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A new member of the IS605 transposable element family, designated ISHp608, was found by subtractive hybridization in Helicobacter pylori. Like the three other insertion sequences (ISs) known in this gastric pathogen, it contains two open reading frames (orfA and orfB), each related to putative transposase genes of simpler (one-gene) elements in other prokaryotes; orfB is also related to the Salmonella virulence gene gapA. PCR and hybridization tests showed that ISHp608 is nonrandomly distributed geographically: it was found in 21% of 194 European and African strains, 14% of 175 Bengali strains, 43% of 131 strains from native Peruvians and Alaska natives, but just 1% of 223 East Asian strains. ISHp608 also seemed more abundant in Peruvian gastric cancer strains than gastritis strains (9 of 14 versus 15 of 45, respectively; P = 0.04). Two ISHp608 types differing by ~11% in DNA sequence were identified: one was widely distributed geographically, and the other was found only in Peruvian and Alaskan strains. Isolates of a given type differed by ≤2% in DNA sequence, but several recombinant elements were also found. ISHp608 marked with a resistance gene was found to (i) transpose in Escherichia coli; (ii) generate simple insertions during transposition, not cointegrates; (iii) insert downstream of the motif 5’-TTAC without duplicating target sequences; and (iv) require orfA but not orfB for its transposition. ISHp608 represents a widespread family of novel chimeric mobile DNA elements whose further analysis should provide new insights into transposition mechanisms and into microbial population genetic structure and genome evolution.

The hundreds of known insertion sequences (ISs) of prokaryotes are diverse in overall structure and in detailed mechanism, specificity, and regulation of transposition (8, 21, 36). Typically, an IS found in one bacterial strain will be absent from many other strains of that species yet be closely related to ISs in other microbes. This pattern bears witness to a rich heritage of interspecies DNA transfer, which has resulted in the spread of individual ISs to taxonomic groups remote from those in which they arose. Their abundance also reflects the action of transposase proteins, which mediate IS movement without need for extensive DNA sequence homology; to the ability of these elements to proliferate in host genomes by transposition itself; and to contributions that these elements or genes associated with them (e.g., in composite drug resistance transposons) sometimes make to bacterial fitness.

Many ISs and other transposable elements specify just one transposase protein that acts as a multimer on matched (inverted repeat) sequences at each element end. Others, exemplified by Tn7, specify two different proteins that form a more complex transposase. With Tn7, each transposase protein has a distinct role in transposition, and additional Tn7-encoded proteins help select insertion sites and affect the efficiency of transposition, in part through interactions with host proteins (9, 31). A third type of element, exemplified by IS605 of the gastric pathogen Helicobacter pylori, contains just two transposition-related genes, each of which has protein level homology to the single putative transposase genes of other simpler one-gene ISs and thus may be of different phylogenetic origin (18, 19). The termini of these latter two types of elements tend to have less inverted repeat character, which implies that each end may be acted on differently by proteins of the transposition complex.

Each of the two other ISs found to date in \textit{H. pylori} (IS606 and IS607) is related to IS605 by protein level homologies in orfB (some 25 to 35% amino acid sequence identity) (18, 19), as is the new ISHp608 element described here (Fig. 1A). This orfB (putative transposase gene) is also related to \textit{gipA}, a gene found in a \textit{Salmonella} prophage that contributes to virulence during murine infection (32).

Two branches of the IS605 family can be distinguished based on protein level homologies in orfA, one represented by IS607, whose transposition was found to be orfA dependent and orfB independent (19), and the other represented by IS605 and IS606 (18) and also the present ISHp608. The orfAs of IS605, IS606, and ISHp608 are protein level homologs of the putative transposase gene of IS200, an element that is abundant in natural isolates of \textit{Salmonella} and \textit{Escherichia coli}. Only a few cases of IS200 transposition have ever been detected, however, and the mechanisms of its movement are not understood (8, 21, 36).

Sequences of IS605, IS606, and IS607 were each found in...
only a subset of *H. pylori* strains, always in chimeric (two-gene) elements, as depicted in Fig. 1A. In particular, none of the several hundred *H. pylori* strains tested to date was found to contain only *orfA* (or only *orfB*) without the cognate *orfB* (or *orfA*), positioned as in Fig. 1A (18, 19). Formal models to explain such association include (i) *OrfA* and *OrfB* proteins serving together as the functional transposase, at least in *H. pylori*; (ii) transposition mediated by just one of these proteins but regulated (in terms of efficiency or specificity) by the other; (iii) each protein mediating transposition in a different set of bacterial species (implying relatively recent acquisition by *H. pylori*); or (iv) one gene needed for transposition and the other contributing to fitness.

As background to the present studies, *H. pylori* chronically infects more than half of all people worldwide and is implicated in peptic ulcer disease and gastric cancer. Infections often begin in infancy, resulting from preferential transmission within the family or perhaps the local community, and tend to last for decades once established; new adult infections are rare (10, 12, 27, 38). *H. pylori* is also one of the most genetically diverse of bacterial species: independent isolates typically differ from one another by some 3 to 5% in DNA sequence in essential genes and can also differ markedly in gene content and chromosomal gene arrangement (1, 3, 4). This diversity is enhanced by recombination. In general, little if any linkage is found between alleles at different polymorphic sites in collections of strains from the same geographic region, a pattern referred to as free recombination (33).

