

Biochemistry and Comparative Genomics of SxxK Superfamily Acyltransferases Offer a Clue to the Mycobacterial Paradox: Presence of Penicillin-Susceptible Target Proteins versus Lack of Efficiency of Penicillin as Therapeutic Agent

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INTRODUCTION

The bacterial wall peptidoglycans are covalently closed, net-like polymers (77, 203). The glycan chains are made of alternating β -1,4-linked *N*-acetylglucosamine and *N*-acetylmuramic acid residues. The D-lactyl groups of the muramic acid residues are amidated with L-alanyl- γ -D-glutamyl-(L)-diaminoacyl-D-alanine stem tetrapeptides and L-alanyl- γ -D-glutamyl-L-diaminoacid stem tripeptides. The stem peptides can be branched, in which case the ω amino group of the diaminoacid residue is substituted by an additional amino acid residue or a short peptide. Unbranched or branched stem peptides belonging to adjacent glycan strands are covalently linked, resulting in polymeric peptidoglycan.

All peptidoglycan-containing bacteria in the exponential phase of growth manufacture a (4 \rightarrow 3) peptidoglycan (Fig. 1) in

a penicillin-susceptible manner. Peptidoglycan crosslinking extends from the carboxy-terminal D-alanine residue at position 4 of a stem tetrapeptide to the lateral amino group at position 3 of another, unbranched or branched, stem peptide. The (4 \rightarrow 3) interpeptide linkages or cross-bridges are made by specialized acyltransferases which are immobilized by penicillin in the form of stable, enzymatically inactive penicillin-binding proteins (PBPs) (214). *Escherichia coli*, the mycobacteria, and leprosy-derived corynebacteria (111) produce unbranched (4 \rightarrow 3) peptidoglycans with meso-diaminopimelic acid at position 3 of the stem peptides. Peptidoglycan crosslinking is mediated by direct (D)-alanyl-(D)-meso-diaminopimelic acid interpeptide linkages (Fig. 1). In mycobacteria, however, muramic acid is either acylated or glycolylated (12). The α -carboxylate of D-glutamic acid can be amidated. A glycine residue substitutes for the L-alanine residue at the amino end of the stem peptides in *Mycobacterium leprae*.

Bacteria also exist which have the dual ability to manufacture a peptidoglycan of the (4 \rightarrow 3) type and another peptidoglycan of the (3 \rightarrow 3) type (Fig. 2). The (3 \rightarrow 3) interpeptide

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Mycobacterium, Corynebacterium, Bacillus, E.coli

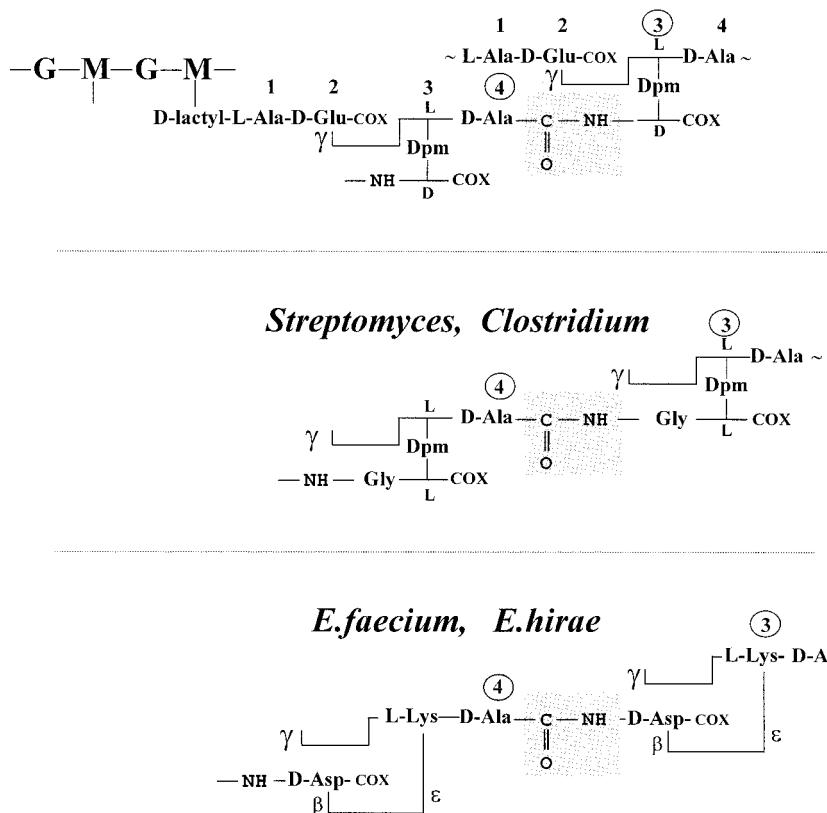


FIG. 1. (4→3) peptidoglycans. D-Alanyl-diaminoacyl interpeptide linkages and cross-bridges (boxed) between L-Ala-γ-D-Glu-(L)-diaminoacyl-d-alanine stem peptides. The diamino acid residues are meso-diaminopimelic acid (Dpm), L, L-diaminopimelic acid, or L-lysine. The stem peptides are unsubstituted at position 3 in *Mycobacterium*, *Corynebacterium*, and *Bacillus* spp. and *E. coli*. They are substituted at position 3 by one or several additional amino acid residues ($\alpha\alpha$) in the other organisms shown. G, N-acetylglucosamine; M, N-acetylmuramic acid (see Fig. 4); COX = COOH or CONH₂. In *S. pneumoniae*, the cross-bridges are N^{ϵ} -(L-alanyl-L-alanyl)- or (L-alanyl-L-seryl)-L-lysine. In *S. aureus*, the cross-bridges comprise five glycine residues or three glycine and two L-serine residues.

linkages or cross-bridges extend from the α -carbonyl of the diaminoacid residue (L-center) at position 3 of a stem peptide to the lateral amino group at position 3 of another, unbranched or branched, stem peptide.

The identification in the early 1970s of the occurrence of (3→3) peptidoglycans in *Streptomyces albus* G, *Clostridium perfringens* (139), *Mycobacterium smegmatis* ATCC 21732, and *Mycobacterium tuberculosis* BCG Pasteur strain (244) came as an exclamation point. Mycobacteria have a highly crosslinked peptidoglycan. About 70% to 80% of the total meso-diaminopimelic acid residues are involved in interpeptide linkages. In *Mycobacterium smegmatis* grown in Sauton's medium in Roux bottles for 9 days at 37°C, the (4→3) and (3→3) peptidoglycans

occur in the proportion of about 2 to 1. The pattern of distribution of the two peptidoglycans is not known, but (3→3)-linked peptide trimers occur in *M. smegmatis* and *M. tuberculosis* BCG.

Subsequently, it was found that *Escherichia coli* manufactures a (3→3) peptidoglycan (Fig. 2) in increasing proportions of total peptidoglycan and becomes increasingly more resistant to β -lactam antibiotics as the generation time increases (227). Penicillin-induced lysis also causes an increased proportion of (3→3) peptidoglycan (127). The contribution of (3→3) peptidoglycan in *Aeromonas* spp., *Acinetobacter acetoaceticus*, *Agrobacterium tumefaciens*, *Enterobacter cloacae*, *Proteus morganii*, *Pseudomonas aeruginosa*, *Pseudomonas putida*, *Sal-*

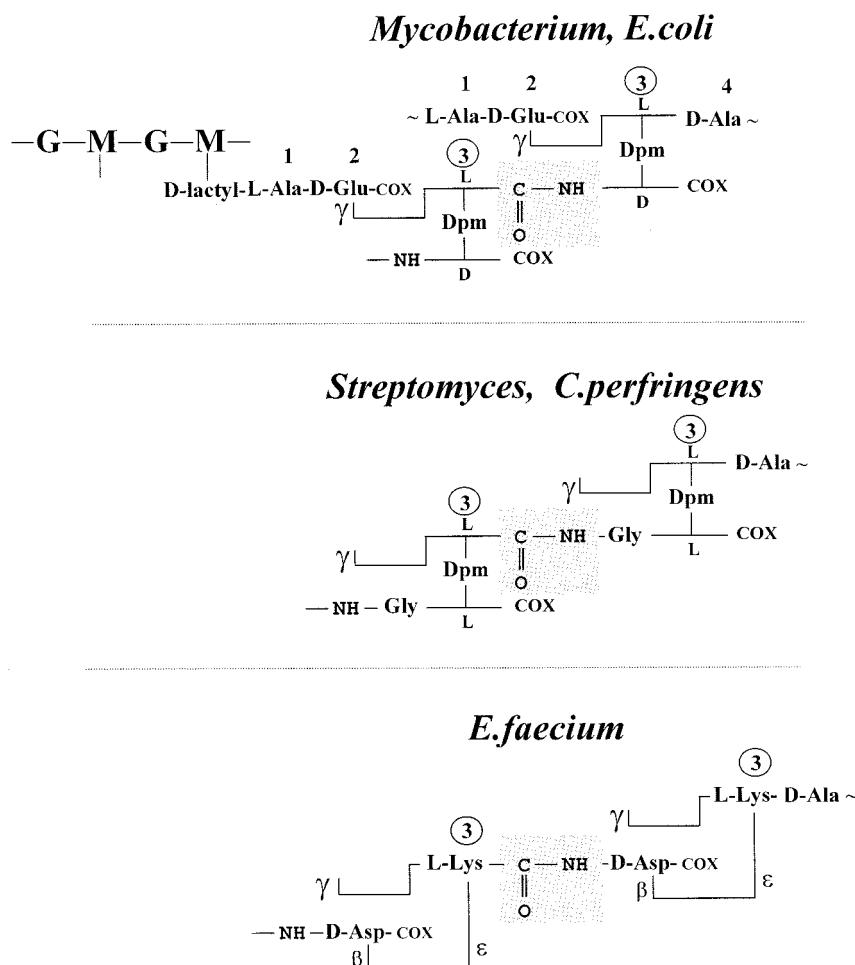


FIG. 2. (3→3) peptidoglycans. Diaminoacyl-diaminoacyl interpeptide linkages and cross-bridges (boxed) between L-Ala-γ-D-Glu-(L)-diaminoacyl-D-alanine stem peptides. The stem peptides are unsubstituted at position 3 in *Mycobacterium* spp. and *E. coli*. They are substituted at position 3 by Gly in *Streptomyces* spp. and *Clostridium perfringens* and by β-D-Asp in *E. faecium*. For details, see the legend to Fig. 1.

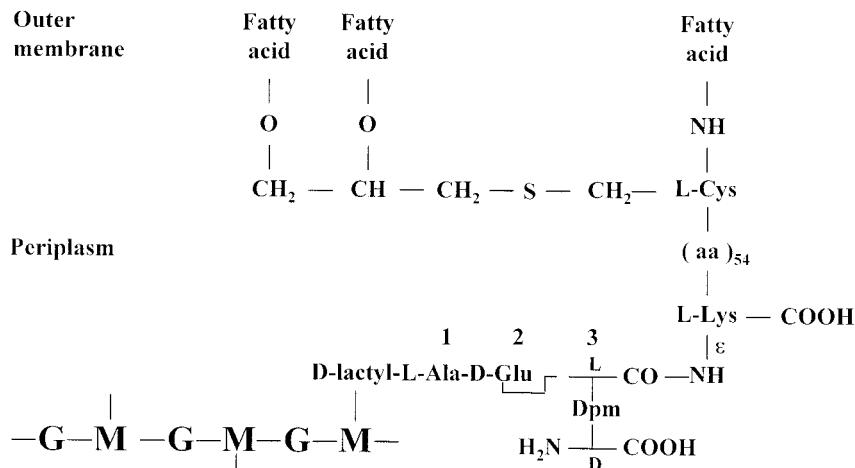
monella enterica serovar Typhimurium, *Vibrio parahaemolyticus*, *Yersinia enterocolitica*, and *E. coli* grown to late exponential phase varies from 1% to 45% of total peptidoglycan (183, 184). Finally, *Enterococcus faecium* is also a (3→3) peptidoglycan manufacturer (Fig. 2). A laboratory mutant has been isolated which resists penicillin in the exponential phase of growth, conditions under which it manufactures a wall peptidoglycan of the (3→3) type exclusively (145, 205).

From the foregoing, it follows that, in all likelihood, (3→3) peptidoglycan crosslinking is carried out by acyltransferases that escape penicillin action and that, in particular genetic backgrounds or under specific growth conditions, the penicillin-resistant (3→3) peptidoglycan assembly molecular machine can substitute for the penicillin-susceptible (4→3) peptidoglycan assembly molecular machine. This conclusion raises questions of fundamental and practical importance, related, in particular, to the mycobacterial pathogens.

E. coli has a 4.60-Mb genome (24). *M. tuberculosis* H37Rv has a 4.41-Mb genome (38). *Mycobacterium leprae* has a 3.27-Mb genome (39) that shows extensive decay and down-sizing. *M. tuberculosis* and *M. leprae* share about 1,500 genes. As stated above, *E. coli* and *Mycobacterium* spp. manufacture

similar, unbranched, *meso*-diaminopimelic acid-containing peptidoglycans of the (4→3) and (3→3) types. The nonpeptidoglycan wall polymers, however, are different. In *E. coli*, several lipoproteins are linked to the peptidoglycan. The 56-amino-acid residue Braun's lipoprotein (29) contains one fatty acid bound as an amide to the amino group of a cysteine residue and two fatty acids bound as esters to the hydroxyl groups of *S*-glycetyl cysteine (Fig. 3). The fatty acids are embedded in the outer membrane. Lipoprotein molecules occur both in a free form and in covalent linkage with the underlying peptidoglycan. The linkage is an amide bond between the ε-amino group of the L-lysine residue at the carboxy end of the lipoprotein and the carbonyl group at the L-center of the *meso*-diaminopimelic acid residue at position 3 of a stem tripeptide of the peptidoglycan.

In mycobacteria, mycolic acid, arabinogalactan, and peptidoglycan form a covalent complex (30). Mycolic acids are 2-alkyl 3-hydroxy branched-chain fatty acids. They are ester linked to arabinogalactan, which itself is linked to carbon C-6 of muramic acid via phosphodiester bonds. One may also note that a large portion of the coding ability to *M. tuberculosis* H37Rv is involved in lipid and polyketide metabolism (38) and

FIG. 3. Braun's lipoprotein-peptidoglycan complex of *E. coli*. For details, see the legend to Fig. 1.

8% of the genome is devoted to the production of two families of glycine-rich proteins of repetitive structure (20, 186).

M. tuberculosis and *M. leprae* have different life styles. *M. tuberculosis* enters host macrophages at cholesterol-rich domains of the plasma membrane (75). It survives in the macrophage by preventing fusion of the mycobacterial phagosomes with lysosomes (50, 201, 218, 236). It may persist inside macrophages lodged in calcified structures of the lungs and be reactivated decades after initial infection (153). It grows in synthetic media with a generation time of 12 to 24 h. Tuberculosis, once almost vanquished, resurged in the late 1980s because of the emergence of multidrug-resistant strains (25, 250). Often acting in deadly combination with AIDS, it kills more than 2 million people each year. *Mycobacterium leprae* enters the body via the mucosal linings of the nose or through open wounds. It shows a marked tropism for myelin-producing Schwann cells, which insulate nerves (187, 188). It may be dormant for years until the body's immune system attacks the infected cells, destroying the nerves and degrading soft tissues and even bones. It is uncultivable but can be grown in the nine-banded armadillo and the footpad of mice with an estimated generation time of about 2 weeks. Leprosy still flourishes in developing countries, where more than 750,000 people contract the disease each year.

Consistent with its ability to manufacture a (4→3) peptidoglycan, *M. tuberculosis* H37Ra (ATCC 25177, an attenuated laboratory strain) produces four major PBPs of 94, 82, 52, and 37 kDa (35). The inactivation of the 94-, 82-, and 52-kDa PBPs in cells grown to mid-exponential phase is associated with antibacterial activity. β-Lactamase production (240), more than the relatively low permeability of the cell envelope, is the major determinant of resistance (35, 63, 112, 185). Consequently, the MICs of ampicillin associated with the β-lactamase inactivator sulbactam are $\approx 0.1 \mu\text{g ml}^{-1}$ for *M. tuberculosis* H37Ra and $\approx 2 \mu\text{g ml}^{-1}$ for *M. tuberculosis* H37Rv (35). The drug combination is bactericidal to *M. tuberculosis* in exponential-phase cultures. It is bactericidal to *M. tuberculosis* multiplying in macrophages (35) and to *M. leprae* multiplying in mouse footpads (180, 189).

Paradoxically in view of the above data, the β-lactam anti-

biotics are not effective therapeutic agents for the treatment of tuberculosis and leprosy (101). This lack of efficiency could be due to a shift of peptidoglycan synthesis from the (4→3) type to the (3→3) type. Information that can help apprehend the problem was sought from comparative genomics of bacterial species (*E. coli*, *Enterobacter faecium*, *Mycobacterium* spp., and others) which have the ability to assemble, presumably from the same pool of precursors, peptidoglycans of the (4→3) and (3→3) types. Searches were also made for the positions of genes on the chromosomes because operons are common in prokaryotes and genes that are neighbors tend to be functionally linked.

PREDICTIVE STUDIES

Data concerning unfinished genome sequences were from the Institute for Genomic Research (TIGR) website at <http://www.tigr.org>. They were produced at TIGR with the support of the National Institute of Allergy and Infectious Diseases for *Mycobacterium avium* 104, at the Sanger Center/Institut Pasteur/VLA Weibrige with the support of MAFF/Beowulf Genomics for *Mycobacterium bovis* AF2122/97, at the Sanger Center/World Health Organization/Public Health Laboratory with the support of Beowulf Genomics for *Corynebacterium diphtheriae* NCTC 13129, and at the Sanger Center/John Innes Center with the support of BBSRC/Beowulf Genomics for *Streptomyces coelicolor* A3 (2).

Amino acid sequence similarities were searched with the Basic Local Alignment Search Tool Blast programs of the National Center for Biotechnology Information (NCBI) from their website at <http://www.ncbi.nlm.nih.gov/Blast> (4). Program BlastP compares a query amino acid sequence against a protein database. Program tBlastN compares a query amino acid sequence against a nucleotide sequence database dynamically translated in all six reading frames. *E. coli*, *M. tuberculosis*, *M. leprae*, *C. diphtheriae*, and *S. coelicolor* proteins were searched in the nonredundant protein database with BlastP and in the finished and unfinished genome database with tBlastN. The Expect value *E* of the NCBI's programs allows estimation of how far from the background noise is the

similarity between aligned sequences (3). For values of ≤ 0.01 , E becomes equivalent to the probability P that the sequences align by chance (118). P values smaller than 10^{-3} to 10^{-4} (depending on the sizes of the databanks) indicate statistically significant similarity.

The P values depend on the number and size of the gaps introduced to optimize the alignments. When the P value is small, equal or close to 0.0, the percentage of identical amino acid residues allows the sequences to be hierarchically classified. Combining the P values, the lengths of the overlapping regions, and the percentages of identities (I) present in the aligned regions gives an estimate of the extent of similarity between the sequences under comparison. Proteins encoded by genes having a common ancestor are homologues (193). Proteins encoded by genes related by duplication within a genome followed by diversification normally evolve new functions (103); they are paralogues. Homologous proteins encoded by genes in different organisms separated by speciation are orthologues (51, 103).

Protein orthologues that are essential and related by small P values and large I values normally retain similar biological functions. This conclusion is especially pertinent when low P values and large I values apply, over the entire sequences, to protein fusions for which the constitutive modules evolved from different protein ancestors. In contrast, little information is obtained when the similarities are restricted to peptide stretches or amino acid groupings. Conserved motifs defining the boundary of a catalytic center give valuable information on the catalytic mechanism, not on the specificity and fate of the reactions catalyzed.

LIPID II PRECURSORS: PEPTIDOGLYCAN ASSEMBLY

Lipid II molecules are the immediate biosynthetic precursors of the wall peptidoglycans (102). A disaccharide peptide exposed on the outer face of the plasma membrane is linked to a $C_{55}H_{89}$ undecaprenyl carrier via a pyrophosphate bridge involving carbon C-1 of *N*-acetylmuramic acid. The stem peptide borne by *N*-acetylmuramic acid is a pentapeptide terminating in D-alanyl-D-alanine. It can be unsubstituted or branched at position 3.

In *E. coli*, the synthesis of lipid II (Fig. 4) involves an interchange of carriers that are compatible with the environments of the cell (231). UDP-*N*-acetylglucosamine is converted into UDP-*N*-acetylglucosamine-enolpyruvate by MurA and from this into UDP-*N*-acetylmuramic acid by MurB. The Ddl (ATP,ADP + P_i) ligase catalyzes the formation of a D-alanyl-D-alanine dipeptide, and the MurC, MurD, MurE, and MurF (ATP,ADP + P_i) ligases catalyze the formation of UDP-*N*-acetylmuramoyl pentapeptide by the sequential additions to UDP-*N*-acetylmuramic acid of L-alanine, D-glutamic acid, meso-diaminopimelic acid, and the preformed D-alanyl-D-alanine dipeptide. Then, MraY transfers the phospho-*N*-acetylmuramoyl pentapeptide from its uridylic carrier to a membrane-bound C_{55} -isoprenoid alcohol phosphate, giving rise to lipid I. MurG transfers the *N*-acetylglucosamine from its uridylic carrier to carbon atom C-4 of *N*-acetylmuramic acid, giving rise to lipid II. Somehow, lipid II flips over the membrane bilayer, and the disaccharide pentapeptide moiety is exposed on the outer face of the membrane (Fig. 4). The genes

ddl, *murC*, *murD*, *murE*, *murF*, *murG*, and *mraY* reside in the *dcw* (or *mra*) cluster at the 2-min region of the chromosome (Fig. 5).

