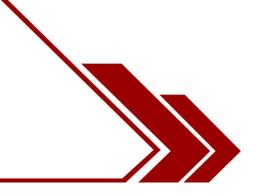


Galapagos NV GLPG.OQ GLPG US

EQUITY: AMERICAS BIOTECHNOLOGY



ABBV Data Update and GLPG Co-Promote

Filgo Emerging as the Safer JAK Inhibitor

This morning, **ABBV** reported positive top-line data from its Phase 3 SELECT-MONOTHERAPY trial evaluating upadacitinib (Upa), an oral JAK1-selective inhibitor, as monotherapy in moderate-severe rheumatoid arthritis (RA) patients with inadequate response to methotrexate (MTX). At 14 weeks, both doses (15mg and 30mg QD) met the primary endpoints of ACR20/50/70 and LDA score. Upa's safety profile was consistent with previously reported studies; however, there was one fatal hemorrhagic stroke in a patient with pre-existing cardiovascular issues in the 15mg cohort; another SAE, pulmonary embolism, was observed in the 15mg cohort in a patient with pre-existing conditions.

- Upa Boasts Comparable Efficacy to Filgo in Monotherapy (Fig. 1), but Safety/Tolerability Are Still Key Competitive Advantage for Fligo. No new safety signals in this study; SAEs occurred at 5% and 3% in the 15 mg and 30 mg groups, respectively, and at 3% in the MTX group. Filgotinib resulted in less reduction of NK cells and a lower rate of opportunistic infection compared with other JAK inhibitors (Figs. 2-3). Also, filgotinib patients experienced an increase in Hb levels.
- GLPG Decides to Opt In to Co-promote Filgotinib, with Partner GILD, in 8 EU countries (Germany, France, Italy, Spain, UK, Netherlands, Belgium, Luxembourg). Outside these territories, Galapagos will be eligible for tiered royalties ranging from 20% to 30% on net filgotinib sales. We anticipated GLPG would opt in to the co-promote, given its desire to build a commercial organization and cash war chest. Deal terms were favorable, with GLPG funding 35% of co-promotion costs at a 50/50 profit split. See p. 5 for model updates.
- Co-Promotion Lock-In Instills Confidence in GILD Collaboration Ahead
 of Equity Investment Lockup Expiration Date. The impending expiration
 of GLPG's lockup and standstill agreement with GILD remains December
 31, 2017. We believe GLPG's decision to opt in on co-promotion of filgotinib
 ahead of the lockup expiration date speaks to management's confidence in
 the potential of its highly selective JAK1 inhibitor.
- Filgotinib's Safety Profile Appears Superior to Upadacitinib's on Paper. However, clinical experience may not differ substantially, potentially leading to marketing prowess trumping data, post-approval.

2016A 2017E 2018E Year-end: Dec EPS (€) Actual Prev. Curr. Cons. Prev. Curr. Cons. 1Q 0.79A -0.29A -0.76E -0.50E N/A N/A N/A 2Q -0.08A -0.72A -0.97E -0.50E N/A N/A N/A 3Q -0.62E -0.52A N/A -0.72A N/A N/A -1.00E 4Q 0.96A N/A -0.64E -0.70E N/A -0.41E -0.54E -1.98E -2.40E Year 1.14A N/A -2.39E N/A -2.10F 1,055,422 1,034,489 Cash & Equivalents (€k) 973,241 1,152,929 1,170,871 N/A N/A

Source: Company data, FactSet, Instinet estimates

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

Instinet, LLC, Equity Research

20 December 2017

Rating Remains	Buy
Target Price Remains	USD 124.00
Closing price 19 December 2017	USD 91.26
Potential upside	+35.9%

Research analysts

Americas Biotech

Christopher Marai, Ph.D. - ILLC Christopher.Marai@Instinet.com + 1 212-310-5466

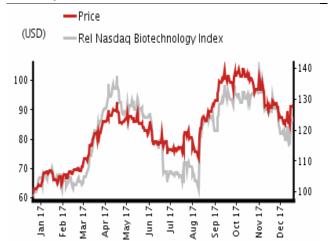
Production Complete: 2017-12-20 21:35 UTC