Superimposed on the extensive recombination between strains from the same region are indications that *H. pylori* gene pools differ geographically. This is based primarily on distributions of DNA sequence motifs in the virulence-associated *vacA* and *cagA* genes and of insertion and deletion motifs downstream of *cagA* (17, 20, 26, 37, 39). Strains from Europe, India, and East Asia were generally easily distinguished from one another using these tests, whereas those from a largely native population in Lima, Peru, seemed closely related to those of Spain. Much of the genetic diversity in local *H. pylori* populations and geographic differences in gene pools can be attributed to *H. pylori*’s patterns of transmission (preferentially to children and within families), the extraordinary chronicity of infection, and the relative isolation of the strain(s) carried by any given person, noted above. In consequence, strains diverge from one another by random genetic drift and by selection for adaptation to the new gastric environments each person may present (15). Despite occasional mixed infection, there is only limited direct competition between *H. pylori* strains of different lineages and no effective selection for just a few genotypes that might be best suited for most people worldwide.

Here we describe the discovery and characterization of *ISHp608*, found first in an *H. pylori* strain from Peru, features of its structure and transposition behavior, and its remarkably nonrandom distribution in human populations.

MATERIALS AND METHODS

**General methods.** Standard procedures were used for *H. pylori* growth in a microaerobic atmosphere on brain heart infusion agar (Difco) containing 10% horse blood (2, 18, 19), *E. coli* growth in L broth or on L agar, genomic DNA isolation, DNA electrophoresis, and transformation of competent *E. coli* cells (29). Antibiotics, when needed, were used at the following concentrations (in

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**FIG. 1. IS element maps.** (A) Structures of ISs of *H. pylori*. *ISHp608* is 1,833 bp long, making it slightly smaller than its three other known *H. pylori* relatives (size range, 1,888 to 2,028 bp [18, 19]). Boxes represent ORFs: those with the same shading pattern are related at the protein level (range of 28 to 36% amino acid sequence identities), *orfA* sizes are 156, 148, 139, and 218 codons and *orfB* sizes are 384, 427, 446, and 420 codons in *ISHp608*, *IS605*, *IS606*, and *IS607*, respectively. *orfA* and *orfB* overlap by 10 codons in *ISHp608*, which is reminiscent of the arrangement in *IS607* (nine-codon overlap). Distances from left and right ends to the 5′ end of *orfA* and 3′ end of *orfB* in a representative *ISHp608* are 171 bp and 75 bp, respectively (accession no. AF357224). GenBank accession nos. for *IS605*, *IS606*, and *IS607* are AC000108, U95957, and AF189015, respectively. (B) Positions of oligonucleotide primers, the subtractive clone that led to the discovery of *ISHp608* and mutations generated in *ISHp608*. The locations and orientations of PCR and sequencing primers are denoted by arrowheads; these primers are described in detail in Table 1. Primers 1 and 20 are specific to sequences in *jhp0924* (an ORF in strain J99 [4]) located to the left and right of *ISHp608*, respectively, in strain PeCan2A. * and ** identify primers specific for *ISHp608* types 2 and 3, respectively (defined in Results). Subtractive clone D7 is a 242-bp *Alul* fragment from strain PeCan18B that begins approximately 30 to 34 bp from the left end of *ISHp608* copy 2, *cam#2*, *cam#3*, *cam#4*, and *cam#5* represent mutational derivatives of *ISHp608* from strain PeCan2A that contain the cam (resistance gene) cassette inserted into (cam#4 and cam#5) or replacing (cam#2 and cam#3) particular *ISHp608* sequences. In *cam#4* (*A′ B″*), cam was inserted just downstream of *orfB* and 68 bp from the rightmost end of *ISHp608* (generated with primers 8 and 18); in *cam#2* (*A′ B″*), 154 bp of *orfA* sequence was replaced by the cam gene (generated with primers 2 and 12); in *cam#3* (*A″ B′*), 948 bp of *orfB* sequence was replaced by cam (generated with primers 4 and 16); in *cam#5* (*A″ B″*), cam was inserted just 22 bp from the *ISHp608* right end, within a 22-bp direct repeat (putative transposase binding site) (generated with primers 9 and 19), as detailed in Results.

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Universal primers for adapter PCR


Blast and Pfam (version 5.3) homology search programs (http://www.ncbi.nlm.nih.gov/)

C (Perkin Elmer 2400). DNA sequence editing and analysis were performed using the Qiagen prep spin miniprep kit (Qiagen, Chatsworth, Calif.).

PCR primers used in these experiments are listed in Table 1, and their positions are diagrammed in Fig. 1B. When needed, the primers were cloned into RV-cleaved pBluescript plasmid DNA containing an additional terminal T (22).

Bacterial strains and plasmids. The H. pylori strains used for subtractive library construction were PeCan18B, from a Peruvian gastric cancer patient (tester), and a pool of SJM180A, SJM184A, and SJM189A, from Peruvian gastritis patients (driver). Peruvian gastric cancer strains (designated with the prefix PeCan) were cultured from biopsy samples provided by Alejandro Basualdo and Juan Combe of the Servicio Universitario de Apoyo, Universidad Peruana Cayetano Heredia (UPCH) in Lima. These and other biopsy samples were from persons with gastroduodenal complaints, were part of a series taken to help in diagnosis of disease, and were obtained with written informed consent under protocols approved by the local (UPCH) human studies committee. Most other H. pylori strains used here were from the Berg laboratory collection and have been described in detail elsewhere (7, 11, 20, 25).

The following notes on the ethnics of patients and medical disease associations may be useful in evaluating the present data. The 45 Peruvian gastritis strains were from persons of primarily Amerindian ancestry living in the shanty town of San Juan de Miraflores in Lima, Peru (strains designated by the prefix PeSM) have been described in detail elsewhere (7, 11, 20, 25).

