



Title	Receptor for Phage $\phi$ 80 on Escherichia coli K12
Author(s)	Matsushiro, Aizo; Hasegawa, Noriyoshi
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Receptor for Phage  $\phi 80$  on *Escherichia coli* K12

Phage  $\phi 80$  showed very strange behavior towards various derivatives of *Escherichia coli* K12. The results obtained are summarized in Table 1.

Table 1. The Behavior of K12 Strains for Phage  $\phi 80$

"Sensitive"	"Resistant"
K12 wild (F <sup>-</sup> , prototroph, $\lambda^+$ , S <sup>s</sup> )	C600 (F <sup>+</sup> , TLB <sub>1</sub> <sup>-</sup> , Lac <sup>-</sup> , $\lambda^-$ , S <sup>r</sup> , T <sup>r</sup> <sub>1,5</sub> )
58-161 (F <sup>+</sup> , M <sup>-</sup> , $\lambda^+$ , S <sup>s</sup> )	W1177 (F <sup>-</sup> , TLB <sub>1</sub> <sup>-</sup> , Lac <sup>-</sup> , $\lambda^+$ , S <sup>r</sup> , T <sup>r</sup> <sub>1,5</sub> )
W-3100 (F <sup>-</sup> , prototroph, $\lambda^+$ , S <sup>s</sup> )	W678 (F <sup>-</sup> , TLB <sub>1</sub> <sup>-</sup> , Lac <sup>-</sup> , $\lambda^+$ , S <sup>r</sup> , T <sup>r</sup> <sub>1,5</sub> )
W-3110 (F <sup>-</sup> , prototroph, $\lambda^-$ , S <sup>s</sup> )	P678 (F <sup>-</sup> , TLB <sub>1</sub> <sup>-</sup> , Lac <sup>-</sup> , $\lambda^-$ , S <sup>r</sup> , A <sub>2</sub> <sup>r</sup> , T <sup>r</sup> <sub>1,5</sub> )
Hfr H (Hfr, B <sub>1</sub> <sup>-</sup> , $\lambda^-$ , S <sup>s</sup> )	112-12 (F <sup>-</sup> , C <sup>-</sup> , H <sup>-</sup> , $\lambda^-$ , S <sup>s</sup> , T <sup>r</sup> <sub>1,5</sub> )
Y70 S (F <sup>-</sup> , TLB <sub>1</sub> <sup>-</sup> , Lac <sup>-</sup> , $\lambda^+$ , S <sup>s</sup> )	

F: compatibility factor. M, T, L, B<sub>1</sub>, C, H: auxotrophic markers for methionine, threonine, leucine, thiamine, cysteine, histidine. Lac: fermentation of lactose. S, A<sub>2</sub>, T<sub>1,5</sub>: resistance to streptomycin, azide, phage T<sub>1</sub> and T<sub>5</sub>.  $\lambda^+$ : lysogenic for phage  $\lambda$ .

First this phage attacks K12 independently of  $\lambda$  and the F factor. Second, strains carrying T<sup>r</sup><sub>1,5</sub> marker are mostly resistant to phage  $\phi 80$ .

Strains 58-161 F<sup>+</sup> and W-1177 F<sup>-</sup> were used in Lederberg's classical cross experiments. The former is originally sensitive to  $\phi 80$  but the latter is resistant. Strain 58-161 is an auxotrophic mutant; F<sup>+</sup>, M<sup>-</sup>, T<sup>+</sup>, L<sup>+</sup>, B<sub>1</sub><sup>+</sup>, Lac<sup>+</sup>, S<sup>s</sup>, T<sup>s</sup><sub>1,5</sub>,  $\phi 80^s$ . Strain W-1177 is a complementary auxotrophic mutant; F<sup>-</sup>, M<sup>+</sup>, T<sup>-</sup>, L<sup>-</sup>, B<sub>1</sub><sup>-</sup>, Lac<sup>-</sup>, S<sup>r</sup>, T<sup>r</sup><sub>1,5</sub>,  $\phi 80^r$ . An attempt to determine the gene locus controlling the receptor for this phage was made by cross experiments on these strains. Recombinants were selected on synthetic media supplemented with 100  $\mu$ g/ml of