The stem pentapeptides can be modified at different stages of the biosynthesis of lipid II (234). Amidation of the α -carboxylate of D-glutamic acid (88) and the carboxylate at the D-center of meso-diaminopimelic acid probably takes place at the level of the UDP-*N*-acetylmuramoyl pentapeptide precursors. In many gram-positive bacteria, the diaminoacid residue at position 3 of the stem pentapeptide is L-lysine. Addition of one or several amino acid residues to the ϵ -amino group of the L-lysine residue, resulting in the formation of branched-stem pentapeptides, probably occurs at the level of lipid II while it is still oriented toward the cytosol.

In *Staphylococcus aureus*, there are four glycyl tRNAs. One is involved in protein synthesis. The three others seem to be used exclusively for peptidoglycan synthesis. Lipid II precursor molecules with one, three, and five glycine residues attached to the ϵ -amino group of the L-lysine residue are formed by the sequential actions of the glycine-adding enzymes FemX (also called FemhB), FemA, and FemB (128, 197). Loss of FemX is lethal. In some strains, the FemAB-like Lif and Epr incorporate an L-serine residue instead of a glycine residue within the branches at positions 3 and 5 (60). In *Streptococcus pneumoniae*, the N^e -(L-alanyl-L-alanyl)-L-lysine and N^e -(L-alanyl-L-seryl)-L-lysine branches are synthesized by MurM and MurN, respectively (64, 242), with MurM being responsible for the addition of the first amino acid residue to the ϵ -amino group of L-lysine.

Vancomycin-resistant enterococci are programmed to produce lipid II precursor molecules which terminate in D-alanyl-D-lactate (8, 136). Five enzymes are necessary and sufficient. The Zn²⁺-dependent VanX dipeptidase hydrolyzes the D-alanyl-D-alanine dipeptide produced by the Ddl ligase. VanH and VanA act sequentially to synthesize the depsipeptide D-alanyl-D-lactate, which is incorporated in lipid II rather than the dipeptide D-alanyl-D-alanine. The production of VanHAX is controlled by a two-component regulatory system that involves the transmembrane sensor kinase VanS and the transcription factor VanR, which becomes active when phosphorylated by VanS (8).

The conversion of the disaccharide peptide (depsipeptide) moiety of lipid II into polymeric (4 \rightarrow 3) peptidoglycan requires the sequential actions of two transferases (Fig. 4). A glycosyltransferase catalyzes attack of carbon C-1 of *N*-acetylmuramic acid of a lipid II molecule by the acceptor nucleophile OH of carbon C-4 of *N*-acetylglucosamine of another lipid II molecule (or at the nonreducing end of a growing glycan chain; not shown). At each transfer, the delivery of a disaccharide peptide unit generates an undecaprenyl pyrophosphate which is dephosphorylated. The C_{55} -isoprenoid alcohol phosphate turns over the membrane bilayer so that the phosphate group faces the cytosol, allowing a new cycle to start.

In turn, the required D,D-acyltransferase (Fig. 4 and reaction I in Fig. 6) catalyzes attack of the carbonyl of the D-alanine residue at position 4 of a pentapeptide (depsipeptide) borne by a glycan chain, by the lateral amino group at position 3 of a peptide borne by another glycan chain. The reaction proceeds via the transitory formation of an *N*-acyl-D-alanyl-enzyme intermediate in which the D-alanine residue is linked as an ester

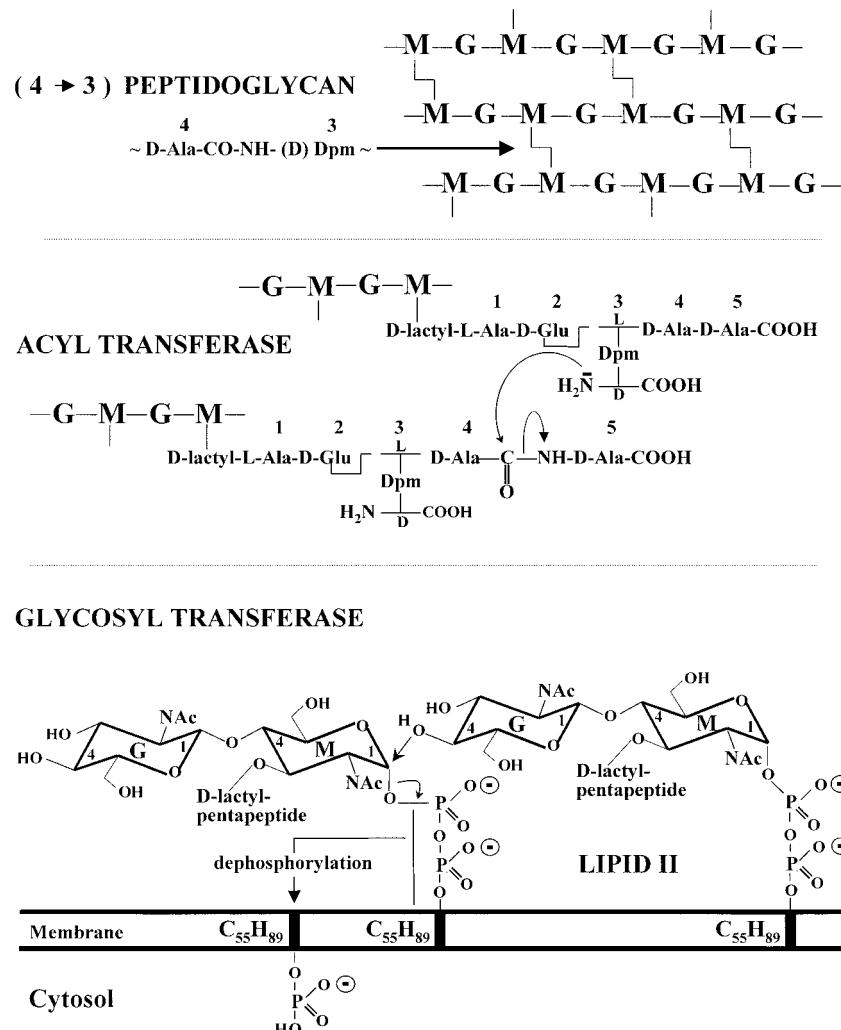


FIG. 4. Lipid II precursor (bottom) and polymeric ($4 \rightarrow 3$) peptidoglycan (top) of *E. coli*. Reactions catalyzed by the glycosyl- and acyltransferases. For details, see the legend to Fig. 1.

to a serine residue at the catalytic center (69, 70, 78). The carbonyl donor involved in enzyme acylation is invariably a D-alanyl-D-alanine(-D-lactic acid) sequence. In contrast, the acceptor involved in enzyme deacylation can be the amino group of a glycine residue, the ϵ -amino group of an L-lysine residue, or an amino group borne by an L- or a D-configured carbon atom (Fig. 1).

Water can also serve as an acceptor. Stem pentapeptides are hydrolyzed into stem tetrapeptides, preventing further ($4 \rightarrow 3$) peptidoglycan crosslinking. The β -lactam antibiotics act as suicide carbonyl donors of the D,D-acyltransferases (224). Rupture of the β -lactam amide bond produces a serine ester-linked acyl (penicilloyl, cephalosporoyl, ...) enzyme, which is a dead end because the bulky acyl moiety is a steric hindrance to the approach of any attacking nucleophile. The catalytic center turns over very slowly, once or less per hour (71, 72), and the inactivated D,D-acyltransferases are detectable as PBPs (214).

No one knows exactly how the ($3 \rightarrow 3$) peptidoglycans are made in a penicillin-resistant manner. The ($3 \rightarrow 3$) interpeptide linkages or cross-bridges could be made through the action of

an L,D-acyltransferase that cleaves the bond between position 3 and position 4 of pentapeptide carbonyl donors, with release of the dipeptide D-alanyl-D-alanine (reaction II in Fig. 6). Alternatively, they could be made through the sequential actions of a D,D-carboxypeptidase catalyzing reaction IIIa (Fig. 6) and an L,D-acyltransferase that cleaves the bond between position 3 and position 4 of tetrapeptide carbonyl donors (reaction III in Fig. 6). Reactions IIIa and III each cause the release of a single D-alanine residue.

SxxK ACYLTRANSFERASE SUPERFAMILY

The serine acyltransferases implicated in wall peptidoglycan assembly are members of a superfamily of SxxK serine enzymes. The term superfamily is used to include proteins with no statistically significant sequence similarities but with similar structures in the classical sequence-based families (169). With x denoting a variable amino acid residue, the acyltransferases of this superfamily have a specific bar code in the form of three motifs, SxxK, SxN (or analogue), and KTG (or analogue). The

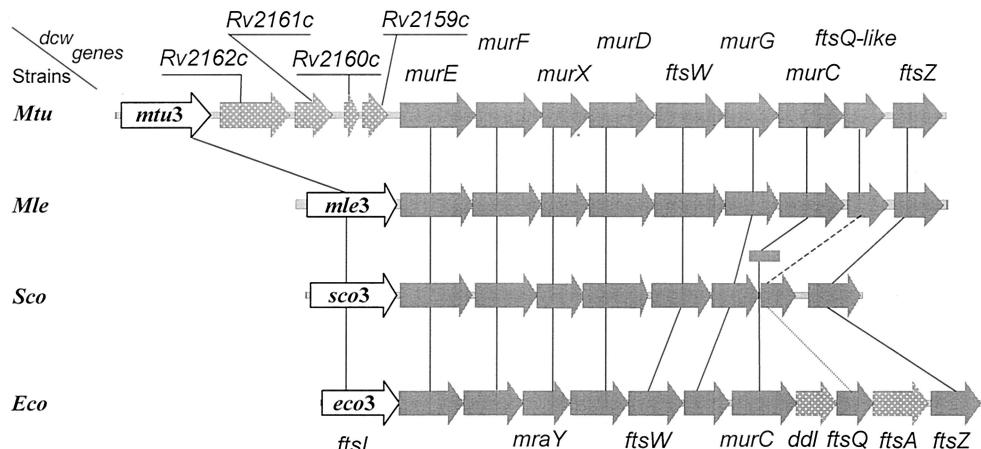


FIG. 5. Organization of the division cell wall (dcw) gene clusters of *M. tuberculosis* (*Mtu*), *M. leprae* (*Mle*), *Streptomyces coelicolor* (*Sco*), and *E. coli* (*Eco*). All the coding sequences are on the same DNA strands. Open arrows, *mtu3*, *mle3*, *sco3*, and *eco3* (synonymous to *ftsI*), coding for cell septation SxxK PBP fusions. Shaded arrows, with *fts* standing for the filamentous phenotype of thermosensitive *E. coli* mutants, *ftsW*, *ftsQ*, *ftsA*, and *ftsZ* code for cell septation-related, non-penicillin-binding proteins. Genes *murE*, *murF*, *murX* (synonymous to *mraY*), *murD*, *murG*, and *murC* code for enzymes of the lipid II-synthesizing pathway. In *S. coelicolor*, *murC* (shaded rectangle) is not in the *dcw* cluster. Stippled grey arrows, genes of a *dcw* cluster having no homologues in the other clusters. In *M. tuberculosis*, the genes inserted between *mtu3* and *murE* probably code for a glycine-rich protein of repetitive structure (*Rv2162c*), a protein of the lincomycin-synthesizing pathway (*Rv2161c*), and proteins of unknown function (*Rv2160c* and *Rv2159c*). A gene homologous to *E. coli* *ddl* (coding for the ligase which catalyzes the formation of D-alanyl-D-alanine) is located elsewhere in the chromosomes of *M. tuberculosis*, *M. leprae*, and *S. coelicolor*. *E. coli* *ftsA* codes for a cell division, actin-like ATPase. Solid, dashed, and dotted lines connecting the *dcw* genes indicate the extents, in decreasing order, of similarity between homologous genes. The *E. coli* cell septation genes *mraZ*, *mraW*, and *ftsL* (not shown) are upstream from *eco3*.

motifs occur at equivalent places and with roughly the same spacing along the polypeptide chains. In the three-dimensional structures, they are brought close to each other at the immediate boundary of the catalytic center between an all- α domain and an α/β domain, the five-stranded β -sheet of which is covered by α -helices (79, 123, 124).

The serine residue of the invariant motif SxxK occupies a central position in the catalytic center and is central to the

enzyme acylation and deacylation steps, which, mechanistically, are similar to the proteolytic reactions of the trypsin protein family. The mature SxxK acyltransferase of *Streptomyces* sp. strain K15 is 262 amino acid residues long. It is one of the smallest members of the superfamily (Fig. 7) (68). Diverging evolution gave rise to a constellation of SxxK acyltransferases with various amino acid sequences. Fusion to other polypeptide chains resulted in a combinatorial system of struc-

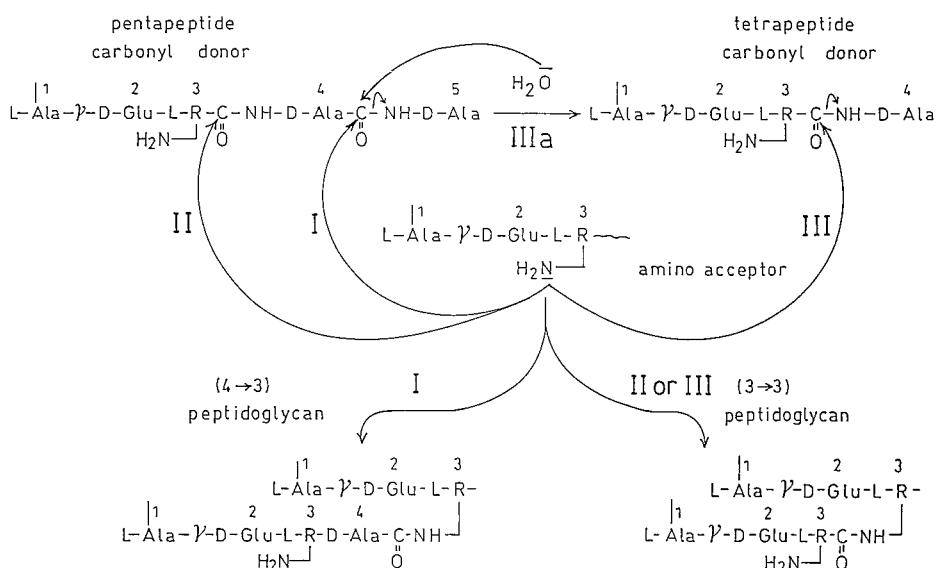


FIG. 6. Acyl transfer reactions. Formation of (4 \rightarrow 3) peptidoglycan interpeptide linkages by penicillin-susceptible D,D-N-acyl-D-alanyl-D-alanine transpeptidases (reaction I). Formation of (3 \rightarrow 3) peptidoglycan interpeptide linkages by penicillin-resistant L,D-N-acyl-L-diaminoacyl-D-dipeptidyl transpeptidases (reaction II) and L,D-N-acyl-L-diaminoacyl-D-alanine transpeptidases (reaction III). Hydrolysis of stem pentapeptides into stem tetrapeptides by penicillin-resistant D,D-N-acyl-D-alanyl-D-alanine carboxypeptidases (reaction IIIa).

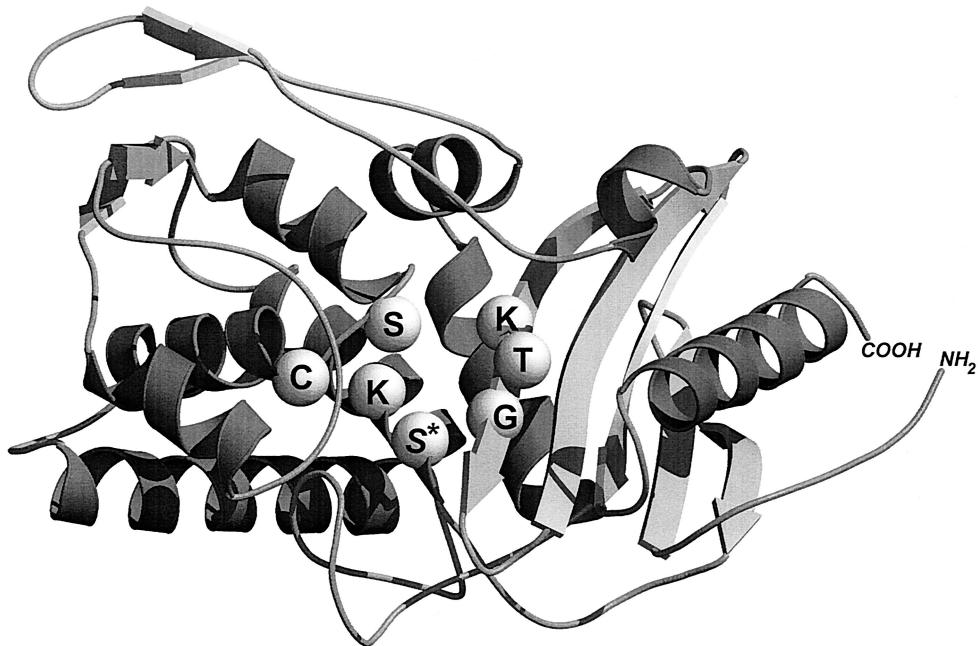


FIG. 7. Basic polypeptide fold of the acyltransferases of the SxxK superfamily and spatial disposition of the three catalytic center-defining amino acid groupings. The structure shown is that of the 262-amino-acid DD-transpeptidase-PBP of *Streptomyces* sp. strain K15. The SxxK acyltransferases comprise an all- α domain (left side) and an α/β domain (right side). The invariant motif 1, S^*xxK , where S^* is the essential serine nucleophile, forms the amino-terminal turn of helix α 2 of the all- α domain. Motif 2, most often SxN or SxD (here SxC), is on a loop connecting two helices of the all- α domain. Motif 3, most often KTG or KSG , is on strand β 3 of the α/β domain. The structure was built with Molscript (130) and Raster3D (155). Illustration courtesy of Paulette Charlier, Eveline Fonzé, Michaël Delmarcelle, and André Piette, Center for Protein Engineering.

tural modules that contributed to a massive increase in functional diversity.

The acyltransferases of the SxxK superfamily are a paradigm of catalytic versatility. They fall into three main groups. The D,D-acyltransferases-PBPs, implicated in (4→3) peptidoglycan synthesis, are of group I. Acyltransferases endowed with diverse functions unrelated to peptidoglycan biochemistry are of group II. Penicillin-resistant acyltransferases that could act as L,D-peptidases in (3→3) peptidoglycan synthesis are of group III. The ensuing sections provide critical coverage of mature topics and speculations on emerging topics. The identifiers of the SxxK acyltransferases are shown in Table 1. They combine the name of the producing bacterial species as a three-letter

prefix flanked by a number or additional letters that specify the protein under consideration (e.g., Eco1a, protein 1a of *E. coli*).

STRUCTURE-ACTIVITY RELATIONSHIPS OF β -LACTAM ANTIBIOTICS

The SxxK D,D-acyltransferases of group I are the target proteins of the β -lactam antibiotics. These compounds are a large group of molecules of which the common structural feature is the presence of a β -lactam (azetidinone) ring (Fig. 8). Currently, they are regarded as a molecular mimic of *N*-acyl-D-alanyl-D-alanine (224), explaining why they are suicide car-

TABLE 1. Prefixes identifying bacterial species

Prefix	Species	Prefix	Species
Aae	<i>Aquifex aeolicus</i>	Mtu	<i>Mycobacterium tuberculosis</i>
Ana	<i>Anabaena</i> sp. strain PCC7120	Nme	<i>Neisseria meningitidis</i>
Bbu	<i>Borrelia burgdorferi</i>	Pae	<i>Pseudomonas aeruginosa</i>
Bsu	<i>Bacillus subtilis</i>	Sau	<i>Staphylococcus aureus</i>
Cdi	<i>Corynebacterium diphtheriae</i>	Scl	<i>Streptomyces clavuligerus</i>
Eco	<i>Escherichia coli</i>	Sco	<i>Streptomyces coelicolor</i>
Efam	<i>Enterococcus faecium</i>	Sgr	<i>Streptomyces griseus</i>
Efas	<i>Enterococcus faecalis</i>	Sor	<i>Streptococcus oralis</i>
Ehi	<i>Enterococcus hirae</i>	Spn	<i>Streptococcus pneumoniae</i>
Hin	<i>Haemophilus influenzae</i>	Ssc	<i>Staphylococcus sciuri</i>
Hpy	<i>Helicobacter pylori</i>	Spy	<i>Streptococcus pyogenes</i>
Mle	<i>Mycobacterium leprae</i>	Sth	<i>Streptococcus thermophilus</i>
Msm	<i>Mycobacterium smegmatis</i>	Syn	<i>Synechocystis</i> sp. strain PCC6203

