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Buchwald Hartwig diversification of unprotected halotryptophans, halotryptophan containing tripeptides and the natural product barettin in aqueous conditions†

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Blending synthetic biology and synthetic chemistry represents a powerful approach to diversity complex molecules. To further enable this, compatible synthetic tools are needed. We report the first Buchwald Hartwig amination reactions with unprotected halotryptophans under aqueous conditions and demonstrate this methodology is applicable also to the modification of unprotected tripeptides and the natural product barettin.

Tryptophan is an essential amino acid, abundant within peptides and proteins, and central to their fluorescence, folding and structure;¹ it is also an important component in many bioactive natural products including non-ribosomal peptides *e.g.*, vancomycin, and a precursor to indole alkaloids *e.g.*, vinblastine and vincristine.² Modulation of tryptophan affords the potential to tune fluorescence, conformation and activity of these (bio)molecules.³ Although there has been significant research exploring the cross-coupling chemistry of nucleosides and the aromatic amino acid phenylalanine, both in their free state or as components of larger systems, the cross-coupling of halotryptophan has seemingly received little attention. Methodology now exists for the modification of halotryptophans, by Sonogashira⁴ and Suzuki-Miyaura^{5,6} but, to the best of our knowledge, the utilisation of Buchwald Hartwig amination (BHA) of halotryptophans and their derivatives remains unexplored.

Whilst BHA conditions have been applied to a wide range of aromatic species including indoles and 6-bromoquinolone (Scheme 1A and B),^{7,8} to the best of our knowledge there is no literature precedent for its use in unprotected biomolecule

functionalisation. Herein we report the application of aqueous BHA coupling to 5-bromoindole (Scheme 1C) and unprotected halotryptophans with a range of substituted aniline coupling partners (Scheme 1D). We then present the BHA derivatisation of tripeptides and the natural product barettin (Scheme 1E and F).

We set out to explore the feasibility of developing aqueous conditions for BHA coupling of halotryptophans, which may be readily accessed through a simple one-step biotransformation of indoles using tryptophan synthase⁹ or through a 4–5 step chemical synthesis.¹⁰ One challenge presented by free amino acids in palladium coupling reactions is the amino acid's ability to chelate and deactivate the palladium catalyst.¹¹ To reduce the likelihood of catalyst deactivation, we selected a precatalyst, L-Pd-G₁ (2) (Fig. 1) with the ^tBu-XPhos ligand, as shown by Buchwald and co-workers.¹² Initially, we examined the coupling of unprotected 5-bromoindole 3 and aniline 4.⁷ Investigating the impact of varying aqueous solvent mixture, catalyst loading, aniline eq. and base screening led us to conditions of 2 eq. of aniline, 2 mol% [Pd(^tBu-XPhos)G₁] catalyst, 1.2 eq. K₂CO₃, THF:water (1:1) at 65 °C for 16 h (Scheme 2) giving an 85% yield of 5.

Translating these conditions directly to the more challenging cross-coupling of 5-bromotryptophan, we saw a substantial drop in yield. This prompted a further round of optimisation and a move to microwave heating resulting in 56% isolated yield of the cross-coupled product 7 (Scheme 2).

Next, we decided to take a closer look at the influence of reaction time under the microwave conditions exploring the impact of heating by microwave from 1 minute to 1.5 h (Table 1).

We found that shorter reaction times gave improved yields, optimum at 8 min (75% yield) (Table 1). It is suspected that longer reaction times cause degradation of the product resulting in lower yields, no clear by-products could be identified. Having established utilisable conditions for the BHA of 5-bromotryptophan, we investigated the reaction with tryptophans substituted with bromo and chloro in varying positions (Table 2). The reaction proceeded well for 5, 6 and 7-bromotryptophans resulting in yields of 75, 51 and 68% respectively. Similarly, the 5, 6 and

