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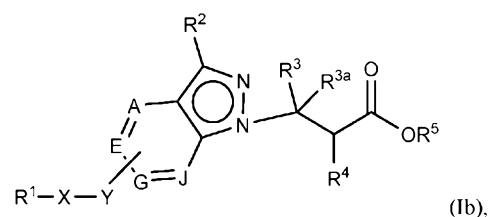
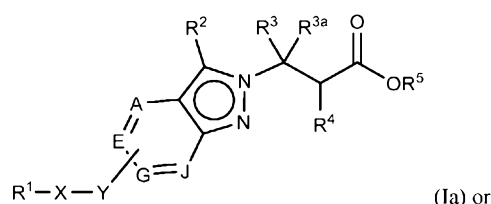
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(54) Title: INDAZOLE DERIVATIVES AS α V INTEGRIN ANTAGONISTS

(57) Abstract: The present invention provides compounds of Formula (Ia) or (Ib): or stereoisomers, tautomers, or pharmaceutically acceptable salts or solvates thereof, wherein all the variables are as defined herein. These compounds are antagonists to α V- containing integrins. This invention also relates to pharmaceutical compositions comprising these compounds and methods of treating a disease, disorder, or condition associated with dysregulation of α V-containing integrins, such as pathological fibrosis, transplant rejection, cancer, osteoporosis, and inflammatory disorders, by using the compounds and pharmaceutical compositions.

INDAZOLE DERIVATIVES AS α_v INTEGRIN ANTAGONISTS

CROSS REFERENCE TO RELATED APPLICATION

This application claims the priority of U.S. Provisional Application serial number 62/418,842 filed November 8, 2016 which is herein incorporated by reference.

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FIELD OF THE INVENTION

The present invention relates to substituted azole amides and amines as α_v integrin antagonists, pharmaceutical compositions comprising such compounds and to their use in therapy, especially in the treatment or prophylaxis of diseases, disorders, and conditions for which an α_v integrin antagonist is indicated in a human.

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BACKGROUND OF THE INVENTION

Integrins belong to a large family of α/β heterodimeric transmembrane proteins that are involved in cell adhesion to a wide variety of extracellular matrix proteins, cell-cell interactions, cell migration, proliferation, survival, and in maintenance of tissue integrity (Barczyk *et al. Cell and Tissue Research* **2010**, 339, 269; Srichai, M. B.; Zent, R. in *Cell-Extracellular Matrix Interactions in Cancer*, **2010**). In mammals, there are 24 α/β integrin heterodimers known from various combinations of 18 alpha and 8 beta subunits. Transforming Growth Factor- β (TGF- β) has a central role in driving a number of pathological processes underlying fibrosis, cell growth, and autoimmune diseases. Alpha V (α_v) Integrins, that include $\alpha_v\beta_1$, $\alpha_v\beta_3$, $\alpha_v\beta_5$, $\alpha_v\beta_6$, and $\alpha_v\beta_8$, are involved in a critical pathway that leads to the conversion of latent TGF- β to its active form (Henderson, N. C.; Sheppard, D. *Biochim, Biophys. Acta* **2013**, 1832, 891). Thus, antagonism of such α_v integrin-mediated activation of latent TGF- β provides a viable therapeutic approach to intervene in TGF- β -driven pathological states (Sheppard, D. *Eur. Resp. Rev.* **2008**, 17, 157; Goodman, S. L.; Picard, M. *Trends Pharmacol. Sciences* **2012**, 33(7), 405; Hinz, B. *Nature Medicine* **2013**, 19(12), 1567; Pozzi, A.; Zent, R. *J. Am. Soc. Nephrol.* **2013**, 24(7), 1034). All five α_v integrins belong to a small subset (8 out of 24) of integrins that recognize the Arginine-Glycine-Aspartic acid (RGD) motif present in

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their native ligands such as fibronectin, vitronectin, and Latency-Associated Peptide (LAP).

The expression of α_v integrin subtypes varies significantly. For example, $\alpha_v\beta_6$ is expressed on epithelial cells at very low levels in healthy tissue but is significantly upregulated during inflammation and wound healing. $\alpha_v\beta_3$ and $\alpha_v\beta_5$ are expressed on osteoclasts, endothelial, smooth muscle, and solid tumor cells, as well as on pericytes and podocytes, while $\alpha_v\beta_1$ is expressed on activated fibroblasts and mesangial cells.

Common fibrotic conditions that represent major unmet medical needs are Idiopathic Pulmonary Fibrosis (IPF), liver and kidney fibrosis, Non-Alcoholic Fatty Liver Disease (NAFLD), Non-Alcoholic Steato-Hepatitis (NASH), as well as systemic sclerosis. Two drugs, pirfenidone and nintedanib, that act by non-integrin-mediated mechanisms, have recently been approved for treatment of IPF. The present invention relates to compounds that inhibit or antagonize the action of one or more of the α_v integrins in the treatment of pathological conditions, such as fibrosis and cancer, mediated by these integrins.

A number of selective or nonselective small molecule, peptidic, and antibody-based antagonists of α_v integrins have been reported in the literature (Kapp, T. G. *et al. Expert Opin. Ther. Patents* **2013**, 23(10), 1273; O'Day, S. *et al. Brit. J. Cancer* **2011**, 105(3), 346; Pickarski, M. *et al. Oncol. Rep.* **2015**, 33, 2737; Wirth, M. *et al. Eur. Urol.* **2014**, 897; Henderson, N. C. *et al. Nature Medicine* **2012**, 19(12), 1617; Horan, G. S. *et al. Am. J. Resp. Crit. Care Med.* **2008**, 177, 56; Puthawala, K. *et al. Am. J. Resp. Crit. Care Med.* **2008**, 177, 82; Reed, N. I. *et al. Sci. Transl. Med.* **2015**, 7(288), 288ra79; Anderson, N. A. *et al.* WO 2014/154725 A1, WO 2016/046225 A1, WO 2016/046226 A1, WO 2016/046230 A1, WO 2016/046241 A1).

SUMMARY OF THE INVENTION

In one aspect, the present invention provides compounds of Formula (Ia), (Ib), (IIa), (IIb), (IIc), (IId), (IIIa), (IIIb), (IIIc), (IIId), (IVa), (IVb), (IVc), (IVd), (IVe) and (IVf) as well as the subgenus and species thereof, including stereoisomers, tautomers,

pharmaceutically acceptable salts, or solvates thereof, which are useful as α_v integrin antagonists.

In another aspect, the present invention also provides processes and intermediates for making the compounds of the present invention.

5 In another aspect, the present invention also provides pharmaceutical compositions comprising a pharmaceutically acceptable carrier and at least one of the compounds of the present invention or stereoisomers, tautomers, pharmaceutically acceptable salts, or solvates thereof.

10 In another aspect, the compounds of the invention may be used in therapy, either alone or in combination with one or more additional therapeutic agents. The compounds of the invention may be used in the treatment of a disease, disorder, or condition associated with dysregulation of α_v -containing integrins in a patient in need of such treatment by administering a therapeutically effective amount of the compound, or a stereoisomer, a tautomer, or a pharmaceutically acceptable salt or solvate thereof, to the
15 patient. The disease, disorder, or condition may be related to pathological fibrosis. The compounds of the invention can be used alone, in combination with one or more compounds of the present invention, or in combination with one or more, e.g., one to two, other therapeutic agents.

The compounds of the invention may be used for the manufacture of a
20 medicament for the treatment of a disease, disorder, or condition associated with dysregulation of α_v -containing integrins in a patient.

Other features and advantages of the invention will be apparent from the following detailed description and claims.

25 DETAILED DESCRIPTION OF THE INVENTION

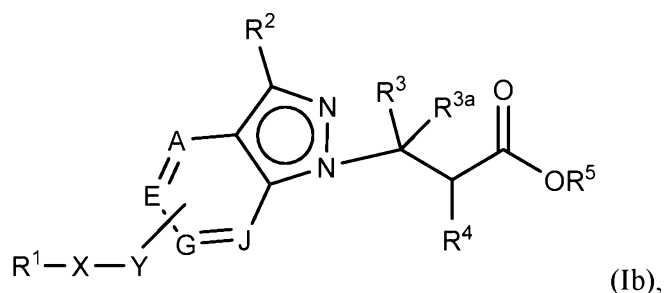
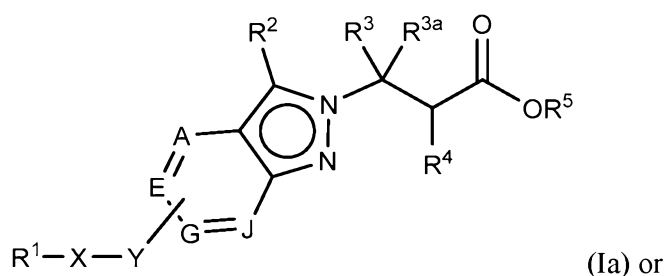
The present application provides compounds, including all stereoisomers, solvates, prodrugs and pharmaceutically acceptable salt and solvate forms thereof, according to Formula (Ia) or (Ib). The present application also provides pharmaceutical compositions containing at least one compound according to Formula (Ia) or (Ib), or a
30 stereoisomer, a tautomer, or a pharmaceutically acceptable salt or a solvate thereof, and optionally at least one additional therapeutic agent. Additionally, the present application

provides methods for treating a patient suffering from an α V Integrin-modulated disease or disorder such as for example, Idiopathic Pulmonary Fibrosis (IPF), liver and kidney fibrosis, Non-Alcoholic Fatty Liver Disease (NAFLD), Non-Alcoholic Steato-Hepatitis (NASH), cardiac fibrosis, and systemic sclerosis, by administering to a patient in need of

5 such treatment a therapeutically effective amount of a compound of the present invention, or a stereoisomer, a tautomer, or a pharmaceutically acceptable salt or a solvate thereof, and optionally in combination with at least one additional therapeutic agent.

In one embodiment, the present invention provides, *inter alia*, a compound of

10 Formula (Ia) or (Ib):



or a stereoisomer, a tautomer, or a pharmaceutically acceptable salt or a solvate thereof, wherein:

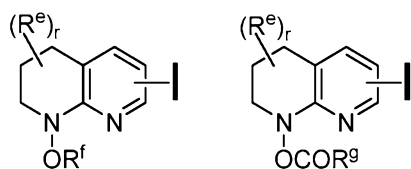
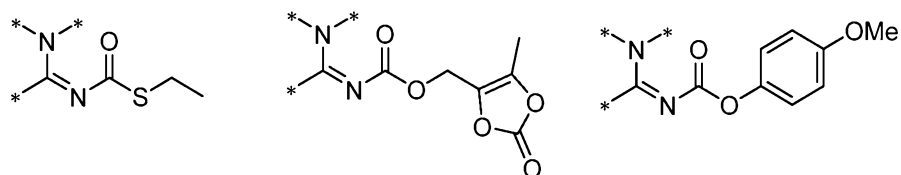
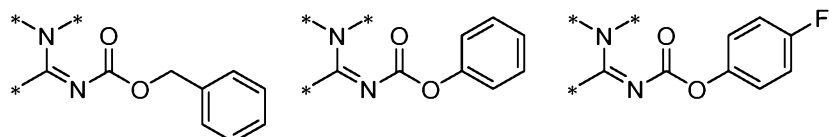
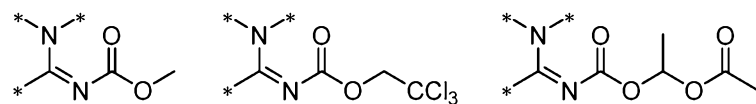
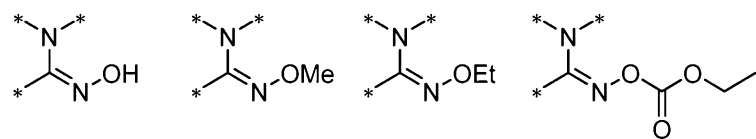
15 A, E, G, and J are independently N, C or CH; with the proviso that at least one of A, E, G, and J is C attached to Y;

X is a C_{1-4} alkylene substituted with 0, 1, or 2 R^{8a} ;

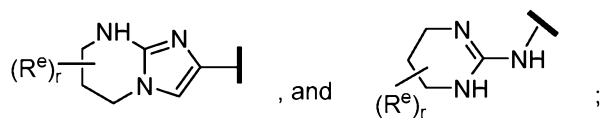
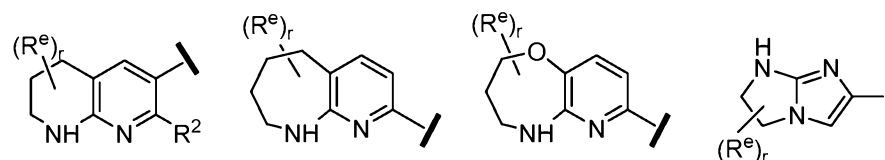
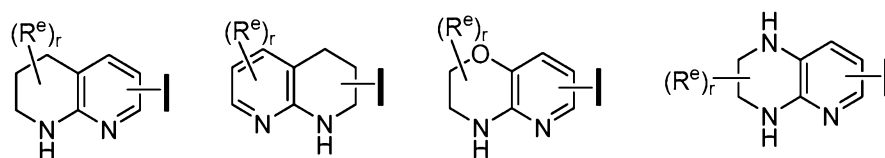
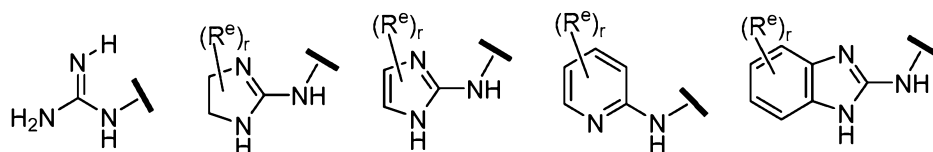
Y is a covalent bond, O, S, NH, -O-(C_{1-3} alkylene)-, -S-(C_{1-3} alkylene)-, or -NH-(C_{1-3} alkylene)-, wherein the C_{1-3} alkylene is each independently substituted with 0,

20 1, or 2 R^{8b} ;

R^1 is an Arginine mimetic moiety selected from the group consisting of



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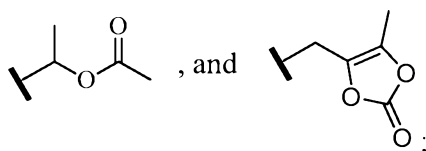
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one of the asterisks in each of the arginine mimetics moiety is an attachment point to X and the other two asterisks are hydrogen;

R^e is OH, amino, amido, carbamate, sulfonamide, C_{1-4} alkyl, halo, C_{1-4} haloalkyl, or C_{3-6} cycloalkyl;

5 R^f = H, CH_3 , CH_2CH_3 , $C(O)OCH_2CH_3$;

R^g = CH_3 , CH_2CCl_3 , phenyl, 4-fluorophenyl, 4-methoxyphenyl, benzyl,



r is an integer of 0, 1, 2, or 3;

R^2 is hydrogen, halo, or C_{1-6} alkyl;

10 R^3 is hydrogen, C_{1-6} alkyl, 3- to 10-membered carbocyclyl, carbocyclylalkyl, 6- to 10-membered aryl, arylalkyl, 3- to 14-membered heterocyclyl, heterocyclylalkyl, 5- to 14-membered heteroaryl, heteroarylalkyl, wherein the alkyl, carbocyclyl, heterocyclyl, aryl, and heteroaryl, by themselves or as part of another group, are each independently substituted with 0, 1, 2, or 3 R^6 ;

15 R^{3a} is hydrogen; or alternatively, R^{3a} and R^3 , together with the atom or atoms to which they are attached, form a 3- to 6-membered carbocyclic or heterocyclic ring which is optionally substituted with one or more groups independently selected from halo, cyano, hydroxyl, amino, C_{1-6} alkyl, haloalkyl, hydroxyalkyl, aminoalkyl, alkoxy, haloalkoxy, amido, carbamate, or sulfonamide;

20 R^4 is hydrogen, C_{1-6} alkyl, 3- to 10-membered carbocyclyl, carbocyclylalkyl, 3- to 10-membered heterocyclyl, heterocyclylalkyl, 6- to 10-membered aryl, arylalkyl, 5- to 14-membered heteroaryl, heteroarylalkyl, $-S(O)_mR^7$, $-C(O)NR^aR^b$, $-NHC(O)OR^a$, $-NHC(O)NR^aR^b$, $-NHC(O)R^7$, $-OC(O)NR^aR^b$, $-OC(O)R^7$, $-NHS(O)_mNR^aR^b$, or $-NHS(O)_mR^7$; wherein the alkyl, carbocyclyl, heterocyclyl, aryl, and heteroaryl, by
25 themselves or as part of another group, are each independently substituted with 0, 1, 2, or 3 R^9 ;

R^6 is halo, cyano, hydroxyl, amino, oxo, nitro, $-S(O)_mR^{12}$, C_{1-6} alkyl, alkoxy, haloalkyl, haloalkoxy, haloaminoalkyl, hydroxyalkyl, aminoalkyl, alkoxycarbonyl, 6- to

10-membered aryl, aryloxy, arylalkoxy, 5- to 10-membered heteroaryl, 3- to 6-membered carbocyclyl, or 3- to 7-membered heterocyclyl; wherein the alkyl, aryl, heteroaryl, carbocyclyl, or heterocyclyl, by themselves or as part of another group, are each independently substituted with 0, 1, or 2 R¹⁰;

5 R⁷ is each independently C₁₋₆ alkyl, C₁₋₆ alkoxy, C₁₋₆ aminoalkyl, C₁₋₆ haloalkyl, 6- to 10-membered aryl, arylalkyl, 5- to 10-membered heteroaryl, cycloalkyl, or heterocycloalkyl; wherein the alkyl, aryl, heteroaryl, cycloalkyl, and heterocycloalkyl, by themselves or as part of another group, are each independently substituted with 0, 1, 2, or 3 R¹¹;

10 R^{8a}, R^{8b}, and R¹¹, at each occurrence, are independently halo, cyano, hydroxyl, amino, oxo, C₁₋₆ alkyl, haloalkyl, hydroxyalkyl, aminoalkyl, alkoxy, or haloalkoxy;

R⁹ is each independently halo, cyano, hydroxyl, amino, oxo, nitro, C₁₋₆ alkyl, alkoxy, haloalkyl, haloalkoxy, haloaminoalkyl, hydroxyalkyl, aminoalkyl, alkoxy carbonyl, 6- to 10-membered aryl, aryloxy, arylalkoxy, 5- to 10-membered
15 heteroaryl, 3- to 6-membered carbocyclyl, or 3- to 7-membered heterocyclyl; wherein the alkyl, aryl, heteroaryl, carbocyclyl, or heterocyclyl, by themselves or as part of another group, are each independently substituted with 0, 1, or 2 R¹³;

R¹⁰ and R¹³ are each independently halo, cyano, hydroxyl, amino, oxo, C₁₋₆ alkyl, haloalkyl, hydroxyalkyl, aminoalkyl, alkoxy, haloalkoxy, amido, carbamate, or
20 sulfonamide;

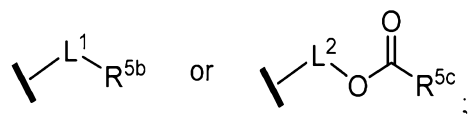
R¹² is -N(R^xR^y), C₁₋₆ alkyl, C₁₋₆ haloalkyl, C₁₋₆ alkoxy, C₁₋₆ haloalkoxy, C₁₋₆ hydroxyalkyl, or C₁₋₆ aminoalkyl; and

R^x and R^y are each independently hydrogen or C₁₋₆ alkyl;

R^a and R^b, at each occurrence, are independently hydrogen, C₁₋₆ alkyl, C₁₋₆ haloalkyl, C₃₋₆ cycloalkyl, 6- to 10-membered aryl, 5- to 10-membered heteroaryl, cycloalkylalkyl, arylalkyl, heteroarylalkyl, or alkoxyalkyl; or alternatively, R^a and R^b,
25 taken together with the atoms to which they are attached, form a 3- to 8-membered carbocyclic or heterocyclic ring which is optionally substituted with one or more groups independently selected from halo, cyano, hydroxyl, amino, oxo, C₁₋₆ alkyl, haloalkyl, hydroxyalkyl, aminoalkyl, alkoxy, haloalkoxy, amido, carbamate, and sulfonamide;
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m is an integer of 1 or 2;

R^{5a} is hydrogen, R^{5b} , or a structural moiety selected from



L^1 and L^2 are each independently C_{1-4} alkylene;

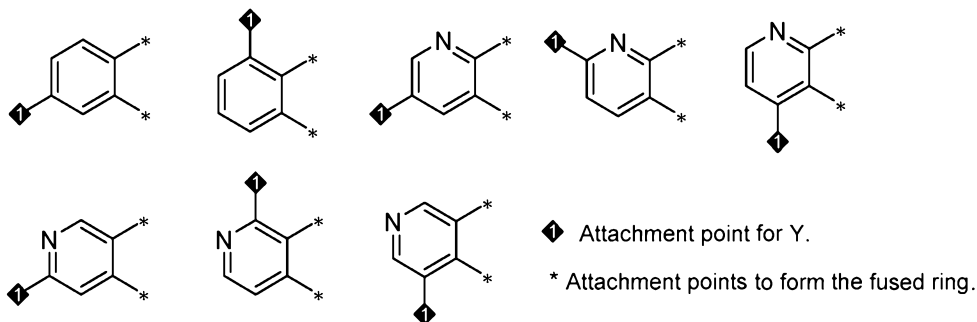
- 5 R^{5a} and R^{5b} are each independently C_{1-6} alkyl, phenyl, benzyl, or 5- to 7-membered heterocyclyl; wherein the alkyl, phenyl, and heterocyclyl are each independently substituted with 0 to 3 R^{5d} ;

R^{5c} is C_{1-6} alkyl or 5- to 7-membered carbocyclyl; wherein the C_{1-6} alkyl, phenyl, and heterocyclyl are each independently substituted with 0 to 3 R^{5d} ; and

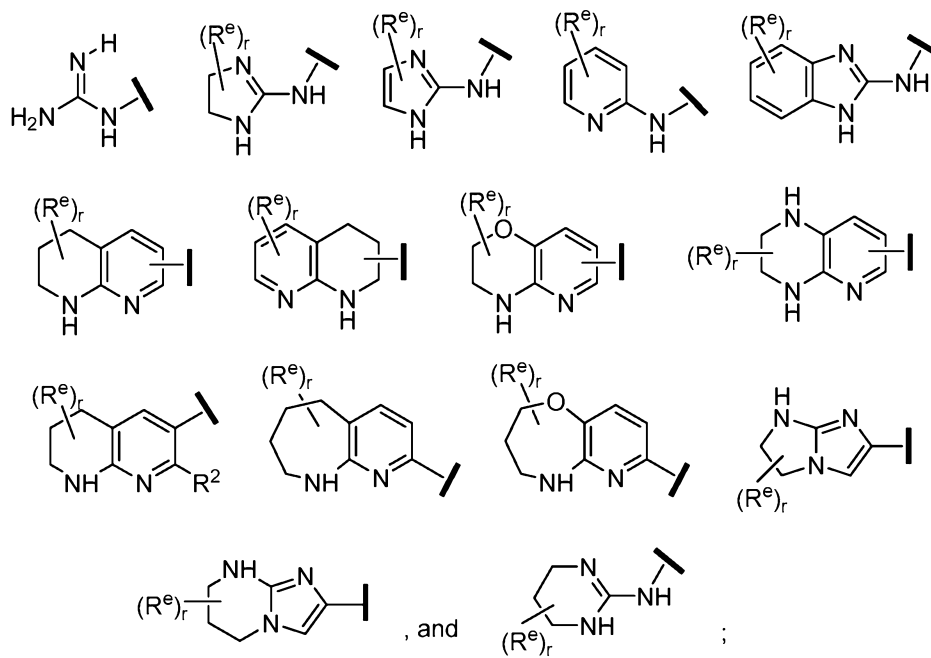
- 10 R^{5d} , at each occurrence, is independently halo, OH, alkoxy, oxo, or alkyl; or alternatively, two adjacent R^{5d} , together with the atoms to which they are attached, form a carbocyclyl moiety.

In one embodiment of Formula (Ia) or (Ib), R^9 is halo, cyano, hydroxyl, amino, oxo, or C_{1-6} alkyl.

- 15 In one embodiment of Formula (Ia) or (Ib), A, E, G, and J, together with the two carbon atoms, form a ring moiety selected from the following structural formula:

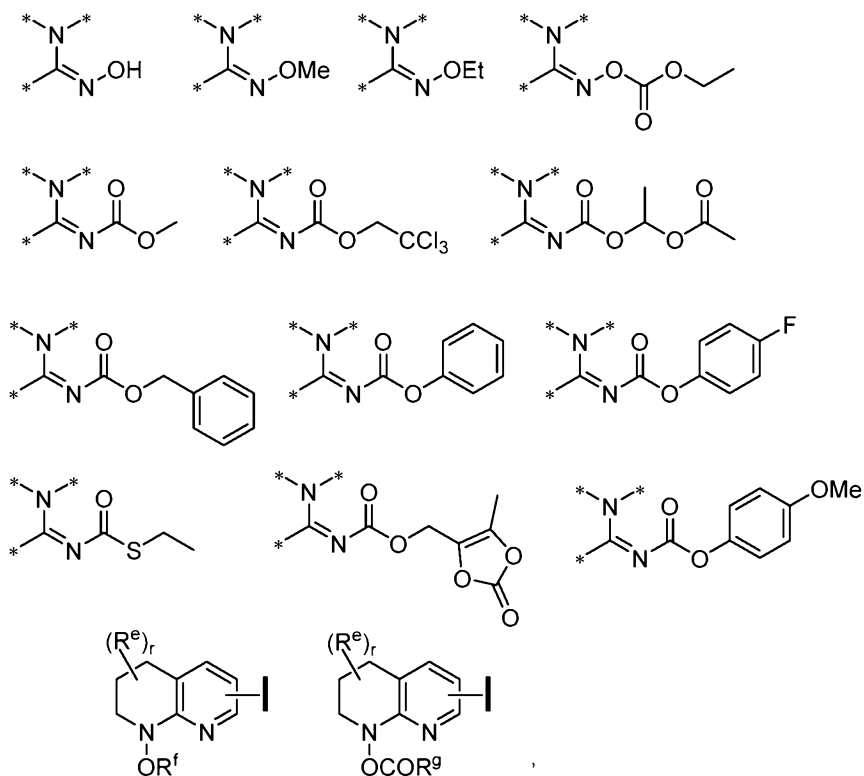


In one embodiment of Formula (Ia) or (Ib), R^1 is an Arginine mimetic moiety selected from the group consisting of



R^e is OH, C_{1-4} alkyl, halo, haloalkyl, or C_{1-4} cycloalkyl; and r is an integer of 0, 1, 2, or 3.

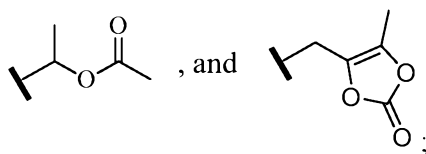
- 5 In one embodiment of Formula (Ia) or (Ib), R^1 is an Arginine mimetic moiety selected from the group consisting of



one of the asterisks in each of the arginine mimetics moiety is an attachment point to X and the other two asterisks are hydrogen;

$R^f = H, Me, Et, COOEt$;

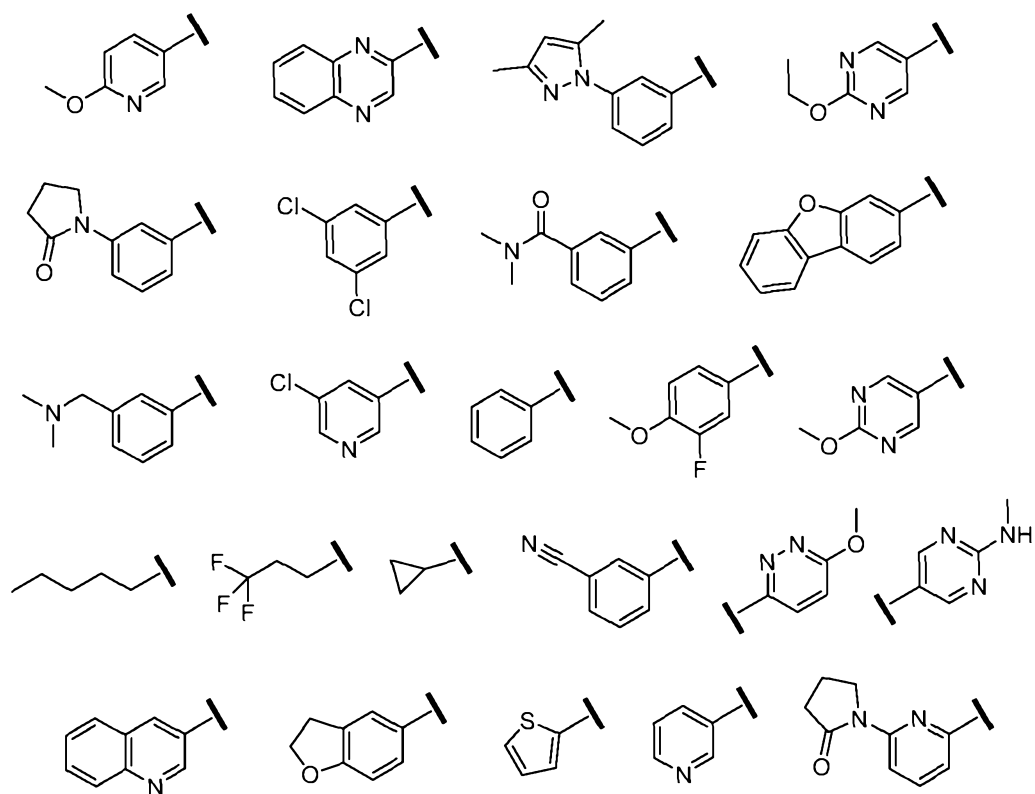
5 $R^g = CH_3, CH_2CH_3, CH_2CCl_3, phenyl, 4-fluorophenyl, 4-methoxyphenyl, benzyl,$

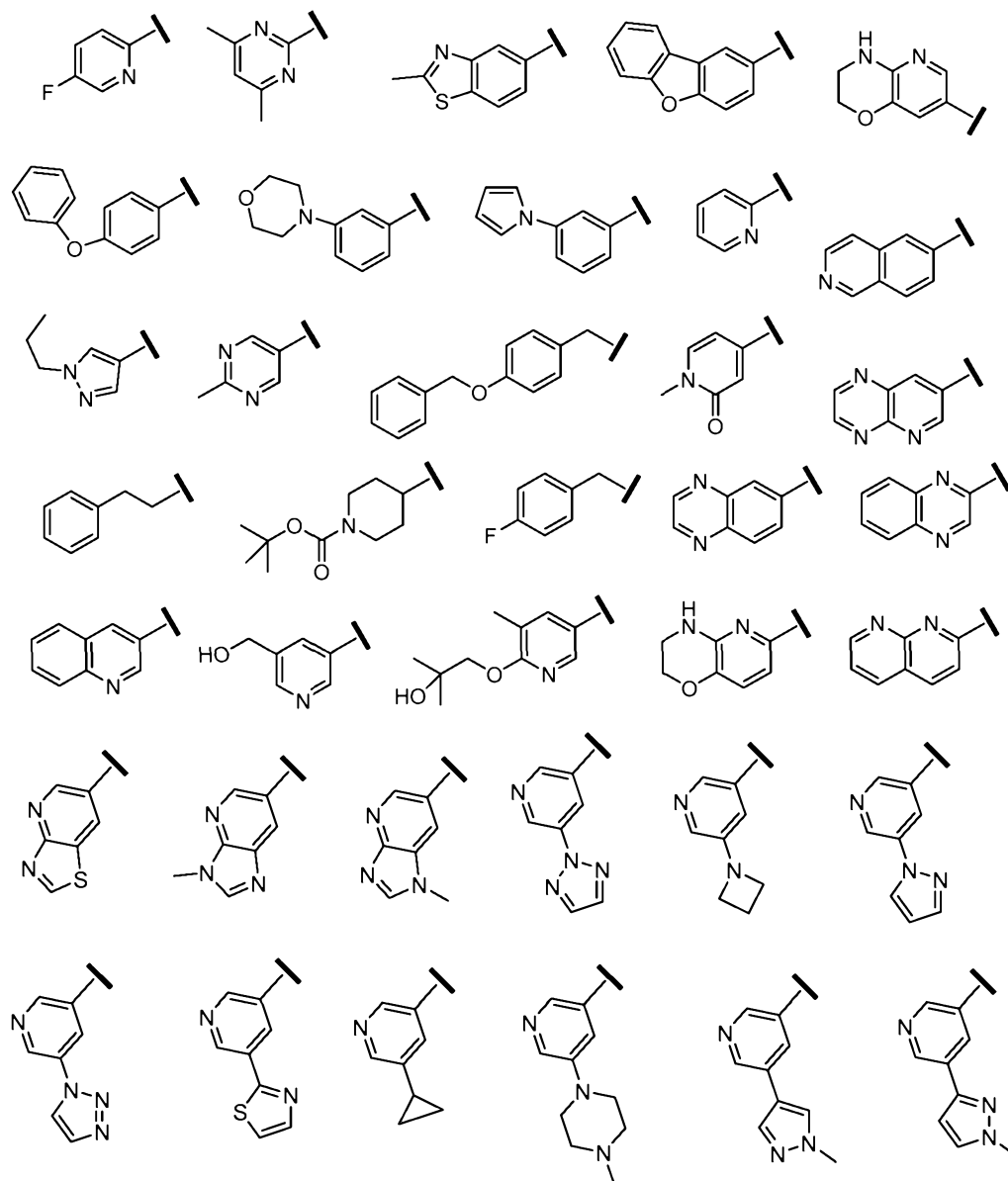


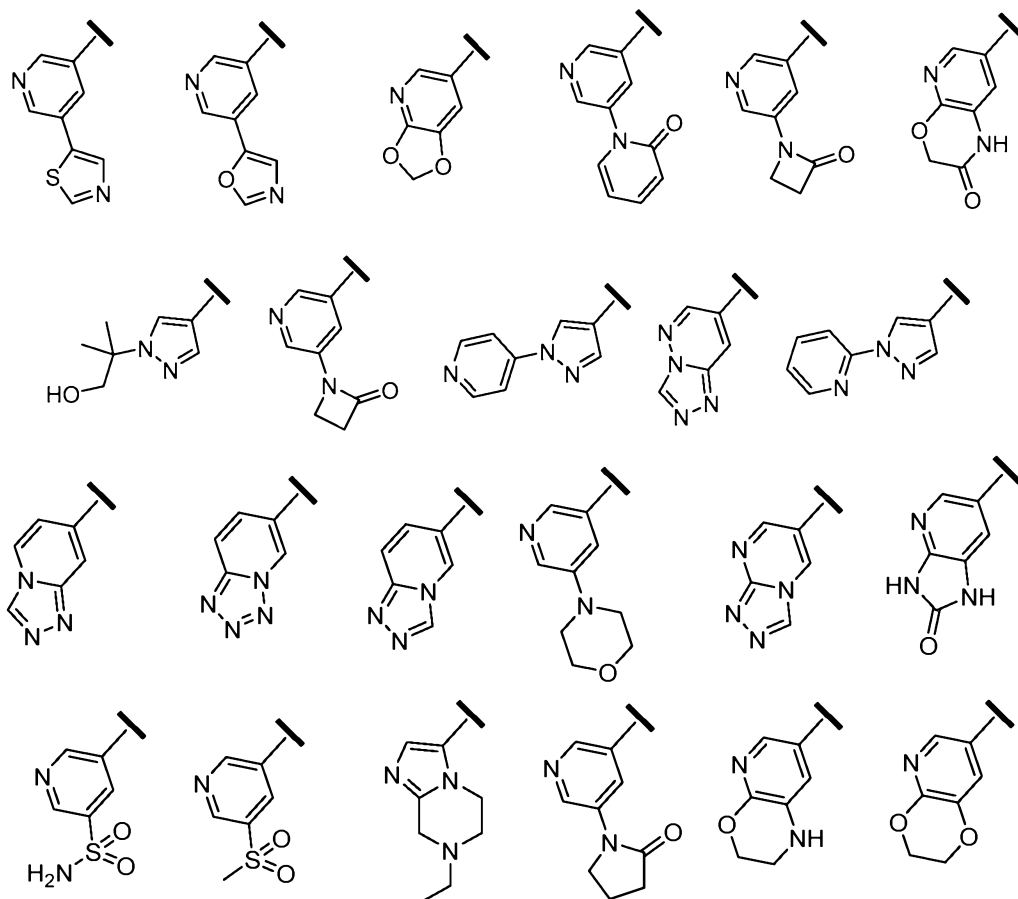
R^e is OH, C_{1-4} alkyl, halo, haloalkyl, or C_{1-4} cycloalkyl; and

r is an integer of 0, 1, 2, or 3.

In one embodiment of Formula (Ia) or (Ib), R^{3a} is hydrogen; and R^3 is hydrogen or
 10 a structural moiety selected from the group consisting of

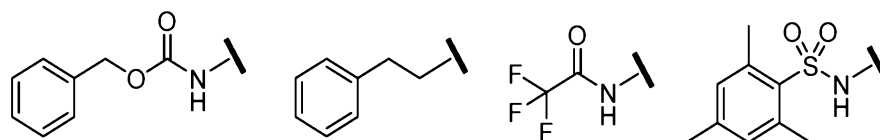






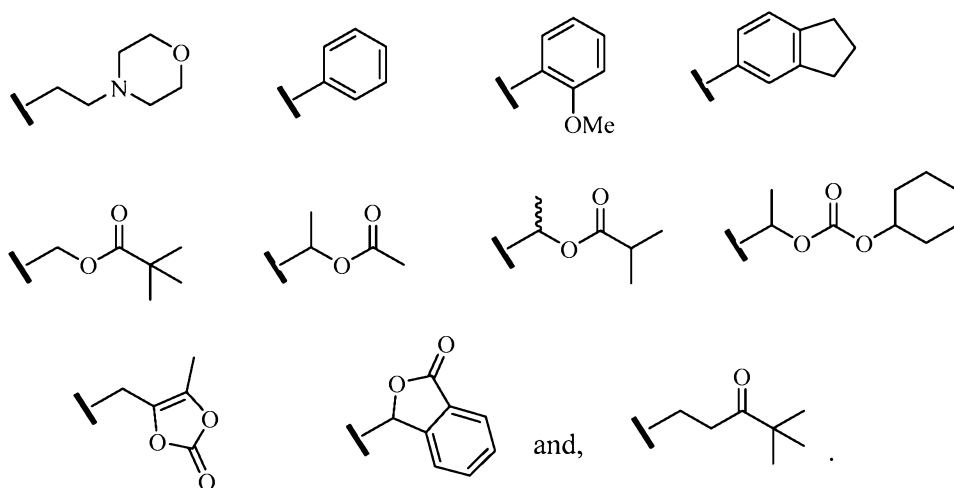
In one embodiment of Formula (Ia) or (Ib), R^{3a} and R^3 , taken together with the atoms to which they are attached, form a C_{3-6} cycloalkyl moiety.

- 5 In one embodiment of Formula (Ia) or (Ib), R^4 is hydrogen or a structural moiety selected from the group consisting of

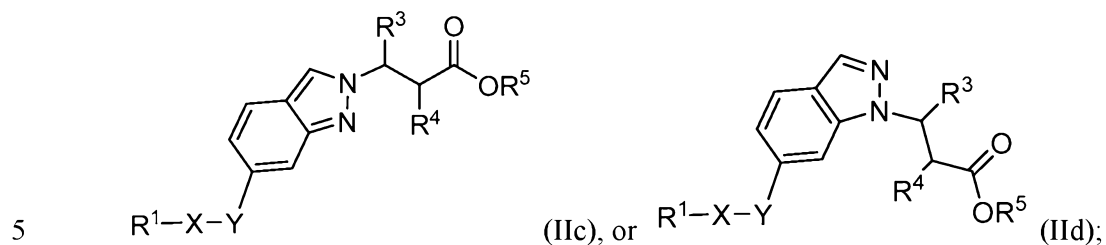
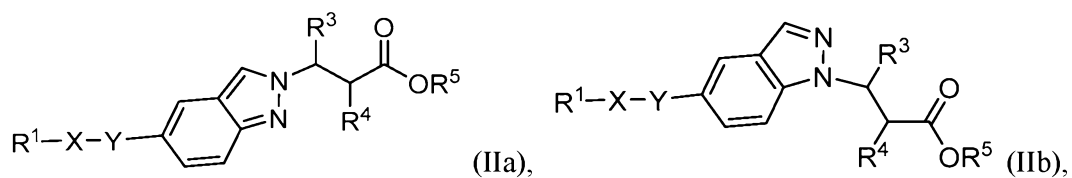


In one embodiment of Formula (Ia) or (Ib), R^4 is H, and R^3 is not H; or alternatively, R^3 is H, and R^4 is not H.

- 10 In one embodiment of Formula (Ia) or (Ib), R^5 is H or R^{5a} ; and R^{5a} is methyl, ethyl, isopropyl, *n*-butyl, isopentyl, or a structural moiety selected from

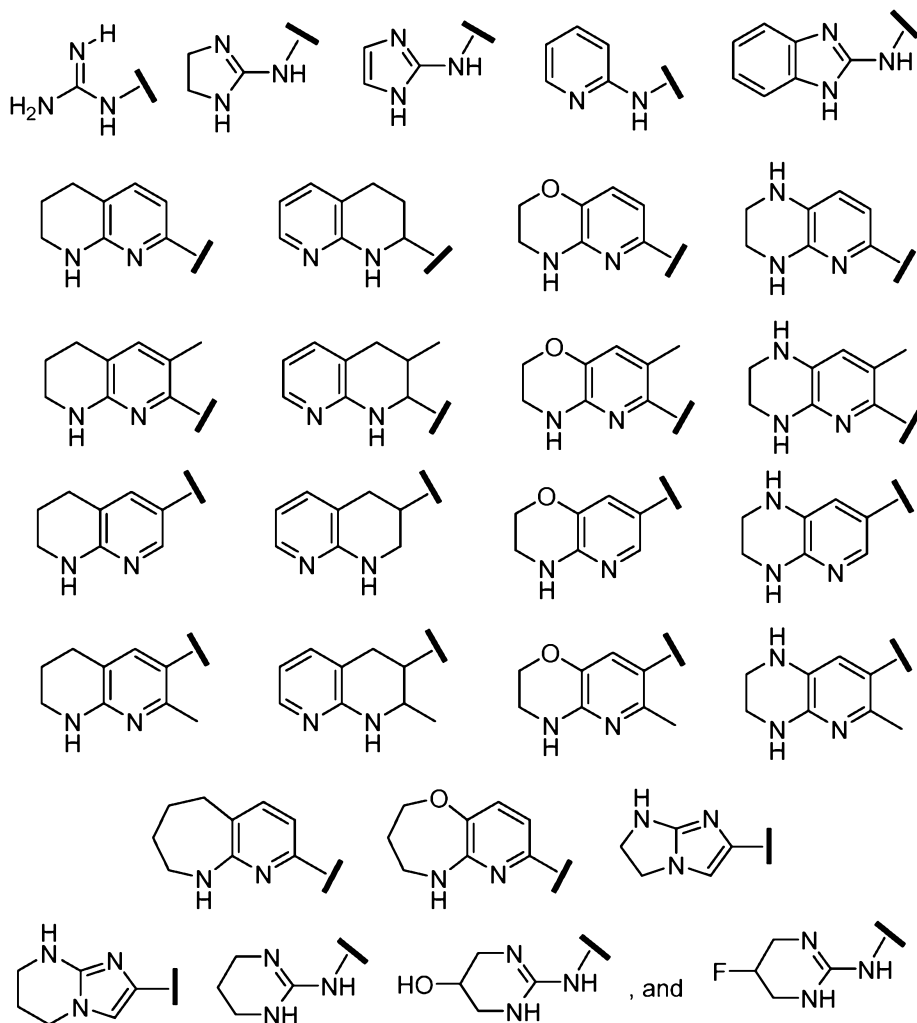


In one embodiment of Formula (Ia) or (Ib), the compound is represented by structural Formula (IIa), (IIb), (IIc), or (IId):



wherein R^1 , X, Y, R^3 , R^4 , and R^5 are the same as defined above.

In one embodiment of Formula (IIa), (IIb), (IIc), or (IId), R^1 is selected from a structural formula selected from the group consisting of



In one embodiment of Formula (IIa), (IIb), (IIc), or (IId), X is C₁₋₄ alkylene; and Y is a covalent bond or O.

5 In one embodiment of Formula (IIa), (IIb), (IIc), or (IId),

R³ is hydrogen, C₁₋₆ alkyl, 3- to 6-membered carbocyclyl, carbocyclylalkyl, 6- to 10-membered aryl, arylalkyl, 3- to 6-membered heterocyclyl, heterocyclylalkyl, 5- to 14-membered heteroaryl, heteroarylalkyl, wherein the alkyl, carbocyclyl, heterocyclyl, aryl, and heteroaryl, by themselves or as part of another group, are each independently

10 substituted with 0, 1, 2, or 3 R⁶;

R^{3a} is hydrogen;

R⁴ is hydrogen;

R⁶ is halo, cyano, hydroxyl, amino, oxo, C₁₋₆ alkyl, alkoxy, haloalkyl, hydroxyalkyl, aminoalkyl, an amide moiety, an ester moiety, 6- to 10-membered aryl, aryloxy, arylalkoxy, 5- to 10-membered heteroaryl, 3- to 6-membered cycloalkyl, or 3- to 6-membered heterocycloalkyl; wherein the alkyl, alkoxy, aminoalkyl, haloalkyl, aryl, 5 aryloxy, heteroaryl, cycloalkyl, or heterocycloalkyl, by themselves or as part of another group, are each independently substituted with 0, 1, or 2 R¹⁰; and

R¹⁰, at each occurrence, is independently halo, cyano, hydroxyl, amino, oxo, or C₁₋₆ alkyl.

In one embodiment of Formula (IIa), (IIb), (IIc), or (IId), R⁴ is hydrogen; R^{3a} and 10 R³, together with the atom or atoms to which they are attached, form a 3- to 6-membered carbocyclic ring.

In one embodiment of Formula (IIa), (IIb), (IIc), or (IId),

R³ is hydrogen;

R^{3a} is hydrogen;

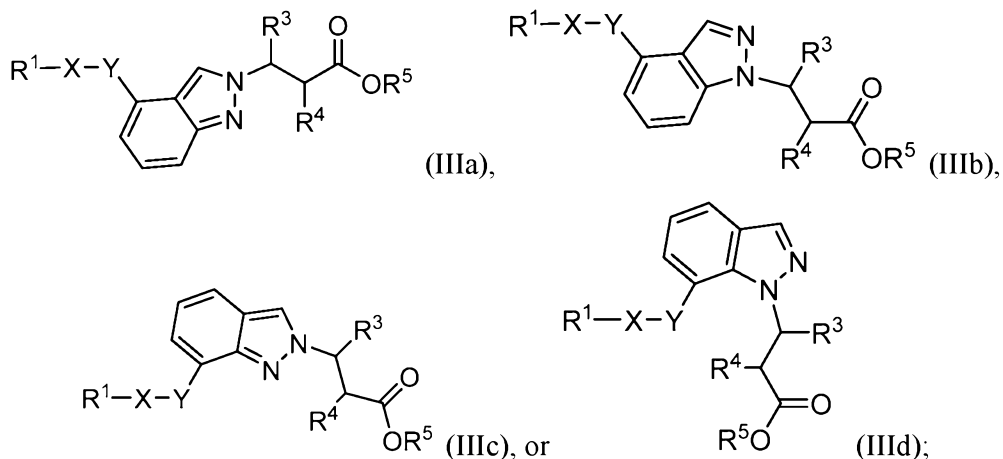
15 R⁴ is C₁₋₆ alkyl, arylalkyl, -S(O)_mR⁷, -C(O)NR^aR^b, -NHC(O)OR^a, -NHC(O)NR^aR^b, -NHC(O)R⁷, -OC(O)NR^aR^b, -OC(O)R⁷, -NHS(O)_mNR^aR^b, or -NHS(O)_mR⁷; wherein the alkyl and arylalkyl are each independently substituted with 0, 1, 2, or 3 R⁹;

R⁷ is each independently C₁₋₆ alkyl, C₁₋₆ alkoxy, C₁₋₆ aminoalkyl, C₁₋₆ haloalkyl, 20 6- to 10-membered aryl, arylalkyl, 5- to 10-membered heteroaryl, cycloalkyl, or heterocycloalkyl; wherein the alkyl, aryl, heteroaryl, cycloalkyl, and heterocycloalkyl, by themselves or as part of another group, are each independently substituted with 0, 1, 2, or 3 R¹¹;

R⁹ and R¹¹, at each occurrence, are independently halo, cyano, hydroxyl, amino, 25 oxo, or C₁₋₆ alkyl.

In one embodiment of Formula (IIa), (IIb), (IIc), or (IId), R⁵ is hydrogen.

In one embodiment of Formula (Ia) or (Ib), the compound is represented by structural Formula (IIIa), (IIIb), (IIIc), or (IIId):



wherein R¹, X, Y, R³, R⁴, and R⁵ are the same as defined above.

In one embodiment of Formula (IIIa), (IIIb), (IIIc), or (IIId), X is C₂₋₄ alkylene;

5 and Y is a covalent bond or O.

In one embodiment of Formula (IIIa), (IIIb), (IIIc), or (IIId),

R³ is C₁₋₆ alkyl, 3- to 6-membered carbocyclyl, carbocyclylalkyl, 6- to 10-membered aryl, arylalkyl, 3- to 6-membered heterocyclyl, heterocyclylalkyl, 5- to 14-membered heteroaryl, heteroarylalkyl, wherein the alkyl, carbocyclyl, heterocyclyl, aryl, and heteroaryl, by themselves or as part of another group, are each independently substituted with 0, 1, 2, or 3 R⁶;

R^{3a} is hydrogen;

R⁴ is hydrogen;

R⁶ is halo, cyano, hydroxyl, amino, oxo, C₁₋₆ alkyl, alkoxy, haloalkyl, hydroxyalkyl, aminoalkyl, an amide moiety, an ester moiety, 6- to 10-membered aryl, aryloxy, arylalkoxy, 5- to 10-membered heteroaryl, 3- to 6-membered cycloalkyl, or 3- to 6-membered heterocycloalkyl; wherein the alkyl, alkoxy, aminoalkyl, haloalkyl, aryl, aryloxy, heteroaryl, cycloalkyl, or heterocycloalkyl, by themselves or as part of another group, are each independently substituted with 0, 1, or 2 R¹⁰; and

R¹⁰, at each occurrence, is independently halo, cyano, hydroxyl, amino, oxo, or C₁₋₆ alkyl.

In one embodiment of Formula (IIIa), (IIIb), (IIIc), or (IIId),

R³ is hydrogen;

R^{3a} is hydrogen;

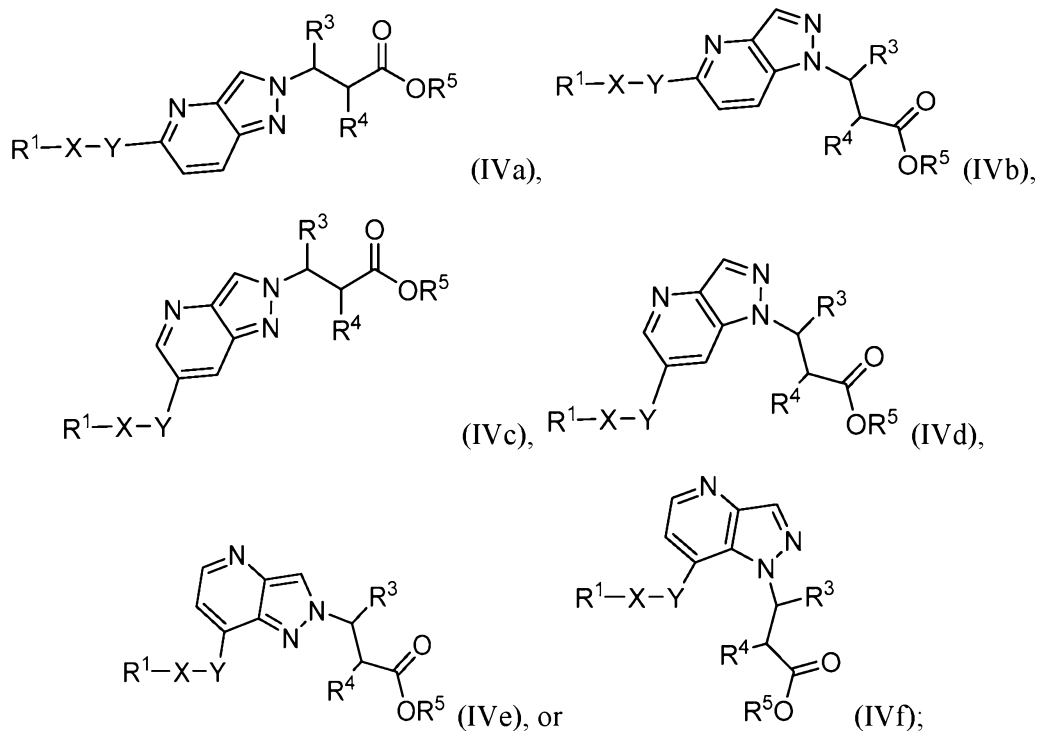
R^4 is C_{1-6} alkyl, arylalkyl, $-S(O)_mR^7$, $-C(O)NR^aR^b$, $-NHC(O)OR^a$, $-NHC(O)NR^aR^b$, $-NHC(O)R^7$, $-OC(O)NR^aR^b$, $-OC(O)R^7$, $-NHS(O)_mNR^aR^b$, or $-NHS(O)_mR^7$; wherein the alkyl and arylalkyl are each independently substituted with 0, 1, 2, or 3 R^9 ;

R^7 is each independently C_{1-6} alkyl, C_{1-6} alkoxy, C_{1-6} aminoalkyl, C_{1-6} haloalkyl, 6- to 10-membered aryl, arylalkyl, 5- to 10-membered heteroaryl, cycloalkyl, or heterocycloalkyl; wherein the alkyl, aryl, heteroaryl, cycloalkyl, and heterocycloalkyl, by themselves or as part of another group, are each independently substituted with 0, 1, 2, or 3 R^{11} ; and

R^9 and R^{11} , at each occurrence, are independently halo, cyano, hydroxyl, amino, oxo, or C_{1-6} alkyl.

In one embodiment of Formula (IIIa), (IIIb), (IIIc), or (IIId), R^5 is hydrogen.

In one embodiment of Formula (Ia) or (Ib), the compound is represented by structural Formula (IVa), (IVb), (IVc), (IVd), (IVe) or (IVf):



wherein R^1 , X , Y , R^3 , R^4 , and R^5 are the same as defined above.

In one embodiment of Formula (IVa), (IVb), (IVc), (IVd), (IVe) or (IVf), X is C₂₋₄ alkylene; and Y is a covalent bond or O.

In one embodiment of Formula (IVa), (IVb), (IVc), (IVd), (IVe) or (IVf),

R³ is C₁₋₆ alkyl, 3- to 6-membered carbocyclyl, carbocyclalkyl, 6- to 10-
 5 membered aryl, arylalkyl, 3- to 6-membered heterocyclyl, heterocyclalkyl, 5- to 14-
 membered heteroaryl, heteroarylalkyl, wherein the alkyl, carbocyclyl, heterocyclyl, aryl,
 and heteroaryl, by themselves or as part of another group, are each independently
 substituted with 0, 1, 2, or 3 R⁶;

R^{3a} is hydrogen;

10 R⁴ is hydrogen;

R⁶ is halo, cyano, hydroxyl, amino, oxo, C₁₋₆ alkyl, alkoxy, haloalkyl,
 hydroxyalkyl, aminoalkyl, an amide moiety, an ester moiety, 6- to 10-membered aryl,
 aryloxy, arylalkoxy, 5- to 10-membered heteroaryl, 3- to 6-membered cycloalkyl, or 3- to
 6-membered heterocycloalkyl; wherein the alkyl, alkoxy, aminoalkyl, haloalkyl, aryl,
 15 aryloxy, heteroaryl, cycloalkyl, or heterocycloalkyl, by themselves or as part of another
 group, are each independently substituted with 0, 1, or 2 R¹⁰; and

R¹⁰, at each occurrence, is independently halo, cyano, hydroxyl, amino, oxo, or
 C₁₋₆ alkyl.

In one embodiment of Formula (IVa), (IVb), (IVc), (IVd), (IVe) or (IVf),

20 R³ is hydrogen;

R^{3a} is hydrogen;

R⁴ is C₁₋₆ alkyl, arylalkyl, -S(O)_mR⁷, -C(O)NR^aR^b, -NHC(O)OR^a,
 -NHC(O)NR^aR^b, -NHC(O)R⁷, -OC(O)NR^aR^b, -OC(O)R⁷, -NHS(O)_mNR^aR^b, or -
 NHS(O)_mR⁷; wherein the alkyl and arylalkyl are each independently substituted with 0, 1,
 25 2, or 3 R⁹;

R⁷ is each independently C₁₋₆ alkyl, C₁₋₆ alkoxy, C₁₋₆ aminoalkyl, C₁₋₆ haloalkyl,
 6- to 10-membered aryl, arylalkyl, 5- to 10-membered heteroaryl, cycloalkyl, or
 heterocycloalkyl; wherein the alkyl, aryl, heteroaryl, cycloalkyl, and heterocycloalkyl, by
 themselves or as part of another group, are each independently substituted with 0, 1, 2, or
 30 3 R¹¹; and

R⁹ and R¹¹, at each occurrence, are independently halo, cyano, hydroxyl, amino, oxo, or C₁₋₆ alkyl.

In one embodiment of Formula (IVa), (IVb), (IVc), (IVd), (IVe) or (IVf), R⁵ is hydrogen.

5 In any one embodiment of Formula (Ia) or (Ib), the compound is selected from any one of the Examples as described in this specification, or a stereoisomer, a tautomer, or a pharmaceutically acceptable salt or solvate thereof.

10 PHARMACEUTICAL COMPOSITIONS, THERAPEUTIC UTILITIES, AND COMBINATIONS

In another embodiment, the present invention provides a composition comprising at least one of the compounds of the present invention, or a stereoisomer, a tautomer, or a pharmaceutically acceptable salt or a solvate thereof.

15 In another embodiment, the present invention provides a pharmaceutical composition comprising a pharmaceutically acceptable carrier and at least one of the compounds of the present invention or a stereoisomer, a tautomer, or a pharmaceutically acceptable salt or a solvate thereof.

20 In another embodiment, the present invention provides a pharmaceutical composition, comprising a pharmaceutically acceptable carrier and a therapeutically effective amount of at least one of the compounds of the present invention or a stereoisomer, a tautomer, or a pharmaceutically acceptable salt or a solvate thereof.

In another embodiment, the present invention provides a process for making a compound of the present invention.

25 In another embodiment, the present invention provides an intermediate for making a compound of the present invention.

In another embodiment, the present invention provides a pharmaceutical composition as defined above further comprising one or more additional therapeutic agents.

30 In another embodiment, the present invention provides a method for the treatment of a disease, disorder, or condition associated with dysregulation of α_v integrins in a

patient in need of such treatment comprising administering a therapeutically effective amount of a compound of the present invention, or a stereoisomer, a tautomer, or a pharmaceutically acceptable salt or solvate thereof, to the patient.

5 In another embodiment, the present invention provides a method for the treatment of the disease, disorder, or condition comprising administering to a patient in need of such treatment a therapeutically effective amount of at least one of the compounds of the present invention, alone, or, optionally, in combination with another compound of the present invention and/or at least one other type of therapeutic agent.

10 In another embodiment, the present invention provides a method for eliciting an integrin receptor antagonizing effect in a patient comprising administering a therapeutically effective amount of a compound of the present invention, or a stereoisomer, a tautomer, or a pharmaceutically acceptable salt or solvate thereof, to the patient. In one embodiment, the integrin receptor antagonizing effect is an antagonizing effect to any of $\alpha_v\beta_6$, $\alpha_v\beta_1$, $\alpha_v\beta_3$, $\alpha_v\beta_5$, and $\alpha_v\beta_8$; or a combination of one or more of
15 $\alpha_v\beta_6$, $\alpha_v\beta_1$, $\alpha_v\beta_3$, $\alpha_v\beta_5$, and $\alpha_v\beta_8$. For example, the integrin receptor antagonizing effect can be an $\alpha_v\beta_6$, $\alpha_v\beta_1$, $\alpha_v\beta_3$, $\alpha_v\beta_5$, and $\alpha_v\beta_8$ antagonizing effect.

In some embodiments, the disease, disorder, or condition is associated with fibrosis, including pulmonary, liver, renal, cardiac, dermal, ocular, and pancreatic fibrosis.

20 In other embodiments, the disease, disorder, or condition is associated with cell-proliferative disorders, such as cancer. In some embodiments, the cancer includes solid tumor growth or neoplasia. In other embodiments, the cancer includes tumor metastasis. In some embodiments, the cancer is of the bladder, blood, bone, brain, breast, central nervous system, cervix, colon, endometrium, esophagus, gall bladder, genitalia,
25 genitourinary tract, head, kidney, larynx, liver, lung, muscle tissue, neck, oral or nasal mucosa, ovary, pancreas, prostate, skin, spleen, small intestine, large intestine, stomach, testicle, or thyroid. In other embodiments, the cancer is a carcinoma, sarcoma, lymphoma, leukemia, melanoma, mesothelioma, multiple myeloma, or seminoma. Examples of diseases, disorders, or conditions associated with the activity of α_v integrins
30 that can be prevented, modulated, or treated according to the present invention include,

but are not limited to, transplant rejection, fibrotic disorders (e. g., idiopathic pulmonary fibrosis (IPF), interstitial lung disease, liver fibrosis, kidney fibrosis, skin fibrosis, systemic sclerosis), inflammatory disorders (e.g., acute hepatitis, chronic hepatitis, non-alcoholic steatohepatitis (NASH), psoriasis, irritable bowel syndrome (IBS), inflammatory bowel disease (IBD)), osteoporosis, as well as cell-proliferative disorders (e.g., cancer, myeloma, fibroma, hepatocarcinoma, leukemia, Kaposi's sarcoma, solid tumors).

The fibrotic disorders, inflammatory disorders, as well as cell-proliferative disorders that are suitable to be prevented or treated by the compounds of the present invention include, but are not limited to, idiopathic pulmonary fibrosis (IPF), interstitial lung disease, non-specific interstitial pneumonia (NSIP), usual interstitial pneumonia (UIP), radiation-induced fibrosis, familial pulmonary fibrosis, airway fibrosis, chronic obstructive pulmonary disease (COPD), diabetic nephropathy, focal segmental glomerulosclerosis, IgA nephropathy, nephropathy induced by drugs or transplantation, autoimmune nephropathy, lupus nephritis, liver fibrosis, kidney fibrosis, chronic kidney disease (CKD), diabetic kidney disease (DKD), skin fibrosis, keloids, systemic sclerosis, scleroderma, virally-induced fibrosis, non-alcoholic fatty liver disease (NAFLD), alcoholic or non-alcoholic steatohepatitis (NASH), acute hepatitis, chronic hepatitis, liver cirrhosis, primary sclerosing cholangitis, drug-induced hepatitis, biliary cirrhosis, portal hypertension, regenerative failure, liver hypofunction, hepatic blood flow disorder, nephropathy, pneumonia, psoriasis, irritable bowel syndrome (IBS), inflammatory bowel disease (IBD), abnormal pancreatic secretion, benign prostatic hyperplasia, neuropathic bladder disease, spinal cord tumor, hernia of intervertebral disk, spinal canal stenosis, heart failure, cardiac fibrosis, vascular fibrosis, perivascular fibrosis, foot-and-mouth disease, cancer, myeloma, fibroma, hepatocarcinoma, leukemia, chronic lymphocytic leukemia, Kaposi's sarcoma, solid tumors, cerebral infarction, cerebral hemorrhage, neuropathic pain, peripheral neuropathy, age-related macular degeneration (AMD), glaucoma, ocular fibrosis, corneal scarring, diabetic retinopathy, proliferative vitreoretinopathy (PVR), cicatricial pemphigoid glaucoma filtration surgery scarring, Crohn's disease or systemic lupus erythematosus; keloid formation resulting from abnormal wound healing; fibrosis occurring after organ transplantation, myelofibrosis,

and fibroids. In one embodiment, the present invention provides a method for the treatment of a fibrotic disorder, an inflammatory disorder, or a cell-proliferative disorder, comprising administering to a patient in need of such treatment a therapeutically effective amount of at least one of the compounds of the present invention, alone, or, optionally, in combination with another compound of the present invention and/or at least one other type of therapeutic agent.

In another embodiment, the present invention provides a compound of the present invention for use in therapy.

In another embodiment, the present invention provides a compound of the present invention for use in therapy for the treatment of a fibrotic disorder, an inflammatory disorder, or a cell-proliferative disorder thereof.

In another embodiment, the present invention also provides the use of a compound of the present invention for the manufacture of a medicament for the treatment of a fibrotic disorder, an inflammatory disorder, or a cell-proliferative disorder thereof.

In another embodiment, the present invention provides a method for the treatment of a fibrotic disorder, an inflammatory disorder, or a cell-proliferative disorder, comprising administering to a patient in need thereof a therapeutically effective amount of a first and second therapeutic agent, wherein the first therapeutic agent is a compound of the present invention.

In another embodiment, the present invention provides a combined preparation of a compound of the present invention and additional therapeutic agent(s) for simultaneous, separate or sequential use in therapy.

In another embodiment, the present invention provides a combined preparation of a compound of the present invention and additional therapeutic agent(s) for simultaneous, separate or sequential use in the treatment of a fibrotic disorder, an inflammatory disorder, or a cell-proliferative disorder.

The compounds of the present invention may be employed in combination with additional therapeutic agent(s), such as one or more anti-fibrotic and/or anti-inflammatory therapeutic agents.

In one embodiment, additional therapeutic agent(s) used in combined pharmaceutical compositions or combined methods or combined uses, are selected from

one or more, preferably one to three, of the following therapeutic agents: inhibitors of TGF β synthesis (for example, pirfenidone), inhibitors of vascular endothelial growth factor (VEGF), platelet-derived growth factor (PDGF) and fibroblast growth factor (FGF) receptor kinases (for example, nintedanib), humanized anti- α v β 6 monoclonal antibody (for example, 3G9), human recombinant pentraxin-2, recombinant human Serum Amyloid P, recombinant human antibody against TGF β -1, -2, and -3, endothelin receptor antagonists (for example, macitentan), interferon gamma, c-Jun amino-terminal kinase (JNK) inhibitor (for example, 4-[[9-[(3*S*)-tetrahydro-3-furanyl]-8-[(2,4,6-trifluorophenyl)amino]-9*H*-purin-2-yl]amino]-trans-cyclohexanol, 3-pentylbenzeneacetic acid (PBI-4050), tetra-substituted porphyrin derivative containing manganese (III), monoclonal antibody targeting eotaxin-2, interleukin-13 (IL-13) antibody (for example, lebrikizumab, tralokinumab), bispecific antibody targeting interleukin 4 (IL-4) and interleukin 13 (IL-13), NK1 tachykinin receptor agonist (for example, Sar⁹, Met(O₂)¹¹-Substance P), Cintredekin Besudotox, human recombinant DNA-derived, IgG1 kappa monoclonal antibody to connective growth factor, and fully human IgG1 kappa antibody, selective for CC-chemokine ligand 2 (for example, carlumab, CCX140), antioxidants (for example, N-acetylcysteine), phosphodiesterase 5 (PDE5) inhibitors (for example, sildenafil), agents for treatment of obstructive airway diseases such as muscarinic antagonists (for example, tiotropium, ipratropium bromide), adrenergic β 2 agonists (for example, salbutamol, salmeterol), corticosteroids (for example, triamcinolone, dexamethasone, fluticasone), immunosuppressive agents (for example, tacrolimus, rapamycin, pimecrolimus), and therapeutic agents useful for the treatment of NALFD, NASH, or systemic sclerosis, such as FXR agonists (for example OCA, GS-9674, and LJN452), LOXL2 inhibitors (for example simtuzumab), LPA1 antagonists (for example SAR 100842), PPAR modulators (for example, elafibrinor, pioglitazone, and saroglitazar, IVA337), SSAO/VAP-1 inhibitors (for example, PXS-4728A and SZE5302), ASK-1 inhibitors (for example GS-4997), ACC inhibitors (for example, CP-640186 and NDI-010976), FGF21 agonist (for example, LY2405319), caspase inhibitors (for example, emricasan), NOX4 inhibitors (for example, GKT137831), MGAT2 inhibitor, and bile acid/fatty acid conjugates (for example aramchol). The α v inhibitors of various

embodiments of the present invention may also be used in combination with one or more therapeutic agents such as CCR2/5 inhibitors (for example, cenicriviroc), Galectin-3 inhibitors (for example, TD-139, GR-MD-02), leukotriene receptor antagonists (for example, tiplukast, montelukast), SGLT2 inhibitors (for example, dapagliflozin, remogliflozin), GLP-1 agonists (for example, liraglutide and semaglutide), FAK inhibitors (for example, GSK-2256098), CB1 inverse agonists (for example, JD-5037), CB2 agonists (for example, APD-371 and JBT-101), autotaxin inhibitors (for example, GLPG1690), prolyl t-RNA synthetase inhibitors (for example, halofuginone), FPR2 agonists (for example, ZK-994), and THR agonists (for example, MGL:3196).

The compounds of this invention can be administered for any of the uses described herein by any suitable means, for example, orally, such as tablets, capsules (each of which includes sustained release or timed release formulations), pills, powders, granules, elixirs, tinctures, suspensions, syrups, and emulsions; sublingually; buccally; parenterally, such as by subcutaneous, intravenous, intramuscular, or intrasternal injection, or infusion techniques (*e.g.*, as sterile injectable aqueous or non-aqueous solutions or suspensions); nasally, including administration to the nasal membranes, such as by inhalation spray; topically, such as in the form of a cream or ointment; or rectally such as in the form of suppositories. They can be administered alone, but generally will be administered with a pharmaceutical carrier selected on the basis of the chosen route of administration and standard pharmaceutical practice.

The term "pharmaceutical composition" means a composition comprising a compound of the invention in combination with at least one additional pharmaceutically acceptable carrier. A "pharmaceutically acceptable carrier" refers to media generally accepted in the art for the delivery of biologically active agents to animals, in particular, mammals, including, *i.e.*, adjuvant, excipient or vehicle, such as diluents, preserving agents, fillers, flow regulating agents, disintegrating agents, wetting agents, emulsifying agents, suspending agents, sweetening agents, flavoring agents, perfuming agents, anti-bacterial agents, anti-fungal agents, lubricating agents and dispensing agents, depending on the nature of the mode of administration and dosage forms. Pharmaceutically acceptable carriers are formulated according to a number of factors well within the purview of those of ordinary skill in the art. These include, without limitation: the type

and nature of the active agent being formulated; the subject to which the agent-containing composition is to be administered; the intended route of administration of the composition; and the therapeutic indication being targeted. Pharmaceutically acceptable carriers include both aqueous and non-aqueous liquid media, as well as a variety of solid and semi-solid dosage forms. Such carriers can include a number of different ingredients and additives in addition to the active agent, such additional ingredients being included in the formulation for a variety of reasons, *e.g.*, stabilization of the active agent, binders, etc., well known to those of ordinary skill in the art. Descriptions of suitable pharmaceutically acceptable carriers, and factors involved in their selection, are found in a variety of readily available sources such as, for example, *Remington's Pharmaceutical Sciences*, 18th Edition (1990).

The terms "treating" or "treatment" as used herein refer to an approach for obtaining beneficial or desired results, including clinical results, by using a compound or a composition of the present invention. For purposes of this invention, beneficial or desired clinical results include, but are not limited to, one or more of the following: decreasing the severity and/or frequency one or more symptoms resulting from the disease, disorder, or condition; diminishing the extent of or causing regression of the disease, disorder, or condition; stabilizing the disease, disorder, or condition (*e.g.*, preventing or delaying the worsening of the disease, disorder, or condition); delay or slowing the progression of the disease, disorder, or condition; ameliorating the disease, disorder, or condition state; decreasing the dose of one or more other medications required to treat the disease, disorder, or condition; and/or increasing the quality of life.

The dosage regimen for the compounds of the present invention will, of course, vary depending upon known factors, such as the pharmacodynamic characteristics of the particular agent and its mode and route of administration; the species, age, sex, health, medical condition, and weight of the recipient; the nature and extent of the symptoms; the kind of concurrent treatment; the frequency of treatment; the route of administration, the renal and hepatic function of the patient, and the effect desired.

By way of general guidance, the daily oral dosage of each active ingredient, when used for the indicated effects, will range between about 0.01 to about 5000 mg per day, preferably between about 0.1 to about 1000 mg per day, and most preferably between

about 0.1 to about 250 mg per day. Intravenously, the most preferred doses will range from about 0.01 to about 10 mg/kg/minute during a constant rate infusion. Compounds of this invention may be administered in a single daily dose, or the total daily dosage may be administered in divided doses of two, three, or four times daily.

5 The compounds are typically administered in admixture with suitable pharmaceutical diluents, excipients, or carriers (collectively referred to herein as pharmaceutical carriers) suitably selected with respect to the intended form of administration, *e.g.*, oral tablets, capsules, elixirs, and syrups, and consistent with conventional pharmaceutical practices.

10 Dosage forms (pharmaceutical compositions) suitable for administration may contain from about 1 milligram to about 2000 milligrams of active ingredient per dosage unit. In these pharmaceutical compositions the active ingredient will ordinarily be present in an amount of about 0.1-95% by weight based on the total weight of the composition.

15 A typical capsule for oral administration contains at least one of the compounds of the present invention (250 mg), lactose (75 mg), and magnesium stearate (15 mg). The mixture is passed through a 60 mesh sieve and packed into a No. 1 gelatin capsule.

 A typical injectable preparation is produced by aseptically placing at least one of the compounds of the present invention (250 mg) into a vial, aseptically freeze-drying
20 and sealing. For use, the contents of the vial are mixed with 2 mL of physiological saline, to produce an injectable preparation.

 The present invention includes within its scope pharmaceutical compositions comprising, as an active ingredient, a therapeutically effective amount of at least one of the compounds of the present invention, alone or in combination with a pharmaceutical
25 carrier. Optionally, compounds of the present invention can be used alone, in combination with other compounds of the invention, or in combination with one or more, preferably one to three, other therapeutic agent(s), *e.g.*, FXR agonists or other pharmaceutically active material.

 The above other therapeutic agents, when employed in combination with the
30 compounds of the present invention may be used, for example, in those amounts

indicated in the *Physicians' Desk Reference*, as in the patents set out above, or as otherwise determined by one of ordinary skill in the art.

Particularly when provided as a single dosage unit, the potential exists for a chemical interaction between the combined active ingredients. For this reason, when the compound of the present invention and a second therapeutic agent are combined in a single dosage unit they are formulated such that although the active ingredients are combined in a single dosage unit, the physical contact between the active ingredients is minimized (that is, reduced). For example, one active ingredient may be enteric coated. By enteric coating one of the active ingredients, it is possible not only to minimize the contact between the combined active ingredients, but also, it is possible to control the release of one of these components in the gastrointestinal tract such that one of these components is not released in the stomach but rather is released in the intestines. One of the active ingredients may also be coated with a material that affects a sustained-release throughout the gastrointestinal tract and also serves to minimize physical contact between the combined active ingredients. Furthermore, the sustained-released component can be additionally enteric coated such that the release of this component occurs only in the intestine. Still another approach would involve the formulation of a combination product in which the one component is coated with a sustained and/or enteric release polymer, and the other component is also coated with a polymer such as a low viscosity grade of hydroxypropyl methylcellulose (HPMC) or other appropriate materials as known in the art, in order to further separate the active components. The polymer coating serves to form an additional barrier to interaction with the other component.

These as well as other ways of minimizing contact between the components of combination products of the present invention, whether administered in a single dosage form or administered in separate forms but at the same time by the same manner, will be readily apparent to those skilled in the art, once armed with the present disclosure.

The compounds of the present invention can be administered alone or in combination with one or more, preferably one to three, additional therapeutic agents. By "administered in combination" or "combination therapy" it is meant that the compound of the present invention and one or more, preferably one to three, additional therapeutic agents are administered concurrently to the mammal being treated. When administered in

combination, each component may be administered at the same time or sequentially in any order at different points in time. Thus, each component may be administered separately but sufficiently closely in time so as to provide the desired therapeutic effect.

The compounds of the present invention are also useful as standard or reference compounds, for example as a quality standard or control, in tests or assays involving the α_v integrins. Such compounds may be provided in a commercial kit, for example, for use in pharmaceutical research involving α_v integrins activity. For example, a compound of the present invention could be used as a reference in an assay to compare its known activity to a compound with an unknown activity. This would ensure the experimenter that the assay was being performed properly and provide a basis for comparison, especially if the test compound was a derivative of the reference compound. When developing new assays or protocols, compounds according to the present invention could be used to test their effectiveness.

The present invention also encompasses an article of manufacture. As used herein, article of manufacture is intended to include, but not be limited to, kits and packages. The article of manufacture of the present invention, comprises: (a) a first container; (b) a pharmaceutical composition located within the first container, wherein the composition, comprises: a first therapeutic agent, comprising a compound of the present invention or a pharmaceutically acceptable salt form thereof; and, (c) a package insert stating that the pharmaceutical composition can be used for the treatment of dyslipidemias and the sequelae thereof. In another embodiment, the package insert states that the pharmaceutical composition can be used in combination (as defined previously) with a second therapeutic agent for the treatment of fibrosis and the sequelae thereof. The article of manufacture can further comprise: (d) a second container, wherein components (a) and (b) are located within the second container and component (c) is located within or outside of the second container. Located within the first and second containers means that the respective container holds the item within its boundaries.

The first container is a receptacle used to hold a pharmaceutical composition. This container can be for manufacturing, storing, shipping, and/or individual/bulk selling. First container is intended to cover a bottle, jar, vial, flask, syringe, tube (*e.g.*, for a cream

preparation), or any other container used to manufacture, hold, store, or distribute a pharmaceutical product.

The second container is one used to hold the first container and, optionally, the package insert. Examples of the second container include, but are not limited to, boxes (e.g., cardboard or plastic), crates, cartons, bags (e.g., paper or plastic bags), pouches, and sacks. The package insert can be physically attached to the outside of the first container via tape, glue, staple, or another method of attachment, or it can rest inside the second container without any physical means of attachment to the first container. Alternatively, the package insert is located on the outside of the second container. When located on the outside of the second container, it is preferable that the package insert is physically attached via tape, glue, staple, or another method of attachment. Alternatively, it can be adjacent to or touching the outside of the second container without being physically attached.

The package insert is a label, tag, marker, etc. that recites information relating to the pharmaceutical composition located within the first container. The information recited will usually be determined by the regulatory agency governing the area in which the article of manufacture is to be sold (e.g., the United States Food and Drug Administration). Preferably, the package insert specifically recites the indications for which the pharmaceutical composition has been approved. The package insert may be made of any material on which a person can read information contained therein or thereon. Preferably, the package insert is a printable material (e.g., paper, plastic, cardboard, foil, adhesive-backed paper or plastic, etc.) on which the desired information has been formed (e.g., printed or applied).

DEFINITIONS

Throughout the specification and the appended claims, a given chemical formula or name shall encompass all stereo and optical isomers and racemates thereof where such isomers exist. Unless otherwise indicated, all chiral (enantiomeric and diastereomeric) and racemic forms are within the scope of the invention. Many geometric isomers of C=C double bonds, C=N double bonds, ring systems, and the like can also be present in the compounds, and all such stable isomers are contemplated in the present invention.

Cis- and *trans*- (or *E*- and *Z*-) geometric isomers of the compounds of the present invention are described and may be isolated as a mixture of isomers or as separated isomeric forms. The present compounds can be isolated in optically active or racemic forms. Optically active forms may be prepared by resolution of racemic forms or by synthesis from optically active starting materials. All processes used to prepare compounds of the present invention and intermediates made therein are considered to be part of the present invention. When enantiomeric or diastereomeric products are prepared, they may be separated by conventional methods, for example, by chromatography or fractional crystallization. Depending on the process conditions the end products of the present invention are obtained either in free (neutral) or salt form. Both the free form and the salts of these end products are within the scope of the invention. If so desired, one form of a compound may be converted into another form. A free base or acid may be converted into a salt; a salt may be converted into the free compound or another salt; a mixture of isomeric compounds of the present invention may be separated into the individual isomers. Compounds of the present invention, free form and salts thereof, may exist in multiple tautomeric forms, in which hydrogen atoms are transposed to other parts of the molecules and the chemical bonds between the atoms of the molecules are consequently rearranged. It should be understood that all tautomeric forms, insofar as they may exist, are included within the invention. As used herein, "a compound of the invention" or "compounds of the invention" means one or more compounds encompassed by Formula (Ia), (Ib), (IIa), (IIb), (IIc), (IId), (IIIa), (IIIb), (IIIc), (IIId), (IVa), (IVb), (IVc), (IVd), (IVe), or (IVf), or a stereoisomer, a tautomer, or a pharmaceutically acceptable salt or solvate thereof.

As used herein, the term "alkyl" or "alkylene" is intended to include both branched and straight-chain saturated aliphatic hydrocarbon groups having the specified number of carbon atoms. For example, "C₁ to C₁₀ alkyl" or "C₁₋₁₀ alkyl" (or alkylene), is intended to include C₁, C₂, C₃, C₄, C₅, C₆, C₇, C₈, C₉, and C₁₀ alkyl groups. Additionally, for example, "C₁ to C₆ alkyl" or "C₁₋₆ alkyl" denotes alkyl having 1 to 6 carbon atoms. Alkyl group can be unsubstituted or substituted with at least one hydrogen being replaced by another chemical group. Example alkyl groups include, but are not

limited to, methyl (Me), ethyl (Et), propyl (*e.g.*, *n*-propyl and isopropyl), butyl (*e.g.*, *n*-butyl, isobutyl, *t*-butyl), and pentyl (*e.g.*, *n*-pentyl, isopentyl, neopentyl). When "C₀ alkyl" or "C₀ alkylene" is used, it is intended to denote a direct bond.

Unless otherwise indicated, the term "lower alkyl" as employed herein alone or as
 5 part of another group includes both straight and branched chain hydrocarbons containing 1 to 8 carbons, and the terms "alkyl" and "alk" as employed herein alone or as part of another group includes both straight and branched chain hydrocarbons containing 1 to 20 carbons, preferably 1 to 10 carbons, more preferably 1 to 8 carbons, in the normal chain, such as methyl, ethyl, propyl, isopropyl, butyl, *t*-butyl, isobutyl, pentyl, hexyl, isohexyl,
 10 heptyl, 4,4-dimethylpentyl, octyl, 2,2,4-trimethylpentyl, nonyl, decyl, undecyl, dodecyl, the various branched chain isomers thereof, and the like as well as such groups including 1 to 4 substituents such as halo, for example F, Br, Cl or I or CF₃, alkyl, alkoxy, aryl, aryloxy, aryl(aryl) or diaryl, arylalkyl, arylalkyloxy, alkenyl, alkynyl, cycloalkyl, cycloalkenyl, cycloalkylalkyl, cycloalkylalkyloxy, hydroxy, hydroxyalkyl, acyl, alkanoyl,
 15 heteroaryl, heteroaryloxy, cycloheteroalkyl, arylheteroaryl, arylalkoxycarbonyl, heteroarylalkyl, heteroarylalkoxy, aryloxyalkyl, aryloxyaryl, alkylamido, alkanoylamino, arylcarbonylamino, nitro, cyano, thiol, haloalkyl, trihaloalkyl and/or alkylthio.

"Heteroalkyl" refers to an alkyl group where one or more carbon atoms have been replaced with a heteroatom, such as, O, N, or S. For example, if the carbon atom of the
 20 alkyl group which is attached to the parent molecule is replaced with a heteroatom (*e.g.*, O, N, or S) the resulting heteroalkyl groups are, respectively, an alkoxy group (*e.g.*, -OCH₃, etc.), an amine (*e.g.*, -NHCH₃, -N(CH₃)₂, etc.), or a thioalkyl group (*e.g.*, -SCH₃). If a non-terminal carbon atom of the alkyl group which is not attached to the parent molecule is replaced with a heteroatom (*e.g.*, O, N, or S) and the resulting heteroalkyl groups are,
 25 respectively, an alkyl ether (*e.g.*, -CH₂CH₂-O-CH₃, etc.), an alkyl amine (*e.g.*, -CH₂NHCH₃, -CH₂N(CH₃)₂, etc.), or a thioalkyl ether (*e.g.*, -CH₂-S-CH₃). If a terminal carbon atom of the alkyl group is replaced with a heteroatom (*e.g.*, O, N, or S), the resulting heteroalkyl groups are, respectively, a hydroxyalkyl group (*e.g.*, -CH₂CH₂-OH), an aminoalkyl group (*e.g.*, -CH₂NH₂), or an alkyl thiol group (*e.g.*, -CH₂CH₂-SH). A
 30 heteroalkyl group can have, for example, 1 to 20 carbon atoms, 1 to 10 carbon atoms, or 1 to

6 carbon atoms. A C₁-C₆ heteroalkyl group means a heteroalkyl group having 1 to 6 carbon atoms.

"Alkenyl" or "alkenylene" is intended to include hydrocarbon chains of either straight or branched configuration having the specified number of carbon atoms and one or more, preferably one to two, carbon-carbon double bonds that may occur in any stable point along the chain. For example, "C₂ to C₆ alkenyl" or "C₂₋₆ alkenyl" (or alkenylene), is intended to include C₂, C₃, C₄, C₅, and C₆ alkenyl groups. Examples of alkenyl include, but are not limited to, ethenyl, 1-propenyl, 2-propenyl, 2-butenyl, 3-butenyl, 2-pentenyl, 3-pentenyl, 4-pentenyl, 2-hexenyl, 3-hexenyl, 4-hexenyl, 5-hexenyl, 2-methyl-2-propenyl, and 4-methyl-3-pentenyl.

"Alkynyl" or "alkynylene" is intended to include hydrocarbon chains of either straight or branched configuration having one or more, preferably one to three, carbon-carbon triple bonds that may occur in any stable point along the chain. For example, "C₂ to C₆ alkynyl" or "C₂₋₆ alkynyl" (or alkynylene), is intended to include C₂, C₃, C₄, C₅, and C₆ alkynyl groups; such as ethynyl, propynyl, butynyl, pentynyl, and hexynyl.

As used herein, "arylalkyl" (a.k.a. aralkyl), "heteroarylalkyl" "carbocyclalkyl" or "heterocyclalkyl" refers to an acyclic alkyl radical in which one of the hydrogen atoms bonded to a carbon atom, typically a terminal or sp³ carbon atom, is replaced with an aryl, heteroaryl, carbocycl, or heterocycl radical, respectively. Typical arylalkyl groups include, but are not limited to, benzyl, 2-phenylethan-1-yl, naphthylmethyl, 2-naphthylethan-1-yl, naphthobenzyl, 2-naphthophenylethan-1-yl and the like. The arylalkyl, heteroarylalkyl, carbocyclalkyl, or heterocyclalkyl group can comprise 4 to 20 carbon atoms and 0 to 5 heteroatoms, *e.g.*, the alkyl moiety may contain 1 to 6 carbon atoms.

