11C02 Fixation: A Renaissance in PET Radiochemistry

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FEATURE ARTICLE
Neil Vasdev et al.
$^{11}$CO$_2$ fixation: a renaissance in PET radiochemistry
Carbon-11 labelled carbon dioxide is the cyclotron-generated feedstock reagent for most positron emission tomography (PET) tracers using this radionuclide. Most carbon-11 labels, however, are installed using derivative reagents generated from $^{11}\text{C}\text{O}_2$. In recent years, $^{11}\text{C}\text{O}_2$ has seen a revival in applications for the direct incorporation of carbon-11 into functional groups such as ureas, carbanates, oxazolidinones, carboxylic acids, esters, and amides. This review summarizes classical $^{11}\text{C}\text{O}_2$ fixation strategies using organometallic reagents and then focuses on newly developed methods that employ strong organic bases to reversibly capture $^{11}\text{C}\text{O}_2$ into solution, thereby enabling highly functionalized labelled compounds to be prepared. Labelled compounds and radiopharmaceuticals that have been translated to the clinic are highlighted.

1. Introduction

Carbon nuclei make up all organic entities and are the most versatile sites for labelling biological molecules of interest without altering their chemical and/or biological profiles. Carbon-11, a nearly pure positron emitter ($t_{1/2} = 20.38$ min, $E_{\text{avg}}(\beta^+) = 0.39$ MeV) is used extensively for developing radiotracers for PET, a non-invasive molecular imaging technique. The development of PET radiopharmaceuticals may provide an ideal methodology to enable diagnosis, monitor disease progression, and evaluate drug...
therapies in vivo without eliciting a pharmacological response.\textsuperscript{1} In addition to serving as a clinical tool for disease diagnosis, PET is increasingly relevant for drug development as it can provide quantitative pharmacokinetic, biodistribution, and receptor occupancy data for a drug candidate. While the longer half-life of fluorine-18 ($t_{1/2} = 109.77$ min, $E_{\text{avg}}(P) = 0.25$ MeV) offers advantages for synthesis, longer imaging times and multi-centre trials, carbon-11 can more frequently be substituted into biological molecules without causing chemical alterations that could influence the outcomes of imaging studies. Furthermore, repeat imaging studies in the same subject over a short duration are possible with carbon-11 due to its shorter half-life.

Dozens of reagents have been employed for incorporation of no-carrier-added carbon-11 ($^{11}$C) nuclei in radiotracers.\textsuperscript{2} Still, the most common strategy used is methylation using $^{11}$C-methyl iodide or $^{11}$C-methyl triflate.\textsuperscript{3-4} Carbon-11 labelled carbon dioxide is the product of proton bombardment of nitrogen-14 in the presence of small amounts of oxygen by the $^{14}$N(p,$p$)$^{11}$C nuclear reaction. Most often, $^{11}$C CO$_2$ is transformed into more reactive species, such as the aforementioned methylating reagents, to facilitate radiolabelling. However, $^{11}$C CO$_2$ itself is an attractive starting material for radiochemists as it is produced directly from the cyclotron in high specific activity. Its use also promises access to high oxidation state functional groups such as carboxylic acids, amides, ureas, carbamates, oxazolidinones and their derivatives without resorting to redox manipulations during radiotracer synthesis. Alternatively, $^{11}$C phosgene (produced from $^{14}$C CO$_2$ (ref. 5 and 6) or $^{11}$C methane\textsuperscript{7-10}) has been used to prepare such functional groups, though it has seen limited adoption due to technical challenges required for its routine production and use.\textsuperscript{11} As described in a recent review, the high reactivity of $^{11}$C CO$_2$ has been exploited to generate $^{11}$C-labelled intermediates such as isocyanates and carbamoyl chlorides from amines, and chlorofromates from alcohols, en route to $^{11}$C-ureas, $^{11}$C-carbamates and alky $^{11}$C-carbonates.\textsuperscript{12} Carbon-11 labelled carbon monoxide, produced by reduction of $^{11}$C CO$_2$,\textsuperscript{13} has been more widely employed for preparation of amides, esters, ureas, carbamates and acids using transition metal or selenium-mediated reactions or photoinitiated radical methods.\textsuperscript{11} Research has focused on overcoming the low solubility of $^{11}$C CO by employing micro-autoclaves,\textsuperscript{14} sequestration reagents such as borane,\textsuperscript{15} microfluidics,\textsuperscript{16-18} or soluble Xe(g) carrier.\textsuperscript{19} This topic has been reviewed elsewhere.\textsuperscript{2,11,20}

