Disclaimer: The following is meant to be an accurate translation from the original Financial Report of Santen Pharmaceutical Co., Ltd., written in Japanese, and is prepared for the information disclosure to the Tokyo Stock Exchange. However, in the case of any discrepancy between the English translation and the Japanese original, the latter shall prevail.

February 5, 2019

Santen Pharmaceutical Co., Ltd. Code : 4536 URL http://www.santen.com Third Quarter Financial Results for the Fiscal Year Ending March 31, 2019 [IFRS] (Consolidated) Akira Kurokawa, Chairman and Chief Executive Officer Contact : Christopher Hohman, General Manager, Corporate Communications Group Tel : +81-6-4802-9360 E-mail : ir@santen.co.jp

(JPY millions)

1. Consolidated performance for the nine months ended December 31, 2018

(1) Operating results

(Core basis *1)

	Nine months ended December 31, 2017	Nine months ended December 31, 2018	% change
Revenue	168,592	173,210	+2.7%
Core operating profit	35,042	35,082	+0.1%
Core net profit for the period	26,231	25,655	(2.2%)
Core net profit for the period attributable to owners of the company	26,216	25,663	(2.1%)
Basic core earnings per share (yen)	64.52	63.06	
Diluted core earnings per share (yen)	64.28	62.87	

(IFRS)

	Nine months ended December 31, 2017	Nine months ended December 31, 2018	% change
Revenue	168,592	173,210	+2.7%
Operating profit	30,087	33,657	+11.9%
Profit before tax	29,539	32,405	+9.7%
Net profit for the period	27,348	23,367	(14.6%)
Net profit for the period attributable to owners of the company	27,332	23,376	(14.5%)
Total comprehensive income for the period	34,602	21,732	(37.2%)
Basic earnings per share (yen)	67.27	57.44	
Diluted earnings per share (yen)	67.02	57.26	

(2) Financial position

	March 31, 2018	December 31, 2018
Total assets	388,463	388,284
Total equity	287,557	298,769
Total equity attributable to owners of the company	285,823	297,122
Total equity attributable to owners of the company ratio	73.6%	76.5%
Equity per share attributable to owners of the company (yen)	702.54	729.98

2. Dividends

	Year to March 2018	Year to March 2019	(Forecasts) Year to March 2019
First quarter dividends per share (yen)	_	-	-
Second quarter dividends per share (yen)	13.00	13.00	-
Third quarter dividends per share (yen)	-	-	-
Year-end dividends per share (yen)	13.00	-	13.00
Annual dividends per share (yen)	26.00		26.00

3. Consolidated forecasts of results for the year ending March 31, 2019

(Core basis *1)

	Year to March 2019	% change
Revenue	237,000	+5.4%
Core operating profit	48,000	+5.8%
Core net profit for the year	35,300	+5.5%
Core earnings per share (yen)	87.26	

(IFRS)

	Year to March 2019	% change
Revenue	237,000	+5.4%
Operating profit	40,700	+5.2%
Profit before tax	41,300	+5.2%
Net profit for the year	30,400	(13.8%)
Basic earnings per share (yen)	75.21	

*Others

(1) Changes in significant subsidiaries during the term (changes in designated subsidiaries resulting in adjustment to the scope of consolidation): No

(2) Changes in accounting policies and accounting estimates

- [i] Changes in the accounting policies required by IFRS: Yes
- [ii] Other changes: No
- [iii] Changes in accounting estimates: No

(3) Number of shares outstanding (common stock):

- [i] Number of shares outstanding at the end of period (including treasury shares) December 31, 2018 : 407,102,254 March 31, 2018 : 406,847,515
- [ii] Number of treasury shares at the end of periodDecember 31, 2018 :7,821March 31, 2018 :7,411

[iii] Average number of outstanding shares (during the fiscal year ended December 31)

Third quarter ended December 31, 2018: 406,966,797

Third quarter ended December 31, 2017: 406,304,134

(Information regarding the implementation of audit procedures)

This financial report is exempt from audit.

(Information regarding presentation currency)

All financial information presented in Japanese yen has been rounded to the nearest million, except when otherwise indicated.

(Caution)

Forecasts in this report are based on the currently available information. Actual results may differ materially depending on a number of factors including adverse economic conditions, delays in new products launch, currency exchange rate, legislative and regulatory developments.

*1(Core basis)

Core results are non-IFRS measures that remove non-core items including amortization on intangible assets associated with products, other income and expenses, finance income and expenses, and temporary expenses of selling, general and administrative expenses in order to provide greater transparency on underlying business performance.

Quarterly consolidated statements of income and comprehensive income

FRS basis		(JPY millions
	Nine months ended December 31, 2017	Nine months ended December 31, 2018
Revenue	168,592	173,210
Cost of sales	(66,150)	(69,814)
Gross profit	102,442	103,397
Selling, general and administrative expenses	(49,504)	(51,224)
Research and development expenses	(17,895)	(17,091)
Amortization on intangible assets associated with products	(4,996)	(5,233)
Other income	368	3,929
Other expenses	(327)	(121)
Operating profit	30,087	33,657
Finance income	809	863
Finance expenses	(1,357)	(2,115
Profit before tax	29,539	32,40
Income tax expenses	(2,192)	(9,037
Net profit for the period	27,348	23,36
Other comprehensive income		
Items that will not be reclassified subsequently to profit of loss		
Remeasurements of defined benefit plans	-	-
Net gain on financial assets measured at fair value through other comprehensive income	2,854	(2,733
Items that may be reclassified subsequently to profit or loss		
Foreign currency translation adjustments	4,400	1,098
Other comprehensive income	7,254	(1,635
Total comprehensive income	34,602	21,732
Profit attributable to		
Owners of the company	27,332	23,376
Non-controlling interests	15	(8
Net profit for the period	27,348	23,367
Total comprehensive income attributable to		
Owners of the company	34,487	21,819
Non-controlling interests	114	(87
Total comprehensive income	34,602	21,732
Earnings per share		
Basic earnings per share (yen)	67.27	57.44
Diluted earnings per share (yen)	67.02	57.26
Core basis		(JPY million
Core basis		(JPY mill

	Nine months ended December 31, 2017	Nine months ended December 31, 2018
Revenue	168,592	173,210
Core operating profit	35,042	35,082
Core net profit for the period	26,231	25,655
Basic core earnings per share (yen)	64.52	63.06
Diluted core earnings per share (yen)	64.28	62.87
Core profit attributable to		
Owners of the company	26,216	25,663
Non-controlling interests	15	(8)
Core net profit for the period	26,231	25,655

