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Additional Information

**Metabolic fingerprinting of Tomato Mosaic Virus infected *Solanum lycopersicum***

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## Abstract

20 <sup>1</sup>H-nuclear magnetic resonance (NMR)-based metabolomics has been applied to study  
the compatible interaction between tomato plants and tomato mosaic virus (ToMV). A  
detailed time course of metabolic fingerprinting of ToMV-inoculated and non-  
inoculated systemically infected tomato leaves has provided a fundamental  
understanding of the metabolic state of the plant not only in response to ToMV  
25 infection, but also under various physiological conditions. By this analytical platform a  
total of 32 metabolites including amino/organic acids, sugars, phenylpropanoids,  
flavonoids and other miscellaneous compounds were detected. Using multivariate data  
analysis, we have identified a subset of metabolites induced during the plant defence  
response and metabolites whose accumulation was dependent on the developmental  
30 stage, the position of the leaf on the stem, and the harvesting time. Specifically, a  
general time-dependent decrease in organic acids, amino acids (excluding asparagine),  
phenylpropanoids and rutin was observed in individual leaves. In addition, metabolite  
alterations were also found to correlate with the developmental stage of the leaf: high  
levels of organic acids, some amino acids, phenylpropanoids, and flavonoids were  
35 found in lower leaves while elevated amounts of sugars were present in the upper ones.  
Moreover, a marked variation in the content of some metabolites was also observed to  
be associated to the asymptomatic ToMV infection both in inoculated and systemically  
infected leaves. While flavonoids accumulated in virus-inoculated leaves, increased  
levels of phenylpropanoids were observed in non-inoculated leaves where ToMV  
40 actively replicates. Finally, diurnal changes in the metabolite content were also  
observed: an increase of amino acids and organic acids (except glutamic acid) were  
observed in the samples collected in the morning, whereas sugars and secondary  
metabolite levels increased in the tomato leaves harvested in the evening.

45 **Keywords:** *Solanum lycopersicum*; *Tomato Mosaic Virus (ToMV)*; *plant-pathogen interaction*; *systemic acquired resistance (SAR)*; *NMR-based metabolomics*

## Introduction

Higher plants are, in most cases, passive receivers of biotic (viroids, viruses, bacteria,  
50 fungi, nematodes or insects) and abiotic (salinity and drought) stresses. Consequently,  
they have evolved a large variety of sophisticated defence mechanisms to resist different  
types of stress (Dixon, 2001; Dangl and Jones, 2001).

Two types of plant-microbe interactions can be defined: incompatible and compatible.  
Incompatible interactions have been extensively studied principally because of their  
55 direct and practical applications in the field (Vlot et al., 2009), whereas much less  
consideration has been paid to the compatible interactions (O'Donnell et al., 2001;  
Huang et al., 2003).

In an incompatible interaction, after an initial perception of the pathogen by the host,  
defences are rapidly activated resulting ultimately in the so-called hypersensitive  
60 response, which culminates in localized cell death at the area of pathogen inoculation  
and, concomitantly, the development of different degrees of necrosis (Ryals et al.,  
1996). The pathogen infection is thus limited to small parts immediately surrounding  
the initially infected cells and signals are generated which stimulate defensive reactions  
in uninfected areas of the plant. These distal sites then become more resistant to  
65 subsequent infections. This phenomenon, known as systemic acquired resistance (SAR),  
results in an immunization against future microbe attacks (Sticher et al., 1997; Durrant  
and Dong, 2004).

Frequently, a defensive response is also induced in a compatible interaction between a  
susceptible host and a virulent pathogen. However, in this case, in the absence of gene-  
70 for-gene resistance, no necrosis occurs (Conejero et al., 1990; Dixon et al., 1994). In  
this type of interaction microbes actively multiply and spread throughout the host  
leading to disease and even the death of the infected plant.

Although differing in the speed of the response to the pathogens, incompatible and compatible interactions share very similar general and common features, such as the  
75 induction of the plant signalling compounds, such as salicylic acid (SA; 2-hydroxybenzoic acid) (Sticher et al., 1997; Vlot et al., 2009) and ethylene, and the synthesis of pathogenesis-defence proteins (van Kan et al., 1992; van Loon et al., 2006). One of the most well established defence responses in plants is the biosynthesis of an often complex array of natural chemicals belonging to phenolic metabolism (Hahlbrock  
80 and Scheel, 1989; Dixon et al., 2002; Abdel-Farid et al., 2009). A pending goal in molecular plant pathology is to characterize the metabolites that play a role in the defence response against pathogens. Traditionally, the analytical techniques employed in the search for these metabolites were basically chromatographic, coupled to spectroscopic detectors or mass spectrometers (Baumert et al., 2001; Shadle et al., 2003;  
85 Tan et al., 2004). Accordingly, the range of metabolites that can be identified using these techniques is restricted based on the case specific experimental procedures employed and the physical-chemical properties of the compounds investigated (Roepenack-Lahaye et al., 2003; Bednarek et al., 2005; Fayos et al., 2006; Zacarés et al., 2007; Bellés et al., 2008; López-Gresa et al., 2011). However, a considerable  
90 amount of biomolecules with different biological functions (and physical-chemical properties), are proposed to be involved in the physiology of plant/pathogen interactions. Therefore, more robust analytical systems need to be developed in order to detect and accurately identify this considerable number of primary and secondary metabolites (Glauser et al., 2010).  
95 The introduction of so-called metabolomics has allowed the relatively rapid, and at the same time thorough, detection of a vast range of metabolites, thus providing an in-depth analysis of the total metabolome of biological processes in the plant. Although it is impossible for one single analytical method to completely characterize the entire plant

metabolite profile, nuclear magnetic resonance spectroscopy (NMR) is an excellent tool  
100 to provide a macroscopic view of the majority of the components of the plant  
metabolome (Verpoorte et al., 2008; Broyart et al., 2010; Kim et al., 2010; Leiss et al.,  
2011). In fact, NMR-based metabolomics has been widely used to decode the broad  
range of chemical compounds that might be implicated in the plant defence against  
microbe attacks (Choi et al., 2006; Ward et al., 2010). Our group has been applying this  
105 technology to the detection and structure elucidation of metabolites in plants upon  
treatment with a wide range of biotic and abiotic stress agents (Choi et al., 2004; Simoh  
et al., 2009; Jahangir et al., 2009; Mirnezhad et al., 2010; Simoh et al., 2010). In recent  
years, very detailed NMR metabolomic studies on both compatible and incompatible  
plant-pathogen interactions have been reported (Lima et al., 2010; Bollina et al., 2010;  
110 Ward et al., 2010; Jones et al., 2011).

Virus diseases cause serious losses worldwide in horticultural and agricultural crops  
(Lomonosoff, 1995). The cultivated tomato is subjected to a range of diseases resulting  
from infection with certain strains of tobacco mosaic virus. The mosaic disease in  
tomato is the most persistent virus in terms of its ability to survive outside plant cells  
115 and in dead tissues (Broadbent, 1976). For this reason and because of its easy  
transmission this virus has a high rate of infectivity. ToMV is the most troublesome  
viral disease of tomatoes, causing distortion of leaves and fruit and stunting of growth.  
However, few metabolomic reports on plant/virus interactions have been described in  
the literature, and a complete analysis of metabolites has been carried out only in the  
120 case of the hypersensitive response of tobacco plants to tobacco mosaic virus (TMV)  
(Choi et al., 2006). In our previous report, some characteristic metabolites of tomato  
plants related to either Citrus Exocortis viroid (CEVd) or *Pseudomonas syringae*  
infection have been identified by NMR-based metabolomics (López-Gresa et al., 2010).

