Identification of a pathogenicity island required for *Salmonella* survival in host cells

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Communicated by Robert K. Selander, Pennsylvania State University, University Park, PA, April 1, 1996 (received for review October 26, 1995)

ABSTRACT We have identified a region unique to the Salmonella typhimurium chromosome that is essential for virulence in mice. This region harbors at least three genes: two (spiA and spiB) encode products that are similar to proteins found in type III secretion systems, and a third (spiR) encodes a putative regulator. A strain with a mutation in spiA was unable to survive within macrophages but displayed wild-type levels of epithelial cell invasion. The culture supernatants of the spi mutants lacked a modified form of flagellin, which was present in the supernatant of the wild-type strain. This suggests that the Spi secretory apparatus exports a protease, or a protein that can alter the activity of a secreted protease. The "pathogenicity island" harboring the spi genes may encode the virulence determinants that set Salmonella apart from other enteric pathogens.

Although virulence genes map to numerous chromosomal locations (1), several large regions-termed "pathogenicity islands"-often define virulence characteristics in enteric bacteria. Pathogenicity islands are found at several positions in the chromosome, and the genes within a given island often determine the specific disease condition that results from infection. For example, uropathogenic (2) and enteropathogenic (3) strains of Escherichia coli harbor different pathogenicity islands at the same chromosomal location (the selC locus). Although the pathogenicity island specific to uropathogenic strains encodes a hemolysin (2), the island in enteropathogenic E. coli encodes a type III secretion system that exports proteins responsible for the attachment and effacing lesions of intestinal cells (4). Therefore, the specific virulence properties of two types of pathogenic E. coli are largely determined by the "cassette" present at the selC locus.

Through the systematic screening of a Salmonella typhimurium DNA library, Fitts (5) recovered numerous clones containing sequences that were apparently unique to the salmonellae. Three of these clones appeared to have been acquired by horizontal gene transfer since they had base compositions much lower than the overall G+C content of 52-54% of the Salmonella genome (6). The restricted phylogenetic distributions of these sequences suggested that they encode biochemical or cellular functions that set Salmonella apart from other enteric species (7). One of these three clones contained a gene cluster-designated spa-that enables Salmonella to invade epithelial cells (8). The spa gene cluster is part of a 40-kb pathogenicity island that encodes a variety of determinants that mediate the entry of Salmonella into nonphagocytic cells (9). These determinants include a type III secretion system and its substrates, which are homologous to an antigen export apparatus on the virulence plasmid of Shigella (10). Another of these clones specified a member of the LysR family of transcriptional regulators-SinR-and has no role in virulence (6).

In a survey of enteric species, only one of the clones— RF333—was strictly confined to Salmonella (6). Here we report the molecular genetic characterization of this clone and the virulence properties of strains harboring mutations in the corresponding region of the chromosome. These analyses define a new pathogenicity island (*spi*) within Salmonella encoding a regulator and a type III secretion system essential for virulence in mice and survival in macrophages. The *spi* island is distinct from the invasion region containing the *spa* locus, suggesting that multiple events of horizontal transfer have been responsible for the acquisition of pathogenic properties in Salmonella.

MATERIALS AND METHODS

Bacterial Strains, Plasmids, and Growth Conditions. All S. typhimurium strains used in this study were derived from the mouse-virulent wild-type strain 14028s. Strains with mutations in the RF333 chromosomal region were constructed from the pMS333 derivatives pEG7186 and pEG7200, which harbor kan insertions in spiA and spiR, respectively (6). The resulting strains-EG5793 and EG5799-contain the SmaI 1.3-kb kan fragment from plasmid pUC4-KIXX (Pharmacia) inserted at the PmeI and NaeI sites of RF333, respectively (Fig. 1). The kan gene is in the same transcriptional orientation as spiA in EG5793 and as spiR in EG5799, and the kan promoter is predicted to transcribe the genes located downstream of spiA and spiR. [Similar mutations within the spa region generated by inserting the same kan cassette were not polar on downstream genes (8)]. The structure of the spi locus in the mutant strains was verified by PCR with RF333-specific primers and by Southern hybridization analysis with both the 5.7-kb BamHI fragment of RF333 and kan-specific probes. Plasmid pMS333 is a pUC19 derivative with the BamHI fragment from RF333 inserted at the BamHI site. Ampicillin was used at 50 μ g/ml and kanamycin at 40 μ g/ml.

DNA Sequencing, Analysis, and Other Molecular Biological Manipulations. Sequencing was carried out on both strands of plasmid pMS333 by the dideoxy chain termination method using the Sequenase kit (United States Biochemical) with ³⁵S-labeled dATP. Additional primers were synthesized as the partial sequences were obtained. Sequence analyses were performed with both GeneWorks (IntelliGenetics) and the GCG package (University of Wisconsin). Restriction endonucleases and phage T4 ligase were purchased from Bethesda Research Laboratories, Boehringer Mannheim, or New England Biolabs, and used according to the supplier's specifications. Other protocols were taken from Maniatis *et al.* (11).