Quarterly consolidated statement of financial position

Assets		(JPY millions)
	March 31, 2018	December 31, 2018
Non-current assets		
Property, plant and equipment	29,706	31,035
Intangible assets	134,495	132,725
Financial assets	35,775	31,730
Deferred tax assets	2,264	2,258
Other non-current assets	2,855	1,685
Total non-current assets	205,095	199,433
Current assets		
Inventories	30,636	31,082
Trade and other receivables	78,654	78,720
Other financial assets	472	256
Other current assets	4,322	3,399
Cash and cash equivalents	69,283	75,394
Total current assets	183,367	188,851
Total assets	388,463	388,284

	March 31, 2018	December 31, 2018
Equity		
Equity attributable to owners of the company		
Share capital	8,032	8,186
Capital surplus	8,657	8,714
Treasury shares	(11)	(12
Retained earnings	249,225	262,404
Other components of equity	19,921	17,830
Total equity attributable to owners of the company	285,823	297,123
Non-controlling interests	1,734	1,647
Total equity	287,557	298,769
Liabilities		
Non-current liabilities		
Financial liabilities	21,244	23,243
Net defined benefit liabilities	1,804	2,24
Provisions	1,367	1,29
Deferred tax liabilities	12,909	12,31
Other non-current liabilities	1,380	1,73
Total non-current liabilities	38,704	40,830
Current liabilities		
Trade and other payables	29,743	27,982
Other financial liabilities	14,404	9,09
Income tax payable	7,656	3,60
Provisions	1,508	64
Other current liabilities	8,890	7,36
Total current liabilities	62,201	48,679
Total liabilities	100,905	89,51
Total equity and liabilities	388,463	388,284

Quarterly consolidated statement of changes in equity

Nine months ended December 31, 2017

(JPY millions)

				Other components of equity		
	Share capital	Capital surplus	Treasury shares	Retained earnings	Remeasurements of defined benefit plans	Net gain or loss on financial assets measured at fair value through other comprehensive income
Balance at April 1, 2016	7,792	8,417	(10)	223,283	_	9,470
Comprehensive income						
Net profit for the period				27,332		
Other comprehensive income						2,854
Total comprehensive income	-	_	-	27,332	-	2,854
Transactions with owners						
Issuance of new shares	119	119				
Acquisition of treasury shares			(1)			
Dividends				(10,563)		
Establishment of subsidiary with non-controlling interests						
Share-based payments						
Other				168		(168)
Total transactions with owners	119	119	(1)	(10,395)	-	(168)
Balance at December 31, 2016	7,911	8,536	(11)	240,220	_	12,156

	Other	components of e	quity	Total equity		
	Foreign currency translation adjustments	Subscription rights to shares	Total	attributable to owners of the company	Non-controlling interests	Total equity
Balance at April 1, 2016	5,332	825	15,628	255,110	819	255,929
Comprehensive income						
Net profit for the period			_	27,332	15	27,348
Other comprehensive income	4,301		7,155	7,155	99	7,254
Total comprehensive income	4,301	_	7,155	34,487	114	34,602
Transactions with owners						
Issuance of new shares		(39)	(39)	200		200
Acquisition of treasury shares			_	(1)		(1)
Dividends			_	(10,563)		(10,563)
Establishment of subsidiary with non-controlling interests			-	-	838	838
Share-based payments		218	218	218		218
Other			(168)	_		-
Total transactions with owners	_	179	11	(10,146)	838	(9,308)
Balance at December 31, 2016	9,633	1,004	22,794	279,451	1,772	281,222

Nine months ended December 31, 2018

(JPY millions)

					Other comp	onents of equity
	Share capital	Capital surplus	Treasury shares	Retained earnings	Remeasurements of defined benefit plans	Net gain or loss on financial assets measured at fair value through other comprehensive income
Balance at April 1, 2017	8,032	8,657	(11)	249,225	_	14,364
Comprehensive income						
Net profit for the period				23,376		
Other comprehensive income						(2,733)
Total comprehensive income	_	_	_	23,376	_	(2,733)
Transactions with owners						
Issuance of new shares	87	87				
Acquisition of treasury shares			(1)			
Dividends				(10,581)		
Share-based payments	67	(29)				
Other				385		(385)
Total transactions with owners	154	57	(1)	(10,196)	_	(385)
Balance at December 31, 2017	8,186	8,714	(12)	262,404	-	11,245

	Other	components of e	quity	Total equity		
	Foreign currency translation adjustments	Subscription rights to shares	Total	attributable to owners of the company	Non-controlling interests	Total equity
Balance at April 1, 2017	4,583	975	19,921	285,823	1,734	287,557
Comprehensive income						
Net profit for the period			-	23,376	(8)	23,367
Other comprehensive income	1,177		(1,557)	(1,557)	(79)	(1,635)
Total comprehensive income	1,177	_	(1,557)	21,819	(87)	21,732
Transactions with owners						
Issuance of new shares		(150)	(150)	23		23
Acquisition of treasury shares			-	(1)		(1)
Dividends			-	(10,581)		(10,581)
Share-based payments			-	38		38
Other			(385)	-		-
Total transactions with owners	_	(150)	(535)	(10,521)	_	(10,521)
Balance at December 31, 2017	5,759	825	17,830	297,122	1,647	298,769

Quarterly consolidated statements of cash flows

		(JPY millions)
	Nine months ended December 31, 2017	Nine months ended December 31, 2018
I . Cash flows from operating activities:		
Net profit for the period	27,348	23,367
Depreciation and amortization	8,092	8,223
Impairment losses	168	-
Gain on disposal of non-current assets	-	(3,592)
Finance expense (income)	(158)	(151)
Income tax expenses	2,192	9,037
Decrease (increase) in trade and other receivables	(6,560)	(370)
Decrease (increase) in inventories	2,043	(735)
Increase (decrease) in trade and other payables	617	(1,696)
Increase (decrease) in provisions and net defined benefit liabilities	346	(380)
Other	323	594
Subtotal	34,411	34,297
Interest received	68	154
Dividends received	580	518
Interest paid	(24)	(7)
Income tax paid	(5,990)	(12,268)
Net cash flows from (used in) operating activities	29,045	22,694
II. Cash flows from investing activities:		
Payments for acquisition of investments	(158)	(767)
Proceeds from sales of investments	514	1,025
Payments for acquisition of property, plant and equipment	(3,489)	(4,468)
Proceeds from sale of property, plant and equipment	—	4,338
Payments for acquisition of intangible assets	(3,051)	(2,165)
Other	(18)	(156)
Net cash flows from (used in) investing activities	(6,202)	(2,193)
II. Cash flows from financing activities:		
-		578
Proceeds from long-term loans	-	
Repayments of long-term loans	(6,862)	(4,098)
Capital contribution from non-controlling interests	838	(40,400)
Dividends paid	(10,471)	(10,493)
Other	196	20
Net cash flows from (used in) financing activities	(16,299)	(13,993)
IV. Net increase (decrease) in cash and cash equivalents	6,544	6,507
V. Cash and cash equivalents at the beginning of period	52,282	69,283
VI. Effect of exchange rate changes on cash and cash equivalents	1,222	(397)
VII. Cash and cash equivalents at the end of period	60,048	75,394

