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FY 2011 Financial Results and Business update

February 27th, 2012

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2011 in review

Eduardo Sanchiz, CEO



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Highlights 2011

Results in line with guidance

Pipeline progress

- Two major filings (aclidinium, linaclotide) advancing well
- Aclidinium + formoterol and Sativex[®] (onco pain) in Phase III
- MABA to commence phase I shortly

Business development achievements

- Aclidinium partnered in Japan
- Ebastine licensed-out in China and Southeast Asia
- Roflumilast (Spain), scitalopram (Italy)

Continued cost focus

- Gross margin improved in a challenging environment
- SG&A declined 6,4% vs 2010
- Plant concentration completed. Toll manufacturing business closed

New products

- Launch of Sativex[®] (MS Spasticity) and Actikerall[®]
- 6 recent launches contributed to nearly € 50 MM in 2011 Sales



Positive momentum in 2012

- ✓ Aclidinium partnering progressing well in Europe
- ✓ Regulatory process ongoing for aclidinium and linaclotide
- ✓ Aclidinium outlicensed in Korea

✓ 2nd MRP wave of Sativex® in EU

✓ Xarelto® co-promotion in Spain



Financial Results 2011

Daniel Martinez, CFO



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FY 2011 Financial Results Summary

- Results in line with guidance
- Gross margin improvement (+20 bps) in a harsh environment
- Continued cost discipline (SG&A -6,4%)
- Solid balance sheet (Equity represents 58%)
- Debt neutral
- Healthy free cash flow generation
- Dividend proposal of €0,18* per share. Opportunity to take cash or shares



Income Statement

€rounded million	YTD Dec 2011	YTD Dec 2010	% Var
Net Sales	768,4	882,4	(12,9%)
Gross Profit	477,3	546,2	(12,6%)
% of sales	62,1%	61,9%	, , ,
Other Income	104,7	119,7	(12,5%)
R&D	(144,5)	(144,9)	(0,3%)
% of sales	(18,8%)	(16,4%)	
SG&A	(340,4)	(363,5)	(6,4%)
% of sales	(44,3%)	(41,2%)	
Other Op. Exp	(0,2)	(3,1)	(93,5%)
% of sales	(0,0%)	(0,4%)	
EBIT	96,9	154,4	(37,2%)
% of sales	12,6%	17,5%	
Depreciation	63,2	61,9	2,1%
% of sales	8,2%	7,0%	
EBITDA	160,1	216,3	(26,0%)
% of sales	20,8%	24,5%	
Sale of noncurrent assets / Other	(2,7)	(0,5)	n.m.
Restructuring costs	(9,9)	(11,6)	(14,7%)
Impairment reversals / (losses)	(7,0)	(14,0)	(50,0%)
Net financial income / (expenses)	(5,3)	(9,1)	(41,8%)
Corporate income tax	12,2	(0,6)	n.m.
Net income	84,2	118,6	(29,0%)
Normalized Net Income	97,9	136,7	(28,4%)
Earnings per share (€) (1)	0,51 €	0,71 €	
Normalized Earnings per share (€) ⁽¹⁾	0,59 €	0,82 €	
No of employees end of period	2.765	2.831	(2,3%)
(1) Number of charge at the and of the period			

- Gross margin improved in a challenging environment
- Continued cost discipline
- In line with guidance

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

Balance Sheet

€rounded million	December 2011	% of BS	December 2010
Goodwill	271,1	18,6%	271,9
Intangible assets	353,1	24,2%	382,8
Property, plant and equipment	152,1	10,4%	154,8
Financial assets	8,5	0,6%	10,2
Other non current assets	213,1	14,6%	189,0
Total Non Current Assets	997,9	68,5%	1.008,7
Inventories	93,2	6,4%	87,9
Accounts receivables	106,0	7,3%	103,8
Cash & equivalents	228,9	15,7%	312,9
Other current assets	30,6	2,1%	23,4
Total Current Assets	458,7	31,5%	528,0
Total Assets	1.456,6		1.536,7
Shareholders equity	854,7	58,7%	819,3
Financial debt	202,2	13,9%	297,5
Non current liabilities	188,3	12,9%	206,8
Current liabilities	211,4	14,5%	213,1
Total Equity and Liabilities	1.456,6		1.536,7

