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(54) Title: COMBINATION OF ANGIOTENSIN CONVERTING ENZYME INHIBITOR, SIDE-EFFECT-REDUCED AMOUNT OF ALDOSTERONE ANTAGONIST AND DIURETIC		
(57) Abstract Combinations of an ACE inhibitor, an aldosterone receptor antagonist and a diuretic, i.e., "triple therapy" combinations, are described for use in treatment of circulatory disorders. Of particular interest are triple therapies using captopril or enalapril co-administered with a low-dose of spironolactone and furosemide. This triple therapy would be particularly useful to treat or prevent the progression of congestive heart failure while avoiding or reducing aldosterone-antagonist-induced side effects such as hyperkalemia.		

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COMBINATION OF ANGIOTENSIN CONVERTING ENZYME INHIBITOR, SIDE-EFFECT-REDUCED
AMOUNT OF ALDOSTERONE ANTAGONIST AND DIURETIC

5

Field of the Invention

Combinations of an angiotensin converting enzyme
10 inhibitor, an aldosterone receptor antagonist and a
diuretic are described for use in treatment of circulatory
disorders, including cardiovascular diseases such as heart
failure, hypertension and congestive heart failure. Of
particular interest are therapies using a spiro lactone-type
15 aldosterone receptor antagonist compound in combination
with an angiotensin converting enzyme inhibitor with a
diuretic, using a side-effect-reduced amount of the
aldosterone receptor antagonist.

20

Background of the Invention

Myocardial (or cardiac) failure, that is, heart
failure ("HF"), whether a consequence of previous
myocardial infarction(s), heart disease associated with
25 hypertension, or primary cardiomyopathy, is a major health
problem of worldwide proportions. The incidence of
symptomatic heart failure has risen steadily over the past
several decades.

30

In clinical terms, decompensated cardiac failure
consists of a constellation of signs and symptoms that
arise from congested organs and hypoperfused tissues to
form congestive heart failure (CHF) syndrome. Congestion
is caused largely by increased venous pressure and by
35 inadequate sodium (Na⁺) excretion, relative to dietary Na⁺
intake, and is importantly related to circulating levels of
aldosterone (ALDO). An abnormal retention of Na⁺ occurs

via tubular epithelial cells throughout the nephron, including the later portion of the distal tubule and cortical collecting ducts, where ALDO receptor sites are present.

5

ALDO is the body's most potent mineralocorticoid hormone. As connoted by the term mineralocorticoid, this steroid hormone has mineral-regulating activity. It promotes Na⁺ reabsorption not only in the kidney, but also
10 from the lower gastrointestinal tract and salivary and sweat glands, each of which represents classic ALDO-responsive tissues. ALDO regulates Na⁺ and water resorption at the expense of potassium (K⁺) and magnesium (Mg²⁺) excretion.

15

ALDO can also provoke responses in non-epithelial cells. Elicited by a chronic elevation in plasma ALDO level that is inappropriate relative to dietary Na⁺ intake, these responses can have adverse consequences on
20 the structure of the cardiovascular system. Hence, ALDO can contribute to the progressive nature of myocardial failure for multiple reasons.

Multiple factors regulate ALDO synthesis and
25 metabolism, many of which are operative in the patient with myocardial failure. These include renin as well as non-renin-dependent factors (such as K⁺, ACTH) that promote ALDO synthesis. Hepatic blood flow, by regulating the clearance of circulating ALDO, helps determine ALDO plasma
30 concentration, an important factor in heart failure characterized by reduction in cardiac output and hepatic blood flow.

The renin-angiotensin-aldosterone system ("RAAS")
35 is one of the hormonal mechanisms involved in regulating pressure/volume homeostasis and also in the development of hypertension, a precursor condition implicated in the

progression of more serious cardiovascular diseases such as congestive heart failure. Activation of the renin-angiotensin-aldosterone system begins with secretion of the enzyme renin from the juxtaglomerular cells in the kidney.

5 The enzyme renin acts on a naturally-occurring substrate, angiotensinogen, to release a decapeptide, Angiotensin I. This decapeptide is cleaved by angiotensin converting enzyme ("ACE") to provide an octapeptide, Angiotensin II, the primary active species of this system. This

10 octapeptide, angiotensin II, is a potent vasoconstrictor and also produces other physiological effects such as stimulating aldosterone secretion, promoting sodium and fluid retention, inhibiting renin secretion, increasing sympathetic nervous system activity, stimulating

15 vasopressin secretion, causing positive cardiac inotropic effect and modulating other hormonal systems.

Emphasis has been placed on minimizing hyperaldosteronism as a basis for optimizing patient

20 treatment. This includes the importance of ALDO-receptor antagonism both in patients treated with conventional diuretic programs and in patients treated with angiotensin-converting enzyme (ACE) inhibitors, who are often constrained to small doses of ACE inhibitor because of

25 orthostatic hypotension. Such patients may demonstrate a recurrence of heart failure symptoms likely related to elevations in plasma ALDO levels.

Many aldosterone receptor blocking drugs and

30 their effects in humans are known. For example, spironolactone is a drug which acts at the mineralocorticoid receptor level by competitively inhibiting aldosterone binding. This steroidal compound has been used for blocking aldosterone-dependent sodium

35 transport in the distal tubule of the kidney in order to reduce edema and to treat essential hypertension and primary hyperaldosteronism [F. Mantero et al, Clin. Sci.

Mol. Med., 45 (Suppl 1), 219s-224s (1973)]. Spironolactone is also used commonly in the treatment of other hyperaldosterone-related diseases such as liver cirrhosis and congestive heart failure [F.J. Saunders et al, 5 Aldactone: Spironolactone: A Comprehensive Review, Searle, New York (1978)]. Progressively-increasing doses of spironolactone from 1 mg to 400 mg per day [i.e., 1 mg/day, 5 mg/day, 20 mg/day] was administered to a spironolactone-intolerant patient to treat cirrhosis-related ascites [P.A. 10 Greenberger et al, N. Eng. Reg. Allergy Proc., 7(4), 343-345 (Jul-Aug, 1986)]. It has been recognized that development of myocardial fibrosis is sensitive to circulating levels of both Angiotensin II and aldosterone, and that the aldosterone antagonist spironolactone prevents 15 myocardial fibrosis in animal models, thereby linking aldosterone to excessive collagen deposition [D. Klug et al, Am. J. Cardiol., 71(3), 46A-54A (1993)].

Spironolactone has been shown to prevent fibrosis in animal models irrespective of the development of left ventricular 20 hypertrophy and the presence of hypertension [C.G. Brilla et al, J. Mol. Cell. Cardiol., 25(5), 563-575 (1993)].

Spironolactone at a dosage ranging from 25 mg to 100 mg daily is used to treat diuretic-induced hypokalemia, when orally-administered potassium supplements or other 25 potassium-sparing regimens are considered inappropriate [Physicians' Desk Reference, 46th Edn., p. 2153, Medical Economics Company Inc., Montvale, N.J. (1992)].

Previous studies have shown that inhibiting ACE 30 inhibits the renin-angiotensin system by substantially complete blockade of the formation of Angiotensin II. Many ACE inhibitors have been used clinically to control hypertension. While ACE inhibitors may effectively control hypertension, side effects are common including chronic 35 cough, skin rash, loss of taste sense, proteinuria and neutropenia.

Moreover, although ACE inhibitors effectively block the formation of Angiotensin II, aldosterone levels are not well controlled in certain patients having cardiovascular diseases. For example, despite continued
5 ACE inhibition in hypertensive patients receiving captopril, there has been observed a gradual return of plasma aldosterone to baseline levels [J. Staessen et al, J. Endocrinol., 91, 457-465 (1981)]. A similar effect has been observed for patients with myocardial infarction
10 receiving zofenopril [C. Borghi et al, J. Clin. Pharmacol., 33, 40-45 (1993)]. This phenomenon has been termed "aldosterone escape". In a side-by-side treatment of two cohorts of rats, one cohort treated with spironolactone sub-cutaneously and the other cohort treated with
15 captopril, spironolactone was found to prevent fibrosis in the hypertensive-rat cohort [C.G. Brilla et al, J. Mol. Cell. Cardiol., 25, 563-575 (1993)].

Combinations of an aldosterone antagonist and an
20 ACE inhibitor have been investigated for treatment of heart failure. It is known that mortality is higher in patients with elevated levels of plasma aldosterone and that aldosterone levels increase as CHF progresses from RAAS activation. Routine use of a diuretic may further elevate
25 aldosterone levels. ACE inhibitors consistently inhibit angiotensin II production but exert only a mild and transient antialdosterone effect.

