Financial Information for the 3rd Quarter of Fiscal Year Ending March 31, 2021

As of February 3, 2021 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 including products under development, but is not intended for advertising or medical advice.

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Summary of Financial Results for the 3rd Quarter of FY2020 Ending March 31, 2021 and Forecasts for FY2020

<Regarding GILENYA Royalty>

As Mitsubishi Tanabe Pharma Corporation (hereinafter, "MTPC") announced on April 24, 2019 in the "Revision to Consolidated Financial Forecasts for Fiscal Year Ending March 31, 2019", MTPC is currently in the arbitration proceedings with Novartis Pharma AG (hereinafter "Novartis"), and among the GILENYA Royalty amounts that MTPC is going to receive from Novartis, MTPC decided not to recognize some of those amounts, which correspond to the clauses in the 1997 License Agreement of which Novartis has protested the validity, as our revenue because such payments do not satisfy one of the requirements under IFRS15, i.e., "Revenue under contract with customers". During the period of the arbitration proceedings, MTPC will continue the same accounting practice as MTPC did in fiscal year 2018. For fiscal year 2020, the forecast is prepared on the assumption that the arbitration procedure to continue. 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. As for the amounts among 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.

1. Summary of Financial Results for the 3rd Quarter of FY2020

				(Amounts less than + 100 minion are rounded)
				[Billion yen]
Revenue	290.2	Y-on-Y	(7.2)	(2.4 %)
Domestic	241.8	Y-on-Y	(5.5)	(2.2 %)
Overseas	48.4	Y-on-Y	(1.7)	(3.5 %)

Domestic ethical drugs sales decreased by 2.0% to ¥235.0 billion, due to NHI price revision in April 2020 and increasing usage of generic drugs, despite increase of SIMPONI for Rheumatoid arthritis (RA) etc. treatment, CANAGLU and CANALIA for type 2 diabetes mellitus treatment and RUPAFIN for allergy treatment, as well as contribution of STELARA additionally approved for ulcerative colitis treatment in March 2020. Royalty revenue, etc. decreased by 8.7% to ¥12.4 billion due to the decline in royalty revenue from GILENYA for multiple sclerosis treatment licensed to Novartis etc.

				[Billion yen]
Core Operating Profit ^{*1}	24.7	Y-on-Y	0.6	2.3 %

Core operating profit increased due to SG&A and R&D expenses contained by voluntary restraint in activities under COVID-19 spread, despite decrease in sales revenue.

				[Billion yen]
Operating Profit ^{*2}	(54.7)	Y-on-Y	(79.7)	-

As for non-recurring items;

Profitability of NeuroDerm's projects for Parkinson's Disease is expected to decline due to delayed clinical study and the competitors' development status. As a result of reviewing the business plan based on the results of recent market research, we recorded an impairment loss of 84.5 billion Japanese yen for intangible assets related to above projects. Gain from fixed assets in transfer of Toda office, etc. booked ¥8.1 billion as well.

				[Billion yen]
Profit before tax for the period ^{*2}	(54.5)	Y-on-Y	(79.1)	-
Net Income Attributable to owners of the Company ^{*2}	(45.3)	Y-on-Y	(63.5)	-

2. Summary of Forecasts for	or FY2020			[Billion yen]
Revenues	373.0	Y-on-Y	(6.8)	(1.8 %)
Core Operating Profit	17.0	Y-on-Y	(2.1)	(10.8 %)
Operating Profit *2	(62.5)	Y-on-Y	(56.4)	-
Profit before Tax ^{*2}	(62.0)	Y-on-Y	(55.5)	-
Net Income Attributable to owners of the Company ^{*2}	(52.5)	Y-on-Y	(52.6)	-

Forecasts announced Nov. 4, 2020 remain unchanged.

*1 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. The Company assumes gain or loss associated with a business transfer, restructuring loss, impairment losses on intangible assets associated with products and others as non-recurring items.

