

# Synthesis and Evaluation of Novologues as C-Terminal Hsp90 Inhibitors with Cytoprotective Activity against Sensory Neuron Glucotoxicity

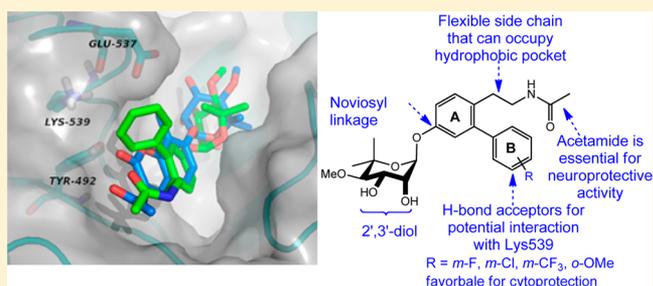
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## S Supporting Information

**ABSTRACT:** Compound **2** (KU-32) is a first-generation novologue (a novobiocin-based, C-terminal, heat shock protein 90 (Hsp90) inhibitor) that decreases glucose-induced death of primary sensory neurons and reverses numerous clinical indices of diabetic peripheral neuropathy in mice. The current study sought to exploit the C-terminal binding site of Hsp90 to determine whether the optimization of hydrogen bonding and hydrophobic interactions of second-generation novologues could enhance neuroprotective activity. Using a series of substituted phenylboronic acids to replace the coumarin lactone of **2**, we identified that electronegative atoms placed at the *meta*-position of the B-ring exhibit improved cytoprotective activity, which is believed to result from favorable interactions with Lys539 in the Hsp90 C-terminal binding pocket. Consistent with these results, a *meta*-3-fluorophenyl substituted novologue (**13b**) exhibited a 14-fold lower ED<sub>50</sub> for protection against glucose-induced toxicity of primary sensory neurons compared to **2**.

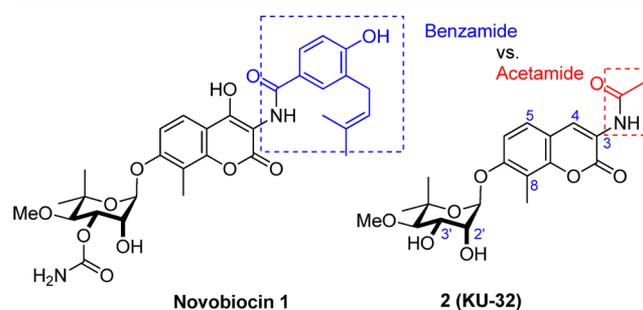


## INTRODUCTION

Approximately 26 million Americans are afflicted with either type 1 or type 2 diabetes. Despite the use of insulin and oral antidiabetic medications to help maintain euglycemia, about 60–70% of these individuals develop diabetic peripheral neuropathy (DPN).<sup>1</sup> To date, approaches toward the treatment of DPN have centered on pathways/targets directly linked to hyperglycemia (i.e., polyol and hexosamine pathways, advanced glycation end products (AGEs), enhanced oxidative stress, and PKC activation).<sup>2</sup> Unfortunately, the contribution of these targets/pathways to the progression of DPN differs between individuals and does not occur with biochemical uniformity, and consequently, these approaches have resulted in little success for the management of DPN. As an alternative approach, we have explored the pharmacologic modulation of molecular chaperones to promote a broad cytoprotective response that may enhance a patient's ability to tolerate hyperglycemic insults and improve the symptoms of DPN.

Molecular chaperones, such as heat shock proteins 90 and 70 (Hsp90, Hsp70), are essential for folding nascent polypeptides into their biologically active structures and for the refolding of aggregated and denatured proteins that occur upon cellular stress.<sup>3,4</sup> Numerous conditions that cause cell stress can also induce the “heat shock response” (HSR): the transcriptional upregulation of antioxidant genes and chaperones such as

Hsp70. Importantly, small molecule inhibition of Hsp90 is sufficient to induce the HSR. Compound **2** (KU-32)<sup>5,6</sup> is a small molecule Hsp90 C-terminal inhibitor that is based on novobiocin **1**, a naturally occurring antimicrobial agent that inhibits DNA gyrase (Figure 1). Although the etiology of DPN

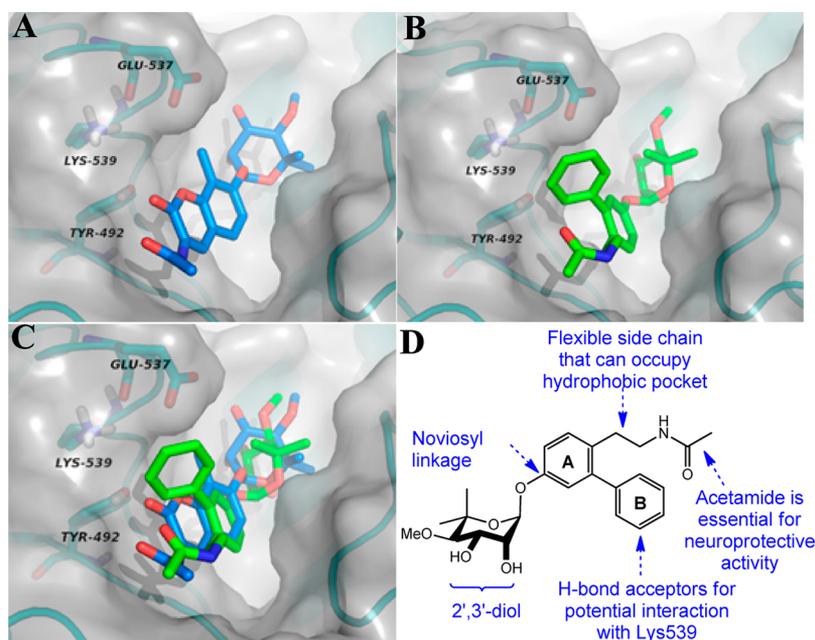


**Figure 1.** Chemical structures of novobiocin **1** and **2** (KU-32).

is unrelated to the accumulation of one specific misfolded or aggregated protein, hyperglycemia can increase oxidative stress and the oxidative modification of amino acids<sup>7</sup> that impair

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**Figure 2.** (A) Compound **2** docked to Hsp90 C-terminal binding site. (B) Novologue docked to Hsp90 C-terminal binding site. (C) Overlay of **2** and novologue docked to Hsp90 C-terminal binding site. All structures docked into Hsp90 $\alpha$  open homology model. (D) Structure of novologue and its attributes.

protein folding,<sup>8</sup> decrease mitochondrial protein import,<sup>9</sup> and promote mitochondrial dysfunction.<sup>2,7a</sup> Even in the absence of a single, disease-specific protein aggregate, we have shown that pharmacologic induction of cytoprotective molecular chaperones can improve myelinated and unmyelinated fiber function in cellular models of glucotoxic stress and animal models of DPN.<sup>10</sup> Mechanistically, compound **2** was ineffective at preventing neuregulin-induced demyelination of myelinated cultures of sensory neurons prepared from Hsp70.1 and 70.3 double knockout mice, indicating that Hsp70 is necessary for the neuroprotective activity manifested by compound **2**. Similarly, weekly treatment with **2** restored normal sensory and motor nerve function in diabetic wild type mice but was unable to reverse multiple clinical indices of DPN in the diabetic Hsp70 knockout mice.<sup>10</sup> Collectively, these studies provide the biological and clinical rationale to support the modulation of molecular chaperones as a viable approach toward treating.

An enviable aspect of **2** is that it induces Hsp70 at concentrations well below those needed to inhibit Hsp90's protein folding ability, thus, it possesses a rather broad therapeutic window that dissociates cytoprotective properties from potentially cytotoxic effects resulting from the degradation of Hsp90-dependent client proteins.<sup>4</sup> We have previously shown that molecules containing a benzamide, as found in novobiocin, exhibit antiproliferative activities, whereas molecules such as **2**, containing an acetamide, manifest neuroprotective properties. However, these prior studies sought to evaluate structure–activity relationships for novobiocin analogues as anticancer agents<sup>11,12</sup> rather than exploring chemical attributes that enhance the neuroprotective properties of novobiocin-based analogues. Therefore, the goal of this work was to determine whether diversification of the scaffold of compound **2** could identify structure–activity relationships (SAR) for novobiocin analogues (novologues) that enhance the neuroprotective properties manifested by Hsp90 C-terminal inhibitors.

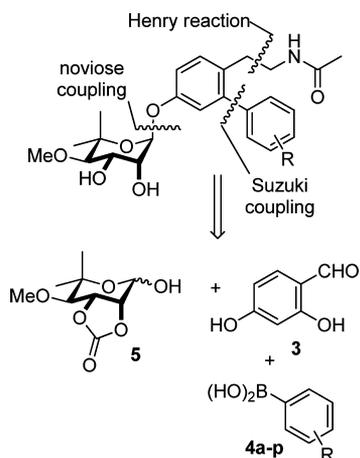
Recently, we used molecular modeling and azide-containing novobiocin derivatives as photoaffinity probes to elucidate, for the first time, the Hsp90 C-terminal binding site.<sup>13</sup> As shown in Figure 2A–C, compound **2** docks to this region and appears to exhibit binding interactions with both the protein backbone and the amino acid side chains similar to those manifested by novobiocin. Interestingly, the coumarin lactone of **2** appears too distant from Lys539 to provide complementary interactions with this residue. In addition, the 3-amido side chain appears to project into a large hydrophobic pocket that could accommodate more flexible linkers. As a consequence of these observations, the novologue scaffold (Figure 2D) was designed to project the B-ring into the pocket wherein Lys539 resides and to serve as a lead compound for further diversification. In addition, the flexible ethyl amide projecting from the A-ring could accommodate a number of orientations that could better occupy the large hydrophobic pocket that remains vacant in the presence of **2**.

On the basis of the novologue design, we envisioned construction of a parallel library to validate this scaffold for use as a neuroprotective agent. The library was designed so that the 3'-carbamate on noviose was omitted, on the basis of prior studies that showed this group to be detrimental to Hsp90 inhibitory activity.<sup>5</sup> In contrast, additional hydrophobic and hydrogen bonding interactions could be provided by the incorporation of functionalities onto the 3-aryl substituent (B-ring), which was designed to provide complementary interactions with Lys539. The 4-ethyl acetamide was included to occupy the binding pocket about the coumarin ring system. Consistent with data obtained from prior studies, the 7-noviosyl linkage was maintained as well as the requisite 2',3'-diol. In this article, we report the parallel synthesis of rationally designed novologues as Hsp90 C-terminal inhibitors and assessment of their neuroprotective activities.

## RESULTS AND DISCUSSION

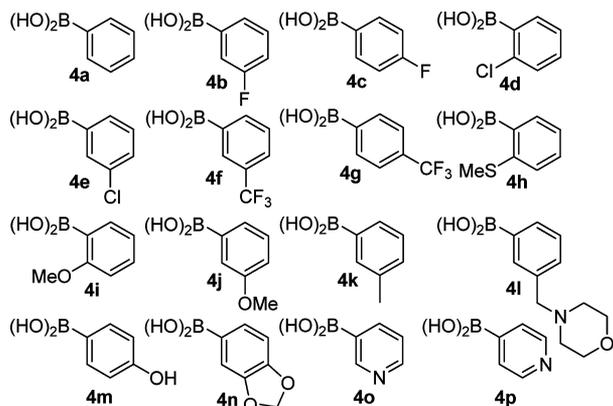
Retrosynthetically, a library of novologues was envisioned for construction via four components (Scheme 1): a resorcinolic

**Scheme 1. Retrosynthetic Analysis for the Construction of Novologues**



benzaldehyde (3), a variety of commercially available boronic acids (4a–p), noviose (5), and the acetamide side chain (Scheme 1). Prior work from our laboratory demonstrated that the trichloroacetimidate of noviose carbonate undergoes rapid coupling with phenols to give the desired  $\alpha$ -anomer in high yield.

The boronic acids chosen for this study contain both electronic and steric moieties that could aid in elucidation of structure–activity relationships and provide crucial interactions with Lys539 and the surrounding pocket. Toward this goal, phenylboronic acids (Figure 3) containing electronegative



**Figure 3.** Boronic acids selected for incorporation into novologue X scaffold.

atoms at the *meta*- and *para*-positions were explored. In addition, hydrogen bond acceptors were included at these locations to provide potential hydrogen bonding interactions with the protonated form of Lys539. To serve as controls, hydrophobic groups (4j, 4k) and a tertiary amine (4l) were included in this series.

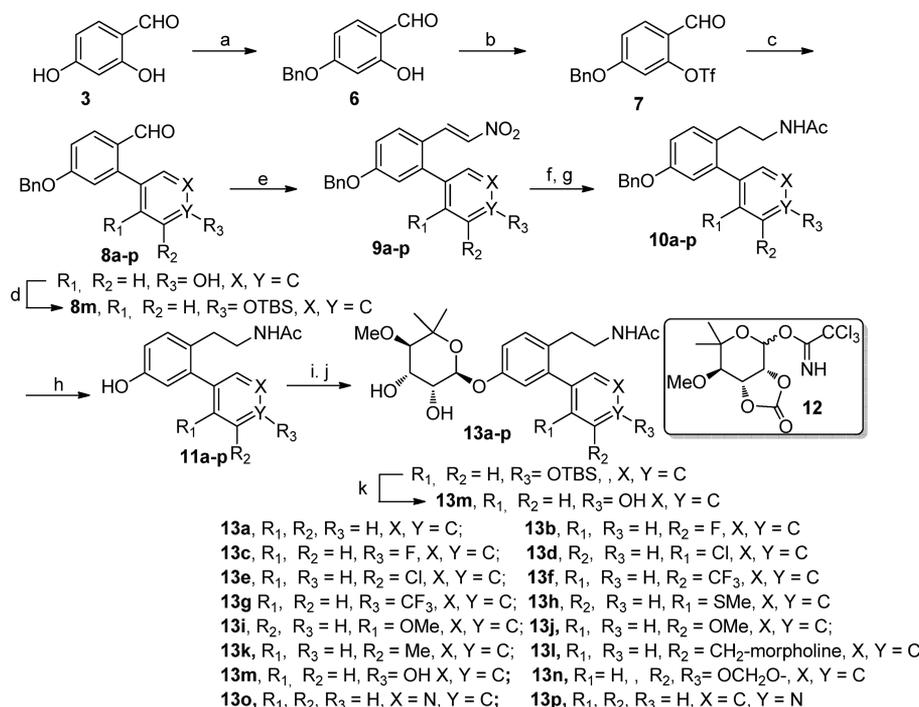
The synthesis of ethyl acetamide side chain containing novologues 13a–p began with commercially available 2,4-dihydroxybenzaldehyde, 3. The 4-phenol of resorcinolic benzaldehyde 3 was protected as the corresponding benzyl

ether 6,<sup>14</sup> and the 2-phenol converted to triflate 7 using trifluoromethanesulfonic anhydride and triethylamine (Scheme 2). Compound 7 was subsequently coupled with commercially available aryl boronic acids (4a–p) under standard Suzuki conditions<sup>15,16</sup> to give biaryl ring systems 8a–p in good yields. Benzaldehydes 8a–p were converted to the corresponding nitrostyrenes (9a–p), following a Henry reaction with nitromethane and ammonium acetate.<sup>17,18</sup> Reduction of the nitro and olefin functionalities with lithium aluminum hydride was followed by acylation of the resulting amines to afford acetamides 10a–p in good yields. The benzyl ether of compounds 10a–p was cleaved under hydrogenolysis conditions to afford phenols 11a–p, which were coupled with the trichloroacetimidate of noviose carbonate 12<sup>5,19</sup> in the presence of a catalytic amount of boron trifluoride etherate. The resulting noviosylated biaryl systems were exposed to methanolic ammonia to solvolyze the cyclic carbonate and give the desired novologues (13a–p) in good to moderate yields.

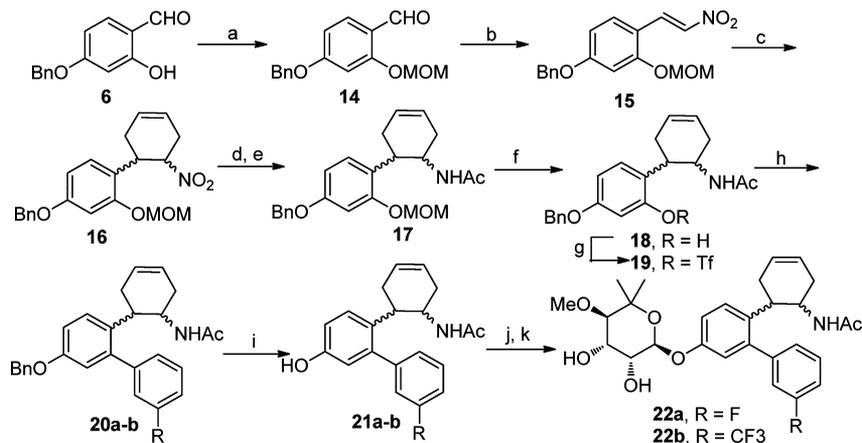
In addition, two cyclohexene analogues 22a–b were pursued to test our hypothesis regarding the region surrounding the flexible side chain (Scheme 3). Although these molecules contain the same linker length, these analogues contain a bulky cyclohexane tether between the biaryl ring system and the acetamide.

Synthesis of cyclohexene analogues 22a–b began with the previously described phenol 6, which was protected as the methoxymethyl (MOM) ether 14<sup>20</sup> before the aldehyde of which was converted to nitrostyrene 15 under Henry conditions.<sup>16</sup> The electron deficient nitrostyrene (15) was subjected to a Diels–Alder cycloaddition with excess butadiene to give an enantiomeric mixture of cyclohexene derivative 16 in excellent yield.<sup>21</sup> The nitro group of 16 was selectively reduced to the amine via zinc dust and acidic isopropyl alcohol,<sup>22</sup> followed by acetylation to afford acetamide 17 in 71% yield over two steps. To construct the biaryl ring system, the MOM-ether was cleaved to give the phenol, which was then converted to the corresponding triflate, 19. A Suzuki reaction between 19 and 3-fluorophenylboronic acid (4b) or 3-(trifluoromethyl)phenylboronic acid (4f) yielded biaryl compounds 20a or 20b, respectively. Finally, boron trifluoride etherate promoted removal of the benzyl ether<sup>23</sup> on compounds 20a–b and gave phenols 21a–b. Lewis acid-catalyzed noviosylation of 21a–b, with activated noviose carbonate (10), followed by methanolysis, afforded an inseparable mixture of diastereomeric products, 22a–b.

**Evaluation of Neuroprotective Efficacy.** Upon synthesis of ethyl acetamide side chain novologues 13a–p that contain various substitutions on the B-ring (hydrogen bond acceptors, hydrogen bond donors, hydrophobic groups, and a tertiary amine), their neuroprotective efficacy against glucose-induced toxicity of embryonic dorsal root ganglion (DRG) sensory neuron cultures was evaluated. As shown in Table 1, *meta*-substituted acetamide novologues (13b, 13e, and 13f) showed significant protection against glucotoxicity and were comparable to that observed with compound 2. Although the corresponding *ortho*- and *para*-substituted (13c, 13d, and 13g) derivatives showed significant protection against glucose-induced cell death, they were modestly less effective than novologues 13b, 13e, and 13f. However, in the case of analogues 13i (*ortho*-OMe) and 13j (*meta*-OMe), the opposite trend was observed. Electronegative atoms at the *meta*-position (F, Cl, CF<sub>3</sub>) exhibited greater cytoprotective activity, which is

Scheme 2. Synthesis of Ethyl Acetamide Side Chain Containing Novologues<sup>a</sup>

<sup>a</sup>Reagents and conditions: (a) NaHCO<sub>3</sub>, BnBr, CH<sub>3</sub>CN, rt, 75%; (b) Tf<sub>2</sub>O, Et<sub>3</sub>N, CH<sub>2</sub>Cl<sub>2</sub>, 60%; (c) **4a-p**, Pd(PPh<sub>3</sub>)<sub>4</sub>, K<sub>2</sub>CO<sub>3</sub>, DMF, 65–92%; (d) TBSCl, imid, CH<sub>2</sub>Cl<sub>2</sub>; (e) CH<sub>3</sub>NO<sub>2</sub>, NH<sub>4</sub>OAc, 90–98%; (f) LiAlH<sub>4</sub>, THF, 0 °C-rt, 30 min; (g) Ac<sub>2</sub>O, Et<sub>3</sub>N, 61–72% over 2 steps; (h) H<sub>2</sub>, Pd/C, MeOH, 69–81%; (i) **12**, BF<sub>3</sub>·OEt<sub>2</sub>, CH<sub>2</sub>Cl<sub>2</sub>, rt; (j) Et<sub>3</sub>N, MeOH, rt, overnight, 55–88% over 2 steps; (k) TBAF, THF, rt, 30 min.

Scheme 3. Synthesis of Cyclohexene Containing Novologues<sup>a</sup>

<sup>a</sup>Reagents and conditions: (a) MOMCl, DIPEA, CH<sub>2</sub>Cl<sub>2</sub>, rt, 84%; (b) CH<sub>3</sub>NO<sub>2</sub>, NH<sub>4</sub>OAc, 84%; (c) 1,4-butadiene, toluene 120 °C, 95%; (d) Zn dust, 1N HCl; (e) Ac<sub>2</sub>O, Et<sub>3</sub>N, 71% over 2 steps; (f) HCl, MeOH; (g) phenyl triflimide, Et<sub>3</sub>N, 85%; (h) **4b** or **4f**, Pd(PPh<sub>3</sub>)<sub>4</sub>, K<sub>2</sub>CO<sub>3</sub>, DMF; (i) HSEtSH, BF<sub>3</sub>·OEt<sub>2</sub>, CH<sub>2</sub>Cl<sub>2</sub>; (j) **12**, BF<sub>3</sub>·OEt<sub>2</sub>, CH<sub>2</sub>Cl<sub>2</sub>; (k) Et<sub>3</sub>N, MeOH, rt, overnight.

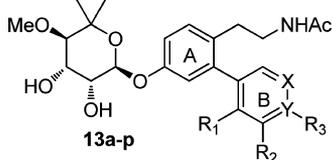
believed to result from favorable interactions with Lys539 in the Hsp90 C-terminal binding pocket. Consistent with this hypothesis, increasing the size of the electronegative atom at the *meta*-position (F to Cl to CF<sub>3</sub>) resulted in a decrease in neuroprotective activity. Analogue **13b** (*meta*-F) was among the most cytoprotective (95% ± 14) compounds evaluated.

Electronegative atoms at the *ortho*- or *para*-position on ring B (**13c**, **13d**, and **13g**) manifested activities comparable to the unsubstituted analogue (**13a**) and were less active than the corresponding *meta*-substituted analogues (**13b**, **13e**, and **13f**). Although novologues **13d** and **13g** manifested protection against neuronal glucotoxicity, they were less effective than our

previous compound **2** and **13b**. Hydrogen bond donors at the *para*-position (**13m**) appeared to be undesired as **13m** (*para*-OH) was unable to provide significant protection against glucotoxicity. It was also somewhat, but not significantly less protective than the unsubstituted analogue (**13a**).

