Substitutions in PBP2b from β-Lactam-resistant Streptococcus pneumoniae Have Different Effects on Enzymatic Activity and Drug Reactivity*

Received for publication, October 25, 2016, and in revised form, December 22, 2016 Published, JBC Papers in Press, January 6, 2017, DOI 10.1074/jbc.M116.764096

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Edited by Gerald W. Hart

Pneumococci resist β-lactams by expressing variants of its target enzymes, the penicillin-binding proteins (PBPs), with many amino acid substitutions. Up to 10% of the sequence can be modified. These altered PBPs have a much reduced reactivity with the drugs but retain their physiological activity of cross-linking the peptidoglycan, the major constituent of the bacterial cell wall. However, because β-lactams are chemical and structural mimics of the natural substrate, resistance mediated by altered PBPs raises the following paradox: how PBPs that react poorly with the drugs maintain a sufficient level of activity with the physiological substrate. This question is addressed for the first time in this study, which compares the peptidoglycan cross-linking activity of PBP2b from susceptible and resistant strains with their inhibition by different β-lactams. Unpredicted, the enzymatic activity of the variants did not correlate with their antibiotic reactivity. This finding indicates that some of the numerous amino acid substitutions were selected to restore a viable level of enzymatic activity by a compensatory molecular mechanism.

Penicillin and other β-lactams are arguably the most important drugs ever, having had a global impact on human health in seven decades of continuous use to fend off bacterial infection (1). β-Lactams hamper formation of the peptidoglycan, which is the main constituent of the bacterial cell wall. Peptidoglycan is a giant polymer encasing the cell and consists of chains of tandemly repeated disaccharides cross-linked by peptide bridges. This cross-linking results from a transpeptidation reaction catalyzed by enzymes that are inhibited by β-lactams, which are mimics of the donor dipeptide of the transpeptidation reaction. The enzymes responsible for the peptidoglycan assembly are called penicillin-binding proteins (PBPs). They all have a penicillin-binding domain that generally catalyzes the transpeptidation but can also act as a carboxypeptidase or endopeptidase in some cases (2). Some PBPs have an additional transglycosylase domain that catalyze the elongation of the glycan strands. The bifunctional transpeptidase and transglycosylase PBPs constitute the class A, whereas the monofunctional transpeptidase PBPs are of class B.

Resistance to antibiotics is recognized as a major threat to human health (3). Resistance to β-lactams, which are the most widely used antimicrobials, is particularly worrisome. A case in point is that of pneumococcus, a major human pathogen that causes otitis, pneumonia, and meningitis and that is estimated to cause over 1.5 million deaths per year (4). Pneumococcus and related oral streptococci resist β-lactams by expression of altered PBPs (5) encoded by mosaic genes that result from multiple events of homologous recombination with genes from close species combined with additional point mutations (6). A similar mechanism of β-lactam resistance operates in Neisseria species (7). Mosaic PBPs from resistant strains typically harbor tens of amino acid substitutions amounting to more than 10% of the primary sequence in some instances (5).

Of the six PBPs in Streptococcus pneumoniae, three are commonly altered in β-lactam resistant strains: the two monofunctional transpeptidase PBP2b and PBP2x and the bifunctional PBP1a (5). The mechanisms responsible for the diminution of the reactivity with the drugs have been investigated biochemically and structurally to various degrees for the three PBPs. For PBPs from a β-lactam-susceptible strain, the common source has been the laboratory strain R6, which has a minimal inhibitory concentration (MIC) below 0.016 μg/ml for penicillin and cefotaxime. The most thoroughly characterized PBPs from resistant pneumococcus are from the highly resistant strain 5204 isolated in France in 1999, with MICs of 6 μg/ml for penicillin and 12 μg/ml for cefotaxime. The monofunctional transpeptidase PBP2x is the most studied of these PBPs. The role of important amino acid substitutions within the active site of PBP2x has been established by kinetics and structural studies. For example, a mutation two residues downstream of the active site serine found in highly resistant strains was shown to change the orientation of the hydroxyl group of the serine, thereby diminishing its reactivity with β-lactams (9). Two other substitutions common in resistant strains were shown to destabilize the structure of a loop.
that lines the active site (10). A comprehensive study of all the substitutions in the transpeptidase domain of PBP2x from the highly resistant strain 5204 determined that only 6 of the 41 substitutions are important for reducing the reactivity with β-lactams (10).

The monofunctional class B PBP2b has not been as extensively studied, although the crystal structure has been solved for the variants from the susceptible laboratory strain R6 and the clinical resistant strain 5204 (11). Like in PBP2x, a loop forming one side or “lip” of the active site is flexible in the variant from the resistant strain. Sequence comparison identified the T446A substitution within the active site as critical for resistance, and an early biochemical study demonstrated that this mutation severely reduces the reactivity with β-lactams, although no reaction kinetics could be measured (12).

Likewise, the crystal structure of the transpeptidase domain of the class A PBP1a was solved for the susceptible strain R6 and the resistant strain 5204 (13, 14). A mutation adjacent to the active site serine was found to modify its orientation, and the loop forming a lip of the active site was destabilized as in PBP2x and PBP2b by a stretch of four substitutions (14).

β-Lactams owe their tremendous medical success to their structural likeness to the terminal dipeptide D-Ala–D-Ala of the donor stem peptide in the transpeptidation reaction (15). The carbonyl of the β-lactam ring that is attacked by the nucleophilic active site serine is analogous to the carbonyl of the peptide bond that links the two terminal d-alanine residues of the peptidoglycan stem peptide. Because of the similarity between the drugs and physiological substrate, we are facing the following paradox: PBP alterations that affect reaction with β-lactams, as briefly presented above, would be expected to impact negatively on their transpeptidase enzymatic function. To solve this paradox, compensatory mechanisms likely mitigate this problem. Compensation could take place at the cellular level to cope with the effectively reduced enzymatic activity of the PBPs in resistant strains. Alternatively, compensation could occur at the molecular level of the PBPs themselves if the consequences of some substitutions are different on the reactivity with the β-lactams and the transpeptidase activity. The two types of compensatory mechanisms are not mutually exclusive and could operate together.

