

THE DESCRIPTION

COMPOUNDS AS HYPOXIA MIMETICS, AND COMPOSITIONS, AND USES THEREOF

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This application claims benefit of priority to Chinese priority patent application serial no. 200910089274.1 filed on July 15, 2009, which is hereby incorporated by reference in its entirety.

FIELD OF THE INVENTION

10 The present invention relates to substituted heterocyclic compounds, and compositions thereof as well as methods of use the same as hypoxia mimetics. This invention further relates to methods of increasing HIF (Hypoxia Inducible Factor) levels or activity in a subject or treating a condition associated with HIF levels or activity such as ischemia, anemia, wound healing, auto-transplantation, allo-transplantation, xenotransplantation, systemic high blood pressure, thalassemia, diabetes, cancer,
15 and an inflammatory disorder.

BACKGROUND OF THE INVENTION

The cellular transcription factor HIF (Hypoxia Inducible Factor) occupies a central position in oxygen homeostasis in a wide range of organisms and is a key regulator of responses to hypoxia. The
20 genes regulated by HIF transcriptional activity can play critical roles in angiogenesis, erythropoiesis, hemoglobin F production, energy metabolism, inflammation, vasomotor function, apoptosis and cellular proliferation. HIF can also play a role in cancer, in which it is commonly upregulated, and in the pathophysiological responses to ischemia and hypoxia.

The HIF transcriptional complex comprises an heterodimer: HIF is a constitutive nuclear protein that
25 dimerizes with oxygen-regulated HIF subunits. Oxygen regulation occurs through hydroxylation of the HIF subunits, which are then rapidly destroyed by the proteasome. In oxygenated cells, the von Hippel-Lindau tumor suppressor protein (pVHL) binds to hydroxylated HIF-subunits, thereby promoting their ubiquitin dependent proteolysis. This process is suppressed under hypoxic conditions, stabilizing HIF and promoting transcriptional activation by the HIF complex.

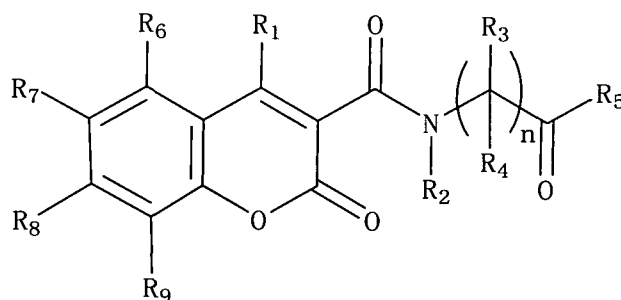
30 Hydroxylation of HIF-subunits can occur on proline and asparagine residues and can be mediated by

a family of 2-oxoglutarate dependent enzymes. This family includes the HIF prolyl hydroxylase isozymes (PHDs), which hydroxylate Pro 402 and Pro 564 of human HIF1, as well as Factor Inhibiting HIF (FIH), which hydroxylates Asn 803 of human HIF1. Inhibition of FIH or the PHDs leads to HIF stabilization and transcriptional activation. See, *e.g.*, Schofield and Ratcliffe, Nature Rev. Mol. Cell Biol., Vol15, pages 343-354 (2004).

Thus, new or improved agents that modulate (such as increasing HIF levels or activity) HIF are continually needed for developing new and more effective pharmaceuticals to treat HIF-associated conditions or diseases or disorders, such as ischemia, anemia, wound healing, auto-transplantation, allo-transplantation, xenotransplantation, systemic high blood pressure, thalassemia, diabetes, cancer, and an inflammatory disorder, to name a few. In discovering new or improved agents that modulate HIF level or activity, it is also desirable but not required to discover agents with improved chemical or biological properties such as solubility, bioavailability, pharmacokinetics, pharmacodynamics, toxicity and/or less side effects such as less cardiovascular side effects. The compounds, compositions, and methods described herein are directed toward these needs and other ends.

SUMMARY OF THE INVENTION

The present invention provides, *inter alia*, compounds of Formula I:



I

a pharmaceutically acceptable salt thereof, a solvate thereof, a chelate thereof, a non-covalent complex thereof, a prodrug thereof, or a mixture of any of the foregoing, wherein constituent members are provided below.

The present invention further provides pharmaceutical compositions comprising a compound of Formula I or pharmaceutically acceptable salt thereof, and at least one pharmaceutically acceptable carrier.

The present invention further provides methods of modulating an activity of HIF comprising

contacting the HIF with a compound of Formula I or pharmaceutically acceptable salt of the same.

The present invention further provides methods of increasing an activity of HIF with a compound of Formula I or pharmaceutically acceptable salt of the same.

5 The present invention further provides methods of modulating (such as increasing) the level of HIF in a subject (such as a cell or a patient) comprising administering to the subject a therapeutically effective amount of a compound of Formula I or pharmaceutically acceptable salt of the same.

The present invention further provides methods of treating one or more of the various HIF-associated conditions, diseases, and disorders named herein by administering to a patient a therapeutically effective amount of a compound of Formula I or pharmaceutically acceptable salt of the
10 same.

The present invention further provides compounds of Formula I or pharmaceutically acceptable salts thereof, for use in therapy.

The present invention further provides use of the compounds of Formula I or pharmaceutically acceptable salts thereof, for the manufacture/preparation of a medicament for use in therapy.

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DESCRIPTION OF DRAWINGS

Figure 1 illustrates certain example compounds' effects on the level of EPO in mice 4 hours after the administration of different compounds (using the assay described in Example B).

20 Figure 2 illustrates certain example compounds' effects on the level of red blood cell counts (RBC) in mice at Day 9 after 7 days of daily dosage of 60 mg/kg (using the assay described in Example C).

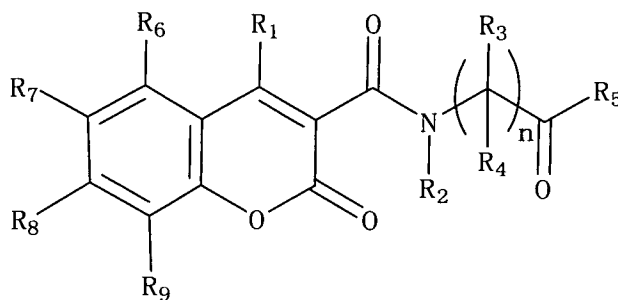
Figure 3 illustrates certain example compounds' effects on the level of blood hemoglobin (HGB) in mice at Day 9 after 7 days of daily dosage of 60 mg/kg (using the assay described in Example C).

Figure 4 illustrates the pharmacokinetic curves in rats for certain example compounds after single oral dosage of 50 mg/kg (using the assay described in Example D).

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DETAILED DESCRIPTION

The present invention provides, *inter alia*, compounds of Formula I:



I

a pharmaceutically acceptable salt thereof, a solvate thereof, a chelate thereof, a non-covalent complex thereof, a prodrug thereof, and mixtures of any of the foregoing, wherein:

n is 1 to 6;

R₁ is selected from OH, SH, NR₃R₄, NHC(O)R₂, NHSO₂R₂ and sulfonyl;

R₂ is selected from H, lower alkyl and substituted lower alkyl;

R₃ and R₄ are independently selected from H, lower alkyl, substituted lower alkyl, lower haloalkyl, substituted lower haloalkyl, or R₃ and R₄ can join together to form a 3 to 6 membered ring or a substituted 3 to 6 membered ring;

R₅ is selected from OH, SH, NH₂, lower alkyl, substituted lower alkyl, lower alkoxy, substituted lower alkoxy, and sulfanyl;

each of R₆, R₇, R₈ and R₉ is independently selected from H, alkyl, substituted alkyl, alkenyl, substituted alkenyl, alkynyl, substituted alkynyl, alkoxy, substituted alkoxy, NR₃R₄, C(O)OH, OR₁₂, SR₁₂, SO₂R₁₂, CN, NO₂, halogen, aryl, substituted aryl, heteroaryl, substituted heteroaryl, arylalkyl, substituted arylalkyl, heteroarylalkyl, substituted heteroarylalkyl, heterocycloalkyl, substituted heterocycloalkyl, alkylsilyl, substituted alkylsilyl, alkenylsilyl, substituted alkenylsilyl, alkynylsilyl, substituted alkynylsilyl, alkoxy carbonyl, substituted alkoxy carbonyl, and -X-R₁₁;

or at least one of adjacent pairs R₁ and R₆, R₆ and R₇, R₇ and R₈, and R₈ and R₉, can join together to form a 4 to 7 membered ring or a substituted 4 to 7 membered ring;

X is selected from -N(R₁₀)-Y- and -Y-N(R₁₀)-;

Y is selected from C(O), SO₂, alkylene, substituted alkylene, alkenylene, substituted alkenylene, alkynylene, and substituted alkynylene;

R₁₀ is selected from H, lower alkyl, and substituted lower alkyl;

R₁₁ is selected from H, cycloalkyl, substituted cycloalkyl, heterocycloalkyl, substituted heterocycloalkyl, aryl, substituted aryl, heteroaryl, and substituted heteroaryl; and

R_{12} is selected from H, alkyl, substituted alkyl, alkenyl, substituted alkenyl, alkynyl, substituted alkynyl and NR_3R_4 .

In some embodiments, R_1 is selected from OH and SH. In some further embodiments, R_1 is OH.

In some embodiments, R_2 is selected from H and lower alkyl (such as methyl or ethyl). In some
5 further embodiments, R_2 is H.

In some embodiments, R_3 and R_4 are each, independently, selected from H, lower alkyl, and lower haloalkyl. In some embodiments, R_3 and R_4 are each, independently, selected from H and lower alkyl (such as methyl or ethyl). In some embodiments, R_3 and R_4 are each H.

In some embodiments, R_3 and R_4 , together with the carbon atom to which they are attached, can
10 form a 3 to 6 membered cycloalkyl or heterocycloalkyl ring, wherein the 3 to 6 membered cycloalkyl or heterocycloalkyl ring is optionally substituted with 1, 2, 3, 4, or 5 substituents each independently selected from lower alkyl, lower alkoxy, lower haloalkyl, lower haloalkoxy, halogen, and OH. In some further embodiments, R_3 and R_4 , together with the carbon atom to which they are attached, form a 3 to 6 membered cycloalkyl optionally substituted with 1, 2, 3, 4, or 5 substituents each independently selected
15 from lower alkyl, lower alkoxy, lower haloalkyl, lower haloalkoxy, halogen, and OH. In some further embodiments, R_3 and R_4 , together with the carbon atom to which they are attached, form a 3 to 6 membered cycloalkyl which is unsubstituted.

In some embodiments, R_3 and R_4 , together with the carbon atom to which they are attached, form a 3 to 6 membered heterocycloalkyl optionally substituted with 1, 2, 3, 4, or 5 substituents each
20 independently selected from lower alkyl, lower alkoxy, lower haloalkyl, lower haloalkoxy, halogen, and OH. In some further embodiments, the heterocycloalkyl formed by R_3 and R_4 , together with the carbon atom to which they are attached comprises at least one or two heteroatoms each independently selected from O, S, and N. In some further embodiments, R_3 and R_4 , together with the carbon atom to which they are attached, form a 3 to 6 membered heterocycloalkyl selected from pyrrolidinyl, piperidinyl,
25 tetrahydrofuranyl, and tetrahydropyranyl.

In some embodiments, R_5 is selected from OH, SH, NH_2 , and lower alkoxy. In some further embodiments, R_5 is selected from OH and SH. In yet further embodiments, R_5 is OH.

In some embodiments, R_5 is selected from OH, NH_2 , and lower alkoxy (such as methoxy, ethoxy and propoxy).

30 In some embodiments, R_5 is selected from OH, lower alkoxy (such as methoxy, ethoxy and propoxy),

and substituted lower alkoxy.

In some embodiments, at least one of adjacent pairs R₆ and R₇, R₇ and R₈, and R₈ and R₉, can join together (and together with the two carbon atoms to which they are attached) to form a 4 to 7 membered ring or a substituted 4 to 7 membered ring. In some further embodiments, the 4 to 7 membered ring or
5 the substituted 4 to 7 membered ring comprises at least one, two, or three heteroatoms.

In some embodiments, R₆ and R₇ are each, independently, selected from H, halogen, OH, lower alkyl, lower haloalkyl, lower alkoxy, lower haloalkoxy. In some further embodiments, R₆ and R₇ are each, independently, selected from H, halogen, lower alkyl (such as methyl or ethyl), and lower haloalkyl (such as trifluoromethyl). In yet embodiments, R₆ and R₇ are each H.

10 In some embodiments, R₈ and R₉ are each, independently, selected from H, halogen, OH, lower alkyl, lower haloalkyl, lower alkoxy, lower haloalkoxy. In some further embodiments, R₈ and R₉ are each, independently, selected from H, halogen, lower alkyl (such as methyl or ethyl), and lower haloalkyl (such as trifluoromethyl). In yet embodiments, R₈ and R₉ are each H.

In some embodiments, at least one of R₆, R₇, R₈ and R₉ is independently selected from halo,
15 haloalkyl and haloalkoxy.

In some embodiments, at least one of R₆, R₇, R₈ and R₉ is independently selected from alkoxy or substituted alkoxy.

In some embodiments, at least one of R₆, R₇, R₈ and R₉ is independently selected from alkylsilyl, substituted alkylsilyl, alkynylsilyl, and substituted alkynylsilyl.

20 In some embodiments, at least one of R₆, R₇, R₈ and R₉ is independently selected from aryl, substituted aryl, heteroaryl, substituted heteroaryl, heterocycloalkyl, and substituted heterocycloalkyl.

In some embodiments, at least one of R₆, R₇, R₈ and R₉ is independently selected from H, alkyl, substituted alkyl, alkenyl, substituted alkenyl, alkynyl, and substituted alkynyl.

In some embodiments, at least one of R₈ and R₉ is heteroaryl or substituted heteroaryl. In some
25 further embodiments, at least one of R₈ and R₉ is heteroaryl selected from pyridyl (e.g., 2-pyridyl, 3-pyridyl, or 4-pyridyl), thienyl, furanyl. In yet further embodiments, at least one of R₈ and R₉ is heteroaryl selected from pyridyl (e.g., 2-pyridyl, 3-pyridyl, or 4-pyridyl).

In some embodiments, at least one of R₈ and R₉ is independently selected from substituted aryl, and substituted heteroaryl.

30 In some embodiments, at least one of R₈ and R₉ is independently selected from substituted phenyl,

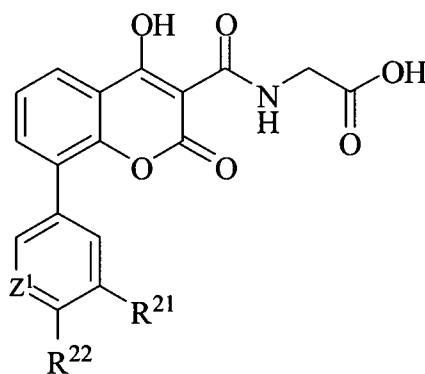
and substituted pyridyl.

In some embodiments, at least one of R_8 and R_9 is phenyl substituted with halogen, alkyl, haloalkyl, alkoxy, and haloalkoxy at the para- or meta- position. In some further embodiments, one of R_8 and R_9 is phenyl substituted with halogen, alkyl, haloalkyl, alkoxy, and haloalkoxy at the para- or meta- position; and the other of R_8 and R_9 is H, halogen, OH, lower alkyl, lower haloalkyl, lower alkoxy, lower haloalkoxy. In yet further embodiments, one of R_8 and R_9 is phenyl substituted with halogen, alkyl, haloalkyl, alkoxy, and haloalkoxy at the para- or meta- position; and the other of R_8 and R_9 is H.

In some embodiments, at least one of R_8 and R_9 is pyridyl substituted with halogen, alkyl, haloalkyl, alkoxy, and haloalkoxy at the para- or meta- position. In some further embodiments, one of R_8 and R_9 is pyridyl substituted with halogen, alkyl, haloalkyl, alkoxy, and haloalkoxy at the para- or meta- position; and the other of R_8 and R_9 is H, halogen, OH, lower alkyl, lower haloalkyl, lower alkoxy, lower haloalkoxy. In yet further embodiments, one of R_8 and R_9 is pyridyl substituted with halogen, alkyl, haloalkyl, alkoxy, and haloalkoxy at the para- or meta- position; and the other of R_8 and R_9 is H.

In some embodiments, n is 1, 2, or 3. In some further embodiments, n is 1 or 2. In yet further embodiments, n is 1.

In some embodiments, the compound of Formula I or pharmaceutically acceptable salt thereof is a compound of Formula II:



II

or pharmaceutically acceptable salt thereof, wherein:

R^{21} is selected from H, halogen, lower alkyl, lower haloalkyl, lower alkoxy, and lower haloalkoxy;

R^{22} is selected from H, halogen, lower alkyl, lower haloalkyl, lower alkoxy, and lower haloalkoxy;

Z^1 is N or CR^{23} ; and

R^{23} is selected from H, halogen, lower alkyl, lower haloalkyl, lower alkoxy, and lower haloalkoxy;

provided that at least one of R^{21} and R^{22} and R^{23} (if present) is selected from halogen, lower alkyl,

lower haloalkyl, lower alkoxy, and lower haloalkoxy.

In some embodiments of compound of Formula II or pharmaceutically acceptable salt thereof, R^{21} is selected from H, Cl, F, methyl, ethyl, trifluoromethyl, methoxy, ethoxy, and trifluoromethoxy.

In some embodiments of compound of Formula II or pharmaceutically acceptable salt thereof, R^{22} is selected from H, Cl, F, methyl, ethyl, trifluoromethyl, methoxy, ethoxy, and trifluoromethoxy.

In some embodiments of compound of Formula II or pharmaceutically acceptable salt thereof, Z^1 is N.

In some embodiments of compound of Formula II or pharmaceutically acceptable salt thereof, Z^1 is CR^{23} .

In some embodiments of compound of Formula II or pharmaceutically acceptable salt thereof, Z^1 is CR^{23} and R^{23} is selected from H, Cl, F, methyl, ethyl, trifluoromethyl, methoxy, ethoxy, and trifluoromethoxy.

In some embodiments of compound of Formula II or pharmaceutically acceptable salt thereof, Z^1 is CR^{23} ; one of R^{21} , R^{22} , and R^{23} is selected from halogen, lower alkyl, lower haloalkyl, lower alkoxy, and lower haloalkoxy; and the other two of R^{21} , R^{22} , and R^{23} are both H.

In some embodiments of compound of Formula II or pharmaceutically acceptable salt thereof, Z^1 is CR^{23} ; one of R^{21} and R^{23} is selected from halogen, lower alkyl, lower haloalkyl, lower alkoxy, and lower haloalkoxy; and the other one of R^{21} and R^{23} is H. In some further embodiments, R^{22} is H.

In some embodiments of compound of Formula II or pharmaceutically acceptable salt thereof, Z^1 is CR^{23} ; two of R^{21} , R^{22} , and R^{23} are each independently selected from halogen, lower alkyl, lower haloalkyl, lower alkoxy, and lower haloalkoxy; and the other one of R^{21} , R^{22} , and R^{23} is H. In some further embodiments, R^{23} is H. In other further embodiments, R^{22} is H.

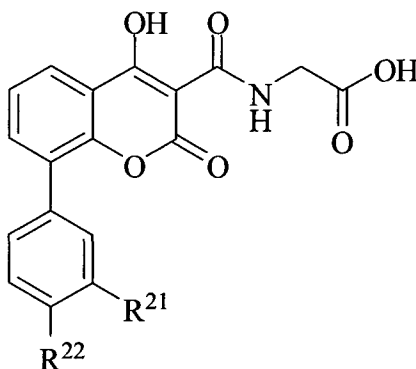
In some embodiments of compound of Formula II or pharmaceutically acceptable salt thereof, Z^1 is CR^{23} ; R^{21} , R^{22} , and R^{23} are each independently selected from halogen, lower alkyl, lower haloalkyl, lower alkoxy, and lower haloalkoxy.

In some embodiments of compound of Formula II or pharmaceutically acceptable salt thereof, Z^1 is N; one of R^{21} and R^{22} is selected from halogen, lower alkyl, lower haloalkyl, lower alkoxy, and lower haloalkoxy; and the other of R^{21} and R^{22} is H.

In some embodiments of compound of Formula II or pharmaceutically acceptable salt thereof, Z^1 is N; R^{21} and R^{22} are each independently selected from halogen, lower alkyl, lower haloalkyl, lower alkoxy,

and lower haloalkoxy.

In some embodiments, the compound of Formula II or pharmaceutically acceptable salt thereof is a compound of Formula IIa



IIa

or pharmaceutically acceptable salt thereof, wherein:

R^{21} is selected from H, halogen, lower alkyl, lower haloalkyl, lower alkoxy, and lower haloalkoxy;

and

R^{22} is selected from H, halogen, lower alkyl, lower haloalkyl, lower alkoxy, and lower haloalkoxy;

provided that at least one of R^{21} and R^{22} is selected from halogen, lower alkyl, lower haloalkyl, lower alkoxy, and lower haloalkoxy.

In some embodiments of compound of Formula IIa or pharmaceutically acceptable salt thereof, R^{21} is H; and R^{22} is selected from halogen, lower alkyl, lower haloalkyl, lower alkoxy, and lower haloalkoxy.

In some embodiments of compound of Formula IIa or pharmaceutically acceptable salt thereof, R^{21} is H; and R^{22} is selected from lower alkyl and lower haloalkyl.

In some embodiments of compound of Formula IIa or pharmaceutically acceptable salt thereof, R^{21} is H; and R^{22} is selected from C_{1-3} alkyl and C_{1-3} haloalkyl. In some further embodiments, R^{22} is selected from C_{1-2} alkyl and C_{1-2} haloalkyl. In yet further embodiments, R^{22} is selected from methyl and C_1 haloalkyl.

In some embodiments of compound of Formula IIa or pharmaceutically acceptable salt thereof, R^{21} is H; and R^{22} is selected from C_{1-3} alkyl. In some further embodiments, R^{22} is selected from methyl or ethyl. In yet further embodiments, R^{22} is methyl.

In some embodiments of compound of Formula IIa or pharmaceutically acceptable salt thereof, R^{22} is H; and R^{21} is selected from halogen, lower alkyl, lower haloalkyl, lower alkoxy, and lower haloalkoxy.

In some embodiments of compound of Formula IIa or pharmaceutically acceptable salt thereof, R^{22}

is H; and R²¹ is selected from lower alkyl and lower haloalkyl.

In some embodiments of compound of Formula IIa or pharmaceutically acceptable salt thereof, R²² is H; and R²¹ is selected from C₁₋₃ alkyl and C₁₋₃ haloalkyl. In some further embodiments, R²¹ is selected from C₁₋₂ alkyl and C₁₋₂ haloalkyl. In yet further embodiments, R²¹ is selected from methyl and C₁ haloalkyl.

In some embodiments of compound of Formula IIa or pharmaceutically acceptable salt thereof, R²² is H; and R²¹ is selected from C₁₋₃ alkyl. In some further embodiments, R²¹ is selected from methyl or ethyl. In yet further embodiments, R²¹ is methyl.

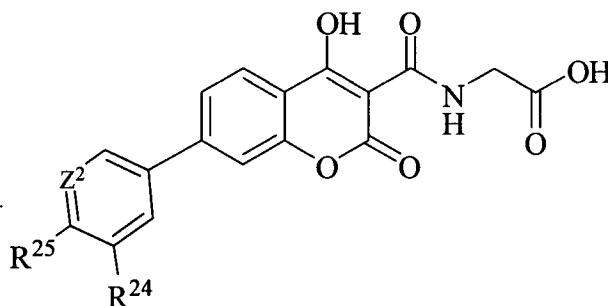
In some embodiments of compound of Formula IIa or pharmaceutically acceptable salt thereof, R²¹ and R²² are each independently selected from halogen, lower alkyl, lower haloalkyl, lower alkoxy, and lower haloalkoxy.

In some embodiments of compound of Formula IIa or pharmaceutically acceptable salt thereof, R²¹ and R²² are each independently selected from lower alkyl and lower haloalkyl.

In some embodiments of compound of Formula IIa or pharmaceutically acceptable salt thereof, R²¹ and R²² are each independently selected from C₁₋₃ alkyl and C₁₋₃ haloalkyl. In some further embodiments, R²¹ and R²² are each independently selected from C₁₋₂ alkyl and C₁₋₂ haloalkyl. In yet further embodiments, R²¹ and R²² are each independently selected from methyl and C₁ haloalkyl.

In some embodiments of compound of Formula IIa or pharmaceutically acceptable salt thereof, R²¹ and R²² are each independently selected from C₁₋₃ alkyl. In some further embodiments, R²¹ and R²² are each independently selected from methyl or ethyl. In yet further embodiments, R²¹ and R²² are each methyl.

In some embodiments, the compound of Formula I or pharmaceutically acceptable salt thereof is a compound of Formula III:



III

or pharmaceutically acceptable salt thereof, wherein:

R^{24} is selected from H, halogen, lower alkyl, lower haloalkyl, lower alkoxy, and lower haloalkoxy;

R^{25} is selected from H, halogen, lower alkyl, lower haloalkyl, lower alkoxy, and lower haloalkoxy;

Z^2 is N or CR^{26} ; and

R^{26} is selected from H, halogen, lower alkyl, lower haloalkyl, lower alkoxy, and lower haloalkoxy;

5 provided that at least one of R^{24} and R^{25} and R^{26} (if present) is selected from halogen, lower alkyl, lower haloalkyl, lower alkoxy, and lower haloalkoxy.

In some embodiments of compound of Formula III or pharmaceutically acceptable salt thereof, R^{24} is selected from H, Cl, F, methyl, ethyl, trifluoromethyl, methoxy, ethoxy, and trifluoromethoxy.

10 In some embodiments of compound of Formula III or pharmaceutically acceptable salt thereof, R^{25} is selected from H, Cl, F, methyl, ethyl, trifluoromethyl, methoxy, ethoxy, and trifluoromethoxy.

In some embodiments of compound of Formula III or pharmaceutically acceptable salt thereof, Z^2 is N.

In some embodiments of compound of Formula III or pharmaceutically acceptable salt thereof, Z^2 is CR^{26} .

15 In some embodiments of compound of Formula III or pharmaceutically acceptable salt thereof, Z^2 is CR^{26} and R^{26} is selected from H, Cl, F, methyl, ethyl, trifluoromethyl, methoxy, ethoxy, and trifluoromethoxy.

