Narrow-host-range IncP Plasmid pH502-1 Lacks a Complete IncP Replication System

By CHRISTOPHER A. SMITH and CHRISTOPHER M. THOMAS
Department of Genetics, University of Birmingham, PO Box 363, Birmingham B15 2TT, UK

(Received 23 February 1987; revised 13 April 1987)

Plasmid pH502-1 shows incompatibility only towards members of the IncP group, but has a narrower host range than typical members of that group. In contrast to other IncP plasmids its replication was not affected by a high-copy-number plasmid carrying the replication origin (oriV) of IncP plasmid RK2. Southern blotting of pH502-1 revealed homology to oriV, consistent with its incompatibility phenotype, but no homology to trfA, the essential replication gene of RK2. Thus it is probable that pH502-1 does not possess a functional IncP replication system, accounting for its restricted host range. A restriction map of pH502-1 was constructed and the mercury-resistance determinant was localized to Tn735, which also carries the trimethoprim-resistance determinant and is related to Tn2I. The presence of a korB-like function on pH502-1 was also demonstrated.

INTRODUCTION

Plasmids which cannot be stably maintained together in the same host cell line are said to be incompatible and to belong to the same incompatibility group; such groups often reflect the possession by their members of related replication systems. Although most plasmids belong to only one incompatibility group, some are members of more than one group, being incompatible with members of two or more groups of plasmids, other members of which are incompatible only with the members of their own group. Such plasmids may possess all or part of two or more replication systems.

Typical plasmids of Escherichia coli incompatibility group P (equivalent to the Pseudomonas group IncP-1) have a broad host range, being capable of self-transfer to and maintenance in members of most Gram-negative genera. These plasmids have been divided into two sub-groups which are relatively distantly evolutionarily related: IncPα, including the indistinguishable isolates RK2, RP1, RP4, R18 and R68, which are the best-studied members of the IncP group, and IncPβ, including R751, R906 and R772 (Chikami et al., 1985; Smith & Thomas, 1987). Although the control circuits of RK2 are complex, only two loci are essential for vegetative replication, oriV, the origin of replication, which is closely linked to major determinants of P-group incompatibility, and trfA, encoding a trans-acting product (Thomas et al., 1985). Incompatibility is also expressed by the region of RK2 encoding korA and korB. These loci are involved in the regulation both of trfA and of a number of kil genes which cannot be cloned in their absence (Figurski et al., 1982; Thomas & Hussain, 1984). Thus the IncP grouping appears to result from the possession of related replication systems by its members.

Certain plasmids are incompatible with IncP plasmids but have much narrower host ranges than the typical members of the group. Examples of IncP plasmids unable to establish themselves in certain hosts when tested for self-transfer include: pMU700–pMU707, in Pseudomonas and Proteus (Grant et al., 1980); pH502, in Acinetobacter, Proteus, Providencia, Pseudomonas and Serratia, and its derivative pH502-1, in Acinetobacter and Pseudomonas (Nugent et al., 1982); and pAV1, which is restricted to certain strains of Acinetobacter calcoaceticus (Hinchliffe & Vivian, 1980). Of these, pMU700–pMU707 and pH502 are also...
incompatible with IncI plasmids, while pAV1 has only been tested for incompatibility with plasmids of group P. In contrast, pHH502-1, which arose from pHH502 by a repeatable spontaneous event apparently involving both deletion and rearrangement, lacks IncI characteristics and does not show incompatibility with members of any other known group (Nugent et al., 1982).

We have investigated pHH502-1, because we are interested in the basis for the broad host range of typical IncP plasmids and hoped that the differences between its replication system and those of other IncP plasmids might reveal features of importance in determining plasmid host range. Here we report evidence that this plasmid both possesses a non-IncP replication system and lacks a complete IncP replication system. We also present a restriction map of pHH502-1 and show that Tn735 carries the Hgr determinant as well as the Tp' determinant of the plasmid and that this transposon is related to Tn21.

