Development of [18F]AldoView as the First Highly Selective Aldosterone Synthase PET Tracer for Imaging of Primary Hyperaldosteronism

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ABSTRACT: The purpose of this study was to synthesize a fluorine-18 labeled, highly selective aldosterone synthase (hCYP11B2) inhibitor, [18F]AldoView, and to assess its potential for the detection of aldosterone-producing adenomas (APAs) with positron emission tomography in patients with primary hyperaldosteronism (PHA). Using dibenzothiophene sulfonium salt chemistry, [18F]AldoView was obtained in high radiochemical yield in one step from [18F]fluoride. In mice, the tracer showed a favorable pharmacokinetic profile, including rapid distribution and clearance. Imaging in the adrenal tissue from patients with PHA revealed diffuse binding patterns in the adrenal cortex, avid binding in some adenomas, and “hot spots” consistent with aldosterone-producing cell clusters. The binding pattern was in good visual agreement with the antibody staining of hCYP11B2 and distinguished areas with normal and excessive hCYP11B2 expression. Taken together, [18F]AldoView is a promising tracer for the detection of APAs in patients with PHA.

INTRODUCTION

Hypertension is a leading cause of premature morbidity and death globally. Although lifestyle changes and medication can be effective, elevated blood pressure due to secondary causes is more difficult to control and has greater morbidity. This is particularly true for primary hyperaldosteronism (PHA), which is characterized by excessive, autonomous aldosterone production by the adrenals. PHA occurs in 5–10% of patients with hypertension and in 15–25% of those with treatment-resistant hypertension. Although PHA is a potentially curable cause of secondary hypertension, it is estimated that >99% of those with PHA remain undiagnosed and on lifelong medication at a high cost to individuals and healthcare budgets. In most cases, PHA is due to either bilateral hyperplasia of the adrenal cortex or an aldosterone-producing adenoma (APA). For patients with a demonstrable unilateral cause, surgical removal of the abnormal gland (adrenalectomy) often results in reduced blood pressure, lower dependence on antihypertensive drugs, and sometimes in the complete cure of hypertension (30–60% of cases). The challenge is how to identify such patients.

Many patients who are >50 years old have adenomas as incidental findings; yet, these are not necessarily a source of excess aldosterone production. Conversely, some APAs are smaller than the limit of detection for computed tomography (CT) or magnetic resonance imaging (MRI) and may be contralateral to an adenoma. This recognition of frequent microadenomas (diameter <1 cm) requires techniques to detect minute functional lesions, in order to identify a surgical opportunity to resect the affected adrenal on one side, and not miss a cause of persistent PHA postadrenalectomy on the other. To avoid removing the wrong adrenal, the current diagnostic pathway to lateralize patients with PHA involves the following: biochemical tests that suggest autonomous, excess aldosterone production, followed by the identification of an adrenal nodule consistent with an adenoma by CT or MRI, followed by adrenal vein sampling (AVS) to assess whether the adrenal nodule is a source of excess aldosterone production. However, AVS is technically challenging, invasive, not widely available, and is often not feasible as it requires patients with severe hypertension to adjust or stop medication for several weeks. This complex diagnostic route is the major reason why so few patients ultimately undergo adrenalectomy, a potentially curative procedure for their hypertension.

Inappropriately high aldosterone production in patients with PHA is due to the increased aldosterone synthase (CYP11B2) activity. CYP11B2 is the only known source of aldosterone production, and a large body of evidence shows overexpression of this enzyme in adrenal tissue from patients with PHA (often >10–100 fold). For these reasons, evidence of abnormal
CYP11B2 expression in adrenal lesions, or the surrounding adrenal cortex, is considered both necessary and sufficient for the histopathological diagnosis of PHA. Imaging of CYP11B2 with positron emission tomography (PET) is an appealing alternative to the aforementioned diagnostic pathway as it could enable the identification and lateralization of adrenal glands with areas of excessive aldosterone production and reveal the functional status of adrenal nodules detected with CT or MRI. However, selective in vivo imaging of CYP11B2 has not yet been achieved due to the close homology between the enzymes involved in aldosterone (CYP11B2) and cortisol (CYP11B1) synthesis. This is a major limitation as CYP11B1 is highly expressed in healthy adrenals and some adenomas, without any apparent correlation with the rate of aldosterone excretion.

