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Characterisation data of simple sequence repeats of phages closely related to T7M



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ABSTRACT

Coliphages T7M and T3, *Yersinia* phage ϕ YeO3-12, and *Salmonella* phage ϕ SG-JL2 share high homology in genomic sequences. Simple sequence repeats (SSRs) are found in their genomes and variations of SSRs among these phages are observed. Analyses on regions of sequences in T7M and T3 genomes that are likely derived from phage recombination, as well as the counterparts in ϕ YeO3-12 and ϕ SG-JL2, have been discussed by Lin in "Simple sequence repeat variations expedite phage divergence: mechanisms of indels and gene mutations" [1]. These regions are referred to as recombinant regions. The focus here is on SSRs in the whole genome and regions of sequences outside the recombinant regions, referred to as non-recombinant regions. This article provides SSR counts, relative abundance, relative density, and GC contents in the complete genome and non-recombinant regions of these phages. SSR period sizes and motifs in the non-recombinant regions of phage genomes are plotted. Genomic sequence changes between T7M and T3 due to insertions, deletions, and substitutions are also illustrated. SSRs and nearby sequences of T7M in the non-recombinant regions are compared to the sequences of ϕ YeO3-12 and ϕ SG-JL2 in the corresponding positions. The sequence variations of SSRs due to vertical evolution are classified into four categories and tabulated: (1) insertion/deletion of SSR units,

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Abbreviations: SSR, simple sequence repeat

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(2) expansion/contraction of SSRs without alteration of genome length, (3) changes of repeat motifs, and (4) generation/loss of repeats.

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Specifications Table

Subject area	<i>Biology</i>
More specific subject area	<i>Genome evolution and sequence mutations</i>
Type of data	<i>Figure, tables</i>
How data was acquired	<i>Analysis of genomic sequences</i>
Data format	<i>Analyzed</i>
Experimental factors	<i>Genome sequences were retrieved from NCBI for analysis.</i>
Experimental features	<i>Software (ClustalW, IMEx) and manual analysis of the sequences, manual characterization and analysis</i>
Data source location	<i>National Chiao Tung University, Hsinchu, Taiwan</i>
Data accessibility	<i>Data are within this article.</i>

Value of the data

- Revealing different types of sequence changes of SSRs by vertical evolution of genomes.
- Detailed SSR distributions may aid in identifying broader patterns of phage evolution.
- Provides a guideline for classification of SSR variations in genome comparisons.
- Variations of SSRs in phages may be applied to phage typing.
- Assists researchers studying T7M, T3, ϕ YeO3-12, and ϕ SG-JL2 related phages in making sequence comparisons.

1. Data

Fig. 1 plots the distribution of SSR period sizes and motifs in the non-recombinant regions of the genomes of phages T7M, T3, ϕ YeO3-12, and ϕ SG-JL2. Table 1 illustrates differences in genomic sequences between T7M and T3. Tables 2 and 3 provide SSR counts, relative abundance, relative density, and GC contents in the complete genomes and non-recombinant regions for T7M, T3, ϕ YeO3-12, and ϕ SG-JL2. The four classes of SSR variations, (1) insertion/deletion of SSR units, (2) expansion/contraction of SSRs without alteration of genome length, (3) changes of repeat motifs, and (4) generation/loss of repeats, in T7M non-recombinant regions relative to counterpart regions of ϕ YeO3-12 and ϕ SG-JL2 are tabulated in Tables 4–9.

2. Experimental design, materials and methods

2.1. Genome sequences and recombinant regions

The genome sequence of T7M is in NCBI under the accession number GenBank: JX421753 [1]. Genome sequences of ϕ YeO3-12, ϕ SG-JL2, and T3 are acquired from GenBank accession numbers

