Solid-Phase Synthesis of Biocompatible N-Heterocyclic Carbene–Pd Catalysts Using a Sub-monomer Approach

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Abstract
Taking inspiration from the assembly of so-called peptoids (N-alkylglycine oligomers) we present a new synthetic methodology whereby N-heterocyclic carbene (NHC) based Pd ligands were assembled using a sub-monomer approach and loaded with Pd via solid-phase synthesis. This allowed the rapid generation a library of NHC–palladium catalysts that were readily functionalised to allow bioconjugation. These catalysts were able to rapidly activate a caged fluorophore and ‘switch-on’ an anticancer prodrug in 3D cell culture.

Introduction
Bioorthogonal reactions enable the selective visualisation and manipulation of biological processes in living systems and have been widely used in a number of applications.1–4 Transition metal mediated bioorthogonal reactions are of particular interest as they enable an array of non-natural chemical transformations that can be used to modulate living systems.5, 6 Reactions mediated by copper,7, 8 iron,9 gold,10–12 ruthenium,12–16 and iridium17 have all found applications in living systems, although palladium is perhaps the most utilised metal in a biological setting.18 Palladium has gained popularity in bioorthogonal chemistry due its ability to perform catalytic cross-coupling reactions, enabling the generation of carbon–carbon and carbon–heteroatom bonds under mild, biological conditions,19–21 and more recently in vivo.22,23 Palladium catalysts have been used to initiate a range of intracellular reactions including dealkylation,24 decaging of propargyloxycarbonyl groups,18,24,25 as well as Suzuki–Miyaura cross-couplings.20,26–28 Thus, palladium mediated reactions have been used to selectively activate enzymes through deprotection of modified amino acids within proteins,21, 29 to synthesise anticancer agents in cellulo from two benign precursors,19 as well as activate produgs;20–22,24 however, to date, the majority of the examples have used palladium nanoparticles entrapped within a polymeric support,19,20,22–24,26,27,30 simple palladium salts such as Pd(OAc)2,31–33 or designed targeted palladium ligands.34 This is in contrast to the highly active palladium catalysts used in conventional organic synthesis that use a variety of stabilising ligands (e.g. phosphines) or N-heterocyclic carbenes (NHC),35, 36 but few have been used in a biological setting. Chen used an NHC–Pd catalyst (with imidazolium-based ligands bearing hydrophilic quaternary ammonium salts) to mediate the Suzuki–Miyaura coupling reaction between a boronic acid functionalised biotin and 4-iodophenylalanine modified cell surface proteins, enabling subsequent imaging of the cell surface with fluorescently labelled streptavidin.33 Recently, we reported a water soluble NHC–Pd catalyst coupled to a cell penetrating peptide, which was able to remove a propargyloxycarbonyl group from a pro-fluorophore in cells, thus demonstrating the first intracellular application of NHC–Pd chemistry.37 While catalyst loading was
carried out on the solid-phase, the synthesis of the NHC ligand required multistep synthesis in solution, purification, and conjugation to a solid support.

Here, we report an efficient microwave assisted solid-phase synthesis of a series of biocompatible NHC–Pd catalysts and their chemistry in cells. To the best of our knowledge, a solid phase approach to the generation of NHC–Pd catalysts has not been reported and, with our approach, these stable NHC–Pd catalysts were prepared in good yields with the majority of the catalysts showing good activity in a biological setting.

**Results and discussion**

A solid-phase synthetic route was developed for the NHC–Pd catalysts by adopting the ‘sub-monomer’ approach, originally developed by Zuckermann for the synthesis of peptoids (N-alkyl glycine oligomers), with sequential acylation and alkylation reactions on a resin.\(^{38,39}\) The NHC ligand chosen here was first reported by Meldal who showed that a resin-bound NHC–Pd complex (conjugated to a hydrophobic dipeptide) showed good catalytic efficiency in Suzuki–Miyaura cross-couplings in water.\(^{40}\) Here, this pyridine-derivatised imidazolium ligand was attached to a range of different amino acid spacers, with the aim of providing biocompatibility, aqueous stability, as well as a handle for biomolecule conjugation.