On the other hand, hydrogen bond acceptors at the *para*-position (**13c** and **13g**) protected against glucose-induced neuronal death but did not display significantly increased protection compared to the novologue containing a *para*-position hydrogen bond donor (**13m**).

Pyridine-containing analogues (**13o-p**) were also synthesized and evaluated for neuroprotective activity. The 3-pyridine

**Table 1.** Cell Viability Data of Ethyl Acetamide Side Chain Containing Novologues

entry	R <sub>1</sub>	R <sub>2</sub>	R <sub>3</sub>	X	Y	% of cell viability <sup>a</sup>
2						86 ± 2
13a	H	H	H	C	C	76 ± 11 <sup>#b</sup>
13b	H	F	H	C	C	95 ± 14 <sup>#b</sup>
13c	H	H	F	C	C	75 ± 27 <sup>#b</sup>
13d	Cl	H	H	C	C	71 ± 21 <sup>#,*,b,c</sup>
13e	H	Cl	H	C	C	90 ± 23 <sup>#b</sup>
13f	H	CF <sub>3</sub>	H	C	C	83 ± 16 <sup>#b</sup>
13g	H	H	CF <sub>3</sub>	C	C	74 ± 19 <sup>#,*,b,c</sup>
13h	SMe	H	H	C	C	83 ± 40 <sup>#b</sup>
13i	OMe	H	H	C	C	92 ± 10 <sup>#b</sup>
13j	H	OMe	H	C	C	78 ± 34 <sup>#b</sup>
13k	H	Me	H	C	C	82 ± 30 <sup>#b</sup>
13l	H	CH <sub>2</sub> -N-morpholine	H	C	C	83 ± 26 <sup>#b</sup>
13m	H	H	OH	C	C	67 ± 10 <sup>*c</sup>
13n	H	-OCH <sub>2</sub> O-		C	C	83 ± 18 <sup>#b</sup>
13o	H	H	H	N	C	61 ± 7 <sup>*c</sup>
13p	H	H	H	C	N	81 ± 12 <sup>#b</sup>

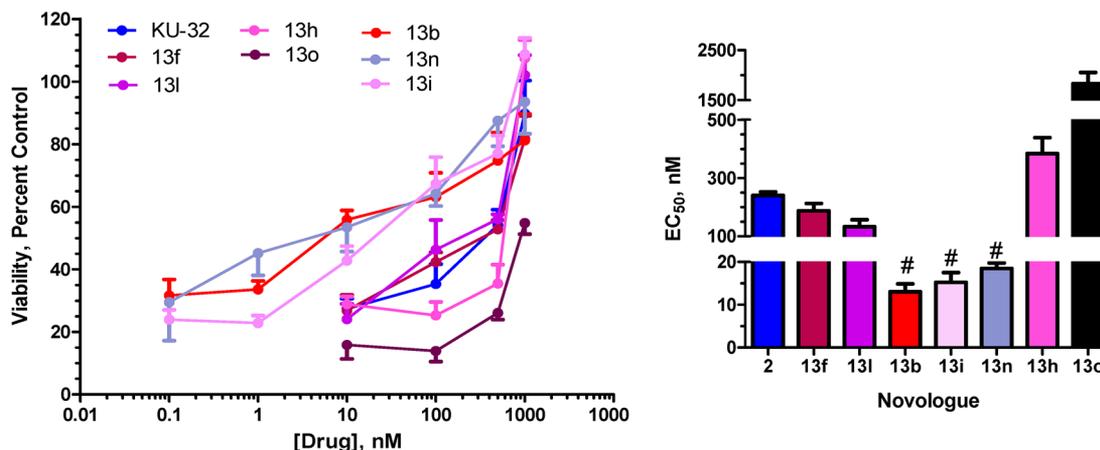
<sup>a</sup>In the presence of 1 μM of each novologue +20 mM excess glucose. Viability in the presence of 20 mM excess glucose + DMSO was 54% ± 2. <sup>b</sup>#, *p* < 0.05 versus glucose + DMSO. <sup>c</sup>\*, *p* < 0.05 versus glucose + compound 2 (*n* = 6–24) per novologue.

analogue (13o) was unable to protect against glucose-induced toxicity and was also significantly less protective than the corresponding 4-pyridine analogue, 13p, 2, and the unsubstituted phenyl analogue, 13a. Although the 4-pyridine-containing analogue (13p) demonstrated a modestly improved neuroprotective activity when compared to the simple phenyl analogue 13a, this difference in efficacy was not significant.

Neuroprotective activity was also determined for the cyclohexene-containing novologues (22a–b) that contain the fluoro or trifluoromethane substituent at the *meta*-position of ring B. In general, cyclohexene-containing analogues 22a–b were less efficacious than the corresponding derivatives that contain a flexible side chain (13b versus 22a, and 13f versus 22b). Although not statistically different, novologue 22a (*meta*-F) exhibited slightly better cytoprotective activity than analogue 22b (*meta*-CF<sub>3</sub>), which follows the same trend observed for flexible acetamide-containing compounds (13b versus 13f). Although these data are inconsistent with our hypothesis that accommodation of the hydrophobic pocket would improve efficacy, the cyclohexene ring may exceed the space allowed in this binding cleft.

The data in Table 1 clearly support that the majority of novologues synthesized decrease neuronal toxicity induced by hyperglycemic stress. Although some of these compounds appear more effective than compound 2 at 1 μM, the differences were relatively minor. Therefore, to further scrutinize their efficacy, compounds exhibiting high neuroprotective activity were further evaluated for determination of EC<sub>50</sub> values. Because the difference in efficacy for novologues with *meta*-F and *meta*-CF<sub>3</sub> substitutions on 13b and 13f were not significantly different from 2 or each other at 1 μM, the EC<sub>50</sub> values for these compounds were determined alongside 13h, 13l, 13n, and 13o. As shown in parts A and B of Figure 4, EC<sub>50</sub> values were significantly improved upon closer inspection and clear distinctions were obtained. Novologue 13b yielded an EC<sub>50</sub> value (13.0 ± 3.6 nM) that was approximately 14-fold lower than compound 2 (240.2 ± 42.5 nM) or 13f (187.7 ± 43.5 nM). Similar results were also observed for novologue 13n, which exhibited an EC<sub>50</sub> value of 18.4 ± 3.2 nM. In contrast, novologue 13h which manifested similar efficacy to compound 2 at 1 μM, exhibited an EC<sub>50</sub> of 384 ± 108 nM, approximately 1.6-fold greater than 2.

The data in Figure 4 demonstrate that novologues 13b, 13i, and 13n are more cytoprotective than the initial lead compound 2. Since we have previously shown that the cytoprotective activity manifested by compound 2 requires Hsp70,<sup>10</sup> we determined the ability of the novologues to induce Hsp70 relative to 2. To this end, a luciferase reporter assay was



**Figure 4.** Determination of EC<sub>50</sub> of select novologues. (Left Panel) DRG sensory neurons were incubated in the absence or presence of 0.1–1000 nM of the indicated novologue overnight and then subjected to 4 h of hyperglycemia. Cell viability was measured as described in General Experimental Methods and the data expressed as percent of normoglycemic controls. Under hyperglycemic conditions and in the absence of any novologues, cell viability was 20% ± 7. (Right Panel) The EC<sub>50</sub> was determined using the EC<sub>anything</sub> function of GraphPad Prism 5.0, and the mean ± SEM (*n* = 3–8) is shown. #, *p* < 0.05 versus compound 2.

developed in which the expression of luciferase is driven by the human Hsp70 promoter containing two heat shock binding elements.<sup>24</sup> Because primary sensory neurons transfect poorly, an immortalized sensory neuron cell line (50B11 cells) was used for transfection.<sup>25</sup> Importantly, 50B11 cells have a very low basal level of Hsp70 expression, similar to primary sensory neurons.<sup>26</sup> Twenty-four hours after transfection with the reporter, cells were reseeded into 12-well plates and incubated for an additional 6 h. The cells were then treated for 16 h with 10–1000 nM of the indicated novologues, cell lysates were prepared, luciferase activity assessed, and luminescence normalized to total protein per well. Consistent with its increased efficacy in protecting against glucotoxicity, **13b** was more effective than **2** at activation of the Hsp70 promoter (Figure 5A) and also increased expression of Hsp70 protein at lower concentrations relative to compound **2** (Figure 5B). Although **13i** had a similar  $EC_{50}$  as **13b** in preventing glucotoxicity, it only activated the Hsp70 promoter at 1  $\mu$ M, and the magnitude of this effect was no better than either **2** or **13b**. However, it was surprising that despite the low  $EC_{50}$  of **13n** in protecting against glucotoxicity, **13n** did not increase luciferase activity at any concentration tested nor did it increase Hsp70 protein expression as effectively as **2** or **13b**. These results suggest that **13n** likely affects Hsp70 levels indirectly and that the mechanism for neuroprotection may be distinct from that of related novologues.

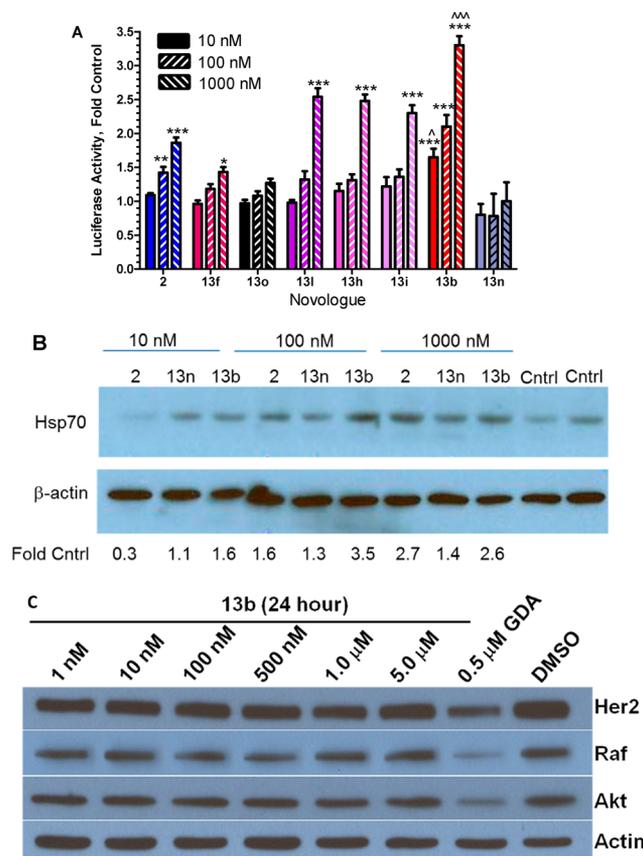
Lastly, we mentioned that an attractive property of the modified novobiocin scaffold of **2** is that it induces Hsp70 at concentrations well below those needed to inhibit Hsp90's protein folding ability.<sup>10</sup> Therefore, to confirm that this new scaffold manifests similar activity, **13b** was evaluated against MCF-7 breast cancer cells that are highly reliant upon the Hsp90 protein folding machinery. As can be observed, no client protein degradation occurred at concentrations up to 5  $\mu$ M, indicating the potential for a large therapeutic window for this scaffold as well.

## CONCLUSION

Using the recently reported model for the Hsp90 C-terminal binding site, a novologue scaffold was designed to afford putative interactions with previously unoccupied regions of the binding pocket, including Lys539. Through systematic replacement of substituents on the novologue B-ring (see Table 2), compound **13b** was identified as a neuroprotective agent that exhibited  $\sim$ 14-fold greater efficacy against glucose-induced toxicity than the lead compound **2**. The concentration of **13b** needed to manifest neuroprotective activity correlated well with its ability to induce Hsp70 levels, therefore linking cytoprotection to Hsp70 induction. When combined, these data demonstrate the rationally designed novologue scaffold provides a promising platform on which diversification of the B-ring can lead to compounds that exhibit better neuroprotective activities than first generation analogues.

## GENERAL EXPERIMENTAL METHODS

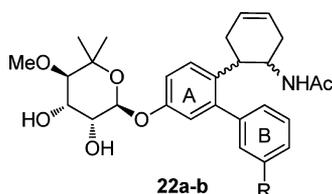
**Preparation of Embryonic Dorsal Root Ganglion (DRG) Neuron Cultures.** DRG from embryonic day 15–18 Sprague–Dawley rat pups were harvested into Leibovitz's L15 medium (L15) and dissociated with 0.25% trypsin for 30 min at 37 °C. The ganglia were sedimented at 1000g for 5 min, resuspended in growth media (phenol red free Neurobasal medium (Gibco, Grand Island, NY) containing 25 mM glucose, 1 $\times$  B-27 additive, 50 ng/mL nerve growth factor (NGF) (Harlan Bioscience, Indianapolis, IN), 4 mM glutamine,



**Figure 5.** Induction of Hsp70 by select novologues in the absence of client protein degradation. (A) 50B11 cells were transfected with a luciferase reporter whose expression was driven by the human Hsp70 promoter. The cells were treated with the indicated concentration of select novologues for 16 h and luciferase activity assessed. \*\*,  $p < 0.01$  and \*\*\*,  $p < 0.001$  versus control; ^  $p < 0.05$  and ^^,  $p < 0.001$  versus compound **2** at same concentration. (B) DRG sensory neurons were incubated in the presence of DMSO (Cntrl) or 10–1000 nM of the indicated novologue overnight and then subjected to 4 h of hyperglycemia. The neurons were harvested and Hsp70, and  $\beta$ -actin levels were determined by immunoblot analysis. Band intensity was quantified using Image J, Hsp70 expression was normalized to the level of  $\beta$ -actin and expressed as a fold control. (C) MCF7 cells were treated with the indicated concentrations of **13b** for 24 h, cell lysates were prepared, and the levels of the Hsp90 client proteins, Her2, Raf, and Akt determined by immunoblot analysis. As a positive control, some cells were treated with 500 nM geldanamycin (GDA) to induce client protein degradation. The level of  $\beta$ -actin verified equivalent protein loading.

100 U/mL penicillin, and 100  $\mu$ g/mL streptomycin) and triturated with a fire-polished glass pipet. The cells were cultured on collagen-coated (0.1 mg/mL collagen followed by overnight air drying in a laminar flow hood) black-walled 96-well plates (Corning Incorporated Corning, NY) at a seeding density of  $(2-3) \times 10^4$  cells per well. DRG neurons were refed the next day with fresh growth media containing 40  $\mu$ M fluorodeoxyuridine and 10  $\mu$ M cytosine  $\beta$ -D-arabinooside (both from Sigma Aldrich, St. Louis, MO) for 2 days to remove proliferating cells. Experiments were performed on DRG neurons on the third day in culture after placing the cells in fresh growth medium.

**Glucotoxicity Assay.** As immature DRG are susceptible to hyperglycemia-induced death,<sup>27</sup> an additional 20 mM glucose was added to the growth medium (yielding a total of 45 mM glucose) for 4 h. Preliminary experiments found that 20 mM excess glucose for 4 h was sufficient to induce a reproducible 40–50% loss in neuronal viability. As a result, the toxicity induced by the acute change in

**Table 2. Cell Viability Data of Cyclohexene Containing Analogues**

entry	R	% of cell viability <sup>a</sup>
2		86 ± 2
22a	F	78 ± 18 <sup>#b</sup>
22b	CF <sub>3</sub>	69 ± 15 <sup>#,*,b,c</sup>

<sup>a</sup>In the presence of 1 μM novologue +20 mM excess glucose. Viability in the presence of 20 mM excess glucose + DMSO was 54% ± 2. <sup>b</sup>#, *p* < 0.05 versus glucose + DMSO. <sup>c</sup>\*, *p* < 0.05 versus glucose + 2 (*n* = 8) per novologue.

glucose concentration makes it a useful model for drug screening.<sup>10,28</sup> Given the short time frame that the neurons are grown in vitro, they are not pure neuronal cultures but instead highly enriched. Importantly, the contaminating SCs that remain in the culture are resistant to glucose-induced death as we and others have reported previously.<sup>6</sup> Unfortunately, the use of highly purified cultures is problematic because the cells extend neurites and establish connections with each other, thus becoming resistant to hyperglycemia-induced death.<sup>6a</sup>

DRG neurons were incubated overnight with the novologues in the presence of Neurobasal medium, 50 ng/mL NGF, and antibiotics only. To monitor the efficiency of the novologues in protecting DRG neurons against glucotoxicity, we made use of Calcein AM (Invitrogen, Carlsbad, CA) to measure cell viability. Hydrolysis of calcein AM to a fluorescent product can only occur in live cells. Excess glucose was added to the cultures for 4 h, and cell viability was measured by incubating the cells with 2 μM calcein AM for 30 min in the dark at 37 °C. Fluorescence was then measured using a plate reader with excitation and emission wavelengths set to 485 nm and 520 nm, respectively. The arbitrary fluorescence readings were normalized to the total amount of protein from each respective well of the neuronal cultures. The protein concentrations in each well were determined using the DC protein assay (Bio-Rad). Significant differences in the efficacy of the novologues for increasing cell viability were determined using a Kruskal–Wallis nonparametric ANOVA and Dunn's post-test.

**Luciferase Reporter Assay and Client Protein Degradation.** A 1.5 kb region upstream of the start codon of the human *HSPA1A* gene was synthesized by GeneArt (Life Technologies, Grand Island, NY) and 5' *KpnI* and 3' *SacI* sites added to direct cloning into the pGL3 basic luciferase reporter plasmid. DNA sequencing verified the integrity of the promoter sequence and the presence of two heat shock elements. 50B11 cells<sup>25</sup> were grown in 10 cm dishes in DMEM containing 25 mM glucose, 10% FCS, and 5 μg/mL blasticidin. The cells were transfected using lipofectamine and after 24 h were reseeded into 24 well plates at a density of 2 × 10<sup>5</sup> cells per well. The cells were permitted to attach to the plate for 6 h in growth medium and treated with the indicated concentrations of the various novologues for 16 h. Luciferase activity was assessed and normalized to the total protein concentration of each well. Results shown are from triplicate wells obtained in at least three separate experiments. Preliminary experiments validated that the reporter was strongly activated as expected by either heat shock (~10-fold) or 250 nM geldanamycin (~4–5-fold). Client protein degradation in MCF7 cells was performed as we have previously described.<sup>10</sup>

**Molecular Modeling.** Surflex-Dock in Sybyl v8.0 was used for molecular modeling and docking studies. A homology model of Hsp90α based on the open HtpG SAXS structure was used as the receptor, while the protomol was generated using docked Novobiocin as described in reference.<sup>13</sup> The energy minimized molecules were

then docked with 10 different starting conformations while rotation of rotatable bonds was unrestricted. Visual interpretation and figure preparation were then carried out in Pymol.

**Chemistry General.** <sup>1</sup>H NMR were recorded at 400 or 500 MHz (Bruker DRX-400 Bruker with a H/C/P/F QNP gradient probe) spectrometer and <sup>13</sup>C NMR spectra were recorded at 125 MHz (Bruker DRX 500 with broadband, inverse triple resonance, and high resolution magic angle spinning HR-MA probe spectrometer); chemical shifts are reported in δ (ppm) relative to the internal reference chloroform-*d* (CDCl<sub>3</sub>, 7.27 ppm). FAB (HRMS) spectra were recorded with a LCT Premier (Waters Corp., Milford, MA). The purity of all compounds was determined to be >95% as determined by <sup>1</sup>H NMR and <sup>13</sup>C NMR spectra, unless otherwise noted. The most active five compounds were verified for >95% purity by HPLC analyses. TLC was performed on glass backed silica gel plates (Uniplate) with spots visualized by UV light. All solvents were reagent grade and, when necessary, were purified and dried by standard methods. Concentration of solutions after reactions and extractions involved the use of a rotary evaporator operating at reduced pressure.

**5-(Benzyloxy)-2-formylphenyl trifluoromethanesulfonate (7).** Triethylamine (1.02 mL, 7.35 mmol) followed by triflic anhydride (1.38 mL, 6.35 mmol) were added simultaneously to a phenol 6 (1.12 g, 4.91 mmol) in anhydrous CH<sub>2</sub>Cl<sub>2</sub> (10 mL) at 0 °C. Upon completion of the reaction, quenched by the addition of water (50 mL), extracted with CH<sub>2</sub>Cl<sub>2</sub> (3 × 15 mL), washed with saturated aqueous sodium chloride solution, dried over anhydrous Na<sub>2</sub>SO<sub>4</sub>, filtered, and concentrated. The residue was purified by column chromatography (SiO<sub>2</sub>, 4:1, Hex:EtOAc) to afford triflate 7 as a yellow oil (1.06 g, 60%).

**General Procedure for Suzuki Coupling Reaction of Triflate 3 and Boronic Acids 4a–p.** 5-(Benzyloxy)-[1,1'-biphenyl]-2-carbaldehyde (**8a**). Tetrakis(triphenylphosphine)palladium(0) (70.4 mg, 0.068 mmol) was added to a mixture of triflate 7 (0.246 g, 0.68 mmol), phenylboronic acid 4a (92 mg, 0.75 mmol), and K<sub>2</sub>CO<sub>3</sub> (0.169 g, 1.2 mmol) in DMF (6.8 mL) under argon atmosphere in a sealed tube. The resulting reaction mixture was sealed and heated to reflux for 16 h. The reaction was cooled to room temperature, quenched with saturated sodium bicarbonate, extracted with EtOAc (3 × 5 mL), washed with saturated aqueous sodium chloride, dried over anhydrous Na<sub>2</sub>SO<sub>4</sub>, filtered, and concentrated. The crude product was purified by column chromatography (SiO<sub>2</sub>, 3:1, Hex:EtOAc) to afford **8a** (0.16 g, 0.56 mmol, 82%) as an amorphous solid. <sup>1</sup>H NMR (400 MHz, CDCl<sub>3</sub>) δ 9.90 (s, 1H), 8.08 (d, *J* = 8.7 Hz, 1H), 7.55–7.34 (m, 10H), 7.11 (d, *J* = 8.7 Hz, 1H), 7.03 (d, *J* = 2.4 Hz, 1H), 5.19 (s, 2H). <sup>13</sup>C NMR (100 MHz, CDCl<sub>3</sub>) δ 191.2, 162.8, 148.6, 137.8, 136.0, 130.0, 128.8, 128.4, 127.6, 116.3, 114.7, 70.4. HRMS (FAB) *m/z* [M + Na<sup>+</sup>] for C<sub>20</sub>H<sub>16</sub>O<sub>2</sub>Na: calcd, 311.1042; found, 311.1046.

