

Results Briefing

Performance Overview

Financial Results for the 3Q of FY2008 (Apr 1, 2008 – Dec 31, 2008)

Jan 30, 2009 Takashi Shoda, President & CEO



Inclusion of Ranbaxy into Daiichi Sankyo consolidated financial results

Transaction Closing Date	2008.11.07
Purchase date of record on	2008.10.01
consolidated accounting basis	(After capital investment
(Inclusion into B/S)	on 2008.10.20)
	B/S: Accounting of Goodwill
Impact on B/S•P/L	P/L: Goodwill Write-down
	B/S•P/L: Goodwill Write-down

Daiichi Sankyo is currently conducting purchase price allocation of our investment in Ranbaxy based on fair value appraisal which will form the basis of the valuations included in our audited full-year FY2008 financial results.

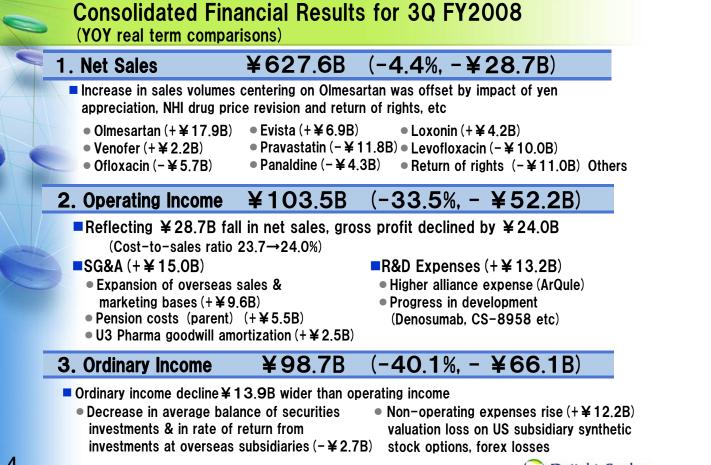


Overview of Consolidated Financial Results for 3Q FY2008 (compared with 30 FY2007)

5									(Billions of yen, r	atio to sales)
	FY2007 3Q YTD Results (Apr-				Dec)	ec)				
			(1)	Non-Pharma Business	Impact of change in the accounting period (EU)	* Special Factors Excluded (2)	(3)	Goodwill Write-down on Ranbaxy	exc. Ranbaxy (4)	Real term Comparison (4)-(2)
	Nat	Calaa	100.0%	100.0%	100.0%	100.0%	100.0%		100.0%	
	Net	Sales	695.8	25.4	14.1	656.3	627.6		627.6	-28.7
		Cost of Color	26.1%	89.2%	25.7%	23.7%	24.0%		24.0%	
		Cost of Sales	181.7	22.7	3.6	155.5	150.8		150.8	-4.7
		SG&A	34.5%	11.5%	58.7%	34.9%	39.8%		38.9%	
		expenses	240.3	2.9	8.3	229.1	249.6	5.5	244.1	15.0
7		R&D	16.8%	2.7%	2.3%	17.7%	20.6%		20.6%	
		expenses	117.0	0.7	0.3	116.0	129.2		129.2	13.2
4		Total Expense	51.4%	14.3%	61.0%	52.6%	60.4%		59.5%	
			357.3	3.6	8.6	345.1	378.8	5.5	373.3	28.2
	0	perating Income	22.5%	-3.4%	13.3%	23.7%	15.6%		16.5%	
	Ope		156.8	-0.9	1.9	155.8	98.1	-5.5	103.5	-52.2
	۲		23.9%	-3.4%	15.4%	25.1%	14.8%		15.7%	
	Ora	inary Income	166.0	-0.9	2.2	164.8	93.2	-5.5	98.7	-66.1
	Not	Income	13.9%	-6.3%	14.3%	14.6%	-47.5%		9.8%	
	Net	income	96.4	-1.6	2.0	96.0	-297.8	-359.5	61.7	-34.3

* Figures of non-pharmaceutical subsidiaries spun off, and the effect of change in the accounting periods of EU subsidiaries are considered as special factors. In order to compare the results in the real term, figures excluding such factors are shown as "Special Factors Excluded."

