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ABSTRACT

Indirect evidence of mitochondrial viruses in plants comes from discovery of genomic fragments integrated into the nuclear and mitochondrial DNA of a number of plant species. Here, we report the existence of replicating mitochondrial virus in plants: from RNAseq data of infected *Chenopodium quinoa*, a plant species commonly used as a test plant in virus host-range experiments, among other virus contigs, we could assemble a 2.7 Kb contig that had highest similarity to mitoviruses found in plant genomes. Northern blot analyses confirmed the existence of plus and minus strand RNA corresponding to the mitovirus genome. No DNA corresponding to the genomic RNA was detected, excluding the endogenization of such virus. We have tested a number of *C. quinoa* accessions, and the virus was present in a number of commercial varieties, but absent from a large collection of Bolivian and Peruvian accessions. The virus could not be transmitted mechanically or by grafting, but it is transmitted vertically through seeds at a 100% rate. Small RNA analysis of a *C. quinoa* line carrying the mitovirus and infected by alfalfa mosaic virus showed that the typical anti-viral silencing response active against cytoplasmic viruses (21-22 nt vsRNA peaks) is not active against CqMV1, since in this specific case the highest accumulating vsRNA length is 16, which is the same as that corresponding to RNA from mitochondrial genes. This is evidence of a distinct viral RNA degradation mechanism active inside mitochondria that could possibly have also an anti-viral effect.

IMPORTANCE

This paper reports the first biological characterization of a bona fide plant mitovirus in an important crop, *Chenopodium quinoa*, providing data supporting that mitoviruses have the typical features of cryptic (persistent) plant viruses. We for the first time demonstrate that plant mitoviruses are associated with mitochondria in plants. In contrast with fungal mitoviruses, plant mitoviruses are not substantially affected by the anti-viral silencing pathway, and the most abundant mitovirus small RNA length is 16 nt.

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INTRODUCTION

The family *Narnaviridae* comprises two genera of positive single-stranded (ss) RNA viruses, 52 the *Narnavirus* and the *Mitovirus*, both originally thought to be mycoviruses, because fungi are the 53 main natural host in which they can replicate. Both are known to be naked viruses presenting only one 54 ORF that encodes one protein, the RNA-dependent RNA polymerase (RdRp) (1). The two genera are 55 distinguished based on phylogenetic analysis (each of them belongs to a distinct, statistically well-56 supported clade) and on subcellular localization: the narnaviruses replicate and stay in cytosol, 57 whereas the mitoviruses typically replicate and persist in the mitochondrion. Evidence of their 58 mitochondrial localization is provided by the fact that they fractionate with the mitochondrial fraction 59 (2) and that for most of them the ORF is translated using the mitochondrial genetic code (1). The 60 mitoviruses described so far are able to infect only mitochondria of filamentous fungi and in some 61 cases they are associated with hypovirulence (1). Sometimes it seems that the mitochondrial 62 morphology is not affected by the virus, whereas in some other cases they can cause a morphological 63 alteration (i.e. fibrous mitochondria) which were possibly associated with the induced hypovirulence 64 (3). 65

Some fungal mitoviruses have the potential to use both the nuclear and mitochondrial genetic 66 code for the translation of their genomes, and based on this, some authors have hypothesized 67 promiscuous replication in both mitochondria and cytoplasm. However, bioinformatic analysis leads 68 to a different explanation: in those host species where mitovirus RdRp can hypothetically be translated 69 using both genetic codes, the mitochondrial genes have a strong bias for the tryptophan codon that is 70 shared with the nuclear genetic code (UGG). This implies that the exclusive use of UGG codon for 71

tryptophan in some mitovirus is not because of a promiscuous lifestyle between mitochondria and 72
cytoplasm, but it reflects the fact that UGA (the only differential codon between the nuclear and 73
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mitochondrial genetic codes in the fungal-plant host species) is not present in general in genes 74
encoded by the mitochondria (4). 75

Intriguingly, species of genus Mitovirus are most closely related to the Leviviridae, the only 76
taxonomically defined group of (+)ssRNA bacterial phages (5, 6). Within the context of the theory of 77
mitochondria derived from an alpha-proteobacterial endosymbiont (7), this suggests that mitoviruses 78
likely derived from an ancestral mitochondrial phage by losing the capsid protein (CP) (8) which is 79
unnecessary due to the absence of an extracellular stage (9). 80

A further interesting observation related to mitoviruses comes from the availability of a great 81
number of complete plant and fungal genomes that were mined to reveal the presence of non-retroviral 82
endogenous RNA viral elements (NERVEs): complete genomes or partial/complete gene sequences of 83
RNA viruses are present in almost all the eukaryotic nuclear genomes (10-13). Specifically, Bruenn 84
and co-authors demonstrated the widespread presence of mitoviral sequences in many plant nuclear 85
and mitochondrial genomes (13). Two different hypotheses can explain this evidence: the first is 86
diverse integration events of a fungal mitovirus, or of a native plant mitochondrial virus, into the 87
mitochondrial genome, and from here to the nucleus as a result of mitochondrial DNA transfer (13, 88
14); the second proposes the possibility of integration of such sequences in plant genomes via fungal 89
mediated horizontal gene transfer (HGT) during the long-term coevolution of fungi and plants (15, 90
16). Very recently, indirect evidence of replicating plant mitoviruses was provided mining the 91
transcriptome of a number of plant species (17). 92

We here report the complete genome sequence and biological characterization of a replicating 93
plant mitovirus detected in a number of *Chenopodium quinoa* accessions and designated as 94
Chenopodium quinoa mitovirus 1 (CqMV1). Furthermore, we provide evidence of differential small 95
RNA (sRNA) processing of this virus compared to a cytoplasmic plant virus infecting the same plant, 96
indicating the possible involvement of a still uncharacterized new differential anti-viral response 97
inside the mitochondria. 98

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RESULTS 100

In silico assembly of a mitovirus sequence from NGS analysis of total RNA from a *C. quinoa* leaf sample. 101
102

In order to identify a mechanically transmitted viral agent from a *Hibiscus rosa-sinensis* plant, 103
we submitted for NGS analysis the total RNA (depleted from ribosomal RNA) of *C. quinoa* leaves 104
showing chlorotic spots that were inoculated with sap from symptomatic *H. rosa-sinensis* plants 105
(Accession SRR8169409). Our bioinformatics pipeline (18) identified two complete virus genomes: 106
BLAST searches of viral databases identified a contig with high similarity to a tobamovirus -*Hibiscus* 107
latent ford pierce virus- (19), and a second contig of 2730 nt, which codes for a single putative 108
protein, from nt 322 to nt 2644 (Fig. 1). A BLAST search identified the latter as a putative mitovirus 109
RNA-dependent RNA polymerase (RdRp). Since such a contig was present also in un-inoculated *C. quinoa*, and not in the original hibiscus plant (data not shown), we decided to provisionally name the 111
putative virus *Chenopodium quinoa mitovirus 1* (CqMV1). Furthermore, to confirm absence of 112
fungal contamination (from endophytes or pathogenic fungi), assembled contigs were analyzed in 113
MEGAN 6 after DIAMOND processing for taxonomical placements of all the assembled contigs from 114
the RNAseq experiment: no fungal contigs were detected (not shown). 115

At the time when we identified the putative mitovirus (Jan 2017, deposited in the databases 116
June 22nd 2017), a tblastn search was performed by using the deduced *C. quinoa* mitovirus RdRp as 117
query, and when the total nr database was used, the first hits were those of a *Solanum tuberosum* 118
mitochondrion gene (XP_006364252; e-value 0.0, 98% query coverage, 53% amino acid identity). A 119
number of other Non-retroviral Endogenized RNA Virus Element (NERVEs), described as mitovirus-120
like sequences, are also present in the list of hits obtained by this search. Repeating the tblastn search 121
at the time of submission (October 2018), when limiting to annotated virus sequences, the highest 122
score is to a recently identified *Beta vulgaris* mitovirus 1 (AVH76945.1; 56% identity at the aa level, 123
82% query cover) (17) that was deposited in the database Dec 21st 2017, and a still uncharacterized 124
Ocimum basilicum RNA virus 2 sequence (YP_009408146; 32% aa identity, 49% query cover) that 125
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was deposited in the database June 3rd 2017. To exclude that such RNA mitoviral sequences resulted 126
from transcription of a full length viral genome endogenized in *C. quinoa*, we investigated the 127
possible existence of a DNA fragment in the mitochondrial or nuclear DNA corresponding to the 128
assembled viral sequence. For this purpose, we designed specific primers on the predicted ORF and 129
we performed a PCR protocol on both DNA and RNA to identify in which nucleic acid fraction the 130

sequence is detectable. *C. quinoa* plants contained the viral sequence in the RNA fraction, but none of 131 them showed any specific band in the DNA fraction (Fig. 1). These results were also confirmed in a 132 more sensitive qRT-PCR protocol (not shown). 133