Revenue of major pharmaceuticals

				Year ended M	arch 31, 2018			Year ending M	larch 31, 2019	(JPY millio
Brand name Generic name/formulation	Therapeutic category	Region	Nine months ended December 31, 2017	Changes from same period of	Year ended March 31 Actual	Changes from same period of	Nine months ended December 31, 2018	Changes from same period of	Year ending March 31 Forecasts	Changes from sam period of
			Actual	previous year		previous year	Actual	previous year		previous ye
		Total	11,434	12.8%	14,944	16.0%	11,513	0.7%	15,443	3.3
Cravit	Bacterial	Japan	3,369	(11.3%)	4,105	(11.9%)	2,569	(23.7%)	3,415	(16.8
evofloxacin/ophthalmic solution	conjunctivitis	Asia	6,820	26.8%	9,225	32.2%	7,794	14.3%	10,154	10.1
		EMEA	1,245	29.6%	1,614	29.7%	1,150	(7.7%)	1,874	16.1
Farivid	Bacterial	Total	1,246	(5.1%)	1,581	2.6%	1,129	(9.4%)	1,507	(4.7
ofloxacin/ophthalmic solution	conjunctivitis	Japan	414	(9.2%)	508	(9.2%)	390	(6.0%)	458	(9.8
	,	Asia	832	(3.0%)	1,073	9.3%	740	(11.1%)	1,049	(2.2
Tapcom		Total	2,966	42.1%	3,892	39.1%	3,612	21.8%	4,950	27.2
afluprost-timolol maleate/	Glaucoma	Japan	1,953	10.9%	2,479	9.9%	2,002	2.5%	2,454	(1.09
combination ophthalmic solution		Asia	109	438.5%	158	317.6%	221	102.3%	289	82.7
		EMEA	904	195.1%	1,255	148.8%	1,389	53.7%	2,208	75.9
		Total	13,791	7.9%	17,844	8.8%	13,942	1.1%	18,883	5.8
Tapros	Glaucoma	Japan	7,590	(0.2%)	9,610	0.2%	7,526	(0.8%)	9,686	0.8
afluprost/ophthalmic solution		Asia	1,343	36.6%	1,807	47.3%	1,571	16.9%	2,026	12.1
		EMEA	4,857	15.9%	6,427	15.0%	4,846	(0.2%)	7,171	11.6
Cosopt		Total	18,622	9.4%	24,200	9.2%	16,956	(8.9%)	21,202	(12.4
lorzolamide hydrochloride-	0	Japan	9,057	0.7%	11,403	0.3%	7,247	(20.0%)	8,957	(21.4)
imolol	Glaucoma	Asia	2,440	30.1%	3,197	19.8%	2,769	13.5%	3,317	3.8
maleate/combination ophthalmic		EMEA	7,125	15.9%	9,600	18.1%	6,939	(2.6%)	8,928	(7.0
solution								. ,		
Timoptol		Total	1,183	1.3%	1,451	(3.8%)	780	(34.1%)	990	(31.8)
imolol maleate/	Glaucoma	Japan	662	(16.0%)	787	(18.7%)	398	(39.9%)	470	(40.3
ophthalmic solution		Asia	86	(4.3%)	116	(2.7%)	88	2.1%	112	(3.3
		EMEA	435	50.0%	548	30.1%	295	(32.3%)	408	(25.6
imoptol XE		Total	1,766	(6.6%)	2,221	(6.8%)	1,485	(15.9%)	1,976	(11.0
imolol maleate/	Glaucoma	Japan	1,144	(12.2%)	1,407	(12.9%)	922	(19.4%)	1,203	(14.5
ong-acting ophthalmic solution	Giauconia	Asia	79	20.1%	105	19.1%	82	4.1%	109	3.6
ong-acting ophilialinic solution		EMEA	544	4.2%	709	4.4%	481	(11.6%)	664	(6.3
		Total	3,587	7.3%	4,677	7.7%	3,436	(4.2%)	4,547	(2.8)
rusopt	01	Japan	1,325	(4.2%)	1,641	(5.2%)	1,175	(11.3%)	1,446	(11.9
lorzolamide hydrochloride/	Glaucoma	Asia	232	11.8%	327	20.9%	300	29.7%	462	41.3
ophthalmic solution		EMEA	2,031	15.9%	2,709	15.7%	1,961	(3.5%)	2,639	(2.6
Rescula		Total	1,200	(9.2%)	1,467	(10.1%)	964	(19.6%)	1,256	(14.4)
sopropyl unoprostone/	Glaucoma	TOtal	1,200	(3.270)	1,407	(10.170)	304	(13.070)	1,200	(14.4)
ophthalmic solution		Japan	1,200	(9.2%)	1,467	(10.1%)	964	(19.6%)	1,256	(14.49
Alesion		Total	7,515	19.4%	16,851	37.7%	7,703	2.5%	17,727	5.2
epinastine hydrochloride/	Allergy									
ophthalmic solution		Japan	7,515	19.4%	16,851	37.7%	7,703	2.5%	17,727	5.2
Flumetholon		Total	2,604	3.4%	3,497	4.5%	2,399	(7.9%)	3,141	(10.29
luorometholone/	Inflammation	Japan	1,566	(4.6%)	2,113	(5.0%)	1,132	(27.7%)	1,652	(21.89
ophthalmic solution		Asia	1,038	18.2%	1,385	23.2%	1,267	22.0%	1,490	7.6
Kary Uni		Total	3,518	9.0%	4,413	7.0%	3,199	(9.1%)	4,398	(0.39
pirenoxine/	Senile	Japan	2,187	(3.7%)	2,741	(3.2%)	2,069	(5.4%)	2,644	(3.6
ophthalmic solution	cataract	Asia	1,331	39.0%	1,672	29.6%	1,130	(15.1%)	1,755	4.9
Oftan Catachrom										
sytochrome C, adenosine,	Senile	Total	2,186	66.6%	2,695	21.2%	1,946	(11.0%)	2,800	3.9
picotinamide/ ophthalmic solution	cataract	EMEA	2,186	66.6%	2,695	21.2%	1,946	(11.0%)	2,800	3.9
Dpegan Hi odium hyaluronate/	Adjuvant for	Total	1,830	1.8%	2,304	0.8%	1,683	(8.1%)	2,109	(8.5
adjuvant for ophthalmic perations	ophthalmic operations	Japan	1,830	1.8%	2,304	0.8%	1,683	(8.1%)	2,109	(8.5
ylea *		Total	40,010	13.1%	51,517	14.1%	43,790	9.4%	54,473	5.7
flibercept/	Intravitreal VEGF inhibitor	Japan	40,010	13.1%	51,517	14.1%	43,790	9.4%	54,473	5.7
soulution for intravitreal injection		Tetal	14 404	1.00/	10 170	2.28/	14.040	E 00/	17 700	(0.5
lyalein	Dec	Total	14,124	1.6%	18,170	3.3%	14,943	5.8%	17,708	(2.5)
odium hyaluronate/ophthalmic	Dry eye	Japan	8,692	(8.8%)	10,772	(9.1%)	6,939	(20.2%)	8,541	(20.7
solution		Asia	5,432	24.4%	7,397	28.8%	8,004	47.3%	9,167	23.9
Diquas	Dev	Total	11,028	20.7%	14,286	19.6%	11,762	6.7%	16,087	12.6
iquafosol sodium/ophthalmic	Dry eye	Japan	9,937	17.0%	12,822	16.4%	10,784	8.5%	14,463	12.8
solution		Asia	1,091	69.6%	1,463	58.3%	978	(10.4%)	1,625	11.0
kervis	_	Total	1,661	99.2%	2,049	57.2%	2,415	45.4%	2,990	45.9
ciclosporin/ophthalmic solution	Dry eye	Asia	4	274.2%	68	- 1	324	- 1	437	537.6
		EMEA	1,657	99.0%	1,981	52.2%	2,091	26.2%	2,553	28.9
		Total	1,638	26.8%	2,092	14.0%	1,873	14.3%	3,196	52.8
Cationorm	Dry eye	Asia	176	104.0%	199	72.3%	118	(32.8%)	173	(13.2
	Dif Cyc	EMEA	1,267	20.0%	1,670	16.1%	1,503	(32.0%)	2,713	62.5
		US	1,207	30.1%	223	(20.4%)	251	28.8%	310	38.8
		Total	11,618	21.6%	14,594	16.3%	11,230	(3.3%)	16,498	13.0
TC pharmaceuticals			11,618	21.6%						
		Japan	11.412	20.6%	14,301	15.1%	11,014	(3.5%)	16,144	12.9