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Overview of Consolidated Forecasts for FY2008

							(Billions of y	en, ratio to sales;
		FY2008 Full year forecast						
		Revised		Lates	Forecast (2009	9-Jan)		forcast
		Forecast (October) (1)		Ranbaxy Oct-Dec Forcast (US GAAP)	Goodwill Write-down on Ranbaxy	exc. Ranbaxy (2)	Change	(2)—(1)
No	t Sales	100.0%	100.0%	100.0%		100.0%		
ne	l Sales	840.0	850.0	40.0		810.0	-24.0	-30.0
	Cost of Sales	24.4%	25.5%	52.5%		24.2%		
	Cost of Sales	205.0	217.0	21.0		196.0	-7.0	-9.0
	SG&A	39.9%	41.7%	38.8%		40.5%		
	expenses	335.0	354.5	15.5	11.0	328.0	14.5	-7.0
	R&D	21.4%	21.0%	6.3%		21.7%		
	expenses	180.0	178.5	2.5		176.0	13.6	-4.0
		61.3%	62.7%	45.0%		62.2%		
	Total Expense	515.0	533.0	18.0	11.0	504.0	28.2	-11.(
0	orating Incomo	14.3%	11.8%	2.5%		13.6%		
Operating Income		120.0	100.0	1.0	-11.0	110.0	-45.1	-10.0
0.	dinamulnaama	13.5%	8.1%	-62.5%		13.0%		
Un	dinary Income	113.0	69.0	-25.0	-11.0	105.0	-62.1	-8.0
Ne	tincomo	7.7%	-37.2%	-27.5%		7.4%		
Ne	t Income	65.0	-316.0	-11.0	-365.0	60.0	-35.1	-5.0

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* Year on year rates are real term comparison excluding the extra 3 months of EU subsidiaries in FY2007.





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Trend of Major Products

			1						(E	Billions of yen)
			FY2007	FY2008 Forecasts			FY2008 3Q YTD Results (Apr-Dec)			pr-Dec)
			3Q YTD	Revised	Latest				Cha	inge
			Results	Forecast (Oct.)	Forecast	from Oct. Forecast	3Q Results	Proceed	simple comparison	Real term Comparison
Ļ	Olmesartan	antihypertensive	150.7	218.0	212.5	-5.5	162.2	76.3%	11.5	17.9
GLOBAL	Levofloxacin	synthetic antibacterial agent	85.1	102.0	99.5	-2.5	75.1	75.5%	-10.0	-10.0
Ū	Pravastatin	antihyperlipidemic agent	61.9	60.5	60.5	0.0	48.5	80.2%	-13.4	-11.8
	Calblock	antihypertensive	7.8	14.0	13.0	-1.0	9.4	72.1%	1.5	1.5
	Artist	antihypertensive	16.5	22.0	23.0	1.0	17.0	73.9%	0.5	0.5
Japan	Kremezin	treatment for chronic renal failure	9.7	14.0	14.0	0.0	9.9	70.9%	0.2	0.2
Jac	Loxonin	anti-inflammatory analgesic	26.2	40.0	41.0	1.0	30.4	74.1%	4.2	4.2
	Omnipaque	contrast agent	25.1	29.0	29.0	0.0	22.6	77.9%	-2.6	-2.6
	Urief	treatment for dysuria	4.0	8.5	8.5	0.0	5.7	66.9%	1.7	1.7
, S	Venofer	treatment for iron deficiency anemia	23.0	30.5	31.5	1.0	25.2	79.9%	2.2	2.2
S.U	Welchol	antihyperlipidemic agent / treatment for type 2 diabetes	17.6	26.0	25.0	-1.0	19.2	76.7%	1.5	1.5

* Accounting period of European subsidiaries in FY2007 3Q were 12 months from Jan-07 to Dec-07 following the change in fiscal year-end.

Figures excluding the extra three months sales are shown for Real term Comparison.



Trend of Major Products (Local Currency Basis)

			FY2007	FY200	8 3Q YTD F	lesults
			3Q YTD	3Q YTD	Cha	nge
-			Results	Results	Real term Comparison	Real term Comparison
		Benicar/Benicar HCT	585	659	74	12.7%
T	Daiichi Sankyo, Inc. <us> (Mil \$)</us>	Azor	16	62	46	288.0%
1		Welchol	150	186	36	24.0%
	Daiichi Sankyo Europe GmbH	Olmetec/Olmetec Plus	190	184	35	23.3%
	(Mil euro)	Sevikar	0	9	9	-
	Luitpold Pharmaceuticals, Inc. <us> (Mil \$)</us>	Venofer	196	245	49	25.0%