*2 Brackets indicate expense and loss

(Amounts less than ¥100 million are rounded)

[Billion yen]

1. Profit and Loss

(1) Profit and Loss

	Q3	Compari	ison to previo	us year	Comparison t	o Forecasts	Notes
	FY2020	Q3 FY2019	Increase (decrease)	Change %	Forecasts*1	Progress %	[Y-on-Y comparison]
Revenue	290.2	297.4	(7.2)	(2.4)	373.0	77.8	Refer to "(2) Sales Revenue of Main Products"
Domestic	241.8	247.3	(5.5)	(2.2)	312.2	77.4	
Overseas	48.4	50.2	(1.7)	(3.5)	60.8	79.6	
Overseas sales ratio	16.7%	16.9%			16.3%		
Cost of sales	147.2	143.1	4.2	2.9	187.5	78.5	Deteriorated by NHI price revision, etc.
Sales cost ratio	50.7%	48.1%			50.3%		
Gross profit	143.0	154.3	(11.4)	(7.4)	185.5	77.1	
SG&A expenses, etc.	118.2	130.2	(12.0)	(9.2)	168.5	70.2	
R&D expenses	50.3	57.6	(7.3)	(12.7)	72.5	69.3	
Core operating profit ^{*2}	24.7	24.2	0.6	2.3	17.0	145.5	
Non-recurring items ^{*3}	(79.5)	0.8	(80.3)	-	(79.5)	-	Gain from sales of Toda Office:7.5, Impairment loss from NeuroDerm projects:84.5, etc.
Operating profit*3	(54.7)	25.0	(79.7)	-	(62.5)	-	
Financial income and loss*3	0.3	(0.4)	0.6	-			
Profit before tax for the period ^{*3}	(54.5)	24.6	(79.1)	-	(62.0)	-	
Income taxes	(6.9)	9.3	(16.2)	-			
Net profit for the period*3	(47.6)	15.3	(62.9)	-	(55.0)	-	
Net profit attributable to owners of the Company ^{*3}	(45.3)	18.2	(63.5)	-	(52.5)	-	

			[Yen]
Exchange rate	Q3 FY2020 average	Q3 FY2019 average	FY2020 planned
USD	105.54	108.89	108.00

Effect of fluctuations in exchange rate for FY2020 Q3: Revenue decreased by ¥0.9 billion and core operating profit increased by ¥1.8 billion.

(2) Sales Revenue of Main Produ	cts				[Billion yen]
	Q3	Compari	ison to previo	us year	Comparison to Forecasts	
	FY2020	Q3 FY2019	Increase (decrease)	Change %	Forecasts*1	Progress %
Domestic ethical drugs	235.0	239.6	(4.7)	(2.0)	302.3	77.7
Priority products	139.2	137.9	1.3	1.0	183.0	76.1
Remicade	35.5	42.5	(7.0)	(16.5)	45.0	78.9
Simponi	32.7	31.7	1.0	3.3	42.7	76.7
Stelara	23.0	20.4	2.6	13.0	31.9	72.1
Tenelia	11.8	12.1	(0.3)	(2.1)	14.9	79.4
Canaglu	8.0	6.6	1.4	21.3	9.8	81.5
Canalia	7.5	5.5	2.0	36.0	9.3	80.6
Vafseo (launched in Aug.)	0.3	-	0.3	-	0.5	65.6
Lexapro	11.9	11.6	0.2	2.1	14.8	80.1
Rupafin	5.2	4.2	1.1	25.3	10.0	52.1
Imusera	3.2	3.4	(0.1)	(3.2)	4.1	79.2
Vaccines*4	36.0	32.9	3.0	9.3	41.6	86.4
Influenza vaccine	13.8	12.4	1.4	11.1	13.2	104.7
Tetrabik	8.2	7.1	1.1	15.3	11.1	73.7
Mearubik	5.1	4.8	0.3	6.1	6.4	79.1
JEBIK V	4.3	4.2	0.2	4.0	5.3	81.9
Varicella vaccine	3.8	3.8	(0.0)	(0.1)	4.8	78.9
Long-listed drugs, etc.*4	59.8	68.9	(9.1)	(13.2)	77.7	77.0
Overseas ethical drugs	37.0	37.5	(0.5)	(1.3)	47.0	78.8
Radicava	15.9	15.9 17.4 (1.4) (8.3)			20.1	79.3
Royalty revenue, etc.	12.4	12.4 13.6 (1.2) (8.7)		(8.7)	15.2	81.8
Royalty from INVOKANA	7.4	6.5	0.9	14.0	Undisclosed	-
Royalty from GILENYA*5	3.1	4.6	(1.5)	(32.7)	Undisclosed	-

*1: Forecasts announced on Nov. 4, 2020.

