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Reversible inactivation of a peptidoglycan transpeptidase by a β -lactam antibiotic mediated by β -lactam-ring recyclization in the enzyme active site

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β -lactam antibiotics act as suicide substrates of transpeptidases responsible for the last cross-linking step of peptidoglycan synthesis in the bacterial cell wall. Nucleophilic attack of the β -lactam carbonyl by the catalytic residue (Ser or Cys) of transpeptidases results in the opening of the β -lactam ring and in the formation of a stable acyl-enzyme. The acylation reaction is considered as irreversible due to the strain of the β -lactam ring. In contradiction with this widely accepted but poorly demonstrated premise, we show here that the acylation of the L,D-transpeptidase Ldt_{fm} from *Enterococcus faecium* by the β -lactam nitrocefin is reversible, leading to limited antibacterial activity. Experimentally, two independent methods based on spectrophotometry and mass spectrometry provided evidence that recyclization of the β -lactam ring within the active site of Ldt_{fm} regenerates native nitrocefin. Ring strain is therefore not sufficient to account for irreversible acylation of peptidoglycan transpeptidases observed for most β -lactam antibiotics.

β -lactams remain the corner stone of antibacterial chemotherapy 76 years after the first therapeutic use of an antibiotic, namely penicillin, in 1941. β -lactams kill bacteria by inactivating the D,D-transpeptidases responsible for the last cross-linking step of peptidoglycan synthesis¹. The latter cell wall polymer is a giant macromolecule of 10⁹ to 10¹⁰ daltons that completely surrounds bacterial cells and plays an essential role in counteracting the osmotic pressure of the cytoplasm during the entire cell cycle including cell division². β -lactams are structure analogues of the peptidoglycan precursors and act as suicide substrates of the transpeptidases³. The enzymes catalyze formation of peptidoglycan cross-links in a two-step reaction⁴. In the first step, the D,D-transpeptidases interact with an acyl donor containing a pentapeptide stem ending in D-Ala⁴-D-Ala⁵, leading to the release of D-Ala⁵ and to the formation of an ester bond between the carbonyl of D-Ala⁴ and the catalytic Ser residue of the enzymes. In the second step, nucleophilic attack of the resulting acyl-enzyme by the acyl acceptor generates the peptidoglycan cross-link. Similar to the first step of the transpeptidation reaction, nucleophilic attack of the carbon carbonyl of β -lactams by the catalytic Ser leads to rupture of the amide bond of the β -lactam ring and inactivation of the D,D-transpeptidases. The resulting acyl-enzymes are highly stable, with typical half-lives in the order of several hours, due to low hydrolysis rates. For this reason, the inactivation reaction is considered as irreversible since recovery of enzyme activity through hydrolysis of the drug-enzyme adduct occurs at a timescale that is too large to compromise antibacterial activity.

The D,D-transpeptidases have been historically referred to as penicillin-binding proteins (PBPs) since these enzymes were routinely identified by covalent labeling with radioactive penicillin followed by gel electrophoresis¹. PBPs belong to a super family of mechanistically and structurally related active-site serine enzymes that also comprise D,D-carboxypeptidases, endopeptidases, and most β -lactamases responsible for β -lactam resistance by drug detoxification^{3,5}. The acyl-enzymes formed by the acylation of the Ser residue of β -lactamases are

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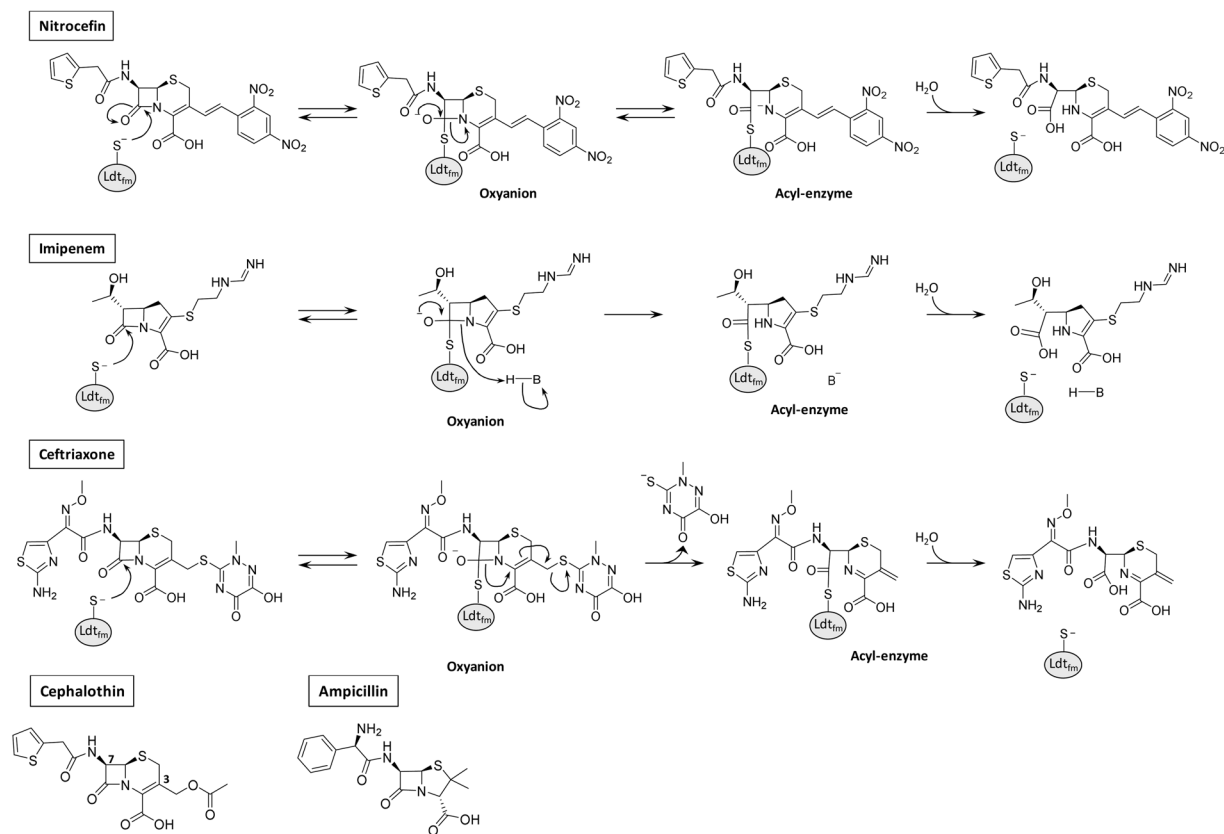


