SECURITIES AND EXCHANGE COMMISSION Washington, D.C. 20549 FORM 10-K

Annual Report pursuant to section 13 or 15(d) of the Securities Exchange Act of 1934 For the fiscal year ended March 31, 1995 MYLAN LABORATORIES INC. Commission File No. 1-9114

(Exact name of registrant as specified in its charter)

Pennsylvania 25-1211621

(State or other jurisdiction of incorporation or organization) No.)

(IRS Employer Identification

130 Seventh Street 1030 Century Building Pittsburgh, Pennsylvania

15222 (Address of principal executive offices) (Zip Code)

Registrant's telephone number, including area code: 412-232-0100 Securities registered pursuant to Section 12(b) of the Act:

Name of Each Exchange

Title of Each Class Common Stock, par value \$.50 per share

on Which Registered New York Stock Exchange

Securities registered pursuant to Section 12(g) of the Act: None
Indicate by checkmark whether the registrant (1) has filed all reports required to be filed by Section 13 or 15(d) of the

Securities Exchange Act of 1934 during the preceding 12 months (or for such shorter period that the registrant was required

to file such reports), and (2) has been subject to such filing requirements for the past 90 days.

Yes X No

Indicate by checkmark if disclosure of delinquent filers pursuant to Item 405 of Regulation S-K is not contained herein.

and will not be contained, to the best of registrant's knowledge, in definitive proxy or information statements incorporated

by reference in Part III of this Form 10-K or any amendment to this Form 10-K.[]

The aggregate market value of voting stock held by persons other than Directors and Officers of the registrant

computed by reference to the closing price of such stock as of May 31, 1995: \$2,200,910,441

The number of shares of Common Stock of the registrant outstanding as of May 31, 1995:

79,514,544

Documents incorporated by reference into this Report are: Parts I and II,

Proxy Statement for 1995 Annual Meeting of Shareholders. Part III. Items 10-13

PART I

Mylan Laboratories Inc., a Pennsylvania corporation incorporated in 1970, and its subsidiaries (herein referred to collectively as the "Company") are engaged in the development, manufacturing and distribution of pharmaceutical products for resale by others. References herein to fiscal 1995, 1994 and 1993 mean the fiscal years ended March 31, 1995, 1994 and 1993, respectively.

Through its subsidiary, Mylan Pharmaceuticals Inc., the Company is recognized as one of the leaders in the generic pharmaceutical industry. Pharmaceutical products initially sold on an exclusive basis are known in the industry as proprietary or branded products. Generic drugs are therapeutically equivalent to their brand name counterparts and are generally sold at prices significantly less than branded products. Accordingly, generics provide a safe, effective and cost efficient alternative to users of these products.

The Company manufactures substantially all of its oral dose products in either its Mylan Pharmaceuticals subsidiary's Morgantown, West Virginia facility or its subsidiary Mylan Inc.'s facility in Caguas, Puerto Rico. To facilitate timely delivery of products to customers in all fifty states the Company operates distribution centers in Greensboro, North Carolina and Reno, Nevada.

Due to the non-exclusive nature of generic products, the generic industry is comprised of numerous competitors including manufacturers who market their products under their own name, distributors who market products manufactured by others and brand name companies, who in recent years market their products under both the brand name and as the generic substitute. This diversity provides significant price competition within the generic pharmaceutical industry which generally results in decreasing prices of generic products over time to those who supply such products to the retail market.

Over the past year, the Company has entered into strategic alliances with several branded pharmaceutical companies. These alliances through distribution and licensing agreements provide the Company with additional products to further broaden the Company's product line. The Company continues to examine other alliances as a way to grow and react in the rapidly changing health care arena.

In June 1989 the Company acquired a 50% interest in Somerset Pharmaceuticals, Inc. ("Somerset"). Pursuant to a license agreement with a Hungarian pharmaceutical company, Somerset has exclusive marketing rights to the product Eldepryl(R) in the United States and certain other countries. Commercial shipments of the product by Somerset commenced in late August 1989.

Under the Orphan Drug Act, Somerset has exclusivity relating to marketing the chemical compound Eldepryl(R) for use as a treatment for late stage Parkinson's disease through June 1996.

Under the Waxman Hatch Act, Somerset had exclusivity for all uses of the chemical compound through June 1994. Somerset is actively involved in research projects regarding additional uses of this and other chemical compounds.

In October 1991, a wholly-owned subsidiary of the Company merged with Dow Hickam Pharmaceuticals, Inc. ("Hickam"), an established branded pharmaceutical company located in Sugar Land, Texas. Hickam currently manufactures and/or markets specialty pharmaceutical products and devices used principally as wound care treatments through its nation-wide sales force.

On February 25, 1993 the Company acquired substantially all of the net assets of Bertek, Inc. ("Bertek"). Bertek, headquartered in St. Albans, Vermont is a manufacturer of transdermal drug delivery systems and also has operations in laminating, coatings and label manufacturing. In addition Bertek provides components, using internally developed technology for transdermal patches marketed by other companies. Bertek is actively involved in development projects to provide new transdermal products.

Products

The information on the Company's product line set forth on pages 17 and 25 of the Annual Report to Shareholders for the year ended March 31, 1995 is incorporated herein by reference. All pharmaceutical products presently manufactured by the Company have been previously developed and marketed by other firms with the exception of Maxzide(R) , Maxzide(R) -25MG and Cystagon TM .

The Company is required to secure and maintain approval from the U.S. Food and Drug Administration ("FDA") for the products and dosage forms which it manufactures. The number of products and dosage forms for which the Company is an approved manufacturer has expanded in recent years. See "New Product Approvals".

During fiscal 1995, 1994 and 1993 approximately \$30,533,000, \$21,648,000 and \$13,524,000 respectively, were expensed by the Company for the development of formulations and procedures for products which it desires to produce, use or sell. The Company's research and development efforts are conducted primarily to qualify the Company to manufacture ethical pharmaceuticals under FDA standards and approval.

New Product Approvals

During fiscal 1995, five approvals were received from the FDA. Subsequent to fiscal 1995 one approval was received from the FDA. The Company presently has requests for approval pending before the FDA representing 19 products of varying strengths. In addition the Company has two Investigational New Drug applications filed with the FDA for new innovator compounds.

Customers and Markets

The Company sells its products to proprietary and ethical pharmaceutical wholesalers and distributors, drug store chains, drug manufacturers and public and governmental agencies. Sales to one customer, McKesson Health & Drug Company, represented 12% of net sales in 1993. No single customer represented more than 10% of net sales in 1995 or 1994.

A majority of the Company's products are marketed to food and drug store chains and to pharmaceutical distributors and wholesalers, who in turn market to retailers, managed care entities, hospitals and government agencies. Certain other products are marketed to institutional accounts who in turn obtain the products from pharmaceutical distributors and wholesalers. The Company's sales activities involve limited public promotion of its products. Approximately 126 employees of the Company are engaged full-time in selling products and servicing customers.

Competition

The Company sells to various markets and classes of customers. With respect to each of the various products it sells, the Company believes it is subject to active competition from numerous firms. The four primary means of competition are services, quality of products, approval for manufacture by the FDA and price. The competition experienced by the Company varies among the markets and classes of customers. The Company has experienced additional competition from brand-name competitors who have entered the generic pharmaceutical industry through the creation of generic subsidiaries, purchasing generic companies or by licensing their products prior to or as their product's patents expire.

Product Liability

Product liability suits by consumers represent a continuing risk to firms in the pharmaceutical industry. The Company strives to minimize such risks by stringent quality control procedures. Although the Company carries insurance, it believes that no reasonable amount of insurance can fully protect it against all such risks because of the potential liability inherent in the business of producing pharmaceuticals for human consumption.

Raw Materials

The chemical ingredients and other materials and supplies used in the Company's pharmaceutical manufacturing operations are generally available and purchased from many different foreign and domestic suppliers. However, some products may have only one source approved by the FDA for certain pharmaceutical ingredients used in their manufacturing process. If this material was no longer available, qualifying a new supplier could delay the manufacturing of such products. During fiscal 1995 there was a limited supply of raw materials to all generic manufacturers of cimetidine. This product had a significant contribution to the Company's net sales and gross profit during fiscal 1995.

With regards to foreign suppliers, recent and pending regulatory action may make obtaining raw materials prior to patent expiration increasingly difficult. This could delay the Company's ability to develop, manufacture and obtain FDA approval to market certain new products.

Regulation

The Company's operations are subject to regulation under the Federal Food, Drug and Cosmetic Act, pursuant to which government standards as to "good manufacturing practice", product content, purity, labeling, effectiveness and recordkeeping (among other things) must be observed. In this regard, the FDA has extensive regulatory powers over the activities of pharmaceutical manufacturers.

The Company is also subject to inspection and regulation under other federal and state legislation relating to drugs, narcotics and alcohol. Many of its suppliers and customers, as well as the drug industry in general, are subject to the same or similar governmental regulations.

In October 1990 Congress passed the Medicaid Prudent Pharmaceutical Purchasing Act of 1990. Under the Act, the Company is required to pay a rebate to each state predicated on the number of prescriptions for the Company's products reimbursed by the states under Medicaid. The Act was effective January 1, 1991.

The President signed into law the Uruguay Round Agreements Act ("URAA") in December 1994. URAA which took effect on June 8, 1995 implemented the General Agreement of Tariffs and Trade ("GATT"). One change in US law required by GATT is the amendment of patent law to reflect a patent term of 20 years from the date of filing the application instead of the current term of 17 years from the date of issuance. URAA extended the requirement by allowing the application of this provision to all patents in force on June 8, 1995.

Congress recognized the potential harm in this requirement and provided that a potential competitor who has already made a "substantial investment" in a competing product could make, use and sell its product after the expiration of the original patent period provided that they pay the patentee "equitable remuneration" through the extended patent period. However, the FDA has taken the position that it cannot approve an Abbreviated New Drug Application ("ANDA"), which certifies the date of patent expiration, until the expiration of the extended patent period. This could delay the launch of future products but has no effect on products presently marketed and sold by the Company.

The Company and other generic drug manufacturers have approached Congress in an attempt to secure passage of legislation which would allow the FDA to approve applications on the passage of the original expiration date. In addition, Mylan has filed suit, as a co-plaintiff, against Bristol Myers Squibb seeking declaratory judgement with respect to these issues in the marketing of captopril.

It is impossible for the Company to predict the extent to which its operations will be affected under the regulations discussed above or any new regulations which may be adopted by regulatory agencies.

Employees

The Company employs approximately 1310 persons, approximately 570 of whom serve in clerical, sales and management capacities. The remainder are engaged in production and maintenance activities.

The production and maintenance employees at the Company's manufacturing facilities in Morgantown, West Virginia, are represented by the Oil, Chemical and Atomic Workers International Union (AFL-CIO) and its Local Union 8-957 under a contract which expires April 5, 1998.

Backlog

At March 31, 1995, the uncompleted portions of the Company's backlog of orders was approximately \$20,979,000 as compared to approximately \$12,543,000 at March 31, 1994. Because of the relatively short lead time required in filling orders for its products, the Company does not believe these interim backlog amounts bear a significant relation to sales or income for any full twelve-month period.

ITEM 2. Properties

The Company operates from various facilities in the United States and Puerto Rico having an aggregate of approximately 1,057,000 square feet.

Mylan Pharmaceuticals owns production, warehouse, laboratory and office facilities in four buildings in Morgantown, West Virginia containing approximately 500,000 square feet. Mylan Pharmaceuticals operates two distribution centers, one in Greensboro, North Carolina containing approximately 64,000 square feet which it owns and one in Reno, Nevada containing approximately 25,000 square feet under a lease expiring in 1996. Currently under construction in Morgantown, West Virginia is a 150,000 square foot research and development facility.

Mylan Inc. owns a production and office facility in Caguas, Puerto Rico containing approximately 115,000 square feet and a production facility in Cidra, Puerto Rico containing approximately 32,000 square feet.

Dow Hickam Pharmaceuticals, Inc. owns production, warehouse and office facilities in two buildings in Sugar Land, Texas containing approximately 70,000 square feet. Hickam also operates a filling and packaging facility in Sugar Land, Texas containing approximately 15,000 square feet under a lease expiring in 1996.

Bertek owns production, warehouse, laboratory and office facilities in five buildings in Swanton and St. Albans, Vermont containing approximately 159,000 square feet. Bertek also operates a coating and extrusion facility in St. Albans containing approximately 71,000 square feet under a lease expiring in 2015.

The Company's production equipment includes that equipment necessary to produce and package tablet, capsule, aerosol, transdermal and powder dosage forms. The Company maintains six analytical testing laboratories for quality control.

The Company's facilities are operated primarily on a two shift basis. Properties and equipment are well maintained and adequate for present operations.

The Company's corporate offices, containing approximately 5,800 square feet, are located at 130 Seventh Street, 1030 Century Building, Pittsburgh, Pennsylvania, and are occupied under a lease expiring in 1996.

ITEM 3. Legal Proceedings

On November 20, 1990 the Company filed a complaint against American Cyanamid Company ("Cyanamid"). Cyanamid, through its Lederle Laboratories Division is the exclusive distributor of the Mylan products Maxzide(R) and Maxzide(R) -25MG pursuant to a March, 1984

agreement. The complaint alleged that Cyanamid underpaid the Company based on the terms of the agreement and that Cyanamid had not met certain obligations under the agreement with regard to marketing of the products. The Company sought general relief in the form of compensatory and punitive damages and also sought certain trademark rights to Maxzide(R) and Maxzide(R) -25MG currently owned by Cyanamid.

On May 30, 1991 Cyanamid filed an Answer and Counterclaim to the complaint filed by the Company on November 20, 1990. The counterclaim included allegations of fraudulent inducement and breach of contract regarding the March 1984 Maxzide contract and allegations of defamation by the Company's former Chairman individually and as an officer of the Company. Cyanamid sought dismissal of the Company's complaint and compensatory and punitive damages. The suit was tried in Federal Court for the Northern District of West Virginia.

During 1994 the jury in the Company's lawsuit against Cyanamid ruled in favor of Cyanamid on the Company's complaint and in favor of the Company on Cyanamid's counterclaims and the judge dismissed the defamation counterclaim. Both parties appealed and the Court of Appeals for the Fourth Circuit affirmed the jury's action in all respects. The judge's decision to dismiss the defamation counterclaim however has been reversed and the case has been remained to the lower court for the resolution of this one remaining issue. The Company and the Estate of the former Chairman have each filed a rit of certiorari with the U.S. Supreme Court with respect to the reinstatement of the defamation counterclaim. Subsequently, this case has been transferred to the U.S. District Court for the District of New Jersey.

On November 24, 1992, Marion Merrell Dow Inc. ("MMD") and Tanabe Seiyaku Co. LTD ("Tanabe") filed suit in Federal District Court for the Western District of Pennsylvania against the Company, its wholly-owned subsidiary Mylan Pharmaceuticals Inc., and another company claiming infringement of Tanabe's patent for the manufacture of diltiazem. MMD and Tanabe seek permanent injunctive relief and treble damages to compensate for the alleged infringement and costs of suits. Shortly thereafter MMD and Tanabe filed a complaint with the International Trade Commission ("ITC"), setting forth the same allegations. On motion by the plaintiff the District Court action was stayed pending the outcome of the ITC proceedings. In May 1995, the Administrative Law Judge for the ITC ruled in favor of the Company on all counts. The ruling was appealed by MMD and Tanabe and in June 1995 the Commission upheld the ruling in Mylan's favor. This case remains subject to appeal by MMD and Tanabe.

On September 7, 1994, Upsher-Smith Laboratories filed suit in Minnesota State Court against the Company and its wholly-owned subsidiary Mylan Pharmaceuticals Inc. The suit alleges breach of contract, breach of implied contract, detrimental reliance and promissory estoppel with respect to the sale and distribution of cimetidine. The Company believes this lawsuit is without merit and intends to vigorously defend its position.

While it is not feasible to predict the ultimate outcome of such proceedings it is the opinion of management that the outcome of these suits will have no material adverse effect on the Company's operation, financial position, or liquidity.

ITEM 4. Submission of Matters to a Vote of Security Holders

Not applicable.

EXECUTIVE OFFICERS OF THE REGISTRANT

The names, ages and positions of the Company's executive officers are as follows:

Milan Puskar	60	Chairman, Chief Executive Officer and
		President
Dana G. Barnett	54	Executive Vice President
Louis J. DeBone	49	Vice President-Operations
Roger L. Foster	48	Vice President-General Counsel
Roderick P. Jackson	55	Senior Vice President
Joseph J. Krivulka	43	Vice President
Dr. John P. O'Donnell	49	Vice President-Research and
		Quality Control
Patricia Sunseri	55	Vice President-Investor and
		Public Relations
C.B. Todd	61	Senior Vice President
Robert W. Smiley	73	Secretary

Mr. Puskar was employed by the manufacturing subsidiary of the Company from 1961 to 1972 and served in various positions including: Secretary-Treasurer, Executive Vice President and member of the Board of Directors. In addition, he has served as a partner in several pharmaceutical firms in foreign countries. Mr. Puskar has served as President of the Company since 1976 and as Chairman and Chief Executive Officer since November of 1993.

Mr. Barnett has been a Vice President of the Company since 1974. His principal occupation since 1966 has been in various management positions with the manufacturing subsidiary of the Company. His responsibilities have covered production, quality control and product development. He became Senior Vice President in 1978 and Executive Vice President in 1987. Since June of 1991, he also serves as President and Chief Executive Officer of Somerset Pharmaceuticals, Inc., a joint venture subsidiary of the Company.

Mr. DeBone has been employed by the Company since September, 1987. Prior to assuming his present position in November, 1991 as Vice President-Operations he served as Vice President-Quality Control. He was previously employed with the Company from March, 1976 until June, 1986 and served as Director of Manufacturing.

Mr. Foster has been employed by the Company since May, 1984. Prior to assuming his present position in June, 1995 as Vice President-General Counsel he served as Director of Legal Services and as Director of Governmental Affairs.

Mr. Jackson has been employed by the Company since April, 1986. Prior to assuming his present position in October, 1992 as Senior Vice President he served as Vice President-Marketing and Sales.

Mr. Krivulka has been employed by the Company since March, 1990. Prior to assuming his present position in April, 1992 as Vice President he served as Assistant to the President. Since April of 1993, he also serves as President of Bertek, Inc., a subsidiary of the Company. From 1989 to 1990 he was employed by Janssen Pharmaceutica, a division of Johnson & Johnson, as Executive Director of Business Unit Management.

Dr. John O'Donnell has been employed by the Company since 1983. Prior to assuming his present position in November, 1991 as Vice President-Research and Quality Control, he served as Vice President-Research and Product Development and as Director of Chemistry and Product Development.

Mrs. Sunseri has been employed by the Company since 1984. Prior to assuming her present position in October, 1989 as Vice President-Investor & Public Relations, she served as Director of Investor Relations.

Mr. Todd has been employed by the Company since 1970. Prior to assuming his present position in October, 1987 as Senior Vice President, Mr. Todd served as Vice President-Quality Control. He also serves as President of Mylan Pharmaceuticals Inc., a subsidiary of the Company.

Mr. Smiley has been Secretary of the Company for approximately twenty years and on December 12, 1975, he was elected to the Board of Directors. His principal occupation is and for approximately forty-one years has been an attorney-at-law in Pittsburgh, Pennsylvania. He was a partner in the law firm of Smiley, McGinty and Steger, general counsel to the Company. On October 1, 1992 Mr. Smiley joined the law firm of Doepken Keevican Weiss & Medved Professional Corporation.

There is no family relationship between any of the above executive officers. Officers of the Company serve at the pleasure of the Board of Directors.

ITEM 5. Market for Registrant's Common Equity and Related Stockholder Matters

The information required by item 5 is hereby incorporated by reference to pp. 36 and 52 of the accompanying Annual Report to Shareholders for the year ended March 31, 1995.

ITEM 6. Selected Financial Data

The information required by item 6 is hereby incorporated by reference to p. 36 of the accompanying Annual Report to Shareholders for the year ended March 31, 1995.

ITEM 7. Management's Discussion and Analysis of Financial Condition and Results of Operations

The information required by item 7 is hereby incorporated by reference to pp. 37-39 of the accompanying Annual Report to Shareholders for the year ended March 31, 1995.

ITEM 8. Financial Statements and Supplementary Data

The information required by item 8 is hereby incorporated by reference to pp. 40-52 of the accompanying Annual Report to Shareholders for the year ended March 31, 1995.

ITEM 9. Changes in and Disagreements with Accountants on Accounting and Financial Disclosure

Not applicable.

ITEM 10. Directors and Executive Officers of the Registrant

The information as to directors required by item 10 is hereby incorporated by reference to pp. 1-3 of the Company's 1995 Proxy Statement. Information concerning executive officers is provided in Part I of this report under the caption "Executive Officers of the Registrant".

ITEM 11. Executive Compensation

The information required by item 11 is hereby incorporated by reference to pp. 3-13 of the Company's 1995 Proxy Statement.

ITEM 12. Security Ownership of Certain Beneficial Owners and Management

The information required by item 12 is hereby incorporated by reference to p. 13 of the Company's 1995 Proxy Statement.

ITEM 13. Certain Relationships and Related Transactions

Not applicable.

ITEM 14. Exhibits, Financial Statement Schedules, and Reports on Form 8-K

(a) 1.	List	of	Financial	Statements Page Number
INCLUDED IN ANNUAL REPORT TO SHAREHOLDERS:				
Consolidated Balance Sheets				40-41
Consolidated Statements of Earnings				42
Consolidated Statements of Shareholders' Equity				43
Consolidated Statements of Cash Flows				44-45
Notes to Consolidated Financial Statements				46-51
Independent Auditors' Report				52

2. Financial Statement Schedules

The information required by this item is incorporated herein by reference to Exhibit 99. All other schedules have been omitted because they are not required.

3. Exhibits

(3)(a) Amended and Restated Articles of Incorporation of the registrant, filed as Exhibit (3)(a) to Form 10-Q for quarter ended June 30,

1992 and incorporated herein by reference.

(b) By-laws of the registrant, as amended to date, filed as Exhibit 3(b) to Form 10-Q for the quarter ended June 30, 1992 and incorporated herein by reference.

- (10)(a) 1986 Incentive Stock Option Plan, as amended to date, filed as Exhibit 10(b) to Form 10-K for fiscal year ended March 31, 1993 and incorporated herein by reference.
 - (b) "Salary Continuation Plan" with Milan Puskar, Dana G. Barnett and C.B. Todd each dated as of January 27, 1995 and filed herewith.

RETIREMENT BENEFIT AGREEMENT

This Retirement Benefit Agreement (the "Agreement") is entered into on this 27th day of January, 1995 (the "Effective Date") by and between:

Mylan Laboratories Inc., a Pennsylvania Corporation, with offices located at 781 Chestnut Ridge Road, Morgantown, WV 26505 (hereinafter referred to as "Mylan" or "Company").

and

Milan Puskar, an employee of Mylan who resides at Lakeview Resort, Route 8, Box 88-A Morgantown WV 26505, (hereinafter referred to as "Employee or "Puskar").

WHEREAS the Company and Employee, in recognition of Employee's long and valuable contribution to the success of the Company, entered into a Salary Continuation Agreement on April 1, 1990; and

WHEREAS Employee continues to perform valuable services for the Company; and

WHEREAS in recognition of his continuing service to Mylan, the Company wishes to provide Employee with financial assistance with respect to certain Contingencies, in addition to that provided for in said April 1, 1990 Agreement; and

WHEREAS the Company and Employee wish to RESCIND, and to REPLACE said Salary Continuation Agreement with this Agreement;

WITNESSETH THEREFORE that in consideration of the additional benefits provided for hereunder, the premises and covenants set forth herein, and other good and valuable consideration, the sufficiency of which is hereby acknowledged, the Company and Employee, intending to be legally bound, agree as follows:

I. DEFINITIONS

Whenever used in the Agreement the following terms shall be defined as follows:

- (a) "Advisor" or "Advisors" shall mean with respect to Employee any person including lawyers, accountants, estate planners and others, with whom he may wish to review and discuss the matters set forth herein.
- (b) "Agreement" shall mean this Retirement Benefit Agreement which is entered into on the $27 \mathrm{th}$ day of January, 1995.
- (c) "At-Will" shall mean with respect to the period of Puskar's employment with Mylan, that the Company is under no obligation to continue to employ Puskar for any period of time, and can terminate his employment at any time without notice, subject to certain statutory and regulatory requirements; and that Employee is under no obligation to remain employed by the Company, and can terminate his employment with Mylan at any time, without notice.
 - (d) "Contingency" shall mean Retirement or death.
- (e) "Mylan" or "Company" shall mean Mylan Laboratories Inc., its subsidiaries and affiliates.
- (f) "Net Present Value" ("NPV") shall mean the present value at any given time of the entire benefit to be paid, discounted at seven percent (7%) per annum.
- (g) "Party" or "Parties" shall mean the Company or Employee, or both the Company and Employee depending upon which term is required by the context in which it is used.
- (h) "Retire" or "Retirement" shall mean the day and date on which Puskar's employment with the Company is terminated by either Party for any reason other than death of the Employee.
- (i) "Successor" shall mean any person, partnership, limited partnership, joint-venture, corporation, trust or any other entity or organization who, subsequent to the Effective Date, comes into possession of or acquires, either directly or indirectly, all or substantially all of the Company's business, assets or voting stock, or the right to direct the business activities and practices of the Company.

II. RESCISSION OF PRIOR AGREEMENT

The Salary Continuation Agreement entered into by the Parties on April 1, 1990 and any and all other agreements, express or implied, which may have been entered into by them prior to the execution of this Agreement, including by way of example and not of limitation, any agreement which addresses or is related to salary continuation, deferred compensation, employment, or similar matters (excluding agreements related to stock options) are hereby RESCINDED by the mutual consent of the Parties hereto upon execution of this Agreement.

THEREFORE THE PARTIES ACKNOWLEDGE THAT ANY RIGHTS AND OBLIGATIONS SET FORTH IN ANY SUCH AGREEMENT, WHETHER EXPRESS OR IMPLIED, ARE FOREVER WAIVED, NULL AND VOID, AND UNENFORCEABLE AT LAW OR IN EQUITY.

III. RETIREMENT

- 3.1 Upon his Retirement from the Company, and if he is eligible to receive payments as provided for elsewhere herein, Employee shall receive an annual retirement benefit equal to the amount set forth below.
- 3.2 Should Employee Retire after the Effective Date but on or before March 31, 1996 he shall receive two hundred fifty thousand dollars (\$250,000.00) each year for fifteen (15) years.
- 3.3 Should Employee Retire after March 31, 1996 he shall receive three hundred thousand dollars (\$300,000.00) each year for fifteen (15) years.
- 3.4 The Company shall pay each of the fifteen (15) annual payments due hereunder in twelve (12) equal or substantially equal installments. The first of any such payments shall be made on the first day of the month following the month in which Employee Retires, and each subsequent payment shall be made on the first day of each successive month until Mylan's obligations with respect to such payments have been satisfied.
- 3.5 However, upon the written request of the Employee, Mylan may pay to Employee the NPV of any amount, or of the balance of any such amount, to which he is entitled hereunder in a lump-sum payment. Said payment shall be paid within thirty (30) days of the date of Employee's request for such payment.

IV. AWARD OF ADDITIONAL RETIREMENT BENEFITS

- 4.1 The Company may increase the retirement benefit to which Employee is entitled under Article III. Any such increases shall be subject to the following conditions:
- (a) the maximum benefit to which Employee is entitled under Article III shall not exceed five hundred thousand dollars (\$500,000.00) per year for fifteen (15) years;
 - (b) no increases shall be granted earlier than April 1, 1996 or later than April 1, 2000;
 - (c) any increase shall be expensed for accounting purposes at the time it is granted;
- (d) the Company's obligation to pay the increased benefit to Employee, if and when such increases are awarded, shall be subject to all the terms and conditions set forth herein; and
- (e) the inclusion herein of any increase in Employee's retirement benefit shall become effective upon Employee's receipt of a notice from the Company informing him of the additional benefit.

V. CAPACITY TO PERFORM DUTIES

- 5.1 Should Employee become unable to perform the material and substantial duties of his position the Company shall pay to Employee five hundred thousand dollars (\$500,000.00) each year for fifteen (15) years in equal or substantially equal payments. However, upon the written request of the Employee, Mylan may pay to Employee the NPV of any amount, or of the balance of any such amount, to which he is entitled hereunder in a lump-sum payment. Said payment shall be paid within thirty (30) days of the date of Employee's request for such payment.
- 5.2 The certification of a licensed physician as to Employee's inability to perform the material and substantial duties of his position shall be conclusive with respect to his status, and the Company agrees to be unequivocally bound by any such certification.
- 5.3 If the Employee receives the benefit described in 5.1 on an annual basis, and during the period in which he receives benefits Employee dies, his beneficiary shall be entitled to the balance, if any, as if Employee had taken payments pursuant to Article III, or as Article III may be amended by Article IV. The beneficiary shall receive the payment provided for in either 3.2 or 3.3 or as 3.3 may be amended by Article IV, for fifteen (15) years, less the number of years during which Employee received payments pursuant to 5.1.