Tomato and Tobamovirus ToMV constitutes another model plant-virus interaction  
125 system. But, as far as we know, no study on metabolic fingerprinting of tomato  
associated with plant development and its interaction with the compatible Tobamovirus  
ToMV has yet been published. Therefore, tomato plants infected by ToMV were  
studied at different times post-inoculation in order to extend our knowledge to  
additional plant/virus interactions.

130 Using the *Solanum lycopersicum*-ToMV interaction as a basic model system, we aim to  
reveal the role of defence metabolites involved in this infection by NMR spectroscopy  
combined with multivariate data analysis. As part of an ongoing global investigation on  
the metabolic state of tomato plants under different biotic or abiotic stress conditions,  
the main aim of this high throughput NMR-based metabolic analysis is geared to  
135 increase the knowledge of the tomato ToMV infected leaf metabolome in diverse states  
of plant development and disease progression. A better understanding of the tomato  
defence response might also provide relevant data for metabolic engineering of tomato  
plants with higher resistance towards pathogens.

## 140 **Experimental**

### *Plant material and inoculation procedure*

Seeds from tomato (*Solanum lycopersicum* cv. Rutgers) (Western Hybrid Seeds Inc.,  
145 Hamilton City, CA, U.S.A) previously surface-sterilized with bleach were used in this  
work. Extreme care was taken in order to maintain controlled conditions of growth to be  
sure that the variables of study were limited to the effect of the infection and the natural  
growth of the plants. Experimental lots of 120 plants of similar morphological and  
physiological characteristics were prepared for the experiment. The plants (one per pot)



150 were grown in 15-cm-diameter pots containing a mixture of peat (Biolan) and  
vermiculite 1:1 and were subirrigated once a day. The tomato plants were maintained in  
a controlled growth chamber at a constant temperature of 24 °C (12 h photoperiod) and  
with relative humidity ranging from 60% (day) to 70% (night). Five-week-old Rutgers  
tomato plants at the five- to six-leaf stage were used in all the experiments described in  
155 this article. Infection of Rutgers tomato plants with the Murakishi PV-0143 (DSMZ)  
strain of Tomato Mosaic Virus was performed with a viral extract obtained from leaves  
of ToMV-infected tomato plants that were homogenized in 10 mM phosphate buffer  
(pH 7.2), 0.5% sodium bisulfite, 0.5% diethyldithiocarbamic acid (1 g of leaf material  
in 20 mL of buffer) as described (Bellés et al., 2006). The third leaf (leaf position  
160 numbered from the base to the apex) of tomato plants were dusted with carborundum  
(particle size 0.037 mm). One ml of viral extract or buffer (mock-inoculated control)  
was applied by gently rubbing the upper face of the third leaf. The third and fourth  
leaves were harvested separately at day 1, 2, 3, 4, 5, 7, 9, 11, 13, and 16 post-  
inoculation at 9 am (9:00 h) and 7 pm (19:00 h) each day. Samples were immediately  
165 frozen in liquid nitrogen, subsequently ground in a cooled mortar and lyophilized for  
metabolomic studies. Three biological replicates from both mock- and ToMV-  
inoculated tomato plants were analyzed for each time point.

#### *Protein extraction and electrophoretic analysis*

170

Leaves from mock-inoculated and ToMV-infected tomato leaves were harvested at the  
indicated times and rapidly frozen in liquid nitrogen and stored at -80 °C until use.  
Protein extracts of tomato leaves were performed by homogenization in extraction  
buffer (50mM Tris, pH 7.5, 15mM mercaptoethanol), as described in (Rodrigo et al.,

175 1993). Proteins were separated by SDS-PAGE and stained with Coomassie Brilliant Blue R-250, as described (Conejero and Semancik, 1977).

#### *Extraction and NMR spectra measurements*

180 Twenty five mg of freeze-dried plant material were extracted in 2 mL-Eppendorf tubes with 1 mL of a mixture of  $\text{KH}_2\text{PO}_4$  buffer (pH 6) in  $\text{D}_2\text{O}$  containing 0.05% trimethylsilane propionic acid sodium salt (TMSP) and  $\text{CH}_3\text{OH-}d_4$  (1:1). The extracts were vortexed, vigorously, sonicated for 20 min and then centrifuged at 13,000 rpm for 10 min. Seven hundred  $\mu\text{L}$  of the supernatant were transferred in 5 mm-NMR tubes for  
185 the spectral analysis.

$^1\text{H}$  NMR spectra were recorded at 25 °C on a 600 MHz Bruker AV 600 spectrometer equipped with cryo-probe operating at a proton NMR frequency of 600 MHz. Methyl signal of  $\text{CH}_3\text{OH-}d_4$  was used as the internal lock. Each  $^1\text{H}$  NMR spectrum consisted of 128 scans requiring 10 min acquisition time with the following parameters: 0.25  
190 Hz/point, pulse width (PW) = 30 ° (10.8  $\mu\text{s}$ ), and relaxation delay (RD) = 1.5 s. A pre-saturation sequence was used to suppress the residual  $\text{H}_2\text{O}$  signal with low power selective irradiation at the  $\text{H}_2\text{O}$  frequency during the recycle delay. Free induction decay (FIDs) were Fourier transformed with Line Broadening (LB) = 0.3 Hz and the spectra were zero-filled to 32 K points. The resulting spectra were manually phased and  
195 baseline corrected, and calibrated to TMSP at 0.0 ppm, using Topspin (version 2.1, Bruker).

#### *Data analysis*

200 <sup>1</sup>H NMR spectra were automatically reduced to ASCII files using AMIX (v. 3.7, Bruker  
Biospin). Spectral intensities were scaled to total intensity TMS<sub>P</sub> and reduced to  
integrated regions of equal width (0.04 ppm) corresponding to the region of δ 0.4 – δ  
10.00. The region of δ 4.7 - δ 4.9 was excluded from the analysis because of the  
residual signal of water as well as δ 3.28 - δ 3.34 for residual methanol. Partial least  
205 square (PLS) was performed with the SIMCA-P software (v. 11.0, Umetrics, Umeå,  
Sweden) using unit variance (UV) scaling method.

