



AU9228454

(12) PATENT ABRIDGMENT (11) Document No. AU-B-28454/92
(19) AUSTRALIAN PATENT OFFICE (10) Acceptance No. 661067

- (54) Title
DIPYRIDAMOLE FOR THE TREATMENT OF PROLIFERATIVE DISEASES
- International Patent Classification(s)
(51)⁵ **A61K 031/505**
- (21) Application No. : **28454/92** (22) Application Date : **18.11.92**
- (30) Priority Data
- (31) Number (32) Date (33) Country
795434 21.11.91 US UNITED STATES OF AMERICA
- (43) Publication Date : **27.05.93**
- (44) Publication Date of Accepted Application : **13.07.95**
- (71) Applicant(s)
ELI LILLY AND COMPANY
- (72) Inventor(s)
RAYMOND FRANCIS KAUFFMAN; JAI PAL SINGH
- (74) Attorney or Agent
SPRUSON & FERGUSON , GPO Box 3898, SYDNEY NSW 2001

- (57) This invention relates to the local delivery of dipyridamole for the treatment of proliferative diseases.

Proliferative diseases such as vascular restenosis, scleroderma, psoriasis, and rheumatoid arthritis share the fundamental mechanism of excessive proliferation of cells in a specific tissue or organ. In each of these diseases, the excessive proliferation of cells contributes significantly to the pathogenesis of the disease.

Claim

1. A method of inhibiting cell proliferation in mammals which comprises the local delivery of an inhibitory amount of dipyridamole.

661067

S & F Ref: 223706

AUSTRALIA

PATENTS ACT 1990

COMPLETE SPECIFICATION

FOR A STANDARD PATENT

ORIGINAL

Name and Address
of Applicant:

Eli Lilly and Company
Lilly Corporate Center
City of Indianapolis State of Indiana
UNITED STATES OF AMERICA

Actual Inventor(s):

Raymond Francis Kauffman and Jai Pal Singh

Address for Service:

Spruson & Ferguson, Patent Attorneys
Level 33 St Martins Tower, 31 Market Street
Sydney, New South Wales, 2000, Australia

Invention Title:

Dipyridamole for the Treatment of Proliferative Diseases

The following statement is a full description of this invention, including the best method of performing it known to me/us:-

Dipyridamole for the
Treatment of Proliferative Diseases

This invention relates to the local delivery of
5 dipyridamole for the treatment of proliferative diseases.

Proliferative diseases such as vascular
restenosis, scleroderma, psoriasis, and rheumatoid
arthritis share the fundamental mechanism of excessive
proliferation of cells in a specific tissue or organ. In
10 each of these diseases, the excessive proliferation of
cells contributes significantly to the pathogenesis of the
disease.

For example, vascular restenosis is
characterized by the reocclusion of coronary arteries
15 following percutaneous transluminal coronary angioplasty
(PTCA), atherectomy, laser angioplasty and arterial bypass
graft surgery. The reocclusion of coronary arteries is
caused in part by the excessive proliferation of vascular
smooth muscle cells. See "Intimal Proliferation of Smooth
20 Muscle Cells as an Explanation for Recurrent Coronary
Artery Stenosis after Percutaneous Transluminal Coronary
Angioplasty," Austin et al., Journal of the American
College of Cardiology 6: 369-375 (Aug. 1985).

Vascular restenosis remains a major long term
25 complication following surgical intervention of blocked
arteries by percutaneous transluminal coronary angioplasty
(PTCA), atherectomy, laser angioplasty and arterial bypass
graft surgery. In about 35% of the patients who undergo
PTCA, reocclusion occurs within three to six months after
30 the procedure. The current strategies for treating
vascular restenosis include mechanical intervention by
devices such as stents or pharmacologic therapies including
heparin, low molecular weight heparin, coumarin, aspirin,
fish oil, calcium antagonist, steroids, and prostacyclin.