Investigating cellular mechanisms that could compensate for lower PBP activity is very difficult because the cascade of events that occur in the pneumococcus following β-lactam challenge remains largely mysterious (16–18). Demonstrating the existence of a compensatory mechanism to maintain the enzymatic activity of mosaic PBPs of S. pneumoniae was not possible until recently because there was no assay to evaluate the transpeptidase activity in vitro. This hurdle was removed because the proper peptidoglycan precursor is now available. The nature of the stem pentapeptide of the membrane-linked precursor (lipid II) varies slightly in different bacterial species. In the pneumococcus, the second residue is a γ-D-iso-glutamine. The discovery of the amido transferase enzyme (MurT/GatD) that modifies the second residue γ-D-glutamate into γ-D-iso-glutamine (19, 20) has allowed the preparation of lipid II that can be used by pneumococcal PBPs to synthesize cross-linked peptidoglycan in vitro (21).

To investigate the paradox raised by PBP-based resistance and the structural similarity between β-lactams and the natural substrate of PBPs, we present here a comparison of four variants of PBP2b, (i) from the susceptible laboratory strain R6, (ii) from the clinical resistant strain 5204 (11, 12), (iii) a hybrid form with the N terminus from strain 5204 and the C terminus of strain R6, and (iv) the T446A point mutant (Figs. 1 and 2). 5204-PBP2b carries 56 substitutions, of which 43 are within the transpeptidase domain. Hybrid-PBP2b carries 28 substitutions, including 15 in the transpeptidase domain, and requires further presentation. The hybrid form was not designed but results from the transformation of the R6 strain with the pph2 gene from strain 5204 and selection with piperacillin. S. pneumoniae is naturally competent and readily recombines foreign homologous DNA. An allele conferring a resistance can therefore easily be introduced and selected, provided that flanking regions are provided to allow recombination. With PBP2b, however, the whole gene could not be introduced because recombination repeatedly occurred within the coding region, resulting in a gene with the 5′-region from the resistant strain and the 3′-region retained from the susceptible strain (12, 18, 22). Characterization of the resulting Hybrid-PBP2B might shed light on this incomplete incorporation of the 5204-pph2b allele in the R6 strain. The T446A substitution, which is found in both 5204- and Hybrid-PBP2b, is the most commonly found in resistant strains and can confer some resistance alone (12).

For the first time, both the in vitro transpeptidase activity and the reactivity with a panel of β-lactams were evaluated for a set of PBPs. We uncovered a new level of complexity because the impacts of substitutions on the transpeptidase activity and reactivity with β-lactams are not fully correlated, revealing that some substitutions have compensatory effects that restore enzymatic activity. We also discovered that variants of PBP2b can display β-lactamase activity that may contribute to the resistance to some β-lactams.

**Results**

**Transpeptidase Activity**—To meaningfully compare their transpeptidase activity, it was necessary to use equivalent amounts of the PBP2b variants. Impurities from Triton X-100 and traces of oxidized DTT prevented the accurate determination of the concentrations of the detergent-solubilized full-length proteins from their UV absorbance. Instead, protein amounts were determined in-gel after electrophoresis by quantifying the fluorescence of Bocillin-labeled PBP2b. The truncated soluble forms of PBP2b were used as standards because their concentrations could be determined from their UV absorbance. Care was taken to use a concentration of Bocillin FL and reaction time sufficient to completely label even the variants with lowest β-lactam reactivity. That this was achieved was shown by the fact that the ratio of fluorescence to subsequent Coomassie-staining was similar for all proteins. Because Bocillin binding implies a functional active site, this observation suggests that the proportion of active enzyme was the same in all cases.

Peptidoglycan assembly was monitored using the polyacrylamide gel electrophoretic assay used previously and presented by a scheme in Fig. 3 (21). The lipid II peptidoglycan precursor...
migrates at the front, glycan chains of various length migrate as a smear within the gel, whereas high molecular mass cross-linked peptidoglycan, the product of transpeptidation, remains at the top of the gel. A small amount of dansylated precursor (10%) is incorporated to allow in-gel fluorescence imaging under UV transillumination. The amount of fluorescent precursor is kept to a minimum as it cannot function as acceptor in the transpeptidation because the dansyl group is attached to the lysine side chain that normally takes part in the reaction. PBP2b is a monofunctional transpeptidase that does not cross-link lipid II stem peptides but requires polymerized glycan chains as substrate. Therefore, the S370A-PBP1a point mutant, which is devoid of transpeptidase activity but retains a functional transglycosylase domain, was included to synthesize glycan chains from lipid II.

The transpeptidase activity of the PBP2b variants was assessed by comparing the amount of fluorescence present at the top of the electrophoretic gel (Fig. 4). In the end point experiment shown in Fig. 4A, R6-PBP2b produced more cross-linked peptidoglycan after 16 h of reaction than Hybrid-PBP2b, followed by 5204- and T446A-PBP2b. The important T446A mutation is in orange and denoted by a triangle. The sequence from 5204-PBP2b is in gray, and that of R6-PBP2b is in blue. The sequence of hybrid-PBP2b is colored according to its two sequences of origin, with the junction positioned arbitrarily midway between unambiguous positions. Residues forming the upper lip of the active site (positions 619–629) are in a darker shade.

Assuming a uniform density of cross-links and considering the amount of cross-linked peptidoglycan after 16 h of reaction, the relative activity of the PBP2b variants can be ranked as follows: R6 (100) > Hybrid (22 ± 3) > 5204 (11 ± 2) > T446A (9 ± 2). The numbers represent the amount of cross-linked peptidoglycan relative to that formed by R6-PBP2b. The error is the standard error to the mean of five independent experiments. Although the difference between the amount of cross-linked peptidoglycan produced by 5204- and T446A-PBP2b is small, the latter was smaller in all five experiments. A one-tailed paired \( t \) test of the data gave a \( p \) value of 0.035, indicating that the small difference is significant. A time course is shown in Fig. 4C. The transpeptidase activity of all PBP2b variants were inhibited with 1 mM penicillin G.

