



Solutions with you in mind

Q3 2010 Financial Results

November 15th, 2010

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Summary – Q3 2010 Financial Results

- Sales (-3.5%) and Normalized Net Income (-4.0%) within guidance.
- Cost discipline and savings a continued priority for 2010 (SG&A: -6.2% YTD).
- Net Debt remains low at x 0.12 EBITDA 2009 and provides strong strategic flexibility.
- Healthy Free Cash Flow generation.

€rounded million	YTD Sep 2010	YTD Sep 2009	% var
Net Sales	676,5	701,3	(3,5%)
EBIT	146,8	162,4	(9,6%)
EBITDA	192,8	210,3	(8,3%)
Normalized Net Income	121,9	127,0	(4,0%)

Income Statement

€rounded million	YTD Sep 2010	YTD Sep 2009	% var
Net Sales	676,5	701,3	(3,5%)
Gross Profit	417,0	443,3	(5,9%)
% of sales	61,6%	63,2%	
Other Income	89,7	77,3	16.0%
R&D	(102,4)	(81,5)	25,6%
% of sales	(15,1%)	(11,6%)	
SG&A	(259,0)	(276,2)	(6,2%)
% of sales	(38,3%)	(39,4%)	
Other Op. Exp	1,5	(0,5)	n.m.
% of sales	0,2%	(0,1%)	
EBIT	146,8	162,4	(9,6%)
% of sales	21,7%	23,2%	
Depreciation	46,0	47,9	(4,0%)
% of sales	6,8%	6,8%	
EBITDA	192,8	210,3	(8,3%)
% of sales	28,5%	30,0%	
Sale of noncurrent assets / Other	0,0	20,5	(100,0%)
Impairment reversals / (losses)	(1,0)	4,0	(125,0%)
Net financial income / (expenses)	(9,4)	(14,8)	(36,5%)
Corporate income tax	(15,4)	(31,4)	(51,0%)
Net income	121,0	140,7	(14,0%)
Normalized Net Income	121,9	127,0	(4,0%)
Earnings per share (€) ⁽¹⁾	0,73 €	0,85 €	
Normalized Earnings per share (€) ⁽¹⁾	0,73€	0,76 €	
Nu. of employees end of period	3.022	3.243	(6,8%)

Sales eroded by Spanish reforms and generic competition.

Driven by higher Eklira[®] and LAS100977 (OD LABA) development expenses.

Cost discipline and savings a key priority in 2010.

Lower EBIT and EBITDA following Gross Margin evolution and higher R&D.

Lower performance driven by € 20 mill. extraordinary item in 2009.

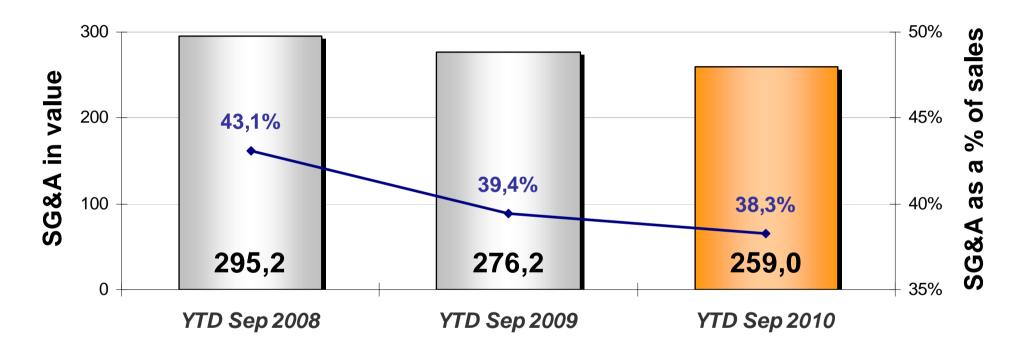
In line with guidance.

