

# Kyowa Kirin Co., Ltd.

# Appendix to the Consolidated Financial Summary (IFRS) Fiscal 2019 Third Quarter

(January 1, 2019 - September 30, 2019)

<sup>-</sup> These materials were made as a supplement to the Kessan Tanshin (Consolidated Financial Summary, IFRS), disclosed at the Tokyo Stock Exchange on October 29, 2019 for the first nine months of Fiscal 2019, from January 1, 2019 to September 30, 2019.

<sup>-</sup> This document is an English translation 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.

<sup>&</sup>quot;FY 2019 forecasts" have not been revised from the consolidated earnings forecasts released on February 5, 2019.

<sup>-</sup> Following the conclusion of an agreement to transfer shares of Kyowa Hakko Bio Co., Ltd., the Bio-Chemicals business is categorized as a discontinued operation, effective the first quarter of Fiscal 2019 for accounting on a consolidated basis. Accordingly, FY 2018 results presented herein have been restated in order to present them similarly to the FY 2019 results.

Note that the amounts after restatement for the "FY 2018 results Jan - Dec" have not been audited by an audit corporation.

<sup>-</sup> Figures presented in these materials have been rounded to the nearest tenth.

<sup>-</sup> 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 2	2018			FY 2019		
			res	ults			forecasts		
		Jan - Mar	Jan - Jun	Jan - Sep	Jan - Dec	Jan - Mar	Jan - Jun	Jan - Sep	Jan - Dec
	USD	110	109	109	110	110	110	109	110
	EUR	134	132	131	131	126	125	123	130
	GBP	152	151	149	148	143	143	140	145

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31.7

10.5

#### I . Consolidated Financial Results

#### 1. Trends in consolidated profit

Dividend payout ratio (%)

ROE (%)

<Accumulative> Unit: Billions of yen FY 2018 results FY 2019 results FY 2019 forecasts Jan - Mar Jan - Jun Jan - Mar Jan - Dec Jan - Sep Jan - Dec Jan - Jun Jan - Sep Change amount Revenue 66.5 134.3 197 7 271.5 75.8 151 4 225.5 27 7 305.0 Cost of sales (19.8) (37.0 (53.9) (73.4) (19.8) (38.6 (57.1) (3.2)(81.0) 46.7 97.4 143.8 198.1 168.4 24.6 224.0 Gross profit to revenue ratio 70.2% 72.5% 73.0% 74.5% 74.79 73.4% 72.79 74.0% (48.3) (72.3) (102.1) (55.3) (83.9) (11.6) (117.5) Selling, general and administrative expenses (23.5) (26.7) (10.2) (32.3) (45.7) (11.9) (24.9) (37.9) (52.5) Research and development expenses (21.6)(5.6)R&D expenses to revenue ratio 15.3% 16.1% 16.3% 16.8% 15.7% 16.4% 16.8% 17.2% Share of profit (loss) of investments accounted for using equity method 1.4 0.9 0.3 (0.1) (0.2) (0.5) (0.9) (1.2) (1.0) 14.4 32.2 45.8 Core operating profit 28.4 39.5 50.3 17.3 6.2 53.0 Core operation profit to revenue ratio 21.6% 21.19 20.09 18.5% 22.9% 21.29 20.39 17.4% Other income 14.4 14.5 18.3 18.6 0.1 0.2 0.3 (18.0) Other expenses (0.1) (0.3) (1.4) (5.3) (10.5) (12.3) (11.9) Finance income (costs) (0.1) (0.4 (0.6) 0.1 (0.0) 28.6 42.2 66.8 12.3 21.8 47.0 56.9 33.5 (23.4) Income tax expense (17.6) (7.7)(10.7) (14.1) (3.0)(3.2)(6.5)7.5 27.0% 24 1% 14.5% Ratio of income tax burder 25.49 24 7% 26.3% 19.6% Profit from continuing operations 20.9 31.5 42.8 492 93 18 7 26.9 (15.9) 37.0 Profit from continuing operations to revenue ratio 31.4% 23.59 21.6% 18.1% 12.3% 12.3% 11.99 12.1% Profit from discontinued operations 1.1 2.7 4.6 5.2 (1.2) 29.4 29.4 24.8 31.0 22.0 34.3 47.4 54.4 8.1 48.1 56.3 8.9 68.0 Profit to revenue ratio 33.1% 25.5% 20.0% 10.7% 31.7% 22.3% 24.0% 25.09 EPS (¥/share) 40.20 62.61 99.40 14.97 89.02 104.48 126.30 86.62 17.86 Annual dividend (¥/share) 35.00 40.00