VI. DEATH BENEFIT

- 6.1 If the Employee dies prior to his Retirement, and while employed by Mylan, the Company shall pay to his beneficiary the sum of one million six hundred thousand dollars (\$1,600,000.00) within thirty (30) days of Employee's death.
- 6.2 In addition to the benefits set forth in 6.1, the Company shall maintain for Employee's benefit during his employment with the Company life insurance policies in the aggregate amount of one million six hundred forty five thousand dollars (\$1,645,000.00).
- 6.3 If the Employee's death occurs after Retirement, but before having received the entire benefit provided for under Article III (or as Article III may be amended by Article IV) or Article V, the balance of the fifteen (15) payments due thereunder shall be paid to Employee's beneficiary in a lump-sum payment equal to the NPV of the remaining payments due.

VII. SUCCESSORSHIP

This Agreement in its entirety shall be binding upon and enforceable against the Company and its Successors.

VIII. NO DUPLICATION OF PAYMENTS

Notwithstanding anything to the contrary which may be set forth elsewhere herein, under no circumstances is Employee or his beneficiary entitled to take benefits under more than any one article included in this Agreement, except in so far as Article III may be amended by Article IV.

IX. EMPLOYEE CONDUCT WITH RESPECT TO COMPETITORS

- 9.1 Employee agrees that he will not, without the prior written consent of the Company, directly or indirectly, whether as an employee, officer, director, independent contractor, consultant, stockholder, partner or otherwise, engage in or assist others to engage in or have any interest in any business which competes with the Company in any geographic area in which the Company markets or has marketed its products during the year preceding termination of Puskar's employment for the greater of:
- (a) the period during which Employee receives monthly payments under this Agreement; or
 - (b) five (5) years following his receipt of a lump-sum payment hereunder.
- 9.2 Notwithstanding anything to the contrary set forth elsewhere herein, stock ownership in a competing business shall not be a breach of this Agreement, provided such stock is traded on a national exchange.
- 9.3 The Parties agree and acknowledge that the time, scope and geographic area and other provisions of this Agreement have been specifically negotiated by the Parties, and Employee specifically hereby agrees that such time, scope and geographic area and other provisions are reasonable under these circumstances. Employee further agrees that if, despite the express agreement of the Parties to this Agreement, a court should hold any portion of this Agreement unenforceable for any reason, the maximum restrictions of time, scope and geographic area reasonable under the circumstances, as determined by the court, will be substituted for the restrictions herein which such court may find to be unreasonable or unenforceable.
- 9.4 The Parties acknowledge that the breach of 9.1 will be such that the Company will not have an adequate remedy at law because the rights of the Company under this Agreement are of a specialized and unique character, and that immediate and irreparable damage will result to the Company if Employee breaches his obligations under 9.1. The Company may, in addition to any other remedies and damages available, seek an injunction in the courts of the State of West Virginia and the United States District Court for the Northern District of West Virginia to restrain any such breach. Employee represents and warrants that his expertise and capabilities are such that his obligations under 9.1 will not prevent him from earning a living.

X. CONSULTING SERVICES

- 10.1 During the ten (10) year period beginning on the day following Employee's Retirement he shall, at the request of the Company, act in the capacity of a consultant for the Company, performing such services as may be consistent with those performed by him during his employment. These services may be designated by the President of the Company, or his authorized representative, and shall be reasonable in scope, duration and frequency.
- 10.2 The Company shall pay the Employee for such consulting services an hourly rate to be determined by the Parties at such time, but not less than two hundred fifty dollars (\$250.00) per hour, payable monthly.
- 10.3 In addition to the foregoing, the Company shall reimburse the Employee monthly for any and all out-of-pocket expenses incurred by the Employee directly for the benefit of the business of the Company.

XI. ELIGIBILITY FOR PAYMENT

- 11.1 Any and all payments due hereunder, may be denied if not already begun, or terminated if they have begun, if in the Company's sole judgment Employee is either not eligible for such payments, or once such payments have begun is found to be or found to have been ineligible.
- 11.2 Employee shall not be eligible for any payments hereunder if the Company, in its sole discretion, finds that during or subsequent to his employment with the Company he:

- (a) breaches, or has breached any term, provision or obligation enumerated herein;
- (b) committed any act by commission or omission which materially and substantially adversely affects the Company's business or reputation; or
- (c) is convicted of any violation of the Federal Food, Drug and Cosmetic Act, or the violation of any other statute relevant to the Company's business.
- 11.3 Should Employee be paid any benefits hereunder and thereafter be found ineligible, or to have been ineligible, he must return to the Company that portion of the benefit paid to him for the period of his ineligibility.

XII. RIGHT TO CONFER

- 12.1 Employee shall have the right, but not the obligation to:
 - (a) Confer with any Advisor of his choice prior to signing the Agreement; and
 - (b) Provide his Advisors with a true and complete copy of the Agreement, and any other pertinent documents which may be of assistance to his Advisors.
- 12.2 Should Employee decline the right to confer with his Advisors prior to executing this Agreement he shall execute an acknowledgement of same which shall then be incorporated herein by reference. (See Exhibit A)

XIII. NO PROMISE OF CONTINUED EMPLOYMENT

13.1 Employee acknowledges his employment with the Company is AT-WILL.

13.2 Nothing set forth herein shall constitute or be construed as a contract of employment except in so far as provided for under 13.1 hereof.

XIV. RESTRICTION OF ALIENABILITY

Benefits payable to the Employee or beneficiary shall not be subject to assignment, transfer, attachment, execution, garnishment, sequestration, or any other seizure under any legal or equitable process, whether on account of the Employee's or beneficiary's act or by operation of the law.

XV. CONTRACT ADMINISTRATOR

The Director of Taxation and Accounting of the Company, or other officer of Mylan designated by the Executive Committee of the Company is hereby named the Contract Administrator for purposes of assuring compliance with the terms and conditions set forth herein.

XVI. MODIFICATION

Except as provided for in Articles IV and IX, this Agreement may not be changed, amended or otherwise modified other than by a written statement; Provided, such statement is signed by both Parties, expresses their intent to change the Agreement, and specifically describes such changes.

XVII. HEADINGS

Except when referenced in the body of this Agreement article headings are set forth herein for the purpose of convenience only. Such headings shall not be considered or otherwise referred to when any question or issue arises with respect to the application or interpretation of any term or condition set forth herein.

XVIII. COUNTERPARTS

This Agreement may be executed in two or more counterparts, each of which is to be considered an original, and taken together as one and the same document.

XIX. GOVERNING LAW

Any an all actions between the Parties regarding the interpretation or application of any term or provision set forth herein shall be governed by and interpreted in accordance with the substantive laws, and not the law of conflicts, of the State of West Virginia. The Company and Employee each do hereby respectively consent and agree that the courts of the State of West Virginia shall have jurisdiction, and venue shall properly lie with the courts of the State of West Virginia, with respect to any and all actions brought hereunder.

XX. SINGULAR OR PLURAL

The singular form of any noun or pronoun shall include the plural when the context in which such word is used is such that it is apparent the singular is intended to include the plural and vice versa.

XXI. ASSIGNMENT

The Agreement may not be assigned by either Party, without the written authorization of the other Party. A Successor shall not be considered an assignee for purposes of this Article.

XXII. ENTIRE AGREEMENT

The terms and conditions set forth herein contain the entire agreement between the Company and Employee, and supersede any and all prior agreements or understandings (whether express or implied) between the Parties with respect to the matters set forth herein.

XXIII. SURVIVAL

Articles I, II, IX, X, XI, XIX and XXIII shall survive any expiration or termination of this Agreement.

XXIV. TERM

The term of this Agreement shall begin on the Effective Date and shall end on the date on which Mylan makes the last payment to which it is obligated hereunder.

IN WITNESS of their agreement to the terms and conditions set forth herein the Company and Employee have caused the following signatures to be affixed hereto: $\frac{1}{2} \left(\frac{1}{2} \right) = \frac{1}{2} \left(\frac{1}{2} \right) \left(\frac$

MYLAN LABORATORIES INC.	MILAN PUSKAR
BY:	BY:
TITLE:	DATE:
DATE:	

ACKNOWLEDGEMENT

I, Milan Puskar, hereby acknowledge and understand that I have been given the opportunity to review and discuss the terms of the Agreement with an Advisor of my choice prior to signing the Agreement. I have decided that it is not necessary for me to discuss this matter with an Advisor, and therefore I decline the opportunity to do so.

No one has discouraged me or otherwise attempted to influence me with regard to my decision to decline the opportunity to discuss this matter with my Advisors. My refusal of this offer is entirely my own and is made without any reservation or conditions whatsoever.

WITNESSED BY:			
	DATE:	BY:	DATE:
	DATE:		

RETIREMENT BENEFIT AGREEMENT

This Retirement Benefit Agreement (the "Agreement") is entered into on this 27th day of January, 1995 (the "Effective Date") by and between:

Mylan Laboratories Inc., a Pennsylvania Corporation, with offices located at 781 Chestnut Ridge Road, Morgantown, WV 26505 (hereinafter referred to as "Mylan" or "Company").

and

Dana G. Barnett, an employee of Mylan who resides at 923 Anchorage Road, Tampa, FL 33602 (hereinafter referred to as "Employee or "Barnett").

WHEREAS the Company and Employee, in recognition of Employee's long and valuable contribution to the success of the Company, entered into a Salary Continuation Agreement on April 1, 1990; and

WHEREAS Employee continues to perform valuable services for the Company; and

WHEREAS in recognition of his continuing service to Mylan, the Company wishes to provide Employee with financial assistance with respect to certain Contingencies, in addition to that provided for in said April 1, 1990 Agreement; and

WHEREAS the Company and Employee wish to RESCIND, and to REPLACE said Salary Continuation Agreement with this Agreement;

WITNESSETH THEREFORE that in consideration of the additional benefits provided for hereunder, the premises and covenants set forth herein, and other good and valuable consideration, the sufficiency of which is hereby acknowledged, the Company and Employee, intending to be legally bound, agree as follows:

I. DEFINITIONS

Whenever used in the Agreement the following terms shall be defined as follows:

- (a) "Advisor" or "Advisors" shall mean with respect to Employee any person including lawyers, accountants, estate planners and others, with whom he may wish to review and discuss the matters set forth herein.
- (b) "Agreement" shall mean this Retirement Benefit Agreement which is entered into on the $27 \mathrm{th}$ day of January, 1995.
- (c) "At-Will" shall mean with respect to the period of Barnett's employment with Mylan, that the Company is under no obligation to continue to employ Barnett for any period of time, and can terminate his employment at any time without notice, subject to certain statutory and regulatory requirements; and that Employee is under no obligation to remain employed by the Company, and can terminate his employment with Mylan at any time, without notice.
 - (d) "Contingency" shall mean Retirement or death.
- (e) "Mylan" or "Company" shall mean Mylan Laboratories Inc., its subsidiaries and affiliates.
- (f) "Net Present Value" ("NPV") shall mean the present value at any given time of the entire benefit to be paid, discounted at seven percent (7%) per annum.
- (g) "Party" or "Parties" shall mean the Company or Employee, or both the Company and Employee depending upon which term is required by the context in which it is used.
- (h) "Retire" or "Retirement" shall mean the day and date on which Barnett's employment with the Company is terminated by either Party for any reason other than death of the Employee.
- (i) "Successor" shall be mean any person, partnership, limited partnership, joint-venture, corporation, trust or any other entity or organization who, subsequent to the Effective Date, comes into possession of or acquires, either directly or indirectly, all or substantially all of the Company's business, assets or voting stock, or the right to direct the business activities and practices of the Company.

II. RESCISSION OF PRIOR AGREEMENT

The Salary Continuation Agreement entered into by the Parties on April 1, 1990 and any and all other agreements, express or implied, which may have been entered into by them prior to the execution of this Agreement, including by way of example and not of limitation, any agreement which addresses or is related to salary continuation, deferred compensation, employment, or similar matters (excluding agreements related to stock options) are hereby RESCINDED by the mutual consent of the Parties hereto upon execution of this Agreement.

THEREFORE THE PARTIES ACKNOWLEDGE THAT ANY RIGHTS AND OBLIGATIONS SET FORTH IN ANY SUCH AGREEMENT, WHETHER EXPRESS OR IMPLIED, ARE FOREVER WAIVED, NULL AND VOID, AND UNENFORCEABLE AT LAW OR IN EQUITY.

III. RETIREMENT

- 3.1 Upon his Retirement from the Company, and if he is eligible to receive payments as provided for elsewhere herein, Employee shall receive an annual retirement benefit equal to the amount set forth below.
- 3.2 Should Employee Retire after the Effective Date but on or before March 31, 1996 he shall receive one hundred fifty thousand dollars (\$150,000.00) each year for fifteen (15) years.
- 3.3 Should Employee Retire after March 31, 1996 he shall receive one hundred eighty thousand dollars (\$180,000.00) each year for fifteen (15) years.
- 3.4 The Company shall pay each of the fifteen (15) annual payments due hereunder in twelve (12) equal or substantially equal installments. The first of any such payments shall be made on the first day of the month following the month in which Employee Retires, and each subsequent payment shall be made on the first day of each successive month until Mylan's obligations with respect to such payments have been satisfied.
- 3.5 However, upon the written request of the Employee, Mylan may pay to Employee the NPV of any amount, or of the balance of any such amount, to which he is entitled hereunder in a lump-sum payment. Said payment shall be paid within thirty (30) days of the date of Employee's request for such payment.

IV. AWARD OF ADDITIONAL RETIREMENT BENEFITS

- 4.1 The Company may increase the retirement benefit to which Employee is entitled under Article III. Any such increases shall be subject to the following conditions:
- (a) the maximum benefit to which Employee is entitled under Article III shall not exceed three hundred thousand dollars (\$300,000.00) per year for fifteen (15) years;
 - (b) no increases shall be granted earlier than April 1, 1996 or later than April 1, 2000;
 - (c) any increase shall be expensed for accounting purposes at the time it is granted;
- (d) the Company's obligation to pay the increased benefit to Employee, if and when such increases are awarded shall be subject to all the terms and conditions set forth herein; and
- (e) the inclusion herein of any increase in Employee's retirement benefit shall become effective upon Employee's receipt of a notice from the Company informing him of the additional benefit.

V. CAPACITY TO PERFORM DUTIES

- 5.1 Should Employee become unable to perform the material and substantial duties of his position the Company shall pay to Employee three hundred thousand dollars (\$300,000.00) each year for fifteen (15) years in equal or substantially equal payments. However, upon the written request of the Employee, Mylan may pay to Employee the NPV of any amount, or of the balance of any such amount, to which he is entitled to hereunder in a lump-sum payment. Said payment shall be paid within thirty (30) days of the date of Employee's request for such payment.
- 5.2 The certification of a licensed physician as to Employee's inability to perform the material and substantial duties of his position shall be conclusive with respect to his status, and the Company agrees to be unequivocally bound by any such certification.
- 5.3 If the Employee receives the benefit described in 5.1 on an annual basis, and during the period in which he receives benefits Employee dies, his beneficiary shall be entitled to the balance, if any, as if Employee had taken payments pursuant to Article III, or as Article III may be amended by Article IV. The beneficiary shall receive the payment provided for in either 3.2 or 3.3 or as 3.3 may be amended by Article IV, for fifteen (15) years, less the number of years during which Employee received payments pursuant to 5.1.

VI. DEATH BENEFIT

- 6.1 The Company shall maintain for Employee's benefit during his employment with the Company life insurance policies in the aggregate amount of one million five hundred thousand dollars (\$1,500,000.00).
- 6.2 If the Employee's death occurs after Retirement, but before having received the entire benefit provided for under Article III (or as Article III may be amended by Article IV) or Article V, the balance of the fifteen (15) payments due thereunder shall be paid to Employee's beneficiary in a lump-sum payment equal to the NPV of the remaining payments due.

VII. SUCCESSORSHIP

This Agreement in its entirety shall be binding upon and enforceable against the Company and its Successors.

VIII. NO DUPLICATION OF PAYMENTS

Notwithstanding anything to the contrary which may be set forth elsewhere herein, under no circumstances is Employee or his beneficiary entitled to take benefits under more than any one article included in this Agreement, except in so far as Article III may be amended by Article IV.

IX. EMPLOYEE CONDUCT WITH RESPECT TO COMPETITORS

- 9.1 Employee agrees that he will not, without the prior written consent of the Company, directly or indirectly, whether as an employee, officer, director, independent contractor, consultant, stockholder, partner or otherwise, engage in or assist others to engage in or have any interest in any business which competes with the Company in any geographic area in which the Company markets or has marketed its products during the year preceding termination of Barnett's employment for the greater of:
- (a) the period during which Employee receives monthly payments under this Agreement; or
 - (b) five (5) years following his receipt of a lump-sum payment hereunder.
- 9.2 Notwithstanding anything to the contrary set forth elsewhere herein, stock ownership in a competing business shall not be a breach of this Agreement, provided such stock is traded on a national exchange.

- 9.3 The Parties agree and acknowledge that the time, scope and geographic area and other provisions of this Agreement have been specifically negotiated by the Parties, and Employee specifically hereby agrees that such time, scope and geographic area and other provisions are reasonable under these circumstances. Employee further agrees that if, despite the express agreement of the Parties to this Agreement, a court should hold any portion of this Agreement unenforceable for any reason, the maximum restrictions of time, scope and geographic area reasonable under the circumstances, as determined by the court, will be substituted for the restrictions herein which such court may find to be unreasonable or unenforceable.
- 9.4 The Parties acknowledge that the breach of 9.1 will be such that the Company will not have an adequate remedy at law because the rights of the Company under this Agreement are of a specialized and unique character, and that immediate and irreparable damage will result to the Company if Employee breaches his obligations under 9.1. The Company may, in addition to any other remedies and damages available, seek an injunction in the courts of the State of West Virginia and the United States District Court for the Northern District of West Virginia to restrain any such breach. Employee represents and warrants that his expertise and capabilities are such that his obligations under 9.1 will not prevent him from earning a living.

X. CONSULTING SERVICES

- 10.1 During the ten (10) year period beginning on the day following Employee's Retirement he shall, at the request of the Company, act in the capacity of a consultant for the Company, performing such services as may be consistent with those performed by him during his employment. These services may be designated by the President of the Company, or his authorized representative, and shall be reasonable in scope duration and frequency.
- 10.2 The Company shall pay the Employee for such consulting services an hourly rate to be determined by the Parties at such time, but not less than one hundred fifty dollars (\$150.00) per hour, payable monthly.
- 10.3 In addition to the foregoing, the Company shall reimburse the Employee monthly for any and all out-of-pocket expenses incurred by the Employee directly for the benefit of the business of the Company.

XI. ELIGIBILITY FOR PAYMENT

11.1 Any and all payments due hereunder, may be denied if not already begun, or terminated if they have begun, if in the Company's sole judgment Employee is either not eligible for such payments, or once such payments have begun is found to be or found to have been ineligible.

- 11.2 Employee shall not be eligible for any payments hereunder if the Company, in its sole discretion, finds that during or subsequent to his employment with the Company he:
 - (a) breaches, or has breached any term, provision or obligation enumerated herein;
- (b) committed any act by commission or omission which materially and substantially adversely affects the Company's business or reputation; or
- (c) is convicted of any violation of the Federal Food, Drug and Cosmetic Act, or the violation of any other statute relevant to the Company's business.
- 11.3 Should Employee be paid any benefits hereunder and thereafter be found ineligible, or to have been ineligible, he must return to the Company that portion of the benefit paid to him for the period of his ineligibility.

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- 12.1 Employee shall have the right, but not the obligation to:

 - (a) Confer with any Advisor of his choice prior to signing the Agreement; and(b) Provide his Advisors with a true and complete copy of the Agreement, and any other pertinent documents which may be of assistance to his Advisors.
- 12.2 Should Employee decline the right to confer with his Advisors prior to executing this Agreement he shall execute an acknowledgement of same which shall then be incorporated herein by reference. (See Exhibit A)

XIII. NO PROMISE OF CONTINUED EMPLOYMENT

- 13.1 Employee acknowledges his employment with the Company is AT-WILL.
- 13.2 Nothing set forth herein shall constitute or be construed as a contract of employment except in so far as provided for under 13.1 hereof.

XIV. RESTRICTION OF ALIENABILITY

Benefits payable to the Employee or beneficiary shall not be subject to assignment, transfer, attachment, execution, garnishment, sequestration, or any other seizure under any legal or equitable process, whether on account of the Employee's or beneficiary's act or by operation of the law.

XV. CONTRACT ADMINISTRATOR

The Director of Taxation and Accounting of the Company, or other officer of Mylan designated by the Executive Committee of the Company is hereby named the Contract Administrator for purposes of assuring compliance with the terms and conditions set forth herein.

XVI. MODIFICATION

Except as provided for in Articles IV and IX, this Agreement may not be changed, amended or otherwise modified other than by a written statement; Provided, such statement is signed by both Parties, expresses their intent to change the Agreement, and specifically describes such changes.

XVII. HEADINGS

Except when referenced in the body of this Agreement article headings are set forth herein for the purpose of convenience only. Such headings shall not be considered or otherwise referred to when any question or issue arises with respect to the application or interpretation of any term or condition set forth herein.

XVIII. COUNTERPARTS

This Agreement may be executed in two or more counterparts, each of which is to be considered an original, and taken together as one and the same document.

XIX. GOVERNING LAW

Any an all actions between the Parties regarding the interpretation or application of any term or provision set forth herein shall be governed by and interpreted in accordance with the substantive laws, and not the law of conflicts, of the State of West Virginia. The Company and Employee each do hereby respectively consent and agree that the courts of the State of West Virginia shall have jurisdiction, and venue shall properly lie with the courts of the State of West Virginia, with respect to any and all actions brought hereunder.

XX. SINGULAR OR PLURAL

The singular form of any noun or pronoun shall include the plural when the context in which such word is used is such that it is apparent the singular is intended to include the plural and vice versa.

XXI. ASSIGNMENT

The Agreement may not be assigned by either Party, without the written authorization of the other Party. A Successor shall not be considered an assignee for purposes of this Article.

XXII. ENTIRE AGREEMENT

The terms and conditions set forth herein contain the entire agreement between the Company and Employee, and supersede any and all prior agreements or understandings (whether express or implied) between the Parties with respect to the matters set forth herein.

XXIII. SURVIVAL

Articles I, II, IX, X, XI, XIX and XXIII shall survive any expiration or termination of this Agreement.

XXIV. TERM

The term of this Agreement shall begin on the Effective Date and shall end on the date on which Mylan makes the last payment to which it is obligated hereunder.

IN WITNESS of their agreement to the terms and conditions set forth herein the Company and Employee have caused the following signatures to be affixed hereto:

MYLAN LABORATORIES INC.	DANA G. BARNETT	
BY:	BY:	
TITLE:	DATE:	
DATE:		

ACKNOWLEDGEMENT

I, Dana G. Barnett, hereby acknowledge and understand that I have been given the opportunity to review and discuss the terms of the Agreement with an Advisor of my choice prior to signing the Agreement. I have decided that it is not necessary for me to discuss this matter with an Advisor, and therefore I decline the opportunity to do so.

No one has discouraged me or otherwise attempted to influence me with regard to my decision to decline the opportunity to discuss this matter with my Advisors. My refusal of this offer is entirely my own and is made without any reservation or conditions whatsoever.

WITNESSED	BY:			
		DATE:	BY:	DATE:
		DATE:		

RETIREMENT BENEFIT AGREEMENT

This Retirement Benefit Agreement (the "Agreement") is entered into on this 27th day of January, 1995 (the "Effective Date") by and between:

Mylan Laboratories Inc., a Pennsylvania Corporation, with offices located at 781 Chestnut Ridge Road, Morgantown, WV 26505 (hereinafter referred to as "Mylan" or "Company").

and

Clarence B. Todd, an employee of Mylan who resides at 38 Lakeview Drive, Morgantown, WV 26505 (hereinafter referred to as "Employee or "Todd").

WHEREAS the Company and Employee, in recognition of Employee's long and valuable contribution to the success of the Company, entered into a Salary Continuation Agreement on April 1, 1990; and

WHEREAS Employee continues to perform valuable services for the Company; and

WHEREAS in recognition of his continuing service to Mylan, the Company wishes to provide Employee with financial assistance with respect to certain Contingencies, in addition to that provided for in said April 1, 1990 Agreement; and

WHEREAS the Company and Employee wish to RESCIND, and to REPLACE said Salary Continuation Agreement with this Agreement;

WITNESSETH THEREFORE that in consideration of the additional benefits provided for hereunder, the premises and covenants set forth herein, and other good and valuable consideration, the sufficiency of which is hereby acknowledged, the Company and Employee, intending to be legally bound, agree as follows:

I. DEFINITIONS

Whenever used in the Agreement the following terms shall be defined as follows:

- (a) "Advisor" or "Advisors" shall mean with respect to Employee any person including lawyers, accountants, estate planners and others, with whom he may wish to review and discuss the matters set forth herein.
- (b) "Agreement" shall mean this Retirement Benefit Agreement which is entered into on the $27 \mathrm{th}$ day of January, 1995.
- (c) "At-Will" shall mean with respect to the period of Todd's employment with Mylan, that the Company is under no obligation to continue to employ Todd for any period of time, and can terminate his employment at any time without notice, subject to certain statutory and regulatory requirements; and that Employee is under no obligation to remain employed by the Company, and can terminate his employment with Mylan at any time, without notice.
 - (d) "Contingency" shall mean Retirement or death.
- (e) "Mylan" or "Company" shall mean Mylan Laboratories Inc., its subsidiaries and affiliates.
- (f) "Net Present Value" ("NPV") shall mean the present value at any given time of the entire benefit to be paid, discounted at seven percent (7%) per annum.
- (g) "Party" or "Parties" shall mean the Company or Employee, or both the Company and Employee depending upon which term is required by the context in which it is used.
- (h) "Retire" or "Retirement" shall mean the day and date on which Todd's employment with the Company is terminated by either Party for any reason other than death of the Employee.
- (i) "Successor" shall mean any person, partnership, limited partnership, joint-venture, corporation, trust or any other entity or organization who, subsequent to the Effective Date, comes into possession of or acquires, either directly or indirectly, all or substantially all of the Company's business, assets or voting stock, or the right to direct the business activities and practices of the Company.

II. RESCISSION OF PRIOR AGREEMENT

The Salary Continuation Agreement entered into by the Parties on April 1, 1990 and any and all other agreements, express or implied, which may have been entered into by them prior to the execution of this Agreement, including by way of example and not of limitation, any agreement which addresses or is related to salary continuation, deferred compensation, employment, or similar matters (excluding agreements related to stock options) are hereby RESCINDED by the mutual consent of the Parties hereto upon execution of this Agreement.

THEREFORE THE PARTIES ACKNOWLEDGE THAT ANY RIGHTS AND OBLIGATIONS SET FORTH IN ANY SUCH AGREEMENT, WHETHER EXPRESS OR IMPLIED, ARE FOREVER WAIVED, NULL AND VOID, AND UNENFORCEABLE AT LAW OR IN EQUITY.

III. RETIREMENT

- 3.1 Upon his Retirement from the Company, and if he is eligible to receive payments as provided for elsewhere herein, Employee shall receive an annual retirement benefit equal to the amount set forth below.
- 3.2 Should Employee Retire after the Effective Date but on or before March 31, 1996 he shall receive one hundred fifty thousand dollars (\$150,000.00) each year for fifteen (15) years.
- 3.3 Should Employee Retire after March 31, 1996 he shall receive one hundred eighty thousand dollars (\$180,000.00) each year for fifteen (15) years.
- 3.4 The Company shall pay each of the fifteen (15) annual payments due hereunder in twelve (12) equal or substantially equal installments. The first of any such payments shall be made on the first day of the month following the month in which Employee Retires, and each subsequent payment shall be made on the first day of each successive month until Mylan's obligations with respect to such payments have been satisfied.
- 3.5 However, upon the written request of the Employee, Mylan may pay to Employee the NPV of any amount, or of the balance of any such amount, to which he is entitled hereunder in a lump-sum payment. Said payment shall be paid within thirty (30) days of the date of Employee's request for such payment.