The low chemical reactivity of carbon dioxide poses a challenge for direct incorporation into organic molecules. Carbon dioxide generally requires highly reactive nucleophiles or catalysts to effect covalent bond formation and typically large excesses of CO$_2$ are used in industrial scale processes with this feedstock.\textsuperscript{21} In contrast, $^{11}$C CO$_2$ is the limiting reagent (10–100 fold excess of precursor) when used in radiochemical transformations. The low amounts of $^{11}$C CO$_2$ in nitrogen carrier gas obtained from the cyclotron target (typically ~100 nmol) necessitate an efficient CO$_2$ trapping solution. Carbon-11 CO$_2$ reactions are further complicated by the presence of oxygen and byproduct nitric oxides in the target gas mixture that often require gas purification steps, particularly if transition metals or sensitive catalysts are to be used. Two predominant strategies exist for purification of $^{11}$C CO$_2$ from target gas. Cryogenic purification consists of depositing the mixture in a vessel cooled by liquid nitrogen. Non-condensable gases are thereby removed, while condensable impurities (such as NO$_2$ species) can be removed using chemical traps.\textsuperscript{22} A second approach is to immobilize the carbon dioxide on a solid material (e.g. molecular sieves such as CarboSpheres\textsuperscript{23}). After stripping off undesired impurities, the CO$_2$ is thermally released from the trap into the reactor. In contrast to $^{11}$C COCL$_2$ and $^{11}$C CO, no additional chemical steps involving elaborate apparatus are required for preparation of $^{11}$C CO$_2$. However, as with any gaseous reagents with short half-lives (e.g., $^{11}$C CO$_2$, $^{11}$C CO, $^{11}$C COCl$_2$, $^{11}$C HCN, $^{11}$C CH$_3$I, $^{11}$C CH$_3$OTf, and $^{19}$F F$_2$) apparatus must be well designed and constructed to allow efficient handling, and be leak-proof. Flow rates must also be appropriate for rapid gas trapping in small volume solutions. Any traps, additives or fixation bases should avoid contamination of the isotope with impurities that could jeopardize the incorporation reaction. Many CO$_2$ fixations proceed rapidly at room temperature and ambient pressure. The reaction set-ups are generally very simple, which contributes to good reproducibility of syntheses and high radiochemical yields relative to starting $^{11}$C CO$_2$ activity are achievable with high specific activity (3–6 Ci μmol$^{-1}$). CO$_2$ is amenable to transformations into commonly found functional groups, and compounds have been recently advanced for first-in-human trials using $^{11}$C CO$_2$ fixation (vide infra). The purpose of this review is to highlight prominent examples of $^{11}$C CO$_2$ fixation that have been used to expand the chemical scope of labelled compounds and radiopharmaceuticals.

2. $^{11}$C CO$_2$ fixation by basic organometallic reagents

2.1 Grignard reagents

Carbon-11 labelled CO$_2$ has been in use for preparation of labelled carboxylic acids since at least the 1940s (Scheme 1A).\textsuperscript{24} Radiolabelled amides have also been synthesized by $^{11}$C CO$_2$ fixation with Grignard reagents. Heating magnesium carboxylate intermediates in the presence of primary or secondary amines has been reported to produce the corresponding $^{11}$C carboxyamides.\textsuperscript{25} The product amides have also been subsequently reduced in the presence of sodium borohydride, giving access to $^{11}$C tertiary amines.\textsuperscript{26} More indirectly, carboxylation can be followed by activation using reagents such as thionyl chloride or phthaloyl dichloride to prepare active acylation intermediates such as $^{11}$C acetyl and $^{11}$C propionyl chloride.\textsuperscript{27-30} While Grignard reagents have proven useful in the synthesis of simple $^{11}$C carboxylic acids and a selection of their derivatives, their high reactivity inherently limits the potential scope of their applications and enforces requirements for careful handling procedures. Since Grignard reagents readily absorb CO$_2$ from the atmosphere and are moisture-sensitive they are ideally prepared fresh and manipulated under an inert atmosphere.
using anhydrous solvents to obtain high specific activities and reproducibility. The propensity for magnesium salts to precipitate from solution poses challenges for automated synthesis and can necessitate delicate and time-consuming filtration steps.

