

**Galapagos NV (GLPG)**  
**Rating: Buy**

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### SELECTION Phase 3 Readout a Couple of Weeks out Plus Our Cheat Sheet on Historic UC Phase 3 Outcomes

Stock Data	05/08/2020
Price	\$217.12
Exchange	NASDAQ
Price Target	\$302.00
52-Week High	\$274.03
52-Week Low	\$110.92
Enterprise Value (M)	\$7,893
Market Cap (M)	\$14,074
Shares Outstanding (M)	64.8
3 Month Avg Volume	191,976
Short Interest (M)	1.21

Balance Sheet Metrics	
Cash (M)	\$6,180.2
Total Debt (M)	\$0.0
Total Cash/Share	\$95.35

General: Currency used is roughly 1 Euro to \$1.08 US. Stock price is US\$ as on NASDAQ

EPS (€) Diluted			
Full Year - Dec	2019A	2020E	2021E
1Q	€(0.89)	€(0.78)A	--
2Q	€(0.86)	€(1.10)	--
3Q	€5.83	€(1.33)	--
4Q	€(1.86)	€1.62	--
FY	€2.49	€(1.58)	€1.80

Revenue (€)			
Full Year - Dec	2019A	2020E	2021E
1Q	€40.9	€106.9A	--
2Q	€67.6	€103.9	--
3Q	€644.0	€103.9	--
4Q	€143.4	€289.1	--
FY	€895.9	€603.9	€874.5



**What's new post 1Q20?** COVID-19 has negatively impacted a few programs, a common refrain for most non-oncology programs. However, 2020 is far from dull with a surfeit of potentially platform and Gilead (GILD; not rated) licensing deal-validating late and mid-stage readouts. Key upcoming deliverables include: (1) topline data from the Phase 3 SELECTION trial, anticipated prior to the end of 2Q20; (2) filgotinib approvals in the U.S., EU, and Japan, in 2H20; (3) Phase 2a data from the PINTA program, i.e., GLPG1205 in IPF during 2H20. Recall, approximately 69 IPF patients have been enrolled in the PINTA, which is evaluating GLPG1205 atop nintedanib or pirfenidone, 33% each, while the remainder are on local standard-of-care; (4) Phase 2a readout in systemic scleroderma, i.e., the NOVESA study of ziritaxestat, during 2H20; and (5) ROCCELLA, the 850+ patients Phase 2a study in OA for GLPG1972, in 2H20. Readouts negatively impacted by COVID-19, include: (1) futility analysis from the ISABELA program pushed into 1H21, about three months behind prior schedule. Note, 1,000 patients been recruited into the dual Phase 3 programs as of April 2020; and (2) first look at the TOLEDO platform, now with GLPG3970 as the lead as opposed to GLPG3312, expected during 1H21, as opposed to 2H20. The switch back to -3970 and futility analysis pushout caused significant hand wringing on May 8, 2020, pulling the stock down 4% while the comp group was up 1.5%, following the 1Q20 print. However, we are not about to second guess our Buy rating on the stock, given the near-to-intermediate term calendar.

**Looking past biologics in UC.** While biologics are the mainstay under current therapeutic algorithms: (1) the cumulative relapse rates vary between 67% and 83% after 10 years; (2) a high percentage of patients have an inadequate response with about 30% being characterized as primary non-responders and a further 50% lose response over the time after initial remission, i.e., secondary non-responders, or develop adverse events that lead to discontinuation. Hence, there is a group of patients in whom remission cannot be achieved and surgery is necessary to control the disease. A recent meta-analysis indicated five- and 10-year risk of surgery at 11.6% and 15.6%, respectively. In North America, the prevalence of UC ranges from 120 to 250 cases per 100,000 people, and the incidence ranges from 8 to 20 cases per 100,000 people. Patients can be diagnosed with UC at any age; however, the disease has an age distribution characterized by two peaks of incidence at 15 to 30 years and 50 to 70 years. Filgotinib, a JAK1i with a cleaner safety profile compared to the other pan- and selective JAKi's (refer to *May the Force Be With Filgotinib; Raising Target to \$150*), could position it as the preferred agent in IBD compared to biologics based on: (1) oral administration, which increases treatment compliance; (2) quick onset of action; (3) lack of immunogenicity; (4) short half-life, (5) linear PK; and (6) a differentiated safety profile from its three RA-centric Phase 3 studies complemented by numerous other Phase 2 programs.