**Solid-phase synthetic route for the NHC–Pd catalysts**

Fmoc-Ahx-OH was coupled to a 2-chlorotrityl chloride linker on a polystyrene resin (mesh 100–200), with subsequent removal of the Fmoc group giving 1, which was acylated with 2-bromoacetic acid (2 M in DMF) using DIC as a coupling reagent at 60 °C under \(\mu\)w irradiation for 20 min (Scheme 1). The bromide in 2 was substituted by imidazole to give 3, with optimised reaction conditions (ESI, Table S1) allowing quantitative N-alkylation via an excess of imidazole (2 M) at 60 °C (\(\mu\)w heating for 40 min) in anhydrous DMSO with 0.5 M AgNO\(_3\). The addition of AgNO\(_3\) was required for full conversion (based on HPLC analysis, ESI Figure S1). The imidazole in 3 was N-alkylated with 2-(bromomethyl)pyridine (1 M) to give 4 with >95% conversion using Et\(_3\)N (1 M) and AgNO\(_3\) (0.5 M) at 60 °C in anhydrous DMF (for optimisation see ESI Table S2) giving the NHC ligand in >95% purity and 65% overall yield over 3 steps (ESI, Figure S2 and S3). Palladium loading of the NHC-ligand 4 on solid-phase was carried out as reported by Meldal\(^{40}\) with minor modifications. In brief, the carbene was generated on the resin from the imidazolium ion with the phosphazene base 2-tert-butylimino-2-diethylamino-1,3-dimethylperhydro-1,3,2-diazaphosphorine (BEMP) in anhydrous DMF. Pd(COD)Cl\(_2\) was added subsequently to generate the NHC–Pd catalyst 5 (Scheme 1), which was cleaved off the resin with 30% HFIP in DCM and purified by semi-preparative RP-HPLC. This allowed the rapid and efficient generation of >100 mg quantities of the catalyst 5 that could be freely stored.\(^{\S}\)
Scheme 1. The microwave assisted solid-phase synthesis of NHC–Pd catalyst 5. a) 0.3 M Fmoc–Ala-OH, 0.5 M DIPEA, anhydrous DCM–DMF (9:1), 1 h; b) 20% piperidine in DMF, 2 × 10 min; c) 2 M BrCH\(_2\)CO\(_2\)H, 1 M DIC, anhydrous DMF, 20 min, 60 °C, µw; d) 2 M imidazole, 0.5 M AgNO\(_3\) in anhydrous DMSO, 40 min, 60 °C, µw; e) 2 M 2-(bromomethyl)pyridine, 1 M Et\(_3\)N, 0.5 M AgNO\(_3\), anhydrous DMF, 90 min, 60 °C, µw; f) BEMP, anhydrous DMF, N\(_2\), 45 min, then Pd(COD)Cl\(_2\) overnight; g) 30% HFIP–DCM, 1 h. All conversions were monitored by cleavage of a small sample from the resin and characterisation by HPLC and NMR. As drawn, the catalyst is Pd(II) with Y most likely formate (from the HPLC purification buffer).

**Synthesis of NHC–Pd catalyst library**

This solid-phase synthesis route was used to generate an NHC–Pd catalyst library (5–12). Using the NHC moiety with different amino acid spacers gave a range of both hydrophilic and hydrophobic groups to compare the potential effect on catalytic activity, as well as the robustness of the synthetic method (Figure 1). The catalysts were synthesised using the 2-chlorotrityl linker, but the methodology was also compatible with the Rink-amide linker (ESI, catalyst 13). The ligands for catalysts 5–12 were fully characterised using NMR, HPLC, and HRMS, and after palladium loading, the catalysts were cleaved off the resin, purified by semi-preparative RP-HPLC, and characterised by ESI-MS and HPLC (ESI, Table S3). The purity of the catalysts was confirmed by analytical HPLC (ESI, Figure S4 and S5 – the “naked ligands” and the Pd-loaded ligands displayed different retention times) and the presence of palladium also established by HRMS (ESI, Figure S6) (the presence of the Pd meant that the \(^1\)H NMR spectra were highly broadened). These catalysts were stable for two weeks at room temperature in ACN/H\(_2\)O, and \(\geq\) 8 weeks at 4 °C.
Screening of catalytic activity

