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Antonio del Vecchio, Alex Talbot, Fabien Caillé, Arnaud Chevalier, Antoine Sallustrau, et al.. Carbon isotope labeling of carbamates by late-stage  $[^{11}\text{C}]$ ,  $[^{13}\text{C}]$  and  $[^{14}\text{C}]$  carbon dioxide incorporation. Chemical Communications, In press, 10.1039/D0CC05031H . cea-02933909

**HAL Id: cea-02933909**

**<https://cea.hal.science/cea-02933909>**

Submitted on 8 Sep 2020

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# Carbon Isotope Labeling of Carbamates by Late-Stage [ $^{11}\text{C}$ ], [ $^{13}\text{C}$ ] and [ $^{14}\text{C}$ ] Carbon Dioxide Incorporation

Received 00th January 20xx,  
Accepted 00th January 20xx

DOI: 10.1039/x0xx00000x

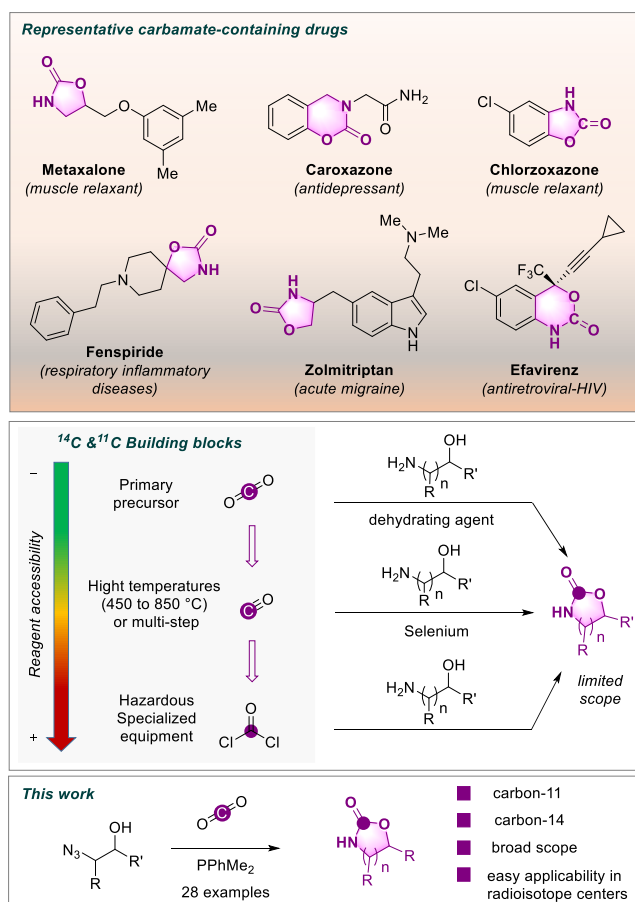
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A general procedure for the late-stage [ $^{11}\text{C}$ ], [ $^{13}\text{C}$ ] and [ $^{14}\text{C}$ ] carbon isotope labeling of cyclic carbamates is reported. This protocol allows the incorporation of carbon dioxide, the primary source of carbon-14 and carbon-11 radioisotopes, in a direct, cost-effective and sustainable manner. This method was applied to a variety of carbamates, including pharmaceuticals. A disconnection / reconnection strategy, involving ring opening / isotopic closure, was also implemented to efficiently broaden the approach.

