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MUTANT SWARMS OF A TOTIVIRUS-LIKE ENTITY ARE PRESENT IN THE
RED MACROALGA *CHONDRUS CRISPUS* AND HAVE BEEN PARTIALLY
TRANSFERRED TO THE NUCLEAR GENOME

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ABSTRACT

Chondrus crispus Stackhouse (Gigartinales) is a red seaweed found on North
Atlantic rocky shores. Electrophoresis of RNA extracts showed a prominent band
with a size of around 6,000 bp. Sequencing of the band revealed several sequences
with similarity to totiviruses, double-stranded RNA viruses that normally infect
fungi. This virus-like entity was named CcV. It should probably be regarded as an
extreme viral quasispecies or a mutant swarm since low identity (<65%) was found
between sequences. Totiviruses typically code for two genes: one capsid gene (*gag*)
and one RNA-dependent RNA polymerase gene (*pol*) with a pseudoknot structure

between the genes. Both the genes and the intergenic structures were found in the CcV sequences. A non-identical *gag* gene was also found in the nuclear genome of *C. crispus*, with associated EST and upstream regulatory features. The gene was presumably horizontally transferred from the virus to the alga. Similar dsRNA bands were seen in extracts from different life cycle stages of *C. crispus* and from all geographical locations tested. In addition, similar bands were also observed in RNA extractions from other red algae; however, the significance of this apparently widespread phenomenon is unknown. No phenotype caused by the infection nor any virus particles, or capsid proteins were identified; thus, the presence of viral particles has not been validated. These findings increase the known host range of totiviruses to include marine red algae.

Key words: *Chondrus*, dsRNA, mutant swarm, Rhodophyta, totivirus, virus

INTRODUCTION

Viruses play a key role in the functioning of marine ecosystems (Fuhrman 1999, Suttle 2005, 2007). Their prevalence and diversity contribute to the control of the abundance, the diversity, and the productivity of many marine organisms and influence global nutrient and energy cycles. For example, it is estimated that viruses are responsible for the mortality of 20% of the oceans' biomass every day and that their abundance is 15 times higher than that of prokaryotes, even if the biomass is less. Because of the increasing understanding of the relevance of marine viruses a substantial research effort has been made to study their ecology, physiology, and molecular biology. Marine viruses have been shown to infect a multitude of various

organisms from cyanobacteria to whales. Viruses have also been suggested to represent the most genetically diverse biological entities in the ocean.

There are various types of viruses that differ in terms of their genetic material: single-stranded DNA (ssDNA), double-stranded DNA (dsDNA), single-stranded RNA (ssRNA), and double-stranded RNA (dsRNA). The sizes of the viral genomes vary between a couple of thousand to several million base pairs with RNA viruses normally having smaller genomes (Philippe et al. 2013). Plant viruses can be divided into four types based on their lifestyle: persistent, chronic, endogenous and acute. Infections with persistent plant viruses typically do not induce any phenotype and are transmitted via gametes. In plants, this type of lifestyle has hitherto been known exclusively from dsRNA viruses. Acute viruses can normally be transported between plants and typically produce a phenotype (Roossinck 2010).

One area where effects of viruses on marine organisms have been more intensely studied is the interaction between viruses and phytoplankton (e.g. Etten et al. 1991, Reisser 1993, Brussaard et al. 2004, Lang et al. 2009). It has been observed (Suttle 2007) that viruses have a key role in controlling the progress, and sometimes the collapse, of many phytoplankton blooms (e.g. *Emiliana huxleyi*, *Phaeocystis globosa* and *Heterosigma akashiwo*). Another system that has been studied in more detail is the ecological consequences of the interaction between the green unicellular alga *Ostreococcus* and its DNA virus (e.g. Derelle et al. 2008, 2015, Moreau et al. 2010). It has therefore been suggested that viral dynamics are one of the explanations to the paradox for the plankton (Fuhrman 1999); successful plankton species, due to their abundance, are more likely to succumb to virus-induced collapses of the populations than are rare species. This preserves the diversity of the phytoplankton.

We have, thus, some information on the interaction between unicellular algae and viruses as well as its importance.

Seaweeds are an important component of the coastal ecosystem; however, much less is known about viruses that infect seaweeds or macroalgae than their unicellular counterparts (Schroeder 2011). One exception is a virus that was found in the brown alga *Ectocarpus siliculosus*. This dsDNA virus, EsV-1, belonging to the Phaeoviruses with a genome of 335,593 bp, becomes associated to the host's genome (Müller et al. 1990, Klein et al. 1993, Delaroque et al. 2001, Cock et al. 2010). The virus particles are most expressed during reproduction and spread through infection of spores and gametes. It is manifested by malformed gametangia and lysis of swollen hyaline cells (Müller et al. 1990, 2000). Similar phaeoviruses have also been found in other brown algae, including, *Feldmannia irregularis*, *F. simplex*, (Müller and Frenzer 1993, Friess-Klebl et al. 1994), *Hincksia hincksiae* (Kapp et al. 1997), *Botrytella micromora* (as *Sorocarpus uvaeformis*) (Oliveira and Bisalputra 1978), *E. fasciculatus* (Sengco et al. 1996), *Streblonema* sp. (La Claire II and West 1977), and *Pylaiella littoralis* (Maier et al. 1998). Virus infections have also been observed in the green coenocytic macroalga *Bryopsis cinicola* (Koga et al. 1998, 2003). Here one 4.5 kbp sequence belonging to the dsRNA virus family Partitiviridae was found associated with mitochondria and for four others of about 2 kbp with chloroplasts (Ishihara et al. 1992).