We then analyzed the CqMV1 amino acid sequence to identify conserved motifs. In addition 134 to the conserved RdRp domain (GDD), we detected all six conserved motifs characteristic of 135 mitoviruses (20). Due to the putative mitochondrial localization of the virus, we expected an AU 136 content > 60% because of the A-U rich nature of mitochondrial genome (21, 22) but the observed A-U 137 content is 58.39%. Whereas fungal mitoviruses typically use UGA to encode tryptophan rather than a 138 stop codon, in our case all tryptophans are encoded by UGG. 139

Stem-loop secondary structures are characteristic of 5' and 3' UTRs sequences of the positive 140 strand mitovirus genomes (3). We here analyzed presence of possible secondary structures with the 141 RNA-Fold software which revealed the presence of these structures at the 5' UTR region ($\Delta G = 33$ 142 kcal/mol; not shown). 143

We then proceeded to carry out a phylogenetic analysis that included representatives of 144 endogenized plant mitoviral sequences, characterized mitoviruses from fungi, and mitoviruses 145 characterized from other plant transcriptomes; as outgroups we included representative of the 146 Narnaviridae family and the recently proposed ourmiaviridae-like family (Fig. 2). As observed by 147 other authors, plant mitoviruses form a well-supported clade within the mitoviruses. Currently, 148 known mitoviral sequences often group according to their specific host, consistent with co-evolution 149 and infrequent interspecific transmission. 150

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CqMV1 is differentially distributed in *C. quinoa* accession and cultivars. *Chenopodium* 151 *quinoa* is known to plant virologists because it is historically a common host-range test plant: it often 152 gives local chlorotic or necrotic lesions upon mechanical inoculation with a number of plant viruses. 153 This species is also a very important agricultural crop (23, 24) because of its resistance to a number of 154 abiotic stresses and for its nutritional value (25, 26). 155

We therefore sought to determine how widespread CqMV1 is within *C. quinoa* germplasm by 156 testing seed batches from different sources for the presence of the virus. For this purpose, initially we 157 compared *C. quinoa* accessions from different plant virology laboratories (personal collection of 158 Gancho Pasev, Maritsa Vegetable Crops Research Institute, Plovdiv, Bulgaria); later, we purchased a 159 variety of accessions from the Leibniz Institute of Plant Genetics and Crop Plant Research (IPK) and 160

from the Collection of the U.S National Plant Germplasm System (NPGS). A sensitive and fast qRT-161 PCR protocol was established to screen newly germinated batches of 10 plantlets for each sample. We 162 found that the *C. quinoa* from our laboratory used as test plants in host range experiment (named in 163 this work IPSP1), some other accessions from the Pasev collection (Supplementary on line table 1), 164 and some common commercial cultivars (cv Regalona and cv Cherry vanilla) were positive for 165 CqMV1. In contrast, CqMV1 did not infect 42 tested accessions from Peru and Bolivia (from the IPK 166 collection) (Supplementary on line table 1). Accession PI614886 (NPGS collection) from Chile is the 167 *C. quinoa* accession used for deriving the recently published genome sequence and it carries CqMV1 168 (27). 169

CqMV1 genomic RNA accumulates preferentially in the mitochondrial fraction. 170

Translation of CqMV1 RdRp could hypothetically occur in both the cytoplasm and the mitochondria 171 since the same protein is encoded using both genetic codes. For this reason, we wanted to investigate 172 if virus RNA accumulated preferentially in mitochondria-enriched preparations, compared to whole 173 cell extracts, or to chloroplast-enriched fractions. We adapted a protocol for spinach mitochondrial 174 enrichment and judged purity of the preparation based on chlorophyll fluorescence (Fig. 3A) and 175 presence of specific genetic markers (ORF-X and S3 for mitochondria, S2 for a mitochondrial 176

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ribosomal protein encoded by a nuclear gene, and Cox for a nuclear encoded gene). RNA was 177 extracted from the different fractions and qRT-PCR was carried out to detect the virus and the mRNA 178 corresponding to each of the marker genes. The virus co-purifies with mitochondrial RNAs. It is >50 179 fold more abundant in the mitochondrial fraction than in either the chloroplastic or the soluble 180 cytoplasmic RNA fractions (Fig. 3B). 181

Evidence of minus strand CqMV1 RNA accumulation in leaf extracts. The recent 182 contention of “contemporary” mitoviruses infecting plants (17) relies on detection of RNA transcripts 183 corresponding to the mitovirus genome (positive RT-PCR, after DNase treatment) in the absence of 184 a corresponding DNA segment (negative PCR). This is robust indirect evidence. Nevertheless, more 185 direct evidence of replicating mitoviruses could come from detection of a negative strand full length 186 RNA corresponding to the viral genome by a PCR-independent method. Therefore, a northern 187 hybridization experiment relying on positive and minus strand run-off transcript probes to detect – 188 strand genomic RNA and + strand genomic RNA respectively, has been performed. We used the 189 accession BO25 as a negative control and Regalona as CqMV1 infected positive control. Our initial 190 experiment with a first pair of + and – strand probes revealed a very abundant accumulation of a 191

specific positive strand CqMV1 genomic RNA band only in the virus-infected line. Attempts to detect 192 a specific full length -strand RNA band from leaf extracts failed; only shorter virus specific RNA 193 species could be detected by the positive strand probe designed close to the 3'end of the genome (Fig 194 4A); the presence of a specific full length -strand viral genomic RNA is likely masked by the 195 unspecific hybridization of the probe with a ribosomal band. We repeated the experiment with a 196 second pair of probes designed in a different region of the genome (Fig. 1), and in this case, we were 197 able to show a faint specific band hybridizing with the -sense genomic RNA after 15 days membrane 198 exposure to film (Fig. 4B), evidence of a minimal CqMV1 replicative activity. 199

Mechanical inoculation, seed transmission and grafting experiments. We then proceeded 200 to investigate some basic biological properties of CqMV1. Most persistent viruses are not 201 mechanically transmissible but are vertically transmissible with 100% rate through seeds. We first 202

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sought to mechanically transmit CqMV1 from infected *C. quinoa* to accession BO25, which had 203 tested negative for CqMV1. Out of 20 inoculated plants, none showed evidence of infection in either 204 the inoculated leaf, or systemically, following standard protocols that gave 100% infection with a 205 mechanically transmissible control virus (alfalfa mosaic virus=AMV; not shown). 206

Next, we investigated the vertical transmission rate of the virus in plantlets germinated from 207 seeds obtained from infected plants (IPSP1 and cv Regalona). We tested individually 100 plantlets for 208 each accession by qRT-PCR and all tested positive. Finally, we tested the possibility of transmitting 209 CqMV1 through grafting, which often overcomes the mechanical transmission limitations of non-210 mechanically transmissible viruses. We grafted healthy BO25 on IPSP1 rootstocks (6 plants). We then 211 tested the scion and graft at 1 and 2 months after grafting. CqMV1 could not be detected in any 212 grafted scion systemically, but its presence was confirmed in all the infected rootstocks. 213

Differential symptom severity of pathogenic virus infections on CqMV1-infected and 214 CqMV1-free *C. quinoa* lines. We focused our experiments on the accessions BO25 and BO78 as 215 negative controls and IPSP1 and Regalona as CqMV1 infected positive controls since they did not 216 show any evident phenotypic differences in our environmental growth conditions. We wanted to test if 217 the presence/absence of the mitovirus has any synergistic or antagonistic effect once the plants are 218 inoculated with a disease-causing plant virus. In particular, the four accessions have been infected 219 with three different pathogenic viruses able to replicate, systemically infect and induce symptoms on 220 *C. quinoa* plants: i) an isolate of AMV, ii) an isolate of lettuce mosaic virus (LMV) and iii) an isolate 221

of hibiscus latent ringspot virus (HLRSV). 222

Leaves of virus-inoculated plants were compared to leaves of mock-inoculated plants 7 days 223
post infection (dpi): for both AMV and HLRV we observed local chlorotic lesions indistinguishable 224
between the CqMV1-infected lines and the CqMV1-free lines (Fig. 5A and Fig. 5C). In contrast, 225
leaves of plants infected with LMV displayed local chlorotic lesions when the leaves came from 226
CqMV1-infected lines, but more severe local necrotic lesions in leaves from CqMV-free lines (Fig. 227
5B). 228

