Financial Results for the 2nd Quarter of Fiscal Year Ending March 31, 2020 < Supplement >

As of October 30, 2019 Mitsubishi Tanabe Pharma Corporation



(Note about forward-looking information)

The statements contained in this presentation is based on a number of assumptions and belief in light of the information currently available to management of the company and is subject to significant risks and uncertainties. It contains information about pharmaceuticals (Include products under development), but is not intended for advertising or medical advice.

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Summary of Financial Results for the 2nd Quarter of FY2019 Ended March 31, 2020 and Forecasts for FY2019

(Amounts less than ¥100 million are rounded off)

1. Summary of Financial Results for the 2nd Quarter of FY2019

[Billion yen]

Revenue	188.1	Y-on-Y	(21.6)	(10.3 %)
Domestic	154.6	Y-on-Y	8.1	5.6 %
Overseas	33.4	Y-on-Y	(29.7)	(47.0 %)

Revenue of domestic ethical drugs increased by 5.4%, year-on-year, to ¥149.1 billion due to the sales growth of priority products contributed by SIMPONI, the treatment agent of Rheumatoid arthritis (RA) and three type 2 diabetes mellitus treatment agents of TENELIA, CANAGLU, and CANALIA, as well as RUPAFIN, an allergy treatment agent with the dismantling of prescription limitation in December 2018 and STELARA, a treatment for Crohn's disease jointly promoted with Janssen Pharmaceutical K.K., updated the co-promotion framework in July 2018.

Royalty revenue, etc. decreased by 74.6%, year-on-year, to ¥9.2 billion due to the decline in royalty revenue from GILENYA, the treatment for multiple sclerosis licensed to Novartis Pharma AG (hereinafter referred to as "Novartis") and INVOKANA and its fixed dose combination with metformin, the treatment for type 2 diabetes mellitus licensed to Janssen Pharmaceuticals, Inc.

With regard to "GILENYA Royalty" amounts, given the arbitration proceedings initiated in February 2019, a part of the royalty income of "GILENYA Royalty" has not been recognized as sales revenue in accordance with IFRS 15. Since the arbitration proceedings were ongoing during the second quarter of the fiscal year ending March 31, 2020, there was a decrease in revenue because of not recognizing a part of the royalty income. The Company maintains it is entitled to receive the full royalty amounts due according to the license agreement with Novartis, and the Company will rigorously pursue its rights in the arbitration. As for the "GILENYA Royalty" amounts which will not be recognized as sales revenue, those will be recognized as revenue at the end of the arbitration, depending on the outcome of the arbitration.

[Billion yen]

Core Operating Profit*	11.6	Y-on-Y	(22.8)	(66.1 %)
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Core operating profit decreased by 66.1%, or ¥22.8 billion, year-on-year, to ¥11.6 billion due to the following results:

- Sales growth of domestic priority products
- The decline of royalty revenue and long listed drugs sales
- Increase in R&D expenses arising from the high level of investments in R&D continuing from the prior year
- *: With adoption of IFRS, the Company, its subsidiaries and its affiliates (collectively, "the Group") has introduced "core operating profit" as a major profit index to demonstrate its recurring profitability and positioned as an important indicator of business management, etc. "Core operating profit" is a profit excluding the income and loss recorded by non-recurring items specified by the Group (hereinafter "non-recurring items") from operating profit. Non-recurring items include gain or loss associated with a business transfer, restructuring loss, impairment losses on intangible assets associated with products, losses on disaster and others.

Operating Profit	12.5	Y-on-Y	(21.9)	(63.6 %)
Profit before Tax for the period	12.1	Y-on-Y	(22.6)	(65.1 %)
Net Income Attributable to owners of the Company	8.3	Y-on-Y	(16.6)	(66.7 %)

2. Summary of Forecasts for FY2019

[Billion yen]

Revenues	376.0	Y-on-Y	(48.7)	(11.5 %)
Core Operating Profit	10.0	Y-on-Y	(45.8)	(82.1 %)
Operating Profit	11.5	Y-on-Y	(38.8)	(77.1 %)
Profit before Tax	12.0	Y-on-Y	(38.4)	(76.2 %)
Net Income Attributable to owners of the Company	5.0	Y-on-Y	(32.3)	(86.6 %)

(Note) Revisions to recently announced consolidated earnings forecasts on May 10, 2019: No

3. Dividends

	FY2019			FY2018			
	End of 1st Half	End of FY2019 (Estimate)	For the Year (Estimate)	End of 1st Half	End of FY2018	For the Year	
Dividends per Share [¥]	28	28	56	28	28	56	
Dividends Payout Ratio	-	-	628.1%	-	-	84.0%	

2 Consolidated Financial Indicators for the 2nd Quarter of FY2019

1. Profit and Loss

(Amounts less than ¥100 million are rounded off)

(1) Profit and Loss

[Billion yen]

		Y-on-Y		Comparison to forecasts			Notes	
1H FY2019		1H FY2018	Increase (decrease)	Change %	Forecasts*1	Increase (decrease)	Change %	Notes [Y-on-Y comparison]
Revenue	188.1	209.7	(21.6)	(10.3)	187.0	1.1	0.6	See "(2) Sales Revenue of Main Products" on page 4
Domestic	154.6	146.4	8.1	5.6	153.6	0.9	0.6	
Overseas Overseas sales ratio	33.4 17.8%	63.2 30.1%	(29.7)	(47.0)	33.3 17.8%	0.1	0.5	
Cost of sales Sales cost ratio	88.5 47.1%	86.1 41.1%	2.3	2.8	87.5 46.8%	1.0	1.2	Increase in the sales cost ratio due to decrease of royalty revenue, etc.
Gross profit	99.6	123.5	(23.9)	(19.4)	99.5	0.1	0.1	
SG&A expenses % of revenue	46.8 24.9%	47.7 22.8%	(0.9)	(1.9)	49.0 26.2%	(2.1)	(4.4)	
R&D expenses % of revenue	39.7 21.2%	39.5 18.9%	0.2	0.6	44.5	(4.7)	(10.6)	
Amortization of intangible assets associated with products	1.2	1.4	(0.2)	(14.5)	1.3	(0.0)	(3.5)	
Other income (expense)*2	(0.0)	(0.3)	0.2	-	(0.2)	0.1	-	
Core operating profit	11.6	34.5	(22.8)	(66.1)	4.5	7.1	159.9	
Non-recurring items*2	0.8	-	0.8	-	0.5	0.3	73.0	
Operating profit	12.5	34.5	(21.9)	(63.6)	5.0	7.5	151.2	
Financial income	0.5	0.5	(0.0)	(4.6)				
Interest income and dividends income	0.5	0.5	(0.0)	(4.3)				
Financial expense	0.9	0.2	0.7	262.3				
Foreign exchange loss	0.7	0.1	0.5	338.9				
Others	0.1	0.0	0.1	517.4				
Profit before tax for the period	12.1	34.8	(22.6)	(65.1)	5.5	6.6	120.9	
Income taxes	5.9	11.6	(5.6)	(48.8)				
Net profit for the period	6.1	23.1	(16.9)	(73.3)	1.0	5.1	518.8	
Net profit attributable to owners of the Company	8.3	24.9	(16.6)	(66.7)	4.0	4.3	107.9	
Total labor cost	38.1	35.8	2.3	6.6	38.4	(0.2)	(0.6)	

^{*1:} The Company announced full year forecasts on May 10, 2019.

^{*2:} Brackets indicate expense and loss

			[Yen]
Exchange rate	1H FY2019 average	1H FY2018 average	FY2019 planed
USD	108.67	110.71	110.00
CAD	81.73	85.08	85.00
EUR	120.91	129.78	125.00

For the 2nd quarter of FY2019, the impact of fluctuations in the foreign exchange rate was as follows;

