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(54) Title: TREATMENT OF INFLAMMATORY DISORDERS OF THE BOWEL AND URINARY BLADDER

(57) **Abrégé/Abstract:**

A method of treating inflammatory disorders of the bowel and inflammatory disorders of the urinary bladder is disclosed. A method of modulating the function of monocytes is also disclosed. The method involves the treatment of a patient's blood with a photoactivatable compound followed by ultraviolet light-activation of the photoactivatable compound. The blood treated as such is returned to the patient in a process known as extracorporeal photopheresis.



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(21) International Application Number: PCT/US98/14118 (22) International Filing Date: 8 July 1998 (08.07.98) (30) Priority Data: 60/052,101 10 July 1997 (10.07.97) US (71) Applicant: THERAKOS, INC. [US/US]; 437 Creamery Way, Exton, PA 19341 (US). (72) Inventors: ZELDIS, Jerome; 437 Creamery Way, Exton, PA 19341 (US). MCLAUGHLIN, Susan; 437 Creamery Way, Exton, PA 19341 (US). STOUCH, Bruce; 437 Creamery Way, Exton, PA 19341 (US). KNOBLER, Robert; 33 W. Putnam Avenue, Greenwich, CT 06832 (US). (74) Agents: CIAMPORCERO, Audley, A., Jr. et al.; Johnson & Johnson, One Johnson & Johnson Plaza, New Brunswick, NJ 08933 (US).		(81) Designated States: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, GM, HR, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, UZ, VN, YU, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG). Published <i>With international search report.</i>
(54) Title: TREATMENT OF INFLAMMATORY DISORDERS OF THE BOWEL AND URINARY BLADDER (57) Abstract <p>A method of treating inflammatory disorders of the bowel and inflammatory disorders of the urinary bladder is disclosed. A method of modulating the function of monocytes is also disclosed. The method involves the treatment of a patient's blood with a photoactivatable compound followed by ultraviolet light-activation of the photoactivatable compound. The blood treated as such is returned to the patient in a process known as extracorporeal photopheresis.</p>		

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TREATMENT OF INFLAMMATORY DISORDERS OF THE BOWEL AND URINARY BLADDER

Background of the Invention

5 Extracorporeal photopheresis is a process where 8-Methoxypsoralen (8-MOP), a naturally occurring light-sensitive compound, is administered orally two hours prior to treatment; blood is then withdrawn from the patient, anti-coagulated, and the white blood cells are separated by centrifugation and collected as a leukocyte enriched fraction. These 8-MOP containing
10 leukocytes are then irradiated with ultraviolet A light (UVA) which binds the 8-MOP to pyrimidine bases in DNA and to intra- and extra-cellular proteins. These treated leukocytes are returned to the patient, and the result is an immunomodulation which has been found to be of clinical benefit in a number of disease states [Edelson RL. Photopheresis: a clinically relevant immunobiologic response modifier. [Review] *Ann NY Acad Sci.* 191;
15 636:154-64].

There are a number of diseases which are felt to primarily involve T-cells or are T-cell mediated. Diseases such as cutaneous T-cell lymphoma, organ
20 allograft rejection after transplantation, progressive systemic sclerosis (PSS), inflammatory bowel disease (IBD), rheumatoid arthritis (RA) and juvenile onset diabetes mellitus (JODM) are thought to be T-cell mediated.

Cutaneous T-cell lymphoma (CTCL) is a malignant disease that is
25 progressive. Therapeutic options are limited. Edelson et al. performed a multi-center trial [Edelson R, Berger C, Gasparro F. et al. Treatment of cutaneous T cell lymphoma by extracorporeal photochemotherapy: Preliminary results. *N Engl J Med* 1987;316:297-303] which showed that 24 of 29 (83%) of erythrodermic patients experienced a significant improvement

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in their disease. These positive responses were seen at a median time of 22.4 weeks after initiation of therapy. Of clinical significance, these patients were those whose diseases were resistant to prior therapy which is felt to be a poor prognostic group. In addition, a decrease in the amount of peripheral blood involvement (Sezary cells) was seen. Actuarial data had indicated that median survival was increased to greater than 60 months from the onset of treatment in comparison with a historical median survival time of less than 30 months. In this original group of patients, remissions were sustained in eight of the subjects who were leukemic. Adverse reactions associated with photopheresis were rare.

Autoimmune diseases are characterized by a dysregulation of the immune system, characterized by specific cellular or humoral mediated destruction of specific organs or tissues in the patient. Examples of such diseases are rheumatoid arthritis and progressive systemic sclerosis.

Rheumatoid arthritis (R.A.) is an inflammatory disease that ultimately leads to joint destruction and is a generalized disease involving many organ systems. There are many pharmaceutical agents in use for R.A., however well tolerated agents with disease modifying potential are needed in as much as the disease is lifelong. In particular, a loss of efficacy and disease progression is seen in a high number of patients after starting secondary line therapy for R.A. Many of the second line agents are immunosuppressive and are themselves the reason for the major side effects such as infection. The need for development of a more specific, non-toxic immunomodulating therapy [Malawista S, Trock D, Edelson R. Treatment of rheumatoid arthritis by extracorporeal photochemotherapy: a pilot study. *Arthritis Rheum* 1991;34:646-54] is great.

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Progressive systemic sclerosis (PSS) is a connective tissue disease characterized by inflammatory and fibrotic changes in the skin and viscera. Treatment has been difficult. Anti-inflammatory drugs and corticosteroids are helpful in the early stages of the disease, but do not appear to influence the progression of the disease. Trials with D-penicillamine, methotrexate, cyclosporine, calcium channel blockers and prostagladins are underway, but these agents do not appear to influence the overall progression of the disease. As this disease has been considered to be T-cell mediated, Rock and colleagues have treated PSS patients with photopheresis [Rook AH, Freundlich B, Jegasothy BV, et al. Treatment of systemic sclerosis with extracorporeal photochemotherapy: Results of a multicenter trial. *Arch Dermatol* 1992;128:337-46]. In this trial, 56 patients were enrolled into a randomized non-blinded clinical trial. A significantly higher response rate was seen in the photopheresis treated group (68% response rate) compared to the D-penicillamine (control) group (32% response rate).

Juvenile onset diabetes mellitus (JODM) is felt to be mediated by the immune system resulting in the destruction of the cells in the pancreas responsible for the production of insulin. Patients with this disorder have not only dysregulation of their blood sugar levels, but the disease is characterized by a vasculopathy, resulting in specific organ damage leading to significant morbidity and mortality.

IBD is either limited to the colon (ulcerative colitis) or affects both the colon and the small intestine. In addition, there are intrainestinal manifestations of the disease including pyoderma gangrenosum, erythema nodosum, sclerosing cholangitis, ankylosing spondylitis, hepatitis, arthritis, and uveitis. IBD involves a dysfunction of the immunoregulatory mechanisms that downregulate immune responses to digestion products, while maintaining the

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ability to develop a specific immune response to pathogens. Exposure to methoxsalen, activated by UV light, modulates immunoregulatory function, allowing the mucosal T-cells to mount a lower proliferative response to common microbial antigens than peripheral T-cells.

