

## 1 **Supplementary Text S1**

2 **Dataset of reference viral genomes** Viral genome sequences and taxonomic  
3 information of viruses and their hosts are based on the GenomeNet/Virus-Host DB  
4 (Mihara, et al., 2016). The Virus-Host DB covers viruses with complete genomes stored  
5 in RefSeq/viral and GenBank. The sequence and taxonomic data are downloadable at  
6 <http://www.genome.jp/virushostdb/>. Viral genome sequences and taxonomic  
7 information used in ViPTree will be updated every few months following Virus-Host  
8 DB updates.

9 **Calculation of genomic similarity ( $S_G$ ) and proteomic tree** The original Phage  
10 Proteomic Tree (Rohwer and Edwards, 2002) tested multiple approaches and  
11 parameters to calculate genomic distance for evaluation of compatibility between the  
12 Phage Proteomic Tree and taxonomical system of the International Committee on  
13 Taxonomy of Viruses. In contrast, ViPTree performs proteomic tree calculation based  
14 on tBLASTx as reported previously by other several studies (Bellas, et al., 2015;  
15 Bhunchoth, et al., 2016; Mizuno, et al., 2013). Specifically, normalized tBLASTx scores  
16 ( $S_G$ ;  $0 \leq S_G \leq 1$ ) between viral genomes are calculated as described (Bhunchoth, et al.,  
17 2016). For simple calculation of  $S_G$  of segmented viruses, sequences of each segmented  
18 viral genome were concatenated into one sequence by inserting a sequence of 100  
19 ambiguous nucleotides (N) at each concatenation site. In addition, genome sequences  
20 longer than 100 kb are split into each 100 kb fragment just for faster tBLASTx  
21 calculation. Genomic distance is calculated as  $1 - S_G$ . Proteomic tree generation is  
22 performed by BIONJ (Gascuel, 1997) based on the genomic distance, using the R  
23 package ‘Ape’. The root of a tree is defined by mid-point rooting.

24 **Gene finding and gene annotation** Gene finding of uploaded sequences is performed

25 by GeneMarkS (Besemer, et al., 2001) without using its self-training method and with  
26 using an option “-p 0” to prohibit gene overlaps. Automatic functional annotations of  
27 predicted genes are performed by protein similarity searches against NCBI/nr using  
28 GHOSTX (Suzuki, et al., 2014).