3

Revenue: decrease by \$1.1 billion

Core operating profit: Increase by ± 0.7 billion

			Y-on-Y		Comparison to forecasts		
	1H FY2019	1H FY2018	Increase (decrease)	Change %	Forecasts*1	Increase (decrease)	Change %
Domestic ethical drugs	149.1	141.5	7.5	5.4	147.5	1.5	1.1
Remicade	27.6	29.9	(2.3)	(7.8)	26.9	0.6	2.4
Simponi	20.4	18.5	1.9	10.4	21.2	(0.7)	(3.7)
Stelara	12.5	4.7	7.8	164.1	11.0	1.4	13.5
Tenelia	8.0	7.2	0.8	12.0	8.0	0.0	0.0
Canaglu	4.1	3.0	1.0	34.9	4.6	(0.5)	(12.2)
Canalia	3.7	3.0	0.6	22.2	4.1	(0.3)	(9.3)
Kremezin	3.3	3.3	(0.0)	(0.4)	4.3	(0.9)	(22.3)
Lexapro	7.4	6.8	0.6	9.0	7.4	0.0	0.6
Ceredist	3.8	4.6	(0.8)	(17.4)	4.5	(0.7)	(15.5)
Rupafin	2.4	0.3	2.1	551.4	2.3	0.1	5.0
Talion	2.1	2.5	(0.3)	(15.5)	2.7	(0.5)	(20.8)
Vaccine [BIKEN products]	15.7	15.5	0.1	1.0	14.4	1.3	9.2
Influenza	1.7	0.9	0.8	82.1	1.0	0.7	71.0
Tetrabik	4.5	4.1	0.4	9.7	4.9	(0.4)	(8.2)
Varicella vaccine	2.5	2.6	(0.1)	(4.6)	2.6	(0.1)	(5.8)
Overseas ethical drugs	24.8	27.4	(2.5)	(9.4)	24.1	0.7	3.0
Radicava	11.6	13.9	(2.2)	(16.2)	11.0	0.5	5.3
Herbesser	3.4	3.3	0.1	4.7	3.5	(0.0)	(1.7)
Simponi	1.0	0.9	0.0	5.8	1.0	0.0	1.6
Argatroban	0.9	1.0	0.0	(5.8)	0.8	0.0	9.8
Tanatril	0.7	0.8	(0.1)	(14.1)	0.8	(0.1)	(14.6)
Royalty revenue, etc.	9.2	36.3	(27.0)	(74.6)	9.8	(0.5)	(6.1)
Royalty from Gilenya	3.2	29.9	(26.6)	(89.1)	Undisclosed	-	-
Royalty from INVOKANA	4.0	4.9	(0.8)	(17.7)	Undisclosed	-	-
OTC products	2.3	2.2	0.1	7.0	2.5	(0.1)	(6.0)
Others*3	2.4	2.1	0.3	15.3	2.9	(0.4)	(15.3)
Total sales revenue	188.1	209.7	(21.6)	(10.3)	187.0	1.1	0.6

^{*1:} The Company announced its first half year forecasts for FY2019 on May 10, 2019

^{*2:} Mitsubishi Tanabe Pharma (MTPC) is currently in the arbitration proceedings with Novartis, and among the Gilenya Royalty amounts that MTPC is going to receive from Novartis, MTPC has decided not to recognize some of those amounts as our revenue because such payments do not satisfy one of the requirements under IFRS 15. The same accounting treatment will be continued during the period of the arbitration proceedings. Regardless of the disclosed amounts, MTPC maintains it is entitled to receive the full royalty amounts due according to the 1997 License Agreement with Novartis, and MTPC will rigorously pursue its rights in the arbitration.

^{*3:} Contracted manufacturing products of other companies

2. Financial Statement

(1) Balance Sheet

[Billion Yen]

rent assets rty, plant and equipment will gible assets tments accounted for using method financial assets efined benefit assets non-current assets red tax assets assets	1055.8 465.7 80.8 89.9 203.7 16.1 41.6 22.6 0.1	100.0 44.1 7.7 8.5 19.3 1.5 3.9 2.1	1056.2 467.8 73.3 91.6 206.9 16.2 46.2	(1.7) (3.1) (0.1)	Capital investments, 5.7; Depreciation, (3.5) Impact of the adoption of IFRS 16, 9.7; Transfer to assets held for sale, (5.7) Decrease due to the impact of fluctuation in the foreign exchange rate Decrease due to the impact of fluctuation in the foreign exchange rate
rty, plant and equipment will gible assets tments accounted for using method financial assets efined benefit assets non-current assets red tax assets assets	80.8 89.9 203.7 16.1 41.6 22.6 0.1	7.7 8.5 19.3 1.5 3.9 2.1	73.3 91.6 206.9 16.2	7.4 (1.7) (3.1) (0.1)	Impact of the adoption of IFRS 16, 9.7; Transfer to assets held for sale, (5.7) Decrease due to the impact of fluctuation in the foreign exchange rate Decrease due to the impact of fluctuation in the foreign exchange rate
will gible assets tments accounted for using method financial assets efined benefit assets non-current assets red tax assets	89.9 203.7 16.1 41.6 22.6 0.1	8.5 19.3 1.5 3.9 2.1	91.6 206.9 16.2	(1.7) (3.1) (0.1)	Impact of the adoption of IFRS 16, 9.7; Transfer to assets held for sale, (5.7) Decrease due to the impact of fluctuation in the foreign exchange rate Decrease due to the impact of fluctuation in the foreign exchange rate
gible assets tments accounted for using method financial assets efined benefit assets non-current assets red tax assets	203.7 16.1 41.6 22.6 0.1	19.3 1.5 3.9 2.1	206.9	(3.1)	the foreign exchange rate Decrease due to the impact of fluctuation in the foreign exchange rate
rments accounted for using remethod financial assets refined benefit assets red tax assets	16.1 41.6 22.6 0.1	1.5 3.9 2.1	16.2	(0.1)	the foreign exchange rate
r method financial assets efined benefit assets non-current assets red tax assets assets	41.6 22.6 0.1	3.9			
financial assets efined benefit assets non-current assets red tax assets assets	22.6 0.1	2.1	46.2	(4 E)	
non-current assets red tax assets assets	0.1			(4.5)	Decrease due to fair value remeasurement investment in securities
red tax assets assets			21.4	1.1	
assets	10.6	0.0	0.2	(0.0)	
		1.0	11.6	(1.0)	
	590.1	55.9	588.4	1.6	
ntories	78.6	7.4	75.5	3.0	
e and other receivables*1	114.3	10.8	116.9	(2.6)	
ade receivable rotation number]	[3.65]		[3.35]		
r financial assets	287.4	27.2	271.4	15.9	
r current assets	15.4	1.5	11.0	4.4	
and cash equivalents	88.5	8.4	111.8	(23.3)	See "(2) Statements of Cash Flow" on page
s held for sale	5.7	0.5	1.6	4.1	Mainly assets related to sale of Toda Office
	162.2	15.4	145.9	16.3	
ent liabilities	73.4	7.0	54.2	19.1	
owings	0.1	0.0	0.1	(0.0)	
r financial liabilities	8.5	0.8	2.1	6.4	
lefined benefit liabilities	0.5	0.1	0.6	(0.0)	
sion	6.6	0.6	6.9	(0.2)	
r non-current liabilities	20.4	1.9	5.1	15.3	
red tax liabilities	36.9	3.5	39.2	(2.2)	
iabilities	88.8	8.4	91.6	(2.8)	
e and other payables ^{*2}	33.3	3.2	31.4	1.8	
r financial liabilities	28.7	2.7	27.0	1.7	
ne taxes payable	2.9	0.3	9.5	(6.6)	
sions	2.4	0.2	1.6	0.8	
r current liabilities	20.8	2.0	21.6	(0.7)	
	893.5	84.6	910.3	(16.7)	
e capital	50.0	4.7	50.0	_	
al surplus	449.6	42.6	451.2	(1.6)	
sury shares				0.0	
•					Net profit for the period, 8.3; Payment for
<u>-</u>					dividends, (15.7)
	ade receivable rotation number] financial assets current assets and cash equivalents sheld for sale ent liabilities wings financial liabilities lefined benefit liabilities sion conocurrent liabilities red tax liabilities e and other payables*2 financial liabilities ne taxes payable sions current liabilities e capital al surplus	ade receivable rotation number] financial assets financial assets current assets and cash equivalents sheld for sale financial liabilities financial liab	(addereceivable rotation number) (addereceivabl	(a.65) (a.35) (a.35) (a.35) (a.65) (a.35) (a.65) (a.35) (a.65) (Section Sect

^{*1:} Trade and other receivables = bills + accounts receivable + allowance for doubtful accounts

^{*2:} Trade receivable rotation number = bills (except non - operating bills) + accounts payable

(2) Cash Flow Statement

[Billion yen]

(2) Cash How Statement	1H FY2019	1H FY2018	Increase (decrease)
Cash and cash equivalents at beginning of year	111.8	127.0	(15.1)
Cash flows from operating activities	19.4	23.4	(4.0)
Profit before tax	12.1	34.8	(22.6)
Depreciation and amortization	7.5	5.8	1.6
Reversal of impairment losses	(1.7)	-	(1.7)
Interest and dividends income	(0.5)	(0.5)	0.0
Share of loss(profit) of affiliates accounted for using equity method	(0.0)	(0.0)	0.0
Decrease(increase) in trade and other receivables	2.5	(1.2)	3.8
Decrease(increase) in inventories	(3.8)	8.1	(12.0)
Increase(decrease) in trade and other payables	2.5	(2.1)	4.6
Increase(decrease) in provisions	0.5	0.6	(0.0)
Decrease(increase) in net defined benefit asset	0.3	0.0	0.2
Interest and dividends received	0.5	0.6	(0.0)
Interest paid	(0.1)	(0.1)	0.0
Income taxes paid	(10.4)	(19.3)	8.9
Other	9.9	(3.1)	13.1
Cash flows from investing activities	(21.9)	(16.8)	(5.1)
Payments into time deposits	-	(1.1)	1.1
Proceeds from withdrawal of time deposits	0.4	3.7	(3.3)
Purchase of property, plant and equipment	(6.4)	(2.1)	(4.2)
Proceeds from sales of property, plant and equipment	1.5	0.0	1.5
Purchase of intangible assets	(2.4)	(0.8)	(1.5)
Purchase of investments	(235.7)	(147.6)	(88.0)
Proceeds from sales and redemption of investments	219.7	131.2	88.4
Proceeds from sales of subsidiaries	1.0	-	1.0
Other	(0.1)	0.0	(0.1)
Cash flows from financing activities	(19.7)	(13.3)	(6.3)
Repayments of lease liabilities	(3.9)	(0.0)	(3.8)
Proceeds from share issuance to non-controlling shareholders	-	2.4	(2.4)
Dividends paid	(15.7)	(15.7)	(0.0)
Other	(0.1)	(0.0)	(0.0)
Effect of exchange rate changes on cash and cash equivalents	(1.1)	1.7	(2.9)
Net increase(decrease) in cash and cash equivalents	(23.3)	(4.9)	(18.4)
Increase(decrease) in cash and cash equivalents due to transfer to assets held for sale	0.0	-	0.0
Cash and cash equivalents at the end of period	88.5	122.0	(33.5)

