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J. Majewski, McGill University
Frederick M Cohan, Wesleyan University

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DNA Sequence Similarity Requirements for Interspecific Recombination in Bacillus

Jacek Majewski and Frederick M. Cohan
Department of Biology, Wesleyan University, Middletown, Connecticut 06459
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ABSTRACT

Gene transfer in bacteria is notoriously promiscuous. Genetic material is known to be transferred between groups as distantly related as the Gram positives and Gram negatives. However, the frequency of homologous recombination decreases sharply with the level of relatedness between the donor and recipient. Several studies show that this sexual isolation is an exponential function of DNA sequence divergence between recombining substrates. The two major factors implicated in producing the recombinational barrier are the mismatch repair system and the requirement for a short region of sequence identity to initiate strand exchange. Here we demonstrate that sexual isolation in Bacillus transformation results almost exclusively from the need for regions of identity at both the 5' and 3' ends of the donor DNA strand. We show that, by providing the essential identity, we can effectively eliminate sexual isolation between highly divergent sequences. We also present evidence that the potential of a donor sequence to act as a recombinogenic, invasive end is determined by the stability (melting point) of the donor-recipient complex. These results explain the exponential relationship between sexual isolation and sequence divergence observed in bacteria. They also suggest a model for rapid spread of novel adaptations, such as antibiotic resistance genes, among related species.

CURRENT models of gene transfer in bacteria assume that the RecA protein binds to a single-stranded 3' end of donor DNA and then searches the recipient genome for a homologous segment (Rao et al. 1995). This search is satisfied by an identical region of ~26 bp, the minimum efficiently processed segment (MEPS) believed to be the unit of homologous pairing (Shen and Huang 1986, 1989; Sieh et al. 1992; Rao et al. 1995). After identifying the homologous region, RecA is able to initiate strand invasion, which is followed by branch migration and extension of the donor-recipient heteroduplex molecule. The stringency of the homology search, along with the subsequent action of the mismatch repair system, is believed to cause sexual isolation between divergent bacterial species (Harris-Warrick and Lederberg 1978; Rayssiguier et al. 1989; Datta et al. 1997; Vulic et al. 1997; Majewski and Cohan 1998).

We previously demonstrated that, in Bacillus transformation, sexual isolation is an exponential function of DNA sequence divergence between the donor and the recipient (Roberts and Cohan 1993; Zawadzki et al. 1995). We subsequently showed that, in our system, mismatch repair is not a significant cause of sexual isolation and proposed a model where sexual isolation between divergent sequences is caused by difficulty in finding regions of identity necessary for strand invasion (Majewski and Cohan 1998). While the model explained the observed exponential relationship, it was based on molecular mechanisms identified in Escherichia coli, which may not necessarily extend to Bacillus. Moreover, most of the molecular studies of recombination are performed either in vitro or between artificial substrates (such as plasmid and phage). Little is known about actual in vivo mechanisms governing the major pathways of genetic exchange in bacteria (transformation, conjugation, and transduction).

In this work, we used an in vivo system of natural transformation in Bacillus subtilis to study the DNA sequence requirements for successful genetic exchange between divergent Bacillus species. We followed the approach of Roberts and Cohan (1993) involving transformation of B. subtilis toward resistance to rifampicin (rifR), using mutant rifR alleles of rpoB from related Bacillus species (Roberts and Cohan 1993; Zawadzki and Cohan 1995; Zawadzki et al. 1995; Majewski and Cohan 1998). We used chimeric constructs composed of donor DNA flanked by the recipient's own DNA to show that flanking identity is necessary for successful recombination. We also carried out a detailed analysis of recombinant junctions resulting from transformation with divergent DNA and determined the minimum sequence similarity for junctions of recombination.

MATERIALS AND METHODS

Strains: B. subtilis subsp. subtilis strain 1A96 was obtained from the Bacillus Genetic Stock Center (BGSC; Table 1). The B. subtilis mismatch repair deletion mutant, PB1856 (Ginetti
DNA). The sexual isolation values were then averaged over to allow formation of a chimeric template. The end primers subtracted from the colony counts obtained after transforma- tion. The divergence level between the recipient strains and the donor strains are based on restriction digest data analysis of the gene. The 3361-bp fragment of chimeric mutants, BG125 and BG126 (et al. 1999), and B. atrophaeus (NRRL NRS-213) was obtained from the Agricultural Research Service Culture Collection (ARSCC) at the National Center for Agricultural Utilization Research. Naturally occurring strains of B. subtilis subsp. subtilis, B. licheniformis, and B. atrophaeus were used as recipients in experiments investigating the role of the AddAB enzyme on the sexual isolation values were then averaged over to allow formation of a chimeric template. The end primers subtracted from the colony counts obtained after transformation.)