The term "benzyl", as used herein, refers to a methyl group on which one of the hydrogen atoms is replaced by a phenyl group, wherein said phenyl group may optionally be substituted with 1 to 5 groups, preferably 1 to 3 groups, OH, OCH₃, Cl, F, Br, I, CN, NO₂, NH₂, N(CH₃)H, N(CH₃)₂, CF₃, OCF₃, C(=O)CH₃, SCH₃, S(=O)CH₃, S(=O)₂CH₃, CH₃, CH₂CH₃, CO₂H, and CO₂CH₃. "Benzyl" can also be represented by formula "Bn".

The term "lower alkoxy", "alkoxy" or "alkyloxy", "aryloxy" or "aralkoxy" refers to any of the above alkyl, aralkyl or aryl groups linked to an oxygen atom. "C₁ to C₆ alkoxy" or "C₁₋₆ alkoxy" (or alkyloxy), is intended to include C₁, C₂, C₃, C₄, C₅, and C₆ alkoxy groups. Example alkoxy groups include, but are not limited to, methoxy, ethoxy, propoxy (*e.g.*, *n*-propoxy and isopropoxy), and *t*-butoxy. Similarly, "lower alkylthio", "alkylthio", "thioalkoxy", "arylthio", or "aralkylthio" represents an alkyl, aryl, or aralkyl group as defined above with the indicated number of carbon atoms attached through a sulphur bridge; for example methyl-S- and ethyl-S-.

The term "alkanoyl" or "alkylcarbonyl" as used herein alone or as part of another group refers to alkyl linked to a carbonyl group. For example, alkylcarbonyl may be represented by alkyl-C(O)-. "C₁ to C₆ alkylcarbonyl" (or alkylcarbonyl), is intended to include C₁, C₂, C₃, C₄, C₅, and C₆ alkyl-C(O)- groups.

The term "alkylsulfonyl" or "sulfonamide", as used herein alone or as part of another group, refers to alkyl or amino linked to a sulfonyl group. For example, alkylsulfonyl may be represented by -S(O)₂R', while sulfonamide may be represented by -S(O)₂NR^cR^d. R' is C₁ to C₆ alkyl; and R^c and R^d are the same as defined below for "amino".

The term "carbamate" as used herein alone or as part of another group refers to oxygen linked to an amido group. For example, carbamate may be represented by N(R^cR^d)-C(O)-O-, and R^c and R^d are the same as defined below for "amino".

The term "amido" as used herein alone or as part of another group refers to amino linked to a carbonyl group. For example, amido may be represented by N(R^cR^d)-C(O)-, and R^c and R^d are the same as defined below for "amino".

The term "amino" is defined as -NR^cR^d, wherein R^c and R^d are independently hydrogen or C₁₋₆ alkyl; or alternatively, R^c and R^d, taken together with the atoms to which they are attached, form a 3- to 8-membered carbocyclic or heterocyclic ring which is optionally substituted with one or more groups independently selected from halo, cyano, hydroxyl, amino, oxo, C₁₋₆ alkyl, haloalkyl, hydroxyalkyl, aminoalkyl, alkoxy, haloalkoxy, amido, carbamate, or sulfonamide. When R^c or R^d (or both of them) is C₁₋₆ alkyl, the amino group can also be referred to as alkylamino. Examples of alkylamino

group include, without limitation, -NH₂, methylamino, ethylamino, propylamino, isopropylamino and the like.

The term "aminoalkyl" refers to an alkyl group on which one of the hydrogen atoms is replaced by an amino group. For example, aminoalkyl may be represented by
5 N(R^cR^d)-alkylene-. "C₁ to C₆" or "C₁₋₆" aminoalkyl", is intended to include C₁, C₂, C₃, C₄, C₅, and C₆ aminoalkyl groups.

The term "halogen" or "halo" as used herein alone or as part of another group refers to chlorine, bromine, fluorine, and iodine, with chlorine or fluorine being preferred.

"Haloalkyl" is intended to include both branched and straight-chain saturated
10 aliphatic hydrocarbon groups having the specified number of carbon atoms, substituted with one or more halogens. "C₁ to C₆ haloalkyl" or "C₁₋₆ haloalkyl" (or haloalkyl), is intended to include C₁, C₂, C₃, C₄, C₅, and C₆ haloalkyl groups. Examples of haloalkyl include, but are not limited to, fluoromethyl, difluoromethyl, trifluoromethyl, trichloromethyl, pentafluoroethyl, pentachloroethyl, 2,2,2-trifluoroethyl,
15 heptafluoropropyl, and heptachloropropyl. Examples of haloalkyl also include "fluoroalkyl" that is intended to include both branched and straight-chain saturated aliphatic hydrocarbon groups having the specified number of carbon atoms, substituted with 1 or more fluorine atoms. The term "polyhaloalkyl" as used herein refers to an "alkyl" group as defined above which includes from 2 to 9, preferably from 2 to 5, halo
20 substituents, such as F or Cl, preferably F, such as polyfluoroalkyl, for example, CF₃CH₂, CF₃ or CF₃CF₂CH₂.

"Haloalkoxy" or "haloalkyloxy" represents a haloalkyl group as defined above with the indicated number of carbon atoms attached through an oxygen bridge. For example, "C₁ to C₆ haloalkoxy" or "C₁₋₆ haloalkoxy", is intended to include C₁, C₂, C₃,
25 C₄, C₅, and C₆ haloalkoxy groups. Examples of haloalkoxy include, but are not limited to, trifluoromethoxy, 2,2,2-trifluoroethoxy, and pentafluoroethoxy. Similarly, "haloalkylthio" or "thiohaloalkoxy" represents a haloalkyl group as defined above with the indicated number of carbon atoms attached through a sulphur bridge; for example trifluoromethyl-S-, and pentafluoroethyl-S-. The term "polyhaloalkyloxy" as used herein
30 refers to an "alkoxy" or "alkyloxy" group as defined above which includes from 2 to 9,

preferably from 2 to 5, halo substituents, such as F or Cl, preferably F, such as polyfluoroalkoxy, for example, $\text{CF}_3\text{CH}_2\text{O}$, CF_3O or $\text{CF}_3\text{CF}_2\text{CH}_2\text{O}$.

"Hydroxyalkyl" are intended to include both branched and straight-chain saturated aliphatic hydrocarbon groups having the specified number of carbon atoms, substituted with 1 or more hydroxyl (OH) or amino, respectively. "C₁ to C₆ hydroxyalkyl" (or hydroxyalkyl), is intended to include C₁, C₂, C₃, C₄, C₅, and C₆ hydroxyalkyl groups.

The term "cycloalkyl" refers to cyclized alkyl groups, including mono-, bi- or poly-cyclic ring systems. "C₃ to C₇ cycloalkyl" or "C₃₋₇ cycloalkyl" is intended to include C₃, C₄, C₅, C₆, and C₇ cycloalkyl groups. Example cycloalkyl groups include, but are not limited to, cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, and norbornyl. Branched cycloalkyl groups such as 1-methylcyclopropyl and 2-methylcyclopropyl are included in the definition of "cycloalkyl".

The term "cycloheteroalkyl" refers to cyclized heteroalkyl groups, including mono-, bi- or poly-cyclic ring systems. "C₃ to C₇ cycloheteroalkyl" or "C₃₋₇ cycloheteroalkyl" is intended to include C₃, C₄, C₅, C₆, and C₇ cycloheteroalkyl groups. Example cycloheteroalkyl groups include, but are not limited to, oxetanyl, tetrahydrofuranyl, tetrahydropyranyl, azetidiny, pyrrolidiny, piperidiny, morpholinyl, and piperazinyl. Branched cycloheteroalkyl groups, such as piperidinylmethyl, piperazinylmethyl, morpholinylmethyl, pyridinylmethyl, pyridizylmethyl, pyrimidylmethyl, and pyrazinylmethyl, are included in the definition of "cycloheteroalkyl".

As used herein, the term "azacycyl" refers to a cycloheteroalkyl containing one or more nitrogen atoms in the ring. Example azacycyl groups include, but are not limited to, pyrrolidiny, piperidiny, morpholinyl, and piperazinyl.

As used herein, "carbocycle", "carbocyclyl", or "carbocyclic" is intended to mean any stable 3-, 4-, 5-, 6-, 7-, or 8-membered monocyclic or 5-, 6-, 7-, 8-, 9-, 10-, 11-, 12-, or 13-membered polycyclic (including bicyclic or tricyclic) hydrocarbon ring, any of which may be saturated or partially unsaturated. That is, the term "carbocycle", "carbocyclyl", or "carbocyclic" includes, without limitation, cycloalkyl and cycloalkenyl. Examples of such carbocycles include, but are not limited to, cyclopropyl, cyclobutyl,

cyclobutenyl, cyclopentyl, cyclopentenyl, cyclohexyl, cycloheptenyl, cycloheptyl, cycloheptenyl, adamantyl, cyclooctyl, cyclooctenyl, cyclooctadienyl, [3.3.0]bicyclooctane, [4.3.0]bicyclononane, [4.4.0]bicyclodecane (decalin), [2.2.2]bicyclooctane, fluorenyl, indanyl, adamantyl, and tetrahydronaphthyl (tetralin). As shown above, bridged rings are also included in the definition of carbocycle (*e.g.*, [2.2.2]bicyclooctane). Preferred carbocycles, unless otherwise specified, are cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, indanyl, and tetrahydronaphthyl. A bridged ring occurs when one or more, preferably one to three, carbon atoms link two non-adjacent carbon atoms. Preferred bridges are one or two carbon atoms. It is noted that a bridge always converts a monocyclic ring into a tricyclic ring. When a ring is bridged, the substituents recited for the ring may also be present on the bridge.

Furthermore, the term "carbocyclyl", including "cycloalkyl" and "cycloalkenyl", as employed herein alone or as part of another group includes saturated or partially unsaturated (containing 1 or 2 double bonds) cyclic hydrocarbon groups containing 1 to 3 rings, including monocyclicalkyl, bicyclicalkyl and tricyclicalkyl, containing a total of 3 to 20 carbons forming the rings, preferably 3 to 10 carbons, forming the ring and which may be fused to 1 or 2 aromatic rings as described for aryl, which include cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, cycloheptyl, cyclooctyl, cyclodecyl and cyclododecyl, cyclohexenyl,



any of which groups may be optionally substituted with 1 to 4 substituents such as halogen, alkyl, alkoxy, hydroxy, aryl, aryloxy, arylalkyl, cycloalkyl, alkylamido, alkanoylamino, oxo, acyl, arylcarbonylamino, nitro, cyano, thiol and/or alkylthio and/or any of the alkyl substituents.

As used herein, the term "bicyclic carbocycle" or "bicyclic carbocyclic group" is intended to mean a stable 9- or 10-membered carbocyclic ring system that contains two fused rings and consists of carbon atoms. Of the two fused rings, one ring is a benzo ring fused to a second ring; and the second ring is a 5- or 6-membered carbon ring which is

saturated or partially unsaturated. The bicyclic carbocyclic group may be attached to its pendant group at any carbon atom which results in a stable structure. The bicyclic carbocyclic group described herein may be substituted on any carbon if the resulting compound is stable. Examples of a bicyclic carbocyclic group are, but not limited to, 1,2-dihydronaphthyl, 1,2,3,4-tetrahydronaphthyl, and indanyl.

As used herein, the term "aryl", as employed herein alone or as part of another group, refers to monocyclic or polycyclic (including bicyclic and tricyclic) aromatic hydrocarbons, including, for example, phenyl, naphthyl, anthracenyl, and phenanthryl.

Aryl moieties are well known and described, for example, in Lewis, R.J., ed., *Hawley's*

Condensed Chemical Dictionary, 13th Edition, John Wiley & Sons, Inc., New York (1997). In one embodiment, the term "aryl" denotes monocyclic and bicyclic aromatic groups containing 6 to 10 carbons in the ring portion (such as phenyl or naphthyl including 1-naphthyl and 2-naphthyl). For example, "C₆ or C₁₀ aryl" or "C₆₋₁₀ aryl" refers to phenyl and naphthyl. Unless otherwise specified, "aryl", "C₆ or C₁₀ aryl",

"C₆₋₁₀ aryl", or "aromatic residue" may be unsubstituted or substituted with 1 to 5 groups, preferably 1 to 3 groups, selected from -OH, -OCH₃, -Cl, -F, -Br, -I, -CN, -NO₂, -NH₂, -N(CH₃)H, -N(CH₃)₂, -CF₃, -OCF₃, -C(O)CH₃, -SCH₃, -S(O)CH₃, -S(O)₂CH₃, -CH₃, -CH₂CH₃, -CO₂H, and -CO₂CH₃.

As used herein, the term "heterocycle", "heterocyclyl", or "heterocyclic group" is intended to mean a stable 3-, 4-, 5-, 6-, or 7-membered monocyclic or 5-, 6-, 7-, 8-, 9-, 10-, 11-, 12-, 13-, or 14-membered polycyclic (including bicyclic and tricyclic) heterocyclic ring that is saturated, or partially unsaturated, and that contains carbon atoms and 1, 2, 3 or 4 heteroatoms independently selected from the group consisting of N, O and S; and including any polycyclic group in which any of the above-defined heterocyclic rings is fused to a carbocyclic or an aryl (e.g., benzene) ring. That is, the term "heterocycle", "heterocyclyl", or "heterocyclic group" includes non-aromatic ring systems, such as heterocycloalkyl and heterocycloalkenyl. The nitrogen and sulfur heteroatoms may optionally be oxidized (*i.e.*, N→O and S(O)_p, wherein p is 0, 1 or 2).

The nitrogen atom may be substituted or unsubstituted (*i.e.*, N or NR wherein R is H or

another substituent, if defined). The heterocyclic ring may be attached to its pendant group at any heteroatom or carbon atom that results in a stable structure. The heterocyclic rings described herein may be substituted on carbon or on a nitrogen atom if the resulting compound is stable. A nitrogen in the heterocycle may optionally be quaternized. It is preferred that when the total number of S and O atoms in the heterocycle exceeds 1, then these heteroatoms are not adjacent to one another. It is preferred that the total number of S and O atoms in the heterocycle is not more than 1. Examples of heterocyclyl include, without limitation, azetidiny, piperaziny, piperidiny, piperidony, piperony, pyran, morpholin, tetrahydrofurany, tetrahydroisoquinolin, tetrahydroquinolin, morpholin, dihydrofuro[2,3-*b*]tetrahydrofuran.

As used herein, the term "bicyclic heterocycle" or "bicyclic heterocyclic group" is intended to mean a stable 9- or 10-membered heterocyclic ring system which contains two fused rings and consists of carbon atoms and 1, 2, 3, or 4 heteroatoms independently selected from the group consisting of N, O and S. Of the two fused rings, one ring is a 5- or 6-membered monocyclic aromatic ring comprising a 5-membered heteroaryl ring, a 6-membered heteroaryl ring or a benzo ring, each fused to a second ring. The second ring is a 5- or 6-membered monocyclic ring which is saturated, partially unsaturated, or unsaturated, and comprises a 5-membered heterocycle, a 6-membered heterocycle or a carbocycle (provided the first ring is not benzo when the second ring is a carbocycle).

The bicyclic heterocyclic group may be attached to its pendant group at any heteroatom or carbon atom which results in a stable structure. The bicyclic heterocyclic group described herein may be substituted on carbon or on a nitrogen atom if the resulting compound is stable. It is preferred that when the total number of S and O atoms in the heterocycle exceeds 1, then these heteroatoms are not adjacent to one another. It is preferred that the total number of S and O atoms in the heterocycle is not more than 1. Examples of a bicyclic heterocyclic group are, but not limited to, 1,2,3,4-tetrahydroquinolin, 1,2,3,4-tetrahydroisoquinolin, 5,6,7,8-tetrahydro-quinolin, 2,3-dihydro-benzofurany, chromany, 1,2,3,4-tetrahydro-quinoxaliny, and 1,2,3,4-tetrahydro-quinazolin.

Bridged rings are also included in the definition of heterocycle. A bridged ring occurs when one or more, preferably one to three, atoms (*i.e.*, C, O, N, or S) link two

non-adjacent carbon or nitrogen atoms. Examples of bridged rings include, but are not limited to, one carbon atom, two carbon atoms, one nitrogen atom, two nitrogen atoms, and a carbon-nitrogen group. It is noted that a bridge always converts a monocyclic ring into a tricyclic ring. When a ring is bridged, the substituents recited for the ring may also
 5 be present on the bridge.

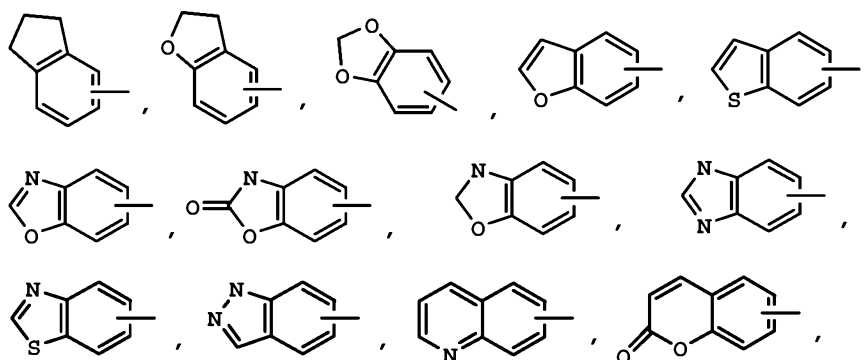
As used herein, the term "heteroaryl" is intended to mean stable monocyclic and polycyclic (including bicyclic and tricyclic) aromatic hydrocarbons that include at least one heteroatom ring member such as sulfur, oxygen, or nitrogen. Heteroaryl groups include, without limitation, pyridyl, pyrimidinyl, pyrazinyl, pyridazinyl, triazinyl, furyl,
 10 quinolyl, isoquinolyl, thienyl, imidazolyl, thiazolyl, indolyl, pyrrolyl, oxazolyl, benzofuryl, benzothienyl, benzthiazolyl, isoxazolyl, pyrazolyl, triazolyl, tetrazolyl, indazolyl, 1,2,4-thiadiazolyl, isothiazolyl, purinyl, carbazolyl, benzimidazolyl, indolinyl, benzodioxolanyl, and benzodioxane. Heteroaryl groups are substituted or unsubstituted. The nitrogen atom is substituted or unsubstituted (*i.e.*, N or NR wherein R is H or another
 15 substituent, if defined). The nitrogen and sulfur heteroatoms may optionally be oxidized (*i.e.*, N→O and S(O)_p, wherein p is 0, 1 or 2).

Examples of heteroaryl include, but are not limited to, acridinyl, azocinyl, benzimidazolyl, benzofuranyl, benzothiofuranyl, benzothiophenyl, benzoxazolyl, benzoxazoliny, benzthiazolyl, benztriazolyl, benztetrazolyl, benzisoxazolyl,
 20 benzisothiazolyl, benzimidazoliny, carbazolyl, 4a*H*-carbazolyl, carbolinyl, chromanyl, chromenyl, cinnoliny, decahydroquinoliny, 2*H*,6*H*-1,5,2-dithiazinyl, furanyl, furazanyl, imidazolidinyl, imidazoliny, imidazolyl, 1*H*-indazolyl, imidazolopyridiny, indolenyl, indolinyl, indoliziny, indolyl, 3*H*-indolyl, isatinoyl, isobenzofuranyl, isochromanyl, isoindazolyl, isoindolinyl, isoindolyl, isoquinoliny, isothiazolyl, isothiazolopyridiny, isoxazolyl, isoxazolopyridiny, methylenedioxyphenyl, naphthyridiny,
 25 octahydroisoquinoliny, oxadiazolyl, 1,2,3-oxadiazolyl, 1,2,4-oxadiazolyl, 1,2,5-oxadiazolyl, 1,3,4-oxadiazolyl, oxazolidiny, oxazolyl, oxazolopyridiny, oxazolidinylperimidiny, oxindolyl, pyrimidinyl, phenanthridiny, phenanthroliny, phenazinyl, phenothiazinyl, phenoxathianyl, phenoxazinyl, phthalazinyl, pteridiny, purinyl, pyrazinyl, pyrazolidiny, pyrazoliny, pyrazolopyridiny, pyrazolyl, pyridazinyl,
 30 pyridooxazolyl, pyridoimidazolyl, pyridothiazolyl, pyridiny, pyrimidinyl, pyrrolidinyl,

pyrrolinyl, 2-pyrrolidonyl, 2H-pyrrolyl, pyrrolyl, quinazolinyl, quinolinyl,
 4*H*-quinoliziny, quinoxaliny, quinuclidiny, tetrazoly, tetrahydrofurany,
 tetrahydroisoquinolinyl, tetrahydroquinolinyl, 6*H*-1,2,5-thiadiaziny, 1,2,3-thiadiazoly,
 1,2,4-thiadiazoly, 1,2,5-thiadiazoly, 1,3,4-thiadiazoly, thianthrenyl, thiazoly, thienyl,
 5 thiazolopyridiny, thienothiazoly, thienooxazoly, thienoimidazoly, thiophenyl, triaziny,
 1,2,3-triazoly, 1,2,4-triazoly, 1,2,5-triazoly, 1,3,4-triazoly, and xanthenyl.


Examples of 5- to 10-membered heteroaryl include, but are not limited to,
 pyridiny, furany, thienyl, pyrazoly, imidazoly, imidazolidiny, indoly, tetrazoly,
 isoxazoly, oxazoly, oxadiazoly, oxazolidiny, thiadiaziny, thiadiazoly, thiazoly,
 10 triaziny, triazoly, benzimidazoly, 1*H*-indazoly, benzofurany, benzothiofurany,
 benzotetrazoly, benzotriazoly, benzisoxazoly, benzoxazoly, oxindoly, benzoxazolinyl,
 benzthiazoly, benzisothiazoly, isatinoy, isoquinolinyl, octahydroisoquinolinyl,
 isoxazolopyridiny, quinazolinyl, quinolinyl, isothiazolopyridiny, thiazolopyridiny,
 oxazolopyridiny, imidazolopyridiny, and pyrazolopyridiny. Examples of 5- to
 15 6-membered heterocycles include, but are not limited to, pyridiny, furany, thienyl,
 pyrroly, pyrazoly, pyraziny, imidazoly, imidazolidiny, indoly, tetrazoly, isoxazoly,
 oxazoly, oxadiazoly, oxazolidiny, thiadiaziny, thiadiazoly, thiazoly, triaziny, and
 triazoly.

Unless otherwise indicated, "carbocyclyl" or "heterocyclyl" includes one to three
 20 additional rings fused to the carbocyclic ring or the heterocyclic ring (such as aryl,
 cycloalkyl, heteroaryl or cycloheteroalkyl rings, for example,






25 and may be optionally substituted through available carbon atoms with 1, 2, or 3 groups
 selected from hydrogen, halo, haloalkyl, alkyl, haloalkyl, alkoxy, haloalkoxy, alkenyl,

trifluoromethyl, trifluoromethoxy, alkynyl, cycloalkyl-alkyl, cycloheteroalkyl, cycloheteroalkylalkyl, aryl, heteroaryl, arylalkyl, aryloxy, aryloxyalkyl, arylalkoxy, alkoxy carbonyl, aryl carbonyl, arylalkenyl, aminocarbonyl aryl, arylthio, arylsulfinyl, arylazo, heteroarylalkyl, heteroarylalkenyl, heteroarylheteroaryl, heteroaryloxy, hydroxy, nitro, cyano, thiol, alkylthio, arylthio, heteroarylthio, arylthioalkyl, alkoxyarylthio, alkyl carbonyl, aryl carbonyl, aminoalkyl carbonyl, arylaminocarbonyl, alkoxy carbonyl, aminocarbonyl, alkyl carbonyloxy, aryl carbonyloxy, alkyl carbonylamino, aryl carbonylamino, arylsulfinyl, arylsulfinylalkyl, arylsulfonylamino and arylsulfonaminocarbonyl and/or any of the alkyl substituents set out herein.

In accordance with a convention used in the art, a bond pointing to a bold line, such as  as used in structural formulas herein, depicts the bond that is the point of attachment of the moiety or substituent to the core or backbone structure.

In accordance with a convention used in the art, a wavy bond in a structural

formula, such as , is used to depict a stereogenic center of the carbon atom to which X', Y', and Z' are attached and is intended to represent both enantiomers in a single figure. That is, a structural formula with such as wavy bond denotes each of the

enantiomers individually, such as  or , as well as a racemic mixture thereof.

It is understood herein that if a carbocyclic, aryl, heterocyclic, or heteroaryl moiety may be bonded or otherwise attached to a designated substrate through differing ring atoms without denoting a specific point of attachment, then all possible points are intended, whether through a carbon atom or, for example, a trivalent nitrogen atom. For example, the term “pyridyl” means 2-, 3- or 4-pyridyl, the term “thienyl” means 2- or 3-thienyl, and so forth.

When a dotted ring is used within a ring structure, this indicates that the ring structure may be saturated, partially saturated or unsaturated.

When a bond to a substituent is shown to cross a bond connecting two atoms in a ring, then such substituent may be bonded to any atom on the ring. When a substituent is

listed without indicating the atom in which such substituent is bonded to the rest of the compound of a given formula, then such substituent may be bonded via any atom in such substituent. Combinations of substituents and/or variables are permissible only if such combinations result in stable compounds.

5 One skilled in the art will recognize that substituents and other moieties of the compounds of the present invention should be selected in order to provide a compound which is sufficiently stable to provide a pharmaceutically useful compound which can be formulated into an acceptably stable pharmaceutical composition. Compounds of the present invention which have such stability are contemplated as falling within the scope of
10 the present invention.

 The term "counter ion" is used to represent a negatively charged species such as chloride, bromide, hydroxide, acetate, and sulfate. The term "metal ion" refers to alkali metal ions such as sodium, potassium or lithium and alkaline earth metal ions such as magnesium and calcium, as well as zinc and aluminum.

15 As referred to herein, the term "substituted" means that at least one hydrogen atom is replaced with a non-hydrogen group, provided that normal valencies are maintained and that the substitution results in a stable compound. When a substituent is keto (*i.e.*, =O), then 2 hydrogens on the atom are replaced. Keto substituents are not present on aromatic moieties. When a ring system (*e.g.*, carbocyclic or heterocyclic) is
20 said to be substituted with a carbonyl group or a double bond, it is intended that the carbonyl group or double bond be part (*i.e.*, within) of the ring. Ring double bonds, as used herein, are double bonds that are formed between two adjacent ring atoms (*e.g.*, C=C, C=N, or N=N).

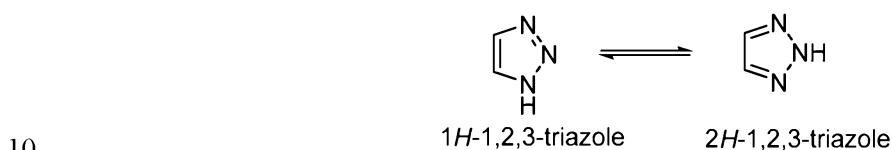
 In cases wherein there are nitrogen atoms (*e.g.*, amines) on compounds of the
25 present invention, these may be converted to N-oxides by treatment with an oxidizing agent (*e.g.*, mCPBA and/or hydrogen peroxides) to afford other compounds of this invention. Thus, shown and claimed nitrogen atoms are considered to cover both the shown nitrogen and its N-oxide (N→O) derivative.

 When any variable occurs more than one time in any constituent or formula for a
30 compound, its definition at each occurrence is independent of its definition at every other occurrence. Thus, for example, if a group is shown to be substituted with 0, 1, 2, or 3 R

groups, then said group be unsubstituted when it is substituted with 0 R group, or be substituted with up to three R groups, and at each occurrence R is selected independently from the definition of R.

Also, combinations of substituents and/or variables are permissible only if such combinations result in stable compounds.

As used herein, the term "tautomer" refers to each of two or more isomers of a compound that exist together in equilibrium, and are readily interchanged by migration of an atom or group within the molecule. For example, one skilled in the art would readily understand that a 1,2,3-triazole exists in two tautomeric forms as defined above:



Thus, this disclosure is intended to cover all possible tautomers even when a structure depicts only one of them.

The phrase "pharmaceutically acceptable" is employed herein to refer to those compounds, materials, compositions, and/or dosage forms that are, within the scope of sound medical judgment, suitable for use in contact with the tissues of human beings and animals without excessive toxicity, irritation, allergic response, and/or other problem or complication, commensurate with a reasonable benefit/risk ratio.

The compounds of the present invention can be present as salts, which are also within the scope of this invention. Pharmaceutically acceptable salts are preferred. As used herein, "pharmaceutically acceptable salts" refer to derivatives of the disclosed compounds wherein the parent compound is modified by making acid or base salts thereof. The pharmaceutically acceptable salts of the present invention can be synthesized from the parent compound that contains a basic or acidic moiety by conventional chemical methods. Generally, such salts can be prepared by reacting the free acid or base forms of these compounds with a stoichiometric amount of the appropriate base or acid in water or in an organic solvent, or in a mixture of the two; generally, nonaqueous media like ether, ethyl acetate, ethanol, isopropanol, or acetonitrile are preferred. Lists of suitable salts are found in *Remington's Pharmaceutical Sciences*,

18th Edition, Mack Publishing Company, Easton, PA (1990), the disclosure of which is hereby incorporated by reference.

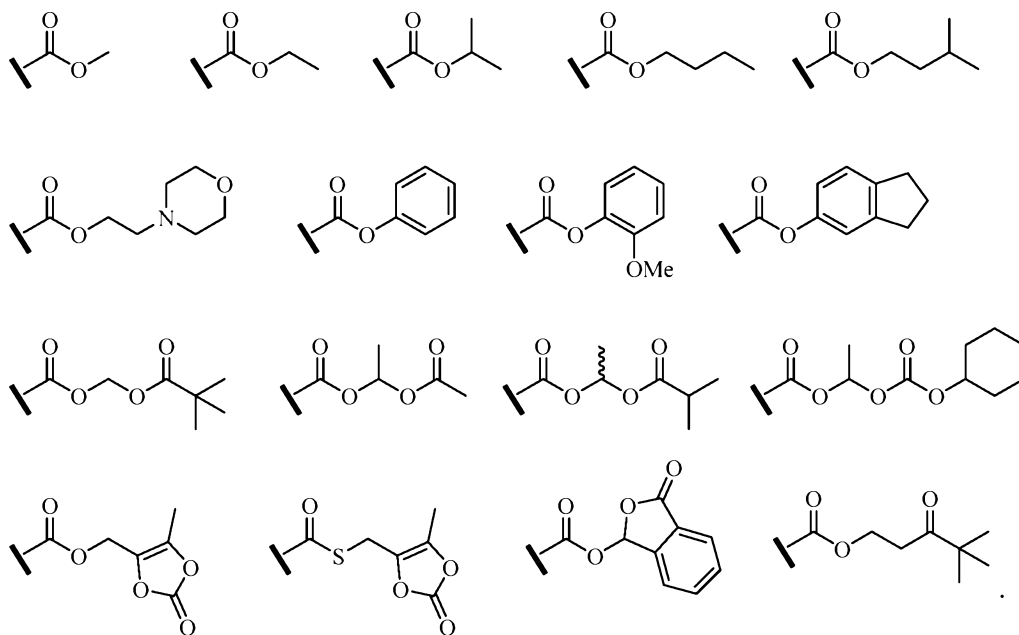
If the compounds of the present invention have, for example, at least one basic center, they can form acid addition salts. These are formed, for example, with strong inorganic acids, such as mineral acids, for example sulfuric acid, phosphoric acid or a hydrohalic acid, with organic carboxylic acids, such as alkanecarboxylic acids of 1 to 4 carbon atoms, for example acetic acid, which are unsubstituted or substituted, for example, by halogen as chloroacetic acid, such as saturated or unsaturated dicarboxylic acids, for example oxalic, malonic, succinic, maleic, fumaric, phthalic or terephthalic acid, such as hydroxycarboxylic acids, for example ascorbic, glycolic, lactic, malic, tartaric or citric acid, such as amino acids, (for example aspartic or glutamic acid or lysine or arginine), or benzoic acid, or with organic sulfonic acids, such as (C₁-C₄) alkyl or arylsulfonic acids which are unsubstituted or substituted, for example by halogen, for example methyl- or p-toluene- sulfonic acid. Corresponding acid addition salts can also be formed having, if desired, an additionally present basic center. The compounds of the present invention having at least one acid group (for example COOH) can also form salts with bases. Suitable salts with bases are, for example, metal salts, such as alkali metal or alkaline earth metal salts, for example sodium, potassium or magnesium salts, or salts with ammonia or an organic amine, such as morpholine, thiomorpholine, piperidine, pyrrolidine, a mono, di or tri-lower alkylamine, for example ethyl, tert-butyl, diethyl, diisopropyl, triethyl, tributyl or dimethyl-propylamine, or a mono, di or trihydroxy lower alkylamine, for example mono, di or triethanolamine. Corresponding internal salts may furthermore be formed. Salts which are unsuitable for pharmaceutical uses but which can be employed, for example, for the isolation or purification of free compounds of Formula I or their pharmaceutically acceptable salts, are also included.

Preferred salts of the compounds of Formula I which contain a basic group include monohydrochloride, hydrogensulfate, methanesulfonate, phosphate, nitrate or acetate.

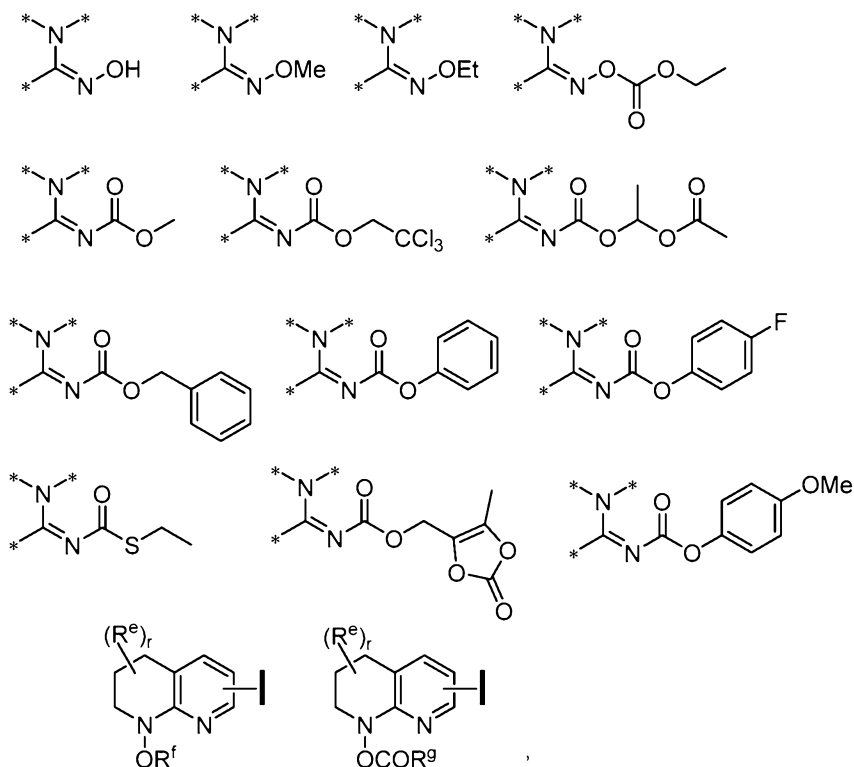
Preferred salts of the compounds of Formula I which contain an acid group include sodium, potassium and magnesium salts and pharmaceutically acceptable organic amines.

In addition, the compounds of the present invention may have prodrug forms. Any compound that will be converted *in vivo* to provide the bioactive agent is a prodrug within the scope and spirit of the invention. The term “prodrug” as used herein encompasses both the prodrugs based on the carboxylic acid residue, i.e., “prodrug esters”, and the prodrugs based on the arginine mimetics moiety, i.e., “prodrugs of arginine mimetics”. Such prodrugs are preferably administered orally since hydrolysis in many instances occurs principally under the influence of the digestive enzymes. Parenteral administration may be used where the ester *per se* is active, or in those instances where hydrolysis occurs in the blood.

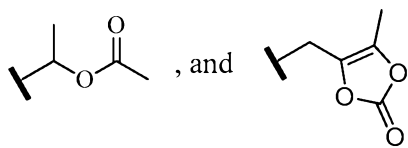
The compounds of the present invention contain a carboxy group which can form physiologically hydrolyzable esters that serve as prodrugs, i.e., “prodrug esters”, by being hydrolyzed in the body to yield the compounds of the present invention *per se*. Examples of physiologically hydrolyzable esters of compounds of the present invention include C₁ to C₆ alkyl, C₁ to C₆ alkylbenzyl, 4-methoxybenzyl, indanyl, phthalyl, methoxymethyl, C₁₋₆ alkanoyloxy-C₁₋₆ alkyl (*e.g.*, acetoxymethyl, pivaloyloxymethyl or propionyloxymethyl), C₁ to C₆ alkoxycarbonyloxy-C₁ to C₆ alkyl (*e.g.*, methoxycarbonyl-oxymethyl or ethoxycarbonyloxymethyl, glycyloxymethyl, phenylglycyloxymethyl, (5-methyl-2-oxo-1,3-dioxolen-4-yl)-methyl), and other well known physiologically hydrolyzable esters used, for example, in the penicillin and cephalosporin arts. Such esters may be prepared by conventional techniques known in the art. The “prodrug esters” can be formed by reacting the carboxylic acid moiety of the compounds of the present invention with either alkyl or aryl alcohol, halide, or sulfonate employing procedures known to those skilled in the art. Examples of such prodrug esters include:



The compounds of the present invention contain an arginine mimetics moiety which can form physiologically hydrolyzable esters that serve as prodrugs, i.e., “prodrugs of arginine mimetics”, by being hydrolyzed in the body to yield the compounds of the present invention *per se*. Representative examples of prodrugs of arginine mimetics include:



wherein, one of the asterisks in each of the arginine mimetics moiety is an attachment point to the parent molecule and the other two asterisks are hydrogen; $R^f = H, Me, Et, COOEt$; $R^g = CH_3, CH_2CCl_3, phenyl, 4-fluorophenyl, 4-methoxyphenyl, benzyl$,



5

; R^e is $OH, C_{1-4} alkyl, halo, haloalkyl, or C_{1-4} cycloalkyl$; and r is an integer of 0, 1, 2, or 3.

Furthermore, various forms of prodrugs are well known in the art. For examples of such prodrug derivatives, see:

Bundgaard, H., ed., *Design of Prodrugs*, Elsevier (1985), and Widder, K. et al., eds., *Methods in Enzymology*, 112:309-396, Academic Press (1985);

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Bundgaard, H., Chapter 5, "Design and Application of Prodrugs", Krosgaard-Larsen, P. et al., eds., *A Textbook of Drug Design and Development*, pp. 113-191, Harwood Academic Publishers (1991);

Bundgaard, H., *Adv. Drug Deliv. Rev.*, 8:1-38 (1992);

Bundgaard, H. et al., *J. Pharm. Sci.*, 77:285 (1988); and

Takeya, N. et al., *Chem. Pharm. Bull.*, 32:692 (1984).

Preparation of prodrugs is well known in the art and described in, for example, King, F.D., ed., *Medicinal Chemistry: Principles and Practice*, The Royal Society of
5 Chemistry, Cambridge, UK (1994); Testa, B. et al., *Hydrolysis in Drug and Prodrug Metabolism. Chemistry, Biochemistry and Enzymology*, VCHA and Wiley-VCH, Zurich, Switzerland (2003); Wermuth, C.G., ed., *The Practice of Medicinal Chemistry*, Academic Press, San Diego, CA (1999).

The present invention is intended to include all isotopes of atoms occurring in the
10 present compounds. Isotopes include those atoms having the same atomic number but different mass numbers. By way of general example and without limitation, isotopes of hydrogen include deuterium and tritium. Isotopes of carbon include ^{13}C and ^{14}C . Isotopically-labeled compounds of the invention can generally be prepared by conventional techniques known to those skilled in the art or by processes analogous to
15 those described herein, using an appropriate isotopically-labeled reagent in place of the non-labeled reagent otherwise employed. Such compounds have a variety of potential uses, e.g., as standards and reagents in determining the ability of a potential pharmaceutical compound to bind to target proteins or receptors, or for imaging compounds of this invention bound to biological receptors *in vivo* or *in vitro*.

20 "Stable compound" and "stable structure" are meant to indicate a compound that is sufficiently robust to survive isolation to a useful degree of purity from a reaction mixture, and formulation into an efficacious therapeutic agent. It is preferred that compounds of the present invention do not contain a N-halo, $\text{S}(\text{O})_2\text{H}$, or $\text{S}(\text{O})\text{H}$ group.

The term "solvate" means a physical association of a compound of this invention
25 with one or more solvent molecules, whether organic or inorganic. This physical association includes hydrogen bonding. The solvent molecules in the solvate may be present in a regular arrangement and/or a non-ordered arrangement. The solvate may comprise either a stoichiometric or nonstoichiometric amount of the solvent molecules. "Solvate" encompasses both solution-phase and isolable solvates. Exemplary solvates

include, but are not limited to, hydrates, ethanlates, methanlates, and isopropanolates. Methods of solvation are generally known in the art.

Abbreviations as used herein, are defined as follows: "1 x" for once, "2 x" for twice, "3 x" for thrice, "°C" for degrees Celsius, "eq" for equivalent or equivalents, "g" for gram or grams, "mg" for milligram or milligrams, "L" for liter or liters, "mL" for milliliter or milliliters, "μL" for microliter or microliters, "N" for normal, "M" for molar, "nM" for nanomolar, "mol" for mole or moles, "mmol" for millimole or millimoles, "min" for minute or minutes, "h" for hour or hours, "rt" for room temperature, "RT" for retention time, "atm" for atmosphere, "psi" for pounds per square inch, "conc." for concentrate, "sat" or "sat'd" for saturated, "MW" for molecular weight, "mp" for melting point, "MS" or "Mass Spec" for mass spectrometry, "ESI" for electrospray ionization mass spectroscopy, "HR" for high resolution, "HRMS" for high resolution mass spectrometry, "LCMS" for liquid chromatography mass spectrometry, "HPLC" for high pressure liquid chromatography, "RP HPLC" for reverse phase HPLC, "TLC" or "tlc" for thin layer chromatography, "NMR" for nuclear magnetic resonance spectroscopy, "nOe" for nuclear Overhauser effect spectroscopy, "¹H" for proton, "δ" for delta, "s" for singlet, "d" for doublet, "t" for triplet, "q" for quartet, "m" for multiplet, "br" for broad, "Hz" for hertz, and "α", "β", "R", "S", "E", and "Z" are stereochemical designations familiar to one skilled in the art.

The compounds of the present invention can be prepared as shown in the following reaction schemes and description thereof, as well as relevant published literature procedures that may be used by one skilled in the art. Exemplary reagents and procedures for these reactions appear hereinafter and in the working Examples.

ABBREVIATIONS

The following abbreviations are employed herein:

Bn = benzyl

t-Bu = tertiary butyl

Boc = *tert*-Butyloxycarbonyl

Boc₂O = di-*tert*-butyl dicarbonate

Cs₂CO₃ = cesium carbonate

DBU = 1,8-Diazabicyclo[5.4.0]undec-7-ene

DCM or CH₂Cl₂ = dichloromethane

DIAD = diisopropyl azodicarboxylate

Dess-Martin periodinane or DMP = 1,1,1-Triacetoxy-1,1-dihydro-1,2-benziodoxol-

5 3(1*H*)-one

DIPEA or *i*-Pr₂NEt = diisopropylethylamine

DMAP = 4-dimethylaminopyridine

DMF = dimethyl formamide

Et = ethyl

10 Et₃N = triethylamine

EtOH = ethanol

Et₂O = diethyl ether

EtOAc = ethyl acetate

HOAc or AcOH = acetic acid

15 K₂CO₃ = potassium carbonate

LiCl = lithium chloride

LiOAc = lithium acetate

LiOH = lithium hydroxide

Me = methyl

20 MeCN or ACN = acetonitrile

MeOH = methanol

MgSO₄ = magnesium sulfate

NaBH₄ = sodium borohydride

NaOH = sodium hydroxide

25 NaHCO₃ = sodium bicarbonate

PBu₃ = tributylphosphine

Ph = phenyl

Pd/C = palladium on carbon

Pd(OAc)₂ = palladium(II) acetate

30 Ph₃P = triphenylphosphine

PtO₂ = platinum dioxide

TBAF = tetra-*n*-butylammonium fluoride

TBDMS = tert-butyldimethylsilyl

TMS = trimethylsilyl

THF = tetrahydrofuran

5 TFA = trifluoroacetic acid

min = minute(s)

hr or hrs = hour(s)

L = liter

mL = milliliter

10 μ L = microliter

g = gram(s)

mg = milligram(s)

mol = moles

mmol = millimole(s)

15 meq = milliequivalent

sat or sat'd = saturated

aq. = aqueous

TLC = thin layer chromatography

HPLC = high performance liquid chromatography

20 LC/MS = high performance liquid chromatography/mass spectrometry

MS or Mass Spec = mass spectrometry

NMR = nuclear magnetic resonance

mp = melting point

25 IV. METHODS OF PREPARATION

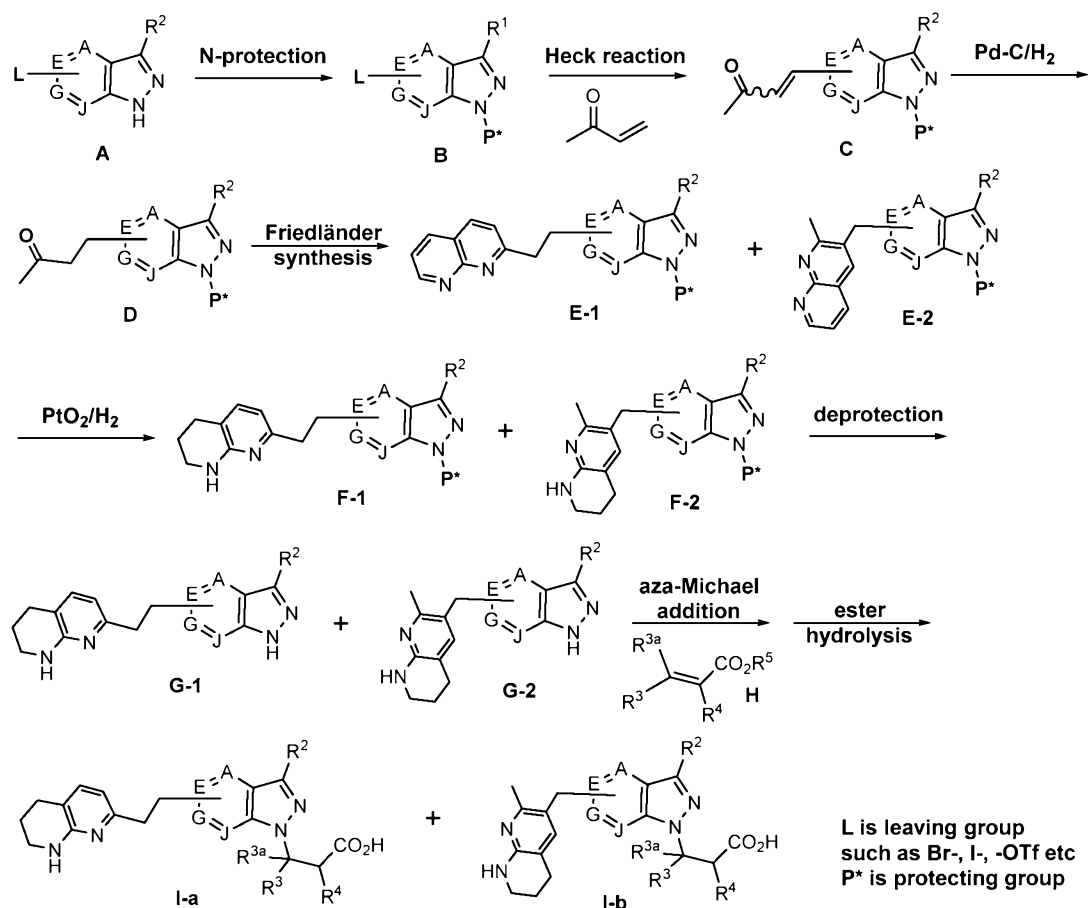
The compounds of Formula (I) may be prepared by the exemplary processes described in the following schemes and working examples, as well as relevant published literature procedures that are used by one skilled in the art. Exemplary reagents and procedures for these reactions appear hereinafter and in the working examples.

30 Protection and de-protection in the processes below may be carried out by procedures generally known in the art (see, for example, Wuts, P.G.M. et al., *Protecting Groups in*

Organic Synthesis, 4th Edition, Wiley (2007)). General methods of organic synthesis and functional group transformations are found in: Trost, B.M. et al., eds., *Comprehensive Organic Synthesis: Selectivity, Strategy & Efficiency in Modern Organic Chemistry*, Pergamon Press, New York, NY (1991); Smith, M.B. et al., *March's Advanced Organic Chemistry: Reactions, Mechanisms, and Structure*, 6th Edition, Wiley & Sons, New York, NY (2007); Katritzky, A.R. et al., eds., *Comprehensive Organic Functional Groups Transformations II*, 2nd Edition, Elsevier Science Inc., Tarrytown, NY (2004); Larock, R.C., *Comprehensive Organic Transformations*, VCH Publishers, Inc., New York, NY (1999), and references therein.

10 The novel compounds of Formula I may be prepared using the reactions and techniques described in this section. The reactions are performed in solvents appropriate to the reagents and materials employed and are suitable for the transformations being effected. Also, in the description of the synthetic methods described below, it is to be understood that all proposed reaction conditions, including solvent, reaction atmosphere, 15 reaction temperature, duration of the experiment and workup procedures, are chosen to be the conditions standard for that reaction, which should be readily recognized by one skilled in the art. One skilled in the art of organic synthesis understands that the functionality present on various portions of the molecule must be compatible with the reagents and reactions proposed. Not all compounds of Formula I falling into a given 20 class may be compatible with some of the reaction conditions required in some of the methods described. Such restrictions to the substituents, which are compatible with the reaction conditions, will be readily apparent to one skilled in the art and alternate methods must be used.

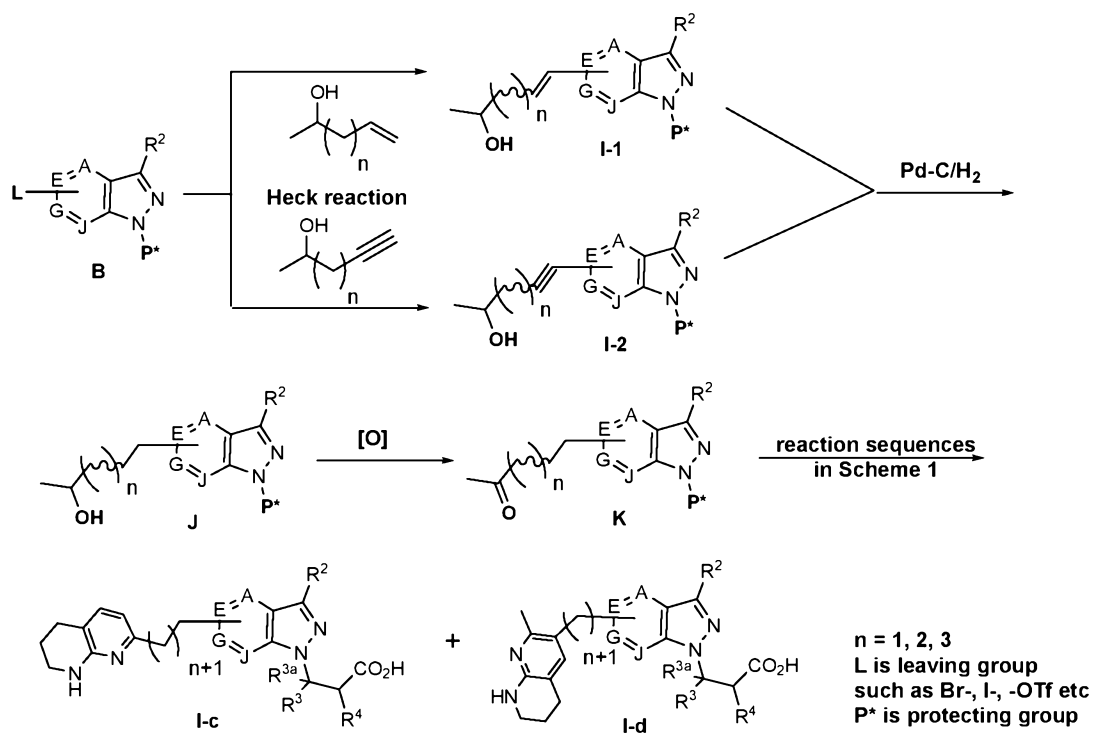
Scheme 1



Scheme 1 describes the preparation of formula I-a and formula I-b, subsets of formula (I). Starting material indazoles **A** bearing a leaving group **L** in various positions are commercially available or can be synthesized by the state of art following literature procedures. For a review of indazole synthesis, see: a) *Science of Synthesis: Houben-Weyl Methods of Molecular Transformations Vol. 12*, **2002**; b) Naděžda Cankařová, Jan Hlaváč, and Viktor Krchňák *Org. Prep. and Proc. Int.*, **2010**, 42, 433–465. A suitable protecting group **P*** (such as Boc-, Ar-SO₂-, MeOCH₂CH₂-, -SEM (2-trimethylsilyl ethoxy methyl), etc.) can be installed onto one of the indazole nitrogen to afford intermediate **B**. In scheme 1, **P*** group is shown at 1H- position as an example. Those skilled in the art will recognize that during the N- protection step, the **P*** group may be installed selectively or unselectively at either N1 or N2 of the 5-membered ring, and that this is inconsequential as this protecting group is removed in a later step. Heck reaction

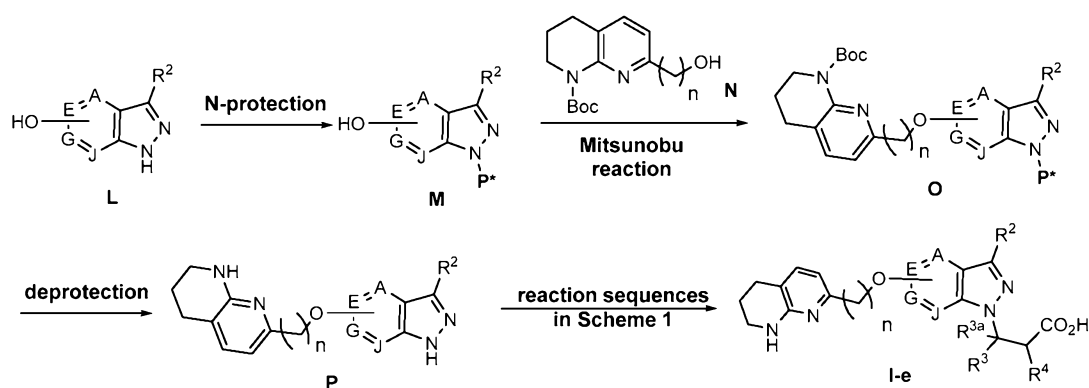
between **B** and methyl vinyl ketone (MVK) under Pd(OAc)₂-catalyzed conditions at elevated temperature affords intermediate **C**, which can be hydrogenated in the presence of Pd-C to intermediate **D**. Friedländer reaction between intermediate **D** and 2-aminonicotinaldehyde can be catalyzed by pyrrolidine or proline to afford intermediate **E-1** as a major product and **E-2** as a minor product. Hydrogenation of **E-1** and **E-2** in the presence of Adams's catalyst (PtO₂) can afford the corresponding tetrahydronaphthyridine **F-1** and **F-2**, where the protecting group P* can be removed to afford **G-1** and **G-2**. Indazoles are reported to be capable of aza-Michael addition to 2-propenamide, α,β -unsaturated ketone, or alkyl acrylate (for the references, see: a) Han, X. *Tetrahedron Lett.* **2007**, 48, 2845-2849; b) Yang, J. et al. *Synthesis* **2016**, 48, 1139-1146; c) An, Y.-L. et al. *Synthesis*, **2015**, 47, 1581-1592). However, no report has been documented about β -substituted α,β -unsaturated ester. We report here that the fully elaborated indazole intermediates such as **G-1** and **G-2** can react with various Michael addition acceptors such as intermediate **H** (where R³ and R^{3a} can be H or aryl or alkyl, R⁴ can be alkyl or N(Boc)₂). The typical reaction condition for such aza-Michael addition is mediated by base (such as DBU, K₂CO₃, Cs₂CO₃ etc) or Lewis acid (such as BF₃ etherate). Upon ester hydrolysis, formula I-a and I-b can be obtained after preparative HPLC purification or reversed phase chromatography purification.

Scheme 2



- Scheme 2 details the preparation of formula I-c and formula I-d, subsets of
 5 formula I. Under Heck reaction condition, intermediate **B** can react with a suitable
 terminal alkene or alkyne to form intermediate **I-1** and **I-2**. After hydrogenation, the
 secondary alcohol **J** can be oxidized to methyl ketone **K**. The transformation of **K** to
 formula I-c and I-d can be achieved by following the reaction sequences in Scheme 1.

Scheme 3

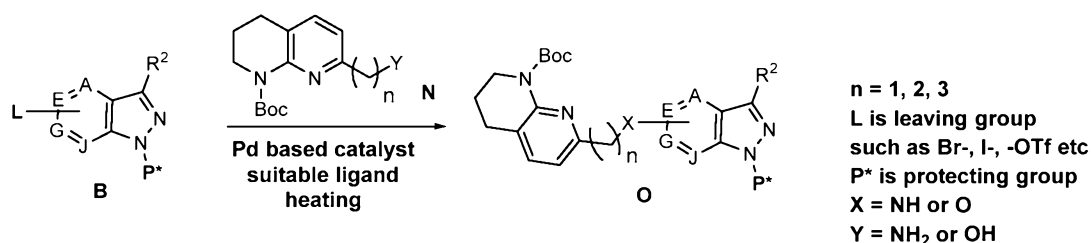


$n = 1, 2, 3$

P^* is protecting group, preferable to be Boc or Ar-SO_2 -

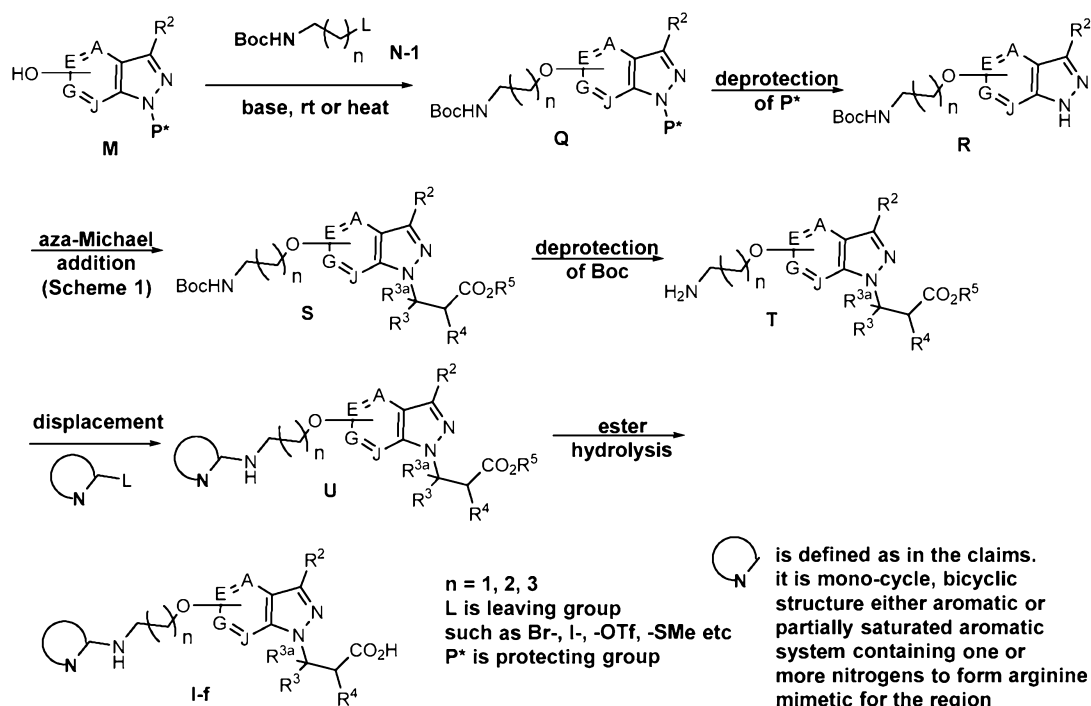
- Scheme 3 describes the preparation of formula I-e, a subset of formula I. The starting hydroxy bearing indazoles **L** are either commercially available or can be obtained
- 5 *via* indazole synthesis by following literature procedures. **L** can be converted to **M** *via* protecting group manipulation (to install a suitable protecting group such as Boc- or ArSO_2 -). The 3,4-dihydro-1,8-naphthyridine alcohols **N** are commercially available or can be synthesized by following literature procedures. Mitsunobu reaction between **M** and **N** can afford **O**, which can be further converted to **P** *via* cleavage of protecting group.
- 10 The key intermediate **P** can be then converted to formula I-e *via* the reaction sequence depicted in Scheme 1.

Scheme 4



- 15 Scheme 4 describes an alternative route to synthesize intermediate **O**. The reaction between **B** and **N** can be catalyzed by Pd, Cu, Ni catalysts and suitable ligands under elevated temperature. For a review of *O*-arylation, see: Muci, A. R.; Buchwald, S. L. *Topics in Current Chemistry* **2002**, 219 (Cross-Coupling Reactions), 131-209.

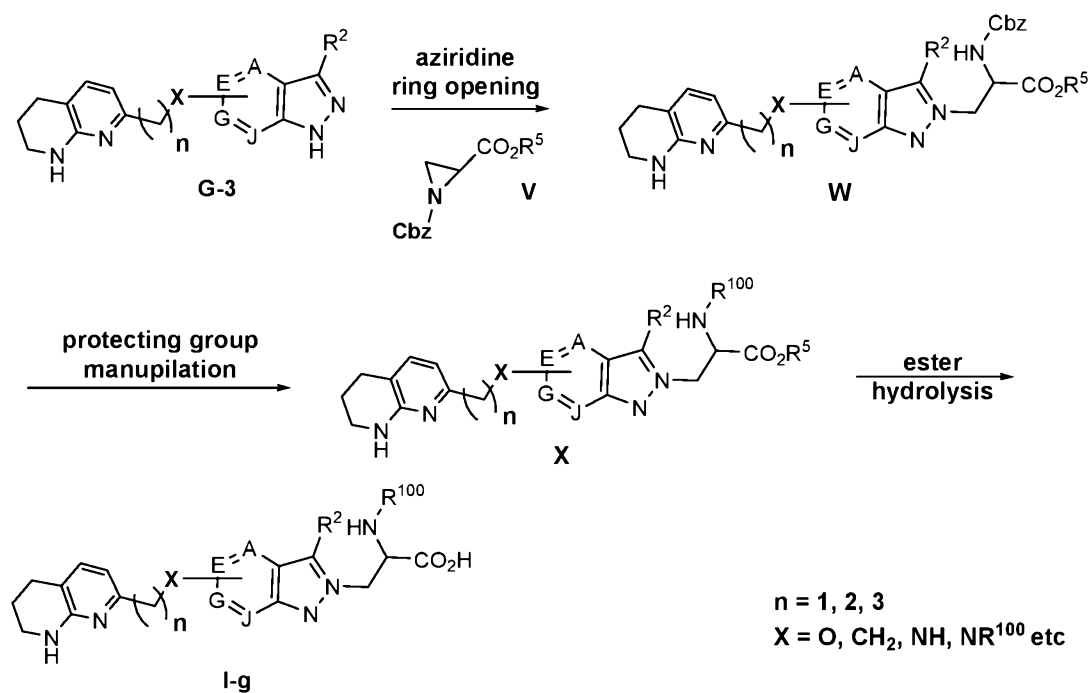
Scheme 5



Scheme 5 describes the preparation of formula I-f, a subset of formula I.

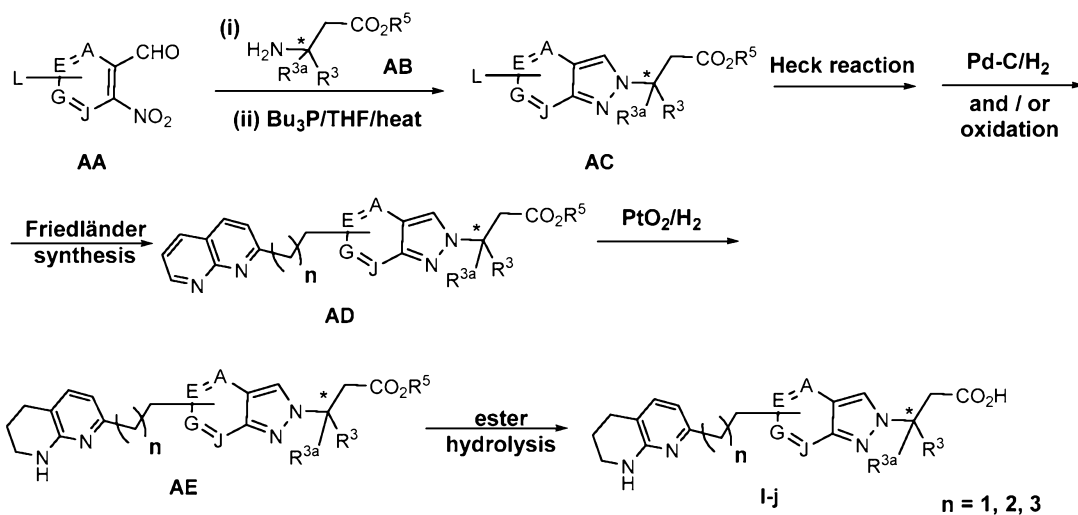
- 5 Displacement reaction of **M** with intermediate **N-1** can be carried out in a suitable solvent in the presence of a base such as K_2CO_3 or Cs_2CO_3 at room temperature or elevated temperature to afford **Q**. Upon selective deprotection of P^* , intermediate **R** can be
- undergo aza-Michael reaction with various acceptors (see Scheme 1) to form **S**. The Boc-protecting group of **S** can be removed under acidic conditions (for example TFA or HCl
- 10 in dioxane) to afford **T**, which reacts with various mono- or bicyclic aromatic or partial aromatic ring systems bearing a leaving group L to form intermediate **U**. Upon ester hydrolysis under aqueous basic condition, formula I-f can be obtained.

Scheme 6



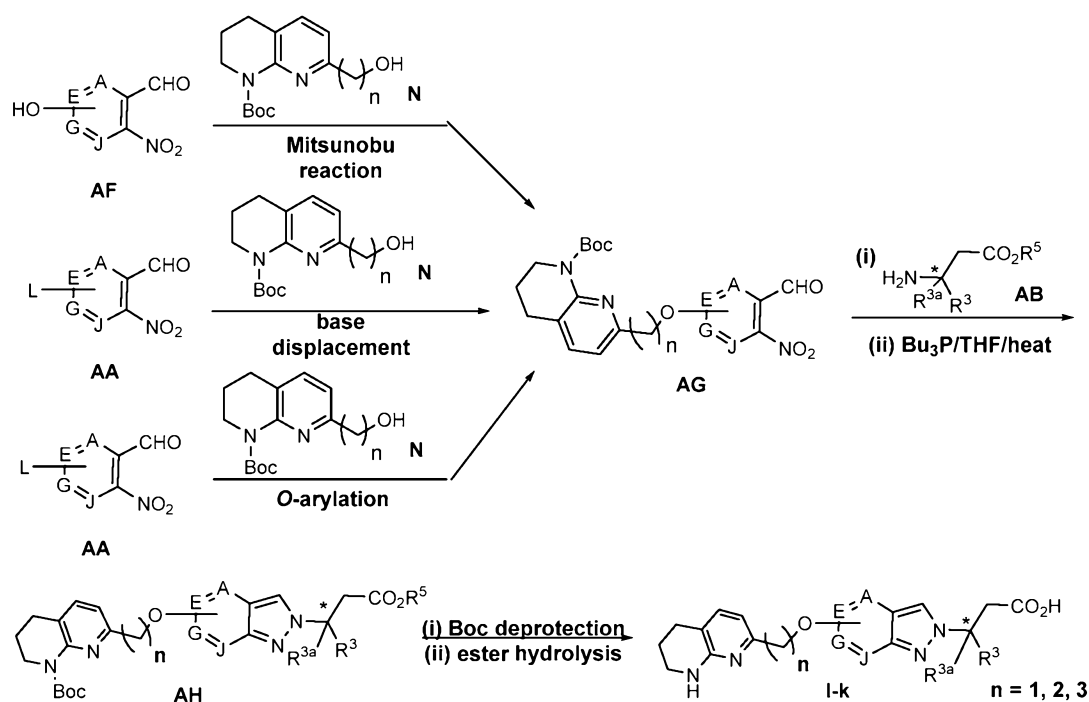
- Scheme 6 describes the preparation of formula I-g, a subset of formula I. The key intermediate **G-3** can be the intermediate **G-1** (from Scheme 1) or any similar
- 5 intermediate from Scheme 2 to Scheme 4. The aziridine ring opening reaction between **G-3** and **V** can be performed selectively in the presence of Lewis acid such as TFA under elevated temperature to afford the 2*H*-indazole **W** as the major product. **W** can then be converted to intermediate **X** ($R^{100} = \text{H}$), which can be further converted to **X** bearing other R^{100} groups (such as $R^{100} = R^{101}\text{SO}_2-$, $R^{101}\text{C(O)}-$, etc; wherein R^{101} can be optionally
- 10 substituted alkyl, aryl, or arylalkyl). Upon hydrolysis in the presence of a base, formula I-g can be obtained.

Scheme 7



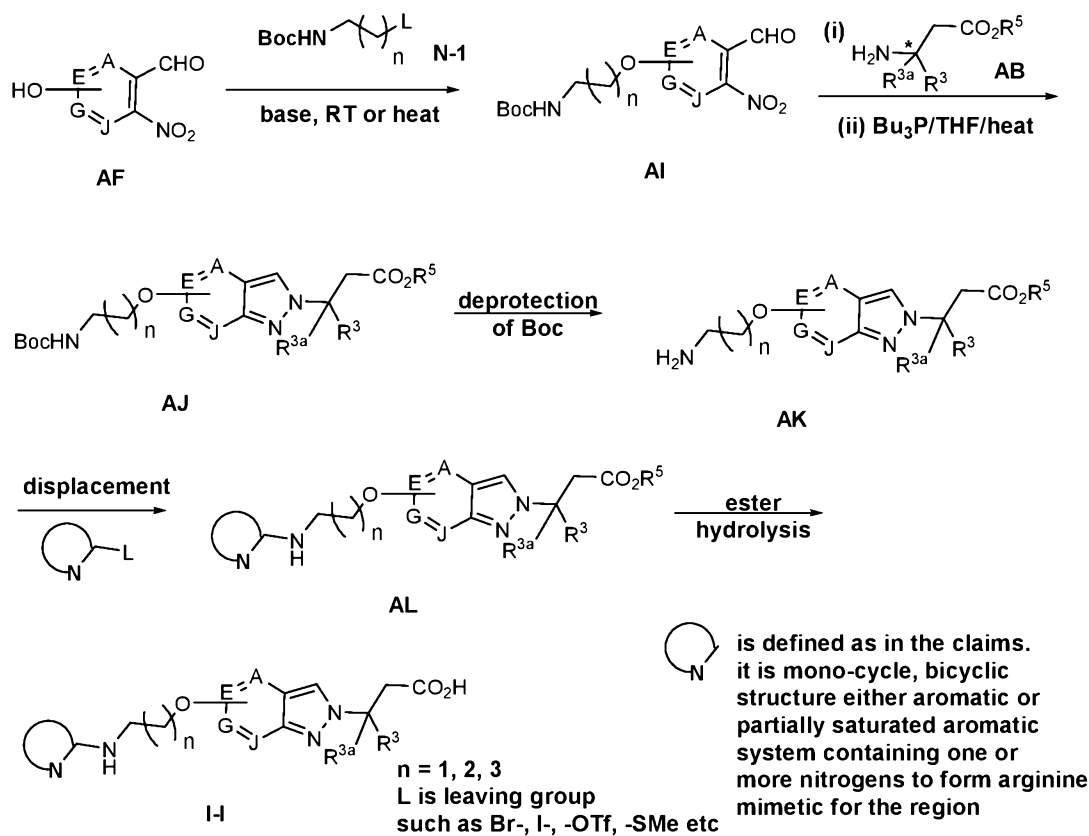
Scheme 7 describes the preparation of formula I-j, a subset of formula I. 2-Nitrobenzaldehyde analogs **AA** are commercially available or can be synthesized using literature procedures. Intermediate **AB** (either chiral or racemic) can be obtained from commercial sources or from synthesis following literature procedures. **AA** can react with **AB** under thermal conditions to form an imine, which can be reduced by Bu₃P in one pot to afford ring-closed product 2H-indazole **AC**. **AC** can be transformed to formula I-j following a sequence analogous to the one described in Scheme 1.

Scheme 8



- Scheme 8 describes the preparation of formula I-k, a subset of formula I. There are three different methods to access intermediate AG. Method 1 is the Mitsunobu reaction between AF and N; Method 2 involves in the base-mediated S_NA2 displacement reaction between hydroxy group of N and leaving group L in AA; Method 3 is the typical O-arylation where hydroxy group of N reacts with AA in the presence of Pd-based catalyst and suitable ligand. The ring closure reaction between AB and AG can occur as described in Scheme 7 to afford AH. Upon Boc deprotection and ester hydrolysis,
- 10 formula I-k can be obtained.

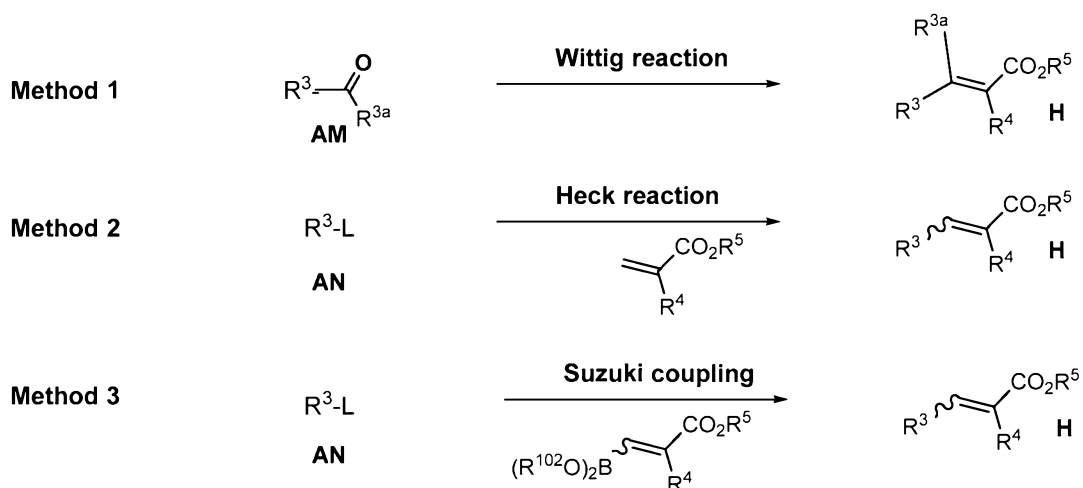
Scheme 9



Scheme 9 describes the preparation of formula I-l, a subset of formula I.

Intermediate **AF** can react with **N-1** under basic condition to form **AI**, which can be subsequently cyclized with **AB** to form **AJ** as described in Scheme 7. The Boc protecting group can be removed under acidic conditions (TFA or HCl in dioxane) to afford **AK**. The transformation of **AK** to formula I-l can occur following a sequence familiar to the one shown in Scheme 5.

Scheme 10

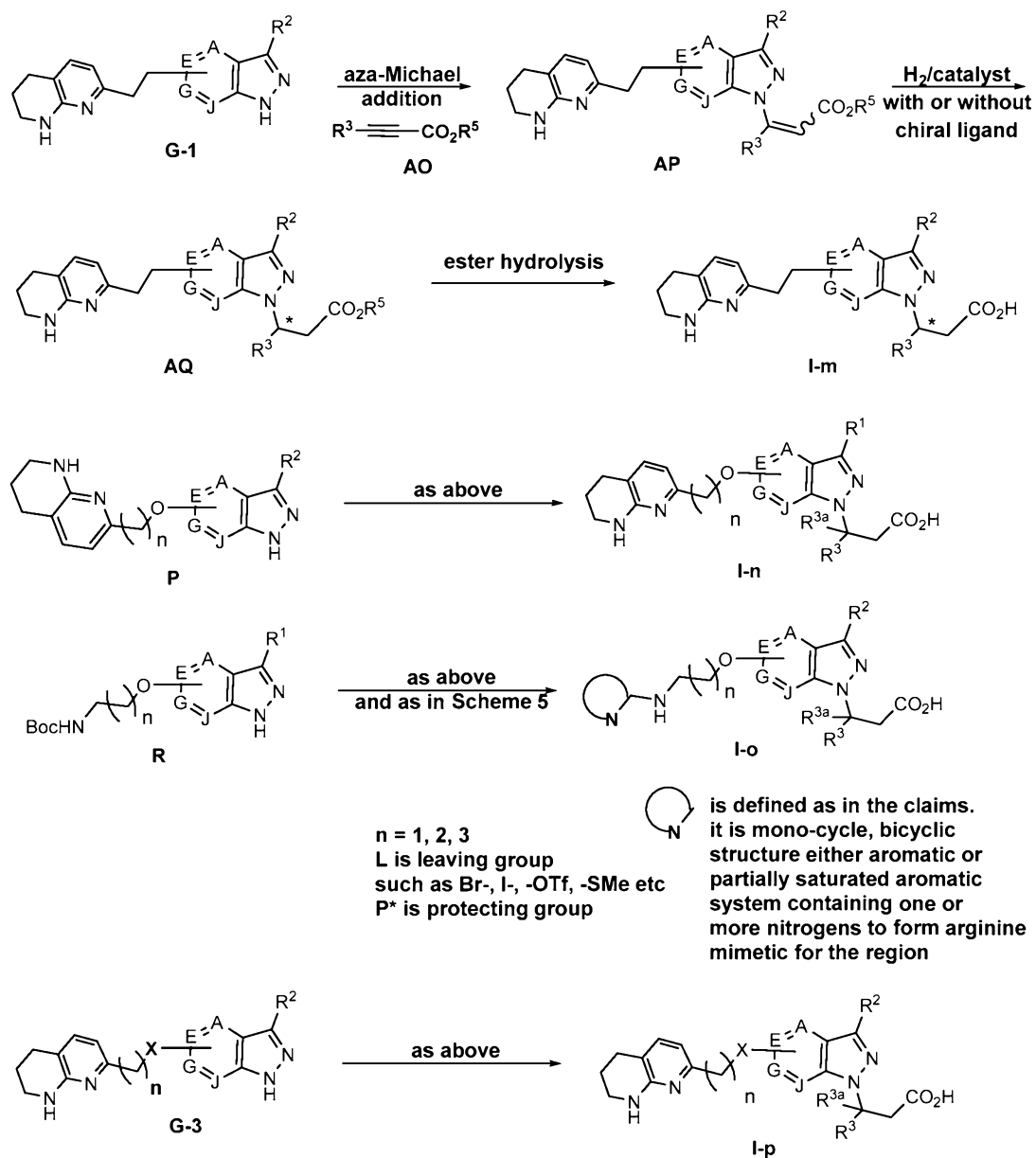


L is leaving group, such as Br, I, Cl, OTf, etc.

R¹⁰² is H, alkyl, aryl, or arylalkyl.

- Scheme 10 describes the synthesis of intermediate **H**, which is used in the above synthetic schemes. For example, appropriate aldehyde **AM** can react with a suitable Wittig reagent to form intermediate **H** (Method 1). Alternatively, **AN** can react with a suitable olefin under Heck reaction conditions (Method 2) or Suzuki coupling reaction conditions (Method 3) to provide intermediate **H**. For Method 2 and 3, **R**³ is typically aryl or heteroaryl, rather than alkyl (less common). However, Method 1 can be used when **R**³ or **R**^{3a} is either alkyl or aryl or heteroaryl.

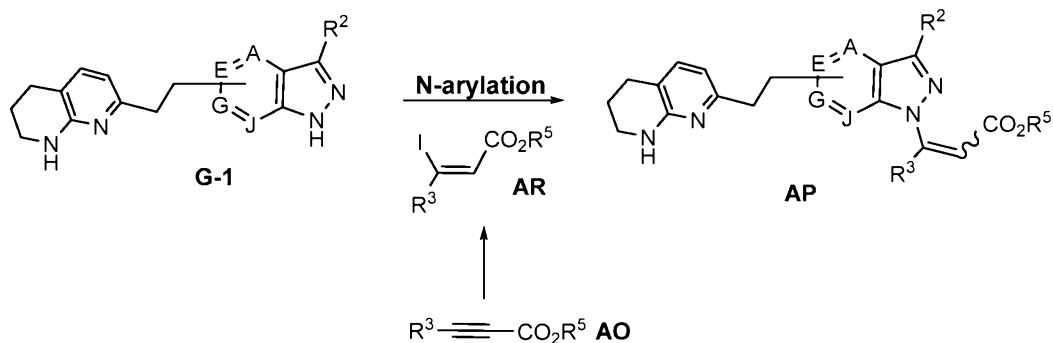
Scheme 11



Scheme 11 describes the preparation of formula I-m, I-n, I-o, and I-p, subsets of
 5 formula I. Intermediate **G-1** (from Scheme 5) can react with alkynyl ester **AO**
 (commercially available or can be synthesized by following literature procedures) to form
 Michael addition adduct **AP**. Upon the hydrogenation of **AP** under Pd catalysis (with or
 without chiral ligand), intermediate **AQ** can be obtained in chiral or racemic form. Ester

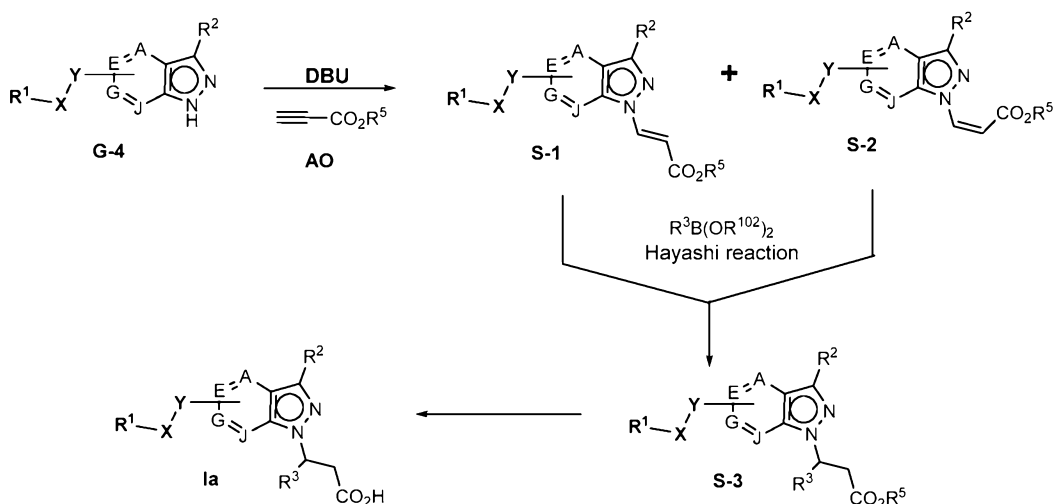
hydrolysis of **AQ** can afford formula I-m. Similarly, intermediates **P**, **R**, **G-3** can undergo the above sequence to afford formula I-n, I-o, and I-p respectively.

Scheme 12



Alternatively, intermediate **AP** can be prepared using N-arylation method detailed in Scheme 12. Thus, alkynyl ester **AO** can be converted to **AR** via treatment with NaI in acetic acid. The reaction between **G-1** and **AR** can be achieved using standard Buckwald N-arylation reaction conditions (see: *PNAS*, **2004**, *101*, 5821–5823).

Scheme 13



10

Scheme 13 describes the preparation of formula Ia using an alternative route. Intermediate **G-4** (see above schemes for preparation) can undergo the aza-Michael addition with alkyl propiolate **AO** to form *trans*-adduct **S-1** (major) and *cis*-adduct **S-2**. The Rh(I)-mediated Hayashi reaction of **S-1** or **S-2** or the mixture with $\text{R}^3\text{B(OR}^{102}\text{)}_2$ can afford **S-3**, which can be further converted to formula Ia upon ester hydrolysis.

15

[0001]

V. EXAMPLES

5 The following Examples are offered as illustrative, as a partial scope and particular embodiments of the invention and are not meant to be limiting of the scope of the invention. Abbreviations and chemical symbols have their usual and customary meanings unless otherwise indicated. Unless otherwise indicated, the compounds described herein have been prepared, isolated and characterized using the schemes and other methods disclosed herein or may be prepared using the same.

10 As appropriate, reactions were conducted under an atmosphere of dry nitrogen (or argon). For anhydrous reactions, DRISOLV® solvents from EM were employed. For other reactions, reagent grade or HPLC grade solvents were utilized. Unless otherwise stated, all commercially obtained reagents were used as received.

15 HPLC/MS AND PREPARATORY/ANALYTICAL HPLC METHODS EMPLOYED IN CHARACTERIZATION OR PURIFICATION OF EXAMPLES

NMR (nuclear magnetic resonance) spectra were typically obtained on Bruker or JEOL 400 MHz and 500 MHz instruments in the indicated solvents. All chemical shifts are reported in ppm from tetramethylsilane with the solvent resonance as the internal
20 standard. ¹HNMR spectral data are typically reported as follows: chemical shift, multiplicity (s = singlet, br s = broad singlet, d = doublet, dd = doublet of doublets, t = triplet, q = quartet, sep = septet, m = multiplet, app = apparent), coupling constants (Hz), and integration.

25 The term HPLC refers to a Shimadzu high performance liquid chromatography instrument with one of following methods:

HPLC-1: Sunfire C18 column (4.6 × 150 mm) 3.5 μm, gradient from 10 to 100% B:A for 12 min, then 3 min hold at 100% B.

Mobile phase A: 0.05% TFA in water:CH₃CN (95:5)

30 Mobile phase B: 0.05% TFA in CH₃CN:water (95:5)

TFA Buffer pH = 2.5; Flow rate: 1 mL/ min; Wavelength: 254 nm, 220 nm.

HPLC-2: XBridge Phenyl (4.6 × 150 mm) 3.5 μm, gradient from 10 to 100% B:A for 12 min, then 3 min hold at 100% B.