IV. AWARD OF ADDITIONAL RETIREMENT BENEFITS

- 4.1 The Company may increase the retirement benefit to which Employee is entitled under Article III. Any such increases shall be subject to the following conditions:
- (a) the maximum benefit to which Employee is entitled under Article III shall not exceed three hundred thousand dollars (\$300,000.00) per year for fifteen (15) years;
 - (b) no increases shall be granted earlier than April 1, 1996 or later than April 1, 2000;
 - (c) any increase shall be expensed for accounting purposes at the time it is granted;
- (d) the Company's obligation to pay the increased benefit to Employee, if and when such increases are awarded shall be subject to all the terms and conditions set forth herein; and
- (e) the inclusion herein of any increase in Employee's retirement benefit shall become effective upon Employee's receipt of a notice from the Company informing him of the additional benefit.

V. CAPACITY TO PERFORM DUTIES

- 5.1 Should Employee become unable to perform the material and substantial duties of his position the Company shall pay to Employee three hundred thousand dollars (\$300,000.00) each year for fifteen (15) years in equal or substantially equal payments. However, upon the written request of the Employee, Mylan may pay to Employee the NPV of any amount, or of the balance of any such amount, to which he is entitled to hereunder in a lump-sum payment. Said payment shall be paid within thirty (30) days of the date of Employee's request for such payment.
- 5.2 The certification of a licensed physician as to Employee's inability to perform the material and substantial duties of his position shall be conclusive with respect to his status, and the Company agrees to be unequivocally bound by any such certification.
- 5.3 If the Employee receives the benefit described in 5.1 on an annual basis, and during the period in which he receives benefits Employee dies, his beneficiary shall be entitled to the balance, if any, as if Employee had taken payments pursuant to Article III, or as Article III may be amended by Article IV. The beneficiary shall receive the payment provided for in either 3.2 or 3.3 or as 3.3 may be amended by Article IV, for fifteen (15) years, less the number of years during which Employee received payments pursuant to 5.1.

VI. DEATH BENEFIT

- 6.1 The Company shall maintain for Employee's benefit during his employment with the Company life insurance policies in the aggregate amount of one million five hundred thousand dollars (\$1,500,000.00).
- 6.2 If the Employee's death occurs after Retirement, but before having received the entire benefit provided for under Article III (or as Article III may be amended by Article IV) or Article V, the balance of the fifteen (15) payments due thereunder shall be paid to Employee's beneficiary in a lump-sum payment equal to the NPV of the remaining payments due.

VII. SUCCESSORSHIP

This Agreement in its entirety shall be binding upon and enforceable against the Company and its Successors.

VIII. NO DUPLICATION OF PAYMENTS

Notwithstanding anything to the contrary which may be set forth elsewhere herein, under no circumstances is Employee or his beneficiary entitled to take benefits under more than any one article included in this Agreement, except in so far as Article III may be amended by Article IV.

IX. EMPLOYEE CONDUCT WITH RESPECT TO COMPETITORS

- 9.1 Employee agrees that he will not, without the prior written consent of the Company, directly or indirectly, whether as an employee, officer, director, independent contractor, consultant, stockholder, partner or otherwise, engage in or assist others to engage in or have any interest in any business which competes with the Company in any geographic area in which the Company markets or has marketed its products during the year preceding termination of Todd's employment for the greater of:
- (a) the period during which Employee receives monthly payments under this Agreement; or
 - (b) five (5) years following his receipt of a lump-sum payment hereunder.
- 9.2 Notwithstanding anything to the contrary set forth elsewhere herein, stock ownership in a competing business shall not be a breach of this Agreement, provided such stock is traded on a national exchange.

- 9.3 The Parties agree and acknowledge that the time, scope and geographic area and other provisions of this Agreement have been specifically negotiated by the Parties, and Employee specifically hereby agrees that such time, scope and geographic area and other provisions are reasonable under these circumstances. Employee further agrees that if, despite the express agreement of the Parties to this Agreement, a court should hold any portion of this Agreement unenforceable for any reason, the maximum restrictions of time, scope and geographic area reasonable under the circumstances, as determined by the court, will be substituted for the restrictions herein which such court may find to be unreasonable or unenforceable.
- 9.4 The Parties acknowledge that the breach of 9.1 will be such that the Company will not have an adequate remedy at law because the rights of the Company under this Agreement are of a specialized and unique character, and that immediate and irreparable damage will result to the Company if Employee breaches his obligations under 9.1. The Company may, in addition to any other remedies and damages available, seek an injunction in the courts of the State of West Virginia and the United States District Court for the Northern District of West Virginia to restrain any such breach. Employee represents and warrants that his expertise and capabilities are such that his obligations under 9.1 will not prevent him from earning a living.

X. CONSULTING SERVICES

- 10.1 During the ten (10) year period beginning on the day following Employee's Retirement he shall, at the request of the Company, act in the capacity of a consultant for the Company, performing such services as may be consistent with those performed by him during his employment. These services may be designated by the President of the Company, or his authorized representative, and shall be reasonable in scope duration and frequency.
- 10.2 The Company shall pay the Employee for such consulting services an hourly rate to be determined by the Parties at such time, but not less than one hundred fifty dollars (\$150.00) per hour, payable monthly.
- 10.3 In addition to the foregoing, the Company shall reimburse the Employee monthly for any and all out-of-pocket expenses incurred by the Employee directly for the benefit of the business of the Company.

XI. ELIGIBILITY FOR PAYMENT

11.1 Any and all payments due hereunder, may be denied if not already begun, or terminated if they have begun, if in the Company's sole judgment Employee is either not eligible for such payments, or once such payments have begun is found to be or found to have been ineligible.

- 11.2 Employee shall not be eligible for any payments hereunder if the Company, in its sole discretion, finds that during or subsequent to his employment with the Company he:
 - (a) breaches, or has breached any term, provision or obligation enumerated herein;
- (b) committed any act by commission or omission which materially and substantially adversely affects the Company's business or reputation; or
- (c) is convicted of any violation of the Federal Food, Drug and Cosmetic Act, or the violation of any other statute relevant to the Company's business.
- 11.3 Should Employee be paid any benefits hereunder and thereafter be found ineligible, or to have been ineligible, he must return to the Company that portion of the benefit paid to him for the period of his ineligibility.

XII. RIGHT TO CONFER

- 12.1 Employee shall have the right, but not the obligation to:

 - (a) Confer with any Advisor of his choice prior to signing the Agreement; and(b) Provide his Advisors with a true and complete copy of the Agreement, and any other pertinent documents which may be of assistance to his Advisors.
- 12.2 Should Employee decline the right to confer with his Advisors prior to executing this Agreement he shall execute an acknowledgement of same which shall then be incorporated herein by reference. (See Exhibit A)

XIII. NO PROMISE OF CONTINUED EMPLOYMENT

- 13.1 Employee acknowledges his employment with the Company is AT-WILL.
- 13.2 Nothing set forth herein shall constitute or be construed as a contract of employment except in so far as provided for under 13.1 hereof.

XIV. RESTRICTION OF ALIENABILITY

Benefits payable to the Employee or beneficiary shall not be subject to assignment, transfer, attachment, execution, garnishment, sequestration, or any other seizure under any legal or equitable process, whether on account of the Employee's or beneficiary's act or by operation of the law.

XV. CONTRACT ADMINISTRATOR

The Director of Taxation and Accounting of the Company, or other officer of Mylan designated by the Executive Committee of the Company is hereby named the Contract Administrator for purposes of assuring compliance with the terms and conditions set forth herein.

XVI. MODIFICATION

Except as provided for in Articles IV and IX, this Agreement may not be changed, amended or otherwise modified other than by a written statement; Provided, such statement is signed by both Parties, expresses their intent to change the Agreement, and specifically describes such changes.

XVII. HEADINGS

Except when referenced in the body of this Agreement article headings are set forth herein for the purpose of convenience only. Such headings shall not be considered or otherwise referred to when any question or issue arises with respect to the application or interpretation of any term or condition set forth herein.

XVIII. COUNTERPARTS

This Agreement may be executed in two or more counterparts, each of which is to be considered an original, and taken together as one and the same document.

XIX. GOVERNING LAW

Any an all actions between the Parties regarding the interpretation or application of any term or provision set forth herein shall be governed by and interpreted in accordance with the substantive laws, and not the law of conflicts, of the State of West Virginia. The Company and Employee each do hereby respectively consent and agree that the courts of the State of West Virginia shall have jurisdiction, and venue shall properly lie with the courts of the State of West Virginia, with respect to any and all actions brought hereunder.

XX. SINGULAR OR PLURAL

The singular form of any noun or pronoun shall include the plural when the context in which such word is used is such that it is apparent the singular is intended to include the plural and vice versa.

XXI. ASSIGNMENT

The Agreement may not be assigned by either Party, without the written authorization of the other Party. A Successor shall not be considered an assignee for purposes of this Article.

XXII. ENTIRE AGREEMENT

The terms and conditions set forth herein contain the entire agreement between the Company and Employee, and supersede any and all prior agreements or understandings (whether express or implied) between the Parties with respect to the matters set forth herein.

XXIII. SURVIVAL

Articles I, II, IX, X, XI, XIX and XXIII shall survive any expiration or termination of this Agreement.

XXIV. TERM

The term of this Agreement shall begin on the Effective Date and shall end on the date on which Mylan makes the last payment to which it is obligated hereunder.

IN WITNESS of their agreement to the terms and conditions set forth herein the Company and Employee have caused the following signatures to be affixed hereto:

MYLAN LABORATORIES INC.	CLARENCE B. TODD	
BY:	BY:	
TITLE:	DATE:	
DATE:		

ACKNOWLEDGEMENT

I, Clarence B. Todd, hereby acknowledge and understand that I have been given the opportunity to review and discuss the terms of the Agreement with an Advisor of my choice prior to signing the Agreement. I have decided that it is not necessary for me to discuss this matter with an Advisor, and therefore I decline the opportunity to do so.

No one has discouraged me or otherwise attempted to influence me with regard to my decision to decline the opportunity to discuss this matter with my Advisors. My refusal of this offer is entirely my own and is made without any reservation or conditions whatsoever.

WITNESSED BY:			
	DATE:	BY:	DATE:
	DATE:		

(c) "Salary Continuation Plan" with Roderick P. Jackson and Louis J. DeBone each dated March 14, 1995 and filed herewith.

RETIREMENT BENEFIT AGREEMENT

This Retirement Benefit Agreement (the "Agreement") is entered into on this 14th day of March, 1995 (the "Effective Date") by and between:

Mylan Laboratories Inc., a Pennsylvania Corporation, with offices located at 781 Chestnut Ridge Road, Morgantown, WV 26505 (hereinafter referred to as "Mylan" or "Company").

and

Roderick P. Jackson, an employee of Mylan who resides at 1456 Lakeland Avenue, Morgantown, WV 26505 (hereinafter referred to as "Employee" or "Jackson").

WHEREAS the Company and Employee, in recognition of Employee 's long and valuable contribution to the success of the Company, entered into a Salary Continuation Agreement on April 1, 1989; and

WHEREAS Employee continues to perform valuable services for the Company; and

WHEREAS in recognition of his continuing service to Mylan, the Company wishes to provide Employee with financial assistance with respect to certain Contingencies, in addition to that provided for in said April 1, 1989 Agreement; and

WHEREAS the Company and Employee wish to RESCIND, and to REPLACE said Salary Continuation Agreement with this Agreement;

WITNESSETH THEREFORE that in consideration of the additional benefits provided for hereunder, the premises and covenants set forth herein, and other good and valuable consideration, the sufficiency of which is hereby acknowledged, the Company and Employee, intending to be legally bound, agree as follows:

I. DEFINITIONS

Whenever used in the Agreement the following terms shall be defined as follows:

(a) "Advisor" or "Advisors" shall mean with respect to Employee any person including lawyers, accountants, estate planners and others, with whom he may wish to review and discuss the matters set forth herein.

- (b) "Agreement" shall mean this Retirement Benefit Agreement which is entered into on the 14th day of March, 1995.
- (c) "At-Will" shall mean with respect to the period of Jackson's employment with Mylan, that the Company is under no obligation to continue to employ Jackson for any period of time, and can terminate his employment at any time without notice, subject to certain statutory and regulatory requirements; and that Employee is under no obligation to remain employed by the Company, and can terminate his employment with Mylan at any time, without notice.
 - (d) "Change of Control" shall mean:
- (1) The acquisition (other than from the Company) by any person, entity or "group", within the meaning of Section 13(d)(3) or 14(d)(2) of the Securities Exchange Act of 1934 (the "Exchange Act"), excluding, for this purpose, the Company or its subsidiaries, or any employee benefit plan of the Company or its subsidiaries which acquires beneficial ownership of voting securities of the Company (within the meaning of Rule 13d-3 promulgated under the Exchange Act), or legal ownership of 20% or more of either the then outstanding shares of common stock or the combined voting power of the Company's then outstanding voting securities entitled to vote generally in the election of directors; or
- (2) Individuals who, as of the date hereof, constitute the Board (as of the date hereof the "Incumbent Board") cease for any reason to constitute at least a majority of the Board, provided that any person becoming a director subsequent to the date hereof whose election, or nomination for election, by the Company's shareholders was approved by a vote of at least a majority of the directors then comprising the Incumbent Board (other than an election or nomination of an individual whose initial assumption of office is in connection with an actual or threatened election contest relating to the election of the Directors of the Company, as such terms are used in Rule 14a-11 of Regulation 14A promulgated under the Exchange Act) shall be, for purposes of this Agreement, considered as though such person were a member of the Incumbent Board: or
- (3) Approval by the shareholders of the Company of a reorganization, merger, consolidation, or other action with respect to which persons who were the shareholders of the Company immediately prior to such reorganization, merger or consolidation, or other action do not, immediately thereafter, own more than 50% of the combined voting power entitled to vote generally in the election of directors of the reorganized, merged or consolidated company's then outstanding voting securities, or of the sale of all or substantially all of the assets of the Company.
 - (e) "Contingency" shall mean Retirement or death.
- (f) "Mylan" or "Company" shall mean Mylan Laboratories Inc., its subsidiaries and affiliates.
- (g) "Net Present Value" ("NPV") shall mean the present value at any given time of the benefit to be paid, discounted at seven percent (7%) per annum.
- (h) "Party" or "Parties" shall mean the Company or Employee, or both the Company and Employee depending upon which term is required by the context in which it is used.
- (i) "Retire" or "Retirement" shall mean the day and date on which Jackson's employment with the Company is terminated by either Party for any reason other than death of the Employee.
- (j) "Successor" shall mean any person, partnership, limited partnership, joint-venture, corporation, trust or any other entity or organization who, subsequent to the Effective Date, comes into possession of or acquires, either directly or indirectly, all or substantially all of the Company's business, assets or voting stock, or the right to direct the business activities and practices of the Company. B

II. RESCISSION OF PRIOR AGREEMENT

The Salary Continuation Agreement entered into by the Parties on April 1, 1989 and any and all other agreements, express or implied, which may have been entered into by them prior to the execution of this Agreement, including by way of example and not of limitation, any agreement which addresses or is related to salary continuation, deferred compensation, employment, or similar matters (excluding agreements related to stock options) are hereby RESCINDED by the mutual consent of the Parties hereto upon execution of this Agreement.

THEREFORE THE PARTIES ACKNOWLEDGE THAT ANY RIGHTS AND OBLIGATIONS SET FORTH IN ANY SUCH AGREEMENT, WHETHER EXPRESS OR IMPLIED, ARE FOREVER WAIVED, NULL AND VOID, AND UNENFORCEABLE AT LAW OR IN EQUITY.

III. RETIREMENT

- 3.1 Upon his Retirement from the Company, and if he is eligible to receive payments as provided for elsewhere herein, Employee shall receive an annual retirement benefit equal to the amount set forth below.
- 3.2 Should Employee Retire after the Effective Date but on or before March 31, 1996 he shall receive thirty six thousand dollars (\$36,000.00) each year for ten (10) years.
- 3.3 Should Employee Retire after March 31, 1996 but on or before March 31, 1997 he shall receive seventy thousand dollars (\$70,000.00) each year for ten (10) years.
- 3.4 Should Employee Retire after March 31, 1997 but on or before March 31, 1998 he shall receive eighty thousand dollars (\$80,000.00) each year for ten (10) years.
- 3.5 Should Employee Retire after March 31, 1998 but on or before March 31, 1999 he shall receive ninety thousand dollars (\$90,000.00) each year for ten (10) years.
- 3.6 Should Employee Retire after March 31, 1999 he shall receive one hundred thousand dollars (\$100,000.00) each year for ten (10) years.

- 3.7 Should Employee become unable to perform the material and substantial duties of his position prior to March 31, 1999, he shall receive, pursuant to 4.1, one hundred thousand dollars (\$100,000.00) each year for ten (10) years in lieu of any benefit specified in Sections 3.2 through 3.6 hereof.
- 3.8 The Company shall pay the amount due hereunder in equal or substantially equal monthly installments. The first of any such payments shall be made on the first day of the month following the month in which Employee Retires, and each subsequent payment shall be made on the first day of each successive month until Mylan's obligations with respect to such payments have been satisfied.
- 3.9 However, upon the written request of the Employee, Mylan may pay to Employee the NPV of any amount, or of the balance of any such amount, to which he is entitled hereunder in a lump-sum payment. If the Company grants the request for a lump sum payment, said payment shall be paid within thirty (30) days of the date of Employee's request.

IV. CAPACITY TO PERFORM DUTIES

4.1 The certification of a licensed physician selected by the Company as to Employee's inability to perform the material and substantial duties of his position shall be conclusive with respect to his status regarding the application of 3.7 hereof.

V. DEATH BENEFIT

- 5.1 The Company shall maintain for Employee's benefit during his employment with the Company life insurance policies in the aggregate amount of one million two hundred fifty thousand dollars (\$1,250,000.00).
- 5.2 If the Employee's death occurs after Retirement, but before having received the entire benefit provided for under Article III hereof, the balance of the payments due thereunder shall be paid to Employee's beneficiary in a lump-sum payment equal to the NPV of the remaining payments.

VI. EFFECT OF CHANGE OF CONTROL

- 6.1 Upon a Change of Control Jackson shall receive, in lieu of the annual payments provided for under Article III, the NPV of One Hundred Thousand Dollars (\$100,000.00) per year for ten (10) years; provided Jackson is employed by the Company at or immediately prior to the Change of Control.
- 6.2 If a Change of Control occurs after his retirement, but before having received the entire benefit provided for under Article III hereof, the balance of the payments due thereunder shall be paid to Employee in a lump-sum payment equal to the NPV of the remaining payments.

VII. SUCCESSORSHIP

This Agreement in its entirety shall be binding upon and enforceable against the Company and its Successors.

VIII. NO DUPLICATION OF PAYMENTS

Notwithstanding anything to the contrary which may be set forth elsewhere herein, under no circumstances is Employee or his beneficiary entitled to take benefits under more than any one article included in this Agreement.

IX. EMPLOYEE CONDUCT WITH RESPECT TO COMPETITORS

- 9.1 Employee agrees that he will not, without the prior written consent of the Company, directly or indirectly, whether as an employee, officer, director, independent contractor, consultant, stockholder, partner or otherwise, engage in or assist others to engage in or have any interest in any business which competes with the Company in any geographic area in which the Company markets or has marketed its products during the year preceding termination of Jackson's employment for the greater of:
- (a) the period during which Employee receives monthly payments under this Agreement; or
 - (b) three (3) years following his receipt of a lump-sum payment hereunder.
- 9.2 Notwithstanding anything to the contrary set forth elsewhere herein, stock ownership in a competing business shall not be a breach of this Agreement, provided such stock is traded on a national exchange.
- 9.3 The Parties agree and acknowledge that the time, scope and geographic area and other provisions of this Agreement have been specifically negotiated by the Parties, and Employee specifically hereby agrees that such time, scope and geographic area and other provisions are reasonable under these circumstances. Employee further agrees that if, despite the express agreement of the Parties to this Agreement, a court should hold any portion of this Agreement unenforceable for any reason, the maximum restrictions of time, scope and geographic area reasonable under the circumstances, as determined by the court, will be substituted for the restrictions herein which such court may find to be unreasonable or unenforceable.
- 9.4 The Parties acknowledge that the breach of 10.1 will be such that the Company will not have an adequate remedy at law because the rights of the Company under this Agreement are of a specialized and unique character, and that immediate and irreparable damage will result to the Company if Employee breaches his obligations under 10.1. The Company may, in addition to any other remedies and damages available, seek an injunction in the courts of the State of West Virginia and the United States District Court for the Northern District of West Virginia to restrain any such breach. Employee represents and warrants that his expertise and capabilities are such that his obligations under 10.1 will not prevent him from earning a living.

CONSULTING SERVICES

- 10.1 During the five (5) year period beginning on the day following Employee's Retirement he shall, at the request of the Company, act in the capacity of a consultant for the Company, performing such services as may be consistent with those performed by him during his Employee's employment. These services may be designated by the President of the Company, or his authorized representative, and shall be reasonable in scope duration and ΓВ
- 10.2 The Company shall pay the Employee for such consulting services an hourly rate to be determined by the Parties at such time, but not less than one hundred fifty dollars (\$150.00) per hour, payable monthly.
- 10.3 In addition to the foregoing, the Company shall reimburse the Employee monthly for any and all out-of-pocket expenses incurred by the Employee directly for the benefit of the business of the Company.

XI. ELIGIBILITY FOR PAYMENT

- 11.1 Any and all payments due hereunder, may be denied if not already begun, or terminated if they have begun, if in the Company's sole judgment Employee is either not eligible for such payments, or once such payments have begun is found to be or found to have been ineligible.
- 11.2 Employee shall not be eligible for any payments hereunder if the Company, in its sole discretion, finds that during or subsequent to his employment with the Company he:
 - (a) breaches, or has breached any term, provision or obligation enumerated herein;
- (b) committed any act by commission or omission which materially and substantially adversely affects the Company's business or reputation; or
- (c) is convicted of any violation of the Federal Food, Drug and Cosmetic Act, or the violation of any other statute of material relevance to the Company's business
- 11.3 Should Employee be paid any benefits hereunder and thereafter be found ineligible, or to have been ineligible, he must return to the Company that portion of the benefit paid to him for the period of his ineligibility.

XII. RIGHT TO CONFER

- 12.1 Employee shall have the right, but not the obligation to:

 - (a) Confer with any Advisor of his choice prior to signing the Agreement; and(b) Provide his Advisors with a true and complete copy of the Agreement, and any other pertinent documents which may be of assistance to his Advisors.
- 12.2 Should Employee decline the right to confer with his Advisors prior to executing this Agreement he shall execute an acknowledgement of same which shall then be incorporated herein by reference. (See Exhibit A)

NO PROMISE OF CONTINUED EMPLOYMENT XTTT.

- 13.1 Employee acknowledges his employment with the Company is AT-WILL.
- 13.2 Nothing set forth herein shall constitute or be construed as a contract of employment except in so far as provided for under 13.1 hereof.

XIV. RESTRICTION OF ALIENABILITY

Benefits payable to the Employee or beneficiary shall not be subject to assignment, transfer attachment, execution, garnishment, sequestration, or any other seizure under any legal or equitable process, whether on account of the Employee's or beneficiary's act or by operation of the law.

XV. CONTRACT ADMINISTRATOR

The Director of Taxation and Accounting of the Company, or other officer of Mylan designated by the Executive Committee of the Company is hereby named the Contract Administrator for purposes of assuring compliance with the terms and conditions set forth herein.

XVI. MODIFICATION

This Agreement may not be changed, amended or otherwise modified other than by a written statement; Provided, such statement is signed by both Parties, expresses their intent to change the Agreement, and specifically describes such changes.

XVII. HEADINGS

Except when referenced in the body of this Agreement article headings are set forth herein for the purpose of convenience only. Such headings shall not be considered or otherwise referred to when any question or issue arises with respect to the application or interpretation of any term or condition set forth herein.

XVTTT. COUNTERPARTS

This Agreement may be executed in two or more counterparts, each of which is to be

considered an original, and taken together as one and the same document.

XIX. GOVERNING LAW

Any an all actions between the Parties regarding the interpretation or application of any term or provision set forth herein shall be governed by and interpreted in accordance with the substantive laws, and not the law of conflicts, of the State of West Virginia. The Company and Employee each do hereby respectively consent and agree that the courts of the State of West Virginia shall have jurisdiction, and venue shall properly lie with the courts of the State of West Virginia, with respect to any and all actions brought hereunder.

XX. SINGULAR OR PLURAL

The singular form of any noun or pronoun shall include the plural when the context in which such word is used is such that it is apparent the singular is intended to include the plural and vice versa.

XXI. ASSIGNMENT

The Agreement may not be assigned by either Party, without the written authorization of the other Party. A Successor shall not be considered an assignee for purposes of this Article.

XXII. ENTIRE AGREEMENT

The terms and conditions set forth herein contain the entire agreement between the Company and Employee, and supersede any and all prior agreements or understandings (whether express or implied) between the Parties with respect to the matters set forth herein.

XXIII. SURVIVAL

Articles I, II, IX, X, XI, XIX, and XXIII shall survive any expiration or termination of this Agreement.

XXIV. TERM

MYLAN LABORATORIES INC.

The term of this Agreement shall begin on the Effective Date and shall end on the date on which Mylan makes the last payment to which it is obligated hereunder.

XXV. AWARD OF ADDITIONAL RETIREMENT BENEFITS

25.1 The Company may increase the retirement benefit to which Employee is entitled under Article III. Any such increases shall be subject to the following conditions:

- (a) the maximum benefit to which Employee is entitled under Article III shall not exceed one hundred fifty thousand dollars (\$150,000.00) per year for ten (10) years;
 - (b) no increases shall be granted earlier than April 1, 2000;
 - (c) any increase shall be expensed for accounting purposes at the time it is granted;
- (d) the Company's obligation to pay the increased benefit to Employee, if and when such increases are awarded shall be subject to all the terms and conditions set forth herein; and
- (e) the inclusion herein of any increase in Employee's retirement benefit shall become effective upon Employee's receipt of a notice from the Company informing him of the additional benefit.

IN WITNESS of their agreement to the terms and conditions set forth herein the Company and Employee have caused the following signatures to be affixed hereto:

RODERICK P. JACKSON

BY:	BY:
TITLE:	DATE:
DATE:	

ACKNOWLEDGEMENT

I, Roderick P. Jackson, hereby acknowledge and understand that I have been given the opportunity to review and discuss the terms of the Agreement with an Advisor of my choice prior to signing the Agreement. I have decided that it is not necessary for me to discuss this matter with an Advisor, and therefore I decline the opportunity to do so.

No one has discouraged me or otherwise attempted to influence me with regard to my decision to decline the opportunity to discuss this matter with my Advisors. My refusal of this offer is entirely my own and is made without any reservation or conditions whatsoever.

WITNESSED BY:			
	DATE:	BY:	DATE:
	DATE:		

RETIREMENT BENEFIT AGREEMENT

This Retirement Benefit Agreement (the "Agreement") is entered into on this 14th day of March, 1995 (the "Effective Date") by and between:

Mylan Laboratories Inc., a Pennsylvania Corporation, with offices located at 781 Chestnut Ridge Road, Morgantown, WV 26505 (hereinafter referred to as "Mylan" or "Company").

and

Louis J. DeBone, an employee of Mylan who resides at 4 Poplar Woods Drive, Morgantown, WV 26505 (hereinafter referred to as "Employee" or "DeBone").

WHEREAS the Company and Employee, in recognition of Employee 's long and valuable contribution to the success of the Company, entered into a Salary Continuation Agreement on April 1, 1989; and

WHEREAS Employee continues to perform valuable services for the Company; and

WHEREAS in recognition of his continuing service to Mylan, the Company wishes to provide Employee with financial assistance with respect to certain Contingencies, in addition to that provided for in said April 1, 1989 Agreement; and

WHEREAS the Company and Employee wish to RESCIND, and to REPLACE said Salary Continuation Agreement with this Agreement;

WITNESSETH THEREFORE that in consideration of the additional benefits provided for hereunder, the premises and covenants set forth herein, and other good and valuable consideration, the sufficiency of which is hereby acknowledged, the Company and Employee, intending to be legally bound, agree as follows:

I. DEFINITIONS

Whenever used in the Agreement the following terms shall be defined as follows:

- (a) "Advisor" or "Advisors" shall mean with respect to Employee any person including lawyers, accountants, estate planners and others, with whom he may wish to review and discuss the matters set forth herein.
 - (b) "Agreement" shall mean this Retirement Benefit Agreement which is entered into

on the 14th day of March, 1995.