Virulence Assays and Preparation of Culture Supernatants. Macrophage survival assays were conducted with the macrophage-like cell line J774 as described (12). Adhesion to and invasion of human intestinal Henle-407 cells were investigated

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Abbreviation: ORF, open reading frame.

Data deposition: The sequence reported in this paper has been deposited in the GenBank database (accession no. U51927). *To whom reprint requests should be addressed.



FIG. 1. Genetic and physical maps of the *S. typhimurium* RF333 (*spi*) region. The arrows indicate the size and direction of transcription of the four largest open reading frames (ORF) contained within RF333. The positions of the *kan* gene insertions in the mutants EG5793 and EG5799 are indicated.

as described (13) except that dilutions of bacterial cells were performed in PBS instead of Luria–Bertani broth. Virulence assays were performed with 7–8-week-old female BALB/c mice inoculated intraperitoneally with 100 μ l of bacteria diluted in PBS. Viability was recorded for at least 30 days in 10 mice per mutant at each dose. Culture supernatants were prepared and analyzed as described (14).

RESULTS

Molecular Genetic Analysis of a Salmonella-Specific Chromosome Segment. We determined the nucleotide sequence of the 5.7-kb BamHI insert of clone RF333 and identified four ORFs, which were designated as spiA, spiB, spiC, and spiR (spi for Salmonella pathogenicity island). The spiA, spiB, and spiC genes are organized in an operon that is divergently transcribed from spiR (Fig. 1). The spiA gene encodes a protein of 497 amino acids, and the spiB ORF is >323 amino acids and extends beyond the BamHI site at the end of the insert. The spiA gene is preceded by the spiC gene, which encodes a protein of 127 or 133 amino acids, depending on the translational start site. (Presumably, the 127-amino acid ORF is translated because the sequence of its putative ribosome binding site—AGGAG—corresponds to the 3' end of the 16s rRNA.) The spiA initiation codon is 1 bp downstream of the stop codon for spiC, which suggests that these proteins are translationally coupled. The initiation codon of spiC is separated by 400 bp from that of the spiR gene, which encodes a protein of >822 amino acids.

SpiR Is Similar to Proteins of the Two-Component Family. Two-component systems generally consist of a sensor protein that, in response to environmental cues, modifies the ability of a regulatory protein to affect transcription of particular genes (15). Several such systems have been implicated in the regulation of virulence functions in the salmonellae (16). The deduced amino acid sequence of the SpiR protein exhibits homology to the subgroup of two-component systems that have both conserved domains-the histidine kinase domain of sensors and the receiver domain of response regulators-in a single molecule (Fig. 2A). The SpiR protein sequence is related to: (i) E. coli BarA, which is encoded by a gene that was isolated as a multicopy number suppressor of an envZ mutation (17); (ii) RscC, which controls capsule synthesis in E. coli (18); (iii) BvgS, a major regulator of virulence determinants in Bordetella (19); as well as two-component systems of plants and yeast (20).

SpiA and SpiB Are Similar to Proteins of Type III Secretion Systems. Type III secretion systems export proteins by a mechanism distinct from either the classical *sec*-dependent pathway or that responsible for secretion of hemolysin in *E. coli* (21). These systems mediate secretion of virulence proteins in both animal and plant pathogens: Homologs include the *Salmonella* Inv/Spa and the *Shigella* Mxi/Spa complexes, which are necessary for host cell invasion (22); the *Yersinia* Ysc/Lcr proteins, which secrete a tyrosine phosphatase and a cytotoxin (23); and proteins in the plant pathogens *Erwinia*, *Pseudomonas*, and *Xanthomonas*, which are involved in the hypersensitive response (24, 25).

As shown in Fig. 2B, SpiA displays the highest degree of sequence similarity to SepC of enteropathogenic E. coli (4), YscC of Yersinia (26), and InvG of Salmonella (27); and SpiB exhibited a low level of similarity to Yersinia YscD (26) and to an unreported ORF adjacent to eaeA in enteropathogenic E. coli (4). The gene order of spiA and spiB on the Salmonella chromosome is the same as that of yscC and yscD on the virulence plasmid of Yersinia (26). But spiC has no homologs in the sequence databases and apparently is a component specific to the spi locus.

Virulence Properties of spi Mutants. To determine the function of the genes present in RF333, we constructed mutations in the spiA and spiR ORFs (Fig. 1; see Materials and Methods). When these mutations were transferred back to the S. typhimurium chromosome, viable colonies were obtained, indicating that the inactivated genes are not essential for growth in complex laboratory media. These mutants behaved like the wild-type parent in their ability to grow at different temperatures and in defined media.

Given the prevalence of type III secretion systems among animal and plant pathogens (21, 22), we hypothesized that strains with mutations in *spi* genes might be unable to export certain virulence determinants and, consequently, would be attenuated for virulence. Indeed, neither *spiR* nor *spiA* mutants killed BALB/c mice when inoculated intraperitoneally at >10,000 times the median lethal dose of the wild-type parent. This finding is in contrast to the phenotypes of the *inv/spa* mutants of *S. typhimurium* that are attenuated only when the bacteria are administered orally (28).