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Other T-cell mediated phenomena include rejection of tissues that are foreign to the host. In the case of organ allograft transplantation, it is desirable to prevent rejection with respect to the transplanted organ, however to otherwise maintain the competence of the immune system, in order to allow the body to combat infection and to allow other normal body defenses. The standard treatments after transplantation are limited as the immunosuppression regimens are used to cause a state of generalized immunosuppression, which leads to microbial or opportunistic infection, the most common adverse reaction to this treatment. Immunomodulation which does not have broad immunosuppressive properties would be more desirable.

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Photopheresis has been shown to be effective, and investigators at Loyola University have been able to successfully treat with photopheresis 13 of 14 cases of cardiac rejection refractory to standard immunosuppressive agents. In a variation of this situation, photopheresis has been successfully used to treat a patient with chronic graft versus host disease [Rossetti *et al.*, 1995, *Transplant*, 59:1, pp.149-151]. This disorder is characterized by an introgenically induced immunoincompetent host, where immune competent cells (bone marrow or peripheral stem cells) are infused into a patient in such situations as treatment for various malignancies and leukemia. Here the transplanted immunocompetent cells attack the patient (the "host"), and the issue is to modulate the immunocompetent cells without causing further broad immunosuppression and the side effects thereof.

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Photopheresis involves the extracorporeal exposure of peripheral blood leukocytes to 8-methoxypsoralen (8-MOP) photoactivated by ultraviolet A light, followed by the reinfusion of the treated white blood cells.

5 8-methoxypsoralen molecules in the blood enter the white blood cell nuclei and intercalate in the double-strand DNA helix. In an extracorporeal circuit, long wave ultraviolet light is directed at the leukocyte-enriched blood fraction within the UVAR® Photopheresis System. The photoactivated drug, responding to the UVA energy, links to the thymidine base in the DNA
10 helix. This results in cross-linking of thymidine bases which prevents the unwinding of the DNA during transcription. The plasma and altered leukocytes are then reinfused into the patient. The reinfusion of the photopheresis damaged leukocytes results in an delayed immune attack against these damaged leukocytes, as well as, otherwise unmodified WBC's
15 displaying the same cell surface antigens.

Methoxsalen is a naturally occurring photoactive substance found in the seed of the Ammi majus (umbelliferae plant). It belongs to a class of compounds known as psoralens or furocoumarins. The chemical name is 9-methoxy-7H-furo[3,2-g][1]-benzopyran-7-one. The formulation of the drug is a sterile liquid
20 at a concentration of 20 mcg/mL in a 10 mL vial. The pharmacokinetic activity of methoxsalen is available.

25 Toxicology studies of extracorporeal photochemotherapy and different dosages of UVADEX® and ultraviolet light in beagle dogs is located in the investigator's brochure.

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UVAR System

The treatment consists of three phases including: 1) the collection of a buffy-coat fraction (leukocyte-enriched), 2) irradiation of the collected buffy coat fraction, and 3) reinfusion of the treated white blood cells. The collection phase has six cycles of blood withdrawal, centrifugation, and reinfusion steps. During each cycle, whole blood is centrifuged and separated in a pediatric pheresis bowl. From this separation, plasma (volume in each cycle is determined by the UVAR® Instrument operator) and 40 ml buffy coat are saved in each collection cycle. The red cells and all additional plasma are reinfused to the patient before beginning the next collection cycle. Finally, a total of 240 ml of buffy coat and 300 ml of plasma are separated and saved for UVA irradiation.

The irradiation of the leukocyte-enriched blood within the irradiation circuit begins during the buffy coat collection of the first collection cycle. The collected plasma and buffy coat are mixed with 200 ml of heparinized normal saline and 200 mcg of UVADEX® (water soluble 8-methoxypsoralen). This mixture flows in a 1.4 mm thick layer through the PHOTOCEPTOR® Photoactivation Chamber, which is inserted between two banks of UVA lamps of the PHOTSETTE®. PHOTSETTE® UVA lamps irradiate both sides of this UVA-transparent PHOTOCEPTOR® chamber, permitting a 180-minute exposure to ultraviolet A light, yielding an average exposure per lymphocyte of 1-2 J/cm². The final buffy coat preparation contains an estimated 20% to 25% of the total peripheral blood mononuclear cell component and has a hematocrit from 2.5% to 7%. Following the photoactivation period, the volume is reinfused to the patient over a 30 to 45 minute period.

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Systems employing these techniques are known whereby extracorporeal treatment of a patient's blood is undertaken. For example, in U.S. Patent No. 4,573,960--Goss, a patient is given a drug that requires photoactivation and the patient's blood is then withdrawn and separated into its components. The untreated components (red blood cells, some plasma, etc.) are returned to the patient. The patient is then disconnected from the treatment apparatus and the separated components, e.g., white blood cells, are exposed to ultraviolet light. Following photoactivation, the treated cells are returned to the patient.

In U.S. Pat. Nos. 4,321,919; 4,398,906; and 4,464,166, issued to Edelson, the external treatment methods for diseases in which there is a pathological increase of lymphocytes, such as cutaneous T-cell lymphoma, have been discussed. In these methods the patient's blood in the presence of a chemical or an antibody is irradiated with ultraviolet light. Ultraviolet light effects a bonding between the lymphocytes and the chemical or antibody thus inhibiting the metabolic processes of the lymphocytes.

A variety of human viruses are able to infect and replicate within mononuclear cells, or infectious viral particles may remain present within the mononuclear cells. The mononuclear cells can act as either a source for viral replication and spread of the virus, or as a reservoir of infectious virus particles which is difficult for the immune system to eliminate. Failure to eliminate these sources of infectious virus may lead to the establishment of a chronic condition. Viruses which can infect, replicate within, or reside in mononuclear cells include, but are not limited to, arthropod borne viruses, enteroviruses, paramyxoviruses (RSV), herpes viruses, cytomegalo-virus (CMV), Epstein-Barr virus (EBV), hepatitis B virus (HBV), hepatitis C virus (HCV), hepatitis D virus (HDV), hepatitis G virus (HGV), and retroviruses (such as HIV).

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5 A variety of human non-viral pathogenic agents are able to infect and replicate within mononuclear cells, or the infectious non-viral pathogenic agents may remain present within the mononuclear cells. The mononuclear cells can act as either a source for replication and spread of the non-viral pathogenic agents, or as a reservoir of infectious non-viral pathogenic agents which is difficult for the immune system to eliminate. Failure to eliminate these sources of infectious non-viral pathogenic agents may lead to the establishment of a chronic condition. Non-viral pathogenic agents which can
10 infect, replicate within, or reside in mononuclear cells include, but are not limited to, bacteria such as arthropod-borne bacteria, mycoplasma species, and mycobacteria species, and parasites such as plasmodium species and other arthropod-borne parasites.