### 2.2 Organolithiums

In the presence of excess methyllithium $[^{11}\text{C}]{\text{CO}}_2$ is readily transformed to the dilithium salt of acetone acetate. Hydrolysis leads to $[2-^{11}\text{C}]$acetone, which is a useful intermediate for radiochemistry (Scheme 1B).\(^{31-33}\) Reports using other organolithiums to trap $[^{11}\text{C}]{\text{CO}}_2$ have been relatively scarce. $[^{11}\text{C}]$pyruvic acid was prepared by tandem methylation-carboxylation of an isocyanide in the presence of methyllithium and $[^{11}\text{C}]{\text{CO}}_2$, followed by hydrolysis.\(^{34}\) Also of note are the syntheses of radiolabelled glycine and two derivative dipeptides using the lithium salt of methylisocyanide. After $[^{11}\text{C}]{\text{CO}}_2$-fixation the isocyanide is hydrolyzed through the formamide to the amine, yielding $[1-^{11}\text{C}]$glycine (Scheme 1C).\(^{35}\) This species could be coupled to an intramolecularly activated amino acid equivalent under basic conditions.

### 2.3 Silanamines

Silylated amines were developed in the 1980s as fixating agents for $[^{11}\text{C}]{\text{CO}}_2$. This work was performed by Ram and co-workers and was applied to various targets including imipramine,\(^{36}\) chlorpromazine,\(^{37}\) SCH-23390,\(^{38}\) and tamoxifen.\(^{39}\) Silanamines were prepared up to one week in advance of radiosynthesis by refluxing a secondary amine with hexamethyldisilazane (HMDS), often in the presence of catalytic quantities of ammonium sulfate. When exposed to $[^{11}\text{C}]{\text{CO}}_2$ at elevated temperatures, O-silyl carbamates were produced, which were subsequently reduced in situ by LiAlH$_4$ to produce labelled tertiary methylamines (Scheme 1D).\(^{40}\) Since silyl groups are electron-donating, the silanamine is rendered more nucleophilic, despite the added steric bulk and weak N–Si bond strength.

The above methods all require the use of an unstable organometallic/organosilicon species for $[^{11}\text{C}]{\text{CO}}_2$ fixation. These reagents are often quite reactive and pose obstacles to reproducibility and automation in a radiochemical setting. Naturally, they also present major chemoselectivity issues during the preparation of PET radiotracers with complex functionalities, often limiting their utility to synthesis of simple prosthetic groups or very small molecules with restricted functionality. These approaches have been previously reviewed\(^{41}\) and stand in contrast to the recent developments described below.

### 3. Recently developed methods of $[^{11}\text{C}]{\text{CO}}_2$ fixation

Over the past 20 years, and under the impetus of “green” chemistry, approaches to CO$_2$ fixation have expanded tremendously.\(^{42}\) Among the most commonly used fixation agents are guanidines,\(^{43}\) amidines,\(^{44,45}\) and alkali carbonates,\(^{46}\) the last of which are not suitable for $[^{11}\text{C}]{\text{CO}}_2$ fixation. Though each of these bases facilitates transamidation of an added amine, the nature of
products of CO₂ trapped by guanidines or amidines have been the subject of some debate. Initial spectroscopic and crystallographic studies suggested that bicarbonate salts are formed in the presence of adventitious water, perhaps by hydrolysis of carbamic intermediates.43,47–49 More recently, the carbamate anion has been confirmed crystallographically in the context of a bicyclic guanidine.50 Practically, Hooker et al. demonstrated that an amidine, diazabicyclo[5.4.0]undec-7-ene (DBU), was highly efficient at trapping cyclotron-produced [¹¹C]CO₂ at ambient temperature and pressure and practical flow rates.51 Wilson et al. later showed that 2-tert-butylimino-2-diethylamino-1,3-dimethyl-perhydro-1,3,2-diazaphosphorine (BEMP) was even more effective in this regard.52 This strategy succeeds by decoupling the [¹¹C]CO₂ capture in solution from subsequent covalent bond formation with substrate.