35.2

8.6

<Quarterly> Unit: Billions of yen FY 2019 results FY 2018 results Jan - Mar Apr - Jun Jul - Sep Oct - Dec Jan - Mar Apr - Jun Jul - Sep Change amount Revenue 66.5 67.9 63.4 73.8 75.8 75.6 74.0 10.7 Cost of sales (19.8) (17.2) (17.0) (19.4) (19.8) (18.9) (18.5) (1.5) Gross profit 46.7 50.7 46.4 54.4 56.1 56.7 55.6 9.2 (23.5) (24.0) (29.8) (28.6) Selling, general and administrative expenses (24.8)(26.7)(28.6)(4.6)Research and development expenses (10.2) (11.4) (10.6) (13.4) (11.9) (13.0) (13.0) (2.3)Share of profit (loss) of investments accounted for using equity method 1.4 (0.5) (0.6) (0.4) (0.2) (0.3) (0.4) 0.2 Core operating profit 14.4 14.0 11.2 10.8 17.3 14.8 2.4 13.6 14.4 0.1 0.1 3.7 0.3 0.1 0.1 (3.6) (0.1) (1.0) Other expenses (0.2) (0.1) (5.3) (5.2) (1.9) (1.7) Finance income (costs) (0.1) (0.3)(0.1)(0.1) 0.1 (0.2) (0.2)(0.1)Profit before tax 28.6 13.6 14.6 10.0 12.3 9.6 11.6 (3.0) Income tax expense (7.7) (3.0) (3.4) (3.5) (3.0) (0.2) (3.4) (0.0) Profit from continuing operations 20.9 10.6 11.3 6.4 9.3 9.3 8.3 (3.0)Profit from discontinued operations 1.1 0.6 30.6 1.6 1.9 (1.2)(1.9)Profit 22.0 12.3 13.1 7.0 8.1 39.9 (4.9) 8.3

### <Breakdown of profit for FY 2019 Jan - Sep from discontinued operations>

Unit: Billions of	yen

		FY 2018	3 results			FY 2019 results	
	Jan - Mar	Jan - Jun	Jan - Sep	Jan - Dec	Jan - Mar	Jan - Jun	Jan - Sep
Revenue	18.2	37.8	56.2	75.0	18.1	18.1	18.1
Gross profit	7.1	14.6	22.2	29.0	6.8	6.8	6.8
Core operating profit	1.9	3.8	6.4	8.4	1.3	1.3	1.3
Profit before tax	1.7	3.5	5.9	6.6	(0.6)	43.2	43.2
Profit	1.1	2.7	4.6	5.2	(1.2)	29.4	29.4

Regarding transactions between continuing operations and discontinued operations, considering the continuity of the transactions in the future, revenue and the expenses generated from transactions between such operations have been eliminated from the results of discontinued operations.

<sup>\*</sup> Profit from discontinued operations is presented separately from continuing operations on the condensed quarterly consolidated statement of profit or loss.

Accordingly, the amounts displayed from revenue through profit from continuing operations are amounts of the continuing operations and exclude the discontinued operations.



2. Revenue by geographic region (continuing operations)

2. Revenue by geographic region (continuing opera	tions)									Unit: Bill	ions of yen
		F	Y 2018 resu	Its			FY 2019	9 results		FY 2019	forecasts
	Jan - Mar	Jan - Jun	Jan - Sep	Jan - Dec	Percentage of consolidated revenue	Jan - Mar	Jan - Jun	Jan - Sep	Percentage of consolidated revenue	Jan - Dec	Percentage of consolidated revenue
Japan	45.7	91.4	135.1	183.5	67.6%	48.8	95.5	140.6	62.4%	185.5	60.8%
International	20.8	43.0	62.7	88.0	32.4%	27.1	55.9	84.9	37.6%	119.5	39.2%
Americas	4.2	10.9	14.6	23.0	8.5%	8.9	20.4	32.8	14.5%	50.0	16.4%
Europe	11.5	21.5	31.8	42.3	15.6%	11.8	22.0	31.7	14.1%	45.0	14.8%
Asia	4.9	10.5	16.2	22.5	8.3%	6.3	13.4	20.3	9.0%	24.0	7.9%
Others	0.2	0.1	0.1	0.2	0.1%	0.0	0.1	0.1	0.0%	0.5	0.2%
Total consolidated revenue	66.5	134.3	197.7	271.5	100%	75.8	151.4	225.5	100%	305.0	100%

<sup>\*</sup> Revenue is classified by region or country based on location of customer.

