

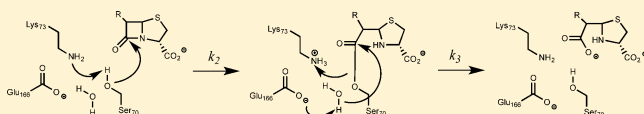
Kinetic Characterization of Hydrolysis of Nitrocefin, Cefoxitin, and Meropenem by β -Lactamase from *Mycobacterium tuberculosis*

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S Supporting Information

ABSTRACT: The constitutively expressed, chromosomally encoded β -lactamase (BlaC) is the enzyme responsible for the intrinsic resistance to β -lactam antibiotics in *Mycobacterium tuberculosis*. Previous studies from this laboratory have shown that the enzyme exhibits an extended-spectrum phenotype, with very high levels of penicillinase and cephalosporinase activity, as well as weak carbapenemase activity [Tremblay, L. W., et al. (2008) *Biochemistry* 47, 5312–5316]. In this report, we have determined the pH dependence of the kinetic parameters, revealing that the maximal velocity depends on the ionization state of two groups: a general base exhibiting a pK value of 4.5 and a general acid exhibiting a pK value of 7.8. Having defined a region where the kinetic parameters are pH-independent (pH 6.5), we determined solvent kinetic isotope effects (SKIEs) for three substrates whose k_{cat} values differ by 5.5 orders of magnitude. Nitrocefin is a highly activated, chromogenic cephalosporin derivative that exhibits steady-state solvent kinetic isotope effects of 1.4 on both V and V/K . Cefoxitin is a slower cephalosporin derivative that exhibits a large SKIE on V of 3.9 but a small SKIE of 1.8 on V/K in steady-state experiments. Pre-steady-state, stopped-flow experiments with cefoxitin revealed a burst of β -lactam ring opening with associated SKIE values of 1.6 on the acylation step and 3.4 on the deacylation step. Meropenem is an extremely slow substrate for BlaC and exhibits burst kinetics in the steady-state experiments. SKIE determinations with meropenem revealed large SKIEs on both the acylation and deacylation steps of 3.8 and 4.0, respectively. Proton inventories in all cases were linear, indicating the participation of a single solvent-derived proton in the chemical step responsible for the SKIE. The rate-limiting steps for β -lactam hydrolysis of these substrates are analyzed, and the chemical steps responsible for the observed SKIE are discussed.



One in three people is infected with *Mycobacterium tuberculosis* worldwide. It is the leading killer of people living with HIV, causing one-quarter of all deaths. Furthermore, the emergence of multidrug resistant and extensively drug resistant strains of *M. tuberculosis* poses a serious problem for cost-effective treatment therapies.¹ The intrinsic resistance of *M. tuberculosis* to β -lactams is attributed to the genomically encoded, Ambler class A β -lactamase (BlaC). β -Lactam antibiotics irreversibly bind to the D,D-transpeptidase domains of penicillin-binding proteins (PBPs) by structurally mimicking the D-Ala-D-Ala C-terminal dipeptide of the pentapeptide of the cell wall peptidoglycan.² In bacteria, β -lactams acylate the active site serine of PBPs, forming stable inactive acyl-enzymes.⁵

The catalytic mechanism of BlaC relies on three highly conserved active site residues, Lys73 and Glu166, which are involved in the activation of the acylating nucleophile Ser70 and the activation of the active site water molecule for deacylation, respectively.¹ Although the class A β -lactamase acylation mechanism is still being debated,⁷ BlaC site-directed mutagenesis and crystallographic data with the cephalosporin cefamandole strongly suggest that Lys73 is involved in the activation of Ser70 for acylation and Glu166 in activating the conserved water molecule for deacylation. The Lys73Ala mutant form of BlaC permitted the structural identification of the Michaelis complex but has no catalytic activity, indicating Lys73 is essential for the acylation step. The Glu166Ala mutant form of BlaC similarly has no hydrolytic activity and forms a

stable acyl-enzyme complex that can be structurally characterized, suggesting that Glu166 is required for deacylation and product release.⁶

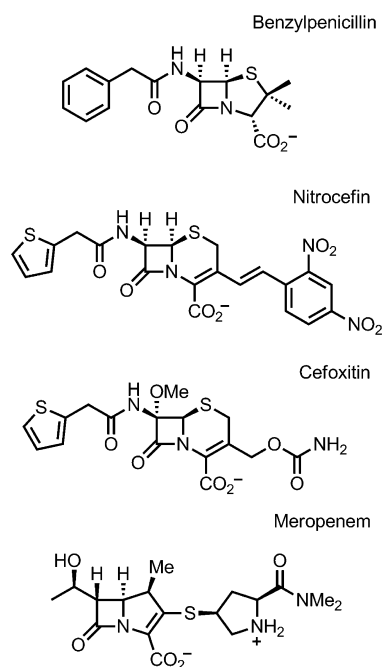
Previously, BlaC has been shown to rapidly hydrolyze penicillin and cephalosporin classes of β -lactams, as well as exhibiting weak carbapenemase activity.² While there has been decades of work in characterizing the mechanism of class A β -lactamases in Gram-negative and Gram-positive bacteria,^{8–12} no extensive studies have been performed to elucidate the rate-limiting steps in the reaction catalyzed by BlaC in *M. tuberculosis*. In this study, we have performed steady-state and pre-steady-state studies to measure solvent kinetic isotope effects on β -lactams that vary in catalytic efficiency. Cephalosporins, nitrocefin and cefoxitin (Scheme 1), are rapidly hydrolyzed by BlaC, whereas the carbapenem meropenem is hydrolyzed extremely slowly. Our comparative kinetic analysis indicates that slow substrates are highly solvent sensitive for both acylation and deacylation reactions; intermediate substrates have a small SKIE on acylation and a larger SKIE on deacylation, and the fast substrates exhibit a small SKIE on acylation only. These findings provide greater mechanistic insight into the rate-limiting steps and chemical

Received: February 12, 2013

Revised: May 8, 2013

Published: May 14, 2013

Scheme 1. Structures of Benzylpenicillin, Nitrocefin, Cefoxitin, and Meropenem



mechanism of the BlaC-catalyzed hydrolysis of β -lactam substrates.

