METHODS OF USING AND COMPOSITIONS COMPRISING IMMUNOMODULATORY COMPOUNDS FOR THE TREATMENT AND MANAGEMENT OF MYELODYSPLASTIC SYNDROMES

1. FIELD OF THE INVENTION

This invention relates to methods of treating, preventing and/or managing myelodysplastic and related syndromes which comprise the administration of immunomodulatory compounds alone or in combination with known therapeutics. The invention also relates to pharmaceutical compositions and dosing regimens. In particular, the invention encompasses the use of immunomodulatory compounds in conjunction with transplantation therapy and/or other standard therapies for myelodysplastic syndromes.

2. BACKGROUND OF THE INVENTION

2.1. PATHOBIOLOGY OF MDS

Myelodysplastic syndrome ("MDS") refers to a diverse group of hematopoietic stem cell disorders. MDS is characterized by a cellular marrow with impaired morphology and maturation (dysmyelopoiesis), peripheral blood cytopenias, and a variable risk of progression to acute leukemia, resulting from ineffective blood cell production. *The Merck Manual* 953 (17th ed. 1999) and List *et al.*, 1990, *J. Clin. Oncol.* 8:1424.

The initial hematopoietic stem cell injury can be from causes such as, but not limited to, cytotoxic chemotherapy, radiation, virus, chemical exposure, and genetic predisposition. A clonal mutation predominates over bone marrow, suppressing healthy stem cells. In the early stages of MDS, the main cause of cytopenias is increased programmed cell death (apoptosis). As the disease progresses and converts into leukemia, gene mutation rarely occurs and a proliferation of leukemic cells overwhelms the healthy marrow. The disease course differs, with some cases behaving as an indolent disease and others behaving aggressively with a very short clinical course that converts into an acute form of leukemia.

The actual incidence of MDS in the U.S. is unknown. MDS was first considered a distinct disease in 1976, and occurrence was estimated at 1500 new cases every year. At that time, only patients with less than five percent blasts were considered to have this disorder. Statistics from 1999 estimated 13,000 new cases per year and about 1000 cases per year in children, surpassing chronic lymphocytic leukemia as the most common form of leukemia in the western hemisphere. The perception that the incidence is increasing may be due to improvements in recognition and criteria for diagnosis. The disease is found worldwide.

An international group of hematologists, the French-American-British (FAB) Cooperative Group, classified MDS disorders into five subgroups, differentiating them from acute myeloid leukemia. *The Merck Manual* 954 (17th ed. 1999); Bennett J.M., *et al.*, *Ann. Intern. Med.* 1985 Oct., 103(4): 620-5; and Besa E.C., *Med. Clin. North Am.* 1992 May, 76(3): 599-617. An underlying trilineage dysplastic change in the bone marrow cells of the patients is found in all subtypes.

There are two subgroups of refractory anemia characterized by five percent or less myeloblasts in bone marrow: (1) refractory anemia (RA) and; (2) RA with ringed sideroblasts (RARS), defined morphologically as having 15% erythroid cells with abnormal ringed sideroblasts, reflecting an abnormal iron accumulation in the mitochondria. Both have a prolonged clinical course and low incidence of progression to acute leukemia. Besa E.C., *Med. Clin. North Am.* 1992 May, 76(3): 599-617.

There are two subgroups of refractory anemias with greater than five percent myeloblasts: (1) RA with excess blasts (RAEB), defined as 6-20% myeloblasts, and (2) RAEB in transformation (RAEB-T), with 21-30% myeloblasts. The higher the percentage of myeloblasts, the shorter the clinical course and the closer the disease is to acute myelogenous leukemia. Patient transition from early to more advanced stages indicates that these subtypes are merely stages of disease rather than distinct entities. Elderly patients with MDS with trilineage dysplasia and greater than 30% myeloblasts who progress to acute leukemia are often considered to have a poor prognosis because their response rate to chemotherapy is lower than *de novo* acute myeloid leukemia patients. The World Health Organization (WHO) classification (1999) proposes to include all cases of RAEB-T, or patients with greater than 20% myeloblasts, in the category of acute leukemia because these patients have similar prognostic outcomes. However, their response to therapy is worse than the *de novo* or more typical acute myelogenous leukemia or acute nonlymphocytic leukemia (ANLL) patient. *Id.*

The fifth type of MDS, the most difficult to classify, is called chronic myelomonocytic leukemia (CMML). This subtype can have any percentage of myeloblasts but presents with a monocytosis of 1000/dL or more. It may be associated with splenomegaly. This subtype overlaps with a myeloproliferative disorder and may have an intermediate clinical course. It is differentiated from the classic chronic myelocytic leukemia (CML) that is characterized by a negative Ph chromosome. The recent WHO classification (1999) proposes that juvenile and proliferative CMML be listed separately from FAB under MDS/myeloproliferative disorder (MPD) with splenomegaly and greater than 13,000 total WBC. CMML is limited to monocytosis, less than 13,000/mm^3 total

leukocytes, and requires trilineage dysplasia. *Id.* Harris N.L., *et al.*, *J. Clin. Oncol.* 1999 Dec., 17(12): 3835-49. Finally, some other international organizations, including WHO, have suggested a sixth class of MDS patients, characterized by a del (5q) abnormality.

MDS is primarily a disease of elderly people, with the median onset in the seventh decade of life. The median age of these patients is 65 years, with ages ranging from the early third decade of life to as old as 80 years or older. The syndrome may occur in any age group, including the pediatric population. Patients who survive malignancy treatment with alkylating agents, with or without radiotherapy, have a high incidence of developing MDS or secondary acute leukemia. About 60-70% of patients do not have an obvious exposure or cause for MDS, and are classified as primary MDS patients.

The most common cases of MDS are primary, or idiopathic. However, a nonspecific history of exposure to indeterminable chemicals or radiation 10-15 years prior to onset of disease may be present in about 50% of patients. This relationship to pathogenesis remains unproved. Compounds such as, but not limited to, benzene, insecticides, weed killers, and fungicides are possible causes of MDS. Goldberg H., *et al.*, *Cancer Res.* 1990 Nov 1; 50(21): 6876-81. Secondary MDS describes development of MDS or acute leukemia after known exposures to chemotherapy drugs that can cause bone marrow damage. These drugs are associated with a high incidence of chromosomal abnormalities following exposure and at the time of MDS or acute leukemia diagnosis.

Further, MDS is associated with complications associated with severe cytopenias. Other complications are development of myelofibrosis, which can accelerate decline in blood counts and increase transfusion requirements. Transformation to acute leukemia accelerates the development of complications such as anemia, bleeding, and infections.

Recently, the International MDS Risk Analysis (IMRA) Workshop proposed an International Prognosis Scoring System (IPSS) to decrease imprecision in predicting survival and AML risk in MDS patients. The IPSS is based on the number of cytopenias, percentage of BM blasts, and type of cytogenetic abnormalities (Table 1). Greenberg *et al.*, *Blood* 1997, 89:2079-88. The latter are categorized into good (normal, -Y, del (5q), del (20q)), intermediate, and poor subgroups (complex or chromosome 7 abnormalities).

Table 1. International Prognostic Scoring System for MDS					
	Score Value				
Prognostic Variable	0	0.5	1.0	1.5	2.0
Bone marrow blasts (%)	<5	5-10	—	11-20	21-30
Karyotype*	Good	Intermediate	Poor		
Cytopenias	0/1	2/3			
*Good, normal, del (5q), del (20q), -Y; Poor, complex (>3) or chromosome 7 abnormalities; Intermediate, +8, and other single or double abnormalities.					

2.2. MDS TREATMENT

The current treatment of MDS is based on the stage and the mechanism of the disease that predominates the particular phase of the disease process. Bone marrow transplantation has been used in patients with poor prognosis or late-stage MDS. Epstein and Slease, 1985, *Surg. Ann.* 17:125. This type of therapy, however, is both painful for donor and recipient, because of the involvement of invasive procedures and can cause severe and even fatal complications to the recipient, particularly with allogeneic transplant and related Graft Versus Host Disease (GVHD) results. Therefore, the risk of GVHD restricts the use of bone marrow transplantation to patients with otherwise fatal diseases. Further, as most patients are elderly and only a few young MDS patients will have a matched donor, the use of bone marrow transplantation is limited.

An alternative approach to therapy for MDS is the use of hematopoietic growth factors or cytokines to stimulate blood cell development in a recipient. Dexter, 1987, *J. Cell Sci.* 88:1; Moore, 1991, *Annu. Rev. Immunol.* 9:159; and Besa E.C., *Med. Clin. North Am.* 1992 May, 76(3): 599-617. The process of blood cell formation, by which a small number of self-renewing stem cells give rise to lineage specific progenitor cells that subsequently undergo proliferation and differentiation to produce the mature circulating blood cells has been shown to be at least in part regulated by specific hormones. These hormones are collectively known as hematopoietic growth factors. Metcalf, 1985, *Science* 229:16; Dexter, 1987, *J. Cell Sci.* 88:1; Golde and Gasson, 1988, *Scientific American*, July:62; Tabbara and Robinson, 1991, *Anti-Cancer Res.* 11:81; Ogawa, 1989, *Environ. Health Presp.* 80:199; and Dexter, 1989, *Br. Med. Bull.* 45:337. The most well characterized growth factors include erythropoietin (EPO), granulocyte macrophage colony stimulating factor (GM-CSF), and granulocyte colony stimulating factor (G-CSF). Apart from inducing proliferation and differentiation of hematopoietic progenitor cells, such cytokines have also

been shown to activate a number of functions of mature blood cells, including influencing the migration of mature hematopoietic cells. Stanley *et al.*, 1976, *J. Exp. Med.* 143:631; Schrader *et al.*, 1981, *Proc. Natl. Acad. Sci.* U.S.A. 78:323; Moore *et al.*, 1980, *J. Immunol.* 125:1302; Kurland *et al.*, 1979, *Proc. Natl. Acad. Sci. U.S.A.* 76:2326; Handman and Burgess, 1979, *J. Immunol.* 122:1134; Vadas *et al.*, 1983, *Blood* 61:1232; Vadas *et al.*, 1983, *J. Immunol.* 130:795; and Weibart *et al.*, 1986, *J. Immunol.* 137:3584.

Unfortunately, hematopoietic growth factors have not proven effective in many clinical settings. Clinical trials of MDS patients treated with recombinant human GM-CSF and G-CSF have shown that while these cytokines can restore granulocytopoiesis in treated patients, their efficacy is restricted to the granulocyte or monocyte lineage with little or no improvement in hemoglobin or platelet counts. Schuster *et al.*, 1990, *Blood* 76 (Suppl.1):318a. When such patients were treated with recombinant human EPO, a sustained improvement in hemoglobin or decrease in transfusion requirement was achieved in only less than 25% of patients. Besa *et al.*, 1990, 76 (Suppl.1):133a; Hellstrom *et al.*, 1990, 76 (Suppl.1):279a; Bowen *et al.*, 1991, *Br. J. Haematol.* 77:419. Therefore, there remains a need for safe and effective methods of treating and managing MDS.

2.3. THALIDOMIDE AND OTHER COMPOUNDS USEFUL IN THE TREATMENT OF DISEASE

Thalidomide is a racemic compound sold under the tradename Thalomid® and chemically named α-(N-phthalimido)glutarimide or 2-(2,6-dioxo-3-piperidinyl)-1H-isoindole-1,3(2H)-dione. Thalidomide was originally developed in the 1950's to treat morning sickness, but due to its teratogenic effects was withdrawn from use. Thalidomide has been approved in the United States for the acute treatment of the cutaneous manifestations of erythema nodosum leprosum in leprosy. Physicians' Desk Reference, 1154-1158 (56th ed., 2002). Because its administration to pregnant women can cause birth defects, the sale of thalidomide is strictly controlled. Id. Thalidomide has reportedly been studied in the treatment of other diseases, such as chronic graft-vs-host disease, rheumatoid arthritis, sarcoidosis, several inflammatory skin diseases, and inflammatory bowel disease. See generally, Koch, H.P., Prog. Med. Chem. 22:165-242 (1985). See also, Moller, D.R., et al., J. Immunol. 159:5157-5161 (1997); Vasiliauskas, E.A., et al., Gastroenterology 117:1278-1287 (1999); Ehrenpreis, E.D., et al., Gastroenterology 117:1271-1277 (1999). It has further been alleged that thalidomide can be combined with other drugs to treat ischemia/repercussion associated with coronary and cerebral occlusion. See U.S. Patent No. 5,643,915, which is incorporated herein by reference.

More recently, thalidomide was found to exert immunomodulatory and anti-inflammatory effects in a variety of disease states, cachexia in AIDS, and opportunic infections in AIDS. In studies to define the physiological targets of thalidomide, the drug was found to have a wide variety of biological activities exclusive of its sedative effect including neurotoxicity, teratogenicity, suppression of TNF-α production by monocytes/macrophages and the accompanying inflammatory toxicities associated with high levels of TNF-α, and inhibition of angiogenesis and neovascularization.

Additionally, beneficial effects have been observed in a variety of dermatological conditions, ulcerative colitis, Crohn's disease, Bechets's syndrome, systemic lupus erythematosis, aphthous ulcers, and lupus. The anti-angiogenic properties of thalidomide in *in vivo* models have been reported. D'Amato *et al.*, *Thalidomide Is An Inhibitor Of Angiogenesis*, 1994, *PNAS, USA* 91:4082-4085.

One of the most therapeutically significant potential uses of thalidomide is in the treatment of cancer. The compound has been investigated in the treatment of various types of cancer, such as refractory multiple myeloma, brain, breast, colon, and prostate cancer, melanoma, mesothelioma, and renal cell carcinoma. *See, e.g.*, Singhal, S., *et al.*, *New England J. Med.* 341(21):1565-1571 (1999); and Marx, G.M., *et al.*, *Proc. Am. Soc. Clin. Oncology* 18:454a (1999). Thalidomide reportedly can also be used to prevent the development of chronic cardiomyopathy in rats caused by doxorubicin. Costa, P.T., *et al.*, *Blood* 92(10:suppl. 1):235b (1998). Other reports concerning the use of thalidomide in the treatment of specific cancers include its combination with carboplatin in the treatment of glioblastoma multiforme. McCann, J., *Drug Topics* 41-42 (June 21, 1999). The use of thalidomide in combination with dexamethasone reportedly was effective in the treatment of patients suffering from multiple myeloma who also received, as supportive care, human granulocyte colony-stimulating factor (G-CSF), ciprofloxacin, and non-absorbable antifungal agents. Kropff, M.H., *Blood* 96(11 part 1):168a (2000); *see also*, Munshi, N. *et al.*, *Blood* 94(10 part 1):578a (1999). Other chemotherapy combinations that comprise thalidomide are disclosed in International Application No. PCT/US01/15326 to R. Govindarjan and A. Zeitlan, and in International Application No. PCT/US01/15327 to J.B. Zeldis, *et al.*

In an effort to provide compounds that have greater therapeutic safety and efficacy than thalidomide, researchers have begun investigating a large number of other compounds, some of which are derivatives of thalidomide. *See, e.g.*, Marriott, J.B., *et al.*, *Expert Opin. Biol. Ther.* 1(4):1-8 (2001); G.W. Muller, *et al.*, Journal of Medicinal Chemistry 39(17): 3238-3240 (1996); and G.W. Muller, *et al.*, Bioorganic & Medicinal Chemistry Letters 8:

2669-2674 (1998). Examples include, but are not limited to, the substituted 2-(2,6-dioxopiperidin-3-yl) phthalimies and substituted 2-(2,6-dloxopiperidm-3-yl)-l-oxoisoindoles described in United States Patent Nos. 6,281,230 and 6,316,471, both to G.W.Mulier, *et al.*

A group of compounds selected for their capacity to potently inhibit TNP-α production by LPS stimulated PBMC has been investigated. L.G. Corral, *et al,* Ann. Rheum. Dis. 58:(Suppl 1) 1107-1113 (1999). These compounds, which are referred to as IMiDs™ or Immunomodulatory Drugs, show not only potent inhibition of TNF-α but also marked inhibition of LPS induced monocyte IL1β and IL12 production. LPS induced IL6 is also inhibited by IMiDs™, *albeit* partially. These compounds are potent stimulators of LPS induced IL10, increasing IL10 levels by 200 to 300%. *Id.*

While many such compounds have shown promise as therapeutic agents, their mechanisms of action and effectiveness are still under investigation. Moreover, there remains a need for therapeutic agents to treat MDS and its related disorders.

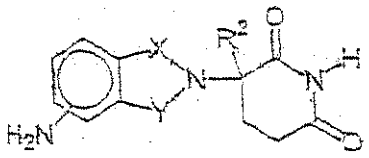
3. SUMMARY OF THE INVENTION

Embodiments of the present disclosure encompass methods of treating or preventing myelodysplastic syndrome ("MDS") which comprise administering to a patient in need thereof a therapeutically or prophylactically effective amount of an immunomodulatory compound of the invention or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof. The disclosure also encompasses methods of managing MDS *(e.g.,* lengthening the time of remission) which comprise administering to a patient in need of such management a therapeutically or prophylactically effective amount of an immunomodulatory compound disclosed herein, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof.

One embodiment of the disclosure encompasses the use of one or more immunomodulatory compounds in combination with conventional therapies presently used to treat, prevent or manage MDS such as hematopoietic growth factors, cytokines, cancer chemotherapeutics, stem cell transplantation and other transplantations.

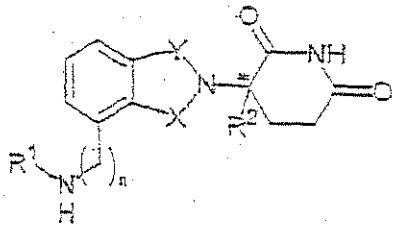
The disclosure further encompasses pharmaceutical compositions, single unit dosage forms, and kits suitable for use in treating, preventing and/or managing MDS, which comprise an immunomodulatory compound disclosed herein, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof.

In a first aspect, there is provided a use of therapeutically or prophylactically effective amount of an immunomodulatory compound, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof for manufacturing a medicament in treating or preventing a myelodysplastic syndrome to a patient in need thereof, wherein the immunomodulatory compound comprises 4-(amino)-2-(2,6-dioxo(3-piperidyl))- isoindoline-1,3-dione, 3 -(4-amino- 1 -oxo- 1,3 -dihydro-isoindol-2- yl)-piperidine-2, 6-dione, formula (I) and (II):



(I)

wherein one of X and Y is C=O, the other of X and Y is C=0 or CH_2, and R^2 is hydrogen or lower alkyl, formula (II):



(II)

wherein

one of X and Y is C=O and the other is CH_2 or C=O;

R^1 is H, (C_1-C_8)alkyl, (C_3-C_7)cycloalkyl, (C_2-C_8)alkenyl, (C_2-C_8)alkynyl, benzyl, aryl, (C_0-C_4)alkyl-(C_1-C_6)heterocycloalkyl, (C_0-C_4)alkyl-(C_2-C_5)heteroaryl, $C(O)R^3$, $C(S)R^3$, $C(O)OR^4$, (C_1-C_8)alkyl-$N(R^6)_2$, (C_1-C_8)alkyl-OR^5, (C_1-C_8)alkyl-$C(O)OR^5$, $C(O)NHR^3$, $C(S)NHR^3$, $C(O)NR^3R^{3'}$, $C(S)NR^3R^{3'}$ or (C_1-C_8)alkyl-$O(CO)R^5$;

R^2 is H, F, benzyl, $(C_1$-$C_8)$alkyl, $(C_2$-$C_8)$alkenyl, or $(C_2$-$C_8)$alkynyl;

R^3 and $R^{3'}$ are independently $(C_1$-$C_8)$alkyl, $(C_3$-$C_7)$cycloalkyl, $(C_2$-$C_8)$alkenyl, $(C_2$-$C_8)$alkynyl, benzyl, aryl, $(C_0$-$C_4)$alkyl-$(C_1$-$C_6)$heterocycloalkyl, $(C_0$-$C_4)$alkyl-$(C_2$-$C_5)$heteroaryl, $(C_0$-$C_8)$alkyl-$N(R^6)_2$, $(C_1$-$C_8)$alkyl-OR^5, $(C_1$-$C_8)$alkyl-$C(O)OR^5$, $(C_1$-$C_8)$alkyl-$O(CO)R^5$, or $C(O)OR^5$;

R^4 is $(C_1$-$C_8)$alkyl, $(C_2$-$C_8)$alkenyl, $(C_2$-$C_8)$alkynyl, $(C_1$-$C_4)$alkyl- OR^5, benzyl, aryl, $(C_0$-$C_4)$alkyl-$(C_1$-$C_6)$heterocycloalkyl, or $(C_0$-$C_4)$alkyl-$(C_2$-$C_5)$heteroaryl;

R^5 is $(C_1$-$C_8)$alkyl, $(C_2$-$C_8)$alkenyl, $(C_2$-$C_8)$alkynyl, benzyl, aryl, or $(C_2$-$C_5)$heteroaryl;

each occurrence of R^6 is independently H, $(C_1$-$C_8)$alkyl, $(C_2$-$C_8)$alkenyl, $(C_2$-$C_8)$alkynyl, benzyl, aryl, $(C_2$-$C_5)$heteroaryl, or $(C_0$-$C_8)$alkyl-$C(O)O$-R^5 or the R^6 groups join to form a heterocycloalkyl group;

n is 0 or 1; and

* represents a chiral-carbon center, a cyano or carboxy derivative of a substituted styrene, a 1-oxo-2-(2,6-dioxo-3 fluoropiperidin-3yl) isoindoline, a 1,3-dioxo-2-(2,6-dioxo-3-fluoropiperidine-3-yl) isoindoline, and a tetra substituted 2-(2,6-dioxopiperdin-3-yl)-1-oxoisoindoline.

In a second aspect, there is provided a use of prophylactically effective amount of an immunomodulatory compound as defined in the first aspect, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof in the manufacture of a medicament for managing a myelodysplastic syndrome to a patient in need thereof.

In a third aspect, there is provided a use of therapeutically or prophylactically effective amount of an immunomodulatory compound as defined in the first aspect, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof, and a therapeutically or prophylactically effective amount of at least one second active ingredient in the manufacture of a medicament for treating, preventing or managing a myelysplastic syndrome to a patient in need thereof.

In a fourth aspect, there is provided a use of therapeutically or prophylactically effective amount of an immunomodulatory compound as defined in the first aspect, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof, in the manufacture of a medicament for treating, preventing or managing myelodysplastic syndromes during, before or after transplanting umbilical cord blood, placental blood, peripheral blood stem cell, hematopoietic stem cell preparation or bone marrow in the patient in need thereof.

In a fifth aspect, there is provided a use of amount of the second active ingredient and a therapeutically or prophylactically effective amount of an immunomodulatory compound as defined in the first, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof in the manufacture of a medicament for reducing or avoiding an adverse effect associated with the administration of a second active ingredient in a patient suffering from a myelodysplastic syndrome.

In a sixth aspect, there is provided a pharmacoutical composition for treating, preventing or managing myelodysplastic syndromes comprising an immunomodulatory compound as defined in the first aspect, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof in an amount effective to treat, prevent or manage myelodysplastic syndromes, and a carrier.

In a seventh aspect, there is provided a pharmaceutical composition for treating, preventing or managing myelodysplastic syndromes comprising an immunomodulatory compound as defined in the first aspect, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof, and a second active ingredient.

In an eighth aspect, there is provided a single unit dosage form for treating, preventing or managing myelodysplastic syndromes comprising an immunornodulatory compound as defined in the first aspect, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof, and a second active ingredient capable of improving blood cell production.

In a ninth aspect, there is provided a single unit dosage form for treating, preventing or managing myelodysplastic syndromes comprising an immunomodulatory compound as defined in the first aspect, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof, and a second active ingredient, wherein the

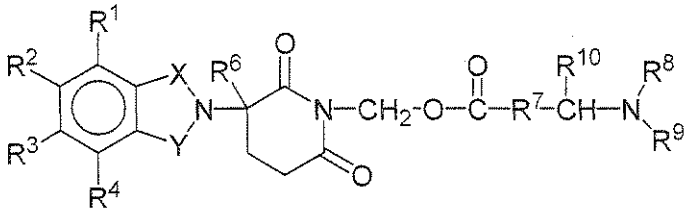
second active ingredient is a cytokine, hematopoietic growth factor, anti-cancer agent, antibiotic, proteasome inhibitor, or immunosuppressive agent.

In a tenth aspect, there is provided a single unit dosage form comprising an immunomodulatory compound as defined in the first aspect, or a pharmacoutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof, and etanercept, imatinib, anti-TNF-ot antibodies, infliximab, G-CSF, GM-CSF, EPO, topotecan, pentoxifylline, ciprofloxacin, irinotecan, vinblastine, dexamethasone, IL2, IL8, IL18, Ara-C, vinorelbine, isotretinoin, 13-cis-retinoic acid, or a pharmacologically active mutant or derivative thereof, or a combination thereof.