*2: COVID-19 impact:¥6.8 billion. Decreased expenses by shrinkage in sales promotion and delay in R&D expenses incurrence overtake sales decrease by consultation restraint.

*3: Brackets indicate expense and loss

*4: Corrected vaccines forecasts from 40.8 to 41.6 and Long-listed drugs, etc. forecasts from 78.5 to 77.7 due to mistaken announcement on Nov.4

*5: MTPC is currently in the arbitration proceedings with Novartis, and among the GILENYA Royalty amounts that MTPC is going to receive from Novartis, MTPC decided not to recognize some of those amounts as our revenue for FY2018 because such payments do not satisfy one of the requirements under IFRS15. 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.

2. Financial Statement

(1) Balance Sheet

[Billion yen]

	End of Q3 FY2020	End of FY2019	Increase (decrease)	Notes
Assets	1,022.0	1,046.3	(24.3)	
Non-current assets	363.3	452.8	(89.5)	
Property, plant and equipment	79.5	86.1	(6.6)	Obtain 15.3, depreciation(8.4), etc.
Goodwill	89.5	89.7	(0.2)	
Intangible assets	91.9	181.3	(89.4)	Impairment loss from NeuroDerm's projects(84.5), etc.
Current assets	658.7	593.5	65.2	
Inventories	74.9	80.3	(5.4)	
Trade and other receivables	133.4	108.6	24.8	
Other financial assets	297.0	300.3	(3.3)	
Cash and cash equivalents	124.3	83.1	41.2	Refer to "(2) Cash Flow Statement"
Liabilities	212.2	188.4	23.9	
Non-current liabilities	96.3	90.3	6.0	
Other non-current liabilities	67.6	40.9	26.7	
Current liabilities	115.9	98.0	17.8	
Trade and other payables	32.5	32.1	0.4	
Equity	809.8	857.9	(48.1)	
Share capital	50.0	50.0	-	
Capital surplus	448.0	448.0	(0.1)	
Retained earnings	313.8	358.4	(44.7)	Net loss for the period 45.3, etc.

(2) Cash Flow Statement

(2) Cash Flow Statement			[Billion yen]
	Q3 FY2020	Q3 FY2019	Increase (decrease)
Cash and cash equivalents at beginning of year	83.1	111.9	(28.8)
Cash flows from operating activities	46.8	32.3	14.5
Profit before tax for the period*	(54.5)	24.6	(79.1)
Depreciation and amortization	11.4	11.3	0.1
Impairment loss	84.5	0.1	84.5
Loss on sales of Property, Plant and Equipment	(8.1)	-	(8.1)
Trade receivable and payable	(24.3)	(5.4)	(18.9)
Other	37.8	1.7	36.0
Cash flows from investing activities	1.4	(10.4)	11.8
Purchase (proceeds from sales) of property, plant and equipment	(0.4)	(8.3)	7.8
Purchase (Proceeds from sales) of investments	64.3	4.5	59.8
Increase in deposits	(65.1)	(0.0)	(65.1)
Other	2.6	(6.6)	9.2
Cash flows from financing activities	(6.8)	(35.7)	28.9
Effect of exchange rate changes on cash and cash equivalents	(0.2)	(0.5)	0.2
Net increase(decrease) in cash and cash equivalents	41.2	(14.3)	55.5
Increase (decrease) by transfer to assets held for sales	-	0.1	(0.1)
Cash and cash equivalents at the end of period	124.3	97.7	26.6

*Brackets indicate loss

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

Investment in Development of Information Systems						
	[Billion yen]					
	Q3	Q3	Increase			
Occurring basis	FY2020	FY2019	(decrease)			
Investment in property, plant and equipment	15.3	9.0	6.3			
Investment in information systems	1.0	0.8	0.1			