To determine the particular stage of infection in which the RF333-encoded determinants are required, we tested the ability of the *spiA* mutant to invade cultured epithelial cells and to survive within macrophages *in vitro*. The *spiA* mutant displayed wild-type levels of epithelial cell invasion but was defective for intramacrophage survival (Fig. 3), a result that further distinguishes the role of the *spi* genes from that of the genes in the *inv/spa* locus. Taken together with the mice virulence data, these experiments establish that the Spi proteins are essential for stages of infection beyond the initial interaction with intestinal cells.

Identification of Proteins Exported by the Spi Secretion System. Because spi apparently encodes components of a type III secretion system, we investigated whether spi mutants were defective in the export of proteins to the growth media. Culture supernatants of wild-type, spiA, and spiR mutant strains were prepared and their protein profiles were compared. The supernatant of the wild-type strain had a 45-kDa protein that was absent from the supernatant of the mutant strains; instead, the mutant supernatant harbored a 47-kDa protein that was absent from the wild-type cultures (Fig. 4). We isolated both the 45- and 47-kDa proteins and determined the amino acid sequences of their first 15 residues. The two sequences were identical to one another and to the S. typhimurium flagellin, and it is likely that the 45-kDa protein is a processed form of the 47-kDa protein. The profile of secreted proteins for the *spiR* mutant was identical to that displayed by the spiA mutant strain, suggesting that SpiR controls transcription of the spiCAB operon.

DISCUSSION

The virulence phenotypes of several enteric pathogens have often been attributed to the presence of DNA segments that are absent from the genomes of nonpathogenic strains. These pathogenicity islands are largely responsible for the virulence properties of enteropathogenic and uropathogenic strains of *E. coli* (2, 3). In *Salmonella*, many of the invasion determinants

