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# Structure-Activity Analysis of Peptidic *Chlamydia* HtrA

## Inhibitors

Ayodeji A. Agbowuro,<sup>a</sup> Jimin Hwang,<sup>a</sup> Emma Peel,<sup>b</sup> Rami Mazraani,<sup>c</sup> Alexandra Springwald,<sup>a</sup> James W. Marsh,<sup>d</sup> Laura McCaughey,<sup>d,c</sup> Allan B. Gamble,<sup>a</sup> Wilhelmina M. Huston,<sup>\*,c</sup> Joel D. A. Tyndall<sup>\*,a</sup>

<sup>a</sup> School of Pharmacy, University of Otago, Dunedin, 9054, New Zealand

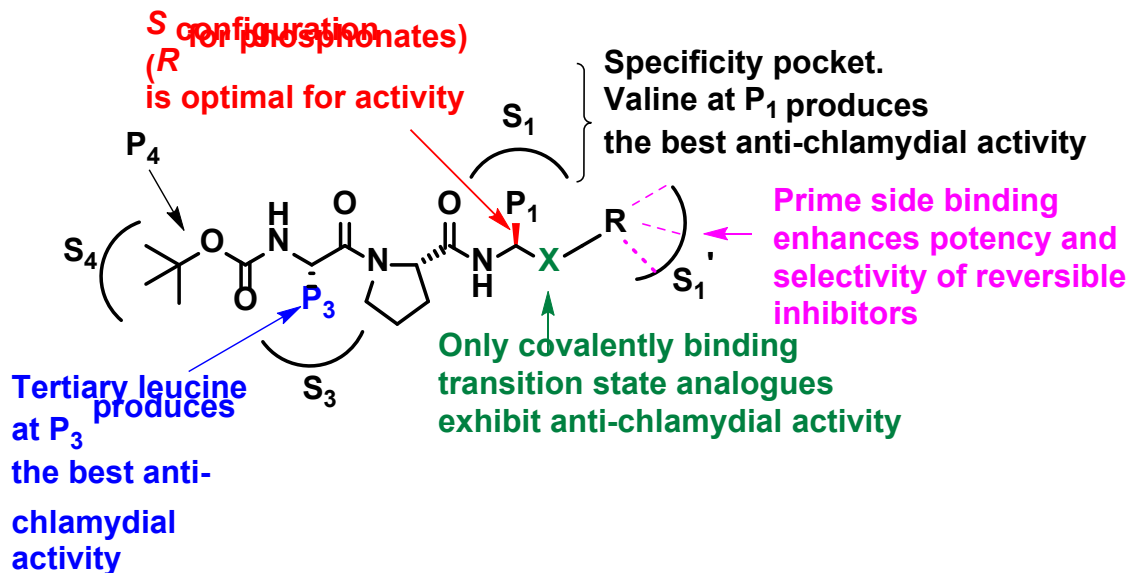
<sup>b</sup> School of Life and Environmental Sciences, The University Sydney, Sydney, NSW 2006, Australia

<sup>c</sup> School of Life Sciences, University of Technology Sydney, 15 Broadway, Ultimo NSW 2007, Australia

<sup>d</sup> The iThree Institute, University of Technology Sydney, 15 Broadway, Ultimo NSW 2007, Australia

<sup>e</sup> Department of Biochemistry, University of Oxford, South Park road, Oxford OX1 3QU, United Kingdom

\*Corresponding authors: [joel.tyndall@otago.ac.nz](mailto:joel.tyndall@otago.ac.nz), [Wilhelmina.Huston@uts.edu.au](mailto:Wilhelmina.Huston@uts.edu.au)



**ABSTRACT:** *Chlamydia trachomatis* high temperature requirement A (CtHtrA) is a serine protease that performs proteolytic and chaperone functions in pathogenic *Chlamydiae*; and is seen as a prospective drug target. This study details the strategies employed in optimizing the irreversible CtHtrA inhibitor JO146 [Boc-Val-Pro-Val<sup>P</sup>(OPh)<sub>2</sub>] for potency and selectivity. A series of adaptations both at the warhead and specificity residues P<sub>1</sub> and P<sub>3</sub> yielded 23 analogues, which were tested in human neutrophil elastase (HNE) and CtHtrA enzyme assays as well as *Chlamydia* cell culture assays. Trypsin and chymotrypsin inhibition assays were also conducted to measure off-target selectivity. Replacing the phosphonate moiety with  $\alpha$ -ketobenzothiazole produced a reversible analogue with considerable CtHtrA inhibition and cell culture activity. Tertiary leucine at P<sub>3</sub> (**8a**) yielded approximately 33-fold increase in CtHtrA inhibitory activity, with an IC<sub>50</sub> = 0.68 ± 0.02  $\mu$ M against HNE, while valine at P<sub>1</sub> retained the best anti-chlamydial activity. This study provides a pathway for obtaining clinically relevant inhibitors.

## 1. Introduction

Members of the phylum Chlamydiae are a unique obligate intracellular bacteria, responsible for a host of diseases with the most notable being *Chlamydia (C.) trachomatis* and *C. pneumoniae*.<sup>1-2</sup> *C. trachomatis* parasitizes only humans, causing complications including ectopic pregnancy, infertility, reversible blindness and infant pneumonia.<sup>3-4</sup> *C. pneumoniae* is the major cause of respiratory infections in children and adults, leading to pneumonia, bronchitis, sinusitis, and it is also a risk factor for atherosclerosis.<sup>5-6</sup> *C. pneumoniae* infects other animals including koalas and horses.

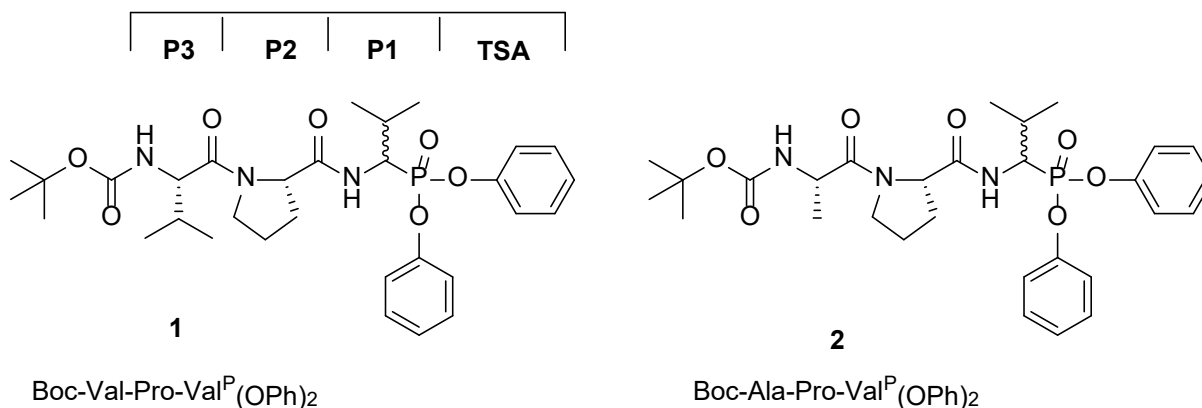
The koala is an iconic Australian marsupial currently listed as vulnerable, with the potential for extinction. It is often plagued by illnesses as a result of *C. pneumoniae* and *C. pecorum* infections.<sup>7-</sup> <sup>8</sup> *C. pecorum* causes distressing ocular and urogenital site infections in the animal,<sup>8</sup> while the (less frequently observed) clinical outcomes of *C. pneumoniae* infection in koala are respiratory in nature and characterized by coughing, sneezing and purulent discharge.<sup>9</sup>

Despite the availability of efficacious antibacterial agents, the management of chlamydiosis remains a challenge. Human chlamydia infections are estimated to have a global incidence of approximately 130 million<sup>10</sup> and clinical treatment failures have been reported<sup>11-12</sup> leading to concern over the effectiveness of current treatment regimens. *C. trachomatis* is often present as a co-infection (*e.g.* with *Neisseria gonorrhoeae* or *Mycoplasma genitalium*) and in these pathogens, resistance to the commonly used anti-chlamydial therapies (macrolides, tetracyclines) is increasingly common,<sup>13-15</sup> hence a drug targeted to *Chlamydia* may reduce the selection of antibiotic resistance in the other pathogens.

We have previously identified the *C. trachomatis* high temperature requirement A (CtHtrA) protein as a serine protease that is expressed at elevated levels under *Chlamydia* stress conditions, which possesses both proteolytic and chaperone capabilities.<sup>16</sup> CtHtrA substrate-specificity studies

revealed a preference for hydrophobic amino acid residues at prime and nonprime positions, with a preference for small hydrophobic residues such as valine (Val) at P<sub>1</sub>, giving the enzyme an elastase-like specificity.<sup>17</sup>

We have previously demonstrated that CtHtrA is essential for *Chlamydia* survival using two selective irreversible CtHtrA inhibitors, JO146 **1** [Boc-Val-Pro-Val<sup>P</sup>(OPh)<sub>2</sub>; IC<sub>50</sub> = 12.5 μM] and JCP83 **2** [Boc-Ala-Pro-Val<sup>P</sup>(OPh)<sub>2</sub>; IC<sub>50</sub> = 47.19 μM; Figure 1] in *in vivo*<sup>18</sup> and *ex vivo*<sup>19</sup> chlamydia models. JO146 was also found to be active against koala *Chlamydia* species (*C. pecorum* and *C. pneumoniae*) with negligible toxicity on koala tissue<sup>19</sup> thus validating CtHtrA as a viable drug target for the development of novel anti-chlamydial drugs.



**Figure 1.** Structures of JO146 (**1**) and JCP83 (**2**) and the P<sub>3</sub>-P<sub>1</sub> protease nomenclature.<sup>18</sup> TSA – Transition State Analogue.

JO146 (**1**, IC<sub>50</sub> = 12.5 μM)<sup>18</sup> is a suitable lead molecule for developing a clinically relevant protease inhibitor for targeting *Chlamydia*. However, it is currently not potent enough, and may be susceptible to chemical and enzymatic degradation due to its peptidic nature. This paper details the strategies employed in optimizing the potency and selectivity of JO146 (**1**). Herein, we describe the design, syntheses and structure activity relationships (SAR) of 23 analogues of JO146 (**1**) tested against both human and koala *Chlamydia* species. The compounds span a series of diphenyl phosphonates as well as reversible transition state analogues; namely α-ketoheterocycles, α-

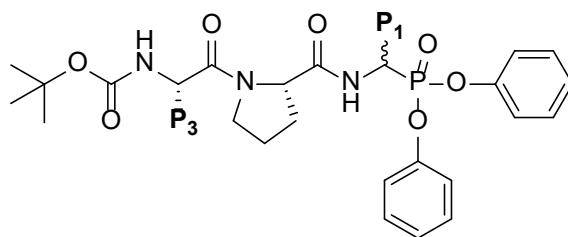
diketones, N-methylamides, valinol,  $\alpha$ -ketoamide, and boronic acid. In addition, the P<sub>1</sub> and P<sub>3</sub> valine residues of JO146 (**1**) were systematically replaced with other natural and non-natural amino acids with the aim of optimizing binding at the respective subpockets of CtHtrA.

## 2. Results and discussion

### 2.1 Development of new CtHtrA inhibitors

The syntheses of targeted tripeptide inhibitors containing various transition state analogues are described in Schemes 1-3, S1 and S2. The diphenyl phosphonates JO146 (**1**) and **8a-h** (Table1) were synthesized in order to identify the optimal subsite specificity for P<sub>3</sub> and P<sub>1</sub>. Compounds **1**, **8a-h** were synthesized according to the procedure reported by Winiarski *et al.*<sup>20</sup> Cbz-protected P<sub>1</sub> precursors **5a-d** were synthesised via an  $\alpha$ -amidoalkylation-type reaction using the appropriate alkyl aldehyde **4a-d**. They were subsequently deprotected to yield the hydrobromide salts **6a-d** using 33% HBr in acetic acid. The final compounds (**1**, **8a-h**) were obtained by coupling **6a-d** to pre-synthesized dipeptide precursors **7a-d** in an HBTU-promoted solution phase peptide coupling. The products were obtained as diastereomeric mixtures and biologically evaluated as such.

**Table 1. Peptidyl diphenyl phosphonate inhibitors of the form Boc-P<sub>3</sub>-Pro-P<sub>1</sub>-(OPh)<sub>2</sub>**

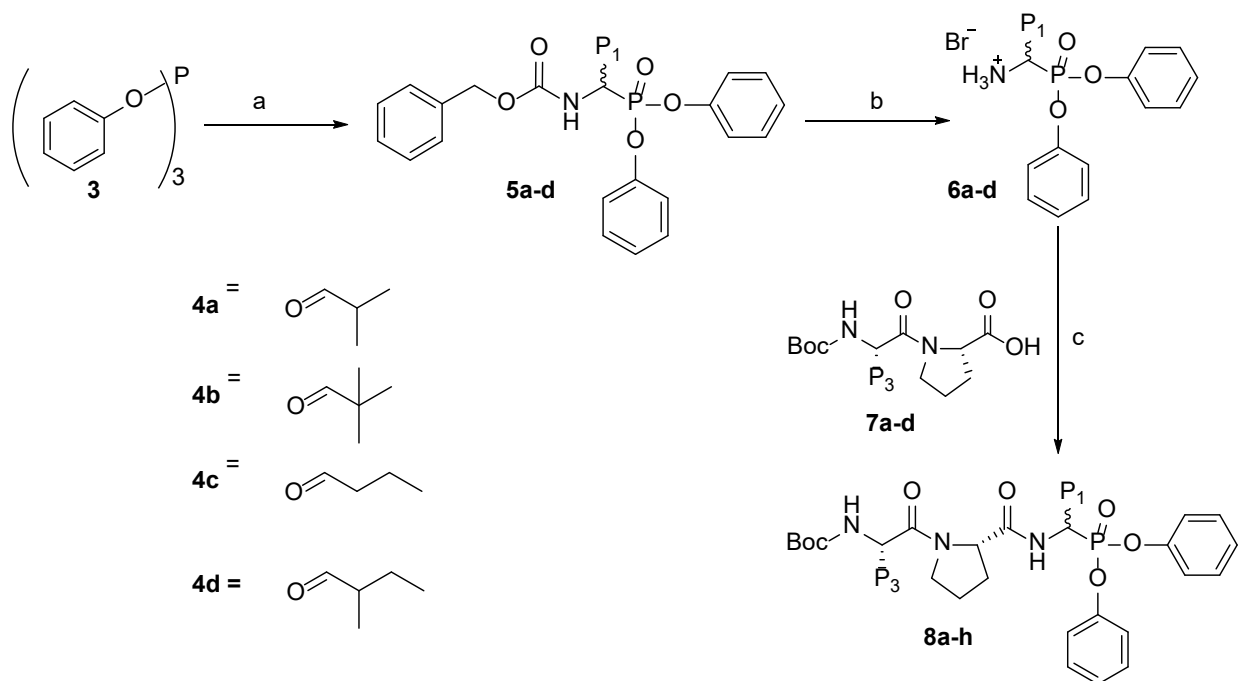


	P <sub>3</sub>	P <sub>1</sub>
JO146 ( <b>1</b> )	Val	Val
<b>2</b>	Ala	Val
<b>8a</b>	Tle	Val
<b>8b</b>	Nva	Val
<b>8c</b>	Ile	Val

<b>8d</b>	Val	Ile
<b>8e</b>	Val	Nva
<b>8f</b>	Val	Tle
<b>8g</b>	Ile	Ile
<b>8h</b>	Tle	Ile

\*A derivative of **1** with Leu at P<sub>1</sub> has previously shown no inhibition (data not shown)

**Scheme 1. Synthesis of the peptidyl diphenyl phosphonates (8a-h)<sup>a</sup>**



<sup>a</sup>Reagents and conditions: (a) Benzylcarbamate, **4a-d**, 80-90 °C, 2 h, 15-72%; (b) 33% HBr/AcOH, 25 °C, 2 h, 92-98%; (c) HBTU, DIEA, DMF, 25 °C, 20 h, 25-77%.

The potential risk of toxicity associated with irreversible inhibition via the diphenyl phosphonate group in JO146 led to the decision to investigate reversible transition state analogues as alternatives. This entailed an exchange of the phosphonate moiety with less reactive groups including  $\alpha$ -ketoheterocycles ( $\alpha$ -ketothiazole,  $\alpha$ -ketobenzothiazole and  $\alpha$ -keto-N-methylimidazole),  $\alpha$ -diketones,  $\alpha$ -ketoamide, boronic acid, and non-reactive N-alkyl methylamides and valinol derivatives.