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(1) Number of shares at the end of the period

SG&A evolution Consistent improvement over time

(€ rounded million)





Q3 2010 vs Q3 2009

	2010	2009	0/	
€rounded million	Q3	Q3	% var.	
Net Sales	207,5	212,5	(2,4%)	
Gross Profit	124,6	128,5	(3,0%)	
% of sales	60,0%	60,5%		
Other Income	28,6	23,0	24,3%	
R&D	(33,0)	(25,8)	27,9%	
% of sales	(15,9%)	(12,1%)		
SG&A	(77,6)	(75,6)	2,6%	
% of sales	(37,4%)	(35,6%)		
Other Op. Exp	(0,5)	0,5		
% of sales	(0,2%)	0,2%		
EBIT	42,1	50,6	(16,8%)	
% of sales	20,3%	23,8%		
Depreciation	15,4	16,1	(4,3%)	
% of sales	7,4%	7,6%		
EBITDA	57,5	66,7	(13,8%)	
% of sales	27,7%	31,4%		
Sale of noncurrent assets / Other	0,1	0,4	(75,0%)	
Net financial income / (expenses)	(6,4)	(6,0)	6,7%	
Тах	(1,7)	(7,7)	(77,9%)	
Net income	34,1	37,3	(8,6%)	
Normalized Net Income	34,5	37,9	(9,0%)	

Key takeaways

- Q3 Sales in 2010 slightly below 2009.
- Gross Margin trends expected to improve in Q4.
- R&D trends expected to persist through Q4.

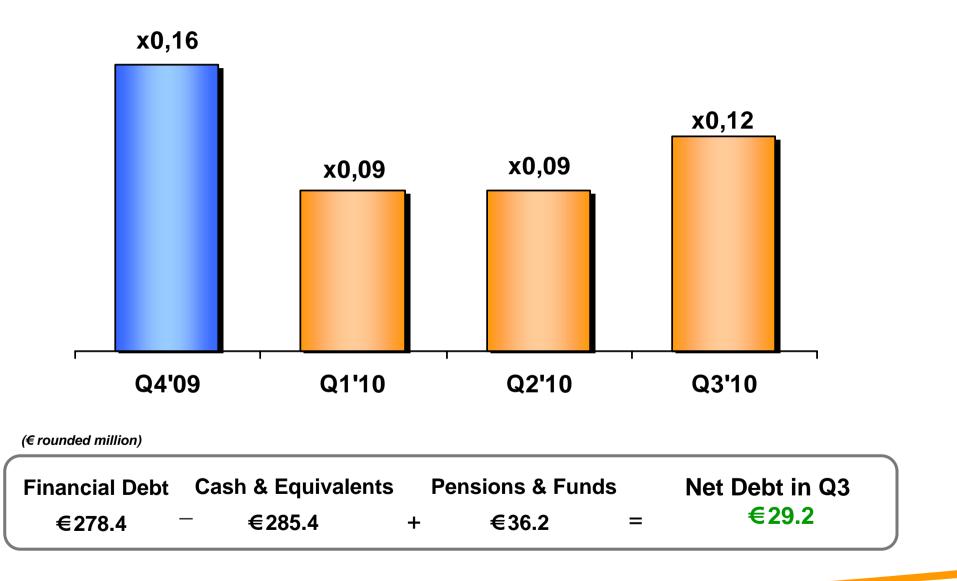
Balance Sheet

€rounded million	September 2010	% of BS	December 2009
Goodwill	272,1	18,1%	272,7
Intangible assets	378,3	25,2%	352,8
Property, plant and equipment	157,0	10,5%	169,1
Financial assets	10,0	0,7%	10,8
Other non current assets	179,4	11,9%	173,6
Total Non Current Assets	996,8	66,4%	979,0
Inventories	97,4	6,5%	97,7
Accounts receivables	109,9	7,3%	120,4
Cash & equivalents	285,4	19,0%	259,7
Other current assets	12,6	0,8%	26,2
Total Current Assets	505,3	33,6%	504,0
Total Assets	1.502,1		1.483,0
Shareholders equity	819,4	54,6%	751,0
Financial debt	278,4	18,5%	265,7
Non current liabilities	210,5	14,0%	228,4
Current liabilities	193,8	12,9%	237,9
Total Equity and Liabilities	1.502,1		1.483,0

Relevant improvement of accounts receivables.