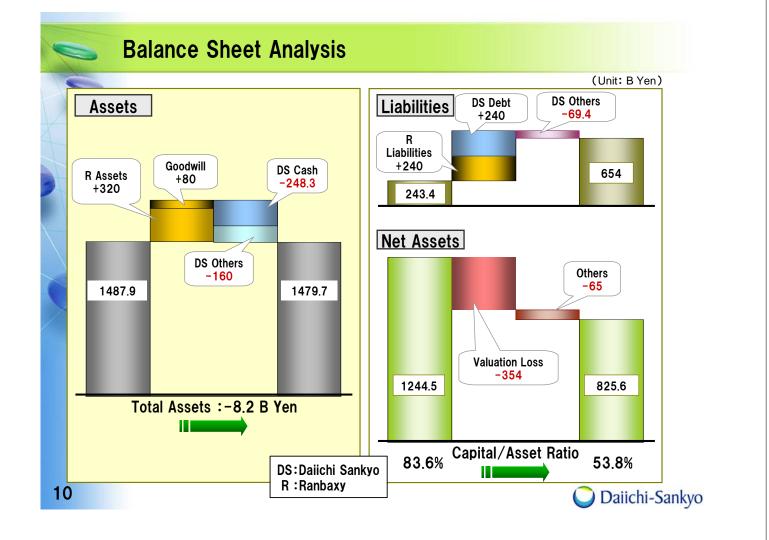
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For Reference: Ranbaxy Results (Based on Indian GAAP)

Zesults idited Jan-Dec '07 100.0% 1 1,61 14.9%	non- Oct-Dec 100.0%	Results audited Jan-Dec '08 100.0% 1,667 -19.2%
'07 100.0% 1 1,61	100.0%	'08 100.0% ' 1,667
1 1,61	19 387	1,667
14.9%	-62.5%	-19.2%
		-13.270
9 24	2 -242	-320
6 11	l2 <mark>41</mark>	131
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R&D Pipeline (Update from October 2008)

New additions ARQ 197

► c-Met Inhibitor (P2)

Progress on Development Phases

Sevikar (Eur) ▶ Olmesartan/Amlodipine combination (Submission→launched) CS-866AZ (Japan) ► Olmesartan/Azelnidipine combination (P3→Submission) Silodosin (China) ► Treatment for dysuria (P3→Submission) DU-176b (Global) ▶ Blood coagulation factor Xa inhibitor ($P2 \rightarrow P3$) CS-8958 (Japan) ► Anti-influenza (P2→P3) CS-7017 (US) ► Anti-tumor PPAR gamma activator (P1→P2) Act-Hib (Japan) ► Act-Hib (Approval→launched) Kuban (Eur) ► Sapropterin (BioMarine: Submission → Approval) Discontinued Projects CS-8080 Treatment for arteriosclerotic disease (did not meet the expected criteria in clinical trial stage) CS-088 Olmesartan glaucoma (did not meet the expected criteria in clinical trial stage)

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Prioritized R&D project 1) Effient/Efient

(Acute Coronary Syndrome-Percutaneous Coronary Intervention)

<u>USA</u> Feb-3th, 2009

Cardiovascular and Renal Drug Advisory Committee Meeting

Europe Dec-18th, 2008

CHMP (European Committee for Medicinal Products for Human Use) recommend approval of Prasugrel NDA

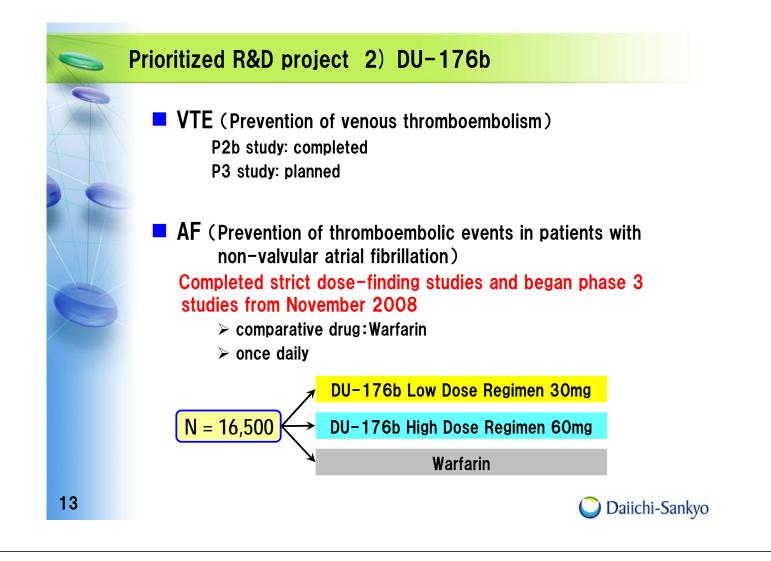
ACS-MM

(Medically Managed without a planned artery-opening procedure)