(4) Depreciation and Amortization Costs

(4) Depreciation and Amortization Costs [Billion yer							
	Q3 FY2020	Q3 FY2019	Increase (decrease)				
Property, plant and equipment	8.4	5.3	3.1				
Intangible assets	0.9	1.0	(0.1)				
Intangible assets with products	2.1	1.9	0.2				

(Amounts less than ¥ 100 million are rounded)

[Billion yen]

(1) Consolidated Forecasts of Profit and Loss

	FY2020	Compa	rison to previou	ıs year	Notes
	forecasts	FY2019	Increase	Change	[Y-on-Y Comparison]
		actual	(decrease)	%	
Revenue	373.0	379.8	(6.8)	(1.8)	Refer to "(2) Sales Revenue Forecasts for Main
Domestic	312.2	314.0	(1.8)	(0.6)	Products"
Overseas	60.8	65.8	(5.0)	(7.7)	
Overseas sales ratio	16.3%	17.3%			
Cost of sales	187.5	181.0	6.5	3.6	Deteriorated by NHI price revision, etc.
Sales cost ratio	50.3%	47.7%			
Gross profit	185.5	198.8	(13.3)	(6.7)	
SG&A expenses, etc.	168.5	179.7	(11.2)		Shrinkage in sales promotion and delay in R&D expenses incurrence
R&D expenses	72.5	79.4	(6.9)	(8.7)	
Core operating profit	17.0	19.1	(2.1)	(10.8)	
Non-recurring items*1	(79.5)	(25.1)	(54.4)	-	Impairment loss from NeuroDerm projects: 84.5 etc.
Operating profit ^{*1}	(62.5)	(6.1)	(56.4)	-	
Profit before tax for the period*1	(62.0)	(6.5)	(55.5)	-	
Net profit for the period*1	(55.0)	(9.4)	(45.6)	-	
Net profit attributable to owners of the Company*1	(52.5)	0.1	(52.6)	-	

Exchange rate		[Yen]
	FY2020	FY2019
	planned	average
USD	108.00	108.95

(2) Sales Revenue Forecasts for Main Products								
		FY2020	Compa	rison to previou	is year			
		forecasts	FY2019	Increase	Change %			
			actual	(decrease)				
	estic ethical drugs	302.3	304.4	(2.1)	(0.7)			
l l	Priority products	183.0	177.1	5.9	3.3			
	Remicade	45.0	53.4	(8.4)	(15.7)			
	Simponi	42.7	41.0	1.7	4.3			
	Stelara	31.9	26.0	5.9	22.6			
	Tenelia	14.9	15.2	(0.3)	(2.1)			
	Canaglu	9.8	8.8	1.0	10.8			
	Canalia	9.3	6.7	2.6	38.1			
	Vafseo (launched in Aug.)	0.5	-	0.5	-			
	Lexapro	14.8	15.0	(0.2)	(1.0)			
	Rupafin	10.0	6.8	3.2	47.7			
	Imusera	4.1	4.2	(0.1)	(3.4)			
	Vaccines*2	41.6	39.0	2.6	6.7			
	Influenza vaccine	13.2	12.6	0.6	4.5			
	Tetrabik	11.1	9.5	1.6	17.0			
	Mearubik	6.4	6.0	0.4	7.5			
	JEBIK V	5.3	5.2	0.1	2.8			
	Varicella vaccine	4.8	4.9	(0.1)	(2.5)			
	Long-listed drugs, etc.*2	77.7	88.3	(10.6)	(12.0)			
Over	seas ethical drugs	47.0	49.7	(2.7)	(5.5)			
Radicava		20.1	23.1	(3.0)	(13.2)			
Roya	lty revenue, etc.	15.2	17.4	(2.2)	(12.8)			
F	Royalty from INVOKANA	Undisclosed	8.5	-	-			
F	Royalty from GILENYA*3	Undisclosed	5.7	-	-			
*1.0	*1: Brackets indicate expense and loss							

*1: Brackets indicate expense and loss

*2: Corrected vaccines forecasts from 40.8 to 41.6 and Long-listed drugs, etc. forecasts from 78.5 to 77.7 due to mistaken announcement on Nov.4

*3: MTPC is currently in the arbitration proceedings with Novartis, and among the GILENYA Royalty amounts that MTPC is going to receive from Novartis, MTPC decided not to recognize some of those amounts as our revenue for FY2018 because such payments do not satisfy one of the requirements under IFRS15. 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.

