Financial Results for the Year Ended March 31, 2015 <Supplement>

As of May 8, 2015 Mitsubishi Tanabe Pharma Corporation



Mitsubishi Tanabe Pharma

Table of Contents

2. Summary of Forecasts for FY2015 3. Dividends	
Consolidated Financial Indicators for FY2014	
. Profit and Loss	
(1) Profit anf Loss (2) Cost of Sales and SG&A Expenses	
(3) Non-operating Income and Loss (4) Extraordinary Income and Loss (5) Taxes	
(6) Sales of Main Products	
P. Financial Statement	
(1) Balance Sheet	
(2) Cash Flow Statement	
(3) Investment in Property, Plant and Equipment and Investment in Development of Information	
Systems (4) Depreciation Costs	
5. Financial Data & Employee Numbers of Major Consolidated Subsidiaries	
Forecasts for FY2015 Ending March 31, 2016	
(1) Consolidated Forecasts of Profit and Loss (2) Sales Forecasts by Segments	
(3) Forecasts of Cost of Sales and SG&A Expenses	
(4) Sales Forecasts for Main Products	
 (5) Forecast for Investment in Property, Plant and Equipment and Information Systems (6) Forecasts for Depreciation Costs 	
(6) Forecasts for Depreciation Costs	
Five-Year Financial Data	
(1) Profit and Loss (2) Balance Sheet (3) Other Financial Data (4) Number of Employees	
Quarterly Trend	
(1) Profit and Loss	
(1) Profit and Loss (2) Sales of Main Products	
(2) Sales of Main Products	
(2) Sales of Main Products State of New Product Development (as of May 8, 2015)	
(2) Sales of Main Products State of New Product Development (as of May 8, 2015) . New Drugs	
(2) Sales of Main Products State of New Product Development (as of May 8, 2015) . New Drugs . Additional Indications	
(2) Sales of Main Products State of New Product Development (as of May 8, 2015) . New Drugs . Additional Indications b. Licensing-out	···· ···
 (2) Sales of Main Products State of New Product Development (as of May 8, 2015) New Drugs Additional Indications Licensing-out Changes Since Previous Announcement on February 2, 2015 	
 (2) Sales of Main Products State of New Product Development (as of May 8, 2015) New Drugs Additional Indications Licensing-out Changes Since Previous Announcement on February 2, 2015 (1) In-house Development (2) Licensing-out 	···· ···
 (2) Sales of Main Products State of New Product Development (as of May 8, 2015) New Drugs Additional Indications Licensing-out Changes Since Previous Announcement on February 2, 2015 (1) In-house Development (2) Licensing-out Additional Information for State of New Product Development 	···· ··· ··· ··· ··· ···
 (2) Sales of Main Products State of New Product Development (as of May 8, 2015) New Drugs Additional Indications Licensing-out Changes Since Previous Announcement on February 2, 2015 (1) In-house Development (2) Licensing-out Additional Information for State of New Product Development (1) New Drugs 	
 (2) Sales of Main Products State of New Product Development (as of May 8, 2015) New Drugs Additional Indications Licensing-out Changes Since Previous Announcement on February 2, 2015 (1) In-house Development (2) Licensing-out Additional Information for State of New Product Development (1) New Drugs (2) Additional Indications 	···· ··· ··· ··· ··· ···
 (2) Sales of Main Products State of New Product Development (as of May 8, 2015) New Drugs Additional Indications Licensing-out Changes Since Previous Announcement on February 2, 2015 (1) In-house Development (2) Licensing-out Additional Information for State of New Product Development (1) New Drugs (2) Additional Indications (3) Licensing-out 	
 (2) Sales of Main Products State of New Product Development (as of May 8, 2015) New Drugs Additional Indications Licensing-out Changes Since Previous Announcement on February 2, 2015 (1) In-house Development (2) Licensing-out Additional Information for State of New Product Development (1) New Drugs (2) Additional Indications (3) Licensing-out Others 	
 (2) Sales of Main Products State of New Product Development (as of May 8, 2015) New Drugs Additional Indications Licensing-out Changes Since Previous Announcement on February 2, 2015 (1) In-house Development (2) Licensing-out Additional Information for State of New Product Development (1) New Drugs (2) Additional Indications (3) Licensing-out Others Subsidiaries and Affiliated Companies 	
 (2) Sales of Main Products State of New Product Development (as of May 8, 2015) New Drugs Additional Indications Licensing-out Changes Since Previous Announcement on February 2, 2015 (1) In-house Development (2) Licensing-out Additional Information for State of New Product Development (1) New Drugs (2) Additional Indications (3) Licensing-out Others Subsidiaries and Affiliated Companies (1) Number of Subsidiaries and Affiliated Companies (2) Consolidated Subsidiaries 	
 (2) Sales of Main Products State of New Product Development (as of May 8, 2015) New Drugs Additional Indications Licensing-out Changes Since Previous Announcement on February 2, 2015 (1) In-house Development (2) Licensing-out Additional Information for State of New Product Development (1) New Drugs (2) Additional Indications (3) Licensing-out Others Subsidiaries and Affiliated Companies (1) Number of Subsidiaries and Affiliated Companies (2) Consolidated Subsidiaries (3) Affiliated Companies Accounted for by the Equity Method 	
 (2) Sales of Main Products State of New Product Development (as of May 8, 2015) New Drugs Additional Indications Licensing-out Changes Since Previous Announcement on February 2, 2015 (1) In-house Development (2) Licensing-out Additional Information for State of New Product Development (1) New Drugs (2) Additional Indications (3) Licensing-out Others Subsidiaries and Affiliated Companies (1) Number of Subsidiaries and Affiliated Companies (2) Consolidated Subsidiaries (3) Affiliated Companies Accounted for by the Equity Method Status of Shareholders 	
 (2) Sales of Main Products State of New Product Development (as of May 8, 2015) New Drugs Additional Indications Licensing-out Changes Since Previous Announcement on February 2, 2015 (1) In-house Development (2) Licensing-out Additional Information for State of New Product Development (1) New Drugs (2) Additional Indications (3) Licensing-out Others Subsidiaries and Affiliated Companies (1) Number of Subsidiaries and Affiliated Companies (2) Consolidated Subsidiaries (3) Affiliated Companies Accounted for by the Equity Method Status of Shareholders (1) Number of Outstanding Shares 	
 (2) Sales of Main Products State of New Product Development (as of May 8, 2015) New Drugs Additional Indications Licensing-out Changes Since Previous Announcement on February 2, 2015 (1) In-house Development (2) Licensing-out Additional Information for State of New Product Development (1) New Drugs (2) Additional Indications (3) Licensing-out Others Subsidiaries and Affiliated Companies (1) Number of Subsidiaries and Affiliated Companies (2) Consolidated Subsidiaries (3) Affiliated Companies Accounted for by the Equity Method Status of Shareholders (1) Number of Outstanding Shares (2) Status of Major Shareholders (3) Ownership and Distribution of Shares 	
 (2) Sales of Main Products State of New Product Development (as of May 8, 2015) New Drugs Additional Indications Licensing-out Changes Since Previous Announcement on February 2, 2015 (1) In-house Development (2) Licensing-out Additional Information for State of New Product Development (1) New Drugs (2) Additional Indications (3) Licensing-out Others Subsidiaries and Affiliated Companies (1) Number of Subsidiaries and Affiliated Companies (2) Consolidated Subsidiaries (3) Affiliated Companies Accounted for by the Equity Method Status of Shareholders (1) Number of Outstanding Shares 	

Summary of Financial Results for FY2014 Ended March 31, 2015 and Forecasts for FY2015

(Amounts less than ¥100 million are rounded.)

1. Summary of Financial Results for FY2014

				[Billion yen]
Net Sales	415.1	Y-on-Y	2.4	0.6 %
Domestic sales	337.2	Y-on-Y	(16.1)	(4.6 %)
Overseas sales	77.9	Y-on-Y	18.6	31.3 %

Net sales increased 0.6%, or ¥2.4 billion, to ¥415.1 billion.

In the domestic sales of ethical drugs, favorable sales growth was recorded by Remicade, an anti-TNF α monoclonal antibody and TENELIA, for the treatment of type2 diabetes mellitus. However, there were the growing impact of generics and NHI price revision in April 2014. As a result, the domestic sales of ethical drugs decreased 5.2%, year-on-year, to ¥323.9 billion.

Overseas sales of ethical drugs were ¥23.0 billion, up 4.6%, year-on-year, due to depreciation of the yen.

Royalty income, etc. increased 60.7%, year-on-year, to ¥60.4 billion due to the increase in royalty revenue from Gilenya, for the treatment of multiple sclerosis, licensed to Novartis and from INVOKANA and the fixed dose combination with metformin (IR), for the treatment of type2 diabetes mellitus, licensed to Janssen Pharmaceuticals.

Operating Income	67.1	Y-on-Y	8.0	13.6 %
------------------	------	--------	-----	--------

Operating income was ¥67.1 billion, up 13.6%, or ¥8.0 billion, year-on-year.

Despite the influence of NHI drug price revision, gross profit increased ¥2.2 billion, year-on-year, to ¥245.5 billion due to the increase in royalty revenue. As a result, the cost of sales ratio improved by 0.1 percentage points, year-on-year, to 40.9%.

SG&A expenses decreased ¥5.8 billion, year-on-year, to ¥178.4 billion due to the decrease in the labor cost accompanying the decrease in retirement benefit expenses and R&D expenses related to the progress of development phase.

				[Billion yen]
Ordinary Income	67.7	Y-on-Y	5.8	9.3 %
Net Income	39.5	Y-on-Y	(5.9)	(13.0 %)

Ordinary income was up 9.3%, or ¥5.8 billion, year-on-year, to ¥67.7 billion, and net income was down 13.0%, or ¥5.9 billion, year-on-year, to ¥39.5 billion.

Foreign exchange gain decreased to ¥0.4 billion (foreign exchange gain was ¥2.5 billion in the previous fiscal year). As a result, non-operating income and loss worsened by ¥2.2 billion, year-on-year.

Extraordinary income was ¥13.7 billion, mainly because the Company recorded gain on sales of property, plant and equipment, such as sales of former Nihonbashi Building. In the previous fiscal year, the Company recorded extraordinary income of ¥15.3 billion, such as profit on arbitration award.

Extraordinary loss was ¥18.6 billion, including restructuring expenses, such as sales of Kashima Plant and the closing of Kazusa office, related to one of the strategic challenges of Medium-Term Management Plan; "accelerating operational and structural reforms." In the previous fiscal year, the Company recorded extraordinary loss of ¥4.8 billion, such as special retirement expenses.

