

Galapagos NV GLPG.OQ GLPG US

EQUITY: AMERICAS BIOTECHNOLOGY

Increasing TP to \$121

Baricitinib CRL Improves Filgotinib Positioning

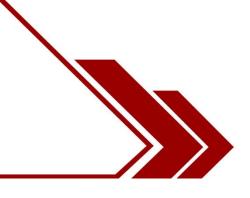
We are increasing our Target Price for GLPG to \$121 from \$87. We view baricitinib's CRL (April 14) as a positive for GLPG, which will ultimately improve filgotinib's opportunity in the competitive RA market.

- Baricitinib (LLY/INCY) CRL May Limit First-Mover Advantage vs. Filgotinib. We now anticipate at least a 1-year delay in baricitinib's approval, slimming the potential lead time on US market over filgotinib. Filgotinib top-line P3 RA results YE18. Additionally, baricitinib CRL-related dose/safety questions may impair the product label and marketing efforts. We now estimate filgotinib can more deeply penetrate the RA market; increasing market penetration in RA to 5% from previously 4%.
- Filgotinib's Higher Specificity May Confer Improved Risk/Benefit vs. Baricitinib. Filgotinib' s selective JAK1-inhibition may confer improved risk/benefit vs. baricitinib and tofacitinib. Xeljanz (tofacitinib), a marketed JAK1/3 inhibitor, remains limited by tolerability and mild efficacy. Baricitinib specificity (JAK2 and JAK1) demonstrated an improved safety profile compared to tofacitinib. We expect that the specific JAK1-inhibition of filgotinib will have lower off-target activity, resulting in an improved safety profile of filgotinib, particularly in combination.
- Filgotinib's FINCH Phase 3 Program Includes a Larger Database in Combination with Widely Used MTX. The FINCH program for filgotinib in RA incorporated 3,273 patients similar to baricitinib's RA P3 program (n=3,388). However, in contrast to baricitinib, filgotinib's program includes two studies that evaluate the combo of filgotinib (high and low dose) and MTX in a total number of 2,850 patients, providing a large safety database for this combo and across doses.
- Next Filgotinib Landscape Data Flow: (1) ABT-494 Crohn's data in 210 pts (4-doses) with inadequate response to anti-TNFs in May 2017; (2) filgotinib's DARWIN3 LOE study is ongoing, top-line data at EULAR (Jun. 14-17); (3) data from CELG's ozanimod in CD, mongersen in UC 1H17.
- Increasing Target Price to \$121 from \$87. We now anticipate a higher penetration into the Rheumatoid Arthritis market from 4% to 5% and raise our TP to \$121 from \$87. GLPG's CF triple combo program represents \$30/share.

Summary of Financials (욥hs except per share)									
	2016A	2017E	2018E						
Total Revenues	€151,612	€166,055	€199,836						
Operating Expense	€163,103	€385,451	€426,043						
Operating Income	(€11,491)	(€219,396)	(€226,207)						
Net Income, (GAAP)	€54,012	(€211,373)	(€226,442)						
Diluted EPS, (GAAP)	€1.14	(€4.35)	(€4.68)						
Cash (mn)	€973	€831	€637						
Diluted Shares Outstanding (th)	47,308	48,624	48,434						

Source: Company data, Instinet estimates

Key company data: See next page for company data and detailed price/index chart.



Instinet, LLC, Equity Research

17 April 2017

Rating Remains	Buy
Target Price Increased from 87.00	USD 121.00
Closing price 13 April 2017	USD 89.96
Potential upside	+34.5%

Research analysts

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Key data on Galapagos NV

Rating

Stock	Buy
Sector	Not rated

Relative performance chart



Source: Thomson Reuters, Instinet research

Performance

(%)	1M	3M	12M
Absolute	15.2	30.0	107.3
Relative to Nasdaq	18.5	06.4	102.0
Biotechnology Index	10.0	26.1	103.8
Stock price data			
Current stock price (\$)			89.96

Current stock price (\$)	89.90
Market cap (\$ - mn)	4,161.2
52-week low (\$)	43.20
52-week high (\$)	90.46
Shares outstanding (mn)	46.26