Combining an ACE inhibitor and spironolactone has been
30 suggested to provide substantial inhibition of the entire RAAS. For example, a combination of enalapril and a 25 mg daily dose of spironolactone has been administered to ambulatory patients with monitoring of blood pressure [P. Poncelet et al, Am. J. Cardiol., 65(2), 33K-35K (1990)].
35 In a 90-patient study, a combination of spironolactone in a range from 50 mg/day to 100 mg/day (average 73 mg/day) and captopril was administered and found effective to control refractory CHF without serious incidents of hyperkalemia

[U. Dahlstrom et al, Am. J. Cardiol., 71, 29A-33A (21 Jan 1993)]. Spironolactone dosage at 100 mg/day coadministered with an ACE inhibitor was reported to be highly effective in 13 of 16 patients afflicted with congestive heart failure, with a 25 mg/day to 50 mg/day spironolactone maintenance dosage given at trial completion to compensated patients being treated with an ACE inhibitor and loop diuretic [A.A. van Vliet et al, Am. J. Cardiol., 71, 21A-28A (21 Jan 1993)]. Clinical improvements have been reported for patients receiving a co-therapy of spironolactone and the ACE inhibitor enalapril, although this report mentions that controlled trials are needed to determine the lowest effective doses and to identify which patients would benefit most from combined therapy [F. Zannad, Am. J. Cardiol., 71(3), 34A-39A (1993)].

Summary of Drawing Figures

Fig. 1 shows urinary aldosterone levels at different rates of spironolactone administration (12.5 mg, 25 mg, 50mg, 75mg), as compared to placebo, co-administered with stable doses of ACE inhibitor and loop diuretic.

Fig. 2 shows plasma renin activity at different rates of spironolactone administration (12.5 mg, 25 mg, 50mg, 75mg), as compared to placebo, co-administered with stable doses of ACE inhibitor and loop diuretic.

Fig. 3 shows N-Terminal ANF levels at different rates of spironolactone administration (12.5 mg, 25 mg, 50mg, 75mg), as compared to placebo, co-administered with stable doses of ACE inhibitor and loop diuretic.

Fig. 4 shows changes in supine blood pressure at different rates of spironolactone administration (12.5 mg, 25 mg, 50mg, 75mg), as compared to placebo, co-administered with stable doses of ACE inhibitor and loop diuretic.

Fig. 5 shows changes in supine heart rate at different rates of spironolactone administration (12.5 mg, 25 mg, 50mg, 75mg), as compared to placebo, co-administered with stable doses of ACE inhibitor and loop diuretic.

Description of the Invention

Treatment or prevention of circulatory disorders, including cardiovascular disorders such as heart failure, hypertension and congestive heart failure, is provided by a combination therapy comprising a therapeutically-effective amount of an angiotensin converting enzyme ("ACE") inhibitor along with a therapeutically-effective amount of a spironolactone-type aldosterone receptor antagonist and a diuretic. Preferably, the spironolactone-type aldosterone receptor antagonist is administered in the combination therapy at a low dose, that is, at a dose lower than has been conventionally used in clinical situations.

The phrase "angiotensin converting enzyme inhibitor" ("ACE inhibitor") is intended to embrace an agent or compound, or a combination of two or more agents or compounds, having the ability to block, partially or completely, the rapid enzymatic conversion of the physiologically inactive decapeptide form of angiotensin ("Angiotensin I") to the vasoconstrictive octapeptide form of angiotensin ("Angiotensin II"). Blocking the formation of Angiotensin II can quickly affect the regulation of fluid and electrolyte balance, blood pressure and blood volume, by removing the primary actions of Angiotensin II.

Included in these primary actions of Angiotensin II are stimulation of the synthesis and secretion of aldosterone by the adrenal cortex and raising blood pressure by direct constriction of the smooth muscle of the arterioles.

5

The phrase "aldosterone receptor antagonist" embraces an agent or compound, or a combination of two or more of such agents or compounds, which agent or compound binds to the aldosterone receptor as a competitive inhibitor of the action of aldosterone itself at an aldosterone receptor site, such as typically found in the renal tubules, so as to modulate the receptor-mediated activity of aldosterone. Typical of such aldosterone receptor antagonists are spiro lactone-type compounds. The term "spiro lactone-type" is intended to characterize a steroidal structure comprising a lactone moiety attached to a steroid nucleus, typically at the steroid "D" ring, through a spiro bond configuration.

20

The phrase "combination therapy" (or "co-therapy"), in defining use of an ACE inhibitor agent, an aldosterone receptor antagonist agent and a diuretic, is intended to embrace administration of each agent in a sequential manner in a regimen that will provide beneficial effects of the drug combination, and is intended as well to embrace co-administration of these agents in a substantially simultaneous manner, such as in a single capsule having a fixed ratio of these active agents or in multiple, separate capsules for each agent.

30

The phrase "therapeutically-effective" is intended to qualify the amount of each agent for use in the combination therapy which will achieve the goal of improvement in cardiac sufficiency by reducing or preventing, for example, the progression of congestive heart failure, while avoiding adverse side effects

35

typically associated with each agent.

The phrase "low-dose amount", in characterizing a therapeutically-effective amount of the aldosterone
5 receptor antagonist agent in the combination therapy, is intended to define a quantity of such agent, or a range of quantity of such agent, that is capable of improving cardiac sufficiency while reducing or avoiding one or more aldosterone-antagonist-induced side effects, such as
10 hyperkalemia. A dosage of spironolactone which would accomplish the therapeutic goal of favorably enhancing cardiac sufficiency, while reducing or avoiding side effects, would be a dosage that substantially avoids inducing diuresis, that is, a substantially non-diuresis-effective dosage.

15 A preferred combination therapy would consist essentially of three active agents, namely, an ACE inhibitor agent, aldosterone receptor antagonist agent and a diuretic. For the ACE inhibitor and an ALDO antagonist
20 combination the agents would be used in combination in a weight ratio range from about 0.5-to-one to about twenty-to-one of the angiotensin converting enzyme agent to the aldosterone receptor antagonist agent. A preferred range of these two agents (ACE inhibitor-to-ALDO antagonist)
25 would be from about one-to-one to about fifteen-to-one, while a more preferred range would be from about one-to-one to about five-to-one, depending ultimately on the selection of the ACE inhibitor and ALDO antagonist. The diuretic agent may be present in a ratio range of 0.1-to-one to
30 about ten to one (ACE Inhibitor to diuretic).

Examples of ACE inhibitors which may be used in the combination therapy are shown in the following four categories.

35

A first group of ACE inhibitors consists of the following compounds: AB-103, ancovenin, benazeprilat, BRL-36378, BW-A575C, CGS-13928C, CL-242817, CV-5975, Equaten,

EU-4865, EU-4867, EU-5476, foroxymithine, FPL 66564, FR-900456, Hoe-065, I5B2, indolapril, ketomethylureas, KRI-1177, KRI-1230, L-681176, libenzapril, MCD, MDL-27088, MDL-27467A, moveltipril, MS-41, nicotianamine, pentopril,
5 phenacein, pivopril, rentiapril, RG-5975, RG-6134, RG-6207, RGH-0399, ROO-911, RS-10085-197, RS-2039, RS 5139, RS 86127,
RU-44403, S-8308, SA-291, spiraprilat, SQ-26900, SQ-28084, SQ-28370, SQ-28940, SQ-31440, Synecor, utibapril,
10 WF-10129, Wy-44221, Wy-44655, Y-23785, Yissum P-0154, zabicipril and zofenopril.

A second group of ACE inhibitors of interest consists of the following compounds: Asahi Brewery AB-47,
15 alatriopril, BMS 182657, Asahi Chemical C-111, Asahi Chemical C-112, Dainippon DU-1777, mixanpril, Prentyl, zofenoprilat and 1-(-(1-carboxy-6-(4-piperidinyl)hexyl) amino)-1-oxopropyl octahydro-1H-indole-2-carboxylic acid.

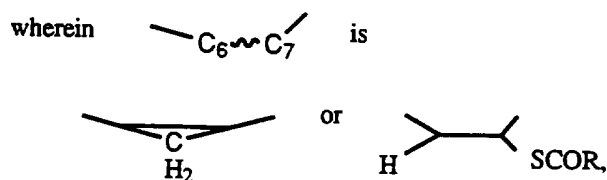
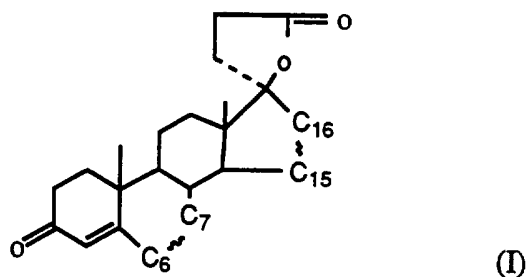
20 A third group of ACE inhibitors of greater interest consists of the following compounds: Bioproject BP1.137, Chiesi CHF 1514, Fisons FPL-66564, idrapril, Marion Merrell Dow MDL-100240, perindoprilat and Servier S-5590.

25 A fourth group of ACE inhibitors of highest interest consists of the following compounds: alacepril, benazepril, captopril, cilazapril, delapril, enalapril, enalaprilat, fosinopril, fosinoprilat, imidapril, lisinopril, perindopril, quinapril, ramipril, saralasin
30 acetate, temocapril, trandolapril, ceranapril, moexipril, quinaprilat and spirapril.