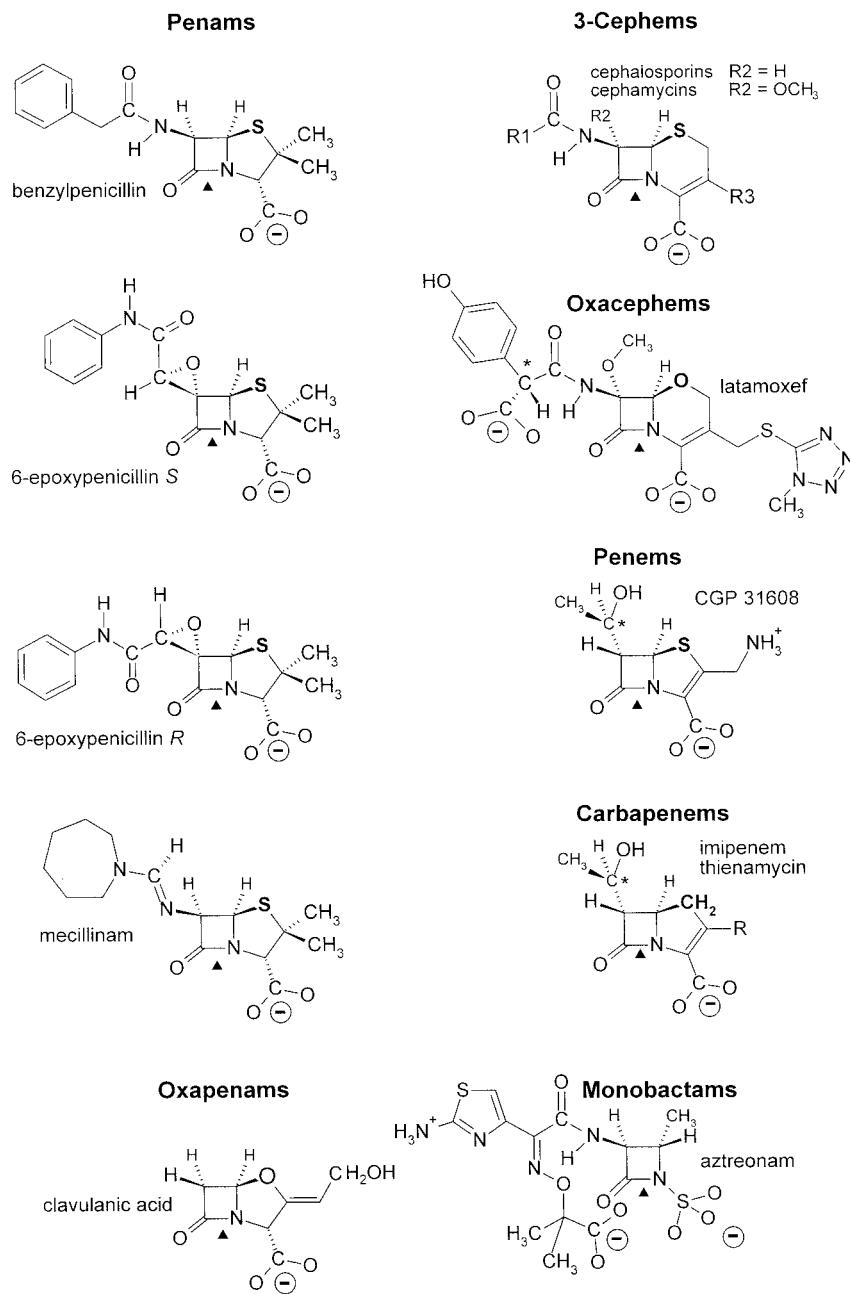


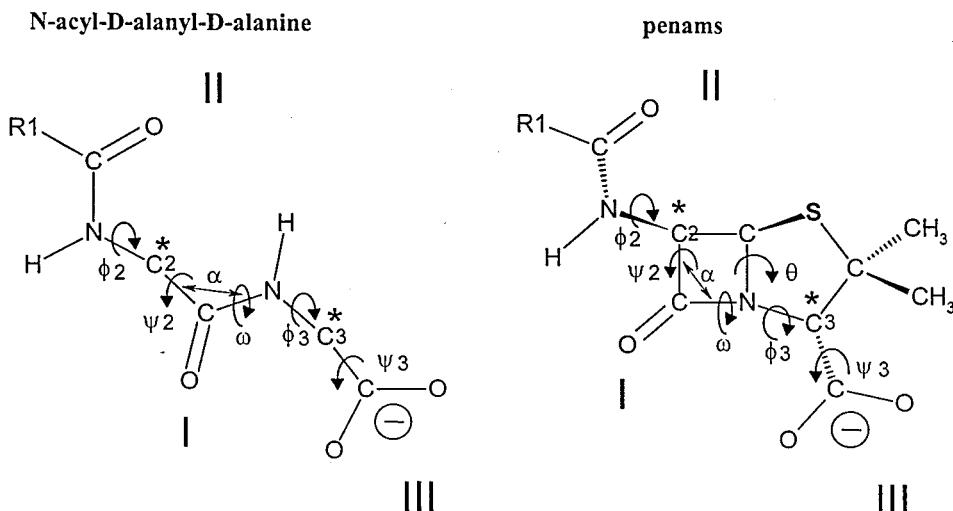
FIG. 8. β -Lactam antibiotic family. Only 6-epoxypenicillin S is active. *, D-configured carbon atom. ("Mecillinam" is another name for amdinocillin.)

bonyl donors of the D,D-acyltransferases–PBPs implicated in (4 \rightarrow 3) peptidoglycan synthesis (Fig. 4 and reaction I in Fig. 6).

The penams, 3-cephem, and N-acyl-D-alanyl-D-alanine-terminated peptides (in the extended conformation) have a common backbone, CON-C₂-CON-C₃-COO⁻ (Fig. 9, upper part). Carbon atoms C-2 and C-3 (marked by asterisks) have a D-configuration. They are the pivots that connect the central CON amide plane at position I to the CON amide plane at position II (through the rotation angles ψ_2 and ϕ_2) and the carboxylate at position III (through the rotation angles ϕ_3 and ψ_3). The scissile CON bonds at position I, however, are far from being isosteric. The nitrogen atom is planar in the pep-

tides. It is pyramidal in the azetidinones. The bond angle α is 117° in the peptides and 90° in the azetidinones. The rotation angle ω is 180° in the peptides, 155° in the 3-cephem, and 135° in the penams.

Rather than being related to similarities between linked atoms, structural analogy between the most stable D-alanyl-D-alanine-terminated peptide, penam, and 3-cephem conformers relies on the spatial disposition of the electrostatic negative wells created by the carbonyl CO at position I, the carbonyl CO at position II, and the carboxylate COO⁻ at position III (80, 131, 132). These three groupings are almost coplanar, and the spanning distances between the oxygen atoms at position I and

Common backbone CON-C₂-CON-C₃-COO⁻

Three-dimensional electrostatic potentials

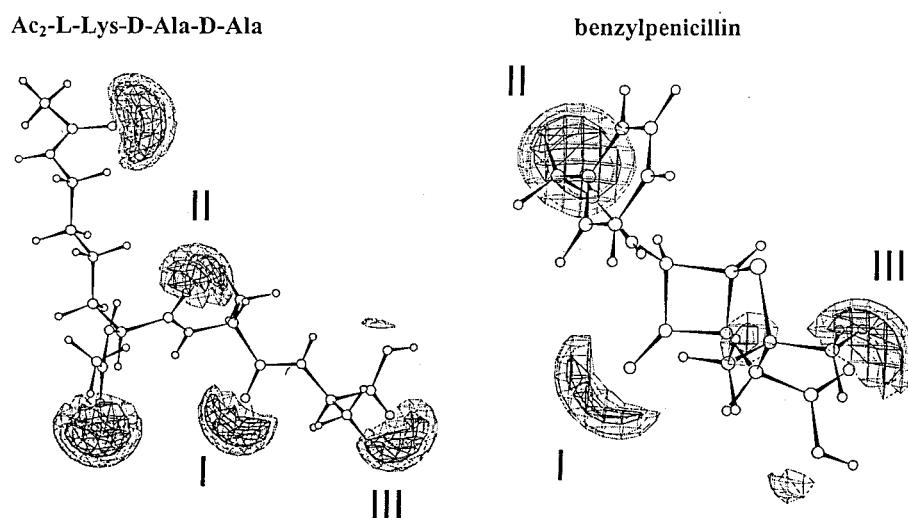


FIG. 9. Common backbone of extended *N*-acyl-*D*-alanyl-*D*-alanine peptides and penams (top) and ab initio electrostatic potentials of bisacetyl-*L*-lysyl-*D*-alanyl-*D*-alanine and benzylpenicillin (bottom) optimized at level AM1 (the terminal carboxylates are protonated). The electrostatic negative wells I, II, and III are shown at levels of -40 kcal (solid contours) and -30 kcal (dotted contours). They are coplanar. The negative wells generated by the acetyl substituents of the α - and ϵ -amino groups of the *L*-lysine residue of bisacetyl-*L*-lysyl-*D*-alanyl-*D*-alanine are above and below the plane, respectively. The negative well of small amplitude seen between well I and well III of benzylpenicillin is due to the pair of free electrons of the nitrogen atom of the azetidinone ring. Illustration courtesy of Georges Dive, Center for Protein Engineering.

position II ($\approx 5 \text{ \AA}$) and between the oxygen atom at position I and the carbon atom of the carboxylate at position III (3.5 to 4.5 \AA) are roughly similar. Figure 9 (lower part) shows the three-dimensional electrostatic potentials of the optimized benzylpenicillin and extended bisacetyl-*L*-lysyl-*D*-alanyl-*D*-alanine molecules. The two additional negative wells seen in the peptide are due to the acetyl groupings that substitute the α - and ϵ -amino groups of the *L*-lysine residue.

As a result of the coplanarity of the negative wells at posi-

tions I, II, and III, the α -face of the azetidinone ring of the β -lactam antibiotics is well exposed, facilitating the attack of the electrophilic center at position I by the serine nucleophile of the SxxK D,D-acyltransferases. The backbone $\text{CON-C}_2\text{-CON-C}_3\text{-COO}^-$ is modified into $\text{CON-C}_2\text{-CON-SO}_3^-$ in the monobactam aztreonam (Fig. 8). It is modified into $\text{C=N-C}_2\text{-CON-C}_3\text{-COO}^-$ in the penam amdinocillin. The carbon atom C-2 of the carbapenem imipenem has an L-configuration. In spite of these structural variations, the disposition of the sulfa-

mate in aztreonam is roughly comparable to that of the carboxylate at position III in the bicyclic penams and 3-cephems. The II environment of the CH=N amidino bond of amdinocillin generates a negative well at position II, albeit of reduced amplitude. Because the rotation angle θ (Fig. 9, upper part) is less open in the carbapenems than in the penams and 3-cephems, the carboxylate at position III of imipenem is moved upward, ensuring coplanarity with the oxygen atom at position I and the alcohol oxygen atom at position II. Of the two 6-epoxypenicillin isomers (Fig. 8), the side chain of which, at position II, has a frozen conformation due to the epoxy cycle, the *S* isomer is active, but the *R* isomer is not.

Depending on the (noncyclic, monocyclic, or bicyclic) framework and conformation of the backbone and the presence of bulky, ionized, and electron-withdrawing side chains, the electrostatic negative wells at positions II and III around the electrophilic center at position I of the carbonyl donors can vary in shape and strength, be displaced along the reference plane, be better expressed in other sections of the molecules, be fused, or be masked. In turn, the SxxK acyltransferases are large bodies studded with positively and negatively charged magnets. Upon binding of a carbonyl donor to the catalytic center, a dense hydrogen bonding network is formed, and a multimembered ring that is both enzyme and ligand specific is created, in which the scissile CO-N amide bond at position I of the carbonyl donor is connected to the catalytic serine γ OH of motif SxxK of the acyltransferase by one or several water molecules and the side chains of several amino acid residues (53, 54, 80). This multimembered ring is utilized as a motorway, allowing the proton of the serine γ OH to be transferred via a transition state to the nitrogen atom of the scissile bond, resulting in enzyme acylation. For the reaction to reach completion, the serine ester-linked acyl enzyme must adopt a conformation that allows entry of an amino group or a water molecule and formation of a new multimembered ring that performs enzyme deacylation.

Potential energy hypersurfaces best portray the geometric rearrangements, electronic redistributions, and free energy barriers that occur along the reaction pathways (80). Depending on the freedom of the water molecules, the ease with which the ligands undergo deformation, and the ease with which the enzyme backbone undergoes relaxation, numerous possible routes for proton transfer exist. The energetically most favorable route dictates the specificity of the SxxK acyltransferases and the completeness (protein binding, protein acylation, protein acylation, and deacylation) and productiveness of the catalyzed reactions.

The D,D-acyltransferases-PBPs of group I perform multiple functions related to (4 \rightarrow 3) peptidoglycan assembly (see below). Changes in protein structure, with conservation of the overall fold, could go as far as a change from D,D specificity to L,D specificity (see the chapter on SxxK acyltransferases of group III).

GROUP I SxxK ACYLTRANSFERASES: IMPLICATED IN (4 \rightarrow 3) PEPTIDOGLYCAN BIOCHEMISTRY

The SxxK D,D-acyltransferases-PBPs of group I occur as independent entities referred to as free-standing PBPs. They also occur as modules of PBP fusions. A D,D-acyltransferase

module of class A or class B is fused, in a class-dependent manner, to polypeptides having their own bar codes and three-dimensional structures (78, 84). Traces of similarity between the free-standing PBPs and the acyltransferase modules of class A or B are generally limited to the three active-site defining motifs. Similarity between the acyltransferase modules of class A and class B is marginal ($P \geq 1 \times 10^{-5}$).

The PBPs are bound to the plasma membrane, with the bulk of the polypeptide chains exposed on the outer face. The PBP fusions are synthesized with an amino-terminal hydrophobic segment that functions as both a signal sequence for secretion and a stop transfer signal that serves as a membrane anchor. Despite a great divergence in the amino acid sequences, the bar codes SxxK, SxN, and KTG of the free-standing PBPs and the acyltransferase modules of the PBP fusions are conserved except that, occasionally, SxD substitutes for SxN and KSG substitutes for KTG.

The core of an SxxK acyltransferase is defined as the sequence starting about 70 amino acid residues upstream from the SxxK motif and terminating about 70 amino acid residues downstream from the KT(S)G motif. Adducts may occur as inserts and/or as carboxy-terminal extensions. The free-standing PBP5 of *E. coli* (45) and the class B PBP fusion Snp2x of *Streptococcus pneumoniae* (86, 173) are of known structure. They each have carboxy-terminal extensions. That of *E. coli* PBP5 is made of β -structures that form a loose β -barrel; that of Snp2x is made of two $\alpha/\beta/\beta/\beta$ domains.

Class A PBP Fusions: Nascent (4 \rightarrow 3) Peptidoglycan

The class A PBP fusions consist of a penicillin-binding acyltransferase module of class A linked to the carboxy end of a glycosyltransferase module, itself linked to the carboxy end of the membrane anchor (Fig. 10). The full bar code comprises motifs 1 to 5 of the glycosyltransferase module, motif 6 of the intermodule junction, and motifs 7 to 9 of the acyltransferase module (84). By combining a glycosyltransferase and an acyltransferase in a single polypeptide chain, class A PBP fusions catalyze the polymerization of the disaccharide pentapeptide units borne by lipid II precursor molecules into nascent (4 \rightarrow 3) peptidoglycans (Fig. 4) (162, 222, 232, 235).

The structure of the glycan backbones of the wall peptidoglycans is invariant. Consistently, the glycosyltransferase modules of the class A PBP fusions form a continuum of sequences (Fig. 10), indicating steady divergence and functional conservation. In contrast, the structures of the peptidoglycan interpeptide linkages and cross-bridges are variable (Fig. 1 and 2). Consistently, the acyltransferase modules of class A PBP fusions cluster into several subclasses (Fig. 10), indicating functional diversification. Modules of the same subclass are related by P values smaller than $\approx 1 \times 10^{-30}$. Subclasses A1 and A2 are typical of gram-negative bacteria. Subclasses A3, A4, and A5 are typical of gram-positive bacteria.

E. coli produces two class A PBP fusions, Eco1a and Eco1b (Fig. 11). The Eco1a-encoding gene, at the 75-min region of the chromosome, and the Eco1b-encoding gene, at the 4-min region, are not linked to particular operons. An insert occurs downstream from the membrane anchor in Eco1b. Inserts occur downstream from the intermodule junction and between motifs 8 and 9 in Eco1a. Eco1b occurs in three forms, each of

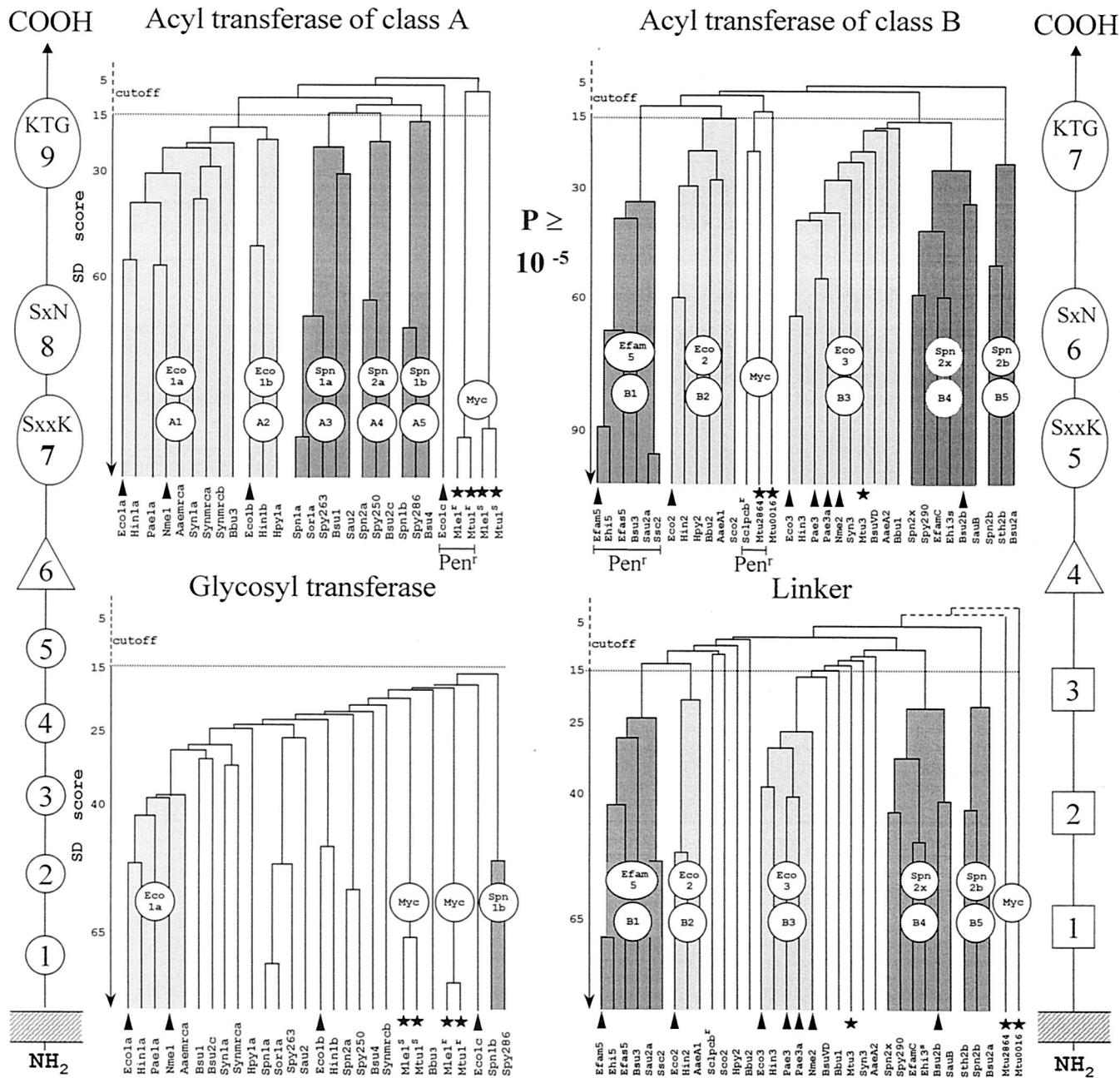


FIG. 10. Hierarchical distribution of the SxxK PBP fusions and Pen^r protein fusions of classes A and B. The occurrence of class-specific motifs 1 to 9 (class A) and 1 to 7 (class B) along the polypeptide chains is shown. Adapted from reference 84. Scores (vertical axis of the dendograms) are the standard deviation values above that expected from a run of 100 randomized pairs of sequences with the same amino acid composition as the two sequences under comparison. The protein identifiers (bottom of the dendograms) are defined in Table 1. The clusters are labeled by two circles, one of which defines a particular subclass (A1 to A5 and B1 to B5) and the other the prototypic protein. Solid arrowheads help identify proteins that are discussed in the text. Solid stars help identify the mycobacterial proteins. The Pen^r acyltransferase modules are underlined.

which is functional and encoded by the same gene (219). The α form, shown in Fig. 11, has a cytosolic amino-terminal tail which is 70 amino acid residues long. The glycosyltransferase modules of Eco1a and Eco1b are related by a *P* value of 4×10^{-33} (identity [*I*], 35%). The acyltransferase modules of Eco1a and Eco1b are distantly related by a *P* value of 2×10^{-15} (identity, 27%) for overlaps 190 amino acid residues long. They belong to distinct subclasses. Eco1a (subclass A1)

and Eco1b (subclass B2) are produced in large quantities, amounting together to several thousand molecules per cell. In *in vitro* assays in the absence of preformed peptidoglycan primer, they each catalyze the conversion of the disaccharide peptide moiety of lipid II into polymeric (4→3) peptidoglycan.