**Estimate of sequence divergence at rpoB:** The reported values of sequence divergence between the recipient strains and the donor strains are based on restriction digest data analysis (Robert et al. 1993). The divergence level between B. licheniformis and the recipient was confirmed by sequencing 3320 bp of the B. licheniformis gene.

**Transformation:** The recipient strains were induced to be competent and were transformed toward rifampicin resistance with 3 μg/ml DNA (Cohan et al. 1991; Roberts and Cohan 1993). This DNA concentration is above saturating, so as to prevent any effect of errors in concentration measurement on transformation frequencies. For each experimental replicate, sexual isolation (ρ) was calculated as the ratio of the homogamic transformation frequency (i.e., using rifR DNA derived from a mutant of the recipient strain) to the heterogamic transformation frequency (using a divergent donor’s rifR DNA). The sexual isolation values were then averaged over all experimental trials. Frequencies of transformation were calculated as the fraction of colony-forming units that were resistant to rifampicin, after accounting for spontaneous mutation toward rifampicin resistance. The spontaneous rate of resistance was determined for each experimental trial and subtracted from the colony counts obtained after transformation.

**PCR amplification of the rpoB gene:** The 3361-bp fragment was amplified using primers extending from 9 to 29 (5′-TCAACTAGTTCAGATGGACG-3′) and 3369 to 3349 (5′-ACCTGGTTCAGGACATGTC-3′) of the B. subtilis rpoB sequence (Boor et al. 1995). For the 5′ fragment of chimeric constructs the primers were 9-29 (as above) and 1532-1332 (5′-TACGTGACGCTATTTGTA-3′; Figure 1b), or 1464-1441 (5′-TCGGACAGCGCTGTATGCT-3′; Figure 1c). For the fragment encompassing the rifR region, the primers were 1332-1352 (5′-ATCAAATTACGCCCTACAGGCTGTAT-3′) or 1441-1475 (5′-ACGTAAAGCCTGTGCTTG-3′), and 2533-2512 (5′-TACCAA CAAGAACATCTCCGTC-3′). The primers for the 3′ end of the gene were 2512-2533 (5′-GACGAGATCTTTGTTGGTA-3′) and 3369-3349 (as above). All of the above PCRs were carried out using Vent DNA polymerase (New England Biolabs, Beverly, MA), at 52°C annealing temperature, under standard conditions, for 25 cycles. The fragments were purified by 0.8% agarose gel electrophoresis and extracted from gels using the Qiagen (Chatsworth, CA) gel purification kit. The DNA concentration was determined by comparison to a standardized concentration of λ phage BstEII marker.

**Overlap PCR and construction of chimeric genes:** Chimeric genes were constructed using a method adapted from Upender et al. (1995; Ge and Rudolph 1997). The following overlapping gene fragments were obtained in the primary PCRs (above): 5′ end of the recipient gene, rifR region of the donor gene, and the 3′ end of the recipient gene. These fragments were mixed in equimolar quantities, and 3 ng of the mixture was used as template in the secondary PCR. This reaction was carried out using Taq DNA polymerase (Promega, Madison, WI) for 2 cycles, at 52°C and 2.5 min. extension time to allow formation of a chimeric template. The end primers (9-29 and 3369-3349, see above) were then added to the mixture and the reaction was carried out for an additional 25 cycles, with 3.5 min extension time. The chimeric products were gel purified, adjusted to a standard concentration, and used for transformation. The accuracy of the constructs was confirmed by restriction mapping.

**Sequencing:** We sequenced the rpoB gene from the B. licheniformis donor strain and obtained 3320 bp of double-stranded DNA.
B. subtilis using the chimeric PCR constructs greatly reduced the invasion takes place, there might exist a difficulty in

...sexual isolation decreased from...  

