

Kyowa Hakko Kirin Co., Ltd.

Appendix to the Consolidated Financial Summary (IFRS) Fiscal 2018 Third Quarter

(January 1, 2018 - September 30, 2018)

- These materials were made as a supplement to the Kessan Tanshin (Consolidated Financial Summary, IFRS), disclosed at the Tokyo Stock Exchange on October 30, 2018 for the first nine months of Fiscal 2018, from January 1, 2018 to September 30, 2018.
- This document is an English translation of parts of the Japanese-language original. The statements, including earnings forecasts, contained in these materials are based on the information currently available to the Company and on certain assumptions deemed to be reasonable by management. As such, they do not constitute guarantees by the Company of future performance. Actual results may differ materially from these projections for a wide variety of reasons.
- Figures presented in these materials have been rounded to the nearest tenth.
- Figures inside parenthesis presented in these materials indicate negative values.

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The average exchange rates for each period were as follows:

Unit: Yen

	FY 2017 results		FY 2018 results	FY 2018 forecasts
	Jan - Sep	Jan - Dec	Jan - Sep	Jan - Dec
USD	112	112	109	110
EUR	124	126	131	130
GBP	142	144	149	150

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I. Consolidated Financial Results (IFRS)

1. Trends in consolidated profit

Unit: Billions of yen

	FY 2017 results		FY 2018 results		FY 2018 forecasts	
	Jan - Sep	Jan - Dec	Jan - Sep	Change amount	Jan - Dec	%
Revenue	261.3	353.4	253.9	(7.3)	335.0	95%
Cost of sales	(96.3)	(129.1)	(88.0)	8.3	(114.0)	88%
Gross profit	165.0	224.3	166.0	1.0	221.0	99%
Selling, general and administrative expenses	(81.4)	(113.0)	(85.9)	(4.5)	(117.0)	104%
Research and development expenses	(35.8)	(49.2)	(34.4)	1.4	(49.5)	101%
Share of profit (loss) of investments accounted for using equity method	(3.1)	(4.5)	0.3	3.4	(0.5)	11%
Core operating profit	44.7	57.7	45.9	1.3	54.0	94%
Other income	1.2	6.6	18.5	17.3		
Other expenses	(2.3)	(8.4)	(1.1)	1.2		
Finance income (costs)	0.2	(0.1)	(0.5)	(0.7)		
Profit before tax	43.7	55.8	62.8	19.0	71.0	127%
Income tax expense	(14.3)	(13.0)	(15.4)	(1.1)		
Profit	29.5	42.9	47.4	18.0	52.0	121%

EPS (¥/share)	53.83	78.38	86.62	32.79	94.99	-
Annual dividend (¥/share)		27.00			30.00	-
Dividend payout ratio (%)		34.4			31.6	-
ROE (%)		7.2				-

* "FY 2018 forecasts" in these materials, excluding those of share of profit (loss) of investments accounted for using equity method, core operating profit, profit before tax, profit, EPS, and dividend payout ratio, which have been revised, still remain unchanged from the earnings forecasts released on February 8, 2018.

2. Revenue by geographic region

Unit: Billions of yen

	FY 2017 results			FY 2018 results		FY 2018 forecasts	
	Jan - Sep	Jan - Dec	Percentage of consolidated revenue	Jan - Sep	Percentage of consolidated revenue	Jan - Dec	Percentage of consolidated revenue
Japan	178.8	240.9	68.2%	164.9	65.0%	221.2	66.0%
International	82.5	112.5	31.8%	89.0	35.0%	113.8	34.0%
Americas	16.6	24.7	7.0%	21.6	8.5%	29.7	8.9%
Europe	42.4	56.0	15.8%	42.7	16.8%	55.2	16.5%
Asia	22.4	30.2	8.6%	24.0	9.4%	28.0	8.3%
Others	1.0	1.6	0.5%	0.6	0.2%	0.9	0.3%
Total consolidated revenue	261.3	353.4	100%	253.9	100%	335.0	100%

* Revenue is classified by region or country based on location of customer.