TABLE 1. Primers used in this study

| Type and primer no. | Primer name | Sequence | Location
|---|---|---|---
| 1 | jhp0924R | 5’-TGGCCGTAGTGGCAAAATTAC | Left of ISHp608 in PeCan2A
| 20 | jhp0924F | 5’-TTATTAAGCTTAAATGGCAATT | Right of ISHp608 in PeCan2A

| Leftward-facing ISHp608-specific | 2 | 608-3a | 5’-GATATGGTATTTTGCAAGATAGAC | 235 bp from left end of ISHp608
| 3 | 608-1 | 5’-ACTAAATCCATTCGGCTGTCA | 208 bp from left end of ISHp608
| 4 | 608-F1 | 5’-CCATACCACCTTAATGTTGTC | 685 bp from left end of ISHp608
| 5 | 608-F | 5’-ATCCCTCTAGTCAATCTGTCAT | 851 bp from left end of ISHp608
| 6 | 608a112-R" | 5’-TTTTGTCATCTTACCTTTA | 958 bp from left end of ISHp608
| 7 | 608a113-R" | 5’-TAGGAGCGTGTGAAGCGACGA | 1444 bp from left end of ISHp608
| 8 | 608neutral-Fnew | 5’-CAAGCAGCTACAGCTCATTAG | 1772 bp from left end of ISHp608
| 9 | 608neutral-Fl | 5’-CTAAAGCTAGGGGAGATTTAGC | 1815 bp from left end of ISHp608

| Rightward-facing ISHp608-specific | 10 | 608a112-R" | 5’-TTATATATTTTTAAAACCAAACCACCA | 1647 bp from right end of ISHp608
| 11 | 608-out | 5’-GGCTACTTCTTGCAAATACATATCG | 1624 bp from right end of ISHp608
| 12 | 608-R | 5’-CAAGCCTTGGAGTGATGAGTGTC | 1446 bp from right end of ISHp608
| 13 | 608-out | 5’-GTAGCCAACTACAGCTTCATCAC | 921 bp from right end of ISHp608
| 14 | 608a113-R" | 5’-CCGAGCCATTCCGAACACCA | 620 bp from right end of ISHp608
| 15 | 608-out | 5’-TAGCAGGATTTGGAAGATCCTT | 343 bp from right end of ISHp608
| 16 | 608-out | 5’-AAGAGAAGAAATACCTCACTGCA | 200 bp from right end of ISHp608
| 17 | 608-out | 5’-CTATCCACATTCCATGAGGTTC | 140 bp from right end of ISHp608
| 18 | 608-neutral-Rnew | 5’-TTGCTTGATCCAACTTTTCCTT | 67 bp from right end of ISHp608
| 19 | 608-neutral-R | 5’-TTGATGTGAGGGCTGTAC | 22 bp from right end of ISHp608

Universal primers for adapter PCR

C1 | 5’-GTACATATTGCTTCGTAAGCGG | Adapter-specific primer (TaKaRa)
C2 | 5’-TAATACGACTCACTATAGGGAA | Adapter-specific primer (TaKaRa)

a Where specified in base pairs, location refers to position of 5’ end of primer in sequence of ISHp608 from H. pylori strain PeCan2A (GenBank accession no. AF357223).

b jhp094 refers to an ORF of unknown function in the genome of H. pylori strain 99 (4). The ISHp608 element found in strain PeCan2A is inserted in this ORF.

c Left and right ends of ISHp608 are diagrammed in Fig. 1.

d ISHp608 type 2 specific.

e ISHp608 type 3 specific.

micrograms per milliliter): ampicillin (Amp), 100; chloramphenicol (Cam), 25; and streptomycin (Str), 150. Plasmid DNAs were isolated from E. coli cultures using the Qiagen prep spin miniprep kit (Qiagen, Chatsworth, Calif.).

High-molecular-weight genomic DNA was isolated from H. pylori and E. coli by a hexadecyltrimethylammonium method (5). Restriction digestion and ligation were carried out as recommended by the manufacturers (generally New England Biolabs, Beverly, Mass.). DNA fragments used for cloning and hybridization were purified from 1% agarose gels using the GeneClean II kit (Bio 101, Vista City, Calif.). A subtractive DNA library was made using the PCR-Select bacterial genome subtraction kit (Clontech) (2). Genomic DNA from a strain of a gastric cancer patient was used as the driver, and a pool of DNAs from three strains from gastritis-only patients was used as the tester. Southern blotting and hybridization were performed using Hybrid N+ membranes (Amersham) according to the manufacturer’s recommendations. DNA was labeled using the ECL kit (Amersham).

Specific PCR was carried out in 25-μl volumes, containing 5 to 10 ng of DNA, 1 U of Taq polymerase (Biosale, Madrid, Spain), 0.25 mM of each deoxynucleoside triphosphate (dNTP), in a standard buffer for 30 cycles with the following cycling parameters: denaturation for 30 s at 94°C; annealing for 30 s at a temperature appropriate for the primer (generally 55°C); and DNA synthesis for 1 min or as appropriate (1 min per kb) at 72°C. PCR primers used in these experiments are listed in Table 1, and their positions are diagrammed in Fig. 1B. When needed, PCR products were cloned into EcoRV-cleaved pBluescript plasmid DNA containing an additional terminal T (22).

DNA sequencing was carried out using a Big Dye Terminator DNA sequencing kit (Perkin-Elmer) and ABI automated sequencers. Direct sequencing on 4 min at 60°C (Perkin Elmer 2400). DNA sequence editing and analysis were performed with programs in the Genetics Computer Group (Madison, Wis.), package, programs and data in H. pylori genome sequence databases (4, 35), and Blast and Pfam (version 5.3) homology search programs (http://www.ncbi.nlm.nih.gov/blast/blast.cgi and http://pfam.wustl.edu/hmmsearch.shtml).