Source: Thomson Reuters, Instinet research

Company Description

Galapagos NV is a clinical-stage biotechnology company focused on the discovery and development of small-molecule therapeutics in an area of disease with a considerable unmet need. The company's lead asset, filgotinib, is a JAK1-specific inhibitor for the treatment of inflammatory diseases. Filgotinib is being studied in three Phase 3 studies for the treatment of rheumatoid arthritis and two Phase 3 studies in inflammatory bowel disease. In addition, the company is developing a triple-combination therapy for the treatment of cystic fibrosis. One of the components of this combo demonstrated positive results in a Phase 2 study. The two additional compounds are in Phase 2 and 1 studies. Galapagos plans to initiate a triple-combo study in patients during 2H17.

Valuation

We change our valuation to top-line revenue multiple from DCF, a method widely used in early-stage biotechnology valuation. We also increase filgotinib peak penetration into the RA market to 5% from previously 4%, due to the delayed approval of baricitinib in this indication. We arrive at our target price of \$121 by using a 6x multiple for EU profit share on filgotinib across inflammatory indications and a 16x multiple for US royalties on filgotinib. In filgotinib for RA we apply a 15% discount rate, reflecting a lower development risk as the target, JAK, is already validated by an approved drug in RA. In filgotinib in UC and Crohn's, we apply at 20% discount rate, reflecting a slightly higher risk for these indications for which no JAK inhibitor is approved. For the Cystic Fibrosis program we use an 18x multiple, reflecting a higher value for the higher-margin orphan program and a 25% discount that reflects a higher development risk.

Fig. 1: Top-Line Revenue Valuation

		Expected	Peak	Valuation		Discounted			
Drug/Indication		Expected Launch	Sales/Royalty Est (\$MM)	Year Sales	Multiple	Asset Value	Value / Share	Discount Rate	Partner
Filgotinib - Profit Split EU Big 5									
RA		2021	\$306	2025	6	\$690	\$14	15%	Gilead
Crohns		2022	\$76	2025	6	\$127	\$3	20%	Gilead
UC		2023	\$34	2025	6	\$57	\$1	20%	Gilead
	Sub Total		\$416			\$874	\$18		
Filgotinib - US Royalties									
RA		2020	\$275	2025	16	\$1,654	\$34	15%	Gilead
Crohns		2021	\$192	2025	16	\$857	\$18	20%	Gilead
UC		2022	\$28	2025	16	\$125	\$3	20%	Gilead
	Sub Total		\$495			\$2,636	\$54		
CF Triple Combo		2021	\$370	2025	18	\$1,477	\$30	25%	Abbvie
Pipeline Value						\$4,988	\$103		
Net Cash (YE:2017)						\$831	\$18		
Total Equity Value							\$121		
Diluted Shares Outstanding Use	d for Valuation	(MM)						48.6	
Numbers may not add up due to	rounding.								