- (c) "At-Will" shall mean with respect to the period of DeBone's employment with Mylan, that the Company is under no obligation to continue to employ DeBone for any period of time, and can terminate his employment at any time without notice, subject to certain statutory and regulatory requirements; and that Employee is under no obligation to remain employed by the Company, and can terminate his employment with Mylan at any time, without notice.
 - (d) "Change of Control" shall mean:
- (1) The acquisition (other than from the Company) by any person, entity or "group", within the meaning of Section 13(d)(3) or 14(d)(2) of the Securities Exchange Act of 1934 (the "Exchange Act"), excluding, for this purpose, the Company or its subsidiaries, or any employee benefit plan of the Company or its subsidiaries which acquires beneficial ownership of voting securities of the Company (within the meaning of Rule 13d-3 promulgated under the Exchange Act), or legal ownership of 20% or more of either the then outstanding shares of common stock or the combined voting power of the Company's then outstanding voting securities entitled to vote generally in the election of directors; or
- (2) Individuals who, as of the date hereof, constitute the Board (as of the date hereof the "Incumbent Board") cease for any reason to constitute at least a majority of the Board, provided that any person becoming a director subsequent to the date hereof whose election, or nomination for election, by the Company's shareholders was approved by a vote of at least a majority of the directors then comprising the Incumbent Board (other than an election or nomination of an individual whose initial assumption of office is in connection with an actual or threatened election contest relating to the election of the Directors of the Company, as such terms are used in Rule 14a-11 of Regulation 14A promulgated under the Exchange Act) shall be, for purposes of this Agreement, considered as though such person were a member of the Incumbent Board; or
- (3) Approval by the shareholders of the Company of a reorganization, merger, consolidation, or other action with respect to which persons who were the shareholders of the Company immediately prior to such reorganization, merger or consolidation, or other action do not, immediately thereafter, own more than 50% of the combined voting power entitled to vote generally in the election of directors of the reorganized, merged or consolidated company's then outstanding voting securities, or of the sale of all or substantially all of the assets of the Company.
 - (e) "Contingency" shall mean Retirement or death.
- (f) "Mylan" or "Company" shall mean Mylan Laboratories Inc., its subsidiaries and affiliates.
- (g) "Net Present Value" ("NPV") shall mean the present value at any given time of the benefit to be paid, discounted at seven percent (7%) per annum.
- (h) "Party" or "Parties" shall mean the Company or Employee, or both the Company and Employee depending upon which term is required by the context in which it is used.
- (i) "Retire" or "Retirement" shall mean the day and date on which DeBone's employment with the Company is terminated by either Party for any reason other than death of the Employee.
- (j) "Successor" shall mean any person, partnership, limited partnership, joint-venture, corporation, trust or any other entity or organization who, subsequent to the Effective Date, comes into possession of or acquires, either directly or indirectly, all or substantially all of the Company's business, assets or voting stock, or the right to direct the business activities and practices of the Company.

II. RESCISSION OF PRIOR AGREEMENT

The Salary Continuation Agreement entered into by the Parties on April 1, 1989 and any and all other agreements, express or implied, which may have been entered into by them prior to the execution of this Agreement, including by way of example and not of limitation, any agreement which addresses or is related to salary continuation, deferred compensation, employment, or similar matters (excluding agreements related to stock options) are hereby RESCINDED by the mutual consent of the Parties hereto upon execution of this Agreement.

THEREFORE THE PARTIES ACKNOWLEDGE THAT ANY RIGHTS AND OBLIGATIONS SET FORTH IN ANY SUCH AGREEMENT, WHETHER EXPRESS OR IMPLIED, ARE FOREVER WAIVED, NULL AND VOID, AND UNENFORCEABLE AT LAW OR IN EQUITY.

III. RETIREMENT

- 3.1 Upon his Retirement from the Company, and if he is eligible to receive payments as provided for elsewhere herein, Employee shall receive an annual retirement benefit equal to the amount set forth below.
- 3.2 Should Employee Retire after the Effective Date but on or before March 31, 1996 he shall receive thirty six thousand dollars (\$36,000.00) each year for ten (10) years.
- 3.3 Should Employee Retire after March 31, 1996 but on or before March 31, 1997 he shall receive seventy thousand dollars (\$70,000.00) each year for ten (10) years.
- 3.4 Should Employee Retire after March 31, 1997 but on or before March 31, 1998 he shall receive eighty thousand dollars (\$80,000.00) each year for ten (10) years.
- 3.5 Should Employee Retire after March 31, 1998 but on or before March 31, 1999 he shall receive ninety thousand dollars (\$90,000.00) each year for ten (10) years.
- 3.6 Should Employee Retire after March 31, 1999 he shall receive one hundred thousand dollars (\$100,000.00) each year for ten (10) years.
 - 3.7 Should Employee become unable to perform the material and substantial duties of

his position prior to March 31, 1999, he shall receive, pursuant to 4.1, one hundred thousand dollars (\$100,000.00) each year for ten (10) years in lieu of any benefit specified in Sections 3.2 through 3.6 hereof.

- 3.8 The Company shall pay the amount due hereunder in equal or substantially equal monthly installments. The first of any such payments shall be made on the first day of the month following the month in which Employee Retires, and each subsequent payment shall be made on the first day of each successive month until Mylan's obligations with respect to such payments have been satisfied.
- 3.9 However, upon the written request of the Employee, Mylan may pay to Employee the NPV of any amount, or of the balance of any such amount, to which he is entitled hereunder in a lump-sum payment. If the Company grants the request for a lump sum payment, said payment shall be paid within thirty (30) days of the date of Employee's request.

IV. CAPACITY TO PERFORM DUTIES

4.1 The certification of a licensed physician selected by the Company as to Employee's inability to perform the material and substantial duties of his position shall be conclusive with respect to his status regarding the application of 3.7 hereof.

V. DEATH BENEFIT

- 5.1 The Company shall maintain for Employee's benefit during his employment with the Company life insurance policies in the aggregate amount of one million two hundred fifty thousand dollars (\$1,250,000.00).
- 5.2 If the Employee's death occurs after Retirement, but before having received the entire benefit provided for under Article III hereof, the balance of the payments due thereunder shall be paid to Employee's beneficiary in a lump-sum payment equal to the NPV of the remaining payments.

VI. EFFECT OF CHANGE OF CONTROL

- 6.1 Upon a Change of Control DeBone shall receive, in lieu of the annual payments provided for under Article III, the NPV of One Hundred Thousand Dollars (\$100,000.00) per year for ten (10) years; provided DeBone is employed by the Company at or immediately prior to the Change of Control.
- 6.2 If a Change of Control occurs after his retirement, but before having received the entire benefit provided for under Article III hereof, the balance of the payments due thereunder shall be paid to Employee in a lump-sum payment equal to the NPV of the remaining payments.

VII. SUCCESSORSHIP

This Agreement in its entirety shall be binding upon and enforceable against the Company and its Successors.

VIII. NO DUPLICATION OF PAYMENTS

Notwithstanding anything to the contrary which may be set forth elsewhere herein, under no circumstances is Employee or his beneficiary entitled to take benefits under more than any one article included in this Agreement.

IX. EMPLOYEE CONDUCT WITH RESPECT TO COMPETITORS

- 9.1 Employee agrees that he will not, without the prior written consent of the Company, directly or indirectly, whether as an employee, officer, director, independent contractor, consultant, stockholder, partner or otherwise, engage in or assist others to engage in or have any interest in any business which competes with the Company in any geographic area in which the Company markets or has marketed its products during the year preceding termination of DeBone's employment for the greater of:
- (a) the period during which Employee receives monthly payments under this Agreement; or
 - (b) three (3) years following his receipt of a lump-sum payment hereunder.
- 9.2 Notwithstanding anything to the contrary set forth elsewhere herein, stock ownership in a competing business shall not be a breach of this Agreement, provided such stock is traded on a national exchange.
- 9.3 The Parties agree and acknowledge that the time, scope and geographic area and other provisions of this Agreement have been specifically negotiated by the Parties, and Employee specifically hereby agrees that such time, scope and geographic area and other provisions are reasonable under these circumstances. Employee further agrees that if, despite the express agreement of the Parties to this Agreement, a court should hold any portion of this Agreement unenforceable for any reason, the maximum restrictions of time, scope and geographic area reasonable under the circumstances, as determined by the court, will be substituted for the restrictions herein which such court may find to be unreasonable or unenforceable.
- 9.4 The Parties acknowledge that the breach of 10.1 will be such that the Company will not have an adequate remedy at law because the rights of the Company under this Agreement are of a specialized and unique character, and that immediate and irreparable damage will result to the Company if Employee breaches his obligations under 10.1. The Company may, in addition to any other remedies and damages available, seek an injunction in the courts of the State of West Virginia and the United States District Court for the Northern District of West Virginia to restrain any such breach. Employee represents and warrants that his expertise and capabilities are such that his obligations under 10.1 will not prevent him from earning a living.

10.1 During the five (5) year period beginning on the day following Employee's Retirement he shall, at the request of the Company, act in the capacity of a consultant for the Company, performing such services as may be consistent with those performed by him during his Employee's employment. These services may be designated by the President of the Company, or his authorized representative, and shall be reasonable in scope duration and frequency.

[B

- 10.2 The Company shall pay the Employee for such consulting services an hourly rate to be determined by the Parties at such time, but not less than one hundred fifty dollars (\$150.00) per hour, payable monthly.
- 10.3 In addition to the foregoing, the Company shall reimburse the Employee monthly for any and all out-of-pocket expenses incurred by the Employee directly for the benefit of the business of the Company.

XI. ELIGIBILITY FOR PAYMENT

- 11.1 Any and all payments due hereunder, may be denied if not already begun, or terminated if they have begun, if in the Company's sole judgment Employee is either not eligible for such payments, or once such payments have begun is found to be or found to have been ineligible.
- 11.2 Employee shall not be eligible for any payments hereunder if the Company, in its sole discretion, finds that during or subsequent to his employment with the Company he:
 - (a) breaches, or has breached any term, provision or obligation enumerated herein;
- (b) committed any act by commission or omission which materially and substantially adversely affects the Company's business or reputation; or
- (c) is convicted of any violation of the Federal Food, Drug and Cosmetic Act, or the violation of any other statute of material relevance to the Company's business.
- 11.3 Should Employee be paid any benefits hereunder and thereafter be found ineligible, or to have been ineligible, he must return to the Company that portion of the benefit paid to him for the period of his ineligibility.

XII. RIGHT TO CONFER

- 12.1 Employee shall have the right, but not the obligation to:
 - (a) Confer with any Advisor of his choice prior to signing the Agreement; and
 - (b) Provide his Advisors with a true and complete copy of the Agreement, and any other pertinent documents which may be of assistance to his Advisors.
- 12.2 Should Employee decline the right to confer with his Advisors prior to executing this Agreement he shall execute an acknowledgement of same which shall then be incorporated herein by reference. (See Exhibit A)

XIII. NO PROMISE OF CONTINUED EMPLOYMENT

- 13.1 Employee acknowledges his employment with the Company is AT-WILL.
- 13.2 Nothing set forth herein shall constitute or be construed as a contract of employment except in so far as provided for under 13.1 hereof.

XIV. RESTRICTION OF ALIENABILITY

Benefits payable to the Employee or beneficiary shall not be subject to assignment, transfer, attachment, execution, garnishment, sequestration, or any other seizure under any legal or equitable process, whether on account of the Employee's or beneficiary's act or by operation of the law.

XV. CONTRACT ADMINISTRATOR

The Director of Taxation and Accounting of the Company, or other officer of Mylan designated by the Executive Committee of the Company is hereby named the Contract Administrator for purposes of assuring compliance with the terms and conditions set forth herein.

XVI. MODIFICATION

This Agreement may not be changed, amended or otherwise modified other than by a written statement; Provided, such statement is signed by both Parties, expresses their intent to change the Agreement, and specifically describes such changes.

XVII. HEADINGS

Except when referenced in the body of this Agreement article headings are set forth herein for the purpose of convenience only. Such headings shall not be considered or otherwise referred to when any question or issue arises with respect to the application or interpretation of any term or condition set forth herein.

XVIII. COUNTERPARTS

This Agreement may be executed in two or more counterparts, each of which is to be considered an original, and taken together as one and the same document.

XIX. GOVERNING LAW

Any an all actions between the Parties regarding the interpretation or application of any term or provision set forth herein shall be governed by and interpreted in accordance with the substantive laws, and not the law of conflicts, of the State of West Virginia. The Company and Employee each do hereby respectively consent and agree that the courts of the State of West Virginia shall have jurisdiction, and venue shall properly lie with the courts of the State of West Virginia, with respect to any and all actions brought hereunder.

XX. SINGULAR OR PLURAL

The singular form of any noun or pronoun shall include the plural when the context in which such word is used is such that it is apparent the singular is intended to include the plural and vice versa.

XXI. ASSIGNMENT

The Agreement may not be assigned by either Party, without the written authorization of the other Party. A Successor shall not be considered an assignee for purposes of this Article.

XXII. ENTIRE AGREEMENT

The terms and conditions set forth herein contain the entire agreement between the Company and Employee, and supersede any and all prior agreements or understandings (whether express or implied) between the Parties with respect to the matters set forth herein.

XXIII. SURVIVAL

Articles I, II, IX, X, XI, XIX, and XXIII shall survive any expiration or termination of this Agreement.

XXIV. TERM

The term of this Agreement shall begin on the Effective Date and shall end on the date on which Mylan makes the last payment to which it is obligated hereunder.

IN WITNESS of their agreement to the terms and conditions set forth herein the Company and Employee have caused the following signatures to be affixed hereto:

MYLAN LABORATORIES INC.	LOUIS J. DEBONE	
BY:	BY:	
TITLE:	DATE:	
DATE:		

EXHIBIT A

ACKNOWLEDGEMENT

I, Louis J. DeBone, hereby acknowledge and understand that I have been given the opportunity to review and discuss the terms of the Agreement with an Advisor of my choice prior to signing the Agreement. I have decided that it is not necessary for me to discuss this matter with an Advisor, and therefore I decline the opportunity to do so.

No one has discouraged me or otherwise attempted to influence me with regard to my decision to decline the opportunity to discuss this matter with my Advisors. My refusal of this offer is entirely my own and is made without any reservation or conditions whatsoever.

WITNESSED BY:			
	DATE:	BY:	DATE:
	DATE:		

- (d) Employment contract with Milan Puskar dated April 28, 1983, as amended to date, filed as Exhibit 10(e) to Form 10-K for fiscal year ended March 31, 1993 and incorporated herein by reference.
- (e) Split Dollar Life Insurance Arrangement with McKnight Irrevocable Trust filed as Exhibit 10(g) to Form 10-K for fiscal year ended March 31, 1994 and incorporated herein by reference.

- (f) 1992 Nonemployee Director Stock Option Plan filed as Exhibit 10(g) to Form 10-K for fiscal year ended March 31, 1993 and incorporated herein by reference.
- (g) "Service Benefit Agreement" with Laurence S. DeLynn, John C. Gaisford, M.D., Richard A. Graciano and Robert W. Smiley, Esq. each dated January 27, 1995 and filed herewith.

SERVICE BENEFIT AGREEMENT

This Service Benefit Agreement (the "Agreement") is entered into on this 27th day of January, 1995 (the "Effective Date") by and between:

Mylan Laboratories Inc., a Pennsylvania Corporation, with offices located at 781 Chestnut Ridge Road, Morgantown, WV 26505 (hereinafter referred to as "Mylan" or "Company").

and

Laurence S. DeLynn, a Director of Mylan who resides at 113 Maple Avenue, Morgantown, WV 26505 (hereinafter referred to as "Director or "DeLynn").

WHEREAS Director has and continues to perform valuable services for the Company; and

WHEREAS in recognition of Director's past and continuing service to Mylan, the Company wishes to provide him with financial assistance with respect to certain Contingencies; WITNESSETH THEREFORE that in consideration of the benefits to be provided hereunder, the premises and covenants set forth herein, and other good and valuable consideration, the sufficiency of which is hereby acknowledged, the Company and Director, intending to be legally bound, agree as follows:

I. DEFINITIONS

Whenever used in the Agreement the following terms shall be defined as follows:

(a) "Advisor" or "Advisors" shall mean with respect to Director any person including lawyers, accountants, estate planners and others, with whom he may wish to review and discuss the matters set forth herein.

- (b) "Agreement" shall mean this Service Benefit Agreement which is entered into on the 27th day of January, 1995.
 - (c) "Contingency" shall mean retirement or death.
- (d) "Mylan" or "Company" shall mean Mylan Laboratories Inc., its subsidiaries and affiliates.
- (e) "Net Present Value" ("NPV") shall mean the present value at any given time of the entire benefit to be paid, discounted at seven percent (7%) per annum.
- (f) "Party" or "Parties" shall mean the Company or Director, or both the Company and Director depending upon which term is required by the context in which it is used.
- (g) "Successor" shall mean any person, partnership, limited partnership, joint-venture, corporation, trust or any other entity or organization who, subsequent to the Effective Date, comes into possession of or acquires, either directly or indirectly, all or substantially all of the Company's business, assets or voting stock, or the right to direct the business activities and practices of the Company.
- (h) "Termination of Service" shall mean the day and date on which Director's service to the Company is terminated by either Party for any reason other than death of the Director.

II. TERMINATION OF SERVICE

- 2.1 Upon his Termination of Service to the Company, and if he is eligible to receive payments as provided for elsewhere herein, Director shall receive an annual benefit equal to the amount set forth below.
- 2.2 Should Director cease to serve as a director of Mylan after the Effective Date he shall receive eighteen thousand dollars (\$18,000.00) each year for ten (10) years.
- 2.3 The Company shall pay each of the ten (10) annual payments due hereunder in twelve (12) equal or substantially equal installments. The first of any such payments shall be made on the first day of the month following the month in which Director ceases to serve as a director of Mylan, and each subsequent payment shall be made on the first day of each successive month until Mylan's obligations with respect to such payments have been satisfied.
- until Mylan's obligations with respect to such payments have been satisfied.

 2.4 However, upon the written request of the Director, Mylan may pay to him the NPV of any amount, or of the balance of any such amount, to which he is entitled hereunder in a lump-sum payment. Said payment shall be paid within thirty (30) days of the date of Director's request for such payment.

III. DEATH BENEFIT

- 3.1 If the Director's death occurs prior to his Termination of Service his beneficiary shall receive, in a lump-sum payment, the NPV of the benefit to which Director was entitled under 2.2 hereof.
- 3.2 If Director's death occurs after Termination of Service his beneficiary shall receive, in a lump-sum payment, the NPV of the remaining benefits to which Director was entitled under 2.2 hereof at the time of his death.

IV. SUCCESSORSHIP

This Agreement in its entirety shall be binding upon and enforceable against the Company and its Successors.

V. NO DUPLICATION OF PAYMENTS

Notwithstanding anything to the contrary which may be set forth elsewhere herein, under no circumstances is Director or his beneficiary entitled to take benefits under more than any one article included in this Agreement.

VI. ELIGIBILITY FOR PAYMENT

- 6.1 Any and all payments due hereunder, may be denied if not already begun, or terminated if they have begun, if in the Company's sole judgment Director is either not eligible for such payments, or once such payments have begun is found to be or found to have been ineligible.
- 6.2 Director shall not be eligible for any payments hereunder if the Company, in its sole discretion, finds that during or subsequent to his service to the Company he:
 - (a) breaches, or has breached any term, provision or obligation enumerated herein;
- (b) committed any act by commission or omission which materially and substantially adversely affects the Company's business or reputation; or
- (c) is convicted of any violation of the Federal Food, Drug and Cosmetic Act, or the violation of any other statute relevant to the Company's business.

6.3 Should Director be paid any benefits hereunder and thereafter be found ineligible, or to have been ineligible, he must return to the Company that portion of the benefit paid to him for the period of his ineligibility.

RIGHT TO CONFER VII.

- 7.1 Director shall have the right, but not the obligation to:

 - (a) Confer with any Advisor of his choice prior to signing the Agreement; and(b) Provide his Advisors with a true and complete copy of the Agreement, and any other pertinent documents which may be of assistance to his Advisors.
- 7.2 Should Director decline the right to confer with his Advisors prior to executing this Agreement he shall execute an acknowledgement of same which shall then be incorporated herein by reference. (See Exhibit A)

CONTRACT ADMINISTRATOR VIII.

The Director of Taxation and Accounting of the Company, or other officer of Mylan designated by the Executive Committee of the Company is hereby named the Contract Administrator for purposes of assuring compliance with the terms and conditions set forth herein.

IX. MODIFICATION

This Agreement may not be changed, amended or otherwise modified other than by a written statement; Provided, such statement is signed by both Parties, expresses their intent to change the Agreement, and specifically describes such changes.

X. HEADINGS

Except when referenced in the body of this Agreement article headings are set forth herein for the purpose of convenience only. Such headings shall not be considered or otherwise referred to when any question or issue arises with respect to the application or interpretation of any term or condition set forth herein.

XI. COUNTERPARTS

This Agreement may be executed in two or more counterparts, each of which is to be considered an original, and taken together as one and the same document.

XII. GOVERNING LAW

Any an all actions between the Parties regarding the interpretation or application of any term or provision set forth herein shall be governed by and interpreted in accordance with the substantive laws, and not the law of conflicts, of the Commonwealth of Pennsylvania. The Company and Director each do hereby respectively consent and agree that the courts of the Commonwealth of Pennsylvania shall have jurisdiction, and venue shall properly lie with the courts of the Commonwealth of Pennsylvania, with respect to any and all actions brought bereunder.

XIII. SINGULAR OR PLURAL

The singular form of any noun or pronoun shall include the plural when the context in which such word is used is such that it is apparent the singular is intended to include the plural and vice versa.

XIV. ASSIGNMENT

The Agreement may not be assigned by either Party, without the written authorization of the other Party. A Successor shall not be considered an assignee for purposes of this Article.

XV. ENTIRE AGREEMENT

The terms and conditions set forth herein contain the entire agreement between the Company and Director, and supersede any and all prior agreements or understandings (whether express or implied) between the Parties with respect to the matters set forth herein.

XVI. SURVIVAL

Articles I, VI, XII and XVI shall survive any expiration or termination of this Agreement.

XVII. TERM

The term of this Agreement shall begin on the Effective Date and shall end on the date on which Mylan makes the last payment to which it is obligated hereunder.

IN WITNESS of their agreement to the terms and conditions set forth herein the Company and Director have caused the following signatures to be affixed hereto: $\frac{1}{2} \left(\frac{1}{2} \right) = \frac{1}{2} \left(\frac{1}{2} \right) \left(\frac$

MYLAN LABORATORIES INC.	LAURENCE S. DELYNN	
BY:	BY:	
TITLE:	DATE:	
DATE:		

ACKNOWLEDGEMENT

I, Laurence S. DeLynn, hereby acknowledge and understand that I have been given the opportunity to review and discuss the terms of the Agreement with an Advisor of my choice prior to signing the Agreement. I have decided that it is not necessary for me to discuss this matter with an Advisor, and therefore I decline the opportunity to do so.

No one has discouraged me or otherwise attempted to influence me with regard to my decision to decline the opportunity to discuss this matter with my Advisors. My refusal of this offer is entirely my own and is made without any reservation or conditions whatsoever.

WITNESSED BY:			
	DATE:	BY:	DATE:
	DATE:		

SERVICE BENEFIT AGREEMENT

This Service Benefit Agreement (the "Agreement") is entered into on this 27th day of January, 1995 (the "Effective Date") by and between:

Mylan Laboratories Inc., a Pennsylvania Corporation, with offices located at 781 Chestnut Ridge Road, Morgantown, WV 26505 (hereinafter referred to as "Mylan" or "Company").

and

John C. Gaisford, M.D., a Director of Mylan who resides at 910 Waldhiem Road West, Pittsburgh, PA 15215 (hereinafter referred to as "Director or "Gaisford").

WHEREAS Director has and continues to perform valuable services for the Company; and

WHEREAS in recognition of his past and continuing service to Mylan, the Company wishes to provide him with financial assistance with respect to certain Contingencies;

WITNESSETH THEREFORE that in consideration of the benefits to be provided hereunder, the premises and covenants set forth herein, and other good and valuable consideration, the sufficiency of which is hereby acknowledged, the Company and Director, intending to be legally bound, agree as follows:

I. DEFINITIONS

(a) "Advisor" or "Advisors" shall mean with respect to Director any person including lawyers, accountants, estate planners and others, with whom he may wish to review and discuss the matters set forth herein.

- (b) "Agreement" shall mean this Service Benefit Agreement which is entered into on the 27th day of January, 1995.
 - (c) "Contingency" shall mean retirement or death.
- (d) "Mylan" or "Company" shall mean Mylan Laboratories Inc., its subsidiaries and affiliates.
- (e) "Net Present Value" ("NPV") shall mean the present value at any given time of the entire benefit to be paid, discounted at seven percent (7%) per annum.
- (f) "Party" or "Parties" shall mean the Company or Director, or both the Company and Director depending upon which term is required by the context in which it is used.
- (g) "Successor" shall mean any person, partnership, limited partnership, joint-venture, corporation, trust or any other entity or organization who, subsequent to the Effective Date, comes into possession of or acquires, either directly or indirectly, all or substantially all of the Company's business, assets or voting stock, or the right to direct the business activities and practices of the Company.
- (h) "Termination of Service" shall mean the day and date on which Director's service to the Company is terminated by either Party for any reason other than death of the Director.

II. TERMINATION OF SERVICE

- 2.1 Upon his Termination of Service to the Company, and if he is eligible to receive payments as provided for elsewhere herein, Director shall receive an annual benefit equal to the amount set forth below.
- 2.2 Should Director cease to serve as a director of Mylan after the Effective Date he shall receive six thousand dollars (\$6,000.00) each year for ten (10) years.
- 2.3 The Company shall pay each of the ten (10) annual payments due hereunder in twelve (12) equal or substantially equal installments. The first of any such payments shall be made on the first day of the month following the month in which Director ceases to serve as a director of Mylan, and each subsequent payment shall be made on the first day of each successive month until Mylan's obligations with respect to such payments have been satisfied.
- until Mylan's obligations with respect to such payments have been satisfied.

 2.4 However, upon the written request of the Director, Mylan may pay to him the NPV of any amount, or of the balance of any such amount, to which he is entitled hereunder in a lump-sum payment. Said payment shall be paid within thirty (30) days of the date of Director's request for such payment.

III. AWARD OF ADDITIONAL RETIREMENT BENEFITS

- 3.1 The Company may increase the retirement benefit to which Director is entitled under Article II. Any such increases shall be subject to the following conditions:
- (a) the maximum benefit to which Director is entitled under Article II shall not exceed eighteen thousand dollars (\$18,000.00) per year for ten (10) years;
 - (b) no increases shall be granted earlier than April 1, 1996;
 - (c) any increase shall be expensed for accounting purposes at the time it is granted;
- (d) the Company's obligation to pay the increased benefit to Director, if and when such increases are awarded shall be subject to all the terms and conditions set forth herein; and
- (e) the inclusion herein of any increase in Director's retirement benefit shall become effective upon Director's receipt of a notice from the Company informing him of the additional benefit.

IV. DEATH BENEFIT

- 4.1 If the Director's death occurs prior to his Termination of Service his beneficiary shall receive, in a lump-sum payment, the NPV of the benefit to which Director was entitled under 2.2 hereof.
- 4.2 If Director's death occurs after Termination of Service his beneficiary shall receive, in a lump-sum payment, the NPV of the remaining benefits to which Director was entitled under 2.2 hereof at the time of his death.

V. SUCCESSORSHIP

This Agreement in its entirety shall be binding upon and enforceable against the Company and its Successors.

VI. NO DUPLICATION OF PAYMENTS

Notwithstanding anything to the contrary which may be set forth elsewhere herein, under no circumstances is Director or his beneficiary entitled to take benefits under more than any one article included in this Agreement.

VII. ELIGIBILITY FOR PAYMENT

- 7.1 Any and all payments due hereunder, may be denied if not already begun, or terminated if they have begun, if in the Company's sole judgment Director is either not eligible for such payments, or once such payments have begun is found to be or found to have been ineligible.
- 7.2 Director shall not be eligible for any payments hereunder if the Company, in its sole discretion, finds that during or subsequent to his service to the Company he:
 - (a) breaches, or has breached any term, provision or obligation enumerated herein;
- (b) committed any act by commission or omission which materially and substantially adversely affects the Company's business or reputation; or
- (c) is convicted of any violation of the Federal Food, Drug and Cosmetic Act, or the violation of any other statute relevant to the Company's business.
- 7.3 Should Director be paid any benefits hereunder and thereafter be found ineligible, or to have been ineligible, he must return to the Company that portion of the benefit paid to him for the period of his ineligibility.

VIII. RIGHT TO CONFER

- 8.1 Director shall have the right, but not the obligation to:
 - (a) Confer with any Advisor of his choice prior to signing the Agreement; and
 - (b) Provide his Advisors with a true and complete copy of the Agreement, and any other pertinent documents which may be of assistance to his Advisors.
- 8.2 Should Director decline the right to confer with his Advisors prior to executing this Agreement he shall execute an acknowledgement of same which shall then be incorporated herein by reference. (See Exhibit A)

IX. CONTRACT ADMINISTRATOR

The Director of Taxation and Accounting of the Company, or other officer of Mylan designated by the Executive Committee of the Company is hereby named the Contract Administrator for purposes of assuring compliance with the terms and conditions set forth herein.

