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A novel amide synthesis methodology is described using amines, CO_2 and Grignard reagents and Mitsunobu reagents. The method was applied to carbon-11 radiochemistry to label amides using cyclotron-produced [¹¹C]CO₂. The synthetic utility of the one-pot labelling methodology was demonstrated by producing [¹¹C]melatonin. The incorporation of [¹¹C]CO₂ into [¹¹C]melatonin was 36% – determined by radioHPLC 2 min post [¹¹C]CO₂ delivery.

Positron emission tomography (PET) is a non-invasive imaging technology used for *in vivo* molecular imaging.¹ Carbon-11 (¹¹C) is a short-lived positron-emitting radionuclide widely used in labelling radiopharmaceuticals for medical research, diagnosis, staging and monitoring treatments. Its physical and nuclear characteristics (radioactive half-life = 20.4 min) and its orthology with carbon-12 make ¹¹C an attractive radionuclide for labelling and molecular imaging. The development of rapid and reliable chemical methods for incorporating carbon-11 into organic molecules is required to expand the repertoire of available radiotracers for *in vivo* imaging studies. Cyclotron-produced [¹¹C]CO₂ is obtained by the proton bombardment of nitrogen-14 gas via the $^{14}N(p,\alpha)^{11}C$ nuclear reaction. Because of its low reactivity and solubility in organic solvents, [¹¹C]CO₂ is usually converted into a more reactive secondary precursor (e.g. $[^{11}C]$ iodomethane²) in order to assemble a radiolabelled molecule of interest.³ The production of a secondary precursor results in significant losses due to non-quantitative yields and additional processing times. Methods for the direct incorporation of $[^{11}C]CO_2$ into molecules of interest would therefore have significant advantages over traditional carbon-11 labelling methodologies, since the processing times and losses due to technical handling could be minimised.

Significant efforts have been made to improve the solubility of $[^{11}C]CO_2$ in organic solvents, resulting in two approaches for

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the fixation and trapping of $[^{11}C]CO_2$:⁴ (1) trapping of $[^{11}C]CO_2$ using highly reactive organometallic reagents, such as Grignard or organolithium reagents to produce $[^{11}C]carboxylic acids and$ derivatives (*e.g.*acid chlorides);⁴ (2) utilisation of trappingagents such as 1,8-diazabicyclo[5.4.0]undec-7-ene (DBU)⁵ and2-tertbutylimino-2-diethylamino-1,3-dimethyl-perhydro-1,3,2 $diazaphosphorine (BEMP)⁶ to form a labile bond with <math>[^{11}C]CO_2$. These approaches have been applied to radiolabel $[^{11}C]$ carboxylic acids,⁷ $[^{11}C]$ amides,⁸ $[^{11}C]$ amines,⁹ $[^{11}C]$ acyl chlorides,¹⁰ $[^{11}C]$ ureas,^{5,11} $[^{11}C]$ carbamates^{6,12} and $[^{11}C]$ isocyanates.¹³

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Current methods for the preparation of ¹¹C-labelled amides utilise the coupling between [¹¹C]carboxylic acids and amines at high temperatures,⁸ assisted by microwave irradiation¹⁴ or through activation to [¹¹C]acyl chlorides using thionyl choride¹⁵ or phthaloyl chloride¹⁶ (Scheme 1). These methodologies often lead to low molar radioactivities since the reagents used readily react with atmospheric CO₂ and have a synthesis time ranging from 5–35 minutes.^{8,14–16} In order to minimise this isotopic dilution, extreme efforts are required to control reagent stoichiometry, stability and exclusion of moisture and atmospheric CO₂ from the reagents and reaction system.

Boronic esters have higher stability to air and moisture compared with Grignard reagents and are able to react with $[^{11}C]CO_2$ to form $[^{11}C]$ amides in two steps within 10–15 min from $[^{11}C]CO_2$. However, this method is limited to the production of aromatic $[^{11}C]$ amides (R = Ar, Scheme 1).⁷



Scheme 1 Different approaches available for the preparation of [¹¹C]amides using Grignard reagents or boronic esters.