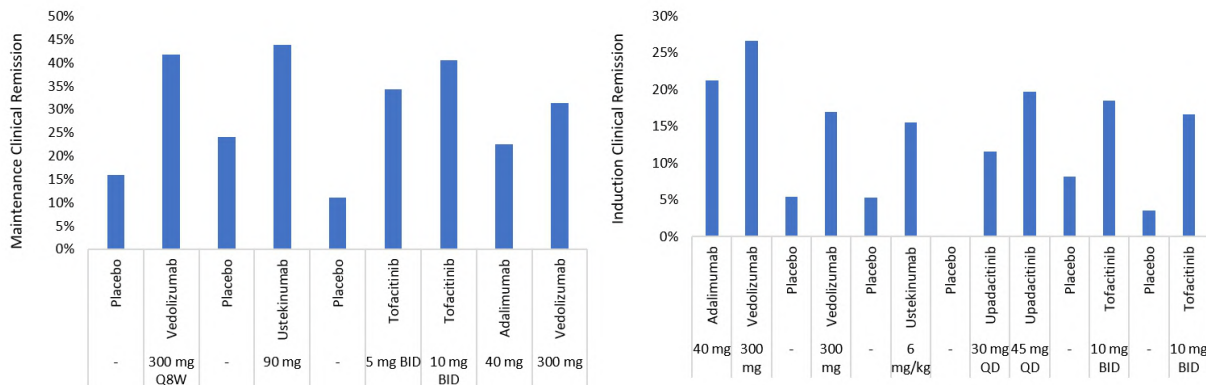
H.C. Wainwright 1868

**We expect the SELECTION study to readout positively.** (1) the prior history of JAKi's; and (2) compelling filgotinib data from CD, i.e., the FITZROY study. Recall, patients in the FITZROY trial were required to have a total SES-CD  $\geq 7$  at trial entry with ulceration scores of  $\geq 1$  in at least one ileocolonic segment. This to us remains a key differentiator vs. the Pfizer (PFE; not rated) experience with Xeljanz in the same indication, where enrollment of mild to moderate patients potentially doomed the trial. Furthermore, the placebo remission rate in the FITZROY program was in line with successful biologics, i.e., about 23% and endoscopic placebo remission rates were 2-7%. Hence, FITZROY success was not due to an underperformance of the placebo cohort, but reflects an active and selective JAK1i; (3) these successes mimicked in to other indications including PsA, and AS, for which registration studies are underway with readouts in 2021 and beyond.

**Baseline characteristics matter in UC.** An analysis of multiple prior Phase 3 programs in UC lead us to the following observation with regard to patient characteristics that either trended, or were significantly associated, with higher clinical remission rates compared with placebo: (1) gender, males have a better response; (2) age, patients older than 41 responded better; (3) baseline or concomitant corticosteroid use, users responded better; (4) baseline or concomitant immunosuppressant use, users responded better; (5) prior anti-TNF exposure, no exposure equates to better response; and (6) baseline hs-CRP level,  $<5$  mg/L correlates with better outcomes. Surprisingly, baseline Mayo score and duration of disease did not improve clinical remission rates relative to placebo.

**Normalized outcomes are consistent across multiple Phase 3 programs.** Clinical remission rates increases from a range of 15% to 20% during induction therapy to 25% to 40% during maintenance therapy, see Exhibit 1, 2, and 3. Oral JAKi appeared to have similar efficacy compared with biologics after adjusting for patient baseline characteristics. For example, on paper adalimumab and vedolizumab appear superior to upadacitinib and tofacitinib induction therapies, but the percent of patients with prior anti-TNF use in the VARSITY trial was only 21%, whereas in the studies with JAKi these rates were markedly higher, i.e., from 53% to 79%, implying a more advanced population. These, in turn, might have contributed to the higher infection rates compared with the VARSITY study. A concerning trend we found with upadacitinib is the higher rates of infections and cardiovascular events, relative to tofacitinib and mAbs; however, these rates are derived from a relatively small number of patients, and could evolve with more patients. We note the inclusion and exclusion criteria for the Phase 3 SELECTION 1 study of filgotinib in UC are the same as peers, and use the same primary endpoints, see Exhibit 4. One metric to watch we think is the rates of Herpes Zoster (HZ) infection rates in the SELECTION program. Recall, in a pooled analysis of Phase 2, 3, maintenance, and OLE global tofacitinib data on 1,157 UC patients, which encompasses about 1,612.8 patient-years (PY) of exposure, indicated that 5.6% patients developed 69 events of HZ infection with incidence rate of 4.1 per 100 PY. The risk of HZ with tofacitinib 10 mg BID was higher (6.6 per 100 PY; 95% CI 3.2–12.2), compared with 5 mg BID (2.1 per 100 PY; 95% CI 0.4–6.0), and placebo (1.0 per 100 PY; 95% CI 0–5.4), suggesting a dose-response relationship.

#### Exhibit 1: Clinical Remission From Maintenance (Left) and Induction (Right) Therapy in UC



Source: H.C. Wainwright & Co. research adapted from *N Engl J Med* 2017; 376:1723-1736 DOI: 10.1056/NEJMoa1606910, *Gastroenterology*. 2020 Feb 22. pii: S0016-5085(20)30241-9. doi: 10.1053/j.gastro.2020.02.030.