X. MODIFICATION

This Agreement may not be changed, amended or otherwise modified other than by a written statement; Provided, such statement is signed by both Parties, expresses their intent to change the Agreement, and specifically describes such changes.

XI. HEADINGS

Except when referenced in the body of this Agreement article headings are set forth herein for the purpose of convenience only. Such headings shall not be considered or otherwise referred to when any question or issue arises with respect to the application or interpretation of any term or condition set forth herein.

XII. COUNTERPARTS

This Agreement may be executed in two or more counterparts, each of which is to be considered an original, and taken together as one and the same document.

XIII. GOVERNING LAW

Any an all actions between the Parties regarding the interpretation or application of any term or provision set forth herein shall be governed by and interpreted in accordance with the substantive laws, and not the law of conflicts, of the Commonwealth of Pennsylvania. The Company and Director each do hereby respectively consent and agree that the courts of the Commonwealth of Pennsylvania shall have jurisdiction, and venue shall properly lie with the courts of the Commonwealth of Pennsylvania, with respect to any and all actions brought hereunder.

XIV. SINGULAR OR PLURAL

The singular form of any noun or pronoun shall include the plural when the context in which such word is used is such that it is apparent the singular is intended to include the plural and vice versa.

XV. ASSIGNMENT

The Agreement may not be assigned by either Party, without the written authorization of the other Party. A Successor shall not be considered an assignee for purposes of this Article.

XVI. ENTIRE AGREEMENT

The terms and conditions set forth herein contain the entire agreement between the Company and Director, and supersede any and all prior agreements or understandings (whether express or implied) between the Parties with respect to the matters set forth herein.

XVII. SURVIVAL

Articles I, VII, XIII and XVII shall survive any expiration or termination of this Agreement.

XVIII. TERM

The term of this Agreement shall begin on the Effective Date and shall end on the date on which Mylan makes the last payment to which it is obligated hereunder.

IN WITNESS of their agreement to the terms and conditions set forth herein the Company and Director have caused the following signatures to be affixed hereto:

MYLAN LABORATORIES INC.	JOHN C. GAISFORD, M.D.
BY:	BY:
TITLE:	DATE:
DATE:	

EXHIBIT A

ACKNOWLEDGEMENT

I, John C. Gaisford, M.D., hereby acknowledge and understand that I have been given the opportunity to review and discuss the terms of the Agreement with an Advisor of my choice prior to signing the Agreement. I have decided that it is not necessary for me to discuss this matter with an Advisor, and therefore I decline the opportunity to do so.

No one has discouraged me or otherwise attempted to influence me with regard to my decision to decline the opportunity to discuss this matter with my Advisors. My refusal of this offer is entirely my own and is made without any reservation or conditions whatsoever.

WITNESSED BY:			
	DATE:	BY:	DATE:
	DATE:		

SERVICE BENEFIT AGREEMENT

This Service Benefit Agreement (the "Agreement") is entered into on this 27th day of January, 1995 (the "Effective Date") by and between:

Mylan Laboratories Inc., a Pennsylvania Corporation, with offices located at 781 Chestnut Ridge Road, Morgantown, WV 26505 (hereinafter referred to as "Mylan" or "Company").

and

Richard A. Graciano, a Director of Mylan who resides at 3302 Scathelock Road, Pittsburgh, PA 15235 (hereinafter referred to as "Director or "Graciano").

WHEREAS Director has and continues to perform valuable services for the Company; and

WHEREAS in recognition of Director's past and continuing service to Mylan, the Company wishes to provide him with financial assistance with respect to certain Contingencies; WITNESSETH THEREFORE that in consideration of the benefits to be provided hereunder, the premises and covenants set forth herein, and other good and valuable consideration, the sufficiency of which is hereby acknowledged, the Company and Director, intending to be legally bound, agree as follows:

I. DEFINITIONS

Whenever used in the Agreement the following terms shall be defined as follows:

(a) "Advisor" or "Advisors" shall mean with respect to Director any person including lawyers, accountants, estate planners and others, with whom he may wish to review and discuss the matters set forth herein.

- (b) "Agreement" shall mean this Service Benefit Agreement which is entered into on the 27th day of January, 1995.
 - (c) "Contingency" shall mean retirement or death.
- (d) "Mylan" or "Company" shall mean Mylan Laboratories Inc., its subsidiaries and affiliates.
- (e) "Net Present Value" ("NPV") shall mean the present value at any given time of the entire benefit to be paid, discounted at seven percent (7%) per annum.
- (f) "Party" or "Parties" shall mean the Company or Director, or both the Company and Director depending upon which term is required by the context in which it is used.
- (g) "Successor" shall mean any person, partnership, limited partnership, joint-venture, corporation, trust or any other entity or organization who, subsequent to the Effective Date, comes into possession of or acquires, either directly or indirectly, all or substantially all of the Company's business, assets or voting stock, or the right to direct the business activities and practices of the Company.
- (h) "Termination of Service" shall mean the day and date on which Director's service to the Company is terminated by either Party for any reason other than death of the Director.

II. TERMINATION OF SERVICE

- 2.1 Upon his Termination of Service to the Company, and if he is eligible to receive payments as provided for elsewhere herein, Director shall receive an annual benefit equal to the amount set forth below.
- 2.2 Should Director cease to serve as a director of Mylan after the Effective Date he shall receive eighteen thousand dollars (\$18,000.00) each year for ten (10) years.
- 2.3 The Company shall pay each of the ten (10) annual payments due hereunder in twelve (12) equal or substantially equal installments. The first of any such payments shall be made on the first day of the month following the month in which Director ceases to serve as a director of Mylan, and each subsequent payment shall be made on the first day of each successive month until Mylan's obligations with respect to such payments have been satisfied.
- until Mylan's obligations with respect to such payments have been satisfied.

 2.4 However, upon the written request of the Director, Mylan may pay to him the NPV
 of any amount, or of the balance of any such amount, to which he is entitled hereunder in a lump-sum payment.
 Said payment shall be paid within thirty (30) days of the date of Director's request
 for such payment.

III. DEATH BENEFIT

- 3.1 If the Director's death occurs prior to his Termination of Service his beneficiary shall receive, in a lump-sum payment, the NPV of the benefit to which Director was entitled under 2.2 hereof.
- 3.2 If Director's death occurs after Termination of Service his beneficiary shall receive, in a lump-sum payment, the NPV of the remaining benefits to which Director was entitled under 2.2 hereof at the time of his death.

IV. SUCCESSORSHIP

This Agreement in its entirety shall be binding upon and enforceable against the Company and its Successors.

V. NO DUPLICATION OF PAYMENTS

Notwithstanding anything to the contrary which may be set forth elsewhere herein, under no circumstances is Director or his beneficiary entitled to take benefits under more than any one article included in this Agreement.

VI. ELIGIBILITY FOR PAYMENT

- 6.1 Any and all payments due hereunder, may be denied if not already begun, or terminated if they have begun, if in the Company's sole judgment Director is either not eligible for such payments, or once such payments have begun is found to be or found to have been ineligible.
- 6.2 Director shall not be eligible for any payments hereunder if the Company, in its sole discretion, finds that during or subsequent to his service to the Company he:
 - (a) breaches, or has breached any term, provision or obligation enumerated herein;
- (b) committed any act by commission or omission which materially and substantially adversely affects the Company's business or reputation; or
- (c) is convicted of any violation of the Federal Food, Drug and Cosmetic Act, or the violation of any other statute relevant to the Company's business.

6.3 Should Director be paid any benefits hereunder and thereafter be found ineligible, or to have been ineligible, he must return to the Company that portion of the benefit paid to him for the period of his ineligibility.

VII. RIGHT TO CONFER

- 7.1 Director shall have the right, but not the obligation to:
 - (a) Confer with any Advisor of his choice prior to signing the Agreement; and
 - (b) Provide his Advisors with a true and complete copy of the Agreement, and any other pertinent documents which may be of assistance to his Advisors.
- 7.2 Should Director decline the right to confer with his Advisors prior to executing this Agreement he shall execute an acknowledgement of same which shall then be incorporated herein by reference. (See Exhibit A)

VIII. CONTRACT ADMINISTRATOR

The Director of Taxation and Accounting of the Company, or other officer of Mylan designated by the Executive Committee of the Company is hereby named the Contract Administrator for purposes of assuring compliance with the terms and conditions set forth herein.

IX. MODIFICATION

This Agreement may not be changed, amended or otherwise modified other than by a written statement; Provided, such statement is signed by both Parties, expresses their intent to change the Agreement, and specifically describes such changes.

X. HEADINGS

Except when referenced in the body of this Agreement article headings are set forth herein for the purpose of convenience only. Such headings shall not be considered or otherwise referred to when any question or issue arises with respect to the application or interpretation of any term or condition set forth herein.

XI. COUNTERPARTS

This Agreement may be executed in two or more counterparts, each of which is to be considered an original, and taken together as one and the same document.

XII. GOVERNING LAW

Any an all actions between the Parties regarding the interpretation or application of any term or provision set forth herein shall be governed by and interpreted in accordance with the substantive laws, and not the law of conflicts, of the Commonwealth of Pennsylvania. The Company and Director each do hereby respectively consent and agree that the courts of the Commonwealth of Pennsylvania shall have jurisdiction, and venue shall properly lie with the courts of the Commonwealth of Pennsylvania, with respect to any and all actions brought bereunder.

XIII. SINGULAR OR PLURAL

The singular form of any noun or pronoun shall include the plural when the context in which such word is used is such that it is apparent the singular is intended to include the plural and vice versa.

XIV. ASSIGNMENT

The Agreement may not be assigned by either Party, without the written authorization of the other Party. A Successor shall not be considered an assignee for purposes of this Article.

XV. ENTIRE AGREEMENT

The terms and conditions set forth herein contain the entire agreement between the Company and Director, and supersede any and all prior agreements or understandings (whether express or implied) between the Parties with respect to the matters set forth herein.

XVI. SURVIVAL

Articles I, VI, XII and XVI shall survive any expiration or termination of this Agreement.

XVII. TERM

The term of this Agreement shall begin on the Effective Date and shall end on the date on which Mylan makes the last payment to which it is obligated hereunder.

IN WITNESS of their agreement to the terms and conditions set forth herein the Company and Director have caused the following signatures to be affixed hereto: $\frac{1}{2} \left(\frac{1}{2} \right) = \frac{1}{2} \left(\frac{1}{2} \right) \left(\frac$

MYLAN LABORATORIES INC.	RICHARD A. GRACIANO	
BY:	BY:	
TITLE:	DATE:	
DATE:		

ACKNOWLEDGEMENT

I, Richard A. Graciano, hereby acknowledge and understand that I have been given the opportunity to review and discuss the terms of the Agreement with an Advisor of my choice prior to signing the Agreement. I have decided that it is not necessary for me to discuss this matter with an Advisor, and therefore I decline the opportunity to do so.

No one has discouraged me or otherwise attempted to influence me with regard to my decision to decline the opportunity to discuss this matter with my Advisors. My refusal of this offer is entirely my own and is made without any reservation or conditions whatsoever.

WITNESSED BY:			
	DATE:	BY:	DATE:
	DATE:		

SERVICE BENEFIT AGREEMENT

This Service Benefit Agreement (the "Agreement") is entered into on this 27th day of January, 1995 (the "Effective Date") by and between:

Mylan Laboratories Inc., a Pennsylvania Corporation, with offices located at 781 Chestnut Ridge Road, Morgantown, WV 26505 (hereinafter referred to as "Mylan" or "Company").

and

Robert W. Smiley, Esq., a Director of Mylan who resides at 526 Sangree Road, Pittsburgh, PA 15237 (hereinafter referred to as "Director or "Smiley").

WHEREAS Director has and continues to perform valuable services for the Company; and

WHEREAS in recognition of Director's past and continuing service to Mylan, the Company wishes to provide him with financial assistance with respect to certain Contingencies; WITNESSETH THEREFORE that in consideration of the benefits to be provided hereunder, the premises and covenants set forth herein, and other good and valuable consideration, the sufficiency of which is hereby acknowledged, the Company and Director, intending to be legally bound, agree as follows:

I. DEFINITIONS

Whenever used in the Agreement the following terms shall be defined as follows:

(a) "Advisor" or "Advisors" shall mean with respect to Director any person including lawyers, accountants, estate planners and others, with whom he may wish to review and discuss the matters set forth herein.

- (b) "Agreement" shall mean this Service Benefit Agreement which is entered into on the 27th day of January, 1995.
 - (c) "Contingency" shall mean retirement or death.
- (d) "Mylan" or "Company" shall mean Mylan Laboratories Inc., its subsidiaries and affiliates.
- (e) "Net Present Value" ("NPV") shall mean the present value at any given time of the entire benefit to be paid, discounted at seven percent (7%) per annum.
- (f) "Party" or "Parties" shall mean the Company or Director, or both the Company and Director depending upon which term is required by the context in which it is used.
- (g) "Successor" shall mean any person, partnership, limited partnership, joint-venture, corporation, trust or any other entity or organization who, subsequent to the Effective Date, comes into possession of or acquires, either directly or indirectly, all or substantially all of the Company's business, assets or voting stock, or the right to direct the business activities and practices of the Company.
- (h) "Termination of Service" shall mean the day and date on which Director's service to the Company is terminated by either Party for any reason other than death of the Director.

II. TERMINATION OF SERVICE

- 2.1 Upon his Termination of Service to the Company, and if he is eligible to receive payments as provided for elsewhere herein, Director shall receive an annual benefit equal to the amount set forth below.
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MYLAN LABORATORIES INC.	ROBERT W. SMILEY, ESQ.	
BY:	BY:	
TITLE:	DATE:	
DATE:		

ACKNOWLEDGEMENT

I, Robert W. Smiley, hereby acknowledge and understand that I have been given the opportunity to review and discuss the terms of the Agreement with an Advisor of my choice prior to signing the Agreement. I have decided that it is not necessary for me to discuss this matter with an Advisor, and therefore I decline the opportunity to do so.

No one has discouraged me or otherwise attempted to influence me with regard to my decision to decline the opportunity to discuss this matter with my Advisors. My refusal of this offer is entirely my own and is made without any reservation or conditions whatsoever.

WITNESSED BY:			
	DATE:	BY:	DATE:
	DATE:		

(13) Fiscal 1995 Annual Report to the Shareholders (only those portions which are incorporated in this Report by reference are being filed herewith).

Mylan Laboratories Inc. 1995 Annual Report

Mylan Laboratories Inc. 1030 Century Building 130 Seventh Street Pittsburgh, Pennsylvania 15222

Description of Business

Mylan Laboratories Inc. and its subsidiaries are engaged primarily in manufacturing a variety of pharmaceutical products in finished tablet, capsule and powder dosage forms.

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About The Cover
 This beautiful child is Natalie . . . and Natalie has Cystinosis! In a world of breakthrough chemical therapies, we often assume that the drugs we need will be there when we need them. Yet, for Natalie and roughly 400 other children worldwide who suffer from a rare genetic disorder known as Cystinosis, the availability of lifesaving medicines was, until recently, something that could never be taken for granted. In patients with Nephropathic Cystinosis, the protein called cystine accumulates in the body's tissues, forming crystalline deposits which disrupt cell functions with devastating results. Invariably, Cystinosis leads to a progressive decline in renal functions and . . . ultimately . . . to the development of end-stage renal failure by the close of the first decade of life. In the early 1970's, researchers discovered that drug therapy employing the chemical cystemmine could dramatically reduce or halt the accumulation of cystine in living cells. Yet, while limited quantities of cystemmine were available throughout the clinical research of pioneering physicians Dr. Jess Thoene of the University of Michigan, Dr.

Jerry Schneider of the University of California at San Diego and Dr. William Gahl of the National Institutes of Health . . . efforts by pharmaceutical companies to develop an approved form of cysteamine were unsuccessful. Cysteamine was subsequently designated an orphan drug as defined under the Orphan Drug Act of 1983. Then, in August of 1992, Mylan assumed responsibility for developing the drug. In less than a year, Mylan's research team succeeded in creating a capsule form of cysteamine which represented a significant improvement over the investigational liquid. In August of 1993, Mylan submitted an orphan drug application to the FDA, and was awarded a new drug approval in August of 1994. The new product, called Cystagon(TM), will be marketed worldwide through a number of specialized agents. Natalie and her sister, Alexandra, enjoy playing together, laughing together and sharing childhood secrets. Now they can look forward to growing up together, laughing together and sharing adulthood secrets. With the introduction of Cystagon(TM), Mylan is helping to transform the children of Cystinosis into "the children of hope"!

To Our Shareholders

As we look back over the past twelve months and review its challenges and accomplishments, we are proud to report that fiscal 1995 has been a great year!

Our record sales and earnings are the result of a family of dedicated employees going the extra step to make sure a quality product is ready to ship when our customers need it.

We had five approvals from the FDA this past fiscal year, increasing our product line to

79 different compounds covering 23 therapeutic categories.

23 ANDA's (generic drugs) are presently submitted to the food and drug administration for approval with over 40 more in various stages of product development. We have several innovator compounds in our pipeline including two IND's (Investigational New Drugs) in progress this year, with at least two more to be filed.

During this past year, we have announced more product alliances with Eli Lilly and Company as well as an alliance with Roche. These are for products that extend our line of generics. We have also signed licensing agreements for additional products for our Dow Hickam Wound Care Division, and we are continually looking at additional licensing agreements and strategic alliances to continue our success and growth.

Our year was a record in more than just revenues... our annual shareholders meeting in June was attended by approximately 1,139 investors! We were a bit overwhelmed but extremely pleased to meet and greet everyone. An increase of the Quarterly Dividend was announced at the meeting, raising the annual total to twenty cents per share from sixteen cents. Then in December, we announced a special one time dividend of ten cents per share in addition to the Quarterly five cents, so, the January 13th payment was for a total of fifteen cents per share. Mylan has distributed \$22,208,000 in cash dividends this year. Also, shareholders equity has grown from \$379,969,000 to \$482,728,000.

We are proud of the growth Mylan has experienced over this past year and thank each and every one of our employees for their hard work and dedication in helping us to accomplish

this record year and in maintaining our position of leadership in the industry.

But we do not rest on last years' laurels. As hard as we work at being the best, we work harder at planning for the future.

I want to personally assure you that the entire Mylan Team is dedicated to the continued growth and prosperity of this company!

Most sincerely,

Milan Puskar Chairman, C.E.O. and President

Company History

Mylan began in 1961 as a privately owned company founded by our Chairman, C.E.O. and President, Milan Puskar, and an associate in White Sulphur Springs, West Virginia. Initially the company did not manufacture product, but operated as a distributor' buying finished goods and reselling to pharmacies, doctors, and etc.

In 1963 Mylan relocated to Princeton, West Virginia and then in 1965 to its present location in Morgantown. The company began manufacturing vitamins in 1965, and in 1966 received approval to start manufacturing Penicillin G tablets. In July 1968 production was expanded to include Tetracycline after receiving FDA approval.

It was at this point that Mylan became recognized in the industry as a very competent manufacturer of quality products and in 1969 Parke-Davis was the first major drug company to purchase finished goods from Mylan. The company continued to expand its list of approved products with the addition of Erythromycin in 1971 and Ampicillin in 1973, and its list of major drug companies purchasing product under private label continued to increase.

But Mylan's growth was not limited to product line and customers, for on February 15, 1973 the first shares of stock were traded on the Over-the-Counter Market, and Mylan became a public' company.

Under the expert guidance of the present management team, which took over May 13, 1976, the company experienced unbelievable growth and soon became eligible to be traded on the National-Over-the-Counter (NASDAQ) Market. The marriage' of Mylan and NASDAQ was a phenomenal combination. The rapidly growing generic manufacturer rewarded its shareholders with six stock splits in the short time span from July, 1979 through February, 1985, gaining the title of "The Nation's Largest Independent Drug Manufacturer". With 24 million shares of stock trading in the marketplace, management felt continued growth could only happen through a broader and more liquid yet closely regulated market. On April 14, 1986 Mylan became a member of the Big Board - THE NEW YORK STOCK EXCHANGE - and its new symbol became MYL.

Once again management's decision proved to be a wise one and with the new stable growth enjoyed on NYSE shareholders were rewarded with the seventh stock split in eight years. On August 1, 1986 a three-for-two stock dividend was forwarded to all Mylan investors and the company's outstanding shares grew to 36 million.

In 1984 Mylan introduced its first proprietary product - MAXZIDE(R), an antihypertensive drug that competes successfully with Smith Kline Beecham's Dyazide(R), one of the most widely prescribed drugs in the United States. In May of 1988, after three years of clinical testing, Mylan received approval on half strength Maxzide(R), known as Maxzide(R)-25MG. This product further strengthens Mylan's position in the mild to moderate hypertensive market. Both of these Maxzide(R) products were licensed to Lederle Laboratories Division of American Cyanamid for distribution.

In 1989 a second manufacturing facility was opened in Caguas, Puerto Rico totaling approximately 60,000 square feet. In the fall of 1993 the size of the Caguas facility was doubled and shortly thereafter a second facility was purchased in the town of Cidra, Puerto Rico.

As the company continued to grow, management could foresee the need for a distribution center to be able to ship product promptly, and in June of 1988 opened a 24,000 square foot facility in Greensboro, North Carolina. That proved to be so successful that the company opened a second facility in Reno, Nevada in June of 1991.

These are two of the reasons Mylan is known for having the fastest shipping record throughout the pharmaceutical industry.

With the opening of the Greensboro facility in June of 1988, Mylan became the only independent drug manufacturer with its own distribution center for servicing wholesale and chain drug accounts, regardless of their size.

In November of 1988 Mylan announced the joint-venture purchase of Somerset Pharmaceuticals with another generic company. Somerset is a small research company who was working on a drug for the treatment of Parkinson's disease. The product, Eldepryl(R), was approved by the Food and Drug Administration in June of 1989 and was launched in mid-September. It is an extremely effective drug widely recognized by physicians and is adding nicely to the Mylan bottom line.

In May of 1989, Mylan's then Chairman and C.E.O. Roy McKnight testified before the House Oversight and Investigations Committee on behalf of the company regarding improprieties at the FDA uncovered by Mylan's investigative work. This prompted an all out investigation of the generic drug industry and exposed cheating, payoffs and fraud. Overall the industry was shaken, but one thing remains solid . . . the honesty and integrity of Mylan! For thirty-four years the company has had one creed - we either do it right or we don't do it at all. At the annual shareholders meeting in June of 1991 Mylan announced the signing of a

At the annual shareholders meeting in June of 1991 Mylan announced the signing of a definitive agreement and plan of merger with Dow B. Hickam of Sugar Land, Texas. They are an established high quality branded pharmaceutical company with a highly skilled, aggressive marketing force . . . A necessary element in Mylan's strategic plan to become a fully integrated pharmaceutical company. The merger was completed on October 30, 1991.

One June 25, 1992 at the annual shareholders meeting, Mylan announced still another stock split . . . its eighth in thirteen years. This two for one split became effective August 1, 1992. Another strategic step was taken in January of 1993 when Mylan announced that it had

Another strategic step was taken in January of 1993 when myian announced that it had reached a definitive agreement to acquire the assets of Bertek, Inc. headquartered in St. Albans, Vermont.

Bertek is an important manufacturer and innovator of state-of-the-art transdermal drug delivery systems. They also have substantial operations in laminating and label manufacturing. The acquisition was completed February 25, 1993.

Mylan's code of ethics has positioned it as a leader in the pharmaceutical industry with a record of solid growth in a very unpredictable market.

The other secret' to Mylan's success is its family' of employees whose dedication to

The other secret' to Mylan's success is its family' of employees whose dedication to their work and pride in the company have been the backbone of this remarkable story. From maintenance to management it has been a blend of ideas, hard work and mutual respect, and continues to be the key to Mylan's ongoing success and growth.

Mylan Laboratories Incorporated

Fiscal 1995 has been an exciting year for Mylan Laboratories and its family of companies. We have reached new financial highs . . . sales have increased to a record \$396,120,000 and our net earnings have climbed to a record \$120,869,000.

When most people think of Mylan, they automatically think "generics". That's only natural since Mylan Pharmaceuticals is the nation's largest manufacturer of generic prescription drugs. But in reality the Mylan formula for success is multifaceted. We are innovators in the development of generic and brand-name health care products as well as in manufacturing, marketing and distribution. We have achieved synergy through acquisitions, mergers and alliances with other health care companies reflecting a diversity of products.

For the last three decades, Mylan has built its business and its reputation on the manufacture of affordable, bio-equivalent substitutes for brand-name prescription drugs. Yet, while generics remain the vital focus of Mylan operations, they are only one facet of a new, more dynamic and integrated Mylan.

Mylan has implemented a series of strategic initiatives designed to broaden its presence in related health care businesses, in ways that would effectively leverage the company's core strengths . . . while establishing Mylan in a position of competitive leadership.

We have grown in order to better meet the emerging needs of health care markets and in the process, we have evolved from a small distributor . . . into a fully-integrated health care company that presently ranks among America's top 1,000 growth businesses.

In addition to our headquarters in Pittsburgh, Pennsylvania the Mylan family of companies consists of a 335,000 square foot manufacturing plant and office complex in Morgantown, West Virginia; two production plants in Caguas and Cidra, Puerto Rico; manufacturing, packaging and office facilities at both our Hickam wound care division in Sugar Land, Texas and our Bertek division in St. Albans and Swanton, Vermont, which produces transdermal patches as well as forms and labels.

Mylan's manufacturing operations are supported by a pair of state-of-the-art distribution terminals located in Greensboro, North Carolina and Sparks, Nevada. Strategic placement of these ultramodern terminals, combined with on-line order entry and processing capabilities, allows Mylan to ship orders anywhere in North America with turnaround time of as little as one day.

Research and development has always been and always will be vital to Mylan's success, providing a stream of new products which fuels our growth. Dr. John O'Donnell, Vice President of Research and Quality Control oversees an extremely aggressive corporate research program wherein all divisions are pursuing projects in their particular field of expertise.

wherein all divisions are pursuing projects in their particular field of expertise.

Combined, these capabilities comprise the Mylan family of health care services, helping Mylan to break away from its generics only' image and break through to a larger universe of health care customers and markets.

Today, Mylan's MYL symbol trades proudly on the New York Stock Exchange . . . a visible sign of the company's exceptional performance and growth.

visible sign of the company's exceptional performance and growth.

Forbes Magazine regards Mylan as one of the best small companies in America and ranks us number 88 out of 200 in their November 7, 1994 issue. Financial World chooses us as one of America's 50 best Mid-Cap companies in their June, 1994 listing and in their January 31, 1995 issue, we are named among their Market Cap 500 which ranks the largest U.S. Corporations in terms of market value as of December 30, 1994.

Executive Report Magazine has favored us with inclusion in two of their issues this year touting us as one of the top 10 Growth Companies as well as one of the top 10 in Revenue increases. We are most grateful for their recognition as well as the recognition by Fortune

Magazine in their March issue as one of the favored stocks on Wall Street.

Our shareholders are extremely important to the Mylan family, so we were particularly pleased when Better Investing' magazine, a publication of the National Association of Investors Corporation, named us as one of the most popular stocks in their listing of the top 100 for 1995.

We are also very proud of the honor bestowed upon us by the Chain Drug Store Industry . . they have voted Mylan as the "Supplier of the Decade" and rewarded us with the coveted REX award . . . truly a great commendation.

Cardinal Health and Smith Drug have each rewarded our year's efforts by voting us Vendor of the Year' and Amerisource has voted us their Manufacturing Partner of the Year'. Gold Circle has elected Mylan as an official member of the Gold Circle Club, and the NWDA has chosen us as the runner up for their Diana Award..the second year in a row we have been

recognized by this fine group.

Excellence is its own reward . . . but it is also very rewarding to be acknowledged by your

s . . . and we at Mylan are very grateful for the honors.
Mylan's evolution from a single location small generic manufacturing facility into a multi location fully integrated, financially strong industry leader has not been just luck' rather it is the reward of careful planning by a management team with the foresight to look into the future and plan for the changes that were bound to occur, and the wisdom and courage to take advantage of those opportunities without compromising their integrity . . . it takes loyal, dedicated employees who work hard and take pride in their accomplishments .

directors who are knowledgeable, honest and supportive.

Our board is comprised of intelligent, skilled and accomplished men who realize the gravity of their responsibility and are dedicated to the common good of this company, its employees and its shareholders. These principles have been the primary basis for all of their directives, and we thank them for their input and guidance.

It has been a year of change and challenge in the industry . . . there has been chaos . chaos creates opportunity. Mylan has met the challenge and taken advantage of the opportunities that were presented . . . we have enjoyed a period of growth . . . and we continue to build for the future as we move forward.

The entire Mylan team . . from maintenance to management . . . is dedicated to the integrity and hard work necessary to continue the growth and prosperity of this company. We look forward to the new year!

