Abstract: Hypovirulence of phytopathogenic fungi are often conferred by mycovirus(es) infections and for this reason many mycoviruses have been characterized, contributing to a better understanding of virus diversity. In this study, three strains of Fusarium head blight fungus (Fusarium boothii) were isolated from Ethiopian wheats as dsRNA-carrying strains: hypovirulent Ep-BL13 (>10, 3 and 2.5 kbp dsRNAs), and virulent Ep-BL14 and Ep-N28 (3 kbp dsRNA each) strains. The 3 kbp-dsRNAs shared 98% nucleotide identity and have single ORFs encoding a replicase when applied to mitochondrial codon usage. Phylogenetic analysis revealed these were strains of a new species termed Fusarium boothii mitovirus 1 in the genus Mitovirus. The largest and smallest dsRNAs in Ep-BL13 appeared to possess single ORFs and the smaller was originated from the larger by removal of its most middle part. The large dsRNA encoded a replicase sharing the highest amino acid identity (35%) with that of Botrytis virus F, the sole member of the family Gammaflexiviridae. Given that the phylogenetic placement, large genome size, simple genomic and unusual 3′-terminal RNA structures were far different from members in the order Tymovirales, the virus termed Fusarium boothii large flexivirus 1 may form a novel genus and family under the order.

Keywords: Fusarium head blight; mycovirus; RNA genome; mitovirus; Tymovirales; Ethiopia
and information on mycovirus diversity has expanded and deepened dramatically through intensive mycovirus hunting aiming at screening for useful virocontrol agents [1]. However, while mycovirus hunting has been extensive in Asian, European, American and Oceanian countries, its exploration in the African continent has been limited: Diaporthe ambigua RNA virus (an unclassified RNA virus), Diplodia scrobiculata RNA virus 1 (a fusagravirus) and Sphaeropsis sapinea RNA virus 1 and 2 (victoriviruses) are reported in South Africa [12–16] with the remainder being mycovirus-like sequences obtained by metagenomic virome approaches [17].

Fusarium head blight (FHB) is an economically important disease of wheat caused by members of the *Fusarium graminearum* species complex (FGSC) that consists of at least 16 species including *F. boothii*. The disease symptoms are reduced yields, shrivelled kernels, reduced seed quality, and diminished seed weight. In addition, the pathogen leaves mycotoxins such as deoxynivalenol (DON) and nivalenol, type B trichothecenes, on grain surface which can cause diarrhoea and vomiting in animals ingesting them. An increasing number of mycoviruses have been reported to infect various *Fusarium* species, including FGSC from different regions of the world. These viruses include members of *Chrysoviridae*, *Hypoviridae*, *Partitiviridae*, *Narnaviridae*, *Totiviridae*, and *Mymonaviridae* families; proposed *Alternaviridae* and *Fusariviridae* families; and other unclassified virus groups [9,18–27]. Those with the greatest potential as virocontrol agents are: *Fusarium graminearum* virus 1 strain DK21 (FgV1-DK21), *Fusarium graminearum* hypovirus 2 strain JS16, and *Fusarium virguliforme* dsRNA mycovirus 1 and 2 [18,21,23,28].

As abovementioned, mitoviruses in the genus *Mitovirus* in the family *Narnaviridae* are omnipresent in filamentous fungi and the members have approximately 3 kb positive sense single stranded RNA ((+)ssRNA) genomes that possess the single open reading frame (ORF) encoding an RNA-dependent RNA polymerase (RdRp) [29]. The ORF accommodates "UGA" triplets that is an interrupting stop codon when applied nuclear gene code but translated as tryptophan in the mitochondrial gene code. *Cryphonectria parasitica* mitovirus 1 (CpMV1) is the first mitovirus described, and co-presence of virus-derived double stranded RNA (dsRNA; considered as a replicative form) with subcellularly fractionated mitochondria suggests the localization of CpMV1 in mitochondria [30]. Khalifa and Pearson [31] observed the accumulation of filamentous structures in mitochondria of *Sclerotinia sclerotiorum* under electron microscope, specifically when a mitovirus infected to the host. Therefore, mitoviruses are expected to uniquely replicate in the host mitochondria. *Botrytis cinerea* mitovirus 1, *Thanatephorus cucumeris* mitovirus (TeMV, known also as *Rhizoctonia* M2 virus), *Sclerotinia sclerotiorum* mitovirus 1/KL-1, *Sclerotinia sclerotiorum* mitovirus 2/KL-1, *Sclerotinia sclerotiorum* mitovirus 3/NZ1, and *Sclerotinia sclerotiorum* mitovirus 4/NZ1 were shown to have a hypovirulence-conferring ability but effects of most mitoviruses on host fungal phenotypes have not been elucidated because mitoviruses are generally hard to cure and introduce [7,31–35]. Several mitoviruses have also been reported from *Fusarium* species including FGSC but biological properties of those are largely unknown [25,36,37].