| A | | | | | | | 1997 (See 1997) | | | | |
|---|---|--|--|---|--|---|---|--|---|---|--|
| SpiR | 371 TE | NKV | A | ERTOALNE | AKKRAERANK | RESTHUTVIS | HELRTPMNGV | LGAIE. LLO | TTPLNIEOOG | LADTARNCTL | 442 |
| BarA | 247 LAAVH | FEMOHNTDOA | TSDLRETLEO | METONVELDI. | AKKRAOFAAR | TKSEFLANMS | HELRTPLNGV | TGETR LTT. | KTELTPTORD | HUNTTERSAN | 339 |
| ResC | 408 LOISE | VHSRVRNENV | ATCVLVDVSS | RVKMEESLOE | MACAAECASO | SKSMELATUS | HELRTPLYCT | TGNLD. LLO | TKELPKGVDR | LUTAMNNSSS | 500 |
| LemA | 229 LONAO | EELOLSTDOA | TEDVRONLET | TETONTELDL | ARKEALEASR | TKSEFLANMS | HETRTPLNGT | LGETH. LLO | KSELTPROFD | YLGTIEKSAD | 321 |
| Byas | 677 TPYCD | SLGELKGIIG | GWIDITERAE | LLRE | AKESADAANR | AKTTELATMS | HETRTPMNAT | TGMLELALLR | PTDOEPD, RO | STOVAYDSAR | 767 |
| Etrl | 307. LSHAA | TLEE | SMRARDL | LMEONVALDL | ARREAETATR | ARNDFLAVMN | HEMRTPMHAT | TAL. SSLLO | ETELTPEORL | MVETILKSSN | 390 |
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| SpiR | SLLAIINNLL | DFSRIESGHF | TLHMEETALL | PLLDOAMOTI | OGPAOSKKLS | LRTFVGOHVP | LYFHTDSIRL | ROILVNLLGN | AVKFTETGGI | RLTVK | 537 |
| BarA | NLLAIINDVL | DESKLEAGKL | ILESIPFPLR | STLDEVVTLL | AHSSHDKGLE | LTLNIKSDVP | DNVIGDPLRL | OOIITNLVGN | AIKFTENGNI | DILVEKRALS | 439 |
| RcsC | LLLKIISDIL | DFSKIESEOL | KIEPREFSPR | EVMNHITANY | LPLVVRKOLG | LYCFIEPDVP | VALNGDPMRL | OOVISNLLSN | AIKFTDTGCI | VLHVR | 595 |
| LemA | NLLSIINEIL | DESKIEAGKL | VLDNIPFNLR | DLLODTLTIL | APAAHAKOLE | LVSLVYRDTP | LALSGDPLRL | ROILTNLVSN | AIKFTREGTI | VARAMLEDET | 421 |
| BygS | SLLELIGDIL | DIAKIEAGKE | DLAPVRTALR | VLPEGAIRVF | DGLAROKGIE | LVLKTDIVGV | DDVLIDPLRM | KOVLSNLVGN | AIKFTTEGOV | .VLAVTARPD | 866 |
| Etrl | LLATLMNDVL | DLSRLEDGSL | OLELGTENLH | TLFREVLNLI | KPIAVVKKLP | TTLNLAPDLP | EFVVGDEKRL | MOIILNIVGN | AVKESKOGSI | SVTALVTKSD | 490 |
| | | | | | | | | | ana ana ana ana ana ana | | |
| SpiR | RHEEOLIFLV | SDSGKGIEIO | OOSOIFTAFY | OADTNS | O.GTGIGLTI | ASSLAKMMGG | NLTLKSVP | GVGTCVSLVL | PLOEYOPPOP | IK.GTLSAPF | 629 |
| BarA | NTKVOIEVOI | RDTGIGIPER | DOSRLFOAFR | QADASISRRH | G.GTGLGLVI | TOKLVNEMGG | DISFHSOPNR | GSTFWFHINL | DLNPNIIIEG | PSTQCLAGKR | 538 |
| RcsC | ADGDYLSIRV | RDTGVGIPAK | EVVRLFDPFF | OVGTGVORNF | Q.GTGLGLAI | CEKLISMMDG | DISVDSEP | GMGSOFTVRI | PLYGAQYPOK | KGVEGLSGKR | 692 |
| LemA | EEHAOLRISV | ODTGIGLSSO | DVRALFOAFS | OADNSLSROP | G.GTGLGLVI | SKRLIEOMGG | EIGVDSTPGE | GSEFWISLKL | PKAREDKEES | LNIP.LGGLR | 519 |
| BvqS | GDAAHVOFSV | SDTGCGISEA | DOROLFKPFS | OVGGSAEAGP | APGTGLGLSI | SRRLVELMGG | TLVMRSAP | GVGTTVSVDL | RLT | | 947 |
| Etrl | TRAV | KDSGAGINPQ | DIPKIFTKFA | OTOSLATRSS | G.GSGLGLAI | SKRFVNLMEG | NIWIESDG.L | GKGCTAIFDV | KLG | | 565 |
| | | | | | - Interference - the | 100 100 000 | | | | | |
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| SpiR | 689LQILL | VDDADINRDI | IGKMLVSLGQ | HVTIAASSNE | ALTLSQQQRF | DLVLIDIRMP | EIDGIEC | VRLWHDEPNN | LDPDCMFVAL | SASVATEDIH | 780 |