...can eliminate the barrier completely. These results show that sexual isolation in Bacillus is predominantly caused by difficulty in strand invasion, with only a slight contribution from mismatch repair.

Analysis of fragments integrated during transformation: To confirm that the residual sexual isolation observed using chimeric PCR constructs results from the construct ends being removed prior to recombination, we analyzed the transformant rifR DNA. We extracted genomic DNA from the transformants obtained with the B. licheniformis construct and then PCR amplified the transformant rifR regions (1159 bp). We used restriction mapping to determine the length of foreign donor DNA that actually integrated into the chromosome. We chose B. licheniformis for this analysis, since its high degree of sequence divergence from the recipient allowed us to generate detailed restriction maps and to determine the size of the donor insert with high confidence. We analyzed 12 independent transformants of the wild-type recipient and ρ = 1.18 in the mismatch repair mutant (Table 4). Hence, by providing flanking regions of identity, we can almost entirely remove the recombinational barrier for a divergent DNA segment. By further removing the mismatch repair system, we can eliminate the barrier completely. These results show that sexual isolation in Bacillus is predominantly caused by difficulty in strand invasion, with only a slight contribution from mismatch repair.

Results

Transformation with chimeric DNA fragments: We used overlap-extension PCR to construct DNA fragments consisting of the rifR region from a divergent donor species, flanked by 5′ and 3′ ends of the B. subtilis recipient (Figure 1). Such chimeric genes provide the recombining rifR fragment with regions of perfect DNA sequence identity necessary for initiation of heteroduplex formation. We transformed two recipients (a wild-type B. subtilis and its mismatch-repair-deficient derivative) with chimeric constructs (Figure 1) and purely donor-derived PCR-amplified segments (Figure 1). The results are shown in Tables 2 and 3. Sexual isolation obtained with the constructs was drastically reduced relative to that of simple PCR products (Figure 2). For B. licheniformis (14.5% divergence at rifR) as donor, the sexual isolation decreased from ρ = 471 (simple PCR fragment) to ρ = 3.6 (chimeric construct) for the wild-type recipient and from ρ = 354 (simple PCR) to ρ = 2.4 (construct) for the mismatch repair mutant.

Retransformation with chimeric genomic DNA: While using the chimeric PCR constructs greatly reduced the recombinational barrier between B. subtilis and B. li-

Figure 1.—Substrates used for transformation. Shading indicates the portion of the gene derived from a divergent donor. Simple PCR (a) was used to amplify rpoB genes from B. subtilis subsp. subtilis strain 1A96 (recipient), B. subtilis subsp. subtilis RO-NN-1 (1% divergent at rifR), B. subtilis subsp. spizizeni RO-E-2 (3%), B. mojavensis RO-C-2 (4.5%), B. atrophaeus NRRL N5-213 (7%), B. amyloliquefaciens ATCC 23350 (8%), B. licheniformis ATCC 14580 (14.5%). Chimeric genes (b and c) were constructed using a method adapted from Upender et al. (1995). The strains used in the chimeric constructs (b) were from B. mojavensis, B. atrophaeus, and B. licheniformis. Only the B. licheniformis strain was used for the modified construct (c). Chromosomal DNA (d) was isolated from a 1A96 transformant containing the 1159-bp rifR segment of B. licheniformis DNA. The asterisk indicates the position of the rifR mutation, an H → Y substitution at amino-acid position 482 of B. subtilis RpoB.

sequence (bases 30–3349 of rpoB, GenBank accession no. AF172323). All sequencing reactions were carried out directly from PCR products, at the University of Pennsylvania DNA sequencing facility.

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to include the rif R mutation (Figure 1). We reasoned the results above, we isolated 43 clones resulting from 12 transformants of each mutant recipient and found donor fragment, as has been suggested by.