Captions p.9

With its blend of quality generic and innovator products, Mylan has grown into a fully integrated pharmaceutical firm, ranking among America's top 1,000 growth companies.

Roderick P. Jackson -Senior Vice President

Business Development: L to R Front: Heather Kirby, Dr. Tom Clark, Robert Lombardi; L to R Back: Bruce Battini, Rod Jackson, Lynn Cayton

Dr. Tom Clark - Medical Director Dr. John P. O'Donnell -Vice President

Research and Quality Control

Captions p11 Mylan's recognition in major publications continues

Captions p.12

Captions p.10

Mylan received numerous awards of excellence throughout the year including the coveted Rex Award for Supplier of the Decade.

Mylan Laboratories Corporate Staff

Seated L. to R. JoAnne Carrozza, Sally Stone, Patricia Sunseri - Vice President Investor and Public Relations; Standing L. to R. Brad Zukowski, Kristine Hartman King, Patti Ventura, Vicki Sidell

Captions p. 13

Board of Directors Mylan Laboratories Inc.: L to R Front: Milan Puskar, Dana G. Barnett; L to R Back: Richard A. Graciano, John C. Gaisford, M.D., C. B. Todd, Robert W. Smiley, Esq., Laurence S. DeLynn

Mylan Pharmaceuticals Incorporated

At Mylan Pharmaceuticals, the generic arm of the corporate family, the question has never been how to achieve excellence, but how to maintain it and move beyond it.

In our modern 335,000 square foot manufacturing facility in Morgantown, West Virginia, that commitment is both visible and growing.

We pride ourselves on having state-of-the-art equipment which we continually update . . . and all of the products manufactured here are made on the same equipment . . . they are developed with that in mind. That's what makes us so efficient and cost effective.

One of the things that keeps Mylan unique in its quality manufacturing is that we have three simple processes . . . basically, a blend mill process, a dry granulation process and a wet granulation process. We apply these simple processes to over 70 products which makes it very simple for product-to-product change-overs, simple for training new employees, very simple for record documentation and very simple for tracking quality control procedures.

Our plant is home to one of the most demanding quality assurance programs in the industry . . a program that combines highly evolved quality control processes with a battery of high-technology safeguards.

Mylan also maintains a center of excellence for research in Morgantown . . . a center of excellence in several different areas: analytical chemistry, dosage form development, pharmacokinetics, regulatory affairs and clinical investigation. The benefit is that, strategically, we can move more quickly and efficiently to bring products to market.

While new technology is a crucial part of Morgantown's long term strategy, Mylan is also aggressively developing the critical mass' of scientific and technical talent that will be required as the company moves ahead into a new century of manufacturing and product leadership.

Our on-site R & D staff includes more than 80 full-time degreed biologists, chemists and

pharmacologists . . . including 30 staff researchers of Master's Degree or Ph. D. caliber. At present we have 23 ANDA's submitted to the FDA for approval, and we have more than 40 other generic products in various stages of development.

We also have several innovator compounds in our pipeline including two IND's in progress with at least two more to be filed.

We have introduced eleven products to our line in Fiscal 1995: Gemfibrozil tablets, Bumetanide, Cefaclor, Cephalexin, Cimetidine, Flurbiprofen, Glipizide, Naproxen Sodium, Propoxyphene HCL, Propoxyphene Compound and Tolmetin Sodium 600 mg.

Six of these products are approvals obtained from the Food and Drug Administration, the other five are the result of our alliances with Eli Lilly and Roche.

Presently we are exploring more opportunities for co-promotion of other products in various therapeutic categories. These actions and relationships will continue to broaden our product portfolio and enhance our strong distribution network.

More than 5 billion tablets and capsules have been produced in this facility during fiscal 1995.

According to the February IMS National Prescription Audit, we have moved to second place among all pharmaceutical companies in the number of prescriptions dispensed . . . and we continue to gain market share.

Meclofenamate

Availability of raw material to meet production demands can be as important as the approval, especially when launching a new product. Mylan's purchasing group works very closely with product development to ensure that there will be an ample supply for development, and once the approval is received, for production. Many times, their expertise in acquiring the raw material has given Mylan the competitive edge.

and once the approval is received, for production. Many times, their expertise in acquiring the raw material has given Mylan the competitive edge.

In fact, it is that dedication by all of our employees in every department that has helped Mylan Pharmaceuticals become the leading independent generic drug manufacturer. It is that same employee strength and conviction that projects the brightest of futures for Mylan!

MYLAN PHARMACEUTICALS INC. GENERIC PRODUCT LINE	
Generic Name	Trade Name
Analgesics Indomethacin	Indocin(R)
** Propoxyphene HCL	Darvon(Ř)
** PropoxypheneCompound	Darvon 65(R)
Propoxyphene HCL &Acetaminophen Propoxyphene Napsylate &	Wygesic(R)
Darvocet-Acetaminophen	N-100(R)
Antiangina	Atenolol Tenormin(R)
Nadolol	Corgard(R)
Nitroglycerin Transdermal System (Patch)	Nitro Bid(R)
Verapamil HCL	Isoptin(R)
Antianxiety	
Alprazolam Diazepam	Xanax(R) Valium(R)
Lorazepam	Ativan(R)
Perphenazine &	
Amitriptyline HCL	Triavil(R)
Antibiotics	
Amoxicillin	Amoxil(R)
Ampicillin ** Cefaclor	Polycillin(R) Ceclor(R)
** Cephalexin	Keflex(R)
Doxycycline Hycate	Vibramycin(R)
Erythromycin Ethylsuccinate	E.E.S.(R)
Erythromycin	Erythrocin
Stearate Paniaillin V Patassium	Stearate(R)
Penicillin V Potassium Tetracycline HCL	V-cillin-K(R) Achromycin(R)
·	, ,
Antidepressant Amitriptyline	HCL Elavil(R)
Chlordiazepoxide &	not travit(K)
Amitriptyline	Limbitrol(R)
Doxepin HCL	Adapin(R) Sineguan(R)
Maprotiline HCL	Ludiomil(R)
Nortriptyline	Pamelor(R)
Antidiabetic	
Chlorpropamide	Diabinese(R)
* Glipizide Tolazamide	Glucotrol(R)
Tolbutamide	Tolinase(R) Orinase(R)
	,
Antidiarrheal Diphenoxylate HCL &	
Atropine Sulfate	Lomotil(R)
Loperamide HCL	<pre>Imodium(R)</pre>
Antigout	
Allopurinol	Zyloprim(R)
Antihistamine	
Cyproheptadine	Periactin(R)
	, ,
ANTIHYPERLIPIDEMIC * Gemfibrozil	Lopid(R)
Antihypertensive	
Amiloride HCL & Hydrochlorothiazide	Moduretic(R)
Clonidine HCL	Catapres(R)
Clonidine HCL & Chlorthalidone	Combipres(R)
Methyldopa	Aldomet(R)
Methyldopa &	. ,
Hydrochlorothiazide Metoprolol	Aldoril(R) Lopressor(R)
Prazosin HCL	Minipres(Ř)
Propranolol	Inderal(R)´
Propranolol HCL & Hydrochlorothiazide	Inderide(R)
, 3. 3011201 0111242240	111001 100(N)
Anti-Inflammatory	Nolfor(D)
Fenoprofen * Flurbiprofen	Nalfon(R) Ansaid(R)
Ibuprofen	Motrin(R)
	Rufen(R)

Rufen(R)

Meclomen(R)

Naprosyn(R) Naproxen Naproxen Sodium Anaprox(R) . Piroxicam Feldene(R) Sulindac Clinoril(R) Tolmetin Sodium Tolectin(R) Tolmetin Sodium Tolectin(R) 600 Antineoplastic Methotrexate Methotrexate(R) Rheumatrex(R) Antipsychotic Fluphenazine HCL Prolixin(R) Haloperidol Haldol(R) Thioridazine HCL Mellaril(R) Thiothixene Navane(R) Anxiolytic Clorazepate Dipotassium Tranxene(R) Beta Blocker Atenolol and Chlorthalidone Tenoretic(R) Pindolol Visken(R) Timolol Maleate Blocadren(R) Bronchial Dilator Albuterol Sulfate Proventil(R) Calcium ChannelBlocker Diltiazem HCL Cardizem(R) Diuretics Bumetanide Bumex(R) Chlorothiazide Diuril(R) Chlorthalidone Hygroton(R) Furosemide Lasix(R) Methyclothiazide Enduron(R) Reserpine & Chlorothiazide Diupres(R) Spironolactone Aldactone(R) Spironolactone & Hydrochlorothiazide Aldactazide(R) Hypnotic Agent Flurazepam Dalmane(R) Temazepam Restoril(R) **H2** Antagonist Cimetidine Tagamet(R) Muscle Relaxant Cyclobenzaprine HCL Flexeril(R) Uricosuric Probenecid Benemid(R) Captions p. 15 Left: Sonny Todd - President, Mylan Pharmaceuticals Center: High Speed Tableting Machine Captions, p. 16 Richard F. Stupar - Vice President, Purchasing

Bottom Right: Mylan Pharmaceuticals Plant, Morgantown, West Virginia Louis J.DeBone - Executive Vice President, Mylan Pharmaceuticals Morgantown, West Virginia

Mylan maintains a center of excellence for research in Morgantown

Mylan Incorporated

Mylan broke ground for its first manufacturing facility in Caguas, Puerto Rico on October 8, 1986, and less than one year later, that 60,000 square foot plant was completed and ready for production.

The success of this operation has been so outstanding that we have doubled the size of the

Caguas facility and purchased a second plant in the town of Cidra.

Puerto Rico is only about a third the size of the state of Vermont, yet it is home to one of the greatest concentrations of pharmaceutical manufacturing capabilities in the world.

Overall, the island boasts more than 70 individual manufacturing and processing plants,

representing nearly every major pharmaceutical company in the world.

The majority of manufacturers have established plants in Puerto Rico to take advantage of the island's exceptional work force and favorable tax structure. But Mylan is doing more than merely taking advantage of a good thing.

With its fully equipped facilities at Caguas and Cidra, Mylan has seized the initiative to push manufacturing technology to the limit, and create a new standard of excellence for quality and integrity in pharmaceutical production.

At the Caguas plant, our state-of-the-art manufacturing facilities produce a wide range of Mylan generics, as well as Somerset's proprietary anti-Parkinson's drug Eldepryl(R). Since it became operational in 1987, the Caguas facility has achieved an excellent record for both product

approval and regulatory compliance.

At its nearby Cidra facility, Mylan has created highly advanced capabilities for handling pharmaceutical products with special handling and manufacturing safety requirements. These capabilities permit Mylan to manufacture Cidra's principal product, Methotrexate.

The unique demands of these products - including expert application of specialized manufacturing equipment, strict adherence to rigid procedures, and constant use of sophisticated protective gear - create daily challenges which must be met and overcome.

But Cidra's dedicated staff has proven equal to the task.

With enormous energy and commitment, they are facing the immediate challenge of satisfying market demand for Mylan products today . . . and laying the foundation for Mylan's entry into significant new market segments tomorrow.

Carlos Machin, President and General Manager of Mylan's Puerto Rico operations commented: "The philosophy of Mylan is quality and integrity . . . the entire Mylan family is built around this, and Puerto Rico is no exception. Our employees are extremely proud of the quality we turn out, of our relationship with the community and with Puerto Rico. We have been a landmark in Puerto Rico, not only for the level of production . . . but for quality and integrity. We are extremely proud to be part of the Mylan family."

p.19 Captions

Left: Mylan Inc. Facility Caguas, Puerto Rico Lower Left: Mylan Inc. facility Cidra, Puerto Rico Top Right: Carlos Machin President and General Manager of Puerto Rico Operations Constant surveillance assures continued high quality Caguas, Puerto Rico

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p. 20 Captions

Francisco Torres - Production Manager, Cidra Plant

A wide range of Mylan's generic products are manufactured at the Caguas Plant

Facing the challenge of meeting market demand

p. 21 Captions

The Cidra plant features special handling and manufacturing capabilities

Dow Hickam Pharmaceuticals Incorporated

Dow Hickam Pharmaceuticals Inc. of Sugar Land, Texas was acquired by Mylan Laboratories Inc. in October 1991 and has become a vital part of a corporation well positioned to take advantage of the ever changing health care industry. This acquisition was another step by Mylan in their commitment to customers, shareholders, and employees to aggressively continue their evolution into a fully integrated pharmaceutical company. Hickam's history and dedication to quality and service make it a perfect fit with Mylan.

Dow Hickam specializes in the manufacturing and marketing of specialty pharmaceutical products and medical devices for use in hospitals, nursing homes, and home health care.

Dow Hickam's wound and burn care products are promoted and sold by its 80 person sales force. With this sizable presence, Hickam is able to reach the nation's 42 most populated states, covering 97 percent of the hospital and nursing home patients in the United States.

The Company recently increased its number of sales regions from seven to nine to support its expanding product line and growth into the managed care market place. The additional coverage gives Hickam the opportunity to grow the sales force when additional products are acquired or developed and will enable Hickam field managers to form stronger relationships with corporate accounts, national and regional buying groups, and managed care companies.

Hickam continues to evolve within the health care market place by acquiring and adding new items to its existing product line and by strengthening its position in the managed care market. During fiscal 1995 the Company successfully completed its launch of Flexzan(R), a highly conformable, polyurethane foam dressing, and in fiscal 1995 Hickam will be adding Flexfilm (TM),

an ultra-thin, highly flexible film dressing, to its line. Both of these products are the result of a license and supply agreement with Polymedica Industries, Inc. for the exclusive U.S. sales and marketing rights.

In December 1994 another exclusive licensing agreement was signed with Smith & Nephew Ltd. for the rights to manufacture, market, and sell a topical anesthetic. The product is currently in development at Hickam and will be manufactured in their Sugar Land facilities.

These new products complement the Company's current line and meet its strategic objective of acquiring unique and innovative products to meet its customers' needs.

In addition to expanding its product line, the Company continues to grow within the managed care market place. In September of 1994 the "Mylan Managed Health Care Program" was launched. The Hickam sales force promotes this program to its institutional pharmacy customers.

The Mylan family of companies is in a unique and quite envious position. Few companies have the strength of the Mylan line of quality generic products, coupled with the manpower of Hickam's national sales force and the Bertek line of pharmaceutical labels and forms.

This union of Mylan/ Hickam/Bertek is extremely effective in providing quality products and services from one source and truly makes Mylan a standout' in America's health care industry.

DOW HICKAM PHARMACEUTICALS INC. PRODUCT LINE GRANULEX(R)
A topical aerosol spray used for management of Stage I-IV

pressure ulcers.

PRODERM(R)
A non-prescription topical aerosol spray used for management of Stage I and II Pressure ulcers.

UNIFIBER(R)

A tasteless, non-gelling dietary fiber supplement.

QUICK!(R)

A topical cleanser for urine or fecal incontinence.

SORBSAN(R)

A highly absorbent calcium alginate wound dressing used in the management of exuding wounds.

FLEXZAN(R)

An ultra-thin, highly conformable, semi-occlusive polyurethane foam adhesive wound dressing.

FLEXFILM(TM)

An ultra-thin, highly flexible film wound dressing.

SULFAMYLON(R) CREAM

A topical antibacterial cream used in the treatment of burn wounds.

BIOBRANE(R)

An adherent biosynthetic temporary wound dressing used in the management of burn wound and donor sites.

p. 22 Captions

Left: Dow Hickam Headquarters, Sugar Land, Texas

Right: William W. Richardson President Dow Hickam Pharmaceuticals, Inc.

David W. Satter Executive Vice President and Chief Financial Officer, Dow Hickam Pharmaceuticals Inc.

Hickam Sales Managers

Hickam's Sales Regions

Bertek Incorporated

Bertek Incorporated, headquartered in St. Albans and Swanton Vermont, is a leading manufacturer in transdermal drug delivery systems technology, with coating, laminating, extrusion and labeling operations.

In February 1993, the company was acquired by Mylan Laboratories Inc. The acquisition provides Mylan with five patents for patch and drug delivery technology, wound care, and other related products to enhance the generic and branded divisions of Mylan. Bertek also provides Mylan with the third component of the "Mylan Managed Health Care Program," their innovative specialty and computer, generated forms and labels.

The health care industry is becoming increasingly complex every day. Health care reform, patient counseling, and patient outcomes all increase the amount of patient information to track and keep on file, placing greater demands on providers for accountability. Bertek technology helps simplify the complex with their pressure sensitive label systems. The development of the "Mylan Managed Health Care Program," enables Bertek to provide Managed Health Care with individualized specialty forms and labels combined with the complete line of Mylan generics and the national sales force of Dow Hickam Pharmaceuticals to better compete in todays ever changing health care arenas.

As a leader in the manufacturing of thin gauge plastic cards, Bertek's client base includes Blue Cross and Blue Shield, and numerous staff model HMO's. Bertek also offers a perfect fit as the complete supplier for computer generated labels and package inserts to Mylan's generic and proprietary pharmaceutical divisions.

Bertek has state of the art technology for producing extruding, coating and laminated pharmaceutical products into transdermal delivery of drugs to patients. These patented patches are also used for wound care. Transdermal osmotic absorption has become a significant advance in drug delivery systems, transdermal drug delivery increases patient compliance and reduces the risk of mismedicating, and in many products reduces side effects. Each day, new methods and applications for transdermal therapeutic systems evolve, and Bertek has helped pioneer that growth. By combining their extensive R&D and pharmacology capabilities with comprehensive GMP manufacturing, including clean room coating and laminating of the finished patch and all its components, Bertek is the complete source from the initial concept through final manufacturing.

Bertek currently provides components using internally developed technology for transdermal patches marketed by other companies under contract. Mylan is actively involved in research and development projects using Bertek technology to provide new products for marketing by its subsidiaries, including, but not limited to, developing generic patches, new chemical entities, line extensions of existing drugs and collaborating with Somerset Pharmaceuticals to develop an Eldepryl(R) patch.

The strategic fit of Bertek and Mylan will enable Mylan to be in the fore front of the ever changing health care market in 1995 and beyond.

Bertek INC. PRODUCT LINE

Transdermal Drug Delivery Systems

Bertek's Medical Products Division, a leader in Transdermal Drug Delivery Systems, represents a unique integration of R & D and the manufacturing know-how and full integration of production facilities to make raw materials and finished patches.

Wound Care Products

Bertek now stands as an established leader in the design, development and manufacture of both critical component materials and custom-designed products for use in wound management. The MEDIFILM(R) SERIES of extruded, controlled high moisture vapor permeable films offers a complete range of design flexibility for use in wound and I.V. site dressings, ulcer dressings, burn dressings, surgical drape and ostomy barrier applications.

Health Care Products and Materials

Surgical Incise Drape

Prolonged surgical procedures require the use of securely adhered incise drape films with a high degree of breathability to eliminate the possibility of perspiration-induced channeling and contamination of the wound site.

Films & Laminates for Ostomy Care & Skin Barriers

Bertek has developed a family of soft, conformable urethane and copolyester skin barrier films specifically for ostomy care.

GMP Converting and Labeling

As a printer of pharmaceutical labels, Bertek has established full capabilities for designing labels from typesetting to finished artwork by our in-house design team.

p. 27 captions

Left: Joseph J. Krivulka - President

Center: Pouching trials

Right: Label inspection

John Campbell - Vice President Pharmaceutical Manufacturing

Patch Die Cutting

p. 28 Captions

Inspecting package inserts

p. 29 Captions

Bertek is an innovator in the design and manufacture of forms and labels

Mylan's participation in Somerset began in 1988 when Mylan agreed to purchase fifty percent of the shares of Somerset. In June 1989 Somerset received FDA approval to market Eldepryl(R) as an Orphan Drug for the treatment of Parkinson's disease. More than 75,000 Parkinsonian patients are currently receiving the benefits of Eldepryl(R).

Sales of Eldepryl(R) have increased each year since market introduction to a level of \$125,000,000 in calendar year 1994. Net profits have increased each year and have generally tracked increases in sales particularly if adjustments are made for the significant increases in resources dedicated to research and development efforts.

In late 1991 Somerset realigned its business goals to include expanded efforts in research and development. As such, Somerset is pursuing line extensions of Eldepryl(R) as well as other related and unrelated new products to complement the sales of Eldepryl(R) which retains its market exclusively through June 5, 1996. To this end, Somerset has allocated resources equivalent to approximately ten percent of the net sales to fund these research efforts.

As a part of its expanded research and development efforts, Somerset located its Research and Development facility in Tampa in a 24,000 square feet laboratory building which is equipped with the latest in research and development equipment including: state-of-the-art analytical equipment such as an LC-MS-MS (liquid chromatograph, dual mass spectrophotometer), and high tech processing equipment such as a Glatt Fluidized Bed Granulator.

Personnel, equipment and facilities are in place at the Tampa location which allow Somerset to add orally administered product line extensions for Eldepryl(R), to pursue a patented Selegline Transdermal System (STS), and to develop an osteoporosis drug product for which Somerset has an exclusive license for the U.S. and Canadian markets. Additional development capabilities in Somerset's laboratory include the design, formulation and production of pilot quantities of liquids, tablets, capsules, ointments and creams. As such, Somerset has directed its development efforts toward expanding its product lines by pursuing drug delivery systems in the above named delivery systems.

Phase II clinical trials are in process assessing the Selegline Transdermal System (STS). Potential therapeutics uses of particular interest to Somerset for the STS include Alzheimers disease, clinical depression, attention deficit hyperactivity disorder and Parkinson disease.

An Investigational New Drug Application (IND) has been submitted for the osteoporotic agent and clinical trials designed to confirm the safety and efficacy of the osteoporotic agent are under way in Europe. The osteoporotic agent is specifically identified as an osteoclast inhibitor of the non-bisphosphonates class. This osteoporotic agent is currently marketed in Hungary, Italy, Japan, and Argentina by our licensor.

On the marketing front, Somerset is committed not only to providing pharmacological agents for the treatment of Parkinson's disease and other disease states, but to educating physicians and patients about how to get the most from their medications. As part of a comprehensive patient education program Somerset and its Eldepryl(R) co-marketing partner, Sandoz, launched two innovating programs in 1995.

In conjunction with the American Parkinson's Disease Association (APDA), Somerset is sponsoring the first ever national patient satellite conference to be held on June 14, 1995. Hosted in New York and featuring three internationally renowned movement disorder experts, the program will be beamed live to six other cities which will have local expert speakers to provide commentary and answer questions.

Also in 1995, an Eldepryl(R) VIP (Value in Persistency) program will be initiated. This program will provide educational videos and booklets on Parkinson's Disease and the various medications used to treat this disorder. The program also provides tips for daily living, exercise and coupons for valuable discounts.

Success with the above research, development, and marketing efforts will allow Somerset to continue to contribute to Mylan's future.

p. 31 Captions

Left: Somerset's R&D Facility Center: Somerset Corporate Office Right: Dana G. Barnett President Dr. Cheryl Blume - Vice President p. 32 Captions

Somerset's State-of-the-Art Analytical Equipment

Somerset Administrative Staff

P. 33 Captions

Mylan Manufactures Somerset's Eldepryl(R) in Puerto Rico Somerset's High Tech Processing Equipment

Financial Highlights

March 31 Net Sales	\$ 1995 396,120,000	\$ 1994 251,773,000
Net Earnings	\$ 120,869,000	\$ 73,067,000
Earnings Per Share	\$ 1.52	\$.93
Working Capital	\$ 275,032,000	\$ 191,647,000
Current Ratio	5.9 to 1	11.7 to 1
Total Assets	\$ 546,201,000	\$ 403,325,000
Shareholders' Equity	\$ 482,728,000	\$ 379,969,000
Book Value Per Share	\$ 6.09	\$ 4.81

70.6 73.1 120.9	1993 1994 1995
Shareholders' Equity (In millions) 167.5 203.5 296.0 380.0 482.7	(BAR GRAPH) Fiscal year 1991 1992 1993 1994 1995
Net Sales (In millions) 104.5 131.9 212.0 251.8 396.1	(BAR GRAPH) 1991 1992 1993 1994 1995

1992

Selected Financial Data

40.1

Year ended March 31 Net Sales	\$ 1995 396,120	\$ 1994 251,773	\$ 1993 211,964	\$ 1992 131,936	\$ 1991 104,524	\$ 1990 107,435	\$ 1989 99,558
Net Earnings	\$ 120,869	\$ 73,067	\$ 70,621	\$ 40,114	\$ 32,952	\$ 26,573	\$ 19,265
Earnings Per Share	\$ 1.52	\$.93	\$.92	\$.52	\$.43	\$. 35	\$. 25
Shares Used In Computation	79,309	78,949	77,101	76,484	76,368	76,226	76,140
At year end Working Capital	\$ 275,032	\$ 191,647	\$ 154,000	\$ 102,105	\$ 81,571	\$ 65,393	\$ 73,022
Total Assets	\$ 546,201	\$ 403,325	\$ 351,105	\$ 226,720	\$ 186,955	\$ 156,911	\$ 131,246
Long-Term Obligations	\$ 7,122	\$ 4,609	\$ 5,125	\$ 3,600	\$ 3,398	\$ 2,705	\$ 3,946
Shareholders' Equity	\$ 482,728	\$ 379,969	\$ 295,972	\$ 203,452	\$ 167,531	\$ 141,262	\$ 117,945
Book Value Per Share	\$ 6.09	\$ 4.81	\$ 3.84	\$ 2.66	\$ 2.19	\$ 1.85	\$ 1.55

Numbers in thousands except per share amounts.

From June of 1985 through June of 1990 the Company paid a semi-annual cash dividend of \$.05 per share per year. From June of 1990 through July of 1992 the Company had a quarterly dividend program totaling \$.10 per share per year. From October of 1992 to July of 1993 the Company had a quarterly dividend program totaling \$.12 per share per year. From October of 1993 to July of 1994 the Company had a quarterly dividend program totaling \$.16 per share per year. Since October of 1994 the Company has a quarterly dividend program totaling \$.20 per share per year. In addition the Company paid a special cash dividend of \$.10 per share in January of 1995.

The above Financial data gives retroactive effect to the October 30, 1991 business combination of Mylan Laboratories Inc. and Dow B. Hickam, Inc. and the two-for-one stock split effective August 1, 1992.

Management's Discussion and Analysis of Results Of Operations and Financial Position

Overview

By any measure fiscal 1995 was a rewarding year for the Company and our shareholders. Years of commitment to Mylan's standards of quality manufacturing, customer service and product development all contributed to the significant growth in net sales and net earnings. The Company's premier position in the generic industry provided a platform for unprecedented success in new product launches in fiscal 1995, including five products developed internally and five products licensed from brand name manufacturers. Additionally the strength of the Company's distribution capabilities was a key factor in the establishment of an arrangement with a subsidiary of Eli Lilly and Company relating to cefaclor. While the Company recognized revenue on a fee for services basis and not to the extent of actual sales to customers, this arrangement provides a significant enhancement to the Mylan product line.

The extraordinary growth in sales and gross profits has enabled the Company to accelerate and expand our investment into new product development and solidify our infrastructure. During fiscal 1995 the Company broke ground on a 150,000 square foot research and development facility. Scheduled for completion in late 1995, this state-of-the-art facility will not only enhance our product development effort, but will also provide for expansion of our manufacturing and administrative facilities. These steps will help the Company to meet the many challenges which lie ahead in the constantly changing pharmaceutical industry.

Mylan shareholders realized significant rewards from the Company's financial results in fiscal 1995. The Company declared dividends of just over \$23 million during the year, a 94% increase over the previous year and the market price for the Company's common stock appreciated significantly.

Results of Operations

Net Sales and Gross Margin

The following table outlines net sales, gross margins and the corresponding increase over the previous year.

(dollars in millions)

Year Ended	Net Sales	Gross Margin	Gross Margin as	
	March 31,	Dollars Change	Dollars Change	% of Net Sales
1995	\$ 396.1 57%	\$ 226.5 80%	57%	
1994	\$ 251.8 19%	\$ 126.1 3%	50%	
1993	\$ 212.0 61%	\$ 122.6 97%	58%	

The changes in net sales, gross margins and gross margin as a % of net sales are indicative of the highly competitive nature of the generic pharmaceutical industry and the Company's history of obtaining new product approvals. Generic products generally yield higher gross margins as a percent of sales in the short term period after introduction, and are subject to, sometimes severe, pricing deterioration as other competitors enter the market.

The Company added eleven products to our generic line in fiscal 1995, eight in 1994, four in 1993 and five in 1992. Of the eleven products added in fiscal 1995, three, cimetidine, glipizide and flurbiprofen had minimal competition and thus provided

significant sales and gross profit dollars. The Company experienced increased price competition on these products during the fourth quarter and expects continued competition throughout fiscal

In addition to new generic products the changes in sales and gross margins from 1993 to 1994 were affected by sales from Bertek (acquired in February 1993) which generally provides lower gross margin rates.