Figure 1. Reactions catalyzed by the L,D-transpeptidase Ldt_{fm} with nitrocefin, imipenem, and ceftriaxone. The figure also shows the structures of cephalothin and ampicillin. The base (B) that provides a proton to the β-lactam nitrogen of imipenem has not been identified^{23,26}.

rapidly hydrolyzed leading to turnover and drug detoxification. More recently, the family of β-lactam-interacting enzymes has been enriched by the detection of active-site cysteine L,D-transpeptidases that by-pass the D,D-transpeptidase activity of PBPs in β-lactam-resistant mutants of *Enterococcus faecium* and *Escherichia coli*^{6,7}. These L,D-transpeptidases were also found to be the main peptidoglycan cross-linking enzymes in *Clostridium difficile* and in mycobacteria⁸. The L,D-transpeptidases are structurally unrelated to active-site Ser PBPs and cleave the L-Lys³-D-Ala⁴ or diaminopymelate³-D-Ala⁴ peptide bond of an acyl donor containing a tetrapeptide stem^{9,10}. In spite of this difference in substrate specificity, L,D-transpeptidases interact with β-lactams belonging to the carbapenem class¹¹ (Fig. 1) that are currently being evaluated for the treatment of tuberculosis¹² and of pulmonary infections due to *Mycobacterium abscessus* in cystic fibrosis patients¹³. The L,D-transpeptidases are also acylated by β-lactams belonging to the cephalosporin and penam classes but this does not lead to target inactivation and antibacterial activity since the corresponding adducts are prone to hydrolysis¹⁴.

Due to the strain of the four-membered ring of β-lactams, the acylation reaction is considered as irreversible in the sense that re-sealing of the scissile amide bond of the β-lactam ring is not thought to occur at any significant rate^{15,16}. In this study, we have critically evaluated this postulate by studying the acylation of the L,D-transpeptidase Ldt_{fm} from *E. faecium* by nitrocefin, a chromogenic cephalosporin. We show that this drug-enzyme combination displays high acylation and low hydrolysis rates leading to acyl-enzyme accumulation. This offered the possibility to assay for recyclization of the β-lactam ring by two approaches based on the use of a competing β-lactam or a competing enzyme. We provide evidence that recyclization of the β-lactam ring of nitrocefin in the acyl-enzyme regenerates the native drug that can freely diffuse out of the Ldt_{fm} active site. These data indicate that the irreversible nature of the acylation reaction is not exclusively driven by the strain of the four-membered ring of β-lactams.

Results

Modification of the absorbance spectrum of nitrocefin following acylation of the L,D-transpeptidase Ldt_{fm}. Hydrolysis of the β-lactam ring of nitrocefin (50 μM) by the β-lactamase BlaC (1 μM) in 100 mM sodium phosphate buffer (pH 6.0) produced the expected large increase in the absorbance at 486 nm (Fig. 2A). Under these conditions, the variation in the molar extinction coefficient ($\Delta\epsilon$) was 15,200 M⁻¹ cm⁻¹ (1,400 M⁻¹ cm⁻¹ versus 16,600 M⁻¹ cm⁻¹ for native and hydrolyzed nitrocefin, respectively). Incubation of Ldt_{fm} (22 μM) with increasing concentrations of nitrocefin (0 to 56 μM) also led to an increase of the absorbance at 486 nm (Fig. 2B). The increase in the absorbance was linear until the concentration of nitrocefin exceeded that of the enzyme. This result indicates that the increase in the absorbance at 486 nm is due to the rupture of