Net Debt as of 31 December 2011:

€14,6 MM *

x 0,09 EBITDA 2011

^{*} **Net Debt =** €202,2 MM Financial Debt **-** €228,9 MM Cash and Equivalents **+** €41,3 MM Pension Liabilities



Cash Flow

	YTD	YTD	
€rounded million	Dec 2011	Dec 2010	
Profit Before Tax	72,0	119,2	
Depreciation and amortisation	63,2	61,9	
Change in working capital	(6,6)	21,8	
Other adjustments	(28,5)	(28,3)	
Cash Flow from Operating Activities (I)	100,1	174,6	
Financial Income	8,5	4,3	
Investments	(38,5)	(93,0)	
Divestments	2,7	0,9	
Other cash flows	0,0	2,8	
Cash Flow from Investing Activities (II)	(27,3)	(85,0)	
Finance Expense	(14,7)	(16,9)	
Dividends distribution	(47,4)	(55,1)	
Debt increase/ (decrease)	(90,4)	36,5	
Other cash flows	(4,3)	(0,8)	
Cash Flow from Financing Activities	(156,8)	(36,3)	
Cash Flow generated during the period	(84,0)	53,3	
Free Cash Flow (III) = (I) + (II)	72,8	89,6	

Steady Free Cash Flow generation



Pipeline & Regulatory Update

Bertil Lindmark, CSO



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Regulatory Update

Aclidinium bromide

 Monotherapy submitted in 2011 for approval in US and EU.

- US:
 - Positive FDA AdCom voting on Feb 23rd
 - PDUFA in Q2'12
- EU: 120-days completed

Linaclotide

- Submitted for approval in 2011 for IBS-C in Europe
- First-in-class treatment with unique efficacy and safety profile
- No IBS-C product approved by EMA so far
- 120-days response received



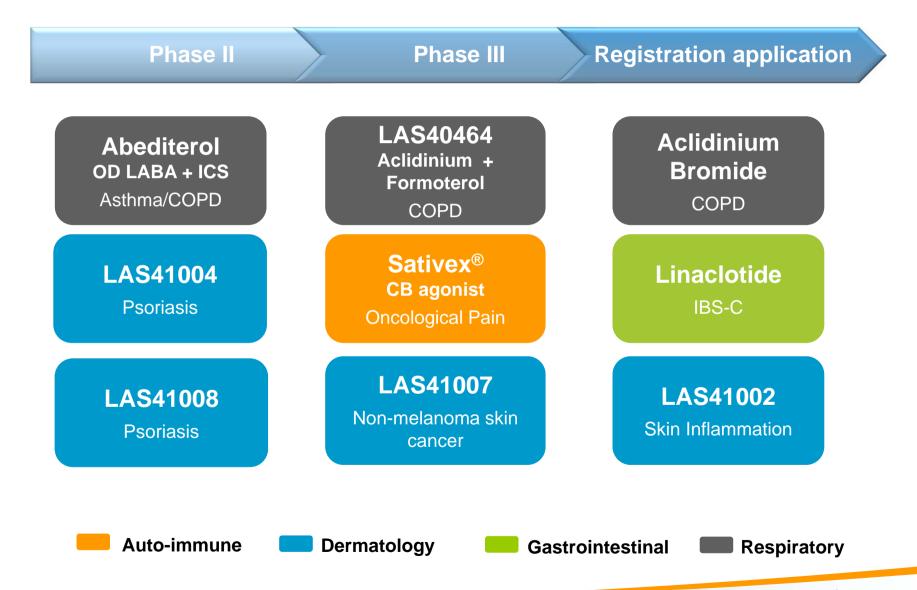
Aclidinium AdCom FDA Voting

	Questions	AdCom Voting
1	Do the efficacy data provide substantial evidence of a clinically meaningful benefit for aclidinium 400 mcg twice daily in the maintainance treatment of bronchospasm associated with Chronic Obstructive Pulmonary Disease (COPD)?	14 to 0 in favor
2	Has the safety of aclidinium been adequately assessed for the proposed indication?	10 to 3 in favor (1 abstention)
3	Do the efficacy and safety data provide substantial evidence to support approval of aclidinium 400 mcg twice daily for the maintainance treatment of bronchospasm associated with COPD?	12 to 2 in favor



