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Azetidinimines as a novel series of non-covalent broad-spectrum inhibitors of β -lactamases with submicromolar activities against carbapenemases of classes A, B and D.

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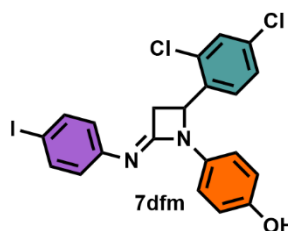
ABSTRACT

The rise of resistances in Gram negative bacteria is reaching an extremely worrying situation and one of the main causes of resistance is the massive spread of very efficient β -lactamases, which render most β -lactam antibiotics useless. Herein, we report the development of a series of imino-analogs of β -lactams (namely azetidinimines) as efficient non-covalent inhibitors of β -lactamases. Despite the structural and mechanistic differences between serine- β -lactamases KPC-2 and OXA-48 and metallo-betalactamase NDM-1, all three enzymes can be inhibited at a submicromolar level by compound **7dfm**, which can also repotentiate imipenem against a resistant strain of *Escherichia coli* expressing NDM-1. We show that this compound

can efficiently inhibit not only the three main clinically-relevant carbapenemases of Ambler classes A, B and D, but also β -lactamases of all four classes (A, B, C and D). Our results pave the way for the development of a new structurally original family of non-covalent broad-spectrum inhibitors of β -lactamases.

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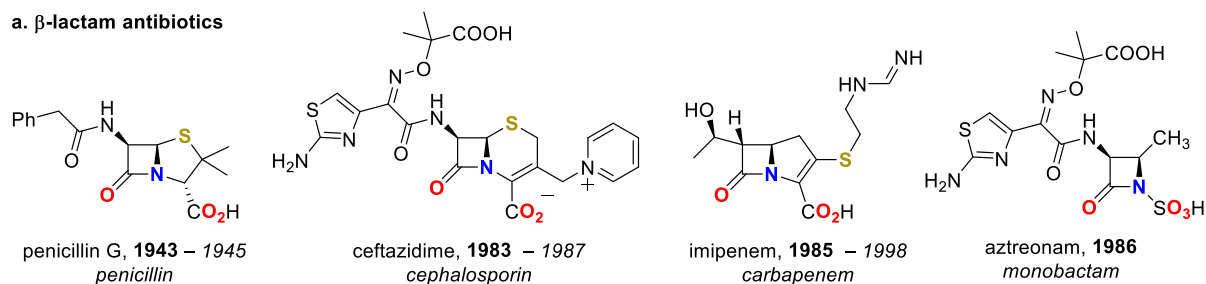
	IC₅₀
<i>NDM-1, class B (metallo-β-lactamase) :</i>	0.1 μM
<i>KPC-2, class A carbapenemase :</i>	0.4 μM
<i>OXA-48, class D carbapenemase :</i>	0.6 μM



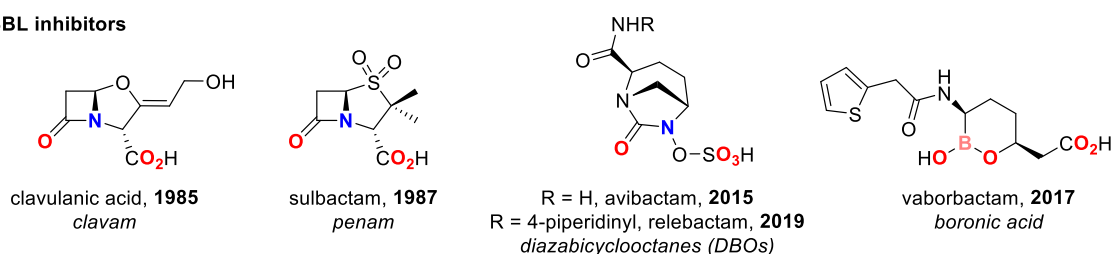
The discovery of penicillin was the start of a golden age for antibiotherapy that is now threatened by exponentially increasing antibioresistance phenomena.¹ Because β -lactams (Figure 1a) have been, and still are, the most prescribed antibiotics worldwide, resistance against them is particularly alarming. Gram negative bacteria (GNB), the main mechanism of β -lactam resistance is due to the production of β -lactamases, enzymes capable of hydrolyzing β -lactams. Even carbapenems, the most powerful β -lactams, are not spared by metallo- β -lactamases (MBLs) such as NDM-1, IMP-1 or VIM-1 (class B) and/or by some clinically worrisome serine β -lactamases (SBLs) such as KPC-2 or OXA-48 (classes A and D, respectively).^{2,3} The discovery of novel antibiotics acting on novel targets is difficult to foresee and strategies to overcome β -lactam resistance^{4,5} especially *via* enzyme inhibition^{6,7} may preserve our current therapeutic arsenal. This strategy was implemented more than 30 years ago with the development of β -lactamase inhibitors (BLIs) including clavulanic acid or sulbactam (Figure 1b), but was not actively pursued until recently. Since 2012 several new broad-spectrum inhibitors of class A and class C β -lactamases have emerged. Of significant interest, avibactam^{8,9} and vaborbactam¹⁰ (Figure 1b) were approved by the FDA for clinical use whilst many of their congeners, in particular diazabicyclooctanes,¹¹⁻¹⁵ are currently going through preclinical or clinical development. These compounds are mainly able to efficiently inhibit SBLs, including carbapenemases, of class A, C and sometimes D, but generally do not inhibit MBLs (class B). In parallel, continuous efforts for the development of efficient MBL inhibitors¹⁶ have recently led to the identification of promising molecules that can bind to zinc atoms of the active site of MBLs, such as thiols¹⁷ – including the clinically available antihypertensive agent L-captopril¹⁸ –, aspergillomarasmine A,¹⁹ rhodanines²⁰ and their thienolate derivatives²¹ or heteroaryl-carboxylic acids such as ANT431²² (Figure 1c). Yet there is still no MBL inhibitor available for clinical use. Moreover, the emergence of bacterial isolates

producing two or even three different carbapenemases of different classes now dictates the development of inhibitors capable of simultaneously inhibiting SBLs and MBLs. So far, apart from some polyphenolic derivatives with moderate activities,²³ only boronic acids have been reported to efficiently inhibit both SBLs and MBLs (Figure 1d).

