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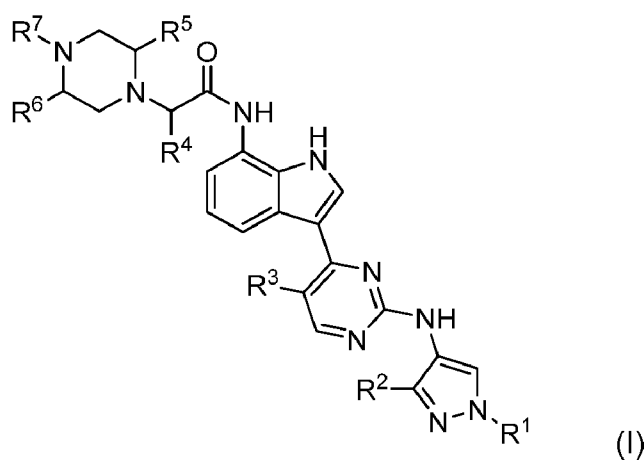
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(54) Title: COMPOUNDS AND METHODS FOR INHIBITING JAK



(57) **Abstract:** Disclosed are compounds of formula (I), pharmaceutical compositions comprising such compounds and methods/uses of using the same, for example, for treating a JAK-related disorder, such as cancer, cancer cachexia or an immune disorder: (I) wherein R¹ is methyl or ethyl; R² is selected from methyl, ethyl, methoxy and ethoxy; R³ is selected from hydrogen, chlorine, fluorine, bromine and methyl; R⁴ is selected from methyl, ethyl and -CH₂OCH₃; R⁵ and R⁶ are each individually methyl or hydrogen; and R⁷ is selected from methyl, ethyl, -(CH₂)₂OH and -(CH₂)₂OCH₃, or a pharmaceutically acceptable salt thereof.



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COMPOUNDS AND METHODS FOR INHIBITING JAK

Background

The JAK (Janus-associated kinase) family consists of four non-receptor tyrosine kinases, JAK1, JAK2, JAK3 and Tyk2, which play a critical role in cytokine and growth factor mediated signal transduction (Schindler C, and Darnell JE Jr., *Annu. Rev. Biochem.* 1995;64;621-651). Cytokine and/or growth factor binding to cell-surface receptors facilitates activation of receptor-associated JAK kinases by autophosphorylation. Activated JAKs directly phosphorylate members of the STAT (signal transducers and activators of transcription) family of transcription factors (STAT1, 2, 3, 4, 5a, 5b and 6) promoting their translocation to the nucleus and the transcriptional activation of target genes.

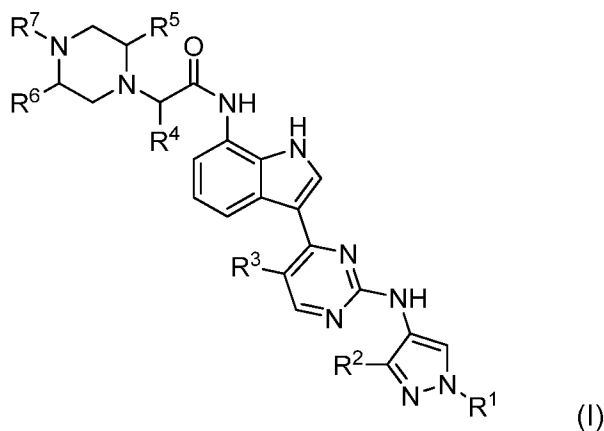
Constitutive activation (*i.e.*, tyrosine phosphorylation) of members of the STAT family, in particular STAT3, has been documented in a wide range of cancers and hyperproliferative disorders, and associated with poor prognosis in several cancers (Yu H, Jove R., *Nat. Rev. Cancer* 2004;4:97–105). Persistently activated STAT3 has been shown to be oncogenic (Bromberg JF, *et al. Cell* 1999;98:295–303) and to drive the expression of cellular proteins contributing to central processes in cancer progression (survival, proliferation, invasion, angiogenesis) (Yu and Jove, 2004, *supra*). One common mechanism of STAT3 activation in cancer cells is via autocrine or paracrine stimulation of JAK/STAT3 signaling by cytokines, typically members of the interleukin-6 (IL-6) cytokine family (Grivennikov, S. and Karin, M. *Cancer Cell* 2008;13;7-9; Bromberg J. and Wang TC. *Cancer Cell* 2009;15;79-80). This is primarily mediated by JAK1, the key JAK kinase responsible for STAT3 activation (Guschin *et al.*, *Embo J* 1995;14;1421-1429., Kim SM, *et al.*, *Mol. Cancer Ther.* 2012;11;2254-2264; Song *et al.*, *Mol. Cancer Ther.* 2011;10;481-494). Inactivation of negative regulatory proteins, such as the SOCS (suppressors of cytokine signalling) or PIAS (protein inhibitor of activated STATs) proteins have also been shown to influence the activation status of the JAK/STAT signalling pathway in cancer (Mottok *et al.*, *Blood* 2007;110;3387-90; Ogata *et al.*, *Gastroenterology* 2006;131;179-193., Lee *et al.*, *Mol. Cancer Ther.* 2006;5;8-19, Brantley *et al.*, *Clin. Cancer Res.* 2008;14;4694-4704).

In addition to basal activation of JAK1/STAT3 signaling in multiple human tumors, the pathway has also been shown to be activated as a feedback resistance mechanism in response to inhibition of driver oncogenic pathways in cancer cells, such as the mutated epidermal growth factor receptor (EGFR) in non-small cell lung cancer (NSCLC), or the MAPK pathway in KRAS mutant tumors (Lee *et al.*, *Cancer Cell* 2014;26;207-221.; VanSchaeybroeck *et al.*, *Cell reports*

Also, cancer cachexia is a significant contributor to increased mortality and poor response to chemotherapy in patients with advanced cancer. Elevated levels of inflammatory cytokines, such as IL-6, which signal through the JAK/STAT pathway have been shown to play a causal role, indicating the potential benefit of JAK1 inhibition in ameliorating cancer cachexia.

Summary

In one embodiment, disclosed are compounds of formula (I):



R¹ is methyl or ethyl;
R² is selected from methyl, ethyl, methoxy and ethoxy;
R³ is selected from hydrogen, chlorine, fluorine, bromine and methyl;
R⁴ is selected from methyl, ethyl and -CH₂OCH₃;
R⁵ and R⁶ are each individually methyl or hydrogen; and
R⁷ is selected from methyl, ethyl, -(CH₂)₂OH and -(CH₂)₂OCH₃, or a pharmaceutically acceptable salt thereof.

In one embodiment, disclosed are pharmaceutical compositions comprising a compound of Formula (I), or a pharmaceutically acceptable salt or solid form thereof, and a pharmaceutically acceptable diluent, excipient or carrier.

5 In another embodiment, disclosed are methods of treating a JAK-related disorder in a subject in need thereof comprising administering to the subject an effective amount of a compound of Formula (I), or a pharmaceutically acceptable salt or solid form thereof.

In another embodiment, disclosed is a compound of Formula (I), or a pharmaceutically acceptable salt or solid form thereof, for use in treating a JAK-related disorder.

10 In another embodiment, disclosed are pharmaceutical compositions comprising a compound of Formula (I), or a pharmaceutically acceptable salt or solid form thereof, for use in treating a JAK-related disorder.

In another embodiment, disclosed is the use of a compound of Formula (I), or a pharmaceutically acceptable salt or solid form thereof, in the manufacture of a medicament for treating a JAK-related disorder.

15 In another embodiment, disclosed are methods of treating cancer in a subject in need thereof comprising administering to the subject an effective amount of a compound as of Formula (I), or a pharmaceutically acceptable salt or solid form thereof, in combination with an anti-cancer therapeutic agent, or a pharmaceutically acceptable salt thereof.

20 In another embodiment, disclosed is a compound of Formula (I), or a pharmaceutically acceptable salt or solid form thereof, in combination with an anti-cancer therapeutic agent, or a pharmaceutically acceptable salt thereof, for use in treating cancer.

In another embodiment, disclosed are pharmaceutical compositions comprising a compound of Formula (I), or a pharmaceutically acceptable salt or solid form thereof, in combination with an anti-cancer therapeutic agent, or a pharmaceutically acceptable salt thereof, for use in treating cancer.

25 In another embodiment, disclosed is the use of a compound of Formula (I), or a pharmaceutically acceptable salt or solid form thereof, in combination with an anti-cancer therapeutic agent, or a pharmaceutically acceptable salt thereof, in the manufacture of a medicament for treating cancer.

30 In another embodiment, disclosed are methods of treating cancer cachexia in a subject in need thereof comprising administering to the subject an effective amount of a compound of Formula (I), or a pharmaceutically acceptable salt or solid form thereof.

In another embodiment, disclosed is a compound of Formula (I), or a pharmaceutically acceptable salt or solid form thereof, for use in treating cancer cachexia.

In another embodiment, disclosed are pharmaceutical compositions comprising a compound of Formula (I), or a pharmaceutically acceptable salt or solid form thereof, for use in treating cancer cachexia.

5 In another embodiment, disclosed is the use of a compound of Formula (I), or a pharmaceutically acceptable salt or solid form thereof, in the manufacture of a medicament for treating cancer cachexia.

In another embodiment, disclosed are methods of treating an immune disorder in a subject in need thereof comprising administering to the subject an effective amount of a compound of Formula (I), or a pharmaceutically acceptable salt or solid form thereof.

10 In another embodiment, disclosed is a compound of Formula (I), or a pharmaceutically acceptable salt or solid form thereof, for use in treating an immune disorder.

In another embodiment, disclosed are pharmaceutical compositions comprising a compound of Formula (I), or a pharmaceutically acceptable salt or solid form thereof, for use in treating an immune disorder.

15 In another embodiment, disclosed is the use of a compound of Formula (I), or a pharmaceutically acceptable salt or solid form thereof, in the manufacture of a medicament for treating an immune disorder.

20 In another embodiment, disclosed are methods of inhibiting JAK in a subject in need thereof comprising administering to the subject an effective amount of a compound of Formula (I), or a pharmaceutically acceptable salt or solid form thereof.

In another embodiment, disclosed is a compound of Formula (I), or a pharmaceutically acceptable salt or solid form thereof, for use in inhibiting JAK.

25 In another embodiment, disclosed are pharmaceutical compositions comprising a compound of Formula (I), or a pharmaceutically acceptable salt or solid form thereof, for use in inhibiting JAK.

In another embodiment, disclosed is the use of a compound of Formula (I), or a pharmaceutically acceptable salt or solid form thereof, in the manufacture of a medicament for inhibiting JAK.

30 **Brief Description of the Drawings**

Figure 1 illustrates the powder X-ray diffraction diagram of Form A (2R)-N-(3-{2-[(3-methoxy-1-methyl-1H-pyrazol-4-yl)amino]pyrimidin-4-yl}-1H-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide.

Figure 2 illustrates the differential scanning calorimetry (DSC) and thermogravimetric

analysis (TGA) traces of Form A (2R)-N-(3-{2-[(3-methoxy-1-methyl-1H-pyrazol-4-yl)amino]pyrimidin-4-yl}-1H-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide.

Figure 3 illustrates the powder X-ray diffraction diagram of Form B (2R)-N-(3-{2-[(3-methoxy-1-methyl-1H-pyrazol-4-yl)amino]pyrimidin-4-yl}-1H-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide.

Figure 4 illustrates the differential scanning calorimetry (DSC) and thermogravimetric analysis (TGA) traces of Form B (2R)-N-(3-{2-[(3-methoxy-1-methyl-1H-pyrazol-4-yl)amino]pyrimidin-4-yl}-1H-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide.

Figure 5 illustrates the powder X-ray diffraction diagram of Form C (2R)-N-(3-{2-[(3-methoxy-1-methyl-1H-pyrazol-4-yl)amino]pyrimidin-4-yl}-1H-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide.

Figure 6 illustrates the differential scanning calorimetry (DSC) and thermogravimetric analysis (TGA) traces of Form C (2R)-N-(3-{2-[(3-methoxy-1-methyl-1H-pyrazol-4-yl)amino]pyrimidin-4-yl}-1H-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide.

Figure 7 illustrates the powder X-ray diffraction diagram of Form D (2R)-N-(3-{2-[(3-methoxy-1-methyl-1H-pyrazol-4-yl)amino]pyrimidin-4-yl}-1H-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide.

Figure 8 illustrates the differential scanning calorimetry (DSC) and thermogravimetric analysis (TGA) traces of Form D (2R)-N-(3-{2-[(3-methoxy-1-methyl-1H-pyrazol-4-yl)amino]pyrimidin-4-yl}-1H-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide.

Figure 9 illustrates the powder X-ray diffraction diagram of Form A (2R)-N-(3-{2-[(3-methoxy-1-methyl-1H-pyrazol-4-yl)amino]pyrimidin-4-yl}-1H-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide saccharine salt.

Figure 10 illustrates the differential scanning calorimetry (DSC) and thermogravimetric analysis (TGA) traces of Form A (2R)-N-(3-{2-[(3-methoxy-1-methyl-1H-pyrazol-4-yl)amino]pyrimidin-4-yl}-1H-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide saccharine salt.

Figure 11 illustrates the powder X-ray diffraction diagram of Form B (2R)-N-(3-{2-[(3-methoxy-1-methyl-1H-pyrazol-4-yl)amino]pyrimidin-4-yl}-1H-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide saccharine salt.

Figure 12 illustrates the differential scanning calorimetry (DSC) and thermogravimetric analysis (TGA) traces of Form B (2R)-N-(3-{2-[(3-methoxy-1-methyl-1H-pyrazol-4-yl)amino]pyrimidin-4-yl}-1H-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide saccharine salt.

Figure 13 illustrates the powder X-ray diffraction diagram of Form C (2R)-N-(3-{2-[(3-methoxy-1-methyl-1H-pyrazol-4-yl)amino]pyrimidin-4-yl}-1H-indol-7-yl)-2-(4-methylpiperazin-1-

yl)propanamide saccharine salt.

Figure 14 illustrates the powder X-ray diffraction diagram of Form D (2R)-N-(3-{2-[(3-methoxy-1-methyl-1H-pyrazol-4-yl)amino]pyrimidin-4-yl}-1H-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide saccharine salt.

Figure 15 illustrates the powder X-ray diffraction diagram of Form E (2R)-N-(3-{2-[(3-methoxy-1-methyl-1H-pyrazol-4-yl)amino]pyrimidin-4-yl}-1H-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide saccharine salt.

Figure 16 illustrates the powder X-ray diffraction diagram of (2R)-N-(3-{2-[(3-methoxy-1-methyl-1H-pyrazol-4-yl)amino]pyrimidin-4-yl}-1H-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide saccharine hydrochloride salt.

Figure 17 illustrates the powder X-ray diffraction diagram of (2R)-N-(3-{2-[(3-methoxy-1-methyl-1H-pyrazol-4-yl)amino]pyrimidin-4-yl}-1H-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide napadisyllic salt.

Figure 18 illustrates the powder X-ray diffraction diagram of (2R)-N-(3-{2-[(3-methoxy-1-methyl-1H-pyrazol-4-yl)amino]pyrimidin-4-yl}-1H-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide trimesic salt.

Figure 19 illustrates NCI-H1975 tumor volumes after treatment with vehicle, osimertinib (2.5 mg/kg QD) administered as a single agent, Example 32 (12.5 mg/kg BID, 25 mg/kg BID or 50 mg/kg BID) as a single agent, and osimertinib (2.5 mg/kg QD) in combination with Example 32 (12.5 mg/kg BID, 25 mg/kg BID or 50 mg/kg BID). ● represents vehicle; ■ represents osimertinib administered as a single agent; ▲ represents 50 mg/kg BID Example 32 administered as a single agent; ◆ represents 25 mg/kg BID Example 32 administered as a single agent; ● represents 12.5 mg/kg BID Example 32 administered as a single agent; ▲ represents osimertinib administered in combination with 50 mg/kg BID Example 32; ◆ represents osimertinib administered in combination with 25 mg/kg BID Example 32; and ● represents osimertinib administered in combination with 12.5 mg/kg BID Example 32.

Figure 20 illustrates body weights after treatment with vehicle, osimertinib (2.5 mg/kg QD) as a single agent, Example 32 (12.5 mg/kg BID, 25 mg/kg BID or 50 mg/kg BID) as a single agent, and osimertinib (2.5 mg/kg QD) in combination with Example 32 (12.5 mg/kg BID, 25 mg/kg BID or 50 mg/kg BID). ● represents vehicle; ■ represents osimertinib administered as a single agent; ▲ represents 50 mg/kg BID Example 32 administered as a single agent; ◆ represents 25 mg/kg BID Example 32 administered as a single agent; ● represents 12.5 mg/kg BID Example 32 administered as a single agent; ▲ represents 50 mg/kg BID Example



32 administered in combination with osimertinib;  represents 25 mg/kg BID Example 32 administered in combination with osimertinib; and  represents 12.5 mg/kg BID Example 32 administered in combination with osimertinib.


Figure 21 illustrates knockdown of pSTAT3 in NCI-H1975 tumors after treatment with vehicle, AZD1480 as a single agent, osimertinib (2.5 mg/kg QD) as a single agent, Example 32 (12.5 mg/kg BID, 25 mg/kg BID or 50 mg/kg BID) as a single agent, and osimertinib (2.5 mg/kg QD) administered in combination with Example 32 (12.5 mg/kg BID, 25 mg/kg BID or 50 mg/kg BID).  represents pSTAT3 and bars represent plasma levels of Example 32.






Figure 22 illustrates PC-9 tumor volumes after treatment with vehicle, gefitinib (IRESSA®, 6.25 mg/kg QD) administered as a single agent and gefitinib (IRESSA®, 6.25 mg/kg QD) administered in combination with Example 32 (12.5 mg/kg BID, 50 mg/kg BID and 50 mg/kg BID dosed 2 days on/5 days off/wk).  represents vehicle;  represents gefitinib (IRESSA®) administered as a single agent;  represents gefitinib (IRESSA®) administered in combination with 50 mg/kg BID Example 32;  represents gefitinib (IRESSA®) administered in combination with 12.5 mg/kg BID Example 32; and  represents gefitinib (IRESSA®) administered in combination with 50 mg/kg BID with Example 32 dosed 2 days on/5 days off/wk.









Figure 23 illustrates body weights after treatment with vehicle, gefitinib (IRESSA®, 6.25 mg/kg QD) administered as a single agent and gefitinib (IRESSA®, 6.25 mg/kg QD) administered in combination with Example 32 (12.5 mg/kg BID, 50 mg/kg BID and 50 mg/kg 2 dosed days on/5 days off/wk).  represents vehicle;  represents gefitinib (IRESSA®) administered as a single agent;  represents gefitinib (IRESSA®) administered in combination with 50 mg/kg BID Example 32;  represents gefitinib (IRESSA®) administered in combination with 12.5 mg/kg BID Example 32; and  represents gefitinib (IRESSA®) administered in combination with 50 mg/kg BID Example 32 dosed 2 days on/5 days off/wk.

Figure 24 illustrates knockdown of pSTAT3 in PC-9 tumors after treatment with vehicle, gefitinib (IRESSA®, 6.25 mg/kg QD) administered as a single agent and gefitinib (IRESSA®, 6.25 mg/kg QD) administered in combination with Example 32 (12.5 mg/kg BID and 50 mg/kg BID).

Figure 25 illustrates NCI-H1650 tumor volumes after treatment with vehicle, gefitinib (IRESSA®, 6.25 mg/kg QD) administered as a single agent, Example 32 (25 mg/kg BID or 50 mg/kg BID) administered as a single agent and gefitinib (IRESSA®, 6.25 mg/kg QD) administered in combination with Example 32 (25 mg/kg BID or 50 mg/kg BID).  represents vehicle;  represents gefitinib (IRESSA®) administered as a single agent;  represents 25




mg/kg BID Example 32 administered as a single agent;  represents 50 mg/kg BID Example 32 administered as a single agent;  represents gefitinib (IRESSA®) administered in combination with 25 mg/kg BID Example 32; and  represents gefitinib (IRESSA®) administered in combination with 50 mg/kg BID Example 32.







Figure 26 illustrates body weights after treatment with vehicle, gefitinib (IRESSA®, 6.25 mg/kg QD) administered as a single agent, Example 32 (25 mg/kg BID or 50 mg/kg BID) administered as a single agent and gefitinib (IRESSA®, 6.25 mg/kg QD) administered in combination with Example 32 (25 mg/kg BID or 50 mg/kg BID).  represents vehicle;  represents gefitinib (IRESSA®) administered as a single agent;  represents 25 mg/kg BID Example 32 administered as a single agent;  represents 50 mg/kg BID Example 32 administered as a single agent;  represents gefitinib (IRESSA®) administered in combination with 25 mg/kg BID Example 32; and  represents gefitinib (IRESSA®) administered in combination with 50 mg/kg BID Example 32.


Figure 27 illustrates knockdown of pSTAT3 in NCI-H1650 tumors after treatment with vehicle, AZD1408 administered as a single agent, gefitinib (IRESSA®, 6.25 mg/kg QD) administered as a single agent, Example 32 (25 mg/kg BID or 50 mg/kg BID) administered as a single agent and gefitinib (IRESSA®, 6.25 mg/kg QD) administered in combination with Example 32 (25 mg/kg BID or 50 mg/kg BID).  represents pSTAT and bars represent plasma levels of Example 32.






Figure 28 illustrates LG1049 tumor volumes after treatment with vehicle (); osimertinib (25 mg/kg QD) administered as a single agent for 28 days (), Example 32 (25 mg/kg BID) administered as a single agent for 18 days (); osimertinib (25 mg/kg QD) administered in combination with Example 32 (25 mg/kg BID) dosed 7 days, then 3 days on/4 days off/wk until day 28 ().  represents mice treated with the combination for 28 days, then with Example 32 (25 mg/kg BID) alone for 3 days on/4 days off/wk until the end of the study.






Figure 29 illustrates body weights after treatment LG1049 tumor volumes after treatment with vehicle (), osimertinib (25 mg/kg QD) administered as a single agent for 28 days (), Example 32 (25 mg/kg BID) administered as a single agent (), osimertinib (25 mg/kg QD) administered in combination with Example 32 (25 mg/kg BID) dosed 7 days, then 3 days on/4 days off until day 28 ().  represents mice treated with the combination for 28 days, then with Example 32 (25 mg/kg BID) alone for 3 days on/4 days off/wk until the end of the study

Figure 30 illustrates knockdown of pSTAT3 and pEGFR in LG1049 tumors after five days

of treatment with vehicle, osimertinib (25 mg/kg QD) administered as a single agent, Example 32 (25 mg/kg BID) administered as a single agent and osimertinib (25 mg/kg QD) administered in combination with Example 32 (25 mg/kg BID).

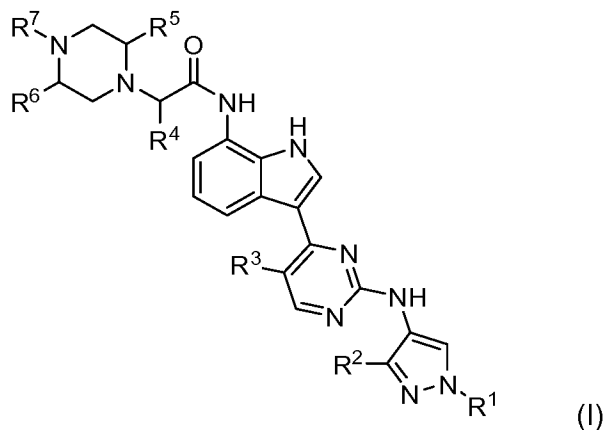
Figure 31 illustrates NCI-H1975 tumor volumes after treatment with vehicle, Example 32 administered as a single agent, osimertinib administered as a single agent, and Example 32 administered in combination with osimertinib. **Figure 31A** illustrates tumor volume over time after continuous dosing with vehicle for 19 days (●); 50 mg/kg BID Example 32 administered as a single agent for 19 days (▲); 2.5 mg/kg QD osimertinib administered as a single agent for 26 days (■); 2.5 mg/kg QD osimertinib administered in combination with 12.5 mg/kg BID Example 32 dosed for 26 days (▲); and 2.5 mg/kg QD osimertinib administered in combination with 50 mg/kg BID Example 32 dosed for 26 days (▽). **Figure 31B** illustrates tumor volume over time after dosing with vehicle (●) for 19 days; 50 mg/kg BID Example 32 administered as a single agent for 19 days (▲); 2.5 mg/kg QD osimertinib administered as a single agent for 26 days (■); 2.5 mg/kg QD osimertinib administered for 26 days in combination with 25 mg/kg BID Example 32 dosed for 7 days (▲); and 2.5 mg/kg QD osimertinib administered for 26 days in combination with 50 mg/kg BID Example 32 dosed for 7 days (▽). **Figure 31C** illustrates tumor volume over time after dosing with vehicle for 19 days (●); 50 mg/kg BID Example 32 as a single agent for 19 days (▲); 2.5 mg/kg QD osimertinib administered as a single agent for 26 days (■); 2.5 mg/kg QD osimertinib administered for 29 days in combination with 25 mg/kg BID Example 32 dosed 7 days on/7 days off/2wk (▲); and 2.5 mg/kg QD osimertinib administered for 29 days in combination with 50 mg/kg BID Example 32 dosed 7 days on/7 days off/2wk (▽). **Figure 31D** illustrates tumor volume over time after dosing with vehicle for 19 days (●); 50 mg/kg BID Example 32 administered as a single agent for 19 days (▲); 2.5 mg/kg QD osimertinib administered as a single agent for 26 days (■); 2.5 mg/kg QD osimertinib administered for 29 days in combination with 25 mg/kg BID Example 32 dosed 4 days on/3 days off/wk (▲); and 2.5 mg/kg QD osimertinib administered for 29 days in combination with 50 mg/kg BID Example 32 dosed for 4 days on/3 days off/wk (▽). **Figure 31E** illustrates tumor volume over time after dosing with vehicle for 19 days (●); 50 mg/kg BID Example 32 for 19 days (▲); 2.5 mg/kg QD osimertinib dosed for 26 days (■); 2.5 mg/kg QD osimertinib dosed for 29 days in combination with 25 mg/kg BID Example 32 dosed for 2 days on/5 days off/wk (▲); and 2.5 mg/kg QD osimertinib dosed for 29 days in combination with 50 mg/kg BID Example 32 dosed 2 days on/5 days off/wk (▽).

Figure 32 illustrates body weight over time after treatment with vehicle, Example 32 as a single agent, osimertinib as a single agent, and Example 32 in combination with osimertinib. ● represents vehicle dosed for 19 days; ■ represents 2.5 mg/kg QD osimertinib dosed for 26 days; ◆ represents 50 mg/kg BID Example 32 dosed for 19 days; ▲ represents 2.5 mg/kg QD osimertinib administered in combination with 12.5 mg/kg BID Example 32 dosed for 26 days; ● represents 2.5 mg/kg QD osimertinib administered in combination with 50 mg/kg BID Example 32 dosed for 26 days; ▼ represents 2.5 mg/kg QD osimertinib administered for 26 days in combination with 25 mg/kg BID Example 32 dosed for 7 days; ○ represents 2.5 mg/kg QD osimertinib administered for 26 days in combination with 50 mg/kg BID Example 32 dosed for 7 days; □ represents 2.5 mg/kg QD osimertinib administered for 29 days in combination with 50 mg/kg BID Example 32 dosed for 7 days on/7 days off for one week; ◇ represents 2.5 mg/kg QD osimertinib administered for 29 days in combination with 50 mg/kg BID Example 32 dosed for 7 days on/7 days off for 2 weeks; ▲ represents 2.5 mg/kg QD osimertinib administered for 29 days in combination with 25 mg/kg BID Example 32 dosed for 4 days on/3 days off for one week; ○ represents 2.5 mg/kg QD osimertinib administered for 29 days in combination with 50 mg/kg BID Example 32 dosed for 4 days on/3 days off for one week; ▼ represents 2.5 mg/kg QD osimertinib administered for 29 days in combination with 25 mg/kg BID Example 32 dosed for 2 days on/5 days off for one week; and ■ represents 2.5 mg/kg QD osimertinib administered for 29 days in combination with 50 mg/kg BID Example 32 dosed for 2 days on/5 days off for one week.

Detailed Description

Compounds

In one embodiment, disclosed are compounds of formula (I):



wherein

R¹ is methyl or ethyl;

R² is selected from methyl, ethyl, methoxy and ethoxy;

R³ is selected from hydrogen, chlorine, fluorine, bromine and methyl;

5 R⁴ is selected from methyl, ethyl and -CH₂OCH₃;

R⁵ and R⁶ are each individually methyl or hydrogen; and

R⁷ is selected from methyl, ethyl, -(CH₂)₂OH and -(CH₂)₂OCH₃, or a pharmaceutically acceptable salt thereof.

10 In some embodiments, R¹ is methyl; R² is methoxy, ethoxy, methyl or ethyl; R³ is hydrogen, fluorine methyl, chlorine or bromine; R⁴ is methyl, ethyl or -CH₂OCH₃; R⁵ is hydrogen or methyl; R⁶ is hydrogen or methyl; and R⁷ is methyl, -(CH₂)₂OCH₃, ethyl or -(CH₂)₂OH.

In some embodiments, R¹ is ethyl; R² is methoxy or ethoxy; R³ is methyl; R⁴ is methyl; R⁵ is hydrogen; R⁶ is hydrogen and R⁷ is methyl.

15 In some embodiments, R² is methoxy; R¹ is methyl or ethyl; R³ is hydrogen, fluorine, methyl, chlorine or bromine; R⁴ is methyl, ethyl or -CH₂OCH₃; R⁵ is hydrogen or methyl; R⁶ is hydrogen or methyl; and R⁷ is methyl, ethyl, -(CH₂)₂OCH₃ or -(CH₂)₂OH.

In some embodiments, R² is ethoxy; R¹ is methyl or ethyl; R³ is fluorine, methyl or chlorine; R⁴ is methyl, ethyl or -CH₂OCH₃; R⁵ is methyl or hydrogen; R⁶ is methyl or hydrogen; and R⁷ is ethyl, methyl or -(CH₂)₂OCH₃.

20 In some embodiments, R² is methyl; R¹ is methyl; R³ is hydrogen, methyl or fluorine; R⁴ is methyl, ethyl or -CH₂OCH₃; R⁵ is hydrogen; R⁶ is hydrogen and R⁷ is methyl.

In some embodiments, R² is ethyl; R¹ is methyl; R³ is methyl; R⁴ is methyl; R⁵ is hydrogen; R⁶ is hydrogen and R⁷ is methyl.

25 In some embodiments, R³ is hydrogen; R¹ is methyl; R² is methoxy or methyl; R⁴ is methyl, ethyl or -CH₂OCH₃; R⁵ is methyl or hydrogen; R⁶ is hydrogen or methyl and R⁷ is methyl, -(CH₂)₂OCH₃ or ethyl.

In some embodiments, R³ is fluorine; R¹ is methyl; R² is methoxy or ethoxy; R⁴ is methyl, ethyl or -CH₂OCH₃; R⁵ is hydrogen or methyl; R⁶ is hydrogen or methyl and R⁷ is methyl, ethyl or -(CH₂)₂OCH₃.

30 In some embodiments, R³ is methyl; R¹ is methyl or ethyl; R² is methoxy, ethoxy, methyl or ethyl; R⁴ is methyl, ethyl or -CH₂OCH₃; R⁵ is hydrogen or methyl; R⁶ is hydrogen or methyl and R⁷ is methyl, ethyl, -(CH₂)₂OCH₃ or -(CH₂)₂OH.

In some embodiments, R³ is chlorine; R¹ is methyl; R² is methoxy or ethoxy; R⁴ is methyl; R⁵ is hydrogen; R⁶ is hydrogen and R⁷ is methyl.

In some embodiments, R³ is bromine; R¹ is methyl; R² is methoxy; R⁴ is methyl; R⁵ is hydrogen; R⁶ is hydrogen and R⁷ is methyl.

In some embodiments, R⁴ is methyl; R¹ is methyl or ethyl; R² is methoxy, ethoxy, methyl or ethyl; R³ is hydrogen, fluorine, methyl, chlorine or bromine; R⁵ is hydrogen or methyl; R⁶ is hydrogen or methyl and R⁷ is methyl, ethyl, $-(CH_2)_2OCH_3$ or $-(CH_2)_2OH$.

In some embodiments, R⁴ is ethyl; R¹ is methyl; R² is methoxy, methyl or ethoxy; R³ is methyl, hydrogen or fluorine; R⁵ is hydrogen; R⁶ is hydrogen and R⁷ is methyl.

In some embodiments, R⁴ is $-CH_2OCH_3$; R¹ is methyl; R² is methoxy, methyl or ethoxy; R³ is methyl, fluorine or hydrogen; R⁵ is hydrogen; R⁶ is hydrogen and R⁷ is methyl.

In some embodiments, R⁵ is hydrogen; R¹ is methyl or ethyl; R² is methoxy, ethoxy, methyl or ethyl; R³ is hydrogen, fluorine, methyl, chlorine or bromine; R⁴ is methyl, ethyl or $-CH_2OCH_3$; R⁶ is hydrogen or methyl and R⁷ is methyl, ethyl, $-(CH_2)_2OCH_3$ or $-(CH_2)_2OH$.

In some embodiments, R⁵ is methyl; R¹ is methyl; R² is methoxy or ethoxy; R³ is hydrogen, fluorine or methyl; R⁴ is methyl; R⁶ is hydrogen and R⁷ is methyl.

In some embodiments, R⁶ is hydrogen; R¹ is methyl or ethyl; R² is methoxy, ethoxy, methyl or ethyl; R³ is hydrogen, fluorine, methyl, chlorine or bromine; R⁴ is methyl, ethyl or $-CH_2OCH_3$; R⁵ is hydrogen or methyl and R⁷ is methyl, ethyl, $-(CH_2)_2OCH_3$ or $-(CH_2)_2OH$.

In some embodiments, R⁶ is methyl; R¹ is methyl; R² is methoxy or ethoxy; R³ is fluorine, methyl or hydrogen; R⁴ is methyl; R⁵ is hydrogen and R⁷ is methyl.

In some embodiments, R⁷ is methyl; R¹ is methyl or ethyl; R² is methyl, ethyl, methoxy or ethoxy; R³ is hydrogen, chlorine, fluorine, bromine or methyl; R⁴ is methyl, ethyl or $-CH_2OCH_3$; R⁵ is hydrogen or methyl; and R⁶ is hydrogen or methyl.

In some embodiments, R⁷ is ethyl; R¹ is methyl; R² is methoxy or ethoxy; R³ is fluorine, methyl or hydrogen; R⁴ is methyl; R⁵ is hydrogen and R⁶ is hydrogen.

In some embodiments, R⁷ is $-(CH_2)_2OCH_3$; R¹ is methyl; R² is methoxy or ethoxy; R³ is fluorine, methyl or hydrogen; R⁴ is methyl; R⁵ is hydrogen and R⁶ is hydrogen.

In some embodiments, R⁷ is $-(CH_2)_2OH$; R¹ is methyl, R² is methoxy; R³ is methyl; R⁴ is methyl; R⁵ is hydrogen and R⁶ is hydrogen.

In some embodiments, R¹ is methyl; R² is methoxy; R³ is hydrogen; R⁴ is methyl; R⁵ is methyl; R⁶ is hydrogen and R⁷ is methyl.

In some embodiments, R¹ is methyl; R² is methoxy; R³ is fluorine; R⁴ is methyl; R⁵ is hydrogen; R⁶ is methyl and R⁷ is methyl.

In some embodiments, R¹ is methyl; R² is methoxy; R³ is fluorine; R⁴ is methyl; R⁵ is hydrogen; R⁶ is hydrogen and R⁷ is $-(CH_2)_2OCH_3$.

In some embodiments, R¹ is methyl; R² is methoxy; R³ is fluorine; R⁴ is methyl; R⁵ is hydrogen; R⁶ is hydrogen and R⁷ is ethyl.

In some embodiments, R¹ is methyl; R² is methoxy; R³ is fluorine; R⁴ is methyl; R⁵ is methyl; R⁶ is hydrogen and R⁷ is methyl.

5 In some embodiments, R¹ is methyl; R² is ethoxy; R³ is fluorine; R⁴ is methyl; R⁵ is hydrogen; R⁶ is hydrogen and R⁷ is ethyl.

In some embodiments, R¹ is methyl; R² is ethoxy; R³ is fluorine; R⁴ is methyl; R⁵ is hydrogen; R⁶ is hydrogen and R⁷ is $-(CH_2)_2OCH_3$.

10 In some embodiments, R¹ is methyl; R² is ethoxy; R³ is fluorine; R⁴ is methyl; R⁵ is methyl; R⁶ is hydrogen and R⁷ is methyl.

In some embodiments, R¹ is methyl; R² is ethoxy; R³ is fluorine; R⁴ is methyl; R⁵ is hydrogen; R⁶ is methyl and R⁷ is methyl.

In some embodiments, R¹ is methyl; R² is ethoxy; R³ is fluorine; R⁴ is methyl; R⁵ is methyl; R⁶ is hydrogen and R⁷ is methyl.

15 In some embodiments, R¹ is methyl; R² is methoxy; R³ is methyl; R⁴ is methyl; R⁵ is hydrogen; R⁶ is methyl and R⁷ is methyl.

In some embodiments, R¹ is methyl; R² is methoxy; R³ is methyl; R⁴ is methyl; R⁵ is hydrogen; R⁶ is hydrogen and R⁷ is $-(CH_2)_2OH$.

20 In some embodiments, R¹ is methyl; R² is methoxy; R³ is methyl; R⁴ is methyl; R⁵ is hydrogen; R⁶ is hydrogen and R⁷ is ethyl.

In some embodiments, R¹ is methyl; R² is methoxy; R³ is methyl; R⁴ is methyl; R⁵ is hydrogen; R⁶ is hydrogen and R⁷ is $-(CH_2)_2OCH_3$.

In some embodiments, R¹ is methyl; R² is methoxy; R³ is methyl; R⁴ is methyl; R⁵ is methyl; R⁶ is hydrogen and R⁷ is methyl.

25 In some embodiments, R¹ is methyl; R² is ethoxy; R³ is methyl; R⁴ is methyl; R⁵ is hydrogen; R⁶ is hydrogen and R⁷ is ethyl.

In some embodiments, R¹ is methyl; R² is ethoxy; R³ is methyl; R⁴ is methyl; R⁵ is hydrogen; R⁶ is hydrogen and R⁷ is $-(CH_2)_2OCH_3$.

30 In some embodiments, R¹ is methyl; R² is ethoxy; R³ is methyl; R⁴ is methyl; R⁵ is methyl; R⁶ is hydrogen and R⁷ is methyl.

In some embodiments, R¹ is methyl; R² is ethoxy; R³ is methyl; R⁴ is methyl; R⁵ is hydrogen; R⁶ is methyl and R⁷ is methyl.

In some embodiments, R¹ is methyl; R² is methoxy; R³ is fluorine; R⁴ is methyl; R⁵ is hydrogen; R⁶ is hydrogen and R⁷ is methyl.

In some embodiments, R¹ is methyl; R² is methoxy; R³ is hydrogen; R⁴ is methyl; R⁵ is hydrogen; R⁶ is methyl and R⁷ is methyl.

In some embodiments, R¹ is methyl; R² is methoxy; R³ is hydrogen; R⁴ is methyl; R⁵ is hydrogen; R⁶ is hydrogen and R⁷ is $-(CH_2)_2OCH_3$.

5 In some embodiments, R¹ is methyl; R² is methoxy; R³ is hydrogen; R⁴ is methyl; R⁵ is hydrogen; R⁶ is hydrogen and R⁷ is ethyl.

In some embodiments, R¹ is methyl; R² is methoxy; R³ is hydrogen; R⁴ is methyl; R⁵ is methyl; R⁶ is hydrogen and R⁷ is methyl.

10 In some embodiments, R¹ is methyl; R² is methoxy; R³ is hydrogen; R⁴ is methyl; R⁵ is hydrogen; R⁶ is methyl and R⁷ is methyl.

In some embodiments, R¹ is methyl; R² is ethoxy; R³ is methyl; R⁴ is methyl; R⁵ is hydrogen; R⁶ is methyl and R⁷ is methyl.

In some embodiments, R¹ is methyl; R² is methoxy; R³ is hydrogen; R⁴ is methyl; R⁵ is hydrogen; R⁶ is hydrogen and R⁷ is methyl.

15 In some embodiments, R¹ is methyl; R² is methoxy; R³ is methyl; R⁴ is methyl; R⁵ is hydrogen; R⁶ is hydrogen and R⁷ is methyl.

In some embodiments, R¹ is methyl; R² is ethoxy; R³ is methyl; R⁴ is methyl; R⁵ is hydrogen; R⁶ is hydrogen and R⁷ is methyl.

20 In some embodiments, R¹ is methyl; R² is methoxy; R³ is chlorine; R⁴ is methyl; R⁵ is hydrogen; R⁶ is hydrogen and R⁷ is methyl.

In some embodiments, R¹ is methyl; R² is methoxy; R³ is bromine; R⁴ is methyl; R⁵ is hydrogen; R⁶ is hydrogen and R⁷ is methyl.

In some embodiments, R¹ is methyl; R² is ethoxy; R³ is fluorine; R⁴ is methyl; R⁵ is hydrogen; R⁶ is hydrogen and R⁷ is methyl.

25 In some embodiments, R¹ is methyl; R² is ethoxy; R³ is chlorine; R⁴ is methyl; R⁵ is hydrogen; R⁶ is hydrogen and R⁷ is methyl.

In some embodiments, R¹ is methyl; R² is methyl; R³ is hydrogen; R⁴ is methyl; R⁵ is hydrogen; R⁶ is hydrogen and R⁷ is methyl.

30 In some embodiments, R¹ is methyl; R² is methoxy; R³ is methyl; R⁴ is ethyl; R⁵ is hydrogen; R⁶ is hydrogen and R⁷ is methyl.

In some embodiments, R¹ is methyl; R² is methoxy; R³ is methyl; R⁴ is $-CH_2OCH_3$; R⁵ is hydrogen; R⁶ is hydrogen and R⁷ is methyl.

In some embodiments, R¹ is methyl; R² is methoxy; R³ is hydrogen; R⁴ is ethyl; R⁵ is hydrogen; R⁶ is hydrogen and R⁷ is methyl.

In some embodiments, R¹ is methyl; R² is methyl; R³ is fluorine; R⁴ is ethyl; R⁵ is hydrogen; R⁶ is hydrogen and R⁷ is methyl.

In some embodiments, R¹ is methyl; R² is methoxy; R³ is fluorine; R⁴ is ethyl; R⁵ is hydrogen; R⁶ is hydrogen and R⁷ is methyl.

5 In some embodiments, R¹ is methyl; R² is methyl; R³ is fluorine; R⁴ is -CH₂OCH₃; R⁵ is hydrogen; R⁶ is hydrogen and R⁷ is methyl.

In some embodiments, R¹ is methyl; R² is ethyl; R³ is methyl; R⁴ is methyl; R⁵ is hydrogen; R⁶ is hydrogen and R⁷ is methyl.

10 In some embodiments, R¹ is methyl; R² is methyl; R³ is fluorine; R⁴ is methyl; R⁵ is hydrogen; R⁶ is hydrogen and R⁷ is methyl.

In some embodiments, R¹ is methyl; R² is methyl; R³ is hydrogen; R⁴ is methyl; R⁵ is hydrogen; R⁶ is hydrogen and R⁷ is methyl.

In some embodiments, R¹ is ethyl; R² is methoxy; R³ is methyl; R⁴ is methyl; R⁵ is hydrogen; R⁶ is hydrogen and R⁷ is methyl.

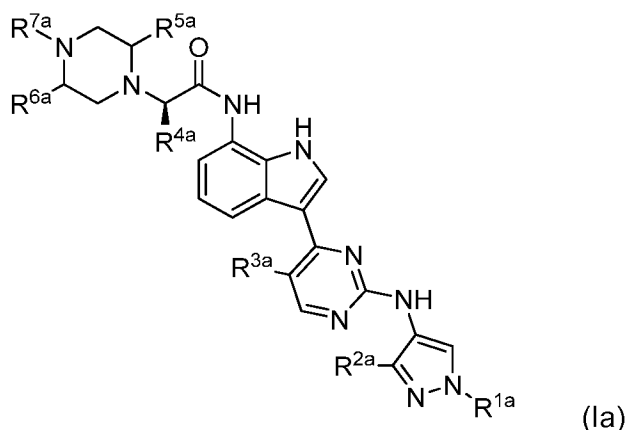
15 In some embodiments, R¹ is ethyl; R² is ethoxy; R³ is methyl; R⁴ is methyl; R⁵ is hydrogen; R⁶ is hydrogen and R⁷ is methyl.

In some embodiments, R¹ is methyl; R² is methoxy; R³ is hydrogen; R⁴ is -CH₂OCH₃; R⁵ is hydrogen; R⁶ is hydrogen and R⁷ is methyl.

20 In some embodiments, R¹ is methyl; R² is ethoxy; R³ is methyl; R⁴ is ethyl; R⁵ is hydrogen; R⁶ is hydrogen and R⁷ is methyl.

In some embodiments, R¹ is methyl; R² is ethoxy; R³ is methyl; R⁴ is -CH₂OCH₃; R⁵ is hydrogen; R⁶ is hydrogen and R⁷ is methyl.

In one embodiment, the compounds of formula (I) are compounds of formula (Ia):



25 wherein

R^{1a} is methyl or ethyl;

R^{2a} is selected from methyl, ethyl, methoxy and ethoxy;

R^{3a} is selected from hydrogen, chlorine, fluorine, bromine and methyl;

R^{4a} is selected from methyl, ethyl and -CH₂OCH₃;

R^{5a} and R^{6a} are each individually methyl or hydrogen; and

5 R^{7a} is selected from methyl, ethyl, -(CH₂)₂OH and -(CH₂)₂OCH₃, or a pharmaceutically acceptable salt thereof.

In some embodiments, R^{1a} is methyl; R^{2a} is methoxy, ethoxy, methyl or ethyl; R^{3a} is hydrogen, fluorine methyl, chlorine or bromine; R^{4a} is methyl, ethyl or -CH₂OCH₃; R^{5a} is hydrogen or methyl; R^{6a} is hydrogen or methyl; and R^{7a} is methyl, -(CH₂)₂OCH₃, ethyl or
10 -(CH₂)₂OH.

In some embodiments, R^{1a} is ethyl; R^{2a} is methoxy or ethoxy; R^{3a} is methyl; R^{4a} is methyl; R^{5a} is hydrogen; R^{6a} is hydrogen and R^{7a} is methyl.

In some embodiments, R^{2a} is methoxy; R^{1a} is methyl or ethyl; R^{3a} is hydrogen, fluorine, methyl, chlorine or bromine; R^{4a} is methyl, ethyl or -CH₂OCH₃; R^{5a} is hydrogen or methyl; R^{6a} is
15 hydrogen or methyl; and R^{7a} is methyl, ethyl, -(CH₂)₂OCH₃ or -(CH₂)₂OH.

In some embodiments, R^{2a} is ethoxy; R^{1a} is methyl or ethyl; R^{3a} is fluorine, methyl or chlorine; R^{4a} is methyl, ethyl or -CH₂OCH₃; R^{5a} is methyl or hydrogen; R^{6a} is methyl or hydrogen; and R^{7a} is ethyl, methyl or -(CH₂)₂OCH₃.

In some embodiments, R^{2a} is methyl; R^{1a} is methyl; R^{3a} is hydrogen, methyl or fluorine; R^{4a}
20 is methyl, ethyl or -CH₂OCH₃; R^{5a} is hydrogen; R^{6a} is hydrogen and R^{7a} is methyl.

In some embodiments, R^{2a} is ethyl; R^{1a} is methyl; R^{3a} is methyl; R^{4a} is methyl; R^{5a} is hydrogen; R^{6a} is hydrogen and R^{7a} is methyl.

In some embodiments, R^{3a} is hydrogen; R^{1a} is methyl; R^{2a} is methoxy or methyl; R^{4a} is methyl, ethyl or -CH₂OCH₃; R^{5a} is methyl or hydrogen; R^{6a} is hydrogen or methyl and R^{7a} is
25 methyl, -(CH₂)₂OCH₃ or ethyl.

In some embodiments, R^{3a} is fluorine; R^{1a} is methyl; R^{2a} is methoxy or ethoxy; R^{4a} is methyl, ethyl or -CH₂OCH₃; R^{5a} is hydrogen or methyl; R^{6a} is hydrogen or methyl and R^{7a} is methyl, ethyl or -(CH₂)₂OCH₃.

In some embodiments, R^{3a} is methyl; R^{1a} is methyl or ethyl; R^{2a} is methoxy, ethoxy, methyl
30 or ethyl; R^{4a} is methyl, ethyl or -CH₂OCH₃; R^{5a} is hydrogen or methyl; R^{6a} is hydrogen or methyl and R^{7a} is methyl, ethyl, -(CH₂)₂OCH₃ or -(CH₂)₂OH.

In some embodiments, R^{3a} is chlorine; R^{1a} is methyl; R^{2a} is methoxy or ethoxy; R^{4a} is methyl; R^{5a} is hydrogen; R^{6a} is hydrogen and R^{7a} is methyl.

In some embodiments, R^{3a} is bromine; R^{1a} is methyl; R^{2a} is methoxy; R^{4a} is methyl; R^{5a} is hydrogen; R^{6a} is hydrogen and R^{7a} is methyl.

In some embodiments, R^{4a} is methyl; R^{1a} is methyl or ethyl; R^{2a} is methoxy, ethoxy, methyl or ethyl; R^{3a} is hydrogen, fluorine, methyl, chlorine or bromine; R^{5a} is hydrogen or methyl; R^{6a} is hydrogen or methyl and R^{7a} is methyl, ethyl, $-(CH_2)_2OCH_3$ or $-(CH_2)_2OH$.

In some embodiments, R^{4a} is ethyl; R^{1a} is methyl; R^{2a} is methoxy, methyl or ethoxy; R^{3a} is methyl, hydrogen or fluorine; R^{5a} is hydrogen; R^{6a} is hydrogen and R^{7a} is methyl.

In some embodiments, R^{4a} is $-CH_2OCH_3$; R^{1a} is methyl; R^{2a} is methoxy, methyl or ethoxy; R^{3a} is methyl, fluorine or hydrogen; R^{5a} is hydrogen; R^{6a} is hydrogen and R^{7a} is methyl.

In some embodiments, R^{5a} is hydrogen; R^{1a} is methyl or ethyl; R^{2a} is methoxy, ethoxy, methyl or ethyl; R^{3a} is hydrogen, fluorine, methyl, chlorine or bromine; R^{4a} is methyl, ethyl or $-CH_2OCH_3$; R^{6a} is hydrogen or methyl and R^{7a} is methyl, ethyl, $-(CH_2)_2OCH_3$ or $-(CH_2)_2OH$.

In some embodiments, R^{5a} is methyl; R^{1a} is methyl; R^{2a} is methoxy or ethoxy; R^{3a} is hydrogen, fluorine or methyl; R^{4a} is methyl; R^{6a} is hydrogen and R^{7a} is methyl.

In some embodiments, R^{6a} is hydrogen; R^{1a} is methyl or ethyl; R^{2a} is methoxy, ethoxy, methyl or ethyl; R^{3a} is hydrogen, fluorine, methyl, chlorine or bromine; R^{4a} is methyl, ethyl or $-CH_2OCH_3$; R^{5a} is hydrogen or methyl and R^{7a} is methyl, ethyl, $-(CH_2)_2OCH_3$ or $-(CH_2)_2OH$.

In some embodiments, R^{6a} is methyl; R^{1a} is methyl; R^{2a} is methoxy or ethoxy; R^{3a} is fluorine, methyl or hydrogen; R^{4a} is methyl; R^{5a} is hydrogen and R^{7a} is methyl.

In some embodiments, R^{7a} is methyl; R^{1a} is methyl or ethyl; R^{2a} is methyl, ethyl, methoxy or ethoxy; R^{3a} is hydrogen, chlorine, fluorine, bromine or methyl; R^{4a} is methyl, ethyl or $-CH_2OCH_3$; R^{5a} is hydrogen or methyl; and R^{6a} is hydrogen or methyl.

In some embodiments, R^{7a} is ethyl; R^{1a} is methyl; R^{2a} is methoxy or ethoxy; R^{3a} is fluorine, methyl or hydrogen; R^{4a} is methyl; R^{5a} is hydrogen and R^{6a} is hydrogen.

In some embodiments, R^{7a} is $-(CH_2)_2OCH_3$; R^{1a} is methyl; R^{2a} is methoxy or ethoxy; R^{3a} is fluorine, methyl or hydrogen; R^{4a} is methyl; R^{5a} is hydrogen and R^{6a} is hydrogen.

In some embodiments, R^{7a} is $-(CH_2)_2OH$; R^{1a} is methyl, R^{2a} is methoxy; R^{3a} is methyl; R^{4a} is methyl; R^{5a} is hydrogen and R^{6a} is hydrogen.

In some embodiments, R^{1a} is methyl; R^{2a} is methoxy; R^{3a} is hydrogen; R^{4a} is methyl; R^{5a} is methyl; R^{6a} is hydrogen and R^{7a} is methyl.

In some embodiments, R^{1a} is methyl; R^{2a} is methoxy; R^{3a} is fluorine; R^{4a} is methyl; R^{5a} is hydrogen; R^{6a} is methyl and R^{7a} is methyl.

In some embodiments, R^{1a} is methyl; R^{2a} is methoxy; R^{3a} is fluorine; R^{4a} is methyl; R^{5a} is hydrogen; R^{6a} is hydrogen and R^{7a} is $-(CH_2)_2OCH_3$.

In some embodiments, R^{1a} is methyl; R^{2a} is methoxy; R^{3a} is fluorine; R^{4a} is methyl; R^{5a} is hydrogen; R^{6a} is hydrogen and R^{7a} is ethyl.

In some embodiments, R^{1a} is methyl; R^{2a} is methoxy; R^{3a} is fluorine; R^{4a} is methyl; R^{5a} is methyl; R^{6a} is hydrogen and R^{7a} is methyl.

5 In some embodiments, R^{1a} is methyl; R^{2a} is ethoxy; R^{3a} is fluorine; R^{4a} is methyl; R^{5a} is hydrogen; R^{6a} is hydrogen and R^{7a} is ethyl.

In some embodiments, R^{1a} is methyl; R^{2a} is ethoxy; R^{3a} is fluorine; R^{4a} is methyl; R^{5a} is hydrogen; R^{6a} is hydrogen and R^{7a} is $-(CH_2)_2OCH_3$.

10 In some embodiments, R^{1a} is methyl; R^{2a} is ethoxy; R^{3a} is fluorine; R^{4a} is methyl; R^{5a} is methyl; R^{6a} is hydrogen and R^{7a} is methyl.

In some embodiments, R^{1a} is methyl; R^{2a} is ethoxy; R^{3a} is fluorine; R^{4a} is methyl; R^{5a} is hydrogen; R^{6a} is methyl and R^{7a} is methyl.

In some embodiments, R^{1a} is methyl; R^{2a} is ethoxy; R^{3a} is fluorine; R^{4a} is methyl; R^{5a} is methyl; R^{6a} is hydrogen and R^{7a} is methyl.

15 In some embodiments, R^{1a} is methyl; R^{2a} is methoxy; R^{3a} is methyl; R^{4a} is methyl; R^{5a} is hydrogen; R^{6a} is methyl and R^{7a} is methyl.

In some embodiments, R^{1a} is methyl; R^{2a} is methoxy; R^{3a} is methyl; R^{4a} is methyl; R^{5a} is hydrogen; R^{6a} is hydrogen and R^{7a} is $-(CH_2)_2OH$.

20 In some embodiments, R^{1a} is methyl; R^{2a} is methoxy; R^{3a} is methyl; R^{4a} is methyl; R^{5a} is hydrogen; R^{6a} is hydrogen and R^{7a} is ethyl.

In some embodiments, R^{1a} is methyl; R^{2a} is methoxy; R^{3a} is methyl; R^{4a} is methyl; R^{5a} is hydrogen; R^{6a} is hydrogen and R^{7a} is $-(CH_2)_2OCH_3$.

In some embodiments, R^{1a} is methyl; R^{2a} is methoxy; R^{3a} is methyl; R^{4a} is methyl; R^{5a} is methyl; R^{6a} is hydrogen and R^{7a} is methyl.

25 In some embodiments, R^{1a} is methyl; R^{2a} is ethoxy; R^{3a} is methyl; R^{4a} is methyl; R^{5a} is hydrogen; R^{6a} is hydrogen and R^{7a} is ethyl.

In some embodiments, R^{1a} is methyl; R^{2a} is ethoxy; R^{3a} is methyl; R^{4a} is methyl; R^{5a} is hydrogen; R^{6a} is hydrogen and R^{7a} is $-(CH_2)_2OCH_3$.

30 In some embodiments, R^{1a} is methyl; R^{2a} is ethoxy; R^{3a} is methyl; R^{4a} is methyl; R^{5a} is methyl; R^{6a} is hydrogen and R^{7a} is methyl.

In some embodiments, R^{1a} is methyl; R^{2a} is ethoxy; R^{3a} is methyl; R^{4a} is methyl; R^{5a} is hydrogen; R^{6a} is methyl and R^{7a} is methyl.