Virus infections in red algae are much less studied than in brown algae; however, virus-like particles have been found, by electron microscopy-studies, in the cytoplasm of the freshwater red alga *Sirodotia tenuissima* (Lee 1971) and in the marine species *Acrochaetium* (*Audouinella*) *saviana* (Pueschel 1995), *Gracilaria epihippisor*a (Apt and Gibor 1991), and *G. verrucosa* (Tripodi and Beth 1976). In

addition, gall-like structures have been found in several species of *Bostrychia* including *B. kelanensis*, *B. radicata*, *B. simpliciuscula*, and *B. tenella*, and in some cases the galls have been associated with virus-like particles (West et al. 2013). For all of these possible red algal viruses the type of virus is unknown, no sequences are known, and the only reported phenotype is the formation of galls. Thus, the prevalence and importance of viruses in red algal biology is virtually unknown.

Chondrus crispus, or Irish moss, is a common, often intertidal red macroalga that has been promoted as a model species for florideophyte red algae (Collén et al. 2014). It has a triphasic life cycle with morphologically similar tetrasporophytes (diploid), male, and female gametophytes (haploid) and morphologically different carposporophytes (diploid) on the female gametophyte. The 105-Mbp genome was recently sequenced and codes for 9,606 genes (Collén et al. 2013). During the purification of RNA for the *Chondrus* genome project a prominent band of unusual size was seen on agarose gels; sequencing of this band showed sequence similarities to totiviruses, dsRNA viruses that normally infect fungi.

MATERIALS AND METHODS

Algal material and culture. Unless explicitly stated otherwise studies were carried out on a strain of the red alga *Chondrus crispus* Stackhouse (Gigartinales) grown in unialgal culture (Collén et al. 2013) at the *Station Biologique de Roscoff*, from a male gametophyte collected at Peggy's Cove, Nova Scotia, Canada (44°29'N, 63°55'W) in 1985 by Juan Correa. For the other specified *C. crispus* life cycle stages, samples were collected near Roscoff or in other specified locations.

120 *RNA extraction and purification. Chondrus crispus* RNA was extracted based on a
121 protocol adapted from Apt et al. (1995). Briefly, frozen tissue (~1 g) was ground
122 with a small amount of sand and liquid N₂ in a mortar and pestle. The powder was
123 transferred to a 50 ml Teflon tube, and extracted with 15 ml extraction buffer (100
124 mM Tris-HCl pH 7.5, 1.5 M NaCl, 50 mM EDTA, 50 mM DTT, 2 % CTAB) for 60
125 min, with continuous shaking at 200 rpm at room temperature. Cellular debris was
126 removed after centrifugation with one volume of chloroform-isoamyl alcohol
127 (24:1) at 10,000 g and 4 °C for 20 min. The aqueous phase was transferred to a
128 Teflon tube and polysaccharides were precipitated with 0.25 volumes of 100 %
129 ethanol and centrifuged with one volume of chloroform-isoamyl alcohol (24:1) at
130 10,000 g and 4 °C for 20 min. Total RNA was precipitated overnight with 3 M LiCl
131 and 1 % of β-mercaptoethanol; after centrifugation, the pellet was dissolved in TE
132 buffer (10 mM Tris-HCl pH 7.5, 1 mM EDTA). RNA quality was assessed on 1.5 %
133 agarose gel stained with 0.1 µg/ml ethidium bromide (EtBr) and quantified using a
134 NanoDrop ND-2000 spectrophotometer (NanoDrop products, Wilmington, DE,
135 USA). CsCl separation of total nucleic acids was used to purify dsRNA in 2 ml
136 polyallomer tubes with 0.95 g/ml CsCl in TE and 50 µl of 10 mg/ml EtBr solution.
137 The solution was centrifuged for 24 h at 90,000 rpm. The dsRNA band was
138 recovered and the EtBr was extracted with TE-saturated butanol. The remaining
139 aqueous phase containing the dsRNA was diluted three times and isopropanol-
140 precipitated over-night at -20 °C. After centrifugation for 45 min at 18,000 g and 4
141 °C, the pellet was washed with 200 µL 80 % ethanol and recentrifuged for 30 min at
142 18,000 g and 4 °C; the pellet was finally dissolved in 100 µL of TE and kept at -20
143 °C until use.