The T446A mutant is the least active, despite having a single substitution that is also present in Hybrid- and 5204-PBP2b. This observation implies that some of the substitutions found in the latter sequences are compensatory and increase the transpeptidase activity that is severely impacted by the T446A mutation. Most interestingly, Hybrid-PBP2b, which is a laboratory selected chimeric sequence, has a better enzymatic activity than its parental clinical strain.

Reaction with Bocillin FL and Comparison between Full-length and Soluble Truncated PBP2bs—It is easier to produce and manipulate soluble proteins than detergent-solubilized membrane proteins. Because we aimed to characterize the acylation of PBP2b by several \( \beta \)-lactams, it was highly desirable to use a truncated soluble form of the proteins in the absence of detergent. However, to ensure the relevance of such measure-
ments, we first compared the reactivity of some detergent-solubilized full-length and soluble truncated PBP2bs with Bocillin FL monitored by polyacrylamide gel electrophoresis (Fig. 5).

The reaction of PBPs with β-lactams can be described kinetically with the three-step model (23) (Equation 1). A non-cova-

lent complex \(\text{PBP} \leftrightarrow \text{H} \rightarrow \text{L}\) is formed between the enzyme and the inhibitor \(\text{H} \rightarrow \text{L}\), with the dissociation constant \(K_D\), from which acylation proceeds to form the covalent complex \(\text{PBP} \rightarrow \text{H} \rightarrow \text{L}\) with the rate \(k_2\). \(\text{PBP} \rightarrow \text{H} \rightarrow \text{L}\) is finally hydrolyzed with the rate \(k_3\) to regenerate the enzyme and an inactivated product \(\text{P}\). The rate described by \(k_3\) is usually negligible on the time scale of a bac-

terial generation. The second order rate constant \(k_2/K_D\) is the efficiency of acylation, which allows calculation of the overall acylation rate at a given concentration of antibiotic. Note that the inhibitory potency of a particular β-lactam for a PBP is given by the \(c_{50}\) which is the antibiotic concentration resulting in the inhibition of half the PBP molecules at steady state (i.e. when the acylation and deacylation reactions proceed at the same rate). The value of \(c_{50}\) is equal to the ratio \(k_3/(k_2/K_D)\).

\[
PBP + \beta L \rightleftharpoons PBP-\beta L \rightarrow PBP-\beta L \rightarrow PBP + \beta (Eq. 1)
\]

Purified recombinant PBP2b variants were incubated with various concentration of Bocillin FL (a fluorescently labeled penicillin V) in large excess. After various time intervals, aliquots were withdrawn, and the reaction was stopped by the addition of a denaturing solution (0.2% SDS final) and heating at 100 °C. Samples were analyzed by polyacrylamide gel electrophoresis, which allowed the separation of PBP2b-bound Bocillin from the free form (Fig. 5A). We checked that the inactivation procedure was effective by adding 1 mM Bocillin FL to the denaturing solution. Minimal labeling of PBP2b occurred in this way (Fig. 5A, sample Ctl). The in-gel fluorescence of PBP2b-bound Bocillin was quantified and normalized with respect to the amount of PBP2b in each sample, as quantified after subsequent Coomassie staining. The second order rate constant \(k_2/K_D\) was obtained from global fitting of the time course data at the different drug concentrations using a second order reaction model described in Equation 2 (Fig. 5B). With R6-PBP2b, which is highly reactive, using low drug concentrations in the same range as that of the protein was necessary to measure sufficiently slow reactions. For this reason, we fitted the data without making the usual first order approximation that is appropriate when the drug concentration is in large excess and can be considered constant.
The acylation efficiency of R6-PBP2b by Bocillin FL was found in the range of \(10^4\) M\(^{-1}\) s\(^{-1}\), which is similar to values reported with PBP3 from *Acinetobacter baumanii* and *Pseudomonas aeruginosa* (24). Reaction with 5204-PBP2b was 200-fold slower, whereas it was reduced 1000-fold with Hybrid-PBP2b. It is surprising that Hybrid-PBP2b is even less reactive with Bocillin than 5204-PBP2b, despite the fact that it exhibits a better transpeptidase activity.

No difference of reactivity with Bocillin was found between the full-length and truncated proteins (Fig. 5). It was assumed to be true of all \(-\)lactams and further kinetic data were obtained with the soluble forms only.

In contrast to the other monofunctional PBP2x, the reaction of PBP2b with \(-\)lactams has not been characterized in details, because of the absence of modification of the optical spectra upon acylation by the drugs (12). To palliate this lack of data, we measured the acylation efficiency by several \(-\)lactams by monitoring the chromogenic reaction with nitrocefin. The absorbance spectrum of nitrocefin changes upon opening of the \(-\)lactam ring, and the reactions were followed by the rise in absorbance at 490 nm. With nitrocefin alone, the data collected at different drug concentrations could be globally fitted to a second order reaction model to extract the acylation efficiency \(k_2/K_D\) (Equation 3).
Concentration of the include errors on the concentrations of drugs and proteins. Table 1 are standard errors from the regression, which do not and deacylation rate constant \( k_2 \), while using the value of \( k_2/K_D \) for nitrocefin measured before. This was the case for R6- and T446A-PBP2b with amoxicillin, penicillin G, piperacillin, and cefotaxime, as well as Hybrid-PBP2b with penicillin. No reaction of 5204- and Hybrid-PBP2b was observed with cefotaxime.