MATERIALS AND METHODS

Materials. All chemicals were of analytical or reagent grade and were used without further purification; 99.9% deuterated water was from Cambridge Isotope Laboratories. The remaining chemicals were purchased from Sigma-Aldrich.

Production and Purification of BlaC. The *blaC* gene was amplified from genomic DNA and cloned into pET28a(+) using NdeI and HindIII, as previously described.¹³ The expression and purification of BlaC followed the protocol previously reported.² On the basis of the stoichiometric inhibition of BlaC by clavulanate,² the active site concentration of BlaC was determined by clavulanate titration.

pH–Rate Profiles. The pH dependence of the kinetic parameters was determined by measuring initial rates at varying concentrations of cefoxitin or nitrocefin. The experiments were conducted at 25 °C in the following buffers (each at 100 mM) at the indicated pH values: sodium acetate (pH 4.0–5.5), MES (pH 5.5–6.7), HEPES (pH 6.7–7.8), and TAPS (pH 7.8–8.7). The resulting k_{cat} data were fit to eq 1 to obtain the $\text{p}K_{\text{a}}$, the negative log of the acid dissociation constant, and the $\text{p}K_{\text{b}}$, the negative log of the base dissociation constant

$$\log k_{\text{cat}} = \log [c / (1 + 10^{\text{p}K_{\text{b}} - \text{pH}} + 10^{\text{pH} - \text{p}K_{\text{a}}})] \quad (1)$$

where c is the pH-independent plateau value. The $k_{\text{cat}}/K_{\text{m}}$ data were fit to eq 2, which describes the pH dependence of a single group whose ionization decreases $k_{\text{cat}}/K_{\text{m}}$ at high pH values.

$$\log k_{\text{cat}}/K_{\text{m}} = \log [c / (1 + 10^{\text{pH} - \text{p}K_{\text{a}}})] \quad (2)$$

Steady-State Kinetics. The steady-state rate of hydrolysis of β -lactam substrates was monitored as a decrease in the absorbance in the UV region, as previously described.¹⁴ Assays using the chromogenic substrate nitrocefin were performed at 486 nm ($\Delta\epsilon = 20500 \text{ M}^{-1} \text{ cm}^{-1}$). Assays using cefoxitin were

performed at 270 nm ($\Delta\epsilon = 8380 \text{ M}^{-1} \text{ cm}^{-1}$). All assays were performed at 25 °C in 100 mM MES/NaCl (pH 6.3). Reactions were initiated by addition of enzyme at concentrations varying from 2 to 300 nM, depending on the substrate used. Initial velocity kinetic data were fit to eq 3

$$\nu = (VS)/(K_{\text{m}} + S) \quad (3)$$

where ν is the initial velocity, V is the maximal velocity, and K_{m} is the Michaelis constant for substrate S .

Solvent Kinetic Isotope Effects. Steady-state solvent kinetic isotope effects were measured at varying concentrations of cefoxitin or nitrocefin and globally fit to eq 4

$$\nu = \frac{VS}{K_{\text{m}}\{1 + F_1(E_k) + S[1 + F_1(E_v)]\}} \quad (4)$$

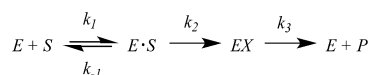
where ν is the initial velocity, V is the maximal velocity, K_{m} is the Michaelis constant for substrate S , E_k is the SKIE on V/K (–1), E_v is the SKIE on V (–1), and $F_1 = 0$ for H_2O and $F_1 = 0.95$ for D_2O . Viscosity effects on V or V/K were evaluated by comparing rates obtained in H_2O or 9% (w/w) glycerol, which mimics the viscosity increase caused by D_2O ($\eta_{\text{r}} = 1.24$).¹⁵ Proton inventories were determined at saturating concentrations of cefoxitin and nitrocefin at 10% volume increments of D_2O from 0 to 90%. SKIEs for steady-state and proton inventory are not reported for meropenem hydrolysis because of a very low K_{m} value, a very low k_{cat} value, and a small extinction coefficient for hydrolysis.

Pre-Steady-State Kinetics. Prior to pre-steady-state kinetic studies, saturation assays were performed to confirm that a sufficient amount of substrate was used to produce pseudo-first-order conditions. A large excess of substrate drives the enzyme to the Michaelis complex form by significantly increasing the rate of binding ($k_1[S]$) so that the acylation (k_2) and deacylation (k_3) rates can be accurately determined.