Potent peptidyl  $\alpha$ -ketoheterocycle inhibitors of the human neutrophil elastase (HNE) as well as thrombin have been reported.<sup>20-21</sup> They possess the ability to form reversible covalent bonds as well as making key interactions with the enzyme active site around P<sub>1</sub>'. Peptidyl  $\alpha$ -ketoheterocycles

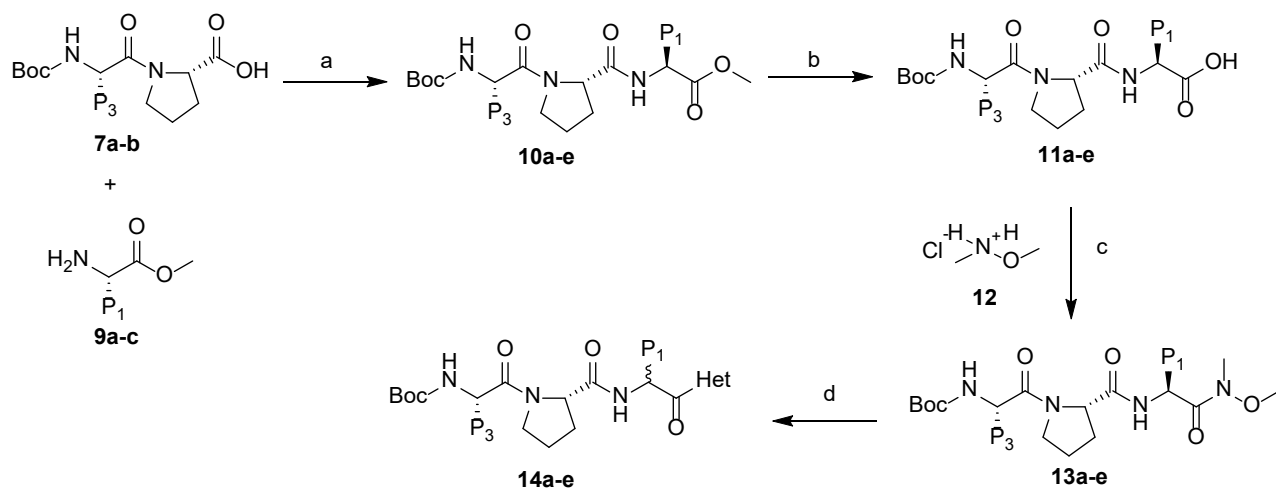


**14a-e** (Table 2), were prepared via Weinreb amides **13a-e** by first coupling the tripeptides **11a-e** to *N,O*-dimethyl-hydroxylamine hydrochloride **12** using HBTU as the reagent (Scheme 2). This was followed by reacting **13a-e** with the respective lithiated heterocycles. The final products **14a-e** were obtained as isomeric mixtures, characterized by additional peaks in the <sup>1</sup>H and <sup>13</sup>C NMR spectra and two distinct peaks in the HPLC trace (see SI), and were likely the result of an *n*-butyllithium-induced epimerization at the alpha-carbon (relative to the heterocycle).

**Table 2. Peptidyl  $\alpha$ -ketoheterocycle inhibitors**

Compound	P <sub>3</sub>	P <sub>1</sub>	Heterocycle
<b>14a</b>	Val	Val	Thiazole
<b>14b</b>	Val	Nva	Thiazole
<b>14c</b>	Val	Val	Benzothiazole
<b>14d</b>	Tle	Ile	Benzothiazole
<b>14e</b>	Val	Val	N-methylimidazole

## Scheme 2. Synthesis of the peptidyl $\alpha$ -ketoheterocycles **14a-e**<sup>a</sup>

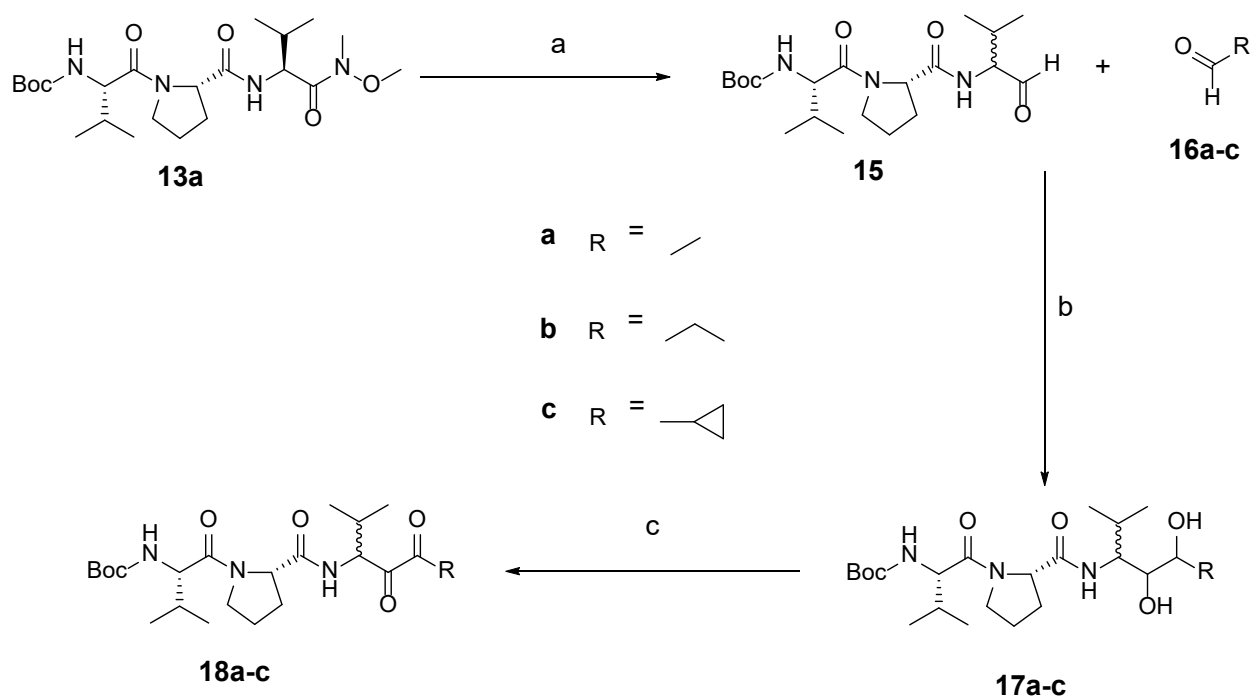


<sup>a</sup>Reagents and conditions: (a) HBTU, DIEA, DMF, 25 °C, 20 h, 45-54%; (b) LiOH.H<sub>2</sub>O, THF, H<sub>2</sub>O, 25 °C, 2 h, 78-90%; (c) HBTU, DIEA, DMF, 25 °C, 20 h, 28-63%; (d) heterocycle, n-BuLi, THF, -78 °C, 28-54%.  $P_3$  = Val or Tle;  $P_1$  = Val, Nva or Ile

Compounds with a 1,2-dicarbonyl moiety such as  $\alpha$ -ketoamides and  $\alpha$ -diketones, are very good serine traps and have been incorporated in potent and safe hepatitis C NS3 protease inhibitors used clinically.<sup>22-24</sup> Previous studies into CtHtrA specificity suggest a preference for hydrophobic residues at  $P_1'$ ,<sup>16-17</sup> identifying glycine and isoleucine as the most favored residues. In the present study, we utilized the ability of  $\alpha$ -diketones to interact reversibly with the  $S_1'$  pocket of CtHtrA and the  $\alpha$ -diketones **18a-c** were synthesized with methyl, ethyl and cyclopropyl substituents at  $P_1'$ . The synthesis of compounds in the  $\alpha$ -diketone series (Scheme 3) was conducted as reported by Han *et al.*<sup>22</sup> The aldehyde **15** was generated according to the method described by Dekhane and Dodd.<sup>25</sup> A solution of the Weinreb amide **13a** in THF was treated with LiAlH<sub>4</sub> under nitrogen, which formed a stable metal complex, restricting further hydride addition that otherwise produces the unwanted alcohol product. Upon a mild acidic workup, the complex dissociated, yielding the

desired aldehyde **15**. After column purification, **15** was recovered as a diastereomeric mixture, as indicated by the additional peaks in the  $^1\text{H}$  and  $^{13}\text{C}$  NMR spectra. Peptide aldehydes are reported to epimerize during purification, and this occurs irrespective of the purification method employed.<sup>26</sup> Compound **15** was then used as the starting material for the synthesis of the 1,2-diols **17a-c** by pinacol cross coupling of **15** with the aliphatic aldehydes **16a-c** as reported by Freudenberger *et al.*<sup>27</sup> Oxidation of the 1,2-diols with Dess-Martin periodinane (DMP) afforded the peptidyl  $\alpha$ -diketones **18a-c**. Compounds **17a-c** and **18a-c** were all obtained as diastereomeric mixtures by virtue of their enantiomerically impure aldehyde precursor (**15**).

**Scheme 3. Synthesis of the peptidyl  $\alpha$ -diketones (18a-c)<sup>a</sup>**



<sup>a</sup>Reagents and conditions: (a) LiAlH<sub>4</sub>, THF, -30 °C, 1 h, 83%; (b) VCl<sub>3</sub>(THF)<sub>3</sub>, Zn, DCM, 25 °C, 2 h, 36-56%; (c) Dess-Martin periodinane (DMP), DCM, 25 °C, 20 min, 20-67%.

The peptidic N-methylamides **24a-d**, and the valinol derivative **28** were synthesized as noncovalent binders of CtHtrA. Compounds **24a-d** were specifically designed with methyl, ethyl, cyclopropyl and allyl side chains for exploring P<sub>1</sub>'. The N-methylamide series (**24a-d**) were synthesized as outlined in Scheme S1 with a view to rapidly explore P<sub>1</sub>'. The N-alkylamide precursors, **22a-d** were prepared by coupling Boc-L-valine **19** to the appropriate N-methylamine hydrochloride **21a-d** via HATU-promoted amide coupling. Compounds **22a-d** were then subjected to Boc deprotection using TFA, followed by treatment with 10% KOH solution to liberate the free amines **23a-d**. Finally, **24a-d** were obtained by coupling **23a-d** to the pre-synthesized dipeptide **7a** using HBTU. Conformational isomerization about the amide bond in the N-methyl amide series **24a-d** was observed by NMR spectroscopy, thus purity of the final compounds was confirmed by HPLC (see SI).

The peptidic valinol derivative **28** was also synthesized as a replacement for the phosphonate moiety (Scheme S2). Benzyl protected L-valinol **26** was coupled to **7a** using HBTU to afford **27**, which was converted to the final compound **28** by hydrogenolysis using Pd catalysis. In CDCl<sub>3</sub>, compound **28** was observed as an isomeric mixture, with additional <sup>1</sup>H and <sup>13</sup>C NMR peaks evident in the spectra. However, HPLC analysis revealed only a single peak at t<sub>r</sub> = 32.5 min, indicative of a conformation isomer, and not diastereomers.

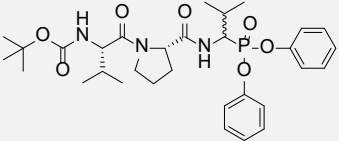
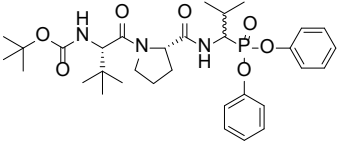
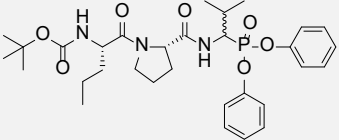
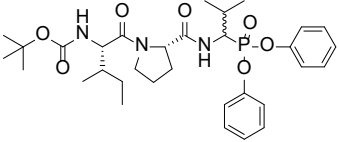
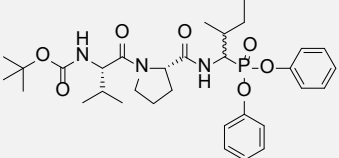
Compounds **29** and **30** with a boronic acid and  $\alpha$ -ketoamide transition state analogue respectively (both present in protease inhibitors used clinically) were purchased and tested to investigate their applicability in this system.<sup>28-29</sup>

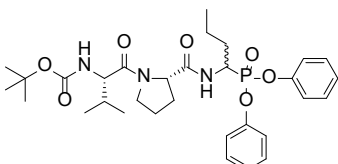
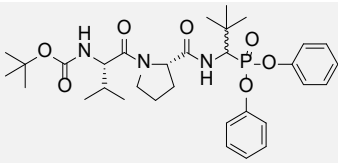
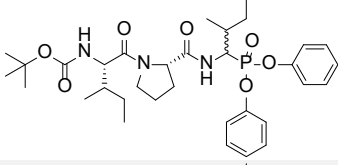
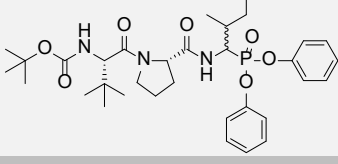
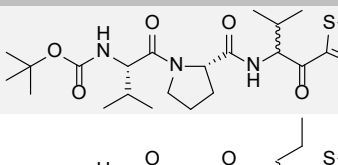
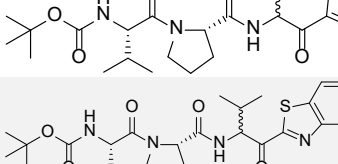
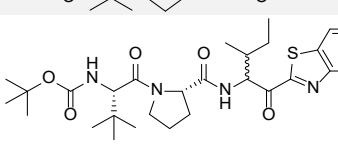

## 2.2 Enzyme inhibition and anti-chlamydial activity

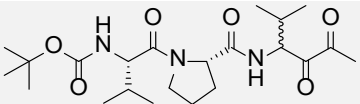
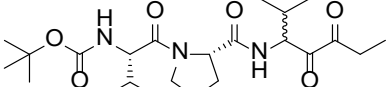
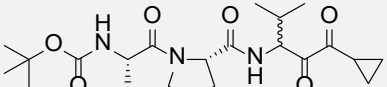
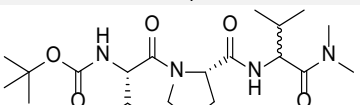
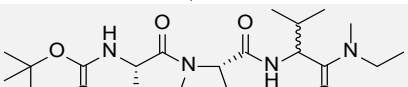
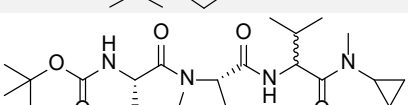
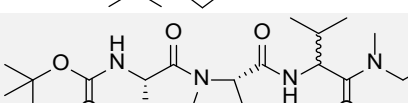
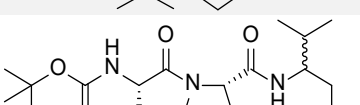
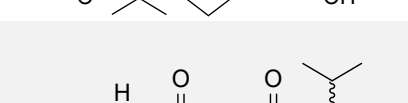
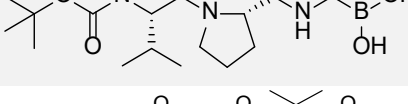
We set out to optimize the binding affinity of the inhibitor at the nonprime positions of JO146 by stepwise replacement of the P<sub>1</sub> and P<sub>3</sub> valine residues (Table 1) with norvaline (Nva), isoleucine and tertiary leucine (Tle). Compounds **8a-f**, **14a-c**, and **18a-c** were thus synthesized and tested for their biological activity. A bio-assessment of the compounds was conducted via protease inhibition assays initially against human neutrophil elastase (HNE) and CtHtrA along with off-target assessment carried out against the human proteases trypsin and chymotrypsin (Table 3). HNE possesses a similar substrate specificity to CtHtrA where substrates consist of mainly small hydrophobic residues such as valine and alanine, particularly at P<sub>1</sub>. Activity against HNE provides a good prediction for anti-CtHtrA activity and this was used as an indicator of CtHtrA activity.<sup>30-</sup>

<sup>31</sup> The *in vitro* susceptibility of *C. trachomatis* and koala *C. pecorum* isolates to all compounds was measured by their ability to significantly reduce the pathogens' infectious progeny, measured as inclusion forming units (IFU/mL; Table 3; Figures 2 and 3).