## Results and Discussion

### 210 *Progression of ToMV infection*

The ToMV strain employed in this work failed to induce visible symptoms in the  
tolerant tomato cultivar Rutgers during the 16 day experiment. Only subtle curling,  
slight distortions and malformations of the leaves and a very limited stunting of the  
215 plants were observed at long periods post-inoculation. To monitor the progress of the  
ToMV disease in the infected tomato plants, a polyacrylamide gel electrophoresis  
(SDS-PAGE) of protein extracts from the third-inoculated and systemically fourth-  
infected leaves was performed to detect the viral capsid accumulation. The knowledge  
of the virus accumulation pattern at each time point allows us to relate the metabolic  
220 changes observed during the infection with the amount of virus present. ToMV actively  
multiplied in the inoculated leaf, and the 17-kDa ToMV coat protein could be clearly  
detected at day three post-inoculation, as assessed by electrophoretic analysis (Fig. 1).  
In the inoculated leaves, ToMV accumulation increased in a time-dependent manner,  
reaching its maximum level at 5 day post-inoculation, and decreasing thereafter (data  
225 not shown). The virus started to spread up to the fourth leaf by day 7, following a

similar induction kinetic pattern to the inoculated leaves and remaining constant from day 11 until the end of the 16-day experiment (Fig. 1). The phenotypic alterations observed consisted mainly of gradual development of brown necrotic spots at the inoculation site (produced by rubbing the leaves with carborundum in the inoculation zone) in both the inoculated and mock-inoculated leaves.

### **Alterations of the metabolic profiles of leaves induced by ToMV infection and developmental conditions**

Diverse factors such as light intensity, temperature, water, nutritional status can produce notable effects on the metabolome during the life cycle of the plant. In metabolic studies it is crucial to be able to differentiate the changes that result from uncontrolled factors and those that are genuinely caused by the experimental manipulation being studied. In the present work, we studied the leaf metabolome affected by ToMV infection. The experimental conditions were carefully designed and controlled to ensure that the metabolic differences identified were restricted to only the effect of ToMV infection. However, in addition, we expanded our study to include other non-controllable factors such as plant aging and development stage. Interestingly, these factors had very important effects on the metabolome of tomato leaves.

Plant metabolites have very diverse physical-chemical properties and are present in a wide range of quantities (Wolfender, 2009). Therefore, the solvent extraction used for their isolation from the plant tissues should be able to rapidly and efficiently dissolve metabolites in an unbiased manner. In this work, tomato leaf metabolites were extracted using a single-step extraction method consisting of a polar solvent mixture of methanol and a buffered aqueous solution. This method is practical, is not time consuming and has been shown to possess a high extraction efficiency constitutes an optimized

extraction protocol for polar untargeted metabolites (Hendrawati et al., 2006; Verpoorte et al., 2007; Verpoorte et al., 2008). NMR spectroscopy in combination with multivariable data analysis was performed on control and ToMV-infected tomato leaves  
255 in order to determine the metabolic changes during this plant-pathogen interaction.

The first step of NMR metabolomic studies is the analysis of the plant extract spectra (Choi et al., 2006). <sup>1</sup>H NMR spectra, 2D NMR experiments (2D *J*-resolved, COSY and HMBC), our in-house database of reference compounds and previous reports (López-Gresa et al., 2010) were used for the signal assignment of metabolites from the tomato  
260 leaves. Table 1 shows a list of 32 the identified metabolites including 13 aminoacids, 5 organic acids (3 citrate-cycle intermediates), 3 sugars, 4 phenylpropanoids, 1 flavonoid, and 6 miscellaneous compounds with their characteristic chemical shifts and coupling constants.

After the chemical analysis of the spectra, the following step in a metabolomic study is  
265 the comparison of all the samples through multivariate data analysis (MVDA).

The first step of the statistical analysis of these large data sets (in this case 240 samples), was the application of a principal component analysis (PCA) in order to group the plants according to the NMR signals characteristic for the metabolic changes caused by the ToMV infection. In the PCA score plot, healthy and  
270 infected samples were grouped together, showing that only slight metabolic changes were produced by the infection. Other parameters, such as time (from 1 to 16 days post inoculation (dpi), hereafter considered as age), position in the stem (lower and upper leaves) or collection time (9:00 h and 19:00 h) play an important role in determining the metabolome, even more than the infection.

275 According to this observation, a partial least square analysis (PLS) was carried out defining as the X variable the area of binned <sup>1</sup>H-NMR signals and as step-wise Y variables age, developmental stage, infection, and sample collection

time. A clear separation was obtained between: a) age of the leaves by component 1 (Fig 2); b) leaves located at different positions in the stem by component 2 (Fig 2); c) healthy control and ToMV infected leaves by component 3 (Fig 3); and d) samples collected at 9:00 h and 19:00 h by component 4 (Fig 4). In order to differentiate the identified compounds, a loading plot was used in which the correlation between grouping and correlated metabolites was shown (Table 1).

285

The first component of PLS explained the changes in the chemical composition during the time course of the experiment (age of the plant: 1 to 16 dpi). Analysis of loading column plot of PLS component 1 showed a time-dependent increase in the content of sugars and a decline of the organic and amino acids, except for asparagine. This observation is in agreement with the results obtained examining the amount of several amino acids in different lines of *Arabidopsis* during the leaf senescence (Diaz et al., 2005). Only asparagine was higher in “old” leaves (from 9 to 16 dpi), in accordance with the well-known decrease of asparaginase activity observed during leaf maturation (Sieciechowicz et al., 1988) (Fig 2 and Table1). The increase of sugars observed in the “old” leaves may be correlated to the increase of total surface area in all leaves with age, which augments the rate of photosynthesis, thus producing a greater accumulation of carbohydrates (Abdel-Farid et al., 2009). Important concentrations of phenylpropanoids and the flavonoid, rutin were found in “young” leaves (from 1 to 9 dpi). These secondary metabolites are associated with defence mechanisms and their presence in the “young” leaves reflects the need of this tissues to increase their defensive potential (Brown et al., 2003; Baker et al., 2010). According to the optimal plant defence theory, plants tend to allocate more defence-associated metabolites to the

valuable plant parts during development, to protect them from stress factors and to maximize their fitness (Kaur et al., 2010).

305 Examination of the component 2-PLS loading plot showed the metabolites which strongly contributed to the separation of the metabolic profiles based on the position of the leaf on the stem. Lower (third) leaves were placed on the positive side of PLS component 2, whereas immediately upper (fourth) leaves were on the negative side (Fig 2). When comparing the metabolome of the leaves located at different positions in the stem, it was observed that organic acids (malic, citric, succinic, fumaric, and formic 310 acid), amino acids (GABA, glutamic, glutamine, aspartic, asparagine, tryptophan, and phenylalanine), phenylpropanoids, and flavonoids were induced in lower leaves, while upper leaves showed higher amount of sugars, acetic acid, choline, trigonelline and the aliphatic amino acids arginine, alanine, and valine (Table 1). Many primary metabolites 315 were induced as a consequence of the abrasive carborundum applied during inoculation process, together with secondary metabolites, in local leaves (Dixon and Paiva, 1995). However, sucrose and glucose levels decreased in these lower leaves. Presumably, carbohydrate consumption is required for production of energy to support the biosynthesis of defensive phenolic compounds induced by wounding (Broeckling et al., 320 2005). Since the metabolite content varies throughout the plant (Choi et al., 2004), it is crucial to harvest the leaves in the same leaf position in order to accurately compare metabolite fingerprinting of infected with the corresponding equivalent control leaves. Focused on our aim, which is to understand the tomato defence reaction against ToMV infection, a detailed metabolic study was performed examining component 3 of the 325 PLS. The score plot for component 1 versus component 3 of PLS from the  $^1\text{H}$  NMR spectra shows that ToMV-infected leaves are separated from healthy ones, independently of whether they were directly inoculated with the virus or systemically infected (Fig 3). The separation is mainly due to component 3, however, both aging and

stem position produce a shift along component 1. To exclude differences due to this  
330 biological variability, partial least square-discriminant analysis (PLS-DA) was applied  
separately on 1) mock-inoculated (third) versus the ToMV locally-infected (third)  
leaves until eleven dpi (Fig 5A) and 2) upper-mock-inoculated (fourth) versus  
systemically-infected (fourth) leaves from five to thirteen dpi (Fig 5B). Applying this  
statistical analysis, a good separation was observed between local and systemically  
335 infected leaves.