Cloning and sequencing. Cloning of RNAs was adapted from Djikeng et al. (2008) using random-tagged primers and RNA purified from agarose gels. Briefly, 100 ng of RNA and 2 pmol of random tagged primers were denaturated for 5 min at 70 °C. The first cDNA strand was synthesized using the Improm-RT polymerase Kit (Promega), after annealing 5 min at 25 °C, elongation 1 h at 38 °C, and inactivation 15 min at 70 °C the RNA strand was degraded with 2 units of RNase H, 20 min at 37 °C. The second strand was generated using 2 pmol of random tagged primer and 2.5 units of Klenow. After purification cDNAs were amplified by PCR and cloned in TOPO-TA (Invitrogen). Race PCR was performed to obtain longer sequences using 75 ng of gel-purified viral RNA with 1.2 µM specific primers, drawn from a first sequence alignment in the polymerase region, and 1.2 µM universal primers. Reverse transcription was carried out at 42 °C for 90 min, using SmartScribe reverse transcriptase Protocol (Clontech). After a 1/12.5 dilution, 1 µl of cDNA was amplified in a total volume of 25 µl, with 400 nM of each specific and universal primers, 0.2 mM dNTP, Pfu buffer, using 1.25 units Pfu DNA polymerase (Invitrogen). After a denaturation step for 4 min at 94 °C, samples were cycled in a ABI2700 thermocycler as following: 30 sec at 94 °C, 30 sec at 60 °C, 6 min at 72 °C for 5 cycles followed by 5 cycles of 30 sec at 94 °C, 30 sec at 56 °C, 6 min at 72 °C and 25 cycles of 30 sec at 94 °C, 30 sec at 52 or 56 °C, depending on the considered specific primer, 6 min at 72 °C and a final step of 10 min at 72 °C. PCR products (2 µL) were directly cloned with TOPO-TA cloning (Invitrogen) and transformants were screened for size by PCR and sequenced using universal primers M13Forward and M13Reverse. Tail-PCR experiments were performed using combined random and specific primers; 200 ng RNA were reverse transcribed using 200 nM GSP1 and 2 µM random primer RP1, using one step Access quick RT-PCR 2-fold master Mix

(Promega) in a 20 µl total volume, according to the manufacturer specifications. The second step included 2.5 µl of the first 50-fold diluted amplified sample, 2 µM of RP1 and 200 nM internal GSP2, 3 mM MgCl₂, 0.2 mM dNTP, Go-Taq buffer and 1 unit Go-Taq polymerase (Promega). The third step is consisted in the same protocol as the second step, including a 50-fold dilution of the sample and 2 µM of RP1 and 200 nM of internal GSP3.

The cloning and sequencing effort generated only small sequence fragments in the capsid part and in the polymerase sequence region; in addition, race or tail-PCR never permitted amplification of the intergenic part, which can be described as a complex pseudoknot structured sequence. Because we demonstrated that at least seven different sequences could be cloned, with regard to the partial sequences motif obtained, we decided to try to extend some of the contigs obtained; using specific primers and one-step RT-PCR cross amplification between ends of the capsid part and the RdRp part to test whether extremities were from the same virus fragment or not. We started from 30 ng of CsCl purified dsRNA, denaturated for 4 min at 99 °C, reverse transcribed for 45 min at 45 °C and the PCR cycling parameters were programmed as follows: 4 min at 95 °C, and 32 cycles of 30 sec at 95 °C, 30 sec at 54 °C, 3 min at 72 °C in a 2700 ABI thermocycler. Sequencing was performed using capillary electrophoresis on an ABI prism 3100 sequencer (Applied Biosystems, Foster City, CA, USA) with 50-cm capillaries.

DNase and RNase essays. To discriminate between DNA and RNA, we compared DNase and RNase digestion on total nucleic acid extracts. We subjected 1.5 µg of total nucleic acid to 2 units of Turbo DNaseI (Ambion) or 1 unit of RNaseA (Macherey Nagel) for 1h at 40 °C. After precipitation, the samples were loaded on 1 % agarose gel in TAE stained with EtBr.

194

195 **RESULT AND DISCUSSION**

196 Native agarose gel electrophoresis of RNA extracts from *Chondrus crispus* showed,
197 in addition to rRNA and mRNA (and some DNA contamination), one prominent
198 band with a size of approximately 6,000 bp. This band was not affected by DNase
199 treatment but disappeared after treatment with RNase (Figure 1A). The bands were
200 much less visible after denaturation of the RNA extract on a denatured gel (Figure
201 1B) than on a native gel. This sensitivity to RNase indicated that the bands were
202 made of RNA and likely double-stranded since the intensity was greatly diminished
203 on a denatured gel. This is caused by the stronger affinity of the ethidium bromide
204 stain to double-stranded RNA than single-stranded RNA (Sambrook and Russell
205 2006).