The order *Tymovirales* contains viruses that have a non-segmented (+)ssRNA genome with single or multiple ORFs and its members mostly infect plants. The order consists of five families—*Alphaflexiviridae*, *Betaflexiviridae*, *Gammaflexiviridae*, *Deltaflexiviridae* and *Tymoviridae* in International Committee on Taxonomy of Viruses (ICTV)—based on the genome organisation, phylogenetic relationships and virion morphology. As of now, *Alpha*, *Gamma* and *Deltaflexiviridae* include mycoviruses and, to our knowledge, at least 10 such mycoviruses have been characterised including the ones associated with hosts that exhibit hypovirulent phenotypes [20,27,38–45]. *Botrytis cinerea* virus F (BVF, *Gammaflexiviridae*) was isolated from a hypovirulent strain of *Botrytis cinerea* as the first mycovirus taxonomically classified in the order *Tymovirales*, and was followed by *Botrytis* virus X (BVX, *Alphaflexiviridae*) [39,40]. *S. sclerotiorum* also hosts mycoviruses of this order, namely *Sclerotinia sclerotiorum* debilitation-associated RNA virus (SsDRV, *Alphaflexiviridae*) and *Sclerotinia sclerotiorum* deltaflexivirus 1 (SsDFV1, *Deltaflexiviridae*). The former is associated with hypovirulence of its host and the latter is recognised as the first mycovirus in the order
Tymovirales that has no capsid protein (CP) [41,42]. Recently, Rhizoctonia solani flexivirus 1 (RsFV-1, unclassified deltaflexivirus-like) was reported to have the largest genome in the order and its unique genomic organisation broadened our perception of genome plasticity within the order [38]. In addition, two mycoviruses, Fusarium graminearum mycotymovirus 1 (FgMTV1, unclassified tymovirus-like) and Fusarium graminearum deltaflexivirus 1 (FgDFV1, Deltaflexiviridae), have been isolated from the FHB pathogen, *F. graminearum* [20,27]. Biological assays demonstrated that FgMTV1 infection does not alter the host virulence, but decreases the growth rate and DON production. Finally, although oyster mushroom spherical virus (OMSV) was earlier reported to be a tyloo-like mycovirus [44], its taxonomic placement is inconclusive till date.

This study characterised a novel member of Tymovirales, potentially forming a new genus or family, and members of a new species in the genus Mitovirus in the family Narnaviridae. These virus classes were identified for the first time in *Fusarium boothii* strains from the African continent. It is noteworthy that the tyloo-like virus possesses the largest genome size (over 12.5 kb) among the currently reported Tymovirales members. This study contributes to our understanding of Tymovirales virus and mitovirus diversity and evolution.

2. Materials and Methods

2.1. Fungal Strains and Culture

*F. boothii* strains (Ep-BL13, Ep-BL14 and Ep-N28) were isolated from wheats with the FHB symptoms in Hadiya region of southern Ethiopia. A Japanese isolate of *F. graminearum* sensu stricto (s.str.) strain was also used as a reference. All strains were maintained at 28 °C on potato dextrose agar (PDA, BD Difco Laboratories, Detroit, MI, USA) and synthetic low nutrient agar (SNA) and stored at 4 °C in the dark. Mycelia were cultured in potato dextrose broth (PDB, BD Difco Laboratories, Detroit, MI, USA) liquid medium at room temperature for 4–7 days for nucleic acid extractions.

2.2. Biological Assessment of FGSC Strains

Growth rate and colony morphology. Mycelial plugs were transplanted on PDA and SNA plates (three replicates) and incubated at 20 °C for 1 week. The longest and shortest diameters of each colony were measured 3–7 days post-transplanting and the mean diameter was used for the calculation of colony area.

Pathogenicity test. Conidial spores (1 × 10⁵ spores/mL) suspended in a 1% (v/v) Triton solution were used for inoculation, and the FHB-sensitive wheat cultivar Apogee was grown at 27 °C in a chamber (KG-50HLA; Koito Electric Industries, Yokohama, Japan) to avoid possible release of inocula into the environment [46,47]. Briefly, 10 µL of spore suspension was injected into wheat spikelets, and the number of diseased spikelets was counted at 15 days post-inoculation. Alternatively, the spore suspension was sprayed three times (500 µL/push) to a wheat head, and disease severity was evaluated according to Ban and Suenaga’s criteria (index, mild to severe; 0, 5, 10, 20, 30, 50, 60, 80, 100) at 15 days post-inoculation [48]. Inoculated plants were placed in a transparent humid box for one day (injection method) or two days (spray method) to promote fungal germination, prior to the maintenance in the chamber.

2.3. Genomic Polymerase Chain Reaction

Fungal genomic DNA was purified by conventional CTAB (cetyl trimethylammonium bromide) extraction. The partial internal transcribed spacer (ITS) of rDNA and transcription elongation factor 1-α gene (*TEF1-α*) sequences were PCR-amplified with the oligonucleotide primers reported by O’Donnel et al. [49,50]. Primers used in this study were: primer 1 (5'-TCATAAGTGGTAGGA(A/G)ACAGAGAC-3') and primer 2 (5'-GCCTGGA(G/A)GTAC CAGT(G/C)ATCATGTT-3') for ITS; and primer 3 (5'-GTGGGCGATTACCCCGCC-3') and primer 4
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(5′-GAGTGGCGGGGTAAATGCC-3′) for TEF1α. Fungal species were identified by homology search with and by phylogenetic analysis with sequences from public databases.

2.4. dsRNA Extraction and cDNA Library Construction

dsRNA fraction was extracted from fungal mycelia by conventional cellulose column chromatography. Filtrated mycelia were homogenised in the presence of liquid nitrogen, and total RNA fractions were obtained by treatment with one round each of phenol, phenol–chloroform–isoamylalcohol (25:24:1), and chloroform–isoamylalcohol (24:1) extraction. dsRNAs were further isolated from the total RNA fractions by using Cellulose Powder C (Advantec, Tokyo, Japan). To eliminate fungal chromosomal DNA and ssRNA species, these fractions were treated with RNase-free RQ1 DNase I (Promega, Madison, WI, USA) and S1 Nuclease (Promega). Conventional cDNA library construction was performed for viral sequence determination. The dsRNAs extracted from F. boothii Ep-BL13, Ep-BL14 and EP-N28 and the 3 kbp- and 2.5-kbp dsRNA bands were gel-purified using a Zymoclean™ Gel RNA Recovery kit (Zymo Research, Irvine, CA, USA). cDNA libraries of each dsRNA fragment were constructed using a non-PCR and a PCR-based method. After denaturation of dsRNA templates at 99 °C for 5 min, cDNA was synthesised using ReverTra Ace-α-(Toyobo, Osaka, Japan) with an adapter-tagged primer (5′-CCTGAATTCGGATCCTCCNNNNNNN-3′). The resulting cDNA fragments were purified from an agarose gel with a size exclusion (1.0–2.0 kb) and amplified by PCR with an adapter primer (5′-CCTGAATTCGGATCCTCC-3′). The PCR conditions are as follows: 94 °C (2 min), then 1 cycle at 94 °C (2 min)/65 °C (1 min)/72 °C (1 min), followed by 35 cycles at 94 °C (40 s)/52 °C (30 s)/72 °C (3 min), and a final extension at 72 °C for 7 min. DNA fragments were cloned into the pGEM-Teasy (Promega) or pCR-Blunt (Thermo Fisher, Waltham, MA, USA) cloning vectors and used for the transformation of Escherichia coli strain DH5α for Sanger sequencing analyses.