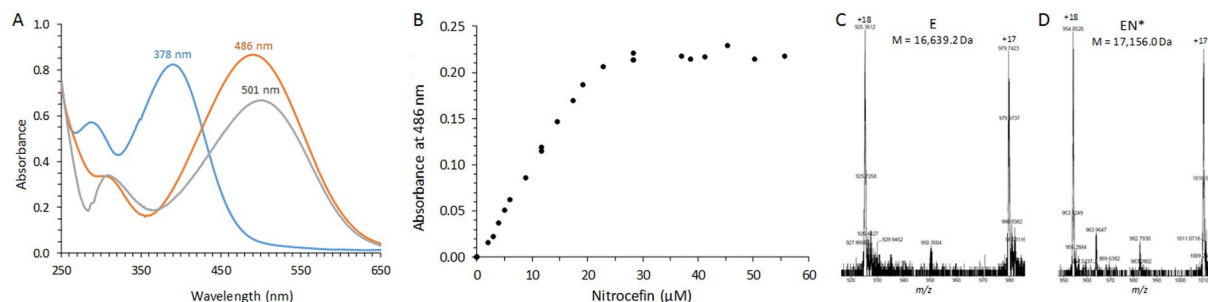


Figure 2. Determination by spectrophotometry of the acylation of Ldt_{fm} by nitrocefim. **(A)** Absorbance spectra of various forms of nitrocefim ($50 \mu\text{M}$). Blue, native nitrocefim; orange, nitrocefim hydrolyzed by BlaC; grey, nitrocefim in the acyl-enzyme formed with Ldt_{fm} . **(B)** Titration of the active site of Ldt_{fm} ($22 \mu\text{M}$) by nitrocefim. **(C and D)** Mass spectra of Ldt_{fm} and of the acyl-enzyme formed with nitrocefim. The average mass of Ldt_{fm} (E) and of the acyl-enzyme (EN*) were deduced from the m/z ratios of the $[M + 17H]^{17+}$ and $[M + 18H]^{18+}$ ions. The calculated mass of nitrocefim, Ldt_{fm} , and of the acyl-enzyme were 516.5 Da , $16,639.3 \text{ Da}$, and $17,155.8 \text{ Da}$, respectively.

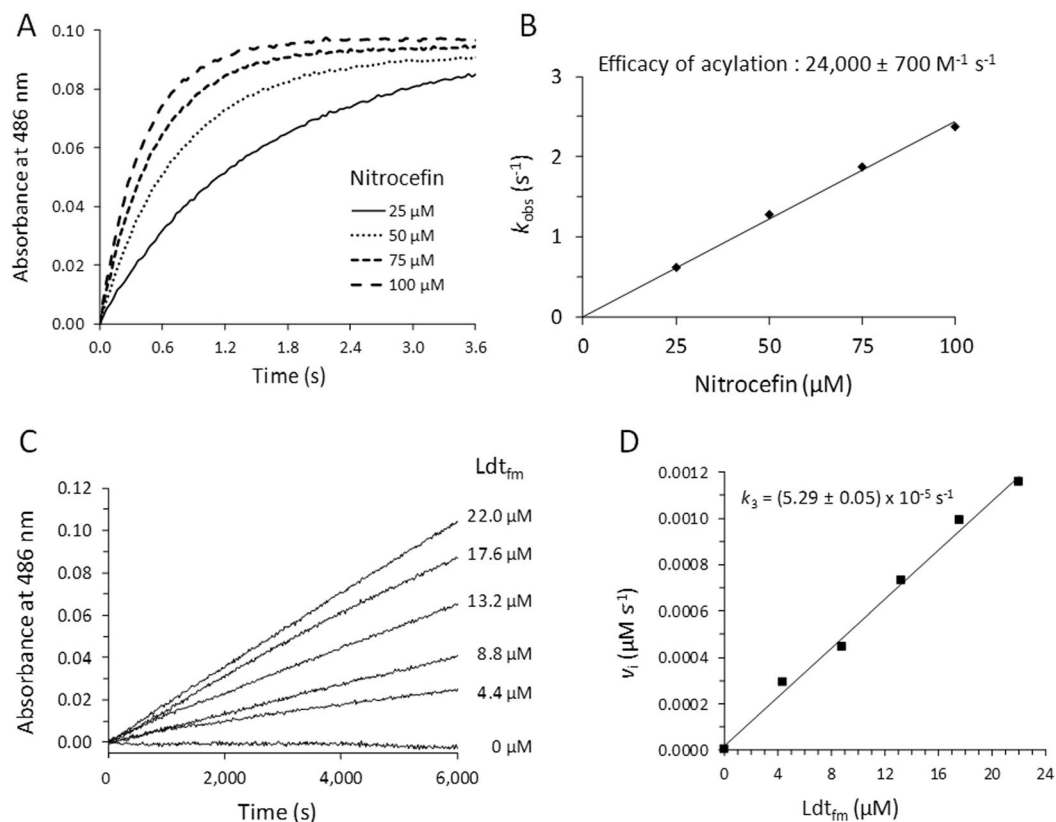
the β -lactam ring of nitrocefim upon acylation of the catalytic Cys residue of Ldt_{fm} . In agreement, a covalent adduct was detected by mass spectrometry (Fig. 2C). The spectra of nitrocefim hydrolyzed by BlaC and of nitrocefim linked to Ldt_{fm} in the acyl-enzyme were similar with two minor differences. Nitrocefim in the acyl-enzyme absorbed less than hydrolyzed nitrocefim ($\epsilon = 9,900 \text{ M}^{-1} \text{ cm}^{-1}$ versus $16,600 \text{ M}^{-1} \text{ cm}^{-1}$ at 486 nm , respectively). A minor red shift might occur in the absorbance maximum (λ_{max} of 501 nm versus 486 nm , respectively). Thus, nitrocefim provides a sensitive assay to titrate the active site of Ldt_{fm} .