Following the first phase of the reaction with 5204-PBP2, a linear increase of the absorbance was observed with amoxicillin, penicillin G, and piperacillin, which was greater than with nitrocefin alone, indicating an additional hydrolysis turnover of these \( \beta \)-lactams. Data could be fitted with a model including the four reactions described in Equations 2–5, while introducing the \( k_2/K_D \) and \( k_3 \) measured independently for nitrocefin.

\[
\text{PBP-\(\beta\)L} \xrightarrow{(k_3/K_D)_\text{NCF}} \text{PBP-NCF} \quad \text{(Eq. 3)}
\]

\[
\text{PBP-NCF} \xrightarrow{(k_3)_\text{NCF}} \text{PBP + P1} \quad \text{(Eq. 4)}
\]

\[
\text{PBP - P1} \xrightarrow{(k_2)_\text{NCF}} \text{PBP - P2} \quad \text{(Eq. 5)}
\]

To obtain the acylation efficiency for other non-chromogenic \( \beta \)-lactams, we performed competition experiments where each drug was added to PBP2b together with nitrocefin. Absorbance data, which measure the disappearance of nitrocefin, were fitted with a model that includes the two competing reactions described in Equations 2 and 3, while using the value of \( k_2/K_D \) for nitrocefin measured before. Figure 6. Reaction of soluble PBP2b variants with nitrocefin, alone or in competition with amoxicillin, at pH 7.5 and 25 °C. The change in absorbance that occurs upon opening of the \( \beta \)-lactam ring of nitrocefin was recorded at 490 nm. A fraction of the data points is shown (between 5 and 20%). The solid lines are global fits to the data using the reaction models described in the text. Rate constants are given in Table 1. A, 18 \( \mu \)M of R6-PBP2b was incubated in the presence of 20 (●), 40 (■), 60 (▲), 80 (○), and 100 (▼) \( \mu \)M of nitrocefin. B, 20 \( \mu \)M of T446A-PBP2b was incubated in the presence of 35 (●), 50 (■), 70 (▲), 85 (○), and 100 (▼) \( \mu \)M of nitrocefin. C, 30 \( \mu \)M of Hybrid-PBP2b and 70 \( \mu \)M of nitrocefin were incubated in the presence of 0 (▼), 15 (●), 30 (▲), 600 (■), and 1200 (○) \( \mu \)M of amoxicillin. D, 20 \( \mu \)M of 5204-PBP2b and 50 \( \mu \)M of nitrocefin were incubated with 0 (▼), 75 (●), 150 (▲), 300 (■), and 600 (○) \( \mu \)M of amoxicillin.

The accuracy of the measured acylation efficiencies is ~20%. With PBP2b, nitrocefin and Bocillin FL are the most reactive drugs of our panel, followed by amoxicillin, piperacillin, and penicillin G. Cefotaxime is much less reactive. R6-PBP2b has an efficiency of acylation by penicillin G of 930 M\(^{-1}\) s\(^{-1}\), which is 100-fold slower than the other monofunctional PBP2x (9). With cefotaxime, R6-PBP2b is 4 \times 10\(^{-2}\)-fold less reactive than PBP2x (9). Qualitatively, the reactivities measured in vitro here and in other studies for R6 PBPs (9, 10, 14) are in agreement with the acylations observed in vivo (18, 26).

The single T446A substitution accounts for most of the diminution in acylation efficiency with a 170–8-fold reduction with nitrocefin or piperacillin, respectively. The additional 55 substitutions in 5204-PBP2b collectively only modestly affect the reactivity, whereas removal of the 27 C-terminal substitutions, as in Hybrid-PBP2b, produces a further 2-fold reduction (Table 1).

Amoxicillin Turnover Evidenced by Further Reaction with Bocillin FL—Given the possible significance for resistance of the deacylation of 5204-PBP2b with several \( \beta \)-lactams and the indirect way it was measured with nitrocefin, we conducted an alternative time course experiments with amoxicillin. Full-length R6- and 5204-PBP2b were first acylated by incubation with amoxicillin at a concentration sufficient to ensure full acylation. Then, at the start of the deacylation time course, a large excess of fluorescent Bocillin FL was added, at a concentration...
at which reaction can be considered instantaneous on the time scale of the experiment. Amoxicillin was still present, but its concentration was much lower than that of Bocillin FL. Reactivation by amoxicillin was therefore negligible, and the time course of labeling by Bocillin FL reflects the deacylation rate of the PBP2b-amoxicillin complex.

A very slow deacylation of R6-PBP2b was detected. After 80 min, less than 25% of the enzyme had been regenerated and able to react with Bocillin FL (Fig. 7A). The data did not allow the determination of a reliable \( k_3 \) but implied a value smaller than \( 6 \times 10^{-6} \) s\(^{-1}\) or a half-life of the acyl-enzyme greater than 3 h. A comparable value of \( 3.8 \times 10^{-3} \) s\(^{-1}\) has been reported for the deacylation of a radioactive penicillin G acyl-enzyme with a soluble form of R6-PBP2b (12). Such rates are negligible on the bacterial generation time scale.

In contrast, a significant deacylation was measured with 5204-PBP2b (Fig. 7, B and C). Nearly complete turnover was observed after 4 min, which yielded the deacylation rate \( k_3 = (10 \pm 1) \times 10^{-3} \) s\(^{-1}\) and a half-life of the inhibitory complex of 70 s. These values are clearly significant for growing bacteria. The value of \( k_3 \) measured for full-length 5204-PBP2b by the time course of reacylation with Bocillin FL is in good agreement with the value of \( 6.2 \times 10^{-5} \) s\(^{-1}\) measured by competition with nitrocefin for the soluble form of the protein (Table 1).