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 $\label{eq:scheme 2} \begin{array}{l} \mbox{Radiosynthetic approach to radiolabelled $[^{11}C]$ ureas5 and $[^{11}C]$ anides (this work) from cyclotron-produced $[^{11}C]$ CO_2. \end{array}$

To address the shortcomings of current $\begin{bmatrix} 1^{11}C \end{bmatrix}$ amide labelling methodologies, we sought to develop an efficient method for producing [¹¹C]amides in short synthesis times, high molar radioactivities and with applicability to a range of amide derivatives bearing alkyl and aryl groups on R and R¹ positions (Scheme 2). The synthetic strategy was inspired by our previous work on the labelling of $[^{11}C]$ ureas,⁵ which involved: (1) reaction of an amine with $[^{11}C]CO_2$ to form a $[^{11}C]$ carbamate anion in the presence of DBU and acetonitrile (MeCN) as a solvent (Scheme 2); (2) conversion from $\begin{bmatrix} 11 \\ C \end{bmatrix}$ carbamate anion to an ^{[11}C]isocyanate or an ^{[11}C]oxyphosphonium intermediate using Mitsunobu reagents (tri-n-butyl phosphine and di-tert-butyl azodicarboxylate, Bu_3P and DBAD; (3) reaction of the [¹¹C]isocyanate intermediate with another molecule of amine to yield the corresponding [¹¹C]urea derivative. Based on this pathway, it was hypothesised that the synthesis of [¹¹C]amides might be achieved by coupling the intermediate $\begin{bmatrix} 11 \\ C \end{bmatrix}$ isocyanate with a Grignard reagent. To test this hypothesis, the formation of amides via the Mitsunobu reaction was tested using both macroscopic amounts of non-radioactive CO2 and tracer amounts of radiolabelled $[^{11}C]CO_2$.

The synthesis of **1A** was chosen as model reaction (Table 1). Initial experiments were performed adapting previously established conditions for the synthesis of urea derivatives.¹⁷ CO₂ was bubbled into a solution of benzylamine (**1**) and DBU in MeCN at room temperature (r.t.) for 40 min. DBU facilitates the

Table 1 Optimization of substrates and reaction conditions for the synthesis of ${\bf 1A}$

N ^{-H} H 1		i) CO ₂ , DBU, MeCN ii) DBAD, Bu ₃ P, MeCh iii)MgBr A		O N H H			
Entry ^a	DBU (equiv.)	DBAD (equiv.)	Bu ₃ P (equiv.)	Yield of $\mathbf{1A}^{b}$ (%)			
1	0.1	2	2	0			
2	0.1	3.8	3.8	0			
3	0.2	3.8	3.8	0			
4	0.05	1	1	0			
5	0.05	2	2	46 ± 8^c			
6	0.05	3.8	3.8	44^d			
7	0.05	72	7 2	0			

^{*a*} Reaction conditions: CO₂ was bubbled in a solution of **1** (138.6 µmol, 1.0 equiv.), DBU (0.05–0.2 equiv.) in MeCN (1 mL), r.t. for 40 min. Mitsunobu reagents (7.2–1 equiv.) in MeCN (0.5 mL) were added and the solution stirred for 10 min. **A** (7.2 equiv. of a 0.5 M solution in THF) was added and quenched after 30 min. ^{*b*} Yield of isolated **1A** calculated from compound **1**. ^{*c*} N = 4. ^{*d*} N = 1.

formation and stabilisation of the carbamate anion. Mitsunobu reagents were subsequently added and the reaction stirred for 10 min before adding an excess of 1-propynylmagnesium bromide (A - as a 0.5 M solution in THF, 7.2 equiv.). The reaction was quenched after 30 min and the conversion to compound 1A was evaluated. Using the reaction conditions previously used for the synthesis of ureas, led to no observable quantities of 1A (Table 1, entry 1). Increasing the concentration of Mitsunobu reagents or DBU had no beneficial effect on the reaction yield (Table 1, entries 2 and 3). As next step, lowering the amount of DBU from 0.1 to 0.05 equivalents different Mitsunobu reagent concentrations were tested (Table 1, entries 4-7). 1A was obtained in 46% and 44% yield using 2 and 3.8 equivalents of Mitsunobu reagent, respectively (Table 1, entries 5 and 6). Using an equimolar or an excess of Mitsunobu reagents did not vield 1A (Fig. S4, ESI[†]).