In an eleventh aspect, there is provided a kit comprising: a pharmaceutical composition for treating myelopdysplastic syndrome comprising an immunomodulatory compound as defined in the first aspect, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof; and umbilical cord blood, placental blood, peripheral blood stem cell, hematopoietic stem cell preparation or bone marrow.

In a twelfth aspect, there is provided a kit comprising: a pharmaceutical composition comprising an immunomodulatory compound as defined in the first aspect, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof; a pharmaceutical composition comprising a second active ingredient selected from the group consisting of a cytokine, hematopoietic growth factor, anti-cancer agent, antibiotic, proteasome inhibitor, and immunosuppressive agent; and umbilical cord blood, placental blood, peripheral blood stem cell, hematopoletic stem cell preparation or bone marrow.

In a thirteenth aspect, there is provided the use of an immunomodulatory compound, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof for manufacturing a medicament for treating, preventing or managing a myolodysplastic syndrome, wherein said immunomodulatory compound is of formula:



wherein:

one of X and Y is C=O and the other of X and Y is C=O or CH_2;

(i) each of R^1, R^2, R^3, or R^4, independently of the others, is halo, alkyl of 1 to 4 carbon atoms, or alkoxy of 1 to 4 carbon atoms or (ii) one of R^1, R^2, R^3, and R^4 is -NHR^5 and the remaining of R^1, R^2, R^3, and R^4 are hydrogen;

R^5 is hydrogen or alkyl of 1 to 8 carbon atoms;

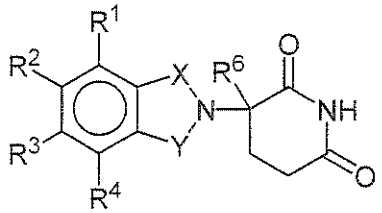
R^6 is hydrogen, alkyl of 1 to 8 carbon atoms, benzo, chloro, or fluoro;

R^7 is m-phenylene or p-phenylene or -(C_nH_{2n})- in which n has a value of 0 to 4;

each of R^8 and R^9 taken independently of the other is hydrogen or alkyl of 1 to 8 carbon atoms, or R^8 and R^9 taken together are tetramethylene, pentamethylene, hexamethylene, or -CH_2CH_2 $X^1CH_2CH_2$- in which X^1 is -O-, -S-, or -NH-;

R^{10} is hydrogen, alkyl of to 8 carbon atoms, or phenyl.

In a fourteenth aspect, there is provided the use of an immunomodulatory compound, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof for manufacturing a medicament for treating, preventing or managing a myolodysplastic syndrome, wherein said immunomodulatory compound is of formula:



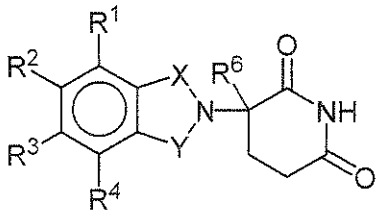
in which

one of X and Y is C=O and the other of X and Y is C=O or CH_2;

each of R^1, R^2, R^3, and R^4, independently of the others, is halo, alkyl of 1 to 4 carbon atoms, or alkoxy of 1 to 4 carbon atoms or (ii) one of R^1, R^2, R^3, and R^4 is nitro or protected amino and the remaining of R^1, R^2, R^3, and R^4 are hydrogen; and

R^6 is hydrogen, alkyl of 1 to 8 carbon atoms, benzo, chloro, or fluoro.

In a fifteenth aspect, there is provided the use of an immunomodulatory compound, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof for manufacturing a medicament for treating, preventing or managing a myolodysplastic syndrome, wherein said immunomodulatory compound is of formula:



in which:

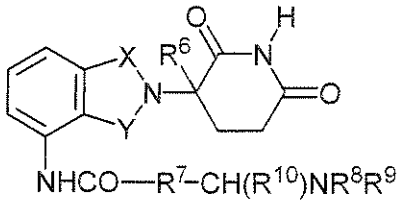
one of X and Y is C=O and the other of X and Y is C=O or CH_2;

(i) each of R^1, R^2, R^3, and R^4, independently of the others, is halo, alkyl of 1 to 4 carbon atoms, or alkoxy of 1 to 4 carbon atoms or (ii) one of R^1, R^2, R^3, and R^4 is -NHR^5 and the remaining of R^1, R^2, R^3, and R^4 are hydrogen;

R^5 is hydrogen, alkyl of 1 to 8 carbon atoms, or CO-R^7-$CH(R^{10})NR^8R^9$ in which each of R^7, R^8, R^9, and R^{10} is as herein defined; and

R^6 is alkyl of 1 to 8 carbon atoms, benzo, chloro, or fluoro.

In a sixteenth aspect, there is provided the use of an immunomodulatory compound, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof for manufacturing a medicament for treating, preventing or managing a myolodysplastic syndrome, wherein said immunomodulatory compound is of formula:



in which:

one of X and Y is C=O and the other of X and Y is C=O or CH_2;

R^6 is hydrogen, alkyl of 1 to 8 carbon atoms, benzyl, chloro, or fluoro;

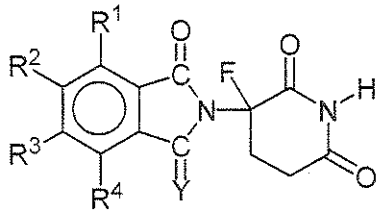
R^7 is m-phenylene, p-phenylene or -(C_nH_{2n})- in which n has a value of 0 to 4;

each of R^8 and R^9 taken independently of the other is hydrogen or alkyl of 1 to 8 carbon atoms, or R^8 and R^9 taken together are tetramethylene, pentamethylene, hexamethylene, or -$CH_2CH_2X^1CH_2CH_2$- in which X^1 is -O-, -S- or -NH-; and

R^{10} is hydrogen, alkyl of 1 to 8 carbon atoms, or phenyl.

In a seventeenth aspect, there is provided the use of an immunomodulatory compound, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof for manufacturing a medicament for treating, preventing or managing a

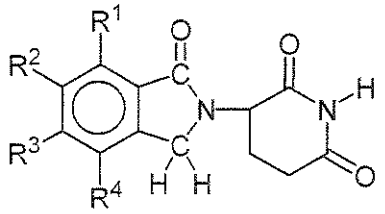
myolodysplastic syndrome, wherein said immunomodulatory compound is of formula:



wherein Y is oxygen or H^2 and

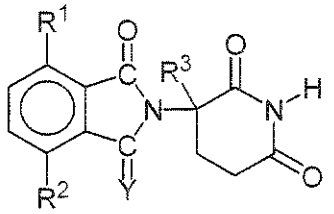
each of R^1, R^2, R^3, and R^4, independently of the others, is hydrogen, halo, alkyl of 1 to 4 carbon atoms, alkoxy of 1 to 4 carbon atoms, or amino.

In an eighteenth aspect, there is provided the use of an immunomodulatory compound, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof for manufacturing a medicament for treating, preventing or managing a myolodysplastic syndrome, wherein said immunomodulatory compound is of formula:



wherein each of R^1, R^2, R^3, and R^4, independently of the others, is halo, alkyl of 1 to 4 carbon atoms, or alkoxy of 1 to 4 carbon atoms.

In a nineteenth aspect, there is provided the use of an immunomodulatory compound, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof for manufacturing a medicament for treating, preventing or managing a myolodysplastic syndrome, wherein said immunomodulatory compound is of formula:



in which

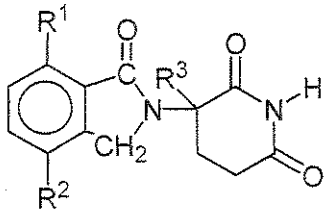
Y is oxygen or H_2,

a first of R^1 and R^2 is halo, alkyl, alkoxy, alkylamino, dialkylamino, cyano, or

carbamoyl, the second of R^1 and R^2, independently of the first, is hydrogen, halo, alkyl, alkoxy, alkylamino, dialkylamino, cyano, or carbamoyl, and

R^3 is hydrogen, alkyl, or benzyl.

In a twentieth aspect, there is provided the use of an immunomodulatory compound, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof for manufacturing a medicament for treating, preventing or managing a myolodysplastic syndrome, wherein said immunomodulatory compound is of formula:

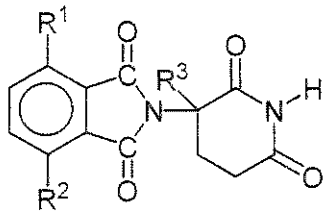


wherein

a first of R^1 and R^2 is halo, alkyl of from 1 to 4 carbon atoms, alkoxy of from 1 to 4 carbon atoms, dialkylamino in which each alkyl is of from 1 to 4 carbon atoms, cyano, or carbamoyl; the second of R^1 and R^2, independently of the first, is hydrogen, halo, alkyl of from 1 to 4 carbon atoms, alkoxy of from 1 to 4 carbon atoms, alkylamino in which alkyl is of from 1 to 4 carbon atoms, dialkylamino in which each alkyl is of from 1 to 4 carbon atoms, cyano, or carbamoyl, and

R^3 is hydrogen, alkyl of from 1 to 4 carbon atoms, or benzyl.

In a twenty-first aspect, there is provided the use of an immunomodulatory compound, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof for manufacturing a medicament for treating, preventing or managing a myolodysplastic syndrome, wherein said immunomodulatory compound is of formula:

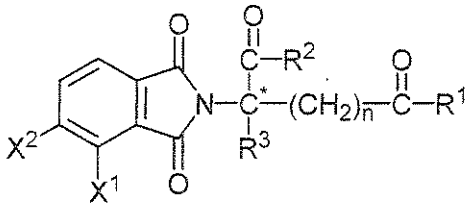


wherein

a first of R^1 and R^2 is halo, alkyl of from 1 to 4 carbon atoms, alkoxy of from 1 to 4 carbon atoms, dialkylamino in which each alkyl is of from 1 to 4 carbon atoms, cyano, or carbamoyl, the second of R^1 and R^2, independently of the first, is hydrogen, halo, alkyl of from 1 to 4 carbon atoms, alkoxy of from 1 to 4 carbon atoms, alkylamino in which alkyl is of from 1 to 4 carbon atoms, dialkylamino in which each alkyl is of from 1 to 4 carbon atoms, cyano, or carbamoyl, and

R^3 is hydrogen, alkyl of from 1 to 4 carbon atoms, or benzyl.

In a twenty-second aspect, there is provided the use of an immunomodulatory compound, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof for manufacturing a medicament for treating, preventing or managing a myolodysplastic syndrome, wherein said immunomodulatory compound is of formula:



in which

the carbon atom designated C* constitutes a center of chirality (when n is not zero and R^1 is not the same as R^2);

one of X^1 and X^2 is amino, nitro, alkyl of one to six carbons, or NH-Z, and the other of X^1 or X^2 is hydrogen;

each of R^1 and R^2 independent of the other, is hydroxy or NH-Z;

R^3 is hydrogen, alkyl of one to six carbons, halo, or haloalkyl;

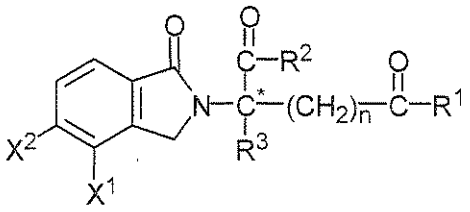
Z is hydrogen, aryl, alkyl of one to six carbons, formyl, or acyl of one to six carbons; and

n has a value of 0, 1, or 2;

provided that if X^1 is amino, and n is 1 or 2, then R^1 and R^2 are not both hydroxy; and the salts thereof.

In a twenty-third aspect, there is provided the use of an immunomodulatory compound, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof for manufacturing a medicament for treating, preventing or managing a

myolodysplastic syndrome, wherein said immunomodulatory compound is of formula:



in which

the carbon atom designated C* constitutes a center of chirality when n is not zero and R^1 is not R^2;

one of X^1 and X^2 is amino, nitro, alkyl of one to six carbons, or NH-Z, and the other of X^1 or X^2 is hydrogen;

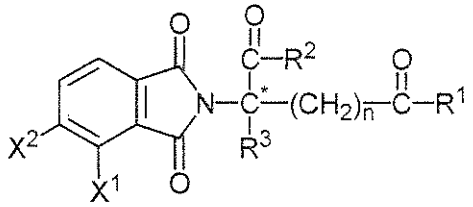
each of R^1 and R^2 independent of the other, is hydroxy or NH-Z;

R^3 is alkyl of one to six carbons, halo, or hydrogen;

Z is hydrogen, aryl or an alkyl or acyl of one to six carbons; and

n has a value of 0, 1, or 2.

In a twenty-fourth aspect, there is provided the use of an immunomodulatory compound, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof for manufacturing a medicament for treating, preventing or managing a myolodysplastic syndrome, wherein said immunomodulatory compound is of formula:



in which

the carbon atom designated C* constitutes a center of chirality when n is not zero and R^1 is not R^2;

one of X^1 and X^2 is amino, nitro, alkyl of one to six carbons, or NH-Z, and the other of X^1or X^2 is hydrogen;

each of R^1 and R^2 independent of the other, is hydroxy or NH-Z;

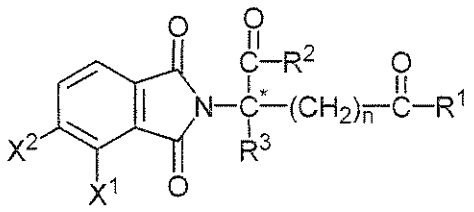
R^3 is alkyl of one to six carbons, halo, or hydrogen;

Z is hydrogen, aryl, or an alkyl or acyl of one to six carbons; and

n has a value of 0, 1, or 2;

and the salts thereof.

In a twenty-fifth aspect, there is provided the use of an immunomodulatory compound, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof for manufacturing a medicament for treating, preventing or managing a myolodysplastic syndrome, wherein said immunomodulatory compound is of formula:



wherein

one of X^1 and X^2 is nitro, or NH-Z, and the other of X^1 or X^2 is hydrogen;

each of R^1 and R^2, independent of the other, is hydroxy or NH-Z;

R^3 is alkyl of one to six carbons, halo, or hydrogen;

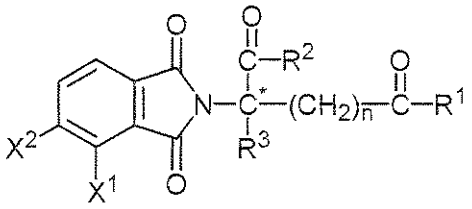
Z is hydrogen, phenyl, an acyl of one to six carbons, or an alkyl of one to six carbons; and

n has a value of 0, 1, or 2;

provided that if one of X^1 and X^2 is nitro, and n is 1 or 2, then R^1 and R^2 are other than hydroxy; and

if $-COR^1$ and $-(CH_2)_nCOR^2$ are different, the carbon atom designated C^* constitutes a center of chirality.

In a twenty-sixth aspect, there is provided the use of an immunomodulatory compound, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof for manufacturing a medicament for treating, preventing or managing a myolodysplastic syndrome, wherein said immunomodulatory compound is of formula:



wherein

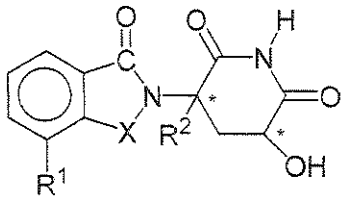
one of X^1 and X^2 is alkyl of one to six carbons;

each of R^1 and R^2, independent of the other, is hydroxy or NH-Z;

R^3 is alkyl of one to six carbons, halo, or hydrogen;

Z is hydrogen, phenyl, an acyl of one to six carbons, or an alkyl of one to six carbons; and
n has a value of 0, 1, or 2; and
if $-COR^1$ and $-(CH_2)_nCOR^2$ are different, the carbon atom designated C^* constitutes a center of chirality.

In a twenty-seventh aspect, there is provided the use of an immunomodulatory compound, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof for manufacturing a medicament for treating, preventing or managing a myolodysplastic syndrome, wherein said immunomodulatory compound is of formula:



wherein:
the carbon atoms designated * constitute centers of chirality;
X is -C(O)- or $-CH_2-$;
R^1 is alkyl of 1 to 8 carbon atoms or $-NHR^3$;
R^2 is hydrogen, alkyl of 1 to 8 carbon atoms, or halogen;and
R^3 is hydrogen, alkyl of 1 to 8 carbon atoms, unsubstituted or substituted with alkoxy of 1 to 8 carbon atoms, halo, amino, or alkylamino of 1 to 4 carbon atoms, cycloalkyl of 3 to 18 carbon atoms, phenyl, unsubstituted or substituted with alkyl of 1 to 8 carbon atoms, alkoxy of 1 to 8 carbon atoms, halo, amino, or alkylamino of 1 to 4 carbon atoms, benzyl, unsubstituted or substituted with alkyl of 1 to 8 carbon atoms, alkoxy of 1 to 8 carbon atoms, halo, amino, or alkylamino of 1 to 4 carbon atoms, or $-COR^4$ in which
R^4 is hydrogen, alkyl of 1 to 8 carbon atoms, unsubstituted or substituted with alkoxy of 1 to 8 carbon atoms, halo, amino, or alkylamino of 1 to 4 carbon atoms, cycloalkyl of 3 to 18 carbon atoms, phenyl, unsubstituted or substituted with alkyl of 1 to 8 carbon atoms, alkoxy of 1 to 8 carbon atoms, halo, amino, or alkylamino of 1 to 4 carbon atoms, or benzyl, unsubstituted or substituted with alkyl of 1 to 8 carbon atoms, alkoxy of 1 to 8 carbon atoms, halo, amino, or alkylamino of 1 to 4 carbon atoms.

In a twenty-eighth aspect, there is provided the use of an immunomodulatory compound, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof for manufacturing a medicament for treating, preventing or managing a

myolodysplastic syndrome, wherein said immunomodulatory compound is selected from the group consisting of:
[2-(2,6-dioxo-piperidin-3-yl)-1,3-dioxo-2,3-dihydro-1H-isoindol-4-ylmethyl]-amide; (2-(2,6-dioxo-piperidin-3-yl)-1,3-dioxo-2,3-dihydro-1*H*-isoindol-4-ylmethyl)-carbamic acid *tert*-butyl ester; 4-(aminomethyl)-2-(2,6-dioxo(3-piperidyl))-isoindoline-1,3-dione; *N*-(2-(2,6-dioxo-piperidin-3-yl)-1,3-dioxo-2,3-dihydro-1*H*-isoindol-4-ylmethyl)-acetamide; *N*-{(2-(2,6-dioxo(3-piperidyl)-1,3-dioxoisoindolin-4-yl)methyl}cyclopropyl-carboxamide; 2-chloro-*N*-{(2-(2,6-dioxo(3-piperidyl))-1,3-dioxoisoindolin-4-yl)methyl}acetamide; *N*-(2-(2,6-dioxo(3-piperidyl))-1,3-dioxoisoindolin-4-yl)-3-pyridylcarboxamide; 3-{1-oxo-4-(benzylamino)isoindolin-2-yl}piperidine-2,6-dione; 2-(2,6-dioxo(3-piperidyl))-4-(benzylamino)isoindoline-1,3-dione; *N*-{(2-(2,6-dioxo(3-piperidyl))-1,3-dioxoisoindolin-4-yl)methyl}propanamide; *N*-{(2-(2,6-dioxo(3-piperidyl))-1,3-dioxoisoindolin-4-yl)methyl}-3-pyridylcarboxamide; *N*-{(2-(2,6-dioxo(3-piperidyl))-1,3-dioxoisoindolin-4-yl)methyl}heptanamide; *N*-{(2-(2,6-dioxo(3-piperidyl))-1,3-dioxoisoindolin-4-yl)methyl}-2-furylcarboxamide; {N-(2-(2,6-dioxo(3-piperidyl))-1,3-dioxoisoindolin-4-yl)carbamoyl}methyl acetate; *N*-(2-(2,6-dioxo(3-piperidyl))-1,3-dioxoisoindolin-4-yl)pentanamide; *N*-(2-(2,6-dioxo(3-piperidyl))-1,3-dioxoisoindolin-4-yl)-2-thienylcarboxamide; N-{[2-(2,6-dioxo(3-piperidyl))-1,3-dioxoisoindolin-4-yl] methyl}(butylamino)carboxamide; N-{[2-(2,6-dioxo(3-piperidyl))-1,3-dioxoisoindolin-4-yl] methyl}(octylamino)carboxamide; and N-{[2-(2,6-dioxo(3-piperidyl))-1,3-dioxoisoindolin-4-yl] methyl}(benzylamino)carboxamide; 1-oxo-2-(2,6-dioxopiperidin-3-yl) isoindolines or 1,3-dioxo-2-(2,6-dioxopiperidin-3-yl) isoindolines; isoindoline-1-one substituted in the 2-position with 2,6-dioxo-3-hydroxypiperidin-5-yl; or isoindoline-1,3-dione substituted in the 2-position with 2,6-dioxo-3-hydroxypiperidin-5-yl.

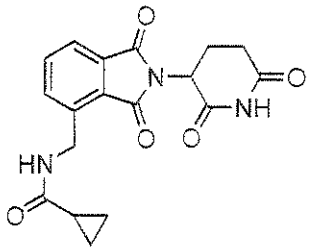
In a twenty-ninth aspect, there is provided the use of an immunomodulatory compound, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof and a second active ingredient for manufacturing a medicament for reducing or avoiding an adverse effect associated with the administration of the second active ingredient in a patient in need of treating, preventing or managing a myelodysplastic syndrome, wherein said immunomodulatory compound is the compound as defined above.

In a thirtieth aspect, there is provided a pharmaceutical composition comprising an immunomodulatory compound, or a pharmaceutically acceptable salt, solvate, hydrate,

stereoisomer, clathrate, or prodrug thereof in an amount effective to treat, prevent or manage myelodysplastic syndromes, and a carrier, wherein said immunomodulatory compound is the compound as defined above.

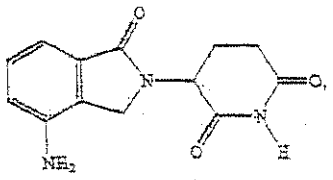
In a thirty-first aspect, there is provided a kit comprising: a pharmaceutical composition comprising an immunomodulatory compound, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof, wherein said immunomodulatory compound is the compound as defined above; and umbilical cord blood, placental blood, peripheral blood stem cell, hematopoietic stem cell preparation or bone marrow.

In a thirty-second aspect, there is provided the use of N-{[2-(2,6-dioxo(3-piperidyl)-1,3-dioxoisoindolin-4-yl]methyl}cyclopropyl-carboxamide having the formula:



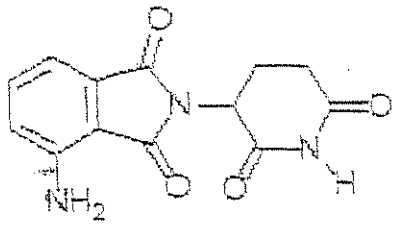
or a pharmaceutically acceptable salt, solvate or stereoisomer thereof in manufacturing a medicament for treating a myelodysplastic syndrome, wherein said medicament is in a daily dosage form comprising said compound in an amount of about 5mg to about 100 mg.

In a thirty-third aspect, there is provided the use of from about 0.1 to about 10 mg per of day of 3-(4-amino-1-oxo-1,3-dihydro-isoindol-2-yl)-piperidine-2,6-dione having the formula:



or a pharmaceutically acceptable salt, solvate or stereoisomer thereof for the manufacturing of a medicament for treatment of a myelodysplastic syndrome.

In a thirty-fourth aspect, there is provided the use of from about 0.1 to about 10 mg per of day of 4-(amino)-2-2,6-dioxo (3-piperidy;))-isoindoline-1,3-dione having the formula:



or a pharmaceutically acceptable salt, solvate or steroisomer thereof for the manufacturing of a medicament for treatment of a myelodysplastic syndrome.

4. DETAILED DESCRIPTION OF THE INVENTION

A first embodiment of the invention encompasses methods of treating or preventing MDS which comprise administering to a patient in need of such treatment or prevention a therapeutically or prophylactically effective amount of an immunomodulatory compound, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof. The embodiment encompasses the treatment, prevention or management of specific sub-types of MDS such as refractory anemia, refractory anemia with ringed sideroblasts, refractory anemia with excess blasts, refractory anemia with excess blasts in transformation and chronic myelomonocytic leukemia.

As used herein, the term "myelodysplastic syndromes" or "MDS" means hematopoietic stem cell disorders characterized by one or more of the following: ineffective blood cell production, progressive cytopenias, risk of progression to acute leukemia or cellular marrow with impaired morphology and maturation (dysmyelopoiesis). The term "myelodysplastic syndromes" or "MDS" unless otherwise noted includes: refractory anemia, refractory anemia with ringed sideroblasts, refractory anemia with excess blasts, refractory anemia with excess blasts in transformation and chronic myelomonocytic leukemia.

Another embodiment of the invention encompasses methods of managing MDS which comprises administering to a patient in need of such management a prophylactically effective amount of an immunomodulatory compound, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof.

Another embodiment of the invention encompasses a pharmaceutical composition comprising an immunomodulatory compound, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof.