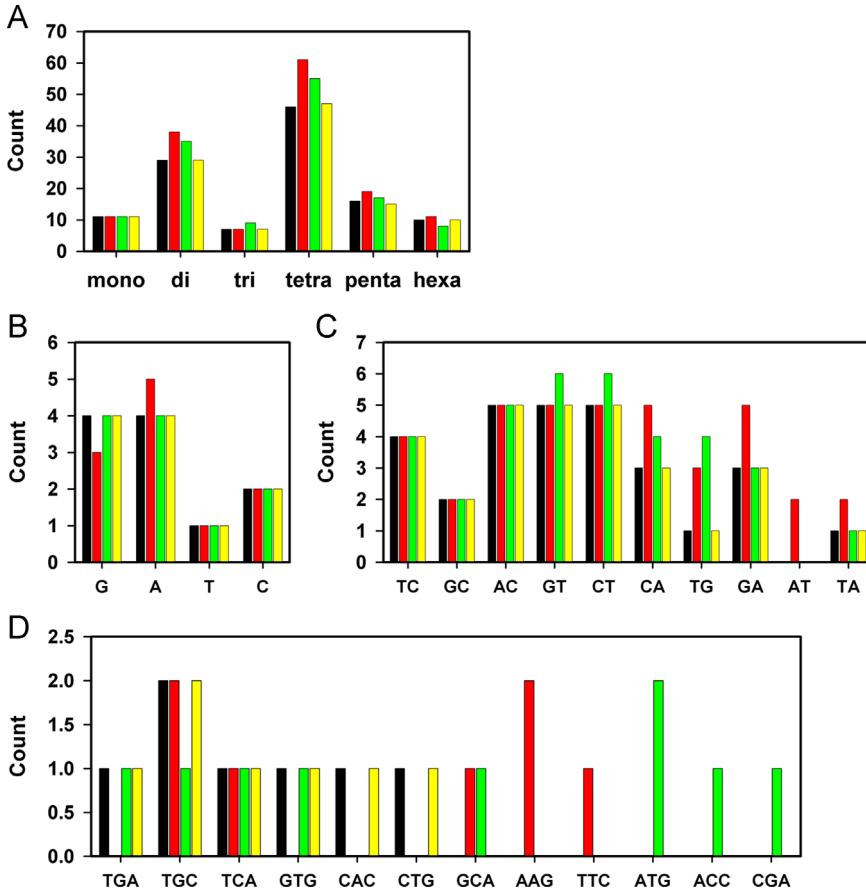


Fig. 1. The distribution of SSR period sizes and motifs in the non-recombinant regions of phage genomes. SSRs in the non-recombinant regions of T7M and T3 as well as the counterparts in ϕ YeO3-12 and ϕ SG-JL2 are compared. (A) Counts of mono- to hexanucleotide SSRs. (B) Mononucleotide motifs. (C) Dinucleotide motifs. (D) Trinucleotide motifs. T7M, black; ϕ YeO3-12, red; ϕ SG-JL2, green; T3, yellow.

Table 1

Difference in genomic sequences between T7M and T3.

T7M nt	T7M→T3 change	Location	Amino acid change ^a
26-27	Insertion of C	Terminal repeat	
9606-9607	Deletion of CG	Gene 3	GVRKVG→CTQGR
9627	Deletion of G	Gene 3	
9971	Deletion of G	Gene 3	WL→GV
9975-9976	Insertion of G	Gene 3	
22153	C→T	Gene 10B	T→I
22171	C→T	Gene 10B	T→I
23105	G→A	Gene 12	A→T
23156	C→A	Gene 12	L→I
24245	A→G	Gene 12	N→D
24659	G→A	Gene 12	G→R
25496-25497	Insertion of AGGGGGG	Between ϕ 13 and gene 13	
37998-37999	Insertion of C	Terminal repeat	

^a Change from T7M to T3 is shown by single letter codes of amino acids.

Table 2

SSR counts, relative abundance, and relative density in the complete genome and non-recombinant regions.

	Size bp	SSR count	RA ^a kb ⁻¹	RD ^b bp/kb	Size bp	SSR count	RA ^a kb ⁻¹	RD ^b bp/kb
	Complete genome				Non-recombinant regions ^c			
T7M	38202	192	5.0	39.7	25664	119	4.6	37.4
φYeO3-12	39600	207	5.2	40.8	26813	147	5.5	43.5
φSG-JL2	38815	195	5.0	39.3	26335	135	5.1	40.3
T3	38208	192	5.0	39.9	25670	119	4.6	37.6

^a Relative abundance; number of SSRs present in per kb of sequence.^b Relative density; the total length (bp) contributed by SSRs per kb of sequence.^c Excluding the two recombination regions in T7M and T3, and the counterpart regions in φYeO3-12 and φSG-JL2.**Table 3**Nucleotide compositions and GC contents of genomic sequences and SSRs in the complete genome versus non-recombinant regions^a of phages.