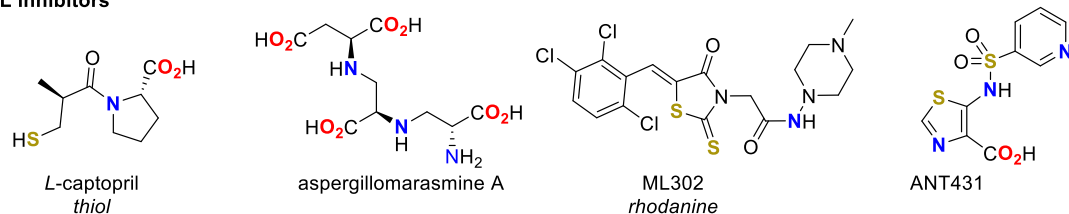
a. β -lactam antibiotics



b. SBL inhibitors



c. MBL inhibitors



d. dual MBL/SBL inhibitors

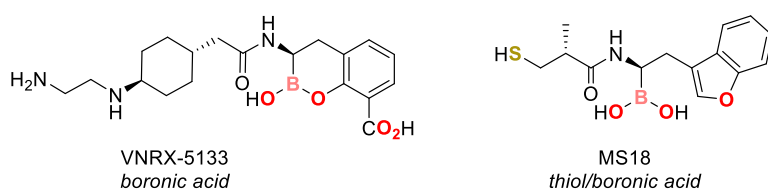


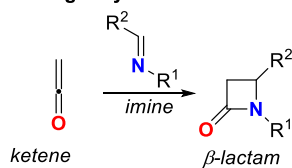
Figure 1. Representative structures of (a) β -lactam antibiotics (in **bold**: year of entry to the market; in *italic*: year of resistance appearance); (b) serine- β -lactamase inhibitors (in **bold**: year of entry to the market); (c) metallo- β -lactamase inhibitors; (d) dual serine-/metallo- β -lactamase inhibitors.

Rigid cyclic analogues of vaborbactam such as VNRX-5133 (currently in phase 3 clinical trials) were the first molecules shown to inhibit all classes of β -lactamases (Figure 1d).^{24–27} VNRX-5133 exhibits submicromolar inhibitory activity against most MBLs and SBLs, being only slightly less active against IMP-1 (class B) and OXA-48 (class D). Very recently, thiol/boronic acid hybrids (such as MS18) were developed and demonstrated interesting dual inhibition properties against SBLs and MBLs.²⁸ MS18 and its congeners also possess a rather broad scope but show limited effects against NDM-1 (class B) and OXA-48.

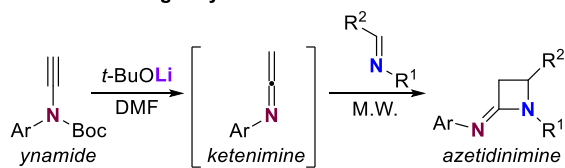
In this context, we sought to develop novel inhibitors that could block the activities of both SBLs and MBLs with comparable efficiency, specifically targeting the three most clinically-relevant carbapenemases: KPC-2 (class A), NDM-1 (class B) and OXA-48 (class D).

We addressed this challenge by exploring uncharted chemical space around the β -lactam nucleus. Synthetically, this four-membered ring can be obtained by a [2+2] cycloaddition between a ketene and an imine,²⁹ a reaction named the Staudinger synthesis after its discoverer (Scheme 1a).³⁰ We recently reported that carefully substituted ynamides^{31,32} can be used as precursors for the *in situ* generation of ketenimines^{33,34} under mild conditions, which can be intercepted by various heterocyclic nucleophiles³⁵ or can undergo a microwave assisted [2+2] cycloaddition with imines.³⁶ By replacing the ketene (C=C=O) with the *in situ* generated iminoketene (C=C=NR), the reaction directly led to an azetidimine (Scheme 1b). Such imino- β -lactams have only been rarely studied with respect to their synthesis^{37,38} and never with respect to their biological and particularly, antibiotic activities. Taking into account their structural similarities with β -lactams, we thus decided to evaluate their therapeutic potential against carbapenemases.

a. Staudinger synthesis



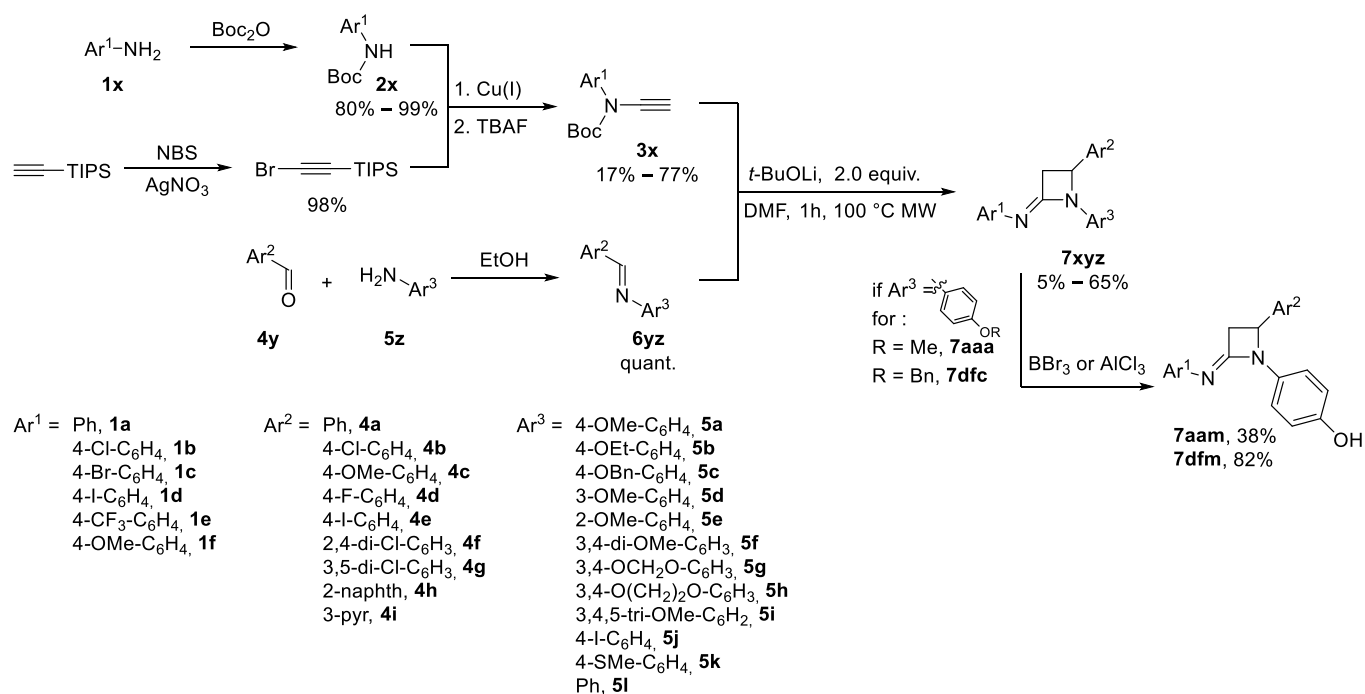
b. imino-Staudinger synthesis



Scheme 1. (a) Staudinger synthesis to access β -lactams; (b) imino-Staudinger synthesis to access azetidinimines.

