
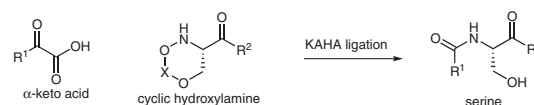


Synthesis and Evaluation of Cyclic Acetals of Serine Hydroxylamine for Amide-Forming KAHA Ligations

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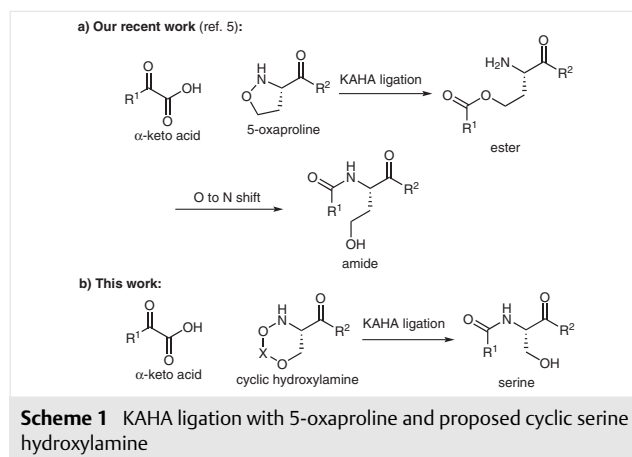
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Abstract The α -ketoacid–hydroxylamine (KAHA) ligation allows the coupling of unprotected peptide segments. The most widely used variant employs a 5-membered cyclic hydroxylamine that forms a homoserine ester as the primary ligation product. While very effective, monomers that give canonical amino acid residues are in high demand. In order to preserve the stability and reactivity of cyclic hydroxylamines, but form a canonical amino acid residue upon ligation, we sought to prepare cyclic derivatives of serine hydroxylamine. An evaluation of several cyclization strategies led to cyclobutanone ketals as the leading structures. The preparation, stability, and amide-forming ligation of these serine-derived ketals are described.

Key words ligation, hydroxylamines, acetals, amides, peptides

In 2006, we reported the α -ketoacid–hydroxylamine (KAHA) amide-forming ligation,¹ which complemented the well-established native chemical ligation² (NCL) for the chemical synthesis of proteins. These methods allow chemoselective couplings of unprotected peptide segments for the synthesis of peptides and proteins.³ Currently the most used hydroxylamine for KAHA ligations is 5-oxaproline, due to its high stability and chemoselective reactivity with α -ketoacids to form a homoserine residue at the ligation site.⁴ The primary product of this reaction is an ester, which rearranges to an amide under basic conditions (Scheme 1).⁵ In 2015, we developed a novel oxazetidine acid which gave serine at the ligation site, exclusively as the amide product.⁶ Unfortunately, the synthesis of this monomer is rather long and it is not stable in its unprotected form.

In this report, we describe the synthesis of a six-membered cyclic hydroxylamine, which directly yields a native

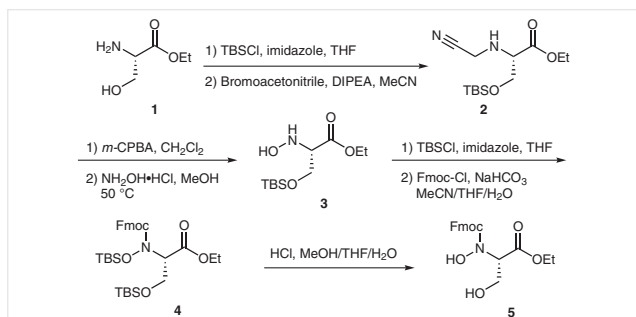


serine residue at the ligation site (Scheme 1). Furthermore, we describe our efforts to incorporate these novel monomers into a peptide segment.

In designing an alternative, serine-forming ligation monomer we sought to identify a larger ring structure that we hoped would be more easily prepared. This consideration led us to consider cyclic variants of L-serine hydroxylamine, including cyclic carbonates, cyclic sulfonates, and cyclic acetals. To begin, L-serine was converted into the corresponding ethyl ester **1**. TBS protection of the free alcohol and subsequent treatment with bromoacetonitrile in the presence of a base gave **2**.

This was treated with a slightly modified protocol from Fukuyama to give the *N*-hydroxyl intermediate **3**.⁷ Initially we envisioned selective Boc-protection of the nitrogen, but this was unsuccessful under various conditions.⁸ We instead elected to first protect the free hydroxyl group using TBSCl followed by N-protection with Fmoc-Cl to give **4**. Although initial attempts to remove the TBS protecting group

using TBAF led mainly to decomposition, treatment of **4** with concentrated HCl gave clean conversion into diol **5** (Scheme 2).



Scheme 2 Synthesis of core structure **5**

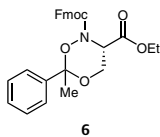
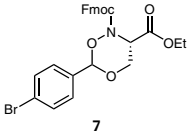
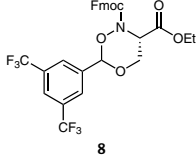
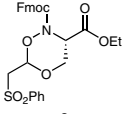
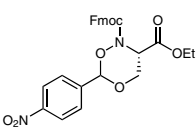
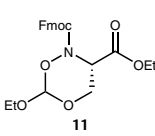
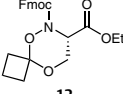
We explored the formation of the corresponding cyclic boronic esters, silanes, carbonates, sulfites, and phosphoric acid esters; however, we were not able to isolate the corresponding six-membered cyclic hydroxylamines. As an alternative, we considered cyclic acetals, as similar structures have proven to be well suited for KAHA ligation.⁹ For use in protein synthesis, however, these acid-labile functional groups would need to survive cleavage from the resin under standard TFA conditions. Prior to in-depth investigations of suitable acetals, we carried out preliminary tests to check that cyclic acetals of **5** could be formed and that Fmoc-removal was feasible. We treated compound **5** with 1,1-dimethoxyethylbenzene in the presence of *p*-toluenesulfonic acid to give **6** without problem. Removal of the Fmoc group under basic conditions also proceeded smoothly. With this promising result we sought to prepare and evaluate various cyclic acetals.

We initially targeted a methylene acetal, as these are known to be more resistant to acid cleavage than other structures.¹⁰ Unfortunately, all attempts to form the desired compound from **5** with various methylene sources using different catalysts or activation using Brønsted or Lewis acids did not yield the desired product. Despite this setback, we continued to synthesize various acetals and ketals by using either a catalytic amount of *p*-toluenesulfonic acid for aldehyde-derived compounds, or $\text{In}(\text{OTf})_3$ for the ketone derivatives. Selected acetals and ketals prepared are shown in Table 1. However, when these compounds were tested for their stability under SPPS conditions [resin cleavage conditions, TFA/DODT/ H_2O (95:2.5:2.5 v/v) for 2 h], only cyclobutane-derivative **12** showed reasonable stability (Table 1).

Despite the poor acid stability, we tested these substrates for activity in KAHA ligation. For this purpose, the Fmoc protecting group was removed with piperidine and the monomers were allowed to react with simple α -ketoac-

id **13** to give amide **14** (Table 2). Commonly reported KAHA conditions use DMSO/ H_2O at 60 °C, but a HFIP/AcOH mixture showed much better solubility and the reaction was observed to proceed at 45 °C.¹¹ Therefore all ligation studies were carried out under these conditions. Almost all monomers tested showed good activity in the KAHA ligation. After a period of 12 hours, α -ketoacid **13** was consumed. The benzylic acetals were stable under the acidic ligation conditions, and electron-poor nitrobenzyl acetal **16** gave slightly better conversion compared to bromobenzyl acetal **15**.

Table 1 Synthesized Monomers and Their Stability^a

Monomer	Stability to SPPS cleavage conditions
	not stable
	not stable
	not stable
	not stable
	not stable
	not stable
	stable

^a Stability was tested by treatment with TFA/DODT/ H_2O (95:2.5:2.5 v/v) for 2 h.

Table 2 KAHA Ligation with Selected Monomers^a

Monomer	Ligation yield after purification
<p>15</p>	64%
<p>16</p>	77%
<p>17</p>	no product observed
<p>18</p>	72%

monomer 16^b

monomer 18^c

^a KAHA ligation carried out at 30 mM at 45 °C in HFIP/AcOH with 1% H₂O for 12 h.

^b HPLC monitoring of the KAHA ligation of α -ketoacid **13** and hydroxylamine **16**, and purified amide **14**.

^c HPLC monitoring of the KAHA ligation of α -ketoacid **13** and hydroxylamine **18**, and purified amide **14**.

Cyclobutanone-derived acetal **18** was also stable during the ligation conditions and gave good conversion of the α -ketoacid. Only **17** did not perform KAHA ligation under HFIP/AcOH conditions because the deprotected monomer was not stable under the ligation conditions (Table 2).

This experiment also indicated that open, unsubstituted serine hydroxylamine does not react under these conditions and that the cyclic structure is essential. With these promising results, we sought to remove the ethyl ester in

the presence of Fmoc and the acetal, for the purpose of applying this monomer in Fmoc-SPPS. Despite all our efforts, the various conditions only led to total decomposition or returned starting material without removing the Fmoc protecting group.

We instead introduced a benzyl ester at the beginning of the synthesis. All previously developed steps were compatible with the benzyl-protected starting material, and we were able to synthesize the benzyl ester analogue **19**. Unfortunately, deprotection of the benzyl ester using Pd/C under a H₂ atmosphere gave mainly decomposition and only traces of product was observed. We were pleased to see that exchanging Pd/C for Pd(OH)₂ allowed a clean reaction.

Free acid **26** was coupled onto a short peptide segment, but proved not to be stable upon resin cleavage. Therefore we prepared and evaluated a number of other ketals from small rings. Although we were not successful in forming the product from oxetan-3-one, other cyclobutanone derivatives proved tractable. Brominated derivative **20** was readily formed and carboxylic esters **21** and **22** could be prepared from the corresponding acid followed by ester formation. The stability of compounds **19–22** was tested on the monomer and on a peptide segment (Table 3).