Lysozyme cleaves glycosidic bonds with net retention of configuration of the anomeric center via a covalent glycosyl-enzyme intermediate that involves Asp52 (239). The glycosyl-

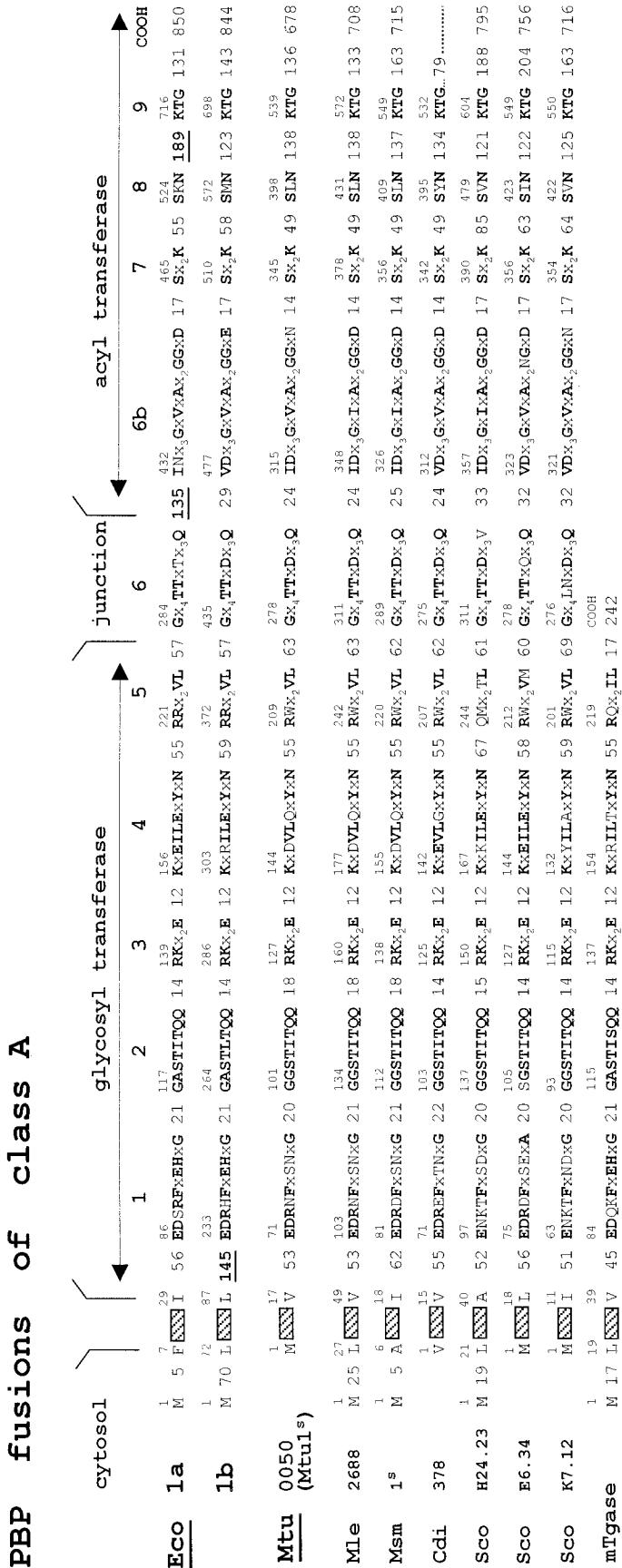


FIG. 11. Class A PBP fusions. Bar code, motifs 1 to 9 and inserts (underlined). For protein identifiers, see Table 1. mtgase, free-standing transglycosylase of *E. coli*.

transferase of class A PBP fusions make new glycosidic bonds with inversion of configuration at carbon C-1, from the α -configuration in lipid II to the β -configuration in the peptidoglycan (Fig. 4). The essential Glu233 of Eco1b, at the amino end of motif 1 of the glycosyltransferase module (Fig. 11), is involved in proton donation to the oxygen atom of the scissile phosphoester bond, resulting in the formation of a muramic oxocarbonium intermediate (222). Asp234 of motif 1 and Glu290 of motif 3 could be responsible for the activation of the 4-OH of the nucleophile *N*-acetylglucosamine and the attachment on the β -face of *N*-acetylmuramic acid.

Crosslinking between peptide-substituted glycan chains is an acylation-deacylation reaction that is strictly dependent on the serine residue of motif SxxK of the acyltransferase module. It is aided by amino acid residues of the other catalytic center-defining motifs and one or several catalytic water molecules. In *in vitro* assays with lipid II, Eco1b transfers the carbonyl of the D-alanine residue at position 4 of stem pentapeptides to water (reaction products, tetrapeptide monomers) and to the D-amino group of the *meso*-diaminopimelic acid residue at position 3 of pentapeptide and tetrapeptide monomers [reaction products: (4→3)-linked tetrapeptide-pentapeptide and tetrapeptide-tetrapeptide dimers]. Peptide oligomers larger than dimers are not produced in detectable amounts, indicating that, *in vitro*, the stem pentapeptide of the tetrapeptide-pentapeptide dimers is not used as a carbonyl donor for further crosslinking.

The acyltransferase of Eco1b is inert on exogenous *N*-acyl-D-alanyl-D-alanine-terminated peptides (222). However, it catalyzes hydrolysis and aminolysis of ester and thiolester analogues, indicating that the D-alanyl-D-alanine-cleaving activity is glycosyltransferase dependent. Conversely, Eco1b, the acyltransferase module of which is inactivated by penicillin, catalyzes glycan chain elongation, indicating that the glycosyltransferase module is acyltransferase independent.

Loss of Eco1a and Eco1b is fatal, but loss of either Eco1a or Eco1b is tolerated, indicating that Eco1a and Eco1b can substitute for each other (119, 220, 252). PBP fusions of subclass A2 are dispensable in some gram-negative bacteria. The coccus-shaped *Neisseria meningitidis*, the genome sequence of which is known (223), produces a single class A PBP fusion of subclass A1 (199). Consistently, the cluster formed by the acyltransferase modules of subclass A1 (to which Eco1a belongs) is more populated, i.e., comprises more sequences, than the cluster formed by the acyltransferase modules of subclass A2 (to which Eco1b belongs) (Fig. 10). It remains possible that Eco1a and Eco1b cause distinct, subtle traits in peptidoglycan crosslinking in *E. coli* that are difficult to detect (34).

Class B PBP Fusions: Morphogenetic Apparatus

The class B PBP fusions are components of the morphogenetic apparatus that are dynamic in abundance and composition. They control wall expansion, ensure cell shape maintenance, and direct and carry out septum formation and cell division (55, 146, 163). The class B PBP fusions consist of a penicillin-binding acyltransferase module of class B linked to the carboxy end of a module which does not have the bar code of a transglycosylase and is itself linked to the carboxy end of

a membrane anchor (Fig. 10). The non-penicillin-binding module mediates protein-protein interactions critical for peptidoglycan assembly in a cell cycle-dependent manner. For this reason, it is referred to as a linker (or protein recognition) module. The full bar code of the class B PBP fusions comprises motifs 1 to 3 of the linker module, motif 4 of the intermodule junction, and motifs 5 to 7 of the acyltransferase module.

The acyltransferase and linker modules diverged in concert (Fig. 10). In gram-negative bacteria, an acyltransferase module of subclass B2 or B3 is fused to a linker module of subclass B2 or B3, respectively. In gram-positive bacteria, an acyltransferase module of subclass B4 or B5 is fused to a linker module of subclass B4 or B5, respectively. *E. coli* produces two class B PBP fusions, Eco2 of subclass B2 and Eco3 of subclass B3 (Fig. 10 and 12). The 588-amino-acid Eco3 is processed into M1-V577 Eco3 (161).

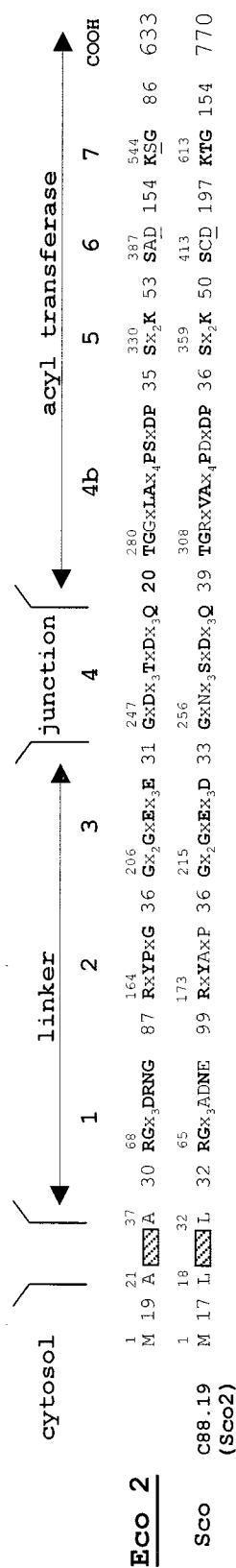
The cell shape apparatus of *E. coli* comprises at least six proteins encoded by genes located at the 14-min and the 71-min regions of the chromosome. Genes of the 14-min cluster code for the PBP fusion Eco2, RodA, and the free-standing PBP Eco5. RodA is an integral membrane protein (108). It belongs to a large protein family called SEDS for shape, elongation, division, and sporulation (100) or SFR for SpoVE, FtsW, and RodA (32). Eco2, RodA, and ribosomal activities are coordinated by a chain of relaying elements, one of which is regulated by the alarmone ppGpp (115, 191). Genes of the 71-min cluster code for MreB, MreC, and MreD (241). MreB and actin, a central component of the eukaryotic cytoskeleton, are very similar in three dimensions (229). MreB-like proteins are widely distributed among rod-shaped, filamentous and helical bacteria, suggesting that an MreB cytoskeleton is important to generate a nonspherical shape.

The cell septation apparatus of *E. coli*, known as the divisome, is a Lego-like interlocking of a set of at least 10 proteins that assemble into a septal ring at midcell. With Fts standing for temperature-sensitive filamentous phenotype, the PBP fusion Eco3 (or FtsI), FtsW, FtsQ, FtsA, FtsZ are encoded by genes of the 2-min *dcw* cluster, which also contains several lipid II synthetase-encoding genes (Fig. 5) (162, 237). FtsK, ZipA, FtsN, and YgbQ are encoded by genes occurring at different places on the chromosome. Like Eco3, FtsL, YgbQ, FtsQ, and FtsN are bitopic membrane proteins with a short cytoplasmic amino tail and a relatively large periplasmic domain (31, 36).

FtsL and YgbQ have a leucine-like zipper motif in the periplasmic domain. They belong to a family of proteins that exhibit a great propensity to form coiled-coil structures (31). FtsW is a homologue of RodA (28). By analogy with FtsW of *S. pneumoniae* (76), the *E. coli* FtsW probably comprises 10 transmembrane segments, with a large extracytoplasmic loop extending between segments 7 and 8. FtsA and the eukaryotic actin are members of the same ATPase domain protein superfamily (to which Hsp70 also belongs) (237). FtsZ is a GTPase homologue of tubulin, the building block of the eukaryotic cell division microtubules (97, 142, 143).

FtsZ arrives first at mid-cell; it serves as scaffold holding the other proteins of the divisome together, and it provides the driving force for cytokinesis (2). FtsA and ZipA localize to the septum independently but in an FtsZ-dependent manner. The other proteins then assemble in a sequential dependency order as follows: FtsK, FtsQ, FtsL, YgbQ, FtsW, Eco3, and FtsN. It

PBP fusions of class B, subclass B2



PBP fusions of class B, subclass B3

Eco 3	1 M 22 F 39 L 25 RGx3DRSG	71 RxYPxG 15 Gx2GxxEx3D	31 AxNx3SxDx3Q	229 TGE5xLAX4PSxDP	23 Sx2K 48 SSN 132 KTG	91 588
Mtu 2163 (Mtu3)	1 M 91 A 109 L 25 RGx3DRNN	99 RxYPxG 15 Gx2GxxEx3D	33 GxTx3TxDx3Q	27 TGE5xLAX4NTxDP	24 Sx2K 51 SSN 148 KTG	84 679
Mle 0908 (Mle3)	1 M 86 V 107 V 23 RGx3DRNN	99 RxYPxG 15 Gx2GxxEx3D	33 GxTx3TxDx3Q	27 TGE5xLAX4NTxDP	24 Sx2K 51 SSN 148 KTG	84 675
Cdi 101	1 R L 135 L 25 RGx3DNTG	126 RxYPxG 15 Gx2GxxEx3D	33 GxNx3TxDx3Q	27 TGE5xLAX4NTxDP	24 Sx2K 51 SSN 148 KTG	84 675
Sco 4A10.23 (Sco3)	1 M 58 M 74 V 30 RGx3DRNG	107 RxYPxG 15 Gx2GxxEx3D	32 GxDx3TxDx3Q	27 TGE5xLAX4PGxDP	23 Sx2K 51 SSN 149 KTG	82 654

PBP fusions of class B-like I

Mtu 0016	1 M 6 I 21 L 54	76 RxYPxP 18 Lx2AxxDx3N	21 GxNx3TxNx3Q	25 TGKxLAX4PSxDP	38 Sx2K 55 SCN 140 KTG	64 491
Mle 0018	1 M 6 I 21 L 54	76 RxYPxP 18 Lx2AxxDx3N	21 GxNx3TxNx3Q	26 TGKxLAX4PSxDP	38 Sx2K 55 SCN 140 KTG	64 492
Cdi 16	1 M 5 F 21 V 54	76 RxVTxP 18 Mx2NxNx3S	21 GxNx3TxNx3Q	26 TGKxLAX4PSxDP	38 Sx2K 55 SCN 140 KTG	64 492
Sco H69.17	1 M 6 I 21 I 54	76 RxYKxG 18 Lx2IxxDx3T	21 GxNx3TxNx3Q	22 TGKxLAX4PSxDP	39 Sx2K 55 SCN 139 KTG	61.....
Sco 6G9.32	1 M 6 A 21 V 54	76 RxYAXG 18 Lx2AxxDx3S	21 GxDx3TxHx3Q	22 TGKxLsx4PSxDP	39 Sx2K 54 SCN 139 KTG	66 490
Sco 2st10A7	1 M 4 A 21 I 55	77 RxYVxG 18 Lx2TxDx3T	21 AxSx3TxDx3Q	23 TGKxLAX4PSxDP	37 Sx2K 55 SCN 144 KTG	65 494
Sgr pbpa	1 M 6 I 21 I 54	76 RxWKxG 18 Lx2LxDx3T	21 GxNx3TxNx3Q	22 SGAxLAX4PSxDP	37 Sx2K 54 SCN 139 KTG	64 484

FIG. 12. Class B PBP fusions. Bar code, motifs 1 to 7. For protein identifiers, see Table 1.

is likely that class A PBP fusions are also components of the divisome (233). The class A PBP fusion Bsu1 (subclass A3) of *Bacillus subtilis* is localized at the division septum in vegetative cells (175).

The cell shape PBP fusion Eco2 of subclass B2 and the cell septation PBP fusion Eco3 of subclass B3 are produced in small amounts: a few tens and about 200 molecules per cell, respectively. Their acyltransferase modules are distantly related by a *P* value of 6×10^{-15} (identity, 29%). Loss of Eco2 results in a block of cell division, transformation of the *E. coli* cells into spherical bodies, and cell death. Loss of Eco3 causes a block of cell septation, formation of multigenomic filaments, and cell death. Loss of either Eco2 or Eco3 is fatal (211). PBP fusions of subclass B2, however, are dispensable in some gram-negative bacteria. *Neisseria meningitidis* possesses a single PBP fusion of class B that belongs to subclass B3 (255). This PBP fusion may fulfill functions comparable to those of Eco2 and Eco3 in *E. coli* (14). Consistently, the clusters formed by the linker and acyltransferase modules of the PBP fusions of subclass B3 are more populated than the clusters formed by the corresponding modules of the PBP fusions of subclass B2 (Fig. 10).

Both Eco2 and Eco3 are implicated in the exponential phase of growth of *E. coli* cells (19), conditions under which the peptidoglycan precursors are incorporated all over the lateral wall in a diffuse way (49). Newly formed (4→3) peptidoglycan is mixed with existing (4→3) peptidoglycan except at the time of cell septation. At this stage, peptidoglycan synthesis is strictly localized to the septum in an Eco3-dependent manner (27). In the course of the hierarchical assembly of the divisome, FtsZ, FtsQ, FtsL, and YgbQ are required for the septal localization of FtsW, and FtsW is essential for the subsequent recruitment of Eco3 (154).

Consistently, the linker module of Eco3 appears to be designed in such a way that the acyltransferase module is positioned, in an active form, within the divisome where it needs to be (148) (Fig. 13). The membrane anchor (243) and the segment upstream from motif 1 of the linker module (148) have the information ensuring that Eco3 localizes at the cell septation site. Motifs 1, 2, and 3 and other peptide segments which form the core of the linker module have the information ensuring that Eco3 folds correctly and that the acyltransferase catalytic center adopts the active conformation. The Glu206-Val217 peptide segment at the surface of the linker module has the information ensuring that Eco3 fulfills its cell septation activity within the fully complemented divisome by interacting with other components of the morphogenetic apparatus.

In *in vitro* assays, Eco3, which lacks glycosyltransferase activity, is inert on lipid II. As observed with the PBP fusion Eco1b of class A, Eco3 is inert on *N*-acyl-D-alanyl-D-alanine-terminated peptides, but it catalyzes the hydrolysis and aminolysis of thiolester analogues of the peptides (1), indicating that *in vivo*, Eco3 identifies *N*-acyl-D-alanyl-D-alanine sequences as carbonyl donors. As derived from studies carried out both *in vivo* and with ether-permeabilized cells (179), the acceptor of the Eco3-catalyzed transfer reaction could be the lateral amino group at position 3 of tripeptide-derived precursors, resulting in the formation of (4→3)-linked tetrapeptide-tripeptide dimers. The required stem tripeptides could be brought into the periplasm in the form of incomplete (disac-

charide tripeptide) lipid II molecules lacking the D-alanyl-D-alanine dipeptide normally incorporated by the ligase MurF.

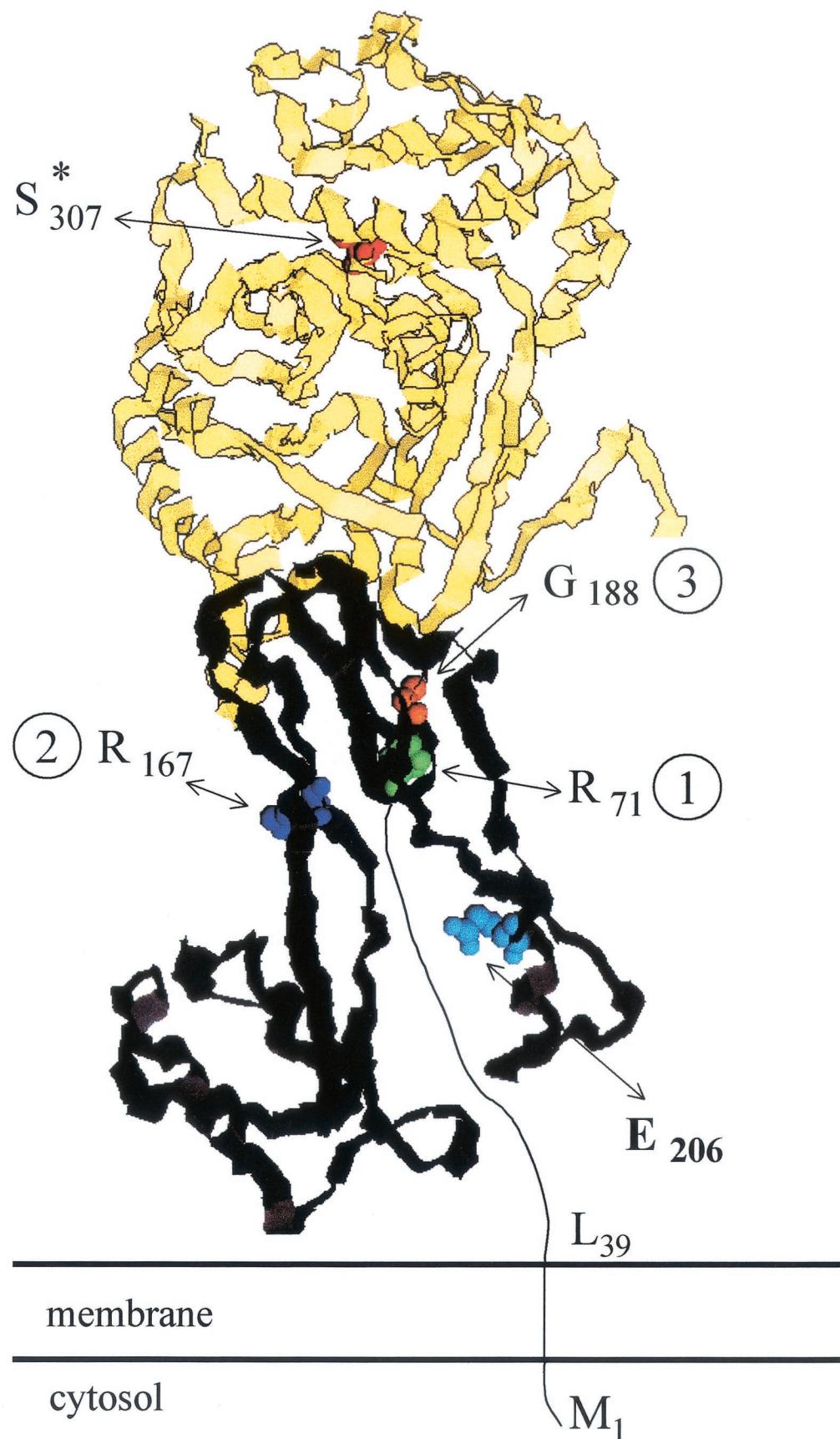
E. coli produces, in the cytosol, penicillin-resistant peptidases that hydrolyze the bond between *meso*-diaminopimelic acid (L-center) and D-alanine at positions 3 and 4 of the stem peptides. An L,D-endopeptidase (85) and an L,D-carboxypeptidase I (156) act on the nucleotides UDP-*N*-acetylmuramoyl-(D-alanyl-D-alanine-terminated) pentapeptide and UDP-*N*-acetylmuramoyl-(D-alanine-terminated) tetrapeptide, respectively. The L,D carboxypeptidase LdcA (221, 228) is made without signal peptide. Loss of LdcA causes cell lysis, and the effect is restricted to the onset of the stationary phase. LdcA does not belong to the SxxK peptidase family. The required stem tripeptides can also be produced, in the periplasm, by the action of the L,D-carboxypeptidase II (17). LdcII acts on stem tetrapeptides of nascent peptidoglycan at the time of cell division. It may be present in nondividing cells, but then its activity is masked. LdcII has not been characterized biochemically.

Eco2 is essential to multiplying rod-shaped *E. coli* cells. It may be also important in the stationary phase (23). In rounded *E. coli* cells, in which Eco2 is impaired, the incorporation of the peptidoglycan precursors is a zonal process (49). There is no mixing of new peptidoglycan and old peptidoglycan, and the wall polymer cannot undergo remodeling. *In vitro* assays that would help identify the reaction that the acyltransferase module of Eco2 performs *in vivo* have not been developed. Eco2 binds benzylpenicillin and ampicillin with high affinity, suggesting that the acyltransferase module is targeted to *N*-acyl-D-alanyl-D-alanine sequences. However, Eco2 is also very susceptible to the nonclassical penam amdinocillin, the carbapenem thienamycin, and the oxapenem clavulanic acid (213) (Fig. 8).