The fluorogenic probe based on 2,7-dichlorofluorescein (DCF) was used to evaluate the activity of the catalysts. The profluorophore O-propargylated DCF-1 was converted to the fluorescent molecule DCF-2 ($\lambda_{Ex/Em}$ 480/520 nm) upon Pd-catalysed cleavage of the propargyl group (Figure 2A). Catalyst screening was carried out at 37 °C in PBS and in MCF-7 cell lysate to evaluate catalytic activity under biologically relevant conditions. Catalysts 5–12 (0.8 mol%) were incubated with DCF-1 (10 μM) and the increase in fluorescence measured over 4 h with an additional measurement at 20 h (ESI, Figure S7). With the exception of 7 and 11, all the catalysts were active in PBS with 8 and 9 showing comparable activity to 0.8 mol% Pd(OAc)$_2$ (Figure 2B). The decaging reactions were notably slower in cell lysate with 8 and 9 again showing the best catalytic efficiency under these conditions and outperforming Pd(OAc)$_2$ (ESI, Figure S8A). Although shorter hydrophobic spacers seemed to be preferred in PBS (8 > 9 > 10), clear structure–activity relationships could not be established for the catalytic activity, particularly in the cell lysate, highlighting the need for methods to rapidly generate combinatorial catalyst libraries.

Catalyst 8 was further evaluated by monitoring the decaging of DCF-1 (50 μM) by HPLC, with the experiments performed in PBS and in human plasma to mirror a cellular environment rich in proteins. In PBS, 8 (2 mol%) gave 92% conversion to DCF-2 after 5 h (Figure 2C), whereas in plasma 97% conversion was observed (ESI, Figure S8B).
Prodrug activation in cancer cells and in cancer cell spheroids

The catalyst 8 was evaluated for its efficiency to activate the caged anticancer drug N-propargyl protected 5-fluorouracil\textsuperscript{24} (Pro-5-FU) (Figure 3A). In PBS, 5 mol\% of 8 was able to convert Pro-5-FU (100 µM) into the active drug 5-fluorouracil (5-FU) within 48 h, as monitored by HPLC (Figure 3B). The co-treatment of MCF-7 cell first with Pro-5-FU (100 µM) for 24 h followed by catalyst 8 (10 mol\%) for 4 days resulted in comparable cytotoxicity (MTT assay) to 5-FU (Figure 3C), with the drug activation likely taking place both intra- and extracellularly (Pro-5-FU is readily taken up by cells\textsuperscript{30}). Catalyst 8 did not show notable cytotoxicity with 84% cell viability at 10 µM concentration.

Next, MCF-7 spheroids were treated with the prodrug Pro-5-FU (100 µM) and catalyst 8 (10 mol\%) for 5 days, stained with the LIVE/DEAD™ Cell Imaging Kit (488/570), and analysed by fluorescence microscopy. Spheroids treated with both the prodrug and catalyst resulted in cell death comparable to cells treated with 5-FU, whereas treatment with only the catalyst or the prodrug Pro-5-FU had no effect on the viability of the 3D spheroids (Figure 3D). These results demonstrate that the catalyst was able to decage a protected prodrug in a more representative 3D cancer model, which may have applications in future anticancer prodrug therapies.
Conclusions

A highly efficient microwave assisted solid-phase synthesis of NHC–Pd catalysts, based on the ‘sub monomer’ approach was developed and used to generate biocompatible catalysts on scale. Catalyst 8, comprising of the NHC–Pd moiety linked to a glycine, was the most robust catalyst in the series and was able to activate a fluorogenic probe in a biological setting and decage the protected anticancer drug (5-fluoro-1-propargyluracil) in a 3D cancer cell culture resulting in comparable cell death to 5-fluorouracil. These robust NHC-Pd catalysts have a carboxylic acid that can be readily converted to a stable active ester thus providing a handle for bioconjugation and offering applications for both specific cell targeting ligands and bioorthogonal prodrug activations.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