2.5. RNA Seq, Reverse Transcription PCR and Rapid Amplification of cDNA Ends

RNA seq. The F. boothii Ep-BL13 virome was established on an Illumina-Hiseq 2500 system (Illumina, San Diego, CA, USA) by a pair-end sequencing run (2 × 100 bp) using 10 ng of dsRNA as a template. cDNA and library construction were outsourced to BGI Japan. Fastqc version 0.11.5 software (https://www.bioinformatics.babraham.ac.uk/projects/fastqc/) was used to check the quality of the read data [51]. Reads were filtered using Sickle version 1.33 software (https://github.com/najoshi/sickle) by requiring an average Phred quality (Q score) of at least 30 and to only retain reads of 20 nucleotides (nt) or greater in length [52]. De novo assembly was carried out using SPAdes version 3.11.1 software (http://cab.spbu.ru/software/spades/) at the default settings [53].

Gap-filling RT-PCR. Target cDNAs were produced using ReverTra Ace and a gene-specific primer set from given dsRNA fragments and amplified with DNA polymerases such as PrimeSTAR (Takara, Shiga, Japan), KOD FX Neo (Toyobo) or GO-Taq (Promega).

3′-RLM-RACE (RNA ligase mediated RACE). The terminal sequences of dsRNAs were determined by this method. Pre-denatured dsRNAs in dimethyl sulphoxide (DMSO) (90%, 65 °C) were ligated at their 3′-ends with a 5′-phosphorylated oligodeoxynucleotide, 3′-RACE-adaptor (5′-CAATAACCTTCTGACCATGACGTAGCTCAGCATG-3′), using T4 RNA ligase (Takara) at 16 °C for 16 h. Ligated DNA–RNA strands were DMSO-denatured in the presence of the oligonucleotide 3′-RACE-1st (5′-CATGCTGACTGTCACTGCATG-3′) and used as templates for cDNA synthesis. The resulting cDNA was amplified by PCR with 3′-RNAce-2nd (5′-TGATGGCTAGATGGTAT-3′) and gene-specific primers. Plasmid clones and genomic- or RT-PCR amplicons were purified with spin columns (GeneElute, Sigma, St. Louis, MO, USA; Nucleospin, Takara) and used for BigDye (ABI, ThermoFisher) sequencing by following the manufacturer’s instructions. Sanger sequencing was carried out on a 3100-Avant sequencer (ABI/HITACHI). All virus-specific oligonucleotide primers used in this study were designed based on sequences obtained from cDNA library sequencing and Illumina sequencing.
2.6. Bioinformatics Analyses

Sequence assemblies were manipulated using GeneStudio (http://genestudio.com/), SPAdes and Genetyx (Genetyx, Tokyo, Japan) software. Sequence homology searches were performed with the Basic Local Alignment Search Tool (BLAST) programme provided by the National Center for Biotechnology Information (NCBI, http://www.ncbi.nlm.nih.gov/) and the Genetyx. Motif searching was conducted using the online tool, InterPro: protein sequence analysis & classification (https://www.ebi.ac.uk/interpro/) [54]. For secondary RNA structure prediction, the 3'- and 5'-terminal viral sequences were submitted to mfold (http://unafold.rna.albany.edu/?q=mfold/rna-folding-form) and potential stem-loops were predicted [55].

2.7. Phylogenetic Analysis

Multiple amino acid or nucleotide sequence alignments were constructed using MUSCLE [56] and with the MEGA X [57]. Phylogenetic analysis was conducted by using PhyML ver. 3.0 (http://www.atgc-ontpellier.fr/phyml/) with the best-fit models suggested by Smart Model Selection in PhyML (http://www.atgc-montpellier.fr/sms/) [58], and the results were visualised with FigTree and further enhanced in Adobe Illustrator.

3. Results

3.1. Detection of dsRNAs in F. boothii Strains

The fungal strains Ep-BL13, Ep-BL14 and Ep-28 were isolated from wheats exhibiting FHB in Ethiopia by using Fusarium-selecting medium, which were found to be dsRNA-positive. Agarose gel electrophoresis of dsRNA fractions extracted from these strains showed a 3 kbp common to all three strains and additional >10 and 2.5 kbp bands only in the Ep-BL13 strain (Figure 1A), thereby suggesting infections by unknown RNA mycoviruses. These fungal strains were identified as F. boothii based on comparison of partial TEF1-α and ITS sequences (data not shown) and by phylogenetic analysis of partial TEF1-α sequences (Figure S1).