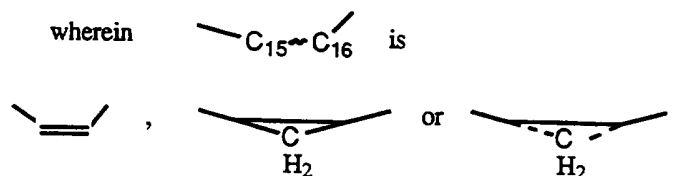
Many of these ACE inhibitors are commercially available, especially those listed in the fourth group,
35 above. For example, a highly preferred ACE inhibitor, captopril, is sold by E.R. Squibb & Sons, Inc., Princeton, N.J., now part of Bristol-Myers-Squibb, under the trademark

"CAPOTEN", in tablet dosage form at doses of 12.5 mg, 50 mg and 100 mg per tablet. Enalapril, or Enalapril Maleate, and Lisinopril are two more highly preferred ACE inhibitors sold by Merck & Co, West Point, Pa. Enalapril is sold under the trademark "VASOTEC" in tablet dosage form at doses of 2.5 mg, 5 mg, 10 mg and 20 mg per tablet. Lisinopril is sold under the trademark "PRINIVIL" in tablet dosage form at doses of 5 mg, 10 mg, 20 mg and 40 mg per tablet.

- 10 A family of spirolactone-type compounds of interest is defined by Formula I



- 15 wherein R is lower alkyl of up to 5 carbon atoms,
and



- 20 Lower alkyl residues include branched and unbranched groups, preferably methyl, ethyl and n-propyl.

Specific compounds of interest within Formula I are the following:

7 α -Acetylthio-3-oxo-4,15-androstadiene-[17(β -1')-
spiro-5']perhydrofuran-2'-one;

3-Oxo-7 α -propionylthio-4,15-androstadiene-[17((β -1')-
spiro-5')]perhydrofuran-2'-one;

5 6 β ,7 β -Methylene-3-oxo-4,15-androstadiene-[17((β -1')-
spiro-5')]perhydrofuran-2'-one;

15 α ,16 α -Methylene-3-oxo-4,7 α -propionylthio-4-
androstene[17(β -1')-spiro-5']perhydrofuran-2'-one;

6 β ,7 β ,15 α ,16 α -Dimethylene-3-oxo-4-androstene
10 [17(β -1')-spiro-5']perhydrofuran-2'-one;

7 α -Acetylthio-15 β ,16 β -Methylene-3-oxo-4-androstene-
[17(β -1')-spiro-5']perhydrofuran-2'-one;

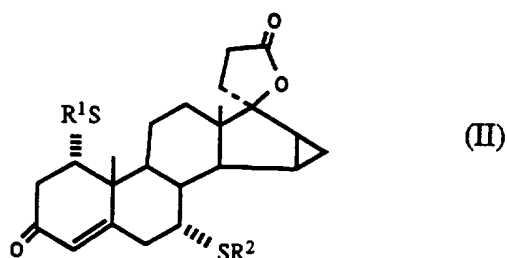
15 β ,16 β -Methylene-3-oxo-7 β -propionylthio-4-androstene-
[17(β -1')-spiro-5']perhydrofuran-2'-one; and

15 6 β ,7 β ,15 β ,16 β -Dimethylene-3-oxo-4-androstene-[17(β -
1')-spiro-5']perhydrofuran-2'-one.

Methods to make compounds of Formula I are described in
U.S. Patent No. 4,129,564 to Wiechart et al issued on 12
20 December 1978.

13

A second family of spirolactone-type compounds of interest is defined by Formula II:



5

wherein R^1 is C_{1-3} -alkyl or C_{1-3} acyl and R^2 is H or C_{1-3} -alkyl.

Specific compounds of interest within Formula II are the following:

10

1 α -Acetylthio-15 β ,16 β -methylene-7 α -methylthio-3-oxo-17 α -pregn-4-ene-21,17-carbolactone; and

15 β ,16 β -Methylene-1 α ,7 α -dimethylthio-3-oxo-17 α -pregn-4-ene-21,17-carbolactone.

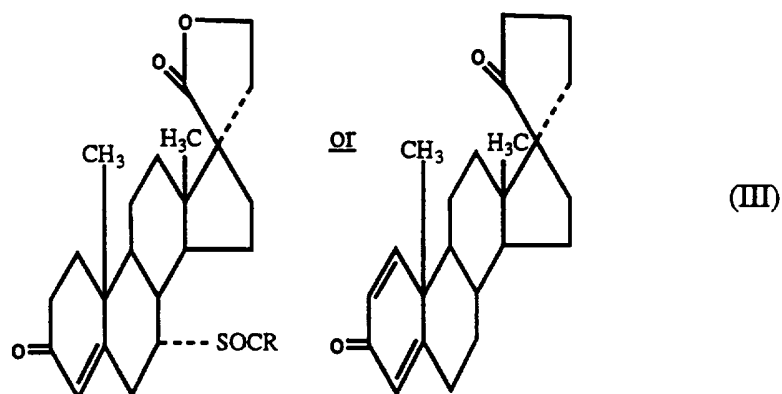
15

Methods to make the compounds of Formula II are described in U.S. Patent No. 4,789,668 to Nickisch et al which issued 6 December 1988.

20

A third family of spirolactone-type compounds of interest is defined by a structure of Formula III:

14



wherein R is lower alkyl, with preferred lower alkyl groups being methyl, ethyl, propyl and butyl. Specific compounds of interest include:

5

3 β ,21-dihydroxy-17 α -pregna-5,15-diene-17-carboxylic acid γ -lactone;

3 β ,21-dihydroxy-17 α -pregna-5,15-diene-17-carboxylic acid γ -lactone 3-acetate;

10

3 β ,21-dihydroxy-17 α -pregn-5-ene-17-carboxylic acid γ -lactone;

3 β ,21-dihydroxy-17 α -pregn-5-ene-17-carboxylic acid γ -lactone 3-acetate;

15

21-hydroxy-3-oxo-17 α -pregn-4-ene-17-carboxylic acid γ -lactone;

21-hydroxy-3-oxo-17 α -pregna-4,6-diene-17-carboxylic acid γ -lactone;

21-hydroxy-3-oxo-17 α -pregna-1,4-diene-17-carboxylic acid γ -lactone;

20

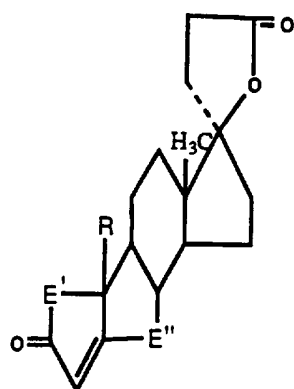
7 α -acylthio-21-hydroxy-3-oxo-17 α -pregn-4-ene-17-carboxylic acid γ -lactones; and

7 α -acetylthio-21-hydroxy-3-oxo-17 α -pregn-4-ene-

17-carboxylic acid γ -lactone.

Methods to make the compounds of Formula III are described
in U.S. Patent No. 3,257,390 to Patchett which issued 21
5 June 1966.

A fourth family of compounds of interest is
represented by Formula IV:



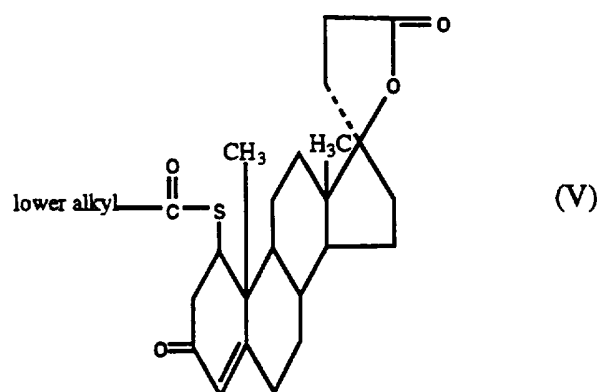
(IV)

10

wherein E' is selected from the group consisting of
ethylene, vinylene and (lower alkanoyl)thioethylene
radicals, E'' is selected from the group consisting of
ethylene, vinylene, (lower alkanoyl)thioethylene and (lower
15 alkanoyl)thiopropylene radicals; R is a methyl radical
except when E' and E'' are ethylene and (lower alkanoyl)
thioethylene radicals, respectively, in which case R is
selected from the group consisting of hydrogen and methyl
radicals; and the selection of E' and E'' is such that at
20 least one (lower alkanoyl)thio radical is present.