3. Capital expenditures and intangible assets investment (continuing operations)

Ur	nit: Billions of yen
	FY 2019 forecasts
Jan - Sep	Jan - Dec
5.7	10.6
	Jan - Sep

FY 2019 resu

Jan - Jun

Jan - Mar

Jan - Jun

Jan - Mar

FY 2018 results

Jan - Sep

Jan - Dec

4. Depreciation and amortization (continuing operation)	ons)						Ur	nit: Billions of yen
	FY 2018 results				FY 2019 results			FY 2019 forecasts
	Jan - Mar	Jan - Jun	Jan - Sep	Jan - Dec	Jan - Mar	Jan - Jun	Jan - Sep	Jan - Dec
Depreciation (property, plant and equipment)	1.6	3.5	5.4	7.3	2.5	5.5	8.1	9.3
Amortization (intangible assets)	2.1	4.2	6.2	8.9	2.2	4.5	6.4	8.6
Total	3.8	7.7	11.7	16.2	4.7	10.0	14.5	17.9

#### II. Consolidated Statement of Cash Flows

II. Consolidated Statement of Cash Flows							Ui	nit: Billions of yen
		FY 2018	results			FY 2019	results	
	Jan - Mar	Jan - Jun	Jan - Sep	Jan - Dec	Jan - Mar	Jan - Jun	Jan - Sep	Change amoun
Cash flows from operating activities	19.5	29.8	46.4	56.2	9.2	19.8	38.6	(7.8
Of which, cash flows of discontinued operations	1.6	2.1	3.5	6.8	6.3	6.3	6.3	2.8
Cash flows from investing activities	(14.5)	(23.0)	(30.8)	(39.9)	29.3	10.6	4.1	35.0
Of which, cash flows of discontinued operations	(2.3)	(4.3)	(5.2)	(6.4)	(1.9)	103.2	103.2	108.4
Cash flows from financing activities	(8.3)	(8.4)	(16.5)	(16.5)	(34.5)	(35.2)	(46.6)	(30.1)
Of which, cash flows of discontinued operations	(0.3)	(0.3)	(0.2)	(0.2)	(0.0)	(0.0)	(0.0)	0.1
Effect of exchange rate changes on cash and cash equivalents	(0.1)	(0.6)	(0.4)	0.4	(0.4)	(0.2)	0.1	0.4
Net increase (decrease) in cash and cash equivalents	(3.5)	(2.2)	(1.3)	0.1	3.7	(4.9)	(3.8)	(2.5)
Transfer to assets held for sale	1.1	1.1	1.1	1.1	(3.6)	-	-	(1.1
Cash and cash equivalents at beginning of period	14.7	14.7	14.7	14.7	15.9	15.9	15.9	1.2
Cash and cash equivalents at end of period*	12.3	13.6	14.5	15.9	15.9	10.9	12.1	(2.4
* Cash reserves at end of period								
Cash and cash equivalents at end of period	12.3	13.6	14.5	15.9	15.9	10.9	12.1	(2.4
+ Loans receivable from parent in excess of three months	168.0	172.4	179.6	181.3	144.6	283.7	283.4	103.8
+ Time deposits whose maturity periods exceed three months	0.0	0.0	0.0	0.0	-	0.8	0.8	0.7
Cash reserves at end of period	180.3	186.0	194.1	197.2	160.5	295.4	296.2	102.1

Capital expenditures (property, plant and equipment) 0.8 3.3 4.5 9.0 Intangible assets investment 0.6 4.4 5.7 3.4 3.7 13.2 11.5 Total 1.4 6.2 9.0 13.5 6.0 8.2 18.9 22.1

<sup>\*</sup> Acquisitions of right-of-use assets are not included.



