

# Synthesis of a Biotin-Derived Alkyne for Pd-Catalyzed Coupling Reactions

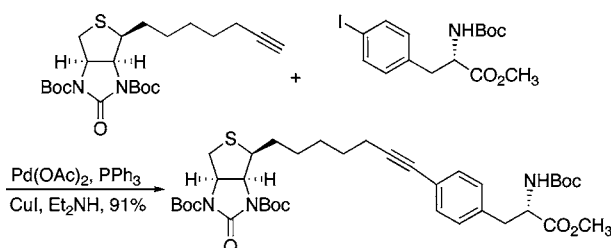
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## ABSTRACT



An efficient synthesis of a terminal alkyne derived from *d*-biotin was developed to provide an alternative for carboxyl-based biotinylation. This approach was illustrated by the preparation of alkyne- and alkene-linked phenylalanine derivatives using palladium-catalyzed Sonogashira and Oh methodology. (Strept)avidin binding was observed using soluble and immobilized receptors. These results demonstrate the applicability of carbon-linked biotin derivatives for use in affinity-based purification and bioanalytical applications.

The interaction between biotin (Vitamin H) **1** and the glycoprotein avidin is one of the strongest noncovalent associations known ( $K_a \sim 2.5 \times 10^{15} \text{ M}^{-1}$ ). Avidin is tetrameric, and each subunit is capable of binding a biotin ligand. The high binding affinity and exceptionally slow dissociation rates of biotin result from a network of hydrogen bonds between the receptor and the heterocyclic 7-oxo-3-thia-6,8-diazabicyclo[3.3.0]oct-4-yl ligand. The ureido nitrogens of biotin form hydrogen bonds with Thr35 and Asn118, the oxygen contacts of Ser16 and Tyr33. Additional hydrogen bonding occurs between the biotin carboxylic acid and avidin residues Ala39, Thr40, and Ser75. The hydrophobic tetrahydrothiophene interacts with Phe79, Trp97, and Trp110.<sup>1</sup>

Avidin and the related tetrameric protein streptavidin share ~33% of the conserved amino acids and a strong biotin binding affinity ( $K_a \sim 1.0 \times 10^{12} \text{ M}^{-1}$ ).<sup>2</sup> Specific interactions include hydrogen bonding between biotin urea nitrogens with Ser45 and Asp128 and oxygen contacts with Asn23, Ser27,

and Tyr43. Interactions also exist between Asn49 and Ser88 and the biotin carboxylic acid group. This network of hydrogen bonding in conjunction with hydrophobic interactions with four Trp residues (Trp 79, 92, 108, 120) and the tetrahydrothiophene result in high binding affinity.<sup>3</sup>

Biotin-(strept)avidin systems have been used for a variety of applications such as affinity isolation and purification, immunoassay, diagnostics, and localization.<sup>4–6</sup> Nearly any biological entity can be labeled with biotin including peptides, proteins, oligonucleotides, and antibodies. Antibody-(strept)avidin conjugates are used in the pretargeting approach to deliver radiolabeled biotin therapeutics and imaging agents.<sup>7</sup> Other applications include drug delivery<sup>8</sup> and material science.<sup>9</sup>

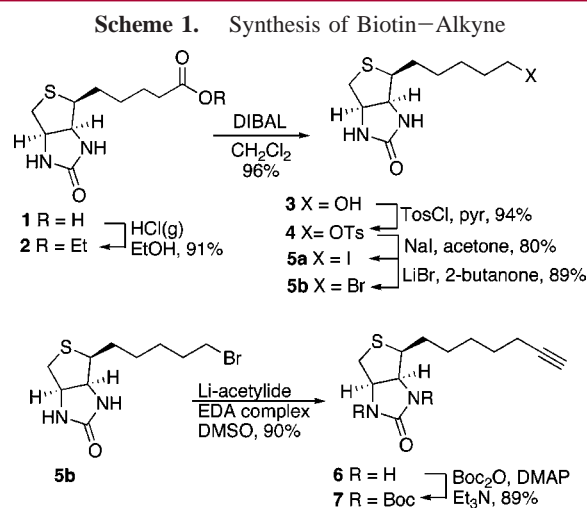
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A variety of biotin-labeling agents are commercially available.<sup>10</sup> Activated esters (*N*-hydroxysuccinimide ester and *p*-nitrophenyl ester), maleimide, and iodoacetyl derivatives are frequently used for coupling to substrates possessing amine or thiol groups.<sup>11</sup> In vivo applications of amide-linked biotin derivatives can be problematic because of cleavage by the endogenous enzyme biotinidase.<sup>12,13</sup>