3.2. Biological Characteristics of F. boothii Strains

Colony morphology of the F. boothii strains on nutrient-rich (PDA) and -poor (SNA) solid media varied as shown in Figure 1B,C. The Ep-BL13 strain grew most slowly on PDA, followed by Ep-BL14 and then Ep-28 (Figure 1B,D). The strains also exhibited different phenotypes on SNA, with Ep-BL13 and Ep-N28 growing faster and slower than those on PDA, respectively (Figure 1C). The pathogenicity of the three F. boothii isolates and of a reference F. graminearum s.str. strain on wheat head was investigated with injection and spray methods. Plants inoculated with F. boothii Ep-BL14 and Ep-28 exhibited typical FHB symptoms similar to that of the plant infected by F. graminearum s.str., whereas the Ep-BL13 showed apparently weak virulence (Figure 1E,F). The effects of the dsRNA elements on host fungal growth and pathogenicity to wheats are yet to be determined.

3.3. Analysis of Viral Sequences: A Typho-Like Virus

The genome organisation of the coding strand of the largest dsRNA segment harboured in Ep-BL13 is shown in Figure 2A. The complete genomic sequence of the virus was 12,579 nt in length and the G + C content of the dsRNA is 64.93% (accession No. LC425115). A BLASTX search of the whole nucleotide sequence showed that the highest amino acid (aa) identity (35%) was to the replicase of BVF, which is the sole member of the family Gammaflexiviridae (Table 1). Based on this result, the virus was tentatively named Fusarium boothii large flexivirus 1 (FbLFV1). The whole genome of FbLFV1 encompasses a single large ORF putatively encoding a polypeptide of 4070 aa (162–12,374 nt) with a calculated molecular mass of 443.6 kDa. However, BVF, the closest virus to FbLFV1 in a BLAST search, possesses two ORFs encoding a putative replicase (a potential readthrough product) and a putative
CP. Methyltransferase (Met), helicase (Hel) and RNA-dependent RNA polymerase (RdRp) domains of the FbLFV1 polypeptide showed low aa identities (22–36%, 20–38% and 25–37%, respectively) to those of representative viruses in the order Tymovirales and related mycoviruses, as shown in Table 1. Three replicase domains, Met, Hel and RdRp, are located in this order, from N to C terminus, and this is typical of the “alphavirus-like” superfamily that includes many other plant-infecting viruses (Figure 2A) [59]. There were two regions in the polypeptide with similarity to a PHA03247 domain (N-terminal part, E-value: $3.94 \times 10^{-6}$; C-terminal part, E-value: $5.01 \times 10^{-3}$), which was found in the large tegument protein UL36 of herpesviruses (a large dsDNA virus), although the function of PHA03247-like domains in FbLFV1 is unknown.

Figure 1. Biological properties of F. boothii strains and dsRNA-banding profiles. (A) The dsRNA-banding profiles of each F. boothii strain (Ep-BL13, Ep-BL14 or Ep-28) and a reference pathogenic strain. F. graminearum s.str. Three F. boothii strains all carried dsRNA fragments of approximately 3 kbp. Aside from the 3-kbp dsRNA band, Ep-BL13 harboured two additional dsRNA bands of over 10 kbp and about 2.5 kbp. (B,C) The colony morphologies of F. boothii strains and F. graminearum s.str. on PDA nutrition-rich media (B) and SNA nutrition-poor media (C) that were photographed at four days after transplanting. The three F. boothii strains exhibited various colony morphologies in terms of colony size, the amount of aerial mycelium and pigmentation. (D) Growth rate of Fusarium strains on PDA media. Colony sizes were measured at 3–7 days post-transplantation. (E) Pathogenicity test I. A wheat spikelet was inoculated with Fusarium strains by injection and photographed at 15 days post-inoculation. The number of spikelets exhibiting symptoms is presented. (F) Pathogenicity test II. Wheat photographs taken as in (E), but after spray inoculation. Disease indices (0–100) as per Ban and Suenaga (2000) [48] are shown.
Figure 2. Schematic representation of the genome structure of the large F. boothii mycovirus. (A) Genome maps of FbLFV1. Thick bars are non-coding sequences and open boxes represent ORFs. Colour-highlighted boxes in ORFs represent the predicted protein domains as indicated. The length of the whole genome, ORF and UTR are given. The deleted region of FbLFV1 in defective RNA (D-RNA) is denoted by the dashed lines and nucleotide positions. Met, methyltransferase domain; Hel, helicase domain; RdRp, RNA-dependent RNA polymerase domain, FS, frame-shift. (B) Comparison of nucleotide contexts surrounding potential initiation codons (red bold letters, potential initiation codon; red underlined letters, surrounding nucleotides) of FbLFV1. Favourable contexts of initiation codon in vertebrate, filamentous fungi and yeast are shown as references. (C) Structure of terminal, non-coding RNA regions of FbLFV1. 5'- and 3'-terminal sequences of the genomes of the FbLFV1 were analysed with secondary structure prediction tools. Representatives of putative stem-loop structures were depicted. Dots in different colors indicate hydrogen bonds between different base pairs: blue, A-U pairs; red, G-C pairs; green, G-U pair.

The initiation codon at 162–164 nt of the large ORF was in a favourable context, CACCAUGGa, for filamentous fungi, although another potential initiation codon at 27–29 nt appeared in frame to the ORF with a weak context uugCAUGcG (capital and small letters represent good and poor residues in the context, respectively; underlined letters are the putative initiation codon; Figure 2B) [60–62]. FbLFV1 had short 5' and 3' untranslated regions (UTRs) that are 161 and 205 nt, respectively (Figure 2A). Although experimental evidence has not been obtained, the presence of a Met domain suggested the capping of 5'-end of FbLFV1 genomic RNA. The Met domain of the members of the order Tymovirales is generally located at the extreme N-terminal region of the replicase but that of FbLFV1 was distant from N-terminus of the polypeptide (aa positions 1261–1528) (Figure 2A).
Table 1. List of viruses in the order Tymovirales and related unassigned viruses with similarity to FbLFV1.

