# Financial Results for the 1Q of FY2009

(April 1, 2009 - June 30, 2009)

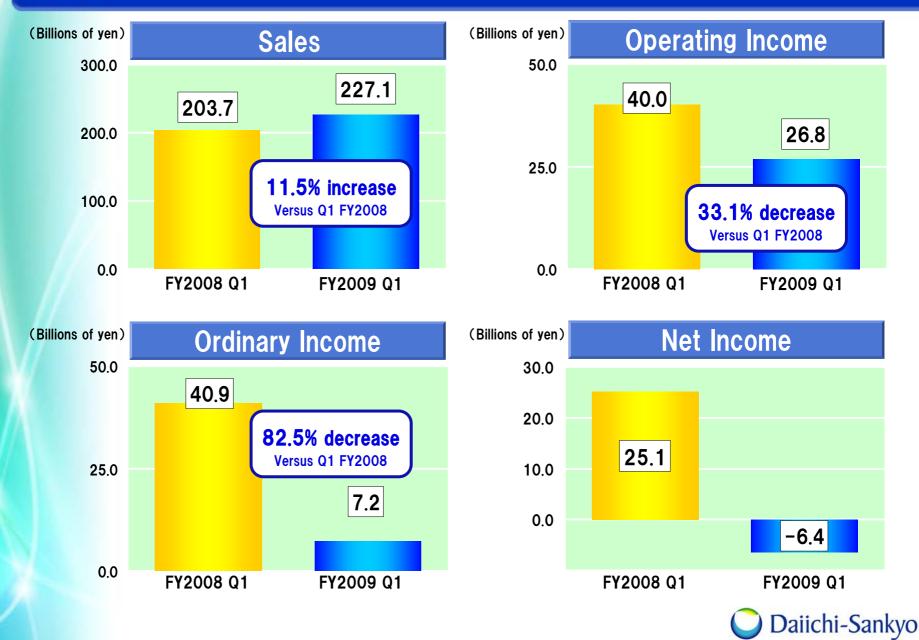
July 31, 2009







### Overview of FY2009 Q1 Results



### Overview of FY2009 Q1 Results - compared with FY2008 Q1 results -

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							(Billions of yen,	allo to salesj
					FY2009 Q	1 Results		
		FY2008 Q1 Results (1)	(2)	Simple Comparison (2)-(1)	Ranbaxy Jan-Mar Results (US GAAP)	Goodwill amortization etc. on Ranbaxy	exc.Ranbaxy <b>(3)</b>	Real term Comparison (3)-(1)
Not	Sales	100.0%	100.0%		100.0%		100.0%	
Net	Sales	203.7	227.1	23.4	29.7		197.4	-6.3
	Cost of Sales	23.9%	27.5%		57.7%		23.0%	
	Cost of Sales	48.6	62.5	13.9	17.1		45.4	-3.3
	SG&A	38.4%	40.8%		41.3%		40.0%	
	expenses	78.2	92.7	14.6	12.3	1.5	79.0	0.8
	R&D	18.1%	19.9%		7.1%		21.8%	
	expenses	36.9	45.1	8.2	2.1		43.0	6.1
	Tetel Francisco	56.5%	60.7%		48.4%		61.8%	
	Total Expense	115.1	137.8	22.8	14.4	1.5	122.0	6.9
On	orating Income	19.6%	11.8%		-6.1%		15.2%	
Ope	erating Income	40.0	26.8	-13.2	-1.8	-1.5	30.0	-10.0
0.0	linaryIncome	20.1%	3.2%		-79.2%		16.3%	
Und	linary Income	40.9	7.2	-33.7	-23.5	-1.5	32.2	-8.7
Net	Income	12.3%	-2.8%		-57.0%		5.1%	
Net	ncome	25.1	-6.4	-31.5	-16.9	0.5	10.0	-15.1



### **Consolidated Year-on-Year Comparisons**

### 1. Net Sales

### ¥227.1 billion (+¥23.4 billion, +11.5%)

Sales from continuing operations (-¥6.3 billion)

●Olmesartan (+ ¥5.3 billion) ●Loxonin (+¥2.8 billion)

●Levofloxacin (-¥6.0 billion) ●Azor (one-time inclusion in 2008 -¥4.7 billion)

Impact of yen appreciation

Ranbaxy (+¥29.7 billion)

Ranbaxy sales by geographic segment:

North America (26%), India (21%), Europe (18%), other regions (35%)

### **2.** Operating Income $\exists 26.8 \text{ billion} (- \exists 13.2 \text{ billion}, -33.1\%)$

Operating income from continuing operations (-¥10.0 billion)

Gross profit decline owing to lower sales (-¥3.0 billion)

SG&A expenses (+¥800 million)

- Expenses higher at Daiichi Sankyo Europe and Luitpold Pharmaceuticals
- U3 Pharma goodwill amortization (+¥1.3 billion) Expenses lower at parent and Daiichi Sankyo, Inc
- •R&D expenses (+¥6.1 billion)
  - Higher spend for projects such as Edoxaban, Denosumab, and Prasugrel
    Rivoglitazone expenses down

### Operating income related to Ranbaxy (-¥3.2 billion)

Ranbaxy operating loss (-¥1.8 billion)

Goodwill and intangible assets amortization (-¥1.5 billion)



### **Consolidated Year-on-Year Comparisons**

### **3.** Ordinary Income \$7.2 billion (-\$33.7 billion, -82.5%)

Non-operating income (+¥200 million)

■Non-operating expenses (+¥20.7 billion)

Ranbaxy-related (+¥22.7 billion)

•Loss on valuation of derivatives (+¥12.9 billion) •Forex loss on loans (+¥8.7 billion)

• Unrelated to Ranbaxy (-¥2.0 billion)

•Forex losses (-¥1.6 billion) •Valuation loss on synthetic stock options (-¥700 million)

### **4.** Net Loss -¥6.4 billion (-¥31.5 billion)

Extraordinary income (+¥2.1 billion) 

 Gain on sales of investment securities (+¥1.8 billion)

 Extraordinary losses (-¥700 million)

### Income Taxes (+¥9.5 billion)

Consolidation adjustment for Ranbaxy

(non-application of tax-effect accounting) (+¥7.0 billion)

- Revision of income taxes for the previous fiscal year (+¥7.5 billion)
- Non-application of R&D tax credits on parent, etc.
- Minority interests in loss (-¥8.9 billion): 63.92% stake in Ranbaxy



### **Overview of FY2009 Forecast**

FY2009 Original

	(Billions of yen, ratio to sales)
FY2009 Latest Forecast	
1st half Forecast	Full year

	Forecast (as of May)			1st half l	Full year				
		1st half (1)	Full year (2)	Q1 Results (3)	Progress (3)/(1)	(4)	(4)-(1)	Forec (5)	cast (5)-(2)
Mat	Calaa	100.0%	100.0%	100.0%		100.0%		100.0%	
Net	Sales	465.0	960.0	227.1	48.8%	465.0	0.0	960.0	0.0
	Cost of Sales	29.0%	29.0%	27.5%		29.0%		29.0%	
	Cost of Sales	135.0	278.0	62.5	46.3%	135.0	0.0	278.0	0.0
	SG&A	42.2%	40.8%	40.8%		42.2%		40.8%	
	expenses	196.0	392.0	92.7	47.3%	196.0	0.0	392.0	0.0
	R&D	20.0%	20.2%	19.9%		20.0%		20.2%	
	expenses	93.0	194.0	45.1	48.5%	93.0	0.0	, 194.0	0.0
	Total Expense	62.2%	61.0%	60.7%		62.2%		61.0%	
	i otai Experise	289.0	586.0	137.8	47.7%	289.0	0.0	586.0	0.0
One	erating Income	8.8%	10.0%	11.8%		8.8%		10.0%	
Obe	rating income	41.0	96.0	26.8	65.3%	41.0	0.0	96.0	0.0
Ord	inary Income	3.7%	7.2%	3.2%		8.0%		7.2%	
	mary meome	17.0	69.0	7.2	42.2%	37.0	20.0	69.0	0.0
Not	Income	1.7%	4.2%	-2.8%		2.4%		4.2%	
Net	lincollie	8.0	40.0	-6.4	-80.5%	11.0	3.0	40.0	0.0

### **Partial Revision of Forecasts**

- **1.** No revisions through operating income level
- 2. Partial revision of H1 forecasts ordinary income (+¥20.0 billion) net income (+¥3.0 billion)
- Main drivers for H1 revisions
  - Ranbaxy has already posted its H1 results (January through June).
    Based on these results, Daiichi Sankyo has revised its Group H1 forecasts
    - to reflect material items from that subsidiary.
      - Ranbaxy's non-operating expenses
        - •¥22.7 billion in Q1  $\rightarrow$  Reduced by appox.¥19.0 billion in Q2
        - Related consolidation adjustments
  - Revision of income taxes for the previous fiscal year