20 In some embodiments of compound of Formula III or pharmaceutically acceptable salt thereof, Z^2 is CR^{26} ; one of R^{24} , R^{25} , and R^{26} is selected from halogen, lower alkyl, lower haloalkyl, lower alkoxy, and lower haloalkoxy; and the other two of R^{24} , R^{25} , and R^{26} are both H.

In some embodiments of compound of Formula III or pharmaceutically acceptable salt thereof, Z^2 is CR^{26} ; one of R^{24} and R^{26} is selected from halogen, lower alkyl, lower haloalkyl, lower alkoxy, and lower haloalkoxy; and the other one of R^{24} and R^{26} is H. In some further embodiments, R^{25} is H.

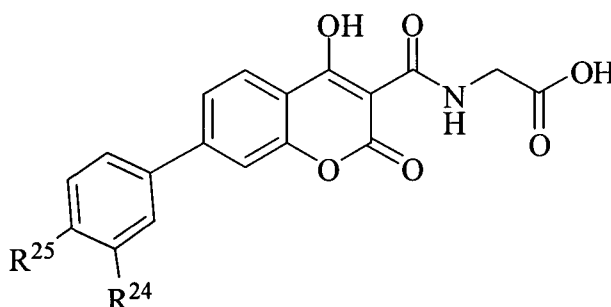
25 In some embodiments of compound of Formula III or pharmaceutically acceptable salt thereof, Z^2 is CR^{26} ; two of R^{24} , R^{25} , and R^{26} are each independently selected from halogen, lower alkyl, lower haloalkyl, lower alkoxy, and lower haloalkoxy; and the other one of R^{24} , R^{25} , and R^{26} is H. In some further embodiments, R^{26} is H. In other further embodiments, R^{25} is H.

30 In some embodiments of compound of Formula III or pharmaceutically acceptable salt thereof, Z^1 is CR^{26} ; R^{24} , R^{25} , and R^{26} are each independently selected from halogen, lower alkyl, lower haloalkyl, lower alkoxy, and lower haloalkoxy.

In some embodiments of compound of Formula III or pharmaceutically acceptable salt thereof, Z^1 is N; one of R^{24} and R^{25} is selected from halogen, lower alkyl, lower haloalkyl, lower alkoxy, and lower haloalkoxy; and the other of R^{24} and R^{25} is H.

In some embodiments of compound of Formula III or pharmaceutically acceptable salt thereof, Z^1 is N; R^{24} and R^{25} are each independently selected from halogen, lower alkyl, lower haloalkyl, lower alkoxy, and lower haloalkoxy.

In some embodiments, the compound of Formula III or pharmaceutically acceptable salt thereof is a compound of Formula IIIa



IIIa

or pharmaceutically acceptable salt thereof, wherein:

R^{24} is selected from H, halogen, lower alkyl, lower haloalkyl, lower alkoxy, and lower haloalkoxy; and

R^{25} is selected from H, halogen, lower alkyl, lower haloalkyl, lower alkoxy, and lower haloalkoxy;

provided that at least one of R^{24} and R^{25} is selected from halogen, lower alkyl, lower haloalkyl, lower alkoxy, and lower haloalkoxy.

In some embodiments of compound of Formula IIIa or pharmaceutically acceptable salt thereof, R^{24} is H; and R^{25} is selected from halogen, lower alkyl, lower haloalkyl, lower alkoxy, and lower haloalkoxy.

In some embodiments of compound of Formula IIIa or pharmaceutically acceptable salt thereof, R^{24} is H; and R^{25} is selected from lower alkyl and lower haloalkyl.

In some embodiments of compound of Formula IIIa or pharmaceutically acceptable salt thereof, R^{24} is H; and R^{25} is selected from C_{1-3} alkyl and C_{1-3} haloalkyl. In some further embodiments, R^{25} is selected from C_{1-2} alkyl and C_{1-2} haloalkyl. In yet further embodiments, R^{25} is selected from methyl and C_1 haloalkyl.

In some embodiments of compound of Formula IIIa or pharmaceutically acceptable salt thereof, R^{24} is H; and R^{25} is selected from C_{1-3} alkyl. In some further embodiments, R^{25} is selected from methyl or

ethyl. In yet further embodiments, R²⁵ is methyl.

In some embodiments of compound of Formula IIIa or pharmaceutically acceptable salt thereof, R²⁵ is H; and R²⁴ is selected from halogen, lower alkyl, lower haloalkyl, lower alkoxy, and lower haloalkoxy.

In some embodiments of compound of Formula IIIa or pharmaceutically acceptable salt thereof, R²⁵ is H; and R²⁴ is selected from lower alkyl and lower haloalkyl.

In some embodiments of compound of Formula IIIa or pharmaceutically acceptable salt thereof, R²⁵ is H; and R²⁴ is selected from C₁₋₃ alkyl and C₁₋₃ haloalkyl. In some further embodiments, R²⁴ is selected from C₁₋₂ alkyl and C₁₋₂ haloalkyl. In yet further embodiments, R²⁴ is selected from methyl and C₁ haloalkyl.

In some embodiments of compound of Formula IIIa or pharmaceutically acceptable salt thereof, R²⁵ is H; and R²⁴ is selected from C₁₋₃ alkyl. In some further embodiments, R²⁴ is selected from methyl or ethyl. In yet further embodiments, R²⁴ is methyl.

In some embodiments of compound of Formula IIIa or pharmaceutically acceptable salt thereof, R²⁴ and R²⁵ are each independently selected from halogen, lower alkyl, lower haloalkyl, lower alkoxy, and lower haloalkoxy.

In some embodiments of compound of Formula IIIa or pharmaceutically acceptable salt thereof, R²⁴ and R²⁵ are each independently selected from lower alkyl and lower haloalkyl.

In some embodiments of compound of Formula IIIa or pharmaceutically acceptable salt thereof, R²⁴ and R²⁵ are each independently selected from C₁₋₃ alkyl and C₁₋₃ haloalkyl. In some further embodiments, R²⁴ and R²⁵ are each independently selected from C₁₋₂ alkyl and C₁₋₂ haloalkyl. In yet further embodiments, R²⁴ and R²⁵ are each independently selected from methyl and C₁ haloalkyl.

In some embodiments of compound of Formula IIIa or pharmaceutically acceptable salt thereof, R²⁴ and R²⁵ are each independently selected from C₁₋₃ alkyl. In some further embodiments, R²⁴ and R²⁵ are each independently selected from methyl or ethyl. In yet further embodiments, R²⁴ and R²⁵ are each methyl.

Also provided herein is a pharmaceutical composition comprising at least one pharmaceutically acceptable excipient, adjuvant, or carrier, and a therapeutically effective amount of at least one compound described herein.

Further provided are pharmaceutical compositions comprising at least one pharmaceutically acceptable excipient, adjuvant, or carrier, and a therapeutically effective amount of at least one compound

described herein in combination with at least one additional compound such as an erythropoiesis stimulating agent or chemotherapeutic agent.

Additionally provided herein is a method of by administering to the subject at least one compound described herein or pharmaceutically acceptable salt thereof, or use of a compound described herein or pharmaceutically acceptable salt thereof in the manufacturing a medicament for, modulating (such as increasing) HIF levels or activity in a subject.

Further provided is a method of by administering to the subject at least one compound described herein or pharmaceutically acceptable salt thereof, or use of a compound described herein or pharmaceutically acceptable salt thereof in the manufacturing a medicament for, treating a condition where it is desired to modulate HIF activity comprising administering to a subject at least one compound described herein.

Also provided is a method of by administering to the subject at least one compound described herein or pharmaceutically acceptable salt thereof, or use of a compound described herein or pharmaceutically acceptable salt thereof in the manufacturing a medicament for, treating a anemic or ischemic related disorder in a subject comprising administering to a subject at least one compound described herein.

Also provided is a method of by administering to the subject at least one compound described herein or pharmaceutically acceptable salt thereof, or use of a compound described herein or pharmaceutically acceptable salt thereof in the manufacturing a medicament for, treating ischemia, anemia, wound healing, auto-transplantation, allo-transplantation, xeno-transplantation, systemic high blood pressure, thalassemia, diabetes, cancer or an inflammatory disorder, or a combination of two or more thereof in a subject comprising administering to a subject at least one compound described herein.

Also provided is a method of by administering to the subject at least one compound described herein or pharmaceutically acceptable salt thereof, or use of a compound described herein or pharmaceutically acceptable salt thereof in the manufacturing a medicament for, treating anemia in a subject comprising administering to a subject at least one compound described herein.

Further provided is a method of by administering to the subject at least one compound described herein or pharmaceutically acceptable salt thereof, or use of a compound described herein or pharmaceutically acceptable salt thereof in the manufacturing a medicament for, modulating the amount of HIF in a cell comprising contacting the cell with at least one compound described herein.

Additionally provided is a method of by administering to the subject at least one compound

described herein or pharmaceutically acceptable salt thereof, or use of a compound described herein or pharmaceutically acceptable salt thereof in the manufacturing a medicament for, increasing the amount of hemoglobin F in a subject comprising administering to the subject at least one compound described herein.

5 Also provided is a method of by administering to the subject at least one compound described herein or pharmaceutically acceptable salt thereof, or use of a compound described herein or pharmaceutically acceptable salt thereof in the manufacturing a medicament for, modulating angiogenesis in a subject comprising administering to the subject at least one compound described herein.

10 Additionally provided is a method of by administering to the subject at least one compound described herein or pharmaceutically acceptable salt thereof, or use of a compound described herein or pharmaceutically acceptable salt thereof in the manufacturing a medicament for, treating at least one disease in a patient in need of such treatment comprising administering to the patient a therapeutically effective amount of at least one compound described herein.

15 Also provided is a method of by administering to the subject at least one compound described herein or pharmaceutically acceptable salt thereof, or use of a compound described herein or pharmaceutically acceptable salt thereof in the manufacturing a medicament for, inhibiting HIF hydroxylation in a subject comprising administering to the subject at least one compound described herein.

Additional embodiments of the invention are set forth in the description which follows, or may be learned by practice of the invention.

20 Unless otherwise indicated, all numbers expressing quantities of ingredients, reaction conditions, and so forth used in the specification and claims are to be understood as being modified in all instances by the term "about." Accordingly, unless indicated to the contrary, the numerical parameters set forth in the following specification and attached claims are approximations that may vary depending upon the standard deviation found in their respective testing measurements.

25 As used herein, when any variable occurs more than one time in a chemical formula, its definition on each occurrence is independent of its definition at every other occurrence. The compounds of the present disclosure may contain one or more chiral centers and/or double bonds and therefore, may exist as stereoisomers, such as double-bond isomers (i.e., geometric isomers), enantiomers or diastereomers. Accordingly, any chemical structures within the scope of the specification depicted, in whole or in part,
30 with a relative configuration encompass all possible enantiomers and stereoisomers of the illustrated

compounds including the stereoisomerically pure form (e.g., geometrically pure, enantiomerically pure or diastereomerically pure) and enantiomeric and stereoisomeric mixtures. Enantiomeric and stereoisomeric mixtures can be resolved into the component enantiomers or stereoisomers using separation techniques or chiral synthesis techniques well known to the skilled artisan.

5 Compounds of Formula I, II, IIa, III and IIIa include, but are not limited to optical isomers of compounds of Formula I, II, IIa, III and IIIa, racemates, and other mixtures thereof. In those situations, the single enantiomers or diastereomers, i.e., optically active forms, can be obtained by asymmetric synthesis or by resolution of the racemates. Resolution of the racemates can be accomplished, for example, by conventional methods such as crystallization in the presence of a resolving agent, or
10 chromatography, using, for example a chiral high-pressure liquid chromatography (HPLC) column. In addition, compounds of Formula I, II, IIa, III and IIIa include Z- and E- forms (or cis- and trans- forms) of compounds with double bonds. Where compounds of Formula I, II, IIa, III and IIIa exists in various tautomeric forms, chemical entities of the present invention include all tautomeric forms of the compound.

15 Compounds of the present disclosure include, but are not limited to compounds of Formula I and all pharmaceutically acceptable forms thereof. Pharmaceutically acceptable forms of the compounds recited herein include pharmaceutically acceptable salts, solvates, crystal forms (including polymorphs and clathrates), chelates, non-covalent complexes, prodrugs, and mixtures thereof. In certain embodiments, the compounds described herein are in the form of pharmaceutically acceptable salts. As used henceforth,
20 the term "compound" encompasses not only the compound itself, but also a pharmaceutically acceptable salt thereof, a solvate thereof, a chelate thereof, a non-covalent complex thereof, a prodrug thereof, and mixtures of any of the foregoing.

As noted above, prodrugs also fall within the scope of chemical entities, for example, ester or amide derivatives of the compounds of Formula I, II, IIa, III and IIIa. The term "prodrugs" includes any
25 compounds that become compounds of Formula I, II, IIa, III and IIIa when administered to a patient, e.g., upon metabolic processing of the prodrug. Examples of prodrugs include, but are not limited to, acetate, formate, and benzoate and like derivatives of functional groups (such as alcohol or amine groups) in the compounds of Formula I, II, IIa, III and IIIa.

The term "solvate" refers to the compound formed by the interaction of a solvent and a compound.
30 Suitable solvates are pharmaceutically acceptable solvates, such as hydrates, including monohydrates and

hemi-hydrates.

At various places in the present specification, substituents of compounds of the invention are disclosed in groups or in ranges. It is specifically intended that the invention include each and every individual subcombination of the members of such groups and ranges. For example, the term "C₁₋₃ alkyl" is specifically intended to individually disclosed methyl, ethyl, and C₃ alkyl (including n-propyl and isopropyl).

For compounds of the invention in which a variable appears more than once, each variable can be a different moiety selected from the Markush group defining the variable. For example, where a structure is described having two R groups that are simultaneously present on the same compound; the two R groups can represent different moieties selected from the Markush group defined for R.

It is further appreciated that certain features of the invention, which are, for clarity, described in the context of separate embodiments, can also be provided in combination in a single embodiment. Conversely, various features of the invention which are, for brevity, described in the context of a single embodiment, can also be provided separately or in any suitable subcombination.

The term "n-membered" where n is an integer typically describes the number of ring-forming atoms in a moiety where the number of ring-forming atoms is n. For example, pyridine is an example of a 6-membered heteroaryl ring and thiophene is an example of a 5-membered heteroaryl group.

"Alkyl" refers to a saturated, branched or straight-chain monovalent hydrocarbon group derived by the removal of one hydrogen atom from a single carbon atom of a parent alkane. Typical alkyl groups include, but are not limited to, methyl, ethyl, propyls such as propan-1-yl, propan-2-yl, and cyclopropan-1-yl, butyls such as butan-1-yl, butan-2-yl, 2-methyl-propan-1-yl, 2-methyl-propan-2-yl, cyclobutan-1-yl, tert-butyl, and the like. In certain embodiments, an alkyl group comprises from 1 to 20 carbon atoms. As used herein the term "lower alkyl" refers to an alkyl group comprising from 1 to 6 carbon atoms.

"Alkenyl" refers to an unsaturated branched, straight-chain or cyclic alkyl group having at least one carbon-carbon double bond derived by the removal of one hydrogen atom from a single carbon atom of a parent alkene. The group may be in either the Z- or E- forms (or *cis* or *trans* conformation) about the double bond(s). Typical alkenyl groups include, but are not limited to, ethenyl; propenyls such as prop-1-en-1-yl, prop-1-en-2-yl, prop-2-en-1-yl (allyl), prop-2-en-2-yl, cycloprop-1-en-1-yl; cycloprop-2-en-1-yl; butenyls such as but-1-en-1-yl, but-1-en-2-yl, 2-methyl-prop-1-en-1-yl,

but-2-en-1-yl, but-2-en-2-yl, buta-1,3-dien-1-yl, buta-1,3-dien-2-yl, cyclobut-1-en-1-yl, cyclobut-1-en-3-yl, cyclobuta-1,3-dien-1-yl; and the like. In certain embodiments, an alkenyl group has from 2 to 20 carbon atoms and in other embodiments, from 2 to 6 carbon atoms, i.e. "lower alkenyl."

"Alkynyl" refers to an unsaturated branched or straight-chain having at least one carbon-carbon triple bond derived by the removal of one hydrogen atom from a single carbon atom of a parent alkyne. Typical alkynyl groups include, but are not limited to, ethynyl; propynyl; butynyl, 2-pentynyl, 3-pentynyl, 2-hexynyl, 3-hexynyl and the like. In certain embodiments, an alkynyl group has from 2 to 20 carbon atoms and in other embodiments, from 2 to 6 carbon atoms (i.e. "lower alkynyl").

"Alkoxy" refers to a radical -OR where R represents an alkyl. Representative examples include, but are not limited to, methoxy, ethoxy, propoxy, butoxy, cyclohexyloxy, and the like.

"Alkoxy carbonyl" refers to a radical -C(O)-OR where R represents an alkyl as defined herein.

"Aryl" refers to a monovalent aromatic hydrocarbon group derived by the removal of one hydrogen atom from a single carbon atom of a parent aromatic ring system. Aryl encompasses 5- and 6-membered carbocyclic aromatic rings, for example, benzene; bicyclic ring systems wherein at least one ring is carbocyclic and aromatic, for example, naphthalene, indane, and tetralin; and tricyclic ring systems wherein at least one ring is carbocyclic and aromatic, for example, fluorene. For example, aryl includes 5- and 6-membered carbocyclic aromatic rings fused to a 5- to 7-membered heterocycloalkyl ring containing 1 or more heteroatoms selected from N, O, and S. In certain embodiments, an aryl group can comprise from 6 to 10 carbon atoms. Aryl, however, does not encompass or overlap in any way with heteroaryl, separately defined below. Hence, if one or more carbocyclic aromatic rings is fused with a heterocycloalkyl aromatic ring, the resulting ring system is heteroaryl, not aryl, as defined herein.

"Arylalkyl" or "aralkyl" refers to an acyclic alkyl group in which one of the hydrogen atoms bonded to a carbon atom, typically a terminal or *sp*³ carbon atom, is replaced with an aryl group. Typical arylalkyl groups include, but are not limited to, benzyl, 2-phenylethan-1-yl, 2-phenylethen-1-yl, naphthylmethyl, 2-naphthylethan-1-yl, 2-naphthylethen-1-yl, naphthobenzyl, 2-naphthophenylethan-1-yl and the like. Where specific alkyl moieties are intended, the nomenclature arylalkyl, arylalkenyl, and/or arylalkynyl is used. In certain embodiments, an arylalkyl group can be (C₆₋₃₀) arylalkyl, e.g., the alkyl group of the arylalkyl group can be (C₁₋₁₀) and the aryl moiety can be (C₅₋₂₀).

"Carbonyl" refers to a radical -C(O) group.

"Carboxy" refers to the radical -C(O)OH.

"Cyano" refers to the radical -CN.

"Cycloalkyl" refers to a saturated or unsaturated, but non-aromatic, cyclic alkyl group. Where a specific level of saturation is intended, the nomenclature "cycloalkanyl" or "cycloalkenyl" is used. Typical cycloalkyl groups include, but are not limited to, groups derived from cyclopropane, cyclobutane, cyclopentane, cyclohexane, and the like. In certain embodiments, the cycloalkyl group can be C₃₋₁₀ cycloalkyl, such as, for example, C₃₋₆ cycloalkyl.

"Heterocycloalkyl" refers to a saturated or unsaturated, but non-aromatic, cyclic alkyl group in which one or more carbon atoms (and any associated hydrogen atoms) are independently replaced with the same or different heteroatom and its associated hydrogen atoms, where appropriate. Typical heteroatoms to replace the carbon atom(s) include, but are not limited to, N, P, O, S, and Si. Where a specific level of saturation is intended, the nomenclature "heterocycloalkanyl" or "heterocycloalkenyl" is used. Typical heterocycloalkyl groups include, but are not limited to, groups derived from epoxides, imidazolidine, morpholine, piperazine, piperidine, pyrazolidine, pyrrolidine, quinuclidine, tetrahydrofuran, tetrahydropyran and the like. Substituted heterocycloalkyl also includes ring systems substituted with one or more oxo (=O) or oxide (-O⁻) substituents, such as piperidinyl N-oxide, morpholinyl-N-oxide, 1-oxo-1-thiomorpholinyl and 1,1-dioxo-1-thiomorpholinyl.

"Disease" refers to any disease, disorder, condition, symptom, or indication.

"Halo" refers to a fluoro, chloro, bromo, or iodo group.

"Heteroaryl" refers to a monovalent heteroaromatic group derived by the removal of one hydrogen atom from a single atom of a parent heteroaromatic ring system. Heteroaryl encompasses: 5- to 7-membered aromatic, monocyclic rings containing one or more, for example, from 1 to 4, or in certain embodiments, from 1 to 3, heteroatoms selected from N, O, and S, with the remaining ring atoms being carbon; and polycyclic heterocycloalkyl rings containing one or more, for example, from 1 to 4, or in certain embodiments, from 1 to 3, heteroatoms selected from N, O, and S, with the remaining ring atoms being carbon and wherein at least one heteroatom is present in an aromatic ring.

For example, heteroaryl includes a 5- to 7-membered heteroaromatic ring fused to a 5- to 7-membered cycloalkyl ring and a 5- to 7-membered heteroaromatic ring fused to a 5- to 7-membered heterocycloalkyl ring. For such fused, bicyclic heteroaryl ring systems wherein only one of the rings contains one or more heteroatoms, the point of attachment may be at the heteroaromatic ring or the cycloalkyl ring. When the total number of S and O atoms in the heteroaryl group exceeds 1, those

heteroatoms are not adjacent to one another. In certain embodiments, the total number of S and O atoms in the heteroaryl group is not more than 2. In certain embodiments, the total number of S and O atoms in the aromatic heterocycle is not more than 1. Heteroaryl does not encompass or overlap with aryl as defined above. Typical heteroaryl groups include, but are not limited to, groups derived from acridine, 5 arsindeole, carbazole, β -carboline, chromane, chromene, cinnoline, furan, imidazole, indazole, indole, indoline, indolizine, isobenzofuran, isochromene, isoindole, isoindoline, isoquinoline, isothiazole, isoxazole, naphthyridine, oxadiazole, oxazole, perimidine, phenanthridine, phenanthroline, phenazine, phthalazine, pteridine, purine, pyran, pyrazine, pyrazole, pyridazine, pyridine, pyrimidine, pyrrole, pyrrolizine, quinazoline, quinoline, quinolizine, quinoxaline, tetrazole, thiadiazole, thiazole, thiophene, 10 triazole, xanthene, and the like. In certain embodiments, the heteroaryl group can be between 5 to 20 membered heteroaryl, such as, for example, a 5 to 10 membered heteroaryl. In certain embodiments, heteroaryl groups can be those derived from thiophene, pyrrole, benzothiophene, benzofuran, indole, pyridine, quinoline, imidazole, oxazole, and pyrazine.

"Heteroarylalkyl" or "heteroaralkyl" refers to an acyclic alkyl group in which one of the hydrogen 15 atoms bonded to a carbon atom, typically a terminal or sp^3 carbon atom, is replaced with a heteroaryl group. Where specific alkyl moieties are intended, the nomenclature heteroarylalkanyl, heteroarylalkenyl, and/or heteroarylalkynyl is used. In certain embodiments, the heteroarylalkyl group can be a 6 to 30 membered heteroarylalkyl, e.g., the alkanyl, alkenyl or alkynyl moiety of the heteroarylalkyl can be 1 to 10 membered and the heteroaryl moiety can be a 5 to 20-membered heteroaryl.

20 "Sulfonyl" refers to a radical $-S(O)_2R$ where R is an alkyl, substituted alkyl, substituted cycloalkyl, substituted heterocycloalkyl, substituted aryl, or substituted heteroaryl group as defined herein. Representative examples include, but are not limited to methylsulfonyl, ethylsulfonyl, propylsulfonyl, butylsulfonyl, and the like.

"Sulfanyl" refers to a radical $-SR$ where R is an alkyl, substituted alkyl, substituted cycloalkyl, 25 substituted heterocycloalkyl, substituted aryl, or substituted heteroaryl group as defined herein that may be optionally substituted as defined herein. Representative examples include, but are not limited to, methylthio, ethylthio, propylthio, butylthio, and the like.

"Pharmaceutically acceptable" refers to generally recognized for use in animals, and more particularly in humans.

30 "Pharmaceutically acceptable salt" refers to a salt of a compound that is pharmaceutically acceptable

and that possesses the desired pharmacological activity of the parent compound. Such salts include: (1) acid addition salts, formed with inorganic acids such as hydrochloric acid, hydrobromic acid, sulfuric acid, nitric acid, phosphoric acid, and the like; or formed with organic acids such as acetic acid, propionic acid, hexanoic acid, cyclopentanepropionic acid, glycolic acid, pyruvic acid, lactic acid, malonic acid, succinic acid, malic acid, maleic acid, fumaric acid, tartaric acid, citric acid, benzoic acid, 3-(4-hydroxybenzoyl) benzoic acid, cinnamic acid, mandelic acid, methanesulfonic acid, and the like; or (2) salts formed when an acidic proton present in the parent compound either is replaced by a metal ion, e.g., an alkali metal ion, an alkaline earth ion, or an aluminum ion; or coordinates with an organic base such as ethanolamine, diethanolamine, triethanolamine, N-methylglucamine, dicyclohexylamine, and the like.

"Pharmaceutically acceptable excipient," "pharmaceutically acceptable carrier," or "pharmaceutically acceptable adjuvant" refer, respectively, to an excipient, carrier or adjuvant with which at least one compound of the present disclosure is administered.

"Pharmaceutically acceptable vehicle" refers to any of a diluent, adjuvant, excipient or carrier with which at least one compound of the present disclosure is administered.

"Stereoisomer" refers to an isomer that differs in the arrangement of the constituent atoms in space. Stereoisomers that are mirror images of each other and optically active are termed "enantiomers," and stereoisomers that are not mirror images of one another and are optically active are termed "diastereoisomers."

"Subject" includes mammals and humans. The terms "human" and "subject" are used interchangeably herein.

