ORIGINAL ARTICLE

Genetic diversity of the movement and coat protein genes of South American isolates of *Prunus* necrotic ringspot virus

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Abstract *Prunus* necrotic ringspot virus (PNRSV) is distributed worldwide, but no molecular data have been previously reported from South American isolates. The nucleotide sequences corresponding to the movement (MP) and coat (CP) proteins of 23 isolates of PNRSV from Chile, Brazil, and Uruguay, and from different *Prunus* species, have been obtained. Phylogenetic analysis performed with full-length MP and CP sequences from all the PNRSV isolates confirmed the clustering of the isolates into the previously reported PV32-I, PV96-II and PE5-III phylogroups. No association was found between specific sequences and host, geographic origin or symptomatology. Comparative analysis showed that both MP and CP have phylogroup-specific amino acids and all of the motifs previously characterized for both proteins. The study of the distribution of synonymous and nonsynonymous changes along both open reading frames revealed that most amino acid sites are under the effect of negative purifying selection.

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Introduction

The South American production of stone fruit trees represents 6.3% of the world fruit production (FAO 2005). Stone fruit trees are affected by a large number of viruses, which cause significant economic losses [38]. Among these many viruses, the most economically important ilarvirus is *Prunus* necrotic ringspot virus (PNRSV). PNRSV has a worldwide distribution and affects *Prunus* spp. including commercial peach, nectarine, cherry, apricot, almond and plum trees [19, 37]. The typical symptoms induced by PNRSV are chlorosis, necrosis, leaf deformation, and growth defects. Chlorosis usually forms rings, bands, lines, mottles, mosaics and arabesques. Ring necrosis mainly affects leaves but also appears on branches and emerging buds in the case of aggressive strains. The foliar rings, initially chlorotic, become necrotic and drop off, resulting in perforations of the leaf [13, 37, 47]. PNRSV is transmitted by pollen, seed and routine vegetative propagation methods [6, 12, 20, 21, 30].

PNRSV belongs to the genus *Ilarvirus* of the family *Bromoviridae*, and its genome is organized into three plus-strand RNAs. RNAs 1 and 2 encode for proteins involved in viral replication, whereas RNA 3 encodes a 5′-proximal movement protein (MP) and a 3′-proximal coat protein (CP), which is expressed via a subgenomic RNA 4 [9]. Many different isolates of PNRSV have been characterized and phylogentically grouped into three subgroups named PV32-I, PV96-II and PE5-III [3, 4, 13, 22, 46, 48]. Analysis of the different isolates revealed a lack of association between specific nucleotide or amino acids patterns and the host origin, symptomatology or geographic area. Nonetheless, the majority of isolates from phylogroups PV96-II and PV32-I tend to exhibit latent/mild or chlorotic/necrotic symptoms, respectively [4, 22, 23]. Molecular analyses of

the coding sequences of CP and MP revealed that both proteins are highly conserved despite the host and geographical origin of the isolates being compared [3]. The PNRSV MP has been assigned to the 30 K superfamily, a group of twenty genera the MPs of which are related to the MP of tobacco mosaic virus (TMV) [36]. Analyses performed with the MP of PNRSV and the closely related alfalfa mosaic virus (AMV) have allowed characterization of an RNA-binding domain upstream of the transmembrane motif [26, 27, 45], the region responsible for tubule formation [41] and the C-terminal portion of the protein that specifically interacts with the cognate CP [40]. Two different motifs have been characterized for the CP genes of both PNRSV and AMV: an RNA-binding domain at the N-terminus that is necessary for different viral processes (e.g. genome activation, virus replication and translation) [2, 7, 9], and the C-terminal region responsible for dimer formation [5, 10].

PNRSV has been detected in many South American countries and in different host species. The prevalence of the virus in stone fruit trees in the central zone of Chile is 9.7% [28], while it reaches up to 25% in temperate zones in Argentina [14]. In Brazil, PNRSV has been detected in Rio Grande do Sul with a prevalence of 17% [16], while in Uruguay its prevalence is even higher, 48% [35]. In spite of the economic importance of stone fruit crops in South America, the molecular characterization of PNRSV isolates is scarce, except for one Argentinean isolate [24]. In the present work, the MP and CP genes of 23 PNRSV isolates from Chile, Brazil and Uruguay have been characterized and analyzed.

Materials and methods

Plant material and viral RNA extraction

Table 1 shows the different PNRSV isolates included in this study, with specific reference to the source tree variety and the symptoms they presented. Total nucleic acid