In some embodiments, R^{1a} is methyl; R^{2a} is methoxy; R^{3a} is fluorine; R^{4a} is methyl; R^{5a} is hydrogen; R^{6a} is hydrogen and R^{7a} is methyl.

In some embodiments, R^{1a} is methyl; R^{2a} is methoxy; R^{3a} is hydrogen; R^{4a} is methyl; R^{5a} is hydrogen; R^{6a} is methyl and R^{7a} is methyl.

In some embodiments, R^{1a} is methyl; R^{2a} is methoxy; R^{3a} is hydrogen; R^{4a} is methyl; R^{5a} is hydrogen; R^{6a} is hydrogen and R^{7a} is $-(CH_2)_2OCH_3$.

5 In some embodiments, R^{1a} is methyl; R^{2a} is methoxy; R^{3a} is hydrogen; R^{4a} is methyl; R^{5a} is hydrogen; R^{6a} is hydrogen and R^{7a} is ethyl.

In some embodiments, R^{1a} is methyl; R^{2a} is methoxy; R^{3a} is hydrogen; R^{4a} is methyl; R^{5a} is methyl; R^{6a} is hydrogen and R^{7a} is methyl.

10 In some embodiments, R^{1a} is methyl; R^{2a} is methoxy; R^{3a} is hydrogen; R^{4a} is methyl; R^{5a} is hydrogen; R^{6a} is methyl and R^{7a} is methyl.

In some embodiments, R^{1a} is methyl; R^{2a} is ethoxy; R^{3a} is methyl; R^{4a} is methyl; R^{5a} is hydrogen; R^{6a} is methyl and R^{7a} is methyl.

In some embodiments, R^{1a} is methyl; R^{2a} is methoxy; R^{3a} is hydrogen; R^{4a} is methyl; R^{5a} is hydrogen; R^{6a} is hydrogen and R^{7a} is methyl.

15 In some embodiments, R^{1a} is methyl; R^{2a} is methoxy; R^{3a} is methyl; R^{4a} is methyl; R^{5a} is hydrogen; R^{6a} is hydrogen and R^{7a} is methyl.

In some embodiments, R^{1a} is methyl; R^{2a} is ethoxy; R^{3a} is methyl; R^{4a} is methyl; R^{5a} is hydrogen; R^{6a} is hydrogen and R^{7a} is methyl.

20 In some embodiments, R^{1a} is methyl; R^{2a} is methoxy; R^{3a} is chlorine; R^{4a} is methyl; R^{5a} is hydrogen; R^{6a} is hydrogen and R^{7a} is methyl.

In some embodiments, R^{1a} is methyl; R^{2a} is methoxy; R^{3a} is bromine; R^{4a} is methyl; R^{5a} is hydrogen; R^{6a} is hydrogen and R^{7a} is methyl.

In some embodiments, R^{1a} is methyl; R^{2a} is ethoxy; R^{3a} is fluorine; R^{4a} is methyl; R^{5a} is hydrogen; R^{6a} is hydrogen and R^{7a} is methyl.

25 In some embodiments, R^{1a} is methyl; R^{2a} is ethoxy; R^{3a} is chlorine; R^{4a} is methyl; R^{5a} is hydrogen; R^{6a} is hydrogen and R^{7a} is methyl.

In some embodiments, R^{1a} is methyl; R^{2a} is methyl; R^{3a} is hydrogen; R^{4a} is methyl; R^{5a} is hydrogen; R^{6a} is hydrogen and R^{7a} is methyl.

30 In some embodiments, R^{1a} is methyl; R^{2a} is methoxy; R^{3a} is methyl; R^{4a} is ethyl; R^{5a} is hydrogen; R^{6a} is hydrogen and R^{7a} is methyl.

In some embodiments, R^{1a} is methyl; R^{2a} is methoxy; R^{3a} is methyl; R^{4a} is $-CH_2OCH_3$; R^{5a} is hydrogen; R^{6a} is hydrogen and R^{7a} is methyl.

In some embodiments, R^{1a} is methyl; R^{2a} is methoxy; R^{3a} is hydrogen; R^{4a} is ethyl; R^{5a} is hydrogen; R^{6a} is hydrogen and R^{7a} is methyl.

In some embodiments, R^{1a} is methyl; R^{2a} is methyl; R^{3a} is fluorine; R^{4a} is ethyl; R^{5a} is hydrogen; R^{6a} is hydrogen and R^{7a} is methyl.

In some embodiments, R^{1a} is methyl; R^{2a} is methoxy; R^{3a} is fluorine; R^{4a} is ethyl; R^{5a} is hydrogen; R^{6a} is hydrogen and R^{7a} is methyl.

5 In some embodiments, R^{1a} is methyl; R^{2a} is methyl; R^{3a} is fluorine; R^{4a} is -CH₂OCH₃; R^{5a} is hydrogen; R^{6a} is hydrogen and R^{7a} is methyl.

In some embodiments, R^{1a} is methyl; R^{2a} is ethyl; R^{3a} is methyl; R^{4a} is methyl; R^{5a} is hydrogen; R^{6a} is hydrogen and R^{7a} is methyl.

10 In some embodiments, R^{1a} is methyl; R^{2a} is methyl; R^{3a} is fluorine; R^{4a} is methyl; R^{5a} is hydrogen; R^{6a} is hydrogen and R^{7a} is methyl.

In some embodiments, R^{1a} is methyl; R^{2a} is methyl; R^{3a} is hydrogen; R^{4a} is methyl; R^{5a} is hydrogen; R^{6a} is hydrogen and R^{7a} is methyl.

In some embodiments, R^{1a} is ethyl; R^{2a} is methoxy; R^{3a} is methyl; R^{4a} is methyl; R^{5a} is hydrogen; R^{6a} is hydrogen and R^{7a} is methyl.

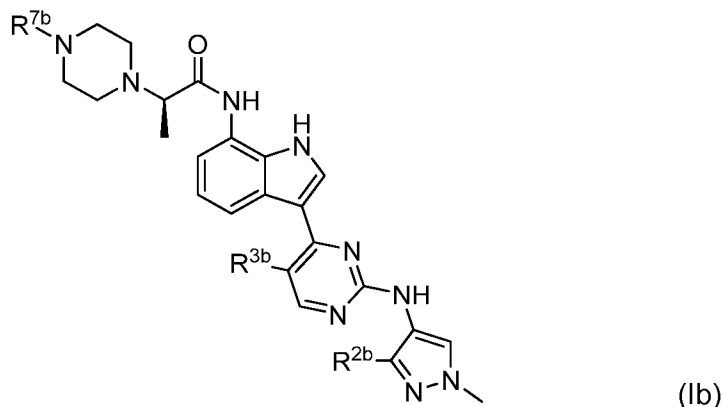
15 In some embodiments, R^{1a} is ethyl; R^{2a} is ethoxy; R^{3a} is methyl; R^{4a} is methyl; R^{5a} is hydrogen; R^{6a} is hydrogen and R^{7a} is methyl.

In some embodiments, R^{1a} is methyl; R^{2a} is methoxy; R^{3a} is hydrogen; R^{4a} is -CH₂OCH₃; R^{5a} is hydrogen; R^{6a} is hydrogen and R^{7a} is methyl.

20 In some embodiments, R^{1a} is methyl; R^{2a} is ethoxy; R^{3a} is methyl; R^{4a} is ethyl; R^{5a} is hydrogen; R^{6a} is hydrogen and R^{7a} is methyl.

In some embodiments, R^{1a} is methyl; R^{2a} is ethoxy; R^{3a} is methyl; R^{4a} is -CH₂OCH₃; R^{5a} is hydrogen; R^{6a} is hydrogen and R^{7a} is methyl.

In one embodiment, the compounds of formula (I) are compounds of formula (Ib):



25 wherein

R^{2b} is selected from methyl, ethyl, methoxy and ethoxy;

R^{3b} is selected from hydrogen, chlorine, fluorine, bromine and methyl; and

R^{7b} is selected from methyl, ethyl, -(CH₂)₂OH and -(CH₂)₂OCH₃, or a pharmaceutically acceptable salt thereof.

In some embodiments, R^{2b} is methoxy; R^{3b} is hydrogen, fluorine, methyl, chlorine or bromine; and R^{7b} is methyl, ethyl, -(CH₂)₂OCH₃ or -(CH₂)₂OH.

5 In some embodiments, R^{2b} is ethoxy; R^{3b} is fluorine, methyl or chlorine; and R^{7b} is ethyl, methyl or -(CH₂)₂OCH₃.

In some embodiments, R^{2b} is methyl; R^{3b} is hydrogen, methyl or fluorine; and R^{7b} is methyl.

In some embodiments, R^{2b} is ethyl; R^{3b} is methyl; and R^{7b} is methyl.

10 In some embodiments, R^{3b} is hydrogen; R^{2b} is methoxy or methyl; and R^{7b} is methyl, -(CH₂)₂OCH₃ or ethyl.

In some embodiments, R^{3b} is fluorine; R^{2b} is methoxy or ethoxy; R^{7b} is methyl, ethyl or -(CH₂)₂OCH₃.

15 In some embodiments, R^{3b} is methyl; R^{2b} is methoxy, ethoxy, methyl or ethyl; and R^{7b} is methyl, ethyl, -(CH₂)₂OCH₃ or -(CH₂)₂OH.

In some embodiments, R^{3b} is chlorine; R^{2b} is methoxy or ethoxy; and R^{7b} is methyl.

In some embodiments, R^{3b} is bromine; R^{2b} is methoxy; and R^{7b} is methyl.

In some embodiments, R^{7b} is methyl; R^{2b} is methyl, ethyl, methoxy or ethoxy; and R^{3b} is hydrogen, chlorine, fluorine, bromine or methyl.

20 In some embodiments, R^{7b} is ethyl; R^{2b} is methoxy or ethoxy; and R^{3b} is fluorine, methyl or hydrogen.

In some embodiments, R^{7b} is -(CH₂)₂OCH₃; R^{2b} is methoxy or ethoxy; and R^{3b} is fluorine, methyl or hydrogen.

In some embodiments, R^{7b} is -(CH₂)₂OH; R^{2b} is methoxy; and R^{3a} is methyl.

25 In some embodiments, R^{2b} is methoxy; R^{3b} is fluorine; and R^{7b} is -(CH₂)₂OCH₃.

In some embodiments, R^{2b} is methoxy; R^{3b} is fluorine; and R^{7a} is ethyl.

In some embodiments, R^{2b} is ethoxy; R^{3b} is fluorine; and R^{7a} is ethyl.

In some embodiments, R^{2b} is ethoxy; R^{3b} is fluorine; and R^{7b} is -(CH₂)₂OCH₃.

In some embodiments, R^{2b} is methoxy; R^{3b} is methyl; and R^{7b} is -(CH₂)₂OH.

30 In some embodiments, R^{2b} is methoxy; R^{3b} is methyl; and R^{7b} is ethyl.

In some embodiments, R^{2b} is methoxy; R^{3b} is methyl; and R^{7b} is -(CH₂)₂OCH₃.

In some embodiments, R^{2b} is ethoxy; R^{3b} is methyl; and R^{7b} is ethyl. In some embodiments, R^{2b} is ethoxy; R^{3b} is methyl; and R^{7b} is -(CH₂)₂OCH₃.

In some embodiments, R^{2b} is methoxy; R^{3b} is fluorine; and R^{7b} is methyl.

In some embodiments, R^{2b} is methoxy; R^{3b} is hydrogen; and R^{7b} is $-(CH_2)_2OCH_3$.

In some embodiments, R^{2b} is methoxy; R^{3b} is hydrogen; and R^{7b} is ethyl.

In some embodiments, R^{2b} is methoxy; R^{3b} is hydrogen; and R^{7b} is methyl.

In some embodiments, R^{2b} is methoxy; R^{3b} is methyl; and R^{7b} is methyl.

5 In some embodiments, R^{2b} is ethoxy; R^{3b} is methyl; and R^{7b} is methyl.

In some embodiments, R^{2b} is methoxy; R^{3b} is chlorine; and R^{7b} is methyl.

In some embodiments, R^{2b} is methoxy; R^{3b} is bromine; and R^{7b} is methyl.

In some embodiments, R^{2b} is ethoxy; R^{3b} is fluorine; and R^{7b} is methyl.

In some embodiments, R^{2b} is ethoxy; R^{3b} is chlorine; and R^{7b} is methyl.

10 In some embodiments, R^{2b} is methyl; R^{3b} is hydrogen; and R^{7b} is methyl.

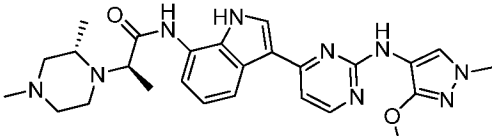
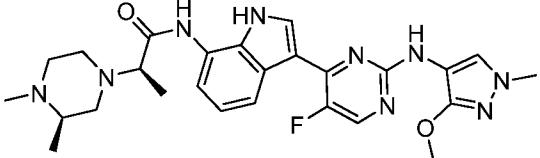
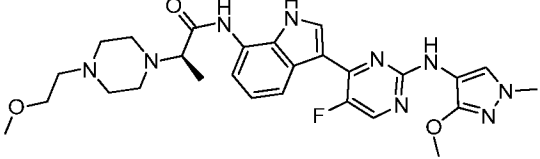
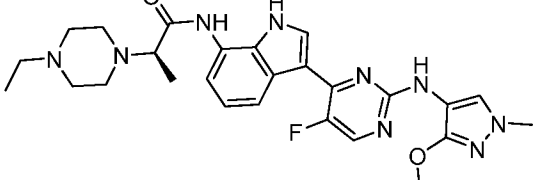
In some embodiments, R^{2b} is ethyl; R^{3b} is methyl; and R^{7b} is methyl.

In some embodiments, R^{2b} is methyl; R^{3b} is fluorine; and R^{7b} is methyl.

In some embodiments, R^{2b} is methyl; R^{3b} is hydrogen; and R^{7b} is methyl.

15 In some embodiments, disclosed are the compounds of Table 1, or a pharmaceutically acceptable salt thereof:

Table 1

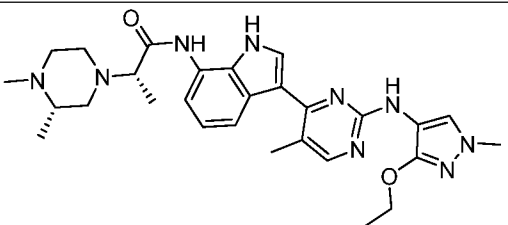
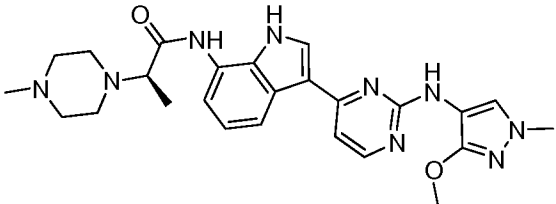
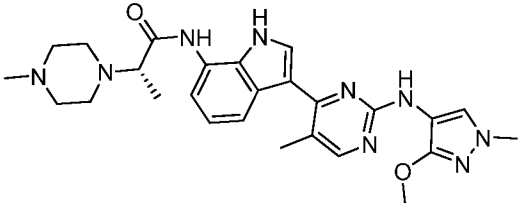
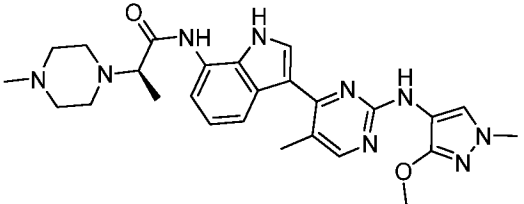
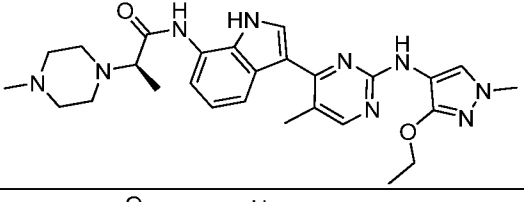
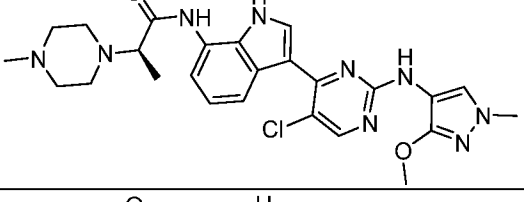
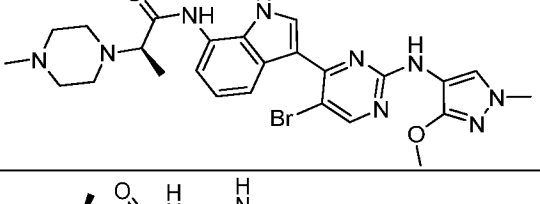
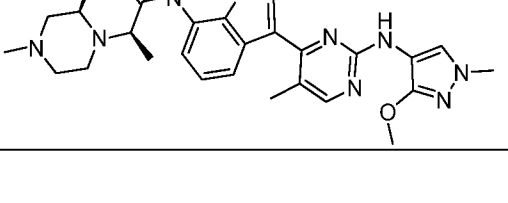
Example No.	Chemical Structure	Name
Example 1		(2 <i>R</i>)-2-[(2 <i>S</i>)-2,4-dimethylpiperazin-1-yl]- <i>N</i> -(3-{2-[(3-methoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]pyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)propanamide
Example 2		(2 <i>R</i>)-2-[(3 <i>R</i>)-3,4-dimethylpiperazin-1-yl]- <i>N</i> -(3-{5-fluoro-2-[(3-methoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]pyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)propanamide
Example 3		(2 <i>R</i>)- <i>N</i> -(3-{5-fluoro-2-[(3-methoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]pyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)-2-[4-(2-methoxyethyl)piperazin-1-yl]propanamide
Example 4		(2 <i>R</i>)-2-(4-ethylpiperazin-1-yl)- <i>N</i> -(3-{5-fluoro-2-[(3-methoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]pyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)propanamide

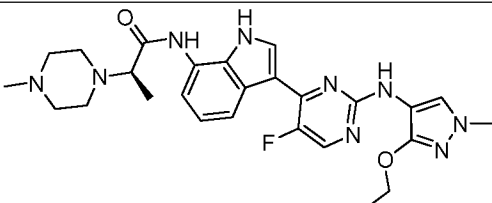
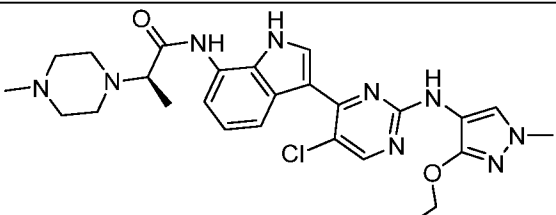
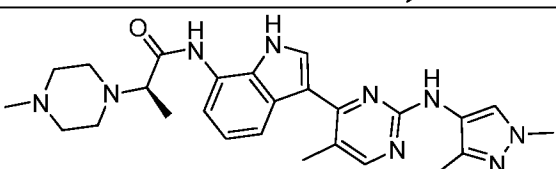
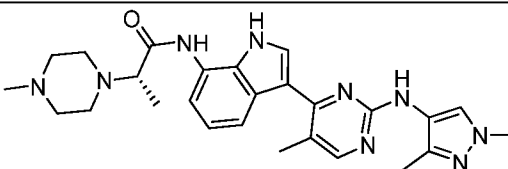
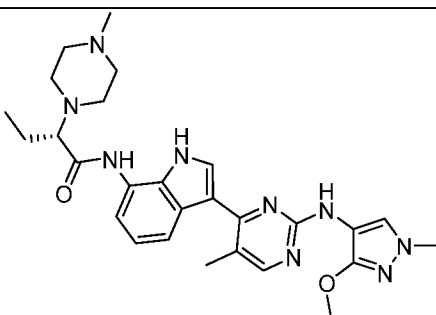
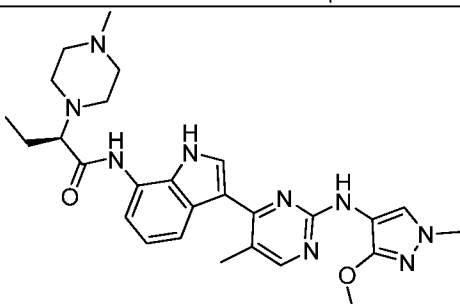
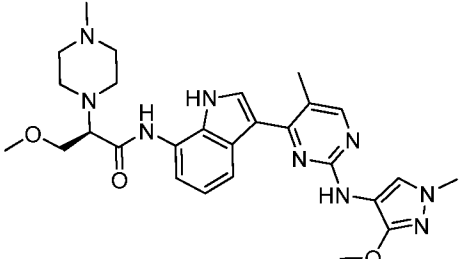
Example 5		(2 <i>R</i>)-2-[(3 <i>S</i>)-3,4-dimethylpiperazin-1-yl]- <i>N</i> -(3-{5-fluoro-2-[(3-methoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]pyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)propanamide
Example 6		(2 <i>R</i>)-2-[(2 <i>R</i>)-2,4-dimethylpiperazin-1-yl]- <i>N</i> -(3-{5-fluoro-2-[(3-methoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]pyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)propanamide
Example 7a		(2 <i>R</i>)-2-[(2 <i>S</i>)-2,4-dimethylpiperazin-1-yl]- <i>N</i> -(3-{5-fluoro-2-[(3-methoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]pyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)propanamide
Example 7b		(2 <i>S</i>)-2-[(2 <i>S</i>)-2,4-dimethylpiperazin-1-yl]- <i>N</i> -(3-{5-fluoro-2-[(3-methoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]pyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)propanamide
Example 8		(2 <i>R</i>)- <i>N</i> -(3-{2-[(3-ethoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]-5-fluoropyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)-2-(4-ethylpiperazin-1-yl)propanamide
Example 9		(2 <i>R</i>)- <i>N</i> -(3-{2-[(3-ethoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]-5-fluoropyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)-2-[4-(2-methoxyethyl)piperazin-1-yl]propanamide
Example 10a		(2 <i>R</i>)-2-[(2 <i>S</i>)-2,4-dimethylpiperazin-1-yl]- <i>N</i> -(3-{2-[(3-ethoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]-5-fluoropyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)propanamide
Example 10b		(2 <i>S</i>)-2-[(2 <i>S</i>)-2,4-dimethylpiperazin-1-yl]- <i>N</i> -(3-{2-[(3-ethoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]-5-fluoropyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)propanamide

Example 11a		(2 <i>R</i>)-2-[(3 <i>S</i>)-3,4-dimethylpiperazin-1-yl]- <i>N</i> -(3-{2-[(3-ethoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]-5-fluoropyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)propanamide
Example 11b		(2 <i>S</i>)-2-[(3 <i>S</i>)-3,4-dimethylpiperazin-1-yl]- <i>N</i> -(3-{2-[(3-ethoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]-5-fluoropyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)propanamide
Example 12		(2 <i>R</i>)-2-[(2 <i>R</i>)-2,4-dimethylpiperazin-1-yl]- <i>N</i> -(3-{2-[(3-ethoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]-5-fluoropyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)propanamide
Example 13		(2 <i>R</i>)-2-[(3 <i>R</i>)-3,4-dimethylpiperazin-1-yl]- <i>N</i> -(3-{2-[(3-ethoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]-5-fluoropyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)propanamide
Example 14		(2 <i>R</i>)-2-[(3 <i>S</i>)-3,4-dimethylpiperazin-1-yl]- <i>N</i> -(3-{2-[(3-methoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)propanamide
Example 15		(2 <i>R</i>)-2-[4-(2-hydroxyethyl)piperazin-1-yl]- <i>N</i> -(3-{2-[(3-methoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)propanamide
Example 16		(2 <i>R</i>)-2-(4-ethylpiperazin-1-yl)- <i>N</i> -(3-{2-[(3-methoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)propanamide
Example 17		(2 <i>R</i>)-2-[4-(2-methoxyethyl)piperazin-1-yl]- <i>N</i> -(3-{2-[(3-methoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)propanamide

Example 18		(2 <i>R</i>)-2-[(3 <i>R</i>)-3,4-dimethylpiperazin-1-yl]- <i>N</i> -(3-{2-[(3-methoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)propanamide
Example 19		(2 <i>R</i>)-2-[(2 <i>S</i>)-2,4-dimethylpiperazin-1-yl]- <i>N</i> -(3-{2-[(3-methoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)propanamide
Example 20		(2 <i>R</i>)- <i>N</i> -(3-{2-[(3-ethoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)-2-(4-ethylpiperazin-1-yl)propanamide
Example 21		(2 <i>R</i>)- <i>N</i> -(3-{2-[(3-ethoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)-2-[4-(2-methoxyethyl)piperazin-1-yl]propanamide
Example 22		(2 <i>R</i>)-2-[(2 <i>S</i>)-2,4-dimethylpiperazin-1-yl]- <i>N</i> -(3-{2-[(3-ethoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)propanamide
Example 23		(2 <i>R</i>)-2-[(2 <i>R</i>)-2,4-dimethylpiperazin-1-yl]- <i>N</i> -(3-{2-[(3-ethoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)propanamide
Example 24		(2 <i>R</i>)-2-[(3 <i>R</i>)-3,4-dimethylpiperazin-1-yl]- <i>N</i> -(3-{2-[(3-ethoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)propanamide

Example 25		(2 <i>R</i>)- <i>N</i> -(3-{5-Fluoro-2-[(3-methoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]pyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide
Example 26		(2 <i>R</i>)-2-[(3 <i>S</i>)-3,4-dimethylpiperazin-1-yl]- <i>N</i> -(3-{2-[(3-methoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]pyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)propanamide
Example 27		(2 <i>R</i>)-2-[4-(2-methoxyethyl)piperazin-1-yl]- <i>N</i> -(3-{2-[(3-methoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]pyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)propanamide
Example 28		(2 <i>R</i>)-2-(4-ethylpiperazin-1-yl)- <i>N</i> -(3-{2-[(3-methoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]pyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)propanamide
Example 29		(2 <i>R</i>)-2-[(2 <i>R</i>)-2,4-dimethylpiperazin-1-yl]- <i>N</i> -(3-{2-[(3-methoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]pyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)propanamide
Example 30		(2 <i>R</i>)-2-[(3 <i>R</i>)-3,4-dimethylpiperazin-1-yl]- <i>N</i> -(3-{2-[(3-methoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]pyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)propanamide
Example 31a		(2 <i>R</i>)-2-[(3 <i>S</i>)-3,4-dimethylpiperazin-1-yl]- <i>N</i> -(3-{2-[(3-ethoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)propanamide

Example 31b		(2 <i>S</i>)-2-[(3 <i>S</i>)-3,4-dimethylpiperazin-1-yl]- <i>N</i> -(3-{2-[(3-ethoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)propanamide
Example 32		(2 <i>R</i>)- <i>N</i> -(3-{2-[(3-methoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]pyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide
Example 33		(2 <i>S</i>)- <i>N</i> -(3-{2-[(3-methoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide
Example 34		(2 <i>R</i>)- <i>N</i> -(3-{2-[(3-Methoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide
Example 35		(2 <i>R</i>)- <i>N</i> -(3-{2-[(3-ethoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide
Example 36		(2 <i>R</i>)- <i>N</i> -(3-{5-chloro-2-[(3-methoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]pyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide
Example 37		(2 <i>R</i>)- <i>N</i> -(3-{5-bromo-2-[(3-methoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]pyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide
Example 38		(2 <i>R</i>)-2-[(2 <i>R</i>)-2,4-dimethylpiperazin-1-yl]- <i>N</i> -(3-{2-[(3-methoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)propanamide

Example 39		(2 <i>R</i>)- <i>N</i> -(3-{2-[(3-ethoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]-5-fluoropyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide
Example 40		(2 <i>R</i>)- <i>N</i> -(3-{5-chloro-2-[(3-ethoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]pyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide
Example 41		(2 <i>R</i>)- <i>N</i> -(3-{2-[(1,3-dimethyl-1 <i>H</i> -pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide
Example 42		(2 <i>S</i>)- <i>N</i> -(3-{2-[(1,3-dimethyl-1 <i>H</i> -pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide
Example 43		(2 <i>S</i>)- <i>N</i> -(3-{2-[(3-Methoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)-2-(4-methylpiperazin-1-yl)butanamide
Example 44		(2 <i>R</i>)- <i>N</i> -(3-{2-[(3-Methoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)-2-(4-methylpiperazin-1-yl)butanamide
Example 45		(2 <i>R</i>)-3-methoxy- <i>N</i> -(3-{2-[(3-methoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide

Example 46		(2 <i>S</i>)-3-methoxy- <i>N</i> -(3-{2-[(3-methoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide
Example 47		(2 <i>R</i>)- <i>N</i> -(3-{2-[(3-Methoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]pyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)-2-(4-methylpiperazin-1-yl)butanamide
Example 48		(2 <i>S</i>)- <i>N</i> -(3-{2-[(3-Methoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]pyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)-2-(4-methylpiperazin-1-yl)butanamide
Example 49		(2 <i>R</i>)- <i>N</i> -(3-{2-[(1,3-dimethyl-1 <i>H</i> -pyrazol-4-yl)amino]-5-fluoropyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)-2-(4-methylpiperazin-1-yl)butanamide
Example 50		(2 <i>S</i>)- <i>N</i> -(3-{2-[(1,3-dimethyl-1 <i>H</i> -pyrazol-4-yl)amino]-5-fluoropyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)-2-(4-methylpiperazin-1-yl)butanamide
Example 51		(2 <i>R</i>)- <i>N</i> -(3-{5-fluoro-2-[(3-methoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]pyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)-2-(4-methylpiperazin-1-yl)butanamide

Example 52		(2 <i>S</i>)- <i>N</i> -(3-{5-fluoro-2-[(3-methoxy-1 <i>H</i> -pyrazol-4-yl)amino]pyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)-2-(4-methylpiperazin-1-yl)butanamide
Example 53		(2 <i>R</i>)- <i>N</i> -(3-{2-[(1,3-dimethyl-1 <i>H</i> -pyrazol-4-yl)amino]-5-fluoropyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)-3-methoxy-2-(4-methylpiperazin-1-yl)propanamide
Example 54		(2 <i>S</i>)- <i>N</i> -(3-{2-[(1,3-dimethyl-1 <i>H</i> -pyrazol-4-yl)amino]-5-fluoropyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)-3-methoxy-2-(4-methylpiperazin-1-yl)propanamide
Example 55		(2 <i>R</i>)- <i>N</i> -(3-{2-[(3-ethyl-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide
Example 56		(2 <i>R</i>)- <i>N</i> -(3-{2-[(1,3-dimethyl-1 <i>H</i> -pyrazol-4-yl)amino]-5-fluoropyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide
Example 57		(2 <i>R</i>)- <i>N</i> -(3-{2-[(1,3-dimethyl-1 <i>H</i> -pyrazol-4-yl)amino]pyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide
Example 58		(2 <i>R</i>)- <i>N</i> -(3-{2-[(1-ethyl-3-methoxy-1 <i>H</i> -pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide
Example 59		(2 <i>R</i>)- <i>N</i> -(3-{2-[(3-ethoxy-1-ethyl-1 <i>H</i> -pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide

Example 60		(2 <i>R</i>)-3-Methoxy- <i>N</i> -(3-{2-[(3-methoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]pyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide
Example 61		(2 <i>S</i>)-3-Methoxy- <i>N</i> -(3-{2-[(3-methoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]pyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide
Example 62		(2 <i>R</i>)- <i>N</i> -(3-{2-[(3-ethoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)-2-(4-methylpiperazin-1-yl)butanamide
Example 63		(2 <i>S</i>)- <i>N</i> -(3-{2-[(3-ethoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)-2-(4-methylpiperazin-1-yl)butanamide
Example 64		(2 <i>R</i>)- <i>N</i> -(3-{2-[(3-ethoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)-3-methoxy-2-(4-methylpiperazin-1-yl)propanamide
Example 65		(2 <i>S</i>)- <i>N</i> -(3-{2-[(3-ethoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)-3-methoxy-2-(4-methylpiperazin-1-yl)propanamide

In one aspect, the disclosed compounds are obtainable by any process described in the Examples. In one embodiment, the disclosed are the intermediate compounds described in the Examples.

The language "pharmaceutically acceptable salt" includes acid addition or base salts that
5 retain the biological effectiveness and properties of the compounds of Formula (I), (Ia), (Ib) and Table 1 and, which typically are not biologically or otherwise undesirable. In many cases, Formula (I), (Ia), (Ib) and Table 1 are capable of forming acid and/or base salts by virtue of the presence of amino and/or carboxyl groups or groups similar thereto.

Pharmaceutically acceptable acid addition salts can be formed with inorganic acids and
10 organic acids, *e.g.*, acetate, aspartate, benzoate, besylate, bromide/hydrobromide, bicarbonate/carbonate, bisulfate/sulfate, camphorsulfonate, chloride/hydrochloride, chlorotheophyllonate, citrate, ethandisulfonate, fumarate, gluceptate, gluconate, glucuronate, hippurate, hydroiodide/iodide, isethionate, lactate, lactobionate, laurylsulfate, malate, maleate, malonate, mandelate, mesylate, methylsulphate, napadisylate, naphthoate, napsylate,
15 nicotinate, nitrate, octadecanoate, oleate, oxalate, palmitate, palmoate, phosphate/hydrogen phosphate/dihydrogen phosphate, polygalacturonate, propionate, stearate, succinate, subsalicylate, tartrate, tosylate, trimesate and trifluoroacetate salts. Inorganic acids from which salts can be derived include, for example, hydrochloric acid, hydrobromic acid, sulfuric acid, nitric acid, phosphoric acid, and the like. Organic acids from which salts can be derived include,
20 for example, acetic acid, propionic acid, glycolic acid, oxalic acid, maleic acid, malonic acid, succinic acid, fumaric acid, tartaric acid, trimesic acid, citric acid, benzoic acid, mandelic acid, methanesulfonic acid, napadisyllic acid, ethanesulfonic acid, toluenesulfonic acid, trifluoroacetic acid, sulfosalicylic acid, and the like.

Pharmaceutically acceptable base addition salts can be formed with inorganic and organic
25 bases. Inorganic bases from which salts can be derived include, for example, ammonium salts and metals from columns I to XII of the periodic table. In certain embodiments, the salts are derived from sodium, potassium, ammonium, calcium, magnesium, iron, silver, zinc, and copper; particularly suitable salts include ammonium, potassium, sodium, calcium and magnesium salts. Organic bases from which salts can be derived include, for example, primary,
30 secondary, and tertiary amines, substituted amines including naturally occurring substituted amines, cyclic amines, basic ion exchange resins, and the like. Certain organic amines include isopropylamine, benzathine, choline, diethanolamine, diethylamine, lysine, meglumine, piperazine and tromethamine.

The pharmaceutically acceptable salts of the compounds Formula (I), (Ia), (Ib) and Table 1 can be synthesized from a basic or acidic moiety, by conventional chemical methods. Generally, such salts can be prepared by reacting free acid forms of these compounds with a stoichiometric amount of the appropriate base (such as Na⁺, Ca²⁺, Mg²⁺, or K⁺ hydroxide, carbonate, bicarbonate or the like), or by reacting free base forms of these compounds with a stoichiometric amount of the appropriate acid. Such reactions are typically carried out in water or in an organic solvent, or in a mixture of the two. Generally, use of non-aqueous media like ether, ethyl acetate, ethanol, isopropanol, or acetonitrile is desirable, where practicable. Lists of additional suitable salts can be found, *e.g.*, in "Remington's Pharmaceutical Sciences," 20th ed., Mack Publishing Company, Easton, Pa., (1985); and in "Handbook of Pharmaceutical Salts: Properties, Selection, and Use" by Stahl and Wermuth (Wiley-VCH, Weinheim, Germany, 2002).

Any formula given herein is also intended to represent unlabeled forms as well as isotopically labeled forms for the compounds of Formula (I), (Ia), (Ib) and Table 1. Isotopically labeled compounds have structures depicted by the formulas given herein except that one or more atoms are replaced by an atom having a selected atomic mass or mass number. Examples of isotopes that can be incorporated into the compounds of Formula (I), (Ia), (Ib) and Table 1 include isotopes of hydrogen, carbon, nitrogen, oxygen, phosphorous, fluorine and chlorine, such as ²H, ³H, ¹¹C, ¹³C, ¹⁴C, ¹⁵N, ¹⁶F, ³¹P, ³²P, ³⁵S, ³⁶Cl and ¹²⁵I. The compounds of Formula (I), (Ia), (Ib) and Table 1 may include various isotopically labeled compounds into which radioactive isotopes, such as ²H, ³H, ¹³C and ¹⁴C, are present. Isotopically labeled compounds of formula (I), (Ia) and (Ib) can generally be prepared by convention techniques known to those skilled in the art or by processes analogous to those described in the accompanying Examples using appropriate isotopically labeled reagents in place of the non-labeled reagents previously employed.

The compounds of formula (I), (Ia), (Ib) and Table 1 may have different isomeric forms. The language "optical isomer" or "stereoisomer" refers to any of the various stereoisomeric configurations which may exist for a given compound of formula (I), (Ia), (Ib) and Table 1. It is understood that a substituent may be attached at a chiral center of a carbon atom and, therefore, the disclosed compounds include enantiomers, diastereomers and racemates. The term "enantiomer" includes pairs of stereoisomers that are non-superimposable mirror images of each other. A 1:1 mixture of a pair of enantiomers is a racemic mixture. The term is used to designate a racemic mixture where appropriate. The terms "diastereomers" or "diastereoisomers" include stereoisomers that have at least two asymmetric atoms, but which

are not mirror images of each other. The absolute stereochemistry is specified according to the Cahn-Ingold-Prelog R-S system. When a compound is a pure enantiomer, the stereochemistry at each chiral center may be specified by either *R* or *S*. Resolved compounds whose absolute configuration is unknown can be designated (+) or (-) depending on the direction (dextro- or levorotatory) which they rotate plane polarized light at the wavelength of the sodium D line. Certain of the compounds of Formula (I), (Ia), (Ib) and Table 1 contain one or more asymmetric centers or axes and may thus give rise to enantiomers, diastereomers or other stereoisomeric forms that may be defined, in terms of absolute stereochemistry, as (*R*)- or (*S*)-. The present disclosure is meant to include all such possible isomers, including racemic mixtures, optically pure forms and intermediate mixtures. Optically active (*R*)- and (*S*)-isomers may be prepared using chiral synthons or chiral reagents, or resolved using conventional techniques well known in the art, such as chiral HPLC.

Solid Forms

In some embodiments, disclosed are solid forms of the compounds of Formula (I), (Ia) and (Ib), or a pharmaceutically acceptable salt thereof. The term "solid form" includes polymorphs, crystalline salts, solvates, hydrates and amorphous forms of the compounds of Formula (I), (Ia) and (Ib). In some embodiments, disclosed are solid forms of (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide, or a pharmaceutically acceptable salt thereof. The term "polymorph" includes crystalline materials that have the same chemical composition but different molecular packing. The term language "crystalline salt" includes crystalline structures with the same chemical materials, but incorporating acid or base addition salts within the molecular packing of the crystalline structure. The term "solvate" includes crystalline structures of the same chemical material, but incorporating molecules of solvent within the molecular packing of the crystalline structure. The term "hydrates" includes crystalline structures of the same chemical material, but incorporating molecules of water within the molecular packing of the crystalline structure. The language "amorphous form" includes compounds of the same molecular material but without the molecular order of a crystalline structure (e.g., polymorph, crystalline salt, solvate or hydrate) of the same molecular material.

It is generally known that solid materials may be characterized using conventional techniques such as X-Ray Powder Diffraction (XRPD), Differential Scanning Calorimetry (DSC), Thermal Gravimetric Analysis (TGA), Diffuse Reflectance Infrared Fourier Transform (DRIFT) spectroscopy, Near Infrared (NIR) spectroscopy, solution and/or solid state nuclear magnetic

resonance spectroscopy. The water content of such solid materials may be determined by Karl Fischer analysis.

The solid forms described herein provide XRPD patterns substantially the same as the XRPD patterns shown in the Figures, and have the various 2-theta (2θ) values as shown in the Tables included herein. One skilled in the art will understand that an XRPD pattern or diffractogram may be obtained which has one or more measurement errors depending on the recording conditions, such as the equipment or machine used. Similarly, it is generally known that intensities in an XRPD pattern may fluctuate depending on measurement conditions or sample preparation as a result of preferred orientation. Persons skilled in the art of XRPD will further realize that the relative intensity of peaks can also be affected by, for example, grains above 30 μm in size and non-unitary aspect ratios. The skilled person understands that the position of reflections can be affected by the precise height at which the sample sits in the diffractometer, and also the zero calibration of the diffractometer. The surface planarity of the sample may also have a small effect.

As a result of these considerations, the diffraction pattern data presented are not to be taken as absolute values (Jenkins, R & Snyder, R.L. 'Introduction to X-Ray Powder Diffractometry' John Wiley & Sons 1996; Bunn, C.W. (1948), 'Chemical Crystallography', Clarendon Press, London; Klug, H. P. & Alexander, L. E. (1974), 'X-Ray Diffraction Procedures'). It should also be understood that the solid forms embodied herein are not limited to those that provide XRPD patterns that are identical to the XRPD pattern shown in the Figures, and any solid forms providing XRPD patterns substantially the same as those shown in the Figures fall within the scope of the corresponding embodiment. A person skilled in the art of XRPD is able to judge the substantial identity of XRPD patterns. Generally, a measurement error of a diffraction angle in an XRPD is approximately $2\theta (\pm 0.2^\circ)$, and such degree of a measurement error should be taken into account when considering the X-ray powder diffraction pattern in the Figures and when reading data contained in the Tables included herein.

A person skilled in the art also understands that the value or range of values observed in a particular compound's DSC thermogram will show variation between batches of different purities. Therefore, whilst for one compound the range may be small, for others the range may be quite large. Generally, a measurement error of a diffraction angle in DSC thermal events is approximately plus or minus 5 $^\circ\text{C}$, and such degree of a measurement error should be taken into account when considering the DSC data included herein. TGA thermograms show similar variations, such that a person skilled in the art recognizes that measurement errors should be taken into account when judging substantial identity of TGA thermograms.

In some embodiments, disclosed is a solid form of (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide, or a pharmaceutically acceptable salt thereof.

In some embodiments, disclosed is an amorphous form of (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide, or a pharmaceutically acceptable salt thereof.

Form A

In some embodiments, disclosed is Form A (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide.

In some embodiments, Form A (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide has an XRPD pattern comprising at least one peak expressed as 2θ ($\pm 0.2^\circ$) selected from the peaks listed in Table 17.

In some embodiments, Form A (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide has an XRPD pattern substantially similar to **Figure 1**.

In some embodiments, Form A (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide has a DSC thermogram comprising an endotherm with a desolvation onset at about 110 °C and a peak at about 113 °C.

In some embodiments, Form A (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide has a DSC thermogram substantially similar to **Figure 2**.

In some embodiments, Form A (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide has a TGA thermogram exhibiting a mass loss of about 7.8% upon heating from about 25 °C to about 150 °C.

In some embodiments, Form A (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide has a TGA thermogram substantially similar to **Figure 2**.

Form B

In some embodiments, disclosed is Form B (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide.

In some embodiments, Form B (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide has an XRPD pattern comprising at least one peak expressed as 2θ ($\pm 0.2^\circ$) selected from the peaks listed in Table 18.

In some embodiments, Form B (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide has an XRPD pattern substantially similar to **Figure 3**.

In some embodiments, Form B (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide has a DSC thermogram comprising an endotherm with a desolvation onset at about 112 °C and a peak at about 117 °C.

In some embodiments, Form B (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide has a DSC thermogram substantially similar to **Figure 4**.

In some embodiments, Form B (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide has a TGA thermogram exhibiting a mass loss of about 10.0% upon heating from about 25 °C to about 200 °C.

In some embodiments, Form B (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide has a TGA thermogram substantially similar to **Figure 4**.

Form C

In some embodiments, disclosed is Form C (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide.

In some embodiments, Form C (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide has an XRPD pattern comprising at least one peak expressed as 2θ ($\pm 0.2^\circ$) selected from the peaks listed in Table 19.

In some embodiments, Form C (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide has an XRPD pattern substantially similar to **Figure 5**.

In some embodiments, Form C (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide has a DSC thermogram comprising an endotherm with a desolvation onset at about 112 °C and a peak at about 114 °C.

In some embodiments, Form C (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide has a DSC thermogram substantially similar to **Figure 6**.

In some embodiments, Form C (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide has a TGA thermogram exhibiting a mass loss of about 9.2% upon heating from about 25 °C to about 175 °C.

In some embodiments, Form C (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide has a TGA thermogram substantially similar to **Figure 6**.

Form D

In some embodiments, disclosed is Form D (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide.

In some embodiments, Form D (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide has an XRPD pattern comprising at least one peak expressed as 2θ ($\pm 0.2^\circ$) at about 21.8°.

In some embodiments, Form D (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide has an XRPD pattern comprising at least one peak expressed as 2θ ($\pm 0.2^\circ$) at about 6.4°.

In some embodiments, Form D (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide has an XRPD pattern comprising at least one peak expressed as 2θ ($\pm 0.2^\circ$) at about 16.6°.

In some embodiments, Form D (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide has an XRPD pattern comprising at least one peak expressed as 2θ ($\pm 0.2^\circ$) at about 8.9°.

In some embodiments, Form D (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide has an XRPD pattern comprising at least one peak expressed as 2θ ($\pm 0.2^\circ$) at about 8.1° .

5 In some embodiments, Form D (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide has an XRPD pattern comprising at least one peak expressed as 2θ ($\pm 0.2^\circ$) selected from about 21.8° and 6.4° .

10 In some embodiments, Form D (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide has an XRPD pattern comprising at least one peak expressed as 2θ ($\pm 0.2^\circ$) selected from about 21.8° and 16.6° .

15 In some embodiments, Form D (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide has an XRPD pattern comprising at least one peak expressed as 2θ ($\pm 0.2^\circ$) selected from about 21.8° and 8.9° .

In some embodiments, Form D (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide has an XRPD pattern comprising at least one peak expressed as 2θ ($\pm 0.2^\circ$) selected from about 21.8° and 8.1° .

20 In some embodiments, Form D (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide has an XRPD pattern comprising at least one peak expressed as 2θ ($\pm 0.2^\circ$) selected from about 6.4° and 16.6° .

25 In some embodiments, Form D (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide has an XRPD pattern comprising at least one peak expressed as 2θ ($\pm 0.2^\circ$) selected from about 6.4° and 8.9° .

In some embodiments, Form D (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide has an XRPD pattern comprising at least one peak expressed as 2θ ($\pm 0.2^\circ$) selected from about 6.4° and 8.1° .

30 In some embodiments, Form D (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide has an XRPD pattern comprising at least one peak expressed as 2θ ($\pm 0.2^\circ$) selected from about 16.6° and 8.9° .

In some embodiments, Form D (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide has an XRPD pattern comprising at least one peak expressed as 2θ ($\pm 0.2^\circ$) selected from about 16.6° and 8.1° .

5 In some embodiments, Form D (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide has an XRPD pattern comprising at least one peak expressed as 2θ ($\pm 0.2^\circ$) selected from about 8.1° and 8.9° .

10 In some embodiments, Form D (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide has an XRPD pattern comprising at least one peak expressed as 2θ ($\pm 0.2^\circ$) selected from about 21.8° , 6.4° and 16.6° .

15 In some embodiments, Form D (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide has an XRPD pattern comprising at least one peak expressed as 2θ ($\pm 0.2^\circ$) selected from about 21.8° , 6.4° and 8.9° .

In some embodiments, Form D (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide has an XRPD pattern comprising at least one peak expressed as 2θ ($\pm 0.2^\circ$) selected from about 21.8° , 6.4° and 8.1° .

20 In some embodiments, Form D (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide has an XRPD pattern comprising at least one peak expressed as 2θ ($\pm 0.2^\circ$) selected from about 21.8° , 16.6° and 8.9° .

25 In some embodiments, Form D (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide has an XRPD pattern comprising at least one peak expressed as 2θ ($\pm 0.2^\circ$) selected from about 21.8° , 16.6° and 8.1° .

30 In some embodiments, Form D (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide has an XRPD pattern comprising at least one peak expressed as 2θ ($\pm 0.2^\circ$) selected from about 21.8° , 8.9° and 8.1° .

In some embodiments, Form D (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide has an XRPD

pattern comprising at least one peak expressed as 2θ ($\pm 0.2^\circ$) selected from about 16.6° , 8.9° and 8.1° .

In some embodiments, Form D (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide has an XRPD
5 pattern comprising at least one peak expressed as 2θ ($\pm 0.2^\circ$) selected from about 21.8° , 6.4° , 16.6° and 8.9° .

In some embodiments, Form D (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide has an XRPD
10 pattern comprising at least one peak expressed as 2θ ($\pm 0.2^\circ$) selected from about 21.8° , 6.4° , 16.6° and 8.1° .

In some embodiments, Form D (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide has an XRPD
pattern comprising at least one peak expressed as 2θ ($\pm 0.2^\circ$) selected from about 6.4° , 16.6° , 8.9° and 8.1° .

15 In some embodiments, Form D (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide has an XRPD
pattern comprising at least one peak expressed as 2θ ($\pm 0.2^\circ$) selected from about 21.8° , 6.4° , 16.6° , 8.9° and 8.1° .

In some embodiments, Form D (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide has an XRPD
20 pattern comprising at least one peak expressed as 2θ ($\pm 0.2^\circ$) selected from the peaks listed in Table 20.

In some embodiments, Form D (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide has an XRPD
25 pattern substantially similar to **Figure 7**.

In some embodiments, Form D (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide has a DSC
thermogram comprising an endotherm with a desolvation onset at about 116°C and a peak at
about 119°C .

30 In some embodiments, Form D (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide has a DSC
thermogram substantially similar to **Figure 8**.

In some embodiments, Form D (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide has a TGA

thermogram exhibiting a mass loss of about 8.0% upon heating from about 25 °C to about 200 °C.

In some embodiments, Form D (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide has a TGA thermogram substantially similar to **Figure 8**.

Form A-Saccharine Salt

In some embodiments, disclosed is Form A (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide saccharine salt.

In some embodiments, Form A (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide saccharine salt has an XRPD pattern comprising at least one peak expressed as 2θ (±0.2°) selected from the peaks listed in Table 21.

In some embodiments, Form A (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide saccharine salt has an XRPD pattern substantially similar to **Figure 9**.

In some embodiments, Form A (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide saccharine salt has a DSC thermogram comprising an endotherm with a melting point onset at about 163 °C and a peak at about 169 °C.

In some embodiments, Form A (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide saccharine salt has a DSC thermogram substantially similar to **Figure 10**.

In some embodiments, Form A (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide saccharine salt has a TGA thermogram exhibiting a mass loss of about 3.1% upon heating from about 25 °C to about 150 °C.

In some embodiments, Form A (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide saccharine salt has a TGA thermogram substantially similar to **Figure 10**.

Form B-Saccharine Salt

In some embodiments, disclosed is Form B (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide saccharine salt.

5 In some embodiments, Form B (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide saccharine salt has an XRPD pattern comprising at least one peak expressed as 2θ ($\pm 0.2^\circ$) selected from the peaks listed in Table 22.

10 In some embodiments, Form B (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide saccharine salt has an XRPD pattern substantially similar to **Figure 11**.

15 In some embodiments, Form B (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide saccharine salt has a DSC thermogram comprising an endotherm with a broad desolvation peak at about 53 °C and two endotherm events with an onset at about 153 °C and a peak at 162 °C and an onset at about 176 °C and a peak at about 182 °C.

In some embodiments, Form B (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide saccharine salt has a DSC thermogram substantially similar to **Figure 12**.

20 In some embodiments, Form B (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide saccharine salt has a TGA thermogram exhibiting a mass loss of about 2.7% upon heating from about 25 °C to about 100 °C.

25 In some embodiments, Form B (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide saccharine salt has a TGA thermogram substantially similar to **Figure 12**.

Form C-Saccharine Salt

30 In some embodiments, disclosed is Form C (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide saccharine salt.

In some embodiments, Form C (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide saccharine salt

has an XRPD pattern comprising at least one peak expressed as 2θ ($\pm 0.2^\circ$) selected from the peaks listed in Table 23.

In some embodiments, Form C (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide saccharine salt
5 has an XRPD pattern substantially similar to **Figure 13**.

Form D-Saccharine Salt

In some embodiments, disclosed is Form D (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide
10 saccharine salt.

In some embodiments, Form D (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide saccharine salt has an XRPD pattern comprising at least one peak expressed as 2θ ($\pm 0.2^\circ$) selected from the peaks listed in Table 24.

15 In some embodiments, Form D (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide saccharine salt has an XRPD pattern substantially similar to **Figure 14**.

Form E-Saccharine Salt

20 In some embodiments, disclosed is Form E (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide saccharine salt.

In some embodiments, Form E (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide saccharine salt
25 has an XRPD pattern comprising at least one peak expressed as 2θ ($\pm 0.2^\circ$) selected from the peaks listed in Table 25.

In some embodiments, Form E (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide saccharine salt
30 has an XRPD pattern substantially similar to **Figure 15**.

Hydrochloride Saccharine Salt

In some embodiments, disclosed is (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide hydrochloride saccharine salt.

In some embodiments, (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide hydrochloride saccharine salt has an XRPD pattern comprising at least one peak expressed as 2θ ($\pm 0.2^\circ$) selected from the peaks listed in Table 26.

5 In some embodiments, (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide hydrochloride saccharine salt has an XRPD pattern substantially similar to **Figure 16**.

Napadisyllic Salt

10 In some embodiments, disclosed is (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide napadisyllic salt.

In some embodiments, (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide hydrochloride napadisyllic salt has an XRPD pattern comprising at least one peak expressed as 2θ ($\pm 0.2^\circ$)
15 selected from the peaks listed in Table 27.

In some embodiments, (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide napadisyllic salt has an XRPD pattern substantially similar to **Figure 17**.

Trimesic Salt

20 In some embodiments, disclosed is (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide trimesic salt.

In some embodiments, (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide hydrochloride
25 trimesic salt has an XRPD pattern comprising at least one peak expressed as 2θ ($\pm 0.2^\circ$) selected from the peaks listed in Table 28.

In some embodiments, (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide trimesic salt has an XRPD pattern substantially similar to **Figure 18**.

30

Pharmaceutical Compositions

In some embodiments, disclosed are pharmaceutical compositions comprising a compound of formula (I), (Ia), (Ib) or of Table 1, and a pharmaceutically acceptable excipient, carrier or diluent.

The language "pharmaceutically acceptable excipient, carrier or diluent" includes compounds, materials, compositions, and/or dosage forms which 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, or other problem or complication, as
5 ascertained by one of skill in the art.

The disclosed compositions may be in a form suitable for oral use (for example, as tablets, lozenges, hard or soft capsules, aqueous or oily suspensions, emulsions, dispersible powders or granules, syrups or elixirs), for topical use (for example, as creams, ointments, gels, or aqueous or oily solutions or suspensions), for administration by inhalation (for example, as a
10 finely divided powder or a liquid aerosol), for administration by insufflation (for example, as a finely divided powder) or for parenteral administration (for example, as a sterile aqueous or oily solution for intravenous, subcutaneous, intramuscular or intramuscular dosing or as a suppository for rectal dosing).

The disclosed compositions may be obtained by conventional procedures using
15 conventional pharmaceutical excipients well known in the art. Thus, compositions intended for oral use may contain, for example, one or more coloring, sweetening, flavoring and/or preservative agents.

Suitable pharmaceutically acceptable excipients for a tablet formulation include, for example, inert diluents such as lactose, sodium carbonate, calcium phosphate or calcium
20 carbonate; granulating and disintegrating agents such as corn starch or algenic acid; binding agents such as starch; lubricating agents such as magnesium stearate, stearic acid or talc; preservative agents such as ethyl or propyl p-hydroxybenzoate; and anti-oxidants, such as ascorbic acid. Tablet formulations may be uncoated or coated either to modify their disintegration and the subsequent absorption of the active ingredient within the gastrointestinal
25 tract, or to improve their stability and/or appearance using conventional coating agents and procedures well known in the art.

Compositions for oral use may be in the form of hard gelatin capsules in which the active ingredient is mixed with an inert solid diluent, for example, calcium carbonate, calcium phosphate or kaolin, or as soft gelatin capsules in which the active ingredient is mixed with
30 water or oil, such as peanut oil, liquid paraffin, or olive oil.

Aqueous suspensions generally contain the active ingredient in finely powdered form or in the form of nano or micronized particles together with one or more suspending agents, such as sodium carboxymethylcellulose, methylcellulose, hydroxypropylmethylcellulose, sodium alginate, polyvinyl-pyrrolidone, gum tragacanth and gum acacia; dispersing or wetting agents

such as lecithin or condensation products of an alkylene oxide with fatty acids (for example polyoxyethylene stearate), or condensation products of ethylene oxide with long chain aliphatic alcohols, for example heptadecaethyleneoxycetanol, or condensation products of ethylene oxide with partial esters derived from fatty acids and a hexitol such as polyoxyethylene sorbitol monooleate, or condensation products of ethylene oxide with long chain aliphatic alcohols, for example heptadecaethyleneoxycetanol, or condensation products of ethylene oxide with partial esters derived from fatty acids and a hexitol such as polyoxyethylene sorbitol monooleate, or condensation products of ethylene oxide with partial esters derived from fatty acids and hexitol anhydrides, for example polyethylene sorbitan monooleate. The aqueous suspensions may also contain one or more preservatives such as ethyl or propyl p-hydroxybenzoate; anti-oxidants such as ascorbic acid; coloring agents; flavoring agents; and/or sweetening agents such as sucrose, saccharine or aspartame.