These strategies have failed to curb the reocclusion rate and have been ineffective for the treatment and prevention of vascular restenosis. See "Prevention of Restenosis after Percutaneous Transluminal Coronary Angioplasty: The Search for a 'Magic Bullet'," Hermans et al., American Heart Journal 122: 171-187 (July 1991).

The excessive proliferation of fibroblast and mesenchymal cells is associated with rheumatoid arthritis and psoriasis. The inflammatory process that is characteristic of rheumatoid arthritis results in the release of growth factors that induce active proliferation of mesenchymal cells. This proliferation is associated with the production of excessive amounts of enzymes capable of destroying the connective tissues that comprise the joint. Pharmacologic agents that inhibit the proliferative response would be effective in repressing some of the destructive potential of rheumatoid arthritis. See "Recent Insights into the Pathogenesis of the Proliferative Lesion of Rheumatoid Arthritis," Harris, Arthritis and Rheumatism 19: 68-72 (January-February 1976).

Scleroderma (systemic sclerosis) is a multisystem disease affecting primarily the vascular, cutaneous, musculoskeletal, gastrointestinal, pulmonary, cardiac, and renal systems. The apparent diffuse clinical features of systemic sclerosis are thought to be linked by a distinctive vascular lesion in the various target organs. This vascular lesion has inflammatory, proliferative, and indurative phases and is clearly related to the proliferation of the fibroblast and cells capable of fibroblast activity. Controlling this mechanism of fibroblastic activation and proliferation may be useful in

treating or preventing systemic sclerosis. See
"Pathogenesis of Systemic Sclerosis: A Vascular
Hypothesis," Campbell et al., Seminars in Arthritis and
Rheumatism 4: 351-368 (May, 1975).

5 In the pathogenesis of proliferative diseases,
excessive cell proliferation occurs as a result of the
presence of various growth factors and cytokines such as
platelet-derived growth factor (PDGF), epidermal growth
factor (EGF), fibroblast growth factor (FGF) and
10 interleukin-1 (IL-1). For example, growth factors produced
by cellular constituents in the blood and the damaged
arterial vessel wall mediate the proliferation of smooth
muscle cells in vascular restenosis. A novel method of
administering dipyridamole to inhibit cellular
15 proliferation caused by various growth factors is therefore
useful for the treatment of proliferative diseases such as
psoriasis, rheumatoid arthritis, scleroderma, and vascular
restenosis. The American Journal of Medicine 70: 1231-1236
(June 1981).

20 Dipyridamole is commonly prescribed as an
antiplatelet or phosphodiesterase inhibitor. It has been
studied independently or in conjunction with aspirin and/or
prostacyclin for the treatment of vascular restenosis. The
results of these studies have demonstrated that
25 dipyridamole, when systemically administered, is
ineffective in treating or preventing vascular restenosis
in patients. Hermans et al., American Heart Journal 122:
171-187 (July, 1991); Harker et al., Arteriosclerosis 10:
828a (September-October, 1990); and FitzGerald, The New
30 England Journal of Medicine 316:1247-57 (May, 1987). The
use of dipyridamole for the treatment of restenosis has
been ineffective in these studies due to the systemic
method of administration. Serum reduces the effectiveness

of systemically administered dipyridamole as an inhibitor of cell proliferation.

Only upon the observation of the effects of serum on systemically administered dipyridamole did it finally become possible to discover the use of dipyridamole as an antiproliferative agent. The invention discloses the local delivery of dipyridamole as a method of inhibiting cell proliferation and is useful for the treatment of proliferative diseases such as restenosis, scleroderma, psoriasis, and rheumatoid arthritis.

This invention provides a method of inhibiting cell proliferation in mammals which comprises the local delivery of an inhibitory amount of dipyridamole.

Dipyridamole is a well known compound used extensively as a coronary vasodilator. The Merck Index Tenth Edition: 3366 (1983). Its chemical name is 2,6-bis(diethanol-amino)-4,8-dipiperidinopyrimido[5,4-d]pyrimidine. Its preparation is disclosed in British patent 807,826 (1959 to Thomae).