Also encompassed by the invention are single unit dosage forms comprising an immunomodulatory compound, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof.

Another embodiment of the invention encompasses a kit comprising: a pharmaceutical composition comprising an immunomodulatory compound, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof and a second active or dexamethasone or instructions for use. The invention further encompasses kits comprising single unit dosage forms.

One embodiment of the invention encompasses a method of treating, preventing and/or managing MDS, which comprises administering to a patient in need of such treatment, prevention and/or management a therapeutically or prophylactically effective

amount of an immunomodulatory compound, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof, and a therapeutically or prophylactically effective amount of a second active agent.

The second active agent is preferably a hematopoietic growth factor, a cytokine, an anti-cancer agent, an antibiotic, an anti-fungal, an anti-inflammatory, an immunosuppressive agent such as a cyclosporin, conventional therapy for MDS, or other chemotherapeutic agent found for example in the Physician's Desk Reference 2002. Preferred anti-cancer or cancer chemotherapeutics are apoptosis inducing agents, topoisomerase inhibitors, anti-angiogenesis compounds, microtubule stabilizing agents, alkylating agents and other known conventional cancer chemotherapy. Most preferred second active agents are those capable of affecting or improving blood production. Second active agents can be large molecules (*e.g.*, proteins) or small molecules (*e.g.*, synthetic inorganic, organometallic, or organic molecules). The examples of specific second active agent include, but are not limited to, etanercept (Enbrel®), imatinib (Glivec®), anti-TNF-α antibodies, infliximab (Remicade®), G-CSF, GM-CSF, EPO, topotecan, irinotecan, pentoxifylline, ciprofloxacin, dexamethasone, IL2, IL8, IL18, Ara-C, vinorelbine, vinblastine, isotretinoin, and13-cis-retinoic acid. This invention also encompasses the use of native, naturally occurring, and recombinant proteins. The invention further encompasses mutants and derivatives (*e.g.*, modified forms) of naturally occurring proteins that exhibit, *in vivo*, at least some of the pharmacological activity of the proteins upon which they are based. Examples of mutants include, but are not limited to, proteins that have one or more amino acid residues that differ from the corresponding residues in the naturally occurring forms of the proteins. Also encompassed by the term "mutants" are proteins that lack carbohydrate moieties normally present in their naturally occurring forms (*e.g.*, nonglycosylated forms). Examples of derivatives include, but are not limited to, pegylated derivatives and fusion proteins, such as proteins formed by fusing IgG1 or IgG3 to the protein or active portion of the protein of interest. *See, e.g.*, Penichet, M.L. and Morrison, S.L., *J. Immunol. Methods* 248:91-101 (2001). Vaccines that cause the secretion of proteins disclosed herein as well as pharmacologically active mutants, derivatives, and fusion thereof are also encompassed by the invention.

Without being limited by theory, it is believed that certain immunomodulatory compounds and proteins can act in complementary or synergistic ways in the treatment or management of MDS. It is also believed that certain proteins may reduce or eliminate particular adverse effects associated with some immunomodulatory compounds, thereby allowing the administration of larger amounts of an immunomodulatory compound to

patients and/or increasing patient compliance. It is further believed that some immunomodulatory compounds may reduce or eliminate particular adverse effects associated with some protein-based MDS therapies, thereby allowing the administration of larger amounts of protein to patients and/or increasing patient compliance.

Another embodiment of the invention encompasses a method of reversing, reducing or avoiding an adverse effect associated with the administration of a chemotherapeutics or therapeutics used to treat cancer or MDS in a patient suffering from MDS, which comprises administering to a patient in need thereof a therapeutically or prophylactically effective amount of an immunomodulatory compound, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof.

As inevitable leukemic transformation develops in certain stages of MDS, transplantation of peripheral blood stem cells, hematopoietic stem cell preparation or bone marrow may be necessary. It is believed that the combined use of an immunomodulatory compound and transplantation of stem cells in a patient suffering from MDS provides a unique and unexpected synergism. In particular, without being limited by theory, it is believed that an immunomodulatory compound exhibits immunomodulatory activity that may provide additive or synergistic effects when given concurrently with transplantation therapy. Immunomodulatory compounds can work in combination with transplantation therapy reducing complications associated with the invasive procedure of transplantation and risk of related Graft Versus Host Disease (GVHD). Therefore, this invention encompasses a method of treating, preventing and/or managing MDS, which comprises administering to a patient (*e.g.*, a human) an immunomodulatory compound, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof, before, during, or after transplantation therapy.

The invention also encompasses pharmaceutical compositions, single unit dosage forms, and kits which comprise one or more immunomodulatory compounds, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof, a second active ingredient, and/or blood or cells for transplantation therapy. For example, the kit may contain one or more compounds of the invention, stem cells for transplantation and an immunosuppressive agent, antibiotic or other drug, each of which is to be used to treat the MDS patient.

4.1. IMMUNOMODULATORY COMPOUNDS

Compounds of the invention can either be commercially purchased or prepared according to the methods described in the patents or patent publications disclosed herein.

Further, optically pure compositions can be asymmetrically synthesized or resolved using known resolving agents or chiral columns as well as other standard synthetic organic chemistry techniques. Compounds used in the invention may include immunomodulatory compounds that are racemic, stereomerically enriched or stereomerically pure, and pharmaceutically acceptable salts, solvates, stereoisomers, clathrates, and prodrugs thereof.

As used herein, unless otherwise indicated, the term "solvates" includes hydrates of the compounds of the invention.

Preferred compounds used in the invention are small organic molecules having a molecular weight less than about 1,000 g/mol, and are not proteins, peptides, oligonucleotides, oligosaccharides or other macromolecules.

As used herein and unless otherwise indicated, the terms "immunomodulatory compounds" and "IMiDs™" (Celgene Corporation) encompasses small organic molecules that markedly inhibit TNF-α, LPS induced monocyte IL1ß and IL12, and partially inhibit IL6 production. Specific immunomodulatory compounds are discussed below.

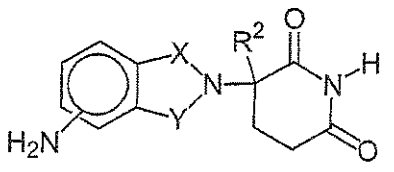
TNF-α is an inflammatory cytokine produced by macrophages and monocytes during acute inflammation. TNF-α is responsible for a diverse range of signaling events within cells. TNF-α may play a pathological role in cancer. Without being limited by theory, one of the biological effects exerted by the immunomodulatory compounds of the invention is the reduction of synthesis of TNF-α. Immunomodulatory compounds of the invention enhance the degradation of TNF-α mRNA.

Further, without being limited by theory, immunomodulatory compounds used in the invention may also be potent co-stimulators of T cells and increase cell proliferation dramatically in a dose dependent manner. Immunomodulatory compounds of the invention may also have a greater co-stimulatory effect on the CD8+ T cell subset than on the CD4+ T cell subset. In addition, the compounds preferably have anti-inflammatory properties, and efficiently co-stimulate T cells.

Specific examples of immunomodulatory compounds, include, but are not limited to, cyano and carboxy derivatives of substituted styrenes such as those disclosed in U.S. patent no. 5,929,117; 1-oxo-2-(2,6-dioxo-3-fluoropiperidin-3yl) isoindolines and 1,3-dioxo-2-(2,6-dioxo-3-fluoropiperidine-3-yl) isoindolines such as those described in U.S. patent nos. 5,874,448 and 5,955,476; the tetra substituted 2-(2,6-dioxopiperdin-3-yl)-1-oxoisoindolines described in U.S. patent no. 5,798,368; 1-oxo and 1,3-dioxo-2-(2,6-dioxopiperidin-3-yl) isoindolines (*e.g.*, 4-methyl derivatives of thalidomide), including, but not limited to, those disclosed in U.S. patent nos. 5,635,517, 6,476,052, 6,555,554, and 6,403,613; 1-oxo and 1,3-dioxoisoindolines substituted in the 4- or 5-position of the

indoline ring (*e.g.*, 4-(4-amino-1,3-dioxoisoindoline-2-yl)-4-carbamoylbutanoic acid) described in U.S. patent no. 6,380,239; isoindoline-1-one and isoindoline-1,3-dione substituted in the 2-position with 2,6-dioxo-3-hydroxypiperidin-5-yl (*e.g.*, 2-(2,6-dioxo-3-hydroxy-5-fluoropiperidin-5-yl)-4-aminoisoindolin-1-one) described in U.S. patent no. 6,458,810; a class of non-polypeptide cyclic amides disclosed in U.S. patent nos. 5,698,579 and 5,877,200; aminothalidomide, as well as analogs, hydrolysis products, metabolites, derivatives and precursors of aminothalidomide, and substituted 2-(2,6-dioxopiperidin-3-yl) phthalimides and substituted 2-(2,6-dioxopiperidin-3-yl)-1-oxoisoindoles such as those described in U.S. patent nos. 6,281,230 and 6,316,471; and isoindole-imide compounds such as those described in U.S. patent application no. 09/972,487 filed on October 5, 2001, U.S. patent application no. 10/032,286 filed on December 21, 2001, and International Application No. PCT/US01/50401 (International Publication No. WO 02/059106). The entireties of each of the patents and patent applications identified herein are incorporated herein by reference. Immunomodulatory compounds do not include thalidomide.

Other specific immunomodulatory compounds of the invention include, but are not limited to, 1-oxo-and 1,3 dioxo-2-(2,6-dioxopiperidin-3-yl) isoindolines substituted with amino in the benzo ring as described in U.S. Patent no. 5,635,517 which is incorporated herein by reference. These compounds have the structure I:



I

in which one of X and Y is C=O, the other of X and Y is C=O or CH_2, and R^2 is hydrogen or lower alkyl, in particular methyl. Specific immunomodulatory compounds include, but are not limited to:

1-oxo-2-(2,6-dioxopiperidin-3-yl)-4-aminoisoindoline;

1-oxo-2-(2,6-dioxopiperidin-3-yl)-5-aminoisoindoline;

1-oxo-2-(2,6-dioxopiperidin-3-yl)-6-aminoisoindoline;

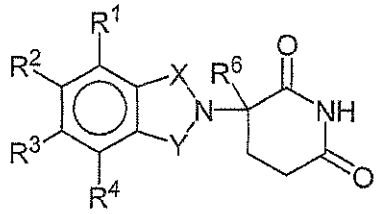
1-oxo-2-(2,6-dioxopiperidin-3-yl)-7-aminoisoindoline;

1,3-dioxo-2-(2,6-dioxopiperidin-3-yl)-4-aminoisoindoline; and

1,3-dioxo-2-(2,6-dioxopiperidin-3-yl)-5-aminoisoindoline.

Other specific immunomodulatory compounds of the invention belong to a class of substituted 2-(2,6-dioxopiperidin-3-yl) phthalimides and substituted 2-(2,6-dioxopiperidin-3-yl)-1-oxoisoindoles, such as those described in U.S. patent nos. 6,281,230; 6,316,471; 6,335,349; and 6,476,052, and International Patent Application No. PCT/US97/13375

(International Publication No. WO 98/03502), each of which is incorporated herein by reference. Representative compounds are of formula:



in which:

one of X and Y is C=O and the other of X and Y is C=O or CH_2;

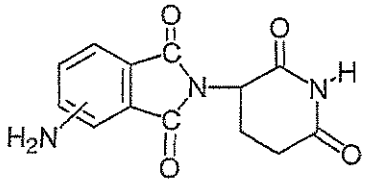
(i) each of R^1, R^2, R^3, and R^4, independently of the others, is halo, alkyl of 1 to 4 carbon atoms, or alkoxy of 1 to 4 carbon atoms or (ii) one of R^1, R^2, R^3, and R^4 is $-NHR^5$ and the remaining of R^1, R^2, R^3, and R^4 are hydrogen;

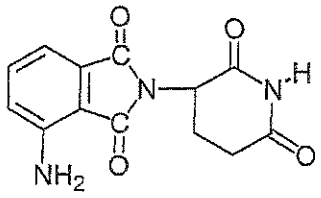
R^5 is hydrogen or alkyl of 1 to 8 carbon atoms;

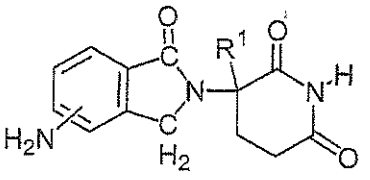
R^6 is hydrogen, alkyl of 1 to 8 carbon atoms, benzyl, or halo;

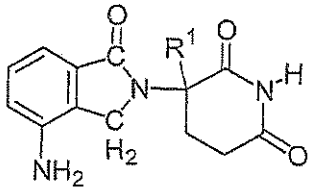
provided that R^6 is other than hydrogen if X and Y are C=O and (i) each of R^1, R^2, R^3, and R^4 is fluoro or (ii) one of R^1, R^2, R^3, or R^4 is amino.

Compounds representative of this class are of the formulas:



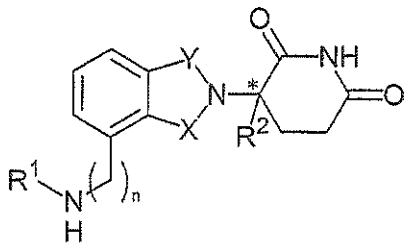






wherein R^1 is hydrogen or methyl. In a separate embodiment, the invention encompasses the use of enantiomerically pure forms (*e.g.* optically pure (R) or (S) enantiomers) of these compounds.

Still other specific immunomodulatory compounds of the invention belong to a class of isoindole-imides disclosed in U.S. Patent Application Publication Nos. US 2003/0096841 and US 2003/0045552, and International Application No. PCT/US01/50401 (International Publication No. WO 02/059106), each of which are incorporated herein by reference. Representative compounds are of formula II:



II

and pharmaceutically acceptable salts, hydrates, solvates, clathrates, enantiomers, diastereomers, racemates, and mixtures of stereoisomers thereof, wherein:

one of X and Y is C=O and the other is CH_2 or C=O;

R^1 is H, $(C_1–C_8)$alkyl, $(C_3–C_7)$cycloalkyl, $(C_2–C_8)$alkenyl, $(C_2$-$C_8)$alkynyl, benzyl, aryl, $(C_0$-$C_4)$alkyl–$(C_1$-$C_6)$heterocycloalkyl, $(C_0$-$C_4)$alkyl–$(C_2$-$C_5)$heteroaryl, $C(O)R^3$, $C(S)R^3$, $C(O)OR^4$, $(C_1$-$C_8)$alkyl–$N(R^6)_2$, $(C_1$-$C_8)$alkyl–OR^5, $(C_1$-$C_8)$alkyl–$C(O)OR^5$, $C(O)NHR^3$, $C(S)NHR^3$, $C(O)NR^3R^{3'}$, $C(S)NR^3R^{3'}$ or $(C_1$-$C_8)$alkyl–$O(CO)R^5$;

R^2 is H, F, benzyl, $(C_1$-$C_8)$alkyl, $(C_2$-$C_8)$alkenyl, or $(C_2$-$C_8)$alkynyl;

R^3 and $R^{3'}$ are independently $(C_1$-$C_8)$alkyl, $(C_3$-$C_7)$cycloalkyl, $(C_2$-$C_8)$alkenyl, $(C_2$-$C_8)$alkynyl, benzyl, aryl, $(C_0$-$C_4)$alkyl–$(C_1$-$C_6)$heterocycloalkyl, $(C_0$-$C_4)$alkyl–$(C_2$-$C_5)$heteroaryl, $(C_0$-$C_8)$alkyl–$N(R^6)_2$, $(C_1$-$C_8)$alkyl–OR^5, $(C_1$-$C_8)$alkyl–$C(O)OR^5$, $(C_1$-$C_8)$alkyl–$O(CO)R^5$, or $C(O)OR^5$;

R^4 is $(C_1$-$C_8)$alkyl, $(C_2$-$C_8)$alkenyl, $(C_2$-$C_8)$alkynyl, $(C_1$-$C_4)$alkyl–OR^5, benzyl, aryl, $(C_0$-$C_4)$alkyl–$(C_1$-$C_6)$heterocycloalkyl, or $(C_0$-$C_4)$alkyl–$(C_2$-$C_5)$heteroaryl;

R^5 is $(C_1$-$C_8)$alkyl, $(C_2$-$C_8)$alkenyl, $(C_2$-$C_8)$alkynyl, benzyl, aryl, or $(C_2$-$C_5)$heteroaryl;

each occurrence of R^6 is independently H, $(C_1$-$C_8)$alkyl, $(C_2$-$C_8)$alkenyl, $(C_2$-$C_8)$alkynyl, benzyl, aryl, $(C_2$-$C_5)$heteroaryl, or $(C_0$-$C_8)$alkyl–$C(O)O–R^5$ or the R^6 groups can join to form a heterocycloalkyl group;

n is 0 or 1; and

* represents a chiral-carbon center.

In specific compounds of formula II, when n is 0 then R^1 is (C_3-C_7)cycloalkyl, (C_2-C_8)alkenyl, (C_2-C_8)alkynyl, benzyl, aryl, (C_0-C_4)alkyl–(C_1-C_6)heterocycloalkyl, (C_0-C_4)alkyl-(C_2-C_5)heteroaryl, $C(O)R^3$, $C(O)OR^4$, (C_1-C_8)alkyl–$N(R^6)_2$, (C_1-C_8)alkyl–OR^5, (C_1-C_8)alkyl–$C(O)OR^5$, $C(S)NHR^3$, or (C_1-C_8)alkyl–$O(CO)R^5$;

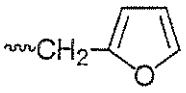
R^2 is H or (C_1-C_8)alkyl; and

R^3 is (C_1-C_8)alkyl, (C_3-C_7)cycloalkyl, (C_2-C_8)alkenyl, (C_2-C_8)alkynyl, benzyl, aryl, (C_0-C_4)alkyl–(C_1 –C_6)heterocycloalkyl, (C_0-C_4)alkyl–(C_2-C_5)heteroaryl, (C_5-C_8)alkyl–$N(R^6)_2$; (C_0-C_8)alkyl–NH–C(O)O–R^5; (C_1-C_8)alkyl–OR^5, (C_1-C_8)alkyl–$C(O)OR^5$, (C_1-C_8)alkyl–$O(CO)R^5$, or $C(O)OR^5$; and the other variables have the same definitions.

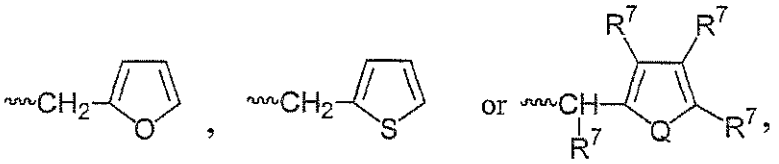
In other specific compounds of formula II, R^2 is H or (C_1-C_4)alkyl.

In other specific compounds of formula II, R^1 is (C_1-C_8)alkyl or benzyl.

In other specific compounds of formula II, R^1 is H, (C_1-C_8)alkyl, benzyl, CH_2OCH_3, $CH_2CH_2OCH_3$, or



In another embodiment of the compounds of formula II, R^1 is



wherein Q is O or S, and each occurrence of R^7 is independently H,(C_1-C_8)alkyl, (C_3-C_7)cycloalkyl, (C_2-C_8)alkenyl, (C_2-C_8)alkynyl, benzyl, aryl, halogen, (C_0-C_4)alkyl–(C_1-C_6)heterocycloalkyl, (C_0-C_4)alkyl–(C_2-C_5)heteroaryl, (C_0-C_8)alkyl–$N(R^6)_2$, (C_1-C_8)alkyl–OR^5, (C_1-C_8)alkyl–$C(O)OR^5$, (C_1-C_8)alkyl–$O(CO)R^5$, or $C(O)OR^5$, or adjacent occurrences of R^7 can be taken together to form a bicyclic alkyl or aryl ring.

In other specific compounds of formula II, R^1 is $C(O)R^3$.

In other specific compounds of formula II, R^3 is (C_0-C_4)alkyl–(C_2-C_5)heteroaryl, (C_1-C_8)alkyl, aryl, or (C_0-C_4)alkyl–OR^5.

In other specific compounds of formula II, heteroaryl is pyridyl, furyl, or thienyl.

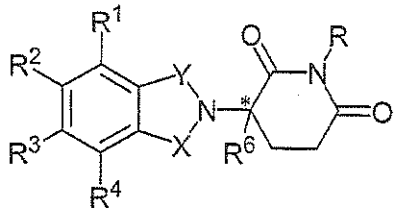
In other specific compounds of formula II, R^1 is $C(O)OR^4$.

In other specific compounds of formula II, the H of C(O)NHC(O) can be replaced with (C_1-C_4)alkyl, aryl, or benzyl.

Further examples of the compounds in this class include, but are not limited to: [2-(2,6-dioxo-piperidin-3-yl)-1,3-dioxo-2,3-dihydro-1H-isoindol-4-ylmethyl]-amide; (2-(2,6-dioxo-piperidin-3-yl)-1,3-dioxo-2,3-dihydro-1*H*-isoindol-4-ylmethyl)-carbamic acid *tert*-

butyl ester; 4-(aminomethyl)-2-(2,6-dioxo(3-piperidyl))-isoindoline-1,3-dione; *N*-(2-(2,6-dioxo-piperidin-3-yl)-1,3-dioxo-2,3-dihydro-1*H*-isoindol-4-ylmethyl)-acetamide; *N*-{(2-(2,6-dioxo(3-piperidyl)-1,3-dioxoisoindolin-4-yl)methyl}cyclopropyl-carboxamide; 2-chloro-*N*-{(2-(2,6-dioxo(3-piperidyl))-1,3-dioxoisoindolin-4-yl)methyl}acetamide; *N*-(2-(2,6-dioxo(3-piperidyl))-1,3-dioxoisoindolin-4-yl)-3-pyridylcarboxamide; 3-{1-oxo-4-(benzylamino)isoindolin-2-yl}piperidine-2,6-dione; 2-(2,6-dioxo(3-piperidyl))-4-(benzylamino)isoindoline-1,3-dione; *N*-{(2-(2,6-dioxo(3-piperidyl))-1,3-dioxoisoindolin-4-yl)methyl}propanamide; *N*-{(2-(2,6-dioxo(3-piperidyl))-1,3-dioxoisoindolin-4-yl)methyl}-3-pyridylcarboxamide; *N*-{(2-(2,6-dioxo(3-piperidyl))-1,3-dioxoisoindolin-4-yl)methyl}heptanamide; *N*-{(2-(2,6-dioxo(3-piperidyl))-1,3-dioxoisoindolin-4-yl)methyl}-2-furylcarboxamide; {N-(2-(2,6-dioxo(3-piperidyl))-1,3-dioxoisoindolin-4-yl)carbamoyl}methyl acetate; *N*-(2-(2,6-dioxo(3-piperidyl))-1,3-dioxoisoindolin-4-yl)pentanamide; *N*-(2-(2,6-dioxo(3-piperidyl))-1,3-dioxoisoindolin-4-yl)-2-thienylcarboxamide; N-{[2-(2,6-dioxo(3-piperidyl))-1,3-dioxoisoindolin-4-yl] methyl}(butylamino)carboxamide; N-{[2-(2,6-dioxo(3-piperidyl))-1,3-dioxoisoindolin-4-yl] methyl}(octylamino)carboxamide; and N-{[2-(2,6-dioxo(3-piperidyl))-1,3-dioxoisoindolin-4-yl] methyl}(benzylamino)carboxamide.

Still other specific immunomodulatory compounds of the invention belong to a class of isoindole-imides disclosed in U.S. Patent Application Publication Nos. US 2002/0045643, International Publication No. WO 98/54170, and United States Patent No. 6,395,754, each of which is incorporated herein by reference. Representative compounds are of formula III:



III

and pharmaceutically acceptable salts, hydrates, solvates, clathrates, enantiomers, diastereomers, racemates, and mixtures of stereoisomers thereof, wherein:

one of X and Y is C=O and the other is CH_2 or C=O;

R is H or CH_2OCOR';

(i) each of R^1, R^2, R^3, or R^4, independently of the others, is halo, alkyl of 1 to 4 carbon atoms, or alkoxy of 1 to 4 carbon atoms or (ii) one of R^1, R^2, R^3, or R^4 is nitro or $-NHR^5$ and the remaining of R^1, R^2, R^3, or R^4 are hydrogen;

R^5 is hydrogen or alkyl of 1 to 8 carbons

R^6 hydrogen, alkyl of 1 to 8 carbon atoms, benzo, chloro, or fluoro;

R' is R^7-CHR^{10}-$N(R^8R^9)$;

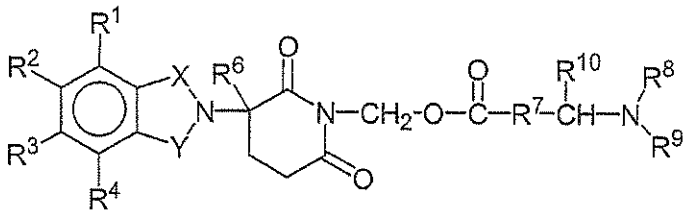
R^7 is m-phenylene or p-phenylene or -(C_nH_{2n})- in which n has a value of 0 to 4;

each of R^8 and R^9 taken independently of the other is hydrogen or alkyl of 1 to 8 carbon atoms, or R^8 and R^9 taken together are tetramethylene, pentamethylene, hexamethylene, or -$CH_2CH_2X_1CH_2CH_2$– in which X_1 is -O-, -S-, or -NH-;

R^{10} is hydrogen, alkyl of to 8 carbon atoms, or phenyl; and

* represents a chiral-carbon center.