RESULTS AND DISCUSSION

Chemistry. Taking into consideration the scope and the fact that only aryl groups can be easily incorporated in our previously developed methodology,³⁶ we devised a convergent synthetic plan to access as many structural variations as possible on the azetidinimine scaffold. The Ar¹ group originated from aniline **1x**, which was protected by a Boc group to give **2x** (Scheme 2). Coupling of **2x** with brominated triisopropylsilylacetylene provided ynamide **3x** after TBAF-promoted desilylation. Condensation of benzaldehyde **4y** with aniline **5z** afforded imines **6yz**.



Scheme 2. General synthetic plan to prepare tri-arylated azetidinimines **7xyz**.

The key [2+2] cycloaddition was performed under microwave heating and gave azetidinimines **7xyz** with concomitant loss of the Boc protecting group. For *para*-methoxy derivatives **7aaa**

and **7aba**, as well as *para*-benzyloxy derivative **7dfc**, their corresponding *para*-phenols **7aam**, **7abm** and **7dfm** were obtained by ether cleavage using BBr₃ (for OMe) or AlCl₃ (for OBn). More than forty compounds were prepared using this route.

First results. The first compound evaluated was **7aaa**, bearing a *p*-anisyl group on the endocyclic nitrogen. This aryl group was initially chosen for synthetic reasons as it would electronically favor the reaction and could also be easily derivatized. Initial assessment showed that in the presence of 10 μM of **7aaa**, the hydrolysis of imipenem by NDM-1 was inhibited by 90% with an IC₅₀ estimated to be between 2 and 5 μM (Table 1). This very encouraging initial result prompted us to try and rationalize this inhibitory activity by performing *in silico* molecular modeling studies (Figure 2). It appeared that the azetidinimine does not behave like a β-lactam, whose carbonyl group is chelated by the zinc ions during the hydrolysis process. Here, the methoxy group is coordinated by both zinc ions in the active site and the four-membered ring acts as a scaffold to position the two phenyl rings in hydrophobic regions of the enzyme active site. The Ar¹ phenyl substituent is surrounded by Ser251, Asp212, Ala 215 and Ser217 for both enantiomers, whereas the Ar² phenyl substituent is positioned near Asn220 in the *R* enantiomer and in the proximity of Val73, Ile 35, Met67 and Phe70 in the *S* enantiomer (Figure 2). The two enantiomers of **7aaa** were separated by chiral chromatography and then tested separately, both showing similar inhibition of NDM-1.

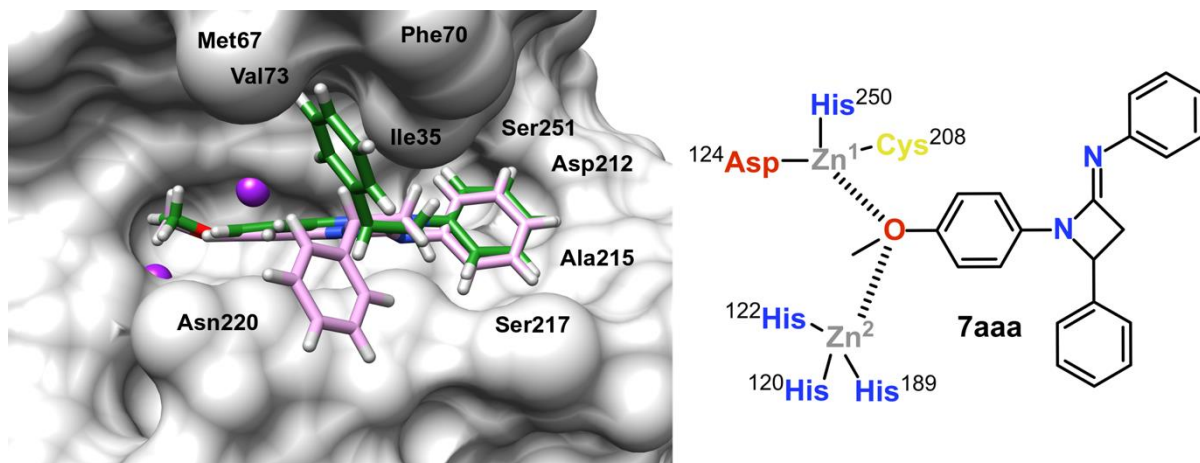
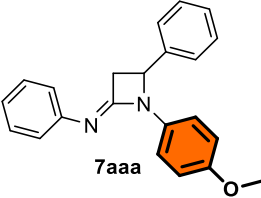
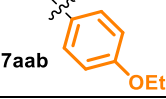
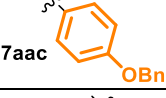
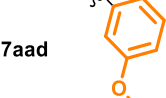
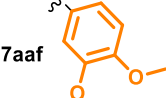
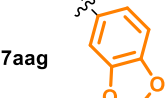
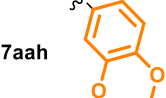
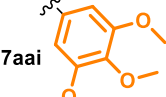

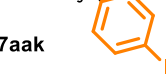




Figure 2. Docking of compound **7aaa** in the active site of NDM-1 (left) and schematic drawing of its key interactions with the zinc ions (right). The enantiomers *R* and *S* of **7aaa** are colored in pink and green, respectively, and the zinc ions are colored in purple.

SAR for enzyme inhibitory activities. A systematic structure-activity relationship was then undertaken by varying the aromatic groups around the azetidinimine nucleus and targeting three carbapenemases: NDM-1, OXA-48 and KPC-2. In addition to its NDM-1 inhibitory activity, **7aaa** was found to be moderately active against KPC-2 but did not have any activity against OXA-48 (Table 1). Variations of the Ar³ ring showed that minimal modifications such as replacing the methoxy of **7aaa** by an ethoxy (**7aab**) or a benzyloxy (**7aac**) led to an improvement of the activity against all three enzymes, **7aac** in particular displaying a sub-micromolar activity against NDM-1 (0.4 μM). Similar activities to **7aab** and **7aac** were observed when the methoxy group was in the meta position (**7aad**). However, the presence of two methoxy groups (**7aaf**), or a benzo[d][1,3]dioxole moiety (**7aag**), led to a slight decrease of efficiency against NDM-1 and a loss of activity for OXA-48 and KPC-2. This was partially restored by incorporation of a 2,3-dihydrobenzo[b][1,4]dioxane, as **7aah** was found to be quite similar to **7aad** in terms of inhibition profile. While 3,4,5-trimethoxy derivative **7aai** only maintained a low activity for NDM-1, aniline-derived compound **7aaj** was found to inhibit all three enzymes with IC₅₀'s of 10.0, 7.0 and 3.5 μM, respectively. The incorporation of an iodine atom (**7aak**) on the *para* position led to solubility issues preventing data from being obtained against OXA-48 and KPC-2 although the anti NDM-1 activity could nevertheless be evaluated to be in the 4.0-5.0 μM range. The thiomethyl derivative **7aal** was also poorly soluble in water, but its strong affinity for the active site Zn ions led to a high inhibition of NDM-1. Finally, the free phenol **7aam** was moderately active against KPC-2 and OXA-48 but again exhibited high inhibitory properties against NDM-1 (0.8 μM). To conclude in this series of Ar³ variations, the best *pan*-carbapenemase inhibitors were *p*-oxyphenyl compounds and especially those bearing a benzyloxy (**7aac**) and a free hydroxy (**7aam**).