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

				[Billion yen]
	1H FY2019	1H FY2018	Increase (decrease)	FY2018
Investment in property, plant and equipment / occurring basis	5.7	2.0	3.6	6.8
Investment in information systems / occurring basis	0.5	0.9	(0.4)	1.7

[Billion yen]

Major investment in property, plant and equipment in 1H FY2019		Major investment in development of information systems in 1H FY2019	
Mitsubishi Tanabe Pharma	0.7	Mitsubishi Tanabe Pharma	0.3
Medicago	3.5		
Mitsubishi Tanabe Pharma Factory	1.0		

(4) Depreciation and Amortization Costs

[Billion yen]

	1H FY2019	1H FY2018	Increase (decrease)	FY2018
Property, plant and equipment	5.6	3.6	2.0	7.1
Intangible assets (except for Intangible assets with products)	0.6	0.7	(0.1)	1.4
Intangible assets with products	1.2	1.4	(0.2)	2.9

3. Financial Data & Employee Numbers of Major Consolidated Subsidiaries

[Billion yen]

	Companies	Mitsubishi Tanabe Pharma Factory Ltd.	Mitsubishi Tanabe Pharma Holdings America, Inc.	Medicago, Inc.	NeuroDerm Ltd.	Tianjin Tanabe Seiyaku Co., Ltd.	Mitsubishi Tanabe Pharma Korea Co., Ltd.
	1H FY2019	14.1	14.9	0.0	-	2.9	3.1
Revenue	FY2018	26.3	33.9	0.6	-	5.8	6.4
	1H FY2018	13.3	16.7	0.3	-	2.9	3.1
	1H FY2019	1.2	0.9	(6.3)	(4.9)	0.5	0.3
Operating profit	FY2018	1.6	3.4	(13.6)	(7.7)	0.4	0.5
	1H FY2018	0.5	0.9	(4.9)	(3.6)	0.2	0.3
	1H FY2019	0.8	0.6	(6.3)	(5.0)	0.2	0.3
Net profit	FY2018	1.2	2.9	(13.7)	(7.7)	0.1	0.4
	1H FY2018	0.3	0.7	(4.9)	(3.5)	0.1	0.2
	1H FY2019	0.5	2.0	6.4	4.9	0.0	-
R&D expenses	FY2018	0.8	4.0	14.2	7.7	0.0	-
	1H FY2018	0.5	2.0	5.3	3.5	0.0	-
Depreciation of	1H FY2019	1.2	0.2	0.3	0.0	0.0	0.0
property, plant and	FY2018	2.4	0.1	0.5	0.0	0.2	0.0
equipment	1H FY2018	1.1	0.0	0.2	0.0	0.1	0.0
	End of 1H FY2019	45.9	42.0	39.9	133.0	5.9	4.3
Total assets	End of FY2018	45.1	54.0	38.9	137.5	5.6	4.7
	End of 1H FY2018	44.6	47.1	38.0	141.0	5.6	4.4
	End of 1H FY2019	39.2	23.0	27.7	99.0	3.3	3.3
Total equity	End of FY2018	39.0	22.9	26.3	103.5	3.2	3.5
	End of 1H FY2018	38.2	21.2	27.1	105.6	3.3	3.5
	End of 1H FY2019	603	263	451	114	523	149
Number of employees	End of FY2018	633	265	421	100	508	143
. ,	End of 1H FY2018	652	269	365	88	517	143

Note: Prior to elimination of internal transactions

3 Forecasts for FY2019 Ending March 31, 2020

(Amounts less than ¥ 100 million are rounded off)

(1) Consolidate Forecasts of Profit and Loss

[Billion yen]

						[Billion yen]
		FY2019 forecasts*1		Comparison to previous fiscal year		Notes [Y-on-Y
		F12019 TOTECASES	FY2018 actual	Increase (decrease)	Change %	Comparison]
Rev	venue	376.0	424.7	(48.7)	(11.5)	See p9 "(2) Sales Forecasts for Main Products"
	Domestic	308.3	307.7	0.6	0.2	. roudets
	Overseas	67.6	117.0	(49.3)	(42.2)	
	Overseas sales ratio	18.0%	27.6%			
Cos	st of sales	178.5	180.6	(2.1)	(1.2)	Increase due to product mix change
	Sales cost ratio	47.5%	42.5%			
Gro	oss profit	197.5	244.1	(46.6)	(19.1)	
S	G&A expenses	99.0	98.2	0.7	0.8	
	% of revenue	26.3%	23.1%			
R	R&D expenses	85.5	86.5	(1.0)	(1.2)	
	% of revenue	22.7%	20.4%			
a	mortization of intangible ssets associated with roducts	2.5	2.9	(0.4)	(14.8)	
	Other income expense) ^{*2}	(0.5)	(0.5)	0.0	-	
Cor	re operating profit	10.0	55.8	(45.8)	(82.1)	
Noi	n-recurring items ^{*2}	1.5	(5.5)	7.0	-	
Ор	erating profit	11.5	50.3	(38.8)	(77.1)	
Pro	fit before tax	12.0	50.4	(38.4)	(76.2)	
Net	profit for the period	4.0	32.2	(28.2)	(87.6)	
	profit attributable to ners of the Company	5.0	37.3	(32.3)	(86.6)	
Tot	al labor cost	74.5	74.1	0.3	0.5	

st1: The Company announced full year forecasts on May 10, 2019

Exchange rate

[Yen]

	FY2019 planned	FY2018 average
USD	110.00	111.07
CAD	85.00	84.47
EUR	125.00	128.26

^{*2:} Brackets indicate expense and loss

	*1	Comparison to previous fiscal year		
	FY2019 forecasts*1	FY2018 actual	Increase (decrease)	Change %
Domestic ethical drugs	298.1	298.7	(0.6)	(0.2)
Remicade	51.5	58.8	(7.2)	(12.3)
Simponi	42.2	37.4	4.7	12.6
Stelara	21.6	15.2	6.4	42.4
Tenelia	15.0	15.2	(0.1)	(0.9)
Canaglu	10.4	6.7	3.6	54.4
Canalia	7.2	7.4	(0.2)	(2.7)
Kremezin	8.3	6.6	1.6	24.7
Lexapro	14.7	14.0	0.7	5.6
Ceredist	8.5	8.9	(0.4)	(4.5)
Rupafin	7.5	3.4	4.0	118.7
Talion	5.4	6.4	(0.9)	(14.6)
Vaccine [BIKEN products]	36.2	37.3	(1.0)	(2.9)
Influenza	10.7	10.2	0.5	5.0
Tetrabik	10.0	8.5	1.4	17.3
Varicella vaccine	5.1	5.1	0.0	1.7
Overseas ethical drugs	49.6	55.1	(5.4)	(9.9)
Radicava	22.0	27.0	(5.0)	(18.6)
Herbesser	7.2	6.8	0.3	5.3
Simponi	2.0	2.0	0.0	4.8
Argatroban	1.7	1.9	(0.1)	(6.6)
Tanatril	1.6	1.5	0.1	7.4
Royalty revenue, etc.	19.2	63.1	(43.8)	(69.5)
Royalty from Gilenya*2	Undisclosed	49.7	-	-
Royalty from INVOKANA	Undisclosed	10.5	-	-
OTC products	4.3	3.7	0.5	14.6
Others*3	4.6	3.9	0.6	16.2
tal sales revenue	376.0	424.7	(48.7)	(11.5)

^{*1:} The Company announced forecasts for FY2019 on May 10, 2019. As for the impact of the NHI drug price revision accompanying the consumption tax increase in October 2019, it is reflected in the overall sales forecast announced on May 10. Therefore, it remains the same. On the other hand, the individual domestic product forecasts have been changed to reflect the impact of the NHI drug price revision this time.

^{*2:} Mitsubishi Tanabe Pharma (MTPC) is currently in the arbitration proceedings with Novartis, and among the Gilenya Royalty amounts that MTPC is going to receive from Novartis, MTPC has decided not to recognize some of those amounts as our revenue because such payments do not satisfy one of the requirements under IFRS 15. The same accounting treatment will be continued during the period of the arbitration proceedings. Regardless of the disclosed amounts, MTPC maintains it is entitled to receive the full royalty amounts due according to the 1997 License Agreement with Novartis, and MTPC will rigorously pursue its rights in the arbitration.