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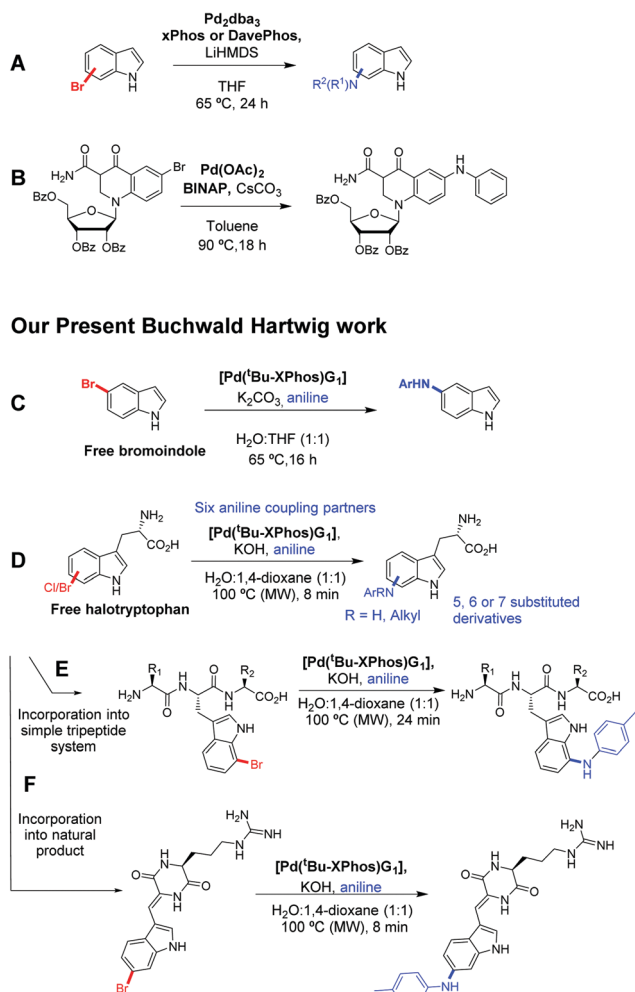
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Related Buchwald Hartwig chemistries



Scheme 1 Buchwald Hartwig related work: examples of relevant prior work are outlined along with work within this study. (A) Example literature conditions for BHA on a bromoindole from Buchwald.⁷ (B) Buchwald Hartwig coupling for a 6-bromoquinolone nucleoside, Wicke *et al.*⁸ (C) Reported BHA modification for 5-bromoindole. (D) Reported BHA modifications for a range of halotryptophans. (E) Reported BHA modification of a bromotryptophan containing tripeptide. (F) Reported BHA modification the natural product baretin.

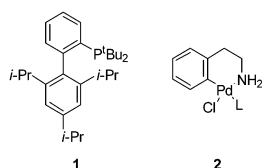
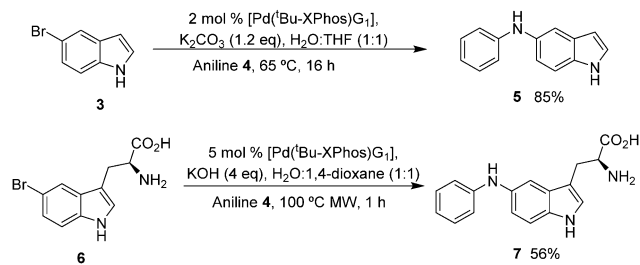


Fig. 1 Buchwald ligand ^tBu-XPhos **1** and precatalyst L-Pd-G1 **2**.

7-chlorotryptophans also worked well (67, 60 and 31% yields respectively).

Having assessed reactivity of the various halotryptophans with aniline, we next sought to examine the effects of steric and electronic properties of the aniline coupling partner. We selected six anilines, **10** to **15**, (Scheme 3 and Table 3) and



Scheme 2 Buchwald Hartwig Amination reaction of 5-bromoindole (**3**) and 5-bromotryptophan (**6**) with aniline (**4**).

