

SECURITIES AND EXCHANGE COMMISSION
WASHINGTON, D.C. 20549

FORM 10-Q/A
Amendment No. 1

(MARK ONE)

QUARTERLY REPORT PURSUANT TO SECTION 13 OR 15(d) OF THE SECURITIES
--- EXCHANGE ACT OF 1934.

For the quarterly period ended March 31, 1995
OR

--- TRANSACTION REPORT PURSUANT TO SECTION 13 OR 15(D) OF THE
SECURITIES EXCHANGE ACT OF 1934

For the transition period from _____ to _____

COMMISSION FILE NUMBER 1-10113

HALSEY DRUG CO., INC.

(Exact name of registrant as specified in its charter)

New York

11-0853640

(State or other Jurisdiction of
incorporation or organization)

(I.R.S. Employer Identification No.)

1827 Pacific Street
Brooklyn, New York

11233

(Address of Principal executive officer)

(Zip Code)

(718) 467-7500

(Registrants telephone number, including area code)

Not Applicable

(Former name, former address and former fiscal year, if changed since last
report.)

Indicate by check mark 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 50 days. (Form 10-K for the year ended December 31,
1994, and Forms 10-Q for the quarters ended June 30, 1994 and September 30, 1994
were filed late.)

YES NO X
--- ---

As of May 10, 1995 the registrant had 8,109,537 shares of Common Stock, \$0.1 par
value, outstanding.

HALSEY DRUG CO., & SUBSIDIARIES

INDEX

PART I. FINANCIAL INFORMATION

Item 1. Financial Statements (Unaudited)	Page #

Condensed Consolidated Sheets- March 31, 1995 and December 31, 1994	3
Condensed Consolidated Statements of Operations - Three months ended March 31, 1995 and March 31, 1994	5
Condensed Consolidated Statements of Cash Flows - Three months ended March 31, 1995 and March 31, 1994	6
Condensed Consolidated Statements of Stockholders Equity - Three months ended March 31, 1995	7
Notes to Condensed Consolidated Financial Statements	8
Item 2. Management's Discussion and Analysis of Financial Condition and Results of Operations	14
PART II. OTHER INFORMATION	

Item 6. Exhibits and Reports on Form 8-K	
SIGNATURES	18

PART I. FINANCIAL INFORMATION
HALSEY DRUG CO., INC. AND SUBSIDIARIES
CONSOLIDATED BALANCE SHEETS

	(UNAUDITED)	
	MARCH 31,	DECEMBER
	1995	1994

CURRENT ASSETS		
Cash	\$172,000	\$28,000
Accounts Receivable - trade, net of allowances for doubtful accounts of \$524,000 and \$755,000 1995 and 1994, respectively	2,262,000	2,326,000
Inventories	7,224,000	6,835,000
Prepaid insurance and other current assets	283,000	496,000
Deferred income tax	-----	296,000
Total current assets	9,941,000	9,981,000
PROPERTY PLANT & EQUIPMENT, NET	8,091,000	8,561,000
OTHER ASSETS	604,000	734,000
	-----	-----
	\$18,636,000	\$19,276,000
	-----	-----
	-----	-----

The accompanying notes are an integral part of these statements

PART I. FINANCIAL INFORMATION
 HALSEY DRUG CO., INC. AND SUBSIDIARIES
 CONSOLIDATED BALANCE SHEETS

	(UNAUDITED) MARCH 31, 1995	DECEMBER 1994
<hr style="border-top: 1px dashed black;"/>		
LIABILITIES AND STOCKHOLDERS' EQUITY		
CURRENT LIABILITIES		
Bank overdraft	\$ 365,000	\$ 218,000
Current maturities of long-term debt	3,778,000	4,850,000
Department of Justice settlement	2,044,000	2,013,000
Accounts payable	2,597,000	4,414,000
Accrued expenses and other liabilities	1,785,000	1,823,000
Advances from minority stockholders	206,000	418,000
Income taxes Payable	178,000	196,000
Deferred income		500,000
	-----	-----
Total current liabilities	10,953,000	14,432,000
LONG-TERM DEBT	2,518,000	2,492,000
LITIGATION SETTLEMENT	3,000,000	3,000,000
CONTINGENCIES		
STOCKHOLDERS' EQUITY (DEFICIT)		
Common stock - \$01 par value; authorized 20,000,000 shares; issued and outstanding 8,109,537 shares in 1995 and 7,609,537 in 1994 respectively	81,000	76,000
Additional paid-in capital	11,139,000	10,162,000
Accumulated deficit	(9,055,000)	(10,886,000)
	-----	-----
	2,165,000	(648,000)
	-----	-----
	\$18,636,000	\$19,276,000
	-----	-----
	-----	-----

