

Financial Results for the 3Q of FY2007

(April 1, 2007 - December 31, 2007)

January 31, 2008





Overview of Consolidated Financial Results for 3Q FY2007 (compared with 3Q FY2006 ①)

		FY2006	3Q YTD F	Results (A	Apr-Dec)	FY2007 3Q YTD Results (Apr-Dec) Change						
		1	Non-Pharma Business	Change in the accounting period (U.S.)	* Special Factors Excluded ②	3	Non-Pharma Business	Change in the accounting period (EU)	* Special Factors Excluded 4	Simple Comparison 3-1	Real term Comparison 4-2	
	Net Sales	730.6	77.6	31.5	621.6	695.8	25.4	14.1	656.3	-34.8	34.8	
	Cost of Sales	208.4	50.4	5.1	152.9	181.7	22.7	3.6	155.5	-26.6	2.6	
	SG&A Expenses	265.0	17.5	15.6	231.9	240.3	2.9	8.3	229.1	-24.6	-2.8	
	R&D Expenses	125.3	4.3	1.8	119.2	117.0 0.7		0.3	116.0	-8.3	-3.2	
	Total Expense	390.3	21.7	17.4	351.1	357.3	3.6	8.6	345.1	-33.0	-6.0	
Oı	perating Income	132.0	5.4	9.0	117.6	156.8	-0.9	1.9	155.8	24.8	38.2	
	Non-operating income (expenses)	14.1	0.1	1.5	12.5	9.3	0.0	0.3	9.0	-4.9	-3.5	
0	rdinary Income	146.1	5.5	10.6	130.0	166.0	-0.9	2.2	164.8	19.9	34.7	
	Extraordinary gains (losses)	-20.4	45.0	-1.0	-64.4	-4.9	1.6	0.0	-6.5	15.6	57.9	
	Income tax and Minority interests	48.0	19.8	3.8	24.4	64.8	2.4	0.1	62.3	16.8	37.8	
	Net Income	77.7	30.6	5.8	41.2	96.4	-1.6	2.0	96.0	18.7	54.8	

^{*} Figures of non-pharmaceutical subsidiaries to be spun off, and the effect of change in the accounting periods of U.S./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."





Overview of Consolidated Financial Results for 3Q FY2007 (compared with 3Q FY2006 ②)

;	Special Factors <u>Excluded</u>	FY2006 3Q YTD (Apr-Dec)	FY2007 3Q YTD (Apr-Dec)	Real term change	Remarks
	Net Sales	621.6	656.3	34.8	Olmesartan +36.2 (including 1.9 from AZOR), Levofloxacin +3.8 Welchol +3.6, Urief +2.3, Loxionin +2.3, Pravastatin -16.2
	Cost of Sales	152.9	155.5	2.6	Cost-to-sales ratio 24.6→23.7% Product mix improvement
	SG&A Expenses	231.9	229.1	-2.8	
	R&D Expenses	119.2	116.0	-3.2	R&D expenses ratio 19.2→17.7%
	Total Expense	351.1	345.1	-6.0	Decrease in personnel cost due to personnel downsizing in domestic companies
Ol	perating Income	117.6	155.8	38.2	Operating income to sales 18.9→23.7%
	Non-operating income (expenses)	12.5	9.0	-3.5	
O	ordinary Income	130.0	164.8	34.7	Ordinary income to sales 20.9→25.1%
	Extraordinary gains (losses)	-64.4	-6.5	57.9	gains -4.6 (sales of property, plant and equipment -1.6, sale of investment securities -2.0) losses -62.5 (loss on business integration -54.7, loss on business restructuring -7.7)
	Income tax and Minority interests	24.4	62.3	37.8	Increase in income before taxes and minority interests
	Net Income	41.2	96.0	54.8	Net income to sales 6.6→14.6%