**Exhibit 2: Detailed Comparison of Moderate to Severe UC Clinical Trials for Maintenance Therapy**

Clinical Trials	GEMINI 1		UNIFI		Octave Sustain			VARSITY,	
	-	300 mg Q8W	-	90 mg	-	5 mg BID	10 mg BID	40 mg	300 mg
Drug	Placebo	Vedolizumab	Placebo	Ustekinumab	Placebo	Tofacitinib	Tofacitinib	Adalimumab	Vedolizumab
Study Phase	3	3	3	3	3	3	3	3b	3b
% Concomitant Corticosteroids	57%	57%	54%	54%	51%	51%	44%	36%	36%
Median Age	40.3	41	42	39.5	43.4	41.9	42.9	40.5	40.8
% Male	55%	57%	61%	53%	59%	52%	56%	56%	61%
Baseline weight	74.7	78.2	71.7	72				73.4	72.7
Baseline CRP mg/L			3.4	4	1	0.7	0.9		
% Prior anti-TNF Use	37%	41%	50%	51%	47%	46%	51%	21%	21%
<b>Safety</b>									
Safety Patient N	126	122	175	176	198	198	197	386	385
% with SAE	16%	8%	10%	9%	7%	5%	6%	14%	11%
% Any Infections	71%	71%	49%	46%	24%	36%	40%	4%	5%
% Cardiovascular Events					0%	1%	1%	0%	0%
% Malignancy	2%	1%	1%	0%	1%	0%	2%		
<b>Efficacy</b>									
Patient N	126	122	175	176	198	198	197	386	385
Maintenance Endpoint (Weeks)	52	52	52	52	52	52	52	52	52
% Achieving 6-Month Steroid Free Maintenance	14%	31%	23%	42%					
% Clinical Remission Mayo, Maintenance	16%	42%	24%	44%	11%	34%	41%	23%	31%
% Histologic Remission Maintenance								3%	10%
% Achieving 6-Month Steroid Free Maintenance								22%	13%

Source: *N Engl J Med* 2013;369:699-710 DOI: 10.1056/NEJMoa1215734, *N Engl J Med* 2019;381:1201-14 DOI: 10.1056/NEJMoa1900750, *N Engl J Med* 2017; 376:1723-1736 DOI: 10.1056/NEJMoa1606910, *N Engl J Med* 2019;381:1215-26. DOI: 10.1056/NEJMoa1905725.

**Exhibit 3: Detailed Comparison of Moderate to Severe UC Clinical Trials for Induction Therapy**

Study	VARSITY		GEMINI 1		UNIFI		NCT02819635			Octave 1		Octave 2	
	40 mg	300 mg	-	300 mg	-	6 mg/kg	-	30 mg QD	45 mg QD	-	10 mg BID	-	10 mg BID
Drug	Adalimumab	Vedolizumab	Placebo	Vedolizumab	Placebo	Ustekinumab	Placebo	Upadacitinib	Upadacitinib	Placebo	Tofacitinib	Placebo	Tofacitinib
Study Phase	3b	3b	3	3	3	3	2b	2b	2b	3	3	3	3
% Pt Using Corticosteroids	36%	36%	56%	56%	49%	53%	54%	48%	50%	48%	45%	49%	46%
Median Age	40.5	40.8	41.2	40.1	41.2	41.7	40	42	37	41.8	41.3	40.4	41.4
% Male	56%	61%	62%	59%	62%	61%	63%	60%	66%	63%	58%	49%	60%
% White	88%	90%	77%	81%									
Baseline weight	73.4	72.7	72.4	72.4	72.9	73							
Baseline CRP mg/L					4.7	4.8	5.4	6.7	6.3	4.7	4.4	5	4.6
% Prior anti-TNF Use	21%	21%	49%	42%	50%	51%	48%	79%	70%	53%	53%	58%	55%
<b>Safety</b>													
Safety Patient N	386	385	149	225	319	320	46	52	56	122	476	112	429
% with SAE	14%	11%	7%	2%	7%	3%	11%	6%	5%	4%	3%	8%	4%
% Any Infections	4%	5%	15%	14%	15%	16%	35%	12%	23%	16%	23%	15%	18%
% Herpes Zoster							0%	0%	2%	1%	1%	0%	0.5%
% Cardiovascular Events	0%	0%					0%	0%	2%	0%	0.4%	0%	0.5%
<b>Efficacy</b>													
Patient N	386	385	149	225	319	322	46	52	56	122	476	112	429
Induction Endpoint (Weeks)	14	14	6	6	8	8				8	8	8	8
% Clinical Remission Mayo, Induction	21%	27%	5%	17%	5%	16%	0%	12%	20%	8%	19%	4%	17%
% Endoscopic Remission Induction							0%	10%	18%	2%	7%	2%	7%
% Histologic Remission Induction	3%	5%											