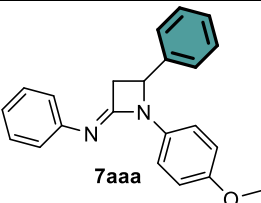
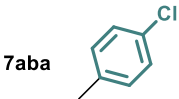
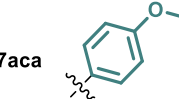
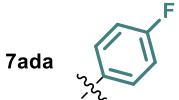
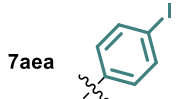
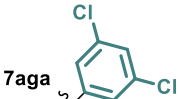
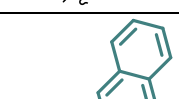
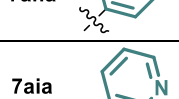
Table 1. Influence of Ar³ on NDM-1, OXA-48 and KPC-2 inhibitory activities compared to 7aaa.

Compound	IC ₅₀ (μM) or % inhibition		
	NDM-1	OXA-48	KPC-2
 7aaa	2.0–5.0	N.E.	55% inhibition at 10 μM
 7aab	2.0–5.0	45% inhibition at 10 μM	2.0–5.0
 7aac	0.4	31% inhibition at 10 μM	1.6
 7aad	1.3	28% inhibition at 10 μM	5.0–10.0
 7aaf	5.0–10.0	9% inhibition at 10 μM	17% inhibition at 10 μM
 7aag	5.0	20% inhibition at 10 μM	6% inhibition at 10 μM
 7aah	1.6	37% inhibition at 10 μM	5.0–10.0
 7aai	78% inhibition at 50 μM	N.E.	N.E.
 7aaj	10.0	7.0	3.5
 7aak	4.0–5	N.D.	N.D.
 7aal	<1.0	15% inhibition at 10 μM	82% inhibition at 20 μM
 7aam	0.8	26% inhibition at 10 μM	45% inhibition at 10 μM

N.E.: no effect (at 10 μM); N. D.: not determined

Keeping a *para*-methoxyphenyl group as Ar³, variations of Ar² were then examined for activity and compared to **7aaa** (Table 2). The *p*-chloro analogue (**7aba**) maintained a similar activity against NDM-1 and though some anti-OXA-48 activity was witnessed, its anti-KPC-2 effect was largely lost. Both *p*-methoxy and *p*-fluoro analogues **7aca** and **7ada** were rather inefficient on all three enzymes while the *p*-iodo, 3,5-dichloro and 2-naphthyl derivatives (**7aea**, **7aga** and **7aha** respectively) presented an interesting profile, being almost equipotent against NDM-1 and KPC-2 but somewhat less effective against OXA-48. In contrast, pyridinyl compound **7aia** was almost inactive on all three enzymes.

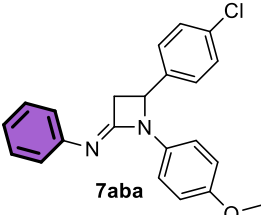
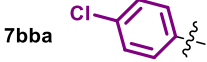
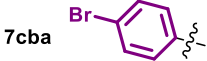

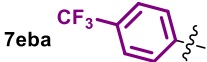
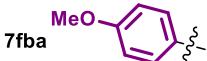
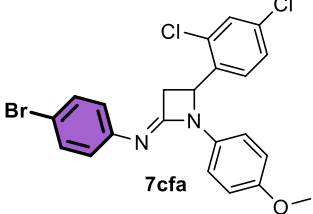

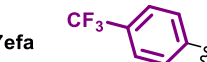
Table 2. Influence of Ar² on NDM-1, OXA-48 and KPC-2 inhibitory activities compared to **7aaa**.

Compound	IC ₅₀ (μM) or % inhibition		
	NDM-1	OXA-48	KPC-2
 7aaa	2.0–5.0	N.E.	55% inhibition at 10 μM
 7aba	2.0–5.0	21% inhibition at 50 μM	44% inhibition at 50 μM
 7aca	67% inhibition at 10 μM	3% inhibition at 10 μM	7% inhibition at 10 μM
 7ada	9% inhibition at 10 μM	N.D.	10% inhibition at 10 μM
 7aea	1.1	33% inhibition at 10 μM	1.1
 7aga	2.0–5.0	15% inhibition at 5 μM	2.0–5.0
 7aha	2.0–5.0	10.0-20.0	5.0-7.0
 7aia	23% inhibition at 100 μM	N.E.	12% inhibition at 100 μM

N.E.: no effect (at 10 μM); N. D.: not determined

From these results, two Ar² groups were selected: *para*-chloro and 2-naphthyl (as in **7aba** and **7aha**). At this stage, we wished to evaluate the influence of Ar¹ in both series (4-chloro in Table 3 and 2-naphthyl in Table 4). It is worth noting that 4-chloro derivatives generally exhibit a greater ease of synthesis due to the superior reactivity of imine **6bz** (or **6fz**) in the [2+2] cycloaddition. Compared to **7aba**, the introduction of a substituent at the *para* position of Ar¹ – whether a chloro (**7bba**), a bromo (**7cba**), an iodo (**7dba**), a trifluoromethyl (**7eba**) and to a lesser extent a methoxy group (**7fba**) – was highly beneficial for the anti-NDM-1 activity, with all compounds possessing IC₅₀ values in the 0.5-0.8 μM range (Table 3).

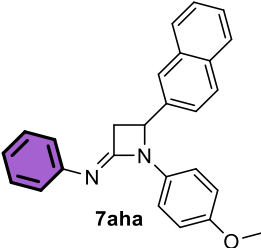
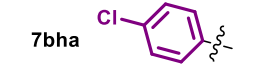
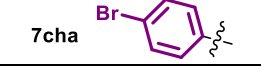
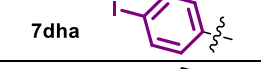
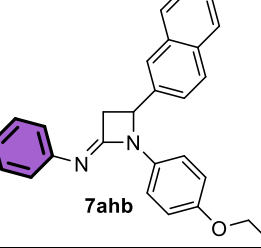
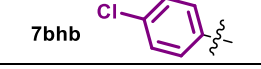
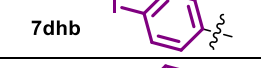

Table 3. Influence of Ar¹ on NDM-1, OXA-48 and KPC-2 inhibitory activities compared to **7aba** and **7cfa**.