Other representative compounds are of formula:



wherein:

one of X and Y is C=O and the other of X and Y is C=O or CH_2;

(i) each of R^1, R^2, R^3, or R^4, independently of the others, is halo, alkyl of 1 to 4 carbon atoms, or alkoxy of 1 to 4 carbon atoms or (ii) one of R^1, R^2, R^3, and R^4 is -NHR^5 and the remaining of R^1, R^2, R^3, and R^4 are hydrogen;

R^5 is hydrogen or alkyl of 1 to 8 carbon atoms;

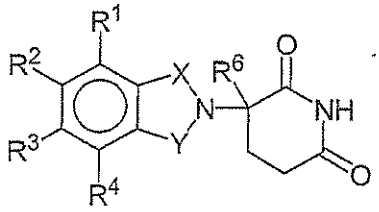
R^6 is hydrogen, alkyl of 1 to 8 carbon atoms, benzo, chloro, or fluoro;

R^7 is m-phenylene or p-phenylene or -(C_nH_{2n})- in which n has a value of 0 to 4;

each of R^8 and R^9 taken independently of the other is hydrogen or alkyl of 1 to 8 carbon atoms, or R^8 and R^9 taken together are tetramethylene, pentamethylene, hexamethylene, or -$CH_2CH_2XCH_2CH_2$- in which X^1 is -O-, -S-, or -NH-;

R^{10} is hydrogen, alkyl of to 8 carbon atoms, or phenyl.

Other representative compounds are of formula:



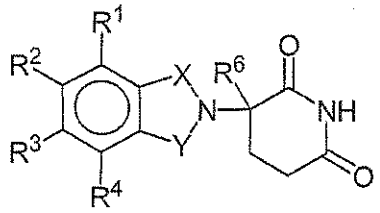
in which

one of X and Y is C=O and the other of X and Y is C=O or CH_2;

each of R^1, R^2, R^3, and R^4, independently of the others, is halo, alkyl of 1 to 4 carbon atoms, or alkoxy of 1 to 4 carbon atoms or (ii) one of R^1, R^2, R^3, and R^4 is nitro or protected amino and the remaining of R^1, R^2, R^3, and R^4 are hydrogen; and

R^6 is hydrogen, alkyl of 1 to 8 carbon atoms, benzo, chloro, or fluoro.

Other representative compounds are of formula:



in which:

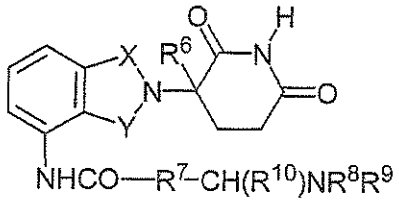
one of X and Y is C=O and the other of X and Y is C=O or CH_2;

(i) each of R^1, R^2, R^3, and R^4, independently of the others, is halo, alkyl of 1 to 4 carbon atoms, or alkoxy of 1 to 4 carbon atoms or (ii) one of R^1, R^2, R^3, and R^4 is $-NHR^5$ and the remaining of R^1, R^2, R^3, and R^4 are hydrogen;

R^5 is hydrogen, alkyl of 1 to 8 carbon atoms, or $CO-R^7-CH(R^{10})NR^8R^9$ in which each of R^7, R^8, R^9, and R^{10} is as herein defined; and

R^6 is alkyl of 1 to 8 carbon atoms, benzo, chloro, or fluoro.

Specific examples of the compounds are of formula:



in which:

one of X and Y is C=O and the other of X and Y is C=O or CH_2;

R^6 is hydrogen, alkyl of 1 to 8 carbon atoms, benzyl, chloro, or fluoro;

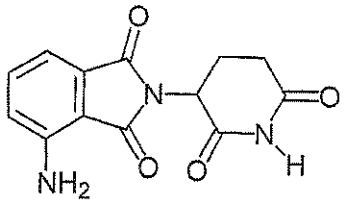
R^7 is m-phenylene, p-phenylene or $-(C_nH_{2n})-$ in which n has a value of 0 to 4;

each of R^8 and R^9 taken independently of the other is hydrogen or alkyl of 1 to 8 carbon atoms, or R^8 and R^9 taken together are tetramethylene, pentamethylene, hexamethylene, or $-CH_2CH_2X^1CH_2CH_2-$ in which X^1 is -O-, -S- or -NH-; and

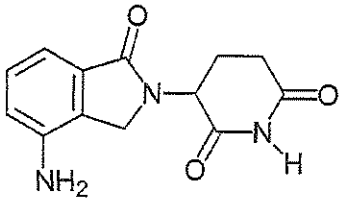
R^{10} is hydrogen, alkyl of 1 to 8 carbon atoms, or phenyl.

The most preferred immunomodulatory compounds of the invention are 4-(amino)-2-(2,6-dioxo(3-piperidyl))-isoindoline-1,3-dione and 3-(4-amino-1-oxo-1,3-dihydro-isoindol-2-yl)-piperidine-2,6-dione. The compounds can be obtained via standard, synthetic methods (*see e.g.*, United States Patent No. 5,635,517, incorporated herein by reference).

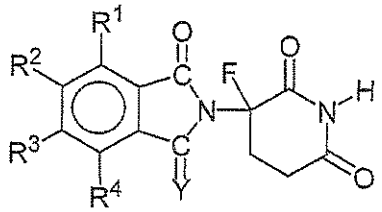
The compounds are available from Celgene Corporation, Warren, NJ. 4-(Amino)-2-(2,6-dioxo(3-piperidyl))-isoindoline-1,3-dione has the following chemical structure:



The compound 3-(4-amino-1-oxo-1,3-dihydro-isoindol-2-yl)-piperidine-2,6-dione has the following chemical structure:



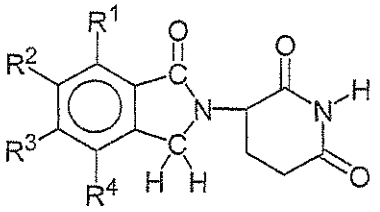
Other specific immunomodulatory compounds of the invention include, but are not limited to, 1-oxo-2-(2,6-dioxo-3-fluoropiperidin-3yl) isoindolines and 1,3-dioxo-2-(2,6-dioxo-3-fluoropiperidine-3-yl) isoindolines such as those described in U.S. patent nos. 5,874,448 and 5,955,476, each of which is incorporated herein by reference. Representative compounds are of formula:



wherein Y is oxygen or H^2 and

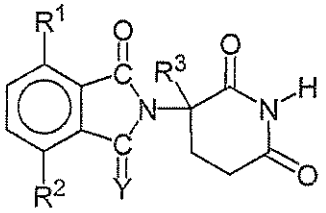
each of R^1, R^2, R^3, and R^4, independently of the others, is hydrogen, halo, alkyl of 1 to 4 carbon atoms, alkoxy of 1 to 4 carbon atoms, or amino.

Other specific immunomodulatory compounds of the invention include, but are not limited to, the tetra substituted 2-(2,6-dioxopiperdin-3-yl)-1-oxoisoindolines described in U.S. patent no. 5,798,368, which is incorporated herein by reference. Representative compounds are of formula:



wherein each of R^1, R^2, R^3, and R^4, independently of the others, is halo, alkyl of 1 to 4 carbon atoms, or alkoxy of 1 to 4 carbon atoms.

Other specific immunomodulatory compounds of the invention include, but are not limited to, 1-oxo and 1,3-dioxo-2-(2,6-dioxopiperidin-3-yl) isoindolines disclosed in U.S. patent no. 6,403,613, which is incorporated herein by reference. Representative compounds are of formula:



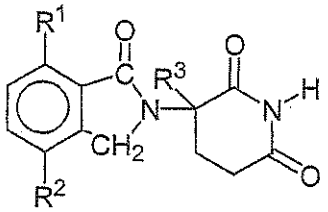
in which

Y is oxygen or H_2,

a first of R^1 and R^2 is halo, alkyl, alkoxy, alkylamino, dialkylamino, cyano, or carbamoyl, the second of R^1 and R^2, independently of the first, is hydrogen, halo, alkyl, alkoxy, alkylamino, dialkylamino, cyano, or carbamoyl, and

R^3 is hydrogen, alkyl, or benzyl.

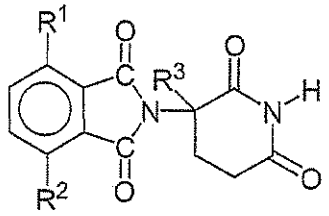
Specific examples of the compounds are of formula:



wherein a first of R^1 and R^2 is halo, alkyl of from 1 to 4 carbon atoms, alkoxy of from 1 to 4 carbon atoms, dialkylamino in which each alkyl is of from 1 to 4 carbon atoms, cyano, or carbamoyl,

the second of R^1 and R^2, independently of the first, is hydrogen, halo, alkyl of from 1 to 4 carbon atoms, alkoxy of from 1 to 4 carbon atoms, alkylamino in which alkyl is of from 1 to 4 carbon atoms, dialkylamino in which each alkyl is of from 1 to 4 carbon atoms, cyano, or carbamoyl, and

R^3 is hydrogen, alkyl of from 1 to 4 carbon atoms, or benzyl. Other representative compounds are of formula:

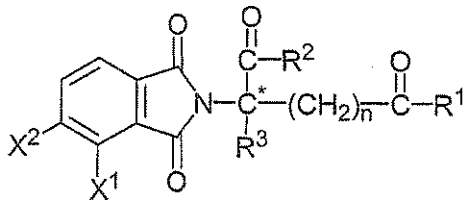


wherein a first of R^1 and R^2 is halo, alkyl of from 1 to 4 carbon atoms, alkoxy of from 1 to 4 carbon atoms, dialkylamino in which each alkyl is of from 1 to 4 carbon atoms, cyano, or carbamoyl,

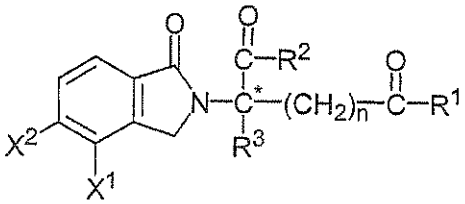
the second of R^1 and R^2, independently of the first, is hydrogen, halo, alkyl of from 1 to 4 carbon atoms, alkoxy of from 1 to 4 carbon atoms, alkylamino in which alkyl is of from 1 to 4 carbon atoms, dialkylamino in which each alkyl is of from 1 to 4 carbon atoms, cyano, or carbamoyl, and

R^3 is hydrogen, alkyl of from 1 to 4 carbon atoms, or benzyl.

Other specific immunomodulatory compounds of the invention include, but are not limited to, 1-oxo and 1,3-dioxoisoindolines substituted in the 4- or 5-position of the indoline ring described in U.S. patent no. 6,380,239, which is incorporated herein by reference. Representative compounds are of formula:



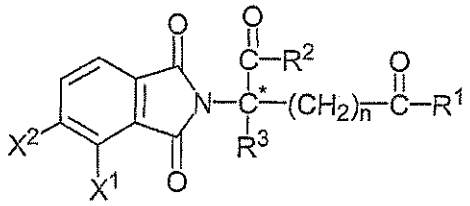
in which the carbon atom designated C* constitutes a center of chirality (when n is not zero and R^1 is not the same as R^2); one of X^1 and X^2 is amino, nitro, alkyl of one to six carbons, or NH-Z, and the other of X^1 or X^2 is hydrogen; each of R^1 and R^2 independent of the other, is hydroxy or NH-Z; R^3 is hydrogen, alkyl of one to six carbons, halo, or haloalkyl; Z is hydrogen, aryl, alkyl of one to six carbons, formyl, or acyl of one to six carbons; and n has a value of 0, 1, or 2; provided that if X^1 is amino, and n is 1 or 2, then R^1 and R^2 are not both hydroxy; and the salts thereof. Further representative compounds are of formula:



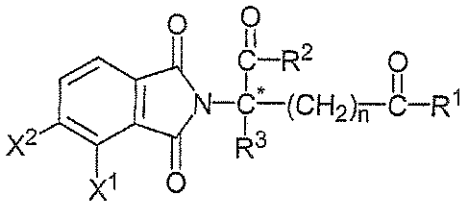
in which the carbon atom designated C* constitutes a center of chirality when n is not zero and R^1 is not R^2; one of X^1 and X^2 is amino, nitro, alkyl of one to six carbons, or

NH-Z, and the other of X^1 or X^2 is hydrogen; each of R^1 and R^2 independent of the other, is hydroxy or NH-Z; R^3 is alkyl of one to six carbons, halo, or hydrogen; Z is hydrogen, aryl or an alkyl or acyl of one to six carbons; and n has a value of 0, 1, or 2.

Other representative compounds are of formula:



in which the carbon atom designated C* constitutes a center of chirality when n is not zero and R^1 is not R^2; one of X^1 and X^2 is amino, nitro, alkyl of one to six carbons, or NH-Z, and the other of X^1or X^2 is hydrogen; each of R^1 and R^2 independent of the other, is hydroxy or NH-Z; R^3 is alkyl of one to six carbons, halo, or hydrogen; Z is hydrogen, aryl, or an alkyl or acyl of one to six carbons; and n has a value of 0, 1, or 2; and the salts thereof. Specific examples of the compounds are of formula:



wherein one of X^1 and X^2 is nitro, or NH-Z, and the other of X^1 or X^2 is hydrogen;

each of R^1 and R^2, independent of the other, is hydroxy or NH-Z;

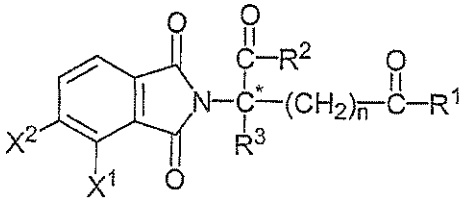
R^3 is alkyl of one to six carbons, halo, or hydrogen;

Z is hydrogen, phenyl, an acyl of one to six carbons, or an alkyl of one to six carbons; and

n has a value of 0, 1, or 2;

provided that if one of X^1 and X^2 is nitro, and n is 1 or 2, then R^1 and R^2 are other than hydroxy; and

if $-COR^1$ and $-(CH_2)_nCOR^2$ are different, the carbon atom designated C* constitutes a center of chirality. Other representative compounds are of formula:



wherein one of X^1 and X^2 is alkyl of one to six carbons;

each of R^1 and R^2, independent of the other, is hydroxy or NH-Z;

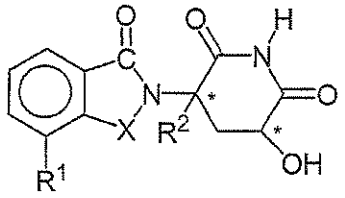
R^3 is alkyl of one to six carbons, halo, or hydrogen;

Z is hydrogen, phenyl, an acyl of one to six carbons, or an alkyl of one to six carbons; and

n has a value of 0, 1, or 2; and

if -COR^1 and -$(CH_2)_nCOR^2$ are different, the carbon atom designated C^* constitutes a center of chirality.

Still other specific immunomodulatory compounds of the invention include, but are not limited to, isoindoline-1-one and isoindoline-1,3-dione substituted in the 2-position with 2,6-dioxo-3-hydroxypiperidin-5-yl described in U.S. patent no. 6,458,810, which is incorporated herein by reference. Representative compounds are of formula:



wherein:

the carbon atoms designated * constitute centers of chirality;

X is -C(O)- or -CH_2-;

R^1 is alkyl of 1 to 8 carbon atoms or -NHR^3;

R^2 is hydrogen, alkyl of 1 to 8 carbon atoms, or halogen;

and

R^3 is hydrogen,

alkyl of 1 to 8 carbon atoms, unsubstituted or substituted with alkoxy of 1 to 8 carbon atoms, halo, amino, or alkylamino of 1 to 4 carbon atoms,

cycloalkyl of 3 to 18 carbon atoms,

phenyl, unsubstituted or substituted with alkyl of 1 to 8 carbon atoms, alkoxy of 1 to 8 carbon atoms, halo, amino, or alkylamino of 1 to 4 carbon atoms,

benzyl, unsubstituted or substituted with alkyl of 1 to 8 carbon atoms, alkoxy of 1 to 8 carbon atoms, halo, amino, or alkylamino of 1 to 4 carbon atoms, or -COR^4 in which

R^4 is hydrogen,

alkyl of 1 to 8 carbon atoms, unsubstituted or substituted with alkoxy of 1 to 8 carbon atoms, halo, amino, or alkylamino of 1 to 4 carbon atoms,

cycloalkyl of 3 to 18 carbon atoms,

phenyl, unsubstituted or substituted with alkyl of 1 to 8 carbon atoms, alkoxy of 1 to 8 carbon atoms, halo, amino, or alkylamino of 1 to 4 carbon atoms, or

benzyl, unsubstituted or substituted with alkyl of 1 to 8 carbon atoms, alkoxy of 1 to 8 carbon atoms, halo, amino, or alkylamino of 1 to 4 carbon atoms.

Compounds of the invention can either be commercially purchased or prepared according to the methods described in the patents or patent publications disclosed herein. Further, optically pure compounds can be asymmetrically synthesized or resolved using known resolving agents or chiral columns as well as other standard synthetic organic chemistry techniques.

As used herein and unless otherwise indicated, the term "pharmaceutically acceptable salt" encompasses non-toxic acid and base addition salts of the compound to which the term refers. Acceptable non-toxic acid addition salts include those derived from organic and inorganic acids or bases know in the art, which include, for example, hydrochloric acid, hydrobromic acid, phosphoric acid, sulfuric acid, methanesulphonic acid, acetic acid, tartaric acid, lactic acid, succinic acid, citric acid, malic acid, maleic acid, sorbic acid, aconitic acid, salicylic acid, phthalic acid, embolic acid, enanthic acid, and the like.

Compounds that are acidic in nature are capable of forming salts with various pharmaceutically acceptable bases. The bases that can be used to prepare pharmaceutically acceptable base addition salts of such acidic compounds are those that form non-toxic base addition salts, *i.e.*, salts containing pharmacologically acceptable cations such as, but not limited to, alkali metal or alkaline earth metal salts and the calcium, magnesium, sodium or potassium salts in particular. Suitable organic bases include, but are not limited to, N,N-dibenzylethylenediamine, chloroprocaine, choline, diethanolamine, ethylenediamine, meglumaine (N-methylglucamine), lysine, and procaine.

As used herein and unless otherwise indicated, the term "prodrug" means a derivative of a compound that can hydrolyze, oxidize, or otherwise react under biological conditions (*in vitro* or *in vivo*) to provide the compound. Examples of prodrugs include, but are not limited to, derivatives of immunomodulatory compounds of the invention that comprise biohydrolyzable moieties such as biohydrolyzable amides, biohydrolyzable esters, biohydrolyzable carbamates, biohydrolyzable carbonates, biohydrolyzable ureides, and biohydrolyzable phosphate analogues. Other examples of prodrugs include derivatives of immunomodulatory compounds of the invention that comprise -NO, $-NO_2$, -ONO, or $-ONO_2$ moieties. Prodrugs can typically be prepared using well-known methods, such as those described in 1 *Burger's Medicinal Chemistry and Drug Discovery*, 172-178, 949-982 (Manfred E. Wolff *ed.*, 5th ed. 1995), and *Design of Prodrugs* (H. Bundgaard ed., Elselvier, New York 1985).

As used herein and unless otherwise indicated, the terms "biohydrolyzable amide," "biohydrolyzable ester," "biohydrolyzable carbamate," "biohydrolyzable carbonate," "biohydrolyzable ureide," "biohydrolyzable phosphate" mean an amide, ester, carbamate,

carbonate, ureide, or phosphate, respectively, of a compound that either: 1) does not interfere with the biological activity of the compound but can confer upon that compound advantageous properties *in vivo*, such as uptake, duration of action, or onset of action; or 2) is biologically inactive but is converted *in vivo* to the biologically active compound. Examples of biohydrolyzable esters include, but are not limited to, lower alkyl esters, lower acyloxyalkyl esters (such as acetoxylmethyl, acetoxyethyl, aminocarbonyloxymethyl, pivaloyloxymethyl, and pivaloyloxyethyl esters), lactonyl esters (such as phthalidyl and thiophthalidyl esters), lower alkoxyacyloxyalkyl esters (such as methoxycarbonyl-oxymethyl, ethoxycarbonyloxyethyl and isopropoxycarbonyloxyethyl esters), alkoxyalkyl esters, choline esters, and acylamino alkyl esters (such as acetamidomethyl esters). Examples of biohydrolyzable amides include, but are not limited to, lower alkyl amides, α-amino acid amides, alkoxyacyl amides, and alkylaminoalkylcarbonyl amides. Examples of biohydrolyzable carbamates include, but are not limited to, lower alkylamines, substituted ethylenediamines, amino acids, hydroxyalkylamines, heterocyclic and heteroaromatic amines, and polyether amines.

Various immunomodulatory compounds of the invention contain one or more chiral centers, and can exist as racemic mixtures of enantiomers or mixtures of diastereomers. This invention encompasses the use of stereomerically pure forms of such compounds, as well as the use of mixtures of those forms. For example, mixtures comprising equal or unequal amounts of the enantiomers of a particular immunomodulatory compounds of the invention may be used in methods and compositions of the invention. These isomers may be asymmetrically synthesized or resolved using standard techniques such as chiral columns or chiral resolving agents. *See, e.g.*, Jacques, J., *et al., Enantiomers, Racemates and Resolutions* (Wiley-Interscience, New York, 1981); Wilen, S. H., *et al., Tetrahedron* 33:2725 (1977); Eliel, E. L., *Stereochemistry of Carbon Compounds* (McGraw-Hill, NY, 1962); and Wilen, S. H., *Tables of Resolving Agents and Optical Resolutions* p. 268 (E.L. Eliel, Ed., Univ. of Notre Dame Press, Notre Dame, IN, 1972).

As used herein and unless otherwise indicated, the term "stereomerically pure" means a composition that comprises one stereoisomer of a compound and is substantially free of other stereoisomers of that compound. For example, a stereomerically pure composition of a compound having one chiral center will be substantially free of the opposite enantiomer of the compound. A stereomerically pure composition of a compound having two chiral centers will be substantially free of other diastereomers of the compound. A typical stereomerically pure compound comprises greater than about 80% by weight of one stereoisomer of the compound and less than about 20% by weight of other

stereoisomers of the compound, more preferably greater than about 90% by weight of one stereoisomer of the compound and less than about 10% by weight of the other stereoisomers of the compound, even more preferably greater than about 95% by weight of one stereoisomer of the compound and less than about 5% by weight of the other stereoisomers of the compound, and most preferably greater than about 97% by weight of one stereoisomer of the compound and less than about 3% by weight of the other stereoisomers of the compound. As used herein and unless otherwise indicated, the term "stereomerically enriched" means a composition that comprises greater than about 60% by weight of one stereoisomer of a compound, preferably greater than about 70% by weight, more preferably greater than about 80% by weight of one stereoisomer of a compound. As used herein and unless otherwise indicated, the term "enantiomerically pure" means a stereomerically pure composition of a compound having one chiral center. Similarly, the term "enantiomerically enriched" means a stereomerically enriched composition of a compound having one chiral center.

It should be noted that if there is a discrepancy between a depicted structure and a name given that structure, the depicted structure is to be accorded more weight. In addition, if the stereochemistry of a structure or a portion of a structure is not indicated with, for example, bold or dashed lines, the structure or portion of the structure is to be interpreted as encompassing all stereoisomers of it.

4.2. SECOND ACTIVE AGENTS

One or more second active ingredients can be used in the methods and compositions of the invention together with an immunomodulatory compound of the invention. In a preferred embodiment, the second active agents are capable of affecting or improving the process of blood cell production. Specific second active agents also stimulate the division and differentiation of committed erythroid progenitors in cells *in vitro* or *in vivo*.