Recently, Merck & Co. reported a class of highly selective aldosterone synthase inhibitors, derived from benzimidazole. Using a dibenzothiophene sulfonium salt—a new fluorine-18 chemistry platform developed by our group—we labeled the benchmark benzimidazole ligand to give the first highly selective CYP11B2 PET tracer (aka [18F]AldoView; IC50 4.7 nM vs 435 nM for CYP11B1). Herein, we describe the synthesis of [18F]AldoView and its characterization in preclinical studies.

**RESULTS**

**Chemistry.** The nonradioactive benchmark compound 3 was synthesized following previously reported procedures (Scheme 1). The [18F]AldoView precursor for labeling with fluorine-18 was prepared in three steps starting from the diamine 2. Condensation of 2 with 5-bromonicotinaldehyde in the presence of oxone gave the bromo-pyridine 4 in 73% yield. Subsequent Suzuki cross-coupling of 4 with the protected biaryl 5 (prepared as described previously) provided the thioether 6 in 94% yield. Ring closure of 6 initiated by calcium hypochlorite led to the formation of the dibenzothiophene sulfonium salt 7. The purity of the labeling precursor 7 was >97% as determined by high-performance liquid chromatography (HPLC) (Figure S1).

**Radiochemistry.** The reaction of the dibenzothiophene sulfonium salt 7 with [18F]fluoride was carried out using a low precursor load of 2 mg per reaction. The reaction conditions were designed to suit automated radiosynthesis following good manufacturing practice (Scheme 2). [18F]AldoView was obtained in 42 ± 8% (decay-corrected; n = 12) radiochemical yield, with a radiochemical purity of >99%, and with a molar activity of 31 ± 3 GBq/μmol (n = 4) when starting from 2 GBq of [18F]fluoride (Figure S2).

**PET/CT Imaging.** Whole-body imaging of [18F]AldoView in female BALB/c mice (n = 6) showed a marked liver uptake at the earlier time points (2−5 and 5−15 min, summation images, Figure 1A). At 15−30 min, the distribution was dominated by the activity in the bladder, and at the later time point (30−60 min), uptake in the bone tissue became evident. Dynamic analysis (Figure 1B) showed that the tracer uptake in major organs such as the brain, lungs, and heart peaked at 1−2 min postinjection and then rapidly decreased. In muscle (region of interest across knee extensors), the level of activity remained low (<1.5% ID/g) throughout the duration of the scan (1 h). The kidney time−activity curve appeared to be biphasic, indicating initial perfusion at 0−5 min postinjection, followed by low and constant radioactivity levels of 5−8% ID/g thereafter. By far the highest level of activity was observed in...
the liver, which peaked at 6 min postinjection with $48 \pm 11\% \text{ID/g}$ and subsequently decreased to half the peak value within 30 min. At later time points, the bone surpassed the liver as the tissue with the highest levels of activity. No dynamic data were recorded for the adrenals due to the minuscule size of these organs in mice.

**Biodistribution.** [18F]AldoView was administered by tail vein injection in healthy female BALB/c mice ($n = 3-4$ per time point), and the distribution of radioactivity was measured in selected organs at 15, 30, 60, and 120 min postinjection (Figure 1C). At the 15 min time point, the highest activity levels were observed in the liver ($11.8 \pm 4.1\% \text{ID/g}$), followed by the adrenals ($6.1\% \text{ID/g}$, $n = 2$), kidneys ($5.2 \pm 3.5\% \text{ID/g}$), and small intestines ($4.3 \pm 4.7\% \text{ID/g}$). The activity levels in blood and the other organs investigated (stomach, spleen, heart, lungs, brain, and bone) were comparatively low. In the adrenals, the activity levels gradually decreased over time ($4.0 \pm 2.5\% \text{ID/g}$ at 30 min to $1.8 \pm 1.1\% \text{ID/g}$ at 60 min). The uptake in the kidneys and liver peaked at 30 min ($5.7 \pm 2.9$ and $12.4 \pm 5.9\% \text{ID/g}$, respectively) and then cleared, more rapidly so from the kidneys than from the liver. The ratio between the tracer uptake in the adrenals and kidneys was 1.2 at 15 min postinjection and subsequently equilibrated at 0.6–0.7 (inset in Figure 1C). The activity levels in the intestines remained practically unchanged over the course of the experiment ($4\text{--}6\% \text{ID/g}$). In bone, the activity levels increased from 15 to 60 min, suggesting that the tracer may undergo metabolic defluorination in vivo.