The local delivery of inhibitory amount of dipyridamole for the treatment of cell proliferation can be by a variety of techniques which administer the dipyridamole at or near the proliferative site. Examples of local delivery techniques are not intended to be limiting but to be illustrative of the techniques available. Examples include local delivery catheters, site specific carriers, implants, direct injection, or direct applications.

Local delivery by a catheter allows the administration of a pharmaceutical agent directly to the proliferative lesion. Examples of local delivery using a balloon catheter are described in EPO 383 492 A2 and U.S. Patent 4,636,195 (Wolinsky, January 13, 1987).

Local delivery by an implant describes the surgical placement of a matrix that contains the pharmaceutical agent into the proliferative lesion. The implanted matrix releases the pharmaceutical agent by diffusion, chemical reaction, or solvent activators. Langer, Science 249: 1527-1533 (September, 1990). An example of local delivery by an implant is the use of a stent. Stents are designed to mechanically prevent the collapse and reocclusion of the coronary arteries. Incorporating a pharmaceutical agent into the stent delivers the drug directly to the proliferative site. Local delivery by this technique is described in Kohn, Pharmaceutical Technology (October, 1990). A second example is a delivery system in which a polymer that contains the pharmaceutical agent is injected into the lesion in liquid form. The polymer then cures to form the implant in situ. This technique is described in PCT WO 90/03768 (Donn, April 19, 1990). Another example is the delivery of a pharmaceutical agent by polymeric endoluminal sealing. This technique uses a catheter to apply a polymeric implant to the interior surface of the lumen. The pharmaceutical agent incorporated into the biodegradable polymer implant is thereby released at the surgical site. It is described in PCT WO 90/01969 (Schindler, August 23, 1989). A final example of local delivery by an implant is by direct injection of vesicles or microparticulates into the proliferative site. These microparticulates may be composed of substances such as proteins, lipids, carbohydrates or synthetic polymers. These microparticulates have the pharmaceutical agent incorporated throughout the microparticle or over the microparticle as a coating. Delivery systems incorporating

microparticulates are described in Lange, Science 249: 1527-1533 (September, 1990) and Mathiowitz, et al., J. App. Poly. Sci., 26:809 (1981).

Local delivery by site specific carriers
5 describes attaching the pharmaceutical agent to a carrier which will direct or link the drug to the proliferative cells. Examples of this delivery technique includes the use of carriers such as a protein ligand or a monoclonal antibody or a membrane anchored linker. Lange, Science 249: 1527-1533 (September, 1990); Langworth, Genetic Engineering News (September, 1990).

Local delivery by direct application includes
the use of topical applications. An example of a local delivery by direct application is applying the
15 pharmaceutical agent directly to the arterial bypass graft during the surgical procedure.

Local delivery by direct injection describes
injecting fine particles of dipyridamole suspended in an inert carrier such as sterile saline solution directly into
the proliferative site.

The dosage of dipyridamole required to produce
the therapeutic effect is dependent upon the method of administration and the particular circumstances of the patient. A therapeutic dosage of dipyridamole is an amount
25 sufficient to inhibit the proliferation of cells. The preferred dosage range is defined to be about 1 µg/day to about 100,000 µg/day delivered at or near the proliferative site.

The term "treatment" includes the administration
30 of a compound of present invention to prevent the onset of the symptoms, alleviating the symptoms, or eliminating the disease, condition, or disorder.

5 Microparticles for use in conjunction with local delivery catheter can be prepared using the following ingredients: biological surfactant (0.1-3 % by volume), dextrose (2 - 5 % by volume), dipyridamole (milled to 2- 10 micron particles, 0.1-10 mg/ml), and water.

The above ingredients are mixed and injected into the proliferative lesion. The following example of a formulation for the local delivery of dipyridamole is illustrative only and are not intended to limit the scope of the invention in any way.