Table 1 Assessment of microwave reaction time on yield

Time of reaction (min)	Yield (%)
1	43 ^a
5	53 ^a
8	75 ^a
15	71 ^a
30	71 ^a
60	56 ^b
90	35 ^b

Conditions: 5-bromotryptophan (1 eq.), [Pd(^tBu-XPhos)G₁] (5 mol%), KOH (4.0 eq.), aniline (2.0 eq.), H₂O/1,4-dioxane (1 : 1, 0.6 mL), microwave heating, sealed tube, under argon, 100 °C. ^a Isolated yield of the product after reverse phase purification. ^b Yield assessed by ¹H-NMR using 1,3,5-tri-*tert*-butylbenzene as an internal standard.

Table 2 Reaction of a series of halotryptophans with aniline

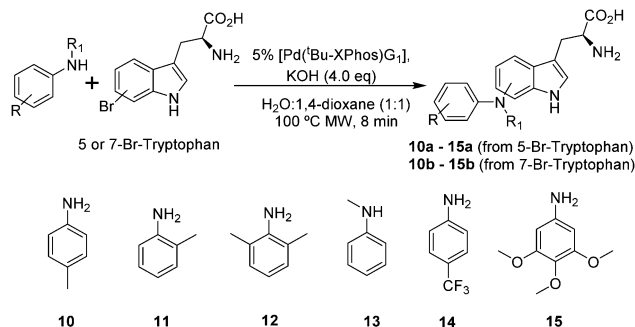
Entry	Tryptophan	Product	Isolated yield/%
1	5-Br	7	75
2	6-Br	8	51
3	7-Br	9	68
4	5-Cl	7	67
5	6-Cl	8	60
6	7-Cl	9	31

Conditions: halotryptophan (1 eq.), [Pd(^tBu-XPhos)G₁] (5 mol%), KOH (4.0 eq.), aniline (2.0 eq.), H₂O/1,4-dioxane (1 : 1, 0.6 mL), microwave heating, sealed tube, under argon, 8 min, 100 °C.

reacted these with 5- and 7-bromotryptophan; the two most reactive regioisomers according to our positional scoping.

These results show that cross-coupling proceeded even in the presence of moderate steric hindrance: 2-methylaniline (**11**) reacted with 5- and 7-bromotryptophans nearly as efficiently as unsubstituted aniline, with yields of 63% and 62% respectively. However, it is striking that two methyl groups in close proximity to the aryl amine, as in 2,6-dimethylaniline (**12**), led to complete suppression of the reaction and recovery of starting materials. Interestingly, *N*-methylaniline (**13**) is partially tolerated yielding 4% product for the 7-bromotryptophan, but no product could be obtained for the 5-bromotryptophan.





Scheme 3 Investigating the reaction of 5-bromo and 7-bromotryptophans with a selection of anilines.

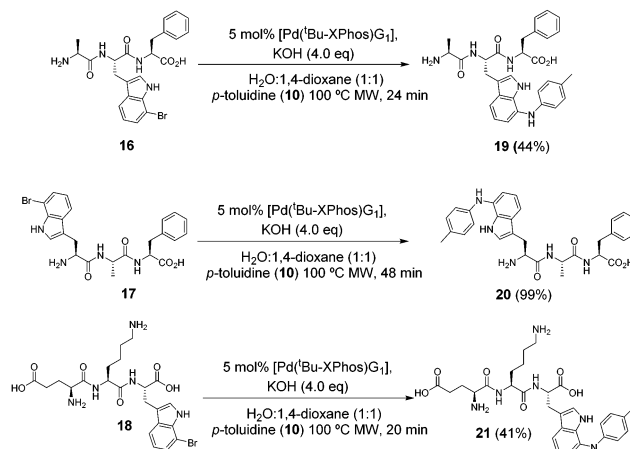
Table 3 Investigating the reactivity of 5-bromo and 7-bromotryptophans with a selection of anilines

Entry	Tryptophan	Product	Isolated yield/%
1	5-Br	10a	59
2	7-Br	10b	53
3	5-Br	11a	63
4	7-Br	11b	62
5	5-Br	12a	No reaction
6	7-Br	12b	No reaction
7	5-Br	13a	No reaction
8	7-Br	13b	4
9	5-Br	14a	7
10	7-Br	14b	15
11	5-Br	15a	67
12	7-Br	15b	59

Conditions: bromotryptophan (1 eq.), [Pd(Bu-XPhos)G₁] (5 mol%), KOH (4.0 eq.), anilines (2.0 eq.), H₂O/1,4-dioxane (1:1, 0.6 mL), microwave heating, sealed tube, under argon, 8 min, 100 °C.