**5-(Benzyloxy)-3'-fluoro-[1,1'-biphenyl]-2-carbaldehyde (8b).** Using 3-fluorophenylboronic acid. <sup>1</sup>H NMR (500 MHz, CDCl<sub>3</sub>) δ 9.85 (d, *J* = 0.7 Hz, 1H), 8.03 (d, *J* = 8.7 Hz, 1H), 7.49–7.33 (m, 6H), 7.20–7.13 (m, 2H), 7.13–7.08 (m, 2H), 7.03 (d, *J* = 2.5 Hz, 1H), 5.15 (s, 2H). <sup>13</sup>C NMR (120 MHz, CDCl<sub>3</sub>) δ 190.7, 162.9, 161.7, 147.2, 140.1, 136.0, 130.5, 129.0, 128.6, 127.8, 126.0, 117.1, 116.9, 116.4, 115.5, 115.1, 70.6. HRMS *m/z* [M + Na<sup>+</sup>] for C<sub>20</sub>H<sub>15</sub>FO<sub>2</sub>Na: calcd, 329.0948; found, 329.0952.

**5-(Benzyloxy)-4'-fluoro-[1,1'-biphenyl]-2-carbaldehyde (8c).** Using 4-fluorophenylboronic acid. <sup>1</sup>H NMR (400 MHz, CDCl<sub>3</sub>) δ 9.84 (s, 1H), 8.06 (dd, *J* = 8.7, 1.0 Hz, 1H), 7.49–7.40 (m, 4H), 7.40–7.32 (m, 3H), 7.21–7.13 (m, 2H), 7.12–7.06 (dd, *J* = 8.0, 2.5 Hz, 1H), 7.03 (d, *J* = 2.2 Hz, 1H), 5.17 (s, 2H). <sup>13</sup>C NMR (100 MHz, CDCl<sub>3</sub>) δ 190.9, 162.8, 147.4, 136.0, 131.7, 131.6, 130.5, 128.8, 128.5, 127.7, 127.6, 116.5, 115.6, 115.4, 114.7, 70.4. HRMS *m/z* [M + Na<sup>+</sup>] for C<sub>20</sub>H<sub>15</sub>FO<sub>2</sub>Na: calcd, 329.0948; found, 329.0944.

**5-(Benzyloxy)-2'-chloro-[1,1'-biphenyl]-2-carbaldehyde (8d).** Using 2-chlorophenylboronic acid. <sup>1</sup>H NMR (500 MHz, CDCl<sub>3</sub>) δ 9.70 (s, 1H), 8.08 (d, *J* = 8.7 Hz, 1H), 7.55–7.49 (m, 1H), 7.49–7.32 (m, 8H), 7.17–7.12 (dd, *J* = 8.6, 2.5 Hz, 1H), 6.99 (d, *J* = 2.6 Hz, 1H), 5.16 (s, 2H). <sup>13</sup>C NMR (125 MHz, CDCl<sub>3</sub>) δ 190.3, 162.9, 145.1, 136.8, 135.9, 133.5, 131.6, 130.0, 129.8, 129.6, 128.8, 128.4, 127.6,

127.6, 126.9, 116.7, 115.1, 70.4. HRMS  $m/z$  [ $M + Na^+$ ] for  $C_{20}H_{15}ClO_2Na$ , calcd, 345.0658; found, 345.0653.

**5-(Benzyloxy)-3'-chloro-[1,1'-biphenyl]-2-carbaldehyde (8e).** Using 3-chlorophenylboronic acid.  $^1H$  NMR (400 MHz,  $CDCl_3$ )  $\delta$  9.85 (s, 1H), 8.04 (d,  $J = 8.7$  Hz, 1H), 7.49–7.33 (m, 8H), 7.26 (m, 1H), 7.13–7.07 (dd,  $J = 8.3, 2.8$  Hz, 1H), 6.96 (d,  $J = 2.5$  Hz, 1H), 5.17 (s, 2H).  $^{13}C$  NMR (100 MHz,  $CDCl_3$ )  $\delta$  190.4, 162.8, 146.8, 139.7, 135.9, 134.5, 130.5, 129.8, 129.7, 128.8, 128.5, 128.4, 128.3, 127.6, 127.5, 116.3, 115.0, 70.4. HRMS  $m/z$  [ $M + Cl^-$ ] for  $C_{20}H_{15}Cl_2O_2$ : calcd, 341.0505; found, 341.0508.

**5-(Benzyloxy)-3'-(trifluoromethyl)-[1,1'-biphenyl]-2-carbaldehyde (8f).** Using 3-(trifluoromethyl)phenylboronic acid.  $^1H$  NMR (400 MHz,  $CDCl_3$ )  $\delta$  9.82 (s, 1H), 8.05 (m, 1H), 7.72 (m, 1H), 7.67–7.64 (td,  $J = 1.6, 0.8$  Hz, 1H), 7.64–7.53 (m, 2H), 7.50–7.35 (m, 5H), 7.15–7.11 (dd,  $J = 8.7, 2.2$  Hz, 1H), 6.96 (d,  $J = 2.5$  Hz, 1H), 5.19 (s, 2H).  $^{13}C$  NMR (100 MHz,  $CDCl_3$ )  $\delta$  190.4, 163.0, 146.8, 138.8, 135.9, 133.4, 131.0, 130.9, 129.0, 129.0, 128.6, 127.8, 127.6, 126.6, 126.5, 125.2, 116.7, 115.2, 70.6. HRMS  $m/z$  [ $M + Na^+$ ] for  $C_{21}H_{15}F_3O_2Na$ : calcd, 379.0922; found, 379.0926.

**5-(Benzyloxy)-4'-(trifluoromethyl)-[1,1'-biphenyl]-2-carbaldehyde (8g).** Using 4-(trifluoromethyl)phenylboronic acid.  $^1H$  NMR (400 MHz,  $CDCl_3$ )  $\delta$  9.84 (s, 1H), 8.06 (d,  $J = 8.7$  Hz, 1H), 7.75 (d,  $J = 8.0$  Hz, 2H), 7.55–7.49 (m, 2H), 7.49–7.34 (m, 6H), 7.17–7.12 (dd,  $J = 9.1, 2.2$  Hz, 1H), 6.98 (d,  $J = 2.5$  Hz, 1H), 5.19 (s, 2H).  $^{13}C$  NMR (100 MHz,  $CDCl_3$ )  $\delta$  190.2, 162.9, 146.7, 141.7, 135.9, 130.8, 130.3, 128.9, 128.6, 127.7, 127.5, 125.5, 125.4, 122.8, 116.6, 115.1, 70.5. HRMS  $m/z$  [ $M + H^+$ ] for  $C_{21}H_{16}F_3O_2$ : calcd, 357.1097; found, 357.1096.

**5-(Benzyloxy)-2'-(methylthio)-[1,1'-biphenyl]-2-carbaldehyde (8h).** Using 2-(methylthio)phenylboronic acid.  $^1H$  NMR (400 MHz,  $CDCl_3$ )  $\delta$  9.62 (s, 1H), 8.05 (d,  $J = 8.7$  Hz, 1H), 7.47–7.32 (m, 6H), 7.30–7.23 (m, 2H), 7.24–7.20 (m, 1H), 7.13–7.09 (m, 1H), 6.93–6.90 (m, 1H), 5.17 (s, 2H), 2.36 (d,  $J = 1.1$  Hz, 3H).  $^{13}C$  NMR (100 MHz,  $CDCl_3$ )  $\delta$  190.8, 163., 146.3, 138.4, 136.2, 136.1, 130.4, 129.5, 129.1, 128.8, 128.4, 127.8, 127.7, 124.7, 124.6, 116.4, 115.3, 70.4, 15.6. HRMS  $m/z$  [ $M + H^+$ ] for  $C_{21}H_{18}O_2SNa$ : calcd, 357.0920; found, 357.0923.

**5-(Benzyloxy)-2'-methoxy-[1,1'-biphenyl]-2-carbaldehyde (8i).** Using 2-methoxyphenylboronic acid.  $^1H$  NMR (500 MHz,  $CDCl_3$ )  $\delta$  9.73 (s, 1H), 8.07 (d,  $J = 8.7$  Hz, 1H), 7.48–7.39 (m, 5H), 7.37 (d,  $J = 6.5$  Hz, 1H), 7.32–7.27 (m, 1H), 7.13–7.07 (m, 2H), 7.02 (d,  $J = 8.3$  Hz, 1H), 6.98–6.95 (dd,  $J = 2.4, 1.1$  Hz, 1H), 5.15 (s, 2H), 3.75 (s, 3H).  $^{13}C$  NMR (125 MHz,  $CDCl_3$ )  $\delta$  191.5, 163.1, 156.6, 144.5, 136.2, 131.4, 130.1, 129.2, 128.8, 128.4, 127.9, 127.7, 126.8, 121.0, 116.9, 114.5, 110.8, 70.3, 55.5. HRMS  $m/z$  [ $M + H^+$ ] for  $C_{21}H_{19}O_3$ : calcd, 319.1329; found, 319.1333.

**5-(Benzyloxy)-3'-methoxy-[1,1'-biphenyl]-2-carbaldehyde (8j).** Using 3-methoxyphenylboronic acid.  $^1H$  NMR (400 MHz,  $CDCl_3$ )  $\delta$  9.93 (s, 1H), 8.06 (d,  $J = 9.0$  Hz, 1H), 7.52–7.35 (m, 6H), 7.10 (d,  $J = 8.6$  Hz, 1H), 7.05–6.93 (m, 4H), 5.20 (s, 2H), 3.89 (s, 3H). HRMS  $m/z$  [ $M + Na^+$ ] for  $C_{21}H_{18}O_3Na$ : calcd, 341.1154; found, 341.1150.

**5-(Benzyloxy)-3'-methyl-[1,1'-biphenyl]-2-carbaldehyde (8k).** Using 3-methylphenylboronic acid.  $^1H$  NMR (500 MHz,  $CDCl_3$ )  $\delta$  9.85 (d,  $J = 0.9$  Hz, 1H), 8.03 (d,  $J = 8.6$  Hz, 1H), 7.49–7.39 (m, 3H), 7.39–7.32 (m, 2H), 7.27 (d,  $J = 8.1$  Hz, 1H), 7.22–7.16 (m, 2H), 7.09–7.05 (ddd,  $J = 8.8, 2.6, 0.9$  Hz, 1H), 6.98 (d,  $J = 2.5$  Hz, 1H), 5.15 (s, 2H), 2.43 (s, 3H).  $^{13}C$  NMR (125 MHz,  $CDCl_3$ )  $\delta$  191.4, 162.8, 148.9, 138.3, 137.9, 136.2, 130.9, 130.1, 129.2, 128.9, 128.5, 127.8, 127.3, 116.3, 114.8, 70.5, 21.7. HRMS  $m/z$  [ $M + H^+$ ] for  $C_{21}H_{18}O_2Na$ : calcd, 325.1205; found, 325.1217.

**5-(Benzyloxy)-3'-(morpholinomethyl)-[1,1'-biphenyl]-2-carbaldehyde (8l).** Using 3-(4-morpholinomethyl)phenylboronic acid pinacol ester.  $^1H$  NMR (400 MHz,  $CDCl_3$ )  $\delta$  9.87 (s, 1H), 8.83 (d,  $J = 8.7$  Hz, 1H), 7.47–7.31 (m, 7H), 7.32–7.24 (m, 1H), 7.12–7.04 (dd,  $J = 8.7, 2.5$  Hz, 1H), 7.05 (d,  $J = 2.5$  Hz, 1H), 5.17 (s, 2H), 3.79–3.68 (t,  $J = 4.6$  Hz, 4H), 3.56 (s, 3H), 2.49 (d,  $J = 6.5$  Hz, 4H).  $^{13}C$  NMR (100 MHz,  $CDCl_3$ )  $\delta$  191.0, 162.7, 148.5, 138.3, 137.8, 136.0, 130.7, 130.2, 129.1, 128.8, 128.4, 127.6, 127.6, 116.4, 114.5, 70.4, 67.1, 63.2, 53.7. HRMS  $m/z$  [ $M + Na^+$ ] for  $C_{25}H_{25}NO_3Na$ , calcd: 410.1726; found, 410.1730.

**5-(Benzyloxy)-4'-hydroxy-[1,1'-biphenyl]-2-carbaldehyde (8m).** Used 4-hydroxyphenylboronic acid. Partially purified biaryl phenol was treated with TBSCl (1.2 equiv) and imidazole (3 equiv) in  $CH_2Cl_2$  and stirred for 2 h at room temperature. After reaction was completed by TLC, the resulting reaction mixture was concentrated. The crude product was purified by column chromatography ( $SiO_2$ , 4:1, Hex:EtOAc) to afford **8m** (94%) as an amorphous solid.  $^1H$  NMR (500 MHz,  $CDCl_3$ )  $\delta$  9.89 (s, 1H), 8.03 (d,  $J = 8.7$  Hz, 1H), 7.52–7.33 (m, 5H), 7.26 (dd,  $J = 6.6, 1.8$  Hz, 2H), 7.05 (dd,  $J = 8.7, 2.3$  Hz, 1H), 7.02–6.93 (m, 3H), 5.17 (s, 2H), 1.05 (s, 9H), 0.29 (s, 6H).  $^{13}C$  NMR (125 MHz,  $CDCl_3$ )  $\delta$  191.2, 162.7, 156.0, 148.4, 136.1, 131.2, 130.6, 130.0, 128.7, 128.3, 127.6, 127.5, 120.0, 116.1, 114.3, 70.3, 25.7, 18.3, 4.3. ESI-HRMS  $m/z$  [ $M + Na^+$ ] for  $C_{26}H_{30}NaO_3Si$ : calcd, 441.5899, found 441.5896.

**2-(Benzo[d][1,3]dioxol-5-yl)-4-(benzyloxy)benzaldehyde (8n).** Using 3,4-(methylenedioxy)phenylboronic acid.  $^1H$  NMR (500 MHz,  $CDCl_3$ )  $\delta$  9.90 (s, 1H), 8.08 (d,  $J = 8.7$  Hz, 1H), 7.48–7.39 (m, 4H), 7.39–7.35 (m, 1H), 7.06 (d,  $J = 8.6$  Hz, 1H), 6.97 (d,  $J = 2.5$  Hz, 1H), 6.91–6.86 (m, 2H), 6.83–6.79 (m, 1H), 6.03 (s, 2H), 5.15 (s, 2H).  $^{13}C$  NMR (125 MHz,  $CDCl_3$ )  $\delta$  191.2, 162.8, 148.2, 147.9, 147.9, 136.1, 131.6, 130.2, 128.8, 128.4, 127.7, 127.6, 124.0, 116.2, 114.5, 110.3, 108.3, 101.5, 70.4. HRMS (FAB)  $m/z$  [ $M + Na^+$ ] for  $C_{21}H_{16}O_4Na$ : calcd, 355.0941; found, 355.0935.

**4-(Benzyloxy)-2-(pyridin-3-yl)benzaldehyde (8o).**  $^1H$  NMR (400 MHz,  $CDCl_3$ )  $\delta$  9.79 (s, 1H), 8.65 (dd,  $J = 5.1, 8.3$  Hz), 8.01 (d, 1H,  $J = 8.8$  Hz), 7.67 (m, 1H), 7.48–7.26 (m, 6H), 7.09 (dd, 1H,  $J = 2.4, 8.7$  Hz), 6.93 (d, 1H,  $J = 2.4$  Hz), 5.14 (s, 2H).  $^{13}C$  NMR (125 MHz,  $CDCl_3$ )  $\delta$  187.8, 165.3, 160.5, 135.8, 131.2, 129.0, 128.7, 127.8, 120.0, 109.5, 102.1, 91.0, 70.8. HRMS (FAB)  $m/z$  [ $M + H^+$ ] for  $C_{19}H_{16}NO_2$ : calcd, 290.1181; found, 290.1177.

**4-(Benzyloxy)-2-(pyridin-4-yl)benzaldehyde (8p).**  $^1H$  NMR (500 MHz,  $CDCl_3$ )  $\delta$  9.82 (s, 1H), 8.67 (d,  $J = 5.9$  Hz, 2H), 8.02 (d,  $J = 8.7$  Hz, 1H), 7.49–7.33 (m, 6H), 7.30 (d,  $J = 6.0$  Hz, 1H), 7.15–7.10 (dd,  $J = 8.6, 2.6$  Hz, 1H), 6.95 (d,  $J = 2.6$  Hz, 1H), 5.15 (s, 2H).  $^{13}C$  NMR (125 MHz,  $CDCl_3$ )  $\delta$  189.7, 162.9, 149.8, 145.8, 145.2, 135.7, 131.0, 128.8, 128.5, 127.6, 127.1, 124.6, 116.3, 115.4, 70.5. HRMS (FAB)  $m/z$  [ $M + H^+$ ] for  $C_{19}H_{16}NO_2$ : calcd, 290.1181; found, 290.1183.

#### General Procedure for Henry Reaction of Compounds 8a–p.

**(E)-5-(Benzyloxy)-2-(2-nitrovinyl)-1,1'-biphenyl (9a).** Nitromethane (1.4 mL) was added to a mixture of aldehyde **8a** (0.16 g, 0.56 mmol) and ammonium acetate (77 mg, 1.0 mmol) and heated to 50 °C. Upon completion (~15–30 min), the reaction mixture was cooled to RT and purified without workup by column chromatography ( $SiO_2$ , 3:1, Hex:EtOAc) to afford nitrostyrene **9a** as a yellow oil (182 mg, 0.55 mmol, 98%).  $^1H$  NMR (400 MHz,  $CDCl_3$ )  $\delta$  8.02 (d,  $J = 13.6$  Hz, 1H), 7.64 (d,  $J = 9.5$  Hz, 1H), 7.50–7.35 (m, 10H), 7.31 (d,  $J = 2.1$  Hz, 2H), 7.04 (d,  $J = 2.5$  Hz, 1H), 5.15 (s, 2H).  $^{13}C$  NMR (100 MHz,  $CDCl_3$ )  $\delta$  161.8, 146.1, 138.1, 136.4, 136.3, 135.5, 131.8, 131.7, 129.9, 129.2, 128.8, 128.0, 121.3, 117.3, 116.3, 116.0, 115.6, 70.7. HRMS (FAB)  $m/z$  [ $M + Na^+$ ] for  $C_{21}H_{18}NO_3$ : calcd, 332.1281; found, 332.1290.

**(E)-5-(Benzyloxy)-3'-fluoro-2-(2-nitrovinyl)-1,1'-biphenyl (9b).**  $^1H$  NMR (400 MHz,  $CDCl_3$ )  $\delta$  8.07 (d,  $J = 13.5$  Hz, 1H), 7.65 (d,  $J = 8.7$  Hz, 1H), 7.49–7.35 (m, 7H), 7.20–7.13 (ddd,  $J = 9.3, 7.9, 2.6$  Hz, 1H), 7.09–7.03 (m, 2H), 7.02 (d,  $J = 2.8$  Hz, 2H), 5.16 (s, 2H).  $^{13}C$  NMR (100 MHz,  $CDCl_3$ )  $\delta$  164.0, 161.5, 145.4, 141.4, 137.6, 136.1, 136.0, 130.5, 130.4, 129.6, 128.6, 127.7, 125.7, 121.0, 116.9, 116.6, 115.6, 115.4, 70.5. HRMS  $m/z$  [ $M + H^+$ ] for  $C_{21}H_{17}FNO_3$ : calcd, 350.1187; found, 350.1185.

**(E)-5-(Benzyloxy)-4'-fluoro-2-(2-nitrovinyl)-1,1'-biphenyl (9c).**  $^1H$  NMR (400 MHz,  $CDCl_3$ )  $\delta$  8.08 (d,  $J = 13.6$  Hz, 1H), 7.64 (d,  $J = 8.7$  Hz, 1H), 7.50–7.34 (m, 6H), 7.32–7.24 (m, 2H), 7.23–7.14 (t,  $J = 8.3$  Hz, 2H), 7.10–7.00 (m, 2H), 5.17 (s, 2H).  $^{13}C$  NMR (100 MHz,  $CDCl_3$ )  $\delta$  161.5, 145.7, 137.8, 136.1, 136.0, 131.5, 131.4, 129.6, 128.9, 128.5, 127.7, 121.0, 117.0, 115.9, 115.7, 115.3, 70.4. HRMS  $m/z$  [ $M + Na^+$ ] for  $C_{21}H_{16}FNO_3Na$ : calcd, 372.1006; found, 372.1011.

**(E)-5-(Benzyloxy)-2'-chloro-2-(2-nitrovinyl)-1,1'-biphenyl (9d).**  $^1H$  NMR (500 MHz,  $CDCl_3$ )  $\delta$  7.85–7.75 (m, 1H), 7.74–7.66 (m, 1H), 7.55 (m, 1H), 7.53–7.34 (m, 8H), 7.31 (d,  $J = 5.3$  Hz, 1H), 7.17 (d,  $J = 8.3$  Hz, 1H), 7.01 (t,  $J = 2.0$  Hz, 1H), 5.20–5.11 (m, 2H).  $^{13}C$  NMR

(125 MHz, CDCl<sub>3</sub>)  $\delta$  161.4, 143.8, 137.7, 137.0, 135.9, 133.2, 131.4, 130.0, 130.0, 129.3, 128.7, 128.3, 127.6, 127.1, 123.4, 121.5, 117.1, 115.6, 70.3. HRMS  $m/z$  [M + H<sup>+</sup>] for C<sub>21</sub>H<sub>17</sub>ClNO<sub>3</sub>: calcd, 366.0892; found, 366.0895.

**5-(Benzyloxy)-3'-chloro-2-(2-nitrovinyl)-1,1'-biphenyl (9e).** <sup>1</sup>H NMR (400 MHz, CDCl<sub>3</sub>)  $\delta$  7.95 (d,  $J$  = 13.5 Hz, 1H), 7.64 (d,  $J$  = 8.8 Hz, 1H), 7.50–7.36 (m, 8H), 7.33 (s, 1H), 7.18 (d,  $J$  = 7.0 Hz, 1H), 7.09–7.04 (m, 1H), 7.00 (d,  $J$  = 2.6 Hz, 1H), 5.17 (s, 2H). <sup>13</sup>C NMR (125 MHz, CDCl<sub>3</sub>)  $\delta$  145.1, 141.1, 140.9, 137.4, 136.1, 134.7, 129.9, 129.6, 129.6, 129.5, 129.0, 128.8, 128.5, 128.4, 128.0, 127.6, 120.9, 116.9, 115.5, 109.9, 70.4. HRMS  $m/z$  [M + Cl<sup>-</sup>] for C<sub>21</sub>H<sub>16</sub>Cl<sub>2</sub>NO<sub>3</sub>: calcd, 400.0513; found, 400.0505.

**(E)-5-(Benzyloxy)-2-(2-nitrovinyl)-3'-(trifluoromethyl)-1,1'-biphenyl (9f).** <sup>1</sup>H NMR (400 MHz, CDCl<sub>3</sub>)  $\delta$  7.90 (d,  $J$  = 13.5 Hz, 1H), 7.78–7.70 (m, 1H), 7.69–7.55 (m, 3H), 7.51–7.34 (m, 7H), 7.13–7.05 (dd,  $J$  = 8.8, 2.6 Hz, 1H), 7.02 (d,  $J$  = 2.6 Hz, 1H), 5.17 (s, 2H). <sup>13</sup>C NMR (100 MHz, CDCl<sub>3</sub>)  $\delta$  161.6, 155.7, 152.1, 145.1, 140.6, 140.0, 137.2, 136.4, 136.0, 133.2, 129.7, 129.3, 129.0, 128.6, 127.7, 121.0, 117.1, 115.8, 70.6. HRMS  $m/z$  [M + H<sup>+</sup>] for C<sub>22</sub>H<sub>17</sub>F<sub>3</sub>NO<sub>3</sub>: calcd, 400.1161; found, 400.1157.

