



Securities Code: 4523

2013.12 Reference Data

Third Quarter Ended December 31, 2013

February 3, 2014

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Forward-Looking Statements and Risk Factors

Materials and information provided in this financial disclosure may contain "forward-looking statements" based on current expectations, forecasts, estimates, business goals and assumptions that are subject to risks and uncertainties, which could cause actual outcomes and results to differ materially from these statements. Risks and uncertainties include general industry and market conditions, and general domestic and international economic conditions such as interest rate and currency exchange fluctuations.

Risks that may cause significant fluctuations in the consolidated results of the Eisai Group or have a material effect on investment decisions are described below. These are risk factors that have been identified and assessed as of the disclosure date of the Financial Report.

Risk factors associated with our business include, but are not limited to, challenges arising in overseas operations, uncertainties in new drug development, as well as risks related to dependency on specific products, strategic alliances with partner companies, medical cost-containment measures, generic drug products, intellectual property, possible occurrence of side effects, laws and regulations, litigation, closure or shutdown of production plants, safety and quality of raw materials, outsourcing, environmental issues, IT security and information management, financial market conditions and currency movement, internal control systems, and disasters.

Contents

1.	Consolidated Financial Highlights	 1
2.	Consolidated Statement of Income	 3
3.	Consolidated Statement of Cash Flows	 5
4.	Financial Results by Reporting Segment	 6
5.	Sales Forecasts by Reporting Segment	 12
6.	Consolidated Balance Sheet	 13
7.	Changes in Consolidated Quarterly Results	 15
8.	Nonconsolidated Financial Highlights	 19
9.	Major News Releases	 20
10.	Major R&D Pipeline	 22

* Revisions have been made to the full-year consolidated forecast announced previously. The revised parts are underlined.

* All amounts are rounded to the nearest specified unit.

* The exchange rates used in the reference data are noted in the table below.

* All overseas profit and loss amounts have been converted into yen based on the average exchange rates for the periods shown in the table below.

Currency Exchange Rates

	US	Europe	UK	China
	(JPY/USD)	(JPY/EUR)	(JPY/GBP)	(JPY/RMB)
(Apr. 2012—Dec. 2012) Nine Months Average Rate	79.99	102.17	127.12	12.70
(Dec. 31, 2012) Third Quarter End Rate	86.58	114.71	139.52	13.91
(Apr. 2012—Mar. 2013) Fiscal Year Average Rate	83.10	107.14	131.13	13.25
(Mar. 31, 2013) Fiscal Year End Rate	94.05	120.73	143.16	15.16
(Apr. 2013—Dec. 2013) Nine Months Average Rate	99.38	132.22	155.88	16.24
(Dec. 31, 2013) Third Quarter End Rate	105.39	145.05	173.76	17.36
Fiscal Year Ending March 31, 2014 Fourth Quarter Forecast Rate	<u>102.00</u>	<u>139.00</u>	<u>167.00</u>	<u>16.80</u>

About Indicators in This Reference Data

The Eisai Group believes that cash-generating ability is the most intrinsic element determining the true value of a company. Based upon this belief, in order to reflect our true earnings capacity, we focus on disclosing "cash income" and "cash EPS," adjusted in consideration of non-cash profit-and-loss items, such as depreciation of property, plant and equipment (PP&E) and intangible assets, amortization of goodwill produced by the acquisition of companies, loss on impairment of noncurrent assets (including loss on devaluation of investment securities), and in-process R&D expenses.

*Cash income

Cash income is the total amount of cash available for investment in future growth, return to shareholders, repayment of borrowings, and other expenditures. We consider cash income as an indicator to assess corporate growth potential and strategies.

Cash income = Net income + Depreciation of PP&E and amortization of intangible assets + In-process R&D expenses + Amortization of goodwill + Loss on impairment of noncurrent assets (including loss on devaluation of investment securities)

*Cash income per share (Cash EPS)

Cash EPS = Cash income / Average number of outstanding shares for the period (after deduction of treasury stock)

Segment Information

The Eisai Group classifies its reporting segments into two business categories, namely, Pharmaceutical business and Other business. Furthermore, effective from the fiscal year ending March 31, 2014, the Group has defined the following segments as new reporting segments for its Pharmaceutical business: Japan (Prescription drugs, Generic drugs and Diagnostics), Americas (North, Central and South America), Asia (mainly China, South Korea, Taiwan, India and ASEAN), EMEA (Europe, the Middle East, Africa and Oceania), and Consumer Healthcare Business—Japan (mainly OTC drugs). In line with this change, figures listed in this report for each segment for the fiscal year ended March 31, 2013 are based on the new reporting segments.

1. Consolidated Financial Highlights

I) IIICOIIIE Statement Data	1	Income	Statement Data	
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1) Income Statement Data					(billi	ons of yen)	
	Nine mo	onths ended	I Dec. 31		Full Year		
	FY2012	FY2013	YOY	FY2012	FY2013 forecast		
			%		Revised forecast	Previous forecast	
Net sales	431.6	448.0	103.8	573.7	<u>587.0</u>	578.0	
Cost of sales	128.1	141.6	110.5	174.1	<u>185.0</u>	180.5	
R&D expenses	87.2	106.2	121.7	120.4	<u>130.0</u>	127.5	
SG&A expenses	162.0	157.5	97.2	208.7	<u>197.5</u>	191.5	
Operating income	54.1	42.8	79.1	70.5	<u>74.5</u>	78.5	
Ordinary income	50.2	39.2	78.1	65.6	<u>70.0</u>	74.9	
Net income	34.0	29.5	86.7	48.3	<u>38.5</u>	53.2	
Cash income	72.9	66.3	91.0	100.7	<u>89.5</u>	100.0	
Cash income (adjusted)	-	-	-	-	100.0	-	
Comprehensive income	51.1	77.4	151.4	95.2	-	-	
			Diff.				
Dividend per share (DPS, yen)	-	-	-	150.0	150.0	150.0	
Earnings per share (EPS, yen)	119.3	103.4*	(15.9)	169.4	<u>135.0*</u>	186.6*	
Cash income per share (Cash EPS, yen)	255.8	232.5*	(23.2)	353.5	<u>313.9*</u>	350.8*	
Cash income (adjusted) per share (Cash EPS, yen)	-	-	-	-	350.7*	-	

* The Company's stock held through the Trust for Officers' Compensation Board Incentive Plan (105,400 shares) is included in the average number

of shares outstanding as treasury stocks which are deducted from the basis of the calculation of basic earnings per share. * "Cash income (adjusted)" is the amount of cash income excluding the impact of expenses related to transformation of the R&D organization

and tax rate changes accompanying the abolishment of the special reconstruction corporate tax in Japan a year ahead of schedule.

* "Cost of sales" includes "Provision for (reversal of) sales returns-net."

2) Cash Flow Statement Data

billions (billions)						
	Nine n	Nine months ended Dec. 31				
	FY2012	FY2013	Diff.	FY2012		
Net cash provided by (used in) operating activities	53.5	58.9	5.4	73.2		
Net cash provided by (used in) investing activities	20.9	17.1	(3.9)	21.7		
Net cash provided by (used in) financing activities	(60.8)	(95.6)	(34.8)	(81.8)		
Cash and cash equivalents at end of period	133.3	140.9	7.6	142.5		
Free cash flow	38.9	43.6	4.7	54.5		

* "Free cash flow" = "Net cash provided by (used in) operating activities" - "Capital expenditures, etc. (cash basis)"

3) Balance Sheet Data

	<u>201</u>	1 <u>3</u>	
	March 31	Dec. 31	Diff.
Total assets	990.2	978.8	(11.5)
Liabilities	515.9	470.5	(45.4)
Borrowings	234.0	227.4	(6.7)
Commercial paper	-	10.0	10.0
Bonds and debentures	80.0	30.0	(50.0)
Equity	474.3	508.2	33.9
Shareholders' equity	469.4	504.1	34.7
Shareholders' equity ratio (%)	47.4	51.5	4.1
Liabilities ratio (Net DER / times)	0.27	0.21	(0.06)

* "Liabilities ratio (Net DER)"=("Interest-bearing debt" ("Borrowings" + "Bonds and debentures") - "Cash and deposits"-

"Short-term investments") / "Shareholders' equity"

(hillions of yen)

(billions of yen)

4) Capital Expenditures and Depreciation/Amortization			(billio	ns of yen)	
	Nine mo	Nine months ended Dec. 31			
	FY2012	FY2013	Diff.	FY2012	
Capital expenditures	15.0	16.8	1.8	20.5	
Property, plant and equipment	5.1	5.1	(0.0)	9.2	
Intangible assets	9.9	11.7	1.9	11.3	
Depreciation and amortization	31.7	29.3	(2.4)	43.3	

* "Depreciation and amortization" includes amortization of "Intangible assets."

5) Financial Results by Business Segment (1) Consolidated Net Sales by Reporting Segment

(1) Consolidated Net Sales by Reporting Segment			(billio	ns of yen)	
	Nine mo	Nine months ended Dec. 31			
	FY2012	FY2013	YOY	FY2012	
			%		
Japan Pharmaceutical Business	234.2	240.3	102.6	307.8	
Americas Pharmaceutical Business	114.5	106.8	93.3	153.3	
U.S. Pharmaceutical Business	114.2	106.3	93.1	153.0	
Asia Pharmaceutical Business	29.7	43.3	146.0	41.3	
EMEA Pharmaceutical Business	18.9	24.3	128.8	25.8	
Consumer Healthcare Business—Japan (mainly OTC drugs)	15.8	16.0	101.1	21.1	
Other	18.5	17.2	93.1	24.4	
Consolidated net sales	431.6	448.0	103.8	573.7	

* Each of the segments above displays net sales to external customers only.

