

Optimization of Chemical Functionalities of Indole-2-carboxamides To Improve Allosteric Parameters for the Cannabinoid Receptor 1 (CB1)

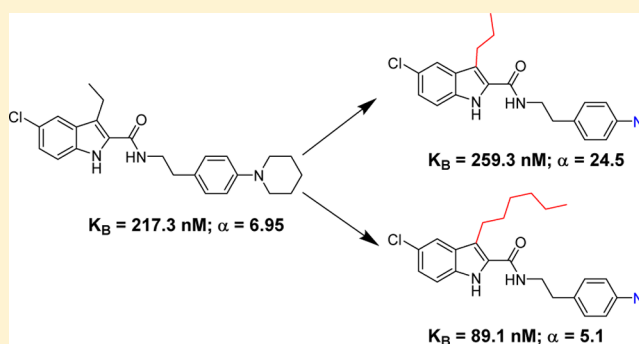
Leepakshi Khurana,[†] Hamed I. Ali,[‡] Teresa Olszewska,[‡] Kwang H. Ahn,[†] Aparna Damaraju,[‡] Debra A. Kendall,^{*,†} and Dai Lu^{*,‡}

[†]Department of Pharmaceutical Sciences, University of Connecticut, Storrs, Connecticut, United States

[‡]Rangel College of Pharmacy, Health Science Center, Texas A&M University, Kingsville, Texas, United States

ABSTRACT: 5-Chloro-3-ethyl-*N*-(4-(piperidin-1-yl)phenethyl)-1*H*-indole-2-carboxamide (**1**; ORG27569) is a prototypical allosteric modulator for the cannabinoid type 1 receptor (CB1). Here, we reveal key structural requirements of indole-2-carboxamides for allosteric modulation of CB1: a critical chain length at the C3-position, an electron withdrawing group at the C5-position, the length of the linker between the amide bond and the phenyl ring B, and the amino substituent on the phenyl ring B. These significantly impact the binding affinity (K_B) and the binding cooperativity (α). A potent CB1 allosteric modulator 5-chloro-*N*-(4-(dimethylamino)phenethyl)-3-propyl-1*H*-indole-2-carboxamide (**12d**) was identified. It exhibited a K_B of 259.3 nM with

a strikingly high binding α of 24.5. We also identified 5-chloro-*N*-(4-(dimethylamino)phenethyl)-3-hexyl-1*H*-indole-2-carboxamide (**12f**) with a K_B of 89.1 nM, which is among the lowest K_B values obtained for any allosteric modulator of CB1. These positive allosteric modulators of orthosteric agonist binding nonetheless antagonized the agonist-induced G-protein coupling to the CB1 receptor, yet induced β -arrestin mediated ERK1/2 phosphorylation.



INTRODUCTION

The cannabinoid type 1 (CB1) receptor is the most abundant G-protein coupled receptor (GPCR) expressed in the central nervous system (CNS), where it attenuates the release of excitatory and inhibitory neurotransmitters.^{1–3} The CB1 receptor is also present in lower concentrations in a variety of peripheral tissues, including, spleen, tonsil, gastrointestinal tract, liver, kidney, and heart.^{4–6} It regulates a variety of physiological functions including neuronal development, neuromodulatory processes, metabolism, nociception, and cardiovascular as well as reproductive functions.^{1,7,8} While CB1 preferentially couples to $G_{i/o}$ type G proteins, it can interact with G_s ⁹ or G_q ¹⁰ under some conditions. The CB1 receptor also modulates the activation of mitogen-activated protein kinases (MAPKs),¹¹ inhibits N- and P/Q-type voltage-gated Ca^{2+} channels, and activates A-type and inwardly rectifying K^+ channels.¹² Moreover, the CB1 receptor can interact with non-G protein partners such as β -arrestins, adaptor protein AP-3, GPCR-associated sorting protein 1 (GASP1), and the adaptor protein FAN to control receptor signaling or trafficking.^{13,14} The complex signaling network of the CB1 receptor suggests the existence of finely controlled modulatory mechanisms of receptor functions.

Traditionally, the functions of the CB1 receptor is regulated through various agonists, partial agonists, antagonists, and

inverse agonists,¹⁵ which bind to the orthosteric site where the endogenous cannabinoids bind. Recently, several allosteric modulators of the CB1 receptor have been identified, which bind to sites that are topologically distinct from the orthosteric binding site. These include 5-chloro-3-ethyl-*N*-(4-(piperidin-1-yl)phenethyl)-1*H*-indole-2-carboxamide (**1**, ORG27569),¹⁶ 1-(4-chlorophenyl)-3-(3-(6-(pyrrolidin-1-yl)pyridin-2-yl)phenyl)urea (PSNCBAM-1),¹⁷ 3-(4-chlorophenyl)-5-(8-methyl-3-*p*-tolyl-8-azabicyclo[3.2.1]octan-2-yl)isoxazole (RTI-371),¹⁸ and the endogenous ligand (*5S,6R,7E,9E,11Z,13E,15S*)-5,6,15-trihydroxyicosanoic acid (lipoxin A4).¹⁹ Allosteric modulators typically work cooperatively with orthosteric ligands and stabilize the receptor in various biological conformations that may be difficult to achieve by the orthosteric ligands.²⁰ This increases the possibility of regulating receptor activities in more sophisticated ways than with orthosteric ligands. Thus, allosteric modulation can significantly expand the pharmacological repertoire for a given receptor.^{21,22} Additionally, allosteric sites are less structurally conserved than the corresponding orthosteric site and thus provide new opportunities for the development of more selective therapeutics.^{8,23}

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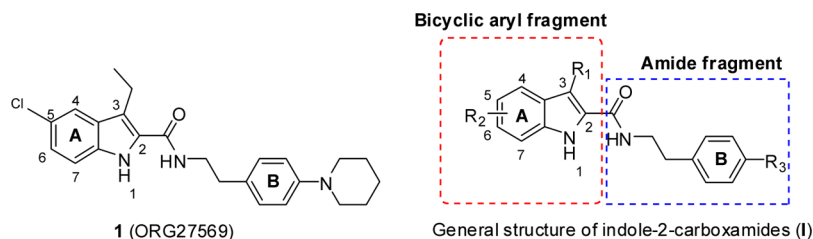


Figure 1. Compound **1** and general structure of indole-2-carboxamides.

The discovery of CB1 allosteric modulators lays the foundation for receptor-selective and signaling-pathway-selective therapies.

Compound **1** was the first allosteric modulator identified for the CB1 receptor.¹⁶ It augments specific binding of the CB1 agonist 2-[(1*R*,2*R*,5*R*)-5-hydroxy-2-(3-hydroxypropyl) cyclohexyl]-5-(2-methyloctan-2-yl)phenol (³H]CP55,940) but decreases the binding of the inverse agonist 5-(4-chlorophenyl)-1-(2,4-dichlorophenyl)-4-methyl-*N*-(piperidin-1-yl)-1*H*-pyrazole-3-carboxamide (³H]SR141716A) in membranes from cells expressing the CB1 receptor.^{16,24} Despite acting as an enhancer of agonist binding, it antagonizes agonist-induced G-protein coupling to the receptor.^{16,24} It was further demonstrated that **1** in the absence of any orthosteric agonist can induce cellular internalization of the CB1 receptor and downstream activation of ERK signaling mediated by β -arrestins^{24–26} as a consequence of CB1 receptor activation. This indicates that allosteric modulators of the CB1 receptor offer the potential to develop drugs capable of generating therapeutic effects via ligand-biased signaling pathways.

Following the discovery of **1**,¹⁶ structure–activity relationship (SAR) studies have revealed that the indole-2-carboxamide scaffold is a viable template for developing CB1 allosteric modulators.^{25,27,28} The general structure of this class of compounds can be divided into two moieties comprising the bicyclic aryl fragment and the amide fragment (Figure 1).

Earlier results from us^{25,28} and others²⁷ have identified several key SARs within this class of compounds (**I**). These include (1) the indole ring of **I** (Figure 1) impacts the ligand's capability to bind to the allosteric site more than the ligand's capability to modulate the orthosteric site;²⁸ (2) the presence of a linear alkyl group at the C3 position of the indole ring is instrumental and its length has a profound influence on the allosteric modulation of the orthosteric binding site;^{25,28} (3) the amide functionality at the C2 position of the indole ring is critical for the allosteric effects on the orthosteric site;²⁷ (4) shortening the linker between the amide bond and the phenyl ring **B** abolished the allosteric modulation of the orthosteric binding site;²⁸ (5) replacing the piperidinyl group of **1** with various functional groups^{27,28} generally led to reduced allosteric modulation except for the methylamino²⁷ and dimethylamino groups.^{27,28} Along with **1**, a few other indole-2-carboxamides^{16,25,27} have also shown allosteric modulation of the CB1 receptor. These molecules, 5-chloro-3-pentyl-*N*-(4-(piperidin-1-yl)phenethyl)-1*H*-indole-2-carboxamide (**2**, ICAM-b),²⁵ *N*-(4-(dimethylamino)phenethyl)-3-ethyl-5-fluoro-1*H*-indole-2-carboxamide (**3**, ORG27759),¹⁶ 5-chloro-*N*-(4-(dimethylamino)phenethyl)-3-ethyl-1*H*-indole-2-carboxamide (**4**),²⁷ and 5-chloro-*N*-(4-(dimethylamino)phenethyl)-3-pentyl-1*H*-indole-2-carboxamide (**5**),²⁸ are shown in Figure 2.

In this context, we expanded our SAR studies of indole-2-carboxamides. Our efforts include elongation of the linker between the amide bond and the phenyl ring **B**, further

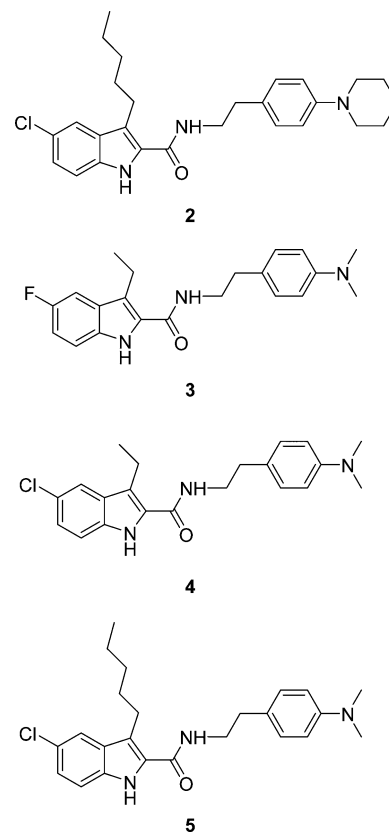
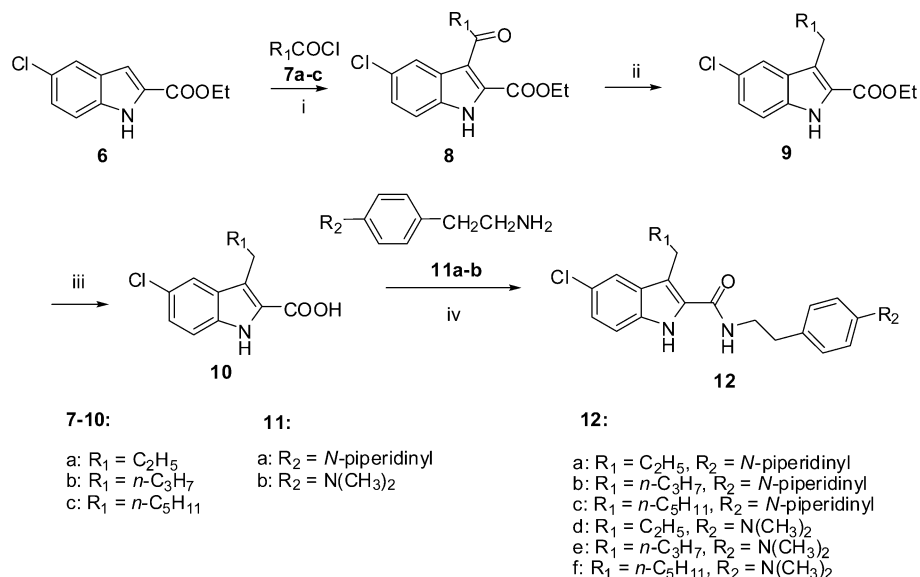
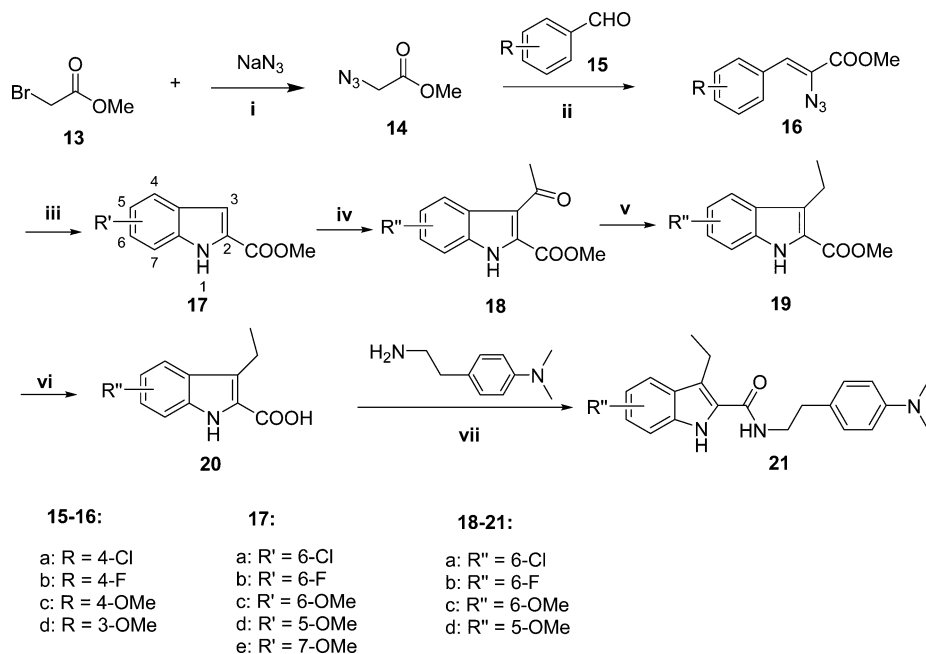