Analysis of recombination junctions and requirement for successful recombination: To determine whether invasive ends need to be present on only the 3' end of the donor strand (as in E.coli; Rao et al. 1995; Reddy et al. 1995) or on both ends, as suggested by the results above, we isolated 43 clones resulting from transforming the wild-type recipient with PCR-amplified segments of purely B. licheniformis-derived rpoB (Figure 1a). We used restriction mapping to determine the exact size and integration junctions of the 43 clones (Figure

Such a mechanism has been proposed as a barrier to transduction between Salmonella species (Zahrt and Maloy 1997): while branch migration stalls at mismatched regions, the RecBCD enzyme is believed to digest the free DNA strand, reducing the amount of genetic material available for integration. The B. subtilis addA and addB are homologues of E. coli recB and recC, and the AddAB enzyme is functionally active in E. coli strains deficient in RecBCD activity (Kooistra et al. 1993). We used the B. licheniformis construct to transform B. subtilis strains PB124 and PB125, with deletions in the addA and addB genes, respectively. We analyzed 12 transformants of each mutant recipient and found that the sizes of integrated foreign inserts were not significantly different than those in the wild-type recipient. Thus, AddA and AddB are not responsible for reducing the size of recombining substrates in Bacillus transformation.

To test whether some other factor might be responsible for clipping the free end of DNA during branch migration, we created a modified construct, where the 5' region of identity was extended further downstream to include the rif mutation (Figure 1). We reasoned that if only a small fragment directly adjacent to the identity region can be integrated, then extending the identity toward the 3' end should result in transformants extending further 3'-ward into the B. licheniformis insert.

However, on analyzing the transformants obtained with the new constructs, we found that the donor fragments were now smaller (<200 bp) and that most of them (22 out of 24) did not extend past base 1564 of rpoB. This was the same cutoff position observed with the previous construct, suggesting that there might be a recombination “hot spot” around position 1564. On examining the donor-recipient sequence alignment in this region, we found two highly conserved sequences (1534–1565 and 1495–1523; Figure 3) that may constitute such a recombination hot spot. These results suggest that regions of identity may be necessary at both ends of the donor fragment, as has been suggested by Harris-Warrick and Lederberg (1978). This requirement, rather than difficulty in heteroduplex extension past mismatched regions, may be responsible for the small size of integrated fragments.

### TABLE 2
Transformation frequencies and levels of sexual isolation using purely donor-derived PCR-amplified DNA

<table>
<thead>
<tr>
<th>Donor</th>
<th>Frequency of transformation (×10⁻³)</th>
<th>Sexual isolation</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Wild-type recipient (1A96)</td>
<td>ΔMutSL recipient (PB 1856)</td>
</tr>
<tr>
<td>B. subtilis subsp. subtilis A196</td>
<td>1.575 ± 0.428</td>
<td>2.893 ± 0.401</td>
</tr>
<tr>
<td>B. subtilis subsp. subtilis RONN1</td>
<td>1.743 ± 0.617</td>
<td>2.819 ± 0.491</td>
</tr>
<tr>
<td>B. subtilis subsp. spizizeni ROE2</td>
<td>0.804 ± 0.167</td>
<td>1.983 ± 0.225</td>
</tr>
<tr>
<td>B. mojavensis ROC2</td>
<td>0.767 ± 0.196</td>
<td>1.609 ± 0.329</td>
</tr>
<tr>
<td>B. atrophaeus NRRL NRS-213</td>
<td>0.153 ± 0.052</td>
<td>0.491 ± 0.157</td>
</tr>
<tr>
<td>B. amyloliquefaciens ATCC 23350</td>
<td>0.025 ± 0.006</td>
<td>0.173 ± 0.044</td>
</tr>
<tr>
<td>B. licheniformis ATCC 14580</td>
<td>0.0032 ± 0.0008</td>
<td>0.0069 ± 0.0017</td>
</tr>
</tbody>
</table>

* Mean and standard error values for four experimental trials.

### TABLE 3
Transformation frequencies and levels of sexual isolation using chimeric PCR-amplified DNA

<table>
<thead>
<tr>
<th>Donor (source of chimera’s insert)</th>
<th>Frequency of transformation (×10⁻³)</th>
<th>Sexual isolation</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Wild-type recipient (1A96)</td>
<td>ΔMutSL recipient (PB 1856)</td>
</tr>
<tr>
<td>B. subtilis subsp. subtilis A196</td>
<td>1.869 ± 0.214</td>
<td>3.663 ± 0.757</td>
</tr>
<tr>
<td>B. mojavensis ROC2</td>
<td>1.404 ± 0.189</td>
<td>3.249 ± 0.796</td>
</tr>
<tr>
<td>B. atrophaeus NRRL NRS-213</td>
<td>0.834 ± 0.069</td>
<td>3.010 ± 0.723</td>
</tr>
<tr>
<td>B. licheniformis ATCC 14580</td>
<td>0.514 ± 0.031</td>
<td>1.551 ± 0.353</td>
</tr>
</tbody>
</table>