2. Summary of Forecasts for FY2014

				[Billion yen]
Net Sales	396.0	Y-on-Y	(19.1)	(4.6 %)
Operating Income	67.5	Y-on-Y	0.4	0.5 %
Ordinary Income	67.0	Y-on-Y	(0.7)	(1.0 %)
Net Income	40.5	Y-on-Y	1.0	2.5 %

3. Dividends

	FY2015 (Estimate)	FY20	14
	End of 1st Half	For the Year	End of 1st Half	For the Year
Dividends per Share (¥)	22	44	20	42
Dividends Payout Ratio	-	60.9%	-	59.6%
prior to amortization of goodwill	-	48.8%	-	47.6%

2 Consolidated Financial Indicators for FY2014

1. Profit and Loss

(Amounts less than ¥100 million are rounded.)

(1) P	(1) Profit and Loss									
	Y-on-Y Comparison to Previous Forecasts									
		FY2014	FY2013	Increase (Decrease)	Change %	Forecast*	Increase (Decrease)	Change %	Notes [Y-on-Y Comparison]	
Net	sales	415.1	412.7	2.4	0.6	406.0	9.1	2.2	See "Sales of Main Products" on page 5. Overseas sales ratio FY2013: 14.4%	
	Domestic sales	337.2	353.3	(16.1)	(4.6)	335.7	1.5	0.4	FY2013. 14.4% FY2014: 18.8% Average exchange rate	
	Overseas sales	77.9	59.4	18.6	31.3	70.3	7.6	10.9	FY2013:1US\$=¥100.49 FY2014: 1US\$=¥110.62	
С	ost of sales	169.6	169.4	0.2	0.1	164.0	5.6	3.4		
	Sales cost ratio	40.9%	41.0%			40.4%				
Gro	ss operation profit	245.5	243.3	2.2	0.9	242.0	3.5	1.5		
S	G&A expenses	178.4	184.2	(5.8)	(3.2)	182.0	(3.6)	(2.0)		
	% of net sales	43.0%	44.6%			44.8%				
Оре	erating income	67.1	59.1	8.0	13.6	60.0	7.1	11.9		
Ord	inary income	67.7	61.9	5.8	9.3	61.5	6.2	10.0		
Extra	ordinary income and loss	(5.0)	10.6	(15.5)	-	0.0	(5.0)	-		
Net	income	39.5	45.4	(5.9)	(13.0)	40.5	(1.0)	(2.5)		

(2) Cost of Sales and Selling, General and Administrative Expenses

[Billion yen]

		Y-on-Y		Comparison to Previous Forecasts				
	FY2014	FY2013	Increase (Decrease)	Change %	Forecast*	Increase (Decrease)	Change %	Notes [Y-on-Y Comparison]
Cost of sales	169.6	169.4	0.2	0.1	164.0	5.6	3.4	
% of Net sales	40.9%	41.0%			40.4%			
SG&A expenses % of Net sales	178.4 43.0%	184.2 44.6%	(5.8)	(3.2)	182.0 44.8%	(3.6)	(2.0)	
R&D expenses % of Net sales	69.6 16.8%	70.4 17.1%	```	(1.1)	72.5 17.9%	(2.9)	(4.0)	
Except R&D expenses	108.8	113.8	(5.0)	(4.4)	109.5	(0.7)	(0.7)	
Labor cost	46.8	48.4	(1.6)	(3.3)	46.2	0.6	1.2	Decrease in retirement benefit expenses, etc.
Amortization of goodwill	10.9	10.6	0.3	2.6	10.8	0.1	1.1	
Others	51.1	54.8	(3.7)	(6.8)	52.5	(1.4)	(2.7)	
Total labor cost	81.1	85.0	()	. ,		1.1	1.3	

*: Published forecasts announced on October 29, 2014 in the financial results of Q2 FY2014

(3) Non-operating Income and Loss

(3) N	(3) Non-operating Income and Loss							
		FY2014	FY2013	Increase (Decrease)	Notes			
Non-	operating income	3.8	6.9	(3.1)				
Ir	nterest income	1.6	1.5	0.1				
D	Dividend income	0.8	0.8	(0.1)				
E	equity in earnings of affiliates	0.0	0.6	(0.6)				
F	oreign exchang gain	0.4	2.5	(2.1)				
R	Rent income	0.2	0.3	(0.1)				
С	Others	0.8	1.0	(0.3)				
Noi	n-operating expenses	3.2	4.1	(0.9)				
Ir	nterest expense	0.2	0.1	0.1				
	djustment for salaries for employees n secondment	0.1	0.8	(0.7)				
D	Oonations	1.5	0.7	0.9				
С	Others	1.4	2.6	(1.2)				

(4) Extraordinary Income and Loss

[Billion yen]

				[2			
	FY2014	FY2013	Increase (Decrease)	Notes			
Extraordinary income	13.7	15.3	(1.7)				
Gain on sales of property, plant and equipment	12.0	1.0	11.0	Gain on the sale of a vacant lot of the former Nihonbashi building			
Gain on sales of investment in securities	1.1	2.4	(1.3)				
Gain on sales of shares of subsidiaries and affiliates	0.6	-	0.6	Gain on the sales of shares of API Corporation and CMIC CMO, Ashikaga			
Profit on arbitration award	-	11.0	(11.0)	FY2013: Reimbursed as the overpayment caused by the arbitration award of Remicade, etc			
Gain on step acquisitions	-	0.9	(0.9)	FY2013: Gain on market value of stock holdings accompanied with acquisition of Medicago's shares			
Extraordinary Loss	18.6	4.8	13.9				
Restructuring expenses	12.3	-	12.3	Close of Kazusa office, transfer of Kashima plant (Ibraki), withdrawal of business of Mitsubishi Pharma (Guangzhou) and losses according to relocation of head office and Kashima office (Osaka)			
Amortization of goodwill	3.5	-	3.5	One-time amortization of goodwill of Bipha			
Impairment loss	2.6	1.4	1.2	FY2014: Toda dormitory, etc. FY2013: Former research center in Yoshitomi, etc.			
Loss on valuation of investment in securities	0.1	0.6	(0.5)				
Loss on sale of investments in securities	0.1	0.0	0.1				
Special retirement expenses	-	2.6	(2.6)	FY2013: Premium retirement expenses accroding to transfer to Japan Blood Products Organization and CMIC CMO, Ashikaga, etc.			
Other	0.1	0.2	(0.1)				

(5) Taxes

(5) Taxes				[Billion yen]
	FY2014	FY2013	Increase (Decrease)	Notes
Income before income taxes and			(20010000)	
minority interests	62.7	72.4	(9.8)	
Income taxes-current	29.8	22.4	7.4	
Income taxes-deferred	(4.4)	4.7	(9.1)	
Minority interests	(2.2)	0.0	(2.2)	
Net Income	39.5	45.4	(5.9)	

(6) Sales of Main Products		[Billion yer					
			Y-on-Y		Comparison to Forecasts		
	FY2014	FY2013	Increase (Decrease)	Change %	Forecasts *1	Increase (Decrease)	Change %
Ethical drugs	410.7	407.2	3.5	0.9	401.3	9.4	2.3
Ethical drugs domestic sales	323.9	341.7	(17.8)	(5.2)	322.5	1.4	0.4
Remicade	70.6	76.3	(5.7)	(7.5)	70.3	0.3	0.4
Talion	16.0	13.7	2.3	16.7	16.4	(0.4)	(2.4)
Ceredist	15.7	17.8	(2.1)	(11.9)	15.4	0.3	1.7
Maintate	14.1	15.5	(1.3)	(8.5)	14.5	(0.3)	(2.1)
Venoglobulin IH	11.6	11.1	0.5	4.5	11.5	0.2	1.3
Simponi	10.5	9.4	1.1	11.6	11.5	(1.1)	(9.4)
Kremezin	10.5	12.6	(2.0)	(16.1)	10.5	0.0	0.1
Urso	10.0	12.4	(2.4)	(19.5)	9.9	0.1	1.1
Anplag	8.3	11.2	(2.9)	(25.7)	8.4	(0.1)	(0.8)
Depas	8.1	9.8	(1.7)	(17.7)	8.1	0.0	(0.4)
Lexapro	8.0	6.5	1.5	23.4	8.0	0.0	(0.4)
Radicut	7.4	10.9	(3.6)	(32.5)	7.2	0.2	2.6
Tenelia	6.2	0.8	5.4	684.1	6.5	(0.2)	(3.7)
Herbesser	5.5	6.9	(1.3)	(19.5)	5.5	0.0	(0.2)
Tanatril	4.6	6.2	(1.5)	(25.1)	4.6	0.0	0.3
BIKEN products [vaccine]	30.3	28.4	1.9	6.5	28.2	2.1	7.3
Tetrabik	7.5	6.7	0.8	11.9	7.1	0.5	6.5
Influenza	7.4	7.2	0.2	2.3	7.6	(0.2)	(2.7)
Varicella vaccine	7.2	3.6	3.6	99.6	5.2	1.9	36.9
Tanabe Seiyaku Hanbai products *2	13.6	14.1	(0.5)	(3.2)	13.7	(0.1)	(0.7)
Ethical drugs overseas sales	23.0	22.0	1.0	4.6	22.0	1.1	4.9
Herbesser	6.5	5.8	0.7	12.1	6.0	0.5	7.7
Argatroban (Novastan)	2.1	2.7	(0.5)	(19.7)	2.2	(0.1)	(4.7)
Tanatril	1.8	1.8	0.0	1.2	2.1	(0.3)	(12.1)
Contracted manufacturing products *3	3.4	5.8	(2.5)	(42.2)	3.3	0.1	1.6
Royalty income, etc.	60.4	37.6	22.8	60.7	53.6	6.8	12.7
Royalty from Gilenya	43.9	32.2	11.8	36.7	Undisclosed	-	-
Royalty from INVOKANA	9.8	Undisclosed	-	-	Undisclosed	-	-
OTC products 4.0 4.5 (0.5) (10.5) 4.3				(0.3)	(6.7)		
Total pharmaceuticals	415.1	412.7	2.4	0.6	406.0	9.1	2.2

(6) Sales of Main Products

[Billion yen]

*1: Published forecasts announced on October 29, 2014 in the financial results of Q2 FY2014.

*2: Tanabe Seiyaku Hanbai Products are composed of generic drugs and the long-listed drugs which were transferred from MTPC.

 *3 :Active pharmaceutical ingredients and others ordered by other companies.