The linker modules of Eco2 and Eco3 have the same conserved motifs 1, 2, and 3. They probably adopt the same fold. As a corollary, peptide segments other than the conserved motifs must serve as recognition sites specifying the morphogenetic apparatus to which Eco2 and Eco3 attach. Cell division and viability in the absence of Eco2 but in the presence of Eco3 are restored by increasing the pool of ppGpp or the level of FtsQAZ (115, 238). Hence Eco2, but not Eco3, is dispensable in particular genetic backgrounds. It is likely that the morphogenetic apparatus is not composed of fixed structures and proteins can belong to several morphogenetic apparatuses simultaneously or at different times.

Free-Standing PBPs: Auxiliary Cell Cycle Proteins

Free-standing PBPs are autonomous folding and catalysis entities. They utilize water as the acceptor nucleophile for the deacylation of the serine ester-linked *N*-acyl-D-alanyl enzyme intermediate. They are implicated, one way or another, in cell morphogenesis. *E. coli* produces multiple free-standing PBPs (Fig. 14). Eco5, Eco6, and Eco6b (13) have carboxy-terminal membrane-anchoring domains, and the bulks of the polypeptide chains are exposed in the periplasm. These D-alanyl-D-alanine-cleaving carboxypeptidases, together with other factors, control the balance between different peptidoglycan precursors and help determine whether *E. coli* cells will elongate or divide (19). The similarities between the pair Eco5-Eco6 (*P*, 10^{-141} ; identity, 60%) and the pair Eco5-Eco6b (*P*, 10^{-104} ; identity, 47%) are great, indicating that Eco5, Eco6,



and Eco6b arose by gene duplication with a high level of sequence conservation. Eco5 and Eco6, however, are not functionally redundant. Loss of Eco5 but not of Eco6 suppresses the block in cell division caused by a single base change in the gene encoding FtsK, a protein that performs a septation function and a chromosome partition function (18). Overproduction of Eco5 (147) but not of Eco6 (230) results in growth of wild-type *E. coli* as spherical cells.

The free-standing PBPs Eco4 (129) and Eco7 (198) hydrolyze D-alanyl-(D)-meso-diaminopimelic acid interpeptide linkages made by transpeptidation. Eco7 is peculiar. It is released from whole cells by osmotic shock. It hydrolyzes the peptidoglycan saccus but is inert on isolated disaccharide peptide dimers. It is inactivated by low concentrations of penems with an L,D-configured scissile bond (Fig. 8), and its inactivation causes lysis of nonmultiplying cells. Eco7, Eco5, Eco6, and Eco6b belong to the S11 family in the database MEROPS (190). Eco4 belongs to the S13 family. One may note here that *E. coli* also produces a penicillin-resistant D-alanyl-(D)-meso-diaminopimelic acid-cleaving peptidase, MepA (120). MepA is secreted in the periplasm. Surprisingly, a fivefold superproduction does not modify the overall extent of peptidoglycan cross-linking. MepA does not belong to the SxxK peptidase family.

E. coli produces two additional chromosome-encoded free-standing SxxK acyltransferases, AmpC and AmpH (99). They are similar in sequence. AmpC is a β-lactamase (see Group II SxxK Acyltransferases below) or a PBP, depending on the structure of the β-lactam antibiotic with which it reacts (74). AmpH is a high-affinity PBP for benzylpenicillin and cephalosporin C (99).

The free-standing PBPs are nonessential auxiliary cell cycle proteins. A triple deletion of Eco4, Eco5, and Eco6 has no obvious effects apart from a slightly longer generation time and slightly altered cell morphology (59). The genes encoding Eco4, Eco5, Eco6, Eco6b, Eco7, AmpC, and AmpH have been deleted in every possible combination; all the deletions give rise to viable *E. coli* cells. Some of them, however, cause significant morphological aberrations (48, 164, 251).

E. coli PBP Fusions as Targets for β-Lactam Antibiotics

To kill bacteria, a β-lactam antibiotic must inactivate the acyltransferase module of one of several PBP fusions at therapeutically achievable concentrations in time periods shorter than the generation time of the bacterium. Currently, 50% and 90% inhibitory concentration (IC_{50} and IC_{90}) values, i.e., the concentrations (in micrograms of antibiotic per milliliter) at which a β-lactam compound inhibits the PBPs by 50% or 90%, respectively, after a given time of incubation with isolated membranes, are used to estimate the inactivating efficiency of the drug. IC values are not intrinsic parameters of the inter-

action. With K (molarity) denoting the dissociation constant of the protein-β-lactam Michaelis complex and k_{+2} (per second) denoting the first-order rate constant of protein acylation, the second-order rate constant k_{+2}/K expresses the inactivating efficiency of a β-lactam antibiotic if the value of the rate of breakdown of the acyl enzyme is small.

Under the experimental conditions defined in reference 82, k_{+2}/K is equal to $-\ln X/([\beta\text{-lactam}] \cdot t)$, where X is the percentage of the target protein left intact, $[\beta\text{-lactam}]$ is the molar drug concentration, and t is the incubation time in seconds. Benzylpenicillin, at $1 \mu\text{g ml}^{-1}$ (i.e., $3.0 \times 10^{-6} \text{ M}$) concentration, inactivates a target protein by 50% in 600 s if the k_{+2}/K value is $\approx 400 \text{ M}^{-1} \text{ s}^{-1}$. At the same concentration and in the same time period, benzylpenicillin inactivates another target protein by 95% if the k_{+2}/K value is $1,600 \text{ M}^{-1} \text{ s}^{-1}$.

Table 2 gives the k_{+2}/K values of the interactions between β-lactam antibiotics and the isolated free-standing PBP Eco4* (194), the His-tagged (Met46-Asn844) PBP fusion Eco1b* (222), the OmpA signal peptide-transported (Gly57-Val577) PBP fusion Eco3* (1), and the membrane-bound PBPs of *E. coli*, as calculated from published IC_{50} values (41). Binding of a β-lactam antibiotic to isolated membranes is a competition between multiple target proteins with various affinities for the drug. Traces of a β-lactamase can interfere with the assays. Hence, the k_{+2}/K values of the isolated Eco4*, Eco1b*, and Eco3* are considerably larger than those derived from the corresponding IC_{50} values, yet conversion of the IC_{50} values into k_{+2}/K values allows the β-lactams to be roughly classified in the order of their acylating potencies.

E. coli is killed via cell spheroplasting and lysis as a result of the selective inactivation of both Eco1a and Eco1b by cefaloridine and cefsulodin; via transformation of the cells into round bodies as a result of the selective inactivation of Eco2 by amdinocillin; via cell filamentation as a result of the selective inactivation of Eco3 by mezlocillin, cefaperazone, cefotaxime, cefuroxime, cephalothin, and aztreonam (data not shown); or via different combinations of these morphological alterations by ampicillin, benzylpenicillin, carbenicillin, and cefoxitin.

The membrane-bound free-standing PBPs Eco4, Eco5, and Eco6 are, in general, less susceptible to the β-lactam antibiotics than the acyltransferase modules of the membrane-bound PBP fusions. However, Eco4 and the PBP fusions Eco1a, Eco1b, Eco2, and Eco3 have comparable susceptibilities to ampicillin, benzylpenicillin, and carbenicillin. Eco5, Eco6, and the PBP fusions Eco1a and Eco1b have comparable susceptibilities to cefoxitin.

From the foregoing, it follows that the essential class A and B PBP fusions present in a bacterial cell each has its own specificity profile for β-lactam antibiotics. Most often, they perform different functions and belong to different subclasses. As shown in the ensuing section, variants of a PBP fusion can

FIG. 13. Schematic structure of the membrane-bound SxxK subclass B3 PBP fusion protein Eco3 of *E. coli*. Spatial disposition along the polypeptide chain of the essential S*307 of the SxxK motif of the acyltransferase module (see Fig. 7) and of motifs 1, 2, and 3 (identified by the residue at the amino side of the sequences) and segment E206 to V217 of the linker module. The acyltransferase module is in yellow, with S*307 of the SxxK motif in red. The linker module is in black with motif 1 (R71 to G79) in green, motif 2 (R167 to G172) in dark blue, motif 3 (G188 to D197) in orange, and segment E206 to V217 in light blue. Motifs 1, 2, and 3 and other peptide segments form the core of the linker module in interaction with a noncatalytic groove of the acyltransferase module. The peptide segment M1 to R71 is of unknown structure. Adapted from reference 148. Illustration courtesy of Robert Brasseur, Faculté Universitaire des Sciences Agronomiques, Gembloux.

arise. The variants belong to the same subclass and presumably perform the same function as the wild-type PBP fusion, but their susceptibilities to the drugs are changed.

Mosaic PBP Fusions

Streptococcus pneumoniae is of known genome sequence (105). It is a paradigm of genetic plasticity and adaptation to the environment via natural transformation (37). DNA sequences that are likely to be functional in a bacterium donor are introduced by homology-dependent recombination in a bacterium acceptor. Factors regulating genetic competence are involved in this process.

S. pneumoniae strains are normally very susceptible to benzylpenicillin (MIC, 0.02 µg of antibiotic ml⁻¹). Determinants conferring reduced susceptibility to β-lactam antibiotics have evolved in commensal *Streptococcus* spp., presumably by the accumulation of point mutations in genes that code for PBP fusions of classes A and B. The shuffling and capture of DNA sequences from strains that have a low susceptibility to the drugs give rise to *S. pneumoniae* isolates that carry genes which code for mosaic PBP fusion variants of classes A and/or B having a decreased affinity for one or several β-lactam antibiotics (56, 89, 90, 257). In the laboratory, several independent mutations are necessary for a penicillin-susceptible PBP fusion to be converted into a mosaic PBP fusion.

In nature, *Streptococcus oralis* and *Streptococcus mitis* are likely to act as DNA donors (6). As the result of one transformation event with chromosomal DNA from an *S. mitis* strain having a low susceptibility to penicillin, *S. pneumoniae* produces an assortment of mosaic PBP fusions that substitute for the wild-type PBP fusions Spn1a, Spn2a, and Spn1b of subclasses A3, A4, and A5 and the PBP fusions Spn2x and Spn2b of subclasses B4 and B5 (91). Mosaic and wild-type PBP fusions with different susceptibilities to β-lactam antibiotics but of the same subclass differ in amino acid residues by up to 15%.

S. pneumoniae strains producing mosaic PBP fusions often manufacture a peptidoglycan which is enriched in interpeptide cross-bridges (64, 65). The inactivation of *murMN* in *S. pneumoniae* Pen6 (MIC, 6 µg of benzylpenicillin ml⁻¹), with loss of the enzymes of the pathway to branched peptidoglycan precursors (see the section on lipid II precursors), restores susceptibility to the drug (MIC, 0.03 µg of benzylpenicillin ml⁻¹). *N. gonorrhoeae* strains that have decreased susceptibility to penicillin have also emerged (212). The mosaic PBP fusions that they produce arise by amino acid substitution, insertion, and exchange of regions of the penicillin-susceptible acyltransferase modules with the homologous regions of resistant PBPs of closely related species.

GROUP II SxxK ACYLTRANSFERASES

The SxxK acyltransferases of group II are indirectly or not implicated in wall peptidoglycan synthesis and metabolism. Many of them help, in one way or another, the SxxK PBP fusions of group I escape penicillin action. Others have very diverse functions. The SxxK acyltransferases of group II occur as free-standing polypeptides and as modules of protein fusions. They illustrate exemplarily the concept of protein superfamily: polypeptide chains adopting the same fold can perform

1. Code : SxxK, SxN, KTG

Eco 4	1 M 60	62 <u>Sx₂K</u> 240	306 SDN 108	417 KTG	COOH 57 477
5	1 M 71	73 <u>Sx₂K</u> 62	139 SGN 100	242 KTG	158 403
6	1 M 64	66 <u>Sx₂K</u> 62	132 SGN 100	235 KTG	162 400
6b	1 M 63	65 <u>Sx₂K</u> 62	131 SGN 100	234 KTG	153 390
7	1 M 70	72 <u>Sx₂K</u> 51	127 SEN 104	234 KTG	76 313

Mtu 3627	1 M 112	114 <u>Sx₂K</u> 177	295 SDN 110	408 KTG	50 461
Mle 0211	1 M 112	114 <u>Sx₂K</u> 177	295 SDN 110	408 KTG	50 461

Mtu 3330	1 M 119	121 <u>Sx₂K</u> 51	176 SGN 103	282 KTG	120 405
Mle 0691	1 M 129	131 <u>Sx₂K</u> 51	186 SGN 103	292 KTG	117 411
Mtu 2911	1 M 67	69 <u>Sx₂K</u> 51	124 SGN 103	230 KTG	58 291

2. Code : SxxK, YxN, HxG

Mtu 0907	1 L 118	120 <u>Sx₂K</u> 87	211 YTN 147	361 HSG	168 532
Mtu 1730	1 L 110	112 <u>Sx₂K</u> 88	204 YNN 146	353 HGG	161 517
Mtu 1922	1 M 83	85 <u>Sx₂K</u> 104	193 YSN 131	327 HAG	41 371

3. Incomplete code : SxxK, YxH

Mtu 1367	1 M 33	35 <u>Sx₂K</u> 108	147 YSH	227	377
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4. Code : Class A β - lactamase

Mtu 2068	1 M 82	84 <u>Sx₂K</u> 54	142 / \ 250 SDG 106	182 Ex ₂ LN	54 307
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FIG. 14. Occurrence of catalytic center-defining motifs along the sequences of free-standing SxxK polypeptides of *E. coli*, *M. tuberculosis*, and *M. leprae*. Polypeptides of group 1 bear the bar codes SxxK, SxN, and KTG. They are auxiliary cell cycle PBPs in *E. coli*. Inserts are underlined. The *M. tuberculosis* polypeptides of groups 2 and 3 have no equivalent in *E. coli*. Their bar codes are modified or incomplete. It is likely that they are not implicated in wall peptidoglycan metabolism. The *M. tuberculosis* β-lactamase of class A (group 4) has a class-specific Ex₂LN motif. *E. coli* can produce two β-lactamases, one of which (plasmid coded) is of class A (Fig. 15).

extremely various functions. Often, the SxN motif and/or the KTG motif is modified. The SxxK motif, however, is strictly conserved (Fig. 15).

Streptomyces sp. strain K15 produces a free-standing PBP which acts as a D,D-transpeptidase. SxC substitutes for the SxN motif (171). The mature enzyme is membrane associated but is secreted when overproduced. In aqueous media which contain 55.5 M H₂O, proper amino compounds, at millimolar concentrations, compete successfully with water as acceptors of the transfer reaction. With the carbonyl donor-amino acceptor

TABLE 2. Second-order rate constant k_{+2}/K values of acylation by β -lactam antibiotics of the membrane-bound PBPs of *E. coli* and *M. tuberculosis* RvH37Ra, the water-soluble Met46-Asn844 Eco1b* and Gly57-Val577 Eco3*, and purified Eco4*

β -Lactam	k_{+2}/K ($M^{-1} s^{-1}$)													
	<i>E. coli</i> PBPs						<i>M. tuberculosis</i> PBPs							
	1a	1b	1b*	2	3	3*	4	4*	5	6	94 kDa	82 kDa	52 kDa	37 kDa
Cephaloridine	2,000	200	350	10	65		30		<2	<2	600	200	400	<5
Cefsulodin	1,300	130		<2	<2		<2		<2	<2				
Mecillinam	<2	<2		<2,000	<2		<2		<2	<2				
Mezlocillin	400	80		700	27,000	290,000	<30		<30	<30				
Cefoperazone	1,500	500	1,800	830	15,000	80,000	<15		<15	<15	30	20	200	<5
Cefotaxime	10,500	800	2,600	100	<10,500	80,000	20		<10	<10				
Cefuroxime	4,000	300	700	40	6,000	66,000	3		<2	3				
Cephalothin	>1,800	30		10	440		7		4	4	80	30	300	<5
Ampicillin	300	100	130	500	450	5,000	200		3	40	2,000	300	250	<5
Benzylpenicillin	800	100	600	480	430	4,000	385	>100,000	16	20	≈600	≈600	≈600	≈30
Carbenicillin	200	100	120	100	200	1,500	120		3	3				
Cefoxitin	5,000	150	900	<2	90	1,000	75		800	550	1,200	600	100	800
Imipenem											6,000	2,000	400	250
Clavulanate											230	90	10	10
Sulbactam											330	150	<3	<3

system bisacetyl-L-lysyl-D-alanyl-D-alanine/glycylglycine, the product of the reaction is almost exclusively bisacetyl-L-lysyl-D-alanyl-glycyl-glycine (140, 165). *Streptomyces* sp. strain R61 and *Actinomadura* sp. strain R39 secrete free-standing PBPs which act mainly as D,D-carboxypeptidases. In the R61 PBP, YxN substitutes for the SxN motif and HTG substitutes for the KTG motif (57). In the R39 PBP, motifs SxxK, SxN, and KTG are conserved, but a large insert occurs between the SxxK motif and the SxN motif (87).

The primary response of *Streptomyces* spp. and other soil bacteria to exposure to β -lactam antibiotics might have been to develop a protective mechanism through the secretion of free-standing PBPs with increasing affinities for the drugs (122). Thus, the k_{+2}/K values of the interactions with benzylpenicillin are $135 M^{-1} s^{-1}$ for the K15 enzyme, $18,000 M^{-1} s^{-1}$ for the R61 enzyme, and $300,000 M^{-1} s^{-1}$ for the R39 enzyme.

The conversion of free-standing PBPs into SxxK β -lactamases gave rise to a defense mechanism of great efficiency (125, 126). The SxxK β -lactamases have a very low activity on N-acyl-D-alanyl-D-alanine peptides because the enzyme acylation step is severely rate limiting (195). They react with and rupture the β -lactam amide bond and produce serine ester-linked acyl (penicilloyl, cephalosporoyl, ...) enzyme intermediates that are hydrolytically labile. On good β -lactam substrates, the catalytic centers of the serine β -lactamases can turn over 1,000 times or more per second. In β -lactamases of class A, the SxxK, SxN, and KTG motifs are conserved, but an additional motif, ExxLN, occurs in the catalytic center. In β -lactamases of classes C and D, YxN and SxV substitute for the SxN motif, respectively.

Penicillin sensory transducers are components of regulatory pathways leading to the inducible synthesis of penicillin resistance determinants (96, 254). BlaR proteins are implicated in β -lactamase synthesis in *Bacillus licheniformis* and *Staphylococcus aureus*. The MecR protein is implicated in synthesis of the penicillin-resistant SxxK protein fusion Sau2a in *S. aureus* (see next section). The tripartite BlaR and MecR proteins comprise a signal receptor which is at the carboxy end of the polypeptide

chains and is exposed on the outer face of the plasma membrane; a four α -helix bundle signal transmitter which is embedded in the membrane; and a cytosolic signal emitter which possesses the HisGluLeuTyrHis consensus of a Zn^{2+} -dependent peptidase.

The Met346-Arg601 penicillin receptor of the *B. licheniformis* BlaR is related to the Oxa2 β -lactamase by a *P* value of 6×10^{-36} (identity, 36%). As an independent entity, the BlaR receptor lacks detectable peptidase and β -lactamase activities and behaves as a high-affinity PBP (96), yet signal reception by the full-size BlaR leading to transcription of the β -lactamase-encoding gene does not involve penicilloylation of the serine residue of the SxxK motif of the receptor. In *S. aureus*, unblocking transcription of the Sau2a-encoding gene is the result of site-specific proteolytic cleavage of the zinc peptidase of the transmitter of MecR, which is activated, and of the repressor, which is inactivated (254).

There are SxxK acyltransferases which are not targeted to peptide bonds extending between two D-alanine residues in an α -position to a carboxylate. *Bacillus cereus* secretes a peptidase ADP which hydrolyzes the D-phenylalanyl-D-phenylalanyl bond in an endopeptidase (10). The peptidase has some β -lactamase activity. Its bar code is similar to that of the *Streptomyces* sp. strain R61 D,D-carboxypeptidase-PBP. The two proteins are related by a *P* value of 10^{-48} (identity, 36%). *Ochrobaculum anthropi* produces intracellularly a D-aminopeptidase, DAP, which acts on D-alanine amide and peptides with a D-alanine residue at the amino end (11, 26). The carbonyl side only of the scissile bond is borne by a D-configured carbon atom. β -Lactam compounds and the tripeptide bisacetyl-L-lysyl-D-alanyl-D-alanine behave as competitive inhibitors.