Compound	IC ₅₀ (μM) or % inhibition		
	NDM-1	OXA-48	KPC-2
 7aba	2.0–5.0	21% inhibition at 50 μM	44% inhibition at 50 μM
 7bba	0.5	27% inhibition at 10 μM	2.9
 7cba	0.6	7.6	2.4
 7dba	0.7	7.5	2.7
 7eba	0.7	32% inhibition at 10 μM	3.6
 7fba	1.9	31% inhibition at 10 μM	N.E.
 7cfa	1.2	46% inhibition at 10 μM	30% inhibition at 10 μM
 7dfa	0.5	8.5	2.9
 7efa	0.8	5.9	8.7

N.E.: no effect (at 10 μM)

The IC₅₀'s against KPC-2 were also improved with values between 2.0 and 3.6 μM, except for (**7fba**). Additionally, the IC₅₀'s against OXA-48 dropped under the 10 μM threshold for two compounds (**7cba** and **7dba**). A similar trend could be observed with an additional chloro at the 2 position of Ar² (2,4-dichloro series): the *para*-iodo (**7dfa**) and *para*-trifluoromethyl (**7efa**) were found to be the most potent *pan*-inhibitors of carbapenemases NDM-1, OXA-48 and KPC-2 in this series.

Table 4. Influence of Ar¹ on NDM-1, OXA-48 and KPC-2 inhibitory activities compared to **7aha** and **7ahb**.

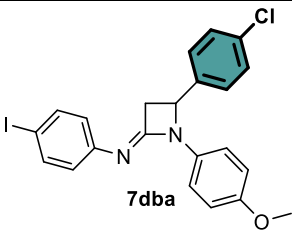
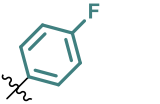
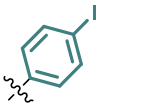
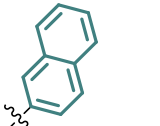
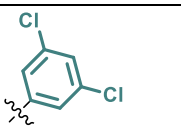
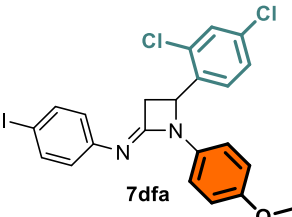
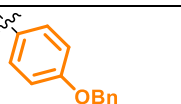
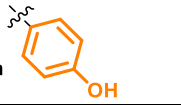
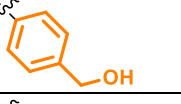
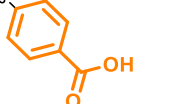
Compound	IC ₅₀ (μM) or % inhibition		
	NDM-1	OXA-48	KPC-2
 7aha	2.0–5.0	10.0-20.0	5.0-7.0
 7bha	1.6	18% inhibition at 10 μM	5.7
 7cha	1.1	3.6	7.5
 7dha	0.5	7.6	2.3
 7ahb	5.0–7.0	N.D.	N.D.
 7bhb	0.6	30% inhibition at 10 μM	2.8
 7dhb	0.5	6.2	2.5
 7fhb	0.6	7.4	2.4

N. D.: not determined

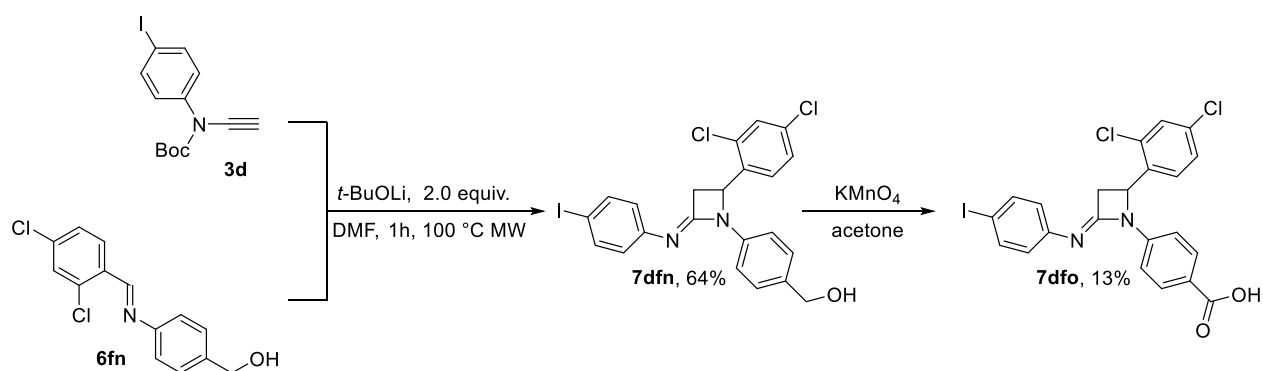
Similarly, excellent anti-NDM-1 activities were observed in the Ar² = naphthyl series (Table 4), with four compounds (**7dha**, **7bhb**, **7dhb** and **7fhb**) having submicromolar IC₅₀'s. Activities against KPC-2 and OXA-48 were also significantly improved, being always under 10 μM except in the case of *para*-chlorinated derivatives **7bha** and **7bhb** (for OXA-48). Despite their high activity, the increased lipophilicity brought by the naphthyl group (for **7dhb** logP_{theor} = 6.94) caused products in this series to be very poorly soluble and to easily form aggregates in the assay media. Thus further studies within this series were stopped.

Having witnessed the highly favorable effect of the *para* substitution on Ar¹ and keeping in mind the importance of the *para*-alkoxy moiety on Ar³, further study of the influence of Ar² was carried out with **7dba** as reference (Table 5). All compounds in this series offered a rather broad-spectrum of activity with most of them inhibiting all carbapenemases with an IC₅₀ below 10 μM and even submicromolar for NDM-1. In contrast with the other compounds, *p*-fluoro (**7dda**), *p*-iodo (**7dea**) and 3,5-dichloro (**7dga**) analogues were more active against OXA-48 than against KPC-2. Finally, keeping Ar¹ = *p*-iodophenyl and Ar² = 2,4-dichlorophenyl (**7dfa**), variations on Ar³ were then further investigated. Surprisingly (compared with **7aaa** and **7aac** in Table 1), the *p*-benzyloxyphenyl derivative (**7dfc**) proved to be less active than **7dfa** (Table 5). **7dfc** was then converted into a free phenol (**7dfm**) by AlCl₃-mediated cleavage of the ether bond (see Scheme 3). This compound exhibited submicromolar IC₅₀ activity against the three enzymes: 0.1 μM for NDM-1 (which could be explained by stronger interactions of the OH function with the zinc ions of the active site), 0.4 μM for OXA-48 and 0.6 μM for KPC-2. While the homologous benzylic alcohol analogue **7dfn** was found to be less active, the corresponding carboxylic acid **7dfo** (obtained by oxidation of **7dfn** with KMnO₄, see Scheme 3) was the most active compound on NDM-1 so far (0.07 μM) but with a complete loss of activity against OXA-48.