| Taxon               | Virus Name (Abbreviation) | Accession       | Met (%) | Hel (%) | RdRp (%) |
|---------------------|---------------------------|-----------------|---------|---------|----------|
| **Alphaflexiviridae** |                           |                 |         |         |          |
| Allexivirus         | Shallot virus X (SVX)     | NC_003795       | 58/198  | 37/154  | 74/221   |
| Betrexiidae         | Botrytis virus X (BVX)    | NC_003512       | 60/196  | 45/176  | 76/221   |
| Lolavirus           | Lolium latent virus (LLV) | NC_010434       | 64/193  | 50/241  | 80/238   |
| Mandarivirus         | Indian citrus ringspot virus (ICRV) | NC_003093 | 59/162  | 63/244  | 84/229   |
| Platyphovirus       | Donkey orchid symptomless virus (DOSV) | NC_002894 | 60/201  | 49/189  | 69/225   |
| Potexvirinae        | Potato virus X (PVX)      | NC_011620       | 60/198  | 58/245  | 79/227   |
| Sclerodarnavirus    | Sclerotinia sclerotiorum debilitation-associated RNA virus (SsDARV) | NC_007415 | 63/198  | 49/191  | 71/227   |
| **Betaflexiviridae** |                           |                 |         |         |          |
| Carlavirus          | Potato virus M (PVM)      | NC_001361       | 55/217  | 35/127  | 62/224   |
| Foveavirinae        | Apple stem pitting virus (ASPV) | NC_001749 | 60/222  | 17/45   | 63/222   |
| Rabievirinae        | Cherry necrotic rusty mottle virus (CNRMV) | NC_002468 | 52/224  | 44/159  | 61/219   |
| **Trivirinae**      |                           |                 |         |         |          |
| Capillovirus        | Apple stem grooving virus (ASGV) | NC_003462 | 64/214  | 33/97   | 72/231   |
| Chordovirinae       | Carrot Ch virus 1 (CCChV) | NC_002469       | 54/214  | 33/118  | 66/225   |
| Citrovirus          | Citrus leaf blotch virus (CLBV) | NC_003887 | 46/205  | 46/160  | 68/223   |
| Divovirus           | Durius virus A (DVA)      | NC_019029       | 53/217  | 26/90   | 66/224   |
| Prunerevirus        | Apricot vein clearing associated virus (AVCSV) | NC_002395 | 49/183  | 24/62   | 66/224   |
| Tepovirus           | Potato virus T (PTV)      | NC_011062       | 56/186  | 34/115  | 67/219   |
| Trichovirus         | Apple chlorotic leaf spot virus (ACLSV) | NC_001409 | 62/195  | 43/172  | 64/218   |
| Vitovirus           | Grapevine virus A (GVA)   | NC_003604       | 50/201  | 22/57   | 65/218   |
| **Gammaflexiviridae** |                           |                 |         |         |          |
| Mycoflexivirinae    | Botrytis virus F (BVF)    | NC_002604       | 63/193  | 71/252  | 82/224   |
| **Deltaflexiviridae** |                           |                 |         |         |          |
| Deltaflexivirinae   |                           |                 |         |         |          |
| Sclerotinia sclerotiorum deltaflexivirus 1 (SsDFV1) | NC_038977 | 54/202 (26%) | 61/253 (24%) | 45/174 (25%) |
| Sclerotinia sclerotiorum deltaflexivirus 2 (SsDFV2) | MH299910 | 63/222 (28%) | 44/184 (23%) | 59/207 (28%) |
| Fusarium graminearum deltaflexivirus 1 (FgDFV1) | FT690226 | 67/204 (32%) | 37/157 (23%) | 57/211 (27%) |
| **Tymoviridae**     |                           |                 |         |         |          |
| Maculavirus         | Grapevine fleck virus (GFV) | NC_003347 | 57/203 (28%) | 43/144 (29%) | 54/143 (37%) |
| Mansfieldavirus      | Maize rayado fino virus (MRFV) | NC_002786 | 58/197 (29%) | 57/201 (28%) | 45/140 (32%) |
| Tymovirus           | Turnip yellow mosaic virus (TUMV) | NC_004063 | 56/196 (28%) | 58/214 (27%) | 61/204 (29%) |
| **Unassigned**      |                           |                 |         |         |          |
| Mycoymovirus        | Fusarium graminearum mycoymovirus 1 (FgMTV1) | KT360947 | 60/196 (30%) | 57/197 (26%) | 73/219 (33%) |
| Rhizoctonia solani flexivirus 1 (RnfFV1) | NC_005063 | 57/198 (28%) | 53/228 (23%) | 65/224 (29%) |
| Rhizoctonia solani flexivirus 2 (RfnFV1) | KX349069 | N/A ** | N/A ** | 50/177 (28%) |
| Fusarium boothii large flexivirus 1 (BlFV1) | LC425115 | 48/189 (25%) | 61/237 (25%) | 73/219 (33%) |

* Results of homology search against FbLFV1 for amino acid cover region and identity are shown. ** N/A indicates “not applicable” due to the unavailability of homologous sequences.
3.4. Presence of a Defective Segment of FbFV1 in the Ep-BL13

The cDNA-library-based sequencing of the approximately 2.5 kbp dsRNA revealed that the segment was of a defective RNA (D-RNA) of FbLFV1. The total length of the dsRNA was 2408 bp, and the positive-sense RNA retained an ORF encoding polypeptide with a potential molecular mass of 76.8 kDa (716 aa; Figure 2A) (accession No. LC425116). This occurred because of the loss of aa 303–3693 of the FbLFV1 large polypeptide (removal of FbLFV1 genome positions 1066–11,235 in-frame) and the addition of 36 aa because of a frame-shift from a single nucleotide deletion at −5 nt position relative to the original termination codon, leading to a new downstream stop codon. Since the RNA sequence was almost identical to the corresponding region of FbLFV1 and the junction of the upstream/downstream parts was uniform, this D-RNA of FbLFV1 might have arisen very recently, but at least before the isolation of Ep-BL13. The segment has the potential to facilitate viral replication because the coded protein retained partial RdRp motifs (122 aa residues of C-terminal end; Figure 2A); however, it is also possible that it may interfere with replication. As BVF is reported to carry similar D-RNA species with unknown function [39], it is interesting to investigate the significance of D-RNAs in life cycles of mycoviruses in the order Tymovirales.