METHODS

Bacterial and plasmid strains. The bacterial host strain used for pHH502-1 was MV10, a trpE5 derivative of E. coli K12 strain C600 (thr-1 leu-6 thi-1 lacY1 supE44 tonA22). pHH502-1 (Nugent et al., 1982) confers resistance to trimethoprim and to mercuric salts, and low-level resistance to sulphonamides. RK2 (Ingram et al., 1973) confers resistance to penicillin, kanamycin and tetracycline. R772 (Coetzee et al., 1979) confers resistance to kanamycin and carries an inactive mercury-resistance determinant (Smith & Thomas, 1987). R906 (Hedges et al., 1974) confers resistance to oxacillin, streptomycin, sulphonamides and mercuric salts. R751 (Jobanputra & Datta, 1974) confers resistance to trimethoprim. References to plasmids constructed by in vitro recombination are given below.

Testing for sensitivity to P-group incompatibility. MV10(pHH502-1) was transformed with the ColE1-derived plasmids pCT7 and pCT8, which both confer resistance to kanamycin and are isogenic except that pCT7 carries the oriV region of RK2 (Thomas et al., 1980). In each case 100 kanamycin-resistant colonies were tested for trimethoprim resistance, indicating retention of pHH502-1, as described for R751 (Smith & Thomas, 1985).

Testing for KorA and KorB phenotypes. Attempts were made to transform MV10(pRK259.2) and MV10(pHH502-1) with the KilA+ plasmid pCAS54, selecting for penicillin resistance, and with the KilB+ plasmid pCAS51, selecting for tetracycline resistance. Transformant colonies are expected only if a functional copy of the korA or korB gene, respectively, is already present in the recipient; both genes are carried by pRK259.2 (Smith & Thomas, 1983).

Southern blotting. Isolation of plasmid DNA, restriction analysis, agarose gel electrophoresis, preparation and radio-labeling of probe fragments, transfer of DNA fragments to nylon membranes, hybridization and autoradiography were done as described by Smith & Thomas (1987). The RK2-derived probe fragments carrying the oriV region and the C- and N-terminal parts of the trfA gene correspond respectively to fragments 5, 6 and 7 of Smith & Thomas (1987), obtained from EcoRI + HindIII digests of plasmids pCAS220, pCAS226 and pCAS228. The probe fragments derived from Tn501 were the two internal EcoRI fragments of the transposon, corresponding to fragments 9 and 10 of Smith & Thomas (1987).

RESULTS AND DISCUSSION

Restriction map

We constructed a restriction map of pHH502-1 (Fig. 1), which confers resistance to trimethoprim (Tp') and mercuric ions (Hgr') and a partial resistance to sulphonamides, designated [Su'] by Nugent et al. (1982). Nugent et al. (1982) reported that three of the EcoRI fragments of pHH502-1, with sizes of 3·2, 2·8 and 0·85 MDa (4·8, 4·2 and 1·3 kb), are contained within Tn735, a transposable element of 9·85 MDa (15 kb) which carries the Tp' determinant of the plasmid. These fragments, together with two small EcoRI fragments not previously reported, span coordinates 13·1 kb to 23·7 kb of our map. The distribution of cleavage sites on the map of pHH502-1 is non-uniform, with 16 of the 25 sites lying within Tn735 (Fig. 1). Such clustering of restriction sites in the regions encoding resistance determinants is also found in the case of typical IncP plasmids (Smith & Thomas, 1987).

Location of mercury-resistance determinant

In order to locate the mercury-resistance determinant of pHH502-1 we used the two internal EcoRI fragments of the mercury-resistance transposon Tn501 as probes against Southern blots of digests of the plasmid (Fig. 2a, b). Each of these fragments carries part of the mercury-resistance operon (mer) of Tn501 (Brown et al., 1983). The regions of overlap between the
Fig. 1. Restriction map of pHH502-1. The region between coordinates 8 kb and 26 kb is expanded for greater clarity, and the probable end-points of Tn735 are indicated. The regions containing homology to the probes used are indicated as follows: RK2 oriV by hatching; Tn301 shorter fragment (mer) by cross-hatching; Tn501 larger fragment (mer and tnp) by stippling.

