

1st Half of Fiscal 2010 Financial Results

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Forward-Looking Statements

- This presentation contains forward-looking statements. These statements are based on expectations in light of the information currently available, assumptions that are subject to risks and uncertainties which could cause actual results to differ materially from these statements.
- Risks and uncertainties include general domestic and international economic conditions such as general industry and market conditions, and changes of interest rate and currency exchange rate. These risks and uncertainties particularly apply with respect to product-related forward-looking statements. Product risks and uncertainties include, but are not limited to, completion and discontinuation of clinical trials; obtaining regulatory approvals; claims and concerns about product safety and efficacy; technological advances; adverse outcome of important litigation; domestic and foreign healthcare reforms and changes of laws and regulations. Also for existing products, there are manufacturing and marketing risks, which include, but are not limited to, inability to build production capacity to meet demand, unavailability of raw materials and entry of competitive products.
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Overview of 1st Half FY2010 Financial Results

Financial Results (Consolidated)



(Units: billion yen)

<consolidated></consolidated>	1st Half FY2010 Forecasts	1 st Half FY2010 Results	Achievement (%)	1 st Half FY2009 Results	Y on Y Change (%)	Change
Sales	142.8	143.3	100.4	132.6	8.1	10.7
Operating income	19.7	19.2	97.6	17.8	7.8	1.4
Ordinary income	18.7	17.6	94.6	16.3	7.9	1.3
Net income	11.0	6.8	62.4	11.5	(40.8)	(4.7)

Due to change in the accounting periods, 1st half results of FY2010 include 9 months from Jan.1 to Sep. 30 for the US subsidiaries.

Financial Position and Cash Flows (Consolidated)

(Units: billion yen)

<financial position=""></financial>	As of Sep. 30, 2010	As of Mar. 31, 2010	Change
Total assets	516.7	540.7	(24.0)
Net assets	326.3	341.9	(15.6)
Equity ratio (%)	63.1	63.2	(0.1)
Net assets per share (yen)	973	1,019	(46)

(Units: billion yen)

<cash flows=""></cash>	1 st Half FY2010 Results	1st Half FY2009 Results	Change
Net cash provided by operating activities	30.9	20.6	10.3
Net cash provided by investing activities	(16.3)	(5.2)	(11.1)
Net cash provided by financing activities	(13.1)	8.0	(21.1)
Net increase (decrease)	(0.9)	23.7	(24.6)
Cash and cash equivalents at end of period	96.7	75.3	-

Sales by Segments (Consolidated)

S-O-N-G

(Units: billion yen)

	1 st Half FY2010 Forecasts	1 st Half FY2010 Results	Achievement (%)	1 st Half FY2009 Results	Y on Y Change (%)	Y on Y Change
Prescription drugs	74.9	75.1	100.3	74.7	0.6	0.4
Crestor	14.5	13.7	94.4	11.5	18.7	2.2
Irbetan	3.8	3.3	86.9	1.2	170.6	2.1
Cymbalta	0.8	0.9	110.9	-	-	0.9
Total of 3 strategic products	19.1	17.9	93.6	12.8	40.2	5.1
OxyContin	4.7	4.8	102.0	4.2	13.6	0.6
Finibax	2.1	1.8	85.8	1.8	2.8	0.0
Differin	1.9	1.5	76.8	1.0	43.5	0.5
Pirespa	1.4	1.3	89.6	0.6	107.5	0.7
Rapiacta	0.0	0.0	-	-	-	0.0
Total of 8 new products	29.2	27.2	93.1	20.3	33.6	6.9
Flomox	9.0	10.1	112.4	11.3	(10.5)	(1.2)
Rinderon	4.8	5.0	104.9	5.1	(1.9)	(0.1)
Flumarin	3.6	3.9	108.3	4.7	(17.0)	(0.8)
Claritin	3.4	3.1	90.2	3.4	(10.5)	(0.3)
Vancomycin	2.2	2.6	118.4	3.4	(24.3)	(0.8)
Export/Overseas subsidiaries	26.9	26.0	96.8	23.9	9.0	2.1
Shionogi Inc.	22.6	21.0	92.9	18.3	14.3	2.7
Doripenem	1.8	2.5	138.0	2.9	(14.6)	(0.4)
Contract manufacturing	1.5	1.8	121.7	3.4	(46.8)	(1.6)
OTC and quasi-drugs	2.9	2.8	97.9	3.0	(4.0)	(0.2)
Diagnostics	1.3	1.5	112.8	1.6	(6.1)	(0.1)
Royalty income	34.3	34.8	101.7	24.0	45.3	10.8
Crestor	32.5	32.8	100.8	22.5	45.3	10.3
Others	1.0	1.2	118.0	2.1	(43.7)	(0.9)
Total	142.8	143.3	100.4	132.6	8.1	10.7