The accompanying notes are an integral part of these statements

PART I. FINANCIAL INFORMATION
 HALSEY DRUG CO., INC. AND SUBSIDIARIES
 CONSOLIDATED STATEMENTS OF OPERATIONS
 (UNAUDITED)

THREE MONTHS ENDED
MARCH 31

	1995	1994
	-----	-----
Net Sales	\$6,873,000	\$7,324,000
Cost of good sold	5,114,000	5,733,000
	-----	-----
Gross profit	1,759,000	1,591,000
Research & Development	156,000	32,000
Selling, general and administrative expenses	1,538,000	1,709,000
	-----	-----
Earnings (loss) from operations	65,000	(150,000)
Gain on the sale of assets	2,288,000	
Interest expense	(226,000)	(160,000)
	-----	-----
Earnings (loss) before income taxes and cumulative effect of accounting change	2,127,000	(310,000)
Provision for income taxes	296,000	
	-----	-----
NET EARNINGS (LOSS)	\$1,831,000	(\$310,000)
	=====	=====
Per Share Amounts:		
Net earnings (loss)	\$.23	(\$0.04)
	=====	=====
Average number of outstanding shares	7,859,537	7,109,537
	=====	=====

The accompanying notes are an integral part of these statements

5

PART I. FINANCIAL INFORMATION
HALSEY DRUG CO., AND SUBSIDIARIES
CONSOLIDATED STATEMENTS OF CASH FLOWS
(UNAUDITED)

	THREE MONTHS ENDED MARCH 31	
	1995	1994
	-----	-----
CASH FLOWS FROM OPERATING ACTIVITIES		
Net Earnings(Loss)	\$1,831,000	(\$310,000)
	-----	-----
Adjustments to reconcile net (loss) earnings to net cash (used in) provided by operating activities		
Depreciation and amortization	419,000	506,000
Provision for losses on accounts receivable	6,000	
Gain on sale of property, plant and equipment	(2,288,000)	
Accrued Department of Justice interest	30,000	
Deferred income taxes	296,000	
Changes in assets and liabilities		

Accounts receivable	58,000	(421,000)
Inventories	(389,000)	1,001,000
Income taxes receivable		
Prepaid insurance and other current assets	213,000	250,000
Accounts payable	(1,817,000)	(1,249,000)
Accrued expenses	(21,000)	(367,000)
Income taxes payable	(18,000)	(22,000)
	-----	-----
Total adjustments	(3,511,000)	(302,000)
	-----	-----
Net cash (used in) provided by operating activities	(1,680,000)	(612,000)
	-----	-----
Cash flows from investing activities		
Capital expenditures	(50,000)	
Increase in other assets	20,000	115,000
Proceeds from sale of property, plant & equipment	2,000,000	
	-----	-----
Net cash provided by (used in) investing activities	1,970,000	115,000
	-----	-----
Cash flows from financing activities		
Payment of long term debt	(1,064,000)	(47,000)
Proceeds from issuance of common stock	982,000	
Payment to Department of Justice		(87,000)
Bank overdraft	146,000	252,000
Advances from minority stockholder	(212,000)	378,000
	-----	-----
Net cash (used in) provided by financing activities	(146,000)	496,000
	-----	-----
NET DECREASE IN CASH AND CASH EQUIVALENTS	144,000	(1,000)
Cash and cash equivalents at beginning of period	28,000	32,000
	-----	-----
Cash and cash equivalents at end of period	\$172,000	\$ 31,000
	=====	=====