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We then repeated symptoms observation on whole plants at 14 dpi. The CqMV1-infected lines 229
IPSP1 and Regalona displayed milder systemic symptoms (Fig. 6D-E-F and Fig. 6J-K-L) compared to 230
the CqMV1-free lines BO25 and BO78 (Fig. 6G-H-I and Fig. 6A-B-C). All four lines showed 231
comparable mild growth impairment, leaf malformation and mild mottling in upper uninoculated 232
leaves. Nevertheless, the most evident specific differential phenotype associated to CqMV1 233
presence/absence is the red-violet stem pigmentation observed in CqMV1-free lines (Fig. 6M). 234
Conversely, mock inoculated or CqMV1-infected plants did not show stem pigmentation (Fig. 6N). 235
Differential small RNA accumulation and processing of AMV and CqMV1 in infected C. 236
quinoa plants. A number of fungal mitoviruses have been discovered through sRNA characterization, 237
suggesting that--at least in fungi--they are subject to RNAi processing (28, 29). In order to 238
characterize the sRNA present in C. quinoa we decided to compare three libraries of sRNA: i) from 239
CqMV1-free BO25 infected by AMV (Accession SRR8169660); ii) mock-inoculated cv. Regalona 240
carrying CqMV1 (Accession SRR8169658); iii) cv Regalona mechanically infected with AMV 241
(Accession SRR8169659). Using a bioinformatic pipeline that was previously used to assemble de 242
novo virus genomes (18, 30), we first confirmed that accession BO25 does not carry any virus other 243
than AMV, whereas the same pipeline could assemble CqMV1 from cv Regalona and AMV from 244
AMV-infected cultivars. When we looked at the percentage of total sRNA that mapped to the two 245
viral full length sequences we noticed that, for plant mitovirus, the percentage is very low compared to 246
AMV (Table 1), even if the overall genomic RNA is actually more abundant based on qRT-PCR 247
assessment and northern blot (not shown). We then proceeded to look at the size distribution of the 248
sRNA. The availability of the C. quinoa genome (27, 31) allowed us to assess reads mapping to the 249
host genome (with a characteristic peak at 24 nt length common to most plants) and to each of the two 250
viruses. In the case of AMV a typical peak corresponding to 21 nt length (47% of the reads) and a 251
minor 22 nt length peak (30% of the reads) are likely the hallmark of a Dicer and RISC mediated anti-252

viral response (Fig. 7). Surprisingly, in the case of reads mapping to CqMV, a sharp peak for the 16 nt 253 long reads corresponding to 30% of the reads was present, whereas the peaks corresponding to 21 and 254

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22 nt long reads were both below 10% (Fig. 7). Availability of *C. quinoa* chloroplast (32), and nuclear 255 (27) genomes and our own selection of a number of mitochondrial genes from our transcriptome data, 256 allowed us to check the read-distribution lengths of host sRNA mapping to those genomes. Small 257

RNA mapping onto the chloroplast genome had a major peak at 27 nt length (22%), whereas the 258 second peak is at the 22 nt; in the case of sRNA mapping to mitochondrial genes (in this case only the 259 putative coding sequences were used), a sharp peak is present for 16 nt length (67% of the sRNA). 260

The N terminal nucleotide distribution of reads mapping to CqMV and AMV were A=30.5% 261

C=12.9% G=37.7% U=18.9% and A=25.6% C=16.3% G=10.2% U=47.9%, respectively. 262

We then performed an sRNA miner analysis on reads mapping on the CqMV1 genome to 263

reveal clustered organellar sRNA (cosRNA), which are putative footprints of RNA binding proteins 264

(RBP): using conservative settings we were able to detect eight different footprints from 4 distinct 265

peaks (Supplementary on line Fig. 1). Therefore, sRNA analysis seems to confirm that most of the 266

CqMV1 RNA is likely protected from dicer/RISC mediated viral anti-silencing response inside the 267

mitochondria, where distinct processing occurs resulting mostly in 16 nt length sRNA. 268

269

DISCUSSION 270

Plant mitoviruses, a new class of cryptic (persistent) plant viruses 271

Our work confirms the recent finding that plant mitochondrial viruses exist not only as 272

widespread endogenized sequences in mitochondrial or nuclear genomes (13), but also as true virus 273

encoded RdRp-dependant replicating RNA elements (17). Here we provide further molecular 274

evidence through detection of minus strand RNA. We can also exclude that such a replicating virus is 275

carried by a fungal endophyte based on RNAseq analysis, as our dataset lacks of any fungal reads. 276

Our work is the first to provide evidence that bona fide plant mitoviruses are enriched in plant 277

mitochondria. 278

This is not the first case of a virus associated with plant mitochondria. The plant virus 279

carnation Italian ringspot virus replicates on mitochondrial external membranes and causes 280

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multivesicular body alterations of mitochondria (33) but particles and RNA readily accumulate in the 281 cytoplasm, and replicases (p35 and co-terminal p95) targeted to the external membrane of 282 mitochondria have N and C termini on the cytosolic side (33), so this virus is best viewed as 283 cytoplasmic. 284

We also note that viruses have already been characterized from chloroplasts and mitochondria 285 of the green alga *Bryopsis* sp. (34), specifically, a mitochondrial virus related to fungal totiviruses (35) 286 and a chloroplastic virus related to the partitiviruses (36); both viruses have a typical dsRNA genome 287 and have no phylogenetic relationship with any known bacterial. In contrast, plant mitoviruses, 288 including CqMV1, are instead phylogenetically related to the phage family Leviviridae. 289

Our phylogenetic analysis would suggest a comprehensive review of the taxonomy of 290 mitoviruses. Inclusion of a number of well-characterized fungal and plant narna-like viruses and 291 members of the family Leviviridae in the phylogenetic analysis makes it evident that mitoviruses are 292 monophyletic. Their wide diversity warrants the establishment of a family taxon called Mitoviridae, 293 separated from the family Narnaviridae. The newly established family would comprise a number of 294 subfamilies and genera, including the very distantly related mitovirus species characterized from the 295 arbuscular mycorrhizal fungus *Gigaspora margarita* which are included in a clade basal to existing 296 characterized mitoviruses (37). As already observed by other authors (17), plant mitovirus are nested 297 in a specific fungal mitovirus clade, therefore raising questions about the evolutionary trajectory of 298 plant mitoviruses which has been discussed at length elsewhere (17). 299

We provide here for the first time a basic biological characterization of a plant mitovirus, 300 which has the typical features of cryptic viruses: they cannot be transmitted horizontally by 301 mechanical inoculation or grafting, whereas they are transmitted vertically at a 100% rate through 302 seeds. Lack of transmission through grafting is indeed an expected result for mitoviruses that replicate 303 in plant mitochondria, since movement of mitochondria through the plant is likely minimal and will 304 not replace existing population of mitochondria. This is somewhat different to what observed in fungi 305

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where “grafting” (hyphal fusion through anastomoses) was shown to allow mitovirus transmission in 306 some cases (38). 307

The high level of seed transmission observed in this study is also expected, as seeds formed on 308 infected plants must inherit their parents’ infected mitochondria. Plants have mechanisms to ensure 309 that mitochondria are only maternally inherited and to maintain low heteroplasmy levels. 310

Nevertheless, there is evidence of biparental transmission of mitochondria in plants (39). *C. quinoa* is normally self-pollinating in nature, but studies of mitochondrial inheritance could potentially be carried out through artificial mechanical emasculation and forced crosses (40). An important avenue of future work will therefore be creation of reciprocal crosses with mitovirus-infected and mitovirus-free maternal and paternal lines to allow isolation of quasi-isogenic plants differing only for the mitochondrial/viral content.

Absence of a movement protein in mitovirus genomes and lack of a movement-complementing virus in CqMV1 infected *C. quinoa* also supports the idea that they can infect all type of cells, including meristematic ones, as is the case of other cryptic viruses (41).

The existence of cryptic viruses in plants was known for at least four decades, since their discovery in a number of different plant species at the end of the 1970s (41). Currently known persistent/cryptic plant viruses include members of five families: Partitiviridae (42-44), Totiviridae (45, 46), Chrisoviridae (47, 48) Endornaviridae (49) and Amalgaviridae (50-53). We here provide convincing evidence that plant mitoviruses should be also defined as cryptic (persistent) viruses. The fact that CqMV1 is present in commercial varieties of *C. quinoa* (Regalona and Cherry Vanilla), but that a number of other accessions from Bolivia and Peru do not carry any mitovirus, seems to support the idea that they cause no specific harm to their host. Moreover, the widespread occurrence of mitoviruses in domesticated material raises the possibility of their beneficial role in specific agro-ecological niches. Recent reviews and metagenomics studies unveiled the widespread occurrence of cryptic (persistent) viruses in plants and other authors have discussed a possible beneficial role for their host (54-56). In particular some studies have looked at the presence of endornavirus as it relates

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to the domestication of pepper (49). There is also growing evidence that other plant viruses, which are not defined as cryptic since in some specific instances they can cause obvious symptoms, can indeed provide advantages to their host in resistance to abiotic stress: an example is grapevine rupestris stem pitting associated virus (GRSPaV) where a molecular mechanism has also been proposed (58, 59).