Equity represents nearly 55% of Total Assets.

Net Debt vs EBITDA 2009



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Cash Flow Statement

€rounded million	YTD	YTD	
	Sep 2010		
Profit Before Tax	136,4	172,0	
Depreciation and amortisation	46,0	47,9	
Change in working capital	(1,8)	(49,8)	
Other adjustments	(44,6)	(11,2)	
Cash Flow from Operating Activities (I)	136,0	158,9	
Financial Income	2,3	1,8	
Investments	(60,4)	(55,0)	
Divestments	0,7	19,4	
Other cash flows	0,5	(1,1)	
Cash Flow from Investing Activities (II)	(56,9)	(34,9)	
Finance Expense	(12,7)	(14,8)	
Dividends distribution	(55,1)	(52,5)	
Debt increase/ (decrease)	15,2	(28,6)	
Other cash flows	(0,7)	(4,1)	
Cash Flow from Financing Activities	(53,3)	(100,0)	
Cash Flow generated during the period	25,8	24,0	
Free Cash Flow (III) = (I) + (II)	79,1	124,0	

Improvement of inventories and accounts receivables vs last year.

Driven by corporate income tax calendar payments.

Includes Sativex[®] and linaclotide downpayments.

Includes Toctino[®] and Meda payments.

Includes debt cancellation in Q1'10 and slight debt increase seen in Balance Sheet.

Net Sales by Business and Region

Net Sales breakdown by Business Model

€rounded million	YTD	YTD	% var
	Sep 2010	Sep 2009	70 Val
Own sales network (within Spain)	383,0	401,4	(4,6%)
Own sales network (other countries)	218,5	214,5	1,9%
Marketing with licensees	52,5	57,7	(9,1%)
Corporate	22,5	27,6	(18,5%)
Total	676,5	701,3	(3,5%)

Net Sales breakdown by Geographic Area

€rounded million	YTD	YTD	% var	
	Sep 2010	Sep 2009	70 Val	
Spain	383,0	401,4	(4,6%)	
Europe & Middle East	218,9	216,9	0,9%	
America, Africa & Asia Pacific	52,1	55,2	(5,9%)	
Corporate	22,5	27,6	(18,5%)	
Total	676,5	701,3	(3,5%)	

 Spanish sales eroded due to healthcare reforms and generics. Underlying sales are +2% excluding Prevencor[®].

• Steady growth in our affiliates (+1.9%) led by Mexico, UK, France and Germany.

• Gradual reduction of toll manufacturing business drives Corporate sales evolution.

America, Africa & Asia-Pacific influenced by lower ebastine sales in Japan.

Highlights

Net Sales breakdown by main Therapeutic Area

€rounded million	YTD Sep 2010	YTD Sep 2009	% var
Respiratory	146,0	146,3	(0,2%)
CNS	127,0	126,8	0,1%
Cardiovascular	122,9	142,7	(13,9%)
Gastrointestinal	117,5	106,2	10,7%
Dermatology	92,0	86,2	6,8%
Osteomuscular	48,5	53,7	(9,6%)
Urological	12,9	14,9	(13,6%)
Other ther. specialties	9,7	24,5	(60,5%)
Total	676,5	701,3	(3,5%)

Highlights

- Positive evolution in gastrointestinal and dermatology.
- Cardiovascular lost *momentum* driven by trends in Prevencor[®].