3.5. Predicted RNA Structure of the Terminal Regions of FbLFV1 Genome

The UTRs of RNA viruses generally carry important cis-elements that form secondary/tertiary structures and/or distant base-pairing. In many cases, these are essential for encapsidation, transcription, translation and replication. To investigate whether FbLFV1 contains such structures, UTR sequences were subjected to secondary structure prediction using mfold software. The 5′ UTR of FbLFV1 did not consistently exhibit stem loops in several outputs, except for the one shown in Figure 2C. Similarly, those of the 3′ UTR were mostly inconsistent but a stem-loop predicted near the 3′-end was the most probable (Figure 2C). It is known that some members of Tymoviridae, such as turnip yellow mosaic virus (TYMV), have tRNA-like structures (TLS) in their 3′-UTR that play an important role in viral replication [63,64]. Given that TLSs are difficult to predict (e.g., the TYMV-TRL (6234–6318 nt, NC_004063) was not detected by tRNAscanning software [65]), more extensive base-pairing analysis of the FbLFV1 3′-end should be conducted before drawing any definitive conclusions.

3.6. Molecular Phylogenetic Analysis of FbLFV1

Amino acid sequences of whole FbLFV1 replicase and representative mycoviruses in the order Tymovirales, including taxonomically unassigned members, were aligned and the Met, Hel and RdRp domains were extracted (Table 1 and Figure 3). These three domains were found to be well conserved in the deduced protein sequence of FbLFV1 replicase. The evolutionarily conserved relationship of FbLFV1 to the Tymovirales members was characterised by maximum likelihood (ML) phylogenetic analyses based on the alignments of each of the three domains (Figure 4). As expected from BLAST analyses, the FbLFV1 Met Hel and RdRp domains were distantly associated with those of viruses in the order Tymovirales, with relatively close relationship to a gammaflexivirus (Figure 4A–C). The concatenate sequence of these domains finally revealed that FbLFV1 was placed independently from the clades of established families in the order Tymovirales, although the branching support value was not sufficiently high in the ML tree (Figure 4D). Based on these phylogenetic analyses, FbLFV1 is expected to belong to a novel virus species “Fusarium boothii large flexivirus 1” that may form an independent genus, or even a new virus family, in the order Tymovirales.
Figure 3. Amino acid alignment of Met, Het and RdRp domains of FbLFV1. The positions of the motifs are indicated by lines above the sequence alignments with the motif numbers from I to VI. Identical residues are colour-highlighted with blue and indicated by asterisks; conserved and semi-conserved amino acid residues are colour-highlighted with yellow and gray, and by colons and dots, respectively.

(A–C) Deduced aa sequences corresponding to Met (A), Hel (B) and RdRp (C) domains were extracted from the FbLFV1 large polypeptide and aligned with those from mycoviruses of the order Tymovirales. Abbreviations of viral names are the same as in Table 1.
Figure 4. Phylogenetic analyses of FbLFV1. (A–C) Phylogenetic trees were constructed based on the alignment of aa sequences of Met (A), Hel (B) and RdRp (C) domains. The ML trees with the best model and rooting methods are shown. Numbers at the nodes represent branch supporting values with an aLRT SH-like method (>0.9 is significant). (D) Phylogenetic tree of Met + Hel + RdRp flanking sequences. Yellow dots indicate mycoviruses and stars indicate FbLFV1. The sequences used for (A–C) were joined and subjected to alignment and phylogenetic analyses as in (A–C). Abbreviations of viral names and accession numbers are listed in Table 1.

3.7. Analysis of Viral Sequences: Three Mitoviruses

The full-length sequences of the 3 kbp-dsRNAs in F. boothii strains Ep-BL13, Ep-BL14 and Ep-N28 were obtained by cDNA library sequencing and 3’ RLM-RACE analyses (accession Nos. LC425112, LC425113 and LC425114, respectively). The viral dsRNAs accumulated in strains Ep-BL13 and Ep-N28
were 2802 bp long and that in strain Ep-BL14 was 2801 bp long. Each positive sense strand sequence possessed a single ORF potentially encoding RdRp when using mitochondrial codon usage (Figure 5A). The length of the ORF and 3′-UTRs was the same among three sequences, being 2472 nt (823 aa) and 136 nt, respectively (Figure S2). A few nucleotide deletions at different positions in the 5′-UTRs were found in the three viral sequences; nevertheless, the 5′-UTR of Ep-BL13 and Ep-N28 were of the same length (194 nt), whereas that of strain Ep-BL14 was 1 nt shorter (193 nt; Figure S2). The nucleotide similarities between these three viruses were 98% in any combinations. A BLASTP search found that the highest aa sequence identity of the RdRp coded by three viral RNAs of Ep-BL13, Ep-BL14 and Ep-N28 were 41% (E-value = 1 × e\(^{-158}\)), 40% (E-values = 8 × e\(^{-158}\)) and 41% (E-value = 2 × e\(^{-159}\)), respectively, to the RdRp sequence of a tentative mitovirus of uncertain origin, soybean leaf-associated mitovirus 5. The highest aa identity with a definitive mitovirus species was 30% to Ophiostoma novo-ulmi mitovirus 3a (OnuMV3a). Thus, these 3 kbp-dsRNAs are apparently replication intermediates of new mitovirus strains that are closely related to each other, and these viruses were tentatively termed Fusarium boothii mitovirus 1 (FbMV1).