Source: *N Engl J Med* 2019;381:1215-26. DOI: 10.1056/NEJMoa1905725, *N Engl J Med* 2013;369:699-710 DOI: 10.1056/NEJMoa1215734, *N Engl J Med* 2017; 376:1723-1736 DOI: 10.1056/NEJMoa1606910, *Gastroenterology*. 2020 Feb 22. pii: S0016-5085(20)30241-9. doi: 10.1053/j.gastro.2020.02.030, *N Engl J Med* 2017; 376:1723-1736 DOI: 10.1056/NEJMoa1606910.

**Exhibit 4: Trial Design for Phase 2/3b SELECTION Program Evaluating Filgotinib in UC**

Component	SELECTION1
Phase	2b/3
Patient N	1351
Design	Randomized, Placebo Controlled
Doses	100 mg, 200 mg
Inclusion Criteria 1	Diagnosis of UC for at least 6 months with endoscopic and histopathologic evidence
Inclusion Criteria 2	Moderately to Severely Active UC
Inclusion Criteria 3	Inadequate clinical response to corticosteroids, immunomodulators, anti-TNF, or vedolizumab
Inclusion Criteria 4	Absence of Crohns disease, or other types of colitis
Induction Primary Endpoints	Percent Achieving Clinical Remission Based on Components of Mayo Clinic Score at Week 10
Secondary Endpoints	Percent Achieving Endoscopic Subscore of 0 at Week 10
Secondary Endpoints	Percent Achieving Histologic Remission at Week 10
Maintenance Secondary Endpoints	Percent Achieving Clinical Remission Based on Components of Mayo Clinic Score at Week 58
Secondary Endpoints	Percent Achieving 6-Month Corticosteroid-Free Remission Based on Components of MCS at Week 58
Secondary Endpoints	Percent Achieving Endoscopic Subscore of 0 at Week 58
Secondary Endpoints	Percent Achieving Histologic Remission at Week 58

Source: Adapted from [Clinicaltrials.gov](http://Clinicaltrials.gov).

**1Q20 housekeeping.** Galapagos reported a net loss of €50.6M vs. our estimate of €80.9M, with GAAP EPS of €0.78 vs. our estimate of €1.30. Total revenue for the quarter was €106.9M vs. our estimate of €103.0M. Galapagos exited the quarter with about \$6.2B (€5.7B) in cash and equivalents, which should be sufficient to fund operations for the foreseeable future. Recall, commercial launch of filgotinib in RA is expected during 4Q20, with additional indications to follow during 2021 and beyond. Due to the negative impact of COVID-19 on the clinical programs, cash burn guidance has been revised lower. FY20 operational cash requirement is now expected to be in the range of €400 and €430M, compared to €420 and €450M, previously. For FY20, we now expect a net loss of €103.0M vs. prior €212.8, and GAAP EPS of (€1.58) vs. prior (€3.40), the revision reflects potential revenue recognition from its Gilead partnership during 4Q20.

**Valuation and risks to our investment thesis.** We reiterate our Buy rating and 12-month price target of \$302 on shares of Galapagos. Our target is derived from a 12-year DCF-based, sum-of-the-parts analysis, which includes a beta of 1.41, terminal growth rate of -3.0%, risk premium of 4.93%, calculated WACC of 8.2%, and tax rate of 20% beginning in FY 2025. Filgotinib (82%), GLPG1690 and GLPG1972 (2% each) together make up about 86% of our value, with the remainder derived from the probability-adjusted, filgotinib-related milestone payments. For filgotinib, we assume POS in the range of: 80% for RA, 65% for UC, and 60% for CD, PsA and AS each, whereas for '1690 and '1972, we assign a 35% and 10% POS, respectively. Key risks include: emergence of safety concerns, clinical risks, regulatory risks, COVID-19 disruptions, and financial risks. Furthermore, regulatory and commercial strategy for filgotinib is under the control of partner, Gilead, not an established player in autoimmune indications. Hence, Gilead may not be able to drive rapid adoption of filgotinib, especially if the overall profile is relatively undifferentiated from AbbVie's (ABBV; not rated) upadacitinib, in our view. Hence, our estimates could be negatively impacted if AbbVie successfully leverages its market positioning with Humira during the launch of RINVOQ.