Key data on Galapagos NV

Rating

Stock	Buy
Sector	Not rated

Relative performance chart



Source: Thomson Reuters, Instinet research

Performance as of 19 December 2017

(%)	1M	3M	12M
Absolute	0.5	-10.1	52.6
Relative to Nasdaq	-1.8	-6.6	34.0
Biotechnology Index			

Market data

Current Stock Price (\$)	91.26
Market Cap (\$mn)	4,648.5
52-week Low (\$)	60.19
52-week High (\$)	104.09
Shares Outstanding (mn)	50.94

Source: Thomson Reuters, Instinet research

Valuation

Year-end: Dec	2016A	2017E	2018E
EV/Sales (€k)	N/A	N/A	N/A

Source: Company data, Instinet estimates

Summary Income Statement

Year-end: Dec; €k	2016A	2017E	2018E
Revenue	151,612	149,898	183,696
Income Tax	235	265	0
Net Income (adj.)	65,046	-103,143	-102,245
GAAP EPS	1.14	-2.39	-2.40
EPS (adj.)	1.37	-2.04	-2.00
Diluted Shares (k)	47,308	50,476	53,758

Summary Balance Sheet

2016A	2017E	2018E
973,241	1,152,929	1,055,422
14,961	31,282	53,553
1,083,338	1,279,348	1,204,112
63	63	63
324,637	327,956	354,965
758,701	951,392	849,146
1,083,338	1,279,348	1,204,112
	973,241 14,961 1,083,338 63 324,637 758,701	973,241 1,152,929 14,961 31,282 1,083,338 1,279,348 63 63 324,637 327,956 758,701 951,392

Summary Cash Flow Statement

€k	2016A	2017E	2018E
Cash from Operations	239,405	-96,083	-67,416
Change in Working Capital	234,450	3,319	27,009
Cash from Investing	-7,287	-20,061	-30,092
Capital Expenditures	-4,458	-20,061	-30,092
Cash from Financing	395,996	295,833	0
Free Cash Flow	234,450	3,319	27,009

Other Metrics

	2016A	2017E	2018E
Enterprise Value (€k)	N/A	N/A	N/A

Fig. 1: Filgotinib Monotherapy vs. Upadacitinib – Efficacy and Safety Results

Filgotinib (12 wk)	ACR 20	ACR 50	ACR 70	Serious AE
Placebo	31%	11%	4%	1%
50 mg QD	67%	36%	8%	1%
100 mg QD	66%	34%	19%	0%
200 mg QD	73%	44%	13%	4%

Upadacitinib (14 wk)	ACR 20	ACR 50	ACR 70	Serious AE
MTX	41%	15%	3%	3%
15 mg QD	68%	42%	23%	5%
30mg QD	71%	52%	33%	3%

Grey shadow indicates statistically significant response MTX = methotrexate

Source: Company data, Instinet research

Fig. 2: Adverse Events Across JAK Inhibitors in the Clinic

	Filgotinib			Baricitinib		Upadacitinib					
% patients with reduction in								3 mg	6 mg	12 mg	18 mg
neutrophil count	Placebo	100 mg	200 mg	Placebo	2 mg	4 mg	Placebo	BID	BID	BID	BID
Grade 1	1.6%	0.6%	3.9%	<1.0%	2.0%	5.0%					
Grade 2	2.4%	0.0%	3.2%	<1.0%	2.0%	3.0%	2.0%	0.0%	6.0%	7.0%	13.0%
Grade 3	0.0%	0.6%	0.7%	0.0%	<1%	0.0%	0.0%	0.0%	0.0%	4.0%	2.0%
Grade 4	0.0%	0.6%	0.0%				0.0%	0.0%	0.0%	2.0%	0.0%
% patients with decreased Hb											
Grade 1	22.5%	11.0%	11.0%	15.0%	16.0%	18.0%					
Grade 2	3.9%	2.9%	1.9%	3.0%	8.0%	5.0%	0.0%	6.0%	9.0%	13.0%	20.0%
Grade 3	0.0%	0.0%	0.0%	0.0%	<1.0%	0.0%	0.0%	6.0%	0.0%	9.0%	11.0%
Grade 4	0.0%	0.0%	0.0%	<1.0%	0.0%	0.0%	0.0%	0.0%	7.0%	4.0%	4.0%