Table 5. Influence of Ar² compared to **7dba** and of Ar³ compared to **7dfa** on NDM-1, OXA-48 and KPC-2 inhibitory activities

Compound	IC ₅₀ (μM)		
	NDM-1	OXA-48	KPC-2
 7dba	0.7	7.5	2.7
 7dda	1.0	4.2	9.5
 7dea	0.7	4.4	8.5
 7dha	0.5	7.6	2.3
 7dga	4.5	4.8	42% inhibition at 10 μM
 7dfa	0.5	8.5	2.9
 7dfc	0.8	41% inhibition at 10 μM	5.5
 7dfm	0.1	0.4	0.6
 7dfn	3.0	N.E.	6.0
 7dfo	0.07	N.E.	6.0

N.E.: no effect (at 10 μM); N. D.: not determined



Scheme 3. Synthesis of compounds **7dfn** and **7dfo**.

Complementary assays. Having found a lead (**7dfm**) in our novel azetidinimine BLI series, we then performed complementary assays to ascertain its therapeutic potential. First, it was screened at 10 μ M against a wider panel of BLs (Table 6). At this concentration a complete inhibition of three NDM variants (NDM-4, NDM-7 and NDM-9) and of VIM-1 (Class B) was observed. In contrast, VIM-52 (class B) was not affected. The extended-spectrum β -lactamase CTX-M-15 (class A) and the cephalosporinase CMY-2 (class C) were also inhibited (83% and 86%, at 10 μ M, respectively). These latter results demonstrate that compound **7dfm** can inhibit not only *carbapenemases from 3 classes* but, more generally, **BLs from all four classes**.

Table 6. Additional enzymatic inhibitory activities (% at 10 μ M) for **7dfm**

Compound	NDM-4	NDM-7	NDM-9	VIM-1	VIM-52	CTX-M-15	CMY-2
7dfm	100%	100%	100%	100%	0%	83%	86%

The metabolic stability of **7dfm** was evaluated and the compound was found to possess an excellent stability profile with a Cl_{int} of μ L/min/mg protein in mouse hepatic cells (Table 7). The toxicity of **7dfm** was measured against both normal cells (MRC-5) and cancer cells (HCT-116) and was found to be in the 20-30 μ M range, that is, almost two orders of magnitude higher than its IC_{50} against carbapenemases. Finally, compound **7dfm** was evaluated for the repotentialization of imipenem against the clinical strain of *E. coli* that expresses NDM-1 (amongst other genes of resistance).³⁹ In the absence of inhibitor the MIC was above the resistance

threshold at 16 $\mu\text{g/mL}$, in the presence of 20 μM of **7dfm** it was divided by 2, while using 50 μM of **7dfm** it was divided by 4, bringing it down to the intermediate/resistant limit.

Table 7. Metabolic stability, cytotoxicity and imipenem repotiation for **7dfm**

	Stability		IC_{50} (μM) ^c		MIC ($\mu\text{g/mL}$) ^d		
	Non-NADPH ^a	CL_{int} ^b	MRC-5	HCT-116	0 μM	20 μM	50 μM
7dfm	96%	13.9	19.9	32.3	16	8	4

^a Stability after 45 min in the presence of mouse hepatic microsome in the absence of NADPH; ^b CL_{int} was determined using mouse hepatic microsomes and is given in $\mu\text{L}/\text{min}/\text{mg}$ protein; ^c inhibition of cell proliferation; ^d Minimum inhibitory concentration for imipenem using *E. coli* GUE-NDM1³⁹ clinical strain.

Molecular modeling Compound **7dfm** was selected for a more in depth study of the interaction with the three main clinically-relevant carbapenemases, NDM-1, KPC-2 and OXA-48. The docking of these three enzymes with the two enantiomers of **7dfm** was performed using GOLD⁴⁰ and the results are presented in Figure 4.

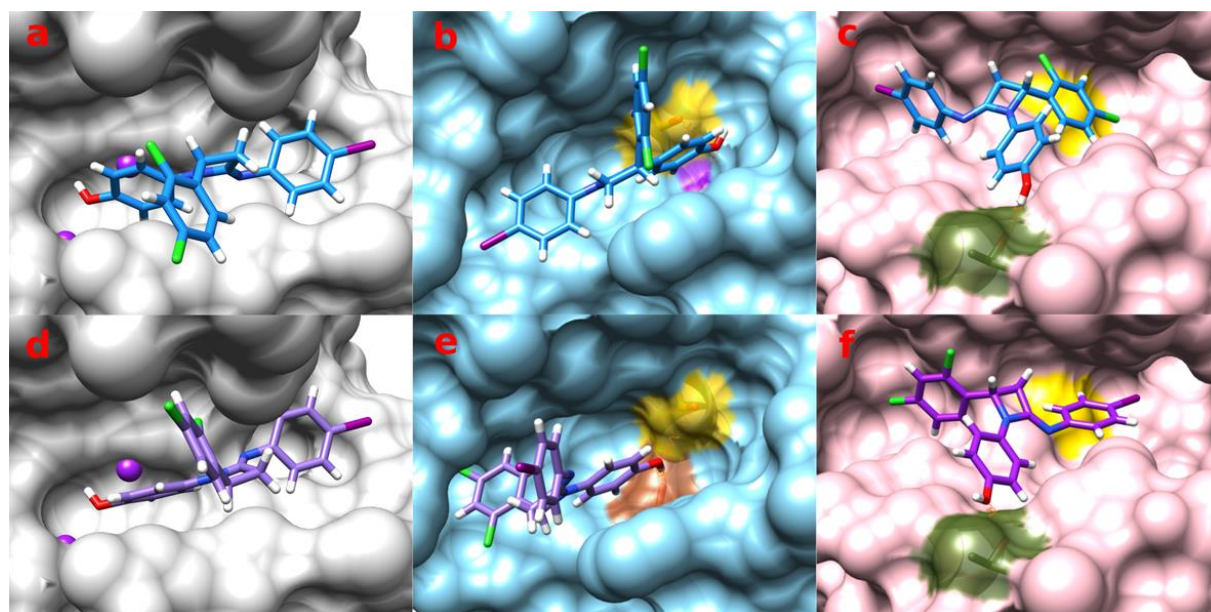


Figure 4: Docking conformations of the enantiomers *R* (a, b, c) and *S* (d, e, f) of compound **7dfm**, in the active site of NDM-1 (class B) represented as gray surface (a, d), KPC-2 (class A) represented as light blue surface (b, e) and OXA-48 (class D) represented as pink surface (c, f). For NDM-1, the zinc ions are colored in purple, for KPC-2 the residues Ser70, Lys73 and Ser130 are colored in yellow, purple and brown, respectively, and for OXA-48 the residues Ser70 and Thr104 are colored in yellow and olive, respectively. Hydrogen bonds are represented as springs colored in orange.