**(E)-5-(Benzyloxy)-2-(2-nitrovinyl)-4'-(trifluoromethyl)-1,1'-biphenyl (9g).** Pushed through plug of SiO<sub>2</sub>. TS1–189: <sup>1</sup>H NMR (400 MHz, CDCl<sub>3</sub>)  $\delta$  7.98–7.90 (m, 1H), 7.80 (d,  $J$  = 8.0 Hz, 2H), 7.68 (d,  $J$  = 8.8 Hz, 1H), 7.52–7.37 (m, 8H), 7.11 (d,  $J$  = 8.8 Hz, 1H), 7.04 (s, 1H), 5.19 (s, 2H). <sup>13</sup>C NMR (100 MHz, CDCl<sub>3</sub>)  $\delta$  161.4, 147.8, 144.9, 144.3, 139.8, 138.6, 137.1, 136.4, 135.8, 133.5, 131.2, 129.5, 129.1, 128.8, 128.5, 127.6, 124.2, 120.8, 120.4, 117.0, 115.6, 70.4. HRMS  $m/z$  [M + H<sup>+</sup>] for C<sub>22</sub>H<sub>17</sub>F<sub>3</sub>NO<sub>3</sub>: calcd, 400.1155; found, 400.1151.

**(E)-5-(Benzyloxy)-2'-(2-nitrovinyl)-[1,1'-biphenyl]-2-yl(methyl)sulfane (9h).** <sup>1</sup>H NMR (400 MHz, CDCl<sub>3</sub>)  $\delta$  7.71 (d,  $J$  = 13.6 Hz, 1H), 7.62 (d,  $J$  = 8.6 Hz, 1H), 7.45–7.31 (m, 7H), 7.31–7.29 (m, 1H), 7.25–7.19 (t,  $J$  = 7.2 Hz, 1H), 7.13–6.99 (m, 2H), 6.95 (d,  $J$  = 2.8 Hz, 1H), 5.09 (s, 2H), 2.35 (s, 3H). <sup>13</sup>C NMR (100 MHz, CDCl<sub>3</sub>)  $\delta$  161.5, 144.9, 138.0, 137.5, 137.2, 136.1, 135.7, 130.0, 129.4, 129.3, 128.8, 128.4, 127.7, 125.0, 124.9, 121.6, 117.0, 115.8, 70.3, 15.6. HRMS  $m/z$  [M + K<sup>+</sup>] for C<sub>22</sub>H<sub>19</sub>NO<sub>3</sub>S: calcd, 416.0718; found, 416.0756.

**(E)-5-(Benzyloxy)-2'-methoxy-2-(2-nitrovinyl)-1,1'-biphenyl (9i).** <sup>1</sup>H NMR (500 MHz, CDCl<sub>3</sub>)  $\delta$  7.86 (d,  $J$  = 13.8 Hz, 1H), 7.65 (d,  $J$  = 8.7 Hz, 1H), 7.57–7.34 (m, 7H), 7.24–7.17 (m, 1H), 7.16–6.99 (m, 4H), 5.15 (s, 2H), 3.74 (s, 3H). <sup>13</sup>C NMR (125 MHz, CDCl<sub>3</sub>)  $\delta$  161.6, 156.4, 143.7, 138.8, 136.3, 135.3, 131.4, 130.4, 128.9, 128.4, 127.7, 122.0, 121.1, 117.5, 115.1, 111.4, 70.4, 55.6. HRMS  $m/z$  [M + H<sup>+</sup>] for C<sub>22</sub>H<sub>19</sub>NO<sub>4</sub>: calcd, 362.1387; found, 362.1389.

**(E)-5-(Benzyloxy)-3'-methoxy-2-(2-nitrovinyl)-1,1'-biphenyl (9j).** <sup>1</sup>H NMR (500 MHz, CDCl<sub>3</sub>)  $\delta$  8.04 (d,  $J$  = 13.6 Hz, 1H), 7.62 (d,  $J$  = 9.5 Hz, 1H), 7.46–7.37 (m, 6H), 7.07–7.02 (m, 3H), 7.02–6.97 (ddd,  $J$  = 8.2, 2.6, 0.9 Hz, 1H), 6.88–6.84 (m, 1H), 6.84–6.80 (dd,  $J$  = 2.6, 1.6 Hz, 1H), 5.15 (s, 2H), 3.85 (s, 3H). <sup>13</sup>C NMR (125 MHz, CDCl<sub>3</sub>)  $\delta$  161.5, 159.8, 146.8, 140.6, 138.2, 136.2, 135.9, 129.9, 129.5, 129.0, 128.6, 127.7, 122.3, 121.1, 116.8, 115.4, 115.4, 114.1, 70.5, 55.6. HRMS  $m/z$  [M + Na<sup>+</sup>] for C<sub>22</sub>H<sub>19</sub>NO<sub>4</sub>Na: 384.1212; found, 384.1218.

**(E)-5-(Benzyloxy)-3'-methyl-2-(2-nitrovinyl)-1,1'-biphenyl (9k).** <sup>1</sup>H NMR (500 MHz, CDCl<sub>3</sub>)  $\delta$  8.01 (d,  $J$  = 13.6 Hz, 1H), 7.62 (m, 1H), 7.48–7.39 (m, 7H), 7.39–7.33 (t,  $J$  = 7.7 Hz, 1H), 7.14–7.07 (m, 2H), 7.05–6.99 (m, 2H), 5.15 (s, 2H), 2.43 (s, 3H). <sup>13</sup>C NMR (125 MHz, CDCl<sub>3</sub>)  $\delta$  138.4, 135.8, 130.4, 129.5, 129.3, 128.9, 128.7, 128.5, 127.8, 127.8, 126.9, 121.1, 116.8, 115.3, 77.5, 77.4, 77.2, 77.0, 70.5, 21.7. HRMS  $m/z$  [M + Na<sup>+</sup>] for C<sub>22</sub>H<sub>19</sub>NO<sub>3</sub>N: calcd, 368.1263; found, 368.1257.

**(E)-4-((5'-(Benzyloxy)-2'-(2-nitrovinyl)-[1,1'-biphenyl]-3-yl)methyl)morpholine (9l).** <sup>1</sup>H NMR (400 MHz, CDCl<sub>3</sub>)  $\delta$  7.98 (d,  $J$  = 13.6 Hz, 1H), 7.63 (d,  $J$  = 9.5 Hz, 1H), 7.48–7.33 (m, 8H), 7.33 (d,  $J$  = 1.7 Hz, 1H), 7.23–7.20 (dd,  $J$  = 6.7, 1.8 Hz, 1H), 7.08–6.99 (m, 2H), 5.15 (d,  $J$  = 1.6 Hz, 2H), 3.79–3.67 (t,  $J$  = 4.1 Hz, 4H), 3.56 (s, 2H), 2.55–2.40 (dd,  $J$  = 5.7, 3.4 Hz, 4H). <sup>13</sup>C NMR (100 MHz, CDCl<sub>3</sub>)  $\delta$  161.5, 146.9, 139.2, 138.5, 138.1, 136.1, 135.8, 130.6, 129.5, 129.3, 128.9, 128.8, 128.5, 128.4, 127.7, 121.0, 116.9, 115.1, 70.4, 67.1,

63.3, 53.8. HRMS  $m/z$  [M + H<sup>+</sup>] for C<sub>26</sub>H<sub>27</sub>N<sub>2</sub>O<sub>4</sub>: calcd, 431.1971; found, 431.1974.

**(E)-((5'-(Benzyloxy)-2'-(2-nitrovinyl)-[1,1'-biphenyl]-4-yl)oxy)(tert-butyl)dimethylsilane (9m).** <sup>1</sup>H NMR (400 MHz, CDCl<sub>3</sub>)  $\delta$  8.03 (d,  $J$  = 13.7 Hz, 1H), 7.61 (d,  $J$  = 8.3 Hz, 1H), 7.49–7.33 (m, 6H), 7.17 (d,  $J$  = 8.4 Hz, 2H), 7.02 (s, 2H), 6.95 (d,  $J$  = 8.5 Hz, 2H), 5.15 (s, 2H), 1.04 (s, 9H), 0.30 (s, 6H). <sup>13</sup>C NMR (100 MHz, CDCl<sub>3</sub>)  $\delta$  161.5, 156.2, 146.8, 138.5, 136.2, 135.8, 132.2, 131.0, 129.6, 128.9, 128.5, 127.7, 121.1, 120.4, 116.8, 115.0, 70.4, 25.9, 18.4, -4.1. HRMS (FAB)  $m/z$  [M + Na<sup>+</sup>] for C<sub>27</sub>H<sub>31</sub>NO<sub>4</sub>SiNa: calcd, 484.1914; found, 484.1936.

**(E)-5-(5-(Benzyloxy)-2-(2-nitrovinyl)phenyl)benzo[d][1,3]dioxole (9n).** <sup>1</sup>H NMR (400 MHz, CDCl<sub>3</sub>)  $\delta$  8.03 (d,  $J$  = 13.6 Hz, 1H), 7.59 (d,  $J$  = 8.0 Hz, 1H), 7.50–7.33 (m, 6H), 7.05–6.98 (m, 2H), 6.92–6.85 (m, 1H), 6.79 (s, 1H), 6.71 (d,  $J$  = 7.9 Hz, 1H), 6.03 (s, 2H), 5.17 (s, 2H). <sup>13</sup>C NMR (100 MHz, CDCl<sub>3</sub>)  $\delta$  161.4, 148.0, 147.9, 146.5, 138.1, 136.1, 135.7, 132.9, 129.5, 128.8, 128.4, 127.6, 123.6, 121.0, 116.7, 115.0, 109.9, 108.5, 101.5, 70.3. HRMS (FAB)  $m/z$  [M + H<sup>+</sup>] for C<sub>22</sub>H<sub>18</sub>NO<sub>5</sub>: calcd, 376.1185; found, 376.1160.

**(E)-3-(5-(Benzyloxy)-2-(2-nitrovinyl)phenyl)pyridine (9o).** <sup>1</sup>H NMR (400 MHz, CDCl<sub>3</sub>)  $\delta$  8.70 (dd,  $J$  = 4.8, 1.6 Hz, 1H), 8.59 (d,  $J$  = 1.6 Hz, 1H), 7.89 (d,  $J$  = 13.5 Hz, 1H), 7.68–7.60 (m, 2H), 7.47–7.32 (m, 8H), 7.12–7.06 (dd,  $J$  = 8.7, 2.5 Hz, 1H), 7.00 (d,  $J$  = 2.6 Hz, 1H), 5.15 (s, 2H). <sup>13</sup>C NMR (100 MHz, CDCl<sub>3</sub>)  $\delta$  161.5, 149.9, 149.6, 142.8, 136.9, 136.8, 136.3, 135.8, 134.8, 129.7, 128.8, 128.5, 127.6, 123.4, 121.1, 117.1, 115.8, 70.4. HRMS (FAB)  $m/z$  [M + Na<sup>+</sup>] for C<sub>20</sub>H<sub>17</sub>N<sub>2</sub>O<sub>3</sub>: 333.1239; found, 333.1234.

**(E)-4-(5-(Benzyloxy)-2-(2-nitrovinyl)phenyl)pyridine (9p).** <sup>1</sup>H NMR (500 MHz, CDCl<sub>3</sub>)  $\delta$  8.74 (dd, 2H,  $J$  = 1.6, 4.4 Hz), 7.91 (d, 1H,  $J$  = 13.6 Hz), 7.67 (d, 1H,  $J$  = 8.8 Hz), 7.48 (d, 1H,  $J$  = 13.4 Hz), 7.41 (m, 5H), 7.25 (dd, 2H,  $J$  = 1.6, 4.4 Hz), 7.11 (dd, 1H,  $J$  = 2.6, 8.7 Hz), 7.01 (d, 1H,  $J$  = 2.5 Hz), 5.17 (s, 2H). <sup>13</sup>C NMR (125 MHz, CDCl<sub>3</sub>)  $\delta$  161.2, 150.2, 147.0, 143.7, 136.7, 136.6, 135.8, 128.9, 127.6, 124.5, 120.7, 116.8, 116.1, 70.6. ESI-HRMS  $m/z$  [M + H]<sup>+</sup> calcd for C<sub>20</sub>H<sub>17</sub>N<sub>2</sub>O<sub>3</sub>: 333.1239, found 333.1249.

**General Procedure for Preparation of 10a–p from 9a–p. N-(2-(5-(Benzyloxy)-[1,1'-biphenyl]-2-yl)ethyl)acetamide (10a).** Nitro-styrene **9a** (182 mg, 0.55 mmol) in THF (0.7 mL) was added dropwise to a solution of lithiumaluminum hydride (42 mg, 1.12 mmol) in THF (2 mL) under argon atmosphere at RT. Upon completion (nearly immediately), the reaction was quenched by the addition of water (42  $\mu$ L), 3 M NaOH (42  $\mu$ L), and water (84  $\mu$ L). The resulted mixture was filtered through a plug of Celite, washed with CH<sub>2</sub>Cl<sub>2</sub>, and dried over K<sub>2</sub>CO<sub>3</sub>. Upon filtration, the mixture was concentrated to oil and used without further purification. Acetic anhydride (58  $\mu$ L, 0.62 mmol) and triethylamine (93  $\mu$ L, 0.67 mmol) were added to a solution of the crude amine in CH<sub>2</sub>Cl<sub>2</sub> (5.6 mL) under an organ atmosphere at RT. After 3 h, the reaction was quenched with saturated aqueous ammonium chloride and extracted with CH<sub>2</sub>Cl<sub>2</sub> (3  $\times$  10 mL); combined organic fractions were washed with saturated aqueous sodium chloride, dried over Na<sub>2</sub>SO<sub>4</sub>, filtered, and concentrated. The residue was purified by column chromatography (SiO<sub>2</sub>; 3:1, Hex:EtOAc) to afford acetamide **10a** (0.12 g, 0.35 mmol, 64%). <sup>1</sup>H NMR (400 MHz, CDCl<sub>3</sub>)  $\delta$  7.50–7.38 (m, 8H), 7.38–7.30 (m, 2H), 7.23 (d,  $J$  = 8.4 Hz, 1H), 7.01–6.95 (dd,  $J$  = 8.4, 2.7 Hz, 1H), 6.93 (d,  $J$  = 2.7 Hz, 1H), 5.71 (br s, NH), 5.08 (s, 2H), 3.42–3.16 (q,  $J$  = 7.0 Hz, 2H), 2.89–2.64 (t,  $J$  = 7.2 Hz, 2H), 1.85 (s, 3H). <sup>13</sup>C NMR (100 MHz, CDCl<sub>3</sub>)  $\delta$  170.2, 157.2, 143.4, 141.4, 137.0, 130.8, 129.1, 128.7, 128.6, 128.4, 128.0, 127.6, 127.2, 116.6, 114.2, 70.1, 40.7, 31.9, 23.2. HRMS  $m/z$  [M + K<sup>+</sup>] for C<sub>23</sub>H<sub>23</sub>NO<sub>2</sub>K: calcd, 384.1361; found, 384.1359.

**N-(2-(5-(Benzyloxy)-3'-fluoro-[1,1'-biphenyl]-2-yl)ethyl)acetamide (10b).** <sup>1</sup>H NMR (400 MHz, CDCl<sub>3</sub>)  $\delta$  7.48–7.30 (m, 6H), 7.24–7.18 (d,  $J$  = 8.4 Hz, 1H), 7.12–7.04 (m, 2H), 7.04–6.92 (ddd,  $J$  = 18.6, 8.2, 2.5 Hz, 2H), 6.85 (d,  $J$  = 2.7 Hz, 1H), 5.34 (br s, NH), 5.05 (s, 2H), 3.32–3.21 (q,  $J$  = 6.4, 5.9 Hz, 2H), 2.79–2.68 (t,  $J$  = 7.1 Hz, 2H), 1.86 (s, 3H). <sup>13</sup>C NMR (100 MHz, CDCl<sub>3</sub>)  $\delta$  170.3, 157.3, 143.7, 142.2, 136.9, 131.0, 130.1, 123.0, 128.8, 128.6, 128.2, 127.7, 125.0, 116.5, 116.4, 114.6, 114.4, 70.2, 40.8, 32.0, 23.3. HRMS  $m/z$  [M + H<sup>+</sup>] for C<sub>23</sub>H<sub>23</sub>FNO<sub>2</sub>: calcd, 364.1713; found, 364.1705.

*N*-(2-(5-(Benzyloxy)-4'-fluoro-[1,1'-biphenyl]-2-yl)ethyl)acetamide (**10c**).  $^1\text{H}$  NMR (400 MHz,  $\text{CDCl}_3$ )  $\delta$  7.44–7.31 (m, 6H), 7.27–7.22 (dd,  $J = 8.4$ , 5.5 Hz, 1H), 7.21–7.17 (d,  $J = 8.4$  Hz, 1H), 7.12–7.05 (m, 3H), 6.96–6.91 (dd,  $J = 8.3$ , 3.0 Hz, 1H), 5.83 (br s, NH), 5.05 (s, 2H), 3.33–3.15 (q,  $J = 6.7$  Hz, 2H), 2.78–2.66 (t,  $J = 7.2$  Hz, 2H), 1.87 (s, 3H).  $^{13}\text{C}$  NMR (100 MHz,  $\text{CDCl}_3$ )  $\delta$  170.5, 157.3, 142.4, 137.0, 130.9, 130.8, 130.7, 128.7, 128.7, 128.2, 127.7, 116.8, 115.5, 115.3, 114.3, 70.2, 40.8, 32.0, 23.1. HRMS  $m/z$  [ $\text{M} + \text{Na}^+$ ] for  $\text{C}_{23}\text{H}_{22}\text{FNO}_2\text{Na}$ : calcd, 386.1527; found, 386.1529.

*N*-(2-(5-(Benzyloxy)-2'-chloro-[1,1'-biphenyl]-2-yl)ethyl)acetamide (**10d**).  $^1\text{H}$  NMR (500 MHz,  $\text{CDCl}_3$ )  $\delta$  7.52–7.45 (m, 1H), 7.45–7.40 (m, 2H), 7.40–7.35 (m, 3H), 7.35–7.29 (m, 3H), 7.25–7.21 (m, 1H), 7.05–6.95 (dd,  $J = 8.5$ , 2.8 Hz, 1H), 6.82 (d,  $J = 2.7$  Hz, 1H), 5.93 (d,  $J = 5.4$  Hz, 1H), 5.05 (s, 2H), 3.36–3.19 (ddq,  $J = 19.3$ , 13.0, 6.1 Hz, 2H), 2.67–2.49 (m, 2H), 1.93 (s, 3H).  $^{13}\text{C}$  NMR (125 MHz,  $\text{CDCl}_3$ )  $\delta$  175.7, 171.0, 157.1, 140.4, 139.8, 136.9, 133.1, 131.3, 130.4, 129.6, 129.0, 128.6, 128.0, 127.6, 126.8, 116.4, 114.9, 70.1, 40.3, 31.8, 22.9. HRMS  $m/z$  [ $\text{M} + \text{H}^+$ ] for  $\text{C}_{23}\text{H}_{23}\text{ClNO}_2$ : calcd, 380.1417; found, 380.1415.

*N*-(2-(5-(Benzyloxy)-3'-chloro-[1,1'-biphenyl]-2-yl)ethyl)acetamide (**10e**).  $^1\text{H}$  NMR (500 MHz,  $\text{CDCl}_3$ )  $\delta$  7.47–7.28 (m, 8H), 7.25–7.17 (m, 2H), 6.99–6.92 (dd,  $J = 8.5$ , 2.7 Hz, 1H), 6.84 (d,  $J = 2.8$  Hz, 1H), 5.46 (br s, NH), 5.06 (s, 2H), 3.34–3.25 (m, 2H), 2.83–2.68 (t,  $J = 7.3$  Hz, 2H), 2.03 (s, 3H).  $^{13}\text{C}$  NMR (125 MHz,  $\text{CDCl}_3$ )  $\delta$  171.6, 157.5, 143.2, 142.1, 136.9, 134.3, 131.1, 129.9, 129.3, 128.8, 128.3, 127.7, 127.6, 127.5, 116.7, 114.8, 70.3, 46.1, 41.3, 31.7, 22.5, 8.8. HRMS  $m/z$  [ $\text{M} + \text{H}^+$ ] for  $\text{C}_{23}\text{H}_{23}\text{ClNO}_2$ : calcd, 380.1412; found, 380.1414.

*N*-(2-(5-(Benzyloxy)-3'-(trifluoromethyl)-[1,1'-biphenyl]-2-yl)ethyl)acetamide (**10f**).  $^1\text{H}$  NMR (400 MHz,  $\text{CDCl}_3$ )  $\delta$  7.64 (d,  $J = 7.7$  Hz, 1H), 7.59–7.54 (m, 2H), 7.55–7.49 (t,  $J = 7.3$  Hz, 1H), 7.47–7.32 (m, 5H), 7.24 (d,  $J = 8.5$  Hz, 1H), 7.01–6.96 (dd,  $J = 8.5$ , 2.7 Hz, 1H), 6.87 (d,  $J = 2.7$  Hz, 1H), 5.90 (br s, NH), 5.06 (s, 2H), 3.34–3.23 (q,  $J = 6.9$  Hz, 2H), 2.79–2.68 (t,  $J = 7.3$  Hz, 2H), 1.99 (s, 3H).  $^{13}\text{C}$  NMR (100 MHz,  $\text{CDCl}_3$ )  $\delta$  170.7, 157.4, 142.2, 141.9, 136.9, 132.6, 131.1, 129.0, 128.8, 128.5, 128.2, 127.7, 124.2, 116.7, 114.8, 70.3, 40.8, 31.9, 23.0. HRMS  $m/z$  [ $\text{M} + \text{H}^+$ ] for  $\text{C}_{24}\text{H}_{23}\text{F}_3\text{NO}_2$ : calcd, 414.1676; found, 414.1681.

*N*-(2-(5-(Benzyloxy)-4'-(trifluoromethyl)-[1,1'-biphenyl]-2-yl)ethyl)acetamide (**10g**).  $^1\text{H}$  NMR (400 MHz,  $\text{CDCl}_3$ )  $\delta$  7.66 (d,  $J = 8.1$  Hz, 2H), 7.46–7.23 (m, 8H), 6.99–6.94 (dd,  $J = 8.5$ , 2.7 Hz, 1H), 6.84 (d,  $J = 2.7$  Hz, 1H), 6.03 (t,  $J = 5.5$  Hz, 1H), 5.06 (s, 2H), 3.33–3.19 (dd,  $J = 14.3$ , 6.4 Hz, 2H), 2.76–2.68 (dd,  $J = 8.3$ , 6.6 Hz, 2H), 1.85 (s, 3H).  $^{13}\text{C}$  NMR (100 MHz,  $\text{CDCl}_3$ )  $\delta$  170.3, 157.1, 145.1, 141.8, 136.8, 130.9, 129.5, 129.1, 128.6, 128.6, 127.5, 125.6, 125.2, 125.2, 122.9, 116.4, 114.6, 70.1, 40.6, 31.9. HRMS  $m/z$ : [ $\text{M} + \text{Na}^+$ ] for  $\text{C}_{24}\text{H}_{22}\text{F}_3\text{NO}_2\text{Na}$ : calcd, 436.1495; found, 436.1489.