* Co-promoted product of Bayer Yakuhin, Ltd. (MAH)

Exchange rate (yen)	Major currency	3rd quarter ended December 31, 2017	Fiscal year ended March 31, 2018	3rd quarter ended December 31, 2018	Fiscal year to March 31, 2019(Forecasts)
	US dollar	111.75	110.94	111.15	110.00
	Euro	128.90	129.92	129.51	130.00
	CNY	16.64	16.84	16.57	17.00

Forecasts in this report are based on the currently available information. Actual results may differ materially depending on a number of factors including adverse economic conditions, etc.

As of February 5

■ Pipeline of prescription pharmaceuticals (Clinical Stage)

Generic name	Dev. code	Indication	Original/Licensor	Region	P1	P2	P3	NDA Filed	Approved Laund
diquafosol sodium	DE-089	Dry eye	Merck Sharp & Dohme Corp. (U.S.)	China					Sep-20
A dry eye treatment which	h stimulates se	ecretion of mucin and aqueous	components from the corneal	and conjunc	tival epith	elium. Its r	nechanisr	n of action is	different from exis
reatments. Launched in	December 201	0 in Japan. Launched in Octo	ber 2013 in Korea. Launched	in February	2016 in V	'ietnam. La	aunched i	n April 2016	in Thailand. Curre
seeking sequential approv	als for marketi	ng in Asia. Launched in Septer	mber 2018 in China.						
0							1 50		, <u>, , , , , , , , , , , , , , , , , , </u>
Generic name	Dev. code	Indication	Original/Licensor	Region	P1	P2	P3	NDA Filed	Approved Laund
				U.S.				-	
sirolimus	DE-109	Uveitis	Original	Japan				_	
			-	Europe					· · · · · · · · · · · · · · · · · · ·
				Asia				Apr-2015	L
-			c effect, etc. Started an additio	nal Phase 3	in Decen	1ber 2018	and planr	ing to compl	ete in January ~ J
2021 in the U.S. NDA filed	a in April 2015	in Asia.							
Generic name	Dev. code	Indication	Original/Licensor	Region	P1	P2	P3	NDA Filed	Approved Launc
tafluprost/		Glaucoma/	Co-development with	Ĵ					
timolol maleate	DE-111	Ocular hypertension	AGC	China					
A fixed dose combination	n drug of a pr	ostaglandin F2α derivative an	d a beta-adrenergic receptor	blocker. La	unched in	Japan in	Novembe	er 2014. Lau	nched successively
European countries since	January 2015	. Launched successively in As	ian countries since April 2016.	Started Pha	ise 3 in Ja	anuary 201	9 and pla	nning to com	plete in the 1st ha
FY2020 in China.									
						-		_	
Generic name	Dev. code	Indication	Original/Licensor	Region	P1	P2	P3	NDA Filed	Approved Launc
	201.0000								
epinastine		Allergic conjunctivitis	Nippon Boehringer	Japan				Sep-2018	
epinastine hydrochloride	DE-114A	Allergic conjunctivitis	Nippon Boehringer Ingelheim	Japan				Sep-2018	
hydrochloride	DE-114A				rug. Filed	for manufa			approval in Septem
hydrochloride An H ₁ receptor antagonis	DE-114A st with membra		Ingelheim tment for allergic conjunctivitis.		rug. Filed	for manufa			approval in Septerr
hydrochloride An H ₁ receptor antagonis 2018 and planning to rece	DE-114A st with membra	ane-stabilizing function, as treat	Ingelheim tment for allergic conjunctivitis. an.	High dose d			cturing an	d marketing	
hydrochloride An H ₁ receptor antagonis	DE-114A st with membra	ane-stabilizing function, as treat	Ingelheim tment for allergic conjunctivitis.	High dose d	rug. Filed P	for manufa		d marketing	approval in Septem
hydrochloride An H ₁ receptor antagonis 2018 and planning to rece	DE-114A st with membra sive approval in Dev. code	ane-stabilizing function, as treat	Ingelheim tment for allergic conjunctivitis. an.	High dose d Region U.S.			cturing an	d marketing	Approved Launc
hydrochloride An H, receptor antagoni 2018 and planning to rece Generic name	DE-114A st with membra	ane-stabilizing function, as treat a July ~ December 2019 in Japa Indication	Ingelheim tment for allergic conjunctivitis. an. Original/Licensor	High dose d Region U.S. Japan			cturing an	d marketing	
hydrochloride An H, receptor antagonis 2018 and planning to rece Generic name omidenepag isopropyl	DE-114A st with membra ive approval in Dev. code DE-117	ane-stabilizing function, as treat a July ~ December 2019 in Japa Indication Glaucoma/ Ocular hypertension	Ingelheim tment for allergic conjunctivitis. an. Original/Licensor Co-development with Ube Industries	High dose d Region U.S. Japan Asia	P1	P2	cturing an	d marketing	Approved Launco
hydrochloride An H ₁ receptor antagonis 2018 and planning to rece Generic name omidenepag isopropyl An EP2 receptor agonist v	DE-114A st with membravel in eive approval in Dev. code DE-117 with a new mee	ane-stabilizing function, as treat a July ~ December 2019 in Japa Indication Glaucoma/ Ocular hypertension	Ingelheim tment for allergic conjunctivitis. an. Original/Licensor Co-development with Ube Industries se 3 in September 2018 and pla	High dose d Region U.S. Japan Asia Inning to cor	P1	P2	cturing an	d marketing	Approved Launco
hydrochloride An H ₁ receptor antagonis 2018 and planning to rece Generic name omidenepag isopropyl An EP2 receptor agonist v	DE-114A st with membravel in eive approval in Dev. code DE-117 with a new mee	ane-stabilizing function, as treat a July ~ December 2019 in Japa Indication Glaucoma/ Ocular hypertension	Ingelheim tment for allergic conjunctivitis. an. Original/Licensor Co-development with Ube Industries	High dose d Region U.S. Japan Asia Inning to cor	P1	P2	cturing an	d marketing	Approved Launco
hydrochloride An H ₁ receptor antagonis 2018 and planning to rece Generic name omidenepag isopropyl An EP2 receptor agonist v	DE-114A st with membravel in eive approval in Dev. code DE-117 with a new mee	ane-stabilizing function, as treat a July ~ December 2019 in Japa Indication Glaucoma/ Ocular hypertension	Ingelheim tment for allergic conjunctivitis. an. Original/Licensor Co-development with Ube Industries se 3 in September 2018 and pla	High dose d Region U.S. Japan Asia Inning to cor	P1	P2	cturing an	MDA Filec	Approved Launco
hydrochloride An H, receptor antagonia 2018 and planning to rece Generic name omidenepag isopropyl An EP2 receptor agonist v 2018 in Japan. Started Ph Generic name	DE-114A st with membra ive approval in Dev. code DE-117 with a new mee asse 3 in Decen	ane-stabilizing function, as treat a July ~ December 2019 in Japa Indication Glaucoma/ Ocular hypertension chanism of action. Started Phase mber 2016 and planning to com	Ingelheim tment for allergic conjunctivitis. an. Original/Licensor Co-development with Ube Industries se 3 in September 2018 and pla splete in the 2nd half of FY2018	High dose d Region U.S. Japan Asia Inning to cor in Asia. Region	P1	P2	P3	MDA Filec	Approved Launc Nov-20 aunched in Noverr
hydrochloride An H ₁ receptor antagonis 2018 and planning to rece Generic name omidenepag isopropyl An EP2 receptor agonist v 2018 in Japan. Started Ph	DE-114A st with membra ive approval in Dev. code DE-117 with a new mec nase 3 in Decer	ane-stabilizing function, as treat a July ~ December 2019 in Japa Indication Glaucoma/ Ocular hypertension chanism of action. Started Phase mber 2016 and planning to com	Ingelheim tment for allergic conjunctivitis. an. Original/Licensor Co-development with Ube Industries se 3 in September 2018 and pla splete in the 2nd half of FY2018 Original/Licensor	High dose d Region U.S. Japan Asia Inning to cor in Asia.	P1	P2 January ~ 、	P3	MDA Filec	Approved Launc Nov-20 aunched in Noverr