Source: Company data, Instinet research

Fig. 3: Safety Filgotinib vs. Other JAK Pathway Inhibitors

	Tofacitinib Phase 3			Baricitinib -Phase 3			Filgotinib Phase 2b			Upadacitinib Phase 2b			
	Placebo	5 mg	10 mg	Placebo	2 mg	4 mg	Placebo	100 mg	200 mg	Placebo	12 mg BID	18 mg BID	24 mg QD
Total AE	54.9%	51.0%	56.7%	56.5%	57.0%	63.5%	46.8%	38.7%	51.6%	35.0%	62.0%	60.0%	35.0%
Serious AE	4.9%	0.4%	2.0%	4.0%	2.0%	4.0%	3.9%	2.6%	2.5%	1.0%	6.0%	3.0%	4.0%
Dicontinuation rate due to AE	4.1%	0.8%	2.4%	3.0%	3.5%	4.5%	4.7%	3.2%	2.6%	3.0%	3.0%	7.0%	2.0%
Malignancies (no. of cases)	1	2	3	0	0	3	0	0	0	0	0	0	0
Serious infection	0.0%	0.0%	0.4%	1.0%	1.0%	2.0%	0.8%	1.9%	1.1%	1.0%	1.0%	0.0%	0.0%
Notes	Boxed warning contains serious infections, opportunistic infections, and malignancy. LDL levels were increased			experienced reduction neutrophil						Infections we	ere the most o	ommon AE	

Source: Company data, Instinet research

Galapagos NV Opt-In on Co-Promotion of Filgotinib w/ GILD in 8 EU Countries Supports Filgotinib Market Potential

Last Friday, GLPG announced a decision to opt in to co-promote filgotinib with partner GILD in 8 EU countries (Germany, France, Italy, Spain, the UK, Netherlands, Belgium, and Luxembourg).

Model Updates

- Outside these territories, Galapagos will be eligible for tiered royalties ranging from 20% to 30% on net filgotinib sales.
 - We anticipated GLPG would opt in to the co-promote, given its desire to build a commercial organization and cash war chest.
- Deal Terms Favorable, 35% Co-promotion Costs, 50/50 Profit Split; Model Updates.
 - In the eight European countries mentioned, GLPG will assume 35% of the promotional efforts and split profits equally with GILD.
 - For Netherlands, Belgium, and Luxembourg (Belenux region), GLPG will book the sales. Accordingly, we have made updates to our model estimates and recording format (see Figs. 4-6).

Co-Promotion Lock-In Instills Confidence In GILD Collab, Ahead of Equity Investment Lock-Up Expiration Date.

- The impending expiration of GLPG's lockup and standstill agreement with GILD is December 31, 2017.
- We believe that GLPG's decision to opt in on co-promotion of filgotinib ahead of the lockup expiration date speaks to management's confidence in the potential of its highly selective JAK1 inhibitor.