A preferred family of compounds within Formula IV
is represented by Formula V:

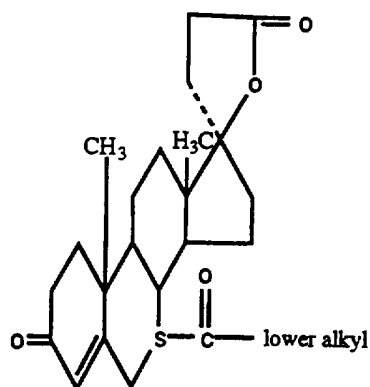
16



A more preferred compound of Formula V is
 1-acetylthio-17 α -(2-carboxyethyl)-17 β -hydroxy-androst-4-en-
 5 3-one lactone.

17

Another preferred family of compounds within Formula IV is represented by Formula VI:



(VI)

5

More preferred compounds within Formula VI include the following:

7 α -acetylthio-17 α -(2-carboxyethyl)-17 β -hydroxy-
10 androst-4-en-3-one lactone;

7 β -acetylthio-17 α -(2-carboxyethyl)-17 β -hydroxy-
androst-4-en-3-one lactone;

1 α ,7 α -diacetylthio-17 α -(2-carboxyethyl)-17 β -
hydroxy-androsta-4,6-dien-3-one lactone;

15 7 α -acetylthio-17 α -(2-carboxyethyl)-17 β -hydroxy-
androsta-1,4-dien-3-one lactone;

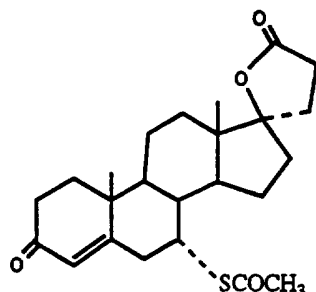
7 α -acetylthio-17 α -(2-carboxyethyl)-17 β -hydroxy-
19-norandrost-4-en-3-one lactone; and

7 α -acetylthio-17 α -(2-carboxyethyl)-17 β -hydroxy-
20 6 α -methylandrost-4-en-3-one lactone;

In Formula IV-VI, the term "alkyl" is intended to embrace linear and branched alkyl radicals containing one to about eight carbons. The term "(lower alkanoyl)thio"

25 embraces radicals of the formula lower alkyl $\overset{\text{O}}{\parallel} \text{C}-\text{S}$.

Of particular interest is the compound spironolactone having the following structure and formal name:



"spironolactone": 17-hydroxy-7 α -mercapto-3-oxo-17 α -pregn-4-ene-21-carboxylic acid γ -lactone acetate

10 Methods to make compounds of Formula IV-VI are described in U.S. Patent No. 3,013,012 to Cella et al which issued 12 December 1961. Spironolactone is sold by G.D. Searle & Co., Skokie, Illinois, under the trademark "ALDACTONE", in tablet dosage form at doses of 25 mg, 50 mg
15 and 100 mg per tablet.

 The diuretic agent which is used with the combination of ACE inhibitor and aldosterone receptor antagonist may be selected from several known classes, such
20 as thiazides and related sulfonamides, potassium-sparing diuretics, loop diuretics and organic mercurial diuretics.

 Examples of thiazides are bendroflumethiazide, benzthiazide, chlorothiazide, cyclothiazide,
25 hydrochlorothiazide, hydroflumethiazide, methyclothiazide, polythiazide and trichlormethiazide.

 Examples of sulfonamides related to thiazides are chlorthalidone, quinethazone and metolazone.

30 Examples of potassium-sparing diuretics are triameterene and amiloride.

Examples of loop diuretics, i.e., diuretics acting in the ascending limb of the loop of Henle of the kidney, are furosemide and ethynacrylic acid.

5

Examples of organic mercurial diuretics are mercaptomerin sodium, merethoxylline, procaine and mersalyl with theophylline.

10

Biological Evaluation

Human Clinical Trials

15

A triple therapy of ACE inhibitor, spironolactone and loop diuretic was evaluated in humans as described in the following clinical trials.

Patients: Two-hundred fourteen (214) patients with symptomatic heart failure had an ejection fraction $\leq 35\%$, a history of New York Heart Association (NYHA) functional classification III-IV six months prior to enrollment, and current classification II-IV were randomized among five treatment groups. Patients were assigned to receive either spironolactone 12.5 mg (41 patients), 25 mg (45 patients), 50 mg (47 patients), 75 mg (41 patients), or placebo (40 patients) once a day for 12 weeks. Two patients that were randomized failed to take the study medication and were excluded from the analysis. All patients were taking a stable dose of ACE inhibitor, loop diuretic, and optional digitalis for 30 days prior to the first dose of study medication. Potassium supplement therapy that was stable for 14 days prior to the first dose of study medication was also allowed. Informed consent was obtained from all patients, and the protocol was approved by each ethical committee. At enrollment all patients had normal serum potassium values (<5.5 mmol/L) and creatinine values of

≤2.0 mg/dL or ≤180 mmol/L. Patients were excluded from enrollment if they: (1) were diagnosed with either an acute life-threatening disease (included patients with automatic implantable cardioverter/ defibrillator),
5 valvular disease, unstable angina, insulin-dependent diabetes, cancer (without a reoccurrence within the last five years), or primary hepatic failure; (2) were on a waiting list for a heart transplant or experienced a myocardial infarction 30 days prior to the first dose of
10 study medication; (3) had laboratory values for hematology or biochemistry considered abnormal and clinically significant prior to the first dose of study medication; (4) received a potassium spacing diuretic within 30 days prior to the first dose of study medication.; (5) were
15 receiving, on a regular basis, either non-steroidal anti-inflammatory drugs or aspirin >325 mg/day, steroids, dopamine agonists or antagonists, insulin or heparin; (6) were on any investigational medication within 30 days of the first dose of medication.

20

Study Design: This was a multinational, double-blind, randomized, parallel group study.

Laboratory Measurements: The following information was
25 obtained from each patient at baseline:

1. Concurrent medication within the past 30 days.
2. 12-lead ECG
3. Cardiac assessments that included blood pressure, pulse,
30 sodium retention score (general assessment of a patient's edematous state was derived from the summation of scores obtained from Table I), NYHA classification, and
4. Signs and symptoms within the past 30 days.

35

Table I
Sodium Retention Score

5	<u>Parameters</u>	<u>Grade</u>	<u>Assessment</u>
	Rales	0	Absent
		1	In lower 1/3 of lungs
		2	In lower 2/3 of lungs
10		3	In all lung fields
	Peripheral	0	Absent
	Pitting Edema	1	Trace
		2	Limited to ankles
15		3	Not limited to ankles
		4	Anasarca
	Weight Change	-1	Decreased
		0	Unchanged
20		1	Increased
	Hepatomegaly	0	Absent
		1	Present
	S3 Gallop	0	Absent
25		1	Present
	Increased Jugular	0	Absent
	Venous Pressure	1	Present
30	The following laboratory values were obtained at the pretreatment visit:		
35	Hematology:	White blood cell count (WBC), hematocrit, hemoglobin, platelet count.	
40	Biochemistry:	Creatinine, potassium, AST, SGOT, urinary sodium/ potassium ratio, bicarbonate, calcium, chloride, creatinine, creatinine clearance, magnesium, glucose, urea, uric acid.	
45	Neurohormones:	Plasma renin activity, pro-atrial natriuretic factor, urinary aldosterone.	
50	Blood and urine samples were centrally analyzed at SciCor Laboratories. Laboratory values for urinary aldosterone and renin levels were done at the Ohio State University Laboratory in Columbus, Ohio. Pro-atrial natriuretic factor samples were evaluated at the University of Oslo Laboratory in Oslo, Norway. Patients were evaluated 9 days		

after beginning study medication. Documented changes in concurrent medications, signs and symptoms and drug compliance were recorded. These procedures were repeated at Week 4 and Week 8 visits. Patient information and
5 procedures on the final visit (Week 12) was identical to the pre-treatment visit.

Statistical Analysis: Analysis of cardiac assessment changes in patient therapy and vital signs were performed
10 for both the Intent-to-Treat (ITT) and evaluable patient groups. Analysis of demographic variables, adverse events and clinical laboratory values were performed in the ITT group. For each efficacy variable, results of each visit were examined separately. An appropriate trend test was
15 used to test for overall dose-response. Pair-wise comparisons were made for each active dose to placebo. Significant levels for pair-wise comparisons were adjusted using the Hochberg-Bonferromi method to maintain the overall Type I error rate. All statistical methods were
20 two-sided.

Recruitment: Two-hundred and fourteen patients were recruited from 22 study sites in eleven countries.

25 **Patient Characteristics:** Patient demographic, vital signs, and cardiac status at baseline are summarized in Table II.