15 Extracorporeal photopheresis (ECP) has been successfully used to treat HIV infection (U.S. Patent No. 4,960,408) and psoralen compounds with long wavelength ultraviolet light have been shown to inactivate certain viruses in vitro, such as HIV (Quinnan, G.V. et al., 1986, Transfusion, 26, pp 481; Bisaccia, A. et al., 1990, Am. Intern. Med., 113, pp 270; Bisaccia, A. et al.,
20 1991, Ann. NY Acad. Sci., 636, pp 321), and influenza virus and herpes simplex virus (Redfield, D.C. et al., 1981; Infect. and Immun., 32, pp 1216). Bisaccia has studied ECP in a pilot trial as therapy for patients with AIDS-related complex. The rationale was that a combination of psoralen with UVA activation could damage HIV *in vitro* and that reinfusion of the
25 damaged virus may initiate an immune response. The authors found that ECP produced an increase in the HIV-Ab production, increase in the CD8(+) lymphocytes, a decrease in the p24 antigen titer and the inability to culture HIV in 3 patients. Eleven of the 20 patients had improvement in their skin test antigen reactivities.

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In addition, a reduced incidence of infection episodes was reported in patients receiving photopheresis treatment for immunosuppression following transplant surgery (Meiser, B.M. et al., 1994, Transplantation, 57, pp. 563).

However, the results observed for the transplant surgery patients did not correlate with photopheresis treatment since infection episodes in general were recorded including patients who received a variety of treatments to prevent rejection of the transplanted organ.

Summary of the Invention

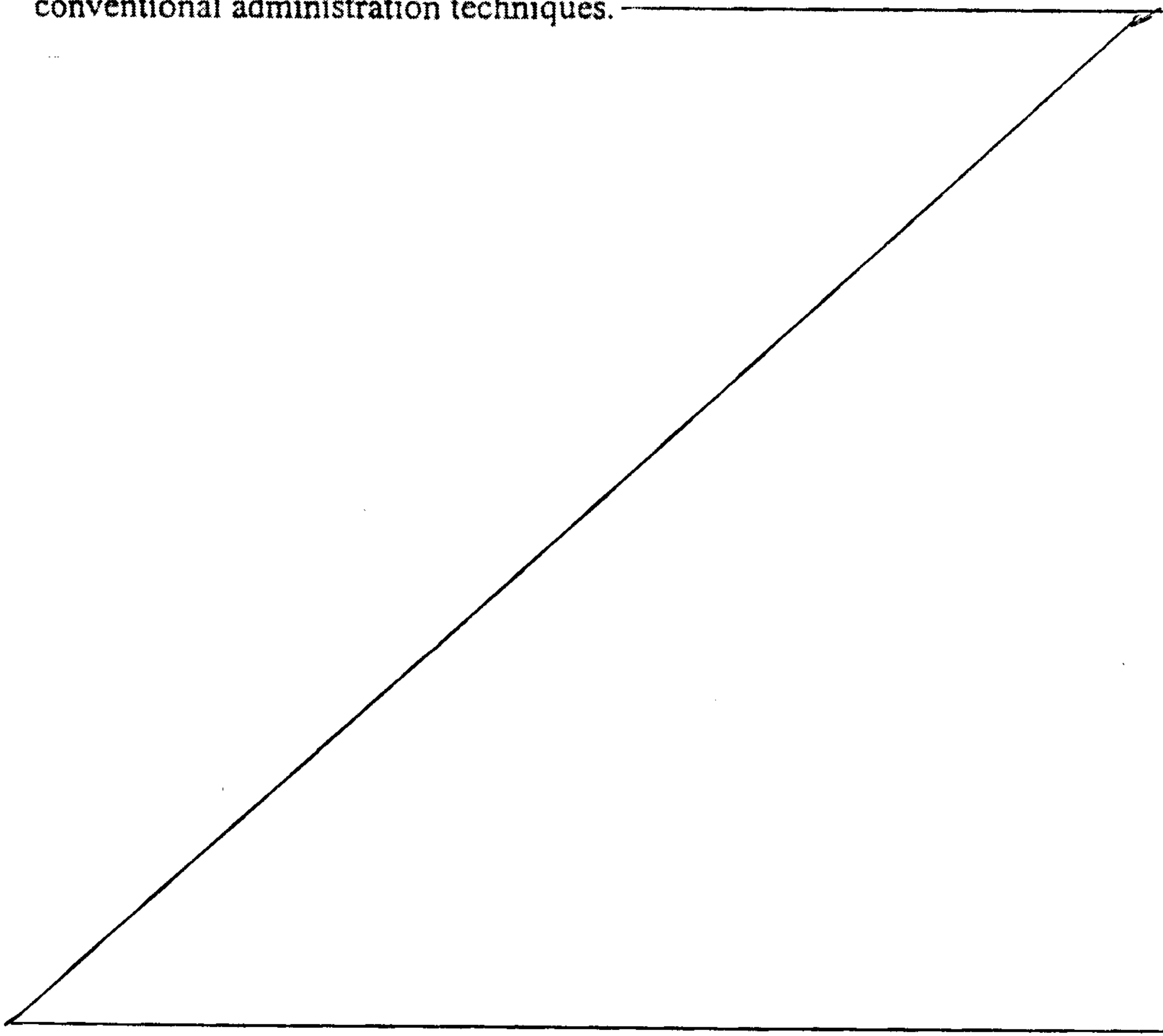
In one aspect, the present invention provides the use of a photoactivatable compound comprising psoralen or a psoralen derivative together with light in a wavelength that activates said photoactivatable compound in an extracorporeal photopheresis treatment, for treating an inflammatory bowel disorder in a patient.

In another aspect, the present invention provides the use of a photoactivatable compound comprising psoralen or a psoralen derivative together with a light in a wavelength that activates said photoactivatable compound in an extracorporeal photopheresis treatment, for treating an inflammatory disorder of the urinary bladder in a patient.

The present invention is drawn to a method of treating inflammatory bowel disorders, including including but not limited to IBD, Crohn's Disease, and ulcerative colitis, using extracorporeal photopheresis. In addition, the present invention is drawn to the treatment of inflammatory diseases of the urinary bladder, including but not limited to cystitis, such as interstitial cystitis. In particular, patients with IBD, Crohn's Disease, ulcerative colitis or interstitial

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cystitis are treated by the method of the present invention. The present invention is also drawn to the alteration or modulation of monocyte function in patients having inflammatory bowel disorders or inflammation of the urinary bladder through the use of the method of photopheresis. The method of treatment in the present invention involves the treatment of a patient's blood with a photoactivatable or photosensitive compound which is capable of binding to nucleic acids in infected nucleated cells upon activation of the compound by ultraviolet light. The photoactivatable or photosensitive compound may be administered to the patient's blood in vitro or in vivo by conventional administration techniques.



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A portion of the patient's blood is then treated extracorporeally using photopheresis, which comprises subjecting the blood to ultraviolet light, preferably long wavelength ultraviolet light in the wavelength range of 320nm to 400nm, commonly called UVA light. The treated blood, or a fraction thereof, is returned to the patient following extracorporeal photopheresis to modulate monocyte function and/or stimulate an immunological response by the patient's immune system. The cellular genetic material is damaged by this treatment which may result in the alteration or modulation of monocyte function as disclosed herein.