206 The probably double-stranded RNA band was purified, reverse transcribed using
207 random priming, and partially sequenced. Several sequences were attained which
208 showed very high sequence diversity and could, thus, not be assembled in a
209 continuous sequence. Several fragments were found to be similar to
210 *Xanthophyllomyces dendrorhous* virus (XdV) (Figure 2) and this sequence was used
211 to align the sequences from *C. crispus*. XdV is a double-stranded RNA virus,
212 belonging to the totivirus (Baeza et al. 2012). The genome of totiviruses, typically
213 with a genome size 4.6-6.3 kbp, normally codes for two proteins, one capsid (*gag*)
214 gene and one RNA-dependent RNA polymerase (*pol*) gene with partially over-
215 lapping open reading frames (ORFs). The sequences from the tentative *C. crispus*
216 virus (CcV), contained partial sequences from both genes (Figure 3 and 4).
217 Considerable efforts were made to try to obtain a full sequence, but each approach,

including SISPA method, race- or tail PCR, with degenerated, specific, or random PCR primers gave sequences that corresponded to new viral sequences instead of one specific sequence. This indicated a considerable sequence diversity within the CcV. It should be noted that the host strain of *C. crispus* used was artificially forced through a bottleneck of reduced size to reduce the possibility of the strain being a chimera before the genome sequencing (Collén et al. 2013) and thereafter grew vegetatively. The origin of the strain represents milligrams of tissue so that no nuclear sequence diversity was detected during genome sequencing.

The high error rate of viral replication with an RNA-dependent RNA polymerase compared to a DNA polymerase generates sequence diversity within viral populations that is higher than in cellular organisms. This causes the formation of quasispecies or mutant swarms (Andino & Domingo 2015). The CcV seems to be an extreme example of this with very high sequence diversity, represented by the 20-65% amino acid identity found between sequences (Table 1). Compared with other dsRNA viruses, such as the polio virus with two nucleotide mutations per genome or less than 0.03% difference in nucleotidic sequences (Vignuzzi et al. 2006) and a similarly low diversity in the St. Louis encephalitis virus (Ciota et al. 2011). The genetic diversity of CcV is thus several orders of magnitude higher. One explanation for this higher sequence diversity could be the absence (or rarity) of horizontal spreading of totiviruses compared to the polio and encephalitis viruses. To our knowledge, the sequence diversity of totiviruses within one host has only been reported as different viruses e.g. the *X. dendrorhous* virus XdVL1 and XdVL2 (Baeza et al. 2012) which are less divergent than the CcV sequences. It should be noted that despite the large sequence variability only one distinct band was seen, implying sequence diversity rather than size diversity and also that size is conserved

even though the sequence is not. In addition, the acquired sequences typically coded for open reading frames. The CcV system is therefore a potentially interesting system to study the biology of quasispecies and a model to study viral evolution with previously unknown features.

Compared to totiviruses from, for example, *X. dendrorhous*, *Saccharomyces cerevisiae* and *Tuber aestivum*, the major conserved amino acid residues described in the consensus sequence of the totivirus *pol* gene (Baeza et al. 2012) are present in the CcV sequence (Figure 3). This is also true for domains 5 and 6, which have been demonstrated to be essential for viral function (Ribas 1992, Routhier 1998). Together, this strongly indicates that functional dsRNA from a totivirus-like entity is present in *C. crispus*.

The totiviruses belong to the mycoviruses which are widespread in yeasts and filamentous fungi but have not been reported from algae. Typically, they do not possess extracellular infections modes, are not associated with obvious disease symptoms, and do not exhibit harmful effects (Ghabrial and Suzuki, 2009). Only a few reports of phenotype modifications like hypovirulence or antifungal activity have been described (Pearson et al, 2009, Magliani 1997). Moreover, natural vectors are so far unknown (Ghabrial and Suzuki, 2009). It has been suggested that the dominant mode of diversification in most totiviruses is codivergence, parallel diversification between host and parasite (Göker et al, 2011). The large sequence diversity found in CcV indicates that other phenomena are probably occurring in the interaction between *C. crispus* and CcV.

Viruses from the totiviridae family are mainly found in fungi and protozoa. Penaeid shrimp infectious myonecrosis virus (IMNV), isolated from the Pacific white shrimp,

was the first totivirus described infecting a host other than a fungus or a protozoan (Nibert, 2007). Another virus, DTV, genetically related to IMNV was found to infect a *Drosophila* cell line (Wu et al., 2010). Totiviruses have also been isolated from mosquitoes (*Armigeres subalbatus*) (Zhai et al., 2010, Isawa et al., 2011). The viruses isolated from mosquitoes are genetically similar to IMNV and DTV (Isawa et al., 2011). A totivirus, related to the Giardiavirus, has also been found in salmon. (Haugland et al., 2011). Together with our data presented here, this shows that the host range for totivirus is relatively large within the eukaryotes.

The totivirus genome organization is characterized by a non-segmented dsRNA, with two genes, *gag* and *pol*, encoding two overlapping proteins, a capsid protein (CP) and an RNA-dependent RNA polymerase (RdRp). In most totiviruses, the (+) strand viral transcript is flanked by a 5' and a 3'-UTR and directs the translation of the CP protein and a smaller CP-RdRp fusion protein via a -1 ribosomal frameshift. The ribosomal -1 frameshift site consists of a heptamer with the consensus motif XXXYYYYZ, (X represents any nucleotide, Y represents A or U, and Z represents A, C or U) preceded by a GC dinucleotide (Jacks et al., 1988; Bekaert et al., 2003). The heptamer is followed by a short region preceding a stable RNA secondary structure, such as a pseudoknot or hairpin. The translational efficiency has been demonstrated to be directly related to the stability of these structures (Bidou et al. 1997). Slippery sites can promote a low level of frameshifting and downstream pseudoknot structures stimulate this process (Giedroc et al. 2000). In the 166 bp overlapping part of the sequence of CcV2 a putative slippery site, GGGUUUU, is located 18 nucleotides upstream a pseudoknot type structure. The structure has an estimated free energy at -18.85 kcal/mol (Figure 4). Thus, the CcV genome is organized as described in other totiviruses with two overlapping ORF including a 5'-ORF coding a capsid protein

and a 3'-ORF coding an RNA-dependent RNA polymerase. This involves an ORF disruption with an internal stop codon and -1 ribosomal frameshift.