	T7M	φYeO3-12	φSG-JL2	T3
Complete genome				
% in complete genomic sequence				
A	26.4	26.2	26.0	26.4
T	23.7	23.2	23.2	23.7
G	26.5	27.0	27.0	26.5
C	23.4	23.6	23.8	23.4
GC	49.9	50.6	50.9	49.9
% in SSRs				
A	23.5 (-2.9)	25.2 (-1.0)	22.6 (-3.4)	23.4 (-3.0)
T	24.6 (1.0)	22.1 (-1.1)	23.8 (0.6)	24.5 (0.9)
G	26.0 (-0.5)	27.0 (0.0)	27.1 (0.1)	26.2 (-0.3)
C	25.8 (2.4)	25.7 (2.1)	26.5 (2.7)	25.9 (2.5)
GC	51.8 (1.9)	52.7 (2.2)	53.6 (2.8)	52.0 (2.1)
Non-recombinant regions^a				
% in non-recombinant regions of genome				
A	26.1	26.2	26.2	26.1
T	23.5	23.3	23.2	23.5
G	26.6	26.6	26.8	26.6
C	23.8	23.9	23.9	23.8
GC	50.4	50.5	50.6	50.4
% in SSRs				
A	22.8 (-3.3)	25.6 (-0.7)	22.0 (-4.2)	22.7 (-3.4)
T	24.6 (1.1)	22.0 (-1.3)	23.0 (-0.2)	24.5 (1.0)
G	25.7 (-1.0)	25.5 (-1.1)	27.7 (1.0)	25.9 (-0.7)
C	26.9 (3.1)	26.9 (3.1)	27.3 (3.5)	26.9 (3.2)
GC	52.6 (2.1)	52.4 (2.0)	55.0 (4.4)	52.9 (2.4)

Only the sequences of sense strands are considered. The number in parenthesis indicates the percent change compared to the complete genomes or the non-recombinant regions of genomes.

^a Excluding the two recombination regions in T7M and T3, and the counterpart regions in φYeO3-12 and φSG-JL2.

Table 4Indels of SSR repeat units in the non-recombinant regions of T7M and counterparts in ϕ YeO3-12 and ϕ SG-JL2.

T7M nt	Sequence in phage	
	T7M	ϕ YeO3-12
26	CCCCCC	CCCCC-
25497	GGGGGGGG	----GGGG
37998	CCCCCC	CCCCC-
T7M nt	Sequence in phage	
	T7M	ϕ SG-JL2
26	CCCCCC	CCCCC-
7704	ACACACAC	ACACAC-
25497	GGGGGGGG	----GGGG
37998	CCCCCC	CCCCC-

Table 5Repeat expansion/contraction without alteration of sequence length in the T7M non-recombinant regions and counterparts of ϕ YeO3-12 and ϕ SG-JL2.

T7M nt	Sequence in phage	
	T7M	ϕ YeO3-12
8183	<u>TCACACACGG</u>	TCTC <u>ACTG</u>
10777	<u>GTTGTG</u>	GCCT <u>GTG</u>
17930	<u>CACCACACCA</u>	CACCGC <u>ACCA</u>
26004	<u>GCGCGG</u>	<u>GCGCGAG</u>
T7M nt	Sequence in phage	
	T7M	ϕ SG-JL2
6218	<u>CTGATGATGATGG</u>	CTAAT <u>GATGATGG</u>
8183-8192	<u>TCACACACGG</u>	TCGA <u>ACACAG</u>
8525-8530	<u>CGGGG</u>	AAG <u>GGG</u>
11576-11584	<u>GTGGTG</u>	<u>GTGGTG</u>
17930-17940	<u>CACCACACCA</u>	CACCGC <u>ACCA</u>
26004-26010	<u>GCGCGG</u>	<u>GCGCGAG</u>

Repeat unit is underlined.

Table 6Repeat motif changes in the non-recombinant regions of T7M compared to counterpart regions of ϕ YeO3-12.