Detailed Description of the Invention

The present invention is directed to the use of photopheresis to treat patients having an inflammatory bowel disorder, including but not limited to IBD, Crohn's Disease, Ulcerative colitis or patients having an inflammatory disorder of the urinary bladder, including but not limited to cystitis, such as interstitial cystitis. While it is not intended that the scope of the present invention be limited by any specific theory of operation, it is believed that inflammatory bowel disorders and inflammatory disorders of the urinary bladder of a patient can be treated by using a photopheresis treatment according to the present invention.

Extracorporeal photochemotherapy using methoxsalen (photopheresis) is believed to cause an immunization against the abnormal (cancerous, in the case of CTCL) T-cells. During photopheresis, methoxsalen enters the white blood cell nuclei and intercalates in the double-strand DNA helix. In an extracorporeal circuit, long wave ultraviolet light is directed at the leukocyte-enriched blood volume. The methoxsalen, responding to the UVA energy, links to the thymidine base in the DNA helix. This results in the cross-linking of thymidine bases which prevent the unwinding of the DNA during

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transcription. Ultraviolet A light (UVA) damages abnormal T-cells rendering them more immunogenic. After cells are photoactivated, reinfusion of these altered T-cells causes an immunological reaction that targets T-cells carrying the same surface antigens[Edelson RL. Photopheresis: a clinically relevant immunobiologic response modifier. [Review] *Ann NY Acad Sci.* 191; 636:154-64]. This results in the production of a highly specific immune response against the abnormal cells (either a cancer clone or perhaps T-cells which express viral antigens on their surface). It is estimated that approximately 25-50% of the total peripheral blood mononuclear cell 5 compartment is treated per photopheresis session (2 consecutive days schedule).

Work by Vowels [Vowels BR, Cassin M, Boufal MH, et al. Extracorporeal photochemotherapy induces the production of tumor necrosis factor-alpha by monocytes: Implications for the treatment of cutaneous T cell lymphoma and systemic sclerosis. *J Invest Dermatol* 1992;98:686-92]. demonstrated monocytes treated in an extracorporeal circuit of plasma containing 8-methoxypsoralen and exposure to ultraviolet-A light (photopheresis) releases tumor necrosis factor-alpha, IL-1, IL-6, and possibly IL-8. It is believed that photopheresis modulates the activity of peripheral blood monocytes/macrophages.

According to the claimed methods, a photoactivatable or photosensitive compound is first administered to the blood of a patient who has an inflammatory bowel disorder or an inflammatory disorder of the urinary bladder. The photoactivatable or photosensitive compound may be administered in vivo (e.g. orally or intravenously) or may be administered in vitro to a portion of the patient's blood which has been removed from the patient by employing conventional blood withdrawal techniques.

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In accordance with the present invention, the photoactivatable or photosensitive compound should be capable of binding to nucleic acids upon activation by exposure to electromagnetic radiation of a prescribed spectrum, e.g., ultraviolet light.

Next, the portion of the patient's blood to which the photoactive compound has been administered is treated by subjecting the portion of the blood to photopheresis using ultraviolet light. The photopheresis step is preferably carried out in vitro using an extracorporeal photopheresis apparatus. The photopheresis step in accordance with the present invention may also be carried out in vivo. A presently preferred extracorporeal photopheresis apparatus for use in the methods according to the invention is currently manufactured by Therakos, Inc., under the name UVAR®. A description of such an apparatus may be found in U.S. Patent No. 4,683,889. The exposure of blood to ultraviolet light in a photopheresis apparatus is within the ability of persons having ordinary skill in the art.

When the photopheresis step is carried out in vitro, at least a fraction of the treated blood is returned to the patient. Preferably, the treatment method described herein is repeated at an interval of about once per week to about once every four weeks. Preferred photoactive compounds for use in accordance with the present invention are compounds known as psoralens (or furocoumarins) as well as psoralen derivatives such as those described in U.S. Patent No. 4,321,919 and U.S. Patent No. 5,399,719. Alternatively, the patient's blood can be separated on a standard apheresis-type device and photoactivated on a separate device.

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The preferred photoactivatable or photosensitive compounds for use in accordance with the present invention include, but are not limited to, the following:

5 Psoralen and psoralen derivatives; 8-methoxypsoralen; 4,5',8-trimethylpsoralen; 5-methoxypsoralen; 4-methylpsoralen; 4,4-dimethylpsoralen; 4-5'-dimethylpsoralen; 4'-aminomethyl-4,5',8-trimethylpsoralen; 4'-hydroxymethyl-4,5',8-trimethylpsoralen; 4',8-methoxypsoralen; and a 4'-(omega-amino-2-oxa) alkyl-4,5',8-trimethylpsoralen, including but not limited to 4'-(4-amino-2-oxa)butyl-10 4,5',8-trimethylpsoralen. The most particularly preferred photosensitive compound for use in accordance with the invention is 8-methoxypsoralen.

15 The photosensitive compound, when administered to the patient's blood in vivo is preferably administered orally, but also can be administered intravenously and/or by other conventional administration routes. The preferred oral dosage of the photosensitive compound is in the range of about 0.3 to about 0.7 mg/kg. most preferably about 0.6 mg/kg.

20 When administered orally, the photosensitive compound should preferably be administered at least about one hour prior to the photopheresis treatment and no more than about three hours prior to the photopheresis treatment. If administered intravenously, the times would be shorter.

25 Alternatively the photosensitive compound may be administered to the patient's blood following its withdrawal from the patient, and prior to or contemporaneously with exposure to ultraviolet light. The photosensitive compound may be administered to whole blood or a fraction thereof provided that the target blood cells or blood components receive the photosensitive compound.

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The photopheresis treatment in the treatment methods according to the present invention is preferably carried out using long wavelength ultraviolet light (UVA) at a wavelength within the range of 320 to 400 nm. The exposure to ultraviolet light during the photopheresis treatment preferably has a duration of sufficient length to deliver about 1-2 J/cm² to the blood.

When the photopheresis treatment according to the present invention is carried out in vivo, careful attention should be paid to controlling the maximum radiant exposure so as to avoid unnecessary injury to the patient. Methods for calculating maximum radiant exposure to ultraviolet light are known in the art.

First, a photosensitive compound as described hereinabove is administered to at least a portion of the donor's blood either prior to removal of the blood, either orally or intravenously, or after removal from the patient in which case it is administered in vitro. Optionally, a portion of the donor's blood could first be processed using known methods to substantially remove the erythrocytes and the photoactive compound is then administered to the resulting enriched leukocyte fraction.

In any case, the portion of blood (or enriched leukocyte fraction thereof) to which the photosensitive compound has been administered is subjected to a photoactivation treatment using ultraviolet light, preferably UVA in the manner previously described. The treated blood or the treated enriched leukocyte fraction (as the case may be) is then administered back to the donor.

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Current treatment for inflammatory bowel disorders include chemotherapeutic agents such as aminosalicylates, corticosteroids, and immunosuppressants. Significant side effects from these chemotherapeutic treatments occur. Many patients develop side effects from aminosalicylates before achieving a therapeutic dose. The side effects of nausea, malaise, headache, and myalgia often result in patient non-compliance. The side effects of long term corticosteroids use (Cushingoid appearance, hypertension, cataract formation, osteoporosis and aseptic necrosis) are well documented. In addition, although corticosteroids may be effective in treating a flare in disease, the ability of these drugs to prevent relapse has never been proven. An increased risk of developing cancer, neutropenia and bone marrow suppression is well documented in the organ transplant recipient population.