Table 1 Listing and properties of PNRSV isolates

Isolate	Source	Origin	Origin of the variety	Symptomatology	Group	Sequence code	Accession number CP	Accession number MP
453	Almond cv. Carmel	Chile	USA	Chlorotic pattern	PV96-II	AlmCl.car1	EF565247	EF565224
553	Cherry cv. Corazón de Paloma	Chile	Unknown	Asymptomatic	PV32-I	ChrCl.cor1	EF565248	EF565225
564	Cherry cv. Bing	Chile	USA	Chlorotic pattern	PV96-II	ChrCl.bin1	EF565249	EF565226
585	Cherry cv. Sweet Hard	Chile	Canada	Asymptomatic	PV32-I	ChrCl.swe1	EF565250	EF565227
775	Cherry cv. Royal Down	Chile	USA	Ugly fruit	PV96-II	ChrCl.roy1	EF565251	EF565228
113	Nectarine cv. Early John	Chile	USA	Asymptomatic	PV96-II	NctCl.ear1	EF565252	EF565229
73	Nectarine cv. August Fire	Chile	USA	Asymptomatic	PE5-III	NctCl.aug1	EF565253	EF565230
26	Peach cv. Pomona	Chile	USA	Chlorotic pattern	PV96-II	PchCl.pom1	EF565254	EF565231
437	Peach cv. Loadell	Chile	USA	Chlorotic pattern	PV96-II	PchCl.loa1	EF565255	EF565232
435	Peach cv. Loadell	Chile	USA	Chlorotic pattern	PV96-II	PchCl.loa2	EF565256	EF565233
487	Peach cv. Rich Lady	Chile	USA	Chlorotic pattern	PV96-II	PchCl.ric1	EF565257	EF565234
518	Peach cv. August Lady	Chile	USA	Chlorotic pattern	PV96-II	PchCl.aug1	EF565258	EF565235
526	Peach cv. Summer Lady	Chile	USA	Chlorotic pattern	PV32-I	PchCl.sum1	EF565259	EF565236
532	Plum cv. Mirabolan	Chile	USA	Arabesques	PV96-II	PlmCl.mrb1	EF565260	EF565237
474	Plum cv. Black Ambar	Chile	USA	Chlorotic pattern	PV32-I	PlmCl.bla1	EF565261	EF565238
478	Plum cv. Friar	Chile	USA	Chlorotic pattern	PV96-II	PlmCl.fri1	EF565262	EF565239
444	Plum cv. D'Agen	Chile	France	Chlorotic pattern	PV96-II	PlmCl.dag1	EF565263	EF565240
MS3	Peach cv. Marli	Brazil	Brazil	Asymptomatic	PV32-I	PchBr.unk1	EF565264	EF565241
MS4	Peach cv. Marli	Brazil	Brazil	Asymptomatic	PV32-I	PchBr.unk2	EF565265	EF565242
M1	Peach cv. Marli	Brazil	Brazil	Asymptomatic	PV96-II	PchBr.unk3	EF565266	EF565243
15	Plum cv. Golden Japan	Uruguay	Unknown	Not Available	PV96-II	PlmUy.gol1	EF565267	EF565244
19	Peach cv. Early Grande	Uruguay	Unknown	Not Available	PV32-I	PchUy.ear1	EF565268	EF565245
36	Peach cv. June Gold	Uruguay	Unknown	Chlorotic pattern	PV32-I	PchUy.jun1	EF565269	EF565246
1	Peach cv. Criollo	Argentina	Unknown	Not Available	PV32-I	PchAr.cri1	AY007217	
PV96						PV96	S78312	
PV32						PV32	Y07568	
PE5						PE5	L38823	

extraction was performed from 0.2 g of leaves using the silica capture extraction protocol [34]. The extracted nucleic acids were stored at −80°C until use.

Amplification and cloning of genome sequences

Primers flanking the MP and CP genes were designed from the sequence available in the database [42] (accession number Y07568; Table 1). The reverse transcription reaction was done in 20 μl containing 1 μl of 10% PVP, 1 mM of each dNTP, 40 U of ribonuclease inhibitor (Amersham International, Cleveland), 10 U of RevertAid™ Minus M-MuLV RT (MBI Fermentas), 100 pmol of each antisense primer (see Table 2) and 1 μg of total nucleic acids extracted from infected tissue. PCR reactions were done in 50 μl containing 1 μl of 10% PVP, 3 μl of the reverse transcription reaction, 0.5 mM of each dNTP, 5 μl of 10× buffer (166 mM $(NH_4)_2SO_4$, 670 mM Tris–HCl at pH 8.8 and Tween 20), 1 U of EcoTaq DNA polymerase (Ecogen SRL, London), and 100 pmol of each primer (Table 2). PCR reaction was performed in a Perkin Elmer 2400 thermal cycler with an incubation at 94°C for 2 min followed by thirty cycles at 94°C for 30 s, 52°C for 30 s and 72°C for 30 s and a final incubation at 72°C for 10 min. Amplified products were electrophoresed in 1% agarose gels in TAE (40 mM Tris–acetate, 1 mM EDTA, pH 8.0), stained with ethidium bromide and extracted with the QIAGEN Tip-20 (QIAGEN Inc., USA). The PCR fragments were directly sequenced by the dideoxynucleotide chain-termination method in an automated sequencer (ABI 3100 Genetic Analyzer; Perkin Elmer Applied Biosystem) using the primers employed for the amplification of both genes. In the case of the MP gene, two additional internal primers were used to sequence the full gene (VP239 and VP951; Table 2).

Sequence alignments and phylogenetic analyses

Amino acid alignments for MP and CP were obtained using MUSCLE version 3.52 [15]. Nucleotide sequences were then aligned by concatenating triplets according to the amino acid sequence alignment using the DAMBE version 4.2.13 [49]. The phylogenetic analysis of the MP and CP genes were inferred in a multi-step process: in the first step, the model of amino acid substitution that maximized the likelihood of the observations was inferred using the PROTTEST version 1.0.6 [1]. The JTT empirical matrix [29] was chosen as the model of amino acid substitution that better explained the observed pattern of genetic variability for both proteins. In a second step, neighbor-joining phylogenetic trees, using the JTT model, were obtained using the MEGA version 3.1 [32]. Third, the statistical reliability of the constructed trees was assessed by the bootstrap method based on 1,000 pseudoreplicates. The homologous sequences of apple mosaic virus (ApMV) were used for rooting purposes (accession number U15608). Finally, to further explore the reliability of certain branches in the trees, the four-cluster likelihood-mapping technique [44] was employed, as implemented in TREEPUZZLE version 5.2, with 10,000 puzzling steps.

The difference between nonsynonymous (d_N) and synonymous (d_S) substitution rates ($d_N − d_S$) was employed as a proxy to seek the sign and intensity of the natural selection acting on different amino acids sites. This difference was estimated for each position in the alignments using the random-effects likelihood (REL) method [39] implemented in the HYPHY server (http://www.datamonkey.org). Standard errors were computed by the bootstrap method (based on 1,000 pseudoreplicates). A value >0 is taken as evidence for positive or directional selection, whereas values <0 are a signature for negative or purifying selection.