hydrochloride An H, receptor antagonis 2018 and planning to rece Generic name omidenepag isopropyl An EP2 receptor agonist v 2018 in Japan. Started Ph Generic name carotuximab	DE-114A st with membra ive approval in Dev. code DE-117 with a new mer nase 3 in Decer Dev. code DE-122	ane-stabilizing function, as treat a July ~ December 2019 in Japa Indication Glaucoma/ Ocular hypertension chanism of action. Started Phase mber 2016 and planning to com Indication Wet Age-related macular degeneration	Ingelheim tment for allergic conjunctivitis. an. Original/Licensor Co-development with Ube Industries se 3 in September 2018 and pla splete in the 2nd half of FY2018 Original/Licensor TRACON Pharmaceuticals	High dose d Region U.S. Japan Asia Inning to cor in Asia. Region U.S.	P1 mplete in C P1 (Pt	P2 lanuary ~ . P2 nase 2a)	P3	MDA Filec	Approved Launc Nov-20 aunched in Novem
hydrochloride An H, receptor antagonis 2018 and planning to rece Generic name omidenepag isopropyl An EP2 receptor agonist v 2018 in Japan. Started Ph Generic name carotuximab	DE-114A st with membra ive approval in Dev. code DE-117 with a new mer nase 3 in Decer Dev. code DE-122	ane-stabilizing function, as treat a July ~ December 2019 in Japa Indication Glaucoma/ Ocular hypertension chanism of action. Started Phase mber 2016 and planning to com Indication Wet Age-related macular degeneration	Ingelheim tment for allergic conjunctivitis. an. Original/Licensor Co-development with Ube Industries se 3 in September 2018 and pla nplete in the 2nd half of FY2018 Original/Licensor TRACON	High dose d Region U.S. Japan Asia Inning to cor in Asia. Region U.S.	P1 mplete in C P1 (Pt	P2 lanuary ~ . P2 nase 2a)	P3	MDA Filec	Approved Launc Nov-20 aunched in Novem
hydrochloride An H, receptor antagonia 2018 and planning to rece Generic name omidenepag isopropyl An EP2 receptor agonist v 2018 in Japan. Started Ph Generic name carotuximab An intravitreal injection of	DE-114A st with membra ive approval in Dev. code DE-117 with a new mec ase 3 in Decer Dev. code DE-122 anti-endoglin a	ane-stabilizing function, as treat a July ~ December 2019 in Japa Indication Glaucoma/ Ocular hypertension chanism of action. Started Phas mber 2016 and planning to com Indication Wet Age-related macular degeneration untibody. Started Phase 2a in Ju	Ingelheim Iment for allergic conjunctivitis. an. Original/Licensor Co-development with Ube Industries se 3 in September 2018 and pla aplete in the 2nd half of FY2018 Original/Licensor TRACON Pharmaceuticals uly 2017 and planning to comple	High dose d Region U.S. Japan Asia Inning to cor in Asia. Region U.S. U.S. ete in the 2n	P1 nplete in 3 P1 (Pt d half of F	P2 lanuary ~ P2 nase 2a) Y2019 for	P3	MDA Filec	Approved Launce Nov-20 aunched in Noverr Approved Launce
hydrochloride An H, receptor antagonis 2018 and planning to rece Generic name omidenepag isopropyl An EP2 receptor agonist v 2018 in Japan. Started Ph Generic name carotuximab	DE-114A st with membra ive approval in Dev. code DE-117 with a new mer nase 3 in Decer Dev. code DE-122	ane-stabilizing function, as treat a July ~ December 2019 in Japa Indication Glaucoma/ Ocular hypertension chanism of action. Started Phase mber 2016 and planning to com Indication Wet Age-related macular degeneration Intibody. Started Phase 2a in Junition	Ingelheim tment for allergic conjunctivitis. an. Original/Licensor Co-development with Ube Industries se 3 in September 2018 and pla nplete in the 2nd half of FY2018 Original/Licensor TRACON Pharmaceuticals uly 2017 and planning to complete Original/Licensor	High dose d Region U.S. Japan Asia Inning to cor in Asia. Region U.S. ete in the 2n Region	P1 nplete in . P1 (Pr d half of F	P2 lanuary ~ 、 P2 nase 2a) Y2019 for P2	P3	MDA Filec	Approved Launc Nov-20 aunched in Novem
hydrochloride An H, receptor antagonia 2018 and planning to rece Generic name omidenepag isopropyl An EP2 receptor agonist v 2018 in Japan. Started Ph Generic name carotuximab An intravitreal injection of	DE-114A st with membra ive approval in Dev. code DE-117 with a new mec ase 3 in Decer Dev. code DE-122 anti-endoglin a	ane-stabilizing function, as treat a July ~ December 2019 in Japa Indication Glaucoma/ Ocular hypertension chanism of action. Started Phase mber 2016 and planning to com Indication Wet Age-related macular degeneration Intibody. Started Phase 2a in Junitibody. Started Phase 2a in Juniti	Ingelheim tment for allergic conjunctivitis. an. Original/Licensor Co-development with Ube Industries se 3 in September 2018 and pla nplete in the 2nd half of FY2018 Original/Licensor TRACON Pharmaceuticals uly 2017 and planning to completed Original/Licensor ONO	High dose d Region U.S. Japan Asia Asia Inning to cor in Asia. Region U.S. tete in the 2n Region U.S.	P1 mplete in . P1 (Pr d half of F P1 (Pr	P2 January ~ (P2 hase 2a) Y2019 for P2 hase 2b)	P3	MDA Filec	Approved Launce Nov-20 aunched in Noverr Approved Launce
hydrochloride An H, receptor antagonis 2018 and planning to rece Generic name omidenepag isopropyl An EP2 receptor agonist v 2018 in Japan. Started Ph Generic name carotuximab An intravitreal injection of Generic name sepetaprost	DE-114A st with membra ive approval in Dev. code DE-117 with a new mec nase 3 in Decen Dev. code DE-122 anti-endoglin a Dev. code DE-126	ane-stabilizing function, as treat a July ~ December 2019 in Japa Indication Glaucoma/ Ocular hypertension chanism of action. Started Phase mber 2016 and planning to com Indication Wet Age-related macular degeneration intibody. Started Phase 2a in Junition Glaucoma/ Ocular hypertension	Ingelheim Iment for allergic conjunctivitis. an. Original/Licensor Co-development with Ube Industries se 3 in September 2018 and pla plete in the 2nd half of FY2018 Original/Licensor TRACON Pharmaceuticals uly 2017 and planning to comple Original/Licensor ONO PHARMACEUTICAL	High dose d Region U.S. Japan Asia Inning to corrin Asia. Region U.S. Lust Region U.S. Interview Region U.S. Interview Region U.S. Japan	P1 mplete in . P1 (Pr d half of F P1 (Pr (Pr	P2 lanuary ~ . P2 nase 2a) Y2019 for r P2 nase 2b) nase 2b)	P3	MDA Filec	Approved Launce Nov-20 aunched in Nover Approved Launce
hydrochloride An H, receptor antagonis 2018 and planning to rece Generic name omidenepag isopropyl An EP2 receptor agonist v 2018 in Japan. Started Ph Generic name carotuximab An intravitreal injection of Generic name sepetaprost A prostaglandin analogue	DE-114A st with membra ive approval in Dev. code DE-117 with a new mer iase 3 in Decer Dev. code DE-122 anti-endoglin a Dev. code DE-126 eye drop drug	ane-stabilizing function, as treat a July ~ December 2019 in Japa Indication Glaucoma/ Ocular hypertension chanism of action. Started Phase mber 2016 and planning to com Indication Wet Age-related macular degeneration Indication Glaucoma/ Ocular hypertension product with a novel mode of a	Ingelheim tment for allergic conjunctivitis. an. Original/Licensor Co-development with Ube Industries se 3 in September 2018 and pla nplete in the 2nd half of FY2018 Original/Licensor TRACON Pharmaceuticals uly 2017 and planning to completed Original/Licensor ONO	High dose d Region U.S. Japan Asia Inning to corrin Asia. Region U.S. Lust Region U.S. Interview Region U.S. Interview Region U.S. Japan	P1 mplete in . P1 (Pr d half of F P1 (Pr (Pr	P2 lanuary ~ . P2 nase 2a) Y2019 for r P2 nase 2b) nase 2b)	P3	MDA Filec	Approved Launce Nov-20 aunched in Nover Approved Launce