"Substituted" refers to a group in which one or more hydrogen atoms are each independently replaced with the same or different substituent(s). Typical substituents include, but are not limited to, -X, -R₃₃, -OH, =O, -OR₃₃, SR₃₃, -SH, =S, -NR₃₃R₃₄, =NR₃₃, -CX₃, -CF₃, -CN, -NO₂, -S(O)₂R₃₃, -OS(O₂)OH, -OS(O)₂R₃₃, -OP(O)(OR₃₃)(OR₃₄), -C(O)R₃₃, -C(S)R₃₃, -C(O)OR₃₃, -C(O)NR₃₃R₃₄, -C(O)OH, -C(S)OR₃₃, -NR₃₅C(O)NR₃₃R₃₄, -NR₃₅C(S)NR₃₃R₃₄, -NR₃₅C(NR₃₃)NR₃₃R₃₄, -C(NR₃₃)NR₃₃R₃₄, -S(O)₂NR₃₃R₃₄, -NR₃₅S(O)₂R₃₃, -NR₃₅C(O)R₃₃, and S(O)R₃₃ where each X is independently a halo; each R₃₃ and R₃₄ are independently hydrogen, alkyl, substituted alkyl, aryl, substituted aryl, arylalkyl, substituted arylalkyl, cycloalkyl, substituted cycloalkyl, heterocycloalkyl, substituted heterocycloalkyl, heteroaryl, substituted heteroaryl, heteroarylalkyl, substituted heteroarylalkyl, -NR₃₅R₃₆, -C(O)R₃₅ or -S(O)₂R₃₅ or optionally R₃₃

and R₃₄ together with the atom to which R₃₃ and R₃₄ are attached form one or more heterocycloalkyl, substituted heterocycloalkyl, heteroaryl, or substituted heteroaryl rings; and R₃₅ and R₃₆ are independently hydrogen, alkyl, substituted alkyl, aryl, substituted aryl, arylalkyl, substituted arylalkyl, cycloalkyl, substituted cycloalkyl, heterocycloalkyl, substituted heterocycloalkyl, heteroaryl, substituted heteroaryl, heteroarylalkyl or substituted heteroarylalkyl, or optionally R₃₅ and R₃₆ together with the nitrogen atom to which R₃₅ and R₃₆ are attached form one or more heterocycloalkyl, substituted heterocycloalkyl, heteroaryl, or substituted heteroaryl rings. In certain embodiments, a tertiary amine or aromatic nitrogen may be substituted with one or more oxygen atoms to form the corresponding nitrogen oxide.

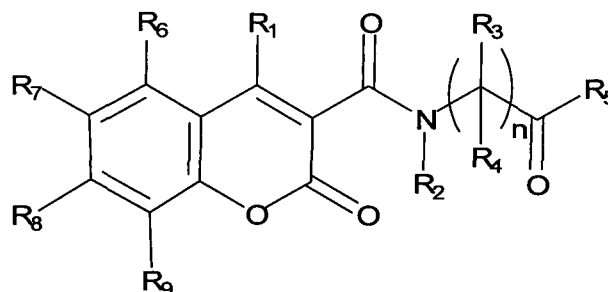
"Therapeutically effective amount" refers to the amount of a compound that, when administered to a subject for treating a disease, or at least one of the clinical symptoms of a disease or disorder, is sufficient to affect such treatment for the disease, disorder, or symptom. The "therapeutically effective amount" can vary depending on the compound, the disease, disorder, and/or symptoms of the disease or disorder, severity of the disease, disorder, and/or symptoms of the disease or disorder, the age of the subject to be treated, and/or the weight of the subject to be treated. An appropriate amount in any given instance can be readily apparent to those skilled in the art or capable of determination by routine experimentation.

"Treating" or "treatment" of any disease or disorder refers to arresting or ameliorating a disease, disorder, or at least one of the clinical symptoms of a disease or disorder, reducing the risk of acquiring a disease, disorder, or at least one of the clinical symptoms of a disease or disorder, reducing the development of a disease, disorder or at least one of the clinical symptoms of the disease or disorder, or reducing the risk of developing a disease or disorder or at least one of the clinical symptoms of a disease or disorder. "Treating" or "treatment" also refers to inhibiting the disease or disorder, either physically, (e.g., stabilization of a discernible symptom), physiologically, (e.g., stabilization of a physical parameter), or both, or inhibiting at least one physical parameter which may not be discernible to the subject. Further, "treating" or "treatment" refers to delaying the onset of the disease or disorder or at least symptoms thereof in a subject which may be exposed to or predisposed to a disease or disorder even though that subject does not yet experience or display symptoms of the disease or disorder.

Reference will now be made in detail to embodiments of the present disclosure. While certain embodiments of the present disclosure will be described, it will be understood that it is not intended to limit the embodiments of the present disclosure to those described embodiments. To the contrary,

reference to embodiments of the present disclosure is intended to cover alternatives, modifications, and equivalents as may be included within the spirit and scope of the embodiments of the present disclosure as defined by the appended claims.

Certain embodiments of the present invention are directed to at least one compound of Formula I:



I

a pharmaceutically acceptable salt thereof, a solvate thereof, a chelate thereof, a non-covalent complex thereof, a prodrug thereof, and mixtures of any of the foregoing, wherein:

n is 1 to 6;

R₁ is selected from OH, SH, NR₃R₄, NHC(O)R₂, NHSO₂R₂ and sulfonyl,;

R₂ is selected from H, lower alkyl and substituted lower alkyl;

R₃ and R₄ are independently selected from H, lower alkyl, substituted lower alkyl, lower haloalkyl, substituted lower haloalkyl, or R₃ and R₄ can join together to form a 3 to 6 membered ring or a substituted 3 to 6 membered ring;

R₅ is selected from OH, SH, NH₂, lower alkyl, substituted lower alkyl, lower alkoxy, substituted lower alkoxy, and sulfanyl;

each of R₆, R₇, R₈ and R₉ is independently selected from H, alkyl, substituted alkyl, alkenyl, substituted alkenyl, alkynyl, substituted alkynyl, alkoxy, substituted alkoxy, NR₃R₄, C(O)OH, OR₁₂, SR₁₂, SO₂R₁₂, CN, NO₂, halogen, aryl, substituted aryl, heteroaryl, substituted heteroaryl, arylalkyl, substituted arylalkyl, heteroarylalkyl, substituted heteroarylalkyl, heterocycloalkyl, substituted heterocycloalkyl, alkylsilyl, substituted alkylsilyl, alkenylsilyl, substituted alkenylsilyl, alkynylsilyl, substituted alkynylsilyl, alkoxycarbonyl, substituted alkoxycarbonyl, and -X-R₁₁,;

or at least one of adjacent pairs R₁ and R₆, R₆ and R₇, R₇ and R₈, and R₈ and R₉, can join together to form a 4 to 7 membered ring or a substituted 4 to 7 membered ring;

X is selected from -N(R₁₀)-Y- and -Y-N(R₁₀)-;

Y is selected from C(O), SO₂, alkylene, substituted alkylene, alkenylene, substituted alkenylene,

alkynylene, and substituted alkynylene;

R₁₀ is selected from H, lower alkyl, and substituted lower alkyl,

R₁₁ is selected from H, cycloalkyl, substituted cycloalkyl, heterocycloalkyl, substituted heterocycloalkyl, aryl, substituted aryl, heteroaryl, and substituted heteroaryl; and

5 R₁₂ is selected from H, alkyl, substituted alkyl, alkenyl, substituted alkenyl, alkynyl, substituted alkynyl and NR₃R₄.

In certain embodiments of compounds of Formula I, R₁ is selected from OH and SH.

In certain embodiments of compounds of Formula I, R₂ is H.

In certain embodiments of compounds of Formula I, R₃ and R₄ are each independently selected from
10 H, lower alkyl such as methyl or ethyl, and substituted lower alkyl (such as lower alkyl substituted with hydroxyl, for example hydroxymethyl).

In certain embodiments of compounds of Formula I, R₅ is selected from OH, a lower alkoxy such as methoxy, ethoxy and propoxy, and a substituted lower alkoxy.

In certain embodiments of compounds of Formula I, R₃ and R₄ join together to form a 3 to 6
15 membered ring or a substituted 3 to 6 membered ring. The 3 to 6 membered rings can comprise at least one heteroatom, such as at least two heteroatoms.

In certain embodiments of compounds of Formula I, R₁ and R₆ can join together to form a 4 to 7 membered ring or a substituted 4 to 7 membered ring. The 4 to 7 membered rings can comprise at least one heteroatom, such as at least two heteroatoms, and at least three heteroatoms.

20 In certain embodiments of compounds of Formula I, at least one of R₆, R₇, R₈ and R₉ is independently selected from halo and a moiety substituted with at least one halogen, such as trifluoromethyl.

In certain embodiments of compounds of Formula I, at least one of R₆, R₇, R₈ and R₉ is independently selected from alkoxy or substituted alkoxy.

25 In certain embodiments of compounds of Formula I, at least one of R₆, R₇, R₈ and R₉ is independently selected from alkylsilyl, substituted alkylsilyl, alkynylsilyl, and substituted alkynylsilyl.

In certain embodiments of compounds of Formula I, at least one of R₆, R₇, R₈ and R₉ is independently selected from aryl, substituted aryl, heteroaryl, substituted heteroaryl, heterocycloalkyl, and substituted heterocycloalkyl, such as substituted pyridines, substituted pyrimidines, substituted
30 pyrazines, substituted pyridazines, substituted tetrahydrofurans, and substituted piperidines.

In certain embodiments of compounds of Formula I, at least one of R₆, R₇, R₈ and R₉ is independently selected from H, alkyl, substituted alkyl, alkenyl, substituted alkenyl, alkynyl, and substituted alkynyl, such as isopropyl, cyclohexane, cyclopentane, cyclohexene and cyclopentene.

Examples of individual representative compounds of the present disclosure, and compounds
5 comprised in compositions of the present disclosure, and used in methods of the present disclosure are listed in Table 1. Each compound listed in Table 1, i.e., Examples 1-38, contains information directed to its structure, name, molecular weight, hydrogen NMR data and at least one method of synthesis.

In certain embodiments, compounds of the present disclosure inhibit prolyl hydroxylases such as HIF prolyl hydroxylases. Variety of assays may be used to determine the prolyl hydroxylase inhibitory
10 activity of a compound.

In certain embodiments, compounds of the present disclosure modulate HIF levels or activity, for example, by stabilizing HIF.

Furthermore, compounds of the present disclosure can contain one or more chiral centers. Such compounds can be prepared or isolated as pure stereoisomers, i.e., as individual enantiomers or
15 diastereomers, or as stereoisomer-enriched mixtures. All such stereoisomers, and enriched mixtures thereof, are included within the scope of the present disclosure. Pure stereoisomers, and enriched mixtures thereof, can be prepared using, for example, optically active starting materials or stereoselective reagents well-known in the art. Alternatively, racemic mixtures of such compounds can be separated using, for example, chiral column chromatography, chiral resolving agents and the like.

Certain embodiments of the present disclosure are directed to a pharmaceutical composition
20 comprising at least one pharmaceutically acceptable excipient, adjuvant or carrier, and a therapeutically effective amount of at least one compound described herein. The at least one compound can be present in an amount effective for the treatment of at least one disease selected from ischemia, anemia, wound healing, auto-transplantation, allo-transplantation, xeno-transplantation, systemic high blood pressure, thalassemia, diabetes, cancer and an inflammatory disorder.
25

Other embodiments of the present disclosure are directed to a method of treating a condition where it is desired to modulate HIF activity comprising administering to a subject at least one compound described herein. The condition can be selected from at least one of ischemia, anemia, wound healing, auto-transplantation, allo-transplantation, xenotransplantation, systemic high blood pressure, thalassemia,
30 diabetes, cancer and an inflammatory disorder.

A further embodiment is directed to a method of treating at least one disease in a patient in need of such treatment comprising administering to the patient a therapeutically effective amount of at least one compound described herein. The at least one disease can be selected from ischemia, anemia, wound healing, auto-transplantation, allo-transplantation, xeno-transplantation, systemic high blood pressure, thalassemia, diabetes, cancer and an inflammatory disorder.

Synthesis

Compounds of the invention, including salts thereof, can be prepared using known organic synthesis techniques and can be synthesized according to any of numerous possible synthetic routes.

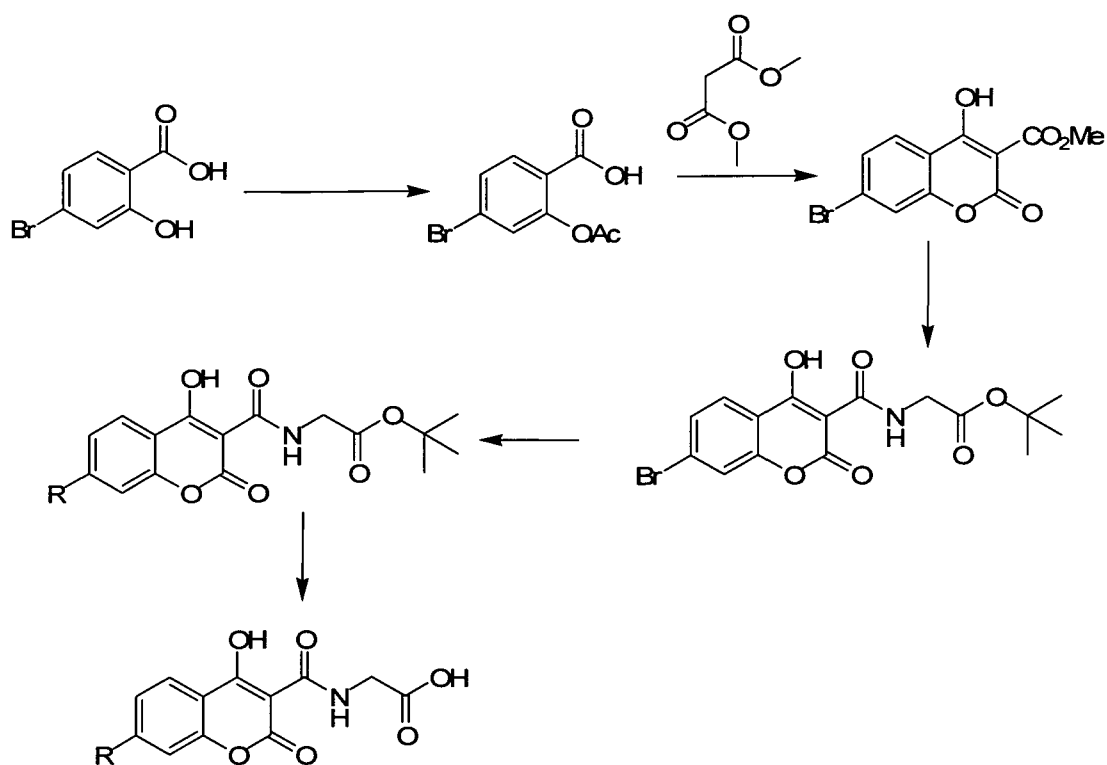
The reactions for preparing compounds of the invention can be carried out in suitable solvents which can be readily selected by one of skill in the art of organic synthesis. Suitable solvents can be substantially non-reactive with the starting materials (reactants), the intermediates, or products at the temperatures at which the reactions are carried out, *e.g.*, temperatures which can range from the solvent's freezing temperature to the solvent's boiling temperature. A given reaction can be carried out in one solvent or a mixture of more than one solvent. Depending on the particular reaction step, suitable solvents for a particular reaction step can be selected by the skilled artisan.

Preparation of compounds of the invention can involve the protection and deprotection of various chemical groups. The need for protection and deprotection, and the selection of appropriate protecting groups, can be readily determined by one skilled in the art. The chemistry of protecting groups can be found, for example, in T. W. Greene and P. G. M. Wuts, *Protective Groups in Organic Synthesis*, 3rd Ed., Wiley & Sons, Inc., New York (1999), which is incorporated herein by reference in its entirety.

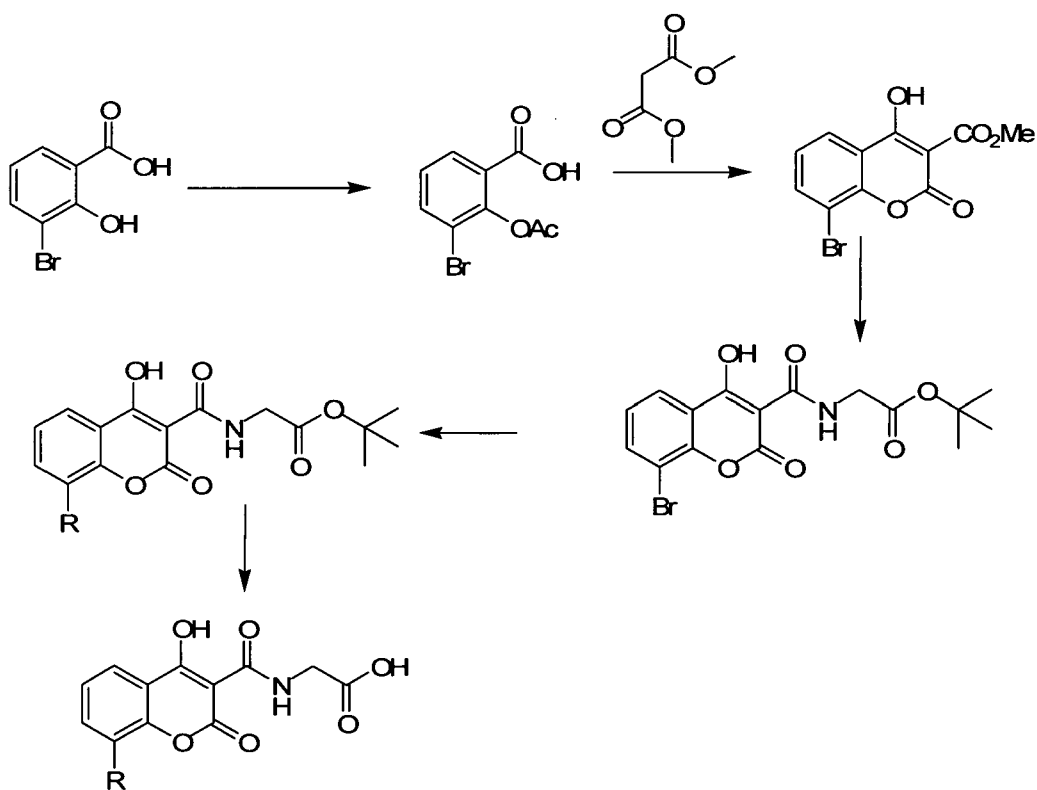
Reactions can be monitored according to any suitable method known in the art. For example, product formation can be monitored by spectroscopic means, such as nuclear magnetic resonance spectroscopy (*e.g.*, ¹H or ¹³C), infrared spectroscopy, spectrophotometry (*e.g.*, UV-visible), mass spectrometry, or by chromatographic methods such as high performance liquid chromatography (HPLC) or thin layer chromatography (TLC).

The compounds of invention can be prepared, for example, according to one or more of the following general reaction schemes and techniques described below.

General Scheme I

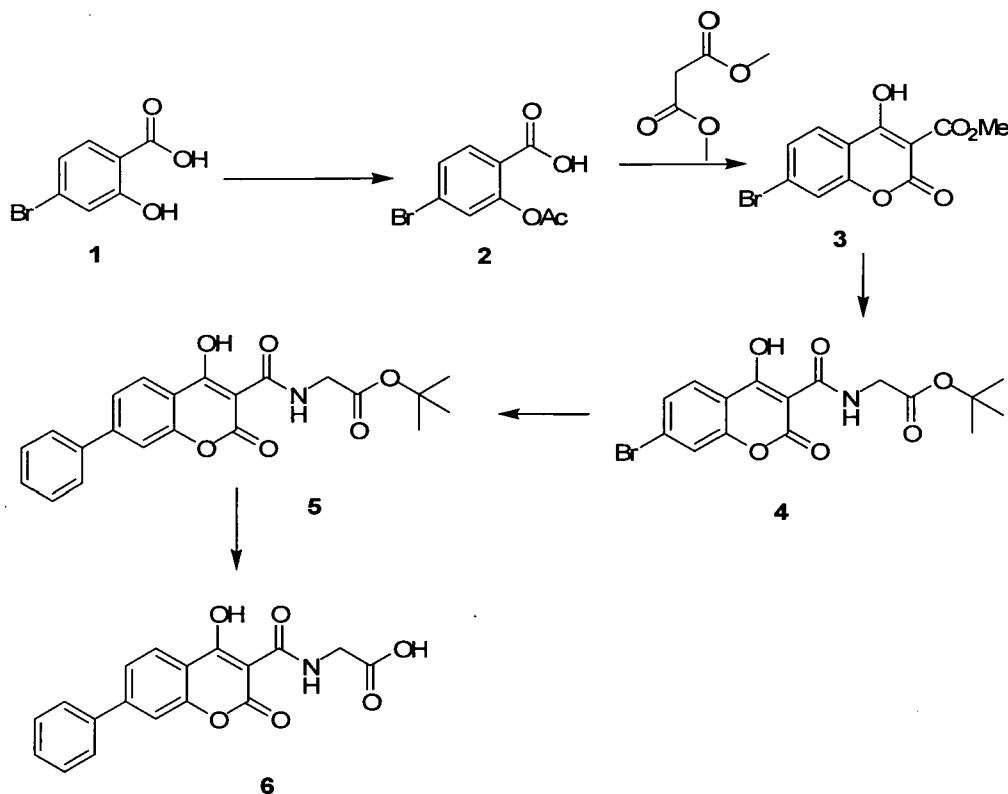


General Scheme II



5 Examples

The present invention is further exemplified, but not limited, by the following examples that illustrate the preparation of compounds of Formulas I, II, IIa, III, or IIIa, according to the invention.

Example 1**Synthesis of N-[(4-hydroxy-2-oxo-7-phenyl-2H-3-chromenyl)carbonyl]glycine**

Reagent 1 (42 g, 33.6 mmol) was dissolved in 200 mL of AC_2O with addition of 1 mL of H_3PO_4 , and was heated at 50°C . After its completion, the reaction mixture was cooled to room temperature, and 500 mL of water was then added in and stirred at 50°C until hydrolyzation is complete. The reaction mixture was then cooled to 0°C and filtered. The resulting solid, Compound 2, was further dried (47 g).

Compound 2 (25.9 g, 100 mmol) and 1-Hydroxybenzotriazole anhydrous (HOBt, 13.5 g) was dissolved in 400 mL of THF. Dicyclohexylcarbodiimide (DCC, 20.9 g) was gradually added in at 0°C and stirred over night at a temperature below 10°C . The reaction mixture was filtered and the Filtrate A was collected. 13.2 g (100 mmol) of dimethyl malonate was dissolved in 800 mL of THF first, then 7.2 g sodium hydride (70% dispersion) was added in. Filtrate A was added while stirred and left to react for 2 hours at room temperature. After THF removed by vacuum distillation, 400 mL of methanol and 400 mL of 10% HCl were added and stirred over night. After filtration, the resulting solid was washed with 400 mL of methanol and dried, yielding 14 g of Compound 3.

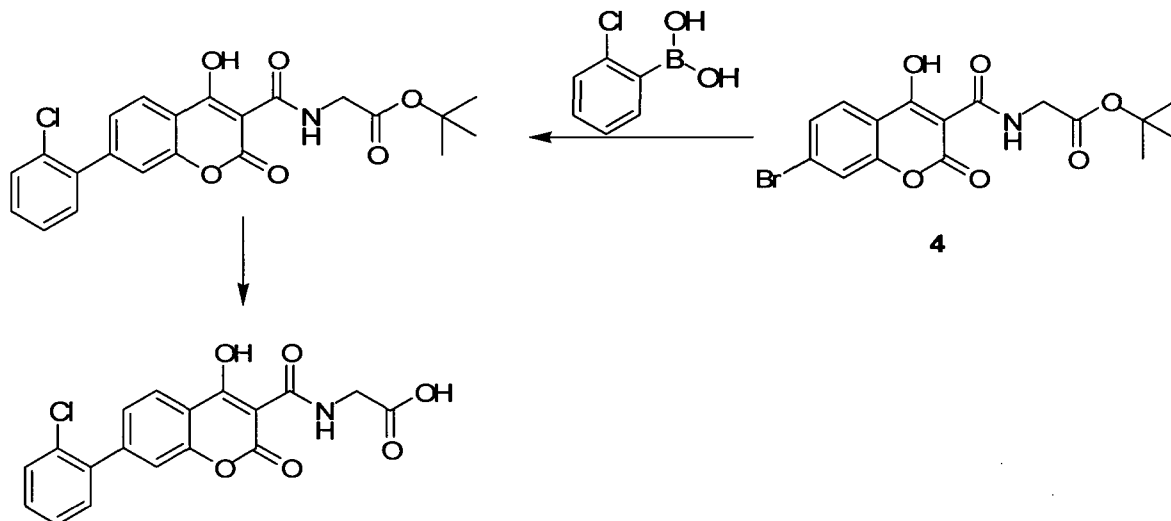
Glycine t-butyl ester HCl salt (13.4 g) and sodium methoxide (4.4 g) were suspended in 200 mL of methanol. After stirred to a homogeneous suspension, it was distilled to remove all methanol. 200 mL of THF and Compound 3 (6.0 g) were added in and the reaction was run at 60°C over night. THF was then

removed by vacuum distillation and 400 mL of methanol was added and stirred for 2 hours. After filtration and drying, it gave 4.5 g of Compound 4.

Compound 4 (240 mg, 0.6 mmol), Pd(PPh₃)₄ (140 mg, 0.12 mmol) and phenyl boronic acid (85.4 mg, 0.7 mmol) were dissolved in 1 mL of 2M Na₂CO₃ aqueous solution and 4 mL of DMF. The resulting solution was heated to 80°C under nitrogen gas overnight. After its completion, the reaction mixture was cooled to room temperature. 100L of water and 100mL of ethyl acetate were added and stirred. The organic layer was retained and washed twice more by water, followed by quickly passing through a silicon gel column to remove solvent. 5mL of dichloromethane and 5mL of trifluoroacetic acid were then added and stirred at room temperature for 4 hrs. After its completion, the reaction mixture was evaporated *in vacuo*. The resulting product was further purified by recrystallization using CH₃OH-THF to arrive at the title compound 6, with the LC-MS [M-H]⁻ m/z 338, and ¹H-NMR (300 MHz, (CD₃)₂SO) δ 13.01 (br s, 1H), 9.55 (br s, 1H), 8.05 (d, 1H, J = 9.0 Hz), 7.85 (d, 1H, J = 9.0 Hz), 7.83 (t, 1H, J = 7.5 Hz), 7.82 (dd, 2H, J = 7.5, 7.5 Hz), 7.54 (dd, 2H, J = 7.5, 1.5 Hz), 7.46 (s, 1H), 4.14 (d, 2H, J = 6.0 Hz).