3. Other trends

(1) Research and development expenses

Unit: Billions of yen

	FY 2017 results		FY 2018 results	FY 2018 forecasts
	Jan - Sep	Jan - Dec	Jan - Sep	Jan - Dec
Research and development (R&D) expenses <i>R&D expenses to revenue ratio</i>	35.9 13.7%	49.2 13.9%	34.5 13.6%	49.5 14.8%
Pharmaceuticals <i>R&D expenses to revenue ratio</i>	33.6 16.5%	46.1 16.7%	32.3 16.3%	46.5 17.7%
Bio-Chemicals	2.3	3.1	2.2	3.0

(2) Capital expenditures (property, plant and equipment)

Unit: Billions of yen

	FY 2017 results		FY 2018 results	FY 2018 forecasts
	Jan - Sep	Jan - Dec	Jan - Sep	Jan - Dec
Capital expenditures	9.9	14.3	8.1	12.5
Pharmaceuticals	5.3	6.6	3.3	5.3
Bio-Chemicals	4.6	7.7	4.8	7.2

(3) Depreciation and amortization (property, plant and equipment and intangible assets)

Unit: Billions of yen

	FY 2017 results		FY 2018 results	FY 2018 forecasts
	Jan - Sep	Jan - Dec	Jan - Sep	Jan - Dec
Depreciation and amortization <i>Amortization of intangible assets</i>	16.4 5.8	22.0 7.8	16.0 6.3	24.6 9.1
Pharmaceuticals <i>Amortization of intangible assets</i>	11.4 5.7	15.3 7.7	11.7 6.2	17.3 9.1
Bio-Chemicals	5.0	6.7	4.4	7.3

II. Consolidated Statement of Cash Flows (IFRS)

Unit: Billions of yen

	FY 2017 results		FY 2018 results	
	Jan - Sep	Jan - Dec	Jan - Sep	Change amount
Cash flows from operating activities	49.1	64.9	46.4	(2.7)
Cash flows from investing activities	(35.0)	(45.3)	(30.8)	4.2
Cash flows from financing activities	(13.9)	(18.3)	(16.5)	(2.6)
Effect of exchange rate changes on cash and cash equivalents	1.1	1.3	(0.4)	(1.5)
Net increase (decrease) in cash and cash equivalents	1.3	2.7	(1.3)	(2.6)
Transfer to assets held for sale	(0.0)	(1.1)	1.1	1.1
Cash and cash equivalents at beginning of period	13.1	13.1	14.7	1.6
Cash and cash equivalents at end of period*	14.4	14.7	14.5	0.1

* Cash reserves at end of period

Cash and cash equivalents at end of period	14.4	14.7	14.5	0.1
+ Loans receivable from parent in excess of three months	135.2	143.2	179.6	44.4
+ Time deposits whose maturity periods exceed three months	0.0	0.0	0.0	0.0
Cash reserves at end of period	149.6	157.9	194.1	44.5

III. Core Operating Profit or Loss by Segment (IFRS)
1. Pharmaceuticals segment
Unit: Billions of yen

	FY 2017 results		FY 2018 results		FY 2018 forecasts	
	Jan - Sep	Jan - Dec	Jan - Sep	Change amount	Jan - Dec	%
Revenue	203.0	275.8	197.7	(5.3)	262.0	95%
Revenue from external customers	202.3	274.8	196.9	(5.4)	261.0	95%
Japan	147.1	197.8	134.3	(12.8)	181.0	91%
International	55.2	76.9	62.6	7.4	80.0	104%
International revenue ratio	27%	28%	32%	-	31%	-
Americas	9.3	15.0	14.6	5.3	20.8	138%
Europe	31.4	41.5	31.8	0.4	39.8	96%
Asia	14.1	19.6	16.2	2.1	18.8	96%
Others	0.5	0.8	0.1	(0.4)	0.6	82%
Inter-segment revenue	0.7	1.0	0.8	0.0	1.0	101%
Cost of sales	(58.8)	(79.1)	(53.9)	4.9	(70.0)	89%
Gross profit	144.2	196.7	143.8	(0.4)	192.0	98%
Selling, general and administrative expenses	(68.3)	(95.5)	(72.3)	(4.0)	(99.0)	104%
Research and development expenses	(33.6)	(46.1)	(32.3)	1.3	(46.5)	101%
Share of profit (loss) of investments accounted for using equity method	(3.1)	(4.5)	0.3	3.4	(0.5)	11%
Core operating profit	39.2	50.5	39.5	0.3	46.0	91%

* Revenue is classified by region or country based on location of customer.

* The figures represent amounts before elimination of inter-segment transactions.