The accompanying notes are an integral part of these statements

6

PART I. FINANCIAL INFORMATION
HALSEY DRUG CO., INC. AND SUBSIDIARIES
CONSOLIDATED STATEMENTS OF STOCKHOLDERS' EQUITY
THREE MONTHS ENDED MARCH 31, 1995

	(UNAUDITED)				
	Common Stock \$0.1 par value Shares	Amount	Additional Paid-In Capital	Accumulated Deficit	Total
Balance at December 31, 1994	7,609,537	\$76,000	\$10,162,000	\$(10,886,000)	\$(648,000)
Net earnings for the three months ended March 31, 1995				1,831,000	1,831,000
Issuance of common stock	500,000	5,000	977,000	-0-	982,000
	-----	-----	-----	-----	-----
Balance at March 31, 1995	8,109,537	\$81,000	\$11,139,000	\$(9,055,000)	\$2,165,000
	=====	=====	=====	=====	=====

The accompanying notes are an integral part of these statements

7

NOTES TO CONDENSED CONSOLIDATED FINANCIAL STATEMENTS

(UNAUDITED)

NOTE 1 - Basis of Presentation

The accompanying unaudited condensed consolidated financial statements of Halsey Drug Co., Inc. (the "Company") have been prepared in accordance with generally accepted accounting principles for interim financial information and with the instructions to Form 10-Q and rule 10-01 of Regulation S-X. Accordingly, they do not include all of the information and footnotes required by generally accepted accounting principles for complete financial statements. In the opinion of management, all adjustments considered necessary for the three month period ended March 31, 1995 have been made, but the financial results for the three months period ended March 31, 1995 are not necessarily indicative of the results that may be expected for the full year ended December 31, 1995. The unaudited condensed consolidated financial statements should be read in conjunction with the consolidated financial statements and footnotes thereto for the year ended December 31, 1994 included in the Company's Annual Report on Form 10-K.

Note 2 - Inventories

Inventories consists of the following:

	March 31, 1995	December 31, 1994
	-----	-----
Finished Goods	\$2,088,000	\$1,990,000
Work in Process	1,368,000	\$1,301,000
Raw Materials	3,768,000	\$3,544,000
	-----	-----
	\$7,224,000	\$6,835,000
	=====	=====

NOTE 3 - Lines of Credit and Long-Term Debt

Lines of Credit

In December 1992, the Company entered into a new credit agreement providing for borrowings of up to \$7,000,000 at the prime rate plus an initial margin of 1/2%, originally maturing in December 1994. Upon certain conditions, as defined in the agreement, the margin rate increases by 2%. Borrowings under the line were available for working capital purposes based upon a percentage of the parent company's eligible accounts receivable and are collateralized by such accounts receivable. The agreement contains certain financial covenants, including minimum interest coverage and working capital ratios, tangible net worth, limitations on capital expenditures, and maximum debt-to-equity ratios. As of March 31, 1995, the Company was not in compliance with the above covenants.

The Company and its banks amended the credit agreement to include the stock of certain subsidiaries, the accounts receivable of Houba, Inc., and the parent company's inventory and equipment as additional collateral, to increase the initial margin rate to 2% (10.5% at March 31,

1995), to restrict certain payments made by the Company, to require payment to be made by the Company to the banks of any income tax refunds received by the Company, to extend the maturity date to August 31, 1995, and to agree in principal to modify the financial covenants at a later date. In addition, if the outstanding borrowings are not repaid by August 31, 1995, the Company is required to pay an additional 3% of the ten outstanding principal due to the banks.

As consideration for the above amendments and the Company's continued borrowings in excess of the borrowing formula, the Company has issued to the banks stock warrants, expiring December 31, 1999, to purchase up to 600,271 shares of the Company's common stock at exercise prices ranging from \$2.25 to \$2.375 per share (subject to the anti-dilution provisions of the credit

agreement, as amended). The fair value of the warrants, \$200,000, as determined by the Company's Board of Directors, has been recorded by the Company as additional paid-in capital and a discount to bank debt which is being amortized through the maturity date, August 31, 1995.