Figure 2. Representative indole-2-carboxamides showing CB1 allosteric.

investigation of the requirement of the C3 alkyl group (i.e., R_1) and modification of the substitutions (i.e., R_2) on the phenyl ring **A** of the bicyclic aryl fragment. To date, most of the reported indole-2-carboxamides showing CB1 allosteric were developed from a 5-chloro-indole-2-carboxamide template^{16,25,27,28} except **3**.¹⁶ Hence, we investigated the impact of different substitutions of the phenyl ring **A** (Figure 1) on the allosteric effects. This effort led to the compounds **21a–d** (Table 2). The allosteric effects of these compounds were evaluated by two essential parameters:^{23,29} the equilibrium dissociation constant (K_B), which reflects the binding affinity of the ligands to the allosteric site, and the binding cooperativity factor (α) that denotes the allosteric interaction between the orthosteric and allosteric ligands when they both occupy the receptor, i.e., it quantifies the direction of and magnitude by which the affinity of one ligand is changed by the other ligand when both are bound to the receptor to form the ternary complex.³⁰ When α is 1.0, the test modulator does not alter orthosteric ligand binding. If α is less than 1.0, the test modulator reduces orthosteric ligand binding (negative allosteric modulation of orthosteric ligand binding). If α is

Scheme 1. Synthesis of 3-Alkyl-5-chloroindole-2-carboxamides 12a–f.^a

^aReagents and conditions: (i) AlCl₃, 1,2-dichloromethane, reflux, 2–3 h; (ii) (Et)₃SiH, CF₃COOH, 0 °C–rt, 4–12 h; (iii) 3N NaOH, EtOH, reflux, 2 h; (iv) BOP, DIPEA, DMF, rt, 4–12 h.

Scheme 2. Synthesis of 3-Ethyl Indole-2-carboxamides 21a–d^a

^aReagents and conditions: (i) NaN₃, DMF, rt, 1.5 h; (ii) NaOCH₃/CH₃OH, –20 °C, 5 h; (iii) xylene, reflux, 3 h; (iv) AlCl₃, 1,2-dichloromethane, reflux, 2.5 h; (v) (Et)₃SiH, CF₃COOH, 0 °C–rt, 4 h; (vi) 3 N NaOH, EtOH, reflux, 2 h; (vii) BOP, DIPEA, DMF, rt, 4–12 h.

greater than 1.0, the modulator increases orthosteric ligand binding (positive allosteric modulation of orthosteric ligand binding).²³ The α and K_B values were analyzed according to the allosteric ternary complex model.¹⁶ Selected allosteric modulators were assessed for their effects on agonist-induced G-protein coupling activity and β -arrestin mediated ERK1/2 phosphorylation.

CHEMISTRY

The syntheses of C3-alkylated indole-2-carboxamides (12a–f) were achieved through the methods illustrated in Scheme 1.

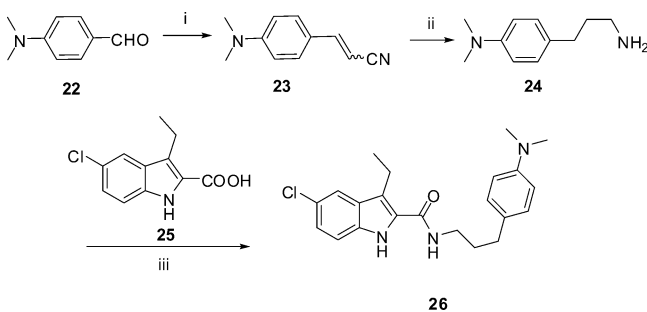
The C3 substituents were introduced through Friedel–Crafts acylation of the ethyl 5-chloroindole-2-carboxylate (6), which is commercially available. Acylation of 6 with various selected acyl chlorides (7a–c) provided the desired 3-acyl-5-chloroindole-2-carboxylates (8a–c). Reduction of their ketone groups by triethylsilane generated the C3 alkylated 5-chloroindole-2-carboxylates (9a–c), which were then hydrolyzed in basic conditions to yield the key intermediate indole-2-carboxylic acids (10a–c). The final compounds (12a–f) were prepared by coupling commercially available amines (11a,b) with the acids (10a–c) individually in the presence of BOP and diisopropyl-

lethyl amine (DIPEA) in anhydrous DMF at room temperature.

To access the indole-2-carboxamides with different substituents on the phenyl ring A (**21a–d**), we employed Hemetsberger–Knittel indole synthesis,³¹ by which the required indole-2-carboxylate **17** can be obtained through Knoevenagel condensation of methyl 2-azidoacetate **14** with the substituted benzaldehyde **15** followed by a thermolysis of the azide of the resultant methyl-2-azidocinnamate **16** and an electrophilic cyclization. The synthesis of indole-2-carboxylate **17** through Hemetsberger–Knittel reaction depends on the reaction conditions, which include the reaction temperature and stoichiometry of reactants in the Knoevenagel condensation (step ii, Scheme 2) and the concentration of reactant in the subsequent thermolytic cyclization (step iii, Scheme 2). We optimized the reaction conditions to obtain **16** and **17** in good yield to proceed with the synthesis (see the discussion in Experimental Section). The cyclization of **16d** provided two regioisomers, the 5- and 7-substituted indole-2-carboxylates (**17d** and **17e**), with the 5-regioisomer (**17d**) being slightly favored over the 7-regioisomer (**17e**). The structures of the two regioisomers were assigned by comparison of their ¹H NMR with the reported data.^{32,33} Acylation of **17a–d** by Friedel–Crafts reaction led to the major product as the desired 3-acyl-indol-2-carboxylates **18a–d**. However, acylation of **17e** generated the 4-acyl-indole-2-carboxamide as the major product, which is not suitable for SAR study in this series; hence, the 4-acylated product was not further pursued in the synthesis of corresponding indole-2-carboxamide **21**. Following the preparation of 3-acylated indole-2-carboxylates **18a–d**, we synthesized the corresponding final products **21a–d** by following Scheme 1 (steps i–iv).

The indole-2-carboxamide **26** was synthesized according to the route illustrated in Scheme 3. It was prepared through

Scheme 3. Synthesis of Indole-2-carboxamide **26**^a



^aReagents and conditions: (i) CH₃CN, KOH, reflux, 10 min; (ii) THF, LiAlH₄, AlCl₃, reflux, 1 h; (iii) BOP, DIPEA, DMF, rt, 4–12 h.

coupling of 4-(3-aminopropyl)-*N,N*-dimethyl aniline **24** with 5-chloro-3-ethyl-1*H*-indole-2-carboxylic acid **25**, which was synthesized according to the reported method.²⁸ For the synthesis of **24**, 4-dimethylaminobenzaldehyde **22** was condensed with acetonitrile through aldol condensation in strong basic conditions to yield a mixture of (*E*)- and (*Z*)-3-(4-(dimethylamino)phenyl)acrylonitrile **23**, which was reduced to yield amine **24**.

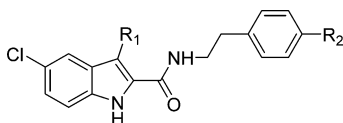
RESULTS AND DISCUSSION

Our previous investigations revealed that structural variation at the C3 position of indole-2-carboxamide has a profound

influence on CB1 allostery. We found that the C3 position prefers a linear alkyl group.²⁸ When the C3 ethyl group of **1** was replaced with a *n*-pentyl group (**2**), it led to the enhancement of the allosteric effect, which is reflected by an improvement in the cooperativity factor α from 6.9 (**1**) to 17.6 (**2**).²⁵ However, further increasing the length of the C3 alkyl to *n*-heptyl (**12g**) and *n*-nonyl (**12h**) groups did not improve the allosteric effects on CB1.²⁸ Here, we elaborated our investigation of the C3 position with variations of the linear alkyl groups such as *n*-propyl, *n*-butyl, and *n*-hexyl groups. The allosteric parameters of the analogues are presented in Table 1. The results reflect that a specific length of the linear alkyl group is required at the C3 position. Increasing the length of the C3 alkyl group of **1** to *n*-propyl (**12a**), *n*-butyl (**12b**), and *n*-pentyl (**2**)²⁵ led to the significant enhancement of binding cooperativity (α). When the length was further elongated to *n*-hexyl (**12c**), *n*-heptyl (**12g**),²⁸ and *n*-nonyl (**12h**),²⁸ the binding cooperativity (α) decreased to a level comparable to **1**. Notably, the significant increase of the binding cooperativity factor (α) of **12a** and **12b** was accompanied by reduced binding affinities to the allosteric site. This reflects that an allosteric modulator can induce a receptor conformation that enhances orthosteric ligand binding despite having a relatively low affinity for the allosteric site. As has been shown previously, the affinity of an allosteric modulator (K_B) and the allostery (α) it exhibits for the orthosteric compound are not necessarily correlated.^{34,35} In this series of modifications (entries 1–7, Table 1), the C3 *n*-propyl provided markedly enhanced allosteric modulation of the orthosteric site (binding cooperativity factor $\alpha = 26.7$). We further assessed the effects of the length of C3 alkyl chain on allosteric properties using **4** as a scaffold because replacing the *N*-piperidinyl group of **1** with a dimethylamino group resulted in improvement of the allosteric effects of indole-2-carboxamides.^{27,28} This effort led to the analogues **12d–f** with improved allosteric parameters. The results suggest that the dimethylamino group on the phenyl ring B is superior to the *N*-piperidinyl group of the indole-2-carboxamides, yielding modulators with higher binding affinity for the allosteric site and greater cooperativity to the orthosteric site (e.g., **12d** vs **12a**, **12e** vs **12b**, **5** vs **2**, and **4** vs **1**). Strikingly, the *n*-hexyl substituent (**12f**) improved the equilibrium dissociation constant (K_B) to 89.1 nM with an α comparable to **1**.

The indole-2-carboxamide **4**^{27,28} was used as a reference compound to vary the substitutions (Table 2) on the phenyl ring A (Figure 1). The C5-chloro group of **4** is an electron withdrawing group (EWG) inductively and is electron donating by resonance. To evaluate the impact of the substitution position, we moved the C5-chloro group of **4** to the C6 position. This modification (**21a**) drastically reduced the binding affinity to the allosteric site ($K_B = 3673$ nM) but not the allosteric modulation of the orthosteric ligand binding ($\alpha = 16.0$). Because a fluoro group has a greater electron-withdrawing inductive effect than a chloro group, we replaced the C6-chloro group with a C6-fluoro group (**21b**), and this modification did not improve the binding affinity to the allosteric site ($K_B = 1580$ nM) while the allosteric modulation on orthosteric ligand binding is well preserved ($\alpha = 22.9$) in comparison with **4** ($\alpha = 19.7$). The result from **21b** along with an earlier result of **3**, which is a 5-fluoro-indole-2-carboxamide,¹⁶ suggested that fluoro as a substituent on ring A is more suboptimal than a chloro group. Taken the fact that chloro group is an EWG inductively and is electron donating

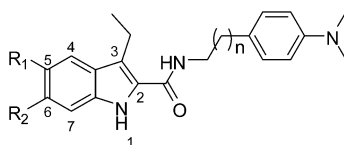
Table 1. Allosterism of Indole-2-carboxamides 12a–f and Some Referenced Compounds



entry	compd	R ¹	R ₂	K _B (nM) ^a	α ^b
1	1 ^c	C ₂ H ₅	N-piperidinyl	217.3 (170.3–277.2)	6.9
2	12a	n-C ₃ H ₇	N-piperidinyl	1746 (377.8–8065)	26.7
3	12b	n-C ₄ H ₉	N-piperidinyl	1985 (775.1–5082)	17.7
4	2 ^d	n-C ₅ H ₁₁	N-piperidinyl	469.9 (126.2–1750)	17.6
5	12c	n-C ₆ H ₁₃	N-piperidinyl	310.6 (110.5–873.2)	4.6
6	12g ^e	n-C ₇ H ₁₅	N-piperidinyl	651.2 (81.51–5203)	7.4
7	12h ^e	n-C ₉ H ₁₉	N-piperidinyl	259.7 (87.56–770)	6.8
8	4 ^e	C ₂ H ₅	N(CH ₃) ₂	207.4 (155.9–2759)	19.7
9	12d	n-C ₃ H ₇	N(CH ₃) ₂	259.3 (19.8–3365)	24.5
10	12e	n-C ₄ H ₉	N(CH ₃) ₂	209.0 (62.7–696.7)	12.8
11	5 ^e	n-C ₅ H ₁₁	N(CH ₃) ₂	167.3 (23.39–1197)	16.5
12	12f	n-C ₆ H ₁₃	N(CH ₃) ₂	89.1 (47.08–168.4)	5.1

^aK_B: equilibrium dissociation constant of a potential allosteric ligand. ^bα: binding cooperativity factor for the tested allosteric modulator. Both parameters were tested using [³H]CP55,940 as the orthosteric ligand. ^cData cited for 1 are from our earlier report²⁴ and are given for comparison. ^dData cited for 2 are from our earlier report²⁵ and are given for comparison. ^eData cited for compounds 4, 5, 12g, and 12h are from our earlier report²⁸ and are given for comparison.

