

## Determination of the Dominant Variants of *Hop Stunt Viroid* in Two Different Cachexia Isolates from North and South of Iran

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### ABSTRACT

Citrus plants are hosts of several viroid species, among which, pathogenic variants of *Hop Stunt Viroid* (HSVd) induce citrus cachexia disease. Stunting, chlorosis, gumming of the bark, stem pitting and decline are symptoms of cachexia in mandarins and their hybrids as susceptible hosts. Based on the pathogenic properties on citrus, HSVd variants are divided in two distinct groups: those that are symptomless on sensitive citrus host species and those that induce cachexia disease. In this study, two cachexia isolates were selected and biological indexing was performed in a controlled temperature greenhouse (40°C day and 28°C night) using Etrog citron (*Citrus medica*) grafted on Rough lemon (*C. jambiri*), as a common indicator for citrus viroids. The plants were inoculated with the inocula from a severe symptomatic tree of a newly declining orchard of Jiroft, Kerman province and a mild symptomatic tree from Mazandaran province. Presence of HSVd was confirmed with sPAGE, Hybridization by DIG-labeled probes and RT-PCR using specific primers of HSVd. Primary and secondary structures of the isolates were studied. The consensus sequence of RT-PCR amplicons of the severe isolate (JX430796) presented 97% identity with the reference sequence of a IIb variant of HSVd (AF213501) and an Iranian isolate of the viroid (GQ923783) deposited in the gene bank. The mild isolate (JX430798) presented 100% homology with the HSVd-IIc variant previously reported from Iran (GQ923784). Both isolates were shown to be cachexia inducing according to their sizes, sequences and lack of “non-cachexia expression motif” structures.

**Keywords:** Biological indexing, Cachexia, Citrus, HSVd, Viroid.

### INTRODUCTION

Viroids are small molecules of single strand, covalently closed RNA and their genome size varies between 246 to 401 nucleotides. They belong to two families, the *Pospiviroidae* and the *Avsunviroidae* (King, *et al.* 2012). Citrus plants are natural hosts of several viroids, all belonging to the *Pospiviroidae* family: *Citrus Exocortis Viroid* (CEVd), *Hop Stunt Viroid* (HSVd), *Citrus Bent Leaf Viroid* (CBLVd), *Citrus Dwarfing Viroid* (CDVd), *Citrus Bark Cracking Viroid* (CBCVd), *Citrus Viroid V* (CVd-V) and *Citrus Viroid VI* (CVd-VI) (Eiras *et al.*,

2013). Among the important citrus diseases, CEVd and pathogenic variants of HSVd are the causal agents of exocortis and cachexia, respectively. They induce symptoms in susceptible host species whereas others remain symptomless. Stunting, chlorosis, gumming of the bark, stem pitting and decline are symptoms of cachexia that appear in mandarins (*Citrus reticulata*) and their hybrids as cachexia susceptible hosts. Severely affected trees are stunted and may even die (Duran-Vila *et al.*, 2000).

Cachexia was first described in 1950 as a disease of Orlando tangelo (Childs, 1950). *Hop Stunt Viroid* (HSVd) with a size of 295–

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303 nucleotides, the only member of the genus *Hostuvioid* within the family *Pospiviroidae* (Flores et al., 2005), was later identified as the causal agent of cachexia (Semancik et al., 1988). Five structural domains termed as central, variable, pathogenic, terminal left and right were characterized in the rod-like secondary structure of the viroids (Keese and Symons, 1985). There are three *HSVd* variants in citrus, variant IIa is non pathogenic, IIb and IIc variants, are pathogenic (Sano et al., 1988; Semancik et al., 1988; Levy and Hadidi, 1993). Five nucleotides in the cachexia-expression motif located in the variable domain, have been demonstrated to differentiate between pathogenic and non-pathogenic variants of *HSVd* (Palacio-Bielsa et al., 2004; Reanwarakorn and Semancik, 1998). Since the beginning of 2010, a widespread disease with decline symptoms appeared in citrus trees of Jiroft region in Kerman province. Several cases of declining *Minneola Tangelo* (*C. paradisi* X *C. reticulata*) trees with typical cachexia symptoms were noticed during surveys and the present research was conducted as a part of etiological studies (Banihashemian and Bani Hashemian, 2012).