A preliminary experiment to detect differential symptom reaction to a panel of viruses systemically infecting *C. quinoa* did not reveal any major synergistic or antagonistic effect caused by CqMV1 mitochondrial infection. Nevertheless, some interesting features in specific virus-plant interactions should be pointed out: the enhanced necrotic hypersensitive reaction and the stem pigmentation in the absence of CqMV1 implies that infection with the mitovirus can somehow ameliorate the symptoms of at least some other viruses. We can speculate that the presence of CqMV1 in the mitochondria

might alter the oxidative stress cellular signaling resulting in necrosis and pigment accumulation. 342
Further classes of biotic and abiotic stress should be tested, in view of the fact that mitochondria are 343
central in a number of stress related phenomena in plants, particularly in the roots mediating tolerance 344
to harsh environments (60). 345
Numerous attempts to isolate CqMV1 dsRNA from infected plants have failed, using a protocol 346
previously described (61) and using as positive control tomato mosaic virus infected *C. quinoa* and the 347
fungal isolate MUT4330 previously described (61). Lack of detectable dsRNA in CqMV1-infected *C. 348*
quinoa could be due to the extreme toxicity of dsRNA to plant mitochondria, as recently observed for 349
dsRNA expressed in mitochondria in human cell lines: a specific degradation pathway prevents the 350
export of dsRNA and the onset of a general anti-viral defense driven by MDA5 dependent anti-viral 351
signaling (62). 352
Differential RNA processing inside mitochondria and evidence of a yet undefined anti-viral 353
response 354
Our sRNA analysis indicates that the mitochondrial virus present in *C. quinoa* is not subject to 355
the Dicer/argonaute dependent anti-viral silencing response that typically targets plant viruses (63). 356
This is not due to a defective silencing response in *C. quinoa*, because we provided evidence that the 357
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typical Dicer/argonaute processing occurs in the same AMV-infected *C. quinoa* plant extract. This 358
raises the question of what might limit mitochondrial virus replication in plants and fungi, where 359
mitoviruses occur. Silencing of fungal mitoviruses is not well characterized, but two recent studies 360
show that mitoviral sRNA generated in fungi is not different from that generated from cytoplasmic 361
viruses in either quality or quantity (28, 29). Our discovery that plants accumulate very low amounts 362
of mitovirus sRNA and that these are most frequently 16 nt in length is a major difference with fungal 363
systems and raises the question of what molecular pathway generates such a specific sRNA size 364
distribution. Furthermore, the significant differences in the N terminal nucleotide of reads mapping to 365
CqMV and to AMV also point to distinct degradation machineries for the two viruses. Our data 366
suggest that the 16nt mitoviral sRNA likely result from a non-viral specific RNA degradation process, 367
since size distribution of sRNA generated from mRNAs expressed from the mitochondrial genome 368
shows the same peak at 16nt. In this respect, the combined roles of exoribonucleases such as PNPase 369
and RNR1 and RNA binding proteins in leaving footprints of various lengths could be at the basis of 370
this differential sRNA accumulation, consistent with recent studies of size distribution footprints in 371

chloroplast and mitochondrial RNA transcripts in *Arabidopsis thaliana* (64). Previous authors 372
hypothesized a possible regulatory role of the sRNA resulting from processing by Pentatricopeptide 373
Repeat proteins (PPR proteins, a subset of RBP proteins typical of plants) (64, 65), providing 374
therefore a testable model of anti-viral defense based on intra-mitochondrial sRNA generation. 375
From an evolutionary perspective, it would be interesting to look at bacterial anti-viral 376
responses against RNA viruses (RNA phages). Recent work has shown that RNA bacteriophage 377
diversity is much higher than previously thought (66): while current taxonomy has only two family of 378
prokaryotic RNA viruses, the Leviviridae and the Cystoviridae, indirect evidence indicates that some 379
Picobirnaviridae are bacterial viruses (67). In this respect also new anti-viral defense systems are 380
constantly unveiled (68). Although RNA-guided RNA cleavage by a specific CRISPR RNA-Cas 381
system is known (69), its role in specific anti-viral response in natural systems has yet to be shown 382
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(70, 71); nevertheless, in an in vitro heterologous system a type III-A CRISPR-Cas system will restrict 383
MS2 RNA phage infection (72). 384
Future work will pursue further biochemical characterization of the sRNA response to 385
mitovirus infection in different biological systems (fungi and plants) and the analysis of possible 386
differential physiological reactions linked to mitovirus infection in plants experiencing harsh 387
environmental stress conditions. 388
389

MATERIALS AND METHODS 390

Plant material and RNA sequencing. Plant seeds used in this study (accession, seeds from 391
personal collections and from public repositories) are described in detail in Supplementary online 392
Table 1. 393
Total RNA extraction from IPSP1 plants was performed using the Spectrum Plant Total RNA Kit 394
(Sigma-Aldrich, Saint Louis, MO, USA) following manufacturer instructions. RNA quantification and 395
quality were tested using NanoDrop 2000 Spectrophotometer (Thermoscientific, Waltham, MA, 396
USA). Macrogen Europe (Amsterdam, Netherlands) performed ribosomal RNA (rRNA) depletion 397
using the Ribo-Zero™ Plant Kit (Epicentre, Madison, WI, USA), library construction and sequencing 398
using an Illumina HiSeq4000. 399
For mechanical inoculation experiments we used three different virus species, belonging to 400
different families, that systemically infect *C. quinoa*: the LMV, family Potyviridae, genus Potyvirus 401
(Dim 60, PLAVIT collection), AMV, family Bromoviridae, genus Alfamovirus (IFA 30, PLAVIT 402

collection) and HLRSV, family Secoviridae, genus Nepovirus (VE 453, PLAVIT collection). 403
Bioinformatics analysis and molecular validation. Raw reads obtained from total RNA 404
sequencing were assembled into contigs using Trinity 2.3.2 (73) and virus were identified as 405
already described (18) using Blast+ suite 2.6.0 (74), BWA 0.7.15-r1140 (75) and samtools 1.3 (76). 406
Once viral sequences were identified specific primers were designed (Supplementary on line 407
Tab. 3) to reveal the molecular nature (if DNA, RNA or both). To detect viral sequence possibly 408
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integrated into in the host genome (both nuclear or mitochondrial) we performed a total nucleic acid 409
extraction using a phenol-chloroform protocol (77). We then performed, on half of the volume, a 410
digestion with RNase A (four hours) to completely remove any trace of RNAs. The second half of the 411
total nucleic acid extraction was subjected to a four-hour DNase treatment in order to completely 412
remove all traces of DNA. The DNased RNA was then used in a retrotranscription (RT) reaction in 413
order to obtain cDNA suitable for PCR. We then performed PCR with specific primers for both an 414
internal control (COX) and for the viral sequence using as template the obtained DNA and cDNA 415
samples from the four C. quinoa lines (IPSP1, Regalona, BO25 and BO78). The same templates were 416
used in quantitative retrotranscriptase PCR (qRT-PCR) with specific primers to evaluate presence and 417
quantities of the internal control and the viral sequence in the four different lines. 418
PCR products for viral sequence were cleaned with DNA clean & concentrator™-5 kit 419
(ZymoResearch, CA, USA), cloned into pGEM-T easy vector (Promega, Madison, WI, USA) and 420
sequenced at Bio-Fab Research (Rome, Italy). 421
The 5' and 3' terminal sequences were obtained through the RACE protocol. Presence of possible 422
secondary structure was evaluated using RNAfold (78). 423
To identify possible contamination with fungal sequences in our RNAseq experiments we 424
analyzed taxonomic placement of all the assembled contigs using MEGAN6 software (79). 425
Small-RNA sequencing and analysis. To detect CqMV1 small RNAs (sRNAs) we started 426
from total RNA extraction of mock inoculated accession Regalona and Regalona infected by AMV. In 427
parallel BO78 accession infected by AMV was used as negative CqMV1 control. Total RNA from the 428
three samples were sent to the Italian Institute for Genomic Medicine (IIGM, Turin, Italy) where 429
sRNAs were isolated, library constructed and then sequenced using a MiSeq System (Illumina Inc., 430
San Diego, CA, USA). 431
Raw reads were cleaned from adaptor, quality filtered using the FASTX-toolkit 432

(http://hannonlab.cshl.edu/fastx_toolkit/index.html) and then assembled using Velvet (80) and Oases 433 (81). Contigs were used in blastx and blastn searches against a custom database to identify viral 434 on February 4, 2019 by guest<http://jvi.asm.org/>Downloaded from Nerva et al. J. Virol.

sequences. To determine the sRNA size distribution on viral genomes (both CqMV1 and AMV) we 435 used BWA (75) and samtools (76) to map raw reads against the two viral genomes and we then 436 filtered for reads length through a custom Perl script. Due to the unexpected pattern of sRNA mapping 437 on CqMV1 genome we decided to map also raw reads against C. quinoa genes. For nuclear and 438 chloroplast genes representation we used the genome assembly GCF_001683475.1 and NC_034949.1. 439 Due to absence of available mitochondrial genome assembly for C. quinoa we created a custom 440 database containing gene sequences of mitochondrial origin retrieved from the total RNA sequencing 441 by comparing with Arabidopsis thaliana mitochondrial genome (NC_037304.1). The relative 442 frequency of each nucleotide at 5' N-terminal position of the small RNAs was calculated using Galaxy 443 tools (82). 444