| BarA | 668. MTVMA | VDDNPANLKL | IGALLEDMVQ | HVELCDSGHQ | AVERAKOMPF | DLILMDIQMP | DMDGIRACEL | IHQLPHQ | QQTPVIAV | TAHAMAGQKE | 757 |
| RcsC | 809. MMILV | VDDHPINRRL | LADOLGSLGY | OCKTANDGVD | ALNVLSKNHI | DIVLSDVNMP | NMDGYR | LTORIRO | LGLTLPVIGV | TANALAEEKQ | 896 |
| LemA | 657PRVLC | VDDNPANLLL | VQTLLEDMGA | EVVAVEGGYA | AVNAVQQEAF | DLVLMDVQMP | GMDGRQATEA | IRAWEAERNQ | SSLPIVAL | TAHAMANEKR | 749 |
| BvgS | 973. LRVLV | VDDHKPNLML | LRQQLDYLGQ | RVIAADSGEA | ALALWREHAF | DVVITDCNMP | GISGYELARR | IRAAEAAPGY | GRTRCILFGF | TASAQMDEAQ | 1067 |
| Etrl | 593LKVLV | MDENGVSRMV | TKGLLVHLGC | EVTTVSSNEE | CLRV.VSHEH | KVVFMDVCMP | GVENYQIALR | IH EKFTKQ | RHQRPLLVAL | SGNTDKSTKE | 684 |
| | 2009 | * | | | | | | | | | |
| SpiR | RCKKNGIHHY | ITKPVTLATL | ARYISIAA// | 808 | | | | | | | |
| BarA | KLLGAGMSDY | LAKPIEEERL | HNLLLRYK// | 785 | | | | | | | |
| ResC | RCLESGMDSC | LSKPVTLDVI | KOSLTLYA// | 924 | | | | | | | |
| LemA | SLLOSGMDDY | LTKPISEROL | AOVVLKWT// | 777 | | | | | | | |
| BvqS | ACRAAGMDDC | LFKPIGVDAL | RORLNEAV// | 1095 | | | | | | | |
| Ltrl | KCMSFGLDGV | LLKPVSLDNI | RDVLSDLL// | 712 | | | | | | | |
| | | NAME AND ADDRESS OF AD | | | | | | | | | |
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| | | | | | | | | | | | |
| В | | | | | | | | | | | |
| B | MUUN | KRI.TI.TI.IFT | LNTAKSD | FLSWKGNDFT | LVAROMPLAF | VI.HI.I.SENVD | TATTISPI.TT | AT PSGKTPP | GPPUDTI.NNT. | AAOYDLLTWF | 9.0 |
| B SpiA | MVVN | KRLILILLFI | LNTAKSD | ELSWKGNDFT | LYARQMPLAE | VLHLLSENYD | TAITISPLIT | AT.FSGKIPP | GPPVDILNNL EKPVKVLEKL | AAQYDLLTWF | 90 |
| B SpiA SepC | MVVN MKKI MKTHILI | KRLILILLFI SFFIFTALFC | LNTAKSD CSAQAAPSSL | ELSWKGNDFT EKRLGKNEYF | LYARQMPLAE IITKSSPVRA | VLHLLSENYD ILNDFAANYS | TAITISPLIT IPVFISSSVN FPUTUSKMAA | AT.FSGKIPP DD.FSGEIKN | GPPVDILNNL EKPVKVLEKL | AAQYDLLTWF SKLYHLTWYY SLOLGLIWYF | 90 93 94 |
| B SpiA SepC InvG MyiD | MVVN MKKI MKTHILL MKKENIK | KRLILILLFI SFFIFTALFC ARVLACAALV SLTLLIVLLP | LNTAKSD CSAQAAPSSL LVTPGYSSEK | ELSWKGNDFT EKRLGKNEYF IPVTGSG | LYARQMPLAE IITKSSPVRA FVAKDDSLRT | VLHLLSENYD ILNDFAANYS FFDAMALQLK | TAITISPLIT IPVFISSSVN EPVIVSKMAA | AT.FSGKIPP DD.FSGEIKN RKKITGNFEF | GPPVDILNNL EKPVKVLEKL HDPNALLEKL SNDFFMLEKI | AAQYDLLTWF SKLYHLTWYY SLQLGLIWYF TLLVGLIWYK | 90 93 94 97 |
| B SepC InvG MxiD | MVVN MKKI MKTHILL MKKFNIK | KRLILILLFI SFFIFTALFC ARVLACAALV SLTLLIVILP PVLTCTLILI | LNTAKSD CSAQAAPSSL LVTPGYSSEK LIVNANNIDS SSY SWAO | ELSWKGNDFT EKRLGKNEYF IPVTGSG HLLEQNDIAK FLDWLDIDYV | LYARQMPLAE IITKSSPVRA FVAKDDSLRT YVAQSDTVGS | VLHLLSENYD ILNDFAANYS FFDAMALQLK FFERFSALLN LLTDECANYD | TAITISPLIT IPVFISSSVN EPVIVSKQAA YPIVVSKQAA | AT.FSGKIPP DD.FSGEIKN RKKITGNFEF KKRISGEFDL | GPPVDILNNL EKPVKVLEKL HDPNALLEKL SNPEEMLEKL DNPODELOHI | AAQYDLLTWF SKLYHLTWYY SLQLGLIWYF TLLVGLIWYK ASLYNIUWYY | 90 93 94 97 |