A summary description of certain inflammatory bowel disorders which are treated in the method of the present invention, include but are not limited to:

1. Ulcerative Colitis

Ulcerative colitis is a mucosal disorder which affects the colon and is associated with significant morbidity and mortality. There are associated extraintestinal manifestations of this disease which include arthritis, ankylosing spondylitis, hepatitis, uveitis, pyoderma gangrenosum, erythema nodosum, and sclerosing cholangitis. Cancer can occur in areas of chronic inflammation. Specific symptoms of ulcerative colitis include diarrhea, bowel bleeding, weight loss, and abdominal pain and cramping. This disease can affect individuals of any age. Children with this disease fail to grow and develop normally.

2. Crohn's Disease

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Crohn's disease is characterized by focal, transmural and asymmetric inflammation involving one or more segments of the alimentary canal. extending anywhere from the mouth to the anus. The most commonly involved areas include the distal ileum and right colon. Exclusive colonic involvement occurs in about 20% and disease limited to the small bowel occurs in 15-20% in patients affected with Crohn's disease. The ulcerations associate with Crohn's disease extend linearly, often isolating pockets of normal mucosa, giving this disease a characteristic cobblestone appearance.

A summary description of certain inflammatory disorders of the urinary bladder which are treated in the method of the present invention, include but are not limited to:

1. **Interstitial Cystitis (IC)**

Interstitial cystitis (IC) is a chronic severe bladder disorder similar to IBD in that it is an autoimmune disease that manifests itself as an attack on the bladder wall rather than the bowel wall. Like IBD, IC histologically can be identified as a unifocal or multifocal inflammatory infiltration of the bladder wall with mucosal ulceration and scarring. It results in contraction of the smooth muscle, diminished urinary capacity, symptoms of frequency, hematuria, urgency, nocturia, painful urination and suprapubic pelvic pain. Treatments such as amitriptyline, hydroxyzine, dimethyl sulfoxide, chlorpactin and heparin may improve symptoms but do not alter the long term course of the disease.

The result of the treatment method of the present invention for inflammatory bowel disorders, including but not limited to IBD, Crohn's Disease and for inflammatory disorders of the urinary bladder including but not limited to cystitis, such as IC, is an elimination of the symptoms of the disease,

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reflected as a reduction in the activity index for the disease and/or a significant reduction in steroid dose (particularly for those who are steroid dependent) and/or the avoidance of surgery (removal of the affected tissue) and increasing symptom-free intervals in these patients.

5

The following Examples are provided to illustrate the present invention and are not to be construed as a limitation thereon.

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EXAMPLE 1**EXTRACORPOREAL PHOTOCHEMOTHERAPY IN THE TREATMENT OF PATIENTS WITH ULCERATIVE COLITIS**

15

I. Description of the Patient Population

Patients who have an acute flair of their ulcerative colitis were the subjects of this clinical investigation. To be included for inclusion into this protocol, the following inclusion/ exclusion criteria had to be met:

20

- Patients must be 40 kg
- Females of childbearing potential must be urine pregnancy test negative within 24 hours prior to starting the study treatment. Effective birth control must be practiced from one month prior to the study treatment to the end of treatment phase of the protocol. Females may not be lactating.

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- Pre-study laboratory values of:
 - WBC greater than 3.0 cmm
 - Hemoglobin greater than 6.5 mg/dL

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- Platelet count > 75,000/UL
- Prothrombin time not more than 3 seconds beyond the normal

range

- Able to provide informed consent
- Adequate venous access
- Able and willing to comply with the study protocol and medication schedules

The diagnosis of ulcerative colitis must be confirmed by the history and physical exam, endoscopic and histologic evaluation. The extent of disease must be at least 10 cm from the anal verge.

- To qualify for the study:

The patient must be experiencing an acute, moderate to severe relapse of their UC and will meet at least three of the seven criteria listed:

- Diarrhea >6/day
- Gross blood in the stool
- Fever >37.7
- Heart rate >90/min
- Anemia less than 75% of the normal
- ESR >30
- Steroid dependent

-or-

- Candidate for surgical intervention

II. Description of the Treatment Frequency and Efficacy Parameters

Photopheresis treatments were performed on two consecutive days for four weeks (8 photopheresis treatments) and then were performed on two consecutive days every other week for the next eight weeks (8 additional

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treatments) for a total of 16 photopheresis procedures over 12 weeks. (Defined as treatment period I). The goal of treatment period I was to assess the primary efficacy parameters (disease activity index [DAI], assess need for surgical intervention, assess ability of patients to wean from steroids or other immunosuppressive medications used to control UC symptoms and to assess the response to treatment on an endoscopical/histological level). For the purpose of this protocol, a successful response to treatment was defined as:

A 50% decrease in the disease activity index (DAI), the avoidance of surgery or the reduction/elimination of steroids and improvement in endoscopy grade and histology are all considered to be clinically significant.

After treatment phase I was completed, the patient was then followed for nine months upon the discontinuation of treatment. Flares (as evidenced by increase in ulcerative colitis symptomatology [diarrhea, bleeding and pain]) during this follow-up period were treated with photopheresis, at the Investigator's discretion. The goal of this follow-up protocol was to calculate: (1) disease free intervals, (2) to determine whether flares in disease can be controlled by photopheresis, and (3) to provide data on weaning from photopheresis therapy.

III. Description of the Efficacy Assessments:

During the treatment and follow-up periods, the patient had the following efficacy assessments:

A. **Disease activity index:** This is a modified Truelove and Witts scale - _____ (Table 1). This scale assesses

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the number of bloody bowel movements, presence of nocturnal diarrhea, incontinence, abdominal pain, general well-being, and abdominal tenderness. The minimum score is 0. The maximum score is 21. Scores of 12 or higher are considered to be in the severe category. Significant improvement indicates a decrease in DAI by 50%. Scores less than 4 indicate clinical remission.

Table 1: Disease Activity Index

Disease Activity	Score
Diarrhea (number of daily stools)	
0-2	0
3-4	1
5-6	2
7-9	3
>10	4
Nocturnal diarrhea	
No	0
Yes	1
Visible blood in stool (% of BM)	
0%	0
<50%	1
>50%	2
100%	3
Fecal incontinence	
No	0
Yes	1
Abdominal pain	
None	0
Mild	1
Moderate	2
Severe	3
General well-being	
Perfect	0
Very Good	1
Good	2
Average	3
Poor	4
Terrible	5
Abdominal tenderness	
None	0
Mild and localized	1
Mild to moderate and diffuse	2
Severe or rebound	3
Need for anti-diarrheals	
No	0
Yes	1

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B. Endoscopic findings were scored according to a _____ scale _____ which assesses severity of mucosal granularity, vascular pattern, friability, ulceration, and mucopus. Analysis of endoscopic findings were analyzed separately from the symptom assessment scores (Table 2). Included in the endoscopic assessment were determination of the length of endoscopic disease and its corresponding mucosal severity

Table 2: Endoscopic Mucosal Severity Score

Activity	Score	Appearance
Quiescent	0	<ul style="list-style-type: none"> • Distorted or absent mucosal vascular pattern • Granularity
Mild	1	<ul style="list-style-type: none"> • Continuous or focal erythema • Friability
Moderate	2	<ul style="list-style-type: none"> • Mucopurulent exudate • Single or multiple ulcers (<5mm), fewer than 10/10cm segment
Severe	3	<ul style="list-style-type: none"> • Large ulcers (>5mm), more than 10 per 10cm segment • Spontaneous bleeding

C. Histology: were scored by one Pathologist. The histology grading score is a system which grades both activity and severity of disease. Histology index (or score) were expressed by adding the values for all five criteria and this total was expressed as the total out of a maximum total score of fifteen.