^{*3:} Contracted manufacturing products of other companies.

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

[Billion yen]

	FY2019 forecasts	FY2018 actual	Increase (decrease)	Change %
Investment in property, plant and equipment / occurring basis	22.3	6.8	15.4	224.9
Investment for information systems / occurring basis	1.7	1.7	(0.0)	(3.8)

[Billion yen]

Major investment in property, plant and equipment in FY2019		Major investment for information systems in FY2019		
Production facilities	19.7	Production related systems	0.3	
[Medicago, Inc.]	[13.5]	R&D related systems	0.4	
[Mitsubishi Tanabe Pharma Factory Ltd.]	[2.5]	Others	1.0	
Facilities & equipment for R&D	1.7			
Others	0.9			

(4) Forecasts for Depreciation and Amortization Costs

[Billion yen]

- ·				
	FY2019 forecasts	FY2018 actual	Increase (decrease)	Change %
Property, plant and equipment*	11.0	7.1	3.8	53.9
Intangible assets (except for intangible assets with products)	1.5	1.4	0.0	3.6
Intangible assets with products	2.5	2.9	(0.4)	(14.8)

^{*:} Including the impact for the application of IFRS 16

4 Five-Year Financial Data

Japan GAAP (Amounts less than ¥100 million are rounded)

(1) Profit and Loss

[Billion yen]

	FY2014	FY2015
Net sales	415.1	431.7
Cost of sales	169.6	155.8
Gross operation profit	245.5	275.9
SG&A expenses	178.4	181.0
R&D expenses	69.6	75.3
Operating income	67.1	94.9
Ordinary income	67.7	94.8
Extraordinary income	13.7	14.1
Extraordinary loss	18.6	24.6
Net income attributable to shareholders of the Company	39.5	56.4

(2) Balance Sheet

[Billion yen]

	End of FY2014	End of FY2015
Total assets	929.3	930.2
Current assets	603.6	657.3
Fixed assets	325.7	273.0
Total liabilities	128.9	113.5
Current liabilities	105.4	91.3
Fixed liabilities	23.5	22.2
Net assets	800.4	816.7

(3) Other Financial Data

[Billion yen]

	FY2014	FY2015
Cash flows from operating activities	68.2	65.2
Cash flows from investing activities	(59.8)	(26.6)
Cash flows from financing activities	(21.9)	(22.2)
Investments in property, plant and equipment	15.7	11.2
Investments for development of information systems	1.6	0.9
Depreciation costs	9.0	8.8
Equity ratio (%)	84.9	86.6
ROE (%)	5.1	7.1
Net income per share (¥)	70.41	100.60
Net assets per share (¥)	1,406.41	1,436.63

IFRS (Amounts less than ¥100 million are rounded off)

(1) Profit and Loss

[Billion yen]

	FY2015	FY2016	FY2017	FY2018	1H FY2019	FY2019 forecasts
Revenues	425.7	423.9	433.8	424.7	188.1	376.0
Cost of sales	155.8	164.3	169.7	180.6	88.5	178.5
Gross operation profit	269.9	259.5	264.1	244.1	99.6	197.5
SG&A expenses	96.3	98.3	104.0	98.2	46.8	99.0
R&D expenses	64.6	64.7	79.0	86.5	39.7	85.5
Core operating profit	106.9	94.5	78.5	55.8	11.6	10.0
Operating income	81.8	94.0	77.2	50.3	12.5	11.5
Profit before income taxes	83.2	96.0	78.7	50.4	12.1	12.0
Net profit for the period	57.0	68.9	53.9	32.2	6.1	4.0
Net profit attributable to owners of the Company	59.3	71.2	57.9	37.3	8.3	5.0

(2) Balance Sheet

[Billion yen]

	End of FY2015	End of FY2016	End of FY2017	End of FY2018	End of 1H FY2019
Assets	958.4	984.5	1,048.4	1,056.2	1,055.8
Non-current assets	308.2	300.7	462.9	467.8	465.7
Current assets	650.1	683.7	585.5	588.4	590.1
Liabilities	132.1	113.1	153.6	145.9	162.2
Non-current liabilities	33.2	24.7	55.4	54.2	73.4
Current liabilities	98.9	88.4	98.1	91.6	88.8
Equity	826.3	871.4	894.8	910.3	893.5

(3) Other Financial Data

[Billion yen]

	FY2015	FY2016	FY2017	FY2018	1H FY2019	FY2019 forecasts
Cash flows from operating activities	80.8	59.7	66.9	41.4	19.4	-
Cash flows from investing activities	(42.2)	(10.5)	(19.1)	(31.2)	(21.9)	-
Cash flows from financing activities	(22.2)	(24.4)	(32.5)	(25.8)	(19.7)	-
Investments in property, plant and equipment	11.2	12.6	4.4	6.8	5.7	22.3
Investments for development of information systems	0.9	1.8	1.6	1.7	0.5	1.7
Depreciation and Amortization Costs	10.3	10.4	11.5	11.5	7.5	15.0
Ratio of equity attributable to owners of the Company to total assets [%]	85.1	87.4	84.2	85.0	83.5	-
ROE [%]	7.4	8.5	6.6	4.2	1.9	-
Basic earnings per share [¥]	105.72	127.03	103.35	66.64	14.83	8.92
Equity attributable to owners of the Company per share [¥]	1,453.71	1,533.91	1,574.26	1,600.64	1,571.99	-

(4) Number of Employees

	End of FY2014	End of FY2015	End of FY2016	End of FY2017	End of FY2018	End of 1H FY2019	Forecasts for end of FY2019
Consolidated	8,457	8,125	7,280	7,187	7,228	7,100	7,200
Non-consolidated	4,844	4,780	4,239	4,222	4,111	3,924	3,960

5 Quarterly Trend

(Amounts less than ¥ 100 million are rounded off)

(1) Profit and Loss

[Billion yen]

			FY2018				FY2019	[Billion yen]
	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.	Full year forecasts ^{*1}
Revenue	105.3	104.3	122.7	92.2	424.7	98.1	89.9	376.0
	24.8%	24.6%	28.9%	21.7%	100.0%	26.1%	23.9%	100.0%
Domestic	74.1	72.3	89.9	71.3	307.7	80.7	73.8	308.3
Domestic	24.1%	23.5%	29.2%	23.2%	100.0%	26.2%	24.0%	100.0%
Overseas	31.1	32.0	32.8	20.9	117.0	17.3	16.1	67.6
Overseas	26.6%	27.4%	28.1%	17.9%	100.0%	25.6%	23.8%	100.0%
Cost of sales	42.3	43.7	53.0	41.4	180.6	44.7	43.7	178.5
Sales cost ratio	40.2%	42.0%	43.2%	44.9%	42.5%	45.6%	48.6%	47.5%
Cross profit	63.0	60.5	69.7	50.8	244.1	53.3	46.2	197.5
Gross profit	25.8%	24.8%	28.6%	20.8%	100.0%	27.0%	23.4%	100.0%
SC&A aynansas	23.1	24.5	25.4	25.0	98.2	22.9	23.8	99.0
SG&A expenses	23.6%	25.0%	25.9%	25.5%	100.0%	23.2%	24.1%	100.0%
R&D expenses	19.6	19.9	22.3	24.6	86.5	19.9	19.8	85.5
R&D expenses	22.7%	23.0%	25.8%	28.5%	100.0%	23.3%	23.2%	100.0%
Amortization of intangible assets associated with	0.7	0.7	0.7	0.7	2.9	0.6	0.6	2.5
products	25.0%	25.0%	25.0%	25.0%	100.0%	25.9%	24.3%	100.0%
Other income (expense)*2	(0.1)	(0.1)	(0.0)	(0.1)	(0.5)	(0.0)	0.0	(0.5) -
Care encusting puefit	19.3	15.1	21.0	0.2	55.8	9.7	1.9	10.0
Core operating profit	34.6%	27.2%	37.7%	0.5%	100.0%	97.5%	19.5%	100.0%
On austing mustic	19.3	15.1	21.9	(6.1)	50.3	9.6	2.9	11.5
Operating profit	38.4%	30.2%	43.6%	(12.2%)	100.0%	83.7%	25.6%	100.0%
Profit before tax	19.7	15.0	21.7	(6.1)	50.4	9.2	2.9	12.0
Profit before tax	39.1%	29.9%	43.1%	(12.1%)	100.0%	76.7%	24.5%	100.0%
Net profit attributable to	13.9	11.0	16.4	(4.0)	37.3	6.8	1.4	5.0
owners of the Company	37.4%	29.5%	44.1%	(11.0%)	100.0%	137.6%	28.7%	100.0%

Note: The each figure (excluding "cost of sales") in the lower displays the progress rate.