Solvent KIEs were measured under multiple-turnover conditions. Solvent kinetic isotope effects were measured in an SX-20 (Applied Photophysics) stopped-flow, rapid-mixing spectrophotometer in the absorbance monitoring mode in either H_2O or 95% D_2O . Experiments were performed under pseudo-first-order conditions with at least 50-fold excess of substrate over limiting BlaC with a fixed concentration of 13–27 μM , depending on the substrate used. Because of the absorbance detection limit of 2 AU, assays for cefoxitin and meropenem were not conducted at the wavelength of their maximal extinction coefficients but at 293 nm ($\Delta\epsilon_{\text{app}} = 1960 \text{ M}^{-1} \text{ cm}^{-1}$) and 317 nm ($\Delta\epsilon_{\text{app}} = 4700 \text{ M}^{-1} \text{ cm}^{-1}$), respectively, to obtain rates for k_2 and k_3 . Time courses ranged from 5 to 500 s depending on the substrate for the full observation of the burst and linear phases. Raw absorbance values were divided by the final enzyme concentration to accommodate fitting. Pre-steady-state kinetic studies were not performed with nitrocefin because the k_{cat} value for hydrolysis is too high to observe a burst phase (the acylation rate is faster than the 0.5 ms dead time of the spectrophotometer).

Data Analysis. All data were fit to the appropriate equations using the nonlinear regression function of SigmaPlot 11 (SPSS, Inc.). The general mechanism of BlaC hydrolysis can be modeled as shown in Scheme 2, where k_1 and k_{-1} represent the rates of reversible binding of the substrate to and dissociation of the substrate from BlaC, respectively, k_2 represents the rate of acylation of enzyme BlaC by attack on the β -lactam ring, and k_3 represents the rate of deacylation or hydrolysis of the BlaC– β -lactam adduct. Using this model, rate

Scheme 2



constants for k_2 and k_3 were determined by fitting burst kinetic traces to eq 5

$$\frac{[P]}{[E]_0} = A_0(1 - e^{-k_{\text{obs}}t}) + k_{\text{cat}}t \quad (5)$$

where A_0 is the burst amplitude, $[E]_0$ is the final enzyme concentration, k_{obs} is the overall rate constant, and t is the time¹⁷ when the binding step is not rate-limiting with respect to the overall rate. The variables can be expressed in terms of k_2 and k_3 and substituted into eq 5 to produce eq 6 for fitting the progress curves. Estimated values for k_2 and k_3 from the fitting were verified by the apparent A_0 and k_{cat} values, which are comparable to steady-state turnover numbers (Table S1 of the Supporting Information).

$$A_0 = \left(\frac{k_2}{k_2 + k_3} \right)^2 \quad k_{\text{obs}} = k_2 + k_3 \quad k_{\text{cat}} = \frac{k_2 k_3}{k_2 + k_3}$$

$$\frac{[P]}{[E]_0} = \left(\frac{k_2}{k_2 + k_3} \right)^2 [1 - e^{-(k_2 + k_3)t}] + \left(\frac{k_2 k_3}{k_2 + k_3} \right) t \quad (6)$$

RESULTS AND DISCUSSION

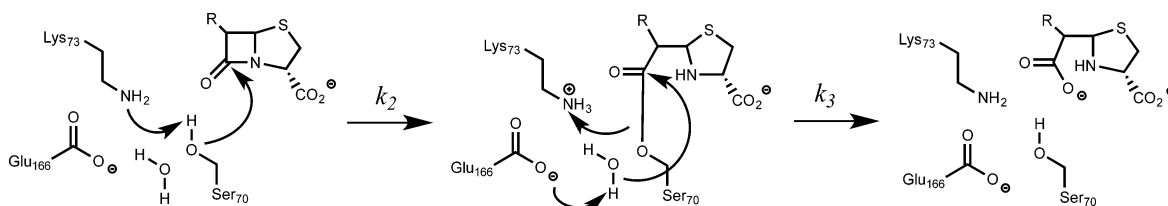
BlaC is the chromosomally encoded β -lactamase in *M. tuberculosis* originally identified after the genome sequence was published in 1998.¹⁸ The full-length enzyme contains an N-terminal 40-amino acid membrane insertion sequence that allows for its export into the periplasm where peptidoglycan cross-linking occurs and anchoring in the outer aspect of the inner membrane. All mechanistic and structural studies have been performed with a soluble construct containing an N-terminal truncation of the first 40 residues, as first reported for the structure of the enzyme in 2006.¹³ Using this construct, it was shown that BlaC is an extended-spectrum β -lactamase (ESBL) with extremely broad substrate specificity for penicillins, cephalosporins, and even carbapenems,³ β -lactams specifically designed to be resistant to hydrolysis by ESBLs.

BlaC is included in the class A Ambler classification of β -lactamases, based on the similarity of its sequence with those of others in this class.¹⁹ These enzymes all contain a conserved triad of catalytic residues, including Ser70, Lys73, and Glu166 (Ambler numbering is used throughout rather than actual sequence numbers). These residues are proposed to act as the nucleophiles responsible for β -lactam ring opening, and the

accessory general acid and base. Mechanistically, they are similar to serine proteases and use a two-step reaction in which the enzyme is first acylated by the putative nucleophilic attack of Ser70 on the β -lactam ring with formation of a transient covalent ester bond, followed by hydrolysis of this bond by the attack of water on the scissile bond (deacylation).