Second active agents can be large molecules (*e.g.*, proteins) or small molecules (*e.g.*, synthetic inorganic, organometallic, or organic molecules). The second active agents include but are not limited to hematopoietic growth factors, cytokines, anti-cancer agents, antibiotics, proteasome inhibitors, immunosuppressive agents and other therapeutics discussed herein. Particular agents include, but are not limited to, G-CSF, GM-CSF, EPO, dexamethasone, topotecan, pentoxifylline, irinotecan, ciprofloxacin, vinorelbine, IL2, IL8, IL18, Ara-C, isotretinoin, 13-cis-retinoic acid, 12-O-tetradecanoylphorbol-13-acetate (TPA), 5-AZA2'-deoyxcytidine, 9-nitrocamp-tothecin, transretinoic acid, amifostine, amphotericin B and liposomal amphotericin B, anti-CD-20 monoclonal antibody, anti-

thymocyle globulin (ATG), arsenic trioxide, azacytidine, bevacizumab, bismuth monoclonal antibody, bryostatin, busulfan, caspofungin acetate, celocoxib, cladribine, cyclophosphamide, cyclosporine, cytarabine, cytosine, daunorubicin, depsipeptide, etoposide, farresy transferase inhibitor, flavopiridol, Flt3 ligand, fludarabine, gentuzumab ozogomicin (mylotarg), etanercept (Enbrel®), imatinib (Glivec®), anti-TNF-α antibodies, infliximab (Remicade®), humanized monoclonal anti-VEGF antibody, idarubicine, leucovorin, melphalan, mitoxantrone, monoclonal antibody ABX-CBL, monoclonal antibody CD52, mycophenolate mofetil, oblimersen, omega-3 fatty acids, pentostatin, phenylbutyrate, PR1 leukemia peptide vaccine, montanide, proteasome inhibitor, sodium phenyl-butyrate, sodium salicylate, temozolomide, thymoglobulin, troxatyl, tumor necrosis factor receptor IgG chimera, Yttrium Y 90 humanized monoclonal antibody M195. In a specific embodiment of the invention, an immunomodulatory compound of the invention is used in combination with pentoxifylline, ciprofloxacin, and/or dexamethasone.

This invention also encompasses the use of native, naturally occurring, and recombinant proteins. The invention further encompasses mutants and derivatives (*e.g.*, modified forms) of naturally occurring proteins that exhibit, *in vivo*, at least some of the pharmacological activity of the proteins upon which they are based. Examples of mutants include, but are not limited to, proteins that have one or more amino acid residues that differ from the corresponding residues in the naturally occurring forms of the proteins. Also encompassed by the term "mutants" are proteins that lack carbohydrate moieties normally present in their naturally occurring forms (*e.g.*, nonglycosylated forms). Examples of derivatives include, but are not limited to, pegylated derivatives and fusion proteins, such as proteins formed by fusing IgG1 or IgG3 to the protein or active portion of the protein of interest. *See, e.g.*, Penichet, M.L. and Morrison, S.L., *J. Immunol. Methods* 248:91-101 (2001).

Recombinant and mutated forms of G-CSF can be prepared as described in U.S. patent nos. 4,810,643; 4,999,291; 5,528,823; and 5,580,755; all of which are incorporated herein by reference. Recombinant and mutated forms of GM-CSF can be prepared as described in U.S. patent nos. 5,391,485; 5,393,870; and 5,229,496; all of which are incorporated herein by reference. In fact, recombinant forms of G-CSF and GM-CSF are currently sold in the United States for the treatment of symptoms associated with specific chemotherapies. A recombinant form of G-CSF known as filgrastim is sold in the United States under the trade name NEUPOGEN®. NEUPOGEN® is known to stimulate division and maturation of granulocytes, mostly neutrophils, in MDS patients and to enhance erythroid response in combination with EPO. *Physicians' Desk Reference*, 587-592 (56th ed., 2002).

A recombinant form of GM-CSF known as sargramostim is also sold in the United States under the trade name LEUKINE®. LEUKINE® is known to stimulate division and maturation of earlier myeloid and macrophage precursor cells and has been reported to increase granulocytes. *Physicians' Desk Reference*, 1755-1760 (56th ed., 2002). A recombinant form of EPO known as epoetin alfa is sold in the United States under the trade name EPOGEN®. EPOGEN® is used to stimulate red cell production by stimulating division and maturation of committed red cell precursor cells. EPOGEN® has been reported to be effective in 20-26% of MDS patient when administered by itself and in as many as 48% of patients when combined with G-CSF or GM-CSF. *Physicians' Desk Reference*, 582-587 (56th ed., 2002).

A growth-factor or cytokine such as G-CSF, GM-CSF and EPO can also be administered in the form of a vaccine. For example, vaccines that secrete, or cause the secretion of, cytokines such as G-CSF and GM-CSF can be used in the methods, pharmaceutical compositions, and kits of the invention. *See, e.g.*, Emens, L.A., *et al.*, *Curr. Opinion Mol. Ther.* 3(1):77-84 (2001).

Other compounds that can be administered or used in combination with an immunomodulatory compound of the invention include those disclosed in U.S. provisional patent application no. 60/380,842, filed May 17, 2002, and U.S. provisional patent application no. 60/380,843, filed May 17, 2002, both of which are incorporated herein by reference.

4.3. METHODS OF TREATMENT AND MANAGEMENT

Methods of this invention encompass methods of preventing, treating and/or managing various types of MDS. As used herein, unless otherwise specified, the term "preventing" includes but is not limited to, inhibition or the averting of symptoms associated with MDS. The symptoms associated with MDS include, but are not limited to, anemia, thrombocytopenia, neutropenia, cytopenia, bicytopenia (two deficient cell lines), and pancytopenia (three deficient cell lines). As used herein, unless otherwise specified, the term "treating" refers to the administration of a composition after the onset of symptoms of MDS, whereas "preventing" refers to the administration prior to the onset of symptoms, particularly to patients at risk of MDS. As used herein and unless otherwise indicated, the term "managing" encompasses preventing the recurrence of MDS in a patient who had suffered from MDS, lengthening the time a patient who had suffered from MDS remains in remission, and/or preventing the occurrence of MDS in patients at risk of suffering from MDS.

The invention encompasses methods of treating or preventing patients with primary and secondary MDS. It further encompasses methods treating patients who have been previously treated for MDS, as well as those who have not previously been treated for MDS. Because patients with MDS have heterogenous clinical manifestations and varying clinical outcomes, it has become apparent that staging the patients according to their prognosis and approaching therapy depending on the severity and stage is necessary. Indeed, the methods and compositions of this invention can be used in various stages of treatments for patients with one or more types of MDS including, but not limited to, refractory anemia (RA), RA with ringed sideroblasts (RARS), RA with excess blasts (RAEB), RAEB in transformation (RAEB-T), or chronic myelomonocytic leukemia (CMML). The invention also contemplates treating patients diagnosed using the IPSS for MDS discussed above. Greenberg *et al.*, *Blood* 1997 (89):2079-88.

Methods encompassed by this invention comprise administering an immunomodulatory compound of the invention, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof to a patient (*e.g.*, a human) suffering, or likely to suffer, from MDS. Specific patient populations include the elderly, *i.e.*, ages 60 and above as well as those over 35 years of age. Patients with familial history of MDS or leukemia are also preferred candidates for preventive regimens.

In one embodiment of the invention, an immunomodulatory compound of the invention is administered orally and in a single or divided daily doses in an amount of from about 0.10 to about 150 mg/day. In a particular embodiment, 4-(amino)-2-(2,6-dioxo(3-piperidyl))-isoindoline-1,3-dione (Actimid™) is administered in an amount of from about 0.1 to about 1 mg per day, or alternatively about 5 mg every other day. 3-(4-amino-1-oxo-1,3-dihydro-isoindol-2-yl)-piperidine-2,6-dione (Revimid™) can be preferably administered in an amount of from about 5 to 25 mg per day, or alternatively from about 25 to about 50 mg every other day.

4.3.1 Combination Therapy With A Second Active Agent

Particular methods of the invention comprise comprises administering 1) an immunomodulatory compound of the invention, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof, and 2) a second active agent or active ingredient. Examples of immunomodulatory compounds of the invention are disclosed herein (*see, e.g.*, section 4.1); and examples of the second active agents are also disclosed herein (*see, e.g.*, section 4.2).

Administration of the immunomodulatory compounds and the second active agents to a patient can occur simultaneously or sequentially by the same or different routes of administration. The suitability of a particular route of administration employed for a particular active agent will depend on the active agent itself (*e.g.*, whether it can be administered orally without decomposing prior to entering the blood stream) and the disease being treated. A preferred route of administration for an immunomodulatory compound is oral. Preferred routes of administration for the second active agents or ingredients of the invention are known to those of ordinary skill in the art. *See, e.g., Physicians' Desk Reference*, 1755-1760 (56th ed., 2002).

In one embodiment, the second active agent is administered intravenously or subcutaneously and once or twice daily in an amount of from about 1 to about 1000 mg, from about 5 to about 500 mg, from about 10 to about 350 mg, or from about 50 to about 200 mg. The specific amount of the second active agent will depend on the specific agent used, the type of MDS being treated or managed, the severity and stage of MDS, and the amount(s) of immunomodulatory compounds of the invention and any optional additional active agents concurrently administered to the patient. In a particular embodiment, the second active agent is GM-CSF, G-CSF, EPO, transretinoic acid, dexamethasone, topotecan, pentoxifylline, ciprofloxacin, dexamethasone, IL2, IL8, IL18, Ara-C, vinorelbine, or a combination thereof. GM-CSF is administered in an amount of from about 60 to about 500 mcg/m^2 intravenously over 2 hours, or from about 5 to about 12 mcg/m^2/day subcutaneously. G-CSF is administered subcutaneously in an amount of about 1 mcg/kg/day initially and can be adjusted depending on rise of total granulocyte counts. The maintenance dose is 300 (in smaller patients) or 480 mcg subcutaneously. EPO is administered subcutaneously in an amount of 10,000 Unit 3 times per week.

4.3.2 Use With Transplantation Therapy

In still another embodiment, this invention encompasses a method of treating, preventing and/or managing MDS, which comprises administering the immunomodulatory compound of the invention, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof, in conjunction with transplantation therapy. As discussed elsewhere herein, the treatment of MDS is based on the stages and mechanism of the disease. As inevitable leukemic transformation develops in certain stages of MDS, transplantation of peripheral blood stem cells, hematopoietic stem cell preparation or bone marrow may be necessary. The combined use of the immunomodulatory compound of the invention and transplantation therapy provides a unique and unexpected synergism. In

particular, an immunomodulatory compound of the invention exhibits immunomodulatory activity that may provide additive or synergistic effects when given concurrently with transplantation therapy in patients with MDS. An immunomodulatory compound of the invention can work in combination with transplantation therapy reducing complications associated with the invasive procedure of transplantation and risk of related Graft Versus Host Disease (GVHD). This invention encompasses a method of treating, preventing and/or managing MDS which comprises administering to a patient (*e.g.*, a human) an immunomodulatory compound of the invention, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof, before, during, or after the transplantation of umbilical cord blood, placental blood, peripheral blood stem cell, hematopoietic stem cell preparation or bone marrow. Examples of stem cells suitable for use in the methods of the invention are disclosed in U.S. provisional patent application no. 60/372,348, filed April 12, 2002 by R. Hariri *et al.*, the entirety of which is incorporated herein by reference.

4.3.3. Cycling Therapy

In certain embodiments, the prophylactic or therapeutic agents of the invention are cyclically administered to a patient. Cycling therapy involves the administration of a first agent for a period of time, followed by the administration of the agent and/or the second agent for a period of time and repeating this sequential administration. Cycling therapy can reduce the development of resistance to one or more of the therapies, avoid or reduce the side effects of one of the therapies, and/or improves the efficacy of the treatment.

In a particular embodiment, prophylactic or therapeutic agents are administered in a cycle of about 16 weeks, about once or twice every day. One cycle can comprise the administration of a therapeutic or prophylactic agent and at least one (1) or three (3) weeks of rest. The number of cycles administered is from about 1 to about 12 cycles, more typically from about 2 to about 10 cycles, and more typically from about 2 to about 8 cycles.

4.4. PHARMACEUTICAL COMPOSITIONS AND SINGLE UNIT DOSAGE FORMS

Pharmaceutical compositions can be used in the preparation of individual, single unit dosage forms. Pharmaceutical compositions and dosage forms of the invention comprise an immunomodulatory compound of the invention, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof. Pharmaceutical compositions and dosage forms of the invention can further comprise one or more excipients.

Pharmaceutical compositions and dosage forms of the invention can also comprise one or more additional active ingredients. Consequently, pharmaceutical compositions and dosage forms of the invention comprise the active ingredients disclosed herein (*e.g.*, an immunomodulatory compound of the invention, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof, and a second active ingredient). Examples of optional additional active ingredients are disclosed herein (*see, e.g.*, section 4.2).

Single unit dosage forms of the invention are suitable for oral, mucosal (*e.g.*, nasal, sublingual, vaginal, buccal, or rectal), or parenteral (*e.g.*, subcutaneous, intravenous, bolus injection, intramuscular, or intraarterial), transdermal or transcutaneous administration to a patent. Examples of dosage forms include, but are not limited to: tablets; caplets; capsules, such as soft elastic gelatin capsules; cachets; troches; lozenges; dispersions; suppositories; powders; aerosols (*e.g.*, nasal sprays or inhalers); gels; liquid dosage forms suitable for oral or mucosal administration to a patient, including suspensions (*e.g.*, aqueous or non-aqueous liquid suspensions, oil-in-water emulsions, or a water-in-oil liquid emulsions), solutions, and elixirs; liquid dosage forms suitable for parenteral administration to a patient; and sterile solids (*e.g.*, crystalline or amorphous solids) that can be reconstituted to provide liquid dosage forms suitable for parenteral administration to a patient.

The composition, shape, and type of dosage forms of the invention will typically vary depending on their use. For example, a dosage form used in the acute treatment of a disease may contain larger amounts of one or more of the active ingredients it comprises than a dosage form used in the chronic treatment of the same disease. Similarly, a parenteral dosage form may contain smaller amounts of one or more of the active ingredients it comprises than an oral dosage form used to treat the same disease. These and other ways in which specific dosage forms encompassed by this invention will vary from one another will be readily apparent to those skilled in the art. *See, e.g., Remington's Pharmaceutical Sciences*, 18th ed., Mack Publishing, Easton PA (1990).

Typical pharmaceutical compositions and dosage forms comprise one or more excipients. Suitable excipients are well known to those skilled in the art of pharmacy, and non-limiting examples of suitable excipients are provided herein. Whether a particular excipient is suitable for incorporation into a pharmaceutical composition or dosage form depends on a variety of factors well known in the art including, but not limited to, the way in which the dosage form will be administered to a patient. For example, oral dosage forms such as tablets may contain excipients not suited for use in parenteral dosage forms. The suitability of a particular excipient may also depend on the specific active ingredients in the

dosage form. For example, the decomposition of some active ingredients may be accelerated by some excipients such as lactose, or when exposed to water. Active ingredients that comprise primary or secondary amines are particularly susceptible to such accelerated decomposition. Consequently, this invention encompasses pharmaceutical compositions and dosage forms that contain little, if any, lactose other mono- or di-saccharides. As used herein, the term "lactose-free" means that the amount of lactose present, if any, is insufficient to substantially increase the degradation rate of an active ingredient.

Lactose-free compositions of the invention can comprise excipients that are well known in the art and are listed, for example, in the *U.S. Pharmacopeia* (USP) 25-NF20 (2002). In general, lactose-free compositions comprise active ingredients, a binder/filler, and a lubricant in pharmaceutically compatible and pharmaceutically acceptable amounts. Preferred lactose-free dosage forms comprise active ingredients, microcrystalline cellulose, pre-gelatinized starch, and magnesium stearate.

This invention further encompasses anhydrous pharmaceutical compositions and dosage forms comprising active ingredients, since water can facilitate the degradation of some compounds. For example, the addition of water (*e.g.*, 5%) is widely accepted in the pharmaceutical arts as a means of simulating long-term storage in order to determine characteristics such as shelf-life or the stability of formulations over time. *See, e.g.*, Jens T. Carstensen, *Drug Stability: Principles & Practice*, 2d. Ed., Marcel Dekker, NY, NY, 1995, pp. 379-80. In effect, water and heat accelerate the decomposition of some compounds. Thus, the effect of water on a formulation can be of great significance since moisture and/or humidity are commonly encountered during manufacture, handling, packaging, storage, shipment, and use of formulations.

Anhydrous pharmaceutical compositions and dosage forms of the invention can be prepared using anhydrous or low moisture containing ingredients and low moisture or low humidity conditions. Pharmaceutical compositions and dosage forms that comprise lactose and at least one active ingredient that comprises a primary or secondary amine are preferably anhydrous if substantial contact with moisture and/or humidity during manufacturing, packaging, and/or storage is expected.

An anhydrous pharmaceutical composition should be prepared and stored such that its anhydrous nature is maintained. Accordingly, anhydrous compositions are preferably packaged using materials known to prevent exposure to water such that they can be included in suitable formulary kits. Examples of suitable packaging include, but are not

limited to, hermetically sealed foils, plastics, unit dose containers (*e.g.*, vials), blister packs, and strip packs.

The invention further encompasses pharmaceutical compositions and dosage forms that comprise one or more compounds that reduce the rate by which an active ingredient will decompose. Such compounds, which are referred to herein as "stabilizers," include, but are not limited to, antioxidants such as ascorbic acid, pH buffers, or salt buffers.

Like the amounts and types of excipients, the amounts and specific types of active ingredients in a dosage form may differ depending on factors such as, but not limited to, the route by which it is to be administered to patients. However, typical dosage forms of the invention comprise an immunomodulatory compound of the invention, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof in an amount of from about 0.10 to about 150 mg. Typical dosage forms comprise an immunomodulatory compound of the invention, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof in an amount of about 0.1, 1, 2, 5, 7.5, 10, 12.5, 15, 17.5, 20, 25, 50, 100, 150 or 200 mg. In a particular embodiment, a preferred dosage form comprises 4-(amino)-2-(2,6-dioxo(3-piperidyl))-isoindoline-1,3-dione (Actimid™) in an amount of about 1, 2, 5, 10, 25 or 50mg. In a specific embodiment, a preferred dosage form comprises 3-(4-amino-1-oxo-1,3-dihydro-isoindol-2-yl)-piperidine-2,6-dione (Revimid™) in an amount of about 5, 10, 25 or 50mg. Typical dosage forms comprise the second active ingredient in an amount of 1 to about 1000 mg, from about 5 to about 500 mg, from about 10 to about 350 mg, or from about 50 to about 200 mg. Of course, the specific amount of the second active ingredient will depend on the specific agent used, the type of MDS being treated or managed, and the amount(s) of immunomodulatory compounds of the invention, and any optional additional active agents concurrently administered to the patient.

4.4.1 ORAL DOSAGE FORMS

Pharmaceutical compositions of the invention that are suitable for oral administration can be presented as discrete dosage forms, such as, but are not limited to, tablets (*e.g.*, chewable tablets), caplets, capsules, and liquids (*e.g.*, flavored syrups). Such dosage forms contain predetermined amounts of active ingredients, and may be prepared by methods of pharmacy well known to those skilled in the art. *See generally, Remington's Pharmaceutical Sciences*, 18th ed., Mack Publishing, Easton PA (1990).

Typical oral dosage forms of the invention are prepared by combining the active ingredients in an intimate admixture with at least one excipient according to conventional

pharmaceutical compounding techniques. Excipients can take a wide variety of forms depending on the form of preparation desired for administration. For example, excipients suitable for use in oral liquid or aerosol dosage forms include, but are not limited to, water, glycols, oils, alcohols, flavoring agents, preservatives, and coloring agents. Examples of excipients suitable for use in solid oral dosage forms (*e.g.*, powders, tablets, capsules, and caplets) include, but are not limited to, starches, sugars, micro-crystalline cellulose, diluents, granulating agents, lubricants, binders, and disintegrating agents.

Because of their ease of administration, tablets and capsules represent the most advantageous oral dosage unit forms, in which case solid excipients are employed. If desired, tablets can be coated by standard aqueous or nonaqueous techniques. Such dosage forms can be prepared by any of the methods of pharmacy. In general, pharmaceutical compositions and dosage forms are prepared by uniformly and intimately admixing the active ingredients with liquid carriers, finely divided solid carriers, or both, and then shaping the product into the desired presentation if necessary.

For example, a tablet can be prepared by compression or molding. Compressed tablets can be prepared by compressing in a suitable machine the active ingredients in a free-flowing form such as powder or granules, optionally mixed with an excipient. Molded tablets can be made by molding in a suitable machine a mixture of the powdered compound moistened with an inert liquid diluent.

Examples of excipients that can be used in oral dosage forms of the invention include, but are not limited to, binders, fillers, disintegrants, and lubricants. Binders suitable for use in pharmaceutical compositions and dosage forms include, but are not limited to, corn starch, potato starch, or other starches, gelatin, natural and synthetic gums such as acacia, sodium alginate, alginic acid, other alginates, powdered tragacanth, guar gum, cellulose and its derivatives (*e.g.*, ethyl cellulose, cellulose acetate, carboxymethyl cellulose calcium, sodium carboxymethyl cellulose), polyvinyl pyrrolidone, methyl cellulose, pre-gelatinized starch, hydroxypropyl methyl cellulose, (*e.g.*, Nos. 2208, 2906, 2910), microcrystalline cellulose, and mixtures thereof.

Suitable forms of microcrystalline cellulose include, but are not limited to, the materials sold as AVICEL-PH-101, AVICEL-PH-103 AVICEL RC-581, AVICEL-PH-105 (available from FMC Corporation, American Viscose Division, Avicel Sales, Marcus Hook, PA), and mixtures thereof. An specific binder is a mixture of microcrystalline cellulose and sodium carboxymethyl cellulose sold as AVICEL RC-581. Suitable anhydrous or low moisture excipients or additives include AVICEL-PH-103™ and Starch 1500 LM.

Examples of fillers suitable for use in the pharmaceutical compositions and dosage forms disclosed herein include, but are not limited to, talc, calcium carbonate (*e.g.*, granules or powder), microcrystalline cellulose, powdered cellulose, dextrates, kaolin, mannitol, silicic acid, sorbitol, starch, pre-gelatinized starch, and mixtures thereof. The binder or filler in pharmaceutical compositions of the invention is typically present in from about 50 to about 99 weight percent of the pharmaceutical composition or dosage form.

Disintegrants are used in the compositions of the invention to provide tablets that disintegrate when exposed to an aqueous environment. Tablets that contain too much disintegrant may disintegrate in storage, while those that contain too little may not disintegrate at a desired rate or under the desired conditions. Thus, a sufficient amount of disintegrant that is neither too much nor too little to detrimentally alter the release of the active ingredients should be used to form solid oral dosage forms of the invention. The amount of disintegrant used varies based upon the type of formulation, and is readily discernible to those of ordinary skill in the art. Typical pharmaceutical compositions comprise from about 0.5 to about 15 weight percent of disintegrant, preferably from about 1 to about 5 weight percent of disintegrant.

Disintegrants that can be used in pharmaceutical compositions and dosage forms of the invention include, but are not limited to, agar-agar, alginic acid, calcium carbonate, microcrystalline cellulose, croscarmellose sodium, crospovidone, polacrilin potassium, sodium starch glycolate, potato or tapioca starch, other starches, pre-gelatinized starch, other starches, clays, other algins, other celluloses, gums, and mixtures thereof.

Lubricants that can be used in pharmaceutical compositions and dosage forms of the invention include, but are not limited to, calcium stearate, magnesium stearate, mineral oil, light mineral oil, glycerin, sorbitol, mannitol, polyethylene glycol, other glycols, stearic acid, sodium lauryl sulfate, talc, hydrogenated vegetable oil (*e.g.*, peanut oil, cottonseed oil, sunflower oil, sesame oil, olive oil, corn oil, and soybean oil), zinc stearate, ethyl oleate, ethyl laureate, agar, and mixtures thereof. Additional lubricants include, for example, a syloid silica gel (AEROSIL200, manufactured by W.R. Grace Co. of Baltimore, MD), a coagulated aerosol of synthetic silica (marketed by Degussa Co. of Plano, TX), CAB-O-SIL (a pyrogenic silicon dioxide product sold by Cabot Co. of Boston, MA), and mixtures thereof. If used at all, lubricants are typically used in an amount of less than about 1 weight percent of the pharmaceutical compositions or dosage forms into which they are incorporated.

A preferred solid oral dosage form of the invention comprises an immunomodulatory compound of the invention, anhydrous lactose, microcrystalline cellulose, polyvinylpyrrolidone, stearic acid, colloidal anhydrous silica, and gelatin.

4.4.2 DELAYED RELEASE DOSAGE FORMS

Active ingredients of the invention can be administered by controlled release means or by delivery devices that are well known to those of ordinary skill in the art. Examples include, but are not limited to, those described in U.S. Patent Nos.: 3,845,770; 3,916,899; 3,536,809; 3,598,123; and 4,008,719, 5,674,533, 5,059,595, 5,591,767, 5,120,548, 5,073,543, 5,639,476, 5,354,556, and 5,733,566, each of which is incorporated herein by reference. Such dosage forms can be used to provide slow or controlled-release of one or more active ingredients using, for example, hydropropylmethyl cellulose, other polymer matrices, gels, permeable membranes, osmotic systems, multilayer coatings, microparticles, liposomes, microspheres, or a combination thereof to provide the desired release profile in varying proportions. Suitable controlled-release formulations known to those of ordinary skill in the art, including those described herein, can be readily selected for use with the active ingredients of the invention. The invention thus encompasses single unit dosage forms suitable for oral administration such as, but not limited to, tablets, capsules, gelcaps, and caplets that are adapted for controlled-release.