2. Financial Statement

	End of FY2014	Composition %	End of FY2013	Increase (Decrease)	Notes
otal Aseets	929.3	100.0	886.5	42.8	
Current Assets	603.6	65.0	540.5	63.2	
Cash and deposits	50.2	5.4	27.2	23.0	See Page 7, (2) Cash Flows Statement
Marketable securities	118.8	12.8	106.5	12.3	Increase in negotiable deposit, etc.
Notes and accounts receivable*1	130.3	14.0	123.5	6.8	
[Months/Revolution]	3.77		3.59	0.18	
Inventories	85.1	9.2	93.7	(8.6)	
Deposits	192.8	20.7	172.1	20.6	
Deferred income taxes	8.3	0.9	8.2	0.2	
Other	18.1	2.0	9.3	8.8	
Fixed Assets	325.7	35.0	346.0	(20.3)	
Property, plant and equipment	92.5	10.0	98.3	(5.8)	Investment for plant and equipment, 15.7; Depreciation, (7. Disposal, sale, impairment and others, (9.4), etc.
Intangible fixed assets	116.9	12.6	133.1	(16.2)	Investment for information system, 1.6; Depreciation, (1.6). Amortization of goodwill of the merger, (10.0), Amortization goodwill of Bipha, (3.8)etc.
Investment in securities	76.3	8.2	71.6	4.7	Increase of fair market value, decrease due to sales of shar API Corporation, etc.
Deferred income taxes	0.8	0.1	0.7	0.1	
Net defined benefit asset	15.7	1.7	16.3	(0.6)	
Other	23.4	2.5	26.0	(2.6)	
otal Liabilities	128.9	13.9	108.6	20.2	
Current Liabilities	105.4	11.3	81.8	23.6	
Notes and accounts payable*2 Short-term debt	34.6	3.7	34.0	0.6	
	-	-	1.2	(1.2)	
Current maturities of long-term debt	0.1	0.0	0.1	0.0	
Accounts payable, other	25.4	2.7	16.8	8.6	
Income taxes payable	19.8	2.1	10.2	9.6	
Reserve for employees' bonuses	10.0	1.1	10.2	(0.2)	
Other	15.5	1.7	9.4	6.2	
Long-term Liabilities	23.5	2.5	26.8	(3.3)	
Long-term debts	0.9	0.1	1.0	(0.1)	
Deferred income taxes	9.8	1.1	13.4	(3.6)	
Reserve for health management allowances for HIV compensation Reserve for health management	1.7	0.2	1.6	0.1	
allowances for SMON compensation	2.7	0.3	3.0	(0.2)	
Reserve for HCV litigation	2.0	0.2	2.6	(0.6)	
Net defined benefit liability	2.5	0.3	2.1	0.3	
Other	3.9	0.4	3.2	0.7	
let Assets	800.4	86.1	777.8	22.6	
Shareholders' equity	776.0	83.5	767.3	8.7	
Common stock	50.0	5.4	50.0	-	
Capital surplus	451.2	48.6	451.2	-	
Retained earnings	275.3	29.6	266.6	8.8	Net income, 39.5; Payment for dividends, (22.4)
Treasury stock, at cost	(0.5)	(0.1)	(0.5)	0.0	
Accumulated other comprehensive loss	13.0	1.4	(1.2)	14.2	
Unrealized holding (losses) gains on securities	14.9	1.6	8.7	6.2	
Deffered (losses) gains on hedges	0.1	0.0	0.5	(0.4)	
Translation adjustments	0.1	0.0	(2.4)	2.5	
Remeasurements of defined benefit plans	(2.2)	(0.2)	(8.1)	5.9	
Minority interests	11.5	1.2	11.8	(0.3)	

*1: Notes and accounts receivable = Bills + Accounts receivable

*2: Notes and account payable=Bills(except non-operating bills)+Accounts payable

2) Cash Flow Statement	[Billion yen			
	FY2014	FY2013	Increase (Decrease)	
Cash and cash equivalents at beginning of year	85.0	58.7	2	
Cash flows from operating activities	68.2	69.9	(*	
Income before income taxes and minority interests	62.7	72.4	(
Depreciation and amortization	9.0	9.1		
Impairment loss	2.6	1.4	·	
Amortization of goodwill	14.4	10.6		
Increase (decrease) in accrued retirement benefit for employees	-	(9.4)		
	(0.5)	7.9		
Increase (decrease) in net defined benefit liability	(0.0)			
Decrease (increase) in prepaid pension expenses	-	36.9	(3	
Decrease (increase) in net defined benefit asset	(3.9)	(34.5)		
Increase (decrease) in reserve for HCV litigation	(0.6)	(1.0)		
Interest and dividend income	(2.4)	(2.4)		
Loss (gain) on sales of shares of subsidiaries and affiliates	(0.6)	-		
Loss (gain) on sales and disposal of fixed assets	(11.8)	(0.7)	(*	
Restructuring expenses	12.3	-		
Profit on arbitration award	-	(11.0)		
Loss (gain) on step acquisitions	-	(0.9)		
Loss (gain) on sale of investment in securities	(1.0)	(2.4)		
Loss (gain) on valuation of investment in securities	0.1	0.6		
Equity in losses (earnings) of affiliates	0.0	(0.6)		
Decrease(increase) in notes and accounts receivable-trade	(6.7)	6.6	(
Decrease (increase) in inventories	7.8	(0.7)		
Decrease(increase) in notes and accounts payable, trade	0.5	(4.1)		
Increase(decrease) in accounts payable, other	5.9	0.8		
Interest and dividends received	2.4	3.5		
Proceeds from arbitration award	-	12.2	(
Income taxes paid	(20.0)	(28.1)		
Other, net	(2.1)	3.7		
ash flows from investing activities	(59.8)	(24.3)	(
Purchase/sales etc. of marketable securities	(26.4)	22.4	(*	
	(20.2)	(1.9)	(
Increase/decrease in time deposits	(20.6)	(1.3)	(
Increase in deposits		(20.7)		
Purchase/sales of property, plant and equipment	(1.3)	()		
Purchase of intangible fixed assets	(1.5)	(2.0)		
Purchase/sales of investment in securities	1.1	8.9		
Purchase of investment in subsidiaries	-	(3.7)		
Proceeds from sales of shares of subsidiaries and associates	7.6	-		
Purchase of investment in subsidiaries resulting in consolidation scope change	-	(17.9)		
Proceeds from sales of shares of subsidiaries resulting in change in scope of consolidation	1.5	-		
Other, net	0.0	(0.1)		
ash flows from financing activities	(21.9)	(21.1)		
Increase (decrease) in short-term debt, net	(1.2)	(0.2)		
Increase in long-term debt	-	1.0		
Proceeds from share issuance to minority shareholders	2.6	0.6		
Cash dividends paid	(22.4)	(22.4)		
Cash dividends paid to minority shareholders	(0.6)	0.0		
Other, net	(0.2)	(0.1)		
fect of exchange rate change on cash and cash equivalents	1.9	1.8		
et increase (decrease) in cash and cash equivalents	(11.6)	26.2	(3	
Cash and cash equivalents at end of the year	73.3	85.0	(1	

The Reconciliation of Cash and Cash Equivalents in the Consolidated Balance Sheets and Cash and Cash Equivalents in the Consolidated Statements of Cash Flows at the End of the Period [Billion yen]

	FY2014	FY2013
Cash and time deposits	50.2	27.2
Time deposits maturing after three months	(25.6)	(4.8)
Short-term investments in marketable securities maturing within three months of acquisition	28.0	42.0
Cash equivalents included in short-term loans receivable*	0.7	0.6
Cash equivalents included in deposits	20.0	20.0
Cash and cash equivalents in the consolidated statements of cash flows *: Short,term loans are included in "Others, Current Assets" on page 6	73.3	85.0

*: Short-term loans are included in "Others, Current Assets" on page 6.

(3) Investment in Property, Plant and Equipment and Investment in Development of Information Systems

			[Billion yen]
	FY2014	FY2013	Increase
	112011	112010	(Decrease)
Investment in property, plant and equipment / occuring basis	15.7	12.6	3.1
Investment in information systems/ occuring basis	1.6	2.1	(0.5)

		[B	illion yen]
Major investment in property, plant and equip	Major investment in property, plant and equipment		
in FY2014	in FY2014		
Mitsubishi Tanabe Pharma	9.3	Mitsubishi Tanabe Pharma	1.3
[Construction of new head office and Kashima office building]	[5.9]		
Mitsubishi Tanabe Pharma Factory	4.0		
[Construction of new manufacturing facility at Yoshitomi plant]	[1.8]		

(4) Depreciation Costs	[Billion yen]		
	FY2014	FY2013	Increase (Decrease)
Property, plant and equipment	7.5	7.9	(0.4)
Intangible fixed assets	1.6	1.3	0.3

3. Financial Data & Employee Numbers of Major Consolidated Subsidiaries

[Billion yen] Mitsubishi Mitsubishi Tanabe Seiyaku Tianjin Tanabe P.T. Tanabe Companies Tanabe Pharma Tanabe Pharma Medicago, Inc. Hanbai Co., Ltd. Seiyaku Co., Ltd. Indonesia Korea Co., Ltd. Factory Ltd. FY2014 39.4 13.6 4.4 1.1 4.5 2.3 Net sales FY2013 47.2 14.1 4.1 0.0 3.6 2.3 FY2014 3.6 (0.1) 0.2 (5.1) 0.5 0.1 Operating income FY2013 1.2 0.4 0.3 (1.3) 0.1 0.3 FY2014 3.6 (0.1) 0.3 (5.2) 0.5 0.1 Ordinary income FY2013 1.1 0.4 0.4 (1.2) 0.1 0.3 FY2014 0.7 (0.1) 0.2 (5.3) 0.3 0.0 Net income FY2013 0.7 0.3 0.3 (1.2) 0.0 0.2 FY2014 1.3 6.2 0.2 0.0 --R&D expenses FY2013 1.2 --1.4 0.0 0.0 FY2014 2.2 0.0 0.1 0.4 0.1 0.1 Depreciation of property, plant and equipment FY2013 2.4 0.0 0.1 0.1 0.1 0.1 End of FY2014 48.2 6.0 3.7 35.6 3.1 6.1 Total assets End of FY2013 57.6 6.3 3.3 36.5 4.4 3.6 End of FY2014 36.9 0.3 3.0 24.2 3.9 1.7 Net assets End of FY2013 39.8 0.5 24.1 3.0 1.6 2.6 End of FY2014 1,087 168 121 263 537 441 Number of employees End of FY2013 1,394 172 125 189 456 480

Note: Prior to elimination of internal transaction

3 Forecasts for FY2015 Ending March 31, 2016

(1) Consolidated Forecasts of Profit and Loss

(Amounts less than ¥100 million are rounded.)

h		1st Half of FY2015 Forecasts	1st Half of FY2014 Actual	Increase (Decrease)	Change %	FY2015 Forecasts	FY2014 Actual	Increase (Decrease)	Change %	Notes
Net	Sales	191.5	198.9	(7.4)	(3.7)	396.0	415.1	(19.1)	(4.6)	Overseas sales ratio FY2014: 18.8% FY2015 estimation:
	Domestic	144.3	165.0	(20.7)	(12.5)	301.5	337.2	(35.7)	(10.6)	
	Overseas	47.2	33.9	13.3	39.3	94.5	77.9	16.6	21.2	planned: 1US\$=¥120
	Cost of Sales	70.5	78.2	(7.7)	(9.8)	147.0	169.6	(22.6)	(13.3)	
	Sales cost ratio	36.8%	39.3%			37.1%	40.9%			
Gro	ss Operatin Profit	121.0	120.7	0.3	0.2	249.0	245.5	3.5	1.4	
SG	& A Expenses	93.0	85.8	7.2	8.5	181.5	178.4	3.1	1.7	
	% of Net Sales	48.6%	43.1%			45.8%	43.0%			
Оре	erating Income	28.0	35.0	(7.0)	(19.9)	67.5	67.1	0.4	0.5	
Ord	inary Income	28.0	35.5	(7.5)	(21.0)	67.0	67.7	(0.7)	(1.0)	
	Extraordinary Income and loss	0.0	10.9	(10.9)	-	(7.5)	(5.0)	(2.5)	-	
Net	Income	19.0	32.5	(13.5)	(41.6)	40.5	39.5	1.0	2.5	