Transition metals have been extensively used for both CO₂ trapping in solution and as fixation catalysts for carbon–carbon bond formation.53,54 The mode and efficiency of trapping are likely to be dependent on the nature of the metal centre, as well as the ligand and solvent systems. These topics have been the subjects of reviews in recent years.55,56

4. Functional groups prepared by [¹¹C]CO₂ fixation

4.1 Urea

4.1.1 Symmetrical ureas. While early examples of [¹¹C]CO₂ fixation centred around formation of carboxylic acids and amides, higher oxidation state functional groups such as ureas and carbamates have become research targets in the past 20 years. Chakraborty et al. reported the first synthesis of [¹¹C]urea by bubbling [¹¹C]CO₂ through a THF solution of LHMDS, followed by hydrolysis using aqueous ammonium chloride (Scheme 2A).57 [¹¹C]urea could then be condensed with diethyl malate in sulfuric acid to yield [¹¹C]uracil.

The synthesis of both ureas and carbamates from carbon dioxide often requires CO₂ to react twice as an electrophile. While we have seen above that in the presence of strong nucleophiles carbon dioxide will react as an electrophile, the resulting carboxylate or carbamate is not electrophilic. It is for this reason that [¹¹C]phosgene has previously been used to prepare labelled ureas.12 An alternative approach is to utilize activating reagents such as phosphoryl chloride or thionyl chloride to generate an acid chloride or isocyanate, which are both highly electrophilic. In its earliest iterations, this strategy was applied to prepare [¹¹C-carbonyl]phenyl isocyanate, which promptly reacted further to form both [¹¹C]diphenyl urea and [¹¹C]diphenyl carbodiimide (Scheme 2B).58 While observation of these products is clear evidence of the desired reactivity, they also illustrate the practical challenges in harnessing it for preparation of unsymmetrical ureas in high radiochemical yield.

4.1.2 Unsymmetrical ureas. By using dilute and carefully balanced solutions of trapping amine, POCl₃, and nucleophile, unsymmetrical [¹¹C]ureas can be produced with high selectivity (Scheme 2B).59 Simply using a large excess of POCl₃ was found to suppress attack of remaining trapping amine on the [¹¹C]-isocyanate generated in situ, but also required even larger excess of the nucleophile, which is also consumed. This results

Scheme 2 Synthesis of [¹¹C]ureas by [¹¹C]CO₂-fixation. (A) Synthesis of parent [¹¹C]urea using LHMDS and aqueous ammonium chloride. (B) Synthesis of symmetrical and unsymmetrical substituted [¹¹C]ureas by activation of a carbamate intermediate with POCl₃. The utility of this reaction was greatly expanded by judicious control of reaction conditions to favour unsymmetrical products. TEA: triethylamine; TMS: trimethylsilyl; asterisk denotes [¹¹C].
in a high concentration reaction mixture and consequent challenging purification. Fortunately, it was found that the concentration of trapping amine could be reduced without negatively impacting the yield or prolonging the reaction time beyond two minutes. Aliphatic primary amines are ideal substrates for isocyanate formation, while aniline reacts sluggishly. Cyclic secondary amines are also well tolerated, presumably through an $^{11}$C-carbamoyl chloride intermediate, though more hindered secondary amines must be used in higher concentrations to achieve useful conversions. The scope of amine nucleophiles for attack on $^{14}$Csocyanates or $^{11}$C-carbamoyl chlorides was limited primarily to dimethylamine for the discovery investigations, but high yields were also obtained with 4-(2-methoxyphenyl)piperazine.