Five-Year Financial Data

(Amounts less than ¥100 million are rounded)

[Billion yen]

[Billion yen]

(1) Profit and Loss

4

(1) Profit and Loss								
	FY2016	FY2017	FY2018	FY2019	Q3 FY2020	FY2020 forecasts		
Revenues	424.0	433.9	424.8	379.8	290.2	373.0		
Cost of sales	164.4	169.8	180.6	181.0	147.2	187.5		
Gross profit	259.6	264.1	244.1	198.8	143.0	185.5		
SG&A expenses, etc.	165.1	185.6	188.3	179.7	118.2	168.5		
R&D expenses	64.8	79.1	86.5	79.4	50.3	72.5		
Core operating profit	94.5	78.5	55.8	19.1	24.7	17.0		
Operating profit	94.1	77.3	50.3	(6.1)	(54.7)	(62.5)		
Profit before tax	96.1	78.8	50.4	(6.5)	(54.5)	(62.0)		
Net profit for the period	68.9	54.0	32.2	(9.4)	(47.6)	(55.0)		
Net profit attributable to owners of the Company	71.3	58.0	37.4	0.1	(45.3)	(52.5)		

(2) Balance Sheet

End of Q3 End of FY2016 End of FY2017 End of FY2018 End of FY2019 FY2020 984.5 1,048.4 1,056.3 Assets 1,046.3 1,022.0 Non-current assets 300.8 462.9 467.9 452.8 363.3 Current assets 683.8 585.5 588.4 593.5 658.7 Liabilities 113.1 153.6 146.0 212.2 188.4 Non-current liabilities 24.7 54.3 90.3 96.3 55.4 Current liabilities 98.2 91.7 98.0 115.9 88.4 871.4 857.9 809.8 Equity 894.8 910.3

(3) Other Financial Data

FY2020 Q3 FY2016 FY2017 FY2018 FY2019 FY2020 forecasts Cash flows from operating activities 59.8 66.9 41.5 49.4 46.8 Cash flows from investing activities (10.6)(19.2)(31.2)(39.2) 1.4 Cash flows from financing activities (24.4)(32.5) (25.9) (37.9) (6.8) Investments in property, 14.5 6.2 8.6 15.5 16.2 18.2 plant and equipment Depreciation and Amortization Costs 10.5 11.5 11.5 10.9 11.4 15.7 Property, plant and equipment 7.3 7.6 7.1 7.0 8.4 11.4 Intangible assets including intangible assets 3.1 4.0 4.4 4.0 3.0 4.3 with products Ratio of equity attributable to owners of 87.4 84.2 85.0 81.4 78.5 the Company to total assets [%] 4.2 ROE [%] 8.5 6.6 0.0 Basic earnings per share [¥] 127.03 103.35 66.64 0.26 Equity attributable to owners of the 1,429.53 1,533.91 1,574.26 1,600.64 1,519.22 Company per share [¥]

(4) Number of Employees

	End of FY2016	End of FY2017	End of FY2018	End of FY2019	End of Q3 FY2020	Forecasts for end of FY2020
Consolidated	7,280	7,187	7,228	6,987	6,805	7,000
Non-consolidated	4,239	4,222	4,111	3,764	3,449	3,450

5 Quarterly Trend

(Amounts less than ¥ 100 million are rounded)