Convertible Subordinated Promissory Note

Pursuant to the Zatpack, Inc. ("Zatpack") agreement (Note N), the Company issued a convertible subordinated promissory note dated December 31, 1994, to Zatpack, for the cancellation of trade payables and advances by Zuellich to the Company's subsidiaries, in the amount of \$1,292,242, bearing interest at 8% per annum, compounded annually, due December 1, 1997. The outstanding principal, plus all accrued and unpaid interest, can be converted, at the option of Zatpack, into the Company's common stock at the rate of one share of common stock for every \$2.50 of principal and interest being converted (the \$2.50 is subject to the anti-dilution provisions of the promissory note). The note is subordinated to bank debt.

Subordinated Promissory Note

On March 21, 1995, the Company satisfied certain accounts payable by issuing a subordinated promissory note to Mallinckrodt Chemical Acquisition, Inc. ("Mallinckrodt") for \$1,200,000, bearing interest at 8% per annum, with interest and principal payable at the earlier of: (i) receipt by Mallinckrodt of all necessary authorization from the FDA or (ii) September 21, 1997 (Note N). The note is collateralized by substantially all of the assets of the Company and is subordinated to future bank indebtedness of up to \$8,000,000. The \$1,200,000 note represents the deferral of payment by the Company of a portion of its December 31, 1994 accounts payable due to an affiliate of Mallinckrodt.

Borrowings under lines of credit and long-term debt consist of the following at March 31, 1995 and December 31, 1994.

	1995 ----	1994 ----
Borrowing under lines of credit	\$3,778,000	\$4,850,000
Convertible subordinated promissory note	1,318,000	1,292,000
Subordinated promissory note	1,200,000	1,200,000
	-----	-----
	\$6,296,000	\$7,342,000
Less current maturities, including amounts in default	\$3,778,000	\$4,850,000
	-----	-----
	\$2,518,000	\$2,492,000
	-----	-----
	-----	-----

NOTE 4 - Gain on Sale of Assets

On March 21, 1995, the Company sold its abbreviated new drug application ("ANDA") for 5mg Oxycodone HCl/325mg and Acetaminophen Tablets ("Tablets") and certain equipment used in the production of the Tablets for up to \$5.4 million to Mallinckrodt. The Company received \$500,000 of the proceeds in July 1994, which was recorded as deferred income on the Company's December 31, 1994 consolidated balance sheet. Mallinckrodt also paid the Company \$2,000,000 on March 21, 1995 and the remainder will be payable as follows: (i) \$1,000,000 upon the Company receiving general clearance from the FDA for unrestricted operations at its Brooklyn facility and written notice from the FDA that it is in compliance with certain provisions of the consent decree dated June 29, 1993 and (ii) \$1,900,000 at the earlier of (a) Mallinckrodt receiving certain authorizations from the FDA or (b) September 21, 1997 ("Deferred Payments"). Mallinckrodt also agreed to defer \$1,200,000 of the Company's trade debt due to an affiliate of Mallinckrodt. For the three months ended March 31, 1995, the Company has recorded a gain of \$2,288,000 for the sale of the ANDA and related equipment net of expenses related to the sale.

In connection with the agreement, the Company agreed to manufacture Tablets for Mallinckrodt for a period of three years through March 31, 1998 and Mallinckrodt agreed to order a minimum number of Tablets from the Company for two years ending March 21, 1997. The Company and Mallinckrodt entered into a non competition agreement pursuant to which the Company agreed not to compete with Mallinckrodt and its affiliates with respect to the Tablets ANDA until March 21, 2000. If, prior to the time it is possible for Mallinckrodt to commence production under the Tablets ANDA or any new Tablets ANDA at its own facility, and the

10

Company ceases or is forced to cease or substantially curtail production under the Tablets ANDA, as a consequence of (i) any action or communication by the FDA or any other regulatory or governmental authority or (ii) any financial or other business difficulty, then Mallinckrodt has the right to cancel payment of any yet unpaid portion of the Deferred Payment (\$1.9 million) and shall further have the right to a full refund of any portion of the Deferred Payment already made to the Company.

In addition, the Company issued to Mallinckrodt the option to purchase the ANDA for oxycodone/acetaminophen capsules at an exercise price equal to 3/4 of annual net capsule revenue, as defined. Upon exercise of the option, the Company and Mallinckrodt would enter into agreements pursuant to which the Company would (i) manufacture oxycodone/acetaminophen capsules for Mallinckrodt for a period of time and (ii) be prohibited from competing with Mallinckrodt and its affiliates with respect to the production of capsules.