Table 3 Synthesized Monomers and Their Stability^a

Monomer	Stability to SPPS cleavage conditions	Stability to SPPS cleavage conditions on peptide
<p>19</p>	stable	not stable
<p>20</p>	stable	stable
<p>21</p>	stable	stable
<p>22</p>	stable ^b	stable ^b

^a Stability was tested by treatment with TFA/DODT/H₂O (95:2.5:2.5 v/v) for 2 h.

^b Observed as free acid on the cyclobutane ring.

All benzyl-protected monomers were also tested for their performance in the KAHA ligation. Fmoc removal occurred smoothly in all cases and the deprotected monomers underwent ligation with α -ketoacid **13** in HFIP/AcOH (Table 4). The cyclobutanone ketal monomers could be converted into the corresponding acid by treatment with

$\text{Pd}(\text{OH})_2$ under H_2 atmosphere (Scheme 3) and all compounds tolerated the coupling conditions to introduce the monomer onto the N-terminus of a peptide segment. Since OMe ester **21** showed better conversion in the KAHA ligation, we selected this monomer for introduction onto a short peptide sequence.

Table 4 KAHA Ligation with Selected Monomers^a

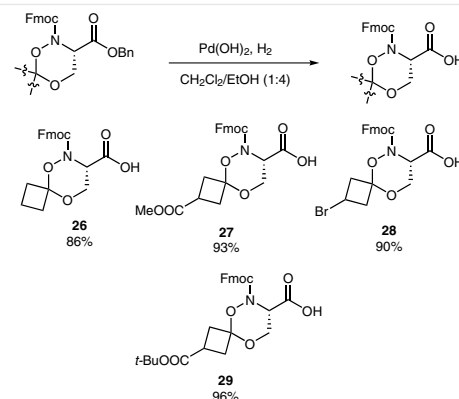
Monomer	Ligation yield after purification
 24	79%
 25	39%
13 and 24^b	13 and 25^c

^a KAHA ligation carried out at 30 mM at 45 °C in HFIP/AcOH with 1% H_2O for 12 h.

^b HPLC monitoring of the KAHA ligation of α -ketoacid **13** and hydroxylamine **24**, and purified amide **23**.

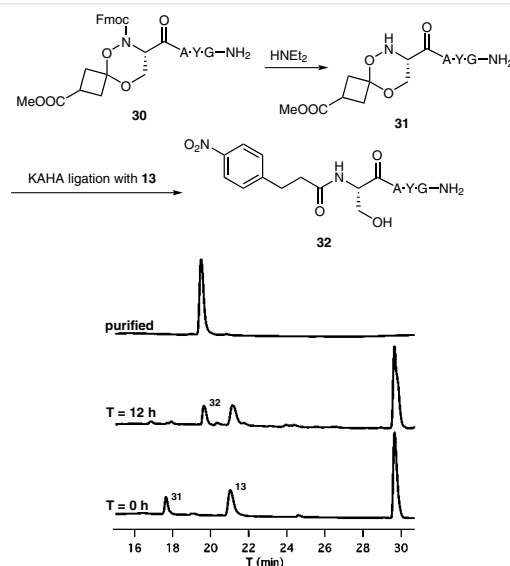
^c HPLC monitoring of the KAHA ligation of α -ketoacid **13** and hydroxylamine **25**, and purified amide **23**.

A short peptide segment with three amino acid residues, prepared by standard SPPS conditions on a Rink amide resin, was chosen as a model peptide. After deprotection of the terminal amine, the free acid of monomer **27** was coupled with HATU/NMM conditions to the resin and cleaved with TFA/DODT/ H_2O (95:2.5:2.5 v/v) for 2 hours. TFA was removed at 40 °C under reduced pressure and the product was isolated by preparative HPLC. The Fmoc protecting group was removed with *N,N*-diethylamine and the



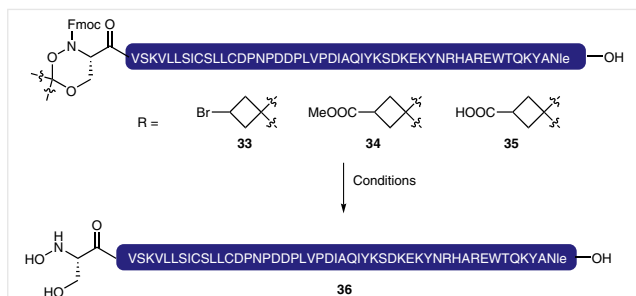
Scheme 3 Benzyl deprotection

KAHA ligation performed directly on the unprotected product without further purification using α -ketoacid **13** in HFIP/AcOH. The free hydroxylamine **31** was observed by LCMS and was completely consumed within 12 hours (Scheme 4).



Scheme 4 Ligation test on a small peptide segment

After this positive result, we expanded to a larger peptide fragment consisting of 50 amino acid residues. Although the Fmoc group could be removed cleanly, with a peak-to-peak conversion by HPLC, the acetal protecting group unfortunately did not survive the two-step procedure of basic Fmoc-removal conditions and acidic purification. The obtained product was **36**, the free hydroxylamine of serine at the N-terminus (Scheme 5).



Scheme 5 Attempts to remove the Fmoc protecting group on a longer peptide fragment

The results were the same regardless of the choice of base used to remove the Fmoc protecting group, including basic aqueous solutions. To date, we were not able to overcome this problem and longer peptides containing the cyclic ketal serine hydroxylamines cannot be isolated. A search for a more stable acetal protecting group is currently under investigation.

In conclusion, we devised a synthetic route to new hydroxylamine monomers that yield serine residues upon KAH ligations. We found that substituted cyclobutane ketals are stable under acidic conditions and are excellent ligation partners. Further investigations are necessary to apply this monomer on larger peptides, however this strategy provides a promising new approach to KAH ligation that forms canonical amino acid residues at the ligation site.

Fmoc-amino acids with suitable side-chain protecting groups, HATU (1-[bis(dimethylamino)methylene]-1*H*-1,2,3-triazolo[4,5-*b*]pyridinium 3-oxide hexafluorophosphate) were purchased from Peptides International (Louisville, KY, USA) and ChemImpex (Wood Dale, IL, USA). Solvents for flash chromatography (EtOAc, hexanes, MeOH) were of technical grade and distilled prior to use. HPLC grade MeCN from Sigma-Aldrich was used for analytical and preparative HPLC purification. DMF (>99.8%) from Sigma-Aldrich was directly used without further purification for solid phase peptide synthesis. Other commercially available reagents and solvents were purchased from Sigma-Aldrich (Buchs, Switzerland), Acros Organics (Geel, Belgium), and TCI Europe (Zwijndrecht, Belgium). DODT = 2,2'-(ethylenedioxy)diethanethiol. ¹H and ¹³C NMR spectra were recorded on Bruker DRX400, Bruker AVIII400 and Bruker AVIII600 spectrometers. HRMS were recorded by the Mass Service of the Laboratory of Organic Chemistry at ETH Zurich either with a Bruker maXis instrument (ESI-MS measurements) equipped with an ESI source and a Qq-TOF detector or with a Bruker solarix instrument (MALDI-FTICR-MS) using 4-hydroxy- α -cyanocinnamic acid as matrix. All reactions were performed using standard techniques under an atmosphere of N₂. Reactions and fractions from flash chromatography were monitored by TLC using aluminum plates (Merck, TLC Silica gel 60 W F₂₅₄) and visualized by staining with basic KMnO₄ solution or acidic ninhydrin solution. Flash chromatography was performed on Silicycle SiO₂ Type F60 (230–400 mesh) using a forced flow of air at 0.5–1.0 bar. Unless otherwise stated, peptides and protein segments were analyzed and purified by RP-HPLC on Jasco analytical and preparative instruments equipped with dual pumps, mixer and in-line degasser, a variable

wavelength UV detector (simultaneous monitoring of the eluent at 220 nm, 254 nm, and 301 nm) and a Rheodyne injector fitted with a 20 or 1000 μ L injection loop. If required, the columns were heated using a column heater or a water bath (preparative HPLC). The mobile phase for RP-HPLC was Milipore-H₂O containing 0.1% (v/v) TFA and HPLC grade MeCN containing 0.1% (v/v) TFA. Analytical HPLC was performed on Shiseido Capcell Pak C18 MGII (5 μ m, 4.6 mm i.d. \times 250 mm) or Shiseido Capcell Pak C18 (UG 80, 5 μ m, 4.6 mm i.d. \times 250 mm) columns at a flow rate of 1 mL/min. Preparative HPLC was performed on Shiseido Capcell Pak MGII (5 μ m, 20 mm i.d. \times 250 mm) or Vydac 248MS C18 (10 μ m, 22 mm i.d. \times 250 mm) columns at a flow rate of 10 mL/min.

Ethyl *O*-(*tert*-Butyldimethylsilyl)-*N*-(cyanomethyl)-*L*-serinate (2)

IR (neat): 2954, 2930, 2857, 1733, 1471, 1251, 1195 cm⁻¹.

¹H NMR (400 MHz, CDCl₃): δ = 4.19 (q, *J* = 7.1 Hz, 2 H), 3.92–3.84 (m, 2 H), 3.80–3.59 (m, 2 H), 3.48 (dt, *J* = 8.2, 4.4 Hz, 1 H), 2.28 (q, *J* = 7.0 Hz, 1 H), 1.27 (t, *J* = 7.2 Hz, 3 H), 0.85 (s, 9 H), 0.03 (d, *J* = 5.7 Hz, 6 H).

¹³C NMR (101 MHz, CDCl₃): δ = 171.54 (CO), 117.73 (CN), 64.41 (CH₂), 61.61 (CH), 61.39 (CH₂), 36.07 (CH₂), 25.82 (3 CH₃), 18.27 (C), 14.29 (CH₃), –5.41 (CH₃), –5.55 (CH₃).