Table 2. Allosterism of Indole-2-carboxamides 21a–d and 26



entry	compd	R ₁	R ₂	n	K _B (nM) ^a	α ^b
13	4 ^c	Cl	H	1	207.4 (155.9–2759)	19.7
14	21a	H	Cl	1	3673 (1048–12880)	16.0
15	21b	H	F	1	1580 (328.7–7599)	22.9
16	21c	OCH ₃	H	1	2708 (973.4–7535)	6.2
17	21d	H	OCH ₃	1	4084 (1213–13750)	11.9
18	26	Cl	H	2	ND ^d	ND ^d

^aK_B: equilibrium dissociation constant of a potential allosteric ligand. ^bα: binding cooperativity factor for the tested allosteric modulator. Both parameters were tested using [³H]CP55,940 as the orthosteric ligand. ^cData cited for 4 are from our earlier report²⁸ and are given for comparison. ^dND: no detectable modulation of [³H]CP55,940 binding using up to 32 μM of test compound.

group (EDG) by resonance, we replaced it with a methoxy group, which is a stronger EDG. This modification led to 21c, which exhibited a significantly decreased cooperativity factor (α = 6.2) and binding affinity (K_B = 2708 nM) in comparison with 4 (α = 19.7, K_B = 207.4 nM). Moving the methoxy group to the C6-position (21d) also reduced the allosteric effect on the orthosteric site and the binding affinity to the allosteric site. This series of compounds (entries 13–17) suggested that the nature and the position of the substituent on the phenyl ring A are critical for both the binding affinity (K_B) to the allosteric site and the binding cooperativity with the orthosteric site.

In line with our earlier finding that the one carbon linker between the amide bond and the phenyl ring B (Figure 1) abolished the allosterism of this class of compounds on CB1 receptor,²⁸ the loss of allosteric modulation of orthosteric agonist CP55,940 binding with 26 indicated the critical role of the 2-carbon linker between the amide bond and the phenyl ring B of the amide fragment within the structure of indole-2-carboxamides (I, Figure 1).

The two robust allosteric modulators 12d and 12f were further tested for their effect on CP55,940-induced G-protein

coupling activity. It was found that both the compounds showed a concentration-dependent inhibition of agonist-induced GTPγS binding as shown in Figure 3.

We previously demonstrated that ERK1/2 can be activated via CB1 in a G-protein-dependent manner by CP55,940 alone and a G-protein-independent manner in the presence of 1 or 2 and CP55,940.^{24,25,36} In agreement, CP55,940-induced ERK1/2 phosphorylation was substantially attenuated by pertussis toxin (PTX) but not by β-arrestin knockdown indicating alone it is G_i-protein mediated (Figure 4A). In contrast, yet in line with the observed inhibition of G protein coupling activity (Figure 3), ERK1/2 phosphorylation due to 12f treatment was PTX insensitive (Figure 4A), suggesting utilization of a β-arrestin mediated pathway. Furthermore, Figure 4B–D show that cotreatment of 2, 12d, or 12f and CP55,940 induce concentration-dependent ERK1/2 phosphorylation that is β-arrestin 1 sensitive; the β-arrestin 1 knockdown resulted in substantial inhibition of ERK1/2 phosphorylation induced by 2, 12d, and 12f. These results indicate that 2, 12d, and 12f are functionally positive allosteric modulators, at least for ERK1/2 phosphorylation. Interestingly, 12d and 12f reached a plateau at

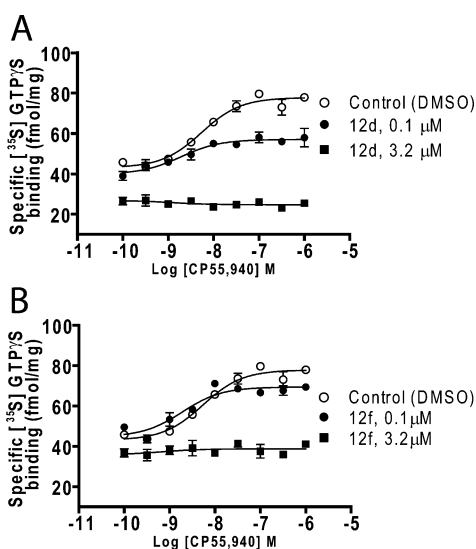


Figure 3. Dose–response curves for CP55,940-induced [³⁵S]GTP γ S binding to HEK293 cell membranes expressing the CB1 receptor in the absence and presence of compounds **12d** (A) and **12f** (B) at the indicated concentrations. Nonspecific binding was determined in the presence of 10 μ M unlabeled GTP γ S. Data is presented as specific binding of GTP γ S (fmol/mg) to the membranes. Each data point represents the mean \pm SE (error bars) of at least three independent experiments performed in duplicate.

5 μ M of the modulator whereas **2** required 10 μ M to achieve comparable levels of ERK1/2 phosphorylation (Figure 4E). This may reflect the higher binding affinity of **12d** and **12f** relative to that of **2** (Table 1).

CONCLUSION

The results from the newly synthesized indole-2-carboxamides along with our earlier findings²⁸ elucidated key structural requirements of indole-2-carboxamides for allosteric modulation of the CB1 receptor. The critical structural factors include: (1) the chain length of the C3-alkyl group is critical with *n*-propyl being preferred for allosteric modulation of orthosteric ligand binding and *n*-hexyl being preferred for enhancing affinity of the allosteric modulator to the CB1 receptor, (2) an electron withdrawing group needs to reside at the C5 position of the indole ring, (3) the linker between the amide bond and the phenyl ring B must be an ethylene group whereas shortening or elongating the linker abolishes allosteric effects, and (4) the substituent on the phenyl ring B explicitly influences both the binding to the allosteric site and the binding cooperativity with the orthosteric ligand, with the *N,N*-dimethyl amino group being preferred over the piperidinyl functionality of the prototypical CB1 allosteric modulator **1**. These SARs will guide the future design and synthesis of more potent CB1 allosteric modulators based on the indole-2-carboxamide scaffold. The therapeutic usefulness of CB1 allosteric modulators is becoming evident. For instance, the CB1 allosteric modulator PSNCBAM-1 exhibits acute hypophagic effects¹⁷ and antagonism of neuronal excitability,³⁷ which have the potential for the treatment of obesity and some CNS disorders. The endogenous CB1 allosteric modulator Lipoxin A4 is capable of protecting neuronal cells from β -amyloid-induced neurotoxicity¹⁹ that has been implied in the etiology of Alzheimer's disease. Given the nature of biased signaling of some CB1 allosteric modulators from the indole-2-carboxamide

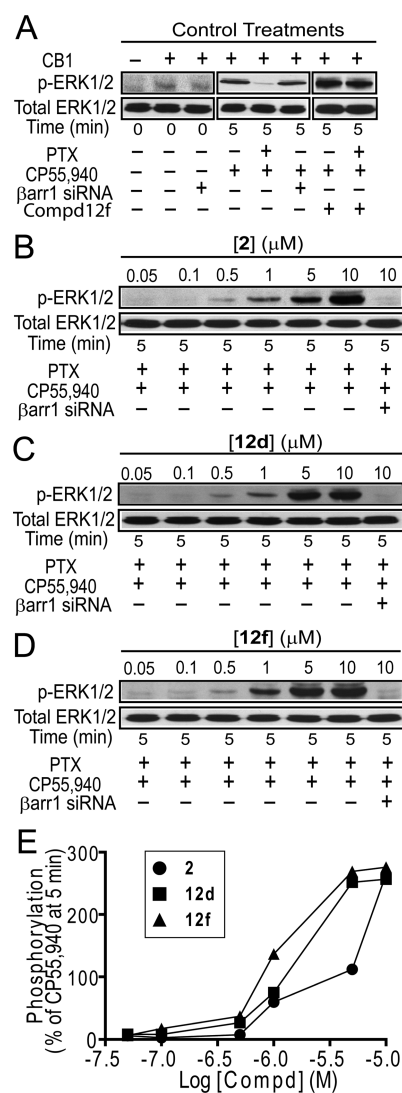


Figure 4. Effect of **12d** and **12f** on ERK1/2 phosphorylation. (A) Mock-transfected and treatment conditions for CP55,940 (0.2 μ M), PTX, and siRNA knockdown of HEK293 cells expressing CB1 are indicated and shown for comparison. (B–D) HEK293 cells expressing CB1 receptors were exposed to 0, 0.05, 0.1, 0.5, 1, 5, and 10 μ M of **2** (B), of **12d** (C), or of **12f** (D) in the presence of 0.2 μ M CP55,940 for 5 min with PTX pretreatment for 16 h. Cell lysates were separated on SDS-PAGE and analyzed by Western blots probed with phospho-ERK1/2 (p-ERK1/2). The total level of ERK1/2 was detected for comparison. Note that the two bands correspond to the predominant isoforms, p42 (ERK2) and p44 (ERK1), for ERK1/2. (E) Graphs provide the quantified ERK1/2 phosphorylation levels induced by each compound for 5 min. Data represent the mean \pm SE and are expressed as a percent of the level of CP55,940-induced ERK1/2 phosphorylation.

class, selective regulation of signaling-pathway-specific functions of the CB1 receptor is possible and this may be therapeutically beneficial. The angiotensin II receptor, for example, may exhibit G-protein dependent or β -arrestin-dependent signaling. Biased agonism of this receptor to promote β -arrestin mediated effects only may provide a beneficial cytoprotective response and obviate a deleterious increase in blood pressure that is observed in a G-protein manner.³⁸ This offers tremendous opportunities for developing drugs for many disorders that have been linked to the CB1 receptor in the CNS and periphery.

EXPERIMENTAL SECTION

Compounds. Tested compounds (12a–f and 21a–d) were synthesized for this study except 1, which was purchased from Tocris Bioscience (Minneapolis, MN). Compounds 2²⁵ and 12g, 12h, and 5 have previously been reported by us²⁸ and were cited in this report for comparison. Compound 4 was first reported by Piscitelli et al.²⁷ and was resynthesized and tested in our laboratories for comparative purposes.

CB1 Expression and Membrane Preparation. HEK 293 cells were maintained in Dulbecco's modified Eagle's medium supplemented with 10% fetal bovine serum and 3.5% mg/mL glucose at 37 °C in 5% CO₂. One day prior to transfection, cells were seeded at approximately 900000 cells/100 mm dishes. The cells were transiently transfected by the calcium phosphate precipitation method.³⁹ At 24 h post transfection, the cells were harvested and washed twice with phosphate buffered saline (PBS). The cells were resuspended in PBS solution containing mammalian protease inhibitor cocktail ((4-2-aminoethyl) benzene-sulfonyl fluoride, pepstatin A, E-64, bestatin, leupeptin, and aprotinin) (Sigma-Aldrich, St. Louis, MO) and lysed by nitrogen cavitation at 750 psi for 5 min using a Parr cell disruption bomb. The lysate was spun at 500g for 10 min at 4 °C, and the supernatant was subsequently spun at 100000g for 45 min at 4 °C. The membrane-containing pellet was resuspended in TME buffer (25 mM Tris-HCl, 5 mM MgCl₂, and 1 mM EDTA, pH 7.4) containing 7% w/v sucrose. For immunoblotting studies, HEK 293 cells were transfected using lipofectamine (Invitrogen, Carlsbad, CA) according to the manufacturer's instructions. Then 24 h post-transfection, cells were washed and incubated for an additional 18 h in serum-free growth media. siRNA transfection was carried out as previously described.³⁶ Briefly, HEK293 cells in a 6-well plate were transfected with the plasmid encoding CB1 and 2.6 μg of siRNA (Qiagen, Valencia, CA) targeting β-arrestin 1 or nonsilencing RNA duplex for control.