The catalytic module of the peptidase DAP and the free-standing D,D-carboxypeptidase-PBP of *Streptomyces* sp. strain R61 are distantly related by a *P* value of 3×10^{-18} (identity, 26%), yet the two polypeptides adopt the same fold (26). The noncatalytic carboxy-terminal module of the peptidase DAP comprises two eight-stranded β -barrels. A loop protruding from the last β -barrel interacts with the catalytic module. This

Free-standing PBPs		motifs	1	2	3		
<i>Streptomyces</i> K15 *	M	62	<u>64</u> Sx₂K	57	<u>125</u> SG<u>C</u>	114	242 KTG
<i>Streptomyces</i> R61 *	M	91	<u>93</u> Sx₂K	93	<u>190</u> YS<u>N</u>	136	329 HTG
<i>Actinomadura</i> R39	M	96	<u>98</u> Sx₂K	<u>245</u>	<u>347</u> SSN	109	458 KTG
β-lactamases					164 <u>Ex₂LN</u>		
Class A <i>E. coli</i> TEM *	M	66	<u>68</u> Sx₂K	56	<u>128</u> SD<u>N</u>	101	232 K<u>S</u>G
Class C <i>E. cloacae</i> P99 *	M	82	<u>84</u> Sx₂K	82	<u>170</u> Y<u>A</u>N	162	335 KTG
Class D <i>P. aeruginosa</i> OXA 10*	M	65	<u>67</u> Sx₂K	44	<u>115</u> SA<u>V</u>	98	205 KTG
<i>B. licheniformis</i> penicillin sensory transducer BLAR	M	<u>400</u>	<u>402</u> Sx₂K	44 70	<u>450</u> ST<u>T</u> 476 YGN	87 60	539 KTG
<i>B. cereus</i> endopeptidase ADP	M	102	<u>104</u> Sx₂K	94	<u>201</u> Y<u>S</u>N	139	342 H<u>G</u>G
D-Phe-D-Phe-D-Phe-D-Phe							
<i>O. anthropi</i> D-aminopeptidase DAP *	M	60	<u>62</u> Sx₂K	87	<u>153</u> Y<u>C</u>N	131	287 H<u>G</u>G
D-alanyl-amide							

FIG. 15. Acyltransferases of the SxxK superfamily of diverse functions not related to wall peptidoglycan metabolism. The proteins are free-standing polypeptides except BlaR of *B. licheniformis* and DAP of *O. anthropi*, which are protein fusions. The SxxK acyltransferase module forms the carboxy-terminal domain of BlaR and the amino-terminal domain of DAP. The three-dimensional structures of the proteins marked with an asterisk are known. Motif 1, SxxK, is invariant. Motifs 2 and/or 3 are modified except in PBP R39. Amino acid changes and inserts are underlined. Motif 2 of BlaR is ambiguous.

feature is somewhat reminiscent of the activation of the *Streptomyces* sp. strain R61 D,D-carboxypeptidase-PBP by elimination of the 26-residue carboxy-terminal segment of the protein precursor, suggesting prosegment occlusion of the catalytic center (62).

Amino acid sequences change more rapidly on the evolutionary time scale than the three-dimensional structures. The *Streptomyces* sp. strain K15 D,D-transpeptidase-PBP (68), the *Streptomyces* sp. strain R61 D,D-carboxypeptidase-PBP (124), the TEM β-lactamase of class A (5, 113, 217), the P99 β-lactamase of class C (168), the Oxa-10 β-lactamase of class D (170), the acyltransferase modules of the D-aminopeptidase DAP (26), and the PBP fusion Spn2x of subclass B4 (173) all adopt the same overall fold, indicating a common ancestral origin. The *Streptomyces* sp. strain K15 D,D-transpeptidase-PBP and the β-lactamases of class A belong to the MEROPS S11 family (190). The *Streptomyces* sp. strain R61 D,D-carboxypeptidase-PBP, β-lactamases of class C, the *B. cereus* peptidase ADP, and the catalytic module of the *O. anthropi* D-aminopep-

tidase belong to the S12 family. The *Actinomadura* sp. strain R39 D,D-carboxypeptidase-PBP belongs to the S13 family.

GROUP III SxxK ACYLTRANSFERASES: PENICILLIN-RESISTANT PROTEIN FUSIONS

The enormous variations in the catalytic properties of the SxxK acyltransferases of groups I and II are the result of alterations of the polypeptide chains, sometimes with a great divergence in the amino acid sequences but always with conservation of the polypeptide fold and positioning of secondary structures even well away from the catalytic centers. In light of this versatility, acyltransferases might have gone as far as a change from D,D specificity to L,D specificity. SxxK L,D-acyltransferases catalyzing reactions II and III shown in Fig. 6 are expected to resist most β-lactam antibiotics. The roles that they may play in (3→3) peptidoglycan assembly are discussed in the ensuing sections.

Class B Pen^r Protein Fusions of Gram-Positive Bacteria

Enterococci and staphylococci produce, in addition to their normal sets of PBP fusions of classes A and B, one protein fusion which has the bar code of class B PBP fusions (Fig. 16) but has a very low affinity for the β -lactam antibiotics (67, 98). The class B penicillin-resistant (Pen^r) protein fusions which have been tested are inert on both D-alanyl-D-alanine-terminated peptides and thioester analogues. The k_{+2}/K values of their interactions with β -lactam antibiotics (from $\approx 1 \text{ M}^{-1} \text{ s}^{-1}$ to $\approx 10 \text{ M}^{-1} \text{ s}^{-1}$) are only about 20-fold greater than that observed with bovine serum albumin.

The linker and acyltransferase modules of the Pen^r protein fusions Efam5 of *Enterococcus faecium* (259), Ehi5 and Ehi3^r (plasmid borne) of *Enterococcus hirae* (61, 177), Efas5 of *Enterococcus faecalis* (206), Sau2a or Sau2' of *Staphylococcus aureus* (210), and Ssc2 of *Staphylococcus sciuri* (247, 248) diverged in concert. They fall into a particular subclass, B1 (Fig. 10), to which the protein fusion Bsu3 of *Bacillus subtilis* (160) also belongs (Fig. 10). Orthologues of Efam5 are also present in *Listeria monocytogenes*, *Listeria innocua*, and *Clostridium acetobutylicum*. Their acyltransferase modules are related to that of Efam5 by P values ranging from 10^{-87} to 10^{-66} . The linker modules of Pen^r protein fusions of subclass B1 invariably contain an insert 120 to 130 amino acid residues long upstream from motif 1 (Fig. 16). The inserts of Ehi5 (157) and Sau2a (246) are essential. They probably have their own fold.

Pen^r protein fusions of subclass B1 are penicillin resistance determinants in exponential-phase cultures of streptococci and staphylococci. They allow the strains that produce them to multiply under conditions under which the acyltransferase modules of class A and B PBP fusions are inactivated by penicillin. The level of resistance, however, is almost always below the level that one would expect on the basis of the very low affinity of the Pen^r protein fusion. In addition, this alternative mode of bacterial growth in the presence of penicillin is of decreased efficiency. When Efam5-producing *E. faecalis* cells are collected from penicillin-free medium and inoculated into penicillin-containing medium, 2 to 3 h elapse before the cells start to multiply again. They continue to multiply in the presence of penicillin, but the generation time is twofold greater than that of the control cells in the absence of penicillin (207).

The Pen^r protein fusions Efam5 of *E. faecium* and Sau2a of *S. aureus* have attracted much attention. The Efam5-encoding gene is probably intrinsic to the *E. faecium* species. All known isolates are Efam5 producers, including those strains which are still susceptible to therapeutically achievable concentrations of penicillin and in which Efam5 is produced at a low basal level. The Efam5-encoding gene belongs to an operon which contains a gene that codes for a protein of the SFR family, indicating that Efam5 is a cell cycle protein. The operon also contains a gene that codes for a Psr protein first identified as a possible repressor of transcription. Recent developments argue against the possibility that Psr plays such a role in *E. faecium* (196) and *E. faecalis* (58).

The evolutionary precursor of the Sau2a-encoding gene, *mecA*, of *S. aureus* could be a homologue present in *S. sciuri*, a species found in rodents and primitive mammals (247, 248). Stepwise exposure of a penicillin-susceptible *S. sciuri* strain in which *mecA* is silent (MIC, 4 μg of antibiotic ml^{-1}) to increas-

ing concentrations of methicillin gives rise to a methicillin-resistant mutant (MIC, 200 μg of antibiotic ml^{-1}) which, as a result of a point mutation introduced in the -10 consensus of the *mecA* promoter, produces a protein which reacts with an anti-Sau2a monoclonal antibody. Transduction of *S. sciuri* *mecA* into a methicillin-susceptible *S. aureus* strain (MIC, 4 μg of antibiotic ml^{-1}) results in increased resistance to methicillin, albeit at a lower level (MIC, 12 to 50 μg of antibiotic ml^{-1}) than that of the donor strain. In methicillin-resistant *S. aureus* strains, *mecA* is carried by a mobile element called the staphylococcal cassette chromosome *SCCmec* (110). The *mecA* sequences appear to be highly conserved. Only two nucleotide positions vary in the genes of methicillin-resistant *S. aureus* strains and *S. sciuri*.

The Pen^r protein fusions Sau2a of *S. aureus* and Efam5 of *E. faecium* belong to the same subclass, B1, indicating similarity in function. Their biochemistry audit is far from complete. The mechanisms through which they contribute to the penicillin resistance of cultures in the exponential phase are undetermined. A definite answer will have to take the following observations into account.

E. faecium cells in the exponential phase manufacture a (3→3) peptidoglycan in various proportions of total peptidoglycan (Fig. 1 and 2). As proposed in references 145 and 205, the (3→3) L-lysyl-N^ε-(D-isoasparaginyl)-L-lysine cross-bridges (Fig. 2) could be made through the sequential actions of a Pen^r N-acyl-D-alanyl-D-alanine carboxypeptidase catalyzing reaction IIIa and a Pen^r L,D-transpeptidase catalyzing reaction III (Fig. 6).

The presumed Pen^r D,D-carboxypeptidase could be a Zn²⁺-dependent peptidase of the EntVanY family (8, 9, 245). VanY is produced by vancomycin-resistant enterococci. VanY homologues occur in *Clostridium* and *Bacillus* spp. but not in *E. coli* or *Mycobacterium tuberculosis*. *Streptomyces albus* G secretes a Zn²⁺-dependent N-acyl-D-alanyl-D-alanine carboxypeptidase functionally equivalent to EntVanY. The *Streptomyces* enzyme is of known structure (52, 83). His154, Asp161, and His197 are the zinc ligands. Amino acid sequence alignments reveal an equivalent zinc site in EntVanY (137, 151).

The presumed Pen^r L,D-transpeptidase could be the peptidase responsible for the D-alanine exchange reactions (of the type bisacetyl-L-lysyl-D-alanine + D-[¹⁴C]alanine ⇌ bisacetyl-L-lysyl-[¹⁴C]-D-alanine + D-alanine) which were identified several decades ago in *E. hirae* (called *Streptococcus faecalis* ATCC 9790 in 1974) (40), *Gaffkya homari* (93–95), *Bacillus megaterium* (44), and, recently, *E. faecium* (145). The membrane-associated L,D-peptidase is present at a very low level in wild-type *E. hirae* cells (40). It assembles properly structured carbonyl donor and amino acceptor peptides into (3→3) N^ε-L-lysyl-N^ε-(D-isoasparaginyl)-L-lysyl-linked dimers. It resists benzylpenicillin and is moderately susceptible to cephaloglycine. Almost certainly, it performs covalent catalysis via the formation of an acyl enzyme intermediate. Whether it belongs to the SxxK acyltransferase superfamily or not remains to be established.

The enterococcal Pen^r D,D-carboxypeptidase and Pen^r L,D-transpeptidase, working cooperatively with a glycosyltransferase, could be the basic molecular machine required for the formation of polymeric (3→3) peptidoglycan from lipid II precursor molecules. The implication of the subclass B1 Pen^r

Pen^r protein fusions of class B, subclass B1

	cytosol	linker	junction	acyl transferase
Efam5	1 9 1 G 143 R _x ₂ S ₃ DRNG 81 RxYPxG 30 G _x ₂ G _x ₃ D 31	258 30 ₀ G _x ₂ G _x ₃ TxD _x ₃ Q 31	335 G _x ₂ D _x ₃ TxD _x ₃ Q 20	368 TGDxLAX _x ₄ PSxDP 34
Efass5	1 15 31 L G 143 R _x ₂ S ₃ DRNG 81 RxYPxG 29 G _x ₂ S _x ₃ D 31	265 30 ₀ G _x ₂ TxD _x ₃ Q 20	341 G _x ₂ TxD _x ₃ Q 20	374 KGDxLAX _x ₄ PSxDP 34
Sau2a	1 6 20 R _x ₂ S ₃ DRNN 81 RxYPxE 32 G _x ₂ G _x ₃ D 32	241 27 ₉ G _x ₂ G _x ₃ D 32	321 G _x ₂ TxD _x ₃ Q 20	354 TGE _x LAX _x ₄ PSxDP 34
Bsu3	1 6 20 R _x ₂ S ₃ DRNG 85 RxYPxG 31 G _x ₂ G _x ₃ D 27	253 29 ₁ G _x ₂ G _x ₃ D 27	328 G _x ₂ TxD _x ₃ Q 20	361 TGE _x LAX _x ₄ PSxDP 34
	M 4 I 138 R _x ₂ S ₃ DRNG			

Pen^r protein fusions of class B-like II

Mtu	2864 M 11 13 G M 127 RAX _x EVGG	150 140	299 AxSx _x TxD _x ₃ Q 22	334 TGE _x LAX _x ₄ AGDA 13
Mle	1577 M 12 14 V V 127 RAX _x ELGG	26 140	278 AxSx _x TxD _x ₃ Q 22	338 TGQxLAX _x ₄ AGNA 13
Cdi	230 1 127 RAX _x SSDG	129 144	282 AxAx _x SxNx _x Q 23	318 TGE _x LAX _x ₄ DlxDK 13

Pen^r protein fusions of class B-like III

Sco	H63.18 M 19 21 A A 147 R _x ₂ LVGA	45 57	249 TxD _x ₃ Q 23	279 NGE _x LAX _x ₄ PSxGF 11
Sco	E87.08 M 15 17 V G 147 PVx _x DRNG	41 189 74	271 TxD _x ₃ Q 21	299 TGE _x LAX _x ₄ GHxFN 10
Sco	E87.07 M 5 7 V A	25 216	242 TxSx _x Q 21	270 TGE _x LAX _x ₄ RQxDG 11
sc1	pcb ^r M 24 26 V G 135 RAX _x DANG	47 176 69	253 TxD _x ₃ Q 22	282 TGE _x LAX _x ₄ PSxxx 10

FIG. 16. Class B Pen^r protein fusions. Bar code, motifs 1 to 7 and inserts (framed). For protein identifiers, see Table 1.

protein fusion Efam5 in this multiprotein complex would depend on the genetic backgrounds of the cells. The ampicillin-resistant mutant D344M512 (MIC, 2,000 µg of antibiotic ml⁻¹) and its parental ampicillin-susceptible mutant D344S (MIC, 0.03 µg of antibiotic ml⁻¹) both lack the Efam5-encoding gene (145, 205). In these particular genetic backgrounds, the amounts of (3→3) peptidoglycan made in the absence of Efam5 represent 100% (in mutant D344 M512) and 0.7% (in mutant D344S) of total peptidoglycan. In the genetic background which prevails in wild-type cells, resistance to penicillin is Efam5 concentration dependent. Efam5 might be an essential cell cycle protein that regulates the activities of the (4→3) and (3→3) peptidoglycan assembly molecular machines.

The ability of *S. aureus* to manufacture a (3→3) peptidoglycan has not been established. It is a possibility worthy of consideration. The amounts of (4→3) peptidoglycan made by Sau2a-producing strains grown in the presence of methicillin are drastically reduced compared to the amount made in the absence of the drug (46), suggesting that a spare Sau2a-dependent peptidoglycan which is not of the (4→3) type substitutes for the missing (4→3) peptidoglycan. Loss of FemX, which adds the first glycine residue to the ε-amino group of the L-lysine residue of peptidoglycan precursors, is lethal. In contrast, loss of FemA (197) restores the susceptibility of Sau2a-producing strains to penicillin, suggesting that peptidoglycan precursors having a single glycine substituent at position 3 of the stem peptides lack acceptor activity for the transfer reaction, which is presumably carried out by the acyltransferase module of Sau2a.

Susceptibility to penicillin is also restored by inactivating the gene that codes for the class A PBP fusion Sau2 (176), suggesting that the glycosyltransferase module of the PBP fusion Sau2 and the acyltransferase module of the Per^f protein fusion Sau2a function cooperatively in assembly of the putative spare peptidoglycan from lipid II precursor molecules. The implication of a free-standing glycosyltransferase is possible. Uncoupled glycosyltransferases occur in *S. aureus* and other bacterial species (174, 215).

Class A Pen^r Protein Fusions of Gram-Negative Bacteria

E. coli produces, in addition to its normal set of PBP fusions of classes A and B, a protein fusion, Eco1c, which has the barcode of a class A PBP fusion (Fig. 17) but escapes acylation by most of the β -lactam antibiotics tested. Eco1c, however, can be detected as a PBP with moxalactam and the iodinated Bolton-Hunter derivative of ampicillin (202, 204). Eco1c has glycosyltransferase activity, and this activity is blocked by moenomycin, indicating that Eco1c can form uncrosslinked peptidoglycan chains from lipid II precursor molecules (202).

The glycosyltransferase module of Eco1c falls within the continuum of known glycosyltransferase sequences (Fig. 10). It is related to the glycosyltransferase modules of PBP fusions Eco1a and Eco1b by *P* values of 10^{-26} (identity, 40%) and 7×10^{-18} (identity, 31%), respectively. Motif 6b at the amino-terminal region of the acyltransferase of Eco1c is not complete. The acyltransferase module falls outside subclasses A1, A2, A3, A4, and A5 (Fig. 10), indicating a distinct function. Eco1c is inert on both D-alanyl-D-alanine-terminated peptide and thioester analogues. Orthologues of Eco1c, assumed to resist

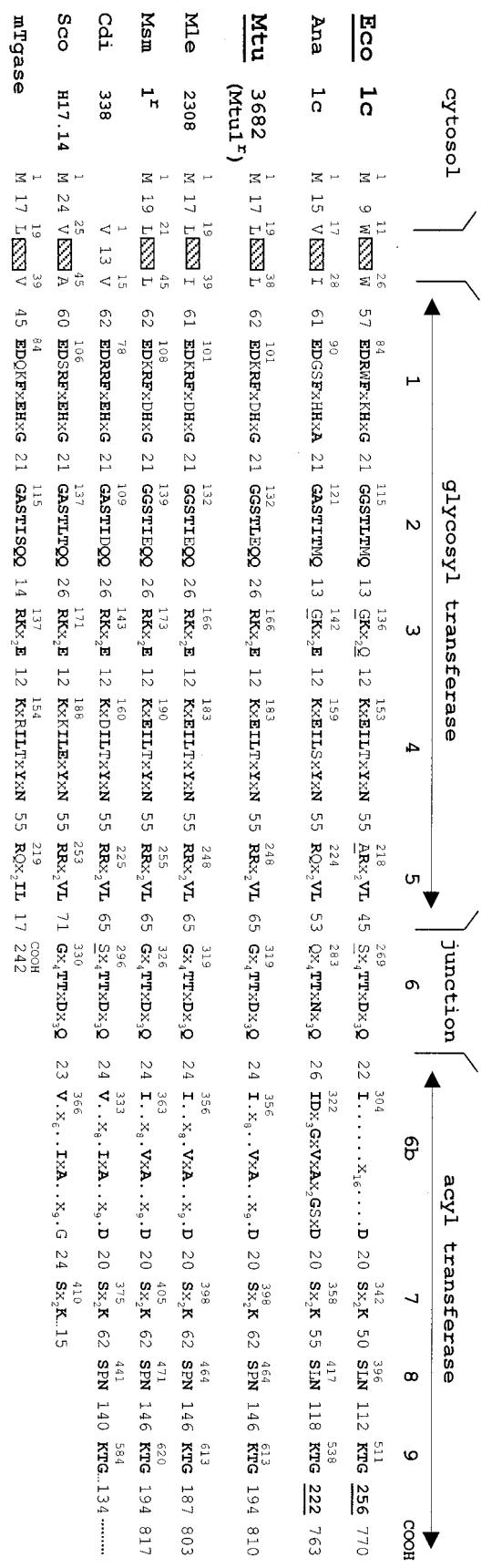


FIG. 17. Class A Pen^r protein fusions. Bar code, motifs 1 to 9 and extensions (underlined). For protein identifiers, see Table 1. mTgase, free-standing glycosyltransferase of *E. coli*.

TABLE 3. Protein orthologues of the SxxK Pen^r protein fusion Eco1c of *E. coli* motif 3: GKxxQ in Eco1c and RKxxE in PBP fusions Eco1a and Eco1b

Bacterial species ^a (reference)	Protein	Config	Motif 3 ^b
<i>Actinobacillus actinomycetemcomitans</i>	Aac1c	118	GKxxQ
<i>Agrobacterium tumefaciens</i> C58	Atu1c ^c		AKxxE
<i>Anabaena</i> 7120	Ana1c		GKxxE
<i>Bordetella bronchiseptica</i>	Bbr1c	1155	AKxxQ
<i>Bordetella parapertussis</i>	Bpa1c	1020	AKxxQ
<i>Bordetella pertussis</i>	Bpe1c	790	AKxxQ
<i>Burkholderia mallei</i>	Bma1c	5014	QKxxQ
<i>Burkholderia pseudomallei</i>	Bps1c	487	QKxxQ
<i>Caulobacter crescentus</i> * (166)	Ccr1c		AKxxQ
<i>Klebsiella pneumoniae</i>	Kpn1c	1192	QRxxA
<i>Mesorhizobium loti</i> * (117)	Mlo1c		SKxxQ
<i>Methylococcus capsulatus</i>	Mca1c	bmc26	QKxxQ
<i>Nostoc punctiforme</i>	Npu1c	615	GKxxE
<i>Pasteurella multocida</i> Pm70* (149)	Pmu1c		GKxxE
<i>Pseudomonas syringae</i> PV tomato	Psylc	E7	GKxxQ
<i>Pseudomonas putida</i> KT 2440	Ppu1c	13,538	GKxxQ
<i>Rickettsia montanensis</i> (7)	Rmo1c		GKxxQ
<i>Rickettsia typhi</i> (7)	Rty1c		GKxxQ
<i>Salmonella enterica</i> serovar Typhi CT 18	Sty1c ^d		GKxxQ
<i>Salmonella enterica</i> serovar Typhimurium LT2	Stym1c	2	GKxxQ
<i>Shewanella putrefaciens</i> 5/9/101	Spu1c	93	GKxxQ
<i>Xylella fastidiosa</i> 9a5c* (208)	Xfa1c		GKxxQ
<i>Xylella fastidiosa</i> Dixon	Xfad1c	286	GKxxQ
<i>Xylella fastidiosa</i> Ann-1	Xfaa1c	81	GKxxQ

^a *, genome sequence known.

^b Motif 3 is GKxxQ in Eco1c and RKxxE in Eco1a and Eco1b.

^c Accession no. gb|AE007870.