29

### 30 **References**

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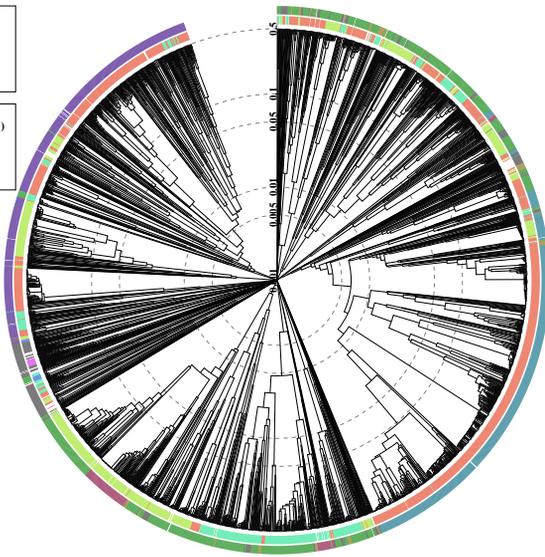
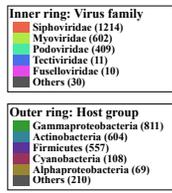
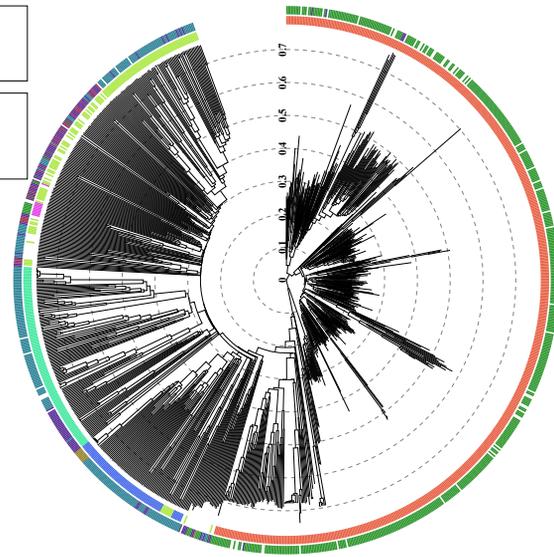
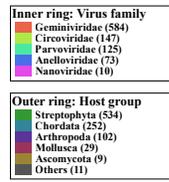
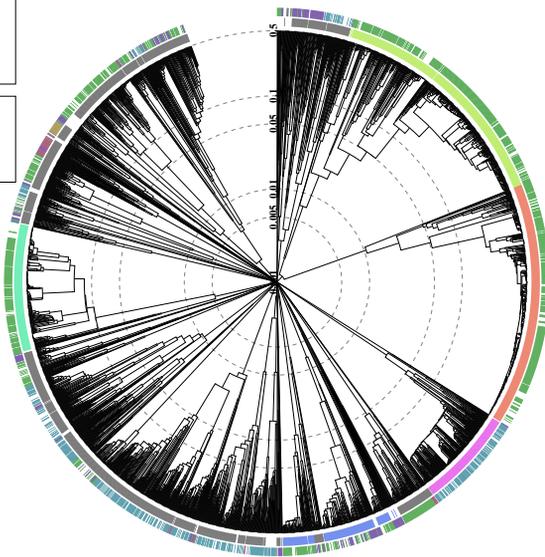
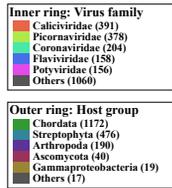
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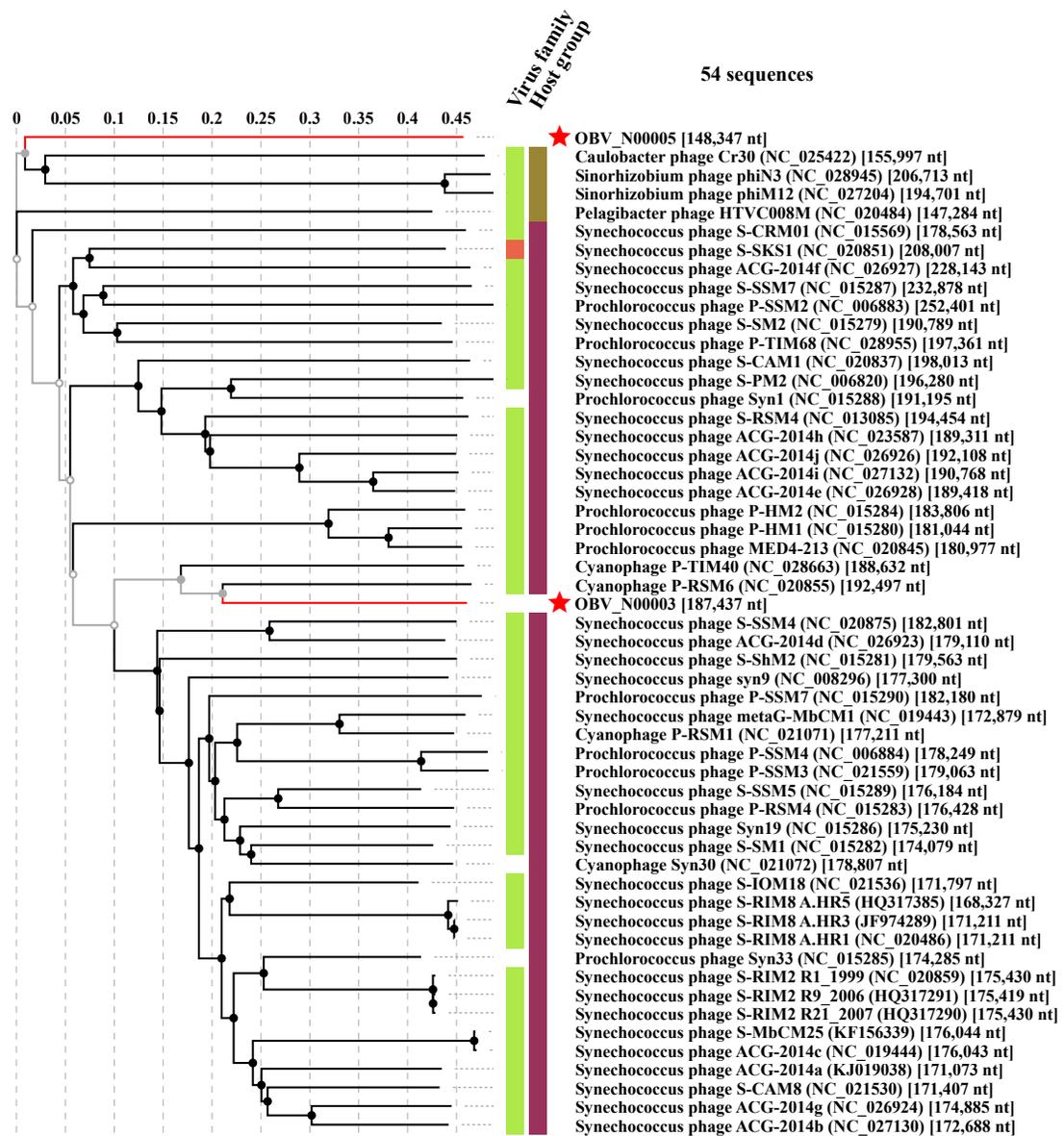
48