HRMS (ESI): *m/z* [M + H]⁺ calcd for C₁₃H₂₇N₂O₃Si: 287.1785; found: 287.1786.

Ethyl *O*-(*tert*-Butyldimethylsilyl)-*N*-hydroxy-*L*-serinate (3)

IR (neat): 2953, 2929, 2857, 1737, 1471, 1252, 1107 cm⁻¹.

¹H NMR (400 MHz, CDCl₃): δ = 6.59 (br s, 1 H), 5.83 (br s, 1 H), 4.21 (qd, *J* = 7.2, 0.8 Hz, 2 H), 3.88 (qd, *J* = 10.2, 4.8 Hz, 2 H), 3.74 (dd, *J* = 5.6, 4.0 Hz, 1 H), 1.27 (t, *J* = 7.1 Hz, 3 H), 0.85 (s, 9 H), 0.03 (d, *J* = 3.3 Hz, 6 H).

¹³C NMR (101 MHz, CDCl₃): δ = 171.21 (CO), 66.83 (CH), 61.21 (CH₂), 61.08 (CH₂), 25.85 (3 CH₃), 18.29 (C), 14.31 (CH₃), –5.46 (CH₃), –5.53 (CH₃).

HRMS (ESI): *m/z* [M + H]⁺ calcd for C₁₁H₂₆NO₄Si: 264.1626; found: 264.1627.

Ethyl *N*-(*tert*-Butyldimethylsiloxy)-*O*-(*tert*-butyldimethylsilyl)-*N*-[(9*H*-fluoren-9-ylmethoxy)carbonyl]-*L*-serinate (4)

IR (neat): 2953, 2928, 2856, 1738, 1463, 1249, 1095 cm⁻¹.

¹H NMR (400 MHz, CDCl₃): δ = 7.79 (dt, *J* = 7.6, 0.9 Hz, 2 H), 7.69 (dt, *J* = 7.5, 1.0 Hz, 2 H), 7.42 (tt, *J* = 7.5, 0.8 Hz, 2 H), 7.32 (tt, *J* = 7.4, 1.1 Hz, 2 H), 4.72 (dd, *J* = 7.9, 6.3 Hz, 1 H), 4.55–4.39 (m, 2 H), 4.31 (t, *J* = 7.1 Hz, 1 H), 4.21 (qd, *J* = 7.1, 3.2 Hz, 2 H), 4.10 (dd, *J* = 7.2, 1.8 Hz, 2 H), 1.28 (t, *J* = 7.1 Hz, 3 H), 0.96 (s, 9 H), 0.91 (s, 9 H), 0.21 (d, *J* = 22.0 Hz, 6 H), 0.09 (s, 6 H).

¹³C NMR (101 MHz, CDCl₃): δ = 168.68 (CO), 159.38 (CO), 143.89 (C), 143.82 (C), 141.44 (C), 141.42 (C), 127.87 (CH), 127.86 (CH), 127.21 (2 CH), 125.49 (CH), 125.43 (CH), 120.10 (CH), 120.09 (CH), 68.36 (CH₂), 66.93 (CH), 61.45 (CH₂), 59.85 (CH₂), 47.13 (CH), 26.09 (3 CH₃), 26.00 (3 CH₃), 18.41 (2 C), 14.32 (CH₃), –4.70 (CH₃), –4.73 (CH₃), –5.21 (CH₃), –5.29 (CH₃).

HRMS (ESI): *m/z* [M + H]⁺ calcd for C₃₂H₅₀NO₆Si₂: 600.3171; found: 600.3174.

Ethyl *N*-[(9*H*-Fluoren-9-ylmethoxy)carbonyl]-*N*-hydroxy-*L*-serinate (5)

IR (neat): 2956, 1730, 1700, 1450, 1312, 1111, 1047 cm⁻¹.

^1H NMR (400 MHz, CDCl_3): δ = 7.78 (dt, J = 7.6, 1.0 Hz, 2 H), 7.62 (t, J = 6.2 Hz, 2 H), 7.42 (tt, J = 7.5, 1.3 Hz, 2 H), 7.36–7.29 (m, 2 H), 4.86 (s, 1 H), 4.57–4.42 (m, 2 H), 4.35–4.21 (m, 3 H), 4.21–4.06 (m, 3 H), 2.74 (s, 1 H), 1.29 (td, J = 7.1, 5.9 Hz, 3 H).

^{13}C NMR (101 MHz, CDCl_3): δ = 158.33 (CO), 143.54 (2 C)*, 143.48 (2 C), 141.43 (2 C)*, 141.43 (2 C), 128.02 (2 CH)*, 128.02 (2 CH), 127.32 (2 CH)*, 127.30 (2 CH), 125.25 (2 CH)*, 125.20 (2 CH), 120.19 (2 CH), 69.06 (CH_2), 64.07 (CH), 62.29 (CH_2), 60.57 (CH_2), 47.02 (CH), 14.35 (CH_3)*, 14.26 (CH_3); * signals of rotamers.

HRMS (MALDI): m/z [$\text{M} + \text{Na}$]⁺ calcd for $\text{C}_{20}\text{H}_{21}\text{NO}_6\text{Na}$: 394.1261; found: 394.1261.

Benzyl O-(tert-Butyldimethylsilyl)-N-(cyanomethyl)-L-serinate (benzyl-2)

IR (neat): 3674, 2987, 2956, 1736, 1463, 1406, 1252, 1083 cm^{-1} .

^1H NMR (400 MHz, CDCl_3): δ = 7.40–7.31 (m, 5 H), 5.19 (s, 2 H), 3.97–3.87 (m, 2 H), 3.81–3.71 (m, 1 H), 3.71–3.53 (m, 2 H), 2.31 (br s, 1 H), 0.86 (s, 9 H), 0.02 (d, J = 10.2 Hz, 6 H).

^{13}C NMR (101 MHz, CDCl_3): δ = 171.48 (CO), 135.39 (C), 128.76 (2 CH), 128.60 (CH), 128.49 (2 CH), 117.71 (CN), 67.17 (CH_2), 64.37 (CH_2), 61.68 (CH), 36.07 (CH_2), 25.85 (3 CH_3), 18.29 (C), –5.40 (CH_3), –5.51 (CH_3).

HRMS (ESI): m/z [$\text{M} + \text{Na}$]⁺ calcd for $\text{C}_{18}\text{H}_{28}\text{N}_2\text{NaO}_3\text{Si}$: 371.1761 found 371.1761.

Benzyl O-(tert-Butyldimethylsilyl)-N-hydroxy-L-serinate (benzyl-3)

IR (neat): 3674, 2987, 2900, 1739, 1462, 1252, 1192, 1105, 1066 cm^{-1} .

^1H NMR (400 MHz, CDCl_3): δ = 7.40–7.29 (m, 5 H), 5.87 (br s, 1 H), 5.58 (br s, 1 H), 5.21 (s, 2 H), 3.97–3.87 (m, 2 H), 3.81 (dd, J = 5.2, 4.1 Hz, 1 H), 0.86 (s, 9 H), 0.03 (s, 3 H), 0.02 (s, 3 H).

^{13}C NMR (101 MHz, CDCl_3): δ = 171.19 (CO), 135.58 (C), 128.74 (2 CH), 128.52 (CH), 128.41 (2 CH), 66.97 (CH_2), 66.90 (CH), 61.18 (CH_2), 25.89 (3 CH_3), 18.33 (C), –5.43 (CH_3), –5.48 (CH_3).

HRMS (ESI): m/z [$\text{M} + \text{H}$]⁺ calcd for $\text{C}_{16}\text{H}_{28}\text{NO}_4\text{Si}$: 326.1782; found: 326.1779.

Benzyl N-(tert-Butyldimethylsiloxy)-O-(tert-butyldimethylsilyl)-N-[(9H-fluoren-9-ylmethoxy)carbonyl]-L-serinate (benzyl-4)

IR (neat): 3674, 2956, 1740, 1450, 1361, 1250, 1098, 1076, 1005 cm^{-1} .

^1H NMR (400 MHz, CDCl_3): δ = 7.75 (dq, J = 7.5, 0.9 Hz, 2 H), 7.63–7.59 (m, 2 H), 7.46–7.18 (m, 9 H), 5.23–5.10 (m, 2 H), 4.71 (dd, J = 7.6, 6.5 Hz, 1 H), 4.49–4.35 (m, 2 H), 4.19 (t, J = 7.0 Hz, 1 H), 4.14–4.07 (m, 2 H), 0.93 (s, 9 H), 0.88 (s, 9 H), 0.20 (s, 3 H), 0.15 (s, 3 H), 0.07 (s, 3 H), 0.06 (s, 3 H).

^{13}C NMR (101 MHz, CDCl_3): δ = 168.55 (CO), 159.37 (CO), 143.85 (2 C)*, 143.78 (2 C), 141.41 (2 C)*, 141.38 (2 C), 135.47 (C), 128.62 (2 CH), 128.36 (CH), 128.27 (2 CH), 127.82 (2 CH)*, 127.80 (2 CH), 127.19 (2 CH), 125.40 (2 CH)*, 125.35 (2 CH), 120.06 (2 CH)*, 120.04 (2 CH), 68.29 (CH_2), 67.16 (CH), 67.08 (CH_2), 59.96 (CH_2), 47.08 (CH), 26.08 (3 CH_3), 25.99 (3 CH_3), 18.39 (C), 18.37 (C), –4.74 (2 CH_3), –5.23 (CH_3), –5.30 (CH_3); * signals of rotamers.

HRMS (ESI): m/z [$\text{M} + \text{H}$]⁺ calcd for $\text{C}_{37}\text{H}_{52}\text{NO}_6\text{Si}_2$: 662.3328; found: 662.3323.

Benzyl N-[(9H-Fluoren-9-ylmethoxy)carbonyl]-N-hydroxy-L-serinate (benzyl-5)

IR (neat): 3674, 3305, 2969, 2900, 1738, 1708, 1450, 1262, 1106, 1048 cm^{-1} .