Fig. 2. Southern blots of digests of pHH502-1, R772 and R906. Probe fragments were derived from: (a) and (b) the shorter and longer EcoRI fragments of Tn301 respectively; (c) the oriV region of RK2; (d) the trfA region of RK2. The lanes contained: (1) pHH502-1 digested with SalI + XhoI; (2) pHH502-1 digested with BglII; (3) pHH502-1 digested with EcoRI; (4) R772 digested with PstI; (5) R772 digested with EcoRI; (6) R906 digested with SstII; (7) R906 digested with EcoRI. The sizes (kb) of the fragments in each digest showing homology to each probe were as follows. (1): (a) 17.5; (b) 27 and 17.5; (c) 17.5; (d) none. (2): (a) 9.0; (b) 31 and 9.0; (c) 17.0; (d) none. (3): (a) 3.6; (b) 7.2 and 4.2; (c) 6.1 and 5.9; (d) none. (4): (c) 3.6; (d) 27. (5): (c) 29; (d) 21.5. (7): (c) 27.5; (d) 9.7.
fragments to which each probe hybridized in the digests of pHH502-1 are indicated in Fig. 1. The two probes hybridized to adjacent EcoRI fragments at coordinates 9.9 kb to 13.1 kb and 13.1 kb to 17.2 kb, indicating that the mer genes of the plasmid are in this region, and share a common EcoRI site with those of Tn501. Since one of these EcoRI fragments is internal to Tn735 (Nugent et al., 1982), the transposon must carry the Hg\textsuperscript{r} determinant as well as the Tp\textsuperscript{r} determinant of pHH502-1.

Relatedness of Tn735 to Tn21

The larger of the two Tn502-derived probe fragments hybridized to two regions of pHH502-1 (Fig. 2 b): to the mer region and to a region at the other end of Tn735 (Fig. 1). Since this probe fragment contains the transposition (tnp) genes of Tn501 as well as part of the mer operon (Diver et al., 1983), it appeared possible that the second region of hybridization on pHH502-1 was due to homology between the tnp genes of Tn735 and Tn501. This hypothesis was confirmed by the distribution of BamHI, EcoRI and HindIII sites between coordinates 13.3 kb and 23.7 kb within Tn735. This distribution is identical with that found in the tnp region of Tn21 (and of several other closely related transposons), to which the tnp region of Tn501 is related (Tanaka et al., 1983; Kratz et al., 1983; Diver et al., 1983). Fig. 1 shows the end-points predicted for Tn735 if its outer ends correspond to those of Tn21 and its relatives. These end points give a total size of 15.4 kb for Tn735, in agreement with the size reported by Nugent et al. (1982). The Tp\textsuperscript{r} determinant carried by Tn735 presumably lies within the EcoRI fragment between coordinates 17.2 kb and 22 kb, which does not show homology to Tn501. Although there is no direct evidence concerning the location of the [Su\textsuperscript{r}] determinant of pHH502-1, it is noteworthy that the distribution of SalI, BamHI, BglII and HindIII sites between coordinates 17.4 kb and 19.2 kb within Tn735 is similar to that within the Su\textsuperscript{r} determinant of Tn21 and its relatives (Kratz et al., 1983).

Homology to the oriV region

Fig. 2(c) shows a Southern blot of digests of pHH502-1 and the IncP\(\beta\) plasmids R772 and R906 probed with a fragment carrying the oriV region of the IncP\(\alpha\) plasmid RK2. The region of overlap between the fragments to which this probe hybridized in the digests of pHH502-1 is indicated in Fig. 1. The smaller of the two bands seen in the EcoRI digest is believed to be a partial EcoRI\* product of the larger.