Although the class A β -lactamases have been extensively studied, the assigned roles of the auxiliary general acid/base to either Lys73 or Glu166 have been disputed. Initially, Glu166 was thought to serve as the general base in Ser70 activation. However, an ultra-high-resolution crystallographic study of TEM-1 suggested that Glu166 was protonated at pH 8.0 and thus served the role of general acid.²⁰ Alternatively, *ab initio* quantum mechanics/molecular mechanics calculations suggested that both Lys73 and Glu166 participate in Ser70 activation.⁷ Previous studies from this lab have clarified the role of each residue using site-directed mutagenesis and X-ray crystallography.⁶ The K73A mutant could not be acylated by cefamandole, and the structure of bound cefamandole in the active site revealed a preacylation, noncovalent Michaelis complex. The E166A mutant was crystallized as a covalent, acylated complex with bound cefamandole. Together, these results argue persuasively for the role of Lys73 as the acylation general base and for Glu166 as the deacylation base, as depicted in Scheme 3. To more completely characterize their respective roles in catalysis, we determined the pH dependence of the kinetic parameters for two different cephalosporin substrates.

pH–Rate Profiles. Nitrocefin was developed to be an extremely active, highly chromogenic general substrate for β -lactamases.^{21–23} It contains a vinyl dinitrobenzene substituent attached to the C3 position of the cephalosporin nucleus, providing strong resonance electron withdrawal from the susceptible β -lactam bond. It is a very efficient substrate for BlaC, exhibiting a k_{cat}/K_m value of $\sim 10^5 \text{ M}^{-1} \text{ s}^{-1}$ (Table 1), a value close to the diffusion limit.² Determination of the pH–rate dependence allows for the identification of any ionizable groups involved in the reaction. The pH dependencies of the kinetic parameters for nitrocefin are shown in Figure 1A, where the points are the experimental data and the smooth curves are the fits of the data to eq 1 (k_{cat} ; upper) or eq 2 (k_{cat}/K_m ; lower). The maximal velocity changes as a result of a group that needs to be deprotonated, which exhibits a pK value of 4.5 ± 0.2 , being protonated at low pH values and as the result of a group that needs to be protonated, which exhibits a pK value of 7.8 ± 0.1 , being deprotonated at high pH values. The pH dependence of k_{cat}/K_m varies only at high pH values, where it decreases as a result of the deprotonation of a group exhibiting a pK value of 8.3 ± 0.1 . This latter parameter includes rates for steps between the binding of nitrocefin to the free enzyme and the first irreversible step, which in the case of BlaC is the acylation of

 Scheme 3. Proposed Catalytic Mechanism of BlaC^a


^aActivation of Ser70 by Lys73 to promote β -lactam ring opening (A), formation of the covalent acyl-enzyme intermediate (B), and activation of the conserved water by Glu166 for product release (C).

Table 1. Solvent Kinetic Isotope Effects on BlaC^a

	Nitrocefin	Cefoxitin	Meropenem
K_m (μM)	86 ± 6	255 ± 9	3.4 ± 0.7^b
k_{cat} (s^{-1})	38 ± 2	0.7 ± 0.1	$(7.7 \pm 0.1) \times 10^{-4}^b$
k_{cat}/K_m ($\text{M}^{-1} \text{s}^{-1}$)	$(4.4 \pm 0.1) \times 10^5$	$(2.7 \pm 0.3) \times 10^3$	$(2.3 \pm 0.8) \times 10^2^b$
k_2 (s^{-1})	na ^c	1.4 ± 0.01	$(1.9 \pm 0.3) \times 10^{-2}$
k_3 (s^{-1})	na ^c	0.4 ± 0.001	$(4.0 \pm 1.0) \times 10^{-4}$
$^{\text{D}}V/K$	1.4 ± 0.1	1.8 ± 0.1	na ^c
$^{\text{D}}V$	1.4 ± 0.1	3.9 ± 0.1	na ^c
$^{\text{D}}k_2$	na ^c	1.6 ± 0.1	3.8 ± 0.1
$^{\text{D}}k_3$	na ^c	3.4 ± 0.1	4.0 ± 0.1

^a V/K reports on k_{cat}/K_m ; V reports on k_{cat} . k_2 is the rate of acylation, and k_3 is the rate of deacylation. ^bData obtained from ref 31. ^cNot available.