Differently substituted anilines, bearing electron-withdrawing or electron-donating groups, were subsequently tested to assess the impact of the electronic properties of the nucleophile in our reaction. With the weakly electron donating *para*-methyl substituent, we saw a slight drop in yield from 75% and 68% to 59% and 53% respectively for the 5- and 7-bromotryptophans. It was predicted¹³ that the strong electron withdrawing nature of the CF₃ substituent would reduce the reactivity of the aniline towards amination and this was borne out in the very low isolated yields (7 and 15%, Table 3, entries 9 and 10). Also, as anticipated, the electron rich trimethoxyaniline reacted well (67 and 59% yields Table 3, entries 11 and 12), but did not surpass the yields observed for the unsubstituted anilines reactions. The slightly lower than expected yield for **15** may be due to improved organic solubility of the products resulting in a slight loss of product during the work up.

We next tested the reaction with more complex halotryptophan containing substrates. Tripeptides **16**, **17** and **18** were synthesised (see ESI[†]) and were subjected to our BHA conditions (Scheme 4) using *p*-toluidine (**10**) as the coupling partner and increasing the reaction times to allow for potentially slower reaction with the more complex molecules. Tripeptide **17** reacted well and a 99% isolated yield was achieved. The products of tripeptides **16** and **18** suffered losses during purification. In particular peptide **21** had poor stability in solution and required careful handling to isolate

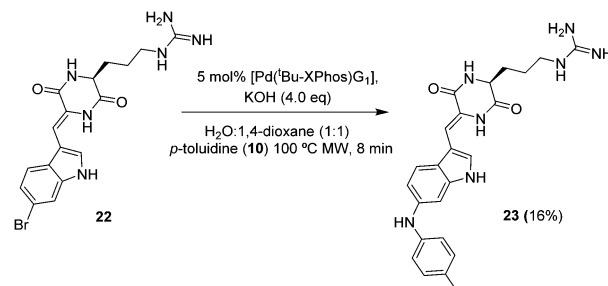


Scheme 4 Investigating the reaction of 7-bromotryptophan containing tripeptides with *p*-toluidine (**10**).

high purity material. These examples demonstrate the successful BHA coupling of halotryptophan present at either terminal or middle locations in an unprotected peptide sequence possessing polar/non-polar side chains.

Encouraged by these results, we next chose derivatisation of baretin a marine natural product reputed to have antioxidant and anti-inflammatory properties and containing a 6-bromotryptophan group. Using our standard conditions (Scheme 5) we observed full consumption of starting material. LCMS showed a distinct peak at 4.3 min with an [M + H]⁺ of 446 corresponding to the desired product. Despite the small scale and purification difficulties, we were able to isolate 1 mg of the product **23** following HPLC purification, (purity > 95%, 16% yield).

We have for the first time demonstrated the use of Buchwald Hartwig Amination to diversify free unprotected 5, 6 and 7-bromo- and chlorotryptophans, and shown the reaction is also effective on short 7-bromotryptophan containing peptides including potentially reactive groups such as glutamic acid and lysine. The reaction products can be isolated by reverse-phase chromatography or HPLC. As seen in our previous work,^{6b} following the cross-coupling reaction, the tripeptides were single peaks by HPLC(LCMS) (see ESI[†]) indicating presence of single diastereomers and therefore no observable epimerisation occurred during the cross-coupling reactions. Our new conditions were also successfully applied to the more complex



Scheme 5 BHA reaction of baretin with *p*-toluidine (**10**).



natural product baretin. This provides an expansion of the reactions that can be carried out with easily accessed⁹ halotryptophans that can be incorporated into more complex peptides and natural products. The incorporation of the halotryptophans and wide range of amine coupling partners could be envisioned to produce a wide variety of natural product analogues for use in structure–activity relationship analyses.