Oily suspensions may be formulated by suspending the active ingredient in a vegetable oil such as arachis oil, olive oil, sesame oil or coconut oil or in a mineral oil such as liquid paraffin. The oily suspensions may also contain a thickening agent such as beeswax, hard paraffin or cetyl alcohol. Sweetening agents such as those set out above, and flavoring agents may be added to provide a palatable oral preparation. These compositions may be preserved by the addition of an anti-oxidant such as ascorbic acid.

Dispersible powders and granules suitable for preparation of an aqueous suspension by the addition of water generally contain the active ingredient together with a dispersing or wetting agent, suspending agent and one or more preservatives. Suitable dispersing or wetting agents and suspending agents are exemplified by those already mentioned above. Additional excipients such as sweetening, flavoring and coloring agents, may also be present.

The pharmaceutical compositions may also be in the form of oil-in-water emulsions. The oily phase may be a vegetable oil, such as olive oil or arachis oil, or a mineral oil, such as for example liquid paraffin or a mixture of any of these. Suitable emulsifying agents may be, for example, naturally-occurring gums such as gum acacia or gum tragacanth, naturally-occurring phosphatides such as soya bean, lecithin, an esters or partial esters derived from fatty acids and hexitol anhydrides (for example sorbitan monooleate) and condensation products of the said partial esters with ethylene oxide such as polyoxyethylene sorbitan monooleate. The emulsions may also contain sweetening, flavoring and preservative agents.

Syrups and elixirs may be formulated with sweetening agents such as glycerol, propylene glycol, sorbitol, aspartame or sucrose, and may also contain a demulcent, preservative, flavoring and/or coloring agent.

The pharmaceutical compositions may also be in the form of a sterile injectable aqueous or oily suspension, which may be formulated according to known procedures using one or more of the appropriate dispersing or wetting agents and suspending agents, which have been mentioned above. A sterile injectable preparation may also be a sterile injectable solution or suspension in a non-toxic parenterally-acceptable diluent or solvent, for example a solution in 1,3-butanediol.

Compositions for administration by inhalation may be in the form of a conventional pressurized aerosol arranged to dispense the active ingredient either as an aerosol containing finely divided solid or liquid droplets. Conventional aerosol propellants such as volatile fluorinated hydrocarbons or hydrocarbons may be used and the aerosol device is conveniently arranged to dispense a metered quantity of active ingredient.

For further information on formulation the reader is referred to Chapter 25.2 in Volume 5 of Comprehensive Medicinal Chemistry (Corwin Hansch; Chairman of Editorial Board), Pergamon Press 1990.

The amount of active ingredient that is combined with one or more excipients to produce a single dosage form will necessarily vary depending upon the host treated and the particular route of administration. For further information on Routes of Administration and Dosage Regimes the reader is referred to Chapter 25.3 in Volume 5 of Comprehensive Medicinal Chemistry (Corwin Hansch; Chairman of Editorial Board), Pergamon Press 1990.

The compounds of Formula (I), (Ia), (Ib) and Table 1 may be administered once, twice, three times a day or as many times in a 24 hour period as medically necessary. One of skill in the art would readily be able to determine the amount of each individual dose based on the subject. In some embodiments, the compounds of Formula (I), (Ia), (Ib) or Table 1 are administered in one dosage form. In some embodiments, the compounds of formula (I), (Ia), (Ib) or Table are administered in multiple dosage forms.

Methods

In one aspect, disclosed are methods for treating a JAK-related disorder in a subject in need thereof, comprising administering to the subject an effective amount of a compound of Formula (I), (Ia), (Ib) or Table 1, or a pharmaceutically acceptable salt or solid form thereof.

In one aspect, disclosed is a compound of Formula (I), (Ia), (Ib) or Table 1, or a pharmaceutically acceptable salt or solid form thereof, for use in treating a JAK-related disorder.

In one aspect, disclosed is the use of a compound of Formula (I), (Ia), (Ib) or Table 1, or a pharmaceutically acceptable salt or solid form thereof, in the manufacture of a medicament for

treating a JAK-related disorder.

In one aspect, disclosed are pharmaceutical compositions comprising a compound of Formula (I), (Ia), (Ib) or Table 1, or a pharmaceutically acceptable salt or solid form thereof, for use in treating a JAK-related disorder.

5 The language "JAK-related disorder" includes cancer, cancer cachexia and immune disorders.

The term "cancer" includes cancers with: (i) an EGFR-related etiology such as non-small cell lung cancer (NSCLC), head and neck: squamous cell cancer (HNSCC) and colorectal cancer; (ii) and activating RAS family mutations such as NSCLC, pancreatic cancer, colorectal cancer, prostate cancer, melanoma, thyroid cancer, bladder cancer, cholangiocarcinoma, and leukemia; (iii) a HER2 amplification or mutation such as breast cancer, gastric cancer, lung cancer; (iv) an ALK gene activation such as lung cancer, breast cancer, colorectal cancer, diffuse large B-cell lymphoma, anaplastic large cell lymphoma; (v) a MET amplification or mutation such as NSCLC, gastric cancer, colorectal cancer, papillary renal cell carcinoma; and
10 (vi) an FGFR-related etiology such as breast cancer, gastric cancer, endometrial cancer, lung cancer. In some embodiments, the cancer is pancreatic cancer, gastrointestinal cancer, breast cancer, a gynecological cancer (e.g., ovarian cancer or cervical cancer), bladder cancer, SCHN, non-small cell lung cancer or small cell lung cancer. In some embodiments, the cancer has metastasized.

20 In one aspect, disclosed are methods for treating cancer in a subject in need thereof comprising administering to the subject an effective amount of a compound of Formula (I), (Ia), (Ib), or Table 1, or a pharmaceutically acceptable salt or solid form thereof in combination with an anti-cancer therapeutic agent, or a pharmaceutically acceptable salt thereof.

In one aspect, disclosed is a compound of Formula (I), (Ia), (Ib) or Table 1, or a
25 pharmaceutically acceptable salt or solid form thereof in combination with anti-cancer therapeutic agent, or a pharmaceutically acceptable salt thereof, for use in treating a cancer.

In one aspect, disclosed is the use of a compound of Formula (I), (Ia), (Ib) or Table 1, or a pharmaceutically acceptable salt or solid form thereof, in combination with an anti-cancer therapeutic agent, or a pharmaceutically acceptable salt thereof, in the manufacture of a
30 medicament for treating cancer.

In one aspect, disclosed are pharmaceutical compositions comprising a compound of Formula (I), (Ia), (Ib) or Table 1, or a pharmaceutically acceptable salt or solid form thereof, in combination with an anti-cancer therapeutic agent, or a pharmaceutically acceptable salt thereof, for use in treating cancer.

The language "in combination with" includes administering the compound of Formula (I), (Ia), (Ib) or Table 1, or a pharmaceutically acceptable salt thereof, and the anti-cancer therapeutic agent, or pharmaceutically acceptable salt thereof, sequentially, separately or simultaneously. In some aspects, the compound of Formula (I), (Ia), (Ib) or Table 1, or a pharmaceutically acceptable salt thereof, and the anti-cancer therapeutic agent, or pharmaceutically acceptable salt thereof, are administered in the same formulation, for example, in a fixed dose formulation. In some embodiments, the compound of Formula (I), (Ia), (Ib) or Table 1, or a pharmaceutically acceptable salt thereof, and the anti-cancer therapeutic agent, or pharmaceutically acceptable salt thereof, are administered in separate formulations, and are administered at substantially the same time, sequentially or separately. In some embodiments, the compound of Formula (I), (Ia), (Ib) or Table 1, or pharmaceutically acceptable salt thereof, is administered for one day, two days, three days, four days, five days, six days, seven days, eight days, nine days, ten days, 11 days, 12 days, 13 days, 14 days, three weeks or one month in a row. In some embodiments, compound of Formula (I), (Ia), (Ib) or Table 1, or pharmaceutically acceptable salt thereof, is administered intermittently, for example, for 7 days followed by a 7 day clearance period (e.g., 7 days on/7 days off), for 1 day followed by a 6 day clearance period (e.g., 1 day on/6 days off), for 2 days followed by a 5 day clearance period (2 days on/5 days off), for 3 days followed by a 4 day clearance period (e.g., 3 days on/4 days off), for 4 days followed by a 3 day clearance period (e.g., 4 days on/3 days off), for 5 days followed by a 2 day clearance period (5 days on/2 days off), or for 6 days followed by a 1 day clearance period (6 days on/1 day off).

The language "anti-cancer therapeutic agent" includes, for example, EGFR inhibitors, MAPK pathway inhibitors, Raf inhibitors, HER2 inhibitors, FGFR inhibitors, antimetabolites, alkylating agents and antimitotic agents, and pharmaceutically acceptable salts thereof.

In some embodiments, a compound of Formula (I), (Ia), (Ib) or Table 1 is administered in combination with one or more EGFR inhibitors. Examples of EGFR inhibitors include EGFR antibodies, ABX-EGF, anti-EGFR immunoliposomes, EGF-vaccine, EMD-7200, ERBITUX® (cetuximab), HR3, IgA antibodies, IRESSA® (gefitinib), TARCEVA® (erlotinib or OSI-774), TP-38, EGFR fusion protein, TYKERB® (lapatinib), TAGRISSO™ (osimertinib or AZD9291), GILOTRIF® (afatinib), CO-1686, WZ4002, PD153035, PF 00299804 and the like. In some embodiments, a compound of Formula (I), (Ia), (Ib) or Table 1 is administered in combination with osimertinib. In some embodiments, a compound of Formula (I), (Ia), (Ib) or Table 1 is administered in combination with gefitinib.

In some embodiments, a compound of Formula (I), (Ia), (Ib) or Table 1 is administered in

combination with one or more MAPK pathway inhibitors. MAPK pathway inhibitors include MEK inhibitors such as Selumetinib, Mekinist® (trametinib), Cobimetinib, PD0325901, Pimasertib, MEK162, Refametinib and the like; Raf and B-Raf inhibitors which include vemurafenib, dabrafenib, Encorafenib (LGX818) and the like.

5 In some embodiments, a compound of Formula (I), (Ia), (Ib) or Table 1 is administered in combination with one or more HER2 inhibitors. HER2 inhibitors include CP-724-714, CI-1033 (canertinib), HERCEPTIN® (trastuzumab), TYKERB® (lapatinib), OMNITARG® (2C4, petuzumab), TAK-165, GW-572016 (ionafarnib), GW-282974, EKB-569, PI-166, dHER2 (HER2 vaccine), APC-8024 (HER-2 vaccine), anti-HER/2neu bispecific antibody, B7.her2lgG3, AS
10 HER2 bifunctional bispecific antibodies, mAB AR-209, mAB 2B-1 and the like.

In some embodiments, a compound of Formula (I), (Ia), (Ib) or Table 1 is administered in combination with one or more ALK inhibitors. ALK inhibitors include crizotinib, ceritinib, and the like.

15 In some embodiments, a compound of Formula (I), (Ia), (Ib) or Table 1 is administered in combination with one or more FGFR inhibitors. FGFR inhibitors include AZD4547, BJJ398, Dovitinib, Lucitanib, MGFR1877S, FP-1039 and the like.

In some embodiments, a compound of Formula (I), (Ia), (Ib) or Table 1 is administered in combination with one or more MET inhibitors. MET inhibitors include Savolitinib, Onartuzumab, Rilotumumab, Cabozantinib, Tivantinib, LY2875358, Ficlatuzumab, Foretinib, Crizotinib,
20 INC280, AMG337, MSC2156119J and the like

In some embodiments, a compound of Formula (I), (Ia), (Ib) or Table 1 is administered in combination with one or more antimetabolites. Antimetabolites include ALIMTA® (pemetrexed disodium, LY231514, MTA), 5-azacitidine, XELODA® (capecitabine), carmofur, LEUSTAT® (cladribine), clofarabine, cytarabine, cytarabine ocfosfate, cytosine arabinoside, decitabine,
25 deferoxamine, doxifluridine, eflornithine, EICAR (5-ethynyl-1-β-D-ribofuranosylimidazole-4-carboxamide), enocitabine, ethynylcytidine, fludarabine, 5-fluorouracil alone or in combination with leucovorin, GEMZAR® (gemcitabine), hydroxyurea, ALKERAN® (melphalan), mercaptopurine, 6-mercaptopurine riboside, methotrexate, mycophenolic acid, nelarabine, nolatrexed, ocfosfate, pelitrexol, pentostatin, pemextred, raltitrexed, Ribavirin, triapine,
30 trimetrexate, S-1, tiazofurin, tegafur, TS-1, vidarabine, UFT and the like.

In some embodiments, a compound of Formula (I), (Ia), (Ib) or Table 1 is administered in combination with one or more alkylating agents. Alkylating agents include altretamine, AMD-473, AP-5280, apaziquone, bendamustine, brostallicin, busulfan, cisplatin, carboplatin, carboquone, carmustine (BCNU), chlorambucil, CLORETAZINE® (laromustine, VNP 40101M),

cyclophosphamide, decarbazine, estramustine, fotemustine, glufosfamide, ifosfamide, KW-2170, lomustine (CCNU), mafosfamide, melphalan, mitobronitol, mitolactol, nimustine, nitrogen mustard N-oxide, nitrosoureas, oxaliplatin, ranimustine, temozolomide, thiotepa, TREANDA® (bendamustine), treosulfan, rofosfamide and the like.

5 In some embodiments, a compound of Formula (I), (Ia), (Ib) or Table 1 is administered in combination with one or more antimitotic agents. Antimitotic agents include batabulin, epothilone D (KOS-862), N-(2-((4-hydroxyphenyl)amino)pyridin-3-yl)-4-methoxybenzenesulfonamide, ixabepilone (BMS 247550), paclitaxel, TAXOTERE® (docetaxel), PNU100940 (109881), patupilone, XRP-9881 (larotaxel), vinflunine, ZK-EPO (synthetic
10 epothilone) and the like.

The language “cancer cachexia” includes a syndrome with symptoms that includes host tissue wasting, anorexia, asthenia and abnormal host intermediary metabolism. In some embodiments, the subject suffering from cancer cachexia has pancreatic cancer or an upper gastrointestinal cancer, for example, esophageal cancer, stomach cancer, gastric cancer, liver
15 cancer, gall bladder cancer, neuroendocrine cancer or Barrett’s esophagus. In some embodiments, the subject suffering from cancer cachexia has terminal cancer.

In one aspect, disclosed are methods for treating cancer cachexia in a subject in need thereof comprising administering to the subject an effective amount of a compound of Formula (I), (Ia), (Ib), or Table 1, or a pharmaceutically acceptable salt or solid form thereof.

20 In one aspect, disclosed is a compound of Formula (I), (Ia), (Ib) or Table 1, or a pharmaceutically acceptable salt or solid form thereof, for use in treating cancer cachexia.

In one aspect, disclosed is the use of a compound of Formula (I), (Ia), (Ib) or Table 1, or a pharmaceutically acceptable salt or solid form thereof, in the manufacture of a medicament for treating cancer cachexia.

25 In one aspect, disclosed are pharmaceutical compositions comprising a compound of Formula (I), (Ia), (Ib) or Table 1, or a pharmaceutically acceptable salt or solid form thereof, for use in treating cancer cachexia.

The language “immune disorder” includes, for example, bone marrow disorders (*e.g.*, myelofibrosis and polycythemia vera), rheumatoid arthritis, psoriasis, irritable bowel disease
30 (IBD), Crohn’s disease, lupus, multiple sclerosis, asthma, autoimmune thyroid disorders (*e.g.*, Hashimoto’s thyroiditis, Graves’ disease or post-partum thyroiditis), ulcerative colitis, Alopecia areata, vitiligo and myositis.

In one aspect, disclosed are methods for treating an immune disorder in a subject in need thereof comprising administering to the subject an effective amount of a compound of Formula

(I), (Ia), (Ib), or Table 1, or a pharmaceutically acceptable salt or solid form thereof.

In one aspect, disclosed is a compound of Formula (I), (Ia), (Ib) or Table 1, or a pharmaceutically acceptable salt or solid form thereof, for use in treating an immune disorder.

5 In one aspect, disclosed is the use of a compound of Formula (I), (Ia), (Ib) or Table 1, or a pharmaceutically acceptable salt or solid form thereof, in the manufacture of a medicament for treating an immune disorder.

In one aspect, disclosed are pharmaceutical compositions comprising a compound of Formula (I), (Ia), (Ib) or Table 1, or a pharmaceutically acceptable salt or solid form thereof, for use in treating an immune disorder.

10 In one aspect, disclosed are methods for inhibiting JAK in a subject in need thereof, comprising administering to the subject an effective amount of a compound of Formula (I), (Ia), (Ib) or Table 1, or a pharmaceutically acceptable salt or solid form thereof.

In one aspect, disclosed is a compound of Formula (I), (Ia), (Ib) or Table 1, or a pharmaceutically acceptable salt or solid form thereof, for use in inhibiting JAK.

15 In one aspect, disclosed is the use of a compound of Formula (I), (Ia), (Ib) or Table 1, or a pharmaceutically acceptable salt or solid form thereof, in the manufacture of a medicament for inhibiting JAK.

20 In one aspect, disclosed are pharmaceutical compositions comprising a compound of Formula (I), (Ia), (Ib) or Table 1, or a pharmaceutically acceptable salt or solid form thereof, for use in inhibiting JAK.

The term "JAK" includes a family of Janus kinases that are intracellular, nonreceptor tyrosine kinases that transduce cytokine-mediated signals via the JAK-STAT pathway. The term JAK includes JAK1, JAK2 and JAK3. In some embodiments, the compounds of Formula (I), (Ia) and (Ib) are selective inhibitors of JAK1, JAK2 and/or JAK3. The language "selective inhibitor" includes compounds that have a greater inhibitory effect (as demonstrated, for example, by a lower IC_{50}) for one or two of the JAK family members over the other JAK family members. For example, a JAK1-selective inhibitor exhibits greater inhibitory effect on JAK1 over JAK2 and JAK3; a JAK2-selective inhibitor exhibits a greater inhibitory effect on JAK2 over JAK1 and JAK3; a JAK3-selective inhibitor exhibits a greater inhibitory effect on JAK3 over JAK1 and JAK2; a JAK1/2-selective inhibitor exhibits a greater inhibitory effect on JAK1 and JAK2 over JAK3; a JAK1/3-selective inhibitor exhibits a greater inhibitory effect on JAK1 and JAK3 over JAK2; and a JAK2/3-selective inhibitor exhibits a greater inhibitory effect on JAK2 and JAK3 over JAK1. In some embodiments, the compounds of Formula (I), (Ia) and (Ib) are JAK1-selective inhibitors. In some embodiments, the compounds of Formula (I), (Ia) and (Ib)

25
30

are JAK1/2-selective inhibitors.

The language “effective amount” includes an amount of a compound of Formula (I), (Ia), (Ib) or Table 1 that will elicit a biological or medical response in a subject, for example, the reduction or inhibition of enzyme or protein activity related to JAK, cancer or an immune disorder; amelioration of symptoms of cancer or an immune disorder; or the slowing or delaying of progression of cancer or an immune disorder. In some embodiments, the language “effective amount” includes the amount of a compound of Formula (I), (Ia), (Ib) or Table 1, that when administered to a subject, is effective to at least partially alleviate, inhibit, and/or ameliorate cancer or an immune disorder or inhibit JAK, and/or reduce or inhibit the growth of a tumor or proliferation of cancerous cells in a subject.

The term “subject” includes warm blooded mammals, for example, primates, dogs, cats, rabbits, rats, and mice. In some embodiments, the subject is a primate, for example, a human. In some embodiments, the subject is suffering from cancer or an immune disorder. In some embodiments, the subject is in need of treatment (*e.g.*, the subject would benefit biologically or medically from treatment). In some embodiments, the subject is suffering from cancer cachexia. In some embodiments, the subject is suffering from cancer. In some embodiments, the subject is suffering from cancer cachexia. In some embodiments, the subject is suffering from an immune disorder. In some embodiments, the subject may have elevated blood levels of inflammatory biomarkers *e.g.*, serum systemic C-reactive protein (CRP), IL-6, TNF α , IL-1, procalcitonin and IL-8. In some embodiments, the subject may be suffering from a high STAT3-positive tumor. In some embodiments, the subject is suffering from a EGFR-M positive cancer (*e.g.*, non-small cell lung cancer). In some embodiments, the EGFR-M positive cancer has a predominately T790M-positive mutation. In some embodiments, the EGFR-M positive cancer has a predominately T790M-negative mutation. In some embodiments, the subject is suffering from a KRAS mutant cancer (*e.g.*, KRAS mutated non-small cell lung cancer). In some embodiments, the subject is suffering from metastatic pancreatic cancer, metastatic gastrointestinal cancer, metastatic breast cancer, a metastatic gynecologic cancer (*e.g.*, metastatic ovarian cancer or metastatic cervical cancer), metastatic bladder cancer, metastatic squamous cell head and neck cancer (SCHN), metastatic non-small cell lung cancer, metastatic haematological cancers (*e.g.*, non-Hodgkin’s lymphoma) or metastatic small cell lung cancer. In some embodiments, the subject suffering from cancer may show evidence of immune inflammation, including, for example, the presence of PDL1, interferon gamma, tumor-infiltrating leukocytes and gene expression signatures indicating increased type I or type II interferon signaling, abnormal levels tumor suppressive cells, such as regulatory T lymphocytes or

myeloid-derived cells, abnormal levels of granulocytes or proteins indicating the presence of granulocytes.

The language "inhibit," "inhibition" or "inhibiting" includes a decrease in the baseline activity of a biological activity or process. In some embodiments, the compounds of Formula (I), (Ia), (Ib) or Table 1 inhibit JAK.

The language "treat," "treating" and "treatment" includes the reduction or inhibition of enzyme or protein activity related to JAK, cancer or an immune disorder in a subject, amelioration of one or more symptoms of a cancer or an immune disorder in a subject, or the slowing or delaying of progression of cancer or an immune disorder in a subject. The language "treat," "treating" and "treatment" also includes the reduction or inhibition of the growth of a tumor or proliferation of cancerous cells in a subject.

Examples

Aspects of the present disclosure can be further defined by reference to the following non-limiting examples, which describe in detail preparation of certain compounds and intermediates of the present disclosure and methods for using compounds of the present disclosure. It will be apparent to those skilled in the art that many modifications, both to materials and methods, can be practiced without departing from the scope of the present disclosure.

Unless stated otherwise:

(i) all syntheses were carried out at ambient temperature, *i.e.* in the range 17 to 25°C and under an atmosphere of an inert gas such as nitrogen unless otherwise stated;

(ii) evaporations were carried out by rotary evaporation or utilising Genevac equipment or Biotage v10 evaporator *in vacuo* and work-up procedures were carried out after removal of residual solids by filtration;

(iii) flash chromatography purifications were performed on an automated Teledyne Isco CombiFlash® Rf or Teledyne Isco CombiFlash® Companion® using prepacked RediSep Rf Gold™ Silica Columns (20-40 µm, spherical particles), GraceResolv™ Cartridges (Davisil® silica) or Silicycle cartridges (40 - 63 µm).

(iv) preparative chromatography was performed on a Gilson prep HPLC instrument with UV collection; alternatively, preparative chromatography was performed on a Waters AutoPurification HPLC-MS instrument with MS- and UV- triggered collection;

(v) chiral preparative chromatography was performed on a Gilson instrument with UV collection (233 injector / fraction collector, 333 & 334 pumps, 155 UV detector) or a Varian Prep Star instrument (2 x SD1 pumps, 325 UV detector, 701 fraction collector) pump running with

Gilson 305 injection; alternatively, chiral preparative chromatography was performed on a Waters Prep 100 SFC-MS instrument with MS- and UV- triggered collection or a Thar MultiGram III SFC instrument with UV collection.

(vi) yields, where present, are not necessarily the maximum attainable;

(vii) in general, the structures of end-products of the Formula I were confirmed by nuclear magnetic resonance (NMR) spectroscopy; NMR chemical shift values were measured on the delta scale [proton magnetic resonance spectra were determined using a Bruker Avance 500 (500 MHz), Bruker Avance 400 (400 MHz), Bruker Avance 300 (300 MHz) or Bruker DRX (300 MHz) instrument]; measurements were taken at ambient temperature unless otherwise specified; the following abbreviations have been used: s, singlet; d, doublet; t, triplet; q, quartet; m, multiplet; dd, doublet of doublets; ddd, doublet of doublet of doublet; dt, doublet of triplets; bs, broad signal.

(viii) in general, end-products of the Formula I were also characterized by mass spectroscopy following liquid chromatography (LCMS or UPLC); UPLC was carried out using a Waters UPLC fitted with a Waters SQ mass spectrometer (Column temp 40 °C, UV = 220-300 nm or 190-400 nm, Mass Spec = ESI with positive/negative switching) at a flow rate of 1 mL/min using a solvent system of 97% A + 3% B to 3% A + 97% B over 1.50 min (total run time with equilibration back to starting conditions, etc., 1.70 min), where A = 0.1% formic acid or 0.05% trifluoroacetic acid in water (for acidic work) or 0.1% ammonium hydroxide in water (for basic work) and B = acetonitrile. For acidic analysis the column used was a Waters Acquity HSS T3 (1.8 µm, 2.1x 50 mm), for basic analysis the column used was a Waters Acquity BEH C18 (1.7 µm 2.1x50 mm). Alternatively, UPLC was carried out using a Waters UPLC fitted with a Waters SQ mass spectrometer (Column temp 30 °C, UV = 210-400 nm, Mass Spec = ESI with positive/negative switching) at a flow rate of 1mL/min using a solvent gradient of 2 to 98% B over 1.5 mins (total run time with equilibration back to starting conditions 2 min), where A = 0.1% formic acid in water and B = 0.1% formic acid in acetonitrile (for acidic work) or A = 0.1% ammonium hydroxide in water and B = acetonitrile (for basic work). For acidic analysis the column used was a Waters Acquity HSS T3 (1.8 µm, 2.1x30 mm), for basic analysis the column used was a Waters Acquity BEH C18 (1.7 µm, 2.1x30 mm); LCMS was carried out using a Waters Alliance HT (2795) fitted with a Waters ZQ ESCi mass spectrometer and a Phenomenex Gemini–NX C18 (5 µm, 110A, 2.1x50 mm column at a flow rate of 1.1 mL/min 95% A to 95% B over 4 min with a 0.5 min hold where A = 0.1% formic acid and B = 0.1% formic acid in acetonitrile (for acidic work) or A = 0.1% ammonium hydroxide in water and B = acetonitrile (for basic work). Additionally, LCMS was carried out using a Shimadzu UFLC fitted with a Shimadzu

LCMS-2020 mass spectrometer and a Waters HSS C18 (1.8 μm , 2.1x50 mm) or Shim-pack XR-ODS (2.2 μm , 3.0x50 mm) or Phenomenex Gemini-NX C18 (3 μm , 3.0x50 mm) column at a flow rate of 0.7mL/min (for Waters HSS C18 column), 1.0mL/min (for Shim-pack XR-ODS column) or 1.2mL/min (for Phenomenex Gemini-NX C18), 95% A to 95% B over 2.2 min with a 0.6 min hold, where A = 0.1% formic acid or 0.05% trifluoroacetic acid in water (for acidic work) or 0.1% ammonium hydroxide or 6.5 mM ammonium carbonate in water (for basic work) and B = acetonitrile. The reported molecular ion corresponds to the $[\text{M}+\text{H}]^+$ unless otherwise specified; for molecules with multiple isotopic patterns (Br, Cl, etc.) the reported value is the one obtained for the lowest isotope mass unless otherwise specified.

(ix) ion exchange purification was generally performed using an SCX-2 (Biotage) cartridge.

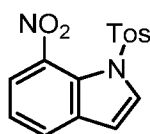
(x) intermediate purity was assessed by thin layer chromatographic, mass spectroscopy, LCMS, UPLC/MS, HPLC (high performance liquid chromatography) and/or NMR analysis;

(xi) the following abbreviations have been used:-

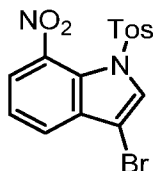
ACN	acetonitrile
BID	twice a day
BSA	bovine serum albumin
DCM	dichloromethane
DMF	<i>N,N</i> -dimethylformamide
DMSO	dimethyl sulphoxide
dppf	1,1'-bis(diphenylphosphino)ferrocene
EA	ethyl acetate
ee	enantiomeric excess
equiv	equivalents
e.r.	enantiomeric ratio
EtOH	ethanol
HATU	(1-[Bis(dimethylamino)methylene]-1H-1,2,3-triazolo[4,5-b]pyridinium 3-oxid hexafluorophosphate)
HCl	hydrochloric acid
HPMC	hydroxypropyl methylcellulose
IPA	Isopropanol
NaOH	sodium hydroxide

NSCLC	non-small cell lung cancer
QD	four times a day
TBME	tert-butyl methyl ether
TEA	triethylamine
5 TFA	trifluoroacetic acid
THF	tetrahydrofuran
Tos	p-toluenesulfonyl
Xantphos	4,5-bis(diphenylphosphino)-9,9-dimethylxanthene

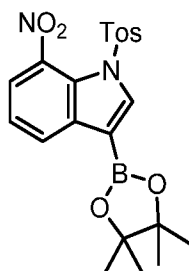
10 **Intermediate 1: 1-[(4-Methylphenyl)sulfonyl]-7-nitro-1H-indole**



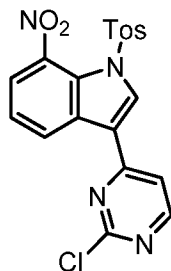
A solution of NaOH (599 g, 14986.55 mmol) in water (1500 mL) was added to a stirred mixture of 7-nitro-1H-indole (243 g, 1498.65 mmol) and tetrabutylammonium hydrogen sulfate (50.9 g, 149.87 mmol) in DCM (3000 mL) at 25°C, over a period of 5 minutes under air. The resulting mixture was stirred at 25 °C for 20 minutes. 4-methylphenylsulfonyl chloride (371 g, 1948.25 mmol) was added under air and the resulting mixture was stirred at 25 °C for 16 hours. The reaction mixture was diluted with DCM (2 L), and washed sequentially with water (500 mL x 2), 10% aqueous K₂CO₃ (500 mL x 2), and 1 M HCl (500 mL x 2) and saturated NaCl (500 mL x 2). The organic layer was dried over Na₂SO₄, filtered and evaporated. When approximately 200 mL DCM was left, 500 mL EA was added. The solvent was removed under reduced pressure. When approximately 200 mL EA was left, 1000 mL TBME was added. The precipitate was collected by filtration, washed with TBME (1 L) and dried under vacuum to afford 1-[(4-methylphenyl)sulfonyl]-7-nitro-1H-indole (402 g, 85 %, **Intermediate 1**) as a white solid, which was used without further purification; ¹H NMR δ (DMSO-d₆, 300 MHz) 2.39 (3H, s), 7.09 (1H, d), 7.40 - 7.55 (3H, m), 7.75 - 7.85 (3H, m), 7.95 - 8.00 (1H, m), 8.06 (1H, d); m/z (ES⁺), [M+H]⁺ = 317.

Intermediate 2: 3-Bromo-1-[(4-methylphenyl)sulfonyl]-7-nitro-1*H*-indole

Bromine (81 mL, 1580 mmol) was added dropwise to 1-[(4-methylphenyl)sulfonyl]-7-nitro-1*H*-indole (50 g, 158 mmol, **Intermediate 1**) in CCl₄ (1000 mL) at 80°C. The resulting solution was stirred at 80°C for 6 hours. The mixture was cooled to room temperature, concentrated and the residue was washed with ethyl acetate to afford 3-bromo-1-[(4-methylphenyl)sulfonyl]-7-nitro-1*H*-indole (53 g, 85%, **Intermediate 2**) as a brown solid; ¹H NMR δ (DMSO-d₆, 300 MHz) 2.41 (3H, s), 7.55 - 7.62 (2H, m), 7.57 (1H, t), 7.85 - 7.92 (3H, m), 7.96 (1H, d), 8.49 (1H, s); m/z (ES⁻), [M-H]⁻ = 393.

Intermediate 3: 1-[(4-Methylphenyl)sulfonyl]-7-nitro-3-(4,4,5,5-tetramethyl-1,3,2-dioxaborolan-2-yl)-1*H*-indole

A solution of 3-bromo-1-[(4-methylphenyl)sulfonyl]-7-nitro-1*H*-indole (200 g, 506 mmol, **Intermediate 2**), 4,4,4',4',5,5,5',5'-octamethyl-2,2'-bi(1,3,2-dioxaborolane) (193 g, 759 mmol), potassium acetate (99 g, 1012 mmol) and PdCl₂(dppf) (18.5 g, 25.3 mmol) in 1,4-dioxane (1500 mL) was degassed with nitrogen three times, then the reaction mixture was stirred at 90°C for 8 hours. The mixture was cooled to room temperature and concentrated. The solids were treated with water and filtered. Washing with methanol and drying in vacuo afforded the 1-[(4-methylphenyl)sulfonyl]-7-nitro-3-(4,4,5,5-tetramethyl-1,3,2-dioxaborolan-2-yl)-1*H*-indole (150 g, 67%, **Intermediate 3**) as a grey solid; ¹H NMR δ (Chloroform-d, 400 MHz) 1.41 (12H, s), 2.47 (3H, s), 7.38 - 7.43 (3H, m), 7.66 (1H, d), 7.87 (2H, d), 8.24 (1H, s), 8.29 - 8.32 (1H, d); m/z (ES⁺), [M+H]⁺ = 443.

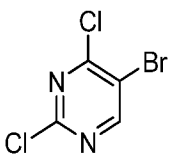
Intermediate 4: 3-(2-Chloro-4-pyrimidinyl)-1-[(4-methylphenyl)sulfonyl]-7-nitro-1H-indole

1-[(4-Methylphenyl)sulfonyl]-7-nitro-3-(4,4,5,5-tetramethyl-1,3,2-dioxaborolan-2-yl)-1H-indole (15 g, 33.9 mmol, **Intermediate 3**), 2,4-dichloropyrimidine (6.6 g, 44.1 mmol), potassium carbonate (14.1 g, 101.7 mmol) and PdCl₂(dppf) (2.5 g, 3.4 mmol) in dioxane (200 mL) and water (40 mL) were stirred under nitrogen at 80°C for 12 hours. The solvent was removed under reduced pressure. The aqueous layer was extracted with THF (4 x 100 mL) and concentrated to give 3-(2-chloro-4-pyrimidinyl)-1-[(4-methylphenyl)sulfonyl]-7-nitro-1H-indole (12 g, 83%, **Intermediate 4**) as a brown solid, which was used without further purification; ¹H NMR δ (DMSO-d₆, 300 MHz) 2.42 (3H, s), 7.52 (2H, d), 7.68 (1H, t), 7.98 (3H, m), 8.31 (1H, d), 8.85 - 8.90 (2H, m), 9.30 (1H, s); m/z (ES⁺), [M+H]⁺ = 429.

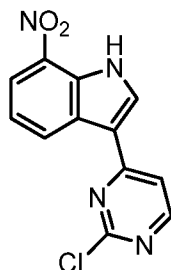
The procedure described above was repeated using the indicated dichloropyrimidine to give Intermediates 5-8 described in Table 2:

Table 2

Intermediate	Dichloropyrimidine	NMR δ (300 MHz)	m/z [M+H] ⁺	Yield %
5 3-(2-chloro-5-fluoro-4-pyrimidinyl)-1-[(4-methylphenyl)sulfonyl]-7-nitro-1H-indole		DMSO-d ₆ 2.43 (3H, s), 7.52 (2H, d), 7.70 (1H, t), 8.00 (3H, m), 8.76 (1H, s), 8.82 (1H, d), 9.04 (1H, d)	447	76
6 3-(2-chloro-5-methyl-4-pyrimidinyl)-1-[(4-methylphenyl)sulfonyl]-7-nitro-1H-indole		DMSO-d ₆ 2.47 (3H, s), 2.50 (3H, s), 7.56 (2H, d), 7.68 (1H, dd), 7.97 (2H, d), 8.04 (1H, d), 8.47 (1H, d), 8.68 (1H, s), 8.85 (1H, s)	443	83
7 3-(2,5-dichloro-4-pyrimidinyl)-1-[(4-methylphenyl)sulfonyl]-7-nitro-1H-indole		DMSO-d ₆ 2.40 (3H, s), 7.52 (2H, d), 7.65 (1H, d), 7.91 (2H, d), 8.00 (1H, d), 8.50 (1H, d), 8.88 (1H, s), 9.06 (1H, s)	463	82

Intermediate	Dichloropyrimidine	NMR δ (300 MHz)	m/z [M+H] ⁺	Yield %
8 3-(5-bromo-2-chloro-4-pyrimidinyl)-1-[(4-methylphenyl)sulfonyl]-7-nitro-1 <i>H</i> -indole		DMSO-d ₆ 2.38 (3H, s), 7.46 (2H, d), 7.62 (1H, t), 7.88 (2H, d), 7.95 (1H, d), 8.35 (1H, d), 8.88 (1H, s), 9.16 (1H, s)	509	51

Intermediate 9: 3-(2-Chloro-4-pyrimidinyl)-7-nitro-1*H*-indole



3-(2-Chloro-4-pyrimidinyl)-1-[(4-methylphenyl)sulfonyl]-7-nitro-1*H*-indole (1 g, 2.3 mmol, **Intermediate 4**) and sodium hydroxide (1.86 g, 46.6 mmol) in THF (10 mL) and water (5 mL) was stirred at 50°C for 2 hours. The solvent was removed under reduced pressure. The crude product was purified by flash silica chromatography, elution gradient 0 to 10% methanol in ethyl acetate. Pure fractions were evaporated to dryness to afford 3-(2-chloro-4-pyrimidinyl)-7-nitro-1*H*-indole (0.52 g, 81%, **Intermediate 9**) as a yellow solid; ¹H NMR δ (DMSO-d₆, 300 MHz) 7.45 (1H, t), 8.10 (1H, s), 8.19 (1H, d), 8.60 (1H, d), 8.66 (1H, s), 8.94 (1H, d), 12.70 (1H, s); m/z (ES⁺), [M+H]⁺ = 275.

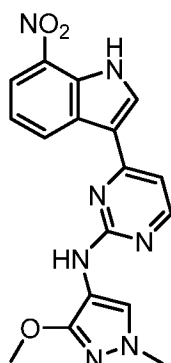
The procedure described above was repeated using the indicated Starting Intermediate to give Intermediates 10-13 described in Table 3:

Table 3

Intermediate	Starting Intermediate	NMR δ (300 MHz)	m/z [M+H] ⁺	Yield %
10 3-(2-chloro-5-fluoro-4-pyrimidinyl)-7-nitro-1 <i>H</i> -indole	5	DMSO-d ₆ 7.42 (1H, t), 8.14 (1H, d), 8.33 (1H, d), 8.69 (1H, d), 8.92 (1H, d) 12.72 (1H, s)	293	89
11 3-(2-chloro-5-methyl-4-pyrimidinyl)-7-nitro-1 <i>H</i> -indole	6	DMSO-d ₆ 2.52 (3H, s), 7.46 (1H, t), 8.20 (1H, s), 8.24 (1H, d), 8.62 (1H, s), 8.89 (1H, d), 12.65 (1H, s)	289	77
12 3-(2,5-dichloro-4-pyrimidinyl)-7-nitro-1 <i>H</i> -indole	7	DMSO-d ₆ 7.50 (1H, t), 8.24 (1H, d), 8.65 (1H, m), 8.89 (1H, s), 8.92 (1H, d), 12.78 (1H, s)	309	70

13 3-(5-bromo-2-chloro-4-pyrimidinyl)-7-nitro-1 <i>H</i> -indole	8	DMSO-d ₆ 7.48 (1H, t), 8.25 (1H, d), 8.75 (1H, s), 8.83 (1H, d), 8.97 (1H, d), 12.77 (1H, s)	355	80
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Intermediate 14: N-(3-Methoxy-1-methyl-1*H*-pyrazol-4-yl)-4-(7-nitro-1*H*-indol-3-yl)pyrimidin-2-amine

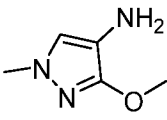
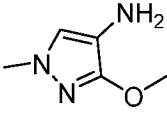
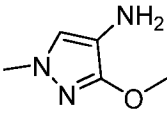
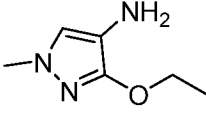
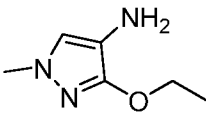


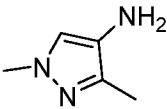
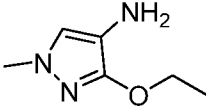
3-(2-Chloro-4-pyrimidinyl)-7-nitro-1*H*-indole (300 mg, 1.1 mmol, **Intermediate 9**), 3-methoxy-1-methyl-1*H*-pyrazol-4-amine dihydrochloride (328 mg, 1.64 mmol) and 4-methylbenzenesulfonic acid monohydrate (623 mg, 3.28 mmol) were dissolved in isopropanol (16 mL) and sealed into a microwave tube. The reaction was heated at 130°C for 2 hours in the microwave reactor and cooled to room temperature. The reaction was concentrated under reduced pressure and then filtered to give N-(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)-4-(7-nitro-1*H*-indol-3-yl)pyrimidin-2-amine (300 mg, 75%, **Intermediate 14**) as a yellow solid and was used in the next step directly without further purification; ¹H NMR δ (DMSO-d₆, 300 MHz) 3.77 (3H, s), 3.83 (3H, s), 7.39 - 7.49 (1H, m), 7.70 (1H, d), 7.83 (1H, s), 8.25 - 8.43 (2H, m), 8.71 (1H, d), 9.33 (1H, br s), 10.26 (1H, br s), 12.91 (1H, s); m/z (ES⁺), [M+H]⁺ = 366.

The procedure described above was repeated using the indicated aminopyrazole and Starting Intermediate to give Intermediates 15-22 described in Table 4:

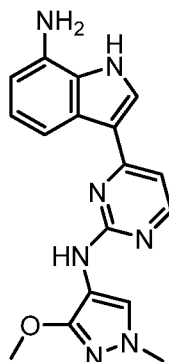
Table 4

Intermediate	Starting Intermediate	Aminopyrazole	NMR δ (300 MHz)	m/z [M+H] ⁺	Yield %
15 5-fluoro-N-(3-methoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)-4-(7-nitro-1 <i>H</i> -indol-3-yl)pyrimidin-2-amine	10		DMSO-d ₆ 3.71 (3H, s), 3.78 (3H, s), 7.32 (1H, t), 7.68 (1H, s), 8.13 (1H, d), 8.17 - 8.27 (1H, m), 8.37 (1H, d), 8.55 (1H, s), 12.52 (1H, s)	384	84

Intermediate	Starting Intermediate	Aminopyrazole	NMR δ (300 MHz)	m/z [M+H] ⁺	Yield %
16 <i>N</i> -(3-methoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)-5-methyl-4-(7-nitro-1 <i>H</i> -indol-3-yl)-2-pyrimidinamine	11		DMSO-d ₆ 2.36 (3H, s), 3.68 (3H, s), 3.80 (3H, s), 7.26 (1H, t), 7.65 (1H, s), 8.01 (1H, s), 8.15 – 8.23 (2H, m), 8.93 (1H, br s), 12.35 (1H, br s)	380	95
17 5-chloro- <i>N</i> -(3-methoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)-4-(7-nitro-1 <i>H</i> -indol-3-yl)pyrimidin-2-amine	12		Not obtained	400	55
18 5-bromo- <i>N</i> -(3-methoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)-4-(7-nitro-1 <i>H</i> -indol-3-yl)pyrimidin-2-amine	13		Not obtained	444	80
19 <i>N</i> -(3-ethoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)-5-fluoro-4-(7-nitro-1 <i>H</i> -indol-3-yl)pyrimidin-2-amine	10		DMSO-d ₆ 1.25 (3H, t), 3.73 (3H, s), 4.18 (2H, q), 7.20 - 7.53 (1H, m), 7.68 (1H, s), 8.11 - 8.30 (2H, m), 8.35 (1H, d), 8.43 (1H, s), 9.04 (1H, s)	398	74
20 <i>N</i> -(3-ethoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)-5-methyl-4-(7-nitro-1 <i>H</i> -indol-3-yl)pyrimidin-2-amine	11		DMSO-d ₆ 1.25 (3H, t), 2.36 (3H, s), 3.68 (3H, s), 4.16 (2H, q), 7.27 (1H, t), 7.64 (1H, s), 7.95 - 8.15 (2H, m), 8.20 (2H, m), 8.94 (1H, br s), 12.35 (1H, s)	394	74

Intermediate	Starting Intermediate	Aminopyrazole	NMR δ (300 MHz)	m/z [M+H] ⁺	Yield %
21 <i>N</i> -(1,3-dimethyl-1 <i>H</i> -pyrazol-4-yl)-5-methyl-4-(7-nitro-1 <i>H</i> -indol-3-yl)pyrimidin-2-amine	11		DMSO-d ₆ 2.09 (3H, s), 2.47 (3H, s), 3.81 (3H, s), 7.38 (1H, s), 7.87 (1H, s), 8.19 - 8.33 (3H, m), 9.66 (1H, s), 12.79 (1H, s)	364	62
22 5-chloro- <i>N</i> -(3-ethoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)-4-(7-nitro-1 <i>H</i> -indol-3-yl)pyrimidin-2-amine	12		Not obtained	414	100

Intermediate 23: 3-{2-[(3-Methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-amine



5

Iron (0.46 g, 8.2 mmol) was added to *N*-(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)-4-(7-nitro-1*H*-indol-3-yl)pyrimidin-2-amine (0.6 g, 1.6 mmol, **Intermediate 14**) and ammonium chloride (0.88 g, 16.4 mmol) in THF (100 mL) and water (50 mL) at 25°C under nitrogen. The resulting mixture was stirred at 80°C for 2 hours. The reaction mixture was filtered through diatomaceous earth. The solvent was removed under reduced pressure and the crude product was purified by flash C18 silica chromatography, elution gradient 30 to 80% methanol in water. Pure fractions were evaporated to dryness to afford 3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-amine (0.41 g, 74%, **Intermediate 23**) as a yellow solid; ¹H NMR δ (DMSO-d₆, 400 MHz) 3.68 (3H, s), 3.72 (3H, s), 5.10 (2H, s), 6.40 (1H, d), 6.82 (1H, t), 7.05 (1H, d), 7.60 - 7.73 (2H, m), 8.05 (1H, s), 8.10 - 8.21 (2H, m), 11.29 (1H, s); m/z (ES⁺), [M+H]⁺ = 336.

10

15

The procedure described above was repeated using the indicated Starting Intermediate to

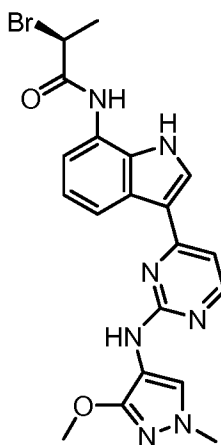
give Intermediates 24-31 described in Table 5:

Table 5

Intermediate	Starting Intermediate	NMR δ (300 MHz)	m/z [M+H] ⁺	Yield %
24 3-{5-fluoro-2-[(3-methoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]pyrimidin-4-yl}-1 <i>H</i> -indol-7-amine	15	DMSO-d ₆ 3.71 (3H, s), 3.79 (3H, s), 5.17 (2H, s), 6.44 (1H, d), 6.83 (1H, t), 7.67 (1H, s), 7.79 (1H, s), 8.04 - 8.10 (1H, m), 8.19 - 8.26 (2H, m), 11.48 (1H, s)	354	79
25 3-{2-[(3-methoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1 <i>H</i> -indol-7-amine	16	DMSO-d ₆ 2.32 (3H, s), 3.66 (3H, s), 3.79 (3H, s), 5.10 (2H, br s), 6.40 (1H, dd), 6.79 (1H, t), 7.68 (2H, s), 7.81 - 7.92 (2H, m), 8.11 (1H, s), 11.23 (1H, s)	350	61
26 3-{5-chloro-2-[(3-methoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]pyrimidin-4-yl}-1 <i>H</i> -indol-7-amine	17	DMSO-d ₆ 3.77 (3H, s), 3.81 (3H, s), 5.12 (2H, s), 6.40 (1H, d), 6.79 (1H, d), 6.88 (1H, s), 7.55 - 7.60 (1H, m), 8.26 (1H, s), 8.35 (2H, s), 11.41 (1H, s)	370	67
27 3-{5-bromo-2-[(3-methoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]pyrimidin-4-yl}-1 <i>H</i> -indol-7-amine	18	DMSO-d ₆ 3.63 (3H, s), 3.78 (3H, s), 5.12 (2H, s), 6.40 (1H, d), 6.90 (1H, t), 7.60 - 7.65 (2H, m), 8.37 (2H, s), 8.43 (1H, d), 11.37 (1H, s)	414	92
28 3-{2-[(1,3-dimethyl-1 <i>H</i> -pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1 <i>H</i> -indol-7-amine	21	Not obtained	334	70
29 3-{2-[(3-ethoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]-5-fluoropyrimidin-4-yl}-1 <i>H</i> -indol-7-amine	19	DMSO-d ₆ 1.25 (3H, t), 3.68 (3H, s), 4.14 (2H, q), 5.15 (2H, s), 6.43 (1H, d), 6.82 (1H, t), 7.64 (1H, s), 7.78 (1H, s), 8.01 - 8.17 (2H, m), 8.22 (1H, d), 11.46 (1H, s)	368	67
30 3-{2-[(3-ethoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1 <i>H</i> -indol-7-amine	20	Methanol-d ₄ 1.38 (3H, t), 2.39 (3H, s), 3.72 (3H, s), 4.26 (2H, q), 6.64 (1H, d), 6.96 (1H, t), 7.74 (3H, m), 8.14 (1H, s) - four exchangeable protons not observed	364	93

Intermediate	Starting Intermediate	NMR δ (300 MHz)	m/z [M+H] ⁺	Yield %
31 3-{5-chloro-2-[(3-ethoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]pyrimidin-4-yl}-1 <i>H</i> -indol-7-amine	22	Not obtained	384	56

Intermediate 32: (2*S*)-2-Bromo-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)propanamide



1-Propanephosphonic acid cyclic anhydride (25.6 g, 40.3 mmol) was added dropwise to 3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-amine (4.5 g, 13.4 mmol, **Intermediate 23**), (*S*)-2-bromopropanoic acid (4.1 g, 26.8 mmol) and pyridine (3.3 mL, 40.3 mmol) in ethyl acetate (100 mL) at -50°C over a period of 30 minutes under nitrogen. The resulting mixture was stirred at -50°C for 1 hour. The reaction was allowed to warm up to -15°C and stirred for 16 hour. The reaction mixture was quenched with ice water (100 mL), extracted with ethyl acetate (3 x 200 mL), the organic layer was dried, filtered and evaporated to afford a tan solid. The crude product was purified by flash silica chromatography, elution gradient 100 to 0% petroleum ether in ethyl acetate. Pure fractions were evaporated to dryness to afford (2*S*)-2-bromo-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)propanamide (4.9 g, 78%, **Intermediate 32**) as a yellow solid; ¹H NMR δ (DMSO-*d*₆, 400 MHz) 1.76 (3H, d), 3.60 (1H, m), 3.62 (3H, s), 3.71 (3H, s), 4.83 (1H, q), 7.01 - 7.16 (2H, m), 7.41 (1H, d), 7.71 (1H, s), 8.20 (1H, d), 8.26 (1H, d), 8.27 (1H, s), 10.12 (1H, s), 11.26 (1H, s); m/z (ES⁺), [M+H]⁺ = 470.

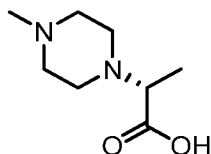
The procedure described above was repeated using the indicated Starting Intermediate to give Intermediates 33-36 described in Table 6:

Table 6

Intermediate	Starting Intermediate	NMR δ (300 MHz)	m/z [M+H] ⁺	Yield %
33 (2 <i>S</i>)-2-bromo- <i>N</i> -(3-{5-fluoro-2-[(3-methoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]pyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)propanamide	24	DMSO-d ₆ 1.85 (3H, d), 3.72 (3H, s), 3.79 (3H, s), 4.81 (1H, q), 7.09 (1H, t), 7.50 (1H, d), 7.67 (1H, s), 8.19 (1H, t), 8.29 (1H, d), 8.39 (2H, d), 10.22 (1H, s), 11.43 (1H, s)	490	94
34 (2 <i>S</i>)-2-bromo- <i>N</i> -(3-{2-[(3-methoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)propanamide	25	DMSO-d ₆ 1.84 (3H, d), 2.33 (3H, s), 3.68 (3H, s), 3.78 (3H, s), 4.80 (1H, q), 7.03 (1H, t), 7.45 (1H, d), 7.66 (1H, s), 7.99 (1H, s), 8.00 (1H, s), 8.15 (1H, s), 8.26 (1H, s), 10.14 (1H, s), 11.22 (1H, s)	484	66
35 (2 <i>S</i>)-2-bromo- <i>N</i> -(3-{2-[(3-ethoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]-5-fluoropyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)propanamide	29	DMSO-d ₆ 1.24 (3H, t), 1.84 (3H, d), 3.70 (3H, s), 4.14 (2H, q), 4.80 (1H, q), 7.07 (1H, t), 7.49 (1H, d), 7.65 (1H, s), 8.18 (1H, t), 8.28 (3H, d), 10.21 (1H, s), 11.27 - 11.63 (1H, m)	502	86
36 (2 <i>S</i>)-2-bromo- <i>N</i> -(3-{2-[(3-ethoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1 <i>H</i> -indol-7-yl)propanamide	30	DMSO-d ₆ 1.30 (3H, t), 1.86 (3H, d), 2.35 (3H, s), 3.60 - 3.65 (1H, m), 3.68 (3H, s), 4.11 (2H, q), 4.84 (1H, q), 7.03 (1H, t), 7.47 (1H, d), 7.66 (1H, s), 7.95 (1H, s), 8.17 (1H, s), 8.28 (1H, s), 10.15 (1H, s), 11.23 (1H, s)	500 (Br isotope value)	62

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Intermediate 37: (*R*)-2-(4-Methylpiperazin-1-yl)propanoic acid dihydrochloride

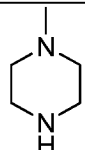
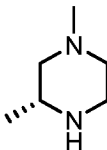


Trifluoromethanesulfonic anhydride (53.6 mL, 317 mmol) was added dropwise to (*S*)-methyl 2-hydroxypropanoate (30 g, 288 mmol) and 2,6-lutidine (37 mL, 317 mmol) in DCM (500 mL) at -78°C over a period of 1 hour. The resulting solution was stirred at -78°C for 0.5 hours.

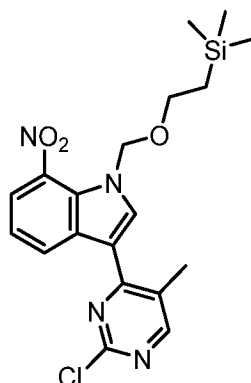
The solution was then warmed to room temperature for 1 hour. The organic phase was washed with 1N HCl (aq.) (2 x 100 mL) and dried over sodium sulfate, then filtered and evaporated. The residue was dissolved in DCM (500 mL), cooled to 0°C, then 1-methylpiperazine (65 g, 646 mmol) was added slowly. Potassium carbonate (212 g, 1537 mmol) in water (700 mL) was added dropwise at 0°C. The solution was stirred at 25°C overnight, then washed with brine, dried over sodium sulfate, filtered and evaporated to give a yellow oil. 6N (aq.) HCl (270 mL, 1625 mmol) was added in one portion at 25°C and the resulting mixture was stirred at 110°C for 18 hours. The solution was evaporated and the product washed with acetonitrile (200 mL) to afford an off-white solid. This solid was suspended in isopropanol (1000 mL) and was stirred for 3 hours at 100°C and then stirred for 16 hours at room temperature. The precipitate was collected by filtration, washed with isopropanol (150 mL) and dried under vacuum to afford (2*R*)-2-(4-methylpiperazin-1-yl)propanoic acid dihydrochloride (15 g, 48%, **Intermediate 37**) as a white solid; ¹H NMR δ (D₂O, 400 MHz) 1.51 (3H, d), 2.94 (3H, s), 3.48 - 4.13 (9H, m); m/z (ES⁺), [M+H]⁺ = 173.