Source: Company reports, Instinet estimates

Upcoming Catalysts

Fig. 2: Potential Upcoming Catalysts

Date	Program	Indication	Impact	Event Description
1H17	Filgotinib	RA	+++	Competitor: Baricitinib PDUFA date. Interested in whether baricitinib will receive the same label as tofaitinib
1H17	Mongersen	UC	++	Competitor: Potential Phase 2 study results in UC
1H17	Ozanimod	CD	++	Competitor: Potential Phase 2 study results in CD
1H17	Otezla	CD	++	Competitor: Potential Phase 2 study results in UC
1H17	filgotinib	RA	++	Interim analysis of the LTE study in RA will be presented at the EULAR,
1H17	Filgotinib	CD	+	Additional data presentation from the FITZROY studies at the DDW conference
1H17/ACR	GLPG1972	Osteoarthritis	+	Data from first in humans study will disclose cartilage breakdown and target
1H17	GLPG1972	Osteoarthritis	+	US IND and initiation of a Phase 1b study
1H17	ABT-494	CD	++	Competitor: Phase 2 readout from AbbVie's ABT-494 in CD at DDW (May 6-7)
1H17	VX-661	CF	++	Competitor: Vertex results from a Phase 3 study with VX-661+ivacaftor
3Q17	Triple combo	CF	++	Initiation of a triple combo study in patients
2H17	2451+2222	CF	+++	Results of a combo study in healthy volunteers
2H17	Filgotinib	CD	+	Additional data presentation from the FITZROY studies at the UEGW conference
2H17	filgotinib	RA	+	Additional data presentation at the ACR conference
2H17	VX-440/VX-152	CF	++	Competitor: Vertex triple combo data from a Phase 2 study in CF patients
2H17	VX-659	CF	+	Competitor: Vertex triple combo data from a Phase 1 study in CF patients
2H17	2222	CF	+++	Results of Phase 2 in patients study in combination with Kalydeco
2H17	2451	CF	++	Results from a FIH study
3Q17	Triple combo	CF	+++	Results of the triple combo in healthy volunteers
1Q18	Triple combo	CF	+++	Results of the in patients study with the triple combo
Early2018	Filgotinib	UC	+++	Interim analysis of the Phase 2/3 study
2H18	Mongersen	CD	+++	Competitor: Phase 3 study results
2H18	Ozanimod	UC	+++	Competitor: Phase 3 study results

Source: Company reports, Instinet research

Fig. 3: Income statement

GLPG Income Statement

(€1000s, except per share data) [FY - Dec]	2015	1Q16	2Q16	3Q16	4Q16A	2016A	1Q17	2Q17	3Q17	4Q17	2017E	2018E
US Filgotinib Sales (RA Only)	0	0	0	0	0	0	0	0	0	0	0	0
Intl Filgotinib Sales (RA Only)	0	0	0	0	0	Ŭ	0	0	0	0	Ű	0
US Filgotinib Sales (Crohn's) Intl Filgotinib Sales (Crohn's)	0	0	0	0	0	0	0	0	0	0	0	0
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US Filgotinib Sales (UC only) Intl Filgotinib Sales (UC only)	0	0	0	0	0	0	0	0 0	0 0	0	0	0
Total Filgotinib Royalties	0	0	0	0	0	0	0	0	0	0	0	0
US GLPG- Triple Combo Royalties (CF)	0	0	0	0	0	0	0	0	0	0	0	0
Intl GLPG-Triple Combo Royalties (CF)	0	0	0	0	0	0	0	0	0	0	0	0
Total GLPG1837+Corrector Royalties	0	0	0	0	0	0	0	0	0	0	0	0
R&D revenue	39,563	10,121	28,674	11,214	79,510	129,519	22,274	28,956	37,643	48,935	137,807	151,588
Other Income	21,017	4,696	5,273	5,062	7,062	22,093	7,062	7,062	7,062	7,062	28,248	48,248
Total Revenues	60,579	14,817	33,947	16,276	86,572	151,612	29,336	36,018	44,705	55,997	166,055	199,836
Costs & Expenses:			· ·				· ·					
Cost of Goods Sold	0	0	0	0	0	0	0	0	0	0	0	0
R&D	129,714	27,818	34,594	34,327	42,834	139,573	55,684	72,389	94,106	122,338	344,518	378,970
SG&A	19,127	3,972	5,854	5,685	8,018	23,529	8,820	9,702	10,672	11,739	40,933	47,073
Restructuring and integration costs	1,182	422	454	396			0	0	0	0	0	0
Total Operating Expenses	150,023	32,212	40,902	40,408	50,852	163,103	64,504	82,091	104,778	134,077	385,451	426,043
Operating Income	(89,444)	(17,395)	(6,955)	(24,132)	35,720	(11,491)	(35,168)	(46,073)	(60,074)	(78,080)	(219,396)	(226,207)
Interest and Other Income (Expense), net	(30,632)	57,479	0	0	0	57,479	0	0	0	0	0	0
Pretax Income (Loss)	(119,627)	35,950	(4,371)	(23,439)	46,106	54,246	(33,104)	(44,009)	(58,009)	(76,016)	(211,138)	(226,207)
Income tax expense (Benefit)	1,218	0	24	(95)	(164)	(235)	(59)	(59)	(59)	(59)	(235)	(235)
Net Income (Loss) as reported	(118,410)	35,950	(3,721)	(24,091)	45,874	54,012	(33,163)	(44,068)	(58,068)	(76,074)	(211,373)	(226,442)
Stock option expense	5,036	1,092	3,150	2,959	3,833	11,034	3,225	4,105	5,239	6,704	19,273	29,823
Other	0	0	0	0	0	(1,605)	0	0	0	0	0	0
Net Income (Loss) Non-GAAP	(113,374)	37,042	(571)	(21,132)	49,707	63,441	(29,937)	(39,963)	(52,829)	(69,370)	(192,100)	(196,619)
Diluted Earnings Per Share Non-GAAP	(€2.90)	€0.83	(€0.01)	(€0.46)	€1.01	€1.34	(€0.62)	(€0.82)	(€1.07)	(€1.43)	(€3.95)	(€4.29)
Basic Earnings Per Share Non-GAAP	(€2.81)	€0.81	(€0.01)	(€0.44)	€1.07	€1.39	(€0.64)	(€0.87)	(€1.14)	(€1.52)	(€4.16)	(€4.29)
Diluted Earnings Per Share	(€3.03)	€0.81	(€0.10)	(€0.53)	€0.93	€1.14	(€0.69)	(€0.91)	(€1.18)	(€1.57)	(€4.35)	(€4.68)
Basic Earnings Per Share as reported	(€2.94)	€0.79	(€0.10)	(€0.52)	€0.99	€1.18	(€0.71)	(€0.96)	(€1.25)	(€1.66)	(€4.58)	(€4.94)
Basic Shares Outstanding (th)	39,076	44,425	45,229	45,527	46,450	45,696	46,496	45,742	46,543	45,787	46,142	45,833
Diluted Shares Outstanding (th)	40,308	45,492	46,756	48,328	49,251	47,308	48,023	48,542	49,344	48,588	48,624	48,434