(1) Profit and Loss

(1) Profit and Loss [Billion yen									Billion yen]
	FY2019						FY2	020	
	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.	Full-year Forecasts
Davianua	98.1	90.0	109.3	82.4	379.8	91.8	95.5	102.9	373.0
Revenue	25.8%	23.7%	28.8%	21.7%	100.0%	24.6%	25.6%	27.6%	100.0%
Demostic	80.8	73.9	92.6	66.7	314.0	75.7	79.3	86.7	312.2
Domestic	25.7%	23.5%	29.5%	21.3%	100.0%	24.3%	25.4%	27.8%	100.0%
Overseas	17.4	16.1	16.7	15.7	65.8	16.1	16.2	16.1	60.8
overseus	26.4%	24.5%	25.3%	23.8%	100.0%	26.4%	26.7%	26.5%	100.0%
Cost of sales	44.8	43.7	54.6	38.0	181.0	45.6	49.2	52.4	187.5
Sales cost ratio	45.6%	48.6%	49.9%	46.1%	47.7%	49.7%	51.6%	50.9%	50.3%
	53.3	46.3	54.7	44.5	198.8	46.2	46.3	50.5	185.5
Gross profit	26.8%	23.3%	27.5%	22.4%	100.0%	24.9%	24.9%	27.2%	100.0%
SG&A expenses,	43.6	44.3	42.3	49.6	179.7	36.6	41.3	40.3	168.5
etc.	24.3%	24.7%	23.5%	27.6%	100.0%	21.7%	24.5%	23.9%	100.0%
	19.9	19.9	17.8	21.9	79.4	15.3	18.6	16.4	72.5
R&D expenses	25.1%	25.0%	22.4%	27.6%	100.0%	21.1%	25.7%	22.6%	100.0%
Cours anounting ausfit*	9.8	1.9	12.5	(5.1)	19.1	9.6	5.0	10.2	17.0
Core operating profit*	51.2%	10.2%	65.5%	(26.9%)	100.0%	56.3%	29.3%	59.9%	100.0%
*	9.6	2.9	12.4	(31.1)	(6.1)	17.7	(79.6)	7.2	(62.5)
Operating profit [*]	-	-	-	-	-	-	-	-	-
*	9.2	2.9	12.5	(31.1)	(6.5)	17.8	(79.4)	7.1	(62.0)
Profit before tax [*]	-	-	-	-	-	-	-	-	-
Net profit attributable to	6.9	1.4	9.9	(18.1)	0.1	11.5	(62.4)	5.6	(52.5)
owners of the Company *	-	-	-	-	-	-	-	-	-

Note: The progress rates show in the lower of each cell, except for "cost of sales".