Radioligand Binding Assay. Ligand binding assays were performed as previously described to determine the cooperativity between the orthosteric and allosteric ligands.²⁴ Briefly, 6 μg of membrane preparation was incubated for 60 min with a fixed concentration of tracer [³H]CP55940 (141 Ci/mmol, PerkinElmer Life Sciences (Boston, MA)), typically at its K_d, which was determined from a saturation binding isotherm in a total volume of 200 μL of TME buffer containing 0.1% fatty acid-free BSA. Ligand depletion was avoided by adjusting the amount of membrane sample and total assay volume to keep the bound ligand less than 10% of the total. At least nine concentrations of unlabeled test compound (ranging between 100 pM and 100 μM) were used for the binding assays as described previously.²⁴ Nonspecific binding was determined in the presence of unlabeled CP55,940 (1 μM). The reaction was terminated by addition of 250 μL of TME buffer containing 5% BSA followed by filtration with a Brandell cell harvester through Whatman GF/C filter paper followed by washing with ice-cold TME buffer. Radioactivity was measured using liquid scintillation counting.

GTPγS Binding Assay. GTPγS binding assays were performed as described previously.²⁴ Briefly, 7.5 μg of membranes were incubated for 60 min at 30 °C in a total volume of 200 μL of GTPγS binding assay buffer (50 mM Tris-HCl, pH 7.4, 3 mM MgCl₂, 0.2 mM EGTA, and 100 mM NaCl) with unlabeled CP55,940 (at least nine different concentrations were used ranging between 100 pM and 100 μM), 0.1 nM [³⁵S]GTPγS (1250 Ci/mmol; PerkinElmer Life Sciences, Boston, MA), 10 μM GDP (Sigma, St. Louis, MO), and 0.1% (w/v) BSA in the absence and presence of varying concentrations of the allosteric compounds as indicated. Nonspecific binding was determined with 10 μM unlabeled GTPγS (Sigma, St. Louis, MO). The reaction was terminated by rapid filtration through Whatman GF/C filters. The radioactivity trapped in the filters was determined by liquid scintillation counting.

Ligand and GTPγS Binding Data Analysis. All ligand binding assays were carried out in duplicate. Data are presented as the mean ± SE or the mean with the corresponding 95% confidence limits from at least three independent experiments. The interactions between the orthosteric radiolabeled agonist [³H]CP55,940 and the test modu-

lators were analyzed by nonlinear regression using Prism 6.0 (Graphpad Software Inc., San Diego, CA) as previously described.²⁴

Immunoblotting Studies. Cells expressing the CB1 receptor and siRNA targeting β-arrestin 1 or nonsilencing RNA duplex were washed twice with PBS and exposed to varying concentrations of allosteric modulators (2, 12d, or 12f) in the presence of 0.2 μM CP55,940 for 5 min. To observe the effect of the modulators on pertussis toxin (PTX)-insensitive ERK1/2 phosphorylation, cells were treated with 5 ng/mL PTX for 16 h at 37 °C prior to compound treatment. The media were aspirated, and the cells were washed with ice-cold PBS and lysed in ice-cold lysis buffer (150 mM NaCl, 1.0% IGEPAL CA-630, 0.5% sodium deoxycholate, 0.1% SDS, 50 mM Tris, pH 7.5) and a protease inhibitor cocktail (4-(2-aminoethyl)benzenesulfonyl fluoride (AEBSF), pepstatin A, E-64, bestatin, leupeptin, and aprotinin; Sigma, St. Louis, MO). Solubilized cell extracts were centrifuged at 18500g for 15 min at 4 °C, and the supernatant was transferred to a new tube and heated at 95 °C for 3 min. Then 12 μg of total protein was resolved by 10% SDS-PAGE and transferred to polyvinylidene fluoride (PVDF) membrane. After washing with blocking reagent (Fisher Scientific, Pittsburgh, PA), the membrane was incubated for 1h at rt with the primary antibody (1:3000 phospho-p44/42 and p44/42 antibodies; Cell Signaling Technology, Danvers, MA). After washing with PBS, the membrane was incubated with antirabbit peroxidase-conjugated secondary antibody (1:5000; Cell Signaling Technology, Danvers, MA) for 60 min at rt. Immunoreactivity was visualized and quantified as reported earlier.³⁶

Synthesis. All chemical reagents and solvents were purchased from Sigma-Aldrich Chemical Co. unless specified otherwise and used without further purification. All anhydrous reactions were performed under a static argon atmosphere in dried glassware using anhydrous solvents. Organic phases in the work up were dried over anhydrous Na₂SO₄ and removed by evaporation under reduced pressure. The crude compounds were purified by a Combiflash Rf chromatography system (Teledyne Technologies, Inc., Thousand Oaks, CA) unless specified otherwise. Purities of the intermediates were established by thin-layer chromatography (TLC), melting point, ¹H NMR, and mass spectrometry. Analytical thin-layer chromatography (TLC) was run on precoated silica gel TLC aluminum plates (Whatman, UV₂₅₄, layer thickness 250 μm), and the chromatograms were visualized under ultraviolet (UV) light. Melting points were determined on a capillary Electrothermal melting point apparatus and are uncorrected. ¹H NMR spectra of intermediates were recorded on a Bruker Avance DPX-300 spectrometer operating at 300 MHz. ¹H NMR spectra of the final compounds were recorded on a Bruker AV-500 spectrometer operating at 500 MHz. All NMR spectra were recorded using CDCl₃ or DMSO-*d*₆ as solvent unless otherwise stated, and chemical shifts are reported in ppm (parts per million) relative to tetramethylsilane (TMS) as an internal standard. Multiplicities are indicated as br (broadened), s (singlet), d (doublet), t (triplet), q (quartet), m (multiplet), and bs (broadened singlet), and coupling constants (*J*) are reported in hertz (Hz). Low resolution mass spectra were performed at the School of Chemical Sciences, University of Illinois at Urbana–Champaign. The purity of each tested compound was analyzed by combustion elemental analysis and was confirmed to be greater than 99%. (Roberson Microlit laboratories, Madison, NJ).

The Hemesberger–Knittel Reaction. The Hemesberger–Knittel reaction (steps ii and iii, Scheme 2) is postulated to proceed via a highly electrophilic singlet nitrene species, which then inserts into the phenyl ring to form the indole derivatives.^{31,40,41} In some circumstances, the Knoevenagel condensation (step ii, Scheme 2) requires highly excessive quantities of the reactant azidoacetate and the catalytic base (e.g., benzaldehyde:azidoacetate:methoxide = 1:10:10 molar ratio) to achieve high yields.⁴² We tried different stoichiometries of the reactants in the condensation of 14 with 15 and found when the molar ratio of reactants (benzaldehyde:azidoacetate:methoxide) is 1:3:3, it provided an acceptable yield ranging from 58% to 74% for the products 16a–d. When the molar ratio of reactants used was benzaldehyde:azidoacetate:methoxide = 1:10:10, the yields for 16a and 16b were increased from 59% and 58% to 93% (16a) and 69.6% (16b), respectively. Additionally, the yield of the 2-azidocinnamate

from the Knoevenagel condensation also depends on the reaction temperature.⁴³ We found that optimal yields can be obtained when the reaction was first carried out at $-20\text{ }^{\circ}\text{C}$ for 30 min and then at -5 to $0\text{ }^{\circ}\text{C}$ for 6–19 h (depending on the benzaldehyde employed). To obtain the indole-2-carboxylates **17** via thermolytic cyclization of **16** (step iii, Scheme 2), there are many thermolysis conditions reported, which include carrying the reaction with microwave and flow chemistry facilitated thermolytic cyclization^{41,44,45} and employment of various catalysts to facilitate the indole ring formation.^{46,47} We tried various conditions including carrying the thermolytic reaction in different solvents such as regular xylene, anhydrous xylene, anhydrous toluene, and anhydrous THF heated in a pressure tube, as well as using iron(II) triflate to catalyze the reaction in THF.⁴⁶ We found that carrying the reaction in dilute and freshly prepared anhydrous xylene solution (i.e., 1 g of azidocinnamate **16** in ~ 100 mL of xylene, approximately 40 mM) for 30 min generally led to good yield of the products (88.6–94.6%) except for **17b** (53%).

General Procedure A: Preparation of Ethyl 3-Acyl-5-chloro-1H-indole-2-carboxylates (8a–8c). To the solution of ethyl 5-chloro-1H-indole-2-carboxylate (**6**; 10 mmol) in anhydrous 1,2-dichloroethane (25 mL) was added anhydrous aluminum chloride (10 mmol) powder and the selected acyl chloride liquid (**7**; 11.5 mmol) dropwise at room temperature. The reaction mixture was stirred and refluxed under argon atmosphere for 2–3 h. The completion of the reaction was monitored by TLC (25–33% ethyl acetate in hexane). Upon completion, the reaction mixture was poured into ice-cold water (150 mL) then treated with 4N HCl to pH 2. The mixture was extracted with ethyl acetate twice. The combined organic phase was washed with water and brine successively. The organic layer was separated and dried over anhydrous sodium sulfate. Filtration and removal of the solvent in vacuo provided the corresponding crude product, which was purified by Combiflash chromatography (0–40% of ethyl acetate in hexane) which yielded the desired products (**8a–8c**).

General Procedure B: Preparation of Ethyl 3-Alkyl-5-chloro-1H-indole-2-carboxylates (9a–c). Liquid triethylsilane (1.18 mL, 7.40 mmol) was added dropwise to the solution of ethyl 3-acyl-5-chloro-1H-indole-2-carboxylate **8** (3.76 mmol) in trifluoroacetic acid (2.88 mL) at $0\text{ }^{\circ}\text{C}$. The reaction mixture was then gradually warmed up to room temperature and stirred for 4–12 h. The reaction is monitored by TLC (30% of ethyl acetate in hexane). Upon completion of the reaction, the mixture was poured onto ice and treated with saturated sodium carbonate aqueous solution to pH 7 and then extracted with ethyl acetate. The combined organic phase was washed with water and brine and dried over anhydrous sodium sulfate. Filtration and removal of solvent provided the crude product, which was purified by Combiflash silica gel column chromatography (0–30% of ethyl acetate in hexane), which yielded the corresponding products (**9a–c**).

General Procedure C: Preparation of 3-Alkyl-5-chloro-1H-indole-2-carboxylic Acids (10a–c). To the solution of ethyl 5-chloro-3-alkyl-1H-indole-2-carboxylate **9** (1.9 mmol) in anhydrous ethanol (20 mL) was added 3 N sodium hydroxide aqueous solution (3.8 mL, 11.44 mmol). The reaction mixture was stirred and refluxed for 2 h. Upon completion of the hydrolysis monitored by TLC (30% ethyl acetate in hexane), it was cooled to room temperature and treated with 0.1 N HCl to pH 2. The white solid precipitate was filtered. The cake was washed with a minimal amount of cold water and hexane (2 mL) and then dried in a vacuum oven overnight to provide the white solid products (**10a–c**).

General Procedure D: Preparation of 5-Chloroindole-2-carboxamides (12a–f and 21a–d). The solution of an appropriate amine **11** (0.82 mmol) in 1 mL of anhydrous DMF was added to the solution of 5-chloroindole-2-carboxylic acid (**10** or **20**; 0.68 mmol), BOP (502 mg, 1.02 mmol), and *N,N*-diisopropylethylamine (0.71 mL, 4.08 mmol) in anhydrous DMF (4 mL). The reaction mixture was stirred at room temperature for 4–12 h. Upon completion of the coupling reaction, which was monitored by TLC (30% ethyl acetate in hexane), the mixture was poured into cold water (40 mL) and extracted with ethyl acetate twice. The organic layer was separated and

washed with water and brine and dried over anhydrous sodium sulfate. Filtration and removal of solvent in vacuo provided the crude product, which was then purified either by trituration followed by recrystallization in the solvent as specified or by flash chromatography to generate the final compounds (**12a–f**) or (**21a–d**), respectively.

Ethyl 5-Chloro-3-propionyl-1H-indole-2-carboxylate (8a). The title compound was prepared according to the general procedure A by acylation of ethyl 5-chloro-1H-indole-2-carboxylate (**6**, 3.0 g, 13.41 mmol) with propionyl chloride **7a** (1.17 mL, 13.41 mmol). The crude product was purified by Combiflash chromatography (0–20% ethyl acetate in hexane) to yield 1.28 g (34%) of white solid product; mp $120\text{--}122\text{ }^{\circ}\text{C}$. $^1\text{H NMR}$ (300 MHz, chloroform-*d*): δ 9.14 (bs, 1H), 7.99 (s, 1H), 7.40–7.30 (m, 2H), 4.47 (q, $J = 7.3$ Hz, 2H), 3.10 (q, $J = 7.5$ Hz, 2H), 1.45 (t, $J = 7.3$ Hz, 3H), 1.29–1.21 (m, 3H). MS (EI) $m/z = 278.9$ (M^+).