Breakdown of the core business

€rounded million	YTD	YTD	%	Prese	ence
	Sep 2010	Sep 2009	Variation	Spain	Intl.
Ebastel [®] and others <i>(ebastine)</i>	93,3	94,2	(0,9%)	\checkmark	\checkmark
Prevencor [®] (atorvastatin)	62,4	87,1	(28,4%)	\checkmark	
Esertia [®] <i>(escitalopram)</i>	50,4	48,1	4,9%	\checkmark	
Plusvent [®] (salmeterol & fluticasone)	45,4	44,4	2,3%	\checkmark	
Almogran [®] (<i>almotriptan</i>)	38,6	39,6	(2,6%)	\checkmark	\checkmark
Parapres [®] (candesartan cilexetile)	35,7	32,3	10,7%	\checkmark	
Airtal [®] and others (aceclofenac)	29,5	32,7	(9,6%)	\checkmark	\checkmark
Opiren [®] (lansoprazole)	25,8	26,1	(1,0%)	\checkmark	
Dobupal [®] (venlafaxine)	23,8	26,0	(8,6%)	\checkmark	
Solaraze [®] (diclofenac sodium)	18,6	17,1	9,2%		\checkmark
Tesavel [®] (sitagliptin) + Efficib [®] (sitagliptin+metformin)	17,6	6,3	177,7%	\checkmark	
Almax [®] (almagate)	16,4	16,3	0,8%	\checkmark	\checkmark
Balneum [®] <i>(soya oil)</i>	13,8	13,0	5,6%		\checkmark
Pantopan [®] (pantoprazole)	13,6	15,4	(11,6%)		\checkmark
Decoderm Tri [®] (flupredniden)	11,4	10,1	11,9%		\checkmark
Other	180,2	192,7	(6,5%)	\checkmark	\checkmark
Total	676,5	701,3	(3,5%)		



First-in-class endocannabinoid system modulator for treating resistant spasticity in multiple sclerosis (MS)



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- Launched in the Canada and UK, approved in Spain (currently under price and reimbursement process), MRP* ongoing in several EU countries.
- Almirall holds pan-European rights (ex-UK).
- Its launch will provide the first new therapeutic solution for the treatment of this symptom in over 10 years.
- Spasticity (muscle rigidity or stiffness, with or without spasms) is one of the most common symptoms of MS, affecting approx. 80% of the people with MS**.
- There are approximately 670,000 people with MS in Europe of those 500,000 in the Top-5 EU countries and 1/3 of them suffer moderate to severe spasticity***.
- Sativex[®] aims to treat (high need) patients who have previously failed to gain adequate benefit from currently available anti-spasticity treatments.

2010 Launches update

Strengthening the core business

	Silodyx®	Conbriza®	Toctino®	Sativex®
Compound	Silodosin	Bazedoxifene	Alitretinoin	Tetrahydro- cannabinol / cannabidiol
Indication	Benign prostatic hyperplasia (BPH)	Osteoporosis	Severe chronic hand eczema (CHE)	
Almirall's Commercial Rights	Spain	Spain	Austria, Belgium, Czech Republic, Italy, Luxembourg, Mexico, the Netherlands, Poland, Portugal, Slovakia and Spain	
Update	Launched in Q3	Launched in Q3	Launched in Austria Q3, Italy Q4	Approved in Spain (under P&R*). MRP** ongoing in other European countries.

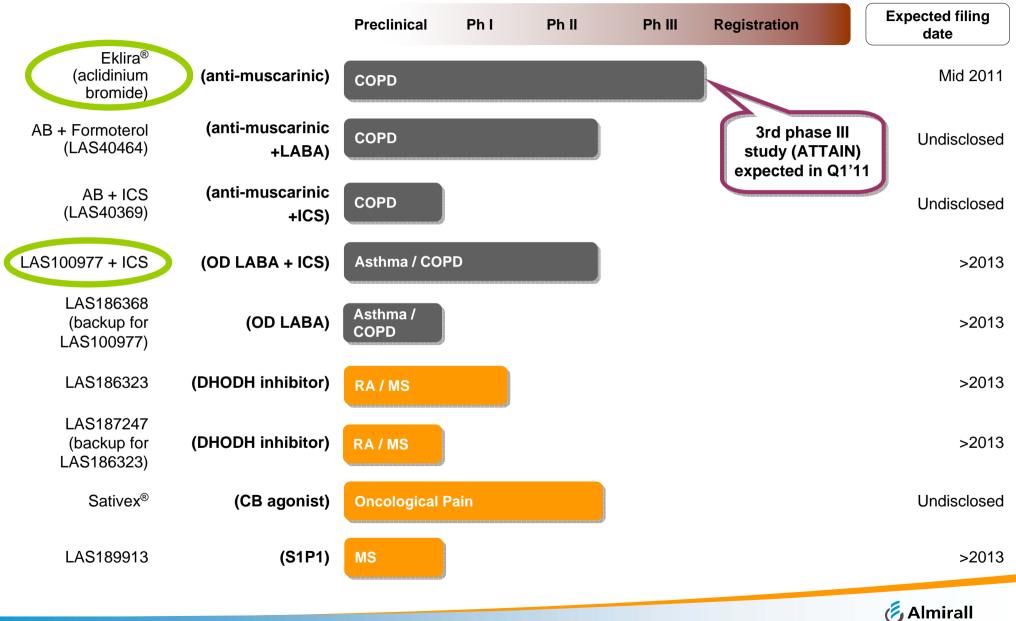
* Price and reimbursement.