| B SepC InvG MxiD YscC | MVVN MKKI MKTHILL MKKFNIK MAFPLHSFFK | KRLILILLFI SFFIFTALFC ARVLACAALV SLTLLIVLLP RVLTGTLLLL | LNTAKSD CSAQAAPSSL LVTPGYSSEK LIVNANNIDS SSYSWAQ | ELSWKGNDFT EKRLGKNEYF IPVTGSG HLLEQNDIAK ELDWLPIPYV | LYARQMPLAE IITKSSPVRA FVAKDDSLRT YVAQSDTVGS YVAKGESLRD | VLHLLSENYD ILNDFAANYS FFDAMALQLK FFERFSALLN LLTDFGANYD | TAITISPLIT IPVFISSSVN EPVIVSKMAA YPIVVSKQAA ATVVVSDKIN | AT.FSGKIPP DD.FSGEIKN RKKITGNFEF KKRISGEFDL DK.VSGQFEH | GPPVDILNNL EKPVKVLEKL HDPNALLEKL SNPEEMLEKL DNPQDFLQHI | AAQYDLLTWF SKLYHLTWYY SLQLGLIWYF TLLVGLIWYK ASLYNLVWYY | 90 93 94 97 96 |
| B SpiA SepC InvG MxiD YscC SpiA | MVVN MKKI MKTHILL MKKFNIK MAFPLHSFFK DGSMLVVYPA | KRLILILFI SFFIFTALFC ARVLACAALV SLTLLIVLLP RVLTGTLLLL SLLKHOVITF | LNTAKSD CSAQAAPSSL LVTPGYSSEK LIVNANNIDS SSYSWAQ NILSTGRFIH | ELSWKGNDFT EKRLGKNEYF IPVTGSG HLLEQNDIAK ELDWLPIPYV YURSONILSS | LYARQMPLAE IITKSSPVRA FVAKDDSLRT YVAQSDTVGS YVAKGESLRD PGCEVKEITG | VLHLLSENYD ILNDFAANYS FFDAMALQLK FFERFSALLN LLTDFGANYD TKAVEVSGVP | TAITISPLIT IPVFISSSVN EPVIVSKMAA YPIVVSKQAA ATVVVSDKIN SCLTRISOLA | AT.FSGKIPP DD.FSGEIKN RKKITGNFEF KKRISGEFDL DK.VSGQFEH | GPPVDILNNL EKPVKVLEKL HDPNALLEKL SNPEEMLEKL DNPQDFLQHI KRKDSAVSVS | AAQYDLLTWF SKLYHLTWYY SLQLGLIWYF TLLVGLIWYK ASLYNLVWYY LYTLKYATAM | 90 93 94 97 96 |
| B SepC InvG MxiD YscC SpiA SepC | MVVN MKKI MKTHILL MKKFNIK MAFPLHSFPK DGSMLYVYPA DENILVIYKT | KRLILILFI SFFIFTALFC ARVLACAALV SLTLLIVLLP RVLTGTLLLL SLLKHQVITF NEISESITTP | LNTAKSD CSAQAAPSSL LVTPGYSSEK LIVNANNIDS SSYSWAQ NILSTGRFIH TYLDIDSLLK | ELSWKGNDFT EKRLGKNEYF IPVTGSG HLLEQNDIAK ELDWLPIPYV YLRSQNILSS YLSDTISVNK | LYARQMPLAE IITKSSPVRA FVAKDDSLRT YVAQSDTVGS YVAKGESLRD PGCEVKEITG NSCNVRKITT | VLHLLSENYD ILNDFAANYS FFDAMALQLK FFERFSALLN LLTDFGANYD TKAVEVSGVP FNS TEVRGVP | TAITISPLIT IPVFISSSVN EPVIVSKMAA YPIVVSKQAA ATVVVSDKIN SCLTRISQLA ECIKYITSIS | AT.FSGKIPP DD.FSGEIKN RKKITGNFEF KKRISGEFDL DK.VSGQFEH SVLDNALI FSLDKEAQ. | GPPVDILNNL EKPVKVLEKL HDPNALLEKL SNPEEMLEKL DNPQDFLQHI KRKDSAVSVS SKAKNKDVVK | AAQYDLLTWF SKLYHLTWYY SLQLGLIWYF TLLVGLIWYK ASLYNLVWYY IYTLKYATAM VFKLNYASAT | 90 93 94 97 96 188 191 |
| B SepC InvG MxiD YscC SpiA SepC nvG | MVVN MKKI MKTHILL MKKFNIK MAFPLHSFFK DGSMLYVYPA DENILYIYKT DGOAIVIYDA | KRLILILFI SFFIFTALFC ARVLACAALV SLTLIVILP RVLTGTLILL SLLKHQVITF NEISRSITTP SEMENAVVSI | LNTAKSD CSAQAAPSSL LVTPGYSSEK LIVNANNIDS SSYSWAQ NILSTGRFIH TYLDIDSLLK RNVSLNEPNN | ELSWKGNDFT EKRLGKNEYF IPVTGSG HLLEQNDIAK ELDWLPIPYV YLRSQNILSS YLSDTISVNK FLKE SCIN | LYARQMPLAE IITKSSPVRA FVAKDDSLRT YVAQSDTVGS YVAKGESLRD PGCEVKEITG NSCNVRKITT KNYPLRGDNR | VLHLLSENYD ILNDFAANYS FFDAMALQLK FFERFSALLN LLTDFGANYD TKAVEVSGVP FNSIEVRGVP KCTFVVSGPP | TAITISPLIT IPVFISSSVN EPVIVSKMAA YPIVVSKQAA ATVVVSDKIN SCLTRISQLA ECIKYITSLS VYVDMVVNAA | AT.FSGKIPP DD.FSGEIKN RKKITGNFEF KKRISGEFDL DK.VSGQFEH SVLDNALI ESLDKEAQ TMMDKOND | GPPVDILNNL EKPVKVLEKL HDPNALLEKL SNPEEMLEKL DNPQDFLQHI KRKDSAVSVS SKAKNKDVVK GIELGRKIG | AAQYDLLTWF SKLYHLTWYY SLQLGLIWYF TLLVGLIWYF ASLYNLVWYY IYTLKYATAM VFKLNYASAT VMRLNYTVG | 90 93 94 97 96 188 191 191 |