With NDM-1, both enantiomers of **7dfm** interact in a similar manner as **7aaa** (see Figure 2). The phenol moiety is coordinated with the two zinc ions and the 4-iodo-phenyl substituent is positioned in the same subpocket in both cases, possibly establishing a stabilizing halogen bond with the side chain of Asp212. The 2,4-dichloro-phenyl substituent is positioned in a hydrophobic environment bordered by residues Val73, Ile35, Met67 and Phe70 for the *S* enantiomer (Figure 4d) and in the vicinity of Asn220 for the *R* enantiomer (Figure 4a).

The docking pose of KPC-2 with the *R* enantiomer (Figure 4b) shows two hydrogen bonds between the phenol group of **7dfm** and the side chains of Ser70 and Lys73. The 4-iodo-phenyl substituent is positioned in a subpocket formed by the side chains of Asn218, His219 and Glu276 and may form a halogen bond with the side chain of Asn218, whereas the 2,4-dichloro-phenyl substituent is more solvent exposed. The *S* enantiomer makes hydrogen bonds with the side chains of Ser70 and Ser130 through the phenol substituent (Figure 4e), and the positions of the other two substituents are inverted compared with the *R* enantiomer.

The interaction of OXA-48 with the *R* enantiomer of **7dfm** (Figure 4c) shows a hydrogen bond between the phenol group and the side chain of Thr104, at the upper extremity of the binding site, whereas the two other substituents are positioned more deeply in the binding site groove. The *S* enantiomer establishes the same hydrogen bond between the phenol group and the side chain of Thr104, with the positions of the 4-iodo-phenyl and 2,4-dichloro-phenyl substituents inverted.

Conclusions

The evaluation of azetidinimines, imino-analogues of β -lactams, as β -lactamase inhibitors led to the development of a new family of non-covalent carbapenemase inhibitors. Structural

optimization led to the identification of phenol **7dfm** as the lead compound, which can strongly inhibit MBL NDM-1 (0.1 μM), class A SBL KPC-2 (0.4 μM), and class D SBL OXA-48 (0.6 μM). Molecular modeling showed that this compound does not mimic a β -lactam within the enzyme active sites and that both enantiomers can interact with the enzymes. Provided one key interaction (such as complexation with the zinc ions in NDM-1 or hydrogen bonds in KPC-2 and OXA-48) is established, the rigid and compact structure of the central four-membered ring indeed allows the other aryl groups to be allocated almost interchangeably in the enzyme pocket to increase the affinity. Further refinement of the structure of these molecules will be pursued to improve the efficiency of these β -lactamase *pan* inhibitors. The metabolic stability of the hit compound and its efficient repotentialization of imipenem against a resistant *E. coli* strain will serve as the basis for future *in vivo* studies.

Materials and Methods

General Procedure for Azetidimine Synthesis. The imine (1.0 equiv.), the ynamide (2.0 equiv.) and the additive – silica gel (1.0 equiv.) or ZnOTf_2 (10 mol%) – were successively added in a microwave sealable tube and placed under argon before the addition of *t*-BuOLi – solid or 2.2 M in solution in THF – (2.0 equiv.) followed by extra dry DMF (0.3M). The sealed tube was placed in a microwave apparatus for 1 h at 100 °C. The crude material was purified by flash chromatography on silica gel or with preparative TLC. See Supporting Information for detailed synthetic procedures and characterizations of the compounds.

***In vitro* β -lactamase inhibition assay.** IC_{50} values were determined against the panel of purified carbapenemases: OXA-48, KPC-2 and NDM-1 by spectrophotometric assay, using ULTROSPEC 2000 UV spectrophotometer and the SWIFT II software (GE Healthcare, Velizy-

Villacoublay, France). Compounds were dissolved in DMSO stock solutions at 10 mM; more dilute stocks were subsequently prepared as necessary by dissolving them also in DMSO. Assay conditions were as follows: 100 mM phosphate buffer, pH 7 (supplemented with 50 μ M Zn²⁺ when testing NDM-1, and with 50 mM NaHCO₃ when testing OXA-48), 100 μ M imipenem (Sigma-aldrich, Saint-Quentin Fallavier, France). The reaction was monitored at 297 nm, time course 600 seconds at 25°C with 3 min of incubation (compound/carbapenemase). Each inhibitor compound was assayed at seven different concentrations, in triplicate for calculating an error value with 95% confidence interval ($p < 0.05$). IC₅₀ values were determined using the equation $IC_{50} = ((1/0.5 \times v_0) - m)/q$, where v_0 is the rate of hydrolysis of the reporter substrate (v_0 being the rate measured in the absence of inhibitor), q the y axis intercept and m the slope of the resulting linear regression. Percentage of inhibition were obtained with a concentration of 10 μ M of compound.

Minimal Inhibition concentrations. MIC values were determined by broth microdilution, in triplicate, in cation-adjusted Mueller Hinton broth according to the Clinical Laboratory Standards Institute (CLSI, <https://clsi.org/>) guidelines. The enterobacterial clinical strain *E. coli* NDM-1 GUE expressing the carbapenemase NDM-1 was used.³⁹ Experiments were performed in microtiter plates containing the medium with imipenem and inhibitors (dissolved in DMSO). Three inhibitor concentrations were tested: 50, 100 and 200 μ M. Plates were incubated overnight at 37°C for 18–24 h.

Incubations in hepatic microsomes. Compounds (5 μ M) were incubated in 0.5 mg/mL of pooled male mouse liver microsomes (from Biopredic, France), in 0.1 M phosphate buffer at pH 7.4 at 37 °C. After prewarming the mixture for 5 min, reactions were initiated by the addition of NADPH (1 mM). Incubations (400 μ L) were performed at 37 °C for 0, 5, 15, 30 and

45 min in duplicate and the reaction immediately terminated by adding 200 μL of cold acetonitrile. Samples (including the control to evaluate non NADPH-dependent stability) were centrifuged and the supernatant fractions analyzed by UPLC-MS/MS with multiple reaction monitoring (MRM). Diphenhydramine was used as positive control in mouse liver microsomes test. The MRM area response of the analyte was set to 100% with the T0 incubation, the relative decrease in MRM area ratio intensity over time against that of the control (percent parent decrease) was used to determine the half-life ($t_{1/2}$) of compounds in the incubation. Half-life values were calculated from the relationship:

$T_{1/2}$ (min) = $0.693/k$, where k is the slope of the \ln concentration vs time curve. The intrinsic clearance (CL_{int}) was calculated as: $CL_{int} = (0.693 \times \text{incubation volume } (\mu\text{L})) / (t \text{ (min)} \times \text{mg of microsomal protein})$.