^1H NMR (400 MHz, CDCl_3): δ = 7.73 (ddt, J = 7.6, 2.1, 0.9 Hz, 2 H), 7.53 (t, J = 8.3 Hz, 2 H), 7.37 (tdt, J = 7.5, 3.3, 0.8 Hz, 2 H), 7.35–7.17 (m, 7 H), 5.18 (s, 2 H), 4.88 (br s, 1 H), 4.47–4.34 (m, 2 H), 4.22–4.07 (m, 3 H), 3.13 (br s, 1 H); 1 H not observed.

^{13}C NMR (101 MHz, CDCl_3): δ = 168.65 (CO), 158.95 (CO), 143.39 (2 C), 141.37 (2 C), 134.97 (C), 128.76 (2 CH), 128.65 (CH), 128.22 (2 CH), 127.99 (2 CH)*, 127.97 (2 CH), 127.31 (2 CH)*, 127.29 (2 CH), 125.21 (2 CH)*, 125.16 (2 CH), 120.14 (2 CH), 69.16 (CH_2), 67.70 (CH_2), 64.31 (CH), 59.96 (CH_2), 46.91 (CH); * signals of rotamers.

HRMS (ESI): m/z [$\text{M} + \text{Na}$]⁺ calcd for $\text{C}_{25}\text{H}_{23}\text{NNaO}_6$: 456.1418; found: 456.1419.

Ketal Formation with In(OTf)₃; General Procedure

In(OTf)₃ (0.30 equiv) was added to a solution of **5** (1.00 equiv) and ketone (5.00 equiv) in CH_2Cl_2 (0.15 M) at 0 °C. The reaction was allowed to warm up to r.t. and was stirred overnight. The solution was diluted with CH_2Cl_2 and 10% citric acid solution was added. The phases were separated and the aqueous layer was extracted with CH_2Cl_2 (2 ×). The combined organic layers were washed with sat. aq. NaHCO_3 solution and brine and dried (Na_2SO_4). The drying agent was removed by filtration and the solvent was evaporated. The residue was purified by flash chromatography.

3-Ethyl 2-(9H-Fluoren-9-ylmethyl) (3S)-6-Methyl-6-phenyl-1,5,2-dioxazinane-2,3-dicarboxylate (6)

IR (neat): 2987, 2900, 1713, 1449, 1304, 1098, 1065 cm^{-1} .

^1H NMR (500 MHz, CDCl_3): δ = 7.78 (ddt, J = 7.6, 5.7, 0.9 Hz, 2 H), 7.77–7.62 (m, 2 H), 7.56–7.52 (m, 2 H), 7.46–7.27 (m, 7 H), 4.76–4.65 (m, 2 H), 4.45 (dd, J = 10.5, 7.7 Hz, 1 H), 4.38 (dd, J = 3.9, 1.6 Hz, 2 H), 4.33 (t, J = 7.1 Hz, 1 H), 4.20 (qd, J = 7.1, 1.6 Hz, 2 H), 1.78 (s, 3 H), 1.20 (t, J = 7.1 Hz, 3 H).

^{13}C NMR (126 MHz, CDCl_3): δ = 168.07 (CO), 156.52 (CO), 144.05 (C), 143.43 (C), 141.57 (C), 141.46 (C), 140.67 (C), 128.82 (CH), 128.30 (2 CH), 127.97 (2 CH)*, 127.92 (2 CH), 127.25 (2 CH)*, 127.20 (2 CH), 125.66 (CH), 125.35 (2 CH), 125.24 (CH), 120.14 (2 CH)*, 120.10 (2 CH), 103.89 (C), 68.23 (CH_2), 62.16 (CH_2), 59.91 (CH_2), 57.53 (CH), 47.14 (CH), 21.58 (CH_3), 14.22 (CH_3); * signals of rotamers/diastereomers.

HRMS (MALDI): m/z [$\text{M} + \text{H}$]⁺ calcd for $\text{C}_{28}\text{H}_{28}\text{NO}_6$: 474.1911; found: 474.1907.

7-Ethyl 6-(9H-Fluoren-9-ylmethyl) (S)-5,9-Dioxo-6-azaspiro[3.5]nonane-6,7-dicarboxylate (12)

IR (neat): 3674, 2987, 2901, 1714, 1450, 1408, 1313, 1058 cm^{-1} .

^1H NMR (500 MHz, CDCl_3): δ = 7.83–7.78 (m, 2 H), 7.78–7.62 (m, 2 H), 7.48–7.41 (m, 2 H), 7.38–7.31 (m, 2 H), 4.66 (dd, J = 10.3, 6.6 Hz, 1 H), 4.55 (s, 1 H), 4.43 (t, J = 9.2 Hz, 1 H), 4.40–4.33 (m, 2 H), 4.30 (q, J = 7.1 Hz, 2 H), 4.05 (dd, J = 11.5, 3.8 Hz, 1 H), 2.49–2.36 (m, 2 H), 2.36–2.24 (m, 2 H), 1.99–1.80 (m, 2 H), 1.29 (t, J = 7.1 Hz, 3 H).

^{13}C NMR (126 MHz, CDCl_3): δ = 168.06 (CO), 156.43 (CO), 144.05 (2 C)*, 143.37 (2 C), 141.50 (2 C)*, 141.43 (2 C), 127.98 (2 CH)*, 127.93 (2 CH), 127.25 (2 CH)*, 127.12 (2 CH), 125.58 (2 CH)*, 125.18 (2 CH), 120.15 (2 CH)*, 120.13 (2 CH), 104.99 (C), 68.50 (CH_2), 62.34 (CH_2), 60.27 (CH_2), 57.22 (CH), 47.03 (CH), 32.70 (CH_2), 30.37 (CH_2), 14.23 (CH_3), 11.59 (CH_2); * signals of rotamers/diastereomers.

HRMS (ESI): m/z [M + Na]⁺ calcd for C₂₄H₂₅NNaO₆: 446.1574; found: 446.1575.

7-Benzyl 6-(9H-Fluoren-9-ylmethyl) (S)-5,9-Dioxa-6-azaspiro[3.5]nonane-6,7-dicarboxylate (19)

IR (neat): 3674, 2987, 2900, 1746, 1716, 1451, 1407, 1260, 1098, 1065 cm⁻¹.

¹H NMR (500 MHz, CDCl₃): δ = 7.77 (dq, J = 7.6, 1.1 Hz, 2 H), 7.75–7.54 (m, 2 H), 7.47–7.21 (m, 9 H), 5.25 (s, 2 H), 4.56 (dd, J = 10.4, 6.7 Hz, 2 H), 4.40–4.31 (m, 2 H), 4.22 (t, J = 7.4 Hz, 1 H), 4.17–4.00 (m, 1 H), 2.50–2.19 (m, 4 H), 1.99–1.78 (m, 2 H).

¹³C NMR (126 MHz, CDCl₃): δ = 168.05 (CO), 156.45 (CO), 144.47 (2 C)*, 144.05 (2 C)*, 143.34 (2 C), 141.66 (2 C)*, 141.49 (2 C*), 141.41 (2 C), 135.22 (C), 128.63 (2 CH), 128.44 (CH), 128.09 (2 CH), 127.97 (2 CH)*, 127.91 (2 CH)*, 127.73 (2 CH), 127.24 (2 CH)*, 127.11 (2 CH), 125.60 (2 CH)*, 125.18 (2 CH), 124.84 (2 CH), 120.20 (2 CH)*, 120.13 (2 CH)*, 120.10 (2 CH), 105.02 (C), 68.47 (CH₂), 67.87 (CH₂), 60.30 (CH₂), 57.34 (CH), 46.96 (CH), 32.72 (CH₂), 30.41 (CH₂), 11.60 (CH₂); * signals of rotamers/diastereomers.

HRMS (ESI): m/z [M + NH₄]⁺ calcd for C₂₉H₃₁N₂O₆: 503.2177; found: 503.2174.

7-Benzyl 6-(9H-Fluoren-9-ylmethyl) (S)-2-Bromo-5,9-dioxa-6-azaspiro[3.5]nonane-6,7-dicarboxylate (20)

IR (neat): 3674, 2987, 2900, 1732, 1720, 1406, 1393, 1275, 1065, 1056 cm⁻¹.

¹H NMR (400 MHz, CDCl₃): δ = 7.76 (ddd, J = 8.4, 2.7, 1.0 Hz, 2 H), 7.55 (dd, J = 33.2, 7.5 Hz, 2 H), 7.43–7.22 (m, 9 H), 5.19 (s, 2 H), 4.62–4.32 (m, 4 H), 4.24–4.04 (m, 2 H), 3.90 (dd, J = 11.6, 3.7 Hz, 1 H), 3.03–2.87 (m, 2 H), 2.70–2.46 (m, 2 H).

¹³C NMR (101 MHz, CDCl₃): δ = 167.56 (CO), 156.01 (CO), 143.81 (2 C)*, 143.15 (2 C), 141.53 (2 C)*, 141.44 (2 C), 135.04 (C), 128.65 (2 CH), 128.52 (CH), 128.12 (2 CH), 128.03 (2 CH)*, 128.01 (2 CH), 127.26 (2 CH)*, 127.19 (2 CH), 125.19 (2 CH)*, 124.94 (2 CH), 120.20 (2 CH)*, 120.12 (2 CH), 102.37 (C), 67.97 (CH₂), 65.98 (CH₂), 60.86 (CH₂), 57.26 (CH), 47.07 (CH), 45.50 (CH₂), 43.97 (CH₂), 31.60 (CH); * signals of rotamers/diastereomers.

HRMS (MALDI/ESI): m/z [M + Na]⁺ calcd for C₂₉H₂₆BrNNaO₆: 586.0836; found: 586.0835.