**Metabolite Analysis.** To assess the in vivo stability of [18F]AldoView, the metabolic profile in blood was determined using radio-HPLC (Figure 1D). At 15, 30, and 60 min postinjection, the intact parent fractions of the tracer were 59, 29, and 21%, respectively. The tracer was still detectable in blood at 120 min, although at low levels. A single, highly polar metabolite accounted for the remaining activity throughout the course of the experiment.

**Human Tissue Imaging.** The ability of [18F]AldoView to depict CYP11B2 expression was assessed by quantitative phosphorimaging in human tissue sections from surgically resected adrenal glands from five patients diagnosed with PHA. Tissue sections from three patients with CYP11B2-negative adrenal lesions were included as controls. Where feasible, several surgical specimens were collected from each patient to include the respective lesions (adenoma/tumor), as well as the adrenal cortex. The expression patterns of CYP11B2 were confirmed independently using immunohistochemical (IHC) staining in adjacent tissue sections. The binding of [18F]AldoView (Figures 2, S3) was visually consistent with the IHC staining of CYP11B2 and demarcated areas with dense expression in some of the larger adenomas (e.g., specimens 2b/c in Figure 2), adrenal cortex (e.g., specimen 3b in Figure 2), and aldosterone-producing cell clusters (APCCs) (e.g., specimen 1b in Figure 2). Specific tracer binding in these CYP11B2-positive areas ranged from 8.6 to 19.1 kBq/cm². In specimens designated as APAs upon postsurgical visual examination (characterized by a typical golden color and dense morphology of the tissue, e.g., specimens 1a, 2a, and 3a...
in Figure 2), tracer binding ranged from 2.5 to 19.7 kBq/cm² (n = 6). Notably, tracer binding was evenly distributed across APA tissue sections, in contrast to adrenal cortex sections (e.g., specimens 1b and 3b in Figure 2), for which different expression patterns and “hot spots” were seen. There was no evidence of elevated tracer uptake in CYP11B2-negative specimens. In contrast, tracer binding was consistently low in specimens from control subjects (2.6 ± 1.8 kBq/cm²; n = 3) and in CYP11B2-negative areas in specimens from patients with PHA (3.2 ± 1.1 kBq/cm²; n = 4).

**DISCUSSION**

PHA is a common type of secondary hypertension, which affects as much as 1% of the population, and causes significant morbidity and premature mortality due to both hypertension and the high levels of aldosterone. Although medical treatment can prevent some of the adverse effects of PHA, in patients with unilateral disease, surgery is often more effective in reducing the blood pressure, morbidity, and risk of mortality and gives a higher quality of life. A practical, scalable, and expeditious diagnosis will allow many more PHA patients to be identified for surgery and potentially cured. PET imaging of CYP11B2 expression in the adrenal glands can meet this need by identifying and lateralizing APAs noninvasively.

The parent compound of [18F]AldoView was reported as a benchmark for the Merck & Co. benzimidazole series of aldosterone synthase inhibitors due to its high potency and selectivity for human CYP11B2 (IC₅₀ 4.7 nM vs 435 nM for hCYP11B1), lack of “off target” binding, high metabolic stability, and favorable pharmacokinetic profile. The presence