### 3. Maintain full-year forecasts

- Basis for maintaining full-year forecasts unchanged
  - The major driver for the revisions to H1 forecasts the reduction of Ranbaxy's non-operating expenses – may be subject to change to the volatility of the dollar/rupee exchange rates
  - Currently, Daiichi Sankyo does not see any changes to major items large enough to require revisions to our full-year Group forecasts
     Daiichi-Sankyo

### Sales of Major Products (Calculated in Yen)

									(Billions of yen)
				FY2009 Late	est Forecast	(no-change	from origin	al forecast)	
				1s <sup>-</sup>	t half Foreca	ist		Full year	
			Q1 Results					Fore	cast
			(1)	Progress (2)/(1)	Comparison with FY08 Q1	(2)	Comparison with FY08 1H		Comparison with FY08
	Olmesartan	antihypertensive	56.0	49.9%	5.3	112.2	7.6	235.5	24.4
GLOBAL	Levofloxacin	synthetic antibacterial agent	19.8	46.4%	-6.0	42.7	-6.6	92.0	-5.7
GLO	Pravastatin	antihyperlipidemic agent	14.7	50.7%	-1.4	29.0	-2.9	55.0	-5.8
	Prasugrel	antiplatelet * alliance revenue	0.0	-	-	-	-	-	-
	Calblock	antihypertensive	3.5	49.3%	0.4	7.0	1.0	15.0	2.9
	Artist	antihypertensive	6.0	50.3%	0.4	12.0	1.0	24.0	2.1
Japan	Kremezin	treatment for chronic renal failure	3.4	47.9%	0.2	7.0	0.6	14.0	1.2
Jap	Loxonin	anti-inflammatory analgesic	11.2	46.8%	2.8	24.0	5.1	51.0	12.3
	Omnipaque	contrast agent	7.1	49.3%	-0.1	14.5	-0.1	28.0	-0.3
	Urief	treatment for dysuria	2.2	49.5%	0.5	4.5	1.0	10.0	2.1
U.S.	Venofer	treatment for iron deficiency anemia	8.0	59.3%	-0.3	13.5	-3.3	27.5	-4.5
Ū.	Welchol	antihyperlipidemic agent / treatment for type 2 diabetes	6.9	51.0%	0.5	13.5	1.0	28.5	4.0

\* Sales forecast for Prasugrel not disclosed



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### Sales of Major Products (Local Currency Basis)

		FY2009 Latest Forecast (no-change from original forecast)							
				1st half Forecast					
		Q1					Fore	cast	
		Results (1)	Progress (3)/(1)	Comparison with FY08 Q1	(2)	Comparison with FY08 1H		Comparison with FY08	
	Benicar/Benicar HCT	227	50.7%	3	447	19	905	36	
Daiichi Sankyo, Inc. <us> (Mil \$)</us>	Azor	31	45.6%	18	68	33	158	72	
	Welchol	71	49.8%	9	142	24	300	56	
Daiichi Sankyo Europe GmbH	Olmetec/Olmetec Plus	63	44.2%	12	142	27	304	43	
(Mil euro)	Sevikar	7	37.2%	7	18	18	38	22	
Luitpold Pharmaceuticals, Inc. <us> (Mil \$)</us>	Venofer	82	57.9%	3	142	-17	289	-29	