(S)-7-(Benzyloxycarbonyl)-6-[(9H-fluoren-9-ylmethoxy)carbon-yl]-5,9-dioxa-6-azaspiro[3.5]nonane-2-carboxylic Acid (21a)

IR (neat): 3674, 2987, 2900, 1736, 1450, 1393, 1264, 1105, 1065 cm⁻¹.

¹H NMR (400 MHz, CDCl₃): δ = 10.67 (br s, 1 H), 7.81–7.73 (m, 2 H), 7.72–7.52 (m, 2 H), 7.48–7.21 (m, 9 H), 5.25 (s, 2 H), 4.72–4.15 (m, 5 H), 4.11–3.91 (m, 1 H), 3.09–2.95 (m, 1 H), 2.84–2.38 (m, 4 H).

¹³C NMR (101 MHz, CDCl₃): δ = 179.80 (CO)*, 179.74 (CO), 167.74 (CO)*, 167.67 (CO), 156.57 (CO)*, 156.20 (CO), 143.80 (2 C)*, 143.74 (2 C)*, 143.32 (2 C)*, 143.14 (2 C), 141.43 (2 C)*, 141.34 (2 C)*, 141.29 (2 C)*, 141.22 (2 C), 135.03 (C)*, 134.99 (C), 128.57 (2 CH)*, 128.55 (2 CH), 128.42 (CH)*, 128.40 (CH), 128.02 (2 CH)*, 128.01 (2 CH), 127.95 (2 CH)*, 127.93 (2 CH)*, 127.87 (2 CH)*, 127.83 (2 CH), 127.24 (2 CH)*, 127.19 (2 CH)*, 127.10 (2 CH), 125.52 (2 CH)*, 125.31 (2 CH)*, 125.26 (2 CH)*, 124.95 (2 CH), 120.10 (2 CH)*, 120.08 (2 CH)*, 119.96 (2 CH), 101.56 (C), 68.25 (CH₂)*, 67.90 (CH₂)*, 67.87 (CH₂), 60.53 (CH₂)*, 60.42 (CH₂), 57.44 (CH), 46.91 (CH)*, 46.72 (CH), 36.34 (CH₂)*, 36.15 (CH₂), 34.65 (CH₂)*, 34.21 (CH₂), 28.84 (C), 28.71 (CH); * signals of rotamers/diastereomers.

HRMS (ESI): m/z [M + H]⁺ calcd for C₃₀H₂₈NO₈: 530.1809; found: 530.1808.

7-Benzyl 6-(9H-Fluoren-9-ylmethyl) 2-Methyl (S)-5,9-Dioxa-6-azaspiro[3.5]nonane-2,6,7-tricarboxylate (21)

IR (neat): 3674, 2987, 2900, 1729, 1450, 1406, 1285, 1090, 1064 cm⁻¹.

¹H NMR (400 MHz, CDCl₃): δ = 7.80–7.75 (m, 2 H), 7.62 (ddd, J = 41.3, 13.0, 8.3 Hz, 2 H), 7.46–7.20 (m, 9 H), 5.24 (d, J = 3.1 Hz, 2 H), 4.65–4.34 (m, 3 H), 4.36–4.15 (m, 2 H), 4.09–3.94 (m, 1 H), 3.71 (d, J = 23.0 Hz, 3 H), 3.11–2.94 (m, 1 H), 2.85–2.40 (m, 4 H).

¹³C NMR (101 MHz, CDCl₃): δ = 174.67 (CO)*, 174.35 (CO), 171.26 (CO), 167.85 (CO)*, 167.75 (CO), 143.93 (2 C)*, 143.37 (2 C)*, 143.25 (2 C), 141.50 (2 C)*, 141.43 (2 C)*, 141.33 (2 C), 135.11 (C), 128.65 (2 CH)*, 128.63 (2 CH), 128.50 (CH)*, 128.48 (CH), 128.12 (2 CH), 128.01 (2 CH)*, 127.97 (2 CH)*, 127.93 (2 CH), 127.36 (2 CH)*, 127.28 (2 CH)*, 127.26 (2 CH)*, 127.15 (2 CH), 125.72 (2 CH)*, 125.37 (2 CH)*, 125.05 (2 CH), 120.17 (2 CH)*, 120.14 (2 CH)*, 120.08 (2 CH)*, 120.05 (2 CH), 101.79 (C), 101.57 (C), 68.90 (CH₂)*, 68.36 (CH₂), 67.98 (CH₂)*, 67.95 (CH₂), 60.63 (CH₂)*, 60.53 (CH₂), 57.57 (CH)*, 57.37 (CH), 52.26 (CH₃), 46.99 (CH)*, 46.84 (CH), 36.57 (CH₂)*, 36.42 (CH₂), 34.73 (CH₂)*, 34.28 (CH₂), 28.85 (C), 28.67 (CH); * signals of rotamers/diastereomers.

HRMS (MALDI/ESI): m/z [M + Na]⁺ calcd for C₃₁H₂₉NNaO₈: 566.1785; found: 566.1786.

7-Benzyl 2-tert-Butyl 6-(9H-Fluoren-9-ylmethyl) (S)-5,9-Dioxa-6-azaspiro[3.5]nonane-2,6,7-tricarboxylate (22)

IR (neat): 3674, 2974, 2900, 1722, 1451, 1366, 1283, 1157, 1089, 1012 cm⁻¹.

¹H NMR (400 MHz, MeOD): δ = 7.79 (ddt, J = 7.6, 3.1, 0.9 Hz, 2 H), 7.61 (dt, J = 27.2, 7.6 Hz, 2 H), 7.51–7.14 (m, 9 H), 5.39–5.02 (m, 3 H), 4.67–3.89 (m, 5 H), 2.98–2.14 (m, 5 H), 1.51–1.40 (m, 9 H).

¹³C NMR (101 MHz, MeOD): δ = 175.06 (CO)*, 174.93 (CO), 169.28 (CO)*, 169.17 (CO), 157.46 (CO), 145.13 (2 C)*, 144.99 (2 C)*, 144.70 (2 C), 142.69 (2 C)*, 142.63 (2 C)*, 142.56 (2 C)*, 142.51 (2 C), 136.75 (C)*, 136.69 (C), 129.53 (2 CH)*, 129.49 (2 CH)*, 129.47 (2 CH), 129.28 (CH), 129.08 (2 CH)*, 129.03 (2 CH)*, 128.96 (2 CH), 128.91 (2 CH)*, 128.90 (2 CH), 128.31 (2 CH)*, 128.26 (2 CH)*, 128.22 (2 CH)*, 128.17 (2 CH), 126.52 (2 CH)*, 126.29 (2 CH)*, 126.15 (2 CH)*, 125.96 (2 CH), 120.98 (2 CH)*, 120.94 (2 CH), 102.82 (C)*, 102.75 (C), 81.99 (C)*, 81.93 (C), 69.71 (CH), 69.02 (CH₂)*, 68.62 (CH₂), 68.56 (CH₂)*, 67.91 (CH₂), 61.73 (CH₂)*, 61.44 (CH₂), 59.36 (CH)*, 58.99 (CH), 48.22 (CH)*, 47.98 (CH), 37.03 (CH₂)*, 36.86 (CH₂), 35.34 (CH₂)*, 34.85 (CH₂), 31.19 (CH)*, 30.93 (CH)*, 30.59 (CH), 28.29 (3 CH₃)*, 28.25 (3 CH₃); * signals of rotamers/diastereomers.

HRMS (MALDI/ESI): m/z [M + Na]⁺ calcd for C₃₄H₃₅NNaO₈: 608.2255; found: 608.2257.

6-[(9H-Fluoren-9-yl)methyl] 2-Methyl (S)-7-[(S)-3-[2-[(S)-1-[(2-Amino-2-oxoethyl)amino]-3-(4-hydroxyphenyl)-1-oxopropan-2-yl]hydrazineyl]-2-oxobutanoyl]-5,9-dioxa-6-azaspiro[3.5]nonane-2,6-dicarboxylate (30)

HRMS (ESI): m/z [M + Na]⁺ calcd for C₃₈H₄₁N₅NaO₁₁: 766.2695; found: 766.2696.

HPLC (Shiseido Capcell Pak UG80 C18 column (4.6 × 250 mm), heated to 60 °C, 10 to 90% CH₃CN with 0.1% TFA in 20 min): t_R = 24.8 min.

Segment 33

HRMS (ESI): m/z [M]⁺ calcd for C₂₈₂H₄₃₁BrN₇₀O₈₃S₂: 6269.0282; found: 6269.0453.

HPLC (Shiseido Capcell Pak UG80 C18 column (4.6 × 250 mm), heated to 60 °C, 10 to 90% CH₃CN with 0.1% TFA in 20 min): *t_R* = 24.9 min.

Segment 34

HRMS (ESI): *m/z* [M]⁺ calcd for C₂₈₄H₄₃₄N₇₀O₈₅S₂: 6249.1231; found: 6249.0065.

HPLC (Shiseido Capcell Pak UG80 C18 column (4.6 × 250 mm), heated to 60 °C, 10 to 90% CH₃CN with 0.1% TFA in 20 min): *t_R* = 24.5 min.

Segment 35

HRMS (ESI): *m/z* [M]⁺ calcd for C₂₈₄H₄₃₃N₆₉O₈₅S₂: 6234.1122; found: 6235.1309.

HPLC (Shiseido Capcell Pak UG80 C18 column (4.6 × 250 mm), heated to 60 °C, 10 to 90% CH₃CN with 0.1% TFA in 20 min): *t_R* = 24.1 min.

Acetal Formation with *p*-Toluenesulfonic Acid Monohydrate; General Procedure

Dimethyl acetal (2.00 equiv) and *p*-toluenesulfonic acid monohydrate (0.10 equiv) were added to **5** (1.00 equiv) in DMF (0.05 M). The solution was stirred at 50 °C under vacuum for 4 h. The resulting dark brown viscous gel was diluted with CH₂Cl₂ and sat. aq NaHCO₃ solution was added to the solution. The aqueous layer was extracted with CH₂Cl₂ (2 ×). The combined organic layers were washed with brine and dried (MgSO₄). The drying agent was removed by filtration and the solvent was evaporated. The residue was purified by flash chromatography.