## MATERIALS AND METHODS

### Plant Materials and HSVd Sources

Two *HSVd* isolates were collected from a severe (HH3) and a mild (HI3) cachexia

symptomatic *Minneola tangelo* tree from Kerman and Mazandaran provinces respectively (Figure 1). Biological indexing was performed using Etrog citron 861-S1 (*C. medica*) grafted on Rough lemon (*C. jambhiri*) rootstock, as the common indicator for citrus viroids. An Italian source of cachexia (HG3) containing three *HSVd* variants (kindly provided by Dr. K. Djelouah, IAMB, Italy) was used for inoculation of positive controls. Blocks of five seedlings were singly-inoculated with two graft patches from each viroid isolate and grown under greenhouse conditions (40°C day and 28°C night) for nine months (Banihashemian and Bani Hashemian, 2012). Non-inoculated plants were used as negative controls.

### RNA Extraction Methods

SDS-potassium acetate method (Bernard and Duran-Vila, 2006) was used as the RNA extraction method for Reverse Transcription-Polymerase Chain Reaction (RT-PCR). Briefly, tissue samples (500 mg of leaf and bark) were placed in sealed plastic bags in the presence of 5 ml of extraction buffer (0.1M Tris-HCl, pH 8.0; 50 mM EDTA; 0.5M NaCl; 10 mM mercaptoethanol) and were homogenized using a pestle. The homogenate was subjected to alkaline denaturation. Standard viroid extraction designed to yield high viroid titers (Bernard and Duran-Vila, 2006), was applied for other detection methods. Tissue samples (5 g of young leaves and



**Figure 1.** Severe symptoms of cachexia, including gumming and pitting of bark and wood (A) of declining *Minneola tangelo* orchard from Jiroft, Kerman Province (B) in comparison with mild symptoms of cachexia in the same variety from Mazandaran Province (C).

barks) from Etrog plants were homogenized in 20 ml of extraction medium containing 15 ml phenol and 5 ml buffer [0.4M Tris-HCl, pH 8.9; 1% (w/v) Sodium Dodecyl Sulfate (SDS); 5 mM EDTA, pH 7.0; 4% (v/v) mercaptoethanol]. The total nucleic acids were partitioned in 2M LiCl, and the soluble fraction was concentrated by ethanol precipitation and resuspended in TKM buffer [10 mM Tris-HCl; pH 7.4; 10 mM KCl, 0.1 mM MgCl<sub>2</sub>]. Standard extracts from citrons infected *HSVd*, *CEVd*, *CDVd*, *CBCVd* and *HSVd-IIa* variants (kindly provided by Dr. N. Duran-Vila, IVIA, Spain) were used for hybridization and electrophoresis.

#### Sequential Polyacrylamide Gel Electrophoresis (sPAGE)

Twenty µl of the nucleic acid preparations from standard extraction method were subjected to two consecutive rounds of polyacrylamide gel electrophoresis in 5% gels using a vertical electrophoresis system, first gel under non-denaturing and the second under denaturing conditions (Rivera-Bustamante *et al.*, 1986). The circular forms of the viroids were viewed by silver staining (Igloi, 1983).

#### Northern Blot Hybridization

The RNAs separated by sPAGE were electroblotted (313 mA for 2 hours) in a transfer system to positively charged nylon membranes (Roche Applied Science) using TBE buffer (90 mM Tris, 90 mM boric acid, and 2 mM EDTA, pH 8.0) and immobilized 2 hours at 80°C in oven. Prehybridization (60°C for 2h) and hybridization (50°C overnight) were performed (Murcia *et al.*, 2009) using digoxigenin (DIG)-labeled viroid-specific probes of *HSVd*, *CEVd*, *CBCVd* and *CDVd* generated by PCR from plasmids containing the full-length viroid sequence kindly provided by Dr. N. Duran-

Vila. The reaction was detected using a Dig-detection kit (Roche Applied Science).