Inhibitors of glycogen synthase kinase 3β (GSK-3β) are currently under exploration for diverse cancers and neurological illnesses, with the “GSK-3 hypothesis of Alzheimer’s disease” sparking further medicinal chemistry efforts in this area. Vasdev et al. prepared the first PET imaging agent for this target, $^{11}$C-methoxy$\text{AR-A014418}$. The room temperature synthesis of an isotopologue of this unsymmetrical urea, $^{11}$C-carbonyl$\text{AR-A014418}$, demonstrates a $^{11}$C$\text{CO}_2$ fixation approach to this tracer. In this case the terminal nucleophile was the aromatic 2-amino-5-nitrothiazole. This is the first successful example of aromatic amines being employed in this role. The total time for synthesis and formulation was 28 minutes, and 70 ± 33 mCi of the tracer could be prepared in 8.3 ± 3.9% uncorrected radiochemical yield (RCY) relative to starting $^{11}$CO$_2$ with high (4.0 ± 1.1 Ci μmol$^{-1}$) specific activity at the end of synthesis (Fig. 1). Though AR-A014418 was found to be less potent than initially reported, this methodology affords a general approach to synthesize arrays of $^{11}$C-labelled urea-based GSK-3β inhibitors, which would not be possible using standard $^{11}$C-methylation strategies.

Pfizer’s potent and irreversible fatty acid amide hydrolase (FAAH) inhibitor, PF-04457845, has advanced to clinical trials and has generated recent interest as a scaffold for PET radiotracer development. Whereas a $^{18}$F-fluoroethyl derivative, $^{18}$FPF-9811, was recently reported, the isotopologue $^{11}$C-carbonyl$\text{PF-04457845}$ was prepared by $^{11}$C$\text{CO}_2$ fixation (Fig. 1). The precursors for $^{11}$C$\text{CO}_2$ fixation are a non-nucleophilic primary aromatic amine and a cyclic secondary aliphatic amine. To compensate for the higher reactivity of the latter compound, the aromatic amine was employed in 20-fold excess relative to the secondary amine. $^{11}$CO$_2$ was bubbled into the vial containing the precursors and BEMP in anhydrous CH$_3$CN. POCl$_3$ was later added, followed by an aqueous quench and purification to provide $^{11}$C-carbonyl$\text{PF-04457845}$ in 4.5 ± 1.3% uncorrected radiochemical yield (Fig. 1). The total synthesis time was 25 ± 2 min and the product had a specific activity of 2.0 ± 0.2 Ci μmol$^{-1}$. All operations were performed at room temperature. In contrast to the preparation of $^{11}$C$\text{AR-A014418}$, both amines were present in the reaction mixture during $^{11}$C$\text{CO}_2$ fixation. Presumably, the reactive intermediate is a $^{11}$C-carbamoyl chloride or a mixed phosphate $^{11}$C-anhydride. Promising preclinical results, coupled with the known pharmacology and toxicology of PF-04457845, should facilitate clinical translation of this radiotracer.

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**Fig. 1** Cyclotron produced $^{11}$CO$_2$ has been recently applied to radiolabelling of structurally complex carboxylic acids and amides, carbamates, oxazolidinones, and ureas. *Trapping is typically done with molecular sieves or liquid nitrogen and release is achieved by heating.*
An alternative approach to $^{11}$C-isocyanates is by the condensation of phosphinimines with $^{11}$C$\mathrm{CO}_2$. Phosphinimines can be prepared from azides or primary amines, and have varying degrees of stability. Commercially available phenyltriphenylphosphinimine was used for $^{11}$C$\text{urea}$ preparation with aliphatic and aromatic amines. 65 The radionuclide was trapped in a solution of THF at $-60\,^\circ\text{C}$ in the presence of the phosphinimine and amine. The reaction was heated to $60\,^\circ\text{C}$ for 6 minutes to complete the synthesis (Scheme 3). By radio-HPLC, aliphatic amines gave decay corrected unpurified radiochemical conversion yields ranging from 45–49%, while aniline unsurprisingly gave a lower yield of $8\pm1\%$.