The Company has revised the gain recorded on the sale of assets to Mallinckrodt and will not recognize the Deferred Payment until the earlier of (i) Mallinckrodt receiving certain authorizations from the FDA or (ii) March 31, 1998. The effect of the adjustments on the accompanying financial statements is as follows (in thousand, except for per share amounts):

As of March 31, 1995:

	As Previously recorded	As Restated
Net earnings	\$ 3,731,000	\$1,831,000
Net earnings per common share	\$.48	\$.23
Long Term Receivable	1,900,000	-----
Accumulated Deficit	(7,155,000)	(9,055,000)

NOTE 5 - Sale of Common Stock

On March 30, 1995, the Company entered into an Agreement with Zatpack for the purchase of 500,000 shares of common stock of the Company by Zatpack, with registration rights, in consideration of \$1,000,000. The \$1,000,000 consideration consists of the cancellation of indebtedness (incurred by the Company's subsidiaries for the purchase of raw materials delivered or being delivered from affiliates of Zuellig) and shares of Indiana Fine Chemicals Corporation. As a result of the above transaction, the Company owns 100% of Indiana Fine Chemical Corporation. In addition, the Company issued a convertible promissory note to Zatpack, dated December 1, 1994 (Note E). Zatpack has acquired the above assets from Zuellig and its subsidiaries.

NOTE 6 - Contingencies

The Company currently is a defendant in several lawsuits involving product liability claims. The Company's insurance carriers have assumed the defense for all of these actions. Management is of the opinion that final disposition of these lawsuits will not result in a material adverse impact on the financial condition of the Company.

11

On June 29, 1993, the Company entered into a consent decree with the U.S. Attorney for the Eastern District of New York on behalf of the FDA that resulted from the FDA's investigation. Under the terms of the consent decree, the Company was enjoined from shipping any solid dosage drug products, (i.e.,

excluding liquid drug formulations) manufactured at the Company's facilities until the Company established, to the satisfaction of the FDA, that the methods used in, and the facilities and controls to be used for, manufacturing, processing, packing, labeling and holding any drug are established, operated and administered in conformity with the Federal Food, Drug and Cosmetic Act and the FDA's Current Good Manufacturing Practice regulations. As part of satisfying the foregoing requirements, the Company is required to validate the manufacturing process for each solid dosage drug product prior to manufacturing consent decree to manufacture and ship from its facilities six identified drug products at its own risk provided that (i) at least twice per month, the Company's independent expert certifies that each batch of drug products upon validation will have been manufactured in accordance with the FDA Regulations and the formulation described in the drug products approved NDA ("New Drug Application") or ANDA, until such time as validation is completed for these products; and (ii) for any batches of these products that have already been manufactured, such certification will include certification by a Company representative with personal knowledge of the records relating to such drug that they are accurate and complete and a certification signed by an independent expert that he has personally reviewed the records provided and that in his professional opinion, the foregoing requirement concerning validation has been met. The Company commenced shipments of five of the six solid dosage products under the foregoing certification process. After review by the Company and its consultants one of the Company's six core products, hydrocodone bitartrate 5mg and acetaminophen 500 mg tablet, discrepancies were discovered with some of the data in the Company's ANDA. This resulted in a voluntary recall of this product and the withdrawal of the ANDA.

On June 21, 1993, the Company entered into a plea agreement with the Department of Justice to resolve the government's investigation. Under the terms of the plea agreement, the Company agreed to resolve the government's investigation. Under the terms of the plea agreement, the Company agreed to plead guilty to five counts of adulteration of a single drug product shipped in interstate commerce and related recordkeeping violation. The plea agreement also requires the Company to pay fine of \$2,500,000 over five years in quarterly installments of \$125,000 beginning September 15, 1993. As of March 31, 1995, the Company has paid two quarterly installments and several partial payments. The plea agreement stipulated that if the Company does not make timely payments, the entire fine becomes due and payable. As a result, the entire Department of Justice Settlement has been reclassified as a current liability in the 1994 consolidated balance sheets. At the present time, no action has been initiated by the Department of Justice to require payment of the entire amount.