Overview of Consolidated Forecasts and 3Q Progress

		FY2007 Forecasts (no change from disclosure in Nov 2007)				FY2007 3Q YTD Results (Apr-Dec)				
			Non- Pharma Business	Change in the accounting period (EU)	* Special Factors Excluded ①		Non- Pharma Business	Change in the accounting period (EU)	* Special Factors Excluded ②	Progress ②/①
	Net Sales	876.0	16.4	14.1	845.5	695.8	25.4	14.1	656.3	77.6%
	Cost of Sales	221.0	14.6	3.6	202.8	181.7	22.7	3.6	155.5	76.7%
	SG&A Expenses	323.5	2.0	8.3	313.2	240.3	2.9	8.3	229.1	73.1%
	R&D Expenses	171.5	0.5	0.3	170.7	117.0	0.7	0.3	116.0	68.0%
	Total Expense	495.0	2.5	8.6	483.9	357.3	3.6	8.6	345.1	71.3%
Op	perating Income	160.0	-0.7	1.9	158.8	156.8	-0.9	1.9	155.8	98.1%
	Non-operating income (expenses)	11.0	0.0	0.3	10.7	9.3	0.0	0.3	9.0	
Ordinary Income		171.0	-0.7	2.2	169.5	166.0	-0.9	2.2	164.8	97.2%
	Extraordinary gains (losses)	-5.0	1.6	0.0	-6.6	-4.9	1.6	0.0	-6.5	
	Income tax and Minority interests	66.0	0.0	0.1	65.9	64.8	2.4	0.1	62.3	
	Net Income	100.0	0.9	2.0	97.0	96.4	-1.6	2.0	96.0	98.9%





Trend of Major Products

		FY 2006	FY 2006 FY2007 Forecasts		FY2007 3Q YTD Results (Apr-Dec)				
Product name			3Q YTD Results	(no change from disclosure in	3Q Results	Proceed	Over the previous year		
			1	Nov 2007)	2		2-1	Ref.*	
AL	Olmesartan	<antihypertensive></antihypertensive>	123.7	202.0	150.7	74.6%	26.9	36.2	
OB	Levofloxacir	synthetic antibacterial agent>	81.3	110.0	85.1	77.3%	3.8		
G	Pravastatin	<antihyperlipidemic agent=""></antihyperlipidemic>	76.6	79.0	61.9	78.4%	-14.6	-16.2	
	Calblock	<antihypertensive></antihypertensive>	6.9	11.5	7.8	68.2%	1.0		
	Artist	<antihypertensive></antihypertensive>	15.4	22.0	16.5	75.2%	1.2		
pan	Kremezin	<pre><treatment chronic="" failure="" for="" renal=""></treatment></pre>	9.5	13.0	9.7	74.7%	0.2		
Jak	Loxonin	<pre><non-steroidal agent="" analgesic="" and="" anti-inflammatory=""></non-steroidal></pre>	23.9	35.0	26.2	74.8%	2.3		
	Omnipaque	<contrast agent=""></contrast>	26.2	32.0	25.1	78.6%	-1.1		
	Urief	<treatment dysuria="" for=""></treatment>	1.7	7.5	4.0	53.7%	2.3		
S.	Venofer	<pre><treatment anemia="" deficiency="" for="" iron=""></treatment></pre>	29.1	28.5	23.0	80.6%	-6.1	0.7	
j	Welchol	<pre><antihyperlipidemic 2="" agent="" diabetes="" for="" treatment="" type=""></antihyperlipidemic></pre>	17.9	24.0	17.6	73.4%	-0.3	3.6	

^{*} Accounting periods of U.S. subsidiaries in FY2006 was15 months from January 2006 to March 2007 following a change in fiscal year-end.

Accounting periods of European subsidiaries will be 15 months in FY2007. Figures excluding the extra 3 months sales are shown for reference.





R&D Pipeline (Change from disclosure in November 2007)

Development Code	Before Change	After Change	Remarks		
CS-747 (Prasugrel)	U.S.: Phase3	U.S.: Application	Anti-platelet agent		
CS-8080	-	U.S./EU: Phase1	Treatment for arteriosclerosis		
Welchol DM	U.S.: Application	U.S.: Approval	Treatment for type 2 diabetes		
DU-6859a (Gracevit)	Japan: Application	Japan: Approval	New quinolone		
Levofloxacin High-dose	Japan: Phase3	Japan: Application	New quinolone		
Levofloxacin Injection	Japan: Phase2	Japan: Phase3	New quinolone		
DX-619	U.S./EU/JP: Phase1	-	New quinolone Withdrawal from project based on R&D portfolio		
CS-758	U.S./EU: Phase1	-	Azole antifungal Withdrawal from project based on R&D portfolio		
SUN 0588r	U.S.: Application	U.S.: Approval	Treatment for Hyperphenylalaninemia		
SUN 03001	-	EU: Application	Out-licensed to BioMarin		



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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