hydrochloride An H, receptor antagonis 2018 and planning to rece Generic name omidenepag isopropyl An EP2 receptor agonist v 2018 in Japan. Started Ph Generic name carotuximab An intravitreal injection of Generic name sepetaprost A prostaglandin analogue	DE-114A st with membra ive approval in Dev. code DE-117 with a new mer iase 3 in Decer Dev. code DE-122 anti-endoglin a Dev. code DE-126 eye drop drug	ane-stabilizing function, as treat a July ~ December 2019 in Japa Indication Glaucoma/ Ocular hypertension chanism of action. Started Phase mber 2016 and planning to com Indication Wet Age-related macular degeneration Indication Glaucoma/ Ocular hypertension product with a novel mode of a	Ingelheim Iment for allergic conjunctivitis. an. Original/Licensor Co-development with Ube Industries se 3 in September 2018 and pla plete in the 2nd half of FY2018 Original/Licensor TRACON Pharmaceuticals uly 2017 and planning to comple Original/Licensor ONO PHARMACEUTICAL	High dose d Region U.S. Japan Asia Inning to corrin Asia. Region U.S. Lust Region U.S. Interview Region U.S. Interview Region U.S. Japan	P1 mplete in . P1 (Pr d half of F P1 (Pr (Pr	P2 lanuary ~ . P2 nase 2a) Y2019 for r P2 nase 2b) nase 2b)	P3	MDA Filec	Approved Launce Nov-20 aunched in Nover Approved Launce
hydrochloride An H, receptor antagonia 2018 and planning to rece Generic name omidenepag isopropyl An EP2 receptor agonist v 2018 in Japan. Started Ph Generic name carotuximab An intravitreal injection of Generic name sepetaprost A prostaglandin analogue Started Phase 2b in July 2	DE-114A st with membra sive approval in Dev. code DE-117 with a new meo nase 3 in Decer Dev. code DE-122 anti-endoglin a Dev. code DE-126 eye drop drug 2017 in the U.S	ane-stabilizing function, as treat a July ~ December 2019 in Japa Indication Glaucoma/ Ocular hypertension chanism of action. Started Phase mber 2016 and planning to com Indication Wet Age-related macular degeneration Intibody. Started Phase 2a in Ju Indication Glaucoma/ Ocular hypertension product with a novel mode of a 5 and Japan.	Ingelheim Iment for allergic conjunctivitis. an. Original/Licensor Co-development with Ube Industries se 3 in September 2018 and pla uplete in the 2nd half of FY2018 Original/Licensor TRACON Pharmaceuticals uly 2017 and planning to comple Original/Licensor ONO PHARMACEUTICAL action that is both FP and EP3	High dose d Region U.S. Japan Asia Inning to cor in Asia. Region U.S. dete in the 2n Region U.S. Japan C.S. Japan C.S. Japan C.S. C.	P1 mplete in a P1 (Pr d half of F P1 (Pr (Pr (Pr (Pr (Pr al agonist	P2 lanuary ~ (P2 nase 2a) Y2019 for P2 nase 2b) for the tre	P3	d marketing	Approved Launce Nov-20 aunched in Noverr Approved Launce Approved Launce d ocular hypertens
hydrochloride An H, receptor antagonis 2018 and planning to rece Generic name omidenepag isopropyl An EP2 receptor agonist v 2018 in Japan. Started Ph Generic name carotuximab An intravitreal injection of Generic name sepetaprost	DE-114A st with membra ive approval in Dev. code DE-117 with a new mer iase 3 in Decer Dev. code DE-122 anti-endoglin a Dev. code DE-126 eye drop drug	ane-stabilizing function, as treat a July ~ December 2019 in Japa Indication Glaucoma/ Ocular hypertension chanism of action. Started Phase mber 2016 and planning to com Indication Wet Age-related macular degeneration Indication Glaucoma/ Ocular hypertension product with a novel mode of a	Ingelheim Iment for allergic conjunctivitis. an. Original/Licensor Co-development with Ube Industries se 3 in September 2018 and pla plete in the 2nd half of FY2018 Original/Licensor TRACON Pharmaceuticals uly 2017 and planning to comple Original/Licensor ONO PHARMACEUTICAL action that is both FP and EP3 I Original/Licensor	High dose d Region U.S. Japan Asia Inning to corrin Asia. Region U.S. Lust Region U.S. Interview Region U.S. Interview Region U.S. Japan	P1 mplete in . P1 (Pr d half of F P1 (Pr (Pr	P2 lanuary ~ . P2 nase 2a) Y2019 for r P2 nase 2b) nase 2b)	P3	d marketing	Approved Launce Nov-20 aunched in Nover Approved Launce
hydrochloride An H, receptor antagonis 2018 and planning to rece Generic name omidenepag isopropyl An EP2 receptor agonist v 2018 in Japan. Started Ph Generic name carotuximab An intravitreal injection of Generic name sepetaprost A prostaglandin analogue Started Phase 2b in July 2 Generic name	DE-114A st with membra ive approval in Dev. code DE-117 with a new meo iase 3 in Decen Dev. code DE-122 anti-endoglin a Dev. code DE-126 eye drop drug 2017 in the U.S	ane-stabilizing function, as treat a July ~ December 2019 in Japa Indication Glaucoma/ Ocular hypertension chanism of action. Started Phase mber 2016 and planning to com Indication Wet Age-related macular degeneration Indication Glaucoma/ Ocular hypertension product with a novel mode of a a and Japan.	Ingelheim Iment for allergic conjunctivitis. an. Original/Licensor Co-development with Ube Industries se 3 in September 2018 and pla plete in the 2nd half of FY2018 Original/Licensor TRACON Pharmaceuticals uly 2017 and planning to comple Original/Licensor ONO PHARMACEUTICAL action that is both FP and EP3 i Original/Licensor Singapore Health	High dose d Region U.S. Japan Asia Inning to cor in Asia. Region U.S. Ete in the 2n Region U.S. Japan receptors du Region	P1 mplete in a P1 (Pr d half of F P1 (Pr (Pr (Pr (Pr (Pr al agonist	P2 lanuary ~ (P2 nase 2a) Y2019 for P2 nase 2b) for the tre	P3	d marketing	Approved Launce Nov-20 aunched in Noverr Approved Launce Approved Launce d ocular hypertens
hydrochloride An H, receptor antagonia 2018 and planning to rece Generic name omidenepag isopropyl An EP2 receptor agonist v 2018 in Japan. Started Ph Generic name carotuximab An intravitreal injection of Generic name sepetaprost A prostaglandin analogue Started Phase 2b in July 2	DE-114A st with membra sive approval in Dev. code DE-117 with a new meo nase 3 in Decer Dev. code DE-122 anti-endoglin a Dev. code DE-126 eye drop drug 2017 in the U.S	ane-stabilizing function, as treat a July ~ December 2019 in Japa Indication Glaucoma/ Ocular hypertension chanism of action. Started Phase mber 2016 and planning to com Indication Wet Age-related macular degeneration Intibody. Started Phase 2a in Ju Indication Glaucoma/ Ocular hypertension product with a novel mode of a 5 and Japan.	Ingelheim Iment for allergic conjunctivitis. an. Original/Licensor Co-development with Ube Industries se 3 in September 2018 and pla plete in the 2nd half of FY2018 Original/Licensor TRACON Pharmaceuticals uly 2017 and planning to comple Original/Licensor ONO PHARMACEUTICAL action that is both FP and EP3 I Original/Licensor	High dose d Region U.S. Japan Asia Inning to cor in Asia. Region U.S. dete in the 2n Region U.S. Japan C.S. Japan C.S. Japan C.S. C.	P1 mplete in a P1 (Pr d half of F P1 (Pr (Pr (Pr (Pr (Pr al agonist	P2 lanuary ~ (P2 nase 2a) Y2019 for P2 nase 2b) for the tre	P3	d marketing	Approved Launce Nov-20 aunched in Noverr Approved Launce Approved Launce d ocular hypertens