Source: Company reports, Instinet estimates

Fig. 4: Balance Sheet

(€1000s, except per share data) [FY - Dec]	2015A	2016A	2017E	2018E
ASSETS				
Current assets:				
Cash and cash equivalents	340,314	973,241	831,362	637,430
Current restricted cash	6,857	6,570	6,570	6,570
Current R&D incentives receivables	9,161	10,154	10,154	10,154
Current financial assests from share subscription agreement	8,371	0	-	
Short term marketable securities	0		0	0
Trade & other receivables	3,931	9,728	2,918	2,918
Inventory	325	300	300	300
Prepaid expenses and other current assets	5,512	7,239	7,239	7,239
Total current assets	374,470	1,007,232	858,544	664,612
Property and equipment, net	13,782	14,961	17,908	23,461
Goodwill	0	0	0	0
Intangible assets	1,550	1,023	1,023	1,023
Deferred tax assets/receivables	1,726	1,957	1,957	1,957
Non-current R&D incentives receivables	49,384	54,188	54,188	54,188
Non-current restricted cash	1,046	1,098	1,098	1,098
Other non-current assets	557	2,880	2,879	2,879
Total assets	442,514	1,083,338	937,596	749,218
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LIABILITIES AND STOCKHOLDER'S EQUITY				
Current liabilities:				
Trade and other payables	29,482	31,269	77,090	85,209
Current obligations under finance lease	52	54	54	54
Current tax payable	2,583	1,022	1,022	1,022
Accrued charges	490	619	1,156	1,022
Deferred income	39,806	70,827	70,827	70,827
Other current liabilities	00,000	0,021	0,021	0
Total current liabilities	72,412	103,791	150,150	158,390
	,	,	,	,
Long term debt	0	0	0	0
Obligations under finance lease	63	9	9	9
Deferred Revenue	0	214,785	214,785	214,785
Provisions	55	63	63	63
Pension liabilities	2,693	3,520	3,520	3,520
Other liabilities	2,291	2,469	2,469	2,469
Total liabilities	77,515	324,637	370,996	379,236
Stockholders' equity:				
Common Stock	185,399	223,928	223,928	223,928
Additional paid in capital	357,402	649,135	649,135	649,135
Other reserves	(18)	(1,000)	(1,000)	(1,000)
Translation differences	(467)	(1,090)	(1,090)	(1,090)
Accumulated other comprehensive loss	0	0	0	0
Accumulated Deficit	(177,319)	(112,272)	(304,372)	(500,991)
Total stockholders' equity	364,999	758,701	566,601	369,982
Total liabilities and stockholders' equity	442,514	1,083,338	937,596	749,218