4.2 Carbamate

4.2.1 Syntheses using alkyl electrophiles. Similar to ureas, carbamates are an attractive functional group for radiolabelling due to their stability in vivo and role as a versatile linker for ligand fragments. This functional group has previously been labelled using $^{11}$C$\text{phosgene}$ and $^{11}$C$\text{carbon monoxide}$, 11 so a synthetic strategy based around $^{11}$C$\text{CO}_2$ would have extensive applications for the development of PET radiopharmaceuticals. In contrast to ureas, syntheses of $O$-alkyl carbamates do not necessarily require activation of the carbamate salt intermediate if reactive electrophiles are used. This strategy was exemplified by employing benzyl chloride in a solution of DBU, benzylamine, DMF and bubbled $^{11}$C$\text{CO}_2$ (Scheme 4). 51 The total reaction time was 10 minutes at $75\,^\circ\text{C}$ and achieved a radiochemical yield of $85\%$. Similar to some of the urea results discussed above, reducing the concentration of benzylamine in this reaction was better tolerated than reducing the concentrations of other reagents. Secondary and sterically hindered amines gave good yields, while anilines were much less reactive. The best electrophiles were benzyl and allyl chlorides since analogous bromides displayed undesired reactivity with amines and DBU. n-Pentyl bromide showed desired reactivity, while more hindered aliphatic electrophiles were troublesome. This technology was used to prepare $^{11}$C$\text{metergoline}$ in $32\%$ decay corrected RCY in high specific activity (up to $5\,\text{Ci}\,\mu\text{mol}^{-1}$) in a single step. $^{11}$C$\text{MS-275}$, an isotopologue of the histone deacetylase inhibitor entinostat, was also prepared in this way and showed poor blood–brain barrier penetration in non-human primates. 66 The tracer was synthesized in $25\%$ decay corrected RCY with specific activities of $2.7–6.2\,\text{Ci}\,\mu\text{mol}^{-1}$.

Synthesis and purification required 50 min and afforded $>20\,\text{mCi}$ per synthesis.

Methyl $^{11}$C-carbonyl]carbamates have also been prepared following a similar strategy with methylating agents such as dimethylsulfate (DMS), methyl iodide (CH$_3$I), or methyl tosylate (CH$_3$OTs) (Scheme 4). 52 In this case, BEMP in DMF was found to be the best trapping solution for $^{11}$C$\text{CO}_2$. The reactions proceeded with primary or secondary amines. Electron rich 4-methoxyaniline was also tolerated, while acceptable yields of electron deficient 4-nitroaniline could be achieved with increased amine concentration. The reactions also proceeded very rapidly, with maximum yields being reached using one minute of mixing prior to addition of the methylating agent and only 10 seconds afterwards. Increasing the concentration of methylating agent had a negative effect on radiochemical yield. Reversing the order of addition (i.e., first DMS, followed by amine) reportedly produced very low RCYs of product, suggesting the intermediacy of the carbamate salt.

The utility of this process was demonstrated by labelling GR103545, a selective high affinity agonist for the $\kappa$-opioid receptor, by $^{11}$C$\text{CO}_2$ fixation. The methyl carbamate has previously been labelled using $^{11}$C$\text{methylchloroformate}$, 67 phosgene with $^{11}$C$\text{CH}_3\text{OH}$, 68 and either $^{11}$C$\text{CH}_3\text{I}$69 or $^{11}$C$\text{CH}_3\text{OTf}$70 with cold CO$_2$ fixation. In this iteration, the “loop” method was employed by lining the inside of a narrow-bore steel tube with the precursor and BEMP in DMF. Without requiring heating, 70–103 mCi of 2.9–4.4 Ci $\mu$mol$^{-1}$ $^{11}$C-carbonyl]GR103545.

![Scheme 3](image)

Scheme 3 Synthesis of $^{11}$C$\text{phenylisocyanate}$ and $^{11}$C$\text{urea}$ by fixation with $N$-phenyl(triphenylphosphin)imine. Asterisk denotes $^{11}$C.

![Scheme 4](image)

Scheme 4 $^{11}$C-carbonyl]carbamates can be synthesized by nucleophilic alkylation of the trapped $^{11}$C$\text{CO}_2$ followed by amine substitution (left), or through activation of an intermediate carbamate salt using POCl$_3$ (right). DBU: 1,8-diazabicyclo[5.4.0]undec-7-ene; Q: fixation base; R: H, phenyl, vinyl, alkyl; X: Cl, Br, I, OTs, sulfate; asterisk denotes $^{11}$C.
(Fig. 1) was produced using 0.1 mg of precursor, an uncorrected
13% RCY at end-of-synthesis (EOS).\textsuperscript{52} The total synthesis time
was 23 minutes.