Mobile phase A: 0.05% TFA in water:CH₃CN (95:5)

5 Mobile phase B: 0.05% TFA in CH₃CN:water (95:5)

TFA Buffer pH = 2.5; Flow rate: 1 mL/ min; Wavelength: 254 nm, 220 nm.

HPLC-3: Chiralpak AD-H, 4.6 × 250 mm, 5 μm.

Mobile Phase: 30% EtOH-heptane (1:1) / 70% CO₂

10 Flow rate = 40 mL/min, 100 Bar, 35 °C; Wavelength: 220 nm

HPLC-4: Waters Acquity UPLC BEH C18, 2.1 x 50 mm, 1.7-μm particles;

Mobile Phase A: 5:95 CH₃CN:water with 10 mM NH₄OAc;

Mobile Phase B: 95:5 CH₃CN:water with 10 mM NH₄OAc;

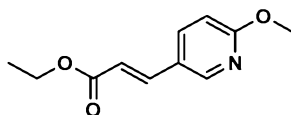
15 Temperature: 50 °C; Gradient: 0-100% B over 3 min, then a 0.75-min hold at 100% B;
Flow: 1.11 mL/min; Detection: UV at 220 nm.

HPLC-5: Waters Acquity UPLC BEH C18, 2.1 x 50 mm, 1.7-μm particles;

Mobile Phase A: 5:95 CH₃CN:water with 0.1% TFA;

20 Mobile Phase B: 95:5 CH₃CN:water with 0.1% TFA;

Temperature: 50 °C; Gradient: 0-100% B over 3 min, then a 0.75-min hold at 100% B; Flow: 1.11 mL/min; Detection: UV at 220 nm.



Intermediate 1A

25 Intermediate **1A** can be synthesized using three different methods: Heck reaction, Wittig reaction, or Suzuki coupling reaction. The following procedures serve as the examples for all acrylates synthesized and used in this application.

Method 1. To a solution of 6-methoxynicotinaldehyde (3 g, 21.88 mmol) in THF (45 mL) was added 15 g of Molecular sieves 4Å followed by ethyl 2-(diethoxyphosphoryl)acetate (5.25 mL, 26.3 mmol) and LiOH (0.629 g, 26.3 mmol). The reaction was stirred at rt overnight. The reaction mixture was filtered through a pad of celite and the volatiles were removed under reduced pressure. The residue was dissolved in EtOAc (25 mL) and washed with 10% NaHCO₃ (aqueous, 12 mL) followed by brine (12 mL). The organic layer was dried (Na₂SO₄), filtered, and concentrated under reduced pressure. The residue was purified *via* flash column chromatography (220 g silica gel, 0 to 20% hexane/ethyl acetate) to afford Intermediate **1A** (4.5 g, 21.72 mmol, 99% yield) as a yellow solid. ¹H NMR (500MHz, CDCl₃) δ 8.28 (d, *J* = 2.5 Hz, 1H), 7.77 (dd, *J* = 8.8, 2.5 Hz, 1H), 7.64 (d, *J* = 16.2 Hz, 1H), 6.77 (d, *J* = 8.8 Hz, 1H), 6.34 (d, *J* = 16.0 Hz, 1H), 4.27 (q, *J* = 7.2 Hz, 2H), 3.98 (s, 3H), 1.34 (t, *J* = 7.2 Hz, 3H). LCMS (ES): *m/z* 208.1 [M+H]⁺.

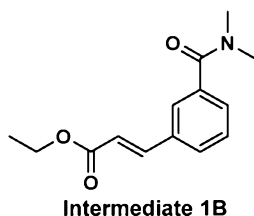
Method 2. A solution of 5-bromo-2-methoxypyridine (1.03 mL, 7.98 mmol), ethyl acrylate (3.0 mL, 27.9 mmol), Et₃N (3.0 mL, 21.54 mmol), Pd(OAc)₂ (0.202 g, 0.899 mmol) and tri-*o*-tolylphosphine (0.404 g, 1.327 mmol) in ACN (2.0 mL) was degassed with argon for 10 min. The mixture was heated at 90 °C for 12 h. The solvent was removed under reduced pressure. Toluene (1.5 mL) was added and the mixture was concentrated again under reduced pressure. Ether (10 mL) was added and the mixture was filtered through a pad of silica gel eluting with ether. The solvent was removed and the residue was purified *via* flash chromatography (80 g silica gel, 0 to 100% hexanes/ethyl acetate) to afford Intermediate **1A** (1.59 g, 7.67 mmol, 96% yield) as a yellow solid.

Method 3. To a degassed solution of 5-bromo-2-methoxypyridine (2 g, 10.64 mmol), (*E*)-ethyl 3-(4,4,5,5-tetramethyl-1,3,2-dioxaborolan-2-yl)acrylate (2.405 g, 10.64 mmol) and K₂CO₃ (4.41 g, 31.9 mmol) in 1,4-dioxane (30 mL) and water (10 mL) was added Pd(PPh₃)₄ (0.492 g, 0.425 mmol). The reaction mixture was stirred in a seal vial at 100 °C overnight. After cooled to rt, the mixture was diluted with water (15 mL), and extracted with CH₂Cl₂ (3 x 10 mL), the combine organics were dried (Na₂SO₄), filtered, and concentrated. the residue was purified *via* flash chromatography (80 g silica gel, 0 to

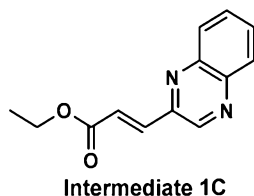
100% hexane/ethyl acetate) to afford Intermediate **1A** (662 mg, 3.19 mmol, 30% yield) as a yellow solid.

The following intermediates can be synthesized using one of the methods above or the procedures in the literature known to the one skilled of the art.

5

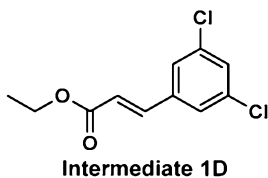


Intermediate **1B**: ^1H NMR (400 MHz, chloroform-*d*) δ 7.70 (dt, $J = 16.0, 0.5$ Hz, 1H), 7.62 – 7.54 (m, 2H), 7.50 – 7.38 (m, 2H), 6.48 (d, $J = 16.0$ Hz, 1H), 4.29 (q, $J = 7.1$ Hz, 2H), 3.15 (s, 3H), 3.01 (s, 3H), 1.36 (t, $J = 7.1$ Hz, 3H). LCMS (ES): m/z 248.2

10 $[\text{M}+\text{H}]^+$.

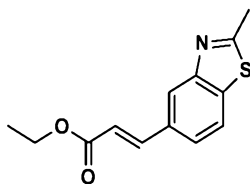
Intermediate **1C**: ^1H NMR (500 MHz, chloroform-*d*) δ 8.94 (s, 1H), 8.08 – 7.99 (m, 2H), 7.83 (d, $J = 15.9$ Hz, 1H), 7.73 (td, $J = 6.3, 5.8, 3.3$ Hz, 2H), 7.08 (d, $J = 15.9$ Hz, 1H), 4.26 (q, $J = 7.1$ Hz, 2H), 1.31 (t, $J = 7.1$ Hz, 3H). LCMS (ES): m/z 229.1

15 $[\text{M}+\text{H}]^+$.



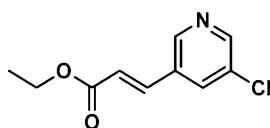
Intermediate **1D**: ^1H NMR (400 MHz, chloroform-*d*) δ 7.64 – 7.51 (d, $J = 16.0$ Hz, 1H), 7.43 – 7.33 (m, 3H), 6.46 (d, $J = 16.0$ Hz, 1H), 4.29 (q, $J = 7.2$ Hz, 2H), 1.36 (t, $J = 7.1$ Hz, 3H). LCMS (ES): m/z 245.1 $[\text{M}+\text{H}]^+$.

20

**Intermediate 1E**

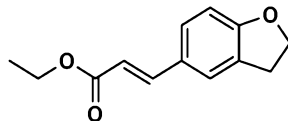
Intermediate **1E**: ^1H NMR (500 MHz, chloroform-*d*) δ 8.10 (d, $J = 1.7$ Hz, 1H), 7.89 – 7.80 (m, 2H), 7.56 (dd, $J = 8.4, 1.6$ Hz, 1H), 6.54 (d, $J = 16.0$ Hz, 1H), 4.31 (q, $J = 7.1$ Hz, 2H), 2.88 (s, 3H), 1.38 (t, $J = 7.1$ Hz, 3H). LCMS (ES): m/z 248.0 $[\text{M}+\text{H}]^+$.

5

**Intermediate 1F**

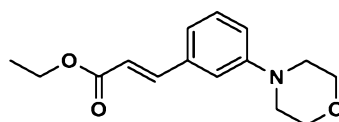
Intermediate **1F**: ^1H NMR (400 MHz, chloroform-*d*) δ 8.61 (dd, $J = 19.0, 2.0$ Hz, 2H), 7.84 (t, $J = 2.1$ Hz, 1H), 7.64 (d, $J = 16.1$ Hz, 1H), 6.54 (d, $J = 16.1$ Hz, 1H), 4.31 (q, $J = 7.2$ Hz, 2H), 1.37 (t, $J = 7.1$ Hz, 3H). LCMS (ES): m/z 211.9 $[\text{M}+\text{H}]^+$.

10

**Intermediate 1G**

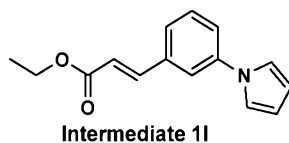
Intermediate **1G**: ^1H NMR (400 MHz, chloroform-*d*) δ 7.66 (d, $J = 16.0$ Hz, 1H), 7.43 (d, $J = 1.7$ Hz, 1H), 7.33 (dd, $J = 8.6, 1.8$ Hz, 1H), 6.81 (d, $J = 8.3$ Hz, 1H), 6.30 (d, $J = 16.0$ Hz, 1H), 4.65 (t, $J = 8.7$ Hz, 2H), 4.28 (q, $J = 7.1$ Hz, 2H), 3.25 (t, $J = 8.7$ Hz, 2H), 1.36 (t, $J = 7.1$ Hz, 3H). LCMS (ES): m/z 219.1 $[\text{M}+\text{H}]^+$.

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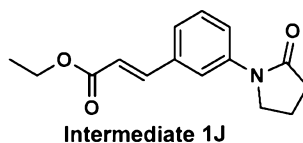
**Intermediate 1H**

Intermediate **1H**: ^1H NMR (500 MHz, chloroform-*d*) δ 7.68 (d, $J = 16.1$ Hz, 1H), 7.31 (t, $J = 7.9$ Hz, 1H), 7.08 (dt, $J = 7.6, 1.1$ Hz, 1H), 7.06 (t, $J = 2.1$ Hz, 1H), 6.97 (dd, J

= 8.1, 2.5 Hz, 1H), 6.44 (d, $J = 16.0$ Hz, 1H), 4.29 (q, $J = 7.1$ Hz, 2H), 3.93 – 3.82 (m, 4H), 3.25 – 3.13 (m, 4H), 1.36 (t, $J = 7.1$ Hz, 3H). LCMS (ES): m/z 262.0 $[M+H]^+$.



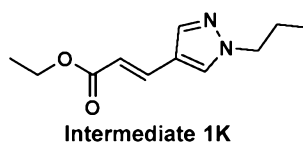
5 Intermediate **1I**: ^1H NMR (500 MHz, chloroform- d) δ 7.73 (d, $J = 16.0$ Hz, 1H), 7.56 (t, $J = 1.8$ Hz, 1H), 7.51 – 7.39 (m, 3H), 7.13 (t, $J = 2.2$ Hz, 2H), 6.51 (d, $J = 16.0$ Hz, 1H), 6.40 (t, $J = 2.1$ Hz, 2H), 4.31 (q, $J = 7.1$ Hz, 2H), 1.38 (t, $J = 7.1$ Hz, 3H). LCMS (ES): m/z 242.1 $[M+H]^+$.



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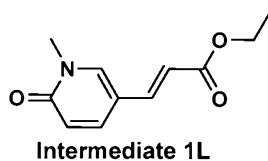
Intermediate **1J**: ^1H NMR (500 MHz, chloroform- d) δ 7.82 (t, $J = 2.0$ Hz, 1H), 7.74 – 7.66 (m, 2H), 7.41 (t, $J = 7.9$ Hz, 1H), 7.34 (dt, $J = 7.7, 1.3$ Hz, 1H), 6.48 (d, $J = 16.0$ Hz, 1H), 4.29 (q, $J = 7.1$ Hz, 2H), 3.91 (t, $J = 7.1$ Hz, 2H), 2.66 (t, $J = 8.1$ Hz, 2H), 2.28 – 2.14 (m, 2H), 1.37 (t, $J = 7.1$ Hz, 3H). LCMS (ES): m/z 260.1 $[M+H]^+$.

15



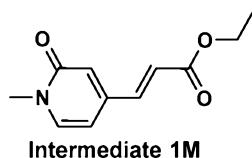
Intermediate **1K**: ^1H NMR (500 MHz, chloroform- d) δ 7.72 (s, 1H), 7.62 – 7.54 (m, 2H), 6.18 (d, $J = 16.0$ Hz, 1H), 4.25 (q, $J = 7.1$ Hz, 2H), 4.10 (t, $J = 7.0$ Hz, 2H), 1.92 (h, $J = 7.3$ Hz, 2H), 1.34 (t, $J = 7.1$ Hz, 3H), 0.95 (t, $J = 7.4$ Hz, 3H). LCMS (ES): m/z 209.2 $[M+H]^+$.

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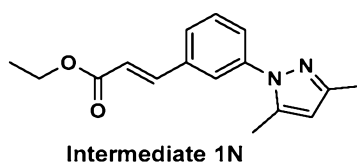
Intermediate **1L**: ^1H NMR (500 MHz, chloroform-*d*) δ 7.61 (dd, $J = 9.6, 2.5$ Hz, 1H), 7.47 (d, $J = 2.5$ Hz, 1H), 7.42 (d, $J = 15.9$ Hz, 1H), 6.64 (d, $J = 9.5$ Hz, 1H), 6.17 (d, $J = 15.8$ Hz, 1H), 4.27 (q, $J = 7.1$ Hz, 2H), 3.60 (s, 3H), 1.35 (t, $J = 7.1$ Hz, 3H). LCMS (ES): m/z 208.1 $[\text{M}+\text{H}]^+$.

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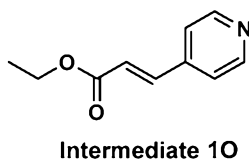
Intermediate **1M**: ^1H NMR (500 MHz, chloroform-*d*) δ 7.44 (d, $J = 15.9$ Hz, 1H), 7.30 (d, $J = 9.1$ Hz, 1H), 6.66 (d, $J = 1.8$ Hz, 1H), 6.45 (d, $J = 15.9$ Hz, 1H), 6.31 (dd, $J = 7.1, 2.0$ Hz, 1H), 4.30 (q, $J = 7.1$ Hz, 2H), 3.57 (s, 3H), 1.36 (t, $J = 7.1$ Hz, 3H). LCMS (ES): m/z 208.1 $[\text{M}+\text{H}]^+$.

10



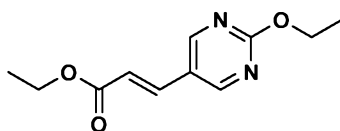
Intermediate **1N**: ^1H NMR (500 MHz, chloroform-*d*) δ 7.72 (d, $J = 16.0$ Hz, 1H), 7.64 (d, $J = 2.2$ Hz, 1H), 7.54 – 7.44 (m, 3H), 6.50 (d, $J = 16.0$ Hz, 1H), 6.04 (s, 1H), 4.29 (q, $J = 7.1$ Hz, 2H), 2.35 (s, 3H), 2.33 (s, 3H), 1.36 (t, $J = 7.1$ Hz, 3H). LCMS (ES): m/z 271.1 $[\text{M}+\text{H}]^+$.

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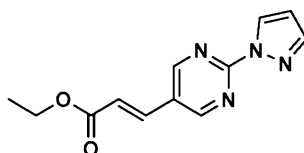
Intermediate **1O**: ^1H NMR (500 MHz, chloroform-*d*) δ 8.71 – 8.65 (m, 2H), 7.62 (d, $J = 16.0$ Hz, 1H), 7.41 – 7.35 (m, 2H), 6.62 (d, $J = 16.0$ Hz, 1H), 4.32 (q, $J = 7.1$ Hz, 2H), 1.38 (t, $J = 7.1$ Hz, 3H). LCMS (ES): m/z 178.2 $[\text{M}+\text{H}]^+$.

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**Intermediate 1P**

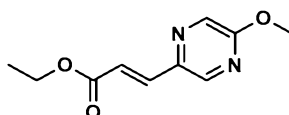
Intermediate **1P**: ^1H NMR (500 MHz, chloroform-*d*) δ 8.68 (s, 2H), 7.59 (d, J = 16.1 Hz, 1H), 6.46 (d, J = 16.2 Hz, 1H), 4.50 (q, J = 7.1 Hz, 2H), 4.30 (q, J = 7.1 Hz, 2H), 1.47 (t, J = 7.0 Hz, 3H), 1.36 (t, J = 7.1 Hz, 3H). LCMS (ES): m/z 223.2 $[\text{M}+\text{H}]^+$.

5

**Intermediate 1Q**

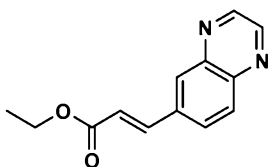
Intermediate **1Q**: ^1H NMR (500 MHz, chloroform-*d*) δ 8.90 (s, 2H), 8.63 (d, J = 2.6 Hz, 1H), 7.89 (d, J = 1.4 Hz, 1H), 7.66 (d, J = 16.2 Hz, 1H), 6.60 (d, J = 16.1 Hz, 1H), 6.56 (dd, J = 2.7, 1.6 Hz, 1H), 4.33 (q, J = 7.1 Hz, 2H), 1.38 (t, J = 7.1 Hz, 3H).

10 LCMS (ES): m/z 245.1 $[\text{M}+\text{H}]^+$.

**Intermediate 1R**

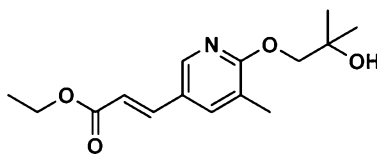
Intermediate **1R**: ^1H NMR (500 MHz, chloroform-*d*) δ 8.26 (d, J = 1.3 Hz, 1H), 8.18 (d, J = 1.3 Hz, 1H), 7.67 (d, J = 15.7 Hz, 1H), 6.86 (d, J = 15.6 Hz, 1H), 4.30 (q, J = 7.1 Hz, 2H), 4.03 (s, 3H), 1.36 (t, J = 7.1 Hz, 3H). LCMS (ES): m/z 209.1 $[\text{M}+\text{H}]^+$.

15

**Intermediate 1S**

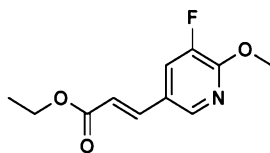
Intermediate **1S**: ^1H NMR (500 MHz, chloroform-*d*) δ 8.90 (d, J = 1.9 Hz, 1H), 8.88 (d, J = 1.8 Hz, 1H), 8.23 (d, J = 2.0 Hz, 1H), 8.14 (d, J = 8.7 Hz, 1H), 7.99 (dd, J =

8.7, 2.0 Hz, 1H), 7.91 (d, $J = 16.0$ Hz, 1H), 6.67 (d, $J = 16.0$ Hz, 1H), 4.34 (q, $J = 7.1$ Hz, 2H), 1.40 (t, $J = 7.1$ Hz, 3H). LCMS (ES): m/z 229.2 $[M+H]^+$.



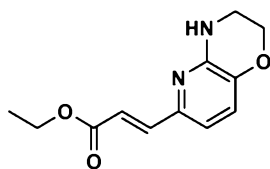
Intermediate 1T

5 Intermediate **1T**: ^1H NMR (500 MHz, chloroform- d) δ 7.53 (dd, $J = 2.6, 1.3$ Hz, 1H), 7.47 – 7.37 (m, 2H), 6.19 (d, $J = 15.9$ Hz, 1H), 4.27 (q, $J = 7.1$ Hz, 2H), 4.07 (s, 2H), 3.59 (s, 1H), 2.22 (d, $J = 1.2$ Hz, 3H), 1.35 (t, $J = 7.1$ Hz, 3H), 1.30 (s, 6H). LCMS (ES): m/z 280.2 $[M+H]^+$.



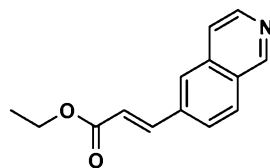
Intermediate 1U

10 Intermediate **1U**: ^1H NMR (500 MHz, chloroform- d) δ 8.07 (d, $J = 2.1$ Hz, 1H), 7.64 (dd, $J = 16.0, 1.6$ Hz, 1H), 7.53 (dd, $J = 10.8, 2.1$ Hz, 1H), 6.34 (d, $J = 16.0$ Hz, 1H), 4.30 (q, $J = 7.1$ Hz, 2H), 4.09 (s, 3H), 1.37 (t, $J = 7.1$ Hz, 3H). LCMS (ES): m/z 226.2 $[M+H]^+$.

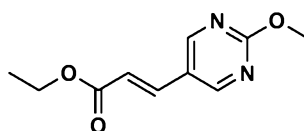


Intermediate 1V

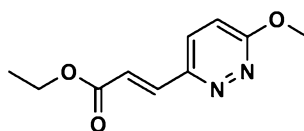
15 Intermediate **1V**: ^1H NMR (500 MHz, chloroform- d) δ 7.49 (d, $J = 15.5$ Hz, 1H), 6.96 (d, $J = 7.9$ Hz, 1H), 6.74 (d, $J = 7.9$ Hz, 1H), 6.68 (d, $J = 15.6$ Hz, 1H), 4.33 – 4.20 (m, 5H), 3.59 (td, $J = 4.5, 2.3$ Hz, 2H), 1.34 (t, $J = 7.1$ Hz, 3H). LCMS (ES): m/z 235.2
20 $[M+H]^+$.

**Intermediate 1W**

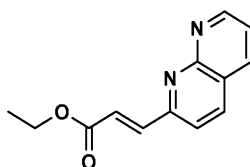
Intermediate **1W**: ^1H NMR (500 MHz, chloroform-*d*) δ 9.28 (s, 1H), 8.60 (d, J = 5.7 Hz, 1H), 8.01 (d, J = 8.5 Hz, 1H), 7.93 (d, J = 1.5 Hz, 1H), 7.86 (d, J = 16.1 Hz, 1H), 7.81 (dd, J = 8.6, 1.7 Hz, 1H), 7.69 (d, J = 5.6 Hz, 1H), 6.64 (d, J = 16.0 Hz, 1H), 4.33 (q, J = 7.1 Hz, 2H), 1.39 (t, J = 7.1 Hz, 3H). LCMS (ES): m/z 228.1 $[\text{M}+\text{H}]^+$.

**Intermediate 1X**

Intermediate **1X**: ^1H NMR (500 MHz, chloroform-*d*) δ 8.69 (s, 2H), 7.60 (d, J = 16.2 Hz, 1H), 6.47 (d, J = 16.2 Hz, 1H), 4.30 (q, J = 7.1 Hz, 2H), 4.08 (s, 3H), 1.37 (t, J = 7.1 Hz, 3H). LCMS (ES): m/z 209.2 $[\text{M}+\text{H}]^+$.

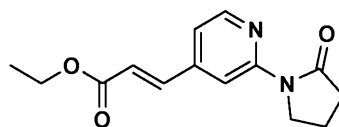
**Intermediate 1Y**

Intermediate **1Y**: ^1H NMR (500 MHz, chloroform-*d*) δ 7.86 (d, J = 16.2 Hz, 1H), 7.58 (d, J = 9.2 Hz, 1H), 7.02 (d, J = 9.1 Hz, 1H), 6.78 (d, J = 16.2 Hz, 1H), 4.32 (q, J = 7.1 Hz, 2H), 4.21 (s, 3H), 1.38 (t, J = 7.1 Hz, 3H). LCMS (ES): m/z 209.3 $[\text{M}+\text{H}]^+$.

**Intermediate 1Z**

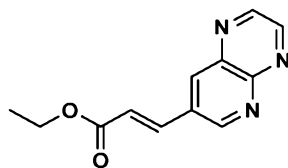
Intermediate **1Z**: ^1H NMR (500 MHz, chloroform-*d*) δ 9.19 (dd, J = 4.2, 2.1 Hz, 1H), 8.26 (d, J = 8.3 Hz, 1H), 8.22 (dd, J = 8.1, 2.0 Hz, 1H), 7.93 (d, J = 15.8 Hz, 1H),

7.71 (d, $J = 8.3$ Hz, 1H), 7.53 (dd, $J = 8.1, 4.2$ Hz, 1H), 7.22 (d, $J = 15.9$ Hz, 1H), 4.33 (q, $J = 7.1$ Hz, 2H), 1.31 (t, $J = 7.1$ Hz, 3H). LCMS (ES): m/z 229.2 $[M+H]^+$.



Intermediate 1AA

5 Intermediate **1AA**: ^1H NMR (500 MHz, chloroform- d) δ 8.60 – 8.56 (m, 1H), 8.39 (d, $J = 5.2$ Hz, 1H), 7.65 (d, $J = 16.1$ Hz, 1H), 7.14 (dd, $J = 5.3, 1.5$ Hz, 1H), 6.64 (d, $J = 16.1$ Hz, 1H), 4.30 (q, $J = 7.1$ Hz, 2H), 4.18 – 4.08 (m, 2H), 2.71 (t, $J = 8.1$ Hz, 2H), 2.23 – 2.10 (m, 2H), 1.37 (t, $J = 7.1$ Hz, 3H). LCMS (ES): m/z 261.2 $[M+H]^+$.

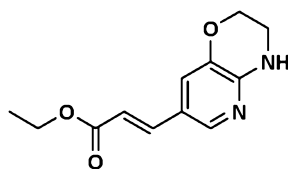


Intermediate 1AB

10

Intermediate **1AB**: ^1H NMR (500 MHz, chloroform- d) δ 9.38 (d, $J = 2.4$ Hz, 1H), 9.10 (d, $J = 1.7$ Hz, 1H), 9.01 (d, $J = 1.7$ Hz, 1H), 8.57 (d, $J = 2.5$ Hz, 1H), 7.92 (d, $J = 16.2$ Hz, 1H), 6.80 (d, $J = 16.1$ Hz, 1H), 4.36 (q, $J = 7.1$ Hz, 2H), 1.41 (t, $J = 7.1$ Hz, 3H). LCMS (ES): m/z 230.2 $[M+H]^+$.

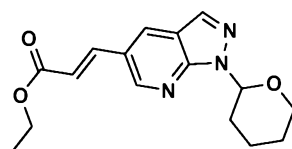
15



Intermediate 1AC

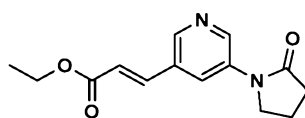
Intermediate **1AC**: ^1H NMR (500 MHz, chloroform- d) δ 7.82 (d, $J = 2.1$ Hz, 1H), 7.58 (d, $J = 16.0$ Hz, 1H), 7.19 (d, $J = 2.1$ Hz, 1H), 6.22 (d, $J = 15.9$ Hz, 1H), 5.34 (s, 1H), 4.31 – 4.20 (m, 4H), 3.63 (td, $J = 4.6, 2.2$ Hz, 2H), 1.35 (t, $J = 7.1$ Hz, 3H). LCMS (ES): m/z 235.2 $[M+H]^+$.

20



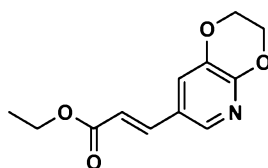
Intermediate 1AD

Intermediate **1AD**: ^1H NMR (500 MHz, chloroform-*d*) δ 8.75 (d, $J = 2.0$ Hz, 1H), 8.22 (d, $J = 2.2$ Hz, 1H), 8.13 (s, 1H), 7.82 (d, $J = 16.0$ Hz, 1H), 6.53 (d, $J = 16.0$ Hz, 1H), 6.14 (dd, $J = 10.5, 2.6$ Hz, 1H), 4.31 (q, $J = 7.1$ Hz, 2H), 4.18 – 4.10 (m, 1H), 3.85 (tt, $J = 11.5, 2.5$ Hz, 1H), 2.25 – 2.11 (m, 1H), 2.05 – 1.97 (m, 1H), 1.89 – 1.75 (m, 2H), 1.73 – 1.61 (m, 2H), 1.37 (t, $J = 7.1$ Hz, 3H). LCMS (ES): m/z 302.1 $[\text{M}+\text{H}]^+$.



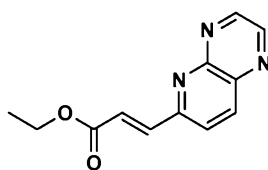
Intermediate 1AE

Intermediate **1AE**: ^1H NMR (500 MHz, chloroform-*d*) δ 8.72 (d, $J = 2.5$ Hz, 1H), 8.54 (d, $J = 1.9$ Hz, 1H), 8.48 (t, $J = 2.3$ Hz, 1H), 7.71 (d, $J = 16.2$ Hz, 1H), 6.57 (d, $J = 16.1$ Hz, 1H), 4.31 (q, $J = 7.1$ Hz, 2H), 3.95 (t, $J = 7.1$ Hz, 2H), 2.68 (t, $J = 8.1$ Hz, 2H), 2.32 – 2.21 (m, 2H), 1.37 (t, $J = 7.1$ Hz, 3H). LCMS (ES): m/z 261.2 $[\text{M}+\text{H}]^+$.



Intermediate 1AF

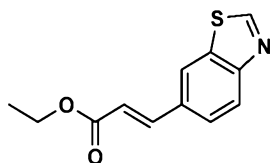
Intermediate **1AF**: ^1H NMR (500 MHz, chloroform-*d*) δ 7.98 (d, $J = 2.2$ Hz, 1H), 7.61 (d, $J = 16.0$ Hz, 1H), 7.38 (d, $J = 2.1$ Hz, 1H), 6.35 (d, $J = 16.0$ Hz, 1H), 4.53 – 4.47 (m, 2H), 4.34 – 4.23 (m, 4H), 1.36 (t, $J = 7.1$ Hz, 3H). LCMS (ES): m/z 236.0 $[\text{M}+\text{H}]^+$.



Intermediate 1AG

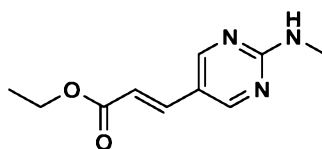
Intermediate **1AG**: ^1H NMR (500 MHz, chloroform-*d*) δ 9.13 (d, $J = 1.7$ Hz, 1H), 8.97 (d, $J = 1.8$ Hz, 1H), 8.53 (d, $J = 8.6$ Hz, 1H), 7.99 – 7.90 (m, 2H), 7.23 (d, $J = 15.9$ Hz, 1H), 4.35 (q, $J = 7.1$ Hz, 2H), 1.40 (t, $J = 7.1$ Hz, 3H). LCMS (ES): m/z 230.2 $[\text{M}+\text{H}]^+$.

5

Intermediate **1AH**

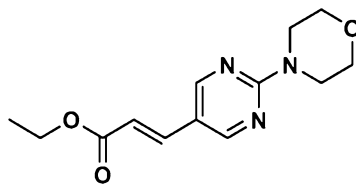
Intermediate **1AH**: ^1H NMR (500 MHz, chloroform-*d*) δ 9.07 (s, 1H), 8.16 (d, $J = 8.5$ Hz, 1H), 8.13 (d, $J = 1.7$ Hz, 1H), 7.83 (d, $J = 16.0$ Hz, 1H), 7.74 (dd, $J = 8.5, 1.8$ Hz, 1H), 6.55 (d, $J = 16.1$ Hz, 1H), 4.32 (q, $J = 7.1$ Hz, 2H), 1.38 (td, $J = 7.2, 1.4$ Hz, 3H).

10 LCMS (ES): m/z 234.1 $[\text{M}+\text{H}]^+$.

Intermediate **1AI**

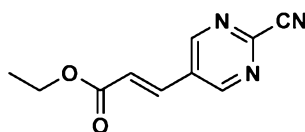
Intermediate **1AI**: ^1H NMR (500 MHz, chloroform-*d*) δ 8.49 (s, 2H), 7.52 (d, $J = 16.2$ Hz, 1H), 6.33 (d, $J = 16.1$ Hz, 1H), 5.40 (s, 1H), 4.28 (q, $J = 7.1$ Hz, 2H), 3.08 (d, $J = 5.1$ Hz, 3H), 1.36 (t, $J = 7.1$ Hz, 3H). LCMS (ES): m/z 208.3 $[\text{M}+\text{H}]^+$.

15

Intermediate **1AJ**

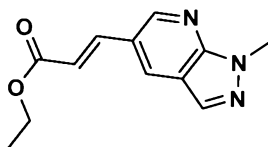
Intermediate **1AJ**: ^1H NMR (500 MHz, chloroform-*d*) δ 8.50 (s, 2H), 7.52 (d, $J = 16.0$ Hz, 1H), 6.33 (d, $J = 16.0$ Hz, 1H), 4.28 (q, $J = 7.1$ Hz, 2H), 3.90 (dd, $J = 5.7, 4.0$

Hz, 4H), 3.79 (dd, $J = 5.7, 4.1$ Hz, 4H), 1.36 (t, $J = 7.1$ Hz, 3H). LCMS (ES): m/z 264.2 $[M+H]^+$.



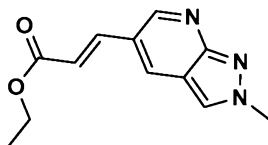
Intermediate 1AK

- 5 Intermediate **1AK**: ^1H NMR (500 MHz, chloroform- d) δ 8.98 (s, 2H), 7.64 (d, $J = 16.2$ Hz, 1H), 6.70 (d, $J = 16.2$ Hz, 1H), 4.34 (q, $J = 7.1$ Hz, 2H), 1.39 (t, $J = 7.1$ Hz, 3H). LCMS (ES): m/z 204.4 $[M+H]^+$.



Intermediate 1AL

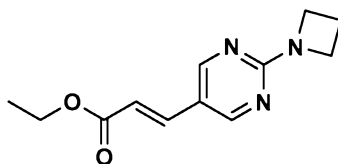
- 10 Intermediate **1AL**: ^1H NMR (500 MHz, chloroform- d) δ 8.75 (d, $J = 2.0$ Hz, 1H), 8.22 (d, $J = 2.1$ Hz, 1H), 8.07 (s, 1H), 7.84 (d, $J = 16.1$ Hz, 1H), 6.54 (d, $J = 16.1$ Hz, 1H), 4.32 (q, $J = 7.2$ Hz, 2H), 4.20 (s, 3H), 1.39 (t, $J = 7.1$ Hz, 3H). LCMS (ES): m/z 232.4 $[M+H]^+$.



Intermediate 1AM

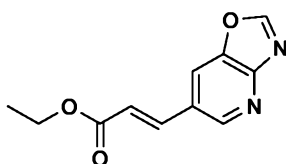
- 15 Intermediate **1AM**: ^1H NMR (500 MHz, chloroform- d) δ 8.93 (d, $J = 2.2$ Hz, 1H), 8.15 (d, $J = 2.2$ Hz, 1H), 7.99 (s, 1H), 7.79 (d, $J = 16.0$ Hz, 1H), 6.55 (d, $J = 16.0$ Hz, 1H), 4.35 – 4.30 (m, 2H), 4.30 (s, 3H), 1.38 (t, $J = 7.1$ Hz, 3H). LCMS (ES): m/z 232.4 $[M+H]^+$.

20

**Intermediate 1AN**

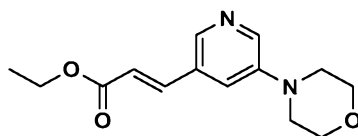
Intermediate **1AN**: ^1H NMR (500 MHz, chloroform-*d*) δ 8.48 (s, 2H), 7.51 (d, J = 16.0 Hz, 1H), 6.31 (d, J = 16.0 Hz, 1H), 4.27 (dt, J = 15.1, 7.3 Hz, 6H), 2.45 (p, J = 7.6 Hz, 2H), 1.35 (t, J = 7.1 Hz, 3H). LCMS (ES): m/z 234.4 $[\text{M}+\text{H}]^+$.

5

**Intermediate 1AO**

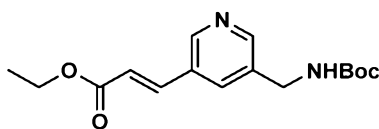
Intermediate **1AO**: ^1H NMR (500 MHz, chloroform-*d*) δ 8.81 (d, J = 1.9 Hz, 1H), 8.41 (s, 1H), 8.08 (d, J = 1.9 Hz, 1H), 7.84 (d, J = 16.2 Hz, 1H), 6.59 (d, J = 16.1 Hz, 1H), 4.33 (q, J = 7.1 Hz, 2H), 1.39 (t, J = 7.1 Hz, 3H). LCMS (ES): m/z 219.2 $[\text{M}+\text{H}]^+$.

10

**Intermediate 1AP**

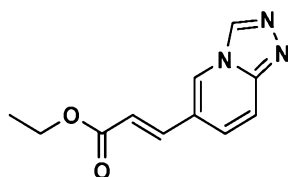
Intermediate **1AP**: ^1H NMR (500 MHz, chloroform-*d*) δ 8.32 (d, J = 2.8 Hz, 1H), 8.28 (d, J = 1.8 Hz, 1H), 7.66 (d, J = 16.0 Hz, 1H), 7.27 (t, J = 2.3 Hz, 1H), 6.50 (d, J = 16.0 Hz, 1H), 4.30 (q, J = 7.1 Hz, 2H), 3.98 – 3.84 (m, 4H), 3.28 – 3.20 (m, 4H), 1.37 (t, J = 7.1 Hz, 3H). LCMS (ES): m/z 263.2 $[\text{M}+\text{H}]^+$.

15

**Intermediate 1AR**

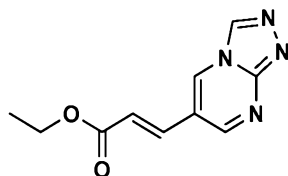
Intermediate **1AR**: ^1H NMR (500 MHz, chloroform-*d*) δ 8.67 (d, J = 2.2 Hz, 1H), 8.55 (d, J = 2.0 Hz, 1H), 7.78 (t, J = 2.2 Hz, 1H), 7.69 (d, J = 16.1 Hz, 1H), 6.53 (d, J =

16.0 Hz, 1H), 4.96 (s, 1H), 4.39 (d, $J = 6.2$ Hz, 2H), 4.31 (q, $J = 7.1$ Hz, 2H), 1.49 (s, 9H), 1.37 (t, $J = 7.1$ Hz, 3H). LCMS (ES): m/z 307.1 $[M+H]^+$.



Intermediate 1AS

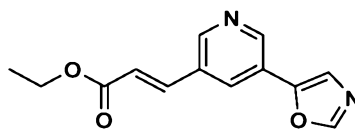
5 Intermediate **1AS**: ^1H NMR (500 MHz, chloroform- d) δ 8.87 (s, 1H), 8.26 (s, 1H), 7.84 (d, $J = 9.6$ Hz, 1H), 7.63 (d, $J = 15.9$ Hz, 1H), 7.52 (dd, $J = 9.6, 1.6$ Hz, 1H), 6.51 (d, $J = 15.9$ Hz, 1H), 4.32 (q, $J = 7.1$ Hz, 2H), 1.38 (t, $J = 7.1$ Hz, 3H). LCMS (ES): m/z 218.4 $[M+H]^+$.



Intermediate 1AT

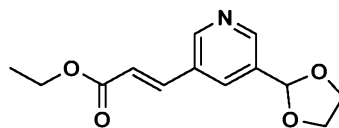
10

Intermediate **1AT**: LCMS (ES): m/z 219.2 $[M+H]^+$.



Intermediate 1AU

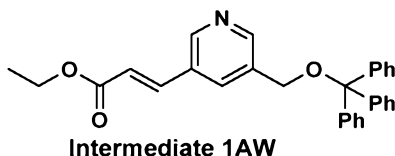
15 Intermediate **1AU**: ^1H NMR (500 MHz, chloroform- d) δ 8.94 (d, $J = 2.1$ Hz, 1H), 8.73 (d, $J = 2.0$ Hz, 1H), 8.09 (t, $J = 2.1$ Hz, 1H), 8.03 (s, 1H), 7.73 (d, $J = 16.2$ Hz, 1H), 7.54 (s, 1H), 6.62 (d, $J = 16.0$ Hz, 1H), 4.33 (q, $J = 7.1$ Hz, 2H), 1.39 (t, $J = 7.1$ Hz, 3H). LCMS (ES): m/z 245.4 $[M+H]^+$.



Intermediate 1AV

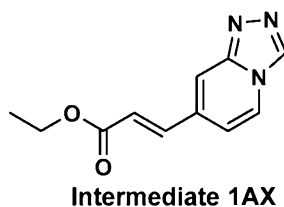
Intermediate **1AV**: ^1H NMR (500 MHz, $\text{MeOH-}d_4$) δ 8.74 (d, $J = 2.1$ Hz, 1H), 8.62 (d, $J = 2.0$ Hz, 1H), 8.13 (d, $J = 2.2$ Hz, 1H), 7.71 (d, $J = 16.1$ Hz, 1H), 6.68 (d, $J = 16.4$ Hz, 1H), 5.87 (s, 1H), 4.31 – 4.22 (m, 2H), 4.19 – 4.00 (m, 4H), 1.34 (t, $J = 7.2$ Hz, 3H). LCMS (ES): m/z 250.2 $[\text{M}+\text{H}]^+$.

5



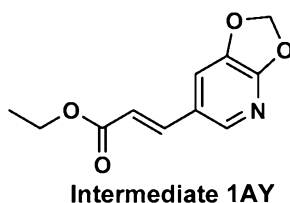
Intermediate **1AW**: ^1H NMR (400 MHz, $\text{chloroform-}d$) δ 8.64 (d, $J = 2.1$ Hz, 1H), 8.56 (d, $J = 2.0$ Hz, 1H), 7.78 – 7.74 (m, 1H), 7.67 (d, $J = 16.1$ Hz, 1H), 7.54 – 7.45 (m, 6H), 7.36 – 7.30 (m, 6H), 7.28 (t, $J = 1.4$ Hz, 1H), 7.26 (m, 2H), 6.49 (d, $J = 16.2$ Hz, 1H), 4.29 (m, 4H), 1.35 (t, $J = 7.1$ Hz, 3H). LCMS (ES): m/z 450.4 $[\text{M}+\text{H}]^+$.

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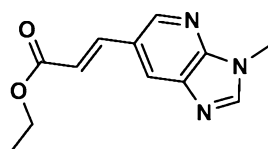
Intermediate **1AX**: ^1H NMR (500 MHz, $\text{chloroform-}d$) δ 8.86 (s, 1H), 8.15 (d, $J = 7.2$ Hz, 1H), 7.89 (s, 1H), 7.69 (d, $J = 16.0$ Hz, 1H), 7.07 (dd, $J = 7.2, 1.4$ Hz, 1H), 6.54 (d, $J = 16.0$ Hz, 1H), 4.33 (q, $J = 7.1$ Hz, 2H), 1.39 (t, $J = 7.1$ Hz, 3H). LCMS (ES): m/z 218.4 $[\text{M}+\text{H}]^+$.

15

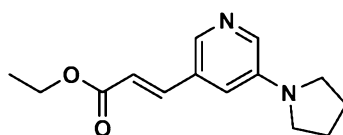


Intermediate **1AY**: ^1H NMR (400 MHz, $\text{MeOH-}d_4$) δ 7.76 (d, $J = 1.9$ Hz, 1H), 7.61 (d, $J = 16.0$ Hz, 1H), 7.48 (d, $J = 1.8$ Hz, 1H), 6.45 (d, $J = 16.0$ Hz, 1H), 6.16 (s, 2H), 4.23 (q, $J = 7.1$ Hz, 2H), 1.31 (t, $J = 7.1$ Hz, 3H). LCMS (ES): m/z 222.2 $[\text{M}+\text{H}]^+$.

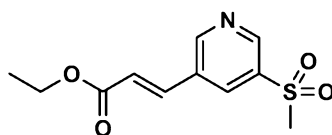
20

**Intermediate 1AZ**

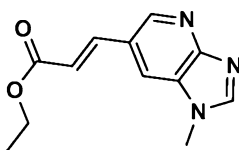
Intermediate **1AZ**: ^1H NMR (500 MHz, chloroform-*d*) δ 8.57 (d, $J = 2.0$ Hz, 1H), 8.24 (d, $J = 1.9$ Hz, 1H), 8.07 (s, 1H), 7.84 (d, $J = 16.0$ Hz, 1H), 6.53 (d, $J = 16.1$ Hz, 1H), 4.29 (q, $J = 7.1$ Hz, 2H), 3.93 (s, 3H), 1.36 (t, $J = 7.2$ Hz, 3H). LCMS (ES): m/z 232.2 $[\text{M}+\text{H}]^+$.

**Intermediate 1BA**

Intermediate **1BA**: ^1H NMR (500 MHz, chloroform-*d*) δ 8.06 (d, $J = 1.7$ Hz, 1H), 7.98 (d, $J = 2.7$ Hz, 1H), 7.63 (d, $J = 16.0$ Hz, 1H), 6.88 (t, $J = 2.3$ Hz, 1H), 6.47 (d, $J = 16.0$ Hz, 1H), 4.27 (q, $J = 7.1$ Hz, 2H), 3.37 – 3.26 (m, 4H), 2.09 – 2.00 (m, 4H), 1.34 (t, $J = 7.1$ Hz, 3H). LCMS (ES): m/z 247.2 $[\text{M}+\text{H}]^+$.

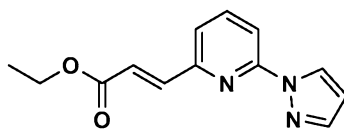
**Intermediate 1BB**

Intermediate **1BB**: ^1H NMR (500 MHz, chloroform-*d*) δ 9.12 (d, $J = 2.1$ Hz, 1H), 8.98 (d, $J = 2.1$ Hz, 1H), 8.34 (t, $J = 2.2$ Hz, 1H), 7.71 (d, $J = 16.1$ Hz, 1H), 6.63 (d, $J = 16.2$ Hz, 1H), 4.30 (q, $J = 7.1$ Hz, 2H), 3.14 (s, 3H), 1.36 (t, $J = 7.1$ Hz, 3H). LCMS (ES): m/z 256.1 $[\text{M}+\text{H}]^+$.

**Intermediate 1BC**

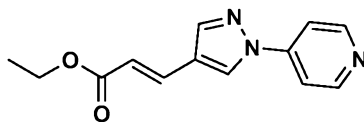
Intermediate **1BC**: ^1H NMR (500 MHz, chloroform-*d*) δ 8.75 (d, $J = 2.2$ Hz, 1H), 8.13 (s, 1H), 7.87 (d, $J = 2.0$ Hz, 1H), 7.84 (d, $J = 16.0$ Hz, 1H), 6.55 (d, $J = 16.0$ Hz, 1H), 4.29 (q, $J = 7.1$ Hz, 2H), 3.91 (s, 3H), 1.36 (t, $J = 7.1$ Hz, 3H). LCMS (ES): m/z 232.2 $[\text{M}+\text{H}]^+$.

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Intermediate **1BD**

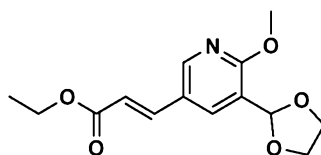
Intermediate **1BD**: ^1H NMR (400 MHz, chloroform-*d*) δ 8.65 (dd, $J = 2.6, 0.7$ Hz, 1H), 7.99 (dd, $J = 8.3, 0.9$ Hz, 1H), 7.83 (dd, $J = 8.3, 7.5$ Hz, 1H), 7.75 (dd, $J = 1.7, 0.8$ Hz, 1H), 7.66 (d, $J = 15.5$ Hz, 1H), 7.28 (dd, $J = 7.6, 0.9$ Hz, 1H), 7.01 (d, $J = 15.6$ Hz, 1H), 6.48 (dd, $J = 2.6, 1.7$ Hz, 1H), 4.30 (q, $J = 7.1$ Hz, 2H), 1.37 (t, $J = 7.1$ Hz, 3H). LCMS (ES): m/z 244.1 $[\text{M}+\text{H}]^+$.

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Intermediate **1BE**

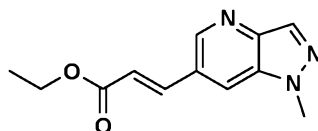
Intermediate **1BE**: ^1H NMR (400 MHz, chloroform-*d*) δ 8.74 – 8.67 (m, 2H), 8.17 (s, 1H), 7.97 (s, 1H), 7.65 – 7.62 (m, 2H), 7.59 (d, $J = 16.0$ Hz, 1H), 6.32 (d, $J = 16.0$ Hz, 1H), 4.27 (q, $J = 7.1$ Hz, 2H), 1.34 (t, $J = 7.1$ Hz, 3H). LCMS (ES): m/z 244.1 $[\text{M}+\text{H}]^+$.

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Intermediate **1BF**

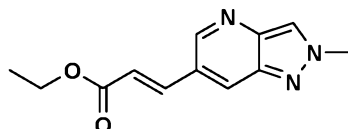
Intermediate **1BF**: LCMS (ES): m/z 232.1 $[\text{M}+\text{H}]^+$.

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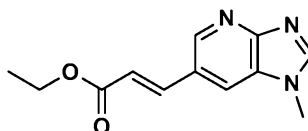
**Intermediate 1BG**

Intermediate **1BG**: ^1H NMR (500 MHz, chloroform-*d*) δ 8.79 (d, $J = 1.9$ Hz, 1H), 8.25 (d, $J = 1.0$ Hz, 1H), 7.91 – 7.81 (m, 2H), 6.66 (d, $J = 16.2$ Hz, 1H), 4.33 (q, $J = 7.1$ Hz, 2H), 4.15 (s, 3H), 1.39 (t, $J = 7.1$ Hz, 3H). LCMS (ES): m/z 232.1 $[\text{M}+\text{H}]^+$.

5

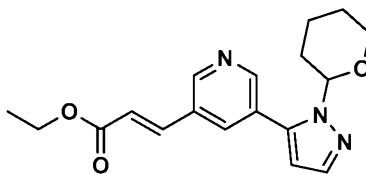
**Intermediate 1BH**

Intermediate **1BH**: ^1H NMR (500 MHz, chloroform-*d*) δ 8.79 (d, $J = 1.9$ Hz, 1H), 8.21 (s, 1H), 8.14 (d, $J = 1.8$ Hz, 1H), 7.83 (d, $J = 16.0$ Hz, 1H), 6.62 (d, $J = 16.1$ Hz, 1H), 4.33 (q, $J = 7.1$ Hz, 2H), 4.31 (s, 3H), 1.39 (t, $J = 7.1$ Hz, 3H). LCMS (ES): m/z

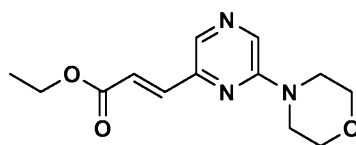
10 232.1 $[\text{M}+\text{H}]^+$.**Intermediate 1BI**

Intermediate **1BI**: ^1H NMR (500 MHz, chloroform-*d*) δ 8.80 - 8.74 (m, 1H), 8.19 - 8.14 (m, 1H), 7.93 - 7.82 (m, 2H), 6.61 - 6.52 (m, 1H), 4.32 (q, $J = 7.2$ Hz, 2H), 3.93 (s, 3H), 1.38 (t, $J = 7.0$ Hz, 3H). LCMS (ES): m/z 232.2 $[\text{M}+\text{H}]^+$.

15

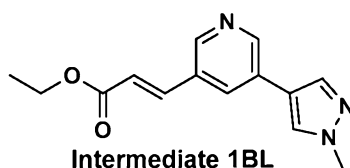
**Intermediate 1BJ**

Intermediate **1BJ**: LCMS (ES): m/z 328.1 $[\text{M}+\text{H}]^+$.

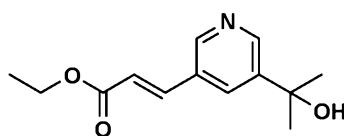
**Intermediate 1BK**

Intermediate **1BK**: ^1H NMR (500 MHz, chloroform-*d*) δ 8.16 - 8.11 (m, 1H), 8.00 - 7.92 (m, 1H), 7.60 - 7.50 (m, 1H), 7.00 - 6.87 (m, 1H), 4.28 (q, $J=7.3$ Hz, 2H), 3.89 - 3.80 (m, 4H), 3.70 - 3.59 (m, 4H), 1.35 (t, $J=7.2$ Hz, 3H). LCMS (ES): m/z 264.1

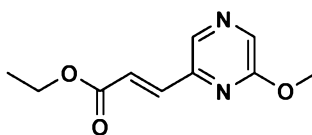
5 [M+H] $^+$.

**Intermediate 1BL**

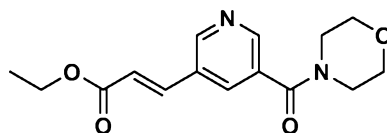
Intermediate **1BL**: ^1H NMR (400 MHz, chloroform-*d*) δ 8.72 (d, $J = 2.1$ Hz, 1H), 8.58 (d, $J = 2.0$ Hz, 1H), 7.86 (t, $J = 2.2$ Hz, 1H), 7.81 (d, $J = 0.9$ Hz, 1H), 7.69 (m, 1H), 7.69 (d, $J = 16.2$ Hz, 1H), 6.55 (d, $J = 16.1$ Hz, 1H), 4.29 (q, $J = 7.1$ Hz, 2H), 3.98 (s, 3H), 1.36 (t, $J = 7.1$ Hz, 3H). LCMS (ES): m/z 258.1 [M+H] $^+$.

**Intermediate 1BM**

Intermediate **1BM**: ^1H NMR (400 MHz, chloroform-*d*) δ 8.73 - 8.66 (m, 1H), 8.62 - 8.55 (m, 1H), 8.05 - 7.98 (m, 1H), 7.67 (d, $J=16.1$ Hz, 1H), 6.60 - 6.45 (m, 1H), 4.33 - 4.20 (m, 2H), 1.63 (s, 6H), 1.34 (t, $J=7.2$ Hz, 3H). LCMS (ES): m/z 236.1 [M+H] $^+$.

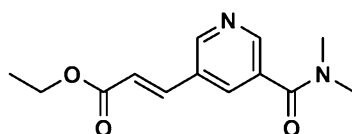
**Intermediate 1BN**

Intermediate **1BN**: ^1H NMR (400 MHz, chloroform-*d*) δ 8.25 - 8.17 (m, 2H), 7.67 - 7.60 (m, 1H), 7.04 (d, $J=15.4$ Hz, 1H), 4.37 - 4.27 (m, 2H), 4.07 - 3.98 (m, 3H), 1.38 (t, $J=7.2$ Hz, 3H). LCMS (ES): m/z 209.0 $[\text{M}+\text{H}]^+$.



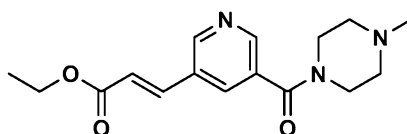
Intermediate **1BO**

Intermediate **1BO**: ^1H NMR (500 MHz, chloroform-*d*) δ 8.86 - 8.74 (m, 1H), 8.70 - 8.58 (m, 1H), 7.96 - 7.84 (m, 1H), 7.75 - 7.62 (m, 1H), 6.62 - 6.49 (m, 1H), 4.36 - 4.24 (m, 2H), 3.97 - 3.40 (m, 8H), 1.42 - 1.31 (m, 3H). LCMS (ES): m/z 291.1 $[\text{M}+\text{H}]^+$.



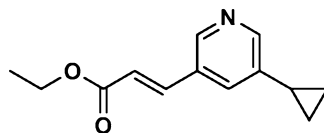
Intermediate **1BP**

Intermediate **1BP**: ^1H NMR (500 MHz, chloroform-*d*) δ 8.79 (d, $J=1.9$ Hz, 1H), 8.67 (d, $J=1.9$ Hz, 1H), 7.93 (t, $J=1.9$ Hz, 1H), 7.69 (d, $J=16.2$ Hz, 1H), 6.56 (d, $J=16.2$ Hz, 1H), 4.35 - 4.25 (m, 2H), 3.25 - 3.10 (m, 3H), 3.10 - 3.00 (m, 3H), 2.99 - 2.87 (m, 2H), 1.37 (t, $J=7.2$ Hz, 3H). LCMS (ES): m/z 249.1 $[\text{M}+\text{H}]^+$.

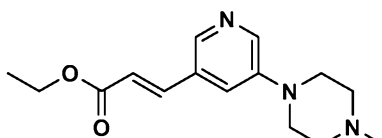


Intermediate **1BQ**

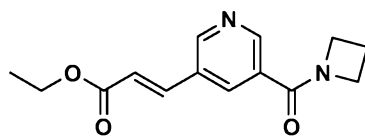
Intermediate **1BQ**: ^1H NMR (500 MHz, chloroform-*d*) δ 8.84 - 8.76 (m, 1H), 8.68 - 8.62 (m, 1H), 7.91 (t, $J=2.1$ Hz, 1H), 7.75 - 7.65 (m, 1H), 6.61 - 6.51 (m, 1H), 4.35 - 4.27 (m, 2H), 3.95 - 3.78 (m, 2H), 3.55 - 3.38 (m, 2H), 2.61 - 2.38 (m, 4H), 2.36 (s, 3H), 1.37 (t, $J=7.2$ Hz, 3H). LCMS (ES): m/z 304.1 $[\text{M}+\text{H}]^+$.

**Intermediate 1BR**

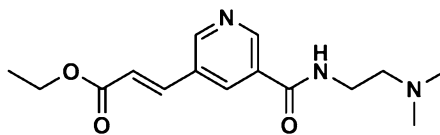
Intermediate **1BR**: ^1H NMR (500 MHz, chloroform-*d*) δ 8.59 - 8.47 (m, 1H), 8.47 - 8.37 (m, 1H), 7.70 - 7.61 (m, 1H), 7.49 - 7.42 (m, 1H), 6.57 - 6.43 (m, 1H), 4.30 (q, $J=7.2$ Hz, 2H), 2.01 - 1.89 (m, 1H), 1.42 - 1.32 (m, 3H), 1.13 - 1.07 (m, 2H), 0.81 - 0.75 (m, 2H). LCMS (ES): m/z 218.2 $[\text{M}+\text{H}]^+$.

**Intermediate 1BS**

Intermediate **1BS**: ^1H NMR (500 MHz, chloroform-*d*) δ 8.39 - 8.30 (m, 1H), 8.30 - 8.21 (m, 1H), 7.72 - 7.61 (m, 1H), 7.35 - 7.24 (m, 1H), 6.49 (d, $J=16.0$ Hz, 1H), 4.30 (q, $J=7.2$ Hz, 2H), 3.35 - 3.23 (m, 4H), 2.70 - 2.55 (m, 4H), 2.39 (s, 3H), 1.36 (t, $J=7.2$ Hz, 3H). LCMS (ES): m/z 276.1 $[\text{M}+\text{H}]^+$.

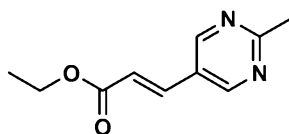
**Intermediate 1BT**

Intermediate **1BT**: ^1H NMR (500 MHz, chloroform-*d*) δ 8.88 - 8.74 (m, 2H), 8.23 - 8.10 (m, 1H), 7.77 - 7.61 (m, 1H), 6.73 - 6.49 (m, 1H), 4.50 - 4.21 (m, 6H), 2.41 (quin, $J=7.8$ Hz, 2H), 1.35 (t, $J=7.2$ Hz, 3H). LCMS (ES): m/z 261.1 $[\text{M}+\text{H}]^+$.

**Intermediate 1BU**

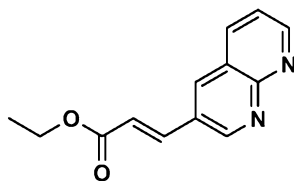
Intermediate **1BU**: ^1H NMR (500 MHz, chloroform-*d*) δ 9.04 - 8.95 (m, 1H), 8.90 - 8.82 (m, 1H), 8.36 - 8.25 (m, 1H), 7.78 - 7.67 (m, 1H), 7.09 - 6.97 (m, 1H), 6.69 -

6.54 (m, 1H), 4.32 (q, $J=7.2$ Hz, 2H), 3.57 (q, $J=5.4$ Hz, 2H), 2.58 (t, $J=5.9$ Hz, 2H), 2.37 - 2.29 (m, 6H), 1.38 (t, $J=7.0$ Hz, 3H). LCMS (ES): m/z 292.1 $[M+H]^+$.



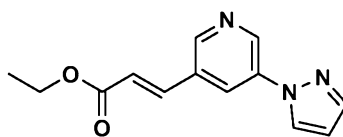
Intermediate 1BV

5 Intermediate **1BV**: ^1H NMR (400 MHz, DMSO- d_6) δ 9.04 (s, 2H), 7.63 (d, $J = 16.1$ Hz, 1H), 6.87 (d, $J = 16.3$ Hz, 1H), 4.21 (q, $J = 7.2$ Hz, 2H), 2.64 (s, 3H), 1.26 (t, $J = 7.2$ Hz, 3H). LCMS (ES): m/z 193.1 $[M+H]^+$.



Intermediate 1BW

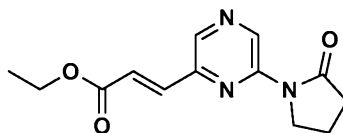
10 Intermediate **1BW**: ^1H NMR (500 MHz, chloroform- d) δ 9.40 - 9.31 (m, 1H), 9.24 - 9.15 (m, 1H), 8.32 - 8.28 (m, 1H), 8.28 - 8.23 (m, 1H), 7.88 (d, $J=16.2$ Hz, 1H), 7.57 (dd, $J=8.1, 4.3$ Hz, 1H), 6.73 (d, $J=16.0$ Hz, 1H), 4.34 (q, $J=7.2$ Hz, 2H), 1.40 (t, $J=7.2$ Hz, 3H). LCMS (ES): m/z 229.2 $[M+H]^+$.



Intermediate 1BX

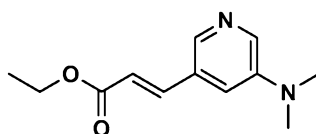
15

Intermediate **1BX**: LCMS (ES): m/z 244.0 $[M+H]^+$.

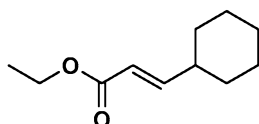


Intermediate 1BY

Intermediate **1BY**: LCMS (ES): m/z 262.1 $[M+H]^+$.

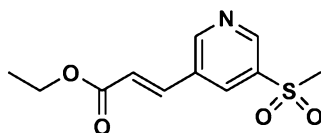
**Intermediate 1BZ**

Intermediate **1BZ**: LCMS (ES): m/z 249.1 $[M+H]^+$.

**Intermediate 1CA**

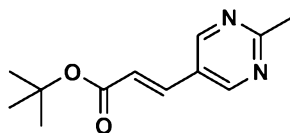
5

Intermediate **1CA**: ^1H NMR (500 MHz, chloroform- d) δ 6.96 - 6.72 (m, 1H), 5.84 - 5.71 (m, 1H), 4.19 - 3.99 (m, 2H), 2.24 - 2.05 (m, 1H), 1.82 - 1.54 (m, 5H), 1.35 - 1.00 (m, 8H).

**Intermediate 1CB**

10

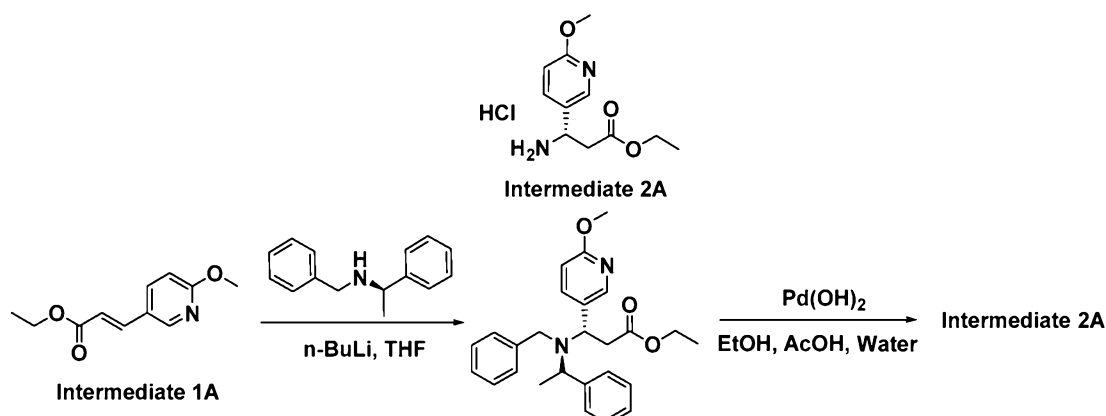
Intermediate **1CB**: ^1H NMR (500 MHz, chloroform- d) δ 9.18 - 9.12 (m, 1H), 9.04 - 8.97 (m, 1H), 8.41 - 8.33 (m, 1H), 7.79 - 7.69 (m, 1H), 6.71 - 6.61 (m, 1H), 4.33 (q, $J=7.2$ Hz, 2H), 3.17 (s, 3H), 1.38 (t, $J=7.2$ Hz, 3H). LCMS (ES): m/z 256.1 $[M+H]^+$.

**Intermediate 1CC**

15

Intermediate **1CC**: ^1H NMR (500 MHz, chloroform- d) δ 8.77 (s, 2H), 7.55 - 7.44 (m, 1H), 6.57 - 6.39 (m, 1H), 2.85 - 2.72 (m, 3H), 1.61 - 1.47 (m, 9H).

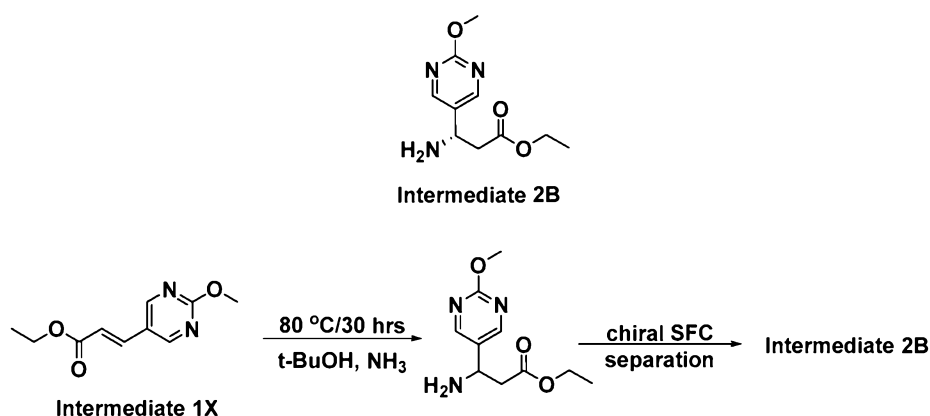
20 **Intermediate 2A. Ethyl (S)-3-amino-3-(3-fluoro-4-methoxyphenyl)propanoate**



Intermediate **2A** was prepared according to the procedure described in:

Hutchinson, J. H. *et. al.*, *J. Med. Chem.* **2003**, 46, 4790. ¹H NMR (500MHz, chloroform-*d*) δ 8.16 (d, *J* = 2.2 Hz, 1H), 7.67 (dd, *J* = 8.5, 2.5 Hz, 1H), 6.76 (d, *J* = 8.5 Hz, 1H), 4.47 (dd, *J* = 8.8, 5.0 Hz, 1H), 4.00 - 3.92 (m, 3H), 2.92 - 2.64 (m, 2H). LCMS (ES): *m/z* 225.0 [M+H]⁺. ¹H NMR (500MHz, CDCl₃) δ 8.28 (d, *J* = 2.5 Hz, 1H), 7.77 (dd, *J* = 8.8, 2.5 Hz, 1H), 7.64 (d, *J* = 16.2 Hz, 1H), 6.77 (d, *J* = 8.8 Hz, 1H), 6.34 (d, *J* = 16.0 Hz, 1H), 4.27 (q, *J* = 7.2 Hz, 2H), 3.98 (s, 3H), 1.34 (t, *J* = 7.2 Hz, 3H).

Intermediate 2B. Ethyl (S)-3-amino-3-(2-methoxypyrimidin-5-yl)propanoate



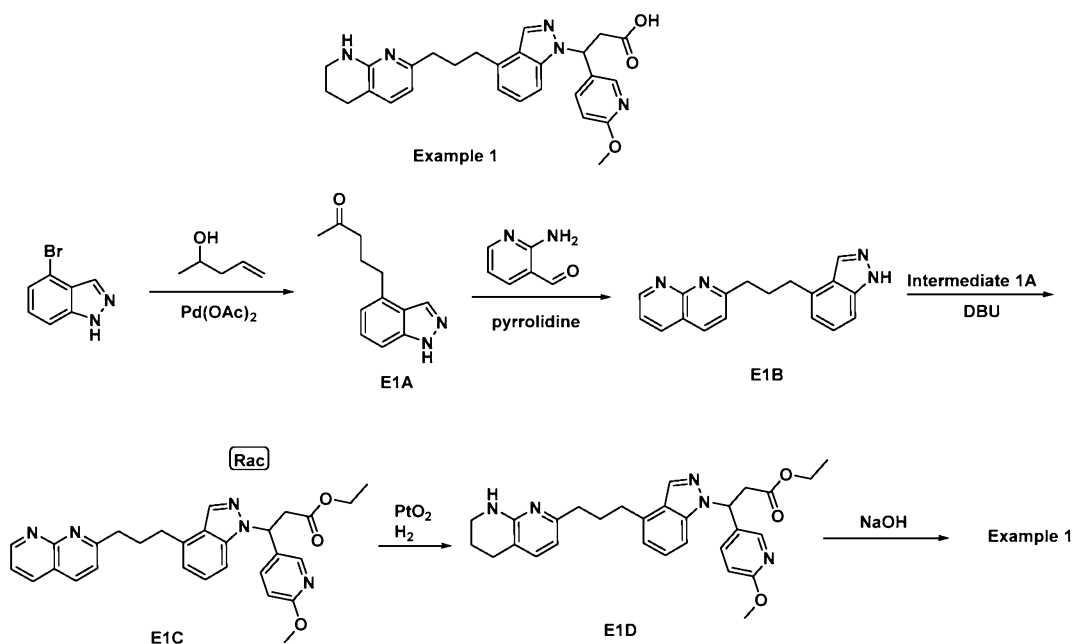
NH₃ gas was bubbled into a cooled *t*-BuOH (300 mL) for 1 hr. It was then added together with (*E*)-ethyl 3-(2-methoxypyrimidin-5-yl)acrylate (20 g, 96 mmol) into an 1 L autoclave. The mixture was heated at 80 °C for 30 hrs. The mixture was concentrated under reduced pressure. The residue was purified via flash column chromatography (5% methanol in chloroform) to afford racemic ethyl-3-amino-3-(2-methoxypyrimidin-5-

yl)propanoate. It was further purified in chiral SFC (Chiralpak IA, 0.4% DEA in EtOH) to afford Intermediate **2B** (2.3 g, 9.80 mmol, 10.2% yield). LCMS (ES): m/z 226.8 $[M+H]^+$.

- 5 Other β -aminoacids were prepared analogously using the procedure described for Intermediates **2A** and **2B** above.

Example 1

3-(6-Methoxypyridin-3-yl)-3-(4-(3-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)propyl)-1H-indazol-1-yl)propanoic acid



- 15 Intermediate **E1A**: To a solution of 4-bromoindazole (0.35 g, 1.776 mmol) in DMF (5.59 mL) was added pent-4-en-2-ol (0.28 mL, 2.66 mmol), $\text{Pd}(\text{OAc})_2$ (0.199 g, 0.888 mmol), LiCl (75 mg, 1.776 mmol), tetra-*n*-butylammonium chloride (0.99 g, 3.55 mmol), and LiOAc (0.294 g, 4.46 mmol). The mixture was heated at 100 °C for 72 hrs. The mixture was diluted with H_2O (10 mL) and extracted with EtOAc (3 x 5 mL). The combined organic layers were washed with H_2O (3 mL) and brine (5 mL). It was concentrated and purified *via* flash chromatography (SiO_2) to give Intermediate **E1A** (82 mg, 23%). ^1H NMR (500 MHz, chloroform-*d*) δ 8.17 (s, 1H), 7.40 - 7.25 (m, 2H), 6.94

(d, $J = 6.9$ Hz, 1H), 2.96 (t, $J = 7.6$ Hz, 2H), 2.48 (t, $J = 7.3$ Hz, 2H), 2.12 (s, 3H), 2.11 - 2.01 (m, 2H).

Intermediate **E1B**: To a solution of Intermediate **E1A** (82 mg, 0.406 mmol) in CH_2Cl_2 (203 μL) and MeOH (610 μL) was added pyrrolidine (8.4 μL , 0.102 mmol),
5 followed by 2-aminonicotinaldehyde (49.6 mg, 0.406 mmol). The mixture was stirred at room temperature overnight. The reaction was evaporated under the reduced pressure. The residue was purified via flash chromatography (SiO_2) to give Intermediate **E1B** (93 mg, 80%). ^1H NMR (500 MHz, chloroform- d) δ 9.09 (dd, $J = 4.3, 2.1$ Hz, 1H), 8.15 (dd, $J = 8.1, 2.0$ Hz, 1H), 8.12 - 8.06 (m, 2H), 7.44 (dd, $J = 8.1, 4.2$ Hz, 1H), 7.36 (dd, $J =$
10 8.3, 6.6 Hz, 2H), 7.30 - 7.23 (m, 1H), 6.98 (d, $J = 6.9$ Hz, 1H), 3.18 - 3.11 (m, 2H), 3.08 (t, $J = 7.6$ Hz, 2H), 2.48 - 2.33 (m, 2H).

Intermediate **E1C**: A mixture of **E1B** (31 mg, 0.108 mmol), Intermediate **1A** (44.6 mg, 0.215 mmol), and DBU (16.20 μL , 0.108 mmol) in acetonitrile (717 μL) was heated at 100 $^\circ\text{C}$ overnight. The solvent was removed under reduced pressure. The residue was
15 purified via preparative HPLC (Phenomenex Luna Axia 5 μ C18 30 x 100 mm; 10 min gradient from 30% A: 70% B to 0% A:100% B (A = 90% H_2O /10% MeOH + 0.1% TFA); (B = 90% MeOH/10% H_2O + 0.1% TFA); detection at 220 nm) to yield Intermediate **E1C** (17 mg, 32%). LCMS (ES): m/z 492.2 $[\text{M}+\text{H}]^+$.

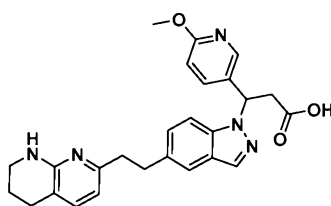
Intermediate **E1D**: To a solution of **E1C** (19 mg, 0.038 mmol) in ethanol (1.0 mL)
20 was added PtO_2 (1.74 mg, 7.67 μmol). It was purged with N_2 , and then charged with H_2 balloon. The mixture was stirred at rt overnight. It was filtered through a pad of celite. Solvent was removed and the residue was used in the next step without further purification. LCMS (ES): m/z 500.5 $[\text{M}+\text{H}]^+$.

Example 1: To a solution of Intermediate **E1D** (16 mg, 0.032 mmol) in ethanol
25 (582 μL) was added NaOH (aq, 1 N, 96 μL , 0.096 mmol) and the mixture was stirred at room temperature for two hours. It was neutralized with AcOH (0.1 mL). The solvent was removed under reduced pressure and the residue was purified *via* preparative LC/MS (Column: XBridge C18, 19 x 200 mm, 5- μm particles; Mobile Phase A: 5:95 acetonitrile: water with 0.1% TFA; Mobile Phase B: 95:5 acetonitrile: water with 0.1%
30 TFA; Gradient: 10-50% B over 20 minutes, then a 5-minute hold at 100% B; Flow: 20 mL/min) to give Example 1 (4.8 mg, 30%). ^1H NMR (500 MHz, $\text{DMSO}-d_6$) δ 8.20 (d, J

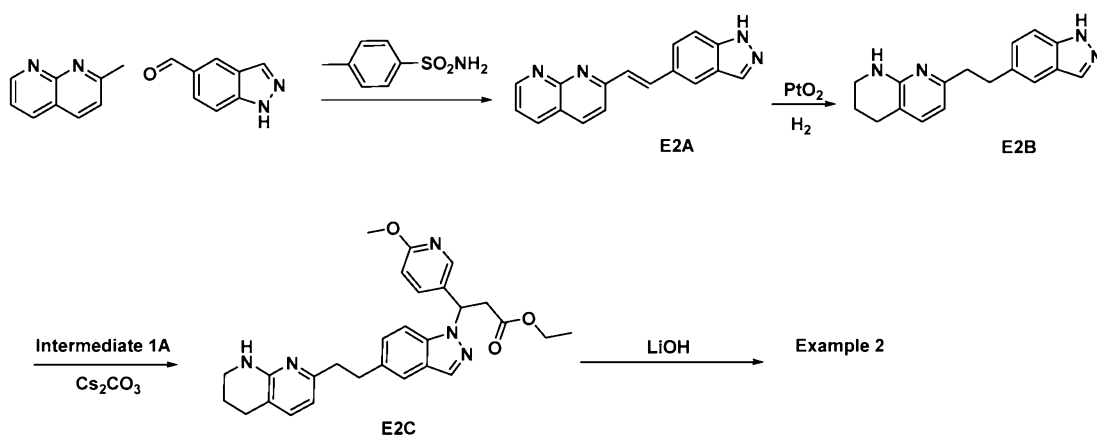
= 2.5 Hz, 1H), 8.12 (s, 1H), 7.67 (dd, $J = 8.8, 2.5$ Hz, 1H), 7.56 (d, $J = 8.6$ Hz, 1H), 7.25 (t, $J = 7.7$ Hz, 1H), 7.03 (d, $J = 7.3$ Hz, 1H), 6.89 (d, $J = 7.0$ Hz, 1H), 6.71 (d, $J = 8.6$ Hz, 1H), 6.26 (d, $J = 7.3$ Hz, 1H), 6.15 (dd, $J = 9.4, 5.3$ Hz, 1H), 3.74 (dt, $J = 14.7, 7.1$ Hz, 2H), 3.54 (dd, $J = 16.5, 9.4$ Hz, 1H), 3.27 – 3.07 (m, 3H), 2.84 (t, $J = 7.7$ Hz, 2H),
 5 2.56 (q, $J = 7.8, 7.0$ Hz, 2H), 2.45 (t, $J = 7.7$ Hz, 2H), 1.91 (dd, $J = 14.3, 6.5$ Hz, 2H), 1.77 – 1.62 (m, 2H). LC/MS (m/z) = 472.0 ($M+H$)⁺. Human α V β 6 IC₅₀ (nM) = 1600.

Example 2

3-(6-Methoxypyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethyl)-
 10 1H-indazol-1-yl)propanoic acid



Example 2



Intermediate **E2A**: A mixture of 2-methyl-1,8-naphthyridine (250 mg, 1.734 mmol), 1H-indazole-5-carbaldehyde (253 mg, 1.734 mmol) and 4-methylbenzenesulfonamide (297 mg, 1.734 mmol) in toluene (4 mL) was heated at 110
 15 °C overnight. The reaction was cooled to rt and diluted with EtOAc (15 mL). The solid was collected via filtering and rinsed with EtOAc (2 x 2 mL) and dried in vacuum to

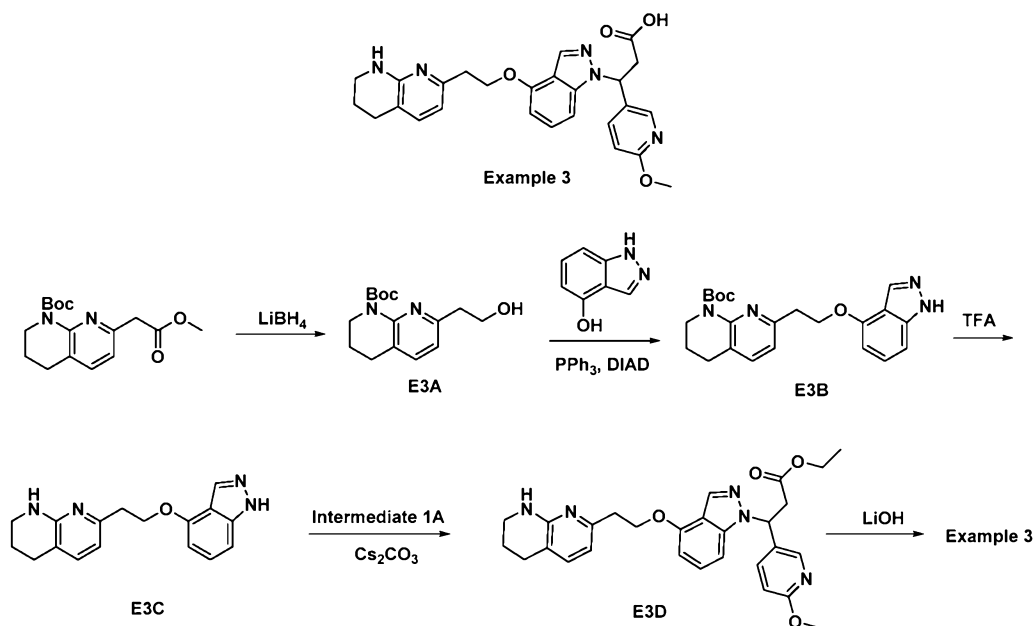
afford Intermediate **E2A** (415 mg, 1.524 mmol, 88% yield). The crude product was used in the next reaction without further purification. LCMS (ES): m/z 273.2 $[M+H]^+$.

Intermediate **E2B**: To a degassed solution of Intermediate **E2A** (200 mg, 0.734 mmol) in EtOH (5 mL) was added PtO_2 (33.4 mg, 0.147 mmol). The mixture was
5 charged with a H_2 balloon and stirred at rt overnight. The reaction was filtered and concentrated to obtain Intermediate **E2B** (203 mg, 0.729 mmol, 99% yield), and the product was used in the next reaction. LCMS (ES): m/z 279.2 $[M+H]^+$.

Intermediate **E2C**: To a solution of Intermediate **E2B** (25 mg, 0.09 mmol) in acetonitrile (0.5 mL) was added cesium carbonate (29.3 mg, 0.09 mmol). After stirring at
10 rt for 5 min, Intermediate **1A** (18.61 mg, 0.09 mmol) was added. The resulting mixture was stirred at 80 °C overnight. The mixture was cooled to rt, filtered, and concentrated. The residue was purified *via* preparative HPLC (Phenomenex Luna Axia 5 μ C18 30 x 100 mm; 10 min gradient from 75% A: 25% B to 0% A:100% B (A = 90% H_2O /10% MeOH + 0.1% TFA); (B = 90% MeOH/10% H_2O + 0.1% TFA); detection at 220 nm) to
15 give Intermediate **E2C** (19 mg, 0.039 mmol, 43.6% yield). LCMS (ES): m/z 486.4 $[M+H]^+$.

Example 2: To a solution of Intermediate **E2C** (19 mg, 0.039 mmol) in THF (0.5 mL) was added a solution of LiOH (aqueous, 1 N, 0.12 mL, 0.12 mmol). After stirring at
rt for 5 hrs, the mixture was neutralized with TFA (50 μ L), filtered, and concentrated.
20 The residue was purified *via* preparative LC/MS with the following conditions: Column: XBridge C18, 19 x 200 mm, 5- μ m particles; Mobile Phase A: 5:95 acetonitrile: water with 10-mM ammonium acetate; Mobile Phase B: 95:5 acetonitrile: water with 10-mM ammonium acetate; Gradient: 10-50% B over 30 minutes, then a 5-minute hold at 100% B; Flow: 20 mL/min to yield Example 2 (5.5 mg, 30% yield). LCMS (ES): m/z 458.2
25 $[M+H]^+$. 1H NMR (500MHz, DMSO- d_6) δ 8.23 (br. s., 1H), 8.00 (s, 1H), 7.67 (t, J = 8.5 Hz, 2H), 7.51 (s, 1H), 7.24 (d, J = 8.6 Hz, 1H), 7.01 (d, J = 7.3 Hz, 1H), 6.72 (d, J = 8.6 Hz, 1H), 6.28 (d, J = 7.2 Hz, 1H), 6.17 (br. s., 1H), 3.76 (s, 3H), 3.23 (br. s., 2H), 3.16 (br. s., 2H), 2.98 - 2.88 (m, 2H), 2.72 (t, J = 7.8 Hz, 2H), 2.58 (t, J = 5.8 Hz, 2H), 1.73 (br. s., 2H). Human $\alpha V\beta 6$ IC₅₀ (nM) = 110.

30

Example 3**3-(6-Methoxypyridin-3-yl)-3-(4-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid**

5 Intermediate **E3A**: To a solution of *tert*-butyl 7-(2-methoxy-2-oxoethyl)-3,4-dihydro-1,8-naphthyridine-1(2H)-carboxylate (1 g, 3.26 mmol) in THF (20 mL) was added a solution of lithium borohydride (2 M, 2.12 mL, 4.24 mmol) in THF. The reaction was stirred at rt overnight. Water (15 mL) was added slowly to the reaction. After stirring at rt for 10 min, the mixture was diluted with EtOAc (12 mL), and extracted with EtOAc

10 (3 x 8 mL). The combine organic layers were dried (Na₂SO₄), filtered, and concentrated. The residue was purified *via* flash chromatography (SiO₂) to give Intermediate **E3A** (782 mg, 86%). LCMS (ES): *m/z* 279.1 [M+H]⁺.

Intermediate **E3B**: To a solution of Intermediate **E3A** (340 mg, 1.221 mmol), 1H-indazol-4-ol (164 mg, 1.22 mmol), and Ph₃P (336 mg, 1.283 mmol) in THF (10 mL) was

15 added DIAD (0.249 mL, 1.283 mmol) slowly. The reaction was stirred at rt for 3 hrs. The mixture was washed with NaHCO₃ solution (aqueous, saturated, 10 mL), the aqueous layer was back extracted with EtOAc (3 x 5 mL). The combine organic layers were washed with brine (10 mL), and then dried over Na₂SO₄. The mixture was filtered and

concentrated under reduced pressure. The residue was purified via flash chromatography to give Intermediate **E3B** (178 mg, 37%). LCMS (ES): m/z 395.3 $[M+H]^+$.