We thank the European commission for the Marie Skłodowska-Curie Individual Fellowship for D.C. (MSCA-IF-2014-EF, MetalCell 659489), and the Engineering and Physical Sciences Research Council and Medical Research Council CDT Optima (grant number EP/L016559/1) and the Rosetrees Trust for funding. We also thank the SIRCAMS at University of Edinburgh for the HRMS analysis and the School of Chemistry NMR facility.
The solid-phase approach used here offers advantages over more traditional solution-phase methods, namely the use of mass-action to drive the chemistries and the removal of intermediary purification steps. The ligand was previously synthesised in three steps (in solution) with a combined reaction time of 7.5 h (compared to 3 h in this work) requiring both silica column chromatography and preparative RP-HPLC purification. \(^{40}\)

1. J. Li and P. R. Chen, *Nat Chem Biol*, 2016, **12**, 129-137.
2. J. G. Rebelein and T. R. Ward, *Curr Opin Biotechnol*, 2018, **53**, 106-114.
3. E. M. Sletten and C. R. Bertozzi, *Angew Chem Int Ed Engl*, 2009, **48**, 6974-6998.
4. J. J. Soldevila-Barreda and N. Metzler-Nolte, *Chem Rev*, 2019, **119**, 829-869.
5. M. Martinez-Calvo and J. L. Mascarenas, *Chimia*, 2018, **72**, 791-801.
6. Y. Bai, J. Chen and S. C. Zimmerman, *Chem Soc Rev*, 2018, **47**, 1811-1821.
7. J. Clavadetscher, S. Hoffmann, A. Lilienkampf, L. Mackay, R. M. Yusop, S. A. Rider, J. J. Mullins and M. Bradley, *Angew Chem Int Ed Engl*, 2016, **55**, 15662-15666.
8. Y. Bai, X. Feng, H. Xing, Y. Xu, B. K. Kim, N. Baig, T. Zhou, A. A. Gewirth, Y. Lu, E. Oldfield and S. C. Zimmerman, *J of the Am Chem Soc*, 2016, **138**, 11077-11080.
9. P. K. Sasmal, S. Carregal-Romero, A. A. Han, C. N. Streu, Z. Lin, K. Namikawa, S. L. Elliott, R. W. Koster, W. J. Parak and E. Meggers, *Chembiochem*, 2012, **13**, 1116-1120.
10. K. Tsubokura, K. H. Vong, A. R. Pradipta, A. Ogura, S. Urano, T. Tahara, S. Nozaki, H. Onoe, Y. Nakao, R. Sibgatullina, A. Kurbangalieva, Y. Watanabe and K. Tanaka, *Angew Chem Int Edit*, 2017, **56**, 3579-3584.
11. A. M. Perez-Lopez, B. Rubio-Ruiz, V. Sebastian, L. Hamilton, C. Adam, T. L. Bray, S. Irusta, P. M. Brennan, G. C. Lloyd-Jones, D. Sieger, J. Santamaria and A. Unciti-Broceta, *Angew Chem Int Edit*, 2017, **56**, 12548-12552.
12. C. Vidal, M. Tomas-Gamasa, P. Destito, F. Lopez and J. L. Mascarenas, *Nat Commun*, 2018, **9**, 1913.
13. C. Streu and E. Meggers, *Angew Chem Int Ed Engl*, 2006, **45**, 5645-5648.
14. M. Tomas-Gamasa, M. Martinez-Calvo, J. R. Couceiro and J. L. Mascarenas, *Nat Commun*, 2016, **7**, 12538.
15. T. Volker, F. Dempwolff, P. L. Graumann and E. Meggers, *Angew Chem Int Ed Engl*, 2014, **53**, 10536-10540.
16. M. I. Sanchez, C. Penas, M. E. Vazquez and J. L. Mascarenas, *Chem Sci*, 2014, **5**, 1901-1907.
17. S. Bose, A. H. Ngo and L. H. Do, *J Am Chem Soc*, 2017, **139**, 8792-8795.