We performed an analysis for clustered organellar sRNA using sRNA miner, implemented in 445 R/Bioconductor (83) using as parameter 40 for the minimal reads for end and 0.85 for the sharpness of 446 end. 447

Phylogenetic analysis. Predicted ORF encoding for RdRp were used to build an alignment 448 using MUSCLE algorithm implemented in MEGA 6 (84). Phylogenetic tree was built using the 449 maximum likelihood method, aligned protein sequences were used to estimate the best substitution 450 rate and parameter with MEGA 6. Substitution pattern and rates were estimated under the model 451 designed by Dimmic (+Gamma +Invar +Freq) (85). One thousand bootstrap replicates were 452 performed and branches with bootstrap value under 50 have been collapsed. A list of the accession 453 numbers of the viruses contained in the tree is shown in supplementary table 2. 454

Mitochondrial and Chloroplast enrichment protocol and quantitative evaluation of 455 CqMV1 and marker genes. Chloroplast and mitochondria enriched fractions were obtained by a 456 modified protocol already used for cucumber plants (86, 87). Ten grams of C. quinoa leaves were 457 homogenized in the ratio 1:10 in chilled extraction buffer (Sucrose 0.45 M, MOPS 15mM, EGTA 458 1.5mM, pH 7.4 with KOH) added with PVP 0.6%, BSA 0.2%, DTT 10mM and PMSF 0.2mM. After 459 filtration on Miracloth (Calbiochem) the homogenate was centrifuged at 2000g for 5 min to separate 460 on February 4, 2019 by guest<http://jvi.asm.org/>Downloaded from Nerva et al. J. Virol.

chloroplasts and cell debris. Pellet was used for the subsequent chloroplast purification; instead, the 461 supernatant was again centrifuged at 13000g for 30 min to obtain the mitochondrial fraction in the 462 pellet. From this point, the protocol followed two different ways. 463

Crude chloroplasts pellet were suspended in 1 ml of sorbitol resuspension buffer (SRM, 464 Sorbitol 1.65M, Hepes 250 mM pH8 with KOH) and layered on a Percoll® (Sigma-Aldrich) gradient 465 (35% buffer and 80% in SRM) and centrifuged at 2600g in a swing out rotor (SW41 Beckman) for 10 466 min. After centrifugation, the fraction with chloroplast was collected with a Pasteur pipette, diluted in 467 30 ml of SRM buffer and centrifuged at 2000g for 5 min, to remove all the Percoll. This step was 468 repeated for two times. Pellet was then suspended in resuspension buffer (RB, Mannitol 0.4 M, Mops 469 10 mM, EGTA 1 mM, pH 7.2) added with PMSF 0.2 mM, and checked on the microscope to evaluate 470 the purity of the preparation. Crude mitochondria pellet was resuspended in 1 ml of washing buffer 471 (WB; sucrose 0.6 M, Mops 20 mM, EGTA 2 mM, pH 7.2 with KOH) added with PMSF 0.2 mM, 472 layered on a Percoll gradient (18%, 23% and 40% in WB) and centrifuged at 12000g in SW21 473 (Beckmann) for 45 min. Mitochondrial fraction, between the 23% and 40% interface, was collected 474 with a Pasteur pipette and two wash-step were performed as already described for chloroplasts in WB. 475 We resuspended pellet in about 0.1 ml and checked the quality of purification by observation with 476 fluorescent microscope. Mitochondrial and chloroplast enriched fractions were stored at -80°C until 477 use for RNA extraction and qRT-PCR analysis. In order to check which fraction contained virus 478 genome enrichments, RNA was extracted using the Spectrum™ plant total RNA kit (Sigma-Aldrich, 479 Saint Louis, MO, USA) from preparations representing normalized amounts of each fraction. 480

Complementary DNA (cDNA) was synthesized using the High Capacity cDNA Reverse Transcription 481 Kit (Applied Biosystems, Foster city, CA, USA). We then tested the presence of virus genomic RNA 482 and mRNA corresponding to a number of marker genes with quantitative real time PCR using a 483 CFX96™ apparatus (Biorad, Hercules, CA, USA) and iTaq™ Universal Probes and iTaq™ Universal 484 SYBR Supermixes (Biorad) following protocols previously described (88). The marker genes 485 corresponded to sequences of the Cytochrome P450 oxydase (ANY30855.1); the S2 ribosomal 486

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protein, present in the mitochondria but encoded by the nuclear genome (89); the S3 ribosomal 487 protein, encoded by plant mitochondrial genomes (90); the ORF-X protein, also encoded by 488 mitochondrial genomes (91); C. quinoa sequences related to the latter three genes were retrieved from 489 our RNAseq database. Oligonucleotides used for qRT-PCR are displayed in Supplementary on line 490 table 3. 491

Northern blot analysis. For Northern blot analyses, total RNA from leaves of different age 492
was prepared using Total Spectrum RNA Reagent (Sigma-Aldrich) as suggested by the manufacturer. 493
RNA samples were separated in denaturing conditions (glyoxal method) as detailed, using HEPES-494
EDTA buffer (92). Radio-labeled probes were prepared from linearized plasmid containing the cDNA 495
clones (producing probes in both orientation) through T7 transcription using the Maxiscript T7 kit 496
reagents (Thermo Fisher Scientific Inc., Waltham, MA, USA) as suggested by manufacturer. 497
qRT-PCR fast screening method. For a fast screening of viral infection we applied 498
modifications of a simple qRT-PCR protocol that uses crude extracts as template (93). We placed 30 499
seeds to germinate in 90 mm diameter Petri dishes with wet filter paper and let it germinate for 3 days 500
(or single plantlets in the case of seed transmission assay). Plantlets of all the accession tested were 501
placed in extraction bags (Bioreba, Reinach, Switzerland) and diluted 1:20 (w/v) with carbonate buffer 502
pH 9.6 (94) added with 2% PVP40, 0.2% BSA, 1% sodium metabisulfite and 0.05% tween 20. Raw 503
extract has been diluted 1:10 in sterile water and boiled 10 min at 95 °C. qRT-PCR screening was 504
performed using a CFX96™ Real-Time PCR Detection System (Biorad), PCR mix was prepared with 505
iTaq™ Universal Probes Supermix (Biorad) adding 3 U of reverse transcriptase from High-Capacity 506
RNA-to-cDNA Kit (Thermo Fisher Scientific) for each sample. Reactions have been performed in 10 507
µl of total volume adding 1 µl of boiled extract to 9 µl of PCR mix. The qRT-PCR protocol has a 30 508
min step at 37°C to perform the reverse transcription of the viral genome, then is followed by 1 min at 509
94 °C and 40 steps of denaturation at 95 °C for 10 sec, annealing and extension at 60 °C for 30 sec. 510
Data availability. The GenBank/eMBL/DDBJ accession numbers of the sequences reported 511
in this paper is MF375475. 512

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FIGURE LEGENDS 724

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FIG 1 *Chenopodium quinoa* mitovirus 1 (CqMV1) genome organization and its replicative nature. (A) genome 726
organization of the single positive strand genome with the RdRp ORF (blue box). The position of the two DNA segments 727
amplified by reverse transcriptase PCR and used as probes in northern hybridization experiments are shown as black lines 728
(Probe 1 and Probe 2). nt= nucleotide position on the genome. (B) Ethidium bromide stained TAE gel (1%) to separate 729
PCR products from RNA template and DNA template extracted from *Chenopodium quinoa* IPSP1. M= molecular weight 730
marker; RT= reverse transcriptase; no-RT= RNA template without reverse transcriptase. Kb=kilobases. 731

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FIG 2 *Chenopodium quinoa* mitovirus 1 phylogenetic placement. Predicted ORF encoding RdRp were used to build an 733
alignment using MUSCLE implemented in MEGA 6 (Tamura et al. 2013). Phylogenetic tree was built using the maximum 734
likelihood method with 1000 bootstrap replicates. Branches with bootstrap value <50 have been collapsed. The analysis 735
involved 125 amino acid sequences. All positions with less than 90% site coverage were eliminated. There were a total of 736
457 positions in the final dataset. A list of the accession numbers of the viruses contained in the tree is shown in 737
supplementary on line Table 2. The diamond symbol represents a node that was collapsed, which includes 27 RdRp 738
sequences from a number of invertebrate and plant ourmiaviruses, still awaiting a taxonomical classification. Asterisks 739
indicate endogenized plant mitochondrial sequences. 740