Local infection was characterized by component 1 (Fig 5A), in which rutin, glucose,  
choline, GABA, glutamine, asparagine, phenylalanine, glutamic, aspartic, malic, citric,  
formic, and fumaric acid, were found to be the major contributing metabolites (Table 1).  
Determination of the relative abundance of several amino acids in ToMV-infected  
340 tomato leaves revealed a decrease in the content of some of them and an increase in  
others. The results previously found in *P syringae*-infected-tomato leaves, in which in  
general high levels of some amino acids were found, (Pérez Garcia et al., 1998; López-  
Gresa et al., 2010), are in contrast with those found in this work upon viral infection  
which produced a general decrease in the pool of amino acids. Only  
345 glutamine/glutamate and asparagine/aspartate were clearly induced in leaves inoculated  
with ToMV. It is well established that glutamine may act as substrate for asparagine  
biosynthesis, so their levels could be interconnected (Eason et al., 1996). These results  
are in accordance with previous reports suggesting that glutamate is a precursor of  
stress-related compounds in plants (Hothorn et al., 2006) and that glutamine content  
350 was higher in *Vitis vinifera* plants resistant to mildew (Figueiredo et al., 2008). In this  
context, it is pertinent to point out that opposite results have been reported in some plant  
pathogen interactions. Thus, as observed in the present work (Table 1), and previously  
(Pérez-García et al., 1998; López-Gresa et al., 2010), the alanine content significantly  
decreases in tomato leaves after systemic infection with viral or bacterial pathogens. By



355 contrast, alanine exhibited the largest increase by far in rice plants upon infection with  
the fungal pathogen *Magnaporthe oryzae* (Jones et al. 2010), and as such, alanine is  
considered as a key marker in this compatible interaction.

GABA induction in plants in response to an abiotic stress treatment is a widely  
observed phenomenon (Wallace et al., 1984; Mayer et al., 1990; Allan et al., 2008). Our  
360 results agree with a previously published study of the metabolic changes in vitis plants  
affected by the Esca disease. In this report, the authors identified a marked increase in  
GABA as a clear indicator of healthy or diseased leaves (Lima et al., 2010). Moreover,  
increases in phenylalanine, which is the substrate of the key enzyme of the  
phenylpropanoid biosynthesis pathway, phenylalanine ammonia lyase (PAL), correlated  
365 well with the concomitant induction of phenylpropanoid pathway products. The positive  
relationship between sucrose and UDP-glucose may be explained because both  
metabolites can be reversibly formed by the sucrose synthase enzyme. The primary  
metabolites, malic and citric acid were found in the present study to be altered in both  
locally inoculated and systemically -infected tomato leaves. Our results are consistent  
370 with those of Choi et al. (2006), studying the hypersensitive interaction between  
tobacco and tobacco mosaic virus (TMV). The authors reported that these organic acids  
also accumulated in both TMV-inoculated and in SAR tobacco leaves.

The study of phenolic compounds during the progression of infection is very relevant  
based on their proposed defensive role in plants. The flavonoid, rutin, which has been  
375 related to the enhancement of the defence system against some stress conditions (Suzuki  
et al., 2005), clearly increased in ToMV-infected tomato leaves. Previously, we had also  
found that citrus exocortis viroid and *P. syringae* also elicited the synthesis of this  
compound in tomato leaves (López-Gresa et al., 2010). Unexpectedly, caffeoyl esters of  
polyhydroxy compounds (glucaric acid) did not accumulate in locally infected tomato  
380 leaves, even though phenylpropanoid metabolism is activated under various stress

conditions, such as UV irradiation, mechanical wounding, or pathogen attack (Lawton and Lamb, 1987; Dixon and Lamb, 1990). In fact, TMV-infected tobacco plants showed, among the phenylpropanoids, only 5-*O*-caffeoylquinic acid accumulation in the locally infected leaves (Choi et al., 2006). Additionally, a decrease content of

385 tryptophan was observable in these ToMV-inoculated leaves. Products of the tryptophan pathway are metabolized into auxin, glucosinolates, phytoalexins, alkaloids, and other indolic compounds, which play diverse roles in plant biological processes, including plant-pathogen interaction (Radwanski and Last, 1995). Previously, NMR spectroscopy-based metabolomics had shown that the levels of a natural derivative of salicylic acid,

390 2,5-dihydroxybenzoic acid (gentisic acid), markedly increased as a conjugated form named gentisic acid 5-*O*- $\beta$ -D-xyloside in tomato leaves infected by the citrus exocortis viroid. Infection with this viroid causes strong symptoms (extreme epinasty, curling, and rugosity of leaf and leaflets) and the intensity of symptoms exhibited a good correlation with gentisic acid content in viroid- and virus-infected plants (Bellés et al.,

395 1999; Bellés et al., 2006). It is therefore not surprising that the present asymptomatic infection caused by ToMV did not induce a significant accumulation of gentisic acid. Viruses produce a vast array of symptoms in plants, and many of them appear to be common even among combinations between diverse virus and plants. Some virus-infected plants may show no symptoms, as in the case of Rutgers tomato infected by

400 this strain of ToMV, and consequently, are excellent virus carriers, as they go unnoticed in the greenhouse. Symptoms could also be influenced by several factors such as environmental conditions, plant age at infection or virulence of the virus. The present NMR based metabolomic study not only gives important and broad physiological information, but also points out that care must be taken to select virus-free plants when

405 performing metabolic analysis.

Metabolic alterations of non-inoculated systemic leaves were compared with the corresponding controls analogously to locally infected leaves by PLS-DA. In this case, upper healthy and ToMV systemically infected leaves from five to eleven thirteen dpi were used as discrete classes (Fig 5B). The selected time course was based on the time  
410 required to detect viral capsid accumulation in distal leaves (Fig1). In this case, for the identification of the metabolites involved in systemic defence response after virus inoculation, the loading plot of component 1 was analyzed. High concentrations of phenylpropanoids, rutin, choline, organic acids, sucrose, GABA, tryptophan, phenylalanine, glutamic, and aspartic acid were characteristic of systemically infected  
415 leaves. Only glutamine and asparagine were not induced in these leaves compared with inoculated ones. Unexpectedly, in the upper control leaves there was an increase in the levels of some metabolites with signals at  $\delta$  8.02, 7.90, 7.42, 6.78, 6.14, and 6.10, belonging to some flavonoid glycosides, which probably have a 4'-OH in the B ring (Fig 6).

420 All these results suggest that flavonoids might be involved in the local response, while phenylpropanoids could play a defensive role in the distal leaves.