T7M nt	T7M	ϕ YeO3-12
1930	<u>ACGCAGGCAGCAGG</u>	<u>ACGCAGGCAGCAGG</u>
4125	<u>GTATCTATC</u>	<u>GTATATACC</u>
5919	<u>CAACGAAATGAAATC</u>	<u>CAACGAAACGAAATC</u>
6218	<u>CTGATGATGATGG</u>	<u>CTAATAATGATGG</u>
8178	<u>GTCACACACA</u>	<u>GCTACTCTCA</u>
11627	<u>CTTTCGTCCTCA</u>	<u>CGTTCGTTCTCA</u>
12316	<u>GGAGAAGGAGAAGGAGA</u>	<u>GAAGAAGGAGAAGGAGA</u>
12700	<u>AATCAATCAAGCAC</u>	<u>AGTCAATCACTCAC</u>
17742	<u>GACATAACATAG</u>	<u>GTCAATAGCATAG</u>
19669	<u>TGCTGCTGCCA</u>	<u>TGCAGCAGCAC</u>
20456	<u>CTGCTGCTGCTG</u>	<u>CGGCTCGGCTG</u>
21313	<u>CTGGCTGGTCTTGT</u>	<u>CTTGGTGGTCTGTT</u>
24066	<u>ACCCATACCCCTTCCTT</u>	<u>ACCCATACCCATCCGTT</u>
24935	<u>AAGGGTAGGGT</u>	<u>AAGGGTAGAGT</u>
26592	<u>TCCGGGGGA</u>	<u>TCAAAGGTA</u>

SSRs and surrounding sequences are listed. Repeats in ϕ YeO3-12 that have at least 3 copies for a mononucleotide or 2 copies for longer repeat periods, but different motifs from those in T7M, are considered. The repeat units with differing motifs between the two phages are underlined.

Table 7

SSR generation in the non-recombinant regions of T7M compared to counterpart regions of ϕ YeO3-12.

T7M nt	T7M	ϕ YeO3-12
1857	<u>GACCGACC</u>	GGATGAAC
7220	<u>GCTGACTGAA</u>	ACTGAGTGAA
9237	<u>CCAAGACAAGAA</u>	CCAAGATAAGAA
9965	<u>AGTGGCGTGGCT</u>	GGTGGAGTGGCT
10159	<u>GGCTGGCTGG</u>	GGCTGGTTAG
11106	<u>TCTGGICTGGTGGT</u> ^a	TCTGGTCTGGCCGT
11576	<u>GTGGTGGTG</u>	GTGGAGGCG
19278	<u>AATGCAATTGC</u>	AACTGCAATTGC
20211	<u>GCAGGCAG</u>	GCAGGCCG
20350	<u>TCAGGTCAAG</u>	TCCGGTCAGG
25654	<u>GCTGTGCTGTC</u>	GCTGTGTTGGC
25892	<u>GTCAATTTCAATT</u>	GTCAATTTCAACT
26016	<u>CAGACAGA</u>	CAGACCGA
36359	<u>CCAACCAAC</u>	TCAACCGAC
37140	<u>GCGTITAGCGTTAG</u>	GCGTITAGCATTGG

The newly generated repeat unit in T7M is underlined. The repeat sequence displays at least 3 iterations of a mononucleotide repeat unit or 2 contiguous iterations of a di- to hexanucleotide repeat unit. Repeat sequences in ϕ YeO3-12 that are also present in T7M are not considered.

^a The sequence has a newly generated GGT repeat in addition to a motif change CTGGT, and both are underlined in this table.

Table 8

Repeat motif changes in the non-recombinant regions of T7M compared to counterpart regions of ϕ SG-JL2.

T7M nt	T7M	ϕ SG-JL2
4125	<u>GTATCTATC</u>	GTGTCTACC
5088	<u>AGCTGCTGGCTGCTG</u>	<u>AGCTGCTAGCTGCTG</u>
11627	<u>CTTTCGTCGGTCA</u>	<u>CGTTCGTTCTGTC</u>
12316	<u>GGAGAAGGAGAAGGAGA</u>	<u>GAAGAAGGAGAAGGAGA</u>
17593	<u>CGATGACGATGA</u>	<u>CGATGATGACGA</u>
17742	<u>GACATAACATAG</u>	<u>GTCATAGCATAG</u>
19669	<u>TGCTGCTGCCA</u>	<u>TGCAGCAGCAC</u>
20456	<u>CTGCTGCTGCTG</u>	<u>CGGCTGCGGCTG</u>
21313	<u>CTGGCTGGTCTTGT</u>	<u>CTGGCTGGICTGGT</u>
24066	<u>ACCCATACCTTCCTT</u>	<u>ACCCATACCCATCCTT</u>
24935	<u>AAGGGTAGGGT</u>	<u>AGGGTAGAGT</u>
26592	<u>TCCGGGGGA</u>	<u>TCAAAGGTA</u>
37648	<u>TACTTACTGCT</u>	<u>TACTTCTGCT</u>