**Table 3 CtHtrA Inhibitory activity and selectivity assessment**

#	Compound structure	Site Change <sup>a</sup>	CtHtrA relative inhibition	Elastase IC <sub>50</sub> (μM)	Trypsin IC <sub>50</sub> (μM)	Chymotrypsin IC <sub>50</sub> (μM)	<i>In vitro</i> Ctrach activity <sup>b</sup> IFU/mL
<b>1</b> <b>(JO146)</b>			1	0.83 ± 0.01	> 500	> 500	4.34 × 10 <sup>2</sup> ± 9.0 × 10 <sup>-4</sup>
<b>Control</b>							DMSO 6.2 × 10 <sup>6</sup> ± 8.5 × 10 <sup>5</sup>
<b>Nonprime Site Optimization</b>							
<b>8a</b>		P <sub>3</sub>	33.33× more active	0.68 ± 0.02	> 500	> 500	1.5 × 10 <sup>3</sup> ± 216
<b>8b</b>		P <sub>3</sub>	9.34× less active	0.45 ± 0.09	> 500	> 500	1.7 × 10 <sup>6</sup> ± 1.5 × 10 <sup>5</sup>
<b>8c</b>		P <sub>3</sub>	14.29× more active	0.40 ± 0.03	> 500	> 500	5.0 × 10 <sup>4</sup> ± 7252
<b>8d</b>		P <sub>1</sub>	20× more active	0.36 ± 0.02	> 500	38.45 ± 0.51	9.2 × 10 <sup>3</sup> ± 2660

<b>8e</b>		P <sub>1</sub>	NA	0.68 ± 0.02	> 500	> 500	1.5 × 10 <sup>5</sup> ± 30297
<b>8f</b>		P <sub>1</sub>	NA	> 500	> 500	> 500	2.1 × 10 <sup>4</sup> ± 4220
<b>8g</b>		P <sub>1</sub> P <sub>3</sub>	13.09× less active	0.84 ± 0.05	> 500	1.28 ± 0.06	1.1 × 10 <sup>4</sup> ± 11565
<b>8h</b>		P <sub>1</sub> P <sub>3</sub>	NA	2.35 ± 0.22	> 500	0.53 ± 0.04	1.1 × 10 <sup>4</sup> ± 2947
Prime sites							
<b>14a</b>		TSA	NA	ND	ND	ND	3.31 × 10 <sup>6</sup> ± 3.9 × 10 <sup>5</sup>
<b>14b</b>		TSA	NA	> 150	> 500	> 500	ND
<b>14c</b>		TSA	1.06 × more active	2.60 ± 0.15	> 500	> 500	2.5 × 10 <sup>5</sup> ± 4.3 × 10 <sup>4</sup>
<b>14d</b>		TSA	NA	4.83 ± 0.08	> 500	> 500	2.8 × 10 <sup>5</sup> ± 4.6 × 10 <sup>4</sup>

<b>18a</b>		P <sub>1</sub> ,	NA	1.35 ± 0.17	> 500	> 500	8.6 × 10 <sup>6</sup> ± 1.1 × 10 <sup>5</sup>
<b>18b</b>		P <sub>1</sub> ,	NA	4.71 ± 0.66	> 500	> 500	5.7 × 10 <sup>6</sup> ± 5.2 × 10 <sup>5</sup>
<b>18c</b>		P <sub>1</sub> ,	NA	ND	> 500	> 500	5.5 × 10 <sup>6</sup> ± 1.5 × 10 <sup>5</sup>
<b>24a</b>		P <sub>1</sub> ,	NA	ND	ND	ND	ND
<b>24b</b>		P <sub>1</sub> ,	NA	ND	ND	ND	ND
<b>24c</b>		P <sub>1</sub> ,	NA	ND	ND	ND	ND
<b>24d</b>		P <sub>1</sub> ,	NA	ND	ND	ND	ND
<b>28</b>		TSA - valinol	NA	ND	ND	ND	ND
<b>29<sup>a</sup></b>		TSA - boronic acid	Activates enzyme before inhibiting <i>not</i> <i>able to fit</i> <i>curve</i>	ND	ND	ND	9.9 × 10 <sup>3</sup> ± 1.6 × 10 <sup>3</sup>
<b>30</b>		TSA - α- ketoamide	NA	ND	ND	ND	ND

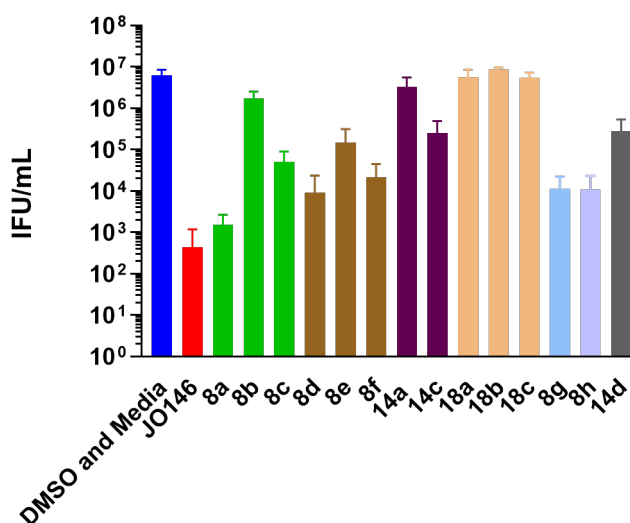


<sup>a</sup>Compounds differed from **1** by varying either the P<sub>3</sub>, P<sub>1</sub> residue, the transition state analogue or a combination thereof. <sup>b</sup>*C. trachomatis* susceptibility to inhibitors at a dose of 100 μM, 16 h PI. NA = not active in CtHtrA assay (see table S3 for details). ND = not determined, Data are presented as the mean ± standard error of the mean (SEM) of triplicate readings. TSA= transition state analogue.

### 2.3 P<sub>3</sub>-P<sub>1</sub> optimization

**P<sub>3</sub> Optimization:** Replacing valine with norvaline at P<sub>3</sub> (**8b**) showed better activity at HNE (IC<sub>50</sub> = 0.45 ± 0.09 μM; JO146 IC<sub>50</sub> = 0.83 ± 0.01 μM) but reduced both CtHtrA (9.34 × less active, Table 3) and cellular activity (*C. trachomatis*), with approximately 10<sup>3.5</sup> IFU reduction compared with JO146 (**1**) at 100 μM (Figure 2). Isoleucine at P<sub>3</sub> (**8c**) was more potent against HNE IC<sub>50</sub> = 0.40 ± 0.03 μM) and approximately fourteen times more active than JO146 (**1**) (CtHtrA relative inhibition; Table 3), but with less cellular activity (*C. trachomatis*) causing a reduction in IFU/mL of 10<sup>2</sup> at 100 μM (Figure 2). Compound **8a** with tertiary leucine at P<sub>3</sub> showed improved HNE activity (IC<sub>50</sub> = 0.68 ± 0.02 μM) over JO146 (**1**) and was the most active P<sub>3</sub>-optimized analogue yielding a 33-fold increase in relative CtHtrA inhibition compared to JO146 (**1**) (Table 3). The cellular potencies of JO146 (**1**) and **8a** were comparable (Figure 2).

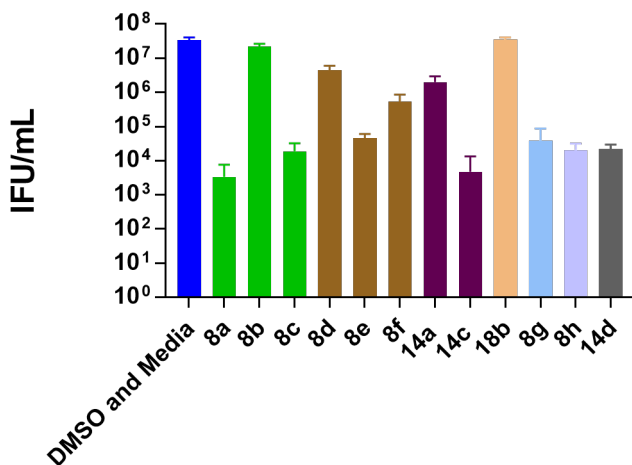
*C. pecorum* displayed greater susceptibility to P<sub>3</sub>-optimized compounds **8a** and **8c** compared to *C. trachomatis*. Against *C. pecorum*, **8a** remained the most active analogue with approximately 10<sup>4</sup> reduction in IFU counts at 100 μM relative to the DMSO control. Similarly, **8c** produced a substantial reduction in IFU counts, accounting for 10<sup>3</sup> loss of infectious progeny at 100 μM. However, compound **8b** with norvaline at P<sub>3</sub> showed only marginal activity against *C. pecorum* at 100 μM.



**Figure 2** *C. trachomatis* susceptibility to representative inhibitors (100  $\mu$ M, 16 h PI) of different covalent and noncovalent transition state analogues during mid-replicative phase of the pathogen's developmental cycle. Inclusion forming units counts (IFU/mL) were determined upon completion of developmental cycle. Green (P<sub>3</sub> variants); brown (P<sub>1</sub> variants); purple ( $\alpha$ -ketoheterocyclic variants); beige ( $\alpha$ -diketone variants); light blue and lilac (P<sub>3</sub> and P<sub>1</sub> variants); grey (P<sub>3</sub> and P<sub>1</sub>  $\alpha$ -ketoheterocycle). Error bars indicate the standard error of the mean obtained from experimental replicates (minimum of 5). \*Note in this assay 10<sup>3</sup> is the threshold for detection and can be equivalent to too few to accurately quantify.

**2.4 P<sub>1</sub> Optimization:** Of the compounds synthesized for P<sub>1</sub> optimization, the analogue with isoleucine at P<sub>1</sub> (**8d**) showed potent HNE activity (IC<sub>50</sub> = 0.36 ± 0.02  $\mu$ M) and produced the highest CtHtrA relative inhibition; exhibiting twenty-fold increase in potency compared to JO146 (**1**, Table 3). However, this activity was not extended to the cellular assays of the molecule at *C. trachomatis* (Figure 2), as it only produced approximately 10<sup>3</sup> loss in infectious progeny at 100  $\mu$ M, which was less potent than JO146 (**1**). In addition, it also showed chymotrypsin inhibition (IC<sub>50</sub> = 38.45 ± 0.51  $\mu$ M) which is not surprising due to the preference of chymotrypsin for larger hydrophobic

residues such as isoleucine at P<sub>1</sub>.<sup>32</sup> The replacement of valine at P<sub>1</sub> with norvaline (**8e**) and tertiary leucine (**8f**), on the other hand, was completely devoid of any CtHtrA inhibition activity and the latter also failed to inhibit HNE (Table 3). However, both **8e** and **8f** displayed promising cellular activity at 100 μM. Compound **8f** produced ~10<sup>2.5</sup> reduction in inclusion forming units (IFU) counts against *C. trachomatis* while **8e** yielded approximately 10<sup>1.5</sup> reduction in IFU counts relative to the untreated DMSO control (Figure 2). *C. pecorum* was generally less susceptible to compounds with P<sub>1</sub> modification, yielding appreciably lower reduction in IFU counts for all three compounds relative to their activity against *C. trachomatis*. Although **8e** produced a surprising 10<sup>3</sup> reduction in *C. pecorum* IFU counts at 100 μM (Figure 3), against the 10<sup>1.5</sup> reduction recorded at *C. trachomatis* (Figure 2). The lack of *in vitro* CtHtrA activity by **8e** and **8f** but with both molecules displaying considerable cellular activity, suggests they may act via a different mechanism of action.



**Figure 3:** Susceptibility of *C. pecorum* to CtHtrA inhibitors (100 μM; 16 h PI) during mid-replicative phase of the pathogen’s developmental cycle. Infectious yield counts (IFU/mL) were determined upon completion of developmental cycle. Colours the same as above in Figure 2. Error

bars indicate the standard error of the mean obtained from experimental replicates (minimum of 5).

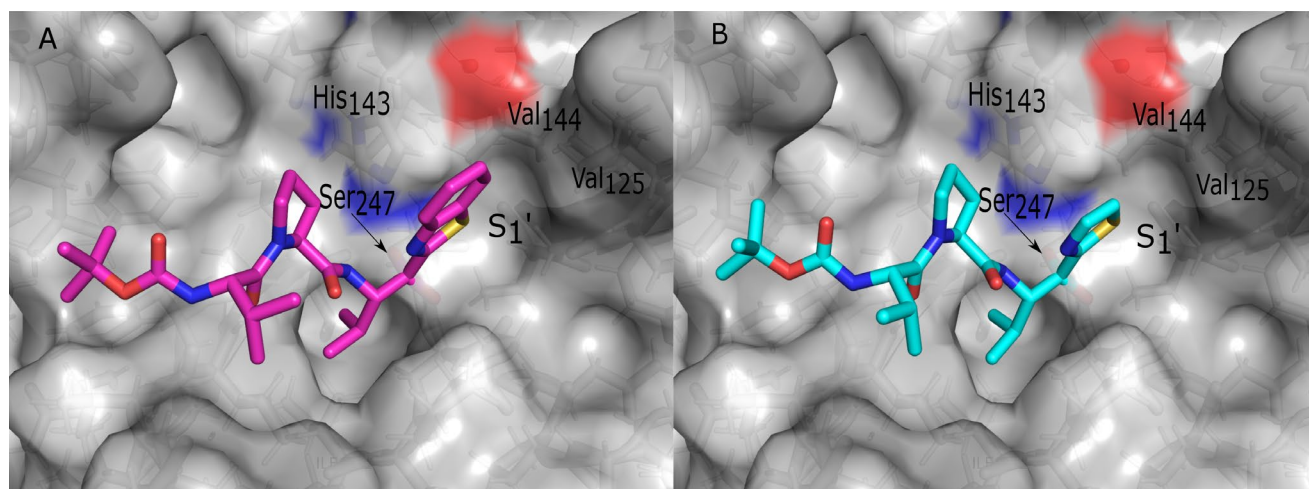
## 2.5 Replacement of Diphenylphosphonate

The effects of substituting the phosphonate moiety in JO146 (**1**) with reversible analogues were evaluated by testing compounds **14a-e**, **18a-c**, **24a-d**, and **28-30** for *in vitro* CtHtrA and HNE inhibition, and trypsin and chymotrypsin to assess off-target selectivity (Table 3).

Among the reversible derivatives of JO146 (**1**), only the benzothiazole  $\alpha$ -ketoheterocycle transition state analogue (**14c**) showed viable anti-chlamydial activity when simply replacing the diphenyl phosphonate group. The compound displayed CtHtrA relative inhibition comparable or slightly better than JO146 (**1**,  $1.06 \times$  more active; Table 3) It was less active at the closely related HNE ( $IC_{50} = 2.60 \pm 0.15 \mu\text{M}$ ; JO146 ( $IC_{50} = 0.83 \pm 0.01$ )) implying improved selectivity over HNE which is desirable. The inhibitor produced some cellular activity against *C. trachomatis*. Against *C. pecorum* it produced a considerably greater reduction with approximately 4 log loss of infectious bodies at 100  $\mu\text{M}$  (Figure 3). The lack of anti-chlamydial activity displayed by the thiazole (**14a**) and N-methyl imidazole (**14e**; data not shown) derivatives, is traceable to their inability to sufficiently activate the ketone carbonyl carbon towards a nucleophilic attack by the catalytic Ser<sub>247</sub> of CtHtrA. These results align with previous studies by Edwards *et al.*<sup>33</sup> which identified peptidyl ketoheterocycle HNE inhibitors with imidazoles as incapable of forming a covalent adduct and supports the necessity of a covalent interaction between the inhibitor and the active site of CtHtrA for anti-chlamydial activity.

To understand the structural basis for the activity exhibited by the **14c** benzothiazole  $\alpha$ -ketoheterocycle compared to **14a**, both molecules were docked into a homology model of CtHtrA.

The docking results (Figure 4) revealed more interaction between the benzothiazole moiety of **14c** and the S<sub>1</sub>' pocket of CtHtrA, with the heterocycle engaging in van der Waals interactions with Val<sub>125</sub> and Val<sub>144</sub> that line the walls of the pocket, as well as a potential edge-to-face pi-stacking interaction with His<sub>143</sub>. Compound **14a** lacks the benzene extension and is therefore unable to make these interactions. Any possible hydrogen bonding between the heterocyclic (thiazole) nitrogen and the catalytic His<sub>143</sub> of CtHtrA was revealed to be unfavourable, as the orientation of the imidazole ring of His<sub>143</sub> relative to the inhibitor thiazole ring makes it unlikely for hydrogen bonding to occur. This may also contribute to the difference in the anti-chlamydial activity between compounds **14a** and **14c**.



**Figure 4.** Binding poses of **14c**, magenta (A) and **14a**, cyan (B) to CtHtrA (transparent surface). Oxygen is shown in red, nitrogen blue and sulphur in yellow.

**2.6 P<sub>1</sub>' Optimization:** In addition to replacing the phosphonate group, expansion and optimization of P<sub>1</sub>' could be probed using  $\alpha$ -diketone based inhibitors. Three inhibitors were synthesized (**18a-c**) containing a methyl, ethyl or cyclopropyl group at P<sub>1</sub>' and showed no CtHtrA inhibition or cellular activity (Table 3; Figures 2 and 3). This result is surprising given both **18a** and **18b** showed inhibition against HNE albeit modest ( $IC_{50} = 1.35 \pm 0.17$  and  $IC_{50} = 4.71 \pm 0.66 \mu\text{M}$  respectively) and the  $\alpha$ -diketone is present in some clinical HCV protease inhibitors.

## 2.7 Combined optimization

Based on the *in vitro* HNE and CtHtrA relative inhibition results above, where isoleucine and tertiary leucine were identified as optimal residues at P<sub>1</sub> and P<sub>3</sub> respectively, compounds retaining the diphenyl phosphonate combining P<sub>3</sub> and P<sub>1</sub> optimisation **8g**, **8h** along with an  $\alpha$ -keto heterocycle transition state analogue **14d** were synthesized. Compound **8h** produced approximately 10<sup>2.5</sup> reduction in IFU counts at 100  $\mu$ M (*C. trachomatis*) and a 10<sup>3</sup> reduction in IFU counts against *C. pecorum* at 100  $\mu$ M (Figure 3).