* Mean and standard error values for four experimental trials.
We found that all the mismatched recombinogenic regions were characterized by a relatively high GC content (i.e., GC > 52%, in a gene with 45% GC). Hence, we hypothesized that there might exist a critical stability of the bond (which may be quantified by its melting temperature) between the invasive end and the recipient DNA, necessary to initiate recombination. To determine the critical melting point ($T_c$), we scanned the donor-recipient sequence alignment, identifying all oligonucleotides between 18 and 27 bp long (which is believed to be the MEPS size in E. coli; Hsieh et al. 1992) and calculated their melting temperatures. We tested several critical melting temperatures for the degree of association between the theoretical MEPS criterion and the empirically observed transformant junctions. The temperature $T_c = 51.2^\circ$ explained the greatest number of observed junctions, while minimizing the number of unused putative MEPS sequences (Figure 5). This melting temperature criterion, which takes into consideration the number of mismatches, the GC content, and the length of the oligonucleotide, gave a better explanation of the observed data than any criterion.

Finally, we determined the sequence similarity requirements for invasive ends. Many of the recombination junctions fell within regions of perfect sequence identity and could be easily explained. However, 44 junctions ended in regions interrupted by one to several mismatches. Moreover, some of the most recombinogenic regions were virtually identical in degree of mismatch to nearby regions producing no junctions (Figure 3). We found that 84 out of the 86 junctions analyzed ended in relatively mismatch-free stretches of 20–30 bp containing at most one or two mismatches (Figure 5). These results demonstrate that invasive ends are necessary on both ends of the donor strand. This differs from the findings in E. coli, where it is believed that a single-stranded invasive 3' end is sufficient for successful recombination between divergent sequences.

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**DISCUSSION**

**Mechanisms and barriers to recombination:** The molecular mechanisms of recombination in Bacillus have not yet been fully elucidated, yet it is clear that they differ from those observed in E. coli. In this article, we show that recombination in Bacillus requires short regions of conserved sequences to be present at both ends of the donor-recipient DNA. This is in contrast to the case of E. coli, where a conserved sequence at only one end is sufficient (Shen and Huang 1986). We show that the mismatch repair system is relatively ineffective in preventing recombination between divergent sequences and that the main barrier to recombination is caused by the difficulty in locating conserved regions necessary for strand invasion. Thus, highly divergent donor segments (up to 15% in this study) recombine with no resistance, if they are flanked with sequences identical to the recipient. This result implies that recom-

### TABLE 4

Transformation frequencies and levels of sexual isolation using genomic transformant DNA

<table>
<thead>
<tr>
<th>Donor of genomic DNA</th>
<th>Frequency of transformation ($\times10^{-3}$)</th>
<th>Sexual isolation</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Wild-type recipient (1A96)</td>
<td>$\Delta$MutSL recipient (PB 1856)</td>
</tr>
<tr>
<td>1A96 spontaneous rif$^+$ mutant</td>
<td>$10.3 \pm 1.8^a$</td>
<td>$10.0 \pm 1.1^a$</td>
</tr>
<tr>
<td>1A96 containing a 1159-bp B. licheniformis insert</td>
<td>$5.6 \pm 0.8$</td>
<td>$8.4 \pm 1.0$</td>
</tr>
</tbody>
</table>

$^a$ Mean and standard error values for four experimental trials.
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exponentially with the DNA sequence divergence between the donor and the recipient. In view of the new results, we now require that such conserved sequences be present on both ends of the donor strand. The modified model still leads to an exponential relationship between sexual isolation and sequence divergence. However, we now obtain a different estimate for the length \( n \) of the MEPS,

\[
n = 2\phi_h \times \ln(10),
\]

where \( \phi_h \) is the empirically determined sensitivity of sexual isolation to sequence divergence, in the absence of mismatch repair. (This is the same as Equation 5 of Majewski and Cohan (1998), except for the additional factor of 2 to account for the need of identity at both ends]. Using \( \phi_h = 17.85 \) (Majewski and Cohan 1998), the model predicts that the length of the MEPS should be 21 bp. This prediction is in excellent agreement with the MEPS sequences observed in our analysis of recombination junctions, where we show that regions of identity as short as 20 bp may promote recombination between divergent species.

