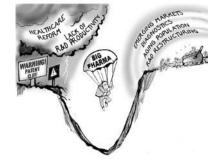
Sector View

Pharmaceuticals

LLY's solanezumab fails in mild dementia due to AD. Negative for MOR, ROG and GFS, but positive for AZN

	1 M	3 M	6 M	31/12/15
Healthcare	-5.6%	-9.4%	-7.3%	-15.8%
DJ Stoxx 600	-1.0%	-0.8%	1.2%	-6.8%
*Stoxx Sector Indices				

Companies o	overed			
ACTELION		BUY	CHF194	
Last Price	CHF155.1	Market Cap.	CHF16,714m	
ASTRAZENEO	CA	BUY	5100p	
Last Price	4168p	Market Cap.	GBP52,727m	
BAYER		NEUTRAL	EUR98	
Last Price	EUR88.34	Market Cap.	EUR73,053m	
GLAXOSMITHKLINE		BUY	1930p	
Last Price	1505p	Market Cap.	GBP73,386m	
GRIFOLS		NEUTRAL	EUR20	
Last Price	EUR17.985	Market Cap.	EUR11,419m	
IPSEN		BUY	EUR72	
Last Price	EUR61.83	Market Cap.	EUR5,156m	
NOVARTIS		NEUTRAL	CHF81	
Last Price	CHF69.1	Market Cap.	CHF181,534m	
NOVO NORE	DISK	NEUTRAL	DKK270	
Last Price	DKK220.7	Market Cap. DKK444,173r		
ROCHE HOLDING		BUY CHF285		
Last Price	CHF222.8	Market Cap.	CHF156,531m	
SANOFI		NEUTRAL	EUR84	
Last Price	EUR74.17	Market Cap.	EUR95,637m	
SHIRE PLC		BUY	6800p	
Last Price	4596p	Market Cap.	GBP41,525m	
SOBI		SELL	SEK90	
Last Price	EUR93.75	Market Cap.	EUR25,349m	
UCB		NEUTRAL	EUR80	
Last Price	EUR61.5	Market Cap.	EUR11,962m	



Lilly's solanezumab (an anti- β amyloid monoclonal antibody) failed to bring a statistically significant benefit to patients with mild dementia due to Alzheimer's disease (AD) in a large Phase III study. We believe the street will increasingly question the viability of MOR/ROG's gantenerumab as well as GFS' Albutein; both being tested as treatments for Alzheimer's disease in different Phase III trials... and both targeting β amyloid. On the other hand, the read-across could be positive for AZN's BACE inhibitor as we believe LLY will probably decide to reprioritize it following solanezumab's failure.

ANALYSIS

- Another failed anti-β amyloid trial. Lilly's solanezumab (an anti-β amyloid monoclonal antibody) failed to bring a benefit in patients with mild dementia due to Alzheimer's disease (AD) in a large Phase III study (EXPEDITION 3). Patients treated with solanezumab did not experience a statistically significant slowing in cognitive decline compared to patients treated with placebo (p=0.095), as measured by the ADAS-Cog (Alzheimer's Disease Assessment Scale-Cognitive subscale). And while the study results (including a range of different secondary clinical endpoints) directionally favoured solanezumab, the magnitudes of treatment differences are said to be quite small.
- Lilly to discontinue solanezumab's development for the whole AD indication? Admittedly, this
 development involved a condition related to AD... But we believe the potential impact of this
 failure could be broader than what the street actually expects, especially since LLY's management
 clearly stated during its Q3 2016 conference call that they would re-evaluate all the "sola"
 programs should EXPEDITION 3 be a failure (and that might include the developments in
 prodromal AD).

VALUATION

- Negative for MOR/ROG, GFS. As underlined in our reports dealing with Morphosys (<u>"Back for MORe"</u>) and Grifols (<u>iEl consenso al borde de un ataque!</u>), we were/are quite pessimistic about the different options developed in AD, and especially the ones targeting the β amyloid protein... So our base-case scenarios are unchanged. Having said that, we believe the street will increasingly question the viability of MOR/ROG's gantenerumab as well as GFS' Albutein; both being tested as AD treatments in different Phase III trials... and both targeting β amyloid.
- Regarding GFS more specifically, we continue to ask ourselves whether the design of the ongoing AMBAR Phase III study is really satisfactory from a regulatory standpoint (less than 500 patients enrolled whereas millions of people are known to be affected by this highly heterogeneous condition).
- But potentially positive for AZN. On the other hand, the read-across could be positive for AZN's BACE inhibitor as we believe LLY will probably decide to reprioritize it following solanezumab's failure.

NEXT CATALYSTS

- 8th December: Presentation of detailed data from the EXPEDITION 3 trial at the CTAD (Clinical Trials on Alzheimer's disease) meeting.
- H1 2017: Phase III results for Grifols' Albutein in AD (AMBAR study).

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Stock rating

BUY	Positive opinion for a stock where we expect a favourable performance in absolute terms over a period of 6 months from the publication of a				
Der	recommendation. This opinion is based not only on the FV (the potential upside based on valuation), but also takes into account a number of				
	elements that could include a SWOT analysis, momentum, technical aspects or the sector backdrop. Every subsequent published update on the				
will feature an introduction outlining the key reasons behind the opinion.					

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Distribution of stock ratings

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NEUTRAL ratings 32.9%

SELL ratings 11.4%

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