| B SpiA SepC InvG MxiD YscC SpiA SepC nvG xiD | MVVN MKKI MKTHILL MKKFNIK MAFPLHSFFK DGSMLYVYPA DENILYIYKT DGQAIYIYDA DCNALYIYDS | KRLILLILFI SFFIFTALFC ARVLACAALV SLTLLIVLLP RVLTGTLLLL SLLKHQVITF NEISRSIITP SEMRNAVVSL GELISKVILI | LNTAKSD CSAQAAPSSL LVTPGYSSEK LIVNANNIDS SSYSWAQ NILSTGRFIH TYLDIDSLLK RNVSLNEFNN ENTSLNVLIO | ELSWKGNDFT EKRLGKNEYF IPVTGSG HLLEQNDIAK ELDWLPIPYV YLRSQNILSS YLSDTISVNK FLKR.SGLYN | LYARQMPLAE IITKSSPVRA FVARDDSLRT YVAQSDTVGS YVAKGESLRD PGCEVKEITG NSCNVRKITT KNYPLRGDNR | VLHLLSENYD ILNDFAANYS FFDAMALQLK FFERFSALLN LLTDFGANYD TKAVEVSGVP FNSIEVRGVP KGTFYVSGPP DKTFY1SGPD | TAITISPLIT IPVFISSSVN EPVIVSKMAA YPIVVSKQAA ATVVVSDKIN SCLTRISQLA ECIKYITSLS VYVDMVVNAA | AT. PSGKIPP DD. PSGEIKN RKKITGNFEF KKRISGEFDL DK. VSGQFEH SVLDNALI ESLDKEAQ TMMDKQND TLIDKOVS | GPPVDILNNL EKPVKVLEKL HDPNALLEKL SNPEEMLEKL DNPQDFLQHI KRKDSAVSVS SKAKNKDVVK GIELGRQKIG SIGTDKVNFG | AAQYDLLTWF SKLYHLTWYY SLQLGLIWYF TLLVGLIWYK ASLYNLVWYY IYTLKYATAM VFKLNYASAT VMRLNNTFVG VIKIKNTFVS | 90 93 94 97 96 188 191 191 |
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| B SpiA SepC InvG MxiD YscC SpiA SepC nvG xiD scC | MVVN MKKI MKTHILL MKKTNIK MAFPLHSFFK DGSMLYVYPA DENILYIYKA DGNALYIYDS DGNULYIFKN | KRLILILDFI SFFIFTALFC ARVLACAALV SLTLLIVILP RVLTGTLLLL SLLKHQVITF NEISRSITF SEMRNAVVSL GELISKVILL SEVASRLIRL | LNTAKSD CSAQAAPSSL LVTPGYSSEK LIVNANNIDS SSYSWAQ NILSTGRFIH TYLDIDSLLK RNVSLNEPNN ENISLNYLIQ QESEAAELKQ | ELSWKGNDFT EKRLGKNEYF IPVTGSG HLLEQNDIAK ELDWLPIPYV YLSDTISVNK FLKR.SGLYN YLKD.ANLYD AL.QRSGIWE | LYÄRQMPLAE IITKSSPVRA FVAKDDSLRT YVÅQSDTVGS YVÅKGESLRD PGCEVKEITG NSCNVRKITT KNYPLRGDNR HRYPIRGNIS PRFGWRPDAS | VLHLLSENYD IILNDFAANYS FFDAMALQLK FFERFSALLN LLTDFGANYD TKAVEVSGVP FNSIEVRGVP KGTFYVSGPP DKTFYISGPP NRLVYVSGPP | TAITISPLIT IPVFISSSVN EPVIVSKAAA ATVVVSDKIN SCLTRISQLA ECIKYITSLS VYVDMVVNAA ALVELVANTA RYLELVEQTA | AT. FSGKIPP DD. FSGEIKN RKKITGNFEF KKRISGEFDL DK. VSGQFEH SVLDNALI ESLDKEAQ THMDRQND TLLDKQVS AALEQQTQIR | GPPVDILNNL EKPVKVLEKL HDPNALLEKL SNPEEMLEKL DNPQDFLQHI KRKDSAVSVS SKAKNKDVVK GIELGRQKIG SIGTDKVNFG SEKTGALAIE | AAQYDLLTWF SKLYHLTWYY SLQLGLIWYF TLLVGIIWYK ASLYNLVWYY IYTLKYATAM VFKLNYASAT VMRLNNTFVG VIKLKNTFVS IFPLKYASAS | 90 93 94 97 96 188 191 191 194 195 |
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FIG. 2. Comparison of *Salmonella* Spi with related proteins. The deduced amino acid sequences of proteins encoded by the *S. typhimurium spi* island are shown in the single-letter code. (A) Comparison of SpiR to members of the two-component family. (B) Comparison of SpiA to components of type III secretion systems. Alignment was conducted using the PILEUP program (GCG). Asterisks indicate residues conserved in all members of this family. For any given comparison, residues present in the majority of related proteins are highlighted.