The procedure described above was repeated using the indicated 2-hydroxypropanoate and piperazine to give the Intermediates 38 and 39 described in Table 7:

Table 7

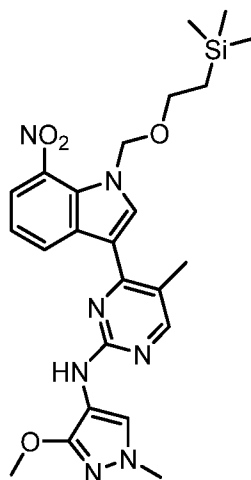
Intermediate	Piperazine	Hydroxy-propanoate	NMR δ (300 MHz)	m/z [M+H] ⁺	Yield %
38 (2 <i>S</i>)-2-(4-methylpiperazin-1-yl)propanoic acid dihydrochloride		(<i>R</i>)-methyl 2-hydroxypropanoate	DMSO-d ₆ 1.54 (3H, d), 2.83 (3H, s), 3.69 (8H, d), 4.30 (1H, s), 11.84 (1H, s)	173	100
39 (2 <i>R</i>)-2-[(2 <i>R</i>)-2,4-dimethylpiperazin-1-yl]propanoic acid dihydrochloride		(<i>S</i>)-methyl 2-hydroxypropanoate	Methanol-d ₄ 1.26 (3H, d), 1.43 (3H, d), 2.90 (4H, s), 3.05 - 3.21 (2H, m), 3.34 - 3.40 (2H, m), 3.42 - 3.49 (1H, m), 3.53 (1H, br s), 4.07 (1H, q)	187	95

Intermediate 40: 3-(2-Chloro-5-methyl-4-pyrimidinyl)-7-nitro-1-[[2-(trimethylsilyl)ethoxy]methyl]-1H-indole



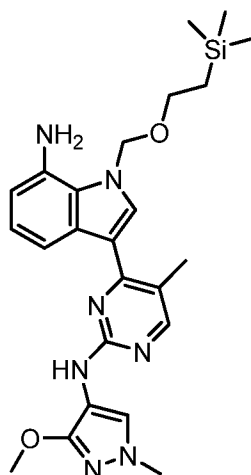
Sodium hydride (60% dispersion in mineral oil) (1.3 g, 33 mmol) was added portion-wise
 5 to a stirred suspension of 3-(2-chloro-5-methyl-4-pyrimidinyl)-7-nitro-1H-indole (6.4 g, 22 mmol, **Intermediate 11**) in anhydrous THF (150 mL) at 0°C. After stirring for 25 minutes, (2-(chloromethoxy)ethyl)trimethylsilane (4.1 mL, 23 mmol) was added rapidly dropwise. After 5 minutes the cooling bath was removed and the reaction left to stir at ambient temperature for 1.5 hours. Additional sodium hydride (60% dispersion in mineral oil) (130 mg, 3.3 mmol) and (2-
 10 (chloromethoxy)ethyl)trimethylsilane (0.4 mL, 2.3 mmol) were added. The reaction was stirred for an additional 40 minutes then quenched with saturated aqueous NaHCO₃ and the pale yellow mixture was diluted with ether. The layers were separated and the aqueous layer extracted with ether. The combined organic layer was washed with brine, dried over magnesium sulfate, filtered and evaporated. The residue was dissolved in chloroform and was subject to
 15 silica gel chromatography using 5 - 45% ethyl acetate - hexane as eluent to afford 3-(2-chloro-5-methyl-4-pyrimidinyl)-7-nitro-1-[[2-(trimethylsilyl)ethoxy]methyl]-1H-indole (9.2 g, 100%, **Intermediate 40**) as a yellow solid; ¹H NMR δ (DMSO-d₆, 400 MHz) -0.16 (9H, s), 0.60 - 0.73 (2H, m), 2.51 - 2.52 (3H, m), 3.11 - 3.22 (2H, m), 5.72 (2H, s), 7.48 (1H, t), 7.94 (1H, dd), 8.57 (1H, s), 8.64 (1H, s), 8.84 (1H, dd); m/z (ES⁺), [M+H]⁺ = 419.

Intermediate 41: N-(3-Methoxy-1-methyl-1*H*-pyrazol-4-yl)-5-methyl-4-(7-nitro-1-{[2-(trimethylsilyl)ethoxy]methyl}-1*H*-indol-3-yl)pyrimidin-2-amine



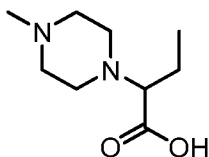
A mixture of dioxane and water (10:1, 44 mL) was added to a mixture of 3-(2-chloro-5-methyl-4-pyrimidinyl)-7-nitro-1-{[2-(trimethylsilyl)ethoxy]methyl}-1*H*-indole (1.59 g, 3.8 mmol, **Intermediate 40**), 3-methoxy-1-methyl-1*H*-pyrazol-4-amine dihydrochloride (1.7 g, 10.4 mmol), palladium(II) acetate (0.085 g, 0.4 mmol), Xantphos (0.22 g, 0.4 mmol) and cesium carbonate (4.95 g, 15.2 mmol) under nitrogen. The mixture was then heated at 110°C for 3.5 hours under nitrogen. The mixture was allowed to cool to ambient temperature, diluted with ethyl acetate, filtered through diatomaceous earth and concentrated. The resultant gum was subject to silica gel chromatography using 30 - 100% ethyl acetate - hexane as eluent to afford N-(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)-5-methyl-4-(7-nitro-1-{[2-(trimethylsilyl)ethoxy]methyl}-1*H*-indol-3-yl)pyrimidin-2-amine (1.3 g, 67%, **Intermediate 41**) as a pale yellow solid; ¹H NMR δ (DMSO-d₆, 400 MHz) -0.17 (9H, s), 0.60 - 0.74 (2H, m), 2.35 (3H, s), 3.05 - 3.20 (2H, m), 3.68 (3H, s), 3.79 (3H, s), 5.69 (2H, s), 7.23 - 7.36 (1H, m), 7.65 (1H, s), 7.86 (1H, d), 8.19 (1H, br s), 8.23 (1H, s), 8.35 (1H, s), 8.67 - 9.02 (1H, m); m/z (ES⁺), [M+H]⁺ = 510.

Intermediate 42: 3-{2-[(3-Methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1-{[2-(trimethylsilyl)ethoxy]methyl}-1*H*-indol-7-amine



A mixture of N-(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)-5-methyl-4-(7-nitro-1-{[2-(trimethylsilyl)ethoxy]methyl}-1*H*-indol-3-yl)pyrimidin-2-amine (1.26 g, 2.5 mmol, **Intermediate 41**) in methanol - ethyl acetate (1:1, 20 mL) was subject to hydrogenation at atmospheric pressure in the presence of 10% palladium on carbon (w/w) (0.26 g, 0.25 mmol) at ambient temperature for 23 hours. The slurry was diluted with ethyl acetate and filtered through diatomaceous earth, then concentrated to afford 3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1-{[2-(trimethylsilyl)ethoxy]methyl}-1*H*-indol-7-amine (1.1 g, 93%, **Intermediate 42**) as a pale yellow solid; ¹H NMR δ (DMSO-d₆, 400 MHz) -0.05 (9H, s), 0.86 - 0.96 (2H, m), 2.30 (3H, s), 3.51 - 3.62 (2H, m), 3.65 (3H, s), 3.79 (3H, s), 4.97 - 5.05 (2H, m), 5.72 (2H, s), 6.53 (1H, d), 6.83 (1H, t), 7.65 (1H, s), 7.70 (1H, br s), 7.90 (1H, s), 7.95 (1H, s), 8.13 (1H, s); m/z (ES⁺), [M+H]⁺ = 480.

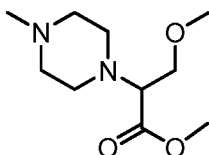
Intermediate 43: 2-(4-Methylpiperazin-1-yl)butanoic acid



Ethyl 2-bromobutanoate (30 g, 154 mmol) was added dropwise to 1-methylpiperazine (61.6 g, 615 mmol) in THF (500 mL) at 0°C over a period of 30 minutes under nitrogen. The resulting mixture was stirred at 25°C for 12 hours. The solvent was removed under reduced pressure. The mixture was made basic with saturated aqueous potassium carbonate and extracted with ethyl acetate (3 x 150 mL). The combined organic layers were evaporated to give a yellow oil which was added dropwise to 6N (aq.) HCl (200 mL, 1200 mmol) at 0°C over a

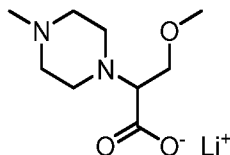
period of 10 minutes under air. The resulting mixture was stirred at 100°C for 16 hours. The reaction mixture was cooled to room temperature before being washed with ethyl acetate (100 mL). The water was removed under reduced pressure to give 2-(4-methylpiperazin-1-yl)butanoic acid hydrochloride (35 g, 96%, **Intermediate 43**) as a white solid that was used in the next step directly without further purification; ¹H NMR δ (DMSO-d₆, 300MHz) 0.93 (3H, t), 1.76 - 2.09 (2H, m), 2.80 (3H, s), 3.29 - 3.69 (8H, m), 3.99 (1H, br s), 11.84 (1H, s); m/z (ES⁺), [M+H]⁺ = 187.

Intermediate 44: Methyl 3-methoxy-2-(4-methylpiperazin-1-yl)propanoate



Potassium carbonate (1.38 g, 10.1 mmol) was added to a stirred solution of 1-methylpiperazine (0.93 mL, 8.3 mmol) and methyl 2-bromo-3-methoxypropanoate (1.7 g, 8.6 mmol) in acetonitrile (20 mL) under a nitrogen atmosphere. The pale yellow mixture was then warmed to 60°C for 21 hours. The reaction was cooled to ambient temperature, diluted with ethyl acetate and filtered. Concentration afforded methyl 3-methoxy-2-(4-methylpiperazin-1-yl)propanoate (1.69 g, 94%) as an orange oil which was used without further purification; ¹H NMR δ (DMSO-d₆, 400 MHz) 2.12 (3H, s), 2.20 - 2.36 (4H, m), 2.48 - 2.56 (5H, m), 3.21 - 3.25 (3H, s), 3.39 (1H, dd), 3.48 - 3.53 (1H, m), 3.63 (3H, s); m/z (ES⁺), [M+H]⁺ = 217.

Intermediate 45: Lithium 3-methoxy-2-(4-methylpiperazin-1-yl)propanoate



A solution of lithium hydroxide (52 mg, 2.2 mmol) in water (3 mL) was added to a stirred solution of methyl 3-methoxy-2-(4-methylpiperazin-1-yl)propanoate (0.47 g, 2.2 mmol, **Intermediate 44**) in THF (3 mL) at ambient temperature. After stirring for 21 hours the reaction was warmed to 40°C for 22 hours. A few drops of methanol were added, clarifying the pale yellow solution and heating continued. After 2 hours, additional lithium hydroxide (16 mg, 0.7 mmol) was added and the reaction left to stir for 4 days. The solvent was removed under reduced pressure and the aqueous solution lyophilized to afford the lithium 3-methoxy-2-(4-methylpiperazin-1-yl)propanoate (0.45 g, 98%) as an off-white solid; ¹H NMR δ (DMSO-d₆, 400

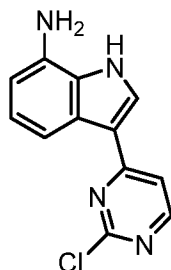
MHz) 2.10 (3H, s), 2.27 (4H, br s), 2.51 - 2.60 (4H, m), 2.87 (1H, t), 3.19 (3H, s), 3.50 - 3.60 (2H, m); m/z (ES+), [M+H]⁺ = 203.

The procedure described for Example 32 was repeated using the indicated Starting Intermediates to give Intermediates 46 and 47 described in Table 8:

Table 8

Intermediate	Starting Intermediates	NMR δ (400 MHz)	m/z [M+H] ⁺	Yield %
46 <i>N</i> -(3-{2-[(3-methoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1-{2-[(trimethylsilyl)ethoxy]ethyl}-1 <i>H</i> -indol-7-yl)-2-(4-methylpiperazin-1-yl)butanamide	42 and 43	DMSO-d ₆ -0.07 (9H, s), 0.86 (2H, dd), 0.98 (3H, t), 1.68 - 1.84 (2H, m), 2.33 (4H, s), 2.80 (2H, br s), 3.07 (3H, m), 3.17 (4H, s), 3.40 - 3.52 (4H, m), 3.67 (3H, s), 3.79 (3H, s), 5.66 - 5.75 (2H, m), 7.10 (1H, t), 7.34 (1H, d), 7.65 (1H, s), 8.05 (1H, br s), 8.11 (1H, s), 8.18 (1H, s), 9.33 (1H, br s), 9.72 (1H, s)	648	81
47 3-methoxy- <i>N</i> -(3-{2-[(3-methoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1-{2-[(trimethylsilyl)ethoxy]methyl}-1 <i>H</i> -indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide	42 and 45	DMSO-d ₆ -0.08 (9H, s), 0.83 - 0.94 (2H, m), 2.33 (3H, s), 2.58 - 3.14 (11H, m), 3.41 - 3.51 (3H, m), 3.67 (3H, s), 3.70 - 3.86 (6H, m), 5.64 - 5.86 (3H, m), 7.09 (1H, t), 7.30 (1H, d), 7.64 (1H, s), 8.05 (1H, s), 8.10 (1H, s), 8.18 (1H, s), 8.30 (1H, br s), 9.36 (1H, br s), 9.84 (1H, s)	664	82

Intermediate 48: 3-(2-Chloropyrimidin-4-yl)-1*H*-indol-7-amine



Ammonium chloride (7.8 g, 146 mmol) was added to 3-(2-chloro-4-pyrimidinyl)-7-nitro-1*H*-indole (4 g, 14.6 mmol, **Intermediate 9**) and iron (4.1 g, 72 mmol) in THF (200 mL) and water (100 mL) at 25°C under nitrogen. The resulting mixture was stirred at 80°C for 12 hours. The reaction mixture was filtered through diatomaceous earth. The organic phase was separated and the aqueous phase was extracted with THF (2 x 100 mL). The organic phases were

combined and concentrated to afford 3-(2-chloropyrimidin-4-yl)-1*H*-indol-7-amine (3 g, 84%, **Intermediate 48**) as a green solid; ¹H NMR δ (DMSO-d₆, 400 MHz) 5.42 (2H, s), 6.44 (1H, d), 6.91 (1H, m), 7.60 (1H, d), 7.78 - 7.91 (1H, m), 8.36 (1H, s), 8.45 (1H, d)-NH proton obscured; m/z (ES⁺), [M+MeCN]⁺ = 286.

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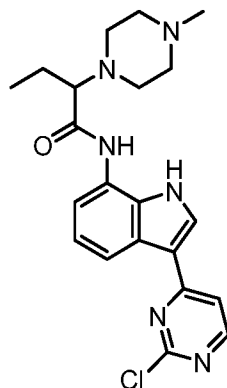
The procedure described above for Intermediate 48 was repeated using the indicated Starting Intermediate to give Intermediates 49 and 50 described in Table 9:

Table 9

Intermediate	Starting Intermediate	NMR δ (400 MHz)	m/z [M+H] ⁺	Yield %
49 3-(2-chloro-5-methylpyrimidin-4-yl)-1 <i>H</i> -indol-7-amine	11	DMSO-d ₆ 2.50 (3H, s), 5.26 (2H, s), 6.47 (1H, dd), 6.92 (1H, t), 7.76 (1H, d), 8.12 (1H, d), 8.45 (1H, s), 11.71 (1H, d)	259	80
50 3-(2-chloro-5-fluoropyrimidin-4-yl)-1 <i>H</i> -indol-7-amine	10	Methanol-d ₄ 6.68 (1H, dd), 7.06 (1H, t), 8.04 (1H, dd), 8.18 (1H, d), 8.39 (1H, d) - three exchangeable protons not observed	263	79

10

Intermediate 51: N-[3-(2-Chloropyrimidin-4-yl)-1*H*-indol-7-yl]-2-(4-methylpiperazin-1-yl)butanamide



1-Propanephosphonic acid cyclic anhydride (7.8 g, 12.3 mmol) was added dropwise to 3-(2-chloropyrimidin-4-yl)-1*H*-indol-7-amine (1 g, 4.1 mmol, **Intermediate 48**), 2-(4-methylpiperazin-1-yl)butanoic acid dihydrochloride (1.3 g, 4.9 mmol, **Intermediate 43**) and pyridine (2 mL, 25 mmol) in DMF (100 mL) at 0°C over a period of 10 minutes under nitrogen. The resulting mixture was stirred at 25°C for 2 hours. The solvent was removed under reduced pressure and the crude product purified by reverse phase silica (C18) gel chromatography using 0 - 100% methanol in water to give N-[3-(2-chloropyrimidin-4-yl)-1*H*-indol-7-yl]-2-(4-methylpiperazin-1-yl)butanamide (0.24 g, 14%, **Intermediate 51**) as a yellow solid; ¹H NMR δ

20

(Methanol-d₄, 300 MHz) 1.08 (3H, t), 1.79 - 2.02 (2H, m), 2.32 (3H, s), 2.58 (4H, s), 2.84 (4H, m), 3.10 - 3.27 (1H, m), 7.25 (2H, m), 7.77 (1H, d), 8.27 (1H, s), 8.37 (1H, dd), 8.44 (1H, d)-two exchangeable protons not observed; m/z (ES⁺), [M+H]⁺ = 413.

5 The procedure described above for Intermediate 51 was repeated using the indicated Starting Intermediates to give Intermediates 52-55 described in Table 10:

Table 10

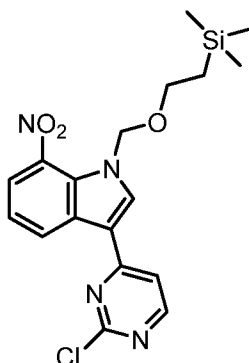
Intermediate	Starting Intermediates	NMR δ (300 MHz)	m/z [M+H] ⁺	Yield %
52 N-[3-(2-chloro-5-fluoropyrimidin-4-yl)-1H-indol-7-yl]-2-(4-methylpiperazin-1-yl)butanamide	50 and 43	Not obtained	431	89
53 N-[3-(2-chloro-5-fluoropyrimidin-4-yl)-1H-indol-7-yl]-3-methoxy-2-(4-methylpiperazin-1-yl)propanamide	50 and 45	DMSO-d ₆ 2.27 (3H, s), 2.41 (4H, br s), 2.64 (2H, m), 2.77 (2H, m), 3.29 (3H, s), 3.50 - 3.73 (2H, m), 3.79 (1H, dd), 7.23 (1H, t), 7.59 (1H, dd), 8.29 - 8.44 (2H, m), 8.71 (1H, d), 9.97 (1H, s), 11.85 (1H, s)	447	47
54 ^{1,2} (R)-N-[3-(2-chloro-5-methylpyrimidin-4-yl)-1H-indol-7-yl]-2-(4-methylpiperazin-1-yl)propanamide	49 and 37	DMSO-d ₆ 1.28 (3H, d), 2.24 (3H, s), 2.51 - 2.71 (11H, m), 3.39 (1H, m), 7.18 (1H, t), 7.50 (1H, d), 8.19 (1H, s), 8.31 (1H, d), 8.52 (1H, s), 9.76 (1H, s), 11.69 (1H, s)	413	81
55 ³ (R)-N-[3-(2-chloro-5-fluoropyrimidin-4-yl)-1H-indol-7-yl]-2-(4-methylpiperazin-1-yl)propanamide	50 and 37	DMSO-d ₆ 1.27 (3H, d), 2.27 (3H, s), 2.35 - 2.68 (8H, m), 3.39 (1H, q), 7.24 (1H, t), 7.53 (1H, d), 8.34 - 8.37 (2H, m), 8.72 (1H, d), 9.83 (1H, s), 11.90 (1H, s)	417	79

¹H NMR analysis was performed using a Bruker Avance 400 (400 MHz) spectrometer.

² The indicated amino acid (1.5 equiv) and 7-amino-indole intermediates were reacted in the presence of HATU (2 equiv) and diisopropylethylamine (4 equiv) in DMF at room temperature.

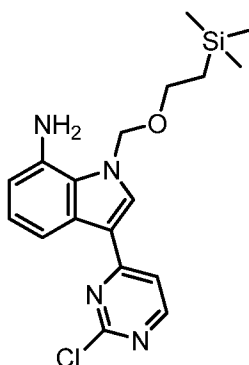
³ The indicated amino acid (1.3 equiv) and 7-amino-indole intermediates were reacted in the presence of HATU (1.5 equiv) and diisopropylethylamine (5 equiv) in DMF at room temperature.

Intermediate 56: 3-(2-Chloropyrimidin-4-yl)-7-nitro-1-{[2-(trimethylsilyl)ethoxy]methyl}-1H-indole



3-(2-Chloropyrimidin-4-yl)-7-nitro-1H-indole (4.4 g, 16 mmol, **Intermediate 9**) was dissolved in THF (60 mL) and cooled to 0°C. Sodium hydride (1.2 g, 29 mmol) was then added and the reaction mixture was warmed to room temperature. (2-(Chloromethoxy)ethyl)trimethylsilane (4.3 mL, 24 mmol) was then slowly added and the reaction mixture was allowed to stir for 1.5 hours. The reaction mixture was quenched with aqueous sodium bicarbonate, and extracted with ethyl acetate. The organic layers were dried over sodium sulfate, filtered, and concentrated giving the crude product as a red oil which was purified via silica gel column chromatography using 0 - 40% ethyl acetate - hexanes as eluent to give 3-(2-chloropyrimidin-4-yl)-7-nitro-1-{[2-(trimethylsilyl)ethoxy]methyl}-1H-indole (5.7 g, 88%, **Intermediate 56**) as a yellow solid; ¹H NMR δ (DMSO-d₆, 300 MHz) -0.16 (9H, s), 0.67 (2H, t), 3.19 (2H, t), 5.66 (2H, s), 7.51 (1H, s), 7.88 - 8.01 (2H, m), 8.71 (1H, d), 8.88 (1H, s), 8.91 (1H, s); m/z (ES⁺), [M+H]⁺ = 405.

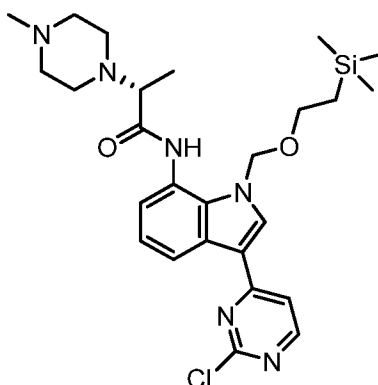
Intermediate 57: 3-(2-Chloropyrimidin-4-yl)-1-{[2-(trimethylsilyl)ethoxy]methyl}-1H-indol-7-amine



3-(2-Chloropyrimidin-4-yl)-7-nitro-1-{[2-(trimethylsilyl)ethoxy]methyl}-1H-indole (5.7 g, 14.1 mmol, **Intermediate 56**) was dissolved in methanol (47 mL), THF (47 mL) and water (47 mL).

The solution was then heated to 60°C and ammonium chloride (32.8 g, 612 mmol) was added, followed by iron (34.4 g, 617 mmol). The solution was then allowed to stir for 2 hours at 60°C. The reaction mixture was partitioned between water and diethyl ether. The organic layers were extracted and combined, dried over sodium sulfate, filtered, and concentrated to give 3-(2-chloropyrimidin-4-yl)-1-[[2-(trimethylsilyl)ethoxy]methyl]-1*H*-indol-7-amine (5.2 g, 98%,
 5 **Intermediate 57**) as a yellow oil; ¹H NMR δ (DMSO-d₆, 300 MHz) -0.04 (9H, m), 0.92 (2H, t), 3.60 (2H, t), 5.15 (2H, s), 5.71 (2H, s), 6.59 - 6.62 (1H, m), 7.00 (1H, t), 7.70 (1H, dd), 7.80 (1H, d), 8.50 (1H, s), 8.55 (1H, d); m/z (ES⁺), [M+H]⁺ = 375.

10 **Intermediate 58: (2*R*)-N-[3-(2-Chloropyrimidin-4-yl)-1-[[2-(trimethylsilyl)ethoxy]methyl]-1*H*-indol-7-yl]-2-(4-methylpiperazin-1-yl)propanamide**

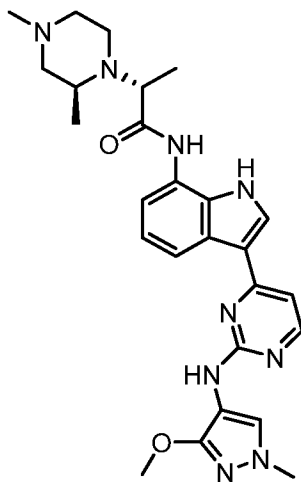


(*R*)-2-(4-Methylpiperazin-1-yl)propanoic acid dihydrochloride (2.45 g, 10 mmol, **Intermediate 37**) was dissolved in DMF (15 mL) and di(1*H*-imidazol-1-yl)methanone (1.3 g, 8 mmol) was added. Gas evolved, and the reaction mixture was allowed to stir under nitrogen at room temperature until the reaction mixture became homogeneous. 3-(2-Chloropyrimidin-4-yl)-1-[[2-(trimethylsilyl)ethoxy]methyl]-1*H*-indol-7-amine (1.5 g, 4 mmol, **Intermediate 57**) in DMSO (11 mL) was then added and the reaction mixture was stirred overnight. The reaction was quenched with 10% potassium carbonate solution and extracted with ethyl acetate. The organic
 20 layers were combined, dried over sodium sulfate, filtered, and concentrated. The crude product was purified via silica gel column chromatography using 100% ethyl acetate then 0 - 20% methanol - DCM as eluent to give (2*R*)-N-[3-(2-chloropyrimidin-4-yl)-1-[[2-(trimethylsilyl)ethoxy]methyl]-1*H*-indol-7-yl]-2-(4-methylpiperazin-1-yl)propanamide (0.92 g, 43%, **Intermediate 58**) as a yellow solid; ¹H NMR δ (Chloroform-d, 300 MHz) -0.10 (9H, s), 0.96 - 1.08 (2H, m), 1.40 (3H, d), 2.36 (3H, m), 2.47 - 2.92 (8H, m), 3.27 (1H, q), 3.51 - 3.64 (2H, m), 5.54 - 5.79 (2H, m), 7.36 (1H, t), 7.58 (1H, d), 7.79 (1H, d), 7.96 (1H, s), 8.12 (1H, d), 8.54 (1H, d), 9.72 (1H, br s); m/z (ES⁺), [M+H]⁺ = 529.

Intermediate 59: (2*R*)-N-[3-(2-Chloropyrimidin-4-yl)-1*H*-indol-7-yl]-2-(4-methylpiperazin-1-yl)propanamide

(2*R*)-N-[3-(2-Chloropyrimidin-4-yl)-1-[[2-(trimethylsilyl)ethoxy]methyl]-1*H*-indol-7-yl]-2-(4-methylpiperazin-1-yl)propanamide (0.38 g, 0.7 mmol, **Intermediate 58**) was dissolved in DMSO (7 mL) and cesium fluoride (0.32 g, 2.1 mmol) was added. The reaction mixture was then heated at 100°C and allowed to stir for 2 hours. The reaction mixture was diluted with ethyl acetate and water. The organic layer was separated and the aqueous layer was extracted with ethyl acetate. The organic layers were combined, dried over sodium sulfate, filtered, and concentrated to give (2*R*)-N-[3-(2-chloropyrimidin-4-yl)-1*H*-indol-7-yl]-2-(4-methylpiperazin-1-yl)propanamide (0.17 g, 59%) as a yellow solid; ¹H NMR δ (Chloroform-d, 300 MHz) 1.43 (3H, d), 2.32 - 3.02 (11H, m), 3.39 (1H, m), 6.84 (1H, m), 7.22 - 7.25 (1H, m), 7.52 (1H, d), 8.03 (1H, d), 8.34 (1H, d), 8.47 (1H, d), 9.81 (1H, br. s), 11.58 (1H, br s); m/z (ES+), [M+H]⁺ = 399.

Example 1: (2*R*)-2-[(2*S*)-2,4-Dimethylpiperazin-1-yl]-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)propanamide

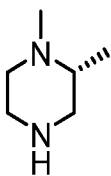
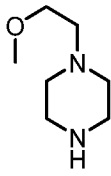
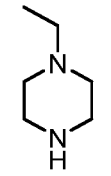
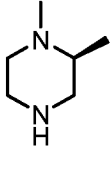


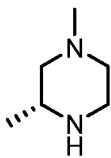
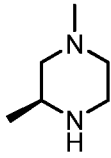
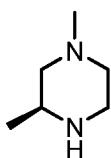
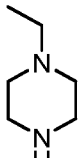
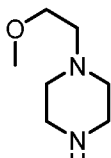
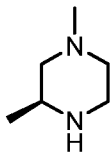
(*S*)-1,3-Dimethylpiperazine dihydrochloride (0.16 g, 0.85 mmol) was added in one portion to (2*S*)-2-bromo-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)propanamide (0.2 g, 0.43 mmol, **Intermediate 32**) and potassium carbonate (0.24 g, 1.7 mmol) in DMF (2 mL) at 0°C. The resulting solution was stirred at 25°C for 16 hours. The crude product was purified by preparative HPLC (X Bridge C18, 5 μm, 19x150 mm; Mobile Phase A: water/0.05% TFA, Mobile Phase B: acetonitrile; Flow rate: 20 mL/min; Gradient: 20%B to 70%B in 10 min; 254 nm) to afford (2*R*)-2-[(2*S*)-2,4-dimethylpiperazin-1-yl]-N-(3-{2-[(3-

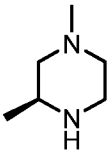
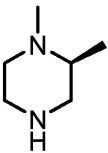
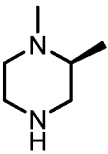
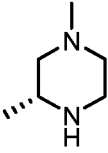
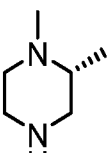
methoxy-1-methyl-1H-pyrazol-4-yl)amino]pyrimidin-4-yl}-1H-indol-7-yl)propanamide (49 mg, 23%, **Example 1**) as a white solid; ¹H NMR δ (Methanol-d₄, 400 MHz) 1.15 (3H, d), 1.43 (3H, d), 2.06 (1H, t), 2.32 (3H, s), 2.41 (1H, m), 2.75 - 2.92 (3H, m), 3.01 (2H, m), 3.79 - 3.83 (4H, m), 3.94 (3H, s), 7.12 - 7.19 (3H, dt), 7.73 (1H, s), 8.09 (1H, s), 8.19 (1H, d), 8.27 (1H, s); m/z (ES⁺), [M+H]⁺ = 504; chiral HPLC (ChiralPak ADH, 5 μm, 0.46x10 cm, mobile phase: 1:1 hexanes (modified with 0.1% TEA) and ethanol at 1.0mL/min) indicates 99:1 e.r., retention time = 9.24 min.

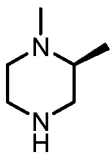
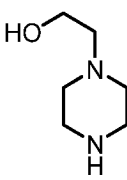
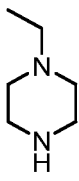
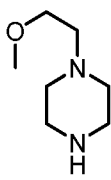
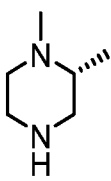
The procedure described above for Example 1 was repeated using the indicated piperazine and Starting Intermediate to give the compounds described in Table 11:

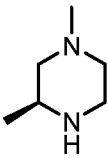
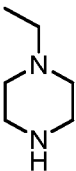
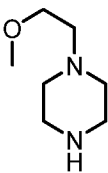
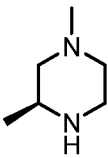
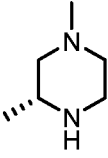
Table 11

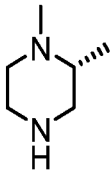
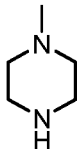
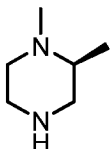
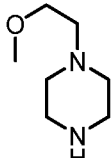
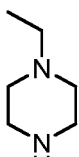
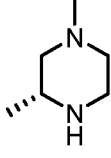
Example	Starting Intermediate	Piperazine	NMR δ (300 MHz)	m/z [M+H] ⁺	Yield %
2	33		Methanol-d ₄ 1.14 (3H, d), 1.42 (3H, d), 2.26 - 2.38 (5H, m), 2.52 (2H, m), 2.86 - 3.01 (3H, m), 3.39 (1H, m), 3.79 (3H, s), 3.92 (3H, s), 7.10 - 7.22 (2H, m), 7.68 (1H, s), 8.14 (2H, dd), 8.42 (1H, s)	522	47
3 ¹	33		Methanol-d ₄ 1.42 (3H, d), 2.61 - 2.84 (10H, m), 3.36 (4H, s), 3.58 (2H, t), 3.79 (3H, s), 3.91 (3H, s), 7.10 - 7.22 (2H, m), 7.68 (1H, s), 8.14 (2H, dd), 8.42 (1H, s)	552	70
4 ¹	33		Methanol-d ₄ 1.15 (3H, t), 1.43 (3H, d), 2.51 (2H, q), 2.56 - 2.90 (8H, m), 3.34 - 3.45 (1H, m), 3.79 (3H, s), 3.92 (3H, s), 7.10 - 7.22 (2H, m), 7.68 (1H, s), 8.14 (2H, dd), 8.42 (1H, s)	522	62
5	33		Methanol-d ₄ 1.14 (3H, d), 1.42 (3H, dd), 2.18 (1H, t), 2.36 (4H, s), 2.50 (1H, m), 2.61 - 2.72 (1H, m), 2.86 - 3.00 (3H, m), 3.33 - 3.44 (1H, m), 3.79 (3H, s), 3.92 (3H, s), 7.10 - 7.22 (2H, m), 7.68 (1H, s), 8.14 (2H, dd), 8.42 (1H, s)	522	60

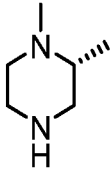
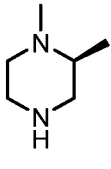
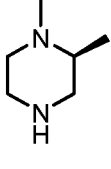
Example	Starting Intermediate	Piperazine	NMR δ (300 MHz)	m/z [M+H] ⁺	Yield %
6	33		Methanol-d4 1.23 (3H, d), 1.30 (3H, d), 2.20 (1H, m), 2.35 (3H, s), 2.42 (1H, m), 2.68 (1H, m), 2.79 (1H, m), 2.85 (3H, m), 3.80 (3H, s), 3.92 (3H, s), 4.06 (1H, m), 7.07 - 7.20 (2H, m), 7.68 (1H, s), 8.14 (2H, dd), 8.43 (1H, s)	522	22
7a ²	33		Methanol-d4 1.17 (3H, d), 1.45 (3H, d), 2.10 (1H, t), 2.33 (3H, s), 2.41 (1H, s), 2.72 - 2.96 (3H, m), 3.02 (2H, m), 3.73 - 3.87 (4H, m), 3.92 (3H, s), 7.16 (2H, d), 7.69 (1H, s), 8.15 (2H, dd), 8.43 (1H, s)	522	35
7b ²	33		Methanol-d4 1.23 (3H, d), 1.31 (3H, d), 2.17 (1H, m), 2.34 (4H, m), 2.60 - 2.94 (5H, m), 3.80 (3H, s), 3.92 (3H, s), 4.07 (1H, d), 7.13 (2H, m), 7.69 (1H, s), 8.15 (2H, dd), 8.44 (1H, s)	522	15
8 ¹	35		Methanol-d4 1.13 (3H, t), 1.32 (3H, t), 1.42 (3H, d), 2.49 (2H, d), 2.55 - 2.94 (8H, m), 3.40 (1H, m), 3.77 (3H, s), 4.23 (2H, q), 7.08 - 7.21 (2H, m), 7.66 (1H, s), 8.13 (2H, dd), 8.41 (1H, d)	536	62
9 ¹	35		Methanol-d4 1.32 (3H, t), 1.41 (3H, d), 2.52 - 2.95 (10H, m), 3.33 - 3.44 (4H, m), 3.56 (2H, t), 3.77 (3H, s), 4.23 (2H, q), 7.04 - 7.26 (2H, m), 7.66 (1H, s), 8.12 (2H, dd), 8.41 (1H, m)	566	57
10a ³	35		Methanol-d4 1.19 (3H, d), 1.34 (3H, t), 1.46 (3H, d), 2.25 (1H, t), 2.44 (3H, s), 2.58 (1H, m), 2.85 - 3.02 (3H, m), 3.03 - 3.13 (2H, m), 3.80 (3H, s), 3.82 - 3.90 (1H, m), 4.25 (2H, q), 7.09 - 7.20 (2H, m), 7.69 (1H, s), 8.15 (2H, dd), 8.43 (1H, s)	536	51

Example	Starting Intermediate	Piperazine	NMR δ (300 MHz)	m/z [M+H] ⁺	Yield %
10b ³	35		Methanol-d4 1.19 - 1.40 (9H, m), 2.17 (1H, t), 2.34 (4H, s), 2.60 - 2.91 (5H, m), 3.79 (3H, s), 4.06 (1H, m), 4.25 (2H, q), 7.06 - 7.22 (2H, m), 7.69 (1H, s), 8.14 (2H, dd), 8.44 (1H, d)	536	12
11a ⁴	35		Methanol-d4 1.13 (3H, d), 1.32 (3H, t), 1.41 (3H, d), 2.16 (1H, t), 2.33 (4H, s), 2.47 (1H, td), 2.65 (1H, td), 2.81 - 2.99 (3H, m), 3.32 - 3.43 (1H, m), 3.77 (3H, s), 4.23 (2H, q), 7.07 - 7.22 (2H, m), 7.67 (1H, s), 8.13 (2H, dd), 8.41 (1H, d)	536	84
11b ⁴	35		Methanol-d4 1.11 (3H, d), 1.32 (3H, t), 1.40 (3H, d), 2.22 - 2.37 (5H, m), 2.42 - 2.59 (2H, m), 2.84 - 3.02 (3H, m), 3.31 - 3.44 (1H, m), 3.77 (3H, s), 4.23 (2H, q), 7.07 - 7.22 (2H, m), 7.67 (1H, s), 8.13 (2H, dd), 8.41 (1H, s)	536	12
12	35		Methanol-d4 1.21 (3H, d), 1.26 - 1.38 (6H, m), 2.16 (1H, t), 2.33 (4H, s), 2.57 - 2.95 (5H, m), 3.77 (3H, s), 4.04 (1H, q), 4.23 (2H, q), 6.91 - 7.22 (2H, m), 7.67 (1H, s), 8.12 (2H, dd), 8.42 (1H, d)	536	33
13	35		Methanol-d4 1.11 (3H, d), 1.32 (3H, t), 1.40 (3H, d), 2.22 - 2.40 (5H, m), 2.42 - 2.59 (2H, m), 2.90 (3H, dd), 3.35 - 3.42 (1H, m), 3.77 (3H, s), 4.23 (2H, q), 7.06 - 7.21 (2H, m), 7.67 (1H, s), 8.13 (2H, dd), 8.42 (1H, br s)	536	88

Example	Starting Intermediate	Piperazine	NMR δ (300 MHz)	m/z [M+H] ⁺	Yield %
14 ⁵	34		Methanol-d ₄ 1.13 (3H, d), 1.41 (3H, d), 2.16 (1H, t), 2.36 (6H, d), 2.54 - 2.41 (1H, m), 2.66 (1H, t), 2.91 (3H, q), 3.72 (3H, s), 3.92 (3H, s), 7.21 - 7.04 (2H, m), 7.70 (1H, s), 7.87 (1H, s), 8.14 (1H, s), 8.20 (1H, d)	518	19
15	34		DMSO-d ₆ 1.26 (3H, d), 2.33 (3H, s), 2.35 - 2.78 (10H, m), 3.33 - 3.35 (1H, m), 3.42 - 3.65 (2H, m), 3.67 (3H, s), 3.79 (3H, s), 4.36 (1H, s), 7.00 (1H, t), 7.41 (1H, d), 7.66 (1H, s), 7.95 (1H, s), 7.97 (1H, s), 8.14 (1H, s), 8.16 (1H, br s), 9.65 (1H, s), 11.28 (1H, s)	534	39
16	34		Chloroform-d 1.17 (3H, t), 1.40 (3H, d), 2.35 (3H, s), 2.39 - 3.02 (10H, m), 3.37 (1H, d), 3.70 (3H, s), 3.98 (3H, s), 6.52 (1H, s), 6.81 (1H, d), 7.12 (1H, t), 7.66 (1H, d), 7.82 (1H, s), 8.14 - 8.30 (2H, m), 9.78 (1H, s), 11.17 (1H, s)	518	75
17	34		Methanol-d ₄ 1.40 (3H, d), 2.35 (3H, s), 2.75 (10H, m), 3.37 (4H, s), 3.57 (2H, t), 3.70 (3H, s), 3.98 (3H, s), 6.54 (1H, s), 6.81 (1H, d), 7.12 (1H, t), 7.66 (1H, d), 7.81 (1H, s), 8.17 - 8.32 (2H, m), 9.80 (1H, s), 11.17 (1H, s)	548	78
18	34		Methanol-d ₄ 1.25 (3H, s), 1.39 (3H, d), 2.35 (3H, s), 2.48 (4H, d), 2.62 (3H, s), 2.94 (3H, d), 3.45 (1H, s), 3.70 (3H, s), 3.98 (3H, s), 6.53 (1H, s), 6.82 - 7.19 (2H, m), 7.64 (1H, d), 7.81 (1H, s), 8.13 - 8.37 (2H, m), 9.68 (1H, s), 11.19 (1H, s)	518	22

Example	Starting Intermediate	Piperazine	NMR δ (300 MHz)	m/z [M+H] ⁺	Yield %
19	34		Methanol-d ₄ 1.11 (3H, d), 1.39 (3H, d), 1.95 - 2.11 (1H, m), 2.31 (7H, d), 2.61 - 2.89 (3H, m), 2.92 - 3.07 (2H, m), 3.68 (4H, s), 3.88 (3H, s), 6.98 - 7.17 (2H, m), 7.66 (1H, s), 7.82 (1H, s), 8.09 (2H, s)	518	6
20	36		DMSO-d ₆ 0.98 (3H, t), 1.20 - 1.31 (6H, m), 2.31-2.78 (13H, m), 3.31 (1H, s), 3.65 (3H, s), 4.15 (2H, q), 7.00 (1H, t), 7.41 (1H, d), 7.64 (1H, s), 7.86 - 7.98 (2H, m), 8.14 (1H, s), 8.22 (1H, s), 9.66 (1H, s), 11.30 (1H, s)	532	44
21	36		DMSO-d ₆ 1.20 - 1.31 (6H, m), 2.33 (3H, s), 2.45 - 2.70 (10 H, m), 3.21 (3H, s), 3.26 - 3.30 (1H, m), 3.41 (2H, t), 3.65 (3H, s), 4.15 (2H, q), 7.00 (1H, t), 7.41 (1H, d), 7.64 (1H, s), 7.86 - 7.99 (2H, m), 8.14 (1H, s), 8.22 (1H, s), 9.65 (1H, s), 11.29 (1H, s)	562	62
22	36		Chloroform-d 1.13 (3H, d), 1.44-1.62 (6H, m), 2.13 - 2.24 (1H, m), 2.25 - 2.63 (6H, m), 2.64 - 2.68 (2H, m), 2.71 - 2.83 (1H, m), 2.84 - 2.98 (3H, m), 3.53 (1H, s), 3.72 (3H, s), 4.32 (2H, q), 6.63 (1H, s), 6.83 (1H, s), 7.17 (1H, t), 7.34 (1H, s), 7.68 (1H, d), 7.87 (1H, s), 8.25 (2H, t), 11.18 (1H, s)	532	28
23	36		Methanol-d ₄ 1.20 - 1.43 (9H, m), 2.19 (1H, t), 2.38 (7H, d), 2.66 - 2.90 (5H, m), 3.73 (3H, s), 4.02 - 4.14 (1H, m), 4.26 (2H, q), 7.12 (2H, d), 7.73 (1H, s), 7.88 (1H, s), 8.13 - 8.27 (2H, m)	532	12

Example	Starting Intermediate	Piperazine	NMR δ (300 MHz)	m/z [M+H] ⁺	Yield %
24	36		Methanol-d4 1.14 (3H, d), 1.29 - 1.47 (6H, m), 2.28 - 2.45 (8H, m), 2.52 (2H, t), 2.86 - 3.04 (3H, m), 3.40 (1H, d), 3.73 (3H, d), 4.26 (2H, q), 7.15 (2H, dt), 7.73 (1H, d), 7.89 (1H, s), 8.13 - 8.27 (2H, m)	532	46
25	33		Methanol-d4 1.42 (3H, d), 2.34 (3H, s), 2.48 - 2.89 (8H, m), 3.40 (1H, m), 3.79 (3H, s), 3.91 (3H, s), 7.10 - 7.22 (2H, m), 7.68 (1H, s), 8.14 (2H, dd), 8.42 (1H, br s)	508	72
26	32		Methanol-d4 1.14 (3H, d), 1.41 (3H, d), 2.15 - 2.18 (1H, m), 2.36 (4H, m), 2.50 - 2.53 (1H, m), 2.64 - 2.71 (1H, m), 2.88 - 2.94 (3H, m), 3.40 (1H, m), 3.79 (3H, s), 3.94 (3H, s), 7.12 - 7.17 (3H, m), 7.73 (1H, s), 8.09 (1H, s), 8.19 (1H, d), 8.29 (1H, br s)	504	56
27 ¹	32		Methanol-d4 1.38 (3H, d), 2.59 - 2.78 (10H, m), 3.33 (3H, s), 3.38 (1H, m), 3.52 (2H, m), 3.75 (3H, s), 3.90 (3H, s), 7.09 - 7.13 (3H, m), 7.69 (1H, s), 8.05 (1H, s), 8.15 (1H, d), 8.25 (1H, br s)	534	45
28	32		Methanol-d4 1.08 (3H, t), 1.38 (3H, d), 2.43 - 2.51 (1H, q), 2.61 - 2.77 (8H, m), 3.37 (1H, m), 3.75 (3H, s), 3.90 (3H, s), 7.08 - 7.13 (3H, m), 7.68 (1H, s), 8.04 (1H, s), 8.14 (1H, d), 8.24 (1H, br s)	504	54
29	32		Methanol-d4 1.14 (3H, d), 1.19 (3H, d), 2.08 - 2.15 (1H, m), 2.29 - 2.41 (4H, m), 2.60 - 2.83 (5H, m), 3.75 (3H, s), 3.90 (3H, s), 4.02 (1H, m), 7.08 - 7.14 (3H, m), 7.68 (1H, s), 8.04 (1H, s), 8.14 (1H, d), 8.24 (1H, d)	504	43

Example	Starting Intermediate	Piperazine	NMR δ (300 MHz)	m/z [M+H] ⁺	Yield %
30⁶	32		Methanol-d4 1.12 (3H, d), 1.40 (3H, d), 2.27 - 2.48 (5H, m), 2.51- 2.57 (2H, m), 2.88 - 2.99 (3H, m), 3.40 (2H, m), 3.79 (3H, s), 3.94 (3H, s), 7.12 - 7.17 (3H, m), 7.73 (1H, s), 8.09 (1H, s), 8.19 (1H, d), 8.29 (1H, s) - two exchangeable protons not observed	504	42
31a⁷	36		Methanol-d4 1.15 (3H, d), 1.31 - 1.48 (6H, m), 2.16 - 2.24 (1H, m), 2.38 (6H, d), 2.52 - 2.56 (1H, m), 2.65- 2.69 (1H, m), 2.92 - 2.99 (3H, m), 3.33 (2H, m), 3.73 (3H, s), 4.86 (2H, q), 7.17 (2H, m), 7.73 (1H, s), 7.89 (1H, s), 8.16 (1H, s), 8.21 (1H, d) - two exchangeable protons not observed	532	22
31b⁷	36		Methanol-d4 1.14 (d, 3H), 1.30-1.50 (m, 6H), 2.25-2.38 (m, 4H), 2.41(s, 3H), 2.45- 2.60 (m, 2H), 2.85-3.10 (m, 3H), 3.35-3.46 (m, 2H), 3.73 (s, 3H), 4.25 (q, 2H), 7.05- 7.25 (m, 2H), 7.73 (s, 1H), 7.89 (s, 1H), 8.16 (s, 1H), 8.25 (d, 1H)	532	33

¹ The appropriate piperazine and 2-bromo-acetamide intermediate were combined in DMF according to the procedure of **Example 1**. Potassium carbonate was not used.

² The indicated piperazine and 2-bromo-acetamide intermediate were reacted under the conditions described by the procedure for **Example 1**. Chiral purification on a preparative chiral-HPLC using a Lux Cellulose-4 column (isocratic elution with 50% EtOH in isohexane modified with 0.2% IPA) to afford **Example 7a** (isolated as the second eluting peak, retention time = 9.45 min) and **Example 7b** (isolated as the first eluting peak, retention time = 7.54 min). Stereochemical assignment of the enantiomers was made based on the major product formation of the reaction and validated by biological activity against JAK1 in the Enzyme Inhibition Studies, as shown in **Example 66**.

³ The indicated piperazine and 2-bromo-acetamide intermediate were reacted under the conditions described by the procedure for **Example 1**. The crude product was purified by preparative chiral-HPLC on a Lux Cellulose-4 column, isocratic with 25% EtOH in isohexane (modified with 0.2% IPA) as eluent to afford **Example 10a** (first eluting peak, retention time = 5.02 min) and **Example 10b**, (second eluting peak, retention time = 6.68 min). Stereochemical assignment of the enantiomers was made based on the major product formation of the reaction and validated by biological activity against JAK1 in the Enzyme Inhibition Studies, as shown in **Example 66**.

⁴ The indicated piperazine and 2-bromo-acetamide intermediate were reacted under the conditions described by the procedure for **Example 1**. The crude product was purified by preparative chiral-HPLC (ADH column, isocratic with 50% EtOH in isohexane (modified with 0.2% IPA) as eluent) to give **Example**

11a (first eluting peak, retention time = 3.61 min) and **Example 11b** (second eluting peak, retention time = 4.60 min). Stereochemical assignment of the enantiomers was made based on the major product formation of the reaction and validated by biological activity against JAK1 in the Enzyme Inhibition Studies, as shown in **Example 66**.

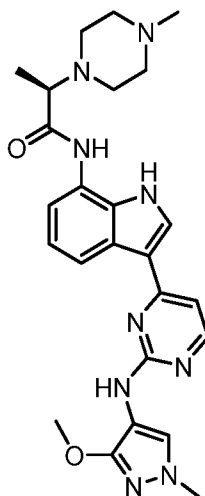
⁵The appropriate piperazine (2 equiv) and racemic 2-bromo-acetamide intermediate (1 equiv) were combined in 1,4-dioxane in the presence of silver oxide (4 equiv). Chiral-HPLC separation (ChiralPak IA column, isocratic with 50% EtOH in n-hexane (modified with 0.1% diethylamine) as eluent) was performed to afford **Example 14** (retention time = 3.99 min).

Stereochemical assignment of the enantiomers was made based on the biological activity against JAK1 in the Enzyme Inhibition Studies, as shown in **Example 66**.

⁶The e.r. was determined to be 94:6 by chiral-HPLC analysis (ChiralPak IA-3, 3 μ m, 0.46x5 cm, mobile phase: 50% ethanol in hexanes (modified with 0.2% IPA) at 1.0mL/min), retention time = 2.99 min.

⁷The indicated piperazine and racemic 2-bromo-acetamide intermediate were reacted in the presence of silver oxide (8.0 equivalents) in 1,4-dioxane at room temperature for 2 hours. The crude product was purified by preparative chiral-HPLC (Chiralpak IB column, isocratic with 50% Hexanes in EtOH (modified with 0.1% TEA) as eluent) to afford **Example 31a** (analytical chiral-HPLC: Chiralpak IA, 5 μ silica, 0.46x25 cm column, hexanes (0.1% TEA):EtOH (60:40) at 1.0 mL/min as the eluent, retention time = 8.18 min) isolated as the first eluting peak and **Example 31b** (analytical chiral-HPLC: Chiralpak IA, 5 μ silica, 0.46x25 cm column, hexanes (0.1% TEA):EtOH (60:40) at 1.0 mL/min as the eluent, retention time = 9.55 min) isolated as the second eluting peak. Stereochemical assignment of the enantiomers was made based on the biological activity against JAK1 in the Enzyme Inhibition Studies, as shown in **Example 66**.

Example 32: (2*R*)-N-(3-{2-[(3-Methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide



3-{2-[(3-Methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-amine (180 mg, 0.54 mmol, **Intermediate 23**), (*R*)-2-(4-methylpiperazin-1-yl)propanoic acid dihydrochloride (158 mg, 0.64 mmol, **Intermediate 37**) and HATU (408 mg, 1.1 mmol) in THF (5 mL) were stirred together to give an orange solution. Diisopropylethylamine (0.38 mL, 2.2 mmol) was added at 25°C. The resulting suspension was stirred at 25°C for 3 hours. The reaction mixture was diluted with ethyl acetate (100 mL), and washed with saturated aqueous Na₂CO₃ (50 mL),

water (50 mL) and brine (50 mL). The organic layer was dried, filtered and evaporated to afford crude product. The crude product was purified by preparative HPLC (XSelect CSH Prep C18 OBD column, 5 μ m, 19x150 mm), employing a gradient of 30-70% acetonitrile in 0.03% aqueous ammonia as eluents. Fractions containing the desired compound were evaporated to dryness to afford (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide (125 mg, 48%, **Example 32**) as a white solid; ¹H NMR δ (DMSO, 400MHz) 1.26 (3H, d), 2.16 (3H, s), 2.25 -2.45 (4H, m), 2.51 - 2.70 (4H, m), 3.71 (3H, s), 3.80 (3H, s), 7.05 (1H, t), 7.13 (1H, d), 7.38 (1H, d), 7.70 (1H, s), 8.16 - 8.31 (4H, m), 9.62 (1H, s), 11.35 (1H, s) - the α -proton to the amide is masked by the residual water peak; m/z (ES+), [M+H]⁺ = 490.

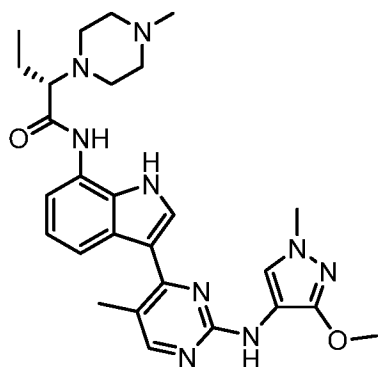
The procedure described above for Example 32 was repeated using the indicated Intermediates to give Examples 33-42 described in Table 12:

Table 12

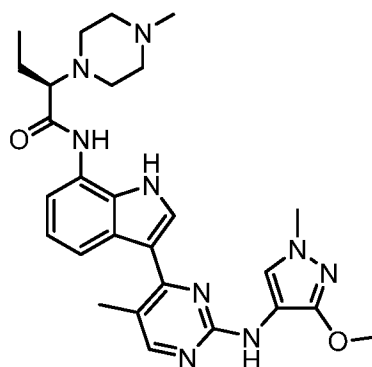
Example	Starting Intermediates	NMR δ (400 MHz)	m/z [M+H] ⁺	Yield %
33	25 and 38	DMSO-d6 with D2O 1.28 (3H, d), 2.27 (3H, s), 2.73 (3H, s), 2.85 - 3.34 (8H, m), 3.44 (1H, q), 3.63 (3H, s), 3.74 (3H, s), 7.04 (1H, t), 7.19 (1H, d), 7.55 (1H, s), 7.91 (1H, s), 8.08 (2H, s), 8.26 (1H, s) –two exchangeable protons not observed	504	13
34	25 and 37	DMSO-d6 1.26 (3H, d), 2.16 (3H, s), 2.33 (3H, s), 2.38 (4H, s), 2.57 - 2.62 (4H, m), 3.33 (1H, q), 3.67 (3H, s), 3.79 (3H, s), 7.00 (1H, t), 7.41 (1H, d), 7.66 (1H, s), 7.96 (2H, t), 8.14 (1H, s), 8.22 (1H, s), 9.65 (1H, s), 11.28 (1H, s)	504	72
35	30 and 37	Methanol-d4 1.34 (3H, t), 1.40 (3H, d), 2.32 (3H, s), 2.37 (3H, s), 2.50 - 2.80 (8H, m), 3.38 (1H, q), 3.69 (3H, s), 4.34 (2H, q), 7.05 - 7.20 (2H, m), 7.69 (1H, s), 7.85 (1H, s), 8.23 (1H, s), 8.17 (1H, d)-three exchangeable protons not observed	518	16
36	26 and 37	DMSO-d6 1.26 (3H, d), 2.27 (3H, s), 2.24 - 2.52 (4H, m), 2.53 - 2.70 (4H, m), 3.30 - 3.36 (1H, m), 3.69 (3H, s), 3.78 (3H, s), 7.02 (1H, s), 7.40 (1H, d), 7.65 (1H, s), 8.32 (1H, s), 8.48 (1H, s), 9.69 (1H, s), 11.42 (1H, s)	524	48

Example	Starting Intermediates	NMR δ (400 MHz)	m/z [M+H] ⁺	Yield %
37	27 and 37	DMSO-d ₆ 1.26 (3H, d), 2.17 (3H, s), 2.23 - 2.45 (4H, m), 2.46 - 2.71 (4H, m), 3.30 - 3.32 (1H, m), 3.68 (3H, s), 3.78 (3H, s), 7.01 (1H, s), 7.37 (1H, d), 7.64 (1H, s), 8.42 (1H, s), 8.45 - 8.56 (2H, m), 9.70 (1H, s), 11.36 (1H, s)	568	49
38	25 and 39	Chloroform-d 1.19 (3H, d), 1.35 (3H, d), 2.10 (1H, m), 2.26 (1H, m), 2.38 (6H, m), 2.69 (2H, t), 2.89 (3H, m), 3.72 (3H, s), 3.91 (1H, q), 4.00 (3H, s), 6.57 (1H, s), 6.80 (1H, d), 7.15 (1H, t), 7.68 (1H, d), 7.84 (1H, s), 8.06 - 8.36 (2H, m), 9.88 (1H, s), 11.15 (1H, s)	518	19
39	29 and 37	Methanol-d ₄ 1.34 (3H, t), 1.43 (3H, d), 2.35 (3H, s), 2.50 - 2.85 (8H, m), 3.41 (1H, q), 3.79 (3H, s), 4.24 (2H, q), 7.10 - 7.22 (2H, m), 7.68 (1H, s), 8.13 (1H, d), 8.16 (1H, d), 8.43 (1H, s)-three exchangeable protons not observed	522	25
40	31 and 37	Methanol-d ₄ 1.33 (3H, t), 1.42 (3H, d), 2.35 (3H, s), 2.63 - 2.71 (4H, m), 2.77 - 2.81 (4H, m), 3.42 (1H, q), 3.76 (3H, s), 4.26 (2H, q), 7.10 - 7.20 (2H, m), 7.70 (1H, s), 8.28 (2H, m), 8.48 (1H, m)-three exchangeable protons not observed	538	22
41	28 and 37	Chloroform-d 1.41 (3H, d), 2.29 (3H, s), 2.36 (3H, s), 2.42 (3H, s), 2.67 - 2.80 (8H, m), 3.38 (1H, q), 3.80 (3H, s), 6.42 (1H, s), 6.82 (1H, d), 7.12 (1H, t), 7.69 (1H, d), 7.88 (1H, s), 8.21 (2H, m), 9.74 (1H, s), 11.18 (1H, s)	488	36
42	28 and 38	DMSO-d ₆ 1.27 (3H, d), 2.12 (3H, s), 2.17 (3H, s), 2.35 (3H, s), 2.40 (4H, s), 2.57 - 2.63 (4H, m), 3.72 (3H, s), 7.03 (1H, t), 7.43 (1H, d), 7.81 (1H, s), 7.97 (1H, d), 8.19 (2H, m), 8.37 (1H, s), 9.68 (1H, s), 11.33 (1H, s)	488	4

Examples 43 and 44: (2*S*)-N-(3-{2-[(3-Methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)butanamide and (2*R*)-N-(3-{2-[(3-Methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)butanamide



Example 43



Example 44

Cesium fluoride (143 mg, 0.94 mmol) was added to a stirring solution of N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1-{2-[(trimethylsilyl)ethoxy]ethyl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)butanamide (203 mg, 0.31 mmol, **Intermediate 46**) in anhydrous DMSO (3 mL). The mixture was heated at 80°C under nitrogen for 4 hours then allowed to cool to ambient temperature. The reaction was diluted with ethyl acetate then water and the phases separated. The aqueous phase was extracted with ethyl acetate then the combined organic phase was washed with brine, dried over magnesium sulfate, and concentrated. The resultant residue was subject to silica gel chromatography using 5 - 20% methanol - DCM as eluent to afford a pale tan solid (101 mg). Chiral separation was performed by chiral-HPLC: Chiralpak ID, 4.6 x 50mm, 3, 50% hexane 50% 1:1 methanol - ethanol (modified with 0.1% diethylamine) to give (2*S*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)butanamide (17 mg, 11%, **Example 43**); Chiral HPLC: >99:1 e.r., retention time = 2.34 min; ¹H NMR δ (Dichloromethane-d₂, 400MHz) 1.09 (3H, t), 1.84 - 1.93 (2H, m), 2.29 (3H, s), 2.36 (3H, s), 2.51 (4H, br s), 2.67 (2H, br s), 2.75 (2H, m), 3.03 (1H, t), 3.67 (3H, s), 3.93 (3H, s), 6.49 (1H, s), 6.82 (1H, d), 7.12 (1H, t), 7.72 (1H, d), 7.79 (1H, s), 8.20 (1H, s), 8.24 (1H, d), 9.62 (1H, s), 11.06 (1H, br s); m/z (ES⁺) [M+H]⁺ = 518; followed by (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)butanamide (18 mg, 11%, **Example 44**); Chiral HPLC: ~99:5 e.r., retention time = 2.78 min; ¹H NMR δ (Dichloromethane-d₂, 400MHz) 1.09 (3H, t), 1.84 - 1.93 (2H, m), 2.29 (3H, s), 2.36 (3H, s), 2.51

(4H, br s), 2.67 (2H, br s), 2.75 (2H, m), 3.03 (1H, t), 3.67 (3H, s), 3.93 (3H, s), 6.49 (1H, s), 6.82 (1H, d), 7.12 (1H, t), 7.72 (1H, d), 7.79 (1H, s), 8.20 (1H, s), 8.24 (1H, d), 9.62 (1H, s), 11.06 (1H, br s); m/z (ES+), [M+H]⁺ = 518. Stereochemical assignment of the enantiomers was made based on the biological activity against JAK1 in the Enzyme Inhibition Studies, as shown in **Example 66**.