The benzothiazole analogue **14d** failed to show *in vitro* CtHtrA inhibition with only moderate HNE inhibition, but showed some cellular activity against *C. trachomatis* (Figure 2), and a slightly improved activity at *C. pecorum* (10<sup>3</sup> reduction in IFU 100  $\mu$ M; Figure 3). However, in contrast to the phosphonate compounds **8g** and **8h** which exhibited high chymotrypsin inhibition with IC<sub>50</sub> values of 1.28  $\pm$  0.06  $\mu$ M and 0.53  $\pm$  0.04  $\mu$ M respectively (Table 2); **14d** showed no chymotrypsin inhibition (IC<sub>50</sub> > 500  $\mu$ M) despite differing only at the transition state analogue (Table 3).

As a measure of off-target effect, no inhibition of trypsin and chymotrypsin (IC<sub>50</sub> > 500  $\mu$ M) was observed for most of the active compounds except **8d**, **8g** and **8h** which have isoleucine at P<sub>1</sub> and the latter compound showing appreciable activity such to exclude it from further study. Compound **8g** showed higher potency compared to **8h** against HNE (IC<sub>50</sub> = 0.84  $\pm$  0.05  $\mu$ M; IC<sub>50</sub> = 2.35  $\pm$  0.22  $\mu$ M) but only at a similar level to JO146.

The dicarbonyl-containing compounds **18a-c** and **30** ( $\alpha$ -diketone and  $\alpha$ -ketoamide respectively) exhibited an unexpected lack of biological activity. Both molecules lacked *in vitro* CtHtrA inhibition. This was extended to negligible cellular performance of the peptidyl  $\alpha$ -ketoamide **30** against *C. trachomatis*.

Unusual activity was observed for the boronic acid inhibitor **29**. The compound activated CtHtrA at doses less than 50  $\mu\text{M}$ , but displayed enzyme inhibition at higher doses (Figure S1). At 100  $\mu\text{M}$ , it produced a 3 log reduction in IFU count against *C. trachomatis* (data not shown). Whilst the precise mechanism for the initial CtHtrA activation is unclear, the presence of the boronic acid moiety mimics a carboxylate terminus of a peptide substrate, which would bind to the PDZ1 domain of CtHtrA and thus activating it towards proteolysis.<sup>17</sup>

Generally, the noncovalent binders (peptidyl N-methylamides **24a-d**, and the peptidyl valinol **28**) lacked both biochemical ( $\text{IC}_{50} > 500\mu\text{M}$ ) and cellular activity, indicating a covalent interaction between the inhibitor and the active site of CtHtrA is crucial for optimal activity.

### 3. Conclusion

We have found that inhibitors are required to interact with CtHtrA via a covalent interaction, either through reversible or irreversible transition state analogues. This is evident in both *in vitro* and *in vivo* data. The most potent compound identified was **8a** (CtHtrA relative inhibition) which has retained the diphenylphosphonate moiety with the related compound **8d** showing the highest activity against HNE. The identification of the  $\alpha$ -ketobenzothiazole moiety as a selective reversible transition state analogue with *in vitro* activity against both human and koala pathogenic *Chlamydia* species, opens up another possible avenue to obtain more “drug-like” inhibitors of CtHtrA. This moves away from the previously identified irreversible diphenylphosphonate inhibitors. It also indicates that P<sub>1</sub>' optimization via the use of  $\alpha$ -ketoheterocycles is a potential strategy for improved potency and selectivity.

Optimization of binding affinity at CtHtrA subpockets via modification to inhibitor residues P<sub>1</sub> and P<sub>3</sub> produced considerable improvement in potency and/or selectivity. Selectivity was obtained



over, trypsin and chymotrypsin and likely HNE, in addition to retaining cellular activity comparable with JO146 (**1**), when tertiary leucine was tested as the P<sub>3</sub> residue (**8a**). This is also a viable means to obtain molecules with favourable pharmacokinetic properties, given that tertiary leucine is a non-natural amino acid, and may avoid recognition by degradative enzymes, thus potentially enhancing inhibitor performance *in vivo*. After testing numerous derivatives, valine remains the best residue at P<sub>1</sub> for *in vivo* activity. Outside of selectivity, there seems no explanation as to why the  $\alpha$ -diketoamide transition state analogues and the related  $\alpha$ -diketone derivatives lacked activity given the use of the former in clinical protease inhibitors. It may be related to the more polar character as compared to the diphenyl phosphonates.

CtHtrA is a complicated enzyme, which makes simple comparisons difficult. The enzyme is known to exist in at least 3 oligomeric states, an inactive hexamer and, following activation of the PDZ domain by a peptide terminal carboxylate, active 12-mers and 24-mers.<sup>16-17, 34</sup> In addition, the covalent nature of the inhibitors complicates the use of explicit IC<sub>50</sub> values hence the use of relative values.<sup>35</sup> The allosteric activation of multiple monomers in a hexamer to then accommodate inhibitor binding adds further complexity. This is illustrated with the behaviour of the boronic acid derivative (**29**) which acted as both an activator and inhibitor (Figure S1).

JO146 remains the most active inhibitor in terms of cellular performance, however potent analogues with malleable structural features and promising pharmacokinetic prospects have been identified. Further optimisation at P<sub>2</sub> (proline) and P<sub>4</sub> (Boc) to increase potency and reduce peptidic character is the next obvious step in moving towards clinically relevant protease inhibitors for the treatment of human and animal chlamydiosis.

## 4. Experimental

### 4.1. Compound Synthesis

**4.1.1. General Methods.** All solvents and reagents were commercially procured (Sigma-Aldrich, Merck, AK Scientific, Thermo Scientific, Acros Organics, BDH, and Cambridge Isotope) and used without further purification. All reactions were conducted in standard glassware and in air or under nitrogen gas unless otherwise stated. Organic solvent extracts were dried with  $\text{MgSO}_4$  and subjected to rotary evaporation, and finally dried at  $10^{-1}$  mbar using a high vacuum pump.

Silica gel 60 (0.040-0.063 mm, 200-400 mesh) was used for flash column chromatography with all solvent systems expressed as volume to volume (v/v) ratios. Solids were recrystallized from a minimum amount of hot solvent and collected by vacuum filtration. Analytical thin layer chromatography (TLC) was performed on Merck TLC aluminium plates coated with 0.2 mm silica gel 60 F<sub>254</sub>. Spots were generally detected by UV (254 nm) and/or permanganate staining

$^1\text{H}$  and  $^{13}\text{C}$  NMR spectra were recorded at room temperature on Varian 400 or 500 MHz spectrometers. Samples were prepared in deuterated solvents, chloroform ( $\delta$  7.26, 77.16 ppm) or acetonitrile ( $\delta$  1.95, 118.69 and 1.72 ppm) with the respective  $^1\text{H}$  and  $^{13}\text{C}$  chemical shifts shown in brackets. Chemical shifts ( $\delta$ ) are expressed in parts per million (ppm) and coupling constants ( $J$ ) in Hertz (Hz), both measured against the residual solvent peak or internal standard tetramethylsilane (TMS). High resolution mass spectrometry were recorded on a Bruker microTOF-Q spectrometer with an electrospray ionization (ESI) source.

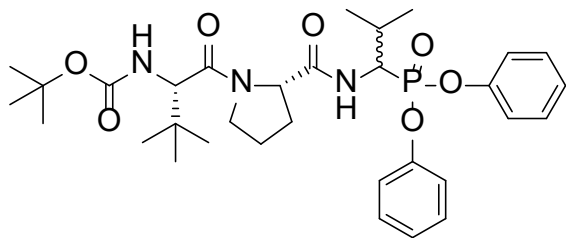
The purity of compounds was determined by reverse-phase high performance liquid chromatography (HPLC) carried out on an Agilent HPLC with a Gemini 5  $\mu\text{m}$  C18 110 Å column (250  $\times$  4.6 mm, Phenomenex, New Zealand). Compounds were eluted using solvent A: 0.1% trifluoroacetic acid (TFA) in water and solvent B: 0.1% TFA in acetonitrile (ACN) over a linear

gradient (40% ACN to 85% ACN over 45 min). Compounds were detected at 254/210 nm with a flow rate of 1.0 mL/min. The purity of compound **28** was determined on a Shimadzu HPLC with a Gemini 5  $\mu\text{m}$  C18 110 Å column (250  $\times$  4.6 mm, Phenomenex, New Zealand). The compound was eluted using solvent A: 0.1% formic acid in water and solvent B: 0.1% formic acid in ACN in a linear gradient from 30 to 60% ACN over 60 min. The flow rate was 1.2 mL/min and compounds were detected at the wavelength of 224 nm. Final compounds including those obtained as isomeric mixtures, are >95% pure unless otherwise stated, see supporting information for HPLC traces of the compounds.

#### 4.1.2. General Procedure for Synthesis of peptidyl phosphonates (**1**, **8a-h**)<sup>20</sup>

Compounds **7a-d** (1 equiv.) were dissolved in 3 mL DMF. To this *O*-(Benzotriazol-1-yl)-*N,N,N',N'*-tetramethyluronium hexafluoro-phosphate (HBTU) (1.2 equiv.) and *N,N*-Diisopropylethylamine (DIEA) (2.5 equiv.) were added to generate the activated esters. After 10 min, compounds **6a-d** (1.5 equiv.) were added and the reaction mixtures were stirred under nitrogen overnight. The mixtures were diluted with EtOAc (25 mL) and an equal amount of sodium bicarbonate (NaHCO<sub>3</sub>), washed with brine (6  $\times$  25 mL), dried (MgSO<sub>4</sub>) and concentrated *in vacuo*. The crude products were purified by column chromatography (100% EtOAc). This yielded **1**, **8a-h**, as diastereomeric mixtures with characteristic <sup>13</sup>C-<sup>31</sup>P coupling. We have previously reported the full NMR and mass spectrometric characterization of JO146 **1**, as part of the stereochemical investigation of its two diastereomers.<sup>36</sup> The analogues **8a-h** are characterized as follows with <sup>13</sup>C-<sup>31</sup>P *J* values reported where clearly evident.

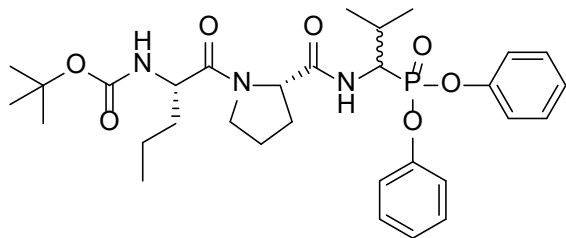
***Tert*-butyl-*N*-[(2*S*)-1-[(2*S*)-2-[[1-(diphenoxyphosphoryl)-2-methylpropyl]carbamoyl]pyrrolidin-1-yl]-3,3-dimethyl-1-oxobutan-2-yl]carbamate (**8a**)**



Using general procedure above, 0.17 g (0.52 mmol) of **7b** was coupled to 0.30 g (0.78 mmol) of **6a** to yield a brown oil, which formed a brown solid foam under reduced pressure (0.08 g, 25%). ~94% purity

by HPLC;  $t_r = 38.234$  and  $38.651$  min.  $^1\text{H}$  NMR (400 MHz,  $\text{CDCl}_3$ )  $\delta$  7.37 – 7.24 (m, 5H), 7.20 – 7.11 (m, 5H), 5.44 (d,  $J = 9.9$  Hz, 0.5H), 5.25 (d,  $J = 9.9$  Hz, 0.5H), 4.80 – 4.70 (m, 1H), 4.55 – 4.48 (m, 1H), 4.35 – 4.26 (m, 1H), 3.83 – 3.51 (m, 2H), 2.48 – 2.33 (m, 2H), 2.10 – 1.81 (m, 3H), 1.44 – 1.40 (m, 9H), 1.15 – 1.04 (m, 6H), 1.00 – 0.91 (m, 9H).  $^{13}\text{C}$  NMR (101 MHz,  $\text{CDCl}_3$ )  $\delta$  172.9, 171.9, 171.7 (d,  $J_{\text{C-P}} = 14.0$  Hz), 171.4 (d,  $J_{\text{C-P}} = 5.8$  Hz), 155.72, 155.65, 150.4 (d,  $J_{\text{C-P}} = 10.0$  Hz), 150.2, 150.1, 150.0, 130.9, 129.8 (d,  $J_{\text{C-P}} = 1.0$  Hz), 129.71 (d,  $J_{\text{C-P}} = 1.0$  Hz), 129.65 (d,  $J_{\text{C-P}} = 0.9$  Hz), 129.60 (d,  $J_{\text{C-P}} = 0.9$  Hz), 125.24, 125.14 (d,  $J_{\text{C-P}} = 1.1$  Hz), 125.07 (d,  $J_{\text{C-P}} = 0.8$  Hz), 120.8 (d,  $J_{\text{C-P}} = 4.0$  Hz), 120.64 (d,  $J_{\text{C-P}} = 4.4$  Hz), 120.63 (d,  $J_{\text{C-P}} = 4.0$  Hz), 120.4 (d,  $J_{\text{C-P}} = 4.3$  Hz), 79.7, 79.5, 60.4, 60.1, 58.3, 51.2 (d,  $J_{\text{C-P}} = 154.8$  Hz), 51.0 (d,  $J_{\text{C-P}} = 155.6$  Hz), 48.61, 48.58, 35.7, 35.5, 35.2, 29.7, 29.33 (d,  $J_{\text{C-P}} = 3.9$  Hz), 29.25 (d,  $J_{\text{C-P}} = 3.9$  Hz), 28.37, 28.35, 28.3, 27.9, 27.6, 26.4, 26.3, 26.2, 25.4, 25.0, 20.57 (d,  $J_{\text{C-P}} = 13.9$  Hz), 20.28 (d,  $J_{\text{C-P}} = 13.8$  Hz), 18.0 (d,  $J_{\text{C-P}} = 4.7$  Hz), 17.9 (d,  $J_{\text{C-P}} = 4.4$  Hz). ESI-MS ( $m/z$ ): calcd for  $\text{C}_{32}\text{H}_{46}\text{N}_3\text{NaO}_7\text{P}$  (M + Na)  $m/z$  638.2971 found 638.2918.

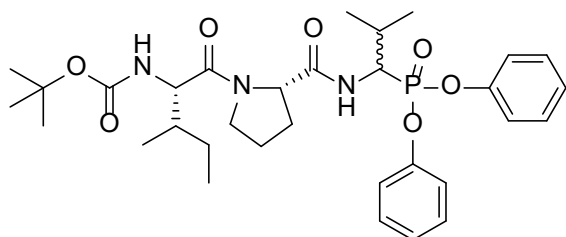
***Tert*-butyl-*N*-[(2*S*)-1-[(2*S*)-2-[[1-(diphenoxyphosphoryl)-2-methylpropyl]carbamoyl]pyrrolidin-1-yl]-1-oxopentan-2-yl]carbamate (**8b**)**



Using general procedure above, 0.08 g (0.25 mmol) of **7c** was coupled to 0.15 g (0.38 mmol) of **6a** to yield a brown oil which formed a brown solid foam under reduced pressure (0.10 g, 67%). HPLC  $t_r$  =

33.679 and 34.581 min.  $^1\text{H}$  NMR (400 MHz,  $\text{CDCl}_3$ )  $\delta$  7.56 – 7.44 (m, 1H), 7.34 – 7.09 (m, 10H), 5.27 (d,  $J = 9.3$  Hz, 0.5H), 5.12 (d,  $J = 9.0$  Hz, 0.5H), 4.78 – 4.65 (m, 2H), 4.56 – 4.32 (m, 1H), 3.74 – 3.52 (m, 2H), 3.45 – 3.36 (m, 1H), 2.47 – 2.27 (m, 2H), 2.20 – 2.08 (m, 1H), 2.02 – 1.75 (m, 3H), 1.64 – 1.50 (m, 2H), 1.46 – 1.39 (m, 9H), 1.13 – 0.97 (m, 6H), 0.93 – 0.65 (m, 3H).  $^{13}\text{C}$  NMR (101 MHz,  $\text{CDCl}_3$ )  $\delta$  173.9, 173.1, 171.0, 170.9 (d,  $J_{\text{C-P}} = 7.1$  Hz), 155.6, 155.5, 150.42 (d,  $J_{\text{C-P}} = 9.1$  Hz), 150.3, 150.2, 150.0 (d,  $J_{\text{C-P}} = 9.1$  Hz), 129.83 (d,  $J_{\text{C-P}} = 1.1$  Hz), 129.76 (d,  $J_{\text{C-P}} = 1.1$  Hz), 129.73 (d,  $J_{\text{C-P}} = 1.1$  Hz), 129.67 (d,  $J_{\text{C-P}} = 0.8$  Hz), 129.63, 125.32 (d,  $J_{\text{C-P}} = 2.0$  Hz), 125.26 (d,  $J_{\text{C-P}} = 1.3$  Hz), 125.10, 120.73 (d,  $J_{\text{C-P}} = 4.1$  Hz), 120.72 (d,  $J_{\text{C-P}} = 4.1$  Hz), 120.4 (d,  $J_{\text{C-P}} = 4.4$  Hz), 79.66, 59.90, 59.69, 51.1 (d,  $J_{\text{C-P}} = 154.6$  Hz), 50.9 (d,  $J_{\text{C-P}} = 130.6$  Hz), 47.4, 47.3, 35.6, 35.1, 29.2 (d,  $J_{\text{C-P}} = 4.1$  Hz), 28.34, 28.33, 26.83, 26.81, 25.2, 25.0, 20.5 (d,  $J_{\text{C-P}} = 13.8$  Hz), 20.3 (d,  $J_{\text{C-P}} = 14.1$  Hz), 18.77, 18.75, 17.9 (d,  $J_{\text{C-P}} = 4.6$  Hz), 17.8 (d,  $J_{\text{C-P}} = 4.3$  Hz), 13.7, 13.4. ESI-MS ( $m/z$ ): calcd for  $\text{C}_{31}\text{H}_{44}\text{N}_3\text{NaO}_7\text{P}$  ( $\text{M} + \text{Na}$ )  $m/z$  624.2815 found 624.2784.