## Introduction

Radiolabeling with carbon isotopes is an attractive strategy in human/veterinary drug development, as well as in plant protecting agent research.<sup>1</sup> Because of the ubiquitous presence of carbon in nature, replacement of carbon-12 with its radioactive isotopes is possible on virtually any organic molecule and structural alterations on the non-labeled original molecule are negligible. Interestingly, the two relevant radioactive isotopes of carbon display antipodean physical properties, which usually require *ad hoc* strategies for their insertion into organic structures. Carbon-14 ( $^{14}\text{C}$ ) is a long-lived  $\beta$  emitter, whose relevance is well-established in the determination of (pre)clinical drug metabolism and biodistribution.<sup>2</sup> Nowadays,  $^{14}\text{C}$  radio-synthesis remains challenging due to the limited number of available raw materials, their prohibitive cost and the generation of long lasting waste ( $t_{1/2}$  5730 years).<sup>3</sup> On the other hand, carbon-11 ( $^{11}\text{C}$ ) is a short-lived positron emitter ( $\beta^+$ ), whose implication in positron emission tomography (PET) can hardly be overstated. Challenges related to the 20.4 minutes half-life and the limited amounts of isotope generated by a cyclotron dramatically affect the chemical space tackled in PET centers, on a routine basis.<sup>4</sup> In such a dichotomy, the two isotopes share the same primary chemical source: carbon dioxide ( $^{14}\text{C}$ CO<sub>2</sub> and  $^{11}\text{C}$ CO<sub>2</sub>). Given the complementarity of their applications, drug discovery would highly benefit from general procedures easily transmutable to both  $^{11}\text{C}$  and  $^{14}\text{C}$  radioisotopes. Unfortunately, recent developments in late-stage carbon isotope labeling are specific and often unsuitable for application to both radioisotopes, either due to practical and/or safety related issues.<sup>5,6</sup> Carbamates are a fundamental functional group, largely found in biologically active compounds and marketed pharmaceuticals (Scheme 1, top).<sup>7</sup> In particular, cyclic carbamates are generally metabolically stable and the carbonyl moiety represents a suitable position for labeling.<sup>8</sup> Due to its high reactivity,

the use of [ $^{14}\text{C}$ ] and [ $^{11}\text{C}$ ]phosgene has been investigated, but shortcomings related to safety and the requirement for specialized apparatus have largely limited its utilization (Scheme 1, bottom).<sup>9</sup>



**Scheme 1.** Top: Relevant examples of cyclic carbamates containing pharmaceuticals; middle: synthetic strategies for cyclic carbamate radiosynthesis; bottom: Staedinger / aza-Wittig approach.

In 2002, Långström reported a selenium-mediated carbamylation reaction in presence of [ $^{11}\text{C}$ ]carbon monoxide.<sup>10</sup> The procedure was applied to simple carbamate derivatives, and later on one drug (Zolmitriptan).<sup>11,12</sup> Methods that utilize [ $^{11}\text{C}$ ]CO<sub>2</sub> and

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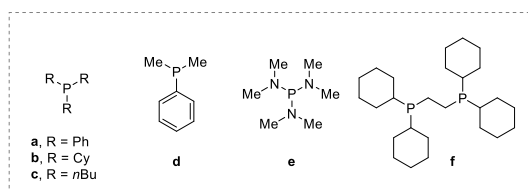
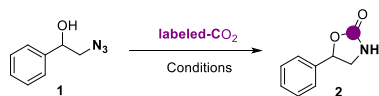
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[ $^{14}\text{C}$ ]CO $_2$  are appealing, but their application was mainly focused on linear carbamates.<sup>13</sup> Vasdev and co-workers labeled an cyclic carbamate ([ $^{11}\text{C}$ ]SL25.1188) using BEMP as fixing base in presence of POCl $_3$  as dehydrating reagent.<sup>14</sup> In 2019, Horkka *et al.* reported a procedure where the dehydration step was performed under Mitsunobu conditions, and labeled one cyclic carbamate, benzoxazol-2-one.<sup>15</sup> Very recently, Jakobsson *et al.* reported the synthesis of [ $^{11}\text{C}$ ]carbonyl difluoride.<sup>16</sup> While this reagent has a similar reactivity to phosgene, it is generated from [ $^{11}\text{C}$ ]CO,<sup>17</sup> a secondary precursor obtained from [ $^{11}\text{C}$ ]CO $_2$  with technically challenging and time-consuming conditions which are unpractical to implement in most PET centers.

The current state of the art highlights the need for general procedures that would prove effective on elaborate molecules. Here, we report on a late-stage labeling protocol of cyclic carbamates that utilized CO $_2$  as isotope source. This procedure is suitable to all carbon isotopes, including  $^{13}\text{C}$ ,  $^{11}\text{C}$  and  $^{14}\text{C}$ , on a variety of substrates including pharmaceuticals.