The synthesis of primary, secondary and aromatic amides was explored to test the scope of the reaction (Table 2). When using the secondary amine *N*-methylbenzylamine (2), amide 2A was not obtained (Table 2). The poorly nucleophilic aromatic amine (3) did not form the target product, probably due to its inability to form an isocyanate as reported previously.¹⁸ The activated aromatic amine (4) however gave 4A in good yield



Reaction conditions: (i) 1–6 (1 equiv.) and DBU (0.05 equiv.) in MeCN or Et₂O, CO₂ (1 atm) r.t. for 40 min; (ii) DBAD (2 equiv.) and Bu₃P (2 equiv.) in MeCN or Et₂O, 10 min; (iii) A–C (as a 0.5–1.0 M solution in THF, 7.2 equiv.) and quenching after 30 min. Yield of isolated compound calculated from the corresponding amine derivative (N = 4 for 1A and N = 1 for 4A, 1B, 1C, 5C, 6C).

(37%, Table 2) due to its ability to form an isocyanate intermediate in agreement with the results obtained by others.¹⁸

The reactivity of different Grignard reagents in MeCN or Et_2O was also investigated. Reaction of **1** with ethylmagnesium bromide (**B** – as a 1.0 M solution in THF, 7.2 equiv.) in MeCN gave **1B** in a good yield (57%, Table 2), whereas when using phenylmagnesium bromide (**C** – as a 1.0 M solution in THF, 7.2 equiv.) in MeCN, the target amide was not obtained. Amide formation between isocyanates and bulky Grignard reagents such as **C** has been successfully observed using Et_2O as solvent at 0 °C.¹⁹ Encouragingly, substituting the MeCN with diethyl ether (Et_2O) under these conditions we obtained **1C**, **5C** and **6C** in yields of 5–28% (Table 2). These results indicate that primary and activated aromatic amines are able to form isocyanate intermediates and subsequently react with a broad range of Grignard reagents forming amides.

Our amide preparation methodology developed using nonradioactive CO₂ was subsequently applied to the synthesis of $[^{11}C]$ amides using $[^{11}C]CO_2$. Translating the reaction conditions from synthetic chemistry to carbon-11 chemistry, we were unable to obtain $[^{11}C]$ **1A** (Table 3, entry 1), with only $[^{11}C]$ dibenzylurea present as a by-product. The solution trapped the cyclotronproduced $[^{11}C]CO_2$ in high efficiency (98%). As the $[^{11}C]CO_2$ chemistry is performed on a nano-molar scale the $[^{11}C]$ amide reaction conditions were further investigated. In an effort to reduce the amount of $[^{11}C]$ dibenzylurea by-product and optimise

Table 3 Radiosynthesis of [¹¹ C]1A							
$(1) \begin{bmatrix} 1^{11}C]CO_2, DBU, MeCN \\ ii) DBAD, Bu_3P, MeCN \\ H \\ iii) = MgBr \\ MgBr \\ I \\ I^{11}C]IA$							
Entry ^a	DBU (equiv.)	Mitsunobu reagents (equiv.)	$T(^{\circ}C)$	RCY of [¹¹ C] 1A ^b (%)			
1 ^{<i>c</i>}	0.05	2	20	0^g			
2^{c}	1	1	20	2^f			
3^d	1	1	20	5^{f}			
4^d	2.5	1	20	4^f			
5^d	5	1	20	8^f			
6^d	10	1	20	9.5 ± 7^g			
7^d	10	1	30	10 ± 2^g			
8^d	10	1	40	20 ± 4^g			
9^d	10	1	50	18 ± 7^g			
10^d	10	1	80	27 ± 8^g			
11^d	10	2	20	10 ± 4^g			
12^d	10	2	50	36 ± 4^g			
13^d	10	2	80	$45~\pm~5^g$			
14^d	10	4	80	50 ± 9^h			
15^d	10	8	80	50 ± 5^h			
16^d	20	2	80	28 ± 5^g			
17^d	30	4	80	30 ± 2^g			
18^e	10	8	80	0^f			