**Thermofluor Assay**—To gain insight on the interactions between the PBP2b variants and the various \( \beta \)-lactams that may account for the different kinetic behavior, we investigated the effect of the acylation on the thermal stability of the soluble proteins. The thermofluor assay measure the fluorescence emitted by a probe (SYPRO® Orange) that binds to hydrophobic surfaces exposed during protein unfolding induced by rising the temperature (27, 28). Fluorescence of the probe is enhanced in the hydrophobic environment. Thermofluor data without or with penicillin or piperacillin are shown in Fig. 8. Three observations can be made from the melting temperatures \( T_{m} \) given in Table 2. First, acylation of R6-PBP2b by penams such as amoxicillin, penicillin G, and piperacillin stabilizes the protein, whereas acylation by the cephalosporin cefotaxime is destabilizing. Second, in the absence of bound antibiotic, R6-PBP2b is thermally less stable than its variants from resistant strains. 5204- and Hybrid-PBP2b have the same higher \( T_{m} \). Third, on the contrary, variants from the resistant strains in their acylated form are less stable than acylated R6-PBP2b. This effect is more important for 5204- than for Hybrid-PBP2b. Note that this does not apply with cefotaxime, which presumably did not acylate the resistance variants at the concentration used (1 mM).

**Discussion**

The properties of the PBP2b variants measured in vitro, both the transpeptidase activity and the reactivity with \( \beta \)-lactams, are consistent with the impossibility to introduce the full PBP2b sequence from a clinical resistant strain (5204) into the susceptible strain R6. Indeed, the Hybrid-PBP2b resulting from the splicing of the N-terminal part of 5204-PBP2b and the C-terminal part of R6-PBP2b has better enzymatic activity and lower drug reactivity than the full 5204-PBP2b. The biochemical properties of Hybrid-PBP2b explain why it is repeatedly selected by \( \beta \)-lactams in laboratory transformation experi-

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**TABLE 1**

| Kinetic parameters of the reaction between \( \beta \)-lactams and PBP2 variants |
|-------------------------|-------------------|------------------|-------------------|
| Nitrocein | R6 | T446A | Hybrid |
| \( k_d/K_i \) \( \text{(M}^{-1} \text{s}^{-1}) \) | 12,700 ± 350 | 74 ± 13 | 230 ± 2 |
| \( k_e \) \( \text{(s}^{-1}) \) | ND | ND | 8 ± 2 | 3.5 × 10⁻⁷ |
| ND | ND | 3.2 × 10⁻⁷ |
| Amoxicillin | 3000 ± 130 | 222 ± 10 | 70 ± 2 |
| PN | ND | ND | 6.2 ± 0.1 | 8.9 × 10⁻⁵ |
| Penicillin G | 930 ± 25 | 33 ± 1 | 35 ± 1 |
| | ND | ND | 2.0 ± 0.05 | 5.7 × 10⁻⁵ |
| | ND | ND | 19 ± 0.3 |
| Pi | 2400 ± 94 | 280 ± 8 | 190 ± 6 |
| Cef | 5 ± 2 | 1.0 ± 0.1 | <10⁻³ |
| Bocillin FL | 13,700 ± 111 | ND | 68 ± 11 |
| | ND | 12 ± 1 |
Transpeptidase Activity and Reactivity of Pneumococcal PBP2b

FIGURE 8. Thermal denaturation of PBP2b variants in apo and acylated forms (Thermofluor). Soluble truncated R6- (A), Hybrid- (B), and 5204-PBP2b (C) were incubated without (continuous line) or with 1 mM penicillin (line with long dashes) or piperacillin (line with short dashes) prior to thermal denaturation in the presence of the fluorescent probe SYPRO® Orange. The first derivative of the fluorescence as a function of increasing temperature is shown.

TABLE 2
Melting temperatures ($T_m$) of PBP2b in the absence or presence of 1 mM β-lactam $T_m$ (°C) were determined as the minimum of the first derivative of the melting curves. The mean of two experiments is given. Standard deviation was at most 0.5 °C.

| Variant       | AMX  | PEN  | PIP  | CTX  |
|---------------|------|------|------|------|
| None          | 49.5 | 59.0 | 57.5 | 59.5 | 46.0 |
| R6-PBP2b      | 52.0 | 53.0 | 45.5 | 51.5 | 52.0 |
| 5204-PBP2b    | 52.0 | 56.0 | 49.0 | 54.5 | 52.5 |
| Hybrid-PBP2b  |      |      |      |      |      |

Note that if the $c_{so}$ is considered, 5204-PBP2b is slightly less susceptible to β-lactam than Hybrid-PBP2b. The level of transpeptidase activity would therefore be the main factor favoring the selection of Hybrid-PBP2b instead of 5204-PBP2b by β-lactams.

Examination of the modeled structure of Hybrid-PBP2b shows that the missing substitutions compared with 5204-PBP2b are distributed in the “upper lip” of the active site and at the “back” of the protein (Fig. 2). The “upper lip” substitutions A619G, D625G, Q628E, and T630N are the most likely to cause the severe diminution of transpeptidase activity of 5204-PBP2b because of their proximity to the active site residues at the entrance of the cleft. The region spanning these substitutions forms a loop connecting strands β3 and β4 that is mobile and not visible in the crystal structure of 5204-PBP2b (11). However, the role of these substitutions in reducing the reactivity with the drugs is likely limited, contrary to what was thought previously (11), because Hybrid-PBP2b with a probable “stiff upper lip” is even less reactive with β-lactams than the more “relaxed” 5204-PBP2b. In contrast, substitutions in the lower lip of the active site S412P, N422Y, T426Q, and Q427L likely contribute the most to the decrease in reactivity with the drugs. Interestingly and in agreement with the PBP2b observations, substitutions in the lower lip of PBP2e were found to reduce β-lactam reactivity, whereas substitutions in the β3-β4 upper lip had no effect or even increased the reactivity with the drug (10). Note that the β3-β4 upper lip in PBP2e protrudes in the solvent and makes no contact with a bound cefuroxime molecule (29). The lower lip on the contrary is held tightly against the active site and contacts the bound antibiotic. Unfortunately, no crystal structure of PBP2b with bound antibiotic and no data on the transpeptidase activity of PBP2e variants are available to further compare both proteins.