(2) Consolidated Operating Income by Reporting Segment		(billion	is of yen)
	Nine mo	onths ended	Dec. 31
	FY2012	FY2013	YOY
			%
Japan Pharmaceutical Business	102.9	122.4	119.0
Americas Pharmaceutical Business	25.7	9.3	36.4
Asia Pharmaceutical Business	6.1	10.3	168.3
EMEA Pharmaceutical Business	1.3	3.5	271.8
Consumer Healthcare Business—Japan (mainly OTC drugs)	2.9	2.9	99.1
Other	9.2	8.0	87.1
R&D expenses	87.2	106.2	121.7
Non-allocated SG&A expenses	6.7	7.4	110.8
Consolidated operating income	54.1	42.8	79.1

2. Consolidated Statement of Income

								s of yen)
			months e				Full year	
	FY2012		FY2013	Sales	YOY	Diff.	FY2012	Sales
Net sales	431.6	% 100.0	448.0	% 100.0	% 103.8	16.5	573.7	% 100.0
Cost of sales	128.1	29.7	141.6	31.6	110.5	13.4	174.1	30.3
Gross profit	303.4	70.3	306.5	68.4	101.0	3.1	399.6	69.7
R&D expenses	87.2	20.2	106.2	23.7	121.7	18.9	120.4	21.0
SG&A expenses	162.0	37.5	157.5	35.1	97.2	(4.6)	-	36.4
Personnel expenses	50.8	11.8	54.6	12.2	107.3	3.7	68.4	11.9
Selling expenses	76.9	17.8	61.8	13.8	80.4	(15.1)		16.2
Administrative and other expenses	34.3	7.9	41.1	9.2	119.9	6.8	47.7	8.3
Operating income	54.1	12.5	42.8	9.6	79.1	(11.3)		12.3
Nonoperating income	1.7	0.4	1.5	0.3		(0.1)	2.3	0.4
Nonoperating expenses	5.6	1.3	5.1	1.1		(0.4)	7.2	1.2
Ordinary income	50.2	11.6	39.2	8.8	78.1	(11.0)	65.6	11.4
Special gain	2.9	0.7	7.2	1.6		4.3	7.5	1.3
Special loss	1.6	0.4	1.8	0.4		0.2	1.7	0.3
Income before income taxes and minority interests	51.5	11.9	44.6	10.0	86.6	(6.9)	71.4	12.5
Income taxes—current	23.0	5.3	21.2	4.7		(1.8)	30.6	5.3
Income taxes—deferred	(5.7)	(1.3)	(6.3)	(1.4)		(0.5)	(7.7)	(1.3)
Income before minority interests	34.2	7.9	29.7	6.6		(4.5)	48.5	8.5
Minority interests in income	0.2	0.1	0.2	0.0		(0.0)	0.3	0.0
Net income	34.0	7.9	29.5	6.6	86.7	(4.5)	48.3	8.4
* "Cost of sales" includes "Provision for (reversal of) sales returns-net."	1							
Cash income								
Net income	34.0	7.9	29.5	6.6	86.7	(4.5)	48.3	8.4
Depreciation of PP&E and amortization of intangible assets	18.5		18.7			0.2	24.9	
Amortization of intangible assets obtained through acquisition	13.2		10.6			(2.6)	18.3	
Amortization of goodwill	5.7		7.0			1.4	7.8	
Loss on impairment of noncurrent assets (including loss on devaluation of investment securities)	1.5		0.5			(1.0)	1.4	

Cash income

* PP&E: propterty, plant and equipment

Notes	
	Primarily increase in net sales of growth drivers such as Humira, Halaven and Lyrica
Net sales <reason for="" increase=""></reason>	Growth in the Asia Pharmaceutical Business (mainly in China), and in Generic Drugs within the Japan Pharmaceutical Business.
Cost of sales to net sales <reason for="" increase=""></reason>	Change in product mix due to decrease in net sales of Aricept and Aciphex/Pariet
R&D expenses <reason for="" increase=""></reason>	Primarily milestone payments associated with progress made in collaborative R&D themes and lump-sum payment related to acquisition of global development and marketing rights for antiobesity agent lorcaserin (brand name in the U.S.: Belviq)
SG&A expenses <reason decrease="" for=""></reason>	Decrease in alliance fees paid to promotion partners
Special gain/loss	Primarily sale of property, plant and equipment, sale of investment securities, and special retirement payments related to transformation of R&D organization

72.9

16.9

66.3

14.8

91.0

(6.6)

100.7

17.6

Consolidated Statement of Comprehensive Income				(billio	ons of yen)		
	Nin	e months en	ded Dec. 31				
	FY2012	FY2013	YOY	Diff.	FY2012		
			%				
Income before minority interests	34.2	29.7	86.7	(4.5)	48.5		
Other comprehensive income (loss)	16.9	47.7	282.6	30.8	46.6		
Valuation difference on available-for-sale securities	(0.4)	0.9		1.3	3.1		
Deferred gain (loss) on derivatives under hedge accounting	(0.0)	0.2		0.2	0.1		
Foreign currency translation adjustments	17.3	46.6		29.3	43.4		
Comprehensive income (loss)	51.1	77.4	151.4	26.3	95.2		
(Breakdown)							
Comprehensive income (loss) attributable to shareholders of the parent company	51.0	77.2	151.4	26.2	95.0		
Comprehensive income (loss) attributable to minority interests	0.1	0.2	184.6	0.1	0.2		

3. Consolidated Statement of Cash Flows

		-	s of yen
		ths ended D	
	FY2012	FY2013	Diff
Income before income taxes and minority interests	51.5	44.6	(6.9
Depreciation and amortization / Amortization of goodwill	37.4	36.3	(1.0
Gain on negative goodwill	(2.0)	(0.2)	1.7
(Gain) loss on sales and disposal of noncurrent assets	(0.5)	(2.8)	(2.3
(Gain) loss on sales of securities	(0.1)	(3.4)	(3.2
Decrease (increase) in notes and accounts receivable-trade, trade payables and inventories	8.8	6.6	(2.3
Increase (decrease) in accounts payable—other / Accrued expenses, etc.	(18.8)	(5.6)	13.2
Other	5.4	7.7	2.3
[Sub-total]	81.7	83.1	1.4
Interest received (paid), etc.	(3.6)	(2.6)	0.9
Income taxes paid	(24.7)	(21.6)	3.1
Net cash provided by (used in) operating activities	53.5	58.9	5.4
Capital expenditures (cash basis)	(14.6)	(15.3)	(0.7
Purchases, proceeds from sales and redemptions of investment securities	0.4	5.9	5.4
Proceeds from sales of investments in subsidiaries resulting in change in scope of consolidation	-	0.9	0.9
Proceeds from sales of investment in subsidiaries in the previous fiscal year	6.1	-	(6.1
Net (increase) decrease in time deposits exceeding three months	29.1	25.1	(3.9
Other	(0.1)	0.4	0.5
Net cash provided by (used in) investing activities	20.9	17.1	(3.9
Net increase (decrease) in short-term borrowings	(2.4)	7.6	10.1
Net increase (decrease) in commercial paper	25.0	10.0	(15.0
Repayments of long-term borrowings	(40.0)	(19.9)	20.1
Redemption of corporate bond	-	(50.0)	(50.0
Dividends paid	(42.7)	(42.8)	(0.0
Other—net	(0.6)	(0.6)	0.1
Net cash provided by (used in) financing activities	(60.8)	(95.6)	(34.8
Foreign currency translation adjustments on cash and cash equivalents	7.1	18.1	11.0
Net increase (decrease) in cash and cash equivalents	20.7	(1.6)	(22.3
Cash and cash equivalents at the beginning of period	112.6	142.5	29.9
Cash and cash equivalents at the end of period	133.3	140.9	7.6
Free cash flow	38.9	43.6	4.7

* "Free cash flow" = "Net cash provided by (used in) operating activities" - "Capital expenditures, etc. (cash basis)"

Notes Net cash provided by (used in) investing activities

Increase from reversal of time deposits exceeding three months as the fund source for redemption of matured bond and debentures

Net cash provided by (used in) financing activities

Decrease from redemption of matured bond and debentures, repayment of long-term borrowings and payment of dividends

1) Japan Pharmaceutical Business	(billio			ons of yen)
	Nine mo	nths ended	Dec. 31	Full year
	FY2012	FY2013	YOY	FY2012
			%	
Net sales	234.2	240.3	102.6	307.8
Segment profit	102.9	122.4	119.0	
Net Sales Breakdown				
Japan Net Sales				
Prescription Drugs	216.4	218.8	101.1	282.2
Generic Drugs (Elmed Eisai Co., Ltd.)	13.5	17.1	127.4	19.6
Diagnostics (EIDIA Co., Ltd.)	4.3	4.4	102.0	6.0
Japan prescription drugs—major products				
Anti-Alzheimer's agent Aricept	56.0	52.3	93.3	72.4
Proton pump inhibitor Pariet	38.6	36.9	95.7	50.1
Fully human anti-TNF-α monoclonal antibody Humira	18.1	22.3	123.3	24.1
Peripheral neuropathy treatment Methycobal	20.1	19.6	97.6	26.1
Pain treatment (neuropathic pain, fibromyalgia) Lyrica*	10.3	13.8	133.5	13.9
Oral anticoagulant Warfarin	7.7	7.5	97.1	10.1
Osteoporosis treatment Actonel	6.9	6.0	86.8	9.1
Gastritis / gastric ulcer treatment Selbex	6.0	5.3	87.5	7.8
Anticancer agent Halaven	4.1	5.0	120.5	5.5