^{*1:} The Company announced full year forecasts on May 10, 2019

^{*2:} Brackets indicate expense and loss

(2) Sales Revenue of Main Products

[Billion yen]

				FY2018				FY2019			
				Q1	Q2	Q3	Q4	Full year	Q1	Q2	Full year
				Apr. to Jun.	Jul. to Sep.	Oct. to Dec.	Jan. to Mar.	actual	Apr. to Jun.	Jul. to Sep.	forecasts*1
	Do	mestic ethica	al drugs	71.6	69.9	87.6	69.5	298.7	78.1	71.0	298.1
			·	24.0%	23.4%	29.3%	23.3%	100.0%	26.2%	23.8%	100.0%
		Remicade		15.1	14.8	16.0	12.8	58.8	14.4	13.1	51.5
	-			25.7% 9.0	25.2% 9.5	27.2% 10.2	21.9% 8.7	100.0% 37.4	28.0% 10.5	25.5% 9.9	100.0% 42.2
		Simponi		24.0%	25.4%	27.3%	23.3%	100.0%	24.9%	23.5%	100.0%
		Stelara		0.2	4.5	5.6	4.7	15.2	6.1	6.3	21.6
	_	Stelara		1.4%	30.0%	37.3%	31.3%	100.0%	28.6%	29.5%	100.0%
		Tenelia		4.4	2.7	3.9	4.0	15.2	4.7	3.3	15.0
	-			29.5% 1.4	18.0% 1.5	25.8% 1.9	26.7%	100.0% 6.7	31.2% 2.1	22.5%	100.0%
		Canaglu		22.2%	22.9%	29.4%	25.5%	100.0%	20.7%	18.6%	100.0%
		Canalia		1.4	1.6	2.3	2.0	7.4	2.2	1.5	7.2
	_	Carialia		19.1%	22.3%	31.1%	27.5%	100.0%	30.5%	21.5%	100.0%
		Kremezin		1.7	1.6	1.8	1.4	6.6	1.7	1.6	8.3
	-			25.5% 3.4	24.9% 3.4	27.6% 3.8	22.0% 3.2	100.0% 14.0	21.0% 3.8	19.3% 3.5	100.0% 14.7
		Lexapro		24.4%	24.4%	27.8%	23.4%	100.0%	26.3%	24.1%	100.0%
		Cauadiat		2.4	2.2	2.4	1.8	8.9	2.1	1.7	8.5
		Ceredist		27.7%	24.6%	27.4%	20.3%	100.0%	25.4%	19.9%	100.0%
		Rupafin		0.1	0.2	0.5	2.4	3.4	1.2	1.2	7.5
	-			5.0%	6.1%	16.7%	72.2%	100.0%	16.6%	16.6%	100.0%
		Talion		1.4 22.3%	1.1 17.9%	1.5 24.7%	2.2 35.1%	6.4 100.0%	1.2 22.6%	0.9 17.2%	5.4 100.0%
	-	Vaccines		8.8	6.7	14.8	6.8	37.3	7.3	8.4	36.2
		BIKEN produ	uctel								
		[DIKEN produ	uctsj	23.7%	18.1%	39.9%	18.4%	100.0%	20.2%	23.3%	100.0%
		Influenza		(0.1)	1.0	8.5	0.7	10.2	(0.0)	1.8	10.7
				(1.1%)	10.6%	83.4%	7.0%	100.0%	(0.2%)	16.8%	100.0%
		Tetrabik		2.2	1.9	2.3	2.0	8.5	2.3	2.2	10.0
				25.7%	23.0%	26.9%	24.4%	100.0%	23.6%	21.9%	100.0%
		Varicella	vaccine	1.4	1.2	1.3	1.1	5.1	1.2	1.2	5.1
				27.7%	23.8%	25.7%	22.9%	100.0%	24.8%	23.4%	100.0%
	Ov	erseas ethica	al drugs	12.9	14.5	14.4	13.1	55.1	12.5	12.2	49.6
	١,		3	23.5%	26.3%	26.3%	23.9%	100.0%	25.4%	24.7%	100.0%
		Radicava		6.4	7.4	6.7	6.4	27.0	6.1	5.5	22.0
	-			23.7%	27.7%	25.0%	23.7%	100.0%	27.8%	25.0%	100.0%
		Herbesser		1.6 24.4%	1.6 23.9%	1.7 24.9%	1.8 26.7%	6.8 100.0%	1.7 24.7%	1.7 23.4%	7.2 100.0%
	-			0.4	0.5	0.4	0.5	2.0	0.5	0.5	2.0
		Simponi		24.2%	25.0%	24.8%	26.1%	100.0%	24.5%	25.2%	100.0%
		Argatroban		0.5	0.4	0.5	0.3	1.9	0.4	0.4	1.7
		, a gata obail		29.4%	24.5%	26.7%	19.3%	100.0%	27.0%	27.4%	100.0%
		Tanatril		0.3 23.7%	0.4	0.4 27.1%	0.2	1.5	0.3	0.3	1.6 100.0%
					30.7%		18.5%	100.0%	21.7% 5.0	21.7% 4.1	100.0%
	Ro	yalty revenu	ie, etc.	18.5	17.7	18.6	8.1	63.1			
	١,	•	•	29.3%	28.2%	29.6%	12.9%	100.0%	26.2%	21.6%	100.0%
		Royalty from	Gilenva*2	15.3	14.5	14.7	5.0	49.7	1.6	1.5	Undisclosed
	_	rioyaley iroini	Circitya	30.9%	29.3%	29.6%	10.2%	100.0%	-	-	-
		Royalty from	INVOKANA	2.4	2.4	3.2	2.3	10.5	2.0	2.0	Undisclosed
		,,	2111 010	23.6%	23.4%	30.5%	22.5%	100.0%	-	-	-
	ОТ	C products		1.2	0.9	1.0	0.5	3.7	1.2	1.0	4.3
	O1	C products		31.9%	26.4%	26.8%	14.9%	100.0%	30.0%	24.5%	100.0%
	O+۱	ners*3		1.0	1.1	0.9	0.8	3.9	1.0	1.4	4.6
	Otl	ICI 2		25.9%	28.8%	22.9%	22.4%	100.0%	22.0%	32.3%	100.0%
Ta	tal.	salos rover	110	105.3	104.3	122.7	92.2	424.7	98.1	89.9	376.0
10	ldl S	sales reven	ue	24.8%	24.6%	28.9%	21.7%	100.0%	26.1%	23.9%	100.0%
		1.6		vs the progress i							

Note: The each figure in the lower displays the progress rate.

^{*1:} The Company announced forecasts for FY2019 on May 10, 2019. As for the impact of the NHI drug price revision accompanying the consumption tax increase in October 2019, it is reflected in the overall sales forecast announced on May 10. Therefore, it remains the same. On the other hand, the individual domestic product forecasts have been changed to reflect the impact of the NHI drug price revision this time.

^{*2:} Mitsubishi Tanabe Pharma (MTPC) is currently in the arbitration proceedings with Novartis, and among the Gilenya Royalty amounts that MTPC is going to receive from Novartis, MTPC has decided not to recognize some of those amounts as our revenue because such payments do not satisfy one of the requirements under IFRS 15. The same accounting treatment will be continued during the period of the arbitration proceedings. Regardless of the disclosed amounts, MTPC maintains it is entitled to receive the full royalty amounts due according to the 1997 License Agreement with Novartis, and MTPC will rigorously pursue its rights in the arbitration.

^{*3:} Contracted manufacturing products of other companies.

6 State of New Product Development (as of October 25, 2019)

i. Immuno-inflammation

Development code Product name (Generic name)	Category (Indications)	Region Stage	Origin/licensee	
MT-5547 (Fasinumab)	Fully human anti-NGF monoclonal antibody (Osteoarthritis)	Japan Phase 2/3	Licensed from Regeneron (US)	
MT-1303	S1P receptor functional antagonist (Multiple sclerosis)	Europe Phase 2	In-house	
(Amiselimod)	(Crohn's disease)	Japan Phase 2	III House	
MT-7117	Dermatologicals, etc. (Erythropoietic protoporphyria)	Global Phase 2	In-house	
MT-2990	Fully human anti-interleukin-33 (IL-33) monoclonal antibody (Endometriosis)	Global Phase 2	In-house	
	(Seasonal Allergic Rhinitis)	Phase 1		

ii. Diabetes and kidney

Development code Product name (Generic name)	Category (Indications)	Region Stage	Origin/licensee
TA-7284	SGLT2 inhibitor (Type 2 diabetes mellitus)	Asia Filed	In-house
Canaglu/INVOKANA (Canagliflozin)	(Diabetic nephropathy)	Europe Filed (Jul. 2019)	Licensed to Janssen Pharmaceuticals (US)
(curiagiiioziii)	(Біавейс першораспу)	Japan Phase 3	In-house
MP-513		Asia Filed	
Tenelia (Teneligliptin)	DPP-4 inhibitor (Type 2 diabetes mellitus)	China Filed (Sep. 2019)	In-house
(Tellenginpany)		Europe Phase 2	
MT-6548 (Vadadustat)	Hypoxia inducible factor prolyl hydroxylase inhibitor (Renal anemia)	Japan Filed (Jul. 2019)	Licensed from Akebia (US)
	Selective mineralocorticoid receptor antagonist	Europe Phase 2	
MT-3995 (Apararenone)	(Diabetic nephropathy)	Japan Phase 2	In-house
	(Non-alcoholic steatohepatitis: NASH)	Japan Phase 2	