^{*a*} Reaction conditions: [¹¹C]CO₂ was bubbled in a solution of 1 (32 μmol, 1 equiv.), DBU (0.05–30 equiv.) in MeCN (160 μL) at room temperature. Then, the reaction mixture was heated (20–80 °C) for 30 seconds. Mitsunobu reagents (1–8 equiv.) in MeCN (100 μL) were added and stirred for 10 s. A (8 equiv. of a 0.5 M solution in THF) was added and quenching after 1 min. ^{*b*} RCY determined by radio-HPLC not decay-corrected. ^{*c*} 138.5 µmol of 1. ^{*d*} 32 µmol of 1. ^{*e*} 32 µmol of 1 in DMF. ^{*f*} N = 1. ^{*g*} N = 2. ^{*h*} N = 3.

the radiochemical yield (RCY) of $[^{11}C]$ **1A**, the influence of changing reagent concentrations (amine, DBU and Mitsunobu reagents), temperature and solvent were studied.

Initial experiments were performed with higher amounts of DBU. A low RCY²⁰ (2%, Table 3, entry 2) of $[^{11}C]$ **1A** was observed using 1 equiv. of DBU with $[^{11}C]$ dibenzylurea as a major by-product. When the reaction was carried out at a lower amine concentration, a slight increase in RCY was observed (5%, Table 3, entry 3). Interestingly, increasing the DBU concentration further to 1, 2.5, 5 and 10 equiv. led to a RCY of 10% (Table 3, entries 3–6).

We have previously reported that the RCY of $[^{11}C]$ ureas is dependent on reaction temperature.⁵ A similar trend was observed for the synthesis of $[^{11}C]$ amides. Indeed, varying the temperature from 20 to 30, 40, 50 and 80 °C improved the RCY (Table 3, entries 6–13). The RCY increased from 9% at 20 °C to 27% at 80 °C (Table 3, entries 6–10, Fig. S5, ESI†). Similarly by adding 2 equiv. of Mitsunobu reagents the RCY increased from 10% at 20 °C to 45% at 80 °C (Table 3, entries 11–13, Fig. S5, ESI†).

Interestingly, the increase of Mitsunobu reagents from 1 to 2, 4 and 8 equiv. (Table 3, entries 10 and 13–15), significantly increased the RCY from 27% to 50%. Increasing DBU concentration from 10 to 30 equivalents did not improve the RCY further (28–30%, Table 3, entries 16–17). Good RCY's of the desired [¹¹C]amide were observed when MeCN was used as a solvent (50%) while DMF was detrimental for the RCY (0%, Table 3, entry 18).

To demonstrate the potential utility of this novel labelling strategy, the endogenous neurotransmitter melatonin, was radiolabeled with carbon-11 (Scheme 3). The incorporation of $[^{11}C]CO_2$ into $[^{11}C]melatonin$ was 36% – determined by radioHPLC 2 min from end of $[^{11}C]CO_2$ delivery with an estimated molar radioactivity of 70–100 GBq µmol⁻¹, consistent with the molar radioactivities obtained for other ¹¹C-labelled tracers at our institution.²¹

In summary a rapid one-pot methodology for the synthesis of amides has been successfully developed. Readily available CO_2 was incorporated into primary or activated aromatic amines in the presence of Mitsunobu reagents to form the isocyanate intermediate. Grignard reagents were then reacted with the isocyanate to form the corresponding amides. The approach was optimised for the radiolabelling of [¹¹C]amides using [¹¹C]CO₂ directly from the cyclotron. The synthesis time from end of [¹¹C]CO₂ delivery was less than 2 minutes. Radiolabelling of a biologically relevant biomolecule, [¹¹C]melatonin, was achieved. A limitation of the proposed technique is the restricted number of commercially available Grignard reagents



Scheme 3 Reaction conditions: [¹¹C]CO₂ was bubbled in a solution of **7** (32 µmol, 1 equiv.), DBU (10 equiv.) in MeCN (160 µL) at room temperature. Then, the reaction mixture was heated (50 °C) for 30 seconds. Mitsunobu reagents (8 equiv.) in MeCN (100 µL) were added and stirred for 10 s. **D** (8 equiv., 3.0 M in Et₂O) was added. The reaction was quenched after 1 min.

and their chemical stability. However these reagents can easily be produced in-house upon demand. As some functional groups are not compatible with Grignard reagents this should be borne in mind when using this methodology. In conclusion, this alternative approach offers new possibilities for the rapid and efficient carbon-11 labelling of drugs containing amide functional groups for PET imaging applications.