-	Dev. code	Indication	Original/Licensor	Region	P1 P2 P3 NDA Filed Approve		Approved	Launched		
	DE-128	Glaucoma	Original	U.S.						
	(MicroShunt)		Onginar	Europe						
In August 2016, Santen	acquired InnFo	ocus, developer of MicroShunt	, a drainage implant device de	esigned to	lower and	sustain intr	aocular pre	ssure (IOP) for the tre	eatment of
primary open-angle glaucoma through the drainage of aqueous humor. Conducting Phase 2/3 in the U.S. and Europe in advance of application to FDA. Planning to complete PMA										
rolling submission in 2019	rolling submission in 2019 and launch in 2020 in U.S. Received CE Mark in Europe.									

Generic name	Dev.code	Indication	Original/Licensor	Region	P1	P2	P3	NDA Filed	Approved	Launched
ai at a sa a si a	DE-076B	Severe keratitis in	Original	U.S.						
ciclosporin	(Cyclokat)	patients with dry eye	Original	Asia					ec-2017	
An ophthalmic emulsion	to treat severe	keratitis in adult patients with	dry eye through an immunos	uppressive	effect. Cat	ionic emuls	sion techno	logy has er	nhanced oc	ular tissue
penetration. Launched in July 2015 in Germany and U.K. with successive launches following in European countries. Launched in December 2017 in Thailand and Korea with										
successive launches follo	wing in Asian d	countries.								