Fig. 4: GLPG Income Statement

(€1000s, except per share data) [FY - Dec]	2015	2016	1Q17A	2Q17A	3Q17A	4Q17E	2017E	1Q18E	2Q18E	3Q18E	4Q18E	2018E	2019E	2020E	2021E	2022E	2023E	2024E	2025E
US Filgotinib Sales (RA Only) Intl Filgotinib Sales (RA Only)	0	0	0	0	0	0	0	0	0	0	0	0	0	19,225	70,246 22,601	169,192 60,632	243,418 114,366	250,149 161,074	257,066 243,060
US Filgotinib Sales (Crohn's)	0	0	0	0	0	n.	0	0	0	0	0	0	0	0	40,245	82,717	127,506	174,708	179,539
Intl Filgotinib Sales (Crohn's)	0	0	0	0	0	0	0	0	0	0	0	o o	0	0	0	14,817	29,811	59,981	60,341
US Filgotinib Sales (UC only)	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	4,799	11,095	25,338	26,039
Intl Filgotinib Sales (UC only)	ő	0	0	0	0	0	0	0	0	0	0	0	0	0	0	4,733	4,869	11,896	26,751
Total Filgotinib Royalties/Profit Share	0	0	0	0	0	0	0	0	0	0	0	0	0	19,225	133,093	332,156	531,067	683,146	792,795
US GLPG- Triple Combo Royalties (CF)	0		0	0	0			0	0	0	0			0	7,314	111,645	181,772	184.966	188,217
Intl GLPG-Triple Combo Royalties (CF)	0	0	0	0	0	0	0	0	0	0	0	0	0	0	7,314	23,103	46,995	101,570	159,513
Total GLPG1837+Corrector Royalties	0	0	0	0	0	0	0	0	0	0	0	0	0	0	7,314	134,748	228,767	286,536	347,730
													_	_					
Fligotinib (Belenux Region)													0	0	5,909	19,725	38,967	60,902	86,314
R&D revenue	39,563	129,519	33,992	26,933	26,945	37,166	125,036	37,391	29,626	29,640	40,883	137,540	115,383	117,690	120,044	91,834	62,447	31,848	32,485
Other Income	21,017	22,093	5,871	6,235	6,378	6,378	24,862	6,442	6,506	6,571	26,637	46,156	84,862	94,862	94,862	94,862	94,862	94,862	94,862
Total Revenues	60,579	151,612	39,863	33,168	33,323	43,544	149,898	43,833	36,132	36,211	67,520	183,696	200,245	231,778	355,314	653,600	917,143	1,096,392	1,267,872
Costs & Expenses:			0		0	0		0	0	0			0		205		4 0 4 0	0.045	4040
Cost of Goods Sold	400.744	100 570		0	-	ŭ	0	•	-		0	0	· ·	004.000	295	986	1,948	3,045	4,316
R&D	129,714 19,127	139,573	44,930	47,984 6,327	56,313	61,944	211,169	65,041	68,293	70,342	71,045	274,721	288,457	294,226 43,557	300,111 47,912	306,113	312,235 52,572	318,480	324,849 56,856
G&A		21,744	5,603		5,853	6,438	24,220	6,760	7,098	7,453	7,825	29,135	37,875			50,308		55,200	11,530
Sales and Marketing	1,182	1,785	556	535	808	606	2,504	2,504	2,504	2,504	2,504	2,629	2,761	2,899	3,490	4,871	6,795	8,989	
Collaboration Split (Belenux Region)	0	0	0	0	0	0	0					0	0	0	1,935	6,460	12,762	19,945	28,268
Total Operating Expenses	150,023	163,103	51,088	54,845	62,973	68,988	237,894	74,305	77,895	80,298	81,374	306,485	329,093	340,681	353,743	368,738	386,312	405,660	425,819