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Notes and references

- 1 (a) G. Antoni, J. Labelled Compd. Radiopharm., 2015, 58, 65-72; (b) J. S. Fowler and A. P. Wolf, Acc. Chem. Res., 1997, 30, 181-188.
- (a) B. Langstrom and H. Lundqvist, *Int. J. Appl. Radiat. Isot.*, 1976, 27, 357–363; (b) C. Marazano, M. Maziere, G. Berger and D. Comar, *Int. J. Appl. Radiat. Isot.*, 1977, 28, 49–52.
- 3 P. W. Miller, N. J. Long, R. Vilar and A. D. Gee, *Angew. Chem., Int. Ed.*, 2008, **47**, 8998–9033.
- 4 B. H. Rotstein, S. H. Liang, J. P. Holland, T. L. Collier, J. M. Hooker, A. A. Wilson and N. Vasdev, *Chem. Commun.*, 2013, **49**, 5621–5629.

- 5 (*a*) A. K. Haji Dheere, N. Yusuf and A. Gee, *Chem. Commun.*, 2013, **49**, 8193–8195; (*b*) A. K. Dheere, S. Bongarzone, C. Taddei, R. Yan and A. D. Gee, *Synlett*, 2015, 2257–2260.
- 6 A. A. Wilson, A. Garcia, S. Houle and N. Vasdev, *Org. Biomol. Chem.*, 2010, **8**, 428–432.
- 7 P. J. Riss, S. Lu, S. Telu, F. I. Aigbirhio and V. W. Pike, *Angew. Chem., Int. Ed.*, 2012, **51**, 2698–2702.
- 8 C. Aubert, C. Huard-Perrio and M.-C. Lasne, J. Chem. Soc., Perkin Trans. 1, 1997, 2837-2842.
- 9 (a) C. Perrio-Huard, C. Aubert and M.-C. Lasne, J. Chem. Soc., Perkin Trans.
 1, 2000, 311–316; (b) F. Liger, T. Eijsbouts, F. Cadarossanesaib,
 C. Tourvieille, D. Le Bars and T. Billard, Eur. J. Org. Chem., 2015, 6434–6438.
- 10 D. Le Bars, S. K. Luthra, V. W. Pike and C. L. Duc, Int. J. Rad. Appl. Instrum. A, 1987, 38, 1073–1077.
- 11 E. W. van Tilburg, A. D. Windhorst, M. van der Mey and J. D. M. Herscheid, J. Labelled Compd. Radiopharm., 2006, **49**, 321–330.
- 12 J. M. Hooker, A. T. Reibel, S. M. Hill, M. J. Schueller and J. S. Fowler, *Angew. Chem., Int. Ed.*, 2009, **48**, 3482–3485.
- 13 A. A. Wilson, A. Garcia, S. Houle, O. Sadovski and N. Vasdev, Chemistry, 2011, 17, 259-264.
- 14 S.-Y. Lu, J. Hong and V. W. Pike, J. Labelled Compd. Radiopharm., 2003, 46, 1249–1259.
- 15 D. R. Hwang, N. R. Simpson, J. Montoya, J. J. Man and M. Laruelle, *Nucl. Med. Biol.*, 1999, 26, 815–819.
- 16 S. K. Luthra, V. W. Pike and F. Brady, J. Chem. Soc., 1985, 1423–1425.
- 17 S. L. Peterson, S. M. Stucka and C. J. Dinsmore, *Org. Lett.*, 2010, **12**, 1340–1343.
- 18 D. Saylik, M. J. Horvath, P. S. Elmes, W. R. Jackson, C. G. Lovel and K. Moody, J. Org. Chem., 1999, 64, 3940–3946.
- 19 G. Schafer, C. Matthey and J. W. Bode, *Angew. Chem., Int. Ed.*, 2012, 51, 9173–9175.
- 20 The radiochemical yields (RCY) determined by radio-HPLC of the crude product.
- 21 An example of clinal tracer produced in our instute is [¹¹C]Methionine produced by ¹¹C-methylation using [¹¹C]CH₃I obtained *via* the "gas-phase" method.