Example 2

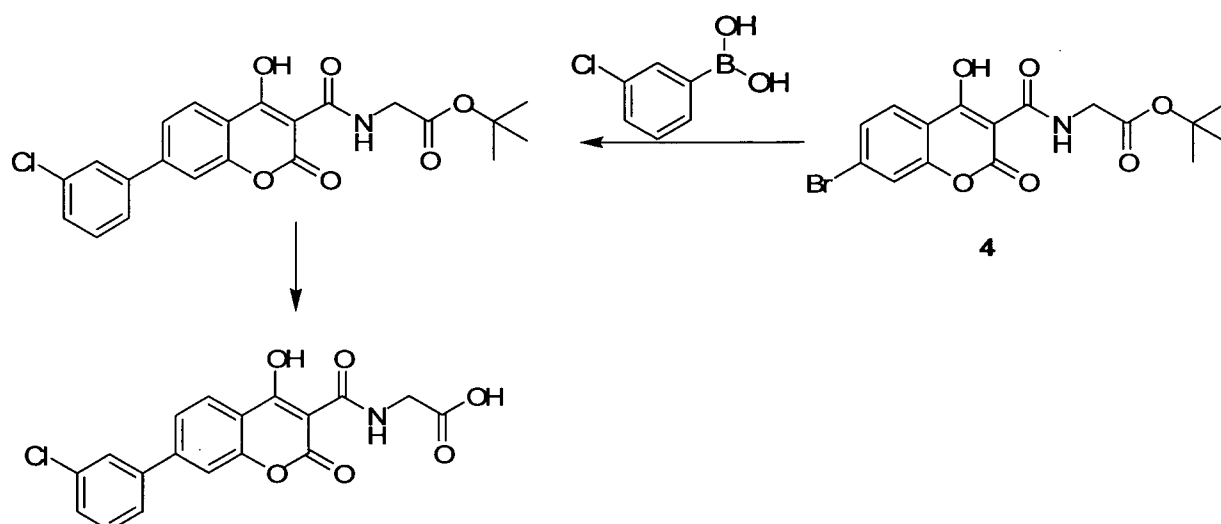
Synthesis of N-[(4-hydroxy-2-oxo-7-(2-chloro-phenyl)-2H-3-chromenyl)carbonyl]glycine



Similar procedure to the previous example was followed to arrive at the title compound, with the LC-MS [M-H]⁻ m/z 373, and ¹H-NMR (300 MHz, (CD₃)₂SO) δ 13.05 (br s, 1H), 9.57 (br s, 1H), 8.08 (d, 1H, J = 8.4 Hz), 7.66-7.32 (m, 6H), 4.15 (d, 2H, J = 6.0 Hz).

Example 3

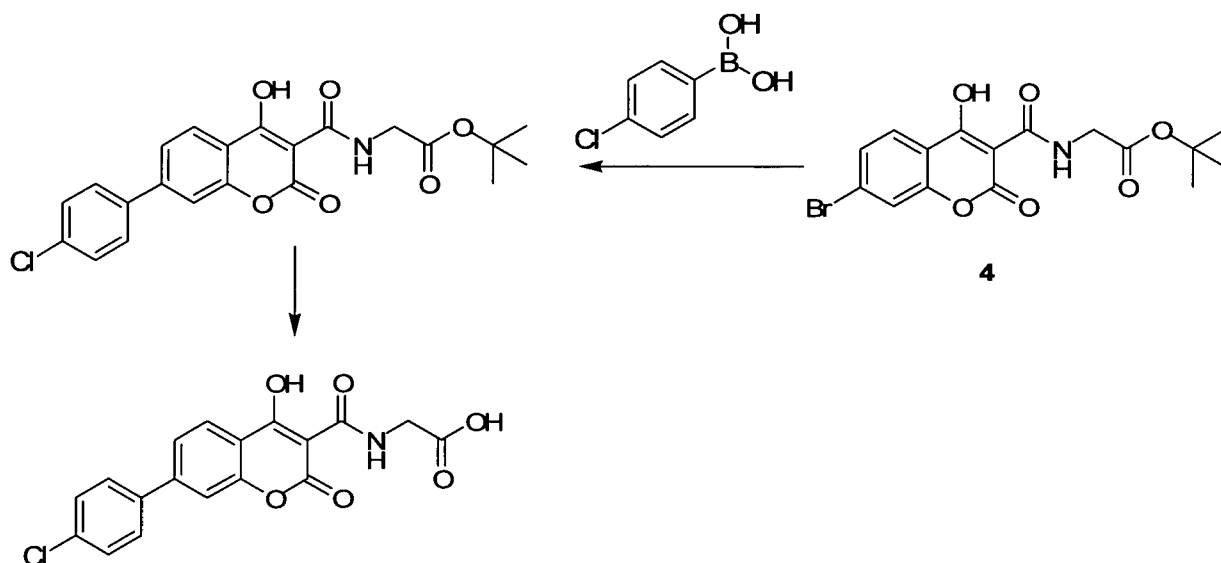
Synthesis of N-[(4-hydroxy-2-oxo-7-(3-chloro-phenyl)-2H-3-chromenyl)carbonyl]glycine



Similar procedure to the previous example was followed to arrive at the title compound, with the LC-MS $[M-H]^-$ m/z 373, and 1H -NMR (300 MHz, $(CD_3)_2SO$) δ 13.05 (br s, 1H), 9.55 (br s, 1H), 8.04 (d, 1H, $J = 8.4$ Hz), 7.92-7.81 (m, 4H), 7.59-7.55 (m, 2H), 4.14 (d, 2H, $J = 6.0$ Hz).

Example 4

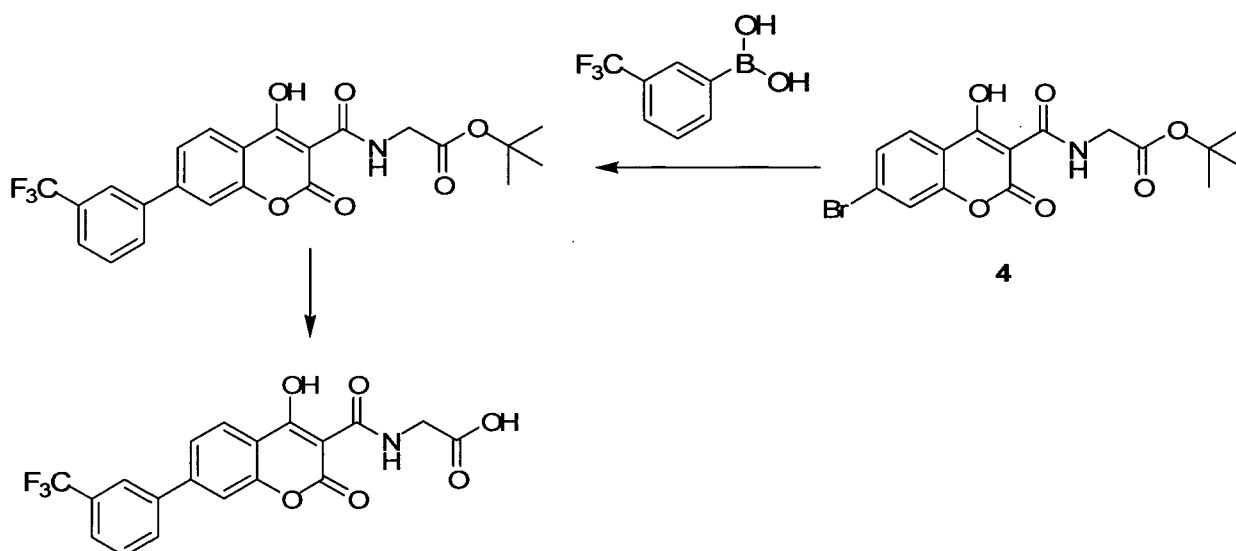
Synthesis of N-[(4-hydroxy-2-oxo-7-(4-chloro-phenyl)-2H-3-chromenyl)carbonyl]glycine



Similar procedure to the previous example was followed to arrive at the title compound, with the LC-MS $[M-H]^-$ m/z 373, and 1H -NMR (300 MHz, $(CD_3)_2SO$) δ 9.55 (br s, 1H), 8.05 (d, 1H, $J = 8.4$ Hz), 7.88 (d, 2H, $J = 8.7$ Hz), 7.84-7.79 (m, 2H), 7.60 (d, 2H, $J = 8.7$ Hz), 4.14 (d, 2H, $J = 6.0$ Hz).

Example 5

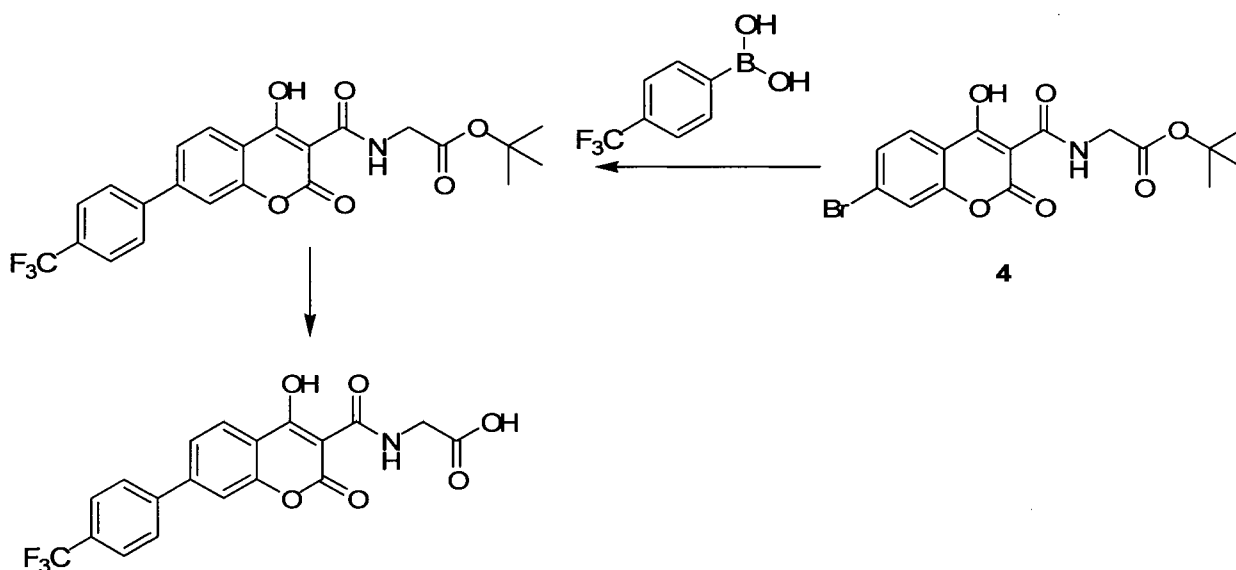
Synthesis of N-[(4-hydroxy-2-oxo-7-(3-trifluoromethyl-phenyl)-2H-3-chromenyl)carbonyl]glycine



Similar procedure to the previous example was followed to arrive at the title compound, with the LC-MS $[M-H]^-$ m/z 406, and 1H -NMR (300 MHz, $(CD_3)_2SO$) δ 9.58 (br s, 1H), 8.18 (s, 1H), 8.16 (s, 1H), 8.09 (d, 1H, $J = 8.4$ Hz), 7.97-7.76 (m, 4H), 4.15 (d, 2H, $J = 6.0$ Hz).

5 Example 6

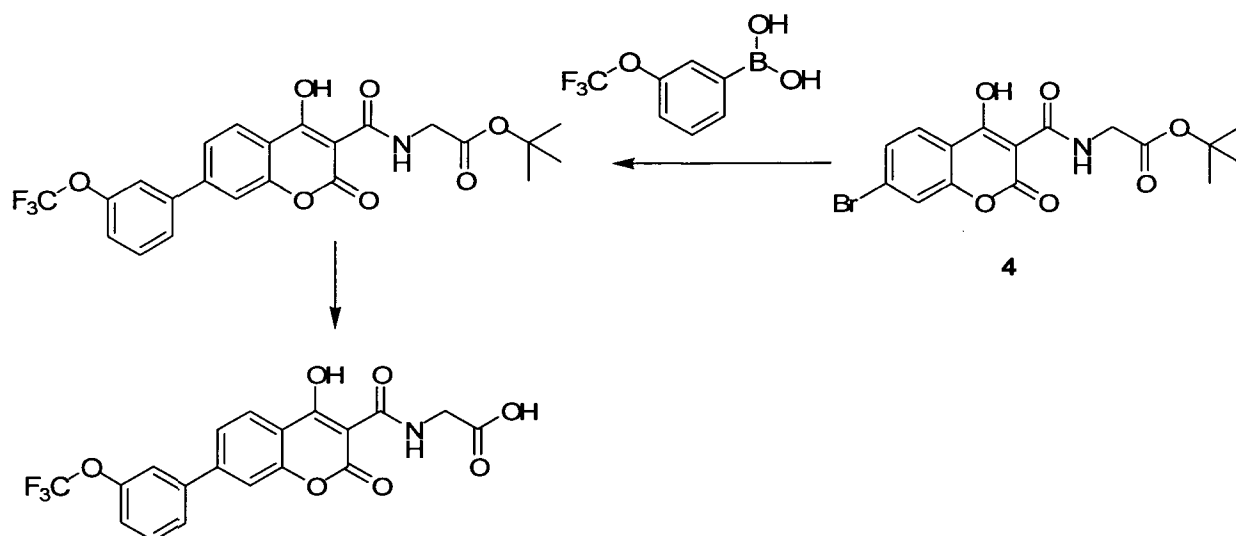
Synthesis of N-[(4-hydroxy-2-oxo-7-(4-trifluoromethyl-phenyl)-2H-3-chromenyl)carbonyl]glycine



Similar procedure to the previous example was followed to arrive at the title compound, with the LC-MS $[M-H]^-$ m/z 406, and 1H -NMR (300 MHz, $(CD_3)_2SO$) δ 9.56 (br s, 1H), 8.11-8.05 (m, 3H), 7.91-7.84 (m, 4H), 4.15 (d, 2H, $J = 6.0$ Hz).

Example 7

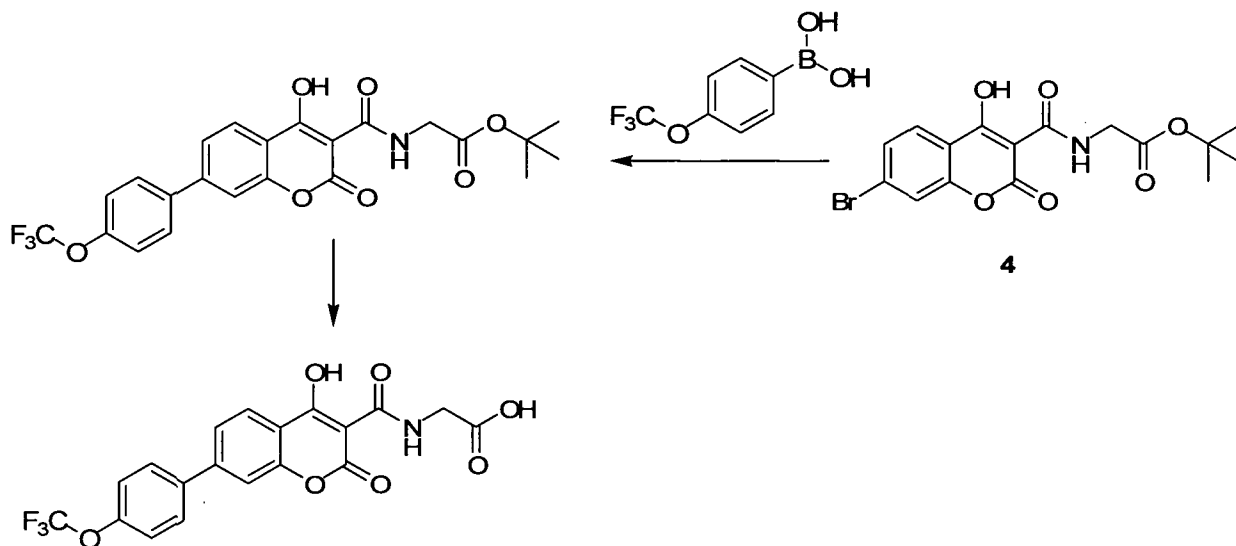
Synthesis of N-[(4-hydroxy-2-oxo-7-(3-trifluoromethoxy-phenyl)-2H-3-chromenyl)carbonyl]glycine



Similar procedure to the previous example was followed to arrive at the title compound, with the LC-MS $[M-H]^-$ m/z 422, and 1H -NMR (300 MHz, $(CD_3)_2SO$) δ 9.57 (br s, 1H), 8.08 (d, 1H, $J = 8.1$ Hz), 7.92-7.85 (m, 4H), 7.69 (t, 1H), 7.50 (d, 1H, $J = 8.4$ Hz), 4.15 (d, 2H, $J = 6.0$ Hz).

5 Example 8

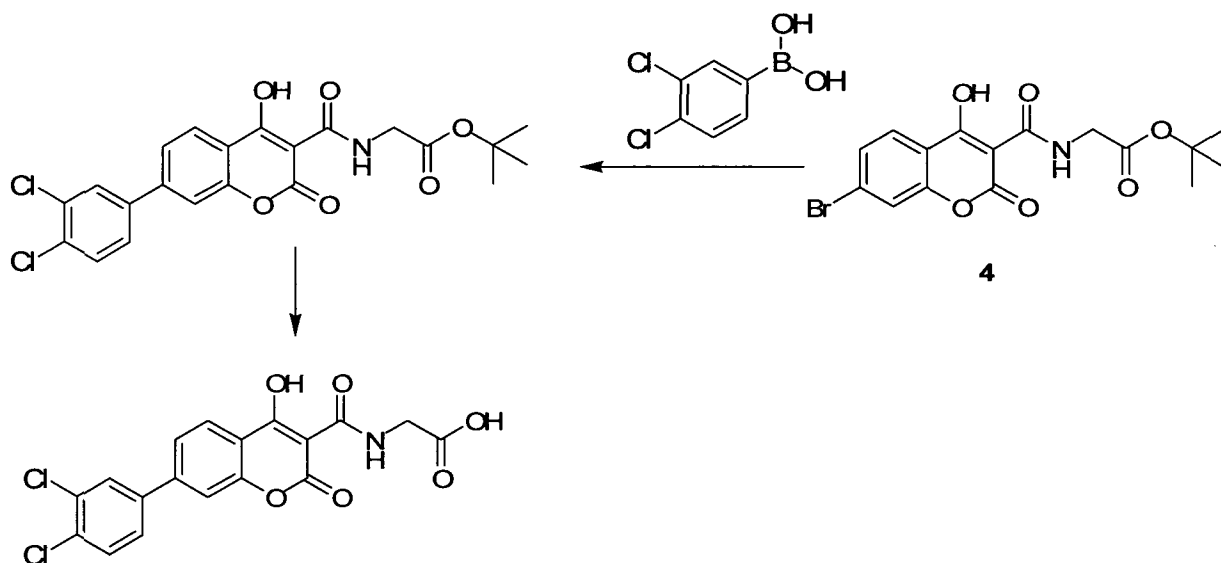
Synthesis of N-[(4-hydroxy-2-oxo-7-(4-trifluoromethoxy-phenyl)-2H-3-chromenyl)carbonyl]glycine



Similar procedure to the previous example was followed to arrive at the title compound, with the LC-MS $[M-H]^-$ m/z 422, and 1H -NMR (300 MHz, $(CD_3)_2SO$) δ 9.56 (br s, 1H), 8.09-7.81 (m, 5H), 7.54 (d, 2H, $J = 8.4$ Hz), 4.15 (d, 2H, $J = 6.0$ Hz).

Example 9

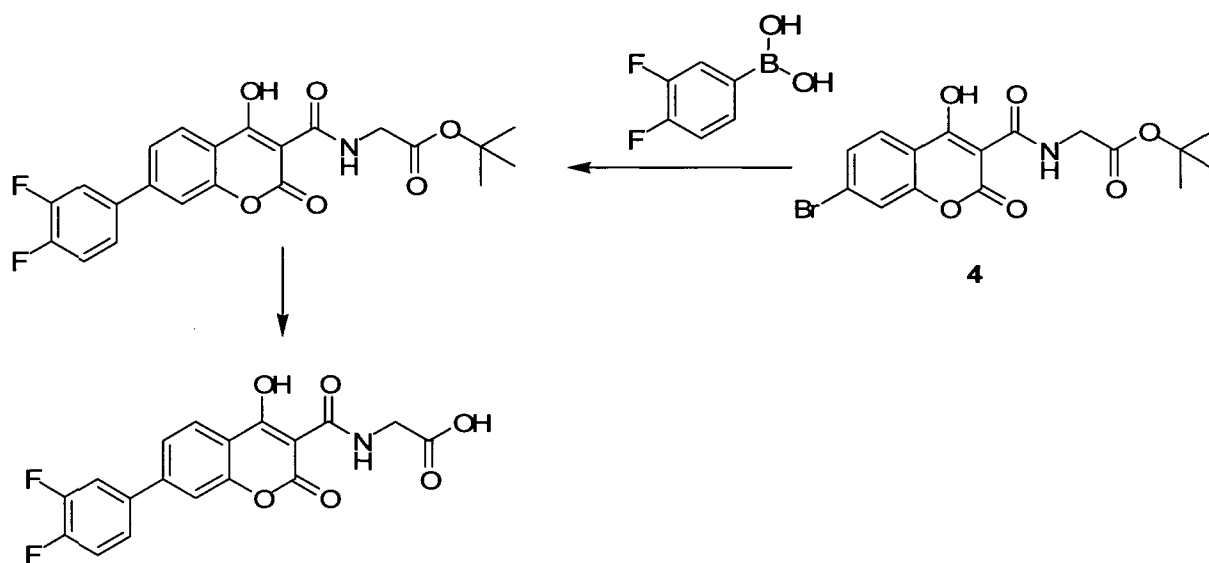
Synthesis of N-[(4-hydroxy-2-oxo-7-(3,4-dichloro-phenyl)-2H-3-chromenyl)carbonyl]glycine



Similar procedure to the previous example was followed to arrive at the title compound, with the LC-MS $[M-H]^-$ m/z 407, and 1H -NMR (300 MHz, $(CD_3)_2SO$) δ 9.57 (br s, 1H), 8.15 (s, 1H), 8.05 (d, 1H, $J = 8.1$ Hz), 7.92-7.78 (m, 4H), 4.15 (d, 2H, $J = 6.0$ Hz).

5 Example 10

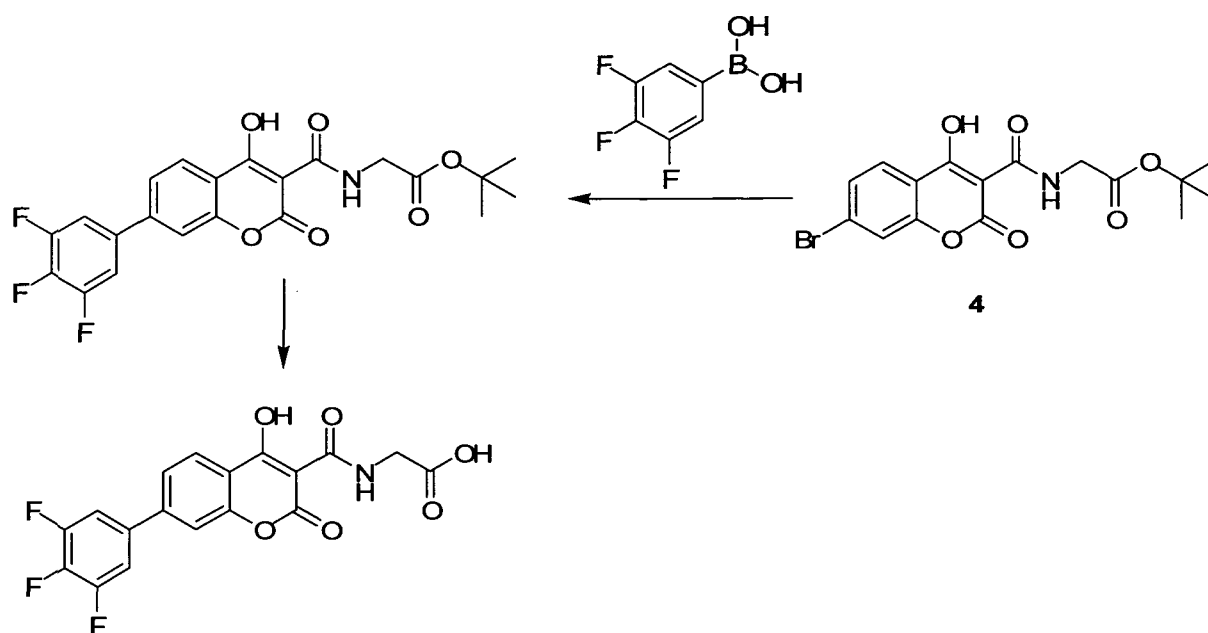
Synthesis of N-[(4-hydroxy-2-oxo-7-(3,4-difluoro-phenyl)-2H-3-chromenyl)carbonyl]glycine



Similar procedure to the previous example was followed to arrive at the title compound, with the LC-MS $[M-H]^-$ m/z 374, and 1H -NMR (300 MHz, $(CD_3)_2SO$) δ 9.55 (br s, 1H), 8.07-7.98 (m, 2H), 7.88-7.82 (m, 2H), 7.66-7.59 (m, 2H), 4.14 (d, 2H, $J = 6.0$ Hz).