Statements of Income (Non-Consolidated)



(Units: billion yen)

	1 st Half FY2010 Forecasts	1 st Half FY2010 Results	Achievement (%)	1 st Half FY2009 Results	Y on Y Change (%)	Y on Y Change
Sales	117.9	120.3	102.1	108.3	11.1	12.0
[Royalty income]	[34.3]	[34.8]	[101.7]	[24.0]	[45.3]	[10.8]
Cost of sales	27.7 (39.0)	28.0 (39.4)		29.5 (37.9)		
Cost of sales	32.6	33.7	103.4	32.0	5.3	1.7
Gross profit	85.3	86.6	101.6	76.3	13.5	10.3
SG&A expenses	^{49.7} 58.6	49.7 59.7	102.0	54.9 59.4	0.5	0.3
Selling & general expenses	34.0	33.9	99.7	33.6	0.7	0.3
R&D expenses	24.6	25.8	105.1	25.8	0.2	0.0
Operating income	^{22.6} 26.7	^{22.3} 26.8	100.7	15.6 16.8	59.3	10.0

Small numbers in red are percentages of sales, and numbers in red provided in parentheses are percentages of royalty excluded sales.



(Units: upper/million dollar, lower/billion yen)

Financial Results of US Operations

	1 st Half FY2010 Forecasts*	1 st Half FY2010 Results*	Achievement (%)	Jan.1-Mar.31 FY2010 Results	Apr.1-Jun.30 FY2010 Results	Jul.1-Sep.30 F'Y2010 Results
Sales	248	235	94.8	110	56	69
	22.6	21.0	92.9	10.1	5.1	5.8
Cost of sales	53	56	105.7	18	23	15
	4.8	5.0	104.2	1.6	2.1	1.3
SG&A expenses	222	214	96.4	76	83	54
	20.3	19.1	94.1	7.0	7.6	4.6
Operating income	(27) (2.5)	(35) (3.2)	-	16 1.5	(51) (4.6)	(1) (0.0)
Extra- ordinary loss	L 25 L 2.2	L 72 L 6.5	-	-	L 25 L 2.3	L 47 L 4.2

<Current Situation>

*: 9 months from Jan. 1 to Sep. 30

- Stabilize current P&L through review of product lineup and cost structure
 - > Streamline product portfolio by reviewing product lineup and priority reassessment of clinical development
 - Reductions of field force in Primary Care and associated commercial and marketing personnel
 - **▶** Reductions of administrative and R&D personnel
- Incurred Extraordinary Loss
 - **▶** Personnel reduction expenses
 - ➤ Impairment loss on intangible assets of discontinued products
- 3.3 billion yen (approx.)
- 3.1 billion yen (approx.)