Generic name	Dev. code	Indication	Original/Licensor	Region	P1	P2	P3	NDA Filed	Approved	Launched
	DE-076C	Vereel		Europe	e Oct					oct-2018
ciclosporin		Vernal keratoconjunctivitis	Original	Asia			N	lov-2018		
	(Vekacia)	keratoconjunctivitis		Others				D	ec-2018	
An ophthalmic emulsion which improves vernal keratoconjunctivitis by immunosuppressive effect. Cationic emulsion technology has enhanced ocular tissue penetration. Received										

the Marketing Authorization Application approval from the European Commission Agency in July 2018 and launched in October 2018 in U.K. Filed for manufacturing and marketing approval in November 2018 and planning to receive approval in July ~ December 2019 in Asia. Received marketing approval in December 2018 and planning to launch in 2019 in Canada.

Generic name	Dev. code	Indication	Original/Licensor	Region	egion P1 P2 F		P3	NDA Filed	Approved	Launched
latananraat	DE-130A	Glaucoma/	Original	Furana						
latanoprost	(Catioprost)	Ocular hypertension	Original	Europe						
An ophthalmic emulsion of	An ophthalmic emulsion of a prostaglandin F ₂₀ derivative, for the treatment of glaucoma and ocular hypertension.									

Changes from Q2 FY18 (November 7, 2018)

Dev. code	Changes
DE-109	Started an additional Phase 3 in December 2018 in the U.S.
DE-111	Started Phase 3 in January 2019 in China.
DE-117	Launched in November 2018 in Japan.
DE-076C (Vekacia)	Filed in November 2018 in Asia and received marketing approval in December 2018 in Canada.

Other consolidated information

Capital expenditures (JPY millions) Year Year Nine months ended Nine months ended ended ending December 31, 2017 December 31, 2018 March 31, 2018 March 31, 2019 Actual Forecast Consolidated 4,460 5,445 5,072 9,000

Depreciation and amortization

	Nine months endedYear endedNine months endedDecember 31, 2017ended March 31, 2018December 31, 2018			Year ending March 31, 2019
		Actual		Forecast
Manufacturing cost	1,445	1,950	1,397	2,040
Selling, general and administrative expenses	1,076	1,453	1,105	1,520
R&D expenses	574	752	489	700
Consolidated total	3,095	4,155	2,991	4,260

Note: Excluding amortization on intangible assets associated with products and long-term advance expense.

Amortization on intangible assets associated with products

	Nine months ended December 31, 2017	Year ended March 31, 2018	Nine months ended December 31, 2018	Year ending March 31, 2019
		Actual		Forecast
Intangible assets (Merck products)	4,140	5,592	4,356	5,810
Intangible assets (Ikervis)	548	736	550	740
Other	308	412	327	380
Consolidated total	4,996	6,740	5,233	6,930

Research & Development expenses

	Nine months ended December 31, 2017	Year ended March 31, 2018	Nine months ended December 31, 2018	Year ending March 31, 2019
		Forecast		
Consolidated	17,895	24,398	17,091	25,000
Percent of revenue	10.6%	10.8%	9.9%	10.5%

*Forecasts in this report are based on the currently available information. Actual results may differ materially depending on a number of factors including adverse economic conditions, etc.

(JPY millions)

(JPY millions)

(JPY millions)