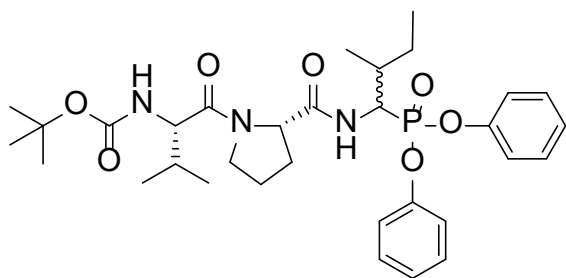
***Tert*-butyl-*N*-[(2*S*)-1-[(2*S*)-2-[[1-(diphenoxyphosphoryl)-2-methylpropyl]carbamoyl]pyrrolidin-1-yl]-3-methyl-1-oxopentan-2-yl]carbamate (**8c**)**



Using general procedure above, 0.25 g (0.76 mmol) of **7d** was coupled to 0.35 g (1.1 mmol) of **6a** to yield a brown oil which formed a brown solid foam under reduced pressure (0.23 g, 49%). HPLC  $t_r$  = 28

37.496 and 37.939 min.  $^1\text{H}$  NMR (400 MHz,  $\text{CDCl}_3$ )  $\delta$  7.40 (d,  $J = 10.4$  Hz, 0.5H), 7.27 – 7.17 (m, 5H), 7.17 – 7.08 (m, 5.5H), 5.27 (d,  $J = 9.5$  Hz, 0.5H), 5.11 (d,  $J = 9.5$  Hz, 0.5H), 4.84 – 4.64 (m, 2H), 4.51 – 4.43 (m, 1H), 4.31 – 4.13 (m, 1H), 3.78 – 3.50 (m, 2H), 2.42 – 2.04 (m, 2H), 2.00 – 1.63 (m, 3H), 1.60 – 1.47 (m, 2H), 1.43 – 1.32 (m, 9H), 1.11 – 0.97 (m, 6H), 0.92 (m, 2H), 0.85 – 0.76 (m, 3H), 0.71 – 0.65 (m, 1H).  $^{13}\text{C}$  NMR (101 MHz,  $\text{CDCl}_3$ )  $\delta$  172.6, 171.3, 171.2, 155.7, 155.6, 150.4, 150.3, 150.2, 150.1 (d,  $J_{\text{C-P}} = 9.0$  Hz), 150.0, 129.7, 129.62, 129.57, 125.25 (d,  $J_{\text{C-P}} = 1.0$  Hz), 125.20 (d,  $J_{\text{C-P}} = 1.2$  Hz), 125.07 (d,  $J_{\text{C-P}} = 0.8$  Hz), 125.04 (d,  $J_{\text{C-P}} = 1.1$  Hz), 120.7 (d,  $J_{\text{C-P}} = 4.1$  Hz), 120.6 (d,  $J_{\text{C-P}} = 4.0$  Hz), 120.42, 120.37, 120.3, 79.6, 79.5, 60.3, 60.0, 56.2, 56.1, 51.3 (d,  $J_{\text{C-P}} = 154.6$  Hz), 51.0 (d,  $J_{\text{C-P}} = 154.6$  Hz), 48.0, 47.8, 37.83, 37.75, 29.3 (d,  $J_{\text{C-P}} = 3.6$  Hz), 29.2 (d,  $J_{\text{C-P}} = 3.8$  Hz), 28.30, 28.29, 27.4, 25.2, 25.0, 24.3, 24.0, 20.5 (d,  $J_{\text{C-P}} = 13.3$  Hz), 20.2 (d,  $J_{\text{C-P}} = 13.7$  Hz), 18.0 (d,  $J_{\text{C-P}} = 5.1$  Hz), 17.9 (d,  $J_{\text{C-P}} = 4.6$  Hz), 15.6, 15.2, 11.1, 10.9. ESI-MS ( $m/z$ ): calcd for  $\text{C}_{32}\text{H}_{46}\text{N}_3\text{NaO}_7\text{P}$  ( $\text{M} + \text{Na}$ )  $m/z$  638.2971 found 638.2911.

***Tert*-butyl-N-[(2*S*)-1-[(2*S*)-2-[[1-(diphenoxyphosphoryl)-2-methylbutyl]carbamoyl]pyrrolidin-1-yl]-3-methyl-1-oxobutan-2-yl]carbamate (**8d**)**

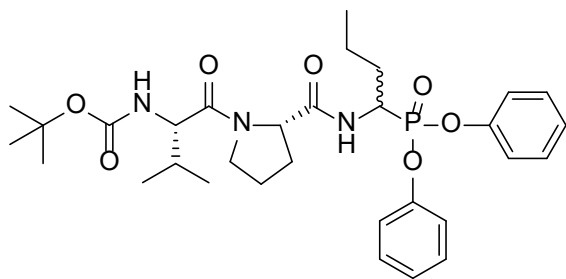


Using general procedure above, 0.10 g (0.32 mmol) of **7a** was coupled to 0.15 g (0.48 mmol) of **6d** to yield a brown oil which formed a brown solid foam under reduced pressure (0.06 g, 31%). HPLC  $t_r =$

37.845, 38.674 and 39.275 min.  $^1\text{H}$  NMR (400 MHz, Chloroform-*d*)  $\delta$  7.46 – 7.35 (m, 1H), 7.32 – 7.10 (m, 10H), 5.36 (d,  $J = 9.6$  Hz, 0.5H), 5.20 (d,  $J = 9.4$  Hz, 0.5H), 4.96 – 4.68 (m, 2H), 4.52 – 4.44 (m, 1H), 4.32 – 4.21 (m, 1H), 3.79 – 3.41 (m, 2H), 2.47 – 2.37 (m, 0.5H), 2.20 – 2.03 (m, 2.5H), 2.02 – 1.74 (m, 4H), 1.49 – 1.36 (m, 9H), 1.16 – 1.02 (m, 3H), 0.99 – 0.78 (m, 9H).  $^{13}\text{C}$  NMR (101 MHz,  $\text{CDCl}_3$ )  $\delta$  173.43, 173.37, 172.44, 172.39, 171.3 (d,  $J_{\text{C-P}} = 6.5$  Hz), 171.22 (d,  $J_{\text{C-P}} = 6.5$  Hz), 171.19 (d,  $J_{\text{C-P}} = 5.7$  Hz), 171.11, 171.06, 155.83, 155.80, 150.49, 150.47, 150.39,

150.37, 150.27, 150.17, 150.09, 150.07, 150.00, 129.73, 129.67, 129.57 (d,  $J_{C-P} = 0.8\text{Hz}$ ), 129.55 (d,  $J_{C-P} = 0.8\text{ Hz}$ ), 125.30 (d,  $J_{C-P} = 1.3\text{ Hz}$ ), 125.27 (d,  $J_{C-P} = 1.5\text{ Hz}$ ), 125.10, 125.09, 125.07, 120.74, 120.70, 120.67, 120.65, 120.64, 120.60, 120.47 (d,  $J_{C-P} = 1.2\text{ Hz}$ ), 120.42 (d,  $J_{C-P} = 1.2\text{ Hz}$ ), 120.38 (d,  $J_{C-P} = 4.3\text{Hz}$ ), 120.36 (d,  $J_{C-P} = 4.3\text{ Hz}$ ), 79.67, 79.59, 60.27, 60.25, 59.9, 56.8, 51.11 (d,  $J_{C-P} = 154.1\text{ Hz}$ ), 51.08 (d,  $J_{C-P} = 155.1\text{ Hz}$ ), 49.93, 49.90, 48.39, 48.35, 47.70, 47.68, 36.2 (d,  $J_{C-P} = 4.0\text{ Hz}$ ), 35.5 (d,  $J_{C-P} = 3.8\text{ Hz}$ ), 31.4, 31.2, 29.7, 28.34, 28.31, 28.22, 27.60, 27.48, 27.40, 27.34, 27.31, 27.25, 27.16, 26.98, 26.82, 25.24, 25.23, 24.95, 24.80, 24.75, 24.70, 19.6, 19.5, 17.4, 17.30, 17.28, 16.5 (d,  $J_{C-P} = 12.2\text{ Hz}$ ), 16.1 (d,  $J_{C-P} = 11.8\text{ Hz}$ ), 15.14 (d,  $J_{C-P} = 3.0\text{ Hz}$ ), 15.06 (d,  $J_{C-P} = 2.8\text{ Hz}$ ), 11.72, 11.71, 11.6, 11.4. ESI-MS ( $m/z$ ): calcd for  $\text{C}_{32}\text{H}_{46}\text{N}_3\text{NaO}_7\text{P}$  (M + Na)  $m/z$  638.2971 found 638.2940.

***Tert*-butyl-*N*-[(2*S*)-1-[(2*S*)-2-{{1-(diphenoxyphosphoryl)butyl}carbamoyl}pyrrolidin-1-yl]-3-methyl-1-oxobutan-2-yl]carbamate (8e)**

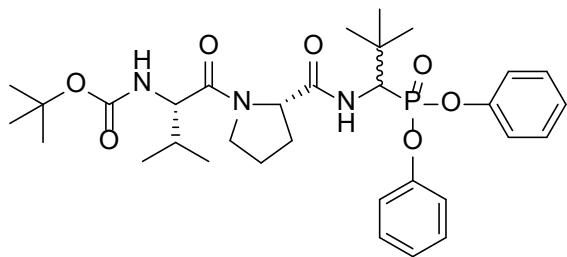


Using general procedure above, 0.13 g (0.41 mmol) of **7a** was coupled to 0.19 g (0.62 mmol) of **6c** to yield a colorless oil which formed a white solid foam under reduced pressure (0.12 g, 48%).

HPLC  $t_r = 33.974$  and  $34.523$  min.  $^1\text{H}$  NMR (400 MHz,  $\text{CDCl}_3$ )  $\delta$  7.84 – 7.79 (m, 1H), 7.43 – 7.04 (m, 10H), 5.28 (d,  $J = 9.6\text{ Hz}$ , 0.5H), 5.20 (d,  $J = 9.3\text{ Hz}$ , 0.5H), 4.85 – 4.70 (m, 1H), 4.53 – 4.44 (m, 1H), 4.25 (dd,  $J = 9.3, 6.1\text{ Hz}$ , 1H), 3.76 – 3.47 (m, 2H), 2.42 – 2.31 (m, 0.5H), 2.25 – 1.61 (m, 4.5H), 1.62 – 1.49 (m, 2H), 1.48 – 1.37 (m, 9H), 1.33 – 1.16 (m, 2H), 0.94 – 0.79 (m, 9H).  $^{13}\text{C}$  NMR (101 MHz,  $\text{CDCl}_3$ )  $\delta$  173.2, 171.03, 170.97, 155.8, 150.4, 150.3, 150.0 (d,  $J_{C-P} = 9.3\text{ Hz}$ ) 130.9, 129.8, 129.74, 129.65 (d,  $J_{C-P} = 0.8\text{ Hz}$ ), 126.7, 125.3 (d,  $J_{C-P} = 1.4\text{ Hz}$ ) 125.19 (d,  $J_{C-P} = 1.3\text{ Hz}$ ), 120.62 (d,  $J_{C-P} = 4.0\text{ Hz}$ ), 120.40 (d,  $J_{C-P} = 4.4\text{ Hz}$ ), 115.4, 114.7, 113.5, 79.7, 60.3, 60.0, 56.9, 56.8, 47.72, 46.71, 45.1, 31.89, 31.86, 31.4, 31.3, 29.7, 29.4, 28.34, 28.32, 27.5, 27.2,

25.2, 25.0, 22.7, 19.5, 19.4, 19.1, 19.0, 17.3, 14.1, 13.53 (d,  $J_{C-P} = 1.0$  Hz), 13.45. ESI-MS ( $m/z$ ): calcd for  $C_{31}H_{44}N_3NaO_7P$  ( $M + Na$ )  $m/z$  624.2815 found 624.2843.

***Tert*-butyl-N-[(2*S*)-1-[(2*S*)-2-[[1-(diphenoxyphosphoryl)-2,2-dimethylpropyl]carbamoyl]pyrrolidin-1-yl]-3-methyl-1-oxobutan-2-yl]carbamate (**8f**)**

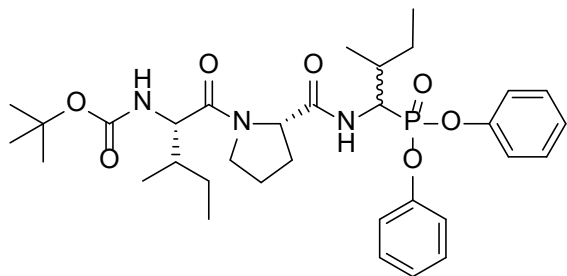


Using general procedure above, 0.10 g (0.32 mmol) of **7a** was coupled to 0.15 g (0.48 mmol) of **6b** to yield a colorless oil which formed a white solid foam under reduced pressure (0.15 g, 77%). HPLC

$t_r = 37.308$  and  $38.316$  min.  $^1H$  NMR (400 MHz,  $CDCl_3$ )  $\delta$  7.53 – 7.49 (m, 0.5H), 7.36 – 7.32 (m, 0.5H), 7.32 – 7.20 (m, 5H), 7.20 – 7.09 (m, 5H), 5.33 (d,  $J = 9.3$  Hz, 0.5H), 5.22 (d,  $J = 9.5$  Hz, 0.5H), 4.80 – 4.74 (m, 1H), 4.68 – 4.58 (m, 1H), 4.48 – 4.43 (m, 1H), 4.32 – 4.21 (m, 1H), 3.72 – 3.48 (m, 2H), 2.19 – 1.92 (m, 4H), 1.85 – 1.75 (m, 1H), 1.46 – 1.36 (m, 9H), 1.24 – 1.09 (m, 9H), 0.98 – 0.90 (m, 3H), 0.88 – 0.78 (m, 3H).  $^{13}C$  NMR (101 MHz,  $CDCl_3$ )  $\delta$  173.4, 172.6, 171.0 (d,  $J_{C-P} = 5.9$  Hz), 155.82, 155.76, 150.6 (d,  $J_{C-P} = 10.2$  Hz), 150.3 (d,  $J_{C-P} = 10.2$  Hz), 150.2, 150.07, 150.0, 129.7, 129.6, 125.2 (d,  $J_{C-P} = 1.3$  Hz), 125.1 (d,  $J_{C-P} = 1.2$  Hz), 125.0 (d,  $J_{C-P} = 2.8$  Hz), 120.8 (d,  $J_{C-P} = 4.0$  Hz), 120.7 (d,  $J_{C-P} = 4.1$  Hz), 120.34 (d,  $J_{C-P} = 4.5$  Hz), 120.27 (d,  $J_{C-P} = 4.5$  Hz), 115.4, 114.5, 113.5, 79.63, 79.58, 60.4, 60.0, 56.8, 56.7, 55.7 (d,  $J_{C-P} = 150.8$  Hz), 55.0 (d,  $J_{C-P} = 151.6$  Hz), 47.8, 47.7, 35.2 (d,  $J_{C-P} = 4.9$  Hz), 34.9 (d,  $J_{C-P} = 5.2$  Hz), 31.31, 31.26, 30.3, 28.3, 28.3, 27.60, 27.55, 27.48, 27.45, 27.39, 25.2, 24.9, 19.7, 19.6, 17.3. ESI-MS ( $m/z$ ): calcd for  $C_{32}H_{46}N_3NaO_7P$  ( $M + Na$ )  $m/z$  638.2971 found 638.2923.