Statements of Income (Consolidated)



(Units: billion yen)

	1 st Half FY2010 Forecasts	1 st Half FY2010 Results	Achievement (%)	1 st Half FY2009 Results	Y on Y Change (%)	Y on Y Change
Sales	142.8	143.3	100.4	132.6	8.1	10.7
[Royalty income]	[34.3]	[34.8]	[101.7]	[24.0]	[45.3]	[10.8]
Cost of sales	26.2 (34.5)	27.2 (36.0)		27.8 (33.9)		
Cost of sales	37.4	39.0	104.4	36.8	6.0	2.2
Gross profit	105.4	104.3	99.0	95.7	8.9	8.6
SG&A expenses	60.0 85.7	59.3 85.0	99.3	58.8 77.9	9.2	7.1
Selling & general expenses	57.4	56.9	99.3	50.0	13.9	6.9
R&D expenses	28.3	28.0	99.3	27.9	0.6	0.1
Operating income	13.8 19.7	13.4 19.2	97.6	13.4 17.8	7.8	1.4

Small numbers in red are percentages of sales, and numbers in red provided in parentheses are percentages of royalty excluded sales.



Review of 1st Flatf FY2010

- **Domestic sales (Prescription Drugs)**
 - Under the influence of average 6.1% NHI drug price reduction in FY2010, the sales of prescription drugs increased compared with previous fiscal year, and achieved 1st half forecast.
 - Three and eight strategic products in all increased by 40.2% and 33.6% in year-on-year term respectively.
- **US Operations**
 - Clarify leader ship in US operations by the establishment of "Shionogi Inc.", and speed up decision-making and achieve efficient operations
 - Stabilize current P&L through review of product lineup and cost structure
- **♦** R&D/Licensing
 - S-349572 (HIV infection): Reported good results in Phase IIb
 - Aggressive licensing activities; agreement with Stallergenes SA about sublingual tablets of house-dust mite allergen extract for immunotherapy, and collaboration with GlaxoSmithKline about novel antibiotics targeting drugresistant Gram-negative bacteria





Revision of FY2010 Financial Forecasts (Consolidated)

(Units: billion yen)

<consolidated></consolidated>	Original Forecasts	Revised Forecasts	Change from Original	FY2009 Results	Y on Y Change (%)	Change
Sales	290.0	283.0	(7.0)	278.5	1.6	4.5
Operating income	57.0	54.0	(3.0)	52.4	3.0	1.6
Ordinary income	55.0	51.5	(3.5)	50.5	1.9	1.0
Net income	36.0	30.0	(6.0)	38.6	(22.3)	(8.6)

Due to change in the accounting periods, FY2010 forecasts include 15 months from Jan. 1, 2010 to Mar. 30, 2011 for the US subsidiaries.

Revision of FY2010 Financial Forecasts (Sales by Segments) (Units: billion yen)

	(Units. binion)					mion yen
	FY2010 Original	FY2010 Revised Forecasts	Change from Original	FY2009 Results	Y on Y Change (%)	Y on Y Change
Prescription drugs	155.2	155.4	0.2	152.5	1.9	2.9
Crestor	30.0	29.2	(0.8)	24.2	20.8	5.0
Irbetan	8.3	7.8	(0.5)	3.8	107.4	4.0
Cymbalta	1.8	2.6	0.8	-	-	2.6
Total of 3 strategic products	40.1	39.6	(0.5)	27.9	41.7	11.7
OxyContin	9.5	9.8	0.3	8.5	15.2	1.3
Finibax	4.3	4.0	(0.3)	3.4	19.1	0.6
Differin	3.9	3.0	(0.9)	2.2	37.8	0.8
Pirespa	3.4	3.0	(0.4)	1.5	97.0	1.5
Rapiacta	3.0	1.8	(1.2)	0.6	227.2	1.2
Total of 8 new products	64.2	61.2	(3.0)	44.1	38.9	17.1
Flomox	19.0	20.1	1.1	24.0	(16.2)	(3.9)
Rinderon	8.9	9.1	0.2	9.5	(3.9)	(0.4)
Claritin	8.1	8.2	0.1	9.0	(9.2)	(0.8)
Flumarin	6.6	6.9	0.3	8.7	(20.7)	(1.8)
Vancomycin	3.9	4.3	0.4	6.1	(30.0)	(1.8)
Export/Overseas subsidiaries	52.1	44.2	(7.9)	49.1	(9.9)	(4.9)
Shionogi Inc.	41.8	33.9	(7.9)	38.6	(12.3)	(4.7)
Doripenem	5.4	5.4	0.0	5.2	3.1	0.2
Contract manufacturing	3.1	3.4	0.3	7.4	(54.2)	(4.0)
OTC and quasi-drugs	5.5	5.5	0.0	5.4	1.1	0.1
Diagnostics	2.8	2.8	0.0	2.9	(4.9)	(0.1)
Royalty income	68.3	68.3	0.0	57.0	19.8	11.3
Crestor	65.5	63.5	(2.0)	50.0	27.0	13.5
Others	3.0	3.4	0.4	4.1	(16.4)	(0.7)
Total	290.0	283.0	(7.0)	278.5	1.6	4.5