June 2008 Start of Phase3



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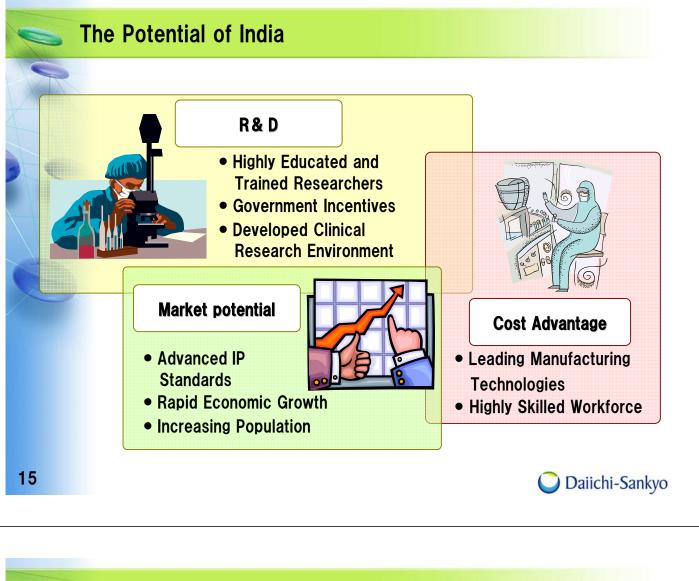
	Daiichi Sankyo R&D Pipeline						
R		Phase 1	Phase 2	Phase 3	Application		
	Cardiovascular diseases	- DB-772d	- Olmetec/diuretic Combo (#)	 <u>DU-176b</u> <u>Prasugrel (ACS-MM)</u> <u>CS-8635</u> Olmetec additional indication (#) <diabetic nephropathy=""></diabetic> 	- <u>Prasugrel (ACS-PCI)</u> - Olmetec/Calblock Combo (#)		
	Glucose metabolic disorders	- CS-1036 (#)		- Rivoglitazone			
	Infectious diseases			- Levofloxacin inj (#) - CS-8958	- Levofloxacin high-dose (#)		
	Malignant neoplasm	- U3-1287	- Nimotuzumab (#) - CS-1008 - ARQ 197 - CS-7017				
-	Immunological allergic diseases	- CS-0777	- SUN 13834				
	Bone / joint diseases			- <u>Denosumab (#)</u> - Loxonin gel (#)			
	Others		- Human ghrelin	- Memantine hydrochloride (#)	- Feron/Ribavirin combination therapy (#) - Silodosin		
	Total	4	7	10	5		

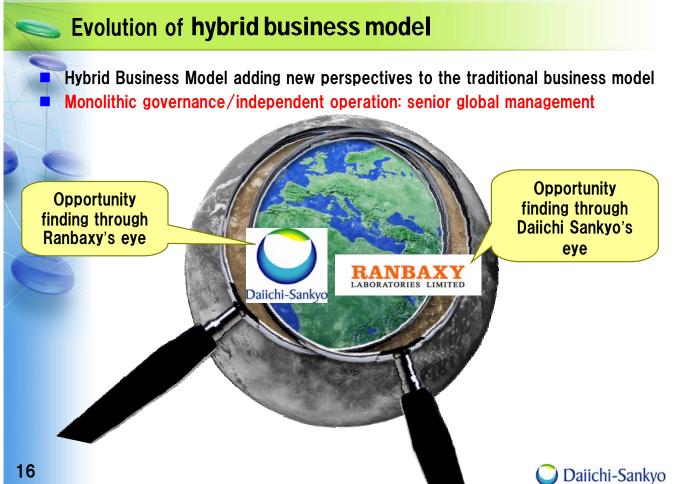
• Only the most advanced stages are described for the projects under global development

• Projects with highest priority are underlined (blue)

• # : Developed only in JPN







	Date (JST)
R&D Meeting	March 31, 2009 14:00-16:00 at Keidanren Kaikan
Financial results for FY2008	Filing to TSE: May 12, 13:00 Meeting for Media: May 13, 2009, 13:00 Meeting for Investor/analyst: May 13, 2009, 15:00





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