#### **III.** Revenue from Main Products

<Accumulative> Unit: Billions of yen FY 2019 FY 2018 results FY 2019 results forecasts Indication / Product name Jan - Mai Jan - Jun Jan - Sep Jan - Dec Jan - Mar Jan - Jun Jan - Dec % Renal anemia treatment drug Nesp 12.0 25.6 39.2 53.7 11.8 25.6 32.0 (7.2)Darbepoetin Alfa Injection Syringe [KKF] 5.6 5.6 53.7 12.0 25.6 48.4 Total 39.2 11.8 25.6 37.6 90% (1.6)Secondary hyperparathyroidism 3.6 Regpara 7.8 10.7 13.3 1.8 3.8 5.2 (5.4)5.1 Secondary hyperparathyroidism Orkedia 0.4 1.1 2.4 1.2 3.0 4.8 3.7 9.5 397% Secondary hyperparathyroidism Rocaltrol 0.9 1.9 2.8 3.8 8.0 1.7 2.7 (0.1)3.6 96% Type-2 diabetes 3.6 3.7 Onglyza 1.6 5.4 7.4 1.7 5.5 0.1 7.6 102% Cardiovascular (Hypertension & angina pectoris) Coniel 2.0 1.2 2.5 3.6 4.8 1.0 3.0 (0.7)3.9 81% Agent for decreasing the incidence of febrile neutropenia G-Lasta 4.3 9.5 14.8 20.7 5.3 11.5 18.3 3.5 22.8 110% Transdermal persistent pain 26 (0.5)**Fentos** 1.2 4.0 5.4 1.1 2.3 3.4 4.8 89% Anticancer Poteligeo 0.4 0.9 1.3 1.8 0.4 1.0 1.5 0.2 1.7 96% Anticance Rituximab BS [KHK] 0.3 1.1 2.4 4.3 1.8 4.2 6.8 4.3 8.4 195% Chronic idiopathic thrombocytopenic purpura Romiplate 0.7 1.5 2.4 3.2 8.0 1.6 3.0 0.6 4.4 136% Antiallergenic Allelock 4.6 7.5 9.7 12.6 4.0 6.4 8.5 (1.3)9.3 74% Antiallergic eyedrops Patanol 7.7 9.7 11.5 13.4 8.5 99 11.7 0.2 11.3 84% Ulcerative colitis Asacol 0.7 1.5 2.2 2.9 0.6 1.2 1.8 (0.4)2.2 77% Psoriasis vulgaris Dovobet 2.8 5.9 1.5 5.0 7.8 12 42 3 4 0.8 132% Psoriasis Lumicef 0.5 0.4 0.9 2.0 1.2 1.8 0.5 2.7 134% 1.4 Parkinson's disease Nouriast 1.9 4.4 6.8 9.4 2.2 4.8 7.3 0.5 10.0 107% Antiepileptic Depakene 1.3 2.7 4.0 5.3 2.3 3.3 (0.6) 11 43 82% **Technology out-licensing** 1.5 1.7 3.7 1.5 162% X-linked hypophosphatemia (XLH) Crysvita 8.0 3.2 7.7 5.7 13.4 21.6 18.4 Anticancer Poteligeo 2.1 2.4 5.4 8.0 8.0 10.0 478% Cancer pain 6.5 Abstral 3.4 9.5 12.8 3.1 5.8 8.3 (1.2)12.3 96% Cancer pair Pecfent 1.0 2.0 3.2 1.9 (0.3)5.0 113% 4.4 1.1 2.9 Chemotherapy-induced nausea and vomiting drug Sancuso 0.6 1.3 2.1 3.0 0.7 1.4 2.3 0.2 2.9 99% Opioid-induced constipation (OIC) Moventig 0.6 0.3 1.0 1.4 0.5 1.0 1.4 0.5 2.5 174% Replacement therapy with testosterone for male hypogonadism Tostran/Fortesta 0.6 1.4 2.1 2.8 0.6 1.2 1.6 (0.5)1.9 68% Osteoporosis drug Adcal-D3 0.9 1.8 2.7 3.5 8.0 1.6 2.3 (0.3)3.1 90% Anticancer Mitomycin-C 0.6 1.3 1.7 2.3 0.6 1.0 (0.2)88% 1.5 2.1 Renal anemia treatment drug 1.5 3.1 4.7 6.4 1.7 3.3 4.7 0.0 5.9 93% Nesp Secondary hyperparathyroidism 3.2 1.5 Regpara 0.7 15 2.3 11 24 38 37 114% Agent for decreasing the incidence of febrile neutropenia Neulasta/Peglasta 0.4 1.0 2.4 1.9 173% 1.7 1.0 2.7 1.0 4.2 Neutropenia treatment drug 27 Gran 14 42 5.4 1.5 3 1 48 0.6 5.6 104% 6.5 12.4 13.9 15.8 5.2 7.9 (6.0)12.9 Technology out-licensing 3.2 82% Of which, Benralizumab royalty 0.0 0.5 1.7 3.3 2.0 3.8 6.1 4.4

<sup>\*</sup> Revenue is classified as Japan or International (other than Japan) based on customer location.

<sup>\*</sup> Revenue from main products does not include revenue from the Early Access Program (EAP).

<sup>\*</sup> 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.

<sup>\*</sup> Benralizumab royalty only refers to the royalty on sales of Fasenra by AstraZeneca (including the Company's own estimates).