## Results and discussion

Our recent results on the late-stage labeling of cyclic ureas<sup>18</sup> encouraged us to investigate a Staudinger aza-Wittig approach (SAW) for the  $^{14}\text{C}$ - and  $^{11}\text{C}$ -labeling of cyclic carbamates, directly from CO $_2$ . Reaction optimization was conducted at room temperature in 5 minutes on model substrate **1**, using stoichiometric [ $^{13}\text{C}$ ]CO $_2$ , as  $^{14}\text{C}$ -surrogate, and its precise addition was monitored utilizing a RC Tritac manifold.

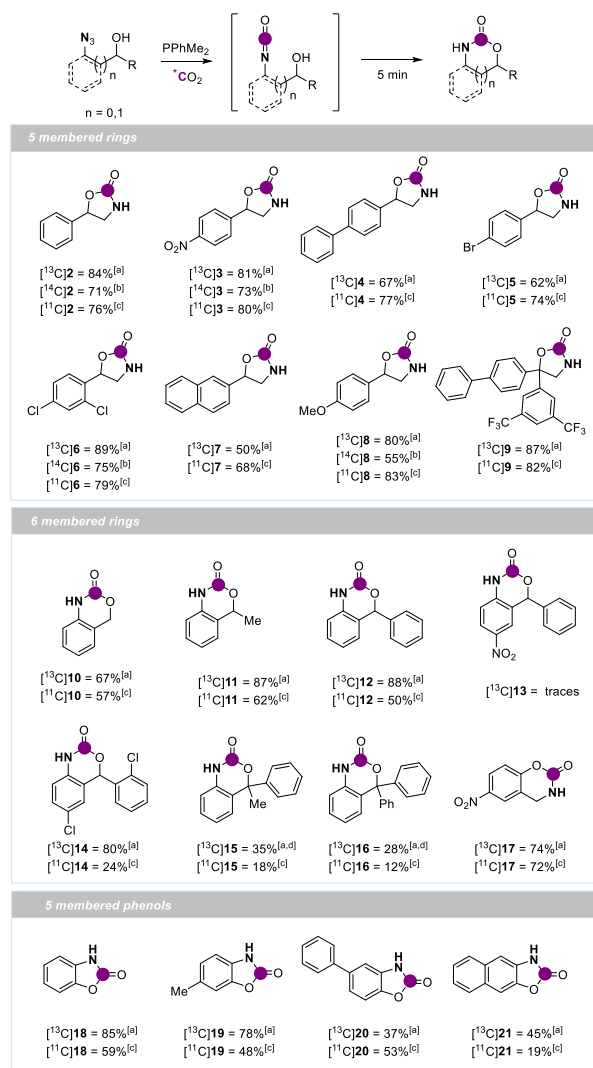


Entry	$^*\text{CO}_2$	Phosphine	Temperature	Time	Yield (%)
1	[ $^{13}\text{C}$ ]CO $_2$	<b>a</b>	25 °C	5 min	63
2	[ $^{13}\text{C}$ ]CO $_2$	<b>b</b>	25 °C	5 min	46
3	[ $^{13}\text{C}$ ]CO $_2$	<b>c</b>	25 °C	5 min	62
4	[ $^{13}\text{C}$ ]CO $_2$	<b>d</b>	25 °C	5 min	84
5	[ $^{13}\text{C}$ ]CO $_2$	<b>e</b>	25 °C	5 min	0
6	[ $^{13}\text{C}$ ]CO $_2$	<b>f</b>	25 °C	5 min	0
7	[ $^{14}\text{C}$ ]CO $_2$	<b>d</b>	25 °C	5 min	71 <sup>[a]</sup>
8	[ $^{11}\text{C}$ ]CO $_2$	<b>d</b>	70 °C	5 min	76 <sup>[b]</sup>

**Table 1.** Optimization of the Staudinger aza-Wittig procedure [a] radiochemical yield; [b] decay-corrected radiochemical conversion. Carbon-13 and -14 experiments were performed in 0.7 mL DMF on a 0.1 mmol scale, in presence of stoichiometric amounts of [ $^{13}\text{C}$ ] and [ $^{14}\text{C}$ ]CO $_2$ ; for carbon-11 labeling, precursor **1** and the phosphine were typically in a 100-fold excess compared to [ $^{11}\text{C}$ ]CO $_2$  (see SI for details).