#### RT-PCR Protocol and Viroid Characterization

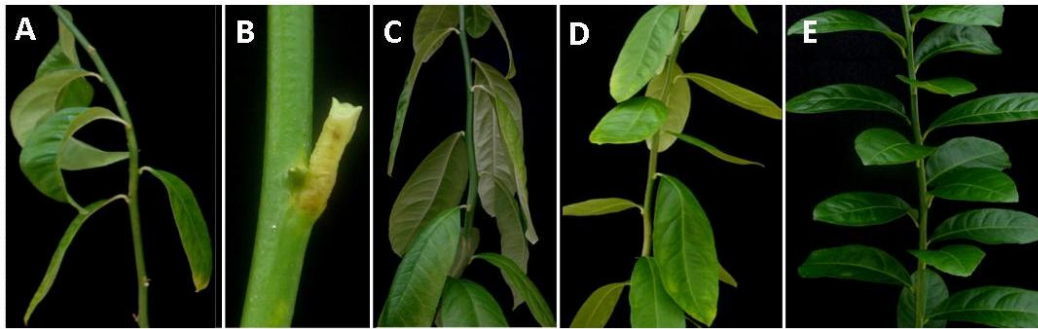
RT-PCR was performed in two steps using Revert Aid Kit (Fermentas) and *HSVd* specific primers HSVd-RT (5'GTGTTGCCCGGGGCTCCTTTCTCTGG-3'), HSVd-F1 (5'GGGGCAACTCTTCTCAGAATCC-3') and HSVd-R1 (5'-GGGGCTCCTTTCTCAGGTAAGTC-3') (Bernard and Duran-Vila, 2006). The viroid template obtained from SDS potassium acetate extracts and the HSVd-RT primer were denatured at 95°C for 5 minutes. The reaction mixture (20 µl final volume) was incubated at 42°C for 1 hour. PCR amplification was performed with the *HSVd* specific primers (HSVd-F1 and HSVd-R1) using a PCR Master Mix (Fermentas).

#### Sequence Analysis and Determination of Secondary Structures

The RT-PCR products of *HSVd* isolates were sequenced and compared with other *HSVd* sequences from the gene bank. Multiple alignments of *HSVd* sequences were obtained using Clustal W (Thompson *et al.*, 1994). The most stable secondary structure was obtained with the RNA structure software (version 4.6).

## RESULTS AND DISCUSSION

Symptom expression of typical epinasty (Figure 2-A) and petiole browning (Figure 2-B) were observed in citrons inoculated with the positive control, HG3, after nine months. A mild leaf epinasty (Figures 2-C and -D) was noticed in the plants inoculated with two cachexia isolates HH3 and HI3 (Banihashemian and Bani Hashemian, 2012). Biological indexing has been

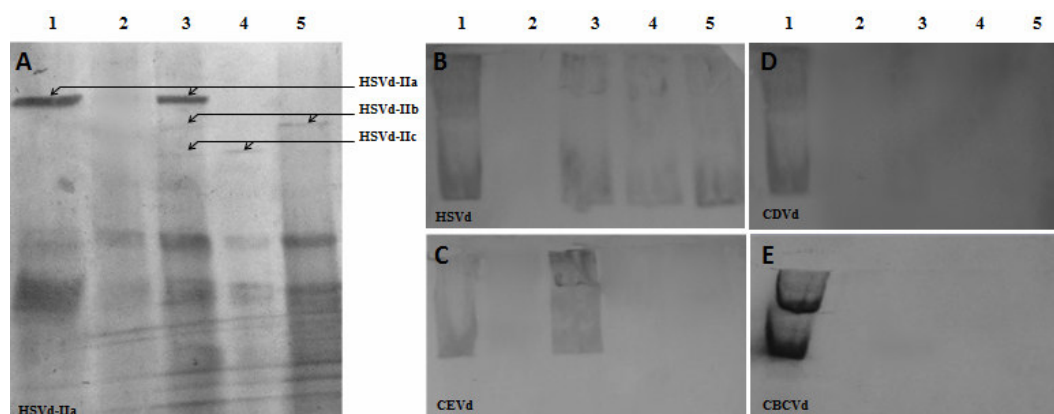


**Figure 2.** Symptoms of Etrog citron indicator plant inoculated with HG3 (A and B), HH3 (C), HI3 (D) isolates compared with the healthy control plant (E).

