

Automated Synthesis of Peptoids and Peptide-Peptoid Hybrids



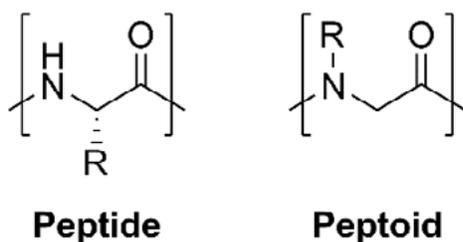
Summary

- The Liberty Blue™ automated microwave peptide synthesizer allows quick and efficient access to peptides, peptoids, and peptoid-peptide hybrids
- Peptide-peptoid hybrid Pro-Glu-(NLeu)-(NPhe)-Gly-(NLys)-NH₂ synthesized in 81% purity in under 2 hr.

Introduction

Peptoids are polymers of various N-substituted glycines. Though similar in structure to peptides (**Figure 1**), peptoids are resistant to proteolytic degradation, attributed to the complete substitution of their amide bonds. The increased stability of peptoids in vivo makes them an attractive peptidomimetic target for drug discovery and development.^{1,2}

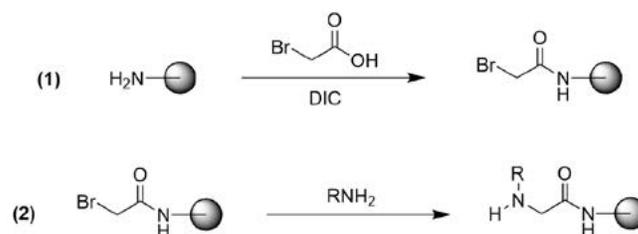
Figure 1



Comparison of peptide and peptoid structure

Peptoids and peptoid-peptide hybrids are typically synthesized through a “sub-monomer” process, which consists of two steps: (1) acylation with bromoacetic acid and N,N’-diisopropylcarbodiimide (DIC) and (2) nucleophilic displacement with a monosubstituted amine (**Figure 2**).^{1,3}

Figure 2



Typical synthesis of peptoids

Because many structurally diverse monosubstituted amines are commercially available, peptoids with a wide variety of side chains can be readily synthesized.^{2,3} However, conventional synthesis of peptoids can take up to three hours per residue.¹ Microwave irradiation has been shown to significantly reduce this time, making production of peptoid libraries and peptoid-peptide hybrids much more viable.¹⁻³

Materials and Methods

Reagents

All amino acids were obtained from CEM Corporation (Matthews, NC) and contained the following side chain protecting groups: Glu(OtBu) and Lys(Boc). Oxyma Pure and Rink Amide ProTide™ LL resin were obtained from CEM Corporation (Matthews, NC). Bromoacetic acid, N,N-diisopropylcarbodiimide (DIC), benzylamine, β-alanine t-butyl ester hydrochloride, piperidine, trifluoroacetic acid (TFA), 3,6-dioxa-1,8-octanedithiol (DODT), triisopropylsilane (TIS), and acetic acid were obtained from Sigma-Aldrich (St. Louis, MO). N-Boc-1,4-diaminobutane and isobutylamine were obtained

from Alfa Aesar (Ward Hill, MA). Dichloromethane (DCM), *N,N*-dimethylformamide (DMF), and anhydrous diethyl ether (Et₂O) were obtained from VWR (West Chester, PA). HPLC-grade water (H₂O), and HPLC-grade acetonitrile (MeCN) were obtained from Fisher Scientific (Waltham, MA).

Peptoid-Peptide Hybrid Synthesis: Pro-Glu-(NLeu)-(NPhe)-Gly-(NLys)-NH₂

The peptoid-peptide hybrid (**Figure 3**) was prepared at 0.1 mmol scale using the CEM Liberty Blue automated microwave peptide synthesizer on Rink Amide ProTide LL resin (0.18 meq/g substitution). For peptoid residues, deprotection was performed with piperidine in DMF, acylation was performed with bromoacetic acid and DIC, and nucleophilic displacement was performed with monosubstituted amine in DMF. For peptide residues, deprotection was performed with piperidine in DMF, and coupling reactions were performed with a 5-fold excess of Fmoc-AA-OH, DIC in DMF and Oxyma Pure in DMF. Cleavage was performed using the CEM Razor high-throughput peptide cleavage system with TFA/H₂O/TIS/ DODT. Following cleavage, the peptide was precipitated in Et₂O and lyophilized overnight.