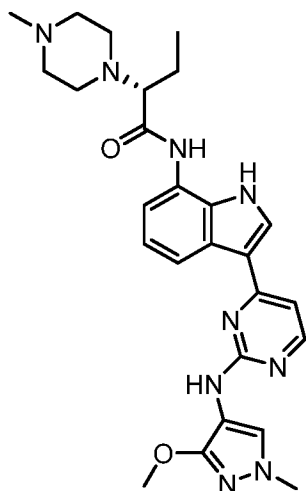
The procedure described above for Examples 43 and 44 was repeated using the indicated Starting Intermediates to give Examples 45 and 46 described in Table 13:

Table 13

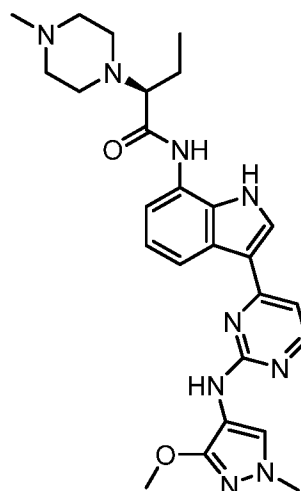
Example	Starting Intermediate	NMR δ (400 MHz)	m/z [M+H] ⁺	Yield %
45 ¹	47	Methanol-d ₄ 2.31 (3H, s), 2.37 (3H, s), 2.58 (4H, br s), 2.81 (2H, br s), 2.86 - 2.99 (2H, m), 3.41 (3H, s), 3.49 (1H, t), 3.70 (3H, s), 3.79 - 3.87 (1H, m), 3.88 - 3.96 (4H, m), 7.06 - 7.12 (1H, m), 7.13 - 7.18 (1H, m), 7.69 (1H, s), 7.86 (1H, s), 8.12 (1H, s), 8.19 (1H, d)-three exchangeable protons not observed	534	26
46 ¹	47	Methanol-d ₄ 2.31 (3H, s), 2.37 (3H, s), 2.58 (4H, br s), 2.81 (2H, br s), 2.86 - 2.99 (2H, m), 3.41 (3H, s), 3.49 (1H, t), 3.70 (3H, s), 3.79 - 3.87 (1H, m), 3.88 - 3.96 (4H, m), 7.06 - 7.12 (1H, m), 7.13 - 7.18 (1H, m), 7.69 (1H, s), 7.86 (1H, s), 8.12 (1H, s), 8.19 (1H, d)-three exchangeable protons not observed	534	28

¹Chiral separation was performed by preparative chiral-SFC (Chiralcel OD, 5 μ m, 4.6x100 mm) with 35% MeOH (modified with 0.1% dimethylethylamine) as eluent at 5 mL/min at 40 °C, to afford **Example 46** (first eluting peak, retention time = 2.54 min) and **Example 45** (second eluting peak, retention time = 3.10 min). Stereochemical assignment of the enantiomers was made based on the biological activity against JAK1 in the Enzyme Inhibition Studies, as shown in **Example 66**.

Examples 47 and 48: (2*R*)-N-(3-{2-[(3-Methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)butanamide and (2*S*)-N-(3-{2-[(3-Methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)butanamide



Example 47



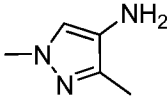
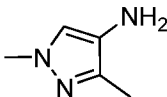
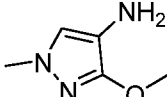
Example 48

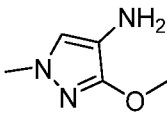
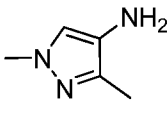
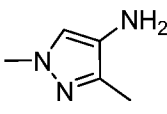
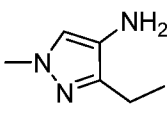
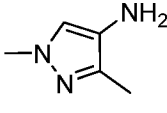
N-[3-(2-Chloropyrimidin-4-yl)-1*H*-indol-7-yl]-2-(4-methylpiperazin-1-yl)butanamide (0.22 g, 0.53 mmol, **Intermediate 51**), 3-methoxy-1-methyl-1*H*-pyrazol-4-amine dihydrochloride (0.16 g, 0.8 mmol) and 4-methylbenzenesulfonic acid monohydrate (0.2 g, 1.1 mmol) were dissolved in isopropanol (6 mL) and sealed into a microwave tube. The reaction was heated at 120°C for 2 hours in the microwave reactor and cooled to room temperature. The crude product was purified by preparative HPLC (XBridge Prep C18 OBD column, 5μ silica, 19x150 mm), using decreasingly polar mixtures of water (containing 0.2% formic acid) and acetonitrile as eluents. Fractions containing the desired compound were evaporated to dryness to afford racemic N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)butanamide (90 mg, 34%) as a white solid; *m/z* (ES⁺), [M+H]⁺ = 504. The product was purified by preparative chiral-HPLC on an IC-3 column, isocratically with 30% ethanol in isohexane (modified with 0.2% isopropanol) as eluent. The fractions containing the desired compound were evaporated to dryness to afford firstly (2*S*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)butanamide (32 mg, 35%, **Example 48**) as a white solid; ¹H NMR δ (Methanol-d₄, 300 MHz) 1.08 (3H, t), 1.89 (2H, dt), 2.33 (3H, s), 2.59 (4H, br s), 2.83 (4H, br s), 3.21 (1H, dd), 3.80 (3H, s), 3.94 (3H, s), 7.10 - 7.27 (3H, m), 7.74 (1H, s), 8.11 (1H, s), 8.21 (1H, d), 8.30 (1H, s)-three exchangeable protons not observed; *m/z* (ES⁺), [M+H]⁺ = 504; chiral HPLC: 100% ee, retention time = 4.48 min; followed by (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-

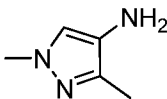
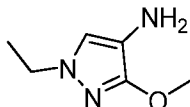
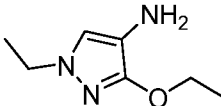
yl)-2-(4-methylpiperazin-1-yl)butanamide (32 mg, 35%, **Example 47**) as a white solid; ^1H NMR δ (Methanol- d_4 , 300 MHz) 1.08 (3H, t), 1.89 (2H, dt), 2.33 (3H, s), 2.59 (4H, br s), 2.83 (4H, br s), 3.21 (1H, dd), 3.80 (3H, s), 3.94 (3H, s), 7.10 - 7.27 (3H, m), 7.74 (1H, s), 8.11 (1H, s), 8.21 (1H, d), 8.30 (1H, s)-three exchangeable protons not observed; m/z (ES $^+$), $[\text{M}+\text{H}]^+ = 504$; Chiral
 5 HPLC: 100% ee, retention time = 5.69 min. Stereochemical assignment of the enantiomers was made based on the biological activity against JAK1 in the Enzyme Inhibition Studies, as shown in **Example 66**.

The procedure described above for Examples 47 and 48 was repeated using the indicated
 10 Starting Intermediates and aminopyrazole to give Examples 49-59 described in Table 14:

Table 14

Example	Starting Intermediate	Aminopyrazole	NMR δ (400 MHz)	m/z $[\text{M}+\text{H}]^+$	Yield %
49¹	52		Methanol- d_4 1.07 (3H, t), 1.89 (2H, m), 2.21 (3H, s), 2.33 (3H, d), 2.59 (4H, s), 2.83 (4H, d), 3.20 (1H, dd), 3.88 (3H, s), 7.12 (1H, t), 7.21 (1H, d), 7.74 (1H, s), 8.16 (2H, d), 8.37 (1H, s)-three exchangeable protons not observed	506	38
50¹	52		Methanol- d_4 1.07 (3H, t), 1.89 (2H, m), 2.21 (3H, s), 2.33 (3H, d), 2.59 (4H, s), 2.83 (4H, d), 3.20 (1H, dd), 3.88 (3H, s), 7.12 (1H, t), 7.21 (1H, d), 7.74 (1H, s), 8.16 (2H, d), 8.37 (1H, s)-three exchangeable protons not observed	506	38
51^{2,3}	52		DMSO- d_6 0.92 (3H, t), 1.65 - 1.82 (2H, m), 2.14 (3H, s), 2.34 (4H, m), 2.65 (4H, m), 3.16 (1H, t), 3.71 (3H, s), 3.78 (3H, s), 7.05 (1H, t), 7.54 (1H, d), 7.66 (1H, s), 8.16 (1H, t), 8.23 - 8.35 (3H, m), 9.75 (1H, s), 11.36 (1H, s)	522	48

Example	Starting Intermediate	Aminopyrazole	NMR δ (400 MHz)	m/z [M+H] ⁺	Yield %
52 ^{2,3}	52		DMSO-d ₆ 0.92 (3H, t), 1.65 - 1.82 (2H, m), 2.14 (3H, s), 2.34 (4H, m), 2.65 (4H, m), 3.16 (1H, t), 3.71 (3H, s), 3.78 (3H, s), 7.05 (1H, t), 7.54 (1H, d), 7.66 (1H, s), 8.16 (1H, t), 8.23 - 8.35 (3H, m), 9.75 (1H, s), 11.36 (1H, s)	522	47
53 ⁴	53		Methanol-d ₄ 2.21 (3H, s), 2.34 (3H, s), 2.61 (4H, m), 2.83 (2H, s), 2.93 (2H, s), 3.43 (3H, s), 3.51 (1H, t), 3.80 - 3.92 (4H, m), 3.95 (1H, m), 7.08 - 7.21 (2H, m), 7.74 (1H, s), 8.15 (2H, t), 8.38 (1H, s)-three exchangeable protons not observed	522	26
54 ⁴	53		Methanol-d ₄ 2.21 (3H, s), 2.34 (3H, s), 2.61 (4H, m), 2.83 (2H, s), 2.93 (2H, s), 3.43 (3H, s), 3.51 (1H, t), 3.80 - 3.92 (4H, m), 3.95 (1H, m), 7.08 - 7.21 (2H, m), 7.74 (1H, s), 8.15 (2H, t), 8.38 (1H, s)-three exchangeable protons not observed	522	26
55 ^{3,5}	54		Methanol-d ₄ 1.31 (3H, t), 1.42 (3H, d), 2.26 (3H, s), 2.35 (3H, s), 2.52 - 2.70 (6H, m), 2.73 - 2.88 (4H, m), 3.40 - 3.42 (1H, m), 3.85 (3H, s), 7.06 - 7.18 (2H, m), 7.77 (1H, s), 7.91 (1H, s), 8.14 - 8.19 (2H, m)-three exchangeable protons not observed	502	19
56 ⁵	55		Methanol-d ₄ 1.43 (3H, d), 2.21 (3H, s), 2.34 (3H, s), 2.51 - 2.91 (8H, m), 3.41 (1H, q), 3.88 (3H, s), 7.07 - 7.21 (2H, m), 7.74 (1H, s), 8.15 (2H, dd), 8.37 (1H, s)-three exchangeable protons not observed	492	17

Example	Starting Intermediate	Aminopyrazole	NMR δ (400 MHz)	m/z [M+H] ⁺	Yield %
57 ⁵	59		DMSO-d ₆ 1.27 (3H, d), 2.14 (3H, s), 2.18 (3H, s), 2.26 - 2.75 (8H, m), 3.25 - 3.41 (1H, m), 3.77 (3H, s), 7.07 (1H, t), 7.14 (1H, d), 7.40 (1H, d), 7.83 (1H, s), 8.08 - 8.36 (3H, m), 8.47 (1H, s), 9.63 (1H, s), 11.36 (1H, br s)	474	60
58 ⁵	54		Dichloromethane-d ₂ 1.17 - 1.44 (6H, m), 2.36 (3H, s), 2.39 (3H, s), 2.57 - 2.90 (8H, m), 3.37 (1H, q), 3.80 - 4.08 (5H, m), 6.65 (1H, s), 6.86 (1H, d), 7.18 (1H, t), 7.75 (1H, d), 7.87 (1H, s), 8.24 (1H, s), 8.32 (1H, d), 9.62 (1H, s), 11.23 (1H, s)	518	50
59 ⁵	54		Chloroform-d 1.36 - 1.50 (9H, m), 2.36 (3H, s), 2.39 (3H, s), 2.55 - 2.90 (8H, m), 3.36 (1H, m), 3.95 (2H, q), 4.32 (2H, q), 6.73 (1H, s), 6.84 (1H, d), 7.13 (1H, t), 7.67 (1H, d), 7.90 (1H, s), 8.13 - 8.40 (2H, m), 9.78 (1H, s), 11.17 (1H, br s)	532	82

¹Chiral separation was performed by chiral-HPLC (Chiralcel OD-H column, isocratic with 10% IPA in hexanes) to give **Example 50** (analytical chiral-HPLC: Chiralpak OD-H 5 μ silica, 0.46x10 cm column, hexanes (modified with 0.2% IPA):EtOH (90:10) at 1.0 mL/min as the eluent, retention time = 9.02 min) isolated as the first eluting peak and **Example 49** (analytical chiral-HPLC: Chiralpak OD-H 5 μ silica, 0.46x10 cm column, hexanes (modified with 0.2% IPA):EtOH (90:10) at 1.0 mL/min as the eluent, retention time = 11.35 min) isolated as the second eluting peak. Stereochemical assignment of the enantiomers was made based on the biological activity against JAK1 in the Enzyme Inhibition Studies, as shown in **Example 66**.

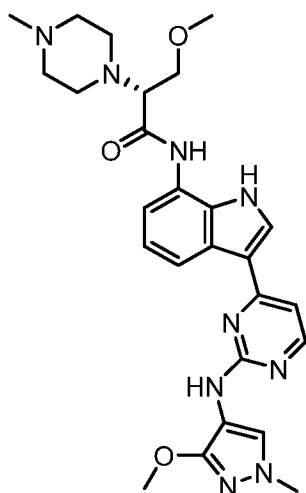
²Chiral separation was performed by preparative chiral-HPLC (Chiralcel OD-H, 20x250 mm column, isocratic with 10% ethanol in hexanes (modified with 0.2% diethylamine) at 20 mL/min as an eluent) to afford **Example 52** (first eluting peak, retention time = 15.87 min) and **Example 51** (second eluting peak, retention time = 21.29 min). Stereochemical assignment of the enantiomers was made based on the biological activity against JAK1 in the Enzyme Inhibition Studies, as shown in **Example 66**.

³¹H NMR analysis was performed using a Bruker Avance 300 (300 MHz) spectrometer.

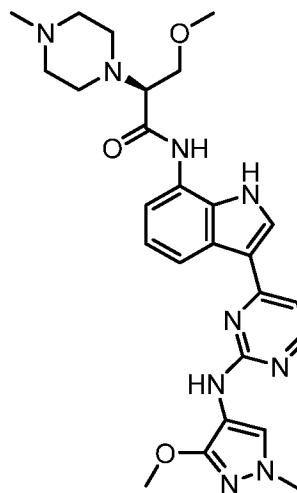
⁴Chiral separation was performed by preparative chiral-HPLC (Chiralcel IC column, isocratic with 40% ethanol in hexanes as an eluent) to afford **Example 54** (analytical chiral-HPLC: Lux Cellulose-4 3 μ silica, 0.46x5 cm column, hexanes (modified with 0.1% TEA):EtOH (60:40) at 1.0 mL/min as the eluent, retention time = 2.69 min) isolated as the first eluting peak and **Example 53** (analytical chiral-HPLC: Lux Cellulose-4 3 μ silica, 0.46x5 cm column, hexanes (modified with 0.1% TEA):EtOH (60:40) at 1.0 mL/min as the eluent, retention time = 3.62 min) isolated as the second eluting peak. Stereochemical assignment of the enantiomers was made based on the biological activity against JAK1 in the Enzyme Inhibition Studies, as shown in **Example 66**.

⁵Enantiopure starting material utilized - final product not subject to chiral-HPLC purification.

Examples 60 and 61: (2*R*)-3-Methoxy-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide and (2*S*)-3-Methoxy-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide



Example 60



Example 61

Diisopropylethylamine (1.25 mL, 7.2 mmol) was added to 3-(2-((3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino)pyrimidin-4-yl)-1*H*-indol-7-amine (0.4 g, 1.2 mmol, **Intermediate 23**), lithium 3-methoxy-2-(4-methylpiperazin-1-yl)propanoate (0.72 g, 3.6 mmol, **Intermediate 45**) and HATU (1.4 g, 3.6 mmol) in DMF (18 mL) at 25°C under nitrogen. The resulting mixture was stirred at 25°C for 1 hour. The crude product was purified by preparative HPLC (XBridge Prep C18 OBD column, 5μ silica, 19 mm diameter, 150 mm length), using decreasingly polar mixtures of water (containing 0.2% ammonia) and acetonitrile as eluents. Fractions containing the desired compound were evaporated to dryness to afford racemic 3-methoxy-N-(3-(2-((3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino)pyrimidin-4-yl)-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide (0.15 g, 24%) as a white solid; *m/z* (ES⁺), [M+H]⁺ = 520. The crude product was purified by preparative chiral-HPLC on a Lux Cellulose-4 column, eluting isocratically with 50% ethanol in isohexane (modified with 0.1% triethylamine) as eluent. The fractions containing the desired compound were evaporated to dryness to afford firstly (2*S*)-3-methoxy-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide (53 mg, 35%, **Example 61**) as a white solid; ¹H NMR δ (Methanol-*d*₄, 300 MHz) 2.33 (3H, s), 2.60 (4H, s), 2.78 - 2.99 (4H, m), 3.43 (3H, s), 3.51 (1H, t), 3.76 - 4.00 (8H, m), 7.08 - 7.22 (3H, m), 7.72 (1H, s), 8.09 (1H, s), 8.19 (1H, d), 8.30 (1H, s)-three exchangeable protons not observed; *m/z* (ES⁺), [M+H]⁺ = 520; Chiral HPLC: 100% ee, *R*_t = 4.072 min;

followed by (2*R*)-3-methoxy-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide (57 mg, 36%, **Example 60**) as a white solid; ¹H NMR δ (Methanol-d₄, 300 MHz) 2.30 (3H, s), 2.57 (4H, s), 2.87 - 2.80 (4H, m), 3.39 (3H, s), 3.47 (1H, t), 3.75 - 3.98 (8H, m), 7.04 - 7.18 (3H, m), 7.68 (1H, s), 8.05 (1H, s), 8.15 (1H, d), 8.26 (1H, s)-three exchangeable protons not observed; m/z (ES⁺), [M+H]⁺ = 520; Chiral HPLC: 99.2% ee, R_t = 5.376 min. Stereochemical assignment of the enantiomers was made based on the biological activity against JAK1 in the Enzyme Inhibition Studies, as shown in **Example 66**.

The procedure described above for Examples 60 and 61 was repeated using the indicated Starting Intermediates to give the compounds described in Table 15:

Table 15

Example	Starting Intermediates	NMR δ (300 MHz)	m/z [M+H] ⁺	Yield %
62 ¹	30 and 43	DMSO-d ₆ 0.86 (3H, t), 1.20 (3H, t), 1.38 – 1.88 (2H, m), 2.07 (3H, s), 2.26 (7H, m), 2.45 – 2.74 (4H, m), 3.12 (1H, t), 3.59 (3H, s), 4.09 (2H, q), 6.93 (1H, t), 7.47 (1H, d), 7.58 (1H, s), 7.77 (1H, s), 7.92 (1H, s), 8.08 (1H, s), 8.18 (1H, br s), 9.58 (1H, s), 11.13 (1H, s)	532	24
63 ¹	30 and 43	DMSO-d ₆ 0.86 (3H, t), 1.20 (3H, t), 1.45 – 1.76 (2H, m), 2.07 (3H, s), 2.26 (7H, s), 2.54 – 2.73 (4H, m), 3.10 (1H, t), 3.59 (3H, s), 4.09 (2H, q), 6.94 (1H, s), 7.45 (1H, d), 7.58 (1H, s), 7.78 (1H, s), 7.91 (1H, s), 8.08 (1H, s), 8.18 (1H, br s), 9.55 (1H, s), 11.10 (1H, s)	532	16
64 ²	30 and 45	DMSO-d ₆ 1.19 (3H, t), 2.10 (3H, s), 2.33 – 2.38 (7H, m), 2.56 – 2.72 (4H, m), 3.34 (3H, s), 3.64 (1H, t), 3.69 – 3.73 (5H, m), 4.09 (2H, q), 6.94 (1H, t), 7.39 (1H, d), 7.58 (1H, s), 7.80 (1H, s), 7.91 (1H, d), 8.08 (1H, s), 8.18 (1H, br s), 9.69 (1H, s), 11.08 (1H, br s)	548	32
65 ²	30 and 45	DMSO-d ₆ 1.27 (3H, t), 2.15 (3H, s), 2.34 – 2.38 (7H, m), 2.60 – 2.76 (4H, m), 3.33 (3H, s), 3.69 (1H, t), 3.68 – 3.73 (5H, m), 4.16 (2H, q), 6.96 (1H, t), 7.45 (1H, d), 7.60 (1H, s), 7.83 (1H, s), 7.90 (1H, s), 8.10 (1H, s), 8.18 (1H, br s), 9.71 (1H, br s), 11.10 (1H, br s)	548	30

¹Chiral separation was performed by preparative chiral-SFC (Chiralcel OD, 5μm, 21x250 mm) eluting isocratically with 20% MeOH (modified with 0.1% dimethylethylamine) at 75 mL/min at 40 °C, to afford **Example 63** (first eluting peak, retention time = 7.89 min) and **Example 62** (second eluting peak,

retention time = 8.81 min). Stereochemical assignment of the enantiomers was made based on the biological activity against JAK1 in the Enzyme Inhibition Studies, as shown in **Example 66**.

²Chiral separation was performed by preparative chiral-SFC (Chiralcel OD, 5 μ m, 21x250 mm) eluting isocratically with 25% MeOH (modified with 0.1% dimethylethylamine) at 75 mL/min at 40 °C, to afford

Example 65 (first eluting peak, retention time = 4.84 min) and **Example 64** (second eluting peak, retention time = 5.95 min). Stereochemical assignment of the enantiomers was made based on the biological activity against JAK1 in the Enzyme Inhibition Studies, as shown in **Example 66**.

Example 66: Enzyme Inhibition Studies

Enzyme inhibition studies were performed using recombinant JAK1 (amino acids 866-1154, Life Technologies, #PV4774, Carlsbad, CA), JAK2 (amino acids 831-1132), or JAK3 (amino acids 781-1124) under buffer conditions of 50 mM HEPES pH 7.3, 1 mM DTT, 0.01% Tween® 20, 50 μ g/mL BSA, and 10 mM MgCl₂. JAK enzyme was expressed as an N-terminal GST fusion in insect cells and purified by glutathione-affinity and size-exclusion chromatographies. Enzymes were assayed both at their respective ATP K_m (JAK1: 55 μ M, JAK2: 15 μ M, JAK3: 3 μ M) and the approximated high end of physiological ATP concentration of 5 mM, in the presence of inhibitor dosed at 30, 3, 0.3, 0.03, 0.003 and 0 μ M final test concentrations. For JAK1, 6 nM of enzyme (for K_m ATP assay) or 4 nM enzyme (for high ATP assay) was incubated with 1.5 μ M peptide substrate (FITC-C6-KKHTDDGYMPMSPGVA-NH₂ (SEQ ID NO:1), Intonation, Boston, MA). For JAK2, 0.8 nM of enzyme (for K_m ATP assay) or 0.3 nM enzyme (for high ATP assay) was incubated with 1.5 μ M peptide substrate (5FAM-GEEPLYWSFPAKKK-NH₂ (SEQ ID NO:2), Intonation, Boston, MA). For JAK3, 0.2 nM of enzyme (for K_m ATP assay) or 0.1 nM enzyme (for high ATP assay) was incubated with 1.5 μ M peptide substrate (5FAM-GEEPLYWSFPAKKK-NH₂ (SEQ ID NO:2), Intonation, Boston, MA). Phosphorylated and unphosphorylated peptides were separated and quantified by a Caliper LC3000 system (Caliper Life Sciences, MA) for calculating percent inhibition. The results of this assay are shown in Table 16 and indicate that the compounds of Formula (I), (Ia), (Ib) and Table 1 exhibit preferential inhibition of JAK1 over JAK2 (in many cases demonstrating over 100 times selectivity for inhibition of JAK1 vs. JAK2).

Example 67: Cellular pSTAT3 assay

NCI-H1975 cells were plated onto Costar #3701 96 or 384 well tissue-culture treated plates at 5,000 cells/well in 30 μ L medium (RPMI, 10% FBS, supplemented with L-glutamine) and incubated overnight at 37 °C in 5% CO₂. Phospho STAT3 signal was quantitated utilizing Cell Signaling Technology #7146B Pathscan 97phosphor STAT3 antibody pair kit, following manufacturer's instructions.

Cells were dosed with compound and incubated at 37 °C in 5% CO₂ for 2 hours, after which medium and compound were aspirated and cells lysed with 35uL cold 1x Cell Signaling Lysis buffer and chilled at 4°C for 1-2 hours. Lysate was incubated on STAT3 capture plates at 4°C overnight, washed 3x with Tris-Buffered Saline with 0.05% Tween® 20 (TBST), then 98 phosphor STAT3 detection antibody was applied for 2 hours. Following washing with TBST (3x), HRP-secondary antibody was applied for 2 hours. After additional washing, signal was detected using TMB and Stop solution and read at 450 nm using Tecan Infinite M100.

IC₅₀ values (the concentration that causes 50% inhibition) were calculated by plotting percent inhibition of the phosphor-signal relative to untreated sample (maximum signal) and positive control treated sample (maximum inhibition/minimum signal), using Xlfit4 version 4.2.2 for Microsoft Excel. The results of this assay, shown in Table 16, demonstrate good correlation between cellular inhibition of STAT3 phosphorylation in NCI-H1975 cells and JAK1 enzyme inhibition.

Table 16

Example	JAK1 (IC ₅₀ , μM)	JAK2 (IC ₅₀ , μM)	JAK3 (IC ₅₀ , μM)	NCI-H1975 pSTAT3 (IC ₅₀ , μM)
1	0.847	>30	>30	0.643
2	0.085	12.8	>30	0.227
3	0.191	26.8	>30	0.302
4	0.043	6.57	>30	0.0986
5	0.024	10.4	>30	0.0872
6	0.030	9.62	>30	0.111
7a	0.359	>30	>30	0.509
7b	3.60	>30	>30	>3
8	0.110	23.6	>30	0.191
9	0.426	>30	>30	0.594
10a	0.846	>30	>30	0.531
10b	7.60	>30	>30	>3
11a	0.040	>30	>30	0.209
11b	1.85	>30	>30	2.28
12	0.068	22.6	>30	0.246
13	0.166	>30	>30	0.222
14	0.009	6.39	>30	0.0703
15	0.007	2.53	>30	0.131
16	0.019	6.24	>30	0.105
17	0.107	18.1	>30	0.229
18	0.057	17.3	>30	0.175
19	0.296	>30	>30	0.470
20	0.051	17.2	>30	0.184
21	0.184	>30	>30	0.703
22	0.490	>30	>30	0.550
23	0.029	25.7	>30	0.410
24	0.113	>30	>30	0.370

25	0.019	6.62	>30	0.115
26	0.069	16.7	>30	0.256
27	0.575	>30	>30	0.609
28	0.106	11.8	>30	0.100
29	0.097	20.4	>30	0.227
30	0.304	>30	>30	0.359
31a	0.015	19.8	>30	0.153
31b	0.454	>30	>30	0.330
32	0.073	>14.7	>30	0.161
33	0.272	>30	>30	0.308
34	0.010	5.37	>30	0.128
35	0.020	17.0	>30	0.153
36	0.007	1.76	19.1	0.0579
37	0.008	1.58	15.4	0.0942
38	0.013	6.48	>30	0.108
39	0.055	25.5	>30	0.191
40	0.024	8.93	>30	0.160
41	<0.005	2.58	>30	0.0923
42	0.106	21.1	>30	0.172
43	0.079	21.1	>30	0.176
44	<0.004	1.95	>30	0.0577
45	<0.003	0.790	>30	0.0603
46	0.986	>30	>30	1.28
47	0.013	6.95	>30	0.136
48	1.21	>30	>30	0.863
49	<0.004	1.77	>30	0.0751
50	0.265	24.1	>30	0.999
51	<0.003	3.19	>30	0.111
52	0.296	>30	>30	0.548
53	<0.003	0.322	>30	0.0566
54	1.67	>30	>30	>3
55	0.027	4.22	>30	0.116
56	0.028	5.15	>30	0.126
57	0.069	9.26	>30	0.155
58	0.017	8.85	>30	0.106
59	0.069	>30	>30	0.245
60	0.004	1.53	>30	0.0973
61	2.90	>30	>30	1.27
62	<0.006	5.01	>30	0.104
63	0.192	>30	>30	0.433
64	<0.005	2.04	>30	0.112
65	2.53	>30	>30	1.03

Example 68: Solid Forms of (2*R*)-*N*-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide

Methods

5 X-Ray Powder Diffraction (XRPD) Analysis

XRPD analysis was performed using a Bruker D4 (or D8) diffractometer, which is commercially available from Bruker AXS IncTM (Madison, Wisconsin). The XRPD spectra were obtained by mounting a sample (approximately 20 mg) of the material for analysis on a single silicon crystal wafer mount (e.g., a Bruker silicon zero background X-ray diffraction sample holder) and spreading out the sample into a thin layer with the aid of a microscope slide. The sample was spun at 30 revolutions per minute (to improve counting statistics) and irradiated with X-rays generated by a copper long-fine focus tube operated at 40 kV and 40 mA with a wavelength of 1.5406 angstroms (i.e., about 1.54 angstroms). The sample was exposed for 1 second per 0.02 degree 2-theta increment (continuous scan mode) over the range 5 degrees (or 10 2 degrees) to 40 degrees 2-theta in theta-theta mode. The running time was ~17 min for D4 and ~15 min for D8.

XRPD 2θ values may vary with a reasonable range, e.g., in the range ± 0.2° and that XRPD intensities may vary when measured for essentially the same crystalline form for a variety of reasons including, for example, preferred orientation. Principles of XRPD are described in 20 publications, such as, for example, Giacobozzo, C. et al. (1995), Fundamentals of Crystallography, Oxford University Press; Jenkins, R. and Snyder, R. L. (1996), Introduction to X-Ray Powder Diffractometry, John Wiley & Sons, New York; and Klug, H. P. & Alexander, L. E. (1974), X-ray Diffraction Procedures, John Wiley and Sons, New York.

25 DSC Analysis

DSC analysis was performed on samples prepared according to standard methods using a Q SERIESTM Q1000 DSC calorimeter available from TA INSTRUMENTS® (New Castle, Delaware). A sample (approximately 2 mg) was weighed into an aluminum sample pan and transferred to the DSC. The instrument was purged with nitrogen at 50 mL/min and data 30 collected between 22 °C and 300 °C, using a dynamic heating rate of 10 °C/minute. Thermal data was analyzed using standard software, e.g., Universal v.4.5A from TA INSTRUMENTS®.

Thermogravimetry Analysis (TGA)

TGA was performed on samples prepared according to standard methods using a Q SERIES™ Q5000 thermogravimetry analyzer available from TA Instruments INSTRUMENTS® (New Castle, Delaware). A sample (approximately 5 mg) was placed into an aluminum sample pan and transferred to the TGA furnace. The instrument was purged with nitrogen at 50 mL/min and data collected between 25 °C and 300 °C, using a dynamic heating rate of 10 °C/minute. Thermal data was analyzed using standard software, e.g., Universal v.4.5A from TA INSTRUMENTS®.

Example 68A: Form A (2R)-N-(3-{2-[(3-methoxy-1-methyl-1H-pyrazol-4-yl)amino]pyrimidin-4-yl}-1H-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide

Method 1: 50 mg of off-white amorphous (2R)-N-(3-{2-[(3-methoxy-1-methyl-1H-pyrazol-4-yl)amino]pyrimidin-4-yl}-1H-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide was dissolved in 0.4 ml of TBME in a 4 mL vial. The solid precipitated out from the solution after 30 minutes. The slurry was stirred under ambient conditions overnight. The resulting white solid material was identified as Form A by XRPD analysis.

Method 2: 500 mg of amorphous (2R)-N-(3-{2-[(3-methoxy-1-methyl-1H-pyrazol-4-yl)amino]pyrimidin-4-yl}-1H-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide and approximately 50 mg of crystalline seeds obtained from Method 1 were mixed in a 20 mL vial. To the mixture, 5 mL of TBME was added to form a slurry. The slurry was stirred under ambient conditions overnight and a homogenous slurry formed. The slurry was filtered, and the resulting solid was washed with TBME and dried in air. 498 mg of a white crystalline solid was obtained and identified as Form A by XRPD analysis.

Form A (Method 2) was analyzed by XRPD and the results are tabulated below (Table 17) and shown in Figure 1.

Table 17

Angle (2θ±0.2°)	Intensity (%)
21.6	100.0
6.4	74.9
16.4	56.5
8.7	54.2
20.4	45.8
7.9	42.8
22.2	36.2
18.8	34.6

16.8	29.6
9.3	29.2
20.1	26.3
16.1	24.5
19.8	23.2
26.5	22.1
13.8	20.9
19.2	20.8
13.2	19.5
12.0	17.2
23.8	16.9
21.1	15.4
25.4	14.3
28.5	14.3
12.9	13.7
8.3	12.7
10.9	11.7
25.9	11.7
24.4	11.5
22.7	11.4
32.0	10.9
15.7	10.7
14.8	10.1
25.0	9.2
27.0	8.7
9.9	8.2
11.6	8.2
29.2	7.7
29.9	7.6
17.4	7.4
17.8	7.4
27.8	7.2
30.2	6.8
32.6	6.2
31.1	5.5

Form A (Method 2) was analyzed by thermal techniques. DSC analysis indicated that Form A has an endotherm event of desolvation with an onset at 110 °C and a peak at 113 °C.

TGA indicated that Form A exhibits a mass loss of about 7.8 % upon heating from about 25 °C to about 150 °C. A representative DSC/TGA thermogram of Form A is shown in Figure 2.

Example 68B: Form B (2R)-N-(3-{2-[(3-methoxy-1-methyl-1H-pyrazol-4-yl)amino]pyrimidin-4-yl}-1H-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide

Approximately 100 mg of light yellow amorphous (2R)-N-(3-{2-[(3-methoxy-1-methyl-1H-pyrazol-4-yl)amino]pyrimidin-4-yl}-1H-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide was suspended in 0.5 mL of toluene with Form B seeds. The slurry was stirred at room temperature overnight. The slurry was evaporated and dried in the ambient condition. A white crystalline solid was obtained and identified as Form B by XRPD.

Form B was analyzed by XRPD and the results are tabulated below (Table 18) and shown in Figure 3.

Table 18

Angle (2θ±0.2°)	Intensity (%)
21.6	100.0
6.3	93.7
8.8	72.2
19.0	53.5
8.0	53.1
16.3	47.3
9.5	43.4
22.4	41.2
20.4	34.8
19.7	30.2
13.9	27.1
26.4	26.7
28.6	25.2
16.7	24.7
15.9	19.5
10.8	19.0
11.9	18.5
13.4	18.4
23.9	17.6
31.8	15.8
24.4	14.7
25.5	14.6
25.1	14.3
28.1	13.5

32.7	12.9
14.6	12.6
29.8	11.9
37.5	11.7
29.3	10.6
15.0	10.4

Single crystals of Form B were obtained from slow evaporation of a toluene solution.

Single crystal structure analysis confirmed that Form B is a hemi-toluene solvate form.

Crystallographic data: Space group triclinic *P*1, unit cell dimensions: *a* = 14.1919(8) Å, *b* =

5 14.2964(8) Å, *c* = 14.7632(8) Å, α = 82.283(1)°, β = 77.596(1)°, γ = 85.567(1)°, *V* = 2895.3(3) Å³.

Form B was analyzed by thermal techniques. DSC analysis indicated that Form B has an endotherm event of desolvation with an onset at 112 °C and a peak at 117 °C. TGA indicated that Form B exhibits a mass loss of about 10.0 % upon heating from about 25 °C to about 200 °C. A representative DSC/TGA thermogram of Form B is shown in Figure 4.

Example 68C: Form C (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide

Approximately 100 mg of amorphous (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide was dissolved in 1 mL of isopropyl acetate to yield a clear solution. The solution was placed in the freezer overnight and a solid precipitated out. The slurry was stirred at room temperature for 4 hours to obtain a white solid in the slurry. The slurry was evaporated and dried under ambient condition. A white crystalline solid was obtained and was identified as Form C by XRPD.

Form C was analyzed by XRPD and the results are tabulated below (Table 19) and shown in Figure 5.

Table 19

Angle (2θ±0.2°)	Intensity (%)
8.7	100.0
21.5	68.0
6.3	58.6
16.3	53.8
22.2	31.5
18.9	30.3
7.9	28.3

20.4	26.3
13.8	22.0
19.7	21.5
26.4	19.3
9.3	18.1
16.6	12.6
23.6	12.1
19.9	11.8
11.9	11.3
13.2	11.1
28.4	11.1
25.3	10.7
16.0	9.3
18.4	8.9
31.8	8.1
26.0	8.0
24.2	7.9
10.7	6.8
17.5	6.8
20.9	6.6
12.8	6.3
29.8	6.0
15.7	5.8
26.9	5.8
14.7	5.5

Form C was analyzed by thermal techniques. DSC analysis indicated that Form C has an endotherm event of desolvation with an onset at 112 °C and a peak at 114 °C. TGA indicated that Form C exhibits a mass loss of about 9.2 % upon heating from about 25 °C to about 175 °C. A representative DSC/TGA thermogram of Form C is shown in Figure 6.

Example 68D: Form D (2R)-N-(3-{2-[(3-methoxy-1-methyl-1H-pyrazol-4-yl)amino]pyrimidin-4-yl}-1H-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide

Method 1: Approximately 100 mg of light yellow amorphous (2R)-N-(3-{2-[(3-methoxy-1-methyl-1H-pyrazol-4-yl)amino]pyrimidin-4-yl}-1H-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide was dissolved in 1 ml of EtOAC to get a clear solution. The solution was placed in the freezer overnight and solid was precipitated out. The slurry was stirred at room temperature for 4 hours to get a white solid in the slurry. The slurry was evaporated and dried

under ambient conditions. An off-white crystalline solid was obtained and identified as Form D by XRPD analysis.

Method 2: 5.01g of light brown amorphous (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide was dissolved in 10 mL of EtOAc to yield a light brown solution and a brown gel. 0.10 g of (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide-Form D seed from Method 1 was added and the solution became a wet cake within 5 minutes. 10 mL of EtOAc was added to form a slurry. The slurry was stirred under ambient conditions overnight. The brown gel disappeared to obtain a slurry of an off-white solid with brown solution. The slurry was filtered, and the solid was washed twice with EtOAc. The off-white solid was dried under ambient conditions. 4.78 g of an off-white crystalline solid was obtained and identified as Form D by XRPD analysis.

Form D (Method 2) was analyzed by XRPD and the results are tabulated below (Table 20) and shown in Figure 7.

Table 20

Angle (2 θ ±0.2°)	Intensity (%)
21.8	100.0
6.4	74.8
16.6	59.2
8.9	50.1
22.5	48.3
8.1	43.0
19.1	40.4
19.9	40.0
20.6	36.8
26.6	30.2
9.5	26.6
16.0	24.8
14.0	20.9
24.1	19.8
28.9	19.2
18.6	19.0
13.4	18.8
25.8	17.1
25.4	15.4
26.3	14.8
32.1	13.5

10.9	12.6
12.1	12.2
14.8	12.2
28.4	11.9
30.0	11.4
33.0	11.4
38.9	10.2
36.5	10.1
15.2	9.3

Single crystals of Form D were obtained from slow evaporation of an EtOAc solution. Single crystal structure analysis confirmed that Form D is a hemi-EtOAc solvate form.

Crystallographic data: Space group triclinic *P*1, unit cell dimensions: *a* = 14.051(2) Å, *b* = 14.289(2) Å, *c* = 14.756(2) Å, α = 81.174(5)°, β = 77.476(5)°, γ = 85.331(6)°, *V* = 2854.5(8) Å³.

Form D was analyzed by thermal techniques. DSC analysis indicated that Form D has an endotherm event of desolvation with an onset at 116 °C and a peak at 119 °C. TGA indicated that Form D exhibits a mass loss of about 8.0 % upon heating from about 25 °C to about 200 °C. A representative DSC/TGA thermogram of Form D is shown in Figure 8.

Example 68E: Form A (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide Saccharine Salt

25.1 mg of (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide (0.05 mmol) was dissolved in 1 mL of MeOH, and 2.0 mL (0.10 mmol) of 0.05 M of saccharin MeOH solution was added to yield a light yellow solution. To the solution, 1 mL of ACN was added and the resulting solution was evaporated under ambient conditions. A partial yellow crystalline material was obtained. Approximately 10 mg of the resulting material was dissolved in 2 mL of ACN and the resulting yellow solution was evaporated slowly to obtain yellow needle crystals identified as Form A (2*R*)-N-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide saccharin salt by XRPD analysis.

Form A of the saccharin salt was analyzed by XRPD and the results are tabulated below (Table 21) and shown in Figure 9.

Table 21

Angle (2 θ ±0.2°)	Intensity (%)
14.4	100.0
10.7	52.0
7.1	48.1
21.4	44.3
10.1	42.1
8.5	38.9
8.2	36.5
17.3	36.5
16.3	27.3
9.4	26.0
11.6	25.2
17.8	23.0
20.8	22.5
20.1	18.2
18.5	16.8
18.8	16.7
25.4	16.3
24.9	15.9
21.8	15.6
27.4	15.2
27.8	15.2
26.7	13.9
24.4	13.6
23.4	13.4
15.4	13.3
28.9	13.1
30.4	12.2
35.3	11.9
31.1	11.2
12.5	11.1
19.4	10.5

Form A of the saccharin salt was analyzed by thermal techniques. DSC analysis indicated that Form A has an endotherm event of melting point with an onset at 163 °C and a peak at 169 °C. TGA indicated that Form A exhibits a mass loss of about 3.1 % upon heating from about 25 °C to about 150 °C. A representative DSC/TGA thermogram of Form D is shown in Figure 10.

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Example 68F: Form B (2R)-N-(3-{2-[(3-methoxy-1-methyl-1H-pyrazol-4-yl)amino]pyrimidin-4-yl}-1H-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide Saccharine Salt

246 mg of light yellow (2R)-N-(3-{2-[(3-methoxy-1-methyl-1H-pyrazol-4-yl)amino]pyrimidin-4-yl}-1H-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide (0.5 mmol) and 184 mg (1.0 mmol) of saccharine was dissolved in 3 mL of acetonitrile and 1 mL of MeOH to yield a clear yellow solution. The solution was evaporated to have about 1.0 mL of solvent and a yellow crystalline material precipitated. The suspension was stirred for 30 minutes and filtered. The solid was dried under ambient conditions. A yellow crystal material was obtained and identified as Form B (2R)-N-(3-{2-[(3-methoxy-1-methyl-1H-pyrazol-4-yl)amino]pyrimidin-4-yl}-1H-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide saccharine salt by XRPD.

Form B of the saccharin salt was analyzed by XRPD and the results are tabulated below (Table 22) and shown in Figure 11.

Table 22

Angle (2θ±0.2°)	Intensity (%)
6.6	100.0
13.4	99.8
25.3	85.8
18.1	82.7
8.0	76.8
17.2	71.0
9.1	65.3
21.8	64.2
9.9	64.0
26.0	58.9
16.4	58.7
26.4	57.9
21.3	57.3
20.2	54.5
27.9	48.6
14.7	48.1
16.2	47.6
24.1	46.8
20.4	45.0
23.0	45.0
24.7	43.3
18.7	42.4

11.0	39.9
15.2	39.2
15.7	35.1
29.7	34.4

Form B of the saccharin salt was analyzed by thermal techniques. DSC analysis indicated that Form B has a broad endotherm event of desolvation with a peak at 53 °C, followed by two endotherm events, one with an onset at 153 °C and a peak at 162 °C and the other with an onset at 176 °C and a peak at 182 °C. TGA indicated that Form B exhibits a mass loss of about 2.7 % upon heating from about 25 °C to about 100 °C. A representative DSC/TGA thermogram of Form B is shown in Figure 12.

Example 68G: Form C (2R)-N-(3-{2-[(3-methoxy-1-methyl-1H-pyrazol-4-yl)amino]pyrimidin-4-yl}-1H-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide Saccharine Salt

Approximately 200 mg of Form B (2R)-N-(3-{2-[(3-methoxy-1-methyl-1H-pyrazol-4-yl)amino]pyrimidin-4-yl}-1H-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide saccharin salt was slurried in acetone for 3 days, and the resulting slurry was evaporated under ambient conditions. A yellow crystal material was obtained and identified as Form C (2R)-N-(3-{2-[(3-methoxy-1-methyl-1H-pyrazol-4-yl)amino]pyrimidin-4-yl}-1H-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide saccharine salt by XRPD.

Form C of the saccharin salt was analyzed by XRPD and the results are tabulated below (Table 23) and shown in Figure 13.

Table 23

Angle (2θ±0.2°)	Intensity (%)
5.5	100.0
17.0	82.5
26.3	82.0
8.2	80.3
14.9	76.9
24.9	73.4
16.0	72.9
14.4	70.6
18.1	57.6
12.4	55.8
9.4	53.0
20.7	52.4

28.0	46.8
19.2	46.6
19.8	45.3
21.5	43.0
23.8	40.2
15.4	36.7
13.4	33.1
22.8	32.6
29.8	28.0

Example 68H: Form D (2R)-N-(3-{2-[(3-methoxy-1-methyl-1H-pyrazol-4-yl)amino]pyrimidin-4-yl}-1H-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide Saccharine Salt

Approximately 15 mg of Form B or Form C (2R)-N-(3-{2-[(3-methoxy-1-methyl-1H-pyrazol-4-yl)amino]pyrimidin-4-yl}-1H-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide saccharin salt was slurried in 0.5 mL of water. The resulting slurry was dried in the sample holder and was measured by XRPD analysis and Form D (2R)-N-(3-{2-[(3-methoxy-1-methyl-1H-pyrazol-4-yl)amino]pyrimidin-4-yl}-1H-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide saccharin salt was identified.

Form D of the saccharin salt was analyzed by XRPD and the results are tabulated below (Table 24) and shown in Figure 14.

Table 24

Angle (2θ±0.2°)	Intensity (%)
19.2	100.0
5.4	96.7
20.1	95.7
6.8	93.7
15.4	83.0
7.7	74.1
20.8	72.1
13.8	66.8
17.3	61.4
13.4	58.5
23.1	58.3
10.4	43.4
9.3	42.5
14.3	33.4
16.9	30.9
11.5	27.2

27.2	22.8
26.2	21.1

Example 68l: Form E (2R)-N-(3-{2-[(3-methoxy-1-methyl-1H-pyrazol-4-yl)amino]pyrimidin-4-yl}-1H-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide Saccharine Salt

About 15 mg of Form C (2R)-N-(3-{2-[(3-methoxy-1-methyl-1H-pyrazol-4-yl)amino]pyrimidin-4-yl}-1H-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide saccharin salt was slurried in 0.5 mL of EtOH. The resulting slurry was evaporated under ambient conditions. A yellow powder was obtained and was identified as Form E (2R)-N-(3-{2-[(3-methoxy-1-methyl-1H-pyrazol-4-yl)amino]pyrimidin-4-yl}-1H-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide saccharin salt by XRPD analysis.

Form E of the saccharin salt was analyzed by XRPD and the results are tabulated below (Table 25) and shown in Figure 15.

Table 25

Angle (2 θ ±0.2°)	Intensity (%)
15.4	100.0
19.3	96.9
5.5	86.4
14.7	78.5
23.2	76.8
6.7	73.3
20.1	71.0
7.8	66.8
26.3	58.4
17.7	53.3
25.2	53.0
18.2	52.5
24.6	52.0
24.0	50.3
13.5	50.1
27.3	49.4
9.3	45.7
17.1	44.5
21.4	42.1
16.4	37.2
29.2	33.9
11.8	33.5

Example 68J: (2R)-N-(3-{2-[(3-methoxy-1-methyl-1H-pyrazol-4-yl)amino]pyrimidin-4-yl}-1H-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide Saccharine Hydrochloride Salt

249 mg (0.50 mmol) of (2R)-N-(3-{2-[(3-methoxy-1-methyl-1H-pyrazol-4-yl)amino]pyrimidin-4-yl}-1H-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide was dissolved in 4 mL of MeOH to yield a light brown solution. 0.5 ml of 1N HCl aqueous solution (0.50 mmol) was added and the color of the solution turned to yellow. To the yellow solution, 0.5 mmol of saccharine was added, and the saccharine gradually dissolved in the solution to yield a yellow solution. The solution was evaporated under ambient conditions to dry. The resulting solid was slurried in 4 mL of acetone overnight, then filtered and washed with acetone. The yellow solid was dried in air and identified as (2R)-N-(3-{2-[(3-methoxy-1-methyl-1H-pyrazol-4-yl)amino]pyrimidin-4-yl}-1H-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide saccharine hydrochloride salt by XRPD analysis.

The saccharin hydrochloride salt was analyzed by XRPD and the results are tabulated below (Table 26) and shown in Figure 16.

Table 26

Angle (2θ±0.2°)	Intensity (%)
13.7	100.0
20.2	84.0
26.0	82.7
17.3	80.8
8.4	59.9
25.4	57.4
20.6	54.8
27.6	53.9
10.6	52.1
26.6	49.4
18.4	48.5
20.9	47.1
24.3	44.8
15.0	43.9
15.7	42.3
12.9	39.7
25.1	39.3
18.8	39.2
12.5	37.7
22.6	36.2
16.1	33.1
29.4	33.1

23.3	32.9
27.2	32.9
28.9	32.3
21.7	31.0
21.2	27.1
32.0	26.3
22.2	22.9
32.8	21.5
33.6	21.1
29.9	19.7
30.5	19.6
31.2	18.8

Example 68K: (2R)-N-(3-{2-[(3-methoxy-1-methyl-1H-pyrazol-4-yl)amino]pyrimidin-4-yl}-1H-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide Napadosylic Salt

35.8 mg of off-white amorphous (2R)-N-(3-{2-[(3-methoxy-1-methyl-1H-pyrazol-4-yl)amino]pyrimidin-4-yl}-1H-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide was dissolved in 0.8 mL of EtOH:water mixture (70:30), and 29.02 mg of napadisyllic acid tetrahydrate was dissolved in 0.5 mL of the same solvent. The counter ion solution was added to the solution of (2R)-N-(3-{2-[(3-methoxy-1-methyl-1H-pyrazol-4-yl)amino]pyrimidin-4-yl}-1H-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide dropwise and a yellow precipitate was obtained. The slurry was stirred at the ambient condition for overnight. The slurry was filtered to obtain a crystalline material.

The napadisyllic acid salt was analyzed by XRPD and the results are tabulated below (Table 27) and shown in Figure 17.

Table 27

Angle (2 θ ±0.2°)	Intensity (%)
5.2	100.0
10.4	78.5
7.8	74.0
18.3	68.1
17.3	64.1
15.0	59.3
22.1	58.7
25.6	58.7
20.9	58.2
15.6	57.4
20.3	56.3

16.9	52.9
16.1	51.1
26.5	46.9
24.2	46.6
21.2	46.6
24.0	44.2
27.5	42.7
23.2	42.5
19.9	41.6
14.2	39.0
7.2	38.4
12.1	32.2
13.5	32.2
11.3	31.0
30.6	24.3
31.2	23.8

Example 68L: (2R)-N-(3-{2-[(3-methoxy-1-methyl-1H-pyrazol-4-yl)amino]pyrimidin-4-yl}-1H-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide Trimesic Salt

5 30 mg of off-white amorphous (2R)-N-(3-{2-[(3-methoxy-1-methyl-1H-pyrazol-4-yl)amino]pyrimidin-4-yl}-1H-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide was dissolved in 0.5 mL of EtOH:water mixture (70:30), and 14.16 mg of trimesic acid was dissolved in 0.6 mL of the same solvent. The counter ion solution was added to the solution of (2R)-N-(3-{2-[(3-methoxy-1-methyl-1H-pyrazol-4-yl)amino]pyrimidin-4-yl}-1H-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide dropwise. The resulting solution was subjected to slow evaporation and isolated by centrifugation.

The trimesic acid salt was analyzed by XRPD and the results are tabulated below (Table 28) and shown in Figure 18.

