



CITI Healthcare Conference

New York, March 2nd, 2011

Almirall at a glance



Highlights 2010

Setting the basis for sustainable long-term growth

Results in line with guidance





Significant pipeline progression

- Phase III completion of aclidinium bromide monotherapy and linaclotide. Filings planned in 2011.
- Three additional projects entering phase III.

Challenging environment

- Price pressures hindered sales performance.
- Two healthcare reforms in Spain.

Focused cost optimisation

- SG&A reduction of 6%.
- Strong operational savings targeted: €7MM in 2011, €25MM from 2012 and beyond.

New products

- Launch of Toctino[®], Silodyx[®] and Conbriza[®]
- Excellent ramp-up of Tesavel[®] and Efficib[®]



Summary – 2010 Financial Results

- Sales (-4,7%) and Normalized Net Income (-5,9%) within guidance.
- Strong operational savings in 2010 (SG&A: -6,0%).
- Significant R&D investment to progress pipeline (+ 19,8%).
- Net Debt remains low at 0,10xEBITDA 2010 and provides strong strategic flexibility.
- Healthy Free Cash Flow generation (€89,6 MM).

€rounded million	2010	2009	% variation
Net Sales	882,4	925,5	(4,7%)
EBIT	154,4	179,1	(13,8%)
EBITDA	216,3	243,9	(11,3%)
Normalized Net Income	136,7	145,3	(5,9%)



Strategic axes to foster long term growth

		2007	2010	
1	Growing internationally			
	% of sales	34%	44%	
2	Leveraging own R&D			
	 Projects in clinical development stage 	8	12	
	 Projects in phase III / registration 	5	7	
3	Maintaining leadership in Spain			
	 Largest Spanish pharma company 	✓	✓	
	Ranked among top 5	\checkmark	\checkmark	



Our pipeline

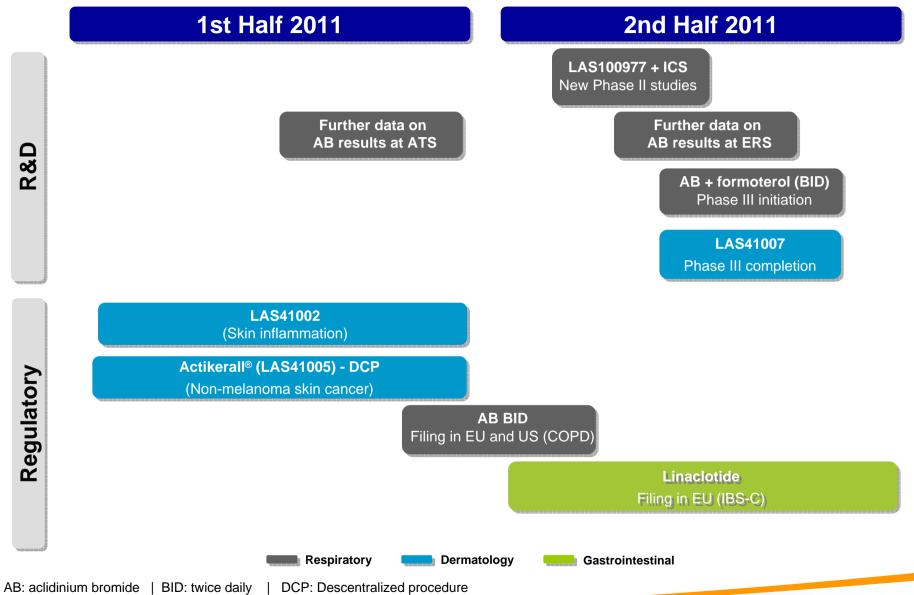


R&D 2010 highlights

- Positive phase III completion of aclidinium bromide monotherapy and linaclotide. Filings expected in 2011.
- Positive phase IIb completion of aclidinium bromide + formoterol.
 Phase III due to start in 2011.
- Reinforcing our respiratory franchise with a new MABA* candidate in preclinical stage.
- Sativex® starts phase III for the oncological pain indication.
- Following regulatory interactions, LAS100977 + ICS to progress in phase II in 2011.
- LAS41007 entered phase III for non melanoma skin cancer.



Newsflow during 2011



Three compounds entering phase III

	Indication	Start date	Compound
AB + formoterol	COPD	H2 2011	LAMA + LABA
Sativex®	Oncological pain	Q4 2010	Tetrahydro- cannabinol / cannabidiol
LAS41007	Non-melanoma skin cancer	Q4 2010	Undisclosed

AB: aclidinium bromide | LAMA: long acting muscarinic antagonist LABA: long acting beta-agonist



Two major filings expected in 2011: aclidinium bromide

- Positively completed phase III showed competitive efficacy and good tolerability.
- Well positioned to be the 2nd LAMA in the COPD market.