performed for many years for detection of citrus viroids. However it is used as an important step in detection of viroid infections, but is not always practical because the technique needs greenhouse facilities (Banihashemian *et al.*, 2012). Parsons Special Mandarin, the specific cachexia indicator requires a long incubation period under special environmental conditions for symptom development (Pina *et al.*, 1991) and for this reason it was not applied in this study. Etrog citron is the common indicator plant for detection of citrus viroids (Roistacher, 1991). Although citrus viroids other than *CEVd* do not produce specific symptoms (Roistacher, 1991) but it is clear that all citrus viroids can well multiply on Etrog Citron (Bani Hashemian *et al.*, 2015). Therefore a

combination of biological indexing using Etrog citron and molecular methods (Duran-Vila *et al.*, 1988 and 1993) was used for detection of citrus viroids and *HSVd* characterization of the study. Epinasty, a symptom related to *CEVd* (Roistacher, 1991), were observed in the Etrog plants inoculated with HG3. The presence of *CEVd* in this isolate was confirmed by hybridization (Figure 3-C).

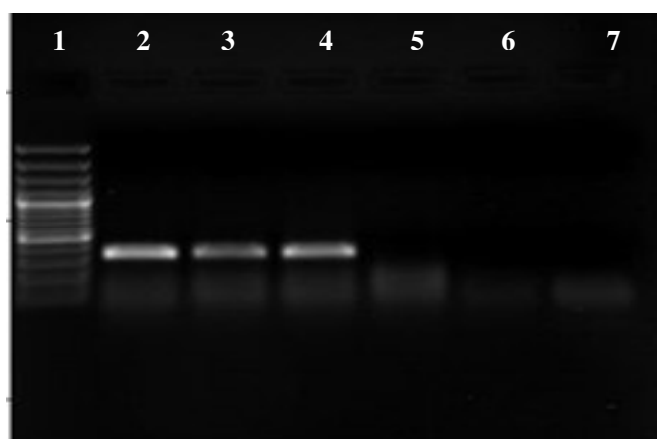
sPAGE analysis of the citron plants inoculated with HG3 demonstrated that the source contained several viroids (data non shown). Presence of *CEVd* and *HSVd* but not *CDVd* and *CBCVd* was confirmed by hybridization in the same isolate (Figures 3-B, -C, -D and -E). Because of similar sequence of variable and terminal left domains of *CEVd* and *CBCVd* (Puchta *et al.*,



**Figure 3.** Stained polyacrylamide gel (A) and hybridization with viroid specific probes of *HSVd* (B), *CEVd* (C), *CDVd* (D) and *CBCVd* (E) containing extracts of positive control (Lane 1), viroid free sample (Lane 2), HG3 (Lane 3), HI3 (Lane 4) and HH3 (Lane 5). Extracts from citrons infected with *HSVd-IIa*, *HSVd*, *CEVd*, *CDVd* and *CBCVd* were used respectively as positive controls of electrophoresis and hybridization.

1991), two bands in hybridization analysis with specific probe of *CBCVd* were seen (Figure 3-E). It shows that the positive extract contains *CEVd* in addition to *CBCVd*. Electrophoresis also revealed presence of three bands related to distinct *HSVd* variants (Figure 3-A). The highest intensive band with the equal mobility of the positive extract of IIa is related to the presence of *HSVd-IIa* in HG3 as a dominant variant. According to negative reaction to *CDVd* probe, two lower bands in electrophoresis of HG3 should be associated with infection to *HSVd-IIb* and *IIC* variants (Duran-vila *et al.*, 1988; 1993). Due to the characteristic mobility of three *HSVd* variants in HG3, the single RNA band found in electrophoresis of HH3 and HI3 (Figure 3-A), could be respectively related to *HSVd-IIb* and *HSVd-IIC* variants. The nucleic acid preparations from Etrog citron inoculated with HG3, HH3 or HI3 had a positive reaction in RT-PCR with *HSVd* specific primers. PCR amplification produced amplicons of ~300bp and no fragments were detected from healthy control samples (Figure 4). Extracts from citrons infected with *HSVd-IIa*, *HSVd*, *CEVd*, *CDVd* and *CBCVd* were used respectively as positive controls of electrophoresis and hybridization.