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as the donor in bacterial conjugation; and MC4100 which is F- and StrR and was used as the recipient in conjugation (6). The plasmid pLibrscript SK- (+/Str- Ag-Tc; SalI) (hereafter called vector A) and the chromophenol resistance cassette (cam), developed initially by D. E. Taylor as a selectable marker for H. pylori, was derived from plasmid pBS103 by restriction with Snal plus EcoRV (as in reference 18) and inserted into ISHp608 to generate mutations and allow selection for ISHp608 transposition. ISHp608 transposition donor plasmids were constructed in the pBS plasmid vector using polymerase to minimize the risk of mutation during PCR.

Plasmid construct 1 [pBS:ISHp608(AB-A')] contained an entire ISHp608 element from strain Perucan2A, a segment obtained by PCR with primers 1 and 20, which are specific to sequences flanking the site of ISHp608 insertion in this strain (Fig. 1B). The PCR product was then cloned into EcoRV-cleaved pBS. Plasmid 2 [pBS:ISHp608orF(A'cam(A' B'))] was constructed by linearizing plasmid 1 above by PCR with outward-facing primers 12 and 2 (Fig. 1B) and then ligating the product with cam, replacing 154 bp of orfA with this selectable marker. Plasmid 3 [pBS:ISHp608orF(ABcam(A' B'))] was generated by linearizing plasmid 1 by PCR with outward-facing primers 16 and 4 (Fig. 1B) and then ligating the product with cam, replacing 948 bp of orfB with this selectable marker. Plasmid 4 [pBS:ISHp608orFneutral(A' B')cam] was generated by linearizing plasmid 1 by PCR with outward-facing primers 18 and 19 (Fig. 1B) and ligating the product with cam, inserting this selectable marker into a noncoding sequence 68 bp from the rightmost ISHp608 end. Plasmid 5 [pBS:ISHp608neutral(1A' B')] was generated by linearizing plasmid 1 by PCR with outward-facing primers 9 and 19 (Fig. 1B) and then ligating the product with cam, inserting this selectable marker in a noncoding sequence 22 bp from the rightmost ISHp608 end.

ISHp608 transposition in E. coli. Strain DB1683 was transformed with plasmids containing marked ISHp608 elements (plasmids 2, 3, 4, and 5, above). Transposition of ISHp608 elements to A factor was then scored in a mating-out assay with selection for the ISHp608 CamR marker. In brief, the cells from single transformant colonies were grown overnight with aeration, diluted 100-fold into 4 ml of LB plus 0.5% glucose in a petri dish, and incubated for another 2 h at 37°C without shaking. They were then mixed with 4 ml of an exponentially growing culture of the MC4100 recipient strain at equal cell density in a total volume of 8 ml and incubated for another 3 to 4 h at 37°C without shaking. One milliliter of mating mix was concentrated by centrifugation, and the pellet was spread on L agar containing Str and Cam. This selected for transposition of the ISHp608cam element to pOX38 and then transfer of pOX38::ISHp608cam to MC4100 (6, 18, 19). The values reported here are expressed as CamR StrR exconjugants per initial donor cell in matings that were carried out under the same conditions in each trial. The F factor plasmid was not marked with a separate drug resistance determinant, and thus the absolute efficiency of F factor transfer was not determined. The selected CamR Str' exconjugants were scored for the presence or absence of the plasmid vector (AmpR or AmpS phenotype, respectively).

Definition of ISHp608 insertion sites. Sites of ISHp608cam insertion in different H. pylori strains were identified by direct chromosomal sequencing, using outward facing primers as detailed above and/or Adapter PCR (PCR in vitro single site amplification and cloning kit; TaKaRa; PanVera, Madison, Wis.). In Adapter PCR, total genomic DNA was digested with restriction enzyme Sau3A and ligated with the Sau3A adapter cassette supplied with the kit. The first nested PCR was carried out with cassette primer Cl and ISHp608 terminus-specific primers (primer 3 for the left end and primer 16 for the right end; Table 1 and Fig. 1B). A second nested PCR was then carried out using cassette primer C2 and second ISHp608 terminus-specific primers (2 for the left end and 17 for the right end; Fig. 1B). Gel-purified PCR fragments were sequenced directly. Sites of ISHp608 insertion in E. coli were identified by direct chromosomal DNA sequencing with outward-facing primers and genomic DNA from ISHp608-containing exconjugants which had been generated by transposition to the pOX38 F factor and transfer of pOX38::ISHp608 by conjugation into the MC4100 F- recipient strain.

Phylogenetic analysis. Sequence data for phylogenetic analyses were aligned and edited using Vector NTI v5.5.2.1 (http://www.informaxinc.com/). Maximum-likelihood analyses were carried out using the Phylip 3.57c package of J. Felsenstein (http://evolution.genetics.washington.edu/phylip.html). Tree plot analysis and plotting were carried out using Treeview (http://taxonomy.zoology.gla.ac.uk/rod/tview.html).

Nomenclature. The ISHp608 element described here was originally accessioned in GenBank as IS608. This element was renamed with the initials Hp to reflect the species in which the element was discovered, as recommended (21), upon finding that Eiichi Ohtsubo and colleagues had independently used IS608 to name an IS from an E. coli pathogen (accession no. AP002563). Their IS608 element is distantly related to IS50 (Tn5) and is not related to ISHp608 from H. pylori that we describe here.

Nucleotide sequence accession numbers. The sequence of ISHp608 copy 1 from strain PeCan18B corresponds to nucleotides (nts) 1 through 1833 of the 1,833-nt sequence in GenBank accession no. AF357224; that of copy 2 from strain PeCan18B (lacking about 30 nts from the left) corresponds to nts 1 through 1787 of 1,787 nts in accession no. AF411947. The other ISHp608 elements were accessioned in GenBank as follows: from strain PeCan2A, nts 41 through 1872 of the 1,897 nts in accession no. AF357223; from strain India49, nts 173 through 2004 of the 2,153 nts in accession no. AF411939; from Lithuanian strain Lith5, nts 5 through 1838 of the 1,838 nts in accession no. AF411940; from strain SpainB53, nts 7 through 1839 of the 1,839 nts in accession no. AF411941; from strain PeruSM92, nts 14 through 1836 of the 1,838 nts in accession no. AF411942; from strain AlaskaP76, nts 7 through 1832 of the 1,832 nts in accession no. AF411943; from strain AlaskaP20, nts 5 through 1827 of the 1,828 nts in accession no. AF411944; from strain India77, nts 168 through 2000 of the 2,088 nts in accession no. AF411945; and from strain PeCan4N, nts 94 through 1898 of the 1,903 nts in accession no. AF411946.