4.2.2 Syntheses via isocyanates. [\textsuperscript{11}C-carbonyl]carbamates
have also been produced through [\textsuperscript{11}C]isocyanate intermediates.\textsuperscript{59}
Again, appropriate stoichiometry must be selected to prevent
symmetrical [\textsuperscript{11}C]urea formation. The isocyanate is quenched
with an alcohol to affix the desired O-substituent. While methanol
and phenols were originally deployed, in our most recent work we
have also successfully used ethanol, isopropanol, and even tert-
butanol and hexafluoropropanol. It is possible that the reaction
is assisted by the strongly basic conditions established by the
presence of BEMP.

The synthesis and function of [\textsuperscript{11}C]CURB (Scheme 5A) is
illustrative of the importance of this labelling strategy. FAAH is
a serine hydrolase that regulates levels of anandamide, an
endocannabinoid, in the central nervous system. FAAH is
found heterogeneously throughout the brain, and is an attrac-
tive target in studies of addiction, obesity, pain, anxiety, and
eating disorders.\textsuperscript{71–73} The O-aryl carbamate scaffold has shown
to be an effective one for design of irreversible inhibitors of
FAAH, as the serine-241 residue in the active site has been
shown to attack the carbonyl of the carbamate, releasing a
phenolic fragment (Scheme 5B). With the potential for
structure–activity relationship (SAR) studies in mind, it can
be appreciated that the conserved carbamate carbonyl is an
ideal position for placing a radiolabel.

[\textsuperscript{11}C]CURB was prepared by [\textsuperscript{11}C]CO\textsubscript{2} fixation using cyclo-
hexylamine and BEMP, followed by [\textsuperscript{11}C]isocyanate formation with POCl\textsubscript{3}.\textsuperscript{59} 2-Phenyldihydroquinone was used to quench the
isocyanate, forming a mixture of regioisomers which are sepa-
rated by HPLC. The radiotracer was isolated in 8% uncorrected
RCY with a specific activity of 2.5 Ci m mol\textsuperscript{−1} in 27 minutes from
end-of-bombardment (EOB). [\textsuperscript{11}C]CURB shows high brain pene-
tration in rats, and selectivity was ascertained by blocking studies
with a known FAAH inhibitor.\textsuperscript{74} The radiotracer has been trans-
lated to human use.\textsuperscript{75} To optimize radiotracer pharmacokinetics,
an SAR study was performed using a small library of O-aryl
carbamates. Eight [\textsuperscript{11}C]carbamates were prepared in a manner
analogous to [\textsuperscript{11}C]CURB.\textsuperscript{76} Each of the radiotracers demon-
strated brain uptake and specificity for FAAH in conscious
rodents. Kinetic analysis in rats showed that [\textsuperscript{11}C]dihydrooxazole
 carbamates had greater brain uptake, lower non-specific binding,
and faster binding to FAAH than the [\textsuperscript{11}C]biphenyl carbamates,
such as [\textsuperscript{11}C]CURB. The results of the SAR study and the human
kinetic studies using [\textsuperscript{11}C]CURB will allow for design and selection
of the optimal FAAH radiotracer.

A camptothecin derivative that displays potent antitumor
activity, irinotecan, was radiolabelled with carbon-11.\textsuperscript{77} Kawamura
\textit{et al.} prepared [\textsuperscript{11}C]irinotecan both by direct [\textsuperscript{11}C]CO\textsubscript{2} fixation and
using [\textsuperscript{11}C]COCl\textsubscript{2} derived from [\textsuperscript{11}C]CO\textsubscript{2} by way of [\textsuperscript{11}C]CH\textsubscript{4} and
[\textsuperscript{11}C]CCl\textsubscript{4}. Using [\textsuperscript{11}C]CO\textsubscript{2} directly, the decay corrected RCY
was 16.9 ± 2.9% with specific activity 35 min from EOB of 2.1–3.7 Ci μmol\textsuperscript{−1}. Using [\textsuperscript{11}C]COCl\textsubscript{2}, the decay corrected RCY
was 8.8 ± 2.0% with specific activity 35 min from EOB of 2.1–5.3 Ci μmol\textsuperscript{−1}. The tracer was used to perform metabolite
analysis of the drug in mice.