Cachexia is an economically important disease that is widespread in great parts of citrus growing areas of the world (Duran-Vila *et al.*, 2000). *HSVd* had been identified as a viroid with a broad host range and two distinct groups of variants in citrus (Loconsole *et al.*, 2013). Based on their pathogenic properties on citrus, variants that induce citrus cachexia disease in the sensitive hosts were shown to be pathogenic (IIb and IIC variants) while those that did not induce symptoms in the same hosts were named as non-pathogenic (IIa variant) (Palacio-Bielsa *et al.*, 2004). Cachexia was first reported from Kerman and Mazandaran provinces of Iran (Habashi and Rahimian, 1984). Two *HSVd* isolates of this study were selected from the mentioned provinces. The severe cachexia isolate (HH3) with typical symptoms of the disease including stunting, chlorosis, gumming of the bark and stem pitting obtained from declining tangelo trees of Kerman province were compared with a mild symptomatic isolate (HI3) from the same variety of Mazandaran province. Since the elevated temperatures favor viroid replication, it has been accepted that warm temperature is very important for maximum expression of cachexia symptoms (Semancik *et al.*, 1988). Hence the indicator plants under index for cachexia and other citrus



**Figure 4.** Electrophoresis of RT-PCR products of *HSVd* in 1% agaros gel from Etrog citron indicator plant. Lane 1: 100 bp Ladder; Lane 2: HG3; Lane 3: HH3; Lane 4: HI3, Lane 5: Negative control (-) included RT-PCR analysis of viroid free samples; Lane 6: RT control without RNA template, and Lane 7: PCR control without cDNA template.







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## تعیین واریانت غالب ویروئید کوتولگی رازک در دو جدایه مختلف کاکسیا از شمال و جنوب ایران

س. ن. بنی هاشمیان، س. م. بنی هاشمیان، و س. م. اشکان

### چکیده

مرکبات میزبان چندین گونه ویروئید است که از آن میان، بیماری کاکسیای مرکبات به وسیله ویروئید های بیماری زای ویروئید کوتولگی رازک (*Hop stunt viroid, HSVd*) ایجاد می شود. کوتولگی، زردی، صمغ زیر پوست، ساقه آبله ای و زوال، علائم کاکسیا هستند که در ارقام نارنگی و دورگ های آن به عنوان میزبان های حساس ظاهر می گردد. بر اساس خصوصیات بیماری زایی در مرکبات، دو گروه مجزا از واریانت های HSVd وجود دارد. آنهایی که در ارقام حساس مرکبات بدون علائم هستند و گروهی که بیماری کاکسیا ایجاد می کنند. در این مطالعه، دو جدایه کاکسیا انتخاب شد و نموده سازی بیولوژیکی در گلخانه تحت شرایط کنترل شده دمایی (روز ۴۰ درجه سانتی-گراد و شب ۲۸ درجه سانتی-گراد) با استفاده از گیاه بالنگ اترانگ (*C. medica*) پیوند شده روی پایه



رافلمون (*C. jambhiri*)، به عنوان گیاه محک عمومی ویروئیدهای مرکبات، انجام شد. گیاهان با منبع آلودگی از درختی با علائم شدید از باغ‌های در حال زوال منطقه جیرفت استان کرمان و درختی با علائم خفیف از استان مازندران، مایه زنی شدند. حضور HSVd با سه روش مولکولی sPAGE، هیبریداسیون با پروب‌های نشاندار شده و RT-PCR با استفاده از پرایمرهای اختصاصی HSVd تأیید گردید. ساختارهای اولیه و ثانویه جدایه‌ها بررسی شد. توالی فرآورده نهایی RT-PCR از جدایه شدید (JX430796)، با توالی مرجع واریانت HSVd-IIb (AF213501) و یک جدایه ایرانی از ویروئید (GQ923783) گزارش شده از بانک ژن، ۹۷ درصد تشابه نشان داد. جدایه خفیف (JX430798) با واریانت HSVd-IIc که قبلاً از ایران گزارش گردید (GQ923784) شباهت کامل داشت. بر اساس اندازه، توالی و فقدان ساختار مشخصه واریانت غیر کاککسیا، هر دو جدایه مورد بررسی به عنوان واریانت‌های مولد کاککسیا تشخیص داده شدند.