*Brackets indicate expense and loss

(2) Sales Revenue of Main Products

		FY2019				FY2020				
		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.	Full-year Forecasts
		78.2	71.0	90.5	64.7	304.4	73.3	77.0	84.7	302
omest	tic ethical drugs	25.7%	23.3%	29.7%	21.3%	100.0%	24.2%	25.5%	28.0%	100.0
	Priority products	46.5	42.2	49.1	39.2	177.1	45.3	44.5	49.4	18
		26.3%	23.8%	27.7%	22.2%	100.0%	24.8%	24.3%	27.0%	100.
	Remicade	14.4 27.1%	13.2 24.7%	14.9 27.9%	10.9 20.4%	53.4 100.0%	11.9 26.4%	11.5 25.6%	12.1 26.8%	4 100.
		10.5	9.9	11.2	9.3	41.0	10.7	10.5	11.5	4
	Simponi	25.7%	24.3%	27.5%	22.6%	100.0%	25.0%	24.7%	27.0%	100
	Stelara	6.2	6.4	7.8	5.7	26.0	7.0	7.0	9.1	1
	Steidid	23.8%	24.6%	29.9%	21.8%	100.0%	21.8%	21.9%	28.4%	100
	Tenelia	4.7	3.4	4.0	3.1	15.2	4.1	3.9	3.9	
		30.9%	22.3%	26.3%	20.5%	100.0%	27.6%	25.9%	26.0%	100
	Canaglu	2.2	1.9	2.5	2.3	8.8	2.5	2.5	3.0	
		24.4%	22.0%	28.1%	25.5%	100.0%	25.9%	25.5%	30.2%	100
	Canalia	2.2	1.6	1.8	1.2	6.7	2.5	2.5	2.5	
		32.7%	23.0%	26.1%	18.2%	100.0%	27.3%	26.6%	26.6%	100
	Vafseo	-	-	-	-	-	-	0.3	0.0	
	(launched in Aug.)	-	-	-	-	-	-	60.6%	5.0%	100
	Lexapro	3.9	3.6	4.2	3.3	15.0	3.9	3.7	4.2	
		26.1%	23.8%	27.9%	22.3% 2.6	100.0%	26.2% 1.7	25.2%	28.7% 2.0	100
	Rupafin	1.3	1.2	1.7		6.8		1.6		
		18.5% 1.1	18.4%	24.6% 1.2	38.5% 0.9	100.0% 4.2	16.6% 1.1	15.6% 1.0	20.0%	100
	Imusera	27.0%	1.0 24.4%	27.7%	0.9 20.9%	4.2 100.0%	1.1 26.8%	25.3%	27.1%	100
		7.3	8.4	17.2	6.1	39.0	7.5	13.6	14.8	
	Vaccines*1	18.7%	21.6%	44.1%	15.6%	100.0%	18.1%	32.7%	35.6%	100
	Influenza vaccine	(0.0)	1.8	10.6	0.2	12.6	(0.0)	6.4	7.5	
	Innuenza vaccine	(0.1%)	14.3%	84.3%	1.6%	100.0%	(0.3%)	48.2%	56.8%	100
	Tetrabik	2.4	2.2	2.5	2.4	9.5	2.7	2.5	3.0	
		25.0%	23.2%	26.5%	25.3%	100.0%	24.1%	22.3%	27.3%	100
	Mearubik	1.9	1.6	1.3	1.2	6.0	1.9	1.8	1.4	
		31.9%	27.1%	21.2%	19.8%	100.0%	29.6%	28.3%	21.3%	100
	JEBIK V	1.5	1.4	1.3	1.0	5.2	1.4	1.5	1.4	
		29.3%	26.6%	25.1%	19.0%	100.0%	27.3%	27.8%	26.8%	100
	Varicella vaccine	1.3	1.2	1.3	1.1	4.9	1.3	1.2	1.3	
		26.2%	24.7%	26.1%	23.1%	100.0%	26.3% 20.4	25.6% 18.9	26.9%	100
1	Long-listed drugs, etc.*1	24.3	20.4	24.2	19.4	88.3			20.5	
		27.6%	23.1%	27.4%	22.0%	100.0%	26.3%	24.3%	26.4%	100
ersea	as ethical drugs	12.6	12.3	12.6	12.2	49.7	12.6	12.5	11.9	
	-	25.3%	24.7%	25.4%	24.6%	100.0%	26.9%	26.6%	25.3%	100
- 1	Radicava	6.1 26.5%	5.5 23.8%	5.7 24.8%	5.8 24.9%	23.1 100.0%	5.6 27.8%	5.5 27.4%	4.9 24.1%	100
_		5.1	4.2	4.4	3.8	100.0 %	3.8	4.1	4.6	100
yalty	revenue, etc.	29.0%	4.2 23.9%	4.4 25.2%	5.8 21.9%	17.4	24.7%	26.8%	4.0 30.3%	100
	Royalty from	29.070	23.970	2.4	21.970	8.5	24.7%	20.0 %	2.8	Undiscle
	INVOKANA	2.1	2.0	2.4 28.3%	2.0	8.5 100.0%	2.0	2.5	2.0	0.101001
	Royalty from	24.2%	23.9%	28.3%	23.7%	5.7	- 1.1	- 0.9	- 1.2	Undisclo
	GILENYA*2	29.3%	27.7%	23.8%	1.1	5.7 100.0%		0.5	1.2	

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

*1: Corrected vaccines forecasts from 40.8 to 41.6 and Long-listed drugs, etc. forecasts from 78.5 to 77.7 due to mistaken announcement on Nov.4

*2: MTPC is currently in the arbitration proceedings with Novartis, and among the GILENYA Royalty amounts that MTPC is going to receive from Novartis, MTPC decided not to recognize some of those amounts as our revenue for FY2018 because such payments do not satisfy one of the requirements under IFRS15. 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.