** Mutual Recognition Procedure.



A pipeline with significant upside (I)

(the right end of each bar represents status of development as of November 15th 2010)



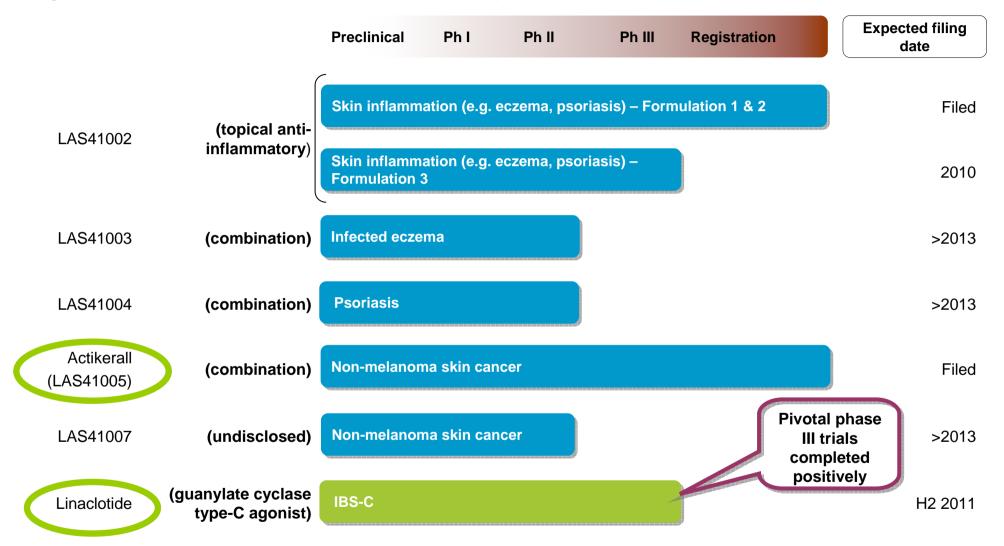
Respiratory Auto-immune

A pipeline with significant upside (II)

Dermatology
Gastrointestinal

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(the right end of each bar represents status of development as of November 15th 2010)



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Actikerall[®] (LAS41005)

Cutaneous solution with 0.5% 5-Fluorouracil and 10% Salicylic Acid

- Once-daily topical treatment.
- A new combination treatment for actinic keratoses (non-melanoma skin cancer).
- Positive phase III results recently presented at the European Academy of Dermatology and Venerology Congress in Gothenburg (Sweden), Sep. 2010.
- All primary and secondary objectives were met and Actikerall[®] showed significant improvements over placebo and Solaraze[®].
- Actikerall[®] was filed in late 2009.
- Decentralized procedure expected to be completed in first half 2011.





- On Oct. 29th, Phase III topline results from ACCORD COPD II were released.
- The improvement from baseline in FEV1 (the primary endpoint) was statistically significant and reached a magnitude compared to placebo of 72mL.
- The bronchodilation effect was less than that observed in three other studies
 - The similarly designed ACCORD COPD I reported in January 2010
 - A previously reported 15-day Phase II trial comparing Eklira[®] to placebo or tiotropium
 - A recently completed 7-day Phase II trial comparing aclidinium to placebo or formoterol

where FEV1 ranged from 124mL to 186mL.