2. Bio-Chemicals segment
Unit: Billions of yen

	FY 2017 results		FY 2018 results		FY 2018 forecasts	
	Jan - Sep	Jan - Dec	Jan - Sep	Change amount	Jan - Dec	%
Revenue	60.7	81.1	58.4	(2.3)	76.0	94%
Revenue from external customers	59.0	78.6	57.0	(2.0)	74.0	94%
Japan	31.7	43.0	30.7	(1.0)	40.2	93%
International	27.3	35.6	26.3	(1.0)	33.8	95%
International revenue ratio	46%	45%	46%	-	46%	-
Americas	7.4	9.7	7.1	(0.3)	9.0	93%
Europe	11.0	14.4	11.0	(0.0)	15.4	107%
Asia	8.3	10.7	7.8	(0.5)	9.2	86%
Others	0.5	0.8	0.5	(0.0)	0.3	32%
Inter-segment revenue	1.7	2.5	1.4	(0.3)	2.0	79%
Cost of sales	(40.0)	(53.3)	(36.5)	3.5	(47.0)	88%
Gross profit	20.6	27.8	21.9	1.2	29.0	104%
Selling, general and administrative expenses	(13.2)	(17.6)	(13.7)	(0.5)	(18.0)	102%
Research and development expenses	(2.3)	(3.1)	(2.2)	0.0	(3.0)	98%
Core operating profit	5.2	7.2	6.0	0.8	8.0	111%

* Revenue is classified by region or country based on location of customer.

* The figures represent amounts before elimination of inter-segment transactions.

IV. Revenue from Main Products of Pharmaceuticals Segment (IFRS)

<Accumulative>

Unit: Billions of yen

Indication / Product name		FY 2017 results		FY 2018 results		FY 2018 forecasts		
		Jan - Sep	Jan - Dec	Jan - Sep	Change amount	Jan - Dec	%	
Japan	Renal anemia treatment drug Nesp	41.0	56.3	39.2	(1.8)	52.4	93%	
	Secondary hyperparathyroidism Regpara	13.9	18.5	10.7	(3.2)	13.2	71%	
	Secondary hyperparathyroidism Orkedia	-	-	1.1	1.1	3.5	-	
	Secondary hyperparathyroidism Rocaltrol	3.0	4.1	2.8	(0.2)	3.5	87%	
	Type-2 diabetes Onglyza	5.2	7.3	5.4	0.1	7.6	103%	
	Cardiovascular (Hypertension & angina pectoris) Coniel	4.7	6.3	3.6	(1.1)	4.9	77%	
	Agent for decreasing the incidence of febrile neutropenia G-Lasta	12.8	18.1	14.8	2.0	20.1	111%	
	Transdermal persistent pain Fentos	4.3	6.0	4.0	(0.4)	5.7	96%	
	Anticancer Poteligeo	1.4	1.9	1.3	(0.0)	1.8	95%	
	Anticancer Rituximab BS [KHK]	-	-	2.4	2.4	3.2	-	
	Chronic idiopathic thrombocytopenic purpura Romiplate	2.5	3.4	2.4	(0.1)	3.4	100%	
	Antiallergenic Allelock	12.2	15.9	9.7	(2.4)	11.7	74%	
	Antiallergic eyedrops Patanol	10.8	12.8	11.5	0.7	12.1	94%	
	Ulcerative colitis Asacol	3.2	4.2	2.2	(1.0)	3.4	80%	
	Psoriasis vulgaris Dovobet	3.9	5.3	4.2	0.3	6.2	117%	
	Psoriasis Lumicef	0.5	1.0	1.4	0.9	2.7	281%	
	Parkinson's disease Nouriastr	6.1	8.5	6.8	0.7	9.4	111%	
	Antiepileptic Depakene	4.8	6.5	4.0	(0.9)	5.4	82%	
	Technology out-licensing		2.8	2.4	2.2	(0.6)	4.6	191%
	International	Renal anemia treatment drug Nesp	4.3	5.7	4.7	0.3	5.8	103%
		Secondary hyperparathyroidism Regpara	1.6	2.2	2.3	0.7	2.8	123%
		Agent for decreasing the incidence of febrile neutropenia Neulasta/Peglasta	1.5	2.0	1.7	0.2	2.0	101%
		Neutropenia treatment drug Gran	3.8	5.2	4.2	0.3	4.4	86%
X-linked hypophosphatemia (XLH) Crysvita		-	-	3.6	3.6	-	-	
Cancer pain Abstral		8.8	11.9	9.5	0.7	13.2	111%	
Cancer pain PecFent		3.1	4.1	3.2	0.2	5.2	128%	
Anticancer Mitomycin-C		2.5	3.4	1.7	(0.7)	2.0	57%	
Chemotherapy-induced nausea and vomiting drug Sancuso		2.0	3.0	2.1	0.0	2.8	92%	
Opioid-induced constipation (OIC) Moventig		0.5	0.8	1.0	0.5	2.4	299%	
Replacement therapy with testosterone for male hypogonadism Tostran/Fortesta		1.7	2.2	2.1	0.4	2.1	93%	
Osteoporosis drug Adcal-D3		2.6	3.6	2.7	0.0	3.3	93%	
Technology out-licensing		10.8	16.0	13.9	3.1	17.3	108%	