(2) Forecasts of Cost of Sales and SG&A Expenses

1st Half of 1st Half of FY2015 FY2014 Increase Increase FY2015 FY2014 Change % Change % (Decrease) Forecasts Actual (Decrease) Forecasts Actual Cost of Sales 70.5 78.2 (7.7) (9.8) 147.0 169.6 (22.6)(13.3)Sales cost ratio 40.9% 36.8% 39.3% 37.1% SG & A Expenses 93.0 85.8 7.2 8.5 181.5 178.4 3.1 1.7 % of Net sales 48.6% 43.1% 45.8% 43.0% 7.6 23.7 74.0 R&D Expenses 39.5 31.9 69.6 4.4 6.3 % of Net sales 20.6% 18.7% 16.8% 16.1% Except R&D Expenses 53.5 53.8 (0.3) (0.6)107.5 108.8 (1.3)(1.2) 23.0 0.0 23.0 (0.1) 46.0 46.8 (0.8) (1.6)Labor Cost 5.3 5.4 (0.1) (1.8) 10.5 10.9 (0.4) (3.8) Amortization of Goodwill 25.2 25.4 (0.8) 51.0 51.1 (0.1) (0.2) Others (0.2) 38.2 39.8 (4.1)77.0 81.1 (4.1) (5.0)Total Labor Cost (1.6)

[Billion yen]

[Billion yen]

(3) Sales Forecasts for Main Products

[Billion	venl
	yenj

,								
	1st Half of FY2015 Forecasts	1st Half of FY2014 Actual	Increase (Decrease)	Change %	FY2015 Forecasts	FY2014 Actual	Increase (Decrease)	Change %
Ethical drugs	189.2	196.5	(7.3)	(3.7)	391.6	410.7	(19.1)	(4.6
Ethical drugs domestic sales	140.8	155.2	(14.4)	(9.3)	294.6	323.9	(29.3)	(9.0
Remicade	35.6	35.2	0.4	1.0	70.7	70.6	0.1	0.1
Talion	6.4	6.1	0.3	5.3	17.1	16.0	1.1	7.0
Ceredist	7.1	7.9	(0.8)	(10.1)	14.1	15.7	(1.6)	(10.0
Simponi	6.2	5.0	1.2	23.1	13.3	10.5	2.8	27.2
Maintate	6.6	7.2	(0.6)	(7.7)	13.2	14.1	(0.9)	(6.
Lexapro	4.1	3.4	0.7	20.2	10.5	8.0	2.5	31.6
Tenelia	4.5	2.7	1.8	68.8	9.6	6.2	3.4	54.2
Kremezin	4.8	5.4	(0.6)	(11.4)	9.3	10.5	(1.2)	(11.
Urso	4.6	5.1	(0.5)	(9.6)	8.8	10.0	(1.2)	(11.
Depas	3.4	4.1	(0.7)	(16.9)	6.8	8.1	(1.3)	(15.
Anplag	3.0	4.4	(1.4)	(31.1)	5.8	8.3	(2.5)	(30.
Radicut	2.9	3.8	(0.9)	(23.6)	5.4	7.4	(2.0)	(26.
BIKEN products [vaccine]	10.6	11.1	(0.5)	(4.5)	26.9	30.3	(3.4)	(11.
Influenza	0.6	0.9	(0.3)	(32.4)	7.9	7.4	0.5	7.
Tetrabik	3.4	3.6	(0.2)	(4.6)	7.1	7.5	(0.4)	(5.
Tanabe Seiyaku Hanbai products *1	6.9	6.4	0.5	7.1	14.4	13.6	0.8	5.
Ethical drugs overseas sales	13.4	11.3	2.1	18.2	24.4	23.0	1.4	5.
Herbesser	3.4	3.5	(0.1)	(1.8)	6.8	6.5	0.3	5.
Argatroban (Novastan)	1.0	1.3	(0.3)	(20.4)	2.5	2.1	0.4	17.
Tanatril	0.8	1.0	(0.2)	(22.3)	1.7	1.8	(0.1)	(7.
Contracted manufacturing products *2	1.1	2.1	(1.0)	(47.1)	2.4	3.4	(1.0)	(28.
Royalty income, etc.	33.9	27.9	6.0	21.7	70.2	60.4	9.8	16.
Royalty from Gilenya	Undisclosed	18.8	-	-	Undisclosed	43.9	-	
Royalty from INVOKANA	Undisclosed	Undisclosed	-	-	Undisclosed	9.8	-	
OTC products	2.1	2.2	(0.1)	(3.8)	4.0	4.0	0.0	0.
Total pharmaceuticals	191.5	198.9	(7.4)	(3.7)	396.0	415.1	(19.1)	(4.

*3: Tanabe Seiyaku Hanbai Products are composed of generic drugs and the long-listed drugs which were transferred from MTPC.

*4:Active pharmaceutical ingredients and others ordered by other companies.

(4) Forecasts of Investment for Property, Plant and Equipment and Information Systems

	•	- .. ,				,		[Billion yen]
	1st Half of FY2015 Forecasts	1st Half of FY2014 Actual	Increase (decrease)	Change %	FY2015 Forecasts	FY2014 Actual	Increase (decrease)	Change %
Investment in property, plant and equipment/occuring basis	4.1	6.2	(2.1)	(33.7)	10.3	15.7	(5.4)	(34.6)
Investment for information systems/occuring basis	1.2	0.8	0.4	56.3	2.1	1.6	0.5	32.4

[Billion yen]

Major investment in property, plant an in FY2015	d equipment	Major investment for information systems in FY2015		
Production facilities	6.1	R&D related systems	0.6	
Facilities & equipment for R&D	3.2	Production related system	0.2	
Others	1.0	Others	1.3	

(5) Forecasts for Depreciation Costs

	1st Half of FY2015 Forecasts	1st Half of FY2014 Actual	Increase (decrease)	Change %	FY2015 Forecasts	FY2014 Actual	Increase (decrease)	Change %
Property, plant and equipment	3.5	3.7	(0.2)	(5.0)	7.2	7.5	(0.3)	(3.7)
Intangible fixed assets	0.8	0.8	0.0	2.4	1.6	1.6	0.0	3.2

[Billion yen]

Five-Year Financial Data 4

(Amounts less than ¥100 million are rounded.)

(1) Profit and Loss						[Billion yen]
	FY2010	FY2011	FY2012	FY2013	FY2014	Forecast for FY2015
Net sales	409.5	407.2	419.2	412.7	415.1	396.0
Cost of sales	154.6	152.3	166.4	169.4	169.6	147.0
Gross operation profit	255.0	254.9	252.8	243.3	245.5	249.0
SG&A expenses	178.4	185.8	183.8	184.2	178.4	181.5
R&D expenses	65.8	70.2	66.5	70.4	69.6	74.0
Operating income	76.6	69.0	69.0	59.1	67.1	67.5
Ordinary income	76.7	68.8	69.4	61.9	67.7	67.0
Extraordinaly income	0.6	1.2	4.2	15.3	13.7	(7.5)
Extraordinaly loss	13.2	6.1	5.9	4.8	18.6	(7.3)
Net income	37.7	39.0	41.9	45.4	39.5	40.5

(2) Balance Sheet

(2) Balance Sheet					[Billion yen]
	End of FY2010	End of FY2011	End of FY2012	End of FY2013	FY2014
Total assets	818.7	819.9	866.8	886.5	929.3
Current assets	391.6	419.7	476.7	540.5	603.6
Fixed assets	427.1	400.3	390.1	346.0	325.7
Total liabilities	122.7	98.4	113.9	108.6	128.9
Current liabilities	87.7	69.6	86.1	81.8	105.4
Fixed liabilities	35.0	28.9	27.7	26.8	23.5
Net assets	696.0	721.5	752.9	777.8	800.4

(3) Other Financial Data

(-)						. , .
	End of FY2010	End of FY2011	FY2012	FY2013	FY2014	Forecast for FY2015
Cash flows from operating activities	59.1	37.2	60.6	69.9	68.2	-
Cash flows from investing activities	(7.7)	(63.2)	(35.0)	(24.3)	(59.8)	-
Cash flows from financing activities	(15.4)	(17.2)	(23.7)	(21.1)	(21.9)	-
Investments in property, plant and equipment	10.2	7.1	9.2	12.6	15.7	10.3
Investments for development of information systems	0.8	1.2	2.2	2.1	1.6	2.1
Depreciation costs	12.4	12.5	8.4	9.2	9.0	8.8
Equity ratio (%)	84.3	87.3	86.3	86.4	84.9	-
ROE (%)	5.5	5.5	5.7	6.0	5.1	-
Net income per share (¥)	67.27	69.54	74.67	80.92	70.41	72.19
Net assets per share (¥)	1,230.16	1,275.85	1,333.22	1,365.52	1,406.41	-

(4) Number of Employees

	End of FY2010	End of FY2011	End of FY2012	End of FY2013	End of FY2014	Forecast for End of FY2015
Consolidated	9,198	9,180	8,835	9,065	8,457	8,359
Non-consolidated	4,957	4,826	4,850	4,867	4,844	4,838

[Billion yen]

5 Quarterly Trend

(Amounts less than ¥100 million are rounded.)