All controlled-release pharmaceutical products have a common goal of improving drug therapy over that achieved by their non-controlled counterparts. Ideally, the use of an optimally designed controlled-release preparation in medical treatment is characterized by a minimum of drug substance being employed to cure or control the condition in a minimum amount of time. Advantages of controlled-release formulations include extended activity of the drug, reduced dosage frequency, and increased patient compliance. In addition, controlled-release formulations can be used to affect the time of onset of action or other characteristics, such as blood levels of the drug, and can thus affect the occurrence of side (*e.g.*, adverse) effects.

Most controlled-release formulations are designed to initially release an amount of drug (active ingredient) that promptly produces the desired therapeutic effect, and gradually and continually release of other amounts of drug to maintain this level of therapeutic or prophylactic effect over an extended period of time. In order to maintain this constant level of drug in the body, the drug must be released from the dosage form at a rate that will replace the amount of drug being metabolized and excreted from the body. Controlled-release of an active ingredient can be stimulated by various conditions including, but not

limited to, pH, temperature, enzymes, water, or other physiological conditions or compounds.

4.4.3 PARENTERAL DOSAGE FORMS

Parenteral dosage forms can be administered to patients by various routes including, but not limited to, subcutaneous, intravenous (including bolus injection), intramuscular, and intraarterial. Because their administration typically bypasses patients' natural defenses against contaminants, parenteral dosage forms are preferably sterile or capable of being sterilized prior to administration to a patient. Examples of parenteral dosage forms include, but are not limited to, solutions ready for injection, dry products ready to be dissolved or suspended in a pharmaceutically acceptable vehicle for injection, suspensions ready for injection, and emulsions.

Suitable vehicles that can be used to provide parenteral dosage forms of the invention are well known to those skilled in the art. Examples include, but are not limited to: Water for Injection USP; aqueous vehicles such as, but not limited to, Sodium Chloride Injection, Ringer's Injection, Dextrose Injection, Dextrose and Sodium Chloride Injection, and Lactated Ringer's Injection; water-miscible vehicles such as, but not limited to, ethyl alcohol, polyethylene glycol, and polypropylene glycol; and non-aqueous vehicles such as, but not limited to, corn oil, cottonseed oil, peanut oil, sesame oil, ethyl oleate, isopropyl myristate, and benzyl benzoate.

Compounds that increase the solubility of one or more of the active ingredients disclosed herein can also be incorporated into the parenteral dosage forms of the invention. For example, cyclodextrin and its derivatives can be used to increase the solubility of an immunomodulatory compound of the invention, and its derivatives. *See, e.g.*, U.S. Patent No. 5,134,127, which is incorporated herein by reference.

4.4.4 TOPICAL AND MUCOSAL DOSAGE FORMS

Topical and mucosal dosage forms of the invention include, but are not limited to, sprays, aerosols, solutions, emulsions, suspensions, or other forms known to one of skill in the art. *See, e.g., Remington's Pharmaceutical Sciences*, 16th and 18th eds., Mack Publishing, Easton PA (1980 & 1990); and *Introduction to Pharmaceutical Dosage Forms*, 4th ed., Lea & Febiger, Philadelphia (1985). Dosage forms suitable for treating mucosal tissues within the oral cavity can be formulated as mouthwashes or as oral gels.

Suitable excipients (*e.g.*, carriers and diluents) and other materials that can be used to provide topical and mucosal dosage forms encompassed by this invention are well known to those skilled in the pharmaceutical arts, and depend on the particular tissue to which a

given pharmaceutical composition or dosage form will be applied. With that fact in mind, typical excipients include, but are not limited to, water, acetone, ethanol, ethylene glycol, propylene glycol, butane-1,3-diol, isopropyl myristate, isopropyl palmitate, mineral oil, and mixtures thereof to form solutions, emulsions or gels, which are non-toxic and pharmaceutically acceptable. Moisturizers or humectants can also be added to pharmaceutical compositions and dosage forms if desired. Examples of such additional ingredients are well known in the art. *See, e.g., Remington's Pharmaceutical Sciences*, 16th and 18th eds., Mack Publishing, Easton PA (1980 & 1990).

The pH of a pharmaceutical composition or dosage form may also be adjusted to improve delivery of one or more active ingredients. Similarly, the polarity of a solvent carrier, its ionic strength, or tonicity can be adjusted to improve delivery. Compounds such as stearates can also be added to pharmaceutical compositions or dosage forms to advantageously alter the hydrophilicity or lipophilicity of one or more active ingredients so as to improve delivery. In this regard, stearates can serve as a lipid vehicle for the formulation, as an emulsifying agent or surfactant, and as a delivery-enhancing or penetration-enhancing agent. Different salts, hydrates or solvates of the active ingredients can be used to further adjust the properties of the resulting composition.

4.4.5 KITS

Typically, active ingredients of the invention are preferably not administered to a patient at the same time or by the same route of administration. This invention therefore encompasses kits which, when used by the medical practitioner, can simplify the administration of appropriate amounts of active ingredients to a patient.

A typical kit of the invention comprises a dosage form of an immunomodulatory compound of the invention, or a pharmaceutically acceptable salt salt, solvate, hydrate, stereoisomer, prodrug, or clathrate thereof. Kits encompassed by this invention can further comprise additional active ingredients such as G-CSF, GM-CSF, EPO, topotecan, pentoxifylline, ciprofloxacin, dexamethasone, IL2, IL8, IL18, Ara-C, vinorelbine, isotretinoin, 13-cis-retinoic acid, or a pharmacologically active mutant or derivative thereof, or a combination thereof. Examples of the additional active ingredients include, but are not limited to, those disclosed herein (*see, e.g.*, section 4.2).

Kits of the invention can further comprise devices that are used to administer the active ingredients. Examples of such devices include, but are not limited to, syringes, drip bags, patches, and inhalers.

Kits of the invention can further comprise cells or blood for transplantation as well as pharmaceutically acceptable vehicles that can be used to administer one or more active ingredients. For example, if an active ingredient is provided in a solid form that must be reconstituted for parenteral administration, the kit can comprise a sealed container of a suitable vehicle in which the active ingredient can be dissolved to form a particulate-free sterile solution that is suitable for parenteral administration. Examples of pharmaceutically acceptable vehicles include, but are not limited to: Water for Injection USP; aqueous vehicles such as, but not limited to, Sodium Chloride Injection, Ringer's Injection, Dextrose Injection, Dextrose and Sodium Chloride Injection, and Lactated Ringer's Injection; water-miscible vehicles such as, but not limited to, ethyl alcohol, polyethylene glycol, and polypropylene glycol; and non-aqueous vehicles such as, but not limited to, corn oil, cottonseed oil, peanut oil, sesame oil, ethyl oleate, isopropyl myristate, and benzyl benzoate.

5. EXAMPLES

The following studies are intended to further illustrate the invention without limiting its scope.

Excessive production of the growth inhibitory cytokine TNF-α is demonstrated in bone marrow plasma of patients with MDS, implicating TNF-α as a critical negative regulator of erythroid progenitor survival in the disorder. As a result, a study with an immunomodulatory compound of the invention was conducted.

5.1. PHARMACOLOGY AND TOXICOLOGY STUDIES

A series of non-clinical pharmacology and toxicology studies have been performed to support the clinical evaluation of an immunomodulatory compound of the invention in human subjects. These studies were performed in accordance with internationally recognized guidelines for study design and in compliance with the requirements of Good Laboratory Practice (GLP), unless otherwise noted.

The pharmacological properties of 3-(4-amino-1-oxo-1,3-dihydro-isoindol-2-yl)-piperidine-2,6-dione, including activity comparisons with thalidomide, have been characterized in *in vitro* studies. Studies examined the effects of 3-(4-amino-1-oxo-1,3-dihydro-isoindol-2-yl)-piperidine-2,6-dione or thalidomide on the production of various cytokines. In all studies, 3-(4-amino-1-oxo-1,3-dihydro-isoindol-2-yl)-piperidine-2,6-dione was at least 50 times more potent than thalidomide. In addition, a safety pharmacology study of 3-(4-amino-1-oxo-1,3-dihydro-isoindol-2-yl)-piperidine-2,6-dione has been conducted in dogs and the effects of 3-(4-amino-1-oxo-1,3-dihydro-isoindol-2-yl)-

piperidine-2,6-dione on ECG parameters were examined further as part of three repeat-dose toxicity studies in primates. The results of these studies are described below.

5.2. MODULATION OF CYTOKINE PRODUCTION

Inhibition of TNF-α production following LPS-stimulation of human PBMC and human whole blood by 3-(4-amino-1-oxo-1,3-dihydro-isoindol-2-yl)-piperidine-2,6-dione or thalidomide was investigated *in vitro* (Muller *et al.*, *Bioorg. Med. Chem. Lett.* 9:1625-1630, 1999). The IC_{50}'s of 3-(4-amino-1-oxo-1,3-dihydro-isoindol-2-yl)-piperidine-2,6-dione for inhibiting production of TNF-α following LPS-stimulation of PBMC and human whole blood were ~100 nM (25.9 ng/mL) and ~480 nM (103.6 ng/mL), respectively. Thalidomide, in contrast, had an IC_{50} of ~194 μM (50.2 μg/mL) for inhibiting production of TNF-α following LPS-stimulation of PBMC.

In vitro studies suggest a pharmacological activity profile for 3-(4-amino-1-oxo-1,3-dihydro-isoindol-2-yl)-piperidine-2,6-dione that is similar to, but 50 to 2000 times more potent than, thalidomide. The pharmacological effects of 3-(4-amino-1-oxo-1,3-dihydro-isoindol-2-yl)-piperidine-2,6-dione derive from its action as an inhibitor of cellular response to receptor-initiated trophic signals (*e.g.*, IGF-1, VEGF, cyclooxygenase-2), and other activities. As a result, 3-(4-amino-1-oxo-1,3-dihydro-isoindol-2-yl)-piperidine-2,6-dione suppresses the generation of inflammatory cytokines, down-regulates adhesion molecules and apoptosis inhibitory proteins (*e.g.*, cFLIP, cIAP), promotes sensitivity to death-receptor initiated programmed cell death, and suppresses angiogenic response. The studies show that 3-(4-amino-1-oxo-1,3-dihydro-isoindol-2-yl)-piperidine-2,6-dione abrogates mitogenic response to VEGF in AML cells by extinguishing ligant-induced Akt-phosphorylation, and selectively suppresses MDS vs normal bone marrow proginitor formation in pre-clinical models.

5.3. CLINICAL STUDIES IN MDS PATIENTS

Protocol

An immunomodulatory compound of the invention, such as 4-(amino)-2-(2,6-dioxo(3-piperidyl))-isoindoline-1,3-dione and 3-(4-amino-1-oxo-1,3-dihydro-isoindol-2-yl)-piperidine-2,6-dione, is administered in an amount of from about 0.1 to about 25 mg per day to patients with MDS for 16 weeks, who are subsequently evaluated for a hematological response. Response rates are assessed in cohorts stratified by the likelihood of an MDS subtype to transform to leukemia according to the International Prognostic Scoring System

(IPSS)-defined risk groups (*i.e.*, IPSS Low and Intermediate I; *versus* IPSS Intermediate II and High).

For example, fifteen patients are enrolled in the first cohort and receive treatment with 25 mg per day of 3-(4-amino-1-oxo-1,3-dihydro-isoindol-2-yl)-piperidine-2,6-dione. The number of patients who subsequently experience an erythroid response (major or minor response) by week 16 is evaluated. If no responses are observed, the study is terminated due to lack of efficacy. If, however, 4 or more patients respond, the study is terminated due to promising clinical activity. In the intermediate case (*e.g.*, 1, 2 or 3 patients respond), a second cohort of 10 patients is enrolled. If after the completion of treatment by the second cohort, 4 or more patients respond among the 25 patients treated, it is concluded that the 3-(4-amino-1-oxo-1,3-dihydro-isoindol-2-yl)-piperidine-2,6-dione shows promising clinical activity.

Clinical Study

Clinical studies were performed for the remitting potential of 3-(4-amino-1-oxo-1,3-dihydro-isoindol-2-yl)-piperidine-2,6-dione in MDS patients with red blood cell transfusion-dependence (>4 units/8 weeks) or symptomatic anemia (Hgb<10 g/dl). Patients received continuous treatment with 3-(4-amino-1-oxo-1,3-dihydro-isoindol-2-yl)-piperidine-2,6-dione at a oral dose of 25 mg daily. Responses were assessed according to IWG criteria after 16 weeks of treatments. Among 15 patients receiving the treatments, 11 patients were evaluable for toxicity, nine patients were evaluable for response (>8 wks therapy), and three patients discontinued the therapy prematurely (<2 weeks) due to cholecystitis, autoimmune hemolytic anemia, or patient refusal. Median age of the patients was 78 years ranging from 51 to 82 years. FAB types of the MDS patients include RA [4 patients], RARS [4 patients], RAEB [6 patients], and RAEB-T [1patient] with corresponding IPSS categories of Low/Int-1 in 11 patients and Int-2/High in four patients. Myelosuppression, which was characterized by higher than grade 3 common toxicity criteria or 50% decrease in leukocyte and platelet counts [9 patients], and grade 3 fatigue [1 patient], necessitated dose reduction to 10 mg in the initial ten patients. All subsequent patients initiated oral administrations with 10 mg daily. Grade 1,2 drug-related adverse effects were limited to the 25 mg dose and included pruritus or itchy scalp [6 patients] and myalgia [1 patient]. Six (66%) of nine evaluable patients experienced hematologic benefit (dual lineage, 1 patient), including 6/7 (86%) patients with IPSS Low/Int-1. Hematologic responses included RBC transfusion-independence [4 patients], decrease in RBC transfusions of more than 50% [1 patient], increase in Hgb of more than

1.5 g [1 patient], and one minor platelet response (increase of more than 30,000/μL). Among five patients evaluable for cytogenetic response, three patients achieved either a complete or partial (decrease in abnormal metaphases of more than 50%) remission. Responses were associated with normalization of blast percentage [1 patient], reduced grade of BM cytologic dysplasia, and 50% to more than 40 times improvement in BM multipotent progenitor (CFU-GEMM) and erythroid burst (BFU-E) formation. Correlation with changes in apoptotic index, angiogenic features (cellular/plasma VEGF, microvessel density), cytokine generation, and proliferative fraction (Ki67) are in progress. The results of this study indicate that 3-(4-amino-1-oxo-1,3-dihydro-isoindol-2-yl)-piperidine-2,6-dione has remarkable erythropoietic and cytogenetic remitting activity in patients with low/intermediate-1 risk MDS. Clinical benefit appears greatest in patients with low/intermediate-1 disease or the 5q-syndrome, associated with resolution of cytology dysplasia. The increase in apoptotic index, restoration of CFC, and suppression of karyotypic abnormalities suggest that the compound accelerates extinction of myelodysplastic clones. Based upon these data, the study has been expanded to treat additional subjects. Treatment with 10 mg as a continuous oral daily dose is well-tolerated with minimal myelosuppression.

Expanded Study

The clinical study was expanded with additional 16 MDS patients for at least eight weeks. According to the IPSS, 13 of these patients were categorized as low- or intermediate-1-risk patients and three patients were grouped as intermediate-2- or high-risk patients. According to the FAB classification, there were 11 patients with refractory anemia (RA) or RA with ringed sideroblasts (RARS), and five patients with RA with excess blasts (RAEB), RAEB in transformation (RAEB-T). The starting dose of 3-(4-amino-1-oxo-1,3-dihydro- isoindol-2-yl)-piperidine-2,6-dione was 25 mg daily for the first 13 patients and 10 mg daily for the remaining three patients. All patients receiving the starting dose of 25 mg required dose reduction by the completion of eight weeks therapy. Among these 16 patients who completed at least 8 weeks of monitoring, nine patients achieved an erythroid response as assessed by the International MDS Working Group Criteria. The erythroid responses consisted of transfusion independence in seven previously transfusion-dependent patients, a >2 g/dL rise in blood hemoglobin concentration in one patient in with transfusion-independent anemia, and a >50 % decrease in RBC transfusion requirement in one transfusion-dependent patient. Therefore, a major erythroid response developed in eight of 16 patients and a minor erythroid response was observed in one

patient. All of nine patients who showed erythroid response were low- or intermediate-1-risk patients. One patient also had a minor platelet response. In addition, complete cytogenetic responses developed in five in eight patients with abnormal karyotypes at baseline. These five patients with complete cytogenetic responses all had the Del5q31-33 abnormality, which has been discovered to be a good prognostic factor for MDS. Indeed, all five patients who enrolled in this study with 5q-syndrome achieved a complete cytogenetic response and a major erythroid response. The study also indicated an association of this therapy with an increased apoptotic index for myelodysplastic progenitors and recovery of normal hematopoietic progenitor cells.

5.4. CYCLING THERAPY IN MDS PATIENTS

As mentioned above, immunomodulatory compounds of the invention can be cyclically administered to patients with MDS. Cycling therapy involves the administration of a first agent for a period of time, followed by the administration of the agent and/or the second agent for a period of time and repeating this sequential administration. Cycling therapy can reduce the development of resistance to one or more of the therapies, avoid or reduce the side effects of one of the therapies, and/or improves the efficacy of the treatment.

Example 1

In a specific embodiment, prophylactic or therapeutic agents are administered in a cycle of about 16 weeks, about once or twice every day. One cycle can comprise the administration of a therapeutic on prophylactic agent and at least one (1), two (2), or three (3) weeks of rest. The number of cycles administered is from about 1 to about 12 cycles, more typically from about 2 to about 10 cycles, and more typically from about 2 to about 8 cycles.

Example 2

The objectives of the study are to evaluate the efficacy and safety of oral administration of 3-(4-amino-1-oxo-1,3-dihydro-isoindol-2-yl)-piperidine-2,6-dione in patients with MDS. Patients receive the compound in an amount of 10 mg/d or 15 mg/d for 21 days every 28 days in 4-week cycles for 16 weeks (4 cycles) or 24 weeks (6 cycles). The subject population comprises patients with low- or interemediate-1-risk MDS (International Prognostic Scoring System) with red blood cell transfusion-dependent anemia who have received at least two units of RBCs within 8 week of baseline (first day of study treatment).

In addition to hematological laboratory monitoring, bone marrow aspirates/biopsies with cytogenic analyses are obtained at baseline, after the completion of 3 cycles and after the completion of 6 cycles. The bone marrow, safety and efficacy data are reviewed to assess benefit-to-risk considerations throughout the study. The study reviews red blood cell transfusion independence and major erythroid response according to the International MDS Working Group Criteria. Further, the study observes red blood cell transfusion independence in the subgroup of patients with the 5q deletion cytogenetic abnormality; platelet, neutrophil, bone marrow and cytogenetic responses; and minor erythroid response of ≥50 % but <100 % reduction in red blood cell transfusion requirement over an 8 week period. The study further monitors adverse events, hematological tests, serum chemistries, TSH, urinalysis, urine or serum pregnancy tests, vital signs, ECG and physical examinations.

Example 3

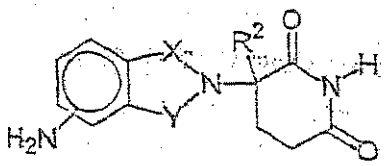
The objectives of the study are to compare the efficacy and safety of oral administration of 3-(4-amino-1-oxo-1,3-dihydro-isoindol-2-yl)-piperidine-2,6-dione to that of placebo plus standard care in patients with MDS. Patients receive the therapy in 4-week cycles for 16 weeks (4 cycles) or 24 weeks (6 cycles). The subject population comprise patients with low- or interemediate-1-risk MDS (International Prognostic Scoring System) with red blood cell transfusion-dependent anemia who have received at least two units of RBCs within 8 week of baseline (first day of study treatment). The study visits to assess safety and efficacy occur every 4 weeks and hematologic laboratory monitoring is performed every 2 weeks. Bone marrow aspirates/biopsies with cytogenetic analyses are obtained at baseline after the completion of 3 cycles and after the completion of 6 cycles. Bone marrow findings, safety and efficacy data are reviewed to assess benefit-to-risk considerations throughout the study. An extension study of continued treatments with the administration of the compound is available for patients who derive clinical benefit from 6 cycles of the therapy and to provide an opportunity for subjects who were randomized to placebo to cross over to the therapy.

Embodiments of the invention described herein are only a sampling of the scope of the invention. The full scope of the invention is better understood with reference to the attached claims.

CLAIMS

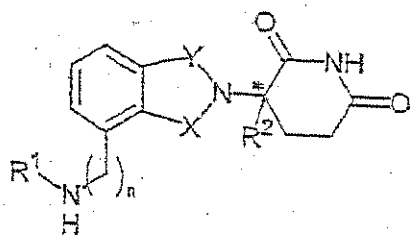
What is claimed is:

1. A use of therapeutically or prophylactically effective amount of an immunomodulatory compound, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof for manufacturing a medicament in treating or preventing a myelodysplastic syndrome to a patient in need thereof, wherein the immunomodulatory compound comprises 4-(amino)-2-(2,6-dioxo(3-piperidyl))- isoindoline-1,3-dione, 3 -(4-amino- 1 -oxo- 1,3 -dihydro-isoindol-2- yl)-piperidine-2, 6-dione, formula (I) and (II):



(I)

wherein one of X and Y is C=O, the other of X and Y is C=0 or CH_2, and R^2 is hydrogen or lower alkyl, formula (II):



(II)

wherein

one of X and Y is C=O and the other is CH_2 or C=O;

R^1 is H, (C_1-C_8)alkyl, (C_3-C_7)cycloalkyl, (C_2-C_8)alkenyl, (C_2-C_8)alkynyl, benzyl, aryl, (C_0-C_4)alkyl-(C_1-C_6)heterocycloalkyl, (C_0-C_4)alkyl-(C_2-C_5)heteroaryl, $C(O)R^3$, $C(S)R^3$,$C(O)OR^4$, (C_1-C_8)alkyl-$N(R^6)_2$, (C_1-C_8)alkyl-OR^5, (C_1-C_8)alkyl-$C(O)OR^5$, $C(O)NHR^3$, $C(S)NHR^3$, $C(O)NR^3R^{3'}$, $C(S)NR^3R^{3'}$ or (C_1-C_8)alkyl-$O(CO)R^5$;

R^2 is H, F, benzyl, $(C_1$-$C_8)$alkyl, $(C_2$-$C_8)$alkenyl, or $(C_2$-$C_8)$alkynyl;

R^3 and $R^{3'}$ are independently $(C_1$-$C_8)$alkyl, $(C_3$-$C_7)$cycloalkyl, $(C_2$-$C_8)$alkenyl, $(C_2$-$C_8)$alkynyl, benzyl, aryl, $(C_0$-$C_4)$alkyl-$(C_1$-$C_6)$heterocycloalkyl, $(C_0$-$C_4)$alkyl-$(C_2$-$C_5)$heteroaryl, $(C_0$-$C_8)$alkyl-$N(R^6)_2$, $(C_1$-$C_8)$alkyl-OR^5, $(C_1$-$C_8)$alkyl-$C(O)OR^5$, $(C_1$-$C_8)$alkyl-$O(CO)R^5$, or $C(O)OR^5$;

R^4 is $(C_1$-$C_8)$alkyl, $(C_2$-$C_8)$alkenyl, $(C_2$-$C_8)$alkynyl, $(C_1$-$C_4)$alkyl- OR^5, benzyl, aryl, $(C_0$-$C_4)$alkyl-$(C_1$-$C_6)$heterocycloalkyl, or $(C_0$-$C_4)$alkyl-$(C_2$-$C_5)$heteroaryl;

R^5 is $(C_1$-$C_8)$alkyl, $(C_2$-$C_8)$alkenyl, $(C_2$-$C_8)$alkynyl, benzyl, aryl, or $(C_2$-$C_5)$heteroaryl;

each occurrence of R^6 is independently H, $(C_1$-$C_8)$alkyl, $(C_2$-$C_8)$alkenyl, $(C_2$-$C_8)$alkynyl, benzyl, aryl, $(C_2$-$C_5)$heteroaryl, or $(C_0$-$C_8)$alkyl-$C(O)O$-R^5 or the R^6 groups join to form a heterocycloalkyl group;

n is 0 or 1; and

* represents a chiral-carbon center, a cyano or carboxy derivative of a substituted styrene, a 1-oxo-2-(2,6-dioxo-3 fluoropiperidin-3yl) isoindoline, a 1,3-dioxo-2-(2,6-dioxo-3-fluoropiperidine-3-yl) isoindoline, and a tetra substituted 2-(2,6-dioxopiperdin-3-yl)-1-oxoisoindoline.

2. A use of prophylactically effective amount of an immunomodulatory compound recited in claim 1, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof in the manufacture of a medicament for managing a myelodysplastic syndrome to a patient in need thereof.

3. A use of therapeutically or prophylactically effective amount of an immunomodulatory compound recited in claim 1, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof, and a therapeutically or prophylactically effective amount of at least one second active ingredient in the manufacture of a medicament for treating, preventing or managing a myelodysplastic syndrome to a patient in need thereof.