3-Ethyl 2-(9*H*-Fluoren-9-ylmethyl) (3*S*)-6-(4-Bromophenyl)-1,5,2-dioxazinane-2,3-dicarboxylate (**7**)

IR (neat): 3674, 2986, 2900, 1719, 1449, 1304, 1209, 1084 cm⁻¹.

¹H NMR (400 MHz, CDCl₃): δ = 7.79–7.20 (m, 12 H), 5.66 (s, 1 H), 4.80–4.63 (m, 3 H), 4.49 (t, *J* = 9.9 Hz, 1 H), 4.34–4.23 (m, 3 H), 4.20–4.06 (m, 1 H), 1.26 (t, *J* = 7.1 Hz, 3 H).

¹³C NMR (101 MHz, CDCl₃): δ = 167.75 (CO), 156.42 (CO), 143.92 (C), 143.11 (C), 141.54 (C), 141.42 (C), 133.85 (C), 131.67 (2 CH), 128.42 (2 CH), 128.02 (2 CH)*, 127.97 (2 CH), 127.22 (2 CH), 125.66 (2 CH)*, 125.39 (2 CH), 124.21 (CBr), 120.16 (2 CH), 103.06 (CH), 68.35 (CH₂), 66.12 (CH₂), 62.41 (CH₂), 57.26 (CH), 47.12 (CH), 14.27 (CH₃); * signals of rotamers/diastereomers.

HRMS (ESI): *m/z* [M + NH₄]⁺ calcd for C₂₇H₂₈BrN₂O₆: 555.1125; found: 555.1118.

3-Ethyl 2-(9*H*-Fluoren-9-ylmethyl) (3*S*)-6-[3,5-Bis(trifluoromethyl)phenyl]-1,5,2-dioxazinane-2,3-dicarboxylate (**8**)

IR (neat): 2970, 2901, 1727, 1450, 1277, 1175, 1132 cm⁻¹.

¹H NMR (400 MHz, CDCl₃): δ = 8.22–7.21 (m, 9 H), 5.82 (br s, 1 H), 4.90–4.66 (m, 3 H), 4.58 (dd, *J* = 10.6, 7.4 Hz, 1 H), 4.38–4.24 (m, 3 H), 4.22–4.09 (m, 1 H), 1.33–1.20 (m, 3 H); 2 H_{arom} not observed.

¹³C NMR (101 MHz, CDCl₃): δ = 167.51 (CO), 141.54 (C), 132.02 (q, *J* = 33.9 Hz, 2 CF₃), 128.08 (CH), 127.27 (CH), 125.30 (CH), 123.83 (CH), 120.15 (CH), 101.54 (CH), 68.71 (CH₂), 66.24 (CH₂), 62.62 (CH₂), 57.55, 47.14 (CH), 14.24 (CH₃); several C not assigned due to mixture of diastereomers and rotamers.

¹⁹F NMR (377 MHz, CDCl₃): δ = –62.80.

HRMS (ESI): *m/z* [M + NH₄]⁺ calcd for C₂₉H₂₇F₆N₂O₆: 613.1768; found: 613.1769.

3-Ethyl 2-(9*H*-Fluoren-9-ylmethyl) (3*S*)-6-[(Phenylsulfonyl)methyl]-1,5,2-dioxazinane-2,3-dicarboxylate (**9**)

IR (neat): 2987, 1736, 1447, 1306, 1147, 1116, 1065 cm⁻¹.

¹H NMR (500 MHz, CDCl₃): δ = 8.11–7.24 (m, 13 H), 5.05 (ddd, *J* = 20.8, 6.6, 4.1 Hz, 1 H), 4.50 (ddd, *J* = 14.1, 10.4, 7.3 Hz, 1 H), 4.45–4.29 (m, 2 H), 4.28–4.19 (m, 2 H), 4.17–4.00 (m, 0.5 H), 3.95 (dd, *J* = 10.1, 4.2 Hz, 0.5 H), 3.54 (dt, *J* = 14.5, 7.4 Hz, 1 H), 3.39 (dt, *J* = 14.3, 4.2 Hz, 1 H), 3.31 (d, *J* = 6.1 Hz, 2 H), 1.27 (q, *J* = 7.1 Hz, 3 H).

¹³C NMR (126 MHz, CDCl₃): δ = 168.26 (CO), 157.70 (CO), 143.93 (C), 143.70 (C)*, 141.42 (C), 141.36 (C), 139.67 (C), 139.48 (C), 134.18 (CH), 129.47 (2 CH), 128.15 (2 CH), 128.03 (CH), 127.89 (CH), 127.26 (2 CH), 125.62 (CH), 125.42 (CH), 120.08 (2 CH), 98.92 (CH)*, 97.72 (CH), 68.79 (CH₂)*, 68.72 (CH₂), 62.22 (CH₂)*, 62.19 (CH₂)*, 61.94 (CH₂), 59.35 (CH₂)*, 58.27 (CH₂), 53.83 (CH₂), 47.05 (CH), 14.27 (CH₃); * signals of rotamers/diastereomers.

HRMS (ESI): *m/z* [M + H]⁺ calcd for C₂₈H₂₈NO₈S: 538.1530; found: 538.1524.

3-Ethyl 2-(9*H*-Fluoren-9-ylmethyl) (3*S*)-6-(4-Nitrophenyl)-1,5,2-dioxazinane-2,3-dicarboxylate (**10**)

IR (neat): 3674, 2987, 2901, 1724, 1524, 1450, 1349, 1242, 1080 cm⁻¹.

¹H NMR (500 MHz, CDCl₃): δ = 8.26–8.21 (m, 2 H), 7.81–7.52 (m, 6 H), 7.45–7.27 (m, 4 H), 5.67 (s, 1 H), 4.74 (s, 2 H), 4.59 (dd, *J* = 10.6, 7.2 Hz, 1 H), 4.28 (dq, *J* = 14.2, 6.9 Hz, 3 H), 4.19–4.04 (m, 2 H), 1.25 (t, *J* = 7.1 Hz, 3 H).

¹³C NMR (126 MHz, CDCl₃): δ = 167.55 (CO), 156.33 (CO), 148.85 (CNO₂), 143.90 (2 C)*, 143.09 (2 C), 141.59 (2 C)*, 141.44 (C), 141.00 (2 C), 128.07 (2 CH)*, 128.04 (2 CH), 127.77 (2 CH), 127.24 (2 CH), 125.42 (2 CH)*, 125.29 (2 CH), 123.63 (2 CH), 120.21 (2 CH)*, 120.17 (2 CH), 101.98 (CH), 68.23 (CH₂), 66.17 (CH₂), 62.51 (CH₂), 57.34 (CH), 47.19 (CH), 14.26 (CH₃); * signals of rotamers/diastereomers.

HRMS (ESI): *m/z* [M + Na]⁺ calcd for C₂₇H₂₄N₂NaO₈: 527.1425; found: 527.1425.

3-Ethyl 2-(9*H*-Fluoren-9-ylmethyl) (3*S*)-6-Ethoxy-1,5,2-dioxazinane-2,3-dicarboxylate (**11**)

IR (neat): 2674, 2986, 2901, 1741, 1450, 1355, 1176, 1057 cm⁻¹.

¹H NMR (500 MHz, CDCl₃): δ = 7.86–7.61 (m, 4 H), 7.47–7.31 (m, 4 H), 5.43 (s, 1 H), 4.72–4.61 (m, 2 H), 4.57–4.31 (m, 2 H), 4.28 (q, *J* = 7.1 Hz, 2 H), 4.14–4.04 (m, 1 H), 3.88 (ddq, *J* = 44.5, 9.6, 7.1 Hz, 2 H), 1.37–1.24 (m, 6 H); 1 H not observed.

¹³C NMR (126 MHz, CDCl₃): δ = 167.46 (CO), 156.01 (CO), 143.85 (2 C)*, 143.31 (2 C), 141.50 (2 C)*, 141.44 (2 C), 128.02 (2 CH)*, 127.98 (2 CH), 127.26 (2 CH)*, 127.24 (2 CH), 125.50 (2 CH)*, 125.28 (2 CH), 120.15 (2 CH), 113.06 (CH), 68.54 (CH₂), 64.90 (CH₂), 63.20 (CH₂), 62.45 (CH₂), 47.07 (CH), 15.28 (CH₃), 14.24 (CH₃); * signals of rotamers/diastereomers.; 1 C not observed.

HRMS (ESI): *m/z* [M + NH₄]⁺ calcd for C₂₃H₂₉N₂O₇: 445.1969; found: 445.1966.

Fmoc Deprotection of Acid-Protected Monomer; General Procedure

Piperidine (5.00 equiv) was added dropwise to Fmoc-protected monomer (1.00 equiv) in CH₂Cl₂ (0.025 M). The reaction was stirred for 1 h at r.t. The solution was diluted with CH₂Cl₂ and sat. aq NH₄Cl solution was added. The phases were separated and the aqueous layer was extracted with CH₂Cl₂ (2 ×). The combined organic layers were

washed with brine and dried (Na_2SO_4). The drying agent was removed by filtration and the solvent was evaporated. The residue was purified by flash chromatography.

Ethyl (3S)-6-(4-Bromophenyl)-1,5,2-dioxazinane-3-carboxylate (15)

IR (neat): 3674, 2986, 2901, 1732, 1490, 1369, 1210, 1043 cm^{-1} .

^1H NMR (500 MHz, CDCl_3): δ = 7.50–7.45 (m, 2 H), 7.32–7.28 (m, 2 H), 6.86 (br s, 1 H), 5.80 (s, 1 H), 4.66 (dd, J = 11.4, 1.4 Hz, 1 H), 4.38–4.26 (m, 3 H), 3.52 (dd, J = 3.2, 1.4 Hz, 1 H), 1.34 (t, J = 7.1 Hz, 3 H).