have been localized to a 40-kb segment of the chromosome specifying at least 25 genes encoding a secretion system, its effectors, and the regulators controlling their expression (9). Because these genes are ancestral to all eight subgroups of *Salmonella enterica*, it has been hypothesized that the acquisition of this segment was an essential step in the evolution of invasion by *Salmonella* (14). However, the presence of structurally and functionally equivalent genes in the *Shigella flexneri* virulence plasmid (22) indicates that this invasion island does

not confer the specific properties that distinguish *Salmonella* from other enteric pathogens.

We have identified a new gene cluster essential for Salmonella virulence that includes the spiCAB operon, encoding components of a putative type III secretion apparatus, and a regulatory gene, spiR. Because the SpiA and SpiB proteins were similar to components of secretion apparatuses, we examined the profiles of proteins exported by spi and wild-type strains. The most conspicuous difference was the presence of



FIG. 3. Intramacrophage survival (A) and invasion (B) properties of an *spiA* mutant of S. *typhimurium*. Values represent the mean of triplicate samples \pm SD. Properties of *phoP* and *pho-24* strains are shown for comparative purposes.

two distinct forms of flagellin: 45 kDa in wild-type and 47 kDa in the *spi* mutants (Fig. 4). Thus, Spi may export a protease or a protein that modifies the activity of a secreted protease. However, flagellin is probably not the physiologically relevant substrate of this protease because flagellin is not essential for virulence and *spi* mutants are motile.

Although the Spi secretion system is reminiscent of that encoded within the *spa* invasion island of *Salmonella*, strains with mutations in the *spi* genes displayed wild-type levels of invasion (Fig. 3). Unlike *invA* mutants, which are only attenuated upon oral inoculation (28), the *spi* mutants failed to cause lethal infections in mice even when inoculated intraperitoneally. The *spi* genes were required for intramacrophage survival—a trait fundamental to *Salmonella* pathogenesis (29, 30)—and not surprisingly, homologous sequences have not been detected in other enteric species (6). Thus, Spi constitutes a novel type III secretion system that exports proteins that permit the survival of *Salmonella* within phagocytic cells, perhaps by modifying host factors required for phagosome– lysosome fusion (31) or phagosome acidification (32).

The spi gene cluster has an anomalous base composition of only 42.1% G+C, which is much lower than the overall G+C content of 52–54% of the Salmonella genome (6), and suggests that it was acquired by horizontal transfer. Regions acquired through horizontal processes often harbor the genes encoding both structural and regulatory elements in a contiguous DNA segment (9, 33); and within the 5.7-kb region analyzed in this study, we identified genes encoding products similar to components of a secretion apparatus (spiA and spiB) and a transcriptional regulator (spiR). SpiR exhibits broad-scale similarity to proteins of the two-component regulatory family and is likely to control the expression of adjacent loci in the spi pathogenicity island. Indeed, the spiR mutant displayed a



FIG. 4. Protein profile of supernatants prepared from wild-type, *spiA*, and *spiR* strains of *S. typhimurium*.

profile of exported proteins identical to that of the *spiA* strain (Fig. 4), implying that SpiR regulates the *spiCAB* operon and/or genes encoding the substrates of the Spi secretion system.

The spi locus is situated between 30.5' and 33.5', a region of the S. typhimurium chromosome that is relatively devoid of mapped genes (34). Only two other Salmonella-specific genes have been mapped to this region: sly and ompD. Despite the role of sly in virulence (35), it is not likely to be contained within the spi pathogenicity island due to its base composition—50.1% vs. 42.1% G+C for the spi locus—and its presence in Shigella and enteropathogenic E. coli. The spi genes do not correspond to any previously described macrophagesurvival loci (36). However, it is presently unknown whether spiA is allelic with a virulence gene that has been reported to be homologous to invG because neither the DNA sequence nor the map position of this gene was provided (37).

What is the mechanism by which horizontally acquired DNA segments are incorporated into the bacterial genome? In *E. coli*, a site adjacent to the *selC* tRNA gene is the target of insertion of two different pathogenicity islands (2, 3) and of a retronphage (38). Moreover, the second pathogenicity island present in certain uropathogenic strains of *E. coli* is located next to *leuX*, which, like *selC*, encodes a tRNA gene for rarely used codons (2). On the other hand, a 100-kb virulence region in the *Yersinia pestis* chromosome is flanked by repeated sequences (39). Thus, different mechanisms may mediate the incorporation of foreign DNA sequences. Future analysis of the entire *spi* island should reveal the manner in which the *spi* region was obtained by ancestral *Salmonella*.

The enteric pathogens Salmonella and Shigella employ similar secretion systems to promote entry into host cells (22). The differences in disease pathology and host range displayed by these pathogens most likely result from genes that are speciesspecific. The *spi* pathogenicity island may harbor such genes because: (*i*) it is present in all Salmonella serotypes investigated (5); (*ii*) *spi*-hybridizing sequences have not been detected in the genomes of other enteric species (6); and (*iii*) this region is essential for virulence functions beyond host cell invasion.

We thank Eric Mayer for technical assistance, Christine Smith (Searle/Monsanto) for protein sequencing, and anonymous reviewers for suggestions on a previous version of this manuscript. This research was supported by grants from the National Institutes of Health (GM-54900 and GM-48407). E.A.G. is a recipient of a Junior Faculty Research Award from the American Cancer Society. F.C.S. is a Career Investigator of the Consejo de Investigaciones de la Universidad Nacional de Rosario (CIUNR) (Rosario, Argentina) and is also supported by the Pew Charitable Trusts.

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