Example 11

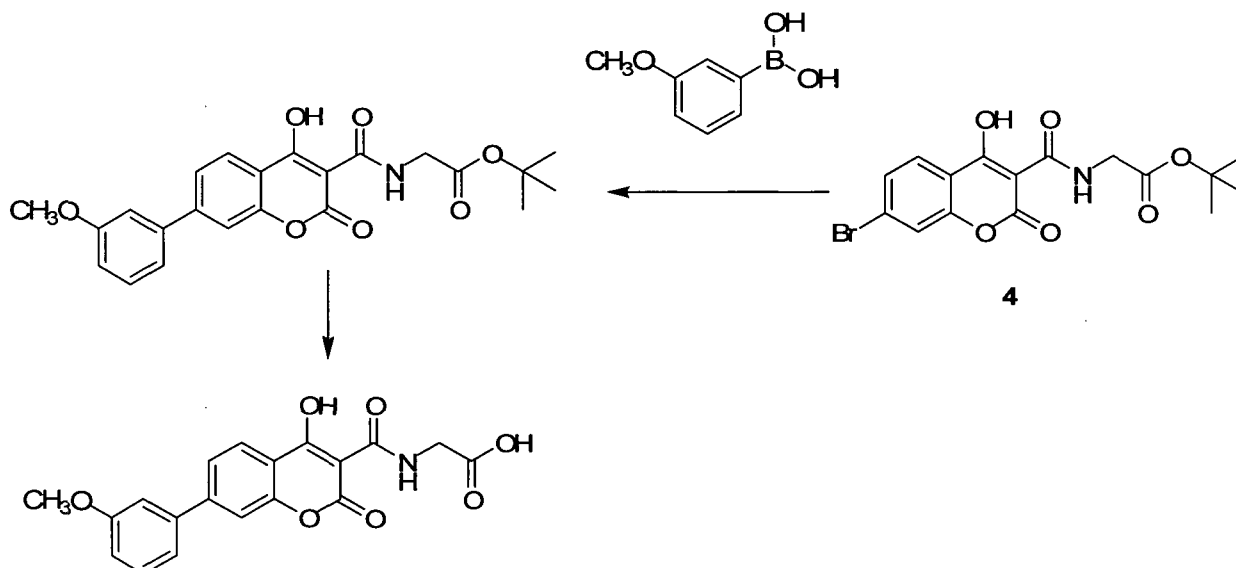
Synthesis of N-[(4-hydroxy-2-oxo-7-(3,4,5-trifluoro-phenyl)-2H-3-chromenyl)carbonyl]glycine



Similar procedure to the previous example was followed to arrive at the title compound, with the LC-MS $[M-H]^-$ m/z 392, and 1H -NMR (300 MHz, $(CD_3)_2SO$) δ 9.55 (br s, 1H), 8.05 (d, 1H, $J = 8.4$ Hz), 7.97-7.85 (m, 4H), 4.14 (d, 2H, $J = 6.0$ Hz).

5 Example 12

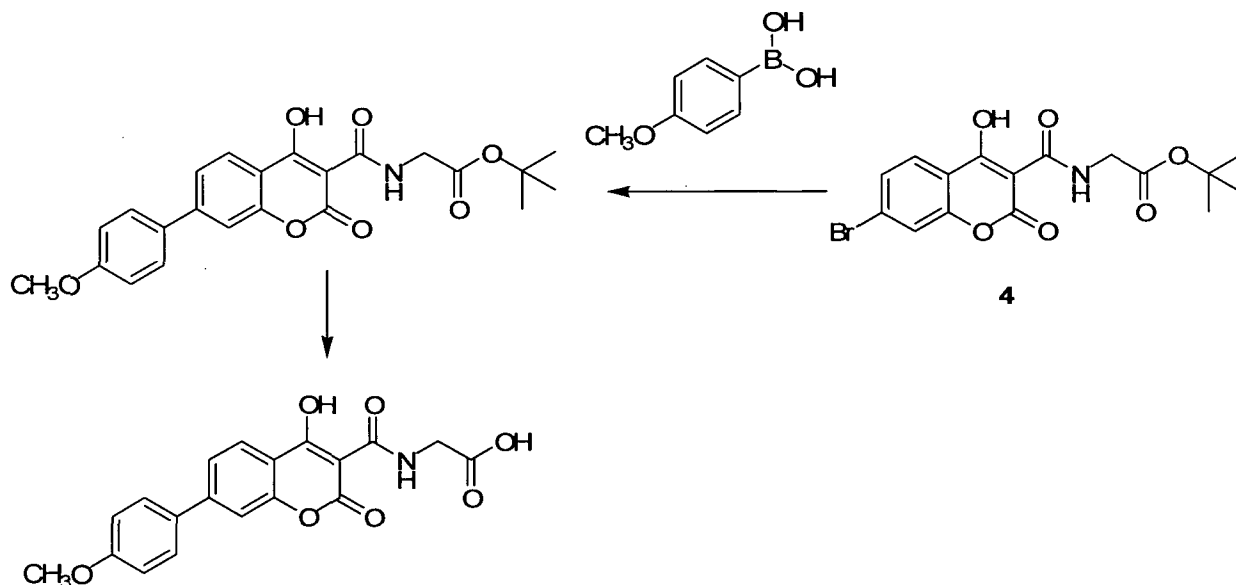
Synthesis of N-[(4-hydroxy-2-oxo-7-(3-methoxy-phenyl)-2H-3-chromenyl)carbonyl]glycine



Similar procedure to the previous example was followed to arrive at the title compound, with the LC-MS $[M-H]^-$ m/z 368, and 1H -NMR (300 MHz, $(CD_3)_2SO$) δ 9.55 (br s, 1H), 8.05 (d, 1H, $J = 8.1$ Hz), 7.86 (s, 1H), 7.83 (d, 1H, $J = 8.1$ Hz), 7.49-7.38 (m, 3H), 7.06 (d, 1H, $J = 8.4$ Hz), 4.15 (d, 2H, $J = 6.0$ Hz).

Example 13

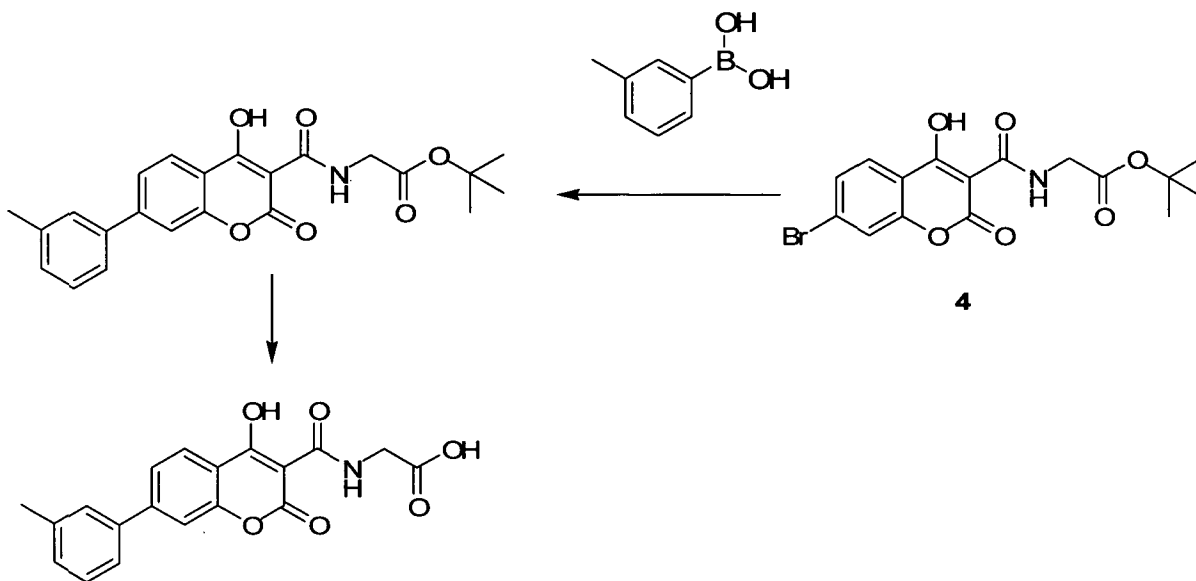
Synthesis of N-[(4-hydroxy-2-oxo-7-(4-methoxy-phenyl)-2H-3-chromenyl)carbonyl]glycine



Similar procedure to the previous example was followed to arrive at the title compound, with the LC-MS $[M-H]^-$ m/z 368, and 1H -NMR (300 MHz, $(CD_3)_2SO$) δ 9.56 (br s, 1H), 8.02 (d, 1H, $J = 8.7$ Hz), 7.85-7.78 (m, 4H), 7.09 (d, 2H, $J = 9.0$ Hz), 4.14 (d, 2H, $J = 6.0$ Hz).

Example 14

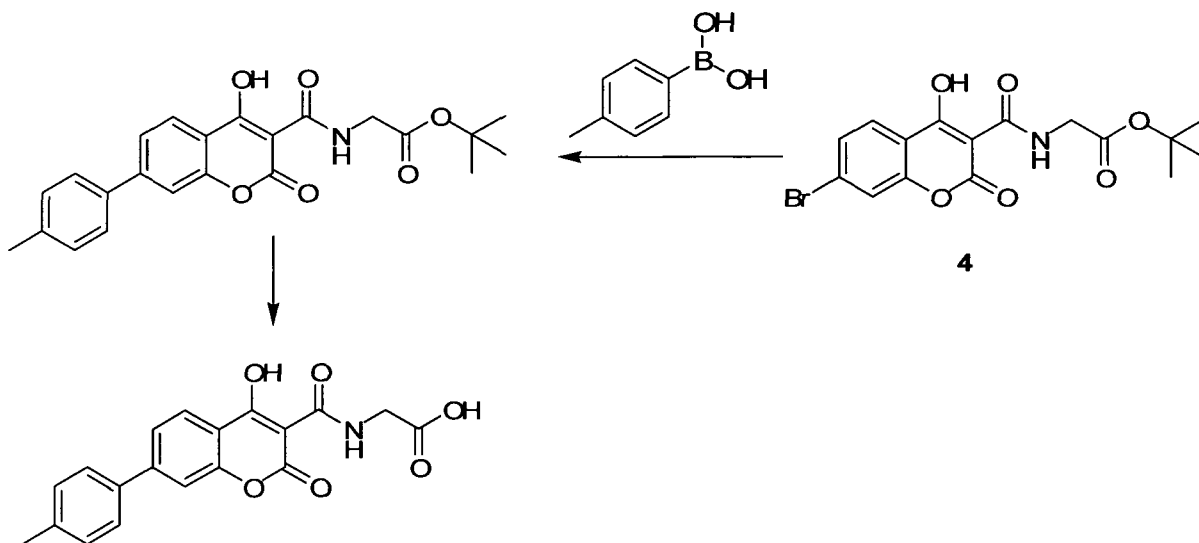
Synthesis of N-[(4-hydroxy-2-oxo-7-(3-methylphenyl)-2H-3-chromenyl)carbonyl]glycine



Similar procedure to the previous example was followed to arrive at the title compound, with the LC-MS $[M-H]^-$ m/z 352, and 1H -NMR (300 MHz, $(CD_3)_2SO$) δ 9.55 (br s, 1H), 8.04 (d, 1H, $J = 8.7$ Hz), 7.80-7.78 (m, 2H), 7.67-7.61 (m, 2H), 7.42 (t, 1H, 7.8 Hz), 7.30 (d, 1H, $J = 7.5$ Hz), 4.14 (d, 2H, $J = 6.0$ Hz), 2.50 (t, 3H, 1.8 Hz).

Example 15

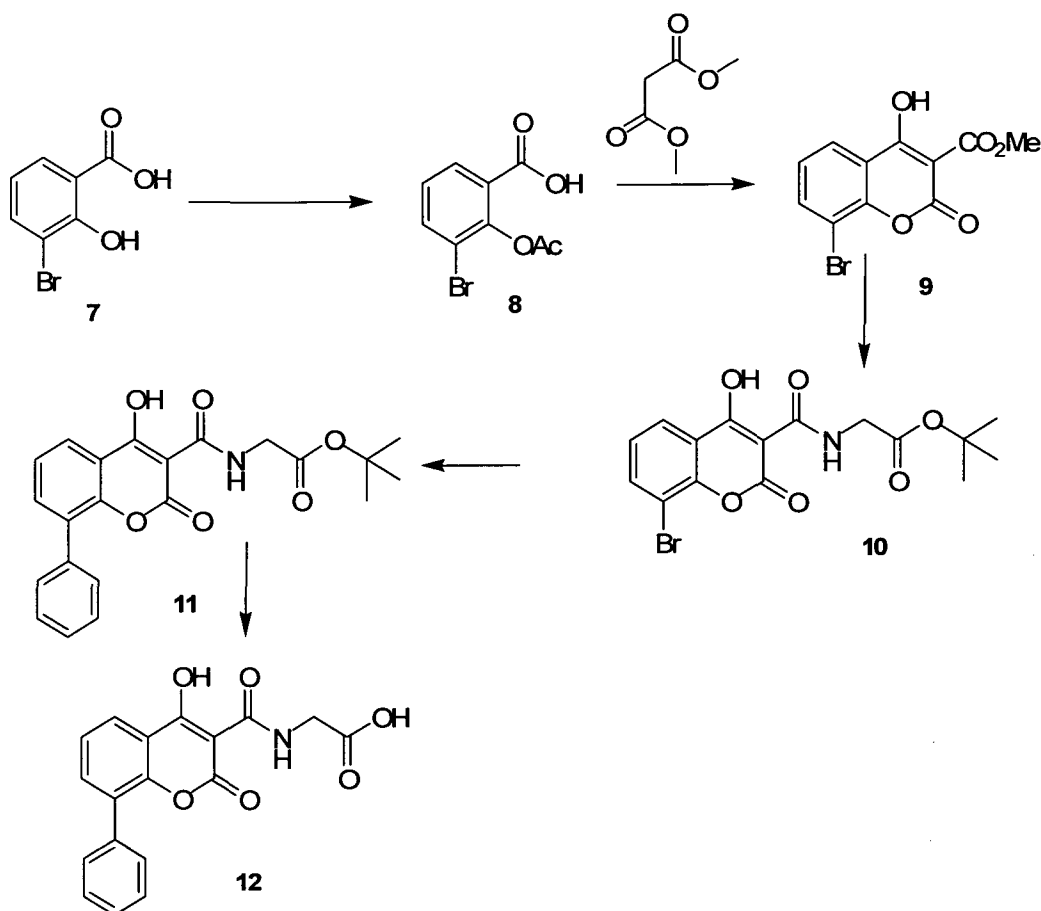
Synthesis of N-[(4-hydroxy-2-oxo-7-(4-methylphenyl)-2H-3-chromenyl)carbonyl]glycine



Similar procedure to the previous example was followed to arrive at the title compound, with the
 5 LC-MS $[M-H]^-$ m/z 352, and 1H -NMR (300 MHz, $(CD_3)_2SO$) δ 9.56 (br s, 1H), 8.02 (d, 1H, $J = 8.7$ Hz), 7.79-7.73 (m, 4H), 7.34 (d, 2H, $J = 8.4$ Hz), 4.13 (d, 2H, $J = 6.0$ Hz), 2.50 (t, 3H, 1.8 Hz).

Example 16

Synthesis of N-[(4-hydroxy-2-oxo-8-phenyl-2H-3-chromenyl)carbonyl]glycine



Similar to the synthetic procedure of Example 1, Reagent 7 (27 g) was dissolved in 130 mL of AC_2O with addition of 1 mL of H_3PO_4 , and was heated at 50°C . After its completion, the reaction mixture was cooled to room temperature, and 500 mL of water was then added in and stirred at 50°C until hydrolyzation is complete. The reaction mixture was then cooled to 0°C and filtered. The resulting solid, Compound 8, was further dried (24.4 g).

Compound 8 (13 g) and 1-Hydroxybenzotriazole anhydrous (HOBt, 6.8 g) was dissolved in 200 mL of THF. Dicyclohexylcarbodiimide (DCC, 10.4 g) was gradually added in at 0°C and stirred over night at a temperature below 10°C . The reaction mixture was filtered and the Filtrate B was collected. 6.6 g (50 mmol) of dimethyl malonate was dissolved in 400 mL of THF first, then 3.8 g sodium hydride (70% dispersion) was added in. Filtrate B was added while stirred and left to react for 2 hours at room temperature. After THF removed by vacuum distillation, 200 mL of methanol and 200 mL of 10% HCl were added and stirred over night. After filtration, the resulting solid was washed with 200 mL of methanol and dried, yielding 6 g of Compound 9.

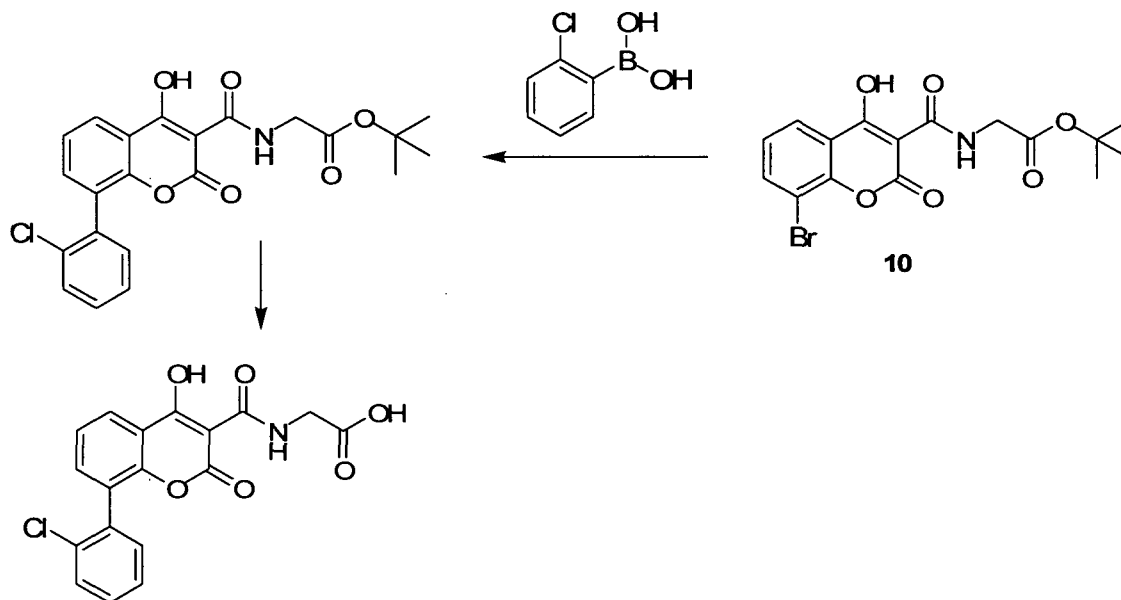
Glycine t-butyl ester HCl salt (13.5 g) and sodium methoxide (4.4 g) were suspended in 200 mL of methanol. After stirred to a homogeneous suspension, it was distilled to remove all methanol. 200 mL of

THF and Compound 9 (6.0 g) were added in and the reaction was run at 60°C over night. THF was then removed by vacuum distillation and 400 mL of methanol was added and stirred for 2 hours. After filtration and drying, it gave 3.5 g of Compound 10.

Compound 10 (240 mg, 0.6 mmol), Pd(PPh₃)₄ (140 mg, 0.12 mmol) and phenyl boronic acid (85.4 mg, 0.7 mmol) were dissolved in 1 mL of 2M Na₂CO₃ aqueous solution and 4 mL of DMF. The resulting solution was heated to 80°C under nitrogen gas overnight. After its completion, the reaction mixture was cooled to room temperature. 100mL of water and 100 mL of ethyl acetate were added and stirred. The organic layer was retained and washed twice more by water, followed by quickly passing through a silicon gel column to remove solvent. 5 mL of dichloromethane and 5 mL of trifluoroacetic acid were then added and stirred at room temperature for 4 hrs. After its completion, the reaction mixture was evaporated *in vacuo*. The resulting product was further purified by recrystallization using CH₃OH-THF to arrive at the title compound 12, with the LC-MS [M-H]⁻ m/z 338, and ¹H-NMR (300 MHz, (CD₃)₂SO) δ 9.51 (br s, 1H), 8.03 (dd, 1H, J = 8.1, 1.5 Hz), 7.86 (dd, 2H, J = 7.5, 1.5 Hz), 7.65-7.45 (m, 6H), 4.15 (d, 2H, J = 6.0 Hz).

Example 17

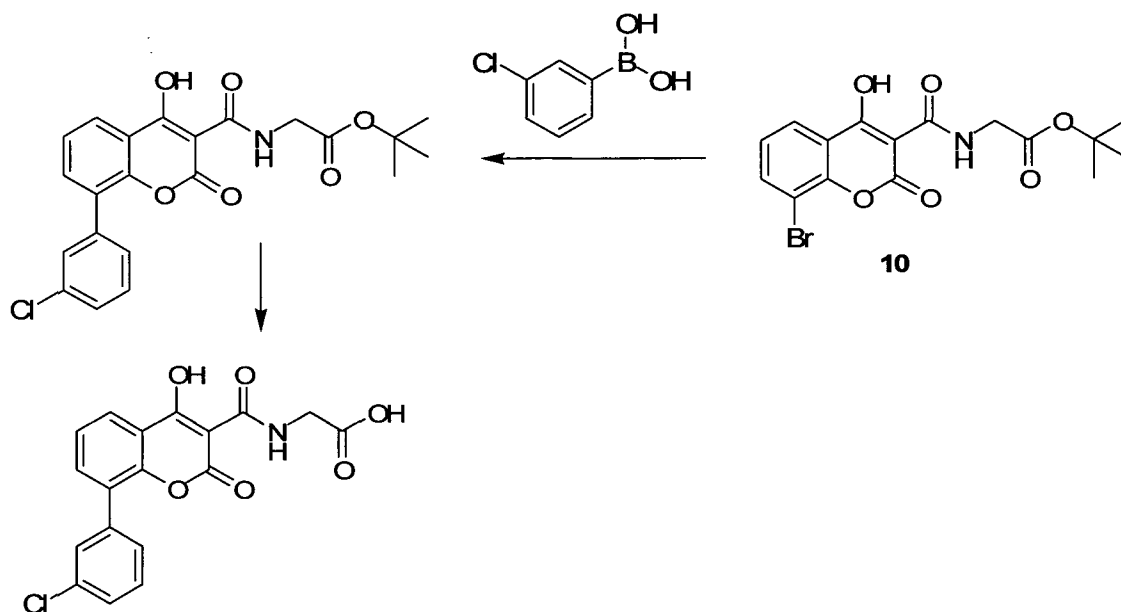
15 Synthesis of N-[(4-hydroxy-2-oxo-8-(2-chlorophenyl)-2H-3-chromenyl)carbonyl]glycine



Similar procedure to the previous example was followed to arrive at the title compound, with the LC-MS [M-H]⁻ m/z 373, and ¹H-NMR (300 MHz, (CD₃)₂SO) δ 9.47 (br s, 1H), 8.10 (m, 1H), 7.77 (m, 1H), 7.67-7.50 (m, 5H), 4.14 (d, 2H, J = 6.0 Hz).

20 Example 18

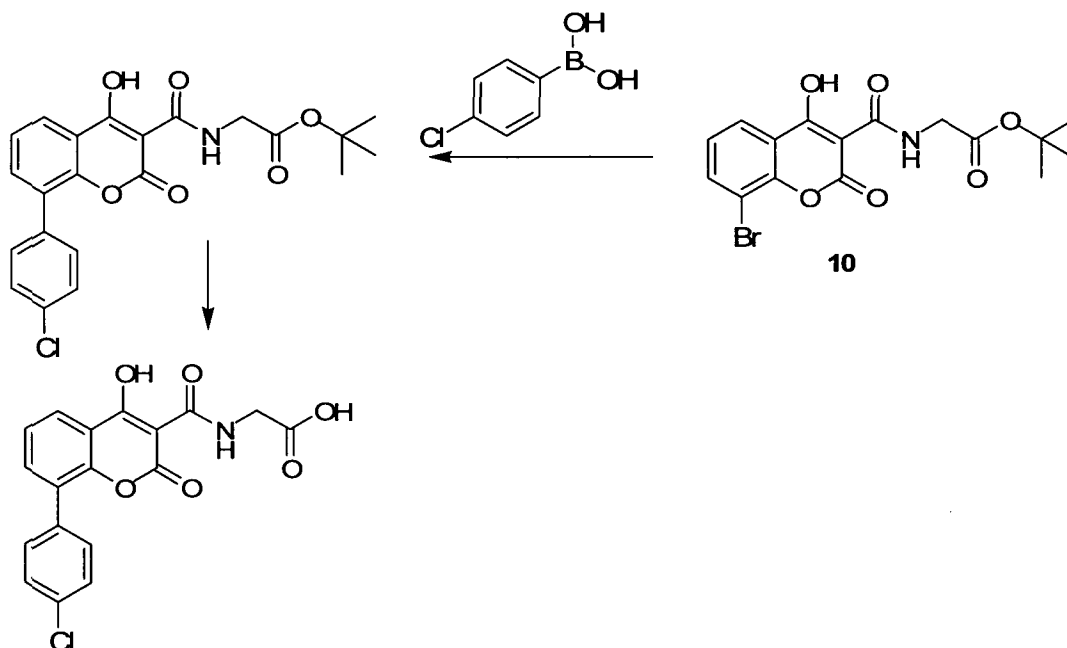
Synthesis of N-[(4-hydroxy-2-oxo-8-(3-chlorophenyl)-2H-3-chromenyl)carbonyl]glycine



Similar procedure to the previous example was followed to arrive at the title compound, with the LC-MS $[M-H]^-$ m/z 373, and 1H -NMR (300 MHz, $(CD_3)_2SO$) δ 9.51 (br s, 1H), 8.06 (dd, 1H, $J = 8.1, 1.5$ Hz), 7.89 (m, 1H), 7.73 (s, 1H), 7.61-7.54 (m, 4H), 4.15 (d, 2H, $J = 6.0$ Hz).

5 Example 19

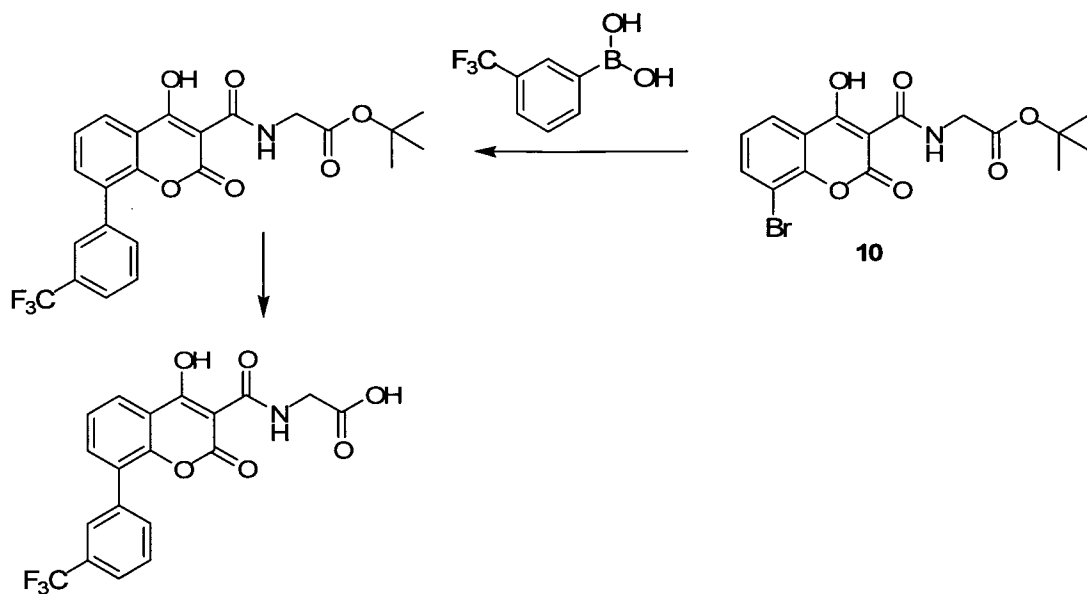
Synthesis of N-[(4-hydroxy-2-oxo-8-(4-chlorophenyl)-2H-3-chromenyl)carbonyl]glycine



Similar procedure to the previous example was followed to arrive at the title compound, with the LC-MS $[M-H]^-$ m/z 373, and 1H -NMR (300 MHz, $(CD_3)_2SO$) δ 9.49 (br s, 1H), 8.04 (dd, 1H, $J = 8.1, 1.5$ Hz), 7.86 (m, 1H), 7.69-7.53 (m, 5H), 4.15 (d, 2H, $J = 6.0$ Hz).