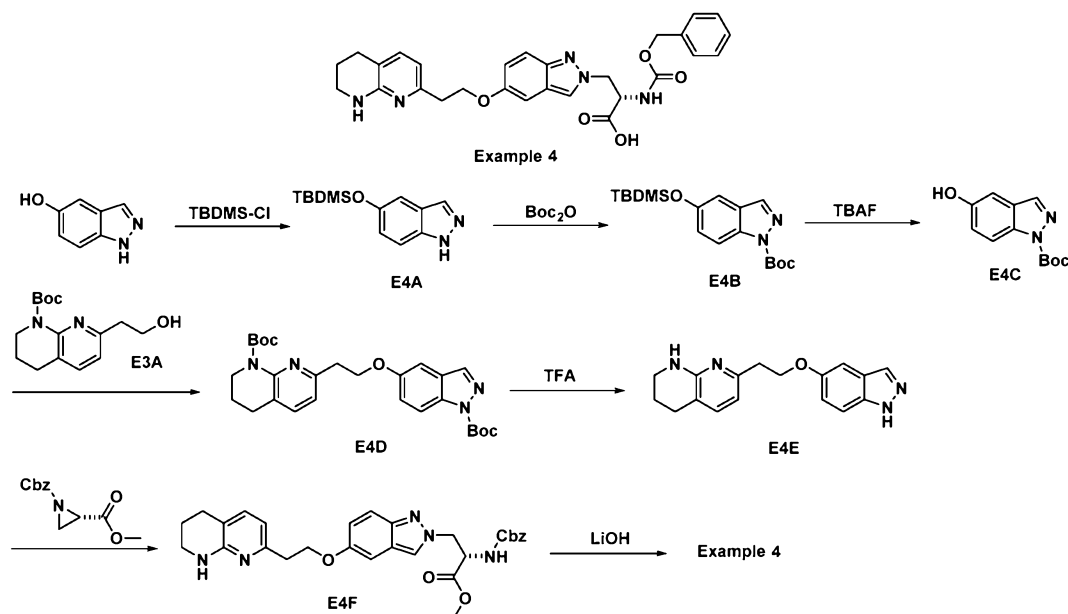
Intermediate **E3C**: To a solution of Intermediate **E3B** (178 mg, 0.451 mmol) in DCM (3 mL) was added TFA (0.174 mL, 2.256 mmol) and the mixture was stirred at rt
5 for 5 hrs. It was concentrated and the crude product was used in the next step without further purification. LCMS (ES): m/z 295.2 $[M+H]^+$.

Intermediate **E3D**: To a solution of Intermediate **E3C** (20 mg, 0.038 mmol) in acetonitrile (0.5 mL) was added cesium carbonate (37.4 mg, 0.115 mmol). After stirring at rt for 5 min, Intermediate **1A** (7.93 mg, 0.038 mmol) was added and the resulting
10 mixture was stirred at 80 °C for 4 hrs. The mixture was cooled to rt, filtered, and concentrated. The residue purified *via* preparative HPLC (Phenomenex Luna Axia 5 μ C18 30 x 100 mm; 10 min gradient from 75% A: 25% B to 0% A:100% B (A = 90% H₂O/10% MeOH + 0.1% TFA); (B = 90% MeOH/10% H₂O + 0.1% TFA); detection at 220 nm) to give Intermediate **E3D** (6 mg, 0.012 mmol, 31.2% yield). LCMS (ES): m/z
15 502.1 $[M+H]^+$.

Example 3: To a solution of Intermediate **E3D** (6 mg, 0.012 mmol) in THF (0.5 mL) was added a solution of LiOH (aqueous, 1 M, 0.036 mL, 0.036 mmol). The reaction mixture was stirred at rt for 6 hrs. The reaction mixture was neutralized with TFA (25 μ L), filtered, and concentrated under reduced pressure. The residue was purified *via*
20 preparative LC/MS with the following conditions: Column: XBridge C18, 19 x 200 mm, 5- μ m particles; Mobile Phase A: 5:95 acetonitrile: water with 10-mM ammonium acetate; Mobile Phase B: 95:5 acetonitrile: water with 10-mM ammonium acetate; Gradient: 10-50% B over 30 minutes, then a 5-minute hold at 100% B; Flow: 20 mL/min to yield Example 3 (5.8 mg, 102% yield). ¹H NMR (500MHz, DMSO-*d*₆) δ 8.22 (br. s., 1H), 8.04 (s, 1H), 7.68 - 7.62 (m, 1H), 7.59 (d, J = 7.3 Hz, 1H), 7.35 (d, J = 8.5 Hz, 1H), 7.29 - 7.23 (m, 1H), 6.74 (dd, J = 13.7, 8.0 Hz, 2H), 6.58 (d, J = 7.7 Hz, 1H), 6.15 (dd, J = 9.8, 5.1 Hz, 1H), 4.42 - 4.32 (m, 2H), 3.76 (s, 2H), 3.39 (br. s., 1H), 3.16 (s, 5H), 2.71 (br. s., 2H), 1.80 (br. s., 2H). LCMS (ES): m/z 473.9 $[M+H]^+$. Human α V β 6 IC₅₀ (nM) = 2300.

Example 4

(*S*)-2-(((Benzyloxy)carbonyl)amino)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-2*H*-indazol-2-yl)propanoic acid



- 5 Intermediate **E4A**: A solution of 1*H*-indazol-5-ol (5.1 g, 38.0 mmol), *tert*-butylchlorodimethylsilane (16.8 g, 111 mmol) and imidazole (12.7 g, 187 mmol) in DCM (200 ml) was stirred at rt overnight. The mixture was diluted with brine (60 mL) and extracted with DCM (3 x 50 mL). The combined organic layers were washed with water (50 mL), then brine (50 mL). It was dried (Na₂SO₄), filtered, and concentrated. The
- 10 residue was purified via flash column (SiO₂) chromatography to afford E4A as a light yellow solid (7.48 g, 30.1 mmol, yield 79%). ¹H NMR (400 MHz, chloroform-*d*) δ 10.27 (s, 1H), 7.99 (d, *J* = 1.1 Hz, 1H), 7.38 (dt, *J* = 8.9, 0.9 Hz, 1H), 7.17 – 7.14 (m, 1H), 7.01 (dd, *J* = 8.9, 2.2 Hz, 1H), 1.04 (s, 9H), 0.24 (s, 6H).

- Intermediate **E4B**: To a solution of **E4A** (5-((*tert*-butyldimethylsilyl)oxy)indazole) (5.56 g, 22.38 mmol) in DCM (100 mL) was added Boc₂O (5.72 mL, 24.62 mmol) followed by DMAP (0.547 g, 4.48 mmol) and Et₃N (3.43 mL, 24.62 mmol). The mixture was stirred at rt overnight. The mixture was concentrated and residue was purified via flash column chromatography (silica gel, hexanes/EtOAc gradient 0 to 25% EtOAc) to give E4B (7.74 g, 22.21 mmol, 99% yield) as a mixture of

isomers: *tert*-butyl 5-((*tert*-butyldimethylsilyl)oxy)indazole-1-carboxylate and *tert*-butyl 5-((*tert*-butyldimethylsilyl)oxy)-2*H*-indazole-2-carboxylate.

Intermediate **E4C**: To a solution of Intermediate **E4B** (*tert*-butyl 5-((*tert*-butyldimethylsilyl)oxy)indazole-1-carboxylate) (7.74 g, 22.21 mmol) in THF (100 mL) was added a solution of TBAF (44.4 mL, 44.4 mmol). The mixture was stirred at rt for 2 hrs. The mixture was diluted with NH₄Cl (aqueous, saturated, 30 mL) and extracted with EtOAc (3 x 25 mL). The combine organic layers were washed with brine (30 mL) and dried (Na₂SO₄), filtered, and concentrated. The residue was purified *via* flash column chromatography to afford Intermediate **E4C** (3.42 g, 14.60 mmol, 65.7% yield) as a mixture of two regioisomers: *tert*-butyl 5-hydroxyindazole-1-carboxylate and *tert*-butyl 5-hydroxy-2*H*-indazole-2-carboxylate.

Intermediate **E4D**: To a solution of **E4C** (3.53 g, 12.68 mmol) and Ph₃P (3.78 g, 14.41 mmol) in THF (70 mL) at 0 °C was added **E3A** (*tert*-butyl 5-hydroxyindazole-1-carboxylate) (2.7 g, 11.53 mmol) followed by (*E*)-diazene-1,2-diylbis(piperidin-1-ylmethanone) (3.64 g, 14.41 mmol). The reaction was stirred and gradually warmed to rt overnight. A solution of NaHCO₃ (aqueous, saturated, 25 mL) was added, and the mixture was extracted with EtOAc (3 x 20 mL). The combined organic layers were washed with water (20 mL) then with brine (20 mL). It was dried (Na₂SO₄), filtered, and concentrated under reduced pressure. The residue was purified *via* flash column chromatography to afford Intermediate **E4D** as a light yellow solid (3.94 g, 7.97 mmol, 69.1% yield). LCMS (ES): *m/z* 495.1 [M+H]⁺.

Intermediate **E4E**: To a solution of **E4D** (3.94 g, 7.97 mmol) in DCM (40 mL) at 0 °C was added TFA (8 mL, 104 mmol). The mixture was gradually warmed to rt and stirred at rt overnight. The reaction was monitored by LCMS. When necessary, additional TFA was added. After completion, the mixture was concentrated under reduced pressure and the residue was purified *via* flash column chromatography (C18 column, 10% ACN in water with 0.1% TFA to 80% ACN in water, 12 min gradient) to give **E4E** (7-(2-((1*H*-indazol-5-yl)oxy)ethyl)-1,2,3,4-tetrahydro-1,8-naphthyridine TFA salt) (2.63 g, 6.44 mmol, 81% yield) as light yellow solid. ¹H NMR (500 MHz, chloroform-*d*) δ 15.60 (s, 1H), 10.31 (s, 1H), 8.01 (s, 1H), 7.40 (d, *J* = 8.9 Hz, 1H), 7.34 (d, *J* = 7.3 Hz, 1H), 7.15 (d, *J* = 2.3 Hz, 1H), 7.11 – 6.99 (m, 1H), 6.53 (d, *J* = 7.2 Hz, 1H), 4.34 (t, *J* = 5.8 Hz,

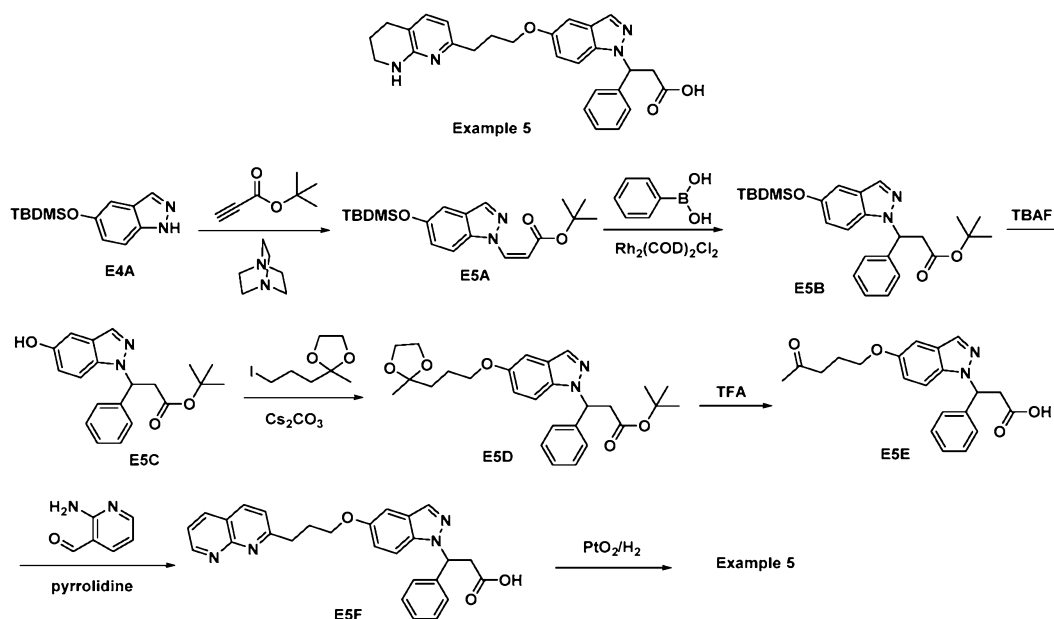
2H), 3.51 (d, $J = 6.0$ Hz, 2H), 3.22 (t, $J = 5.9$ Hz, 2H), 2.77 (t, $J = 6.3$ Hz, 2H), 1.94 (q, $J = 5.9$ Hz, 2H). LCMS (ES): m/z 295.2 $[M+H]^+$.

Intermediate **E4F**: A mixture of Intermediate **E4E** (37 mg, 0.091 mmol) and (*S*)-1-benzyl 2-methyl aziridine-1,2-dicarboxylate (27.7 mg, 0.118 mmol) in toluene (1 mL) was heated at 110 °C overnight. The solution was concentrated and the crude product was purified *via* preparative HPLC (Column: XBridge C18, 19 x 200 mm, 5- μ m particles; Mobile Phase A: 5:95 acetonitrile: water with 10-mM ammonium acetate; Mobile Phase B: 95:5 acetonitrile: water with 10-mM ammonium acetate; Gradient: 12-42% B over 25 minutes, then a 2-minute hold at 42% B; Flow: 20 mL/min.) to give Intermediate **E4F** (16 mg, 0.030 mmol, 33.3% yield). LCMS (ES): m/z 530.0 $[M+H]^+$.

Example 4: To a solution of Intermediate **E4F** (16 mg, 0.030 mmol) in THF (0.5 mL) was added a solution of LiOH (aqueous, 1 M, 0.091 mL, 0.091 mmol). The mixture was stirred at rt for 2 hrs. The mixture was neutralized with TFA (20 μ L) and concentrated under reduced pressure. The residue was purified *via* preparative LC/MS with the following conditions: Column: XBridge C18, 19 x 200 mm, 5- μ m particles; Mobile Phase A: 5:95 acetonitrile: water with 10-mM ammonium acetate; Mobile Phase B: 95:5 acetonitrile: water with 10-mM ammonium acetate; Gradient: 10-60% B over 20 minutes, then a 5-minute hold at 100% B; Flow: 20 mL/min to give Example 4 (11.3 mg, 73%). ^1H NMR (400 MHz, MeOH- d_4) δ 7.87 (s, 1H), 7.44 – 7.33 (m, 2H), 7.22 (d, $J = 9.2$ Hz, 5H), 6.85 (d, $J = 8.9$ Hz, 2H), 6.58 (d, $J = 7.3$ Hz, 1H), 5.07 – 4.89 (m, 2H), 4.88 – 4.65 (m, 2H), 4.58 (s, 1H), 4.16 (t, $J = 6.3$ Hz, 2H), 3.41 (t, $J = 5.7$ Hz, 2H), 3.02 (t, $J = 6.3$ Hz, 2H), 2.73 (t, $J = 6.3$ Hz, 2H), 1.92 – 1.82 (m, 2H). LCMS (ES): m/z 516.3 $[M+H]^+$. Human $\alpha\text{V}\beta 6$ IC₅₀ (nM) = 21.

Example 5

3-Phenyl-3-(5-(3-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)propoxy)-1*H*-indazol-1-yl)propanoic acid



Intermediate **E5A**: To a solution of **E4A** (0.745 g, 3.00 mmol) in DCM (6 mL) at 0 °C under argon was added DABCO (0.034 g, 0.300 mmol), followed by slow addition of *tert*-butyl propiolate (0.51 mL, 3.6 mmol). The mixture was stirred at this temperature for additional 20 min, then gradually warmed to rt overnight. The mixture was acidified with HOAc (34 μ L). The solvent was removed under reduced pressure and the residue was purified via flash column chromatography (SiO₂, hexanes/EtOAc gradient 0 to 40% EtOAc) to give **E5B** (0.19 g, 17% yield). ¹H NMR (500 MHz, chloroform-*d*) δ 8.03 (d, *J* = 0.9 Hz, 1H), 7.27 (d, *J* = 8.8 Hz, 1H), 7.21 (d, *J* = 9.6 Hz, 1H), 7.12 (d, *J* = 2.2 Hz, 1H), 7.03 (dd, *J* = 8.9, 2.2 Hz, 1H), 5.62 (d, *J* = 9.5 Hz, 1H), 1.47 (s, 9H), 1.03 (s, 9H), 0.23 (s, 6H).

Intermediate **E5B**: To a solution of **E5A** (46.2 mg, 0.123 mmol), Et₃N (103 μ L, 0.740 mmol), and phenylboronic acid (30.1 mg, 0.247 mmol) in MeOH (617 μ L) was added chloro(1,5-cyclooctadiene)rhodium(I) dimer (3.04 mg, 6.17 μ mol). The mixture was degassed for 10 min, and heated at 60 °C overnight. The solvent was removed and the residue was purified via flash column chromatography (SiO₂) to give **E5B** (39 mg, 70% yield). ¹H NMR (500 MHz, chloroform-*d*) δ 7.93 (d, *J* = 0.9 Hz, 1H), 7.40 - 7.22 (m, 6H), 7.08 (d, *J* = 2.2 Hz, 1H), 6.92 (dd, *J* = 8.9, 2.3 Hz, 1H), 6.40 (d, *J* = 13.7 Hz,

0H), 6.03 (dd, $J = 9.6, 5.6$ Hz, 1H), 3.65 (dd, $J = 15.9, 9.6$ Hz, 1H), 3.18 (dd, $J = 15.9, 5.7$ Hz, 1H), 1.26 (s, 9H), 1.01 (s, 9H), 0.20 (d, $J = 1.7$ Hz, 6H).

Intermediate **E5C**: To a solution of Intermediate **E5B** (80 mg, 0.177 mmol) in CH_2Cl_2 (353 μL) was added TBAF (265 μL , 0.265 mmol). The mixture was stirred at rt for 3 hrs. It was neutralized with HOAc (150 μL). Solvent was removed and the residue was purified via flash column chromatography (SiO_2) to give Intermediate **E5C** (51 mg, 92% yield). ^1H NMR (400 MHz, chloroform- d) δ 7.88 (d, $J = 0.9$ Hz, 1H), 7.36 - 7.18 (m, 7H), 7.10 - 6.94 (m, 1H), 6.89 (dd, $J = 9.0, 2.3$ Hz, 1H), 6.02 (dd, $J = 9.7, 5.6$ Hz, 1H), 3.65 (dd, $J = 15.9, 9.8$ Hz, 1H), 3.16 (dd, $J = 16.0, 5.6$ Hz, 1H), 1.26 (s, 9H).

Intermediate **E5D**: To a solution of Intermediate **E5C** (32 mg, 0.095 mmol) and 2-(3-iodopropyl)-2-methyl-1,3-dioxolane (36.3 mg, 0.142 mmol) in acetonitrile (946 μL) was added Cs_2CO_3 (92 mg, 0.284 mmol). The mixture was stirred at rt overnight. The solvent was removed and the residue was purified via flash column chromatography (SiO_2) to give **E5D** (37 mg, 84% yield). LCMS (ES): m/z 467.3 $[\text{M}+\text{H}]^+$.

Intermediate **E5E**: To a solution of Intermediate **E5D** (37 mg, 0.079 mmol) in DCM (227 μL) was added TFA (566 μL). The mixture was stirred at rt for 4 hrs. The solvent was removed under reduced pressure and the residue was used in the next reaction without further purification. LCMS (ES): m/z 367.1 $[\text{M}+\text{H}]^+$.

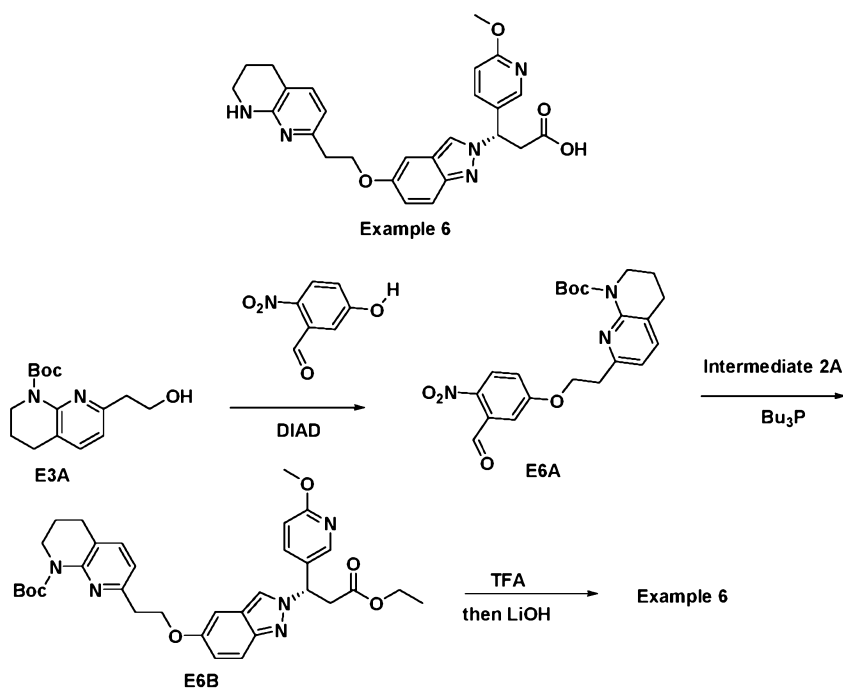
Intermediate **E5F**: To a solution of Intermediate **E5E** (29 mg, 0.079 mmol) in DCM (198 μL) and MeOH (594 μL) was added pyrrolidine (13.1 μL , 0.158 mmol). The mixture was stirred at rt for 15 min. Then 2-aminonicotinaldehyde (11.60 mg, 0.095 mmol) was added. The mixture was stirred at rt overnight. The solvent was removed and the residue was purified *via* preparative HPLC (Phenomenex Luna Axia 5 μ C18 30 x 100 mm; 10 min gradient from 85% A: 15% B to 0% A:100% B (A = 90% H_2O /10% ACN + 0.1% TFA); (B = 90% ACN/10% H_2O + 0.1% TFA); detection at 220 nm) to yield Intermediate **E5F** (11.3 mg, 36% yield). ^1H NMR (500 MHz, MeOH- d_4) δ 9.11 (s, 1H), 8.90 (dd, $J = 8.2, 1.7$ Hz, 1H), 8.80 (d, $J = 8.5$ Hz, 1H), 8.00 (d, $J = 8.5$ Hz, 1H), 7.95 (dd, $J = 8.2, 4.8$ Hz, 1H), 7.91 (d, $J = 0.7$ Hz, 1H), 7.36 - 7.22 (m, 6H), 7.06 (d, $J = 2.2$ Hz, 1H), 6.67 (dd, $J = 9.1, 2.3$ Hz, 1H), 6.12 (dd, $J = 9.9, 5.1$ Hz, 1H), 4.18 (t, $J = 5.7$ Hz, 2H), 3.71 (dd, $J = 16.7, 9.9$ Hz, 1H), 3.44 (t, $J = 7.2$ Hz, 2H), 3.24 (dd, $J = 16.6, 5.1$ Hz, 1H), 2.54 - 2.43 (m, 2H).

Example 5: To a solution of Intermediate **E5F** (11.3 mg, 0.025 mmol) in MeOH (675 μ L) was added sodium bicarbonate (4.20 mg, 0.050 mmol), followed by PtO₂ (1.134 mg, 4.99 μ mol). The mixture was charged with H₂ balloon. It was stirred at room temperature overnight. The mixture was filtered through a pad of celite. The filtrate was purified *via* preparative LC/MS with the following conditions: Column: XBridge C18, 19 x 200 mm, 5- μ m particles; Mobile Phase A: 5:95 acetonitrile: water with 10-mM ammonium acetate; Mobile Phase B: 95:5 acetonitrile: water with 10-mM ammonium acetate; Gradient: 15-55% B over 15 minutes, then a 3-minute hold at 100% B; Flow: 20 mL/min to give Example 5 (3.9 mg, 33%). LCMS (ES): m/z 457.1 [M+H]⁺. ¹H NMR (500 MHz, MeOH-*d*₄) δ 7.95 (d, *J* = 0.8 Hz, 1H), 7.55 (d, *J* = 7.4 Hz, 1H), 7.46 (d, *J* = 9.1 Hz, 1H), 7.30 (d, *J* = 4.3 Hz, 4H), 7.28 - 7.23 (m, 1H), 7.13 (d, *J* = 2.3 Hz, 1H), 6.94 (dd, *J* = 9.1, 2.3 Hz, 1H), 6.64 (d, *J* = 7.3 Hz, 1H), 6.18 (dd, *J* = 9.9, 5.1 Hz, 1H), 4.07 (t, *J* = 5.7 Hz, 2H), 3.72 (dd, *J* = 16.6, 9.9 Hz, 1H), 3.46 (td, *J* = 5.3, 2.3 Hz, 2H), 3.25 (dd, *J* = 16.6, 5.0 Hz, 1H), 2.92 (t, *J* = 7.5 Hz, 2H), 2.80 (t, *J* = 6.3 Hz, 2H), 2.20 (dq, *J* = 13.2, 6.2 Hz, 2H), 1.94 (p, *J* = 6.1 Hz, 2H). Human α V β 6 IC₅₀ (nM) = 440.

Example 6

(*S*)-3-(6-Methoxypyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-2*H*-indazol-2-yl)propanoic acid

20



- Intermediate **E6A**: To a solution of Ph_3P (104 mg, 0.396 mmol) and Intermediate **E3A** (97 mg, 0.349 mmol) in THF (1.86 mL) at 0 °C was added 5-hydroxy-2-nitrobenzaldehyde (53 mg, 0.317 mmol), followed by DIAD (77 μL , 0.396 mmol). The mixture was stirred under argon and gradually warmed to rt overnight. The reaction was diluted with EtOAc (10 mL), washed with NaHCO_3 (aqueous, saturated, 8 mL). The aqueous layer was extracted with EtOAc (3 x 6 mL). The combined organic layers were washed with water (8 mL), then with brine (8 mL). It was dried (Na_2SO_4), filtered, and concentrated under reduced pressure. The residue was purified *via* flash column chromatography (SiO_2 , hexanes/EtOAc gradient 0 to 100% EtOAc) to give Intermediate **E6A** (73.8 mg, 0.173 mmol, 54.4% yield). ^1H NMR (500 MHz, chloroform-*d*) δ 10.49 (s, 1H), 8.16 (d, $J = 9.1$ Hz, 1H), 7.38 - 7.33 (m, 2H), 7.19 (dd, $J = 9.1, 2.9$ Hz, 1H), 6.92 (d, $J = 7.5$ Hz, 1H), 4.55 (t, $J = 6.7$ Hz, 2H), 3.84 - 3.73 (m, 2H), 3.25 (t, $J = 6.7$ Hz, 2H), 2.77 (t, $J = 6.7$ Hz, 2H), 1.95 (p, $J = 6.5$ Hz, 2H), 1.53 (s, 9H).
- Intermediate **E6B**: To a solution of Intermediate **2A** (24.82 mg, 0.111 mmol) in 2-propanol (234 μL) was added **E6A** (43 mg, 0.101 mmol). The mixture was heated at 80 °C for 4 hours. It was cooled to rt. PBu_3 (74.5 μL , 0.302 mmol) was added in one portion. The mixture was heated at 80 °C for 16 hrs. The mixture was cooled to rt, diluted with

EtOAc (5 mL), then washed with ammonium chloride (5 mL), followed by brine (5 mL). The organic layer was dried (MgSO₄), filtered, and concentrated under reduced pressure. The residue was purified *via* flash column chromatography (SiO₂, 0-100% EtOAc/heptane) to afford Intermediate **E6B** (16 mg, 27%). LCMS (ES): m/z 602.8

5 [M+H]⁺.

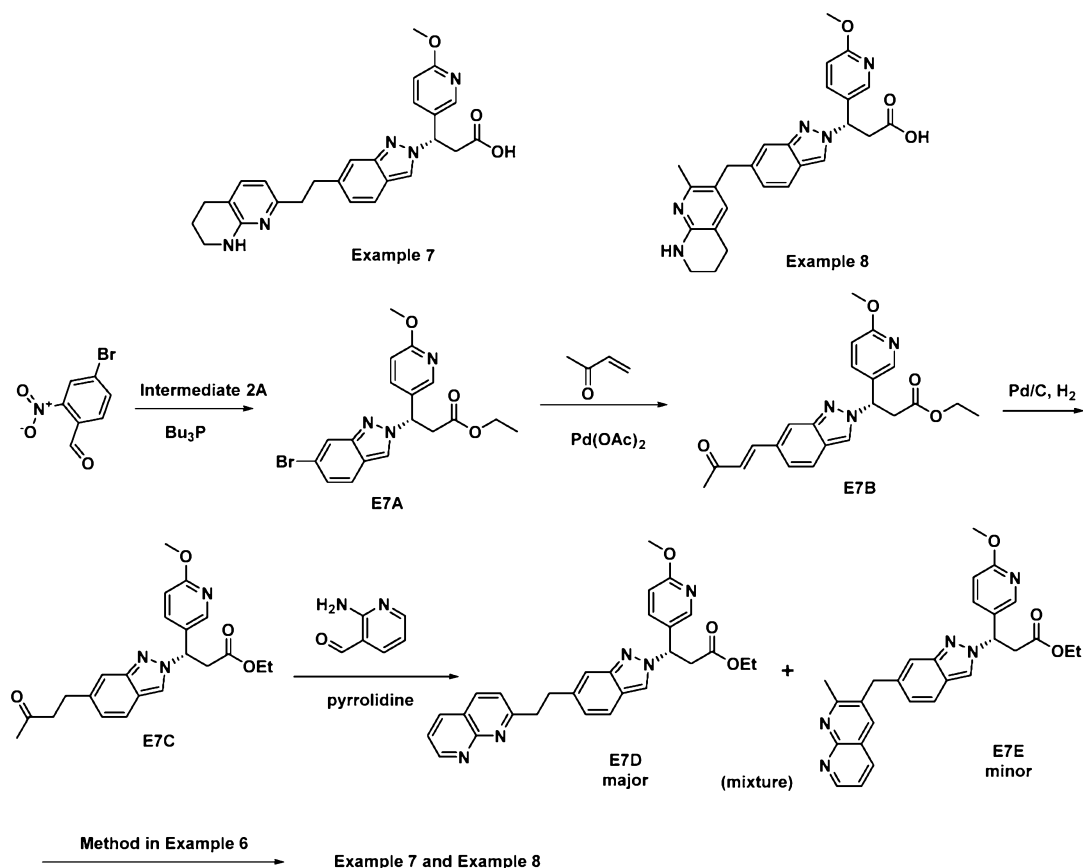
Example 6: To a solution of Intermediate **E6B** (16.3 mg, 0.027 mmol) in CH₂Cl₂ (342 μL) was added TFA (68.4 μL). The mixture was stirred at rt overnight. The solvent was removed under reduced pressure. The above residue was dissolved in THF (342 μL), LiOH (aqueous, 1 N, 81 μL, 0.081 mmol) was added. The mixture was stirred at rt
10 overnight. It was neutralized with HCl (aqueous, 1 N, 100 μL) and purified *via* preparative LC/MS with the following conditions: Column: XBridge C18, 19 x 200 mm, 5-μm particles; Mobile Phase A: 5:95 acetonitrile: water with 10-mM ammonium acetate; Mobile Phase B: 95:5 acetonitrile: water with 10-mM ammonium acetate; Gradient: 3-40% B over 25 minutes, then a 5-minute hold at 100% B; Flow: 20 mL/min to give
15 Example 6 (5 mg, 37%). ¹H NMR (500 MHz, MeOH-*d*₄) δ 8.18 (d, *J* = 0.9 Hz, 1H), 8.15 (d, *J* = 2.6 Hz, 1H), 7.71 (dd, *J* = 8.7, 2.6 Hz, 1H), 7.44 (dt, *J* = 9.0, 1.0 Hz, 1H), 7.30 (dt, *J* = 7.4, 1.1 Hz, 1H), 6.89 - 6.85 (m, 2H), 6.73 (dd, *J* = 8.7, 0.7 Hz, 1H), 6.53 (d, *J* = 7.3 Hz, 1H), 6.10 (dd, *J* = 9.1, 6.1 Hz, 1H), 4.13 (td, *J* = 6.5, 4.2 Hz, 2H), 3.86 (s, 3H), 3.49 (dd, *J* = 16.0, 9.2 Hz, 1H), 3.38 (dd, *J* = 6.5, 4.7 Hz, 2H), 3.18 (dd, *J* = 16.0, 6.1 Hz,
20 1H), 2.98 (t, *J* = 6.3 Hz, 2H), 2.71 (t, *J* = 6.3 Hz, 2H), 1.91 - 1.81 (m, 2H). LCMS (ES): m/z 474.0 [M+H]⁺. Human αVβ6 IC₅₀ (nM) = 600.

Example 7

(*S*)-3-(6-Methoxypyridin-3-yl)-3-(6-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-
25 yl)ethyl)-2*H*-indazol-2-yl)propanoic acid
and

Example 8

(S)-3-(6-Methoxypyridin-3-yl)-3-(6-((2-methyl-5,6,7,8-tetrahydro-1,8-naphthyridin-3-yl)methyl)-2H-indazol-2-yl)propanoic acid



Intermediate **E7A**: To a solution of Intermediate **2A** (350 mg, 1.559 mmol) in 2-propanol (3.3 mL) was added 4-bromo-2-nitrobenzaldehyde (326 mg, 1.417 mmol). The mixture was heated at 80 °C for 4 hrs under argon. It was cooled to rt. PBu₃ (1.1 mL, 4.25 mmol) was added. The mixture was heated at 80 °C for 16 hrs. The mixture was cooled to rt and diluted with EtOAc (5 mL) and washed with ammonium chloride (5 mL), followed by brine (5 mL). The organic layer was dried (MgSO₄), filtered, and concentrated under reduced pressure. The residue was purified *via* flash column chromatography (SiO₂, 0-100% EtOAc/heptane) to afford Intermediate **E7A** (501 mg, 70%). ¹H NMR (400 MHz, chloroform-*d*) δ 8.21 (d, *J* = 2.5 Hz, 1H), 7.95 (d, *J* = 1.0 Hz, 1H), 7.88 (dt, *J* = 1.7, 0.9 Hz, 1H), 7.67 (dd, *J* = 8.7, 2.5 Hz, 1H), 7.47 (dd, *J* = 9.0, 0.8 Hz, 1H), 7.13 (dd, *J* = 8.8, 1.6 Hz, 1H), 6.71 (dd, *J* = 8.6, 0.7 Hz, 1H), 6.01 (dd, *J* = 8.8, 6.0 Hz, 1H), 4.07 (qd, *J* =

7.2, 1.0 Hz, 2H), 3.91 (s, 3H), 3.75 (dd, $J = 16.6, 8.8$ Hz, 1H), 3.19 (dd, $J = 16.5, 6.0$ Hz, 1H), 1.15 (t, $J = 7.1$ Hz, 3H).

Intermediate **E7B**: A solution of Intermediate **E7A** (0.199 g, 0.492 mmol), but-3-en-2-one (0.142 mL, 1.723 mmol), Et₃N (0.185 mL, 1.329 mmol), Pd(OAc)₂ (12 mg, 0.055 mmol) and tri-*o*-tolylphosphine (0.025 g, 0.082 mmol) in ACN (4 mL) was
5 degassed with argon for 10 min. The mixture was then sealed and heated at 120 °C for 12 hrs. The solvent was removed under reduced pressure, and the residue was purified *via* flash column chromatography (SiO₂) to afford Intermediate **E7B** (0.156 g, 0.397 mmol, 81% yield). ¹H NMR (400 MHz, chloroform-*d*) δ 8.23 (d, $J = 2.4$ Hz, 1H), 7.96 (d, $J =$
10 1.0 Hz, 1H), 7.84 (s, 1H), 7.68 (dd, $J = 8.7, 2.6$ Hz, 1H), 7.62 (d, $J = 5.2$ Hz, 1H), 7.59 (d, $J = 2.2$ Hz, 1H), 7.29 (dd, $J = 8.8, 1.4$ Hz, 1H), 6.79 - 6.70 (m, 2H), 6.04 (dd, $J = 8.7, 6.1$ Hz, 1H), 4.08 (qd, $J = 7.2, 0.8$ Hz, 2H), 3.91 (s, 3H), 3.77 (dd, $J = 16.6, 8.8$ Hz, 1H), 3.22 (dd, $J = 16.5, 6.1$ Hz, 1H), 2.40 (s, 3H), 1.15 (t, $J = 7.1$ Hz, 3H).

Intermediate **E7C**: To a solution of Intermediate **E7B** (0.156 g, 0.397 mmol) in
15 EtOAc (3.97 ml) was added Pd/C (10%, 0.021 g, 0.020 mmol). The mixture was purged with H₂ gas, then charged with H₂ balloon. It was stirred at rt overnight. It was filtered through a pad of celite and rinsed with MeOH. The filtrate was concentrated under reduced pressure to afford **E7C**, which was used in the next reaction without further purification. LCMS (ES): m/z 396.1 [M+H]⁺.

20 Intermediate **E7D** and Intermediate **E7E**: To a solution of Intermediate **E7C** (38.7 mg, 0.098 mmol) in CH₂Cl₂ (245 μ L) and MeOH (734 μ L) was added pyrrolidine (16.19 μ L, 0.196 mmol). After stirring at rt for 15 min, 2-aminonicotinaldehyde (14.34 mg, 0.117 mmol) was added. The mixture was stirred at rt overnight. The solvent was removed and the residue was purified *via* chromatography to afford Intermediates **E7D**
25 and **E7E** as a mixture. Intermediate **E7D**: LCMS (ES): m/z 482.0 [M+H]⁺. Intermediate **E7E**: LCMS (ES): m/z 482.0 [M+H]⁺.

Example 7 and Example 8 were prepared from the mixture of Intermediates **E7D** and **E7E** according to the method described in Example 6.

Example 7: ¹H NMR (500 MHz, MeOH-*d*₄) δ 8.28 (s, 1H), 8.15 (d, $J = 2.5$ Hz, 1H), 7.72 (dd, $J = 8.7, 2.6$ Hz, 1H), 7.53 (d, $J = 8.6$ Hz, 1H), 7.30 (d, $J = 7.3$ Hz, 1H),
30 7.17 (s, 1H), 6.87 (dd, $J = 8.7, 1.4$ Hz, 1H), 6.73 (d, $J = 8.6$ Hz, 1H), 6.37 (d, $J = 7.3$ Hz,

1H), 6.13 (dd, $J = 9.4, 5.9$ Hz, 1H), 3.86 (s, 3H), 3.49 (dd, $J = 15.8, 9.4$ Hz, 1H), 3.39 (t, $J = 5.7$ Hz, 2H), 3.18 (dd, $J = 15.8, 5.9$ Hz, 1H), 2.88 (h, $J = 6.1, 5.0$ Hz, 4H), 2.70 (t, $J = 6.3$ Hz, 2H), 1.86 (p, $J = 6.1$ Hz, 2H). LCMS (ES): m/z 458.3 $[M+H]^+$. Human $\alpha V\beta 6$ IC₅₀ (nM) = 6.6. Example 8: ¹H NMR (500 MHz, MeOH-*d*₄) δ 8.32 (s, 1H), 8.16 (d, $J = 2.5$ Hz, 1H), 7.73 (dd, $J = 8.7, 2.6$ Hz, 1H), 7.59 (d, $J = 8.6$ Hz, 1H), 7.29 (s, 1H), 7.22 (s, 1H), 6.87 (dd, $J = 8.6, 1.4$ Hz, 1H), 6.73 (d, $J = 8.7$ Hz, 1H), 6.12 (dd, $J = 8.9, 6.4$ Hz, 1H), 3.90 (s, 2H), 3.85 (s, 3H), 3.48 - 3.35 (m, 3H), 3.16 (dd, $J = 15.6, 6.4$ Hz, 1H), 2.70 (t, $J = 6.3$ Hz, 2H), 2.28 (s, 3H), 1.92 - 1.82 (m, 2H). LCMS (ES): m/z 458.3 $[M+H]^+$. Human $\alpha V\beta 6$ IC₅₀ (nM) = 390.

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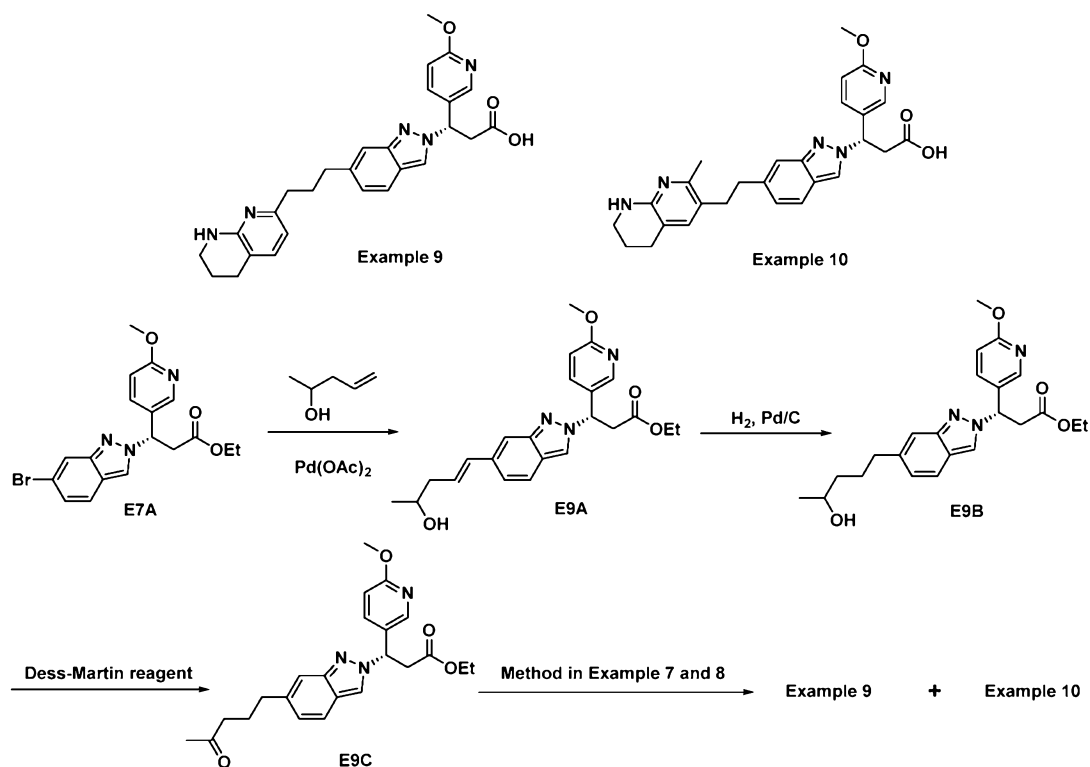
Example 9

(*S*)-3-(6-Methoxypyridin-3-yl)-3-(6-(3-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)propyl)-2*H*-indazol-2-yl)propanoic acid

and

15 **Example 10**

(*S*)-3-(6-Methoxypyridin-3-yl)-3-(6-(2-(2-methyl-5,6,7,8-tetrahydro-1,8-naphthyridin-3-yl)ethyl)-2*H*-indazol-2-yl)propanoic acid



Intermediate **E9A**: A solution of Intermediate **E7A** (0.1295 g, 0.320 mmol), pent-4-en-2-ol (0.117 mL, 1.121 mmol), Et_3N (0.12 mL, 0.865 mmol), $\text{Pd}(\text{OAc})_2$ (8.11 mg, 0.036 mmol) and tri-*o*-tolylphosphine (0.016 g, 0.053 mmol) in ACN (4 mL) was degassed with argon for 10 min. The mixture was sealed and heated at 120 °C for 12 hrs. After cooled to rt, the solvent was removed under reduced pressure, and the residue was purified *via* chromatography to afford Intermediate **E9A** (85 mg, 65%). ^1H NMR (400 MHz, chloroform-*d*) δ 8.21 (d, $J = 2.5$ Hz, 1H), 7.89 (d, $J = 0.9$ Hz, 1H), 7.67 (dd, $J = 8.7, 2.7$ Hz, 1H), 7.59 - 7.48 (m, 2H), 7.21 (dd, $J = 8.8, 1.5$ Hz, 1H), 6.70 (d, $J = 8.6$ Hz, 1H), 6.57 (d, $J = 15.8$ Hz, 1H), 6.25 (ddd, $J = 15.8, 7.8, 6.9$ Hz, 1H), 6.01 (dd, $J = 8.6, 6.3$ Hz, 1H), 4.18 - 4.01 (m, 3H), 3.99 - 3.92 (m, 1H), 3.90 (s, 3H), 3.75 (dd, $J = 16.4, 8.6$ Hz, 1H), 3.21 (dd, $J = 16.5, 6.4$ Hz, 1H), 2.53 - 2.28 (m, 2H), 1.32 - 1.20 (m, 3H), 1.20 - 1.08 (m, 3H).

Intermediate **E9B**: To a solution of Intermediate **E9A** (85.3 mg, 0.208 mmol) in EtOAc (2.1 mL) was added Pd/C (10%, 11.08 mg, 10.42 μmol). The mixture was purged with H_2 gas and then charged with H_2 balloon. The mixture was stirred at rt overnight. It

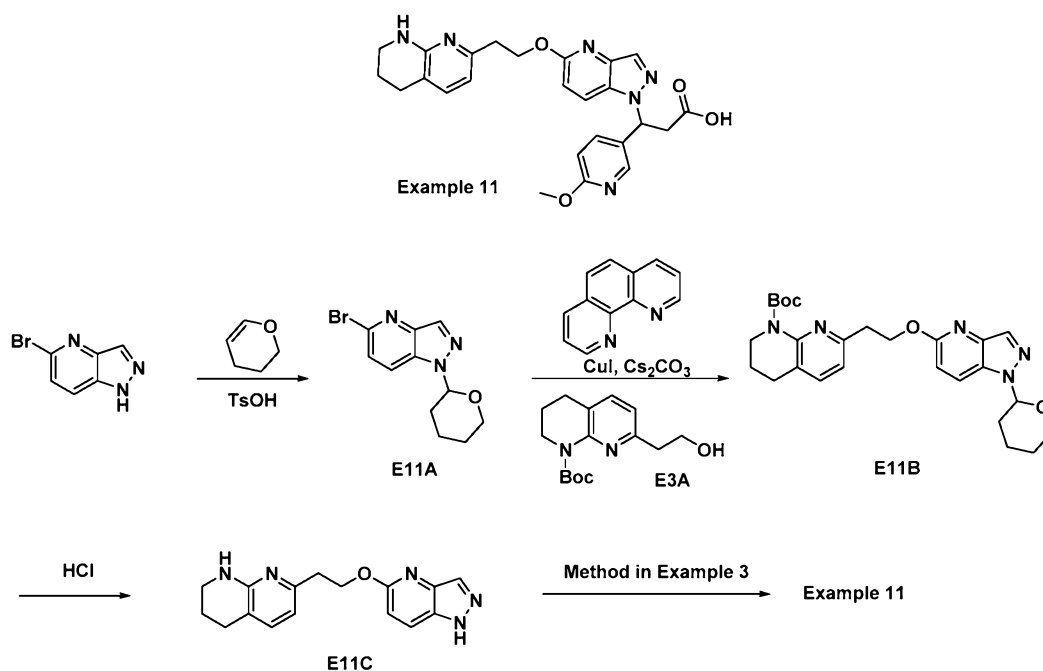
was filtered through a pad of celite and rinsed with MeOH. The filtrate was concentrated and the residue was used in the next reaction without further purification. LCMS (ES): m/z 412.1 $[M+H]^+$.

Intermediate **E9C**: To a solution of Intermediate **E9B** (83 mg, 0.202 mmol) in CH_2Cl_2 (2.1 mL) was added Dess-Martin periodinane (103 mg, 0.242 mmol). After stirred at rt for 1 hr, the mixture was diluted with Et_2O (10 mL), the precipitate was filtered off, and rinsed with Et_2O (10 mL). The filtrate was concentrated under reduced pressure, and the residue was purified *via* flash column chromatography to give Intermediate **E9C** (74 mg, 90%). 1H NMR (500 MHz, chloroform- d) δ 8.23 (d, J = 2.5 Hz, 1H), 7.93 (s, 1H), 7.70 (dd, J = 8.7, 2.5 Hz, 1H), 7.58 - 7.50 (m, 1H), 7.47 (s, 1H), 6.93 (dd, J = 8.5, 1.4 Hz, 1H), 6.72 (d, J = 8.6 Hz, 1H), 6.04 (dd, J = 8.5, 6.3 Hz, 1H), 4.10 (qd, J = 7.1, 1.5 Hz, 2H), 3.92 (d, J = 2.2 Hz, 3H), 3.76 (dd, J = 16.5, 8.5 Hz, 1H), 3.25 (dd, J = 16.5, 6.4 Hz, 1H), 2.72 (t, J = 7.4 Hz, 2H), 2.47 (t, J = 7.4 Hz, 2H), 2.13 (d, J = 1.9 Hz, 3H), 1.97 (p, J = 7.4 Hz, 2H), 1.17 (td, J = 7.2, 1.8 Hz, 3H).

Example 9 and Example 10 were prepared from Intermediate **E9C** according to the method described in Example 7. Example 9: 1H NMR (500 MHz, DMSO- d_6) δ 8.45 (s, 1H), 8.27 (s, 1H), 7.94 (s, 1H), 7.79 (dd, J = 8.6, 2.4 Hz, 1H), 7.56 (d, J = 8.5 Hz, 1H), 7.33 (s, 1H), 7.02 (d, J = 7.3 Hz, 1H), 6.87 (d, J = 8.6 Hz, 1H), 6.76 (d, J = 8.7 Hz, 1H), 6.25 (d, J = 7.3 Hz, 1H), 6.09 (s, 1H), 3.79 (s, 2H), 3.23 (d, J = 23.7 Hz, 3H), 2.88 (m, 1H), 2.72 (m, 1H), 2.62 (t, J = 7.4 Hz, 2H), 2.57 (d, J = 6.3 Hz, 1H), 2.43 (t, J = 7.7 Hz, 2H), 1.78 - 1.67 (m, 3H), 1.22 (m, 2H). LC/MS (m/z) = 472.0 ($M+H$) $^+$. Human $\alpha V\beta 6$ IC₅₀ (nM) = 2.6. Example 10: 1H NMR (500 MHz, DMSO- d_6) δ 8.45 (s, 1H), 8.25 (d, J = 2.5 Hz, 1H), 7.94 (s, 1H), 7.77 (dd, J = 8.6, 2.5 Hz, 1H), 7.58 (d, J = 8.5 Hz, 1H), 7.36 (s, 1H), 6.99 - 6.88 (m, 1H), 6.76 (d, J = 8.7 Hz, 1H), 6.09 (t, J = 7.8 Hz, 1H), 3.80 (s, 2H), 3.60 (s, 3H), 3.25 - 3.12 (m, 2H), 2.89 (s, 1H), 2.82 - 2.70 (m, 2H), 2.65 (dd, J = 10.2, 6.0 Hz, 1H), 2.55 (d, J = 5.8 Hz, 2H), 1.90 (s, 3H), 1.76 (s, 1H), 1.73 (d, J = 6.3 Hz, 2H). LC/MS (m/z) = 472.0 ($M+H$) $^+$. Human $\alpha V\beta 6$ IC₅₀ (nM) = 160.

Example 11

3-(6-Methoxypyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)pyrazolo[4,3-*b*]pyridin-1-yl)propanoic acid



Intermediate **E11A**: To a solution of 5-bromopyrazolo[4,3-*b*]pyridine (0.4395 g, 2.22 mmol) and 3,4-dihydro-2*H*-pyran (0.25 mL, 2.66 mmol) in CH₂Cl₂ (4.1 mL) was added 4-methylbenzenesulfonic acid (38 mg, 0.222 mmol). The mixture was stirred at rt overnight. The mixture was diluted with DCM (20 mL) and washed with water (3 x 8 mL). The organic layer was dried (Na₂SO₄), filtered, and concentrated under reduced pressure. The residue was purified *via* flash column chromatography to give **E11A** (527 mg, 84%). ¹H NMR (500 MHz, chloroform-*d*) δ 8.19 (d, *J* = 1.0 Hz, 1H), 7.89 (dd, *J* = 8.8, 0.9 Hz, 1H), 7.45 (d, *J* = 8.7 Hz, 1H), 5.75 (dd, *J* = 8.7, 2.6 Hz, 1H), 3.99 (dtd, *J* = 11.7, 4.0, 1.4 Hz, 1H), 3.81 - 3.71 (m, 1H), 2.56 - 2.42 (m, 1H), 2.20 - 2.10 (m, 2H), 1.87 - 1.67 (m, 3H).

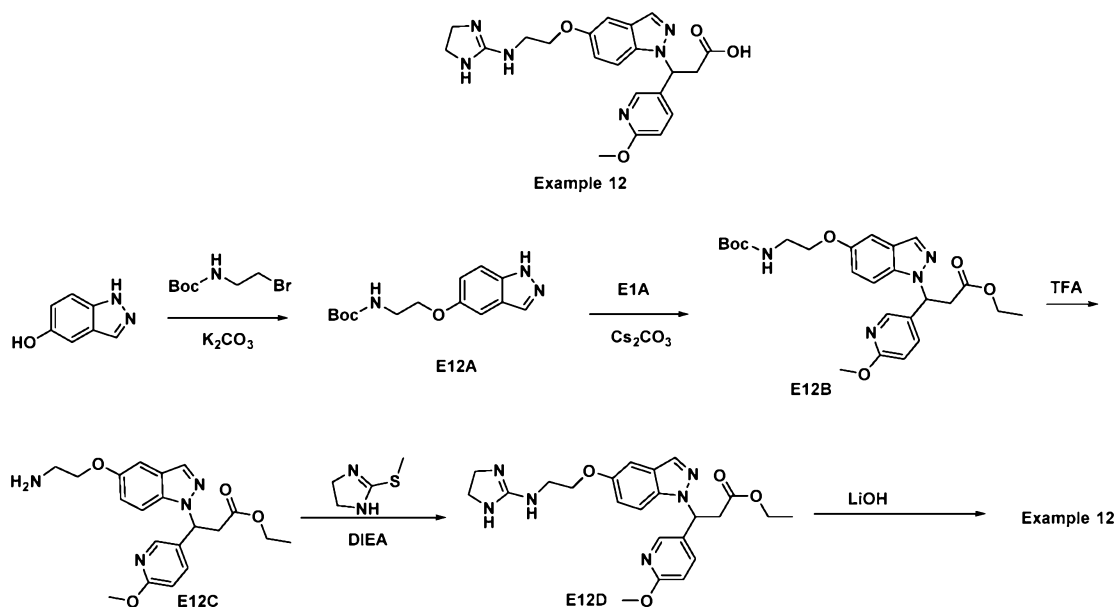
Intermediate **E11B**: A mixture of Intermediate **E11A** (284 mg, 1.021 mmol), 1,10-phenanthroline (30.7 mg, 0.170 mmol), copper(I) iodide (16.2 mg, 0.085 mmol), and Cs₂CO₃ (416 mg, 1.276 mmol) in toluene (1.1 mL) was degassed with argon for 10 min. The mixture was then sealed and heated at 120 °C overnight. The mixture was concentrated under reduced pressure, and the residue was purified *via* flash column chromatography (using 10% EtOAc in hexanes) to give **E11B** (90 mg, 22%). LCMS (ES): *m/z* 480.5 [M+H]⁺.

Intermediate **E11C**: To a solution of **E11B** (90 mg, 0.19 mmol) in MeOH (164 μ L) was added a solution of HCl in dioxane (4 M, 1 mL, 4 mmol). After stirred at rt for 48 hrs, the mixture was concentrated under reduced pressure, and the residue was purified *via* preparative HPLC (Phenomenex Luna Axia 5 μ C18 30 x 100 mm; 10 min gradient from 85% A: 15% B to 0% A:100% B (A = 90% H₂O/10% ACN + 0.1% TFA);
5 (B = 90% ACN/10% H₂O + 0.1% TFA); detection at 220 nm) to give **E11C** (10 mg, 18%). LCMS (ES): m/z 296.3 [M+H]⁺.

Example **11** was prepared from Intermediate **E11C** according to the method described in Example **3**. ¹H NMR (500 MHz, MeOH-*d*₄) δ 8.23 – 8.19 (m, 0H), 8.15 (d, *J* = 2.5 Hz, 1H), 8.04 (d, *J* = 9.1 Hz, 1H), 7.92 (s, 1H), 7.72 (dd, *J* = 8.7, 2.6 Hz, 1H),
10 7.44 (d, *J* = 7.4 Hz, 1H), 6.79 (d, *J* = 9.1 Hz, 1H), 6.73 (d, *J* = 8.8 Hz, 1H), 6.62 (d, *J* = 7.4 Hz, 1H), 6.17 (dd, *J* = 9.8, 5.1 Hz, 1H), 4.65 (ddt, *J* = 14.6, 11.5, 5.7 Hz, 2H), 3.85 (s, 3H), 3.67 (dd, *J* = 16.5, 9.8 Hz, 1H), 3.47 – 3.40 (m, 2H), 3.20 (dd, *J* = 16.6, 5.1 Hz, 1H), 3.13 (t, *J* = 6.1 Hz, 2H), 2.67 (d, *J* = 11.3 Hz, 2H), 1.85 (p, *J* = 6.0 Hz, 2H). LC/MS
15 (m/z) = 475.2 (M+H)⁺. Human α V β 6 IC₅₀ (nM) = 120.

Example 12

3-(5-(2-((4,5-Dihydroimidazol-2-yl)amino)ethoxy)-1H-indazol-1-yl)-3-(6-methoxypyridin-3-yl)propanoic acid



Intermediate **E12A**: To a solution of 1H-indazol-5-ol (1 g, 7.46 mmol) in DMF (5 mL) was added K₂CO₃ (2.06 g, 14.91 mmol) followed by *tert*-butyl (2-bromoethyl)carbamate (2.01 g, 8.95 mmol). The reaction mixture was stirred at rt for 1 day. The mixture was diluted with water (10 mL) and extracted with EtOAc (3 x 5 mL). The combined organic layers were washed with water (10 mL), followed by brine (10 mL). It was dried (Na₂SO₄), filtered, and concentrated under reduced pressure. The residue was purified *via* preparative HPLC (Phenomenex Luna Axia 5μ C18 30 x 100 mm; 10 min gradient from 85% A: 15% B to 0% A:100% B (A = 90% H₂O/10% ACN + 0.1% TFA); (B = 90% ACN/10% H₂O + 0.1% TFA); detection at 220 nm) to give **E12A** (526 mg, 1.897 mmol, 25.4% yield). LCMS (ES): m/z 278.2 [M+H]⁺.

Intermediate **E12B**: To a solution of **E12A** (75 mg, 0.270 mmol) in acetonitrile (1.5 mL) was added cesium carbonate (264 mg, 0.811 mmol) and stirred for 5 min at rt then Intermediate **E1A** (56.0 mg, 0.270 mmol) was added and stirred at 80 °C for 5 hrs. The reaction was cooled to room temperature, filtered and concentrated. The crude product was diluted with MeCN filtered and purified by preparative HPLC (Phenomenex Luna Axia 5μ C18 30 x 100 mm; 10 min gradient from 85% A: 15% B to 0% A:100% B (A = 90% H₂O/10% ACN + 0.1% TFA); (B = 90% ACN/10% H₂O + 0.1% TFA);

detection at 220 nm) to give **E12B** (53 mg, 0.089 mmol, 32.7% yield). LCMS (ES): m/z 485.1 [M+H]⁺.

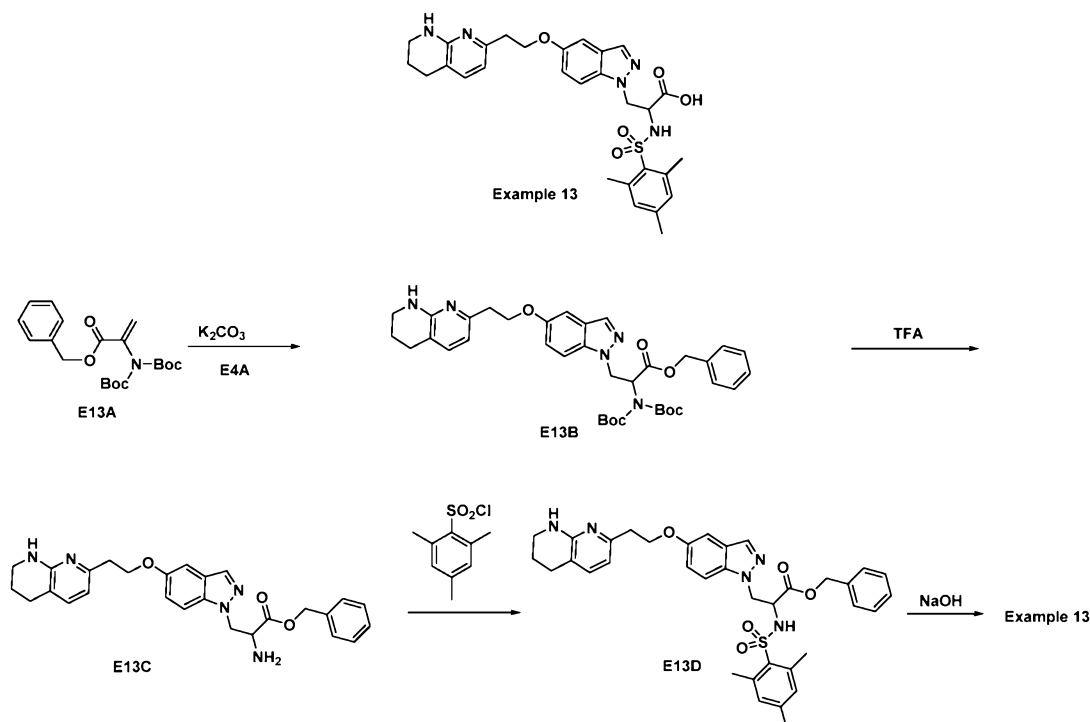
Intermediate **E12C**: To a solution of Intermediate **E12B** (53 mg, 0.089 mmol) in DCM (0.7 mL) was added TFA (0.05 mL, 0.649 mmol) and stirred at rt for 5 hrs. The reaction was concentrated. The crude product was diluted with MeCN filtered and purified by preparative HPLC (Phenomenex Luna Axia 5 μ C18 30 x 100 mm; 10 min gradient from 85% A: 15% B to 0% A:100% B (A = 90% H₂O/10% ACN + 0.1% TFA); (B = 90% ACN/10% H₂O + 0.1% TFA); detection at 220 nm) to give Intermediate **E12C** (57 mg, 0.093 mmol, 105% yield). LCMS (ES): m/z 385.1 [M+H]⁺.

Intermediate **E12D**: A solution of **E12C** (57 mg, 0.093 mmol), 2-(methylthio)-4,5-dihydroimidazole HCl salt (21.31 mg, 0.140 mmol) and DIPEA (0.081 mL, 0.465 mmol) in EtOH (2 mL) was heated to 150 °C in a microwave reactor for 15 min. The crude product was purified *via* preparative HPLC (Phenomenex Luna Axia 5 μ C18 30 x 100 mm; 10 min gradient from 85% A: 15% B to 0% A:100% B (A = 90% H₂O/10% ACN + 0.1% TFA); (B = 90% ACN/10% H₂O + 0.1% TFA); detection at 220 nm) to **E12D** (46 mg, 0.081 mmol, 87% yield). LCMS (ES): m/z 453.4 [M+H]⁺.

Example **12**: To a solution of Intermediate **E12D** (46 mg, 0.081 mmol) in THF (1 mL) was added a solution of LiOH (aqueous, 1 N, 0.244 mL, 0.244 mmol). The reaction mixture was stirred at rt overnight. The mixture was concentrated and purified by preparative LC/MS with the following conditions: Column: XBridge C18, 19 x 200 mm, 5- μ m particles; Mobile Phase A: 5:95 acetonitrile: water with 10-mM ammonium acetate; Mobile Phase B: 95:5 acetonitrile: water with 10-mM ammonium acetate; Gradient: 12-52% B over 25 minutes, then a 3-minute hold at 100% B; Flow: 20 mL/min to give Example **12** (2.5 mg, 7.4%). ¹H NMR (500MHz, chloroform-*d*) δ 7.94 (s, 1H), 7.66 (d, *J* = 9.1 Hz, 1H), 7.34 (d, *J* = 7.2 Hz, 1H), 7.12 - 7.03 (m, 2H), 6.99 (d, *J* = 9.1 Hz, 1H), 6.57 (d, *J* = 8.0 Hz, 1H), 6.50 (d, *J* = 7.2 Hz, 1H), 6.21 (t, *J* = 6.7 Hz, 1H), 4.28 (t, *J* = 5.6 Hz, 2H), 4.19 (dt, *J* = 8.5, 4.4 Hz, 2H), 3.69 - 3.52 (m, 3H), 3.51 - 3.39 (m, 3H), 3.16 (t, *J* = 5.6 Hz, 2H), 2.74 (t, *J* = 6.1 Hz, 2H), 1.97 - 1.85 (m, 2H). LC/MS (m/z) = 501.4 (M+H)⁺. Human α V β 6 IC₅₀ (nM) = 2,000.

Example 13

3-(5-(2-(5,6,7,8-Tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)-2-((2,4,6-trimethylphenyl)sulfonamido)propanoic acid



Intermediate **E13B**: To a solution of **E4A** (55.2 mg, 0.135 mmol) and **E13A** (51.0 mg, 0.135 mmol) in acetonitrile (1.35 mL) was added K_2CO_3 (112 mg, 0.811 mmol). The mixture was stirred at rt for overnight. The mixture was filtered, and rinsed with acetonitrile. The filtrate was concentrated and the residue was purified *via* flash column chromatography to afford **E13B** (55 mg, 60%). LCMS (ES): m/z 672.8 $[M+H]^+$.

Intermediate **E13C**: To a solution of **E13B** (54.9 mg, 0.082 mmol) in CH_2Cl_2 (545 μ l) was added TFA (82 μ L, 1.062 mmol). The mixture was stirred at rt overnight. The solvent was removed and the residue was used in the next reaction without further purification. LCMS (ES): m/z 472.5 $[M+H]^+$.

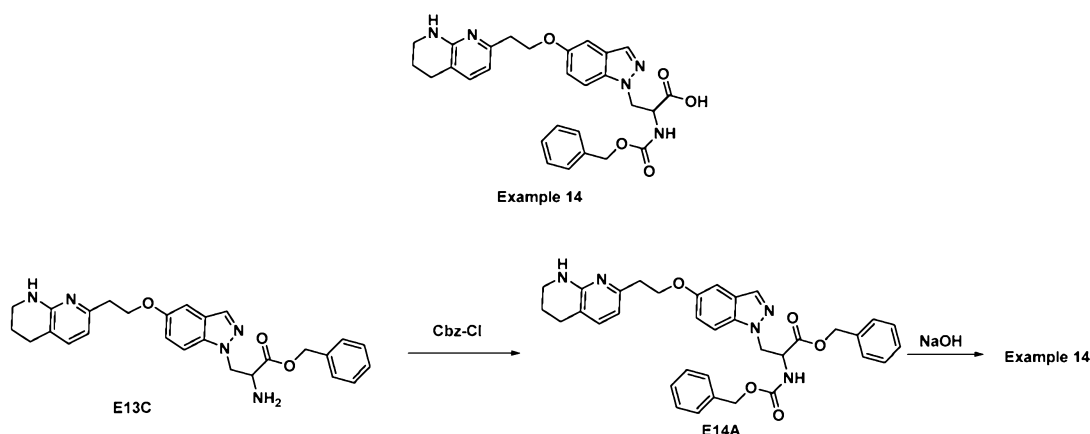
Intermediate **E13D**: To a solution of Intermediate **E13C** (19mg, 0.040 mmol) in THF (403 μ L) was added Et_3N (22.46 μ L, 0.161 mmol), followed by 2,4,6-trimethylbenzene-1-sulfonyl chloride (9.1 mg, 0.04 mmol). The mixture was stirred at rt overnight. The mixture was concentrated and purified *via* preparative HPLC (Sunfire 5 μ C18 30 x 100 mm; 10 min gradient from 95% A: 5% B to 0% A:100% B (A = 90%

H₂O/10% ACN + 0.1% TFA); (B = 90% ACN/10% H₂O + 0.1% TFA); detection at 220 nm) to **E13D** (4.4 mg, 17% yield). LCMS (ES): *m/z* 654.6 [M+H]⁺.

Example **13**: To a solution of Intermediate **E13D** (4.4 mg, 6.73 μmol) in MeOH (122 μL) was added NaOH (aqueous, 1 N, 20.2 μL, 0.020 mmol). The mixture was stirred at rt overnight. The mixture was neutralized with 1N HCl and concentrated. The crude was dissolved in 2 mL MeOH, filtered and purified by preparative LC/MS with the following conditions: Column: XBridge C18, 19 x 200 mm, 5-μm particles; Mobile Phase A: 5:95 acetonitrile: water with 10-mM ammonium acetate; Mobile Phase B: 95:5 acetonitrile: water with 10-mM ammonium acetate; Gradient: 10-50% B over 19 minutes, then a 5-minute hold at 100% B; Flow: 20 mL/min to give Example **13** (1.5 mg, 32%).
¹H NMR (500 MHz, MeOH-*d*₄) δ 7.63 (s, 1H), 7.57 (d, *J* = 7.3 Hz, 1H), 7.33 (d, *J* = 9.0 Hz, 1H), 6.99 (d, *J* = 2.2 Hz, 1H), 6.92 (d, *J* = 9.2 Hz, 1H), 6.72 (d, *J* = 7.4 Hz, 1H), 6.68 (s, 2H), 4.61 (dd, *J* = 14.3, 4.4 Hz, 1H), 4.47 (dd, *J* = 14.3, 8.1 Hz, 1H), 4.27 (t, *J* = 6.0 Hz, 2H), 4.15 (s, 1H), 3.49 (t, *J* = 5.6 Hz, 1H), 3.15 (t, *J* = 6.1 Hz, 2H), 2.81 (t, *J* = 6.3 Hz, 2H), 2.66 (s, 2H), 2.36 (s, 5H), 2.16 (s, 3H), 1.93 (t, *J* = 6.0 Hz, 2H). LC/MS (*m/z*) = 564.4 (M+H)⁺. Human αVβ6 IC₅₀ (nM) = 320.

Example 14

2-(((Benzyloxy)carbonyl)amino)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid



Intermediate **E14A**: To a solution of benzyl 2-amino-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoate (19 mg, 0.04 mmol) in THF

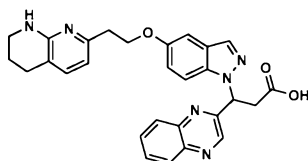
(403 μ L) was added sodium bicarbonate (aqueous, 1 N, 201 μ L, 0.201 mmol), followed by benzyl carbonochloridate (6.87 μ L, 0.048 mmol). The mixture was stirred at rt for 3 hrs. Solvent was removed under reduced pressure. The crude was dissolved in 2 mL MeOH, filtered and purified by preparative HPLC with the following conditions:

- 5 Column: Phenomenex Luna AXIA 5u C18 21.2 x 100 mm; Mobile Phase A: 10:90 MeOH: water with 0.1% TFA; Mobile Phase B: 90:10 MeOH: water with 0.1% TFA; Gradient: 20-100% B over 10 minutes, then a 2-minute hold at 100% B; Flow: 20 mL/min to give Intermediate **E14A** (17.6 mg, 72%). ^1H NMR (400 MHz, MeOH- d_4) δ 7.88 (s, 1H), 7.59 (dt, J = 7.3, 1.3 Hz, 1H), 7.36 (d, J = 9.1 Hz, 1H), 7.31 – 7.12 (m, 11H), 6.98 (dd, J = 9.1, 2.3 Hz, 1H), 6.74 (d, J = 7.3 Hz, 1H), 5.16 – 5.00 (m, 2H), 4.96 (s, 2H), 4.81 – 4.66 (m, 3H), 4.31 (t, J = 5.9 Hz, 2H), 3.48 (dd, J = 6.5, 4.8 Hz, 2H), 3.19 (t, J = 5.9 Hz, 2H), 2.80 (t, J = 6.3 Hz, 2H), 1.98 – 1.87 (m, 2H). LC/MS (m/z) = 606.7 (M+H) $^+$.
- 10

- Example 14: To a solution of **E14A** (benzyl 2-(((benzyloxy)carbonyl)amino)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoate) (17.6 mg, 0.029 mmol) in MeOH (528 μ L) was added NaOH (aqueous, 87 μ L, 1 N, 0.087 mmol). The mixture was stirred at rt overnight. The mixture was neutralized with HCl (aqueous, 1 N, 87 μ L) and concentrated under reduced pressure. The crude was purified *via* preparative LC/MS with the following conditions: Column: XBridge C18, 19 x 200 mm, 5- μ m particles; Mobile Phase A: 5:95 acetonitrile: water with 10-mM ammonium acetate; Mobile Phase B: 95:5 acetonitrile: water with 10-mM ammonium acetate; Gradient: 10-50% B over 20 minutes, then a 4-minute hold at 100% B; Flow: 20 mL/min to give Example 14 (9.3 mg, 62%). ^1H NMR (500 MHz, DMSO- d_6) δ 7.91 (s, 1H), 7.48 (s, 2H), 7.28 (d, J = 7.0 Hz, 4H), 7.17 (d, J = 6.7 Hz, 2H), 7.07 (d, J = 7.2 Hz, 1H), 6.95 (d, J = 9.0 Hz, 1H), 6.38 (d, J = 7.3 Hz, 1H), 6.30 (s, 1H), 4.92 (s, 2H), 4.68 (s, 3H), 4.23 (s, 2H), 3.24 (s, 2H), 3.17 (s, 1H), 2.90 (s, 2H), 2.61 (t, J = 6.3 Hz, 2H), 1.75 (t, J = 6.1 Hz, 2H). LC/MS (m/z) = 516.3 (M+H) $^+$. Human $\alpha\text{V}\beta 6$ IC₅₀ (nM) = 300.
- 15
- 20
- 25

Example 15

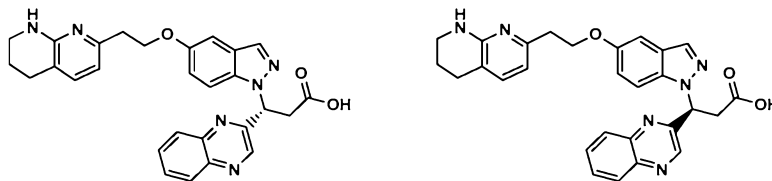
- 30 (±)-3-(Quinoxalin-2-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid



(±)-3-(Quinoxalin-2-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid was synthesized according to the procedure described in Example 3 using Intermediate E4A and ethyl (*E*)-3-(quinoxalin-2-yl)acrylate. ¹H NMR (500MHz, MeOH-*d*₄) δ 8.39 (s, 1H), 8.14 (d, *J* = 8.2 Hz, 1H), 8.02 (dd, *J* = 9.5, 7.9 Hz, 2H), 7.91 - 7.78 (m, 2H), 7.67 (d, *J* = 9.2 Hz, 1H), 7.57 (d, *J* = 7.5 Hz, 1H), 7.22 (d, *J* = 2.0 Hz, 1H), 7.10 (dd, *J* = 9.2, 2.1 Hz, 1H), 6.73 (d, *J* = 7.3 Hz, 1H), 6.62 (d, *J* = 5.5 Hz, 1H), 4.33 (t, *J* = 6.0 Hz, 2H), 3.85 (d, *J* = 17.5 Hz, 1H), 3.52 - 3.44 (m, 2H), 3.17 (t, *J* = 6.0 Hz, 2H), 2.79 (t, *J* = 6.3 Hz, 2H), 1.97 - 1.85 (m, 2H). LC/MS (*m/z*) = 495.1 (M+H)⁺. Human αVβ6 IC₅₀ (nM) = 6.0; Human αVβ1 IC₅₀ (nM) = 270; Human αVβ3 IC₅₀ (nM) = 2.7; Human αVβ5 IC₅₀ (nM) = 0.31; and Human αVβ8 IC₅₀ (nM) = 1,500.

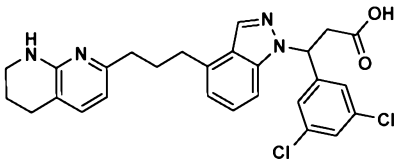
Example 16 and Example 17

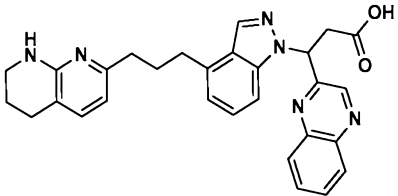
(*R*)-3-(Quinoxalin-2-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid and (*S*)-3-(quinoxalin-2-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid

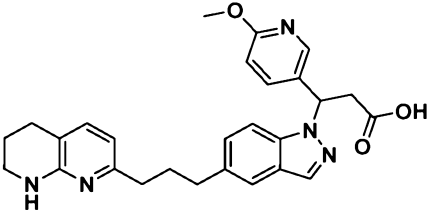


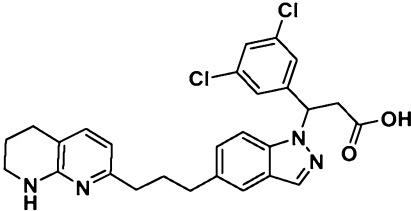
Example 15 (60 mg) was subjected to preparative chiral SFC purification (Column: Chiralpak IA, 21 x 250 mm, 5 micron, BPR Pressure: 120 bars, Temperature: 40 °C, Flow rate: 45 mL/min, Mobile Phase: CO₂/MeOH (60/40), Detector Wavelength: 220 nm) to afford Example 16 (15 mg) and Example 17 (19 mg) as a yellow solid. The enantiomeric excess for both Example 16 and Example 17 is ≥ 99.0%. Example 16: Human αVβ6 IC₅₀ (nM) = 136.76. Example 17: Human αVβ6 IC₅₀ (nM) = 4.2; Human αVβ1 IC₅₀ (nM) = 190; Human αVβ3 IC₅₀ (nM) = 2.1; Human αVβ5 IC₅₀ (nM) = 0.25; and Human αVβ8 IC₅₀ (nM) = 1,900.

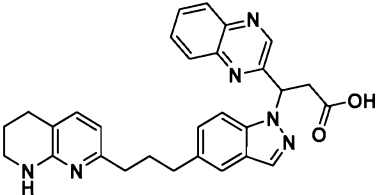
The following examples were prepared using methods analogous to the ones indicated in the table below.