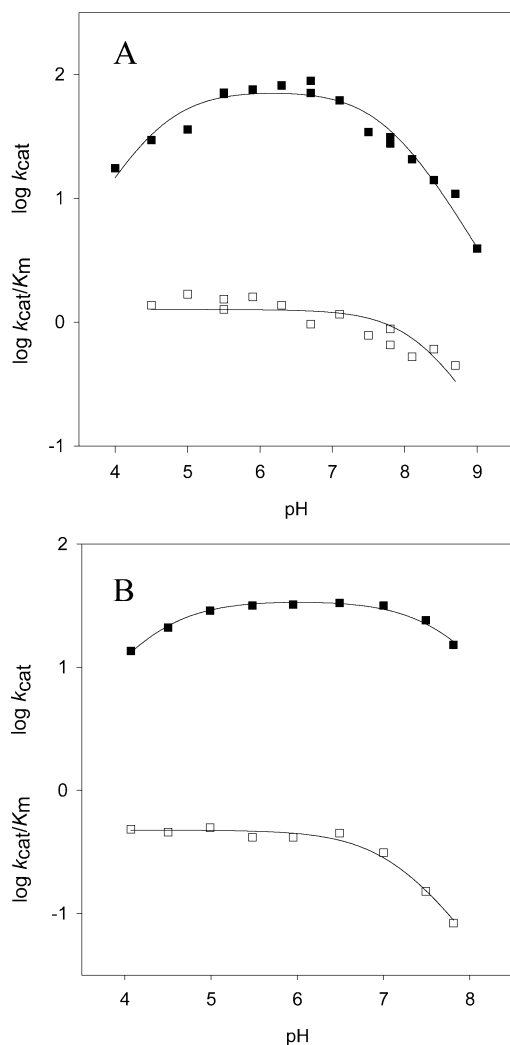
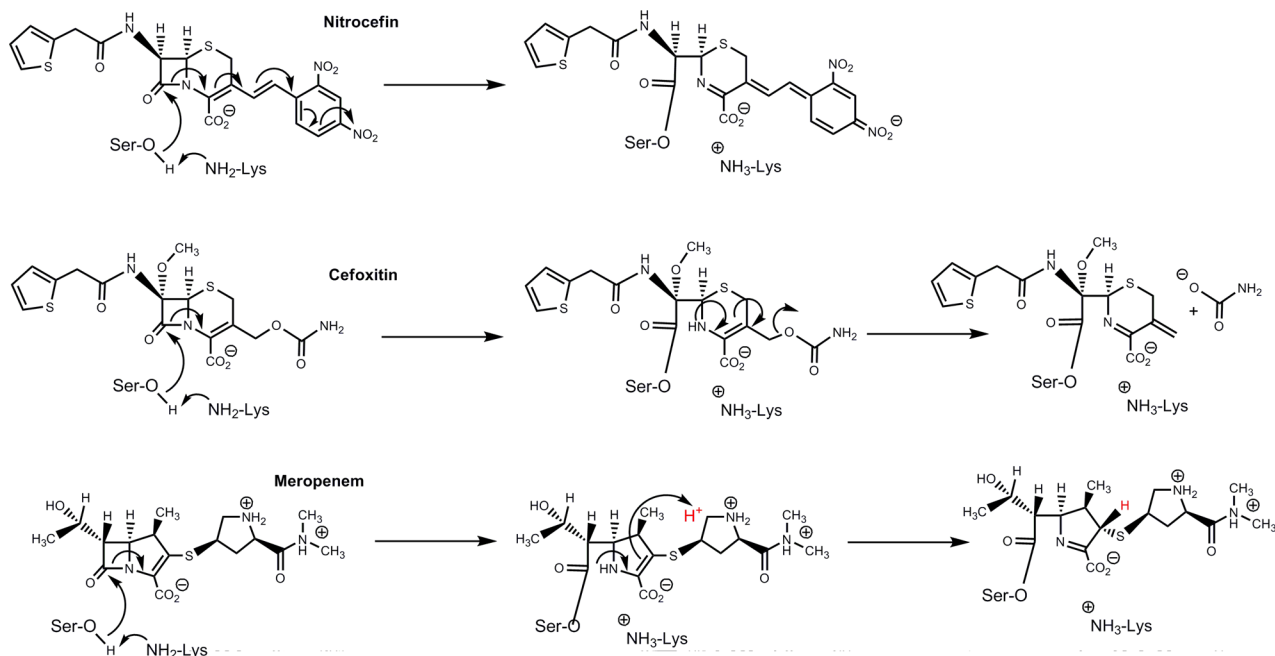


Figure 1. pH dependence of BlaC on log k_{cat} (■) and log k_{cat}/K_m (□) with (A) nitrocefin and (B) cefoxitin. The lines represent fits to eqs 1 and 2. A mixed buffer system was utilized, with 100 mM sodium acetate, 100 mM MES, 100 mM HEPES, and 100 mM TAPS.

the enzyme.¹⁶ The lack of any break at low pH of the k_{cat}/K_m profile suggests this parameter is not sensitive or reporting on rate-limiting chemistry that is dependent on the protonation states of enzyme groups. The decrease at high values may reflect the ionization behavior of a protonated enzyme group involved in nitrocefin binding, likely the carboxylate that is a necessary feature of all β -lactam antibiotics.

The bell-shaped k_{cat} profile suggests that chemistry requires the participation of an active site base and acid. Similar bell-shaped pH profiles have been reported for the *Bacillus cereus* β -lactamase acting on both penicillins and cephalosporins.²³ The maximal velocity includes rate contributions from steps after the formation of the enzyme–substrate complex to the final release of products and, in the context of the model for BlaC catalysis, includes both the acylation rate constant and the deacylation rate constant. For reasons that will be discussed further below, we believe that the acylation step in nitrocefin hydrolysis is slower than deacylation, and that the k_{cat} –pH profile is reporting on the ionization behavior of active site groups involved in this step. This suggests that the acylation base observed is Lys73, which removes a proton from the Ser70 hydroxyl that attacks the β -lactam amide linkage (Scheme 3). The extremely low pK value of 4.5 observed for this lysine likely reflects the highly perturbed active site environment after substrate binding. The observation of a general acid in the acylation reaction of nitrocefin is also unusual, because upon β -lactam ring opening, the resulting anion on the thiazole ring nitrogen is highly resonance-stabilized via the conjugated vinylidinitrobenzene substituent and would not be expected to require nitrogen protonation (Scheme 4). However, similar k_{cat} –pH profiles with this substrate were previously reported for the β -lactamase from *B. cereus*.²³