* Revenue is classified as Japan or International (other than Japan) based on customer location.

* Revenue listed as "Technology out-licensing" specifies revenue from the upfront payment, milestone revenue, and running royalties revenue that are obtained based on licensing agreements recognizing the granting to third parties the rights for development, manufacturing and sales of the Group's pipeline compounds or the use of technology, etc.

IV. Revenue from Main Products of Pharmaceuticals Segment (IFRS)

<Quarterly>

Unit: Billions of yen




Indication / Product name	FY 2017 results			FY 2018 results				
	Jan - Mar	Apr - Jun	Jul - Sep	Jan - Mar	Apr - Jun	Jul - Sep	Change amount	
Japan	Renal anemia treatment drug Nesp	12.5	14.2	14.4	12.0	13.6	13.6	(0.7)
	Secondary hyperparathyroidism Regpara	4.6	4.8	4.5	3.6	4.2	2.9	(1.5)
	Secondary hyperparathyroidism Orkedia	-	-	-	-	0.4	0.8	0.8
	Secondary hyperparathyroidism Rocaltrol	0.9	1.0	1.0	0.9	1.0	0.9	(0.1)
	Type-2 diabetes Onglyza	1.6	1.8	1.9	1.6	1.9	1.8	0.0
	Cardiovascular (Hypertension & angina pectoris) Coniel	1.6	1.6	1.5	1.2	1.3	1.1	(0.4)
	Agent for decreasing the incidence of febrile neutropenia G-Lasta	3.8	4.3	4.6	4.3	5.1	5.3	0.7
	Transdermal persistent pain Fentos	1.3	1.6	1.4	1.2	1.4	1.4	(0.1)
	Anticancer Poteligeo	0.4	0.5	0.4	0.4	0.5	0.4	0.0
	Anticancer Rituximab BS [KHK]	-	-	-	0.3	0.8	1.3	1.3
	Chronic idiopathic thrombocytopenic purpura Romiplate	0.8	0.9	0.8	0.7	0.8	0.8	0.0
	Antiallergenic Allelock	5.6	3.5	3.0	4.6	2.9	2.3	(0.8)
	Antiallergic eyedrops Patanol	7.0	1.8	1.9	7.7	2.0	1.7	(0.2)
	Ulcerative colitis Asacol	1.0	1.1	1.1	0.7	0.8	0.7	(0.4)
	Psoriasis vulgaris Dovobet	1.2	1.4	1.2	1.2	1.6	1.4	0.2
	Psoriasis Lumicef	0.1	0.1	0.3	0.4	0.5	0.5	0.2
	Parkinson's disease Nourias	1.8	2.1	2.1	1.9	2.5	2.4	0.3
	Antiepileptic Depakene	1.6	1.7	1.6	1.3	1.4	1.3	(0.3)
	Technology out-licensing	0.8	1.2	0.8	1.1	0.4	0.8	(0.1)
	International	Renal anemia treatment drug Nesp	1.4	1.5	1.5	1.5	1.6	1.6
Secondary hyperparathyroidism Regpara		0.5	0.5	0.6	0.7	0.8	0.9	0.3
Agent for decreasing the incidence of febrile neutropenia Neulasta/Peglasta		0.5	0.5	0.5	0.4	0.6	0.7	0.2
Neutropenia treatment drug Gran		1.0	1.3	1.5	1.4	1.3	1.5	0.0
X-linked hypophosphatemia (XLH) Crysvita		-	-	-	-	0.8	2.7	2.7
Cancer pain Abstral		2.9	2.8	3.1	3.4	3.1	3.1	0.0
Cancer pain PecFent		1.1	0.9	1.1	1.0	1.0	1.2	0.1
Anticancer Mitomycin-C		0.8	0.8	0.9	0.6	0.6	0.5	(0.4)
Chemotherapy-induced nausea and vomiting drug Sancuso		0.6	0.7	0.7	0.6	0.7	0.8	0.1
Opioid-induced constipation (OIC) Moventig		0.2	0.1	0.2	0.3	0.3	0.4	0.2
Replacement therapy with testosterone for male hypogonadism Tostran/Fortesta		0.4	0.6	0.6	0.6	0.8	0.7	0.1
Osteoporosis drug Adcal-D3		0.8	0.9	0.9	0.9	0.9	0.9	(0.1)
Technology out-licensing		8.0	1.6	1.3	6.5	5.9	1.5	0.2

* Revenue is classified as Japan or International (other than Japan) based on customer location.