TABLE II
Patient Demographics

Demographic	Spironolactone 12.5 mg/d	Spironolactone 25 mg/d	Spironolactone 50 mg/d	Spironolactone 75 mg/d	Placebo	p- Value
Age (years)	63 ±12	61 ±9	62 ±13	62 ±13	61 ±12	N.S.
Caucasian/other (%)	93/7	98/2	93/7	88/12	97/3	N.S.
Male/female (%)	78/22	82/18	74/26	88/12	83/18	N.S.
Vital Signs						
Weight (kg)	74	75	73	78	73	N.S.
Blood pressure (mmHg)						
Systolic	121	120	121	125	121	N.S.
Diastolic	76	76	75	81	74	N.S.
Pulse (bpm)	76	74	76	74	71	N.S.
Cardiac Status						
NYHA (%)						
II	63	60	43	49	38	
III	34	38	55	49	60	
IV	2	2	2	2	2	N.S.
Sodium retention score						
Mean value	1.54	1.62	1.64	1.61	1.78	N.S.
ACE-I (Mean dose)						
Captopril (mg)	57.3	57.5	69.7	59.4	65.4	N.S.
Enalapril (mg)	16.4	13.4	14.5	16.3	10.8	N.S.
Loop Diuretic (Mean dose)						
Furosemide (mg)	58.8	82.8	76.9	84.9	63.2	N.S.
Digoxin (%)						
Potassium supplement (%)	43.9	37.8	34.0	39.0	30.0	N.S.

Patients ranged in age from 26 to 83 years (mean = 60), 81% were male, 94% were Caucasian. At baseline 51% of the patients were NYHA Class II, 47% were Class III.

With respect to sodium retention score, a statistically significant dose response was seen at Day 9 with higher doses showing more reduction in sodium retention score ($p = 0.019$). However, this effect was not seen at later visits ($p > 0.20$). There was an improvement in NYHA Class placebo group and in all the spironolactone groups. Although a trend toward improvement in the spironolactone group was observed, the difference was not statistically significant.

Changes in Patient Therapy: The treatment groups did not differ significantly with respect to changes in dose of ACE inhibitor, digitalis or potassium supplements at any visit ($p \geq 0.11$). The treatment groups did differ significantly with respect to changes in loop diuretic therapy only at Week 8 ($p = 0.004$) in that more patients on the higher doses of spironolactone had decreases in the loop diuretic dose compared to the placebo group. This pattern was not observed at Week 12.

Changes in Vital Signs: Changes from baseline in vital signs at Week 12 are summarized in Table III.

TABLE III

Mean Change in Weight and Vital Signs from Baseline to Week 12

	Spiro nolactone 12.5 mg/d	Spiro nolactone 25 mg/d	Spiro nolactone 50 mg/d	Spiro nolactone 75 mg/d	Placebo	p- Value
Weight	0.59 (3.00)	-0.16 (3.02)	0.62 (2.05)	-0.81 (2.70)	0.11 (2.46)	0.109
Supine systolic BP	1.84 (11.82)	-4.46 (13.97)	-7.04 (15.83)	-5.68 (15.62)	0.22 (13.45)	0.036
Supine diastolic BP	-0.19 (9.13)	-2.74 (9.57)	-5.11 (11.11)	-5.91 (9.05)	1.78 (7.84)	0.014
Supine pulse (BPM)	-3.70 (9.56)	-1.40 (10.00)	-3.21 (11.27)	-1.07 (13.79)	1.42 (9.69)	0.422

B

At all visits the 25 mg, 50 mg, and 75 mg groups had decreases in mean systolic and diastolic blood pressure, while the placebo group had increases in mean systolic and diastolic blood pressure (both standing and supine). Dose response with respect to standing and supine diastolic blood pressure was statistically significant for all visits ($p \leq 0.002$). Dose response with respect to standing and supine blood pressure was statistically significant at Week 4, Week 8, and Week 12 ($p \leq 0.033$), but not at Day 9 ($p \geq 0.12$). No significant between-treatment differences in change from baseline in pulse were observed at any visit (p -values ≥ 0.136). A statistically significant dose response with greater decreases in pulse in the supine position at higher doses was observed at Week 4 (p -value = 0.045). Spironolactone doses of 25 and 50 mg were also significantly different from placebo (p -values ≤ 0.043) (See Fig. 1). At Day 9 and Week 4 visits, there was a statistically significant dose response with respect to changes from baseline in body weight in that patients in the 75 mg dose group experienced more weight loss than other patients. This dose response was not observed at later visits ($p \geq 0.062$).

25

Clinical Laboratory Values: Table IV contains details of the different clinical laboratory values that showed statistically significant treatment differences with respect to mean changes at Week 12 visit compared with their respective baseline value.

30

TABLE IV
Week 12 Mean Change

	Spiro 12.5 mg/d	Spiro 25 mg/d	Spiro 50 mg/d	Spiro 75 mg/d	P- Placebo Value
Urinary aldosterone (nmol/D)	4.21	4.27	8.11	11.13	0.76 0.002
N-Terminal ANF (pmol/L)	-287.30	-294.60	-351.30	-370.60	54.50 0.022
PRA (NgAngl/L/s)	9.90	9.33	13.18	10.23	0.50 0.002
Hematocrit (%)	0.00	-0.02	-0.02	-0.03	0.00 0.002
Hemoglobin (mmol/L/Fe)	0.12	-0.20	-0.31	-0.46	0.00 0.005
Potassium (mmol/L)	0.18	0.37	0.51	0.58	-0.10 0.001
Creatinine (umol/L)	6.83	9.30	14.06	21.90	-1.96 0.001
Sodium (mmol/L)	-1.61	-1.85	-2.52	-3.37	-0.03 0.001

Urinary Aldosterone (See Fig. 1): Urinary aldosterone was determined only for baseline and the 12 week visit.

5 Urinary aldosterone excretion showed mean increases from baseline in all treatment groups ($P \leq 0.012$). Greater increases were seen at higher doses of spironolactone ($p = 0.002$). All pair-wise comparisons between active treatment and placebo were statistically significant ($p \leq 0.009$).

10

Plasma Renin Activity (PRA) (See Fig. 2): A

statistically significant dose-response with respect to change from baseline in PRA was seen at Day 9, Week 4 and
15 Week 12 ($P \leq 0.001$) with higher doses of spironolactone associated with greater increases in PRA. PRA was not measured at Week 8.

N-Terminal Atrial Natriuretic Factor (ANF) (See

20 **Fig. 3):** All active treatments showed decreases from baseline at all treatment visits. Dose-response was statistically significant at Day 9 ($p = 0.048$), Week 4 ($p = 0.005$), and Week 12 ($p = 0.008$). ANF was not measured at Week 8. In comparisons the 50 mg dose group differed
25 significantly from placebo at Week 4 ($p = 0.009$) and Week 12 ($p = 0.006$), while the 75 mg dose group differed significantly from placebo at Week 12 only ($p = 0.007$).

Hematocrit and Hemoglobin: At Day 9 visit a

30 statistically significant mean value difference between placebo and the different active treatments was observed with lower values for the placebo group than the active treatments ($p < 0.001$). At Week 12 a reverse statistically significant difference was observed with lower levels for
35 the active treatment groups for hematocrit ($p = 0.002$) and hemoglobin (0.005).

Serum Potassium: A statistically significant dose-response with respect to change from baseline in serum potassium was seen at all treatment period visits ($p < 0.001$). Higher doses of spironolactone were associated with larger increases in potassium. All doses of active treatment had significantly higher serum potassium levels relative to baseline than placebo ($p \leq 0.034$).

Incidence of Hyperkalemia

Treatment: Placebo	Spironolactone 12.5 mg/d	Spironolactone 25 mg/d	Spironolactone 50 mg/d	Spironolactone 75 mg/d
Patients (%)2(5%)	2(5%)	6(13%)	9(20%)	10(24%)

Predictors of Hyperkalemia: Seven possible predictors of hyperkalemia (potassium ≥ 5.5 mmol/L) were included in a step-wise Cox regression analysis: randomized treatment (treated as a categorical variable), age, baseline NYHA class, baseline serum potassium, baseline PRA, baseline creatinine, baseline urinary aldosterone, and type and dose of ACE-I. Besides the dose of spironolactone, the following predictors of hyperkalemia were statistically significant in the step-wise regression analysis: type of ACE-I (captopril versus other), baseline serum creatinine, and baseline serum potassium. Results are summarized as follows:

<u>Factor</u>	<u>p-value</u>	<u>Risk Ratio</u>
Captopril vs other ACE-I	0.013	0.318
Serum Creatinine > normal	0.038	2.72
Baseline Potassium > median	0.040	2.32

In this analysis, the risk ratio can be thought of as the probability that the patient with the risk factor will develop hyperkalemia, relative to the probability that a patient without the risk factor will develop it. (For example, patients on captopril are about one-third as likely to develop hyperkalemia as a patient on another ACE-I.)

Risk ratios relative to placebo for the various doses of spironolactone are:

	<u>Dose</u>	<u>p-value</u>	<u>Risk Ratio</u>
5	Spironolactone 12.5 mg	0.98	1.02
	Spironolactone 25 mg	0.19	2.91
	Spironolactone 50 mg	0.034	5.32
	Spironolactone 75 mg	0.016	6.66

After adjusting for the above factors, other predictors
 10 included in the step-wise regression analysis were not
 significant (p -values ≥ 0.07). However, the following
 additional factor was significantly related to the
 development of hyperkalemia when considered apart from
 other predictors except the dose of spironolactone.