In a previous study, Niehl et al. (2006) reported a detailed analysis of defence responses during the compatible interaction of potato plants between potato (*Solanum tuberosum* L. cv Desiree) and the Potato Virus X (PVX). As in Rutgers tomato plants infected with  
425 Tomato Mosaic Virus (ToMV), infection of potato with PVX results in a systemic infection of the plants. In agreement with the results presented here, the authors found that low molecular weight secondary metabolites such as  $\beta$ -phenylethylamine-alkaloids were substantially induced upon PVX infection. By contrast, primary metabolism as levels of organic acids (citrate-cycle intermediates), sugars, and aminoacids remained  
430 mostly unchanged in PVX-infected potato leaves, this being in contrast with the results observed in ToMV-infected Rutgers tomato leaves (Table 1). Taken together, all these

results demonstrate that potato and tomato respond to systemic virus infection in a different way depending of the host plant.

Finally, the metabolic alterations due to sample collection time were identified by  
435 studying component 4 of PLS analysis (Fig 4). A clear separation by component 4 was  
observed between tomato leaves collected at 9:00 h and 19:00 h, mainly from the  
youngest plants (from 1 to 9 days post-inoculation). It is well known that fluctuations of  
primary and secondary metabolites are observed when plants are harvested at different  
times of the day. NMR metabolomic analysis of *Cannabis sativa* clearly demonstrates a  
440 clear difference between samples harvested in the morning and afternoon (Kim and  
Verpoorte, 2010). In our study, amino acids and organic acids (except glutamic acid)  
were induced in the samples collected in the morning, while sugars and secondary  
metabolites were induced in the tomato leaves harvested in the evening. This  
carbohydrate production is likely a result of the photosynthesis that takes place during  
445 the day. A complete quantitative analysis of carbohydrates and primary metabolites was  
done through a diurnal period in potato leaves (Urbanczyk-Wochniak et al., 2005).  
Although it is difficult to compare our results with those obtained in potato because we  
have not performed a quantitative analysis, levels of carbohydrates, particularly sucrose,  
and citric acid followed the same trend as in potato leaves.

450 As shown in this work, a broad variety of primary and secondary compounds can be  
detected by using simple extraction procedures and NMR spectroscopy. Multivariate  
data analysis methods, such as partial least square (PLS) and partial least square-  
discriminant analysis (PLS-DA) were applied to characterize the metabolic alterations  
of infected symptomless tomato leaves, taking into account their development stage and  
455 the time of harvesting.

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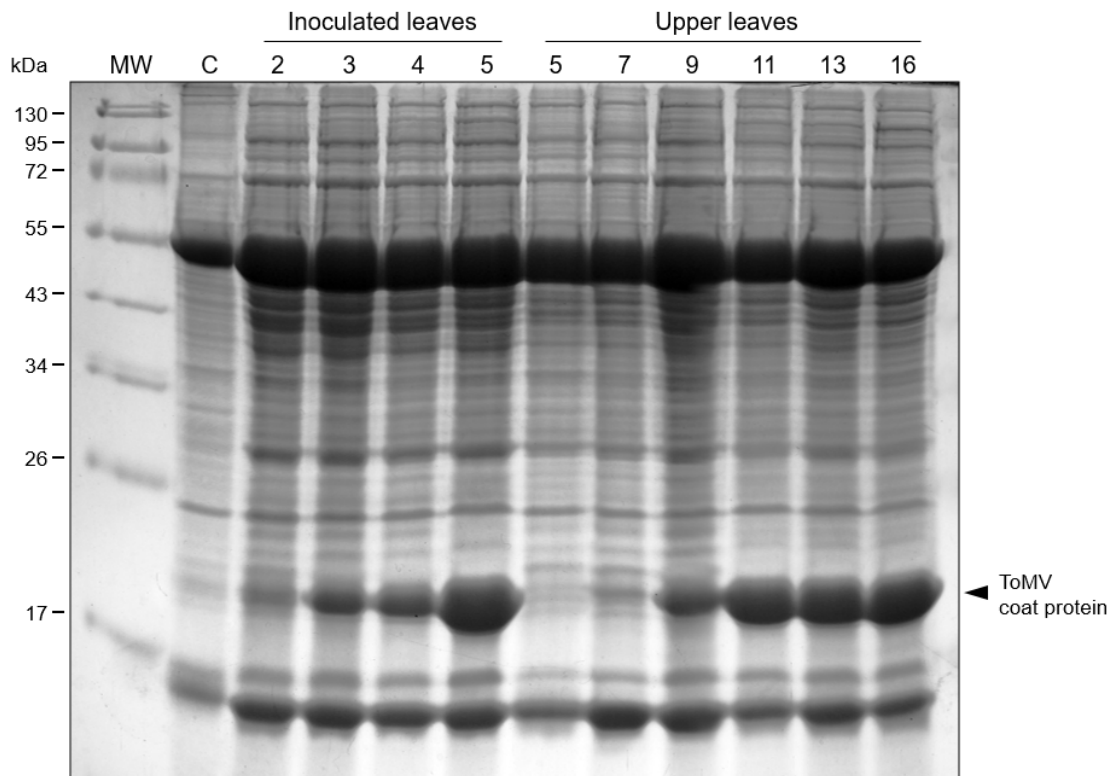
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## Figures

**Figure 1.** SDS-PAGE analysis of soluble proteins in extracts from Tomato Mosaic

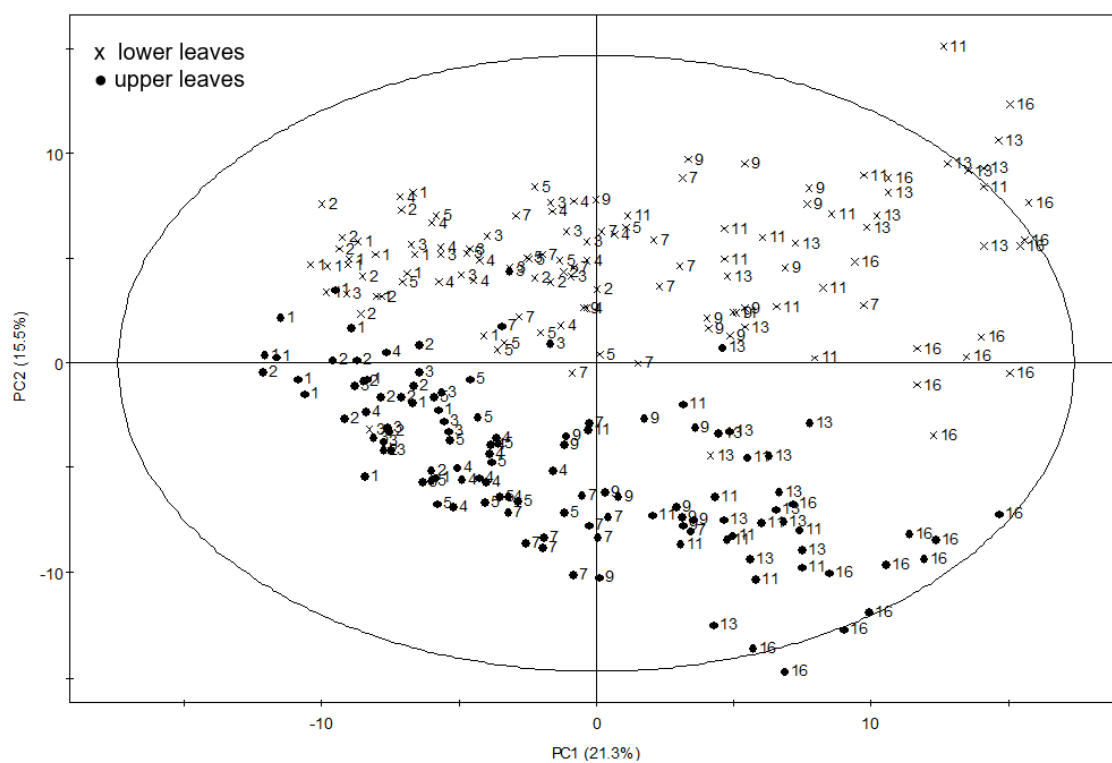
675 Virus (ToMV)-inoculated and the corresponding upper systemically infected leaves at the indicated time (days) post-inoculation. Protein size markers (kDa) are indicated on the left. Arrow on the right indicates the ToMV coat protein.