The requirement for flanking identity can also explain why there is a more significant deviation from the exponential relationship when PCR-amplified DNA fragments are used in transformation, as compared to genomic DNA, which follows the relationship very closely (Robert's and Cohan 1993; Majewski and Cohan 1998). This deviation has been observed by Zawadzki et al. (1995) and in this article (Figure 2). We expect that transformation with small fragments, where most of the strand invasion events take place in the immediate vicinity of the mutation conferring antibiotic resistance, will be very sensitive to the position of the mutation; if the mutation occurs within a highly conserved region, transformation rates will be en-

![Figure 3](image-url)

**Figure 3.** Examples of sequences with low numbers of mismatches permitting (segments 2-6 and 8) and not permitting (segments 1 and 7) recombination junctions. Segments as short as 20 bp with one mismatch may be used as invasive ends, provided they have a high GC content (segment 6). In contrast, larger segments with low GC content (segments 1 and 7) produced no transformants.

![Figure 4](image-url)

**Figure 4.** (A) \( \text{rpoB} \) sequence alignment of \( B. \ subtilis \) and \( B. \ licheniformis \) for the first 2810 bp (the region where transformation was detected). Dark lines represent mismatches. (B) Extent to which melting temperatures exceed the critical temperature of \( T_c = 51.2^\circ \text{C} \) (see Figure 5 for explanation) for each oligonucleotide (18-27 bp) that may serve as an invasive end. The melting temperature was calculated using the formula \( T_m = 81 - 600/\text{length} + 16.6 \log(Na) + 0.41(\%GC) - 0.8(\% \text{ match}) \) and evaluated at 50 mm Na concentration (Sambrook et al. 1989; Malkov et al. 1997). (C) Inserts resulting from transformation of wild-type recipient with pure \( B. \ licheniformis \) PCR-amplified fragment. The end lines represent confidence intervals for transformant junctions. The asterisk represents the rif \( \text{R} \) mutation (nucleotide 1444). All junctions (except those labeled 1, 2, and 3) are associated with high melting point regions. The exceptions, 1 and 2, fall within an AT-rich region containing sequences similar to those implicated in illegitimate recombination in \( B. \ subtilis \) (Meima et al. 1998). Junction 3 falls in a region with a \( T_m \) of 49° (just below critical).
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Figure 5.—Estimation of the critical melting temperature of a MEPS. The vertical axis value is calculated as the mean of the following: (1) the percentage of junctions that are associated with a given critical temperature and (2) the percentage of putative MEPS with the given critical melting temperature that are associated with transformant junctions. Note that using the melting point (and thus including GC content) in MEPS determination gives a better explanation of experimental data than using only length and the number of mismatches. No preference was found for GT-rich regions, believed to be favored in recombination in E. coli and Saccharomyces cerevisiae.

enhanced, whereas mutations in less conserved regions will exhibit reduced frequencies of integration. On the other hand, during transformation with genomic DNA, strand invasion takes place within several kilobases of the mutation (Zawadzki and Cohan 1995), and the exact position of the marker is not as relevant. Hence, for transformation with spontaneous mutant markers (as is the case in our experiments), the exponential relationship will be followed more closely by large DNA fragments.

Our results also suggest that the strength of the bonds formed between the endpoints of the donor-recipient complex may determine whether or not recombination will take place. In our system, sequences with a relatively high GC content are more likely to produce recombinants than similar sequences (with respect to length of identity and number of mismatches) with a lower GC content. Although it has been suggested that the bonds formed between the invading donor strand and the recipient DNA duplex are not Watson-Crick in nature (Reddy et al. 1995), it is possible that their strength might still be dependent on GC pairing. We did not find any preference for recombination in GT-rich regions, which have been shown to preferentially bind both the E. coli RecA and the yeast Rad51 homologue (Tracy et al. 1997).