Due to the competitive nature of the generic pharmaceutical industry, net sales and gross margin percentages recognized in fiscal 1995 are not necessarily indicative of the results to be expected in future periods.

Research and Development

Research and Development expenses were \$30,533,000 compared to \$21,648,000 in 1994 and \$13,524,000 in 1993. These amounts represent approximately 8% of corresponding net sales in 1995, 9% in 1994 and 6% in 1993. Fiscal 1994 and 1995 expenditures include amounts for transdermal delivery system development in addition to increased expenditures for ongoing research and development of both innovative and generic products. The Company continues its commitment to new and increased product development.

Selling and Administrative

Selling and Administrative expenses were \$58,035,000 in fiscal 1995 compared to \$49,173,000 in fiscal 1994 and \$36,714,000 in fiscal 1993, which represents approximately 15%, 20% and 17% of corresponding net sales. The fiscal 1994 and 1995 amounts include approximately \$3,143,000 each year in amortization of intangible assets associated with the Bertek acquisition. In 1994, \$3,229,000 was expensed resulting from the death of Mr. McKnight the former Chairman and Chief Executive Officer of the Company. Other changes from 1994 to 1995 and from 1993 to 1994 are attributable in large part to compensation and related expenses, selling/marketing expenses associated with new products including sales commissions and legal and professional fees associated with the various court actions to which the Company has been involved.

Equity in Earnings of Somerset

Somerset's contribution to the Company's pretax earnings (in thousands) and net earnings per share are as follows:

		1995 Net		1994 Net		1993 Net
Quarter	Pretax	Earnings	Pretax	Earnings	Pretax	Earnings
Ended	Earnings	Per Share	Earnings	Per Share	Earnings	Per Share
6/30	\$ 5,348	\$.06	\$ 5,682	\$.06	\$ 4,309	\$.05
9/30	6,141	. 07	5,727	.07	5,101	.06
12/31	8,330	.09	6,841	.08	6,120	.07
3/31	5,587	.06	5,346	.06	5,606	.06
Fiscal Year	\$25, 406	\$.28	\$ 23,596	\$.27	\$ 21,136	\$.24

Under the Orphan Drug Act, Somerset has exclusivity relating to marketing the chemical compound Eldepryl(R) for use as a treatment for late stage Parkinson's disease through June of 1996. Under the Waxman Hatch Act, Somerset had exclusivity for all uses of the chemical compound through June of 1994. Somerset is actively involved in research projects regarding additional uses of this chemical compound, and other potential products.

Other Income

Other income for the year ended March 31, 1995 was \$7,958,000 compared to \$8,148,000 in 1994 and \$3,879,000 in 1993. The 1994 amount includes \$3,375,000 in legal settlements. Other changes are indicative of market fluctuations effecting the yields on investments, and changes in assets available for investment.

Income Taxes

The effective tax rates for 1995, 1994 and 1993 were 29%, 16% and 27% respectively. The 1994 effective tax rate was reduced by 5% as a result of recording the cumulative effects of changes in financial reporting requirements and changes in the federal tax code.

The Company recognized tax credits which reduced the effective tax rates by approximately 5% in 1995, 8% in 1994 and 2% in 1993. The tax credits result principally from operations in Puerto Rico and also from credits for increasing research and development activities. Changes in the Federal Tax Code enacted in 1993 reduced tax credits otherwise available for operating in Puerto Rico by 40% in fiscal 1995 with additional 5% reductions in the next four fiscal years. In addition recent tax rulings may reduce the amount of credit otherwise available to the Company for research and development activities.

Liquidity and Capital Resources

The Company's balance sheet remains strong with total assets increasing by 35% to \$546,201,000 at March 31, 1995. Working capital of \$275,032,000 represents 57% of net worth at March 31, 1995 versus \$191,647,000 or 50% of net worth at March 31, 1994. The ratio of current assets to current liabilities was 5.9 to 1 at March 31, 1995 versus 11.7 to 1 at March 31, 1994.

Cash flows from operating activities increased significantly from fiscal 1994 to 1995 due principally to the higher level of net earnings and also to the timing of collections of accounts receivable and payment of income taxes.

The Company invested \$17,485,000 in 1995 and \$20,164,000 in 1994 for facilities expansion, including relocation of our Greensboro distribution center, and expansion and renovation of facilities in Puerto Rico and Vermont. During fiscal 1994, the Company also acquired aircraft which were previously leased on

a flight by flight basis. All capital expenditures during fiscal years 1995 and 1994 were made with the general funds of the Company without incurring bank financing. As of March 31, 1995 the Company has commitments for future capital expenditures of approximately \$10,828,000, including the construction of a 150,000 square foot research and development facility in Morgantown, WV.

In addition to investing in property, plant and equipment, the Company has invested considerable amounts over the past two years into marketable securities, principally tax exempt instruments.

Increases in intangible and other assets includes equity investments and payments for long-term licenses to entities with which the Company is jointly developing new products. Payments of long-term obligations in 1994 represents a final settlement with the estate of Roy McKnight in connection with Mr. McKnight's salary continuation agreement. Prior year payments were for obligations assumed in the business combination of Bertek in 1993. The Company paid cash dividends of \$.28 per share in 1995 totaling \$22,208,0000 compared to \$.14 per share in 1994 which totaled \$11,026,000.

Consolidated Balance Sheets

March 31 Assets	1995		1994
Current Assets Cash and cash equivalents Marketable securities Accounts receivable - net Inventories Prepaid income taxes Deferred income tax benefit	\$ 127,280,000 52,575,000 58,343,000 78,205,000	\$	75,526,000 12,925,000 55,430,000 57,996,000 1,265,000 2,082,000
Other current assets Total Current Assets	4,435,000 331,383,000		4,349,000 209,573,000
Property, Plant and Equipment-net of accumulated depreciation Deferred income tax benefit, non-current Marketable Securities, Non-Current Intangible Assets-Net of accumulated amortization Other Assets Investment In and Advances to Somerset	92,299,000 1,043,000 21,958,000 28,518,000 48,945,000 22,055,000		82,514,000 - 12,125,000 33,228,000 48,122,000 17,763,000
Total Assets See notes to consolidated Financial statements.	\$ 546,201,000	;	\$ 403,325,000

Consolidated Balance Sheets

March 31 Liabilities and Shareholders' Equity	1995	1995
Current Liabilities Trade accounts payable Income taxes payable Other current liabilities Cash dividend payable Total Current Liabilities	24,01 17,89 3,97	\$6,000 \$ 6,699,000 .9,000 - .0,000 8,056,000 .6,000 3,171,000 .1,000 17,926,000
Long-Term Obligations Deferred Income Tax Liability	7,12	22,000 4,609,000 - 821,000
Shareholders' Equity Preferred stock, par value \$.50 per share, authorized 5,000,000 shares,		-
and 79,697,295 shares at March 31, 1994 Additional paid-in capital Retained earnings Unrealized gain on investments	57, 57 386, 21	4,000 -
Less treasury stock at cost - 476,523 shares at March 31, 1995 and 495,864 shares at March 31, 1994 Net Worth	,	21,000 2,509,000
Total Liabilities and Shareholders' Equity	\$ 546,20	1,000 \$ 403,325,000

Consolidated Statements of Earnings

Year ended March 31 Net Sales	\$ 1995 396,120,000	\$ 1994 251,773,000	\$ 1993 211,964,000
Cost and Expenses Cost of Sales Research and Development Selling and Administrative	169,590,000 30,533,000 58,035,000 258,158,000	125,631,000 21,648,000 49,173,000 196,452,000	89,400,000 13,524,000 36,714,000 139,638,000
Equity in Earnings of Somerset Other Income Earnings Before income Taxes Income Taxes Net Earnings	\$ 25,406,000 7,958,000 171,326,000 50,457,000 120,869,000	\$ 23,596,000 8,148,000 87,065,000 13,998,000 73,067,000	\$ 21,136,000 3,879,000 97,341,000 26,720,000 70,621,000
Earnings Per Share	\$ 1.52	\$.93	\$.92
Weighted Average Common Shares See notes to consolidated financial statements.	79,309,000	78,949,000	77,101,000

	Common Stock Shares	Common Stock Amount	ı	Additional Paid-In Capital	Retained Earnings	Unrealized Gain n Investments
April 1, 1992 Stock options exercised	38,631,373 713,857	\$ 19,317,000 357,000	\$	7,699,000 \$ 12,732,000	176,789,000	\$ -
Cash dividend \$.115 per share	-	-		12,732,000	(8,902,000)	-
Stock split (2 for 1)	38,654,343	19,327,000		(7,958,000)	(11,369,000)	-
Bertek acquisition	615,880	308,000		17,393,000	-	-
Net earnings	-	-		-	70,621,000	-
March 31, 1993	78,615,453	\$ 39,309,000	\$	29,866,000 \$	227,139,000	\$ -
Stock options exercised	347,747	173,000		4,447,000	-	-
Cash dividend \$.15 per share	-	-		-	(11,849,000)	-
Bertek acquisition	734,095	367,000		19,959,000	-	-
Net earnings	-	-		-	73,067,000	-
March 31, 1994	79,697,295	\$ 39,849,000	\$	54,272,000	\$288,357,000	\$ _
Stock options exercised	274,953	137,000		3,305,000	-	-
Cash dividend \$.29 per share	=	- '		· -	(23,014,000)	-
Net earnings	-	-		-	120,869,000	-
Change in unrealized gain on investments	-	-		-	-	1,374,000
March 31, 1995 See notes to consolidated Financial statem	79,972,248 ents.	\$ 39,986,000	\$	57,577,000	\$386,212,000	\$ 1,374,000

Consolidated Statements of Cash Flows

Year ended March 31	1995	1994	1993
Cash flows from operating activities			
Net earnings	\$ 120,869,000	\$ 73,067,000	\$ 70,621,000
Adjustments to reconcile net earnings to net cas	h provided from oper	ating activities:	
Depreciation and amortization	12,700,000	11,154,000	5,089,000
Deferred income tax benefit	(10,427,000)	(656,000)	(888,000)
Equity in earnings of Somerset	(25,406,000)	(23,596,000)	(21, 136, 000)
Cash received from Somerset	21,114,000	20,676,000	19,966,000
Other noncash expenses	13,252,000	4,192,000	3,758,000
Changes in operating assets and liabilities:			
Accounts receivable	(14,240,000)	(23,485,000)	(9,073,000)
Inventories	(19,590,000)	(12,002,000)	(9,825,000)
Trade accounts payable	3,410,000	207,000	1,911,000
Income taxes payable	25,060,000	(11,111,000)	6,263,000
Other operating assets and liabilities	9,789,000	(2,813,000)	1,651,000
Net cash provided from operating activities	136,531,000	35,633,000	68,337,000
Cash Flows from Investing Activities			
Additions to property, plant and equipment	(17,485,000)	(20,164,000)	(12,294,000)
Increase in intangible and other assets	(8,238,000)	(15, 147, 000)	(10,833,000)
Purchase of marketable securities	(58,491,000)	(12,925,000)	-
Proceeds from sales and maturities	(11, 1, 111)	(, = = , = = ,	
of marketable securities	25,482,000	4,800,000	-
Acquisition-net of cash acquired	(6,432,000)	-	-
Net cash used in investing activities	(65, 164, 000)	(43,436,000)	(23, 127, 000)
See notes to consolidated Financial statements.	(11, 11, 11)	(-,,,	(=, ==: , === ,

Consolidated Statements of Cash Flows

Year ended March 31	1995	1994	1993
Cash flows from Financing activities			
Payments on long-term obligations	\$ (451,000) \$	(4,320,000)	\$ (8,373,000)
Cash dividend paid	(22,208,000)	(11,026,000)	(8,476,000)
Payments on acquisition obligation		(977,000)	
Proceeds from exercise of stock options	3,046,000	1,406,000	9,561,000
Net cash used in financing activities	(19,613,000)	(14,917,000)	(7,288,000)
Net Increase (Decrease) in Cash and Cash Equival	ents 51,754,000	(22,720,000)	37,922,000
Cash and Cash Equivalents-Beginning of Year	75,526,000	98,246,000	60,324,000
Cash and Cash Equivalents-End of Year	\$ 127,280,000 \$	75,526,000	\$ 98,246,000

For purposes of presentation in the statements of cash flows, cash, overnight deposits, certificates of deposit, money market funds and marketable securities with original maturities of less than three months have been classified as cash and cash equivalents. The carrying value of these items

approximates fair value. Cash payments for interest were \$25,000 in 1995, \$30,000 in 1994 and \$64,000 in 1993. Cash payments for income taxes were \$35,822,000 in 1995, \$27,055,000 in 1994 and \$21,345,000 in 1993.

During fiscal 1993 the Company acquired substantially all of the assets of Bertek, Inc. (Bertek) (see note B) for approximately \$39,112,000 and assumed liabilities of approximately \$10,090,000. During the years ended March 31, 1994 and 1993 \$20,326,000 and \$17,701,000 of the purchase price was satisfied through the issuance of the Company's common stock. At the closing of this transaction, the Company repaid with cash approximately \$8,293,000 in long-term debt obligations assumed.

During fiscal 1993 the Company declared a 2 for 1 stock split effected in the form of a stock dividend (see note M).

Certain stock option transactions result in a reduction of income taxes payable and a corresponding increase in additional paid-in capital. The amounts for the years ended March 31, 1995, 1994 and 1993 were \$396,000, \$1,040,000 and \$3,528,000 respectively.

During fiscal 1995 and 1994 the Company received and recorded into treasury stock 659 and 75,658 shares of common stock valued at \$14,000 and \$ 2,174,000 respectively in consideration for the exercise of stock options.

A. Summary of Significant Accounting Policies

1. Principles of Consolidation

The consolidated financial statements include the accounts of Mylan Laboratories Inc. (the Company) and its wholly-owned subsidiaries. All intercompany accounts and transactions have been eliminated in consolidation. The Company's principal line of business is the manufacturing and distribution of pharmaceutical products. The Company had sales to one customer which represented 12% of net sales in 1993.

2. Marketable Securities

Effective April 1, 1994, the Company adopted Statement of Financial Accounting Standards No. 115 (SFAS No. 115), "Accounting for Certain Investments in Debt and Equity Securities."

this Statement, the Company's investments classified as "available for sale" are recorded at current market value with offsetting adjustments to shareholders' equity, net of income taxes. At March 31, 1994, investments were stated at the lower of aggregate cost or market. The adoption of SFAS No. 115 did not have a material impact on the financial position of the Company.

3. Accounts Receivable and Revenue Recognition

The Company recognizes revenue from product sales upon shipment to customers. Provisions for discounts, rebates, returns and other adjustments are provided for in the same period as the related sales are recorded. Accounts receivable are presented net of such provisions which amounted to \$14,777,000 at March 31, 1995 and \$3,449,000 at March 31, 1994.

4. Inventories

Inventories are stated at the lower of cost (principally, first-in, first-out) or market.

5. Property, Plant and Equipment

Property, plant and equipment are stated at cost. Depreciation is provided in amounts sufficient to relate cost of depreciable assets to operations over the estimated service lives, principally on a straight-line basis.

6. Research and Development

Research and development expenses are charged to operations as incurred.

7. Income Taxes

Deferred income taxes reflect the tax consequences on future years of events that have already been recognized by the Company in the financial statements or tax returns. Prior to April 1, 1993 deferred income taxes were provided for the difference between income and expense recognition for financial reporting purposes and income tax purposes.

8. Earnings Per Share

Earnings per share of common stock are based on the weighted average number of shares outstanding during each year. The effect on earnings per share, resulting from the assumed exercise of outstanding stock options, is not material.

9. Reclassification

Certain prior year amounts have been reclassified to conform to the 1995 presentation.

B. Business Combinations

American Triumvirate Insurance Company

On December 21, 1994 the Company acquired the 50% interest in American Triumvirate Insurance Company (ATIC) that it did not previously own. ATIC is a captive insurance company located in Burlington, Vermont. The business combination has been accounted for under the purchase method of accounting. The Company paid \$8,166,000 which equaled 50% of the net book value of ATIC as of December 31, 1994.

Bertek

On February 25, 1993 a wholly-owned subsidiary of the Company acquired substantially all of the net assets of Bertek, Inc. (Bertek). Bertek, headquartered in St. Albans, Vermont, is a manufacturer

of transdermal drug delivery systems and also has operations in laminating and label manufacturing. The business combination has been accounted for under the purchase method of accounting. Goodwill of approximately \$2,686,000 resulting from the acquisition is being amortized on a straight-line basis over a 20 year period.

C. Inventories

Inventories consist of the following components: (in thousands)

March 31,	1995	1994
Raw materials	\$ 29,795	\$ 26,138
Work in process	17,539	14,978
Finished goods	30,871	16,880
ű	\$ 78, 205	\$ 57, 996

D. Property, Plant and Equipment

Property, plant and equipment consist of the following components: (in thousands)

March 31,	1995	1994
Land and land improvements	\$ 5,767	\$ 5,088
Buildings and improvements	48,674	41,705
Machinery and equipment	69,626	59,178
Construction in progress	8,532	9,143
	132,599	115,114
Less accumulated depreciation	40,300	32,600
	\$ 92,299	\$ 82,514

E. Investment in and Advances to Somerset

The Company owns 50% of all the outstanding common stock of Somerset Pharmaceuticals Inc.

(Somerset) and uses the equity method of accounting for its investment. Equity in Earnings of Somerset includes the Company's 50% portion of Somerset's net earnings and expense for amortization of intangible assets resulting from the acquisition of Somerset. Such intangible assets are amortized over a 15 year period. Amortization expense amounted to \$924,000 in 1995, 1994 and 1993. Additionally, the Company's charges to Somerset for management services and product development activities are included in Equity in Earnings of Somerset. These charges have been recorded by Somerset as a reduction of its net earnings. Condensed unaudited balance sheet information of Somerset is as follows: (in thousands)

March 31,	1995	1994	1993
Current assets	\$ 40,539	\$ 27,931	\$ 30,409
Non-current assets	6,854	6,043	2,760
Current liabilities	19,110	14,918	20,675
Payable to owners	1,328	1,002	1,796
Other liabilities	475	642	808

Condensed unaudited income statement information o	f Somerse	et is as	follows	: (in tho	usands	S)
Year ended March 31,		1995		1994		1993
Net sales	\$	125,975	5 \$	111,970	\$	108,518
Cost and expenses		60,270)	50,465		49,872
Income taxes		21,485	5	19,547		21,789
Net earnings	\$	44,220	\$	41,958	\$	36,857

The above information represents 100% of Somerset's operations of which the Company has a 50% interest

Under the Orphan Drug Act, Somerset has exclusivity relating to marketing the chemical compound Eldepryl(R) for use as a treatment for late stage Parkinson's disease through June of

F. Marketable Securities

Effective April 1, 1994 the Company adopted Statement of Financial Accounting Standards No. 115 (SFAS No. 115), "Accounting for Certain Investments in Debt and Equity Securities". Prior years' financial statements have not been restated to apply the provisions of SFAS No. 115. This Standard changes the manner in which certain investments are valued and affects the way in which unrealized gains and losses are recognized for financial reporting purposes. The Company has classified its portfolio of marketable securities as "available for sale". Such securities are recorded at market value with unrealized gains and losses being recognized as a separate component of shareholders' equity, net of income taxes.

The amortized cost and estimated market values at March 31, 1995 are as follows: (in thousands)

	mortized Cost	ss alized ains	Gross Jnrealiz Losse	Market Value
Debt securities:				
U.S. Government obligations	\$ 17,083	\$ 262	\$ 22	\$ 17,323
Municipal obligations	33,047	95	10	33,132
Corporate bonds	3,090	69	75	3,084
Certificates of deposit	15,804	-	-	15,804
Total debt securities	69,024	426	107	69,343
Equity securities	3,400	1,868	78	5,190

Maturities of debt securities at market value are as follows: (in thousands)

Due in one year or less $% \left\{ 1,2,\ldots ,n\right\} =0$ \$ 47,384 Due after one year through five years 16,553 Due after five years 5,406 Total 69,343

Proceeds from sales of marketable securities were \$5,068,000 during 1995. Gross gains of \$14,000 and gross losses of \$142,000 were realized on those sales during 1995. The cost of investments sold is determined by the specific identification method.

G. Intangible Assets

Intangible assets are stated net of accumulated amortization of \$13,874,000 and \$8,874,000 at March 31, 1995 and 1994 respectively. Amortization is provided for on a straight-line basis over periods ranging from 14 to 17 years for patents and technology and 2 to 20 years for other intangible assets.

Values assigned to patents and technology at March 31, 1995 and 1994, were \$20,945,000 and \$22,324,000 respectively. The remaining amounts consist principally of values assigned to licenses, agreements and goodwill.

H. Other Assets

Other assets consist of the following components: (in thousands)

March 31, 1995 1994 Pooled asset funds 14.587 13,758 Cash surrender value 16,377 16,254 Captive insurance company 7,712 Other investments 17,981 10,398 48,945 48,122

Pooled asset funds are carried at the lower of cost or market and include the Company's interest in various funds which invest in common and preferred stocks, bonds, and money market funds. Earnings on these investments included under the caption "Other Income" amounted to \$829,000 in 1005, \$402,000 in 1004, and (\$645,000) in 1005. in 1995, \$402,000 in 1994 and (\$645,000) in 1993. At March 31, 1995 and 1994 the carrying amounts of these investments approximated fair value.

Cash surrender value represents insurance policies on

certain officers and key employees and the value of a split dollar life insurance arrangement with the estate of the former Chairman and Chief Executive Officer of the Company (see note P). Other investments are comprised principally of equity investments in non-publically traded

entities. Such investments are accounted for under the cost method. Prior to December 21, 1994 the Company's interest in a captive insurance company was accounted for using the equity method of accounting (see note B). Earnings from this investment included under the caption "Other Income" amounted to \$454,000 in 1995 prior to acquisition, \$937,000 in 1994 and \$940,000 in 1993.

I. Other Current Liabilities

Other current liabilities includes payroll and employee benefit plan accruals which amounted to \$6,103,000 at March 31, 1995 and \$5,388,000 at March 31, 1994, accruals for Medicaid reimbursements of \$3,640,000 at March 31, 1995 and \$1,479,000 at March 31, 1994 and deferred revenue related to a distribution agreement (see note K) of \$3,500,000 at March 31, 1995.

J. Long-Term Obligations

Long-term obligations represent accruals for post-retirement compensation pursuant to agreements with certain key employees and directors. Under these agreements, benefits are to be paid over periods of 10 to 15 years commencing at retirement.

K. Distribution Agreement

On October 10, 1994 the Company entered into a distribution agreement with STC Pharmaceuticals, Inc. (STC), a wholly owned subsidiary of Eli Lilly and Company (Lilly). Under the terms of the agreement the Company is distributing a generic form of Lilly's oral antibiotic Ceclor(R) on behalf of STC. The Company is being paid a fixed monthly fee for distributing the product. Upon certain events, as defined in the agreement, the fixed monthly fee will convert to a variable amount predicated upon STC's net sales of the product. Revenues and gross profits resulting from this agreement did not have a material impact on operations.

L. Income Taxes

Income taxes consist of the following components: (in thousands)

Year ended March 31, Federal	1995	1994	1993
Current Deferred	\$ 48,851 (8,111) 40,740	\$ 11,888 61 11,949	\$ 25,325 (888) 24,437
State			
Current Deferred	12,033 (2,316) 9,717	2,766 (717) 2,049	2,283 - 2,283
Income taxes	\$ 50,457	\$ 13,998	\$ 26,720
Pre-tax earnings	\$ 171,326	\$ 87,065	\$ 97,341
Effective tax rate	29.5%	16.1%	27.4%

Effective April 1, 1993 the Company adopted Statement of Financial Accounting Standards No. 109 (SFAS No. 109), "Accounting for Income Taxes". The prior year's financial statements have not been restated to apply the provisions of SFAS No. 109. The cumulative effect of adopting this Standard resulted in an increase in net earnings of \$1,124,000 or \$.01 per share in the 1994

Consolidated Statement of Earnings. There was no cash flow impact.
This Standard requires an asset and liability approach to accounting for income taxes. Deferred income tax assets and liabilities reflect the future tax consequences of events that have already been recognized in the financial statements or tax returns. Changes in enacted tax rates or laws will result in adjustments to the recorded tax asset or liabilities in the period that the tax law is enacted.

Temporary differences and carryforwards which give rise to the deferred income tax assets and liabilities are as follows:

(in thousands)

March 31,		1995	1	.994	
Deferred Tax Assets:					
Employee benefits	\$	2,758	\$	1,745	
Deferred revenue		1,881		868	
Asset allowances		5,785		1,279	
Inventory		5,378		829	
Investments		634		1,022	
Total Deferred Tax Assets		16,436		5,743	
Deferred Tax Liabilities:					
Property		2,642		3,379	
Investments		2,206		1,103	
Total Deferred Tax Liabilities		4,848		4,482	
Deferred Tax Assets - Net	\$	11,588	\$	1,261	
Classification in the Consolidated Balance Sheet:					
Deferred Income Tax Benefit - Current	\$	10,545	\$	2,082	
Deferred Income Tax Benefit - Non-Current		1,043		-	
Deferred Income Tax Liability - Non-Current		- '		(821)	
Deferred Tax Assets - Net	\$	11,588	\$	1,261	
A reconciliation of the statutory tax rate to the effect:	ive tax rate	is as fol	lows:		
Year Ended March 31,		1995	1994	1993	
Statutory tax rate		35.0%	35.0%	34.0%	
State income taxes-net		4.2%	1.7%	1.5%	
Tax exempt earnings-					
primarily dividend exclusion		(4.8%)	(7.7%)	(5.9%)
Tax credits		(4.9%)	(7.6%)	(2.2%)
SFAS 109		-	(1.3%)		-
Changes in tax code		-	(3.7%)		
Other items		-	(0.3%)	-	
Effective tax rate		29.5%	16.1%	27.4%	

Tax credits result principally from operations in Puerto Rico. In August of 1993, President Clinton signed into law the Omnibus Budget Reconciliation Act of 1993 ("the Act"). The Act has several provisions which affect the Company's income tax expense including a change in the Federal corporate tax rate and significant changes relating to tax credits for operations in Puerto Rico. As a direct result of the changes in the tax code, the Company reassessed its position on the filing alternatives available under the tax code. Based on the new tax code provisions, the Company made a decision which resulted in a reduction of income tax expense of \$3,225,000. This amount represents management's estimate of the cumulative effect of this change.

M. Common Stock

During fiscal year 1993 the company declared a 2 for 1 stock split effected in the form of a stock dividend. The par value of the new shares issued totaled \$19,327,000 and has been transferred from additional paid-in capital and retained earnings to the common stock account. Per share amounts and stock options have been adjusted for the stock split.

On April 7, 1993, the shareholders of the Company approved an increase in the number of shares of common stock authorized to 300,000,000.

N. Stock Option Plans

On December 1, 1986 the Board of Directors adopted the "Mylan Laboratories Inc. 1986 Incentive Stock Option Plan" ("the Plan") which was approved by the shareholders on June 24, 1987. A total of 6,000,000 shares of the Company's common stock are reserved for issuance upon exercise of stock options. Options, which may be granted at not less than fair market value on the date of the grant may be excercised within ten years from the date of grant. As of March 31, 1995 options for 3,040,300 shares have been granted pursuant to the Plan. On June 23, 1992 the Board of Directors adopted the "1992 Nonemployee Director Stock Option Plan" (the "Directors' Plan") subject to shareholder approval, which was obtained on April 7, 1993. A total of 400,000 shares of the Company's common stock are reserved for issuance upon exercise of stock options which may be granted at not less than fair market value on the date of grant. Shares are granted, based on a formula as described in the Directors' Plan, upon the nonemployee director's initial and subsequent election to the Board of Directors. Options may be exercised within ten years from the date of grant. As of March 31, 1995, 170,000 shares have been granted pursuant to the Directors' Plan. As of March 31, 1995 options for 1,212,299 shares are

exercisable under all plans at option prices ranging from \$4.24\$ to \$28.125\$ per share.A summary of the activity resulting from all plans adjusted for the stock split is as follows:

	Number of shares under option	tion price per share
Outstanding		
April 1, 1992	1,120,504	\$ 4.125-16.00
Options granted	1,778,000	18.00
Options exercised	(713,857)	4.125-18.00
Options cancelled or surrendered	(64,398)	4.75-16.00
Outstanding		
March 31, 1993	2,120,249	\$ 4.125-18.00
Options granted	6,000	26.125-28.125
Options exercised	(347,747)	4.125-18.00
Options cancelled or surrendered	(6,875)	16.00
Outstanding		
March 31, 1994	1,771,627	\$ 4.24-28.125
Options granted	296,000	15.875-22.00
Options exercised	(274, 953)	4.125-18.00
Options cancelled or surrendered	(22,000)	7.25-18.00
Outstanding		
March 31, 1995	1,770,674	\$ 4.24-28.125

O. Profit Sharing and 401(K) Plans

The Company has a noncontributory trusteed profit sharing plan covering essentially all employees who are not covered by 401(k) plans, a 401(k) plan covering essentially all Hickam employees, a profit sharing plan with a 401(k) provision covering all employees of Bertek and a 401(k) plan covering all employees of the bargaining unit.