RESULTS

ISHp608 discovery. Subtractive hybridization was carried out to search for sequences in an H. pylori strain from a native Peruvian with gastric cancer (PeCan18B) that were absent from each of three strains from native Peruvians with gastritis only (SJM180A, SJM184A, and SJM189A, used as a pool). One end of one subtractive clone (D7, 242 bp long) exhibited protein level homology to amino termini of a family of putative transposases: identities of 13 of 16 amino acids to transposases from Yersinia enterocolitica and Enterococcus faecium (AJ238015 and AF125554) and 12 of 16 amino acids to transposases of IS200 of E. coli and IS1004 of Vibrio cholerae (2002282D and AE004380). However, no significant DNA sequence matches to known H. pylori ISs or other database entries were detected in BlastN homology searches. These data suggested that a previously unknown IS had been found, which we provisionally designated ISHp608.

ISHp608 sequence and diversity. Sequencing of ISHp608 was initiated by primer walking, using genomic DNA of PeCan18B and primers based on the sequence of subtractive clone D7. Five primers were used in succession (primers 3, 11, 12, 13, and 15; Fig. 1B). Each of the three internal primers gave clear, unambiguous sequence traces of at least 500 nts, whereas two others (outward-facing primers 3 and 15, near left and right ends) yielded mixed traces after about 100 and 320 nts, respectively. These results suggested that this strain contained two or more copies of ISHp608 that were identical for nearly 1.7 kb and that either (i) the ends of these ISHp608 elements were traversed where sequences became mixed or (ii) the copies of this element differed in sequence at one or both ends. This ISHp608 sequence contained open reading frames (ORFs) of 156 and 384 codons that were protein level homologs of orfA and orfB, respectively, in IS605 (Fig. 1A). However, the two ORFs in ISHp608 were in the same orientation and overlapped by 10 ccdons, whereas their homologs in IS605 and IS606 were divergent. Their parallel orientation in ISHp608 is reminiscent of that in IS607, an element with homology to ISHp608 only in orfB, not orfA.

ISHp608 copy numbers in PeCan18B and four other ISHp608-positive strains (identified in preliminary dot blot tests) were estimated in Southern blots of genomic DNAs that had been singly digested by HindIII and SspI and probed
Hp608

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FIG. 2. ISHp608 terminal sequences in several H. pylori strains. Left and right termini of ISHp608 determined by sequence comparison are shown in uppercase, and flanking DNA and empty sites in reference strains 26065 and 399 (HP and jhp gene designations, respectively [4, 35]) are shown in lowercase letters. The left end of ISHp608 copy 1 in strain PeCan18B and flanking sequence were determined by sequencing adapter PCR products. This left end was located in a gene with low homology to HP0456 of strain 26065 and was absent from strain J99. The sequence of the right end was obtained directly from genomic DNA, but flanking sequence was not determined because this strain contained two ISHp608 copies. The right end of ISHp608 copy 2 from strain PeCan18B and flanking sequence were determined by adapter PCR and sequencing. Attempts to obtain the leftmost ~30 bp of ISHp608 copy 2 by PCR with jhp0942-specific primers and by sequencing directly on genomic DNA were not successful. The left and right ends of ISHp608 in strain PeCan2A and flanking sequences were determined by adapter PCR and by direct chromosomal sequencing using a primer matching the inferred flanking sequence, respectively. The left end of ISHp608 in strain PeCan16A and the flanking sequence were determined by adapter PCR; this left end was located in an IS605 orfA sequence. The right end was identified by genomic DNA sequencing, but the flanking sequence was not determined because this strain contained two ISHp608 copies and because no product was obtained after PCR using primers specific for ISHp608 and for the anticipated flanking IS605 sequence. Both ends and flanking sequences of the single copy of ISHp608 in strain India49 were determined directly from genomic DNA. The element was found inserted into the gene corresponding to jhp0974 of reference strain J99 (4). Both ends and flanking sequences of the single copy of ISHp608 in strain India49 were determined directly from genomic DNA and were found inserted into an intergenic (noncoding) region (between genes jhp1113 and jhp1114 in J99 and HP1188 and HP1189 in 26065).

with an internal 460-bp PCR fragment from ISHp608 (made with primers 5 and 12; Fig. 1B). Two strains (PeCan4A and PeCan10B) yielded just one hybridizing band in each digest, initiating just one ISHp608 copy per strain, whereas PeCan18 and the two other strains (PeCan2A and PeCan16A) each yielded two hybridizing bands, indicating two ISHp608 copies per strain.

The sequences of ISHp608 insertion sites were defined more closely by adapter PCR or direct sequencing of genomic DNAs from several representative ISHp608-carrying strains and identification of corresponding “empty site” sequences (lacking ISHp608) in databases of reference strains 399 and 26065. Sequences of each ISHp608 end and adjacent DNAs were obtained from three strains (PeCan2A, India49, and India77), and additional useful, although less complete, sequences were obtained in three other cases (strains PeCan16A and copies 1 and 2 from PeCan18B) (detailed in Fig. 2 legend). These H. pylori sequence data indicated that ISHp608 inserts preferentially with its left end adjacent to 5’-TTAC in target sequences, that it has little or no specificity for sequences next to its right end, and that it does not duplicate target DNA. This specificity is reminiscent of that seen with IS605 and IS606, which insert preferentially downstream of 5’-TTAA and 5’-TTAT, respectively (18). The organization of sequences within the termini of ISHp608 will be presented in the context of sequence conservation and diversity among ISHp608 elements below.