As expected, the nature of the phosphine had a major impact on the transformation. While the use of triphenylphosphine (**a**) allowed

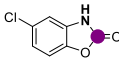

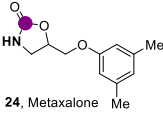
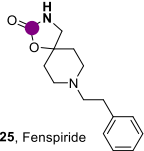
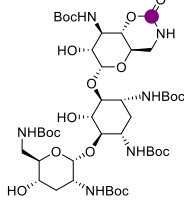
the formation of [ $^{13}\text{C}$ ] **2** in 63% isolated yield, with electron-rich tricyclohexylphosphine (**b**) the yield dropped to 46%. When tri-*n*-butylphosphine (**c**) was utilized, carbamate [ $^{13}\text{C}$ ] **2** was isolated in 62% yield. The best result was obtained with dimethylphenylphosphine (**d**) which provided the corresponding [ $^{13}\text{C}$ ] carbamate with 84% isolated yield. The use of (**e**), reminder of the core structure of commonly used iminophosphoranes trapping agents such as BEMP, was unproductive. While the phosphazide is formed rapidly from (**e**), the subsequent nitrogen release required harsh conditions, as reported by Lyapkalo and co-workers.<sup>19</sup> Finally, the use of bidentate phosphine (**f**) did not provide the desired product.<sup>20</sup> Without further optimization, the procedure was applied to  $^{14}\text{C}$ . Pleasingly, [ $^{14}\text{C}$ ] **2** was isolated in 71% RCY directly from [ $^{14}\text{C}$ ]CO $_2$ . Compared to previous methods, this procedure is cost-effective and drastically minimizes the generation of radioactive waste. Based on the short reaction time required, application to  $^{11}\text{C}$ -labeling seemed feasible. Despite the diametrically different environment and the CO $_2$  stoichiometry, [ $^{11}\text{C}$ ] **2** was labeled in 76% decay-corrected (d.c.) RCC, under slightly different conditions (70 °C). Importantly, the radiosynthesis was carried out in automated modules, according to nuclear safety requirements for PET tracers production.



**Scheme 2.** Late-stage labeling of cyclic carbamates with carbon isotopes. [a] Yield of the isolated product; [b] radiochemical yield; [c] d.c. radiochemical conversion; [d] reaction temperature, 150 °C. See SI for experimental details.