Several structural modifications have been described to prevent biotinidase cleavage; however, these derivatives exhibit increased dissociation rates from (strept)avidin. Biotin amide derivatives containing a small  $\alpha$ -substituent have provided the most effective balance of biotinidase stability and (strept)avidin affinity.<sup>14</sup> Both *N*-methylation of the biotinamide linkage<sup>15</sup> and homologation of the valeric acid chain provided increased resistance to cleavage<sup>16</sup> as did replacement of the biotinamide connection with thiourea. Unfortunately, increased dissociation rates were observed with all of these linkage-modified derivatives.<sup>17</sup>

We have been interested in developing biotin conjugates of hydrophobic receptor ligands for affinity purification. Available biotin-coupling reagents are not suitable for derivatizing compounds that lack reactive functional groups such as amine, carboxylate, or thiol. The installation of polar substituents on a hydrophobic receptor substrate is possible but may decrease receptor-binding affinity because of unfavorable electrostatic interactions, modified solvation characteristics, different lipophilicities (as measured by log *P* octanol/water), and increased steric interactions.<sup>18</sup> As an alternative, we considered using nonpolar linkages synthesized via catalytic C–C bond-coupling methodology. Replacement of the biotin carboxamide with an alkyne or alkene connection would eliminate chemical or enzymatic hydrolysis. The success of this approach depends on the resulting C–C linked biotin substrates maintaining (strept)avidin affinity.

We elected to replace the carboxylic acid group with an alkyne that would allow entry into a wide variety of metal-catalyzed coupling procedures.<sup>19</sup> The synthesis was initiated by acid-catalyzed esterification of biotin. Selective reduction of the ester **2** was accomplished using diisobutylaluminum hydride (DIBAL) at  $-78^\circ\text{C}$ , affording alcohol **3** in 96% yield. Reaction of **3** with toluenesulfonyl chloride in pyridine at  $0^\circ\text{C}$  provided the sulfonate ester **4** in 94% yield (Scheme



1).<sup>20</sup> The iodide **5a** and bromide **5b** were prepared by halide substitution of **4**.

Low yields of the desired alkyne **6** were obtained from direct lithium acetylide substitution reactions of tosylate **4** or iodide **5a**. Fortunately, the alkyl bromide **5b** underwent efficient displacement with lithium acetylide–ethylenediamine in DMSO with careful temperature control  $\leq 15^\circ\text{C}$  and produced the desired alkyne **6** in high yield. This synthesis provided alkyne–biotin derivative **6** with a combined yield of 66% over five steps from biotin.

We evaluated avidin binding of **6** in solution. Competitive displacement of 2-(4'-hydroxyphenylazo)benzoic acid (HABA) with biotin derivatives provides a convenient spectroscopic method for assessing avidin binding. In solution, HABA forms a complex with the biotin binding site of avidin that is characterized by an absorbance band at 500 nm. Displacement of the HABA substrate by biotin results in decreased absorbance at 500 nm. This method has been widely used as a qualitative assay for evaluation of biotinylated substrates.<sup>21</sup>

We constructed a standardized biotin response curve for the HABA–avidin complex in 0.1 M phosphate buffer solution. The decrease in absorbance accompanying additions of  $5\ \mu\text{L}$  aliquots of *d*-biotin reference solution ( $5.0 \times 10^{-4}$  M in 0.10 M  $\text{NaH}_2\text{PO}_4$ ) was measured in triplicate and plotted against the concentration of added biotin. A response curve was generated analogously for alkyne **6** by addition to the standardized HABA–avidin solution. A decrease in absorbance was observed with the addition of **6** demonstrating HABA displacement. From the results of this assay, it can be concluded that the association constant of **6** remained high because HABA ( $K_a = 6 \times 10^6\ \text{M}^{-1}$ ) is displaced.<sup>21</sup> Although the HABA displacement was attenuated relative to biotin, effective binding of **6** was retained despite replacement of the carboxyl group with alkyne.