Table 28

Angle (2θ±0.2°)	Intensity (%)
4.2	100.0
12.4	66.3
7.8	63.0
25.1	27.8
24.9	27.3
2.1	20.6

27.5	19.1
12.1	16.3
15.7	16.2
10.6	14.4
26.1	13.8
16.5	13.2
15.9	12.8
8.3	12.6
19.8	11.8
20.7	11.8
25.7	11.4
29.1	11.0
8.7	10.7
22.1	10.4
17.5	10.0
23.0	9.9
19.2	9.0
13.9	9.0
23.5	8.9
23.8	8.9
28.1	8.2
26.9	8.1
21.2	7.6
14.3	7.4
18.2	7.4
22.7	7.4
15.2	7.3
17.0	6.9
37.7	6.5
18.6	6.3
33.2	6.1
34.3	6.1
39.3	5.6
39.3	5.4
30.6	5.3
11.3	5.0
31.2	4.9

Example 69: Example 32 in combination with Osimertinib-*In Vivo* Efficacy and Pharmacodynamics in the H1975 Xenograft Model

SUMMARY: NCI-H1975 tumor xenografts were grown subcutaneously in female nude mice. The mice were treated by oral dosing with vehicle, Example 32, the EGFR inhibitor osimertinib (AZD9291, TAGRISSO™), an irreversible inhibitor of T790M mutant EGFR, or combinations of Example 32 and osimertinib. Average tumor size at the start of treatment (10 days after implant) was 189 mm³. Tumor volume was measured twice a week. Additional mice were treated with the same doses of Example 32 and osimertinib, and tumors and plasma harvested after one day of treatment, for analysis of pSTAT3 (Y705) levels in tumors and drugs levels in plasma.

The combination of Example 32 with osimertinib resulted in enhanced antitumor activity, compared to treatment with osimertinib alone. There was no significant antitumor activity observed after treatment with Example 32 alone. The enhanced antitumor activity of the combination correlated with pSTAT3 knockdown by Example 32, consistent with a role for JAK/STAT signaling in escape from pEGFR inhibition.

Osimertinib, an irreversible inhibitor of T790M mutant EGFR, overcomes T790M-mediated resistance to EGFR inhibitors such as gefitinib and erlotinib in lung cancer. This study was carried out to evaluate the ability of Example 32 to enhance the antitumor response to osimertinib in mice bearing subcutaneous NCI-H1975 tumor xenografts. The EGFR gene in the NCI-H1975 tumors is mutated at L858R and also contains the T790M resistance mutation.

MATERIALS AND METHODS: NCI-H1975 cells (a human NSCLC cell line with L858R and T790M mutations in the EGFR gene) tumor cells were implanted subcutaneously in female NCr nude mice (Taconic Laboratories), 3 x 10⁶ cells / mouse. Ten days after cell implantation, mice were randomized into 10 groups (6-8 mice/group, average tumor volume 189 mm³, range 152-250 mm³), and were dosed orally with either vehicle (20% captisol), Example 32 as a single agent (12.5 mg/kg, 25 mg/kg, 50 mg/kg), osimertinib as a single agent (2.5 mg/kg), or combinations of Example 32 and osimertinib (osimertinib at 2.5 mg/kg and Example 32 at 12.5 mg/kg, 25 mg/kg and 50 mg/kg) for 18 days. Some of the mice from the tumor implant that were not randomized into these groups were treated for a single day with these compounds for pharmacokinetic and pharmacodynamic analysis (tumor and plasma collected for analysis of drug levels in plasma and pSTAT3 (Y705) levels in tumor lysates, samples collected 2, 8 and 24 hours after the AM dose). Tumor length and width were measured by caliper, and tumor volume calculated using the formula volume = (length x width²)*π/6. Example 32 was formulated in water, adjusted to pH 2 with methane sulfonic acid. Osimertinib was formulated in

0.5% HPMC in water. All formulations were administered by oral gavage, at a volume of 5 ml/kg. Osimertinib was dosed QD (AM), Example 32 was dosed BID (AM and PM, 8 hours apart). In the groups that were dosed with osimertinib in combination with Example 32, the AM dosing of osimertinib occurred 3 hours before Example 32 to minimize exposure interactions.

AZD1480 (5-chloro-N2-[(1S)-1-(5-fluoro-2-pyrimidinyl)ethyl]-N4-(5-methyl-1H-pyrazol-3-yl)-2,4-pyrimidinediamine, *see* U.S. Patent Application Publication No. US20080287475), a JAK1/2 inhibitor and a positive control for 100% pSTAT3 knockdown in the pharmacokinetic/pharmacodynamic experiment was formulated in 0.5% HPMC/0.1% Tween® 80 in water, and administered by oral gavage at a volume of 5 ml/kg. Y705 phosphorylated STAT3 (pSTAT3) levels were measured in tumor lysates using a sandwich ELISA (PathScan Phospho-STAT3 Sandwich ELISA Kit, CST #7146B). Drug levels in plasma were measured by LC/MS, using a Linear Ion Trap Quadrupole LC/MS/MS Mass Spectrometer (QTRAP 5500 model 1024945-BB, AB Sciex Instruments), with separation on a Waters Xbridge C18 column.

RESULTS: The addition of Example 32 enhanced the antitumor activity of osimertinib, compared to treatment with osimertinib alone. Example 32 administered as a single agent had only weak antitumor activity relative to vehicle control treatment (Figure 19). The enhancement of antitumor activity increased with increasing dose of Example 32 (12.5 mg/kg up to 50 mg/kg). At the last day of treatment, the inhibition of tumor growth of all of the combination treatments was statistically significant ($p < 0.05$), compared to single agent osimertinib (Table 29). All treatments were well tolerated, with no significant weight loss (Figure 20) or other outward signs observed over the course of treatment. A dose dependent increase in pSTAT3 knockdown was observed with increasing dose of Example 32 (Figure 21), which correlated with an increasing level of Example 32 in plasma.

Table 29. p values For Antitumor Activity of Osimertinib and Example 32 combinations, After 18 Days of Treatment

Treatment	vs. Osimertinib Single Agent ^a	vs. Vehicle Control ^b
osimertinib, 2.5 mg/kg QD	---	0.0002
osimertinib, 2.5 mg/kg QD + Example 32, 12.5 mg/kg BID	0.0027	< 0.0001
osimertinib, 2.5 mg/kg QD + Example 32, 25 mg/kg BID	< 0.0001	< 0.0001
osimertinib, 2.5 mg/kg QD + Example 32, 50 mg/kg BID	< 0.0001	< 0.0001

^a Two-sided test

^b One-sided test

CONCLUSION: The increased antitumor activity of the osimertinib in combination with Example 32 compared to single agent osimertinib, and the correlation with pSTAT3 knockdown by Example 32, is consistent with a role for STAT3 signaling in escape from, or resistance to, EGFR inhibition in this model of non-small cell lung cancer. This conclusion is further supported by the correlation of increased pSTAT3 knockdown with increased combination antitumor activity over the dose range of Example 32. The results support the hypothesis that inhibition of STAT3 signaling can enhance the antitumor activity of an EGFR inhibitor in T790M EGFR mutant NSCLC.

Example 70: Example 32 in combination with gefitinib-*In Vivo* Efficacy and Pharmacodynamics in the PC-9 Xenograft Model

SUMMARY: PC9 tumor xenografts were grown subcutaneously in female nude mice. The mice were treated, by oral dosing, with vehicle, Example 32, the EGFR inhibitor gefitinib, or combinations of Example 32 and gefitinib. Average tumor size at the start of treatment was 240 mm³. Tumor volume was measured twice a week. At the end of treatment (21 days), tumors were harvested for analysis of pSTAT3 (Y705) levels.

The combination of Example 32 and gefitinib resulted in enhanced antitumor activity, compared to treatment with gefitinib alone. The enhanced antitumor activity of the combination correlated with pSTAT3 knockdown by Example 32, consistent with a role for JAK/STAT signaling in escape from pEGFR inhibition.

Gefitinib is an inhibitor of mutant EGFR, with activity in non-small cell lung cancer patients bearing the del19 and L858R mutations in EGFR. This study was carried out to evaluate the ability of Example 32 to enhance the antitumor response to gefitinib in mice bearing subcutaneous PC-9 tumor xenografts. The EGFR gene in PC-9 tumors contains the del19 mutation.

MATERIALS AND METHODS: PC-9 cells (a human NSCLC cell line with the EGFR del19 mutation), were implanted subcutaneously in female CB17-SCID mice (Charles River Laboratories), 2 x 10⁶ cells/mouse. Thirty-two days after cell implantation, the mice were randomized into 5 groups (7 mice/group, average tumor volume 240 mm³, range 204-298 mm³). Mice were dosed orally with either vehicle (1% Tween® 80, QD), gefitinib as a single agent or combinations of gefitinib and Example 32, for 21 days at the doses and schedules indicated in Figures 22 and 23. On the last day of treatment, tumors were collected (2, 8 and 24 hours after the AM dose) for analysis of pSTAT3 (Y705) levels in tumor lysates. Tumor length and width were measured by caliper, and tumor volume calculated using the formula volume = (length x

width²)* π /6. Example 32 was formulated in water, adjusted to pH 2 with methane sulfonic acid. Gefitinib was formulated in 1% Tween® 80 in water. All formulations were administered by oral gavage, at a volume of 5 ml/kg. Gefitinib was dosed QD (AM), Example 32 was dosed BID (AM and PM, 8 hours apart). In the groups that were dosed with gefitinib in combination with Example 32, the AM dosing of gefitinib occurred before Example 32 (less than 10 minutes between the two). Y705 phosphorylated STAT3 (pSTAT3) levels were measured in tumor lysates (collected 2, 8 and 24 hours after the AM dose) by Western Blot analysis (4-12% tris-glycine PAGE gels and semi-dry transfer to PVDF membrane; immunoblotting for pSTAT3 with Cell Signaling Technologies (CST) #9145 primary antibody and CST #7074 goat anti-rabbit HRP-linked secondary antibody; immunoblotting for beta-actin with CST #3700 primary antibody and CST #7076 goat anti-mouse HRP-linked secondary antibody), with electrochemiluminescence captured using ImageQuant LAS 4000 and analysis with ImageQuant TL software.

RESULTS: The addition of Example 32 enhanced the antitumor activity of gefitinib (iressa), compared to treatment with gefitinib (Figure 22). For treatment with gefitinib in combination with Example 32, the enhancement of antitumor activity increased with increasing dose of Example 32, from 12.5 mg/kg to 50 mg/kg. When Example 32 was dosed 2 days on/5 days off on a weekly cycle, in combination with daily gefitinib, the antitumor activity was similar to that of Example 32 at 12.5 mg/kg administered 7 days a week in combination with gefitinib. On the last day of treatment, the inhibition of tumor growth with gefitinib in combination with 50 mg/kg of Example 32 was statistically significant ($p < 0.05$), compared to single agent gefitinib (Table 30). Gefitinib in combination with 12.5 mg/kg of Example 32 daily and with 50 mg/kg Example 32 dosed 2 days on/5 days off had greater activity than single agent gefitinib, but did not quite achieve statistical significance. All treatments were well tolerated, with no significant weight loss (Figure 23) or other outward signs observed over the course of treatment. A dose dependent increase in the duration of pSTAT3 knockdown was observed with increasing dose of Example 32 (Figure 24).

Table 30. p values for Antitumor Activity of Gefitinib in Combination with Example 32, After 36 Days of Treatment

Treatment	vs. gefitinib single agent ^a	vs. vehicle control ^b
gefitinib, 6.25 mg/kg QD	---	< 0.0001
gefitinib, 6.25 mg/kg QD + Example 32, 12.5 mg/kg BID	0.1263	< 0.0001
gefitinib, 6.25 mg/kg QD + Example 32, 50 mg/kg BID	0.0006	< 0.0001
gefitinib, 6.25 mg/kg QD + Example 32, 50 mg/kg BID, 2 days on / 5 days off / week	0.0685	< 0.0001

^a Two-sided test^b One-sided test

5

CONCLUSION: The increased antitumor activity of Example 32 in combination with gefitinib compared to single agent gefitinib, and the correlation with pSTAT3 knockdown by Example 32, is consistent with a role for STAT3 signaling in escape from, or resistance to, EGFR inhibition in this model of non-small cell lung cancer. This conclusion is further supported by the correlation of increased pSTAT3 knockdown with increased combination antitumor activity over the dose range of Example 32 (12.5 to 50 mg/kg). The results support the hypothesis that inhibition of STAT3 signaling can enhance the antitumor activity of an EGFR inhibitor in NSCLC tumors bearing the exon 19 deletion in the EGFR gene.

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Example 71: Example 32 plus gefitinib combination-*In Vivo* Efficacy and Pharmacodynamics in the H1650 Xenograft Model

SUMMARY: NCI-H1650 tumor xenografts were grown subcutaneously in female nude mice. The mice were treated, by oral dosing, with vehicle, Example 32, the EGFR inhibitor gefitinib, or combinations of Example 32 plus gefitinib. Average tumor size at the start of treatment was 257 mm³. Tumor volume was measured twice a week. Additional tumor bearing mice were treated with the same doses of Example 32 and gefitinib, and tumors and plasma were harvested after one day of treatment for analysis of pSTAT3 levels in tumors and drug levels in plasma.

20

The combination of Example 32 plus gefitinib resulted in enhanced antitumor activity, compared to treatment with gefitinib alone. The enhanced antitumor activity of the combination correlated with pSTAT3 knockdown by Example 32, consistent with a role for JAK/STAT signaling in escape from pEGFR inhibition.

25

Gefitinib is an inhibitor of mutant EGFR, with activity in non-small cell lung cancer patients bearing the del19 and L858R mutations in EGFR. This study was carried out to

evaluate the ability of Example 32 to enhance the antitumor response to gefitinib in mice bearing subcutaneous NCI-H1650 tumor xenografts. The EGFR gene in NCI-H1650 tumors contains the del19 mutation.

MATERIALS AND METHODS: NCI-H1650 cells (a human NSCLC cell line with the EGFR del19 mutation), were implanted subcutaneously in female CB17-SCID mice (Charles River Laboratories), 5×10^6 cells / mouse. Twenty-three days after cell implantation, the mice were randomized into 6 groups (9 mice/group, average tumor volume 257 mm³, range 205-303 mm³). Mice were dosed orally with either vehicle (1% Tween® 80, QD), gefitinib as a single agent, Example 32 as a single agent, or combinations of gefitinib plus Example 32, at the doses and schedules indicated in Figures 25 and 26 for 21 days. Tumor length and width were measured by caliper, and tumor volume calculated using the formula $\text{volume} = (\text{length} \times \text{width}^2) \times \pi / 6$. Example 32 was formulated in water, adjusted to pH 2 with methane sulfonic acid. Gefitinib was formulated in 1% Tween® 80 in water. All formulations were administered by oral gavage, at a volume of 5 ml/kg. Gefitinib was dosed QD (AM), Example 32 was dosed BID (AM and PM, 8 hours apart). In the groups that were dosed with gefitinib in combination with Example 32, the AM dosing of gefitinib occurred before dosing with Example 32 (less than 10 minutes between the two). Additional tumor bearing mice were treated with the same doses of Example 32 and gefitinib, and tumors and plasma were harvested after one day of treatment for analysis of pSTAT3 levels in tumors and drug levels in plasma. AZD1480 (5-chloro-N2-[(1S)-1-(5-fluoro-2-pyrimidinyl)ethyl]-N4-(5-methyl-1H-pyrazol-3-yl)-2,4-pyrimidinediamine, *see* U.S. Patent Application Publication No. US20080287475), a JAK1/2 inhibitor and a positive control for 100% pSTAT3 knockdown in the pharmacokinetic/pharmacodynamic experiment was formulated in 0.5% HPMC/0.1% Tween® 80 in water, and administered by oral gavage at a volume of 5 ml/kg. Y705 phosphorylated STAT3 (pSTAT3) levels were measured in tumor lysates (collected 2, 8 and 24 hours after the AM dose) by Western Blot analysis (4-12% tris-glycine PAGE gels and semi-dry transfer to PVDF membrane; immunoblotting for pSTAT3 with Cell Signaling Technologies (CST) #9145 primary antibody and CST #7074 goat anti-rabbit HRP-linked secondary antibody; immunoblotting for GAPDH with CST #2118 primary antibody and CST #7074 goat anti-rabbit HRP-linked secondary antibody), with electrochemiluminescence captured using ImageQuant LAS 4000 and analysis with ImageQuant TL software. Drug levels in plasma were measured by LC/MS, using a Linear Ion Trap Quadrupole LC/MS/MS Mass Spectrometer (QTRAP 5500 model 1024945-BB, AB Sciex Instruments), with separation on a Waters Xbridge C18 column.

RESULTS: The addition of Example 32 enhanced the antitumor activity of gefitinib, compared to single agent gefitinib (Figure 25). Example 32 administered as a single agent had only modest antitumor activity. For the treatment of gefitinib in combination with Example 32, enhancement of antitumor activity increased with increasing dose of Example 32, from 25 mg/kg to 50 mg/kg. At the last day of treatment, the inhibition of tumor growth with gefitinib in combination with 25 or 50 mg/kg of Example 32 was statistically significant ($p < 0.05$), compared to single agent gefitinib (Table 31). All treatments were well tolerated, with no significant weight loss (Figure 26) or other outward signs observed over the course of treatment. Knockdown of pSTAT3 was observed with Example 32 as single agent and in combination (Figure 27), measured after one day of treatment.

Table 31. p values for Antitumor Activity of Gefitinib Plus Example 32 Combination, After 21 Days of Treatment

Treatment	vs. gefitinib single agent ^a	vs. vehicle control ^b
gefitinib, 6.25 mg/kg QD	---	< 0.0001
Example 32, 25 mg/kg BID	0.0314	0.0002
Example 32, 50 mg/kg BID	0.282	< 0.0001
gefitinib, 6.25 mg/kg QD + Example 32, 25 mg/kg BID	< 0.0001	< 0.0001
gefitinib, 6.25 mg/kg QD + Example 32, 50 mg/kg BID	< 0.0001	< 0.0001

^a Two-sided test

^b One-sided test

CONCLUSIONS: The increased antitumor activity of Example 32 in combination with gefitinib compared to single agent gefitinib and the correlation with pSTAT3 knockdown by Example 32 is consistent with a role for STAT3 signaling in escape from, or resistance to, EGFR inhibition in this model of non-small cell lung cancer. The results support the hypothesis that inhibition of STAT3 signaling can enhance the antitumor activity of an EGFR inhibitor in NSCLC tumors bearing the del19 mutation in the EGFR gene.

Example 72: Example 32 in combination with Osimertinib-*In Vivo* Efficacy and Pharmacodynamics in the LG1049 Xenograft PDX Model

SUMMARY: LG1049 non-small cell lung cancer PDX tumor xenografts were grown subcutaneously in female NSG mice. The mice were treated, by oral dosing, with vehicle, Example 32 as a single agent, the EGFR inhibitor osimertinib (an irreversible inhibitor of T790M mutant EGFR) as a single agent, or combinations of Example 32 with osimertinib. Average

tumor size at the start of treatment was 189 mm³. Tumor volume was measured twice a week. A separate set of mice bearing LG1049 tumors were treated for 5 days and tumors harvested for analysis of pSTAT3 (Y705) and pEGFR levels.

The combination of Example 32 with osimertinib resulted in enhanced tumor regression, compared to treatment with osimertinib alone. There was no significant antitumor activity observed after treatment with Example 32 alone. When treatment was stopped after 28 days, tumors rapidly regrew in mice that had been treated with osimertinib as a single agent or with osimertinib in combination with Example 32. When Example 32 treatment was continued, tumors also regrew, but more slowly. Analysis of tumors taken from mice treated for 5 days confirmed robust knockdown of pSTAT3 and pEGFR by Example 32 and osimertinib, respectively.

Osimertinib, an irreversible inhibitor of T790M mutant EGFR, overcomes T790M-mediated resistance to EGFR inhibitors such as gefitinib and erlotinib in lung cancer. This study was carried out to evaluate the ability of Example 32 to enhance the antitumor response to osimertinib in mice bearing subcutaneous LG1049 tumor xenografts. LG1049 is a non-small cell lung cancer (NSCLC) primary tumor xenograft (PDX) model, in which the EGFR gene contains the T790M resistance mutation.

MATERIALS AND METHODS: LG1049 tumor fragments were implanted subcutaneously in female NSG mice (JAX Stock No. 005557). Once tumor volumes reached ~125-275 mm³, mice were randomized into 5 groups (10 mice/group, average tumor volume 189 mm³, range 138-253 mm³). Mice were dosed orally with either vehicle, Example 32 as a single agent, osimertinib as a single agent, or combinations of Example 32 with osimertinib, at the doses and schedules indicated in Figures 28 and 29 for 28 days (18 days for single agent Example 32). In one of the two groups that received Example 32 in combination with osimertinib, treatment with Example 32 was continued for an additional 14 days. A separate set of tumor bearing mice were treated with vehicle, Example 32 as a single agent, osimertinib as a single agent, or a combination of Example 32 and osimertinib for 5 days, and tumors collected for analysis of pSTAT3 (Y705) and pEGFR levels in tumor lysates. Tumor length and width were measured by caliper, and tumor volume calculated using the formula volume = (length x width²)* π /6. Example 32 was formulated in water, adjusted to pH 2 with methane sulfonic acid. Osimertinib was formulated in 0.5% HPMC in water. All formulations were administered by oral gavage, at a volume of 5 ml/kg. Osimertinib was dosed QD (AM), Example 32 was dosed BID (AM and PM, 8 hours apart). In the groups that were dosed with Example 32 in combination with osimertinib, the AM dosing of osimertinib occurred before the dosing of Example 32 (less

than 10 minutes between the two). Phosphorylated STAT3 and phosphorylated EGFR levels were measured in tumor lysates by Western Blot analysis (4-12% tris-glycine PAGE gels and semi-dry transfer to PVDF membrane; immunoblotting for pSTAT3 with Cell Signaling Technologies (CST) #9145 primary antibody and CST #7074 goat anti-rabbit HRP-linked secondary antibody; immunoblotting for pEGFR pY1173 with Epitomics #1124 primary antibody and CST #7074 goat anti-rabbit HRP-linked secondary antibody; immunoblotting for GAPDH with CST #2118 primary antibody and CST #7074 goat anti-rabbit HRP-linked secondary antibody), with electrochemiluminescence captured using ImageQuant LAS 4000 and analysis with ImageQuant TL software.

RESULTS: Addition of Example 32 enhanced the tumor regression induced by osimertinib, compared to treatment with osimertinib alone. Example 32 administered as a single agent had no significant activity relative to vehicle control treatment (Figure 28). When treatment was stopped, after 28 days of dosing (osimertinib and Example 32 in combination with osimertinib groups), tumors regrew. When osimertinib treatment was continued for an additional 14 days (one of the combination groups) tumors also regrew, but more slowly. Mice treated with vehicle or Example 32 alone experienced significant weight loss (Figure 29), suggesting that the weight loss was the result of tumor growth. The Example 32 single agent group was terminated early due to excessive weight loss. Tumor dependent weight loss was seen in other experiments with this model. The lack of significant weight loss in treatment groups in which tumors regressed (osimertinib as a single agent, and Example 32 in combination with osimertinib), is consistent with weight loss being dependent on tumor growth.

The doses and schedules of Example 32 and osimertinib used in this experiment resulted in robust knockdown of pSTAT3 and pEGFR after 5 days of treatment, measured at 4H after the final dose (Figure 30).

CONCLUSION: The increased antitumor activity of Example 32 in combination with osimertinib compared to single agent osimertinib, and the correlation with pSTAT3 knockdown by Example 32, is consistent with a role for STAT3 signaling in escape from, or resistance to, EGFR inhibition in this model of EGFR T790M mutant non-small cell lung cancer. The results support the hypothesis that inhibition of STAT3 signaling can enhance the antitumor activity of an EGFR inhibitor in T790M EGFR mutant NSCLC.

Example 73: Example 32 in Combination with Osimertinib-*In Vivo* Efficacy of Intermittent Example 32 Dosing Schedules in the H1975 Xenograft Model

SUMMARY: NCI-H1975 tumor xenografts were grown subcutaneously in female nude mice. The mice were treated by oral dosing with vehicle, Example 32 as a single agent, the EGFR inhibitor osimertinib (an irreversible inhibitor of T790M mutant EGFR) as a single agent, or combinations of osimertinib and Example 32 on different intermittent dosing schedules. Average tumor size at the start of treatment was 185 mm³. Tumor volume was measured twice a week.

Compared to treatment with osimertinib alone, the combination of Example 32 and osimertinib resulted in enhanced antitumor activity on all the schedules of the treatment tested. There was a trend to reduced efficacy with less intensive dosing schedules. The enhanced antitumor activity of the combination and the correlation of greater efficacy with more intensive dosing schedules of Example 32 is consistent with a role for JAK/STAT signaling in escape from pEGFR inhibition.

This study was carried out to evaluate the ability of Example 32 to enhance the antitumor response to osimertinib in mice bearing subcutaneous NCI-H1975 tumor xenografts, and to investigate the frequency of target coverage with Example 32 that is required to retain combination activity. The EGFR gene in the NCI-H1975 tumors is mutated at L858R and also contains the T790M resistance mutation.

MATERIALS AND METHODS: NCI-H1975 cells (a human NSCLC cell line with L858R and T790M mutations in the EGFR gene), were implanted subcutaneously in female NCr nude mice (Taconic Laboratories), 3 x 10⁶ cells/mouse. Nine days after cell implantation, mice were randomized into 13 groups (8 mice/group, average tumor volume 185 mm³, range 127-327 mm³), and were dosed orally with either vehicle, Example 32 as a single agent, osimertinib as a single agent, or Example 32 in combination with osimertinib, at the different doses and schedules of Example 32 indicated in Figures 31A-31E, for 19 to 29 days (better responding groups were dosed longer). Tumor length and width were measured by caliper, and tumor volume calculated using the formula volume = (length x width²)* π /6. Example 32 was formulated in water, adjusted to pH 2 with methane sulfonic acid. Osimertinib was formulated in 0.5% HPMC in water. All formulations were administered by oral gavage, at a volume of 5 ml/kg. Osimertinib was dosed QD (AM), and Example 32 was dosed BID (AM and PM, 8 hours apart). In the groups that were dosed with Example 32 in combination with osimertinib, the AM dose of osimertinib was administered 3 hours before Example 32 to minimize exposure interactions.

RESULTS: The addition of Example 32 enhanced the antitumor activity of osimertinib, compared to treatment with osimertinib alone. Example 32 administered as a single agent had no significant activity relative to vehicle control treatment (Figure 31A-31E). For the combination treatment of Example 32 and osimertinib, enhancement of antitumor activity was greater at the higher dose of Example 32 (50 mg/kg BID vs. 25 mg/kg BID). Although the enhancement of osimertinib activity decreased with decreasing schedule intensity of Example 32 (daily > 4on/3off > 7on/7off > 2on/5off > just first 7 days), the difference was statistically significant after 26 days of treatment at all schedules of Example 32, with the exception of the groups in which Example 32 was dosed only for the first 7 days (Table 32). All treatments were well tolerated, with no significant weight loss (Figure 32) or other outward signs observed over the course of treatment.

Table 32. p values For Antitumor Activity of AZD9291 Plus JAK1 Inhibitor combinations, After 26 Days of Treatment

Treatment	vs. osimertinib single agent ^a
osimertinib, 2.5 mg/kg QD	---
osimertinib, 2.5 mg/kg QD + Example 32, 12.5 mg/kg BID, daily	0.0156
osimertinib, 2.5 mg/kg QD + Example 32, 50 mg/kg BID, daily	< 0.0001
osimertinib, 2.5 mg/kg QD + Example 32, 25 mg/kg BID, 4on/3off	0.0002
osimertinib, 2.5 mg/kg QD + Example 32, 50 mg/kg BID, 4on/3off	< 0.0001
osimertinib, 2.5 mg/kg QD + Example 32, 25 mg/kg BID, 7on/7off	0.0079
osimertinib, 2.5 mg/kg QD + Example 32, 50 mg/kg BID, 7on/7off	0.0002
osimertinib, 2.5 mg/kg QD + Example 32, 25 mg/kg BID, 2on/5off	0.0391
osimertinib, 2.5 mg/kg QD + Example 32, 50 mg/kg BID, 2on/5off	0.0010
osimertinib, 2.5 mg/kg QD + Example 32, 25 mg/kg BID, just first 7 days	0.4744
osimertinib, 2.5 mg/kg QD + Example 32, 50 mg/kg BID, just first 7 days	0.4101

^aTwo-sided test

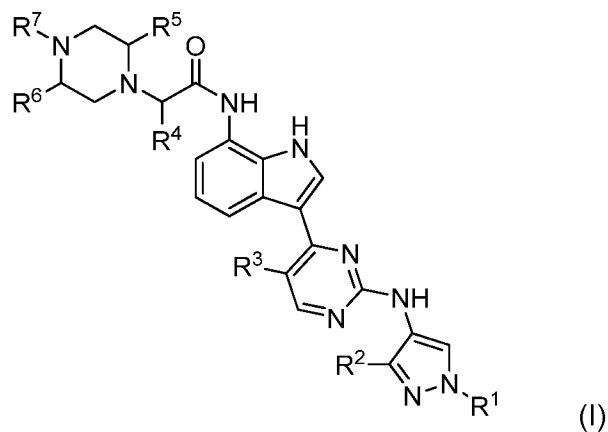
CONCLUSION: The increased antitumor activity of Example 32 in combination with osimertinib compared to single agent Example 32 is consistent with a role for STAT3 signaling in escape from, or resistance to, EGFR inhibition in this model of non-small cell lung cancer.

The results support the hypothesis that inhibition of STAT3 signaling can enhance the antitumor activity of an EGFR inhibitor in T790M EGFR mutant NSCLC. The significant combination activity observed even when Example 32 was administered as infrequently as 2on/5off (*i.e.*, only days 1 and 2 of a weekly cycle), suggests that enhancement of osimertinib activity may be

5 achievable with only intermittent inhibition of pSTAT3 signaling.

Claims

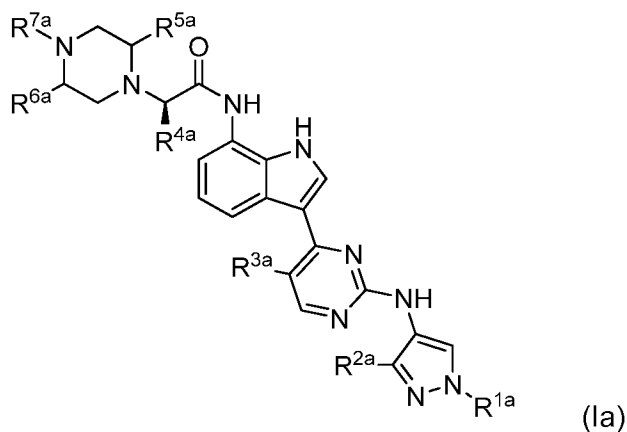
1. A compound of formula (I):



wherein

- 5 R^1 is methyl or ethyl;
 R^2 is selected from methyl, ethyl, methoxy and ethoxy;
 R^3 is selected from hydrogen, chlorine, fluorine, bromine and methyl;
 R^4 is selected from methyl, ethyl and $-\text{CH}_2\text{OCH}_3$;
 R^5 and R^6 are each individually methyl or hydrogen; and
 10 R^7 is selected from methyl, ethyl, $-(\text{CH}_2)_2\text{OH}$ and $-(\text{CH}_2)_2\text{OCH}_3$, or a pharmaceutically acceptable salt thereof.

2. A compound according to claim 1, having the structure of formula (Ia):



- 15 wherein
 R^{1a} is methyl or ethyl;
 R^{2a} is selected from methyl, ethyl, methoxy and ethoxy;
 R^{3a} is selected from hydrogen, chlorine, fluorine, bromine and methyl;

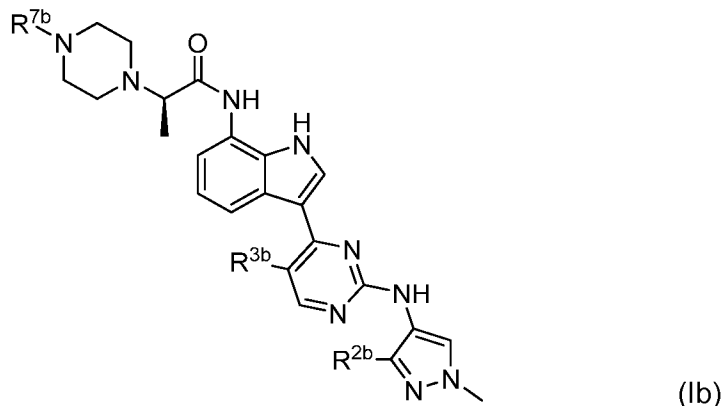
R^{4a} is selected from methyl, ethyl and -CH₂OCH₃;

R^{5a} and R^{6a} are each individually methyl or hydrogen; and

R^{7a} is selected from methyl, ethyl, -(CH₂)₂OH and -(CH₂)₂OCH₃, or a pharmaceutically acceptable salt thereof.

5

3. A compound according to claim 1 having the structure of formula (Ib):



wherein

R^{2b} is selected from methyl, ethyl, methoxy and ethoxy;

10 R^{3b} is selected from hydrogen, chlorine, fluorine, bromine and methyl; and

R^{7b} is selected from methyl, ethyl, -(CH₂)₂OH and -(CH₂)₂OCH₃, or a pharmaceutically acceptable salt thereof.

4. A compound according to any one of claims 1-3 selected from:

15 (2*R*)-2-[(2*S*)-2,4-dimethylpiperazin-1-yl]-*N*-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)propanamide;

(2*R*)-2-[(3*R*)-3,4-dimethylpiperazin-1-yl]-*N*-(3-{5-fluoro-2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)propanamide;

20 (2*R*)-*N*-(3-{5-fluoro-2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-[4-(2-methoxyethyl)piperazin-1-yl]propanamide;

(2*R*)-2-(4-ethylpiperazin-1-yl)-*N*-(3-{5-fluoro-2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)propanamide;

(2*R*)-2-[(3*S*)-3,4-dimethylpiperazin-1-yl]-*N*-(3-{5-fluoro-2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)propanamide;

25 (2*R*)-2-[(2*R*)-2,4-dimethylpiperazin-1-yl]-*N*-(3-{5-fluoro-2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)propanamide;

(2*R*)-2-[(2*S*)-2,4-dimethylpiperazin-1-yl]-*N*-(3-{5-fluoro-2-[(3-methoxy-1-methyl-1*H*-pyrazol-

4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)propanamide;

(2*S*)-2-[(2*S*)-2,4-dimethylpiperazin-1-yl]-*N*-(3-{5-fluoro-2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)propanamide;

(2*R*)-*N*-(3-{2-[(3-ethoxy-1-methyl-1*H*-pyrazol-4-yl)amino]-5-fluoropyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-ethylpiperazin-1-yl)propanamide;

(2*R*)-2-[(2*S*)-2,4-dimethylpiperazin-1-yl]-*N*-(3-{2-[(3-ethoxy-1-methyl-1*H*-pyrazol-4-yl)amino]-5-fluoropyrimidin-4-yl}-1*H*-indol-7-yl)propanamide;

(2*S*)-2-[(2*S*)-2,4-dimethylpiperazin-1-yl]-*N*-(3-{2-[(3-ethoxy-1-methyl-1*H*-pyrazol-4-yl)amino]-5-fluoropyrimidin-4-yl}-1*H*-indol-7-yl)propanamide;

(2*R*)-2-[(3*S*)-3,4-dimethylpiperazin-1-yl]-*N*-(3-{2-[(3-ethoxy-1-methyl-1*H*-pyrazol-4-yl)amino]-5-fluoropyrimidin-4-yl}-1*H*-indol-7-yl)propanamide;

(2*S*)-2-[(3*S*)-3,4-dimethylpiperazin-1-yl]-*N*-(3-{2-[(3-ethoxy-1-methyl-1*H*-pyrazol-4-yl)amino]-5-fluoropyrimidin-4-yl}-1*H*-indol-7-yl)propanamide;

(2*R*)-2-[(2*R*)-2,4-dimethylpiperazin-1-yl]-*N*-(3-{2-[(3-ethoxy-1-methyl-1*H*-pyrazol-4-yl)amino]-5-fluoropyrimidin-4-yl}-1*H*-indol-7-yl)propanamide;

(2*R*)-2-[(3*R*)-3,4-dimethylpiperazin-1-yl]-*N*-(3-{2-[(3-ethoxy-1-methyl-1*H*-pyrazol-4-yl)amino]-5-fluoropyrimidin-4-yl}-1*H*-indol-7-yl)propanamide;

(2*R*)-2-[(3*S*)-3,4-dimethylpiperazin-1-yl]-*N*-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1*H*-indol-7-yl)propanamide;

(2*R*)-2-[4-(2-hydroxyethyl)piperazin-1-yl]-*N*-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1*H*-indol-7-yl)propanamide;

(2*R*)-2-(4-ethylpiperazin-1-yl)-*N*-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1*H*-indol-7-yl)propanamide;

(2*R*)-2-[4-(2-methoxyethyl)piperazin-1-yl]-*N*-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1*H*-indol-7-yl)propanamide;

(2*R*)-2-[(3*R*)-3,4-dimethylpiperazin-1-yl]-*N*-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1*H*-indol-7-yl)propanamide;

(2*R*)-2-[(2*S*)-2,4-dimethylpiperazin-1-yl]-*N*-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1*H*-indol-7-yl)propanamide;

(2*R*)-*N*-(3-{2-[(3-ethoxy-1-methyl-1*H*-pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-ethylpiperazin-1-yl)propanamide;

(2*R*)-*N*-(3-{2-[(3-ethoxy-1-methyl-1*H*-pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1*H*-indol-7-yl)-2-[4-(2-methoxyethyl)piperazin-1-yl]propanamide;

(2*R*)-2-[(2*S*)-2,4-dimethylpiperazin-1-yl]-*N*-(3-{2-[(3-ethoxy-1-methyl-1*H*-pyrazol-4-

yl)amino]-5-methylpyrimidin-4-yl}-1*H*-indol-7-yl)propanamide;

(2*R*)-2-[(2*R*)-2,4-dimethylpiperazin-1-yl]-*N*-(3-{2-[(3-ethoxy-1-methyl-1*H*-pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1*H*-indol-7-yl)propanamide;

(2*R*)-2-[(3*R*)-3,4-dimethylpiperazin-1-yl]-*N*-(3-{2-[(3-ethoxy-1-methyl-1*H*-pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1*H*-indol-7-yl)propanamide;

(2*R*)-*N*-(3-{5-Fluoro-2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide;

(2*R*)-2-[(3*S*)-3,4-dimethylpiperazin-1-yl]-*N*-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)propanamide;

(2*R*)-2-[4-(2-methoxyethyl)piperazin-1-yl]-*N*-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)propanamide;

(2*R*)-2-(4-ethylpiperazin-1-yl)-*N*-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)propanamide;

(2*R*)-2-[(2*R*)-2,4-dimethylpiperazin-1-yl]-*N*-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)propanamide;

(2*R*)-2-[(3*R*)-3,4-dimethylpiperazin-1-yl]-*N*-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)propanamide;

(2*R*)-2-[(3*S*)-3,4-dimethylpiperazin-1-yl]-*N*-(3-{2-[(3-ethoxy-1-methyl-1*H*-pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1*H*-indol-7-yl)propanamide;

(2*S*)-2-[(3*S*)-3,4-dimethylpiperazin-1-yl]-*N*-(3-{2-[(3-ethoxy-1-methyl-1*H*-pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1*H*-indol-7-yl)propanamide;

(2*R*)-*N*-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide;

(2*S*)-*N*-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide;

(2*R*)-*N*-(3-{2-[(3-Methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide;

(2*R*)-*N*-(3-{2-[(3-ethoxy-1-methyl-1*H*-pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide;

(2*R*)-*N*-(3-{5-chloro-2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide;

(2*R*)-*N*-(3-{5-bromo-2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide;

(2*R*)-2-[(2*R*)-2,4-dimethylpiperazin-1-yl]-*N*-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-

yl)amino]-5-methylpyrimidin-4-yl}-1*H*-indol-7-yl)propanamide;

(2*R*)-*N*-(3-{2-[(3-ethoxy-1-methyl-1*H*-pyrazol-4-yl)amino]-5-fluoropyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide;

(2*R*)-*N*-(3-{5-chloro-2-[(3-ethoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide;

(2*R*)-*N*-(3-{2-[(1,3-dimethyl-1*H*-pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide;

(2*S*)-*N*-(3-{2-[(1,3-dimethyl-1*H*-pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide;

(2*S*)-*N*-(3-{2-[(3-Methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)butanamide;

(2*R*)-*N*-(3-{2-[(3-Methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)butanamide;

(2*R*)-3-methoxy-*N*-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide;

(2*S*)-3-methoxy-*N*-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide;

(2*R*)-*N*-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)butanamide;

(2*S*)-*N*-(3-{2-[(3-Methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)butanamide;

(2*R*)-*N*-(3-{2-[(1,3-dimethyl-1*H*-pyrazol-4-yl)amino]-5-fluoropyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)butanamide;

(2*S*)-*N*-(3-{2-[(1,3-dimethyl-1*H*-pyrazol-4-yl)amino]-5-fluoropyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)butanamide;

(2*R*)-*N*-(3-{5-fluoro-2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)butanamide;

(2*S*)-*N*-(3-{5-fluoro-2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)butanamide;

(2*R*)-*N*-(3-{2-[(1,3-dimethyl-1*H*-pyrazol-4-yl)amino]-5-fluoropyrimidin-4-yl}-1*H*-indol-7-yl)-3-methoxy-2-(4-methylpiperazin-1-yl)propanamide;

(2*S*)-*N*-(3-{2-[(1,3-dimethyl-1*H*-pyrazol-4-yl)amino]-5-fluoropyrimidin-4-yl}-1*H*-indol-7-yl)-3-methoxy-2-(4-methylpiperazin-1-yl)propanamide;

(2*R*)-*N*-(3-{2-[(3-ethyl-1-methyl-1*H*-pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1*H*-indol-7-

yl)-2-(4-methylpiperazin-1-yl)propanamide;

(2*R*)-*N*-(3-{2-[(1,3-dimethyl-1*H*-pyrazol-4-yl)amino]-5-fluoropyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide;

(2*R*)-*N*-(3-{2-[(1,3-dimethyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide;

(2*R*)-*N*-(3-{2-[(1-ethyl-3-methoxy-1*H*-pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide;

(2*R*)-*N*-(3-{2-[(3-ethoxy-1-ethyl-1*H*-pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide;

(2*R*)-3-Methoxy-*N*-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide;

(2*S*)-3-Methoxy-*N*-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide;

(2*R*)-*N*-(3-{2-[(3-ethoxy-1-methyl-1*H*-pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)butanamide;

(2*S*)-*N*-(3-{2-[(3-ethoxy-1-methyl-1*H*-pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)butanamide;

(2*R*)-*N*-(3-{2-[(3-ethoxy-1-methyl-1*H*-pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1*H*-indol-7-yl)-3-methoxy-2-(4-methylpiperazin-1-yl)propanamide; and

(2*S*)-*N*-(3-{2-[(3-ethoxy-1-methyl-1*H*-pyrazol-4-yl)amino]-5-methylpyrimidin-4-yl}-1*H*-indol-7-yl)-3-methoxy-2-(4-methylpiperazin-1-yl)propanamide; or a pharmaceutically acceptable salt thereof.

5. The compound of claim 4 which is (2*R*)-*N*-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide, or a pharmaceutically acceptable salt thereof.

6. The compound of claim 5.

7. The pharmaceutically acceptable salt of the compound of claim 5.

8. A solid form of (2*R*)-*N*-(3-{2-[(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)amino]pyrimidin-4-yl}-1*H*-indol-7-yl)-2-(4-methylpiperazin-1-yl)propanamide, or a pharmaceutically acceptable salt thereof.

9. A pharmaceutical composition comprising a compound of any one of claims 1-8, and a pharmaceutically acceptable diluent, excipient or carrier.

5 10. A method of treating a JAK-related disorder in a subject comprising administering to the subject an effective amount of a compound as claimed in any one of claims 1-8, or a pharmaceutically acceptable salt thereof.

10 11. A compound of any one of claims 1-8, or a pharmaceutically acceptable salt thereof, for use in treating a JAK-related disorder.

12. A pharmaceutical composition comprising a compound of any one of claims 1-8, or a pharmaceutically acceptable salt thereof, for use in treating a JAK-related disorder.

15 13. Use of a compound of any one of claims 1-8, or a pharmaceutically acceptable salt thereof, in the manufacture of a medicament for treating a JAK-related disorder.