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**Figure 2.** In vitro autoradiography with [18F]AldoView in tissue sections from surgically resected adrenal glands in comparison with CYP11B2-specific IHC staining in directly adjacent tissue sections. Case 1 is a female PHA patient aged 41 at the time of adrenalectomy, diagnosed with a 17 × 12 mm adenoma and otherwise normal parenchyma (as determined by CT imaging). Autoradiography revealed a high CYP11B2 expression in the APA (specimen 1a) and in APCCs located in the zona glomerulosa of the adrenal cortex (1b). The tracer binding pattern was in good visual agreement with CYP11B2-specific IHC. Case 2 is a male PHA patient aged 55, diagnosed with a 10 mm adenoma and hyperplasia. [18F]AldoView binding comparable to that found in the control tissue was observed in the suspected APA (specimen 2a), suggesting low CYP11B2 expression levels. CYP11B2 IHC was inconclusive due to a lack of contrast. Parts of specimen 2b were identified as APA following CYP11B2 IHC, and subsequent autoradiography revealed high enzyme expression levels. Specimen 2c showed a high CYP11B2 expression adjacent to the adrenal cortex, as determined with autoradiography. The tracer binding pattern was consistent with CYP11B2 IHC. Case 3 is a female PHA patient aged 37 at the time of adrenalectomy, diagnosed with a 9 mm adenoma. [18F]AldoView binding to the APA was found to be moderately increased (specimen 3a). An area in specimen 3b, adjacent to the adrenal cortex, revealed high CYP11B2 expression levels by [18F]AldoView autoradiography and CYP11B2 IHC. Scale bar: 1 cm. NB1: The different shades of brown in the CYP11B2 IHC staining do not reflect enzyme expression levels. NB2: Autoradiography for cases 4–8 can be found in Figure S3.
of multiple fluorine substituents presented a convenient option for labeling with fluorine-18, to give [18F]AldoView, while fully retaining the chemical structure. We have recently developed sulfonylum salts as a new class of leaving groups for the radiofluorination of drug-like molecules, and in this case, incorporation of a dibenzothiophenium salt in the 3-position of the pyridine moiety provided a practical and efficient route to the precursor for labeling. Treatment with [18F]fluoride allowed direct labeling in a single step in 42 ± 8% (decay-corrected; n = 12) radiochemical yield. The use of fluorine-18 (half-life 110 min) for labeling and the high efficiency of the labeling reaction ensure that [18F]AldoView can become widely available for routine diagnostic use.

The pharmacokinetic properties of [18F]AldoView were characterized preclinically with PET imaging and biodistribution studies in mice. PET/CT imaging in mice revealed a high initial uptake in the liver, with rapid clearance, resulting in low background in all tissues surrounding the adrenals glands at later time points. In the biodistribution studies, the liver uptake was less pronounced, whereas the peak uptake and clearance in the kidneys occurred at a later time point. These discrepancies may reflect physiological effects of the anesthesia in the animals imaged with PET. The metabolic stability of the tracer appears adequate, with 21% intact parent fraction measured in blood after 60 min. It should be noted that there are substantial species differences between adrenal enzymes and that rodents do not provide a suitable model for human CYP11B2. In a recent study by Novartis, the potency of aldosterone synthase inhibitors for rat CYP11B2 was shown to inversely correlate with CYP11B2/CYP11B1 selectivity for the human isoforms. The adrenal uptake and clearance profile of [18F]AldoView in mice, which closely resembles that of the kidneys, is therefore encouraging and consistent with the high hCYP11B2 selectivity (B2/B1 ratio of 93) of this ligand.

The binding of the tracer in the adrenal glands resected from patients with PHA was in good visual agreement with IHC staining for CYP11B2 and demarcated areas of intense uptake in some but not all of the designated APAs. In the cortex, diffuse binding patterns were evident, with highly localized “hot spots” consistent with APCCs. In tissue from control cases, that is, from patients with CYP11B2-negative adrenal tumors, tracer binding was consistently low. The ratio of tracer uptake between CYP11B2-positive and -negative areas ranged from 2 to 9, with >95% specific binding in areas with a high tracer uptake. Surgical removal of the adrenal glands from the PHA patients included in our study resulted in good clinical outcomes. In cases 1 and 3, where phosphorimaging with [18F]AldoView revealed APCCs in the adrenal cortex, adenolecotomy resulted in complete cure of the patients as evident from a normalized blood pressure as well as normal renin and aldosterone levels. The subjects 2, 4, and 5 still had elevated but controlled blood pressure postoperatively. Treatment with antihypertensive medication could be reduced from three to one drug in patients 2 and 5, resulting in a blood pressure of 140/90 mm Hg. Patient 4 had a blood pressure of 160/89 mm Hg postoperatively, normalized aldosterone and renin values, and did not continue treatment with any of the six antihypertensive drugs taken preadrenalectomy. Taken together, the results suggest that [18F]AldoView can allow imaging of the CYP11B2 expression in human adrenals with PET.