Microparticles were prepared using the ingredients below:

	Quantity
polyoxyethylenesorbitan	2 % (by volume)
dextrose	5 % (by volume)
dipyridamole (milled)	250 mg
water	50 ml

15 The microcapsules are injected into the proliferative lesion using a catheter.

20 A critical aspect of the present invention provides for the local delivery of dipyridamole to prevent cell proliferation. Cellular proliferation may be induced by cytokines such as interleukin-1 (IL-1) or multiple growth factors such as platelet-derived growth factor (PDGF), epidermal growth factor (EGF), and/or fibroblast growth factor (FGF).

25 Previous attempts to use dipyridamole for the treatment of restenosis have been ineffective due to the systemic method of administration. Systemic administration includes delivery techniques that introduce the

pharmaceutical agent to the entire organism. Examples of systemic delivery include oral and intravenous administration.

5 Serum reduces the effectiveness of systemically administered dipyridamole as an inhibitor of cell proliferation. The effect of serum on the antiproliferative activity of dipyridamole was demonstrated as follows: Smooth muscle cells from rabbit aorta (derived by explant method as described in Ross, Journal of Cell Biology 50:172 (1971)) were seeded in 96 well tissue culture plates in Dulbecco's Modified Eagle's Medium (DMEM) containing 10% fetal bovine serum (Gibco Laboratories, Grand Island, NY), 2mM L-glutamine, 100 U/ml penicillin, and 100 µg/ml streptomycin. Growth arrested, confluent cultures in 96 well microtiter plates were incubated in medium containing 1%, 5% or 20 % serum (Hyclone Laboratories, Logan, Utah), 2 mM L-glutamine, 100 U/ml penicillin, 100 µg/ml streptomycin, 40 ng/ml PDGF (Genzyme, Cambridge, MA), 1 µCi/ml ³H thymidine (DuPont, Boston, MA) and indicated concentrations of dipyridamole (Sigma Chemical, St. Louis, MO). Cells were incubated at 37°C for 24 hours under 5% CO₂/95% air. At the end of 24 hours, cells were fixed in methanol and DNA synthesis was determined by ³H thymidine incorporation as described in Bonin et al., Exp. Cell Res. 181, 475-482 (1989). The following table demonstrates that increasing concentrations of serum attenuate the growth inhibitory effects of dipyridamole.

'Table 1

Effect of serum concentration on inhibition of DNA synthesis by dipyridamole.

Dipyridamole (μ g/ml)	% Inhibition of DNA Synthesis		
	1% Serum	5% Serum	20% Serum
0	0	0	0
0.04	59	52	6
0.08	70	53	0
0.15	79	60	33
0.3	80	71	38
0.6	86	80	60
1.2	90	87	74
2.5	93	92	77

5

The effect of systemically administered dipyridamole upon cell proliferation was demonstrated as follows:

10 Balloon injury to the left common carotid arteries of male Sprague-Dawley rats (350-400g) was accomplished by three passes of an inflated 2F Fogarty balloon catheter (Baxter Healthcare, McGaw Park, IL) as described by Clowes et al., Lab Invest. 49: 208-215 (1983). Animals were anesthetized with Ketamine (80 mg/kg,

intramuscular, Aveco, Ft. Dodge, IA) and Rompun (16 mg/kg, intramuscular, Mobay Corp., Shawnee, KA). Entry of the balloon catheter to the left common carotid artery was made via a nick in the external carotid artery, which was tied off at the end of the surgical procedure. Dipyridamole was systemically administered for two weeks (.03 and .10% wt/wt, as an admixture in the diet, equivalent to approximately 30 and 100 mg/kg/day, respectively). No significant effect upon intimal thickening in the balloon-injured rat carotid arteries was observed as demonstrated in Table 2.