^{13}C NMR (126 MHz, CDCl_3): δ = 170.61 (CO), 135.50 (C), 131.53 (2 CH), 128.30 (2 CH), 123.58 (CBr), 101.88 (CH), 65.27 (CH_2), 62.07 (CH_2), 58.68 (CH), 14.39 (CH_3).

HRMS (ESI): m/z [$\text{M} + \text{H}$] $^+$ calcd for $\text{C}_{12}\text{H}_{15}\text{BrNO}_4$: 316.0179; found: 316.0178.

Ethyl (3S)-6-(4-Nitrophenyl)-1,5,2-dioxazinane-3-carboxylate (16)

IR (neat): 3674, 2987, 2901, 1734, 1522, 1348, 1214, 1044 cm^{-1} .

^1H NMR (500 MHz, CDCl_3): δ = 8.22–8.18 (m, 2 H), 7.64–7.56 (m, 2 H), 6.94 (s, 1 H), 5.93 (s, 1 H), 4.70 (dt, J = 11.4, 1.4 Hz, 1 H), 4.37–4.29 (m, 3 H), 3.56 (dd, J = 3.3, 1.4 Hz, 1 H), 1.34 (t, J = 7.1 Hz, 3 H).

^{13}C NMR (126 MHz, CDCl_3): δ = 170.45 (CO), 148.55 (CNO_2), 142.82 (C), 127.64 (2 CH), 123.56 (2 CH), 100.98 (CH), 65.34 (CH_2), 62.20 (CH_2), 58.73 (CH), 14.40 (CH_3).

HRMS (ESI): m/z [$\text{M} + \text{H}$] $^+$ calcd for $\text{C}_{12}\text{H}_{15}\text{N}_2\text{O}_6$: 283.0925; found: 283.0927.

Ethyl (3S)-6-Ethoxy-1,5,2-dioxazinane-3-carboxylate (17)

IR (neat): 3674, 2987, 2900, 1736, 1393, 1250, 1074, 1056 cm^{-1} .

^1H NMR (500 MHz, CDCl_3): δ = 6.37 (d, J = 119.9 Hz, 1 H), 5.43 (dd, J = 11.2, 0.5 Hz, 1 H), 4.43–4.35 (m, 1 H), 4.26 (p, J = 7.2 Hz, 2 H), 4.09–4.01 (m, 1 H), 3.89–3.66 (m, 3 H), 1.33–1.28 (m, 3 H), 1.28–1.23 (m, 3 H).

^{13}C NMR (126 MHz, CDCl_3): δ = 169.44 (CO)*, 168.41 (CO), 111.02 (CH)*, 110.47 (CH), 62.92 (CH_2)*, 62.69 (CH_2), 62.63 (CH_2)*, 62.46 (CH_2), 61.97 (CH_2)*, 61.89 (CH_2), 57.65 (CH)*, 57.53 (CH), 15.16 (CH_3), 14.28 (CH_3)*, 14.26 (CH_3); * signals of diastereomers.

HRMS (ESI): m/z [$\text{M} + \text{Na}$] $^+$ calcd for $\text{C}_8\text{H}_{15}\text{NNaO}_5$: 228.0842; found: 228.0842.

Ethyl (S)-5,9-Dioxa-6-azaspiro[3.5]nonane-7-carboxylate (18)

IR (neat): 3674, 2987, 2901, 1733, 1451, 1375, 1280, 1079, 1020 cm^{-1} .

^1H NMR (500 MHz, CDCl_3): δ = 4.26 (q, J = 7.2 Hz, 2 H), 4.07 (d, J = 4.4 Hz, 2 H), 3.67 (t, J = 4.4 Hz, 1 H), 2.44–2.14 (m, 4 H), 1.81–1.71 (m, 2 H), 1.30 (t, J = 7.1 Hz, 3 H); NH not observed.

^{13}C NMR (126 MHz, CDCl_3): δ = 169.68 (CO), 103.05 (C), 61.79 (CH_2), 60.59 (CH_2), 58.37 (CH), 32.30 (CH_2), 31.64 (CH_2), 14.28 (CH_3), 11.85 (CH_2).

HRMS (ESI): m/z [$\text{M} + \text{H}$] $^+$ calcd for $\text{C}_9\text{H}_{16}\text{NO}_4$: 202.1074; found: 202.1074.

7-Benzyl 2-Methyl (S)-5,9-Dioxa-6-azaspiro[3.5]nonane-2,7-dicarboxylate (24)

IR (neat): 3674, 2987, 2900, 2253, 1733, 1453, 1393, 1258, 1065, 903 cm^{-1} .

^1H NMR (500 MHz, CDCl_3): δ = 7.45–7.27 (m, 5 H), 6.18 (s, 1 H), 5.29–5.21 (m, 2 H), 4.21–4.03 (m, 2 H), 3.70 (d, J = 1.1 Hz, 3 H), 3.66 (t, J = 3.8 Hz, 1 H), 2.99–2.40 (m, 5 H).

^{13}C NMR (126 MHz, CDCl_3): δ = 175.15 (CO), 169.67 (CO), 135.29 (C), 128.78 (2 CH), 128.66 (CH), 128.39 (2 CH)*, 128.37 (2 CH), 100.02 (C)*, 99.91 (C), 67.47 (CH_2), 60.68 (CH_2)*, 60.61 (CH_2), 58.49 (CH)*, 58.41 (CH), 52.14 (CH)*, 52.12 (CH), 36.54 (CH_2)*, 36.37 (CH_2), 35.54 (CH_2)*, 35.31 (CH_2), 29.09 (CH)*, 28.96 (CH); * signals of rotamers/diastereomers.

HRMS (ESI): m/z [$\text{M} + \text{H}$] $^+$ calcd for $\text{C}_{16}\text{H}_{20}\text{NO}_6$: 322.1285; found: 322.1286.

Benzyl (S)-2-Bromo-5,9-dioxa-6-azaspiro[3.5]nonane-7-carboxylate (25)

IR (neat): 3674, 3296, 2987, 2900, 1738, 1454, 1282, 1198, 1065 cm^{-1} .

^1H NMR (500 MHz, CDCl_3): δ = 7.44–7.33 (m, 5 H), 6.23 (s, 1 H), 5.33–5.23 (m, 2 H), 4.33–4.13 (m, 2 H), 4.10 (ddd, J = 11.5, 3.4, 2.7 Hz, 1 H), 3.68 (dt, J = 7.7, 3.9 Hz, 1 H), 3.37–2.57 (m, 4 H).

^{13}C NMR (126 MHz, CDCl_3): δ = 169.55 (CO)*, 169.45 (CO), 135.23 (C), 128.80 (2 CH), 128.71 (CH), 128.41 (2 CH)*, 128.39 (2 CH), 100.62 (C), 67.55 (CH_2), 60.88 (CH_2), 58.44 (CH)*, 58.41 (CH), 45.95 (CH_2)*, 45.78 (CH_2), 44.90 (CH_2)*, 44.57 (CH_2), 32.98 (CH)*, 32.75 (CH); * signals of rotamers/diastereomers.

HRMS (ESI): m/z [$\text{M} + \text{H}$] $^+$ calcd for $\text{C}_{14}\text{H}_{17}\text{BrNO}_4$: 342.0335; found: 342.0331.

Benzyl Deprotection; General Procedure

$\text{Pd}(\text{OH})_2/\text{charcoal}$ (20% on carbon wetted with ca. 50% water) (6% m/m) was added to a solution of benzyl-protected acid (1.00 equiv) in $\text{CH}_2\text{Cl}_2/\text{EtOH}$ (1:4) at 0 $^\circ\text{C}$. The mixture was stirred under H_2 atmosphere for 1 h while gradually warming up to r.t. The mixture was filtered through a plug of celite and the solvent was evaporated.

(S)-6-[(9H-Fluoren-9-ylmethoxy)carbonyl]-5,9-dioxa-6-azaspiro[3.5]nonane-7-carboxylic Acid (26)

IR (neat): 3674, 2987, 2900, 1717, 1450, 1394, 1266, 1133, 1065 cm^{-1} .

^1H NMR (500 MHz, CDCl_3): δ = 8.65 (br s, 1 H), 7.79–7.73 (m, 2 H), 7.64 (d, J = 45.5 Hz, 2 H), 7.44–7.37 (m, 2 H), 7.31 (tt, J = 7.4, 1.1 Hz, 2 H), 4.66 (dd, J = 10.5, 6.2 Hz, 1 H), 4.49 (s, 2 H), 4.38–4.22 (m, 2 H), 4.01 (s, 1 H), 2.44–2.18 (m, 4 H), 1.87 (dtd, J = 19.3, 12.1, 10.7, 7.0 Hz, 2 H).

^{13}C NMR (126 MHz, CDCl_3): δ = 173.61 (CO), 156.39 (CO), 143.82 (2 C)*, 143.19 (2 C), 141.50 (2 C)*, 141.48 (2 C), 128.05 (2 CH)*, 127.98 (2 CH), 127.39 (2 CH)*, 127.17 (2 CH), 125.42 (2 CH)*, 125.08 (2 CH), 120.14 (2 CH), 105.24 (C), 68.62 (CH_2), 60.08 (CH_2), 56.89 (CH), 46.99 (CH), 32.60 (CH_2), 30.25 (CH_2), 11.47 (CH_2); * signals of rotamers.

HRMS (ESI): m/z [$\text{M} + \text{Na}$] $^+$ calcd for $\text{C}_{22}\text{H}_{21}\text{NNaO}_6$: 418.1261; found: 418.1261.

(S)-6-[(9H-Fluoren-9-ylmethoxy)carbonyl]-2-(methoxycarbonyl)-5,9-dioxa-6-azaspiro[3.5]nonane-7-carboxylic Acid (27)

IR (neat): 3674, 2987, 2900, 1730, 1606, 1450, 1266, 1224, 1065 cm^{-1} .