The decylation rate ($k_d$), which may also contribute to the resistance, is particularly elevated with 5204-PBP2b. It is possible that the flexibility of the upper lip and the replacement of Ala$^{519}$ by a glycine very close to the active site Ser$^{386}$ allow easier access of water to the acyl-enzyme bond.

The T446A substitution is the most important to reduce the acylation by β-lactams (Table 1). Collectively, the 55 other substitutions in 5204-PBP2b cause only a modest additional reduction of the acylation rate (Table 1). Among these, the 27 substitutions at the C terminus must collectively have an opposite effect resulting from the homologous recombination of long stretches of DNA. What is revealed in the present study is that some substitutions are compensatory and participate to the resistance not by restricting the reaction with the drugs but by restoring an acceptable level of physiological enzymatic activity, which would otherwise be severely impacted by mutations that hamper the reaction with β-lactams.

The findings above raise the question of why is 5204-PBP2b present in a clinical resistant strain, since a sequence with fewer substitutions appears biochemically superior? A clue can be found in a laboratory study of the transfer of β-lactam resistance from Streptococcus mitis to S. pneumoniae (22). The full sequence of PBP2b from a resistant S. mitis strain, including the 16 substitutions at the C terminus following position 590, could be transformed and selected by benzylpenicillin in S. pneu-
moniae, only if the mosaic variant of the murM gene from the S. mitis was also introduced.

The murMN operon encodes two cytoplasmic enzymes responsible for the “branching” of the stem peptides of the peptidoglycan (17). The precursor of the peptidoglycan is synthesized with a pentapeptide, the third residue of which is a lysine in pneumococcus. The free amine of the lysine side chain of the donor peptide forms the peptide bond with the fourth residue (D-Ala) carboxyl group of the donor peptide to cross-link glycan chains. Alternatively, MurM and MurN can add successively two residues onto the lysine side chain to produce branched stem peptides with additional Ser-Ala or Ala-Ala dipeptides. It is then the N terminus of the dipeptide on the acceptor that forms the cross-linking peptide bond with the donor stem peptide. Strain R6 and its parental strain R36A were found to have ~45 and 36% branched peptides, respectively (30, 31). In clinical resistant strains that express a MurM variant from a mosaic allele, the proportion of branched stem peptide is increased up to 85% (31), because of a greater enzymatic activity of the MurM variant (32). In the absence of MurM, no branched peptides are detected, and the resistance is abolished despite the presence of altered PBPs (31).

To explain the special relationship between PBP2b and MurM in β-lactam resistance, two plausible explanations can be considered. More branched peptides may have a compensatory role (i) on the enzymatic activity of PBP2b or (ii) on the cell wall metabolism. (i) Branched stem peptides may be better substrates for altered PBP2bs than the linear form, either as donor or acceptor, thus compensating the decreased transpeptidase activity. It is highly desirable to test this hypothesis in vitro, but suitable substrates are unfortunately not available in sufficient amounts. However, the fact that in the absence of antibiotic challenge, a strain devoid of MurM and branched peptide grows normally with altered PBPs (31) argues strongly against PBP2b simply discriminating between different peptide substrates. (ii) PBP2b participates in the elongation of the ovoid pneumococcal cell (33). Addition of peptidoglycan material to the cell wall during elongation requires concomitant opening of the existing peptidoglycan to permit insertion. A peptidoglycan lytic transglycosylase has recently been identified that likely works in concert with PBP2b during elongation of the pneumococcus (34). A decreased transpeptidase activity of PBP2b might create an imbalance with the degradative activity of its associated lytic transglycosylase that may trigger full lysis with the participation of other peptidoglycan hydrolases such as the major autolysin LytA. The greater proportion of branched peptides caused by mosaic MurM may protect against lysis. Indeed, elevated level of branched peptides caused by expression of a mosaic hyperactive MurM also protects against cell lysis normally induced by non-β-lactam antibiotics (17). Also, gradual depletion of PBP2b was found to be tolerated to a large extent due to an increased level of branched peptides. In the absence of MurM, and consequently of branched peptides, depletion of PBP2b was much less tolerated (35).

In conclusion, we propose that the deleterious effect of mutations that greatly diminish the reactivity of PBP2b with β-lactams, such as the T446A, is compensated by a combination of compensatory substitutions within PBP2b that maintain a minimal necessary transpeptidase activity, together with an alteration of the peptidoglycan composition that compensate for the reduced enzymatic activity.

The reactivity with Bocillin FL of the extracellular domain of PBP2b was found to be the same as that of the full-length membrane proteins (Fig. 5). In contrast, the transpeptidase activity was severely impacted by the truncation of the transmembrane segment. Residual activity was detected only with soluble R6-PBP2b (not shown). An influence of the membrane anchor on the active site cannot be ruled out but is unlikely in the light of the reaction with β-lactams. Rather, the glycan chains and the full-length enzymes may be brought into close proximity by their common anchoring into detergent micelles. This efficient concentration effect is lost by truncation of the transmembrane segment. In any case, the in vitro transpeptidase activities reported here and previously are extremely weak and far from realistic physiologic rates. In the future, it will be necessary to study monofunctional PBPs in a membrane environment and/or in the presence of their protein partners (36), in particular SEDS proteins that act as transglycosylases (37).

Additional comments can be made by comparing of the reaction rates with different β-lactams. Cefotaxime does not react significantly with PBP2b, as reported previously (12, 26). This property is not general to cephalosporins, however, because nitrocefin was the most reactive drug of our panel. Examination of the structure of the different cephalosporins that fail to react with PBP2b (26, 38) suggests that the nature of the R2 substituent is important. Bulky and planar substituent in position R2 appear to prevent recognition by PBP2b, whereas smaller and tetrahedral substituent seem compatible. The unfavorable interaction between cefotaxime and PBP2b seems to persist once the β-lactam ring is open and the acyl-enzyme complex is formed, because the thermal stability of the protein is diminished (Table 2).