Kinetic analyses of Ldt_{fm} acylation by nitrocefim. Stopped-flow kinetics were performed to assess the efficacy of acylation of Ldt_{fm} ($11 \mu\text{M}$) by nitrocefim ($25, 50, 75,$ and $100 \mu\text{M}$; Fig. 3A). The concentration of the acyl-enzyme (EI*) was determined based on the difference in the molar extinction coefficient ($\Delta\epsilon_{486 \text{ nm}} = 8,500 \text{ M}^{-1} \text{ cm}^{-1}$) between native nitrocefim ($\epsilon = 1,400 \text{ M}^{-1} \text{ cm}^{-1}$) and nitrocefim in the acyl-enzyme ($\epsilon = 9,900 \text{ M}^{-1} \text{ cm}^{-1}$) (Fig. 2). To determine the value of k_{obs} , as defined in equation 1 (Fig. 3), exponential rise to maximum was fitted to kinetic data. Full acylation of Ldt_{fm} was observed at all the concentrations of nitrocefim that were tested and the value of k_{obs} increased linearly with the concentration of nitrocefim (Fig. 3B). The slope provided an estimate of the efficacy of the acylation reaction ($24,000 \pm 700 \text{ M}^{-1} \text{ s}^{-1}$).

Hydrolysis of the acyl-enzyme. Incubation of nitrocefim ($50 \mu\text{M}$) with various concentrations of Ldt_{fm} (0 to $22 \mu\text{M}$) for an extended time period (14 hours) revealed an increase in the absorbance at 486 nm (Fig. 3C), which was proportional to the enzyme concentration (Fig. 3D). The concentration of hydrolyzed nitrocefim was determined using the value of $\Delta\epsilon$ of $15,200 \text{ M}^{-1} \text{ cm}^{-1}$ obtained above with BlaC at 486 nm (Fig. 2). In these experiments, data were acquired between 1 min and 14 hours due to the lag time required to charge an automatic sample changer. Thus, Ldt_{fm} was fully acylated at the beginning of the recording of the data (see above, Fig. 3A). The initial absorbance was subtracted to generate the kinetics presented in Fig. 3C. In these conditions, the increase in absorbance was due to enzyme turnover resulting in nitrocefim hydrolysis (Scheme 1 in Fig. 3). The observed turnover ($5.29 \pm 0.05 \times 10^{-5} \text{ s}^{-1}$) corresponds to the value of the kinetic parameter k_3 since the bulk of Ldt_{fm} is acylated under these conditions (Fig. 3A). The value of k_3 implies a half-life of 218 min for the acyl-enzyme. The high efficacy of acylation coupled to the low value of k_3 accounts for the steady-state presence of EN* as the main form of Ldt_{fm} in the presence of nitrocefim.

Competitive acylation of Ldt_{fm} by nitrocefim and the carbapenem imipenem. Ldt_{fm} ($22 \mu\text{M}$) was simultaneously exposed to nitrocefim ($50 \mu\text{M}$) and imipenem ($64 \mu\text{M}$) (Fig. 4A). The kinetics without imipenem was determined as a control. Imipenem had little impact on the initial rate of acylation of Ldt_{fm} by nitrocefim, as determined by the increase in the absorbance at 486 nm . Thus, imipenem did not significantly interfere with acylation of Ldt_{fm} by nitrocefim. Full acylation of Ldt_{fm} by nitrocefim was observed in *ca.* 3 s but a longer incubation in the presence of imipenem resulted in a decrease in the absorbance at 486 nm (Fig. 4B). This decrease could not be attributed to the hydrolysis of the acyl-enzyme that would result in an increase rather than a decrease in the absorbance of 486 nm . At this wavelength, reactions involving hydrolysis of imipenem or acylation of Ldt_{fm} by this drug had no impact on the absorbance (data not shown). We therefore concluded that native nitrocefim and nitrocefim bound to the catalytic Cys of Ldt_{fm} in the acyl-enzyme were in equilibrium and that imipenem displaced this equilibrium in favor of the native drug. In the competing acylation step, Ldt_{fm} was initially fully acylated by nitrocefim but imipenem gradually replaced nitrocefim in the enzyme active site since the acylation reaction was irreversible with imipenem but reversible with nitrocefim (Scheme 2). In the control experiment without imipenem, slow hydrolysis of nitrocefim was observed as expected from the kinetics displayed in Fig. 3C.