* Co-promotion income

		Nine mor	ths ended	Dec. 31	Full year
		FY2012	FY2013	YOY	FY2012
				%	
Net sales	Billions JPY	114.5	106.8	93.3	153.3
				<75.1>	
Segment profit	Billions JPY	25.7	9.3	36.4	
Americas Prescription Drugs—major products					
Proton pump inhibitor	Billions JPY	37.8	33.4	88.3	51.4
Aciphex	[Millions USD]	[473]	[336]	<71.0>	[618]
Antiemetic agent Aloxi	Billions JPY	27.1	32.2	118.6 <95.4>	36.7
U.S. Prescription Drugs					
	Billions JPY	27.1	32.2	118.5	36.7
	[Millions USD]	[339]	[324]	<95.4>	[442]
DNA methylation inhibitor	Billions JPY	13.5	11.1	82.0	19.3
Dacogen	[Millions USD]	[169]	[112]	<66.0>	[232]
Anticancer agent Halaven	Billions JPY	8.5	9.9	116.1 <93.5>	11.6
U.S. Prescription Drugs	Billions JPY	8.5	9.7	114.1	11.6
	[Millions USD]	[106]	[98]	<91.8>	[139]
Antiepileptic agent Banzel	Billions JPY	3.8	5.6	148.0 <119.2>	5.2
				<119.ZZ	
U.S. Prescription Drugs	Billions JPY	3.8	5.5	147.3	5.1
	[Millions USD]	[47]	[56]	<118.5>	[62]
Anticoagulant	Billions JPY	7.8	5.2	67.0	9.7
Fragmin	[Millions USD]	[97]	[52]	<54.0>	[116]
Anti-Alzheimer's agent	Billions JPY	9.3	3.8	40.7	11.0
Aricept	[Millions USD]	[117]	[38]	<32.8>	[133]
Antiobesity agent	Billions JPY	-	1.7	-	-
Belviq	[Millions USD]	-	[17]	-	-

2) Americas Pharmaceutical Business (North, Central and South America)

* Sales of Aricept 23 mg tablet out of total sales of Aricept for FY2013 (April 1, 2013 to December 31, 2013) totaled ¥2.3 billion (U.S.\$23 million).

* The U.S. is the only country were Eisai markets Dacogen, Fragmin and Belviq; it is also the only country where Eisai directly markets Aciphex and Aricept. * Year-on-year percentage: figures shown in angle brackets "< >" exclude the effects of foreign currency fluctuations.

		Nine mor	nths ended	Dec. 31	Full year
		FY2012	FY2013	YOY	FY2012
				%	
Net sales	Billions JPY	29.7	43.3	146.0 <116.1>	41.3
Net sales in China	Billions JPY	15.8	23.6	148.9 <116.4>	21.8
Segment profit	Billions JPY	6.1	10.3	168.3	
Asia Prescription Drugs—major products					
Peripheral neuropathy treatment Methycobal	Billions JPY	7.7	12.2	158.9 <125.7>	10.4
China Prescription Drugs	Billions JPY [Millions RMB]	6.8 [534]	10.5 [645]	154.5 <120.8>	9.0 [681]
Anti-Alzheimer's agent Aricept	Billions JPY	5.9	8.9	150.9 <119.7>	8.1
China Prescription Drugs	Billions JPY [Millions RMB]	1.6 [127]	2.8 [170]	170.5 <133.3>	2.2 [163]
Fully human anti-TNF-α monoclonal antibody Humira	Billions JPY	3.5	5.3	152.1 <121.4>	4.9
Proton pump inhibitor Pariet	Billions JPY	3.3	4.2	128.2 <103.0>	4.3
China Prescription Drugs	Billions JPY [Millions RMB]	1.0 [82]	1.6 [99]	153.9 <120.3>	1.3 [99]
Liver disease / Allergic disease agents Stronger Neo-Minophagen C and Glycyron Tablets	Billions JPY	3.5	3.8	110.5 <86.5>	5.3
China Prescription Drugs	Billions JPY [Millions RMB]	3.4 [271]	3.8 [234]	110.4 <86.3>	5.3 [397]
Anticancer agent Halaven	Billions JPY	0.0	0.3	616.9 <503.2>	0.1

3) Asia Pharmaceutical Business (mainly China, South Korea, Taiwan, India and ASEAN)

* Year-on-year percentage: figures shown in angle brackets "<>" exclude the effects of foreign currency fluctuations.

		Nine mor	nths ended	Dec. 31	Full year
		FY2012	FY2013	YOY %	FY2012
Net sales	Billions JPY	18.9	24.3	128.8 <101.0>	25.8
Segment profit	Billions JPY	1.3	3.5	271.8	
EMEA Prescription Drugs—major products					
Anticancer agent Halaven	Billions JPY	3.6	6.3	171.9 <134.5>	5.4
Antiepileptic agent Zonegran	Billions JPY	3.4	5.0	147.0 <114.6>	4.8
Antiepileptic agent Zebinix	Billions JPY	1.2	1.8	152.7 <118.6>	1.8
Anti-Alzheimer's agent Aricept	Billions JPY	2.3	1.4	62.2 <48.4>	2.7
Antiepileptic drug Fycompa	Billions JPY	0.3	1.1	391.9 <309.4>	0.5
Proton pump inhibitor Pariet	Billions JPY	2.4	0.5	20.3 <16.2>	2.7

* Year-on-year percentage: figures shown in angle brackets "< >" exclude the effects of foreign currency fluctuations.

5) Consumer Healthcare Business—Japan (mainly OTC drugs) (billion					
	Nine mor	Nine months ended Dec. 31			
	FY2012	FY2013	YOY	FY2012	
			%		
Net sales	15.8	16.0	101.1	21.1	
Segment profit	2.9	2.9	99.1		
Consumer Healthcare Business—Japan—major products			0		
Vitamin B2 preparation, "Chocola BB Plus," etc. Chocola BB Group	8.6	9.2	106.8	11.2	

6) Sales of Major Products

(1) Oncology-Related Products

		Nine months ended Dec. 31			Full year
		FY2012	FY2013	YOY	FY2012
				%	
Total	Billions JPY	73.8	77.5	105.0	100.4
				<86.5>	
Halaven (Anticancer agent)	Billions JPY	16.4	21.5	131.2	22.6
				<110.7>	
Japan	Billions JPY	4.1	5.0	120.5	5.5
Americas	Billions JPY	8.5	9.9	116.1	11.6
				<93.5>	
U.S. Prescription Drugs	Billions JPY	8.5	9.7	114.1	11.6
	[Millions USD]	[106]	[98]	<91.8>	[139]
Asia	Billions JPY	0.0	0.3	616.9	0.1
				<503.2>	
EMEA	Billions JPY	3.6	6.3	171.9	5.4
				<134.5>	
Aloxi (Antiemetic agent)	Billions JPY	27.1	32.2	118.6	36.7
				<95.4>	
U.S. Prescription Drugs	Billions JPY	27.1	32.2	118.5	36.7
	[Millions USD]	[339]	[324]	<95.4>	[442]
Dacogen (DNA methylation inhibitor)	Billions JPY	13.5	11.1	82.0	19.3
	[Millions USD]	[169]	[112]	<66.0>	[232]
Fragmin (Anticoagulant)	Billions JPY	7.8	5.2	67.0	9.7
	[Millions USD]	[97]	[52]	<54.0>	[116]
Treakisym/Symbenda (Anticancer agent)	Billions JPY	2.7	3.0	111.9	3.5
				<110.8>	
Other	Billions JPY	6.3	4.5	71.3	8.6
				<59.1>	

* The U.S. is the only country where Eisai markets Dacogen and Fragmin.

* Year-on-year percentage: figures shown in angle brackets "<>" exclude the effects of foreign currency fluctuations.

(2) Pariet/Aciphex (Proton pump inhibitor)

		Nine mor	Nine months ended Dec. 31		
		FY2012	FY2013	YOY %	FY2012
Total	Billions JPY	82.1	75.0	91.3 <82.3>	108.4
Japan	Billions JPY	38.6	36.9	95.7	50.1
Americas	Billions JPY [Millions USD]	37.8 [473]	33.4 [336]	88.3 <71.0>	51.4 [618]
Asia	Billions JPY	3.3	4.2	128.2 <103.0>	4.3
EMEA	Billions JPY	2.4	0.5	20.3 <16.2>	2.7

* The U.S. is the only country in the Americas where Eisai directly markets Aciphex.

* Year-on-year percentage: figures shown in angle brackets "<>" exclude the effects of foreign currency fluctuations.

(3) Aricept (Anti-Alzheimer's agent)

		Nine mor	Nine months ended Dec. 31		
		FY2012	FY2013	YOY	FY2012
				%	
Total	Billions JPY	73.6	66.4	90.3	94.3
				<86.3>	
Japan	Billions JPY	56.0	52.3	93.3	72.4
Americas	Billions JPY	9.3	3.8	40.7	11.0
	[Millions USD]	[117]	[38]	<32.8>	[133]
Asia	Billions JPY	5.9	8.9	150.9	8.1
				<119.7>	
EMEA	Billions JPY	2.3	1.4	62.2	2.7
				<48.4>	

* The U.S. is the only country in the Americas where Eisai books the sales of Aricept.

* Year-on-year percentage: figures shown in angle brackets "<>" exclude the effects of foreign currency fluctuations.

(4) Humira (Fully human anti-TNF-α monoclonal antibody)

		Nine mor	Full year		
		FY2012	FY2013	YOY	FY2012
				%	
Total	Billions JPY	21.6	27.6	127.9	29.0
				<123.0>	
Japan	Billions JPY	18.1	22.3	123.3	24.1
Asia	Billions JPY	3.5	5.3	152.1	4.9
				<121.4>	

* Year-on-year percentage: figures shown in angle brackets "< >" exclude the effects of foreign currency fluctuations.