*N*-(2-(5-(Benzyloxy)-2'-(methylthio)-[1,1'-biphenyl]-2-yl)ethyl)acetamide (**10h**).  $^1\text{H}$  NMR (400 MHz,  $\text{CDCl}_3$ )  $\delta$  7.48–7.30 (m, 7H), 7.28–7.18 (m, 2H), 7.14 (s, 1H), 7.03–6.98 (ddd,  $J = 8.5$ , 2.8, 1.0 Hz, 1H), 6.87–6.83 (m, 1H), 5.63 (br s, NH), 5.05 (s, 2H), 3.43–3.16 (ddt,  $J = 42.5$ , 13.3, 6.6 Hz, 2H), 2.66–2.52 (t,  $J = 6.7$  Hz, 2H), 2.39 (d,  $J = 1.0$  Hz, 3H), 1.84 (d,  $J = 1.0$  Hz, 3H).  $^{13}\text{C}$  NMR (100 MHz,  $\text{CDCl}_3$ )  $\delta$  170.3, 157.3, 141.1, 139.1, 137.6, 137.0, 130.6, 129.8, 129.4, 128.7, 128.4, 128.1, 127.7, 124.5, 124.0, 116.5, 115.2, 70.2, 40.1, 31.7, 23.3, 15.2. HRMS  $m/z$  [ $\text{M} + \text{Na}^+$ ] for  $\text{C}_{24}\text{H}_{25}\text{NO}_2\text{SNa}$ : calcd, 414.1504; found, 414.1509.

*N*-(2-(5-(Benzyloxy)-2'-methoxy-[1,1'-biphenyl]-2-yl)ethyl)acetamide (**10i**).  $^1\text{H}$  NMR (400 MHz,  $\text{CDCl}_3$ )  $\delta$  7.47–7.30 (m, 5H), 7.22 (d,  $J = 8.5$  Hz, 1H), 7.17–7.13 (dd,  $J = 7.4$ , 1.9 Hz, 1H), 7.07–6.95 (m, 4H), 6.85 (d,  $J = 2.7$  Hz, 1H), 5.51 (br s, NH), 5.07 (s, 2H), 3.77 (s, 3H), 3.44–3.18 (m, 2H), 2.68–2.56 (td,  $J = 6.8$ , 3.7 Hz, 2H), 1.86 (s, 3H).  $^{13}\text{C}$  NMR (125 MHz,  $\text{CDCl}_3$ )  $\delta$  170.0, 157.2, 156.4, 139.9, 137.1, 131.2, 130.1, 129.2, 128.7, 128.1, 127.8, 120.9, 116.8, 114.4, 111.2, 70.1, 55.8, 40.4, 31.9, 23.5. HRMS  $m/z$  [ $\text{M} + \text{H}^+$ ] for  $\text{C}_{24}\text{H}_{26}\text{NO}_3$ : calcd, 376.1913; found, 376.1902.

*N*-(2-(5-(Benzyloxy)-3'-methoxy-[1,1'-biphenyl]-2-yl)ethyl)acetamide (**10j**).  $^1\text{H}$  NMR (400 MHz,  $\text{CDCl}_3$ )  $\delta$  7.48–7.36 (m, 4H), 7.36–7.30 (m, 3H), 7.21 (d,  $J = 8.4$  Hz, 1H), 6.98–6.92 (m, 1H), 6.92–6.82 (m, 3H), 5.49 (br s, NH), 5.06 (s, 2H), 3.85 (s, 3H), 3.34–3.22 (q,  $J = 6.6$ , 6.2 Hz, 2H), 2.85–2.68 (t,  $J = 7.2$  Hz, 2H), 1.85 (s,

3H).  $^{13}\text{C}$  NMR (100 MHz,  $\text{CDCl}_3$ )  $\delta$  170.1, 159.5, 157.2, 143.3, 142.9, 137.0, 130.8, 129.5, 128.7, 128.1, 128.1, 127.7, 121.6, 116.5, 114.9, 114.3, 112.7, 70.17, 55.4, 40.8, 32.0, 23.3. HRMS  $m/z$  [ $\text{M} + \text{H}^+$ ] for  $\text{C}_{24}\text{H}_{25}\text{NO}_3\text{Na}$ : calcd, 398.1732; found, 398.1725.

*N*-(2-(5-(Benzyloxy)-3'-methyl-[1,1'-biphenyl]-2-yl)ethyl)acetamide (**10k**).  $^1\text{H}$  NMR (400 MHz,  $\text{CDCl}_3$ )  $\delta$  7.45 (m, 3H), 7.40 (m, 3H), 7.37–7.30 (q,  $J = 7.7$ , 7.1 Hz, 1H), 7.21 (d,  $J = 1.4$  Hz, 1H), 7.15–7.10 (m, 2H), 6.96 (d,  $J = 8.1$  Hz, 1H), 6.90 (s, 1H), 5.51 (br s, NH), 5.08 (s, 2H), 3.34–3.24 (q,  $J = 6.5$  Hz, 2H), 2.83–2.71 (t,  $J = 7.0$  Hz, 2H), 2.41 (s, 3H), 1.84 (s, 3H).  $^{13}\text{C}$  NMR (100 MHz,  $\text{CDCl}_3$ )  $\delta$  170.0, 157.2, 143.5, 141.4, 138.0, 137.0, 130.7, 129.9, 128.7, 128.7, 128.3, 128.1, 128.0, 127.6, 126.2, 116.5, 114.2, 70.1, 40.8, 31.9, 23.3, 21.6. ESI-HRMS  $m/z$  [ $\text{M} + \text{Na}^+$ ] calcd for  $\text{C}_{24}\text{H}_{25}\text{NO}_2\text{Na}$ : 382.1777, found 382.1770.

*N*-(2-(5-(Benzyloxy)-3'-(morpholinomethyl)-[1,1'-biphenyl]-2-yl)ethyl)acetamide (**10l**).  $^1\text{H}$  NMR (400 MHz,  $\text{CDCl}_3$ )  $\delta$  7.47–7.30 (m, 7H), 7.28 (s, 1H), 7.24–7.18 (m, 2H), 6.98–6.93 (dd,  $J = 8.4$ , 2.8 Hz, 1H), 6.89 (d,  $J = 2.7$  Hz, 1H), 5.40 (s, 1H), 5.05 (s, 2H), 3.75–3.69 (t,  $J = 4.7$  Hz, 4H), 3.55 (s, 2H), 3.36–3.22 (q,  $J = 6.9$  Hz, 2H), 2.80–2.68 (t,  $J = 7.1$  Hz, 2H), 2.47 (m, 4H), 1.85 (s, 3H).  $^{13}\text{C}$  NMR (100 MHz,  $\text{CDCl}_3$ )  $\delta$  170.0, 157.3, 143.4, 141.5, 138.0, 137.1, 130.9, 123.0, 128.7, 128.7, 128.4, 128.2, 128.2, 128.0, 127.7, 116.8, 114.1, 70.2, 67.1, 63.5, 53.8, 40.6, 32.1, 23.4. HRMS  $m/z$  [ $\text{M} + \text{H}^+$ ] for  $\text{C}_{28}\text{H}_{33}\text{N}_2\text{O}_3$ : calcd, 445.2491; found, 445.2494.

*N*-(2-(5-(Benzyloxy)-4'-(tert-butyl(dimethylsilyloxy)-[1,1'-biphenyl]-2-yl)ethyl)acetamide (**10m**).  $^1\text{H}$  NMR (500 MHz,  $\text{CDCl}_3$ )  $\delta$  7.44 (d,  $J = 7.5$  Hz, 3H), 7.42–7.36 (dt,  $J = 10.5$ , 5.7 Hz, 3H), 7.36–7.31 (m, 1H), 7.21–7.14 (m, 3H), 6.94–6.86 (m, 2H), 5.08 (s, 2H), 3.34–3.23 (q,  $J = 6.7$  Hz, 2H), 2.75 (t,  $J = 7.1$  Hz, 2H), 1.74 (s, 3H), 1.97 (s, 9H), 0.25 (s, 6H).  $^{13}\text{C}$  NMR (125 MHz,  $\text{CDCl}_3$ )  $\delta$  169.9, 157.3, 155.0, 143.3, 137.2, 134.5, 130.8, 130.2, 128.7, 128.1, 127.7, 127.0, 116.8, 114.0, 70.2, 53.6, 40.7, 32.1, 25.8, 23.4, 18.4, -4.2. HRMS (FAB)  $m/z$  [ $\text{M} + \text{Na}^+$ ] for  $\text{C}_{29}\text{H}_{37}\text{NO}_3\text{SiNa}$ : calcd, 498.2440; found, 498.2447.

*N*-(2-(Benzo[d][1,3]dioxol-5-yl)-4-(benzyloxy)phenethyl)acetamide (**10n**).  $^1\text{H}$  NMR (400 MHz,  $\text{CDCl}_3$ )  $\delta$  7.49–7.36 (m, 5H), 7.34 (d,  $J = 4.4$  Hz, 1H), 7.20 (d,  $J = 8.3$  Hz, 1H), 6.96–6.89 (dd,  $J = 8.4$ , 2.8 Hz, 1H), 6.90–6.84 (m, 2H), 6.81–6.73 (m, 1H), 6.00 (s, 2H), 5.69–5.60 (t,  $J = 5.8$  Hz, 1H), 5.06 (s, 2H), 3.42–3.16 (m, 2H), 2.93–2.68 (t,  $J = 7.3$  Hz, 2H), 1.87 (s, 3H).  $^{13}\text{C}$  NMR (100 MHz,  $\text{CDCl}_3$ )  $\delta$  170.4, 157.2, 147.5, 146.8, 143.0, 137.0, 135.2, 130.8, 129.3, 128.8, 128.1, 127.6, 123.2, 122.4, 116.7, 114.1, 109.7, 108.3, 101.2, 70.1, 40.7, 31.9, 23.2. HRMS (FAB)  $m/z$  [ $\text{M} + \text{Na}^+$ ] for  $\text{C}_{24}\text{H}_{23}\text{NO}_4\text{Na}$ : calcd, 412.1519; found, 412.1524.

*N*-(4-(Benzyloxy)-2-(pyridin-3-yl)phenethyl)acetamide (**10o**).  $^1\text{H}$  NMR (400 MHz,  $\text{CDCl}_3$ )  $\delta$  8.69–8.52 (dd,  $J = 18.2$ , 4.0 Hz, 2H), 7.71–7.63 (dt,  $J = 7.8$ , 2.0 Hz, 1H), 7.49–7.31 (m, 7H), 7.06–6.97 (dd,  $J = 8.5$ , 2.8 Hz, 1H), 6.84 (d,  $J = 2.8$  Hz, 1H), 5.06 (s, 2H), 3.36–3.20 (q,  $J = 6.5$  Hz, 2H), 2.78–2.67 (dd,  $J = 8.1$ , 6.6 Hz, 2H), 1.90 (s, 3H).  $^{13}\text{C}$  NMR (125 MHz,  $\text{CDCl}_3$ )  $\delta$  170.1, 157.5, 149.6, 148.5, 139.5, 136.9, 131.2, 129.0, 128.8, 128.3, 127.7, 123.5, 116.9, 115.0, 70.3, 40.7, 32.2, 23.5. HRMS (FAB)  $m/z$ : [ $\text{M} + \text{H}^+$ ] for  $\text{C}_{22}\text{H}_{23}\text{N}_2\text{O}_2$ : calcd, 347.1759; found, 347.1754.

*N*-(4-(Benzyloxy)-2-(pyridin-4-yl)phenethyl)acetamide (**10p**).  $^1\text{H}$  NMR (400 MHz,  $\text{CDCl}_3$ )  $\delta$  8.66 (d,  $J = 5.1$  Hz, 2H), 7.46–7.39 (m, 5H), 7.36 (s, 1H), 7.30 (s, 2H), 7.06–7.01 (m, 1H), 6.84 (d,  $J = 2.7$  Hz, 1H), 5.94 (d,  $J = 4.8$  Hz, 1H), 5.09 (s, 2H), 3.35–3.23 (dd,  $J = 14.5$ , 6.4 Hz, 2H), 2.74 (t,  $J = 7.5$  Hz, 2H), 1.90 (s, 3H).  $^{13}\text{C}$  NMR (100 MHz,  $\text{CDCl}_3$ )  $\delta$  171.4, 158.1, 156.3, 137.2, 132.3, 132.2, 130.8, 128.7, 128.5, 129.7, 127.5, 117.9, 106.2, 103.0, 69.9, 41.1, 29.7, 29.6, 23.1. HRMS (FAB)  $m/z$  [ $\text{M} + \text{Na}^+$ ] for  $\text{C}_{22}\text{H}_{22}\text{N}_2\text{O}_2\text{Na}$ : calcd, 369.1579; found, 369.1573.

**General Hydrogenolysis Procedure for Compounds 10a–p.** *N*-(2-(5-Hydroxy-[1,1'-biphenyl]-2-yl)ethyl)acetamide (**11a**). Palladium on carbon (10%, 5 mg) was added to **10a** (120 mg, 0.35 mmol) in degassed MeOH (3.5 mL), and the solution was placed under an atmosphere of  $\text{H}_2$ . After 12 h, the solution was diluted with  $\text{CH}_2\text{Cl}_2$  and filtered through Celite. The eluent was concentrated to afford a yellow solid, which was purified by column chromatography ( $\text{SiO}_2$ , 100:5,  $\text{CH}_2\text{Cl}_2$ :MeOH) to afford phenol **11a** (64 mg, 0.25 mmol,

79%) as a pale-yellow amorphous solid.  $^1\text{H}$  NMR (400 MHz,  $\text{CDCl}_3$ )  $\delta$  7.25–7.14 (m, 5H), 7.11–7.05 (m, 1H), 6.90 (d,  $J$  = 8.3 Hz, 1H), 6.64 (d,  $J$  = 8.3 Hz, 1H), 6.59 (d,  $J$  = 2.5 Hz, 1H), 5.61 (t,  $J$  = 5.5 Hz, 1H), 3.12–3.02 (m, 2H), 2.55 (t,  $J$  = 7.1 Hz, 2H), 1.66 (s, 3H).  $^{13}\text{C}$  NMR (100 MHz,  $\text{CDCl}_3$ )  $\delta$  171.2, 155.2, 143.4, 141.6, 130.8, 129.1, 128.4, 127.2, 117.4, 115.0, 41.1, 31.8, 23.2. HRMS  $m/z$  [ $\text{M} + \text{Na}^+$ ] for  $\text{C}_{16}\text{H}_{17}\text{NO}_2\text{Na}$ : calcd, 278.1151; found, 278.1155.

*N*-(2-(3'-Fluoro-5-hydroxy-[1,1'-biphenyl]-2-yl)ethyl)acetamide (**11b**).  $^1\text{H}$  NMR (500 MHz, MeOD)  $\delta$  7.88 (s, 1H), 7.39 (d,  $J$  = 7.5 Hz, 1H), 7.16–6.99 (m, 4H), 6.77 (d,  $J$  = 8.1 Hz, 1H), 6.62 (d,  $J$  = 2.6 Hz, 1H), 3.15 (t,  $J$  = 6.6 Hz, 2H), 2.66 (t,  $J$  = 7.4 Hz, 2H), 1.80 (s, 3H).  $^{13}\text{C}$  NMR (125 MHz, MeOD)  $\delta$  173.1, 164.8, 162.9, 156.7, 145.5, 143.3, 132.0, 131.0, 128.3, 126.1, 117.6, 115.9, 114.7, 41.8, 32.8, 22.5. HRMS  $m/z$  [ $\text{M} + \text{Na}^+$ ] for  $\text{C}_{16}\text{H}_{16}\text{FNO}_2\text{Na}$ : calcd, 296.1063; found, 296.1059.

*N*-(2-(4'-Fluoro-5-hydroxy-[1,1'-biphenyl]-2-yl)ethyl)acetamide (**11c**).  $^1\text{H}$  NMR (400 MHz, MeOD)  $\delta$  7.26–7.20 (m, 2H), 7.11–7.03 (m, 3H), 6.71 (dd,  $J$  = 8.3, 2.5 Hz, 1H), 6.56 (d,  $J$  = 2.5 Hz, 1H), 3.07 (t,  $J$  = 7.6 Hz, 2H), 2.60 (t,  $J$  = 7.6 Hz, 2H), 1.78 (s, 3H).  $^{13}\text{C}$  NMR (100 MHz, MeOD)  $\delta$  173.0, 156.7, 143.5, 139.2, 131.9, 131.9, 131.8, 128.5, 117.8, 116.0, 115.8, 115.7, 41.8, 32.9, 22.5. HRMS  $m/z$  [ $\text{M} + \text{Na}^+$ ] for  $\text{C}_{16}\text{H}_{16}\text{FNO}_2\text{Na}$ : calcd, 296.1063; found, 296.1065.

*N*-(2-(2'-Chloro-5-hydroxy-[1,1'-biphenyl]-2-yl)ethyl)acetamide (**11d**).  $^1\text{H}$  NMR (400 MHz,  $\text{CDCl}_3$ )  $\delta$  8.37 (br s, OH), 7.45–7.39 (m, 1H), 7.32–7.24 (m, 2H), 7.21–7.15 (m, 1H), 7.09 (d,  $J$  = 8.3 Hz, 1H), 6.85 (dd,  $J$  = 8.3, 2.5 Hz, 1H), 6.68 (d,  $J$  = 2.6 Hz, 1H), 5.62 (s, 1H), 3.40–3.14 (m, 2H), 2.63–2.44 (dd,  $J$  = 7.1, 5.1 Hz, 2H), 1.86 (s, 3H).  $^{13}\text{C}$  NMR (125 MHz,  $\text{CDCl}_3$ )  $\delta$  171.1, 155.1, 140.5, 140.0, 133.2, 131.4, 130.5, 129.7, 129.0, 127.7, 126.9, 117.3, 115.7, 40.5, 31.8, 23.3. HRMS  $m/z$  [ $\text{M} + \text{H}^+$ ] for  $\text{C}_{16}\text{H}_{17}\text{ClNO}_2$ : calcd, 290.0948; found, 290.0941.

*N*-(2-(3'-Chloro-5-hydroxy-[1,1'-biphenyl]-2-yl)ethyl)acetamide (**11e**).  $^1\text{H}$  NMR (500 MHz,  $\text{CDCl}_3$ )  $\delta$  7.40–7.09 (m, 5H), 6.83–6.76 (dq,  $J$  = 8.1, 4.9, 3.8 Hz, 1H), 6.76–6.67 (dd,  $J$  = 18.3, 2.7 Hz, 1H), 3.34–3.23 (p,  $J$  = 6.6 Hz, 2H), 2.77–2.64 (dt,  $J$  = 14.3, 7.2 Hz, 2H), 1.76 (s, 3H).  $^{13}\text{C}$  NMR (125 MHz,  $\text{CDCl}_3$ )  $\delta$  170.8, 154.9, 143.6, 141.6, 131.0, 130.9, 129.7, 129.2, 128.5, 127.5, 117.4, 115.5, 115.0, 41.0, 32.0, 23.4. HRMS  $m/z$  [ $\text{M} + \text{Na}^+$ ] for  $\text{C}_{16}\text{H}_{16}\text{ClNO}_2\text{Na}$ : calcd, 312.0762; found, 312.0788.

*N*-(2-(5-Hydroxy-3'-(trifluoromethyl)-[1,1'-biphenyl]-2-yl)ethyl)acetamide (**11f**).  $^1\text{H}$  NMR (400 MHz,  $\text{CDCl}_3$ )  $\delta$  7.64–7.39 (m, 4H), 7.07 (s, 1H), 6.82 (s, 1H), 6.73 (s, 1H), 6.00 (s, 1H), 3.34–3.18 (q,  $J$  = 6.8 Hz, 2H), 2.66 (t,  $J$  = 7.0 Hz, 2H), 1.87 (s, 3H).  $^{13}\text{C}$  NMR (100 MHz,  $\text{CDCl}_3$ )  $\delta$  171.3, 155.4, 142.4, 141.8, 132.6, 131.0, 130.8, 128.9, 126.9, 125.8, 125.8, 124.0, 117.3, 115.6, 60.7, 41.0, 21.2. HRMS  $m/z$  [ $\text{M} + \text{Na}^+$ ] for  $\text{C}_{17}\text{H}_{16}\text{F}_3\text{NO}_2\text{Na}$ : calcd, 346.1031; found, 346.1040.

*N*-(2-(5-Hydroxy-4'-(trifluoromethyl)-[1,1'-biphenyl]-2-yl)ethyl)acetamide (**11g**).  $^1\text{H}$  NMR (400 MHz,  $\text{CDCl}_3$ )  $\delta$  7.54 (d, 2H,  $J$  = 8.0 Hz), 7.31 (d, 2H,  $J$  = 8.0 Hz), 7.03 (d, 1H,  $J$  = 8.3 Hz), 6.72 (dd, 1H,  $J$  = 2.5, 8.3 Hz), 6.59 (d, 1H,  $J$  = 2.5 Hz), 4.09 (br s, 2H), 3.10 (t,  $J$  = 7.5 Hz, 2H), 2.56 (t, 2H,  $J$  = 7.5 Hz), 1.76 (s, 3H).  $^{13}\text{C}$  NMR (100 MHz,  $\text{CDCl}_3$ )  $\delta$  170.5, 155.1, 146.3, 141.7, 130.8, 129.5, 127.1, 125.1 (q,  $J$  = 4.2 Hz), 116.9, 116.5, 115.3, 45.6, 40.6, 23.0. HRMS  $m/z$  [ $\text{M} + \text{H}^+$ ] for  $\text{C}_{17}\text{H}_{16}\text{F}_3\text{NO}_2\text{Na}$ : calcd, 346.1031; found, 346.1025.

*N*-(2-(5-Hydroxy-2'-(methylthio)-[1,1'-biphenyl]-2-yl)ethyl)acetamide (**11h**).  $^1\text{H}$  NMR (500 MHz,  $\text{CDCl}_3$ )  $\delta$  7.40–7.34 (m, 1H), 7.25–7.14 (m, 3H), 7.12–7.07 (m, 1H), 6.86–6.82 (dd,  $J$  = 8.4, 2.7 Hz, 1H), 6.68 (d,  $J$  = 2.7 Hz, 1H), 5.51 (br s, NH), 3.42–3.16 (m, 2H), 2.55 (t,  $J$  = 6.8 Hz, 2H), 2.37 (s, 3H), 1.85 (s, 3H).  $^{13}\text{C}$  NMR (125 MHz,  $\text{CDCl}_3$ )  $\delta$  170.5, 154.5, 141.2, 139.1, 137.6, 130.7, 123.0, 128.8, 128.5, 124.6, 124.0, 117.3, 115.6, 40.2, 31.6, 23.4, 15.2. HRMS  $m/z$  [ $\text{M} + \text{Na}^+$ ] for  $\text{C}_{17}\text{H}_{19}\text{NO}_2\text{SNa}$ : calcd, 324.1034; found, 324.1035.