## Valuation: Galapagos (GLPG) Discounted Cash Flow (DCF) Analysis

		Discounted Cash Flow Analysis													TV
		2020	2021	2022	2023	2024	2025	2026	2027	2028	2029	2030	2031		
Ticker	Galapagos														
Period	2020E														
Beta est	1.41	EBIT (000s €)	€ (82,017)	€ 119,960	€ 570,153	€ 1,135,648	€ 1,816,857	€ 2,061,754	€ 2,545,424	€ 2,501,966	€ 2,645,740	€ 2,711,551	€ 2,738,748	€ 2,758,177	
Risk-free rate (R <sub>f</sub> )(10 yr yield)	1.25%	Tax rate	-0.3%	0.0%	0.0%	0.0%	0.0%	20.0%	20.0%	20.0%	20.0%	20.0%	20.0%	20.0%	
Risk premium (R <sub>e</sub> )	4.93%	EBIT*(1-t)	(82,286)	119,960	570,153	1,135,648	1,816,857	1,649,403	2,036,339	2,001,573	2,116,592	2,169,241	2,190,998	2,206,542	
Cost of equity (K <sub>E</sub> )	8.2%	Capital expenditures	(20,866)	(22,953)	(25,248)	(27,773)	(30,550)	(33,605)	(36,965)	(40,662)	(44,728)	(49,201)	(54,121)	(59,533)	
Cost of debt (K <sub>D</sub> )	0.0%	% growth	-6.8%	10.0%	10.0%	10.0%	10.0%	10.0%	10.0%	10.0%	10.0%	10.0%	10.0%	10.0%	
Terminal growth rate	-3.0%	Depreciation	14,689	17,627	18,508	19,434	20,405	21,425	22,497	23,622	24,803	26,043	27,345	28,712	
Terminal value (% of total value)	42.7%	% growth	21.5%	20.0%	5.0%	5.0%	5.0%	5.0%	5.0%	5.0%	5.0%	5.0%	5.0%	5.0%	
Shareholder equity	2,828,109	Change in non-cash working capital	65,981	65,999	70,057	74,341	78,863	22,250	11,434	17,710	19,038	23,570	29,793	36,650	
Debt outstanding	0	% growth	419.6%	0.0%	6.1%	6.1%	6.1%	-71.8%	-48.6%	54.9%	7.5%	23.8%	26.4%	23.0%	
Total capital	2,828,109	Free cash flow to the firm	(112,712)	94,540	543,851	1,108,513	1,788,949	1,682,183	2,084,367	2,048,147	2,167,085	2,220,915	2,242,671	2,258,137	19,554,806
Equity/cap	100.0%	Discount factor	0.97	0.92	0.85	0.79	0.73	0.67	0.62	0.58	0.53	0.49	0.45	0.42	
Debt/cap	0.0%	Present value of cash flows	(109,789)	87,374	464,531	875,071	1,305,173	1,134,255	1,298,911	1,179,597	1,153,496	1,092,546	1,019,626	948,840	7,795,845
WACC (calculated)	8.2%	Value of firm	18,245,476												
WACC (applied)	8.2%	Debt	0												
Shares outstanding	65,144	Value of equity	18,245,476												
		Value per share (\$)	\$ 302.00												

Source: H.C. Wainwright &amp; Co. estimates.