7) Overseas Sales (billions								
	Nine mor	ths ended D	ec. 31	Full year				
	FY2012	FY2013	YOY	FY2012				
			%					
Overseas sales	172.0	181.0	105.2	231.6				
Overseas sales (% of total sales)	39.9	40.4	-	40.4				
* Net sales to external customers for each segment								

Net sales to external customers for each segment

5. Sales Forecasts by Reporting Segment (FY2013)

			(bil	ions of yen)
<u>Ni</u>	ne months ended Dec. 31		Full year	
	FY2013	FY2012	FY2013 Revised forecast	forecast Previous forecast
Japan	240.3	307.8	<u>309.0</u>	310.5
Prescription Drugs	218.8	282.2	<u>280.0</u>	281.0
Anti-Alzheimer's agent Aricept	52.3	72.4	67.0	67.0
Proton pump inhibitor Pariet	36.9	50.1	<u>47.0</u>	45.0
Fully human anti-TNF-α monoclonal antibody Humira	22.3	24.1	30.0	30.0
Peripheral neuropathy treatment Methycobal	19.6	26.1	24.5	24.5
Oral anticoagulant Warfarin	7.5	10.1	10.0	10.0
Anticancer agent Halaven	5.0	5.5	7.0	7.0
Generic Drugs (Elmed Eisai Co., Ltd.)	17.1	19.6	<u>23.0</u>	23.5
Diagnostics (EIDIA Co., Ltd.)	4.4	6.0	6.0	6.0
Americas	106.8	153.3	<u>143.0</u>	138.5
U.S.	106.3	153.0	<u>142.5</u>	137.5
Asia	43.3	41.3	<u>57.5</u>	53.0
China	23.6	21.8	<u>31.0</u>	28.0
EMEA	24.3	25.8	<u>33.0</u>	32.5
Consumer Healthcare Business—Japan (mainly OTC drugs)	16.0	21.1	22.5	22.5
Vitamin B2 preparation, "Chocola BB Plus," etc. Chocola BB Group	9.2	11.2	13.0	13.0
Other	17.2	24.4	<u>22.0</u>	21.0
Consolidated net sales	448.0	573.7	<u>587.0</u>	578.0
Global net sales of major products				
Pariet/Aciphex	75.0	108.4	<u>90.0</u>	84.5
Aricept	66.4	94.3	<u>85.0</u>	81.0
Halaven	21.5	22.6	<u>30.0</u>	34.0

(billions of yen)

* Sales amounts by new reporting segments for FY2012 are provided for reference purposes only.

6. Consolidated Balance Sheet

1) Consolidated Balance Sheet <Assets>

1) Consolidated Balance Sheet <	Assets>				(billio	ns of yen)
	2013		2013			Diff.
	March 31	%	Dec. 31	%	% change	
Total current assets	530.7	53.6	505.8	51.7	95.3	(24.9)
Cash and deposits	88.7		94.3			5.6
Notes and accounts receivable-trade	185.5		186.7			1.2
Short-term investments	98.8		67.0			(31.8)
Inventories	87.6		90.4			2.8
Deferred tax assets	47.1		48.6			1.5
Other	23.2		18.9			(4.3)
Allowance for doubtful accounts	(0.1)		(0.1)			(0.0)
Total noncurrent assets	459.5	46.4	473.0	48.3	102.9	13.4
Total property, plant and equipment	142.2	14.4	142.8	14.6	100.4	0.5
Buildings and structures	85.9		88.5			2.6
Other	56.3		54.2			(2.1)
Total intangible assets	236.0	23.8	246.8	25.2	104.5	10.7
Goodwill	127.3		135.3			7.9
Sales rights	51.4		53.1			1.7
Core technology	43.7		46.2			2.5
Other	13.5		12.2			(1.4)
Total investments and other assets	81.2	8.2	83.5	8.5	102.7	2.2
Investment securities	34.3		32.7			(1.6)
Deferred tax assets	40.7		43.8			3.1
Other	6.3		7.0			0.7
Allowance for doubtful accounts	(0.1)		(0.1)			0.0
Total assets	990.2	100.0	978.8	100.0	98.8	(11.5)

Notes Total assets

Decrease in cash and deposits and short-term investments due to redemption of matured bonds and debentures of ¥50.0 billion and repayment of long-term borrowings of US\$200 million

2) Consolidated Balance Sheet <lial< th=""><th>2013</th><th></th><th></th><th></th><th>(2</th><th>ns of yen)</th></lial<>	2013				(2	ns of yen)
	Z013 March 31	%	2013 Dec. 31	%	% change	Diff.
Total current liabilities	215.7	21.8	211.0	21.6	97.8	(4.8)
Notes and accounts payable—trade	26.1		26.1			0.1
Short-term borrowings	7.6		15.2			7.6
Long-term borrowings (current portion)	18.8		45.5			26.7
Commercial paper	-		10.0			10.0
Bonds and debentures (current portion)	50.0		-			(50.0)
Accounts payable—other / Accrued expenses	82.1		76.5			(5.7)
Income taxes payable	7.4		9.2			1.8
Reserve for sales rebates	15.7		16.4			0.8
Other	8.1		12.0			3.9
Total noncurrent liabilities	300.2	30.3	259.5	26.5	86.5	(40.7)
Bonds and debentures	30.0		30.0			0.0
Long-term borrowings	207.6		166.6			(41.0)
Deferred tax liabilities	19.6		19.6			(0.0)
Liability for retirement benefits	13.8		13.5			(0.3)
Other	29.1		29.7			0.6
Total liabilities	515.9	52.1	470.5	48.1	91.2	(45.4)
Total shareholders' equity	532.5	53.8	519.5	53.1	97.6	(13.0)
Common stock	45.0		45.0			_
Capital surplus	56.9		57.0			0.1
Retained earnings	469.7		456.4			(13.3)
Treasury stock	(39.0)		(38.8)			0.2
Total accumulated other comprehensive income (loss)	(63.2)	(6.4)	(15.4)	(1.6)	-	47.7
Valuation difference on available-for-sale securities	4.3		5.3			0.9
Deferred gain (loss) on derivatives under hedge accounting	(1.0)		(0.7)			0.2
Foreign currency translation adjustments	(66.5)		(20.0)			46.6
Stock options	1.1	0.1	1.1	0.1	101.6	0.0
Minority interests	3.9	0.4	3.0	0.3	79.1	(0.8)
Total equity	474.3	47.9	508.2	51.9	107.2	33.9
Total liabilities and equity	990.2	100.0	978.8	100.0	98.8	(11.5)

Notes

Total liabilities

Decrease due to redemption of matured bonds and debentures of ¥50.0 billion and repayment of long-term borrowings of US\$200 million

Total equity

Increase in yen equivalent amount of equity of overseas subsidiaries due to yen depreciation compared with that of the previous fiscal year end

7. Changes in Consolidated Quarterly Results

		FY20	12		<u> </u>	FY2013	
	1st Quarter	2nd Quarter	3rd Quarter	4th Quarter	1st Quarter	2nd Quarter	3rd Quarter
Net sales	146.9	141.6	143.1	142.1	154.2	153.2	140.6
Cost of sales	43.2	41.8	43.2	46.0	46.3	47.0	48.3
R&D expenses	28.4	29.1	29.8	33.1	39.6	30.8	35.8
SG&A expenses	56.2	52.5	53.3	46.7	54.2	54.3	49.0
Operating income	19.1	18.2	16.8	16.3	14.2	21.2	7.4
Ordinary income	17.9	16.6	15.7	15.4	12.9	19.9	6.5
Net income	11.9	12.6	9.5	14.3	9.4	18.2	1.8
Cash income	24.3	25.9	22.8	27.9	21.9	30.1	14.3
Comprehensive income	(1.1)	8.5	43.7	44.0	26.1	18.1	33.3
Earnings per share (EPS, yen)	41.7	44.2	33.4	50.0	33.0*	64.0*	6.5*
Cash income per share (Cash EPS, yen)	85.1	90.8	79.9	97.7	76.8*	105.6*	50.1*

* The Company's stock held through the Trust for Officers' Compensation Board Incentive Plan (105,400 shares) is included in the average number

of shares outstanding as treasury stocks which are deducted from the basis of the calculation of basic earnings per share. * "Cost of sales" includes "Provision for (reversal of) sales returns—net."