Table 2

Effect of Systemic Administration of Dipyridamole Upon Intimal Thickening

<u>Systemic Administration</u> <u>of Dipyridamole</u> (% in diet, wt/wt)	<u>Area of Intimal</u> <u>Thickening</u> mm ² , (% of control)
0.00	0.120 ± 0.014 (100)
0.03	0.116 ± 0.017 (97.6)
0.10	0.109 ± 0.014 (90.8)

The effect of the present invention to control the cellular proliferation and intimal thickening by the local delivery of dipyridamole has been demonstrated by in vivo studies. The following examples illustrate the present invention and are not intended to limit the same.

Example 1Balloon Injury of Carotid Arteries

Balloon injury to the left common carotid arteries of male Sprague-Dawley rats (350-400g) was accomplished by three passes of an inflated 2F Fogarty balloon catheter as described by Clowes et al., Lab Invest. 49: 208-215 (1983). Animals were anesthetized with Ketamine (80 mg/kg, intramuscular) and Rompun (16 mg/kg, intramuscular). Entry of the balloon catheter to the left common carotid artery was made via a nick in the external carotid artery, which was tied off at the end of the surgical procedure. Continuous local delivery of dipyridamole was accomplished by means of a minosmotic pump implanted subcutaneously in the back of the rat. Pumps were primed before surgery and implanted immediately following balloon injury. Dosing solutions were delivered to the adventitia (exterior) space surrounding the injured carotid artery via a micro-renathane catheter (MRE-40, Baxter Healthcare, Santa Ana, CA) at a rate of 5 μ l per hour. The catheter is sutured in place with two ligatures (4-0 silk) to the left external carotid artery.

Fourteen days post surgery, animals were anesthetized (vide supra) and perfused through the abdominal aorta in a retrograde manner at physiological pressure with a zinc formalin fixative (Anatech LTD., Battle Creek, MI). Middle sections (5 mm) of the carotids were removed from the animals, processed, and embedded in paraffin. Three adjacent cross sections (5 μ m thick) of each vessel were cut, stained with hematoxylin and eosin, and cross-sectional intimal areas quantitated with an image analyzer (Quantimet 970, Cambridge Inst., Cambridge, UK).

The difference between intimal areas of drug-treated vs. control groups were analyzed for statistical significance using Student's t-test as described in

Tallarida et al., Manual of Pharmacologic Calculations with Computer Programs, Springer-Verlag, New York, 1981, p. 51.

P values less than 0.05 were taken to indicate statistical significance. The results are demonstrated in Table 3.

5

Table 3

Effect of Local Administration of Dipyridamole Upon Intimal Thickening

<u>Local, Adventitial</u> <u>Administration of</u> <u>Dipyridamole</u> ($\mu\text{g/day}$)	<u>Area of Intimal</u> <u>Thickening</u> mm^2 , (% of control)
0 (Vehicle)	0.129 ± 0.013 (100)
600	0.087 ± 0.011 (67.4)*

* $P < 0.05$ vs. corresponding control group (i.e., absence of dipyridamole)

Example 2

Dipyridamole also inhibits proliferation of cells of mesenchyme origin. Inhibition of fibroblast growth by dipyridamole is demonstrated as follows: 20,000 Balb/c3T3 fibroblasts (American Tissue Culture Type, CCL-163) were plated in 12 well tissue culture plates in 3 ml DMEM containing 5% fetal bovine serum, 2 mM L-glutamine, 100 U/ml penicillin, and 100 $\mu\text{g/ml}$ streptomycin, and were incubated for 18-24 hours. Cells were then transferred to above medium containing indicated concentrations of dipyridamole. After three days cell growth was determined

20

by counting using a ZM Coulter counter (Coulter Diagnostic, Inc.).