Operating Income	(89,444)	(11,491)	(11,225)	(21,677)	(29,650)	(25,443)	(87,995)	(30,472)	(41,762)	(44,088)	(13,854)	(122,789)	(128,848)	(108,904)	1,571	284,862	530,831	690,732	842,053
Interest and Other Income (Expense), net	(30,632)	57,479	0	0	-	0	0	0	0	-	0	0	0	0	0	0	0	0	0
Interest Income	1,987	9,950	894	1,425	1,345	1,345	5,008	1,352	1,302	1,202	1,152	5,008	5,008	5,008	5,008	5,008	5,008	5,008	5,008
Interest Expense	(1,539)	(1,692)	(3,274)	(15,299)	(8,232)	(8,232)	(35,037)	(9,460)	(9,110)	(8,409)	(8,059)	(35,037)	(35,037)	(35,037)	(35,037)	(35,037)	(35,037)	(35,037)	(35,037
Other Income (Expense)	0						0					0	0	0	0	0	0	0	0
Pretax Income (Loss)	(119,628)	54,246	(13,605)	(35,552)	(36,538)	(32,331)	(118,025)	(38,580)	(49,570)	(51,295)	(20,761)	(122,789)	(128,848)	(108,904)	1,571	284,862	530,831	690,732	842,053
Income tax expense (Benefit)	(1,218)	235	0	92	69	104	265	0	0	0	0	0	0	0	0	0	0	27,629	57,260
Net Income (Loss) as reported	(118,410)	54,012	(13,605)	(35,644)	(36,607)	(32,434)	(118,289)	(38,580)	(49,570)	(51,295)	(20,761)	(122,789)	(128,848)	(108,904)	1,571	284,862	530,831	663,103	784,793
Stock option expense	5,036	11,034	3,023	3,945	4,729	3,449	15,146	3,715	3,895	6,424	6,510	20,544	23,037	23,848	24,762	25,812	27,042	28,396	29,807
Other	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
Net Income (Loss) Non-GAAP	(113,374)	65,046	(10,582)	(31,699)	(31,878)	(28,985)	(103,143)	(34,864)	(45,675)	(44,871)	(14,251)	(102,245)	(105,812)	(85,056)	26,333	310,674	557,872	691,499	814,600
Diluted Earnings Per Share Non-GAAP	(€3.18)	€1.37	(€0.22)	(€0.61)	(€0.63)	(€0.57)	(€2.04)	(€0.65)	(€0.85)	(€0.83)	(€0.26)	(€2.00)	(€2.07)	(€1.66)	€0.49	€5.76	€10.33	€12.79	€15.05
Earnings Per Share, Diluted Fully Taxed	€3.32	€ 0.77					nm					nm	nm	nm	€0.02	€3.61	€6.71	€8.72	€10.62
Basic Earnings Per Share Non-GAAP	(€2.94)	€1.42	(€ 0.23)	(€0.64)	(€0.63)	(€0.57)	(€2.08)	(€0.68)	(€0.89)	(€0.88)	(€0.28)	(€2.00)	(€2.07)	(€1.66)	€ 0.51	€6.05	€10.85	€13.44	€ 15.81
Diluted Earnings Per Share	(€3.32)	€1.14	(€0.29)	(€0.72)	(€0.72)	(€0.64)	(€2.39)	(€0.76)	(€0.97)	(€1.00)	(€0.41)	(€2.40)	(€2.52)	(€2.12)	€0.03	€5.28	€9.83	€12.26	€14.50
Basic Earnings Per Share as reported	(€3.08)	€1.18	(€ 0.29)	(€ 0.72)	(€0.72)	(€0.64)	(€2.39)	(€ 0.76)	(€0.97)	(€1.00)	(€0.41)	(€2.40)	(€2.52)	(€2.12)	€0.03	€5.55	€10.32	€12.88	€ 15.23
Basic Shares Outstanding (th)	35,700	45,696	46,256	49,830	50,902	51,029	49,504	51,080	51,131	51,182	51,234	51,157	51,208	51,259	51,311	51,362	51,413	51,465	51,516
Diluted Shares Outstanding (th)	38,506	47,308	48,330	51,644	50,902	51,029	50,476	53,881	53,932	53,983	54,034	53,758	53,809	53,860	53,911	53,963	54,014	54,065	54,117