Ethyl 3-Butyryl-5-chloro-1H-indole-2-carboxylate (8b). The title compound was prepared by acylation of ethyl 5-chloro-1H-indole-2-carboxylate **6** (3.0 g, 13.41 mmol) with butyryl chloride **7b** (1.39 mL, 13.41 mmol) according to the general procedure A. The crude product was purified by Combiflash chromatography (0–20% ethyl acetate in hexane), which provided 1.70 g (43.3%) of white solid product; mp $119\text{--}120\text{ }^{\circ}\text{C}$. $^1\text{H NMR}$ (300 MHz, chloroform-*d*): δ 9.22 (bs, 1H), 7.99 (s, 1H), 7.40–7.30 (m, 2H), 4.48 (q, $J = 7.1$ Hz, 2H), 3.06 (t, $J = 7.4$ Hz, 2H), 1.78 (h, $J = 7.4$ Hz, 2H), 1.45 (t, $J = 7.1$ Hz, 3H), 1.00 (t, $J = 7.4$ Hz, 3H). MS (EI) $m/z = 293.2$ (M^+).

Ethyl 5-Chloro-3-hexanoyl-1H-indole-2-carboxylate (8c). The title compound was prepared by acylation of ethyl 5-chloro-1H-indole-2-carboxylate **6** (3.0 g, 13.41 mmol) with hexanoyl chloride **7c** (1.87 mL, 13.41 mmol) according to the general procedure A. The crude product was purified by Combiflash chromatography (0–20% ethyl acetate in hexane), which yielded 1.89 g (43.8%) of white solid product; mp $119\text{--}121\text{ }^{\circ}\text{C}$. $^1\text{H NMR}$ (300 MHz, chloroform-*d*): δ 9.17 (bs, 1H), 7.98 (s, 1H), 7.40–7.29 (m, 2H), 4.48 (q, $J = 7.1$ Hz, 2H), 3.08 (t, $J = 7.4$ Hz, 2H), 1.75 (p, $J = 7.4$ Hz, 2H), 1.45 (t, $J = 7.1$ Hz, 3H), 1.41–1.28 (m, 4H), 0.97–0.86 (m, 3H). MS (EI) $m/z = 321.1$ (M^+).

Ethyl 5-Chloro-3-propyl-1H-indole-2-carboxylate (9a). The title compound was prepared according to general procedure B from ethyl 5-chloro-3-propionyl-1H-indole-2-carboxylate **8a** (1.25 g, 4.47 mmol). The crude product was purified by Combiflash chromatography (0–20% ethyl acetate in hexane), which provided 925 mg (77.9%) of white solid product; mp $142\text{--}144\text{ }^{\circ}\text{C}$. $^1\text{H NMR}$ (300 MHz, chloroform-*d*): δ 8.71 (bs, 1H), 7.66 (s, 1H), 7.34–7.24 (m, 2H), 4.43 (q, $J = 7.1$ Hz, 2H), 3.04 (t, $J = 7.5$ Hz, 2H), 1.71 (h, $J = 7.5$ Hz, 2H), 1.44 (t, $J = 7.1$ Hz, 3H), 0.99 (t, $J = 7.5$ Hz, 3H). MS (EI) $m/z = 265.0$ (M^+).

Ethyl 3-Butyl-5-chloro-1H-indole-2-carboxylate (9b). The title compound was prepared from ethyl 5-chloro-3-butyryl-1H-indole-2-carboxylate **8b** (1.67 g, 5.70 mmol) according to the general procedure B. The crude product was purified by Combiflash chromatography (0–20% ethyl acetate in hexane), which provided 1.34 g (84.0%) of white solid product; mp $122\text{--}125\text{ }^{\circ}\text{C}$. $^1\text{H NMR}$ (300 MHz, chloroform-*d*): δ 8.74 (bs, 1H), 7.66 (s, 1H), 7.35–7.24 (m, 2H), 4.43 (q, $J = 7.2$ Hz, 2H), 3.06 (t, $J = 7.6$ Hz, 2H), 1.67 (p, $J = 7.6$ Hz, 2H), 1.51–1.34 (m, 5H), 0.96 (t, $J = 7.6$ Hz, 3H). MS (EI) $m/z = 279.1$ (M^+).

Ethyl 5-Chloro-3-hexyl-1H-indole-2-carboxylate (9c). The title compound was prepared from ethyl 5-chloro-3-hexanoyl-1H-indole-2-carboxylate **8c** (1.86 g, 5.78 mmol) according to the general procedure B. The crude product was purified by Combiflash chromatography (0–20% ethyl acetate in hexane), which provided 1.58 g (88.8%) of white solid product; mp $103\text{--}105\text{ }^{\circ}\text{C}$. $^1\text{H NMR}$ (300 MHz, chloroform-*d*): δ 9.17 (bs, 1H), 7.98 (s, 1H), 7.40–7.29 (m, 2H), 4.48 (q, $J = 7.1$ Hz, 2H), 3.08 (t, $J = 7.4$ Hz, 2H), 1.75–1.73 (m, 2H), 1.45 (t, $J = 7.1$ Hz, 3H), 1.41–1.28 (m, 6H), 0.97–0.86 (m, 3H). MS (EI) $m/z = 306.9$ (M^+).

5-Chloro-3-propyl-1H-indole-2-carboxylic Acid (10a). The title compound was prepared by hydrolysis of ethyl 5-chloro-3-propyl-1H-indole-2-carboxylate **9a** (900 mg, 3.39 mmol) in ethanolic NaOH solution according to the general procedure C, which provided

625 mg (77.6%) of white solid; mp 166–168 °C. ¹H NMR (300 MHz, DMSO-*d*₆): δ 13.03 (bs, 1H), 11.58 (s, 1H), 7.69 (s, 1H), 7.39 (d, *J* = 8.8 Hz, 1H), 7.21 (d, *J* = 8.8 Hz, 1H), 2.99 (t, *J* = 7.4 Hz, 2H), 1.59 (h, *J* = 7.4 Hz, 2H), 0.89 (t, *J* = 7.4 Hz, 3H). MS (EI) *m/z* = 237.0 (*M*⁺).

3-Butyl-5-chloro-1H-indole-2-carboxylic Acid (10b). The title compound was prepared by hydrolysis of ethyl 5-chloro-3-butyl-1H-indole-2-carboxylate **9b** (1.28 g, 4.58 mmol) in ethanolic NaOH solution according to the general procedure C to provide 1.03 g (89.3%) of white solid product; mp 158–160 °C. ¹H NMR (300 MHz, chloroform-*d*): δ 8.80 (s, 1H), 7.69 (s, 1H), 7.42–7.26 (m, 2H), 3.13 (t, *J* = 7.4 Hz, 2H), 1.69 (p, *J* = 7.4 Hz, 2H), 1.45 (h, *J* = 7.4 Hz, 2H), 0.98 (t, *J* = 7.4 Hz, 3H). MS (EI) *m/z* = 250.9 (*M*⁺).

5-Chloro-3-hexyl-1H-indole-2-carboxylic Acid (10c). The title compound was prepared by hydrolysis of ethyl 5-chloro-3-hexyl-1H-indole-2-carboxylate **9c** (1.55 g, 5.04 mmol) in ethanolic NaOH solution according to the general procedure C to provide 965 mg (68.4%) of white solid product; mp 138–140 °C. ¹H NMR (300 MHz, chloroform-*d*): δ 8.78 (s, 1H), 7.69 (s, 1H), 7.39–7.31 (m, 2H), 3.11 (t, *J* = 7.6 Hz, 2H), 1.70 (p, *J* = 7.2 Hz, 2H), 1.50–1.25 (m, 6H), 0.96–0.83 (m, 3H). MS (EI) *m/z* = 278.9 (*M*⁺).

5-Chloro-*N*-(4-(piperidin-1-yl)phenethyl)-3-propyl-1H-indole-2-carboxamide (12a). The title compound was prepared from 5-chloro-3-propyl-1H-indole-2-carboxylic acid **10a** (216 mg, 0.92 mmol) and 2-(4-piperidin-1-ylphenyl)-ethylamine **11a** (224 mg, 1.1 mmol) according to the general procedure D. The crude product was purified by Combiflash chromatography (30% ethyl acetate in hexane), which yielded 142 mg (36.6%) of white solid product; mp 190–192 °C. ¹H NMR (500 MHz, chloroform-*d*): δ 9.13 (bs, 1H), 7.53 (bs, 1H), 7.29 (d, *J* = 8.6 Hz, 1H), 7.20 (d, *J* = 8.6 Hz, 1H), 7.13 (d, *J* = 8.3 Hz, 2H), 6.91 (d, *J* = 8.3 Hz, 2H), 6.00 (s, 1H), 3.78 (q, *J* = 6.2 Hz, 2H), 3.12 (t, *J* = 5.3 Hz, 4H), 2.87 (t, *J* = 6.2 Hz, 2H), 2.66 (t, *J* = 7.6 Hz, 2H), 1.79–1.65 (m, 4H), 1.59–1.45 (m, 2H), 1.37–1.20 (m, 2H), 0.87 (t, *J* = 6.9 Hz, 3H). MS (EI) *m/z* = 423.2 (*M*⁺). Anal. Calcd for (C₂₃H₃₀ClN₃O): C, 70.82; H, 7.13; N, 9.91. Found: C, 71.09; H, 7.24; N, 9.64.

5-Chloro-3-butyl-*N*-(4-piperidin-1-yl)phenethyl)-1H-indole-2-carboxamide (12b). The title compound was prepared from 5-chloro-3-butyl-1H-indole-2-carboxylic acid **10a** (200 mg, 0.79 mmol) and 2-(4-piperidin-1-ylphenyl)-ethylamine **11a** (194 mg, 0.95 mmol) according to the general procedure D. The crude product was purified by Combiflash chromatography (30% ethyl acetate in hexane), which provided 111 mg (31.9%) of white solid product; mp: 218–220 °C. ¹H NMR (500 MHz, chloroform-*d*): δ 9.19 (bs, 1H), 7.53 (bs, 1H), 7.30 (d, *J* = 8.6 Hz, 1H), 7.20 (d, *J* = 8.6 Hz, 1H), 7.12 (d, *J* = 8.2 Hz, 2H), 6.91 (d, *J* = 8.2 Hz, 2H), 6.02 (bs, 1H), 3.78 (q, *J* = 6.3 Hz, 2H), 3.13 (t, *J* = 5.4 Hz, 4H), 2.87 (t, *J* = 6.3 Hz, 2H), 2.69 (t, *J* = 7.7 Hz, 2H), 1.75–1.66 (m, 2H), 1.60–1.53 (m, 6H), 0.86 (t, *J* = 7.7 Hz, 3H). MS (EI) *m/z* = 437.2 (*M*⁺). Anal. Calcd for (C₂₆H₃₂ClN₃O): C, 70.82; H, 7.13; N, 9.91. Found: C, 71.06; H, 7.42; N, 9.49.

5-Chloro-3-hexyl-*N*-(4-piperidin-1-yl)phenethyl)-1H-indole-2-carboxamide (12c). The title compound was prepared from 5-chloro-3-hexyl-1H-indole-2-carboxylic acid **10c** (200 mg, 0.71 mmol) and 2-(4-piperidin-1-ylphenyl)-ethylamine **11a** (175 mg, 0.86 mmol) according to the general procedure D. The crude product was purified by Combiflash chromatography (0–30% ethyl acetate in hexane), which provided 174 mg (52.6%) of white solid product; mp 160–162 °C. ¹H NMR (500 MHz, chloroform-*d*): δ 9.14 (s, 1H), 7.52 (s, 1H), 7.29 (d, *J* = 8.7 Hz, 1H), 7.20 (dd, *J* = 8.7 Hz, 1.8 Hz, 1H), 7.12 (d, *J* = 8.1 Hz, 2H), 6.71 (d, *J* = 8.1 Hz, 2H), 6.02 (s, 1H), 3.78 (q, *J* = 6.3 Hz, 2H), 2.93 (s, 6H), 2.86 (t, *J* = 6.3 Hz, 2H), 2.69 (t, *J* = 6.9 Hz, 2H), 1.44 (q, *J* = 6.9 Hz, 2H), 1.33–1.13 (m, 6H), 0.87 (t, *J* = 6.9 Hz, 3H). MS (EI) *m/z* = 465.2 (*M*⁺). Anal. Calcd for (C₂₈H₃₆ClN₃O): C, 72.16; H, 7.79; N, 9.02. Found: C, 72.24; H, 7.65; N, 9.01.

5-Chloro-*N*-(4-(dimethylamino)phenethyl)-3-propyl-1H-indole-2-carboxamide (12d). The title compound was prepared from 5-chloro-3-propyl-1H-indole-2-carboxylic acid **10a** (200 mg, 0.84 mmol) and 4-(2-aminoethyl)-*N,N*-dimethyl aniline **11b** (166 mg, 1.01 mmol) according to the general procedure D. The crude product was purified by Combiflash chromatography (30% ethyl acetate in hexane), which yielded 51 mg (15.8%) of white solid

product; mp 112–114 °C. ¹H NMR (500 MHz, chloroform-*d*): δ 9.20 (s, 1H), 7.53 (s, 1H), 7.29 (d, *J* = 8.5 Hz, 1H), 7.20 (d, *J* = 8.5 Hz, 1H), 7.13 (d, *J* = 8.2 Hz, 2H), 6.73 (d, *J* = 8.2 Hz, 2H), 6.02 (s, 1H), 3.77 (q, *J* = 6.3 Hz, 2H), 2.93 (s, 6H), 2.87 (t, *J* = 6.3 Hz, 2H), 2.66 (t, *J* = 7.5 Hz, 2H), 1.49 (h, *J* = 7.5 Hz, 2H), 0.83 (t, *J* = 7.2 Hz, 3H). MS (EI) *m/z* = 384.5 (*M*⁺ + 1). Anal. Calcd for (C₂₂H₂₆ClN₃O): C, 68.83; H, 6.83; N, 10.95. Found: C, 68.56; H, 6.63; N, 10.68.