We are currently unable to determine why two conserved regions are necessary for successful recombination in Bacillus. In vitro studies using E. coli proteins show that RecA is able to initiate strand invasion at both 3' and 5' free ssDNA ends, but that branch migration can take place only in the 3' → 5' direction. It is possible that in Bacillus the 5' bond serves either to stabilize the entire complex during branch migration, or is necessary for termination of the process. It is also possible that the polarity of branch migration is different in Bacillus than in E. coli, since the yeast RecA homologue, Rad51, promotes branch migration in the 5' → 3' direction.

Interspecies recombination and adaptation: Consider next the possible adaptive value of the barriers to interspecies recombination observed in Bacillus and other taxa. We have previously shown that recombination across species poses little fitness cost in bacteria (Cohan 1994). Because recombination in bacteria is extremely rare, the cost of incorporating maladaptive foreign alleles should be very low, even if between-species recombination were to occur at the same rate as recombination within species. Given that recombination occurs within Bacillus species, for example, at a rate of 10^-7, the incremental fitness increase of further decreasing the between-species recombination rate is likely very small. Nevertheless, the selection disfavoring between-species recombination is not negligible, owing to the large population sizes for bacteria.

On the other hand, it is occasionally beneficial for bacteria to incorporate DNA from other species. The promiscuous nature of bacterial recombination allows a recipient to acquire adaptations from other species. Bacterial genomes are modular enough to accommodate the expression of some foreign alleles, and the size of recombining fragments is frequently small enough so that a generally adaptive allele can pass between species without the cotransfer of maladaptive alleles at other loci (Zawadzki and Cohan 1995). The average benefit of being able to recombine between species at an increased rate is likely to be quite low, however, since only a small fraction of adaptations would be expected to be functional in different species.

In summary, there are low, but nonzero, selection pressures acting to both increase and decrease the rate of between-species recombination. We expect that the optimum level of interspecies recombination will depend on the relative importance of obtaining other species’ adaptations vs. avoiding uptake of maladaptive foreign alleles.

Can the balance between these selection pressures
explain differences between bacterial species in their resistance to between-species recombination? This study has shown that in B. subtilis transformation, resistance to recombination occurs primarily at the strand invasion stage, with little effect of mismatch repair. That recombination appears to be blocked only by the thermodynamic requirements of donor-recipient stability suggests that the recombinational barrier observed may be a minimum that cannot be further reduced. In contrast, conjugation-mediated recombination between E. coli and Salmonella typhimurium is reduced 100 times more than is the case for transformation between similarly divergent Bacillus species (Vulic et al. 1997). This recombinational barrier is even more pronounced in transduction between S. typhimurium and S. typhi (Zahrt and Maloy 1997), where a 1% DNA sequence divergence can lead to an almost complete inhibition of recombination. This is due to a greater inhibitory effect of mismatch repair on recombination between these enteric species.

The difference between Bacillus and the enteric bacteria in recombinational barriers may result from a difference in the balance between the positive and negative selection pressures acting on interspecies recombination rates. In Bacillus transformation, competence is induced primarily under starvation (Dubnau 1993), a time when the relative importance of taking up new metabolic capabilities might be enhanced. We argue that transformation under unfavorable growth conditions would push the selective balance in favor of increased between-species recombination. Conjugation between Escherichia and Salmonella is not associated with a particular condition of growth, so the optimal level of between-species recombination may be quite low, requiring the inhibitory effect of mismatch repair.

It will be interesting to investigate whether genetic barriers such as mismatch repair may become more effective in Bacillus under nonstressful conditions. Specifically, are the additional barriers actively suppressed during transformation, resulting in the relatively low sexual isolation, or do they simply not exist under any conditions? Experiments such as interspecific transduction between related Bacillus species should be useful in resolving the above issue.

Whether genetic barriers in Bacillus are facultatively suppressed or constitutively weak, natural transformation presents an efficient system for acquisition of new genes and functions. Our findings have important implications in the spread of new adaptations, such as antibiotic resistance or artificially introduced foreign genes, in the bacterial world. We suggest that once a gene for a novel adaptation transfers into a species from a divergent donor, the gene becomes flanked by recipient DNA, and, in the absence of barriers other than resistance to strand invasion, this allows efficient homologous recombination of the adaptation into related species (see also Miller and Levy 1989). Our results suggest that even very divergent and heterologous genes may transfer between species at very high rates via homologous recombination.

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LITERATURE CITED
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