Transmembrane motifs were predicted by the online PREDICTPROTEIN software (http://www.cubic.bioc. columbia.edu/predictprotein/).

Results and discussion

PNRSV is distributed worldwide and is probably one of the plant viruses for which the most different isolates have been characterized, principally from the USA, Europe and the Mediterranean area. However, data for PNRSV isolates

Table 2 List of primers used for PCR amplification and DNA sequencing

Underlining indicates extra nucleotides designed to introduce specific restriction sites

[a] Antisense primer

[b] Sense primer

Primer	Sequence	Location in the viral RNA 3 (nt)	Target gene
MPf[s]	5′-TTTGGATCCGATTGTTGGTTGTCTTTTC	124–142	MP
MPr[a]	5′-TTTAAGCTTATTGCAAATTCGGCAAAAC	1,105–1,123	MP
VP81[b]	5′-AGTGGATCCATGGTTTGCCGAATTTGC	1,103–1,120	CP
VP103[a]	5′-ACATAAGCTTCTAGATCTCAAGCAGGTC	1,765–1,783	CP
VP239[b]	5′-GGCCGTGTATTCCTCGTTTATGTA	438–461	MP
VP951[a]	5′-CACAGCTAGCCGGGAATAAAATAGGATTC	731–749	MP

from South America are scarce. In the present study, the molecular variability of the movement (MP) and coat (CP) protein genes from 23 isolates from Chile, Uruguay, and Brazil has been analyzed.

The deduced amino acids sequences of the MP and CP genes ranged between 283–284 and 224–226 residues, respectively. The average amino acid identities among South American isolates had percentage values of 91.5–100 and 89.7–100, respectively. The alignments for the two genes (Fig. 1) confirm that South American isolates belong to the same three phylogroups of the PNRSV isolates previously described PV32-I, PV96-II and PE5-III [3, 4, 22, 48].

The distribution into phylogroups of the 45 previously characterized PNRSV isolates from North America, Europe and the Mediterranean area was 57.8, 31.1, and 11.1% for PV96-II, PV32-I, and PE5-III, respectively [4]. No significant differences were detected between the frequency distribution of phylogroups among the 24 South American isolates (58.3, 37.5 and 4.2%; $\chi^2 = 3.805$, 2 d.f., $P = 0.149$) and those from the rest of the world. Therefore, it is statistically justified to pool all samples and to compute an overall expected distribution for the members of each phylogroup (58.0% PV96-II, 33.3% PV32-I and 8.7% PE5-III) that should be valid irrespective of the geographic area and host species sampled. Furthermore, this frequency distribution of phylogroups may represent an equilibrium distribution in which the abundance of a given type depends on factors such as its fitness, growth, and transmission rates under different environmental conditions and in different host species. However, it is worth noting the low number of PNRSV isolates assigned to the PE5-III phylogroup (6 out of 69) and the previous observation that the PE5-III phylogroup has an accelerated rate of molecular evolution for the MP and CP genes [11]. Collectively, these observations suggest that selective pressures acting on PE5-III isolates are stronger than on other phylogroups. Which specific pressures act on PE5-III, and particularly, whether its actual frequency is stable or transiently changing toward higher levels, which may suggest a faster rate of molecular evolution, is still a question to be explored.

Functional relevance of the observed variability in MP

In the case of the MP, the intra-group amino acid identities ranged between 96.5–100, 96.8–98.9 and 98.9% among isolates belonging to the PV96-II, PV32-I, and PE5-III phylogroups, respectively, which are in the same range as the previously reported values [4]. The alignment of MP sequences (Fig. 1a) shows a highly conserved protein, except for the C-terminus, which contains the majority of the differences among all isolates. This portion of the

Fig. 1 Amino acid alignment of the movement (*MP*) (**a**) and coat ▶ (*CP*) proteins (**b**) of the 23 South American PNRSV isolates characterized in this work. In *bold*, isolates previously characterized and used to represent the three phylogroups (PV32-I, PV96-II and PE5-III; see Table 1). *Dots* indicate identical residues, whereas *dashes* denote gaps in the sequence. The *dash* at position 258 of the MP is included to maintain the same numbering when all PNRSV isolates sequenced so far are included. The basic region carrying critical residues for the RNA binding of the MP [26] is *underlined* with the basic residues in *bold*. The C-terminal region that aligned with the C-terminal 44 amino acids of AMV MP, which specifically interact with the cognate CP [40], are *doubly underlined*. The basic residues critical for the RNA-binding properties of the CP [7] are *highlighted in bold*. The C-terminal region required for dimer formation is *underlined* [5]. The amino acid sequences of the PNRSV isolates PV96, PV32 and PE5 are included. *Numbers at the top* represent the corresponding residues in the full-length protein