 $\ensuremath{\mathbb{X}}$ Asia: excluding Japan and China

iii. Central nervous system

Development code Product name (Generic name)	Category (Indications)	Region Stage	Origin/licensee
MCI-186 Radicut/Radicava (Edaravone)	Free radical scavenger (Amyotrophic lateral sclerosis: ALS)	Asia Filed	In-house
MP-214 (Cariprazine)	Dopamine D3/D2 receptor partial agonist (Schizophrenia)	Asia Filed	Licensed from Gedeon Richter (Hungary)
MT-210 (Roluperidone)	5-HT2A/Sigma 2 receptor antagonist (Schizophrenia)	US, Europe Phase 3	Licensed to Minerva Neurosciences (US)
MT-5199 (Valbenazine)	Vesicular monoamine transporter type 2 inhibitor (Tardive dyskinesia)	Japan Phase 2/3	Licensed from Neurocrine Biosciences (US)
ND0612 (Levodopa/Carbidopa)	Continuous SC pump Global (Parkinson's disease) Phase 3		In-house
MT-0551 (Inebilizumab)	(Neuromyelitis ontica spectrum disorder:		Licensed from Viela Bio (US)
MT-8554	TRPM8 antagonist (Painful diabetic peripheral neuropathy)	Europe Phase 2	· In-house
(Elismetrep)	(Vasomotor symptoms associated with menopause)	Global Phase 2	In nouse
ND0701 (Apomorphine)	Continuous SC pump (Parkinson's disease)	Phase 1	In-house
MT-1186 (Edaravone)	AT-1186 Free radical scavenger (Amyotrophic lateral sclerosis: ALS / Oral		In-house
MT-6345	Nervous system	Phase 1	Co-developed with Ube Industries (Japan)
MT-3921	Anti-RGMa antibody (Spinal cord injury)	Phase 1	Co-developed with Osaka University (Japan)

iv. Vaccines

Development code Product name (Generic name)	Category (Indications)	Region Stage	Origin/licensee
MT-2355	Combined vaccine (Prophylaxis of pertussis, diphtheria, tetanus, poliomyelitis and prophylaxis of Hib infection in infants)	Japan Phase 3	Co-developed with The Research Foundation for Microbial Diseases of Osaka University (Japan)
MT-2271	Plant-based VLP vaccine (Prophylaxis of seasonal influenza/adults)	Canada Filed (Sep. 2019) US, Europe Phase 3	Medicago product (Canada)
	(Prophylaxis of seasonal influenza/elderly)	US, Europe Phase 3	(3, 3, 3,
MT-8972	Plant-based VLP vaccine (Prophylaxis of H5N1 influenza)	Canada Phase 2	Medicago product (Canada)
MT-7529	Plant-based VLP vaccine (Prophylaxis of H7N9 influenza)	Phase 1	Medicago product (Canada)
MT-5625	Plant-hased VI P vaccine		Medicago product (Canada)

 $\ensuremath{\,\times\,}$ Asia: excluding Japan and China

v. Others

Development code Product name (Generic name)	Category (Indications)	Region Stage	Origin/licensee
TAU-284 Talion (Bepotastine)	Selective histamine H1 receptor antagonist, anti-allergic agent (Allergic rhinitis, Urticaria)	Asia Filed	Licensed from Ube Industries (Japan)
MT-4580 Orkedia (Evocalcet)	Ca sensing receptor agonist (Hypercalcemia in patients with parathyroid carcinoma or primary hyperparathyroidism)	Japan Filed (Apr. 2019)	Licensed to Kyowa Kirin (Japan)
(Lvocaicer)	(Secondary Hyperparathyroidism)	China, Asia Phase 3	
MT-4129	Cardiovascular system, etc.	Phase 1	In-house
MT-8633/TR1801-ADC	Anti-c-Met ADC* (Solid tumor)	Phase 1	In-house Collaborate with Open Innovation Partners (Japan)

*Antibody drug conjugate

 ${\it \divideontimes}$ Asia: excluding Japan and China

Changes Since Previous Announcement

Development code Product name (Generic name)	Category (Indications)	Previous Announcement	As of Oct 25, 2019	Origin / licensee
MCI-186 Radicut/Radicava (Edaravone)	Free radical scavenger (Amyotrophic lateral sclerosis: ALS)	China Filed (Apr. 2019)	China Approved (Jul. 2019)	In-house
MP-214 (Cariprazine)	Dopamine D3/D2 receptor partial agonist (Schizophrenia)	Asia Filed	Thailand Approved (Jul. 2019)	Licensed from Gedeon Richter (Hungary)
TA-7284 Canaglu/INVOKANA	SGLT2 inhibitor	US Filed (Mar. 2019)	US Approved (Sep. 2019)	Licensed to Janssen
(Canagliflozin)	(Diabetic nephropathy)	None	Europe Filed (Jul. 2019)	Pharmaceuticals (US)
MP-513 Tenelia (Teneligliptin)	DPP-4 inhibitor (Type 2 diabetes mellitus)	China Phase 3	China Filed (Sep. 2019)	In-house
MT-2271	Plant-based VLP vaccine (Prophylaxis of seasonal influenza/adults)	None	Canada Filed (Sep. 2019)	Medicago product (Canada)
ND0612 (Levodopa/Carbidopa)	Continuous SC pump (Parkinson's disease)	Global Phase 2	Global Phase 3	In-house
MT-0551 (Inebilizumab)	Humanized anti-CD19 monoclonal antibody (Neuromyelitis optica spectrum disorder: NMOSD)	None	Japan, Asia Phase 3	Licensed from Viela Bio (US)
MT-4580 Orkedia (Evocalcet)	Ca sensing receptor agonist (Secondary Hyperparathyroidism)	None	China, Asia Phase 3	Licensed to Kyowa Kirin (Japan)
MT-8633/TR1801-ADC	Anti-c-Met ADC* (Solid tumor)	None	Phase 1	In-house Collaborate with Open Innovation Partners (Japan)

*Antibody drug conjugate $\mbox{$\%$}$ Asia: excluding Japan and China

7 Others

1. Subsidiaries and Affiliated Companies

(1) Number of Subsidiaries and Affiliated Companies

	End of 1H FY2019	End of FY2018	Increase (Decrease)	Notes
Consolidated subsidiaries	33	34	(1)	Decrease: Tanabe Seiyaku Yoshiki Factory Co., Ltd.
Associates and joint ventures	1	2	(1)	Decrease: Synthelabo-Tanabe Chimie S.A.
Total	34	36	(2)	

(2) Consolidated Subsidiaries

[As of September 30, 2019]