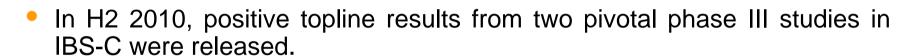
- Convenient Genuair® MDPI device.
- Sizable and growing global COPD market of € 9bn with significant unmet needs.
- FDA and EMA filing expected in mid 2011.
- Phase III fixed dose combination (aclidinium bromide + formoterol) to start in 2011.
- Further aclidinium bromide data to be presented at ATS (May'11) and ERS (September'11).



Two major filings expected in 2011: linaclotide

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

- IBS is a functional gastrointestinal disorder with abdominal pain and altered bowel habits.
- No product approved by EMA so far in this indication.



- 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.
- Almirall plans to file in Europe in second half 2011.



Building a strong respiratory franchise around the Genuair® device

- Aclidinium bromide ready to file in the US and EU.
- The aclidinium + formoterol combination progressing to phase III for US and EU.
- The Once Daily LABA/steroid combination progressing in phase II.
- Preparing for MABA* to enter in phase I.



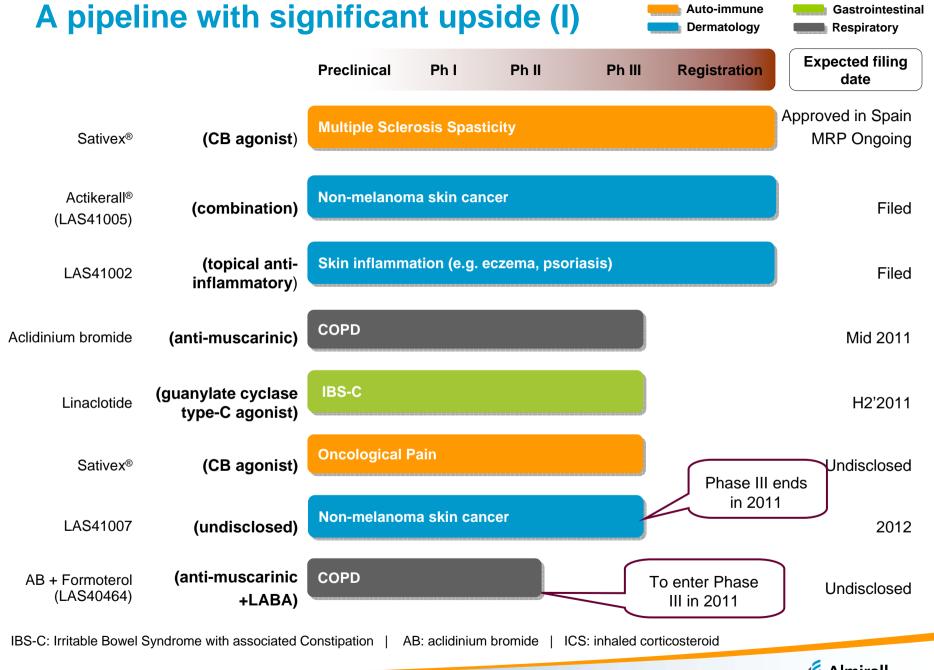
^{*} MABA: muscarinic antagonist beta agonist



Building the dermatology franchise

- Actikerall® (LAS41005) for non-melanoma skin cancer approaching regulatory feedback.
- LAS41002 for skin inflammation approaching regulatory feedback.
- LAS41007 for non-melanoma skin cancer: phase III expected to be completed in second half 2011.
- LAS41003 (infected eczema) + LAS41004 (psoriasis) progressed into phase II during 2010.
- ✓ Strong R&D connectivity with a growing therapeutic area in sales





A pipeline with significant upside (II)



		Preclinical	Ph I	Ph II	Ph III	Registration	Expected filing date
LAS100977 + ICS	(OD LABA + ICS)	Asthma / COF	PD				>2014
LAS41003	(combination)	Infected ecze	ma				2013
LAS41004	(combination)	Psoriasis					2014
LAS186323	(DHODH inhibitor)	RA/MS		То	be licensed		Undisclosed
LAS189913	(S1P1)	MS		To be licens	out		Undisclosed
LAS190792	MABA	COPD		Out			>2014

ICS: inhaled corticosteroid | RA: Rhematoid Arthritis | MS: Multiple Sclerosis | MABA: muscarinic antagonist beta agonist



Key deliverables in 2011



Projected key deliverables in the next 12 months

Operations

Corporate Development

R&D and Regulatory

- Launch of Sativex® in Spain, Denmark, Germany and Sweden.
- Continue roll out of Toctino[®].
- Launch of Solaraze[®] in Spain.
- Launch of LAS41005 (Actikerall®).
- Continued productivity improvements.

- Partnering of aclidinium in Europe and other selected geographies.
- Continue track record in licensing-in around core therapies.
- Explore acquisition opportunities.

- Two major filings: aclidinium and linaclotide.
- Aclidinium + formoterol combination enters phase III in 2011.
- LAS100977 (OD LABA) + ICS progressing in phase II.
- Continue development of MABA*.



^{*} MABA: muscarinic antagonist beta agonist

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