protein has been postulated by covariation analysis to interact with the N-terminus of the cognate CP, in which E256 of the MP could interact with T52 of the CP [11]. Direct evidence of such interaction has been reported for the closely related AMV, in which the C-terminal 44 amino acids of the MP interact with the cognate CP [40]. Interestingly, the C-terminal 44 amino acids of the AMV MP aligned with the variable C-terminal 35 amino acids of the PNRSV MP (Fig. 1a). Most viral MPs have an RNA-binding domain, required for the cell-to-cell movement, that is characterized by an accumulation of basic residues. This domain has been mapped between residues 56–88 of the PNRSV MP, which carries nine basic residues [26]. Mutational analysis of the different basic residues showed that the substitution of three out of nine amino acids is sufficient to reduce the RNA binding affinity of the MP and also to block cell-to-cell transport without affecting other viral properties [27]. Interestingly, the nine basic residues of the RNA-binding domains are conserved in all PNRSV isolates sequenced so far except for isolate NecSp.mur [4], which contains a shorter version of only eight residues. This domain is also highly conserved among all isolates presented herein, including the nine basic residues. Furthermore, all isolates belonging to the PV32-I and PE5-III phylogroups contain an extra basic residue (K81). This peculiarity is a general rule for all isolates belonging to the two groups except for the PV32 isolate. The observation that isolate PchBr.unk1 contains two extra basic residues (R75, K81) allows it to be argued that it is possible that a functional RNA-binding domain requires from eight to eleven basic residues. This observation may reflect a range of biological RNA binding affinities in which extreme RNA interactions (strong or low) are incompatible with other processes during the viral life cycle (e.g., replication, encapsidation, etc.). In this sense, the dissociation constant (K_d) for the PNRSV MP–RNA complex was estimated to be 1.4 μM [26], which is within the range of the values

A

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              30        40        50        60        70        80       240       250       260       270       280
PV96      : LHKLMLSDEMKALPTKGCHILHLVNLPKSNVLRLASKEQKGFLLRQADKVKNKIYRC // VLTIQPKAPIVEEVKDED-EPTGSNG-ENHMEEKTVTVKVGSSGSA
PlmCl.fri1 : ...................................S.............S........ // .....................-.........-.................
PchCl.ric1 : ...................................S.............S........ // .....................-.........-.................
NctCl.ear1 : ...................................S.............S........ // .................N..-.......-G....................
PlmCl.dag1 : ...................................S.............S........ // .................N..-.............................
AlmCl.car1 : ...................................S.............S........ // .............REK.-..I....-........................
PlmCl.mrb1 : ...................................S.............S........ // ...............D.V..-.........-...................
PchBr.unk3 : ...................................S.............S........ // .................N..-..I...-......................
PlmUy.gol1 : ...................................S.............S........ // ...................-..I...-......................
ChrCl.roy1 : ..........R........................S.............S........ // .................N..-..I...-......................
ChrCl.bin1 : ...................................S.............S........ // .................N..-..I...-......................
PchCl.pom1 : ...................................S.............S........ // .................N..-..I...-......................
PchCl.aug1 : ...................................S.............S........ // ...................-..I...-......................
PchCl.loa1 : ..........R........................S.............S........ // ..........V.......I..V....-.......................
PchCl.loa2 : ..........R........................S.............S........ // ..........V.......I..V....-.......................

PV32      : ...................................I.............S........ // ..............I..DV-..L....-......................
ChrCl.cor1 : ..........R........................I.............S....K.... // ....R.........I..DV-..L....-......................
ChrCl.swe1 : ...................................I.............S........ // ..............I..DV-..L...........................
PchBr.unk1 : ..........R........................I...........S.R....K.... // ..........T...I..DV-..L....-......................
PchBr.unk2 : ..........R........................I.............S....K.... // ..........T...I..DV-..L....-......................
PlmCl.bla1 : ...................................I.............S....K.... // .............DV-..L...S-..........................
PchCl.sum1 : ...................................I.............S....K.... // ..............I..DV-..L....-......................
PchUy.ear1 : ..........R........................I.............S....K.... // ..............I..DV-..L....-......................
PchUy.jun1 : ..........R........................I.............S....K.... // ..............I..DV-..L...........................

PE5       : ..N.......RS.............................S.......K..... // .......T..I..L.NDE-..IS.SST.K.....V.A.E..RT...
NctCl.aug1 : ..N.......RS.............................S.......K..... // .......T..I..L.NDE-..I..SST.K.....V.A.E..RT...
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B

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              20        30        40        50        60        70        80       110       120       130       140       150
PV96      : CHPNGALVPLRAQQRAANNPNR--NPNRASSGTGPAVRPQPVVKTTWTVRGPNVPPRIPKGFV // MGQNLTLLTVIVRMNSMSSNGWIGMVEDYKVERPDGPNALS
PlmCl.fri1 : ......................--........V............................... // ..................................Q......
PchCl.ric1 : ......................--........V............................... // ..................................Q......
NctCl.ear1 : ......................--........V............................... // ..................................Q......
PlmCl.dag1 : ......................--........V............................... // ..................................Q......
AlmCl.car1 : ...........H..........--........V............................... // ..................................Q......
PlmCl.mrb1 : ............V.....--..I.........V............................... // ................................Q.N......
PchBr.unk3 : ......I...............--........V............................... // ................................Q.N......
PlmUy.gol1 : ............V.........--........V............................... // ..................................Q......
ChrCl.roy1 : ......................--........V............................... // ..................................Q......
ChrCl.bin1 : ......................--........V............................... // .........................................
PchCl.pom1 : ............V.........--........V............................... // .........................................
PchCl.aug1 : ............V.........--........V............................... // ..................................A......
PchCl.loa1 : ....N.................--.....V...I..V........................... // ..................................DQ.....
PchCl.loa2 : ....N.................--.....V...I..V........................... // ..................................DQ.....

PV32      : ....D...............S.SR...V...V...I.....................Y. // .........M........................DQ.....
PchAr.cri1 : ....D...............NR....V...L...I..............M.Y. // ..................................DQ.....
ChrCl.cor1 : ....D...............NR....V...TI..................Y. // ..................................DQ.....
ChrCl.swe1 : ....D...............GNR...V..S...TI.S.............Y. // ..................................DQ.....
PchBr.unk1 : ....D...............NR....V...I.................A..... // ..................................DQ.....
PchBr.unk2 : ....D...............NR....V...I.................A..... // ..................................DQ.....
PlmCl.bla1 : ....D...............NR....V...I..................Y. // ..................................DQ.....
PchCl.sum1 : ....D...............NR....V...I..................Y. // ..................................DQ.....
PchUy.ear1 : ....D...............NR....V...I..................Y. // ..................................DQ.....
PchUy.jun1 : ....D...............NR....V...I..................Y. // ..................................DQ.....