1. ACTIVITY INDEX

Criterion #1: NEUTROPHIL LEUKOCYTES IN INTACT LAMINA PROPRIA

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Method: The number of neutrophils in one high power (x40) field of the most actively inflamed area of intact mucosa were counted. Ulcer debris and granulation tissue were excluded.

0 = no neutrophils

1 = 1-10/hpf

2 = 11-20/hpf

3 = >21/hpf

Criterion #2 ACUTE CRYPTITIS

Method: Evaluate neutrophil infiltration of crypts, including crypt abscesses and crypt destruction by the acute inflammation.

0 = no cryptitis

1 = crypt infiltration by neutrophils; no crypt abscesses; no active crypt destruction

2 = crypt abscesses +; no active crypt

destruction

3 = active crypt destruction

Criterion #3 EROSION/ULCERATION

0 = no erosion or ulceration

1 = focal partial thickness mucosal erosion

only

2 = presence of acutely inflamed necrotic

tissue

devoid of mucosa (= superficial ulcer)

3 = presence of acutely inflamed granulation

tissue

(= deep ulcer)

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2. DISEASE CHRONICITY INDEX

Criterion #4 CRYPT ARCHITECTURE

0 = normal

1 = mild distortion

2 = moderate distortion with definite branching

3 = severe distortion

**Criterion #5 CHRONIC INFLAMMATION IN
LAMINA PROPRIA**

Method: Assess lymphocyte, plasma cell and
eosinophil numbers (subjective, no count)

0 = normal

1 = mild increase

2 = moderate increase

3 = severe disease

Adverse events were assessed as follows:

- The frequency and severity of adverse events classified by their relationship to the device, drug and disease
- The frequency of abnormal laboratory parameters and new symptomatology at each treatment visit.

IV. Sample Schedule of Assessments

A. Baseline Assessments (within 48 hours of the first photopheresis treatment)

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- CBC
- Chemistry
- Sedimentation rate
- Disease activity index (DAI)
- Concomitant medication listing
- Endoscopy with histological evaluation
- Quality of life questionnaire

B. Prior to each two day photopheresis session

- CBC
- Concomitant medication listing

C. Every two weeks during the treatment phase

- Chemistry panel
- Sedimentation rate
- Disease activity index (DAI)

D. Pre-tx #5

- Quality of life questionnaire

E. Monthly

- Urine pregnancy test (if applicable)

F. At Weeks 6 and 12 of photopheresis treatment

- Endoscopic and histological evaluation
- Quality of life questionnaire
- CD4/CD8 (week #12 only)

G. Post Treatment Evaluation

One Month Following Last Treatment in Period I

- 5
- 10
- CBC
 - Sedimentation rate
 - Chemistry
 - Disease activity index (DAI)
 - CD4/CD8
 - Quality of life questionnaire
 - Concomitant medication listing

Months 3, 6 and 9

- 15
- 20
- CBC
 - Sedimentation rate
 - Chemistry
 - Disease activity index (DAI)
 - Quality of life questionnaire
 - Concomitant medication listing

Months 2, 4, 5, 7 and 8

- 25
- Disease activity index (DAI)

V. Results

Patient #1/1

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Assessment	Screenin g Visit	Pre-Photo Tx #5	Pre-Photo Tx #9	Pre-Photo Tx #11	Pre-Photo #13	Pre-Photo Tx #15
UC Histology Activity Score	6		1			
UC Chronicity Score	3		2			
Endoscopy Score	2		1			
Cm UC involvement from anal verge	20		10			
Quality of Life Questoinnaire	151	169		208.5		
ESR	9	7	7		9	
Disease Activity Index	8 2/20/98	5 3/12/98		2 4/17/98	2 5/2/98	

Patient UC 1/1

5

Summary: 26 year old female with endoscopically and histologically confirmed diagnosis of ulcerative colitis

10

Baseline Evaluation : Entered the protocol having diarrhea greater than 7-9 times per day, fecal incontinence, gross blood in the stool and a fever greater than 37.7. Colonoscopy showed mucosal changes from 20 to 0 cm. The findings included an erythematous appearance, superficial erosions, aphthoid erosions and areas of frank hemorrhage. The histology activity score was 6, the UC chronicity score was 3 and the endoscopy score was 2. The patient's disease activity index was 8 of 20 and the patient's quality of life questionnaire totaled 151 of a possible 224 points. Patient reported on the patient diary to feel "fair" and be experiencing "moderate" pain.

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Two Week Evaluation: The patient's disease activity index fell to 5 of 20, with the major areas of improvement being noted as a decreased number of episodes of diarrhea to 3-4 times per day and no fecal incontinence. The patient's quality of life increased to 169 out of a possible 224 points. The patient reported on the patient diary to feel "well" with only "mild" pain, the patient still reported blood in the stool.

Six Week Evaluation: The patient's disease activity index fell to 2 of 20, with the major area of improvement being noted as a decreased number of episodes of diarrhea to 0-2 times per day and no fecal incontinence. The patient's quality of life increased to 208 out of a possible 224 points. The patient reported on the patient diary to feel "very well" with "no" pain and reported many more days without blood in the stool. Sigmoidoscopy showed mucosal changes in the rectum, with both erythematous and granular changes. The visualized colon (up to 30 cm) were grossly normal. The UC histology score decreased from 6 to 1, UC chronicity score decreased from 3 to 2 and the endoscopy score reduced from 2 to 1.

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Patient #3 RJK

Assessment	Screening Visit	Pre-Photo Tx #1	Pre-Photo Tx #3	Pre-Photo Tx #5
UC Histology Activity Score	4			
UC Chronicity Score	0			
Endoscopy Score	2			
Cm UC involvement from anal verge	40			
Cm UC involvement from anal verge	40			
Quality of Life Questionnaire	Total score = 188			Total score = 212
ESR		15		
Disease Activity Index	3	3		3 4/23/98
	X			
	X			X
Concomitant Med Listing	Prednisone 20 mg QOD	Pred 20 mg QOD	Pred 15 mg QOD	Pred 10 mg QOD

5

Patient UC 1/3

Summary: 42 year old male with a 5+ year history of steroid dependency to control ulcerative colitis symptoms

10

Baseline Evaluation: Entered the protocol with bloody stools, colonoscopy showing mucosal changes from 0 cm to 40 cm. Endoscopy findings included edema, an erythematous and friable appearance, aphthoid erosions, (a rim of erythema, a yellow-gray central crater, absence of a raised margin), a blurred/blunted vascular pattern. DAI was 3 and patient was receiving Prednisone 20 mg every other day. Baseline quality of life was 188 out of a possible 224 points.