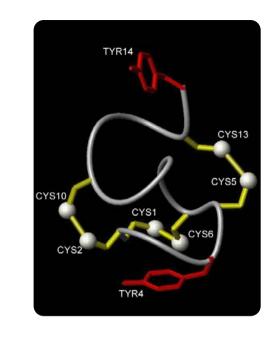
- A Phase III trial (ATTAIN) is currently under way with topline data expected in Q1 2011.
- EU and US submissions expected in mid 2011.



Linaclotide (I)

A first-in-class treatment developed for IBS-C

- IBS is a functional gastrointestinal disorder with abdominal pain and altered bowel habits.
- Patients suffering from IBS can be affected physically, psychologically, socially and economically.
- The overall prevalence of IBS estimated in Europe is 11.5%^{*}.
- No product approved by EMA so far in this indication.



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^{*} P. S. Hungin et al - The prevalence, patterns and impact of irritable bowel syndrome: an international survey of 40,000 subjects - Aliment Pharmacol Ther 2003; 17: 643–650.

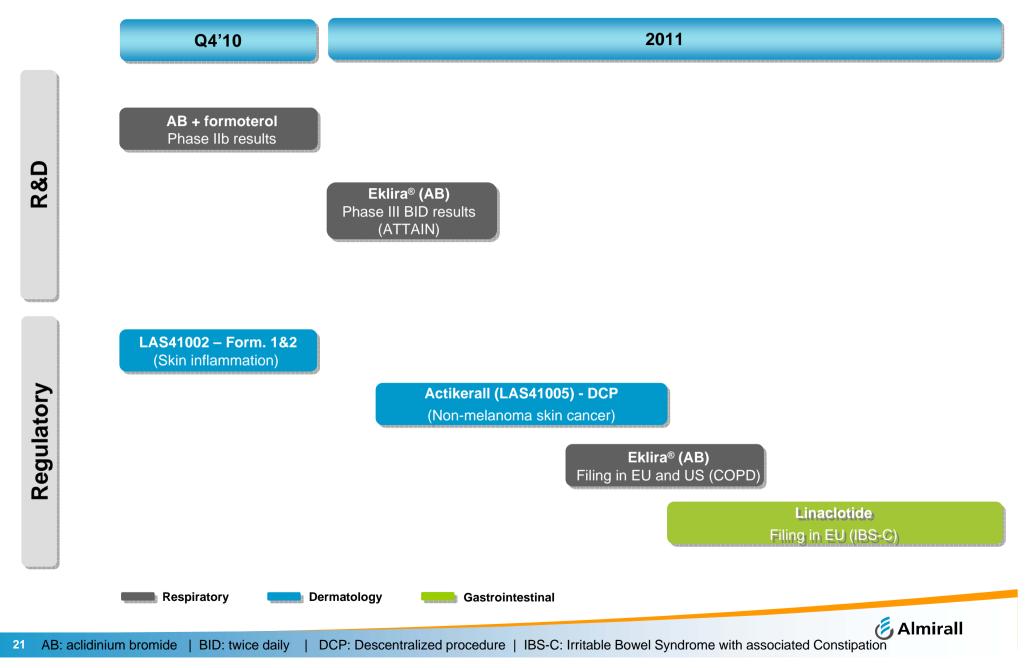
Linaclotide (II)

A first-in-class treatment developed for IBS-C

- Recently, positive topline results from two pivotal phase III studies in IBS-C were released.
- The two co-primary endpoints required by EMA* were met, showing statistical significance and sustained clinically relevant improvement for linaclotide-treated patients.
- All main secondary endpoints were also met, including 26-week endpoints.
- Safety results were consistent with those observed in previous linaclotide clinical studies.
- Almirall plans to file in Europe in second half 2011.



Newsflow during Q4'2010 and 2011



Guidance reiterated

• Sales – Mid single digit decline (%)

- Normalized Net Income Mid single digit decline (%)
- Q4 Restructuring costs planned*

*Only affects Net Income (not applicable to the Normalized Net Income forecast).

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