PE5       : ....D...............--..V.V.NVV......K.............. // .........I.N............I....R.NN.R.....
NctCl.aug1 : ....D...............--..V.V.NVV..V...KS............. // .........I.N............I....R.NN.E.....
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              180       190       200       210       220
PV96      : // THRVVIEFKTEVPAGAKVLVRDLYVVVSDLPRVQIPTDVLLVDEDLLEI
PlmCl.fri1 : // .................................................
PchCl.ric1 : // .................................................
NctCl.ear1 : // .................................................
PlmCl.dag1 : // .................................................
AlmCl.car1 : // .................................................
PlmCl.mrb1 : // .....................N..Q........................
PchBr.unk3 : // .................................................
PlmUy.gol1 : // .................................................
ChrCl.roy1 : // .................................................
ChrCl.bin1 : // .................................................
PchCl.pom1 : // .................................................
PchCl.aug1 : // ............V....................................
PchCl.loa1 : // .................................................
PchCl.loa2 : // .................................................

PV32      : // .................................................
PchAr.cri1 : // .................................................
ChrCl.cor1 : // .................................................
ChrCl.swe1 : // .................................................
PchBr.unk1 : // ...................T.............................
PchBr.unk2 : // ...................T.............................
PlmCl.bla1 : // .................................................
PchCl.sum1 : // .........G.......................................
PchUy.ear1 : // .................................................
PchUy.jun1 : // .................................................

PE5       : // ...I........A....................................
NctCl.aug1 : // ...I.............................................
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reported for other viral MPs such as those from cucumber mosaic virus (CMV) [31], TMV [33], brome mosaic virus (BMV) [18] and cowpea chlorotic mottle virus [17]. The observation that the MPs of PNRSV, CMV, BMV, and TMV are functionally interchangeable with the MP of AMV [40] supports the idea of a similar biological RNA binding affinity in vivo for the abovementioned viral MPs.

The alignment of all of the PNRSV MP sequences sequenced so far highlights the existence of residues representative of the PV32-I, PV96-II and PE5-III phylogroups. Residue D/I257 is specific for the PV96-II group (with the exception of the SW6 isolate from cherry, AF013287) meanwhile, amino acids V257 and L261 are representative of the PV32-I group. The PE5-III group contains a large number of specific residues with N32, S/H41, I250, L253, E/G257, S/I264, T266, V274, R/G281 and T282. Below, we will analyze in more detail the relationship between isolate SW6 and the rest of PNRSV groups. With the exception of residues N32 and S/H41, the rest of the specific residues are located at the variable C-terminal region, which has been postulated herein to interact with the cognate coat protein [11]. Residues V257 and L261 have been identified in the group of eight amino acids that allowed CH9 serotypes causing a rugose mosaic disease to be differentiated from those causing mild mottling in sweet cherry.

Functional relevance of the observed variability in CP

The alignment of CP sequences (Fig. 1b) shows a highly conserved protein, except at its N-terminus, where most of the differences among all isolates are contained. The South American PNRSV isolates grouped into the same PV96-II, PV32-I and PE5-III phylogroups obtained with the MP gene, and the intragroup amino acid identity values ranged between 96–100, 95.6–100 and 97.8%, respectively. The analysis of all isolates sequenced so far shows the existence of certain phylogroup-specific residues for PV32-I and PE5-III but not for PV96-II. Isolates belonging to PV32-I are characterized by the presence of two extra residues (N/S42 and R43) and the amino acid Y81. Only isolates CH38 (AF034991) and CH57 (AF034993) or PchBr.unk1 and PchBr.unk2 from the PV32-I group do not contain the additional amino acids or the Y81 residue, respectively. Residues K59, N121, R139, N142 and I181 are specific of phylogroup PE5-III (Fig. 1b). Remarkably, the extra residues of the PV32-I group are inside of the RNA-binding domain described for PNRSV (amino acids 25–50; Fig. 1b) [7]. This motif contains four R residues (five in the PV32-I group) that are critical for the RNA binding capacity of the CP. All PNRSV isolates sequenced so far have the four basic residues except ChrCl.swe1, which contains only three. The extra R observed in the PV32-I phylogroup

could contribute to the RNA-binding capacity of the CP. The K_d observed for a synthetic peptide corresponding to the RNA-binding domain of an PNRSV (isolate PV32) that carries the five basic residues was estimated to be 17 μM, whereas it was 11.3 nM for the entire protein [7]. This discrepancy was explained by the presence of a putative zinc finger domain at the N-terminus [42]. Replacement of some R residues located at the RNA-binding domain by A significantly reduced the RNA interaction capacity of the synthetic peptide [7]. However, isolate ChrCl.swe1 lacks one of the critical R residues. This observation suggests that at least one of the mutated RNA-binding domains, characterized in vitro as having a low RNA-binding affinity, is still functional in vivo in the context of the full protein, indicating that other domains might contribute to the RNA interaction capacity (e.g. zinc finger domain). The observation that all R amino acids of the domain contribute to the RNA affinity of the CP [7] suggests that there is a functional range, with a K_d of 11.3 nM (five R) [7] or larger (four R). In this sense, the PNRSV CP of isolate PV96 that carries four basic residues is functionally interchangeable with the AMV CP, whose K_d is 500 nM [8, 43]. Covariation analyses of PNRSV CP [11] identified a significant association between the residues at sites V48 and D141.