15	<u>Factor</u>	<u>p-value</u>	<u>Risk Ratio</u>
	High ACE-I Dose	0.050	2.93

Serum Magnesium: Change from baseline in serum
 20 magnesium showed a statistically significant dose-response
 at Day 9 and Week 4 ($p \leq 0.048$), with more patients in the
 placebo group showing decreases in serum magnesium.
 However, this effect was not seen at later visits (p
 ≥ 0.083).

25 **Adverse Effects:** Table V summarizes the twelve most
 common adverse events by different treatment groups. Only
 one symptom, hyperkalemia, showed a clear dose-response in
 term of incidence ($p = 0.001$).

Table V
Incidence of Adverse Events
Spironolactone Dose-Ranging Study
Intent-to-Treat Cohort
(Top Twelve Events)

Adverse Events	Treatment Group (Percentage of Patients)					
	Spironolactone 12.5 mg	Spironolactone 25 mg	Spironolactone 50 mg	Spironolactone 75 mg	Placebo	Total
Dyspnea	22.0	15.6	26.1	24.4	30.0	23.5
Angina Pectoris	19.5	20.0	8.7	14.6	17.5	16.0
Dizziness	12.2	13.3	13.0	17.1	15.0	14.1
Fatigue	12.2	13.3	15.2	14.6	15.0	14.1
Nausea	2.4	17.8	6.5	19.5	12.5	11.7
Diarrhea	4.9	22.2	8.7	14.6	5.0	11.3
Abdominal Pain	7.3	8.9	13.0	7.3	17.5	10.8
Headache	9.8	2.2	15.2	7.3	20.0	10.8
Hyperkalemia	2.4	8.9	15.2	19.5	2.5	9.9
URT Infection	4.9	11.1	8.7	2.4	12.5	8.0
Arthralgia	4.9	4.4	8.7	4.9	7.5	6.1
Coughing	4.9	2.2	4.3	2.4	12.5	5.2

A breakdown of the hospitalizations is as follows:

	<u>Treatment</u>	<u>Placebo</u>	<u>12.5 mg</u>	<u>25 mg</u>	<u>50 mg</u>	<u>75 mg</u>	<u>P</u>
5	Patients (%)	5(12.5%)	3(7.3%)	3(6.6%)	13(27.6%)	6(14.6%)	N.S.

No deaths were reported during the drug treatment period. Three patients died within 30 days after the study was completed. These three patients were previously at the 50

10 mg dose.

Administration of the angiotensin converting enzyme inhibitor, the aldosterone receptor antagonist and diuretic may take place sequentially in separate

15 formulations, or may be accomplished by simultaneous administration in a single formulation or separate formulations. Administration may be accomplished by oral route, or by intravenous, intramuscular or subcutaneous injections. The formulation may be in the form of a bolus,

20 or in the form of aqueous or non-aqueous isotonic sterile injection solutions or suspensions. These solution and suspensions may be prepared from sterile powders or granules having one or more pharmaceutically-acceptable carriers or diluents, or a binder such as gelatin or

25 hydroxypropyl-methyl cellulose, together with one or more of a lubricant, preservative, surface-active or dispersing agent.

For oral administration, the pharmaceutical

30 composition may be in the form of, for example, a tablet, capsule, suspension or liquid. The pharmaceutical composition is preferably made in the form of a dosage unit containing a particular amount of the active ingredient. Examples of such dosage units are tablets or capsules. The

35 ACE inhibitor may be present in an amount from about 1 to 200 mg, preferably from about 2 to 150 mg, depending upon the specific ACE inhibitor selected. A suitable daily dose for a mammal may vary widely depending on the condition of the patient and other factors. The ALDO antagonist may be

40 present in an amount of from about 1 to 400 mg, preferably

from about 2 to 150 mg, depending upon the specific ALDO antagonist compound selected and the specific disease state being targetted for the combination therapy.

5 For disease states which require prevention, reduction or treatment of a cardiovascular disease state without incidence of hyperkalemia, for example, the ALDO antagonist component, typically spironolactone, will be present in the combination therapy in an amount in a range
10 from about 1 mg to about 25 mg per dose. A preferred range for spironolactone would be from about 5 mg to 15 mg per dose. More preferably, the spironolactone dosage would be in a range from about 10 mg to 15 mg per dose per day.

15 The diuretic agent may be present in a range from about 1 mg to about 200 mg, preferably from 5 mg to 150 mg, depending upon the diuretic selected and the disease state targetted.

20 Examples of various fixed combinations of ACE inhibitor and ALDO antagonist, for use with a diuretic in the "triple therapy" of the invention, are as follow:

<u>ACE Inhibitor</u>		<u>ALDO Antagonist</u>
Captopril (mg) ¹	Enalapril (mg) ²	Spironolactone (mg) ²
12.5 to 25	5 to 15	5
12.5 to 25	5 to 15	7.5
12.5 to 25	5 to 15	10
12.5 to 25	5 to 15	12.5
30 12.5 to 25	5 to 15	15
12.5 to 25	5 to 15	17.5
12.5 to 25	5 to 15	20
12.5 to 25	5 to 15	22.5

¹Dose given 1, 2 or 3 times per day

35 ²Dose given once per day

The active ingredients may also be administered by injection as a composition wherein, for example, saline, dextrose or water may be used as a suitable carrier.

The dosage regimen for treating a disease condition with the combination therapy of this invention is selected in accordance with a variety of factors, including the type, age, weight, sex and medical condition of the patient, the severity of the disease, the route of administration, and the particular compound employed, and thus may vary widely.

For therapeutic purposes, the active components of this combination therapy invention are ordinarily combined with one or more adjuvants appropriate to the indicated route of administration. If administered per os, the components may be admixed with lactose, sucrose, starch powder, cellulose esters of alkanolic acids, cellulose alkyl esters, talc, stearic acid, magnesium stearate, magnesium oxide, sodium and calcium salts of phosphoric and sulfuric acids, gelatin, acacia gum, sodium alginate, polyvinylpyrrolidone, and/or polyvinyl alcohol, and then tableted or encapsulated for convenient administration. Such capsules or tablets may contain a controlled-release formulation as may be provided in a dispersion of active compound in hydroxypropylmethyl cellulose. Formulations for parenteral administration may be in the form of aqueous or non-aqueous isotonic sterile injection solutions or suspensions. These solutions and suspensions may be prepared from sterile powders or granules having one or more of the carriers or diluents mentioned for use in the formulations for oral administration. The components may be dissolved in water, polyethylene glycol, propylene glycol, ethanol, corn oil, cottonseed oil, peanut oil, sesame oil, benzyl alcohol, sodium chloride, and/or various buffers. Other adjuvants and modes of administration are well and widely known in the pharmaceutical art.

Pharmaceutical compositions for use in the treatment methods of the invention may be administered in oral form or by intravenous administration. Oral administration of the combination therapy is preferred. Dosing for oral administration may be with a regimen

calling for single daily dose, or for a single dose every other day, or for multiple, spaced doses throughout the day. The active agents which make up the combination therapy may be administered simultaneously, either in a combined dosage form or in separate dosage forms intended for substantially simultaneous oral administration. The active agents which make up the combinatin therapy may also be administered sequentially, with either active component being administered by a regimen calling for multi-step ingestion. Thus, a regimen may call for sequential administration of the active agents with spaced-apart ingestion of the separate, active agents. The time period between the multiple ingestion steps may range from a few minutes to several hours, depending upon the properties of each active agent such a potency, solubility, bioavailability, plasma half-life and kinetic profile of the agent, as well as depending upon the age and condition of the patient. The active agents of the combined therapy whether administered simultaneously, substantially simultaneously, or sequentially, may involve a regimen calling for administration of one active agent by oral route and the other active agent by intravenous route. Whether the active agents of the combined therapy are administered by oral or intravenous route, separately or together, each such active agent will be contained in a suitable pharmaceutical formulation of pharmaceutically-acceptable excipients, diluents or other formulations components. Examples of suitable pharmaceutically-acceptable formulations containing the active components for oral administration are given below. Even though such formulations list both active agents together in the same recipe, it is appropriate for such recipe to be utilized for a formulation containing one of the active components.

Example 1

An oral dosage may be prepared by screening and then mixing together the following list of ingredients in the amounts indicated. The dosage may then be placed in a hard

gelatin capsule.

	<u>Ingredients</u>	<u>Amounts</u>
	captopril	62.0 mg
5	spironolactone	12.5 mg
	furosemide	73.9 mg
	magnesium stearate	10 mg
	lactose	100 mg

10 Example 2

An oral dosage may be prepared by mixing together granulating with a 10% gelatin solution. The wet granules are screened, dried, mixed with starch, talc and stearic acid, screened and compressed into a tablet.