^d Accession no. emb|AL513382.

penicillin, occur in many gram-negative bacterial species (Table 3). A common marker is that the usual Arg residue at the amino terminus of motif 3 of the glycosyltransferase module is replaced by another amino acid residue, most often Gly. The acyltransferase modules are related to that of Eco1c by *P*

values ranging from $10^{-\infty}$ to 10^{-30} (Table 4). They are related to those of the PBP fusions Eco1a and Eco1b by *P* values larger than 10^{-20} .

In *E. coli*, the amount of Eco1c produced 34 h after onset of the stationary phase is low compared to the level produced at mid-exponential phase. It returns to normal after reinitiation of growth (47). The PBP fusions Eco1a, Eco2, and Eco3 behave like Eco1c. Eco1b, however, undergoes less important variations. The production of PBP fusions and free-standing PBPs is frequently growth phase dependent. The gram-negative organism *Pseudomonas aeruginosa* produces two PBP fusions, Pae3 and Pae3a, that belong to the same subclass, B3 (141). Pae3 and Pae3a are closely related to Eco3 by *P* values of 10^{-131} (identity, 45%) and 10^{-111} (identity, 41%), respectively. Pae3 (encoding gene in the *dcw* cluster) is produced maximally during the exponential phase of growth and decreases drastically when cells enter the stationary phase. Pae3a (encoding gene 2 Mbp away from the *dcw* cluster) is produced mainly, if not exclusively, during the stationary phase of growth. Loss of Pae3a is tolerated (141).

In the gram-positive sporeforming organism *Bacillus subtilis*, the genes encoding the PBP fusions Bsu2b of subclass B4 (249) and BsuVD (encoded by *spoVD*) of subclass B3 (42) occur in tandem in the *dcw* cluster (43). The essential Bsu2b is produced during both vegetative growth and sporulation. BsuVD is a mother cell PBP required for spore morphogenesis. These observations suggest that the Pen^r protein fusion Eco1c of class A is important for *E. coli* cells that are actively multiplying. Contrary to expectation, Eco1c is dispensable in the exponential phase of growth. Loss of Eco1c (encoding gene in the 57-min region) is harmless, and Eco1c cannot rescue an Eco1a^{ts}-Eco1b double mutant from death at the restrictive temperature (202).

The above studies refer to a time slice of what is a more global process. They do not include long-term survival and

TABLE 4. Similarity indexes (*P* and *I*) between the SxxK acyltransferase modules of Eco1c (494 amino acid residues) and Eco1b (397 amino acid residues), used as references, and homologous proteins of gram-negative bacteria

Bacterial species	Protein	Vs Eco1c			Vs Eco1b		
		P (10^{-n})	I (%)	Overlap (no. of aa)	P (10^{-n})	I (%)	Overlap (no. of aa)
Eco1c orthologues							
<i>E. coli</i> K-12	Eco1c	∞	100	1-494	7	24	26-283
<i>S. enterica</i> serovar Typhi CT18	Sty1c	∞	80	1-491	8	26	17-283
<i>P. putida</i> KT2440	Ppu1c	153	55	1-491	7	24	17-283
<i>X. fastidiosa</i>	Xfa1c	107	43	1-491	9	28	26-295
<i>M. loti</i>	Mlo1c	60	42	4-301	11	26	23-294
<i>P. multocida</i>	Pmu1c	40	26	5-485	12	25	26-282
<i>Anabaena</i> sp. strain 7120	Ana1c	36	33	4-305	21	27	4-329
<i>R. typhi</i>	Rty1c	32	31	26-280	9	22	14-277
Eco1b orthologues							
<i>X. fastidiosa</i>	Xfa1b	13	28	1-284	49	36	3-310
<i>P. putida</i> KT2440	Ppu1b	13	26	26-339	62	38	3-343
<i>H. influenzae</i>	Hin1b	9	22	21-335	80	44	3-342
<i>P. multocida</i>	Pmu1b	12	23	21-332	84	49	3-339
<i>S. enterica</i> serovar Typhi CT18	Sty1b	7	25	26-267	∞	83	2-397
<i>E. coli</i> K-12	Eco1b	7	24	1-267	∞	100	1-397
<i>E. coli</i> K-12	Eco1a	9	28	3-183	15	27	19-210

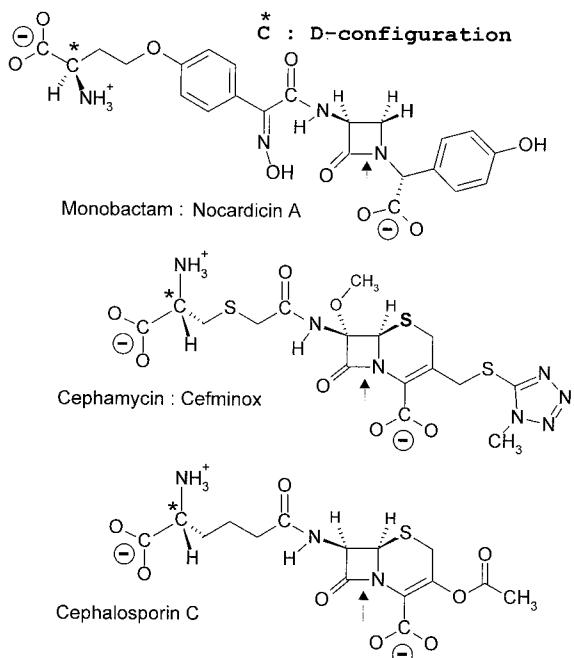


FIG. 18. β -Lactam antibiotics with side chains terminating in a D-configured $\text{NH}_2\text{-C}^*\text{HR-COOH}$ grouping.

growth under stress conditions. As the generation time of *E. coli* cells increases from 0.8 h to 6 h, conditions under which the amounts of Eco1a and Eco1b decrease drastically, the proportion of (4→3) peptidoglycan interpeptide linkages remains roughly unchanged (37.4% of total stem peptides versus 41.7%), the proportion of (3→3) peptidoglycan interpeptide linkages increases considerably (15.2% versus 3.1%), and uncrosslinked stem tripeptides represent about 50% of the total peptidoglycan (227). In stationary-phase cells, the peptidoglycan is hypercrosslinked and richer in bound lipoprotein molecules, and these molecules are attached almost exclusively to (3→3) peptide dimers (178).

The peptidoglycan-lipoprotein assemblage is hampered by nocardicin A, cephalosporin C, and cefminox (Fig. 18), the side chains of which terminate in a D-configured $\text{NH}_2\text{-CHR-COOH}$ grouping. These antibiotics cause rapid lysis of *E. coli* cells even at concentrations below the MICs (226). They are bactericidal for cells at the onset of the stationary phase of growth. They undergo covalent attachment to the peptidoglycan through a transfer reaction in which the D-amino group of the side chain serves as the acceptor. At micromolar concentrations, they prevent the lipoprotein molecules from being linked to the peptidoglycan, destabilizing the bacterial cell envelope.

The formation of the (3→3) peptidoglycan interpeptide linkages and the covalent attachment of the lipoprotein and β -lactam antibiotic molecules to the peptidoglycan both require the rupture of an L,D-peptide bond in the carbonyl donor. The carbonyl group at the L-center of meso-diaminopimelic acid is then transferred to an amino group borne by a carbon atom which is D-configured or, in the case of the lipoprotein attachment, is located at the end of the lateral chain of a L-lysine residue. The ability of Eco1c to perform acyltransferase activ-

ity has not been established, but by combining a glycosyltransferase module and an L,D-acyltransferase module catalyzing reaction II or reaction III (Fig. 6) in a single polypeptide chain, Eco1c could catalyze the three reactions on a competitive basis.

Assuming that the acyltransferase module of Eco1c functions as an N-acyl-L-diaminoacyl-D-dipeptidyl transpeptidase (reaction II in Fig. 6), D-alanyl-D-alanine is released into the periplasm. As discussed in references 136 and 138, the combined actions of a dipeptide transport system, a penicillin-resistant VanX-like Zn^{2+} -dependent D-alanyl-D-alanine dipeptidase, and pyruvate oxidase could enable the import of the released D-alanyl-D-alanine into the cytosol, its hydrolysis into D-alanine, and the oxidation of D-alanine to acetate and CO_2 , thus supplying energy. Alternatively, D-alanyl-D-alanine could be reutilized by the ligase MurF for lipid II recycling. The possibility also exists that the acyltransferase module of Eco1c functions as an N-acyl-L-diaminoacyl-D-alanine transpeptidase catalyzing reaction III (Fig. 6), after prior action of a penicillin-resistant D,D-carboxypeptidase catalyzing reaction IIIa. As discussed above in the section on class B PBP fusions, *E. coli* produces several peptidases which perform hydrolytic versions of reaction II and reaction III.

Gene deletion or disruption highlights the importance, if not the essentiality, of the Pen^r protein class A fusions in gram-negative bacteria. *E. coli* cells lacking Eco1a, Eco4, Eco5, Eco6, Eco6b, Eco7, EcoAmpC, and EcoAmpH and in which Eco2 and Eco3 are inactivated by amdinocillin and aztreonam, respectively, survive as large spherical bodies (48, 251). In particular genetic backgrounds, the combined activities of the PBP fusion Eco1b and the Pen^r protein fusion Eco1c may be sufficient to provide *E. coli* cells with a wall peptidoglycan that is no longer rod-shaped but is of sufficient tensile strength.

The cyanobacterium *Anabaena* sp. strain PCC7120 performs a higher-plant type of oxygenic photosynthesis. Under aerobic conditions in the presence of nitrate, it grows as filaments comprised of cells which all have the same morphology. In the absence of nitrate, some vegetative cells differentiate into nitrogen-fixing heterocysts. *Anabaena* sp. strain PCC7120 produces a panoply of PBP fusions (134). It also produces a protein fusion of class A, Ana1c, called PBP2 in reference 134, which is related to Eco1c by a *P* value of 10^{-69} (Tables 3 and 4; Fig. 17).

Disruption of the Ana1c-encoding gene causes no phenotypic differences under aerobic growth conditions in the presence of nitrate. In the absence of nitrate in a plate under nitrogen, the mutant still grows, but the filaments are shorter, they comprise vegetative cells of unequal size, and the heterocyst-like cells are distorted and have a thin cell envelope. In liquid medium without nitrate under aerobic conditions, the mutant dies, indicating that Ana1c plays an essential role in aerobic nitrogen fixation. Ana1c could be involved in the assembly of a particular peptidoglycan that would provide the heterocysts with a cell envelope capable of protecting the nitrogenase against oxygen. HcwA, a peptidoglycan hydrolase probably endowed with *N*-acetylmuramoyl-L-alanine amidase activity, is also required for heterocyst differentiation (256). The structure of the heterocyst peptidoglycan, however, is not known.

SxxK ACYLTRANSFERASES IN SPECIFIC ORGANISMS

Like *E. coli*, the mycobacteria manufacture peptidoglycans of the (4→3) and (3→3) types (Fig. 1 and 2). The SxxK acyltransferases of *Mycobacterium tuberculosis*, *Mycobacterium avium*, *Mycobacterium bovis*, and *Mycobacterium leprae* have been investigated in light of the accumulated knowledge about SxxK acyltransferases of groups I, II, and III. The acyltransferases of *M. avium* and *M. bovis* are close orthologues of the *M. tuberculosis* acyltransferases, and they are not discussed.

The study has been extended to *Corynebacterium diphtheriae*. Corynebacteria and mycobacteria belong to a suprageneric taxon of mycolic acid-containing actinomycetes. The mycolic acids of corynebacteria are relatively simple mixtures of saturated and unsaturated acids compared to the very complicated mixtures characteristic of mycobacteria (92, 116). *C. diphtheriae* manufactures a *meso*-diaminopimelic-containing (4→3) peptidoglycan similar to that of *M. tuberculosis*. Whether it also manufactures a (3→3) peptidoglycan has not been investigated.

The study has also been extended to *Streptomyces coelicolor*. *Streptomyces* spp. manufacture branched glycol-L,L-diaminopimelic acid peptidoglycans of the (4→3) and (3→3) types (Fig. 1 and 2). They secrete a collection of free-standing PBPs and wall-dissolving enzymes (77). In contrast to mycobacteria, which sometimes branch or form filaments that readily fragment into rods and coccoid bodies, *Streptomyces* spp. undergo very complex developmental processes (104). They produce branching networks of hyphae both on the surface of a solid substratum and into it to form a mycelium. Septa divide the hyphae into long cells containing several nucleoids. Upon growth completion, aerial filaments divide into unigenomic cells that further metamorphose into spores. *S. coelicolor* carries a large 8-Mb linear chromosome (192).

The *M. tuberculosis* (Mtu) and *M. leprae* (Mle) proteins are denoted according to the numbering of the encoding genes, *Rv1* to *Rv3923* and *Ml1* to *Ml2720*, respectively. The *C. diphtheriae* (Cdi) proteins are denoted according to the contig numbering. The *S. coelicolor* (Sco) proteins are denoted by the cosmid name and the CDS numbering. The assignment of the SxxK acyltransferases to different classes and subclasses rests on the bar codes borne by the proteins, the similarity indexes *P* and *I*, the known biochemical properties of several *Mycobacterium* and *Streptomyces* acyltransferases, the PBP pattern of *M. tuberculosis* H37Ra (35), and the environment of the encoding genes. The results are summarized in Table 5.

Class A PBP Fusions of *M. tuberculosis*, *M. leprae*, *M. smegmatis*, and *C. diphtheriae*

M. tuberculosis, *M. leprae*, and *C. diphtheriae* all have the information for a single PBP fusion of class A: Mtu0050 (Mtu1^s), Mle2688 (Mle1^s), and Cdi378, respectively (Table 5; Fig. 11). Mtu1^s (21) and Mle1^s (135) have been produced in *E. coli* with the correct membrane topology. The k_{+2}/K values of the interactions with benzylpenicillin, ampicillin, amoxicillin, cefotaxime, cefuroxime, and ceftriaxone range from $\approx 100,000 \text{ M}^{-1} \text{ s}^{-1}$ to $\approx 10,000 \text{ M}^{-1} \text{ s}^{-1}$. Ticarcillin, piperacilline, methicillin, and moxalactam are not acylating agents. Mtu1^s has been obtained in a soluble ($\Delta G95\text{-Q143}$) truncated

TABLE 5. SxxK acyltransferases of *M. tuberculosis*, *M. leprae*, *C. diphtheriae*, and *S. coelicolor*^a

Type	Class	PBP and Pen ^r protein fusions				Similarity index vs <i>M. tuberculosis</i>						Gene environment	
		<i>M. tuberculosis</i>		<i>M. leprae</i>	<i>C. diphtheriae</i>	<i>P</i> (10^{-n})		<i>I</i> (%)		<i>M. leprae</i>	<i>C. diphtheriae</i>	<i>S. coelicolor</i>	
		Protein	M _r			<i>M. leprae</i>	<i>C. diphtheriae</i>	<i>S. coelicolor</i>	<i>M. leprae</i>				
PBP	A	0050 (Mtu1 ^s)	71,151	94–82	2688 (Mle1 ^s)	378	H24.23 E6.34	8	173	74	83	51	37
B3		2163 (Mtu3)	72,506	94–82	0908 (Mle3)	101	K7.12 4A10.23 (Sco3)	8	163	41	80	51	35
B2							C88.19 (Sco2)	8	107	80	85	45	30
B-like I		0016	51,776	52	0018	16	H69.17 6G9.32 2st10A7			79	78	39	33
												35	<i>dew</i>
													<i>mre, sfr</i>
													<i>sfr</i>
Pen ^r	A	3682 (Mtu1 ^r)	84,636	2308 (Mle1 ^r)	338	H17.14		8	∞	51	86	48	36
B-like II		2864	63,015	1577	230		H63.18	8	118	80	41	27*	27*
B-like III							E87.08					34*	34*
							E87.07					33*	33*

^a Similarity indexes *P* and *I* apply to full-size protein fusions except for *S. coelicolor* protein fusions H63.18, E87.08, and E87.07 (*), where similarity with protein fusion Mtu2864 is restricted to the acyltransferase modules.

form that adopts an active folded state in terms of penicillin binding. Mle1^s is relatively thermolabile, with half-lives of 60 min at 25°C and less than 5 min at 37°C. It catalyzes hydrolysis and aminolysis of thiolester analogues of *N*-acyl-D-alanyl-D-alanine-terminated peptides.

Mycobacterium smegmatis mc²155 is a fast-growing mycobacterium. Cultures in Luria-Bertani medium under aeration enter the stationary phase after 40 h. As a result of screening for transpositional mutants with permeability defects, an *M. smegmatis* strain in which a gene was disrupted at a locus coding for the sequence immediately downstream from motif 5, RWxxVL, of a glycosyltransferase module was obtained (22). This glycosyltransferase module is linked to an acyltransferase module which is a close orthologue of the acyltransferase module of Mtu1^s (P , 10^{-152} ; identity, 79%), indicating that the disrupted gene (accession number AF165523) codes for a PBP fusion, Msm1^s, of class A (Fig. 11). For comparison purposes, the acyltransferase module of Msm1^s and that of Mtu1^r (see the ensuing section) are distantly related by a P value of 9×10^{-13} (identity, 26%).

Compared to the parental strain, the Msm1^s-less mutant has a twofold-higher permeability for glycine; it grows more slowly in the exponential phase; the protein content of the culture at the onset of the stationary phase is reduced by 90%; and the mutant has an increased susceptibility to β-lactam antibiotics (MICs, 15 µg of ampicillin ml⁻¹ versus 300 µg of ampicillin ml⁻¹ for the parental strain in the absence of a β-lactamase inactivator).

Chromosomal duplication in *M. smegmatis* (its genome is nearly 7 Mbp long, i.e., about 50% larger than that of its slow-growing relatives) results in duplicate copies of many genes and is a potent source of genomic dynamics (73). The analysis of the benzylpenicillin-labeled membranes of the Msm1^s-less mutant failed to reveal the presence of any residual Msm1^s PBP fusion. Considering that the locus at which the Msm1^s-encoding gene was disrupted, the amino-terminal region of the disrupted protein is an intact glycosyltransferase module. This module could work cooperatively with the acyltransferase module of a PBP fusion of class B, which itself could be made more readily accessible to the drug because of defects in the cell envelope, explaining why the mutant is viable and is more susceptible to penicillin than the parental strain.

Class A Pen^r Protein Fusions of *M. tuberculosis*, *M. leprae*, *M. smegmatis*, and *C. diphtheriae*

M. tuberculosis, *M. leprae*, and *C. diphtheriae* all have the information for a single Pen^r fusion of class A: Mtu3682 (Mtu1^r), Mle2308 (Mle1^r), and Cdi338, respectively (Table 5 and Fig. 17). The acyltransferase modules of the pair Mtu1^r-Mle1^r and the pair Mtu1^s-Mle1^s are distantly related by a P value of $\cong 10^{-17}$ (identity, 25%). Mle1^r has been produced in *E. coli* (16). The k_{+2}/K values of the interactions are 5 to 10 M⁻¹ s⁻¹ with ceftriaxone and benzylpenicillin and 1 M⁻¹ s⁻¹ and even smaller with ampicillin, amoxicillin, cefoxitin, ticarcillin, temocillin, and cephaloridine. Mle1^r lacks activity on thioesters, which are carbonyl donors of Mle1^s. It has a high thermostability, with no loss of its low affinity for penicillin after 10 min at 60°C. The sequence Leu174 to Arg314 contains a membrane association site (144). The acyltransferase module

can be produced intracellularly in *E. coli* as inclusion bodies which can be refolded in a form that has the same low affinity for β-lactams as the full-size protein fusion (144). Mtu1^r has not been characterized biochemically. In all likelihood, it has the same low affinity for β-lactam antibiotics as its orthologue Mle1^r (P , $10^{-\infty}$; identity, 86%).

M. smegmatis mc²155 survives for very long periods of time, 150 days or more, during either carbon, nitrogen, or phosphorus starvation (209). The viability of cultures in rich Lab-Lemco medium under aeration remains unchanged, about 10⁹ CFU ml⁻¹, for 40 days. It then declines to about 5 × 10⁷ CFU ml⁻¹ at day 120 (121). Like *M. tuberculosis* and *M. leprae*, *M. smegmatis* has the information for a Pen^r protein fusion, Msm1^r, of class A (Fig. 17), which is a close orthologue of Mtu1^r (P , $10^{-\infty}$; identity, 80%). As a result of screening for transpositional mutants with long-term survival defects, an *M. smegmatis* mutant in which the Msm1^r-encoding gene (now identified in contig 2889) is disrupted at a locus coding for motif 1 of the glycosyltransferase module was obtained (121). Compared to the parental strain, cultures of the Msm1^r-less mutant had a much impaired viability, 10⁶ CFU ml⁻¹, i.e., 0.1% of the parental value, at day 40. Subsequently, however, viability returned and reached the level of the parental strain at day 120. Other mutants also impaired in long-term survival but in which the predicted functions of the disrupted genes are in biotin and polyketide biosynthesis behave the same way (121).

M. smegmatis manufactures peptidoglycans of the (4→3) and (3→3) types (see the Introduction). Loss of the PBP fusion Msm1^s of class A is deleterious to *M. smegmatis* cells in the exponential phase, indicating that Msm1^s is an important component of the molecular machine implicated in (4→3) peptidoglycan synthesis from lipid II precursor molecules. Loss of the Pen^r protein fusion Msm1^r of class A is very damaging for cells in long-term stationary phase. A (3→3) peptidoglycan especially adapted to survival under starvation conditions might be synthesized from lipid II by Msm1^r.