^1H NMR (400 MHz, CDCl_3): δ = 7.78–7.73 (m, 2 H), 7.72–7.49 (m, 2 H), 7.45–7.28 (m, 4 H), 4.70–4.25 (m, 5 H), 4.14–3.91 (m, 1 H), 3.71 (d, J = 20.1 Hz, 3 H), 3.10–2.25 (m, 4 H).

^{13}C NMR (101 MHz, CDCl_3): δ = 174.71 (CO)*, 174.37 (CO), 172.75 (CO)*, 172.68 (CO), 158.09 (CO), 143.85 (2 C)*, 143.33 (2 C)*, 143.19 (2 C), 141.50 (2 C)*, 141.49 (2 C)*, 141.43 (2 C)*, 141.40 (2 C), 128.04 (2

CH)*, 127.99 (2 CH), 127.95 (CH), 127.39 (2 CH)*, 127.37 (2 CH), 127.19 (2 CH)*, 127.06 (2 CH)*, 127.04 (2 CH), 125.61 (2 CH)*, 125.32 (2 CH)*, 125.25 (2 CH)*, 125.02 (2 CH), 120.16 (2 CH)*, 120.09 (2 CH), 101.88 (C)*, 101.65 (C), 68.97 (CH₂)*, 68.34 (CH₂), 60.52 (CH₂)*, 60.42 (CH₂), 52.30 (CH₃), 47.06 (CH)*, 46.89 (CH), 36.54 (CH₂)*, 36.37 (CH₂), 34.60 (CH₂)*, 34.20 (CH₂), 28.79 (CH), 28.67 (C); * signals of rotamers/diastereomers.

HRMS (MALDI/ESI): m/z [M + Na]⁺ calcd for C₂₄H₂₃NNaO₈: 476.1316; found: 476.1316.

(S)-2-Bromo-6-[(9H-fluoren-9-ylmethoxy)carbonyl]-5,9-dioxo-6-azaspiro[3.5]nonane-7-carboxylic Acid (28)

IR (neat): 3674, 2987, 2900, 1717, 1450, 1408, 1283, 1264, 1139, 1067 cm⁻¹.

¹H NMR (400 MHz, CDCl₃): δ = 9.23 (br s, 1 H), 7.82–7.73 (m, 2 H), 7.59 (dd, J = 24.6, 7.5 Hz, 2 H), 7.46–7.28 (m, 4 H), 4.67 (d, J = 5.9 Hz, 2 H), 4.36–4.21 (m, 2 H), 4.20–3.97 (m, 1 H), 3.96–3.85 (m, 1 H), 2.95 (d, J = 15.7 Hz, 2 H), 2.70–2.45 (m, 2 H); 1 H not observed.

¹³C NMR (101 MHz, CDCl₃): δ = 173.11 (CO), 155.77 (CO), 143.71 (2 C)*, 143.08 (2 C), 141.55 (2 C)*, 141.51 (2 C), 128.09 (2 CH)*, 128.05 (2 CH), 127.37 (2 CH)*, 127.26 (2 CH), 125.05 (2 CH)*, 124.90 (2 CH), 120.21 (2 CH)*, 120.15 (2 CH), 102.49 (C), 68.01 (CH₂), 60.71 (CH₂), 47.16 (CH), 45.42 (CH₂), 43.85 (CH₂), 31.46 (CH); * signals of rotamers/diastereomers; 1 C not observed.

HRMS (MALDI/ESI): m/z [M + Na]⁺ calcd for C₂₂H₂₀BrNNaO₆: 496.0366; found: 496.0367.

(S)-2-(tert-Butoxycarbonyl)-6-[(9H-fluoren-9-ylmethoxy)carbonyl]-5,9-dioxo-6-azaspiro[3.5]nonane-7-carboxylic Acid (29)

IR (neat): 3674, 2976, 2901, 1719, 1450, 1366, 1283, 1135, 1060 cm⁻¹.

¹H NMR (400 MHz, CDCl₃): δ = 7.80–7.55 (m, 4 H), 7.44–7.28 (m, 4 H), 4.74–3.61 (m, 6 H), 3.03–2.01 (m, 5 H), 1.46 (d, J = 19.5 Hz, 9 H); OH not observed.

¹³C NMR (101 MHz, CDCl₃): δ = 173.54 (CO)*, 173.23 (CO), 173.10 (CO)*, 173.01 (CO), 156.42 (CO), 143.86 (2 C)*, 143.32 (2 C)*, 143.19 (2 C), 141.49 (2 C)*, 141.42 (2 C)*, 141.37 (2 C), 128.03 (2 CH)*, 127.96 (2 CH)*, 127.92 (2 CH), 127.41 (2 CH)*, 127.37 (2 CH)*, 127.19 (2 CH), 125.66 (2 CH)*, 125.34 (2 CH)*, 125.04 (2 CH), 120.16 (2 CH)*, 120.05 (2 CH), 101.97 (C), 101.65 (C), 81.06 (C)*, 81.01 (C), 68.91 (CH₂)*, 68.34 (CH₂), 60.44 (CH₂)*, 60.38 (CH₂), 57.20 (CH), 47.04 (CH), 46.88 (CH), 36.49 (CH₂)*, 36.21 (CH₂), 34.49 (CH₂)*, 34.01 (CH₂), 29.93 (CH)*, 29.70 (CH), 28.18 (3 CH₃); * signals of rotamers/diastereomers.

HRMS (MALDI/ESI): m/z [M + Na]⁺ calcd for C₂₇H₂₉NNaO₈: 518.1785; found: 518.1783.

KAHA Ligation; General Procedure

α -Ketoacid **13** (1.20 equiv) was added to the free hydroxylamine (1.00 equiv) and dissolved in HFIP/AcOH (1:1, 30 mM) containing 1% water. The mixture was heated to 45 °C for 12 h with monitoring by analytical HPLC. The crude mixture was directly purified by preparative HPLC (Shiseido Capcell Pak C18 column, 50 × 250 mm) heated to 60 °C using a gradient of 5 to 80% MeCN in Milipore-H₂O containing 0.1% (v/v) TFA over 35 min. HPLC monitoring of the KAHA ligation was carried out using Shiseido Capcell Pak UG80 C18 column (4.6 × 250 mm), heated to 60 °C, using a gradient of 10 to 90% MeCN in Milipore-H₂O containing 0.1% (v/v) TFA over 20 min.

Ethyl N-[3-(4-Nitrophenyl)propanoyl]-L-serinate (14)

IR (neat): 3384, 2987, 2901, 1734, 1646, 1517, 1451, 1346, 1228, 1056 cm⁻¹.

¹H NMR (500 MHz, MeOD): δ = 8.19–8.13 (m, 2 H), 7.51–7.46 (m, 2 H), 4.48 (dd, J = 5.1, 4.2 Hz, 1 H), 4.18 (qd, J = 7.2, 2.2 Hz, 2 H), 3.88–3.71 (m, 2 H), 3.07 (td, J = 7.4, 1.3 Hz, 2 H), 2.65 (td, J = 7.5, 2.1 Hz, 2 H), 1.25 (t, J = 7.1 Hz, 3 H).

¹³C NMR (126 MHz, MeOD): δ = 174.56 (CO), 171.76 (CO), 150.32 (C), 147.97 (CNO₂), 130.61 (2 CH), 124.53 (2 CH), 62.80 (CH₂), 62.46 (CH₂), 56.27 (CH), 37.50 (CH₂), 32.27 (CH₂), 14.43 (CH₃).

HRMS (ESI): m/z [M + Na]⁺ calcd for C₁₄H₁₈N₂NaO₆: 333.1057; found: 333.1061.

Benzyl N-(3-(4-Nitrophenyl)propanoyl)-L-serinate (23)

IR (neat): 2939, 1737, 1643, 1515, 1454, 1343, 1190 cm⁻¹.

¹H NMR (500 MHz, MeOD): δ = 8.14–8.10 (m, 2 H), 7.47–7.43 (m, 2 H), 7.38–7.28 (m, 5 H), 5.22–5.13 (m, 2 H), 4.55 (dd, J = 5.2, 4.2 Hz, 1 H), 3.90–3.73 (m, 2 H), 3.09–2.99 (m, 2 H), 2.63 (td, J = 7.5, 2.3 Hz, 2 H).

¹³C NMR (126 MHz, MeOD): δ = 174.64 (CO), 171.64 (CO), 150.28 (C), 147.95 (CNO₂), 137.20 (C), 130.57 (2 CH), 129.54 (2 CH), 129.27 (CH), 129.11 (2 CH), 124.52 (2 CH), 68.00 (CH₂), 62.75 (CH₂), 56.35 (CH), 37.47 (CH₂), 32.26 (CH₂).

HRMS (ESI): m/z [M + Na]⁺ calcd for C₁₉H₂₀N₂NaO₆: 395.1214.1057; found: 395.1218.

(S)-N-(2-amino-2-oxoethyl)-2-(2-((2S,5S)-6-hydroxy-5-(3-(4-nitrophenyl)propanamido)-3,4-dioxohexan-2-yl)hydrazineyl)-3-(4-hydroxyphenyl)propanamide (32)

HRMS (ESI): m/z [M + Na]⁺ calcd for C₂₆H₃₂N₆NaO₉: 595.2123; found: 595.2123.

HPLC (Shiseido Capcell Pak UG80 C18 column (4.6 × 250 mm), heated to 60 °C, 10 to 90% CH₃CN with 0.1% TFA in 20 min): t_R = 19.8 min.

Segment 36

HRMS (ESI): m/z [M]⁺ calcd for C₂₆₃H₄₁₈N₇₀O₈₁S₂: 5917.0183; found: 5918.0125.

HPLC (Shiseido Capcell Pak UG80 C18 column (4.6 × 250 mm), heated to 60 °C, 10 to 90% CH₃CN with 0.1% TFA in 20 min): t_R = 22.4 min.

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Supporting Information

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