#### III. Revenue from Main Products

<Quarterly> Unit: Billions of yen FY 2018 results FY 2019 results Indication / Product name Change amount Jan - Mar Apr - Jun Jul - Sep Oct - Dec Jan - Mar Apr - Jun Jul - Sep Renal anemia treatment drug Nesp 12.0 13.6 13.6 14.5 11.8 13.8 6.4 (7.3)Darbepoetin Alfa Injection Syringe [KKF] 5.6 5.6 12.0 13.6 13.6 14.5 11.8 13.8 12.0 Total (1.7)Secondary hyperparathyroidism 3.6 2.9 1.5 Regpara 2.6 1.8 1.9 (1.5)Secondary hyperparathyroidism Orkedia 0.4 8.0 1.3 1.2 1.8 1.8 1.0 Secondary hyperparathyroidism Rocaltrol 0.9 1.0 0.9 1.0 8.0 0.9 0.9 (0.0)Type-2 diabetes Onglyza 1.6 1.9 1.8 2.1 1.7 2.0 1.8 (0.0)Cardiovascular (Hypertension & angina pectoris) Coniel 1.2 1.0 (0.2)1.2 1.3 1.1 0.9 1.1 Agent for decreasing the incidence of febrile neutropenia G-Lasta 4.3 5.1 5.3 5.9 5.3 6.2 6.8 1.5 Transdermal persistent pain 1.2 Fentos 1.2 14 14 14 1.1 1.1 (0.2)Anticancer Poteligeo 0.4 0.5 0.4 0.5 0.4 0.5 0.5 0.1 Anticancer Rituximab BS [KHK] 0.3 8.0 1.3 1.8 1.8 2.3 2.6 1.3 Chronic idiopathic thrombocytopenic purpura Romiplate 0.7 8.0 8.0 0.9 8.0 0.9 1.3 0.5 Antiallergenic Allelock 4.6 2.9 2.3 2.8 4.0 2.4 2.0 (0.2)Antiallergic eyedrops Patanol 7.7 20 17 19 8.5 1.4 1.7 0.0 Ulcerative colitis 0.8 Asacol 0.7 0.7 0.7 0.6 0.6 0.6 (0.1)Psoriasis vulgaris 1.7 1.6 1.5 1.9 1.6 0.2 Dovobet 12 14 Psoriasis Lumicef 0.5 0.6 0.4 0.5 0.5 0.7 0.7 0.1 Parkinson's disease Nouriast 1.9 2.5 2.4 2.6 2.2 2.6 2.5 0.1 Antiepileptic Depakene (0.2)1.3 1.3 1.3 1.2 14 11 1 1 **Technology out-licensing** 0.8 0.5 0.9 0.7 1.3 X-linked hypophosphatemia (XLH) Crysvita 8.0 2.4 4.5 5.7 7.7 8.2 5.9 Anticancer Poteligeo 2.1 2.4 3.0 2.6 2.6 Cancer pain Abstral 3.1 3.4 3.1 3.2 3.1 2.7 2.5 (0.6)Cancer pain Pecfent 1.0 1.0 1.2 1.2 0.9 1.0 (0.2)1.1 Chemotherapy-induced nausea and vomiting drug Sancuso 0.6 0.7 8.0 0.9 0.7 0.7 8.0 0.1 Opioid-induced constipation (OIC) Moventig 0.3 0.3 0.4 0.4 0.5 0.5 0.5 0.1 Replacement therapy with testosterone for male hypogonadism Tostran/Fortesta 0.6 0.8 0.7 0.7 0.6 0.6 0.4 (0.3)Osteoporosis drug Adcal-D3 0.9 0.9 0.9 0.8 8.0 0.8 0.7 (0.1)Anticancer Mitomycin-C 0.6 0.6 0.5 0.6 0.6 0.5 (0.0)0.4 Renal anemia treatment drug 1.7 Nesp 1.5 1.6 1.6 1.7 1.6 1.4 (0.2)Secondary hyperparathyroidism Regpara 0.7 0.8 0.90.91.1 1.3 1.4 0.5 Agent for decreasing the incidence of febrile neutropenia Neulasta/Peglasta 0.4 0.6 0.7 1.0 0.1 0.7 1.0 8.0 Neutropenia treatment drug 12 1.7 1.7 0.2 Gran 1.4 1.3 15 1.5 **Technology out-licensing** 6.5 5.9 1.5 1.9 3.2 2.0 27 1.2 Of which, Benralizumab royalty 0.0 0.5 1.2 1.6 2.0 1.8 2.3 1.1

<sup>\*</sup> Revenue is classified as Japan or International (other than Japan) based on customer location

<sup>\*</sup> Revenue from main products does not include revenue from the Early Access Program (EAP).

<sup>\*</sup> 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.

<sup>\*</sup> Benralizumab royalty only refers to the royalty on sales of Fasenra by AstraZeneca (including the Company's own estimates).