*N*-(2-(5-Hydroxy-2'-methoxy-[1,1'-biphenyl]-2-yl)ethyl)acetamide (**11i**).  $^1\text{H}$  NMR (400 MHz,  $\text{CDCl}_3$ )  $\delta$  7.52 (br s, OH), 7.41–7.31 (m, 1H), 7.14–7.07 (dd,  $J$  = 8.4, 6.4 Hz, 1H), 7.05–6.94 (m, 3H), 6.83–6.76 (dd,  $J$  = 8.3, 2.7 Hz, 1H), 6.70 (d,  $J$  = 2.7 Hz, 1H), 5.55 (s, 1H), 3.76 (s, 3H), 3.41–3.17 (ddt,  $J$  = 34.4, 13.1, 6.5 Hz, 2H), 2.57 (t,  $J$  = 6.9 Hz, 2H), 1.85 (s, 3H).  $^{13}\text{C}$  NMR (125 MHz,  $\text{CDCl}_3$ )  $\delta$  171.0, 156.4, 155.1, 139.9, 131.3, 130.5, 130.1, 129.1, 128.5, 121.0, 117.7, 115.2, 111.4, 55.9, 40.7, 31.7, 23.3. HRMS  $m/z$  [ $\text{M} + \text{Na}^+$ ] for  $\text{C}_{17}\text{H}_{19}\text{NO}_3\text{Na}$ : calcd, 308.1263; found, 308.1264.

*N*-(2-(5-Hydroxy-3'-methoxy-[1,1'-biphenyl]-2-yl)ethyl)acetamide (**11j**).  $^1\text{H}$  NMR (400 MHz,  $\text{CDCl}_3$ )  $\delta$  7.83 (br s, OH), 7.30–7.24 (m, 1H), 7.06 (d,  $J$  = 8.2 Hz, 1H), 6.90–6.70 (m, 5H), 5.59 (t,  $J$  = 5.7 Hz, 1H), 3.79 (s, 3H), 3.33–3.19 (q,  $J$  = 6.9 Hz, 2H), 2.69 (t,  $J$  = 7.1 Hz, 2H), 1.85 (s, 3H).  $^{13}\text{C}$  NMR (100 MHz,  $\text{CDCl}_3$ )  $\delta$  171.1, 159.4, 155.1, 143.3, 143.0, 130.9, 129.5, 127.3, 121.7, 117.2, 115.1, 115.0, 112.6, 55.4, 41.1, 31.8, 23.3. HRMS  $m/z$  [ $\text{M} + \text{H}^+$ ] for  $\text{C}_{17}\text{H}_{20}\text{NO}_3$ : calcd, 286.1443; found, 286.1436.

*N*-(2-(5-Hydroxy-3'-methyl-[1,1'-biphenyl]-2-yl)ethyl)acetamide (**11k**).  $^1\text{H}$  NMR (400 MHz,  $\text{CDCl}_3$ )  $\delta$  7.50 (br s, OH), 7.30–7.24 (m, 1H), 7.15 (d,  $J$  = 7.6 Hz, 1H), 7.09–7.03 (m, 3H), 6.80 (d,  $J$  = 7.6 Hz, 1H), 6.73 (s, 1H), 5.53 (br s, NH), 3.31–3.21 (q,  $J$  = 6.7 Hz, 2H), 2.71 (t,  $J$  = 7.0 Hz, 2H), 2.37 (s, 3H), 1.85 (s, 3H).  $^{13}\text{C}$  NMR (100 MHz,  $\text{CDCl}_3$ )  $\delta$  170.9, 155.0, 143.6, 141.6, 138.1, 130.8, 123.0, 128.3, 128.0, 127.4, 126.3, 117.4, 114.9, 41.1, 31.8, 23.3, 21.7. HRMS  $m/z$  [ $\text{M} + \text{Na}^+$ ] for  $\text{C}_{17}\text{H}_{19}\text{NO}_2\text{Na}$ : calcd, 292.1308; found, 292.1314.

*N*-(2-(5-Hydroxy-3'-(morpholinomethyl)-[1,1'-biphenyl]-2-yl)ethyl)acetamide (**11l**).  $^1\text{H}$  NMR (500 MHz,  $\text{CDCl}_3$ )  $\delta$  7.36–7.23 (m, 4H), 7.16 (d,  $J$  = 7.2 Hz, 1H), 7.07 (d,  $J$  = 8.2 Hz, 1H), 6.74–6.69 (dd,  $J$  = 8.2, 2.7 Hz, 1H), 6.62 (d,  $J$  = 2.6 Hz, 1H), 5.50 (br s, NH), 3.74 (m, 4H), 3.53 (s, 3H), 3.29–3.20 (q,  $J$  = 6.7 Hz, 2H), 2.69 (t,  $J$  = 7.0 Hz, 2H), 2.49 (t,  $J$  = 4.8 Hz, 4H), 1.87 (s, 3H).  $^{13}\text{C}$  NMR (125 MHz,  $\text{CDCl}_3$ )  $\delta$  170.5, 155.0, 143.4, 141.7, 130.9, 130.2, 128.4, 128.2, 117.5, 115.0, 66.9, 63.4, 53.8, 40.8, 32.0, 23.4. HRMS  $m/z$  [ $\text{M} + \text{H}^+$ ] for  $\text{C}_{21}\text{H}_{27}\text{N}_2\text{O}_3$ : calcd, 355.2022; found, 355.2024.

*N*-(2-(4'-((tert-butyl)dimethylsilyloxy)-5-hydroxy-[1,1'-biphenyl]-2-yl)ethyl)acetamide (**11m**).  $^1\text{H}$  NMR (500 MHz,  $\text{CDCl}_3$ )  $\delta$  7.16–7.10 (d,  $J$  = 6.7 Hz, 2H), 7.10–7.06 (d,  $J$  = 8.2 Hz, 1H), 7.00 (br s, OH), 6.91–6.84 (d,  $J$  = 8.4 Hz, 2H), 6.79–6.72 (m, 2H), 5.38 (s, 1H), 3.34–3.21 (q,  $J$  = 6.6 Hz, 2H), 2.78–2.64 (t,  $J$  = 6.9 Hz, 2H), 1.93–1.81 (s, 3H), 1.00 (s, 9H), 0.24 (s, 6H).  $^{13}\text{C}$  NMR (125 MHz,  $\text{CDCl}_3$ )  $\delta$  170.7, 155.0, 154.9, 143.3, 134.6, 130.9, 130.3, 127.8, 120.0, 117.5, 114.7, 41.0, 32.0, 26.0, 23.4, 18.4, –4.1. HRMS (FAB)  $m/z$  [ $\text{M} + \text{Na}^+$ ] for  $\text{C}_{22}\text{H}_{31}\text{NO}_3\text{SiNa}$ : calcd, 408.1965; found, 408.1960.

*N*-(2-(Benzo[d][1,3]dioxol-5-yl)-4-hydroxyphenethyl)acetamide (**11n**).  $^1\text{H}$  NMR (500 MHz,  $\text{CDCl}_3$ )  $\delta$  8.00 (br s, OH), 7.08–6.98 (d,  $J$  = 8.3 Hz, 1H), 6.81–6.73 (m, 2H), 6.73–6.68 (m, 2H), 6.68–6.64 (dd,  $J$  = 7.9, 1.7 Hz, 1H), 5.97–5.92 (s, 2H), 5.70–5.63 (t,  $J$  = 5.7 Hz, 1H), 3.29–3.21 (td,  $J$  = 7.1, 5.6 Hz, 2H), 2.75–2.63 (t,  $J$  = 7.2 Hz, 2H), 1.89–1.81 (s, 3H).  $^{13}\text{C}$  NMR (125 MHz,  $\text{CDCl}_3$ )  $\delta$  171.1, 155.1, 147.5, 146.8, 143.0, 135.4, 130.8, 127.4, 122.4, 117.5, 114.9, 109.8, 108.3, 101.2, 41.1, 31.9, 23.3. HRMS (FAB)  $m/z$  [ $\text{M} + \text{Na}^+$ ] for  $\text{C}_{17}\text{H}_{17}\text{NO}_4\text{Na}$ : calcd, 322.1050; found, 322.1022.

*N*-(4-Hydroxy-2-(pyridin-3-yl)phenethyl)acetamide (**11o**).  $^1\text{H}$  NMR (400 MHz,  $\text{CDCl}_3$ )  $\delta$  8.54 (s, 2H), 7.72 (d,  $J$  = 7.9 Hz, 1H), 7.42–7.34 (dd,  $J$  = 8.0, 4.8 Hz, 1H), 7.14 (d,  $J$  = 8.4 Hz, 1H), 6.90–6.84 (dd,  $J$  = 8.3, 2.7 Hz, 1H), 6.73 (d,  $J$  = 2.7 Hz, 1H), 5.82 (t,  $J$  = 5.9 Hz, 2H), 3.33–3.19 (q,  $J$  = 6.8 Hz, 2H), 2.69 (t,  $J$  = 7.2 Hz, 2H), 1.85 (s, 3H).  $^{13}\text{C}$  NMR (100 MHz,  $\text{CDCl}_3$ )  $\delta$  170.7, 156.1, 149.1, 147.7, 138.8, 138.0, 131.4, 127.3, 123.7, 117.5, 116.4, 100.2, 40.9, 32.0, 23.4. HRMS (FAB)  $m/z$  [ $\text{M} + \text{H}^+$ ] for  $\text{C}_{15}\text{H}_{17}\text{N}_2\text{O}_2$ : calcd, 257.1290; found, 257.1297.

*N*-(4-Hydroxy-2-(pyridin-4-yl)phenethyl)acetamide (**11p**).  $^1\text{H}$  NMR (400 MHz,  $\text{CDCl}_3$ )  $\delta$  8.69–8.60 (m, 2H), 7.25 (d,  $J$  = 1.5 Hz, 2H), 7.17 (d,  $J$  = 8.4 Hz, 1H), 6.90–6.83 (dd,  $J$  = 8.4, 2.7 Hz, 1H), 6.70 (d,  $J$  = 2.7 Hz, 1H), 6.02 (br s, OH), 5.47 (s, 1H), 3.33–3.24 (q,  $J$  = 7.0 Hz, 2H), 2.71 (t,  $J$  = 7.4 Hz, 2H), 1.90 (s, 3H).  $^{13}\text{C}$  NMR (125 MHz,  $\text{CDCl}_3$ )  $\delta$  173.0, 157.1, 152.8, 149.7, 149.6, 141.3, 132.4, 128.0, 126.2, 117.2, 117.1, 116.9, 41.8, 32.8, 22.5. HRMS (FAB)  $m/z$  [ $\text{M} + \text{Na}^+$ ] for  $\text{C}_{15}\text{H}_{16}\text{N}_2\text{O}_2\text{Na}$ : calcd, 279.1104; found, 279.1109.

**General Procedure for Activated Noviose Carbamate Coupling and Followed by Methanolysis of Compounds 11a–p.** Borontrifluoride etherate (6.2  $\mu\text{L}$ , 0.05 mmol) was added to **11a** (0.25 mmol) and activated noviose (0.2 mmol) in 2.5 mL anhydrous  $\text{CH}_2\text{Cl}_2$ . After stirring at room temperature for 2 h, triethylamine (150  $\mu\text{L}$ ) was added and concentrated. The residue was partially purified via column chromatography ( $\text{SiO}_2$ , 100:8  $\text{CH}_2\text{Cl}_2$ :acetone) to give noviose coupled product as a colorless foam, which was used directly for next step.

Triethylamine (0.22 mL, 10%) was added to the cyclic carbonate (100 mg, 0.22 mmol) in MeOH (2.2 mL). After 12 h, the solvent was concentrated and the residue was purified via column chromatography (SiO<sub>2</sub>, 10:1, CH<sub>2</sub>Cl<sub>2</sub>:acetone) to afford inseparable diastereomers **13a** (see following experimental section for diastereoselectivities) as a colorless amorphous solids. Compounds **13b–m** were synthesized using this procedure.

*N*-(2-(5-(((3*R*,4*S*,5*R*)-3,4-Dihydroxy-5-methoxy-6,6-dimethyltetrahydro-2*H*-pyran-2-yl)oxy)-[1,1'-biphenyl]-2-yl)ethyl)acetamide (**13a**). Colorless amorphous solid (63% yield over 2 steps). <sup>1</sup>H NMR (500 MHz, CDCl<sub>3</sub>) δ 7.41–7.28 (m, 3H), 7.28–7.18 (dt, *J* = 5.9, 3.2 Hz, 2H), 7.13 (m, 1H), 6.97 (m, 1H), 6.92–6.78 (dd, *J* = 7.6, 2.7 Hz, 1H), 5.55–5.47 (dd, *J* = 7.7, 2.7 Hz, 1H), 5.39 (m, 1H), 4.14 (m, 2H), 3.58–3.46 (m, 3H), 3.34–3.15 (m, 4H), 3.03 (d, *J* = 5.5 Hz, 1H), 2.77–2.65 (m, 2H), 1.84–1.76 (m, 3H), 1.31 (d, *J* = 4.9 Hz, 3H), 1.21–1.10 (m, 3H). <sup>13</sup>C NMR (125 MHz, CDCl<sub>3</sub>) δ 170.3, 155.4, 143.5, 141.4, 130.8, 129.5, 129.2, 128.5, 127.4, 118.2, 115.2, 98.1, 84.5, 78.4, 71.5, 68.8, 62.0, 40.8, 32.1, 29.2, 23.4, 23.1. HRMS *m/z* [M + H<sup>+</sup>] for C<sub>24</sub>H<sub>32</sub>NO<sub>6</sub>: calcd, 430.2224; found, 430.2227.

*N*-(2-(5-(((3*R*,4*S*,5*R*)-3,4-Dihydroxy-5-methoxy-6,6-dimethyltetrahydro-2*H*-pyran-2-yl)oxy)-3'-fluoro-[1,1'-biphenyl]-2-yl)ethyl)acetamide (**13b**). Colorless amorphous solid (51% yield over 2 steps). <sup>1</sup>H NMR (500 MHz, CDCl<sub>3</sub>) δ 7.39 (dd, 1H, *J* = 7.9, 13.9 Hz), 7.22 (d, 1H, *J* = 8.5 Hz), 7.07 (dd, 2H, *J* = 7.5, 10.5 Hz), 7.02 (dd, 1H, *J* = 2.8, 8.4 Hz), 6.99 (m, 1H), 6.91 (d, 1H, *J* = 2.7 Hz), 5.34 (d, 1H, *J* = 1.3 Hz), 5.28 (s, 1H), 4.20 (d, 1H, *J* = 2.2 Hz), 3.80 (m, 1H), 3.63 (s, 3H), 3.30 (d, 1H), 3.28 (m, 2H), 2.75 (t, 2H, *J* = 7.2 Hz), 2.63 (m, 2H, *J* = 15.9 Hz), 1.87 (s, 3H), 1.41 (s, 3H), 1.28 (s, 3H). <sup>13</sup>C NMR (125 MHz, CDCl<sub>3</sub>) δ 169.9, 163.5–161.6 (d, *J* = 251 Hz) 155.0, 143.2 (d, *J* = 7.8 Hz), 142.1 (d, *J* = 1.8 Hz), 130.9, 130.1, 130.0 (d, *J* = 8.8 Hz), 124.8 (d, *J* = 2.8 Hz), 118.0, 116.0 (d, *J* = 8.8 Hz), 115.4, 114.3 (d, *J* = 21.6 Hz), 93.8, 84.2, 76.0, 71.3, 71.1, 62.0, 40.4, 32.0, 28.6, 23.3, 18.5. HRMS *m/z*: [M + H<sup>+</sup>] for C<sub>24</sub>H<sub>31</sub>FNO<sub>6</sub>: calcd, 448.2180; found, 448.2174. This material was determined to be 95.6% pure (retention time = 6.401) by HPLC (Phenomenex Luna C-18, 5 μm, 10 mm × 250 mm column eluting with 30% CH<sub>3</sub>CN, 70% H<sub>2</sub>O, flow rate 5.0 mL/min).

*N*-(2-(5-(((3*R*,4*S*,5*R*)-3,4-Dihydroxy-5-methoxy-6,6-dimethyltetrahydro-2*H*-pyran-2-yl)oxy)-4'-fluoro-[1,1'-biphenyl]-2-yl)ethyl)acetamide (**13c**). Colorless amorphous solid (57% yield over 2 steps). <sup>1</sup>H NMR (500 MHz, CDCl<sub>3</sub>) δ 7.25 (dd, 2H, *J* = 5.4, 8.6 Hz), 7.18 (d, 1H, *J* = 8.5 Hz), 7.10 (t, 2H, *J* = 8.7 Hz), 7.01 (dd, 1H, *J* = 2.7, 8.5 Hz), 6.87 (d, 1H, *J* = 2.7 Hz), 5.54 (d, 1H, *J* = 2.2 Hz), 5.37 (t, 1H, *J* = 5.2 Hz), 4.20 (dd, 1H, *J* = 3.3, 9.1 Hz), 4.15 (m, 1H), 3.59 (s, 3H), 3.33 (d, 1H, *J* = 9.1 Hz), 3.26 (q, 2H, *J* = 6.9 Hz), 2.97 (s, 1H), 2.81 (s, 1H), 2.72 (t, 2H, *J* = 7.3), 1.87 (s, 3H), 1.36 (s, 3H), 1.22 (s, 3H). <sup>13</sup>C NMR (125 MHz, CDCl<sub>3</sub>) δ 170.2, 163.2–161.3 (d, *J* = 250 Hz), 155.3, 142.3, 137.2 (d, *J* = 3.2 Hz), 130.8, 130.8, 130.7, 129.5, 118.1, 115.4, 115.3, 115.3, 97.9, 84.4, 78.3, 71.4, 68.7, 62.0, 40.6, 32.1, 29.1, 23.4, 23.1. HRMS *m/z* [M + Na<sup>+</sup>] for C<sub>24</sub>H<sub>30</sub>FNO<sub>6</sub>: calcd, 470.1955; found, 470.1958.

*N*-(2-(2'-Chloro-5-(((3*R*,4*S*,5*R*)-3,4-dihydroxy-5-methoxy-6,6-dimethyltetrahydro-2*H*-pyran-2-yl)oxy)-[1,1'-biphenyl]-2-yl)ethyl)acetamide (**13d**). Colorless amorphous solid (62% yield over 2 steps). <sup>1</sup>H NMR (500 MHz, CDCl<sub>3</sub>) δ 7.46 (m, 1H), 7.31 (m, 2H), 7.21 (m, 2H), 7.03 (m, 1H), 6.86 (dd, 1H, *J* = 2.7, 13.2 Hz), 5.55 (m, 1H), 5.42 (s, 1H), 4.20 (dt, 1H, *J* = 3.0, 9.1 Hz), 4.14 (m, 1H), 3.59 (s, 3H), 3.33 (dd, 1H, *J* = 2.5, 9.1 Hz), 3.26 (ddt, 2H, *J* = 4.8, 6.8, 9.3 Hz), 3.11 (s, 1H), 2.93 (s, 1H), 2.58 (tq, 2H, *J* = 7.1, 14.2 Hz), 1.86 (s, 3H), 1.35 (d, 3H, *J* = 2.4 Hz), 1.20 (t, 3H, *J* = 5.8 Hz). <sup>13</sup>C NMR (125 MHz, CDCl<sub>3</sub>) δ 170.2, 155.2, 140.6, 140.5, 139.8, 133.4, 131.4, 130.5, 129.8, 126.9, 118.1, 117.9, 116.05, 97.9, 84.5, 78.4, 71.5, 71.4, 68.7, 62.1, 62.0, 40.2, 40.2, 32.1, 32.1, 29.3, 29.2, 23.5, 23.1, 23.0. HRMS *m/z* [M + Na<sup>+</sup>] for C<sub>24</sub>H<sub>30</sub>ClNO<sub>6</sub>: calcd, 486.1659; found, 486.1652.

*N*-(2-(3'-Chloro-5-(((3*R*,4*S*,5*R*)-3,4-dihydroxy-5-methoxy-6,6-dimethyltetrahydro-2*H*-pyran-2-yl)oxy)-[1,1'-biphenyl]-2-yl)ethyl)acetamide (**13e**). Colorless amorphous solid (55% yield over 2 steps). <sup>1</sup>H NMR (500 MHz, CDCl<sub>3</sub>) δ 7.35 (m, 2H), 7.28 (m, 1H), 7.18 (m, 2H), 7.03 (dd, 1H, *J* = 2.7, 8.5 Hz), 6.87 (d, 1H, *J* = 2.7 Hz), 5.55 (t, 1H, *J* = 2.5 Hz), 5.34 (m, 1H), 4.21 (dd, 1H, *J* = 3.1, 9.1 Hz), 4.16 (m,

1H), 3.60 (s, 3H), 3.34 (dd, 1H, *J* = 1.9, 9.1 Hz), 3.28 (m, 2H), 2.75 (dt, 4H, *J* = 7.3, 14.5 Hz), 1.88 (s, 3H), 1.37 (s, 3H), 1.22 (s, 3H). <sup>13</sup>C NMR (125 MHz, CDCl<sub>3</sub>) δ 170.1, 155.4, 143.5, 142.0, 134.3, 131.0, 130.9, 129.8, 129.4, 128.5, 127.6, 127.4, 118.2, 115.7, 97.9, 84.6, 78.4, 71.5, 68.7, 62.1, 40.8, 32.1, 29.2, 23.6, 23.1. HRMS *m/z* [M + Na<sup>+</sup>] for C<sub>24</sub>H<sub>30</sub>ClNO<sub>6</sub>: calcd, 486.1659; found, 486.1642.