Fig. 5: GLPG Balance Sheet

(€1000s, except per share data) [FY - Dec]	2015A	2016A	2017E	2018E	2019E	2020E	2021E	2022E	2023E	2024E	2025E
ASSETS	_0.000	_,,,,,									
Current assets:											
Cash and cash equivalents	340,314	973,241	1,152,929	1,055,422	905,996	759,299	700,651	904,029	1,304,699	1,719,646	2,248,808
Current restricted cash	6,857	6,570	6,570	6,570	6,570	6,570	6,570	6,570	6,570	6,570	6,570
Current R&D incentives receivables	9,161	10,154	10,154	10,154	10,154	10,154	10,154	10,154	10,154	10,154	10,154
Current financial assests from share subscription agreemen	8,371	0			•		•				
Short term marketable securities	0	0.700	0	0	0	0	0	0	0	0	0
Trade & other receivables	3,931	9,728	9,728	9,728	9,728	9,728	9,728	9,728	9,728	54,820	63,394
Inventory	325	300	300	300	300	300	300	300	300	300	300
Prepaid expenses and other current assets	5,512	7,239	7,239	7,239	7,239	7,239	7,239	13,072	18,343	21,928	25,357
Total current assets	374,470	1,007,232	1,186,920	1,089,413	939,987	793,290	734,642	943,853	1,349,794	1,813,418	2,354,583
Property and equipment, net	13,782	14,961	31,282	53,553	85,302	131,682	200,321	302,579	455,441	684,342	958,846
Goodwill	0	0	0	0	0	0	0	0	0	0	0
Intangible assets	1,550	1,023	1,023	1,023	1,023	1,023	1,023	1,023	1,023	1,023	1,023
Deferred tax assets/receivables	1,726	1,957	1,957	1,957	1,957	1,957	1,957	1,957	1,957	1,957	1,957
Non-current R&D incentives receivables	49,384	54,188	54,188	54,188	54,188	54,188	54,188	54,188	54,188	54,188	54,188
Non-current restricted cash	1,046	1,098	1,098	1,098	1,098	1,098	1,098	1,098	1,098	1,098	1,098
Other non-current assets	557	2,880	2,880	2,880	2,880	2,880	2,880	2,880	2,880	2,880	2,880
Total assets	442,514	1,083,338	1,279,348	1,204,112	1,086,435	986,118	996,109	1,307,577	1,866,381	2,558,906	3,374,575
LIABILITIES AND STOCKHOLDER'S EQUITY											
Current liabilities:											
Trade and other payables	29,482	31,269	34,494	61,297	49,364	34,068	17,687	18,437	19,316	20,283	21,291
Current obligations under finance lease	52	54	54	54	54	54	54	54	54	54	54
Current tax payable	2,583	1,022	1,022	1,022	1,022	1,022	1,022	1,022	1,022	1.022	1,022
Accrued charges	490	619	714	919	987	1,022	1,061	1,106	1,159	1,217	1,277
Deferred income	39,806	70,827	70,827	70,827	70,827	70,827	70,827	70,827	70,827	70,827	70,827
Other current liabilities	03,000	70,027	70,027	70,027	70,027	70,027	70,027	10,021	10,021	70,027	70,027
Total current liabilities	72,412	103,791	107,110	134,119	122,254	106,993	90,651	91,446	92,378	93,403	94,471
Total current habilities	12,412	103,731	107,110	134,113	122,204	100,333	30,031	31,770	32,370	33,403	34,471
Long term debt	0	0	0	0	0	0	0	0	0	0	0
Obligations under finance lease	63	9	9	9	9	0	9	9	9	9	0
•	0	-	-	ŭ	ŭ	044705	-	•	ľ	Ü	04.4.705
Deferred Revenue	-	214,785	214,785	214,785	214,785	214,785	214,785	214,785	214,785	214,785	214,785
Provisions	55	63	63	63	63	63	63	63	63	63	63
Pension liabilities	2,693	3,520	3,520	3,520	3,520	3,520	3,520	3,520	3,520	3,520	3,520
Other liabilities	2,291	2,469	2,469	2,469	2,469	2,469	2,469	2,469	2,469	2,469	2,469
Total liabilities	77,515	324,637	327,956	354,965	343,100	327,839	311,497	312,292	313,224	314,249	315,317
Stockholders' equity:											
Common Stock	185,399	223,928	223,928	223,928	223,928	223,928	223,928	223,928	223,928	223,928	223,928
Additional paid in capital	357,402	649,135	944,968	944,968	944,968	944,968	944,968	944,968	944,968	944,968	944,968
Other reserves	(18)	(1,000)	(1,000)	(1,000)	(1,000)	(1,000)	(1,000)	(1,000)	(1,000)	(1,000)	(1,000)
Translation differences	(467)	(1,000)	(1,000)	(1,000)	(1,000)	(1,000)	(1,000)	(1,000)	(1,000)	(1,000)	(1,000)
Accumulated other comprehensive loss	(107)	(1,090)	(1,090)	(1,090)	(1,090)	(1,090)	(1,090)	(1,090)	(1,090)	(1,090)	(1,090)
Accumulated Deficit	(177,319)	(112,272)	(215,415)	(317,660)	(423,472)	(508,528)	(482,195)	(171,521)	386,351	1,077,850	1,892,451
Total stockholders' equity	364,999	758,701	951,392	849,146	743,335	658,279	(482, 193) 684,611	995,285	1,553,157	2,244,657	3,059,257
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Total liabilities and stockholders' equity	442,514	1,083,338	1,279,348	1,204,112	1,086,435	986,118	996,109	1,307,577	1,866,381	2,558,906	3,374,575