Contributions to the profit sharing plan and the Bertek plan are made at the discretion of the Board of Directors. Contributions to the Hickam plan are based upon a formula matching the employees salary deferral. Contributions to the bargaining unit plan are based upon the union agreement. Total contributions to all plans for the years ended March 31, 1995, 1994 and 1993 were \$3,060,000, \$2,300,000 and \$1,860,000 respectively.

P. Related Party Transactions

Pursuant to a salary continuation agreement between Mr. McKnight, former Chairman and Chief Executive Officer, and the Company, a one time payment of \$4,306,000 was made on March 31, 1994 of which \$2,861,000 was expensed during 1994. The Company also purchased aircraft, which it previously leased on a flight by flight basis, from the estate of Mr. McKnight for \$5,900,000. In addition, the Company will continue to fund life insurance premiums pursuant to a split-dollar life insurance agreement whereby the Company has rights to the cash surrender value of the insurance policies.

O. Legal Matters

The Company is involved in various legal proceedings that are considered normal to its business. While it is not feasible to predict the ultimate outcome of such proceedings it is the opinion of management that the outcome will have no material adverse effect on the Company's operations or financial position.

During fiscal 1994 the Company settled certain legal matters relating to the Company's suit filed under the Federal Antitrust Laws and the Racketeer Influence and Corrupt Organization Act (RICO), for approximately \$3,375,000. Additionally during fiscal 1994 the jury in the Company's lawsuit against American Cyanamid ruled in favor of Cyanamid on the company's claims and in favor of the Company on Cyanamid's counterclaims. No money damages were awarded to either party. The U.S. Court of Appeals for the Fourth Circuit upheld the jury verdicts on all claims.

We have audited the accompanying consolidated balance sheets of Mylan Laboratories Inc. and subsidiaries as of March 31, 1995 and 1994, and the related consolidated statements of earnings, shareholders' equity, and cash flows for each of the three years in the period ended March 31, 1995, appearing on pages 40 through 51. These fiancial statements are the responsibility of the Company's management. Our responsibility is to express an opinion on these financial statements based on our audits.

We conducted our audits in accordance with generally accepted auditing standards. Those standards require that we plan and perform the audit to obtain reasonable assurance about whether the financial statements are free of material misstatement. An audit includes examining, on a test basis, evidence supporting the amounts and disclosures in the financial statements. An audit also includes assessing the accounting principles used and significant estimates made by management, as well as evaluating the overall financial statement presentation. We believe that our audits provide a reasonable basis for our opinion.

In our opinion, such consolidated financial statements present fairly, in all material respects, the financial position of Mylan Laboratories Inc. and subsidiaries as of March 31, 1995 and 1994, and the results of their operations and their cash flows for each of the three years in the period ended March 31, 1995, in conformity with generally accepted accounting principles.

Pittsburgh, Pennsylvania April 28, 1995

Market Prices

Fiscal 1995	High	Low
First Quarter	23	155/8
Second Quarter	273/8	201/2
Third Quarter	297/8	241/2
Fourth Quarter	333/4	243/4
Fiscal 1994	High	Low
First Quarter	30 3/4	23 5/8
Second Quarter	30 3/8	19 5/8
Third Quarter	33 1/4	23 1/2
Fourth Quarter	25 1/8	15 7/8

New York Stock Exchange Symbol: MYL

On April 30, 1995 the Company had approximately 85,213 shareholders.

Quarterly Financ (Amounts in thou					
except per share	e amounts)				
1st Quarter			1995	199	94
Net sales		\$	85,146	\$	58,507
Gross profit			52,150		29,952
Net earnings			27,130		16,108
Earnings per sha	ire		.34		.21
2nd Quarter					
Net sales		\$	96,013	\$	57,756
Gross profit			55,791		27,848
Net earnings			28,658		17,442
Earnings per sha	ıre		.36		. 22
3rd Quarter					
Net sales	\$	5	104,271	\$	66,436
Gross profit			57,569		34,271
Net earnings			31,839		22,123
Earnings per sha	ıre		. 40		. 28
4th Quarter					
Net sales	\$	5	110,690	\$	69,074
Gross profit			61,020		34,071
Net earnings			33,242		17,394
Earnings per sha			.42		. 22
Year	\$	5	396,120		251,773
Gross profit			226,530	1	L26,142
Net earnings			120,869		73,067
Earnings per sha	ire		1.52		. 93

Design: John Brady Design Consultants Inc., Pittsburgh, Pennsylvania Notice of Annual Meeting

The annual meeting of shareholders of the Company will be held on Wednesday, June 28, 1995 at 10:30 AM at the Lakeview Resort & Conference Center, Morgantown, West Virginia. A formal notice together with a proxy statement and form of proxy will be mailed to shareholders entitled to vote in advance of the meeting.

Shareholder Information

A copy of the Mylan Laboratories Inc. Annual Report to the Securities and Exchange Commission on Form 10-K is available to shareholders on request. For a copy of Form 10-K, please write to:

Mylan Laboratories Inc. 1030 Century Building 130 Seventh Street Pittsburgh, Pennsylvania 15222

Shareholder Contact Patricia Sunseri (412) 232-0100

Directors

Milan Puskar Chairman of the Board, C.E.O. and President of the Company Dana G. Barnett Executive Vice President of the Company

Laurence S. DeLynn Retail Consultant Morgantown, West Virginia

John C. Gaisford, M.D. Director of Burn Research West Penn Hospital Pittsburgh, Pennsylvania

Richard A. Graciano Associate/Partner Graciano Enterprises Pittsburgh, Pennsylvania

Robert W. Smiley, Esq. Doepken, Keevican, Weiss & Medved Attorneys-At-Law Pittsburgh, Pennsylvania

C. B. Todd Senior Vice President of the Company

Officers

Milan Puskar Chairman, C.E.O. and President

Dana G. Barnett Executive Vice President

Louis J. DeBone Vice President-Operations

Roderick P. Jackson Senior Vice President

Joseph J. Krivulka Vice President

Dr. John P. O'Donnell Vice President-Research and Quality Control

Robert W. Smiley, Esq. Secretary

Patricia Sunseri Vice President-Investor and Public Relations

C. B. Todd Senior Vice President Corporate Directory

Mylan Laboratories Inc. 1030 Century Building 130 Seventh Street Pittsburgh, Pennsylvania 15222 (412) 232-0100

Registrar and Transfer Agent American Stock Transfer Co. New York, New York

Certified Public Accountants Deloitte & Touche LLP Pittsburgh, Pennsylvania

Financial Consultants PDA Associates, Inc. Ironia, NJ

Securities Traded New York Stock Exchange Mylan Laboratories Inc. Common Stock Symbol: MYL

- (21) Subsidiaries of the registrant, filed herewith.
- (23) Consents of Independent Auditors', filed herewith.

EXHIBIT 23 Independent Auditors' Consent

We consent to the incorporation by reference in Registration Statements No. 33-65916 and No. 33-65918 of Mylan Laboratories Inc. on Form S-8 of our report dated April 28, 1995,

incorporated by reference in this Annual Report on Form 10-K of Mylan Laboratories Inc. for the year ended March 31, 1995.

/s/ Deloitte & Touche LLP Pittsburgh, Pennsylvania June 16,1995

Independent Auditors' Consent

We consent to the incorporation by reference in Registration Statements No. 33-65916 and No. 33-65918 of Mylan Laboratories Inc. on Form S-8 of our report dated February 3, 1995 relating to the consolidated financial statements of Somerset Pharmaceuticals, Inc. and subsidiaries for the three years in the period ended December 31,1994, included in the Annual Report on Form 10-K of Mylan Laboratories Inc. for the year ended March 31,1995.

/s/ Deloitte & Touche LLP Pittsburgh, Pennsylvania June 16, 1995

- (27) Financial Data Schedule, filed herewith.
- (99) Consolidated financial statements of Somerset Pharmaceuticals, Inc. for Years ended December 31, 1994, 1993 and 1992, filed herewith.

SOMERSET PHARMACEUTICALS, INC. AND SUBSIDIARIES

Consolidated Financial Statements for the Years Ended December 31, 1994, 1993 and 1992, and Independent Auditors' Report

INDEPENDENT AUDITORS' REPORT
To the Board of Directors of
 Somerset Pharmaceuticals, Inc.:

We have audited the accompanying consolidated balance sheets of Somerset Pharmaceuticals, Inc. and subsidiaries as of December 31, 1994 and 1993, and the related consolidated statements of income, stockholders' equity, and cash flows for each of the three years in the period ended December 31, 1994. These financial statements are the responsibility of the Company's management. Our responsibility is to express an opinion on these financial statements based on our audits.

We conducted our audits in accordance with generally accepted auditing standards. Those standards require that we plan and perform the audit to obtain reasonable assurance about whether the financial statements are free of material misstatement. An audit includes examining, on a test basis, evidence supporting the amounts and disclosures in the financial statements. An audit also includes assessing the accounting principles used and significant estimates made by management, as well as evaluating the overall financial statement presentation. We believe that our audits provide a reasonable basis for our opinion.

In our opinion, such consolidated financial statements present fairly, in all material respects, the financial position of Somerset Pharmaceuticals, Inc. and subsidiaries as of December 31, 1994 and 1993, and the results of their operations and their cash flows for each of the three years in the period ended December 31, 1994 in conformity with generally accepted accounting principles.

February 3, 1995

SOMERSET PHARMACEUTICALS, INC. AND SUBSIDIARIES

NOTES TO CONSOLIDATED FINANCIAL STATEMENTS YEARS ENDED DECEMBER 31, 1994, 1993 AND 1992 SOMERSET PHARMACEUTICALS, INC. AND SUBSIDIARIES

CONSOLIDATED BALANCE SHEETS

DECEMBER 31, 1994 AND 1993

ASSETS 1994 1993

CURRENT ASSETS:

Cash and cash equivalents	\$17,529,000	\$10,281,000
Investment securities	3,338,000	3,470,000
Accounts receivable (net of allowance for doubtful		
accounts of \$100,000)	20,653,000	16,095,000
Inventories	5,293,000	3,820,000
Prepaid expenses and other current assets	1.957.000	1,582,000

Total current assets 48,770,000 35,248,000

PROPERTY AND EQUIPMENT - Net 4,266,000 2,762,000

INTANGIBLE ASSETS - Net 1,644,000 1,987,000

OTHER ASSETS 470,000 1,416,000

\$55,150,000 \$41,413,000

Liabilities and Stock holder's Equity 1994 1993 Current Liabilities: Accounts payable 292,000 205,000 Note payable 253,000 Accrued marketing costs 11,000,000 9,100,000 Royalty payable 5,850,000 4,780,000 Other accrued expenses 2,833,000 2,070,000 Accrued research and development 1,901,000 2,046,000 Income taxes payable 5,017,000 2,900,000 Amounts due to related paries 2,318,000 2,063,000 Total current liabilities 29,211,000 23,417,000 Deferred revenue 292,000 458,000 Stockholder's Equity: Common stock, \$.01 par value; 13,719 shares authorized, 11,297 shared issued Retained earnings 26,099,000 17,990,000 Less treasury stock, 644 shares at cost (452,000) (452,000) 17,538,000 Total stockholder's equity 25,647,000 \$55,150,000 \$41,413,000

See notes to consolidated financial statements.

SOMERSET PHARMACEUTICALS, INC. AND SUBSIDIARIES

CONSOLIDATED STATEMENTS OF INCOME YEARS ENDED DECEMBER 31,1994, 1993 AND 1992

	1994	1993	1992
NET SALES	\$124,566,000	\$118,998,000	\$104,071,000
COSTS AND EXPENSES: Cost of sales Marketing Research and development Administrative	16,399,000 23,457,000 10,424,000 9,845,000 60,125,000	13,991,000 25,826,000 9,134,000 8,005,000 56,956,000	12,552,000 23,415,000 5,580,000 6,736,000 48,283,000
	64,441,000	62,042,000	55,788,000
OTHER INCOME	568,000	1,131,000	1,017,000
INCOME BEFORE INCOME TAXES 56,805,000	65,009,000	63,173,000	
PROVISION FOR INCOME TAXES 20,736,000	20,900,000	21,408,000	
NET INCOME	\$ 44,109,000	\$ 41,765,000	\$ 36,069,000

See notes to consolidated financial statements.

SOMERSET PHARMACEUTICALS, INC. AND SUBSIDIARIES

CONSOLIDATED STATEMENTS OF STOCKHOLDERS' EQUITY

YEARS ENDED DECEMBER 31, 1994, 1993 AND 1992

BALANCE, DECEMBER 31, 1991 \$ 7,448,000	11,297 \$	-	644	\$ (452,000)	\$ 7,900,000	
Accretion of the carrying value of the redeemable preferred stock Dividends Net income	- - - -	- - -	- - -	- - -	(122,000) (41,207,000) 36,069,000	(122,000) (41,207,000) 36,069,000
BALANCE, DECEMBER 31, 1992 2,188,000	11,297	-	644	(452,000)	2,640,000	
Accretion of the carrying value of the redeemable preferred stock Dividends Net income	- - -	- - -	- - -	- - -	(15,000) (26,400,000) 41,765,000	(15,000) (26,400,000) 41,765,000
BALANCE, DECEMBER 31, 1993 17,538,000	11,297	-	644	(452,000)	17,990,000	
Dividends Net income	-	-	- -	- -	(36,000,000) 44,109,000	(36,000,000) 44,109,000
BALANCE, DECEMBER 31, 1994 \$ 25,647,000	11,297 \$	-	644	\$ (452,000)	\$ 26,099,000	

See notes to consolidated financial statements.

SOMERSET PHARMACEUTICALS, INC. AND SUBSIDIARIES

CONSOLIDATED STATEMENTS OF CASH FLOWS YEARS ENDED DECEMBER 31, 1994, 1993 AND 1992

	1994	1993	1992
CASH FLOWS FROM OPERATING ACTIVITIES: Net income Adjustments to reconcile net income to net cash provided by operating activities:	\$44,109,000	\$41,765,000	\$36,069,000
Depreciation and amortization	587,000	285,000	229,000
Deferred tax expense (benefit)	862,000	,	,
Deferred revenue	(166,000)	, , ,	,
Changes in operating assets and liabilities:	, ,	(- , ,	(,,
Accounts receivable	(4,558,000)	(1,872,000)	(3,989,000)
Inventories	(1,473,000)	(1,578,000)	614,000
Prepaid expenses and other current assets	(375,000)	352,000	(249,000)
Accounts payable	87,000	(227,000)	(1,121,000)
Royalty payable	1,070,000	190,000	901,000
Accrued marketing costs	1,900,000	1,386,000	2,074,000
Accrued research and development	(145,000)		-
Other accrued expenses	763,000	,	, ,
Income taxes payable	2,117,000	570,000	(177,000)
Amounts due to related parties	255,000	278,000	542,000
Net cash provided by operating activities	45,033,000	41,364,000	36,085,000
CASH FLOWS FROM INVESTING ACTIVITIES:			
Net decrease (increase) in investment securities	132,000	2,006,000	(1,685,000)
Purchase of property and equipment	(1,898,000)	(2,690,000)	(127,000)
Decrease (increase) in other assets	234,000	(268,000)	23,000
Net cash used in investing activities	(1,532,000)	(952,000)	(1,789,000)

(Continued)
SOMERSET PHARMACEUTICALS, INC. AND SUBSIDIARIES

CONSOLIDATED STATEMENTS OF CASH FLOWS YEARS ENDED DECEMBER 31, 1994, 1993 AND 1992

	1994	1993	1992
CASH FLOWS FROM FINANCING ACTIVITIES: Redemption of preferred stock Dividends paid on preferred stock Dividends paid on common stock (36,300,000) Net (decrease) increase in note payable Net cash used in financing activities (37,534,000)	\$ - (36,000,000) (253,000) (36,253,000)	253,000	\$ (1,149,000) (85,000)
NET INCREASE (DECREASE) IN CASH AND CASH EQUIVALENTS (3,238,000)	7,248,000	4,140,000	
CASH AND CASH EQUIVALENTS, BEGINNING OF YEAR 9,379,000	10,281,000	6,141,000	

END OF YEAR	\$ 17,529,000	\$ 10,281,000	\$ 6,141,000
SUPPLEMENTAL DISCLOSURES OF CASH FLOW INFORMATION: Cash paid during the year for:			
Interest	\$ 7,000	\$ 5,000	\$ -
Income taxes	\$ 17,683,000	\$ 21,259,000	\$ 20,992,000

SUPPLEMENTAL DISCLOSURE OF NONCASH FINANCING ACTIVITIES:

During 1992, the Company recorded \$8,800,000 of dividends payable on common stock which had not been paid as of the end of that year.

See notes to consolidated financial statements.

CASH AND CASH FOLITIVALENTS

PRINCIPLES OF CONSOLIDATION AND OPERATIONS

The consolidated financial statements include the accounts of Somerset Pharmaceuticals, Inc. (the Company) and its wholly owned subsidiaries, Somerset Pharmaceuticals Holding Company and Somerset Caribe, Inc. The Company is jointly owned by Mylan Laboratories, Inc. and Circa Pharmaceuticals, Inc., with each owning 50% of all the outstanding common stock of the Company. All significant intercompany accounts and transactions have been eliminated in consolidation. The Company, incorporated in February 1986, is engaged in the development, testing and marketing of drugs to be used in the treatment of various human disorders. Currently, the Company manufactures, markets and sells Eldepryl, which is used as a treatment for Parkinson's Disease.

The Company is party to an exclusive 14-year agreement (through November 22, 2003) with Chinoin Pharmaceutical Company (Chinoin) of Budapest, Hungary under which Eldepryl and other new potential drugs resulting from Chinoin research are made available for licensing by the Company. The license agreement requires the Company to pay royalties equal to 7% of net sales of Eldepryl including sub-license revenues. The Company incurred royalty expense of approximately \$9,983,000, \$8,383,000 and \$8,105,000 for the years ended December 31, 1994, 1993 and 1992, respectively.

SUMMARY OF SIGNIFICANT ACCOUNTING POLICIES

- a. Cash and Cash Equivalents The Company generally considers debt instruments purchased with a maturity of three months or less to be cash equivalents.
- b. Investment Securities Effective January 1, 1994, the Company adopted Statement of Financial Accounting Standards (SFAS) No. 115, "Accounting for Certain Investments in Debt and Equity Securities." The effect of adopting SFAS No. 115 on the Company's financial statements was not material. During the current year, gross proceeds from sales and maturities of investments approximated \$797,000 and \$750,000, respectively, and realized gains or losses were not material. At December 31, 1994, the investment securities were available for sale, and there were no material unrealized gains or losses.

At December 31, 1993, investment securities included both debt and equity instruments which were valued at the lower of cost or market with cost approximating market.

- c. Inventories Inventory is stated at the lower of cost or market, with cost determined on a first-in, first-out basis.
- d. Property and Equipment Property and equipment are stated at cost. Depreciation is provided over the estimated useful lives of the assets by the straight-line method. Estimated useful lives are five to seven years for machinery and equipment and thirty-five years for the building.
- e. Intangible Assets Intangible assets are amortized on a straight-line basis over 14 years.
- f. Research and Development Research and development costs are expensed as incurred.

3. INVENTORY

Inventory consists of the following at December 31:

	1994	1993
Raw material Work in process Finished goods	\$ 4,686,000 \$ 375,000 232,000	2,864,000 470,000 486,000
Total	\$ 5,293,000 \$	3,820,000

4. PROPERTY AND EQUIPMENT

Property and equipment consist of the following at December 31:

1994 1993

Land	\$ 300,000	\$ 300,000
Building	2,067,000	1,638,000
Machinery and equipment	2,410,000	963,000
Furniture and fixtures	87,000	65,000
	4,864,000	2,966,000
Less accumulated depreciation	(598,000)	(204,000)

SUB-LICENSE OF RIGHTS

On February 9, 1988, the Company granted a sub-license to its exclusive right and license to use its technology to Draxis Health Inc. (formerly Deprenyl Research Limited) to commercialize certain drugs in Canada for 15 years. The Company receives a royalty of 11% of Draxis Health Inc.'s net sales over the license period.

Royalty income, less related royalty expense to Chinoin, included in other income for the years ended December 31, 1994, 1993 and 1992 was approximately \$199,000, \$357,000 and \$414,000, respectively.

6. INTANGIBLE ASSETS

Intangible assets primarily represent the cost of a modification to the terms of the Chinoin Agreement, less accumulated amortization of \$1,061,000 and \$868,000 at December 31, 1994 and 1993, respectively.

7. CO-PROMOTIONAL AGREEMENT

Effective October 1, 1990, the Company entered into an agreement with Sandoz Pharmaceuticals Corporation (Sandoz) to co-promote the product Eldepryl. Under the terms of the agreement, the Company is required to make certain payments to Sandoz in the event sales of Eldepryl exceed certain predefined minimums. The agreement requires Sandoz, among other things, to expend, at a minimum, a predetermined amount for advertising during each year of the agreement. Once the predetermined levels of sales are exceeded, the Company is required to pay Sandoz for advertising expenditures made on behalf of the Company. After Sandoz's advertising expenses are reimbursed, any additional amounts are shared by Sandoz and the Company based upon the terms of the agreement. During 1994, 1993 and 1992, the Company expensed approximately \$22,360,000, \$24,260,000 and \$22,321,000, respectively, pursuant to the agreement. Additionally, certain co-promotional fees paid by Sandoz at the commencement of the agreement are being recognized ratably by the Company during the term of the agreement (six years), and certain costs associated with the procurement, negotiation and execution of the agreement by the owners of the Company are being incurred by the Company in approximately the same amount.

In December 1994, the Company amended its co-promotional agreement with Sandoz. The amended agreement eliminated certain residual period payments to Sandoz, shortened the term to March 31, 1996, eliminated certain sales force detail requirements and requires certain payments to be made to the Company if a predetermined level of sales is not achieved.

8. NOTE PAYABLE

On June 30, 1993, the Company entered into a one-year \$1,500,000 line of credit agreement with a bank in conjunction with the renovation of the Company's research facility. Interest on amounts drawn on the line of credit was payable monthly at the bank's prime rate (6% at December 31, 1993) less 0.25%. Pursuant to the agreement, all borrowings under this line of credit were paid off in 1994.

9. REDEEMABLE PREFERRED STOCK

The Class A and B redeemable preferred stock were fully redeemed during 1993.

The Class A stock, (\$.10 par value, 1,950 shares authorized and 488 shares outstanding at December 31, 1992) was carried at redemption value plus undeclared dividends.

The Class B stock (\$.10 par value, 2,850 shares authorized and issued and 661 shares outstanding at December 31, 1992) was convertible into common stock based on a conversion price set by the Board of Directors and subject to adjustment from time to time.

10. INCOME TAXES

The Company adopted Statement of Financial Accounting Standards (SFAS) No. 109, "Accounting for Income Taxes" effective January 1, 1993. As permitted by SFAS No. 109, prior-year financial statements have not been restated to reflect the change in accounting method. The cumulative effect of adopting SFAS No. 109 on the Company's financial statements was not material.

The income tax provision consists of the following for the years ended December ${\bf 31:}$

	1994	1993	1992
Current tax expense: Federal State Foreign	\$ 15,025,000 4,899,000 114,000	\$ 17,938,000 4,124,000 146,000	\$ 18,540,000 2,050,000 126,000
	20,038,000	22,208,000	20,716,000
Deferred tax expense (benefit): Federal State	754,000 108,000	(700,000) (100,000)	20,000
	862,000	(800,000)	20,000
Total provision for income taxes	\$ 20,900,000	\$ 21,408,000	\$ 20,736,000

Deferred income taxes reflect the net tax effects of temporary differences between the carrying amounts of assets and liabilities for financial reporting purposes and the amounts used for income tax purposes. The tax effects of significant items comprising the Company's deferred

taxes (which are included in "Other Assets" in the balance sheet) as of December 31, 1994 are as

Deferred tax assets:

Deferred revenue Deferred compensation Chargeback allowance Other	\$ 110,000 228,000 152,000 42,000
	532,000

Deferred tax liabilities:

Excess of tax amortization over reporting amortization 165,000

Net deferred tax assets \$ 367,000

The statutory federal income tax rate is reconciled to the effective tax rate as follows for the years ended December 31:

	1994	1993	1992
Tax at statutory rate State income tax (net of federal benefit) Tax credits Tollgate tax Other	35.0% 3.5 (9.9) 3.9 (.4)	35.0% 4.1 (7.2) 2.0	34.0% 2.4 (.2) - .3
Effective tax rate	32.1%	33.9%	36.5%

Tax credits result principally from operations in Puerto Rico.

11. RELATED PARTY TRANSACTIONS

The Company incurs expenses for ongoing management services and over a six year period for specific services related to the procurement, negotiation and execution of the co-promotion agreement by the owners of the Company. The Company also incurs other expenses from one or both of its owners as detailed below for the years ended December 31:

	1994	1993	1992
Management fees	\$ 6,228,000 \$	5,950,000 \$	5,204,000
Research and development	1,020,000	835,000	239,000
Inventory handling and distribution fees	650,000	750,000	331,000
Rent - equipment and facilities	1,065,000	647,000	- '
Product liability insurance	618,000	675,000	675,000

During 1993, the Company purchased \$696,000 of equipment from one of its owners.

At December 31, 1994 and 1993 the balance of amounts due to related parties represents rent, research and development, distribution and management fees payable to the owners of the Company.

12. SIGNIFICANT CUSTOMERS

The Company had sales to certain customers which individually exceeded 10% of net sales. Three customers represented 57% of net sales for the year ended December 31, 1994 and two customers represented 45% and 41% of net sales for the years ended December 31, 1993 and 1992, respectively.

13. COMMITMENTS

As of December 31, 1994, the Company is committed to fund approximately \$5,160,000 for various research and development studies through 1995.

14. EMPLOYEE BENEFIT PLANS

The Company has a defined contribution profit sharing plan covering substantially all employees. Contributions are made at the discretion of the Board of Directors. Additionally, during 1994, the Company initiated a deferred compensation plan for select key employees. Contributions are based on profitability levels for the year. During 1994, 1993 and 1992, the Company recorded expense of \$755,000, \$100,000 and \$-0- for these plans, respectively.

* * * * * *

(b) Reports on Form 8-K

The Company was not required to file a report on Form 8-K during the quarter ended March 31, 1995.

SIGNATURES

Pursuant to the requirements of section 13 or 15(d) of the Securities Exchange Act of 1934, the registrant has duly caused this Report to be signed on its behalf by the undersigned, thereunto duly authorized.

Date: June 16, 1995 MYLAN LABORATORIES INC.

> by /s/ Milan Puskar Milan Puskar

Chairman, Chief Executive Officer and President

Pursuant to the requirements of the Securities Exchange Act of 1934, this Report has been signed below by the following persons on behalf of the registrant and in the capacities and on the dates indicated.

/s/ Milan Puskar June 16,1995 1995

/s/ Dana G. Barnett June 16,1995

June 16,

Milan Puskar

Chairman, Chief Executive Officer and

Dana G. Barnett Executive Vice President and Director

President

/s/ Laurence S. DeLynn June 16, 1995

Laurence S. DeLynn

/s/ Robert W. Smiley June 16,1995 Robert W. Smiley

Director

Secretary and Director

/s/ Richard A. Graciano June 16, 1995 Richard a. Graciano

Director

/s/ John C. Gaisford M.D. June 16, 1995 John C. Gaisford, M.D.

Director

/s/ C.B. Todd June 16,1995 C.B. Todd

Senior Vice President and Director

/s/ Frank A. DeGeorge June 16, 1995

Frank A. DeGeorge Director of Corporate Finance as

Chief Accounting Officer

EXHIBIT 22

Subsidiaries

Name State of Incorporation

Milan Holding, Inc. Delaware
Mylan Inc. Delaware

Mylan Pharmaceuticals Inc. West Virginia

Dow Hickam Pharmaceuticals, Inc. Texas

Bertek, Inc. West Virginia

American Triumvirate Insurance Company Vermont

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YEAR

MAR-31-1995

MAR-31-1995

127,280,000

52,575,000

58,343,000

14,777,000

78,205,000

331,383,000

40,300,000

546,201,000

56,351,000

0

0

39,986,000

442,742,000

546,201,000

396,120,000

396,120,000

169,590,000

88,568,000

0

25,000

171,326,000

50,457,000

120,869,000

0

120,869,000

1.52

1.52
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