[Billion yen]

(1) Profit and Los	is									[=	sillion yenj
		FY2013				FY2015					
	Q1 Apr. to Jun.	Q2 Jul. to Sep.	Q3 Oct. to Dec.	Q4 Jan. to Mar.	Full year Actual	Q1 Apr. to Jun.	Q2 Jul. to Sep.	Q3 Oct. to Dec.	Q4 Jan. to Mar.	Full year Actual	Forecasts for full-year
Net sales	103.9	98.9	120.4	89.5	412.7	94.6	104.3	120.9	95.3	415.1	396.0
	25.2%	24.0%	29.2%	21.7%	100.0%	22.8%	25.1%	29.1%	23.0%	100.0%	
Domestic	91.4	85.3	103.3	73.4	353.3	79.8	85.3	98.6	73.6	337.2	301.5
Domestic	25.9%	24.1%	29.2%	20.8%	100.0%	23.7%	25.3%	29.2%	21.8%	100.0%	
Overees	12.5	13.7	17.1	16.1	59.4	14.8	19.1	22.3	21.7	77.9	94.5
Overseas	21.0%	23.0%	28.8%	27.1%	100.0%	19.0%	24.5%	28.7%	27.9%	100.0%	
Cost of sales	43.5	38.9	50.6	36.4	169.4	39.2	39.0	49.8	41.6	169.6	147.0
Sales Cost Ratio	41.9%	39.3%	42.0%	40.7%	41.0%	41.4%	37.4%	41.2%	43.6%	40.9%	37.1%
Gross operating	60.4	60.0	69.8	53.1	243.3	55.4	65.3	71.0	53.8	245.5	249.0
profit	24.8%	24.7%	28.7%	21.8%	100.0%	22.6%	26.6%	28.9%	21.9%	100.0%	
	44.2	45.8	44.8	49.5	184.2	41.7	44.1	42.4	50.2	178.4	181.5
SG&A expenses	24.0%	24.9%	24.3%	26.9%	100.0%	23.4%	24.7%	23.8%	28.2%	100.0%	
R&D expenses	17.6	16.7	17.1	19.0	70.4	16.3	15.6	16.2	21.5	69.6	74.0
	24.9%	23.8%	24.3%	27.0%	100.0%	23.5%	22.4%	23.2%	30.9%	100.0%	
	26.6	29.1	27.7	30.5	113.8	25.3	28.5	26.3	28.7	108.8	107.5
Non-R&D expenses	23.4%	25.6%	24.3%	26.8%	100.0%	23.3%	26.2%	24.1%	26.4%	100.0%	
	11.9	12.0	12.4	12.1	48.4	11.1	11.9	11.8	12.0	46.8	46.0
Labor costs	24.5%	24.8%	25.6%	25.1%	100.0%	23.8%	25.5%	25.2%	25.6%	100.0%	
Amortization of	2.6	2.7	2.6	2.8	10.6	2.7	2.7	2.7	2.8	10.9	10.5
goodwill	24.5%	25.0%	24.5%	26.0%	100.0%	24.7%	24.7%	24.7%	25.8%	100.0%	
	12.1	14.4	12.7	15.6	54.8	11.5	13.9	11.8	13.9	51.1	51.0
Others	22.2%	26.3%	23.1%	28.4%	100.0%	22.6%	27.1%	23.1%	27.2%	100.0%	
<u> </u>	16.2	14.2	25.1	3.6	59.1	13.7	21.3	28.6	3.5	67.1	67.5
Operating income	27.5%	24.1%	42.4%	6.1%	100.0%	20.4%	31.7%	42.7%	5.3%	100.0%	
Ordinansia	17.1	15.1	25.6	4.1	61.9	14.6	20.9	28.6	3.6	67.7	67.0
Ordinary income	27.6%	24.5%	41.3%	6.6%	100.0%	21.5%	30.9%	42.3%	5.3%	100.0%	
Natinggma	10.4	18.1	15.3	1.5	45.4	9.6	22.9	15.8	(8.8)	39.5	40.5
Net income	22.9%	39.9%	33.7%	3.4%	100.0%	24.3%	58.1%	39.9%	(22.3%)	100.0%	

The each figure (excluding Cost of sales) in the lower displays the progress rate.

(2) Sales of Main Products

[Billion yen]

) Sales UI											[D	-
				FY2013					FY2014			FY201
		Q1	Q2	Q3	Q4	Full Year	Q1	Q2	Q3	Q4	Full Year	Full Yea
		Apr. to Jun.	Jul. to Sep.	Oct. to Dec.		Actual	Apr. to Jun.	Jul. to Sep.	Oct. to Dec.	Jan. to Mar.	Actual	Forecas
thical drugs		102.3 25.1%	97.3 23.9%	119.1 29.3%	88.4 21.7%	407.2 100.0%	93.2 22.7%	103.3 25.1%	119.8 29.2%	94.4 23.0%	410.7 100.0%	39 [.]
		88.5	82.4	100.5	70.2	341.7	77.5	23.1%	29.2 <i>%</i> 96.8	71.9	323.9	294
Ethical drugs do	omestic sales	25.9%	24.1%	29.4%	20.5%	100.0%	23.9%	24.0%	29.9%	22.2%	100.0%	
Remica	do	19.2	19.9	21.9	15.4	76.3	17.3	17.9	19.3	16.1	70.6	7
Remica	ue	25.1%	26.0%	28.7%	20.2%	100.0%	24.6%	25.3%	27.3%	22.8%	100.0%	
Talion		2.7	2.4	4.4	4.2	13.7	3.3	2.8	4.3	5.7	16.0	1
		19.9%	17.5%	31.9%	30.7%	100.0%	20.3%	17.7%	26.6%	35.4%	100.0%	
Ceredist	t	4.7	4.4	5.0 28.4%	3.6	17.8	4.1	3.8 24.0%	4.4	3.4	15.7	1
— — —		26.5% 4.0	24.9% 3.8	28.4%	20.2% 3.3	100.0% 15.5	26.4% 3.8	24.0%	27.9% 3.8	21.7% 3.2	100.0% 14.1	1
Maintate	e	25.7%	24.3%	28.6%	21.4%	100.0%	26.7%	23.9%	26.8%	22.6%	100.0%	
		2.9	2.7	3.4	2.1	11.1	2.8	2.8	3.5	2.5	11.6	
Venoglo	bulin IH	26.2%	24.6%	30.3%	18.9%	100.0%	24.4%	24.3%	30.1%	21.2%	100.0%	
Simponi		2.1	2.4	2.8	2.1	9.4	2.5	2.6	2.9	2.5	10.5	1
Simpon		22.1%	25.2%	29.8%	22.9%	100.0%	23.5%	24.6%	27.9%	23.9%	100.0%	
Kremez	zin	3.2	3.2	3.5	2.6	12.6	2.8	2.6	2.8	2.3	10.5	
		25.8%	25.2%	28.1%	20.9%	100.0%	26.6%	24.9%	26.8%	21.8%	100.0%	
Urso		3.3	3.1	3.5	2.5	12.4	2.7	2.4	2.7	2.2	10.0	
		26.5% 3.1	25.0% 2.8	28.2% 3.2	20.3%	100.0%	26.7% 2.4	24.2% 2.0	27.0%	22.1% 1.7	100.0% 8.3	
Anplag		27.7%	2.8	28.3%	2.1 19.1%	100.0%	2.4 28.7%	2.0	2.2	20.5%	0.3 100.0%	
		21.170	2.4	20.0%	2.1	9.8	20.7 %	20.0%	27.0%	1.8	8.1	
Depas		26.1%	24.7%	27.4%	21.8%	100.0%	25.9%	24.9%	27.1%	22.2%	100.0%	
Lavanta		1.0	1.4	2.3	1.7	6.5	1.7	1.7	2.7	1.8	8.0	
Lexapro	1	15.9%	21.8%	35.4%	26.9%	100.0%	21.1%	21.7%	34.3%	22.9%	100.0%	
Radicut		3.0	2.7	3.2	2.1	10.9	2.0	1.8	2.1	1.5	7.4	
		27.1%	25.1%	28.9%	18.9%	100.0%	26.7%	24.7%	28.3%	20.3%	100.0%	
Tenelia		0.0	0.0	0.5	0.3	0.8	1.1	1.6	1.9	1.7	6.2	
		0.1% 1.9	3.8%	64.0% 1.9	32.1% 1.4	100.0% 6.9	17.9% 1.5	24.9% 1.3	30.6% 1.5	26.5% 1.2	100.0% 5.5	
Herbess	ser	26.9%	24.7%	28.2%	20.2%	100.0%	27.1%	24.3%	27.3%	21.4%	100.0%	
		1.7	1.5	1.8	1.2	6.2	1.3	1.1	1.2	1.0	4.6	
Tanatril		27.4%	24.6%	28.4%	19.6%	100.0%	27.9%	24.3%	27.0%	20.8%	100.0%	
BIKEN p	products	8.8	5.2	9.6	4.9	28.4	4.9	6.2	14.8	4.4	30.3	2
[vaccines	s]	30.9%	18.3%	33.7%	17.1%	100.0%	16.2%	20.4%	48.8%	14.5%	100.0%	
	Tetrabik	2.9	0.5	1.2	2.2	6.7	1.8	1.8	2.1	1.8	7.5	
		43.1%	7.5%	17.3%	32.2%	100.0%	23.6%	23.7%	28.2%	24.4%	100.0%	
	Influenza	(0.1)	1.2	6.5	(0.4)	7.2	(0.1)	0.9	7.3	(0.8)	7.4	
	Varicella	(0.7%) 1.0	16.2% 0.9	90.6% 0.7	(6.1%) 1.0	100.0% 3.6	(0.7%) 0.6	12.7% 1.3	99.3% 3.5	(11.3%) 1.7	100.0% 7.2	
	vaccine	28.9%	25.0%	19.3%	26.8%	100.0%	8.7%	18.2%	48.8%	24.3%	100.0%	
Tanabe S	eiyaku Hanbai	3.5	3.2	4.1	3.2	14.1	3.4	3.1	3.9	3.3	13.6	1
products *	1	25.0%	22.5%	29.4%	23.1%	100.0%	24.8%	22.5%	28.6%	24.0%	100.0%	
Ethical drugs ov	verseas sales	5.1	5.3	5.9	5.7	22.0	5.3	6.0	5.5	6.2	23.0	2
		23.3%	24.1%	26.9%	25.8%	100.0%	23.0%	26.3%	23.7%	27.1%	100.0%	
Herbe	sser	1.5	1.3	1.5	1.6	5.8	1.6	1.9	1.4	1.6	6.5	
Argatro		25.3% 0.7	22.6%	25.1%	26.9% 0.6	100.0%	24.4% 0.7	29.1%	22.3% 0.5	24.2% 0.4	100.0%	
(Novas		0.7 24.8%	0.7 26.3%	0.7 25.3%	23.7%	2.7 100.0%	32.5%	0.6 26.5%	22.9%	0.4 18.1%	2.1 100.0%	
		0.5	0.4	0.5	0.5	1.8	0.5	0.6	0.4	0.4	1.8	
Tanatr	il	25.8%	20.8%	27.2%	26.3%	100.0%	24.7%	31.4%	21.8%	22.1%	100.0%	
Contracted mar	nufacturing	1.5	1.4	1.4	1.5	5.8	0.9	1.2	0.5	0.8	3.4	
products *2		25.9%	24.5%	24.0%	25.6%	100.0%	27.0%	34.6%	15.1%	23.2%	100.0%	
Royalty incor	me, etc	7.1	8.2	11.3	11.0	37.6	9.5	18.4	17.0	15.5	60.4	7
	, 510.	18.9%	21.8%	30.0%	29.3%	100.0%	15.7%	30.4%	28.2%	25.7%	100.0%	
Royalty f	rom Gilenya	6.5	7.6	9.5	8.6	32.2	7.7	11.1	13.7	11.4	43.9	Undiscl
		20.1%	23.6%	29.6%	26.6%	100.0%	17.6%	25.3%	31.1%	26.0%	100.0%	11
Royalty f		Undisclosed	Undisclosed	Undisclosed	Undisclosed	Undisclosed	Undisclosed	Undisclosed	Undisclosed	3.6	9.8	Undiscl
	AN	- 1.1	- 1.3	- 1.1	- 0.9	- 4.5	- 1.2	- 1.0	- 1.0	37.3% 0.8	100.0% 4.0	
TC products		1.1 25.5%	29.0%	24.4%	0.9 21.1%	4.5 100.0%	1.2 30.4%	1.0 24.2%	24.9%	0.8 20.4%	4.0 100.0%	
		103.9	98.9	120.4	89.5	412.7	94.6	104.3	120.9	95.3	415.1	39
tal sales											÷.,	

The each figure in the lower displays the progress rate.