Some mitoviruses form a long pan-handle structure between the 5′ and 3′ termini [66]. In the terminal regions of FbMV1 genome, tetra nucleotides at extreme termini 5′-GGGG and CCCC-3′ may have potential to form base pairs. Another common characteristic of mitovirus genomes is the presence of stem-loop structures at near the terminal regions. Both the 5′ and 3′ UTRs of the FbMV1 genome clearly folded into two potential stem-loop structures, as shown in Figure 5B. None of the sequences was predicted to form pseudoknot.

![Figure 5](image)

**Figure 5.** Schematic representation of the genome structure of *F. boothii* mitoviruses. (A) Genome maps of FbMV1. Thick bars are non-coding sequences and open boxes represent ORFs. Highlighted boxes in ORFs represent the predicted RdRp (red) and AroE (blue) domains. The length of the whole genome, ORF and UTR are given. (B) Structure of terminal, non-coding RNA regions of FbMV1. 5′- and 3′-terminal sequences of the genomes of FbMV1 Ep-BL13 were analysed with secondary structure prediction tools. Representatives of putative stem-loop structures were depicted. Red lines indicate nucleotide residues that potentially form base-pairing between 5′- and 3′-ends. Dots in different colours indicate hydrogen bonds between different base pairs: blue, A-U pairs; red, G-C pairs; green: G-U pairs.

### 3.8. Molecular Phylogenetic Analysis of Mitoviral RdRps

Phylogenetic analysis of the RdRps of the three new mitoviruses and some representative mitoviruses and narnaviruses (members of the family *Narnaviridae*) was conducted using PhyML (Figure 6 and Table S1). According to Xu et al. [2015] [34], mitoviruses are divided into three major clades I, II and III, and the new mitoviruses were included in clade II accommodating OnuMV3a. Mitoviruses found from *Fusarium* species have been mostly grouped into clade III, however, FbMV1 together with recently reported Fusarium poae mitovirus 3 and 4 (FpMV3 and FpMV4 in Figure 6) [25]
exemplified a broad diversity of *Fusarium* mitoviruses. Taken together, these data suggest that the three viruses characterised in this study are strains of a new virus species *Fusarium boothii* mitovirus 1 in the genus *Mitovirus* in the family *Narnaviridae*. This is consistent with the species demarcation criteria within the genus *Mitovirus* (more than 90% sequence identity with RdRp refers strains) in the 9th report of ICTV [29].

Figure 6. Phylogenetic analyses of FbMV1. A phylogenetic tree was constructed based on the alignment of RdRp domain sequences of narnaviruses, mitoviruses and FbMV1 strains. The midpoint-rooted ML tree is shown. Dot at the nodes represents reliable aLRT values over 0.9 for branch supporting. The best model of the ML tree constructs is indicated at the upper left. Clade clustering is referred to a previous report by Xu et al. (2015) [34]. Blue arrowheads indicate mitoviruses isolated from the *Fusarium* species. Abbreviations of viral names and accession numbers are listed in Table S1. Red letters represent mitoviruses characterized in this study.

4. Discussion

Virus hunting from fungal hosts has been widely conducted, and recent advances in sequencing technologies has accelerated the discovery of novel types of mycoviruses [1,17,25,43,67]. The virus-like sequences mining from NGS-assisted metadata are highly useful in understanding the diversity and evolution of viruses although there is an argument as to what kind of information can validate the presence of “actual viruses” and their classification, even within ICTV [68]. Indeed, recent examples of virus characterization in *Rosellinia necatrix* and *S. sclerotiorum* revealed identification of novel mycoviruses including taxonomically unique individuals, and demonstrated effectiveness of NGS [69,70]. Hence, the accumulation of more virome data from African biological materials will facilitate our understanding of mycovirus diversity and may lead to discovery of unexpectedly unique populations of mycoviruses, as is the case for FbLFV1. There are many barriers in the establishment of effective virocontrol of herbaceous annual plant diseases, including FHB, as discussed by Pearson et al. [71]. Thus, it will be important to screen for virocontrol agents which have much better properties such as higher transmissibility and broader cross-species specificity.
Characterization of mycoviruses has significantly contributed to the taxonomy of viruses including the order Tymovirales. Those include creation of the genus Mycoflexivirus and the family Gammaflexiviridae (BVF); creation of the genus Deltaflexivirus and the family Deltaflexiviridae (SsDFV1, SsDFV2, FgDFV1 and soybean leaf-associated mycoflexivirus 1); creation of the genera Sclerodarnavirus and Botrexvirus under the family Alphaflexiviridae (SsDARV and BVX, respectively); and proposal of new genus Mycotymovirus under the family Tymoviridae (FgMTV1) (Table 1; Figure 7) [20,27,39–42,67]. This study additionally reports a novel member of the order Tymovirales that has a potential to further expand taxonomic diversity in the order.

![Figure 7. Comparison of genomic structures of the Tymovirales members and related unclassified viruses. Representative members of Tymovirales were selected as shown in Table 1 and their genomic structures are illustrated. Viral families, subfamilies and genera are shown in bold letters, and ‘new?’ represents a potential new taxon. Colour differentiated symbols are as indicated. The arrow on BVF replicase indicates potential readthrough translation. The blue triangle points to a known proteolytic cleavage site. Gray triangles at 3′-end of FbLFV1 and DOSV genomes indicate uncommon form that is not polyadenylated or tRNA-like structured.](image-url)
FbLFV1 showed similar characteristic to and distinct features from present five families in the order as follows. The careful assessment of phylogenetic relationship between FbLFV1 and other Tymovirales members revealed its close-yet-distinct correlations to the gammaflexivirus BVF and deltaflexiviruses (Figures 3 and 4). The presence of PHA03247 domain in these viruses further supported their evolutionally close association. However, the genome length and organization of FbLFV1 and BVF differed from each other (FbLFV1, 12,579 nt with single ORF vs. BVF, 6827 nt + poly-A tail with two ORFs) [39]. The absence of CP or CP domain of trichovirus/tymoviruses (see those of apple stem grooving virus, Diuris virus A and maize rayado fino marafivirus in Figure 7) in the FbLFV1 large polypeptide is highly possible and this feature is similar to deltaflexiviruses, an exceptional alphaflexivirus SsDRV and a tymo-like virus FgMTV1, highlighting the uniqueness of capsid-less nature of some mycoviruses, which is often observed (Figure 7) [20,27,38,41,42,45,67].