It is remarkable that substitutions that lead to the thermal stabilization of the apo form of 5204- and Hybrid-PBP2b compared with the apo form of R6-PBP2b have the opposite effect on the acylated forms (Table 2). In the variants from the resistant strains, the covalently bound drugs must make unfavorable interactions with the proteins. Although the “open” forms of the antibiotics that are bound in the active site are different from the original β-lactams, they likely retain interactions that also occur in the precacylation complex or the transition state and may be relevant to the acylation and deacylation kinetics.

Other than nitrocefin, which is not in clinical use, the most important diminution of reactivity measured for 5204-PBP2b compared with R6-PBP2b was with amoxicillin. It is also with amoxicillin that the greatest β-lactamase activity was measured for 5204-PBP2b (Table 1). Amoxicillin has long been a widely prescribed treatment for otitis media and sore throat, even when bacterial infection is not clearly diagnosed (39, 40). The nasopharyngeal flora, including S. pneumoniae has therefore been subjected to a severe selection pressure by amoxicillin, and it is possible that the substitutions found in 5204-PBP2b reflects the clinical practice. Strain 5204 exhibits the highest level of amoxicillin resistance (MIC 6 μg ml⁻¹) in a panel of French clinical isolates, together with isolate 5268, which has a similar PBP2b sequence (8).
The surprising finding that a laboratory selected PBP2b (Hybrid) performs better in vitro than its clinical parental variant (5204) indicates that β-lactam resistance in pneumococcus is a complex process that is not fully described by biochemical reconstitution experiments. Resistance to β-lactams must be considered as a possible physiological response with compensatory mechanisms at play. Such mechanisms should be studied in the future because they may offer novel therapeutic options.

**Experimental Procedures**

**Chemicals**—Lyophilized nitrocefin was purchased from Oxoid (catalog no. SR0112C), reconstituted at 2 mM in DMSO, and stored at −20 °C. Bocillin FL was from Thermofisher Scientific (ref: B13233). Stock solutions were prepared in 50 mM HEPES, pH 7.5, 100 mM NaCl, 10 mM MgCl2, at 50 mM and stored at −20 °C. Amoxicillin (catalog no. A8523), cefotaxime (catalog no. C7912), penicillin G (catalog no. P3032), and pipercillin (catalog no. P8396) were from Sigma and solubilized at 100 mM in the same buffer and stored at −20 °C, except amoxicillin, which was solubilized with the addition of 1% NH4OH and kept for a maximum of 15 days at −80 °C.

Lys-containing lipid II and dansylated lipid II were prepared as described previously (41, 42). Amidation of lipid II and subsequent purification was performed as described previously (21).

**Structure Modeling**—The structural model of Hybrid-PBP2b was constructed by superposing the crystal structures of R6-PBP2b (PDB code 2WAF) from residues 518–682 onto that of 5204-PBP2b (PDB code 2WAD) using LSQ Superpose in COOT (43) from the CCP4 program suite (44). The model structure was then created by replacing residues 518–682 in 5204-PBP2b by those from R6-PBP2b. Because examination of the model found no clash or other anomaly, a simple structure idealization was performed with REFMAC (45).

**Expression Plasmids**—Full-length R6-PBP2b with a C-terminal Strep-tag was expressed from the modified pET-30 plasmid described before (21). The gene encoding Hybrid-PBP2b was PCR-amplified from genomic DNA from laboratory strain R6 ΔlytA pbb2s (18) using primers Nde2b and 2bStopNheSpeBam (21). The gene coding for 5204-PBP2b was amplified from DNA of the clinical strain 5204 (8) using primers Nde2b and 2b5204Bam (GGTCGACGATCATTACATGTGATTGTTGG). A NdeI internal site was silently mutated by PCR using primers 2b5204mutNdeFW (GTATAAATTGGCGTATGGAATCATTACATGTGATTGTTGG) and 2b5204mutNdeRV (GAAAAGATCGATACGCACATTATACAT). The resulting PBP2b coding genes were introduced as NdeI/BamHI fragments into a modified pET-30 expression plasmid following PCR amplification from DNA of the clinical strain 5204 (8) using primers Nde2b and 2bStopNheSpeBam to replace the original pET-30 plasmid, which encodes a C-terminal Strep-tag.

N-terminally truncated versions of the three PBP2b variants starting at residue Met39 were subcloned into the same modified pET-30 expression plasmid following PCR amplification from the parent plasmids encoding the full-length proteins, using primers Nde2bstar (GAGAATTCATATGCGACGGTTTGGAA-CGAAGGATTTCAG) and 2bStopNheSpeBam to replace the original pET-30 plasmid. All final coding sequences were checked.

**Protein Expression and Purification**—Full-length S370A-PBP1a with an N-terminal His tag was produced and purified as described (46). The preparation of PBP2bs was as detailed in the supporting information of previous work (21). Briefly, all pneumococcal proteins were produced in Escherichia coli BL21 (DE3) star. Full-length variants were purified from Triton X-100-solubilized membranes, whereas soluble truncated forms were purified from the soluble fraction of cell lysates. All variants harboring a C-terminal Strep-tag were purified by Strep-Tactin affinity chromatography in 50 mM HEPES, pH 7.5, 150 mM NaCl, 10 mM MgCl2. Proteins were eluted with 2.5 mM desthiobiotin. Full-length protein solutions also contained 0.02% Triton X-100 and 1 mM DTT because the PBP2bs contain a cysteine residue near the N terminus.