#### IV. R&D Pipeline

\*\* antibody Filed • Approved As of Sep. 30, 2019 Code Name In-House Generic Name Formulation Area Mechanism of Action Indication Stage Remarks Renal Anemia (on Dialysis) Filed in CN KRN321 Long-Acting Erythropoiesis Stimulating Agent S) Darbepoetin Alfa Injection Kirin-Amgen Renal Anemia Approved in ID Nephrology Hypercalcemia In Patients With Parathyroid Carcinoma or Primary KHK7580 Evocalcet Oral Licensed from Mitsubishi Tanabe Pharma X Calcimimetic Filed in JP Hyperparathyroidism Filed in KR KHK4827 Brodalumab Injection Filed in MY Immunology /Allergy Anti-IL-17 Receptor A Fully Human Antibody Psoriasis Kirin-Amgen Filed in CN Filed in MO KW-6002 Central Adenosine A<sub>2A</sub> Receptor Antagonist 水 Nervous System Istradefylline Oral Parkinson's Disease Approved in US In-House Filed in TW Filed in CH Filed in KW X-linked Hypophosphatemia (XLH) Filed in SA Human Antibody-Producing Technology Jointly Developed with Ultragenyx in US and EU ©KRN23 Burosumab Injection Anti-FGF23 Fully Human Antibody In-House Other Filed in CN ٧ Filed in HK Filed in SG FGF23-Related Hypophosphatemic Rickets and Osteomalacia Approved in JP Filed in KR

#### Ph Ⅱ~Ⅲ

Ph II∼III Area		Code Name Generic Name	Mechanism of Action	Indication	Stage	In-House or	Remarks
Area		Formulation	Wechanism of Action	indication	Stage	Licensed	Remarks
	άķ	©RTA 402 Bardoxolone Methyl Oral	Antioxidant Inflammation Modulator	Diabetic Kidney Disease	Phase Ⅲ in JP	Licensed from Reata	
Nephrology	*	KHK7580 Evocalcet Oral	Calcimimetic	Secondary Hyperparathyroidism	Phase III in CN, KR, TW and HK	Licensed from Mitsubishi Tanabe Pharma	
	水		NHE3 Inhibitor	Hyperphosphatemia Under Maintenance Dialysis	Phase II in JP	Licensed from Ardelyx	
	**		HDAC Inhibitor	Breast Cancer	Phase II in JP	Licensed from Syndax	
Oncology	<b>Y</b>	KW-0761 Mogamulizumab Injection	Anti-CCR4 Humanized Antibody	Adult T-cell Leukemia/Lymphoma	Phase II in US, EU and others	In-House	POTELLIGENT®
	8	KRN125 Pegfilgrastim Injection	Long-Acting Granulocyte Colony Stimulating Factor	Mobilization of Hematopoietic stem cell into Peripheral blood	Phase II in JP	Kirin-Amgen	
	<b>Y</b>	©KHK4083	Anti-OX40 Fully Human	Ulcerative Colitis	Phase II in US, EU and others	In-House	POTELLIGENT <sup>®</sup>
		Injection	Antibody	Atopic Dermatitis	Phase II in JP, US, CA and EU		Human Antibody-Producing Technology
mmunology/				Axial Spondyloarthritis (axSpA)	Phase III in JP, KR and TW		
Allergy	<b>Y</b>	KHK4827 Brodalumab Injection	Anti-IL-17 Receptor A Fully Human Antibody	Systemic Sclerosis	Phase Ⅲ in JP	Kirin-Amgen	
				Palmoplantar Pustulosis	Phase Ⅲ in JP		
	<b>Y</b>		Anti-CD40 Fully Human Antibody	Recurrence of Focal Segmental Glomerulosclerosis (FSGS) in <i>de</i> novo kidney transplant recipients	Phase II in US	In-House	Human Antibody-Producing Technology Jointly Developed with Astellas
Central Nervous	*	KW-0761 Mogamulizumab Injection	Anti-CCR4 Humanized Antibody	HTLV-1 associated myelopathy (HAM)	Phase Ⅲ in JP	In-House	POTELLIGENT <sup>®</sup>
System	xř.	⊚KW-6356 Oral	Adenosine A <sub>2A</sub> Receptor Antagonist	Parkinson's Disease	Phase II in JP	In-House	
	<b>Y</b>	©KRN23 Burosumab Injection	Anti-FGF23 Fully Human Antibody	Tumor Induced Osteomalacia(TIO)/Epidermal Nevus Syndrome (ENS)	Phase II in US  Phase II in JP and KR	In-House	Human Antibody-Producing Technology Jointly Developed with Ultragenyx in US and EU
Other		AMG531		Aplastic Anemia Who Have Had an Inadequate Response to Conventional Therapy	Phase II/III in KR		
	8	Romiplostim Injection	Thrombopoietin Receptor Agonist	Aplastic Anemia Who Were Previously Untreated with Immunosuppressive Therapy	Phase Ⅱ/Ⅲ in JP, KR and TW	Kirin-Amgen	
				Idiopathic (Immune) Thrombocytopenic Purpura	Phase III in CN		

Updated since Jun. 30, 2019 (Area, Stage, Filed, Approved, etc.)