Revision of FY2010 Financial Forecasts

(Statements of Income: Non-Consolidated)



(Units: billion yen)

	Original Forecasts	Revised Forecasts	Change from Original	FY2009 Results	Y on Y Change (%)	Change
Sales	243.5	244.0	0.5	228.5	6.7	15.5
[Royalty income]	[68.3]	[68.3]	[0.0]	[57.0]	[19.7]	[11.3]
Cost of sales	27.6 (38.4) 67.3	27.7 (38.5) 67.6	0.3	28.6 (38.2) 65.4	3.2	2.2
Gross profit	176.2	176.4	0.2	163.1	8.2	13.3
SG&A expenses	47.6 115.8	47.5 115.8	0.0	49.8	1.7	2.0
Selling & general expenses	67.8	67.8	0.0	67.0	1.1	0.8
R&D expenses	48.0	48.0	0.0	46.7	2.6	1.3
Operating income	24.8 60.4	24.8 60.6	0.2	21.5 49.2	23.0	11.4

Small numbers in red are percentages of sales, and numbers in red provided in parentheses are percentages of royalty excluded sales.



Next Actions and Forecasts of US Operations (Units: upper/million dollar, lower/billion yen)

	FY2010 Original Forecasts	FY2010 Revised Forecasts	Change from Original
Sales	460	395	(65)
	41.8	33.9	(7.9)
Cost of sales	80	83	3
	7.3	7.2	(0.1)
SG&A expenses	352	327	(25)
	32.0	28.1	(3.9)
Operating income	28	(15)	(43)
	2.5	(1.3)	(3.8)
Extra-	L 25	L 140	(115)
ordinary loss	L 2.2	L 12.0	(9.8)

1 st Half FY2010 Results*	2 nd Half FY2010 Revised Forecasts
235	160
21.0	12.9
56	27
5.0	2.1
214	113
19.1	9.0
(35)	20
(3.2)	1.8
L 72	L 68
L 6.5	L 5.6

[•] Measures to further streamline and stabilize US operations

- Enhance organizational efficiencies by further consolidation of US operations to New Jersey headquarters
- ➤ Improvement of quality processes associated with contract manufacturers
- ➤ Aggressive license activities to strengthen product portfolio
- Anticipation of extraordinary loss
 - **Expenses on integration of business locations**

- 1.4 billion yen (approx.)
- Impairment loss on intangible assets of pre-determinate discontinued products coming in 2H 4.2 billion yen (approx.)
- Accelerate steady growth by speed-up of clinical development and licensing-in

^{*: 9} months from Jan. 1 to Sep. 30

Revision of FY2010 Financial Forecasts

(Statements of Income: Consolidated)



(Units: billion yen)