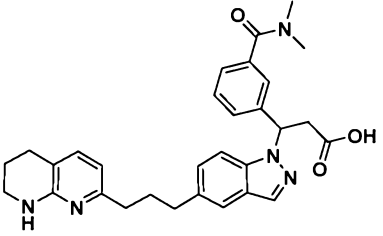
Example No.	Structure & Name	Analytical Data	Method
18	 <p>3-(3,5-Dichlorophenyl)-3-(4-(3-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)propyl)-1H-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, DMSO- <i>d</i> ₆) δ 8.22 (s, 1H), 7.60 (d, <i>J</i> = 8.5 Hz, 1H), 7.47 (d, <i>J</i> = 2.0 Hz, 1H), 7.40 (dd, <i>J</i> = 12.5, 4.6 Hz, 3H), 7.28 (t, <i>J</i> = 7.7 Hz, 1H), 6.94 (d, <i>J</i> = 7.0 Hz, 1H), 6.50 (d, <i>J</i> = 7.3 Hz, 1H), 6.22 (dd, <i>J</i> = 10.1, 4.9 Hz, 1H), 3.68 (s, 1H), 3.23 (dd, <i>J</i> = 16.8, 4.8 Hz, 1H), 2.96 – 2.82 (m, 2H), 2.64 (t, <i>J</i> = 7.2 Hz, 4H), 2.54 (s, 3H), 2.06 – 1.93 (m, 2H), 1.82 – 1.70 (m, 2H). LC/MS (m/z) = 509.1 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 2,700.	Example 1

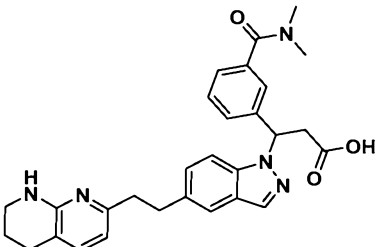
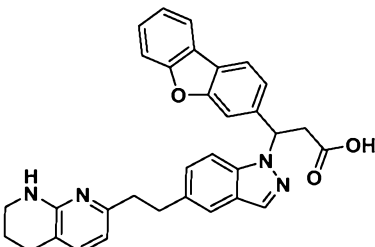
Example No.	Structure & Name	Analytical Data	Method
19	 <p>3-(Quinoxalin-2-yl)-3-(4-(3-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)propyl)-1H-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, DMSO- <i>d</i> ₆) δ 8.66 (s, 1H), 8.17 (s, 1H), 8.07 (dd, <i>J</i> = 16.5, 7.9 Hz, 2H), 7.86 (dt, <i>J</i> = 13.2, 6.9 Hz, 2H), 7.68 (d, <i>J</i> = 8.5 Hz, 1H), 7.32 (t, <i>J</i> = 7.7 Hz, 1H), 7.02 (d, <i>J</i> = 7.3 Hz, 1H), 6.96 (d, <i>J</i> = 7.1 Hz, 1H), 6.62 (t, <i>J</i> = 7.2 Hz, 1H), 3.21 (s, 2H), 2.88 (t, <i>J</i> = 7.6 Hz, 2H), 2.58 (t, <i>J</i> = 6.3 Hz, 2H), 2.55 (m, 2H), 2.47 (d, <i>J</i> = 7.6 Hz, 2H), 1.96 (t, <i>J</i> = 7.8 Hz, 2H), 1.90 (s, 1H), 1.78 – 1.66 (m, 2H). LC/MS (<i>m/z</i>) = 493.1 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 220.	Example 1

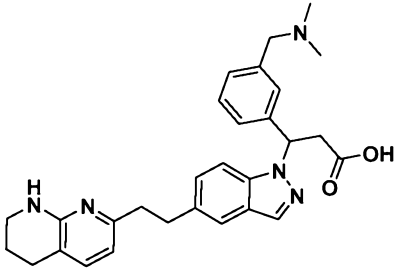
Example No.	Structure & Name	Analytical Data	Method
20	 <p data-bbox="427 801 863 981">3-(6-Methoxypyridin-3-yl)-3-(5-(3-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)propyl)-1H-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, DMSO- <i>d</i> ₆) δ 8.21 (s, 1H), 8.00 (s, 1H), 7.66 (d, <i>J</i> = 8.5 Hz, 2H), 7.48 (s, 1H), 7.21 (d, <i>J</i> = 8.5 Hz, 1H), 7.02 (d, <i>J</i> = 7.3 Hz, 1H), 6.71 (d, <i>J</i> = 8.9 Hz, 1H), 6.25 (d, <i>J</i> = 7.3 Hz, 1H), 6.17 - 6.08 (m, 1H), 3.75 (s, 3H), 3.60 (br. s., 2H), 3.21 (br. s., 2H), 2.63 (t, <i>J</i> = 7.2 Hz, 2H), 2.59 - 2.55 (m, 2H), 2.41 (t, <i>J</i> = 7.6 Hz, 2H), 1.87 - 1.80 (m, 2H), 1.74 - 1.65 (m, 2H). LC/MS (<i>m/z</i>) = 472.4 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 70.	Example 1

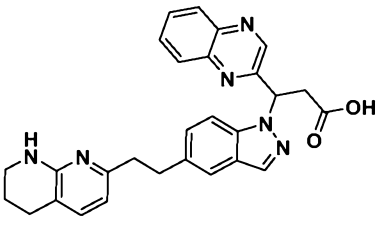
Example No.	Structure & Name	Analytical Data	Method
21	 <p>3-(3,5-Dichlorophenyl)-3-(5-(3-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)propyl)-1H-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, DMSO- <i>d</i> ₆) δ 8.07 (s, 1H), 7.68 (d, <i>J</i> = 8.5 Hz, 1H), 7.51 (s, 1H), 7.46 (s, 1H), 7.39 (s, 2H), 7.23 (d, <i>J</i> = 8.5 Hz, 1H), 7.02 (d, <i>J</i> = 7.3 Hz, 1H), 6.25 (d, <i>J</i> = 7.3 Hz, 1H), 6.23 - 6.17 (m, 1H), 3.59 (br. s., 2H), 3.21 (br. s., 2H), 2.64 (t, <i>J</i> = 7.0 Hz, 2H), 2.57 (t, <i>J</i> = 6.1 Hz, 2H), 2.42 (t, <i>J</i> = 7.6 Hz, 2H), 1.89 - 1.82 (m, 2H), 1.71 (d, <i>J</i> = 5.5 Hz, 2H). LC/MS (m/z) = 509.4 (M+H) ⁺ . Human αVβ6 IC ₅₀ (nM) = 44.	Example 1

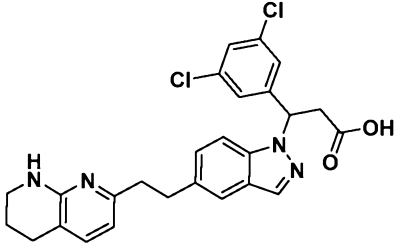
Example No.	Structure & Name	Analytical Data	Method
22	 <p data-bbox="432 835 855 1003">3-(Quinoxalin-2-yl)-3-(5-(3-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)propyl)-1H-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, DMSO- <i>d</i> ₆) δ 8.73 (s, 1H), 8.04 (t, <i>J</i> = 7.8 Hz, 2H), 8.00 (s, 1H), 7.83 (t, <i>J</i> = 7.2 Hz, 2H), 7.74 (d, <i>J</i> = 8.5 Hz, 1H), 7.24 (d, <i>J</i> = 8.2 Hz, 1H), 7.00 (d, <i>J</i> = 6.7 Hz, 1H), 6.56 (br. s., 1H), 6.29 - 6.15 (m, 2H), 3.21 (br. s., 2H), 2.72 - 2.62 (m, 2H), 2.58 (d, <i>J</i> = 5.2 Hz, 2H), 2.43 (t, <i>J</i> = 7.3 Hz, 2H), 1.93 - 1.82 (m, 4H), 1.73 (br. s., 2H). LC/MS (<i>m/z</i>) = 509.4 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 190.	Example 1

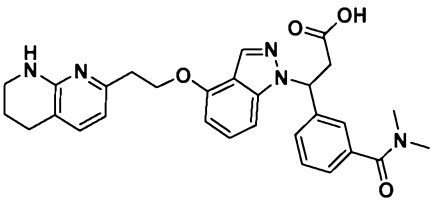
Example No.	Structure & Name	Analytical Data	Method
23	 <p>3-(3-(Dimethylcarbamoyl)phenyl)-3-(5-(3-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)propyl)-1H-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, DMSO- <i>d</i> ₆) δ 8.02 (br. s., 1H), 7.63 (d, <i>J</i> = 8.6 Hz, 1H), 7.55 - 7.46 (m, 1H), 7.40 (br. s., 1H), 7.37 - 7.29 (m, 2H), 7.24 (d, <i>J</i> = 7.4 Hz, 1H), 7.19 (d, <i>J</i> = 8.6 Hz, 1H), 7.01 (d, <i>J</i> = 7.1 Hz, 1H), 6.24 (d, <i>J</i> = 6.6 Hz, 2H), 3.21 (br. s., 2H), 2.93 (br. s., 3H), 2.78 (br. s., 3H), 2.64 (t, <i>J</i> = 7.1 Hz, 2H), 2.58 (t, <i>J</i> = 6.1 Hz, 2H), 2.42 (t, <i>J</i> = 7.4 Hz, 2H), 1.96 - 1.83 (m, 4H), 1.77 - 1.68 (m, 2H). LC/MS (<i>m/z</i>) = 512.2 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 61.	Example 1

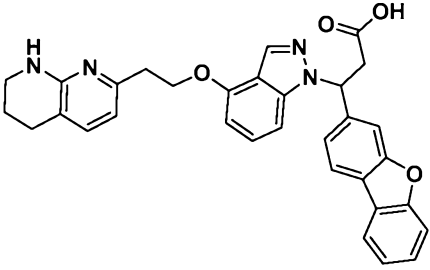
Example No.	Structure & Name	Analytical Data	Method
24	 <p>3-(3-(Dimethylcarbamoyl)phenyl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethyl)-1H-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, DMSO- <i>d</i> ₆) δ 8.05 (s, 1H), 7.67 (d, <i>J</i> = 8.9 Hz, 1H), 7.54 (s, 1H), 7.43 (d, <i>J</i> = 7.7 Hz, 1H), 7.39 – 7.33 (m, 2H), 7.25 (t, <i>J</i> = 8.2 Hz, 2H), 6.27 – 6.17 (m, 1H), 3.38 (m, 2H), 2.96 (m, 4H), 2.79 (m, 2H), 2.64 (m, 2H), 2.55 (s, 6H), 1.77 (s, 2H), 1.24 (s, 2H). LC/MS (<i>m/z</i>) = 458.2 (M+H) ⁺ . Human αVβ6 IC ₅₀ (nM) = 130.	Example 2
25	 <p>3-(Dibenzo[<i>b,d</i>]furan-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethyl)-1H-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, DMSO- <i>d</i> ₆) δ 8.10 - 7.99 (m, 3H), 7.77 - 7.62 (m, 3H), 7.54 - 7.44 (m, 2H), 7.41 - 7.33 (m, 2H), 7.22 (d, <i>J</i> = 8.6 Hz, 1H), 6.99 (d, <i>J</i> = 7.2 Hz, 1H), 6.36 (br. s., 1H), 6.26 (d, <i>J</i> = 7.2 Hz, 1H), 3.68 (br. s., 1H), 3.22 (br. s., 2H), 2.98 - 2.87 (m, 2H), 2.71 (t, <i>J</i> = 7.8 Hz, 2H), 2.58 (t, <i>J</i> = 5.8 Hz, 2H), 1.73 (br. s., 2H). LC/MS (<i>m/z</i>) = 517.2 (M+H) ⁺ . Human αVβ6 IC ₅₀ (nM) = 530.	Example 2

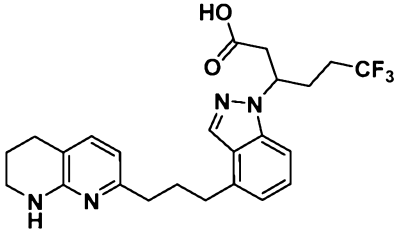
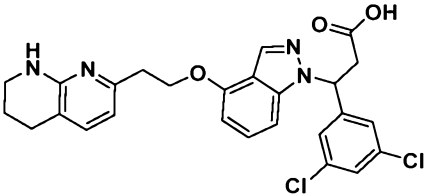
Example No.	Structure & Name	Analytical Data	Method
26	 <p data-bbox="411 857 874 1032">3-(3-((Dimethylamino)methyl)phenyl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethyl)-1H-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, DMSO- <i>d</i> ₆) δ 8.01 (s, 1H), 7.58 (d, <i>J</i> = 8.8 Hz, 1H), 7.51 (s, 1H), 7.28 (s, 1H), 7.21 (d, <i>J</i> = 4.0 Hz, 3H), 7.12 (br. s., 1H), 7.01 (d, <i>J</i> = 7.2 Hz, 1H), 6.27 (d, <i>J</i> = 7.2 Hz, 1H), 6.14 (dd, <i>J</i> = 9.5, 4.9 Hz, 1H), 3.30 (s, 1H), 3.23 (br. s., 2H), 3.18 - 3.11 (m, 1H), 2.98 - 2.90 (m, 2H), 2.78 - 2.68 (m, 2H), 2.59 (t, <i>J</i> = 5.9 Hz, 2H), 2.07 (s, 5H), 1.90 (s, 3H), 1.74 (d, <i>J</i> = 8.3 Hz, 2H). LC/MS (<i>m/z</i>) = 484.3 (<i>M</i> +H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 120.	Example 2

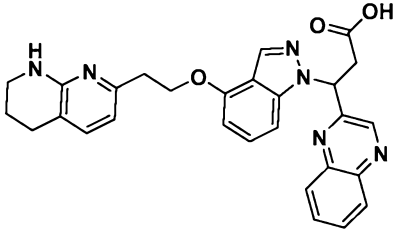
Example No.	Structure & Name	Analytical Data	Method
27	 <p data-bbox="422 806 869 929">3-(Quinoxalin-2-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethyl)-1H-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, DMSO- <i>d</i> ₆) δ 8.65 (s, 1H), 8.12 - 7.98 (m, 3H), 7.85 (t, <i>J</i> = 7.5 Hz, 2H), 7.75 (d, <i>J</i> = 8.7 Hz, 1H), 7.54 (s, 1H), 7.30 (d, <i>J</i> = 8.5 Hz, 1H), 7.01 (d, <i>J</i> = 7.3 Hz, 1H), 6.59 (br. s., 1H), 6.29 (d, <i>J</i> = 7.2 Hz, 1H), 3.50 (br. s., 2H), 3.23 (br. s., 2H), 2.95 (t, <i>J</i> = 8.0 Hz, 2H), 2.79 - 2.70 (m, 2H), 2.59 (t, <i>J</i> = 6.0 Hz, 2H), 1.73 (br. s., 2H). LC/MS (<i>m/z</i>) = 479.1 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 240.	Example 2

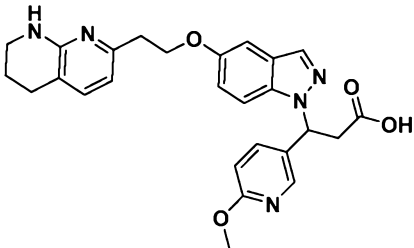
Example No.	Structure & Name	Analytical Data	Method
28	 <p data-bbox="416 869 866 996">3-(3,5-Dichlorophenyl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethyl)-1H-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, DMSO- <i>d</i> ₆) δ 8.10 - 8.02 (m, 1H), 7.71 (br d, <i>J</i> = 8.5 Hz, 1H), 7.56 - 7.50 (m, 1H), 7.50 - 7.44 (m, 1H), 7.42 (s, 2H), 7.26 (br d, <i>J</i> = 8.5 Hz, 1H), 7.07 - 6.99 (m, 1H), 6.29 (br d, <i>J</i> = 7.3 Hz, 1H), 6.24 (br dd, <i>J</i> = 9.0, 5.3 Hz, 1H), 3.22 - 3.14 (m, 2H), 3.04 (br d, <i>J</i> = 7.6 Hz, 1H), 2.99 - 2.92 (m, 2H), 2.86 (br d, <i>J</i> = 7.9 Hz, 1H), 2.81 - 2.69 (m, 2H), 2.60 (br t, <i>J</i> = 6.0 Hz, 2H), 1.80 - 1.71 (m, 1H). LC/MS (<i>m/z</i>) = 495.1 (M+H) ⁺ . Human αVβ6 IC ₅₀ (nM) = 210.	Example 2

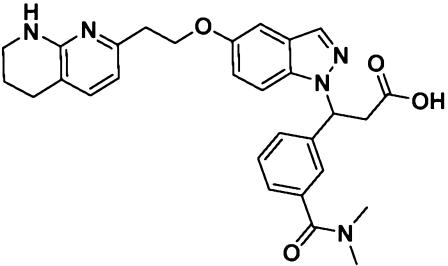
Example No.	Structure & Name	Analytical Data	Method
29	 <p>3-(3-(Dimethylcarbamoyl)phenyl)-3-(4-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, DMSO- <i>d</i> ₆) δ 8.06 (s, 1H), 7.62 (d, <i>J</i> = 7.2 Hz, 1H), 7.43 - 7.23 (m, 6H), 6.77 (d, <i>J</i> = 7.3 Hz, 1H), 6.57 (d, <i>J</i> = 7.5 Hz, 1H), 6.18 (dd, <i>J</i> = 9.8, 4.7 Hz, 1H), 4.37 (t, <i>J</i> = 5.7 Hz, 2H), 3.22 - 3.11 (m, 6H), 2.93 (br. s., 3H), 2.79 (br. s., 3H), 2.75 - 2.68 (m, 2H), 1.80 (br. s., 2H). LC/MS (<i>m/z</i>) = 514.0 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 1,500.	Example 2

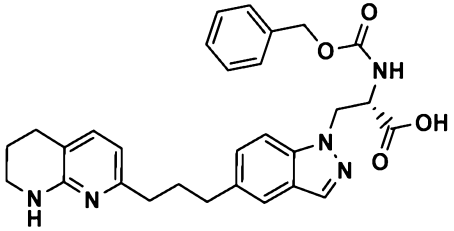
Example No.	Structure & Name	Analytical Data	Method
30	 <p data-bbox="427 943 858 1115">3-(Dibenzo[<i>b,d</i>]furan-3-yl)-3-(4-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1<i>H</i>-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, DMSO- <i>d</i> ₆) δ 8.10 - 7.98 (m, 3H), 7.70 (s, 1H), 7.65 (d, <i>J</i> = 8.2 Hz, 1H), 7.49 (t, <i>J</i> = 7.8 Hz, 1H), 7.41 - 7.33 (m, 2H), 7.30 (d, <i>J</i> = 8.2 Hz, 1H), 7.25 - 7.18 (m, 1H), 7.06 (d, <i>J</i> = 7.3 Hz, 1H), 6.56 (d, <i>J</i> = 7.6 Hz, 1H), 6.41 (d, <i>J</i> = 7.3 Hz, 1H), 6.32 (dd, <i>J</i> = 9.8, 4.6 Hz, 1H), 4.35 (t, <i>J</i> = 6.1 Hz, 2H), 3.69 (dd, <i>J</i> = 16.5, 10.1 Hz, 1H), 3.22 (br. s., 2H), 2.95 (t, <i>J</i> = 6.6 Hz, 2H), 2.59 (t, <i>J</i> = 6.0 Hz, 2H), 1.80 - 1.65 (m, 2H). LC/MS (<i>m/z</i>) = 532.9 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 1,200.	Example 3

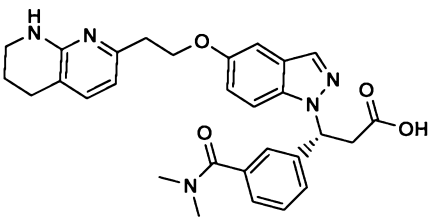
Example No.	Structure & Name	Analytical Data	Method
31	 <p>6,6,6-Trifluoro-3-(4-(3-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)propyl)-1H-indazol-1-yl)hexanoic acid</p>	¹ H NMR (500 MHz, DMSO- <i>d</i> ₆) δ 8.20 (s, 1H), 7.58 - 7.48 (m, 2H), 7.30 (t, <i>J</i> = 7.6 Hz, 1H), 6.94 (d, <i>J</i> = 7.0 Hz, 1H), 6.60 (d, <i>J</i> = 7.3 Hz, 1H), 5.13 (br. s., 1H), 3.02 - 2.86 (m, 5H), 2.78 - 2.66 (m, 4H), 2.30 - 1.96 (m, 6H), 1.80 (br. s., 2H). LC/MS (<i>m/z</i>) = 460.9 (<i>M</i> +H) ⁺ . Human αVβ6 IC ₅₀ (nM) = 3,300.	Example 1
32	 <p>3-(3,5-Dichlorophenyl)-3-(4-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, DMSO- <i>d</i> ₆) δ 8.10 (s, 1H), 7.57 (d, <i>J</i> = 7.3 Hz, 1H), 7.49 (s, 1H), 7.40 (d, <i>J</i> = 1.5 Hz, 2H), 7.36 (d, <i>J</i> = 8.5 Hz, 1H), 7.31 - 7.25 (m, 1H), 6.75 (d, <i>J</i> = 7.3 Hz, 1H), 6.60 (d, <i>J</i> = 7.6 Hz, 1H), 6.21 (dd, <i>J</i> = 10.1, 4.6 Hz, 1H), 4.38 (t, <i>J</i> = 6.1 Hz, 2H), 3.56 (dd, <i>J</i> = 16.8, 10.1 Hz, 1H), 3.17 (t, <i>J</i> = 5.6 Hz, 3H), 2.71 (t, <i>J</i> = 5.8 Hz, 2H), 1.80 (d, <i>J</i> = 5.5 Hz, 2H). LC/MS (<i>m/z</i>) = 511.2 (<i>M</i> +H) ⁺ . Human αVβ6 IC ₅₀ (nM) = 1,300.	Example 3

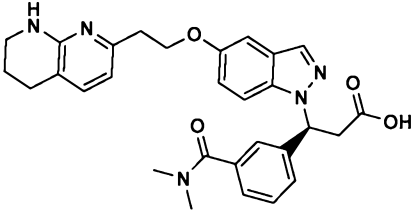
Example No.	Structure & Name	Analytical Data	Method
33	 <p data-bbox="432 824 855 996">3-(Quinoxalin-2-yl)-3-(4-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1<i>H</i>-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, DMSO- <i>d</i> ₆) δ 8.64 (s, 1H), 8.10 - 8.04 (m, 3H), 7.87 (quin, <i>J</i> = 6.8 Hz, 2H), 7.45 (br s, 1H), 7.43 (br d, <i>J</i> = 8.5 Hz, 1H), 7.33 (t, <i>J</i> = 8.0 Hz, 1H), 6.68 (br d, <i>J</i> = 7.0 Hz, 1H), 6.66 - 6.58 (m, 2H), 4.40 (br t, <i>J</i> = 6.0 Hz, 2H), 3.72 (dd, <i>J</i> = 16.8, 5.7 Hz, 1H), 3.61 - 3.52 (m, 1H), 3.13 (br s, 2H), 2.71 - 2.65 (m, 2H), 1.81 - 1.75 (m, 2H). LC/MS (<i>m/z</i>) = 495.3 (M+H) ⁺ . Human αVβ6 IC ₅₀ (nM) = 1,600.	Example 3

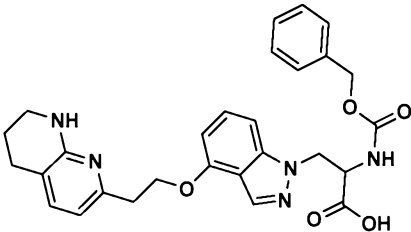
Example No.	Structure & Name	Analytical Data	Method
34	 <p>3-(6-Methoxypyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid, TFA</p>	¹ H NMR (500 MHz, chloroform- <i>d</i>) δ 8.26 (d, <i>J</i> = 2.2 Hz, 1H), 7.95 (s, 1H), 7.64 (dd, <i>J</i> = 8.8, 2.5 Hz, 1H), 7.35 - 7.28 (m, 2H), 7.06 (d, <i>J</i> = 1.9 Hz, 1H), 6.97 (dd, <i>J</i> = 9.1, 2.2 Hz, 1H), 6.73 (d, <i>J</i> = 8.8 Hz, 1H), 6.50 (d, <i>J</i> = 7.2 Hz, 1H), 6.06 (dd, <i>J</i> = 8.5, 5.8 Hz, 1H), 4.27 (t, <i>J</i> = 5.8 Hz, 2H), 3.92 (s, 3H), 3.73 (dd, <i>J</i> = 16.6, 8.7 Hz, 1H), 3.48 (t, <i>J</i> = 5.6 Hz, 2H), 3.29 (dd, <i>J</i> = 16.6, 5.6 Hz, 1H), 3.16 (t, <i>J</i> = 5.8 Hz, 2H), 2.74 (t, <i>J</i> = 6.1 Hz, 2H), 1.92 (quin, <i>J</i> = 5.9 Hz, 2H). LC/MS (<i>m/z</i>) = 474.1 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 12; Human α V β 1 IC ₅₀ (nM) = 95; Human α V β 3 IC ₅₀ (nM) = 2.5; Human α V β 5 IC ₅₀ (nM) = 0.38; and Human α V β 8 IC ₅₀ (nM) = 1,300.	Example 3

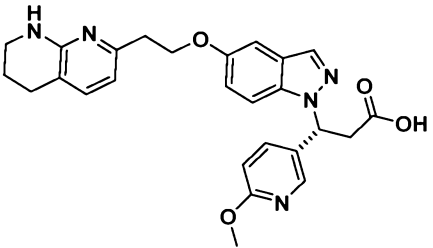
Example No.	Structure & Name	Analytical Data	Method
35	 <p data-bbox="419 965 868 1137">3-(3-(Dimethylcarbamoyl)phenyl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, MeOH- <i>d</i> ₄) δ 7.93 (s, 1H), 7.46 - 7.44 (m, 1H), 7.44 - 7.42 (m, 1H), 7.37 (t, <i>J</i> = 7.7 Hz, 1H), 7.33 (t, <i>J</i> = 1.7 Hz, 1H), 7.30 - 7.26 (m, 2H), 7.02 (d, <i>J</i> = 2.1 Hz, 1H), 6.92 (dd, <i>J</i> = 9.0, 2.3 Hz, 1H), 6.51 (d, <i>J</i> = 7.3 Hz, 1H), 6.24 (dd, <i>J</i> = 9.5, 5.5 Hz, 1H), 4.16 - 4.03 (m, 2H), 3.62 (dd, <i>J</i> = 15.9, 9.5 Hz, 1H), 3.41 - 3.36 (m, 2H), 3.19 (dd, <i>J</i> = 15.9, 5.4 Hz, 1H), 3.05 (s, 3H), 2.95 (t, <i>J</i> = 6.3 Hz, 2H), 2.86 (s, 3H), 2.70 (t, <i>J</i> = 6.2 Hz, 2H), 1.89 - 1.81 (m, 2H). LC/MS (<i>m/z</i>) = 514.0 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 17.	Example 3

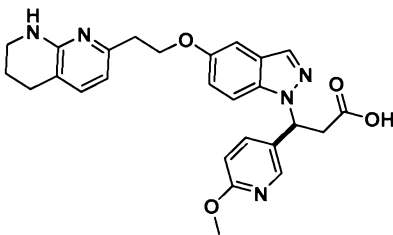
Example No.	Structure & Name	Analytical Data	Method
36	 <p data-bbox="416 757 871 936">(S)-2-(((Benzyloxy)carbonyl)amino)-3-(5-(3-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)propyl)-1H-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, DMSO- <i>d</i> ₆) δ 8.12 (s, 1H), 7.51 (s, 1H), 7.40 (s, 1H), 7.34 – 7.14 (m, 7H), 7.14 – 7.03 (m, 2H), 6.28 (d, <i>J</i> = 7.3 Hz, 1H), 4.95 (s, 2H), 3.62 (m, 4H), 3.23 (s, 3H), 2.60 (t, <i>J</i> = 6.5 Hz, 3H), 2.45 (t, <i>J</i> = 7.7 Hz, 1H), 1.89 (d, <i>J</i> = 14.9 Hz, 5H). LC/MS (<i>m/z</i>) = 514.2 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 77.	Example 4

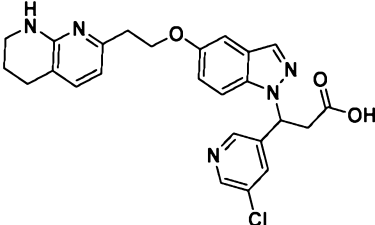
Example No.	Structure & Name	Analytical Data	Method
37	 <p>(R)-3-(3-(Dimethylcarbamoyl)phenyl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, MeOH- <i>d</i> ₄) δ 7.95 (d, <i>J</i> = 0.8 Hz, 1H), 7.50 – 7.43 (m, 2H), 7.39 (t, <i>J</i> = 7.7 Hz, 1H), 7.35 (t, <i>J</i> = 1.8 Hz, 1H), 7.33 – 7.27 (m, 2H), 7.04 (d, <i>J</i> = 2.2 Hz, 1H), 6.94 (dd, <i>J</i> = 9.1, 2.3 Hz, 1H), 6.53 (d, <i>J</i> = 7.3 Hz, 1H), 6.26 (dd, <i>J</i> = 9.5, 5.4 Hz, 1H), 4.11 (ddt, <i>J</i> = 25.1, 9.7, 6.4 Hz, 2H), 3.64 (dd, <i>J</i> = 16.0, 9.4 Hz, 1H), 3.43 – 3.38 (m, 2H), 3.21 (dd, <i>J</i> = 16.0, 5.5 Hz, 1H), 3.07 (s, 3H), 2.96 (t, <i>J</i> = 6.3 Hz, 2H), 2.88 (s, 3H), 2.72 (t, <i>J</i> = 6.3 Hz, 2H), 1.88 (dq, <i>J</i> = 6.9, 5.7 Hz, 2H). LC/MS (<i>m/z</i>) = 514.0 (<i>M</i> + <i>H</i>) ⁺ . Human αVβ6 IC ₅₀ (nM) = 1,100.	Example 16 and 17

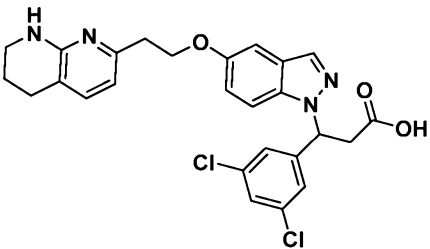
Example No.	Structure & Name	Analytical Data	Method
38	 <p data-bbox="411 958 874 1137">(S)-3-(3-(Dimethylcarbamoyl)phenyl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, MeOH- <i>d</i> ₄) δ 7.95 (d, <i>J</i> = 0.8 Hz, 1H), 7.50 – 7.43 (m, 2H), 7.39 (t, <i>J</i> = 7.7 Hz, 1H), 7.35 (t, <i>J</i> = 1.8 Hz, 1H), 7.33 – 7.27 (m, 2H), 7.04 (d, <i>J</i> = 2.2 Hz, 1H), 6.94 (dd, <i>J</i> = 9.1, 2.3 Hz, 1H), 6.53 (d, <i>J</i> = 7.3 Hz, 1H), 6.26 (dd, <i>J</i> = 9.5, 5.4 Hz, 1H), 4.11 (ddt, <i>J</i> = 25.1, 9.7, 6.4 Hz, 2H), 3.64 (dd, <i>J</i> = 16.0, 9.4 Hz, 1H), 3.43 – 3.38 (m, 2H), 3.21 (dd, <i>J</i> = 16.0, 5.5 Hz, 1H), 3.07 (s, 3H), 2.96 (t, <i>J</i> = 6.3 Hz, 2H), 2.88 (s, 3H), 2.72 (t, <i>J</i> = 6.3 Hz, 2H), 1.88 (dq, <i>J</i> = 6.9, 5.7 Hz, 2H). LC/MS (<i>m/z</i>) = 514.0 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 4.2.	Example 16 and 17

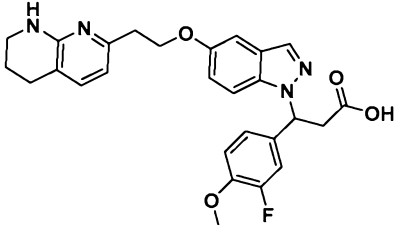
Example No.	Structure & Name	Analytical Data	Method
39	 <p>2-(((Benzyloxy)carbonyl)amino)-3-(4-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, MeOH- <i>d</i> ₄) δ 8.22 (s, 1H), 7.41 (d, <i>J</i> = 7.1 Hz, 1H), 7.33 – 7.05 (m, 6H), 6.63 (d, <i>J</i> = 7.3 Hz, 1H), 6.41 – 6.33 (m, 1H), 5.12 – 4.94 (m, 2H), 4.85 (m, 2H), 4.60 (s, 1H), 4.36 (q, <i>J</i> = 6.0 Hz, 2H), 3.44 – 3.36 (m, 2H), 3.10 (t, <i>J</i> = 6.2 Hz, 2H), 2.69 (d, <i>J</i> = 18.0 Hz, 3H), 1.86 (d, <i>J</i> = 6.4 Hz, 2H). LC/MS (<i>m/z</i>) = 516.3 (M+H) ⁺ . Human αVβ6 IC ₅₀ (nM) = 340.	Example 4

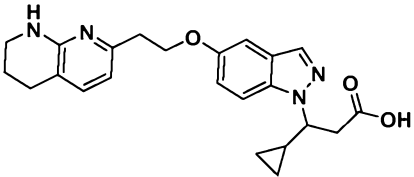
Example No.	Structure & Name	Analytical Data	Method
40	 <p data-bbox="427 958 861 1131">(R)-3-(6-Methoxypyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, MeOH- <i>d</i> ₄) δ 8.12 (d, <i>J</i> = 2.1 Hz, 1H), 7.91 (s, 1H), 7.68 (dd, <i>J</i> = 8.7, 2.4 Hz, 1H), 7.50 (d, <i>J</i> = 9.0 Hz, 1H), 7.28 (d, <i>J</i> = 7.3 Hz, 1H), 7.00 (d, <i>J</i> = 1.8 Hz, 1H), 6.94 (dd, <i>J</i> = 9.0, 2.3 Hz, 1H), 6.69 (d, <i>J</i> = 8.7 Hz, 1H), 6.51 (d, <i>J</i> = 7.3 Hz, 1H), 6.18 (dd, <i>J</i> = 9.1, 5.7 Hz, 1H), 4.19 - 4.01 (m, 2H), 3.84 (s, 3H), 3.56 (dd, <i>J</i> = 15.7, 9.0 Hz, 1H), 3.41 - 3.34 (m, 2H), 3.15 (dd, <i>J</i> = 15.7, 5.6 Hz, 1H), 2.94 (t, <i>J</i> = 6.3 Hz, 2H), 2.68 (t, <i>J</i> = 6.3 Hz, 2H), 1.89 - 1.79 (m, 2H). LC/MS (<i>m/z</i>) = 474.2 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 210.	Example 16 and 17

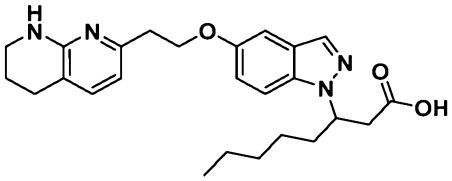
Example No.	Structure & Name	Analytical Data	Method
41	 <p>(S)-3-(6-Methoxypyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, MeOH- <i>d</i> ₄) δ 8.11 (d, <i>J</i> = 2.4 Hz, 1H), 7.91 (s, 1H), 7.68 (dd, <i>J</i> = 8.7, 2.4 Hz, 1H), 7.50 (d, <i>J</i> = 9.2 Hz, 1H), 7.27 (d, <i>J</i> = 7.3 Hz, 1H), 7.01 (d, <i>J</i> = 2.0 Hz, 1H), 6.95 (dd, <i>J</i> = 9.2, 2.3 Hz, 1H), 6.69 (d, <i>J</i> = 8.7 Hz, 1H), 6.51 (d, <i>J</i> = 7.3 Hz, 1H), 6.18 (dd, <i>J</i> = 9.2, 5.8 Hz, 1H), 4.19 - 4.04 (m, 2H), 3.84 (s, 3H), 3.55 (dd, <i>J</i> = 15.8, 9.2 Hz, 1H), 3.41 - 3.34 (m, 2H), 3.15 (dd, <i>J</i> = 15.9, 5.8 Hz, 1H), 2.94 (t, <i>J</i> = 6.3 Hz, 2H), 2.69 (t, <i>J</i> = 6.3 Hz, 2H), 1.85 (dt, <i>J</i> = 11.7, 6.0 Hz, 2H). LC/MS (<i>m/z</i>) = 474.3 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 6.2; Human α V β 1 IC ₅₀ (nM) = 83; Human α V β 3 IC ₅₀ (nM) = 2.3; Human α V β 5 IC ₅₀ (nM) = 0.22; and Human α V β 8 IC ₅₀ (nM) = 510.	Example 16 and 17

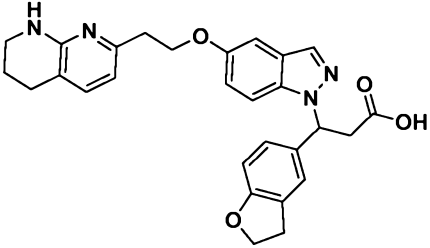
Example No.	Structure & Name	Analytical Data	Method
42	 <p>3-(5-Chloropyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid, TFA</p>	¹ H NMR (500 MHz, MeOH- <i>d</i> ₄) δ 8.46 (d, <i>J</i> = 7.5 Hz, 1H), 8.00 (s, 1H), 7.86 (s, 1H), 7.62 - 7.55 (m, 2H), 7.19 (d, <i>J</i> = 2.1 Hz, 1H), 7.06 (dd, <i>J</i> = 9.2, 2.3 Hz, 1H), 6.74 (d, <i>J</i> = 7.3 Hz, 1H), 6.29 (dd, <i>J</i> = 9.5, 5.5 Hz, 1H), 4.30 (t, <i>J</i> = 6.0 Hz, 2H), 3.69 (dd, <i>J</i> = 16.7, 9.7 Hz, 1H), 3.52 - 3.45 (m, 2H), 3.16 (t, <i>J</i> = 5.9 Hz, 2H), 2.80 (t, <i>J</i> = 6.2 Hz, 2H), 1.97 - 1.87 (m, 2H). LC/MS (m/z) = 477.9 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 4.4.	Example 3

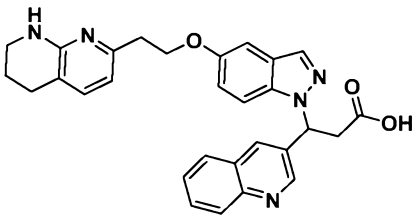
Example No.	Structure & Name	Analytical Data	Method
43	 <p data-bbox="416 931 866 1104">3-(3,5-Dichlorophenyl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1<i>H</i>-indazol-1-yl)propanoic acid, TFA</p>	¹ H NMR (500 MHz, DMSO- <i>d</i> ₆) δ 8.02 (s, 1H), 7.71 (d, <i>J</i> = 9.2 Hz, 1H), 7.49 (t, <i>J</i> = 1.9 Hz, 1H), 7.41 (d, <i>J</i> = 2.0 Hz, 2H), 7.18 (d, <i>J</i> = 2.1 Hz, 1H), 7.05 (d, <i>J</i> = 7.3 Hz, 1H), 7.00 (dd, <i>J</i> = 9.1, 2.4 Hz, 1H), 6.37 (d, <i>J</i> = 7.2 Hz, 1H), 6.31 (br. s., 1H), 6.23 (dd, <i>J</i> = 10.0, 5.0 Hz, 1H), 4.24 (td, <i>J</i> = 6.9, 1.3 Hz, 2H), 3.56 (dd, <i>J</i> = 16.7, 10.0 Hz, 1H), 3.25 - 3.18 (m, 3H), 2.89 (t, <i>J</i> = 6.8 Hz, 2H), 2.61 (t, <i>J</i> = 6.2 Hz, 2H), 1.80 - 1.68 (m, 2H). LC/MS (<i>m/z</i>) = 511.1 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 3.2.	Example 5

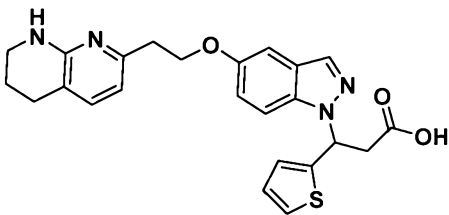
Example No.	Structure & Name	Analytical Data	Method
44	 <p>3-(3-Fluoro-4-methoxyphenyl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid, TFA</p>	¹ H NMR (500 MHz, DMSO- <i>d</i> ₆) δ 7.97 (s, 1H), 7.64 (d, <i>J</i> = 9.2 Hz, 1H), 7.27 - 7.15 (m, 3H), 7.11 - 7.02 (m, 2H), 6.97 (dd, <i>J</i> = 9.1, 2.4 Hz, 1H), 6.48 (d, <i>J</i> = 6.3 Hz, 1H), 6.11 (dd, <i>J</i> = 10.0, 5.0 Hz, 1H), 4.24 (td, <i>J</i> = 6.6, 1.8 Hz, 2H), 3.80 - 3.71 (m, 3H), 3.56 (dd, <i>J</i> = 16.6, 10.0 Hz, 1H), 3.15 (dd, <i>J</i> = 16.6, 5.0 Hz, 1H), 2.96 (t, <i>J</i> = 6.1 Hz, 2H), 2.64 (t, <i>J</i> = 6.0 Hz, 2H), 1.82 - 1.71 (m, 2H). LC/MS (<i>m/z</i>) = 490.9 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 8.71.	Example 3

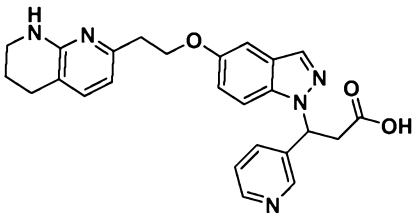
Example No.	Structure & Name	Analytical Data	Method
45	 <p>3-Cyclopropyl-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid, TFA</p>	¹ H NMR (500 MHz, MeOH- <i>d</i> ₄) δ 7.92 (s, 1H), 7.56 (d, <i>J</i> = 7.3 Hz, 1H), 7.49 (d, <i>J</i> = 9.2 Hz, 1H), 7.13 (d, <i>J</i> = 2.1 Hz, 1H), 7.02 (dd, <i>J</i> = 9.1, 2.4 Hz, 1H), 6.72 (d, <i>J</i> = 7.3 Hz, 1H), 4.34 (td, <i>J</i> = 9.6, 4.6 Hz, 1H), 4.28 (td, <i>J</i> = 6.0, 1.4 Hz, 2H), 3.51 - 3.45 (m, 2H), 3.26 (dd, <i>J</i> = 16.2, 9.8 Hz, 1H), 3.14 (t, <i>J</i> = 6.0 Hz, 2H), 3.06 (dd, <i>J</i> = 16.0, 4.6 Hz, 1H), 2.80 (t, <i>J</i> = 6.2 Hz, 2H), 1.97 - 1.88 (m, 2H), 1.50 - 1.39 (m, 1H), 0.72 - 0.60 (m, 1H), 0.47 - 0.34 (m, 2H), 0.28 (dq, <i>J</i> = 9.6, 4.7 Hz, 1H). LC/MS (<i>m/z</i>) = 406.9 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 2,300.	Example 3

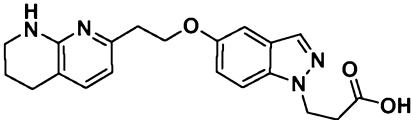
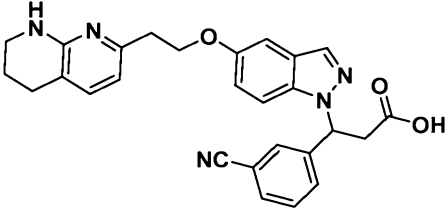
Example No.	Structure & Name	Analytical Data	Method
46	 <p data-bbox="416 920 869 1048">3-(5-(2-(5,6,7,8-Tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)octanoic acid, TFA</p>	¹ H NMR (500 MHz, MeOH- <i>d</i> ₄) δ 7.94 (s, 1H), 7.60 - 7.49 (m, 2H), 7.15 (d, <i>J</i> = 2.1 Hz, 1H), 7.04 (dd, <i>J</i> = 9.2, 2.4 Hz, 1H), 6.72 (d, <i>J</i> = 7.3 Hz, 1H), 4.29 (t, <i>J</i> = 6.0 Hz, 2H), 3.50 - 3.47 (m, 2H), 3.15 (t, <i>J</i> = 6.0 Hz, 2H), 3.03 (dd, <i>J</i> = 16.2, 9.2 Hz, 1H), 2.88 (dd, <i>J</i> = 16.2, 4.9 Hz, 1H), 2.81 (t, <i>J</i> = 6.2 Hz, 2H), 2.09 - 1.99 (m, 1H), 1.94 (dt, <i>J</i> = 11.7, 6.0 Hz, 2H), 1.90 - 1.80 (m, 1H), 1.27 - 1.04 (m, 6H), 0.88 - 0.80 (m, 1H), 0.79 - 0.73 (m, 3H). LC/MS (<i>m/z</i>) = 437.2 (<i>M</i> +H) ⁺ . Human α V β 6 IC (nM) = 330.	Example 3

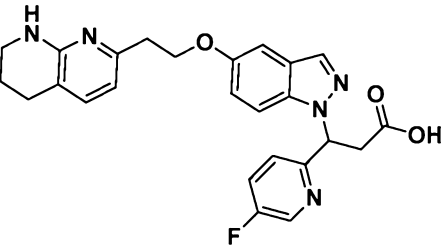
Example No.	Structure & Name	Analytical Data	Method
47	 <p data-bbox="416 860 866 1032">3-(2,3-Dihydrobenzofuran-5-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1<i>H</i>-indazol-1-yl)propanoic acid, TFA</p>	¹ H NMR (500 MHz, MeOH- <i>d</i> ₄) δ 7.90 (s, 1H), 7.48 - 7.41 (m, 2H), 7.17 - 7.09 (m, 2H), 7.05 (dd, <i>J</i> = 8.3, 1.8 Hz, 1H), 6.97 (dd, <i>J</i> = 9.2, 2.3 Hz, 1H), 6.67 - 6.57 (m, 2H), 6.08 (dd, <i>J</i> = 9.6, 5.2 Hz, 1H), 4.47 (td, <i>J</i> = 8.7, 1.0 Hz, 2H), 4.23 (t, <i>J</i> = 6.3 Hz, 2H), 3.62 (dd, <i>J</i> = 16.4, 9.7 Hz, 1H), 3.49 - 3.39 (m, 2H), 3.20 - 3.04 (m, 5H), 2.75 (t, <i>J</i> = 6.2 Hz, 2H), 1.89 (dt, <i>J</i> = 11.7, 6.1 Hz, 2H). LC/MS (<i>m/z</i>) = 485.0 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 7.0.	Example 3

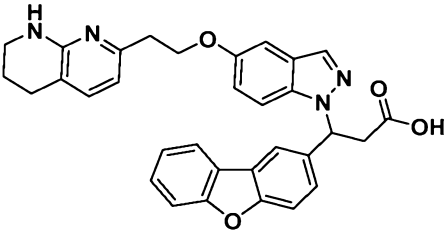
Example No.	Structure & Name	Analytical Data	Method
48	 <p>3-(Quinolin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid, TFA</p>	¹ H NMR (500 MHz, MeOH- <i>d</i> ₄) δ 8.81 (d, <i>J</i> = 2.0 Hz, 1H), 8.34 (d, <i>J</i> = 2.0 Hz, 1H), 8.01 (s, 1H), 7.99 - 7.87 (m, 2H), 7.81 - 7.73 (m, 1H), 7.66 - 7.53 (m, 3H), 7.19 (d, <i>J</i> = 2.1 Hz, 1H), 7.05 (dd, <i>J</i> = 9.2, 2.3 Hz, 1H), 6.73 (d, <i>J</i> = 7.2 Hz, 1H), 6.47 (dd, <i>J</i> = 9.2, 5.4 Hz, 1H), 4.30 (t, <i>J</i> = 6.0 Hz, 2H), 3.79 (s, 1H), 3.52 - 3.41 (m, 3H), 3.21 - 3.11 (m, 2H), 2.77 (t, <i>J</i> = 6.0 Hz, 2H), 1.90 (dt, <i>J</i> = 11.7, 6.1 Hz, 2H). LC/MS (<i>m/z</i>) = 494.0 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 5.5; Human α V β 1 IC ₅₀ (nM) = 270; Human α V β 3 IC ₅₀ (nM) = 3.8; Human α V β 5 IC ₅₀ (nM) = 0.54; and Human α V β 8 IC ₅₀ (nM) = 4,400.	Example 3

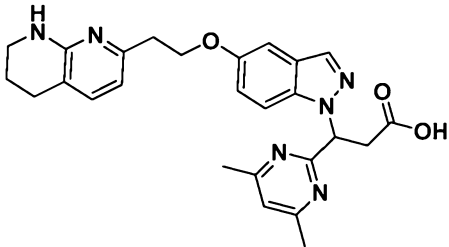
Example No.	Structure & Name	Analytical Data	Method
49	 <p>3-(5-(2-(5,6,7,8-Tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)-3-(thiophen-2-yl)propanoic acid, TFA</p>	¹ H NMR (500 MHz, DMSO- <i>d</i> ₆) δ 7.96 (s, 1H), 7.69 (d, <i>J</i> = 9.3 Hz, 1H), 7.37 (dd, <i>J</i> = 5.0, 1.2 Hz, 1H), 7.16 (dd, <i>J</i> = 6.1, 2.9 Hz, 2H), 7.05 (d, <i>J</i> = 7.3 Hz, 1H), 7.00 (dd, <i>J</i> = 9.1, 2.4 Hz, 1H), 6.91 (dd, <i>J</i> = 5.0, 3.5 Hz, 1H), 6.46 (dd, <i>J</i> = 9.7, 5.1 Hz, 1H), 6.37 (d, <i>J</i> = 7.2 Hz, 1H), 6.31 (br. s., 1H), 4.24 (t, <i>J</i> = 6.9 Hz, 2H), 3.59 (dd, <i>J</i> = 16.3, 9.8 Hz, 1H), 3.26 - 3.20 (m, 3H), 2.90 (t, <i>J</i> = 6.8 Hz, 2H), 2.61 (t, <i>J</i> = 6.2 Hz, 2H), 1.75 (quin, <i>J</i> = 6.0 Hz, 2H). LC/MS (<i>m/z</i>) = 449.0 (<i>M</i> +H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 33.	Example 3

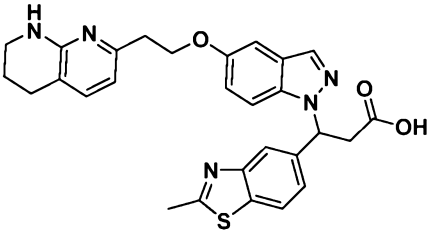
Example No.	Structure & Name	Analytical Data	Method
50	 <p>3-(Pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid, TFA</p>	¹ H NMR (500 MHz, MeOH- <i>d</i> ₄) δ 8.57 (s, 1H), 8.47 (d, <i>J</i> = 4.7 Hz, 1H), 7.98 (s, 1H), 7.94 (d, <i>J</i> = 7.7 Hz, 1H), 7.65 - 7.53 (m, 2H), 7.46 (dd, <i>J</i> = 7.7, 5.2 Hz, 1H), 7.19 (d, <i>J</i> = 2.2 Hz, 1H), 7.05 (dd, <i>J</i> = 9.1, 2.2 Hz, 1H), 6.74 (d, <i>J</i> = 7.4 Hz, 1H), 6.30 (dd, <i>J</i> = 9.1, 5.5 Hz, 1H), 4.31 (t, <i>J</i> = 5.9 Hz, 2H), 3.71 (dd, <i>J</i> = 16.8, 9.4 Hz, 1H), 3.54 - 3.46 (m, 2H), 3.38 - 3.33 (m, 1H), 3.17 (t, <i>J</i> = 5.8 Hz, 2H), 2.80 (t, <i>J</i> = 6.2 Hz, 2H), 1.93 (quin, <i>J</i> = 5.9 Hz, 2H). LC/MS (<i>m/z</i>) = 444.1 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 80.	Example 3

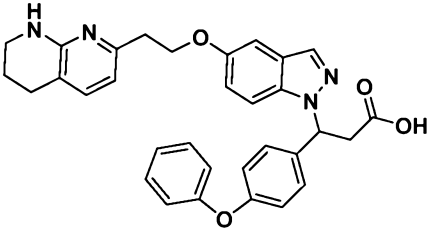
Example No.	Structure & Name	Analytical Data	Method
51	 <p>3-(5-(2-(5,6,7,8-Tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1<i>H</i>-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, DMSO- <i>d</i> ₆) δ 7.90 (s, 1H), 7.55 (d, <i>J</i> = 9.0 Hz, 1H), 7.16 (s, 1H), 7.06 (d, <i>J</i> = 7.0 Hz, 1H), 6.98 (d, <i>J</i> = 9.3 Hz, 1H), 6.38 (d, <i>J</i> = 7.3 Hz, 1H), 6.27 (br. s., 1H), 4.52 (t, <i>J</i> = 6.4 Hz, 2H), 4.23 (t, <i>J</i> = 6.6 Hz, 2H), 3.23 (br. s., 2H), 2.89 (t, <i>J</i> = 6.5 Hz, 2H), 2.80 (t, <i>J</i> = 6.4 Hz, 2H), 2.66 - 2.57 (m, 2H), 1.74 (br. s., 2H). LC/MS (m/z) = 367.1 (M+H) ⁺ . Human αVβ6 IC ₅₀ (nM) = 2,500.	Example 3
52	 <p>3-(3-Cyanophenyl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1<i>H</i>-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, MeOH- <i>d</i> ₄) δ 7.70 - 7.55 (m, 4H), 7.52 - 7.42 (m, 2H), 7.17 (s, 1H), 7.07 (d, <i>J</i> = 7.9 Hz, 1H), 6.73 (d, <i>J</i> = 7.3 Hz, 1H), 6.20 (br. s., 1H), 4.31 (t, <i>J</i> = 5.9 Hz, 2H), 3.54 - 3.43 (m, 2H), 3.16 (t, <i>J</i> = 5.7 Hz, 2H), 2.80 (t, <i>J</i> = 6.2 Hz, 2H), 1.93 (dt, <i>J</i> = 11.7, 6.1 Hz, 2H). LC/MS (m/z) = 468.0 (M+H) ⁺ .	Example 3

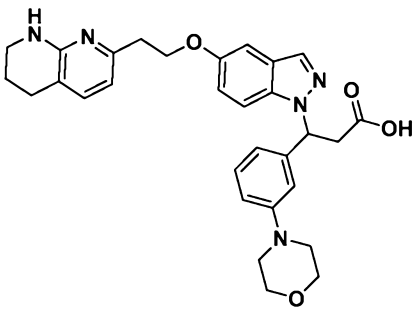
Example No.	Structure & Name	Analytical Data	Method
53	 <p>3-(5-Fluoropyridin-2-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid, TFA</p>	¹ H NMR (500 MHz, MeOH- <i>d</i> ₄) δ 8.38 (d, <i>J</i> = 2.7 Hz, 1H), 7.91 (s, 1H), 7.54 - 7.37 (m, 2H), 7.16 (d, <i>J</i> = 7.3 Hz, 1H), 7.13 (d, <i>J</i> = 2.1 Hz, 1H), 7.05 (dd, <i>J</i> = 8.9, 4.3 Hz, 1H), 6.99 (d, <i>J</i> = 10.8 Hz, 1H), 6.48 (d, <i>J</i> = 7.3 Hz, 1H), 6.33 (t, <i>J</i> = 7.2 Hz, 1H), 4.24 (t, <i>J</i> = 6.7 Hz, 2H), 3.43 - 3.35 (m, 4H), 2.98 (t, <i>J</i> = 6.7 Hz, 2H), 2.70 (t, <i>J</i> = 6.2 Hz, 2H), 1.90 - 1.83 (m, 2H). LC/MS (m/z) = 462.0 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 150.	Example 3

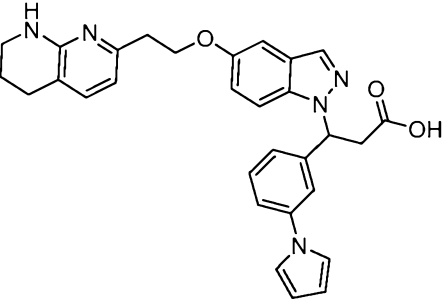
Example No.	Structure & Name	Analytical Data	Method
54	 <p>3-(Dibenzo[<i>b,d</i>]furan-2-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1<i>H</i>-indazol-1-yl)propanoic acid, TFA</p>	¹ H NMR (500 MHz, DMSO- <i>d</i> ₆) δ 8.09 - 8.05 (m, 1H), 8.04 - 8.00 (m, 2H), 7.73 - 7.63 (m, 3H), 7.49 (ddd, <i>J</i> = 8.4, 7.2, 1.4 Hz, 1H), 7.42 - 7.34 (m, 2H), 7.17 (d, <i>J</i> = 2.3 Hz, 1H), 7.03 (d, <i>J</i> = 7.3 Hz, 1H), 6.96 (dd, <i>J</i> = 9.1, 2.4 Hz, 1H), 6.38 - 6.32 (m, 2H), 6.29 (s, 1H), 4.28 - 4.18 (m, 2H), 3.70 (dd, <i>J</i> = 16.6, 10.0 Hz, 1H), 3.24 - 3.18 (m, 2H), 2.88 (t, <i>J</i> = 6.9 Hz, 2H), 2.59 (t, <i>J</i> = 6.3 Hz, 2H), 1.78 - 1.68 (m, 2H). LC/MS (<i>m/z</i>) = 533.0 (<i>M</i> + <i>H</i>) ⁺ . Human αVβ6 IC ₅₀ (nM) = 30.	Example 3

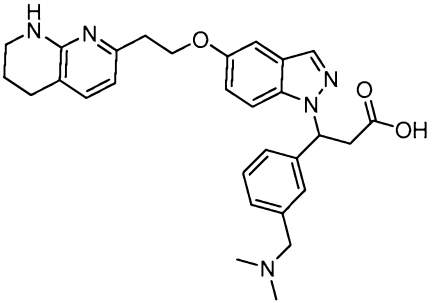
Example No.	Structure & Name	Analytical Data	Method
55	 <p>3-(4,6-Dimethylpyrimidin-2-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid, TFA</p>	¹ H NMR (500 MHz, DMSO- <i>d</i> ₆) δ 7.87 (d, <i>J</i> = 0.6 Hz, 1H), 7.57 (d, <i>J</i> = 9.2 Hz, 1H), 7.18 - 7.11 (m, 2H), 7.06 (d, <i>J</i> = 7.3 Hz, 1H), 6.99 (dd, <i>J</i> = 9.1, 2.4 Hz, 1H), 6.38 (d, <i>J</i> = 7.3 Hz, 1H), 6.30 (s, 1H), 6.14 (dd, <i>J</i> = 8.0, 6.8 Hz, 1H), 4.24 (t, <i>J</i> = 6.9 Hz, 2H), 3.57 (dd, <i>J</i> = 16.7, 6.5 Hz, 1H), 3.38 - 3.34 (m, 1H), 3.26 - 3.19 (m, 2H), 2.90 (t, <i>J</i> = 6.8 Hz, 2H), 2.61 (t, <i>J</i> = 6.3 Hz, 2H), 2.33 (s, 6H), 1.81 - 1.69 (m, 2H). LC/MS (<i>m/z</i>) = 473.0 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 920.	Example 3

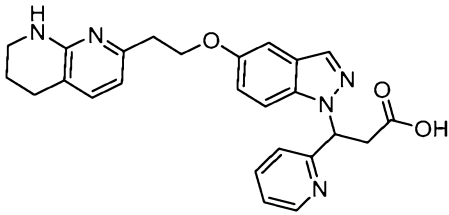
Example No.	Structure & Name	Analytical Data	Method
56	 <p>3-(2-Methylbenzo[d]thiazol-5-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid, TFA</p>	¹ H NMR (500 MHz, DMSO- <i>d</i> ₆) δ 7.96 (s, 1H), 7.86 (d, <i>J</i> = 8.3 Hz, 1H), 7.78 (s, 1H), 7.58 (d, <i>J</i> = 9.2 Hz, 1H), 7.32 (d, <i>J</i> = 8.4 Hz, 1H), 7.14 (s, 1H), 7.04 (d, <i>J</i> = 7.2 Hz, 1H), 6.91 (d, <i>J</i> = 9.2 Hz, 1H), 6.35 (d, <i>J</i> = 7.2 Hz, 1H), 6.28 (t, <i>J</i> = 7.2 Hz, 1H), 6.18 (br. s., 1H), 4.19 (br. s., 2H), 3.21 (br. s., 2H), 3.06 (dd, <i>J</i> = 15.9, 5.8 Hz, 1H), 2.86 (t, <i>J</i> = 6.6 Hz, 2H), 2.72 (s, 3H), 2.58 (t, <i>J</i> = 5.9 Hz, 2H), 1.72 (br. s., 2H). LC/MS (m/z) = 514.0 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 4.1.	Example 3

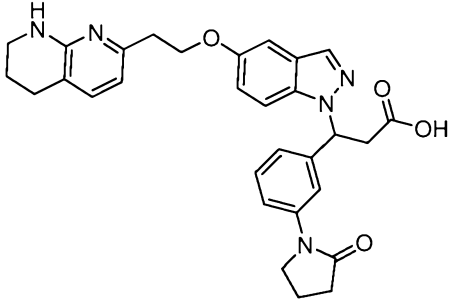
Example No.	Structure & Name	Analytical Data	Method
57	 <p>3-(4-Phenoxyphenyl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1<i>H</i>-indazol-1-yl)propanoic acid, TFA</p>	¹ H NMR (500 MHz, MeOH- <i>d</i> ₄) δ 7.90 (s, 1H), 7.59 - 7.51 (m, 1H), 7.42 (d, <i>J</i> = 9.0 Hz, 1H), 7.33 - 7.22 (m, 3H), 7.15 (s, 1H), 7.11 - 6.98 (m, 3H), 6.89 - 6.79 (m, 4H), 6.74 - 6.66 (m, 1H), 6.14 (dd, <i>J</i> = 9.6, 5.2 Hz, 1H), 4.29 (t, <i>J</i> = 6.0 Hz, 2H), 3.53 - 3.42 (m, 2H), 3.14 (d, <i>J</i> = 5.5 Hz, 2H), 2.78 (t, <i>J</i> = 6.2 Hz, 2H), 1.98 - 1.84 (m, 2H). LC/MS (<i>m/z</i>) = 535.0 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 11.	Example 3

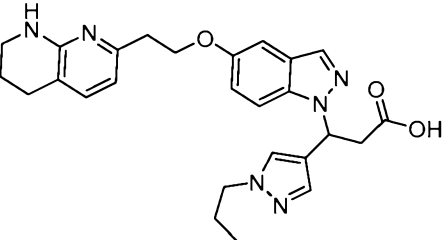
Example No.	Structure & Name	Analytical Data	Method
58	 <p>3-(3-Morpholinophenyl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid, TFA</p>	¹ H NMR (500 MHz, chloroform- <i>d</i>) δ 8.01 (s, 1H), 7.33 (d, <i>J</i> = 7.4 Hz, 1H), 7.22 (dd, <i>J</i> = 18.3, 8.7 Hz, 2H), 7.12 (d, <i>J</i> = 2.2 Hz, 1H), 6.99 (dd, <i>J</i> = 9.1, 2.2 Hz, 1H), 6.83 (dd, <i>J</i> = 8.3, 2.2 Hz, 1H), 6.80 (s, 1H), 6.72 (d, <i>J</i> = 7.4 Hz, 1H), 6.51 (d, <i>J</i> = 7.2 Hz, 1H), 6.00 (dd, <i>J</i> = 9.2, 4.3 Hz, 1H), 4.32 (t, <i>J</i> = 5.9 Hz, 2H), 3.85 (t, <i>J</i> = 4.8 Hz, 4H), 3.79 (dd, <i>J</i> = 16.4, 9.2 Hz, 1H), 3.50 (t, <i>J</i> = 4.8 Hz, 2H), 3.31 (dd, <i>J</i> = 16.4, 4.3 Hz, 1H), 3.24 - 3.17 (m, 2H), 3.11 (q, <i>J</i> = 4.5 Hz, 4H), 2.76 (t, <i>J</i> = 6.1 Hz, 2H), 1.98 - 1.88 (m, 2H). LC/MS (<i>m/z</i>) = 528.2 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 1.7.	Example 3

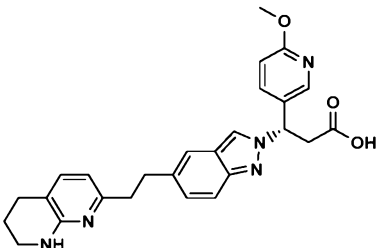
Example No.	Structure & Name	Analytical Data	Method
59	 <p>3-(3-(1<i>H</i>-Pyrrol-1-yl)phenyl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1<i>H</i>-indazol-1-yl)propanoic acid, TFA</p>	¹ H NMR (500 MHz, MeOH- <i>d</i> ₄) δ 7.92 (s, 1H), 7.50 (d, <i>J</i> = 9.3 Hz, 1H), 7.42 (s, 1H), 7.35 - 7.27 (m, 2H), 7.16 (d, <i>J</i> = 7.0 Hz, 2H), 7.13 (d, <i>J</i> = 2.4 Hz, 1H), 7.09 (t, <i>J</i> = 2.2 Hz, 2H), 6.99 (dd, <i>J</i> = 9.2, 2.3 Hz, 1H), 6.48 (d, <i>J</i> = 7.2 Hz, 1H), 6.28 - 6.19 (m, 3H), 4.24 (t, <i>J</i> = 6.7 Hz, 2H), 3.54 - 3.44 (m, 1H), 3.41 - 3.35 (m, 1H), 2.98 (t, <i>J</i> = 6.8 Hz, 2H), 2.69 (t, <i>J</i> = 6.3 Hz, 2H), 1.89 - 1.80 (m, 2H). LC/MS (<i>m/z</i>) = 508.0 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 1.2.	Example 3

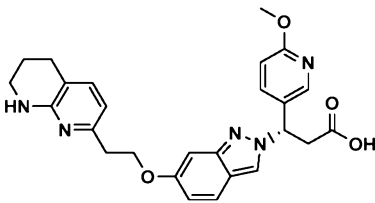
Example No.	Structure & Name	Analytical Data	Method
60	 <p>3-(3-((Dimethylamino)methyl)phenyl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid, TFA</p>	¹ H NMR (500 MHz, MeOH- <i>d</i> ₄) δ 7.90 (d, <i>J</i> = 0.5 Hz, 1H), 7.52 - 7.44 (m, 2H), 7.39 - 7.30 (m, 2H), 7.29 - 7.24 (m, 1H), 7.18 (d, <i>J</i> = 7.3 Hz, 1H), 7.12 (d, <i>J</i> = 2.1 Hz, 1H), 6.97 (dd, <i>J</i> = 9.1, 2.4 Hz, 1H), 6.49 (d, <i>J</i> = 7.3 Hz, 1H), 6.22 (t, <i>J</i> = 7.6 Hz, 1H), 4.22 (t, <i>J</i> = 6.7 Hz, 2H), 4.05 (s, 2H), 3.39 - 3.35 (m, 3H), 2.97 (t, <i>J</i> = 6.7 Hz, 2H), 2.70 (t, <i>J</i> = 6.3 Hz, 2H), 2.65 (s, 6H), 1.90 - 1.81 (m, 2H). LC/MS (<i>m/z</i>) = 500.1 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 5.1.	Example 3

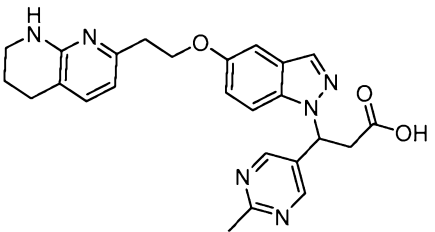
Example No.	Structure & Name	Analytical Data	Method
61	 <p>3-(Pyridin-2-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid, TFA</p>	¹ H NMR (500 MHz, DMSO- <i>d</i> ₆) δ 8.51 (d, <i>J</i> = 4.0 Hz, 1H), 7.98 (s, 1H), 7.66 (t, <i>J</i> = 7.0 Hz, 1H), 7.53 (d, <i>J</i> = 8.9 Hz, 1H), 7.29 - 7.23 (m, 1H), 7.18 (s, 1H), 7.07 (d, <i>J</i> = 7.3 Hz, 1H), 6.97 (d, <i>J</i> = 7.3 Hz, 1H), 6.89 (d, <i>J</i> = 7.9 Hz, 1H), 6.38 (d, <i>J</i> = 7.3 Hz, 1H), 6.23 - 6.14 (m, 1H), 4.22 (t, <i>J</i> = 6.6 Hz, 2H), 3.56 (br. s., 1H), 3.50 - 3.40 (m, 1H), 3.23 (br. s., 2H), 2.89 (t, <i>J</i> = 6.6 Hz, 2H), 2.60 (t, <i>J</i> = 6.0 Hz, 2H), 1.81 - 1.66 (m, 2H). LC/MS (<i>m/z</i>) = 444.0 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 80.	Example 3

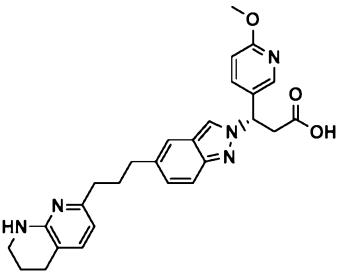
Example No.	Structure & Name	Analytical Data	Method
62	 <p>3-(3-(2-Oxopyrrolidin-1-yl)phenyl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid, TFA</p>	¹ H NMR (500 MHz, MeOH- <i>d</i> ₄) δ 7.94 (d, <i>J</i> = 0.6 Hz, 1H), 7.63 - 7.56 (m, 2H), 7.51 - 7.43 (m, 2H), 7.29 (t, <i>J</i> = 7.9 Hz, 1H), 7.16 (d, <i>J</i> = 2.0 Hz, 1H), 7.08 (d, <i>J</i> = 7.6 Hz, 1H), 7.00 (dd, <i>J</i> = 9.2, 2.3 Hz, 1H), 6.74 (d, <i>J</i> = 7.3 Hz, 1H), 6.16 (dd, <i>J</i> = 9.9, 5.0 Hz, 1H), 4.31 (t, <i>J</i> = 6.0 Hz, 2H), 3.88 - 3.81 (m, 2H), 3.72 (dd, <i>J</i> = 16.6, 9.9 Hz, 1H), 3.52 - 3.44 (m, 2H), 3.25 (dd, <i>J</i> = 16.6, 5.0 Hz, 1H), 3.16 (t, <i>J</i> = 5.9 Hz, 2H), 2.79 (t, <i>J</i> = 6.2 Hz, 2H), 2.61 - 2.52 (m, 2H), 2.20 - 2.09 (m, 2H), 1.98 - 1.86 (m, 2H). LC/MS (m/z) = 526.2 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 37.	Example 3

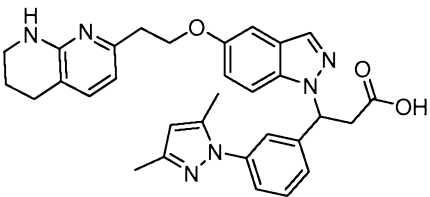
Example No.	Structure & Name	Analytical Data	Method
63	 <p>3-(1-Propylpyrazol-4-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid, TFA</p>	¹ H NMR (500 MHz, MeOH- <i>d</i> ₄) δ 7.89 (s, 1H), 7.56 - 7.51 (m, 2H), 7.34 (s, 1H), 7.24 (d, <i>J</i> = 7.0 Hz, 1H), 7.08 (d, <i>J</i> = 2.1 Hz, 1H), 6.99 (dd, <i>J</i> = 9.1, 2.4 Hz, 1H), 6.51 (d, <i>J</i> = 7.3 Hz, 1H), 6.21 (dd, <i>J</i> = 8.4, 6.4 Hz, 1H), 4.19 (q, <i>J</i> = 6.3 Hz, 2H), 3.99 (t, <i>J</i> = 7.0 Hz, 2H), 3.41 - 3.36 (m, 2H), 3.21 - 3.12 (m, 1H), 2.97 (t, <i>J</i> = 6.6 Hz, 2H), 2.71 (t, <i>J</i> = 6.3 Hz, 2H), 1.91 - 1.83 (m, 2H), 1.82 - 1.72 (m, 2H), 0.83 (t, <i>J</i> = 7.4 Hz, 3H). LC/MS (<i>m/z</i>) = 475.3 (<i>M</i> + <i>H</i>) ⁺ . Human α V β 6 IC ₅₀ (nM) = 55.	Example 3

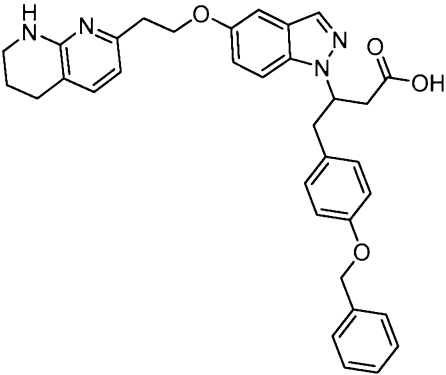
Example No.	Structure & Name	Analytical Data	Method
64	 <p>(S)-3-(6-Methoxypyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethyl)-2H-indazol-2-yl)propanoic acid</p>	¹ H NMR (500 MHz, MeOH- <i>d</i> ₄) δ 8.24 (d, <i>J</i> = 0.9 Hz, 1H), 8.16 (d, <i>J</i> = 2.5 Hz, 1H), 7.71 (dd, <i>J</i> = 8.7, 2.6 Hz, 1H), 7.46 (dt, <i>J</i> = 9.0, 0.9 Hz, 1H), 7.26 – 7.20 (m, 2H), 7.06 (dd, <i>J</i> = 8.9, 1.6 Hz, 1H), 6.73 (dd, <i>J</i> = 8.7, 0.7 Hz, 1H), 6.32 (d, <i>J</i> = 7.3 Hz, 1H), 6.14 (dd, <i>J</i> = 9.3, 5.9 Hz, 1H), 3.86 (s, 3H), 3.48 (dd, <i>J</i> = 15.8, 9.3 Hz, 1H), 3.39 (dd, <i>J</i> = 6.5, 4.7 Hz, 2H), 3.17 (dd, <i>J</i> = 15.8, 6.0 Hz, 1H), 2.91 – 2.78 (m, 4H), 2.69 (t, <i>J</i> = 6.3 Hz, 2H), 1.91 – 1.83 (m, 2H). LC/MS (<i>m/z</i>) = 458.3 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 19.	Example 7

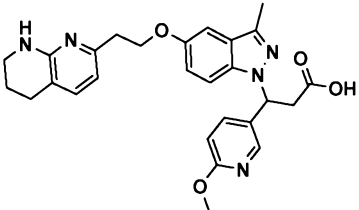
Example No.	Structure & Name	Analytical Data	Method
65	 <p>(S)-3-(6-Methoxypyridin-3-yl)-3-(6-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-2H-indazol-2-yl)propanoic acid</p>	¹ H NMR (500 MHz, MeOH- <i>d</i> ₄) δ 8.25 (s, 1H), 8.15 (d, <i>J</i> = 2.5 Hz, 1H), 7.71 (dd, <i>J</i> = 8.7, 2.6 Hz, 1H), 7.45 (d, <i>J</i> = 9.0 Hz, 1H), 7.34 (d, <i>J</i> = 7.3 Hz, 1H), 6.82 (d, <i>J</i> = 2.1 Hz, 1H), 6.72 (d, <i>J</i> = 8.7 Hz, 1H), 6.63 (dd, <i>J</i> = 9.1, 2.1 Hz, 1H), 6.55 (d, <i>J</i> = 7.3 Hz, 1H), 6.08 (dd, <i>J</i> = 9.2, 6.0 Hz, 1H), 4.16 (t, <i>J</i> = 6.4 Hz, 2H), 3.85 (s, 3H), 3.49 (dd, <i>J</i> = 16.0, 9.3 Hz, 1H), 3.40 (t, <i>J</i> = 5.7 Hz, 2H), 3.18 (dd, <i>J</i> = 15.9, 5.9 Hz, 1H), 3.05 - 2.96 (m, 2H), 2.72 (t, <i>J</i> = 6.3 Hz, 2H), 1.87 (p, <i>J</i> = 6.1 Hz, 2H). LC/MS (<i>m/z</i>) = 474.0 (M+H) ⁺ . Human αVβ6 IC ₅₀ (nM) = 1.2; Human αVβ1 IC ₅₀ (nM) = 2,300; Human αVβ3 IC ₅₀ (nM) = 2.4; Human αVβ5 IC ₅₀ (nM) = 1.0; and Human αVβ8 IC ₅₀ (nM) = 2,300.	Example 6

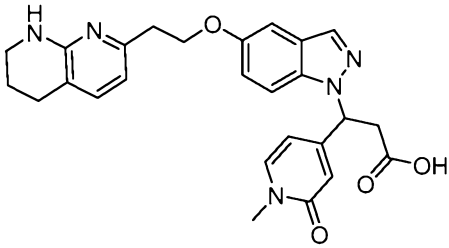
Example No.	Structure & Name	Analytical Data	Method
66	 <p data-bbox="427 824 863 1003">3-(2-Methylpyrimidin-5-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid, TFA</p>	¹ H NMR (500 MHz, chloroform- <i>d</i>) δ 9.63 (br. s., 1H), 8.83 (s, 2H), 7.99 (s, 1H), 7.39 - 7.31 (m, 2H), 7.08 (s, 1H), 7.01 (d, <i>J</i> = 9.1 Hz, 1H), 6.51 (d, <i>J</i> = 7.4 Hz, 1H), 6.13 (t, <i>J</i> = 7.2 Hz, 1H), 4.29 (t, <i>J</i> = 5.6 Hz, 2H), 3.73 (dd, <i>J</i> = 16.8, 8.0 Hz, 1H), 3.48 (t, <i>J</i> = 5.4 Hz, 2H), 3.38 (dd, <i>J</i> = 16.9, 6.5 Hz, 1H), 3.17 (t, <i>J</i> = 5.8 Hz, 2H), 2.81 - 2.70 (m, 5H), 1.92 (quin, <i>J</i> = 5.9 Hz, 2H). LC/MS (<i>m/z</i>) = 459.1 (<i>M</i> +H) ⁺ .	Example 3

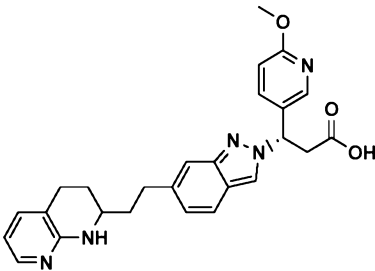
Example No.	Structure & Name	Analytical Data	Method
67	 <p data-bbox="427 965 863 1133">(S)-3-((6-Methoxypyridin-3-yl)-3-(5-(3-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)propyl)-2H-indazol-2-yl)propyl)propanoic acid</p>	¹ H NMR (500 MHz, MeOH- <i>d</i> ₄) δ 8.27 (s, 1H), 8.21 (d, <i>J</i> = 2.5 Hz, 1H), 7.79 (dd, <i>J</i> = 8.7, 2.6 Hz, 1H), 7.51 (d, <i>J</i> = 8.9 Hz, 1H), 7.43 (d, <i>J</i> = 7.4 Hz, 1H), 7.39 (s, 1H), 7.14 (dd, <i>J</i> = 8.9, 1.6 Hz, 1H), 6.76 (d, <i>J</i> = 8.7 Hz, 1H), 6.55 (d, <i>J</i> = 7.4 Hz, 1H), 6.14 (dd, <i>J</i> = 9.2, 6.0 Hz, 1H), 3.87 (s, 3H), 3.66 (dd, <i>J</i> = 16.7, 9.2 Hz, 1H), 3.38 - 3.33 (m, 2H), 2.77 (t, <i>J</i> = 7.1 Hz, 2H), 2.70 (t, <i>J</i> = 7.6 Hz, 2H), 2.61 (td, <i>J</i> = 6.1, 2.7 Hz, 2H), 2.13 - 2.01 (m, 2H), 1.80 (p, <i>J</i> = 6.2 Hz, 2H). LC/MS (<i>m/z</i>) = 472.3 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 740.	Example 9

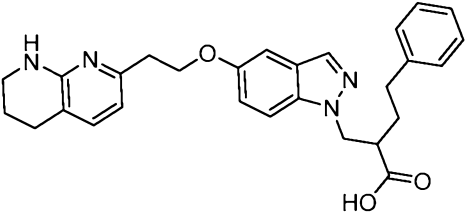
Example No.	Structure & Name	Analytical Data	Method
68	 <p>3-(3-(3,5-Dimethylpyrazol-1-yl)phenyl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid, TFA</p>	¹ H NMR (500 MHz, MeOH- <i>d</i> ₄) δ 7.92 (s, 1H), 7.48 (d, <i>J</i> = 9.1 Hz, 1H), 7.45 - 7.37 (m, 2H), 7.32 (s, 1H), 7.28 (t, <i>J</i> = 6.5 Hz, 2H), 6.99 (s, 1H), 6.91 (dd, <i>J</i> = 9.1, 2.0 Hz, 1H), 6.47 (d, <i>J</i> = 7.3 Hz, 1H), 6.28 (dd, <i>J</i> = 9.7, 5.0 Hz, 1H), 6.02 (s, 1H), 4.12 - 4.04 (m, 1H), 4.04 - 3.96 (m, 1H), 3.66 (dd, <i>J</i> = 15.9, 9.8 Hz, 1H), 3.40 - 3.34 (m, 2H), 3.20 (dd, <i>J</i> = 15.9, 5.0 Hz, 1H), 2.95 - 2.80 (m, 2H), 2.21 (s, 3H), 2.12 (s, 3H), 1.88 - 1.77 (m, 2H). LC/MS (<i>m/z</i>) = 537.4 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 1.5.	Example 3

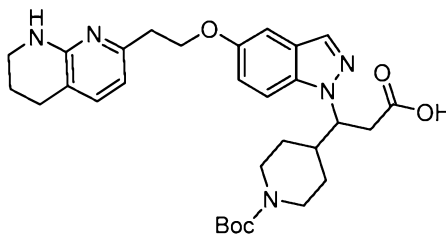
Example No.	Structure & Name	Analytical Data	Method
69	 <p>4-(4-(Benzyloxy)phenyl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)butanoic acid, TFA</p>	¹ H NMR (500 MHz, DMSO- <i>d</i> ₆) δ 7.91 (s, 1H), 7.39 - 7.32 (m, 4H), 7.32 - 7.26 (m, 2H), 7.08 (d, <i>J</i> = 2.0 Hz, 1H), 7.04 (d, <i>J</i> = 7.3 Hz, 1H), 6.88 (d, <i>J</i> = 8.5 Hz, 2H), 6.84 (dd, <i>J</i> = 9.1, 2.2 Hz, 1H), 6.72 (d, <i>J</i> = 8.5 Hz, 2H), 6.39 - 6.30 (m, 2H), 5.11 (d, <i>J</i> = 4.8 Hz, 1H), 4.95 (s, 2H), 4.20 (t, <i>J</i> = 6.8 Hz, 2H), 3.22 (br. s., 2H), 3.11 - 2.95 (m, 3H), 2.92 - 2.79 (m, 3H), 2.60 (t, <i>J</i> = 6.0 Hz, 2H), 1.81 - 1.66 (m, 2H). LC/MS (<i>m/z</i>) = 563.2 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 400.	Example 3

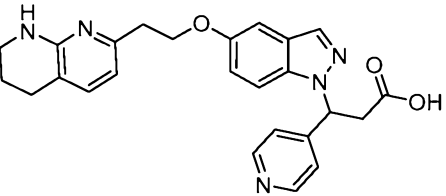
Example No.	Structure & Name	Analytical Data	Method
70	 <p data-bbox="416 958 871 1133">3-(6-Methoxypyridin-3-yl)-3-(3-methyl-5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, DMSO- <i>d</i> ₆) δ 8.22 (d, <i>J</i> = 2.5 Hz, 1H), 7.68 – 7.61 (m, 2H), 7.09 (d, <i>J</i> = 2.3 Hz, 1H), 7.06 (d, <i>J</i> = 7.3 Hz, 1H), 6.96 (dd, <i>J</i> = 9.0, 2.3 Hz, 1H), 6.72 (d, <i>J</i> = 8.6 Hz, 1H), 6.38 (d, <i>J</i> = 7.2 Hz, 1H), 6.34 (s, 1H), 6.08 (dd, <i>J</i> = 9.7, 5.4 Hz, 1H), 4.23 (td, <i>J</i> = 7.1, 2.6 Hz, 2H), 3.77 (s, 3H), 3.55 (dd, <i>J</i> = 16.4, 9.9 Hz, 1H), 3.23 (dq, <i>J</i> = 6.1, 2.8 Hz, 2H), 3.16 (dd, <i>J</i> = 16.4, 5.5 Hz, 1H), 2.90 (t, <i>J</i> = 6.9 Hz, 2H), 2.61 (t, <i>J</i> = 6.3 Hz, 2H), 2.44 (s, 3H), 1.75 (p, <i>J</i> = 6.0 Hz, 2H). LC/MS (<i>m/z</i>) = 488.0 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 953.	Example 3

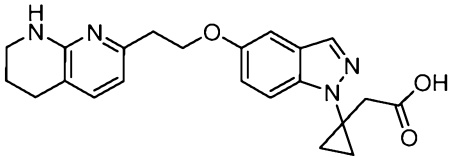
Example No.	Structure & Name	Analytical Data	Method
71	 <p>3-(1-Methyl-2-oxo-1,2-dihydropyridin-4-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid, TFA</p>	¹ H NMR (500 MHz, MeOH- <i>d</i> ₄) δ 7.93 (s, 1H), 7.52 (d, <i>J</i> = 7.0 Hz, 1H), 7.45 (d, <i>J</i> = 9.1 Hz, 1H), 7.23 (d, <i>J</i> = 7.3 Hz, 1H), 7.12 (s, 1H), 6.99 (dd, <i>J</i> = 9.0, 2.0 Hz, 1H), 6.51 (d, <i>J</i> = 7.3 Hz, 1H), 6.32 (d, <i>J</i> = 7.0 Hz, 1H), 6.27 (s, 1H), 6.07 (dd, <i>J</i> = 8.5, 5.8 Hz, 1H), 4.21 (t, <i>J</i> = 6.1 Hz, 2H), 3.52 - 3.42 (m, 4H), 3.41 - 3.37 (m, 2H), 3.11 (dd, <i>J</i> = 15.9, 5.7 Hz, 1H), 2.98 (t, <i>J</i> = 6.4 Hz, 2H), 2.71 (t, <i>J</i> = 6.2 Hz, 2H), 1.91 - 1.82 (m, 2H). LC/MS (<i>m/z</i>) = 474.3 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 64.	Example 3

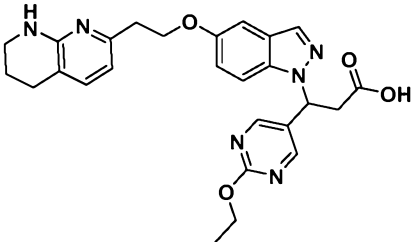
Example No.	Structure & Name	Analytical Data	Method
72	 <p data-bbox="414 1108 869 1232">(3<i>S</i>)-3-(6-Methoxypyridin-3-yl)-3-(6-(2-(1,2,3,4-tetrahydro-1,8-naphthyridin-2-yl)ethyl)-2<i>H</i>-indazol-2-yl)propanoic acid</p>	¹ H NMR (500 MHz, MeOH- <i>d</i> ₄) δ 8.28 (s, 1H), 8.15 (d, <i>J</i> = 2.5 Hz, 1H), 7.72 (dd, <i>J</i> = 8.7, 2.6 Hz, 1H), 7.53 (d, <i>J</i> = 8.6 Hz, 1H), 7.30 (d, <i>J</i> = 7.3 Hz, 1H), 7.17 (s, 1H), 6.87 (dd, <i>J</i> = 8.6, 1.4 Hz, 1H), 6.73 (d, <i>J</i> = 8.6 Hz, 1H), 6.37 (d, <i>J</i> = 7.3 Hz, 1H), 6.13 (dd, <i>J</i> = 9.4, 5.9 Hz, 1H), 3.86 (s, 3H), 3.49 (dd, <i>J</i> = 15.8, 9.4 Hz, 1H), 3.39 (t, <i>J</i> = 5.7 Hz, 2H), 3.18 (dd, <i>J</i> = 15.8, 5.9 Hz, 1H), 2.88 (h, <i>J</i> = 6.1, 5.0 Hz, 4H), 2.70 (t, <i>J</i> = 6.3 Hz, 2H), 1.86 (p, <i>J</i> = 6.1 Hz, 2H). LC/MS (<i>m/z</i>) = 457.9 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 8.2; Human α V β 1 IC ₅₀ (nM) = 740; Human α V β 3 IC ₅₀ (nM) = 4.4; Human α V β 5 IC ₅₀ (nM) = 1.5; and Human α V β 8 IC ₅₀ (nM) = 3,200.	Example 7

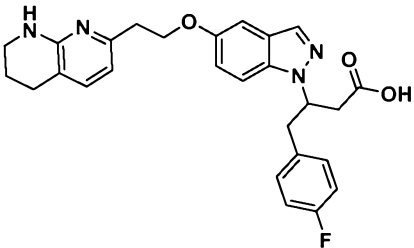
Example No.	Structure & Name	Analytical Data	Method
73	 <p>4-Phenyl-2-((5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)methyl)butanoic acid</p>	¹ H NMR (500 MHz, DMSO- <i>d</i> ₆) δ 7.90 (s, 1H), 7.52 (d, <i>J</i> = 9.1 Hz, 1H), 7.28 - 7.21 (m, 2H), 7.19 - 7.13 (m, 2H), 7.10 (d, <i>J</i> = 7.3 Hz, 2H), 7.06 (d, <i>J</i> = 7.2 Hz, 1H), 6.97 (dd, <i>J</i> = 9.0, 1.9 Hz, 1H), 6.38 (d, <i>J</i> = 7.2 Hz, 1H), 6.31 (br. s., 1H), 4.60 (dd, <i>J</i> = 14.1, 8.1 Hz, 1H), 4.45 (dd, <i>J</i> = 14.1, 6.0 Hz, 1H), 4.23 (t, <i>J</i> = 6.7 Hz, 2H), 3.23 (br. s., 2H), 2.97 - 2.81 (m, 3H), 2.61 (t, <i>J</i> = 5.9 Hz, 3H), 1.87 - 1.63 (m, 4H). LC/MS (<i>m/z</i>) = 471.3 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 1,800.	Example 3

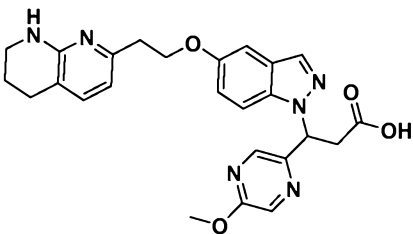
Example No.	Structure & Name	Analytical Data	Method
74	 <p>3-(1-(<i>tert</i>-Butoxycarbonyl)piperidin-4-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1<i>H</i>-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, DMSO- <i>d</i> ₆) δ 7.92 (s, 1H), 7.60 (d, <i>J</i> = 9.1 Hz, 1H), 7.13 (d, <i>J</i> = 1.9 Hz, 1H), 7.06 (d, <i>J</i> = 7.2 Hz, 1H), 6.97 (dd, <i>J</i> = 9.1, 2.1 Hz, 1H), 6.38 (d, <i>J</i> = 7.2 Hz, 1H), 6.34 (br. s., 1H), 4.84 - 4.73 (m, 1H), 4.24 (t, <i>J</i> = 6.8 Hz, 2H), 3.93 (br. s., 1H), 3.76 (br. s., 1H), 3.23 (br. s., 1H), 3.03 - 2.94 (m, 2H), 2.90 (t, <i>J</i> = 6.8 Hz, 2H), 2.61 (t, <i>J</i> = 6.1 Hz, 3H), 1.97 (d, <i>J</i> = 8.2 Hz, 1H), 1.83 - 1.66 (m, 3H), 1.34 (s, 9H), 1.13 - 0.87 (m, 2H), 0.78 (br. s., 1H). LC/MS (<i>m/z</i>) = 550.4 (<i>M</i> + <i>H</i>) ⁺ . Human α V β 6 IC ₅₀ (nM) = 260.	Example 3

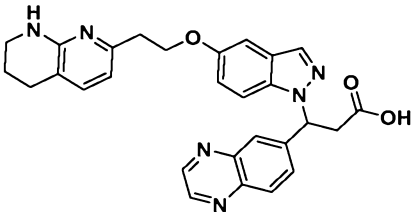
Example No.	Structure & Name	Analytical Data	Method
75	 <p>3-(Pyridin-4-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid, 2 TFA</p>	¹ H NMR (500 MHz, MeOH- <i>d</i> ₄) δ 8.77 (d, <i>J</i> = 6.9 Hz, 2H), 8.07 (s, 1H), 7.99 (d, <i>J</i> = 6.6 Hz, 2H), 7.60 (t, <i>J</i> = 8.9 Hz, 2H), 7.23 (d, <i>J</i> = 2.2 Hz, 1H), 7.08 (dd, <i>J</i> = 9.2, 2.3 Hz, 1H), 6.75 (d, <i>J</i> = 7.4 Hz, 1H), 6.54 (dd, <i>J</i> = 9.4, 5.2 Hz, 1H), 4.32 (td, <i>J</i> = 6.0, 2.6 Hz, 2H), 3.73 (dd, <i>J</i> = 17.1, 9.4 Hz, 1H), 3.53 - 3.44 (m, 3H), 3.19 (t, <i>J</i> = 5.8 Hz, 2H), 2.82 (t, <i>J</i> = 6.1 Hz, 2H), 2.02 - 1.89 (m, 2H). LC/MS (<i>m/z</i>) = 444.1 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 23.	Example 3

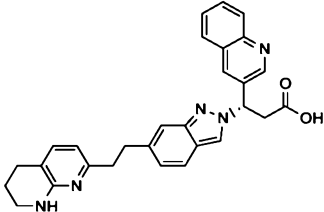
Example No.	Structure & Name	Analytical Data	Method
76	 <p>2-(1-(5-(2-(5,6,7,8-Tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)cyclopropyl)acetic acid, TFA</p>	¹ H NMR (500 MHz, MeOH- <i>d</i> ₄) δ 7.87 (s, 1H), 7.66 (d, <i>J</i> = 9.2 Hz, 1H), 7.50 (d, <i>J</i> = 7.3 Hz, 1H), 7.15 (d, <i>J</i> = 1.9 Hz, 1H), 7.05 (dd, <i>J</i> = 9.1, 2.1 Hz, 1H), 6.69 (d, <i>J</i> = 7.3 Hz, 1H), 4.29 (t, <i>J</i> = 6.1 Hz, 2H), 3.55 - 3.42 (m, 2H), 3.12 (t, <i>J</i> = 6.1 Hz, 2H), 2.87 - 2.73 (m, 4H), 1.93 (quin, <i>J</i> = 5.9 Hz, 2H), 1.35 (s, 4H). LC/MS (<i>m/z</i>) = 393.3 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 3,400.	Example 3