One of two elements from PeCan2A was sequenced next, in part to prepare for functional tests (below). The entire element was PCR amplified with primers specific for flanking sequences (in jhp0924, primers 1 and 20; Fig. 1B) under high-fidelity conditions (Expand High Fidelity polymerase; Boehringer-Mannheim), cloned in a pBS plasmid vector, and then sequenced. This element was 1.83 kb long with only 92% identity (DNA level) to that from strain PeCan18B. However, it also contained two ORFs that were 97% (orfA) and 91% (orfB) identical in amino acid sequence to those of the first isolate (PeCan18B) and that also overlapped by 10 codons.

Eight additional elements were then characterized, primarily by genomic DNA sequencing without PCR or cloning, to better understand phylogenetic relationships among ISHp608 isolates. In overview, each element was closely matched to the others in size and ORF content, and 7 of 11 elements sequenced belonged to two major types (type I from strains PeCan2, India49, SpainB53, and Lith5, and type 2 from PeSJMM2, AlaskaP20, and AlaskaP76). These two types differed by about 11% in DNA sequence, in contrast to ~1 to 2% divergence among isolates of a given type, as diagrammed in Fig. 3A. At the protein level, identities and similarities between types 1 and 2 were 95 and 98%, respectively, in orfA, and 88 and 92%, respectively, in orfB. The other four elements (from strains PeCan4A, India77, and the two elements from PeCan18B) were inferred to be recombinant (Fig. 3B).

In particular, the element in strain PeCan4A consisted primarily of type 2 sequence but with replacement of an internal 600-bp segment with type 1 sequence. The two copies of ISHp608 from strain PeCan18B seemed to contain seven or eight interspersed patches of type 1 and type 2 sequences, suggesting a history of numerous homologous recombination events. The two copies seemed to be identical (as noted above), except for about 160 bp near their left ends. The element from strain India77 consisted primarily of type 1 sequence but also contained an internal 400-bp segment that was only 72 and 75% matched to corresponding sequences in types 1 and 2 elements, respectively. The ≥25% DNA divergence from other ISHp608 sequences suggested that this 400-bp patch was derived from a third type of element.

A well-conserved DDE(K) motif, reminiscent of those found to be important in active sites in several families of transposases (21), was found in orfB of each ISHp608 element sequenced (identified with an asterisk in Fig. 4), in the orfBs of the three other H. pylori ISs, and indeed in each of the 26 significant orfB homologs from other organisms that were found in the public database (Fig. 4). Most of these DDE motifs are unusual, however, in the relatively short distance between D and E (just 18 amino acids, rather than 33 to more than 100 in most other systems analyzed to date) (14, 21).

Sequences near the ends of ISs generally constitute sites of...
transposition protein action. These sites may contain or be included in simple inverted repeats of about 10 to 25 bp in some elements but be more complex in structure (e.g., direct and interrupted inverted repeats) in other elements, depending on the binding specificity of cognate transposition proteins (8, 21, 28). Type 1 and type 2 ISHp608 elements each contained (i) a well-conserved direct repeat of 22 bp in the left and right ends; (ii) an octanucleotide (5′AAGCCCTT) or penta- or hexanucleotide (5′AACGCC or 5′AACGCC) direct repeats in the left ends of type 1 and type 2 elements, respectively; and (iii) conserved decamer (5′-GCTTTAGCTA) or dodecamer (5′-TAGCTTTAGCTA) direct repeats in the right ends of type 1 and type 2 elements, respectively (boxes and horizontal lines in Fig. 5A). These various direct repeat motifs were embedded in a longer and much-interrupted (~37 of 54 bp) inverted repeat (Fig. 5B). Given the apparently strict specificity for insertion of the ISHp608 left end downstream of 5′-TTAC target sequences, it is noteworthy that this left end varied in length and sequence: 5′-CC(A)₅CT or 5′-C(A)₅CT in type 1 elements and 5′-C(A/T)CC(C/T) in type 2 elements, whereas the right end, which seemed to join to target sequences less specifically, was constant in length in the elements analyzed (Fig. 2 and 5A).

**ISHp608 geographic distribution.** The frequency of ISHp608 carriage in various *H. pylori* populations was estimated by dot blot hybridization with 80 strains and PCR with 723 strains. Hybridization was done with a probe containing parts of *orfA* and *orfB* (460 bp, generated with primers 5 and 12; Fig. 1B), and PCR was carried out with primers that generated a ~296-bp product from both type 1 and type 2 ISHp608s (primers 4 and 12; Fig. 1B). Hybridization and PCR results were complementary and entirely consistent; the same 22 strains were scored as positive for ISHp608 in both tests, and no strains were scored as positive in one test but not the other. In particular, hybridization test results with DNAs from 24 Japanese strains supported the conclusion from PCR (below) that ISHp608 is rare or absent in East Asian strains.

ISHp608 was found in some 15 to 20% of strains from Europeans, Africans, Indians, and Bangladeshis, at somewhat higher frequency (41%) in strains from native peoples of the Americas (Peru and Alaska), but in only 1% of East Asian (Japanese, Korean, and Chinese) strains (Table 2). ISHp608 also seemed to be more abundant in our relatively few strains from Peruvian patients with gastric cancer than from those with gastritis only (~64% versus ~33%, respectively; *P* = 0.04, Fischer’s exact probability). No other correlation between ISHp608 carriage and overt disease was observed among cases for which clinical data were available.