	Original Forecasts	Revised Forecasts	Change from Original	FY2009 Results	Y on Y Change (%)	Change
Sales	290.0	283.0	(7.0)	278.5	1.6	4.5
[Royalty income]	[68.3]	[68.3]	[0.0]	[57.0]	[19.8]	[11.3]
Cost of sales	26.4 (34.5) 76.5	27.0 (35.5) 76.3	(0.2)	27.4 (34.4) 76.2	0.0	0.1
Gross profit	213.5	206.7	(6.8)	202.2	2.2	4.5
SG&A expenses	54.0 156.5	54.0 152.7	(3.8)	53.8 149.8	1.9	2.9
Selling & general expenses	102.2	99.2	(3.0)	97.9	1.2	1.3
R&D expenses	54.3	53.5	(0.8)	51.8	3.3	1.7
Operating income	19.7 57.0	19.1 54.0	(3.0)	18.8 52.4	3.0	1.6

Small numbers in red are percentages of sales, and numbers in red provided in parentheses are percentages of royalty excluded sales.



Dividend



Dividend Forecast

	Dividends per Share				
	Half-year	Year-end	Annual		
	Yen	Yen (Forecasts)	Yen (Forecasts)		
FY2010	20.00	20.00	40.00		
FY2009	18.00	18.00	36.00		



Progress of Developmental Compounds (since August 2010)

♦ Approval

- RAPIACTA® (Neuraminidase inhibitor, injection):
 - **Influenza Infection for Pediatric Use**
 - ➤ Japan: Approved October 2010
 - ➤ This drug is expected to be effective against seasonal influenza virus infection of child as well as adult with a single-dose administration and improve compliance.
 - ➤ Shionogi has collected date on all patients who received RAPIACTA to delineate its clinical utilization and safety, and submitted a summary of this experience to the MHLW. As it was judged that additional efforts for tracking safety events at this time were not needed, Shionogi has ended the monitoring.
- KAPVAYTM (Clonidine HCL, Alpha 2 specific adrenergic agonist, Oral): ADHD, Pediatric
 - **► USA: Approved September 2010**
 - ➤ KAPVAY is the first and only ADHD treatment approved with two indications, monotherapy and add-on to stimulants.



Progress of Developmental Compounds (since August 2010)

- Change of Phases in Clinical Studies
 - S-811717 (Oxycodone hydrochloride, Injection): the treatment of moderate to severe pain in patients with cancer pain
 - **▶** Japan: NDA filed September 2010
 - S-349572 (Integrase inhibitor, Oral): HIV infection
 - **▶** Global: Phase III
 - S-707106 (Insulin sensitizer, Oral): Type2 diabetes
 - **USA: Phase IIa**

Progress of Developmental Compounds (since August 2010)

- **♦** Newly added development compound
 - S-265744 LAP (Integrase inhibitor, Injection): HIV infection
 - **USA:** Phase I
 - S-524101 (sublingual tablet of house-dust mite allergen extracts for immunotherapy): Allergic rhinitis caused by house-dust mites allergen
 - ➤ Japan: Phase I in preparation
- **♦** Discontinued development
 - Priority reassessment: Pravastatin/Fenofibrate, Jenloga XR
 - Discontinuation of development: S-444823

LAP: Long acting parenteral formulation



Phase III study Design: S/GSK1349572

- **♦** SPRING-2 Study Design
 - 788 treatment-naïve patients
 - S/GSK1349572 50mg (once-daily) vs. Raltegravir (RAL) 400mg (twice-daily)
 - Primary Objective: antiviral activity of S/GSK1349572 compared to RAL over 48-weeks
- **♦ SAILING Study Design**
 - 688 treatment experienced, but integrase inhibitor-naïve
 - S/GSK1349572 50mg (once-daily) vs. RAL 400mg (twice-daily)
 - Primary Objective:
 antiviral efficacy of S/GSK1349572 compared to RAL at 48-weeks