In the above experiments involving incubation of Ldt_{fm} with imipenem and nitrocefim, the recyclization reaction generates the native form of Ldt_{fm} , which is expected to be competitively acylated by imipenem and nitrocefim. Recyclization followed by acylation with nitrocefim contributes to a futile cycle with no net impact on the concentration of nitrocefim. This implies that the net rate of nitrocefim recyclization should increase with the



$$\text{Equation 1} \quad EN^* = E_0 (1 - e^{-k_{\text{obs}}t})$$

$$\text{Equation 2} \quad v_i = k_3 E$$

Scheme 1

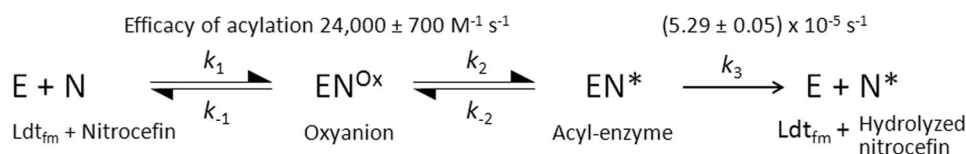
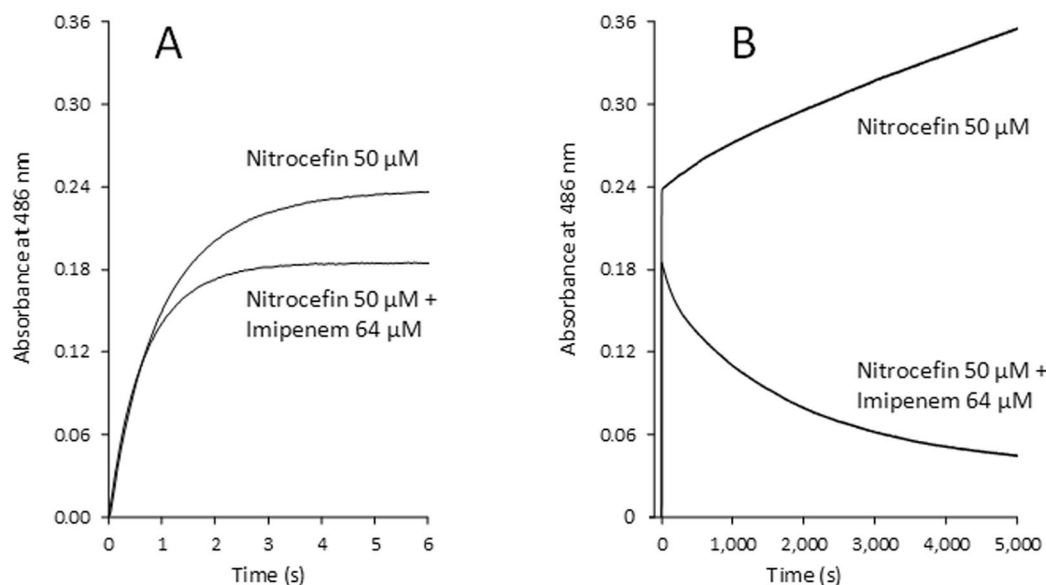


Figure 3. Kinetics of Ldt_{fm} acylation by nitrocefin and of hydrolysis of the resulting acyl-enzyme. (A) Stopped-flow kinetics of acylation of Ldt_{fm} (11 μM) by nitrocefin (25, 50, 75, and 100 μM) in 100 mM sodium phosphate buffer (pH 6.0) at 20 °C. The value of k_{obs} was determined by fitting Equation 1 to data. E_0 , concentration of native Ldt_{fm} at $t = 0$. (B) Variation of k_{obs} as a function of the concentration of nitrocefin. (C) Hydrolysis of a fixed concentration of nitrocefin (50 μM) by Ldt_{fm} at various concentrations (0 to 22 μM). (D) The initial rate of hydrolysis (v_i) was plotted as a function of the concentration of Ldt_{fm} and the value of k_3 was determined by linear regression (Equation 2) considering that Ldt_{fm} is fully acylated under these conditions.

concentration of the competing imipenem. Testing recyclization of nitrocefin (50 μM) in the presence of various concentrations of imipenem (64, 128, 256, and 512 μM) revealed the expected increase in the net rate of nitrocefin recyclization (Supplementary Fig. 1).