2) Cash Flow Segment Data

2) Cash Flow Segment Data						(billior	ns of yen)	
	<u>FY2012</u>				FY2013			
	1st	2nd	3rd	4th	1st	2nd	3rd	
	Quarter	Quarter	Quarter	Quarter	Quarter	Quarter	Quarter	
Net cash provided by (used in) operating activities	28.3	8.6	16.6	19.7	12.3	28.9	17.7	
Net cash provided by (used in) investing activities	7.2	24.2	(10.4)	0.8	29.0	(8.5)	(3.5)	
Net cash provided by (used in) financing activities	(20.6)	(42.6)	2.3	(21.0)	(82.1)	(5.7)	(7.8)	
Cash and cash equivalents at the end of period	123.3	112.3	133.3	142.5	108.3	122.8	140.9	
Free cash flow	22.3	3.3	13.4	15.6	2.7	29.0	12.0	

* "Free cash flow" = "Net cash provided by (used in) operating activities" - "Capital expenditures, etc. (cash basis)"

3) Balance Sheet Data

3) Balance Sheet Data						(billior	is of yen)
		<u>FY20</u>	12		ļ	FY2013	
	June 30	Sep.30	Dec.31	March 31	June 30	Sep.30	Dec.31
Total assets	977.2	921.9	968.2	990.2	939.8	952.2	978.8
Liabilities	577.7	515.8	538.3	515.9	462.1	457.3	470.5
Borrowings	266.1	222.6	225.5	234.0	227.5	221.8	227.4
Commercial paper	-	-	25.0	-	-	-	10.0
Bonds and debentures	80.0	80.0	80.0	80.0	30.0	30.0	30.0
Equity	399.5	406.1	430.0	474.3	477.7	494.9	508.2
Shareholders' equity	392.9	401.4	425.2	469.4	472.7	490.9	504.1
Shareholders' equity ratio (%)	40.2	43.5	43.9	47.4	50.3	51.5	51.5
Liabilities ratio (Net DER / times)	0.39	0.38	0.35	0.27	0.30	0.23	0.21

* "Liabilities ratio (Net DER)"=("Interest-bearing debt" ("Borrowings" + "Bonds and debentures") - "Cash and deposits" - "Short-term investments") / "Shareholders' equity"

4) Capital Expenditures, Depreciation a	nd Amort	ization				(billior	ns of yen)	
		FY20	12		<u>FY2013</u>			
	1st	2nd	3rd	4th	1st Overter	2nd	3rd	
	Quarter	Quarter	Quarter	Quarter	Quarter	Quarter	Quarter	
Capital expenditures	7.0	4.4	3.6	5.5	8.4	3.1	5.3	
Property, plant and equipment	1.3	2.2	1.5	4.1	1.5	2.2	1.4	
Intangible assets	5.6	2.2	2.0	1.5	6.9	0.9	4.0	
Depreciation and amortization	10.2	10.6	10.9	11.5	10.2	9.6	9.6	

* "Depreciation and amortization" includes amortization of "Intangible assets."

5) Sales of Major Products

(1) Oncology-Related Products

			<u>FY20</u>	12		 -	FY2013	
		1st Quarter	2nd Quarter	3rd Quarter	4th Quarter	1st Quarter	2nd Quarter	3rd Quarter
Total	Billions JPY	25.2	23.3	25.3	26.6	27.1	26.7	23.7
Halaven	Billions JPY	5.5	5.3	5.6	6.2	7.0	6.9	7.5
Japan	Billions JPY	1.3	1.4	1.4	1.4	1.6	1.6	1.7
Americas	Billions JPY	3.1	2.7	2.7	3.1	3.3	3.2	3.4
U.S. Prescription Drugs	Billions JPY [Millions USD]	3.1 [39]	2.7 [34]	2.7 [34]	3.1 [33]	3.2 [32]	3.2 [32]	3.3 [33]
Asia	Billions JPY	0.0	0.0	0.0	0.1	0.1	0.1	0.2
EMEA	Billions JPY	1.0	1.2	1.4	1.7	2.0	2.0	2.3
Aloxi	Billions JPY	9.5	8.4	9.3	9.6	10.3	11.1	10.8
U.S. Prescription Drugs	Billions JPY [Millions USD]	9.5 [119]	8.4 [107]	9.3 [114]	9.6 [103]	10.3 [105]	11.1 [112]	10.8 [107]
Dacogen	Billions JPY [Millions USD]	4.4 [55]	4.4 [55]	4.8 [59]	5.8 [63]	5.7 [58]	3.5 [35]	1.9 [19]
Fragmin	Billions JPY [Millions USD]	2.9 [36]	2.5 [32]	2.4 [30]	1.9 [19]	1.8 [18]	2.3 [24]	1.1 [10]
Treakisym/Symbenda	Billions JPY	0.9	0.9	0.9	0.8	1.0	1.0	1.0
Other	Billions JPY	2.0	1.9	2.4	2.2	1.3	1.8	1.4

* The U.S. is the only country where Eisai markets Dacogen and Fragmin.

(2) Pariet/Aciphex

			<u>FY20</u>	12		<u> </u>	<u>-Y2013</u>	
		1st Quarter	2nd Quarter	3rd Quarter	4th Quarter	1st Quarter	2nd Quarter	3rd Quarter
Total	Billions JPY	28.5	24.8	28.8	26.4	29.9	28.9	16.2
Japan	Billions JPY	13.1	12.7	12.8	11.5	12.8	12.4	11.8
Americas	Billions JPY [Millions USD]	13.2 [164]	10.2 [131]	14.4 [178]	13.6 [146]	15.5 [157]	15.1 [152]	2.8 [27]
Asia	Billions JPY	1.1	1.1	1.1	1.1	1.4	1.3	1.4
EMEA	Billions JPY	1.2	0.8	0.5	0.2	0.2	0.1	0.2

* The U.S. is the only country in the Americas where Eisai directly markets Aciphex.

(3) Aricept

			<u>FY20</u>	12		<u> </u>	FY2013	
		1st Quarter	2nd Quarter	3rd Quarter	4th Quarter	1st Quarter	2nd Quarter	3rd Quarter
Total	Billions JPY	27.3	26.2	20.1	20.7	23.2	22.1	21.1
Japan	Billions JPY	21.7	18.6	15.7	16.4	18.3	17.3	16.7
Americas	Billions JPY [Millions USD]	2.4 [30]	5.1 [64]	1.9 [23]	1.7 [16]	1.5 [16]	1.3 [13]	0.9 [9]
Asia	Billions JPY	2.0	1.9	2.0	2.2	2.9	3.0	2.9
EMEA	Billions JPY	1.2	0.6	0.5	0.4	0.4	0.5	0.5

 * The U.S. is the only country in the Americas where Eisai directly markets Aricept.

(4) Humira

			<u>FY20</u>		<u>FY2013</u>			
		1st	2nd	3rd	4th	1st	2nd	3rd
		Quarter	Quarter	Quarter	Quarter	Quarter	Quarter	Quarter
Total	Billions JPY	6.8	7.2	7.6	7.5	8.5	9.3	9.8
Japan	Billions JPY	5.8	6.1	6.3	6.0	6.9	7.4	8.0
Asia	Billions JPY	1.1	1.1	1.3	1.4	1.6	1.8	1.8

8. Nonconsolidated Financial Highlights

1) Nonconsolidated Financial Highlights

	(1)	Income	Statement Data
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(1) Income Statement Data			(billi	ons of yen)
	Nine mor	ths ended D	ec. 31	Full Year
	FY2012	FY2013	YOY	FY2012
			%	
Net sales	264.0	266.0	100.8	348.0
Cost of sales	73.8	76.3	103.5	97.8
R&D expenses	82.1	98.9	120.5	111.0
SG&A expenses	81.0	60.0	74.1	100.5
Operating income	27.1	30.7	113.2	38.7
Ordinary income	24.3	28.2	115.9	34.9
Net income	17.6	23.3	132.2	27.6

* "Cost of sales" includes "Provision for (reversal of) sales returns-net."

(2) Cash Flow Statement Data

(2) Cash Flow Statement Data			(billi	ons of yen)
	Nine months ended Dec. 31			Full Year
	FY2012	FY2013	Diff.	FY2012
Net cash provided by (used in) operating activities	28.1	30.1	1.9	40.6
Net cash provided by (used in) investing activities	27.3	24.3	(3.0)	28.6
Net cash provided by (used in) financing activities	(60.7)	(65.5)	(4.8)	(56.6)
Cash and cash equivalents at end of period	8.3	15.1	6.8	26.1
Free cash flow	19.2	23.2	4.0	28.9

* "Free cash flow" = "Net cash provided by (used in) operating activities" - "Capital expenditures (cash basis)"

(3) Balance Sheet Data

(3) Balance Sheet Data		(billior	billions of yen)	
		<u>2013</u>		
	March 31	Dec. 31	Diff.	
Total assets	891.7	853.1	(38.6)	
Liabilities	375.2	354.2	(21.0)	
Borrowings	202.5	220.0	17.5	
Commercial paper	-	10.0	10.0	
Bonds and debentures	80.0	30.0	(50.0)	
Equity	516.5	498.9	(17.6)	
Shareholders' equity	515.4	497.8	(17.6)	
Shareholders' equity ratio (%)	57.8	58.3	0.5	

2) Net Sales Highlights

2) Net Sales Highlights (billion				
	Nine months ended Dec. 31		Full year	
	FY2012	FY2013	YOY	FY2012
			%	
Net sales	264.0	266.0	100.8	348.0
Prescription drugs	216.3	218.7	101.1	282.1
Consumer healthcare products, etc.	16.0	16.1	101.0	21.2
Industrial property rights, etc.	3.4	4.7	137.2	6.1
Export of pharmaceuticals	27.3	25.7	94.0	37.5
Other	0.9	0.7	82.1	1.2