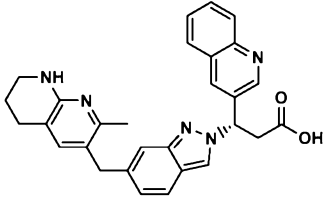
Example No.	Structure & Name	Analytical Data	Method
77	 <p>3-(2-Ethoxypyrimidin-5-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid, TFA</p>	¹ H NMR (500 MHz, chloroform- <i>d</i>) δ 8.51 (s, 2H), 7.96 (s, 1H), 7.36 - 7.30 (m, 2H), 7.06 (d, <i>J</i> = 2.2 Hz, 1H), 7.02 (dd, <i>J</i> = 9.1, 2.2 Hz, 1H), 6.50 (d, <i>J</i> = 7.2 Hz, 1H), 6.01 (dd, <i>J</i> = 8.4, 5.9 Hz, 1H), 4.38 (q, <i>J</i> = 7.2 Hz, 2H), 4.27 (t, <i>J</i> = 5.8 Hz, 2H), 3.73 (dd, <i>J</i> = 16.6, 8.7 Hz, 1H), 3.47 (t, <i>J</i> = 5.4 Hz, 2H), 3.31 (dd, <i>J</i> = 16.5, 5.8 Hz, 1H), 3.16 (t, <i>J</i> = 5.8 Hz, 2H), 2.74 (t, <i>J</i> = 6.2 Hz, 2H), 1.96 - 1.86 (m, 2H), 1.43 - 1.34 (m, 3H). LC/MS (<i>m/z</i>) = 489.1 (<i>M</i> +H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 5.8; Human α V β 1 IC ₅₀ (nM) = 170; Human α V β 3 IC ₅₀ (nM) = 2.8; Human α V β 5 IC ₅₀ (nM) = 0.39; and Human α V β 8 IC ₅₀ (nM) = 5,000.	Example 3

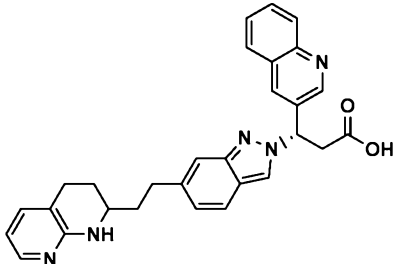
Example No.	Structure & Name	Analytical Data	Method
78	 <p data-bbox="435 862 850 1030">4-(4-Fluorophenyl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)butanoic acid, TFA</p>	¹ H NMR (500 MHz, chloroform- <i>d</i>) δ 9.99 (br. s., 1H), 7.96 (s, 1H), 7.33 (d, <i>J</i> = 7.2 Hz, 1H), 7.01 - 6.92 (m, 2H), 6.89 - 6.81 (m, 3H), 6.80 - 6.73 (m, 2H), 6.49 (d, <i>J</i> = 7.2 Hz, 1H), 5.09 - 4.94 (m, 1H), 4.28 (d, <i>J</i> = 5.0 Hz, 2H), 3.48 (t, <i>J</i> = 5.4 Hz, 2H), 3.32 - 3.21 (m, 2H), 3.19 - 3.12 (m, 3H), 3.03 (dd, <i>J</i> = 16.4, 4.3 Hz, 1H), 2.75 (t, <i>J</i> = 6.2 Hz, 2H), 1.97 - 1.87 (m, 2H). LC/MS (<i>m/z</i>) = 475.1 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 804.48.	Example 3

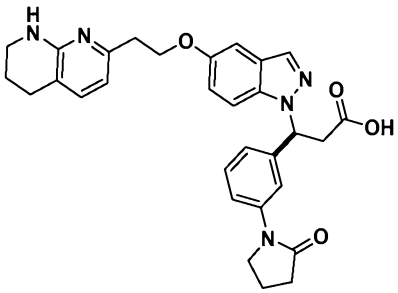
Example No.	Structure & Name	Analytical Data	Method
79	 <p>3-(5-Methoxypyrazin-2-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid, TFA</p>	¹ H NMR (500 MHz, chloroform- <i>d</i>) δ 8.17 (d, <i>J</i> = 1.1 Hz, 1H), 7.99 (s, 1H), 7.65 (s, 1H), 7.40 - 7.30 (m, 2H), 7.10 (d, <i>J</i> = 1.9 Hz, 1H), 7.02 (dd, <i>J</i> = 9.1, 2.2 Hz, 1H), 6.50 (d, <i>J</i> = 7.2 Hz, 1H), 6.20 (dd, <i>J</i> = 8.0, 5.5 Hz, 1H), 4.31 (t, <i>J</i> = 5.9 Hz, 2H), 3.90 (s, 3H), 3.75 - 3.56 (m, 2H), 3.49 (t, <i>J</i> = 5.5 Hz, 2H), 3.19 (t, <i>J</i> = 5.8 Hz, 2H), 2.75 (t, <i>J</i> = 6.2 Hz, 2H), 1.92 (quin, <i>J</i> = 5.9 Hz, 2H). LC/MS (<i>m/z</i>) = 475.1 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 33.	Example 3

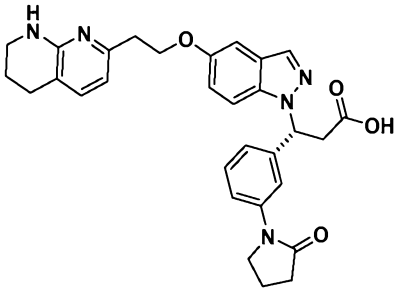
Example No.	Structure & Name	Analytical Data	Method
80	 <p>3-(Quinoxalin-6-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid, TFA</p>	¹ H NMR (500 MHz, chloroform- <i>d</i>) δ 8.85 (s, 2H), 8.08 - 8.01 (m, 2H), 7.97 (d, <i>J</i> = 1.7 Hz, 1H), 7.66 (dd, <i>J</i> = 8.8, 1.9 Hz, 1H), 7.34 - 7.28 (m, 2H), 7.10 (d, <i>J</i> = 1.9 Hz, 1H), 6.97 (dd, <i>J</i> = 9.1, 2.2 Hz, 1H), 6.48 (d, <i>J</i> = 7.2 Hz, 1H), 6.32 (dd, <i>J</i> = 8.9, 5.1 Hz, 1H), 4.28 (t, <i>J</i> = 5.8 Hz, 2H), 3.91 (dd, <i>J</i> = 16.5, 8.8 Hz, 1H), 3.52 - 3.41 (m, 3H), 3.16 (t, <i>J</i> = 5.8 Hz, 2H), 2.73 (t, <i>J</i> = 6.2 Hz, 2H), 1.96 - 1.85 (m, 2H). LC/MS (<i>m/z</i>) = 495.1 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 13.	Example 3

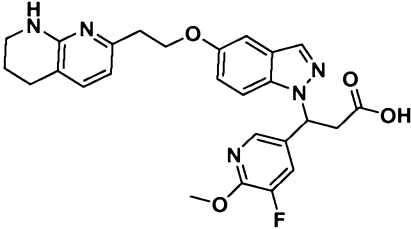
Example No.	Structure & Name	Analytical Data	Method
81	 <p data-bbox="421 891 866 1016">(S)-3-(Quinolin-3-yl)-3-(6-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethyl)-2H-indazol-2-yl)propanoic acid</p>	¹ H NMR (500 MHz, DMSO- <i>d</i> ₆) δ 8.99 (d, <i>J</i> = 2.3 Hz, 1H), 8.56 (s, 1H), 8.40 (d, <i>J</i> = 2.3 Hz, 1H), 7.97 (t, <i>J</i> = 8.1 Hz, 2H), 7.79 – 7.70 (m, 1H), 7.64 – 7.54 (m, 2H), 7.35 (s, 1H), 7.01 (d, <i>J</i> = 7.3 Hz, 1H), 6.91 (d, <i>J</i> = 8.6 Hz, 1H), 6.37 (t, <i>J</i> = 7.5 Hz, 1H), 6.28 (d, <i>J</i> = 7.2 Hz, 1H), 3.22 (s, 2H), 2.91 (dd, <i>J</i> = 17.9, 9.6 Hz, 2H), 2.78 – 2.70 (m, 2H), 2.58 (t, <i>J</i> = 6.3 Hz, 2H), 1.73 (t, <i>J</i> = 5.9 Hz, 2H). LC/MS (<i>m/z</i>) = 478.4 (M+H) ⁺ . Human αVβ6 IC ₅₀ (nM) = 3.2.	Example 7

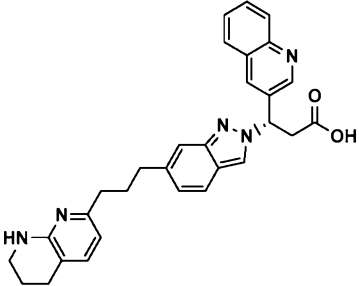
Example No.	Structure & Name	Analytical Data	Method
82	 <p data-bbox="421 808 866 981">(S)-3-(6-((2-Methyl-5,6,7,8-tetrahydro-1,8-naphthyridin-3-yl)methyl)-2H-indazol-2-yl)-3-(quinolin-3-yl)propanoic acid</p>	¹ H NMR (500 MHz, DMSO- <i>d</i> ₆) δ 8.99 (s, 1H), 8.61 (s, 1H), 8.42 (d, <i>J</i> = 2.2 Hz, 1H), 8.02 – 7.94 (m, 2H), 7.75 (t, <i>J</i> = 7.7 Hz, 1H), 7.62 (d, <i>J</i> = 8.4 Hz, 2H), 7.50 (s, 1H), 7.28 (s, 1H), 6.88 (d, <i>J</i> = 8.6 Hz, 1H), 6.38 (t, <i>J</i> = 4.7 Hz, 1H), 3.87 (d, <i>J</i> = 11.5 Hz, 2H), 3.79 – 3.67 (m, 1H), 3.35 (s, 2H), 2.66 (d, <i>J</i> = 6.7 Hz, 2H), 2.34 (s, 3H), 1.77 (m, 2H). LC/MS (<i>m/z</i>) = 478.3 (M+H) ⁺ . Human αVβ6 IC ₅₀ (nM) = 150.	Example 8

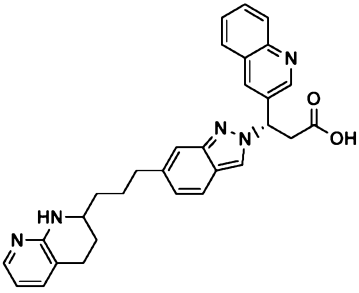
Example No.	Structure & Name	Analytical Data	Method
83	 <p>(3<i>S</i>)-3-(Quinolin-3-yl)-3-(6-(2-(1,2,3,4-tetrahydro-1,8-naphthyridin-2-yl)ethyl)-2<i>H</i>-indazol-2-yl)propanoic acid</p>	¹ H NMR (500 MHz, DMSO- <i>d</i> ₆) δ 8.97 (s, 1H), 8.58 (s, 1H), 8.43 (s, 1H), 7.97 (t, <i>J</i> = 7.1 Hz, 2H), 7.81 – 7.69 (m, 2H), 7.63 (d, <i>J</i> = 8.1 Hz, 3H), 7.43 (s, 1H), 6.96 (d, <i>J</i> = 8.7 Hz, 1H), 6.72 (t, <i>J</i> = 6.8 Hz, 1H), 6.37 (dd, <i>J</i> = 9.4, 5.8 Hz, 1H), 3.56 – 3.45 (m, 2H), 3.16 (s, 2H), 2.83 – 2.66 (m, 4H), 2.00 – 1.85 (m, 2H), 1.85 – 1.74 (m, 1H), 1.62 (d, <i>J</i> = 8.6 Hz, 1H). LC/MS (<i>m/z</i>) = 478.4 (<i>M</i> + <i>H</i>) ⁺ . Human αVβ6 IC ₅₀ (nM) = 4.7; Human αVβ1 IC ₅₀ (nM) = 360; Human αVβ3 IC ₅₀ (nM) = 4.1; Human αVβ5 IC ₅₀ (nM) = 2.2; and Human αVβ8 IC ₅₀ (nM) = 5,000.	Example 7

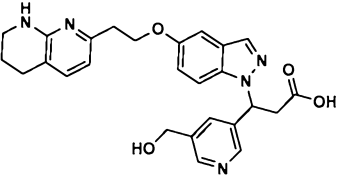
Example No.	Structure & Name	Analytical Data	Method
84	 <p>(S)-3-(3-(2-Oxopyrrolidin-1-yl)phenyl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid, TFA</p>	¹ H NMR (500 MHz, chloroform- <i>d</i>) δ 7.98 (s, 1H), 7.61 (br. s., 1H), 7.44 (d, <i>J</i> = 8.3 Hz, 1H), 7.33 (d, <i>J</i> = 7.4 Hz, 1H), 7.28 - 7.22 (m, 2H), 7.09 (d, <i>J</i> = 1.9 Hz, 1H), 6.96 (dd, <i>J</i> = 8.9, 1.8 Hz, 2H), 6.50 (d, <i>J</i> = 7.2 Hz, 1H), 6.07 (dd, <i>J</i> = 8.5, 5.2 Hz, 1H), 4.28 (t, <i>J</i> = 5.6 Hz, 2H), 3.88 - 3.71 (m, 3H), 3.48 (t, <i>J</i> = 5.5 Hz, 2H), 3.32 (dd, <i>J</i> = 16.5, 4.7 Hz, 1H), 3.17 (t, <i>J</i> = 5.8 Hz, 2H), 2.74 (t, <i>J</i> = 6.2 Hz, 2H), 2.60 (t, <i>J</i> = 8.1 Hz, 2H), 2.14 (quin, <i>J</i> = 7.4 Hz, 2H), 1.97 - 1.87 (m, 2H). LC/MS (<i>m/z</i>) = 526.5 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 120.	Example 16 and 17

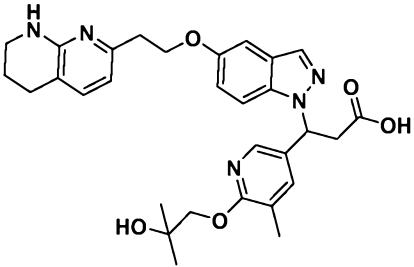
Example No.	Structure & Name	Analytical Data	Method
85	 <p data-bbox="416 952 871 1126">(R)-3-(3-(2-Oxopyrrolidin-1-yl)phenyl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid, TFA</p>	¹ H NMR (500 MHz, chloroform- <i>d</i>) δ 7.98 (s, 1H), 7.59 (s, 1H), 7.43 (d, <i>J</i> = 8.3 Hz, 1H), 7.32 (d, <i>J</i> = 7.2 Hz, 1H), 7.26 - 7.23 (m, 2H), 7.08 (s, 1H), 6.98 - 6.92 (m, 2H), 6.49 (d, <i>J</i> = 7.4 Hz, 1H), 6.06 (dd, <i>J</i> = 8.8, 5.0 Hz, 1H), 4.27 (t, <i>J</i> = 5.6 Hz, 2H), 3.86 - 3.70 (m, 3H), 3.53 - 3.43 (m, 2H), 3.32 (dd, <i>J</i> = 16.4, 4.5 Hz, 1H), 3.16 (t, <i>J</i> = 5.5 Hz, 2H), 2.74 (t, <i>J</i> = 5.9 Hz, 2H), 2.59 (t, <i>J</i> = 8.0 Hz, 2H), 2.20 - 2.08 (m, 2H), 1.96 - 1.85 (m, 2H). LC/MS (<i>m/z</i>) = 526.5 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 2.5.	Example 16 and 17

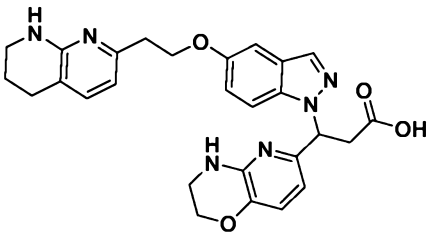
Example No.	Structure & Name	Analytical Data	Method
86	 <p>3-(5-Fluoro-6-methoxypyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid, TFA</p>	¹ H NMR (500 MHz, chloroform- <i>d</i>) δ 7.95 (s, 1H), 7.90 (d, <i>J</i> = 1.9 Hz, 1H), 7.36 - 7.28 (m, 3H), 7.05 (d, <i>J</i> = 1.7 Hz, 1H), 6.99 (dd, <i>J</i> = 9.1, 2.2 Hz, 1H), 6.49 (d, <i>J</i> = 7.4 Hz, 1H), 6.01 (dd, <i>J</i> = 8.8, 5.5 Hz, 1H), 4.25 (t, <i>J</i> = 5.8 Hz, 2H), 3.97 (s, 3H), 3.74 (dd, <i>J</i> = 16.6, 8.9 Hz, 1H), 3.47 (t, <i>J</i> = 5.5 Hz, 2H), 3.25 (dd, <i>J</i> = 16.5, 5.2 Hz, 1H), 3.15 (t, <i>J</i> = 5.6 Hz, 2H), 2.74 (t, <i>J</i> = 6.1 Hz, 2H), 1.95 - 1.84 (m, 2H). LC/MS (<i>m/z</i>) = 492.4 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 7.8.	Example 3

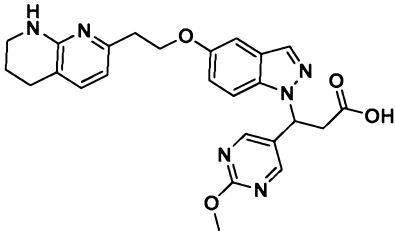
Example No.	Structure & Name	Analytical Data	Method
87	 <p data-bbox="427 1048 858 1216">(S)-3-(Quinolin-3-yl)-3-(6-(3-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)propyl)-2H-indazol-2-yl)propanoic acid</p>	¹ H NMR (500 MHz, DMSO- <i>d</i> ₆) δ 9.02 (t, <i>J</i> = 2.4 Hz, 1H), 8.59 (s, 1H), 8.43 (s, 1H), 7.98 (t, <i>J</i> = 7.8 Hz, 2H), 7.76 (t, <i>J</i> = 7.7 Hz, 1H), 7.67 – 7.55 (m, 2H), 7.35 (s, 1H), 7.02 (d, <i>J</i> = 7.2 Hz, 1H), 6.89 (d, <i>J</i> = 8.6 Hz, 1H), 6.39 (s, 1H), 6.26 (d, <i>J</i> = 7.3 Hz, 1H), 3.74 (dd, <i>J</i> = 17.0, 9.4 Hz, 1H), 3.21 (s, 2H), 2.63 (t, <i>J</i> = 7.6 Hz, 2H), 2.57 (d, <i>J</i> = 6.2 Hz, 2H), 2.44 (t, <i>J</i> = 7.7 Hz, 2H), 1.89 (d, <i>J</i> = 8.3 Hz, 2H), 1.72 (s, 2H). LC/MS (<i>m/z</i>) = 1.30. (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 1.3; Human α V β 1 IC ₅₀ (nM) = 10,000; Human α V β 3 IC ₅₀ (nM) = 2.3; Human α V β 5 IC ₅₀ (nM) = 3.2; and Human α V β 8 IC ₅₀ (nM) = 3,300.	Example 9

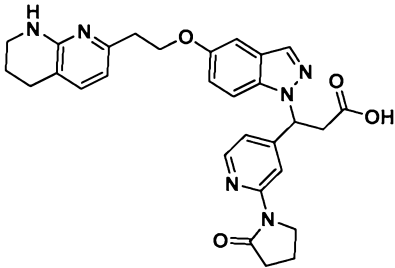
Example No.	Structure & Name	Analytical Data	Method
88	 <p data-bbox="422 974 865 1144">(3<i>S</i>)-3-(Quinolin-3-yl)-3-(6-(3-(1,2,3,4-tetrahydro-1,8-naphthyridin-2-yl)propyl)-2<i>H</i>-indazol-2-yl)propanoic acid</p>	¹ H NMR (500 MHz, MeOH- <i>d</i> ₄) δ 8.86 (d, <i>J</i> = 2.2 Hz, 1H), 8.44 (s, 1H), 8.36 (d, <i>J</i> = 2.2 Hz, 1H), 7.99 (d, <i>J</i> = 8.5 Hz, 1H), 7.92 (d, <i>J</i> = 8.2 Hz, 1H), 7.76 (ddd, <i>J</i> = 8.4, 6.7, 1.4 Hz, 1H), 7.68 – 7.56 (m, 3H), 7.37 (s, 1H), 7.24 (d, <i>J</i> = 7.2 Hz, 1H), 6.98 (d, <i>J</i> = 8.6 Hz, 1H), 6.49 (dd, <i>J</i> = 7.2, 5.3 Hz, 1H), 6.43 (t, <i>J</i> = 7.6 Hz, 1H), 3.54 (dd, <i>J</i> = 15.9, 8.3 Hz, 1H), 3.45 – 3.36 (m, 2H), 2.74 (t, <i>J</i> = 7.5 Hz, 2H), 2.70 (t, <i>J</i> = 6.4 Hz, 2H), 1.79 (p, <i>J</i> = 8.1 Hz, 2H), 1.65 – 1.47 (m, 3H). LC/MS (<i>m/z</i>) = 492.2 (<i>M</i> + <i>H</i>) ⁺ . Human α V β 6 IC ₅₀ (nM) = 20.	Example 9

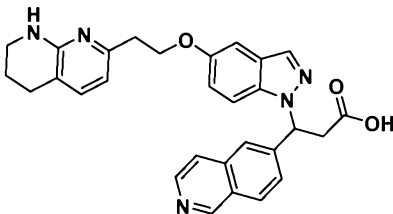
Example No.	Structure & Name	Analytical Data	Method
89	 <p data-bbox="416 965 874 1137">3-(5-(Hydroxymethyl)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, MeOH- <i>d</i> ₄) δ 8.54 (s, 1H), 8.43 (d, <i>J</i> = 2.2 Hz, 1H), 8.37 (s, 1H), 7.91 (s, 1H), 7.83 (s, 1H), 7.53 (d, <i>J</i> = 9.1 Hz, 1H), 7.17 – 7.10 (m, 2H), 7.00 (dd, <i>J</i> = 9.2, 2.4 Hz, 1H), 6.47 (d, <i>J</i> = 7.4 Hz, 1H), 6.27 (t, <i>J</i> = 7.4 Hz, 1H), 4.58 (s, 2H), 4.23 (t, <i>J</i> = 6.8 Hz, 2H), 3.08 – 2.91 (m, 4H), 2.69 (t, <i>J</i> = 6.3 Hz, 2H), 1.92 – 1.81 (m, 3H), 1.29 (t, <i>J</i> = 7.3 Hz, 3H). LC/MS (<i>m/z</i>) = 474.2 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 19; Human α V β 1 IC ₅₀ (nM) = 370; Human α V β 3 IC ₅₀ (nM) = 3.1; Human α V β 5 IC ₅₀ (nM) = 0.42; and Human α V β 8 IC ₅₀ (nM) = 5,000.	Example 3

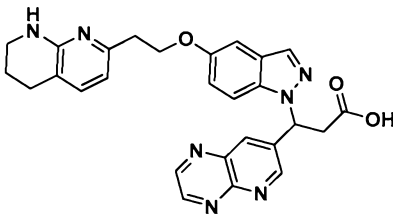
Example No.	Structure & Name	Analytical Data	Method
90	 <p data-bbox="427 920 863 1137">3-(6-(2-Hydroxy-2-methylpropoxy)-5-methylpyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid, TFA</p>	¹ H NMR (500 MHz, MeOH- <i>d</i> ₄) δ 7.90 (s, 1H), 7.61 (s, 1H), 7.54 (d, <i>J</i> = 9.1 Hz, 1H), 7.39 (s, 1H), 7.24 (d, <i>J</i> = 7.2 Hz, 1H), 7.09 (d, <i>J</i> = 1.6 Hz, 1H), 6.99 (dd, <i>J</i> = 9.1, 2.0 Hz, 1H), 6.51 (d, <i>J</i> = 7.3 Hz, 1H), 6.02 - 5.93 (m, 1H), 4.26 - 4.12 (m, 2H), 4.09 - 3.95 (m, 2H), 3.38 (t, <i>J</i> = 5.5 Hz, 3H), 3.10 (dd, <i>J</i> = 15.5, 5.8 Hz, 1H), 2.97 (t, <i>J</i> = 6.4 Hz, 2H), 2.71 (t, <i>J</i> = 6.1 Hz, 2H), 2.02 (s, 3H), 1.90 - 1.81 (m, 2H), 1.14 (d, <i>J</i> = 12.0 Hz, 6H). LC/MS (<i>m/z</i>) = 546.5 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 9.5.	Example 3

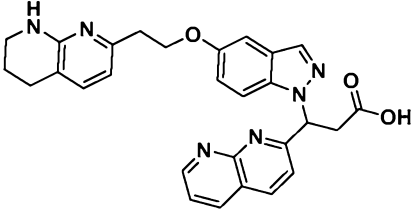
Example No.	Structure & Name	Analytical Data	Method
91	 <p data-bbox="427 853 858 1070">3-(3,4-Dihydro-2H-pyrido[3,2-<i>b</i>][1,4]oxazin-6-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid, 2 TFA</p>	¹ H NMR (500 MHz, chloroform- <i>d</i>) δ 7.94 (s, 1H), 7.66 (d, <i>J</i> = 9.1 Hz, 1H), 7.34 (d, <i>J</i> = 7.2 Hz, 1H), 7.12 - 7.03 (m, 2H), 6.99 (d, <i>J</i> = 9.1 Hz, 1H), 6.57 (d, <i>J</i> = 8.0 Hz, 1H), 6.50 (d, <i>J</i> = 7.2 Hz, 1H), 6.21 (t, <i>J</i> = 6.7 Hz, 1H), 4.28 (t, <i>J</i> = 5.6 Hz, 2H), 4.19 (dt, <i>J</i> = 8.5, 4.4 Hz, 2H), 3.69 - 3.52 (m, 3H), 3.51 - 3.39 (m, 3H), 3.16 (t, <i>J</i> = 5.6 Hz, 2H), 2.74 (t, <i>J</i> = 6.1 Hz, 2H), 1.97 - 1.85 (m, 2H). LC/MS (<i>m/z</i>) = 501.4 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 31.	Example 3

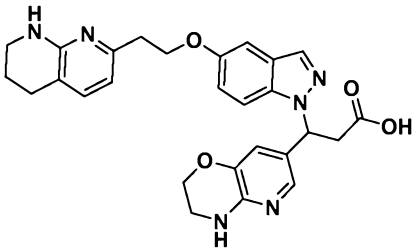
Example No.	Structure & Name	Analytical Data	Method
92	 <p>3-(2-Methoxypyrimidin-5-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid, TFA</p>	¹ H NMR (500 MHz, chloroform- <i>d</i>) δ 8.51 (s, 2H), 7.97 (s, 1H), 7.36 - 7.30 (m, 2H), 7.07 (d, <i>J</i> = 1.9 Hz, 1H), 7.03 (dd, <i>J</i> = 8.9, 2.3 Hz, 1H), 6.50 (d, <i>J</i> = 7.4 Hz, 1H), 6.02 (dd, <i>J</i> = 8.4, 5.9 Hz, 1H), 4.31 (t, <i>J</i> = 5.9 Hz, 2H), 3.97 (s, 3H), 3.74 (dd, <i>J</i> = 16.8, 8.5 Hz, 1H), 3.48 (t, <i>J</i> = 5.4 Hz, 2H), 3.33 (dd, <i>J</i> = 16.6, 5.9 Hz, 1H), 3.17 (t, <i>J</i> = 5.8 Hz, 2H), 2.74 (t, <i>J</i> = 6.1 Hz, 2H), 1.96 - 1.86 (m, 2H). LC/MS (<i>m/z</i>) = 475.5 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 31; Human α V β 1 IC ₅₀ (nM) = 210; Human α V β 3 IC ₅₀ (nM) = 2.4; Human α V β 5 IC ₅₀ (nM) = 0.39; and Human α V β 8 IC ₅₀ (nM) = 5,000.	Example 3

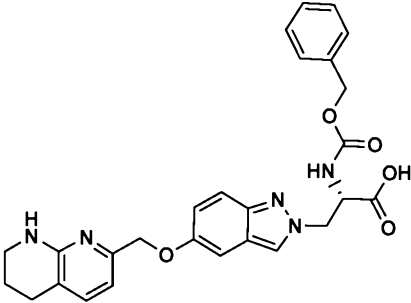
Example No.	Structure & Name	Analytical Data	Method
93	 <p>3-(2-(2-Oxopyrrolidin-1-yl)pyridin-4-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid, TFA</p>	¹ H NMR (500 MHz, chloroform- <i>d</i>) δ 9.82 (br. s., 1H), 8.41 (s, 1H), 8.25 (d, <i>J</i> = 5.2 Hz, 1H), 8.03 (s, 1H), 7.35 (d, <i>J</i> = 7.2 Hz, 1H), 7.12 (d, <i>J</i> = 1.9 Hz, 1H), 7.01 (dd, <i>J</i> = 9.2, 2.3 Hz, 1H), 6.76 (d, <i>J</i> = 5.5 Hz, 1H), 6.52 (d, <i>J</i> = 7.4 Hz, 1H), 6.10 (dd, <i>J</i> = 8.9, 4.8 Hz, 1H), 4.31 (t, <i>J</i> = 5.8 Hz, 2H), 4.08 (t, <i>J</i> = 7.2 Hz, 2H), 3.79 (dd, <i>J</i> = 16.5, 8.8 Hz, 1H), 3.51 (t, <i>J</i> = 5.5 Hz, 2H), 3.37 (dd, <i>J</i> = 16.5, 4.7 Hz, 1H), 3.19 (t, <i>J</i> = 5.8 Hz, 2H), 2.77 (t, <i>J</i> = 6.1 Hz, 2H), 2.71 - 2.66 (m, 2H), 2.22 - 2.09 (m, 2H), 2.00 - 1.89 (m, 2H). LC/MS (<i>m/z</i>) = 527.5 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 8.1.	Example 3

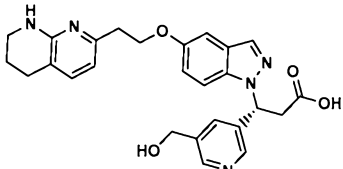
Example No.	Structure & Name	Analytical Data	Method
94	 <p>3-(Isoquinolin-6-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid, 2 TFA</p>	¹ H NMR (500 MHz, MeOH- <i>d</i> ₄) δ 9.57 (br. s., 1H), 8.51 (d, <i>J</i> = 6.3 Hz, 1H), 8.34 (d, <i>J</i> = 8.5 Hz, 1H), 8.26 (d, <i>J</i> = 3.6 Hz, 1H), 8.14 (s, 1H), 8.03 (s, 1H), 7.91 (d, <i>J</i> = 8.5 Hz, 1H), 7.64 - 7.53 (m, 2H), 7.21 (d, <i>J</i> = 1.9 Hz, 1H), 7.03 (dd, <i>J</i> = 9.1, 2.2 Hz, 1H), 6.73 (d, <i>J</i> = 7.2 Hz, 1H), 6.51 (dd, <i>J</i> = 9.5, 5.1 Hz, 1H), 4.31 (td, <i>J</i> = 5.9, 2.8 Hz, 2H), 3.81 (dd, <i>J</i> = 16.8, 9.6 Hz, 1H), 3.51 - 3.42 (m, 3H), 3.17 (t, <i>J</i> = 5.9 Hz, 2H), 2.80 (t, <i>J</i> = 6.2 Hz, 2H), 1.93 (quin, <i>J</i> = 5.9 Hz, 2H). LC/MS (<i>m/z</i>) = 494.2 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 13.	Example 3

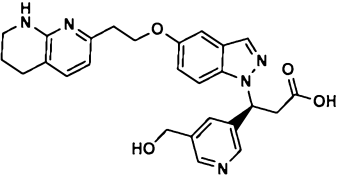
Example No.	Structure & Name	Analytical Data	Method
95	 <p>3-(Pyrido[2,3-<i>b</i>]pyrazin-7-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1<i>H</i>-indazol-1-yl)propanoic acid, 2 TFA</p>	¹ H NMR (500 MHz, chloroform- <i>d</i>) δ 9.26 (br. s., 1H), 9.07 (d, <i>J</i> = 1.4 Hz, 1H), 8.98 (d, <i>J</i> = 1.7 Hz, 1H), 8.47 (d, <i>J</i> = 1.9 Hz, 1H), 8.04 (s, 1H), 7.42 - 7.31 (m, 2H), 7.09 (d, <i>J</i> = 1.9 Hz, 1H), 7.02 (dd, <i>J</i> = 9.1, 2.2 Hz, 1H), 6.51 (d, <i>J</i> = 7.2 Hz, 1H), 6.45 - 6.36 (m, 1H), 4.32 - 4.24 (m, 2H), 3.58 - 3.52 (m, 1H), 3.49 (t, <i>J</i> = 5.5 Hz, 2H), 3.16 (t, <i>J</i> = 5.8 Hz, 2H), 2.75 (t, <i>J</i> = 6.2 Hz, 2H), 1.97 - 1.87 (m, 2H). LC/MS (<i>m/z</i>) = 496.2 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 30; Human α V β 1 IC ₅₀ (nM) = 210; Human α V β 3 IC ₅₀ (nM) = 2.4; and Human α V β 8 IC ₅₀ (nM) = 7,600.	Example 3

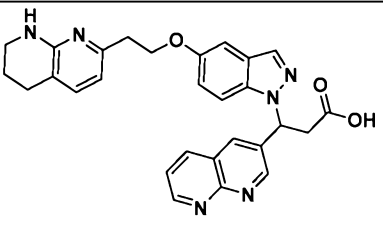
Example No.	Structure & Name	Analytical Data	Method
96	 <p data-bbox="427 965 863 1137">3-(1,8-Naphthyridin-2-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid, 2 TFA</p>	¹ H NMR (500 MHz, chloroform- <i>d</i>) δ 10.09 (br. s., 1H), 9.21 (br. s., 1H), 8.33 (d, <i>J</i> = 7.7 Hz, 1H), 8.12 (d, <i>J</i> = 8.5 Hz, 1H), 8.01 (s, 1H), 7.65 (dd, <i>J</i> = 7.8, 4.0 Hz, 1H), 7.42 (d, <i>J</i> = 9.1 Hz, 1H), 7.33 (d, <i>J</i> = 7.2 Hz, 1H), 7.13 (s, 1H), 7.08 (d, <i>J</i> = 8.3 Hz, 1H), 6.98 (d, <i>J</i> = 7.4 Hz, 1H), 6.58 (t, <i>J</i> = 6.6 Hz, 1H), 6.50 (d, <i>J</i> = 7.2 Hz, 1H), 4.32 (t, <i>J</i> = 5.8 Hz, 2H), 3.99 (dd, <i>J</i> = 16.2, 6.9 Hz, 1H), 3.57 - 3.44 (m, 3H), 3.19 (t, <i>J</i> = 5.8 Hz, 2H), 2.75 (t, <i>J</i> = 6.1 Hz, 2H), 1.92 (quin, <i>J</i> = 5.8 Hz, 2H). LC/MS (<i>m/z</i>) = 495.2 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 7.2.	Example 3

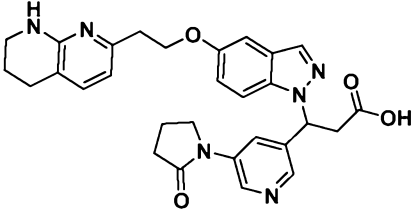
Example No.	Structure & Name	Analytical Data	Method
97	 <p>3-(3,4-Dihydro-2H-pyrido[3,2-b][1,4]oxazin-7-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid, 2 TFA</p>	¹ H NMR (500 MHz, chloroform- <i>d</i>) δ 10.61 (br. s., 1H), 10.12 (br. s., 1H), 7.96 (s, 1H), 7.76 (s, 1H), 7.37 - 7.28 (m, 2H), 7.11 (d, <i>J</i> = 1.9 Hz, 1H), 7.01 (dd, <i>J</i> = 8.9, 1.8 Hz, 1H), 6.49 (d, <i>J</i> = 7.2 Hz, 1H), 5.95 (t, <i>J</i> = 7.0 Hz, 1H), 4.29 (t, <i>J</i> = 4.8 Hz, 2H), 4.24 - 4.09 (m, 2H), 3.67 - 3.53 (m, 3H), 3.51 - 3.36 (m, 3H), 3.17 (t, <i>J</i> = 5.8 Hz, 2H), 2.74 (t, <i>J</i> = 6.1 Hz, 2H), 1.97 - 1.86 (m, 2H). LC/MS (<i>m/z</i>) = 501.2 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 10; Human α V β 1 IC ₅₀ (nM) = 100; Human α V β 3 IC ₅₀ (nM) = 2.6; Human α V β 5 IC ₅₀ (nM) = 0.59; and Human α V β 8 IC ₅₀ (nM) = 5,000.	Example 3

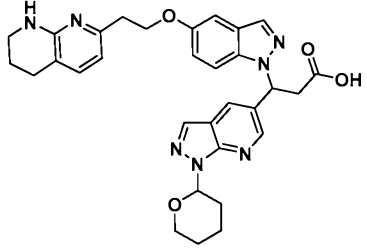
Example No.	Structure & Name	Analytical Data	Method
98	 <p>(S)-2-(((Benzyloxy)carbonyl)amino)-3-(5-((5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)methoxy)-2H-indazol-2-yl)propanoic acid</p>	¹ H NMR (500 MHz, MeOH- <i>d</i> ₄) δ 7.84 (s, 1H), 7.46 (dd, <i>J</i> = 12.5, 9.2 Hz, 1H), 7.36 (d, <i>J</i> = 7.3 Hz, 1H), 7.31 – 7.22 (m, 4H), 7.22 – 7.17 (m, 1H), 7.12 (d, <i>J</i> = 2.3 Hz, 1H), 7.03 (dd, <i>J</i> = 9.1, 2.4 Hz, 1H), 6.75 – 6.69 (m, 1H), 4.79 (dd, <i>J</i> = 14.5, 4.3 Hz, 1H), 4.68 (dd, <i>J</i> = 14.4, 7.4 Hz, 1H), 4.61 – 4.46 (m, 1H), 3.78 (s, 2H), 3.43 (q, <i>J</i> = 5.4 Hz, 2H), 2.77 (t, <i>J</i> = 6.2 Hz, 3H), 1.90 (q, <i>J</i> = 5.9 Hz, 2H). LC/MS (<i>m/z</i>) = 502.1 (M+H) ⁺ . Human αVβ6 IC ₅₀ (nM) = 160.	Example 4

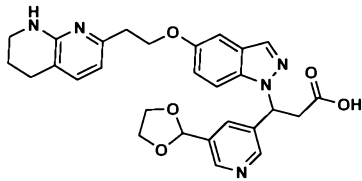
Example No.	Structure & Name	Analytical Data	Method
99	 <p data-bbox="416 824 869 996">(R)-3-(5-(Hydroxymethyl)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, MeOH- <i>d</i> ₄) δ 8.54 (s, 1H), 8.43 (d, <i>J</i> = 2.2 Hz, 1H), 8.37 (s, 1H), 7.91 (s, 1H), 7.83 (s, 1H), 7.53 (d, <i>J</i> = 9.1 Hz, 1H), 7.17 – 7.10 (m, 2H), 7.00 (dd, <i>J</i> = 9.2, 2.4 Hz, 1H), 6.47 (d, <i>J</i> = 7.4 Hz, 1H), 6.27 (t, <i>J</i> = 7.4 Hz, 1H), 4.58 (s, 2H), 4.23 (t, <i>J</i> = 6.8 Hz, 2H), 3.08 – 2.91 (m, 4H), 2.69 (t, <i>J</i> = 6.3 Hz, 2H), 1.92 – 1.81 (m, 3H), 1.29 (t, <i>J</i> = 7.3 Hz, 3H). LC/MS (<i>m/z</i>) = 474.2 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 340.	Example 16 and 17

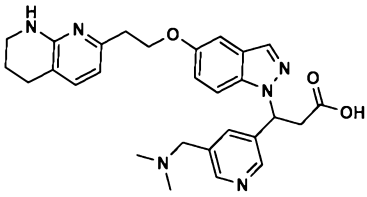
Example No.	Structure & Name	Analytical Data	Method
100	 <p>(S)-3-(5-(Hydroxymethyl)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, MeOH- <i>d</i> ₄) δ 8.54 (s, 1H), 8.43 (d, <i>J</i> = 2.2 Hz, 1H), 8.37 (s, 1H), 7.91 (s, 1H), 7.83 (s, 1H), 7.53 (d, <i>J</i> = 9.1 Hz, 1H), 7.17 – 7.10 (m, 2H), 7.00 (dd, <i>J</i> = 9.2, 2.4 Hz, 1H), 6.47 (d, <i>J</i> = 7.4 Hz, 1H), 6.27 (t, <i>J</i> = 7.4 Hz, 1H), 4.58 (s, 2H), 4.23 (t, <i>J</i> = 6.8 Hz, 2H), 3.08 – 2.91 (m, 4H), 2.69 (t, <i>J</i> = 6.3 Hz, 2H), 1.92 – 1.81 (m, 3H), 1.29 (t, <i>J</i> = 7.3 Hz, 3H). LC/MS (<i>m/z</i>) = 474.2 (<i>M</i> +H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 14; Human α V β 1 IC ₅₀ (nM) = 140; Human α V β 3 IC ₅₀ (nM) = 2.9; Human α V β 5 IC ₅₀ (nM) = 0.41; and Human α V β 8 IC ₅₀ (nM) = 4,400.	Example 16 and 17

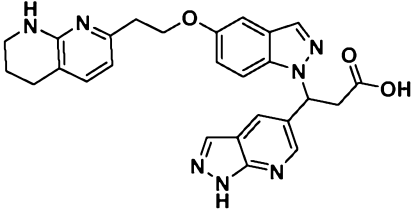
Example No.	Structure & Name	Analytical Data	Method
101	 <p>3-(1,8-Naphthyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid, 2 TFA</p>	¹ H NMR (500 MHz, chloroform- <i>d</i>) δ 10.03 (br s, 1H), 9.29 (d, <i>J</i> = 2.2 Hz, 1H), 9.18 (dd, <i>J</i> = 4.3, 1.8 Hz, 1H), 8.31 - 8.23 (m, 2H), 8.03 (s, 1H), 7.60 (dd, <i>J</i> = 8.1, 4.3 Hz, 1H), 7.40 - 7.31 (m, 2H), 7.11 (d, <i>J</i> = 1.9 Hz, 1H), 7.00 (dd, <i>J</i> = 9.1, 2.2 Hz, 1H), 6.50 (d, <i>J</i> = 7.4 Hz, 1H), 6.41 (t, <i>J</i> = 7.2 Hz, 1H), 4.36 - 4.27 (m, 2H), 3.84 (dd, <i>J</i> = 16.5, 7.7 Hz, 1H), 3.54 (dd, <i>J</i> = 16.8, 6.6 Hz, 1H), 3.48 (br t, <i>J</i> = 5.4 Hz, 2H), 3.18 (t, <i>J</i> = 5.8 Hz, 2H), 2.75 (br t, <i>J</i> = 6.1 Hz, 2H), 1.96 - 1.88 (m, 2H). LC/MS (<i>m/z</i>) = 495.4 (<i>M</i> +H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 5.7; Human α V β 1 IC ₅₀ (nM) = 100; Human α V β 3 IC ₅₀ (nM) = 1.6; and Human α V β 8 IC ₅₀ (nM) = 2,300.	Example 3

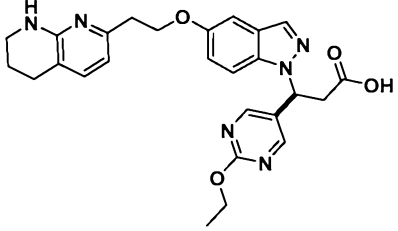
Example No.	Structure & Name	Analytical Data	Method
102	 <p>3-(5-(2-Oxopyrrolidin-1-yl)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid, 2 TFA</p>	¹ H NMR (500 MHz, chloroform- <i>d</i>) δ 9.98 (br s, 1H), 9.16 (d, <i>J</i> = 1.9 Hz, 1H), 8.80 (s, 1H), 8.47 (s, 1H), 8.00 (s, 1H), 7.37 - 7.33 (m, 2H), 7.10 (d, <i>J</i> = 2.2 Hz, 1H), 6.88 (dd, <i>J</i> = 9.1, 2.2 Hz, 1H), 6.50 (d, <i>J</i> = 7.4 Hz, 1H), 6.41 (t, <i>J</i> = 7.0 Hz, 1H), 4.37 - 4.22 (m, 2H), 3.87 - 3.81 (m, 2H), 3.81 - 3.72 (m, 2H), 3.41 (br dd, <i>J</i> = 17.1, 6.3 Hz, 1H), 3.17 (br t, <i>J</i> = 5.9 Hz, 2H), 2.76 (br t, <i>J</i> = 6.1 Hz, 2H), 2.68 - 2.59 (m, 2H), 2.23 (br t, <i>J</i> = 7.7 Hz, 2H), 1.99 - 1.88 (m, 2H). LC/MS (<i>m/z</i>) = 527.5 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 4.3; Human α V β 1 IC ₅₀ (nM) = 39; Human α V β 3 IC ₅₀ (nM) = 2.2; and Human α V β 8 IC ₅₀ (nM) = 1,300.	Example 3

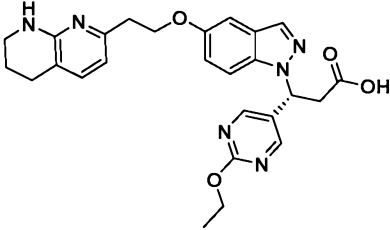
Example No.	Structure & Name	Analytical Data	Method
103	 <p>3-(5-(2-(5,6,7,8-Tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)-3-(1-(tetrahydro-2H-pyran-2-yl)pyrazolo[3,4-b]pyridin-5-yl)propanoic acid, TFA</p>	¹ H NMR (500 MHz, chloroform- <i>d</i>) δ 9.84 (br s, 1H), 8.66 - 8.55 (m, 1H), 8.07 - 7.96 (m, 2H), 7.37 - 7.30 (m, 2H), 7.08 (s, 1H), 6.98 (br d, <i>J</i> = 9.1 Hz, 1H), 6.49 (d, <i>J</i> = 7.4 Hz, 1H), 6.24 (br dd, <i>J</i> = 8.3, 5.8 Hz, 1H), 6.05 (ddd, <i>J</i> = 10.2, 5.1, 2.1 Hz, 1H), 4.28 (br t, <i>J</i> = 5.4 Hz, 2H), 4.09 (br d, <i>J</i> = 10.5 Hz, 1H), 3.90 - 3.75 (m, 2H), 3.48 (br t, <i>J</i> = 5.1 Hz, 2H), 3.39 (dt, <i>J</i> = 16.4, 4.7 Hz, 1H), 3.21 - 3.13 (m, 2H), 2.74 (br t, <i>J</i> = 6.1 Hz, 2H), 2.68 - 2.56 (m, 1H), 2.23 - 2.09 (m, 1H), 2.01 - 1.87 (m, 3H), 1.78 (br t, <i>J</i> = 9.1 Hz, 2H), 1.63 (br d, <i>J</i> = 7.2 Hz, 1H). LC/MS (<i>m/z</i>) = 568.6 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 5.4.	Example 3

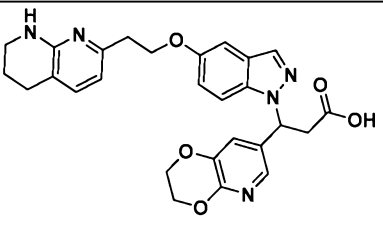
Example No.	Structure & Name	Analytical Data	Method
104	 <p>3-(5-(1,3-Dioxolan-2-yl)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, DMSO- <i>d</i> ₆) δ 8.61 (s, 1H), 8.47 (s, 1H), 8.00 (s, 1H), 7.75 (s, 1H), 7.67 (d, <i>J</i> = 9.2 Hz, 1H), 7.17 (d, <i>J</i> = 2.4 Hz, 1H), 7.06 (d, <i>J</i> = 7.3 Hz, 1H), 7.01 – 6.92 (m, 1H), 6.37 (d, <i>J</i> = 7.3 Hz, 1H), 6.25 (d, <i>J</i> = 8.9 Hz, 1H), 5.72 (s, 1H), 4.22 (d, <i>J</i> = 7.3 Hz, 2H), 3.99 (d, <i>J</i> = 6.4 Hz, 1H), 3.92 (d, <i>J</i> = 6.2 Hz, 1H), 3.68 (d, <i>J</i> = 12.0 Hz, 1H), 3.55 – 3.42 (m, 1H), 3.26 – 3.13 (m, 4H), 2.87 (t, <i>J</i> = 6.7 Hz, 2H), 2.59 (t, <i>J</i> = 6.3 Hz, 2H), 1.78 – 1.63 (m, 2H). LC/MS (<i>m/z</i>) = 516.3 (M+H) ⁺ . Human αVβ6 IC ₅₀ (nM) = 12; Human αVβ1 IC ₅₀ (nM) = 30; Human αVβ3 IC ₅₀ (nM) = 2.61; Human αVβ5 IC ₅₀ (nM) = 0.28; and Human αVβ8 IC ₅₀ (nM) = 1,800.	Example 3

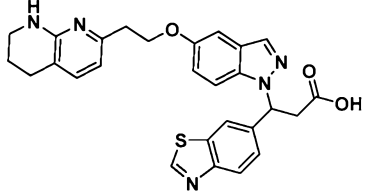
Example No.	Structure & Name	Analytical Data	Method
105	 <p data-bbox="411 996 871 1171">3-(5-((Dimethylamino)methyl)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid</p>	¹ H NMR (400 MHz, MeCN- <i>d</i> ₃) δ 9.51 (s, 1H), 8.67 (d, <i>J</i> = 2.1 Hz, 1H), 8.58 (d, <i>J</i> = 2.0 Hz, 1H), 8.07 (t, <i>J</i> = 2.1 Hz, 1H), 7.96 (d, <i>J</i> = 0.8 Hz, 1H), 7.51 (d, <i>J</i> = 9.1 Hz, 1H), 7.44 (d, <i>J</i> = 7.4 Hz, 1H), 7.17 (d, <i>J</i> = 2.4 Hz, 1H), 7.01 (dd, <i>J</i> = 9.1, 2.3 Hz, 1H), 6.58 (d, <i>J</i> = 7.3 Hz, 1H), 6.22 (dd, <i>J</i> = 9.2, 5.6 Hz, 1H), 4.32 - 4.24 (m, 3H), 4.23 - 4.13 (m, 2H), 3.69 (dd, <i>J</i> = 16.8, 9.2 Hz, 1H), 3.41 (t, <i>J</i> = 5.7 Hz, 2H), 3.34 (dd, <i>J</i> = 16.8, 5.6 Hz, 1H), 3.11 (t, <i>J</i> = 6.2 Hz, 2H), 2.72 (d, <i>J</i> = 7.7 Hz, 2H), 2.70 (s, 6H), 1.86 (dq, <i>J</i> = 7.0, 5.6 Hz, 2H). LC/MS (<i>m/z</i>) = 501.1 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 10.	Example 3

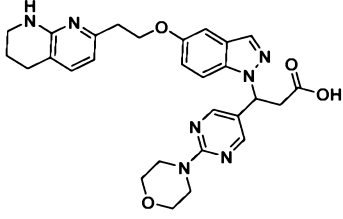
Example No.	Structure & Name	Analytical Data	Method
106	 <p>3-(1<i>H</i>-Pyrazolo[3,4-<i>b</i>]pyridin-5-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1<i>H</i>-indazol-1-yl)propanoic acid, 2 TFA</p>	<p>¹H NMR (500 MHz, MeOH-<i>d</i>₄) δ 8.59 (d, <i>J</i> = 1.9 Hz, 1H), 8.27 (d, <i>J</i> = 1.9 Hz, 1H), 8.08 (s, 1H), 8.00 (s, 1H), 7.66 - 7.58 (m, 2H), 7.20 (d, <i>J</i> = 1.9 Hz, 1H), 7.06 (dd, <i>J</i> = 9.2, 2.3 Hz, 1H), 6.76 (d, <i>J</i> = 7.2 Hz, 1H), 6.41 (dd, <i>J</i> = 9.4, 5.5 Hz, 1H), 4.33 (br t, <i>J</i> = 5.6 Hz, 2H), 3.80 (dd, <i>J</i> = 16.8, 9.4 Hz, 1H), 3.51 - 3.46 (m, 2H), 3.45 - 3.38 (m, 1H), 3.18 (t, <i>J</i> = 5.8 Hz, 2H), 2.80 (br t, <i>J</i> = 6.1 Hz, 2H), 2.01 - 1.89 (m, 2H).</p> <p>LC/MS (<i>m/z</i>) = 484.5</p> <p>(<i>M</i>+<i>H</i>)⁺. Human αVβ6 IC₅₀ (nM) = 23; Human αVβ1 IC₅₀ (nM) = 78; Human αVβ3 IC₅₀ (nM) = 1.3; Human αVβ5 IC₅₀ (nM) = 2.6; and Human αVβ8 IC₅₀ (nM) = 3,000.</p>	Example 3

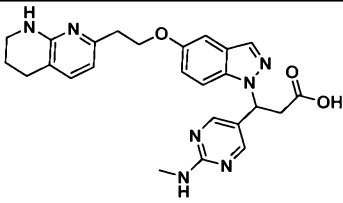
Example No.	Structure & Name	Analytical Data	Method
107	 <p data-bbox="421 667 865 842">(S)-3-(2-Ethoxypyrimidin-5-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid, TFA</p>	¹ H NMR (500 MHz, MeOH- <i>d</i> ₄) δ 8.58 (s, 2H), 8.00 (s, 1H), 7.67 - 7.61 (m, 2H), 7.21 (d, <i>J</i> = 2.2 Hz, 1H), 7.09 (dd, <i>J</i> = 9.1, 2.5 Hz, 1H), 6.77 (d, <i>J</i> = 7.4 Hz, 1H), 6.24 (dd, <i>J</i> = 9.1, 5.8 Hz, 1H), 4.40 (q, <i>J</i> = 7.0 Hz, 2H), 4.34 (t, <i>J</i> = 5.9 Hz, 2H), 3.68 (dd, <i>J</i> = 16.6, 9.2 Hz, 1H), 3.55 - 3.49 (m, 2H), 3.20 (t, <i>J</i> = 5.8 Hz, 2H), 2.83 (t, <i>J</i> = 6.2 Hz, 2H), 1.96 (br dd, <i>J</i> = 6.3, 5.2 Hz, 2H), 1.38 (t, <i>J</i> = 7.2 Hz, 3H). LC/MS (<i>m/z</i>) = 489.5 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 15; Human α V β 1 IC ₅₀ (nM) = 64; Human α V β 3 IC ₅₀ (nM) = 2.4; Human α V β 5 IC ₅₀ (nM) = 0.32; and Human α V β 8 IC ₅₀ (nM) = 300.	Example 16 and 17

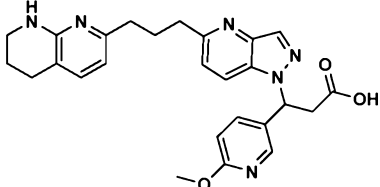
Example No.	Structure & Name	Analytical Data	Method
108	 <p data-bbox="422 667 863 840">(R)-3-(2-Ethoxypyrimidin-5-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid, TFA</p>	¹ H NMR (500 MHz, MeOH- <i>d</i> ₄) δ 8.58 (s, 2H), 8.00 (s, 1H), 7.67 - 7.61 (m, 2H), 7.21 (d, <i>J</i> = 2.2 Hz, 1H), 7.09 (dd, <i>J</i> = 9.1, 2.5 Hz, 1H), 6.77 (d, <i>J</i> = 7.4 Hz, 1H), 6.24 (dd, <i>J</i> = 9.1, 5.8 Hz, 1H), 4.40 (q, <i>J</i> = 7.0 Hz, 2H), 4.34 (t, <i>J</i> = 5.9 Hz, 2H), 3.68 (dd, <i>J</i> = 16.6, 9.2 Hz, 1H), 3.55 - 3.49 (m, 2H), 3.20 (t, <i>J</i> = 5.8 Hz, 2H), 2.83 (t, <i>J</i> = 6.2 Hz, 2H), 1.96 (br dd, <i>J</i> = 6.3, 5.2 Hz, 2H), 1.38 (t, <i>J</i> = 7.2 Hz, 3H). LC/MS (<i>m/z</i>) = 489.5 (<i>M</i> +H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 390.	Example 16 and 17

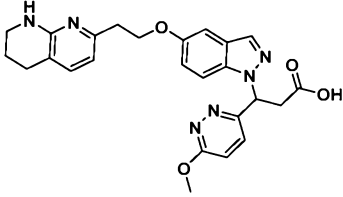
Example No.	Structure & Name	Analytical Data	Method
109	 <p>3-(2,3-Dihydro-[1,4]dioxino[2,3-b]pyridin-7-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid, 2 TFA</p>	<p>¹H NMR (500 MHz, chloroform-<i>d</i>) δ 7.99 (s, 1H), 7.95 (s, 1H), 7.38 - 7.30 (m, 2H), 7.28 - 7.25 (m, 1H), 7.09 (d, <i>J</i> = 2.2 Hz, 1H), 7.00 (dd, <i>J</i> = 9.1, 1.9 Hz, 1H), 6.53 (d, <i>J</i> = 7.2 Hz, 1H), 6.06 (t, <i>J</i> = 7.2 Hz, 1H), 4.48 - 4.40 (m, 2H), 4.34 - 4.27 (m, 2H), 4.27 - 4.20 (m, 2H), 3.74 (dd, <i>J</i> = 16.5, 8.5 Hz, 1H), 3.51 (br t, <i>J</i> = 5.5 Hz, 2H), 3.31 (br dd, <i>J</i> = 16.5, 5.5 Hz, 1H), 3.22 - 3.15 (m, 2H), 2.77 (br t, <i>J</i> = 6.1 Hz, 2H), 1.99 - 1.91 (m, 2H).</p> <p>LC/MS (<i>m/z</i>) = 502.2 (M+H)⁺. Human αVβ6 IC₅₀ (nM) = 15; Human αVβ1 IC₅₀ (nM) = 62; Human αVβ3 IC₅₀ (nM) = 1.3; and Human αVβ8 IC₅₀ (nM) = 9,300.</p>	Example 3

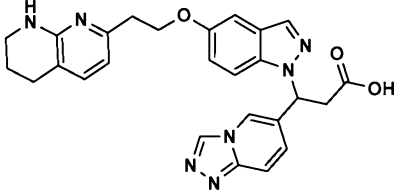
Example No.	Structure & Name	Analytical Data	Method
110	 <p>3-(Benzo[d]thiazol-6-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid, TFA</p>	¹ H NMR (500 MHz, chloroform- <i>d</i>) δ 8.98 (s, 1H), 8.06 (d, <i>J</i> = 8.5 Hz, 1H), 8.04 (s, 1H), 7.79 (s, 1H), 7.44 (dd, <i>J</i> = 8.5, 1.4 Hz, 1H), 7.34 - 7.30 (m, 1H), 7.26 (s, 1H), 7.12 (d, <i>J</i> = 1.7 Hz, 1H), 6.98 (dd, <i>J</i> = 8.9, 2.1 Hz, 1H), 6.48 (d, <i>J</i> = 7.4 Hz, 1H), 6.23 (dd, <i>J</i> = 9.1, 4.7 Hz, 1H), 4.30 (br t, <i>J</i> = 5.8 Hz, 2H), 3.87 (dd, <i>J</i> = 16.4, 9.2 Hz, 1H), 3.48 (br t, <i>J</i> = 5.2 Hz, 2H), 3.40 (dd, <i>J</i> = 16.2, 4.7 Hz, 1H), 3.18 (br t, <i>J</i> = 5.6 Hz, 2H), 2.74 (br t, <i>J</i> = 6.2 Hz, 2H), 1.95 - 1.89 (m, 2H). LC/MS (<i>m/z</i>) = 500.2 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 19.	Example 3

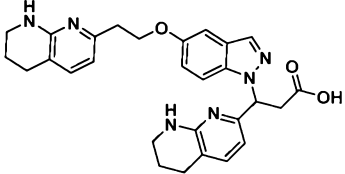
Example No.	Structure & Name	Analytical Data	Method
111	 <p>3-(2-Morpholinopyrimidin-5-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid, TFA</p>	¹ H NMR (500 MHz, chloroform- <i>d</i>) δ 8.60 (s, 2H), 7.99 (s, 1H), 7.40 - 7.33 (m, 2H), 7.10 (d, <i>J</i> = 1.9 Hz, 1H), 6.99 (d, <i>J</i> = 8.4 Hz, 1H), 6.54 (d, <i>J</i> = 7.2 Hz, 1H), 6.05 (t, <i>J</i> = 7.3 Hz, 1H), 4.31 (br t, <i>J</i> = 5.5 Hz, 2H), 3.89 - 3.81 (m, 4H), 3.81 - 3.75 (m, 4H), 3.66 (br d, <i>J</i> = 7.4 Hz, 1H), 3.52 (br t, <i>J</i> = 5.5 Hz, 2H), 3.39 (br dd, <i>J</i> = 16.6, 7.0 Hz, 1H), 3.22 - 3.15 (m, 2H), 2.78 (br t, <i>J</i> = 6.1 Hz, 2H), 2.00 - 1.91 (m, 2H). LC/MS (<i>m/z</i>) = 530.3 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 24.	Example 3

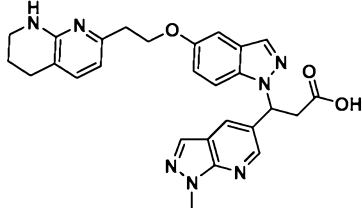
Example No.	Structure & Name	Analytical Data	Method
112	 <p>3-(2-(Methylamino)pyrimidin-5-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid, 2 TFA</p>	¹ H NMR (500 MHz, MeOH-d ₄) δ 8.43 (s, 2H), 7.98 (s, 1H), 7.67 - 7.58 (m, 2H), 7.20 (d, <i>J</i> = 2.2 Hz, 1H), 7.09 (dd, <i>J</i> = 9.1, 2.2 Hz, 1H), 6.75 (d, <i>J</i> = 7.4 Hz, 1H), 6.13 (dd, <i>J</i> = 9.1, 6.1 Hz, 1H), 4.34 (t, <i>J</i> = 5.9 Hz, 2H), 3.62 (dd, <i>J</i> = 16.6, 8.9 Hz, 1H), 3.54 - 3.47 (m, 2H), 3.31 - 3.25 (m, 1H), 3.20 (t, <i>J</i> = 5.9 Hz, 2H), 2.93 (s, 3H), 2.82 (br t, <i>J</i> = 6.2 Hz, 2H), 2.01 - 1.90 (m, 2H). LC/MS (<i>m/z</i>) = 474.2 (<i>M</i> +H) ⁺ . Human αVβ6 IC ₅₀ (nM) = 21; Human αVβ1 IC ₅₀ (nM) = 93; Human αVβ3 IC ₅₀ (nM) = 2.9; and Human αVβ8 IC ₅₀ (nM) = 5,000.	Example 3

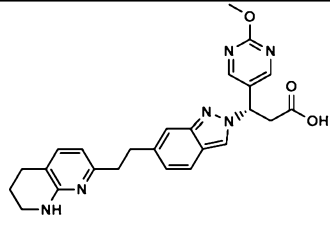
Example No.	Structure & Name	Analytical Data	Method
113	 <p>3-(6-Methoxypyridin-3-yl)-3-(5-(3-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)propyl)-1H-pyrazolo[4,3-b]pyridin-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, DMSO- <i>d</i> ₆) δ 8.27 (d, <i>J</i> = 2.5 Hz, 1H), 8.21 (d, <i>J</i> = 10.4 Hz, 2H), 7.71 (dd, <i>J</i> = 8.7, 2.4 Hz, 1H), 7.28 (d, <i>J</i> = 8.7 Hz, 1H), 7.02 (d, <i>J</i> = 7.1 Hz, 1H), 6.74 (d, <i>J</i> = 8.6 Hz, 1H), 6.30 – 6.19 (m, 2H), 3.78 (s, 3H), 3.54 (s, 1H), 3.19 (d, <i>J</i> = 23.9 Hz, 3H), 2.81 (t, <i>J</i> = 7.6 Hz, 2H), 2.58 (t, <i>J</i> = 6.3 Hz, 2H), 2.45 (t, <i>J</i> = 7.6 Hz, 2H), 1.98 (t, <i>J</i> = 7.5 Hz, 2H), 1.77 – 1.68 (m, 2H). Human αVβ6 IC ₅₀ (nM) = 890.	Example 1

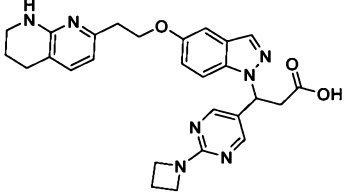
Example No.	Structure & Name	Analytical Data	Method
114	 <p>3-(6-Methoxypyridazin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid, TFA</p>	¹ H NMR (500 MHz, chloroform- <i>d</i>) δ 7.98 (s, 1H), 7.44 (br d, <i>J</i> = 9.1 Hz, 1H), 7.37 (d, <i>J</i> = 7.4 Hz, 1H), 7.30 (d, <i>J</i> = 9.1 Hz, 1H), 7.12 - 7.05 (m, 1H), 7.01 (d, <i>J</i> = 9.1 Hz, 2H), 6.60 - 6.47 (m, 2H), 4.28 (br t, <i>J</i> = 5.8 Hz, 2H), 4.12 (s, 3H), 3.75 - 3.57 (m, 2H), 3.51 (br t, <i>J</i> = 5.4 Hz, 2H), 3.17 (br t, <i>J</i> = 5.6 Hz, 2H), 2.77 (br t, <i>J</i> = 6.1 Hz, 2H), 1.99 - 1.90 (m, 2H). LC/MS (<i>m/z</i>) = 475.2 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 2.6.	Example 3

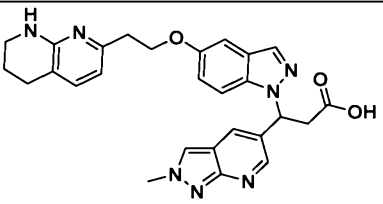
Example No.	Structure & Name	Analytical Data	Method
115	 <p>3-([1,2,4]Triazolo[4,3-<i>a</i>]pyridin-6-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1<i>H</i>-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, DMSO- <i>d</i> ₆) δ 9.23 (s, 1H), 8.66 (s, 1H), 8.03 (s, 1H), 7.93 (s, 1H), 7.73 (d, <i>J</i> = 9.1 Hz, 1H), 7.69 (d, <i>J</i> = 9.7 Hz, 1H), 7.59 (d, <i>J</i> = 7.4 Hz, 1H), 7.39 – 7.32 (m, 1H), 7.29 (s, 1H), 7.23 – 7.16 (m, 2H), 7.09 (s, 1H), 7.01 (dd, <i>J</i> = 9.1, 2.3 Hz, 1H), 6.70 (d, <i>J</i> = 7.4 Hz, 1H), 6.24 (dd, <i>J</i> = 10.1, 4.7 Hz, 1H), 4.26 (q, <i>J</i> = 5.6 Hz, 2H), 3.38 (t, <i>J</i> = 5.6 Hz, 2H), 3.26 (dd, <i>J</i> = 16.7, 4.7 Hz, 1H), 3.11 (t, <i>J</i> = 6.1 Hz, 2H), 2.70 (d, <i>J</i> = 6.4 Hz, 2H), 1.79 (t, <i>J</i> = 5.8 Hz, 2H). LC/MS (<i>m/z</i>) = 484.4 (<i>M</i> + <i>H</i>) ⁺ . Human αVβ6 IC ₅₀ (nM) = 60.	Example 3

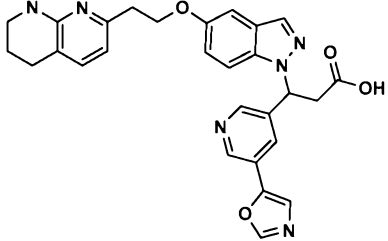
Example No.	Structure & Name	Analytical Data	Method
116	 <p>3-(5,6,7,8-Tetrahydro-1,8-naphthyridin-2-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid, 2 TFA</p>	¹ H NMR (500 MHz, MeOH- <i>d</i> ₄) δ 8.04 (s, 1H), 7.68 – 7.56 (m, 2H), 7.51 (d, <i>J</i> = 7.4 Hz, 1H), 7.23 (d, <i>J</i> = 2.2 Hz, 1H), 7.10 (dd, <i>J</i> = 9.1, 2.5 Hz, 1H), 6.77 (d, <i>J</i> = 7.2 Hz, 1H), 6.62 (d, <i>J</i> = 7.2 Hz, 1H), 6.22 (dd, <i>J</i> = 8.9, 5.6 Hz, 1H), 4.34 (br t, <i>J</i> = 5.9 Hz, 2H), 3.62 (dd, <i>J</i> = 17.1, 9.1 Hz, 1H), 3.55 – 3.46 (m, 4H), 3.41 (br dd, <i>J</i> = 16.9, 5.6 Hz, 1H), 3.25 – 3.18 (m, 2H), 2.84 (br t, <i>J</i> = 6.1 Hz, 2H), 2.80 (br t, <i>J</i> = 6.1 Hz, 2H), 2.00 – 1.88 (m, 4H). LC/MS (<i>m/z</i>) = 499.2 (<i>M</i> + <i>H</i>) ⁺ . Human α V β 6 IC ₅₀ (nM) = 5.9.	Example 3

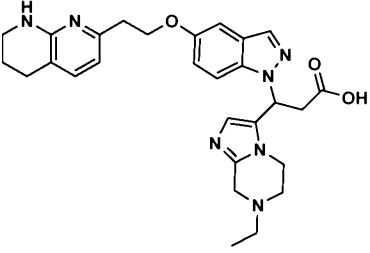
Example No.	Structure & Name	Analytical Data	Method
117	 <p>3-(1-Methyl-1H-pyrazolo[3,4-b]pyridin-5-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid, 2 TFA</p>	¹ H NMR (500 MHz, MeOH-d ₄) δ 8.61 (d, <i>J</i> = 1.9 Hz, 1H), 8.23 (d, <i>J</i> = 2.2 Hz, 1H), 8.04 (s, 1H), 8.00 (s, 1H), 7.64 – 7.58 (m, 2H), 7.19 (d, <i>J</i> = 2.2 Hz, 1H), 7.05 (dd, <i>J</i> = 9.1, 2.2 Hz, 1H), 6.76 (d, <i>J</i> = 7.4 Hz, 1H), 6.41 (dd, <i>J</i> = 9.2, 5.6 Hz, 1H), 4.38 – 4.28 (m, 2H), 4.08 (s, 3H), 3.80 (dd, <i>J</i> = 16.8, 9.4 Hz, 1H), 3.52 – 3.48 (m, 2H), 3.44 – 3.38 (m, 1H), 3.21 – 3.16 (m, 2H), 2.81 (br t, <i>J</i> = 6.1 Hz, 2H), 1.97 – 1.91 (m, 2H). LC/MS (<i>m/z</i>) = 498.2 (M+H) ⁺ . Human αVβ6 IC ₅₀ (nM) = 19; Human αVβ1 IC ₅₀ (nM) = 63; Human αVβ3 IC ₅₀ (nM) = 1.6; Human αVβ5 IC ₅₀ (nM) = 0.46; and Human αVβ8 IC ₅₀ (nM) = 2,200.	Example 3

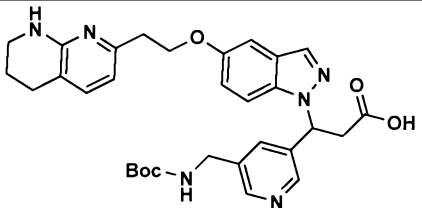
Example No.	Structure & Name	Analytical Data	Method
118	 <p>(S)-3-(2-Methoxypyrimidin-5-yl)-3-(6-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethyl)-2H-indazol-2-yl)propanoic acid</p>	¹ H NMR (500 MHz, DMSO- <i>d</i> ₆) δ 8.74 (s, 2H), 8.50 (s, 1H), 7.95 (s, 1H), 7.61 (d, <i>J</i> = 8.6 Hz, 1H), 7.40 (s, 1H), 7.37 (d, <i>J</i> = 7.3 Hz, 1H), 6.95 (d, <i>J</i> = 8.6 Hz, 1H), 6.50 (d, <i>J</i> = 7.3 Hz, 1H), 6.18 (dd, <i>J</i> = 9.2, 5.7 Hz, 1H), 3.88 (s, 3H), 3.64 (dd, <i>J</i> = 16.9, 9.4 Hz, 1H), 3.40 – 3.31 (m, 1H), 3.02 – 2.95 (m, 2H), 2.92 (d, <i>J</i> = 7.8 Hz, 2H), 2.67 (t, <i>J</i> = 6.2 Hz, 2H), 1.83 – 1.73 (m, 2H). LC/MS (<i>m/z</i>) = 459.0 (M+H) ⁺ . Human αVβ6 IC ₅₀ (nM) = 12; Human αVβ1 IC ₅₀ (nM) = 4,300; Human αVβ3 IC ₅₀ (nM) = 2.0; Human αVβ5 IC ₅₀ (nM) = 1.1; and Human αVβ8 IC ₅₀ (nM) = 3,100.	Example 7

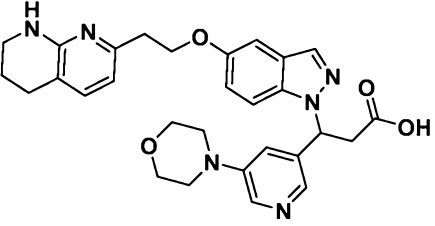
Example No.	Structure & Name	Analytical Data	Method
119	 <p>3-(2-(Azetidin-1-yl)pyrimidin-5-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid, 2 TFA</p>	¹ H NMR (500 MHz, chloroform- <i>d</i>) δ 10.05 (br s, 1H), 8.60 (s, 2H), 7.94 (s, 1H), 7.38 - 7.29 (m, 2H), 7.07 (d, <i>J</i> = 2.2 Hz, 1H), 6.85 (d, <i>J</i> = 8.6 Hz, 1H), 6.48 (d, <i>J</i> = 7.2 Hz, 1H), 6.11 (br t, <i>J</i> = 7.2 Hz, 1H), 4.32 - 4.19 (m, 6H), 3.81 - 3.59 (m, 1H), 3.47 (br t, <i>J</i> = 5.2 Hz, 2H), 3.26 (br dd, <i>J</i> = 16.5, 6.3 Hz, 1H), 3.16 (br t, <i>J</i> = 5.9 Hz, 2H), 2.74 (br t, <i>J</i> = 6.1 Hz, 2H), 2.42 (br t, <i>J</i> = 7.7 Hz, 2H), 1.98 - 1.88 (m, 2H). LC/MS (<i>m/z</i>) = 500.2 (<i>M</i> +H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 10.55; Human α V β 1 IC ₅₀ (nM) = 98; Human α V β 3 IC ₅₀ (nM) = 1.4; Human α V β 5 IC ₅₀ (nM) = 0.60; and Human α V β 8 IC ₅₀ (nM) = 4,400.	Example 3

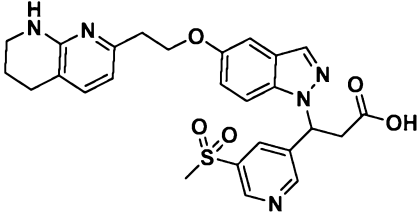
Example No.	Structure & Name	Analytical Data	Method
120	 <p>3-(2-Methyl-2H-pyrazolo[3,4-b]pyridin-5-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid, 2 TFA</p>	<p>¹H NMR (500 MHz, MeOH-<i>d</i>₄) δ 8.63 (br s, 1H), 8.33 - 8.25 (m, 2H), 8.00 (s, 1H), 7.66 - 7.58 (m, 2H), 7.19 (d, <i>J</i> = 2.2 Hz, 1H), 7.06 (dd, <i>J</i> = 9.2, 2.3 Hz, 1H), 6.76 (d, <i>J</i> = 7.2 Hz, 1H), 6.38 (dd, <i>J</i> = 9.4, 5.5 Hz, 1H), 4.33 (br t, <i>J</i> = 5.8 Hz, 2H), 4.24 (s, 3H), 3.77 (dd, <i>J</i> = 16.6, 9.2 Hz, 1H), 3.52 - 3.46 (m, 2H), 3.41 (br dd, <i>J</i> = 16.6, 5.6 Hz, 1H), 3.18 (t, <i>J</i> = 5.9 Hz, 2H), 2.81 (br t, <i>J</i> = 6.1 Hz, 2H), 1.99 - 1.90 (m, 2H).</p> <p>LC/MS (<i>m/z</i>) = 498.2</p> <p>(<i>M</i>+<i>H</i>)⁺. Human αVβ6 IC₅₀ (nM) = 16; Human αVβ1 IC₅₀ (nM) = 200; Human αVβ3 IC₅₀ (nM) = 1.5; Human αVβ5 IC₅₀ (nM) = 0.20; and Human αVβ8 IC₅₀ (nM) = 7,500.</p>	Example 3

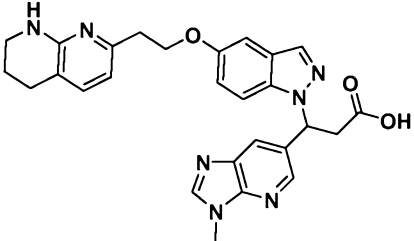
Example No.	Structure & Name	Analytical Data	Method
121	 <p data-bbox="416 678 871 846">3-(5-(Oxazol-5-yl)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid</p>	<p data-bbox="898 421 1217 1301">¹H NMR (500 MHz, DMSO-<i>d</i>₆) δ 8.86 (d, <i>J</i> = 2.1 Hz, 1H), 8.58 (d, <i>J</i> = 2.1 Hz, 1H), 8.51 (s, 1H), 8.08 (t, <i>J</i> = 2.2 Hz, 1H), 8.04 (s, 1H), 7.82 (s, 1H), 7.75 (d, <i>J</i> = 9.2 Hz, 1H), 7.19 (d, <i>J</i> = 2.3 Hz, 1H), 7.12 (d, <i>J</i> = 7.2 Hz, 1H), 7.00 (dd, <i>J</i> = 9.1, 2.4 Hz, 1H), 6.41 (d, <i>J</i> = 7.3 Hz, 1H), 6.33 (dd, <i>J</i> = 10.0, 5.1 Hz, 1H), 4.29 – 4.16 (m, 2H), 3.66 (dd, <i>J</i> = 16.6, 9.9 Hz, 1H), 3.33 (dd, <i>J</i> = 16.7, 5.1 Hz, 1H), 3.25 (s, 2H), 2.92 (t, <i>J</i> = 6.7 Hz, 2H), 2.62 (t, <i>J</i> = 6.2 Hz, 2H), 1.81 – 1.68 (m, 2H).</p> <p data-bbox="898 1323 1217 1682">LC/MS (<i>m/z</i>) = 511.1 (M+H)⁺. Human αVβ6 IC₅₀ (nM) = 8.8; Human αVβ1 IC₅₀ (nM) = 71; Human αVβ3 IC₅₀ (nM) = 3.1; Human αVβ5 IC₅₀ (nM) = 0.46; and Human αVβ8 IC₅₀ (nM) = 2,600.</p>	Example 3

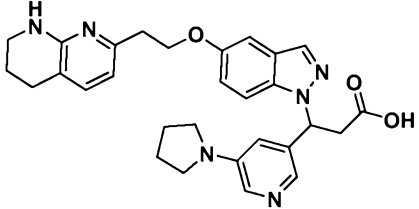
Example No.	Structure & Name	Analytical Data	Method
122	 <p data-bbox="411 689 871 913">3-(7-Ethyl-5,6,7,8-tetrahydroimidazo[1,2-a]pyrazin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, DMSO- <i>d</i> ₆) δ 7.96 (s, 1H), 7.68 (d, <i>J</i> = 9.3 Hz, 1H), 7.34 (d, <i>J</i> = 7.3 Hz, 1H), 7.23 – 7.13 (m, 2H), 7.02 (dd, <i>J</i> = 9.2, 2.4 Hz, 1H), 6.56 (d, <i>J</i> = 7.4 Hz, 1H), 6.29 (t, <i>J</i> = 7.3 Hz, 1H), 4.26 (t, <i>J</i> = 6.5 Hz, 2H), 4.02 (s, 1H), 3.33 (d, <i>J</i> = 7.0 Hz, 2H), 3.02 (t, <i>J</i> = 6.5 Hz, 2H), 2.96 – 2.61 (m, 4H), 1.94 (d, <i>J</i> = 23.9 Hz, 2H), 1.84 – 1.71 (m, 2H), 1.24 (s, 2H), 1.02 (t, <i>J</i> = 7.1 Hz, 4H). LC/MS (<i>m/z</i>) = 516.5 (M+H) ⁺ . Human αVβ6 IC ₅₀ (nM) = 14.	Example 3

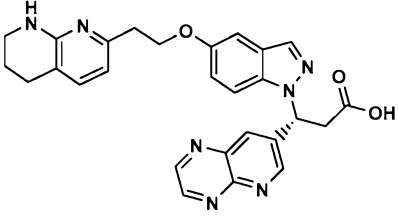
Example No.	Structure & Name	Analytical Data	Method
123	 <p>3-(5-(((<i>tert</i>-Butoxycarbonyl)amino)methyl)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1<i>H</i>-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, MeOH- <i>d</i> ₄) δ 8.83 - 8.65 (m, 1H), 8.65 - 8.52 (m, 1H), 8.42 - 8.16 (m, 1H), 8.03 (s, 1H), 7.62 (br d, <i>J</i> = 8.5 Hz, 2H), 7.21 (d, <i>J</i> = 1.9 Hz, 1H), 7.08 (br d, <i>J</i> = 9.1 Hz, 1H), 6.76 (br dd, <i>J</i> = 6.9, 2.8 Hz, 1H), 6.43 (br s, 1H), 4.33 (br d, <i>J</i> = 3.6 Hz, 4H), 3.72 (br dd, <i>J</i> = 16.9, 9.2 Hz, 1H), 3.55 - 3.48 (m, 2H), 3.48 - 3.37 (m, 1H), 3.24 - 3.16 (m, 2H), 2.83 (br t, <i>J</i> = 5.9 Hz, 2H), 1.96 (quin, <i>J</i> = 5.7 Hz, 2H), 1.43 (br s, 9H). LC/MS (<i>m/z</i>) = 573.3 (<i>M</i> + <i>H</i>) ⁺ . Human α V β 6 IC ₅₀ (nM) = 5.5.	Example 3

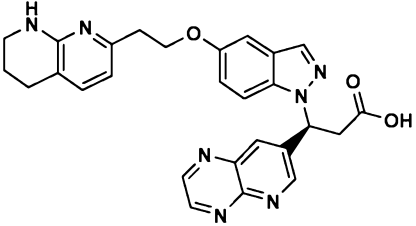
Example No.	Structure & Name	Analytical Data	Method
124	 <p>3-(5-Morpholinopyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid, 3 TFA</p>	¹ H NMR (500 MHz, MeOH- <i>d</i> ₄) δ 8.27 (br s, 1H), 8.13 (s, 1H), 8.11 - 8.02 (m, 2H), 7.68 - 7.56 (m, 2H), 7.22 (d, <i>J</i> = 1.9 Hz, 1H), 7.10 (dd, <i>J</i> = 9.1, 2.2 Hz, 1H), 6.76 (d, <i>J</i> = 7.1 Hz, 1H), 6.38 (br s, 1H), 4.38 - 4.28 (m, 2H), 3.85 (t, <i>J</i> = 5.0 Hz, 4H), 3.72 (br dd, <i>J</i> = 16.9, 9.2 Hz, 1H), 3.56 - 3.48 (m, 2H), 3.46 - 3.39 (m, 1H), 3.39 - 3.35 (m, 4H), 3.24 - 3.16 (m, 2H), 2.83 (br t, <i>J</i> = 6.1 Hz, 2H), 1.96 (quin, <i>J</i> = 5.8 Hz, 2H). LC/MS (<i>m/z</i>) = 529.2 (M+H) ⁺ . Human αVβ6 IC ₅₀ (nM) = 1.9.	Example 3

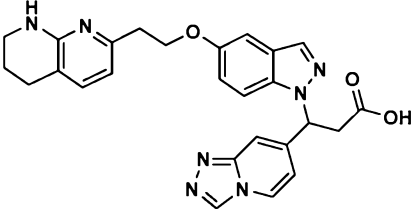
Example No.	Structure & Name	Analytical Data	Method
125	 <p>3-(5-(Methylsulfonyl)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid, 2 TFA</p>	¹ H NMR (500 MHz, MeOH- <i>d</i> ₄) δ 9.00 (br d, <i>J</i> = 2.2 Hz, 1H), 8.92 - 8.82 (m, 1H), 8.34 (br s, 1H), 8.05 - 8.00 (m, 1H), 7.65 - 7.53 (m, 2H), 7.23 - 7.17 (m, 1H), 7.08 (br dd, <i>J</i> = 9.1, 2.2 Hz, 1H), 6.79 - 6.68 (m, 1H), 6.43 (br dd, <i>J</i> = 9.2, 5.6 Hz, 1H), 4.38 - 4.28 (m, 2H), 3.74 (br dd, <i>J</i> = 16.6, 9.2 Hz, 1H), 3.53 - 3.47 (m, 2H), 3.42 (br dd, <i>J</i> = 16.8, 5.5 Hz, 1H), 3.25 - 3.12 (m, 5H), 2.81 (br d, <i>J</i> = 5.8 Hz, 2H), 2.01 - 1.86 (m, 2H). LC/MS (<i>m/z</i>) = 522.2 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 4.7.	Example 3

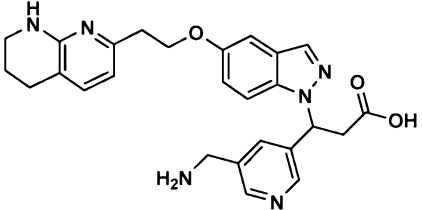
Example No.	Structure & Name	Analytical Data	Method
126	 <p data-bbox="419 683 871 853">3-(3-(3-Methyl-3H-imidazo[4,5-b]pyridin-6-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid, 2 TFA</p>	¹ H NMR (500 MHz, MeOH- <i>d</i> ₄) δ 9.07 - 8.78 (m, 1H), 8.68 (br s, 1H), 8.20 (br s, 1H), 8.01 (s, 1H), 7.64 - 7.55 (m, 2H), 7.22 - 7.15 (m, 1H), 7.05 (br d, <i>J</i> = 9.1 Hz, 1H), 6.77 - 6.70 (m, 1H), 6.47 (br dd, <i>J</i> = 8.4, 5.4 Hz, 1H), 4.36 - 4.27 (m, 2H), 4.04 - 3.96 (m, 3H), 3.80 (dd, <i>J</i> = 16.6, 9.2 Hz, 1H), 3.49 (br t, <i>J</i> = 5.2 Hz, 2H), 3.43 (br dd, <i>J</i> = 16.5, 5.8 Hz, 1H), 3.18 (br t, <i>J</i> = 5.8 Hz, 2H), 2.80 (br s, 2H), 2.01 - 1.85 (m, 2H). LC/MS (<i>m/z</i>) = 498.2 (<i>M</i> +H) ⁺ . Human αVβ6 IC ₅₀ (nM) = 21.	Example 3