Fig. 6: GLPG Cash Flow Statement

(€1000s, except per share data) [FY - Dec]	2015A	2016A	2017E	2018E	2019E	2020E	2021E	2022E	2023E	2024E	2025E
CASH FLOWS FROM OPERATING ACTIVITIES											
Net Income (Loss)	(118,410)	54,012	(118,289)	(122,789)	(128,848)	(108,904)	1,571	284,862	530,831	663,103	784,793
Adjustments											
Tax income/expenses	(1,218)	235	0	0	0	0	0	0	0	0	0
Other net financial income	(448)	(8,258)	0	0	0	0	0	0	0	0	0
Fair value measurment of share subscription	30,632	(57,479)	0	0	0	0	0	0	0	0	0
Depreciation and amortization	3,402	4,182	3,740	7,820	13,388	21,325	32,921	50,080	75,645	113,860	171,085
Net realized loss for foreign exchange transaction	(398)	1,229	0	0	0	0	0	0	0	0	0
Stock based compensation	5,036	11,034	15,146	20,544	23,037	23,848	24,762	25,812	27,042	28,396	29,807
Other	0		0	0	0	0	0	0	0	0	0
Change in assets and liabilities:											
Increase/decrease in provisions	(125)	7	0	0	0	0	0	0	0	0	0
Increase pension liabilities	` 30	244	0	0	0	0	0	0	0	0	0
Gain on sale of fixed assests	(62)	(14)	0	0	0	0	0	0	0	0	0
Inventories	(44)	25	0	0	0	0	0	0	0	0	0
Account receivables	(7,220)	(12,978)	0	0	0	0	0	0	0	(45,092)	(8,574)
Prepaid expenses & other assets	0	0	0	0	0	0	0	(5,833)	(5,271)	(3,585)	(3,430)
Accounts payable and accrued expenses	(26,728)	2,102	3,319	27,009	(11,865)	(15,261)	(16,342)	795	931	1,025	1,068
Interest paid	(49)	(47)	0	0	0	0	0	0	0	0	0
Interest received	1,106	1,066	0	0	0	0	0	0	0	0	0
Income taxes paid/received	(94)	(1,763)	0	0	0	0	0	0	0	0	0
Current obligations under finance lease	0	2	0	0	0	0	0	0	0	0	0
Deferred revenues & other	0	245,806	0	0	0	0	0	0	0	0	0
Net cash provided by (used in) operating activities	(114,590)	239,405	(96,083)	(67,416)	(104,289)	(78,992)	42,912	355,716	629,178	757,709	974,751
Purchase of and expenditure of intangible fixed assets	(565)	(332)		0	0	0	0	0	0	0	0
Proceeds from disposal of PPE	110	18	0	0	0	0	0	0	0	0	0
Increase/decrease in restricted cash	2,258	235	0	0	0	0	0	0	0	0	0
Investments, net Other	0	(2,750)	0	0	0	0	0	0	0	0	0
Net cash used in investing activities	(4,297)	(7,287)	(20,061)	(30,092)	(45,137)	(67,706)	(101,559)	(152,338)	(228,507)	(342,761)	(445,589)
iver cash used in investing activities	(4,237)	(1,201)	(20,001)	(30,092)	(43,137)	(01,100)	(101,559)	(132,330)	(220,301)	(342,701)	(443,309)
CASH FLOWS FROM FINANCING ACTIVITIES											
Proceeds from issuance of shares, net cost	271,413	391,784	295,833	0	0	0	0	0	0	0	0
Exercise of options	0	4,261	0	0	0	0	0	0	0	0	0
Repayment obligations under finance and other debt	(43)	(49)	0	0	0	0	0	0	0	0	0
Repurchase of common stock	0	0	0	0	0	0	0	0	0	0	0
Other	0		0	0	0	0	0	0	0	0	0
Net cash provided by financing activities	271,370	395,996	295,833	0	0	0	0	0	0	0	0
Effect of exchange rate on cash	118	4,816									
Net increase in cash and cash equivalents	152,483	632,927	179,689	(97,507)	(149,426)	(146,698)	(58,647)	203,377	400,670	414,948	529,161
Cash and cash equivalents at beginning of period	187,712	340,314	973,241	1,152,929	1,055,422	905,996	759,299	700,651	904,029	1,304,699	1,719,646
Cash and cash equivalents at end of period	340,314	973,241	1,152,930	1,055,422	905,996	759,299	700,651	904,029	1,304,699	1,719,646	2,248,808