9. Major News Releases

Date	Description
April 2013	• Eisai Establishes Pharma Sales Subsidiary in Moscow Ahead of Planned Direct Sales Launch in Russia
	<issued 5="" april="" on=""></issued>
	Eisai Co-establishes the Global Health Innovative Technology Fund <issued 8="" april="" on=""></issued>
	Eisai Announces Preclinical Research Findings Suggesting Novel Inhibitory Effect on Tumor Metastasis
	for Anticancer Agent Halaven at AACR 104th Annual Meeting <issued 10="" april="" on=""></issued>
	• Eisai Receives Manufacturing and Marketing Authorization for Vascular Embolization Device DC Bead in Japan
	<issued 17="" april="" on=""></issued>
	Eisai to Establish New Parenteral Facility in China <issued 25="" april="" on=""></issued>
	Eisai Supports Earthquake Relief Efforts in Sichuan, China <issued 26="" april="" on=""></issued>
lay	Eisai Files for Indication Expansion of Anticancer Agent Halaven with European Medicines Agency
,	<pre><issued 7="" may="" on=""></issued></pre>
	Publication in Federal Register Tomorrow Moves Belvig Closer to Launch <issued 8="" may="" on=""></issued>
	Notification Regarding the Introduction of a Performance-Related Stock Compensation System in Accordance
	with the Revision of the Compensation System for the Corporate Officers of the Company
	<issued 13="" may="" on=""></issued>
	Notification Regarding the Disposal of Treasury Stock through Third-Party Allotment in Accordance with
	the Introduction of Performance-Related Stock Compensation System <issued 13="" may="" on=""></issued>
	Eisai to Present New Research on Oncology Products and Pipeline at 49th ASCO Annual Meeting
	cissued on May 17>
	AbbVie and Eisai Announce Humira Pre-filled Syringe Has Received Approval for the Treatment of
	Intestinal Behçet's Disease in Japan <issued 24="" may="" on=""></issued>
	• Eisai Announces Launch of Antiepileptic Agent Inovelon Tablets 100 mg, 200 mg in Japan <issued 28="" may="" on=""></issued>
	Notification Regarding the Completion of the Disposal of Treasury Stock Through Third-Party Allotment in
	Accordance with the Introduction of Performance-Related Stock Compensation System <issued 30="" may="" on=""></issued>
	Notification Regarding Partial Amendment to the Articles of Incorporation <issued 30="" may="" on=""></issued>
une	Eisai Presents New Quality of Life Findings in Patients with Metastatic Breast Cancer from Halaven (Eribulin)
	Versus Capecitabine Study at 49th ASCO Annual Meeting <issued 3="" june="" on=""></issued>
	Eisai Confirms Therapeutic Effects of Lenvatinib in Patients with Melanoma in Strategic Collaboration with
	Quintiles to Develop Eisai Anticancer Compounds <issued 10="" june="" on=""></issued>
	Eisai to Launch Chocola BB Sparkling Vitamin Kyutto Lemon Flavor <issued 11="" june="" on=""></issued>
	 AbbVie and Eisai Announce Humira Pre-filled Syringe 40 mg / 0.8 mL, a Fully Human Monoclonal
	Anti-TNF-α Antibody Formulation, Has Received Approval for the Treatment of Moderate to Severe
	Ulcerative Colitis (UC) in Japan <issued 14="" june="" on=""></issued>
	Eisai to Make New Investment in Expansion of Global Packaging Facility at Hatfield Production Plant in U.K.
	<issued 17="" june="" on=""></issued>
	Eisai Announces Launch of New Dry Syrup Formulation of Alzheimer's Disease Treatment Aricept in Japan
	<pre><issued 25="" june="" on=""></issued></pre>
	 Eisai to Suspend Temporarily Commercial Distribution of Antiepileptic Drug Fycompa in Germany
	<issued 25="" june="" on=""></issued>
uly	Notice Regarding Transfer of Shares of Eisai Subsidiary, Eisai Seikaken Co., Ltd. <issued 19="" july="" on=""></issued>
	Eisai Receives Positive Opinion from EMA's CHMP on Use of Antiepileptic Agent Zonegran in Pediatric
	Patients <issued 29="" july="" on=""></issued>
lugust	 Continuation of "Policy for Protection of The Company's Corporate Value and Common Interests of
	Shareholders (Shareholder Rights Plan)" <issued 1="" august="" on=""></issued>
	Eisai Receives Approval to Market Pariet Triple Formulation Packs Rabecure 400 and 800 and Rabefine,
	for Primary and Secondary H. Pylori Eradication Respectively, in Japan <issued 21="" august="" on=""></issued>
	Eisai Announces Launch of Higher-Dose Aricept 23 mg Tablet for Moderate-to-Severe Alzheimer's Disease
	in South Korea <issued 27="" august="" on=""></issued>
	Eisai Receives Prequalification from World Health Organization For Lymphatic Filariasis Medicine
	Diethylcarbamazine <issued 27="" august="" on=""></issued>
	Eisai Enters Community Development Partnership Agreement with City of Yokohama to Promote
	Local Dementia Support Initiatives <issued 29="" august="" on=""></issued>

Date	Description
September	Eisai Announces Launch of Selbelle Ukon 27 Plus Granule in Japan <issued 11="" on="" september=""></issued>
	Eisai Announces Launch of Anticancer Agent Halaven As Company's First Product in Russia
	<issued 12="" on="" september=""></issued>
	Eisai's U.S. Research Subsidiary H3 Biomedicine Enters into Collaborative Agreement with Selvita to
	Create Novel Anticancer Agents <issued 18="" on="" september=""></issued>
	Eisai Selected for Membership of Dow Jones Sustainability Asia Pacific Index, An Index for Socially
	Responsible Investment <issued 25="" on="" september=""></issued>
October	• Eisai Presents Additional Analysis Findings on Halaven (Eribulin) at European Cancer Congress 2013
	<issued 1="" october="" on=""></issued>
	Eisai Announces Launch of Anticancer Agent Halaven in India <issued 1="" october="" on=""></issued>
	• Eisai Receives European Commission Approval on Use of Antiepileptic Agent Zonegran in Pediatric Patients
	<issued 3="" october="" on=""></issued>
	Eisai to Expand Antiobesity Agent Belviq Sales Force in U.S. <issued 16="" october="" on=""></issued>
	Eisai Begins Free Supply of Diethylcarbamazine in Line with Its Global Commitment to Eliminate
	Lymphatic Filariasis <issued 29="" october="" on=""></issued>
	• Eisai Submits Application to Expand Indication of Anti-Alzheimer's Agent Aricept As Treatment for Dementia
	with Lewy Bodies in Japan <issued 31="" october="" on=""></issued>
November	Eisai Expands Marketing and Supply Agreement for Antiobesity Agent Lorcaserin to Include Most Countries
	Worldwide <issued 8="" november="" on=""></issued>
	Eisai Enters Global Agreement with Broad Institute to Develop New Drugs for Neglected Tropical Diseases
	and Tuberculosis <issued 8="" november="" on=""></issued>
	Eisai Makes Donation to Victims of Typhoon Haiyan in Philippines <issued 12="" november="" on=""></issued>
	Eisai Announces Launch of New "Crystal Veil Fuite Bokin 24" Wet Wipes Containing Long-Acting
	Antimicrobial Agent Etak <issued 20="" november="" on=""></issued>
	Eisai Joins Groundbreaking Tuberculosis Drug Accelerator Partnership to Discover New Tuberculosis Drugs
	<issued 25="" november="" on=""></issued>
	Eisai to Receive Japan Marketing Authorization Holder License from Nobelpharma for Antineoplastic
	Agent Gliadel 7.7 mg Implant <issued 25="" november="" on=""></issued>
	Eisai Submits Application for Proton Pump Inhibitor Pariet in Japan Seeking Indication Expansion for
	Prevention of Recurrent Gastric or Duodenal Ulcer Caused by Low-Dose Aspirin Therapy and Approval
	of New Formulation <issued 28="" november="" on=""></issued>
	• Eisai Product Creation Systems (EPCS) Undergoes Transformation with Aim of Further Focusing and
	Strengthening Product Creation Capabilities <issued 29="" november="" on=""></issued>
	Notification Regarding Implementation of Voluntary Retirement Program <issued 29="" november="" on=""></issued>
	Notification Regarding Execution of Business Transfer Agreement to Transfer Business Operations
	at Eisai Misato Plant to Bushu Pharmaceuticals Ltd. <issued 29="" november="" on=""></issued>
December	• Eisai to Present New Research on Halaven (Eribulin) at 36th Annual San Antonio Breast Cancer Symposium
	<issued 5="" december="" on=""></issued>
	Launch of Measurement Reagent Kit "E Test Tosoh II (PIVKA-II)" for Use with Tosoh AIA Systems in Detecting
	Hepatocellular Carcinoma Diagnostic-Aid Marker PIVKA-II/DCP in Japan <issued 13="" december="" on=""></issued>
January	Eisai Announces Launch of Antiepileptic Drug Fycompa In U.S. <issued 6="" january="" on=""></issued>
- and any	Eisai Announces Launch of New "Crystal Veil Mask Bokin 24 Mint" Face Mask Spray Containing Long-Acting
	Antimicrobial Agent Etak <issued 22="" january="" on=""></issued>
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10. Major R&D Pipeline