© New Molecular Entity



## Ⅳ. R&D Pipeline

		Cada Nama	☆ antibody	Protein  sm	all molecule	In H	As of Sep. 30, 201
Area		Code Name Generic Name Formulation	Mechanism of Action	Indication	Stage	In-House or Licensed	Remarks
	×ř	⊚KHK2455	IDO1 Inhibitor	Solid Tumor	Phase I in US	In-House	Combination with KW-0761
Oncology	ताः	Oral	IDOT Inhibitor	Urothelial carcinoma	Phase I in US	In-House	Combination with avelumab
Oricology	*	⊚ME-401 Oral	PI3Kδ Inhibitor	B-cell malignancies	Phase I in JP	Licensed from MEI Pharma	
mmunology/	Y	©KHK4083 Injection	Anti-OX40 Fully Human Antibody	Ulcerative Colitis	Phase I in JP	In-House	POTELLIGENT® Human Antibody-Producing Technology
Allergy	Y	KHK4323 Injection		Atopic Dermatitis	Phase I in JP	In-House	
Central		©KHK6640	Anti-Amyloid Beta		Phase I in EU	Licensed from	
Nervous System	T	Injection	Peptide Antibody	Alzheimer's Disease	Phase I in JP	Immunas Pharma	
Others	8	KW-3357 Antithrombin Gamma Injection	Recombinant Human Antithrombin	Disseminated Intravascular Coagulation, Congenital Antithrombin Deficiency	Phase I in EU	In-House	
Filed ∙ App	rovec	I	New Molecular En	30, 2019 (Area, Stage, ity age, Filed, Approved, e			
Area		Code Name Generic Name	Mechanism of Action	Indication	Stage	In-House or	Remarks
		Formulation				Licensed	
Nephrology	8	KRN321 Darbepoetin Alfa	Long-Acting Erythropoiesis	Renal Anemia	Approved in ID	Kirin-Amgen	
	3	Injection	Stimulating Agent			9	
	<b>Y</b>	Injection KHK4827 Brodalumab Injection	Stimulating Agent  Anti-IL-17 Receptor A Fully Human Antibody	Psoriasis	Filed in MO	Kirin-Amgen	
mmunology	<b>Y</b>	KHK4827 Brodalumab	Anti-IL-17 Receptor A	Psoriasis Parkinson's Disease	Filed in MO Approved in US		
Immunology /Allergy Central Nervous System	<b>Y</b>	KHK4827 Brodalumab Injection  KW-6002 Istradefylline Oral	Anti-IL-17 Receptor A Fully Human Antibody  Adenosine A <sub>2A</sub> Receptor Antagonist  Anti-FGF23 Fully			Kirin-Amgen In-House	
mmunology /Allergy Central Nervous	<b>Y</b>	KHK4827 Brodalumab Injection KW-6002 Istradefylline Oral	Anti-IL-17 Receptor A Fully Human Antibody  Adenosine A <sub>2A</sub> Receptor Antagonist	Parkinson's Disease FGF23-Related Hypophosphatemic Rickets	Approved in US	Kirin-Amgen	Human Antibody-Producing Technolog Jointly Developed with Ultragenyx in US and EU
mmunology /Allergy Central Nervous System	<b>Y</b>	KHK4827 Brodalumab Injection  KW-6002 Istradefylline Oral	Anti-IL-17 Receptor A Fully Human Antibody  Adenosine A <sub>2A</sub> Receptor Antagonist  Anti-FGF23 Fully	Parkinson's Disease  FGF23-Related Hypophosphatemic Rickets and Osteomalacia  X-linked	Approved in US  Approved in JP	Kirin-Amgen In-House	Jointly Developed with Ultragenyx in US
Immunology /Allergy Central Nervous System	<b>Y</b>	KHK4827 Brodalumab Injection  KW-6002 Istradefylline Oral	Anti-IL-17 Receptor A Fully Human Antibody  Adenosine A <sub>2A</sub> Receptor Antagonist  Anti-FGF23 Fully	Parkinson's Disease  FGF23-Related Hypophosphatemic Rickets and Osteomalacia  X-linked	Approved in US  Approved in JP	Kirin-Amgen In-House	Jointly Developed with Ultragenyx in US
mmunology /Allergy  Central Nervous System  Others  Ph II~III  Area	<b>Y</b>	KHK4827 Brodalumab Injection  KW-6002 Istradefylline Oral	Anti-IL-17 Receptor A Fully Human Antibody  Adenosine A <sub>2A</sub> Receptor Antagonist  Anti-FGF23 Fully Human Antibody	Parkinson's Disease  FGF23-Related Hypophosphatemic Rickets and Osteomalacia  X-linked Hypophosphatemia (XLH)	Approved in US  Approved in JP  Filed in SG	Kirin-Amgen In-House In-House In-House or	Jointly Developed with Ultragenyx in US and EU