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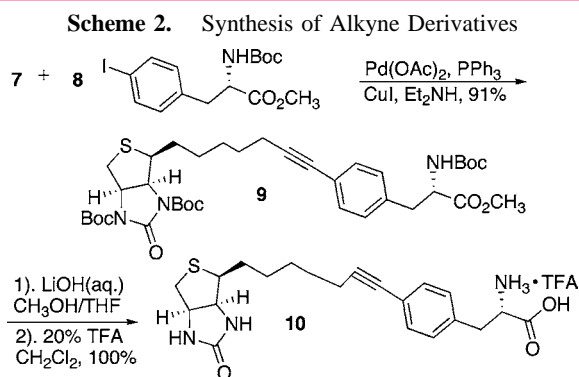
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Encouraged by the affinity exhibited by **6**, we proceeded to synthesize a phenylalanine biotin derivative. Amino acids labeled with biotin are important in peptide synthesis.<sup>8,22,23</sup> Sonogashira cross-coupling reactions have been used to produce alkynyl phenylalanine derivatives.<sup>24,25</sup> To increase the solubility of **6** in organic solvents, we protected the urea as the di-*tert*-butyl carbamate **7**. The 4-iodo-phenylalanine derivative **8** was coupled with alkyne **7** under standard conditions using catalytic Pd(OAc)<sub>2</sub>, PPh<sub>3</sub>, and CuI in diethylamine to give conjugate **9** in 91% yield (Scheme 2).

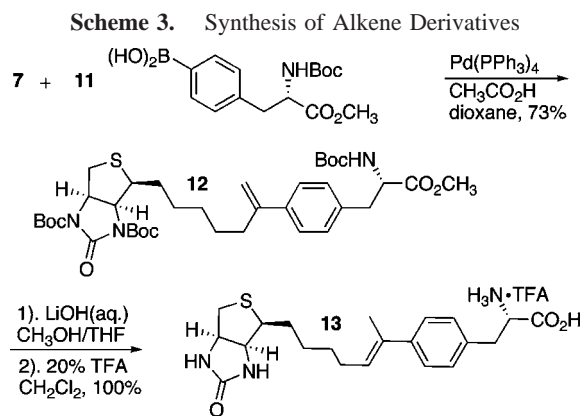


Hydrolysis of the ester with LiOH(aq) in MeOH/THF at 0 °C, followed by deprotection of the <sup>t</sup>Boc groups with trifluoroacetic acid (TFA) in CH<sub>2</sub>Cl<sub>2</sub> gave the alkyne-linked phenylalanine conjugate **10**. Spectroscopic characterization by NMR and HPLC-MS confirmed the product identity.

We wanted to investigate the effect of shortening the linkage between the biotin heterocycle and the phenylalanine on avidin affinity. Recently, Oh et al. reported a palladium-catalyzed insertion of arylboronic acids into terminal alkynes.<sup>26</sup> Following this procedure, alkyne **7** was coupled with protected 4-boronic acid derivative **11** to afford alkene **12** in 73% yield (Scheme 3). The alkene protons appeared as doublets at  $\delta$  5.21 and 5.01 ( $J = 1.3$  Hz) in the <sup>1</sup>H NMR spectrum, consistent with the expected alkene. No other alkene isomers were observed by <sup>1</sup>H NMR.

The ester **12** was hydrolyzed with LiOH, followed by <sup>t</sup>Boc deprotection with TFA/CH<sub>2</sub>Cl<sub>2</sub>. The product was characterized spectroscopically and by HPLC-MS. The exocyclic alkene isomerized to the more substituted (*E*)-alkene **13** during deprotection. The vinyl proton signal was observed as a triplet at  $\delta$  5.78 ( $J = 7.1$  Hz), and the allylic methyl group appeared as a singlet at  $\delta$  1.96.

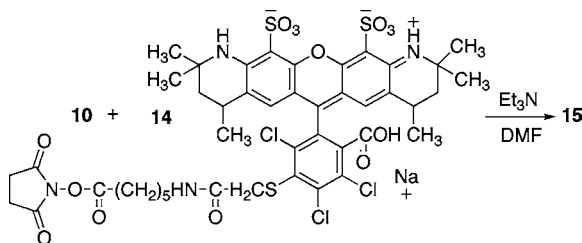
The avidin affinities of **10** and **13** were evaluated using the HABA assay as described for **6**. Both derivatives displaced HABA from avidin with comparable efficiency.



The sp<sup>2</sup>-hybridized carbon in alkene **13** corresponds to the position of the carboxyl in biotinamide derivatives and can be considered to be isosteric although not capable of hydrogen bonding. The linear alkyne connection in **10** is comparable to the overall length of homologated biotinamide conjugates. These minor differences in length and geometry did not significantly alter the observed binding affinity of **10** and **13**. These derivatives exhibited decreased affinity for avidin relative to biotin as expected from the deletion of the carboxyl group.