Example 20

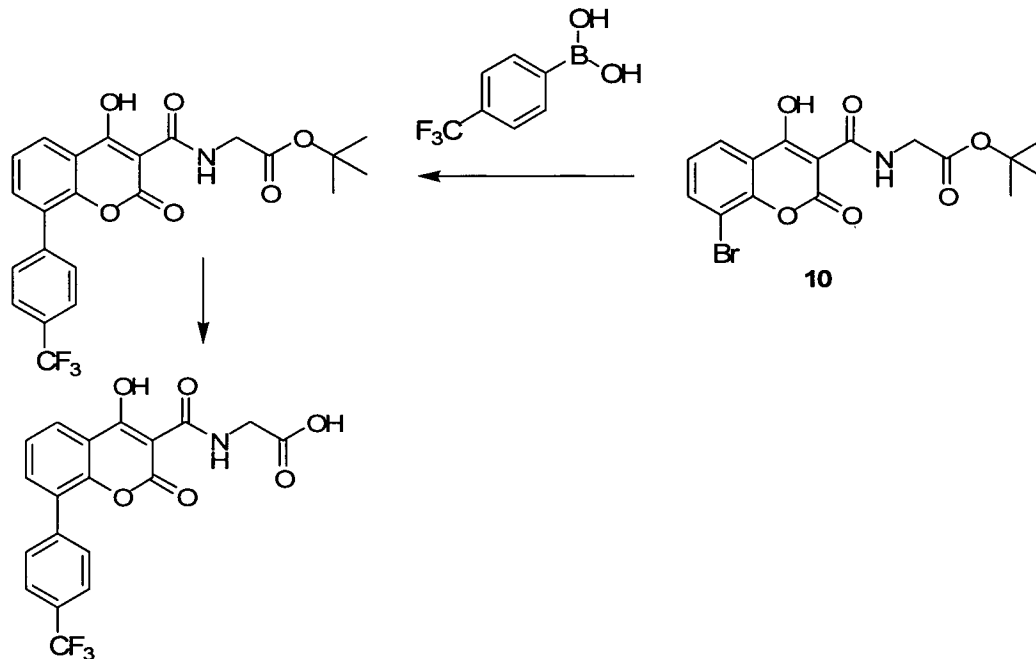
Synthesis of N-[(4-hydroxy-2-oxo-8-(3-trifluoromethylphenyl)-2H-3-chromenyl)carbonyl]glycine



Similar procedure to the previous example was followed to arrive at the title compound, with the LC-MS $[M-H]^-$ m/z 406, and 1H -NMR (300 MHz, $(CD_3)_2SO$) δ 9.52 (br s, 1H), 8.09-7.76 (m, 6H), 7.58 (t, 1H, 7.8 Hz), 4.15 (d, 2H, $J = 6.0$ Hz).

5 Example 21

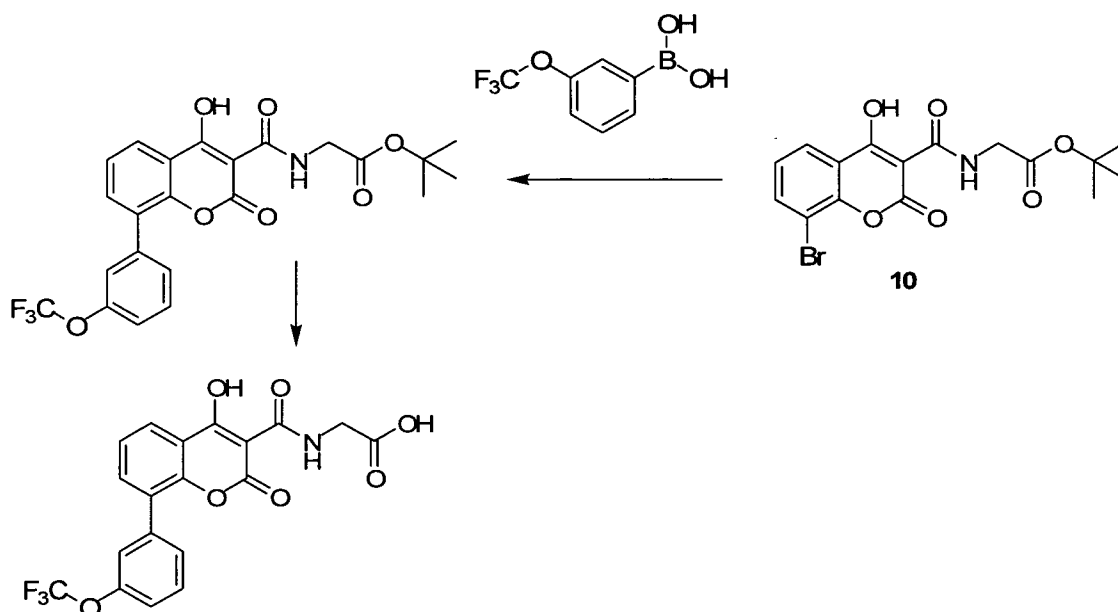
Synthesis of N-[(4-hydroxy-2-oxo-8-(4-trifluoromethylphenyl)-2H-3-chromenyl)carbonyl]glycine



Similar procedure to the previous example was followed to arrive at the title compound, with the LC-MS $[M-H]^-$ m/z 406, and 1H -NMR (300 MHz, $(CD_3)_2SO$) δ 9.50 (br s, 1H), 8.09 (dd, 1H, 7.8, 1.5 Hz), 7.93-7.86 (m, 5H), 7.59 (t, 1H, 7.8 Hz), 4.15 (d, 2H, $J = 6.0$ Hz).

Example 22

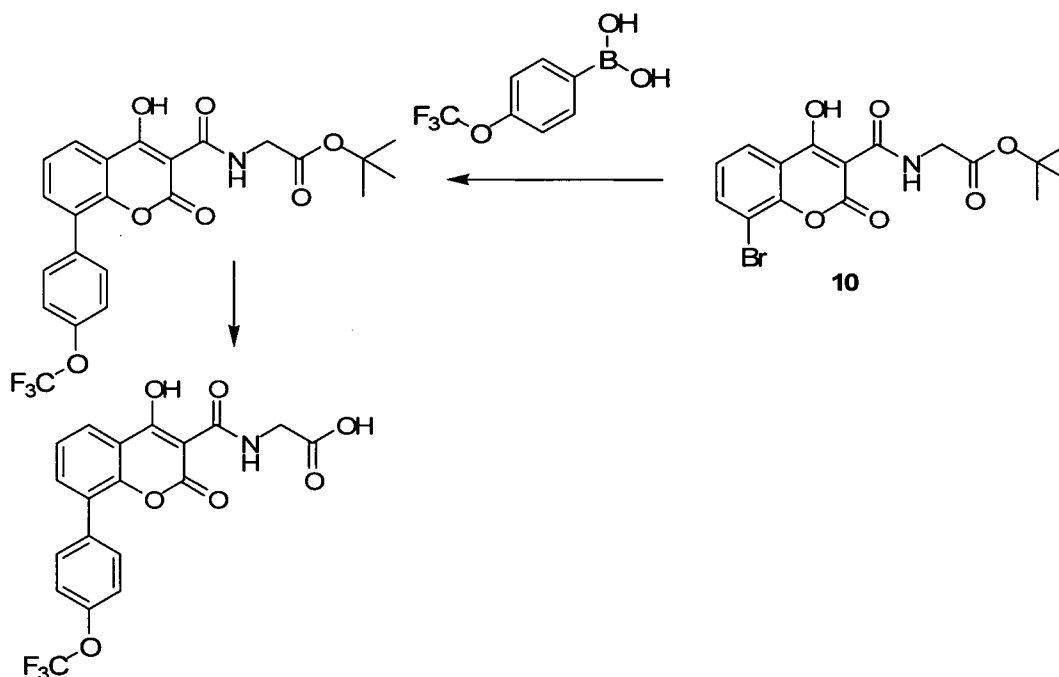
Synthesis of N-[(4-hydroxy-2-oxo-8-(3-trifluoromethoxyphenyl)-2H-3-chromenyl)carbonyl]glycine



Similar procedure to the previous example was followed to arrive at the title compound, with the LC-MS $[M-H]^-$ m/z 422, and 1H -NMR (300 MHz, $(CD_3)_2SO$) δ 9.52 (br s, 1H), 8.07 (dd, 1H, 7.8, 1.5 Hz), 7.91 (dd, 1H, 7.5, 1.5 Hz), 7.70-7.68 (m, 3H), 7.57 (t, 1H, 7.8 Hz), 7.49 (br s, 1H), 4.15 (d, 2H, $J = 6.0$ Hz).

Example 23

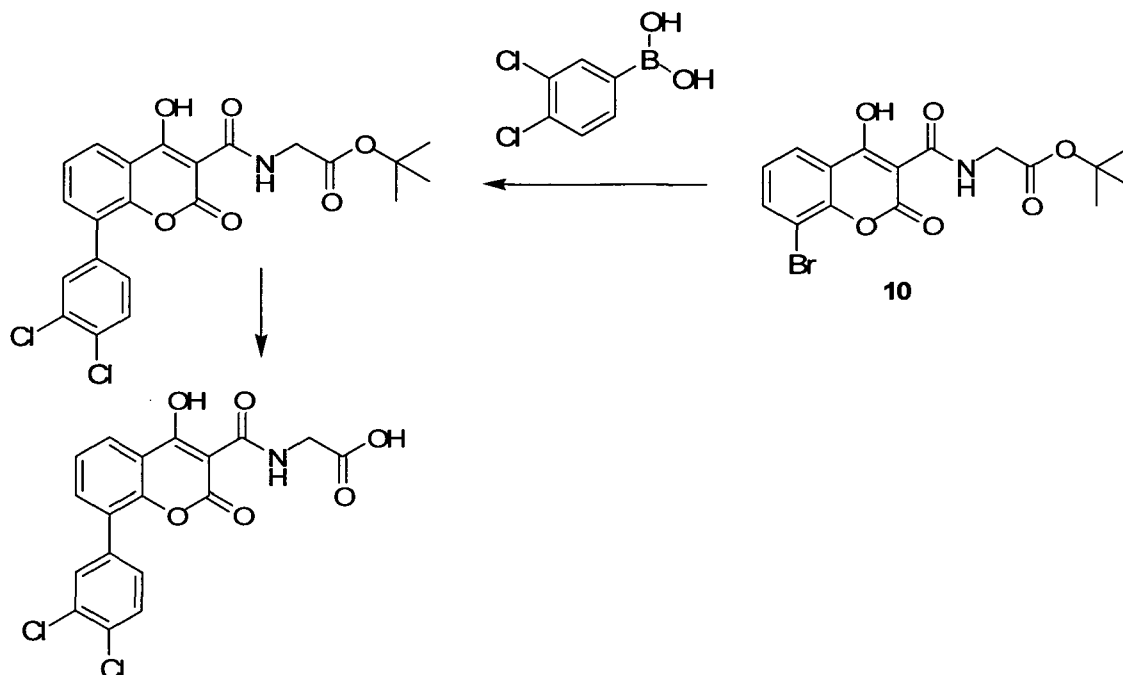
Synthesis of N-[(4-hydroxy-2-oxo-8-(4-trifluoromethoxyphenyl)-2H-3-chromenyl)carbonyl]glycine



Similar procedure to the previous example was followed to arrive at the title compound, with the LC-MS $[M-H]^-$ m/z 422, and 1H -NMR (300 MHz, $(CD_3)_2SO$) δ 9.50 (br s, 1H), 8.06 (dd, 1H, 8.1, 1.5 Hz), 7.89 (m, 1H), 7.78 (d, 2H, $J = 8.7$ Hz), 7.60-7.54 (m, 3H), 4.15 (d, 2H, $J = 6.0$ Hz).

Example 24

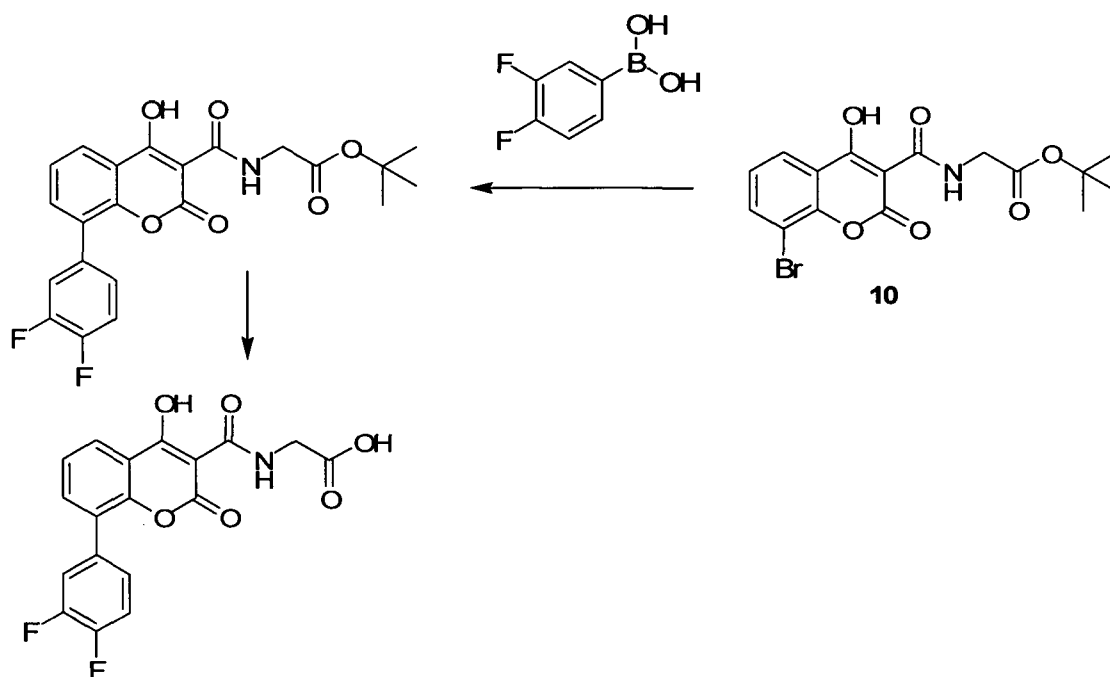
Synthesis of N-[(4-hydroxy-2-oxo-8-(3,4-dichlorophenyl)-2H-3-chromenyl)carbonyl]glycine



Similar procedure to the previous example was followed to arrive at the title compound, with the
 5 LC-MS $[M-H]^-$ m/z 407, and 1H -NMR (300 MHz, $(CD_3)_2SO$) δ 9.51 (br s, 1H), 8.06 (dd, 1H, 8.1, 1.5 Hz), 7.95-7.89 (m, 2H), 7.82 (d, 1H, $J = 8.4$ Hz), 7.65 (dd, 1H, $J = 8.4, 1.5$ Hz), 7.56 (t, 1H, $J = 7.5$ Hz), 4.15 (d, 2H, $J = 6.0$ Hz).

Example 25

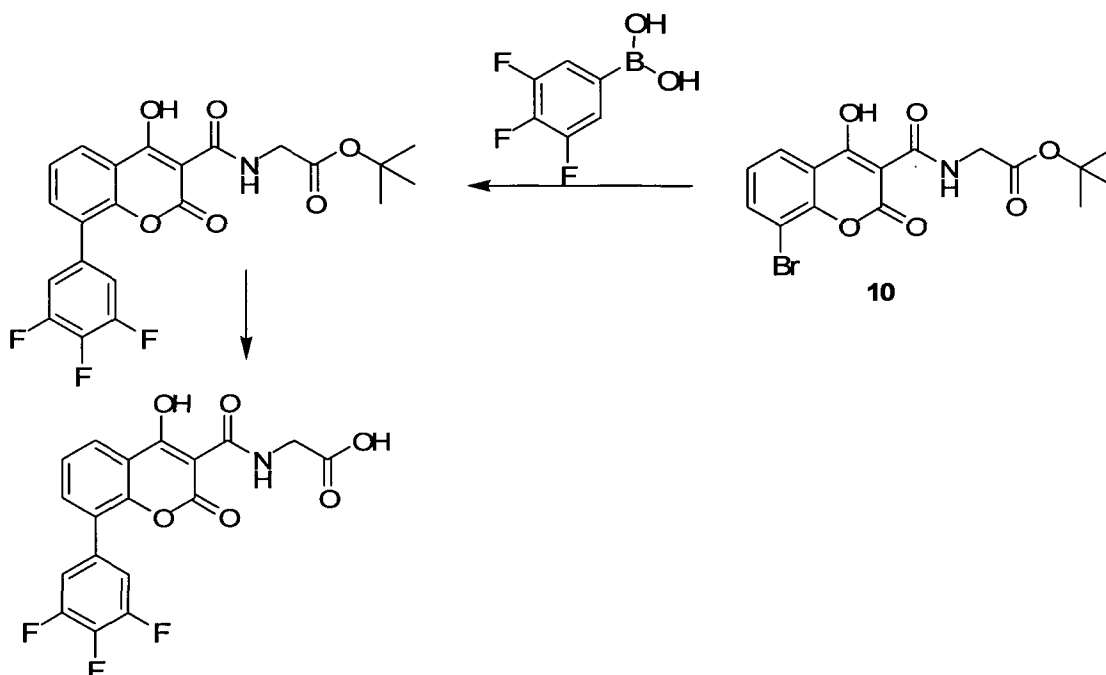
Synthesis of N-[(4-hydroxy-2-oxo-8-(3,4-difluorophenyl)-2H-3-chromenyl)carbonyl]glycine



Similar procedure to the previous example was followed to arrive at the title compound, with the LC-MS $[M-H]^-$ m/z 374, and 1H -NMR (300 MHz, $(CD_3)_2SO$) δ 9.51 (br s, 1H), 7.88 (dd, 1H, $J = 7.5, 1.2$ Hz), 7.82-7.75 (m, 2H), 7.68 -7.53 (m, 3H), 4.15 (d, 2H, $J = 6.0$ Hz).

Example 26

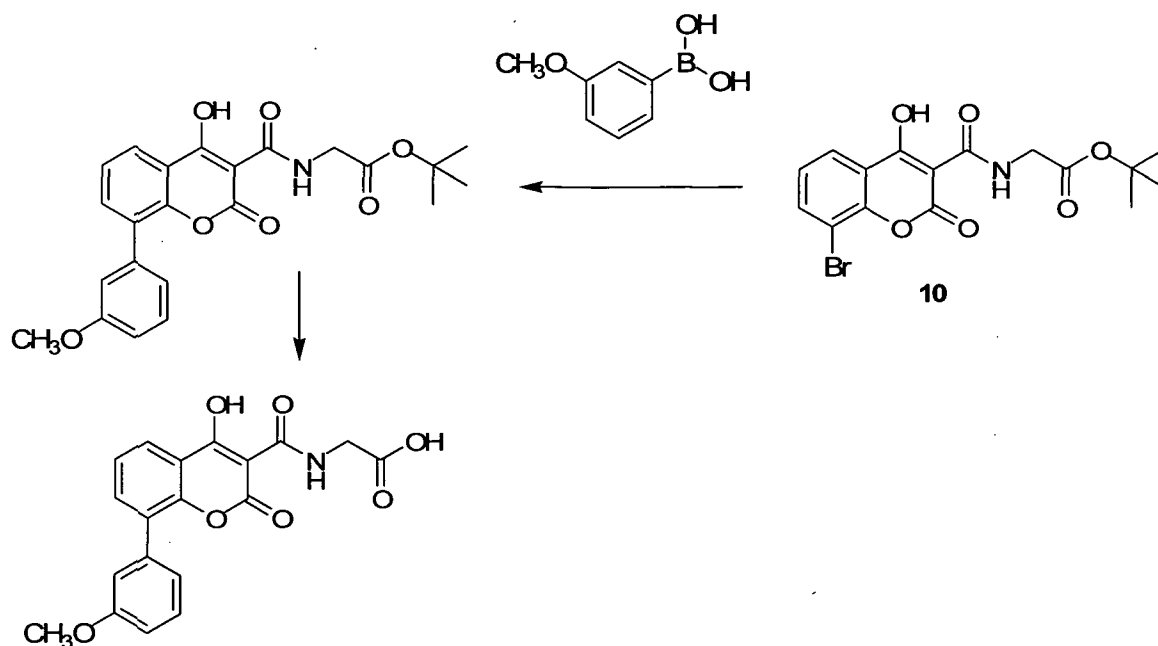
5 Synthesis of N-[(4-hydroxy-2-oxo-8-(3,4,5-trifluorophenyl)-2H-3-chromenyl)carbonyl]glycine



Similar procedure to the previous example was followed to arrive at the title compound, with the LC-MS $[M-H]^-$ m/z 392, and 1H -NMR (300 MHz, $(CD_3)_2SO$) δ 9.51 (br s, 1H), 8.07 (dd, 1H, $J = 7.8, 1.5$ Hz), 8.01-7.97 (m, 1H), 7.92 -7.89 (m, 1H), 7.71-7.66 (m, 1H), 7.63 (t, 1H, $J = 7.8$ Hz), 4.15 (d, 2H, $J = 6.0$ Hz).

Example 27

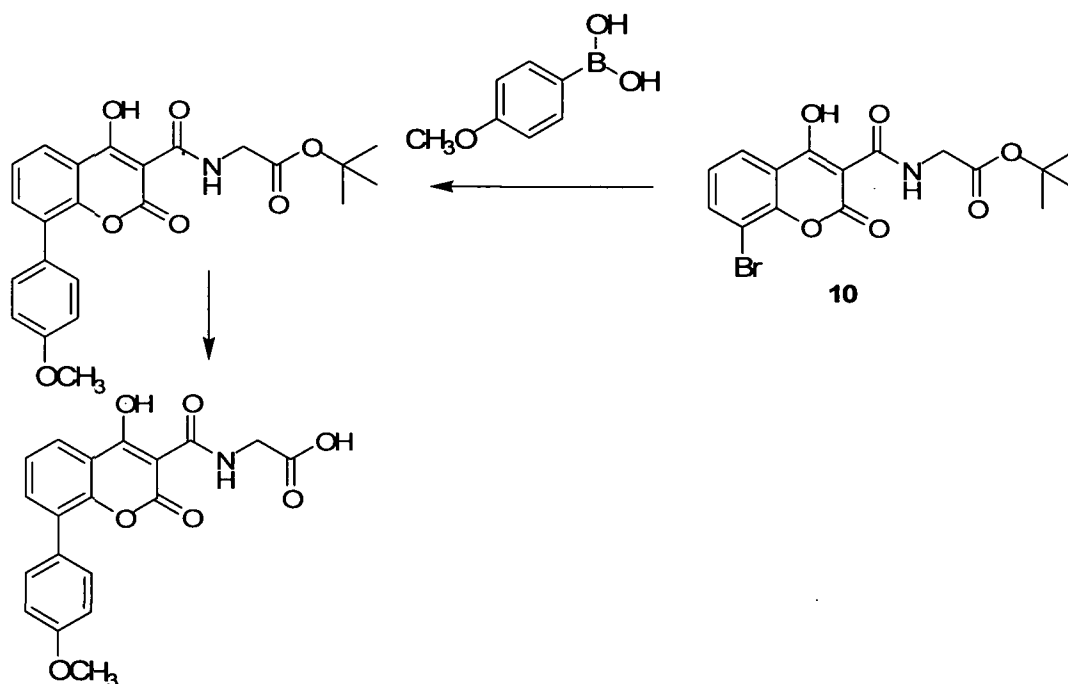
10 Synthesis of N-[(4-hydroxy-2-oxo-8-(3-methoxyphenyl)-2H-3-chromenyl)carbonyl]glycine



Similar procedure to the previous example was followed to arrive at the title compound, with the LC-MS $[M-H]^-$ m/z 368, and 1H -NMR (300 MHz, $(CD_3)_2SO$) δ 9.52 (br s, 1H), 8.07 (dd, 1H, $J = 7.8, 1.5$ Hz), 7.87 (dd, 1H, $J = 7.8, 1.5$ Hz), 7.55 (t, 1H, $J = 7.8$ Hz), 7.45 (t, 1H, $J = 7.8$ Hz), 7.20-7.18 (m, 2H), 7.06-7.03 (m, 1H), 4.15 (d, 2H, $J = 6.0$ Hz).

Example 28

Synthesis of N-[(4-hydroxy-2-oxo-8-(4-methoxyphenyl)-2H-3-chromenyl)carbonyl]glycine

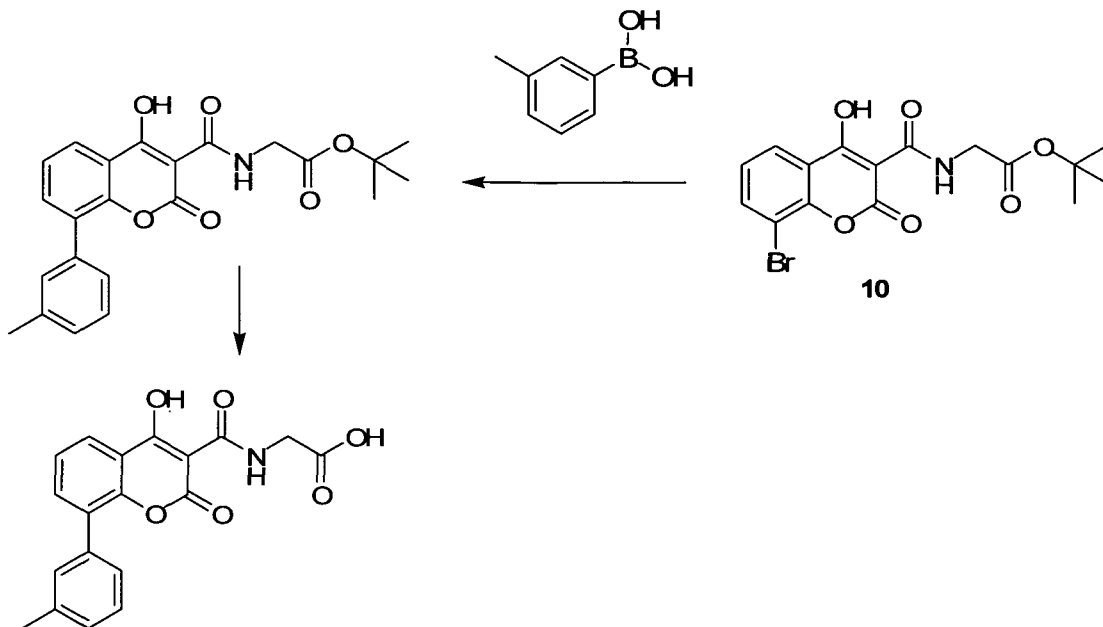


Similar procedure to the previous example was followed to arrive at the title compound, with the LC-MS $[M-H]^-$ m/z 368, and 1H -NMR (300 MHz, $(CD_3)_2SO$) δ 9.51 (br s, 1H), 7.98 (d, 1H, $J = 7.8$ Hz),

7.83-7.80 (m, 1H), 7.59-7.50 (m, 3H), 7.09 (d, 2H, $J = 8.7$ Hz), 4.15 (d, 2H, $J = 6.0$ Hz).