These studies were repeated with cefoxitin, a significantly slower cephalosporin substrate with a k_{cat} of 0.7 s^{-1} and a k_{cat}/K_m of $2.7 \times 10^3 \text{ M}^{-1} \text{ s}^{-1}$ (Table 1). The pH dependence of the kinetic parameters determined using cefoxitin appears to be similar to that exhibited by nitrocefin, and the interpretation of the k_{cat}/K_m profile is the same, a protonated group exhibiting a pK value of 6.8 ± 0.1 responsible for interacting with the carboxylic acid of cefoxitin. However, the interpretation of the pH dependence of k_{cat} is different. The maximal velocity depends on a group that needs to be deprotonated, and exhibits a pK value of 4.3 ± 0.1 , and as the result of a group that needs to be protonated, and exhibits a pK value of 7.7 ± 0.1 , is deprotonated at high pH values. Cefoxitin exhibits “burst” kinetics in stopped-flow experiments (see below), thus requiring that the acylation step be faster than the deacylation step. Because the k_{cat} for this substrate is determined by the deacylation step, the pH dependence of k_{cat} must involve the protonation states of enzyme groups involved in that reaction. Deacylation is initiated by Glu166 abstracting a proton from a conserved water molecule in the active site that hydrolyzes the ester bond. The Ser70 alkoxide is protonated by Lys73, regenerating the protonation state of these residues to that required for the next round of catalysis. Thus, in the deacylation reaction, the function and protonation states of the two

Scheme 4. BlaC Acylation of Nitrocefin, Cefoxitin, and Meropenem^a


^aPyrroline ring isomerization of meropenem results in the incorporation of a solvent exchangeable proton (red) at C3.

auxiliary acid–base groups are reversed, with the group acting as the general base being Glu166 and the group acting as the general acid being Lys73.

One final comment is worth making given the various pH environments that *M. tuberculosis* must confront. After ingestion by host alveolar macrophages, the organism is subjected to increasingly low pH values as the phagosome is slowly acidified and experiences intraphagosomal pH values as low as 5.5.²⁴ BlaC, which is located in the periplasmic space outside of the cell membrane, remains remarkably active at these low pH values as a result of the low pK values of Lys73 in the acylation half-reaction and Glu166 in the deacylation half-reaction and thus can function effectively even as acidification of the phagosome is occurring.

Solvent Kinetic Isotope Effects. The foregoing analysis of the pH dependence of the kinetic parameters allowed us to define a broad pH window in which solvent kinetic isotope effects would be unaffected by solvent isotope composition-dependent changes in the pK values of the enzyme groups involved in catalysis. Solvent kinetic isotope effects are powerful probes of proton transfer events in enzyme-catalyzed reactions and have been used to probe transition-state structures for both acylation and deacylation reactions of mechanism-related proteases.^{25–28} This extends to the analysis of SKIEs on β -lactamases, including important contributions to penicillanic acid sulfone hydrolysis by Brenner and Knowles,²⁹ β -lactam hydrolysis by the *B. cereus* TEM β -lactamase by Hardy and Kirsch,²³ and β -lactam hydrolysis by several related class A and C β -lactamases by Adediran and Pratt.³⁰ We chose three substrates for this analysis with BlaC, which span 5.5 orders of catalytic turnover rates. Viscosity controls showed no effect of added 9% (w/w) glycerol on rates.

Nitrocefin steady-state hydrolysis is only modestly affected by solvent isotopic composition (Figure 2A). Fits of the data points yielded equivalent, small solvent kinetic isotope effects of 1.4 ± 0.1 on V and V/K (Table 1). The small effects on the

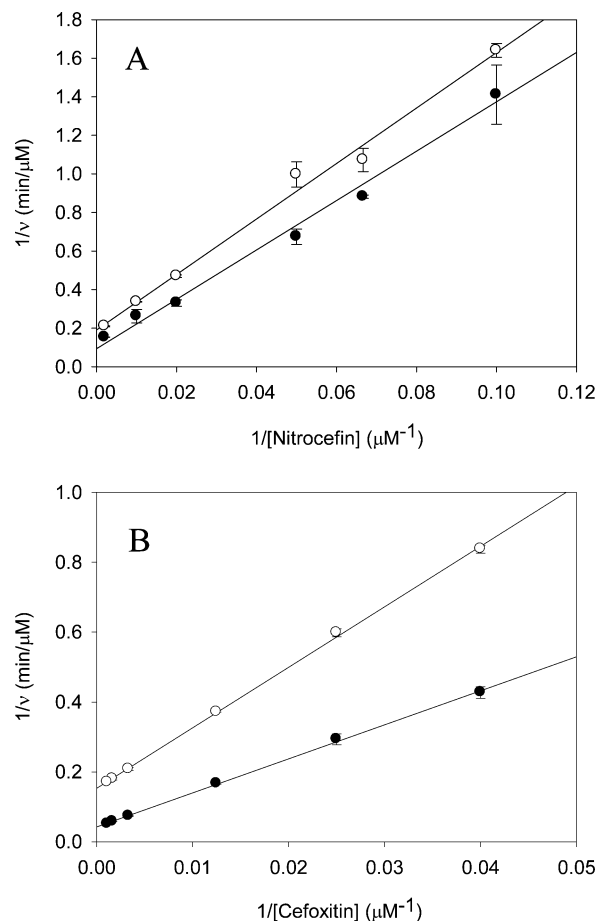


Figure 2. Initial velocity pattern for (A) nitrocefin with 2 nM BlaC and (B) cefoxitin with 350 nM BlaC in H₂O (●) and in D₂O (○) with 100 mM MES (pH 6.3).

hydrolysis of nitrocefin by BlaC are similar to those observed previously for penicillins and cephalosporins by Hardy and Kirsch (1.53 and 1.09, respectively) and Adediran and Pratt (1.04 and 1.39, respectively).^{23,30} Because k_{cat}/K_m reports on only steps between binding of nitrocefin to the free enzyme and the formation of the covalent acyl-enzyme, the steady-state results for nitrocefin hydrolysis revealing equivalent, small solvent KIEs on k_{cat}/K_m and k_{cat} support acylation as the rate-limiting step for this substrate. The proton inventory for nitrocefin hydrolysis is linear (Figure 3A), suggesting a single