4.3 Oxazolidinone
[\textsuperscript{11}C]SL25.1188 (Fig. 1) was developed as a reversibly binding
radiotracer for monoamine oxidase-B (MAO-B). The radiolabel
is placed as a [\textsuperscript{11}C-carbonyl]oxazolidinone. The original synthesis
employed [\textsuperscript{11}C]phosgene, and suffered from a low 3.5–7%
decay-corrected RCY with 1.4–1.9 Ci μmol\textsuperscript{−1} specific activity
after a 30–32 minutes.\textsuperscript{78} The application of the amino-alcohol
precursor for [\textsuperscript{11}C]CO\textsubscript{2} fixation followed by dehydration and
intramolecular cyclization improves the radiosynthesis.\textsuperscript{79} After
optimization of the fixation base, dehydrating agent, and
component concentrations, conditions were developed to
perform the labelling at ambient temperature using BEMP
and POCl\textsubscript{3}. The uncorrected RCY was 11.5 ± 0.9% to
prepare 98 ± 8 mCi of the radiotracer with 1.0 ± 0.05 Ci μmol\textsuperscript{−1}
specific activity after a 30 min synthesis. This tracer has
since been successfully translated for human use and specific
activities of 3.7 ± 0.8 Ci μmol⁻¹ have been achieved by us (unpublished work).

4.4 Carboxylic acid

Transition metal-mediated carboxylation of organic compounds has seen significant development in part due to the reputation of CO₂ as an environmentally benign feedstock. In particular, catalytic and even metal-free conditions using prefunctionalized organoboron and organozinc reagents have been the focus of much attention.⁴⁰⁻⁴² The advantages these reagents hold over Grignard and organolithium compounds are their functional group tolerance and relative stability to storage under ambient conditions. Boronic esters have recently been applied for [¹¹C]CO₂ fixation using a copper catalyst.⁸³ Due to low concentration of [¹¹C]CO₂ available in the reaction mixture, which is in sharp contrast to “cold” (non-radioactive) CO₂ fixation conditions, the reaction parameters required significant optimization efforts. Since alkoxide bases bound [¹¹C]CO₂ tightly, TMEDA was employed as both fixation base and presumably a ligand for the copper catalyst. A soluble fluoride additive was also found to dramatically improve RCY. With optimized conditions at 90–100 °C, various arylboronic esters were efficiently converted to [¹¹C]carboxylic acids (Scheme 6). Halide, formyl, cyano, and nitro substituents were well tolerated, while protic substituents such as hydroxyl and amino groups, and electron poor heterocycles were challenging substrates. Examples of alkyl, vinyl and alkynyl substrates undergoing [¹¹C]carboxylation were also reported.

The [¹¹C]benzoic acids were also elaborated to a methyl ester (using iodomethane), an amide (via an acid chloride), and a succinimide ester (via carbodiimide activation).⁸⁴ An [¹¹C]amide-labelled oxytocin receptor ligand⁸⁴ was also reported in 20% decay-corrected RCY with 1.5 Ci μmol⁻¹ specific activity over 43 minutes synthesis time. Access to various [¹¹C]carboxylic acid derivatives through regiospecific metal catalyzed carboxylation is sure to be widely exploited.

5. Conclusion

The last few years have seen considerable enthusiasm for radiosynthesis using [¹¹C]CO₂. Whereas classical [¹¹C]CO₂ incorporation has relied on highly basic organometallic reagents, modern strategies employ more functional group tolerant reagents and catalysts for activation. This has enabled the synthesis of structurally complex radiopharmaceuticals and precise placement of the carbon-11 label within functional groups such as ureas, carbamates, and oxazolidinones under very mild conditions, as well as carboxylic acids, esters and amides. Most recently, the carbamate radiotracer [¹¹C]CURB and the oxazolidinone, [¹¹C]SL25.1188 have advanced for human neuroimaging studies of FAAH and MAO-B, respectively. We anticipate that future developments in this field will establish mild and robust strategies for labelling a greater number of functional positions, including amides. Enabling technologies such as microfluidics and automated apparatus should prove to be valuable for rapid method optimization and widespread adoption of [¹¹C]CO₂ fixation for clinical translation.

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References