4. The use of claim 3, wherein the second active ingredient is capable of improving blood cell production.

5. The use of claim 3, wherein the second active ingredient is a cytokine, hematopoietic growth factor, anti-cancer agent, antibiotic, proteasome inhibitor, or immunosuppressive agent.

6. The use of claim 3, wherein the second active ingredient is etanercept, imatinib, anti-TNF-o' antibodies, infliximab, G-CSF, GM-CSF, EPO, topotecan, pentoxifylline, ciprofloxacin, irinotecan, vinblastine, dexamethasone, IL2, IL8, IL18, Ara C, vinorelbine, isotretinoin, 13-cis- retinoic acid, or a pharmacologically active mutant or derivative thereof.

7. The use of any one of claims 1 to 3, wherein the myelodysplastic syndrome is refractory anemia, refractory anemia with ringed sideroblasts, refractory anemia with excess blasts, refractory anemia with excess blasts in transformation, or chronic myelomonocytic leukemia.

8. The use of any one of claims 1 to 3, wherein the myelodysplastic syndrome is primary or secondary.

9. The use of any one of claims 1 to 3, wherein the stereoisomer of the immunomodulatory compound is an enantiomer.

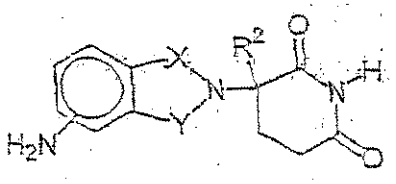
10. The use of any one of claims 1 to 3, wherein the immunomodulatory compound is 4-(amino)-2-(2,6-dioxo(3-piperidyl))- isoindoline-1,3-dione.

11. The use of claim 10, wherein the immunomodulatory compound is enantiomerically pure.

12. The use of any one of claims 1 to 3, wherein the immunomodulatory compound is 3 -(4-amino- 1 -oxo- 1,3 -dihydro-isoindol-2- yl)-piperidine-2, 6-dione.

13. The use of claim 12, wherein the immunomodulatory compound is enantiomerically pure.

14. The use of any one of claims 1 to 3, wherein the immunomodulatory compound has formula (I):

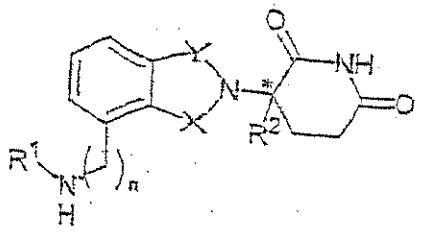


(I)

wherein one of X and Y is C=O, the other of X and Y is C=0 or CH_2, and R^2 is hydrogen or lower alkyl.

15. The use of claim 14, wherein the immunomodulatory compound is enantiomerically pure.

16. The use of any one of claims 1 to 3, wherein the immunomodulatory compound has formula (II):



(II)

wherein

one of X and Y is C=O and the other is CH_2 or C=O;

R^1 is H, $(C_1\text{-}C_8)$alkyl, $(C_3\text{-}C_7)$cycloalkyl, $(C_2\text{-}C_8)$alkenyl, $(C_2\text{-}C_8)$alkynyl, benzyl, aryl, $(C_0\text{-}C_4)$alkyl-$(C_1\text{-}C_6)$heterocycloalkyl, $(C_0\text{-}C_4)$alkyl-$(C_2\text{-}C_5)$heteroaryl, $C(O)R^3$, $C(S)R^3$, $C(O)OR^4$, $(C_1\text{-}C_8)$alkyl-$N(R^6)_2$, $(C_1\text{-}C_8)$alkyl-OR^5, $(C_1\text{-}C_8)$alkyl-$C(O)OR^5$, $C(O)NHR^3$, $C(S)NHR^3$, $C(O)NR^3R^{3'}$, $C(S)NR^3R^{3'}$ or $(C_1\text{-}C_8)$alkyl-$O(CO)R^5$;

R^2 is H, F, benzyl, $(C_1\text{-}C_8)$alkyl, $(C_2\text{-}C_8)$alkenyl, or $(C_2\text{-}C_8)$alkynyl;

R^3 and $R^{3'}$ are independently $(C_1\text{-}C_8)$alkyl, $(C_3\text{-}C_7)$cycloalkyl, $(C_2\text{-}C_8)$alkenyl, $(C_2\text{-}C_8)$alkynyl, benzyl, aryl, $(C_0\text{-}C_4)$alkyl-$(C_1\text{-}C_6)$heterocycloalkyl, $(C_0\text{-}C_4)$alkyl-$(C_2\text{-}C_5)$heteroaryl, $(C_0\text{-}C_8)$alkyl-$N(R^6)_2$, $(C_1\text{-}C_8)$alkyl-OR^5, $(C_1\text{-}C_8)$alkyl-$C(O)OR^5$, $(C_1\text{-}C_8)$alkyl-$O(CO)R^5$, or $C(O)OR^5$;

R^4 is $(C_1\text{-}C_8)$alkyl, $(C_2\text{-}C_8)$alkenyl, $(C_2\text{-}C_8)$alkynyl, $(C_1\text{-}C_4)$alkyl- OR^5, benzyl, aryl, $(C_0\text{-}C_4)$alkyl-$(C_1\text{-}C_6)$heterocycloalkyl, or $(C_0\text{-}C_4)$alkyl-$(C_2\text{-}C_5)$heteroaryl;

R^5 is $(C_1\text{-}C_8)$alkyl, $(C_2\text{-}C_8)$alkenyl, $(C_2\text{-}C_8)$alkynyl, benzyl, aryl, or $(C_2\text{-}C_5)$heteroaryl;

each occurrence of R^6 is independently H, $(C_1\text{-}C_8)$alkyl, $(C_2\text{-}C_8)$alkenyl, $(C_2\text{-}C_8)$alkynyl, benzyl, aryl, $(C_2\text{-}C_5)$heteroaryl, or $(C_0\text{-}C_8)$alkyl-$C(O)O\text{-}R^5$ or the R^6 groups join to form a heterocycloalkyl group;

n is 0 or 1; and

* represents a chiral-carbon center.

17. The use of claim 16, wherein the immunomodulatory compound is enantiomerically pure.

18. The use of any one of claims 1 to 3, wherein the immunomodulatory compound is a cyano or carboxy derivative of a substituted styrene, a 1- oxo-2-(2,6-dioxo-3 fluoropiperidin-3yl) isoindoline, a 1,3-dioxo-2-(2,6- dioxo-3-fluoropiperidine-3-yl) isoindoline, and a tetra substituted 2-(2, 6-dioxopiperdin-3-yl)-1- oxoisoindoline.

19. The use of claim 18, wherein the immunomodulatory compound is enantiomerically pure.

20. A use of therapeutically or prophylactically effective amount of an immunomodulatory compound recited in claim 1, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof, in the manufacture of a medicament for treating, preventing or managing myelodysplastic syndromes during, before or after transplanting umbilical cord blood, placental blood, peripheral blood stem cell, hematopoietic stem cell preparation or bone marrow in the patient in need thereof.

21. A use of amount of the second active ingredient and a therapeutically or prophylactically effective amount of an immunomodulatory compound recited in claim 1, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof in the manufacture of a medicament for reducing or avoiding an adverse effect associated with the administration of a second active ingredient in a patient suffering from a myelodysplastic syndrome.

22. The use of claim 21, wherein the second active ingredient is capable of improving blood cell production.

23. The use of claim 21, wherein the second active ingredient is a cytokine, hematopoletic growth factor, anti-cancer agent, antibiotic, proteasome inhibitor, or immunosuppressive agent.

24. The use of claim 21, wherein the second active ingredient is etanercept, imatinib, anti-TNF-ot antibodies, infliximab, G-CSF, GM- CSF, EPO, topotecan, pentoxifylline, ciprofloxacin, irinotecan, vinblastine, dexamethasone, IL2, IL8, IL18, Ara C, vinorelbine, isotretinoin, 13-cis- retinoic acid, or a pharmacologically active mutant or derivative thereof, or a combination thereof.

25. A pharmacoutical composition for treating, preventing or managing myelodysplastic syndromes comprising an immunomodulatory compound of claim 1, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof in an amount effective to treat, prevent or manage myelodysplastic syndromes, and a carrier.

26. A pharmaceutical composition for treating, preventing or managing myelodysplastic syndromes comprising an immunomodulatory compound of claim 1, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof, and a second active ingredient.

27. The pharmaceutical composition of claim 26, wherein the second active ingredient is capable of improving blood cell production.

28. The pharmaceutical composition of claim 26, wherein the second active ingredient is a cytokine, hematopoietic growth factor, anti- cancer agent, antibiotic, proteasome inhibitor, or immunosuppressive agent.

29. The pharmaceutical composition of claim 26, wherein the second active ingredient is etanercept, imatinib, anti-TNF-ot antibodies, infliximab, G- CSF7 GM-CSF, EPO, topotecan, pentoxifylline, ciprofloxacin, irinotecan, vinblastine, dexamethasone, IL2, IL8, ILLS, Ara-C, vinorelbine, isotretinoin, 13-cis-retinoic acid, or a pharmacologically active mutant or derivative thereof, or a combination thereof.

30. A single unit dosage form for treating, preventing or managing myelodysplastic syndromes comprising an immunornodulatory compound of claim 1, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof, and a second active ingredient capable of improving blood cell production.

31. A single unit dosage form for treating, preventing or managing myelodysplastic syndromes comprising an immunomodulatory compound of claim 1, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof, and a second active ingredient, wherein the second active ingredient is a cytokine, hematopoietic growth factor, anti-cancer agent, antibiotic, proteasome inhibitor, or immunosuppressive agent.

32. A single unit dosage form comprising an immunomodulatory compound of claim 1, or a pharmacoutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof, and etanercept, imatinib, anti-TNF-ot antibodies, infliximab, G-CSF, GM-CSF, EPO, topotecan, pentoxifylline, ciprofloxacin, irinotecan, vinblastine, dexamethasone, IL2, IL8, IL18, Ara-C, vinorelbine, isotretinoin, 13-cis-retinoic acid, or a pharmacologically active mutant or derivative thereof, or a combination thereof.

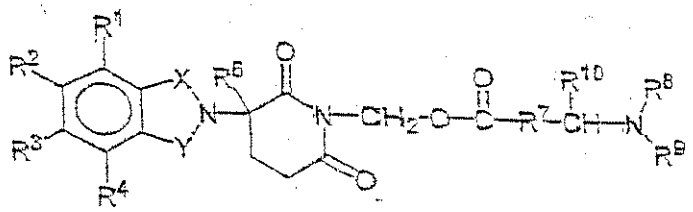
33. The single unit dosage forte to claim 30, 31 or 32, wherein the dosage form is suitable for intravenous or subcutaneous administration to a patient.

34. A kit comprising: a pharmaceutical composition for treating myelopdysplastic syndrome comprising an immunomodulatory compound of claim 1, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof; and umbilical cord blood, placental blood, peripheral blood stem cell, hematopoietic stem cell preparation or bone marrow.

35. A kit comprising: a pharmaceutical composition comprising an immunomodulatory compound of claim 1, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof; a pharmaceutical composition comprising a second active ingredient selected from the group consisting of a cytokine, hematopoietic growth factor, anti-cancer agent, antibiotic, proteasome inhibitor, and immunosuppressive agent; and umbilical cord blood, placental blood, peripheral blood stem cell, hematopoletic stem cell preparation or bone marrow.

36. The kit of claim 34 or 35 which further comprises a device for the administration of the pharmaceutical composition or the single unit dosage form.

37. Use of an immunomodulatory compound, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof for manufacturing a medicament for treating, preventing or managing a myelodysplastic syndrome, wherein said immunomodulatory compound is of formula:



wherein:

one of X and Y is C=O and the other of X and Y is C=O or CH_2;

(i) each of R^1, R^2, R^3, or R^4, independently of the others, is halo, alkyl of 1 to 4 carbon atoms, or alkoxy of 1 to 4 carbon atoms or (ii) one of R^1, R^2, R^3, and R^4 is $-NHR^5$ and the remaining of R^1, R^2, R^3, and R^4 are hydrogen;

R^5 is hydrogen or alkyl of 1 to 8 carbon atoms;

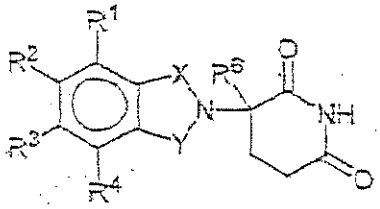
R^6 is hydrogen, alkyl of 1 to 8 carbon atoms, benzo, chloro, or fluoro;

R^7 is m-phenylene or p-phenylene or $-(C_nH_{2n})-$ in which n has a value of 0 to 4;

each of R^8 and R^9 taken independently of the other is hydrogen or alkyl of 1 to 8 carbon atoms, or R^8 and R^9 taken together are tetramethylene, pentamethylene, hexamethylene, or $-CH_2CH_2X^1CH_2CH_2-$ in which X^1 is -O-, -S-, or -NH-;

R^{10} is hydrogen, alkyl of to 8 carbon atoms, or phenyl.

38. Use of an immunomodulatory compound, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof for manufacturing a medicament for treating, preventing or managing a myelodysplastic syndrome, wherein said immunomodulatory compound is of formula:



in which

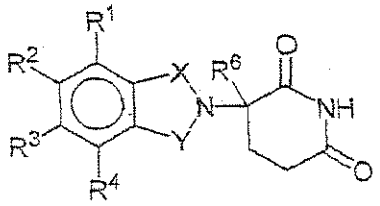
one of X and Y is C=O and the other of X and Y is C=O or CH_2;

each of R^1, R^2, R^3, and R^4, independently of the others, is halo, alkyl of 1 to 4 carbon atoms, or alkoxy of 1 to 4 carbon atoms or (ii) one of R^1, R^2, R^3, and R^4 is nitro or protected

amino and the remaining of R^1, R^2, R^3, and R^4 are hydrogen; and

R^6 is hydrogen, alkyl of 1 to 8 carbon atoms, benzo, chloro, or fluoro.

39. Use of an immunomodulatory compound, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof for manufacturing a medicament for treating, preventing or managing a myolodysplastic syndrome, wherein said immunomodulatory compound is of formula:



in which:

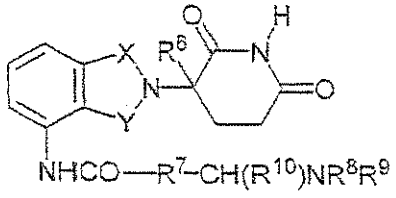
one of X and Y is C=O and the other of X and Y is C=O or CH_2;

(i) each of R^1, R^2, R^3, and R^4, independently of the others, is halo, alkyl of 1 to 4 carbon atoms, or alkoxy of 1 to 4 carbon atoms or (ii) one of R^1, R^2, R^3, and R^4 is $-NHR^5$ and the remaining of R^1, R^2, R^3, and R^4 are hydrogen;

R^5 is hydrogen, alkyl of 1 to 8 carbon atoms, or $CO-R^7-CH(R^{10})NR^8R^9$ in which each of R^7, R^8, R^9, and R^{10} is as herein defined; and

R^6 is alkyl of 1 to 8 carbon atoms, benzo, chloro, or fluoro.

40. Use of an immunomodulatory compound, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof for manufacturing a medicament for treating, preventing or managing a myolodysplastic syndrome, wherein said immunomodulatory compound is of formula:



in which:

one of X and Y is C=O and the other of X and Y is C=O or CH_2;

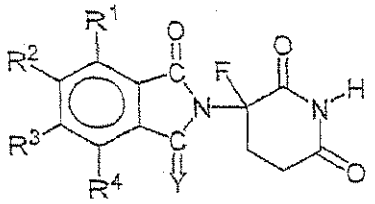
R^6 is hydrogen, alkyl of 1 to 8 carbon atoms, benzyl, chloro, or fluoro;

R^7 is m-phenylene, p-phenylene or $-(C_nH_{2n})-$ in which n has a value of 0 to 4;

each of R^8 and R^9 taken independently of the other is hydrogen or alkyl of 1 to 8 carbon atoms, or R^8 and R^9 taken together are tetramethylene, pentamethylene, hexamethylene, or $-CH_2CH_2X^1CH_2CH_2-$ in which X^1 is -O-, -S- or -NH-; and

R^{10} is hydrogen, alkyl of 1 to 8 carbon atoms, or phenyl.

41. Use of an immunomodulatory compound, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof for manufacturing a medicament for treating, preventing or managing a myolodysplastic syndrome, wherein said immunomodulatory compound is of formula:

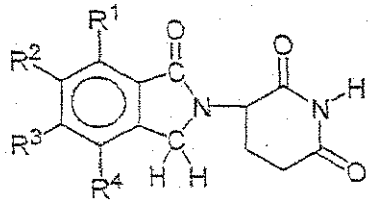


wherein Y is oxygen or H^2 and

each of R^1, R^2, R^3, and R^4, independently of the others, is hydrogen, halo, alkyl of 1 to 4 carbon atoms, alkoxy of 1 to 4 carbon atoms, or amino.

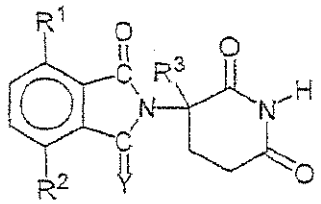
42. The use of claim 41, wherein said immunomodulatory compound is 1-oxo-2-(2,6-dioxo-3-fluoropiperidin-3yl) isoindolines or 1,3-dioxo-2-(2,6-dioxo-3-fluoropiperidine-3-yl) isoindolines.

43. Use of an immunomodulatory compound, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof for manufacturing a medicament for treating, preventing or managing a myolodysplastic syndrome, wherein said immunomodulatory compound is of formula:



wherein each of R^1, R^2, R^3, and R^4, independently of the others, is halo, alkyl of 1 to 4 carbon atoms, or alkoxy of 1 to 4 carbon atoms.

44. Use of an immunomodulatory compound, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof for manufacturing a medicament for treating, preventing or managing a myolodysplastic syndrome, wherein said immunomodulatory compound is of formula:



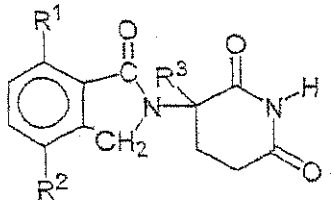
in which

Y is oxygen or H_2,

a first of R^1 and R^2 is halo, alkyl, alkoxy, alkylamino, dialkylamino, cyano, or carbamoyl, the second of R^1 and R^2, independently of the first, is hydrogen, halo, alkyl, alkoxy, alkylamino, dialkylamino, cyano, or carbamoyl, and

R^3 is hydrogen, alkyl, or benzyl.

45. Use of an immunomodulatory compound, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof for manufacturing a medicament for treating, preventing or managing a myolodysplastic syndrome, wherein said immunomodulatory compound is of formula:

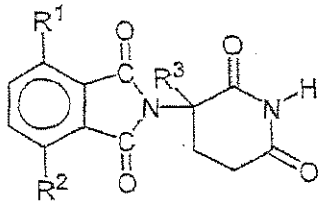


wherein

a first of R^1 and R^2 is halo, alkyl of from 1 to 4 carbon atoms, alkoxy of from 1 to 4 carbon atoms, dialkylamino in which each alkyl is of from 1 to 4 carbon atoms, cyano, or carbamoyl; the second of R^1 and R^2, independently of the first, is hydrogen, halo, alkyl of from 1 to 4 carbon atoms, alkoxy of from 1 to 4 carbon atoms, alkylamino in which alkyl is of from 1 to 4 carbon atoms, dialkylamino in which each alkyl is of from 1 to 4 carbon atoms, cyano, or carbamoyl, and

R^3 is hydrogen, alkyl of from 1 to 4 carbon atoms, or benzyl.

46. Use of an immunomodulatory compound, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof for manufacturing a medicament for treating, preventing or managing a myolodysplastic syndrome, wherein said immunomodulatory compound is of formula:



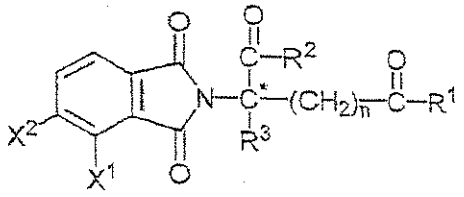
wherein

a first of R^1 and R^2 is halo, alkyl of from 1 to 4 carbon atoms, alkoxy of from 1 to 4 carbon atoms, dialkylamino in which each alkyl is of from 1 to 4 carbon atoms, cyano, or carbamoyl, the second of R^1 and R^2, independently of the first, is hydrogen, halo, alkyl of from 1 to 4 carbon atoms, alkoxy of from 1 to 4 carbon atoms, alkylamino in which alkyl is of from 1 to

4 carbon atoms, dialkylamino in which each alkyl is of from 1 to 4 carbon atoms, cyano, or carbamoyl, and

R^3 is hydrogen, alkyl of from 1 to 4 carbon atoms, or benzyl.

47. Use of an immunomodulatory compound, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof for manufacturing a medicament for treating, preventing or managing a myolodysplastic syndrome, wherein said immunomodulatory compound is of formula:



in which

the carbon atom designated C* constitutes a center of chirality (when n is not zero and R^1 is not the same as R^2);

one of X^1 and X^2 is amino, nitro, alkyl of one to six carbons, or NH-Z, and the other of X^1 or X^2 is hydrogen;

each of R^1 and R^2 independent of the other, is hydroxy or NH-Z;

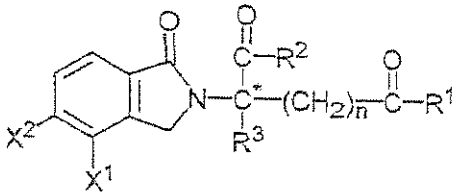
R^3 is hydrogen, alkyl of one to six carbons, halo, or haloalkyl;

Z is hydrogen, aryl, alkyl of one to six carbons, formyl, or acyl of one to six carbons; and

n has a value of 0, 1, or 2;

provided that if X^1 is amino, and n is 1 or 2, then R^1 and R^2 are not both hydroxy; and the salts thereof.

48. Use of an immunomodulatory compound, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof for manufacturing a medicament for treating, preventing or managing a myolodysplastic syndrome, wherein said immunomodulatory compound is of formula:



in which

the carbon atom designated C* constitutes a center of chirality when n is not zero and R^1 is not R^2;

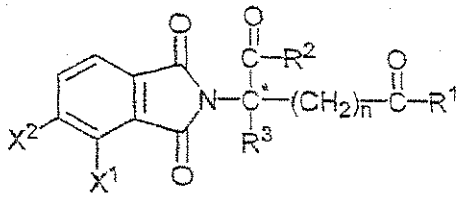
one of X^1 and X^2 is amino, nitro, alkyl of one to six carbons, or NH-Z, and the other of X^1 or X^2 is hydrogen;

each of R^1 and R^2 independent of the other, is hydroxy or NH-Z;

R^3 is alkyl of one to six carbons, halo, or hydrogen;

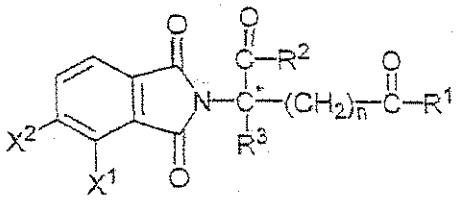
Z is hydrogen, aryl or an alkyl or acyl of one to six carbons; and
n has a value of 0, 1, or 2.

49. Use of an immunomodulatory compound, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof for manufacturing a medicament for treating, preventing or managing a myolodysplastic syndrome, wherein said immunomodulatory compound is of formula:



in which
the carbon atom designated C* constitutes a center of chirality when n is not zero and R^1 is not R^2;
one of X^1 and X^2 is amino, nitro, alkyl of one to six carbons, or NH-Z, and the other of X^1 or X^2 is hydrogen;
each of R^1 and R^2 independent of the other, is hydroxy or NH-Z;
R^3 is alkyl of one to six carbons, halo, or hydrogen;
Z is hydrogen, aryl, or an alkyl or acyl of one to six carbons; and
n has a value of 0, 1, or 2;
and the salts thereof.

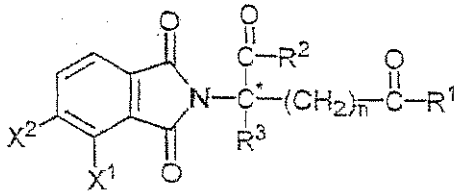
50. Use of an immunomodulatory compound, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof for manufacturing a medicament for treating, preventing or managing a myolodysplastic syndrome, wherein said immunomodulatory compound is of formula:



wherein
one of X^1 and X^2 is nitro, or NH-Z, and the other of X^1 or X^2 is hydrogen;
each of R^1 and R^2, independent of the other, is hydroxy or NH-Z;
R^3 is alkyl of one to six carbons, halo, or hydrogen;
Z is hydrogen, phenyl, an acyl of one to six carbons, or an alkyl of one to six carbons; and
n has a value of 0, 1, or 2;
provided that if one of X^1 and X^2 is nitro, and n is 1 or 2, then R^1 and R^2 are other than hydroxy; and

if -COR^1 and -$(CH_2)_nCOR^2$ are different, the carbon atom designated C^* constitutes a center of chirality.