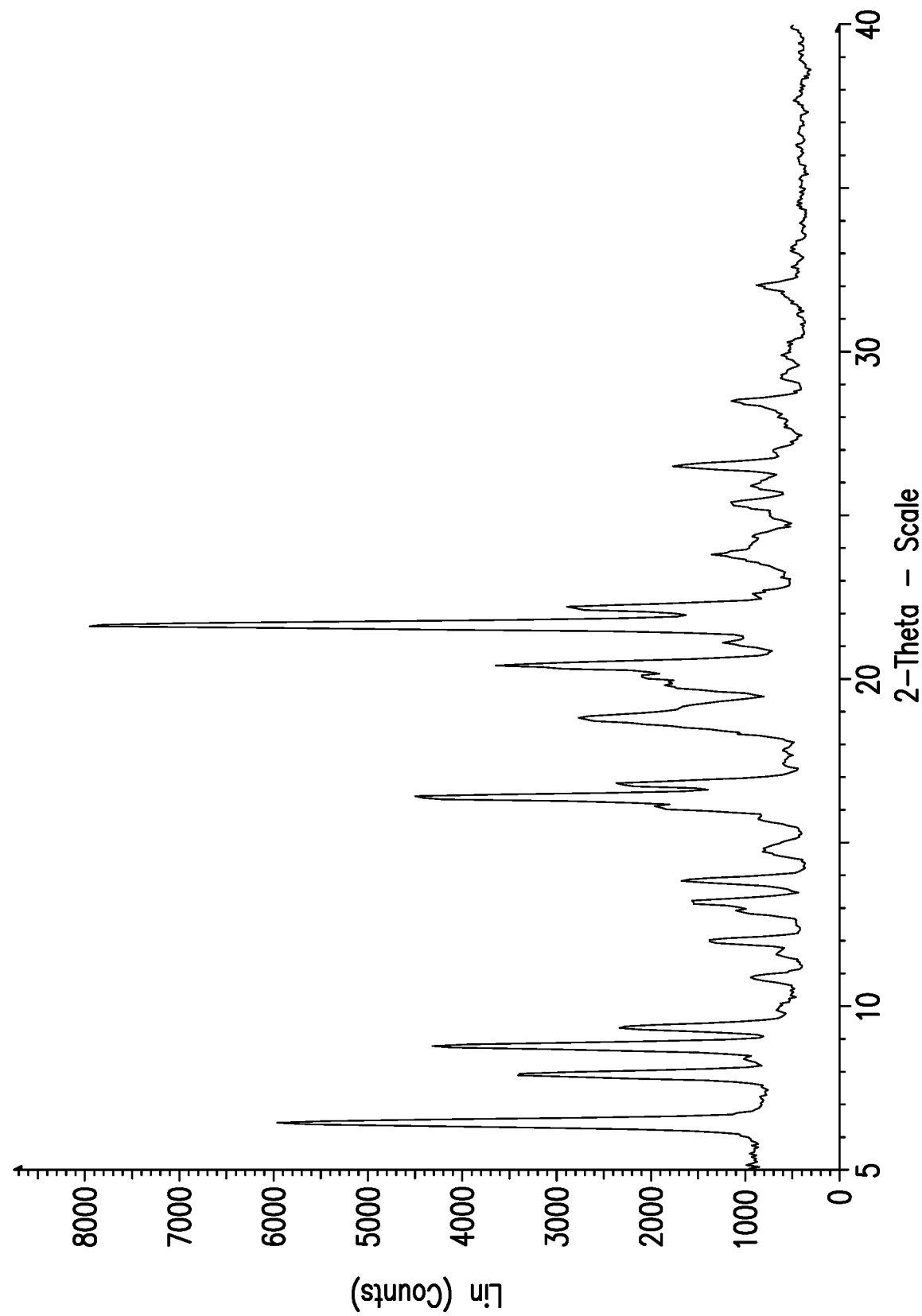


FIG.1

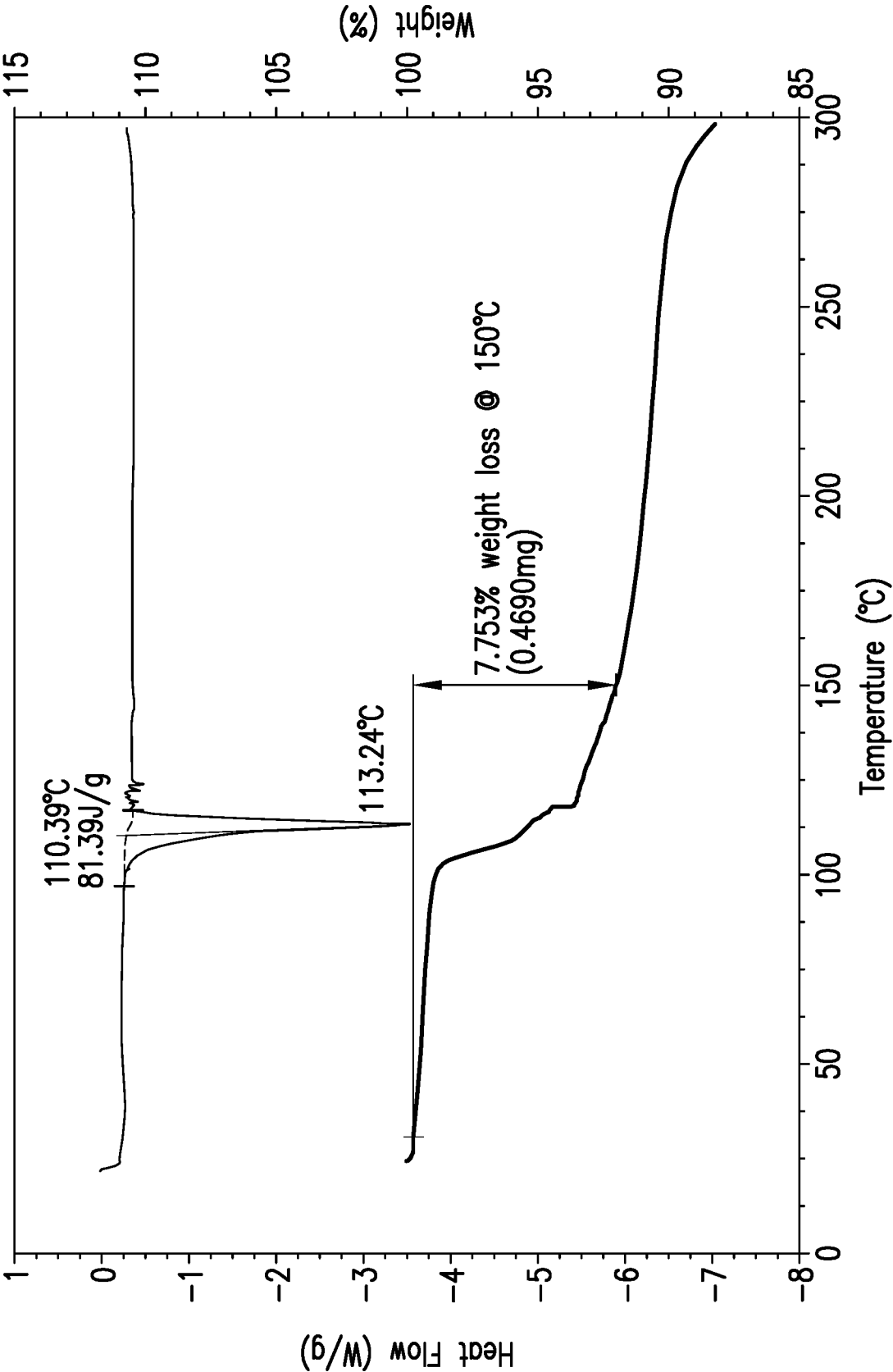


FIG.2

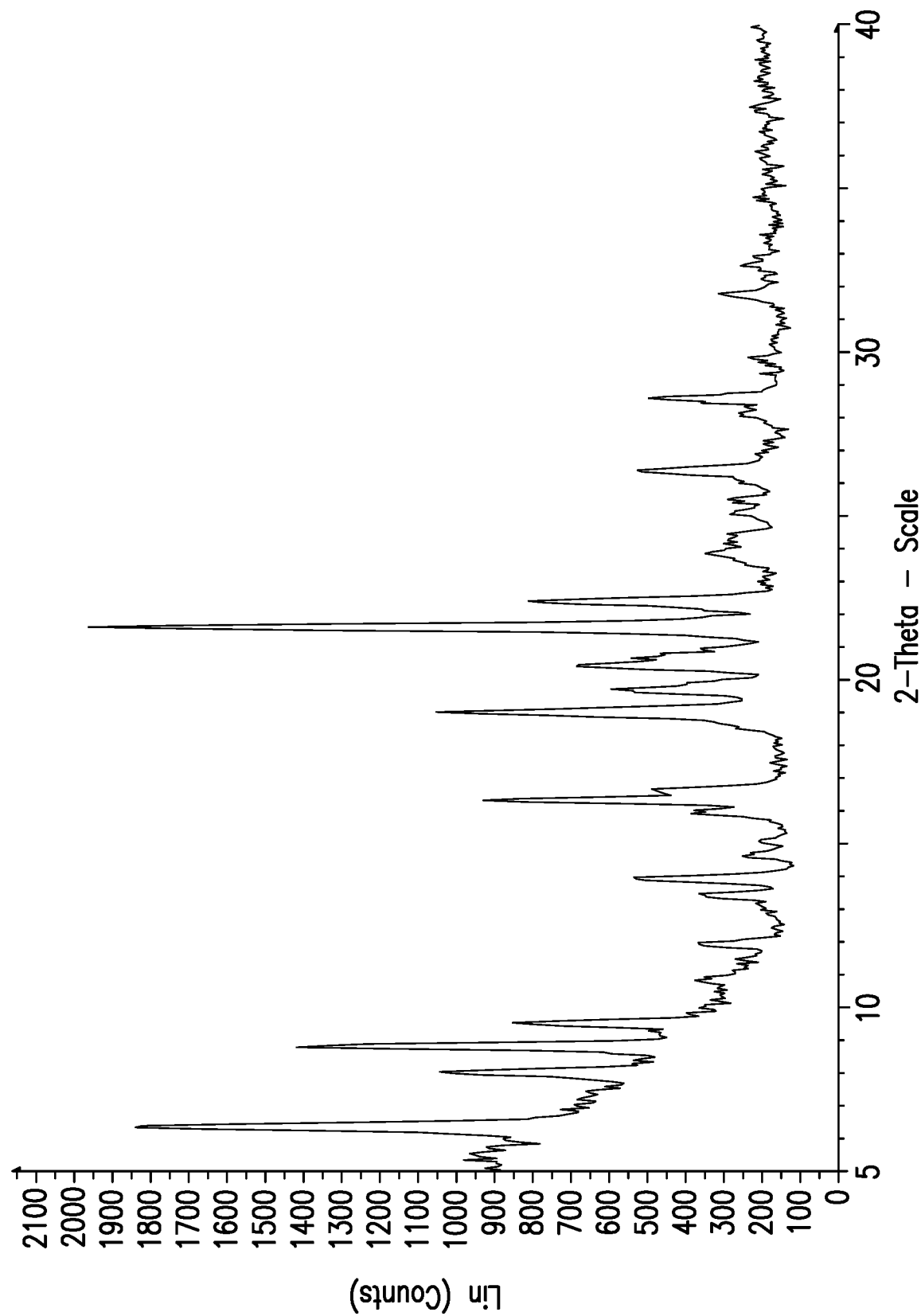


FIG.3

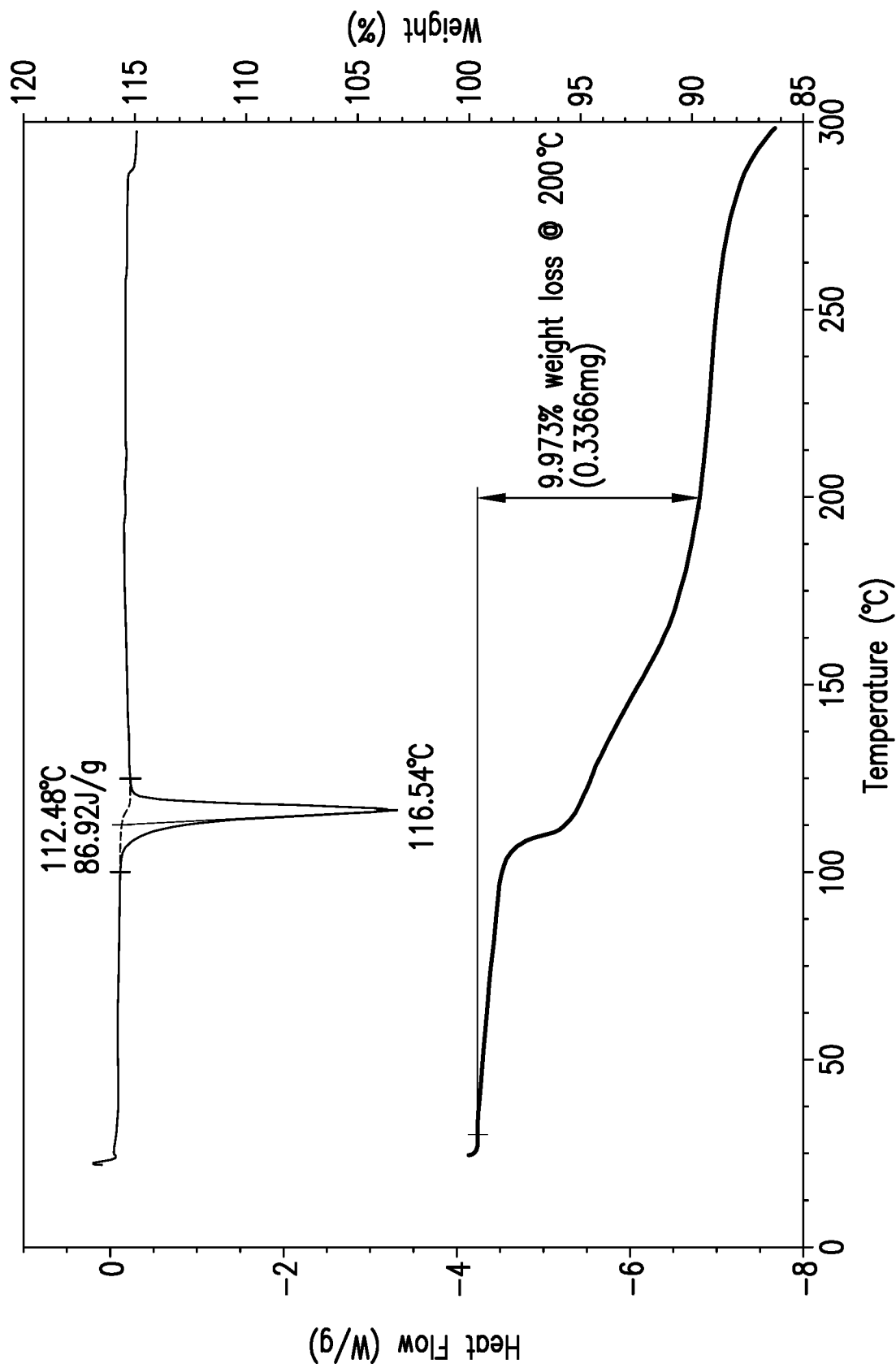


FIG.4

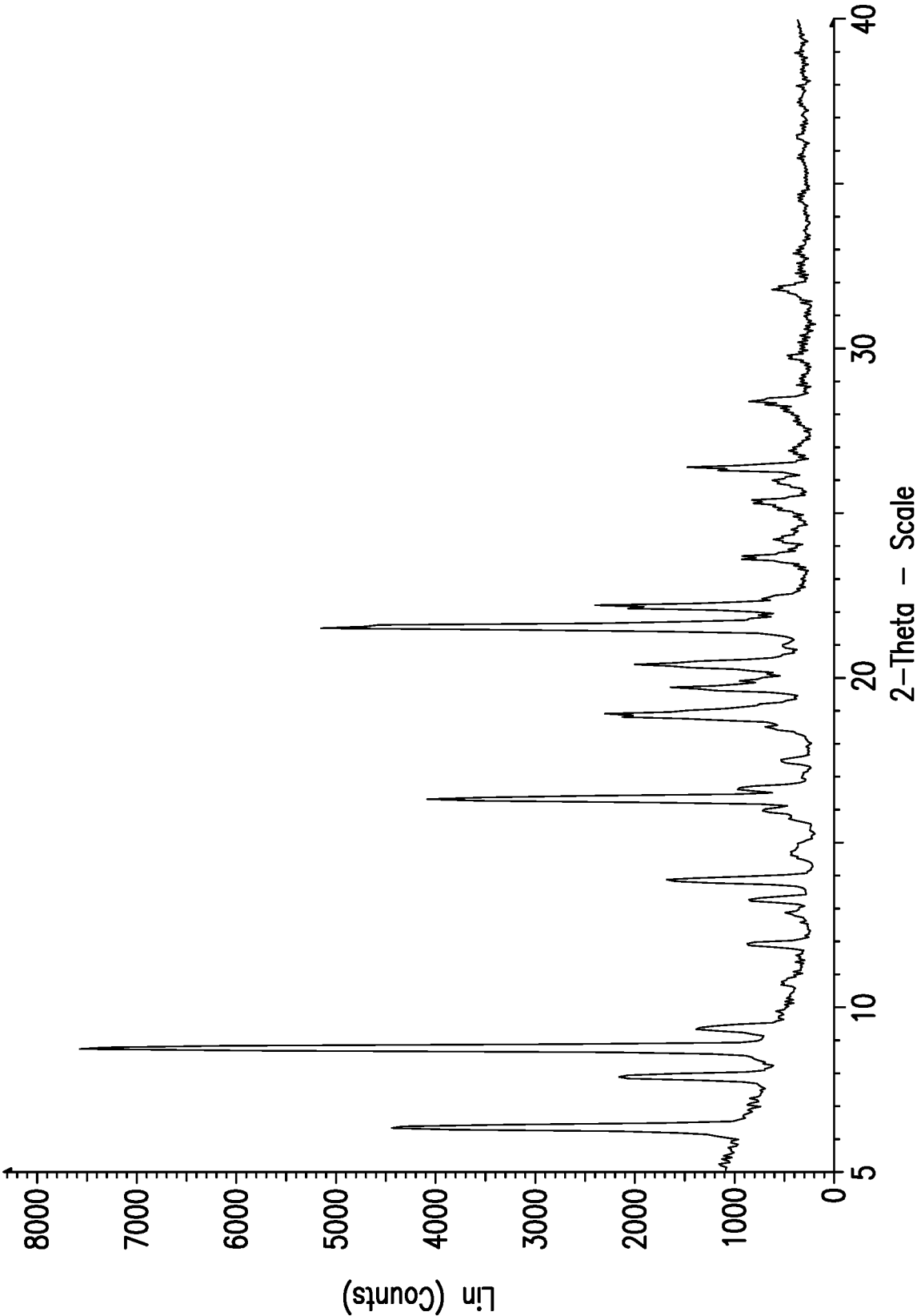


FIG.5

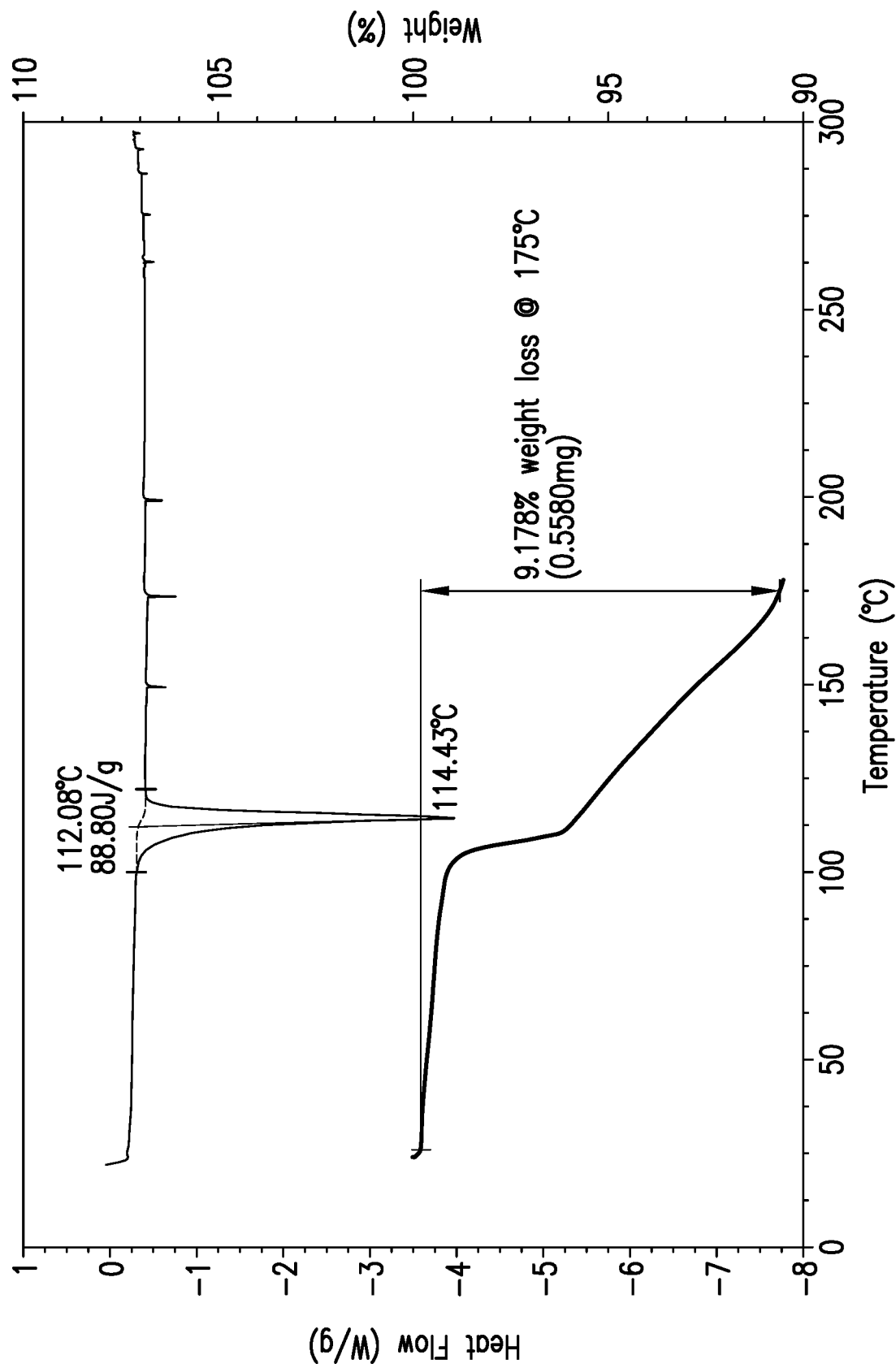
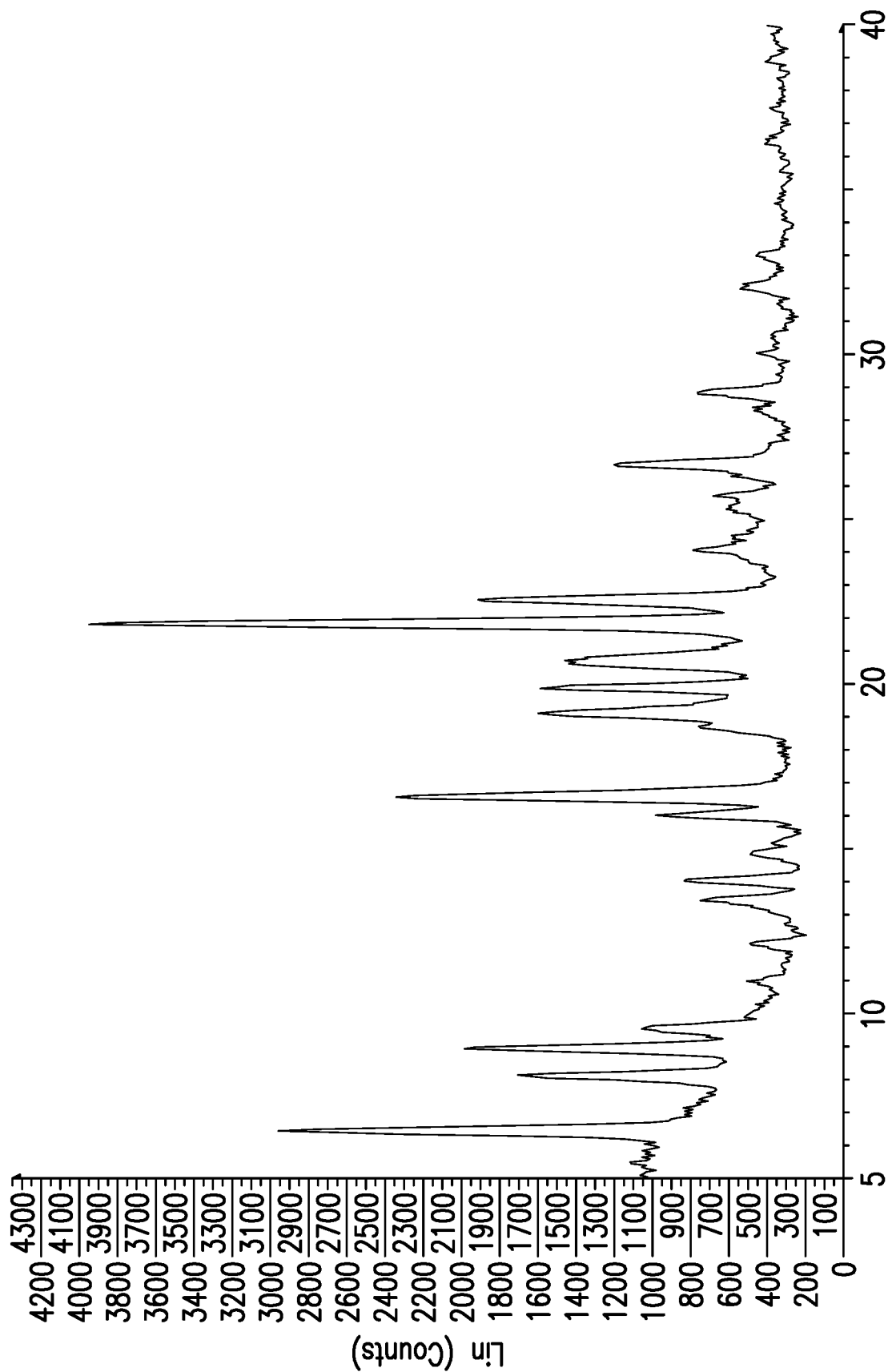


FIG.6



2-Theta - Scale

FIG.7

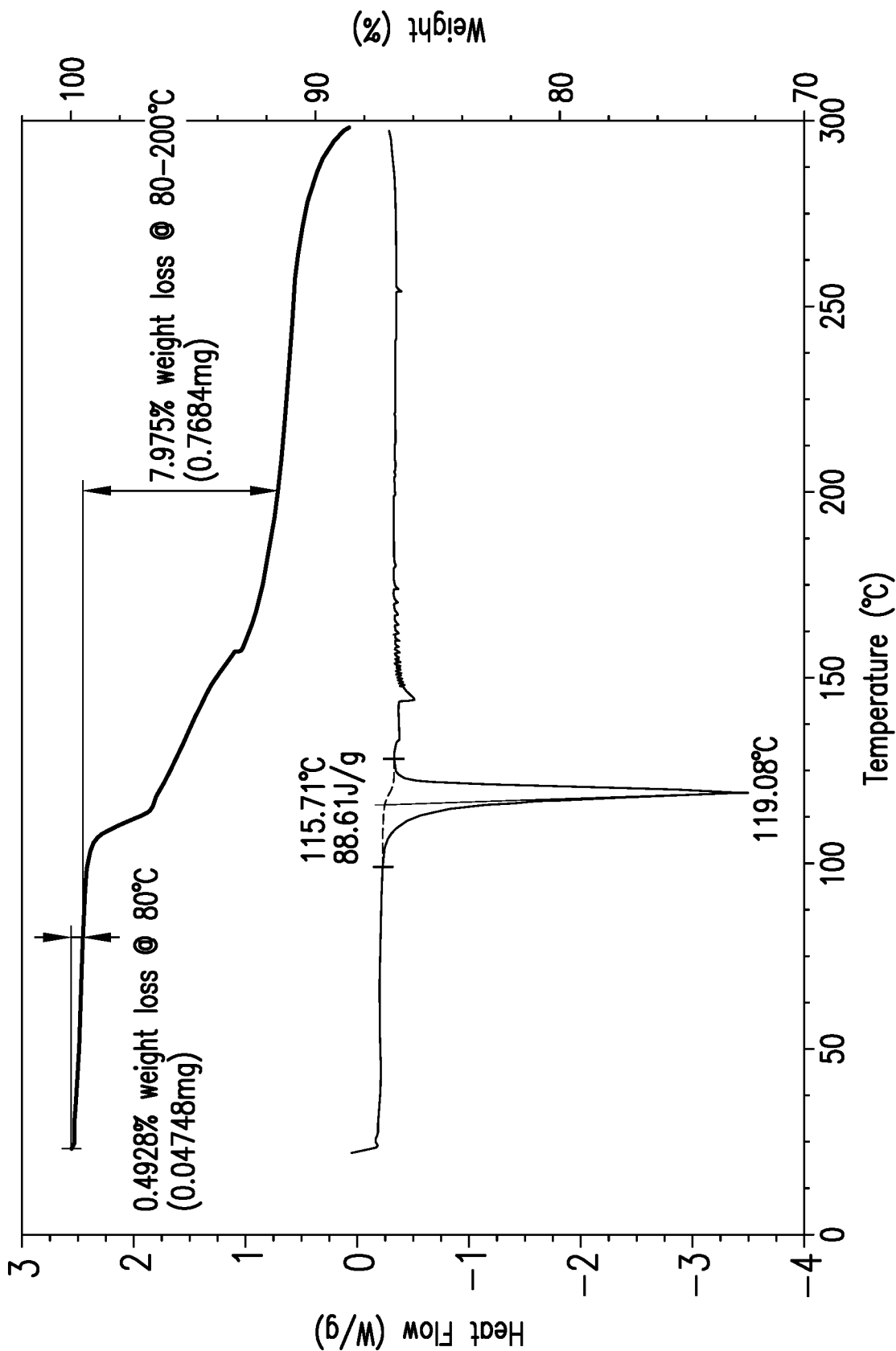


FIG.8

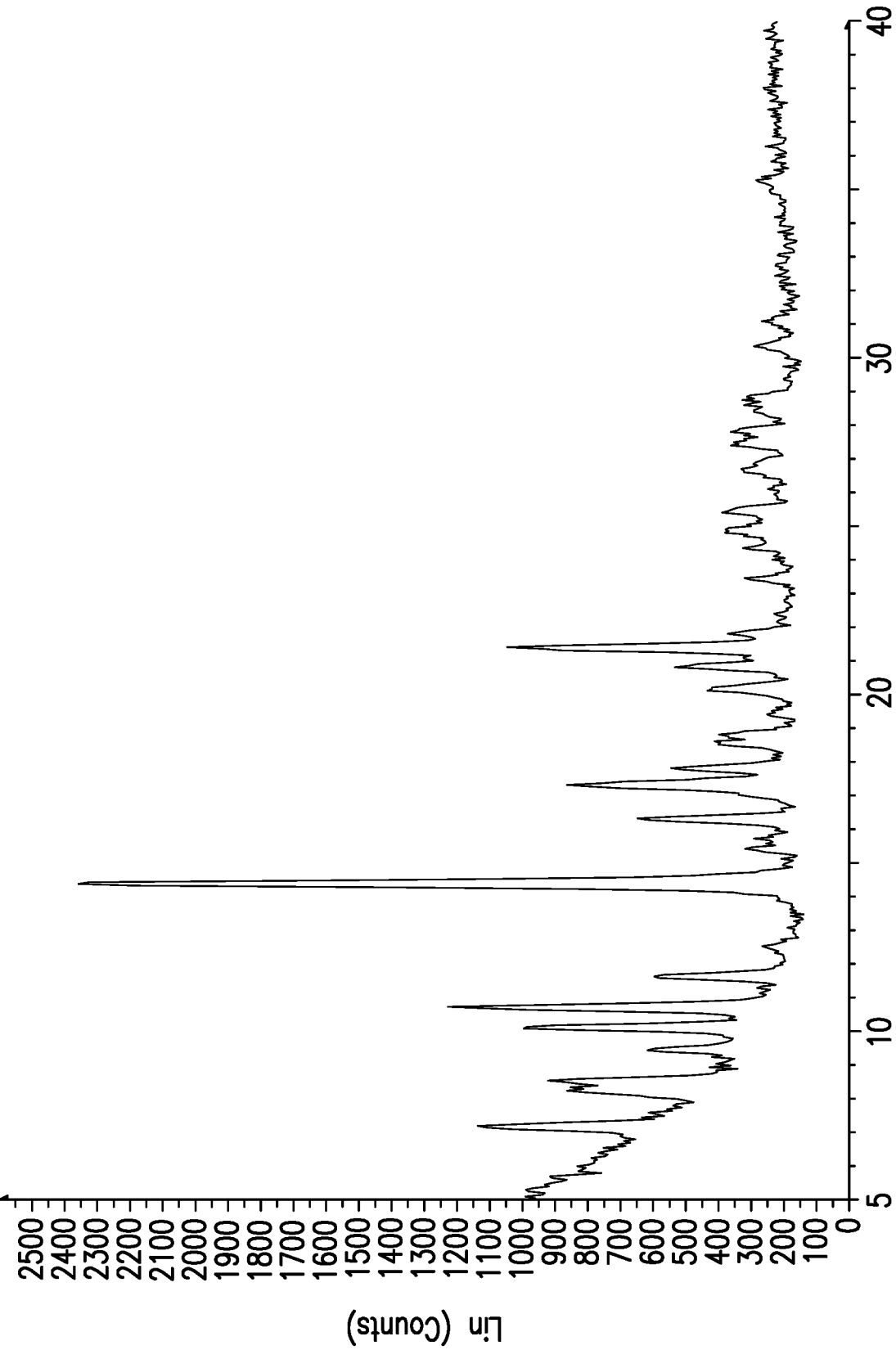


FIG.9

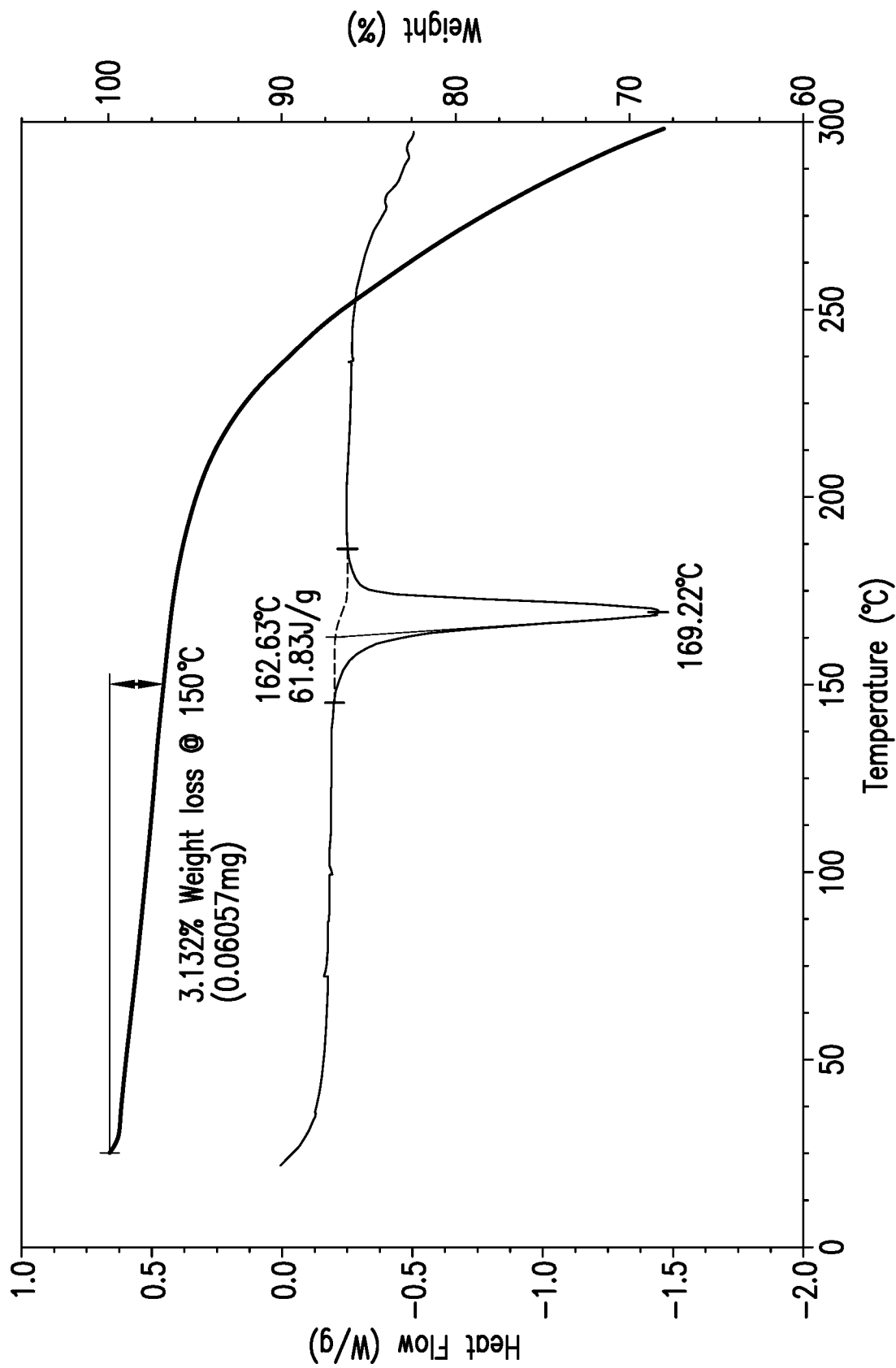
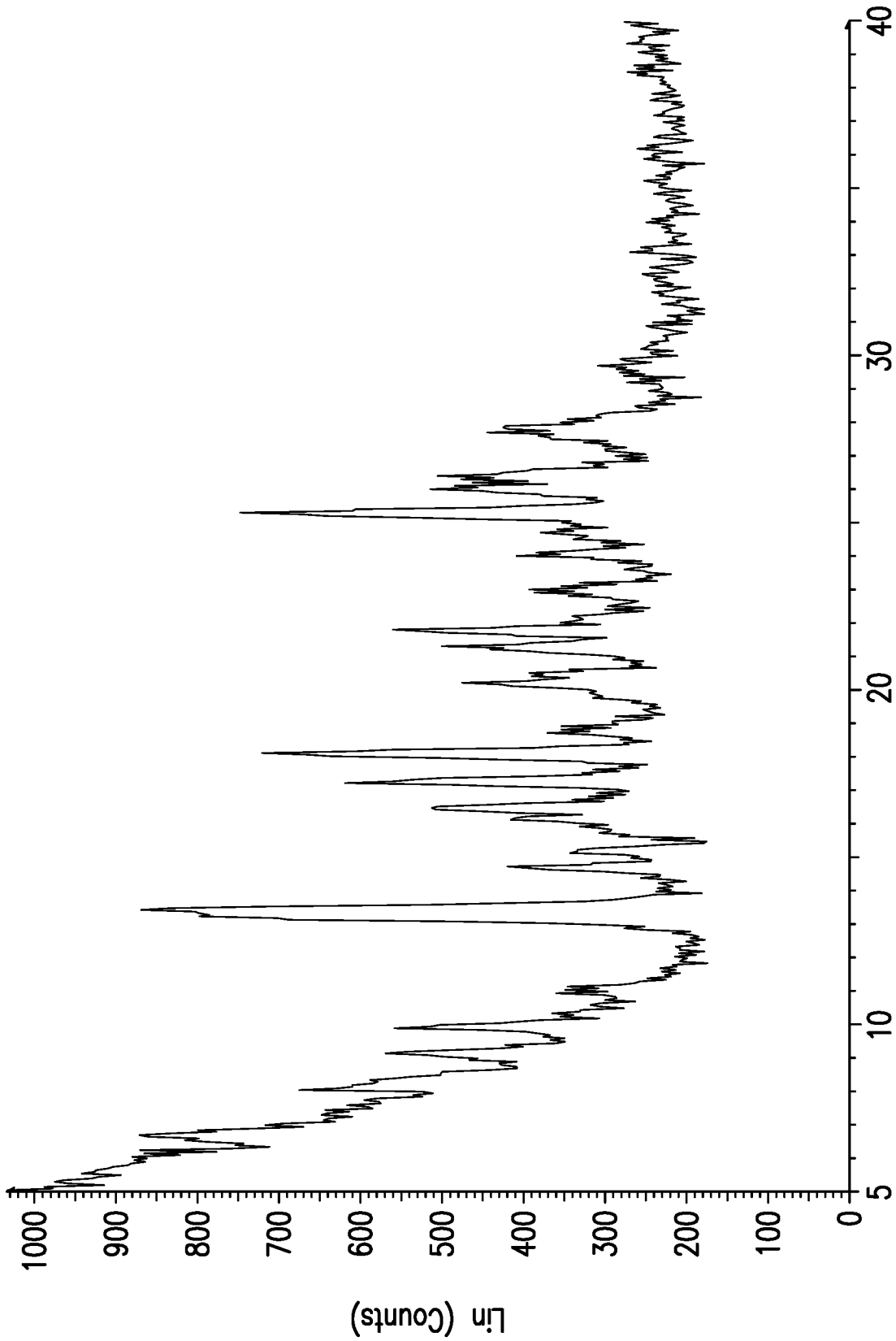


FIG.10



2-Theta - Scale

FIG.11

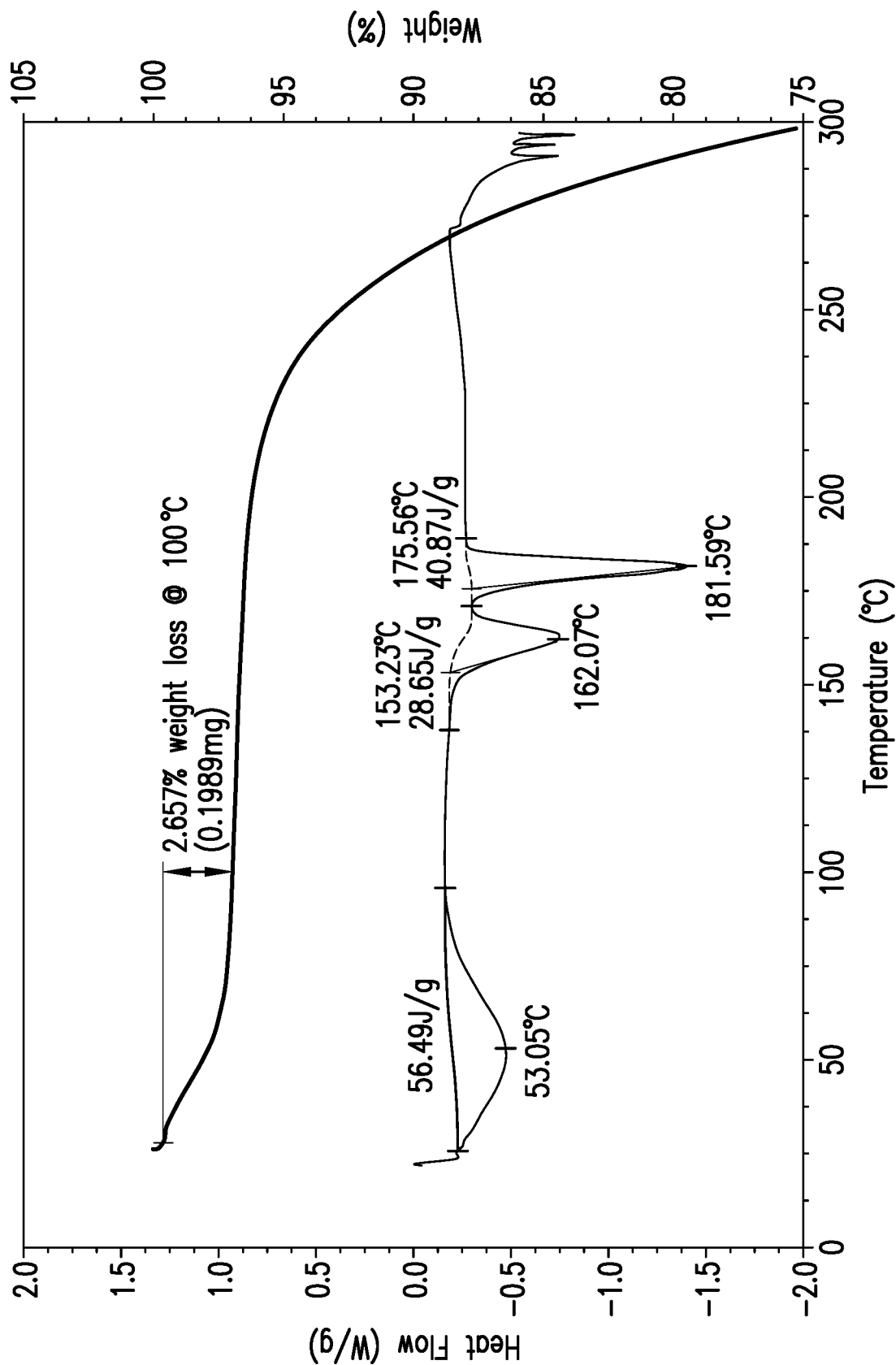
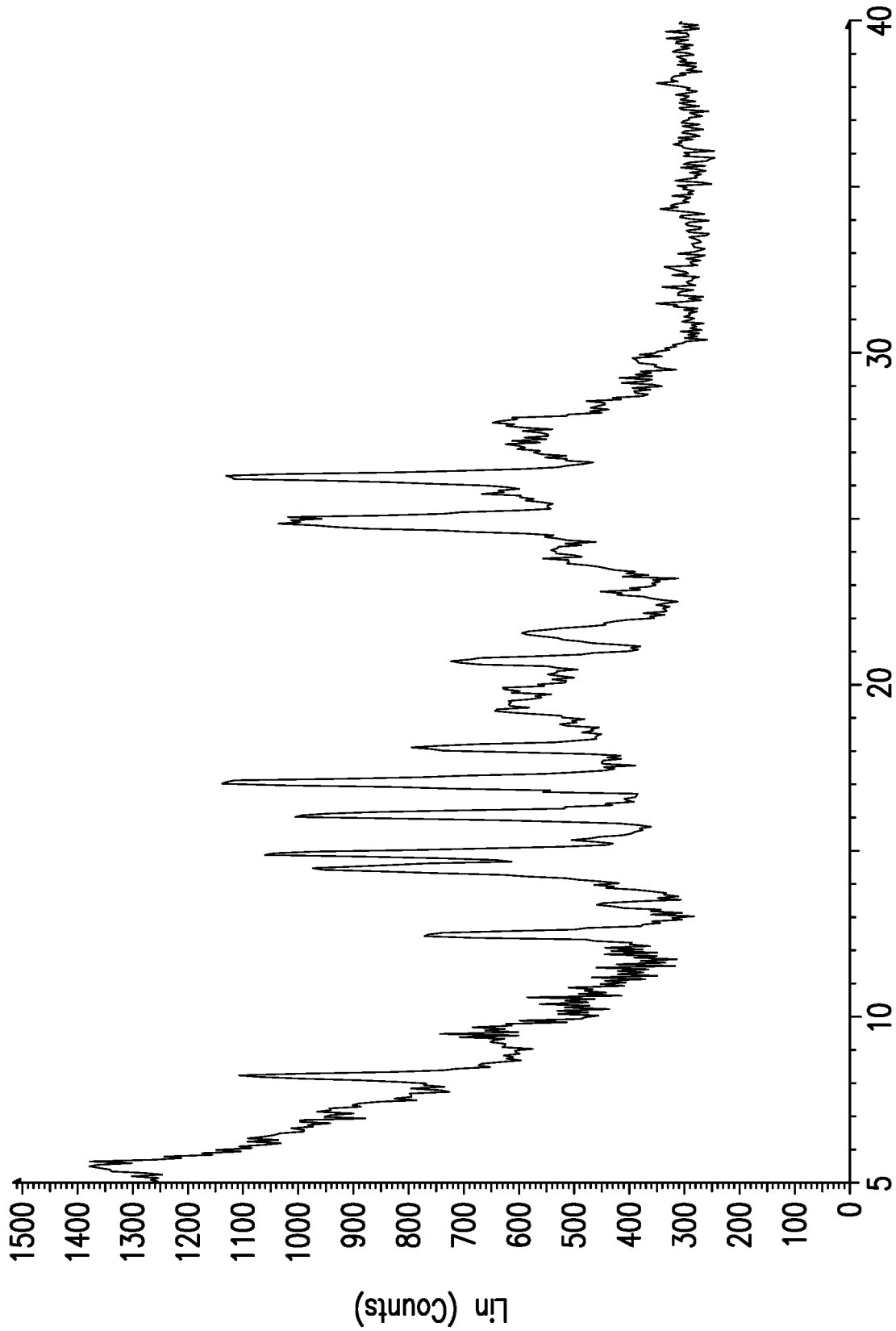
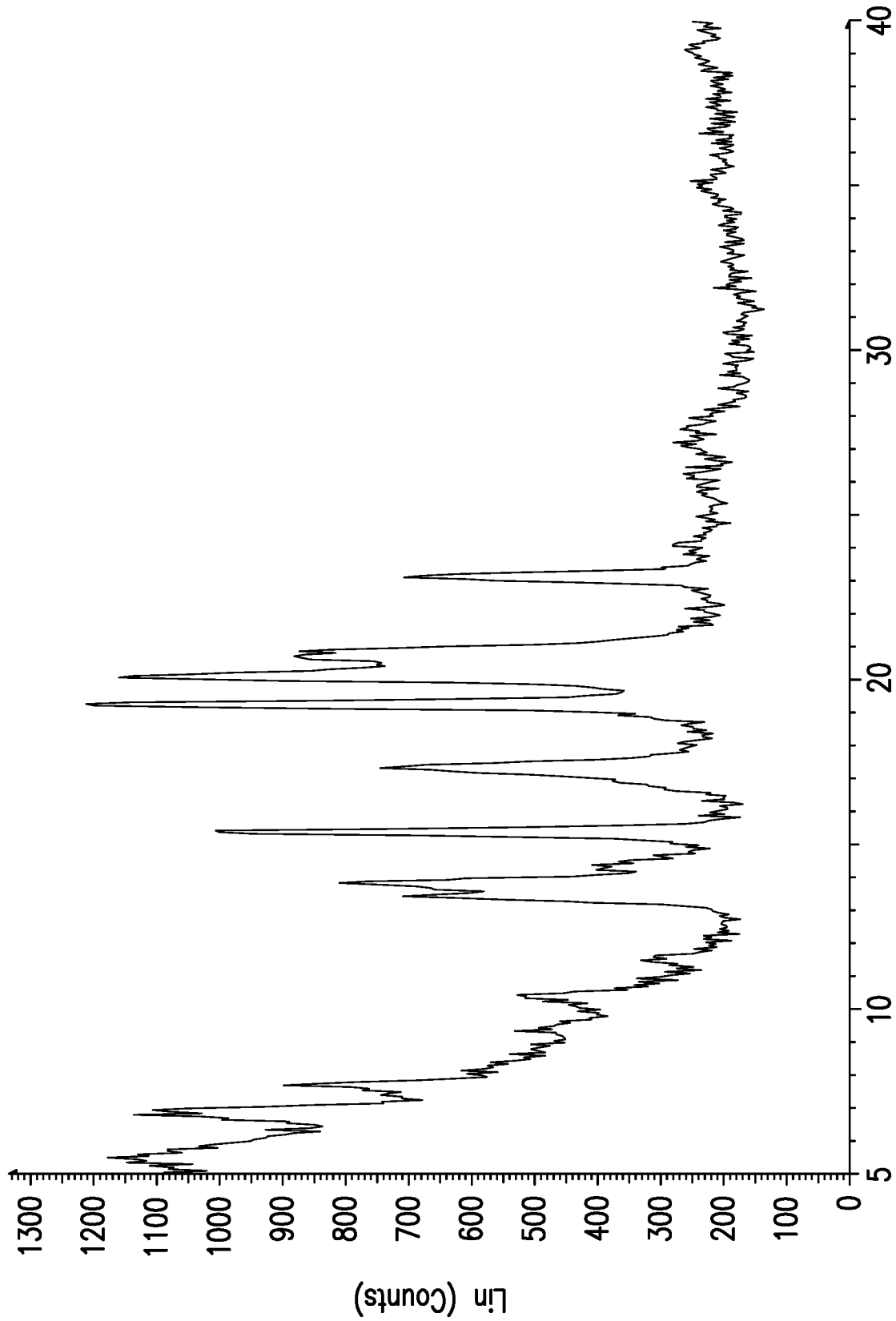


FIG.12



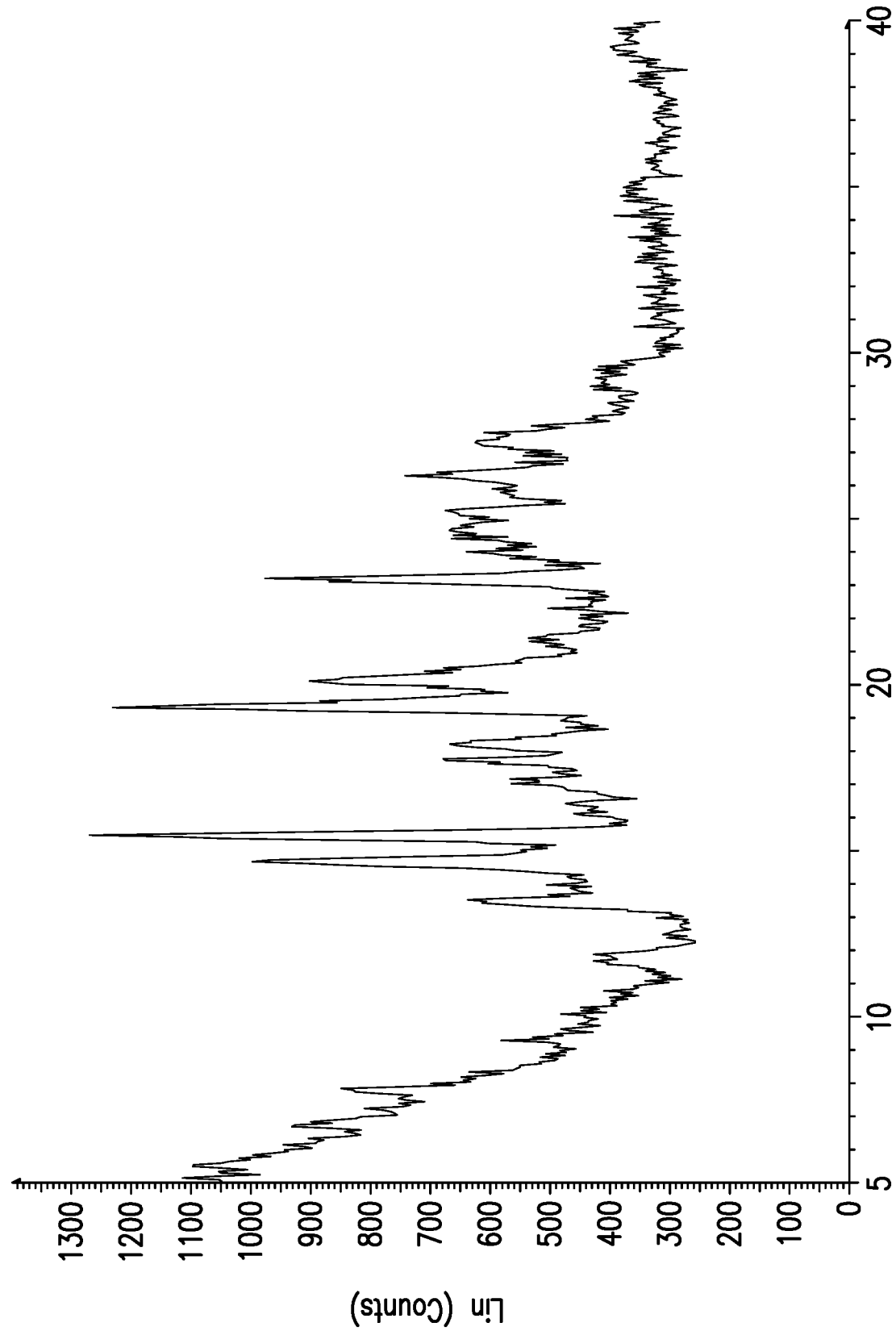
2-Theta - Scale

FIG.13

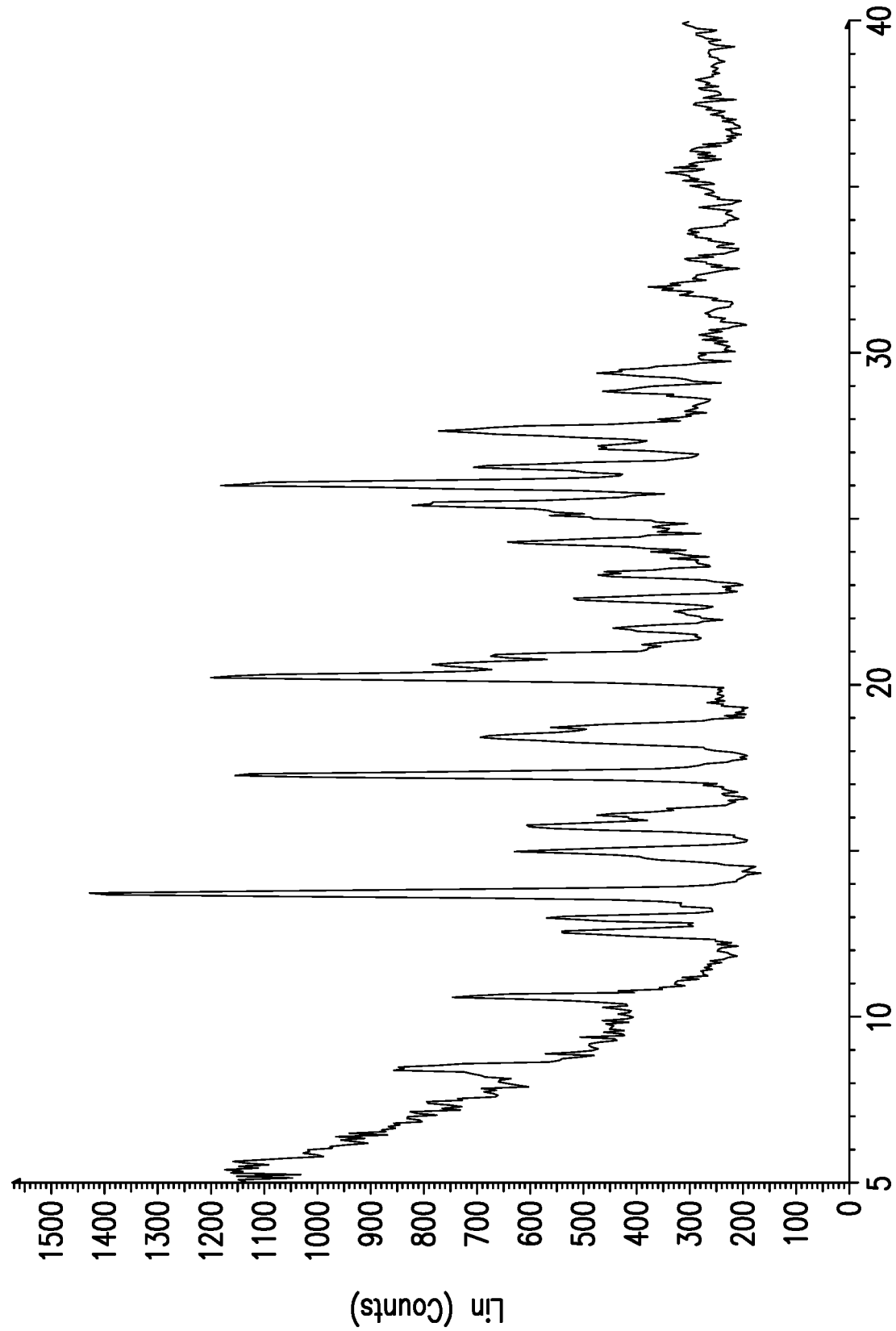


2-Theta - Scale

FIG.14



2-Theta - Scale
FIG.15



2-Theta - Scale

FIG.16

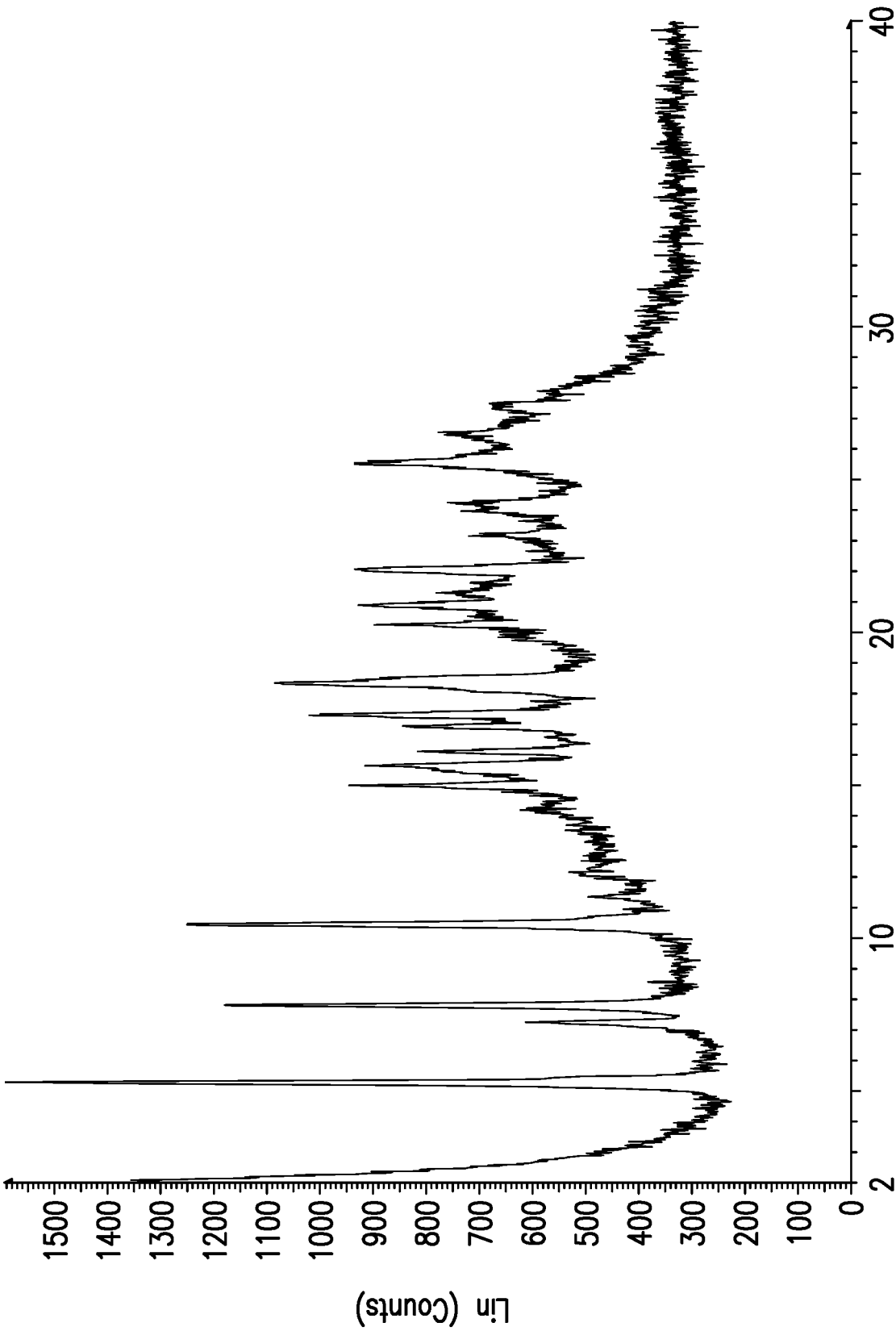


FIG.17

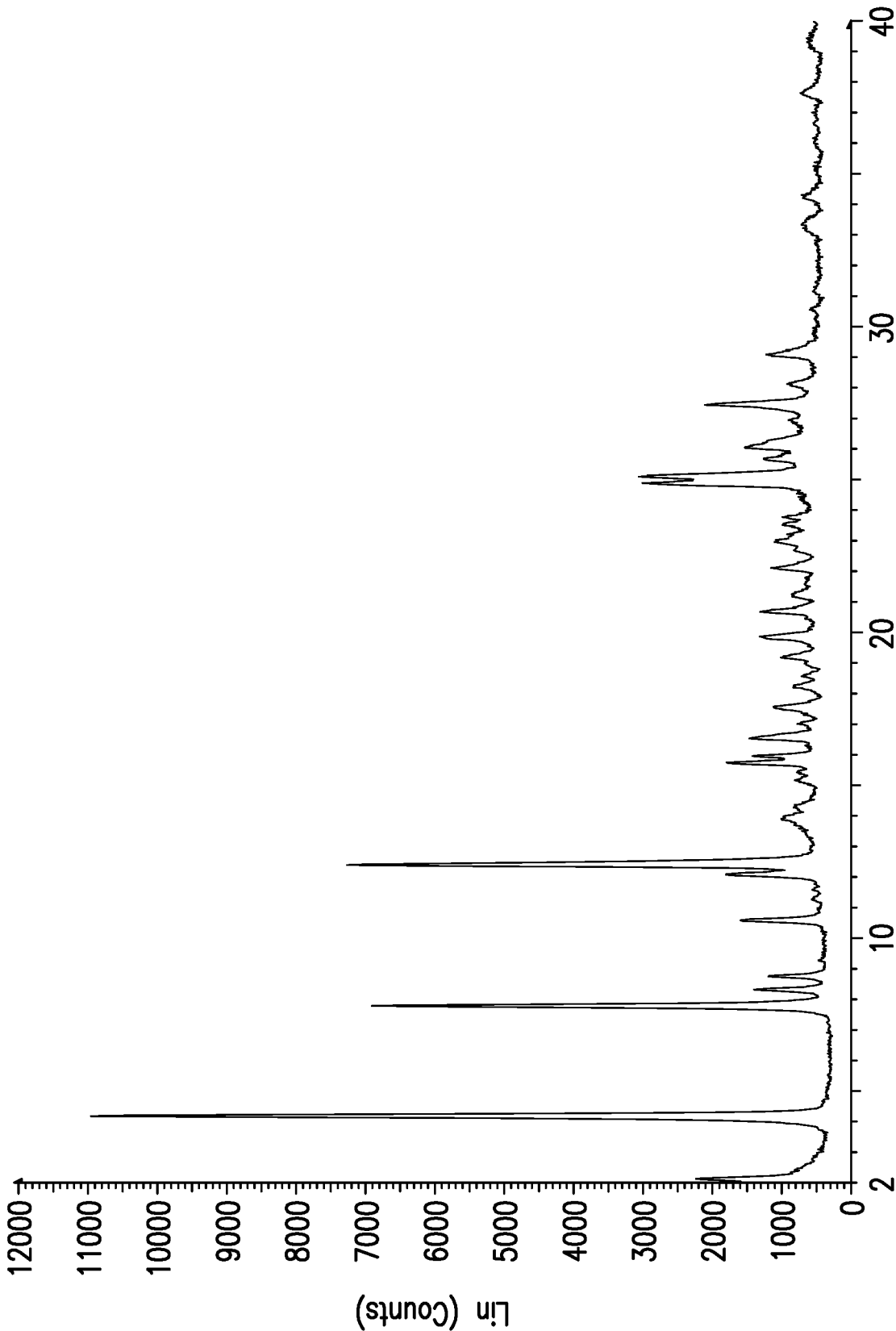
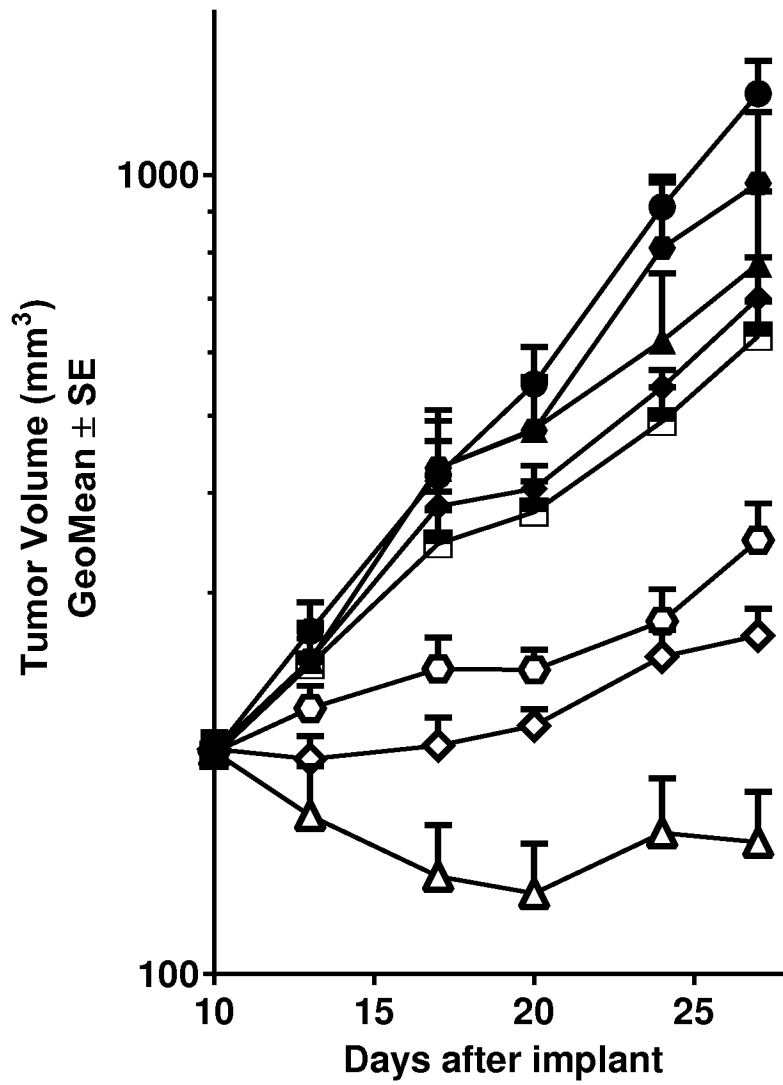
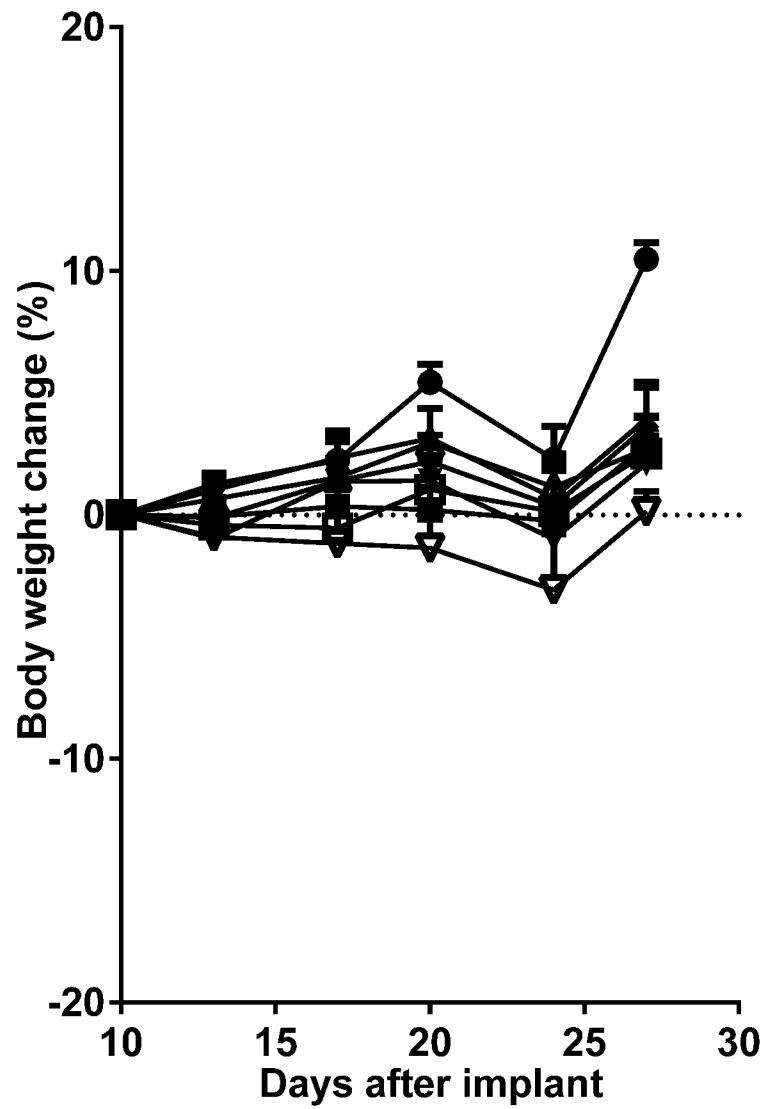