*N*-(2-(5-(((3*R*,4*S*,5*R*)-3,4-Dihydroxy-5-methoxy-6,6-dimethyltetrahydro-2*H*-pyran-2-yl)oxy)-3'-(trifluoromethyl)-[1,1'-biphenyl]-2-yl)ethyl)acetamide (**13f**). Colorless amorphous solid (52% yield over 2 steps). <sup>1</sup>H NMR (500 MHz, CDCl<sub>3</sub>) δ 7.64 (d, 1H, *J* = 7.7 Hz), 7.55 (t, 2H, *J* = 7.6 Hz), 7.49 (m, 1H), 7.23 (d, 1H, *J* = 8.5 Hz), 7.06 (dd, 1H, *J* = 2.7, 8.4 Hz), 6.89 (d, 1H, *J* = 2.7 Hz), 5.56 (d, 1H, *J* = 2.2 Hz), 5.31 (s, 1H), 4.19 (m, 2H), 3.60 (s, 3H), 3.34 (d, 1H, *J* = 9.1 Hz), 3.29 (dd, 2H, *J* = 7.0, 13.3 Hz), 2.72 (t, 2H, *J* = 7.3 Hz), 2.69 (s, 1H), 2.64 (s, 1H), 1.87 (s, 3H), 1.37 (s, 3H), 1.22 (s, 3H). <sup>13</sup>C NMR (125 MHz, CDCl<sub>3</sub>) δ 170.1, 155.4, 142.1, 141.9, 132.6, 131.0, 130.7 (q, *J* = 31.5 Hz), 129.4, 129.0, 125.9 (q, *J* = 3.6, 7.2 Hz), 125.3, 124.2 (q, *J* = 3.6, 7.2 Hz), 123.1, 118.0, 115.8, 97.9, 84.4, 77.4, 71.4, 68.7, 62.0, 40.6, 32.1, 29.8, 29.2, 23.4, 23.0. HRMS *m/z* [M + Na<sup>+</sup>] for C<sub>25</sub>H<sub>30</sub>F<sub>3</sub>NO<sub>6</sub>: calcd, 520.1923; found, 520.1932. This material was determined to be 97.2% pure (retention time = 7.631) by HPLC (Phenomenex Luna C-18, 5 μm, 10 mm × 250 mm column eluting with 30% CH<sub>3</sub>CN, 70% H<sub>2</sub>O, flow rate 5.0 mL/min).

*N*-(2-(5-(((3*R*,4*S*,5*R*)-3,4-Dihydroxy-5-methoxy-6,6-dimethyltetrahydro-2*H*-pyran-2-yl)oxy)-4'-(trifluoromethyl)-[1,1'-biphenyl]-2-yl)ethyl)acetamide (**13g**). Colorless amorphous solid (49% yield over 2 steps). <sup>1</sup>H NMR (400 MHz, CDCl<sub>3</sub>) δ 7.70 (d, *J* = 7.6 Hz, 2H), 7.43 (d, *J* = 7.9 Hz, 2H), 7.24 (d, *J* = 8.4 Hz, 1H), 7.09–7.03 (dd, *J* = 8.6, 2.7 Hz, 1H), 6.90 (d, *J* = 2.7 Hz, 1H), 5.55 (d, *J* = 2.3 Hz, 1H), 5.33 (m, 1H), 4.26–4.11 (m, 2H), 3.60 (s, 3H), 3.36–3.25 (m, 3H), 2.74 (t, *J* = 7.4 Hz, 2H), 2.56 (br s, 2OH), 1.88 (s, 3H), 1.37 (s, 3H), 1.22 (s, 3H). <sup>13</sup>C NMR (125 MHz, MeOD) δ 173.1, 156.8, 146.9, 143.2, 132.1, 130.9, 130.7, 130.5, 130.2, 126.3, 126.2, 124.7, 118.5, 116.8, 100.1, 85.3, 79.5, 72.8, 69.5, 62.1, 41.7, 32.9, 29.2, 23.6, 22.5. HRMS *m/z* [M + Na<sup>+</sup>] for C<sub>25</sub>H<sub>30</sub>F<sub>3</sub>NO<sub>6</sub>: calcd, 520.1923; found, 520.1934.

*N*-(2-(5-(((3*R*,4*S*,5*R*)-3,4-Dihydroxy-5-methoxy-6,6-dimethyltetrahydro-2*H*-pyran-2-yl)oxy)-2'-(methylthio)-[1,1'-biphenyl]-2-yl)ethyl)acetamide (**13h**). Colorless amorphous solid (63% yield over 2 steps). <sup>1</sup>H NMR (400 MHz, CDCl<sub>3</sub>) δ 7.36 (t, 1H, *J* = 7.0 Hz), 7.27 (m, 3H), 7.09 (m, 1H), 7.01 (m, 1H), 6.87 (s, 1H), 5.64 (s, 1H), 5.54 (m, 1H), 4.16 (m, 2H), 3.32 (d, 2H, *J* = 8.8 Hz), 3.27 (m, 2H), 3.06 (s, 1H), 2.56 (t, 2H, *J* = 6.2 Hz), 2.36 (d, 3H, *J* = 7.6 Hz), 1.83 (s, 3H), 1.33 (s, 3H), 1.20 (s, 3H). <sup>13</sup>C NMR (100 MHz, CDCl<sub>3</sub>) δ 170.3, 155.1, 155.0, 141.0, 138.9, 130.5, 130.1, 129.8, 128.4, 124.6, 124.2, 118.3, 116.2, 115.9, 97.9, 84.5, 78.3, 71.5, 68.7, 62.0, 53.6, 40.1, 31.7, 29.3, 23.3, 15.3, 15.2. HRMS *m/z* [M + Na<sup>+</sup>] for C<sub>25</sub>H<sub>33</sub>NO<sub>6</sub>SNa: calcd, 498.1926; found, 498.1925. This material was determined to be 95% pure (retention time = 7.465) by HPLC (Phenomenex Luna C-18, 5 μm, 10 mm × 250 mm column eluting with 30% CH<sub>3</sub>CN, 70% H<sub>2</sub>O, flow rate 5.0 mL/min).

*N*-(2-(5-(((3*R*,4*S*,5*R*)-3,4-Dihydroxy-5-methoxy-6,6-dimethyltetrahydro-2*H*-pyran-2-yl)oxy)-2'-methoxy-[1,1'-biphenyl]-2-yl)ethyl)acetamide (**13i**). Colorless amorphous solid (41% yield over 2 steps). <sup>1</sup>H NMR (500 MHz, CDCl<sub>3</sub>) δ 7.36 (ddd, 1H, *J* = 1.8, 7.6, 8.2 Hz), 7.18 (d, 1H, *J* = 8.3 Hz), 7.12 (t, 1H, *J* = 5.8 Hz), 7.02 (m, 3H), 6.87 (dd, 1H, *J* = 2.3, 11.3 Hz), 5.54 (s, 1H), 5.39 (s, 1H), 4.21 (dt, 1H, *J* = 3.3, 9.0 Hz), 4.15 (m, 1H), 3.77 (d, 3H, *J* = 6.9 Hz), 3.60 (s, 3H), 3.33 (d, 1H, *J* = 8.7 Hz), 3.29 (m, 2H), 2.73 (s, 1H), 2.66 (s, 1H), 2.60 (dd, 2H, *J* = 6.5, 12.8 Hz), 1.84 (s, 3H), 1.37 (s, 3H), 1.24 (s, 3H). <sup>13</sup>C NMR (125 MHz, CDCl<sub>3</sub>) δ 170.0, 156.4, 155.2, 139.9, 131.2, 130.8, 130.2, 130.0, 129.2, 120.9, 118.6, 118.3, 115.7, 115.2, 111.4, 111.2, 98.0, 97.9, 84.5, 78.2, 71.4, 68.7, 62.0, 55.9, 55.9, 40.3, 31.9, 30.2, 29.3, 29.2, 23.4, 23.1. HRMS *m/z* [M + H<sup>+</sup>] for C<sub>25</sub>H<sub>34</sub>NO<sub>7</sub>: calcd, 460.2335; found, 460.2336. This material was determined to be 96.1% pure (retention time = 5.057) by HPLC (Phenomenex Luna C-18, 5 μm, 10 mm × 250 mm column eluting with 30% CH<sub>3</sub>CN, 70% H<sub>2</sub>O, flow rate 5.0 mL/min).

*N*-(2-(5-(((3*R*,4*S*,5*R*)-3,4-Dihydroxy-5-methoxy-6,6-dimethyltetrahydro-2*H*-pyran-2-yl)oxy)-3'-methoxy-[1,1'-biphenyl]-2-yl)ethyl)acetamide (**13j**). Colorless amorphous solid (53% yield over 2 steps). <sup>1</sup>H NMR (400 MHz, CDCl<sub>3</sub>) δ 7.31 (t, *J* = 7.9 Hz, 1H), 7.17 (d, *J* =

8.5 Hz, 1H), 7.02–6.96 (dd,  $J = 8.5, 2.7$  Hz, 1H), 6.92–6.83 (m, 4H), 6.81 (d,  $J = 1.5$  Hz, 2H), 5.54 (d,  $J = 2.2$  Hz, 1H), 5.45 (s, 1H), 4.25–4.16 (dd,  $J = 9.1, 3.2$  Hz, 1H), 4.17–4.10 (dd,  $J = 3.3, 2.2$  Hz, 1H), 3.82 (s, 3H), 3.58 (s, 3H), 3.39–3.20 (m, 3H), 3.24 (br s, OH), 2.97 (br s, OH), 2.75 (t,  $J = 7.1$  Hz, 2H), 1.85 (s, 3H), 1.35 (s, 3H), 1.20 (s, 3H).  $^{13}\text{C}$  NMR (125 MHz,  $\text{CDCl}_3$ )  $\delta$  170.4, 159.5, 155.3, 143.3, 142.8, 130.8, 129.5, 129.5, 121.7, 118.0, 115.3, 115.1, 112.7, 98.1, 84.5, 78.4, 71.5, 68.7, 62.0, 55.4, 40.9, 32.0, 29.1, 23.4, 23.1. HRMS  $m/z$  [ $\text{M} + \text{H}^+$ ] for  $\text{C}_{25}\text{H}_{34}\text{NO}_7$ : calcd, 460.2335; found, 460.2322.

***N*-(2-(5-(((3*R*,4*S*,5*R*)-3,4-dihydroxy-5-methoxy-6,6-dimethyltetrahydro-2*H*-pyran-2-yl)oxy)-3'-methyl-[1,1'-biphenyl]-2-yl)ethyl)acetamide (13k)**. Colorless amorphous solid (44% yield over 2 steps).  $^1\text{H}$  NMR (400 MHz,  $\text{CDCl}_3$ )  $\delta$  7.32–7.27 (m, 1H), 7.16 (d,  $J = 6.6$  Hz, 2H), 7.10–7.04 (m, 2H), 6.99 (d,  $J = 8.5$  Hz, 1H), 6.88 (s, 1H), 5.55 (s, 1H), 5.41 (s, 1H), 4.25–4.08 (m, 2H), 3.57 (s, 3H), 3.37–3.20 (m, 5H), 2.75 (t,  $J = 7.0$  Hz, 2H), 2.39 (s, 3H), 1.83 (s, 3H), 1.35 (s, 3H), 1.20 (s, 3H).  $^{13}\text{C}$  NMR (100 MHz,  $\text{CDCl}_3$ )  $\delta$  170.4, 155.3, 143.6, 141.3, 138.1, 130.8, 130.0, 129.5, 128.3, 128.1, 126.3, 118.1, 115.1, 98.1, 84.5, 78.4, 71.5, 68.7, 62.0, 40.9, 32.0, 29.2, 23.4, 23.1, 21.7. HRMS  $m/z$  [ $\text{M} + \text{H}^+$ ] for  $\text{C}_{25}\text{H}_{33}\text{NO}_6\text{Na}$ : calcd, 466.2206; found, 466.2203.

***N*-(2-(5-(((3*R*,4*S*,5*R*)-3,4-dihydroxy-5-methoxy-6,6-dimethyltetrahydro-2*H*-pyran-2-yl)oxy)-3'-(morpholinomethyl)-[1,1'-biphenyl]-2-yl)ethyl)acetamide (13l)**. Colorless amorphous solid (47% yield over 2 steps).  $^1\text{H}$  NMR (500 MHz,  $\text{CDCl}_3$ )  $\delta$  7.41–7.29 (m, 2H), 7.27 (m, 1H), 7.19 (d,  $J = 8.1$  Hz, 2H), 7.04–6.99 (dd,  $J = 8.5, 2.7$  Hz, 1H), 6.91 (d,  $J = 2.7$  Hz, 1H), 5.55 (d,  $J = 2.4$  Hz, 1H), 5.35 (s, 1H), 4.26–4.18 (dd,  $J = 9.0, 3.3$  Hz, 1H), 4.15 (t,  $J = 2.8$  Hz, 1H), 3.72 (t,  $J = 4.7$  Hz, 4H), 3.59 (s, 3H), 3.56 (s, 2H), 3.34 (d,  $J = 9.0$  Hz, 1H), 3.30–3.21 (q,  $J = 6.7$  Hz, 2H), 2.75 (t,  $J = 7.1$  Hz, 2H), 2.58–2.41 (m, 6H), 1.85 (s, 3H), 1.36 (s, 3H), 1.23 (s, 3H).  $^{13}\text{C}$  NMR (125 MHz,  $\text{CDCl}_3$ )  $\delta$  155.4, 143.4, 141.5, 137.8, 130.9, 130.1, 129.6, 128.5, 128.3, 128.2, 118.2, 115.3, 98.1, 84.6, 78.4, 71.5, 68.8, 67.1, 63.5, 62.0, 53.8, 40.7, 32.2, 29.2, 23.5, 23.2. HRMS (FAB)  $m/z$  [ $\text{M} + \text{Na}^+$ ] for  $\text{C}_{29}\text{H}_{40}\text{N}_2\text{O}_7\text{Na}$ : calcd, 551.2728; found, 551.2734.

***N*-(2-(5-(((3*R*,4*S*,5*R*)-3,4-dihydroxy-5-methoxy-6,6-dimethyltetrahydro-2*H*-pyran-2-yl)oxy)-4'-hydroxy-[1,1'-biphenyl]-2-yl)ethyl)acetamide (13m)**. After cyclic carbonate hydrolysis following the same procedure as compound 13a, the crude TBS protected compound was dissolved in THF (2 mL) and tetrabutylammonium fluoride (1.5 equiv) was added dropwise at 0 °C under argon atmosphere. After 1 h, the reaction was quenched with water and extracted with EtOAc (3 × 10 mL); combined organic fractions were washed with saturated aqueous sodium chloride, dried over anhydrous  $\text{Na}_2\text{SO}_4$ , filtered, and concentrated. The residue was purified by column chromatography ( $\text{SiO}_2$ ; 10:1,  $\text{CH}_2\text{Cl}_2$ :acetone) to afford acetamide 13m as an amorphous solid (40% yield over 3 steps).  $^1\text{H}$  NMR (500 MHz, MeOD)  $\delta$  7.20 (d,  $J = 8.4$  Hz, 1H), 7.15–7.08 (d,  $J = 8.4$  Hz, 2H), 6.96 (dd,  $J = 8.4, 2.6$  Hz, 1H), 6.85–6.79 (m, 3H), 5.45 (d,  $J = 2.4$  Hz, 1H), 4.12 (dd,  $J = 9.3, 3.3$  Hz, 1H), 3.96 (t,  $J = 2.8$  Hz, 1H), 3.59 (s, 3H), 3.21 (d,  $J = 9.3$  Hz, 1H), 3.16 (dd,  $J = 8.5, 6.5$  Hz, 2H), 2.70 (dd,  $J = 8.5, 6.5$  Hz, 2H), 1.84 (s, 3H), 1.32 (s, 3H), 1.18 (s, 3H).  $^{13}\text{C}$  NMR (125 MHz, MeOD)  $\delta$  173.1, 157.7, 156.6, 144.7, 134.0, 131.7, 131.2, 131.1, 118.9, 116.0, 115.7, 100.1, 85.4, 79.4, 72.8, 69.5, 62.1, 41.8, 33.0, 29.2, 23.6, 22.5. HRMS (FAB)  $m/z$  [ $\text{M} + \text{Na}^+$ ] for  $\text{C}_{24}\text{H}_{31}\text{NO}_7\text{Na}$ : calcd, 468.1998; found, 468.1999.

***N*-(2-(Benzo[d][1,3]dioxol-5-yl)-4-(((3*R*,4*S*,5*R*)-3,4-dihydroxy-5-methoxy-6,6-dimethyltetrahydro-2*H*-pyran-2-yl)oxy)phenethyl)acetamide (13n)**. Colorless amorphous solid (51% yield over 2 steps).  $^1\text{H}$  NMR (500 MHz,  $\text{CDCl}_3$ )  $\delta$  7.15 (d,  $J = 8.5$  Hz, 1H), 7.00–6.96 (dd,  $J = 8.5, 2.7$  Hz, 1H), 6.88 (d,  $J = 2.6$  Hz, 1H), 6.84 (d,  $J = 7.9$  Hz, 1H), 6.76 (d,  $J = 1.6$  Hz, 1H), 6.74–6.69 (m, 1H), 6.01 (s, 2H), 5.54 (d,  $J = 2.4$  Hz, 1H), 5.40 (s, 1H), 4.21 (dd,  $J = 9.1, 3.3$  Hz, 1H), 4.14 (t,  $J = 2.7$  Hz, 2H), 3.58 (s, 3H), 3.33 (d,  $J = 9.1$  Hz, 1H), 3.30–3.23 (q,  $J = 6.9$  Hz, 2H), 3.11 (br s, OH), 2.92 (br s, OH), 2.74 (t,  $J = 7.2$  Hz, 2H), 1.86 (s, 3H), 1.34 (s, 3H), 1.20 (s, 3H).  $^{13}\text{C}$  NMR (125 MHz,  $\text{CDCl}_3$ )  $\delta$  170.2, 155.4, 147.7, 147.0, 143.1, 135.3, 130.8, 129.7, 122.6, 118.3, 115.2, 109.9, 108.4, 101.3, 98.1, 84.6, 78.4, 71.5, 68.8, 62.0, 40.8, 32.1, 29.2, 23.4, 23.2. HRMS (FAB)  $m/z$  [ $\text{M} + \text{Na}^+$ ] for  $\text{C}_{25}\text{H}_{31}\text{NO}_8\text{Na}$ : calcd, 496.1947; found, 496.1940. This material was

determined to be 98.4% pure (retention time = 4.384) by HPLC (Phenomenex Luna C-18, 5  $\mu\text{m}$ , 10 mm × 250 mm column eluting with 40%  $\text{CH}_3\text{CN}$ , 60%  $\text{H}_2\text{O}$ , flow rate 5.0 mL/min).

***N*-(4-(((3*R*,4*S*,5*R*)-3,4-dihydroxy-5-methoxy-6,6-dimethyltetrahydro-2*H*-pyran-2-yl)oxy)-2-(pyridin-3-yl)phenethyl)acetamide (13o)**. Colorless amorphous solid (37% yield over 2 steps).  $^1\text{H}$  NMR (500 MHz,  $\text{CDCl}_3$ )  $\delta$  8.55 (d,  $J = 3.9$  Hz, 1H), 8.49 (s, 1H), 7.60 (m, 1H), 7.35 (dd,  $J = 7.8, 4.5$  Hz, 1H), 7.20 (d,  $J = 8.5$  Hz, 1H), 7.05–6.99 (dd,  $J = 8.4, 2.7$  Hz, 1H), 6.85 (d,  $J = 2.6$  Hz, 1H), 5.52 (d,  $J = 2.4$  Hz, 1H), 5.36 (s, 1H), 4.14 (dd,  $J = 3.4, 9.1$  Hz, 1H), 4.10 (t,  $J = 2.7$  Hz, 1H), 3.59 (s, 3H), 3.31 (d,  $J = 9.0$  Hz, 1H), 3.27–3.20 (m, 2H), 2.68 (t,  $J = 7.3$  Hz, 2H), 1.86 (s, 3H), 1.33 (s, 3H), 1.17 (s, 3H).  $^{13}\text{C}$  NMR (125 MHz,  $\text{CDCl}_3$ )  $\delta$  170.2, 155.5, 149.8, 148.7, 139.5, 136.8, 131.1, 131.0, 130.6, 129.8, 123.4, 118.3, 118.2, 116.1, 98.0, 84.5, 78.5, 71.4, 68.7, 62.1, 40.7, 32.2, 29.2, 23.5, 23.1. HRMS (FAB)  $m/z$  [ $\text{M} + \text{Na}^+$ ] for  $\text{C}_{23}\text{H}_{31}\text{N}_2\text{O}_6$ : calcd, 431.2182; found, 431.2194.

***N*-(4-(((3*R*,4*S*,5*R*)-3,4-dihydroxy-5-methoxy-6,6-dimethyltetrahydro-2*H*-pyran-2-yl)oxy)-2-(pyridin-4-yl)phenethyl)acetamide (13p)**. Colorless amorphous solid (42% yield over 2 steps).  $^1\text{H}$  NMR (400 MHz,  $\text{CDCl}_3$ )  $\delta$  8.73–8.63 (dd,  $J = 5.7, 3.9$  Hz, 2H), 7.27–7.23 (m, 3H), 7.11–7.03 (m, 1H), 6.86 (t,  $J = 2.8$  Hz, 1H), 5.55 (d,  $J = 2.3$  Hz, 1H), 5.41–5.31 (m, 2H), 4.26–4.13 (m, 2H), 4.05 (d,  $J = 6.9$  Hz, 1H), 3.61 (s, 3H), 3.36–3.25 (m, 2H), 2.78–2.71 (dd,  $J = 8.3, 6.8$  Hz, 2H), 1.90 (s, 3H), 1.39 (s, 3H), 1.24 (s, 3H).  $^{13}\text{C}$  NMR (125 MHz,  $\text{CDCl}_3$ )  $\delta$  170.1, 155.5, 149.8, 140.5, 131.4, 129.1, 124.4, 117.9, 116.3, 98.0, 94.1, 84.5, 71.5, 71.4, 68.7, 62.1, 40.7, 32.2, 29.2, 28.8, 23.5, 23.1, 18.7. HRMS (FAB)  $m/z$  [ $\text{M} + \text{Na}^+$ ] for  $\text{C}_{23}\text{H}_{30}\text{N}_2\text{O}_6\text{Na}$ : calcd, 453.2001; found, 453.1972.

**(*Z*)-4-(Benzyloxy)-2-(methoxymethoxy)-1-(2-nitrovinyl)benzene (15)**. Nitromethane (11.5 mL) was added to a mixture of aldehyde 14 (1.24 g, 4.6 mmol) and ammonium acetate (0.63 g, 8.2 mmol) and heated to 50 °C. Upon completion (~20 min), the reaction mixture was cooled to room temperature and purified without workup by column chromatography ( $\text{SiO}_2$ , 4:1, Hex:EtOAc) to afford nitrostyrene 15 as a colorless oil (1.22 g, 84%).  $^1\text{H}$  NMR (400 MHz,  $\text{CDCl}_3$ )  $\delta$  8.17 (d,  $J = 13.4$  Hz, 1H), 7.80 (d,  $J = 13.6$  Hz, 1H), 7.50–7.32 (m, 6H), 6.88 (d,  $J = 2.5$  Hz, 1H), 6.67 (m, 1H), 5.30 (s, 2H), 5.12 (s, 2H), 3.52 (s, 3H).  $^{13}\text{C}$  NMR (100 MHz,  $\text{CDCl}_3$ )  $\delta$  163.4, 159.0, 136.1, 136.0, 135.6, 133.5, 128.8, 128.5, 127.7, 127.7, 113.0, 108.6, 102.2, 94.7, 70.5, 56.6. HRMS (FAB)  $m/z$  [ $\text{M} + \text{Na}^+$ ] for  $\text{C}_{17}\text{H}_{17}\text{NO}_5\text{Na}$ : calcd, 338.1004; found, 338.1007.