680 **Figure 2.** Score plot of PLS based on the whole range of the  $^1\text{H}$  NMR signals in the range of  $\delta$  0.3-10.0. The numbers indicate the days after mock and ToMV inoculation.

(x) lower leaves of mock-and ToMV-inoculated tomato plants

(●) upper leaves of healthy control and ToMV-infected tomato plants.

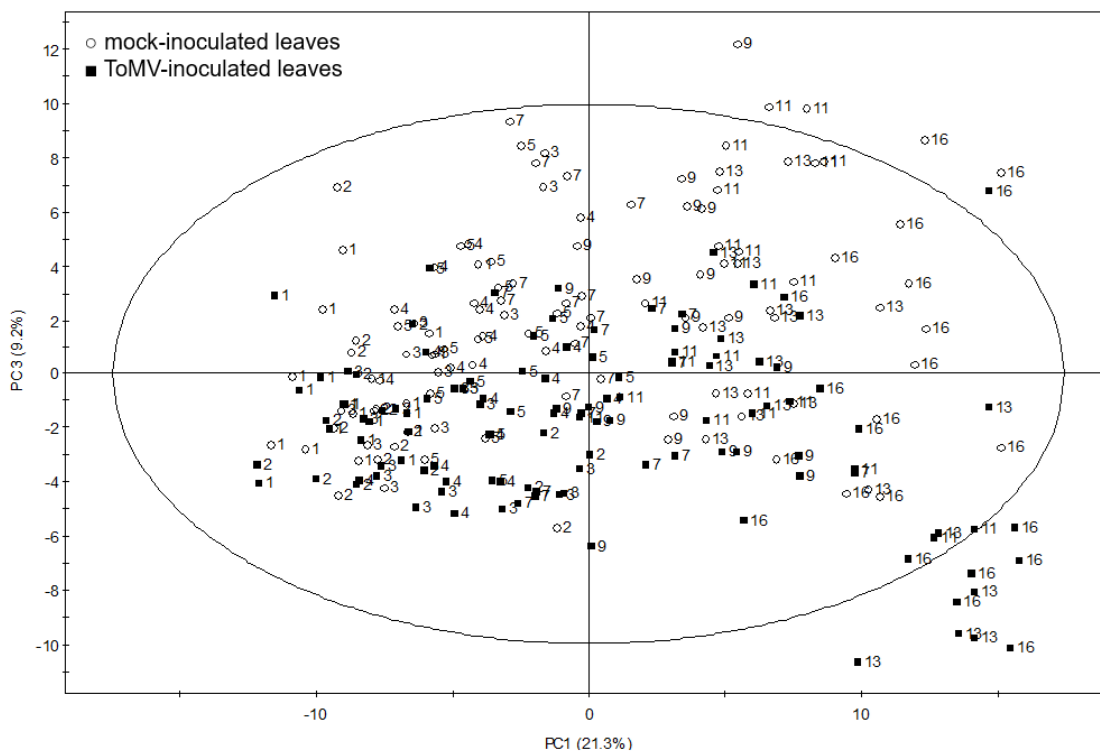


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**Figure 3.** Score plot of PLS based on the whole range of the  $^1\text{H}$  NMR signals in the range of  $\delta$  0.3-10.0. The numbers indicate the days after mock and ToMV inoculation.

(○) lower mock-inoculated and upper leaves of healthy tomato plants

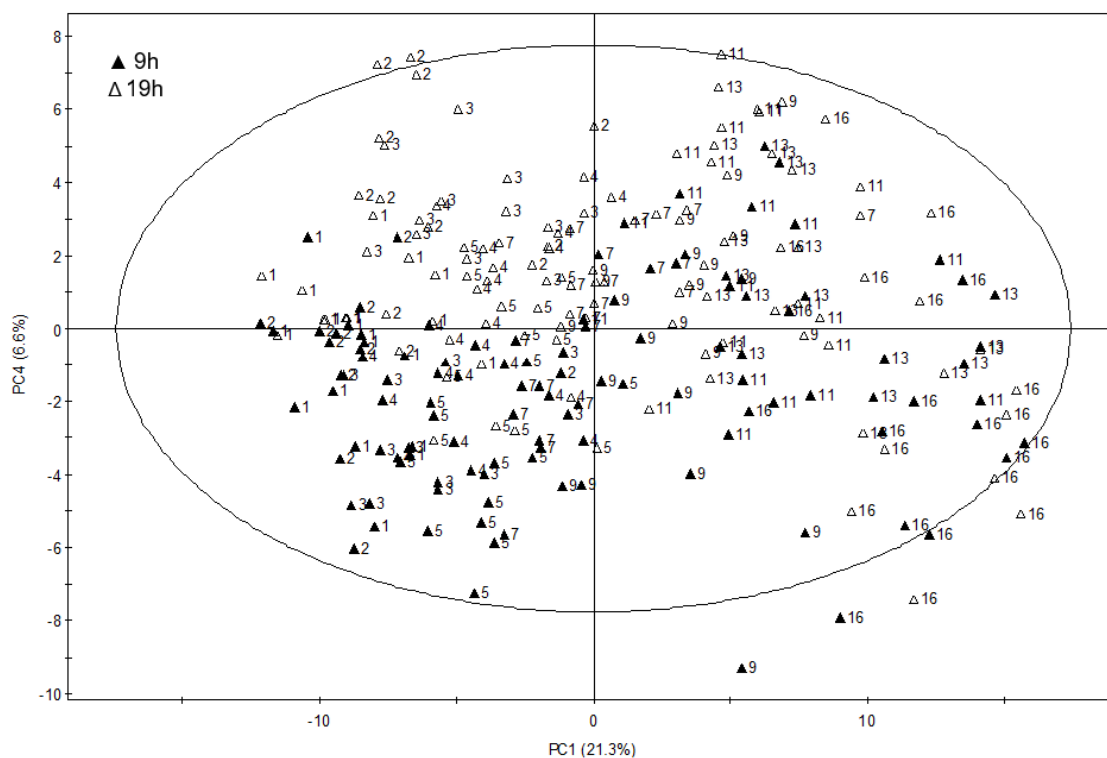
(■) lower ToMV-inoculated and upper systemically-infected leaves of tomato plants.