Competitive acylation of two forms of Ldt_{fm} by nitrocefin. Since the reversibility of the acylation reaction was unprecedented for any β -lactam, our next objective was to demonstrate the reversibility of the acylation of Ldt_{fm} by nitrocefin using a second independent method. We surmise that resealing of the β -lactam ring of nitrocefin should regenerate the native form of the drug that could leave the Ldt_{fm} active site, diffuse in the medium, and competitively acylate another molecule of Ldt_{fm} (Fig. 5A). To bring these reactions to light, we prepared uniformly ^{13}C - and ^{15}N -labeled Ldt_{fm} . The unlabeled and labeled forms of Ldt_{fm} , which were readily identified by mass spectrometry, were expected to have similar catalytic properties since they only differ by their isotopic content. To test for the reversibility of the acylation reaction, a limiting amount of nitrocefin (1.5 nmole) was first incubated with labeled Ldt_{fm} (2.2 nmole) in 100 μl of 5 mM sodium phosphate buffer (pH 6.0). The extent of acylation (ca. 82%) indicated that the bulk of nitrocefin had reacted with Ldt_{fm} (Fig. 5B). Then, unlabeled Ldt_{fm}



Scheme 2

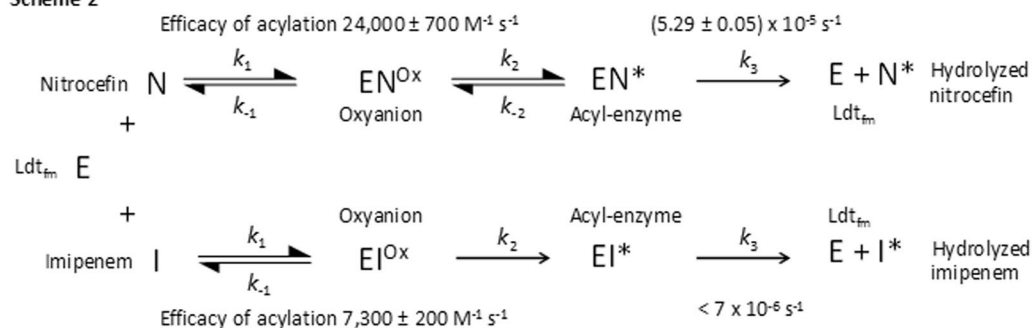


Figure 4. Competitive acylation of Ldt_{fm} by nitrocefim and imipenem. One syringe of the stopped-flow apparatus contained nitrocefim ($100 \mu\text{M}$) and imipenem ($128 \mu\text{M}$) in 100 mM sodium phosphate buffer (pH 6.0). The second syringe contained Ldt_{fm} ($44 \mu\text{M}$) in the same buffer. Equal volumes from syringes 1 and 2 were injected into the cuvette of the spectrophotometer and the absorbance was recorded at 486 nm . Concentrations indicated in the Figure take into account the two-fold dilution. Panels A and B represent the same kinetics for two timescales ($0\text{--}6 \text{ s}$ and $0\text{--}5000 \text{ s}$, respectively). In panel a, the increase in the absorbance at 486 nm mainly results from the rupture of the β -lactam ring of nitrocefim upon acylation of Ldt_{fm} . In panel B, further increase in the absorbance observed in the absence of imipenem (upper curve) results from slow hydrolysis of nitrocefim. In the presence of imipenem, recyclization of the β -lactam ring of nitrocefim accounts for the decrease in the absorbance.

(2.2 nmole) was added and the partition of nitrocefim between the two forms of Ldt_{fm} was determined by mass spectrometry. After 600 s of incubation an equilibrium was reached and 40% of the labeled and unlabeled forms of Ldt_{fm} were acylated by nitrocefim. This result indicates that the β -lactam ring of nitrocefim was resealed within the active site of labeled Ldt_{fm} since this was the only source of native nitrocefim for acylation of unlabeled Ldt_{fm} .

Antibacterial activity of nitrocefim and imipenem resulting from inhibition of Ldt_{fm} . We have previously reported activation of the L,D-transpeptidase pathway in a mutant of *Enterococcus faecium* (M512), which was selected in laboratory conditions on media containing increasing concentrations of ampicillin¹¹. Depending upon the growth conditions, the peptidoglycan of the mutant M512 is exclusively cross-linked by the PBPs or by Ldt_{fm} . In the presence of ampicillin ($32 \mu\text{g/ml}$), the D,D-transpeptidase activity of all PBPs produced by the mutant M512 is inhibited and Ldt_{fm} is the only functional transpeptidase. Antibiotic susceptibility testing in media containing ampicillin ($32 \mu\text{g/ml}$) was therefore used to assay for inhibition of Ldt_{fm} by nitrocefim. The broth microdilution assay revealed a 64-fold difference between the minimal inhibitory concentration (MIC) of nitrocefim ($4 \mu\text{g/ml}$) and that of imipenem ($0.0625 \mu\text{g/ml}$). Thus, nitrocefim was less active than imipenem in spite of a higher rate of acylation of Ldt_{fm} .