**4'-(Benzyloxy)-2'-(methoxymethoxy)-2-nitro-1,2,3,6-tetrahydro-1,1'-biphenyl (16)**. Nitrostyrene 15 (0.65 g, 2.06 mmol) was dissolved in toluene (0.6 mL) in a 2 mL sealed tube and cooled to –78 °C. Butadiene was bubbled into the solution to double the volume, and then the tube was sealed and heated to reflux for 48 h. To prevent bumping of the butadiene gas, the tube was cooled again to –78 °C and used directly in purification by column chromatography ( $\text{SiO}_2$ ; 3:1, Hex:EtOAc) to afford cyclohexene adduct 16 (0.72 g, 95%).  $^1\text{H}$  NMR (500 MHz,  $\text{CDCl}_3$ )  $\delta$  7.40 (m, 4H), 7.36–7.28 (m, 1H), 7.06 (d,  $J = 8.4$  Hz, 1H), 6.80 (d,  $J = 2.4$  Hz, 1H), 6.56 (dd,  $J = 8.4, 2.5$  Hz, 1H), 5.86–5.77 (m, 1H), 5.71 (ddd,  $J = 9.8, 5.1, 2.3$  Hz, 1H), 5.27–5.20 (m, 1H), 5.20 (s, 2H), 5.00 (s, 2H), 3.70 (dt,  $J = 17.0, 8.7$  Hz, 1H), 3.49 (s,  $J = 12.7$  Hz, 3H), 2.84–2.74 (m, 1H), 2.71 (ddd,  $J = 13.2, 8.4, 1.5$  Hz, 1H), 2.45 (m, 2H).  $^{13}\text{C}$  NMR (125 MHz,  $\text{CDCl}_3$ )  $\delta$  159.2, 156.1, 136.9, 129.3, 128.6, 127.0, 122.5, 120.9, 107.8, 120.6, 94.6, 85.6, 70.1, 31.5, 31.3, 29.7.

***N*-(4'-(Benzyloxy)-2'-(methoxymethoxy)-1,2,3,6-tetrahydro-1,1'-biphenyl-2-yl)acetamide (17)**. Nitro compound 16 (0.23 g, 0.62 mmol) was dissolved in isopropyl alcohol (12.4 mL) and aqueous 1 M HCl (6.2 mL). Zinc dust (811 mg, 12.4 mmol) was added sequentially, and the mixture was stirred vigorously at 50 °C for 1.5 h. After cooling to room temperature, saturated  $\text{NaHCO}_3$  (8 mL) was added and the resulting mixture was stirred for an additional 20 min. The solids were removed by filtration, and the remaining solution was extracted with  $\text{CH}_2\text{Cl}_2$  (3 × 20 mL). The organic layers were combined and washed with aqueous saturated sodium chloride solution, dried over  $\text{Na}_2\text{SO}_4$ , filtered, and concentrated to afford amine as a colorless oil (0.20 g, 0.59 mmol, 95%), which was used for next reaction without further purification.

Acetic anhydride (62  $\mu$ L, 0.65 mmol) and triethylamine (95  $\mu$ L, 0.68 mmol) were added to a solution of the amine (0.62 mmol) in  $\text{CH}_2\text{Cl}_2$  (6.2 mL) at room temperature. After 3 h, the reaction was quenched with saturated aqueous ammonium chloride and extracted with  $\text{CH}_2\text{Cl}_2$  ( $3 \times 10$  mL); combined organic fractions were washed with saturated aqueous NaCl, dried over anhydrous  $\text{Na}_2\text{SO}_4$ , filtered, and concentrated. The residue was purified by column chromatography ( $\text{SiO}_2$ ; 3:1, Hex:EtOAc) to afford acetamide **17** (0.17 g, 74%).  $^1\text{H}$  NMR (500 MHz,  $\text{CDCl}_3$ )  $\delta$  7.42 (d,  $J = 7.8$  Hz, 2H), 7.39–7.33 (t,  $J = 7.2$  Hz, 2H), 7.33–7.28 (m, 1H), 7.12 (d,  $J = 8.5$  Hz, 1H), 6.77 (s, 1H), 6.63 (d,  $J = 8.5$  Hz, 1H), 5.90 (d,  $J = 8.4$  Hz, 1H), 5.71 (d,  $J = 36.2$  Hz, 2H), 5.17 (s, 2H), 5.02 (s, 2H), 4.36–4.23 (dtd,  $J = 13.8$ , 10.4, 9.9, 7.2 Hz, 1H), 3.50 (s, 3H), 3.31–3.22 (dd,  $J = 18.6$ , 7.9 Hz, 1H), 2.59 (d,  $J = 17.3$  Hz, 1H), 2.33 (s, 2H), 2.02–1.93 (m, 1H), 1.74 (s, 3H).  $^{13}\text{C}$  NMR (125 MHz,  $\text{CDCl}_3$ )  $\delta$  169.8, 158.3, 156.1, 136.9, 128.5, 128.3, 127.9, 127.6, 126.7, 125.0, 124.4, 108.0, 102.9, 95.6, 70.0, 56.2, 48.8, 37.4, 33.0, 32.6, 23.1. HRMS (FAB)  $m/z$  [ $M + \text{Na}^+$ ] for  $\text{C}_{23}\text{H}_{27}\text{NO}_4\text{Na}$ : calcd, 404.1832; found, 404.1827.

*N*-(4'-(Benzyloxy)-2'-hydroxy-1,2,3,6-tetrahydro-[1,1'-biphenyl]-2-yl)acetamide (**18**). Catalytic amount of concd HCl (few drops) was added to MOM protected phenol **17** (0.27 g, 0.71 mmol) in methanol (7.1 mL) and stirred vigorously at 50  $^\circ\text{C}$  for overnight. Upon completion the reaction, mixture was concentrated and residue was purified by column chromatography ( $\text{SiO}_2$ ; 5:100, MeOH: $\text{CH}_2\text{Cl}_2$ ) to afford phenol **18** (0.19 g, 81%).  $^1\text{H}$  NMR (400 MHz,  $\text{CDCl}_3$ )  $\delta$  8.86 (s, 1H), 7.41–7.25 (m, 5H), 7.01 (d,  $J = 8.5$  Hz, 1H), 6.73 (d,  $J = 2.4$  Hz, 1H), 6.48 (d,  $J = 6.0$  Hz, 1H), 5.73 (m, 1H), 5.65 (m, 1H), 4.96 (s, 2H), 4.26 (m, 1H), 3.42 (m, 1H), 2.55–2.12 (m, 4H), 1.98 (s, 3H).  $^{13}\text{C}$  NMR (100 MHz,  $\text{CDCl}_3$ )  $\delta$  173.2, 158.3, 155.4, 136.9, 128.5, 128.0, 127.9, 127.6, 127.0, 123.9, 121.1, 107.2, 103.4, 69.9, 51.9, 50.0, 36.6, 31.6, 21.0. HRMS (FAB)  $m/z$  [ $M + \text{Na}^+$ ] for  $\text{C}_{21}\text{H}_{23}\text{NO}_3\text{Na}$ : calcd, 360.1576; found, 360.1571.

2'-Acetamido-4-(benzyloxy)-1',2',3',6'-tetrahydro-[1,1'-biphenyl]-2-yl trifluoromethanesulfonate (**19**). A solution of phenol (0.19 g, 0.58 mmol) in anhydrous  $\text{CH}_2\text{Cl}_2$  (5.8 mL) and triethylamine (0.12 mL, 0.87 mmol) followed by *N*-phenyl-bis-(trifluoromethanesulfonimide) (0.31 g, 0.87 mmol) were added at 0  $^\circ\text{C}$ . Upon completion, the reaction was quenched by addition of water (50 mL), washed with saturated aqueous NaCl solution, dried ( $\text{Na}_2\text{SO}_4$ ), filtered, and concentrated. The residue was purified by column chromatography ( $\text{SiO}_2$ ; 3:1, Hex:EtOAc) to afford triflate **19** as a pale-yellow oil (0.23 g, 0.49 mmol, 85%).  $^1\text{H}$  NMR (400 MHz,  $\text{CDCl}_3$ )  $\delta$  7.45–7.31 (m, 6H), 7.00 (d,  $J = 11.2$  Hz, 1H), 6.84 (d,  $J = 2.4$  Hz, 1H), 5.70 (m, 2H), 5.60 (d,  $J = 9.3$  Hz, 1H), 5.04 (s, 2H), 4.53–4.38 (dt,  $J = 15.2$ , 10.2 Hz, 1H), 3.18–3.03 (td,  $J = 11.2$ , 5.2 Hz, 1H), 2.63–2.50 (dd,  $J = 16.2$ , 4.2 Hz, 1H), 2.42–2.32 (m, 1H), 2.28–2.15 (m, 1H), 2.11–1.97 (t,  $J = 14.5$  Hz, 1H), 1.71 (s, 3H).  $^{13}\text{C}$  NMR (100 MHz,  $\text{CDCl}_3$ )  $\delta$  169.7, 158.2, 147.6, 136.0, 129.9, 128.8, 128.5, 128.1, 127.7, 126.1, 125.4, 115.8, 108.2, 70.7, 48.3, 38.5, 34.8, 33.7, 23.2. HRMS (FAB)  $m/z$  [ $M + \text{Na}^+$ ] for  $\text{C}_{22}\text{H}_{22}\text{F}_3\text{NO}_5\text{Na}$ : calcd, 492.106318; found, 492.1067.

*N*-(4'-(Benzyloxy)-3''-fluoro-1,2,3,6-tetrahydro-[1,1':2',1''-terphenyl]-2-yl)acetamide (**20a**). Followed same Suzuki coupling procedure as described above for **8a**.  $^1\text{H}$  NMR (400 MHz,  $\text{CDCl}_3$ )  $\delta$  7.47–7.30 (m, 7H), 7.13–7.05 (t,  $J = 8.9$  Hz, 1H), 7.05–7.00 (t,  $J = 7.2$  Hz, 2H), 6.97 (d,  $J = 10.9$  Hz, 1H), 6.80 (d,  $J = 2.7$  Hz, 1H), 5.72–5.53 (m, 2H), 5.06 (s, 2H), 4.91 (d,  $J = 8.7$  Hz, 1H), 4.36–4.24 (m, 1H), 2.90–2.75 (dd,  $J = 19.2$ , 8.2 Hz, 1H), 2.59–2.45 (dt,  $J = 16.3$ , 4.4 Hz, 1H), 2.36 (m, 2H), 1.75 (s, 3H).  $^{13}\text{C}$  NMR (100 MHz,  $\text{CDCl}_3$ )  $\delta$  169.3, 156.8, 143.9, 142.3, 137.0, 132.6, 130.2, 130.2, 128.8, 128.5, 128.2, 127.8, 126.7, 125.2, 125.0, 116.4, 116.2, 116.0, 115.2, 114.5, 114.3, 70.2, 49.4, 40.5, 35.3, 33.4, 23.5. HRMS (FAB)  $m/z$  [ $M + \text{Na}^+$ ] for  $\text{C}_{27}\text{H}_{26}\text{FNO}_2\text{Na}$ : calcd, 438.1840; found, 438.1818.

*N*-(4'-(Benzyloxy)-3''-(trifluoromethyl)-1,2,3,6-tetrahydro-[1,1':2',1''-terphenyl]-2-yl)acetamide (**20b**). Followed same Suzuki coupling procedure as described above for **8a**.  $^1\text{H}$  NMR (400 MHz,  $\text{CDCl}_3$ )  $\delta$  7.73–7.30 (m, 10H), 7.05 (d,  $J = 8.7$  Hz, 1H), 6.85 (s, 1H), 5.66 (m, 2H), 5.16 (d,  $J = 8.5$  Hz, 1H), 5.08 (s, 2H), 4.43–4.29 (m, 1H), 2.90–2.74 (q,  $J = 10.0$ , 9.0 Hz, 1H), 2.50 (d,  $J = 17.7$  Hz, 1H), 2.40–2.28 (dd,  $J = 6.9$ , 3.9 Hz, 2H), 1.75 (s, 3H).  $^{13}\text{C}$  NMR (101

MHz,  $\text{CDCl}_3$ )  $\delta$  169.7, 156.9, 142.4, 141.9, 136.9, 132.6, 131.1, 130.8, 129.1, 128.7, 128.6, 128.2, 127.7, 126.6, 126.0, 125.9, 125.0, 124.3, 124.2, 116.3, 115.3, 70.2, 49.4, 40.6, 35.2, 33.1, 23.4. HRMS (FAB)  $m/z$  [ $M + \text{Na}^+$ ] for  $\text{C}_{28}\text{H}_{26}\text{F}_3\text{NO}_2\text{Na}$ : calcd, 488.1813; found, 488.1812.

*N*-(3''-Fluoro-4'-hydroxy-1,2,3,6-tetrahydro-[1,1':2',1''-terphenyl]-2-yl)acetamide (**21a**). 1,2-Ethanedithiol (0.22 mL, 2.66 mmol) and  $\text{BF}_3 \cdot \text{OEt}_2$  (0.176 mL, 1.4 mmol) were added to benzyl ether **20a** (64 mg, 0.14 mmol) in  $\text{CH}_2\text{Cl}_2$  (1.8 mL). After 8 h at room temperature, reaction mixture was concentrated and purified by column chromatography ( $\text{SiO}_2$ ; 1:10, MeOH: $\text{CH}_2\text{Cl}_2$ ) to afford phenol **21a** as an amorphous solid (45 mg, 0.12 mmol, 86%).  $^1\text{H}$  NMR (500 MHz,  $\text{CDCl}_3$ )  $\delta$  8.98 (s, 1H), 7.40–7.34 (q,  $J = 7.1$ , 6.2 Hz, 1H), 7.27 (d,  $J = 7.6$  Hz, 1H), 7.10–7.01 (m, 2H), 6.96 (d,  $J = 9.4$  Hz, 1H), 6.84–6.79 (dd,  $J = 8.5$ , 2.6 Hz, 1H), 6.68 (d,  $J = 2.6$  Hz, 1H), 5.73–5.52 (m, 2H), 4.51–4.38 (dt,  $J = 9.9$ , 5.0 Hz, 1H), 2.88–2.77 (q,  $J = 9.5$ , 7.9 Hz, 1H), 2.43 (d,  $J = 17.3$  Hz, 1H), 2.34 (m, 2H), 2.18 (s, 1H), 1.77 (s, 3H).  $^{13}\text{C}$  NMR (125 MHz,  $\text{CDCl}_3$ )  $\delta$  170.5, 163.6–161.7 (d,  $J = 244.0$  Hz), 155.2, 144.2 (d,  $J = 7.6$  Hz), 142.3, 130.8, 130.1 (d,  $J = 8.4$  Hz), 128.2, 127.0, 125.0 (d,  $J = 2.2$  Hz), 124.7, 116.5, 116.3 (d,  $J = 20.3$  Hz), 115.9, 114.2 (d,  $J = 20.3$  Hz, 49.6, 40.8, 35.5, 33.5, 23.2. HRMS (FAB)  $m/z$  [ $M + \text{Na}^+$ ] for  $\text{C}_{21}\text{H}_{25}\text{FNO}_2\text{Na}$ : 348.1376; found, 348.1379.

*N*-(4'-Hydroxy-3''-(trifluoromethyl)-1,2,3,6-tetrahydro-[1,1':2',1''-terphenyl]-2-yl)acetamide (**21b**). Followed same procedure as for **21a**.  $^1\text{H}$  NMR (400 MHz,  $\text{CDCl}_3$ )  $\delta$  9.20 (s, 1H), 7.66–7.56 (m, 4H), 7.29 (d,  $J = 8.5$  Hz, 1H), 6.82–6.72 (d,  $J = 10.5$  Hz, 1H), 6.65 (s, 1H), 5.65 (m, 1H), 5.54 (m, 1H), 5.21 (d,  $J = 9.7$  Hz, 2H), 4.56–4.33 (m, 1H), 2.76–2.61 (m, 1H), 2.46–2.24 (m, 3H), 1.75 (s, 3H).  $^{13}\text{C}$  NMR (100 MHz,  $\text{CDCl}_3$ )  $\delta$  170.5, 155.3, 142.7, 142.0, 132.6, 130.8, 130 (q,  $J = 32.5$  Hz), 128.8, 128.4, 126.9, 126.0 (m), 124.7, 124.0 (m), 116.7, 116.0, 49.6, 40.9, 35.6, 33.5, 23.2. HRMS (FAB)  $m/z$  [ $M + \text{Na}^+$ ] for  $\text{C}_{21}\text{H}_{20}\text{F}_3\text{NO}_2\text{Na}$ : calcd, 398.1344; found, 398.1346.

*N*-(4'-(((3*R*,4*S*,5*R*)-3,4-Dihydroxy-5-methoxy-6,6-dimethyltetrahydro-2*H*-pyran-2-yl)oxy)-3''-fluoro-1,2,3,6-tetrahydro-[1,1':2',1''-terphenyl]-2-yl)acetamide (**22a**). Followed the same noviose coupling procedure as described above for **13a** to afford **22a** as an inseparable mixture of diastereomers.  $^1\text{H}$  NMR (500 MHz,  $\text{CDCl}_3$ )  $\delta$  7.32 (ddd, 1H,  $J = 6.0$ , 7.9, 13.9 Hz), 7.22 (dd, 1H,  $J = 2.8$ , 8.7 Hz), 7.00 (m, 2H), 6.94 (d, 1H,  $J = 7.6$  Hz), 6.87 (m, 1H), 6.77 (dd, 1H,  $J = 2.7$ , 8.7 Hz), 5.59 (m, 1H), 5.52 (m, 1H), 5.49 (d, 1/2H,  $J = 2.4$  Hz), 5.45 (d, 1/2H,  $J = 2.4$  Hz), 4.81 (dd, 1H,  $J = 2.5$ , 8.8 Hz), 4.21 (m, 1H), 4.12 (m, 1H), 4.07 (m, 1H), 3.52 (s, 3H), 3.25 (dd, 1H,  $J = 0.9$ , 9.0 Hz), 2.89 (br s, 1H), 2.76 (m, 1H), 2.67 (s, 1H), 2.26 (m, 2H), 1.69 (m, 1H), 1.65 (s, 3/2H), 1.64 (s, 3/2H), 1.29 (s, 3/2H), 1.28 (s, 3/2H), 1.13 (s, 3/2H), 1.12 (s, 3/2H).  $^{13}\text{C}$  NMR (125 MHz,  $\text{CDCl}_3$ )  $\delta$  169.4, 169.4, 163.7–161.7 (d,  $J = 249.0$  Hz), 154.9, 154.7, 143.0 (dd,  $J = 1.7$ , 8.5 Hz), 142.2 (d,  $J = 1.7$  Hz), 133.4, 133.3, 130.2 (dd,  $J = 1.7$ , 8.5 Hz), 128.4 (d,  $J = 5.0$  Hz), 126.6 (d,  $J = 3.2$  Hz), 125.1 (d,  $J = 3.6$  Hz), 125.0 (m), 117.2, 116.9, 116.6, 116.3 (dd,  $J = 13.4$ , 20.9 Hz), 116.2, 114.3 (dd,  $J = 1.5$ , 20.9 Hz), 98.0, 97.7, 84.5, 84.4, 78.3, 78.3, 77.4, 71.5, 71.4, 68.8, 62.0, 61.9, 49.6, 49.6, 40.5, 40.5, 35.2, 35.1, 33.4, 29.2, 23.6, 23.5, 23.2, 23.1. HRMS (FAB)  $m/z$  [ $M + \text{Na}^+$ ] for  $\text{C}_{28}\text{H}_{34}\text{FNO}_6\text{Na}$ : 522.2262; found, 522.2267.

*N*-(4'-(((3*R*,4*S*,5*R*)-3,4-Dihydroxy-5-methoxy-6,6-dimethyltetrahydro-2*H*-pyran-2-yl)oxy)-3''-(trifluoromethyl)-1,2,3,6-tetrahydro-[1,1':2',1''-terphenyl]-2-yl)acetamide (**22b**). Followed the same noviose coupling procedure as described above for **13a** to afford **22b** as an inseparable mixture of diastereomers.  $^1\text{H}$  NMR (500 MHz,  $\text{CDCl}_3$ )  $\delta$  7.65 (d, 1H,  $J = 8.2$  Hz), 7.56 (t, 1H,  $J = 7.7$  Hz), 7.49 (s, 1H), 7.45 (d, 1H,  $J = 7.6$  Hz), 7.32 (dd, 1H,  $J = 3.0$ , 8.7 Hz), 7.08 (td, 1H,  $J = 2.7$ , 8.6 Hz), 6.85 (dd, 1H,  $J = 2.7$ , 8.5 Hz), 5.65 (m, 1H), 5.59 (m, 1H), 5.57 (d, 1/2H,  $J = 2.4$  Hz), 5.53 (d, 1/2H,  $J = 2.3$  Hz), 4.90 (t, 1H,  $J = 8.2$  Hz), 4.30 (m, 1H), 4.19 (dd, 1H,  $J = 4.3$ , 8.2 Hz), 4.14 (m, 1H), 3.59 (s, 3/2H), 3.59 (s, 3/2H), 3.33 (d, 1H,  $J = 9.0$  Hz), 3.17 (s, 1H), 2.95 (s, 1H), 2.76 (m, 1H), 2.49 (m, 1H), 2.33 (m, 1H), 1.74 (m, 1H), 1.73 (s, 3/2H), 1.72 (s, 3/2H), 1.36 (s, 3/2H), 1.35 (s, 3/2H), 1.21 (s, 3/2H), 1.20 (s, 3/2H).  $^{13}\text{C}$  NMR (100 MHz,  $\text{CDCl}_3$ )  $\delta$  169.4, 169.4, 155.0, 154.8, 142.3, 141.8, 133.4, 133.3, 132.8, 132.6, 131.8 (dq,  $J = 2.2$ , 32.5 Hz), 129.1, 128.6, 126.5, 125.9 (q,  $J = 3.2$ , 7.0 Hz), 125.0, 124.2, 117.6, 117.0, 116.8, 98.1, 97.8, 84.5, 84.4, 78.4, 78.3,

71.3, 71.3, 68.7, 68.7, 61.9, 61.9, 49.4, 49.3, 40.5, 40.5, 35.1, 35.0, 33.1, 29.0, 29.0, 23.4, 23.4, 23.1, 23.0. HRMS (FAB)  $m/z$  [ $M + Na^+$ ] for  $C_{29}H_{34}F_3NO_6Na$ : Calcd, 572.2230; found, 572.2227.

## ■ ASSOCIATED CONTENT

### ■ Supporting Information

$^1H$  and  $^{13}C$  NMR spectra for all new compounds and HPLC traces for most active compounds. This material is available free of charge via the Internet at <http://pubs.acs.org>.

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### Notes

The authors declare no competing financial interest.

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## ■ ABBREVIATIONS USED

DPN, diabetic peripheral neuropathy; DNA, deoxyribonucleic acid; SAR, structure–activity relationships; AGEs, advanced glycation end products enhanced oxidative stress; Hsp90 and Hsp70, heat shock protein 90 and 70

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