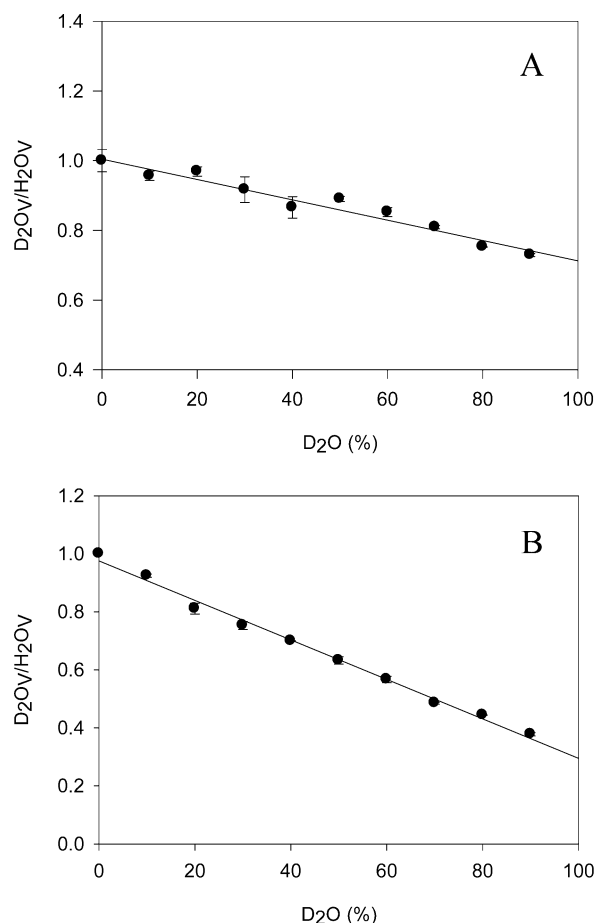


Figure 3. Proton inventories of (A) nitrocefin with 2 nM BlaC and (B) cefoxitin with 350 nM BlaC measuring the initial velocity with varying levels of D₂O in 100 mM MES (pH 6.3).

proton is transferred in the acylation reaction, and we assign this to the exchangeable proton on the Ser70 hydroxyl group to Lys73 (Scheme 4). Nitrocefin is too fast as a substrate for stopped-flow, multiple-turnover kinetics, and thus, we could not obtain independent evidence of the SKIEs on k_2 (acylation) and k_3 (deacylation). We then turned to both steady-state and multiple-turnover SKIE determinations of cefoxitin hydrolysis.

Fits of the steady-state data for SKIEs on cefoxitin hydrolysis yielded a $^{D_2O}V/K$ value of 1.8 ± 0.1 and a ^{D_2O}V value of 3.9 ± 0.1 (Table 1 and Figure 2B). The much larger solvent kinetic isotope effect on V suggests that deacylation is rate-limiting for this substrate. A single proton is involved in this proton transfer step as evidenced by the linear proton inventory obtained at saturating cefoxitin concentrations (Figure 3B). We ascribe this proton transfer as being the deprotonation of the conserved hydrolytic water molecule by Glu166 (Scheme 3). The small

SKIE on V/K is likely to arise from a small contribution from the acylation reaction, as observed with nitrocefin.

Under multiple-turnover conditions, using high concentrations of both BlaC and cefoxitin, a distinct burst of absorbance is observed (Figure 4A), suggesting that the on-

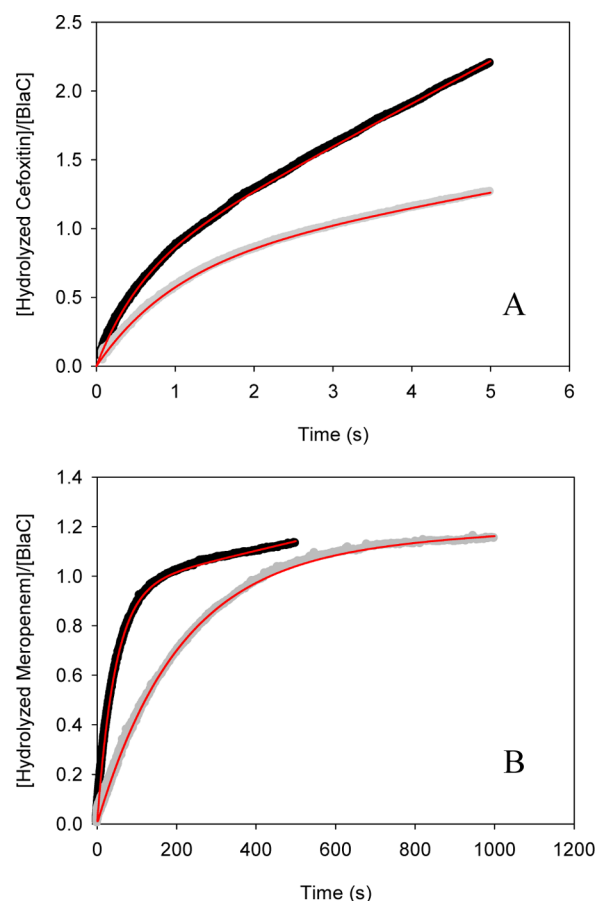


Figure 4. Burst kinetic isotope effects on (A) cefoxitin and (B) Meropenem in H₂O (black) and D₂O (gray) with fits of experimental traces to the k_2 (acylation) and k_3 (deacylation) values listed in Table 1 (red). Cefoxitin studies were conducted with 1800 μM cefoxitin mixed with 27 μM BlaC. Meropenem studies were conducted with 875 μM Meropenem mixed with 13 μM BlaC.