On March 31, 1993, and April 1, 1993, five lawsuits were filed by shareholders against the Company and three or more of the Company's directors. In June 1994, the plaintiffs of the five lawsuits and the two shareholder-derivative lawsuits and the Company agreed to a settlement of these lawsuits. The Company agreed to pay to the plaintiffs \$1,000,000 in cash, which has been paid by the Company's insurance carrier and, at the Company's option, either (i) the issuance of shares of the Company's common stock having a value, as of the date of the distribution of \$3,000,000 or (ii) the payment by the Company of \$3,000,000 in cash or (iii) any combination of issuance of shares or payment by the Company having a combined value as of date of distribution of \$3,000,000. The remainder of the settlement, \$3,000,000, will be paid by the Company after court approval is obtained by plaintiffs attorney for distribution.

On November 12, 1993, the Securities and Exchange Commission ("SEC") requested that the Company provide to the SEC, on a voluntary basis, information and documents regarding the ingredients and filings relating to the following drugs; quinidine gluconate, propylthiouracil, acetaminophen and codeine phosphate, metronidazole, quinidine sulfate, and hydralazine hydrochloride. The SEC advised the Company that the inquiry relates to public information disseminated by the Company and trading in the Company's securities during the period August 1987 through July 1993. The Company is cooperating with the SEC and has made available various documents. These documents relate to the testing, formulations and sale of these drugs which were maintained by the Company at the offices of its counsel in Maryland. In April 1994, the SEC requested additional documentation regarding these matters. The Company has complied with the additional request. On July 5, 1994, the Company made formal submission to the SEC and outlined the parameters of a proposed settlement. An additional submission was made on January 31, 1995 to bring additional information to the SEC. The Company is unable to predict the likelihood of an

unfavorable outcome as a result of this inquiry and, accordingly, no provision has been made for any potential costs.

A lawsuit has been filed by the minority shareholders of H. R. Cenci Laboratories, Inc. ("Cenci") and Cenci Powder Products, Inc. against the Company and several of the officers of the Company. The lawsuit alleges that the Company has breached several representations made during the course of negotiations leading to the Company's purchase of 51% of the stock of Cenci. This action seeks unspecified compensatory damages, as well as punitive damages, rescission, specific performance, reformation and a declaration as to what amount, if any is owed to plaintiff. Because of the early stage of this action, it is not possible at this time to predict with reasonable certainty the ultimate outcome of this matter and, accordingly, no provision has been made for any potential costs relating to this matter.

HALSEY DRUG CO., INC. AND SUBSIDIARIES

I

ITEM 2. MANAGEMENT'S DISCUSSION AND ANALYSIS OF FINANCIAL CONDITION AND RESULTS OF OPERATIONS

	Percentage of Net Sales		Percentage Change
			Year-to-Year
			Increase (decrease)
	-----		-----
	Three months ended March	Three months ended	
	-----		-----
	31	March 31	
	--	-----	
			1995
	1995	1994	to
	----	----	1994
	%	%	----
	-	-	-
Net Sales	100.0	100.0	6.2
Cost of Goods	74.4	78.3	10.8
Gross Profit	25.6	21.7	(10.6)
Research & Development	2.3	.4	(387.5)
Selling, General and administrative expenses	22.4	23.3	10.0
Earnings(loss) from operations	.9	(2.0)	143.3
Gain on the sale of assets	33.3		100.0
Interest expense	(3.3)	(2.2)	26.1
Earnings before income taxes	30.9	(4.2)	786.2
Provision for income taxes	4.3		
Net earnings (loss)	26.6	(4.2)	690.6
	====	=====	=====

Net Sales

The Company's net sales for the three months ended March 31, 1995 of \$6,873,000 represents a decrease of \$451,000 (6.2%) as compared to net sales for the three months ended March 31, 1994 of \$7,324,000. The decrease in 1995 is primarily attributable to the discontinuance of shipments of liquid products subsequent to the second quarter of 1994.