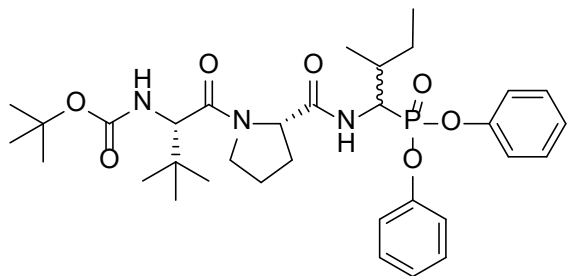
***Tert*-butyl-*N*-[(2*S*)-1-[(2*S*)-2-[[1-(diphenoxyphosphoryl)-2-methylbutyl]carbamoyl]pyrrolidin-1-yl]-3-methyl-1-oxopentan-2-yl]carbamate (8g)**



Using general procedure above, 0.14 g (0.44 mmol) of **7d** was coupled to 0.21 g (0.66 mmol) of **6d** to yield a colorless oil which formed a brown solid foam under reduced pressure (0.10 g, 37%).

HPLC  $t_r$  = 41.076, 41.471 and 42.094 min).  $^1\text{H}$  NMR (400 MHz,  $\text{CDCl}_3$ )  $\delta$  7.47 – 7.40 (m, 0.5H), 7.36 (d,  $J$  = 10.0 Hz, 0.5H), 7.30 – 7.22 (m, 4H), 7.20 – 7.06 (m, 6H), 5.30 – 5.24 (m, 0.5H), 5.11 (d,  $J$  = 9.5 Hz, 0.5H), 4.94 – 4.68 (m, 2H), 4.48 – 4.42 (m, 1H), 4.30 – 4.15 (m, 1H), 3.75 – 3.49 (m, 2H), 2.16 – 2.06 (m, 2H), 1.97 – 1.64 (m, 3H), 1.59 – 1.47 (m, 2H), 1.43 – 1.34 (m, 9H), 1.28 – 1.19 (m, 2H), 1.11 – 1.01 (m, 4H), 0.91 – 0.66 (m, 8H).  $^{13}\text{C}$  NMR (101 MHz,  $\text{CDCl}_3$ )  $\delta$  173.6, 172.61, 172.55, 171.3 (d,  $J_{\text{C-P}}$  = 6.4 Hz), 171.1 (d,  $J_{\text{C-P}}$  = 6.3 Hz), 170.0 (d,  $J_{\text{C-P}}$  = 6.3 Hz), 155.7, 155.6, 150.4 (d,  $J_{\text{C-P}}$  = 9.7 Hz), 150.3, 150.2, 150.10, 150.07, 150.0, 129.8, 129.7, 129.63, 129.56, 125.4, 125.3, 125.2 (d,  $J_{\text{C-P}}$  = 1.2 Hz), 125.0, 120.72 (d,  $J_{\text{C-P}}$  = 3.8 Hz), 120.65 (d,  $J_{\text{C-P}}$  = 3.9 Hz), 120.60, 120.56, 120.4 (d,  $J_{\text{C-P}}$  = 3.0 Hz), 115.4, 79.7, 79.6, 60.3, 59.9, 56.24, 56.15, 52.1, 51.8, 51.73, 51.65, 50.3, 50.2, 50.1, 49.9, 49.8, 48.3, 48.2, 47.9, 47.8, 37.9, 37.8, 37.7, 36.2, 36.14, 36.10, 35.6 (d,  $J_{\text{C-P}}$  = 3.6 Hz), 35.5 (d,  $J_{\text{C-P}}$  = 3.7 Hz), 35.40 (d,  $J_{\text{C-P}}$  = 4.1 Hz), 28.31, 28.29, 27.5, 27.4, 27.2, 27.1, 26.99, 26.96, 26.8, 25.21, 25.20, 24.9, 24.81, 24.76, 24.7, 24.30, 24.28, 24.1, 24.0, 23.0, 16.4 (d,  $J_{\text{C-P}}$  = 12.0 Hz), 16.3 (d,  $J_{\text{C-P}}$  = 11.7 Hz), 16.1 (d,  $J_{\text{C-P}}$  = 11.8 Hz), 15.50, 15.47, 15.3, 15.1 (d,  $J_{\text{C-P}}$  = 2.9 Hz), 15.0, 11.74, 11.73, 11.70, 11.69, 11.53, 11.51, 11.4, 11.1, 10.9. ESI-MS ( $m/z$ ): calcd for  $\text{C}_{33}\text{H}_{48}\text{N}_3\text{NaO}_7\text{P}$  ( $\text{M} + \text{Na}$ )  $m/z$  652.3128 found 652.3096.

**Tert-butyl-N-[(2*S*)-1-[(2*S*)-2-[[1-(diphenoxyphosphoryl)-2-methylbutyl]carbamoyl]pyrrolidin-1-yl]-3,3-dimethyl-1-oxobutan-2-yl]carbamate (**8h**)**



Using general procedure above, 0.50 g (1.50 mmol) of **7b** was coupled to 0.73 g (2.25 mmol) of **6d** to yield a colorless oil which formed a brown solid foam under reduced pressure (0.35 g, 37%).

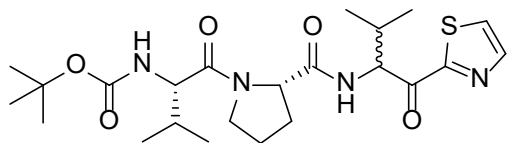
HPLC  $t_r$  = 41.897, 42.294 and 42.817 min.  $^1\text{H}$  NMR (400 MHz,  $\text{CDCl}_3$ )  $\delta$  7.40 (d,  $J$  = 10.4 Hz, 0.5H), 7.28 – 7.18 (m, 5H), 7.16 – 7.09 (m, 5.5H), 5.44 (d,  $J$  = 9.9 Hz, 1H), 4.96 – 4.70 (m, 2H), 4.49 (dd,  $J$  = 8.1, 3.7 Hz, 1H), 4.32 – 4.24 (m, 1H), 3.77 – 3.58 (m, 2H), 2.11 – 1.98 (m, 2H), 1.95 – 1.74 (m, 3H), 1.48 – 1.34 (m, 9H), 1.32 – 1.19 (m, 2H), 1.11 – 1.04 (m, 3H), 0.99 – 0.92 (m, 6H), 0.92 – 0.80 (m, 6H).  $^{13}\text{C}$  NMR (101 MHz,  $\text{CDCl}_3$ )  $\delta$  171.8, 171.7, 171.5, 155.7, 155.6, 150.4 (d,  $J_{C-P}$  = 9.7 Hz), 150.1 (d,  $J_{C-P}$  = 9.3 Hz), 129.79 (d,  $J_{C-P}$  = 1.0 Hz), 129.71, 129.58 (d,  $J_{C-P}$  = 0.9 Hz), 129.55 (d,  $J_{C-P}$  = 0.8 Hz), 129.3, 125.23, 125.19, 125.09, 125.0, 120.76 (d,  $J_{C-P}$  = 4.0 Hz), 120.74, 120.65, 120.61, 120.57, 120.47, 120.4, 120.3, 115.4, 79.6, 79.5 (d,  $J_{C-P}$  = 1.7 Hz), 60.3, 60.0, 58.3, 51.1 (d,  $J_{C-P}$  = 154.5 Hz), 51.0 (d,  $J_{C-P}$  = 154.8 Hz), 49.9, 48.6, 48.3, 36.0, 35.7, 35.64, 35.56, 28.4, 28.3, 28.0, 27.8, 26.9, 26.8, 26.21, 26.16, 25.3 (d,  $J_{C-P}$  = 2.1 Hz), 25.0, 24.7 (d,  $J_{C-P}$  = 5.2 Hz), 16.5 (d,  $J_{C-P}$  = 12.0 Hz), 16.1 (d,  $J_{C-P}$  = 11.8 Hz), 15.2, 14.3, 11.9, 11.71, 11.70, 11.6, 11.3. ESI-MS ( $m/z$ ): calcd for  $\text{C}_{33}\text{H}_{48}\text{N}_3\text{NaO}_7\text{P}$  ( $M + \text{Na}$ )  $m/z$  652.3128 found 652.3071.

**4.1.3. General Procedure for Synthesis of peptidyl  $\alpha$ -ketoheterocycles (**14a-e**).**

To a solution of heterocycle (1 equiv.) in 2 mL dry THF was added n-butyllithium (2.18 M, 8.3 equiv.) slowly over 3 min at  $-78^\circ\text{C}$ , and the mixture was stirred for an additional 10 min. A solution of **13a-e** (3.3 equiv.) in 1.5 mL dry THF was added rapidly over 1 minute and the mixture stirred for 15 minutes at  $-78^\circ\text{C}$ . The reaction mixture was poured into saturated  $\text{NH}_4\text{Cl}$  and extracted with EtOAc. The combined organic extracts were washed with saturated  $\text{NaHCO}_3$ , dried ( $\text{MgSO}_4$ ) and

concentrated *in vacuo*. The crude product was purified by column chromatography eluting with EtOAc: Hexane (1:2) to obtain compounds **14a-e** as mixtures of diastereomers.

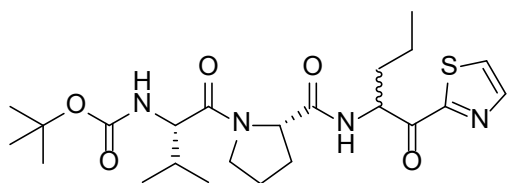
***Tert*-butyl-N-[(2*S*)-3-methyl-1-[(2*S*)-2-[(2*S*)-3-methyl-1-oxo-1-(1,3-thiazol-2-yl)butan-2-yl]carbamoyl]pyrrolidin-1-yl]-1-oxobutan-2-yl]carbamate (**14a**)**



Using the general method described for Peptidyl  $\alpha$ -keto heterocycles, 0.18 g (0.40 mmol) of **13a** was reacted with lithiated thiazole to yield a brown oil which formed

a light brown foam under reduced pressure. Recovered as a mixture of diastereomers (0.036 g, 54%). HPLC  $t_r$  = 22.702 and 23.587 min.  $^1\text{H}$  NMR (400 MHz,  $\text{CDCl}_3$ )  $\delta$  8.01 (d,  $J$  = 3.0 Hz, 1H), 7.67 (dd,  $J$  = 4.5, 3.0 Hz, 1H), 7.42 (d,  $J$  = 8.1 Hz, 1H), 5.58 (dd,  $J$  = 8.2, 4.9 Hz, 1H), 5.30 (d,  $J$  = 9.3 Hz, 1H), 4.63 (dd,  $J$  = 8.1, 2.9 Hz, 1H), 4.28 (dd,  $J$  = 9.4, 6.1 Hz, 1H), 3.75 – 3.57 (m, 2H), 2.43 – 2.25 (m, 2H), 2.14 – 1.87 (m, 4H), 1.41 (s, 9H), 1.04 – 0.94 (m, 6H), 0.93 (d,  $J$  = 6.8 Hz, 3H), 0.85 (d,  $J$  = 6.9 Hz, 3H).  $^{13}\text{C}$  NMR (101 MHz,  $\text{CDCl}_3$ )  $\delta$  191.6, 172.5, 171.0, 170.9, 165.4, 155.8, 145.0, 144.9, 126.6, 126.5, 79.5, 59.93, 59.91, 56.8, 47.7, 31.5, 31.0, 30.9, 28.3, 27.3, 26.9, 25.1, 25.0, 20.1, 19.9, 19.6, 19.6, 17.4, 17.3, 17.2, 17.0. ESI-MS ( $m/z$ ): calcd for  $\text{C}_{23}\text{H}_{36}\text{N}_4\text{NaO}_5\text{S}$  ( $M + \text{Na}$ )  $m/z$  503.2304 found 503.2272.

***Tert*-butyl-N-[(2*S*)-3-methyl-1-oxo-1-[(2*S*)-2-[(2*S*)-1-oxo-1-(1,3-thiazol-2-yl)pentan-2-yl]carbamoyl]pyrrolidin-1-yl]butan-2-yl]carbamate (**14b**)**

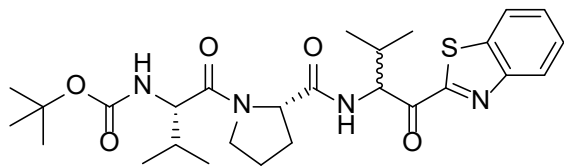


Using the general method described for Peptidyl  $\alpha$ -keto heterocycles, 0.18 g (0.40 mmol) of **13b** was reacted with lithiated thiazole to yield a brown oil which formed

a brown foam under reduced pressure (0.03 g, 45%). HPLC  $t_r$  = 24.086 and 24.852 min.  $^1\text{H}$  NMR (400 MHz,  $\text{CDCl}_3$ )  $\delta$  8.06 – 7.95 (m, 1H), 7.71 – 7.63 (m, 1H), 7.57 (d,  $J$  = 7.8 Hz, 0.5H), 7.45 (d,

$J = 7.5$  Hz, 0.5H), 5.67 – 5.56 (m, 1H), 5.31 – 5.22 (m, 1H), 4.74 – 4.58 (m, 1H), 4.35 – 4.22 (m, 1H), 3.75 – 3.56 (m, 2H), 2.44 – 2.25 (m, 2H), 2.12 – 1.85 (m, 5H), 1.71 – 1.49 (m, 2H), 1.45 – 1.37 (m, 9H), 1.03 – 0.86 (m, 9H).  $^{13}\text{C}$  NMR (101 MHz,  $\text{CDCl}_3$ )  $\delta$  208.9, 208.7, 191.7, 191.4, 172.9, 172.4, 170.8, 170.6, 164.79, 164.77, 155.8, 145.02, 144.97, 126.6, 126.5, 79.5, 59.7, 56.80, 56.77, 55.23, 55.20, 47.6, 47.5, 40.9, 39.5, 39.4, 38.6, 34.6, 33.4, 33.3, 31.4, 31.3, 28.3, 28.2, 27.5, 27.4, 27.2, 27.0, 25.52, 25.51, 25.1, 24.9, 23.9, 23.9, 22.2, 20.9, 20.8, 19.6, 19.4, 18.84, 18.83, 18.6, 18.5, 17.49, 17.47, 17.33, 17.19, 14.6, 13.84, 13.80, 13.79, 13.74, 13.72, 13.67. ESI-MS ( $m/z$ ): calcd for  $\text{C}_{23}\text{H}_{36}\text{N}_4\text{NaO}_5\text{S}$  ( $\text{M} + \text{Na}$ )  $m/z$  503.2304 found 503.2287.

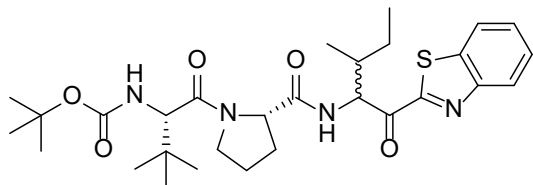
***Tert*-butyl-*N*-[(2*S*)-1-[(2*S*)-2-[(2*S*)-1-(1,3-benzothiazol-2-yl)-3-methyl-1-oxobutan-2-yl]carbamoyl]pyrrolidin-1-yl]-3-methyl-1-oxobutan-2-yl]carbamate (*I4c*)**



Using the general method described for Peptidyl  $\alpha$ -ketoheterocycles, 0.23 g (0.50 mmol) of **13a** was reacted with lithiated benzothiazole to yield a

colourless oil which formed a white foam under reduced pressure (0.10 g, 50%). HPLC  $t_r = 34.802$  and 35.474 min.  $^1\text{H}$  NMR (400 MHz,  $\text{CDCl}_3$ )  $\delta$  8.22 – 8.15 (m, 1H), 7.99 – 7.93 (m, 1H), 7.72 (d,  $J = 8.3$  Hz, 0.5H), 7.58 – 7.47 (m, 2.5H), 5.77 – 5.67 (m, 1H), 5.30 – 5.22 (m, 1H), 4.75 (dd,  $J = 8.1, 2.3$  Hz, 0.5H), 4.65 (dd,  $J = 8.1, 2.9$  Hz, 0.5H), 4.34 – 4.27 (m, 1H), 3.75 – 3.57 (m, 2H), 2.54 – 2.24 (m, 2H), 2.14 – 1.89 (m, 4H), 1.43 (s, 9H), 1.11 – 1.00 (m, 6H), 0.97 – 0.92 (m, 3H), 0.89 (dd,  $J = 8.7, 6.9$  Hz, 3H).  $^{13}\text{C}$  NMR (101 MHz,  $\text{CDCl}_3$ )  $\delta$  193.3, 193.1, 173.2, 172.6, 171.0, 170.9, 164.60, 164.55, 155.8, 153.6, 137.3, 137.2, 127.84, 127.81, 127.0, 125.79, 125.75, 122.31, 122.28, 79.6, 60.0, 59.9, 59.7, 56.8, 47.7, 47.6, 31.5, 31.4, 31.1, 30.9, 28.3, 27.3, 26.8, 25.1, 25.0, 20.0, 19.95, 19.7, 19.6, 17.5, 17.4, 17.3, 17.2. ESI-MS ( $m/z$ ): calcd for  $\text{C}_{27}\text{H}_{38}\text{N}_4\text{NaO}_5\text{S}$  ( $\text{M} + \text{Na}$ )  $m/z$  553.2461 found 553.2416.