3-Butyl-5-chloro-*N*-(4-(dimethylamino)phenethyl)-1H-indole-2-carboxamide (12e). The title compound was prepared from 3-butyl-5-chloro-1H-indole-2-carboxylic acid **10b** (200 mg, 0.79 mmol) and 4-(2-aminoethyl)-*N,N*-dimethyl aniline **11b** (156 mg, 0.95 mmol) according to the general procedure D. The crude product was purified by Combiflash chromatography (30% ethyl acetate in hexane), which provided 80 mg (25.0%) of white solid product; mp 198–200 °C. ¹H NMR (500 MHz, chloroform-*d*): δ 9.19 (s, 1H), 7.52 (s, 1H), 7.29 (d, *J* = 8.5 Hz, 1H), 7.20 (d, *J* = 8.5 Hz, 1H), 7.12 (d, *J* = 7.9 Hz, 2H), 6.72 (d, *J* = 8.2 Hz, 2H), 6.03 (s, 1H), 3.78 (q, *J* = 6.3 Hz, 2H), 2.93 (s, 6H), 2.87 (t, *J* = 6.3 Hz, 2H), 2.69 (t, *J* = 7.6 Hz, 2H), 1.43 (p, *J* = 7.5 Hz, 2H), 1.27–1.16 (m, 3H), 0.84 (t, *J* = 7.2 Hz, 3H). MS (EI) *m/z* = 398.5 (*M*⁺ + 1). Anal. Calcd for (C₂₃H₂₈ClN₃O): C, 69.42; H, 7.09; N, 10.56. Found: C, 69.38; H, 7.22; N, 10.28.

5-Chloro-*N*-(4-(dimethylamino)phenethyl)-3-hexyl-1H-indole-2-carboxamide (12f). The title compound was prepared from 5-chloro-3-hexyl-1H-indole-2-carboxylic acid **10c** (200 mg, 0.71 mmol) and 4-(2-aminoethyl)-*N,N*-dimethyl aniline **11b** (140 mg, 0.85 mmol) according to the general procedure D. The crude product was purified by Combiflash chromatography (30% ethyl acetate in hexane), which yielded 150 mg (49.6%) of white solid product; mp 158–160 °C. ¹H NMR (500 MHz, chloroform-*d*): δ 9.14 (s, 1H), 7.52 (s, 1H), 7.29 (d, *J* = 8.7 Hz, 1H), 7.20 (d, *J* = 8.6 Hz, 1H), 7.12 (d, *J* = 8.2 Hz, 2H), 6.71 (d, *J* = 8.1 Hz, 2H), 6.02 (bs, 1H), 3.78 (q, *J* = 6.3 Hz, 2H), 2.93 (s, 6H), 2.86 (t, *J* = 6.6 Hz, 2H), 2.69 (t, *J* = 7.9 Hz, 2H), 1.50–1.40 (m, 2H), 1.33–1.13 (m, 6H), 0.87 (t, *J* = 6.9 Hz, 3H). MS (EI) *m/z* = 426.6 (*M*⁺ + 1). Anal. Calcd for (C₂₅H₃₂ClN₃O): C, 70.49; H, 7.57; N, 9.86. Found: C, 70.38; H, 7.63; N, 9.79.

Methyl 2-Azidoacetate (14). Methylbromoacetate **13** (40 g, 263.0 mmol) was added dropwise to the suspension of sodium azide (19.0 g, 292.3 mmol) in anhydrous DMF (72 mL) at room temperature. The reaction mixture was stirred at room temperature for 2 h. During the reaction, sodium bromide was formed which precipitated out. Water was added until the NaBr solid was dissolved and then the reaction mixture was extracted three times with diethyl ether (3 × 50 mL). The combined organic layers were washed six times with water (total 360 mL) and then separated and washed with brine and dried over magnesium sulfate. Removal of the solvent in vacuo yielded 25 g (83.3%) of colorless oil. ¹H NMR (300 MHz, chloroform-*d*): δ 3.90 (s, 2H), 3.81 (s, 3H). The NMR result is in agreement with published data.⁴¹ Its purity matches with the commercially available methyl-2-azidoacetate (Sigma-Aldrich), therefore **14** was used without further purification.

(*Z*)-Methyl 2-Azido-3-(4-chlorophenyl)acrylate (16a). To the solution of freshly prepared sodium methoxide (480 mg, 8.89 mmol) in anhydrous methanol (12 mL) cooled to –20 °C was added the solution of 4-chlorobenzaldehyde (500 mg, 3.56 mmol) in anhydrous methanol (2.5 mL) in one portion. The temperature was maintained at –20 °C, and liquid methyl 2-azidoacetate (1.02 g, 8.89 mmol) was added dropwise to the reaction mixture during 30 min. The mixture was then allowed to warm to –5–0 °C and stirred at the same temperature until reaction completion (7 h), which was monitored by TLC. Then ice–water was added to the reaction mixture, which led to the precipitation of a white solid, which was filtered off and then washed with ice water and dried in vacuum oven to give 499 mg (59%) of the product **16a**. Alternatively, **16a** can be prepared by condensation of 4-chlorobenzaldehyde **15a** (3 g, 21.34 mmol) and methyl-2-azidoacetate **14** (24.6 g, 213.4 mmol) in similar conditions as above to afford 4.72 g (93%) of white solid product; mp 35–37 °C. ¹H NMR (300 MHz, chloroform-*d*): δ 7.77 (d, *J* = 8.6 Hz, 2H), 7.37 (d, *J* = 8.6 Hz, 2H), 6.78 (s, 1H), 3.93 (s, 3H). MS (EI) *m/z* = 237.1 (*M*⁺).

(*Z*)-Methyl 2-Azido-3-(4-fluorophenyl)acrylate (16b). To a freshly prepared sodium methoxide (432 mg, 8.00 mmol) in

anhydrous methanol (12 mL) cooled to $-20\text{ }^{\circ}\text{C}$ was added 4-fluorobenzaldehyde **15b** (500 mg, 4.03 mmol) in anhydrous methanol (2.5 mL). The temperature was maintained at $-20\text{ }^{\circ}\text{C}$ and liquid methyl 2-azidoacetate **14** (1.40 g, 12.09 mmol) was added dropwise to the reaction mixture during 30 min. The mixture was allowed to warm to -5 to $0\text{ }^{\circ}\text{C}$ and kept at this temperature for 19 h until the reaction was complete. Then ice-water was added to the reaction mixture, which led to the precipitation of a white solid, which was filtered off and then washed with ice-water and dried in vacuum oven to give 516 mg (58%) of the product. Alternatively, **16b** can be prepared by condensation of 4-fluorobenzaldehyde **15b** (5 g, 40.29 mmol) and methyl-2-azidoacetate **14** (46.4 g, 402.9 mmol) in similar conditions as above to yield 6.2 g (69.6%) of **16b**; mp $33\text{--}34\text{ }^{\circ}\text{C}$. $^1\text{H NMR}$ (300 MHz, chloroform-*d*): δ 7.84 (dd, $J = 8.6\text{ Hz, }5.9\text{ Hz, }2\text{H}$), 7.09 (t, $J = 8.6\text{ Hz, }2\text{H}$), 6.89 (s, 1H), 3.93 (s, 3H). MS (EI) $m/z = 221.1\text{ (M}^+)$.

(Z)-Methyl 2-Azido-3-(4-methoxyphenyl)acrylate (16c). To a freshly prepared solution of sodium methoxide (496 mg, 9.18 mmol) in anhydrous methanol (12 mL) cooled to $-20\text{ }^{\circ}\text{C}$ was added 4-methoxybenzaldehyde **15c** (500 mg, 3.67 mmol) in anhydrous methanol (2.5 mL). The temperature was maintained at $-20\text{ }^{\circ}\text{C}$, and liquid methyl 2-azidoacetate **14** (1.06 g, 9.18 mmol) was added dropwise to the reaction mixture during 30 min. The mixture was allowed to warm to -5 to $0\text{ }^{\circ}\text{C}$ and kept at this temperature for 6 h until the reaction was complete. Then ice-water was added to the reaction mixture, which led to the precipitation of a white solid, which was filtered off and then washed with ice-cooled water and dried in vacuum oven to give 637 mg (74%) of white solid product; mp $70\text{--}72\text{ }^{\circ}\text{C}$. $^1\text{H NMR}$ (300 MHz, chloroform-*d*): δ 7.81 (d, $J = 8.9\text{ Hz, }2\text{H}$), 6.93 (d, $J = 9.4\text{ Hz, }3\text{H}$), 3.91 (s, 3H), 3.86 (s, 3H). MS (EI) $m/z = 233.1\text{ (M}^+)$. Alternatively, **16c** can be prepared by condensation of 4-methoxybenzaldehyde **15c** (5 g, 36.72 mmol) and methyl-2-azidoacetate **7** (21.1 g, 183.6 mmol) in the solution of sodium methoxide (322.29 mmol) in anhydrous methanol (130 mL) in similar conditions as above to yield 4.3 g (50.2%) of white solid.

(Z)-Methyl 2-Azido-3-(3-methoxyphenyl)acrylate (16d). To a freshly prepared sodium methoxide (417 mg, 7.71 mmol) solution in anhydrous methanol (12 mL) cooled to $-20\text{ }^{\circ}\text{C}$ was added 3-methoxybenzaldehyde **15d** (500 mg, 3.67 mmol) in anhydrous methanol (2.5 mL). The temperature was maintained at $-20\text{ }^{\circ}\text{C}$, and liquid methyl 2-azidoacetate **14** (846 mg, 7.34 mmol) was added dropwise to the reaction mixture during 30 min. The mixture was allowed to warm to $-10\text{ }^{\circ}\text{C}$ and stirred at the same temperature for 5 h until the reaction was complete. Then ice-cooled water was added to the reaction mixture, which led to the precipitation of a white solid, which was filtered off and then washed with ice-cooled water and dried in a vacuum oven to give 580 mg (68%) of white solid product; mp $163\text{--}165\text{ }^{\circ}\text{C}$. $^1\text{H NMR}$ (300 MHz, chloroform-*d*): δ 7.77 (d, $J = 8.6\text{ Hz, }2\text{H}$), 7.37 (d, $J = 8.6\text{ Hz, }2\text{H}$), 6.78 (s, 1H), 3.93 (s, 3H). MS (EI) $m/z = 233.1\text{ (M}^+)$.

Methyl 6-Chloro-1H-indole-2-carboxylate (17a). The solution of (*Z*)-methyl 2-azido-3-(4-chlorophenyl)acrylate **16a** (1.1 g, 4.63 mmol) in freshly prepared anhydrous xylene (110 mL) was bubbled through with argon flow for 3 min. The reaction mixture was refluxed for 30 min in argon atmosphere and then cooled to room temperature. The solvent was evaporated in vacuo, and the residue was purified by Combiflash chromatography (0–20% ethyl acetate in hexane) to yield 858 mg (88.6%) of a white solid product; mp $178\text{--}180\text{ }^{\circ}\text{C}$. $^1\text{H NMR}$ (300 MHz, chloroform-*d*): δ 9.02 (bs, 1H), 7.62 (d, $J = 8.6\text{ Hz, }1\text{H}$), 7.44 (s, 1H), 7.21 (s, 1H), 7.15 (d, $J = 8.6\text{ Hz, }1\text{H}$), 3.92 (s, 3H). MS (EI) $m/z = 209.0\text{ (M}^+)$.

Methyl 6-Fluoro-1H-indole-2-carboxylate (17b). The title compound was prepared from (*Z*)-methyl 2-azido-3-(4-fluorophenyl)acrylate **16b** (500 mg, 2.26 mmol) in anhydrous xylene (60 mL) according to the procedure described for **17a**. The crude product was purified by Combiflash chromatography (0–20% ethyl acetate in hexane) to yield 231 mg (53%) of white solid product; mp $149\text{--}151\text{ }^{\circ}\text{C}$. $^1\text{H NMR}$ (300 MHz, chloroform-*d*): δ 9.01 (bs, 1H), 7.64 (dd, $J = 8.6\text{ Hz, }5.5\text{ Hz, }1\text{H}$), 7.22 (s, 1H), 7.11 (d, $J = 9.3\text{ Hz, }1\text{H}$), 6.95 (t, $J = 9.3\text{ Hz, }1\text{H}$), 3.96 (s, 3H). MS (EI) $m/z = 193.0\text{ (M}^+)$.