mmunology /Allergy  Central Nervous System  Others  Area  mmunology /Allergy	Y ** Y	KHK4827 Brodalumab Injection  KW-6002 Istradefylline Oral  ©KRN23 Burosumab Injection  Code Name Generic Name Formulation  KHK4827 Brodalumab	Anti-IL-17 Receptor A Fully Human Antibody  Adenosine A <sub>2A</sub> Receptor Antagonist  Anti-FGF23 Fully Human Antibody  Mechanism of Action  Anti-IL-17 Receptor A	Parkinson's Disease  FGF23-Related Hypophosphatemic Rickets and Osteomalacia  X-linked Hypophosphatemia (XLH)  Indication	Approved in US  Approved in JP  Filed in SG  Stage	Kirin-Amgen  In-House  In-House  or Licensed	Jointly Developed with Ultragenyx in Usand EU
Central Nervous System  Others  Area  Immunology /Allergy	Y ** Y	KHK4827 Brodalumab Injection  KW-6002 Istradefylline Oral  ©KRN23 Burosumab Injection  Code Name Generic Name Formulation  KHK4827 Brodalumab	Anti-IL-17 Receptor A Fully Human Antibody  Adenosine A <sub>2A</sub> Receptor Antagonist  Anti-FGF23 Fully Human Antibody  Mechanism of Action  Anti-IL-17 Receptor A	Parkinson's Disease  FGF23-Related Hypophosphatemic Rickets and Osteomalacia  X-linked Hypophosphatemia (XLH)  Indication	Approved in US  Approved in JP  Filed in SG  Stage	Kirin-Amgen  In-House  In-House  or Licensed	Jointly Developed with Ultragenyx in US and EU
mmunology /Allergy  Central Nervous System  Others  Ph II~III  Area  mmunology /Allergy  Ph I	Y ** Y	KHK4827 Brodalumab Injection  KW-6002 Istradefylline Oral	Anti-IL-17 Receptor A Fully Human Antibody  Adenosine A <sub>2A</sub> Receptor Antagonist  Anti-FGF23 Fully Human Antibody  Mechanism of Action  Anti-IL-17 Receptor A Fully Human Antibody	Parkinson's Disease  FGF23-Related Hypophosphatemic Rickets and Osteomalacia  X-linked Hypophosphatemia (XLH)  Indication  Palmoplantar Pustulosis	Approved in US  Approved in JP  Filed in SG  Stage  Phase III in JP	In-House	Jointly Developed with Ultragenyx in Us and EU  Remarks
mmunology /Allergy  Central Nervous System  Others  Others  Area  mmunology /Allergy  Ph I Area  Oncology	Y ** Y	KHK4827 Brodalumab Injection  KW-6002 Istradefylline Oral	Anti-IL-17 Receptor A Fully Human Antibody  Adenosine A <sub>2A</sub> Receptor Antagonist  Anti-FGF23 Fully Human Antibody  Mechanism of Action  Anti-IL-17 Receptor A Fully Human Antibody	Parkinson's Disease  FGF23-Related Hypophosphatemic Rickets and Osteomalacia  X-linked Hypophosphatemia (XLH)  Indication  Palmoplantar Pustulosis	Approved in US  Approved in JP  Filed in SG  Stage  Phase II in JP	In-House In-House In-House or Licensed In-House or Licensed	Jointly Developed with Ultragenyx in Usand EU  Remarks  Remarks
Immunology /Allergy  Central Nervous System  Others  Ph II ~ III  Area  Immunology /Allergy  Ph II  Area	Y ** Y	KHK4827 Brodalumab Injection  KW-6002 Istradefylline Oral	Anti-IL-17 Receptor A Fully Human Antibody  Adenosine A <sub>2A</sub> Receptor Antagonist  Anti-FGF23 Fully Human Antibody  Mechanism of Action  Anti-IL-17 Receptor A Fully Human Antibody	Parkinson's Disease  FGF23-Related Hypophosphatemic Rickets and Osteomalacia  X-linked Hypophosphatemia (XLH)  Indication  Palmoplantar Pustulosis	Approved in US  Approved in JP  Filed in SG  Stage  Phase II in JP	In-House In-House In-House or Licensed In-House or Licensed	Jointly Developed with Ultragenyx in Us and EU  Remarks  Remarks