**Study Limitations.** Due to species differences in the human and murine adrenal enzymes, the preclinical characterization of [18F]AldoView in mice did not provide direct evidence that the tracer allows imaging of CYP11B2 in vivo. Although imaging in human adrenal sections was in good visual agreement with the IHC staining of CYP11B2, in vitro binding does not directly translate to in vivo imaging. In addition, we were unable to quantify the correlation of tracer binding with enzyme expression due to the lack of independent methods to measure the aldosterone synthase expression. This was particularly challenging in tissues designated as APAs by morphological inspection, for which the IHC staining of CYP11B2 gave inconclusive results due to the lack of contrast, and tracer binding varied considerably between the cases. We were also unable to demonstrate lack of tracer binding to areas with high CYP11B1 expression, as in our hands, antibody staining of this enzyme gave inconclusive results (uniform staining in tissue sections). Finally, we note that, although CYP11B2 is the only known source of aldosterone, regulation of aldosterone secretion is not entirely transcriptional, and therefore, CYP11B2 expression measured with PET may not correlate directly with the rate of aldosterone production in the adrenals.

**Translational Outlook.** The study aim was to address the unmet clinical need for a noninvasive method to better diagnose PHA and to stratify patients with unilateral PHA for surgical treatment. [18F]AldoView, the first highly selective aldosterone synthase tracer, was developed to allow imaging of the adrenal hCYP11B2 expression with PET. In preclinical studies in mice, the PET tracer showed a favorable pharmacokinetic profile and organ distribution, suggesting that in vivo imaging will be feasible. Binding studies in tissue sections from surgically resected adrenals provided evidence for a highly selective tracer binding to hCYP11B2. The results are encouraging and warrant translation to human studies. First-in-human PET scans are due to commence in the near future. The results will determine the diagnostic efficacy of image-derived adrenal enzymatic lateralization with PET/CT.
be >95% in each case. The purity of compound 7, as assessed by HPLC, was >97% (Figure S1).

N-Cyclopropyl-4,5-difluoro-2-nitroaniline (1). To a stirring solution of 2,4,5-trifluorotribenzoic (5.0 g, 3.24 mL, 28.24 mmol, 1.0 equiv) and triethylamine (8.57 g, 11.8 mL, 84.70 mmol, 3.0 equiv) in tetrahydrofuran (25.7 mL) at room temperature, cyclopentalamine (6.65 g, 0.87 mL, 118 mmol, 4.2 equiv) was added dropwise over 1 h (the solution became very thick during the addition). The resulting yellow mixture was stirred at 60 °C for 2 h. After cooling to room temperature, the reaction was quenched with water (20 mL) and diluted with ethyl acetate (20 mL). The organic layer was separated, and the aqueous layer was extracted with ethyl acetate (2 × 30 mL). The combined organic layers were washed with water, brine, dried over MgSO₄, filtered, and concentrated in vacuo. The crude product was purified by flash column chromatography (ethyl acetate/petrol 0 to 10%) on a silica gel to afford the desired product as a yellow solid (690 mg) in 11% yield. ¹H NMR (CDCl₃, 500 MHz): δ 9.31 (d, J = 1.9 Hz, 1H), 8.80 (d, J = 2.2 Hz, 1H), 8.45 (t, J = 2.1 Hz, 1H), 7.57 (dd, J = 10.2, 7.3 Hz, 1H), 7.41 (d, J = 9.7, 7.0 Hz, 1H), 3.57 (tt, J = 6.9, 3.8 Hz, 1H), 1.29–1.21 (m, 2H), and 0.85–0.78 (m, 2H); ¹³C NMR (CDCl₃, 126 MHz): δ 151.8, 151.0, 149.6 (dd, J = 40.2, 15.2 Hz), 147.7, 147.7 (dd, J = 38.9, 17.6 Hz), 138.9, 138.0 (d, J = 24.0 Hz), 132.7 (d, J = 10.8 Hz), 128.0, 120.8, 107.2 (d, J = 19.7 Hz), 99.2 (d, J = 23.2 Hz), and 26.6, 9.2 (2C); ¹⁹F NMR (CDCl₃, 282 MHz): δ −139.41 (d, J = 20.5 Hz) and −142.23 (d, J = 20.5 Hz); and HRMS: C₃₁H₂₁F₄N₉O⁺ (290.0919) calcd, 290.0914, 350.0104. 2-Ethylethyl 3-(′,′,′-Dimethoxy-5-methyl-1,1′-biphenyl-2-yl)thiopropanoate (5). The compound was prepared as previously reported. In brief, heating of a mixture of 2-bromo-1-iodo-4-methylbenzene (1 equiv), 2-ethylhexyl mercaptotripropanoate (1 equiv), triethylamine (2 equiv), tris(dibenzylideneacetone)dipalladium(0) (0.65 mol %), and Xanthos (3 mol %) in toluene at 125 °C for 2 h and subsequent workup by filtration over Celite and flash column chromatography provided 2-ethylthiethyl 3-((2-bromo-4-methylphenyl)thiopropanoate as a colorless oil (96%). This intermediate (1 equiv) was further reacted in a Suzuki cross-coupling reaction with/refined by (3,5-dimethylphenyl)boronic acid (1.25 equiv), potassium carbonate (4 equiv), and PEPPSI-IPr catalyst (3 mol %) in a mixture of toluene and water for 2 h. Filtration over Celite, followed by an aqueous workup and flash column chromatography afforded the title compound as a colorless oil (95%).