Methyl 6-Methoxy-1H-indole-2-carboxylate (17c). The title compound was prepared from (*Z*)-methyl 2-azido-3-(4-methoxyphenyl)acrylate **16c** (500 mg, 3.67 mmol) in anhydrous xylene (55 mL) according to the procedure described for **17a**. The crude product was purified by Combiflash chromatography (0–20% ethyl acetate in hexane) to give 390 mg (87%) of a white solid product; mp $110\text{--}112\text{ }^{\circ}\text{C}$. $^1\text{H NMR}$ (300 MHz, chloroform-*d*): δ 8.91 (bs, 1H), 7.57 (d, $J = 9.5\text{ Hz, }1\text{H}$), 7.18 (s, 1H), 6.89–6.81 (m, 2H), 3.95 (s, 3H), 3.87 (s, 3H). MS (EI) $m/z = 205.1\text{ (M}^+)$.

Methyl 5-Methoxy-1H-indole-2-carboxylate (17d) and Methyl 7-Methoxy-1H-indole-2-carboxylate (17e). The solution of (*Z*)-methyl 2-azido-3-(3-methoxyphenyl)acrylate **16d** (2.7 g, 4.63 mmol) in freshly prepared anhydrous xylene (110 mL) was bubbled through with argon flow for 3 min. The reaction mixture was refluxed for 30 min in argon atmosphere and then cooled to room temperature. The solvent was evaporated in vacuo, and the residue was purified by Combiflash chromatography (0–20% ethyl acetate in hexane) to give 1.3 g (55%) of **17d** as a white solid product and 940 mg (39.6%) of **17e** as a white solid product. **17d**: mp $175\text{--}177\text{ }^{\circ}\text{C}$. $^1\text{H NMR}$ (300 MHz, chloroform-*d*) δ 8.91 (s, 1H), 7.33 (d, $J = 8.9\text{ Hz, }1\text{H}$), 7.16 (s, 1H), 7.10 (s, 1H), 7.02 (d, $J = 9.8\text{ Hz, }1\text{H}$), 3.96 (s, 3H), 3.87 (s, 3H). MS (EI) $m/z = 205.0\text{ (M}^+)$. The $^1\text{H NMR}$ chemical shifts of **17d** are in agreement with the published results.³² **17e**: mp $115\text{--}117\text{ }^{\circ}\text{C}$. $^1\text{H NMR}$ (300 MHz, chloroform-*d*): δ 9.11 (s, 1H), 7.30 (d, $J = 8.3\text{ Hz, }1\text{H}$), 7.22 (bs, 1H), 7.08 (t, $J = 7.9\text{ Hz, }1\text{H}$), 6.74 (d, $J = 7.7\text{ Hz, }1\text{H}$), 3.99 (s, 3H), 3.96 (s, 3H). MS (EI) $m/z = 205.0\text{ (M}^+)$. The $^1\text{H NMR}$ chemical shifts of **17e** are also in agreement with the published data.³³

Methyl 3-Acetyl-6-chloro-1H-indole-2-carboxylate (18a). The title compound was prepared by acylation of methyl 6-chloro-1H-indole-2-carboxylate **17a** (1.7 g, 8.11 mmol) with acetyl chloride (0.72 mL, 10.13 mmol) according to the general procedure A. The crude product was purified by Combiflash chromatography (0–20% ethyl acetate in hexane), which provided 1.2 g (58.8%) of yellow solid product; mp $220\text{--}222\text{ }^{\circ}\text{C}$. $^1\text{H NMR}$ (300 MHz, chloroform-*d*): δ 12.61 (s, 1H), 7.94 (d, $J = 8.7\text{ Hz, }1\text{H}$), 7.52 (s, 1H), 7.23 (d, $J = 8.7\text{ Hz, }1\text{H}$), 3.95 (s, 2H), 2.96 (s, 3H). MS (EI) $m/z = 251.1\text{ (M}^+)$.

Methyl 3-Acetyl-6-fluoro-1H-indole-2-carboxylate (18b). The title compound was prepared by acylation of methyl 6-fluoro-1H-indole-2-carboxylate **17b** (1.5 g, 7.77 mmol) with acetyl chloride (0.72 mL, 10.09 mmol) according to the general procedure A. The crude product was purified by Combiflash chromatography (0–20% ethyl acetate in hexane), which provided 846 mg (46.3%) of white solid product; mp $198\text{--}200\text{ }^{\circ}\text{C}$. $^1\text{H NMR}$ (300 MHz, DMSO-*d*₆): δ 12.54 (s, 1H), 7.95 (dd, $J = 8.8\text{ Hz, }5.7\text{ Hz, }1\text{H}$), 7.23 (d, $J = 9.3\text{ Hz, }1\text{H}$), 7.09 (t, $J = 9.3\text{ Hz, }1\text{H}$), 3.94 (s, 3H), 2.60 (s, 3H). MS (EI) $m/z = 235.1\text{ (M}^+)$.

Methyl 3-Acetyl-6-methoxy-1H-indole-2-carboxylate (18c). The title compound was prepared by acylation of methyl 6-methoxy-1H-indole-2-carboxylate **17c** (1.5 g, 7.31 mmol) with acetyl chloride (0.67 mL, 9.50 mmol) according to the general procedure A. The crude product was purified by Combiflash chromatography (0–20% ethyl acetate in hexane) to provide 1.05 g (58.1%) of white solid product; mp $128\text{--}130\text{ }^{\circ}\text{C}$. $^1\text{H NMR}$ (300 MHz, chloroform-*d*): δ 11.03 (bs, 1H), 7.84 (d, $J = 8.8\text{ Hz, }1\text{H}$), 7.20 (d, $J = 2.3\text{ Hz, }1\text{H}$), 6.92 (d, $J = 8.8\text{ Hz, }1\text{H}$), 4.04 (s, 3H), 3.95 (s, 3H), 2.74 (s, 3H). MS (EI) $m/z = 247.1\text{ (M}^+)$.

Methyl 3-Acetyl-5-methoxy-1H-indole-2-carboxylate (18d). The title compound was prepared by acylation of methyl 5-methoxy-1H-indole-2-carboxylate **17d** (1.2 g, 5.84 mmol) with acetyl chloride (0.44 mL, 6.13 mmol) according to the general procedure A. The crude product was purified by Combiflash chromatography (0–20% ethyl acetate in hexane), which provided 1.0 g (69.3%) of white solid product; mp $187\text{--}189\text{ }^{\circ}\text{C}$. $^1\text{H NMR}$ (300 MHz, acetone-*d*₆): δ 11.33 (bs, 1H), 7.58 (d, $J = 2.4\text{ Hz, }1\text{H}$), 7.48 (d, $J = 9.0\text{ Hz, }1\text{H}$), 7.01 (dd, $J = 9.0\text{ Hz, }2.4\text{ Hz, }1\text{H}$), 3.98 (s, 3H), 3.84 (s, 3H), 2.66 (s, 3H). MS (EI) $m/z = 247.0\text{ (M}^+)$.

Methyl 3-Ethyl-6-chloro-1H-indole-2-carboxylate (19a). The title compound was prepared by reduction of methyl 3-acetyl-6-chloro-1H-indole-2-carboxylate **18a** (500 mg, 1.99 mmol) with triethylsilane (0.57 mL, 4.96 mmol) in trifluoroacetic acid (1.52 mL) according to

the general procedure B. The crude product was purified by Combiflash chromatography (0–30% ethyl acetate in hexane) to yield 460 mg (97.3%) of white solid product; mp 163–165 °C. ¹H NMR (300 MHz, chloroform-*d*): δ 8.72 (bs, 1H), 7.62 (d, *J* = 8.6 Hz, 1H), 7.38 (s, 1H), 7.12 (d, *J* = 8.6 Hz, 1H), 3.98 (s, 3H), 3.11 (q, *J* = 7.5 Hz, 2H), 1.28 (t, *J* = 7.5 Hz, 3H). MS (EI) *m/z* = 237.0 (M⁺).

Methyl 3-Ethyl-6-fluoro-1H-indole-2-carboxylate (19b). The title compound was prepared by reduction of methyl 3-acetyl-6-fluoro-1H-indole-2-carboxylate **18b** (500 mg, 2.13 mmol) with triethylsilane (0.84 mL, 5.31 mmol) in trifluoroacetic acid (2.42 mL) according to the general procedure B. The crude product was purified by Combiflash chromatography (0–30% ethyl acetate in hexane) to yield 446 mg (94.6%) of white solid product; mp 131–133 °C. ¹H NMR (300 MHz, DMSO-*d*₆): δ 8.75 (s, 1H), 7.63 (dd, *J* = 8.3 Hz, 5.5 Hz, 1H), 7.05 (d, *J* = 9.2 Hz, 1H), 6.93 (t, *J* = 9.2 Hz, 1H), 3.97 (s, 3H), 3.11 (q, *J* = 7.5 Hz, 2H), 1.29 (t, *J* = 7.5 Hz, 3H). MS (EI) *m/z* = 221.0 (M⁺).

Methyl 3-Ethyl-6-methoxy-1H-indole-2-carboxylate (19c). The title compound was prepared by reduction of methyl 3-acetyl-6-methoxy-1H-indole-2-carboxylate **18c** (1.0 g, 4.04 mmol) with triethylsilane (1.29 mL, 8.08 mmol) in trifluoroacetic acid (3 mL) according to the general procedure B. The crude product was purified by Combiflash chromatography (0–30% ethyl acetate in hexane), which provided 900 mg (95.5%) of white solid product; mp 150–152 °C. ¹H NMR (300 MHz, chloroform-*d*): δ 8.66 (s, 1H), 7.50 (d, *J* = 8.8 Hz, 1H), 7.19 (s, 1H), 6.91 (d, *J* = 8.8 Hz, 1H), 3.95 (s, 3H), 3.92 (s, 3H), 2.87 (q, *J* = 7.5 Hz, 2H), 1.28 (t, *J* = 7.5 Hz, 3H). MS (EI) *m/z* = 233.1 (M⁺).

Methyl 3-Ethyl-5-methoxy-1H-indole-2-carboxylate (19d). The title compound was prepared by reduction of methyl 3-acetyl-5-methoxy-1H-indole-2-carboxylate **18d** (600 mg, 2.43 mmol) with triethylsilane (0.95 mL, 6.06 mmol) in trifluoroacetic acid (1.86 mL) according to the general procedure B. The crude product was purified by Combiflash chromatography (0–30% ethyl acetate in hexane), which yielded 450 mg (79.4%) of white solid product; mp 125–127 °C. ¹H NMR (300 MHz, chloroform-*d*): δ 8.66 (s, 1H), 7.27 (d, *J* = 8.7, 1H), 7.08 (d, *J* = 8.2 Hz, 1H), 7.02 (d, *J* = 8.5 Hz, 1H), 3.96 (s, 3H), 3.89 (s, 3H), 3.11 (q, *J* = 7.6 Hz, 2H), 1.28 (t, *J* = 7.6 Hz, 3H). MS (EI) *m/z* = 233.1 (M⁺).

3-Ethyl-6-chloro-1H-indole-2-carboxylic Acid (20a). The title compound was prepared by hydrolysis of methyl 3-ethyl-6-chloro-1H-indole-2-carboxylate **19a** (400 mg, 1.68 mmol) according to the general procedure C to yield 300 mg (79.8%) of white solid product; mp 193–195 °C. ¹H NMR (300 MHz, DMSO-*d*₆): δ 13.03 (bs, 1H), 11.50 (s, 1H), 7.67 (d, *J* = 8.6 Hz, 1H), 7.39 (s, 1H), 7.05 (d, *J* = 8.6 Hz, 1H), 3.03 (q, *J* = 7.4 Hz, 2H), 1.17 (t, *J* = 7.4 Hz, 3H). MS (EI) *m/z* = 223.1 (M⁺).

3-Ethyl-6-fluoro-1H-indole-2-carboxylic Acid (20b). The title compound was prepared by hydrolysis of methyl 3-ethyl-6-fluoro-1H-indole-2-carboxylate **19b** (400 mg, 1.81 mmol) according to the general procedure C to yield 300 mg (80.0%) of white solid product; mp 192–194 °C. ¹H NMR (300 MHz, DMSO-*d*₆): δ 12.92 (s, 1H), 11.44 (s, 1H), 7.67 (dd, *J* = 8.6 Hz, 5.7 Hz, 1H), 6.91 (d, *J* = 9.9 Hz, 1H), 6.91 (t, *J* = 9.3 Hz, 1H), 3.03 (q, *J* = 7.4 Hz, 2H), 1.17 (t, *J* = 7.4 Hz, 3H). MS (EI) *m/z* = 207.1 (M⁺).

3-Ethyl-6-methoxy-1H-indole-2-carboxylic Acid (20c). The title compound was prepared by hydrolysis of methyl 3-ethyl-6-methoxy-1H-indole-2-carboxylate **19c** (400 mg, 1.71 mmol) according to the general procedure C to provide 300 mg (80.0%) of white solid product; mp 193–195 °C. ¹H NMR (300 MHz, chloroform-*d*): δ 8.71 (s, 1H), 7.54 (d, *J* = 8.8 Hz, 1H), 7.36 (s, 1H), 6.94 (d, *J* = 8.8 Hz, 1H), 3.93 (s, 3H), 2.84 (q, *J* = 7.5 Hz, 2H), 1.27 (t, *J* = 7.5 Hz, 3H). MS (EI) *m/z* = 219.1 (M⁺).