Example 29

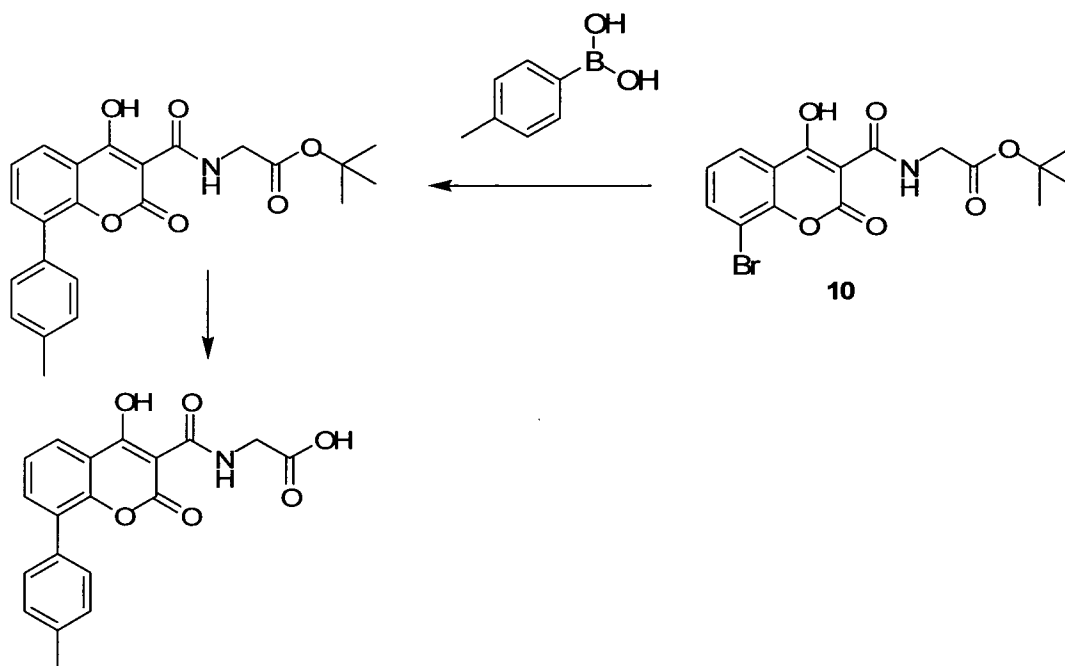
Synthesis of N-[(4-hydroxy-2-oxo-8-(3-methylphenyl)-2H-3-chromenyl)carbonyl]glycine



- 5 Similar procedure to the previous example was followed to arrive at the title compound, with the LC-MS $[M-H]^-$ m/z 352, and 1H -NMR (300 MHz, $(CD_3)_2SO$) δ 9.52 (br s, 1H), 8.03 (dd, 1H, $J = 7.8, 1.5$ Hz), 7.84-7.82 (m, 1H), 7.57-7.52 (m, 1H), 7.43-7.42 (m, 3H), 7.29 (br s, 1H), 4.14 (d, 2H, $J = 6.0$ Hz), 2.51 (t, 3H, $J = 1.8$ Hz).

Example 30

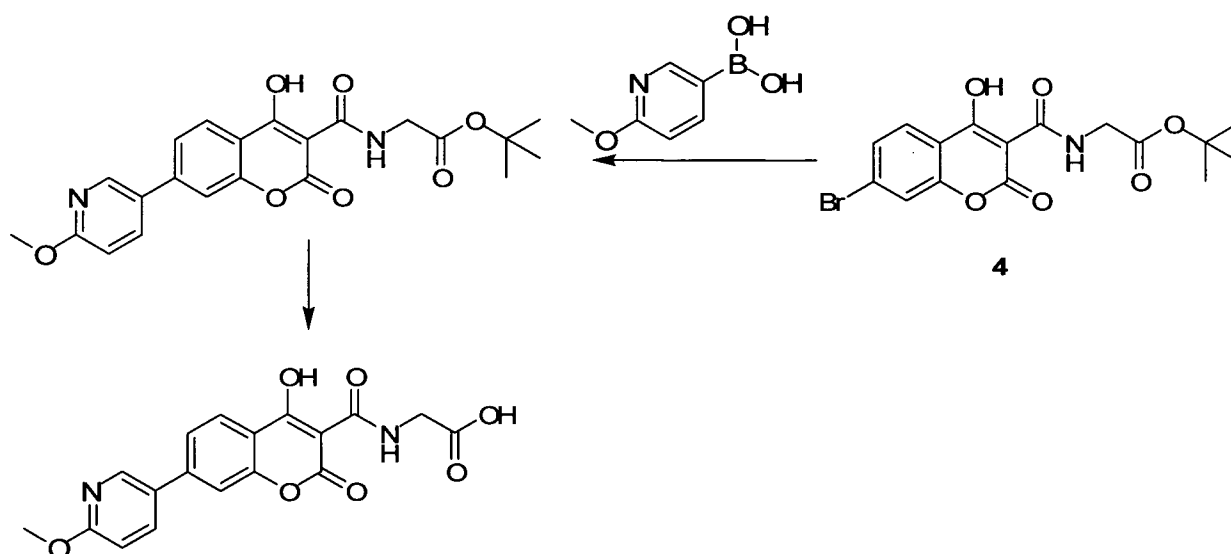
- 10 Synthesis of N-[(4-hydroxy-2-oxo-8-(4-methylphenyl)-2H-3-chromenyl)carbonyl]glycine



Similar procedure to the previous example was followed to arrive at the title compound, with the LC-MS $[M-H]^-$ m/z 352, and 1H -NMR (300 MHz, $(CD_3)_2SO$) δ 9.53 (br s, 1H), 8.02-7.99 (m, 1H), 7.84-7.82 (m, 1H), 7.57-7.51 (m, 3H), 7.36-7.33 (m, 2H), 4.14 (d, 2H, $J = 6.0$ Hz), 2.51 (t, 3H, $J = 1.8$ Hz).

Example 31

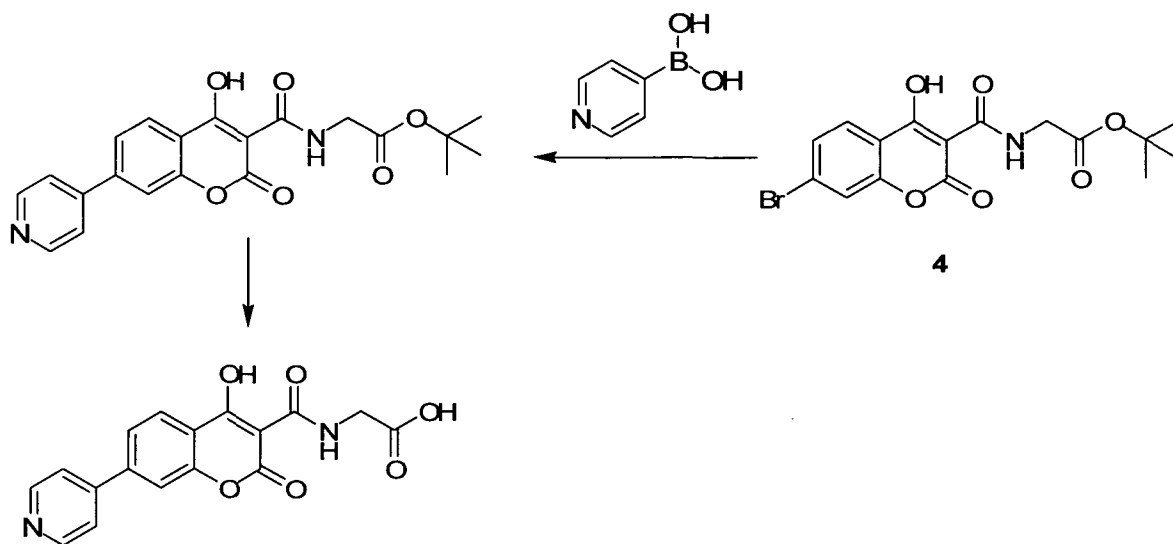
Synthesis of N-[(4-hydroxy-2-oxo-7-(6-methoxypyridin-3-yl)-2H-3-chromenyl)carbonyl]glycine



Similar procedure to Example 1 was followed to arrive at the title compound, with the LC-MS $[M-H]^-$ m/z 369, and 1H -NMR (300 MHz, $(CD_3)_2SO$) δ 9.54 (br s, 1H), 8.69 (d, 1H, $J = 1.8$ Hz), 8.21 (dd, 1H, $J = 8.7, 1.8$ Hz), 8.023 (d, 1H, 8.1 Hz), 7.85-7.80 (m, 2H), 6.98 (d, 1H, 8.7 Hz), 4.14 (d, 2H, $J = 6.0$ Hz), 2.51 (m, 3H).

Example 32

Synthesis of N-[(4-hydroxy-2-oxo-7-(pyridin-4-yl)-2H-3-chromenyl)carbonyl]glycine

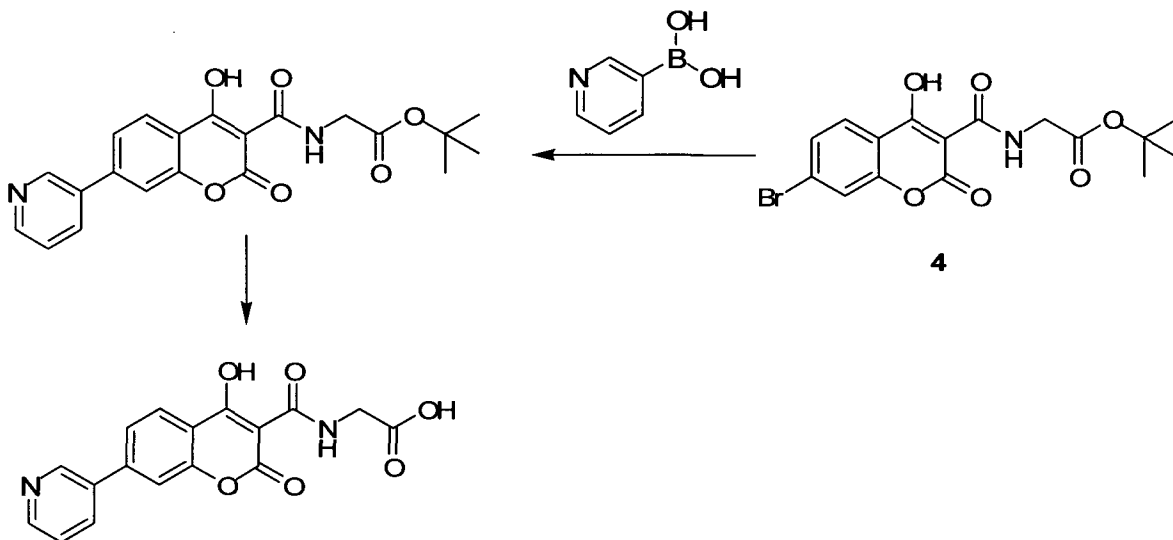


Similar procedure to the previous example was followed to arrive at the title compound, with the

5 LC-MS $[M-H]^-$ m/z 339.

Example 33

Synthesis of N-[(4-hydroxy-2-oxo-7-(pyridin-3-yl)-2H-3-chromenyl)carbonyl]glycine

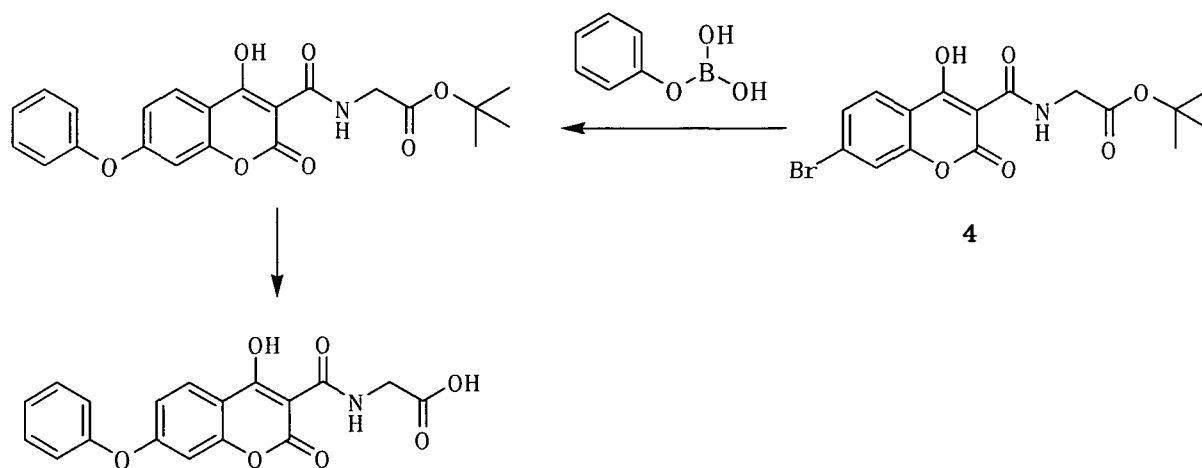


Similar procedure to the previous example was followed to arrive at the title compound, with the

10 LC-MS $[M-H]^-$ m/z 339, and 1H -NMR (300 MHz, $(CD_3)_2SO$) δ 9.55 (br s, 1H), 9.12 (s, 1H), 8.73 (s, 1H), 8.38 (d, 1H, $J = 8.1$ Hz), 8.11 (d, 1H, $J = 8.4$ Hz), 7.97 (s, 1H), 7.90 (d, 1H, $J = 8.4$ Hz), 7.69-7.65 (m, 1H), 4.16 (d, 2H, $J = 6.0$ Hz).

Example 34

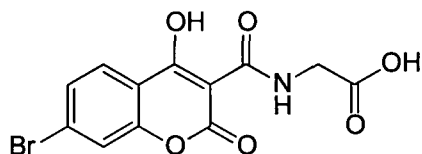
Synthesis of N-[(4-hydroxy-2-oxo-7-phenoxy-2H-3-chromenyl)carbonyl]glycine



Similar procedure to the previous example was followed to arrive at the title compound, with the LC-MS $[M-H]^-$ m/z 354, and 1H -NMR (300 MHz, $(CD_3)_2SO$) δ 9.47 (br s, 1H), 8.99 (d, 1H, 8.7 Hz), 7.55-7.49 (m, 2H), 7.33 (t, 1H, $J = 7.5$ Hz), 7.24-7.21 (m, 2H), 7.05-7.01 (m, 1H), 6.96 (d, 1H, $J = 2.4$ Hz), 4.12 (d, 2H, $J = 6.0$ Hz).

Example 35

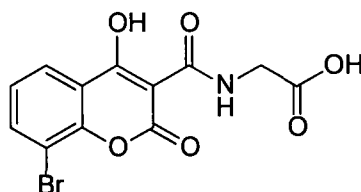
Synthesis of N-[(4-hydroxy-2-oxo-7-bromo-3-chromenyl)carbonyl]glycine



Synthesis was shown as of Compound 4 in Example 1, followed by similar procedure to de-protect the t-butyl group on the carboxylate group to arrive at the title compound with the LC-MS $[M-H]^-$ m/z 341, and 1H -NMR (300 MHz, $(CD_3)_2SO$) δ 9.53 (br s, 1H), 7.93-7.86 (m, 2H), 7.68-7.65 (m, 1H), 4.14 (m, 2H).

Example 36

Synthesis of N-[(4-hydroxy-2-oxo-8-bromo-3-chromenyl)carbonyl]glycine

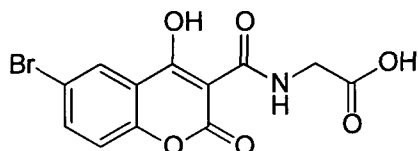


Synthesis was shown as of Compound 10 in Example 16, followed by similar procedure to de-protect the t-butyl group on the carboxylate group to arrive at the title compound with the LC-MS $[M-H]^-$ m/z 341, and 1H -NMR (300 MHz, $(CD_3)_2SO$) δ 9.51 (br s, 1H), 8.12 (d, 1H, $J = 7.8$ Hz), 8.00 (d,

1H, J = 7.8 Hz), 7.40 (t, 1H, J = 7.8 Hz), 4.15 (d, 2H, J = 6.0 Hz).

Example 37

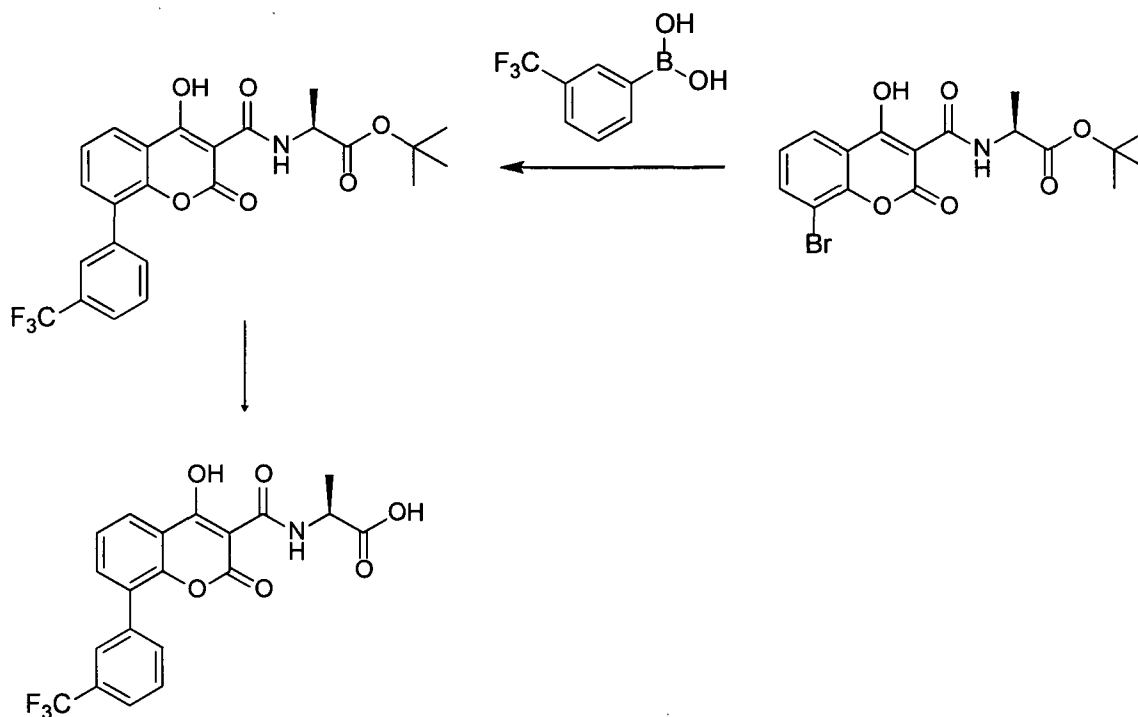
Synthesis of N-[(4-hydroxy-2-oxo-6-bromo-3-chromenyl)carbonyl]glycine



5 Similar procedure to the previous example was followed to arrive at the title compound with the LC-MS [M-H]⁻ m/z 341, and ¹H-NMR (300 MHz, (CD₃)₂SO) δ 9.56 (br s, 1H), 8.07 (d, 1H, J = 2.4 Hz), 7.99 (dd, 1H, J = 9.0, 2.4 Hz), 7.49 (d, 1H, J = 9.0 Hz), 4.14 (d, 2H, J = 6.0 Hz).

Example 38

Synthesis of N-[(4-hydroxy-2-oxo-8-(3-trifluoromethylphenyl)-2H-3-chromenyl)carbonyl]alanine



10 Similar procedure to Example 20 was followed, except that an alanine t-butyl ester was used instead of a glycine t-butyl ester, to arrive at the title compound, with the LC-MS [M-H]⁻ m/z 420, and ¹H-NMR (300 MHz, (CD₃)₂SO) δ 9.59 (br s, 1H), 8.09-8.03 (m, 2H), 7.95 (dd, 2H, J = 7.5, 1.5 Hz), 7.86-7.76 (m, 2H), 7.59 (t, 1H, 7.5 Hz), 4.57 (m, 1H), 1.48 (d, 1H, J = 7.2 Hz).

Example A. Assay of HIF-PHD2 Enzyme Activity

HIF-PHD2 activity was measured using homogeneous TR-FRET technology (see also, US2008/004817; Dao JH et al., Anal Biochem. 2009, 384:213-23). To each well of a 1/2Area 96-well

plate was added 2 μ L of test compound in DMSO and 40 μ L of assay buffer (50 mM Tris PH7.4/0.01% Tween-20/0.1 mg/ml BSA/1 mM Sodium ascorbate/20 μ g/ml Catalase/10 μ M FeSO₄) containing 600 nM full length PHD2. After a 30 min preincubation at room temperature, the enzymatic reactions were initiated by the addition of 8 μ L of substrates(final concentrations of 0.2 μ M 2-oxoglutarate and 0.5 μ M HIF-1 α peptide biotinyI-DLDLEMLAPYIPMDDDFQL). After 2 hr at room temperature, the reactions were terminated and signals were developed by the addition of a 50 μ L quench/detection mix to a final concentration of 1 mM ortho-phenanthroline, 0.1 mM EDTA, 0.5 nM anti-(His)₆LANCE reagent, 100 nM AF647-labeled Streptavidin, and 30 nM (His)₆- VHL-elonginB-elonginC complex. The ratio of time resolved fluorescence signals at 665 and 620 nm was determined, and percent inhibition was calculated relative to an uninhibited control sample run in parallel.

Example B. Determination of Erythropoietin (EPO) Induction in Normal Mice

Eight-week-old male C57BL/6 mice were dosed orally with test compound at 20, 60 and 100 mg/kg. Blood samples were obtained from the orbital venous plexus 6 hours after dosing and serum was collected (see also, Robinson A, et al., Gastroenterology. 2008, 134:145-55; Hsieh MM, et al., Blood. 2007, 110:2140-7). Samples were analyzed for EPO by electrochemiluminescence-based immunoassay (MSD) according to manufacturer's instructions.

Example C. Determination of Hematology in Normal Mice

Eight-week-old male C57BL/6 mice were dosed orally with test compound at 60 mg/kg once a day for a week. Blood samples were obtained from the orbital venous plexus with time points including 1, 3 and 5 days after dosing. An Automated Hematology Analyzer MEK-6318K was used to determine hematological parameters such as erythrocyte counts (RBC), hemoglobin concentration (HGB), hematocrit value (HCT).

Example D. Pharmacokinetics Studies

Test compounds were administered in solution as a single oral dose to fasted male CD (SD) IGS rats (n=6/group) at 50 mg/kg. Plasma samples were obtained from the orbital venous plexus of each individual animal at 15', 30', 1h, 2h, 4h, 6h, 12h and 24h post-administration. Compound blood concentrations in the plasma samples were measured by HPLC.

The IC₅₀ values for certain example compounds of invention in the Enzyme Assay of Example A are provided in Table 1 as follows.

Table 1

COMPOUND	Enzyme Assay IC50 (nM)
Example 1	250
Example 2	150
Example 3	150
Example 4	150
Example 5	150
Example 6	150
Example 7	150
Example 8	150
Example 9	150
Example 10	150
Example 11	150
Example 12	150
Example 13	150
Example 14	300
Example 15	150
Example 16	350
Example 17	200
Example 18	90
Example 19	150
Example 20	100
Example 21	90
Example 22	100
Example 23	150
Example 24	30
Example 25	150
Example 26	150
Example 27	150

Table 1, Continued

STRUCTURE	Enzyme Assay IC₅₀ (nM)
Example 28	80
Example 29	150
Example 30	80
Example 31	-
Example 32	-
Example 33	200
Example 34	150
Example 35	150
Example 36	300
Example 37	700
Example 38	-

Although the present invention has been described in considerable detail with reference to certain preferred versions thereof, other versions are possible. Therefore, the spirit and scope of the invention should not be limited to the description of the preferred versions described herein.

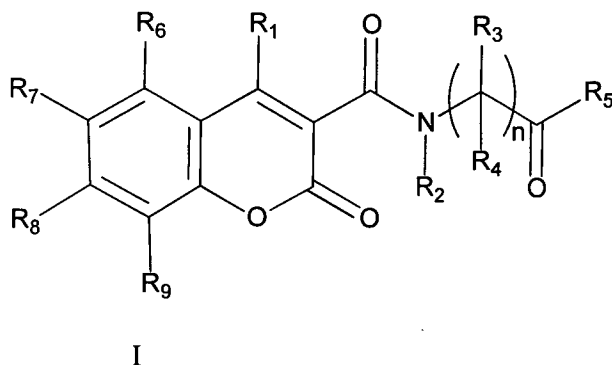
All features disclosed in the specification, including the abstract and drawings, and all the steps in any method or process disclosed, may be combined in any combination, except combinations where at least some of such features and/or steps are mutually exclusive. Each feature disclosed in the specification, including abstract and drawings, can be replaced by alternative features serving the same, equivalent or similar purpose, unless expressly stated otherwise. Thus, unless expressly stated otherwise, each feature disclosed is one example only of a generic series of equivalent or similar features. Various modifications of the invention, in addition to those described herein, will be apparent to those skilled in the art from the foregoing description. Such modifications are also intended to fall within the scope of the appended claims.