* Revenue listed as "Technology out-licensing" specifies revenue from the upfront payment, milestone revenue, and running royalties revenue that are obtained based on licensing agreements recognizing the granting to third parties the rights for development, manufacturing and sales of the Group's pipeline compounds or the use of technology, etc.

V. R&D Pipeline








Filed • Approved



 antibody  protein  small molecule

As of Sep. 30, 2018

Area		Code Name Generic Name Formulation	Mechanism of Action	Indication	Stage	In-House or Licensed	Remarks
Nephrology		KRN321 Darbepoetin Alfa Injection	Long-Acting Erythropoiesis Stimulating Agent	Renal Anemia (on Dialysis)	NDA in preparation in CN	Kirin-Amgen	
				Renal Anemia	Filed in ID		
Oncology		KW-0761 Mogamulizumab Injection	Anti-CCR4 Humanized Antibody	Mycosis Fungoides and Sézary Syndrome	Approved in US	In-House	POTELLIGENT® Additional Indication, Dosage and Administration
				Cutaneous T-cell Lymphoma	Filed in EU Approved in JP		
Immunology/A llergy		KHK4827 Brodalumab Injection	Anti-IL-17 Receptor A Fully Human Antibody	Psoriasis	Filed in KR	Kirin-Amgen	
					Approved in TH		
					Filed in SG		
					Filed in MY Filed in HK		
Central Nervous System		KW-6002 Istradefylline Oral	Adenosine A _{2A} Receptor Antagonist	Parkinson's Disease	NDA in preparation in US	In-House	
Other		© KRN23 Burosumab Injection	Anti-FGF23 Fully Human Antibody	X-linked Hypophosphatemia (XLH)	Filed in CA	In-House	Human Antibody-Producing Technology Jointly Developed with Ultragenyx in US and EU




Phase II, Phase III

Area		Code Name Generic Name Formulation	Mechanism of Action	Indication	Stage	In-House or Licensed	Remarks	
Nephrology		© RTA 402 Bardoxolone Methyl Oral	Antioxidant Inflammation Modulator	Diabetic Kidney Disease	Phase III in JP	Licensed from Reata		
					Phase III in JP			
Oncology		KHK7580 Evocalcet Oral	Calcium Receptor Agonist	Hypercalcemia In Patients With Parathyroid Carcinoma or Primary Hyperparathyroidism	Phase III in JP	Licensed from Mitsubishi Tanabe Pharma		
					Phase II in JP			
Immunology/A llergy		© KHK4083 Injection	Anti-CCR4 Humanized Antibody	Adult T-cell Leukemia/Lymphoma	Phase II in US, EU and others	In-House	POTELLIGENT® Human Antibody-Producing Technology	
					Phase II in US, EU and others			
					Phase III in JP			
					Phase II in JP			
Immunology/A llergy		KHK4563 Benralizumab Injection	Anti-IL-5 Receptor Humanized Antibody	Chronic Obstructive Pulmonary Disease (COPD)	Phase III in JP	In-House	POTELLIGENT® Jointly Developed with AstraZeneca/MedImmune	
				Eosinophilic Chronic Rhinosinusitis (ECRS)	Phase II in JP			
				Anti-IL-17 Receptor A Fully Human Antibody	Phase III in JP, KR and TW			Kirin-Amgen
				Anti-CD40 Fully Human Antibody	Phase II in US			
Central Nervous System		KW-0761 Mogamulizumab Injection	Anti-CCR4 Humanized Antibody	HTLV-1 associated myelopathy (HAM)	Phase III in JP	In-House	POTELLIGENT®	
					© KW-6356 Oral			Adenosine A _{2A} Receptor Antagonist
Other		© KRN23 Burosumab Injection	Anti-FGF23 Fully Human Antibody	X-linked Hypophosphatemia (XLH) in adult patients	Phase III in US, CA, EU, JP and KR	In-House	Human Antibody-Producing Technology Jointly Developed with Ultragenyx in US and EU	
				X-linked Hypophosphatemia (XLH) in pediatric patients	Phase III in US, CA, EU, AU, JP and KR			
				Tumor Induced Osteomalacia (TIO)/Epidermal Nevus Syndrome (ENS)	Phase II in US Phase II in JP and KR			
				Aplastic Anemia	Phase II/III in KR			Kirin-Amgen
Idiopathic (Immune) Thrombocytopenic Purpura	Phase III in CN							