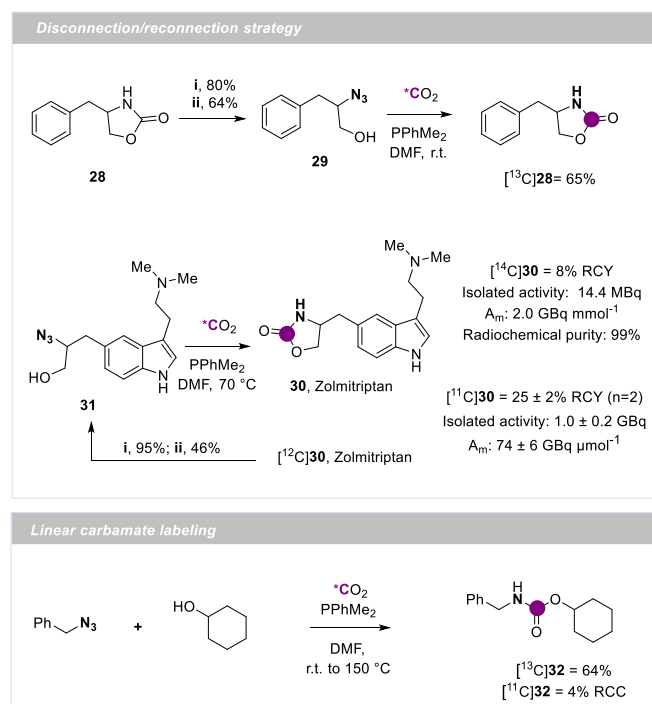
With this optimized set of conditions, we explored the scope of the transformation on a series of substituted azido alcohols (Scheme 2). Aliphatic 5-membered carbamates were labeled with  $^{13}\text{C}$  to provide [ $^{13}\text{C}$ ]3 to [ $^{13}\text{C}$ ]9 in 50 to 89% yield. The electronic nature of substituents on the aromatic ring did not affect the reaction outcome and also a sterically hindered quaternary alcohol precursor delivered **9** in 87% yield. Selected substrates were labeled with radiocarbon, to provide [ $^{14}\text{C}$ ]2, **3**, **6** and **8** in 55 to 75% RCY, under otherwise identical reaction conditions. Pleasingly,  $^{11}\text{C}$  labeling proceeded smoothly in all tested substrates in 68 to 83% d.c. RCC. Next, we looked at the labeling of six membered ring carbamates (Scheme 2). The presence of a methyl or phenyl substituent in *alpha* position to the hydroxyl did not affect its reactivity and 2-benzoxazinones **10–12** could be efficiently labeled with  $^{13}\text{C}$  and  $^{11}\text{C}$  isotopes. These derivatives would be quite challenging to label with current methods using  $\text{CO}_2$ , because of the poor nucleophilicity of anilines.<sup>13</sup> On the other hand, from tertiary alcohol precursors, derivatives **15** and **16** were delivered with a significant drop in the yields and required higher reaction temperature (Scheme 2). In addition, when a strong electron-withdrawing *p*- $\text{NO}_2$  substituent was present on the aromatic azide, only traces amounts of [ $^{13}\text{C}$ ]13 were recovered. The increased electrophilicity of the isocyanate intermediate, which presumably undergoes fast hydrolysis under the reaction conditions, was detrimental to the process. Interestingly, also regioisomeric carbamate **17** could be labeled both with  $^{11}\text{C}$  and  $^{13}\text{C}$  from the corresponding benzylic azide precursor. In this case, the presence of a nitro group in *para* position to the alcohol was well tolerated: [ $^{13}\text{C}$ ]17 and [ $^{11}\text{C}$ ]17 were obtained in 74% and 72% d.c. RCC, respectively. Benzoxazol-2-ones are widely present in pharmaceuticals and their late-stage labeling is a worthwhile endeavour.<sup>21</sup> After some optimization, benzoxazol-2-ones **18–21** could be labeled with  $^{13}\text{C}$ , in presence of 2 equiv. of DIPEA. Interestingly, the use of  $\text{CO}_2$  fixing agents such as DBU, DABCO, DBN and TBD was detrimental to the reaction outcome and did not provide formation of the desired product.<sup>20</sup> On the other hand, when the reaction on model substrate **18** was performed in the absence of DIPEA, the isolated yield dropped to 64%.<sup>20</sup> For short-lived  $^{11}\text{C}$ , the optimal conditions were in absence of additive; gentle heating at 70 °C was required and [ $^{11}\text{C}$ ]18–21 were obtained in 19 to 59% RCC. With these results in hand, we next turned our attention to the labeling of pharmaceutically relevant derivatives (Scheme 3). Chlorzoxazone, a FDA approved drug displaying muscle relaxant properties,<sup>22</sup> was labeled from the corresponding azidoalcohol precursor under standard conditions to isolate [ $^{14}\text{C}$ ]22 in 39% RCY and high molar activity ( $A_m$ ) 2.0 GBq  $\text{mmol}^{-1}$ . The short-lived isotopomer, ready-to-inject [ $^{11}\text{C}$ ]22 was isolated after HPLC purification and formulation in  $37 \pm 2\%$  d.c. RCY and d.c.  $A_m$  of  $85 \pm 4$  GBq/ $\mu\text{mol}$ . Caroxazone precursor **23** was labeled in 30% RCY from [ $^{14}\text{C}$ ]CO<sub>2</sub> and  $25 \pm 5\%$  RCY from [ $^{11}\text{C}$ ]CO<sub>2</sub>. Metaxalone, an aliphatic five-membered carbamate with muscle relaxant properties, was obtained in 59% RCY and  $44 \pm 3\%$  d.c. RCY with carbon-14 and carbon-11, respectively. Next, we focused on Fenspiride, an anti-inflammatory drug, which was recently labeled with tritium.<sup>23</sup> The required *beta*-azido alcohol precursor was obtained in a three-step sequence, including an alkylation, a Corey-Chaykovsky epoxidation and a regioselective ring opening azidation from commercially available materials.<sup>20</sup> This substrate underwent  $^{14}\text{C}$ -labeling in 45% RCY and  $^{11}\text{C}$ -labeling in  $23 \pm 3\%$  d.c. RCY. The procedure was also applied for the  $^{14}\text{C}$  and  $^{11}\text{C}$  labeling of N-Boc protected Tobramycine derivative **26**,<sup>24</sup> on a non-pharmacophoric position, according to previous SAR studies.<sup>25</sup> The SAW approach allowed to isolate the carbamates [ $^{14}\text{C}$ ]26 and [ $^{11}\text{C}$ ]26