	<u>Ingredients</u>	<u>Amounts</u>
	captopril	62.0 mg
	spironolactone	12.5 mg
20	furosemide	73.9 mg
	calcium sulfate dihydrate	100 mg
	sucrose	15 mg
	starch	8 mg
	talc	4 mg
25	stearic acid	2 mg

Example 3

An oral dosage may be prepared by screening and then mixing together the following list of ingredients in the amounts indicated. The dosage may then be placed in a hard gelatin capsule.

	<u>Ingredients</u>	<u>Amounts</u>
35	enalapril	14.3 mg
	spironolactone	12.5 mg
	furosemide	73.9 mg
	magnesium stearate	10 mg
	lactose	100 mg

Example 4

An oral dosage may be prepared by mixing together granulating with a 10% gelatin solution. The wet granules
5 are screened, dried, mixed with starch, talc and stearic acid, screened and compressed into a tablet.

	<u>Ingredients</u>	<u>Amounts</u>
10	enalapril	14.3 mg
	spironolactone	12.5 mg
	furosemide	73.9 mg
	calcium sulfate dihydrate	100 mg
	sucrose	15 mg
15	starch	8 mg
	talc	4 mg
	stearic acid	2 mg

Although this invention has been described with
20 respect to specific embodiments, the details of these embodiments are not to be construed as limitations.

What Is Claimed Is:

1. A combination comprising a therapeutically-effective amount of an angiotensin converting enzyme
5 inhibitor, an aldosterone receptor antagonist and a diuretic, said aldosterone receptor antagonist being present in an amount which is therapeutically effective to antagonize a physiological effect of aldosterone but which amount is not sufficient for said aldosterone receptor
10 antagonist to induce an adverse side effect.

2. The combination of Claim 1 wherein said aldosterone receptor antagonist is a spiro lactone-type compound.
15

3. The combination of Claim 2 wherein said spiro lactone-type compound is spironolactone.

4. The combination of Claim 1 wherein
20 angiotensin converting enzyme inhibitor is selected from the group consisting of alacepril, benazepril, captopril, cilazapril, delapril, enalapril, enalaprilat, fosinopril, fosinoprilat, imidapril, lisinopril, perindopril, quinapril, ramipril, saralasin acetate, temocapril,
25 trandolapril, ceranapril, moexipril, quinaprilat, spirapril, Bioproject BP1.137, Chiesi CHF 1514, Fisons FPL-66564, idrapril, Marion Merrell Dow MDL-100240, perindoprilat and Servier S-5590.

30 5. The combination of Claim 4 wherein said angiotensin converting enzyme inhibitor is selected from the group consisting of alacepril, benazepril, captopril, cilazapril, delapril, enalapril, enalaprilat, fosinopril, fosinoprilat, imidapril, lisinopril, perindopril,
35 quinapril, ramipril, saralasin acetate, temocapril, trandolapril, ceranapril, moexipril, quinaprilat and spirapril.

6. The combination of Claim 1 further characterized by said angiotensin converting enzyme inhibitor and said aldosterone receptor antagonist being present in said combination in a weight ratio range from about 0.1-to-one to about twenty-five-to-one of said angiotensin converting enzyme inhibitor to said aldosterone receptor antagonist.

7. The combination of Claim 6 wherein said weight ratio range is from about 0.5-to-one to about fifteen-to-one.

8. The combination of Claim 7 wherein said weight ratio range is from about 0.5-to-one to about five-to-one.

9. A triple therapy for treating cardiovascular disorders in a subject afflicted with or susceptible to multiple cardiovascular disorders, wherein said triple therapy comprises administering a therapeutically-effective amount of an angiotensin converting enzyme inhibitor and a diuretic agent, and administering an aldosterone receptor antagonist in an amount therapeutically effective to antagonize aldosterone but insufficient to induce an adverse side effect.

10. The triple therapy of Claim 9 wherein said subject is afflicted with or susceptible to heart failure and said subject further requires avoidance of the incidence of hyperkalemia.

11. The triple therapy of Claim 10 wherein said subject is further susceptible to congestive heart failure.

12. The triple therapy of Claim 10 wherein said subject is further susceptible to hypertension.

13. The triple therapy of Claim 10 further characterized by administering said angiotensin converting enzyme inhibitor, said aldosterone receptor antagonist and said diuretic in a sequential manner.

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14. The triple therapy of Claim 10 further characterized by administering said angiotensin converting enzyme inhibitor and said aldosterone receptor antagonist in a substantially simultaneous manner.

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15. The triple therapy of Claim 9 wherein said aldosterone receptor antagonist is a spiro lactone-type compound.

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16. The triple therapy of Claim 15 wherein said spiro lactone-type compound is spironolactone.

17. The triple therapy of Claim 9 wherein said angiotensin converting enzyme inhibitor is selected from the group consisting of alacepril, benazepril, captopril, cilazapril, delapril, enalapril, enalaprilat, fosinopril, fosinoprilat, imidapril, lisinopril, perindopril, quinapril, ramipril, saralasin acetate, temocapril, trandolapril, ceranapril, moexipril, quinaprilat, spirapril, Bioproject BP1.137, Chiesi CHF 1514, Fisons FPL-66564, idrapril, Marion Merrell Dow MDL-100240, perindoprilat and Servier S-5590.

18. The triple therapy of Claim 17 wherein said angiotensin converting enzyme inhibitor is selected from the group consisting of alacepril, benazepril, captopril, cilazapril, delapril, enalapril, enalaprilat, fosinopril, fosinoprilat, imidapril, lisinopril, perindopril, quinapril, ramipril, saralasin acetate, temocapril, trandolapril, ceranapril, moexipril, quinaprilat and spirapril.

19. The triple therapy of Claim 9 further characterized by said angiotensin converting enzyme inhibitor and said aldosterone receptor antagonist being used in said co-therapy in a weight ratio range from about 0.1-to-one to about twenty-five-to-one of said angiotensin converting enzyme inhibitor to said aldosterone receptor antagonist.

20. The triple therapy of Claim 19 wherein said weight ratio range is from about 0.5-to-one to about fifteen-to-one.

21. The triple therapy of Claim 20 wherein said weight ratio range is from about 0.5-to-one to about five-to-one.

22. The triple therapy of Claim 9 wherein said angiotensin converting enzyme inhibitor is captopril, in a dose range from about 30 mg to about 80 mg per dose, or is enalapril in a dose range from about 5 mg to about 25 mg per dose.

23. The triple therapy of Claim 22 wherein said aldosterone receptor antagonist is spironolactone in a dose range from about 1 mg to less than 25 mg per dose.

24. The triple therapy of Claim 9 wherein said diuretic agent is selected from the group consisting of thiazides and related sulfonamides, potassium-sparing diuretics, loop diuretics and organic mercurial diuretics.

25. The triple therapy of Claim 9 wherein said diuretic agent is selected from the group consisting of bendroflumethiazide, benzthiazide, chlorothiazide, hydrochlorothiazide, hydroflumethiazide, methyclothiazide, polythiazide, trichlormethiazide, quinethazone, metolazone, triameterene and amiloride.