S-O-N-G for youl

License Agreement

- ◆ S-524101 (sublingual tablet of house-dust mite allergen extracts for immunotherapy)
 - Licensed from Stallergenes SA (France)
 - Allergic rhinitis caused by house-dust mites allergen
 - Patients (Japan): over 30 million people
 - Characteristics:
 - ➤ Demonstration efficacy after 4 months of treatment and the persistence of its therapeutic effect after only one year of treatment
 - ➤ More convenient treatment option with much lower risk of systemic adverse reactions
 - Phase I in preparation (Japan)
 - Start to develop sublingual allergen immunotherapy tablet products for Japanese cider pollen induced allergen rhinitis with Stallergenes
- ♦ Novel antibiotics targeting drug-resistant Gram-negative bacteria
 - Collaboration with GlaxoSmithKline focused on research, development and commercialization of novel cephem antibiotics targeting Gram-negative bacteria
 - Activity against a wide range of drug resistant bacteria such as multidrug resistant Acinetobacter spp., Escherichia coli, and Klebsiella pneumoniae, in addition to multidrug resistant Pseudomonas aeruginosa.



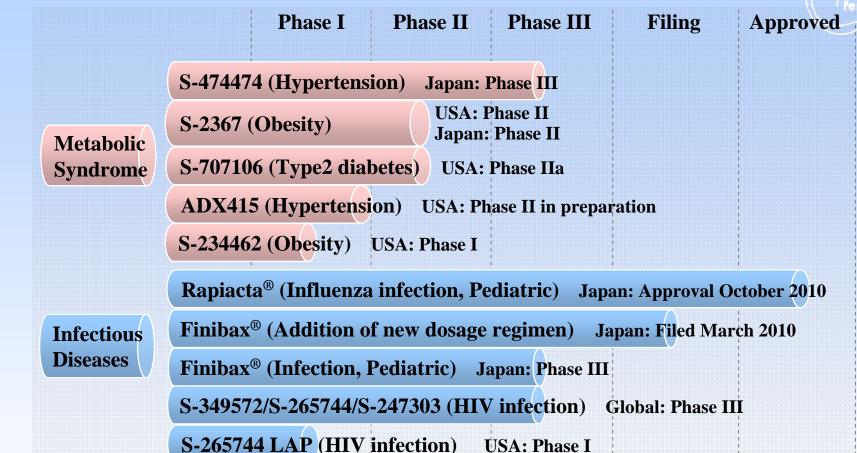


International industry-academic collaboration initiatives

- ◆ Agreement with Sweden on international industry-academic collaboration initiatives
 - Main undertakings by mutual agreed
 - ➤ Matching drug seeds held by Sweden research institutions with drugdiscovery technology and needs of Shionogi and establishing cooperative research relationships with those institutions by embarking on join research for drug seeds which meets demands
 - ➤ Personnel exchanges with Sweden research institutions to facilitate international human resources development
 - Further accelerate the globalization of its activities on international industry-academic collaboration

Pipeline (As of November 2010)

Cymbalta® (DNP)



Pain

S-811717 (Cancer pain) Japan: Filed September 2010

S-297995 (Alleviation of opioid-induced adverse effect) USA: Phase IIa Japan: Phase I

Japan: Filed September 2009



Pipeline (as of November 2010)



Phase I Phase II Phase III **Filing Approved**

Women's Health

PSD502 (Premature ejaculation) USA/EU: Phase III

Ospemifene (Post-menopausal vaginal atrophy) USA: Phase III

Pediatrics

CUVPOSATM (Chronic drooling) USA: Approved July 2010

KAPVAYTM (ADHD) USA: Approved September 2010

S-555739 (Allergic disease)

EU: Proof of Mechanism

Japan: Phase IIa

S-888711 (Thrombocytopenia)

USA/EU: Phase II

Japan: Phase I

Others

S-288310 (Peptide cancer vaccine, Bladder cancer) Japan: Phase I/II

S-222611 (Malignant tumor) EU: Phase Ib

S-488410 (Peptide cancer vaccine, Esophageal cancer) Japan: Phase I/II in preparation

S-524101 (Allergic rhinitis caused by house-dust mite allergen) Japan: Phase I in preparation

<Out-licensed>

Doripenem (Respiratory tract infection) USA: Filed

S-3013 (Acute coronary syndrome) USA/EU: Phase III

S-0373 (Spinocerebellar ataxia) Japan: Phase II



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