**Peptidoglycan Synthesis**—Unlabeled iso-glutamine-containing lipid II and dansylated lipid II in CHCl3-methanol (1:1) were mixed in a 10:1 ratio and dried under nitrogen flow. The dried lipid II mix was then redissolved to a final concentration of 55 μM in the reaction mix typically containing 50 mM HEPES, pH 7.5, 150 mM NaCl, 10 mM MgCl2, 25% (w/v) DMSO, and 0.02% (w/v) Triton X-100, and the PBPs were investigated. The concentrations of buffer, salts, and particularly of detergent contributed by the protein stocks were taken into account. The concentrations of S370A-PBP1a and PBP2bs were 0.3 and 1 μM, respectively. Penicillin G (1 mM) was included as required. The reactions were left to proceed overnight at 30 °C unless otherwise stated. For time course experiments, aliquots were withdrawn after various time intervals, and the reaction was stopped by the addition of penicillin G (1 mM) and monomycin (Flavomycin, Hoechst, 0.5 mM). Samples were analyzed by SDS-PAGE (21) and visualized with blue transillumination using the GelDoc ChemiDoc MP imager (Bio-Rad). Quantification was performed on unsaturated images using the Image Lab software (Bio-Rad).

**Kinetics of PBP2b Reaction with Bocillin FL**—Full-length and truncated soluble variants of PBP2b were incubated in the presence of various concentrations of Bocillin FL at 20 °C in 50 mM HEPES, pH 7.5, 150 mM NaCl, 10 mM MgCl2, and 0.02% (w/v) Triton X-100 for the full-length proteins. The ranges of Bocillin FL concentrations were 5–20 μM for R6-PBP2bs or 0.25–1 mM for Hybrid- and 5204-PBP2bs. Aliquots were withdrawn after various time intervals, and the reaction was stopped by the addition of loading buffer (0.2% (w/v) SDS final concentration) and a heat shock (100 °C, 5 min). The fluorescence of Bocillin-PBP2b complexes was recorded under blue transillumination. Fluorescence data were corrected for the relative amount of PBP2b in each sample after subsequent quantification of Coomassie-stained protein bands. Imaging and quantification were performed using a GelDoc ChemiDoc MP imager and the Image Lab software (Bio-Rad). The data were fitted globally with the DynaFit software (47) (Biokine) using a second order reaction model.

**A moxicillin Deacetylation Kinetics**—Full-length R6- or 5204-PBP2b were incubated at 1 μM for 15 min at 20 °C with 100 μM amoxicillin in 50 mM HEPES, pH 7.5, 150 mM NaCl, 10 mM MgCl2, and 0.02% (w/v) Triton X-100, prior to addition of 1 mM Bocillin FL at time 0 of the deacylation time course. After various time intervals, the reaction was stopped, and samples were analyzed as above. The data for the fraction (f) of Bocillin-labeled protein were fitted to a first order kinetic (f = 1 − exp(−k,t)) using the Kaleidagraph software (Synergy).
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Kinetics of PBP2b Reaction with Nitrocefin and Other β-Lactams—The different soluble PBP2b variants were incubated with either various concentrations of nitrocefin or a fixed concentration nitrocefin and varying concentrations of other β-lactams, in 50 mM HEPES, pH 7.5, 150 mM NaCl, 10 mM MgCl₂. Absorbance at 490 nm was recorded at 25°C in an OPTIMA FLUOstar plate reader where 150 μM of protein solution were automatically injected onto 100 μl of antibiotic solution in a Greiner 96-well plate. PBP2b and drug concentrations were as follows. With 18 μM R6-PBP2b, nitrocefin was varied alone from 20 to 100 μM, whereas it was kept at 10 μM while amoxicillin ranged from 0 to 25 μM, penicillin G from 0 to 800 μM, pipercillin from 0 to 80 μM, and cefotaxime from 0 to 5 mM. With 20 μM 5204-PBP2b, nitrocefin was varied alone from 100 to 500 μM, whereas it was kept at 150 μM while amoxicillin and penicillin G ranged from 0 to 2.5 mM, and pipercillin from 0 to 800 μM. With 30 μM Hybrid-PBP2b, nitrocefin was varied alone from 200 to 800 μM, whereas it was varied from 70 μM while amoxicillin ranged from 0 to 1.2 mM, penicillin G from 0 to 2 mM and pipercillin from 0 to 150 μM. With 20 μM T446A-PBP2b, nitrocefin was varied alone from 70 to 200 μM, whereas it was kept at 35 μM while amoxicillin ranged from 0 to 625 μM, penicillin G from 0 to 2.4 mM, pipercillin from 0 to 300 μM, and cefotaxime from 0 to 4.5 mM. For each PBP2b variant, the data for a range of β-lactam concentrations were fitted globally with the DynaFit software (47) (Biokine) using second order reaction models for the acylations and first order model for the deacylation, as described under “Results.”

Thermofluor Assay—Soluble PBP2b variants at a final concentration of 0.5 mg/ml (~6 μM) were incubated with or without 1 mM of amoxicillin, penicillin G, pipercillin, or cefotaxime in 50 mM HEPES, pH 7.5, 150 mM NaCl, and 10 mM MgCl₂ for 5 min at room temperature prior to the addition of SYPRO® Orange (12× final concentration) on ice. Fluorescence melting curves were recorded in the SYBR channel in a CFX Connect instrument (Bio-Rad) with a 20–100°C temperature ramp (0.5°C steps of 7.2 s) and analyzed with the CFX Manager software.

Author Contributions—P. C. and A. Z. designed and conducted most of the experiments and analyzed the results. E. B. synthesized lipid II. D. I. R. purified PBP1a. M. D. performed the thermal shift assays. C. C.-M. modeled the protein structure. A. Z. wrote the manuscript.

Acknowledgments—We thank Anne-Marie Villard for performing site-directed mutagenesis and Thierry Vernet and Max Maurin for critical discussion of the results.

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Substitutions in PBP2b from β-Lactam-resistant Streptococcus pneumoniae Have Different Effects on Enzymatic Activity and Drug Reactivity
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J. Biol. Chem. 2017, 292:2854-2865.
doi: 10.1074/jbc.M116.764696 originally published online January 6, 2017

Access the most updated version of this article at doi: 10.1074/jbc.M116.764696

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