**Figure 4.** Score plot of PLS based on the whole range of the  $^1\text{H}$  NMR signals in the range of  $\delta$  0.3-10.0. The numbers indicate the days after mock and ToMV inoculation.

(▲) leaves harvested at 9:00 h from mock-inoculated and ToMV-infected tomato

695 (Δ) leaves harvested at 19:00 h from mock-inoculated and ToMV-infected tomato plants.

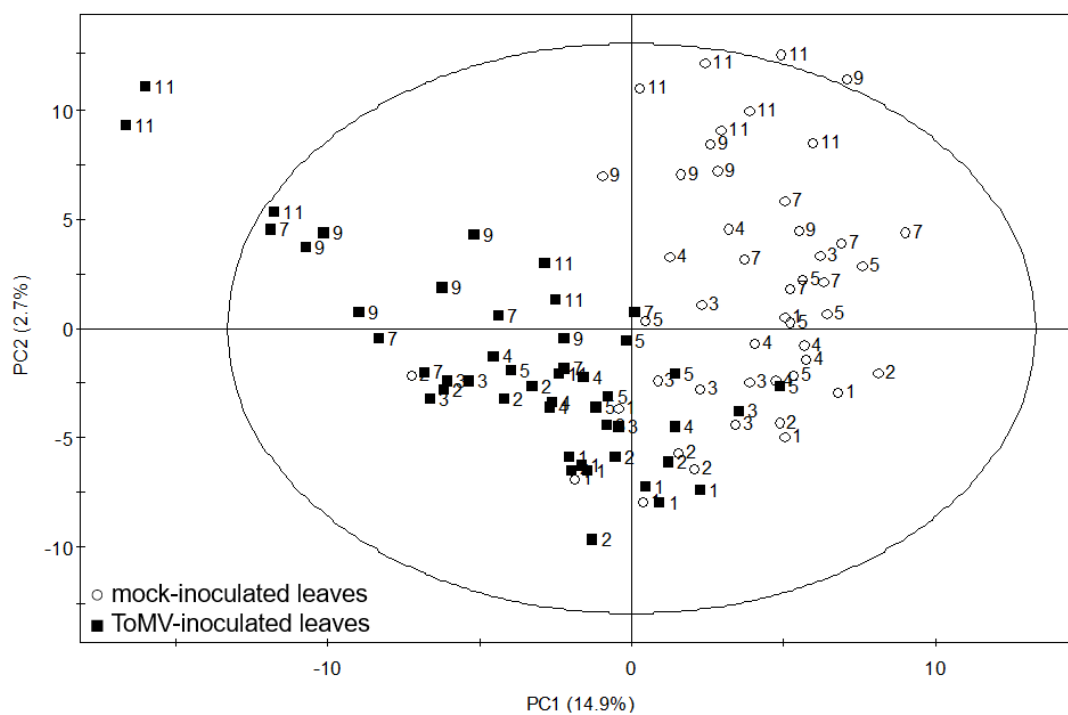


**Figure 5A.** PLS-DA score plot of early local viral infection based on the  $^1\text{H}$  NMR

700 signals in the range of  $\delta$  0.3-10.0. The numbers indicate the days after mock and ToMV inoculation.

(○) mock-inoculated leaves of healthy tomato plants

(■) ToMV-inoculated leaves of infected tomato plants.



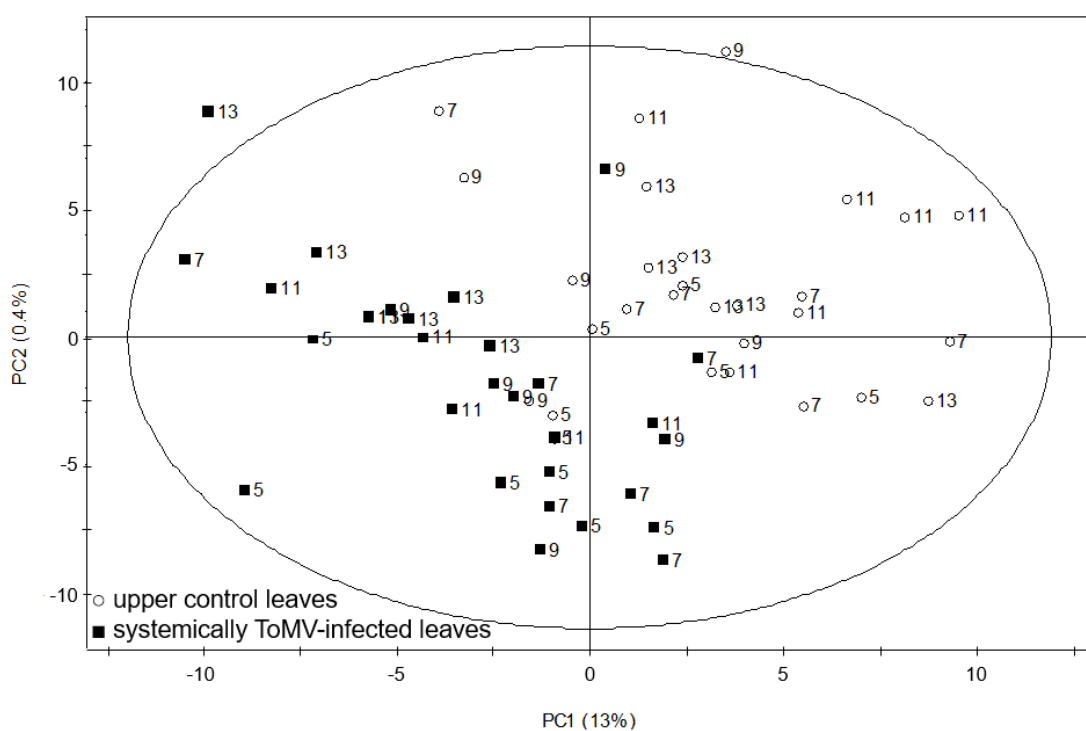
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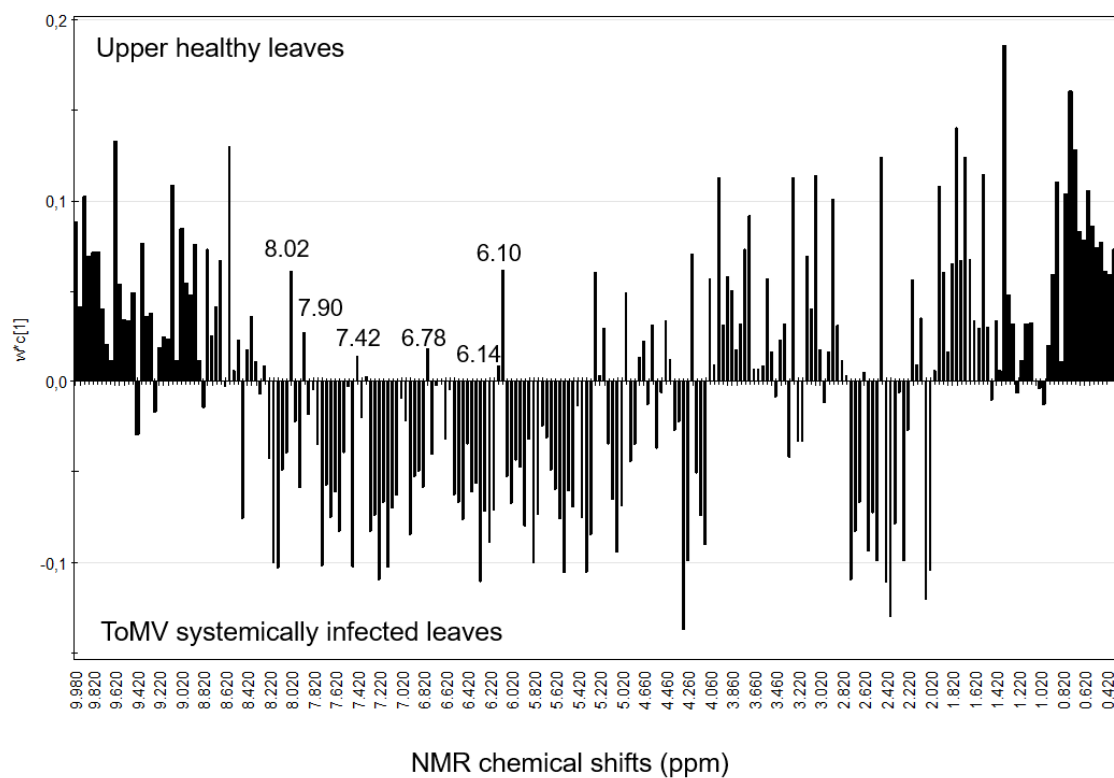
**Figure 5B.** PLS-DA score plot of delayed systemic viral infection on the  $^1\text{H}$  NMR signals in the range of  $\delta$  0.3-10.0. The numbers indicate the days after mock and ToMV inoculation.

