

## Supporting Information © Wiley-VCH 2005

69451 Weinheim, Germany

## Orthogonally Protected Sugar Diamino Acids as Novel Building Blocks for Linear and Branched Oligosaccharide Mimetics

Frank Sicherl and Valentin Wittmann\*

## Experimental Section

17: β-Alanine amide 13 (27 mg, 0.067 mmol) was dissolved in dry DMF (4 mL) and treated with piperidine (1 mL). After 40 min, the solvent was removed under reduced pressure followed by co-evaporation with toluene. The residue was re-dissolved in CHCl<sub>3</sub> (3 mL) and 2 (44 mg, 0.081 mmol) and a solution of HBTU (31 mg, 0.081 mmol) and HOBt (19 mg, 0.121 mmol) in dry DMF (3 mL) were added.  $iPr_2NEt$  (31 μL, 0.202 mmol) was added and the mixture was stirred over night. After dilution with CHCl<sub>3</sub> (15 mL), the organic phase was washed with 0.1 N HCl and sat aq NaHCO<sub>3</sub>, dried with Na<sub>2</sub>SO<sub>4</sub> and evaporated. Purification by flash chromatography (silica, EtOAc/MeOH 95/5) gave 17 (42 mg, 89 %):  $R_F = 0.5$  (EtOAc/MeOH 9/1).

**18**: Peptide **17** (50 mg, 0.071 mmol) was dissolved in dry DMF (4 mL) and treated with piperidine (1 mL) for 30 min. After evaporation and co-evaporation with toluene, the remaining solid was dissolved in CHCl<sub>3</sub> (8 mL) and building block **1** (88 mg, 0.142 mmol), a solution of HATU (54 mg, 0.142 mmol) and HOAt (29 mg, 0.214 mmol) in DMF (8 mL), and iPr<sub>2</sub>NEt (55  $\mu$ L, 0.356 mmol) were added. After stirring over night, aqueous workup as described for **17** followed. Flash chromatography (silica, EtOAc/MeOH 95/5) gave **18** (61 mg, 79 %):  $R_F = 0.53$  (EtOAc/MeOH 9/1).

**19**: To a solution of **18** (50 mg, 0.046 mmol) in THF (4 mL) PMe<sub>3</sub> (278 μL, 1 m in THF) and water (1 mL) were added. After 1 h, the mixture was evaporated and co-evaporated several times with toluene. The subsequent peptide coupling was carried out as described for **17** using 2 eq of **1**. Purification by flash chromatography (silica, EtOAc/MeOH 95/5) gave **19** (56 mg, 73 %):  $R_F = 0.48$  (EtOAc/MeOH 9/1); RP-HPLC (Vydac 218TP54 C<sub>18</sub> reversed-phase column,  $4 \times 250$  mm, flow = 1 mL min<sup>-1</sup>, 20–80 % acetonitrile in water/0.1 % TFA over 30 min):  $t_R = 24.0$  min. HRMS (MALDI-FTICR), calcd for C<sub>83</sub>H<sub>110</sub>N<sub>8</sub>O<sub>27</sub>: 1673.73730 [ $M + Na^{+}$ ], found: 1673.73587,  $\Delta m = 0.8$  ppm.

**20**: Protected oligomer **19** (15 mg, 0.009 mmol) was dissolved in CHCl<sub>3</sub>/THF 1 : 1 (1 mL) and stirred for 1 h. The mixture was evaporated and the remainder was dissolved in MeOH (500  $\mu$ L), treated with 1 N HCl (500  $\mu$ L) for 1 h, and lyophilized. Finally, stirring with 20 % piperidine in DMF (500  $\mu$ L) led to complete deprotection. Purification by RP-HPLC (Vydac 218TP54 C<sub>18</sub> reversed-phase column, 4 × 250 mm, flow = 1 mL min<sup>-1</sup>, 1–100 % acetonitrile in water/0.13 % pentafluoropropionic acid over 30 min,  $t_R$  = 15.5 min) gave **20** • 4 F<sub>3</sub>C-CF<sub>2</sub>-CO<sub>2</sub>H (3.5 mg, 28 %). HRMS (ESI-FTICR, MeCN/H<sub>2</sub>O), calcd for C<sub>31</sub>H<sub>50</sub>N<sub>8</sub>O<sub>13</sub>: 743,35696 [M + H<sup>+</sup>], found: 743.35563,  $\Delta m$  = 1.8 ppm (see Figure S-1).

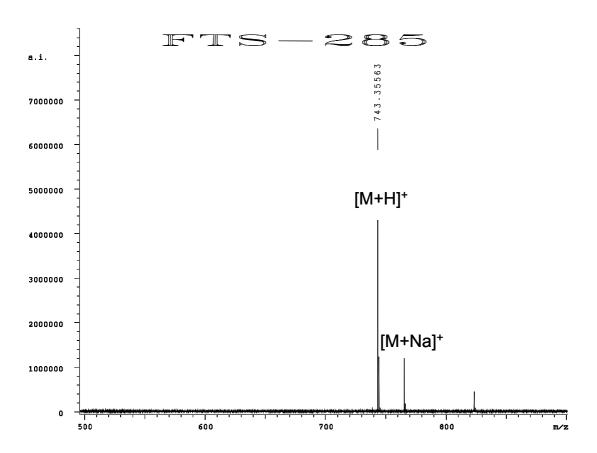


Figure S-1. HRMS (ESI-FTICR, MeCN/H<sub>2</sub>O) of **20**.