FIG.18

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**FIG. 19**

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**FIG. 20**

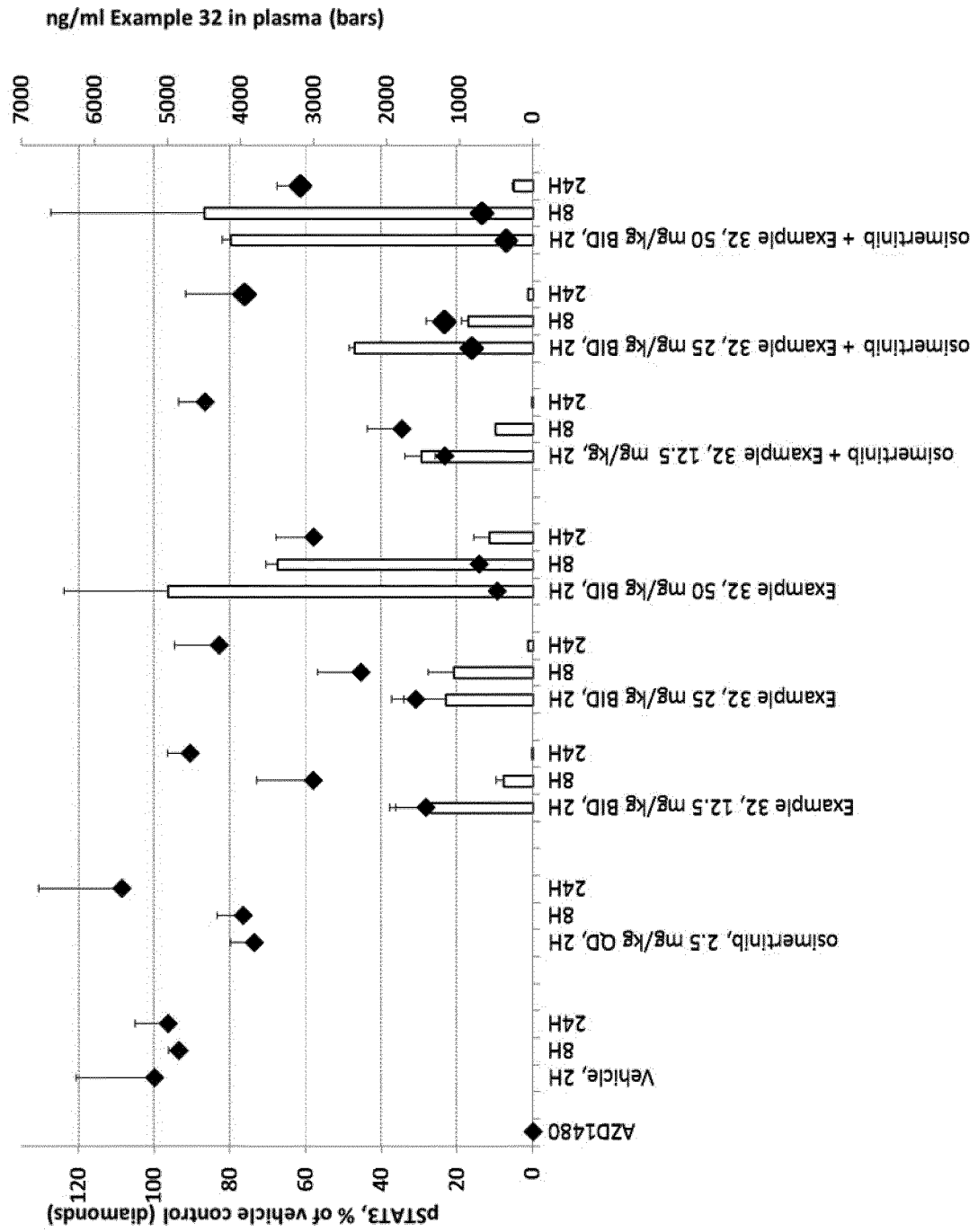
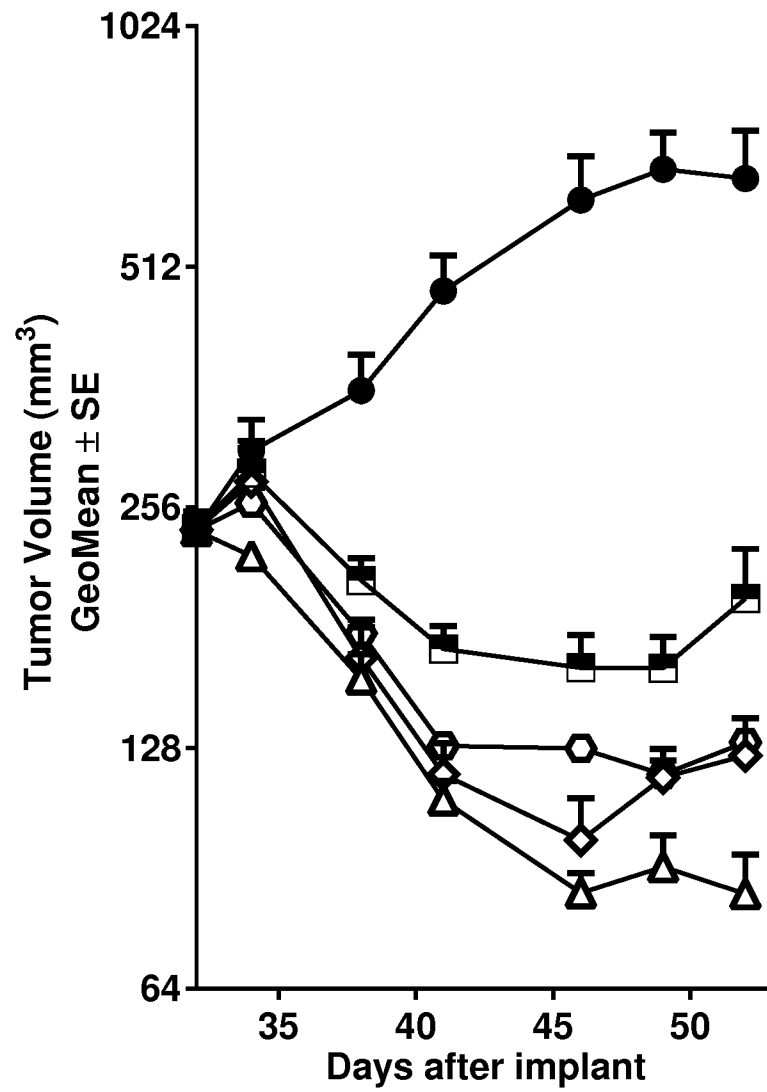
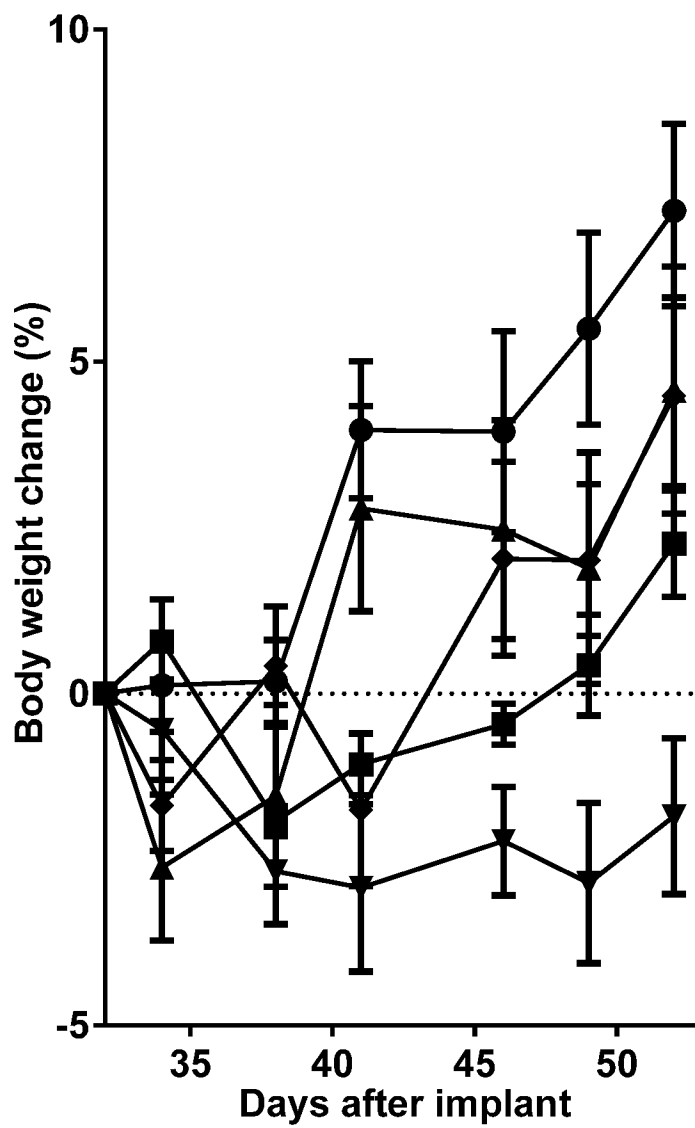


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**FIG. 22**

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**FIG. 23**

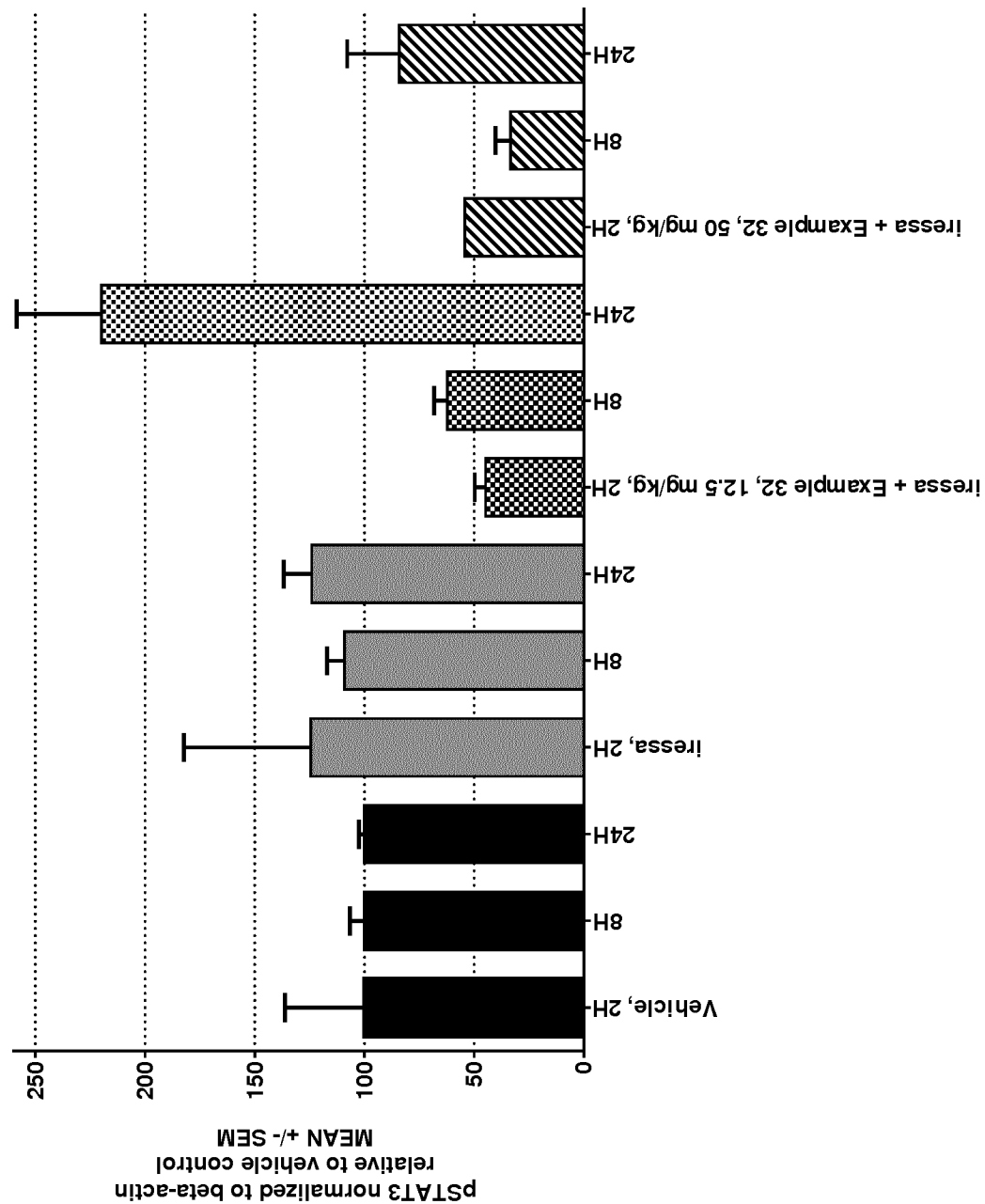
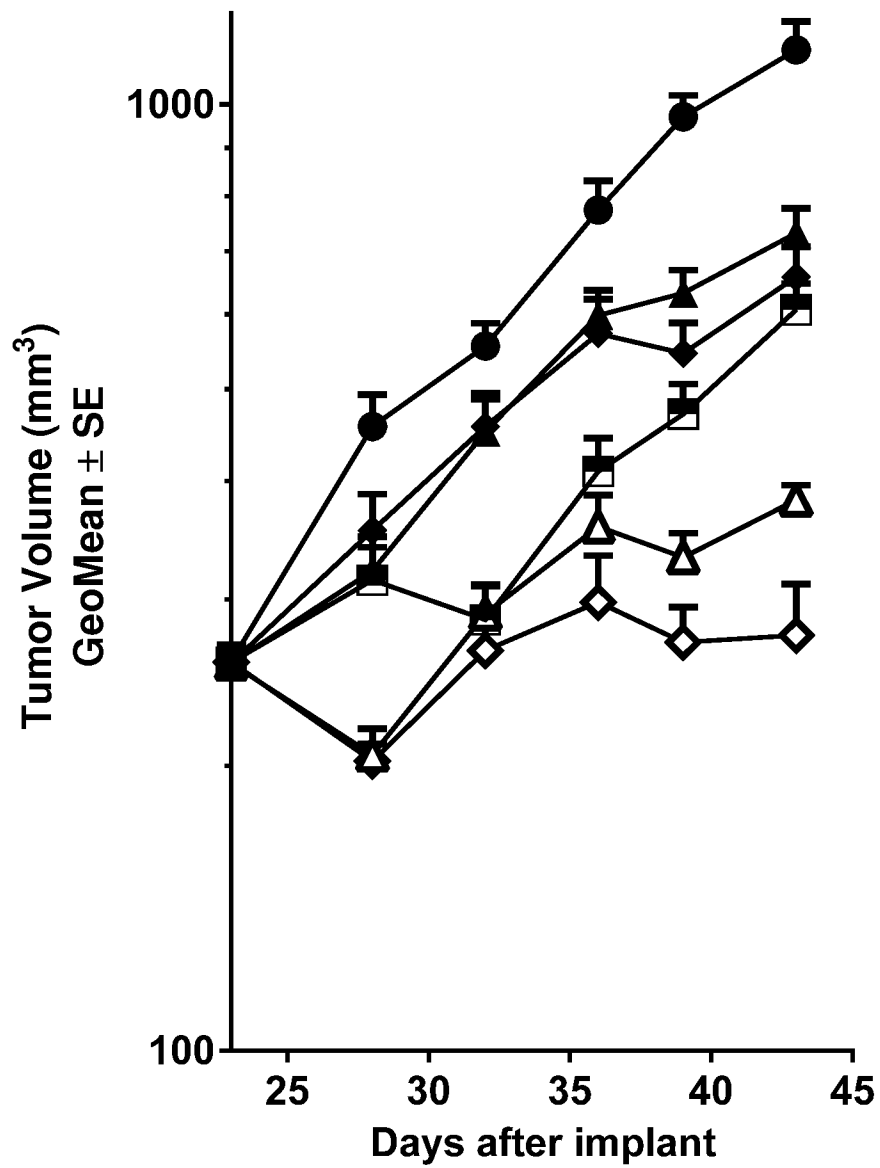
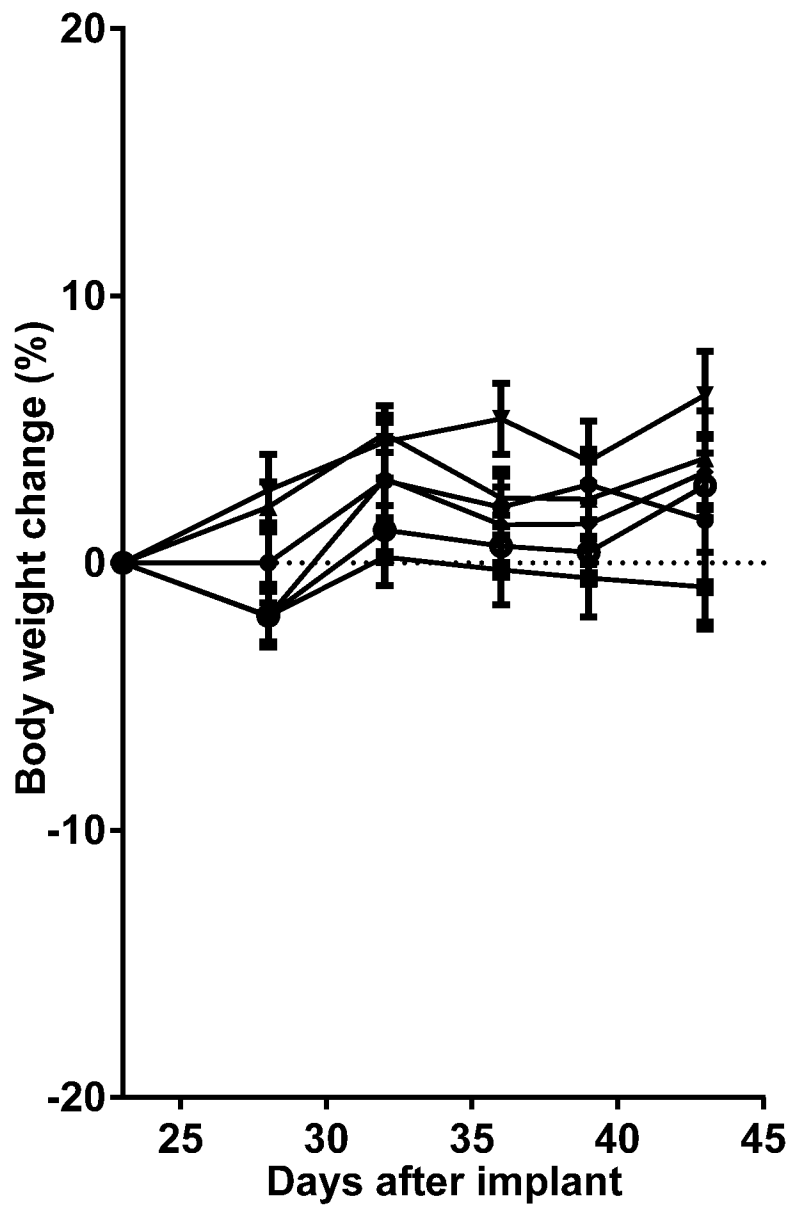


FIG. 24

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**FIG. 25**

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**FIG. 26**

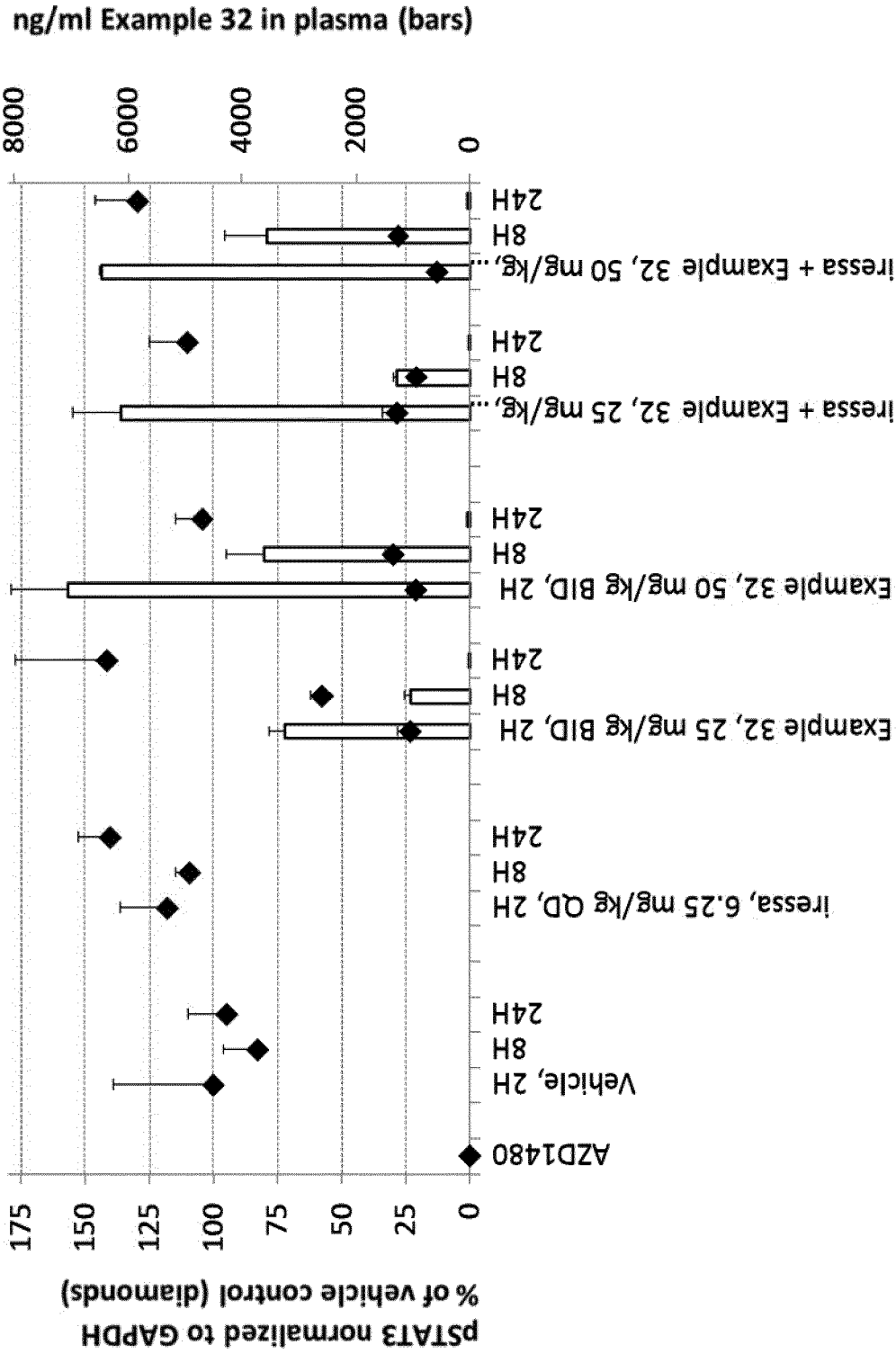


FIG. 27

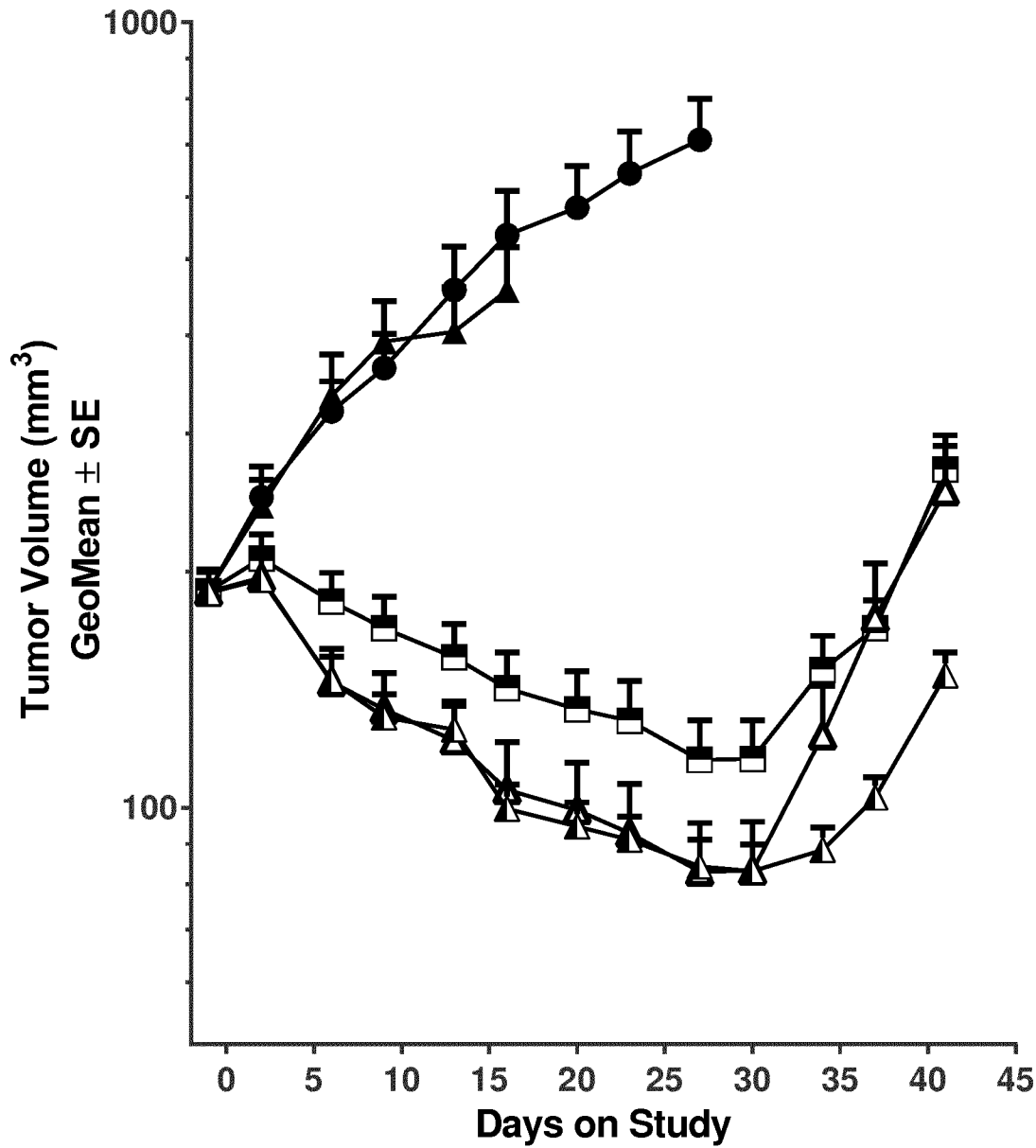
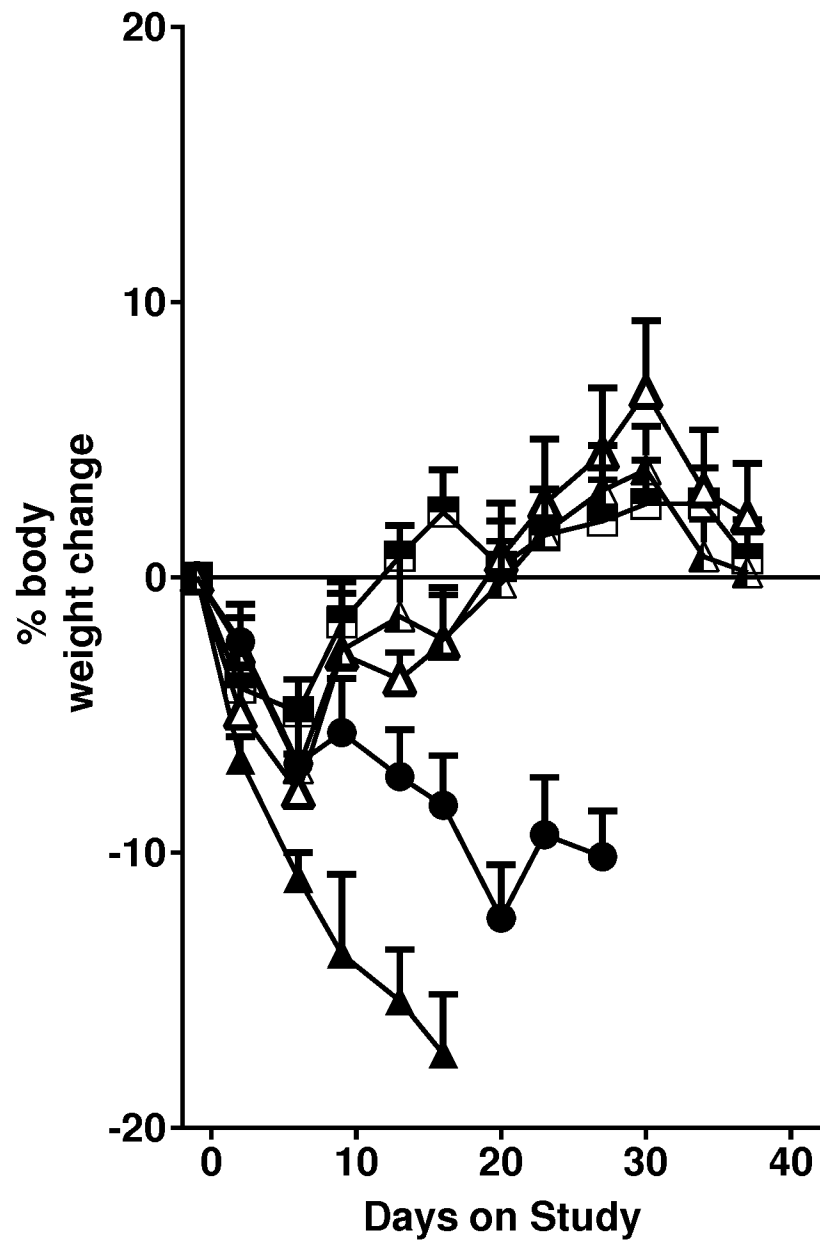


FIG. 28

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**FIG. 29**

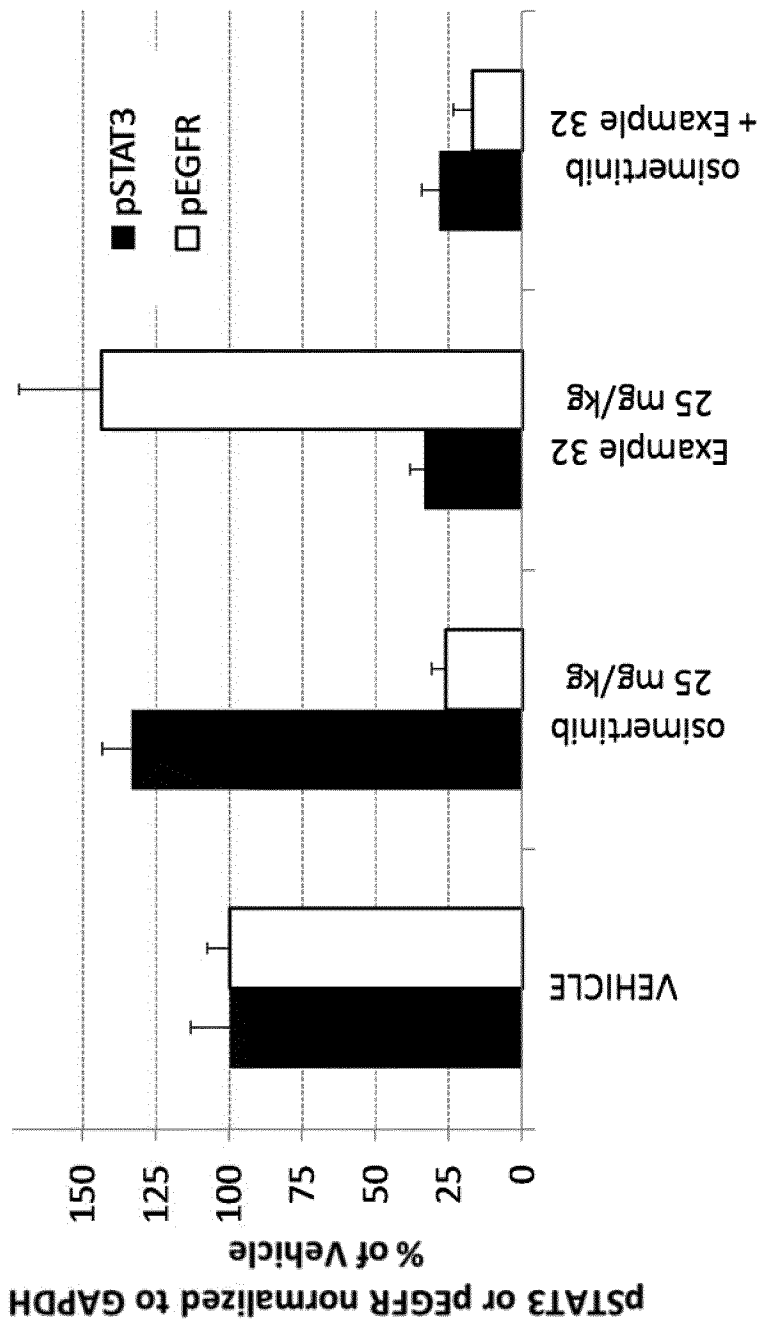
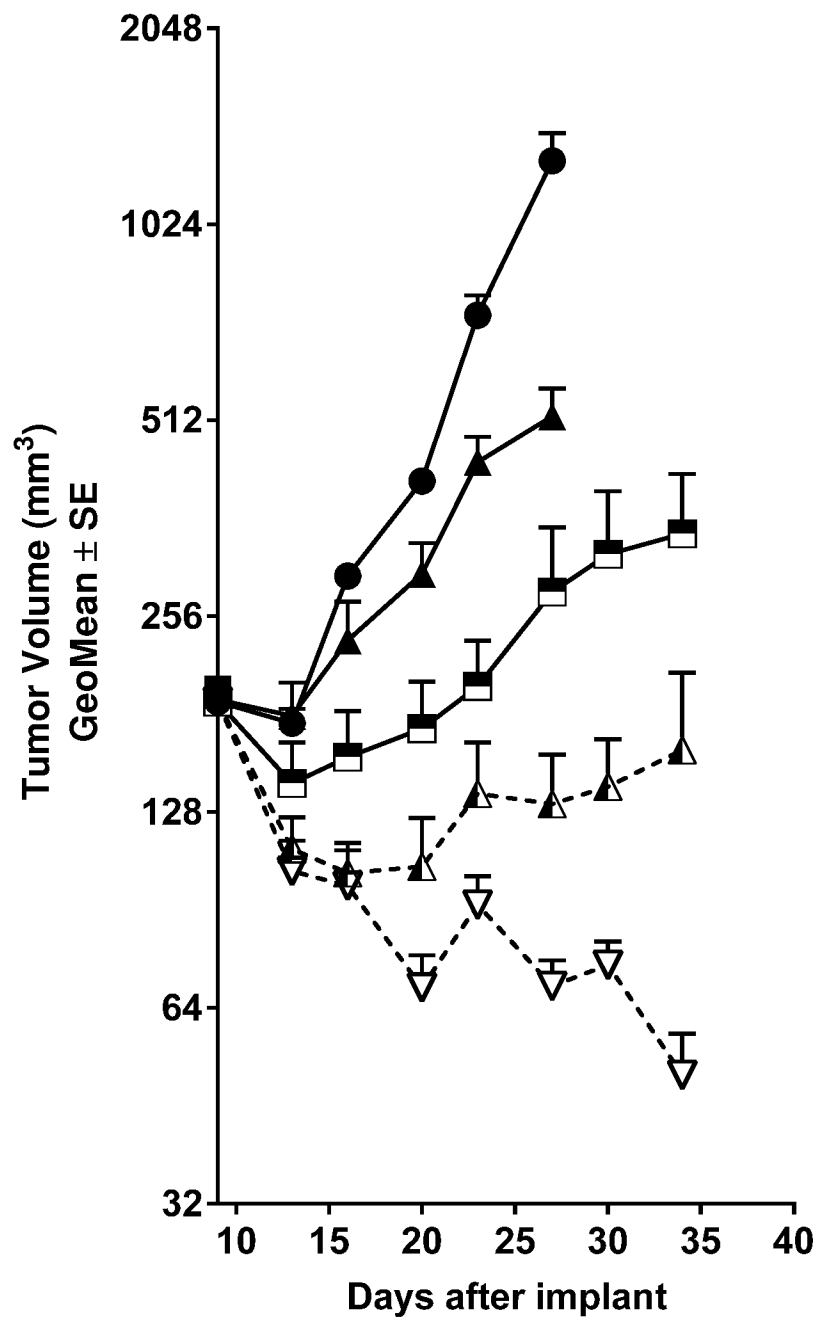
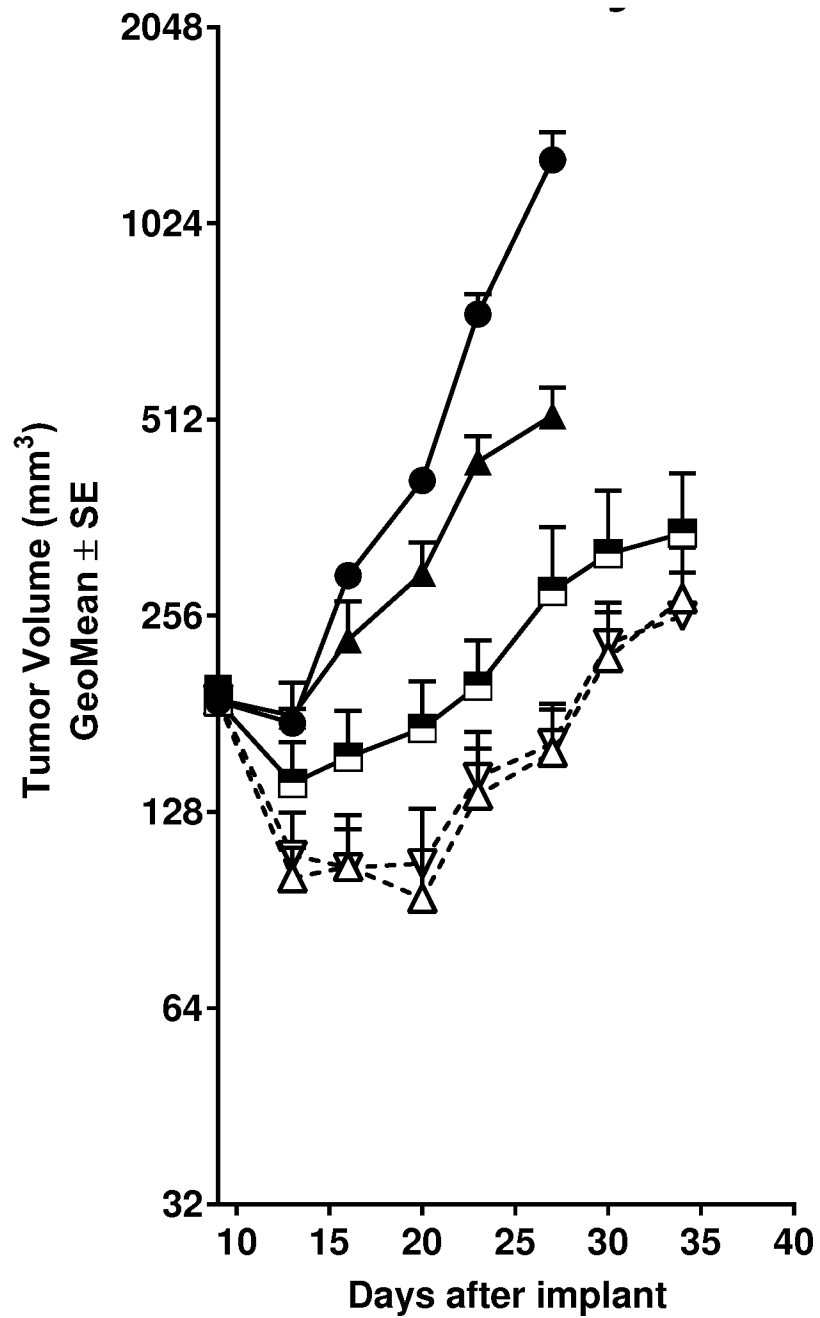


FIG. 30

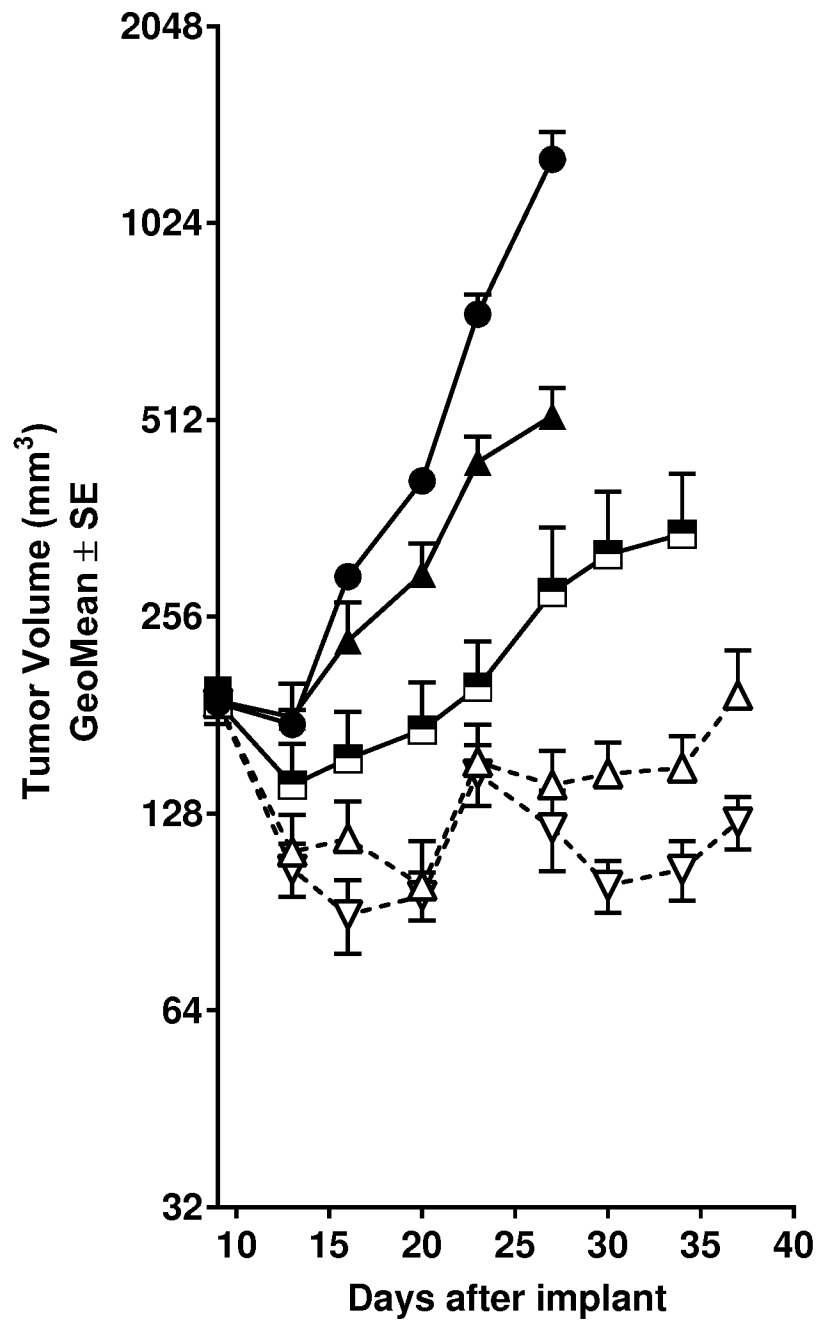
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**FIG. 31A**

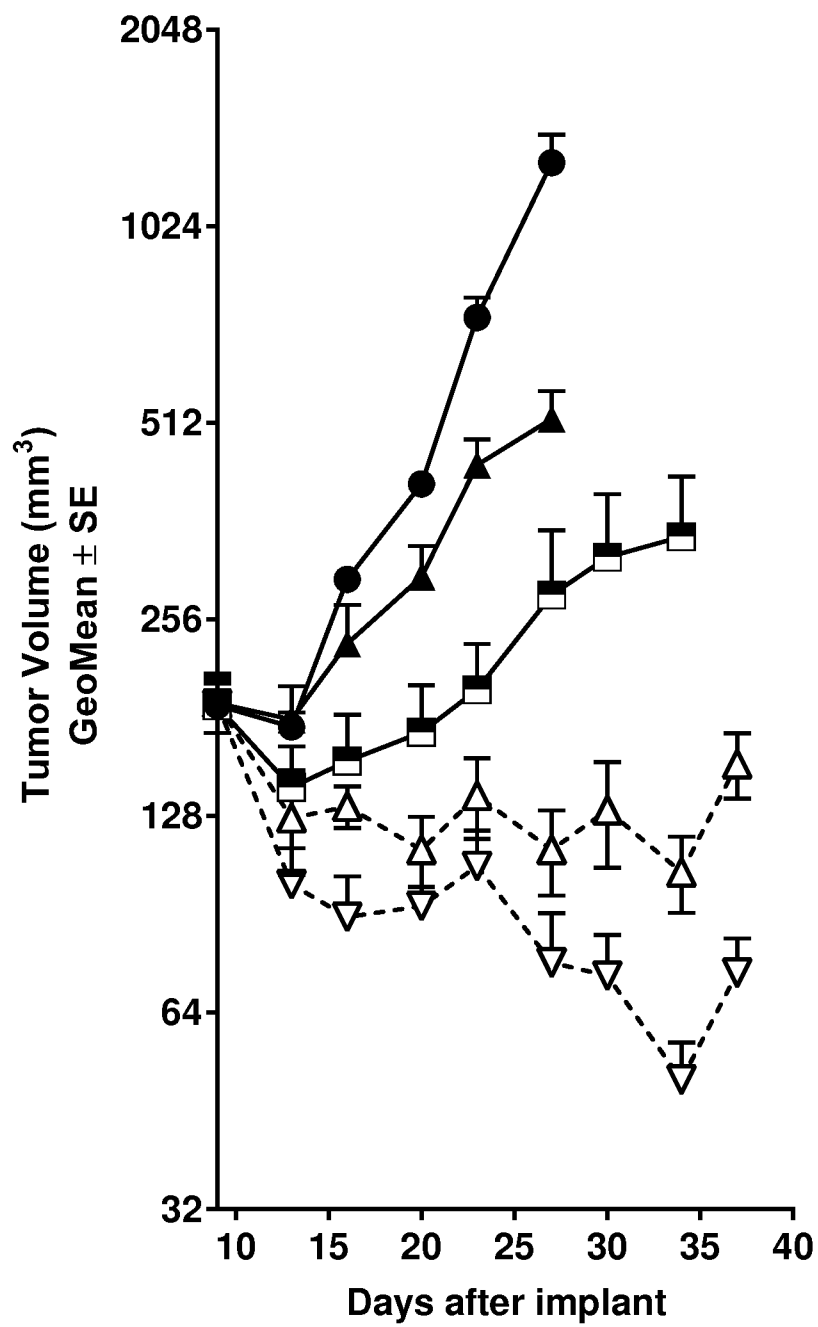
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**FIG. 31B**

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**FIG. 31C**

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**FIG. 31D**

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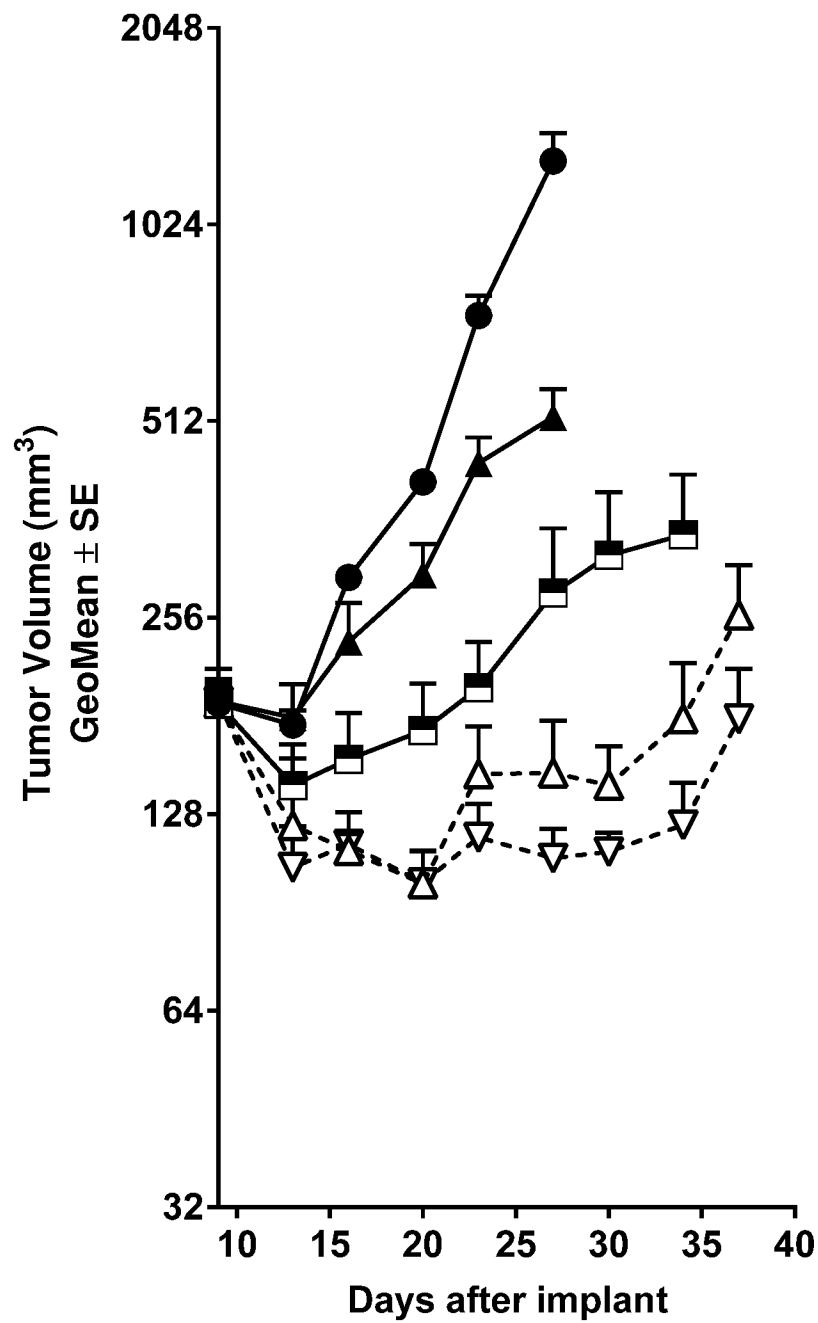
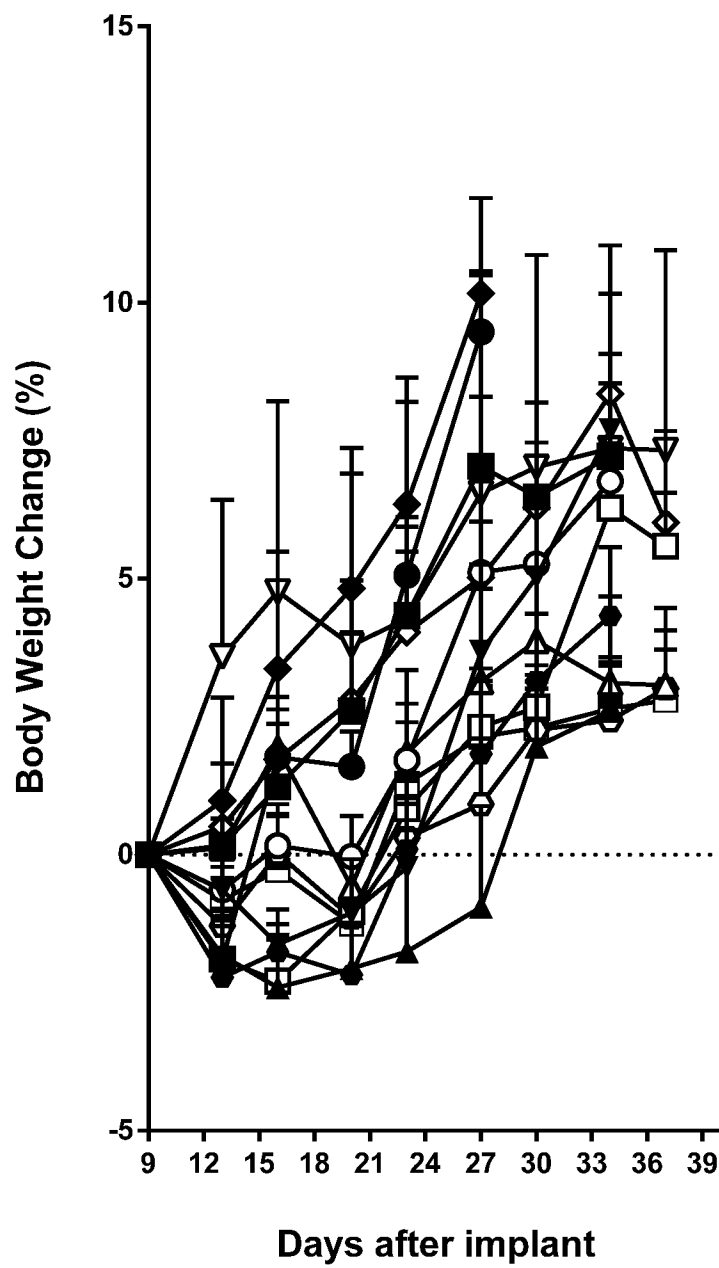


FIG 31E

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**FIG. 32**

INTERNATIONAL SEARCH REPORT

International application No
PCT/EP2016/072616

A. CLASSIFICATION OF SUBJECT MATTER
INV. C07D403/14 A61K31/506 A61P35/00
ADD.

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
C07D

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

EPO-Internal, WPI Data, CHEM ABS Data

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	WO 2012/030910 A1 (AMBIT BIOSCIENCES CORP [US]; HADD MICHAEL J [US]) 8 March 2012 (2012-03-08) paragraph [0002] paragraph [0010] paragraph [0227] claim 1 -----	1-13
A	WO 2012/116247 A1 (SYNTA PHARMACEUTICALS CORP [US]; BLACKMAN RONALD K [US]; FOLEY KEVIN P) 30 August 2012 (2012-08-30) page 2, column 1 - page 4, column 3 claim 1 -----	1-13



Further documents are listed in the continuation of Box C.



See patent family annex.

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"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

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Date of the actual completion of the international search

13 December 2016

Date of mailing of the international search report

21/12/2016

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Fax: (+31-70) 340-3016

Authorized officer

Koch, Kristian

INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No

PCT/EP2016/072616

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 2012030910 A1	08-03-2012	EP 2611793 A1	10-07-2013
		US 2013225615 A1	29-08-2013
		WO 2012030910 A1	08-03-2012

WO 2012116247 A1	30-08-2012	US 2014045908 A1	13-02-2014
		WO 2012116247 A1	30-08-2012