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FIG 3 *Chenopodium quinoa* mitovirus 1 (CqMV1) genomic RNA is enriched in mitochondrial fractions. (A) Fluorescent 742
microscopy observation of mitochondria (a-c) and chloroplasts (d-f) fractions purified from *Chenopodium quinoa* leaves. 743
a, d: bright-field; b, e: chlorophyll fluorescence; c, f: overlay of the two images. Magnification bar: 50 μ m. (B) Real time 744
quantification of RNA corresponding to CqMV1 and two nuclear (Cox and S2) and two mitochondrial genes (OrfX and 745
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FIG 4 Evidence of minus strand genomic RNA accumulation. Time of autoradiography exposure is indicated at the 749
bottom of each panel (exp); RNA samples were extracted from leaves from uninfected *Chenopodium quinoa* accession 750
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In each panel, negative sense probe reacting with the viral positive RNA strand is shown on the left and positive strand 752
probe reacting with the viral negative RNA strand is shown on the right. Asterisks show weak specific signals given by 753
positive probes targeting the viral negative RNA strand. (A) panel shows signal from the two orientations of Probe 1 (see 754
Fig. 1), (B) panel shows signal given by Probe 2 (see Fig. 1). Lower panels are methylene blue stained membranes 755
showing ribosomal RNAs loading (rRNA). 756

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FIG 5 Differential symptom severity of virus infections: local symptoms in leaves. Two *Chenopodium quinoa* mitovirus 1 (CqMV1) infected accessions (cv Regalona and IPSP1) and two CqMV1-free accessions (BO25 and BO75) were used to assess their responses when inoculated with pathogenic viruses. Observations were done at 7 days post inoculation (dpi). In panel A, alfalfa mosaic virus (AMV) locally inoculated leaves did not show differences in term of symptoms severity between CqMV1-infected and CqMV1-free accessions. In panel B, infection with Lettuce mosaic virus (LMV) revealed differences between CqMV1-infected accessions, in which chlorotic lesions were observed, and CqMV1-free accessions, in which necrotic lesion (red arrows) were observed. In panel C, infection with Hibiscus latent ringspot virus (HLRSV) did not revealed symptom differences among the four accessions.

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FIG 6 Differential symptom severity of virus infections: systemic symptoms. Two *Chenopodium quinoa* mitovirus 1 (CqMV1) infected accessions and two CqMV1-free accessions were used to assess their responses when inoculated with three pathogenic viruses. Observations were done at 14 days post inoculation. In vertical rows the virus species used in the experiments: alfalfa mosaic virus (AMV) A,D,G,J; Lettuce mosaic virus (LMV) B,E,H,K; Hibiscus latent ring spot virus (HLRSV) C,F,I,L. In horizontal rows the four *Chenopodium quinoa* accessions BO78, BO25, IPSP1, and Regalona (Regal.) are reported. A negative mock-inoculated plant of the same age is present next to two infected plants in each panel. All accessions shows systemic symptoms of mild growth impairment, malformation and mild mottling. CqMV1-free accessions BO78 (A,B,C) and BO25 (D,E,F), showed red-violet pigmentation on stems (white arrows) whereas CqMV1 infected accessions IPSP1 (G,H,I) and Regalona (J,K,L) and mock inoculated plants did not showed any pigmentation. Inset of a pigmented stem from accession BO78 infected by LMV (panel B) is enlarged in panel M, whereas inset of a stem of cultivar Regalona also infected by LMV (panel K) is enlarged in panel N.

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FIG 7 Small RNA (sRNA) length distribution in virus infected *Chenopodium quinoa* leaves. Reads from small RNA sequencing were mapped against genes encoded by chloroplast, nucleus, mitochondrion, alfalfa mosaic virus (AMV) and *Chenopodium quinoa* mitovirus 1 (CqMV1) genomes. Abundance is expressed as percentage of reads of a particular length and arrows above bars indicates the most abundant sRNA length inside the specific gene set. CqMV1 shared the same sRNA pattern distribution of genes encoded by the mitochondrial genomes, suggesting a mitochondrial localization and a specific but still uncharacterized RNA-degradation pathway inside the mitochondria.

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Table 1. Small RNA reads mapping against each of the corresponding genomes as a percentage of total reads in *Chenopodium quinoa* cv Regalona, cv. Regalona infected with AMV and BO78 infected with AMV.

Table 1. Small RNA reads mapping against each of the corresponding genomes as a percentage of total reads in *Chenopodium quinoa* cv Regalona, cv. Regalona infected with AMV and BO78 infected with AMV.

	Chloroplast	CqMV1	Nucleus	AMV	Mitochondria	Total
Regalona	14.017	0.006	84.071	0.000	1.906	4119721
Regalona AMV	14.783	0.111	78.580	4.583	1.945	7240581
BO78 AMV	15.262	0.000	80.333	2.489	1.916	3758683

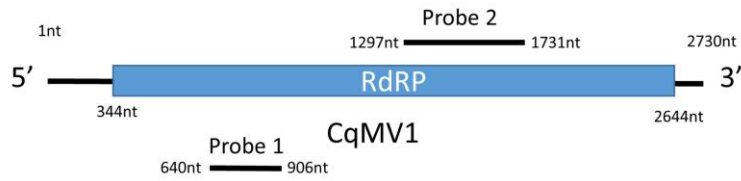
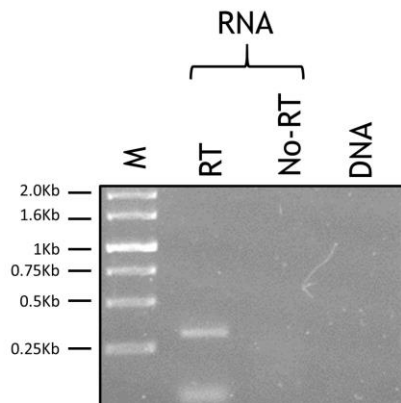
A**B**

FIG 1 *Chenopodium quinoa* mitovirus 1 (CqMV1) genome organization and its replicative nature. (A) genome organization of the single positive strand genome with the RdRp ORF (blue box). The position of the two DNA segments amplified by reverse transcriptase PCR and used as probes in northern hybridization experiments are shown as black lines (Probe 1 and Probe 2). nt= nucleotide position on the genome. (B) Ethidium bromide stained TAE gel (1%) to separate PCR products from RNA template and DNA template extracted from *Chenopodium quinoa* IPSP1. M= molecular weight marker; RT= reverse transcriptase; no-RT= RNA template without reverse transcriptase. Kb=kilobases.