in 35% and  $68 \pm 2\%$  d.c. RCY, respectively. In all cases, consistent high  $A_m$  and radiochemical purities for both radioisotopes were obtained.

Carbon-14	Labeling of drug derivatives	Carbon-11
[ $^{14}\text{C}$ ]22 = 39% RCY Isolated activity: 70.8 MBq $A_m$ : 2.0 GBq $\text{mmol}^{-1}$ Radiochemical purity: 99%	 22, Chlorzoxazone	[ $^{11}\text{C}$ ]22 = $37 \pm 2\%$ RCY (n=2) Isolated activity: $2.8 \pm 0.3$ GBq $A_m$ : $85 \pm 4$ GBq/ $\mu\text{mol}$ Radiochemical purity: 99%
[ $^{14}\text{C}$ ]23 = 30% RCY Isolated activity: 55.4 MBq $A_m$ : 2.0 GBq $\text{mmol}^{-1}$ Radiochemical purity: 99%	 23, Caroxazone precursor	[ $^{11}\text{C}$ ]23 = $25 \pm 5\%$ RCY (n=2) Isolated activity: $0.9 \pm 0.1$ GBq $A_m$ : $75 \pm 10$ GBq $\mu\text{mol}^{-1}$ Radiochemical purity: 99%
[ $^{14}\text{C}$ ]24 = 59% RCY Isolated activity: 111.0 MBq $A_m$ : 2.0 GBq $\text{mmol}^{-1}$ Radiochemical purity: 99%	 24, Metaxalone	[ $^{11}\text{C}$ ]24 = $44 \pm 3\%$ RCY (n=2) Isolated activity: $2.1 \pm 0.4$ GBq $A_m$ : $78 \pm 3$ GBq $\mu\text{mol}^{-1}$ Radiochemical purity: 99%
[ $^{14}\text{C}$ ]25 = 45% RCY Isolated activity: 90.3 MBq $A_m$ : 1.7 GBq $\text{mmol}^{-1}$ Radiochemical purity: 99%	 25, Fenspiride	[ $^{11}\text{C}$ ]25 = $23 \pm 3\%$ RCY (n=3) Isolated activity: $0.8 \pm 0.2$ GBq $A_m$ : $81 \pm 8$ GBq $\mu\text{mol}^{-1}$ Radiochemical purity: 99%
[ $^{14}\text{C}$ ]26 = 35% RCY Isolated activity: 48.5 MBq $A_m$ : 1.9 GBq $\text{mmol}^{-1}$ Radiochemical purity: 99%	 26, Tobramycine derivative	[ $^{11}\text{C}$ ]26 = $68 \pm 2\%$ RCY (n=2) Isolated activity: $1.1 \pm 0.2$ GBq Radiochemical purity: 99%

**Scheme 3.** Late-stage radiolabeling of pharmaceutically relevant carbamates. RCY: radiochemical yield;  $A_m$ : molar activity. For carbon-11, RCY and  $A_m$  are decay-corrected.