enzyme acylation rate is faster than the slower hydrolysis of the cefoxitin–BlaC complex. When the curves are fit to an equation that describes an exponential increase followed by a linear (“steady-state”) rate, SKIEs are again observed on both the burst phase and the linear rate. By fitting progress curves to eq 6, we can dissect the rates of acylation and deacylation and determine the SKIEs on both k_2 and k_3 . Because k_3 is not significantly slower than k_2 (Table 1), we observe an initial burst phase (A_0) equivalent to approximately 70% of the initial active enzyme concentration used to initiate the reaction. The solvent kinetic isotope effect on k_2 , $^{D_2O}k_2$, was 1.6 ± 0.1 , and that on k_3 , $^{D_2O}k_3$, was 3.4 ± 0.1 (Table 1). The near equivalence of the values for ^{D_2O}V and $^{D_2O}k_3$ argues persuasively for the effect of isotope on deacylation being the source of the solvent kinetic isotope effect on V . The combination of the burst and the SKIEs on each phase of the reaction demonstrates that deacylation is rate-limiting. Viscosity controls using 9% (w/w) glycerol for cefoxitin and Meropenem hydrolysis exhibited a slight increase in the linear steady-state rate, as observed

previously with P99 β -lactamase by Adediran and Pratt.³⁰ The presence of glycerol in the active site may act like methanol by accelerating the solvolysis of the slow step (deacylation) and thereby enhancing k_{cat} (see Figure S2 of the Supporting Information).

The very low K_m value for Meropenem, coupled with its weak extinction coefficient and extremely slow turnover rate (3.4 s^{-1}),³¹ made the determination of steady-state SKIEs both difficult and unreliable. Rather, we measured SKIEs at a single high concentration of both BlaC and Meropenem. As seen in Figure 4B, there is an easily observable burst followed by a linear steady-state rate of hydrolysis. Here, we observe a stoichiometric burst phase with the initial enzyme concentration because the deacylation rate is significantly slower than the acylation rate. When the experiment was repeated in D_2O , both rates were quite significantly affected, with a calculated $\text{D}_2\text{O}k_2$ value of 3.8 ± 0.1 and a calculated $\text{D}_2\text{O}k_3$ value of 4.0 ± 0.1 (Table 1). The SKIE on deacylation corresponds almost exactly to that observed with cefoxitin and has the same source, e.g., deprotonation of the water molecule by Glu166 leading to hydrolysis of the ester-linked substrate. The source of the equally large SKIE on k_2 is less obvious and unique to this carbapenem substrate. During the acylation reaction of carbapenems catalyzed by BlaC, the ring opening of the β -lactam yields an initial Δ^2 -pyrroline ring, which subsequently isomerizes to a Δ^1 -pyrroline ring with stereospecific protonation at C3 to generate the final, stable acylated adduct (Scheme 4). The structure of the Meropenem complex with BlaC shows that C3 is clearly sp^3 hybridized,² and structures determined for the related carbapenems, ertapenem and doripenem, reveal similar isomerizations.¹⁶ This reaction is not, however, concerted as the Δ^2 -pyrroline ring form is observed crystallographically at very short soaking times with ertapenem. We propose that it is this solvent-mediated C3 protonation accompanying isomerization that is the source of the solvent KIE on k_2 , the acylation rate constant.

In conclusion, the *M. tuberculosis* blaC-encoded β -lactamase exhibits a very broad pH dependence that allows it to act effectively at a range of pH values that the organism encounters, including the low pH levels that it encounters intracellularly after macrophage capture. The pH dependence of k_{cat} is reporting on the required ionization states of Glu166 and Lys73 for effective acylation of the enzyme (for nitrocefin) or for deacylation of the covalent enzyme- β -lactam complex (for the slower cefoxitin and Meropenem). This interpretation is confirmed by the solvent kinetic isotope effect analysis with relatively slow substrates, such as cefoxitin, where large SKIEs are observed on k_3 , the deacylation rate constant, and not k_2 , the acylation rate constant. However, using Meropenem, an extremely slow substrate, large SKIEs are observed on both acylation and deacylation, the former being the result of a slow isomerization-protonation step accompanying β -lactam ring opening of the carbapenem nucleus. This interpretation has support from structural studies of the Δ^2 - and Δ^1 -pyrroline forms of the enzyme-bound carbapenem.¹⁶

■ ASSOCIATED CONTENT

● Supporting Information

Burst SKIEs in 9% glycerol, saturation curve for cefoxitin burst kinetics, and pre-steady-state fittings for kinetic parameters. This material is available free of charge via the Internet at <http://pubs.acs.org>.

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Funding

This work was supported by the National Institutes of Health Grants AI33696 and AI60899 to J.S.B.

Notes

The authors declare no competing financial interest.

■ ACKNOWLEDGMENTS

We thank Dr. Clarissa Czekster for her guidance in performing the stopped-flow experiments.

■ ABBREVIATIONS

AU, absorbance units; BlaC, β -lactamase; ESBL, extended-spectrum β -lactamase; HEPES, *N*-(2-hydroxyethyl)piperazine-*N'*-2-ethanesulfonic acid; MES, 2-(*N*-morpholino)-ethanesulfonic acid; SKIE, solvent kinetic isotope effect; TAPS, *N*-[tris(hydroxymethyl)methyl]-3-aminopropanesulfonic acid.

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