(○) upper leaves of healthy tomato plants

710 (■) systemically ToMV-infected leaves of inoculated tomato plants.



**Figure 6.** Column loading plot of PLS-DA component 1 on upper healthy and ToMV systemically infected leaves.



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Table 1. Compounds identified from <sup>1</sup>H-NMR and 2D NMR spectra in 50% CH<sub>3</sub>OH-*d*<sub>4</sub> in D<sub>2</sub>O (KH<sub>2</sub>PO<sub>4</sub> buffer pH 6) of control and ToMV infected *S.lycopersicum* leaves.

720 (↑): increase, (↓) decrease, (=) no change, (L): local infection. (S): systemic infection

Metabolites	Chemical shift (δ in ppm) and coupling constant ( <i>J</i> in Hz)	Age (old)	Development stage (distal)	Infection		Collection time (19h)
				L	S	
Amino/organic acids						
Isoleucine	0.95 (t, 7.5), 1.02 (d, 7.0), 1.26 (m), 1.50 (m), 1.98 (m), 3.61 (d, 4.0)	↓	=	↓	=	↓
Leucine	0.97 (d, 6.4), 0.98 (d, 6.4), 1.72 (m)	↓	=	↓	=	↓
Valine	1.00 (d, 7.2), 1.05 (d, 7.2), 2.28 (m), 3.54 (d, 4.0)	↓	↑	↓	=	↓
Threonine	1.33 (d, 6.6), 3.53 (d, 5.0), 4.23 (m)	=	=	↓	↓	↓
Alanine	1.48 (d, 7.2), 3.73 (m)	↓	↑	↓	↓	↓
Arginine	1.67 (m), 1.72 (m), 3.24 (m), 3.67 (t, 5.8)	↓	↑	↓	↓	↓
GABA	1.90 (m), 2.31 (t, 7.2), 3.01 (t, 7.2)	=	↓	↑	↑	↓
Acetic acid	1.94 (s)	↑	↑	↓	↓	↓
Glutamic acid	2.05 (m), 2.13 (m), 2.39 (td, 7.2, 2.4), 3.70 (dd, 7.2, 4.3)	↓	↓	↑	↑	↑
Glutamine	2.14 (m), 2.45 (m), 3.71 (m)	↓	↓	↑	↓	↓
Malic acid	2.47 (dd, 15.8, 8.3), 2.72 (dd, 15.8, 3.9), 4.29 (dd, 8.3, 3.9)	↓	↓	↑	↑	↓
Citric acid	2.53 (d, 16.3), 2.71 (d, 16.3)	↓	↓	↑	↑	↓
Aspartic acid	2.65 (dd, 17.3, 8.8), 2.81 (dd, 17.3, 4.0), 3.84 (dd, 8.8, 4.0)	↓	↓	↑	↑	↓
Asparagine	2.81 (dd, 17.0, 8.2), 2.95 (dd, 17.0, 4.0), 3.94 (dd, 8.2, 4.0)	↑	↓	↑	↓	↓
Tryptophan	7.15 (td, 7.9, 2.0), 7.24 (td, 7.9, 2.0), 7.29 (s), 7.49 (d, 8.0), 7.71 (d, 7.9)	=	↓	↓	↑	↓
Fumaric acid	6.54 (s)	↓	↓	↑	↑	=
Phenylalanine	7.35 (m)	↓	↓	↑	↑	↓
Formic acid	8.46 (s)	↑	↓	↑	↑	↓
Sugars						

Sucrose	4.17 ( <i>d</i> , 8.6), 5.41 ( <i>d</i> , 3.8)	↑	↑	↓	↑	↑
Glucose ( $\alpha,\beta$ form)	5.18 ( <i>d</i> , 3.8), 4.58 ( <i>d</i> , 7.9)	↑	↑	↑	↓	↑
Phenylpropanoids/ Flavonoids						
Caffoyl esters of polyhydroxy compounds (glucaric acid) (4)	5.10 ( <i>d</i> , 4.9), 6.47 ( <i>d</i> , 16.0), 6.98 ( <i>d</i> , 8.2), 7.06 ( <i>dd</i> , 8.2, 2.0), 7.18 ( <i>d</i> , 2.0), 7.66 ( <i>d</i> , 16.0)	↓	↓	↓	↑	↑
Rutin	6.33 ( <i>d</i> , 2.1), 6.54 ( <i>d</i> , 2.1), 6.99 ( <i>d</i> , 8.4), 5.02 ( <i>d</i> , 7.8)	↓	↓	↑	↑	↑
Other compounds						
Dimethylamine	2.97 ( <i>s</i> )	↑	↑	↑	↑	↑
Ethanolamine	3.12 ( <i>t</i> , 5.5)	=	=	=	↓	↓
Choline	3.21 ( <i>s</i> )	↑	↑	↑	↑	↓
Trigonelline	4.46 ( <i>s</i> ), 8.10 ( <i>dd</i> , 1.8, 6.2), 8.87 ( <i>m</i> ), 9.14 ( <i>s</i> )	=	↑	=	↓	↓
UDP-Glucose	5.96 ( <i>d</i> , 8.1), 7.97 ( <i>d</i> , 8.1)	↑	=	↓	↑	↑
Adenosin	5.96 ( <i>d</i> , 4.4), 8.24 ( <i>s</i> ), 8.54 ( <i>s</i> )	↓	↓	↓	=	↓