.

Example:	1 rxn	10 rxns
Direct Master Mix	17,5 µl	175,0 µl
RT-enzyme	0,5 µl	5,0 µl

- Store the **TICV 1-Step master mix** by placing it on ice or at 4°C.

Reaction set-up

- Shake briefly **TICV 1-Step master mix** and spin down the liquid.
- Add 18 µl of **TICV 1-Step master mix** (without RNA template) to each PCR tubes or wells of an optical-grade PCR plate.
- Add 2 µl of RNA template to the **TICV 1-Step 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
RNA template or Positive control or Negative control	2 µl
TICV 1-Step master mix	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)
Reverse transcription	50°C	15 min	1
Enzyme activation	95°C	10 min	1
Denaturation	95°C	15 sec	40
Annealing and elongation	60°C	60 sec	

Results analysis

The reaction for TICV will generate a specific FAM®-labeled amplification curve.

Fig.1: Example of an amplification curve relative to a TICV positive sample.

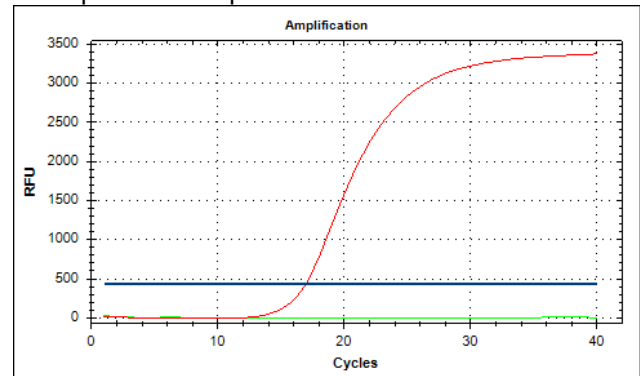


fig.1 shows the amplification curves associated to a TICV-infected sample or **Positive Control** (red curve) and to a healthy sample or **Negative Control** (green curve)

ANALYSIS VALIDATION

The PCR plate is validated only when:

- ✓ the **Positive Control** generates an amplification curve for FAM® fluorophore. The Cycle threshold (Ct) value of the FAM®-labeled amplification curve should be below or equal to 38 (**fig.1**).
- ✓ the **Negative Control** does not generate any amplification curve associated to FAM® fluorophore or a curve is generated for FAM® fluorophore but the Ct is higher than 38 (**fig.1**).

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 FAM® fluorochrome specific to TICV.

Ct	Interpretation
Ct ≤ 38	Positive
Ct > 38	Negative

tab.1 shows the results interpretation

Special handling instructions

This kit 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/RNA and the components of the kit. Do not touch any kit 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 kits with different batch numbers in the same test procedure. Do not interchange reagents with other kits. 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 kit components. Do not use reagents after their expiration date.

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 case of 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 kit 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

Warranty and Responsibilities

Qualiplante SAS guarantees the buyer exclusively concerning the quality of reagents and of the components used to produce the Kits. 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 Kits. The user consciously and under her/his own responsibilities decides for the utilization purposes of the Kits 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 Kits, 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 Kits.

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

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