Implications for MP and CP interactions

When covariation analyses were extended to detect MP and CP interactions, the MP residues V253, E256, D257, and T261 (I261 in the alignment used in [11]) were seen to covary with the CP residues T52 and E141, suggesting a putative MP–CP interaction across these regions. Considering all the PNRSV isolates available, the sequences of the MP and CP covarying residues that are characteristic of the PV32-I group are I/V253-D256-V257-L261 and L/V/M/T/I52-D141, while the covarying residues defining the PV96-II group are V/D253-E/K256-D257-T/I261 and T/A/V/I52-A/E141, except for the SW6, Pch.loa1 and 2 isolates, which have a consensus sequence more similar to that of the PV32 group (I/V253-E256-I/V257-V261 and I52-D/N141). Apparently, the presence of an aliphatic residue at MP position 257 (I or V) correlates with the presence of a D/N at CP position 141. This observation suggests that the putative MP–CP interaction for isolates SW6, Pch.loa1 and Pch.loa2 may be more similar to those, which characterize the PV32-I group than to those which characterize the PV96-III group.

Phylogenetic inferences

Previous phylogenetic analyses of the PNRSV MP and CP sequences suggested a division into three groups whose

representative members are isolates PV32, PV96 and PE5 [3, 4]. A similar grouping was later confirmed by Vasková et al. [48], although they proposed an alternative nomenclature for each phylogroup (I, II and III, respectively). Phylogenetic analysis of the MP and CP sequences from the 23 South American isolates characterized herein plus all of the isolates analyzed by Codoñer et al. [11] further supports the existence of the same three phylogroups, irrespective of the protein employed for the analysis (Fig. 2). However, the statistical support for the internal nodes in the phylogenetic tree is, in general, higher in the MP tree than in the CP tree (Fig. 2). The South American isolates were evenly distributed into the three groups, confirming a lack of clustering by geographical origin. However, considering the pollen transmission of PNRSV and the asexual propagation of stone fruit trees, a North American origin for these isolates cannot be ruled out since most *Prunus* varieties come from this geographic region (Table 1). Similarly, clustering does not reflect host species or symptomatology, thus confirming earlier reports [3, 4, 22, 46, 48]. In this sense, most South American isolates clustered into the PV96-II phylogroup cause not only the characteristic mild symptoms but also severe disorders. Moreover, some of the isolates included in the PV32-I phylogroup, which mostly contains isolates causing severe symptoms, were asymptomatic (Table 1).

The phylogenetic positions of isolates PchCL.loa1 and PchCL.loa2 were not well resolved in the CP tree, with a bootstrap value of 42% (Fig. 2b). In addition, both isolates were closely related to isolate SW6, a cherry variant that was grouped outside of the three PNRSV groups [11, 22]. To further test to which PNRSV group these three isolates belong, an alternative statistical approach, the four-cluster likelihood-mapping method, was employed. Figure 3 shows the results of this analysis. In the case of the MP, 72.5% of all of the quartets supported the branching pattern that groups the three isolates (SW6 cluster) and the PV96-II phylogroup versus the remaining sequences. Somewhat less statistical support for this hypothesis was provided by the CP (59.3%). Therefore, based on likelihood mapping, extra support for the inclusion of isolates PchCL.loa1, PchCL.loa2 and SW6 within the PV96-II phylogroup has been provided. Since the main differences between these isolates and the rest of the PV96-II isolates are located at the putative MP-CP interaction residues, the low statistical support may reflect an interesting biological phenomenon: typical PV96-II MP and CP proteins that are interacting as PV32-I proteins do. It would be very interesting to explore the interchangeability of the CP gene between PV32-I and PV96-II phylogroups to confirm this hypothesis.

Targets of natural selection

Considering the already large number of PNRSV sequences available in the database, we decided to analyze which particular MP and CP amino acid sites were under selective constraints. To this end, the difference between nonsynonymous (d_N) and synonymous (d_S) substitution rates $d_N - d_S$ has been used to evaluate the sign and intensity of selection. For the MP, 40 sites were found to be under negative selection and only one site was found to be under positive selection ($p < 0.05$; Fig. 4). In the case of CP, 13 sites were predicted to be under negative selection (Fig. 4b).

Regarding the MP, four of the selected sites are located at the previously characterized RNA-binding motif (S65, E67, G70, and D77) [26]. Interestingly, none of the negatively selected sites correspond to the basic residues that are critical for the RNA-binding capacity. This observation suggests that selection is probably acting on the maintenance of the right secondary/tertiary conformation of the RNA-binding motif. The next six negatively selected amino acid sites (R89, Y94, I97, I98, Q99, and T102) are located inside of the predicted transmembrane domain [42], with I97 and I98 showing the most negative differences. This transmembrane region is highly conserved across all MPs related to the 30K superfamily (the β-1 and β-2 sections of the MP as described by Melcher et al. [36]). The MP region preceding the C-terminal 35 residues allowed the accumulation of selected sites with large negative differences (V231, D236, R238, T242 and P248). It has to be mentioned that no function has been determined for these domains, although we found that a PNRSV MP mutant carrying the N-terminal 245 residues is still functionally interchangeable with the corresponding gene of AMV while a mutant carrying the N-terminal 235 is not (data not shown). The C-terminal 35 residues have been postulated to interact with the cognate CP (see above; Fig. 1b). This C-terminal portion of the protein contains four sites selected under negative selection (I249, T274, V279 and G280) and the only amino acid site predicted to be under positive selection (D255).

In the case of the CP, the selected sites are not related to any of the previously characterized domains except for the C-terminal region responsible for dimer formation (D206, Q211 and T214), which is highly conserved in all members of the genus *Ilarvirus* sequenced to date [5, 25].