1-Cyclopropyl-2-(3,3′-[′,′-Dimethoxy-5-methyl-[1,1′-biphenyl]-2-yl)thiopridin-3-yl]-5,6-difluoro-1H-benz[d]imidazole (6). To a flame-dried three-necked round-bottom flask equipped with an argon inlet and condenser were added sequentially anhydrous K₂CO₃ (1.96 g, 14.22 mmol, 3.0 equiv), bis(2-diphenylphosphino)biphenyl ether (198 mg, 0.37 mmol, 2 equiv), and Xantphos (3 mol %) in toluene and degassed for 5 min using a balloon filled with argon and a vent needle and then heated at 125 °C for 1 h. After cooling to room temperature, the reaction mixture was concentrated in vacuo. The crude product was purified by flash column chromatography (ethyl acetate/petrol 10 to 60%) on silica gel to afford the desired product as an orange solid (296 mg) in 64% yield. ¹H NMR (CDCl₃, 500 MHz): δ 9.04 (s, J = 1.7 Hz, 1H), 8.61 (d, J = 2.8 Hz, 1H), 8.01 (dd, J = 9.1, 2.8, 1.7 Hz, 1H), 7.57 (dd, J = 10.3, 7.3 Hz, 1H), 7.41 (d, J = 9.7, 7.0 Hz, 1H), 3.58 (tt, J = 6.8, 3.8 Hz, 1H), 1.25 (dd, J = 6.9, 1.2 Hz, 2H), and 0.84–0.79 (m, 2H); ¹³C NMR (CDCl₃, 126 MHz): δ 159.1 (d, J = 258.2 Hz), 151.1, 150.2 (d, J = 239, 15.4 Hz), 146.9 (dd, J = 21.4, 15.4 Hz), 145.5 (d, J = 4.3 Hz), 139.2 (d, J = 23.1 Hz), 137.8 (d, J = 11.0 Hz), 132.6 (d, J = 10.8 Hz), 127.8 (d, J = 4.2 Hz), 123.2 (d, J = 20.0 Hz), 107.6 (d, J = 19.9 Hz), 99.2 (d, J = 23.1 Hz), 26.6, and 9.1 (2C); ¹⁹F NMR (CDCl₃, 282 MHz): δ −125.72, −139.48 (d, J = 20.6 Hz) and −142.30 (d, J = 20.6 Hz); and HRMS: C₂₁H₁₈F₂N₄O⁺ (350.0900) calcd, 350.0900.