In-House R&D Pipeline List

	Product Name / Research Code	Additional Indication, etc.*	Development Stage**	Therapeutic Area
Ne	w Approval			
0	DC Bead (Transcatheter arterial embolization (TAE) of hepatocellular carcinoma)		(JP) approved	Oncology and Supportive Care
0	Zonegran (Pediatric partial-onset seizures)	AI	(EU) approved	Neurology
С	Humira (Intestinal Behçet's disease)	AI	(JP) approved	Vascular and Immunological Reaction
)	Humira (Ulcerative colitis)	AI	(JP) approved	Vascular and Immunological Reaction
C	Pariet (Triple formulation pack for Helicobacter pylori eradication)	AF	(JP) approved	Gastrointestinal and Hepatic Disord
Su	bmitted / Preparing for Submission		1 、 7 …	
	cinitapride (Functional dyspepsia)		(CN) submitted	Gastrointestinal and Hepatic Disord
)	Aricept (Lewy body dementia)	AI	(JP) submitted	Neurology
С	Halaven (Second-line treatment for breast cancer)	AI	(EU) submitted	Oncology and Supportive Care
0	Pariet (Prevention of recurrence of gastric and duodenal ulcers during treatment with low-dosage aspirin, new 5 mg tablet)	AI, AF	(JP) submitted	Gastrointestinal and Hepatic Disord
0	Tambocor (Pediatric fine granule formulation)	AF	(JP) submitted	Vascular and Immunological Reaction
Cli	nical			
	Fycompa (Partial-onset seizures)		(JP/CN/AS) PIII	Neurology
	E5501 (Idiopathic thrombocytopenic purpura (ITP))		(US/EU/AS) PIII	Vascular and Immunological Reaction
٢	E5501 (Thrombocytopenia in chronic liver disease requiring surgery)		(US/EU/AS) PIII	Vascular and Immunological Reacti
	E5564 (Severe sepsis)		(JP/US/EU) PIII	Vascular and Immunological Reacti
)	Halaven (Third-line treatment for breast cancer)		(CN) PIII	Oncology and Supportive Care
	E7080 (Thyroid cancer)		(JP/US/EU/AS) PIII	Oncology and Supportive Care
	E7080 (Hepatocellular carcinoma)		(JP/US/EU/CN/AS) PIII	Oncology and Supportive Care
	MORAb-003 (Platinum-sensitive ovarian cancer)		(JP/US/EU/AS) PIII	Oncology and Supportive Care
	Fycompa (Generalized seizures)	AI	(JP/US/EU/AS) PIII	Neurology
	Halaven (Non-small cell lung cancer)	AI	(JP/US/EU/AS) PIII	Oncology and Supportive Care
	Halaven (Sarcoma)	AI	(US/EU/AS) PIII	Oncology and Supportive Care
0	Halaven (First-/second-line treatment for HER2-negative breast cancer)	AI	(US) PIII	Oncology and Supportive Care
	DC Bead (Transcatheter arterial embolization (TAE) of hypervascular tumors)	AI	(JP) PIII	Oncology and Supportive Care
	Aricept (Severe Alzheimer's disease)	AI	(CN) PIII	Neurology
	Inovelon/Banzel (Pediatric Lennox-Gastaut syndrome)	AI	(US/EU) PIII	Neurology
C	Pariet (Maintenance therapy for proton pump inhibitor (PPI)-resistant reflux esophagitis)	AI	(JP) PIII	Gastrointestinal and Hepatic Disord
	Aricept (Higher dose 23 mg tablet)	ADA, AF	(JP) PIII	Neurology
	E0302 (Amyotrophic lateral sclerosis (ALS))		(JP) PII/III	Neurology
	AS-3201 (Diabetic neuropathy)		(US/EU) PII/III	Neurology
	BAN2401 (Alzheimer's disease)		(US/EU) PII	Neurology
)	E2006 (Insomnia)		(US) PII	Neurology
-	E5501 (Thrombocytopenia during interferon therapy (both initiation and maintenance) for hepatitis C)		(US) PII	Vascular and Immunological Reacti
	E6005 (Atopic dermatitis)		(JP) PII	Vascular and Immunological Reacti
	E7016 (Melanoma)		(US) PII	Oncology and Supportive Care
	E7080 (Endometrial cancer)		(US/EU) PII	Oncology and Supportive Care
	E7080 (Melanoma)		(US/EU) PII	Oncology and Supportive Care
	E7080 (Glioma)		(US) PII	Oncology and Supportive Care
	E7080 (Non-small cell lung cancer)		(JP/US/EU/AS) PII	Oncology and Supportive Care
	E7820 (Colorectal cancer)		(US/EU) PII	Oncology and Supportive Care
	MORAb-003 (Non-small cell lung cancer)		(US/EU) PII	Oncology and Supportive Care
	MORAb-004 (Melanoma)		(US/EU) PII	Oncology and Supportive Care
	MORAb-004 (Colorectal cancer)		(US/EU) PII	Oncology and Supportive Care
	MORAb-004 (Sarcoma)		(US/EU) PII	Oncology and Supportive Care
	MORAb-009 (Mesothelioma)		(US/EU) PII	Oncology and Supportive Care
	Fycompa (Pediatric partial-onset seizures)	AI	(US/EU) PII	Neurology
)	Aricept (Regression symptoms in people with Down syndrome)	AI	(JP) PII	Neurology
	Halaven (Sarcoma)	AI	(JP) PII	Oncology and Supportive Care
	Ontak (Melanoma)	AI	(US) PII	Oncology and Supportive Care
	Dacogen (Pediatric acute myeloid leukemia (AML))	AI	(US) PII	Oncology and Supportive Care
		1	(30)	subology and supportive sale

* Al: Additional Indication, ADA: Additional Dosage & Administration, AF: Additional Formulation

** P: Clinical Phase; JP: Japan, US: United States, EU: Europe, CN: China, AS: Asia (excluding Japan and China)

• Eisai received a non-approval letter from the Chinese regulatory authority for clevudine as a treatment for patients with chronic hepatitis B. Future development plans are currently under review.

• Eisai decided to discontinue development of the multi-kinase inhibitor E6201, which was in a Phase II study in the United States and Europe as a potential treatment for psoriasis.

• After reviewing development plans for Halaven as a second-line treatment for breast cancer in the United States, Eisai has now begun a new Phase III study to investigate the agent as a potential first-/second-line treatment for HER2-negative breast cancer.

O Development progress from April 2013 onwards O Development progress from October 2013 onwards

(1) Oncology and Supportive Care

Product Name: Halaven Research Code: E7389 Generic Name: eribulin (Anticancer agent / microtubule dynamics inhibitor)

Description: A synthetic analog of halichondrin B derived from the marine sponge, *Halichondria okadai*. Believed to exert an antitumor effect by arresting the cell cycle through inhibition of the growth of microtubules. Currently being investigated as a potential treatment for breast cancer and various other solid tumors. Approved in 52 countries including the United States, Singapore, European Union (EU) member states, Japan, and Switzerland.

0	Additional Indication: Second-line treatment for breast cancer	EU: submitted (April 20	113), accepted (April 2013)	Inj.
0	Third-line treatment for breast cancer	CN: PIII		Inj.
O	Additional Indication: First-/second-line treatment for HER2-negative breast cancer	US: PIII		Inj.
	Additional Indication: Non-small cell lung cancer	JP/US/EU/AS: PIII	Submission Target: FY2014	Inj.
	Additional Indication: Sarcoma	US/EU/AS: PIII JP: PII	Submission Target: FY2014	Inj.

• After reviewing development plans for Halaven as a second-line treatment for breast cancer in the United States, Eisai has now begun a new Phase III study to investigate the agent as a potential first-/second-line treatment for HER2-negative breast cancer.

• The submission timeline for non-small cell lung cancer has been reviewed and subsequently changed from FY2013 to FY2014.

Research Code: E7820 (Anticancer agent / alpha 2 integrin suppressant)

Description: An angiogenesis inhibitor that suppresses the expression of alpha 2 integrin, a vascular endothelial cell adhesion molecule.
Colorectal cancer US/EU: PII Oral

Research Code: E7080 Generic Name: lenvatinib

(Anticancer agent / selective tyrosine kinase inhibitor with a novel binding mode)

escription: Selective tyrosine kinase inhibitor with rious solid tumors.	h a novel binding mode. Currently being in	nvestigated as a potential treatm	nent fo
Thyroid cancer	JP/US/EU/AS: PIII	Submission Target: FY2014	Ora
Hepatocellular carcinoma	JP/US/EU/CN/AS: PIII		Ora
Endometrial cancer	US/EU: PII		Ora
Melanoma	US/EU: PII		Ora
Glioma	US: PII		Ora
Non-small cell lung cancer	JP/US/EU/AS: PII		Ora

• The submission timeline for thyroid cancer has been reviewed and subsequently changed from FY2013 to FY2014.

Research Code: E7016 (Anticancer agent / poly (ADP-ribose) polymerase inhibitor)

Description: Poly (ADP-ribose) polymerase (PARP) is an enzyme that is involved in DNA repair. PARP inhibitors exhibit an antitumor effect by inhibiting DNA repair in tumor cells and are expected to enhance the effect of chemotherapy and radiotherapy, both of which damage DNA.

US: PII

Melanoma

Research Code: MORAb-003 Generic Name: farletuzumab (Anticancer agent / humanized anti-FRA monoclonal antibody)

Description: A humanized IgG1 monoclonal antibody that targets folate receptor alpha (FRA). Expected to exhibit an antitumor effect				
against carcinomas that over-express FRA.				
Platinum-sensitive ovarian cancer	JP/US/EU/AS: PIII	Inj.		
Non-small cell lung cancer	US/EU: PII	Inj.		

Oral

Research Code: MORAb-004 (Anticancer agent / humanized anti-endosialin monoclonal antibody)

Description: A humanized IgG1 monoclonal antibody that targets Tumor Endothelial Marker 1 (TEM-1) / endosialin. Expected to exhibit an antitumor effect against carcinomas that express endosialin.				
Melanoma	US/EU: PII	Inj.		
Colorectal cancer	US/EU: PII	Inj.		

US/EU: PII

Research Code: MORAb-009 Generic Name: amatuximab (Anticancer agent / chimeric anti-mesothelin monoclonal antibody)

Description: A chimeric IgG1 monoclonal antibody that blocks the function of mesothelin. Expected to exhibit an antitumor effect against	l
carcinomas that express mesothelin.	J
	i .

Mesothelioma	US/EU: PII	Inj.

Product Name: Dacogen Research Code: E7373 Generic Name: decitabine (DNA methylation inhibitor)

Description: Induces cell differentiation by inhibiting DNA methylation. Currently approved in the United States for the treatment of myelodysplastic syndromes (MDS).

Additional Indication: Pediatric acute myeloid leukemia (AML) US: PII Inj.

Product Name: Ontak Research Code: E7272 Generic Name: denileukin diftitox

(Anticancer agent / interleukin-2 diphtheria toxin fusion protein)

Sarcoma

Description: A fusion protein that combines the interleukin-2 (IL-2) receptor binding domain with diphtheria toxins. Specifically binds to IL-2 receptors on the cell surface, causing diphtheria toxins that have entered cells to inhibit protein synthesis. Already approved in the United States as a treatment for CD25 (a component of the IL-2 receptor) positive cutaneous T-cell lymphoma.