Cost of Goods Sold

For the three months ended March 31, 1995, cost of goods sold decreased by approximately \$619,000 as compared to the three months ended March 31, 1994. The decrease for 1994 is primarily attributable to the suspension of shipments of certain liquid products of Cenci as a result of the Company's review of operations at this location. In an effort to reduce the loss from lower revenues at this subsidiary, the Company has reduced its operating costs through significant reductions in personnel and other expenses at Cenci. The Company's gross margin as a percentage of sales for the three months ended March 31, 1995 was 25.6% as compared to 21.7% for the three months ended March 31, 1994.

Selling, General and Administrative Expenses

Selling, general and administrative expenses as a percentage of sales for the three months ended March 31, 1995 and 1994 were 22.4% and 21.7%, respectively. These expenses decreased by approximately \$171,000 or 10.0% as compared to 1994. The decrease was attributable to cost saving measures effected by management during the year, combined with a reduction in freight costs associated with the reduced sales volume at the subsidiary location.

Gain on Sale of Assets

On March 21, 1995, the Company sold its abbreviated new drug applications ("ANDA") for 5mg Oxycodone HCL/325mg and Acetaminophen Tablets ("Tablets") and certain equipment used in the production of Tablets for up to \$5.4 million to Mallinckrodt. The Company received \$ 500,000 of the proceeds in July 1994, which was recorded as deferred income on the Company's December 31, 1994 consolidated balance sheet. Mallinckrodt also paid the Company \$2,000,000 on March 21, 1995 and the remainder will be payable as follows: (i) \$1,000,000 upon the Company receiving general clearance from the FDA that it is in compliance with certain provisions of the consent decree dated June 29, 1993, and (ii) \$1,900,000 at the earlier of (a) Mallinckrodt receiving certain authorizations from the FDA or (b) September 21, 1997 ("Deferred Payments"). Mallinckrodt also agreed to defer \$1,200,000 of the Company's trade debt due to an affiliate of Mallinckrodt. For the three months ended March 31, 1995, the Company has recorded a gain of \$2,288,000 for the sale of the ANDA and related equipment net of expenses related to the sale. If, prior to the time it is possible for Mallinckrodt to commence production under the Tablets ANDA or any new Tablets ANDA at its own facility, and the Company ceases or is forced to cease or substantially curtail production under the Tablets ANDA, as a consequence of (i) any action or business difficulty, then Mallinckrodt has the right to cancel payment of any yet unpaid portion of the Deferred Payment (\$1.9 million) and shall further have the right to a full refund of any portion of the Deferred Payment already made to the Company.

Interest Expense

Interest expense for 1995 increased by \$65,000 as compared to 1994 as a result of an increase in the prime rate during 1994.

(Benefit) Provision for Income Taxes

The Company had a tax provision of \$296,000 as a result of available net operating loss carryforward. In 1994, the Company had no tax benefit since the available loss carryback to prior years was utilized by the net operating loss for 1993 carryback to the prior three years.

Net Earnings (Loss)

For the three months ended March 31, 1995, the Company had net earnings of \$1,831,000 as compared to a net loss of \$310,000 for the three months ended March 31, 1994. This increase is primarily attributable to the gain on the sale of assets in the amount of \$2,288,000.

Liquidity and Capital Resources

At March 31, 1995, the Company had cash and cash equivalents of \$172,000 as compared to \$27,000 at March 31, 1994. The Company had a working capital deficiency at March 31, 1995 of \$1,012,000 and \$4,451,000 at December 31, 1994.

As a result of the decline in shipments of solid dosage products from the Company's Brooklyn plant following the entry of the consent decree, and as a result of the lack of available borrowing under the Company's credit agreement, the Company's liquidity position has been materially adversely affected since June 30, 1993 and the Company's capital resources have been severely limited. The Company has actively sought to reduce its operating costs at the Brooklyn plant, where it has made significant reductions in personnel at the Brooklyn plant. In addition, the Company's liquidity position has been affected during the second half of 1994 by the discontinuance of shipments of liquid products from its Cenci subsidiary as a result of review completed by the Company of this liquid operation. In an effort to reduce the loss from lower revenues at this subsidiary, the Company has reduced its operating costs at Cenci through significant reductions of personnel and other expenses.