18. S. V. Chankeshwara, E. Indrigo and M. Bradley, *Curr Opin Chem Biol*, 2014, **21**, 128-135.
19. J. Clavadetscher, E. Indrigo, S. V. Chankeshwara, A. Lilienkampf and M. Bradley, *Angew Chem Int Edit*, 2017, **56**, 6864-6868.
20. R. M. Yusop, A. Unciti-Broceta, E. M. Johansson, R. M. Sanchez-Martin and M. Bradley, *Nat Chem*, 2011, **3**, 239-243.
21. J. Li, J. Yu, J. Zhao, J. Wang, S. Zheng, S. Lin, L. Chen, M. Yang, S. Jia, X. Zhang and P. R. Chen, *Nat Chem*, 2014, **6**, 352-361.
22. M. A. Miller, B. Askevold, H. Mikula, R. H. Kohler, D. Pirovich and R. Weissleder, Nat Commun, 2017, 8, 15906.
23. T. L. Bray, M. Salji, A. Brombin, A. M. Perez-Lopez, B. Rubio-Ruiz, L. C. A. Galbraith, E. E. Patton, H. Y. Leung and A. Unciti-Broceta, Chem Sci, 2018, 9, 7354-7361.
24. J. T. Weiss, J. C. Dawson, K. G. Macleod, W. Rybski, C. Fraser, C. Torres-Sanchez, E. E. Patton, M. Bradley, N. O. Carragher and A. Unciti-Broceta, Nat Commun, 2014, 5, 3277.
25. J. T. Weiss, J. C. Dawson, C. Fraser, W. Rybski, C. Torres-Sanchez, M. Bradley, E. E. Patton, N. O. Carragher and A. Unciti-Broceta, J Med Chem, 2014, 57, 5395-5404.
26. E. Indrigo, J. Clavadetscher, S. V. Chankeshwara, A. Lilienkampf and M. Bradley, Chem Commun, 2016, 52, 14212-14214.
27. F. Wang, Y. Zhang, Z. Du, J. Ren and X. Qu, Nat Commun, 2018, 9, 1209.
28. P. Destito, A. Sousa-Castillo, J. R. Couceiro, F. Lopez, M. A. Correa-Duarte and J. L. Mascarenas, Chem Sci, 2019, 10, 2598-2603.
29. J. Wang, S. Zheng, Y. Liu, Z. Zhang, Z. Lin, J. Li, G. Zhang, X. Wang, J. Li and P. R. Chen, J Am Chem Soc, 2016, 138, 15118-15121.
30. G. Y. Tonga, Y. D. Jeong, B. Duncan, T. Mizuhara, R. Mout, R. Das, S. T. Kim, Y. C. Yeh, B. Yan, S. Hou and V. M. Rotello, Nat Chem, 2015, 7, 597-603.
31. J. Li, S. Lin, J. Wang, S. Jia, M. Yang, Z. Hao, X. Zhang and P. R. Chen, J Am Chem Soc, 2013, 135, 7330-7338.
32. C. D. Spicer and B. G. Davis, Chem Commun, 2011, 47, 1698-1700.
33. X. J. Ma, H. X. Wang and W. Z. Chen, J Org Chem, 2014, 79, 8652-8658.
34. M. Martinez-Calvo, J. R. Couceiro, P. Destito, J. Rodriguez, J. Mosquera and J. L. Mascarenas, ACS Catal, 2018, 8, 6055-6061.
35. G. C. Fortman and S. P. Nolan, Chem Soc Rev, 2011, 40, 5151-5169.
36. J. F. Jensen, K. Worm-Leonhard and M. Meldal, Eur J Org Chem, 2008, 3785-3797.
37. E. Indrigo, J. Clavadetscher, S. V. Chankeshwara, A. Megia-Fernandez, A. Lilienkampf and M. Bradley, Chem Commun, 2017, 53, 6712-6715.
38. R. N. Zuckermann, J. M. Kerr, S. B. H. Kent and W. H. Moos, J Am Chem Soc, 1992, 114, 10646-10647.
39. T. S. Burkoth, A. T. Fafarman, D. H. Charych, M. D. Connolly and R. N. Zuckermann, J Am Chem Soc, 2003, 125, 8841-8845.
40. K. Worm-Leonhard and M. Meldal, Eur J Org Chem, 2008, 5244-5253.
41. M. Santra, S. K. Ko, I. Shin and K. H. Ahn, Chem Commun, 2010, 46, 3964-3966.