## Cash Flow Statement

Cash flow statement (€ in thousands, except per share data)	2018A	1Q19A	2Q19A	3Q19A	4Q19A	2019A	1Q20A	2Q20E	3Q20E	4Q20E	2020E	2021E	2022E	2023E	2024E	2025E	2026E	2027E	2028E	2029E	2030E	2031E	
<b>Cash flows from Operating activities:</b>																							
Net gain (loss)	€ (29,259)	€ (48,656)	€ (47,249)	€ 361,234	€ (115,484)	€ 149,845	€ (50,601)	€ (71,651)	€ (86,302)	€ 105,521	€ (103,033)	€ 117,613	€ 567,688	€ 1,133,060	€ 1,814,139	€ 1,647,120	€ 2,033,942	€ 1,999,056	€ 2,113,949	€ 2,166,466	€ 2,188,085	€ 2,203,483	
Adjustments for non-cash transactions, and items to disclose under operating cash flow	€ 17,364	€ 4,007	€ 16,407	€ 107,523	€ 112,359	€ 240,296	€ 22,188	€ 17,050	€ 17,550	€ 18,050	€ 74,838	€ 46,482	€ 51,541	€ 57,126	€ 63,292	€ 70,097	€ 77,608	€ 85,891	€ 95,031	€ 104,591	€ 115,683	€ 127,916	
Adjustment for items to disclose under investing and financing cash flows	(668)	(3)	0	0	(5,059)	(5,061)	(2,596)	0	0	0	(2,596)	(2,622)	(2,648)	(2,675)	(2,701)	(2,728)	(2,756)	(2,783)	(2,811)	(2,839)	(2,868)	(2,896)	
Change in working capital other than deferred income	19,322	(2,294)	(13,624)	57,044	(28,428)	12,698	52,481	5,000	3,500	5,000	65,381	65,998	70,657	74,341	77,863	22,250	11,434	17,710	19,038	23,570	29,790	38,650	
Decrease in deferred income	(153,312)	(25,979)	(27,499)	2,943,764	(86,084)	2,804,202	(91,677)	(100,000)	(100,000)	(100,000)	(391,677)	(391,677)	(391,677)	(391,677)	(175,000)	(175,000)	(150,000)	(150,000)	(150,000)	(100,000)	(75,000)	(25,000)	
Interest paid, and received, net	3,495	1,238	2,000	990	2,466	6,694	2,574	650	650	650	4,524	4,976	5,474	6,021	6,624	7,286	8,015	8,816	9,698	10,667	11,734	12,907	
Income taxes paid	(8)	(11)	(77)	(67)	88	(57)	(1,243)	0	0	0	(1,243)	0	0	0	0	0	0	0	0	0	0	0	
<b>Net cash provided (used) by Operating activities</b>	<b>(142,466)</b>	<b>(71,698)</b>	<b>(70,042)</b>	<b>3,470,498</b>	<b>(120,141)</b>	<b>3,208,617</b>	<b>(68,874)</b>	<b>(148,951)</b>	<b>(164,602)</b>	<b>29,221</b>	<b>(353,206)</b>	<b>(159,229)</b>	<b>300,435</b>	<b>876,197</b>	<b>1,785,216</b>	<b>1,569,025</b>	<b>1,978,242</b>	<b>1,958,689</b>	<b>2,084,904</b>	<b>2,202,455</b>	<b>2,267,427</b>	<b>2,353,060</b>	
<b>Cash flows from Investing activities:</b>																							
Purchases of property and equipment	(10,392)	(2,103)	(2,930)	(12,289)	(5,063)	(22,385)	(2,866)	(6,000)	(6,000)	(6,000)	(20,866)	(22,953)	(25,248)	(27,773)	(30,550)	(33,605)	(36,965)	(40,662)	(44,728)	(49,201)	(54,121)	(59,533)	
Purchase of and expenditure in intangible fixed assets	(3,325)	(1,201)	(2,334)	(1,930)	(17,635)	(23,300)	(10,159)	(1,000)	(1,000)	(1,000)	(13,159)	(13,617)	(14,506)	(15,233)	(15,995)	(16,795)	(17,634)	(18,516)	(19,442)	(20,414)	(21,435)	(22,506)	
Proceeds from disposal of p.p. and e	1	1	1	(1)	(1)	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	
Increase in/ Interest received/ Decrease in financial investments; and decrease in rest	0	0	0	0	(3,718,880)	(3,718,880)	945,334	0	0	0	945,334	0	0	0	0	0	0	0	0	0	0	0	
Acquisition of available-for-sale financial assets	(4,559)	(177)	0	0	0	(177)	(2,669)	0	0	0	(2,669)	0	0	0	0	0	0	0	0	0	0	0	