Cell culture and proliferation assay. Assays were carried out at the Institut de Chimie des Substances Naturelles by the CIBI screening platform. Cell lines were obtained from the American type Culture Collection (Rockville, USA) and were cultured according to the supplier's instructions. Briefly, human MRC-5 cells were grown in DMEM supplemented with 10% fetal calf serum (FCS) and 1% glutamine and HCT116 colorectal carcinoma cells were grown in RPMI 1640 containing 10% FCS and 1% glutamine. All cell lines were maintained at 37 °C in a humidified atmosphere containing 5% CO_2 . Cell growth inhibition was determined by an MTS assay according to the manufacturer's instructions (Promega, Madison, WI, USA). Briefly, the cells were seeded in 96-well plates (2.5×10^3 cells/well) containing 200 μL of growth medium. After 24 h of culture, the cells were treated with the test compounds at different final concentrations. After 72 h of incubation, 40 μL of resazurin was added for 2 h before recording absorbance at 490 nm with a spectrophotometric plate reader. The IC_{50} value

corresponded to the concentration of compound inducing a decrease of 50% in absorbance of drug-treated cells compared with untreated cells. Experiments were performed in triplicate. Paclitaxel was used as the reference compound.

PCR, Cloning, Expression, and DNA Sequencing. Whole-cell DNA of the enterobacterales expressing β -lactamases NDM-1, NDM-4, NDM-7, NDM-9, VIM-1, VIM-52, CTX-M-15, CMY-2 and OXA-48 was extracted, using the QIAamp DNA mini kit (Qiagen, Courtaboeuf, France) and used as a template to amplify all the different genes. The sequences without the peptide signal (predicted by SignalP 4.1 Server, <http://www.cbs.dtu.dk/services/SignalP-4.1/>) encoding for the mature protein, were obtained by PCR amplification, using the forward primers, which included an *NdeI* restriction site, and the reverse primer which included an *XhoI* restriction site and a deletion of the stop codon of the gene to allow the expression of an C_Term His tag. Then, PCR product was cloned into pET41b vector (Invitrogen[®], Life Technologies, Cergy-Pontoise, France), using *NdeI* and *XhoI* restriction enzymes, to obtain a C-Term His_g-tag. The accuracy of the recombinant plasmid was verified by sequencing, using a T7 promoter and T7 terminator with an ABI Prism 3100 automated sequencer (Applied Biosystems, Thermo Fisher Scientific, Les Ulis, France). The nucleotide sequences were analyzed by using software available at the National Center for Biotechnology Information website (<http://www.ncbi.nlm.nih.gov>).

Protein purification. An overnight culture of *E. coli* BL21 DE3 harboring recombinant pET41b plasmids was used to inoculate 2 L of LB medium broth containing 50mg/L kanamycin. Bacteria were cultured at 37 °C until an OD of 0.6 at 600 nm was reached. The expression of the β -lactamase genes was carried out overnight at 22 °C with 0.2 mM IPTG as inducer. Cultures were centrifuged at 6000 g for 15 min and then the pellets were resuspended with the binding

buffer (10 mM imidazole, 25 mM sodium phosphate pH 7.4 and 300 mM NaCl). Bacterial cells were disrupted by sonication and the bacterial pellet was removed by two consecutive centrifugation steps at 10000g for 1h at 4 °C; the supernatant was then centrifuged at 96000g for 1h at 4 C. The soluble fractions were filtered and then passed through a HisTrap™ HP column (GE Healthcare) and proteins were eluted with the elution buffer (500 mM imidazole, 25 mM sodium phosphate pH 7.4 and 300 mM NaCl). Finally, a gel filtration step was performed with 100 mM sodium phosphate buffer pH 7 and 150 mM NaCl with a Superdex 75 column (GE Healthcare). The protein purity was estimated by SDS–PAGE. The pooled fractions were dialyzed against 10 mM Tris-HCl pH 7.6, for NDM and VIM 50 μM of ZnSO₄ was added, then concentrated using Vivaspin columns (Sartorius, Aubagne, France). The concentrations were determined by measuring the OD at 280 nm and with the extinction coefficients obtained from the ProtParam tool (Swiss Institute of Bioinformatics online resource portal).⁴¹

Molecular modeling. The three-dimensional structure of compound **7aaa**³⁶ was retrieved from the Cambridge Structural Database⁴² (CSD refcode KEMJEU) and that of compound **7dfm** were built starting from **7aaa** by manual editing using UCSF Chimera package.⁴³ The structures of enantiomers were generated using an in-house script. Molecular docking was performed using the GOLD suite⁴⁰ (CCDC) and the GoldScore scoring function, with the structures 4HL2,⁴⁴ 2OV5⁴⁵ and 4S2P⁴⁶ as receptors for NDM-1, KPC-2 and OXA-48, respectively. The binding sites were defined as 15 Å radius spheres centered on the Zn1 ion for NDM-1 and on the OG atom of Ser70 for KPC-2 and OXA-48. In agreement with our previous studies^{47–53} showing that an enhanced conformational search is beneficial, especially for large molecules, a search efficiency of 200% was used to better explore the ligand conformational space. All other parameters were used with the default values. Images were generated using UCSF Chimera.⁴³

Supporting Information

The Supporting Information contains detailed synthetic procedures and characterizations of the compounds and copies of the NMR spectra.

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This study was conceived by K. Cariou, B. I. Iorga, T. Naas and R. H. Dodd. Synthesis and characterization of the compounds were performed by E. Romero, M. Benchekroun, S. Ventre, A. C. A. D'Hollander, K. Vijayakumar and C. Minard. Purification of the proteins, inhibition assays and MIC assays were performed by S. Oueslati with C. Exilie, L. Tlili and A. Zavala. Metabolic studies were performed by L. A. Nguyen under the supervision of A. Pruvost. Structural analysis and molecular modeling studies were performed by A. Zavala, E. Elisée, E.

Selwa, P. Retailleau and B. I. Iorga. K. Cariou and B. I. Iorga drafted the manuscript which was amended and commented on by all authors.

E. R. & S. O contributed equally; † M. B. & A. C. A. D. contributed equally

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Conflict of Interest Disclosure: The authors declare no competing financial interest

List of Abbreviations

CTX M 15: Cefotaximase-Munich 15

CMY-2: Cephamycinase-2

DMSO: dimethyl sulfoxide GNB: Gram-negative Bacilli

HCT 116: Human Colorectal Carcinoma cells

KPC: Klebsiella Pneumoniae Carbapenemase

MBL: Metallo- β -lactamase

MIC: Minimum Inhibitory Concentrations

MRC-5: Medical Research Council cell strain 5

ND: Not Determined

NDM: New Delhi metallo- β -lactamase

NE: no effect

OXA: Oxacillinase

SAR: Structure Activity Relationship

SBL: Serine- β -lactamases

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