Table 4

Inhibition of Fibroblast Growth by Dipyridamole

Dipyridamole ($\mu\text{g/ml}$)	% Inhibition of Cell Growth
0.0	0.0
1.0	0
5.0	55
10	72
20	86
40	95

Example 3

Dipyridamole inhibits smooth muscle cell proliferation induced by multiple growth factors. Smooth muscle cells from rabbit aorta (derived by explant method) were seeded in 96 well tissue culture plates in DMEM containing 10% fetal bovine serum, 2mM L-glutamine, 100 U/ml penicillin and 100 $\mu\text{g/ml}$ streptomycin. Growth arrested, confluent cultures in 96 well microtiter plates were incubated in medium containing 1% platelet poor plasma, 2 mM L-glutamine, 100 U/ml penicillin, 100 $\mu\text{g/ml}$ streptomycin, 20 ng/ml PDGF, 3ng/ml EGF (Genzyme), 3 ng/ml FGF (Genzyme), 1 $\mu\text{Ci/ml}$ ^3H thymidine and indicated

concentrations of dipyridamole. Cells were incubated at 37°C for 24 hours under 5% CO₂/95% air. At the end of 24 hours, cells were fixed in methanol. DNA synthesis was determined by ³H thymidine incorporation as previously described. The results in Table show that dipyridamole inhibits cell proliferation induced by PDGF, FGF, and EGF.

Table 5

Inhibition of DNA Synthesis induced by PDGF, FGF or EGF by dipyridamole.

Dipyridamole (μ g/ml)	% Inhibition of DNA Synthesis		
	FGF	EGF	PDGF
0	0	0	0
0.08	67	68	64
0.15	74	76	71
0.32	80	85	82
0.64	87	90	89
1.2	92	94	93
2.5	93	96	96

~~We claim:~~ The claims defining the invention are as follows:

1. A method of inhibiting cell proliferation in mammals which comprises the local delivery of an inhibitory amount of dipyridamole.

5 2. The method of use of Claim 1 wherein the cells inhibited are vascular smooth muscle cells.

3. The method of use of Claim 1 where the local delivery is by means of a local delivery catheter.

10 4. The method of use of Claim 1 where the local delivery is by means of a site specific carrier.

5. The method of use of Claim 1 where the local delivery is by means of an implant.

6. The method of use of Claim 1 where the local delivery is by means of a membrane anchored linker.

15 7. The method of use of Claim 1 where the local delivery is by means of direct injection.

8. The method of use of Claim 2 where the local delivery is by means of a local delivery catheter.

20 9. The method of use of Claim 2 where the local delivery is by means of a membrane anchored linker.

10. The method of use of Claim 2 where the local delivery is by means of an implant.

11. The method of use of Claim 2 where the local delivery is by means of direct injection.

25 12. A method of inhibiting cell proliferation in mammals which comprises the local delivery of an inhibitory amount of dipyridamole wherein said cellular proliferation is induced by growth factors comprising platelet-derived growth factor (PDGF), epidermal growth factor (EGF), and/or
30 fibroblast growth factor (FGF).

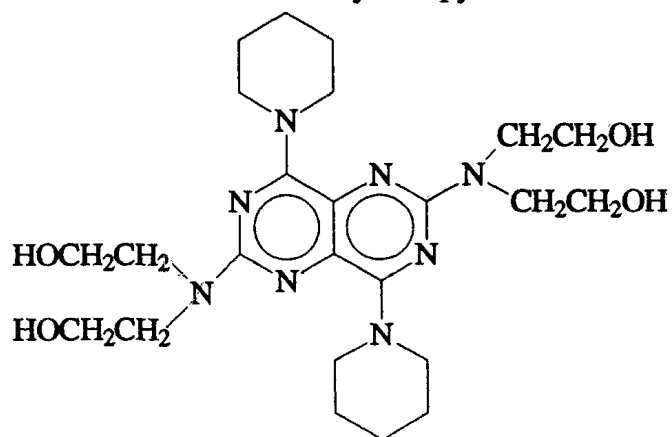
DATED this ELEVENTH day of NOVEMBER 1992
Eli Lilly and Company

Patent Attorneys for the Applicant
SPRUSON & FERGUSON

Dipyridamole for the Treatment of Proliferative Diseases

Abstract

This invention relates to the local delivery of dipyridamole



5 for the treatment of proliferative diseases.