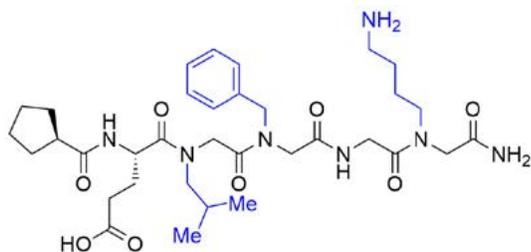


Figure 3
Target Peptoid-Peptide Hybrid: Pro-Glu-(NLeu)-(NPhe)-Gly-(NLys)-NH₂

Peptoid-Peptide Hybrid Analysis

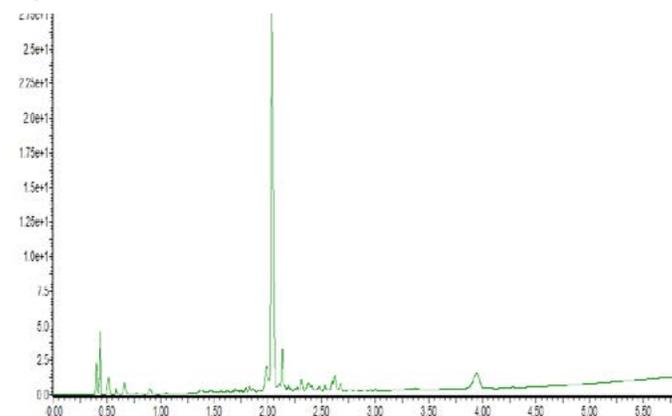
The peptoid-peptide hybrid was analyzed on a Waters Acquity UPLC system with PDA detector equipped with an Acquity UPLC BEH C8 column (1.7 mm and 2.1 x 100 mm). The UPLC system was connected to a Waters 3100 Single Quad MS for

structural determination. Peak analysis was achieved on Waters MassLynx software. Separations were performed with a gradient elution of 0.1% TFA in (i) H₂O and (ii) MeCN.

Results

Microwave-enhanced SPPS of Pro-Glu-(NLeu)-(NPhe)-Gly-(NLys)-NH₂ on the Liberty Blue automated microwave peptide synthesizer produced the target peptide in 81% purity (**Figure 4**).

Figure 4



UPLC Chromatogram of Pro-Glu-(NLeu)-(NPhe)-Gly-(NLys)-NH₂

Conclusion

The CEM Liberty Blue automated microwave peptide synthesizer allows quick and efficient access to peptides, peptoids, and peptoid-peptide hybrids. Microwave-enhanced SPPS produced peptoid-peptide hybrid, Pro-Glu-(NLeu)-(NPhe)-Gly-(NLys)-NH₂, in 81% purity.

References:

- (1) Olivos, H. J.; Alluri, P. G.; Reddy, M. M.; Salony, D.; Kodadek, T. *Org. Lett.* **2002**, *4*, 4057–4059.
- (2) Unciti-Broceta, A.; Diezmann, F.; Ou-Yang, C. Y.; Fara, M. A.; Bradley, M. *Bioorg. Med. Chem.* **2009**, *17*, 959–966.
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United States (Headquarters)

800-726-3331
704-821-7015
Fax: 704-821-7894
info@cem.com

Italy

(39) 35-896224
Fax: (39) 35-891661
info.srl@cem.com

France

33 (01) 69 35 57 80
Fax: 33 (01) 60 19 64 91
info.fr@cem.com

Japan

+81-3-5793-8542
Fax: +81-3-5793-8543
info@cemjapan.co.jp

Germany, Austria, Switzerland

(49) 2842-9644-0
Fax: (49) 2842-9644-11
info@cem.de

United Kingdom

(44) 1280-822873
Fax: (44) 1280-822873
info.uk@cem.com

Ireland

+353 (0) 1 885 1752
Fax: +353 (0) 1 885 1601
info.ireland@cem.com

www.cem.com

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