Source: Company reports, Instinet research

Fig. 5: Cash Flow Statement

(€1000s, except per share data) [FY - Dec]	2015A	2016A	2017E	2018
CASH FLOWS FROM OPERATING ACTIVITIES				
Net Income (Loss)	(118,410)	54,012	(211,373)	(226,442
Adjustments	(110,410)	04,012	(211,070)	(220,442
Tax income/expenses	(1,218)	235	0	C
Other net financial income	(448)	(8,258)	0	(
Fair value measurment of share subscription	30,632	(57,479)	0	(
Depreciation and amortization	3,402	4,182	3,740	4,477
Net realized loss for foreign exchange transaction	(398)	1,229	0	(
Stock based compensation	5,036	11,034	19,273	29,823
Other	0	,	0	,
Change in assets and liabilities:			-	
Increase/decrease in provisions	(125)	7	0	
Increase pension liabilities	30	244	0	(
Gain on sale of fixed assests	(62)	(14)	0	(
Inventories	(02)	(14)	0	
Account receivables	(7,220)	(12,978)	6,810	
Prepaid expenses & other assets	0	0	0	(
Accounts payable and accrued expenses	(26,728)	2,102	46,359	8,240
Interest paid	(49)	(47)	0	
Interest received	1,106	1,066	0	
Income taxes paid/received	(94)	(1,763)	0	(
Current obligations under finance lease	0	2	0	(
Deferred revenues & other	0	245,806	0	(
Net cash provided by (used in) operating activities	(114,590)	239,405	(135,192)	(183,902
CASH FLOWS FROM INVESTING ACTIVITIES				
Purchases of property and equipment	(6,100)	(4,458)	(6,687)	(10,03
Purchase of and expenditure of intangible fixed assets	(565)	(332)	0	(
Proceeds from disposal of PPE	110	18	0	(
Increase/decrease in restricted cash	2,258	235	0	(
Investments, net Other	0	(2,750) 0	0	(
Net cash used in investing activities	(4,297)	(7,287)	(6,687)	(10,03
	(1,=01)	(1,201)	(0,001)	(10,00)
CASH FLOWS FROM FINANCING ACTIVITIES				
Proceeds from issuance of shares, net cost	271,413	391,784	0	(
Exercise of options	0	4,261	0	(
Repayment obligations under finance and other debt	(43)	(49)	0	(
Repurchase of common stock	0	0	0	(
Other	0	205 000	0	(
Net cash provided by financing activities	271,370	395,996	0	
Effect of exchange rate on cash	118	4,816		
Net increase in cash and cash equivalents	152,483	632,927	(141,879)	(193,932
Cash and cash equivalents at beginning of period	187,712	340,314	973,241	831,362
Cash and cash equivalents at end of period	340,314	973,241	831,362	637,430

Source: Company reports, Instinet research

Appendix A-1

Analyst Certification

I, Christopher Marai, hereby certify (1) that the views expressed in this Research report accurately reflect my personal views about any or all of the subject securities or issuers referred to in this Research report, (2) no part of my compensation was, is or will be directly or indirectly related to the specific recommendations or views expressed in this Research report and (3) no part of my compensation is tied to any specific investment banking transactions performed by Nomura Securities International, Inc., Nomura International plc or any other Nomura Group company.