Proceeds from sale of available-for-sale financial assets	2,361	82	0	0	0	82	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	
<b>Net cash provided (used) by Investing activities</b>	<b>(15,914)</b>	<b>(3,398)</b>	<b>(5,263)</b>	<b>(14,220)</b>	<b>(3,741,779)</b>	<b>(3,764,660)</b>	<b>929,640</b>	<b>(7,000)</b>	<b>(7,000)</b>	<b>(7,000)</b>	<b>908,640</b>	<b>(36,770)</b>	<b>(39,755)</b>	<b>(43,006)</b>	<b>(46,545)</b>	<b>(50,399)</b>	<b>(54,600)</b>	<b>(59,178)</b>	<b>(64,170)</b>	<b>(69,615)</b>	<b>(75,556)</b>	<b>(82,039)</b>	
<b>Cash flows from Financing activities:</b>																							
Repayment of obligations under finance leases and other debts	(5)	(1,248)	(896)	(1,690)	(1,257)	(5,091)	(1,425)	0	0	0	(1,425)	0	0	0	0	0	0	0	0	0	0	0	
Proceeds from capital and share premium increases, gross amount	296,188	0	0	960,087	0	960,087	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	
Issue costs paid related to capital and share premium increases	(15,964)	0	0	0	(4,447)	(4,447)	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	
Proceeds from capital and share premium increases from exercise of warrants	7,657	3,481	4,324	6,675	370,722	385,202	5,355	0	0	0	5,355	0	0	0	0	0	0	0	0	0	0	0	
<b>Net cash provided (used) by Financing activities</b>	<b>287,876</b>	<b>2,233</b>	<b>3,428</b>	<b>965,072</b>	<b>365,018</b>	<b>1,335,751</b>	<b>3,930</b>	<b>0</b>	<b>0</b>	<b>0</b>	<b>3,930</b>	<b>0</b>	<b>0</b>	<b>0</b>	<b>0</b>	<b>0</b>	<b>0</b>	<b>0</b>	<b>0</b>	<b>0</b>	<b>0</b>	<b>0</b>	
Effect of exchange rate differences on cash, cash equivalents; and transfer to curr fina	10,089	4,968	(3,101)	30,514	(241,263)	(208,888)	17,261	0	0	0	17,261	18,124	19,030	19,982	20,981	22,030	23,131	24,288	25,502	26,777	28,116	29,522	
<b>Net increase (decrease) in Cash/Cash Equivalents</b>	<b>139,585</b>	<b>(67,895)</b>	<b>(74,978)</b>	<b>4,451,864</b>	<b>(3,738,171)</b>	<b>570,820</b>	<b>881,957</b>	<b>(155,951)</b>	<b>(171,602)</b>	<b>22,221</b>	<b>576,625</b>	<b>(177,874)</b>	<b>279,709</b>	<b>853,173</b>	<b>1,759,652</b>	<b>1,540,655</b>	<b>1,946,773</b>	<b>1,923,799</b>	<b>2,046,237</b>	<b>2,159,618</b>	<b>2,219,888</b>	<b>2,300,543</b>	
<b>Cash &amp; Cash Equivalents at Beginning of Period</b>	<b>1,151,211</b>	<b>1,290,796</b>	<b>1,222,901</b>	<b>1,147,923</b>	<b>5,599,787</b>	<b>1,290,796</b>	<b>1,861,616</b>	<b>2,743,573</b>	<b>2,587,622</b>	<b>2,416,019</b>	<b>1,861,616</b>	<b>2,438,241</b>	<b>2,260,366</b>	<b>2,540,076</b>	<b>3,393,249</b>	<b>5,152,901</b>	<b>6,893,556</b>	<b>8,640,330</b>	<b>10,564,129</b>	<b>12,610,366</b>	<b>14,769,984</b>	<b>16,989,972</b>	
<b>Cash &amp; Cash Equivalents at End of Period</b>	<b>€ 1,290,796</b>	<b>€ 1,222,901</b>	<b>€ 1,147,923</b>	<b>€ 5,599,787</b>	<b>€ 1,861,616</b>	<b>€ 1,861,616</b>	<b>€ 2,743,573</b>	<b>€ 2,587,622</b>	<b>€ 2,416,019</b>	<b>€ 2,438,241</b>	<b>€ 2,438,241</b>	<b>€ 2,260,366</b>	<b>€ 2,540,076</b>	<b>€ 3,393,249</b>	<b>€ 5,152,901</b>	<b>€ 6,893,556</b>	<b>€ 8,640,330</b>	<b>€ 10,564,129</b>	<b>€ 12,610,366</b>	<b>€ 14,769,984</b>	<b>€ 16,989,972</b>	<b>€ 19,290,515</b>	