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Two Week Evaluation: DAI still 3, but patient has been weaned down to Prednisone 10 mg every other day, with no increase in symptoms. Quality of life has increased from 188 to 212.

EXAMPLE 2

EXTRACORPOREAL PHOTOCHEMOTHERAPY IN THE TREATMENT OF PATIENTS WITH CROHN'S DISEASE

I. Description of the Patient Population

Patients who have an acute flair of their Crohn's disease are the subjects of this clinical investigation. To be included for inclusion into this protocol, the following inclusion/exclusion criteria were met:

A. Inclusion Criteria

- Age between 18 and 65
- Diagnosis of Crohn's disease according to Malchow's criteria and Crohn's activity index (CDAI)
- Radiological and/or endoscopic and/or sonographic localization documented within the last 12 months before the start of the study
- Steroid dependent disease course with a documented treatment of at least 10 mg of prednisone per day for at least the last three months needed in order to achieve a remission (remission is defined as a CDAI <200 for two successive weeks). The steroid dependence must be

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confirmed by a history of at least one relapse (CDAI >200) in an attempt to reduce the steroid dose to <10 mg

– Written or oral consent in the presence of witness after the patient has been informed thoroughly by the treating physician

5 – Before beginning treatment, all women of childbearing age must give a documented negative serum beta-HCG pregnancy test within 24 hours before the first treatment. In addition, they must agree to follow an adequate form of contraception during the study

– No known allergy to heparin or to 8-methoxypsoralen

10 **B. Exclusion Criteria**

– Patients unable to cooperate with the screening or treatment procedures

15 – Patients with a high likelihood of needing surgical intervention because of bleeding, abscess or peritonitis

– Patients with clinically relevant stenosis of the gastrointestinal tract

– Patients with intestinal stoma

20 – Concurrent or previous exposure within the last three months of treatment with cyclosporin-A

– No previous exposure (within the last three months) or a stable dose of azathioprine within the last three months

25 – More than three antibiotic treatment phases with ciproxin or metronidazole over a period of fourteen days each within the last three months with confirmed identification of the pathogen

– Nursing or pregnant women

– Patients with a serious concomitant illness

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- Cardiovascular instability which would not allow the required volume of blood to be withdrawn during photopheresis
- Treatment with drugs with known photosensitizing potential
- Patients with poor venous access
- Patients with HIV, Hbs antigen or HCV positivity
- Patients with the following laboratory findings:
 - hemoglobin less than 10.5 g/dL
 - platelets less than $100 \times 10^9/L$
 - WBC $<4000 \times 10^9/l$
 - serum bilirubin >3 mg/dL
 - serum creatinine >2 mg/dL
 - PT $<60\%$, PTT >50 sec

II. Description of the Treatment Frequency and Efficacy Parameters

A. Treatment Frequency and Efficacy Assessments

Preliminary Phase 0

In the preliminary phase of the study, the patients who meet the inclusion criteria or who could meet them in future observation were selected. Retrospective documentation of the steroid-dependent course of the patient's disease must be guaranteed.

Each patient's CDAI was determined six weeks before the start of the study. In the next six weeks, the patient was stabilized at the maintenance dose, which was the lowest dose of steroid that ensures inactivity of the Crohn's symptoms (defined as a CDAI <200 to two successive weeks). The patients were brought into the study at the maintenance dose which must be >10 mg of prednisone per day.

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Further monitoring of the patients took place within the first week before the start of the study (week -1).

Assessment at Week (-1):

- Demographics
- History of Crohn's disease
- Localization of Crohn's disease
- Previous surgical history
- Previous treatment for Crohn's disease
- Documentation of evidence of steroid dependence

according to the definition of the inclusion criteria

- History and physical
- Laboratory assessment (CBC with differential, ESR, CRP, 1-acid glycoprotein, serum electrolytes, creatinine, BUN, SGOT, SGPT, bilirubin, γ -GT, serum glucose, PTT, PT and urinalysis

- Serum frozen for future study
- CDAI
- Documentation of extraintestinal manifestations of

Crohn's disease

- Concomitant medication listing
- Urine β -HCG test (when applicable)
- Compliance check
- Triple permeability test: sucrose, lactulose, mannitol
- Quality of life index

Preliminary Phase 1

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Patients who met the inclusion criteria were introduced into the study during the six-month preliminary phase. Prospective documentation of the disease activity and the cumulative steroid dose in the preliminary phase must be guaranteed.

In the preliminary phase, continuous attempts to reduce steroid dose were made according to a fixed steroid reduction schedule. Repeated attempts were made to reduce the dose to a level lower than the maintenance dose according to the respective schedule. The maintenance dose is was reduced permanently in the actual treatment phase.

The patient's clinical activity, measured by CDAI, was used as the criterion for deciding on whether to wean steroids as per the steroid weaning schedule.

Steroid reduction began with the first day of treatment in phase 1. The dosage was reduced by alternating the initial steroid dose with the next lower dose level of a standard dosage series (i.e., 50 mg, 37.5 mg, 25 mg, 15 mg, 10 mg, 5 mg) for two weeks. For the next two weeks, an attempt was made to reduce the dose to this level. If the clinical remission is maintained (no increase in CDAI >60 above the initial level or >200 after the second week), the steroid dose is reduced by an additional dose level for four weeks, as described above.

A recurrence of the primary illness (increase in CDAI > 60 over the starting value or > 200 over two weeks) was treated by returning the steroid dose to the next higher dose level of the dosage series for two weeks previous to the flare. If clinical remission cannot be achieved with this dose

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(CDAI <200 for two weeks), the steroid dose is increased to 50 mg per day for one week with subsequent reduction of dose according to the standard steroid weaning schedule. However, the dose was only reduced to the dose at which a clinical remission was most recently observed. This dose may be different from the maintenance dose and is referred to as the “remission stabilization dose,” the dose is again reduced according to the steroid weaning schedule. The possibility cannot be ruled out that the maintenance dose at the end of the treatment phase differs from that of the preliminary phase of the study.

The clinical course of the patient’s Crohn’s disease was stable for the last two weeks before the start of the photopheresis treatment phase.

In the last two weeks before the start of the photopheresis treatment phase, the patient was given a Crohn’s diary for documenting the symptoms associated with their disease.

In the preliminary phase, blood sample for CBC with differential, chemistry, CRP and 1-acid glycoprotein was obtained every four weeks.

Treatment Phase

Photopheresis treatments were performed on two consecutive days on a monthly basis for six months.

Prior to the first day of treatment at each one of the monthly visits, the following assessments were performed:

- Assessment of clinical status
- CDAI
- Documentation of extraintestinal symptoms of Crohn’s disease

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– Documentation of steroid dose and other concomitant medications

– Laboratory assessment: blood samples for CBC with differential, chemistry, CRP and 1-acid glycoprotein

5 – Assessment of adverse events

– Quality of life index

– Permeability test (months 0 and 6 only)

– Colonoscopy with biopsy is optional

10 Steroid reduction began with the first day of treatment. The dose reduction amounts to 5 mg of prednisone in four week according to the standard steroid weaning schedule. Relapses of Crohn's disease with CDAI values of more that 450 lead to termination of the study.