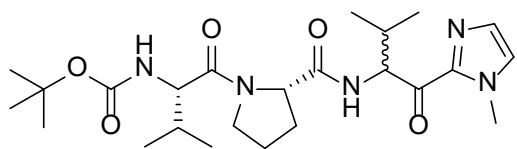
***Tert*-butyl-*N*-[(2*S*)-1-[(2*S*)-2-[(2*S*)-1-(1,3-benzothiazol-2-yl)-3-methyl-1-oxopentan-2-yl]carbamoyl]pyrrolidin-1-yl]-3,3-dimethyl-1-oxobutan-2-yl] carbamate (*14d*)**



Using the general method described for Peptidyl  $\alpha$ -ketoheterocycles, 0.50 g (0.30 mmol) of **13d** was reacted with lithiated benzothiazole to yield a dark

brown oil which formed a dark brown foam under reduced pressure (0.15 g, 28%). HPLC  $t_r$  = 44.367 and 45.645 min.  $^1\text{H}$  NMR (400 MHz,  $\text{CDCl}_3$ )  $\delta$  8.19 – 8.11 (m, 1H), 7.97 – 7.91 (m, 1H), 7.78 (d,  $J$  = 8.3 Hz, 0.5H), 7.57 – 7.47 (m, 2.5H), 5.89 (dd,  $J$  = 8.3, 3.7 Hz, 0.5H), 5.74 (dd,  $J$  = 8.2, 5.4 Hz, 0.5H), 5.30 (dd,  $J$  = 9.4, 5.9 Hz, 1H), 4.77 – 4.68 (m, 0.5H), 4.63 – 4.55 (m, 0.5H), 4.34 – 4.27 (m, 1H), 3.78 – 3.59 (m, 2H), 2.31 – 2.06 (m, 2H), 1.98 – 1.81 (m, 3H), 1.43 – 1.37 (m, 9H), 1.25 – 1.10 (m, 2H), 1.07 – 0.93 (m, 12H), 0.87 – 0.80 (m, 3H).  $^{13}\text{C}$  NMR (101 MHz,  $\text{CDCl}_3$ )  $\delta$  193.6, 192.9, 172.9, 172.1, 171.01, 170.9, 164.8, 164.4, 155.7, 153.6, 153.5, 137.3, 137.2, 127.8, 127.7, 127.0, 126.9, 125.8, 125.7, 122.3, 122.2, 79.6, 59.9, 59.6, 58.3, 48.6, 37.8, 36.9, 35.6, 35.5, 28.3, 27.3, 26.3, 25.2, 25.1, 24.3, 16.2, 14.0, 11.9, 11.4. ESI-MS ( $m/z$ ): calcd for  $\text{C}_{29}\text{H}_{42}\text{N}_4\text{NaO}_5\text{S}$  ( $M + \text{Na}$ )  $m/z$  581.2774 found 581.2748.

***Tert*-butyl-*N*-[(2*S*)-3-methyl-1-[(2*S*)-2-[(2*S*)-3-methyl-1-(1-methyl-1*H*-imidazol-2-yl)-1-oxobutan-2-yl]carbamoyl]pyrrolidin-1-yl]-1-oxobutan-2-yl]carbamate (*14e*)**



Using the general method described for Peptidyl  $\alpha$ -ketoheterocycles, 0.43 g (0.94 mmol) of **13a** was reacted with lithiated *N*-methylimidazole to yield a

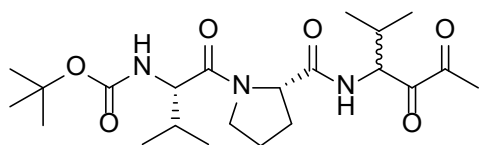
yellow oil which formed a light yellow foam under reduced pressure (0.07 g, 54%). HPLC  $t_r$  = 11.818 and 13.022 min.  $^1\text{H}$  NMR ( $\text{CD}_3\text{CN}$ , 400 MHz)  $\delta$  7.46 – 7.37 (m, 0.5H), 7.27 – 7.22 (m, 1.5H), 7.15 – 7.11 (m, 1H), 5.49 – 5.45 (m, 0.5H), 5.34 (dd,  $J$  = 9.3, 6.0 Hz, 0.5H), 4.53 – 4.44 (m, 1H), 4.25 – 4.18 (m, 1H), 3.95 – 3.90 (m, 3H), 3.74 – 3.61 (m, 2H), 2.33 – 2.25 (m, 1H), 2.08

– 1.88 (m, 6H), 1.44 – 1.38 (m, 9H), 1.00 – 0.96 (m, 3H), 0.94 – 0.87 (m, 6H), 0.84 (dd,  $J = 6.9$ , 1.2 Hz, 3H).  $^{13}\text{C}$  NMR (101 MHz,  $\text{CD}_3\text{CN}$ )  $\delta$  191.5, 191.2, 173.2, 173.1, 172.9, 172.7, 157.1, 143.1, 142.9, 130.2, 129.5, 80.0, 61.62, 61.59, 61.4, 60.6, 58.4, 48.8, 38.0, 37.0, 33.1, 32.7, 32.1, 30.1, 29.6, 29.0, 26.2, 26.0, 20.7, 20.6, 20.3, 20.7, 18.3, 18.1. ESI-MS ( $m/z$ ): calcd for  $\text{C}_{24}\text{H}_{39}\text{N}_5\text{O}_5$  (M + Na)  $m/z$  500.2843; found 500.2815.

#### 4.1.4. General Procedure for synthesis of peptidyl $\alpha$ -diketones (*18a-c*).

Wet DCM was prepared by solvating 2.6  $\mu\text{L}$  (1.1 equiv.) of water in 2.6 mL of DCM, and this was slowly added to a stirring solution of **17a-c** (1 equiv.) and DMP (2 equiv.) in 1.0 mL of dry DCM under nitrogen. The translucent solution grew cloudy towards the end of the reaction which took ~20 min. The reaction mixture was diluted in diethyl ether and concentrated under vacuum. The residue was further dissolved in 5 mL diethyl ether and washed with 5 mL of 1:1 10%  $\text{Na}_2\text{S}_2\text{O}_3$ : saturated  $\text{NaHCO}_3$  followed by 3 mL of water and 3 mL of brine. The aqueous fraction was extracted with diethyl ether (7 mL). The combined organic layers were washed with water and brine, and dried over  $\text{MgSO}_4$  and concentrated under vacuum. The crude product was purified by column chromatography. Compounds **18a-c** were eluted as a mixtures of diastereomers with EtOAc: Hexane (1:1).

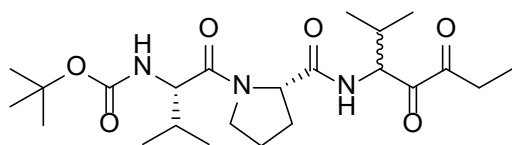
#### *Tert*-butyl-N-[(2*S*)-3-methyl-1-[(2*S*)-2-[(3*S*)-2-methyl-4,5-dioxohexan-3-yl]carbamoyl]pyrrolidin-1-yl]-1-oxobutan-2-yl]carbamate (**18a**)



Using the general method described for Peptidyl  $\alpha$ -diketones, 0.24 g (0.54 mmol) of **17a** was oxidized to yield a bright yellow solid which formed a yellow foam under reduced pressure (0.05 g, 21%). NMR and HPLC spectra revealed this to be a mixture of diastereomers. Purity not determined by HPLC.  $^1\text{H}$  NMR (400 MHz,  $\text{CDCl}_3$ )  $\delta$  7.62 (d,  $J = 6.8$  Hz,

0.5H), 7.48 (d,  $J = 6.2$  Hz, 0.5H), 5.26 – 5.15 (m, 1H), 4.90 – 4.80 (m, 1H), 4.68 – 4.54 (m, 1H), 4.31 – 4.21 (m, 1H), 3.72 – 3.51 (m, 2H), 2.39 – 2.22 (m, 1H), 2.14 – 1.77 (m, 5H), 1.48 – 1.33 (m, 9H), 1.12 – 0.71 (m, 15H).  $^{13}\text{C}$  NMR (101 MHz,  $\text{CDCl}_3$ )  $\delta$  196.77, 196.75, 196.6, 196.5, 173.20, 172.95, 171.03, 171.02, 155.8, 79.7, 59.4, 59.3, 58.3, 58.1, 56.83, 56.79, 47.7, 47.6, 31.4, 31.3, 29.7, 29.6, 28.3, 26.6, 26.5, 25.1, 25.0, 23.85, 23.82, 19.7, 19.6, 19.53, 19.46, 17.9, 17.7, 17.5, 17.3. ESI-MS ( $m/z$ ): calcd for  $\text{C}_{22}\text{H}_{37}\text{N}_3\text{NaO}_6$  ( $\text{M} + \text{Na}$ )  $m/z$  462.2580 found 462.2544.

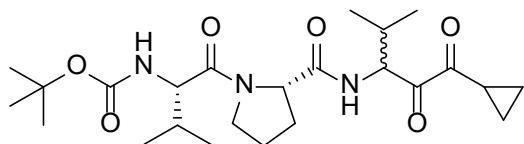
***Tert*-butyl-*N*-[(2*S*)-3-methyl-1-[(2*S*)-2-[(3*S*)-2-methyl-4,5-dioxoheptan-3-yl]carbamoyl]pyrrolidin-1-yl]-1-oxobutan-2-yl]carbamate (18b)**



Using the general method described for Peptidyl  $\alpha$ -diketones, 0.06 g (0.13 mmol) of **17b** was oxidized to yield a yellow solid which formed a yellow foam under

reduced pressure (0.04 g, 67%). Purity not determined by HPLC.  $^1\text{H}$  NMR (400 MHz,  $\text{CDCl}_3$ )  $\delta$  7.09 (d,  $J = 8.3$  Hz, 1H), 5.25 (d,  $J = 9.2$  Hz, 1H), 4.60 – 4.53 (m, 1H), 4.49 (dd,  $J = 8.3, 4.4$  Hz, 1H), 4.29 (dd,  $J = 9.3, 6.1$  Hz, 1H), 3.75 – 3.59 (m, 2H), 2.30 – 2.25 (m, 1H), 2.21 – 1.87 (m, 7H), 1.42 (s, 9H), 1.00 (dd,  $J = 7.0, 3.1$  Hz, 3H), 0.96 – 0.85 (m, 9H), 0.80 (dd,  $J = 6.9, 3.0$  Hz, 3H).  $^{13}\text{C}$  NMR (101 MHz,  $\text{CDCl}_3$ )  $\delta$  199.59, 199.55, 197.0, 196.9, 171.04, 171.00, 155.8, 79.6, 60.8, 59.4, 59.3, 58.5, 58.3, 56.8, 47.7, 47.6, 36.1, 31.4, 31.3, 29.66, 29.65, 29.62, 29.59, 28.3, 28.2, 26.7, 26.5, 25.1, 25.0, 19.8, 19.7, 19.6, 19.5, 18.69, 18.66, 17.8, 17.7, 17.5, 17.4, 17.3, 14.1, 6.80, 6.77. ESI-MS ( $m/z$ ): calcd for  $\text{C}_{23}\text{H}_{39}\text{N}_3\text{NaO}_6$  ( $\text{M} + \text{Na}$ )  $m/z$  476.2737 found 476.2715.

***Tert*-butyl-*N*-[(2*S*)-1-[(2*S*)-2-[(3*S*)-1-cyclopropyl-4-methyl-1,2-dioxopentan-3-yl]carbamoyl]pyrrolidin-1-yl]-3-methyl-1-oxobutan-2-yl]carbamate (18c)**



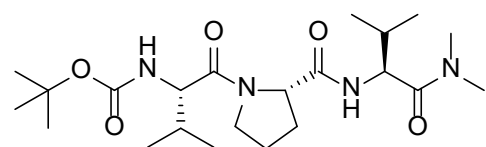
Using the general method described for Peptidyl  $\alpha$ -diketones, 0.18 g (0.38 mmol) of **17c** was oxidized to

yield a yellow solid which formed a yellow foam under reduced pressure (0.035 g, 20%). <95% purity by HPLC;  $t_r = 24.818$  and  $25.584$ .  $^1\text{H}$  NMR (400 MHz,  $\text{CDCl}_3$ )  $^1\text{H}$  NMR (400 MHz, Chloroform- $d$ )  $\delta$  7.59 (d,  $J = 7.4$  Hz, 0.5H), 7.41 (d,  $J = 7.1$  Hz, 0.5H), 5.27 – 5.20 (m, 1H), 5.04 – 4.93 (m, 1H), 4.70 – 4.59 (m, 1H), 4.33 – 4.24 (m, 1H), 3.73 – 3.54 (m, 2H), 2.71 – 2.61 (m, 1H), 2.40 – 2.28 (m, 1H), 2.14 – 1.81 (m, 5H), 1.46 – 1.36 (m, 9H), 1.16 – 1.04 (m, 4H), 1.03 – 0.79 (m, 12H).  $^{13}\text{C}$  NMR (101 MHz,  $\text{CDCl}_3$ )  $\delta$  198.32, 198.27, 196.6, 196.5, 173.2, 172.8, 171.0, 170.9, 155.8, 79.6, 59.6, 59.5, 58.4, 58.3, 56.8, 47.7, 47.6, 31.43, 31.37, 29.9, 29.72, 29.66, 28.3, 28.2, 26.9, 26.6, 25.1, 25.0, 19.8, 19.7, 19.6, 19.5, 17.6, 17.47, 17.45, 17.35, 15.53, 15.50, 13.8, 13.7, 13.49, 13.45. ESI-MS ( $m/z$ ): calcd for  $\text{C}_{24}\text{H}_{39}\text{N}_3\text{NaO}_6$  ( $\text{M}^+ \text{Na}$ )  $m/z$  488.2737 found 488.2698.

#### 4.1.5. General Procedure for synthesis of peptidyl N-methylamides (**24a-d**).

Compound **7a** (1 equiv.) was dissolved in 7 mL DMF. To this was added HBTU (1.2 equiv.) and DIEA (2.5 equiv.) to generate the activated ester. After 10 minutes, **23a-d** (1.5 equiv.) was added and the reaction mixture was stirred under nitrogen overnight. The reaction was diluted with EtOAc (50 mL) and an equal amount of  $\text{NaHCO}_3$ , washed with brine ( $6 \times 50$  mL), dried ( $\text{MgSO}_4$ ) and concentrated under vacuum. The crude product was subjected to column chromatography (100% EtOAc) which yielded **24a-d**.

#### *Tert*-butyl-N-[(2*S*)-1-[(2*S*)-2-[(1*S*)-1-(dimethylcarbamoyl)-2-methylpropyl]carbamoyl]pyrrolidin-1-yl]-3-methyl-1-oxobutan-2-yl]carbamate (**24a**)

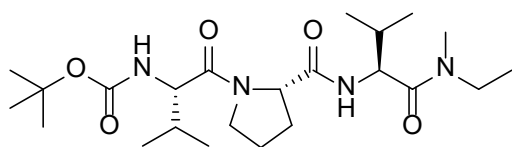


Using the procedure described for the synthesis of Peptidyl N-methylamides, 0.25 g (0.81 mmol) of **7a** was coupled to 0.17 g (1.18 mmol) of **23a** to yield a colourless oil which formed a white solid foam under reduced



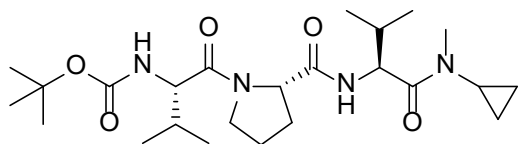
pressure (0.35 g, 99%). HPLC  $t_r$  = 12.232 min.  $^1\text{H}$  NMR (400 MHz,  $\text{CDCl}_3$ )  $\delta$  7.04 (d,  $J$  = 8.8 Hz, 1H), 5.35 (d,  $J$  = 9.3 Hz, 1H), 4.73 (dd,  $J$  = 8.8, 6.3 Hz, 1H), 4.51 (dd,  $J$  = 8.0, 3.4 Hz, 1H), 4.27 (dd,  $J$  = 9.4, 5.9 Hz, 1H), 3.74 – 3.57 (m, 2H), 3.06 (s, 3H), 2.93 (s, 3H), 2.15 – 2.06 (m, 2H), 2.02 – 1.89 (m, 4H), 1.39 (s, 9H), 0.97 (d,  $J$  = 6.8 Hz, 3H), 0.93 – 0.81 (m, 9H).  $^{13}\text{C}$  NMR (101 MHz,  $\text{CDCl}_3$ )  $\delta$  171.8, 171.5, 171.1, 155.9, 79.3, 60.1, 56.7, 53.6, 47.5, 37.3, 35.6, 31.7, 31.4, 28.3, 28.26, 25.1, 19.5, 19.5, 17.6, 17.2. ESI-MS ( $m/z$ ): calcd for  $\text{C}_{22}\text{H}_{40}\text{N}_4\text{NaO}_5$  (M + Na)  $m/z$  463.2896 found 463.2903.