 Updated since Jun. 30, 2018 (Area, Stage, Filed, Approved, etc.)
 New Molecular Entity



V. R&D Pipeline

Ph I

 antibody  protein  small molecule




As of Sep. 30, 2018

Area		Code Name Generic Name Formulation	Mechanism of Action	Indication	Stage	In-House or Licensed	Remarks
Oncology		ⓈKHK2455 Oral	IDO 1 Inhibitor	Solid Tumor	Phase I in US	In-House	Combination with KW-0761
		ⓈKHK2823 Injection	Anti-CD123 Fully Human Antibody	Cancer	Phase I in UK	In-House	POTELLIGENT® Human Antibody-Producing Technology
		KW-0761 Mogamulizumab Injection	Anti-CCR4 Humanized Antibody	Solid Tumor	Phase I / II in US	In-House	POTELLIGENT® Combination with Nivolumab (Jointly Developed with Bristol-Myers Squibb)
Immunology/ Allergy		ⓈKHK4083 Injection	Anti-OX40 Fully Human Antibody	Ulcerative Colitis Atopic Dermatitis	Phase I in JP Phase I in JP	In-House	POTELLIGENT® Human Antibody-Producing Technology
		KHK4827 Brodalumab Injection	Anti-IL-17 Receptor A Fully Human Antibody	Autoimmune Disease	Phase I in JP	Kirin-Amgen	
Central Nervous System		ⓈKHK6640 Injection	Anti-Amyloid Beta Peptide Antibody	Alzheimer's Disease	Phase I in EU Phase I in JP	Licensed from Immunas Pharma	
Others		KW-3357 Antithrombin Gamma Injection	Recombinant Human Antithrombin	Disseminated Intravascular Coagulation, Congenital Antithrombin Deficiency	Phase I in EU	In-House	


 Updated since Jun. 30, 2018 (Area, Stage, Filed, Approved, etc.)
 New Molecular Entity

Updated since Jun. 30, 2018 (Area, Stage, Filed, Approved, etc.)

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Area		Code Name Generic Name Formulation	Mechanism of Action	Indication	Stage	In-House or Licensed	Remarks
Oncology		KW-0761 Mogamulizumab Injection	Anti-CCR4 Humanized Antibody	Mycosis Fungoides and Sézary Syndrome	Approved in US	In-House	POTELLIGENT® Additional Indication, Dosage and Administration
				Cutaneous T-cell Lymphoma	Approved in JP		
Immunology/A llergy		KHK4827 Brodalumab Injection	Anti-IL-17 Receptor A Fully Human Antibody	Psoriasis	Filed in KR	Kirin-Amgen	
					Approved in TH		
Other		AMG531 Romiplostim Injection	Thrombopoietin Receptor Agonist	Aplastic Anemia	Filed in JP	Kirin-Amgen	

Terminated

Area		Code Name Generic Name Formulation	Mechanism of Action	Indication	Stage	In-House or Licensed	Remarks
Oncology		KW-0761 Mogamulizumab Injection	Anti-CCR4 Humanized Antibody	Solid Tumor	Phase I in JP	In-House	POTELLIGENT® Combination with Nivolumab (Jointly Developed with Ono Pharmaceutical)

<Appendix> Fujifilm Kyowa Kirin Biologics co. Ltd., Pipeline of Biosimilar

As of September 30th 2018

Code Name	Generic Name	Stage	Remarks
FKB327	Adalimumab (fully human anti-TNF- α monoclonal antibody)	Approved in EU	Fujifilm Kyowa Kirin Biologics Co., Ltd.
FKB238	Bevacizumab (humanized anti-VEGF monoclonal antibody)	Phase III in US, Europe and others	Centus Biotherapeutics Ltd.(*)

* Centus Biotherapeutics Ltd. is a fifty-fifty joint venture between Fujifilm Kyowa Kirin Biologics Co., Ltd. and AstraZeneca.