Viral sequences from various families are sometimes found integrated into the host organism's genome (Holmes 2011). A search using the CcV sequence in the sequenced *C. crispus* nuclear genome showed that one non-identical copy of the *gag* gene was present (Fig 4B). RNAseq data showed expression of this gene in *C. crispus* (not shown). Sequence analysis suggests that the gene also contains regulatory sequences (see below). This indicates that a transfer has occurred from the virus to the *C. crispus* genome. This type of transfer between endogenous viral elements from non-retroviral RNA viruses is known from insects (Chrochu et al. 2004), other animals (Katzourakis and Gifford 2010), and plants (Chiba et al. 2011). For the viral genes found in plant genomes they represented both single- and double-stranded viruses of various families. There is evidence that totiviral and partitiviral genes are transferred frequently into the nuclear genome of eukaryotes, and that some transferred genes have functions in the recipient genomes since totivirus-like sequences have been identified in, for example, *Medicago truncatula*, *Lotus japonica*, and *Populus trichocarpa*, where they have been matched to the *gag* gene of the *Vicia* cryptic virus M (Liu et al. 2010). ESTs from totiviruses have also been found in a cDNA library from the diatom *Phaeodactylum tricornutum* (Liu et al. 2012), but it is unknown if this was due to incorporation in the genome or caused by the presence of viral entities. Normally, after integration of viral gene sequences in a genome the genes start to degenerate (Zwart et al. 2014). Considering that the *gag* gene in the nuclear genome is full length and has EST support, this suggests that either the gene provides an advantage to *C. crispus* or that the integration is recent. It could be noted that transgenic maize expressing the totiviral protein KP4 is more

317 resistant to fungal infection (Allen et al. 2011) and thus, incorporation of viral genes
318 may have positive effects. No ESTs or presence in the genome were found for the
319 *pol* gene in *C. crispus*.

320 A close analysis of nuclear genome regions around the *gag* gene gave unexpected
321 results (Figure 6). Consensus sequences of prokaryote-type promoters were found in
322 the upstream part of the gene. This includes a sigma70 type promoter located -11 of
323 the TATAAA-box and -38 of the TTGAGA-box preceded by a CAAT-box located -
324 67 before a putative start codon. Using yeast mitochondrial codon table instead of the
325 eukaryote nuclear one results in a unique reading frame for the *cap* gene from the
326 methionine codon found downstream the TATA-box. Interestingly, 35 aa preceding
327 the putative *gag* start codon show similarities with a bacterial protein from *Simkania*
328 *nevegensis*. Furthermore, in the same reading frame 24 aa sequence homologous to
329 the cytochrome P450 (CYP) CYP808A2 (XP_005710778.1) from *C. crispus*.
330 Preceding the putative CYP gene are several putative regulatory boxes, three
331 successive adjacent E-Boxes, followed by a G-Box, and a GCC-Box.

332 An E-Box is a cis-acting element, a conserved DNA sequence involved in regulation
333 of gene expression linked to environmental and physiological signaling. These
334 elements serve as binding sites for transcriptional activators, and have been
335 demonstrated to be important for circadian gene expression (Giuliano et al., 1988,
336 Staiger et al, 1989, Williams et al., 1992). For example, in the green alga
337 *Chlamydomonas reinhardtii*, upregulation of the C3 subunit of the CHLAMY1
338 RNA-binding protein is mediated by an E-box element in its promoter; causing
339 temperature-dependent up-regulation of C3, its circadian expression, and its co-
340 regulation by C1 (Seitz et al, 2010). The G-Box is a conserved DNA sequence in
341 plants that has been described as a putative sugar sensor involved in starch

catabolism (Reinhold et al., 2011). A close proximity of the G-Box and the GCC-box is often found in tobacco and linked to the regulation of gene expression during plant defense responses (Buttner, 1997). GCC-box sequences in plants are promoter elements involved in the regulation of expression of plant defense genes against biotic or abiotic stressors and play an important role in relaying pathogen-initiated signals and the activation of plant defense responses (Solano et al, 1998, Brown et al, 2003; Shenk et al, 2000; Reymond et al, 1998). GCC boxes have also been characterized in tomato endochitinase, glucanase and osmotin genes (Tournier et al, 2003). This gene structure rich in sensor motifs and a viral-type sequence indicates that gene expression is modulated by changes in environmental or physiological conditions.