All strains found to carry ISHp608 were screened by PCR with primers specific for type 2 elements (primers 6 and 10; Fig. 1B). These tests indicated that 8 of 24 native Peruvian and 14 of 32 Alaska native ISHp608 elements were type 2 (or contained type 2 sequences if recombinant), whereas none of the 53 elements from European, Indian subcontinent, or African strains seemed to be (Table 2).

The existence of a probable third type of ISHp608 element with ≥25% DNA divergence from all others was suggested by the sequence of a 400-bp segment in the recombinant element from strain India77 (noted above). No other elements with this type 3 sequence were found by PCR using type 3-specific
other members of this family studied to date (IS\textsuperscript{pOX38}, not cointegrate formation, as is common with the two \textit{Hp608} transposition involved simple separation of the marked genes at average frequencies of 5\%/H11003; AF357223, IS\textit{Hp608} type 1; AF411944, IS\textit{U95957}, IS\textit{606} 607; AF189015, IS\textit{S74909}, F75401, C72305, and A82794. OrfB homolog accession nos. used here are AC000108, E64644, U95957, A33489, AAC97568, CAC03683, G75376, CAB41498, BAA15060, C64895, CAA60219, F70884, E70811, AAB12365, AAF05601, AF357223, JC4292, AAK40078, AAA26505, AAF98319, T36649, AAA83564, S74909, F75401, CT2305, and A82794.

FIG. 4. Multiple alignments of amino acid sequences in region of putative DDE(K) motifs in OrfB and homologs. Asterisks identify residues of the DDE(K) motif that, by extrapolation from other systems, could be critical features of an active site (namely, coordination of divalent cations and nucleophilic attack leading to DNA cleavage early in transposition [14, 21]). The consensus sequence (Cons) was developed by multiple alignment in ClustalW in the NTI vector program. Underlining identifies residues also conserved in the consensus based on 26 sequences of OrfB transposition.

**DISCUSSION**

Prokaryotic transposable elements can affect microbial evolution, serve as sensitive indicators of population genetic structure and evolutionary change, act as “selfish” DNAs that increase in copy number by transposition, and exhibit diversity in detailed mechanisms of transposition. The IS\textit{Hp608} element described here represents a widespread but as yet relatively unstudied family whose members contain two genes (orf\textit{A} and orf\textit{B}) that are each homologs of the single putative transposase genes of simpler (one-gene) ISs. We identified two main IS\textit{Hp608} sequence types and found that IS\textit{Hp608} was randomly distributed in \textit{H. pylori} populations, that it formed sim-
ple insertions with limited target site specificity, and that it required orfA but not orfB for transposition in *E. coli*.

**Population genetics and evolution.** The observed orfA-orfB arrangements in the four known ISs of *H. pylori* (Fig. 1A) and the need for orfA but not orfB for transposition in *E. coli* (19; this study) contributes to interest in the evolutionary history of this mobile DNA family. The two unrelated orfA (e.g., ISHp608 and IS607) and related orfAs in two orientations (e.g., in ISHp608 and IS605) suggest that these elements arose several times, perhaps by transposition of simpler elements or specialized or illegitimate recombination. Explanations for the conservation of a particular orfA-orfB association in each IS species include (i) both genes being important for transposition in *H. pylori*; (ii) a need for just one gene for transposition.

**FIG. 5.** Repeat sequences in the termini of ISHp608 elements. (A) Size and spacing of direct repeats of type 1 and type 2 element termini. Boxed sequences represent a well-conserved 22 (or 23)-bp direct repeat present in both ISHp608 ends. Horizontal lines identify shorter direct repeats: octanucleotide (*5’ AACGCCTT 3’*) and related penta- or hexanucleotides (*5’ AACGC 3’, 5’ AACGCC 3’*) in the left ends of type 1 and type 2 elements, respectively; and decamer (*5’-GCTTTAGCTA-3’) or dodecamer (*5’-TAGCTTTAGCTA-3’) in the right ends of type 1 and type 2 elements, respectively. *TAG* is the stop codon at the 3′ end of orfB. *T* identifies the site of insertion of the *cam* cassette in ISHp608 right end (insertions 4 and 5; see Fig. 1B). Lowercase letters represent flanking sequences. (B) Subterminal imperfect inverted repeats of ISHp608. Left and right ends of ISHp608 (same strand) are aligned to highlight inverted repeat sequence (capital letters).
alleles of the East Asia. These results are reminiscent of findings that East Asian and other strains are generally distinguishable based on alleles of the \textit{vacA} and \textit{cagA} virulence genes and sets of deletion and retention motifs downstream of \textit{cagA} (17, 20, 26, 37, 39). Collectively these findings indicated that the gene pools of East Asian and Western \textit{H. pylori} strains are distinct and that \textit{H. pylori} is not truly a "panmictic" species, as had been postulated (33). Our IS\textit{Hp608} data suggest that IS\textit{Hp608} was transferred into \textit{H. pylori} after the East Asian and Western \textit{H. pylori} populations were separated. IS\textit{Hp608} would then have spread by DNA transfer between strains in a population and by recombination and transposition. This spread would have been accelerated if IS\textit{Hp608} contributed to bacterial fitness. The paucity of IS\textit{Hp608} in East Asian \textit{H. pylori} strains might then reflect a relative lack of \textit{H. pylori} gene flow (due to the thousands of years in which there was rather little human migration) between East Asia and many other parts of the world.