*1: Tanabe Seiyaku Hanbai Products are composed of generic drugs and the long-listed drugs which were transferred from MTPC.

*2 Active pharmaceutical ingredients and others ordered by other companies.

6 State of New Product Development (As of May 8, 2015)

i. New Drugs

Development code (Generic name)	Category (Indications)	Region	Stage	Origin	
TA-650 (Infliximab [recombinant])	Anti-human TNFα monoclonal antibody (Crohn's disease, ulcerative colitis, pediatric Crohn's disease, pediatric ulcerative colitis)	Taiwan	Filed (Sep., 2013)	US:Janssen Biotech	
TA-7284 (Canagliflozin)	SGLT2 inhibitor (Type 2 diabetes mellitus)	Taiwan	Filed (Mar., 2015)	In-house	
MP-513	DPP-4 inhibitor	Indonesia	Filed (Apr., 2015)		
(Teneligliptin)	(Type 2 diabetes mellitus)	Europe US	Phase 2 Phase 1	In-house	
MT-4666	α7nACh receptor agonist (Dementia of Alzheimer's type)	Global clinical trial*	Phase 3	US: FORUM Pharmaceutica	
MT-2412 (Teneligliptin, Canagliflozin)	Fixed-dose combination of DPP-4 inhibitor and SGLT2 inhibitor (Type 2 diabetes mellitus)	Japan	Phase 3	In-house	
MP-214 (Cariprazine)	Dopamine D3/D2 receptor partial agonist (Schizophrenia)	Japan,Asia	Phase 2b/3	Hungary: Gedeo Richter	
		Europe	Phase 2		
MT-3995	Selective mineralocorticoid receptor antagonist (Diabetic nephropathy)	Japan	Phase 2	In-house	
	(US	Phase 1		
	S1P receptor functional antagonist (Multiple sclerosis)	Europe	Phase 2		
MT-1303	(Psoriasis)	Europe	Europe Phase 2		
	(Inflammatory diseases, autoimmune diseases)	Japan,Europe, US	Phase1	-	
MT-2301	Haemophilus influenza type b (Hib) vaccine (Prophylaxis of pediatric Hib infection)	Japan	Phase 2	US: Nuron Biotech	
Influenza vaccine	Plant-based VLP vaccine (Prophylaxis of H5N1 influenza)	Canada	Phase 2	In-house	
Influenza vaccine	Plant-based VLP vaccine (Prophylaxis of seasonal influenza)	US, Canada	Phase 2	In-house	
Influenza vaccine	Plant-based VLP vaccine (Prophylaxis of H7N9 influenza)	Canada	Phase 1	In-house	
GB-1057 (Recombinant human serum albumin)	Recombinant human serum albumin (Stabilizing agent)	US	Phase 1	In-house	
MP-124	PARP inhibitor (Acute ischemic stroke)	US	Phase 1	In-house	
MP-157	Angiotensin type 2 receptor agonist (Hypertension)	Europe	Phase 1	In-house	
MT-0814	CC chemokine receptor 3 antagonist (Age-related macular degeneration)	Japan	Phase 1	In-house	

* Co-developed with FORUM Pharmaceuticals.

ii. Additional Indications

Product name (Generic name)	Category (Indications)	Region	Stage	Origin	Notes
Talion (Bepotastine)	Selective histamine H1 receptor antagonist, anti- allergic agent (Pediatric allergic rhinitis)	Japan	sNDA filed (May, 2014)	Japan: Ube Industries	
, , , , , , , , , , , , , , , , , , ,	(Pediatric atopic dermatitis)		sNDA filed (May, 2014)		
Radicut (Edaravone)	Free radical scavenger (Amyotrophic lateral sclerosis*)	Japan	sNDA filed (Oct., 2014)	In-house	
	Anti-human TNFα monoclonal antibody (Behcet's disease with special lesions*)		sNDA filed (Oct., 2014)		
Remicade	(Refractory Kawasaki disease*)	Japan	Phase 3	US:Janssen Biotech	
(Infliximab [recombinant])	(Pediatric Crohn's disease)		Phase 3		
	(Pediatric ulcerative colitis)		Phase 3		
	(Psoriasis: increased dose)		Phase 3		
Tribik (Adsorbed diphtheria-purified pertussis-tetanus combined vaccine)	Vaccine (Prophylaxis of pertussis, diphtheria, and tetanus; Stage 2 vaccination)	Japan	sNDA filed (Apr., 2015)	Japan:The Research Foundation for Microbial Diseases of Osaka University	Co-developed with The Research Foundation for Microbial Diseases of Osaka University
Telavic (Telaprevir)	NS3-4A protease inhibitor (Chronic hepatitis C, [combination with Feron])	Japan	Phase 3	US:Vertex Pharmaceutic als	
lmusera (Fingolimod)	S1P receptor functional antagonist (Chronic inflammatory demyelinating polyradiculoneuropathy)	Global clinical trial	Phase 3	In-house	Co-developed with Novartis Pharma in Japan, licensed to Novartis overseas
Canaglu (Canagliflozin)	SGLT2 inhibitor (Diabetic nephropathy)	Global clinical trial	Phase 3	In-house	Sponsor: Janssen Research & Development, LLC

* Orphan drug designated

iii. Licensing-out

Development code (Generic name)	Category (Indications)	Region	Stage	Licensee (Notes)
TA-7284	SGLT2 inhibitor (Type2 diabetes mellitus / fixed dose combination with metformin, XR)	US	Phase 3	US: Janssen Pharmaceuticals,
(Canagliflozin)	(Diabetic nephropathy)	Global clinical trial	Phase 3	Inc
FTY720 (Fingolimod)	S1P receptor functional antagonist (Chronic inflammatory demyelinating polyradiculoneuropathy)	Global clinical trial	Phase 3	Switzerland: Novartis (Co-developed with Novartis Pharma in Japan)
Y-39983	ROCK (rho-kinase) inhibitor (Glaucoma)	Japan	Phase 2	Japan: Senju Pharmaceutical
MT-210	5-HT2A/ Sigma 2 receptor antagonist (Schizophrenia)	Europe	Phase 2	US:Minerva Neuroscience
TA-7906	PDE4 inhibitor (Atopic dermatitis)	Japan	Phase 2	Japan: Maruho
MCC-847 (Masilukast)	Leukotriene D4 receptor antagonist (Asthma)	Korea	Phase 2	Korea: SAMA Pharma
TA-8995	CETP inhibitor (Dyslipidemia)	Europe	Phase 2	Netherlands: DEZIMA Pharma
MT-4580	Ca sensing receptor agonist (Secondary hyperparathyroidism in hemodialysis patients)	Japan	Phase 2	Japan: Kyowa Hakko Kirin
sTU-199 (Tenatoprazole)	Proton pump inhibitor (Gastroesophageal reflux disease)	Europe	Phase 1	France: Negma/Sidem
Wf-516	SSRI / 5HT1A receptor antagonists (Depression)	Europe	Phase 1	US:Minerva Neuroscience
Y-803	Bromodomain inhibitor (Hematological cancer)	Europe, Canada	Phase 1	US: Merck [*]
	(Solid cancer)	Europe, Canada	Phase 1	(Development code: OTX015)

* Merck acquired OncoEthix, the licensee, in December 2014.

iv. Changes Since Previous Announcement on February 2, 2015

In-house Development

Development code/product name (Generic name)	Category (Indications)	Region	As of February 2, 2015	As of May 8, 2015
TA-650 (Infliximab [recombinant])	Anti-human TNFα monoclonal antibody (Crohn's disease, Ulcerative colitis, Pediatric Crohn's disease, Pediatric ulcerative colitis)	Taiwan	None	Filed (Sep., 2013)
TA-7284 (Canagliflozin)	SGLT2 inhibitor (Type 2 diabetes mellitus)	Taiwan	None	Filed (Mar., 2015)
MP-513 (Teneligliptin)	DPP-4 inhibitor (Type 2 diabetes mellitus)	Indonesia	None	Filed (Apr., 2015)
Tribik (Adsorbed diphtheria-purified pertussis-tetanus combined vaccine)	Vaccine (Prophylaxis of pertussis, diphtheria, and tetanus; Stage 2 vaccination)	Japan	Phase 3	sNDA filed (Apr., 2015)
MP-424 (Telaprevir)	NS3-4A protease inhibitor (Chronic hepatitis C)	Korea	Phase 1	Discontinued
Telavic (Telaprevir)	NS3-4A protease inhibitor (Chronic hepatitis C, [combination with Pegasys])	Japan	Phase 3	Discontinued
BindRen (Colestilan[INN])	Non-absorbed phosphate binder (Pediatric hyperphosphatemia)	Europe	Phase 3	Discontinued
Cholebine	Bile acid signal regulation (Type 2 diabetes mellitus)	lanan	Phase 2	Discontinue
(Colestimide[JAN])	Non-absorbed phosphate binder (Hyperphosphatemia)	Japan	Phase 1	Discontinued

Licensing-out

Development code (Generic name)	Category (Indications)	Region	As of February 2, 2015	As of May 8, 2015
MP-513 (Teneligliptin)	DPP-4 inhibitor (Type2 diabetes mellitus / fixed dose combination with metformin, XR)	Korea	NDA filed*	Approved (Mar., 2015)
FTY720 (Fingolimod)	S1P receptor functional antagonist (Primary progressive multiple sclerosis)	Global clinical trial	Phase 3	Discontinued

* 20mg/1000mg(teneligliptin/metformin), 10mg/750mg and 10mg/500mg were submitted in Oct., Nov., and Dec. 2014, respectively.