Appendix A-1

Analyst Certification

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Materially mentioned issuers

Issuer	Ticker	Price	Price date	Stock rating	Sector rating	Disclosures
Galapagos NV	GLPG US	USD 91.26	19-Dec-2017	Buy	Not rated	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) USD 91.26 (19-Dec-2017) Buy (Sector rating: Not rated) Rating and target price chart (three year history) 19-Dec-2014 to 18-Dec-2017 130.00 PRICE(USD) 120.00 110.00 100.00 90.00 80.00 70.00 60.00 50.00 30.00 20 00 10.00 Jan-15 Apr-15 Jul-15 Oct-15 Jan-16 Apr-16 Jul-16 Oct-16 Jan-17 Apr-17 Jul-17 - GALAPAGOS NV 🛕 Target Price Change 🌘 Recommendation Changes Source: Thomson Reuters, Nomura research

Date	Rating	Target price	Closing price
10-Aug-17		124.00	81.06
22-Jun-17		108.00	78.56
17-Apr-17		121.00	92.40
01-Mar-17	Buy		71.78
01-Mar-17		87.00	71.78
27-Oct-15	Not Rated		47.69
25-Sep-15		62.00	44.60
08-Jun-15	Buy		55.51
08-Jun-15		73.00	55.51

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

Valuation Methodology For Galapagos NV (GLPG), we use a top-line revenue multiple valuation, a method widely used for early-stage biotech companies. Our target price of \$124 represents a 6x multiple for EU profit share on filgotinib across inflammatory indications and a 16x multiple for U.S. 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, as 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. For the IPF program, we use an 8x multiple, reflecting a higher value for the higher-margin orphan program and a 45% discount that reflects a higher development risk. The benchmark for this stock is the Nasdaq Biotechnology Index.

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 the company 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 on 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, which would affect commercial viability. 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 could take the lion's share of the market. Celgene's mongersen, an 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. This difference in design may result in a smaller efficacy difference between the placebo and treatment arms in the Phase 3 study.

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As at 30 September 2017.

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