1-Cyclopropyl-2-(3,3′-[′,′-Dimethoxy-5-methyl-[1,1′-biphenyl]-2-yl)thiopridin-3-yl]-5,6-difluoro-1H-benz[d]imidazole (7). To a solution of 1-cyclopropyl-2-(3,3′-[′,′-dimethoxy-5-methyl-[1,1′-biphenyl]-2-yl)thiopridin-3-yl)-5,6-difluoro-1H-benz[d]imidazole (6) (322 mg, 0.92 mmol, 1 equiv) in acetone (analytical reagent grade; 3.4 mL) at 0 °C, was added a freshly prepared solution of Ca(OCl₂)₂ (a fresh bottle

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of technical grade reagent (65% w/w) was used; 45.4 mg, 0.21 mmol, 0.4 equiv) in an aqueous acetate buffer (1 M, pH = 4; 3.4 mL). Under stirring, the resulting yellow solution was allowed to come to room temperature. Stirring was then continued for 1 h. The reaction was subsequently diluted with dichloromethane (30 mL) and water (20 mL). The organic layer was separated, and the aqueous layer was extracted with dichloromethane (2 × 30 mL). The combined organic layers were washed with an aqueous solution of sodium trifluoromethanesulfonate (1 M; 10 mL), dried over MgSO₄, and concentrated in vacuo. The crude product was purified by flash column chromatography on silica gel. Initially, an ethyl acetate/petrol gradient from 70% to 100% was used to remove lipophilic impurities. The column was then continued using a methanol/dichloromethane gradient from 1 to 3% to remove more polar impurities before eluting the product fraction at 3% methanol/dichloromethane. The title compound was isolated as a yellow solid (114 mg) in 32% yield (48 mg of the starting material was recovered). ^1H NMR (CD₃CN, 400 MHz): δ 9.39 (d, J = 1.9 Hz, 1H), 9.09 (d, J = 2.3 Hz, 1H), 8.20 (d, J = 1.8, 0.7 Hz, 1H), 8.07–8.02 (m, 2H), 7.60–7.50 (m, 3H), 7.45 (d, J = 2.1 Hz, 1H), 6.78 (d, J = 2.1 Hz, 1H), 4.00 (s, 3H), 3.94 (s, 3H), 3.34 (tt, J = 6.9, 3.8 Hz, 1H), 2.57 (d, J = 0.7 Hz, 3H), 0.92–0.78 (m, 2H), and 0.65–0.52 (m, 2H); ^13C NMR (CD₃CN, 126 MHz): δ 169.9, 159.8, 155.7, 152.1, 151.2, 150.4 (d, J = 32.2, 15.7 Hz), 147.7 (dd, J = 31.1, 15.3 Hz), 147.5, 143.3, 140.8, 138.6 (d, J = 10.7 Hz), 136.8, 133.9 (d, J = 11.4 Hz), 133.9, 129.8, 129.1, 129.0, 126.7, 125.8, 107.8 (d, J = 19.6 Hz), 107.3, 103.3, 101.6, 100.6 (d, J = 23.3 Hz), 58.2, 57.7, 27.0, 21.8, and 9.0 (2C); ^19F NMR (CD₃CN, 282 MHz): δ −79.3, −142.07 (d, J = 20.0 Hz) and −144.97 (d, J = 20.0 Hz); and HRMS: 528.1555 (C₃₀H₂₄F₂N₃O₂S⁺) calcd, 528.1552.

**Radiosynthesis of [¹⁸F]AldoView.** All labeling reactions were performed manually using [¹⁸F]fluoride in [¹⁸O]water. Radio-HPLC was performed on an Agilent 1200 HPLC system equipped with a 1200 Series Diode Array Detector and a GABI Star NaI(Tl) PET/CT Imaging System (GABI Imaging Systems, Budapest, Hungary). Mice (n = 6) were anesthetized with isoflurane (2% v/v in oxygen) and placed on the preheated bed of the scanner (set at 38 °C). [¹⁸F]AldoView (7.3 ± 4.1 MBq, formulated in 100–150 µL saline containing 5% ethanol) was injected into the tail vein via intravenous cannulation. After injection, the catheter was carefully removed. The breathing rate and body temperature of the animals were closely monitored during the dynamic PET scans and, if necessary, the isoflurane dose was adjusted. Scans were recorded over 1 h, and the animals were subsequently sacrificed by cervical dislocation.