Each reference cited in the present application is herein incorporated by reference in its entirety.

THE CLAIMS

What is claimed is:

1. A compound of Formula I:



or a pharmaceutically acceptable salt thereof, or a solvate thereof, or a chelate thereof, or a non-covalent complex thereof, or a prodrug thereof, wherein:

n is 1 to 6;

R₁ is selected from OH, SH, NR₃R₄, NHC(O)R₂, NHSO₂R₂ and sulfonyl;

R₂ is selected from H, lower alkyl and substituted lower alkyl;

R₃ and R₄ are independently selected from H, lower alkyl, substituted lower alkyl, lower haloalkyl, substituted lower haloalkyl, or R₃ and R₄ can join together to form a 3 to 6 membered ring or a substituted 3 to 6 membered ring;

R₅ is selected from OH, SH, NH₂, lower alkyl, substituted lower alkyl, lower alkoxy, substituted lower alkoxy, and sulfanyl;

each of R₆, R₇, R₈ and R₉ is independently selected from H, alkyl, substituted alkyl, alkenyl, substituted alkenyl, alkynyl, substituted alkynyl, NR₃R₄, C(O)OH, OR₁₂, SR₁₂, SO₂R₁₂, CN, NO₂, halo, aryl, substituted aryl, heteroaryl, substituted heteroaryl, heteroarylalkyl, substituted heteroarylalkyl, heterocycloalkyl, substituted heterocycloalkyl, alkylsilyl, substituted alkylsilyl, alkynylsilyl, substituted alkynylsilyl, alkoxy, substituted alkoxy, alkoxycarbonyl, substituted alkoxycarbonyl, and -X-R₁₁;

or at least one of adjacent pairs R₁ and R₆, R₆ and R₇, R₇ and R₈, and R₈ and R₉, joins together to form a 4 to 7 membered ring or a substituted 4 to 7 membered ring;

X is selected from -N(R₁₀)-Y- and -Y-N(R₁₀)-;

Y is selected from C(O), SO₂, alkylene, substituted alkylene, alkenylene, substituted alkenylene, alkynylene, and substituted alkynylene;

R₁₀ is selected from H, lower alkyl, and substituted lower alkyl;

R₁₁ is selected from H, heterocycloalkyl, substituted heterocycloalkyl, aryl, substituted aryl, heteroaryl, and substituted heteroaryl; and

R₁₂ is selected from H, alkyl, substituted alkyl, alkenyl, substituted alkenyl, alkynyl, substituted alkynyl and NR₃R₄.

2. The compound of claim 1 or pharmaceutically acceptable salt thereof, wherein R₁ is selected from OH and SH.

3. The compound of claim 1 or pharmaceutically acceptable salt thereof, wherein R₁ is OH.

4. The compound of according to any one of claims 1-3 or pharmaceutically acceptable salt thereof, wherein R₂ is selected from H and lower alkyl.

5. The compound of according to any one of claims 1-3 or pharmaceutically acceptable salt thereof, wherein R₂ is H.

6. The compound of according to any one of claims 1-5 or pharmaceutically acceptable salt thereof, wherein R₃ and R₄ are each, independently, selected from H, lower alkyl, and lower haloalkyl.

7. The compound of according to any one of claims 1-5 or pharmaceutically acceptable salt thereof, wherein R₃ and R₄ are each, independently, selected from H and lower alkyl.

8. The compound of according to any one of claims 1-5 or pharmaceutically acceptable salt thereof, wherein R₃ and R₄ are each H.

9. The compound of according to any one of claims 1-5 or pharmaceutically acceptable salt thereof, wherein R₃ and R₄ join together to form a 3 to 6 membered ring or a substituted 3 to 6 membered ring.

10. The compound according to claim 9 or pharmaceutically acceptable salt thereof, wherein the 3 to 6 membered ring or the substituted 3 to 6 membered ring comprises at least one heteroatom.

11. The compound according to claim 9 or pharmaceutically acceptable salt thereof, wherein the 3 to 6 membered ring or the substituted 3 to 6 membered ring comprises at least two heteroatoms.

12. The compound of according to any one of claims 1-11 or pharmaceutically acceptable salt thereof, wherein R₅ is selected from OH, SH, NH₂, and lower alkoxy.

13. The compound of according to any one of claims 1-11 or pharmaceutically acceptable salt thereof, wherein R₅ is OH.

14. The compound of according to any one of claims 1-13 or pharmaceutically acceptable salt thereof, wherein at least one of adjacent pairs R₆ and R₇, R₇ and R₈, and R₈ and R₉, joins together to form

a 4 to 7 membered ring or a substituted 4 to 7 membered ring.

15. The compound according to claim 14 or pharmaceutically acceptable salt thereof, wherein the 4 to 7 membered ring or the substituted 4 to 7 membered ring comprises at least one heteroatom.

16. The compound according to claim 14 or pharmaceutically acceptable salt thereof, wherein the 4 to 7 membered ring or the substituted 4 to 7 membered ring comprises at least two heteroatoms.

17. The compound according to claim 14 or pharmaceutically acceptable salt thereof, wherein the 4 to 7 membered ring or the substituted 4 to 7 membered ring comprises at least three heteroatoms.

18. The compound of according to any one of claims 1-13 or pharmaceutically acceptable salt thereof, wherein at least one of R₆, R₇, R₈ and R₉ is selected from halo and a moiety substituted with at least one halo.

19. The compound of according to any one of claims 1-13 or pharmaceutically acceptable salt thereof, wherein at least one of R₆, R₇, R₈ and R₉ is independently selected from alkoxy or substituted alkoxy.

20. The compound of according to any one of claims 1-13 or pharmaceutically acceptable salt thereof, wherein at least one of R₆, R₇, R₈ and R₉ is independently selected from alkylsilyl, substituted alkylsilyl, alkynylsilyl, and substituted alkynylsilyl.

21. The compound of according to any one of claims 1-13 or pharmaceutically acceptable salt thereof, wherein at least one of R₆, R₇, R₈ and R₉ is independently selected from aryl, substituted aryl, heteroaryl, substituted heteroaryl, heterocycloalkyl, and substituted heterocycloalkyl.

22. The compound of according to any one of claims 1-13 or pharmaceutically acceptable salt thereof, wherein at least one of R₆, R₇, R₈ and R₉ is independently selected from H, alkyl, substituted alkyl, alkenyl, substituted alkenyl, alkynyl, and substituted alkynyl.

23. The compound of according to any one of claims 1-22 or pharmaceutically acceptable salt thereof, wherein at least one of R₈ and R₉ is independently selected from substituted aryl, and substituted heteroaryl.

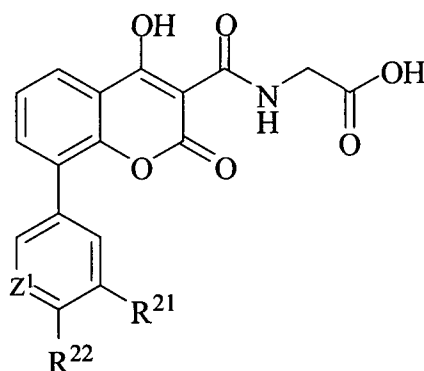
24. The compound of according to any one of claims 1-22 or pharmaceutically acceptable salt thereof, wherein at least one of R₈ and R₉ is independently selected from substituted phenyl, and substituted pyridyl.

25. The compound of according to any one of claims 1-22 or pharmaceutically acceptable salt thereof, wherein at least one of R₈ and R₉ is phenyl substituted with halogen, alkyl, haloalkyl, alkoxy, or

haloalkoxy at the para- or meta- position.

26. The compound of according to any one of claims 1-22 or pharmaceutically acceptable salt thereof, wherein at least one of R_8 and R_9 is pyridyl substituted with halogen, alkyl, haloalkyl, alkoxy, or haloalkoxy at the para- or meta- position.

5 27. The compound according to claim 1 or pharmaceutically acceptable salt thereof, wherein the compound or pharmaceutically acceptable salt thereof is a compound of Formula II:



II

or pharmaceutically acceptable salt thereof, wherein:

10 R^{21} is selected from H, halogen, lower alkyl, lower haloalkyl, lower alkoxy, and lower haloalkoxy;
 R^{22} is selected from H, halogen, lower alkyl, lower haloalkyl, lower alkoxy, and lower haloalkoxy;
 Z^1 is N or CR^{23} ; and
 R^{23} is selected from H, halogen, lower alkyl, lower haloalkyl, lower alkoxy, and lower haloalkoxy;
provided that at least one of R^{21} and R^{22} and R^{23} (if present) is selected from halogen, lower alkyl,
15 lower haloalkyl, lower alkoxy, and lower haloalkoxy.

28. The compound according to any one of claims 27 or pharmaceutically acceptable salt thereof, wherein Z^1 is CR^{23} .

29. The compound according to claim 27 or 28 or pharmaceutically acceptable salt thereof, wherein R^{21} is selected from H, Cl, F, methyl, ethyl, trifluoromethyl, methoxy, ethoxy, and trifluoromethoxy.

20 30. The compound according to claim 27 or 28 or pharmaceutically acceptable salt thereof, wherein R^{22} is selected from H, Cl, F, methyl, ethyl, trifluoromethyl, methoxy, ethoxy, and trifluoromethoxy.

31. The compound according to claim 27 or 28 or pharmaceutically acceptable salt thereof, wherein Z^1 is CR^{23} ; R^{23} is selected from H, Cl, F, methyl, ethyl, trifluoromethyl, methoxy, ethoxy, and trifluoromethoxy.

25 32. The compound according to claim 27 or 28 or pharmaceutically acceptable salt thereof, wherein

Z^1 is CR^{23} ; one of R^{21} , R^{22} , and R^{23} is selected from halogen, lower alkyl, lower haloalkyl, lower alkoxy, and lower haloalkoxy; and the other two of R^{21} , R^{22} , and R^{23} are both H.

33. The compound according to claim 27 or 28 or pharmaceutically acceptable salt thereof, wherein Z^1 is CR^{23} ; one of R^{21} and R^{23} is selected from halogen, lower alkyl, lower haloalkyl, lower alkoxy, and lower haloalkoxy; and the other one of R^{21} and R^{23} is H.

34. The compound according to claim 33 or pharmaceutically acceptable salt thereof, wherein R^{22} is H.

35. The compound according to claim 27 or 28 or pharmaceutically acceptable salt thereof, wherein Z^1 is CR^{23} ; two of R^{21} , R^{22} , and R^{23} are each independently selected from halogen, lower alkyl, lower haloalkyl, lower alkoxy, and lower haloalkoxy; and the other one of R^{21} , R^{22} , and R^{23} is H.

36. The compound according to claim 35 or pharmaceutically acceptable salt thereof, wherein R^{22} is H.

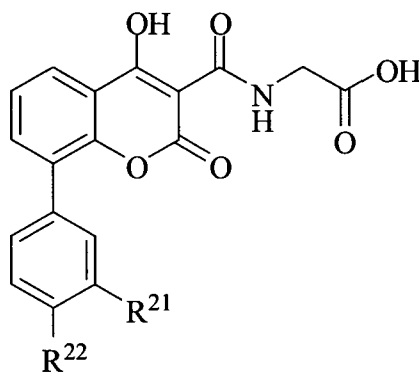
37. The compound according to claim 27 or 28 or pharmaceutically acceptable salt thereof, wherein Z^1 is CR^{23} ; R^{21} , R^{22} , and R^{23} are each independently selected from halogen, lower alkyl, lower haloalkyl, lower alkoxy, and lower haloalkoxy.

38. The compound according to claim 27 or pharmaceutically acceptable salt thereof, wherein Z^1 is N.

39. The compound according to claim 27 or 38, or pharmaceutically acceptable salt thereof, wherein Z^1 is N; one of R^{21} and R^{22} is selected from halogen, lower alkyl, lower haloalkyl, lower alkoxy, and lower haloalkoxy; and the other of R^{21} and R^{22} is H.

40. The compound according to claim 27 or 38, or pharmaceutically acceptable salt thereof, wherein Z^1 is N; R^{21} and R^{22} are each independently selected from halogen, lower alkyl, lower haloalkyl, lower alkoxy, and lower haloalkoxy.

41. The compound according to claim 27 or pharmaceutically acceptable salt thereof, wherein the compound or pharmaceutically acceptable salt thereof is a compound of Formula IIa:



IIa

or pharmaceutically acceptable salt thereof, wherein:

R^{21} is selected from H, halogen, lower alkyl, lower haloalkyl, lower alkoxy, and lower haloalkoxy;

5 and

R^{22} is selected from H, halogen, lower alkyl, lower haloalkyl, lower alkoxy, and lower haloalkoxy;

provided that at least one of R^{21} and R^{22} is selected from halogen, lower alkyl, lower haloalkyl, lower alkoxy, and lower haloalkoxy.

42. The compound according to claim 41 or pharmaceutically acceptable salt thereof, wherein R^{21} is
10 H; and R^{22} is selected from halogen, lower alkyl, lower haloalkyl, lower alkoxy, and lower haloalkoxy.

43. The compound according to claim 41 or pharmaceutically acceptable salt thereof, wherein R^{21} is
H; and R^{22} is selected from lower alkyl and lower haloalkyl.

44. The compound according to claim 41 or pharmaceutically acceptable salt thereof, wherein R^{21} is
H; and R^{22} is selected from C_{1-3} alkyl and C_{1-3} haloalkyl.

15 45. The compound according to claim 44 or pharmaceutically acceptable salt thereof, wherein R^{22} is
selected from methyl and C_1 haloalkyl.

46. The compound according to claim 44 or pharmaceutically acceptable salt thereof, wherein R^{22} is
selected from C_{1-3} alkyl.

47. The compound according to claim 44 or pharmaceutically acceptable salt thereof, wherein R^{22} is
20 selected from selected from methyl or ethyl.

48. The compound according to claim 41 or pharmaceutically acceptable salt thereof, wherein R^{22} is
H; and R^{21} is selected from halogen, lower alkyl, lower haloalkyl, lower alkoxy, and lower haloalkoxy.

49. The compound according to claim 41 or pharmaceutically acceptable salt thereof, wherein R^{22} is
H; and R^{21} is selected from lower alkyl and lower haloalkyl.

25 50. The compound according to claim 49 or pharmaceutically acceptable salt thereof, wherein R^{21} is

selected from methyl and C₁ haloalkyl.

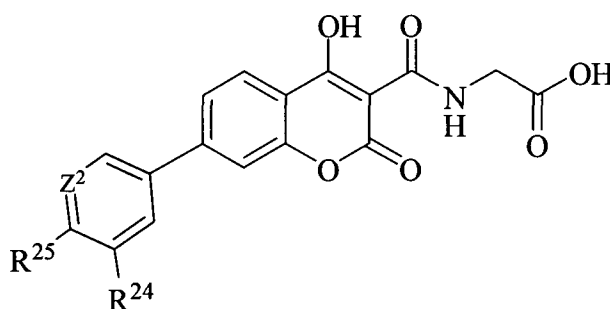
51. The compound according to claim 49 or pharmaceutically acceptable salt thereof, wherein R²¹ is selected from C₁₋₃ alkyl.

52. The compound according to claim 49 or pharmaceutically acceptable salt thereof, wherein R²¹ is selected from methyl or ethyl.

53. The compound according to claim 41 or pharmaceutically acceptable salt thereof, wherein R²¹ and R²² are each independently selected from halogen, lower alkyl, lower haloalkyl, lower alkoxy, and lower haloalkoxy.

54. The compound according to claim 41 or pharmaceutically acceptable salt thereof, wherein R²¹ and R²² are each independently selected from C₁₋₂ alkyl and C₁₋₂ haloalkyl.

55. The compound according to claim 1 or pharmaceutically acceptable salt thereof, wherein the compound or pharmaceutically acceptable salt thereof is a compound of Formula III:



III

or pharmaceutically acceptable salt thereof, wherein:

R²⁴ is selected from H, halogen, lower alkyl, lower haloalkyl, lower alkoxy, and lower haloalkoxy;

R²⁵ is selected from H, halogen, lower alkyl, lower haloalkyl, lower alkoxy, and lower haloalkoxy;

Z² is N or CR²⁶; and

R²⁶ is selected from H, halogen, lower alkyl, lower haloalkyl, lower alkoxy, and lower haloalkoxy;

provided that at least one of R²⁴ and R²⁵ and R²⁶ (if present) is selected from halogen, lower alkyl, lower haloalkyl, lower alkoxy, and lower haloalkoxy.

56. The compound according to claim 55 or pharmaceutically acceptable salt thereof, wherein Z² is CR²⁶.

57. The compound according to claim 55 or 56 or pharmaceutically acceptable salt thereof, wherein R²⁴ is selected from H, Cl, F, methyl, ethyl, trifluoromethyl, methoxy, ethoxy, and trifluoromethoxy.

58. The compound according to any one of claims 55-57 or pharmaceutically acceptable salt thereof,

wherein R^{25} is selected from H, Cl, F, methyl, ethyl, trifluoromethyl, methoxy, ethoxy, and trifluoromethoxy.

59. The compound according to any one of claims 55-58 or pharmaceutically acceptable salt thereof, wherein Z^2 is CR^{26} ; R^{26} is selected from H, Cl, F, methyl, ethyl, trifluoromethyl, methoxy, ethoxy, and trifluoromethoxy.

60. The compound according to claim 55 or 56 or pharmaceutically acceptable salt thereof, wherein Z^2 is CR^{26} ; one of R^{24} , R^{25} , and R^{26} is selected from halogen, lower alkyl, lower haloalkyl, lower alkoxy, and lower haloalkoxy; and the other two of R^{24} , R^{25} , and R^{26} are both H.

61. The compound according to claim 55 or 56 or pharmaceutically acceptable salt thereof, wherein Z^2 is CR^{26} ; one of R^{24} and R^{26} is selected from halogen, lower alkyl, lower haloalkyl, lower alkoxy, and lower haloalkoxy; and the other one of R^{24} and R^{26} is H.

62. The compound according to claim 61 or pharmaceutically acceptable salt thereof, wherein R^{25} is H.

63. The compound according to claim 55 or 56 or pharmaceutically acceptable salt thereof, wherein Z^1 is CR^{26} ; two of R^{24} , R^{25} , and R^{26} are each independently selected from halogen, lower alkyl, lower haloalkyl, lower alkoxy, and lower haloalkoxy; and the other one of R^{24} , R^{25} , and R^{26} is H.

64. The compound according to claim 63 or pharmaceutically acceptable salt thereof, wherein R^{25} is H.

65. The compound according to claim 55 or 56 or pharmaceutically acceptable salt thereof, wherein Z^2 is CR^{26} ; R^{24} , R^{25} , and R^{26} are each independently selected from halogen, lower alkyl, lower haloalkyl, lower alkoxy, and lower haloalkoxy.

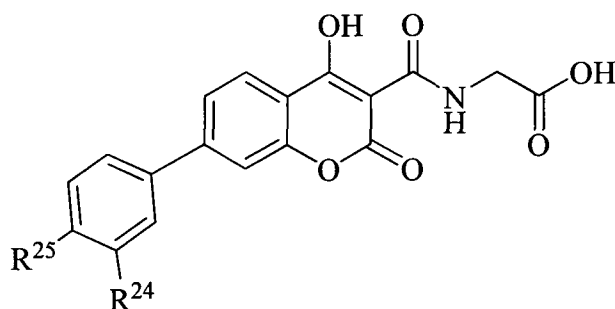
66. The compound according to any one of claims 55, 57-58, or pharmaceutically acceptable salt thereof, wherein Z^2 is N.

67. The compound according to claim 55 or 66, or pharmaceutically acceptable salt thereof, wherein Z^2 is N; one of R^{24} and R^{25} is selected from halogen, lower alkyl, lower haloalkyl, lower alkoxy, and lower haloalkoxy; and the other of R^{24} and R^{25} is H.

68. The compound according to claim 55 or 66, or pharmaceutically acceptable salt thereof, wherein Z^2 is N; R^{24} and R^{25} are each independently selected from halogen, lower alkyl, lower haloalkyl, lower alkoxy, and lower haloalkoxy.

69. The compound according to claim 55 or pharmaceutically acceptable salt thereof, wherein the

compound or pharmaceutically acceptable salt thereof is a compound of Formula IIIa



IIIa

or pharmaceutically acceptable salt thereof, wherein:

- 5 R^{24} is selected from H, halogen, lower alkyl, lower haloalkyl, lower alkoxy, and lower haloalkoxy;
and

R^{25} is selected from H, halogen, lower alkyl, lower haloalkyl, lower alkoxy, and lower haloalkoxy;
provided that at least one of R^{24} and R^{25} is selected from halogen, lower alkyl, lower haloalkyl,
lower alkoxy, and lower haloalkoxy.

- 10 70. The compound according to claim 69 or pharmaceutically acceptable salt thereof, wherein R^{24} is
H; and R^{25} is selected from halogen, lower alkyl, lower haloalkyl, lower alkoxy, and lower haloalkoxy.

71. The compound according to claim 69 or pharmaceutically acceptable salt thereof, wherein R^{24} is
H; and R^{25} is selected from lower alkyl and lower haloalkyl.

- 15 72. The compound according to claim 69 or pharmaceutically acceptable salt thereof, wherein R^{24} is
H; and R^{25} is selected from C_{1-3} alkyl and C_{1-3} haloalkyl.

73. The compound according to claim 72 or pharmaceutically acceptable salt thereof, wherein R^{25} is
selected from methyl and C_1 haloalkyl.

74. The compound according to claim 72 or pharmaceutically acceptable salt thereof, wherein R^{25} is
selected from selected from C_{1-3} alkyl.

- 20 75. The compound according to claim 72 or pharmaceutically acceptable salt thereof, wherein R^{25} is
selected from selected from methyl or ethyl.

76. The compound according to claim 69 or pharmaceutically acceptable salt thereof, wherein R^{25} is
H; and R^{24} is selected from halogen, lower alkyl, lower haloalkyl, lower alkoxy, and lower haloalkoxy.

- 25 77. The compound according to claim 69 or pharmaceutically acceptable salt thereof, wherein R^{25} is
H; and R^{24} is selected from lower alkyl and lower haloalkyl.

78. The compound according to claim 77 or pharmaceutically acceptable salt thereof, wherein R^{24} is

selected from methyl and C₁ haloalkyl.

79. The compound according to claim 77 or pharmaceutically acceptable salt thereof, wherein R²⁴ is selected from selected from C₁₋₃ alkyl.

80. The compound according to claim 77 or pharmaceutically acceptable salt thereof, wherein R²⁴ is
5 selected from selected from methyl or ethyl.

81. The compound according to claim 69 or pharmaceutically acceptable salt thereof, wherein R²⁴ and R²⁵ are each independently selected from halogen, lower alkyl, lower haloalkyl, lower alkoxy, and lower haloalkoxy.

82. The compound according to claim 69 or pharmaceutically acceptable salt thereof, wherein R²⁴
10 and R²⁵ are each independently selected from C₁₋₂ alkyl and C₁₋₂ haloalkyl.

83. A compound selected from Examples 1-15, Examples 17-33, and Examples 35-38, or pharmaceutically acceptable salt thereof.

84. A pharmaceutical composition comprising at least one pharmaceutically acceptable excipient, adjuvant or carrier, and a therapeutically effective amount of at least one compound according to any
15 one of Claims 1-83 or pharmaceutically acceptable salt thereof.

85. A pharmaceutical composition comprising at least one pharmaceutically acceptable excipient, adjuvant, or carrier, and a therapeutically effective amount of at least one compound according to any one of Claims 1-83 or pharmaceutically acceptable salt thereof, with at least one additional compound.

86. The pharmaceutical composition according to claim 85, wherein the said additional compound is
20 selected from an erythropoiesis stimulating agent or chemotherapeutic agent.

87. The pharmaceutical composition according to any one of claims 84-86, wherein the at least one compound is present in an amount effective for the treatment of at least one disease selected from ischemia, anemia, and thalassemia.

88. A method of modulating HIF levels or activity in a subject administering to the subject at least
25 one compound according to any one of Claims 1-83 or pharmaceutically acceptable salt thereof.

89. A method for treating a disease, disorder, or condition in a patient comprising administering the patient a therapeutically effective amount of at least one compound according to any one of Claims 1-83 or pharmaceutically acceptable salt thereof, wherein the disease, disorder, or condition is associated with HIF level or HIF activity.

30 90. Use of a compound according to any one of Claims 1-83 or pharmaceutically acceptable salt

thereof in treatment of a disease, disorder, or condition associated with HIF level or HIF activity.

91. Use of a compound according to any one of Claims 1-83 or pharmaceutically acceptable salt thereof in the manufacturing a medicament for the treatment of a disease, disorder, or condition associated with HIF level or HIF activity.

5 92. A method for treating a disease, disorder, or condition in a patient comprising administering the patient a therapeutically effective amount of at least one compound according to any one of Claims 1-83 or pharmaceutically acceptable salt thereof, wherein the disease, disorder, or condition is ischemia, anemia, wound healing, auto-transplantation, allo-transplantation, xeno-transplantation, systemic high blood pressure, thalassemia, diabetes, cancer or an inflammatory disorder, or a combination of two or
10 more thereof.

93. A method for treating a disease, disorder, or condition in a patient comprising administering the patient a therapeutically effective amount of at least one compound according to any one of Claims 1-83 or pharmaceutically acceptable salt thereof, wherein the disease, disorder, or condition is an anemic or ischemic related disorder.

15 94. A method of modulating the amount of HIF in a cell comprising contacting the cell with at least one compound according to any one of Claims 1-83.

95. Use of a compound according to any one of Claims 1-83 or pharmaceutically acceptable salt thereof in treatment of a disease, disorder, or condition that is an anemic or ischemic related disorder.

20 96. Use of a compound according to any one of Claims 1-83 or pharmaceutically acceptable salt thereof in the manufacturing a medicament for the treatment of a disease, disorder, or condition that is an anemic or ischemic related disorder.

97. Use of a compound according to any one of Claims 1-83 or pharmaceutically acceptable salt thereof in treatment of a disease, disorder, or condition selected from ischemia, anemia, and thalassemia.

25 98. Use of a compound according to any one of Claims 1-83 or pharmaceutically acceptable salt thereof in the manufacturing a medicament for the treatment of a disease, disorder, or condition selected from ischemia, anemia, and thalassemia.

99. The compound according to any one of Claims 1-83, wherein the HIF PHD inhibitory activity IC₅₀ value of the compound is 40 μ M or less.

30 100. The compound according to any one of Claims 1-83, wherein the HIF PHD inhibitory activity IC₅₀ value is 10 μ M or less.