**A****B****C**

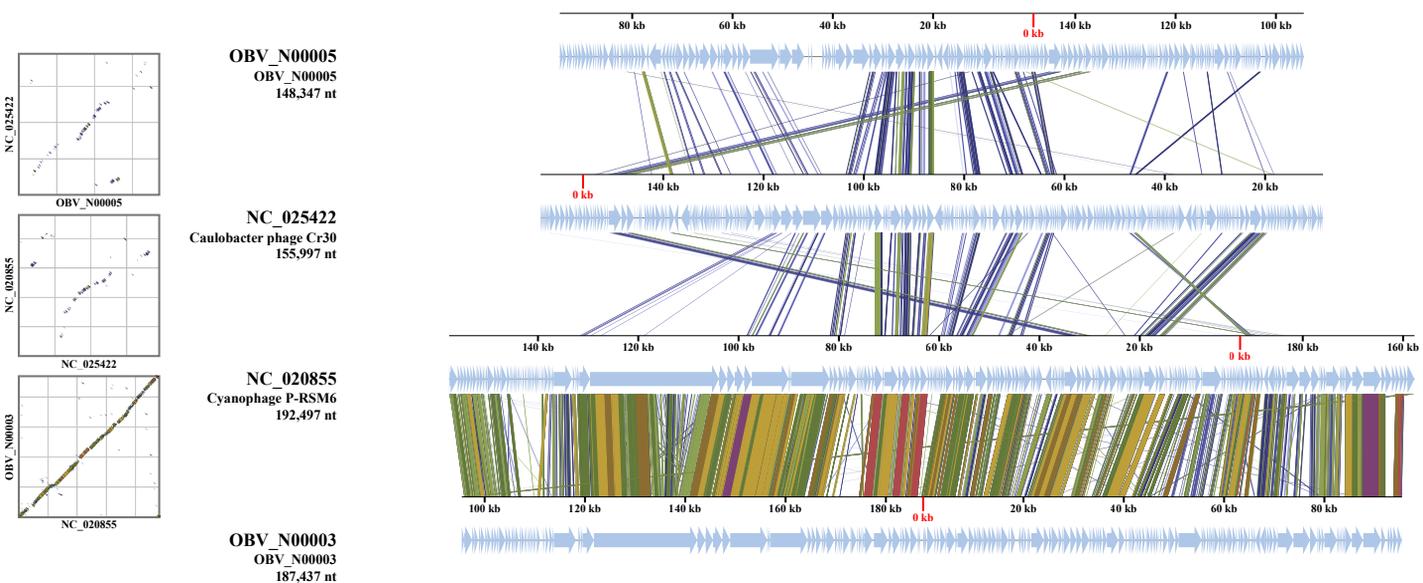
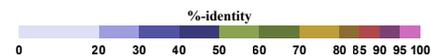
**Supplementary Figure S1.** Viral proteomic trees of reference genomes represented in the circular view. Color rings indicate virus families (inner rings) and host groups (at a level of phylum except for Proteobacteria; outer rings). These trees are calculated by BIONJ based on genomic distance matrixes, and mid-point rooted. Branch lengths are log-scaled (A, C) and linearly scaled (B). The sequence and taxonomic data is based on Virus-Host DB (release 78). (A) A tree of prokaryotic dsDNA viruses (2,384 genomes). (B) A tree of eukaryotic ssDNA viruses (1,013 genomes). (C) A tree of all ssRNA viruses (2,485 genomes).

**Left line: Virus family**  
 Myoviridae (48)  
 Siphoviridae (1)

**Right line: Host group**  
 Cyanobacteria (48)  
 Alphaproteobacteria (4)



**Supplementary Figure S2.** A Proteomic tree of dsDNA viruses represented in the rectangular view. This tree includes 52 reference viral genomes (48 *Myoviridae*, one *Siphoviridae*, and three unclassified viruses) and two uploaded viral genomes that are highlighted by red branches and stars (OBV\_N00003 and OBV\_N00005; BioProject: PRJDB4437). The tree is constructed by BIONJ based on genomic distance matrixes, and mid-point rooted. Branch lengths are linearly scaled. In the ViPTree server, where inner nodes of a tree are shown as filled circles, each of them links to a genomic alignment of sequences included in its subtree.



**Supplementary Figure S3.** An example of the genomic alignment view of four viral genomes (two reference and two environmental viral genomes) that are included in Supplementary Figure S2. Pairwise dot plots of these genomes are also shown. Colored lines in the alignment and the dot plots indicate tBLASTx results ( $E\text{-value} < 1e-2$ ). Grid lines in the dot plots indicate 40 kb intervals. Positions of each sequences are automatically adjusted (i.e., circular permuted and reverse stranded) for clear representation of colinearity between genomes.