**Biodistribution Studies.** [¹⁸F]AldoView (6.6 ± 1.9 MBq, formulated in saline containing 5% ethanol) was administered intravenously into the tail vein of female wild-type BALB/c mice (n = 3–4 per time point) without anesthesia. At designated time points (15, 30, 60, and 120 min postinjection), animals were anesthetized with isoflurane (4% v/v in oxygen) and the blood was taken by cardiac puncture. Mice were subsequently sacrificed by cervical dislocation. The organs of interest were sampled and weighed, and the radioactivity content was measured by automated gamma counting (PerkinElmer Wizard®). Results were expressed as % ID/g body-weight.

**Metabolite Analysis.** Blood metabolite analysis was performed as part of the biodistribution studies (n = 2–3 per time point). Blood samples were collected in heparin-coated tubes, and an aliquot (20 µL) was taken for gamma counting. After centrifugation (5 min, 13,000 rpm), the plasma (~500 µL) was separated. Plasma proteins were subsequently precipitated with ice-cold acetone (1 mL), and samples were centrifuged (5 min, 13,000 rpm). An aliquot of the supernatant (500 µL) was separated from the pellet and diluted with water (500 µL). The recovery of radioactivity in the supernatant was near quantitative (≥95%). Analysis was carried out by radio-HPLC using a Phenomenex Onyx monolithic C-18 column (100 × 10 mm). The mobile phase consisted of water and methanol, and the flow rate was 5 mL/min. An isocratic eluent (10% methanol in water) was used for 3 min, followed by a gradient from 10 to 70% methanol over 7 min.

**Adrenal Tissue.** Adrenal surgical specimens were obtained from patients with Conn’s disease, Cushing’s disease, pheochromocytoma, and adrenal tumors, who had consented to the use of their tissue for research applications. Ethical approval was obtained from the UK Multicentre Research Ethics Committee (06/ Q0104/133). Following routine adenectomy, surgical specimens were immediately flash frozen using a slurry of isopentane and dry ice. Cryosections (20 µm) obtained from the frozen tissue blocks were mounted onlysine-coated microscope slides and stored at −80 °C.

**Quantitative Phosphorimaging.** Thawed tissue sections on microscopic slides were dipped in pentane for 5 s to degrease the tissue. This step was required to avoid nonspecific binding (NSB), in
AldoView in TBS and diluted to give a final concentration of 2 MBq/mL. For the determination of NSB, a solution of the nonradioactive unlabeled AldoView was diluted with TBS to a concentration of 2 MBq/mL. After incubation, unbound [18F]AldoView was removed by washing the sections in ice-cold TBS (three times for 5 min) and water (1 min). Internal standards were prepared by serial dilution of the [18F]AldoView solution used for determination of the TB. Tissue sections and internal standards absorbed on filter paper were left to air-dry and subsequently exposed to a phosphor screen (BAS-IP MS; GE Healthcare) overnight. Phosphorimaging was performed on a Typhoon Trio scanner (GE Healthcare). All phosphorimaging experiments were performed in at least three adjacent tissue sections per case, tissue block, and experimental condition (TB, NSB). Quantification of phosphor images was performed using the analysis software ImageJ (version 1.48). Determination of TB and NSB (in kBq/cm²; Table S1) was based on correlation curves generated from the internal standards. The specific [18F]AldoView binding was calculated as the difference between TB and NSB.

Immunohistochemistry. IHC staining was carried out in a flash-frozen tissue on sequential sections to those used for quantitative phosphorimaging. Tissue sections were incubated with mouse monoclonal anti-human CYP11B2 (gifted by Prof. Celso E. Gomez-Sanchez; 1:1000, overnight, 4 °C), followed by biotinylated antimouse IgG (Vector Laboratories, 1:400, 1 h, room temperature) and VectaStain ABC Kit (Vector Laboratories, 30 min, room temperature). Color was developed with 3,3′-diaminobenzidine and hydrogen peroxide. Counterstaining was carried out using Mayer’s haematoxylin.

ASSOCIATED CONTENT

Supporting Information

The Supporting Information is available free of charge at https://pubs.acs.org/doi/10.1021/acs.jmedchem.1c00539.

NMR spectra of compounds 3, 4, 6, and 7; HPLC trace of the [18F]AldoView labeling precursor; radio-HPLC trace of [18F]AldoView; [18F]AldoView binding in adrenal surgical specimens; and quantitative phosphorimaging with [18F]AldoView in tissue sections from surgically resected adrenal glands (PDF)

Molecular formula strings (CSV)

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