The concept of isotope exchange has been largely exploited for hydrogen,<sup>26</sup> but only recently introduced for carbon and, to our knowledge, strictly limited to carboxylic acids.<sup>5</sup> In order to accelerate the access to labeled carbamates and limit the synthesis of precursors, we envisioned a disconnection/reconnection strategy (DRS).<sup>9a</sup> The DRS was first validated on model compound **28**. After carbamate cleavage under strong basic conditions, the corresponding *beta*-amino alcohol underwent a diazo-transfer reaction to obtain the azido alcohol **29** and further converted into [ $^{13}\text{C}$ ]28 in 65% yield. With this procedure in hand, we next looked to Zolmitriptan, a compound in use for the treatment of acute migraine, commercialized by AstraZeneca. From commercially available unlabeled Zolmitriptan, the azido alcohol precursor **31** was obtained in 43% yield, over two steps. Subsequent SAW reaction allowed to obtain [ $^{14}\text{C}$ ]30 in moderated 8% RCY, nonetheless with a high  $A_m$  of 2 GBq  $\text{mmol}^{-1}$ . Carbon-11 labeled Zolmitriptan was also synthesized in  $25 \pm 2\%$  RCY and  $A_m$   $74 \pm 6$  GBq/ $\mu\text{mol}$ . Compared to the previous  $^{11}\text{C}$ -labeling performed using [ $^{11}\text{C}$ ]CO and selenium,<sup>10,27</sup> this procedure has the advantage to use a more readily accessible carbon-11 source (*i.e.* [ $^{11}\text{C}$ ]CO<sub>2</sub>) and avoid the manipulation of selenium derivatives. Additionally, the DRS was also successfully performed on Fenspiride.<sup>20</sup>



**Scheme 4.** Top: disconnection/reconnection strategy labeling of carbamates with carbon dioxide. Conditions: i) KOH, EtOH/H<sub>2</sub>O (4:1), 100 °C, ii) TfN<sub>3</sub>, DCM, CuSO<sub>4</sub>·5H<sub>2</sub>O, K<sub>2</sub>CO<sub>3</sub>, MeOH, overnight, r.t. (see SI for details). Bottom: labeling of linear carbamates.

Finally, we looked to extend the methodology to linear carbamates. Although metabolically more labile and generally used as prodrugs, linear carbamates have been labeled with <sup>11</sup>C with more or less success and usually under harsh conditions. For a first proof of concept, we selected substrate **32** that proved highly challenging with previous methods.<sup>13c,28</sup> After some optimization of the intermolecular reaction, [<sup>13</sup>C]**32** was labeled in 64% yield, after heating the mixture for 15 min at 150 °C. For the application to <sup>11</sup>C, the product was observed only in 4% d.c. RCC. This result clearly highlights the challenges related to the inherent physical and technical differences between carbon isotopes. Thought preliminary, this work shows the possibility to reach challenging linear carbamate systems.

## Conclusions

To conclude, a general methodology for the radiolabeling of cyclic aliphatic and aromatic carbamates has been developed. The reaction takes place with controlled amounts of CO<sub>2</sub>, the first available building block for <sup>14</sup>C and <sup>11</sup>C radioisotopes, resulting in a late-stage carbon isotope labeling of carbamate-containing drugs and analogues. Additionally, a disconnection/reconnection strategy was successfully implemented thus simplifying the synthesis of precursor and accelerating the whole labeling process. Finally, a proof of concept was obtained with a more challenging linear carbamate.

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgments

This project has received funding from the European Union's Horizon 2020 research and innovation program under the Marie Skłodowska-Curie grant agreement N°675071 and the CEA. The authors thank David-Alexandre Buisson, Amélie Goudet, Sabrina Lebrequier and Elodie Marcon for the excellent analytical support and Dr. Bertrand Kuhnast for helpful discussion.

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