In summary, the results presented herein show that all of the PNRSV isolates characterized in South America perfectly group into the three previously described PNRSV phylogroups and share all of the characterized motifs in both their MP and CP molecules.
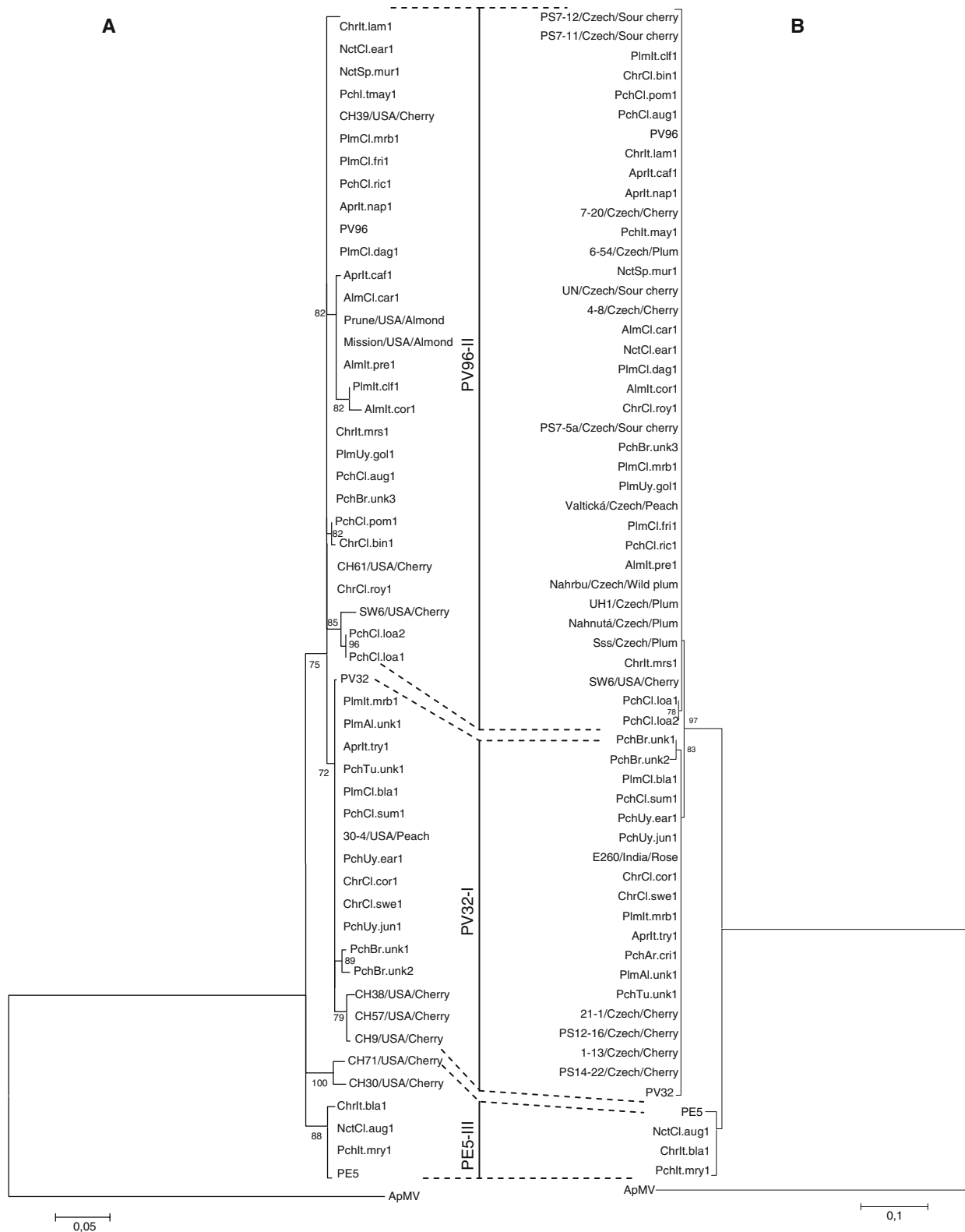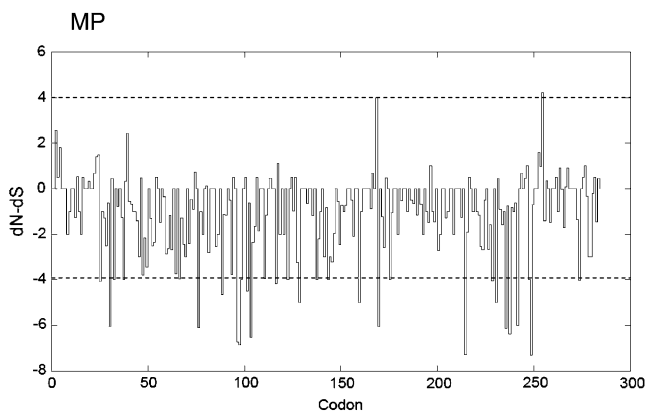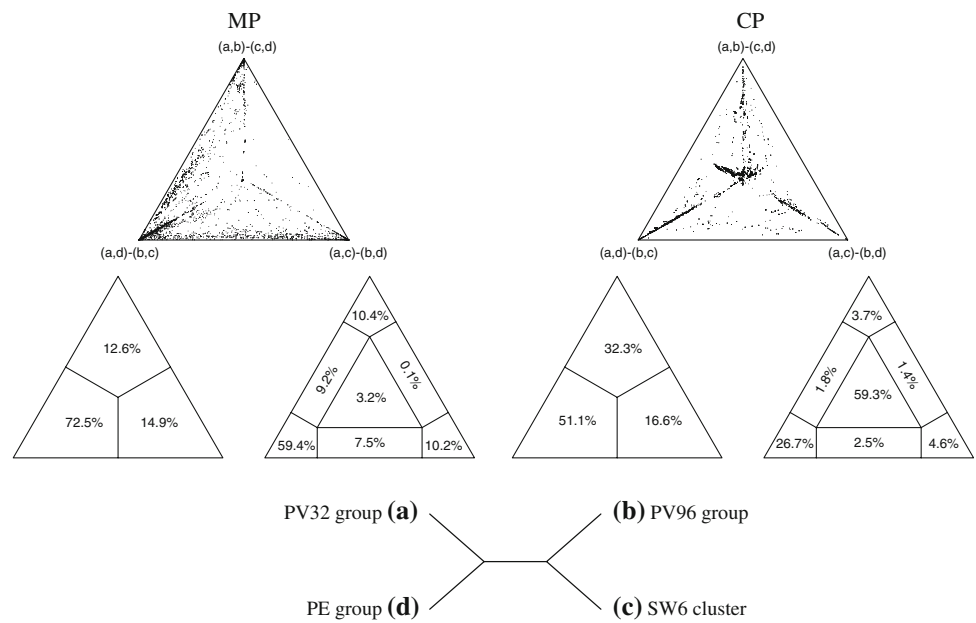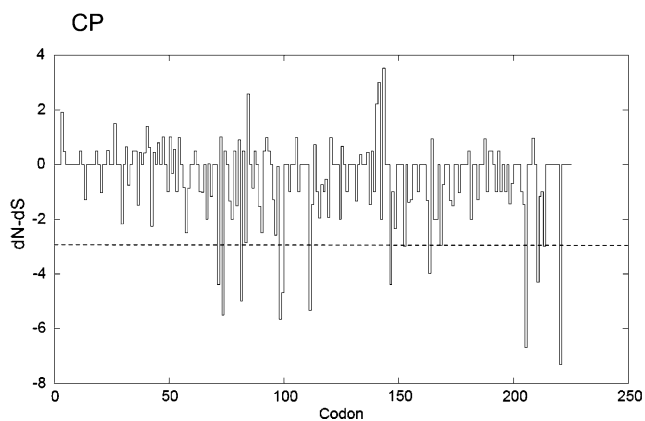
Fig. 2 Phylogenetic trees of MP (**a**) and CP (**b**). The analysis was performed with the 23 PNRSV isolates described here plus isolates analyzed previously by Codoñer et al. [11]. Trees were inferred by the minimum-evolution method. *Numbers* at the nodes represent bootstrap support values based on 10,000 pseudoreplicates. Nodes with bootstrap support <50% have been collapsed. Amino acid sequences representative of the three subgroups are include (PV96, PV32 and PE5). The CP sequence of the Argentine PNRSV isolate (PchAr.cri1, AY007217) has been included in **b**. The trees are rooted to the corresponding amino acids sequences of ApMV (Accession number U15608)