(-)	2) Consolidated Substitutions 30, 2019						
	Company Name	Paid-in Capital	% Voting Control [% Indirect Ownership]		Settling Day	Description of Business	
1	Yoshitomiyakuhin Corporation	JPY 385 million	100.0	[-]	End of Mar.	Provision of information about pharmaceuticals	
2	Mitsubishi Tanabe Pharma Factory Ltd.	JPY 1,130 million	100.0	[-]	End of Mar.	Manufacture and sale of pharmaceuticals	
3	Mitsubishi Tanabe Pharma Provision Co., Ltd.	JPY 100 million	100.0	[-]	End of Mar.	Handling pharmacy information and accounting, general affairs and human resources management etc.	
4	Tanabe Palm Service Co., Ltd.	JPY 10 million	100.0	[100.0]	End of Mar.	Servicing office support, in-house mail and printing.	
5	Stelic Institute & Co., Inc.	JPY 1 million	100.0	[100.0]	End of Mar.	R&D of pharmaceuticals	
6	Mitsubishi Tanabe Pharma Holdings America, Inc.	USD 167	100.0	[-]	End of Mar.	Managing the US business	
7	Mitsubishi Tanabe Pharma Development America, Inc.	USD 200	100.0	[100.0]	End of Mar.	R&D of pharmaceuticals	
8	Mitsubishi Tanabe Pharma America, Inc.	USD 100	100.0	[100.0]	End of Mar.	Sale of pharmaceuticals	
9	MP Healthcare Venture Management, Inc.	USD 100	100.0	[100.0]	End of Mar.	Investments in bio-ventures	
10	Tanabe Research Laboratories U.S.A., Inc.	USD 3 Mill.	100.0	[100.0]	End of Mar.	R&D of pharmaceuticals	
11	Mitsubishi Tanabe Pharma Canada, Inc.	CAD 4 Mill.	100.0	[100.0]	End of Mar.	Sale of pharmaceuticals	
12	MTPC Holdings Canada Inc.	CAD 641.3 Mill.	100.0	[-]	End of Mar.	Investments in Medicago Group	
13	Medicago Inc.	CAD 851.0 Mill.	64.7	[63.0]	End of Mar.	Manufacture and sale of vaccines	
14	Medicago USA Inc.	USD 99	64.7	[64.7]	End of Mar.	Manufacture of vaccines	
15	Medicago R&D Inc.	CAD 500	64.7	[64.7]	End of Mar.	R&D of vaccines	
16	Mitsubishi Tanabe Pharma Development (Beijing) Co., Ltd.	USD 1 Mill.	100.0	[-]	End of Dec.	R&D of pharmaceuticals	
17	Tianjin Tanabe Seiyaku Co., Ltd.	USD 16.2 Mill.	75.4	[-]	End of Dec.	Manufacture and sale of pharmaceuticals	
18	Taiwan Tanabe Seiyaku Co., Ltd.	TWD 90 Mill.	65.0	[-]	End of Mar.	Manufacture and sale of pharmaceuticals	
19	Tai Tien Pharmaceuticals Co., Ltd.	TWD 20 Mill.	65.0	[-]	End of Mar.	Sale of pharmaceuticals	
20	P.T. Mitsubishi Tanabe Pharma Indonesia	USD 2.5 Mill.	99.6	[-]	End of Mar.	Manufacture and sale of pharmaceuticals	
21	Mitsubishi Tanabe Pharma Singapore Pte. Ltd.	SGD 2 Mill.	100.0	[-]	End of Mar.	Managing the ASEAN business	
22	Mitsubishi Tanabe Pharma Malaysia Sdn. Bhd.	MYR 5 Mill.	100.0	[100.0]	End of Mar.	Sale of pharmaceuticals	
23	Mitsubishi Tanabe Pharma (Thailand) Co., Ltd.	THB 103 Mill.	100.0	[2.0]	End of Mar.	Sale of pharmaceuticals	
24	Mitsubishi Tanabe Pharma Korea Co., Ltd.	KRW 2,100 Mill.	100.0	[-]	End of Mar.	Manufacture and sale of pharmaceuticals	
25	NeuroDerm Ltd.	USD 58,000	100.0	[-]	End of Mar.	R&D of pharmaceuticals	
26	Mitsubishi Tanabe Pharma Europe Ltd.	GBP 4.6 Mill.	100.0	[-]	End of Mar.	R&D of pharmaceuticals	
27	Mitsubishi Tanabe Pharma GmbH	EUR 25,000	100.0	[100.0]	End of Mar.	Sale of pharmaceuticals	

Note: Aside from the above, The Company own 5 consolidated subsidiaries. Among them, 2 companies are under the liquidation and 1 company is a dormant company. Besides, the executive compensation BIP Trust is included as one of the consolidated subsidiaries.

(3) Associates and Joint Ventures

[As of September 30, 2019]

	Company Name	Paid-in Capital	% Voting Control [% Indirect Ownership]	Settling Day	Description of Business
1	BIKEN Co., Ltd.	JPY 100 million	33.4 [-]	Fnd of Mar	Manufacture and sale of biological products including vaccines

2. Status of Shareholders

(1) Number of Outstanding Shares

. ,	End of September, 2019	End of March, 2019
Issued	561,417,916	561,417,916
The company's own shares at the end of the period st	631,915	640,305
Number of shares outstanding at the end of the period	560,786,001	560,777,611
Average number of the company's own share in the period	637,599	641,042
Average number of shares outstanding in the period	560,780,317	560,776,874

^{*}The Company introduces the executive compensation BIP Trust. The shares that the trust account holds are included in treasury shares (200,279 shares at the end of September 2019, compared to 208,655 shares at the end of March 2019).

(2) Status of Major Shareholders

		End of September, 2019			End of March, 2019		
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	56.39	1	316,320	56.39	
2	The Master Trust of Japan, Ltd.	26,210	4.67	2	26,596	4.74	
3	Japan Trustee Services Bank, Ltd.	13,226	2.36	3	14,679	2.62	
4	Nippon Life Insurance Company	12,065	2.15	4	12,065	2.15	
5	STATE STREET BANK WEST CLIENT-TREATY 505234	6,596	1.18	5	7,826	1.40	
6	JPMorgan Securities Japan	5,888	1.05	14	3,410	0.61	
7	Japan Trustee Services Bank, Ltd. (Trust Account 9)	5,648	1.01	6	4,627	0.82	
8	STATE STREET BANK AND TRUST COMPANY 505103	4,723	0.84	9	3,928	0.70	
9	Japan Trustee Services Bank, Ltd. (Trust Account 5)	4,645	0.83	7	4,113	0.73	
10	STATE STREET BANK AND TRUST COMPANY 505225	4,010	0.71	8	4,029	0.72	

(3) Ownership and Distribution of Shares

	End of September, 2019			End of March, 2019			
_	Number of Shareholders	Number of Shares (Thousands)	Percentage of Total %	Number of Shareholders	Number of Shares (Thousands)	Percentage of Total %	
Financial institutions	64	89,424	15.93	74	90,932	16.20	
Foreign corporations and others	612	97,311	17.34	624	101,801	18.14	
Individuals and others*	28,009	32,699	5.83	24,964	29,762	5.30	
Other corporations	293	329,666	58.73	285	330,056	58.80	
Securities firms	39	12,201	2.17	44	8,754	1.56	
Total	29,017	561,303	100.00	25,991	561,307	100.00	
Less than trading unit	-	114	-	-	110	-	

Note: The trading unit of the Company's stock is 100 shares

(4) Trend of Dividend and Stock Price

(Japan GAAP)	FY2014	FY2015
Dividends per share [yen]	42	46
Dividend payout ratio [%]	59.6	45.7
[prior to amortization of goodwill]	[47.6]	[38.8]
Stock price at the end of FY [yen]	2,062	1,957
Market capitalization [billion yen]	1,157.6	1,098.7

(IFRS)	FY2015	FY2016	FY2017	FY2018	1H FY2019	FY2019 Estimate
Dividends per share [yen]	46	52	66*	56	28	56
Dividend payout ratio [%]	43.5	40.9	63.9	84.0	-	628.1
Stock price at the end of FY [yen]	1,957	2,318	2,080	1,479	1,185	-
Market capitalization [billion yen]	1,098.7	1,301.4	1,167.7	830.3	665.3	-

^{*} The Company distributed a commemorative dividend of ¥10 to shareholders at the end of 1st half in FY2017 for celebrating its 10th anniversary

^{*} Individuals and Others include treasury stocks (431 thousands shares at the end of September, 2019 and 431 thousands shares at the end of March, 2019)

Reference

Major Ethical Drugs

Remicade (Infliximab)

Launch: May 2002

Category Anti-TNFa monoclonal antibody

Remicade is very fast-acting and its efficacy is sustained for eight weeks with a single administration. It has indications for the treatment of rheumatoid arthritis, Crohn's disease, Behcet's disease with refractory uveoretinitis, psoriasis, ankylosing spondylitis, ulcerative colitis, Entero-Behcet's disease, neuro-Behcet's disease, and vasculo-Behcet's disease and Kawasaki disease. Partial change in dosage and usage (increased dose) for psoriasis was approved in May 2016. And partial change in administration / dosage of a shortened administration interval for Crohn's disease was approved in May 2017. Origin: Janssen Biotech

Simponi (Golimumab)

Launch: Sep. 2011

Category Anti-TNFa monoclonal antibody

Simponi is a human anti-TNFa monoclonal antibody for the treatment of rheumatoid arthritis (including prevention of articular structural damage). It shows a long acting efficacy by subcutaneous injection once every four weeks. Additional indication of ulcerative colitis was approved in March 2017 by Janssen Pharmaceutical. Self injection for rheumatoid arthritis was approved in April 2018. Autoinjector was released in May 2019. Origin: Janssen Biotech

Stelara (Ustekinumab)

Launch:

Category Anti-IL12/23p40 monoclonal antibody Mar. 2011

Stelara is a human anti-IL12/23p40 monoclonal antibody. It shows a long acting efficacy by subcutaneous injection once every 12 weeks. (initial admin. is intravenous drip injection) Additional indication of Crohn's disease was approved in March 2017.

Mitsubishi Tanabe Pharma and Janssen Pharmaceutical jointly promote STELARA on indication of Crohn's disease in Japan from April 2017. For the indication of psoriasis, promotion is handled solely by Janssen Pharmaceutical. Origin: Janssen Biotech

Imusera (Fingolimod)

Launch:

Nov. 2011

Category Agent for treatment for multiple sclerosis (MS)

Imusera is a first-in-class drug that controls inflammation in the brain and spinal cord in MS. It inhibits the receptor function of sphingosine-1-phosphate (S1P) receptor on the lymphocyte, and prevents auto-aggressive lymphocytes from invading the central nervous system. It can be administered orally (once daily), thereby lowering the burden on patients with MS. It was discovered by Mitsubishi Tanabe Pharma and developed jointly by Mitsubishi Tanabe Pharma and Novartis Pharma in Japan. Mitsubishi Tanabe Pharma is marketing this product under the name Imusera, while Novartis Pharma is marketing it under the name Gilenva.