III. Results

The Effect of Photopheresis on 5 Crohn's Disease Patients

(Arithmetic mean standard deviation, median and range)

24-Week Pre-Photopheresis Observation Period				24-Week Photopheresis Treatment Period				Changes observed during the 24-week photopheresis treatment period minus the changes observed during the 24-week pre-photopheresis observation period
Parameter	Baseline	Final	Final Minus Baseline	Baseline	Final	Final Minus Baseline		
CDAI	173.2 ± 72.5 MD = 167 (95 to 278)	183.6 ± 65.4 MD = 181 (115 to 265)	10.4 ± 90.7 MD = -5 (-97 to 137)	166.2 ± 69.1 MD = 181 (53 to 230)	132.8 ± 50.7 MD = 114 (86 to 191)	-33.4 ± 77.0 MD = -19 (-144 to 61)	-43.8 ± 157.9 MD = -62 (-281 to 103)	
Prednisone Usage (Mg)	78.0 ± 59.5 MD = 70 (25 to 175)	101.0 ± 43.8 MD = 85 (70 to 175)	23.0 ± 34.4 MD = 15 (-15 to 70)	93.0 ± 40.2 MD = 90 (55 to 160)	45.5 ± 45.8 MD = 35 (7.5 to 125)	-47.5 ± 14.1 MD = -55 (-62.5 to -30)	-70.5 ± 46.6 MD = 70 (-125 to -15)	
Patient Wt. (Kgs)	67.4 ± 4.8 MD = 67 (62 to 74)	67.8 ± 5.8 MD = 66 (64 to 78)	0.4 ± 3.04 MD = 1 (-4 to 4)	67.6 ± 5.9 MD = 65 (64 to 78)	68.4 ± 7.4 MD = 66 (62 to 81)	0.8 ± 2.4 MD = 2 (-3 to 3)	0.4 ± 1.7 MD = 0 (-1 to 3)	

During this treatment phase, steroids were reduced and Crohn's disease symptoms were controlled (CDAI <200) while receiving photopheresis.

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EXAMPLE 3

10

**EXTRACORPOREAL PHOTOCHEMOTHERAPY IN THE TREATMENT OF
PATIENTS WITH INTERSTITIAL CYSTITIS****I. Description of the Patient Population**

15

Patients who have an acute flair of their interstitial cystitis are the subjects of this clinical investigation. To be included for inclusion into this protocol, the following inclusion/exclusion criteria are to be met:

A. Inclusion Criteria

20

– Age between 18 and 65

– Written or oral consent in the presence of witness after the patient has been informed thoroughly by the treating physician

25

– Before beginning treatment, all women of childbearing age must give a documented negative serum beta-HCG pregnancy test within 24 hours before the first treatment. In addition, they must agree to follow an adequate form of contraception during the study

– No known allergy to heparin or the 8-methoxypsoralen

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– Patient must have pain in bladder upon filling and relief upon emptying bladder

-and-

Patient must have suprapubic, pelvic, urethral, vaginal or perineal pain

-and-

Patient must have waking frequency of micturation >5 time per day and nocturia >2 times per night

-and-

Patient must have glomerulation after hydrodistention on cystoscopy

-OR-

Hunner's ulcer

B. Exclusion Criteria

– Patients unable to cooperate with the screening or treatment procedures

– Patients with benign or malignant bladder tumors

– Patients with radiation, bacterial or cyclophosphamide induced cystitis

– Patients with vaginitis

– Patients with symptomatic urethral diverticulum

– Patients with uterine, cervical, vaginal or urethral

cancers

– Patients with active herpes

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- Patients with bladder or lower urethral calculi
- Symptoms of cystitis are relieved by antibiotics, urinary antiseptics, urinary analgesics
- Duration of symptoms has been less than 12 months
- Patient has involuntary bladder contractions by urodynamics
- At a bladder capacity of 400 mL, there is an absence of sensory urgency
- Nursing or pregnant women
- Patients with a serious concomitant illness
- Cardiovascular instability which would not allow the required volume of blood to be withdrawn during photopheresis
- Treatment with drugs with known photosensitizing potential
- Patients with poor venous access
- Patients with HIV, Hbs antigen or HCV positivity
- Patients with the following laboratory findings:
 - hemoglobin less than 10.5 g/dL
 - platelets less than $100 \times 10^9/L$
 - WBC $<4000 \times 10^9/L$
 - serum bilirubin $>3 \text{ mg/dL}$
 - serum creatinine $>2 \text{ mg/dL}$
 - PT $<60\%$, PTT $>50 \text{ sec}$

II. Description of the Treatment Frequency and Efficacy

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Photopheresis treatments are performed on two consecutive days weekly for four weeks (8 photopheresis treatments) and then are performed on two consecutive days every other week for the next eight weeks (8 additional treatments) for a total of 16 photopheresis procedures over 12 weeks. Photopheresis is weaned after 12 weeks to a frequency of monthly and beyond, depending on the patient's long-term response. The treatment can be reinitiated to treat a disease flare.

A significant reduction in pain, hematuria and frequency of urination is expected resulting from the treatment method of this invention. The type and amount of concomitant medications that the patients require for treatment of IC is reduced, resulting in an overall increase in the quality of life and disease reduction.

The embodiments of the invention in which an exclusive property or privilege is claimed are defined as follows:

1. Use of a photoactivatable compound selected from the group consisting of a psoralen and a psoralen derivative together with light in a wavelength that activates said photoactivatable compound in an extracorporeal photopheresis treatment, for treating an inflammatory bowel disorder in a patient.
2. Use according to claim 1, wherein said inflammatory bowel disorder is an Inflammatory Bowel Disease.
3. The use according to claim 2, wherein said Inflammatory Bowel Disorder is selected from the group consisting of Crohn's Disease and ulcerative colitis.
4. Use according to any one of claims 1 to 3, wherein said psoralen or psoralen derivative is 8-methoxypsoralen.
5. The use of any one of claims 1 to 3 wherein the compound is a psoralen.
6. The use of any one of claims 1 to 3 wherein the compound is a psoralen derivative.
7. Use of a photoactivatable compound selected from the group consisting of a psoralen and a psoralen derivative together with light in a wavelength that activates said photoactivatable compound in an extracorporeal photopheresis treatment, for treating an inflammatory disorder of the urinary bladder in a patient.
8. Use according to claim 7, wherein said inflammatory disorder of the urinary bladder is cystitis.
9. Use according to claim 7, wherein said inflammatory disorder of the urinary bladder is interstitial cystitis.

10. Use according to any one of claims 7 to 9, wherein said psoralen or psoralen derivative is 8-methoxypsoralen.
11. The use of any one of claims 7 to 9 wherein the compound is a psoralen.
12. The use of any one of claims 7 to 9 wherein the compound is a psoralen derivative.