Additional Indication: Melanoma	US: PII	Inj.

Product Name: **DC Bead** Research Code: **E7040** (Embolic bead / medical device)

Description: Contains hydrophilic microspherical particles produced from cross-linked polyvinyl alcohol polymer. These embolic beads are injected through a catheter to selectively embolize targeted blood vessels. The beads are microscopic and uniformly spherical in shape, allowing for sustained embolization of targeted vessels based on vascular diameter and tumor size. Approved in Japan as a device for transcatheter arterial embolization (TAE) therapy in patients with hepatocellular carcinoma.

0	Transcatheter arterial embolization (TAE) of hepatocellular carcinoma	JP: approved	d (April 2013)	Embolic Agent
	Additional Indication: Transcatheter arterial embolization	JP: PIII	Submission Target: FY2014	Embolic
	(TAE) of hypervascular tumors		6	Agent

lnj.

(2) Neurology

Product Name: Aricept Research Code: E2020 Generic Name: donepezil (Anti-Alzheimer's agent)

Description: Increases levels of the neurotransmitter acetylcholine in the brain by inhibiting its breakdown by the enzyme acetylcholinesterase, thereby slowing the overall progression of symptoms associated with Alzheimer's disease (AD). Currently approved in more than 90 countries around the world for the treatment of mild to moderate AD. It is also approved as a treatment for patients with severe AD in numerous countries including the United States, Japan, Canada, and several other Asian and Latin American countries.

0	Additional Indication: Lewy body dementia	JP: submitted (O	ctober 2013)	Oral
	Additional Indication: Severe Alzheimer's disease	CN: PIII		Oral
	Additional Dosage & Administration, Formulation:	JP: PIII Submission Target: FY201		Oral
	Higher dose 23 mg tablet	JF. FIII	Submission rarget. F12014	Olai
\cap	Additional Indication: Regression symptoms in people with	JP: PII		Oral
0	Down syndrome	JF. FII		Ulai

Product Name: Fycompa Research Code: E2007 Generic Name: perampanel (AMPA receptor antagonist)

Description: A selective antagonist against the AMPA receptor (a glutamate receptor subtype). Currently being investigated as a potential adjunctive therapy for partial-onset seizures as well as a treatment for generalized seizures in patients with epilepsy. Approved in 36 countries including in Europe, the United States, and Canada.

Partial-onset seizures	JP/CN/AS: PIII	Submission Target: FY2014	Oral
Additional Indication: Generalized seizures	JP/US/EU/AS: PIII	Submission Target: FY2014	Oral
Additional Indication: Pediatric partial-onset seizures	US/EU: PII		Oral

Research Code: AS-3201 Generic Name: ranirestat (Treatment for diabetic complications / aldose reductase inhibitor)

	Description: An aldose reductase inhibitor that is believed to reduce intracellular accumulation of sorbitol. Currently being investigated		
	as a potential treatment for diabetic neuropathy, one of the most common diabetic complications.		
Diabetic neuropathy US/EU: PII/III			

Product Name: Zonegran Research Code: E2090 Generic Name: zonisamide (Antiepileptic agent)

Description: Believed to exhibit a broad antiepileptic spectrum and is well-tolerated. Currently indicated as an adjunctive therapy and monotherapy for the treatment of partial-onset seizures in adult patients with epilepsy and as an adjunctive therapy for the treatment of partial-onset seizures in pediatric patients with epilepsy.

Image: Second stateAdditional Indication: Pediatric partial-onset seizuresEU: approved (October 2013)Oral

Research Code: E0302 Generic Name: mecobalamin (Amyotrophic lateral sclerosis)

 Description: A mecobalamin (vitamin B₁₂ coenzyme) formulation. Restores damaged peripheral nerves and is widely used for the treatment of peripheral neuropathy. Currently being investigated as a potential treatment for amyotrophic lateral sclerosis (ALS).

 Amyotrophic lateral sclerosis (ALS)
 JP: PII/III
 Submission Target: FY2014
 Inj.

Product Name: Inovelon/Banzel Research Code: E2080 Generic Name: rufinamide (Antiepileptic agent)

Description: A triazole derivative that is structurally unrelated to currently marketed antiepileptic drugs (AEDs). Currently approved in Japan, Europe and the United States as an adjunctive therapy to other AEDs in the treatment of Lennox-Gastaut syndrome (LGS), one of the most severe and intractable forms of childhood-onset epilepsy. The product names are Inovelon in Japan and Europe and Banzel in the United States.

Additional Indication: Pediatric Lennox-Gastaut syndrome (LGS) US/EU: PIII Oral

Research Code: **BAN2401** (Anti-Alzheimer's agent / humanized anti-Aβ protofibrils monoclonal antibody)

$Description: A \ humanized \ IgG1 \ monoclonal \ antibody \ that \ targets \ amyloid \ beta \ (A\beta) \ protofibrils. \ Expected \ to \ be \ effective \ in \ the \ treatment$			
of Alzheimer's disease by halting disease progression through the elimination of Aß protofibrils reported to exhibit neurotoxicity.			
Alzheimer's disease US/EU: PII In			

O Development progress from April 2013 onwards

O Development progress from October 2013 onwards

Research Code: E2006 (Anti-insomnia agent / orexin receptor antagonist)

Description: Anti-insomnia agent with novel mechanism of action. By antagonizing the orexin receptors that maintain wakefulness, it is			
expected to alleviate wakefulness and thereby induce natural sleep.			

O	Insomnia	US: PII	Oral.
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(3) Vascular and Immunological Reaction

Product Name: Humira Research Code: D2E7 Generic Name: adalimumab (Fully human anti-TNFα monoclonal antibody)

Description: A fully human anti-TNF α monoclonal antibody, which neutralizes tumor necrosis factor alpha (TNF α), a type of cytokine that plays a central role in inflammatory reactions in patients with autoimmune diseases. Approved in Japan for the treatment of rheumatoid arthritis, psoriasis, Crohn's disease, ankylosing spondylitis, juvenile idiopathic arthritis, inhibition of structural damage of joints, intestinal Behçet's disease and ulcerative colitis.

0	Additional Indication: Intestinal Behçet's disease	JP: approved (May 2013)	lnj.
0	Additional Indication: Ulcerative colitis	JP: approved (June 2013)	Inj.

Product Name: Tambocor Generic Name: flecainide (anti-tachyarrhythmia agent)

Description: Suppresses tachyarrhythmia by blocking cardiac sodium channels. The agent was approved for the treatment of tachyarrhythmia (paroxysmal atrial fibrillation/flutter and ventricular tachycardia) in adults and tachyarrhythmia (paroxysmal atrial fibrillation/flutter, paroxysmal superventricular tachycardia and ventricular tachycardia) in pediatric patients.

Research Code: E5564 Generic Name: eritoran (Treatment for severe sepsis / endotoxin antagonist)

Description: Exhibits endotoxin antagonist effects that inhibit isolation of inflammatory cytokines. Suppresses various clinical conditions caused by endotoxins.

Severe sepsis

JP/US/EU: PIII

Research Code: E5501/AKR-501 Generic Name: avatrombopag

(Treatment for thrombocytopenia / thrombopoietin receptor agonist)

	scription: A novel, oral thrombopoietin receptor agonist that stin nditions that are associated with thrombocytopenia.	nulates platelet production.	Expected to exhibit effects against
	Idiopathic thrombocytopenic purpura (ITP)	US/EU/AS: PIII	Oral
O	Thrombocytopenia in chronic liver disease requiring surgery	US/EU/AS: PIII	Oral
	Thrombocytopenia during interferon therapy (both initiation and maintenance) for hepatitis C	US: PII	Oral

Research Code: **E6005** (Phosphodiesterase 4 inhibitor)

Description: Inhibits the activity of phosphod	liesterase 4, a cyclic AMP-degrading enzyme that acts a	as an intracellular messenger.
Expected to be effective as a treatment to sup	press the various symptoms associated with atopic diseas	e.
Atopic dermatitis	JP: PII	Topical

• Eisai decided to discontinue the development of the multi-kinase inhibitor E6201, which was in a Phase II study in the U.S. and Europe as a potential treatment for psoriasis.

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(4) Gastrointestinal and Hepatic Disorders

Product Name: Pariet/Aciphex Research Code: E3810 Generic Name: rabeprazole (Proton pump inhibitor)

Description: A proton pump inhibitor approved for the treatment of gastric and duodenal ulcers, reflux esophagitis and eradication of Helicobacter pylori infections, etc. Additional Formulation: Triple formulation pack for Ο JP: approved (August 2013) Oral Helicobacter pylori eradication Additional Indication, Formulation: Prevention of recurrence 0 of gastric and duodenal ulcers during treatment with JP: submitted (November 2013) Oral low-dosage aspirin, new 5 mg tablet Additional Indication: Maintenance therapy for proton pump 0 JP: PIII Oral inhibitor (PPI)-resistant reflux esophagitis Additional Indication: Functional dyspepsia JP: PII Oral

Generic Name: cinitapride (Gastroprokinetic agent)

Description: By stimulating 5-HT₂ and 5-HT₄ receptors found in the gastrointestinal tract, the agent increases acetylcholine release and improves upper gastrointestinal motility. Its antidopaminergic effects also help stimulate the release of acetylcholine by blocking dopamine receptors, thereby improving upper gastrointestinal function.

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• Eisai received a non-approval letter from the Chinese regulatory authority for clevudine as a treatment for patients with chronic hepatitis B. Future development plans are currently under review.

O Development progress from October 2013 onwards