Table 2. The Segregation of Non-selective Characters (Lac and T<sub>1,5</sub>) in Recombinants between 58-161  $\times$  W-1177

Parents	Recombinants			
	Lac <sup>-</sup> $\phi 80^r$	Lac <sup>+</sup> $\phi 80^s$	Lac <sup>-</sup> $\phi 80^r$	Lac <sup>-</sup> $\phi 80^s$
58-161/F <sup>+</sup> $\times$ W-1177/F <sup>-</sup> (TLB <sub>1</sub> <sup>+</sup> , M <sup>-</sup> ) (TLB <sub>1</sub> <sup>-</sup> , M <sup>+</sup> )				
Lac <sup>+</sup> $\phi 80^r$ $\times$ Lac <sup>-</sup> $\phi 80^s$	67 (37.8%)	5 (2.8%)	52 (29.3%)	53 (29.9%)
Lac <sup>-</sup> $\phi 80^s$ $\times$ Lac <sup>-</sup> $\phi 80^r$	2 (2.0%)	37 (37.0%)	31 (31.0%)	30 (30.0%)

streptomycin. From the data shown in Table 2, the genetic factor seems to be located between Lac and T-L. The locus controlling the T<sub>1,5</sub>-receptor is also probably located here.

From these results it seems that these two loci (T<sup>r/s</sup><sub>1,5</sub> and  $\phi 80^{r/s}$ ) are closely

linked to each other or are even identical.

To study this an investigation was made of whether the  $T^s_{1,5} \rightarrow T^r_{1,5}$  mutation followed by a  $\phi 80^s \rightarrow \phi 80^r$  mutation.  $T^s_{1,5}$  strains were all sensitive to  $\phi 80$ , but  $T^r_{1,5}$  mutants selected from these  $T^s_{1,5}$  strains were all also resistant to  $\phi 80$ . The  $\phi 80^s \rightarrow \phi 80^r$  mutation was followed, as might have been expected, by a  $T^s_{1,5} \rightarrow T^r_{1,5}$  mutation (Table 3).

Table 3. The Cross-resistance Interrelationship between  $\phi 80$  and  $T_1, T_5$

	K12	K12/ $\phi 80$	K12/ $T_{1,5}$	K12/ $T_1$ - $tryp$
$\phi 80$	+	—	—	—
$T_1$	+	—	—	—
$T_5$	+	—	—	+

+: sensitive, -: resistance

Recently Yanofsky *et al.* (1959)<sup>1)</sup> have described  $T^r_1$   $tryp^{**}$  mutants, which are rare among  $T^r_1$  isolates in strain K12. All the ten  $T^r_1$   $tryp^-$  mutants isolated by Yura (Virus Institute, Kyoto Univ.) and the author were also resistant to phage  $\phi 80$ . Though these  $tryp$  deletion mutants were found to carry the various  $tryp$  region deletion (Yura, personal communication), all were resistant to  $T_1$  as well as  $\phi 80$  but were sensitive to  $T_5$ .

$T_1$   $tryp$  is independent region upon  $T_{1,5}$  marker on the K12 chromosome. However, from the fact that all the variety of  $T^r_1$  strains are always also the  $\phi 80^r$  strain, it is clear that the  $T_1$  phage receptor gene is not only the same as the receptor gene for  $\phi 80$  phage but also the receptor site on the cellular surface for both phages is common. This receptor site may be controlled by two different genes in the  $T_{1,5}$  and  $T_1$   $tryp$  regions.

Finally it is very interesting that one of the genes controlling the adsorption of this phage is closely linked with the  $tryp$  region, because the  $tryp$  locus is itself also closely linked to this  $\phi 80$  prophage locus on the K-12 chromosome, as will be described elsewhere.

#### REFERENCE

1) Yanofsky, C. and Lennox, E. S. (1959). Transduction and recombination study of linkage relationships among the genes controlling tryptophan synthesis in *Escherichia coli*. *Virology* **8**, 425-447.

AIZO MATSUSIRO  
NORIYOSHI HASEGAWA

Department of parasitology,  
The Research Institute for Microbial Diseases,  
Osaka University, Osaka  
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\*\*  $tryp$ : auxotrophic marker for tryptophan.