A pipeline with significant upside

Preclinical and phase I projects not included





Platforms of growth

Luciano Conde, COO



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Aclidinium differentiation vs existing therapies

- Maximum efficacy from the 1st day of treatment and sustained over time
- Round-the-clock symptom control; early morning, day and night
- Superior bronchodilation during nighttime compared to tiotropium
- Clinically relevant improvements in quality of life and less use of rescue medication
- Reduction in exacerbations rate and improvement in exercise tolerance
- Well tolerated, low incidence of anticholinergic side effects (similar to placebo)
- Novel multi-dose, patient preferred, easy-to-use inhaler



Linaclotide

First-in-class opportunity

- Novel first-in-class compound for the treatment of IBS-C
- Once-a-day oral capsule
- Acts locally in the GI tract and is minimally absorbed
- Clinical studies involving nearly 1,600 patients with IBS-C have demonstrated sustained efficacy and a clean safety profile
- No IBS-C product approved by EMA so far
- Submitted for approval in 2011 for IBS-C in Europe
- Almirall holds commercial Pan-European rights



Additional growth platforms

Sativex®

- Launched by Almirall in three European countries in 2011
- Four additional launches planned in 2012
- Phase III ongoing in oncological pain

Dermatology franchise

- Leading derma player in Europe
- Key franchise that represents 15% of total sales
- Late stage derma pipeline progressing :
 - Skin inflammation
 - Non-melanoma skin cancer
 - Psoriasis



Sativex® Product roll-out status



Launched by Almirall

New launches planned in 2012

2nd MRP Wave moving forward

Germany

Spain

Denmark

Sweden

Italy

Austria

Czech Republic

Regulatory submisson in 10 additional European countries



New product cycle unfolding 49,1 12,3 ✓ Growing contribution over the quarters. 34,3 (in € Mill rounded) 22,1 36,8 9,9 26,5 17,6 8,2 Q1 2011 Q2 2011 Q3 2011 Q4 2011 ■ Tesavel + Efficib ■ Actikerall + Sativex + Silodyx + Toctino



2012 and beyond

Eduardo Sanchiz, CEO



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Guidance 2012-14

Barring unforeseen circumstances

	2012	2013	2014		
Sales (€MM)	Similar trends to 2011	Revenues to accelerate sharply			
Normalized Net Income (€MM)	Lower decline than in 2011	Lag effect on profitability	Profit growth		



Mid term takeaways

International growth : strategic priority

International sales > 70% of total by 2014

Double digit Sales CAGR 2012-14



Wrap-up and outlook 2012

- ✓ Aclidinium partnering progressing well in Europe
- ✓ Regulatory process ongoing for aclidinium and linaclotide
- Aclidinium outlicensed in Korea
- ✓ 2nd MRP wave of Sativex® in EU
- ✓ Xarelto[®] co-promotion in Spain

2012 Catalysts

- Aclidinium EU partnership
- PDUFA in Q2
- EU regulatory feedback for aclidinium and linaclotide



Appendixes

Q4 vs Q4

Crowndad million	2011	2010	0/ Vor	
€rounded million	4Q	4Q	% Var	
Net Sales	176,5	205,9	(14,3%)	
Gross Profit	106,3	129,2	(17,7%)	
% of sales	60,2%	62,7%		
Other Income	28,1	30,0	(6,3%)	
R&D	(47,2)	(42,5)	11,1%	
% of sales	(26,7%)	(20,6%)		
SG&A	(90,4)	(104,5)	(13,5%)	
% of sales	(51,2%)	(50,8%)		
Other Op. Exp	(1,0)	(4,6)	(78,3%)	
% of sales	(0,6%)	(2,2%)		
EBIT	(4,2)	7,6	(155,3%)	
% of sales	(2,4%)	3,7%		
Depreciation	16,3	15,9	2,5%	
% of sales	9,2%	7,7%		
EBITDA	12,1	23,5	(48,5%)	
% of sales	6,9%	11,4%		
Sale of noncurrent assets / Other	(3,7)	(0,5)	n.m.	
Reestructuring costs	(9,9)	(11,6)	(14,7%)	
Impairment reversals / (losses)	0,0	(13,0)	(100,0%)	
Net financial income / (expenses)	(1,6)	0,3	n.m.	
Tax	14,8	14,8	0,0%	
Net income	(4,6)	(2,4)	91,7%	
Normalized Net Income	4,2	14,8	(71,6%)	