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Conflicts of interest

There are no conflicts to declare.

Notes and references

- (a) G. D. Rose, A. R. Geselowitz, G. J. Lesser, R. H. Lee and M. H. Zehfus, *Science*, 1985, **229**, 834–838; (b) D. A. Dougherty, *Science*, 1996, **271**, 163–168.
- R. L. Noble, C. T. Beer and J. H. Cutts, *Ann. N. Y. Acad. Sci.*, 1958, **76**, 882–894.
- (a) R. J. M. Goss, S. Shankar and A. A. Fayad, *Nat. Prod. Rep.*, 2012, **29**, 870–889; (b) K. P. P. Mahoney, D. R. M. Smith, E. J. A. Bogosyan and R. J. M. Goss, *Synthesis*, 2014, 2122–2132; (c) A. E. Osbourn, R. J. M. Goss and G. T. Carter, *Natural Products: Discourse, Diversity, and Design*, Wiley-Blackwell, 2014, ISBN: 978-1-118-29806-0; (d) A. Kirschning, F. Taft and T. Knobloch, *Org. Biomol. Chem.*, 2007, **5**, 3245–3259.
- M. J. Corr, S. V. Sharma, C. Pubill-Ulldemolins, R. T. Bown, P. Poirrot, D. R. M. Smith, C. Cartmell, A. Abou Fayad and R. J. M. Goss, *Chem. Sci.*, 2017, **8**, 2039–2046.
- A. Deb Roy, R. J. M. Goss, G. K. Wagner and M. Winn, *Chem. Commun.*, 2008, 4831–4833.
- (a) A. Deb Roy, S. Gruschow, N. Cairns and R. J. M. Goss, *J. Am. Chem. Soc.*, 2010, **132**, 12243–12245; (b) S. V. Sharma, X. Tong, C. Pubill-Ulldemolins, C. Cartmell, E. J. A. Bogosyan, E. J. Rackham, E. Marelli, R. B. Hamed and R. J. M. Goss, *Nat. Commun.*, 2017, **8**, 229–238.
- M. D. Charles, P. Schultz and S. L. Buchwald, *Org. Lett.*, 2005, **7**, 3965–3968.
- L. Wicke, J. W. Engels, R. Gambari and A. M. Saab, *Arch. Pharm. Chem. Life Sci.*, 2013, **346**, 757–765.
- (a) D. R. M. Smith, T. Willemse, D. S. Gkotsi, W. Schepens, B. U. W. Maes, S. Ballet and R. J. M. Goss, *Org. Lett.*, 2014, **16**, 2622–2625; (b) R. J. M. Goss and P. L. A. Newill, *Chem. Commun.*, 2006, 4924–4925.
- For example, see: (a) Y. Konda-Yamada, C. Okada, K. Yoshida, U. Yasuyuki, S. Arima, N. Sato, T. Kai, H. Takayanagi and Y. Harigaya, *Tetrahedron*, 2002, **58**, 7851–7861; (b) C. Ma, X. Liu, X. Li, J. Flippen-Anderson, S. Yu and J. Cook, *J. Org. Chem.*, 2001, **66**, 4525–4542.
- M. A. Carvalho, B. C. Souza, R. E. F. Paiva, F. R. G. Bergamini, A. F. Gomes, F. C. Gozzo, W. R. Lustrì, A. L. B. Formiga, G. Rigatto and P. P. Corbi, *J. Coord. Chem.*, 2012, **65**, 1700–1711.
- N. C. Bruno, M. T. Tudge and S. L. Buchwald, *Chem. Sci.*, 2013, **4**, 916–920.
- K. H. Hoi, S. Calimsiz, R. D. J. Froese, A. C. Hopkinson and M. G. Organ, *Chem. – Eur. J.*, 2012, **18**, 145–151.