5. Additional Information for State of New Product Development (as of May 8, 2015)

(1) New Drugs

Development code (Generic name)	Information
TA-650 (Infliximab[recombinant])	TA-650 is an anti-human TNFα monoclonal antibody. In Japan, it was launched under the brand name of Remicade® in 2002.
TA-7284 (Canagliflozin)	As a selective SGLT2 inhibitor, TA-7284 decreases blood glucose levels by inhibiting reabsportion of glucose in the kidney. It was launched in Japan for the treatment of type2 diabetes mellitus in September 2014, under the branc name of CANAGLU®.
MP-513 (Teneligliptin)	MP-513 selectively inhibits DPP-4, thus accelerates the insulin secretion after meal intake without effect on the fasting insulin secretion. It was launched in Japan for the treatment of type2 diabetes mellitus in September 2012, under the brand name of TENELIA®.
MT-4666	MT-4666, licensed from FORUM Pharmaceuticals(US), is an α 7nACh receptor agonist, which ameliorates cognitive dysfunction by activation of both the cholinergic system and the glutamatergic system. Clinical stage is Phase 3 for dementia of Alzheimer's type. It is a global clinical trial and co-developed with FORUM Pharmaceuticals.
MT-2412	MT-2412 is a fixed-dose combination of Teneligliptin(DPP-4 inhibitor) and Canagliflozin(SGLT2 inhibitor).
MP-214 (Cariprazine)	MP-214 is a dopamine D3/D2 receptor partial agonist, licensed from Gedeon Richter (Hungary). Efficacy on negative symptoms and cognitive functions in addition to positive symptoms for schizophrenia is expected.
MT-3995	MT-3995 is a selective mineralocorticoid receptor antagonist, which shows renoprotective effect on diabetic nephropathy.
MT-1303	MT-1303 is a sphingosine-1-phosphate receptor functional antagonist, which keeps lymphocytes sequestered in the lymph nodes and prevents them from contributing to autoimmune reactions. It's a successor of Imusera/Gilenya.
MT-2301	MT-2301 is a Haemophilus influenza type b (Hib) vaccine, licensed from Nuron Biotech(US).
Influenza vaccine	Plant-based VLP influenza vaccine for prophylaxis of H5N1 influenza.
Influenza vaccine	Plant-based VLP influenza vaccine for prophylaxis of seasonal influenza.
Influenza vaccine	Plant-based VLP influenza vaccine for prophylaxis of H7N9 influenza.
GB-1057(Recombinant human serum albumin)	GB-1057 is a recombinant human serum albumin.
MP-124	MP-124 is a PARP inhibitor that has neuroprotective effect.
MP-157	MP-157 is an angiotensin type2 receptor agonist.
MT-0814	MT-0814 is a CC chemokine receptor 3 antagonist.

(2) Additional Indications

Product name (Generic name)	Information
Talion (Bepotastine)	Talion is a selective histamine H1 receptor antagonist. It was launched as an anti-allergic agent for adult in 2000.
Radicut (Edaravone)	Radicut is a free radical scavenger. In 2001, it was launched for improvement neurological symptoms at the acute stage of cerebral infarction, interference with activities of daily living and functional disability.
Remicade (Infliximab[recombinant])	Remicade is an anti-human TNFα monoclonal antibody. In Japan, it was launched as a treatment for Crohn's disease in 2002, followed by rheumatoid arthritis, intractable uveoretinitis caused by Behcet's disease, psoriasis, ankylosing spondylitis, and ulcerative colitis.
Tribik (Adsorbed diphtheria-purified pertussis-tetanus combined vaccine)	Tribik is a diphtheria-purified pertussis-tetanus combined vaccine. It has been jointly developed with the Research Foundation for Microbial Diseases of Osaka University.
Telavic (Telaprevir)	Telavic was launched in Japan for the treatment of chronic hepatitis C (genotype1) in 2011, followed by Chronic hepatitis C (genotype2) in September, 2014.
Imusera (Fingolimod)	Imusera is a sphingosine-1-phosphate receptor functional antagonist, which keeps lymphocytes sequestered in the lymph nodes and prevents them from attacking the myelin of the nerve cells in multiple sclerosis. It was launched as a treatment for multiple sclerosis in 2011 in Japan. Imusera had been jointly developed with Novaltis Pharma for the domestic market. Global Phase 3 study for chronic inflammatory demyelinating polyradiculoneuropathy is underway. It has been jointly developed with Novartis Pharma for the domestic market.
CANAGLU (Canagliflozin)	As a selective SGLT2 inhibitor, CANAGLU decreases blood glucose levels by inhibiting reabsportion of glucose in the kidney. It was launched in Japan for the treatment of type2 diabetes mellitus in September, 2014. It was launched for the treatment of type2 diabetes mellitus under the brand name of INVOKANA® by Janssen Pharmaceuticals, Inc. in the US and its affiliate in Europe.

(3) Licensing-out

Development code (Generic name)	Information
TA-7284 (Canagliflozin)	As a selective SGLT2 inhibitor, TA-7284 decreases blood glucose levels by inhibiting reabsorption of glucose in the kidney. It was launched for the treatment of type2 diabetes mellitus under the brand name of INVOKANA® by Janssen Pharmaceuticals, Inc. in the US and its affiliate in Europe. The fixed dose combination with metformin (IR) was approved in Europe (April, 2014) and the US (August, 2014).
FTY720 (Fingolimod)	Sphingosine-1-phosphate receptor functional antagonist. It was launched as a treatment for multiple sclerosis under the brandname of Imusera by Mitsubishi Tanabe Pharma in Japan. It is also marketed under the brand name of Gilenya by Novartis.
Y-39983	Y-39983 is a ROCK (Rho-kinase) inhibitor, which relaxes vascular smooth muscles.
MT-210	MT-210 is a 5-HT2A/ Sigma 2 receptor antagonist.
TA-7906	TA-7906 is a PDE4 inhibitor.
MCC-847 (Masilukast)	MCC-847 is a Leukotriene D4 receptor antagonist.
TA-8995	TA-8995 is a CETP inhibitor, which raises HDL-C levels and lowers LDL-C levels.
MT-4580	MT-4580 is a Ca sensing receptor agonist.
sTU-199 (Tenatoprazole)	sTU-199 is an isomer of TU-199, developed in Japan, and licensed to Negma (France). Pharmacokinetic/pharmacodynamic results from Phase 1 in Europe and the US demonstrated that sTU-199 controlled gastric acid secretion at nighttime in patients receiving this compound once-daily, with the long half-life. It is expected that this compound could reveal rapid improvement for non-erosive reflux disease.
Wf-516	Wf-516 is a SSRI / 5HT1A receptor antagonists.
Y-803	Y-803 is a Bromodomain inhibitor.

Others 7

1 Subsidiaries and Affiliated Companies

(1) Number of Subsidiaries and Affiliated Companies

	End of FY2014	End of FY2013	Increase (Decrease)	Notes
Consolidated subsidiaries	28	31	(3)	Decrease: Mitsubishi Pharma (Guangzhou), Benesis, Mitsubishi Tanabe Pharma America
Non-consolidated subsidiaries	1	2	(1)	Decrease: CMIC CMO Ashikaga
Affiliated companies	1	5		Decrease: API Corporation, Arkema Yoshitomi, Mapic Europe, Mapic India
Total	30	38	(8)	

(2) Consolidated Subsidiaries

[As of March 31, 2015]

(2)	2) Consolidated Subsidiaries [As of March 31, 2015]							
	Company Name	Paid-in Capital	% Voting [% Ind Owners	irect	Settling Day	Description of Business		
1	Mitsubishi Tanabe Pharma Factory Ltd.	JPY 1,130 million	100.0	[-]	End of Mar.	Manufacture and sale of pharmaceuticals		
2	Mitsubishi Tanabe Pharma Korea Co., Ltd.	KRW 2,100,000,000	100.0	[-]	End of Mar.	Manufacture and sale of pharmaceuticals		
3	Tianjin Tanabe Seiyaku Co., Ltd.	USD 16,230,000	75.4	[-]	End of Dec.	Manufacture and sale of pharmaceuticals		
4	Yoshitomiyakuhin Corporation	JPY 385 million	100.0	[-]	End of Mar.	Provision of information about pharmaceuticals		
5	Bipha Corporation	JPY 100 million	100.0	[-]	End of Mar.	Manufacture and sale of pharmaceuticals		
6	Tanabe Seiyaku Yoshiki Factory Co., Ltd.	JPY 400 million	100.0	[-]	End of Mar.	Manufacture and sale of pharmaceuticals		
7	Tanabe Seiyaku Hanbai., Ltd.	JPY 169 million	100.0	[-]	End of Mar.	Sale of generic pharmaceuticals, etc.		
8	Tanabe R&D Service Co., Ltd.	JPY 44 million	100.0	[-]	End of Mar.	Support of R&D regarding pharmaceuticals		
9	Tanabe Total Service Co., Ltd.	JPY 90 million	100.0	[-]	End of Mar.	Real estate management and creation of promotion materials, etc.		
10	MP Healthcare Venture Management, Inc.	USD 100	100.0	[100.0]	End of Mar.	Investments in bio-ventures		
11	Mitsubishi Tanabe Pharma Holdings America, Inc. Mitsubishi Tanabe Pharma Development America,	USD 167	100.0	[-]	End of Mar.	Management of group companies in US		
12	Inc.	USD 100	100.0	[100.0]	End of Mar.	R&D of pharmaceuticals		
13	Tanabe Research Laboratories U.S.A., Inc.	USD 3,000,000	100.0	[100.0]	End of Mar.	R&D of pharmaceuticals		
14	MTPC Holdings Canada Inc.	CAD 242 million	100.0	[-]	End of Mar.	Investments in Medicago Group		
15	Medicago Inc.	CAD 253 million	60.0	[55.9]	End of Mar.	Manufacture and sale of vaccines		
16	Medicago USA Inc.	USD 99	60.0	[60.0]	End of Mar.	Manufacture of vaccines		
17	Medicago R&D Inc.	CAD 500	60.0	[60.0]	End of Mar.	R&D of vaccines		
18	Mitsubishi Tanabe Pharma Development (Beijing) Co., Ltd.	USD 1,000,000	100.0	[-]	End of Dec.	R&D of pharmaceuticals		
19	Guangdong Tanabe Pharmaceutical Co., Ltd.	CNY 7,000,000	100.0	[-]	End of Dec.	Sale of pharmaceuticals		
20	Taiwan Tanabe Seiyaku Co., Ltd.	TWD 90,000,000	65.0	[-]	End of Mar.	Manufacture and sale of pharmaceuticals		
21	Tai Tien Pharmaceuticals Co., Ltd.	TWD 20,000,000	65.0	[-]	End of Mar.	Sale of pharmaceuticals		
22	P.T. Tanabe Indonesia	USD 2,500,000	99.6	[-]	End of Mar.	Manufacture and sale of pharmaceuticals		
23	Mitsubishi Tanabe Pharma Europe Ltd.	GBP 4,632,000	100.0	[-]	End of Mar.	R&D of pharmaceuticals		
24	Mitsubishi Tanabe Pharma GmbH	EUR 25,000	100.0	[100.0]	End of Mar.	Sale of pharmaceuticals		
Not	lote: Aside from the companies mentioned above, there are four consolidated companies under the liquidations.							