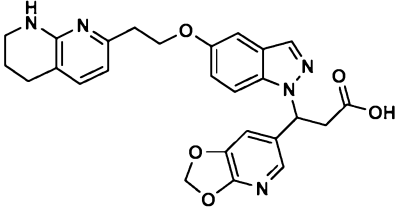
Example No.	Structure & Name	Analytical Data	Method
127	 <p data-bbox="416 645 871 824">3-(5-(Pyrrolidin-1-yl)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid, 3 TFA</p>	¹ H NMR (500 MHz, MeOH- <i>d</i> ₄) δ 8.05 (s, 1H), 7.94 (s, 1H), 7.89 (d, <i>J</i> = 2.2 Hz, 1H), 7.68 - 7.59 (m, 3H), 7.23 (d, <i>J</i> = 1.9 Hz, 1H), 7.10 (dd, <i>J</i> = 9.1, 2.2 Hz, 1H), 6.76 (d, <i>J</i> = 7.4 Hz, 1H), 6.38 (dd, <i>J</i> = 9.2, 5.6 Hz, 1H), 4.38 - 4.28 (m, 2H), 3.73 (dd, <i>J</i> = 16.9, 9.2 Hz, 1H), 3.56 - 3.48 (m, 2H), 3.46 - 3.39 (m, 1H), 3.39 - 3.35 (m, 4H), 3.24 - 3.16 (m, 2H), 2.84 (br t, <i>J</i> = 6.1 Hz, 2H), 2.10 (br t, <i>J</i> = 6.5 Hz, 4H), 1.96 (dt, <i>J</i> = 11.7, 6.0 Hz, 2H). LC/MS (<i>m/z</i>) = 513.2 (M+H) ⁺ . Human αVβ6 IC ₅₀ (nM) = 6.2.	Example 3

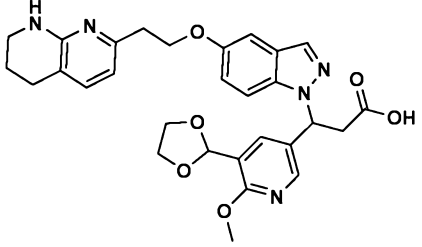
Example No.	Structure & Name	Analytical Data	Method
128	 <p data-bbox="421 651 865 824">(R)-3-(Pyrido[2,3-<i>b</i>]pyrazin-7-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1<i>H</i>-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, chloroform- <i>d</i>) δ 9.26 (br. s., 1H), 9.07 (d, <i>J</i> = 1.4 Hz, 1H), 8.98 (d, <i>J</i> = 1.7 Hz, 1H), 8.47 (d, <i>J</i> = 1.9 Hz, 1H), 8.04 (s, 1H), 7.42 - 7.31 (m, 2H), 7.09 (d, <i>J</i> = 1.9 Hz, 1H), 7.02 (dd, <i>J</i> = 9.1, 2.2 Hz, 1H), 6.51 (d, <i>J</i> = 7.2 Hz, 1H), 6.45 - 6.36 (m, 1H), 4.32 - 4.24 (m, 2H), 3.58 - 3.52 (m, 1H), 3.49 (t, <i>J</i> = 5.5 Hz, 2H), 3.16 (t, <i>J</i> = 5.8 Hz, 2H), 2.75 (t, <i>J</i> = 6.2 Hz, 2H), 1.97 - 1.87 (m, 2H). LC/MS (<i>m/z</i>) = 496.2 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 25.	Example 16 and 17

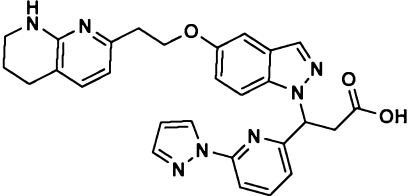
Example No.	Structure & Name	Analytical Data	Method
129	 <p data-bbox="421 667 869 837">(S)-3-(Pyrido[2,3-<i>b</i>]pyrazin-7-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1<i>H</i>-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, chloroform- <i>d</i>) δ 9.26 (br. s., 1H), 9.07 (d, <i>J</i> = 1.4 Hz, 1H), 8.98 (d, <i>J</i> = 1.7 Hz, 1H), 8.47 (d, <i>J</i> = 1.9 Hz, 1H), 8.04 (s, 1H), 7.42 - 7.31 (m, 2H), 7.09 (d, <i>J</i> = 1.9 Hz, 1H), 7.02 (dd, <i>J</i> = 9.1, 2.2 Hz, 1H), 6.51 (d, <i>J</i> = 7.2 Hz, 1H), 6.45 - 6.36 (m, 1H), 4.32 - 4.24 (m, 2H), 3.58 - 3.52 (m, 1H), 3.49 (t, <i>J</i> = 5.5 Hz, 2H), 3.16 (t, <i>J</i> = 5.8 Hz, 2H), 2.75 (t, <i>J</i> = 6.2 Hz, 2H), 1.97 - 1.87 (m, 2H). LC/MS (<i>m/z</i>) = 496.2 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 480.	Example 16 and 17

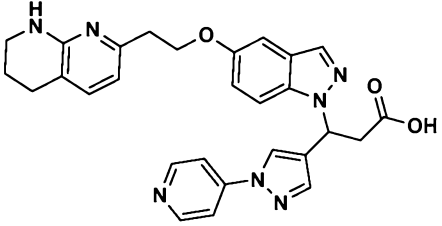
Example No.	Structure & Name	Analytical Data	Method
130	 <p>3-([1,2,4]Triazolo[4,3-a]pyridin-7-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, DMSO- <i>d</i> ₆) δ 9.17 (s, 1H), 8.44 (d, <i>J</i> = 7.3 Hz, 1H), 8.05 (s, 1H), 7.73 (d, <i>J</i> = 9.1 Hz, 1H), 7.68 (s, 1H), 7.44 (d, <i>J</i> = 7.2 Hz, 1H), 7.22 (d, <i>J</i> = 2.4 Hz, 1H), 7.01 (dd, <i>J</i> = 9.1, 2.4 Hz, 1H), 6.87 (d, <i>J</i> = 7.4 Hz, 1H), 6.61 (d, <i>J</i> = 7.2 Hz, 1H), 6.28 (d, <i>J</i> = 8.1 Hz, 1H), 4.26 (q, <i>J</i> = 5.6 Hz, 2H), 3.63 (dd, <i>J</i> = 17.1, 9.8 Hz, 1H), 3.40 – 3.24 (m, 2H), 3.05 (t, <i>J</i> = 6.5 Hz, 2H), 2.93 (s, 1H), 2.68 (t, <i>J</i> = 6.2 Hz, 2H), 1.78 (d, <i>J</i> = 6.4 Hz, 3H), 1.17 (t, <i>J</i> = 7.3 Hz, 2H). LC/MS (<i>m/z</i>) = 484.1 (M+H) ⁺ . Human αVβ6 IC ₅₀ (nM) = 57.	Example 3

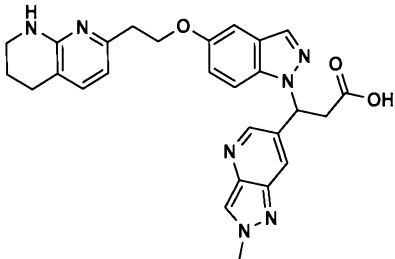
Example No.	Structure & Name	Analytical Data	Method
131	 <p>3-(5-(Aminomethyl)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, methanol- <i>d</i> ₄) δ 8.63 (s, 1H), 8.54 (s, 1H), 8.02 – 7.95 (m, 2H), 7.63 – 7.55 (m, 2H), 7.18 (d, <i>J</i> = 2.3 Hz, 1H), 7.06 (dd, <i>J</i> = 9.1, 2.3 Hz, 1H), 6.73 (d, <i>J</i> = 7.4 Hz, 1H), 6.32 (dd, <i>J</i> = 9.3, 5.6 Hz, 1H), 4.30 (tt, <i>J</i> = 6.2, 3.1 Hz, 2H), 4.14 (s, 2H), 3.70 (dd, <i>J</i> = 16.7, 9.3 Hz, 1H), 3.49 (t, <i>J</i> = 5.7 Hz, 2H), 3.38 – 3.32 (m, 1H), 3.17 (t, <i>J</i> = 5.9 Hz, 2H), 2.81 (t, <i>J</i> = 6.3 Hz, 2H), 1.97 – 1.89 (m, 2H). LC/MS (<i>m/z</i>) = 473.1 (M+H) ⁺ . Human αVβ6 IC ₅₀ (nM) = 30.	Deprotection of Boc- in Example 126 with TFA

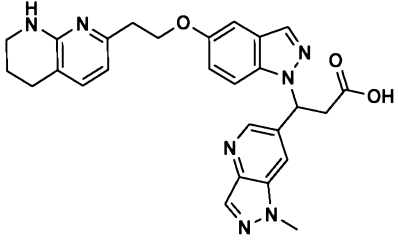
Example No.	Structure & Name	Analytical Data	Method
132	 <p data-bbox="411 645 871 819">3-([1,3]Dioxolo[4,5-<i>b</i>]pyridin-6-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1<i>H</i>-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, DMSO- <i>d</i> ₆) δ 8.01 (d, <i>J</i> = 14.3 Hz, 1H), 7.72 (d, <i>J</i> = 9.1 Hz, 1H), 7.68 (d, <i>J</i> = 1.9 Hz, 1H), 7.61 (d, <i>J</i> = 7.3 Hz, 1H), 7.30 – 7.13 (m, 3H), 7.09 – 6.96 (m, 2H), 6.73 (dd, <i>J</i> = 7.2, 3.3 Hz, 1H), 6.23 – 6.11 (m, 1H), 6.09 (s, 1H), 6.05 (s, 1H), 4.28 (d, <i>J</i> = 8.6 Hz, 3H), 3.61 (dd, <i>J</i> = 16.7, 10.0 Hz, 1H), 3.22 – 3.09 (m, 3H), 2.73 (s, 2H), 1.82 (s, 3H). LC/MS (<i>m/z</i>) = 487.9 (<i>M</i> + <i>H</i>) ⁺ . Human αVβ6 IC ₅₀ (nM) = 22.	Example 3

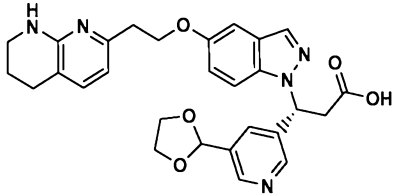
Example No.	Structure & Name	Analytical Data	Method
133	 <p data-bbox="427 683 863 902">3-(5-(1,3-Dioxolan-2-yl)-6-methoxypyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, methanol- <i>d</i> ₄) δ 7.99 (s, 1H), 7.88 (s, 1H), 7.61 (d, <i>J</i> = 7.4 Hz, 1H), 7.49 (d, <i>J</i> = 9.1 Hz, 1H), 7.19 (d, <i>J</i> = 2.4 Hz, 1H), 7.00 (dd, <i>J</i> = 9.1, 2.4 Hz, 1H), 6.93 (s, 1H), 6.76 (d, <i>J</i> = 7.3 Hz, 1H), 6.63 (dd, <i>J</i> = 10.9, 3.6 Hz, 1H), 5.93 (s, 1H), 4.33 (q, <i>J</i> = 5.0, 4.0 Hz, 3H), 4.27 – 4.20 (m, 1H), 4.20 – 4.09 (m, 2H), 3.85 (s, 3H), 3.72 (dd, <i>J</i> = 16.8, 10.8 Hz, 1H), 3.50 (t, <i>J</i> = 5.7 Hz, 2H), 3.27 – 3.14 (m, 3H), 2.81 (t, <i>J</i> = 6.3 Hz, 2H), 1.94 (p, <i>J</i> = 5.8 Hz, 2H). LC/MS (<i>m/z</i>) = 546.1 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 400.	Example 3

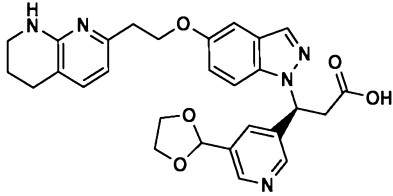
Example No.	Structure & Name	Analytical Data	Method
134	 <p>3-(6-(1H-Pyrazol-1-yl)pyridin-2-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, DMSO- <i>d</i> ₆) δ 8.59 (d, <i>J</i> = 2.4 Hz, 1H), 8.03 (s, 1H), 7.88 – 7.75 (m, 3H), 7.66 (d, <i>J</i> = 9.1 Hz, 1H), 7.51 (d, <i>J</i> = 7.4 Hz, 1H), 7.23 (d, <i>J</i> = 2.3 Hz, 1H), 7.03 (dd, <i>J</i> = 9.0, 2.3 Hz, 1H), 6.77 (d, <i>J</i> = 7.6 Hz, 1H), 6.66 (d, <i>J</i> = 7.4 Hz, 1H), 6.59 (t, <i>J</i> = 2.1 Hz, 1H), 6.33 – 6.24 (m, 1H), 4.27 (t, <i>J</i> = 6.1 Hz, 2H), 3.47 (dd, <i>J</i> = 17.3, 8.8 Hz, 1H), 3.37 (t, <i>J</i> = 5.6 Hz, 2H), 3.09 (t, <i>J</i> = 6.2 Hz, 2H), 2.70 (t, <i>J</i> = 6.2 Hz, 2H), 1.79 (s, 2H), 1.22 (s, 2H). LC/MS (<i>m/z</i>) = 510.3 (M+H) ⁺ . Human αVβ6 IC ₅₀ (nM) = 17.	Example 3

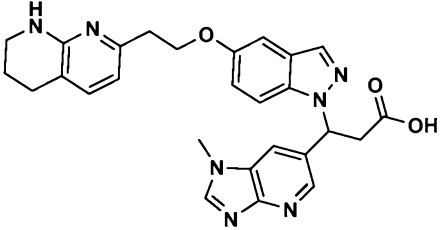
Example No.	Structure & Name	Analytical Data	Method
135	 <p>3-(1-(Pyridin-4-yl)-1H-pyrazol-4-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, DMSO- <i>d</i> ₆) δ 8.76 (s, 1H), 8.68 (d, <i>J</i> = 5.6 Hz, 2H), 7.99 (s, 1H), 7.91 (d, <i>J</i> = 5.6 Hz, 2H), 7.74 (d, <i>J</i> = 9.1 Hz, 1H), 7.61 (d, <i>J</i> = 7.4 Hz, 1H), 7.29 (s, 1H), 7.23 – 7.13 (m, 2H), 7.12 – 7.00 (m, 2H), 6.72 (d, <i>J</i> = 7.3 Hz, 1H), 6.28 – 6.16 (m, 1H), 4.27 (s, 2H), 3.39 (t, <i>J</i> = 5.6 Hz, 2H), 3.28 (dd, <i>J</i> = 16.9, 5.0 Hz, 1H), 3.12 (t, <i>J</i> = 6.2 Hz, 2H), 2.72 (t, <i>J</i> = 5.9 Hz, 2H), 1.80 (s, 2H), 1.22 (s, 2H). LC/MS (<i>m/z</i>) = 510.4 (M+H) ⁺ . Human αVβ6 IC ₅₀ (nM) = 3.8.	Example 3

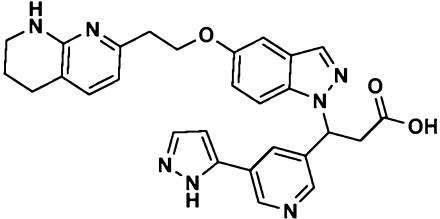
Example No.	Structure & Name	Analytical Data	Method
136	 <p data-bbox="416 913 871 1088">3-(2-Methyl-2H-pyrazolo[4,3-b]pyridin-6-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, DMSO-d ₆) δ 8.60 - 8.48 (m, 2H), 8.00 (br d, <i>J</i> =17.9 Hz, 2H), 7.75 - 7.66 (m, 1H), 7.23 - 7.15 (m, 1H), 7.14 - 7.06 (m, 1H), 7.03 - 6.96 (m, 1H), 6.45 - 6.38 (m, 1H), 6.38 - 6.31 (m, 1H), 4.31 - 4.23 (m, 2H), 4.22 - 4.13 (m, 3H), 3.74 - 3.58 (m, 2H), 2.97 - 2.86 (m, 2H), 2.66 - 2.60 (m, 2H), 2.29 (br t, <i>J</i> =7.4 Hz, 1H), 1.83 - 1.72 (m, 2H), 1.60 - 1.48 (m, 1H), 0.87 (s, 1H) LC/MS (<i>m/z</i>) = 497.9 (M+H) ⁺ . Human αVβ6 IC ₅₀ (nM) = 41.	Example 3

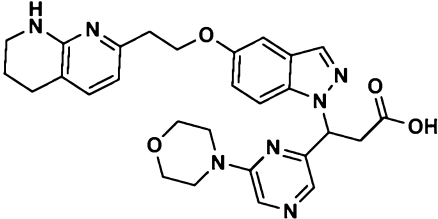
Example No.	Structure & Name	Analytical Data	Method
137	 <p>3-(1-Methyl-1H-pyrazolo[4,3-b]pyridin-6-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, DMSO-d ₆) δ 8.60 - 8.48 (m, 1H), 8.21 - 8.15 (m, 2H), 8.03 (s, 1H), 7.74 (d, <i>J</i> =9.1 Hz, 1H), 7.62 - 7.53 (m, 1H), 7.26 - 7.18 (m, 1H), 7.07 - 6.98 (m, 1H), 6.73 - 6.66 (m, 1H), 6.44 - 6.35 (m, 1H), 4.35 - 4.21 (m, 2H), 4.08 - 3.97 (m, 3H), 3.79 - 3.68 (m, 1H), 3.44 - 3.37 (m, 1H), 3.17 - 3.07 (m, 1H), 2.77 - 2.68 (m, 2H), 1.87 - 1.76 (m, 2H). LC/MS (<i>m/z</i>) = 497.9 (<i>M</i> +H) ⁺ . Human αVβ6 IC ₅₀ (nM) = 7.0.	Example 3

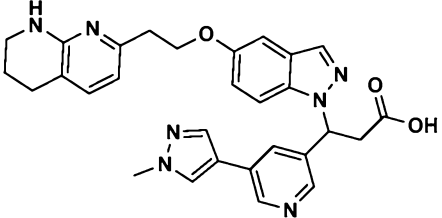
Example No.	Structure & Name	Analytical Data	Method
138	 <p data-bbox="427 857 871 1032">(R)-3-(5-(1,3-dioxolan-2-yl)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid</p>	¹ H NMR (400 MHz, methanol- <i>d</i> ₄) δ 8.54 (d, <i>J</i> = 8.6 Hz, 2H), 7.98 (s, 1H), 7.95 (s, 1H), 7.64 – 7.51 (m, 2H), 7.18 (d, <i>J</i> = 2.3 Hz, 1H), 7.05 (dd, <i>J</i> = 9.1, 2.4 Hz, 1H), 6.74 (d, <i>J</i> = 7.4 Hz, 1H), 6.31 (dd, <i>J</i> = 9.4, 5.3 Hz, 1H), 5.79 (s, 1H), 4.31 (t, <i>J</i> = 5.9 Hz, 2H), 4.10 – 3.95 (m, 4H), 3.77 – 3.62 (m, 1H), 3.48 (dd, <i>J</i> = 6.5, 4.8 Hz, 2H), 3.17 (t, <i>J</i> = 5.9 Hz, 2H), 2.79 (t, <i>J</i> = 6.2 Hz, 2H), 1.97 – 1.87 (m, 3H). LC/MS (<i>m/z</i>) = 516.2 (<i>M</i> + <i>H</i>) ⁺ . Human α V β 6 IC ₅₀ (nM) = 43.	Examples 104, 16, and 17.

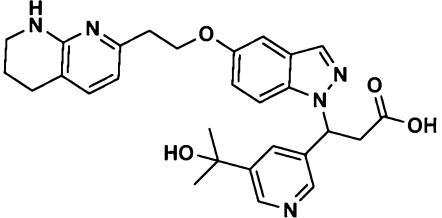
Example No.	Structure & Name	Analytical Data	Method
139	 <p>(S)-3-(5-(1,3-dioxolan-2-yl)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid</p>	¹ H NMR (400 MHz, methanol- <i>d</i> ₄) δ 8.55 (s, 3H), 7.99 (s, 1H), 7.63 – 7.53 (m, 2H), 7.18 (s, 1H), 7.05 (d, <i>J</i> = 8.4 Hz, 1H), 6.75 (d, <i>J</i> = 7.5 Hz, 1H), 6.32 (s, 1H), 5.80 (s, 1H), 4.31 (t, <i>J</i> = 5.9 Hz, 2H), 4.12 – 3.94 (m, 4H), 3.71 (dd, <i>J</i> = 16.8, 9.4 Hz, 1H), 3.48 (td, <i>J</i> = 4.9, 3.2 Hz, 3H), 3.17 (t, <i>J</i> = 5.9 Hz, 2H), 3.13 (p, <i>J</i> = 1.7 Hz, 1H), 2.80 (t, <i>J</i> = 6.2 Hz, 3H), 1.97 – 1.88 (m, 2H). LC/MS (<i>m/z</i>) = 516.1 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 4.7.	Example 104, 16, and 17.

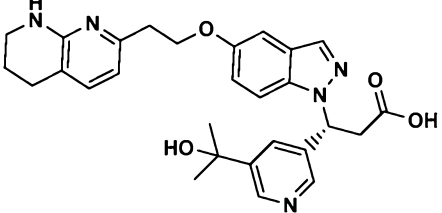
Example No.	Structure & Name	Analytical Data	Method
140	 <p data-bbox="422 667 865 840">3-(1-methyl-1H-imidazo[4,5-b]pyridin-6-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid</p>	<p data-bbox="895 421 1222 1120">¹H NMR (500 MHz, methanol-<i>d</i>₄) δ 8.58 - 8.48 (m, 2H), 8.21 (s, 1H), 8.01 (s, 1H), 7.68 - 7.59 (m, 2H), 7.20 (d, J=2.2 Hz, 1H), 7.06 (dd, J=9.1, 2.2 Hz, 1H), 6.76 (d, J=7.4 Hz, 1H), 6.46 (dd, J=9.4, 5.5 Hz, 1H), 4.38 - 4.29 (m, 2H), 3.95 (s, 3H), 3.83 (dd, J=16.5, 9.4 Hz, 1H), 3.52 - 3.47 (m, 2H), 3.43 (br dd, J=16.9, 5.6 Hz, 1H), 3.21 - 3.15 (m, 2H), 2.81 (br t, J=6.1 Hz, 2H), 1.99 - 1.90 (m, 2H).</p> <p data-bbox="895 1131 1222 1254">LC/MS (m/z) = 498.3 (M+H)⁺. Human αVβ6 IC₅₀ (nM) = 5.7.</p>	Example 3

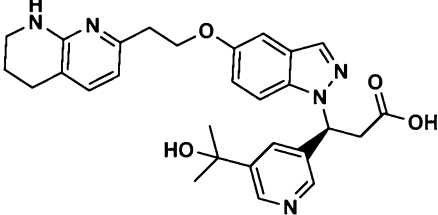
Example No.	Structure & Name	Analytical Data	Method
141	 <p data-bbox="422 660 863 840">3-(5-(1H-pyrazol-5-yl)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, DMSO- <i>d</i> ₆) δ 8.91 - 8.82 (m, 1H), 8.56 - 8.49 (m, 1H), 8.17 - 8.09 (m, 1H), 8.07 - 8.00 (m, 1H), 7.84 - 7.70 (m, 2H), 7.51 - 7.42 (m, 1H), 7.24 - 7.18 (m, 1H), 7.04 - 6.97 (m, 1H), 6.82 - 6.75 (m, 1H), 6.68 - 6.59 (m, 1H), 6.35 - 6.26 (m, 1H), 4.35 - 4.18 (m, 2H), 3.76 - 3.51 (m, 2H), 3.40 - 3.29 (m, 2H), 3.06 (br t, <i>J</i> =5.8 Hz, 2H), 2.73 - 2.64 (m, 2H), 1.84 - 1.73 (m, 2H). LC/MS (<i>m/z</i>) = 510.3 (<i>M</i> + <i>H</i>) ⁺ . Human αVβ6 IC ₅₀ (nM) = 8.2.	Example 3

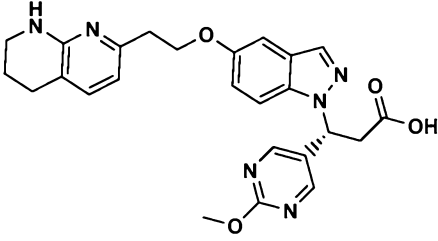
Example No.	Structure & Name	Analytical Data	Method
142	 <p>3-(6-morpholinopyrazin-2-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, DMSO- <i>d</i> ₆) δ 8.19 - 8.11 (m, 1H), 8.02 - 7.94 (m, 1H), 7.67 - 7.60 (m, 1H), 7.43 - 7.33 (m, 1H), 7.23 - 7.18 (m, 1H), 7.18 - 7.10 (m, 1H), 7.04 - 6.96 (m, 1H), 6.48 - 6.40 (m, 1H), 6.17 - 6.10 (m, 1H), 4.30 - 4.20 (m, 2H), 3.73 - 3.62 (m, 3H), 3.52 - 3.46 (m, 1H), 3.31 - 3.22 (m, 1H), 2.97 - 2.88 (m, 2H), 2.67 - 2.58 (m, 2H), 2.56 - 2.52 (m, 2H), 2.50 (br s, 5H), 1.82 - 1.70 (m, 2H). LC/MS (m/z) = 530.3 (M+H) ⁺ . Human αVβ6 IC ₅₀ (nM) = 36.	Example 3

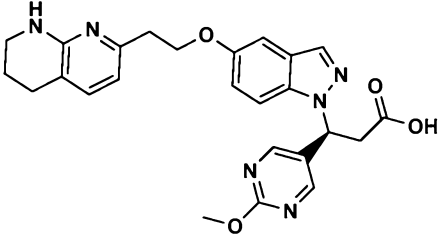
Example No.	Structure & Name	Analytical Data	Method
143	 <p data-bbox="414 660 873 840">3-(5-(1-methyl-1H-pyrazol-4-yl)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, DMSO- <i>d</i> ₆) δ 8.76 - 8.62 (m, 1H), 8.39 - 8.29 (m, 1H), 8.24 - 8.17 (m, 1H), 8.07 - 8.00 (m, 2H), 7.93 - 7.88 (m, 1H), 7.78 - 7.70 (m, 1H), 7.62 - 7.55 (m, 1H), 7.35 - 7.26 (m, 1H), 7.26 - 7.16 (m, 1H), 7.14 - 7.06 (m, 1H), 7.06 - 6.97 (m, 1H), 6.74 - 6.66 (m, 1H), 6.29 - 6.17 (m, 1H), 4.26 (br d, <i>J</i> =3.7 Hz, 2H), 3.86 (s, 3H), 3.73 - 3.52 (m, 1H), 3.43 - 3.36 (m, 2H), 3.35 - 3.27 (m, 1H), 3.15 - 3.07 (m, 2H), 2.75 - 2.67 (m, 2H), 1.86 - 1.73 (m, 2H). LC/MS (<i>m/z</i>) = 524.3 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 9.8.	Example 3

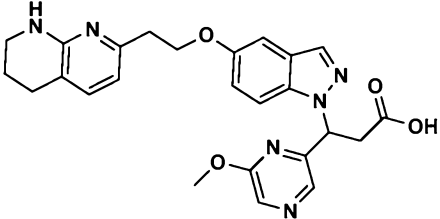
Example No.	Structure & Name	Analytical Data	Method
144	 <p>3-(5-(2-hydroxypropan-2-yl)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, DMSO- <i>d</i> ₆) δ 8.61 - 8.49 (m, 1H), 8.48 - 8.37 (m, 1H), 8.09 - 7.99 (m, 1H), 7.99 - 7.90 (m, 1H), 7.75 - 7.66 (m, 1H), 7.64 - 7.56 (m, 1H), 7.36 - 7.28 (m, 1H), 7.25 - 7.17 (m, 2H), 7.15 - 7.07 (m, 1H), 7.06 - 6.97 (m, 1H), 6.75 - 6.65 (m, 1H), 6.31 - 6.19 (m, 1H), 4.25 (br d, <i>J</i> =2.4 Hz, 2H), 3.68 - 3.55 (m, 1H), 3.43 - 3.33 (m, 2H), 3.31 - 3.21 (m, 1H), 3.15 - 3.06 (m, 2H), 2.75 - 2.66 (m, 2H), 1.84 - 1.74 (m, 2H), 1.43 - 1.31 (m, 6H). LC/MS (<i>m/z</i>) = 502.1 (<i>M</i> +H) ⁺ . Human αVβ6 IC ₅₀ (nM) = 5.0.	Example 3

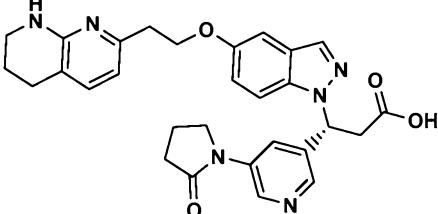
Example No.	Structure & Name	Analytical Data	Method
145	 <p>(R)-3-(5-(2-hydroxypropan-2-yl)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1H-indazol-1-yl)ethoxy)-1H-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, methanol- <i>d</i> ₄) δ 8.56 (d, <i>J</i> =2.0 Hz, 1H), 8.40 (d, <i>J</i> =2.0 Hz, 1H), 7.99 (s, 1H), 7.98 (s, 1H), 7.53 (d, <i>J</i> =9.0 Hz, 1H), 7.32 (d, <i>J</i> =7.3 Hz, 1H), 7.00 (s, 1H), 6.95 (d, <i>J</i> =8.9 Hz, 1H), 6.52 (d, <i>J</i> =7.3 Hz, 1H), 6.31 (dd, <i>J</i> =9.5, 5.5 Hz, 1H), 4.13 - 3.99 (m, 2H), 3.63 (dd, <i>J</i> =15.9, 9.5 Hz, 1H), 3.41 - 3.37 (m, 2H), 3.24 - 3.18 (m, 1H), 2.94 (t, <i>J</i> =6.3 Hz, 2H), 2.70 (t, <i>J</i> =6.2 Hz, 2H), 1.90 - 1.82 (m, 2H), 1.54 - 1.48 (m, 6H). LC/MS (<i>m/z</i>) = 502.3 (<i>M</i> + <i>H</i>) ⁺ . Human α V β 6 IC ₅₀ (nM) = 1,900.	Examples 144, 16, and 17

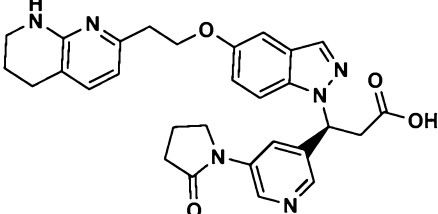
Example No.	Structure & Name	Analytical Data	Method
146	 <p data-bbox="414 660 869 840">(S)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, methanol- <i>d</i> ₄) δ 8.61 - 8.51 (m, 1H), 8.43 - 8.33 (m, 1H), 8.02 - 7.92 (m, 2H), 7.55 - 7.48 (m, 1H), 7.39 - 7.31 (m, 1H), 7.05 - 6.99 (m, 1H), 6.99 - 6.91 (m, 1H), 6.58 - 6.51 (m, 1H), 6.34 - 6.24 (m, 1H), 4.13 - 4.02 (m, 2H), 3.62 (dd, <i>J</i> =15.9, 9.5 Hz, 1H), 3.40 - 3.36 (m, 2H), 3.25 - 3.18 (m, 1H), 2.96 (t, <i>J</i> =6.1 Hz, 2H), 2.70 (t, <i>J</i> =6.2 Hz, 2H), 1.50 - 1.47 (m, 6H). LC/MS (<i>m/z</i>) = 502.3 (M+H) ⁺ . Human αVβ6 IC ₅₀ (nM) = 5.2.	Examples 144, 16, and 17

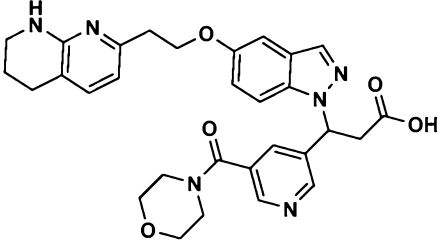
Example No.	Structure & Name	Analytical Data	Method
147	 <p>(R)-3-(2-methoxy pyrimidin-5-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, DMSO- <i>d</i> ₆) δ 8.71 - 8.47 (m, 2H), 8.09 - 7.90 (m, 1H), 7.84 - 7.72 (m, 1H), 7.65 - 7.54 (m, 1H), 7.25 - 7.17 (m, 1H), 7.07 - 6.99 (m, 1H), 6.80 - 6.60 (m, 1H), 6.36 - 6.16 (m, 1H), 4.28 (br s, 2H), 3.85 (s, 3H), 3.68 - 3.55 (m, 1H), 3.30 (dd, <i>J</i> =16.6, 5.0 Hz, 1H), 3.18 - 3.07 (m, 2H), 2.75 - 2.68 (m, 2H), 1.86 - 1.74 (m, 2H), 1.19 (s, 1H). LC/MS (<i>m/z</i>) = 475.2 (M+H) ⁺ . Human αVβ6 IC ₅₀ (nM) = 620.	Examples 92, 16, and 17

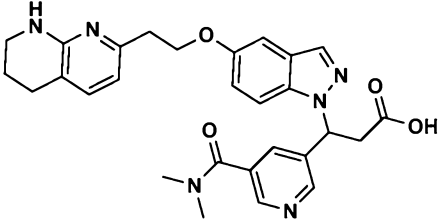
Example No.	Structure & Name	Analytical Data	Method
148	 <p data-bbox="422 674 865 846">(S)-3-(2-methoxypyrimidin-5-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, DMSO- <i>d</i> ₆) δ 8.70 - 8.54 (m, 2H), 8.08 - 7.92 (m, 1H), 7.85 - 7.71 (m, 1H), 7.69 - 7.53 (m, 1H), 7.25 - 7.15 (m, 1H), 7.12 - 6.93 (m, 1H), 6.79 - 6.62 (m, 1H), 6.30 - 6.17 (m, 1H), 4.34 - 4.18 (m, 2H), 3.85 (s, 3H), 3.66 - 3.53 (m, 1H), 3.35 - 3.22 (m, 1H), 3.19 - 3.07 (m, 2H), 2.80 - 2.66 (m, 2H), 1.88 - 1.74 (m, 2H), 1.30 - 1.14 (m, 1H). LC/MS (m/z) = 475.2 (M+H) ⁺ . Human αVβ6 IC ₅₀ (nM) = 18.	Examples 92, 16, and 17

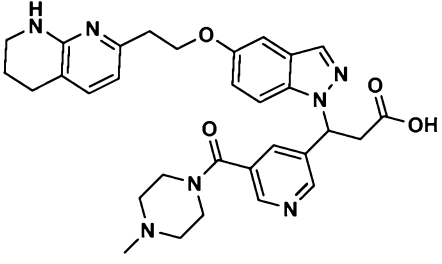
Example No.	Structure & Name	Analytical Data	Method
149	 <p>3-(6-methoxypyrazin-2-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, methanol- <i>d</i> ₄) δ 8.09 (s, 1H), 8.01 (s, 1H), 7.66 - 7.56 (m, 3H), 7.24 (s, 1H), 7.11 (dd, J=9.1, 1.4 Hz, 1H), 6.77 (br s, 1H), 6.34 (t, J=7.4 Hz, 1H), 4.36 (br t, J=5.8 Hz, 2H), 4.02 - 3.95 (m, 3H), 3.63 (d, J=6.1 Hz, 1H), 3.57 (br d, J=8.5 Hz, 1H), 3.53 - 3.46 (m, 2H), 3.21 (br t, J=5.8 Hz, 2H), 2.86 - 2.79 (m, 2H), 2.01 - 1.90 (m, 2H). LC/MS (m/z) = 475.1 (M+H) ⁺ . Human αVβ6 IC ₅₀ (nM) = 2,300.	Example 3

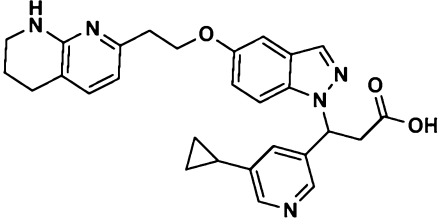
Example No.	Structure & Name	Analytical Data	Method
150	 <p data-bbox="414 660 869 840"><i>(R)</i>-3-(5-(2-oxopyrrolidin-1-yl)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, methanol- <i>d</i> ₄) δ 8.94 - 8.83 (m, 1H), 8.39 - 8.33 (m, 1H), 8.33 - 8.23 (m, 1H), 8.04 - 7.96 (m, 1H), 7.55 (br s, 2H), 7.24 - 7.15 (m, 1H), 7.12 - 7.00 (m, 1H), 6.70 (br s, 1H), 6.36 - 6.26 (m, 1H), 4.38 - 4.22 (m, 2H), 3.99 - 3.82 (m, 2H), 3.78 - 3.66 (m, 1H), 3.54 - 3.46 (m, 2H), 3.38 - 3.33 (m, 1H), 3.21 - 3.12 (m, 2H), 2.86 - 2.77 (m, 2H), 2.66 - 2.50 (m, 2H), 2.26 - 2.13 (m, 2H), 1.93 (dt, <i>J</i> =11.7, 6.0 Hz, 2H). LC/MS (<i>m/z</i>) = 527.2 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 110.	Examples 102, 16, and 17

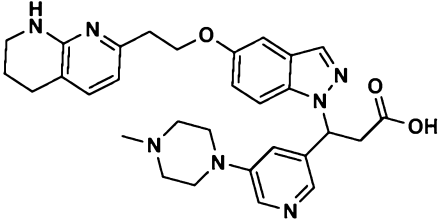
Example No.	Structure & Name	Analytical Data	Method
151	 <p data-bbox="414 660 869 840">(S)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)-3-(5-(2-oxopyrrolidin-1-yl)pyridin-3-yl)propanoic acid</p>	<p data-bbox="901 414 1212 1209">¹H NMR (500 MHz, methanol-<i>d</i>₄) δ 8.94 - 8.83 (m, 1H), 8.39 - 8.33 (m, 1H), 8.33 - 8.23 (m, 1H), 8.04 - 7.96 (m, 1H), 7.55 (br s, 2H), 7.24 - 7.15 (m, 1H), 7.12 - 7.00 (m, 1H), 6.70 (br s, 1H), 6.36 - 6.26 (m, 1H), 4.38 - 4.22 (m, 2H), 3.99 - 3.82 (m, 2H), 3.78 - 3.66 (m, 1H), 3.54 - 3.46 (m, 2H), 3.38 - 3.33 (m, 1H), 3.21 - 3.12 (m, 2H), 2.86 - 2.77 (m, 2H), 2.66 - 2.50 (m, 2H), 2.26 - 2.13 (m, 2H), 1.93 (dt, <i>J</i>=11.7, 6.0 Hz, 2H).</p> <p data-bbox="901 1220 1212 1355">LC/MS (<i>m/z</i>) = 527.2 (<i>M</i>+<i>H</i>)⁺. Human αVβ6 IC₅₀ (nM) = 1.</p>	Examples 102, 16, and 17

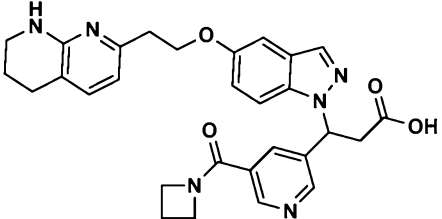
Example No.	Structure & Name	Analytical Data	Method
152	 <p data-bbox="422 683 863 851">3-(5-(morpholine-4-carbonyl)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, methanol- <i>d</i> ₄) δ 8.76 - 8.64 (m, 1H), 8.54 (s, 1H), 8.02 (s, 1H), 7.82 (t, <i>J</i> =1.8 Hz, 1H), 7.66 - 7.56 (m, 2H), 7.21 (d, <i>J</i> =1.9 Hz, 1H), 7.07 (dd, <i>J</i> =9.1, 2.2 Hz, 1H), 6.76 (d, <i>J</i> =7.2 Hz, 1H), 6.35 (dd, <i>J</i> =9.1, 5.8 Hz, 1H), 4.34 (td, <i>J</i> =5.8, 2.1 Hz, 2H), 3.73 (br dd, <i>J</i> =16.8, 9.1 Hz, 5H), 3.55 (br s, 2H), 3.53 - 3.47 (m, 2H), 3.46 - 3.36 (m, 1H), 3.20 (br t, <i>J</i> =5.9 Hz, 2H), 2.83 (br t, <i>J</i> =6.1 Hz, 2H), 1.95 (dt, <i>J</i> =11.6, 6.1 Hz, 2H). LC/MS (<i>m/z</i>) = 557.1 (<i>M</i> +H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 22.	Example 3

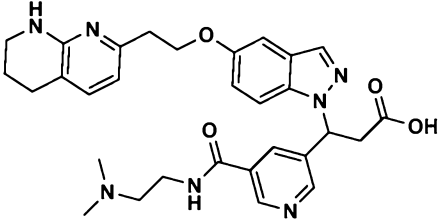
Example No.	Structure & Name	Analytical Data	Method
153	 <p>3-(5-(dimethylcarbamoyl)pyridin-3-yl)- 3-(5-(2-(5,6,7,8-tetrahydro-1,8- naphthyridin-2-yl)ethoxy)-1H-indazol-1- yl)propanoic acid</p>	¹ H NMR (500 MHz, methanol- <i>d</i> ₄) δ 8.67 (s, 1H), 8.55 (s, 1H), 8.01 (s, 1H), 7.93 (s, 1H), 7.60 (d, <i>J</i> =8.5 Hz, 2H), 7.20 (d, <i>J</i> =2.2 Hz, 1H), 7.07 (dd, <i>J</i> =9.1, 2.2 Hz, 1H), 6.75 (d, <i>J</i> =7.4 Hz, 1H), 6.35 (dd, <i>J</i> =9.1, 5.8 Hz, 1H), 4.33 (t, <i>J</i> =5.8 Hz, 2H), 3.73 (dd, <i>J</i> =16.6, 9.2 Hz, 1H), 3.53 - 3.47 (m, 2H), 3.45 - 3.35 (m, 1H), 3.19 (t, <i>J</i> =5.8 Hz, 2H), 3.10 (s, 3H), 2.93 (s, 3H), 2.81 (br t, <i>J</i> =6.1 Hz, 2H), 1.99 - 1.90 (m, 2H). LC/MS (<i>m/z</i>) = 515.1 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 17.	Example 3

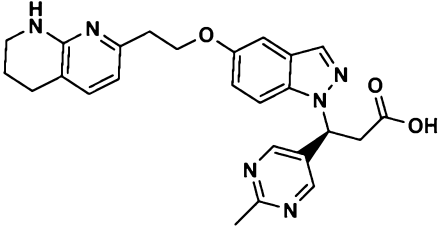
Example No.	Structure & Name	Analytical Data	Method
154	 <p data-bbox="422 694 865 913">3-(5-(4-methylpiperazine-1-carbonyl)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, methanol- <i>d</i> ₄) δ 8.73 (s, 1H), 8.61 (s, 1H), 8.01 (s, 2H), 7.64 - 7.57 (m, 2H), 7.20 (d, <i>J</i> =1.9 Hz, 1H), 7.08 (dd, <i>J</i> =9.1, 2.2 Hz, 1H), 6.74 (d, <i>J</i> =7.4 Hz, 1H), 6.36 (dd, <i>J</i> =8.9, 5.9 Hz, 1H), 4.33 (br t, <i>J</i> =5.5 Hz, 2H), 3.72 (br dd, <i>J</i> =16.6, 9.2 Hz, 2H), 3.55 - 3.47 (m, 3H), 3.46 - 3.35 (m, 2H), 3.24 - 3.15 (m, 3H), 2.96 (s, 3H), 2.81 (t, <i>J</i> =6.2 Hz, 2H), 2.01 - 1.88 (m, 2H). LC/MS (<i>m/z</i>) = 570.2 (<i>M</i> +H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 29.	Example 3

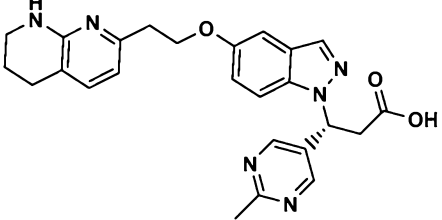
Example No.	Structure & Name	Analytical Data	Method
155	 <p data-bbox="427 660 863 831">3-(5-cyclopropylpyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, methanol- <i>d</i> ₄) δ 8.52 (br s, 1H), 8.44 (br s, 1H), 8.02 (br d, <i>J</i> =9.6 Hz, 2H), 7.66 - 7.55 (m, 2H), 7.27 - 7.13 (m, 1H), 7.07 (dd, <i>J</i> =9.1, 1.9 Hz, 1H), 6.72 (d, <i>J</i> =7.4 Hz, 1H), 6.36 (br dd, <i>J</i> =8.9, 5.6 Hz, 1H), 4.31 (br t, <i>J</i> =5.6 Hz, 2H), 3.68 (br dd, <i>J</i> =16.8, 9.1 Hz, 1H), 3.48 (br t, <i>J</i> =5.5 Hz, 2H), 3.39 (br dd, <i>J</i> =16.8, 5.5 Hz, 1H), 3.18 (br t, <i>J</i> =5.8 Hz, 2H), 2.80 (br t, <i>J</i> =6.1 Hz, 2H), 2.15 - 2.00 (m, 1H), 1.93 (quin, <i>J</i> =5.8 Hz, 2H), 1.25 - 1.12 (m, 2H), 0.95 - 0.77 (m, 2H). LC/MS (<i>m/z</i>) = 484.1 (<i>M</i> + <i>H</i>) ⁺ . Human α V β 6 IC ₅₀ (nM) = 5.3.	Example 3

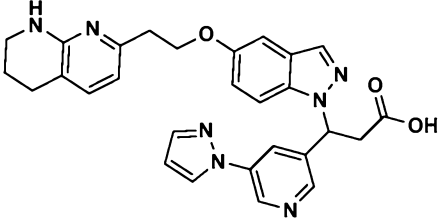
Example No.	Structure & Name	Analytical Data	Method
156	 <p>3-(5-(4-methylpiperazin-1-yl)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, methanol- <i>d</i> ₄) δ 8.42 - 8.33 (m, 1H), 8.21 (s, 1H), 8.07 - 8.01 (m, 2H), 7.67 - 7.57 (m, 2H), 7.21 (d, <i>J</i> =2.2 Hz, 1H), 7.09 (dd, <i>J</i> =9.1, 2.2 Hz, 1H), 6.74 (d, <i>J</i> =7.2 Hz, 1H), 6.37 (dd, <i>J</i> =9.2, 5.6 Hz, 1H), 4.33 (br t, <i>J</i> =5.1 Hz, 2H), 3.73 (br dd, <i>J</i> =16.8, 9.4 Hz, 2H), 3.55 - 3.48 (m, 3H), 3.42 (br dd, <i>J</i> =16.9, 5.6 Hz, 2H), 3.24 - 3.17 (m, 2H), 2.99 (s, 3H), 2.82 (br t, <i>J</i> =6.1 Hz, 2H), 1.99 - 1.88 (m, 2H). LC/MS (<i>m/z</i>) = 542.2 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 8.8.	Example 3

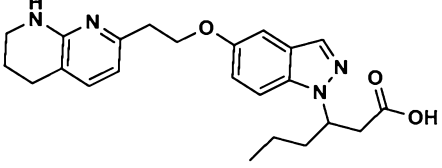
Example No.	Structure & Name	Analytical Data	Method
157	 <p>3-(5-(azetidine-1-carbonyl)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, methanol- <i>d</i> ₄) δ 8.82 - 8.72 (m, 2H), 8.18 - 8.06 (m, 1H), 8.02 (s, 1H), 7.64 - 7.57 (m, 2H), 7.20 (d, <i>J</i> =2.2 Hz, 1H), 7.07 (dd, <i>J</i> =9.1, 2.2 Hz, 1H), 6.74 (d, <i>J</i> =7.4 Hz, 1H), 6.38 (dd, <i>J</i> =8.8, 5.8 Hz, 1H), 4.39 - 4.23 (m, 4H), 4.19 (br t, <i>J</i> =7.7 Hz, 2H), 3.72 (dd, <i>J</i> =16.8, 9.1 Hz, 1H), 3.55 - 3.46 (m, 2H), 3.45 - 3.36 (m, 1H), 3.19 (br t, <i>J</i> =5.8 Hz, 2H), 2.81 (br t, <i>J</i> =5.9 Hz, 2H), 2.43 - 2.31 (m, 2H), 2.00 - 1.89 (m, 2H). LC/MS (<i>m/z</i>) = 527.0 (<i>M</i> + <i>H</i>) ⁺ . Human αVβ6 IC ₅₀ (nM) = 32.	Example 3

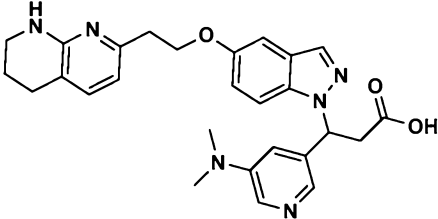
Example No.	Structure & Name	Analytical Data	Method
158	 <p data-bbox="414 660 873 884">3-((5-((2-(dimethylamino)ethyl)carbamoyl)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, methanol- <i>d</i> ₄) δ 8.93 (br s, 1H), 8.75 (br s, 1H), 8.36 (br s, 1H), 8.00 (s, 1H), 7.68 - 7.57 (m, 2H), 7.20 (s, 1H), 7.07 (br d, <i>J</i> =8.5 Hz, 1H), 6.74 (br d, <i>J</i> =7.2 Hz, 1H), 6.37 (br dd, <i>J</i> =8.3, 5.8 Hz, 1H), 4.33 (br s, 2H), 3.85 - 3.69 (m, 3H), 3.50 (br s, 2H), 3.45 - 3.36 (m, 3H), 3.23 - 3.13 (m, 2H), 2.99 (s, 6H), 2.81 (br t, <i>J</i> =5.5 Hz, 2H), 2.00 - 1.89 (m, 2H). LC/MS (<i>m/z</i>) = 558.2 (<i>M</i> + <i>H</i>) ⁺ . Human α V β 6 IC ₅₀ (nM) = 13.	Example 3

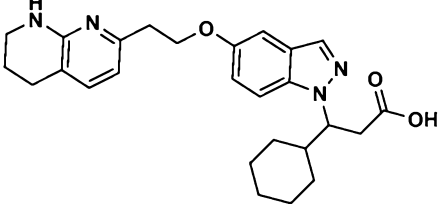
Example No.	Structure & Name	Analytical Data	Method
159	 <p data-bbox="422 667 865 840">(S)-3-(2-methylpyrimidin-5-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, methanol- <i>d</i> ₄) δ 8.70 (s, 2H), 8.05 - 7.98 (m, 1H), 7.67 - 7.56 (m, 2H), 7.24 - 7.17 (m, 1H), 7.13 - 7.04 (m, 1H), 6.81 - 6.69 (m, 1H), 6.34 - 6.25 (m, 1H), 4.38 - 4.30 (m, 2H), 3.75 - 3.65 (m, 1H), 3.55 - 3.48 (m, 2H), 3.42 - 3.35 (m, 1H), 3.23 - 3.16 (m, 2H), 2.86 - 2.77 (m, 2H), 2.69 - 2.57 (m, 3H), 1.98 - 1.91 (m, 2H). LC/MS (<i>m/z</i>) = 459.2 (<i>M</i> +H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 47.	Examples 66, 16, and 17

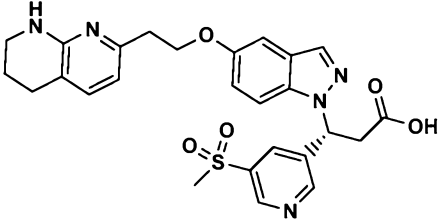
Example No.	Structure & Name	Analytical Data	Method
160	 <p data-bbox="422 667 865 837">(R)-3-(2-methylpyrimidin-5-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, methanol- <i>d</i> ₄) δ 8.70 (s, 2H), 8.05 - 7.98 (m, 1H), 7.67 - 7.56 (m, 2H), 7.24 - 7.17 (m, 1H), 7.13 - 7.04 (m, 1H), 6.81 - 6.69 (m, 1H), 6.34 - 6.25 (m, 1H), 4.38 - 4.30 (m, 2H), 3.75 - 3.65 (m, 1H), 3.55 - 3.48 (m, 2H), 3.42 - 3.35 (m, 1H), 3.23 - 3.16 (m, 2H), 2.86 - 2.77 (m, 2H), 2.69 - 2.57 (m, 3H), 1.98 - 1.91 (m, 2H). LC/MS (<i>m/z</i>) = 459.2 (<i>M</i> +H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 2,600.	Example 3

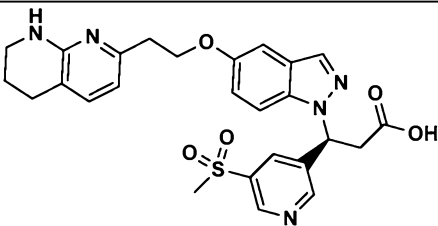
Example No.	Structure & Name	Analytical Data	Method
161	 <p data-bbox="422 660 861 840">3-(5-(1H-pyrazol-1-yl)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, chloroform- <i>d</i>) δ 9.58 - 9.40 (m, 1H), 9.08 - 9.01 (m, 1H), 8.83 - 8.75 (m, 1H), 8.51 - 8.42 (m, 1H), 8.02 - 7.96 (m, 2H), 7.81 - 7.76 (m, 1H), 7.39 - 7.33 (m, 2H), 7.07 (br d, <i>J</i> =2.1 Hz, 1H), 6.97 - 6.94 (m, 1H), 6.57 - 6.54 (m, 1H), 6.52 - 6.49 (m, 1H), 6.37 - 6.29 (m, 1H), 4.31 - 4.22 (m, 2H), 4.01 - 3.99 (m, 1H), 3.84 - 3.77 (m, 1H), 3.51 - 3.46 (m, 3H), 3.19 - 3.13 (m, 2H), 2.78 - 2.72 (m, 2H), 1.96 - 1.89 (m, 2H). LC/MS (<i>m/z</i>) = 510.2 (M+H) ⁺ . Human α V β 6 IC ₅₀ (nM) = 5.5.	Example 3

Example No.	Structure & Name	Analytical Data	Method
162	 <p>3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)hexanoic acid</p>	¹ H NMR (500 MHz, DMSO- <i>d</i> ₆) δ 8.25 - 8.09 (m, 1H), 7.51 - 7.44 (m, 1H), 7.12 - 7.06 (m, 1H), 7.02 - 6.97 (m, 1H), 6.87 - 6.81 (m, 1H), 6.42 - 6.36 (m, 1H), 4.80 (dt, <i>J</i> =9.0, 4.7 Hz, 1H), 4.24 - 4.15 (m, 2H), 3.28 - 3.19 (m, 2H), 3.03 - 2.84 (m, 4H), 2.65 - 2.57 (m, 2H), 1.95 - 1.85 (m, 2H), 1.81 - 1.69 (m, 3H), 1.14 - 1.01 (m, 1H), 0.94 - 0.81 (m, 1H), 0.82 - 0.74 (m, 3H). LC/MS (<i>m/z</i>) = 409.2 (<i>M</i> +H) ⁺ . Human αVβ6 IC ₅₀ (nM) = 730.	Example 3

Example No.	Structure & Name	Analytical Data	Method
163	 <p>3-(5-(dimethylamino)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, DMSO- <i>d</i> ₆) δ 8.01 - 7.96 (m, 1H), 7.95 - 7.93 (m, 1H), 7.85 - 7.78 (m, 1H), 7.68 - 7.63 (m, 1H), 7.19 - 7.14 (m, 1H), 7.08 - 7.04 (m, 2H), 6.99 - 6.93 (m, 1H), 6.39 - 6.31 (m, 2H), 6.17 - 6.10 (m, 1H), 4.28 - 4.17 (m, 2H), 3.64 - 3.55 (m, 1H), 3.26 - 3.20 (m, 1H), 2.92 - 2.83 (m, 8H), 2.64 - 2.58 (m, 2H), 1.77 - 1.71 (m, 2H). LC/MS (m/z) = 487.2 (M+H) ⁺ . Human αVβ6 IC ₅₀ (nM) = 3.8.	Example 3

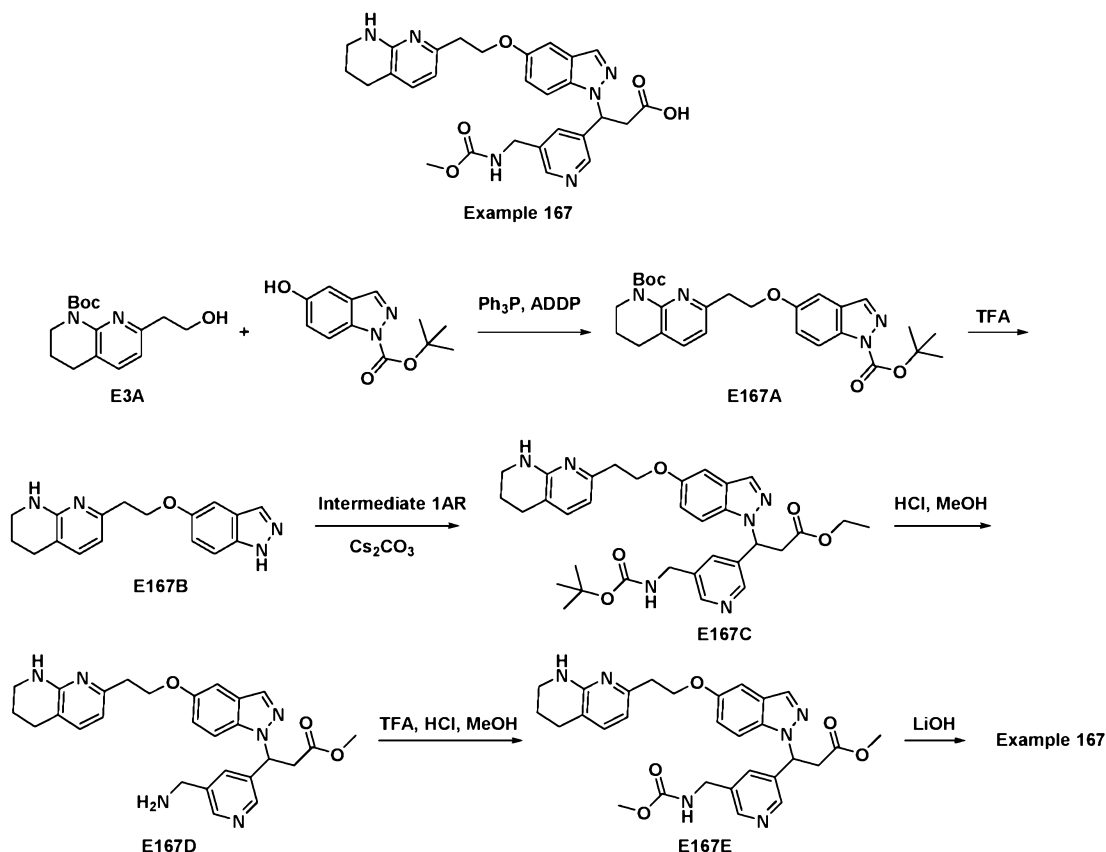
Example No.	Structure & Name	Analytical Data	Method
164	 <p data-bbox="414 651 873 779">3-cyclohexyl-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, DMSO- <i>d</i> ₆) δ 7.73 - 7.67 (m, 1H), 7.36 - 7.30 (m, 1H), 6.92 (s, 1H), 6.86 (d, <i>J</i> =7.0 Hz, 1H), 6.78 - 6.72 (m, 1H), 6.18 (d, <i>J</i> =7.3 Hz, 1H), 6.12 - 6.06 (m, 1H), 4.55 - 4.45 (m, 1H), 4.07 - 3.99 (m, 2H), 3.07 - 2.99 (m, 1H), 2.81 - 2.66 (m, 3H), 2.47 - 2.37 (m, 2H), 1.67 - 1.42 (m, 5H), 1.37 - 1.23 (m, 2H), 1.05 - 0.89 (m, 2H), 0.86 - 0.52 (m, 5H). LC/MS (<i>m/z</i>) = 449.1 (<i>M</i> +H) ⁺ . Human αVβ6 IC ₅₀ (nM) = 3,200.	Example 3

Example No.	Structure & Name	Analytical Data	Method
165	 <p data-bbox="422 660 863 840"><i>(R)</i>-3-(5-(methylsulfonyl)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, DMSO- <i>d</i> ₆) δ 9.00 - 8.91 (m, 2H), 8.30 - 8.23 (m, 1H), 8.11 - 8.03 (m, 1H), 7.83 - 7.75 (m, 1H), 7.65 - 7.57 (m, 1H), 7.26 - 7.19 (m, 1H), 7.07 - 6.99 (m, 1H), 6.76 - 6.69 (m, 1H), 6.49 - 6.40 (m, 1H), 4.33 - 4.22 (m, 2H), 3.93 - 3.86 (m, 1H), 3.71 - 3.60 (m, 1H), 3.32 - 3.28 (m, 1H), 3.19 - 3.08 (m, 2H), 2.76 - 2.68 (m, 2H), 2.54 - 2.52 (m, 3H), 1.86 - 1.75 (m, 2H), 1.29 - 1.19 (m, 1H). LC/MS (m/z) = 522.2 (M+H) ⁺ . Human αVβ6 IC ₅₀ (nM) = 540.	Examples 125, 16, and 17

Example No.	Structure & Name	Analytical Data	Method
166	 <p>(S)-3-(5-(methylsulfonyl)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, DMSO- <i>d</i> ₆) δ 9.00 - 8.91 (m, 2H), 8.30 - 8.23 (m, 1H), 8.11 - 8.03 (m, 1H), 7.83 - 7.75 (m, 1H), 7.65 - 7.57 (m, 1H), 7.26 - 7.19 (m, 1H), 7.07 - 6.99 (m, 1H), 6.76 - 6.69 (m, 1H), 6.49 - 6.40 (m, 1H), 4.33 - 4.22 (m, 2H), 3.93 - 3.86 (m, 1H), 3.71 - 3.60 (m, 1H), 3.32 - 3.28 (m, 1H), 3.19 - 3.08 (m, 2H), 2.76 - 2.68 (m, 2H), 2.54 - 2.52 (m, 3H), 1.86 - 1.75 (m, 2H), 1.29 - 1.19 (m, 1H). LC/MS (m/z) = 522.2 (M+H) ⁺ . Human αVβ6 IC ₅₀ (nM) = 5.2.	Examples 125, 16, and 17

Example 167

3-(5-(((Methoxycarbonyl)amino)methyl)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid, 2 TFA



Intermediate **E167A**: To a solution of Intermediate **E3A** (3.53 g, 12.7 mmol), *tert*-butyl 5-hydroxy-1H-indazole-1-carboxylate [(WO 2016/21043), 2.7 g, 11.5 mmol] and
 5 Ph_3P (3.78 g, 14.4 mmol) in THF (70 mL), maintained in an ice-water bath, was added (*E*)-diazene-1,2-diylbis(piperidin-1-ylmethanone) (3.64 g, 14.41 mmol) dropwise over 5 min. The reaction mixture was allowed to warm to rt and stir for 16 h. The reaction was diluted with NaHCO_3 solution (aqueous, saturated, 30 mL), the resulting aqueous mixture was extracted with EtOAc (3 x 50 mL). The combine organic layers were washed with
 10 brine (10 mL), and then dried over Na_2SO_4 . The mixture was filtered and concentrated under reduced pressure. The residue was purified via flash chromatography (hexanes/ethyl acetate, 0-100 % gradient) to give Intermediate **E167A** (3.94 g, 69%). ^1H NMR (500 MHz, chloroform- d) δ 8.09 - 8.01 (m, 2H), 7.37 - 7.32 (m, 1H), 7.20 - 7.15 (m, 2H), 6.98 - 6.93 (m, 1H), 4.44 (t, J =6.9 Hz, 2H), 3.81 - 3.76 (m, 2H), 3.29 - 3.20 (m,

2H), 2.80 - 2.74 (m, 2H), 1.98 - 1.92 (m, 2H), 1.76 - 1.75 (m, 1H), 1.75 - 1.73 (m, 9H), 1.53 (s, 9H). LCMS (ES): m/z 495.1 $[M+H]^+$.

Intermediate **E167B**: To a solution of Intermediate **E167A** (3.94 g, 7.97 mmol) in DCM (40 mL) was added TFA (8 mL, 104 mmol) and the mixture was stirred at rt for 16
5 hrs. It was concentrated and the crude product was purified using medium pressure reverse phase chromatography (10-90% water 0.1% TFA/ acetonitrile gradient) to afford Intermediate **E167B**, TFA salt (2.63 g, 6.44 mmol, 81 % yield). 1H NMR (500 MHz, chloroform- d) δ 10.39 - 10.23 (m, 1H), 8.17 - 7.88 (m, 1H), 7.45 - 7.38 (m, 1H), 7.37 - 7.31 (m, 1H), 7.19 - 7.12 (m, 1H), 7.11 - 6.96 (m, 1H), 6.59 - 6.48 (m, 1H), 4.48 - 4.23
10 (m, 2H), 3.63 - 3.41 (m, 2H), 3.31 - 3.09 (m, 2H), 2.84 - 2.61 (m, 2H), 2.04 - 1.83 (m, 2H). LCMS (ES): m/z 295.2 $[M+H]^+$.

Intermediate **E167C**: To a solution of Intermediate **E167B**, TFA salt (100 mg, 0.245 mmol) in acetonitrile (2 mL) was added cesium carbonate (239 mg, 0.735 mmol). After stirring at rt for 5 min, Intermediate **1AR** (75 mg, 0.245 mmol) was added and the
15 resulting mixture was stirred at 80 °C for 8 hrs. The mixture was cooled to rt, filtered, and concentrated. The residue purified *via* preparative HPLC (Phenomenex Luna Axia 5 μ C18 30 x 100 mm; 10 min gradient from 75% A: 25% B to 0% A:100% B (A = 90% H₂O/10% MeOH + 0.1% TFA); (B = 90% MeOH/10% H₂O + 0.1% TFA); detection at 220 nm) to give Intermediate **E167C**, bis TFA salt (95 mg, 0.115 mmol, 47 % yield).
20 LCMS (ES): m/z 601.3 $[M+H]^+$.

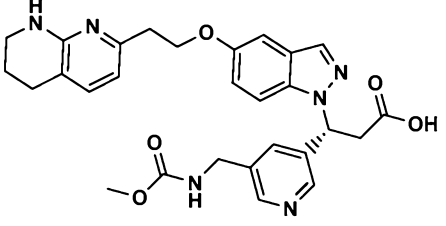
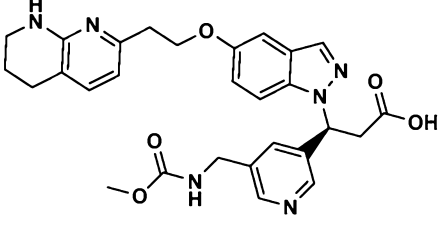
Intermediate **E167D**: To a solution of Intermediate **E167C**, bis TFA salt (95 mg, 0.115 mmol) in methanol (1 mL) was added a 4 M solution of HCl in dioxane (0.115 mL, 0.459 mmol). The reaction mixture was stirred at rt for 3 days. The mixture was diluted with acetonitrile and purified using reverse phase preparative HPLC (Phenomenex
25 Luna Axia 5 μ C18 30 x 100 mm; 10 min gradient from 75% A: 25% B to 0% A:100% B (A = 90% H₂O/10% MeOH + 0.1% TFA); (B = 90% MeOH/10% H₂O + 0.1% TFA); detection at 220 nm) to give Intermediate **E167D**, 3 TFA salt (50 mg, 0.060 mmol, 53 % yield). LCMS (ES): m/z 487.1 $[M+H]^+$.

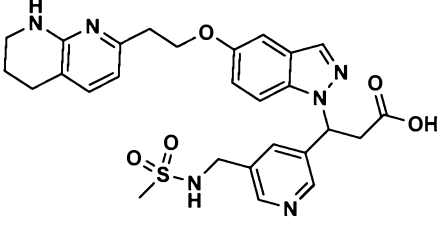
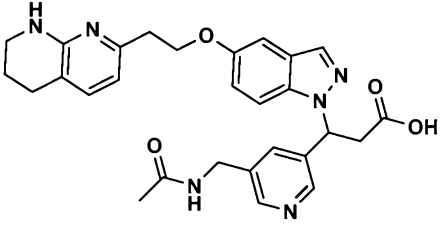
Intermediate **E167E**: To a solution of **E167D**, 3 TFA (16 mg, 0.019 mmol) in
30 DCM (0.5 mL) was added triethylamine (0.013 mL, 0.097 mmol). The mixture was stirred at rt for 10 min and then methyl carbonochloridate (2.74 mg, 0.029 mmol) was

added and stirred at rt for 4 hrs. After 4 hrs, the reaction mixture was diluted with sat aqueous NaHCO₃ solution and then extracted with dichloromethane. The combine organics layers were washed with brine, dried over sodium sulfate, filtered, and concentrated in vacuo. The mixture was diluted with acetonitrile and purified using reverse phase preparative HPLC (Phenomenex Luna Axia 5 μ C18 30 x 100 mm; 10 min gradient from 20% A: 80% B to 0% A:100% B (A = 90% H₂O/10% MeOH + 0.1% TFA); (B = 90% MeOH/10% H₂O + 0.1% TFA); detection at 220 nm) to give Intermediate **E167E**, bis TFA salt (13 mg, 0.017 mmol, 87 % yield). LCMS (ES): m/z 545.1 [M+H]⁺.

Example **167**: To a solution of Intermediate **E167E**, bis TFA salt (13 mg, 0.017 mmol) in THF (0.5 mL) was added a solution of aqueous 1 M LiOH (0.067 mL, 0.067 mmol). The reaction mixture was stirred at rt for 3 hrs. The reaction mixture was neutralized with TFA, filtered, and concentrated under reduced pressure. The residue was diluted purified using reverse phase preparative HPLC (Phenomenex Luna Axia 5 μ C18 30 x 100 mm; 10 min gradient from 20% A: 80% B to 0% A:100% B (A = 90% H₂O/10% MeOH + 0.1% TFA); (B = 90% MeOH/10% H₂O + 0.1% TFA); detection at 220 nm) to give Intermediate **E167**, bis TFA salt (12 mg, 0.016 mmol, 94 % yield). ¹H NMR (500 MHz, methanol-*d*₄) δ 8.55 (br s, 1H), 8.47 (br s, 1H), 8.10 - 7.91 (m, 2H), 7.59 (br t, *J*=9.4 Hz, 2H), 7.20 (d, *J*=2.2 Hz, 1H), 7.07 (dd, *J*=9.1, 2.2 Hz, 1H), 6.77 - 6.72 (m, 1H), 6.34 (br dd, *J*=9.2, 5.4 Hz, 1H), 4.37 - 4.28 (m, 4H), 3.72 (br dd, *J*=16.8, 9.4 Hz, 1H), 3.65 (s, 3H), 3.54 - 3.46 (m, 2H), 3.43 - 3.35 (m, 1H), 3.19 (br t, *J*=5.9 Hz, 2H), 2.82 (br t, *J*=6.1 Hz, 2H), 1.99 - 1.91 (m, 2H). LC/MS (m/z) = 531.1 (M+H)⁺. Human α V β 6 IC₅₀ (nM) = 15.

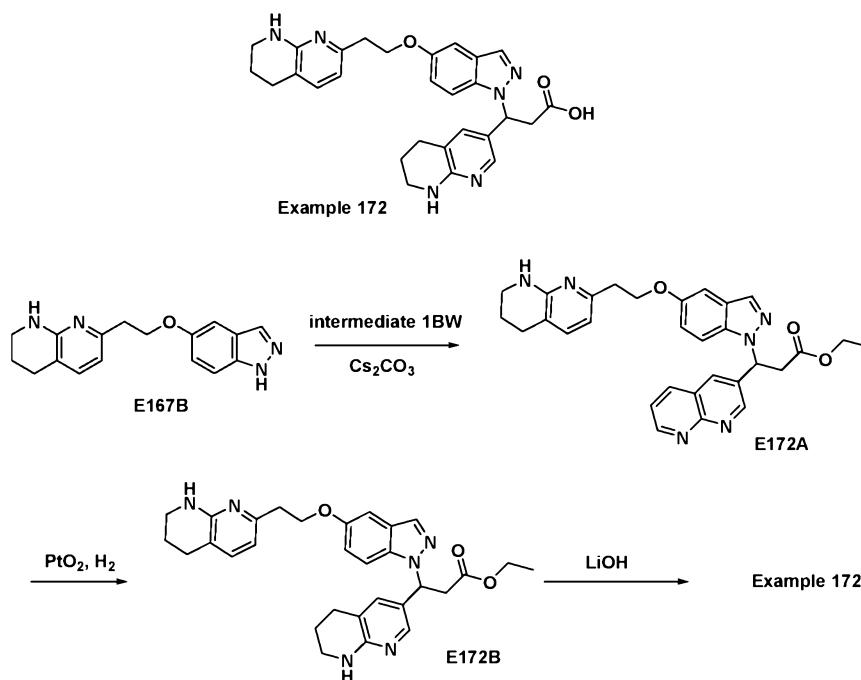
The following examples were prepared using methods analogous to the ones indicated in the table below.

Example No.	Structure & Name	Analytical Data	Method
168	 <p>(<i>R</i>)-3-(5-(((methoxycarbonyl)amino)methyl)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, methanol- <i>d</i> ₄) δ 8.48 - 8.41 (m, 1H), 8.39 - 8.33 (m, 1H), 8.00 - 7.94 (m, 1H), 7.81 - 7.75 (m, 1H), 7.54 - 7.49 (m, 1H), 7.32 - 7.28 (m, 1H), 7.04 - 7.00 (m, 1H), 6.99 - 6.93 (m, 1H), 6.55 - 6.50 (m, 1H), 6.33 - 6.27 (m, 1H), 4.30 - 4.25 (m, 2H), 4.16 - 4.03 (m, 2H), 3.66 - 3.56 (m, 4H), 3.40 - 3.36 (m, 5H), 3.25 - 3.19 (m, 1H), 2.97 - 2.93 (m, 2H), 2.73 - 2.68 (m, 2H), 1.90 - 1.83 (m, 2H). LC/MS (m/z) = 531.4 (M+H) ⁺ . Human αVβ6 IC ₅₀ (nM) = 740.	Examples 167, 16, and 17
169	 <p>(<i>S</i>)-3-(5-(((methoxycarbonyl)amino)methyl)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, methanol- <i>d</i> ₄) δ 8.52 - 8.30 (m, 2H), 8.00 - 7.93 (m, 1H), 7.80 - 7.76 (m, 1H), 7.56 - 7.49 (m, 1H), 7.32 - 7.27 (m, 1H), 7.05 - 7.01 (m, 1H), 6.99 - 6.94 (m, 1H), 6.55 - 6.50 (m, 1H), 6.33 - 6.26 (m, 1H), 4.28 (s, 2H), 4.16 - 4.04 (m, 2H), 3.66 - 3.55 (m, 4H), 3.41 - 3.36 (m, 3H), 3.24 - 3.18 (m, 1H), 2.98 - 2.93 (m, 2H), 2.73 - 2.68 (m, 2H), 1.90 - 1.84 (m, 2H). LC/MS (m/z) = 515.1 (M+H) ⁺ . Human αVβ6 IC ₅₀ (nM) = 3.7.	Examples 167, 16, and 17

Example No.	Structure & Name	Analytical Data	Method
170	 <p>3-(5-(methylsulfonamidomethyl)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, methanol- <i>d</i> ₄) δ 8.59 - 8.45 (m, 2H), 8.07 - 7.93 (m, 2H), 7.68 - 7.55 (m, 2H), 7.20 (d, <i>J</i> =1.9 Hz, 1H), 7.06 (dd, <i>J</i> =9.4, 2.2 Hz, 1H), 6.77 (br d, <i>J</i> =7.4 Hz, 1H), 6.40 - 6.25 (m, 1H), 4.43 - 4.22 (m, 4H), 3.74 (br dd, <i>J</i> =16.8, 9.4 Hz, 1H), 3.51 (br t, <i>J</i> =5.6 Hz, 3H), 3.23 - 3.15 (m, 2H), 2.90 - 2.78 (m, 4H), 2.00 - 1.88 (m, 2H). LC/MS (<i>m/z</i>) = 552.1 (M+H) ⁺ . Human αVβ6 IC ₅₀ (nM) = 18.	Example 167
171	 <p>3-(5-(acetamidomethyl)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid</p>	¹ H NMR (500 MHz, methanol- <i>d</i> ₄) δ 8.63 - 8.49 (m, 1H), 8.49 - 8.43 (m, 1H), 8.07 - 7.84 (m, 2H), 7.65 - 7.55 (m, 2H), 7.21 (d, <i>J</i> =1.9 Hz, 1H), 7.07 (dd, <i>J</i> =9.2, 2.1 Hz, 1H), 6.77 (br d, <i>J</i> =6.1 Hz, 1H), 6.38 - 6.30 (m, 1H), 4.44 - 4.36 (m, 2H), 4.33 (br t, <i>J</i> =5.1 Hz, 2H), 3.97 - 3.89 (m, 1H), 3.72 (dd, <i>J</i> =16.8, 9.4 Hz, 1H), 3.54 - 3.48 (m, 2H), 3.23 - 3.15 (m, 2H), 2.83 (br t, <i>J</i> =6.2 Hz, 2H), 2.00 - 1.91 (m, 5H). LC/MS (<i>m/z</i>) = 515.1 (M+H) ⁺ . Human αVβ6 IC ₅₀ (nM) = 46.	Example 167

Example 172

3-(5-(2-(5,6,7,8-Tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)-3-(5,6,7,8-tetrahydro-1,8-naphthyridin-3-yl)propanoic acid, 2 TFA



Intermediate **E172A**: To a solution of Intermediate **E167B**, TFA salt (55 mg, 0.135 mmol) in acetonitrile (1.2 mL) was added cesium carbonate (132 mg, 0.404 mmol). After stirring at rt for 5 min, Intermediate **1BW** (30.7 mg, 0.135) was added and the resulting mixture was stirred at 80 °C for 6 hrs. The mixture was cooled to rt, filtered, and concentrated. The residue purified *via* preparative HPLC (Phenomenex Luna Axia 5μ C18 30 x 100 mm; 10 min gradient from 20% A: 80% B to 0% A:100% B (A = 90% H₂O/10% MeOH + 0.1% TFA); (B = 90% MeOH/10% H₂O + 0.1% TFA); detection at 220 nm) to give Intermediate **E172A**, bis TFA salt (62 mg, 0.083 mmol, 61.3 % yield). LCMS (ES): m/z 523.1 [M+H]⁺.

Intermediate **E172B**: To a degassed solution of **E172A** in EtOH (0.7 mL) was added platinum(IV) oxide (3 mg, 0.013 mmol). The reaction was stirred at rt for 4 hrs with under an atmosphere of hydrogen (balloon). The reaction was purged with nitrogen, filtered, and concentrated in vacuo. The crude product was diluted with MeCN, filtered, and purified using reverse phase preparative HPLC (Phenomenex Luna Axia 5μ C18 30 x 100 mm; 10 min gradient from 75% A: 25% B to 0% A:100% B (A = 90% H₂O/10% MeOH + 0.1% TFA); (B = 90% MeOH/10% H₂O + 0.1% TFA); detection at 220 nm) to

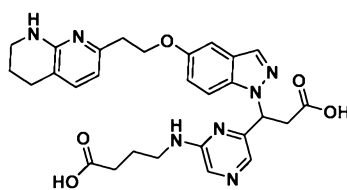
give Intermediate **E172B**, bis TFA salt (22 mg, 0.029 mmol, 35.3 % yield). LCMS (ES): m/z 527.1 $[M+H]^+$.

Example 172: To a solution of Intermediate **E172B**, bis TFA salt (22 mg, 0.029 mmol) in THF (0.5 mL) was added a solution of aqueous 1 M LiOH (0.117 mL, 0.117 mmol). The reaction mixture was stirred at rt overnight. The reaction mixture was neutralized with TFA, filtered, and concentrated under reduced pressure. The residue was diluted purified using reverse phase preparative HPLC (Phenomenex Luna Axia 5 μ C18 30 x 100 mm; 10 min gradient from 20% A: 80% B to 0% A:100% B (A = 90% H₂O/10% MeOH + 0.1% TFA); (B = 90% MeOH/10% H₂O + 0.1% TFA); detection at 220 nm) to give Intermediate **E172**, bis TFA salt (11 mg, 0.014 mmol, 46.7 % yield). ¹H NMR (500 MHz, methanol-*d*₄) δ 7.96 (s, 1H), 7.72 (s, 1H), 7.70 (br s, 1H), 7.65 - 7.52 (m, 2H), 7.18 (s, 1H), 7.06 (br d, J =9.1 Hz, 1H), 6.73 (d, J =7.4 Hz, 1H), 6.10 (br dd, J =8.8, 6.1 Hz, 1H), 4.31 (br t, J =5.6 Hz, 2H), 3.59 (br dd, J =16.5, 9.1 Hz, 1H), 3.53 - 3.39 (m, 4H), 3.30 - 3.22 (m, 1H), 3.18 (br t, J =5.5 Hz, 2H), 2.87 - 2.72 (m, 4H), 2.01 - 1.85 (m, 4H). LC/MS (m/z) = 499.1 $(M+H)^+$. Human α V β 6 IC₅₀ (nM) = 12.

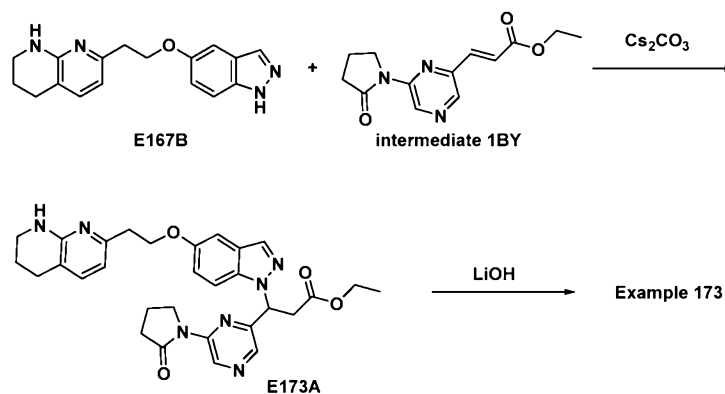
Example 173

4-(((6-(2-Carboxy-1-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)ethyl)pyrazin-2-yl)amino)butanoic acid, TFA

20



Example 173



Intermediate **E173A**: To a solution of Intermediate **E167B**, TFA salt (136 mg, 0.463 mmol) in acetonitrile (2.5 mL) was added cesium carbonate (453 mg, 1.389 mmol). After stirring at rt for 5 min, Intermediate **1BY** (121 mg, 0.463 mmol) was added and the resulting mixture was stirred at 80 °C for 16 hrs. The mixture was cooled to rt, filtered, and concentrated. The residue purified *via* preparative HPLC (Phenomenex Luna Axia 5μ C18 30 x 100 mm; 10 min gradient from 20% A: 80% B to 0% A:100% B (A = 90% H₂O/10% MeOH + 0.1% TFA); (B = 90% MeOH/10% H₂O + 0.1% TFA); detection at 220 nm) to give Intermediate **E173A**, bis TFA salt (19.6 mg, 0.035 mmol, 7 % yield). LCMS (ES): m/z 556.3 [M+H]⁺.

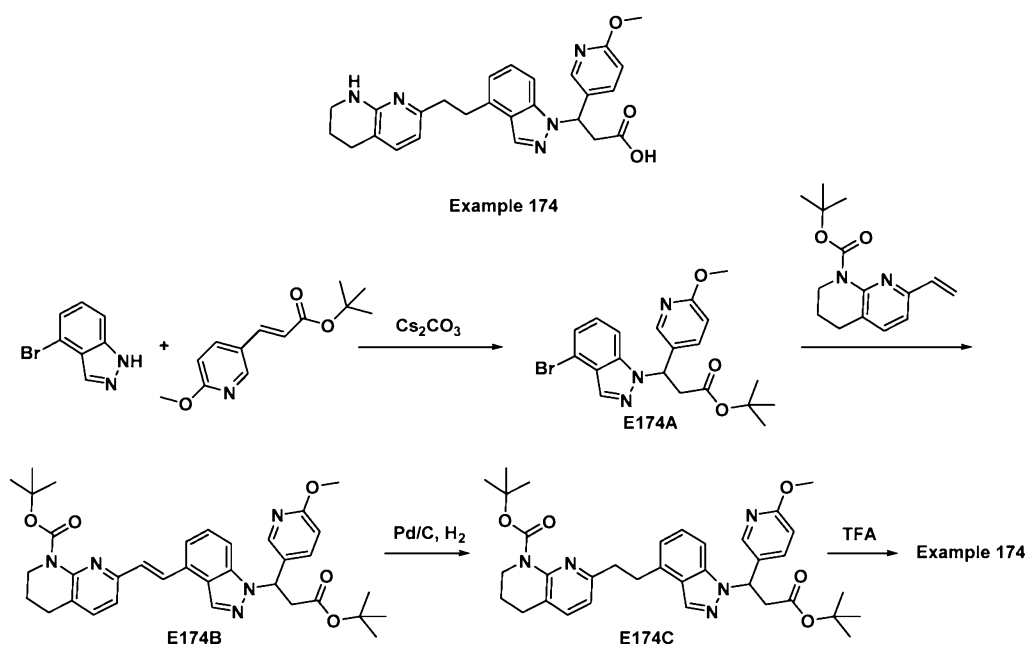
Example **173**: To a solution of Intermediate **E173A**, bis TFA salt (19.6 mg, 0.035 mmol) in THF (1 mL) was added a solution of aqueous 1 M LiOH (0.106 mL, 0.106 mmol). The reaction mixture was stirred at rt overnight. The reaction mixture was concentrated under reduced pressure, diluted with acetonitrile and filtered. The product was purified using reverse phase preparative HPLC (Phenomenex Luna Axia 5μ C18 30 x 100 mm; 10 min gradient from 20% A: 80% B to 0% A:100% B (A = 90% H₂O/10% MeOH + 0.1% TFA); (B = 90% MeOH/10% H₂O + 0.1% TFA); detection at 220 nm) to give Example **173**, TFA salt (8.1 mg, 0.011 mmol, 32 % yield). ¹H NMR (500 MHz,

methanol-*d*₄) δ 8.00 - 7.97 (m, 1H), 7.64 - 7.61 (m, 2H), 7.59 - 7.55 (m, 1H), 7.24 - 7.21 (m, 1H), 7.10 - 7.09 (m, 1H), 7.09 - 7.07 (m, 1H), 6.80 - 6.77 (m, 1H), 6.21 - 6.16 (m, 1H), 4.37 - 4.31 (m, 3H), 3.54 - 3.50 (m, 4H), 3.42 - 3.39 (m, 2H), 3.23 - 3.19 (m, 3H), 2.85 - 2.81 (m, 3H), 2.36 (t, $J=7.4$ Hz, 2H), 1.98 - 1.94 (m, 3H), 1.86 (t, $J=7.2$ Hz, 2H).