Tenelia (Teneligliptin)

Launch: Sep. 2012

Category

Selective DPP-IV inhibitor

-Agent for treatment of type2 diabetes mellitus-

Tenelia, which Mitsubishi Tanabe has created and developed, is the first DPP-4 inhibitor originating in Japan. 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.

Canaglu (Canagliflozin)

Launch: Sep. 2014

Category

SGLT2 Inhibitor

-Agent for treatment of type2 diabetes mellitus-

Canaglu which was discovered by Mitsubishi Tanabe Pharma is a treatment for type 2 diabetes mellitus. It inhibits SGLT2 (sodium glucose co-transporter 2) of kidneys, suppresses the reabsorption of glucose, promotes the excretion of excessive glucose into the urine, and as a result, lowers the blood glucose level. In Overseas markets, licensee Janssen Pharmaceuticals (US) received approval in the US, EU, Australia and more than 80 countries, and this drug is sold under the brand name Invokana.

Canalia

(Teneligliptin/Canagliflozin)

Launch: Sep. 2017

Category

Selective DPP-IV inhibitor/SGLT2 Inhibitor combination tablets -Agent for treatment of type2 diabetes mellitus-

Canalia is the first combination tablets containing DPP-4 inhibitor and SGLT2inhibitor in Japan,containing DPP-4 inhibitor "Tenelia" and SGLT2 inhibitor "Canaglu" which Mitsubishi Tanabe has created and developed. It expects that are long-term good control of blood glucose and improvement of adherence by reducing the number of taking medicine.

Kremezin

Dec. 1991

Category Agent for treatment of Chronic renal failure

Kremezin is an oral absorptive charcoal consisting of porous spherical activated carbon of high purity. It absorbs and excretes uremic toxins out of the body. Kremezin was introduced to the Japanese market in December 1991 as the first pharmaceuticals drug in the world for proactive treatment of chronic renai failure (progressive). In April 2011, the marketing rights were transferred from Daiichi Sankyo to MTPC.In January 2018, Kremezin tablets was released. Origin, Manufacturer and distributor: Kureha.

Lexapro (Escitalopram)

Launch: Aug. 2011

Category Selective serotonin reuptake inhibitor (SSRI)

Lexapro, a highly selective serotonin reuptake inhibitor (SSRI), has been globally approved in 101 countries and regions. It shows good efficacy and tolerability in patients with depressive disorder. Moreover, due to simple dosage and administration, it is expected to improve adherence of the treatment. Social anxiety disorder (SAD) was approved in November 2015.

Origin: H. Lundbeck A/S (Denmark), Manufacturer and distributor: Mochida Pharmaceutical Co., Ltd

Radicut / Radicava (Edaravone)

Launch: Jun. 2001

Category

Free radical scavenger (Cerebral neuroprotectant)

Radicut is the world's first brain protecting agent (free radical scavenger) 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.

It was designated as an orphan drug of amyotrophic lateral sclerosis (ALS) and approved for ALS in Japan in June 2015, followed by approval in Korea (December 2015), the United States (May 2017), Canada (Octorber 2018), Switzerland (January 2019) and China(July 2019).

Ceredist (Taltirelin)

Launch: Sep. 2000

Category Agent for treatment of spinocerebellar degeneration

Thyrotropin releasing hormone (TRH) was known to be effective against ataxia caused by spinocerebellar degeneration, but it was previously administered only through injection. Ceredist is the world's first oral TRH derivative drug by in-house development. An additional formulation, orally disintegrating tablets, was launched in October 2009.

Rupafin (Rupatadine)

Launch: Nov. 2017

Category Agent for treatment of allergic disorders

Rupafin has Anti-PAF activity and anti-Histamine activity. Histamine and PAF lead to early phase reaction and PAF leads to late phase reaction in allergic disorder, Rupafin suppresses PAF and histamine. It has been approved for the treatment of allergic rhinitis, urticaria and itch associated with skin diseases (eczema/dermatitis, cutaneous pruritus).

※PAF: platlet activating factor

Origin: Uriach(Spain), Manufacturer and distributor: Teikoku Seiyaku

Talion (Bepotastine)

Launch: Oct. 2000

Category

Selective histamine H1 receptor antagonist Agent for treatment of allergic disorders

Talion has rapid onset of anti-histamine (H1) effects and has been demonstrated to be effective for allergic rhinitis, urticaria, and pruritus accompanying dermatitis. It has minimal incidence of sedation. An additional formulation, orally disintegrating tablets was launched in July 2007. Pediatric indications (from seven to fifteen years old) was approved in May 2015.

Origin: Ube Industries

Influenza vaccine

Launch: Sep. 1972

Category Viral vaccines

It is for prevention of seasonal influenza. It was changed from trivalent vaccine to quadrivalent vaccine in 2015. Origin, Manufacturer and distributor: BIKEN (The Research Foundation for Microbial Diseases of Osaka University)

Tetrabik

Launch: Oct. 2012

Category Vaccine toxoid combined formulation

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 similar 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)

Varicella vaccine

Launch: Mar. 1987

Category Viral vaccines

It is for prevention of varicella and included in regular vaccination from 2014. An indication for prevention of shingles in people older than 50 was approved in

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

Mearubik

Launch:

Dec. 2005

Category Viral vaccines combined formulation

Mearubik is for prevention of measles and rubellathe. It is combination vaccine for measles and rubella, and children are able to receive both measles and rubella shot at a time with Mearubik, which is used at the 1st term and the 2nd term of its regular vaccination. By both reducing the number of injections and relieving physical pain on people to be vaccinated, It is expected to contribute enhancement of immunization rate for measles and rubella in Japan. Origin, Manufacturer and distributor: BIKEN (The Research Foundation for Microbial Diseases of Osaka University)

JEBIK V

Launch: Jun. 2009

Category Viral vaccines

JEBIK V is for prevention of Japanese encephalitisa. It is freeze-dried preparation containing inactivated Japanese encephalitis virus derived from Vero cells which were used in the manufacturing process as a host to increase the virus. It is used at the 1st term and 2nd term of the regular vaccination. It is expected to reduce the occurrence of ADEM by not using mice's brains in the manufacturing process.

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

News Releases

The major news releases after April, 2019 are as follows. Please refer to the Company's website for the details. (https://www.mt-pharma.co.jp/e/release/index.php)

Date	Contents
April 11, 2019	Notice Regarding Reorganization of Research, Production, and Technology Bases
April 15, 2019	For the ALS patients in the world, we hope to deliver Japan-originated ALS treatment—NMPA accepts our filing for Radicut to treat ALS in China
April 26, 2019	Mitsubishi Tanabe Pharma and Salix enter into a licensing agreement for MT-1303, a therapeutic agent for autoimmune diseases
May 29, 2019	Launch of SIMPONI Subcutaneous Injection 50mg Autoinjector
May 30, 2019	Withdrawal of Marketing Authorization Application of edaravone for ALS in the European Union
July 23, 2019	Submission of Vadadustat (MT-6548) New Drug Application in Japan for renal anemia
August 6, 2019	Commencement of R&D of Gene Therapy Product for Hemophilia B
August 7, 2019	NMPA approved Japan-originated ALS treatment Edaravone in China
August 8, 2019	Regulatory Approval for Marketing Cariprazine (MP-214) for the Treatment of Schizophrenia in Part of the ASEAN (Singapore and Thailand)
August 8, 2019 Phase I Clinical Trial study protocol for MT-3921 in Patients with Spinal Cord Injury submitte	
August 28, 2019 The Start of a Phase 3 Clinical Trial of Continuous Subcutaneous Liquid levodopa/carbidopa Admini (ND0612) for patients with fluctuating Parkinson's disease	
September 4, 2019	Notice regarding the launch of Collategene intramuscular injection 4 mg, an HGF gene therapy product
September 17, 2019	Mitsubishi Tanabe Pharma and Daiichi Sankyo announce alliance for ALS treatment agent edaravone in Brazil
September 20, 2019	Notice Regarding the Conclusion of Joint Sales Promotion Contract Concerning Alesion Ophthalmic Solution 0.05% and Alesion LX Ophthalmic Solution 0.1% between Mitsubishi Tanabe Pharma and Santen Pharmaceutical Co., Ltd.
September 24, 2019	NMPA accepts our filing for TENELIA, a treatment agent for type 2 diabetes mellitus in China
September 30, 2019	Mitsubishi Tanabe Pharma to participate in new screening program through the Global Health Innovative Technology Fund \sim Targeting treatments for infectious diseases that burden the developing countries \sim
October 2, 2019	Notification of acceptance of the New Drug Submission for Scientific Review of VLP Seasonal Influenza Vaccines (MT-2271) by Health Canada
October 9, 2019	Mitsubishi Tanabe Pharma enters into a licensing agreement with Viela Bio for inebilizumab, a treatment agent for neuromyelitis optica spectrum disorder, in Japan and other Asian regions
October 25, 2019	Vadadustat (MT-6548) Japan Phase 3 results for treatment of renal anemia to be presented at ASN Kidney Week 2019