Solid-supported (strept)avidin sorbents have been widely used for batch affinity isolation and purification of biotinylated substrates. Nonbinding components are easily removed by elution, and isolation of biotinylated substrates is accomplished by dissociation from the supported (strept)-avidin.<sup>27</sup> The strong biotin–(strept)avidin interaction presents a problem in these applications because harsh conditions are required to dissociate the biotin from the support matrix. Biotin derivatives with reduced binding affinity such as desthiobiotin and 2-imunobiotin are advantageous because elution occurs with milder conditions. We prepared the fluorescent alkyne-linked phenylalanine **15** to evaluate the affinity for immobilized (strept)avidin (Scheme 4).

**Scheme 4. Synthesis of a Biotin Alexafluor Conjugate**



The  $\alpha$ -amino group of **10** was coupled with Alexafluor 546 succinimidyl ester **14** to give **15**. Compound **15** was dissolved in 0.1 M phosphate buffer, pH 7.2, at a concentra-

(27) Technical Document #25–0524, *Streptavidin Agarose: For Affinity Chromatography and Immunoprecipitation of Biotinylated Proteins*; Invitrogen: Eugene, OR, 2003.

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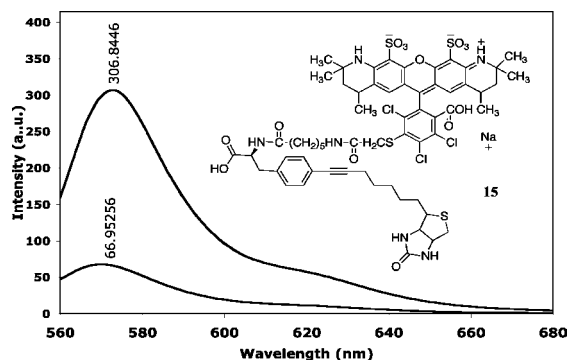
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tion of  $3.27 \times 10^{-6}$  M. A 3 mL aliquot (50% of the manufacturer's recommended loading) was added to a centrifuge tube containing (strept)avidin (5 mL) on agarose support. The tube was agitated for 15 min and centrifuged for 5 min, and the supernate containing unbound **15** was isolated. The tube was washed with 10 bed volumes of water to collect unbound substrate **15**. The concentration of free **15** was determined spectroscopically (Figure 1). The im-



**Figure 1.** Emission spectrum of **15**.

mobilized (strept)avidin retained 84% of **15**, confirming the potential for C-linked derivatives of **6** in batch isolation methods.

The avidin binding of **15** was studied by fluorescence titration. It was observed that **15** displayed higher emission intensity in the presence of avidin. Lo et al. have attributed the observed fluorescence enhancement to greater hydrophobicity within the binding pocket. A standardized solution of solubilized avidin was prepared and titrated with **15**. Fluorescence emission enhancement was monitored to determine the concentration required to saturate the receptor.<sup>28</sup> From this titration experiment, a dissociation constant

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( $K_d$ ) of ca.  $3.8 \times 10^{-9}$  M was determined for avidin-bound **15** using the method described by Srivastava et al.<sup>29</sup> This dissociation constant lies between the values of desthiobiotin ( $K_d = 5.0 \times 10^{-13}$  M)<sup>30</sup> and 2-imunobiotin ( $K_d = 8.0 \times 10^{-6}$  M)<sup>31</sup> and is also comparable to other recently reported biotin-fluorophore conjugates.<sup>28,32</sup>

This study evaluated the potential for carbon-linked biotin conjugates in affinity-based methods. Alkyne **6** was efficiently synthesized from *d*-biotin and exhibited strong binding in the HABA assay despite the deletion of the carboxylic acid. The 'Boc-protected alkyne **7** was effectively coupled using palladium-catalyzed reactions. Synthetic phenylalanine derivatives **10** and **13** and fluorescent probe **15** exhibited affinity for soluble and immobilized (strept)avidin, as expected for binding to the preserved 7-oxo-3-thia-6,8-diazabicyclo[3.3.0]oct-4-yl core. This approach enables conjugation to substrates that cannot be labeled with activated carboxylate derivatives of biotin. The covalent carbon linkage that replaces the carboxamide group of traditional biotin conjugates decreases relative (strept)avidin binding but retains sufficient affinity for bioanalytical applications and provides complete protection from chemical or enzymatic hydrolysis. This approach may be extended to other alkyne-coupling and cycloaddition procedures.

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**Supporting Information Available:** General methods, experimental details, copies of <sup>1</sup>H NMR and <sup>13</sup>C NMR spectra of compounds 1–3, 5–7, 9, 10, 12, 13, and HABA titration curves. This material is available free of charge via the Internet at <http://pubs.acs.org>.

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