5 LC/MS (m/z) = 546.3 ($M+H$)⁺. Human $\alpha V\beta 6$ IC₅₀ (nM) = 370.

Example 174

3-(6-Methoxypyridin-3-yl)-3-(4-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethyl)-1H-indazol-1-yl)propanoic acid, TFA



10

Intermediate **E174A**: To a mixture of *tert*-butyl (*E*)-3-(6-methoxypyridin-3-yl)acrylate [(*J. Org. Chem.* **2004**, *69*, 1959) 0.746 g, 3.17 mmol] and 4-bromo-1H-indazole (0.500g, 2.54 mmol) in acetonitrile (20 mL) at room temperature was added DBU (0.383 mL, 2.54 mmol). The reaction mixture was heated at 50 °C for 48 h. The mixture was diluted with water and extracted with ethyl acetate. The combined organic layers were washed with brine, dried over magnesium sulfate, filtered and concentrated in vacuo. The crude residue was purified using silica gel column chromatography (2% MeOH/dichloromethane) to afford **E174A** (447 mg, 1.034 mmol, 41 % yield). ¹H NMR

15

20

(400 MHz, chloroform-*d*) δ 8.18 (d, $J=2.3$ Hz, 1H), 8.04 (s, 1H), 7.58 (dd, $J=8.8, 2.5$ Hz, 1H), 7.41 (d, $J=8.5$ Hz, 1H), 7.31 - 7.27 (m, 1H), 7.25 - 7.16 (m, 1H), 6.67 (d, $J=8.5$ Hz, 1H), 6.00 (dd, $J=9.2, 5.9$ Hz, 1H), 3.89 (s, 3H), 3.64 (dd, $J=16.1, 9.3$ Hz, 1H), 3.16 (dd, $J=16.1, 6.0$ Hz, 1H), 1.28 (s, 9H). LCMS (ES): m/z 432.2, 434.2 $[M+H]^+$.

- 5 Intermediate **E174B**: Palladium(II) acetate (8.62 mg, 0.038 mmol) was added to a vial charged with a degassed mixture of tert-butyl 7-vinyl-3,4-dihydro-1,8-naphthyridine-1(2H)-carboxylate [(*Eur. J. Med. Chem.* **2007**, 42, 334), 0.100 g, 0.384 mmol], intermediate **E174A** (0.166 g, 0.384 mmol), tri-*o*-tolylphosphine (0.023 g, 0.077 mmol), and triethylamine (0.107 mL, 0.768 mmol) in DMF (3 mL). The vessel's headspace was
10 purged with nitrogen and the vial was sealed. The mixture was heated at 100 °C for 20 h. After cooling to rt, the vessel was uncapped and its contents were diluted with water and extracted with ethyl acetate. The combined organic layers were washed with brine, dried over magnesium sulfate, filtered and concentrated in vacuo. The residue was purified using silica gel column chromatography (5% methanol/dichloromethane) to afford
15 **E174B** (238 mg, 0.389 mmol, 101 % yield). The product was a mixture of cis/trans isomers and was contaminated with minor impurities. The material was used for subsequent chemistry without further purification. LC/MS (m/z) = 612.43 ($M+H$)⁺.

- Intermediate **E174C**: To a flask charged with a solution of **E174B** (238 mg, 0.389 mmol) in methanol under a nitrogen atmosphere was added 10% palladium on carbon
20 (41.4 mg, 0.389 mmol). The vessel was partially evacuated and flushed repeatedly with hydrogen gas. The reaction was left to stir under a hydrogen atmosphere (double balloon). After 24 h, the reaction mixture was purged with nitrogen and filtered through celite. The concentrated filtrate was purified using silica gel column chromatography (5-10 % MeOH/dichloromethane) to afford **E174C** (25 mg, 0.040 mmol, 10 % yield).
25 LC/MS (m/z) = 614.4 ($M+H$)⁺.

- Example 174.** Trifluoroacetic acid (0.5 mL) was added to a solution containing **E174C** (41.8 mg, 0.068 mmol) in dichloromethane (2.5 mL). The resulting mixture was heated at 40 °C for 2 h. The mixture was concentrated under a stream of dry nitrogen. The residue was dissolved in methanol and purified via preparative LC/MS with the following
30 conditions: Column: XBridge C18, 19 x 200 mm, 5- μ m particles, Mobile Phase A: 5:95 acetonitrile: water with 10-mM ammonium acetate; Mobile Phase B: 95:5 acetonitrile:

water with 10-mM ammonium acetate; Gradient: 0-40% B over 20 minutes, then a 5-minute hold at 100% B; Flow: 20 mL/min. Fractions containing the desired product were combined and dried via centrifugal evaporation to afford example **172** (29.4 mg, 0.062 mmol, 91 % yield). ¹H NMR (500 MHz, DMSO-*d*₆) δ 8.30 - 8.22 (m, 1H), 8.22 - 8.15 (m, 1H), 7.74 - 7.65 (m, 1H), 7.65 - 7.56 (m, 1H), 7.31 - 7.22 (m, 1H), 7.07 - 6.99 (m, 1H), 6.97 - 6.90 (m, 1H), 6.77 - 6.69 (m, 1H), 6.38 - 6.27 (m, 2H), 6.25 - 6.13 (m, 1H), 3.82 - 3.73 (m, 3H), 3.63 (br s, 1H), 3.31 - 3.12 (m, 5H), 2.88 - 2.77 (m, 2H), 2.64 - 2.56 (m, 2H), 1.80 - 1.70 (m, 2H). LC/MS (m/z) = 458.2 (M+H)⁺. Human αVβ6 IC₅₀ (nM) = 540.

10

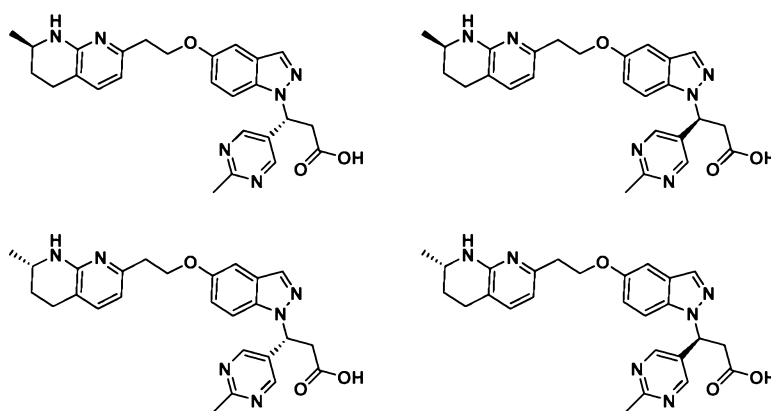
Examples 175-178

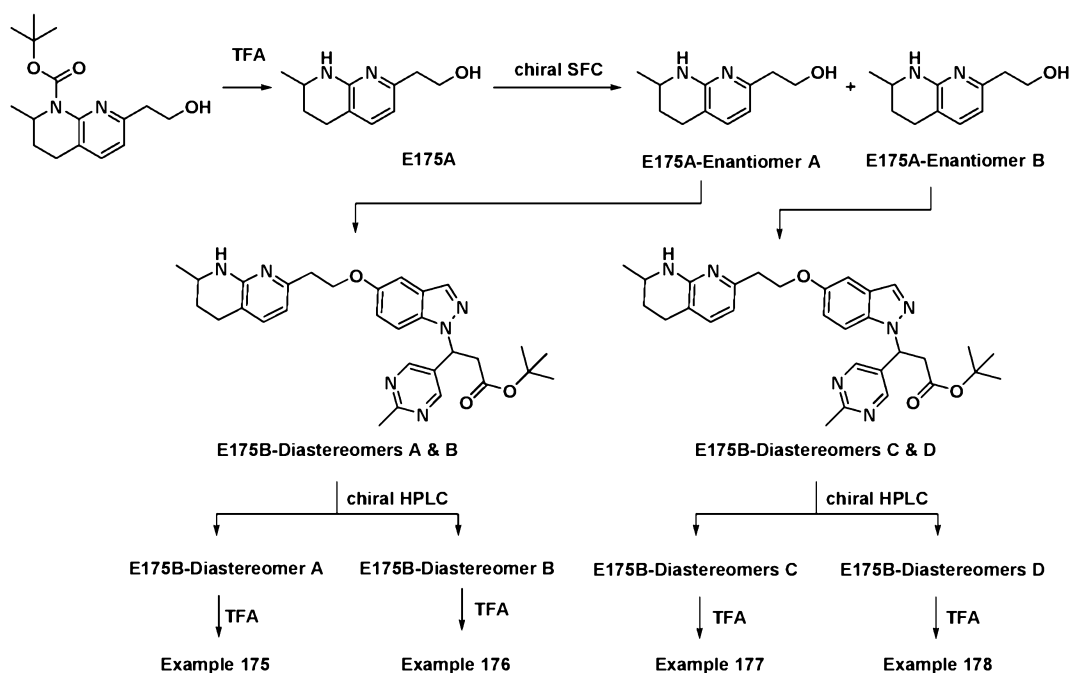
(*R*)-3-(5-(2-((*R*)-7-methyl-5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)-3-(2-methylpyrimidin-5-yl)propanoic acid;

15 **(*S*)-3-(5-(2-((*R*)-7-methyl-5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)-3-(2-methylpyrimidin-5-yl)propanoic acid;**

(*R*)-3-(5-(2-((*S*)-7-methyl-5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)-3-(2-methylpyrimidin-5-yl)propanoic acid;

20 **(*S*)-3-(5-(2-((*S*)-7-methyl-5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)-3-(2-methylpyrimidin-5-yl)propanoic acid**





Intermediate **E175A**: To a solution of *tert*-butyl 7-(2-hydroxyethyl)-2-methyl-3,4-dihydro-1,8-naphthyridine-1(2H)-carboxylate [(WO 2007/141473), 2.509 g, 8.58 mmol] in DCM (20 mL) was added trifluoroacetic acid (5 mL). The resulting mixture was allowed to stir at rt for 24 h. The reaction was concentrated in vacuo and diluted with aqueous saturated sodium bicarbonate solution. The aqueous mixture was extracted with ethyl acetate. The combined organic layers were washed with brine, dried over magnesium sulfate, filtered and concentrated in vacuo. The crude product was purified using silica gel column chromatography (ISCO system, prepacked ISCO silica cartridge, 90:10 dichloromethane/methanol) to afford **E175A** (2.509 g, 8.58 mmol). ¹H NMR (500 MHz, chloroform-*d*) δ 7.38 - 7.31 (m, 1H), 6.45 - 6.37 (m, 1H), 3.93 (s, 2H), 3.74 - 3.61 (m, 1H), 2.93 (t, *J*=5.9 Hz, 2H), 2.84 - 2.69 (m, 2H), 2.06 - 1.97 (m, 1H), 1.65 - 1.55 (m, 1H), 1.37 (d, *J*=6.6 Hz, 3H). LCMS (ES): *m/z* 193.1 [*M*+H]⁺.

Intermediates **E175B-Enantiomers A and B**. A sample of **E175A** (1.3 g) was subjected to preparative chiral SFC purification (Column: Chiralpak AD-H, 30 x 250 mm, 5 micron, BPR pressure: 120 bar, temperature 35 °C, flow rate: 70.0 mL/min, mobile phase: 50% MeCN w/0.1% DEA in CO₂, detector wavelength: 314 nm, stacked injections: 0.5 mL of 110 mg/mL solution) to afford **E175A-Enantiomer A** (374 mg) and

E175A-Enantiomer B (391 mg). LCMS data for each enantiomer was identical to the racemate. The absolute configuration of the separated enantiomers was not determined.

Intermediates **E175B-Diastereomers A & B**. A sample of **E175A-Enantiomer A** was subjected to the Mitsunobu coupling (with *tert*-butyl 5-hydroxy-1H-indazole-1-carboxylate, WO 2016/21043), BOC deprotection, and Michael addition (with
5 **intermediate CC**) employing the methods outlined in **Example 152** to afford a mixture of **E175B-Diastereomers A & B** (106 mg, 11 % yield over 3 steps). LCMS (ES): *m/z* 529.4 [M+H]⁺. A sample of **E175B-Diastereomers A & B** (106 mg) was subjected to preparative chiral HPLC purification (Column: Chiralpak OD, 21 x 250 mm, 10 micron,
10 flow rate: 15 mL/min, mobile phase: 20% ethanol/80% heptane, detector wavelength: 220 nm, injection: 1 mL of 40 mg/mL solution) to afford **E175B-Diastereomer A** (44.6 mg) and **E175B-Diastereomer B** (47.5 mg). LCMS data for the separated diastereomers was identical to the diastereomeric mixture. The relative and absolute stereochemistry of the individual diastereomers was not determined.

15 Intermediates **E175B-Diastereomers C & D**. A sample of **E175A-Enantiomer B** was subjected to the Mitsunobu coupling (with *tert*-butyl 5-hydroxy-1H-indazole-1-carboxylate, WO 2016/21043), BOC deprotection, and Michael addition (with **intermediate CC**) employing the methods outlined in **Example 152** to afford a mixture of **E175B-Diastereomers C & D** (121 mg, 12 % yield over 3 steps). LCMS (ES): *m/z*
20 529.4 [M+H]⁺. A sample of **E175B-Diastereomers C & D** (121 mg) was subjected to preparative chiral HPLC purification (Column: Chiralpak OD, 21 x 250 mm, 10 micron, flow rate: 15 mL/min, mobile phase: 20% ethanol/80% heptane, detector wavelength: 220 nm, injection: 1 mL of 40 mg/mL solution) to afford **E175B-Diastereomer C** (49 mg) and **E175B-Diastereomer D** (47 mg). LCMS data for the separated diastereomers was
25 identical to the diastereomeric mixture. The relative and absolute stereochemistry of the individual diastereomers was not determined.

Example 175. Trifluoroacetic acid (0.5 mL) was added to a flask charged with a stirred solution of **E175B-Diastereomer A** (34.6 mg, 0.065 mmol) in dichloromethane (2.0 mL). The reaction vessel was placed in a 50 °C oil bath for 2 h. The reaction
30 contents were concentrated under a stream of dry nitrogen. The residue was dissolved in a mixture of 1 mL of 28-30% aqueous ammonium hydroxide solution/1 mL DMSO/1 mL

95:5 water (containing 0.5% of 30% aqueous ammonium hydroxide solution):acetonitrile. The clear solution was loaded onto a Waters Sep-Pak C18 Plus Short Cartridge, 360 mg Sorbent per cartridge, 55-105 μM particle size (WAT020515) that had been preconditioned with 10 mL of 2 M ammonia in methanol, then equilibrated with 20 mL of 95:5 water (containing 0.5% of 30% aqueous ammonium hydroxide solution):acetonitrile. After loading, the cartridge was flushed with 40 mL of 95:5 water (containing 0.5% of 30% aqueous ammonium hydroxide solution):acetonitrile at a flow rate equal to a fast drip. The cartridge was next eluted with 5 mL of 2 M ammonia in methanol at the same rate. The salt free product eluted in the first 2.5 mL of methanolic ammonia. The desired fraction was concentrated in vacuo to afford Example 175 (30.9 mg, 92 % yield). ^1H NMR (500 MHz, methanol- d_4) δ 8.76 - 8.65 (m, 2H), 8.02 - 7.95 (m, 1H), 7.57 - 7.48 (m, 1H), 7.35 - 7.29 (m, 1H), 6.96 - 6.88 (m, 2H), 6.53 - 6.47 (m, 1H), 6.32 - 6.23 (m, 1H), 4.09 - 3.99 (m, 1H), 3.96 - 3.88 (m, 1H), 3.63 - 3.48 (m, 2H), 3.25 - 3.15 (m, 1H), 2.95 - 2.87 (m, 2H), 2.76 - 2.64 (m, 2H), 2.64 - 2.58 (m, 3H), 1.98 - 1.86 (m, 1H), 1.49 - 1.37 (m, 1H), 1.26 - 1.19 (m, 1H). LC/MS (m/z) = 473.2 ($\text{M}+\text{H}$) $^+$. Human $\alpha\text{V}\beta 6$ IC_{50} (nM) = 6,000.

Example 176. A sample of **E175B-Diastereomer B** (37.5 mg, 0.071 mmol) was subjected to the deprotection and desalting methods outlined for **Example 175** to afford **Example 176** (33.5 mg, 92 % yield). ^1H NMR (500 MHz, methanol- d_4) δ 8.80 - 8.63 (m, 2H), 8.04 - 7.95 (m, 1H), 7.61 - 7.51 (m, 1H), 7.38 - 7.29 (m, 1H), 7.00 - 6.88 (m, 2H), 6.57 - 6.47 (m, 1H), 6.34 - 6.23 (m, 1H), 4.12 - 4.01 (m, 1H), 4.00 - 3.90 (m, 1H), 3.65 - 3.51 (m, 2H), 3.27 - 3.18 (m, 1H), 2.97 - 2.89 (m, 2H), 2.80 - 2.67 (m, 2H), 2.66 - 2.60 (m, 3H), 1.98 - 1.89 (m, 1H), 1.54 - 1.41 (m, 1H), 1.29 - 1.23 (m, 3H). LC/MS (m/z) = 473.2 ($\text{M}+\text{H}$) $^+$. Human $\alpha\text{V}\beta 6$ IC_{50} (nM) = 33.

Example 177. A sample of **E175B-Diastereomer C** (39 mg, 0.074 mmol) was subjected to the deprotection and desalting methods outlined for **Example 175** to afford **Example 177** (32.9 mg, 93 % yield). ^1H NMR (500 MHz, methanol- d_4) δ 8.75 - 8.64 (m, 2H), 8.02 - 7.92 (m, 1H), 7.58 - 7.51 (m, 1H), 7.33 - 7.27 (m, 1H), 6.98 - 6.91 (m, 2H), 6.54 - 6.47 (m, 1H), 6.32 - 6.22 (m, 1H), 4.13 - 4.02 (m, 1H), 4.02 - 3.92 (m, 1H), 3.58 - 3.47 (m, 2H), 3.26 - 3.15 (m, 1H), 2.97 - 2.87 (m, 2H), 2.74 - 2.65 (m, 2H), 2.62 (s, 3H),

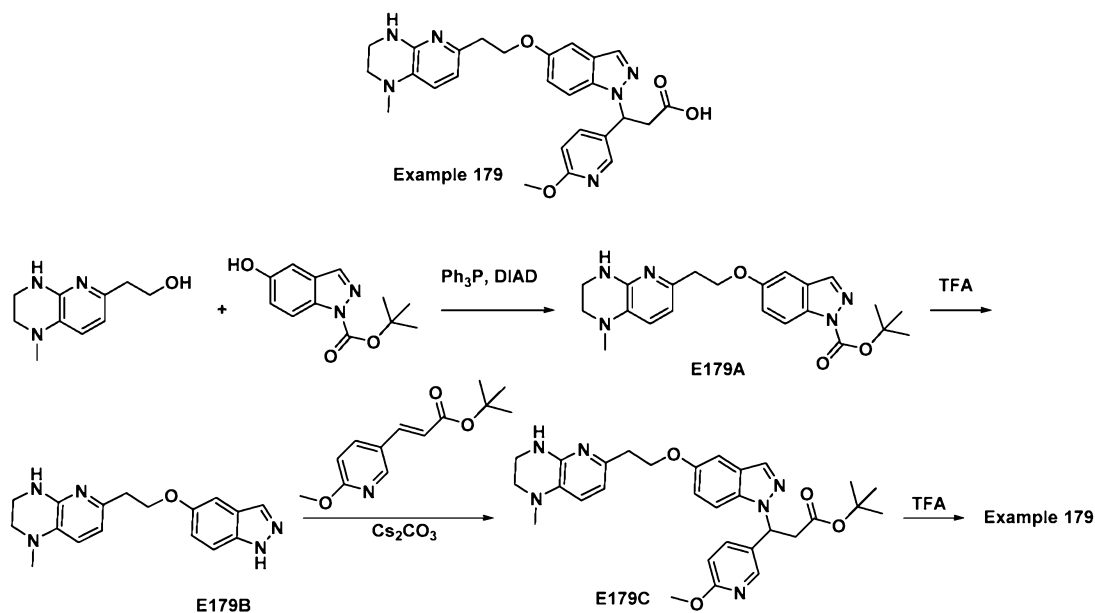
1.97 - 1.88 (m, 1H), 1.45 (dtd, $J=13.1, 9.4, 6.0$ Hz, 1H), 1.27 - 1.21 (m, 3H). LC/MS (m/z) = 473.2 ($M+H$)⁺. Human $\alpha V\beta 6$ IC₅₀ (nM) = 5,000.

Example 178. A sample of **E175B-Diastereomer D** (37 mg, 0.070 mmol) was subjected to the deprotection and desalting methods outlined for **Example 175** to afford

Example 178 (31.9 mg, 95 % yield). ¹H NMR (500 MHz, methanol-*d*₄) δ 8.77 - 8.66 (m, 2H), 8.02 - 7.95 (m, 1H), 7.58 - 7.49 (m, 1H), 7.36 - 7.28 (m, 1H), 6.99 - 6.91 (m, 2H), 6.54 - 6.48 (m, 1H), 6.31 - 6.23 (m, 1H), 4.11 - 4.01 (m, 1H), 4.00 - 3.92 (m, 1H), 3.62 - 3.50 (m, 2H), 3.25 - 3.18 (m, 1H), 2.97 - 2.87 (m, 2H), 2.76 - 2.66 (m, 2H), 1.97 - 1.87 (m, 1H), 1.51 - 1.38 (m, 1H), 1.27 - 1.20 (m, 3H). LC/MS (m/z) = 473.2 ($M+H$)⁺.
Human $\alpha V\beta 6$ IC₅₀ (nM) = 110.

Example 179

3-(6-Methoxypyridin-3-yl)-3-(5-(2-(1-methyl-1,2,3,4-tetrahydropyrido[2,3-b]pyrazin-6-yl)ethoxy)-1H-indazol-1-yl)propanoic acid



Intermediate **E179A**: To a solution of 2-(1-methyl-1,2,3,4-tetrahydropyrido[2,3-b]pyrazin-6-yl)ethan-1-ol [(US 2004/0092538) 50 mg, 0.259 mmol], *tert*-butyl 5-hydroxy-1H-indazole-1-carboxylate [(WO 2016/21043), 60.6 mg, 0.259 mmol] and Ph₃P (71.3 mg, 0.272 mmol) in THF (2270 μ l) was added DIAD (52.8 μ l, 0.272 mmol); the

reaction was stirred at rt overnight. The reaction was diluted with aqueous saturated NaHCO₃ solution. The resulting mixture extracted 3 times with EtOAc. The combine organic layers were washed with water, then with brine, and dried over sodium sulfate. The mixture was filtered and concentrated under reduced pressure. The crude product
5 was purified by silica gel column chromatography using a gradient of 20-100% EtOAc/hexanes). The fractions containing the desired product were collected and the solvent was removed under vacuo to afford **E179A** (52 mg, 49 % yield) which was contaminated with a small amount of triphenylphosphine oxide. The material was carried on without further purification. LCMS (ES): m/z 410.0 [M+H]⁺.

10 Intermediate **E179B**: To a solution of intermediate **E179A** (52 mg, 0.127 mmol) in DCM (2 mL) was added TFA (0.5 mL, 6.5 mmol) and the mixture was stirred at rt for 4 hrs. The reaction mixture was concentrated in vacuo to afford **E179B**, TFA salt (30 mg, 52 % yield). LCMS (ES): m/z 310.0 [M+H]⁺.

Intermediate **E179C**: To a solution of Intermediate **E179B**, TFA salt (30 mg,
15 0.071 mmol) in acetonitrile (0.6 mL) was added cesium carbonate (69.3 mg, 0.213 mmol). After stirring at rt for 5 min, *tert*-butyl (*E*)-3-(6-methoxypyridin-3-yl)acrylate [(*J. Org. Chem.* **2004**, *69*, 1959), 25.1 mg, 0.106 mmol] was added and the resulting mixture was stirred at 80 °C for 16 hrs. The mixture was cooled to rt, filtered, and concentrated. The residue was purified by silica-gel column chromatography (ISCO column, 40g, 30-
20 100% EtOAc/hexanes). The pure fractions were concentrated in vacuo to afford **E179C** (30 mg, 0.055 mmol, 78 % yield). LCMS (ES): m/z 545.2 [M+H]⁺.

Example 179. Trifluoroacetic acid (1 mL) was added to a solution containing **E179C** (40 mg, 0.073 mmol) in dichloromethane (2 mL). The resulting mixture was stirred at rt for 4 h. The mixture was concentrated in vacuo. The residue was dissolved
25 in methanol and purified via preparative LC/MS with the following conditions: Column: XBridge C18, 19 x 200 mm, 5-μm particles; Mobile Phase A: 5:95 acetonitrile: water with 10-mM ammonium acetate; Mobile Phase B: 95:5 acetonitrile: water with 10-mM ammonium acetate; Gradient: 10-100% B over 17 minutes, then a 5-minute hold at 100% B; Flow: 20 mL/min. Fractions containing the desired product were combined and dried
30 via centrifugal evaporation to afford example **179** (29.4 mg, 0.062 mmol, 91 % yield).
¹H NMR (500 MHz, DMSO-*d*₆) δ 8.28 - 8.22 (m, 1H), 8.00 - 7.92 (m, 1H), 7.74 - 7.68

(m, 1H), 7.68 - 7.63 (m, 1H), 7.20 - 7.13 (m, 1H), 7.01 - 6.96 (m, 1H), 6.75 - 6.70 (m, 1H), 6.58 - 6.52 (m, 1H), 6.40 - 6.31 (m, 2H), 6.22 - 6.12 (m, 1H), 4.23 - 4.14 (m, 2H), 3.79 - 3.74 (m, 3H), 3.66 - 3.53 (m, 1H), 3.46 - 3.37 (m, 3H), 3.23 - 3.13 (m, 1H), 3.09 - 3.02 (m, 2H), 2.87 - 2.80 (m, 2H), 2.75 - 2.68 (m, 3H). LC/MS (m/z) = 489.1 ($M+H$)⁺.

5 Human $\alpha V\beta 6$ IC₅₀ (nM) = 47.

Examples 180

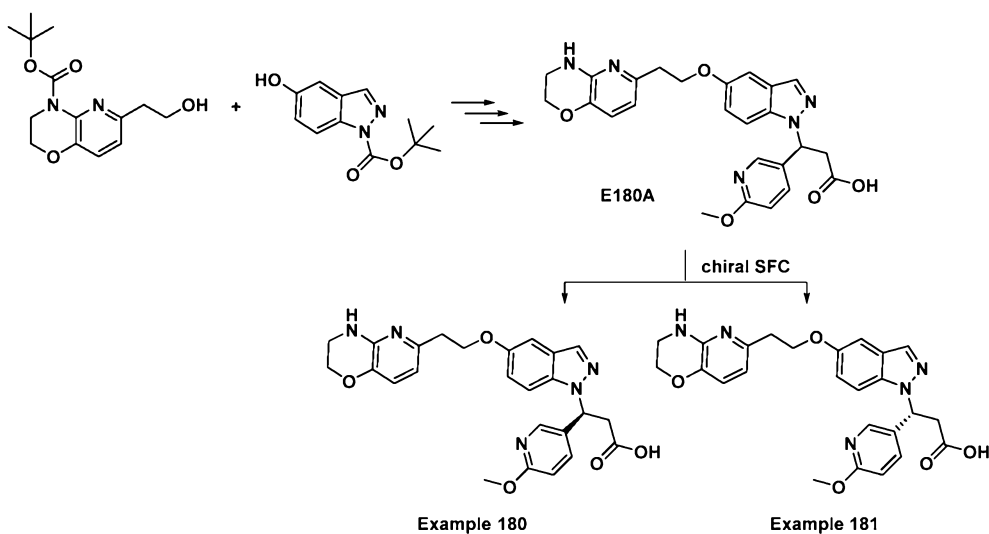
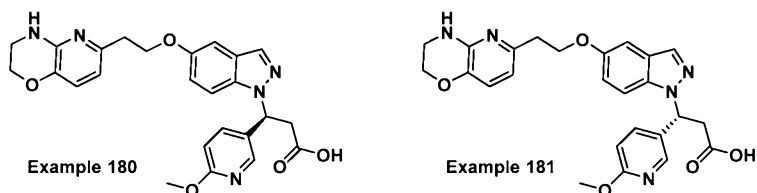
(*S*)-3-(5-(2-(3,4-dihydro-2H-pyrido[3,2-b][1,4]oxazin-6-yl)ethoxy)-1H-indazol-1-yl)-3-(6-methoxypyridin-3-yl)propanoic acid;

10 and

Example 181

(*R*)-3-(5-(2-(3,4-dihydro-2H-pyrido[3,2-b][1,4]oxazin-6-yl)ethoxy)-1H-indazol-1-yl)-3-(6-methoxypyridin-3-yl)propanoic acid

15



Intermediate **E180A**: A sample of *tert*-butyl 6-(2-hydroxyethyl)-2,3-dihydro-4H-pyrido[3,2-b][1,4]oxazine-4-carboxylate (*Bioorg. Med. Chem. Lett.* **2005**, *15*, 2679) was converted in 4 steps using the methods outlined in Example 3 into **E173A**. LCMS (ES): m/z 476.0 $[M+H]^+$.

5 **Examples 180 and Example 181.** A sample of **E180A** (143 mg) was subjected to chiral SFC purification (Column: Chiralpak OJ-H, 30 x 250 mm, 5 micron, BPR pressure: 150 bar, temperature 35 °C, flow rate: 70.0 mL/min, mobile phase: 20% MeOH w/0.1% NH₄OH in CO₂, detector wavelength: 254 nm, stacked injections: 0.5 mL of 24 mg/mL solution) to afford **Example 180** (29 mg) and **E181** (32 mg).

10 Data for **Example 180**: ¹H NMR (400 MHz, DMSO-*d*₆) δ 8.23 (d, J =2.3 Hz, 1H), 7.96 (s, 1H), 7.72 - 7.67 (m, 1H), 7.67 - 7.62 (m, 1H), 7.17 - 7.15 (m, 1H), 7.00 - 6.95 (m, 1H), 6.84 (d, J =7.8 Hz, 1H), 6.71 (d, J =8.5 Hz, 1H), 6.67 - 6.60 (m, 1H), 6.41 (d, J =7.8 Hz, 1H), 6.16 (dd, J =9.7, 5.4 Hz, 1H), 4.21 (br t, J =6.7 Hz, 2H), 4.13 - 3.95 (m, 2H), 3.77 (s, 3H), 3.56 (br dd, J =16.2, 9.7 Hz, 1H), 3.35 (br s, 1H), 3.16 (br dd, J =16.6, 5.3 Hz, 1H), 2.90 (t, J =6.8 Hz, 2H). LC/MS (m/z) = 476.1 (M+H)⁺. Human α V β 6 IC₅₀ (nM) = 15.

20 Data for **Example 181**: ¹H NMR (400 MHz, DMSO-*d*₆) δ 8.24 (d, J =2.3 Hz, 1H), 7.96 (s, 1H), 7.69 (d, J =9.0 Hz, 1H), 7.65 (dd, J =8.5, 2.5 Hz, 1H), 7.16 (d, J =2.3 Hz, 1H), 6.98 (dd, J =9.0, 2.3 Hz, 1H), 6.84 (d, J =7.8 Hz, 1H), 6.71 (d, J =8.5 Hz, 1H), 6.67 - 6.61 (m, 1H), 6.41 (d, J =7.8 Hz, 1H), 6.16 (dd, J =9.5, 5.3 Hz, 1H), 4.21 (br t, J =6.8 Hz, 2H), 4.09 - 4.02 (m, 2H), 3.77 (s, 3H), 3.57 (br dd, J =16.6, 9.8 Hz, 1H), 3.35 (br d, J =2.8 Hz, 2H), 3.18 (br dd, J =16.4, 5.1 Hz, 1H), 2.90 (br t, J =6.9 Hz, 2H). LC/MS (m/z) = 476.1 (M+H)⁺. Human α V β 6 IC₅₀ (nM) = 5,000.

25 BIOLOGICAL EVALUATION

All binding assays used the HTRF (homogeneous time resolved fluorescence) technology from Cisbio International, therefore all assays are described as HTRF binding assays. The assay results for the Examples are listed above together with the characterization data. The HTRF binding assays are established for the following

30 integrins: human α V β 6, human α V β 1, human α V β 3, human α V β 5, and human α V β 8. All assays used the following assay buffer: 20 mM Tris, pH 7.4, 1 mM MgCl₂, 1 mM MnCl₂,

0.01% Tween 20, and 0.01% BSA. Alternatively, a SPA-based assay was used for evaluation of receptor binding.

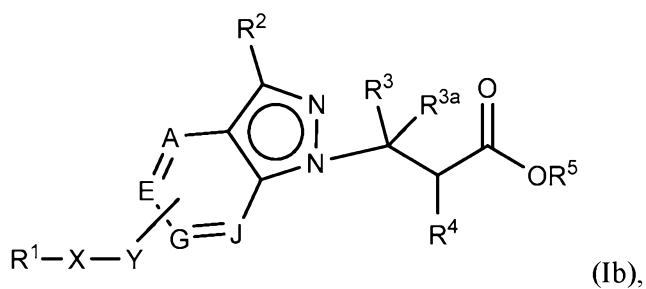
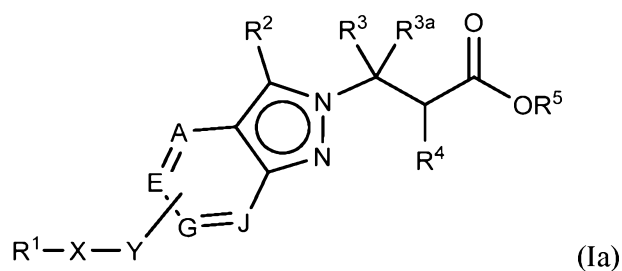
The following describes the components and a representative procedure for the human $\alpha V\beta 6$ HTRF binding assay: Recombinant human $\alpha V\beta 6$ Integrin (R & D systems, 3817-AV) was biotinylated. Biotinylated human $\alpha V\beta 6$ Integrin was added to assay vessel at a final concentration of 1.25 nM. FITC-conjugated fibronectin (Cytoskeleton, FNR02) was then added at the final concentration of 5 nM. The mixture was centrifuged at 600 rpm for three minutes using Thermo Fisher Heraeus Multifuge X3 centrifuge and then incubated at room temperature for an hour. Streptavidin Terbium (Cisbio international 610STLB) was then added at the final concentration of 0.625 nM. The resulting mixture was centrifuged at 600 rpm for three minutes using Thermo Fisher Heraeus Multifuge X3 centrifuge and then incubated at room temperature overnight in dark before reading HTRF signals.

The SPA-based assay was carried out according to the protocol and procedures similar to the ones described in the following reference with appropriate modifications to agents and ligands which are readily understood by one skilled in the art: Pachter JA, Zhang R, Mayer-Ezell R., "Scintillation proximity assay to measure binding of soluble fibronectin to antibody-captured $\alpha V\beta 1$ integrin" *Anal Biochem.* **1995** Sep 1;230(1):101-7.

Other features of the invention should become apparent in the course of the above descriptions of exemplary embodiments that are given for illustration of the invention and are not intended to be limiting thereof. The present invention may be embodied in other specific forms without departing from the spirit or essential attributes thereof. This invention encompasses all combinations of preferred aspects of the invention noted herein. It is understood that any and all embodiments of the present invention may be taken in conjunction with any other embodiment or embodiments to describe additional embodiments. It is also understood that each individual element of the embodiments is its own independent embodiment. Furthermore, any element of an embodiment is meant to be combined with any and all other elements from any embodiment to describe an additional embodiment.

WHAT IS CLAIMED IS:

1. A compound of Formula (Ia) or (Ib):



wherein:

A, E, G, and J are independently N, C or CH with the proviso that at least one of A, E, G, and J is C attached to Y;

L^1 and L^2 are each independently C_{1-4} alkylene;

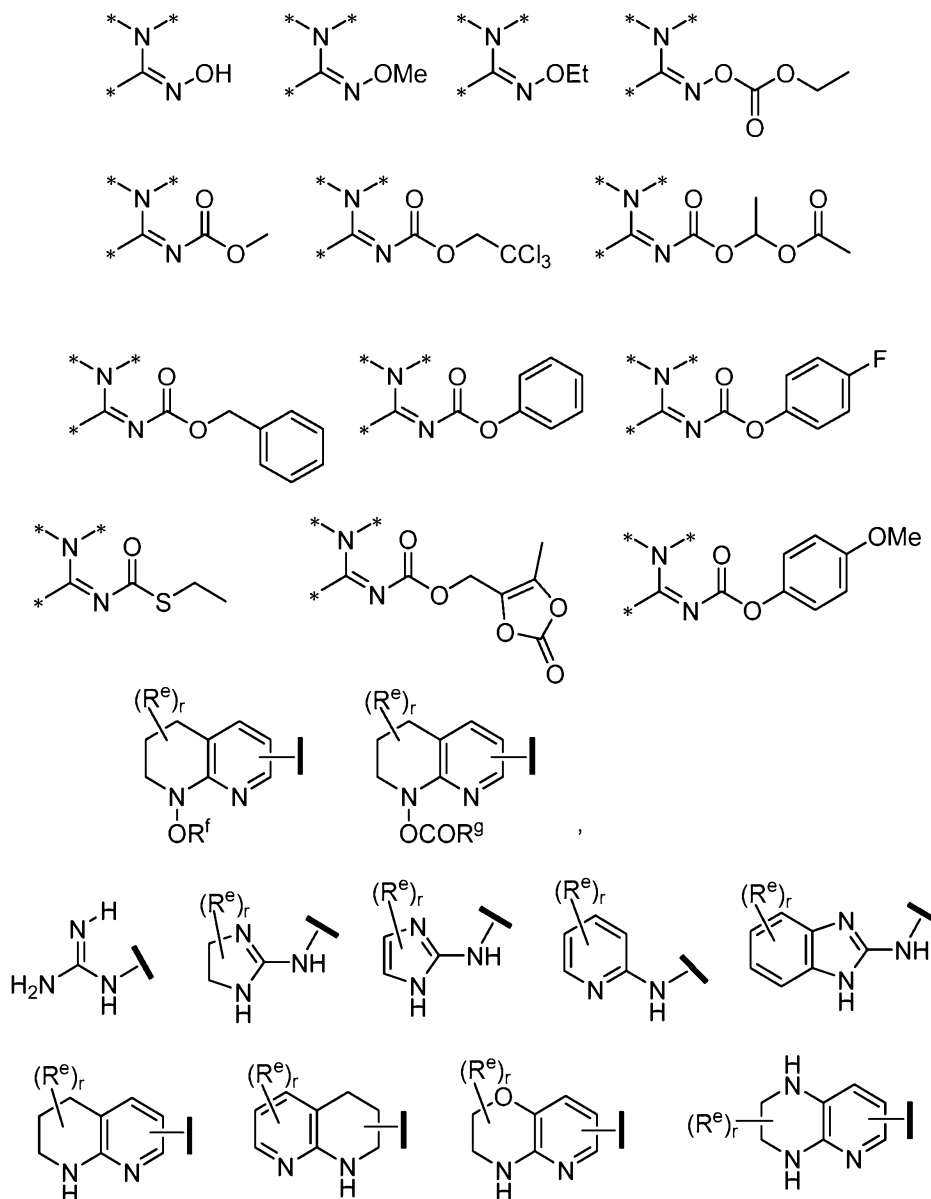
X is a C_{1-4} alkylene substituted with 0, 1, or 2 R^{8a} ;

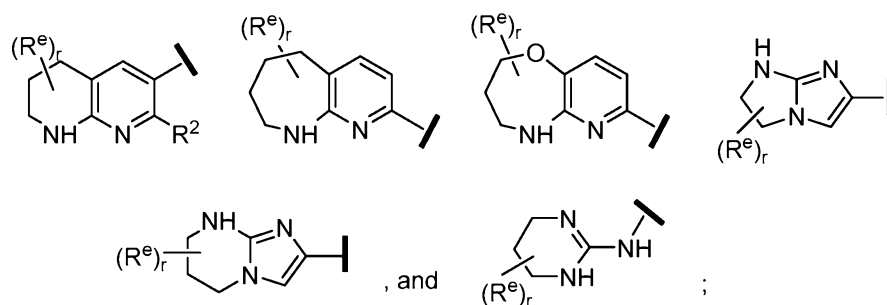
Y is a covalent bond, O, S, NH, -O-(C_{1-3} alkylene)-, -S-(C_{1-3} alkylene)-, or -NH-(C_{1-3} alkylene)-, wherein the C_{1-3} alkylene is each independently substituted with 0, 1, or 2 R^{8b} ;

m is an integer of 1 or 2;

r is an integer of 0, 1, 2, or 3;

R¹ is an Arginine mimetic moiety selected from the group consisting of





wherein one of the asterisks in each of the arginine mimetics moiety is an attachment point to X and the other two asterisks are hydrogen;

R^2 is hydrogen, halo, or C_{1-6} alkyl;

R^3 is hydrogen, C_{1-6} alkyl, 3- to 10-membered carbocyclyl, carbocyclylalkyl, 6- to 10-membered aryl, arylalkyl, 3- to 14-membered heterocyclyl, heterocyclylalkyl, 5- to 14-membered heteroaryl, or heteroarylalkyl, wherein the alkyl, carbocyclyl, heterocyclyl, aryl, and heteroaryl, by themselves or as part of another group, are each independently substituted with 0, 1, 2, or 3 R^6 ;

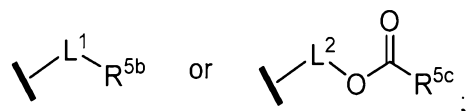
R^{3a} is hydrogen;

or R^{3a} and R^3 taken together with the atom to which they are attached, form a 3- to 6-membered carbocyclic or heterocyclic ring which is optionally substituted with one or more groups independently selected from halo, cyano, hydroxyl, amino, C_{1-6} alkyl, haloalkyl, hydroxyalkyl, aminoalkyl, alkoxy, haloalkoxy, amido, carbamate, and sulfonamide;

R^4 is hydrogen, C_{1-6} alkyl, 3- to 10-membered carbocyclyl, carbocyclylalkyl, 3- to 10-membered heterocyclyl, heterocyclylalkyl, 6- to 10-membered aryl, arylalkyl, 5- to 14-membered heteroaryl, heteroarylalkyl, $-S(O)_mR^7$, $-C(O)NR^aR^b$, $-NHC(O)OR^a$, $-NHC(O)NR^aR^b$, $-NHC(O)R^7$, $-OC(O)NR^aR^b$, $-OC(O)R^7$, $-NHS(O)_mNR^aR^b$, or $-NHS(O)_mR^7$; wherein the alkyl, carbocyclyl, heterocyclyl, aryl, and heteroaryl, by

themselves or as part of another group, are each independently substituted with 0, 1, 2, or 3 R^9 ;

R^5 is hydrogen, R^{5a} , or a structural moiety selected from



R^{5a} and R^{5b} are each independently C_{1-6} alkyl, phenyl, benzyl, or 5- to 7-membered heterocyclyl; wherein the alkyl, phenyl, and heterocyclyl are each independently substituted with 0 to 3 R^{5d} ;

R^{5c} is C_{1-6} alkyl or 5- to 7-membered carbocyclyl; wherein the C_{1-6} alkyl, phenyl, and heterocyclyl are each independently substituted with 0 to 3 R^{5d} ; and

R^{5d} , at each occurrence, is independently halo, OH, alkoxy, oxo, or alkyl; or alternatively, two adjacent R^{5d} , together with the atoms to which they are attached, form a carbocyclyl moiety.

R^6 is halo, cyano, hydroxyl, amino, oxo, nitro, $-S(O)_mR^{12}$, C_{1-6} alkyl, alkoxy, haloalkyl, haloalkoxy, haloaminoalkyl, hydroxyalkyl, aminoalkyl, alkoxy carbonyl, 6- to 10-membered aryl, aryloxy, arylalkoxy, 5- to 10-membered heteroaryl, 3- to 6-membered carbocyclyl, or 3- to 7-membered heterocyclyl; wherein the alkyl, aryl, heteroaryl, carbocyclyl, or heterocyclyl, by themselves or as part of another group, are each independently substituted with 0, 1, or 2 R^{10} ;

R^7 is each independently C_{1-6} alkyl, C_{1-6} alkoxy, C_{1-6} aminoalkyl, C_{1-6} haloalkyl, 6- to 10-membered aryl, arylalkyl, 5- to 10-membered heteroaryl, cycloalkyl, or heterocycloalkyl; wherein the alkyl, aryl, heteroaryl, cycloalkyl, and heterocycloalkyl, by themselves or as part of another group, are each independently substituted with 0, 1, 2, or 3 R^{11} ;

R^{8a} and R^{8b}, at each occurrence, are independently halo, cyano, hydroxyl, amino, oxo, C₁₋₆ alkyl, haloalkyl, hydroxyalkyl, aminoalkyl, alkoxy, or haloalkoxy;

R⁹ is each independently halo, cyano, hydroxyl, amino, oxo, nitro, C₁₋₆ alkyl, alkoxy, haloalkyl, haloalkoxy, haloaminoalkyl, hydroxyalkyl, aminoalkyl, alkoxycarbonyl, 6- to 10-membered aryl, aryloxy, arylalkoxy, 5- to 10-membered heteroaryl, 3- to 6-membered carbocyclyl, or 3- to 7-membered heterocyclyl; wherein the alkyl, aryl, heteroaryl, carbocyclyl, or heterocyclyl, by themselves or as part of another group, are each independently substituted with 0, 1, or 2 R¹³;

R¹⁰ is halo, cyano, hydroxyl, amino, oxo, C₁₋₆ alkyl, haloalkyl, hydroxyalkyl, aminoalkyl, alkoxy, haloalkoxy, amido, carbamate, or sulfonamide;

R¹¹, at each occurrence, is independently halo, cyano, hydroxyl, amino, oxo, C₁₋₆ alkyl, haloalkyl, hydroxyalkyl, aminoalkyl, alkoxy, or haloalkoxy; R¹² is -N(R^xR^y), C₁₋₆ alkyl, C₁₋₆ haloalkyl, C₁₋₆ alkoxy, C₁₋₆ haloalkoxy, C₁₋₆ hydroxyalkyl, or C₁₋₆ aminoalkyl;

R¹³ is halo, cyano, hydroxyl, amino, oxo, C₁₋₆ alkyl, haloalkyl, hydroxyalkyl, aminoalkyl, alkoxy, haloalkoxy, amido, carbamate, or sulfonamide;

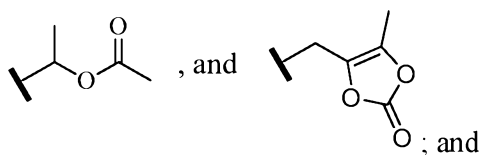
R^a and R^b, at each occurrence, are independently hydrogen, C₁₋₆ alkyl, C₁₋₆ haloalkyl, C₃₋₆ cycloalkyl, 6- to 10-membered aryl, 5- to 10-membered heteroaryl, cycloalkylalkyl, arylalkyl, heteroarylalkyl, or alkoxyalkyl; or alternatively, R^a and R^b, taken together with the atoms to which they are attached, form a 3- to 8-membered carbocyclic or heterocyclic ring; wherein the aryl and heteroaryl, by themselves or as part of another group, are each independently substituted with one or more groups independently selected from halo, cyano, hydroxyl, amino, C₁₋₆ alkyl, haloalkyl, hydroxyalkyl, aminoalkyl, alkoxy, haloalkoxy, amido, carbamate, and sulfonamide; and the carbocyclyl and heterocyclyl, by themselves or as part of another group, are each independently

substituted with one or more groups independently selected from halo, cyano, hydroxyl, amino, oxo, C₁₋₆ alkyl, haloalkyl, hydroxyalkyl, aminoalkyl, alkoxy, haloalkoxy, amido, carbamate, and sulfonamide;

R^e is OH, amino, amido, carbamate, sulfonamide, C₁₋₄ alkyl, halo, C₁₋₄ haloalkyl, or C₃₋₆ cycloalkyl;

R^f is H, CH₃, CH₂CH₃, or C(O)OCH₂CH₃;

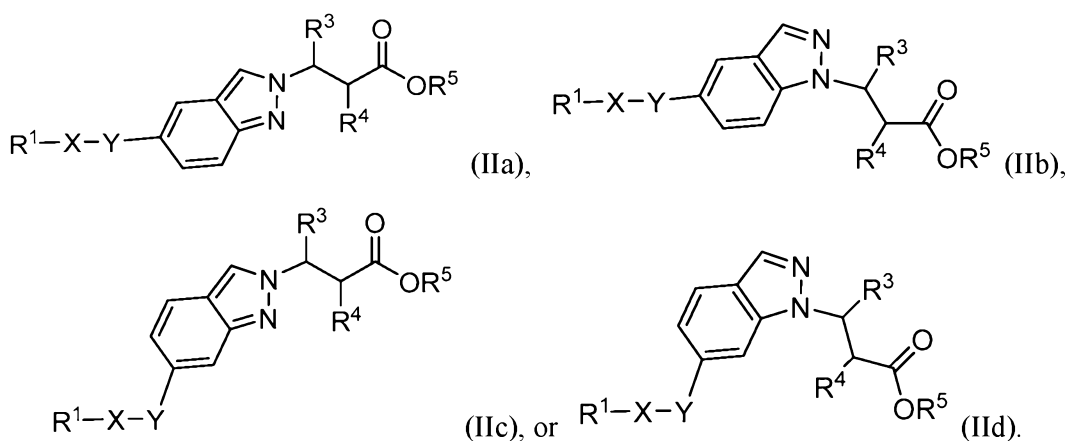
R^g is selected from CH₃, CH₂CCl₃, phenyl, 4-fluorophenyl, 4-methoxyphenyl, benzyl,



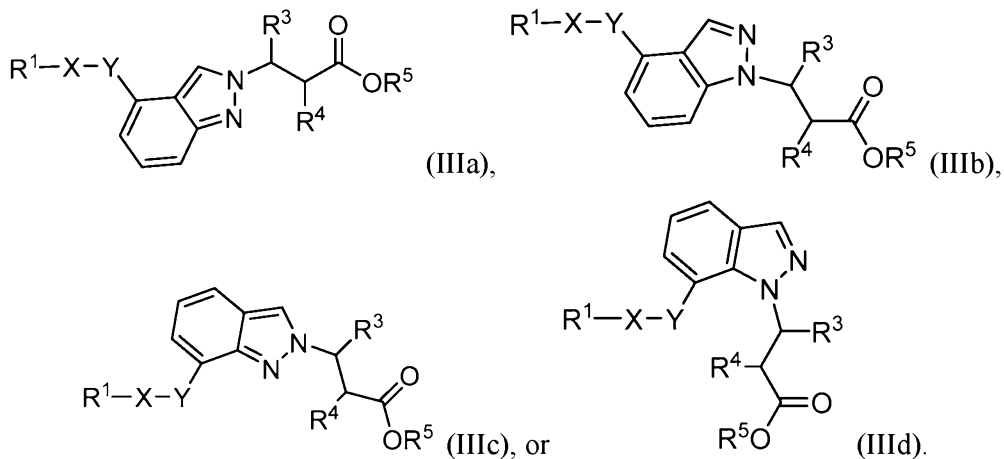
R^x and R^y are each independently hydrogen or C₁₋₆ alkyl;

or a pharmaceutically acceptable salt thereof.

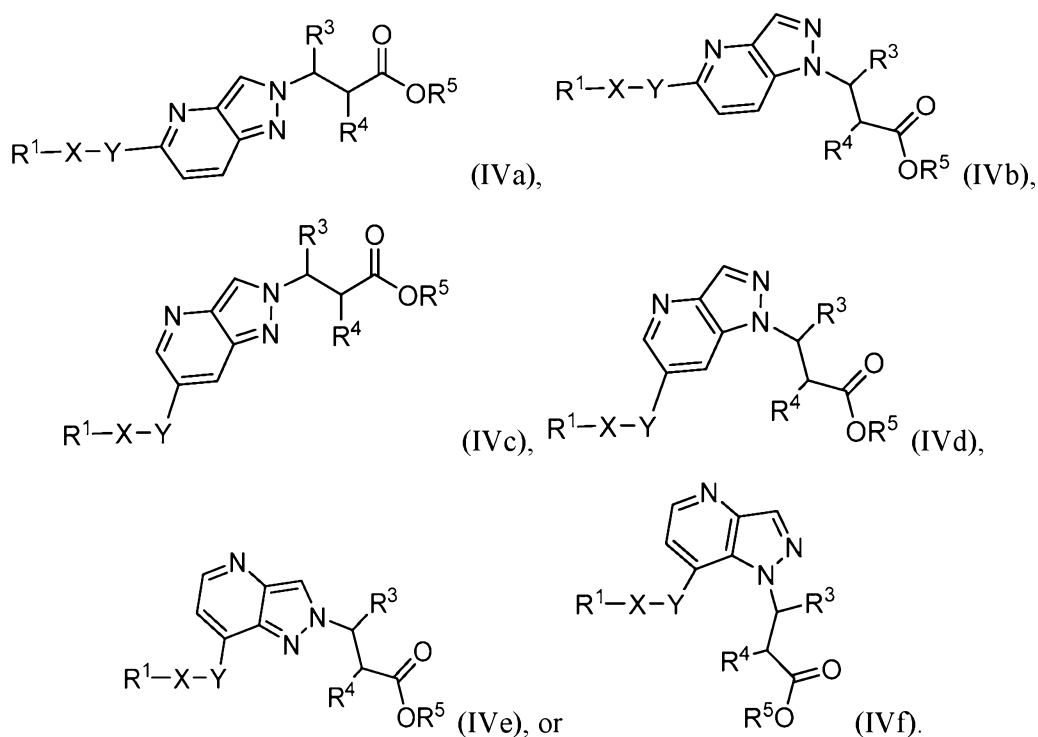
2. A compound of claim 1 wherein the compound of Formula (Ia) or (Ib) is represented by structural Formula (IIa), (IIb), (IIc), or (IId):



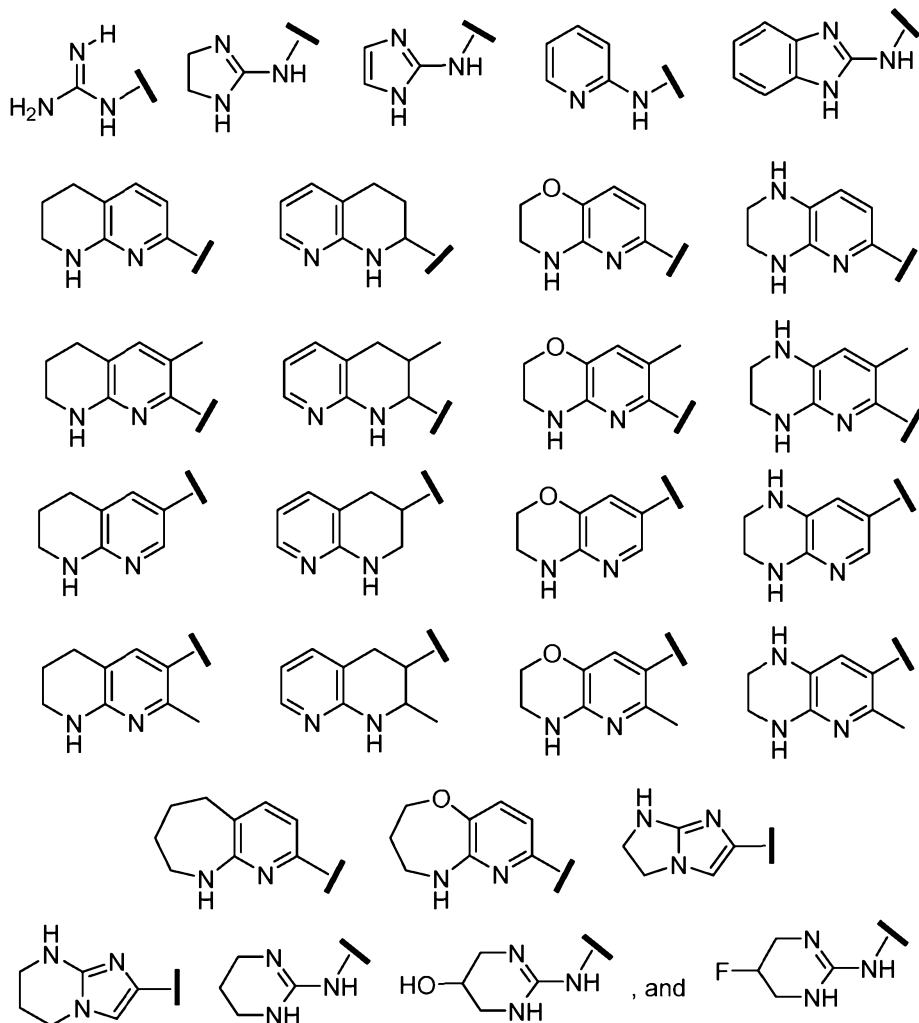
3. A compound of claim 1 wherein the compound of Formula (Ia) or (Ib) is represented by structural Formula (IIIa), (IIIb), (IIIc), or (IIId):



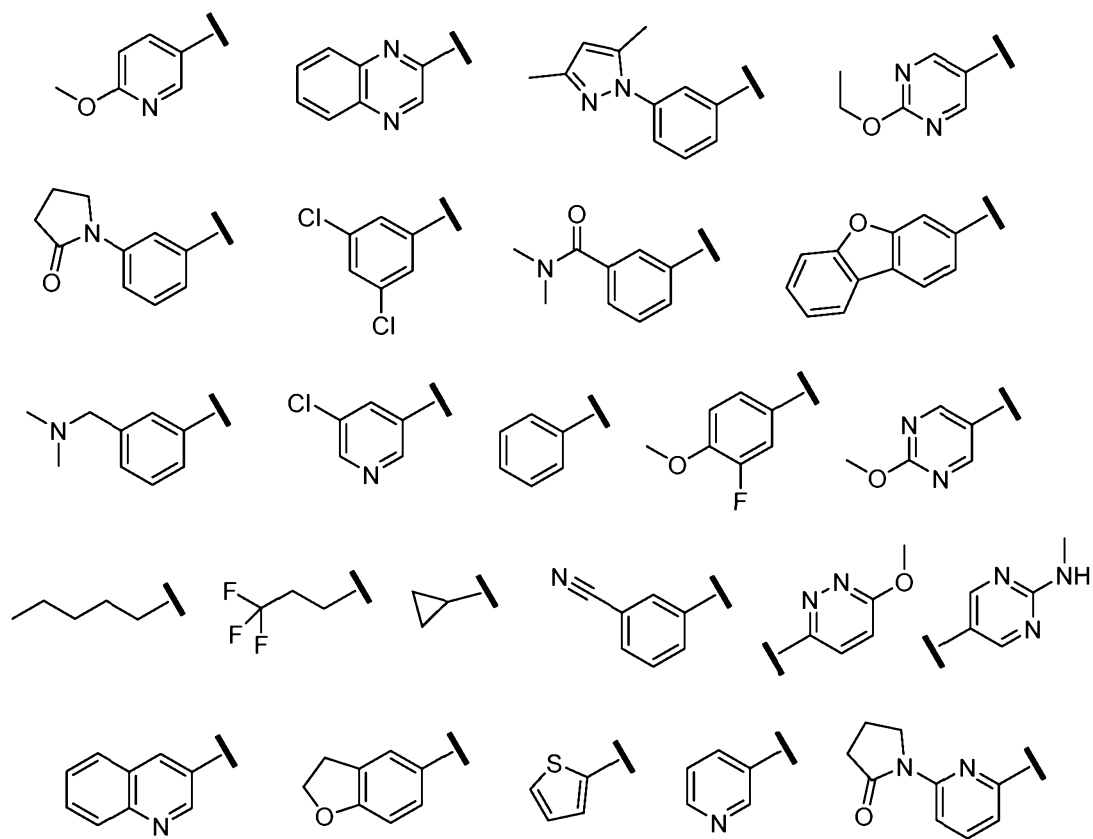
4. A compound of claim 1 wherein the compound of Formula (Ia) or (Ib) is represented by structural Formula (IVa), (IVb), (IVc), (IVd), (IVe) or (IVf):

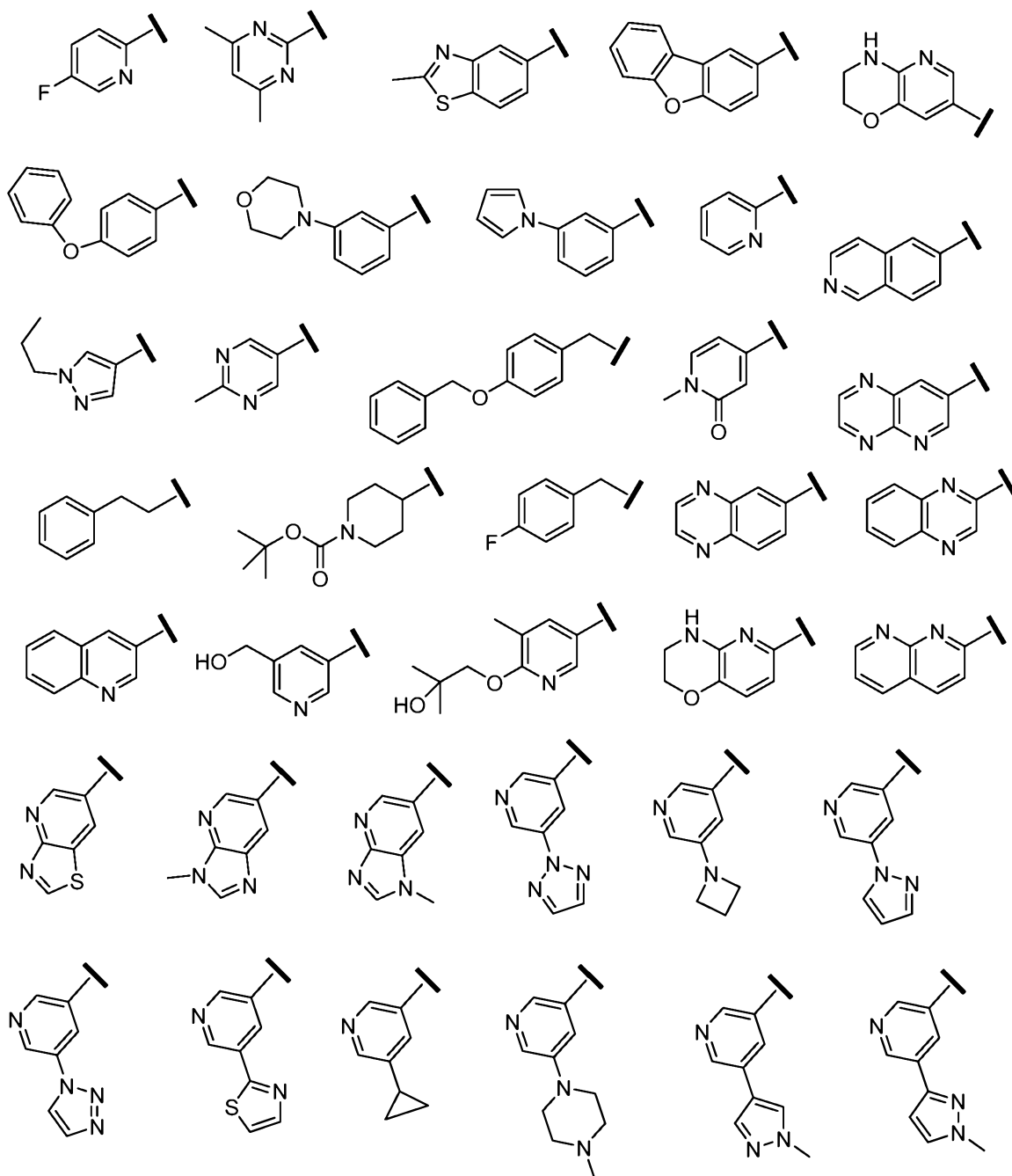


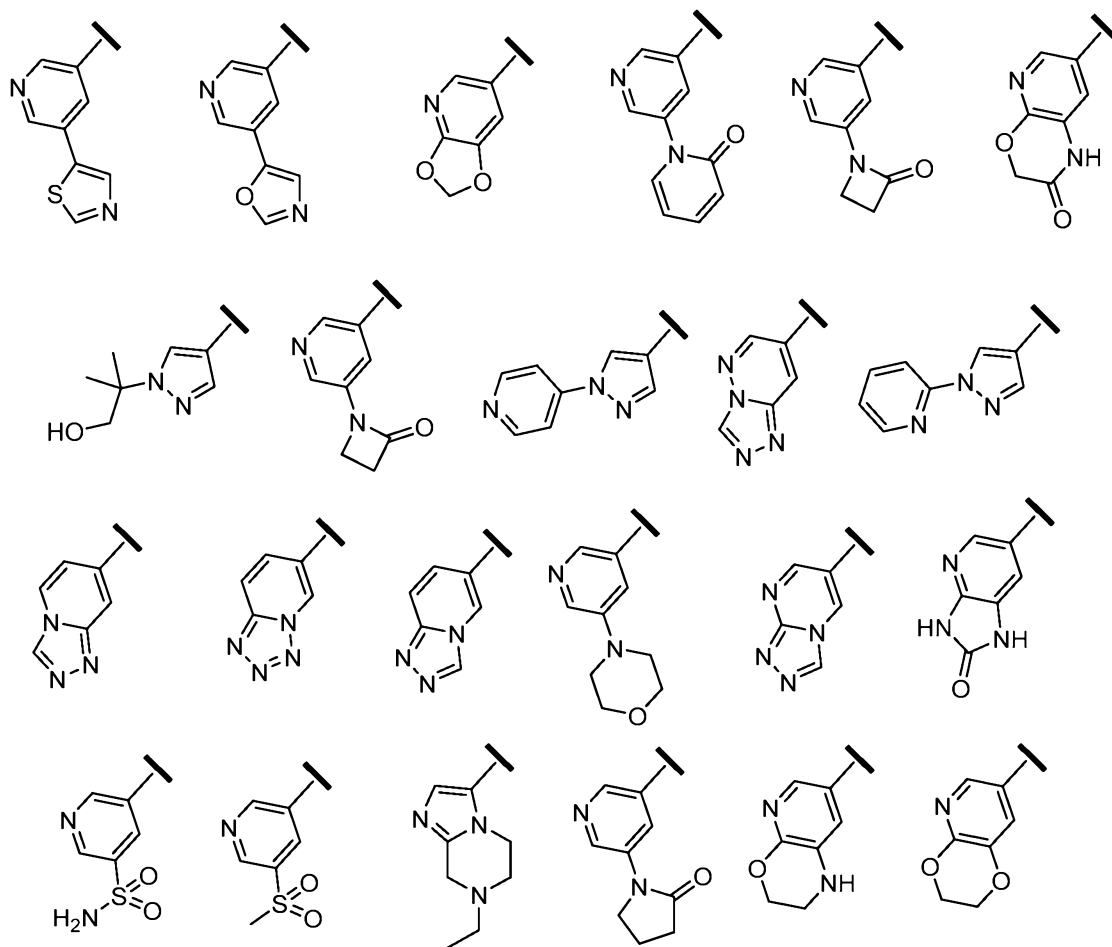
5. A compound of claim 1 wherein R¹ is selected from the group consisting of



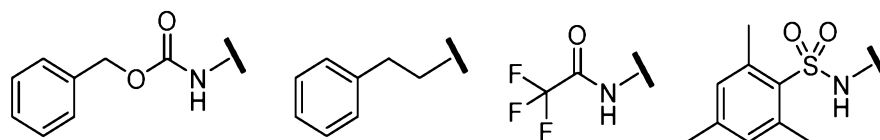
6. A compound of claim 1 wherein R^{3a} is hydrogen; and R³ is hydrogen or a structural moiety selected from the group consisting of



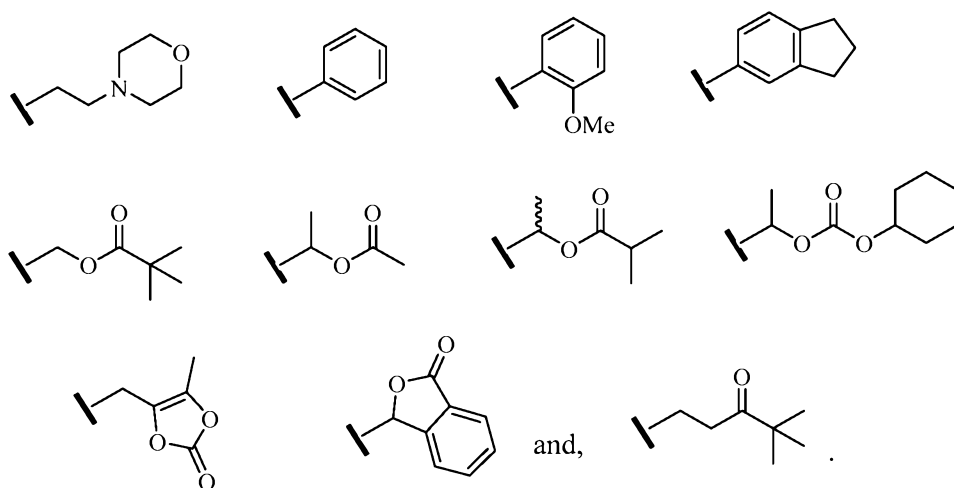




7. A compound of claim 1 wherein R^4 is hydrogen or a structural moiety selected from the group consisting of



8. A compound of claim 1 wherein R^5 is H or R^{5a} , and R^{5a} is methyl, ethyl, isopropyl, *n*-butyl, isopentyl, or a structural moiety selected from



9. A compound of claim 1 wherein X is C₁₋₄ alkylene; and Y is a covalent bond or O.

10. A compound of claim 1 selected from the group consisting of:

3-(6-Methoxypyridin-3-yl)-3-(4-(3-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)propyl)-1H-indazol-1-yl)propanoic acid;

3-(6-Methoxypyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethyl)-1H-indazol-1-yl)propanoic acid;

3-(6-Methoxypyridin-3-yl)-3-(4-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid;

(S)-2-(((Benzyloxy)carbonyl)amino)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-2H-indazol-2-yl)propanoic acid;

3-Phenyl-3-(5-(3-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)propoxy)-1H-indazol-1-yl)propanoic acid;

(*S*)-3-(6-Methoxypyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-2*H*-indazol-2-yl)propanoic acid;

(*S*)-3-(6-Methoxypyridin-3-yl)-3-(6-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethyl)-2*H*-indazol-2-yl)propanoic acid;

(*S*)-3-(6-Methoxypyridin-3-yl)-3-(6-((2-methyl-5,6,7,8-tetrahydro-1,8-naphthyridin-3-yl)methyl)-2*H*-indazol-2-yl)propanoic acid;

(*S*)-3-(6-Methoxypyridin-3-yl)-3-(6-(3-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)propyl)-2*H*-indazol-2-yl)propanoic acid;

(*S*)-3-(6-Methoxypyridin-3-yl)-3-(6-(2-(2-methyl-5,6,7,8-tetrahydro-1,8-naphthyridin-3-yl)ethyl)-2*H*-indazol-2-yl)propanoic acid;

3-(6-Methoxypyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)pyrazolo[4,3-*b*]pyridin-1-yl)propanoic acid;

3-(5-(2-((4,5-Dihydroimidazol-2-yl)amino)ethoxy)-1*H*-indazol-1-yl)-3-(6-methoxypyridin-3-yl)propanoic acid;

3-(5-(2-(5,6,7,8-Tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)-2-((2,4,6-trimethylphenyl)sulfonamido)propanoic acid;

2-(((Benzyloxy)carbonyl)amino)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

3-(Quinoxalin-2-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

(*R*)-3-(Quinoxalin-2-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy))-1*H*-indazol-1-yl)propanoic acid;

(*S*)-3-(Quinoxalin-2-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy))-1*H*-indazol-1-yl)propanoic acid;

3-(3,5-Dichlorophenyl)-3-(4-(3-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)propyl))-1*H*-indazol-1-yl)propanoic acid;

3-(Quinoxalin-2-yl)-3-(4-(3-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)propyl))-1*H*-indazol-1-yl)propanoic acid;

3-(6-Methoxypyridin-3-yl)-3-(5-(3-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)propyl))-1*H*-indazol-1-yl)propanoic acid;

3-(3,5-Dichlorophenyl)-3-(5-(3-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)propyl))-1*H*-indazol-1-yl)propanoic acid;

3-(Quinoxalin-2-yl)-3-(5-(3-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)propyl))-1*H*-indazol-1-yl)propanoic acid;

3-(3-(Dimethylcarbamoyl)phenyl)-3-(5-(3-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)propyl))-1*H*-indazol-1-yl)propanoic acid;

3-(3-(Dimethylcarbamoyl)phenyl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethyl))-1*H*-indazol-1-yl)propanoic acid;

3-(Dibenzo[*b,d*]furan-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethyl))-1*H*-indazol-1-yl)propanoic acid;

3-(3-((Dimethylamino)methyl)phenyl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethyl)-1*H*-indazol-1-yl)propanoic acid;

3-(Quinoxalin-2-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethyl)-1*H*-indazol-1-yl)propanoic acid;

3-(3,5-Dichlorophenyl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethyl)-1*H*-indazol-1-yl)propanoic acid;

3-(3-(Dimethylcarbamoyl)phenyl)-3-(4-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

3-(Dibenzo[*b,d*]furan-3-yl)-3-(4-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

6,6,6-Trifluoro-3-(4-(3-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)propyl)-1*H*-indazol-1-yl)hexanoic acid;

3-(3,5-Dichlorophenyl)-3-(4-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

3-(Quinoxalin-2-yl)-3-(4-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

3-(6-Methoxypyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

3-(3-(Dimethylcarbamoyl)phenyl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

(*S*)-2-(((Benzyloxy)carbonyl)amino)-3-(5-(3-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)propyl)-1*H*-indazol-1-yl)propanoic acid;

(*R*)-3-(3-(Dimethylcarbamoyl)phenyl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

(*S*)-3-(3-(Dimethylcarbamoyl)phenyl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

2-(((Benzyloxy)carbonyl)amino)-3-(4-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

(*R*)-3-(6-Methoxypyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

(*S*)-3-(6-Methoxypyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

3-(5-Chloropyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

3-(3,5-Dichlorophenyl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

3-(3-Fluoro-4-methoxyphenyl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

3-Cyclopropyl-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

3-(5-(2-(5,6,7,8-Tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)octanoic acid;

3-(2,3-Dihydrobenzofuran-5-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

3-(Quinolin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid (48);

3-(5-(2-(5,6,7,8-Tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)-3-(thiophen-2-yl)propanoic acid;

3-(Pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

3-(5-(2-(5,6,7,8-Tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

3-(3-Cyanophenyl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

3-(5-Fluoropyridin-2-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

3-(Dibenzo[*b,d*]furan-2-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

3-(4,6-Dimethylpyrimidin-2-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

3-(2-Methylbenzo[d]thiazol-5-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

3-(4-Phenoxyphenyl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

3-(3-Morpholinophenyl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

3-(3-(1*H*-Pyrrol-1-yl)phenyl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

3-(3-((Dimethylamino)methyl)phenyl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

3-(Pyridin-2-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

3-(3-(2-Oxopyrrolidin-1-yl)phenyl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

3-(1-Propylpyrazol-4-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

(*S*)-3-(6-Methoxypyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethyl)-2*H*-indazol-2-yl)propanoic acid;

(*S*)-3-(6-Methoxypyridin-3-yl)-3-(6-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-2*H*-indazol-2-yl)propanoic acid;

3-(2-Methylpyrimidin-5-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

(*S*)-3-(6-Methoxypyridin-3-yl)-3-(5-(3-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)propyl)-2*H*-indazol-2-yl)propanoic acid;

3-(3-(3,5-Dimethylpyrazol-1-yl)phenyl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

4-(4-(Benzyloxy)phenyl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)butanoic acid;

3-(6-Methoxypyridin-3-yl)-3-(3-methyl-5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

3-(1-Methyl-2-oxo-1,2-dihydropyridin-4-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

(3*S*)-3-(6-Methoxypyridin-3-yl)-3-(6-(2-(1,2,3,4-tetrahydro-1,8-naphthyridin-2-yl)ethyl)-2*H*-indazol-2-yl)propanoic acid;

4-Phenyl-2-((5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)methyl)butanoic acid;

3-(1-(*tert*-Butoxycarbonyl)piperidin-4-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

3-(Pyridin-4-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

2-(1-(5-(2-(5,6,7,8-Tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)cyclopropyl)acetic acid;

3-(2-Ethoxypyrimidin-5-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

4-(4-Fluorophenyl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)butanoic acid;

3-(5-Methoxypyrazin-2-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

3-(Quinoxalin-6-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

(*S*)-3-(Quinolin-3-yl)-3-(6-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethyl)-2*H*-indazol-2-yl)propanoic acid;

(*S*)-3-(6-((2-Methyl-5,6,7,8-tetrahydro-1,8-naphthyridin-3-yl)methyl)-2*H*-indazol-2-yl)-3-(quinolin-3-yl)propanoic acid;

(3*S*)-3-(Quinolin-3-yl)-3-(6-(2-(1,2,3,4-tetrahydro-1,8-naphthyridin-2-yl)ethyl)-2*H*-indazol-2-yl)propanoic acid;

(*S*)-3-(3-(2-Oxopyrrolidin-1-yl)phenyl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

(*R*)-3-(3-(2-Oxopyrrolidin-1-yl)phenyl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

3-(5-Fluoro-6-methoxypyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

(*S*)-3-(Quinolin-3-yl)-3-(6-(3-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)propyl)-2*H*-indazol-2-yl)propanoic acid;

(3*S*)-3-(Quinolin-3-yl)-3-(6-(3-(1,2,3,4-tetrahydro-1,8-naphthyridin-2-yl)propyl)-2*H*-indazol-2-yl)propanoic acid;

3-(5-(Hydroxymethyl)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

3-(6-(2-Hydroxy-2-methylpropoxy)-5-methylpyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

3-(3,4-Dihydro-2*H*-pyrido[3,2-*b*][1,4]oxazin-6-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

3-(2-Methoxypyrimidin-5-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

3-(2-(2-Oxopyrrolidin-1-yl)pyridin-4-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

3-(Isoquinolin-6-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

3-(Pyrido[2,3-*b*]pyrazin-7-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

3-(1,8-Naphthyridin-2-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

3-(3,4-Dihydro-2*H*-pyrido[3,2-*b*][1,4]oxazin-7-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

(*S*)-2-(((Benzyloxy)carbonyl)amino)-3-(5-((5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)methoxy)-2*H*-indazol-2-yl)propanoic acid;

(*R*)-3-(5-(Hydroxymethyl)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

(*S*)-3-(5-(Hydroxymethyl)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

3-(1,8-Naphthyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

3-(5-(2-Oxopyrrolidin-1-yl)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

3-(5-(2-(5,6,7,8-Tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)-3-(1-(tetrahydro-2*H*-pyran-2-yl)pyrazolo[3,4-*b*]pyridin-5-yl)propanoic acid;

3-(5-(1,3-Dioxolan-2-yl)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

3-(5-((Dimethylamino)methyl)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

3-(1*H*-Pyrazolo[3,4-*b*]pyridin-5-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

(*S*)-3-(2-Ethoxypyrimidin-5-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

(*R*)-3-(2-Ethoxypyrimidin-5-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

3-(2,3-Dihydro-[1,4]dioxino[2,3-*b*]pyridin-7-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

3-(Benzo[*d*]thiazol-6-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

3-(2-Morpholinopyrimidin-5-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

3-(2-(Methylamino)pyrimidin-5-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

3-(6-Methoxypyridin-3-yl)-3-(5-(3-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)propyl)-1*H*-pyrazolo[4,3-*b*]pyridin-1-yl)propanoic acid;

3-(6-Methoxypyridazin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

3-([1,2,4]Triazolo[4,3-*a*]pyridin-6-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

3-(5,6,7,8-Tetrahydro-1,8-naphthyridin-2-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

3-(1-Methyl-1*H*-pyrazolo[3,4-*b*]pyridin-5-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

(*S*)-3-(2-Methoxypyrimidin-5-yl)-3-(6-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethyl)-2*H*-indazol-2-yl)propanoic acid;

3-(2-(Azetidin-1-yl)pyrimidin-5-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

3-(2-Methyl-2*H*-pyrazolo[3,4-*b*]pyridin-5-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

3-(5-(Oxazol-5-yl)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

3-(7-Ethyl-5,6,7,8-tetrahydroimidazo[1,2-*a*]pyrazin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

Butoxycarbonylamino)methylpyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

3-(5-Morpholinopyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

3-(5-(Methylsulfonyl)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

3-(3-Methyl-3*H*-imidazo[4,5-*b*]pyridin-6-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

3-(5-(Pyrrolidin-1-yl)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

(*R*)-3-(Pyrido[2,3-*b*]pyrazin-7-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

(*S*)-3-(Pyrido[2,3-*b*]pyrazin-7-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

3-([1,2,4]Triazolo[4,3-*a*]pyridin-7-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

3-(5-(Aminomethyl)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

3-([1,3]Dioxolo[4,5-*b*]pyridin-6-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

3-(5-(1,3-Dioxolan-2-yl)-6-methoxypyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

3-(6-(1*H*-Pyrazol-1-yl)pyridin-2-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

3-(1-(Pyridin-4-yl)-1*H*-pyrazol-4-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1*H*-indazol-1-yl)propanoic acid;

3-(2-Methyl-2H-pyrazolo[4,3-b]pyridin-6-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid;

3-(1-Methyl-1H-pyrazolo[4,3-b]pyridin-6-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid;

(*R*)-3-(5-(1,3-dioxolan-2-yl)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid;

(*S*)-3-(5-(1,3-dioxolan-2-yl)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid;

3-(1-methyl-1H-imidazo[4,5-b]pyridin-6-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid;

3-(5-(1H-pyrazol-5-yl)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid;

3-(6-morpholinopyrazin-2-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid;

3-(5-(1-methyl-1H-pyrazol-4-yl)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid;

3-(5-(2-hydroxypropan-2-yl)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid;

(*R*)-3-(5-(2-hydroxypropan-2-yl)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid;

(*S*)-3-(5-(2-hydroxypropan-2-yl)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid;

(*R*)-3-(2-methoxypyrimidin-5-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid;

(*S*)-3-(2-methoxypyrimidin-5-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid;

3-(6-methoxypyrazin-2-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid;

(*R*)-3-(5-(2-oxopyrrolidin-1-yl)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid;

(*S*)-3-(5-(2-oxopyrrolidin-1-yl)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid;

3-(5-(morpholine-4-carbonyl)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid;

3-(5-(dimethylcarbamoyl)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid;

3-(5-(4-methylpiperazine-1-carbonyl)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid;

3-(5-cyclopropylpyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid;

3-(5-(4-methylpiperazin-1-yl)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid;

3-(5-(azetidine-1-carbonyl)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid;

3-(5-((2-(dimethylamino)ethyl)carbamoyl)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid;

(*S*)-3-(2-methylpyrimidin-5-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid;

(*R*)-3-(2-methylpyrimidin-5-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid;

3-(5-(1H-pyrazol-1-yl)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid;

3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)hexanoic acid;

3-(5-(dimethylamino)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid;

3-cyclohexyl-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid;

(*R*)-3-(5-(methylsulfonyl)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid;

(*S*)-3-(5-(methylsulfonyl)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid;

3-(5-(((methoxycarbonyl)amino)methyl)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid;

(*R*)-3-(5-(((methoxycarbonyl)amino)methyl)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid;

(*S*)-3-(5-(((methoxycarbonyl)amino)methyl)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid;

3-(5-(methylsulfonamidomethyl)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid;

3-(5-(acetamidomethyl)pyridin-3-yl)-3-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)propanoic acid;

3-(5-(2-(5,6,7,8-Tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)-3-(5,6,7,8-tetrahydro-1,8-naphthyridin-3-yl)propanoic acid;

4-(((6-(2-Carboxy-1-(5-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)ethyl)pyrazin-2-yl)amino)butanoic acid;

3-(6-Methoxypyridin-3-yl)-3-(4-(2-(5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethyl)-1H-indazol-1-yl)propanoic acid;

(*R*)-3-(5-(2-((*R*)-7-methyl-5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)-3-(2-methylpyrimidin-5-yl)propanoic acid;

(*S*)-3-(5-(2-((*R*)-7-methyl-5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)-3-(2-methylpyrimidin-5-yl)propanoic acid;

(*R*)-3-(5-(2-((*S*)-7-methyl-5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)-3-(2-methylpyrimidin-5-yl)propanoic acid;

(*S*)-3-(5-(2-((*S*)-7-methyl-5,6,7,8-tetrahydro-1,8-naphthyridin-2-yl)ethoxy)-1H-indazol-1-yl)-3-(2-methylpyrimidin-5-yl)propanoic acid;

3-(6-Methoxypyridin-3-yl)-3-(5-(2-(1-methyl-1,2,3,4-tetrahydropyrido[2,3-*b*]pyrazin-6-yl)ethoxy)-1H-indazol-1-yl)propanoic acid;

(*S*)-3-(5-(2-(3,4-dihydro-2H-pyrido[3,2-*b*][1,4]oxazin-6-yl)ethoxy)-1H-indazol-1-yl)-3-(6-methoxypyridin-3-yl)propanoic acid; and

(*R*)-3-(5-(2-(3,4-dihydro-2H-pyrido[3,2-*b*][1,4]oxazin-6-yl)ethoxy)-1H-indazol-1-yl)-3-(6-methoxypyridin-3-yl)propanoic acid;

or a pharmaceutically acceptable salt thereof.

11. A pharmaceutical composition comprising a compound of claim 1, or a pharmaceutically acceptable salt thereof, and a carrier.

12. A method of treating a disease, disorder, or condition selected from pathological fibrosis, transplant rejection, cancer, osteoporosis, or inflammatory disorders comprising administering a therapeutically effective amount of a compound of claim 1, or a pharmaceutically acceptable salt thereof, to a patient in need thereof.

13. The method of claim 12 wherein the pathological fibrosis is pulmonary, liver, renal, cardiac, dermal, ocular, or pancreatic fibrosis.

14. The method of claim 12 wherein the disease, disorder, or condition is idiopathic pulmonary fibrosis (IPF), nonalcoholic steatohepatitis (NASH), chronic kidney disease, diabetic kidney disease, and systemic sclerosis.

15. The method of claim 12 wherein the cancer is of the bladder, blood, bone, brain, breast, central nervous system, cervix, colon, endometrium, esophagus, gall bladder, genitalia, genitourinary tract, head, kidney, larynx, liver, lung, muscle tissue, neck, oral or nasal mucosa, ovary, pancreas, prostate, skin, spleen, small intestine, large intestine, stomach, testicle, or thyroid.