SSRs and surrounding sequences are listed. Repeats in ϕ SG-JL2 that have at least 3 copies for a mononucleotide or 2 copies for longer repeat periods, but different motifs from those in T7M, are considered. The repeat units with differing motifs between the two phages are underlined.

GenBank: AJ251805 [2], GenBank: NC_010807 [3], and GenBank: AJ318471 [4], respectively. Sequences were aligned by ClustalW [5], and differences between phages are compared. The T7M sequence nt 13245–16687 and 26695–35789 align to T3 nt 13243–16685 and 26700–35794, respectively, and likely arise from a recombination between a ϕ YeO3-12-like phage and a T7-like phage, as suggested for T3 [4]. These regions and the counterparts in ϕ YeO3-12 and ϕ SG-JL2 are referred to as recombinant regions, and the rest of the genomes are referred to as non-recombinant regions [1].

Table 9SSR generation in the non-recombinant regions of T7M compared to counterpart regions of ϕ SG-JL2.

T7M nt	T7M	ϕ SG-JL2
1930	<u>ACGCAGGCAGCAG</u>	ACGCAGGCCAAGG
4996	<u>GGCTGGCTATAT</u>	GGCTGGTTATAT
5582	<u>AACCTGAACCTG</u>	AAGCTGAACCTA
5731	<u>ACTTTCITTA</u>	long ^a
5919	CAACGAAATGAAATC	long ^a
8178	<u>GTCACTCACA</u>	GTCACTCGAA
9237	<u>CCAAGACAAGAA</u>	CCAAGATAAGAA
9965	<u>AGTGGCGTGGCT</u>	GGTGGAGTGGCT
10159	<u>GGCTGGCTGG</u>	GGCTGGTTAG
11106	<u>TCTGGTCTGGTGGT^b</u>	TCTGGTCTGGCGGT
12700	<u>AATCAATCAAG</u>	AGTCAATCACC
16958	<u>ATCAAGCAAGG</u>	ATTAAGCAAGG
19278	<u>AATTGCAATTGC</u>	AACTGCAATTGC
20211	<u>GCAGGCAG</u>	GCAGGCCG
20350	<u>TCAGGTCAGG</u>	TCCGGTCAGG
25654	<u>GCTGTGCTGTC</u>	GCTGTGTTGGC
25892	<u>GTCAAITTCAAATTA</u>	GTCAAITCCAATTA
26016	<u>CAGACAGA</u>	CAGACCGA
26335	<u>CAAGTCAAGTC</u>	CGAGTCAAGTC
36359	<u>CCAACCAAC</u>	TCAACCGAC
37140	<u>GCGTTAGCGTTAG</u>	GCGTTAGCAITGG

The newly generated repeat unit in T7M is underlined. The repeat sequence consists of at least 3 iterations of a mononucleotide or 2 contiguous iterations of a di- to hexanucleotide. Repeat sequences in ϕ SG-JL2 that are also present in T7M are not considered.

^a The sequence is longer in ϕ SG-JL2 and does not align well to that of T7M in this region.

^b The sequence has a newly generated GGT repeat in addition to a motif change CTGGT, and both are underlined in this table.

2.2. Simple sequence repeats

Simple sequence repeats were searched in phage genomes or non-recombinant regions by IMEX [6]. Unless otherwise specified, the minimum repeat units for mono- to hexanucleotide were 5, 3, 3, 2, 2. Repeats sequences were not standardized.

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Transparency document. Supplementary material

Transparency data associated with this article can be found in the online version at <http://dx.doi.org/10.1016/j.dib.2016.06.035>.

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