We used the *gag* and *pol* sequences from CcV and the nuclear *gag* sequence to understand the phylogeny of the CcV. The sequences for the *gag* and *pol* genes were compared with other sequences of totiviruses found in GenBank in December 2015. The *pol* genes from CcV cluster within the totivirus and is most closely related to the XdV gene (Figure 7). It is to be noted is that all of these sequences are from terrestrial organisms and most from Fungi. It is, however, unclear if the sequences obtained from terrestrial plants are due to fungal infections or if they are using the plant cells as hosts (Roossinck 2010). Similarly, the CcV *gag* genes cluster with the XdV gene. However, the genomic *gag* gene did not cluster with the CcV sequences, but rather with the *Saccharomyces* and *Scheffersomyces* virus gene sequences. The *gag* gene it is sometimes reported to be of viral origin and sometimes to be of nuclear origin (Figure 7), but no clear phylogenetic pattern emerged.

Even though the nuclear *gag* gene was expressed and dsRNA coding for the *gag* and *pol* genes were present in *C. crispus*, inspections of TEM images showed no clearly

367 identifiable virus-like particles (not shown) nor were any attempts to purify viral
368 particles successful. Therefore, we are not presently convinced that viral particles are
369 produced in *C. crispus* during normal conditions.

370 After finding the CcV in our gametophytic laboratory strain we wanted to know if
371 this was specific for this strain and this life cycle stage. We therefore collected *C.*
372 *crispus* from various locations and in different life cycle stages. The dsRNA could be
373 visualized in all life stages tested, including tetrasporophytes and male and female
374 gametophytes of *C. crispus* from Roscoff and in each part of the thallus tested. A
375 band could also be seen in *C. crispus* samples from every location tested; in Europe
376 from the northernmost location, Tjärnö in Sweden, via Plymouth in the UK, to the
377 southernmost location, Porto in Portugal (Table 2). The band was also present with
378 similar intensity in *C. crispus* with visible epiphytes and lesions. This indicates that
379 the presence of dsRNA is wide-spread within *C. crispus*, at least along the European
380 coast. For the laboratory strain with an origin in Canada, we cannot exclude an
381 “infection” after the arrival of the strain in the laboratory in 1986.

382 We also wanted to verify if the presence of dsRNA was a phenomenon specific for
383 *C. crispus* or if it was a more general occurrence. We therefore collected samples of
384 other red algae around Roscoff, France (Table 3). A visible band was found in most
385 of the red algae, except *Palmaria palmata* and *Furcellaria lumbricalis*, including one
386 member of the Bangiales, a *Porphyra/Pyropia* species. This indicates that the
387 presence of dsRNA identities of probable viral origin is wide-spread in red algae.

388 When the first sequence signatures showed similarities with totiviruses, which are
389 mainly found in fungi or protozoa, we first suspected a contamination of *C. crispus*
390 by epi- or endophytes in the culture medium. To exclude this, we decided to extract

nucleic acids from protoplasts. In protoplasts, the cell wall is enzymatically digested with cellulase and carrageenase (Le Gall et al, 1990). The total nucleic acid extraction from protoplast was then obtained and a band of similar intensity was seen (Figure 7).

We consider CcV to be associated with *C. crispus* and not other eukaryotic organisms since it was found in extractions of RNA performed simultaneously with the DNA extractions for the genome project and no DNA from other eukaryotes were found during the genome analysis. The cultures of *C. crispus* are not axenic, so it is possible that the dsRNA virus is associated with bacteria. However, the virus was also present in extractions of RNA from protoplasts (Figure 7), where bacteria associated with the cell wall were drastically reduced. Totiviruses are also unknown from bacteria (Ghabrial and Suzuki 2009) and considering the amounts of dsRNA found on the gels we find it is highly likely that CcV is associated with *C. crispus* and not an associated organism. This is, at least, true for the Peggy's Cove strain. For the other samples microscopic analysis to localize endophytic fungi gave negative results (data not shown), but we cannot formally exclude that endophytic fungi associated with algae might be the host of totiviruses for these strains and species.

CONCLUSIONS

A totivirus-like entity closely related to viral strains that mostly infect fungal species exists in the red macroalga *Chondrus crispus*. Double-stranded viral RNA was found in relatively high quantities, but no viral particles were found, nor has any phenotype been identified. It is present in all tested life cycle stages and ubiquitous in *C. crispus* isolates from different geographic areas. In a limited sample very large

sequence diversity was seen. Similar bands of dsRNA are present in other red algae. To our knowledge, this is the first time a virus-related sequence has been reported from a red alga and the first time a totivirus is reported from algae. The CcV genome is organized as described in other totiviruses with two overlapping ORFs including a 5'-ORF coding a capsid protein (CP) and a 3'-ORF coding an RNA-dependent RNA polymerase. This involves an ORF disruption with an internal stop codon and a -1 ribosomal frameshift. A copy of the gene encoding the capsid was identified in the nuclear genome of *C. crispus*. Despite the fact that the presence of dsRNA seems to be widespread in red algae nothing is known about the physiological and ecological consequences of the association with totivirus-like entities.