FIG. 1

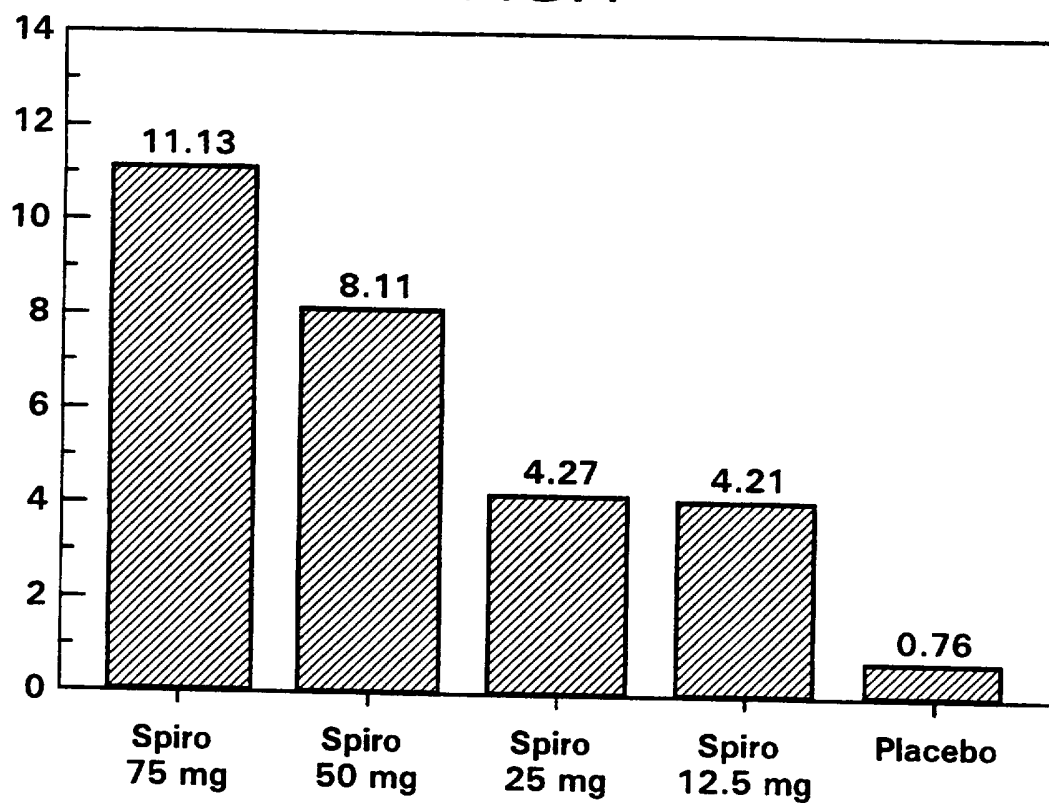


FIG. 2

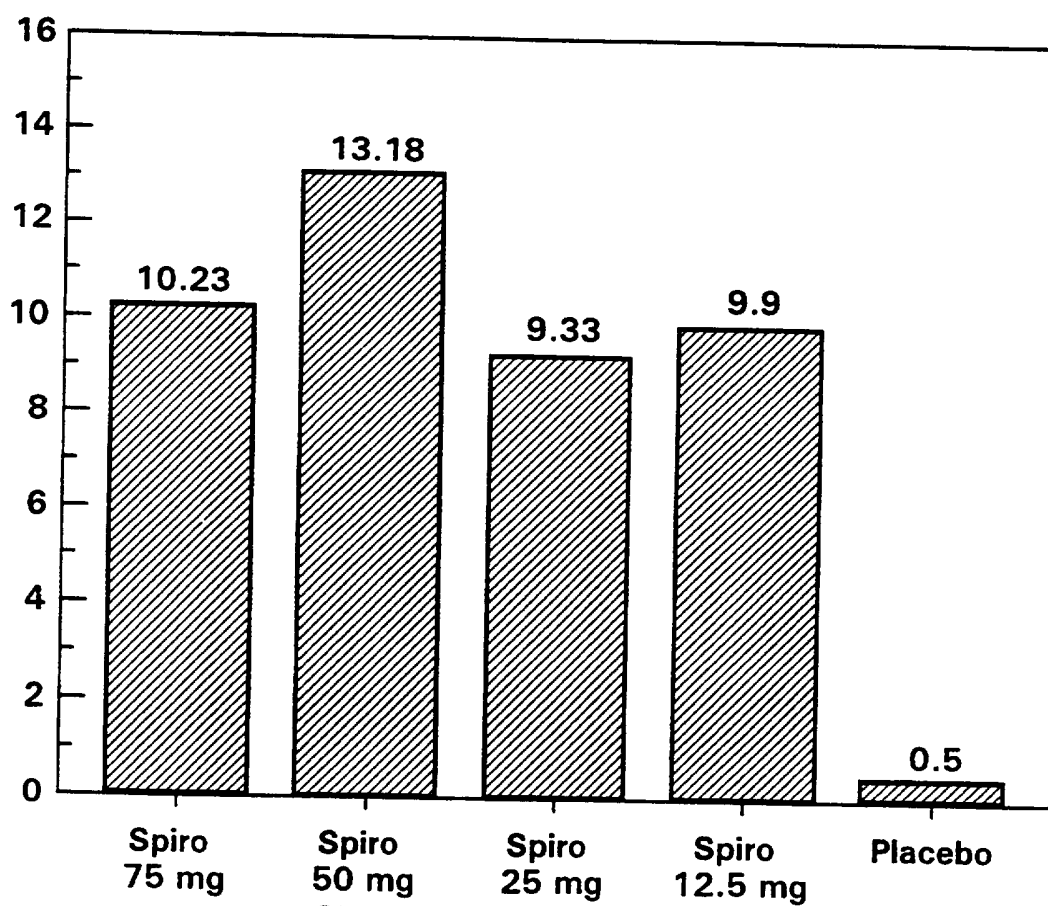


FIG.3

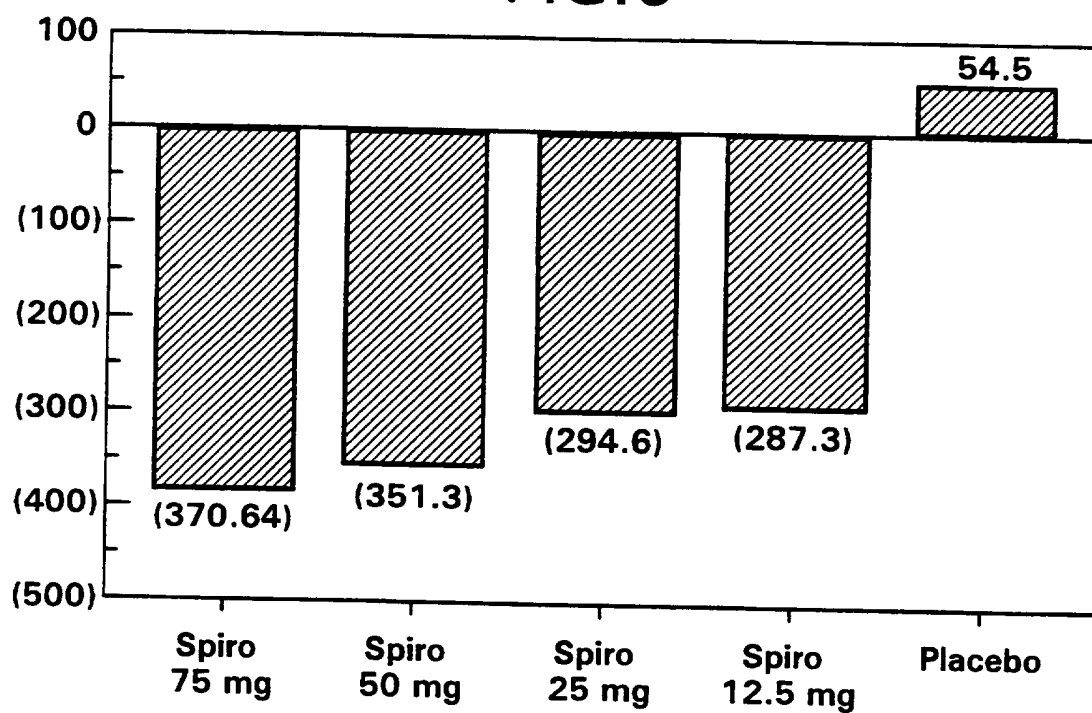
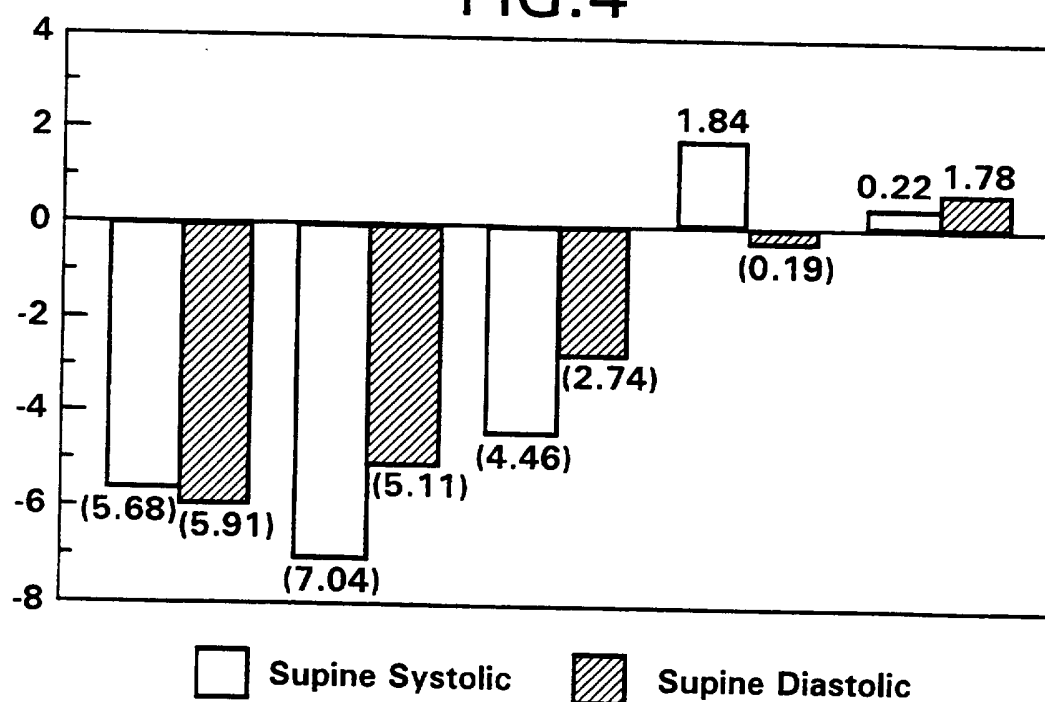


FIG.4



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FIG. 5