6 State of New Product Development (As of January 25, 2021)

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-7117 (Dersimelagon)	Selective melanocortin 1 receptor agonist (Erythropoietic protoporphyria, X-Linked protoporphyria)	Global Phase 3	In-house
MT-0551 (Inebilizumab)	Humanized anti-CD19 monoclonal antibody (IgG4-related disease)	Japan Phase 3	Licensed from Viela Bio (US) and co-developed (Global study ongoing)
MT-1303	S1P receptor functional antagonist (Multiple sclerosis)	Europe Phase 2	In-house
(Amiselimod)	(Crohn's disease)	Japan Phase 2	in nouse
MT-2990	Fully human anti-interleukin-33 (IL-33) monoclonal antibody (Endometriosis)	Global Phase 2	In-house

ii. Diabetes and kidney

Development code Product name (Generic name)	Category (Indications)	Region Stage	Origin/licensee
TA-7284 Canaglu/INVOKANA (Canagliflozin)	SGLT2 inhibitor (Diabetic nephropathy)	Asia Filed Japan Phase 3	In-house
MP-513 Tenelia (Teneligliptin)	DPP-4 inhibitor (Type 2 diabetes mellitus)	Asia Filed China Filed (Sep. 2019) Europe Phase 2	In-house
MT-3995 (Apararenone)	Selective mineralocorticoid receptor antagonist (Diabetic nephropathy) (Non-alcoholic steatohepatitis: NASH)	Europe Phase 2 Japan Phase 2 Japan Phase 2	In-house

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-0551	Humanized anti-CD19 monoclonal antibody (Neuromyelitis optica spectrum disorder: NMOSD)	Japan Filed (June 2020) Asia Filed	Licensed from Viela Bio (US)
(Inebilizumab)	(Myasthenia gravis)	Japan Phase 3	Licensed from Viela Bio (US) and co-developed (Global study ongoing)
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 Asia Filed	Licensed from Neurocrine Biosciences (US)
ND0612 (Levodopa/Carbidopa)	Continuous SC pump (Parkinson's disease)	Global Phase 3	In-house
MT-1186 (Edaravone)	Free radical scavenger (Amyotrophic lateral sclerosis: ALS / Oral suspension)	Global Phase 3	In-house
MT-8554 (Elismetrep)	TRPM8 antagonist Europe (Painful diabetic peripheral neuropathy) Phase 2 p) Global		In-house
MT-3921	(Vasomotor symptoms associated with menopause) Anti-RGMa antibody (Spinal cord injury)	Phase 2 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) Europe Phase 3 Europe	Medicago product (Canada)
MT-2766	(Prophylaxis of seasonal influenza/elderly) Plant-based VLP vaccine (Prophylaxis of COVID-19)	Phase 3 Global Phase 2	Medicago product (Canada)
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-based VLP vaccine (Prophylaxis of rotavirus gastroenteritis)	Phase 1	Medicago product (Canada)
MT-2654	Adjuvanted plant-based VLP vaccine (Prophylaxis of seasonal influenza/elderly)	Phase 1	Medicago product (Canada)

Asia: Excluding Japan and China

v. Others

Development code Product name (Generic name)	Category (Indications)	Region Stage	Origin/licensee
MT-4580 Orkedia (Evocalcet)	Ca sensing receptor agonist (Secondary Hyperparathyroidism)	China, Asia Phase 3	Licensed to Kyowa Kirin (Japan)
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

Changes Since Previous Announcement

Development code Product name (Generic name)	Category (Indications)	Previous Announcement	As of Jan. 25, 2021	Origin/licensee
MT-0551 (Inebilizumab)	Humanized anti-CD19 monoclonal antibody (IgG4-related disease)	None	Japan Phase 3	Licensed from Viela Bio (US) and co-developed (Global study ongoing)
	Humanized anti-CD19 monoclonal antibody (Myasthenia gravis)	None	Japan Phase 3	
MT-2766	Plant-based VLP vaccine (Prophylaxis of COVID-19)	Phase 1	Global Phase 2	Medicago product (Canada)
MT-2654	Adjuvanted plant-based VLP vaccine (Prophylaxis of seasonal influenza/ elderly)	None	Phase 1	Medicago product (Canada)

Asia: Excluding Japan and China