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 671

672 Table 1. Comparisons of % identity between amino acid sequences obtained by
 673 sequencing a dsRNA band from *Chondrus crispus* and comparing multiple
 674 sequences for the *pol* gene. Sequences were translated to amino acids, aligned and
 675 compared pairwise using Clustal Omega (www.ebi.ac.uk).

	CcV1	CcV2	CcvP5	CcvP6	CcvP7	CcvP8	CcvP9
CcV1	100						
CcV2	37	100					
CcvP5	44	52	100				
CcvP6	29	24	28	100			
CcvP7	50	57	55	29	100		
CcvP8	23	21	26	19	26	100	
CcvP9	50	61	52	32	65	27	100

676

677 Table 2. The presence of a detectable dsRNA band around 6 kbp in RNA
 678 extractions from *Chondrus crispus*.

Life cycle stage	Location		Virus detected
Gametophyte male	Roscoff, France	(48° 43' N, 3° 58' W)	Yes
Gametophyte female	Roscoff		Yes
Tetrasporophyte	Roscoff		Yes
Gametophytes stressed*	Roscoff		Yes
Gametophyte	Tjärnö, Sweden	(58° 86' N, 11° 13' E)	Yes
Gametophyte	Plymouth, UK	(50° 39' N, 4° 9' W)	Yes
Gametophyte	Helgoland, Germany	(54° 11' N, 7° 53' E)	Yes
Gametophyte	Chaucre, France	(45° 59' N, 1° 22' W)	Yes
Gametophyte	Lorient, France	(47° 42' N, 3° 23' W)	Yes
Gametophyte	Viana do Castelo, Portugal	(41° 42' N, 8° 51' W)	Yes

679 *Algae with visible epiphytes and lesions.

680

681 Table 3. The presence of a detectable dsRNA band with a size around 6
682 kbp in different species of red macroalgae collected around Roscoff,
683 France.

Species	Class	Band detectable
<i>Dumontia contorta</i> (S.G. Gmelin) Ruprecht	Florideophyceae	Yes
<i>Grateloupia turuturu</i> Yamada	Florideophyceae	Yes
<i>Chondracanthus acicularis</i> (Roth) Fredericq	Florideophyceae	Yes
<i>Porphyra/Pyropia</i> sp	Bangiophyceae	Yes
<i>Laurencia pinnatifida</i> (Hudson) J.V. Lamouroux	Florideophyceae	Yes
<i>Mastocarpus stellatus</i> (Stackhouse) Guiry	Florideophyceae	Yes
<i>Polysiphonia elongata</i> (Hudson) Sprengel	Florideophyceae	Yes
<i>Palmaria palmata</i> (L.) Weber & Mohr	Florideophyceae	No
<i>Corallina elongata</i> J. Ellis & Solander	Florideophyceae	Yes
<i>Furcellaria lumbricalis</i> (Hudson) J.V. Lamouroux	Florideophyceae	No

684

FIGURE LEGENDS

Figure 1. Agarose gel electrophoresis of nucleic acid extract from *Chondrus crispus*:
A. native control (1), native control incubated 1h at 40°C (2), DNase1 treated (3),
RNase treated (4); L Smart Ladder (Eurogentec). B. Native control with 10%
DMSO (1), after denaturing 5 min at 95 °C with 10% DMSO.

Figure 2. Schematic representation of the *Chondrus crispus* viral sequences
compared to the sequences of related viral sequences of *Xanthophyllomyces*
dendrorhous (XdL1b, XdL2) and *Saccharomyces cerevisiae* (ScVL-A(L1)); note the
presence of the viral capsid sequence integrated into the *C. crispus* genome
(CcVGe). The capsid coding open reading frame (ORF) in blue, RNA dependent
RNA polymerase coding ORF in red, and overlapping ORF in purple.

Figure 3. Alignment of domains in the RNA-dependent RNA polymerase (*pol*) gene
from totiviruses as determined in Bruenn (2003) and Koonin (1991) with the
Chondrus crispus dsRNA sequences. The consensus conserved domains described in
Baeza et al. (2012) are highlighted.

Figure 4. Pseudoknot structure prediction of the CcV2 sequence using DotKnot
method (based on Cao and Chen 2006, 2009) and drawn using PseudoViewer3
(Byun et al. 2009).

Figure 5. Genome region of *Chondrus crispus* containing the *gag* gene homolog. CYP: fragment of a cytochrome P-450 gene similar to a *C. crispus* nuclear gene (in blue); E-Box, G-Box and GCC-Box (in grey boxes): cis-regulatory elements located upstream the *gag* gene CcVGe (in red); open reading frame related to an unknown protein found in *Simkania nevegensis* (in green); putative sigma-type promoters (underlined in black); universal genetic code/yeast mitochondrial genetic code (in italics).

Figure 6. A. An unrooted neighbor-joining tree of the capsid sequence (n, reported as nuclear gene; v, reported as viral gene). B. An unrooted neighbor-joining tree of RNA dependent RNA polymerase sequences. The alignment was based on entire sequences using Muscle and neighbor-joining analysis was carried out in MEGA.