51. Use of an immunomodulatory compound, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof for manufacturing a medicament for treating, preventing or managing a myolodysplastic syndrome, wherein said immunomodulatory compound is of formula:



wherein

one of X^1 and X^2 is alkyl of one to six carbons;

each of R^1 and R^2, independent of the other, is hydroxy or NH-Z;

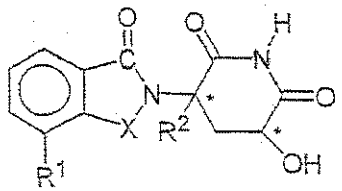
R^3 is alkyl of one to six carbons, halo, or hydrogen;

Z is hydrogen, phenyl, an acyl of one to six carbons, or an alkyl of one to six carbons; and

n has a value of 0, 1, or 2; and

if -COR^1 and -$(CH_2)_nCOR^2$ are different, the carbon atom designated C^* constitutes a center of chirality.

52. Use of an immunomodulatory compound, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof for manufacturing a medicament for treating, preventing or managing a myolodysplastic syndrome, wherein said immunomodulatory compound is of formula:



wherein:

the carbon atoms designated * constitute centers of chirality;

X is -C(O)- or -CH_2-;

R^1 is alkyl of 1 to 8 carbon atoms or -NHR^3;

R^2 is hydrogen, alkyl of 1 to 8 carbon atoms, or halogen;and

R^3 is hydrogen, alkyl of 1 to 8 carbon atoms, unsubstituted or substituted with alkoxy of 1 to 8 carbon atoms, halo, amino, or alkylamino of 1 to 4 carbon atoms, cycloalkyl of 3 to 18 carbon atoms, phenyl, unsubstituted or substituted with alkyl of 1 to 8 carbon atoms, alkoxy of 1 to 8 carbon atoms, halo, amino, or alkylamino of 1 to 4 carbon atoms, benzyl,

unsubstituted or substituted with alkyl of 1 to 8 carbon atoms, alkoxy of 1 to 8 carbon atoms, halo, amino, or alkylamino of 1 to 4 carbon atoms, or $-COR^4$ in which

R^4 is hydrogen, alkyl of 1 to 8 carbon atoms, unsubstituted or substituted with alkoxy of 1 to 8 carbon atoms, halo, amino, or alkylamino of 1 to 4 carbon atoms, cycloalkyl of 3 to 18 carbon atoms, phenyl, unsubstituted or substituted with alkyl of 1 to 8 carbon atoms, alkoxy of 1 to 8 carbon atoms, halo, amino, or alkylamino of 1 to 4 carbon atoms, or benzyl, unsubstituted or substituted with alkyl of 1 to 8 carbon atoms, alkoxy of 1 to 8 carbon atoms, halo, amino, or alkylamino of 1 to 4 carbon atoms.

53. Use of an immunomodulatory compound, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof for manufacturing a medicament for treating, preventing or managing a myolodysplastic syndrome, wherein said immunomodulatory compound is selected from the group consisting of: [2-(2,6-dioxo-piperidin-3-yl)-1,3-dioxo-2,3-dihydro-1H-isoindol-4-ylmethyl]-amide; (2-(2,6-dioxo-piperidin-3-yl)-1,3-dioxo-2,3-dihydro-1*H*-isoindol-4-ylmethyl)-carbamic acid *tert*-butyl ester; 4-(aminomethyl)-2-(2,6-dioxo(3-piperidyl))-isoindoline-1,3-dione; *N*-(2-(2,6-dioxo-piperidin-3-yl)-1,3-dioxo-2,3-dihydro-1*H*-isoindol-4-ylmethyl)-acetamide; *N*-{(2-(2,6-dioxo(3-piperidyl)-1,3-dioxoisoindolin-4-yl)methyl}cyclopropyl-carboxamide; 2-chloro-*N*-{(2-(2,6-dioxo(3-piperidyl))-1,3-dioxoisoindolin-4-yl)methyl}acetamide; *N*-(2-(2,6-dioxo(3-piperidyl))-1,3-dioxoisoindolin-4-yl)-3-pyridylcarboxamide; 3-{1-oxo-4-(benzylamino)isoindolin-2-yl}piperidine-2,6-dione; 2-(2,6-dioxo(3-piperidyl))-4-(benzylamino)isoindoline-1,3-dione; *N*-{(2-(2,6-dioxo(3-piperidyl))-1,3-dioxoisoindolin-4-yl)methyl}propanamide; *N*-{(2-(2,6-dioxo(3-piperidyl))-1,3-dioxoisoindolin-4-yl)methyl}-3-pyridylcarboxamide; *N*-{(2-(2,6-dioxo(3-piperidyl))-1,3-dioxoisoindolin-4-yl)methyl}heptanamide; *N*-{(2-(2,6-dioxo(3-piperidyl))-1,3-dioxoisoindolin-4-yl)methyl}-2-furylcarboxamide; {N-(2-(2,6-dioxo(3-piperidyl))-1,3-dioxoisoindolin-4-yl)carbamoyl}methyl acetate; *N*-(2-(2,6-dioxo(3-piperidyl))-1,3-dioxoisoindolin-4-yl)pentanamide; *N*-(2-(2,6-dioxo(3-piperidyl))-1,3-dioxoisoindolin-4-yl)-2-thienylcarboxamide; N-{[2-(2,6-dioxo(3-piperidyl))-1,3-dioxoisoindolin-4-yl] methyl}(butylamino)carboxamide; N-{[2-(2,6-dioxo(3-piperidyl))-1,3-dioxoisoindolin-4-yl] methyl}(octylamino)carboxamide; and N-{[2-(2,6-dioxo(3-piperidyl))-1,3-dioxoisoindolin-4-yl] methyl}(benzylamino)carboxamide; 1-oxo-2-(2,6-dioxopiperidin-3-yl) isoindolines or 1,3-dioxo-2-(2,6-dioxopiperidin-3-yl) isoindolines; isoindoline-1-one substituted in the 2-position with 2,6-dioxo-3-hydroxypiperidin-5-yl; or isoindoline-1,3-dione substituted in the 2-position with 2,6-dioxo-3-hydroxypiperidin-5-yl.

54. The use of any one of claims 37 to 53, wherein the medicament further comprises at least one second active ingredient.

55. The use of claim 54, wherein the second active ingredient is capable of improving blood cell production.

56. The use of claim 54, wherein the second active ingredient is a cytokine, hematopoietic growth factor, anti-cancer agent, antibiotic, proteasome inhibitor, or immunosuppressive agent.

57. The use of claim 54, wherein the second active ingredient is etanercept, imatinib, anti-TNF-α antibodies, infliximab, G-CSF, GM-CSF, EPO, topotecan, pentoxifylline, ciprofloxacin, irinotecan, vinblastine, dexamethasone, IL2, IL8, IL18, Ara-C, vinorelbine, isotretinoin, 13-cis-retinoic acid, or a phannacologically active mutant or derivative thereof, or a combination thereof.

58. The use of any one of claims 37 to 53, wherein the myelodysplastic syndrome is refractory anemia, refractory anemia with ringed sideroblasts, refractory anemia with excess blasts, refractory anemia with excess blasts in transformation, or chronic myelomonocytic leukemia.

59. The use of any one of claims 37 to 53, wherein the myelodysplastic syndrome is primary or secondary.

60. The use of any one of claims 37 to 53, wherein the stereoisomer of the immunomodulatory compound is an enantiomerically pure.

61. The use of any one of claims 37 to 53, wherein the medicament is administrated to a patient in need of treating, preventing or managing a myelodysplastic syndrome before, during or after transplanting umbilical cord blood, placental blood, peripheral blood stem cell, hematopoietic stem cell preparation or bone marrow in the patient.

62. Use of an immunomodulatory compound, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof and a second active ingredient for manufacturing a medicament for reducing or avoiding an adverse effect associated with the administration of the second active ingredient in a patient in need of treating, preventing or managing a myelodysplastic syndrome, wherein said immunomodulatory compound is the compound in any one of claims 37 to 53.

63. The use of claim 62, wherein the second active ingredient is capable of improving blood cell production.

64. The use of claim 62, wherein the second active ingredient is a cytokine, hematopoietic growth factor, anti-cancer agent, antibiotic, proteasome inhibitor, or immunosuppressive agent.

65. The use of claim 62, wherein the second active ingredient is etanercept, imatinib, anti-

TNF-a antibodies, infliximab, G-CSF, GM-CSF, EPO, topotecan, pentoxifylline, ciprofloxacin, irinotecan, vinblastine, dexamethasone, IL2, IL8, IL18, Ara- C, vinorelbine, isotretinoin, 13-cis-retinoic acid, or a pharmacologically active mutant or derivative thereof, or a combination thereof.

66. A pharmaceutical composition for treating myelodysplastic syndromes, comprising an immunomodulatory compound, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof in an amount effective to treat, prevent or manage myelodysplastic syndromes, and a carrier, wherein said immunomodulatory compound is the compound in any one of claims 37 to 53.

67. The pharmaceutical composition of claim 66, wherein the amount is about 0.1mg to 200mg.

68. The pharmaceutical composition of claim 66, further comprising at least one second active ingredient.

69. The pharmaceutical composition of claim 68, wherein the second active ingredient is capable of improving blood cell production.

70. The pharmaceutical composition of claim 68, wherein the second active ingredient is a cytokine, hematopoietic growth factor, anti-cancer agent, antibiotic, proteasome inhibitor, or immunosuppressive agent.

71. The pharmaceutical composition of claim 68, wherein the second active ingredient is etanercept, imatinib, anti-TNF- antibodies, infliximab, G-CSF, GM-CSF, EPO, topotecan, pentoxifylline, ciprofloxacin, irinotecan, vinblastine, dexamethasone, IL2, IL8, IL18, Ara-C, vinorelbine, isotretinoin, 13-cis-retinoic acid, or a pharmacologically active mutant or derivative thereof, or a combination thereof.

72. The pharmaceutical composition of claim 66, wherein the pharmaceutical composition is in a single unit dosage form suitable for oral, mucosal, parenteral, transdermal or transcutaneous administration.

73. The pharmaceutical composition of claim 72, wherein the single unit dosage form is tablets, caplets, capsules, cachets, troches, lozenges, dispersions, suppositories, powders, aerosols, gels, suspensions, solutions, elixirs, or sterile solids.

74. The pharmaceutical composition of claim 72, wherein the single unit dosage form is rapid release dosage forms or delayed release dosage form.

75. The pharmaceutical composition of claim 72, wherein the amount of the

immunomodulatory compound or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof is about 0.1mg, 1mg, 5mg, 7.5mg, 10mg, 12.5mg, 15mg, 17.5mg, 20mg, 25mg, 50mg, 100mg, 150mg or 200mg.

76. A kit comprising:

a pharmaceutical composition comprising an immunomodulatory compound, or a pharmaceutically acceptable salt, solvate, hydrate, stereoisomer, clathrate, or prodrug thereof, wherein said immunomodulatory compound is the compound in any one of claims 37 to 53; and

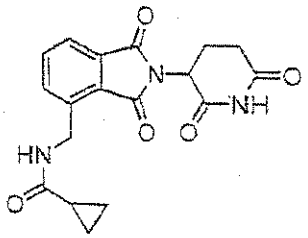
umbilical cord blood, placental blood, peripheral blood stem cell, hematopoietic stem cell preparation or bone marrow.

77. The kit of claim 76, further comprising:

a pharmaceutical composition comprising a second active ingredient selected from the group consisting of a cytokine, hematopoietic growth factor, anti-cancer agent, antibiotic, proteasome inhibitor, and immunosuppressive agent.

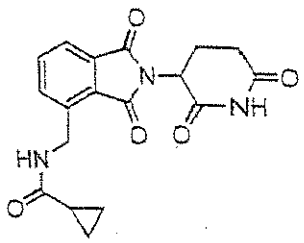
78. The kit of claim 76 or 77 which further comprises a device for the administration of the pharmaceutically composition or the single unit dosage form.

79. Use of N-{[2-(2,6-dioxo(3-piperidyl)-1,3-dioxoisoindolin-4-yl]methyl}cyclopropyl-carboxamide having the formula:



or a pharmaceutically acceptable salt, solvate or stereoisomer thereof in manufacturing a medicament for treating a myelodysplastic syndrome, wherein said medicament is in a daily dosage form comprising said compound in an amount of about 5mg to about 100 mg.

80. The use of claim 79, wherein the compound is N-{[2-(2,6-dioxo(3-piperidyl)-1,3-dioxoisoindolin-4-yl]methyl}cyclopropyl-carboxamide having the formula:



, as a free base.

81. The use of claim 79, wherein the compound is a pharmaceutically acceptable salt.

82. The use of claim 79, wherein the compound is a pharmaceutically acceptable solvate.

83. The use of claim 79, wherein the compound is a pharmaceutically acceptable stereoisomer.

84. The use of claim 83, wherein the stereoisomer is an enantiomerically pure R isomer.

85. The use of claim 83, wherein the stereoisomer is an enantiomerically pure S isomer.

86. The use of claim 79, wherein said medicament which further comprises a therapeutically effective amount of a second active agent.

87. The use of claim 86, wherein the second active agent is capable of improving blood cell production.

88. The use of claim 86, wherein the second active agent is a cytokine, hematopoietic growth factor, an anti-cancer agent, an antibiotic, a proteasome inhibitor, or an immunosuppressive agent.

89. The use of claim 88, wherein the second active agent is etanercept, imatinib, anti-TNF-α antibodies, infliximab, G-CSF, GM-CSF, EPO, topotecan, pentoxifylline,

ciprofloxacin, irinotecan, vinblastine, dexamethasone, IL2, IL8, IL18, Ara-C, vinorelbine, isotretinoin, 13-cis-retinoic acid, arsenic trioxide or a pharmacologically active mutant or derivative thereof.

90. The use of claim 79, wherein the myelodysplastic syndrome is refractory anemia, refractory anemia with ringed sideroblasts, refractory anemia with excess blasts, refractory anemia with excess blasts in transformation, or chronic myelomonocytic leukemia.

91. The use of claim 79, wherein the compound or a pharmaceutically acceptable salt, solvate or stereoisomer thereof is administered before, during or after transplanting umbilical cord blood, placental blood, peripheral blood stem cell, hematopoietic stem cell preparation or bone marrow in the patient.

92. The use of claim 89, wherein the second active agent is dexamethasone.

93. The use of claim 79, wherein the medicament is administrated to a patient in need of treating for a myelodysplastic syndrome, and said patient is not previously treated for a myelodysplastic syndrome.

94. The use of claim 79, wherein the medicament is administrated to a patient who has been previously treated for a myelodysplastic syndrome.

95. The use of claim 79, wherein the medicament is in a dosage form suitable for oral administration.

96. The use of claim 95, wherein the medicament is administered in the form of a capsule or tablet.

97. The use of claim 79, wherein the medicament is administered cyclically.

98. The use of claim 97, wherein the medicament is administered once or twice every day for sixteen or twenty-four weeks.

99. The use of claim 97, wherein in one cycle the medicament is administrated with at least one, two, or three weeks of rest.

100. The use of claim 97, wherein the medicament is administrated from one to twelve cycles.

101. The use of claim 97, wherein the medicament is in a daily dosage form comprising said compound in an amount of from about 5 to about 50 mg, and said medicament is administrated per day for 21 days every 28 days for sixteen or twenty-four weeks.

102. The use of claim 82, wherein said solvate is a hydrate.

103. The use of claim 79, wherein the medicament is in a daily dosage form comprising said compound in an amount of from about 5 mg to about 50 mg.

104. The use of claim 79, wherein the medicament is in a daily dosage form comprising about 5 mg of said compound.

105. The use of claim 79, wherein the medicament is in a daily dosage form comprising about 10 mg of said compound.

106. The use of claim 79, wherein the medicament is in a daily dosage form comprising about 20 mg of said compound.

107. The use of claim 79, wherein the medicament is in a daily dosage form comprising about 25 mg of said compound.

108. The use of claim 79, wherein the medicament is in a daily dosage form comprising about 50 mg of said compound.

109. The use of claim 79, wherein the medicament is in a daily dosage form comprising said compound in an amount of from about 25 mg to about 50 mg, and said medicament is administrated every other day.

110. The use of claim 79, wherein the medicament is administered in a cycle of about 16 weeks and about once or twice every day.

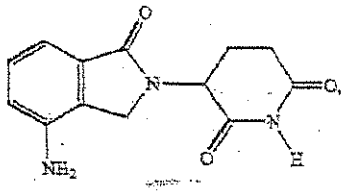
111. The use of claim 110, wherein said cycle comprises at least one (1), two (2), or three (3) weeks of rest.

112. The use of claim 79, wherein the medicament is in a daily dosage form comprising said compound in an amount of about 20 mg or about 50 mg, and said medicament is administrated per day for 21 days out of a block of 28 days.

113. The use of claim 79, wherein the medicament is in a daily dosage form of capsule comprising about 5 mg of said compound.

114. The use of claim 79, wherein the medicament is in the dosage form of capsule comprising about 25 mg.

115. Use of from about 0.1 to about 10 mg per of day of 3-(4-amino-1-oxo-1,3-dihydro-isoindol-2-yl)-piperidine-2,6-dione having the formula:



or a pharmaceutically acceptable salt, solvate or stereoisomer thereof for the manufacturing of a medicament for treatment of a myelodysplastic syndrome.

116. The use of claim 115, wherein the compound is a pharmaceutically acceptable salt.

117. The use of claim 115, wherein the compound is a pharmaceutically acceptable solvate.

118. The use of claim 115, wherein the compound is a pharmaceutically acceptable stereoisomer.

119. The use of claim 118, wherein the stereoisomer is an enantiomerically pure R isomer.

120. The use of claim 118, wherein the stereoisomer is an enantiomerically pure S isomer.

121. The use of claim 115, which further comprises administering a therapeutically effective amount of a second active agent.

122. The use of claim 121, wherein the second active agent is capable of improving blood cell production.

123. The use of claim 121, wherein the second active agent is a cytokine, hematopoietic growth factor, an anti-cancer agent, an antibiotic, a proteasome

inhibitor, or an immunosuppressive agent.

124. The use of claim 123, wherein the second active agent is etanercept, imatinib, anti-TNF-.alpha. antibodies, infliximab, G-CSF, GM-CSF, EPO, topotecan, pentoxifylline, ciprofloxacin, irinotecan, vinblastine, dexamethasone, IL2, IL8, IL18, Ara-C, vinorelbine, isotretinoin, 13-cis-retinoic acid, arsenic trioxide or a pharmacologically active mutant or derivative thereof.

125. The use of claim 115, wherein the myelodysplastic syndrome is refractory anemia, refractory anemia with ringed sideroblasts, refractory anemia with excess blasts, refractory anemia with excess blasts in transformation, or chronic myelomonocytic leukemia.

126. The use of claim 115, wherein the compound or a pharmaceutically acceptable salt, solvate or stereoisomer thereof is administered before, during or after transplanting umbilical cord blood, placental blood, peripheral blood stem cell, hematopoietic stem cell preparation or bone marrow in the patient.

127. The use of claim 124, wherein the second active agent is dexamethasone.

128. The use of claim 115, wherein the patient is not previously treated for a myelodysplastic syndrome.

129. The use of claim 115, wherein the patient has been previously treated for a myelodysplastic syndrome.

130. The use of claim 115, wherein the compound is administered orally.

131. The use of claim 130, wherein the compound is administered in the form of a capsule or tablet.

132. The use of claim 115, wherein the compound is administered cyclically.

133. The use of claim 132, wherein the compound is administered once or twice every day for sixteen or twenty-four weeks.

134. The use of claim 132 wherein one cycle comprises the administration of the compound and at least one, two, or three weeks of rest.

135. The use of claim 132, wherein the number of cycle is from one to twelve cycles.

136. The use of claim 132, wherein 3-(4-amino-1-oxo-1,3-dihydro-isoindol-2-yl)-piperidine-2,6-dione is administered in an amount of from about 5 to about 25 mg per day for 21 days every 28 days for sixteen or twenty-four weeks.

137. The use of claim 117 wherein said solvate is a hydrate.

138. The use of claim 115, wherein the compound is administered in an amount of from about 5 mg per day to about 25 mg per day.

139. The use of claim 115, wherein the compound is administered in an amount of 10 mg per day.

140. The use of claim 115, wherein the compound is administered in an amount of 15 mg per day.

141. The use of claim 115, wherein the compound is administered in an amount of 25 mg per day.

142. The use of claim 115, wherein the compound is administered in an amount of from about 25 mg every other day to about 50 mg every other day.

143. The use of claim 115, wherein the compound is administered in a cycle of about 16 weeks and about once or twice every day.

144. The use of claim 143, wherein said cycle comprises at least one (1), two

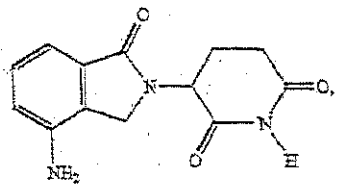
(2), or three (3) weeks of rest.

145. The use of claim 115, wherein the compound is administered in an amount of 10 mg per day or 15 mg per day for 21 days out of a block of 28 days.

146. The use of claim 115, wherein the compound is administered orally in an amount of 5 mg as a capsule per day.

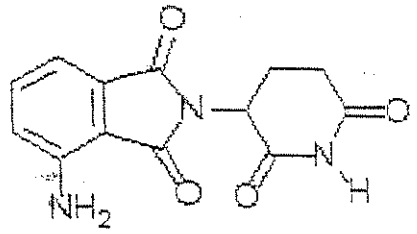
147. The use of claim 115, wherein the compound is administered orally in an amount of 10 mg as a capsule per day.

148. The use of claim 115, wherein the compound is 3-(4-amino- 1-oxo-1,3-dihydro-isoindol-2-yl)-piperidine-2,6-dione, having the formula:



as a free base.

149. Use of from about 0.1 to about 10 mg per of day of 4-(amino)-2-(2,6-dioxo(3-piperidyl))-isoindoline-1,3-dione having the formula:



or a pharmaceutically acceptable salt, solvate or stereoisomer thereof for the manufacturing of a medicament for treatment of a myelodysplastic syndrome.

150. The use of claim 149, wherein the compound is a pharmaceutically acceptable salt.

151. The use of claim 149, wherein the compound is a pharmaceutically acceptable solvate.

152. The use of claim 149, wherein the compound is a pharmaceutically acceptable stereoisomer.

153. The use of claim 152, wherein the stereoisomer is an enantiomerically pure R isomer.

154. The use of claim 152, wherein the stereoisomer is an enantiomerically pure S isomer.

155. The use of claim 149, which further comprises administering a therapeutically effective amount of a second active agent.

156. The use of claim 155, wherein the second active agent is capable of improving blood cell production.

157. The use of claim 155, wherein the second active agent is a cytokine, hematopoietic growth factor, an anti-cancer agent, an antibiotic, a proteasome

inhibitor, or an immunosuppressive agent.

158. The use of claim 157, wherein the second active agent is etanercept, imatinib, anti-TNF-.alpha. antibodies, infliximab, G-CSF, GM-CSF, EPO, topotecan, pentoxifylline, ciprofloxacin, irinotecan, vinblastine, dexamethasone, IL2, IL8, IL18, Ara-C, vinorelbine, isotretinoin, 13-cis-retinoic acid, arsenic trioxide or a pharmacologically active mutant or derivative thereof.

159. The use of claim 149, wherein the myelodysplastic syndrome is refractory anemia, refractory anemia with ringed sideroblasts, refractory anemia with excess blasts, refractory anemia with excess blasts in transformation, or chronic myelomonocytic leukemia.

160. The use of claim 149, wherein the compound or a pharmaceutically acceptable salt, solvate or stereoisomer thereof is administered before, during or after transplanting umbilical cord blood, placental blood, peripheral blood stem cell, hematopoietic stem cell preparation or bone marrow in the patient.

161. The use of claim 158, wherein the second active agent is dexamethasone.

162. The use of claim 149, wherein the patient is not previously treated for a myelodysplastic syndrome.

163. The use of claim 149, wherein the patient has been previously treated for a myelodysplastic syndrome.

164. The use of claim 149, wherein the compound is administered orally.

165. The use of claim 164, wherein the compound is administered in the form of a capsule or tablet.

166. The use of claim 149, wherein the compound is administered cyclically.

167. The use of claim 166, wherein the compound is administered once or twice every day for sixteen or twenty-four weeks.

168. The use of claim 166 wherein one cycle comprises the administration of the compound and at least one, two, or three weeks of rest.

169. The use of claim 166, wherein the number of cycle is from one to twelve cycles.

170. The use of claim 166, wherein 4-(amino)-2-(2,6-dioxo(3-piperidyl))-isoindoline-1,3-dione is administered in an amount of from about 0.1 to about 10 mg per day for 21 days every 28 days for sixteen or twenty-four weeks.

171. The use of claim 151 wherein said solvate is a hydrate.