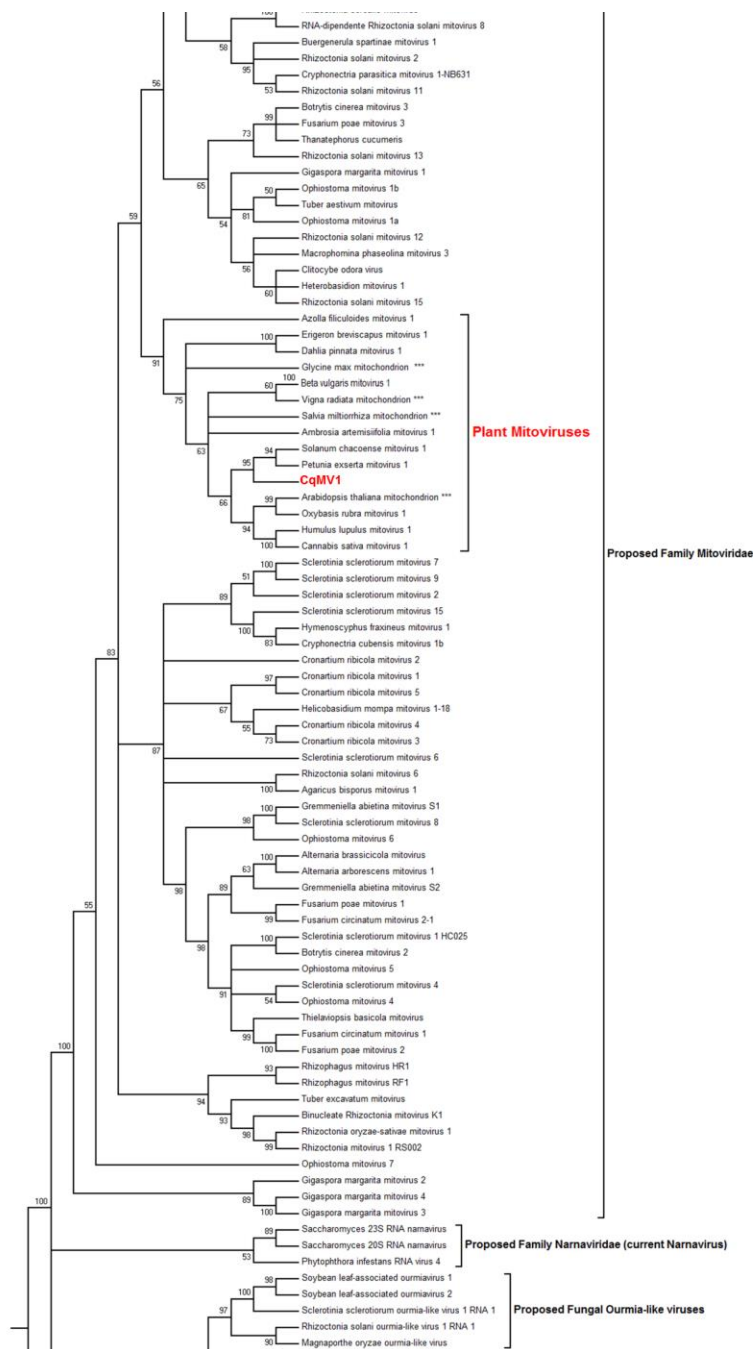


FIG 2 *Chenopodium quinoa* mitovirus 1 phylogenetic placement. Predicted ORF encoding RdRp were used to build an 733 alignment using MUSCLE implemented in MEGA 6 (Tamura et al. 2013). Phylogenetic tree was built using the maximum likelihood method with 1000 bootstrap replicates. Branches with bootstrap value <50 have been collapsed. The analysis involved 125 amino acid sequences. All positions with less than 90% site coverage were eliminated. There were a total of 736 457 positions in the final dataset. A list of the accession numbers of the viruses contained in the tree is shown in 737 supplementary on line Table 2. The diamond symbol represents a node that was collapsed, which includes 27 RdRp 738 sequences from a number of invertebrate and plant ourmiaviruses, still awaiting a taxonomical classification. Asterisks 739 indicate endogenized plant mitochondrial sequences.

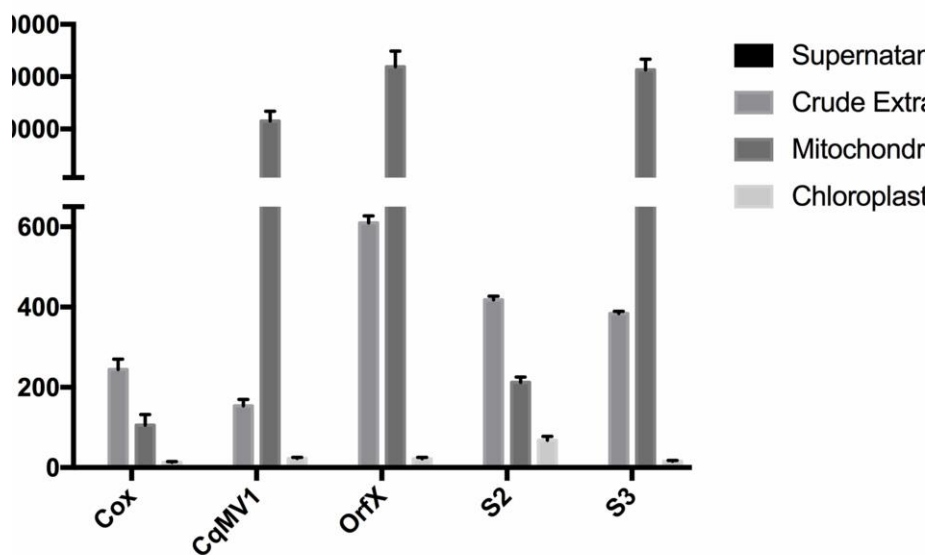
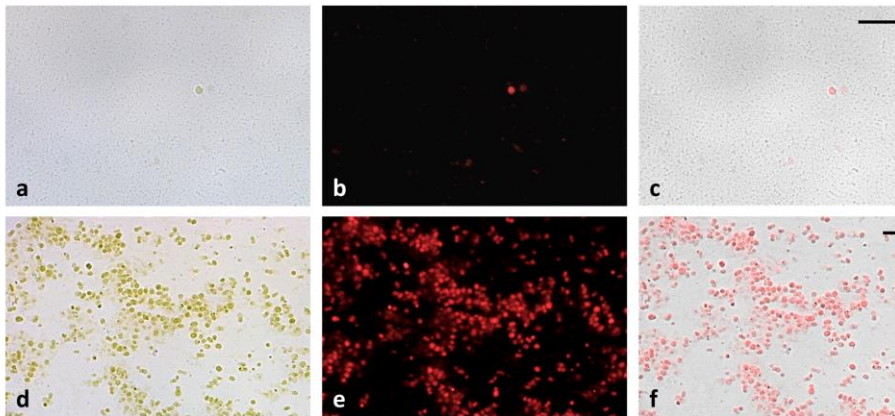


FIG 3 *Chenopodium quinoa* mitovirus 1 (CqMV1) genomic RNA is enriched in mitochondrial fractions. (A) Fluorescent 742 microscopy observation of mitochondria (a-c) and chloroplasts (d-f) fractions purified from *Chenopodium quinoa* leaves. 743 a, d: bright-field; b, e: chlorophyll fluorescence; c, f: overlay of the two images. Magnification bar: 50 μ m. (B) Real time 744 quantification of RNA corresponding to CqMV1 and two nuclear (Cox and S2) and two mitochondrial genes (OrfX and 745 S3). All quantifications are relative to the amount of virus or mRNA present in the Supernatar of the 14K centrifuge run: 746 such amount was arbitrarily established as 1. Error bars represent standard error of the mean (n=3).

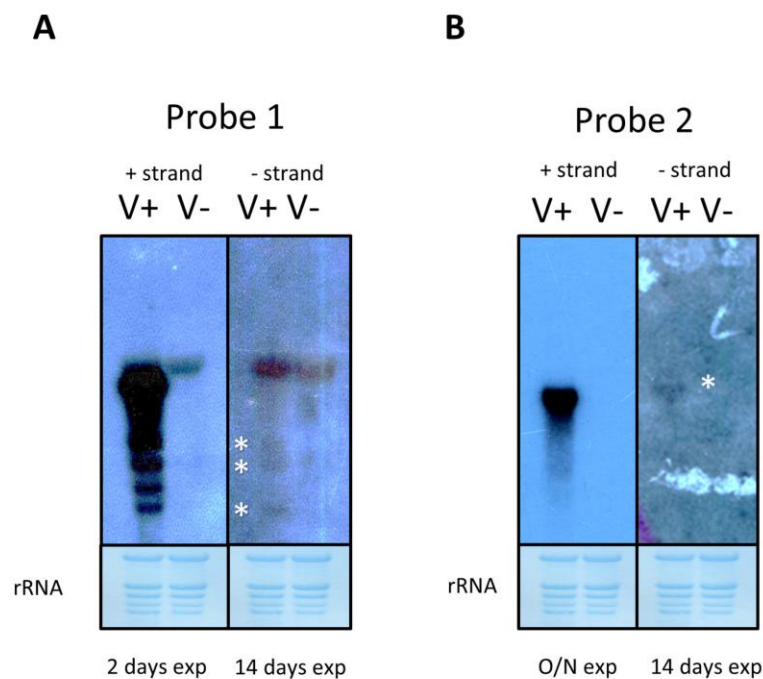


FIG 4 Evidence of minus strand genomic RNA accumulation. Time of autoradiography exposure is indicated at the 749 bottom of each panel (exp); RNA samples were extracted from leaves from uninfected *Chenopodium quinoa* accession 750 BO25 (V-) and from *Chenopodium quinoa* mitovirus 1 (CqMV1)-infected *Chenopodium quinoa* cultivar Regalona (V+). 751 In each panel, negative sense probe reacting with the viral positive RNA strand is shown on the left and positive strand 752 probe reacting with the viral negative RNA strand is shown on the right. Asterisks show weak specific signals given by 753 positive probes targeting the viral negative RNA strand. (A) panel shows signal from the two orientations of Probe 1 (see 754 Fig. 1), (B) panel shows signal given by Probe 2 (see Fig. 1). Lower panels are methylene blue stained membranes 755 showing ribosomal RNAs loading (rRNA).