Figure 7. Gel electrophoresis of nucleic acid extracts from *C. crispus* protoplasts on a 1% agarose gel stained with EtBr. L: smart Ladder Eurogentec

Figure 1

Figure 2

		Domain1	Domain2	Domain3
		XEXXXLXGRXXXXXXEXXXRX	RXXWXXBXX	KXEXGKRXI XXDXXXYXXXXXXXB
CcV1		FEMNLI NRQGTVDVDAEYKRS	SRVDFPTGSI	IKYEWGRRAIYGTDLTSVLTNFAMVCE
CcV2		FEID VLSNRVGGVDVQGEKENRL	ARVDMATGS	IKYEWGRRAIYGTDLTSVLTHFAFYNCE
CcVP9		FEAEVLNRSI GQVDVGEKENRL	ARVDMAGGS	VKYEWGRRAIYGTDLTSVIMAHFAFYNCE
CcVP7		GVDVGEKNRM	ARVDMAGGS	VKYEWGRRAIYGTDLTSVIMSHFAFYNCE
CcVP5		FEID VLI NRYLEPIXESEKRNQ	ARVDMATGS	IKYEWAKRAIYGTDLTSVLTNFVFNCE
X. dendrorhous	V- L1b	FEIDT LI NRD GEVDVWEERDHRV	SRVDMAGGS	IKYEWGRRAIYGTDLTSVLSNFAFYNCE
X. dendrorhous	V- L1a	FELEVLNRD GVDVGEKENRQ	NRVDMAGGS	TKYEWGRRAIYGTDLTSVLSNFAFYNCE
Black raspb erry	F	FELQVLNRD GVDVWNERKHQ	SRVEWPTGSV	VKYEWGRRAIYGTDLTSVITNYAMFCE
T. aestivum	V1	FELTVLLNRD GQVDVGEKENRQ	SRVEWPTGSV	TKYEWGRRAIYGTDLRSTIITN FAMFCE
S. cerevisiae	L- A	FELAVLMNRG GQVDVGEKENRQ	NRVEWPTGSV	TKYEWGRRAIYGTDLRSTIITN FAMFCE
CcVP8		FELQVLNRD GTD VDAEKEKRI	GRSILMPNGSV	V
S. segobiensis	V- L Y	ELNVLVNRD DSDVWKEIDHRT	GRVAMPNGSV	TKYEWGRRALYGGDVSFLHSDFGMTNCE
S. cerevisiae	L- BC	FELNVLENGVDEVDVGEKENRS	GRVAMPNGSV	TKYEWGRRALYGGDVSHTMABGLLQCE
CcVP6		FETEVLNRD GQDLVDVGEQRNRQ	TRVLHVPTGSA	EKYEWGRRAIYGTDEGFILCDIALPSAE

		Domain4	Domain5	Domain6	Domain7	Domain8
		XXDXDFNSQH-X	TLXSGRXTTFXNXLNXXYX	HXGDDX	KXXXXXXEFLRX	YXXRXXXXXW
CcV1		DFDFDFNSQHS	GTLLSGHRLTTR NSVLNRI YL	I HNGDBL	TKCPFGMAEFLRV	QYLTRAVATLV
CcV2		VDFFDFNSQHS	GTUMSGWRLTTTNSVLNRI YS	VHNGDDVM	TKGSFGGLAEFLRV	QYMTNRNI ATLV
CcVP9		IDFFDFNSQHS	GTUMSGWRLTTTNSVLNLI YT			
CcVP7		IDFFDFNSQHS	GTLLSGWRLTTTNTI LNYI YT	VHNGDDM		
CcVP5		LDFFDFNSQHS	ATLLSGWRLTTTNSVLNAVYT	LHNGDDM	SKCAFGGIAEFLRI	QYLTRAI ATLM
X. dendrorhous	V- L1b	LDFFDFNSQHS	GTLLSGWRLTTTNSVLNLI YT	LHNGDDM	SKCAVGAIAEFLRI	QYLSRAVATMR
X. dendrorhous	V- L1a	YDFDDFNAQHS	GTLLSGWRLTTTNTVLNLI YF	VHNGDDM	AKQNVFSIGEFLRV	QYLSRAATLV
Black raspb erry	F	FDYDDFNSQHS	GTLLSGWRLTTTNTVLNLI YF	VHNGDDM	TKQNVLSISEFLRI	QYLSRSATLV
T. aestivum	V1	FDYDDFNSQHS	GTLLSGWRLTTTNTVLNLI YF	VHNGDDM	AKQNVLSISEFLRV	QYLSRSATLV
S. cerevisiae	L- A	FDYDDFNSQHS	GTLLSGWRLTTTNTVLNLI YF	VHNGDDM	AKQNVLSISEFLRV	QYLSRSATLV
CcVP8		FDYDDFNSQHS	GTUMSGWRLTTFMNTVLNRVYL	LHNGDDM	SKTNLGTIGEFLRV	QYLSRSVATLV
S. segobiensis	V- L	YDFDDFNSQHS	GTLLSGWRLTTFMNTALNYCYL	LHNGDDM	TKMNIGTIAEFLRV	QYLTRGIATFT
S. cerevisiae	L- BC	YDFDDFNSQHS				
CcVP6

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Figure 5

Figure 6

Figure 7