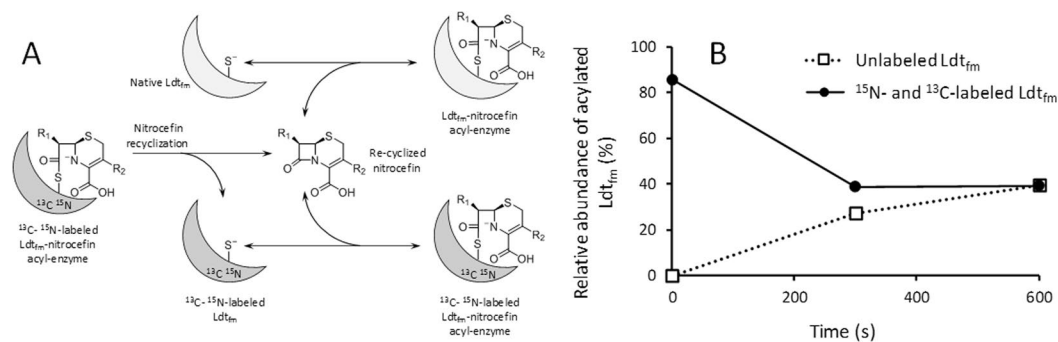


Figure 5. Detection of nitrocefin recyclization based on competitive acylation of unlabeled and labeled Ldt_{fm}. (A) Schematic representation of the reactions. Resealing of the β-lactam ring of nitrocefin regenerates the native form of the drug (Re-cyclized nitrocefin) that leaves the Ldt_{fm} active site and competitively acylates Ldt_{fm} or ¹³C-¹⁵N-labeled Ldt_{fm}. (B) Uniformly ¹³C- and ¹⁵N-labeled Ldt_{fm} (2.2 nmole) was incubated with nitrocefin (1.5 nmole) for 1 min at 20 °C in 100 μl of 5 mM sodium phosphate buffer pH 6.0. The spectrum of the protein revealed ca. 82% acylation (time = 0). Unlabeled Ldt_{fm} (2.2 nmole in 1.46 μl) was added and mass spectra were recorded at 300 s and 600 s. The relative abundance of the acylated forms of Ldt_{fm} was deduced from the relative intensity of the peaks.

Discussion

In conclusion, we provide evidence for reversible acylation of the L,D-transpeptidase Ldt_{fm} by nitrocefin. To the best of our knowledge, this is the first time that reversible acylation is documented for a β-lactam antibiotic. This conclusion is based on two independent approaches. First, we show that the deacylation (recyclization) reaction regenerates the β-lactam ring of nitrocefin based on spectrophotometry, a technique that provided access to kinetic evaluation (Fig. 3). Second, we show that the deacylation reaction regenerates a native molecule of nitrocefin that eventually leaves the Ldt_{fm} active site and acylates another enzyme molecule. This was established by using two preparations of Ldt_{fm} that only differ by their isotopic content.

The irreversibility of the acylation step is generally attributed to the strain of the β-lactam ring although this has not yet been explored by computational investigations¹⁵. In agreement, compounds containing a γ-lactam ring reversibly react with nucleophilic serine enzymes. For example, recyclization has been extensively documented for the 5-membered cyclic urea group of avibactam^{16,17}, a β-lactamase inhibitor recently approved for clinical use in combination with the β-lactam ceftazidime¹⁸.

Our experimental design provides direct evidence that acylation is reversible for certain β-lactams (e.g. nitrocefin) but not for others (e.g. imipenem) since simultaneous incubation of Ldt_{fm} with the two drugs led to the rapid formation of the Ldt_{fm}-nitrocefin acyl-enzyme, which was then slowly replaced by the Ldt_{fm}-imipenem acyl-enzyme. The basis for this difference remains unknown, but it is tempting to speculate that it involves the fate of the negative charge that develops on the β-lactam nitrogen upon rupture of the β-lactam ring (Fig. 1). The determination of the NMR structure of the ertapenem-Ldt_{fm} acyl-enzyme has shown that the nitrogen atom of this carbapenem is protonated¹⁹. In the case of nitrocefin, protonation of the nitrogen atom is not expected to be required for stabilization of the acyl-enzyme since the negative charge is delocalized by the very strong electron withdrawing power of the dinitrophenyl substituent. This could account for reversible acylation of Ldt_{fm} by nitrocefin but not by imipenem. Of note, resealing of the β-lactam ring is thermodynamically more favorable for L,D-transpeptidases than for classical PBPs since the thioester formed with cysteine-containing L,D-transpeptidases has a much higher potential than the corresponding ester formed with serine-containing β-lactamases and D,D-transpeptidases.