Fig. 5

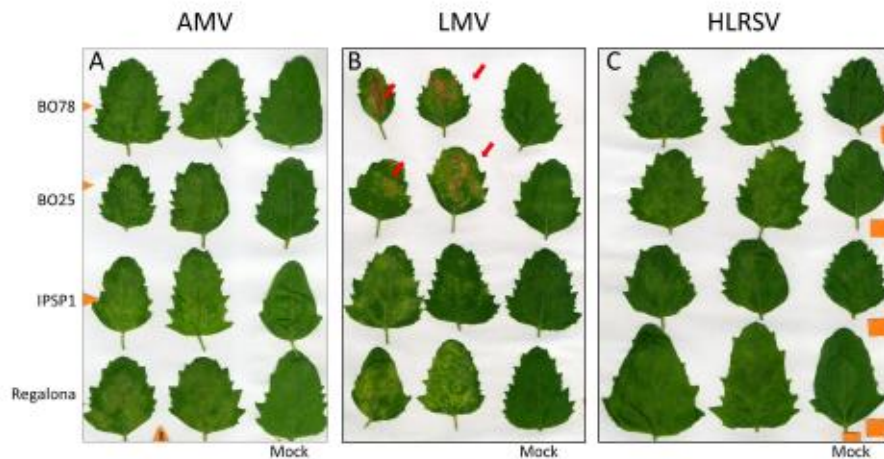


FIG 5 Differential symptom severity of virus infections: local symptoms in leaves. Two *Chenopodium quinoa* mitovirus 1 758 (CqMV1) infected accessions (cv Regalona and IPSP1) and two CqMV1-free accessions (BO25 and BO75) were used to 759 assess their responses when inoculated with pathogenic viruses. Observations were done at 7 days post inoculation (dpi). 760 In panel A, alfalfa mosaic virus (AMV) locally inoculated leaves did not show differences in term of symptoms severity 761 between CqMV1-infected and CqMV1-free accessions. In panel B, infection with Lettuce mosaic virus (LMV) revealed 762 differences between CqMV1-infected accessions, in which chlorotic lesions were observed, and CqMV1-free accessions, 763 in which necrotic lesion (red arrows) were observed. In panel C, infection with Hibiscus latent ringspot virus (HLRSV) did 764 not revealed symptom differences among the four accessions

Figure 6

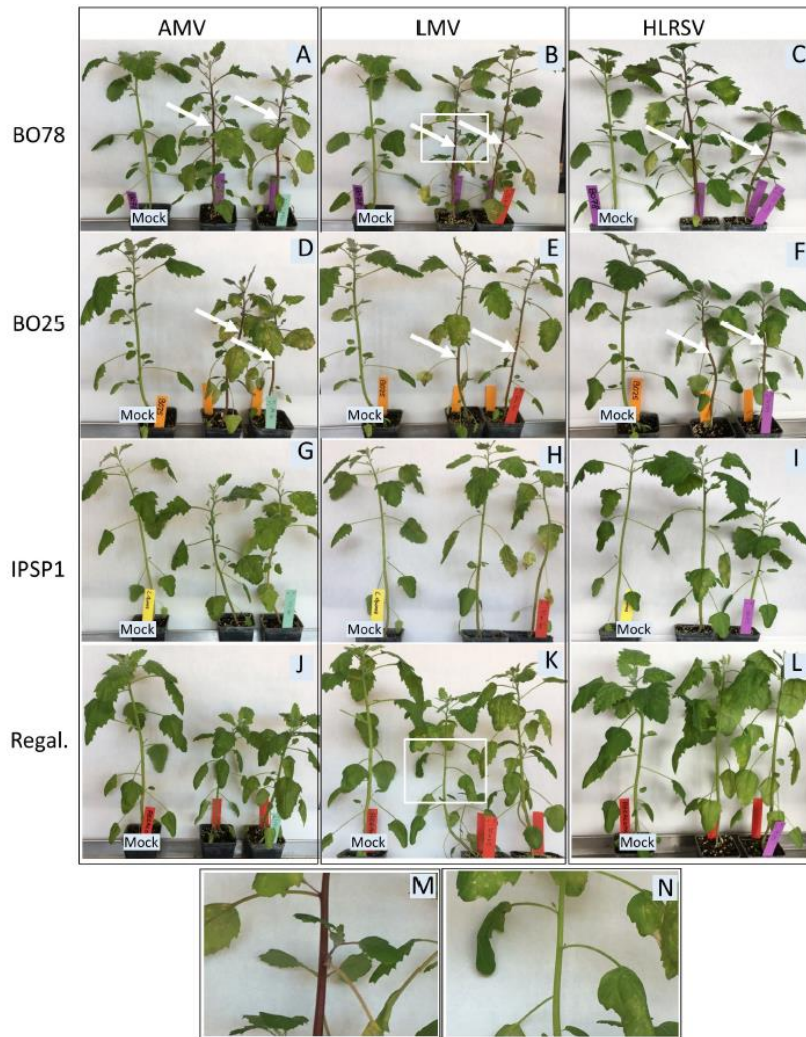


FIG 6 Differential symptom severity of virus infections: systemic symptoms. Two *Chenopodium quinoa* mitovirus 1 767 (CqMV1) infected accessions and two CqMV1-free accessions were used to assess their responses when inoculated with 768 three pathogenic viruses. Observations were done at 14 days post inoculation. In vertical rows the virus species used in the 769 experiments: alfalfa mosaic virus (AMV) A,D,G,J; Lettuce mosaic virus (LMV) B,E,H,K; Hibiscus latent ring spot virus 770 (HLRSV) C,F,I,L. In horizontal rows the four *Chenopodium quinoa* accessions BO78, BO25, IPSP1, and Ragalona 771 (Regal.) are reported. A negative mock-inoculated plant of the same age is present next to two infected plants in each 772 panel. All accessions shows systemic symptoms of mild growth impairment, malformation and mild mottling. CqMV1-773 free accessions BO78 (A,B,C) and BO25 (D,E,F), showed red-violet pigmentation on stems (white arrows) whereas 774 CqMV1 infected accessions IPSP1 (G,H,I) and Regalona (J,K,L) and mock inoculated plants did not showed any 775 pigmentation. Inset of a pigmented stem from accession BO78 infected by LMV (panel B) is enlarged in panel M, whereas 776 inset of a stem of cultivar Regalona also infected by LMV (panel K) is enlarged in panel N.

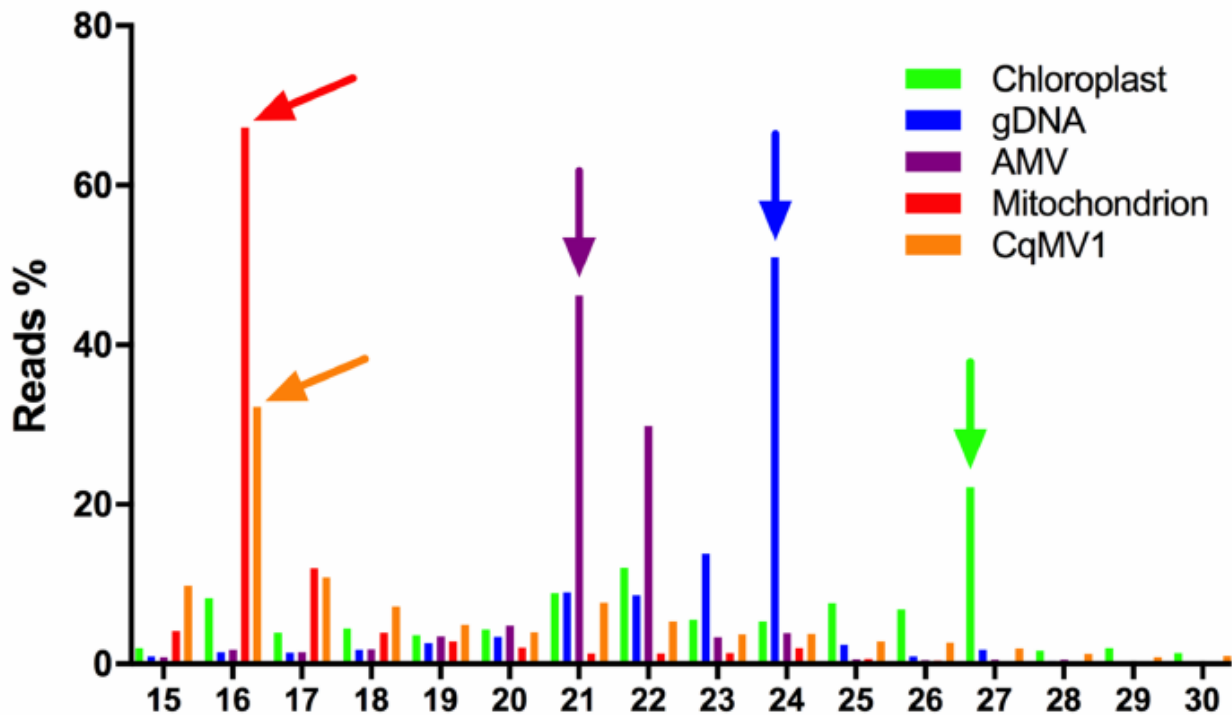


FIG 7 Small RNA (sRNA) length distribution in virus infected *Chenopodium quinoa* leaves. Reads from small RNA 779 (sRNA) sequencing were mapped against genes encoded by chloroplast, nucleus, mitochondrion, alfalfa mosaic virus 780 (AMV) and *Chenopodium quinoa* mitovirus 1 (CqMV1) genomes. Abundance is expressed as percentage of reads of a 781 particular length and arrows above bars indicates the most abundant sRNA length inside the specific gene set. CqMV1 782 shared the same sRNA pattern distribution of genes encoded by the mitochondrial genomes, suggesting a mitochondrial 783 localization and a specific but still uncharacterized RNA-degradation pathway inside the mitochondria.