All Tymovirales viruses possess poly-A tail at the 3′ end with exceptions of those of the genus Tymovirus in the family Tymoviridae, which have TLSs, and of the genus Platypuvirus in the family Alphaflexiviridae, which have no obvious RNA structures (Figure 7) [72]. The 3′ end of FbLFV1 genome does not have a poly-A tail nor TLS, further discriminating FbLFV1 from established genera and families in the order Tymovirales (Figures 2 and 7). In contrast, 5′ ends of Tymovirales virus genomes poorly conserve characteristics, except for the cap structure, across the families. In the genus Tymovirus, 5′ UTRs generally carry one to four protonatable hairpins nearby initiation codons that are probably required for dicistronic expression of overlapping ORFs (Figure 7, Tymovirus) [73]. The stem-loop structure predicted in 5′ UTR of FbLFV1 was placed at 81 nt upstream of start codon (Figure 2C) and there is no potential overlapping ORF, suggesting a distinct role of the structure in multiplication process of FbLFV1.

The genome length of the members in the order Tymovirales ranges from about 5.9 to 9.0 kb [59] whereas the recently reported unassigned member, a Rhizoctonia virus RsFV1, has the largest genome (10,621 nt + poly-A tail) [38]. The discovery of FbFLV1 increases the maximum genome size in the order by about 2 kb. RsFV1 and FbFLV1 share a similar genomic structure: the large RNA genome of over 10 kb; the single large ORF encoding replicase; the presence of a functionally unknown region at N-terminus in the replicase; and a potential of capsid-less nature (Figure 7). The N-terminal regions contain a PHA03247 protein motif that is similar to a portion of the UL36 large tegument protein found in herpesviruses that has also been found from other mycoviruses BVF and SsDRV (Figure 7) [74]. Despite the above-mentioned similarity, RsFV1 is evolutionally closer to deltaflexivirus, whereas FbLFV1 is phylogenetically independent. In addition, a taxonomically floating virus, OMSV, appeared to be monophyletic, despite its apparently tymovirus (Tymoviridae)-like characteristics with respect to genome organisation and spherical virion structure (Figures 4 and 7) [44]. Thus, FbLFV1 and OMSV will become key viruses when taxonomic reconsideration within the order Tymovirales is required.

The strains of FbMV1 were found in three independent F. boothii isolates, and TcMV exhibited a high similarity to FbMV1 (35%). It has been reported that the TcMV RdRp has an AroE domain, which is one of the domains on AROM protein encoded by Saccharomyces cerevisiae [75]. AROM is a pentafunctional polypeptide that catalyses a sequence of five reactions in prechorismate polyaromatic amino acid biosynthesis, which is the former half of shikimate pathway. Although the function of the AroE-like domain in TcMV RdRp is unclear, similar sequences have been found in phylogenetically close relatives such as CpMV1 and OnuMV3a, as well as FbMV1 (Figure 5 and Figure S3), further supporting the involvement of FbMV1 in the mitovirus group of clade II. TcMV was identified in a basidiomycete fungus, Thanatephorus cucumeris (anamorph Rhizoctonia solani), which is phylogenetically very distant from ascomycetous Fusarium species. These points support an earlier hypothesis by Deng et al. [76] that mitoviruses have evolved not only through co-evolution with the viruses strictly confined within their host fungi and close relatives, but also through host-switching and adaptation to phylogenetically distant hosts via unknown routes of transmissions.

Overall, this study sequenced three isolates of a new mitovirus FbMV1 harbour in three independent F. boothii strains, as well as a novel tymo-like virus, FbLFV1, with the largest genome size
thus far described in the order Tymovirales. Detailed analyses of gene expression, potential polyprotein processing, particle formation and pathogenicity of FbLFV1 are the next challenges to understand the properties of this new taxon in the order Tymovirales. Biological property of FbLFV1 may be able to be assessed by protoplast-fusion method as reported for FgV1-DK21 [77].

Supplementary Materials: The following are available online at http://www.mdpi.com/1999-4915/10/11/584/s1, Table S1. List of viruses in the family Narnaviridae and related unassigned members used in the phylogenetic study shown in Figure 6. Figure S1. Phylogenetic analysis of partial TEF1-a gene sequences derived from Fusarium samples. Products from the genomic PCR targeting TEF1-a were subjected to direct sequencing and the resultant nucleotide sequences were aligned in MAFFT ver. 7 with publicly available sequences of various F. graminearum species complex members. The alignment was used for maximum likelihood phylogenetic analysis with the recommended model shown at upper left. Figure S2. Nucleotide sequence alignment of three FbMV1 strains Ep-BL13, Ep-BL14 and Ep-N28. The entire nucleotide sequences of FbMV1 strains aligned with MAFFT ver.7. Small letters, sequence of non-coding region; capital letters, replicate coding region. Figure S3. The putative AROM domains in mitovirus RdRps. The amino acid sequence of AroE domain of AROM protein encoded by S. cerevisiae, and corresponding sequences of FbMV1, TcMV and close relatives were aligned with MAFFT ver.7.

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