Our sequence and PCR analyses indicated that one-third or more of the IS\textit{Hp608s} in Alaskan and Peruvian strains were distinct (type 2), differing by some 11% in DNA sequence from most other IS\textit{Hp608} elements (Table 2 and Fig. 3A). The abundance of type 2 elements in Alaskan strains is difficult to interpret because there is little other information at present on Alaska native \textit{H. pylori} genotypes. The abundance of type 2 elements in Peru was surprising, however, because the Peruvian strains analyzed here seemed to be closely related to Spanish strains in tests of several diagnostic markers (20). Nevertheless, no type 2 elements were found among nine IS\textit{Hp608} elements from Spain or among the 28 other IS\textit{Hp608s} from elsewhere in Europe (United Kingdom and Lithuania). Many explanations seem tenable, given our present limited understanding of \textit{H. pylori} genome evolution, and two will be considered here. In one explanation, the ancestors of modern native Americans might have brought \textit{H. pylori} carrying type 2 IS\textit{Hp608} with them from central and northern Asia thousands of years ago. If those putative original \textit{H. pylori} strains differed genetically from the Spanish-type strains that now predominate, it would imply that type 2 IS\textit{Hp608} had been transferred into the new Spanish-type strains before they had completely displaced the original (Asian-type) Peruvian strains. This explanation would also imply, however, that the original putative Asian-Peruvian \textit{H. pylori} strains had often carried type 2 IS\textit{Hp608} elements, even though IS\textit{Hp608} is very rare now in East Asia (the two East Asian exceptions, being from Hong Kong, might actually be Western in origin). Alternatively, \textit{H. pylori} might have been brought to the Americas by early European conquerors (20). In this case, type 2 IS\textit{Hp608} elements might have been uncommon in the original European \textit{H. pylori} populations but then became abundant in Peru by chance (founder effect) and/or due to special conditions accompanying the spread of \textit{H. pylori} in the New World. For example, rapid person-to-person transmission in the large, rapid, \textit{H. pylori}-free, highly susceptible native population might have created con-
ditions that also fostered interstrain gene transfer and/or transposition of type 2 ISHp608 elements.

The orfB-gipA (Salmonella virulence gene) homology (32) and contributions of the unrelated Tn5 transposase to E. coli fitness (16) serve as reminders that the frequency of an ISHp608 type would be further increased if it contributed to the efficiency of H. pylori transmission to new hosts in any way. Direct tests for effects of ISHp608 elements on H. pylori fitness in culture and in in vivo models and studies of strains from other populations may help resolve these issues.

Transposition mechanism. The transposition of ISHp608 in E. coli was orfA dependent but orfB independent, as was the case with IS607, a family member related to ISHp608 only in orfB (protein level) (19). Essentially all ISHp608 transposition products were simple insertions, not cointegrates (which contain both plasmid vector and ISHp608 DNAs), whereas most IS605 and IS607 transposition products were cointegrates (18, 19). Where tested (ISHp608 and IS607), these outcomes were not affected by orfB inactivation. Simple insertions and cointegrates often result from double- versus single-strand DNA breaks by transposase, resulting in conservative (break-join) and replicative transposition, respectively (6, 28, 30, 36). Studies with Tn7 and IS903 have shown that elements can be switched from conservative to replicative modes by subtle mutations affecting transposase activity or accessibility of cognate DNA sites (23, 34). Equivalent differences might account for the different spectra of transposition products generated with ISHp608 versus IS605 and IS607.

Inspection of the sequences flanking ISHp608 in H. pylori and after transposition in E. coli indicated that (i) ISHp608 inserts with its left (orfA) end immediately downstream of 5′-TTAC; (ii) its right end joins target DNAs nonspecifically; and (iii) it neither duplicates nor deletes sequences at its site of insertion. Similar patterns were evident with IS605 and IS606, except that they inserted next to 5′-TATTAAA and 5′-TATTAT, respectively (18). In contrast, IS607, whose OrfA is unrelated, inserted preferentially between adjacent G nucleotides in target DNA (19). These patterns suggest that OrfA proteins help select DNA targets, a property exhibited by many but not all transposase proteins (13).

The 22-bp direct repeats embedded within the imperfect inverted repeats near each ISHp608 end (Fig. 5) may constitute sites of transposase action. In accord with this view, transposition was blocked by placing a cam cassette within one repeat unit near the right end but not by placing it interior to this repeat (68 bp from the end).

One model to explain transposition in E. coli of orfB mutant derivatives of ISHp608 and related elements holds that their movement is truly independent of any OrfB-like protein. Alternatively, an OrfB-like protein might be needed but be satisfied in E. coli by the chromosomally encoded homolog (B1432; 25% identity and 42% similarity to ISHp608 OrfB, accession no. C64895). In accord with this view are (i) the strong orfA-orfB associations discussed above (18, 19; this study), (ii) the frequent annotation of OrfB homologs as transposases, and (iii) conserved DDE(K) motifs (Fig. 4), reminiscent of those in active sites of transposases of several families.

Different distances can separate D1, D2, and E residues in the linear transposase sequence. In the Tn5 transposase, whose D2-E distance is unusually large (138 amino acids), the intervening peptide contributes importantly to transposase-DNA binding (14). If the DDE(K) motifs of OrfB homologs also form part of transposase active sites, they would, in contrast, be among the most tightly spaced of any known to date (18 amino acids between D2 and E in orfB versus 33 to more than 100 amino acids in others [21]). Based on the Tn5 precedent (14), this short spacing might affect ISHp608 OrfB protein interaction with DNA substrates.

Continued population-level analyses of ISHp608 and related elements and the proteins they encode may help elucidate the origins and modes of evolution of prokaryotic transposable elements, forces that shape their distributions in bacterial populations, and factors important in the evolution of their H. pylori hosts as well. Similarly, detailed biochemical analyses, focused on the two families of OrfA proteins, each needed for transposition although lacking obvious DDE(K) motifs, and also on the universally present OrfB proteins with their tightly spaced DDE(K) motifs (if OrfB ever participates in IS movement), should provide new insights into the diversity of mechanisms of transposition and DNA rearrangement.

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