Lack of homology to the trfA gene

Fig. 2(d) shows the same digests as Fig. 2(c), probed with a fragment carrying the N-terminal part of the tefA gene of RK2. Although the autoradiograph was deliberately overexposed (as can be seen from the bands in the lanes containing digests of IncP\(\beta\) plasmids R772 and R906), no homology between pHH502-1 and this probe was detected; the same result was obtained using a probe-fragment carrying the remainder of the tfeA gene of RK2 (data not shown). This implies that if pHH502-1 possesses a tefA-like gene its sequence has diverged from that of the tefA gene of RK2 too far for the homology to be detected under the hybridization conditions used. Since under the conditions used homology was easily detected between the oriV regions of RK2 and R751 (Smith & Thomas, 1987), which show only 65\% overall sequence homology (Smith & Thomas, 1985), it is more probable that pHH502-1 lacks a tefA-like gene. Since the tefA gene is essential for a functional IncP replication system, the lack of this gene would imply that pHH502-1 possesses a replication system of another incompatibility group.

Insensitivity to IncP incompatibility

When the oriV region of RK2 is inserted into a high-copy-number vector replicon of another incompatibility group it exhibits strong incompatibility against IncP plasmids. Because the maintenance of the vector replicon is not affected by the presence of an IncP plasmid, this incompatibility is unidirectional, resulting in the preferential elimination of the IncP plasmid. We tested the effect of transformation with such a plasmid, pCT7, on resident pHH502-1 and observed 100\% retention of pHH502-1. In contrast, similar experiments showed only 4\%
Plasmid pHH502-1

retention of pRK24 (a kanamycin-sensitive derivative of RK2) and complete loss of IncPβ plasmid R751, but 100% retention of each plasmid following transformation with pCT8, an otherwise isogenic plasmid lacking the oriV region (Thomas et al., 1980; Smith & Thomas, 1985). This result confirmed that pHH502-1 possesses a replication system (of an unknown incompatibility group) which is not susceptible to the oriV-associated IncP determinants. This is consistent with the observation that transfer of pHH502-1 to a strain carrying RP4 results in the elimination of the resident plasmid from a high proportion of clones, while transfer of RP4 does not result in the elimination of resident pHH502-1, but gives unstable clones carrying both plasmids (Nugent et al., 1982).

Presence of a korB-like gene

Figurski et al. (1982) showed that a range of diverse IncP plasmids possess genes which can substitute for korA and korB in the control of the potentially host-lethal kilA and kilB loci of RK2. We tested pHH502-1 for the ability to enable the establishment of compatible plasmids carrying the kilA and kilB loci of RK2. Following transformation of MV10(pHH502-1) with korB-dependent plasmid pCAS51, healthy colonies were obtained on medium selective for the incoming plasmid, indicating that pHH502-1 possesses a korB-like gene. However, transformation with korA-dependent plasmid pCAS54 gave equivocal results; no colonies were obtained from MV10, small slow-growing colonies from MV10(pHH502-1) and healthy colonies from MV10(pRK259.2), the positive control. It is unclear whether or not this result indicates that pHH502-1 possesses an active korA-like gene.

Evolutionary implications

It is likely that pHH502-1 consists of a replicon of unknown incompatibility group carrying some part of an IncP replication system, presumably as a result of a replicon fusion followed by one or more deletion events. The IncP loci present include the oriV region and the korB gene, but not the trfA gene, essential for replication. This provides a clear, although trivial, explanation for the restricted host range of pHH502-1 compared to that of typical IncP plasmids. Since Nugent et al. (1982) found that the pili determined by pHH502-1 differ from those determined by other IncP plasmids serologically, morphologically and in the bacteriophage sensitivity which they confer, it is probable that the regions of the plasmid responsible for conjugal transfer did not form part of the IncP plasmid ancestor of pHH502-1.

Since pHH502, from which pHH502-1 is derived, also has a restricted host range it is probable that it also lacks a functional IncP replication system. The narrow host range of pMU700–pMU707 and of pAV1 may also result from the lack of a complete functional IncP-like replication system.

The work described in this paper was funded and C. A. Smith is supported by MRC project grant G82/2421/3CB awarded to C. M. Thomas.

REFERENCES


Grant, A. J., Bird, P. I. & Pittard, J. (1980). Naturally occurring plasmids exhibiting incompati-
bility with members of incompatibility groups I and P. Journal of Bacteriology 144, 758–765.