### For Reference: Consolidated Results of Ranbaxy (I GAAP Basis)

#### (Rs. Millions, ratio to sales)

	2009				
Indian GAAP	Jan-Mar Q1	Apr-Jun Q2	Jan-Jun H1	Q2 vs Q1	
Sales	100.0% <b>15,584</b>	100.0% 17,953	100.0% <b>33,537</b>	2,369	
Less:Excise Duty	36	34	70	-2	
Net Sales	15,548	17,919	33,467	2,371	
Other Operating Income	223	873	1,096	650	
Cost of Sales	9,115	11,178	20,293	2,063	
SG&A	5,554	5,987	11,541	433	
R&D Expense	1,079	1,058	2,137	-21	
Forex (Gain)/Loss	845	-716	129	-1,561	
Operating Profit before Interest, Depreciation and Amortization	-5.3% -822	7.2% 1,285	1.4% <b>463</b>	2,107	
Interest	246	197	443	-49	
Depreciation & Amortizatio	639	644	1,283	5	
Forex (Gain)/Loss on Loans	1,273	-1,908	-635	-3,181	
Operating Profit before Tax	-19.1% -2,980	13.1% 2,352	-1.9% -628	5,332	
Interest and Other Income	457	400	857	-57	
Exceptional Items	-9,188	8,067	-1,121	17,255	
Profit before Tax	-75.1% -11,711	<sup>60.3%</sup> 10,819	-2.7% -892	22,530	
Тах	-4,101	3,888	-213	7,989	
Profit after Tax	-48.8% <b>-7,61</b> 0	<sup>38.6%</sup> 6,931	-2.0% -679	14,541	

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# **R&D Pipeline Update**



### **R&D Pipeline (Update from May 2009)**

## Progress on Development Phases

■Effient<sup>™</sup> (US)

Anti-platelet agent ACS-PCI<sup>∗</sup> (Application→Approved)

\*Acute Coronary Syndrome-Percutaneous Coronary Intervention

Loxoprofen sodium hydrate/Gel (JP)

Anti-inflammatory and analgesic (P3 $\rightarrow$ Application)

## Discontinued Projects

Rivoglitazone (CS-011)

Glitazone agent that improves insulin resistance (difficult to sufficiently differentiate from existing product)

### CS-866DM (JP)

Olmesartan/Diabetic nephropathy in patients with type 2 DM (did not satisfy the primary endpoint criteria)



# Effient<sup>TM</sup> – Approved in the U.S. on July 10th

### Indications

Effient is indicated for the reduction of thrombotic cardiovascular events (including stent thrombosis) in patients with acute coronary syndrome (ACS) who are to be managed with PCI (ACS-PCI).

- Before the approval of Effient, there were few options for treating patients with ACS-PCI.
- Some of the ACS patients are non-responders against the current standard of care.

### Boxed warning

Highlights certain patient populations that are at high risk for serious bleeding.

### TRITON-TIMI38

- Treatment with Effient produced a highly significant 19 percent reduction in relative risk for the combined endpoint of cardiovascular death, non-fatal heart attacks or non-fatal stroke compared with Plavix.
- These benefits were accompanied by significantly higher serious bleeding in patients treated with Effient compared with patients treated with Plavix.
- When the efficacy benefits were compared with the risk of bleeding events in the entire patient population, for every 1,000 people treated with Effient compared with Plavix, there were 23 fewer heart attacks and six more major bleeding events.

### ACS in the US -> approx.1.5 mil./year

Launch in early August



### **Development Pipeline**

	Phase 1	Phase 2	Phase 3	Application
Cardiovascular diseases	- DB-772d	- Olmetec/diuretic Combo (#)	- <u>Edoxaban</u> - <u>Prasugrel (ACS-MM)</u> - <u>CS-8635</u>	- Olmetec/Calblock Combo (#)
Glucose metabolic disorders	- CS-1036 (#)			
Infectious diseases			- Levofloxacin inj (#) - Laninamivir (CS-8958)	
Malignant neoplasm	- U3-1287	- Tigatuzumab - Nimotuzumab (#) - ARQ 197 - CS-7017		
Immunological allergic diseases	- CS-0777	- SUN 13834		
Bone / joint diseases			- <u>Denosumab (#)</u>	- Loxonin gel (#)
Others		- Sonazoid additional indication (#) <contrast <br="" for="" lesion="" prostatic="">Contrast for mammary lesion&gt;</contrast>	- Human ghrelin - Memantine (#)	- Feron/Ribavirin combination therapy (#) - Silodosin
Total	4	7	8	4

• Only the most advanced stages are described for the projects under global development

• Prioritized projects are <u>underlined (blue)</u>

• # : Developed only in JPN



### FY2009 Daiichi Sankyo Briefings

	Date (JST)
FY2009 Q1 Financial Results	July 31, 2009
FY2009 Q2 Financial Results	October 30, 2009
R&D Meeting	November or December, 2009
FY2009 Q3 Financial Results	January 29, 2010
2nd mid-term business management plan (2010-2012)	March, 2010







Contact address regarding this material

### DAIICHI SANKYO CO., LTD.

### **Corporate Communications Department**

TEL: +81-3-6225-1126 FAX: +81-3-6225-1132

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