Fig. 3 Quartet likelihood-mapping analysis supporting the inclusion of isolates SW6, PchCl.loa1 and PchCl.loa2 (SW6 cluster) as members of the PNRSV phylogroup PV96-II

Fig. 4 Distribution of $d_N - d_S$ along the sequence of the MP and CP coding regions. Values were computed using a random effects likelihood (*REL*) method. The statistical significance of each value

CODON	d_N-d_S	p-value
255 (D)	4,21	0,0484
26 (I)	-4,07	0,0268
31 (H)	-6,05	0,0015
33 (L)	-4	0,0123
38 (E)	-3,98	0,0191
46 (G)	-3	0,037
48 (H)	-3,78	0,0184
50 (L)	-3,43	0,0438
65 (S)	-3,74	0,019
67 (E)	-3,97	0,0193
70 (G)	-3	0,037
77 (D)	-6,09	0,0015
89 (R)	-4,63	0,0183
94 (Y)	-3,76	0,0231
97 (I)	-6,71	0,0006
98 (I)	-6,84	0,0008
99 (Q)	-3,99	0,0437
102 (T)	-4,5	0,0178
104 (G)	-6,53	0,0025
111 (Q)	-3,94	0,0452
117 (E)	-4,17	0,0166

CODON	d_N-d_S	p-value
123 (T)	-4	0,0123
129 (R)	-4,99	0,0041
138 (G)	-4	0,0146
142 (V)	-3	0,037
144 (S)	-4	0,0147
145 (A)	-3	0,037
160 (P)	-5	0,0041
170 (F)	-6,04	0,0015
176 (S)	-4	0,0144
215 (I)	-7,27	0,008
229 (I)	-4,03	0,0272
231 (V)	-5	0,0041
236 (D)	-6,12	0,0014
238 (R)	-6,36	0,0016
242 (T)	-6	0,0013
248 (P)	-4	0,0123
249 (I)	-7,3	0,0125
274 (T)	-4	0,0449
279 (V)	-3	0,037
280 (G)	-3	0,037

CODON	d_N-d_S	p-value
72 (N)	-4,39	0,0117
74 (P)	-5,5	0,0068
82 (V)	-5	0,0041
99 (S)	-5,65	0,003
100 (I)	-4,68	0,018
112 (Q)	-5,32	0,0129
147 (N)	-4,39	0,0117
153 (G)	-3	0,037
164 (F)	-3,98	0,0158
206 (D)	-6,68	0,0033
211 (Q)	-4,29	0,0308
214 (T)	-3	0,037
221 (E)	-7,3	0,0008

was evaluated by its Bayesian posterior probability. Amino acids sites predicted to be under either a negative or positive selection are indicated with their corresponding $d_N - d_S$ and *P*-values

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