Subclass B3 and B-Like I PBP Fusions of *M. tuberculosis*, *M. leprae*, and *C. diphtheriae*

M. tuberculosis, *M. leprae*, and *C. diphtheriae* all have the information for a protein fusion of class B: Mtu2163 (Mtu3), Mle0908 (Mle3), and Cdi101, respectively (Table 5; Fig. 12). The linker and acyltransferase modules of Mtu3 diverged in concert and fall into the same subclass, B3 (Fig. 10). The acyltransferase module of Mtu3 is more closely related to the acyltransferase modules of Eco3 of *E. coli* and Pae3 of *P. aeruginosa* of subclass B3 (P , 10^{-32} and 3×10^{-48} , respectively; identity, 30% and 34%, respectively) than to the acyltransferase module of Eco2 of subclass B2 (P , 9×10^{-13} ; identity, 25%). In *M. tuberculosis*, Rv2163 codes for Mtu3; Rv2150c and Rv2154c code for protein homologues of the *E. coli* cell separation proteins FtsZ and FtsW, respectively; Rv2151c codes for a protein of the FtsQ-like family; and Rv2152c, Rv2153c, Rv2155c, Rv2156c, Rv2157c, and Rv2158c code for protein homologues of the *E. coli* MurC, MurG, MurD, MurX (i.e., MraY), MurF, and MurE, respectively. Altogether, they form the *M. tuberculosis* dcw cluster (Fig. 5). On the basis of these predictive studies (and in the absence of direct biochemical data), Mtu2163 (Mtu3), Mle0908 (Mle3), and

Cdi101 are provisionally identified as PBP fusions of subclass B3.

M. tuberculosis, *M. leprae*, and *C. diphtheriae* all have the information for another protein fusion of class B: Mtu0016, Mle0018, and Cdi16, respectively (Table 5; Fig. 12). The acyltransferase modules of Mtu0016 and Mtu3 of subclass B3 are distantly related by a *P* value of 10^{-13} (identity, 25%), indicating that they belong to different subclasses of class B. In contrast to the linker module of Mtu3, which is 147 amino acid residues long, the linker module of Mtu0016 is 110 amino acid residues long, it lacks motif 1 but contains motifs 2 and 3. Mtu0016 and the 484-amino-acid PBP fusion SgrpbpA of *Streptomyces griseus* (114) are related by a *P* value of 10^{-72} (identity, 36%) over the entire sequences. Both the Mtu0016- and SgrpbpA-encoding genes are located in the neighborhood of a gene that codes for a protein of the SFR family. Disruption of the SgrpbpA-encoding gene has no obvious effects on the growth and sporulation of *S. griseus*. *Streptomyces* spp. have a plethora of PBPs. SgrpbpA and other PBP fusions may substitute for each other. On the basis of these predictive studies, Mtu0016, Mle0018, and Cdi16 are provisionally identified as PBP fusions of a particular subclass, B-like I.

Class B-Like II Pen^r Protein Fusions of *M. tuberculosis*, *M. leprae*, and *C. diphtheriae*

M. tuberculosis, *M. leprae*, and *C. diphtheriae* all have the information for a protein fusion: Mtu2864, Mle1577, and Cdi230, respectively (Table 5, Fig. 16). They each have a not-well-conserved motif 1, lack motifs 2 and 3, and contain a 127-amino-acid insert located between the membrane anchor and the presumed motif 1, a structural feature which is reminiscent of the Pen^r protein fusions of subclass B1. *M. tuberculosis* does not produce any detectable PBPs migrating at a middle distance between the 82-kDa and 52-kDa PBPs on sodium dodecyl sulfate gels (35).

It is likely that Mtu2864 (theoretical *M*_r 63,015) does not bind penicillin. Consistently, its acyltransferase module is distantly related to the acyltransferase module of the PBP fusion Mtu3 by a *P* value of 10^{-15} (identity, 32%). It is related to the acyltransferase module of the Pen^r protein fusion SlcpbR of *Streptomyces clavuligerus* (encoding gene downstream from the isopenicillin synthase-encoding gene of the cephamicin cluster) (172) by a *P* value of 5×10^{-38} (identity, 35%). SlcpbR is essential in stationary-phase cultures, conditions under which cephamicin is produced. On the basis of these predictive studies, Mtu2864, Mle1577, and Cdi230 are provisionally identified as Pen^r protein class B-like II fusions.

Free-Standing PBPs, β-Lactamases, and Related Polypeptides of *M. tuberculosis* and *M. leprae*

The polypeptides fall into four groups (Fig. 14). *M. tuberculosis* has the information for three free-standing polypeptides, Mtu3627, Mtu3330, and Mtu2911, and *M. leprae* has the information for two free-standing polypeptides, Mle0211 and Mle0691, which bear the bar codes SxxK, SxN, and KTG (Fig. 14). They are probably auxiliary cell cycle proteins implicated in (4→3) peptidoglycan synthesis. Mtu3330 and Mle0691 are

orthologues belonging to the MEROPS S11 family. Mtu3627 and Mle0211 are also orthologues, belonging to the MEROPS S13 family. Like Eco4, they contain a large insert between the SxxK motif and the SxN motif. An *M. smegmatis* PBP has been isolated (15). Its apparent molecular mass, 49.5 kDa, is similar to those of Eco4, Mtu3627, and Mle0211. It has a great affinity for benzylpenicillin, ampicillin, and cefoxitine. It catalyzes hydrolysis and aminolysis of bisacetyl-L-lysyl-D-alanyl-D-alanine (15, 159).

M. tuberculosis but not *M. leprae* has the information for three free-standing polypeptides, Mtu0907, Mtu1730, and Mtu1922, which bear the modified bar codes SxxK, YxN, and HxG (Fig. 14), indicating that they are not related to wall peptidoglycan biochemistry. They belong to the MEROPS S12 family. The *P* values for the pair Mtu1730-Mtu0907 and the pair Mtu1730-Mtu1922 are 4×10^{-23} (identity, 30%) and 3×10^{-18} (identity, 29%), respectively. Mtu1730 is related to the free-standing PBP4 of *Nocardia lactamdurans* (encoding gene within the cluster involved in cephamicin biosynthesis) by a *P* value of 10^{-45} (identity, 34%). Mtu1922 is related (for more than 80% of its entire sequence) to the *Streptomyces* sp. strain R61 D,D-carboxypeptidase-PBP (*P*, 2×10^{-19} ; identity, 29%), the endopeptidase ADP of *B. cereus* (*P*, 8×10^{-20} ; identity, 29%); and the catalytic module of the D-aminopeptidase DAP of *O. anthropi* (*P*, 5×10^{-18} ; identity, 28%) (Fig. 15).

M. tuberculosis produces a β-lactamase of class A, Mtu2068 (Fig. 14), which is related to a β-lactamase of *Nocardia lactamdurans* by a *P* value of 9×10^{-62} (identity, 50%). *M. leprae* apparently lacks the information for a β-lactamase of class A similar to Mtu2068. However, *M. leprae* isolated from the tissues of experimentally infected armadillos treated with benzylpenicillin 6 months or more before sacrifice has β-lactamase activity (181).

M. tuberculosis has the information for a free-standing polypeptide, Mtu1367, which has motif 1, SxxK, and motif 2, YxH, but lacks motif 3 (Fig. 14). Mtu1367 is related to a triacylglycerol lipase of *Pseudomonas* sp. strain 109 by a *P* value of 2×10^{-34} (identity, 36%). In *M. leprae*, pseudogene Ml0528 carries the same information.

PBP Fusions and Pen^r Protein Fusions of *S. coelicolor*

S. coelicolor has the information for three class A PBP fusions, ScoH24.23, ScoE6.34, and ScoK7.12 (Fig. 11). They are orthologues of Mtu1^s (*P* ≤ 10^{-40}) (Table 5).

S. coelicolor has the information for one PBP fusion of subclass B3, Sco4A10.23 (Fig. 12). Sco4A10.23 is an orthologue of Mtu3 (*P*, 10^{-82}) (Table 5). In *S. coelicolor*, 4A10.23c codes for Sco4A10.23 (Sco3), and 4A10.22c to 4A10.15c code for protein homologues of *E. coli* MurE, MurF, MraY, MurD, FtsW, MurG, FtsQ, and FtsZ (192). Altogether, they form a cluster equivalent to the cell septation *dcw* cluster of *E. coli* (Fig. 5). An *S. coelicolor* FtsZ null mutant is viable and produces aerial hyphae. It is unable to produce spores, indicating that FtsZ is required for the hyphae to undergo septation into uninucleoid cells that differentiate into spores (152). The manipulation of a developmentally controlled FtsZ promoter also generates a nonsporulating *S. coelicolor* strain (66).

S. coelicolor has the information for three PBP fusions of

subclass B-like I, ScoH69.17, ScoG9.32, and Sco2st10A7 (Fig. 12). They are orthologues of Mtu0016 ($P \approx 10^{-80}$) (Table 5).

S. coelicolor has the information for one PBP fusion of subclass B2, ScoC88.19 (Sco2) (Fig. 12). Sco2 has no equivalent in *M. tuberculosis*, *M. leprae*, or *C. diphtheriae*. The linker and acyltransferase modules of Sco2 diverged in concert in a way that indicates relatedness with Eco2 of subclass B2 (Fig. 10). In addition, the acyltransferase module of Sco2 is more closely related to the acyltransferase module of Eco2 ($P, 6 \times 10^{-31}$; identity, 30%) than to the acyltransferase module of Eco3 of subclass B3 ($P, 7 \times 10^{-11}$; identity, 21%). In *S. coelicolor* (32), C88.19c codes for Sco2; C88.22c, C88.21c, and C88.20c code for protein homologues of the *E. coli* actin-like MreB, MreC, and MreD, respectively; and C88.18c codes for a protein of the SFR family. Altogether, they form a cluster equivalent to the rod shape-determining *mre* cluster of *E. coli* (241).

S. coelicolor has the information for a protein fusion, ScoH17.14 (Fig. 17), which is an orthologue of the Pen^r protein fusion Mtu1^r of class A ($P, 10^{-51}$) (Table 5).

Finally, *S. coelicolor* has the information for three protein fusions, ScoH63.18, ScoE87.08, and ScoE87.07 (Fig. 16), the acyltransferase modules of which are related to the acyltransferase module of the Pen^r protein fusion Mtu2864 of class B-like II (by P values ranging from 10^{-27} to 10^{-38}) (Table 5). The linker modules, however, lack statistically significant similarity with the linker modules of the Pen^r protein fusions of class B-like II. On this basis, ScoH63.18, ScoE87.08, and ScoE87.07 are identified provisionally as Pen^r protein fusions of a particular class B-like III.

Detected versus Predicted PBPs of *M. tuberculosis* H37Ra

M. tuberculosis H37Ra produces four PBPs of 94, 82, 52, and 37 kDa (35). The 94- and 82-kDa PBPs migrate close to each other on sodium dodecyl sulfate gels because of poor resolution of large proteins. Their molecular masses are almost certainly overestimated. Thus, disruption of the *M. smegmatis* gene encoding the 715-amino-acid PBP fusion Msm1^s (theoretical M_r 74,459) causes the disappearance from the gel of a protein band of 95 kDa (22). Identifying the experimentally detected PBPs with the predicted PBP fusions of class A, class B, class B-like I, and free-standing PBPs remains somewhat ambiguous. It is likely that the *M. tuberculosis* 94- to 82-kDa protein bands comprise the PBP fusions Mtu1^s of class A (theoretical M_r 71,119) and Mtu3 of subclass B3 (theoretical M_r 72,506). The 52-kDa PBP sometimes appears as a doublet (35). The 52-kDa protein band probably comprises the PBP fusion Mtu0016 of subclass B-like I (theoretical M_r 51,777) and the free-standing PBP Mtu3627 (theoretical M_r 46,835), the equivalent of Eco4. In turn, the 37-kDa PBP is probably Mtu3330 (theoretical M_r 41,683), the equivalent of Eco5 or Eco6 in *E. coli*.

The specificity profiles of the 94-, 82-, 52-, and 37-kDa PBPs for β -lactam antibiotics have been expressed in k_{+2}/K values (Table 2) from the published IC₅₀ values (35). Imipenem is a potent inactivator of the 94- to 82-kDa PBP fusions (k_{+2}/K values from 2,000 to 6,000 M⁻¹ s⁻¹). The 52-kDa PBP fusion is less susceptible to the β -lactams.

WHERE NEXT?

Bacterial cell walls were isolated for the first time in the early 1950s (200). Many facets of the wall biochemistry and mode of action of penicillin have been disclosed. Bacteria that are actively multiplying, conditions under which they manufacture a wall peptidoglycan of the (4→3) type, are killed by β -lactam antibiotics. SxxK D,D-acyltransferases implicated in the assembly of the polymer are immobilized by the drugs in the form of stable, inactive PBPs. Class A PBP fusions comprised of a glycosyltransferase module and a D,D-acyltransferase module of class A, class B PBP fusions comprised of a linker (peptide recognition) module and a D,D-acyltransferase module of class B, free-standing D,D-acyltransferases-PBPs, and a set of non-penicillin-binding cell cycle proteins assemble into a morphological apparatus that ensures the formation of a nascent (4→3) peptidoglycan from lipid II precursor molecules and the remodeling of the wall polymer throughout cell expansion and division. PBP fusions of classes A and B are globally the lethal targets of the β -lactam antibiotics.

In spite of these advances, we are still in the reductionist approach of dissecting the constituents of the morphogenic apparatus. The known molecular parts will have to be reassembled into functional entities in order to resolve the issues of the process. The modes of internal communication, transmission of the directives eliciting the right responses, and coordination of many events from DNA replication to cell division represent work for years to come. Even the question of how the different assortments of PBPs determine cell growth and division in exponential-phase cultures of different bacterial species is left open.

One PBP fusion each in classes A and B is probably the minimal set required for a bacterium, such as the coccus-shaped *N. meningitidis*, to multiply. The rod-shaped *E. coli* has two class A PBP fusions (they can substitute for each other) and two class B PBP fusions (one of which, Eco3, can substitute for the other, Eco2, in particular genetic backgrounds). *M. tuberculosis*, *M. leprae*, and *C. diphtheriae* have one PBP fusion each in classes A, B, and B-like I. Class B-like I PBP fusions, which have an incomplete linker module, may be typical of the actinomycetes.

S. coelicolor undergoes complex developmental processes. It has the information for three PBPs of class A, one PBP fusion each in subclasses B2 and B3, and three class B-like I PBP fusions. The genus *Chlamydia* is unusual in many respects (81). These gram-negative bacteria are susceptible to penicillin. They undergo binary fission as reticulate bodies within the chlamydial inclusion. They have the information for the synthesis of a lipid II precursor similar to that of *E. coli* (216). However, they are peptidoglycanless. Consistently, they lack the information for class A PBP fusions. They produce two class B PBP fusions which perhaps manufacture a wall glycan-less-peptide polymer in a penicillin-sensitive manner (81).

Phylogenetically distant bacterial species, such as *E. faecium*, *E. coli*, other gram-negative bacteria, *Mycobacterium* spp., and *Streptomyces* spp., have the dual ability to manufacture a (4→3) peptidoglycan in a penicillin-susceptible manner and a (3→3) peptidoglycan in a penicillin-resistant manner. The conjecture is that, basically, the polymeric (3→3) peptidoglycans are formed from lipid II precursor molecules by the combined

activities of a glycosyltransferase and a penicillin-resistant acyltransferase that has undergone a change from the D,D specificity (reaction I in Fig. 6) to the L,D specificity (reaction II or III in Fig. 6). The results of the studies reported here lead to the conclusion that the (3→3) peptidoglycan-synthesizing machines are different in different bacterial species. They raise a host of problems that remain to be worked on.

E. faecium has two acyltransferase activities, a Pen^r N-acyl-D-alanyl-D-alanine D,D-carboxypeptidase and a Pen^r N-acyl-L-lysyl-D-alanine-cleaving transpeptidase, that would be required to make (3→3) L-lysyl-N^ε-(D-isoasparaginyl)-L-lysine cross-bridges (Fig. 2) according to reactions IIIa and III shown in Fig. 6. The activities have been detected, but the enzymes have not been isolated and characterized biochemically; the Pen^r D,D-carboxypeptidase could be a Zn²⁺-dependent hydrolase, and the glycosyltransferase remains hypothetical.

E. faecium also produces a Pen^r SxxK protein fusion, Efam5, of subclass B1. The encoding gene is in the neighborhood of a gene that codes for an SFR protein, suggesting that in exponential-phase cultures, Efam5 plays the role of a cell cycle protein in (3→3) peptidoglycan metabolism similar to the roles that the class B PBP fusions play in (4→3) peptidoglycan metabolism. Consistent with this view, *E. faecium* cells in the exponential phase of growth exhibit various levels of resistance to penicillin in an Efam5 concentration-dependent manner, and they manufacture a (3→3) peptidoglycan in various proportions of total peptidoglycan. In particular genetic backgrounds, however, Efam5 is dispensable. An Efam5-less mutant which manufactures exclusively a (3→3) peptidoglycan during growth has been isolated. It has a MIC for ampicillin of greater than 2 mg of antibiotic ml⁻¹.

What is true for *E. faecium* is probably true for other enterococci and the staphylococci. Like Efam5 in *E. faecium*, the Pen^r SxxK protein fusion Sau2a of subclass B1 is a penicillin resistance determinant in exponential-phase cultures of *S. aureus*. However, the ability of *S. aureus* to manufacture a (3→3) peptidoglycan has not been established.

The (3→3) peptidoglycan-synthesizing machines of *E. coli*, *M. tuberculosis*, *M. leprae*, and *M. smegmatis* appear to be different from that of *E. faecium* in terms of composition and growth phase dependency. *E. coli* and the mycobacteria all produce a Pen^r SxxK protein fusion of class A, Eco1c, Mtu1^r, Mle1^r, and Msm1^r, respectively. By combining a glycosyltransferase module and a Pen^r acyltransferase module having the L,D specificity, the Pen^r protein class A fusions would have the activities required to carry out (3→3) peptidoglycan assembly from lipid II precursor molecules in a penicillin-resistant manner, either directly, according to reaction II shown in Fig. 6, or after the action of a Pen^r D,D-carboxypeptidase, according to reaction III.

The Pen^r protein class A fusions are not penicillin resistance determinants in exponential-phase cultures of *E. coli* and *Mycobacterium* spp., conditions under which they multiply in a penicillin-susceptible manner. Eco1c does not rescue an *E. coli* strain that has the double mutation Eco1a^{ts} Eco1b from death at the nonpermissive temperature. Wild-type *E. coli* cells manufacture a (3→3) peptidoglycan in increasing proportions of total peptidoglycan and become increasingly more resistant to penicillin when they divide at decreasing growth rates and when they enter the stationary phase. An *M. smegmatis* mutant

that lacks Msm1^r shows much impaired viability in long-term survival, conditions under which the wild-type strain manufactures large amounts of (3→3) peptidoglycan (see the Introduction). These observations lead to the plausible conclusion that (3→3) peptidoglycan synthesis and intrinsic resistance to penicillin at the level of the target proteins are dependent on class A Pen^r protein fusions in *E. coli* and mycobacteria exposed to adverse growth conditions.

Stationary phase and long-term survival are not synonymous with no metabolism and no cell division as it occurs in sporulation. The stringent response is a vast transcriptional program that encompasses many genes. It is initiated by increased concentrations of ppGpp and pppGpp, made by RelA and SpoT in *E. coli* and by Rel in *M. tuberculosis* (33, 182). *E. coli* lacking RelA and SpoT cannot grow in minimal medium and hardly survives in stationary phase. One induced regulon concerns genes whose expression depends on the *rpoS*-encoded sigma factor σ^s. Mutations in *rpoS* and other unlinked loci confer on cells in aged cultures a competitive advantage that is restricted to the stationary phase over cells of a young population, so that the mutants take over the bacterial population (253).

Many genes form a defensive system aimed at avoiding the damaging effects of ongoing respiratory activity (167). Superoxide dismutases are important under aerobic carbon starvation. An alkylhydroperoxide reductase is required under aerobic phosphate starvation (158). About 2% of the genome is under the control of the nitrogen regulatory protein C, which activates operons whose products minimize the effects of nitrogen-limiting conditions (258). The peptidyl prolyl isomerase SurA, involved in protein folding in the periplasm, is essential for stationary-phase survival (133, 225).

Likewise, Rel is essential to *M. tuberculosis*. Persistent cells incorporate radioactive uridine into RNA and produce 16S RNA and mRNA transcripts (107). The glyoxylate shunt enzyme isocitrate lyase is essential (153). The mRNA level of the sigma factor SigJ-encoding gene is upregulated 20-fold in 100-day cultures compared to exponential-phase cultures (106). Stationary-phase cultures of *M. smegmatis* are a dynamic population of cells. Growth and cell division occur and mutations accumulate which lead to the emergence of variants more adapted to survival (209). In light of these data, a shift of peptidoglycan synthesis from the (4→3) type to the (3→3) type could be the first known response to starvation and other stress conditions that would affect wall peptidoglycan metabolism.

According to the proposed scenarios, lipid II precursor molecules perform dual functions. The SxxK D,D-acyltransferase modules of the PBP fusions examine the stem pentapeptides, search for D-alanyl-D-alanine sequences, and use them as carbonyl donors for the synthesis of (4→3) interpeptide linkages or cross-bridges in a penicillin-susceptible manner (reaction I in Fig. 6). The SxxK L,D-acyltransferase modules of the Pen^r protein fusions would search for L-diaminoacyl-D-alanyl-D-alanine or L-diaminoacyl-D-alanine sequences and use them as carbonyl donors for the synthesis of (3→3) interpeptide linkages or cross-bridges (reactions II and III in Fig. 6) in a penicillin-resistant manner. These transfer reactions differ from the transfer reactions through which many surface proteins are attached covalently to the peptidoglycan in gram-positive bacteria. In this case, cysteine transferases, called sortases (109, 150), identify the signal sequence LPxTG at the carboxy end of

the surface protein as the carbonyl donor, cleave the threonylglycine bond, and transfer the carbonyl of the threonine residue to the ω -amino group at position 3 of peptidoglycan precursor molecules.

Little is known about the Pen^r SxxK protein fusions beyond their cursory description. Further studies may well reshape our views on peptidoglycan biochemistry, offer a clue to the myco-bacterial paradox, and provide much-needed information for a full understanding of the intrinsic resistance to penicillin of important bacterial pathogens.

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