(3) Affiliated Companies Accounted for by the Equity Method

(3)	Affiliated Companies Accounted for by the Equity M	/lethod			[As of March 31, 2015]
	Company Name	Paid-in Capital (Million yen)	% Voting Control [% Indirect Ownership]	Settling Day	Description of Business
1	Synthelabo-Tanabe Chimie S.A.	EUR 1,600,000	50.0 [-]	End of Dec.	Manufacture and sale of pharmaceuticals

2 Status of Shareholders

(1) Number of Outstanding Shares

	End of March, 2015	End of March, 2014
Issued	561,417,916	561,417,916
The company's own shares at the end of the period	428,340	426,862
Number of shares outstanding at the end of the period	560,989,576	560,991,054
Average number of the company's own share in the period	427,456	425,775
Average number of shares outstanding in the period	560,990,460	560,992,141

(2) Status of Major Shareholders

		End of Ma	rch, 2015	End of March, 2014			
Rank	Name of Shareholders	Number of Shares (Thousands)	Percentage of Total	Rank	Number of Shares (Thousands)	Percentage of Total	
1	Mitsubishi Chemical Holdings Corporation	316,320	56.34%	1	316,320	56.34%	
2	The Master Trust of Japan, Ltd.	24,137	4.30%	2	22,305	3.97%	
3	Nippon Life Insurance Company	12,065	2.15%	3	13,574	2.42%	
4	Japan Trustee Services Bank, Ltd.	10,669	1.90%	4	9,406	1.68%	
5	The Bank of Tokyo-Mitsubishi UFJ, Ltd.	7,254	1.29%	5	7,254	1.29%	
6	STATE STREET BANK AND TRUST COMPANY 505225	5,488	0.98%	10	4,432	0.79%	
7	Employee Stock Ownship Plan	4,505	0.80%	9	4,779	0.85%	
8	STATE STREET BANK CLIENT OMNIBUS OM04	4,172	0.74%	56	921	0.16%	
9	Nipro Corporation	3,821	0.68%	13	3,821	0.68%	
10	STATE STREET BANK WEST CLIENT-TREATY 505234	3,635	0.65%	17	2,767	0.49%	

(3) Ownership and Distribution of Shares

	End of March, 2015			End of March, 2014		
	Number of Shareholders	Number of Shares (Thousands)	Percentage of Total	Number of Shareholders	Number of Shares (Thousands)	Percentage of Total
Financial institutions	75	98,321	17.52%	77	85,620	15.25%
Foreign corporations and others	465	100,650	17.93%	402	110,839	19.75%
Individuals and others	13,972	24,578	4.38%	16,660	28,217	5.03%
Other corporations	251	334,680	59.62%	270	334,919	59.67%
Securities firms	34	3,083	0.55%	28	1,716	0.31%
Total	14,797	561,315	100.00%	17,437	561,314	100.00%
Less than trading unit	-	102	-	-	103	-

* The trading unit of the Company's stock is 100 shares.

* Individuals and Others include treasury stocks (428 thousands shares at the end of March, 2015 and 426 thousands shares at the end of March, 2014)

(4) Trend of Divinded and Stock Price

	FY2010	FY2011	FY2012	FY2013	FY2014	FY2015 Estimate
Dividends per share (yen)	28	35	40	40	42	44
Dividend payout ratio(%)	41.6	50.3	53.6	49.4	59.6	60.9
(prior to amortization of goodwill)	(32.9)	(40.0)	(43.2)	(40.5)	(47.6)	(48.8)
Stock price at the end of FY	1,350	1,161	1,445	1,443	2,062	-
Market capitalization (billion yen)	7,579	6,518	8,112	8,101	11,576	-

Reference

Major Ethical Drugs

Major Ethical Drugs			
Remicade (Infliximab)	Launch: May 2002	Category	Anti-TNFα monoclonal antibody
with a single administration. It h	as indications for and ulcerative coli	the treatment	ant inflammatory cytokine. It is very fast-acting and its efficacy is sustained for eight weeks of rheumatoid arthritis, Crohn's disease, Behcet's disease with refractory uveoretinitis in July 2009 and August 2011, changes in usage/dosage were approved for rheumatoic
Talion (Bepotastine)	Launch: Oct. 2000	Category	Agent for treatment of allergic disorders
			n demonstrated to be effective for allergic rhinitis, urticaria, and pruritus accompanying ulation, orally disintegrating tablets, was approved in March and launched in July 2007.
Ceredist (Taltirelin)	Launch: Sep. 2000	Category	Agent for treatment of spinocerebellar degeneration
			gainst ataxia caused by spinocerebellar degeneration, but it was previously administered s first oral TRH derivative drug. An additional formulation, orally disintegrating tablets, was
Maintate (Bisoprolol)	Launch: Nov. 1990	Category	Selective $\beta 1$ antagonist (Treatment of hypertension, angina pectoris, and arrhythmias)
phamacokinetics profiles. It has	high efficacy and 11, the indication of	safety, and evi of atrial fibrillat	countries around the world. It exhibits high selectivity for β 1 receptor and excellen idence-based cardioprotective action. In addition to the indication of chronic heart failure ion has been newly approved in June, 2013. Maintate is the only β -blocker with both
Simponi (Golimumab)	Launch: Sep. 2011	Category	Anti-TNFα monoclonal antibody
	eutical. It shows a		ent of rheumatoid arthritis (including prevention of articular structural damage), and co- cacy by subcutaneous injection once every four weeks, and currently is under developmer
Kremezin	Launch: Apr. 2011	Category	Agent for treatment of Chronic renal failure
Keremezin was introduced to the	arcoal consisting c Japanese market 1, the marketing ric	in December 1	ical activated carbon of high purity. It absorbs and excretes uremic toxins out of the body 991 as the first pharmaceuticals drug in the world for proactive treatment of chronic rena ferred from Daiichi Sankyo to MTPC.
Urso (Ursodeoxycholic Acid)	Launch: July 1962	Category	Agent for improving hepatic, biliary and digestive functions
	one of the bile acid	Is existing in th	en extracted from blackbear's gallbladder in the past and has been used in the treatment o ne human body. Urso has effects of hapatic protection and indications of improvement o n of gallstones.
Anplag (Sarpogrelate)	Launch: Oct. 1993	Category	5-HT2 blocker (Anti-platelet agent)
associated with chronic arterial c	ed to patients with occlusion. Anplag	especially imp	s obliterans (ASO) to improve ischemic symptoms like as ulcer, pain and coldness of limbs roves the bloodstream of collateral circulation and inhibits platelet aggregation, vascular stic action to serotonin receptor in platelets and vessels.
Depas (Etizolam)	Launch: Mar. 1984	Category	Antianxiety agent
		-	e to its broad pharmacological properties, Depas shows reasonable effectiveness for iscle-contraction headache, depression and sleep disorder.

Lexapro (Escitalopram)	Launch: Aug. 2011	Category	Selective sertonin reuptake inhibitor (SSRI)
Lexapro, a highly selective serotonin re	intake inhibitor (S	SRI) has be	en globally approved in 98 countries and regions. It shows good efficacy and tolerability

Lexapro, a nighty selective serotonin reuptake inhibitor (SSRI), has been globally approved in 98 countries and regions. It shows good emicacy and tolerability in patients with depressive disorder. Moreover, due to simple dosage and administration, it is expected to improve adherence of the treatment. Origin: H. Lundbeck A/S (Denmark), Manufacturer and distributor: Mochida Pharmaceutical Co., Ltd

Dedicut (Education)	Launch:	Category	Free radical scavenger (Cerebral neuroprotectant)
Radicut (Edaravone)	Jun. 2001		

Radicut is the world's first brain protecting agent (free radical scavemger) shown to improve neurological symptoms, interference with activities of daily living, and disability (at hospital discharge) in patients at acute stage of cerebral infarction. Specific indications include the treatment of various types of infarction (cerebral lacunar, atherothrombotic and cardiogenic infarction) It is initiated administration within 24 hours after onset, and is not administrated for more than 14 days. An additional formulation, Radicut bag for I.V. Infusion, was launched in May 2010.

Tenelia (Teneligliptin)	Launch:	Category	Selective DPP-IV inhibitor	
renena (renengiiptin)	Sep. 2012	Category		

Tenelia, which Mitsubishi Tanabe has created and developed, is the first DPP-4 inhibitor originating in Japan that has ever been launched. It inhibits the function of dipeptidyl peptidase-4 (DPP-4), which selectively breaks down glucagon-like peptide-1(GLP-1), a hormone secreted from the gastrointestinal tract in response to food intake. In this way, Tenelia promotes insulin secretion and suppresses glucagon secretion, thereby demonstrating blood glucose lowering action.

Herbesser (Diltiazem)	Launch: Feb. 1974	Category	Calcium antagonist (Treatment of angina pectoris and hypertension)
-----------------------	----------------------	----------	--

Herbesser is a representative calcium antagonist that is used in more than 110 countries around the world. In addition to a blood pressure lowering effect, it has a cardioprotective action in patients with hypertension or angina pectoris by reducing the cardiac load through a heart rate lowering effect and by increasing the oxygen supply through a coronary vasodilating effect.

Tanatril (Imidapril) Launch: Dec. 1993	Category ACE inhibitor (Treatment of hypertension)	
---	--	--

Tanatril shows excellent blood pressure control with effective organ protection as well as minimal incidence of dry cough, a common side effect of ACE inhibitors. With the approval of an additional indication in January 2002, it became the first drug in Japan approved for diabetic nephropathy with type I diabetes mellitus.

TETRABIK	Launch:		
(Absorbed Diphtheria-purified	Oct. 31. 2012	Category	Prevention of Diphtheria, Pertussis, Tetanus and polio
Pertussis-tetanus inactivated polio	000.01.2012		

TETRABIK is a combined vaccine that prevents acute poliomyelitis (polio), pertussis, diphtheria and tetanus. It is used at 1st term (initial 3 times) and 1st term (additional 1 time), in total 4 times, of the regular vaccination. By using TETRABIK, It is expected to avoid the very rare occurrence of paralytic symptoms simila to those in natural polio due to live-attenuated oral polio vaccine.

Origin, Manufacturer and distributor: BIKEN (The Research Foundation for Microbial Diseases of Osaka University)

News Releases

The major news releases after October, 2014 are as follows. Please refer to the Company's website for the details. (http://www.mt-pharma.co.jp/e/release/index.php)

Date	Contents
October 30, 2014	Applications filed for REMICADE for I.V. Infusion 100, an anti-human TNFα monoclonal antibody, for additional indications for entero-Behcet's disease, neuro-Behcet's disease, and vasculo-Behcet's disease
November 6, 2014	Supporting "good blood glucose control" in Japanese patients with diabetes joint promotion with Johnson & Johnson K.K. Medical Company
November 13, 2014	Application filed for additional indication for ALS for RADICUT inj. 30mg and RADICUT BAG for I.V. Infusion 30mg
November 28, 2014	Notice regarding conclusion of the final agreement for the transfer of the Kashima Plant of Mitsubishi Tanabe Pharma Factory Ltd.
December 18, 2014	Reorganization of the U.S. Affiliates to Accelerate the U.S. Operations Expansion
January 29, 2015	-Increasing Production Capacity to Meet Growing Asian Pharmaceutical Markets -Completion of New Manufacturing FacilityTianjin Tanabe Seiyaku Co., Ltd. and P.T. Tanabe Indonesia
February 6, 2015	Completion of Construction of the New Head Office Building
February 25, 2015	Reorganization of research bases in Japan
February 25, 2015	Medicago Develops Alternative Production Process for Ebola Antibodies Contract concluded with U.S. government
April 1, 2015	Execution of License Agreement on VMAT2 Inhibitor NBI-98854