3-Ethyl-5-methoxy-1H-indole-2-carboxylic Acid (20d). The title compound was prepared by hydrolysis of methyl 3-ethyl-5-methoxy-1H-indole-2-carboxylate **19d** (389 mg, 1.68 mmol) according to the general procedure C to provide 318 mg (86.3%) of white solid product; mp 193–195 °C. ¹H NMR (300 MHz, acetone-*d*₆): δ 10.38 (s, 1H), 7.40 (d, *J* = 8.8 Hz, 1H), 7.15 (s, 1H), 6.96 (d, *J* = 8.8 Hz,

1H), 3.85 (s, 1H), 3.14 (q, *J* = 7.4 Hz, 2H), 1.17 (t, *J* = 7.4 Hz, 3H). MS (EI) *m/z* = 219.0 (M⁺).

N-(4-(Dimethylamino)phenethyl)-3-ethyl-6-chloro-1H-indole-2-carboxamide (21a). The title compound was prepared from 3-ethyl-6-chloro-1H-indole-2-carboxylic acid **20a** (200 mg, 0.89 mmol) and 4-(2-aminoethyl)-*N,N*-dimethylaniline **11b** (176 mg, 1.07 mmol) according to the general procedure D. The crude product was purified by Combiflash chromatography (0–20% ethyl acetate in hexane), which yielded 130 mg (39.5%) of white solid product; mp 166–168 °C. ¹H NMR (500 MHz, chloroform-*d*): δ 9.29 (bs, 1H), 7.48 (d, *J* = 8.6 Hz, 1H), 7.37 (d, *J* = 1.8 Hz, 1H), 7.16–7.12 (m, 2H), 7.06 (dd, *J* = 8.6 Hz, 1.8 Hz, 1H), 6.73 (d, *J* = 8.7 Hz, 2H), 6.00 (bs, 1H), 3.78 (q, *J* = 6.4 Hz, 2H), 2.93 (s, 6H), 2.88 (t, *J* = 6.4 Hz, 2H), 2.73 (q, *J* = 7.7 Hz, 2H), 1.09 (t, *J* = 7.7 Hz, 3H). MS (EI) *m/z* = 369.2 (M⁺). Anal. Calcd for (C₂₁H₂₄ClN₃O): C, 68.19; H, 6.54; N, 11.36. Found: C, 68.29; H, 6.60; N, 11.26.

N-(4-(Dimethylamino)phenethyl)-3-ethyl-6-fluoro-1H-indole-2-carboxamide (21b). The title compound was prepared from 3-ethyl-6-fluoro-1H-indole-2-carboxylic acid **20b** (200 mg, 0.97 mmol) and 4-(2-aminoethyl)-*N,N*-dimethylaniline **11b** (190 mg, 1.16 mmol) according to the general procedure D. The crude product was purified by Combiflash chromatography (0–20% ethyl acetate in hexane), which yielded 136 mg (39.7%) of white solid product; mp 160–162 °C. ¹H NMR (500 MHz, chloroform-*d*): δ 9.20 (bs, 1H), 7.49 (dd, *J* = 8.8 Hz, 5.3 Hz, 1H), 7.13 (d, *J* = 8.5 Hz, 2H), 7.04 (dd, *J* = 9.5 Hz, 2.2 Hz, 1H), 6.87 (td, *J* = 9.1 Hz, 2.2 Hz, 1H), 6.73 (d, *J* = 8.7 Hz, 2H), 5.96 (bs, 1H), 3.77 (q, *J* = 6.6 Hz, 2H), 2.93 (s, 6H), 2.88 (t, *J* = 6.6 Hz, 2H), 2.74 (q, *J* = 7.7 Hz, 2H), 1.10 (t, *J* = 7.7 Hz, 3H). MS (EI) *m/z* = 353.2 (M⁺). Anal. Calcd for (C₂₁H₂₄FN₃O): C, 71.36; H, 6.84; N, 11.89. Found: C, 71.45; H, 6.95; N, 11.62.

N-(4-(Dimethylamino)phenethyl)-3-ethyl-6-methoxy-1H-indole-2-carboxamide (21c). The title compound was prepared from 3-ethyl-6-methoxy-1H-indole-2-carboxylic acid **20c** (200 mg, 0.91 mmol) and 4-(2-aminoethyl)-*N,N*-dimethylaniline **11b** (180 mg, 1.10 mmol) according to the general procedure D. The crude product was purified by Combiflash chromatography (0–30% ethyl acetate in hexane), which provided 131 mg (39.4%) of white solid product; mp 158–161 °C. ¹H NMR (500 MHz, chloroform-*d*): δ 8.99 (bs, 1H), 7.42 (d, *J* = 8.8 Hz, 1H), 7.13 (d, *J* = 8.7 Hz, 2H), 6.88 (d, *J* = 8.8 Hz, 1H), 6.75 (d, *J* = 8.7 Hz, 2H), 6.66 (d, *J* = 2.4 Hz, 1H), 6.14 (bs, 1H), 3.90 (s, 3H), 3.70 (q, *J* = 6.8 Hz, 2H), 2.95 (bs, 6H), 2.87–2.81 (m, 4H), 1.23 (t, *J* = 7.6 Hz, 3H). MS (EI) *m/z* = 365.3 (M⁺). Anal. Calcd for (C₂₂H₂₇N₃O₂·¹/₁₀H₂O): C, 71.95; H, 7.46; N, 11.44. Found: C, 71.88; H, 7.63; N, 11.09.

N-(4-(Dimethylamino)phenethyl)-3-ethyl-5-methoxy-1H-indole-2-carboxamide (21d). The title compound was prepared from 3-ethyl-5-methoxy-1H-indole-2-carboxylic acid **20d** (224 mg, 1.02 mmol) and 4-(2-aminoethyl)-*N,N*-dimethylaniline **11b** (201 mg, 1.22 mmol) according to the general procedure D. The crude product was purified by Combiflash chromatography (0–30% ethyl acetate in hexane), which yielded 125 mg (33.5%) of white solid product; mp 183–185 °C. ¹H NMR (500 MHz, chloroform-*d*): δ 9.01 (bs, 1H), 7.28 (d, *J* = 10.0 Hz, 1H), 7.14 (d, *J* = 7.3 Hz, 2H), 6.97 (s, 1H), 6.95 (d, *J* = 10.0 Hz, 1H), 6.74 (d, *J* = 7.3 Hz, 2H), 6.01 (bs, 1H), 3.86 (s, 3H), 3.87 (q, *J* = 6.6 Hz, 2H), 2.94 (s, 6H), 2.88 (t, *J* = 6.6 Hz, 2H), 2.75 (q, *J* = 7.6 Hz, 2H), 1.12 (t, *J* = 7.6 Hz, 3H). MS (EI) *m/z* = 365.2 (M⁺). Anal. Calcd for (C₂₂H₂₇N₃O₂): C, 72.30; H, 7.45; N, 11.50. Found: C, 72.04; H, 7.20; N, 11.42.

(E)- and (Z)-3-(4-(Dimethylamino)phenyl)acrylonitrile (23a and 23b). The title compound was prepared according to a reported procedure with modifications.⁴⁸ A 100 mL 3-necked round bottomed flask equipped with a pressure equalization addition funnel and a reflux condenser was charged with powdered KOH (2.26 g, 40.22 mmol) and anhydrous acetonitrile (50 mL) under argon atmosphere. The mixture was heated to reflux and the solution of 4-dimethylamino-benzaldehyde (5.0 g, 33.5 mmol) was added to 20 mL of acetonitrile in a stream. After the addition, stirring was continued for 10 min and the hot solution was poured onto cracked ice. The mixture was extracted three times with CH₂Cl₂ (3 × 75 mL). The organic layer was separated and washed with brine and dried over anhydrous Na₂SO₄.

After filtration, the filtrate was condensed in vacuo in the water bath maintained at 30 °C to minimize decomposition. The crude product was purified by Combiflash chromatography (0–10% ethyl acetate in hexane), which yielded 3.15 g (54.8%) of the mixture of the *E*- and *Z*-isomers as light-yellow powder. The mixture gives an *E/Z* ratio of 4:1; mp 149–151 °C. ¹H NMR (300 MHz, chloroform-*d*) *Z*-isomer: δ 7.76 (d, *J* = 8.9 Hz, 2H), 6.96 (d, *J* = 12.1 Hz, 1H), 6.73 (d, *J* = 8.9 Hz, 2H), 5.1 (d, *J* = 12.1 Hz, 1H), 3.05 (s, 6H). *E*-isomer: 7.34 (d, *J* = 8.8 Hz, 2H), 7.27 (d, *J* = 16.4 Hz, 1H), 6.67 (d, *J* = 8.5 Hz, 2H), 5.59 (d, *J* = 16.4 Hz, 1H), 3.05 (s, 6H). MS (EI) *m/z* = 172.1 (M⁺).

4-(3-Aminopropyl)-*N,N*-dimethylaniline (24). The title compound was prepared according to a reported procedure with modifications.⁴⁸ The 3-(4-(dimethylamino)phenyl)acrylonitrile **23** (450 mg, 2.61 mmol) was dissolved in anhydrous THF (20 mL) and added dropwise to a freshly prepared mixture suspension of AlCl₃ (1.22 g, 9.144 mmol) and LiAlH₄ (347 mg, 9.144 mmol) in anhydrous THF (20 mL) under argon atmosphere. The reaction mixture was stirred and refluxed for 1 h and then cooled to room temperature. Then 10 mL of water was carefully added to the reaction mixture. Then 0.5 N NaOH (100 mL) was cautiously added to the reaction mixture, which was subsequently extracted with ethyl acetate (3 × 25 mL). The organic layers were separated and combined and washed with brine and dried over sodium sulfate. Filtration and removal of solvent yielded 300 mg (64.4%) of the desired amine as viscous oil which was used directly without further purification. ¹H NMR (300 MHz, DMSO-*d*₆): δ 6.99 (d, *J* = 7.8 Hz, 2H), 6.65 (d, *J* = 7.8 Hz, 2H), 2.95–2.86 (m, 4H), 2.83 (s, 6H), 2.59–2.42 (m, 2H). MS (EI) *m/z* = 178.1 (M⁺).

5-Chloro-*N*-(3-(4-(dimethylamino)phenyl)propyl)-3-ethyl-1*H*-indole-2-carboxamide (26). The title compound was prepared from 5-chloro-3-ethyl-1*H*-indole-2-carboxylic acid **25** (200 mg, 0.89 mmol) and 4-(3-aminopropyl)-*N,N*-dimethyl aniline **24** (192 mg, 1.08 mmol) according to the general procedure D. The crude product was purified by Combiflash chromatography (0–30% ethyl acetate in hexane), which yielded 69 mg (20.2%) of white solid product; mp 164–166 °C. ¹H NMR (500 MHz, chloroform-*d*) δ 9.01 (s, 1H), 7.59 (s, 1H), 7.30 (d, *J* = 8.8 Hz, 1H), 7.22 (d, *J* = 8.8 Hz, 1H), 7.10 (d, *J* = 8.3 Hz, 2H), 6.71 (d, *J* = 8.3 Hz, 2H), 5.98 (bs, 1H), 3.55 (q, *J* = 7.3 Hz, 2H), 2.92 (s, 6H), 2.89 (q, *J* = 7.8 Hz, 2H), 2.67 (t, *J* = 7.3 Hz, 2H), 1.97 (p, *J* = 7.3 Hz, 2H), 1.32 (t, *J* = 7.8 Hz, 3H). MS (EI) *m/z* = 383.2 (M⁺). Anal. Calcd for (C₂₂H₂₆ClN₃O): C, 68.83; H, 6.83; N, 10.95. Found: C, 68.93; H, 6.83; N, 10.68.

AUTHOR INFORMATION

Corresponding Authors

*For D.L.: phone, (361) 221-0745; fax, (361) 221-0793; E-mail, dlu@tamhsc.edu.

*For D.A.K.: phone, (860) 486-1891; E-mail, debra.kendall@uconn.edu.

Notes

The authors declare no competing financial interest.

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ABBREVIATIONS USED

CB1, cannabinoid receptor type 1; CB2, cannabinoid receptor type 2; SAR, structure–activity relationship; [³⁵S]GTPγS, guanosine 5'-*O*-(3-[³⁵S]thio)triphosphate; ERK, extracellular signal-regulated kinases; cAMP, 3'-5'-cyclic adenosine monophosphate; BOP, (benzotriazol-1-yloxy) tris (dimethylamino)-phosphonium hexafluorophosphate; DIPEA, diisopropylethyl amine; DMF, dimethylformamide; rt, room temperature; HEK293, human embryonic kidney 293 cells; BSA, bovine serum albumin; TME, Tris-Mg²⁺-EDTA; EDTA, ethylene

diamine tetraacetate; EGTA, ethylene glycol tetraacetic acid; TLC, thin-layer chromatography; MS, mass spectrometry; PTX, pertussis toxin

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