Zoom in – Other Income

Includes Actonel[®], Conbriza[®], Libertek[®] (roflumilast) and Cipralex[®] (escitalopram)

€rounded million	YTD Dec 2011	YTD Dec 2010	% Var
Revenues under co-promotion agreements	10,9	21,5	(49,1%)
Collaboration in product promotion	18,4	24,6	(25,1%)
Revenues under co-development agreements	68,0	65,8	3,4%
Other	7,2	7,7	(1,4%)
Total	104,7	119,7	(12,5%)

Includes:

€44,9 MM of co-development revenues

€23,1 MM linked to upfront and milestones payments



Sales by Region

€rounded million	YTD Dec 2011	YTD Dec 2010	% var vs LY
Spain	388,0	495,3	(21,7%)
Europe & Middle East	291,0	288,7	0,8%
America, Africa & Asia Pacific	72,9	70,5	3,4%
Corporate	16,4	27,9	(41,1%)
Total	768,4	882,4	(12,9%)



Breakdown of the core business

Proprietary productsIn-licensing products

€rounded million		YTD Dec	YTDDec	% Var	Presence	
		2011	2010	YTD	Spain	Intl.
Ebastel® and others (ebastine)		116,2	117,5	(1,1%)	✓	\checkmark
Plusvent® (salmeterol & fluticasone)		54,3	57,0	(4,9%)	✓	
Almogran [®] and others (almotriptan)		49,7	48,2	3,1%	✓	\checkmark
Parapres® (candesartan cilexetile)		47,9	45,5	5,2%	✓	
Tesavel® (sitagliptine) + Efficib® (sitagliptine+metformine)		36,8	24,6	49,6%	✓	
Airtal [®] and others (aceclofenac)		30,8	39,1	(21,2%)	✓	✓
Esertia® (scitalopram)		30,6	61,3	(50,1%)	✓	
Prevencor® (atorvastatin)		30,5	72,9	(58,2%)	✓	
Solaraze [®] (sodium diclofenac)		26,6	24,7	7,7%	✓	✓
Opiren [®] (lansoprazole)		21,9	32,9	(33,3%)	✓	
Almax [®] (almagate)		19,4	22,8	(14,6%)	✓	✓
Decoderm [®] and others (flupredniden)		18,0	17,5	3,1%		\checkmark
Balneum [®] (urea)		17,8	18,6	(4,0%)	✓	✓
Pantopan [®] (pantoprazole)		17,7	18,6	(4,9%)		✓
Cidine® and others (cinitapride)		14,7	13,8	6,4%	✓	✓
Others		235,5	267,5	(12,0%)	✓	✓
Total		768,4	882,4	(12,9%)		



Net Sales breakdown by main Therapeutic Area

€ rounded million	YTD Dec 2011		
Respiratory	181,1	184,4	(1,8%)
Gastrointestinal and Metabolism	156,3	157,6	(0,8%)
Dermatology	118,2	120,0	(1,5%)
CNS	114,3	157,2	(27,3%)
Cardiovascular	106,2	149,6	(29,0%)
Osteomuscular	53,5	63,6	(15,8%)
Urological	20,2	16,9	19,5%
Other therapeutic specialties	18,6	33,3	(49,0%)
Total Net Sales	768,4	882,4	(12,9%)



For further information, please contact:

Jordi Molina
Investor Relations and Corporate Communication
Ph. +34 93 291 3087
jordi.molina@almirall.com

Or visit our website: www.almirall.com