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Materially mentioned issuers

Issuer	Ticker	Price	Price date	Stock rating	Sector rating	Disclosures
Galapagos NV	GLPG US	USD 89.96	13-Apr-2017	Buy	Not rated	A6

A6 The Nomura Group expects to receive or intends to seek compensation for investment banking services from the subject company in the next three months.

Galapagos NV (GLPG US) Rating and target price chart (three year history)	USD 89.96 (13-Apr-2017)	Buy (Sect	or rating: I	Not rated)	
11-Apr-2014 to 10-Apr-2017	PRICE(USD)	Date	Rating	Target price	Closing price
90.00 85.00 80.00 75.00 65.00 60.00 55.00 55.00 40.00 35.00 30.00 25.00 15.00 30.00 25.00 50.00 55	Worked Worked and Barrow	01-Mar-17 01-Mar-17 27-Oct-15 25-Sep-15 08-Jun-15 08-Jun-15	Buy Not Rated Buy	87.00 62.00 73.00	71.78 71.78 47.69 44.60 55.51 55.51
0.00 Jul-14 Oct-14 Jan-15 Apr-15 Jul-15 Oct-15 Jan-1 — GALAPAGOS NV 🔺 Target Price Change 🌰 R					

For explanation of ratings refer to the stock rating keys located after chart(s)

Valuation Methodology We use top-line revenue multiple valuation, a method widely used in early-stage biotechnology valuation. We arrive at our target price of \$121 by using a 6x multiple for EU profit share on filgotinib across inflammatory indications and a 16x multiple for US royalties on filgotinib. In filgotinib for RA we apply a 15% discount rate, reflecting a lower development risk as the target, JAK, is already validated by an approved drug in RA. In filgotinib in UC and Crohn's we apply a 20% discount rate, reflecting a slightly higher risk for these indications for which no JAK inhibitor is approved. For the Cystic Fibrosis program we use a 18x multiple, reflecting a higher value for the higher-margin orphan program and a 25% discount that reflects a higher development risk.

Risks that may impede the achievement of the target price Regulatory risk: The FDA may require Galapagos to present data on the efficacy of the individual triple combo drugs in the target patient population, which would require Galapagos to conduct a large Phase 2 study. Enrollment of patients in these studies might be challenging due to the low expectation of efficacy from a single compound. For filgotinib, the FDA may issue a class label concerning the risk for serious infections and malignancies. This action will not prevent filgotinib from reaching the market, but it could create a negative perception of the drug among patients and physicians, whih would affect commercial viability of the drug. Competitive risk: Baricitinib, a JAK 1/2 inhibitor, was expected to be approved by January 19, 2017. In clinical studies, the drug presented compelling efficacy, superior

to adalimumab. If baricitinib is found to be safe and approved without a black-box warning, it has the potential to take the lion's share of the market. Celgene's mongersen, a SMAD7 anti-sense RNA, showed compelling safety and efficacy profile in a Phase 2 study in CD patients. The compound is in a Phase 3 study and is set to report top-line data by 2H18. If approved, mongersen would have first-mover advantage as the only orally available DMT for Crohn's. Clinical risk: The Phase 2 study with filgotinib in CD used the CDAI as the primary outcome measure. The Phase 3 study is using the more traditional PRO as the primary outcome measure as may result in a smaller efficacy difference between the placebo and treatment arms in the Phase 3 study.

Important Disclosures

Online availability of research and conflict-of-interest disclosures

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As at 31 March 2017.

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STOCKS

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SECTORS

A 'Bullish' stance, indicates that the analyst expects the sector to outperform the Benchmark during the next 12 months. A 'Neutral' stance, indicates that the analyst expects the sector to perform in line with the Benchmark during the next 12 months. A 'Bearish' stance, indicates that the analyst expects the sector to underperform the Benchmark during the next 12 months. Sectors that are labelled as 'Not rated' or shown as

'N/A' are not assigned ratings. Benchmarks are as follows: United States: S&P 500; Europe: Dow Jones STOXX 600; Global Emerging Markets (ex-Asia): MSCI Emerging Markets ex-Asia. Japan/Asia ex-Japan: Sector ratings are not assigned.

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