Source: H.C. Wainwright &amp; Co. estimates.





## Cash Outlook, Ratio Analysis, and Enterprise Value

CASH QUICK LOOK :	2018A	1Q19A	2Q19A	3Q19A	4Q19A	2019A	1Q20A	2Q20E	3Q20E	4Q20E	2020E	2021E	2022E	2023E	2024E	2025E	2026E	2027E	2028E	2029E	2030E	2031E
Cash burn in period (cash from operations)	€ 362,654	€ 94,161	€ 111,957	€ 153,323	€ 166,157	€ 525,598	€ 151,501	€ 170,586	€ 185,238	€ 178,599	€ 685,924	€ 754,516	€ 829,968	€ 912,965	€ 958,613	€ 987,371	€ 1,016,993	€ 1,047,502	€ 1,068,452	€ 1,081,534	€ 1,094,795	€ 1,108,238
Total cash and cash equivalents	€ 1,290,796	€ 1,222,901	€ 1,147,923	€ 5,599,787	€ 5,780,832	€ 5,780,832	€ 5,722,378	€ 5,566,427	€ 5,394,824	€ 5,417,046	€ 5,417,046	€ 5,239,171	€ 5,518,881	€ 6,372,054	€ 8,131,706	€ 9,672,361	€ 11,619,135	€ 13,542,934	€ 15,589,171	€ 17,748,789	€ 19,988,777	€ 22,269,320
Periods of cash remaining	3.5 yrs	13.0 qts	10.5 qts	36.5 qts	35.0 qts	11.0 yrs	38.0 qts	32.5 qts	29.0 qts	30.5 qts	8.0 yrs	7.0 yrs	6.5 yrs	7.0 yrs	8.5 yrs	10.0 yrs	11.5 yrs	13.0 yrs	14.5 yrs	16.5 yrs	18.0 yrs	20.0 yrs
Ratio analysis	2018A	1Q19A	2Q19A	3Q19A	4Q19A	2019A	1Q20A	2Q20E	3Q20E	4Q20E	2020E	2021E	2022E	2023E	2024E	2025E	2026E	2027E	2028E	2029E	2030E	2031E
EBIT	(44,809)	(53,242)	(44,367)	490,631	(22,730)	370,292	(44,585)	(66,651)	(81,302)	110,521	(82,017)	119,960	570,153	1,135,648	1,816,857	2,061,754	2,545,424	2,501,966	2,645,740	2,711,551	2,738,748	2,758,177
EBITDA	(40,618)	(50,484)	(41,472)	493,815	(19,480)	382,379	(40,396)	(63,151)	(77,802)	114,021	(67,328)	137,587	588,661	1,155,081	1,837,262	2,083,179	2,567,921	2,525,588	2,670,543	2,737,594	2,766,093	2,786,889
EBITDA % of Sales	-13%	-123%	-61%	77%	-14%	43%	-38%	-61%	-75%	39%	-11%	15%	39%	61%	63%	66%	65%	66%	65%	66%	66%	66%
EV/EBITDA multiple	(72x)	(88x)	(122x)	6x	(564x)	29x	(223x)	(175x)	(144x)	99x	(167x)	83x	19x	9x	5x	3x	2x	1x	0x	(0x)	(1x)	(2x)
Gross Profit Margin	100%	100%	100%	100%	100%	100%	100%	100%	100%	100%	100%	100%	96%	93%	93%	92%	92%	91%	91%	91%	91%	91%
OpMargin	-14%	-130%	-66%	76%	-16%	41%	-42%	-64%	-78%	38%	-14%	13%	38%	51%	60%	62%	65%	64%	65%	65%	65%	65%
Net Operating Profit After Tax (NOPAT)	(44,858)	(53,310)	(44,428)	507,459	(39,643)	370,078	(44,921)	(66,651)	(81,302)	110,521	(82,353)	119,960	570,153	1,135,648	1,816,857	1,649,973	2,036,938	2,002,202	2,117,253	2,169,935	2,191,726	2,207,306
Free Cash Flow (FCF)	(152,858)	(73,801)	(72,972)	3,458,209	(125,204)	3,186,232	(71,740)	(154,951)	(170,602)	23,221	(374,072)	(182,182)	275,167	848,424	1,754,666	1,535,420	1,941,276	1,918,028	2,040,176	2,153,254	2,213,306	2,293,527
Book per share	€ 23.25	€ 21.53	€ 20.86	€ 40.92	€ 46.32	€ 47.83	€ 43.81	€ 42.82	€ 41.62	€ 43.37	€ 43.41	€ 45.72	€ 54.96	€ 72.88	€ 101.18	€ 126.92	€ 158.54	€ 189.62	€ 222.44	€ 256.06	€ 290.03	€ 324.27
Net cash per share	€ 24.72	€ 22.39	€ 20.94	€ 90.39	€ 29.99	€ 30.97	€ 42.33	€ 39.84	€ 37.12	€ 37.39	€ 37.43	€ 34.63	€ 38.84	€ 51.78	€ 78.47	€ 101.73	€ 131.05	€ 159.91	€ 190.51	€ 222.69	€ 255.65	€ 289.69
Return on assets (ROA)	-2%	-3%	-3%	6%	-2%	2%	-1%	-1%	-2%	2%	-2%	2%	10%	16%	21%	16%	16%	14%	13%	11%	10%	9%
Return on equity (ROE)	-2%	-4%	-4%	14%	-4%	5%	-2%	-3%	-3%	4%	-4%	4%	16%	24%	27%	25%	19%	16%	14%	13%	11%	10%
Current ratio	6.04	6.31	6.40	8.95	10.26	10.26	9.28	8.73	8.19	7.95	7.95	6.73	6.27	6.43	7.32	7.88	8.61	9.15	9.64	10.07	10.43	10.73
Enterprise Value (MC + Total Debt - Cash)	2,916,600	4,451,576	5,071,758	2,920,723	10,977,923	10,977,923	8,988,670	11,032,112	11,230,954	11,236,027	11,222,407	11,427,603	11,175,269	10,349,527	8,617,360	7,104,245	5,185,068	3,288,919	1,270,389	(861,468)	(3,053,639)	(5,326,309)
Market Cap (MC)	4,207,396	5,674,477	6,219,681	8,520,510	12,839,539	12,839,539	11,732,243	13,619,734	13,646,973	13,674,267	13,660,648	13,687,969	13,715,345	13,742,776	13,770,261	13,797,802	13,825,397	13,853,048	13,880,754	13,908,516	13,936,333	13,964,205
Current Share price €	209.70																					

Source: H.C. Wainwright &amp; Co. estimates.