Under the terms of the plea agreement with the DOJ, the Company has agreed to pay a \$2,500,000 fine, payable in quarterly installments of \$125,000 over five years. Two installments have been paid to date. Only additional partial payments have been paid in the amount of \$30,000. The agreement with the DOJ stipulates if any payments are not made in a timely fashion, the entire amount of the fine shall become due and payable immediately. As a result, the entire amount of the settlement has been classified as current as of December 31, 1994. As of the current date, no action has been initiated to require immediate payment of the entire amount; however, the Company has recently made several partial payments.

In May 1994, the Company and its banks amended the credit agreement to (i) modify the terms of the warrants by adjusting the initial exercise price per share of the warrants to \$2.875; (ii) require the payment of any income tax refunds of the Company and its subsidiaries to an escrow account maintained by a designated agent; (iii) require the maintenance of a consultant for designated duties specified in the agreement; (iv) restrict certain payments made by the Company

or its subsidiaries; and (v) require the reimbursement of certain fees incurred by the banks in connection with the credit agreement.

In July 1994, the Company and its banks further amended the credit agreement to extend the due date to December 31, 1994, to modify certain financial covenants, to restrict the use of proceeds of loans and advances received by the Company including the receipts from the agreement with Mallinckrodt and to require the reimbursement of certain fees to the bank in connection with this agreement. As consideration, the Company issued 77,988 new warrants, at an exercise price of \$3.4375 per share, and agreed to issue additional warrants, for each month the loan remains outstanding through the due date of December 31, 1994. The Company has issued warrants for the purchase of an aggregate of 203,939 shares at exercise prices varying from \$2.875 to \$2.25 per share. Such warrants were valued at \$100,000 in 1993 and \$100,000 in 1994. The fair value of the warrants, \$200,000, as determined by the Company's Board of Directors, has been recorded by the Company as additional paid-in-capital and a discount to bank debt which is being amortized through the extended maturity date of the credit agreement, which is August 31, 1995.

In July 1994, the Company received an income tax refund of \$470,000, net of penalties and interest, which the Company used to reduce the outstanding debt and to pay interest and fees outstanding to the banks.

In March 1995, the Company and its banks restructured the credit agreement to include an extension of the due date to August 31, 1995, modification of the financial covenants, reduction of the exercise prices of all warrants in excess of \$2 3/8 per share and extension of the expiration date of the warrants to December 1999. As consideration for these modifications, the banks received \$1,500,000 of the proceeds received from the transaction with Mallinckrodt. Funds have been applied to reduce outstanding principal by approximately \$1,113,000 to approximately \$3,777,000, to pay accrued interest (approximately \$154,000) and fees (approximately \$233,000).

On March 30, 1995, the Company signed the Zatpack Agreement with Zatpack which provides for the purchase of 500,000 shares of common stock of the Company by Zatpack in consideration of \$1,000,000 (\$982,000 net of expenses).

As previously indicated, the Company has continued to actively pursue financing. At the current time, the Company is discussing with several parties obtaining financing which will replace the Company's banks and provide additional working capital. There can be no assurance that the Company will be able to obtain any such financing on commercially acceptable terms.

17

PART II OTHER INFORMATION

Item 6. Exhibits and Reports on Form 8-K

- (a) Exhibits:
Financial Data Schedule
- (b) Reports on Form 8-K
Current report on Form 8-K dated as of
March 21, 1995: Item 5 - Other Events; and
Item 7 - Financial Statements

18

SIGNATURES

Pursuant to the requirements 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.

HALSEY DRUG CO., INC.

Date: October 3, 1996

BY: /s/ Rosendo Ferran

Rosendo Ferran
President and Chief
Executive Officer

Date: October 3, 1996

BY: /s/ Robert J. Mellage

Robert J. Mellage
Corporate Controller

<ARTICLE> 5

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This schedule contains summary financial information extracted from the Condensed Consolidated Statement of Financial Condition At March 31, 1995 (Unaudited) and the Condensed Consolidated Statement of Income for the Three Months Ended March 31, 1995 (Unaudited) and is qualified in its entirety by reference to such financial statements.

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