***Tert*-butyl-*N*-[(2*S*)-1-[(2*S*)-2-[(1*S*)-1-[ethyl(methyl)carbamoyl]-2-methylpropyl]carbamoyl]pyrrolidin-1-yl]-3-methyl-1-oxobutan-2-yl]carbamate (**24b**)**



Using the procedure described for the synthesis of Peptidyl *N*-methylamides, 0.25 g (0.81 mmol) of **7a** was coupled to 0.19 g (1.21 mmol) of **23b** to yield a colourless oil which formed a white solid foam under reduced pressure (0.24 g, 66%). HPLC  $t_r$  = 15.392 min.  $^1\text{H}$  NMR (400 MHz,  $\text{CDCl}_3$ )  $\delta$  7.02 (d,  $J$  = 8.9 Hz, 1H), 5.38 – 5.26 (m, 1H), 4.71 – 4.64 (m, 1H), 4.53 – 4.45 (m, 1H), 4.27 (dd,  $J$  = 9.4, 5.8 Hz, 1H), 3.74 – 3.57 (m, 2H), 3.52 – 3.22 (m, 2H), 3.05 – 2.85 (2  $\times$  s, 3H), 2.15 – 2.05 (m, 2H), 2.00 – 1.89 (m, 4H), 1.39 (s, 9H), 1.18 – 1.05 (m, 3H), 0.97 (dd,  $J$  = 6.8, 0.9 Hz, 3H), 0.93 – 0.83 (m, 9H).  $^{13}\text{C}$  NMR (101 MHz,  $\text{CDCl}_3$ )  $\delta$  171.82, 171.79, 171.2, 171.13, 171.08, 170.8, 155.9, 79.3, 60.2, 56.7, 53.7, 53.6, 47.5, 44.4, 42.7, 34.8, 32.8, 32.0, 31.7, 31.4, 28.32, 28.27, 25.2, 25.1, 19.64, 19.57, 19.5, 17.6, 17.5, 17.24, 17.21, 12.1. ESI-MS ( $m/z$ ): calcd for  $\text{C}_{23}\text{H}_{42}\text{N}_4\text{NaO}_5$  (M + Na)  $m/z$  477.3053 found 477.3038.

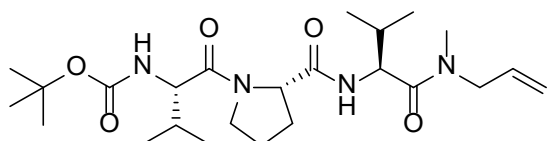
***Tert*-butyl-*N*-[(2*S*)-1-[(2*S*)-2-[(1*S*)-1-[cyclopropyl(methyl)carbamoyl]-2-methylpropyl]carbamoyl]pyrrolidin-1-yl]-3-methyl-1-oxobutan-2-yl]carbamate (**24c**)**



Using the procedure described for the synthesis of Peptidyl N-methylamides, 0.25 g (0.81 mmol) of **7a** was coupled to 0.21 g (1.21 mmol) of **23c** to yield a

light yellow oil which formed a light yellow foam under reduced pressure (0.32 g, 86%). HPLC  $t_r$  = 17.086 min.  $^1\text{H}$  NMR (400 MHz,  $\text{CDCl}_3$ )  $\delta$  6.97 (d,  $J$  = 9.2 Hz, 1H), 5.36 (d,  $J$  = 9.3 Hz, 1H), 5.16 (dd,  $J$  = 9.2, 7.0 Hz, 1H), 4.50 (dd,  $J$  = 8.0, 3.6 Hz, 1H), 4.26 (dd,  $J$  = 9.4, 5.9 Hz, 1H), 3.73 – 3.57 (m, 2H), 2.88 (s, 3H), 2.75 – 2.66 (m, 1H), 2.14 – 2.04 (m, 2H), 2.01 – 1.89 (m, 4H), 1.38 (s, 9H), 0.95 (d,  $J$  = 6.8 Hz, 3H), 0.91 – 0.85 (m, 9H), 0.84 – 0.75 (m, 2H), 0.71 – 0.54 (m, 2H).  $^{13}\text{C}$  NMR (101 MHz,  $\text{CDCl}_3$ )  $\delta$  174.4, 171.8, 171.1, 155.9, 79.3, 60.1, 56.7, 54.7, 47.5, 34.3, 31.9, 31.4, 31.1, 28.3, 28.2, 25.1, 19.6, 19.6, 17.8, 17.2, 10.2, 8.0. ESI-MS ( $m/z$ ): calcd for  $\text{C}_{24}\text{H}_{42}\text{N}_4\text{NaO}_5$  (M + Na)  $m/z$  489.3053 found 489.3008.

***Tert*-butyl-N-[(2*S*)-3-methyl-1-[(2*S*)-2-[(1*S*)-2-methyl-1-[methyl(prop-2-en-1-yl)carbamoyl]propyl]carbamoyl]pyrrolidin-1-yl]-1-oxobutan-2-yl]carbamate (**24d**)**

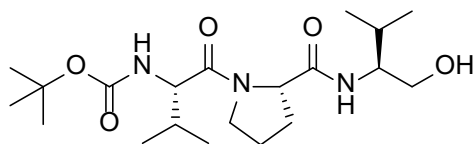


Using the procedure described for the synthesis of Peptidyl N-methylamides, 0.35 g (1.11 mmol) of **7a**

was coupled to 0.28 g (1.67 mmol) of **23a** to yield a light yellow oil which formed a light yellow solid foam under reduced pressure (0.46 g, 88%). HPLC  $t_r$  = 17.612 min.  $^1\text{H}$  NMR (400 MHz,  $\text{CDCl}_3$ )  $\delta$  7.12 – 7.05 (m, 1H), 5.78 – 5.61 (m, 1H), 5.44 – 5.35 (m, 1H), 5.18 – 5.05 (m, 2H), 4.75 – 4.61 (m, 1H), 4.56 – 4.47 (m, 1H), 4.26 (dd,  $J$  = 9.4, 6.0 Hz, 1H), 4.11 – 3.79 (m, 2H), 3.74 – 3.56 (m, 2H), 3.01 – 2.85 (2  $\times$  s, 3H), 2.16 – 2.05 (m, 2H), 2.00 – 1.87 (m, 4H), 1.39 (s, 9H), 0.96 (dd,  $J$  = 6.8, 2.1 Hz, 3H), 0.92 – 0.81 (m, 9H).  $^{13}\text{C}$  NMR (101 MHz,  $\text{CDCl}_3$ )  $\delta$  171.9, 171.7, 171.24, 171.22, 171.1, 155.9, 132.6, 132.5, 117.5, 117.4, 79.3, 60.12, 60.10, 56.7, 53.84, 53.76, 52.1, 50.2, 47.6, 34.8, 33.5, 31.8, 31.6, 31.4, 28.31, 28.27, 25.1, 19.7, 19.5, 19.5, 17.6, 17.5, 17.30, 17.26. ESI-MS ( $m/z$ ): calcd for  $\text{C}_{24}\text{H}_{42}\text{N}_4\text{NaO}_5$  (M + Na)  $m/z$  489.3053 found 489.3014.

#### 4.1.6. Synthesis of peptidyl valinol (**28**).

##### *Tert*-butyl N-[(2*S*)-1-[(2*S*)-2-[(2*S*)-1-hydroxy-3-methylbutan-2-yl]carbonyl]pyrrolidin-1-yl]-3-methyl-1-oxobutan-2-yl]carbamate (**28**)



Pre-synthesized Boc-Val-Pro-valinol benzyl ether **27** (0.19 g, 0.38 mmol) was dissolved in a mixture of methanol/water/acetic acid 9:0.9:0.1 (5 ml). Palladium-carbon 5% (0.11 g, 0.05 mmol) was added to this solution while stirring, then a balloon with hydrogen was attached and the flask was purged with hydrogen gas under vacuum. The mixture was stirred under hydrogen for 17 hr. TLC confirmed the completion of the reaction. The mixture was filtered over celite and the solvent was evaporated under reduced pressure. The crude product was purified by column chromatography (eluent: 100% chloroform) to give **28** (0.132 g, 86%). HPLC  $t_r = 32.542$  min  $^1\text{H}$ . NMR (400 MHz,  $\text{CDCl}_3$ )  $\delta$  7.01 (d,  $J = 7.1$  Hz, 1H), 5.31 (d,  $J = 9.2$  Hz, 1H), 4.55 (dd,  $J = 8.1, 3.2$  Hz, 1H), 4.26 (dd,  $J = 9.3, 6.3$  Hz, 1H), 3.99 (dd,  $J = 8.4, 5.6$  Hz, 1H), 3.76 – 3.56 (m, 5H), 2.36 – 2.24 (m, 1H), 2.05 – 1.76 (m, 5H), 1.40 (d,  $J = 2.1$  Hz, 9H), 1.00 – 0.83 (m, 12H).  $^{13}\text{C}$  NMR (400 MHz,  $\text{CDCl}_3$ )  $\delta$  172.8, 171.8, 171.7, 170.5, 157.0, 155.8, 80.8, 79.6, 63.6, 62.4, 61.2, 60.3, 57.7, 57.59, 57.56, 57.0, 47.8, 46.6, 31.7, 31.3, 30.5, 29.0, 28.8, 28.33, 28.31, 27.5, 25.2, 21.9, 19.7, 19.5, 19.4, 19.3, 18.8, 17.5, 17.4. ESI-MS ( $m/z$ ): calcd for calculated for  $\text{C}_{20}\text{H}_{37}\text{N}_3\text{NaO}_5$  ( $M + \text{Na}$ )  $m/z$  422.2650 found 422.2625.

**4.2. Protein Expression and Purification.** The expression and purification of recombinant CtHtrA was conducted according to our previously reported procedure.<sup>16</sup>

#### 4.3. Protease Assays.

*In vitro* CtHtrA inhibition studies were conducted as previously reported.<sup>17</sup> The assay was carried out in black 96-well plates at an excitation wavelength of 340 nm and emission wavelength of 405

nm on a POLARstar Optima (BMG Labtech) plate reader. Readings were obtained 50 cycles at the minimum cycle time of 10 flashes per well and a gain of 2000 with orbital shaking for 1 sec before each cycle. The activity of CtHtrA and its inhibition were measured by incubating the enzyme (14.4  $\mu$ L; 0.5 mg/mL) with 5  $\mu$ L of different concentrations of inhibitors (dissolved in DMSO) made up to a final volume of 150  $\mu$ L in 50 mM Tris/20 mM MgCl<sub>2</sub> (pH 7.0) buffer. Incubation was carried out at 37 °C for 20 min, this was followed by adding 10  $\mu$ L of the custom-synthesized fluorescent peptide MeOCoum-ENLHLPLPIIF-NH-DNP (Mimotopes) with 7-methoxycoumarin-4-acetic acid (MCA) as fluorophore (MeOCoum) and 2,4-dinitrophenol (DNP) as quencher; made to a concentration of 1 mg/mL in 50% isopropanol, and fluorescence readings were acquired immediately.

The inhibition of HNE, trypsin and chymotrypsin was conducted by modifying an existing method.<sup>37</sup> The activity of the enzymes was measured at 37 °C over a period of 5 min using AAPV-pNA (Sigma-Aldrich M4765), N $\alpha$ -Benzoyl-DL-R-pNA (Sigma-Aldrich B4875) and Suc-AAPF-pNA (Sigma-Aldrich S7388) as HNE, trypsin and chymotrypsin substrates respectively. Absorbance readings were measured at 405 nm and monitored at 15 sec intervals.

Enzyme inhibition was assessed by incubating 50  $\mu$ L of the appropriate enzyme solutions in the assay buffer (0.10 M Tris-HCl, pH 8.1, 0.02 M CaCl<sub>2</sub>) with 1.5  $\mu$ L of the inhibitors dissolved in DMSO, and made up to 100  $\mu$ L in the assay buffer. The mixture was incubated at 37 °C for 15 min before the addition of 50  $\mu$ L of the substrate which initiated the reaction to give an intense yellow coloration for controls and wells without enzyme inhibition.

All experiments were conducted in triplicate and inhibition was measured as the percentage of enzyme activity remaining. Compounds were tested against CtHtrA in 3 separate batches, each time giving a different IC<sub>50</sub> value for JO146 but with low errors across replicates (Table S3). The

trend of the results were consistent with HNE activity and are a reasonable representation of CtHtrA activity. Data analysis was conducted using GraphPad prism.

#### 4.4. *Chlamydia* cellular assays

*In vitro* *C. trachomatis* and *C. pecorum* cell culture assays were conducted according to our previously reported method.<sup>19</sup> *C. trachomatis* D (D/UW-3/Cx) and *C. pecorum* IPA (ATCC VR629) isolates were routinely cultured in McCoy B cells on DMEM, 10% fetal calf serum (FCS) at 37 °C, 5% CO<sub>2</sub>, 0.1 mg/mL streptomycin, and 0.05 mg/ml gentamycin. Inhibitor experiments were routinely conducted in 96-well plates seeded with 20,000 host cells per well 24 h prior to the Chlamydial infection. Infections were routinely conducted at a Multiplicity of Infection (MOI) of 0.3. The Inclusion Forming Units (IFU) were determined from cultures harvested at the completion of the developmental cycle during which inhibitor treatment was conducted. Briefly, JO146 and its analogues at 50 μM, 100 μM and 10 μM doses alongside a DMSO control (inhibitor solvent), were added at 16 h post infection (PI) and left in the cultures until completion of the developmental cycle (44 h PI). Harvested cultures were lysed by vigorous pipetting and serially diluted onto fresh monolayers, fixed and stained for enumeration of IFU/ml. Results obtained from these experiments are presented in tables S1 and S2 in the supporting information.

#### 4.5 Docking

Docking was carried out using GOLD v5.2 with a previously built homology model of CtHtrA.<sup>38-</sup>  
<sup>39</sup> The ligands (**14a** and **14c**) were prepared and docked based on mechanisms described by previous X-ray crystal structures of  $\alpha$ -ketoheterocycles covalently bound to serine proteases.<sup>21, 40</sup> Key hydrogen bonding constraints between the enzyme and ligands were pre-defined. These included a covalent bond between the carbonyl carbon alpha to the heterocycles and the catalytic serine hydroxyl oxygen forming a covalent hemiketal adduct, hydrogen bond between NH of P<sub>1</sub>

amide and C=O of Thr<sub>263</sub> and another hydrogen bond between C=O of P<sub>1</sub> backbone and NH of Ile<sub>265</sub> amide.

## **Abbreviations**

CtHtrA, *Chlamydia trachomatis* high temperature requirement A protease; SAR, structure activity relationship; HBTU, *O*-(Benzotriazol-1-yl)-*N,N,N',N'*-tetramethyluronium hexafluorophosphate; HPLC, high pressure liquid chromatography; DIEA, *N,N*-diisopropylethylamine; DMF, dimethylformamide; DMP, Dess-Martin periodinane; TFA, trifluoroacetic acid; IFU, inclusion forming unit; Val, valine; Tle, tertiary leucine; Pro, proline; Ile, isoleucine; Nva, norvaline; TLC, thin layer chromatography; Cbz, Carboxybenzyl; Cpr, cyclopropyl; PI, post infection; DMSO, dimethyl sulphoxide; MCA, 7-methoxycoumarin-4-acetic acid.

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## **Appendix A. Supplementary data**

Supplementary data to this article can be found online

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## Figure Captions

**Figure 1.** Structures of JO146 (**1**) and JCP83 (**2**) and the P<sub>3</sub>-P<sub>1</sub> protease nomenclature.<sup>18</sup> TSA – Transition State Analogue.

**Figure 2** *C. trachomatis* susceptibility to representative inhibitors (100 μM, 16 h PI) of different covalent and noncovalent transition state analogues during mid-replicative phase of the pathogen's developmental cycle. Inclusion forming units counts (IFU/mL) were determined upon completion of developmental cycle. Green (P<sub>3</sub> variants); brown (P<sub>1</sub> variants); purple (α-ketoheterocyclic variants); beige (α-diketone variants); light blue and lilac (P<sub>3</sub> and P<sub>1</sub> variants); grey (P<sub>3</sub> and P<sub>1</sub> α-ketoheterocycle). Error bars indicate the standard error of the mean obtained from experimental replicates (minimum of 5). \*Note in this assay 10<sup>3</sup> is the threshold for detect and can be equivalent to too few to accurately quantify.

**Figure 3:** Susceptibility of *C. pecorum* to CtHtrA inhibitors (100 μM; 16 h PI) during mid-replicative phase of the pathogen's developmental cycle. Infectious yield counts (IFU/mL) were

determined upon completion of developmental cycle. Colours the same as above in Figure 2.

Error bars indicate the standard error of the mean obtained from experimental replicates

(minimum of 5).

**Figure 4.** Binding poses of **14c**, magenta (A) and **14a**, cyan (B) to CtHtrA (transparent surface).

Oxygen is shown in red, nitrogen blue and sulphur in yellow.