The efficacy of acylation of Ldt_{fm} by nitrocefin ($24,000 \pm 700 \text{ M}^{-1} \text{ s}^{-1}$) (Fig. 3) was higher than that observed for any carbapenem-L,D-transpeptidase combination in previous studies^{14,20–22}. The efficacy of acylation of Ldt_{fm} was found to be lower for cephalosporins such as ceftriaxone ($50 \pm 5 \text{ M}^{-1} \text{ s}^{-1}$) and cephalotin ($8.3 \pm 3.3 \text{ M}^{-1} \text{ s}^{-1}$) or penams such as ampicillin ($8.7 \pm 0.3 \text{ M}^{-1} \text{ s}^{-1}$)¹⁴. Since cephalotin and nitrocefin are two cephalosporins harboring the same side-chain at position C⁷ (Fig. 1), the 2,900-fold difference in the acylation efficacy observed between the two drugs is due to the side-chain at position C³. The NMR structure of Ldt_{fm} acylated by ertapenem indicates that there is little interaction, if any, between the side-chain at C³ and the enzyme²³. In agreement, the efficacy of acylation mainly varies with the β-lactam class (carbapenem, cephalosporin, or penam), with a lesser impact originating from differences in their side-chains^{14,21}. These observations suggest that the high efficacy of acylation of Ldt_{fm} by nitrocefin depends upon an effect of the side-chain on the reactivity of the β-lactam ring, as opposed to a direct interaction of the side-chain with the enzyme. This effect of the side-chain may involve the polarization of the carbon-nitrogen bond of the β-lactam ring of nitrocefin.

In spite of the unexpected high efficacy of acylation, the antibacterial activity of nitrocefin was less than that of imipenem. This observation suggests that the irreversibility of the acylation reaction, rather than its efficacy, is important for antibacterial activity. Optimum antibacterial activity may therefore involve an additional reaction step that follows the acylation and prevents its reversibility. For carbapenems, this additional reaction step could be the protonation of the nitrogen of the β-lactam ring, as discussed above²³. For most cephalosporins,

irreversibility could result from the loss of a leaving group in the C³ side-chain (Fig. 1)^{14,22}. Of note, other types of rearrangements are instrumental to the inhibition of β -lactamases by clavulanate⁵.

In summary, our study shows that the strain of the β -lactam ring is not sufficient to prevent recyclization following acylation of Ldt_{fm}. Drug efficacy may in part rely on a secondary modification of the drug in the active site to prevent the reversibility of the acylation step. These observations illustrate the complexity of parameters that should be taken into account for the optimization of β -lactams.

Methods

Materials. Stock solutions (25 mg/ml) of nitrocefin (Calbiochem) were prepared in 100% DMSO and stored at -20°C . Dilutions were extemporaneously prepared in water. Stock solutions of imipenem (provided by Merck) were prepared extemporaneously in water.

Protein purification. The plasmid constructs for production of the catalytic domain of Ldt_{fm} (residues 341 to 466) and of a soluble fragment of the β -lactamase BlaC from *M. tuberculosis* (residues 39 to 306) have been previously described^{14,24}. BlaC and Ldt_{fm} were produced in *E. coli* BL21 (DE3) cells grown in brain heart infusion broth (BHI)^{14,24}. Labeling of Ldt_{fm} was obtained in M9 minimal medium containing ¹³C glucose and ¹⁵N NH₄Cl¹⁴. Ldt_{fm} and BlaC were purified from clarified lysates by affinity and size-exclusion chromatography in 100 mM sodium phosphate buffer (pH 6.4) and in 25 mM Tris-HCl (pH 7.5) containing 300 mM NaCl, respectively^{24,25}. The purified enzymes were concentrated by ultrafiltration (Amicon Ultra-4 centrifugal filter devices, Millipore) and stored at -65°C in the same buffers.

Spectrophotometry. All kinetics were performed at 20°C in 100 mM sodium phosphate (pH 6.0) either in a spectrophotometer (Cary 100; Varian SA) coupled to a stopped-flow apparatus (RX-2000; Applied Photophysics) or in a spectrophotometer (Cary 300) equipped with an automatic sample changer (Varian SA).

Mass spectrometry. The formation of Ldt_{fm}- β -lactam adducts was tested by incubating Ldt_{fm} with β -lactams at 20°C in 5 mM sodium phosphate buffer (pH 6.0). Five μl of acetonitrile and 1 μl of 1% formic acid were extemporaneously added, and the reaction mixture was directly injected into the mass spectrometer (Qstar Pulsar I; Applied Biosystem) at a flow rate of 0.05 ml/min (acetonitrile, 50%, water, 49.5%, and formic acid, 0.5%; per volume). Spectra were acquired in the positive mode, as previously described¹¹.

Antibiotic susceptibility testing. The microdilution assay was performed in BHI broth containing 32 $\mu\text{g}/\text{ml}$ of ampicillin (Euromedex) in 96-well plates. Each well (200 μl) was inoculated with overnight cultures to obtain 5×10^5 cfu/ml. Plates were incubated for 24 h at 37°C and the minimal inhibitory concentration was defined as the minimal concentration that prevented visible growth.

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Author Contributions

Z.E. performed the experiments. J.-E.H. and M.A. designed the study. All authors participated to data analysis and to the writing of the manuscript.

Additional Information

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